<u>GOVERNMENT OF TELANGANA</u> <u>FISHERIES DEPARTMENT</u>

From,

R.Laxminaryana, M.Sc., HDCM, District Fisheries Officer, Medak District To,

The In-Charge Department of Zoology, Tara Govt. College Sangareddy District

Lr.No.05/Fish/Train/2017

Dated:18-09-2017.

Sir,

Sub:- Fisheries – Memorandum of understanding with Dept. of Zoology, Tara Govt. College, Sangareddy - Reg
Ref:- In-Charge, Dept. of Zoology, Lr.No.474/TGS-SRD/2017, Dt. 18-09-2017

With reference to the subject, I am glad to sing a Memorandum of understanding with Department of Zoology, to give training in the following topics related to Fisheries sector. The training would be given to B.Sc.(BZC) III year students every year a batch of (2°) students in the following activities.

1. Maintenance of different types of ponds Nursury, Rearing and Stocking.

2. Preservation.

3. Process of induced breeding

4. Identifying the diseases fishes

5. Transport and Marketing

Yours faithfully

District Eisheries Officer N: Medal District

TARA GOVERNMENT COLLEGE (AUTONOMOUS)

SANGAREDDY

The following students were selected for training on following techniques as undertaken in the MoU with Department of Fisheries, Medak district.

- 1. Maintenance of different types of ponds, nursery, rearing and stocking techniques.
- 2. Preservation

1 . 18

- 3. Process of Induced breeding.
- 4. Identifying the diseases in fishes.
- 5. Transport and Marketing.

S.No	Name of the student	Roll No.	Course			
1	A.Rajeshwari	605815445503	BZC III Year			
2	A.Ravinder	605815445506	BZC III Year			
3	B.Sandhya	605815445511	BZC III Year			
4	B. Shashirekha	605815445515	BZC III Year			
5	B. Rekha	605815445503	BZC III Year			
6.	Ch. Shailaja	605815445516	BZC III Year			
7.	Ch. Vittal	605815445517	BZC III Year			
8.	D. Ananthalaxmi	605815445523	BZC III Year			
9.	E. Jyothi	605815445524	BZC III Year			
10.	G.Ravi	605815445527	BZC III Year			
11.	H. Harish	605815445530	BZC III Year			
12.	J. Srikanth	605815445532	BZC III Year			
13.	K. Sangeetha	605815445540	BZC III Year			
14.	K. Bhavani	605815445543	BZC III Year			
15.	M. Mamatha	605815445546	BZC III Year			
16.	M. Jayamma	605815445554	BZC III Year			
17.	M. Esther Rani	605815445555	BZC III Year			
18.	T. Premalatha	605815445585	BZC III Year			
19.	T. Sumathi	605815445586	BZC III Year			
20	A.Brahmachary	605813445502	BZC III Year			

Department of Zoology Tara Govt. College Sangareddy - 502 001

Government of Telangana Commissionerate of Collegiate Education

Attendance Certificate

This is to certify that Sri Shankar Naik, Assistant Professor of Public Administration, Tara GDC, Sangareddy has attended duty at O/o CCE from **18 to 27** January, **2018**. He was assigned to coordinate the Jignasa- State Level Presentations of Student Study Projects- 2017-18 as **Coordinator of Tour Committee**. His absence in the College shall be treated as **ON DUTY**.

the the state

For Commissioner of Collegiate Education

Hyderabad 21.01.2018



TELANGANA TRIBAL WELFARE RESIDENTIAL DEGREE COLLEGE (MEN) SANGAREDDY

LETTER OF APPRECIATION

This Certificate is awarded to

Dr. A.V. SHARMA, HOD, ENGLISH

In Appreciation for contributing your service towards the development of Tribal Students of TTWRDC(M) Sangareddy.

On behalf of the Students, We the Staff and The Principal express our Sincere Thanks.

TTWRDC (MEN) College Co

GOVT. DEGREE COLLEGE FOR WOMEN, SIDDIPET

\$

CERTIFICATE OF ACKNOWLEDGEMENT

This is to certify that Sri. A. Vishweshwara Sharma, Assistant Professor of English, Tara Government Degree College, Sangareddy has attended this college on 25/01/2020 as a Resource Person and delivered an Extensive talk on NAAC preparatory works. This certificate is issued as token of acknowledgement of his Service at our college.

Date : 25/01/2020, Place : SIDDIPET.

Govt. Degree College for Women SIL. IPET, Dist: Siddipet-502103.



GOVT. DEGREE COLLEGE FOR WOMEN, SIDDIPET, SIDDIPET DIST.

ATTENDANCE CERTIFICATE

Date: 31/01/2020

This is to certify that Sri A. Vishweshwara Sharma, Assistant Professor of English, at TARA GOVERNMENT COLLEGE SANGAREDDY(A) has attended the college from 28/01/2020 to 31/01/2020 to guide the Staff in its uploading of SSR for its first cycle.

The incumbent's services are exemplary. The undersigned expresses pleasure at the way the incumbent has extended his services. This Certificate has the reference of the commissioner of collegiate Education, Telangana State Proc. File No. CCE-AC/QLTY/NAAC/2/2018-ACADEMIC CELL

· S. Ping

Govt Degree College for Women LSIL SET, Dist: Siddipet-502103.

File No.CCE-AC/QLTY/NAAC/2/2018-ACADEMIC CELL

PROCEEDINGS OF COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA PRESENT: SRI.NAVIN MITTAL, IAS.

Sub:- Accreditation of GDCs-New system of Assessment by NAAC- Status of preparation for NAAC Reaccreditation-Visits to certain GDCs-Reg. Ref:- CCE-AC/QLTY/NAAC/2/2018-ACADEMIC CELL

Vide reference read, certain GDCs due for NAAC reaccreditation have submitted the status of preparation to the Commissioner of Collegiate Education. In this connection, GDC (W) Siddipet has submitted IIQA and is preparing SSR for fresh accreditation.

In view of the above, Principal, GDC (W) Siddipet has requested Commissioner, Collegiate Education for the guidance of Sri.A.Vishweswara Sharma, Asst. Professor of English, GDC (A) Sangaredy, in the submission of data on College website from 28th to 31st January 2020.

In this connection, Principal, GDC (A) Sangareddy is informed that the absence of above mentioned faculty shall be considered as ON DUTY.

The faculty visiting the college have to submit the report on the actual status of preparation in respective GDC on or before 03.02.2020.

Signature valid

Digitally signed by PODILA SALA BHASKAR Date: 2020.01.31 10:49.50 IST Reason: Approved

For Commissioner of Collegiate Education

То

The Principals of GDCs concerned

2.2020

Govt. Model Degree College

Narayankhed, Dist : Sangareddy



Date: 15/02/2020

ATTENDANCE CERTIFICATE

This is to certify that Sri A. Vishweshwara Sharma, Asst Prof of English, Tara Govt. College (Autonomous) Sangareddy has delivered a lecture on *"Preparation of Records and SSR"* on 15-02-2020 at Govt. Model Degree College- Narayankhed, Has part of the Orientation programme conducted by the IQAC of the College. His class is excellent.

Principal Principal Govt Degree College Narayankhed Sangareddy Dist 502288 T S



TO WHOM SO EVER IT MAY CONCERN

This is to certify that A. Vishweshwara Sharma, Assistant Professor of English, Tara Govt. College, Sangareddy (A) has attended the Confidential work at O/o TSPSC, Nampally, Hyderabad, Telangana State for three (3) days i.e., from 23/11/2020 to 25/11/2020.

This is for information.

2020 5/11

ASSISTANT SECRETARY TSPSC: HYDERABAD.

Enter to Learn

Leave to Serve



R & B.G.N.R. Government Arts & Science College

(An Autonomous College under the Jurisdiction of Kakatiya University - Warangal) Re-Accredited at B⁺⁺ by NAAC, Bengaluru & an ISO 9001:2015 Certified Institution

> KHAMMAM - 507 002, TELANGANA STATE Estd: 1956

Mr. K.S.S. RATNAPRASAD., M.A., PGDTE. Principal (FAC) Office: 08742-298876, Mobile: 91548 06767, E-mail: <u>khammamsrbgnr.jkc@gmail.com</u>

CERTIFICATE

This is to certify that, Sri. A. Vishweshwara Sharma, Assistant Professor of English, the Dept. of English, TARA Government College (Autonomous), Sangareddy, has delivered an Extension Lecture on "QUALITATIVE & QUANTITAIVE METRICS OF NAAC" a one-day National Webinar, through Google Meet, the on-line platform as a Resource Person on 29-01-2021, hosted by the IQAC of our College.

his

The Participants are enriched with your Aura of insights & Speech

PRINCIPAL, Chairperson of the Webinar PRINCIPAL (FAC) SR&BGNR Govt. Arts&Science College (A), Khammam

0/10/120 02/2024

Organizers: The College IQAC



Certificate of Appreciation

This is to certify that Sri A. Vishweshwara Sharma, who has been working as an Assistant Professor of English, delivered an extension lecture on "Gender Sensitisation" to the stakeholders of Mahitha Sangareddy on 2 February 2021. The programme was an initiative of orientation programmes for the staff. The incumbent has got good feedback as his lecture was termed exemplary.

Signat

Head Office: #12-13-484/21, Street No.1, Tarnaka, Secunderabad-17, Telangana, India Field Areas: Hyderabad, Adilabad, Nalgonda & Ranga Reddy Districts

Phone: +91-40-27151858/27156231, Email: mahitahyd2002@gmail.com, mahitahyd2002@yahoo.com Website: www.mahita.org | facebook Page: https://www.facebook.com/mahitango

A B V GOVERNMENT DEGREE COLLEGE JANGAON, DT. JANGAON

Accredidated by NAAC 'B' Grade

Affiliated to Kakatiya University, Warangal E-mail:Warangal.jgnjkc@gmail.com Accredidated by NAAC 'B' Grade Phone No. : 08716-220044

ATTENDANCE CERTIFICATE

Date: 07-01-2021

This is to certify that <u>Sri.A.Vishweshwara Sharma</u>, Assistant Professor of English TARA Government College(A), Sangareddy has conducted an orientation session to the faculty of the college on 07-01-2021 on **"NAAC PEER TEAM VISIT**" and gave valuable suggestions for the successful conduct of PEER TEAM VISIT scheduled on 03-02-2021 and 04-02-2021.

ncipal Principal

ABV Govt. Degree College JANGAON, Dist: JANGAON



GOVERNMENT DEGREE COLLEGE FOR WOMEN (Affiliated to Osmania University & Accredited by NAAC with 'B' grade) SANGAREDDY-502 001, SANGAREDDY (Dist.), T.S.

Dr. Humera Sayeed, M.A., Ph.D. Principal(FAC) Phone No: 918455 277312 E-mail:gdcwsrd.jkc@gmail.com

APPRECIATION CERTIFICATE

I am happy to announce that the services rendered by Sri. A.V.Sharma, *Asst. Prof. of English* of **Tara Government Degree College** (A), Sangareddy Telangana towards the "Motivation lecture on NAAC awareness" being organized by IQAC of our college on 9th February 2021. His contribution in this regard is highly appreciated. Since we are going to conduct such many more programmes in the future, we expect for the same kind of contribution and enthusiasm.

Covit. Derre & covies for (W)men, Saaganddody.



Dr. Humera Sayeed, M.A., Ph.D. Principal (FAC) Phone No: 918455 277312 E-mail:gdcwsrd.jkc@gmail.com

CERTIFICATE OF APPRECIATION

This is to certify that A. Vishweswara Sharma, Assistant Professor of English of TARA Government College Sangareddy (Autonomous) has attended as a Resource Person to deliver an Extension lecture on 09.02.2021 held at Government Degree College for Women, Sangareddy, and has given a talk on NAAC preparation. His services are appreciable.

PRINCIPAL

Govt. Desige College C



GOVERNMENT DEGREE COLLEGE FOR WOMEN



(Affiliated to Osmania University & Accredited by NAAC with 'B' grade) SANGAREDDY-502 001, SANGAREDDY (Dist.), T.S.

Dr. Humera Sayeed, м.А., Ph.D. Principal(FAC) Phone No: 918455 277312 E-mail:<u>gdcwsrd.jkc@gmail.com</u>

To The Principal, TARA Govt. Degree College, Sangareddy, Dist. Sangareddy.

Respected Sir/Madam,

Sub: GDC(W) Sangareddy-NAAC-Awareness program- A.V.Sharma -Resource person -Request-Regarding.

With reference to the subject cited above, as we are going for NAAC this year, we would like to organize an awareness program for our faculty on 9th February 2021 and we would like to invite A.V.Sharma, Asst. Prof. of English as a resource person. Hence, I humbly request you to permit A.V.Sharma to be the resource person.

Looking forward to hearing from you soon.

Thanking you,

Yours faithfully,

PRINC Govt. De or Women. ollege (W) Govte ngareddy.



GOVERNMENT DEGREE COLLEGE FOR WOMEN



(Affiliated to Osmania University & Accredited by NAAC with 'B' grade)

SANGAREDDY-502 001, SANGAREDDY (Dist.), T.S.

Dr. Humera Sayeed, M.A., Ph.D. Principal(FAC)

Phone No: 918455 277312 E-mail:<u>gdcwsrd.jkc@gmail.com</u>

To The Principal, TARA Govt. Degree College, Sangareddy, Dist. Sangareddy.

Respected Sir/Madam,

Sub: GDC(W) Sangareddy-NAAC-Awareness program- A.V.Sharma -Resource person -Request-Regarding.

With reference to the subject cited above, as we are going for NAAC this year, we would like to organize an awareness program for our faculty on 9th February 2021 and we would like to invite A.V.Sharma. Asst. Prof. of English as a resource person. Hence, I humbly request you to permit A.V.Sharma to be the resource person.

Looking forward to hearing from you soon.

Thanking you,

Yours faithfully,

Govt. De or Women. Govt e College (W) gareddy.



ch-00 MANISHA SABOO

AVP- DELIVERY HEAD & DC HEAD OF HYD SEZ

NIVAS P C S

LEAD LEARNING & DEVELOPMENT, ETA

CERTIFICATE OF PARTICIPATION

THIS IS PRESENTED TO A Vishweshwara Sharma

Congratulations for successfully completing Faculty Development Enablement Program organized by Infosys Limited from 22-24 February 2021



GOVERNMENT DEGREE COLLEGE (W) SANGAREDDY



CERTIFICATE OF APPRECIATION

14 Aug 2021

This is to certify that Mr. A. Vishweshwara Sharma, Assistant Professor of English of TARA Government college, Sangareddy (A), delivered an extension lecture on "Holistic and Multi-disciplinary education" to the staff and students on 14 August 2021. The services of the resource person are highly appreciable and we wish him very best in his endeavour.

The webinar was organized by UGC committee of the college. The lecture was organized as part of the UGC's initiative 'One year of Transformative Reforms under NEP,2020 (D.O.No.F.1-3/2021 (QIP) dated 29 July 2021).

Convener

e (W



GOVERNMENT DEGREE COLLEGE SADASIVPET CERTIFICATE OF APPRECIATION



06 Aug 2021

This is to certify that Mr. A. Vishweshwara Sharma, Assistant Professor of English of TARA Government college, Sangareddy (A), delivered an extension lecture on "Holistic and Multi-disciplinary education" to the staff and students on 02 August 2021. The services of the resource person are highly appreciated and we wish him very best in his endeavour.

The webinar was organized by the Department of Chemistry of the college. The lecture series was organized as part of the UGC's initiative 'One year of Transformative Reforms under NEP,2020 (D.O.No.F.1-3/2021 (QIP) dated 29 July 2021).

onvener

Principal PRINCIPAL Govt. Degree College Sadasivpet, Dist. Sangareddy.



BJR Government Degree College Narayanaguda, Hyderabad (

(Affiliated to Osmania University) An ISO 9001: 2015 certified college

Certificate of Appreciation

This is to certify that Sri A. Vishweswara Sharma, Assistant Professor of English at Tara Government College Sangareddy (A), has delivered an extension lecture on "Conversation Skills" for UG semester II students on 25 September 2021. The extension lecture was organized by the Department of English. The lecture has got good feedback.

This certificate is issued to him as a token of appreciation for his services to the students and college. We wish him best in his endeavour.

In-Charge

Department of English

Secoly Jaman

Principal PRINCIPAL BJR Govt. Degree College Vittalwadi, Narayanguda, Hyderabad, T.S.-29.



BJR Government Degree College Narayanaguda, Hyderabad

(Affiliated to Osmania University)

An ISO 9001: 2015 certified college

Certificate of Appreciation

This is to certify that Sri A. Vishweswara Sharma, Assistant Professor of English at Tara Government College Sangareddy (A), has conducted a workshop on "NAAC AWARENESS" for teaching staff of the college on 25 September 2021. The workshop was organized by Internal Quality Assurance Cell (IQAC) of the college. The incumbent has got excellent feedback.

This certificate is issued to him as a token of appreciation for his services to the stakeholders of the college.

Internal Quality Assurance Cell

Geellydahan

Principal PRINCIPAL JR Govt. Degree College Vinsiwadi, Narayanguda, Hyderabed, T.S.-29.



GOVERNMENT DEGREE COLLEGE, TANDUR, VIKARABAD DISTRICT TELANGANA - 501141

(Estd : 2008 affiliated to Osmania University

Date: 22-02-2022

ATTENDANCE CERTIFICATE

This is to certify that A.Vishweshwara Sharma, Assistant Professor of English, Tara Government Degree College, Sangareddy has attended Government Degree College, Tandur, Vikarabad District for giving training to staff on NAAC procedures on 22-02-2022.

PRINCIPAL

KINCIPAL t. Degree College idur, Vikarabad Dist Pin Code: 501141 \bigcirc

us.

GOVERNMENT DEGREE COLLEGE FOR WOMEN (Affiliated to Osmania University & Accredited by NAAC with 'B' grade)

SANGAREDDY-502 001, SANGAREDDY (Dist.), T.S.

Dr. Humera Sayeed, M.A., Ph.D. Principal (FAC) Mobile: 9154806840 E-mail:gdcwsrd.jkc@gmail.com

15th March 2022

Certificate of Appreciation

This is to certify that Sri. **A.Vishweshwara Sharma**, IQAC coordinator,Tara GDC(A),Sangareddy attended our college on 15th March 2022 as a resource person to address and guide the faculty of our college regarding NAAC preparation and impeccable presentation on. We are very glad with his presence and wholeheartedly appreciate the guidance provided to

Principal Govt. Degree College for Women Sangareddy- 502,001 (T.S)

<u>GOVERNMENT DEGREE COLLEGE, TANDUR,</u> <u>VIKARABAD DISTRICT</u> <u>TELANGANA – 501141</u>

(Estd : 2008 affiliated to Osmania University

Date: 22-02-2022

ATTENDANCE CERTIFICATE

This is to certify that A.Vishweshwara Sharma, Assistant Professor of English, Tara Government Degree College, Sangareddy has attended Government Degree College, Tandur, Vikarabad District for giving training to staff on NAAC procedures on 22-02-2022.

PRINCIPAL

MKOAL JUNCIPAL JVI. Degree College andur, Vikarabad Dist Pin Code: 501141



TELANGANA SOCIAL WELFARE RESIDENTIAL DEGREE COLLEGEOF COMMERCE FOR WOMEN SANGAREDDY AT BUDHERA



CERTIFICATE OF APPRECIATION

This is to certify that Sri.A.Vishweshwara Sharma, Asst.Prof of English, at Tara Government College Sangareddy (A), has attended the college on 21.06.2022 in the capacity of Resource Person for the Faculty Development Programme Conducted for staff of the college on NAAC Accreditation.

The Resource Person got excellent feedback. We thank him for his services.

NGAREDD gereddy Dist.

GOVERNMENT OF TELANGANA PROCEEDINGS OF THE RETURNING OFFICER, 39-SANGAREDDY ASSEMBLY CONSTITUENCY AND REVENUE DIVISIONAL OFFICER, SANGAREDDY.

PRESENT: SRI S.SREENU

Proc. No. C/17020/2018

Dated:23-11-2018.

Sub: - GENERAL ELECTIONS 2018 – General Elections toTSLA in 39-Sangareddy Assembly Constituency– Allotment of various duties at distribution and reception centers – Orders – Issued.

ORDER:

In view of the General Elections to 39-Sanbgareddy Assembly Constituency the persons shown in the Annexure – I are made in charges for the counters shown against their names for the distribution of materials to the polling personnel and the persons shown in Annexure – II are made in charges for the counters noted against their names for receipt of material from the polling personnel. The details of Polling stations allotted to each counter are noted against their names.

All the staff members shown in Annexure-I & II should be present at the Distribution centre i.e., Dr.B.R.Ambedkar Stadium Grounds, Sangareddy on **05-12-2018 at 11-00 AM** before the undersigned, check the material and will present at distribution centre on **06-12-2018** at 7.00 AM attend and see that the material allotted to the polling stations of their counters should be issued to the polling personnel concerned. On **07-12-2018** after completion of poll the personnel and material will be received at reception centre. Therefore the staff drafted in Annexure – II should report before the undersigned on **07-12-2018** by 4.00 PM at reception centre i.e., Dr.B.R.Ambedkar Stadium Grounds, Sangareddy and receive the material from the polling personnel of the polling stations allotted to them. The acknowledgement of the Electronic Voting Machines, VVPATs and other material should be issued to Polling Personnel after thorough verification.

Any deviation in the matter not only liable for disciplinary action under C.C.A Rules, but also offence under Election Law.

39-Sangareddy Assembly Constituency & Revenue Divisional Officer, Sangareddy.

Encl: (2)

То

All Concerned.

Copy submitted to the District Collector & District Election Officer, Sangareddy for information.

	Hand	ds on experience
5.	 K.Sreedhar, Asst. Prof. Tara Govt.College, Srd. A.Vishweshwar Sharma, Asst. Prof. Tara Govt.College, Srd. Govindaram Principal, GJC(B) Srd. 	Display of the DVD and training to Polling Personnel on usage of EVM.
	VEHIC	LE IN-CHARGES
6.	 Balaraju, DT RDO's Office , Sangareddy. Srinivas, Sr.Asst. T.O Sangareddy. Jawan Narsimulu, VRO Sangareddy. Srikanth, VRA, Ismailkhanpet. 	Allotment of Busses Route wise and pasting routes on wind shield and all other officers of Election duty, obtaining trip sheets, maintenance of all accounts, providing of fuel etc., and all other connected work of vehicles till relieved.
7.	 Vishnu Dy.Tahr O/o T.O. S.S.Pet. Yadaiah, VRO Mollaiah, VRO Mollaiah, VRO Aizaj, Girdavar-I Sangareddy. Harichandra Prasad, Girdavar-II, Sadasivapet 	Obtaining the attendance & correspondence of Micro observers , web casting and video graphers allotment of PS wise

Si. No	Name of the in-charge		Route No.	Polling stations allotted	No.of P.S
1	Srinivas, Panchayath Secretary, Kondapur.	Incharge for Statutory material			12
	Nagaraju, VRO, Sadasivapet. (2) V.R.As.	Incharge for Non- Statutory material	I	19, 190, 18, 16, 12, 13, 14, 15, 3, 1, 2, 4	
2	Kum.Jyotshna, Girdavar-I, T.O. Sadasivapet.	Incharge for Statutory material		198, 199, 200, 197,	11
	Naveen, VRO Sadasivapet. (2) V.R.As.	Incharge for Non- Statutory material	п	196,195,193, 194, 191, 192, 17	
3	Khaja, Panchayath Secretary, MPP Kondapur.	Incharge for Statutory material		174, 186, 187, 188,	12
	Chandraleela, Panchayath Secretary, Girmapur MPP Kondapur.	Incharge for Non- Statutory material	ш	189,201,202, 203, 204, 205, 206, 239	

6. LOGISTIC ARRANGEMENT AT RECEIPTION CENTER:

- 1. Varaprasad, Girdavar-II, Sangareddy along with (10) Village Servants and arrangements of Shamiyanas, Sound System, Dias, Furniture and canteen etc.,
- 2. Santosh, V.R.O. Kalvakunta T.O. Sangareddy.
- 3. Shivaram, V.R.O. Fasalwadi, T.O. Sangareddy.
- 4. Yadagiri, V.R.A. Sangareddy.
- 5. Rajashekar, V.R.A. Sangareddy.

7.DRINKING WATER & SANITATION:

- 1. S/Sri Prasad, Muncipal Commissioner, Sangareddy.
- 2. Sri Vijay Babu, Sanitary Inspector Srd.Municipality.
- 3. Sri Vittal, Srd.Municipality and Sanitary workers for maintaining cleanliness Of the Ground and also arranging continous Drinking water to the polling persons.

8. GENERAL DUTIES AND REPORT:

- 1. Sri Aroundaya Chary, EPIC DT, T.O. Sanagareddy.
- 2. Sri Suresh Kumar, DT(CS) Sangareddy.
- 2. Sri Kartik, Sr.Asst. R.D.O office, Sangareddy.
- 3. Sri Srinivas, VRO, Sangareddy.
- 4. Sri Chandra Shekar, HHRP, RDO office, Sangareddy.
- 5. Sri Chandu, VRA, T.O. Sangareddy.
- 6. Sri Sameer V.R.A. Kandi.
- 7. Sri Srinivas, V.R.A. Kandi.
- 8. Sri Murali, VRA, Sangareddy.
- 9. Sri Vinod Kumar Chary, Comp.Ope. Kondapur

Officer, 39 - Sangareddy Assembly Constituency & Revenue Divisional Officer, Sangareddy.

To

The concerned.

Copy submitted to the District Election Officer & District Collector, Sangareddy for favour of kind information.



ST.ANN'S COLLEGE FOR WOMEN



(Autonomous) Osmania University NAAC Reaccredited with 'A+' Grade (3rd Cycle) College with Potential for Excellence by UGC MEHDIPATNAM, HYDERABAD

This is to certify that

A. Visburesbura Sharma.

Maria Government college (A) (Sangareddy)

Participated in

One Day National seminar

on

"Corporate Academia Partnership - Fostering Innovation & Entrepreneurship"

Organized by Internal Quality Assurance Cell (IQAC), St. Ann's College for Women, Mehdipatnam, Hyderabad on 25th March 2019.

E

Dr. Sr. P. Amrutha Principal 64 Ann's College for Womon Mendipetnam, Hyd-26,

St. Pious X Degree & P.G. College For Women

(Re-Accredited by NAAC with A+ Grade)

In Collaboration with Commissionerate of Collegiate Education, Govt. of Telangana

State Level Workshop on "Mentoring the Mentors on Revised AQAR Format"

27th March, 2019

Certificate of Participation

of TARA Govt. College A Sanga Reddy has participated in a State Level Workshop on "Mentoring the Mentors on Revised AQAR Format" organized by 90.40, This is to certify that Mr. / Ht. Nichweshwaya Shanna

St. Pious X Degree & PG College for Women, Nacharam, Hyderabad

IQAC Coordinator

Principal

COMMISSIONERATE OF COLLEGIATE EDUCATION . Academic Cell, Telangana, Hyderabad

Attendance Certificate

E.S.M

For Commissioner of Collegiate Education

Hyderabad, 20- 0**8**- 2019

NAGARJUNA GOVERNMENT COLLEGE



(AUTONOMOUS)

(RE - ACCREDITED 'A' BY NAAC)

NALGONDA District, TELANGANA



ATTENDENCE CERTIFICATE

one day workshop on Faculty Development Workshop on Role Professor, Tara Degree College, Sanga Reddy has Conducted of IQAC & NAAC in Quality Sustenance on 05-09-2018 as a resource person at NAGARJUNA GOVERNMENT COLLEGE (A), This is to certify that Sri. A. Visweswara Sharma , Asst. NALGONDA.



COLLEGE (A)	aliation (196		of English.	on Role of IQAC & NAAC in Quality Sustance	on. 05-09-20/8	d in acknowledgment of his/her association		Principal Principal Principal Nagarjuna Govt. College (Autonomous) NALGONDA,
NAGARJUNA GOVERNMENT COLLEGE (A) (Re-Accredited 'A' by NAAC) (Re-Accredited 'A' by NAAC) Nalgonda District, Telangana state	Certificate of Apprec	This is to certify that Dr. Mr. Ms A. U. Shweawara. Showma	Professor/ Associate-Professor/Asst. Professor/ Lecturer	delivered lectrone on Role of IQAC &	This Programme was organized by the Department of $ZQAC$	The Presentation is informative and Impressive. This certificate has been issued in acknowledgment of his/her association	and participation with the workshop/Extension Lecture /Programme.	VV - Dry V. V. Suthandone In charge 12 Ac coordinatur



Website: www.gdcarmoor.in,e-mail ID: gdcarmoor@gmail.com,Phone:08463 - 222259 (2(f) & 12(B) of UGC) Reaccredited by NAAC with "B" grade GOVT. DEGREE COLLEGE, ARMOOR **Dist:** Nizamabad

ATTENDANCE CERTIFICATE

Government GDC (A) Sangareddy, attended as CCE nominee to asses the status of This is to certify that Sri . A.Vishweswara Sharma, Asst. Prof of English, NAAC works going on and gave guidance in regarding new NAAC accreditation CCE-AC/QLTY/NAAC/3/2018-ACADEMIC on 25.09.2019 at Government Degree College, Armoor, Nizamabad District. as per the proceedings of system CELL

PRINCIPAL PRINCIPAL Principal Govt Degree College ARMOOR-503 224

File No.CCE-AC/ODWS/4/2019-ACADEMIC CELL

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI.NAVIN MITTAL, I.A.S.

Sub: Collegiate Education-Tara GDC, Sangareddy-A.Vishweshwara Sharma, Assistant Professor of English-Served as a **Resource person** for a day on **11.10.2019** in three day Orientation Programme organized by TTWREIS for newly recruited TS Residential Degree College Lecturers-Sanction of **On Duty**-Orders-Regarding.

Ref: Letter received from Principal, Tara GDC, Sangareddy, dated: 13.10.2019.

Vide references read, the Commissioner of Collegiate Education has accorded On Duty to **A.Vishweshwara Sharma**, Assistant Professor of English, Tara GDC, Sangareddy served as a Resource person on **11.10.2019** in a three day Orientation Programme organized by Telangana Tribal Welfare Residential Educational Institutions Society (TTWREIS) for newly recruited TW residential Degree College Lecturers at Rajendranagar, Hyderabad.

The absence of the above said Assistant Professor shall be treated as **ON DUTY** on above said date. In view of the above, the Principal, Tara GDC, Sangareddy is informed to consider his absence in the college as **On DUTY**.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by PODILA BALA BHASKAR Date: 2019.10.17 10:55:16 IST Reason: Approved

For Commissioner of Collegiate Education

To The Principal of GDC concerned.

File No.CCE-AC/ODWS/4/2019-ACADEMIC CELL

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI.NAVIN MITTAL, I.A.S.

Sub: Collegiate Education-Tara GDC, Sangareddy-**A.Vishweshwara Sharma**, Assistant Professor of English-Served as a **Resource person** for a day on **11.10.2019** in three day Orientation Programme organized by TTWREIS for newly recruited TS Residential Degree College Lecturers-Sanction of **On Duty**-Orders-Regarding.

Ref: Letter received from Principal, Tara GDC, Sangareddy, dated: 13.10.2019.

Vide references read, the Commissioner of Collegiate Education has accorded On Duty to **A.Vishweshwara Sharma**, Assistant Professor of English, Tara GDC, Sangareddy served as a Resource person on **11.10.2019** in a three day Orientation Programme organized by Telangana Tribal Welfare Residential Educational Institutions Society (TTWREIS) for newly recruited TW residential Degree College Lecturers at Rajendranagar, Hyderabad.

The absence of the above said Assistant Professor shall be treated as **ON DUTY** on above said date. In view of the above, the Principal, Tara GDC, Sangareddy is informed to consider his absence in the college as **On DUTY**.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by PODILA BALA BHASKAR Date: 2019.10.17 10:55:16 IST Reason: Approved

For Commissioner of Collegiate Education

To The Principal of GDC concerned.

1.10.1

TELANGANA TRIBAL WELFARE UPGRADED RESIDENTIAL JUNIOR COLLEGE (B), JINNARAM, SANGAREDDY DISTRICT

This is to certify that Mr. A. VISHVESHWARA SHARMA, Asst prof of English at TARA GOVT.DEGREE COLLEGE SANGAREDY, has delivered an extension lecture on "GOAL SETTING" for the students of TTWURJC Jinnaram, on 02/11/2019. His lecture received the best response from the students.

Principal 21115 PRINCIPAL TTWUR JR.COLLEGE (Boys) JINNARAM-802 319. Sangareddy Dist.

GOVT. DEGREE COLLEGE FOR WOMEN, SANGAREDDY (Accredited by NAAC with B Grade) SANGAREDDY Dist. TELANGANA STATE - 502001

ATTENDANCE CERTIFICATE

Date: 19.11.2019

This is to certify that Sri.A.V. Sharma, Assistant Professor of English, Tara Govt College, Sangareddy has been delivered an extension Lecture on "Seven Criteria of NAAC" on 19.11.2019 organized by IQAC at Govt. Degree College for women, Sangareddy.

PRINCIPAL

6

Govt. Degree College (W) Sangareddy.

Phone : 9440379380 Email : gajwel.jkc@gmail.com



GOVERNMENT DEGREE COLLEGE GAJWEL

(Re-Accredited by NAAC with B) GAJWEL – 502278 Siddipet Dist. Telangana

Dr.A. Srinivas Reddy. M. Sc, Ph.D. Principal Date: 13.12.2019

CERTIFICATE OF APPRECIATION

This is to certify that **Mr. A. Vishveshwara Sharma**, Asst. Prof. of English, Tara Government Degree College (A) Dist. Sangareddy, has delivered a talk as Resource Person at **Orientation Programme on the Preparation of Self Study Report** organized by IQAC, GDC Gajwel on 13.12.2019 from 2.00 p.m. to 4.30 p.m.

His Talk has been appreciated immensely by the statt who attended the Orientation Programme.

Principal 1

PRINCIPAL Government Degree College Gajwel, Siddipet-Dist, Telangana.

Government Degree College for Women, Gajwel

Attendance Certificate

This is to state that Sri.A.V.Sharma, Asst. Prof of English, Tara Government Degree College, Sangareddy has conducted a workshop in our college on 13/12/2019 on Assessment and Accreditation process under revised NAAC framework.

PRINCIPAL 13/12/2019 Govt. Degree College for Women GAJWEL, Dist. Siddipet.

Date: 13/12/2019

N.M.GOVT. DEGREE COLLEGE, JOGIPET, SANGAREDDY DIST.

Present: Dr.H.R.ANITHA, M.Sc., Ph.D.

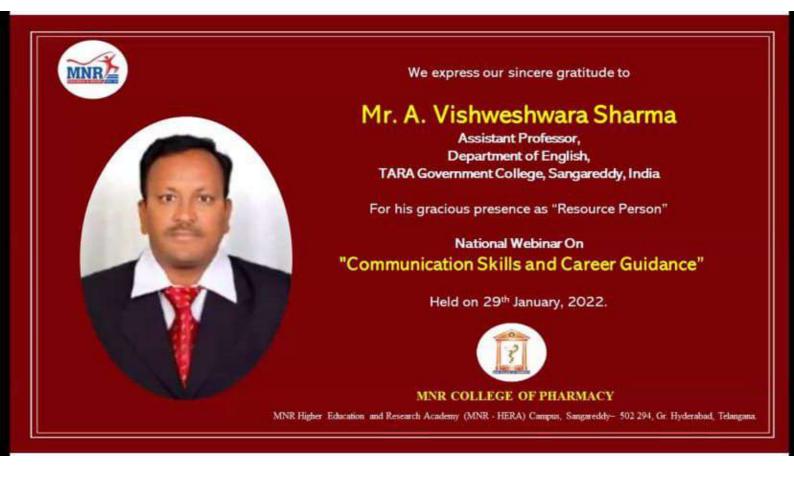
Date: 18-11-2019

ATTENDANCE CERTIFICATE

Sri. A.VISHWESHWARA SHARMA, Asst Professor of ENGLISH, Tara Govt. College, Sangareddy(A), has attended the college as a resource person for an extension lecture for II year students on "PRESENTATION SKILLS" on 18/11/2019 in the afternoon session.

The extension lecture was received with rapt attention by students.

N.M. Govt. Degree College,



SR and BGNR Government Arts and Science College (Autonomous) (Re-Accredited with 'B' Grade by NAAC) Khammam (Dist), Telangana

CERTIFICATE

This is to certify that Mr A. Vishweshwara Sharma, who has been working as an Assistant Professor of English at TARA Government College Sangareddy (Autonomous), Sangareddy district, has been serving as a Panel member on the Board of Studies (BoS) of the Department of English. He has been rendering his services to the board from the academic year 2018-19.

He may utilize this certificate for his API and other academic purposes.

Controller of Examinations

8.G.N

Controller of Examinations Controller of Examinations S.R. & B.G.N.R. Arts & Science College (Autonomous), KHAMMAM.

Tiztza Principa

S.R.& B.G.N.R. Govt. Arts & Science College (NAAC-B++) Autonomous-K H A M M-A*

SR and BGNR Government Arts and Science College (Autonomous) (Re-Accredited with 'B' Grade by NAAC) Khammam (Dist), Telangana

CERTIFICATE

This is to certify that Mr A. Vishweshwara Sharma, who has been working as an Assistant Professor of English at TARA Government College Sangareddy (Autonomous), Sangareddy district, has been serving as a Panel member on the Board of Studies (BoS) of the Department of English. He has been rendering his services to the board from the academic year 2018-19.

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Controller of Examinations Controller of Examinations S.R. & B.G.N.R. Arts & Science College (Autonomous), KHAMMAM.

Tiztza Principa

S.R.& B.G.N.R. Govt. Arts & Science College (NAAC-B++) Autonomous-K H A M M-A*

				Α	nnexure I								
				Commissionerat	e of Collegiate Education	1							
	T-SAT: Nipuna Live Telecast Schedule- 2019-20 11.00 AM to 11.45 AM												
S No	Date	Day	Day Subject	Year and Semester	Торіс	Name of the Presenter and Place of Work	Contact No.						
1	20.07.201 9	Saturda y	Commerce	I Year/I Sem	Bank Reconciliation statement	Tulja Bhavani, GDC Ibrhimpatnam	9100092589						
2	22.07.201 9	Monday	Telugu (Panel Discussion)	I Year/ I sem	Srujanatmaka Bodhana -Bhasha Naipunyaalu	1.Dr.J.Neeraja, O/o CCE TS Hyderabad 2.Dr.Seetha ramarao, GDC (A) khammam 3.Dr. Koyi.Koteswara Rao ,GDC (W) Nalgonda 4.Dr.R.Srinivas, TARA GDC(A) Sangareddy	7660020730 9866563519 9440480274 9848050694						
3	23.07.201 9	Tuesday	Mathematics	II Year /III Sem	Sequences, Limit of Sequences & Examples	D.Venkatesh, KGC,Hanamkonda	9948185643						
4	24.07.201 9	Wednsd ay	Zoology	I Year/I sem	Corals & Coral Formation	Dr.N.Raj Kumar , GDC Hayatnagar	9440697746						
5	25.07.201 9	Thursda y	History	I Year/I Sem	Mauryan Administration	M.Shanthaveni,IPGDC(W),Nampally	9440980955						
6	26.07.201 9	Friday	Computer Science	B.sc (II Year/III Sem)	Boolean Algebra	S.Yamuna Rani , GDC ,Malkajgiri	9908521900						
7	27.07.201 9	Saturda y	Special Live Program	All	Dial the Doctor / Nutritionist								
8	30.07.201 9	Tuesday	Physics	II Year/ III Sem	Kinetic Theory of Gases	B.Srinivas Goud, ABV GDC Jangaon	9959336467						
9	31.07.201 9	Wednsd ay	Botany (Pair teaching)	I Year/I Sem	Cyanobacteria:General Charecters, Bio	1.P.Hari Krishna,SRR GDC Karimnagar, 2.P.Balaraju SRR GDC Karimnagar	9966998858 9490664677						

					fertilisers, Type Studies		
10	01.08.201 9	Thursda y	Political Science	III Year/V Sem	Mao-Tse-Tung	P.Venkata Ramana , GDC Kukatpally	91770 87189
11	02.08.201 9	Friday	Gender Sensitization	All			
12	03.08.201 9	Saturda y	Commerce	III Year/ V Sem	Process Costing	M.Somaiah, KGC Hanamkonda	9948048949
13	05.08.201 9	Monday	Basic Communication	All	Listening Skills	Dr.E.Srinivasa Rao, ABV GDC Jangaon	9848788425
14	06.08.201 9	Tuesday	Chemistry	I Year /I Sem	Aromatic Hydrocarbons	A.Srinivas Reddy,GDC Jammikunta	9949905069
15	07.08.201 9	Wednsd ay	Zoology	I Year/I sem	Polymorphism in Siphanophora	B.Srinivas Reddy,GDC Hayatnagar	9493461555
16	08.08.201 9	Thursda y	Economics(Pair teaching)	I Year/ I Sem	Production Function	1.Dr.A.Venkatesham TARA GDC(A), Sangareddy 2.E.Pavani Govt.City College, Nayapul	9989942298 9959632449
17	13.08.201 9	Tuesday	Mathematics	I Year/I Sem	Lengths of Plane Curves	Dr.D.Pushpa, ABV GDC,Jangaon	9440553078
18	14.08.201 9	Wednsd ay	Biotechnology	II Year/III Sem	Electrophoresis and its application	K.Geetanjali,GDC Kukatpally	9701004242
19	16.08.201 9	Friday	Social Responsibility	All			
20	17.08.201 9	Saturda Y	Commerce	II Year/III Sem	Income from Salary	Dr.Md.Yakub, GDC Mulugu	7396473313
21	19.08.201 9	Monday	Modern Language- English	III Year	Obituary - A.K.Ramanujan	A.Vishweshwara Sharma, TARA GDC Sangareddy	9490932161
22	20.08.201 9	Tuesday	Physics	III Year /V Sem	Electrical Potentail	P.R.Ratan Kumar, TARA GDC(A) Sangareddy	9618125210

23	21.08.201 9	Wednsd ay	Zoology(Pair teaching)	I Year/I sem	Parasitic Adaptations in Helminthes	1.B.Suresh Kumar, SRR GDC Karimnagar 2.P.Raju,SRR GDC Karimnagar	9490580630 9963700000
24	22.08.201 9	Thursda y	Public Administration	I Year/I Sem	Bureaucratic Aproach- Max Weber	Dr.G.Narasimulu, GDC (M) Gajwel	9441301961
25	23.08.201 9	Friday	Computer Science	B.Sc (III Year/ V Sem)	Exception handling and Multithreading in Java	S.Yamuna Rani , GDC ,Malkajgiri	9908521900
26	26.08.201 9	Monday	Telugu(Pair teaching)	I Year/ I sem	Samvaranudi Tapassu	1.Dr.T.Krishnaiah ,NGC(A) Nalgonda 2.Dr. R.Srinivas,TARA GDC(A) Sangareddy	9704731346 , 9848050694
27	27.08.201 9	Tuesday	Chemistry	I Yr /I Sem	Solutions	Dr.A.Dayanand, Govt.City College(A),Nayapul	9492302186
28	28.08.201 9	Wednsd ay	Botany	II Year/III Sem	Bentham & Hooker System of Classification	S.Veeraiah , GDC(W), Nalgonda	9848546345
29	29.08.201 9	Thursda Y	History	I Year/II Sem	Reforms of Allauddhin Khilji	S.Ganapathi Rao,KGC ,Hanamkonda	9030403940
30	30.08.201 9	Friday	Bio Chemistry	I Year/I Sem	Classification & Propertiesof Aminoacids	A.Chandra Shekar,GDC(W),Karimnagar	9963871117
31	31.08.201 9	Saturda y	Special Live Program	All	Environment Protection		
32	03.09.201 9	Tuesda y	Mathematics (Student as Teacher Program)				
33	04.09.201 9	Wednsd ay	Microbiology (Student as Teacher Program)				
34	05.09.201 9	Thursda y	Political Science (Student as Teacher Program)				
35	06.09.201	Friday	ТЅКС	All			

	9						
36	07.09.201 9	Saturda y	<i>Commerce (Student as Teacher Program)</i>				
37	12.09.201 9	Thursda y	Economics (Student as Teacher Program)				
38	13.09.201 9	Friday	Health & Nutrition	All			
39	16.09.201 9	Monday	Telugu(Pair teaching)	II Year / IV Sem	Devarakonda Durgam	1.Dr.Belli Yadaiah, GDC Ramannapet 2.Dr.Mrudula , GDC Narsapur	9848392690 9441408393
40	17.09.201 9	Tuesda y	Physics (Student as Teacher Program)				
41	18.09.201 9	Wednsd ay	Zoology (Student as Teacher Program)				
42	19.09.201 9	Thursda y	Public Administration (Student as Teacher Program)				
43	20.09.201 9	Friday	Computer Science	B.sc (II Year/III Sem)	Trees & Graphs in Data structures	Dr.B.Sambasiva Rao,BJR GDC Narayanguda	9848450439
44	21.09.201 9	Saturda y	Special Live Program	All	Dial your CCE		
45	23.09.201 9	Monday	English	II Year	Media Reports	Dr.Ram Bhasker Raju,KGC Hanamkonda	9849169320
46	24.09.201 9	Tuesda Y	Chemistry (Student as Teacher Program)				
47	25.09.201 9	Wednsd ay	Botany (Student as Teacher Program)				
48	26.09.201 9	Thursda y	History (Student as Teacher Program)				
49	27.09.201 9	Friday	Commerce	III Year/V Sem	Banking Theory & Practise (Over view of	S.Ramesh,GDC(A) Khammam	9490993210

					RBI)		
50	10.10.201 9	Thursda y	Political Science (Pair teaching)	II Year/III Sem	Prime Minister	1.Dr.Jaheeda Bagum, GDC Hayatnagar 2.S.Yadagiri , NGC Nalgonda	9908078786 9492875705
51	11.10.201 9	Friday	Computer Science	B.Sc (III Year/ V Sem)	Process Synchronisation, Deadlocks	Dr.B.Sambasiva Rao,BJR GDC Narayanguda	9848450439
52	14.10.201 9	Monday	English(Student as Teacher Program)				
53	15.10.201 9	Tuesday	Mathematics	III Year /V Sem	Linear transformations & their Properties	M.Rajesh, GDC Agraharam	9849573963
54	16.10.201 9	Wednsd ay	Biotechnology	II Year/III Sem	Enzyme Inhibition Types	D.Annapurna, TARA GDC (A), Sangareddy	9959220195
55	17.10.201 9	Thursda y	Economics	II Year/III Sem	Monoploy	R.Venkata Ramarao,GDC(W) Nalgonda	9603374014
56	18.10.201 9	Friday	Zoology	III Year/V Sem	Enzymes	Dr.P.Ayodhya Reddy, GDC(M) Gajwel	9490550890
57	19.10.201 9	Saturda y	Commerce	III Year/ V Sem	Information Technology Act-2000	PGK.Gajendra Babu,GDC Malkajgiri	9440498388
58	21.10.201 9	Monday	Telugu (Student as Teacher Program)				
59	22.10.201 9	Tuesday	Physics	II Year/III Sem	Production of Low Temperatures	K.Haritha ,GDC Aler	9885286905
60	23.10.201 9	Wednsd ay	Zoology	II Year/III Sem	Distinguished characters of Poisionous & Non Poisonous snakes	J.Swamy, GDC (W) Nalgonda	9848480243
61	24.10.201 9	Thursda y	Public Administration	I Year/II Sem	New Public Administration- Minnowbrook-I	Dr.G.Narasimulu, GDC (M) Gajwel	9441301961
62	25.10.201	Friday	ТЅКС	All			

	9						
					Stress management		
	26.10.201	Saturda			and life skills		
63	9	У	Special Live Program	All	education		
	29.10.201						
64	9	Tuesday	Chemistry	I Year /I Sem	Gaseous State	Dr.B.Ramesh,KGC Hanamkonda	9866962589
					Anamolous Secondary		
					Growth Achyranthus,		
	30.10.201	Wednsd			Draceana, Boerhavia		
65	9	ау	Botany	II Year/IV Sem	stem	Dr.A.Srinivas, SRR GDC,Karimnagar	9490683621
	31.10.201	Thursda	History (Student as				
66	9	У	Teacher Program)				
	01.11.201						
67	9	Friday	Health & Nutrition	All			
	02.11.201	Saturda			Coordination and		
68	9	У	Commerce	I Year/I Sem	Control	S.Narender Reddy, GDC Godavarikani	9440383277
						1.Dr.T.S.Praveen Kumar,GDC Eturunagaram	9396806948
						2.Dr.Ram Bhasker Raju, KGC Hanamkonda	9849169320
	04.11.201		Basic Communication		Right to English (Panel	3.Dr.E.Srinivasa Rao, ABV GDC Jangaon	9848788425
69	9	Monday	(Panel Discussion)	All	Discussion)	4.DSSR Krishna,GDC Hayatnagar	9493212313
	05.11.201				Volumes and Surfaces		
70	9	Tuesday	Mathematics	I Year/I Sem	of revolution	Dr.D.Pushpa, ABV GDC, Jangaon	9440553078
	06.11.201	Wednsd					
71	9	ау	Microbiology	II Year/IV Sem	Transription	Dr.N.Hari Krishna, GDC(M) Gajwel	9966222110
	07.11.201	Thursda	Political Science (Pair		Sovereign	1.Dr.K.Bhasker, GDC Patancheru	9949999690
72	9	У	teaching)	I Year/ I Sem	state:Challenges	2.E.Swarnalatha O/o CCE TS Hyderabad	7660020750
	08.11.201						
73	9	Friday	Gender Sensitization	All			

74	11.11.201 9	Monday	Modern Language- Telugu (Panel discussion)	II Year / III Sem	Sahitya charitra	1.Dr.T.Krishnaiah,NGC(A) Nalgonda 2.Dr.Mrudula , GDC Narsapur 3.Dr.Rakesh Bhavani, GDC Hayatnagar 4.Dr.A.Param Jyothy , GDC Marripeda	9704731346 9441408393 9246607551 7396156568
75	13.11.201 9	Wednsd ay	Biotechnology	IIIYear/ V Sem	Post Transcriptional Modifications	Dr.G.Kishore Kumar,SRR GDC ,Karimnagar	9703663902
76	14.11.201 9	Thursda Y	Economics(Panel discussion)	III Year/V Sem	Issues of Indian Agriculture Sector	1.Dr.A.Venkatesham TARA GDC(A), Sangareddy 2.Dr.Nagoji GDC ,Ibrahimpatnam, 3.T.Bhasker Reddy, GDC(W) Nalgonda 4.B.Kasturi Bai, Pingle GDC (W) Warangal	9989942298 9440571781 9182564607 6302596798
77	15.11.201 9	Friday	Social Responsibility	All			
78	16.11.201 9	Saturda Y	Commerce	II Year/III Sem	Sampling Methods	Dr.D.T.Chary,GDC Parkal	9849610028
79	18.11.201 9	Monday	Basic Communication	All	Writing Skills	Dr.T.S.Praveen Kumar,GDC Eturunagaram	9396806948
80	19.11.201 9	Tuesday	Physics	I Year/I Sem	Central Forces	Rekha Venkateswarlu,GDC(W) Nalgonda	9440493244
81	20.11.201 9	Wednsd ay	Zoology	III Year/V Sem	Urine Formation	R.Naresh,GDC (W) Nalgonda	8919920204
82	21.11.201 9	Thursda Y	Political Science (Panel discussion)	II Year/ III Sem	Globalisation	1.P.Venakata Ramana, GDC Kukatpally 2.Dr.Jaheeda Begum, GDC Hayatnagar, 3.Dr.K.Bhasker GDC ,Patancheru 4.Dr.B.Kavitha, GDC,Wardannapet	9177087189 9908078786 9949999690 9948857326
83	22.11.201 9	Friday	Computer Science	B.Com (III Year/V Sem)	HTML Tags	Y.V.Ramarao,GDC Hayatnagar	9912890463
84	23.11.201 9	Saturda y	Commerce	III Year/V sem	Scope & Importance of Computerised Accounting	K.Linga Reddy, KGC Hanamkonda	9849404642

85	25.11.201 9	Monday	English	All	Soft Skills (Pair Teaching)	1.Dr.Nancy Serena,GDC Kukatpally 2.Dr.T.S Praveen Kumar,GDC Eturunagaram	8332896068 9396806948
86	26.11.201 9	Tuesday	Chemistry (Pair Teaching)	III Yr /V Sem	Molecular Spectroscopy	Dr.P.Adivi Raju, GDC(M) Gajwel	9440867068
87	27.11.201 9	Wednsd ay	Botany(Pair Teaching)	II Year/VI Sem	C3,C4 Cycles (Plant Physiology)	1.Dr.K.Sarojini Chakravarthy,IPGDC, Nampally, 2.D.Srihari Reddy, GDC Patancheru	9849370624 9441564471
88	28.11.201 9	Thursda y	History	II Year/III Sem	Administrative reforms of Shershah	M.Shanthaveni,IPGDC(W),Nampally	9440980955
89	29.11.201 9	Friday	Bio Chemistry (Student as Teacher Program)				
90	30.11.201 9	Saturda y	Special Live Program	All	Methods/ Strategies /Best Practices adopted by the colleges		
91	02.12.201 9	Monday	Telugu(Panel Discussion)	I Year/ II sem	Snehalatha Lekha	1.Dr.J.Neeraja, O/o CCE TS Hyderabad 2.Dr.Koyi Koteswara Rao, GDC (W) Nalgonda 3.Dr.M.Sampathy Kumar reddy,GDC (W) Karimnagar	7660020730 9440480274 9849470792
92	03.12.201 9	Tuesday	Mathematics	III Year /V Sem	Diagonalization	M.Rajesh, GDC Agraharam	9849573963
93	04.12.201 9	Wednsd ay	Microbiology	II Year/IV Sem	Transription	Dr.A.Madhavi, GDC (W) Begampet	9581208104
94	05.12.201 9	Thursda Y	<i>Political Science (Student as Teacher Program)</i>				
95	06.12.201 9	Friday	тѕкс	All			
96	07.12.201 9	Saturda y	<i>Commerce (Student as Teacher Program)</i>				

97	09.12.201 9	Monday	Basic Communication (Panel Discussion)	All	Interview Skills (Panel Discussion)	1.Dr.T.S.Praveen Kumar,GDC Eturunagaram 2.Dr.Ram Bhasker Raju, KGC, Hanamkonda 3.Dr.E.Srinivasa Rao, ABV GDC Jangaon 4.DSSR Krishna,GDC Hayatnagar	9396806948 9849169320 9848788425 9493212313
98	10.12.201 9	Tuesday	Physics	I Year/II Sem	Fundamentals of Vibrations	Rekha Venkateswarlu,GDC(W) Nalgonda	9440493244
99	11.12.201 9	Wednsd ay	Biotechnology (Student as Teacher Program)				
100	12.12.201 9	Thursda y	Economics (Student as Teacher Program)				
101	13.12.201 9	Friday	Health & Nutrition	All			
102	16.12.201 9	Monday	Modern Language- Telugu (Panel Discussion)	III Year/ V Sem	Vimarsha	1.Dr.Seetha Rama Rao , GDC (A) Khammam 2.Dr.Belli Yadaiah GDC Ramannapet 3.Dr.T.Suresh ,GDC Godavarikhnai 4.Dr. A.Paramjyothy, GDC Marripeda	9866563519 , 9848392690 , 8790569436 , 7396156568
103	17.12.201 9	Tuesda y	Chemistry (Student as Teacher Program)				
104	18.12.201 9	Wednsd ay	Zoology (Student as Teacher Program)				
105	19.12.201 9	Thursda y	Public Administration (Student as Teacher Program)				
106	20.12.201 9	Friday	Gender Sensitization	All			
107	21.12.201 9	Saturda y	Commerce	I Year/I Sem	Depreciation Accounts	Dr.A.Bala Chandram, GDC (A) Siddipet	9550559454
108	23.12.201 9	Monday	Basic Communication	All	Review (Book & Film)	DSSR Krishna,GDC Hayatnagar	9493212313

109	24.12.201 9	Tuesday	Mathematics	II Year/III Sem	Radius of Convergence	D.Venkatesh, KGC,Hanamkonda	9948185643
110	27.12.201 9	Friday	Social Responsibility	All			
111	28.12.201 9	Saturda y	Special Live Program	All	Preparation of Competitive exams - Tips & Strategies		
112	30.12.201 9	Monday	English(Student as Teacher Program)				
113	31.12.201 9	Tuesda y	Physics (Student as Teacher Program)				
114	02.01.202	Thursda y	History	III Year/V Sem	First World War- Causes-Consequences	S.Ganapathi Rao,KGC ,Hanamkonda	9030403940
115	03.01.202	Friday	Computer Science	B.sc& B.Com (I Year/ II Sem)	Constructors-C++	Dr.K.Suresh babu,KGC, Hanamkonda	9849664111
116	04.01.202 0	Saturda y	Commerce (Student as Teacher Program)				
117	06.01.202 0	Monday	Telugu (Student as Teacher Program)				
118	07.01.202 0	Tuesday	Chemistry(Pair Teaching)	I Year /II Sem	Inter Halogens,Polyhalides, Pseudo Halogens	1.Dr.V.Srinivas, GDC Mulugu, 2.Dr.M.Aruna, Pingle GDC(W) Warangal	9885059533 , 9985300641
119	08.01.202 0	Wednsd ay	Botany (Student as Teacher Program)				
120	09.01.202 0	Thursda y	Political Science(Panel Discussion)	II Year/IV Sem	Electoral Reforms	1.Dr.K.Hussain GDC (M) Gajwel 2.E.Swarnalatha O/o CCE TS Hyderabad 3.S.Yadagiri NGC Nalgonda 4.Dr.B.Kavitha ,GDC Wardannapet	9885077541 7660020750 9492875705 9948857326
121	10.01.202 0	Friday	тѕкс	All			

122	17.01.202 0	Friday	<i>Computer Science (Student as Teacher Program)</i>				
123	18.01.202 0	Saturda y	Commerce	III Year/VI Sem	Companies Act-2013 An Overview	Dr.Gopala Sudarshanam, GDC (M) Gajwel	9989450086
124	20.01.202 0	Monday	English	All	Indianisms	B.Raghavendra, GDC Narsapur	7901290580
125	21.01.202 0	Tuesda y	<i>Mathematics (Student as Teacher Program)</i>				
126	22.01.202 0	Wednsd ay	Zoology(Pair Teaching)	II Year/IV Sem	Forces of Evolution	1.G.N Radhika,GDC Patancheru, 2.L.Mahesh TARA GDC (A) Sangareddy	9490347231 9440830275
127	23.01.202 0	Thursda y	Political Science	III Year/VI Sem	Basava	Dr.K.Hussain , GDC (M) Gajwel	9885077541
128	24.01.202 0	Friday	Commerce (Panel discussion)	III Year/VI Sem	Overview of GST	1.Gajendra Babu PGK,GDC Malkajgiri 2. S.Ramesh, GDC(A),Khammam 3.M.Somaiah, KGC,Hanamkonda	9440498388 9490993210 9948048949
129	25.01.202 0	Saturda y	Special Live Program	All	Focus on examination preparation and study opportunities in Higher Education.		
130	27.01.202 0	Monday	Telugu(Panel Discussion)	II Year / IV Sem	Mana Grama Naamaalu	1.Dr.M.Sampath Kumar reddy,GDC(W),Karimnagar 2.Dr.T.Suresh ,GDC Godavarikhnai 3.Dr. Rakesh Bhavani, GDC Hayatnagar	9849470792 8790569436 9246607551
131	28.01.202 0	Tuesday	Physics	I Year/II Sem	Damped Oscillations	K.Haritha ,GDC Aler	9885286905
132	29.01.202 0	Wednsd ay	Botany(Pair Teaching)	I Year/II Sem II Year/IV sem	Asclepiadaceae,Lamiac eae(Taxonomy)	1.Dr.K.Usha Rani, O/o CCE TS Hydearabad, 2.Dr. Annie Sheran KGC Hanamkonda	9849172438 9492434304
133	30.01.202 0	Thursda y	Economics (Panel Discussion)	II Year/IV Sem	Tax system in India	1.Dr.Nagoji GDC ,Ibrahimpatnam, 2.E.Pavani , Govt.City College, Nayapul 3.T.Bhasker Reddy, GDC(W) Nalgonda 4.B.Kasturi Bai, Pingle GDC (W) Warangal	9440571781 9959632449 9182564607 6302596798

			Computer Sicence		
	31.01.202		(Student as Teacher		
134	0	Friday	Program)		

For Commissioner of Collegiate Education





Department of Chemistry,

Tara Govt. College(A), Sangareddy.

RECORD OF EXTENSION LECTURES ARRANGED

<u>2018-19</u>

S.No.	Name of Resource	Affiliation	Date	Topic	No. of
	person				students
1	Dr. Sabitha.Y	Managing	15.02.19	Molcular	25
		Director &		Docking	
		Chief Scientist			
		from Ciencia			
		Life Sciences,			
		Hyderabad			





Extension lecture on Molcular Docking by Dr. Sabitha.Y, Managing Director & Chief Scientist from Ciencia Life Sciences, Hyderabad



Attendance Certificate

11 October 2019

Mr A Vishweshwara Sharma, Assistant Professor of English at TARA Government college, Sangareddy (A), has attended the Orientation Programme for newly recruited lecturers of Tribal welfare Residential Degree Colleges on 11 October as a resource person for 3 batches and gave extension lectures on ' Effective Classroom management : creating a platform for Debate League'.

His absence at his college may be treated as On Duty (OD) for one day on 11-10-2019.

File No.CCE-SER2/CLEC/9/2019-SER2

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION TELANGANA STATE :: HYDERABAD

PRESENT: NAVIN MITTAL, I.A.S.,

- Sub:- Contract faculty- Contract Lecturers working in Government Degree Colleges in the State – Identified the surplus Contract Lecturers due to relocation of regular faculty - Relocation of Contract Lecturers Working in Government Degree Colleges - On duty facility to Certain faculty - Orders - Issued.
- Read:- 1. Govt.Memo.No.2859/CE/A1/2015, dated 08.06.2019
 2. File NoCCE-AC/GEN/98-Academic Cell (Computer No.157641) dated 19.12.2019.
 3. Counselling conducted on 13.12.2019 and Options given by the individuals.

The Commissioner of Collegiate Education, Telangana State, Hyderabad have identified the surplus Contract Lecturers due to relocation of regular faculty. In reference 2nd read above a counselling was held on 13.12.2019 at IPGDC(W), Nampally.

In the circumstances reported in references 1 to 3rd read above, the Commissioner of Collegiate Education, Telangana State, Hyderabad is hereby direct to work on "On Duty" till last day of summer vacation 2019-20 Academic Year and join their original place of work on the re-opening day of 2020-21 as shown hereunder:-

	Name of the Contract Lecturer	Subject	Presently working	"On Duty to Work" at
11 9	M.Santosh	Physics	GDC (A) Sangareddy	GDC (A) Mahabubnagar

File No.CCE-SER2/CLEC/9/2019-SER2

Further, the Principals of Government Degree College concerned are requested to report compliance in the matter promptly.

The receipt of these proceedings shall be acknowledged.

(Orders issued with the approval of the Commissioner of Collegiate Education, Telangana State, Hyderabad)

Signature Not Verified Digitally signed by N VIJAYA BHASKAR RAO Date: 2019.12.31 16:05:41 IS Reason: Approved

For COMMISSIONER OF COLLEGIATE EDUCATION

То

The Principal, Government Degree Colleges as mentioned above. Copy to the Individual (through Principal of the College) Copy to the DTO / STO concerned Copy to the GDC-CS concerned

File No.CCE-SER2/TRF/4/2019-SER2

PROCEEDINGS OF THE COMMISSIONERATE OF COLLEGAITE EDUCATION TELANGANA::HYDERABAD

Present: Navin Mittal IAS

- Sub:- Commissionerate of Collegiate Education-Govt. Degree Colleges- Reorganization of courses-Admissions in Govt. Degree Colleges-Workload Calculation-Reorganization of posts and Shifting of Surplus Employees along with post- Orders - Issued.
- Read:1. G.O.Rt.No.65 Higher Education (CE) Dept., dated 13.04.2018-2. FileNo.CCE-AC/GEN/82/2019-AcademicCell19.10.20193. FileNo.CCE-AC/GEN/83/2019-AcademicCelldated

31.10.2019

The Commissioner of Collegiate Education has identified an imbalance in the teaching faculty in the colleges based on the workload calculation. In view of this, surplus regular faculty identified in some Government Degree Colleges are hereby shifted along with the posts to the required colleges based on the workload calculation for optimum utilization of regular teaching staff under administrative grounds.

In the circumstances reported in references 1 to 3rd read above, the Commissioner of Collegiate Education, Telangana State, Hyderabad is hereby shift the Lecturer along with the post as shown hereunder:-

SI.	Name of the	Subject	Presently	Shifting of the
No.	Lecturer		working college	post along with
	Sri/Smt.			the Person to
				the College
18	Rajendra Kumar	Hindi	Government	Government
			Degree College,	Degree College
			Shadnagar	(A), Sangareddy

The Principals of Government Degree Colleges are requested to relieve and admit to duty immediately and report compliance.

The Principals of Government Degree Colleges concerned are requested to make necessary entries in the respective staff register and scale register and report compliance immediately.

The receipt of these Proceedings shall be acknowledged.

(Orders issued with the approval of the Commissioner of Collegiate Education, Telangana State, Hyderabad)

Signature Not Verified Digitally signed by Dr N Vijaya Braskar Rao Date: 2019.11.01 18:17:15 ISP Reason: Approved

For Commissioner of Collegiate Education

То

The Individual Concerned

(through Principal of Government Degree College concerned)

Copy to the Principal, Government Degree College concerned.

File No.CCE-SER2/TRF/4/2019-SER2

PROCEEDINGS OF THE COMMISSIONERATE OF COLLEGAITE EDUCATION TELANGANA::HYDERABAD

Present: Navin Mittal IAS

Sub:- Commissionerate of Collegiate Education-Govt. Degree Colleges- Reorganization of courses-Admissions in Govt. Degree Colleges-Workload Calculation-Reorganization of posts and Shifting of Surplus Employees along with post- Orders - Issued.

Read: File No.CCE-AC/GEN/81/2019-Academic Cell dated 12.10.2019

The Commissioner of Collegiate Education has identified an imbalance in the teaching faculty in the colleges based on the workload calculation. In view of this, surplus regular faculty identified in some Government Degree Colleges are hereby shifted along with the posts to the required colleges based on the workload calculation for optimum utilization of regular teaching staff under administrative grounds.

In the circumstances above, the Commissioner of Collegiate Education, Telangana State, Hyderabad is hereby shift the Lecturer along with the post as shown hereunder:-

SI. No.	Name of the Lecturer	Subject	Presently working college	Shifting of the post along with the Person to the College
20	Ramulu	Public Administratio n	Government Degree College, Sadasivpet	Government Degree College (Co-Ed), Sangareddy

The Principals of Government Degree Colleges are requested to relieve and admit to duty immediately and report compliance.

File No.CCE-SER2/TRF/4/2019-SER2

The Principals of Government Degree Colleges concerned are requested to make necessary entries in the respective staff register and scale register and report compliance immediately.

The receipt of these Proceedings shall be acknowledged.

(Orders issued with the approval of the Commissioner of Collegiate Education, Telangana State, Hyderabad)

Signature Not Verified Digitally signed by Dr C Manjulatha Date: 2019.10.12 18:23:15 IST Reason: Approved

For Commissioner of Collegiate Education

То

The Individual Concerned

(through Principal of Government Degree College concerned)

Copy to the Principal, Government Degree College concerned.

File No.CCE-AC/ODWS/3/2019-ACADEMIC CELL

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI.NAVIN MITTAL, I.A.S.

- Sub: Centre of Excellence for Studies in Classical Telugu, Central Institute of Indian Languages(MHRD), Manasagangothri, Mysore-"A five Day Workshop on "Prachina Telugu Sahitya Adhyayanam" from 16th to 20th September 2019- Permission accorded to the List of Participants from Telugu Faculty from various GDCs to attend the Work Shop-Orders Regarding.
- **Ref:** Letter received from Project Director of Workshop, Centre of Excellence for Studies in Classical Telugu, Central Institute of Indian Languages, Mysore, dated: 06.09.2019

Vide reference read, the Commissioner of Collegiate Education has accorded permission to the list of following Telugu Faculty members from various GDCs of Telangana State to participate in the **Workshop on "Prachina Telugu Sahitya Adhyayanam"** from **16th** to **20th September 2019** which will be held at Central Institute of Indian Languages (MHRD), Manasagangothri, Mysore.

SI.		
No	Name of the College	Name of the Faculty
1	Tara GDC, Sangareddy/GDCW, Nalgonda	Dr.Rapolu Srinivas
2	SRRGDC, Karimnagar	K.Malla Reddy
3	GDC, Siddipet	Nandigama Nirmala Kumari
4	GDCw, Kalwakurthy/GDCW, Jagtial	Goli Sreelatha
5	NG GDC, Nalgonda	Dr.N.Deepika

The absence of the above said faculty members shall be treated as **ON DUTY** for the said date and they are not eligible to claim TA/DA from the Colleges. In view of the above, the Principals of concerned GDCs are informed to relieve them to attend the above said Workshop.

Signature Not Verified Digitally signed by Dr C Manjuatha Date: 2019.09.09 01:20:00

For Commissioner of Collegiate Education

То

The Principals of GDC concerned.

File No.CCE-SER2/TRF/4/2019-SER2

PROCEEDINGS OF THE COMMISSIONERATE OF COLLEGAITE EDUCATION TELANGANA::HYDERABAD

Present: Navin Mittal IAS

Sub:- Commissionerate of Collegiate Education-Govt. Degree Colleges- Reorganization of courses-Admissions in Govt. Degree Colleges-Workload Calculation-Reorganization of posts and Shifting of Surplus Employees along with post- Orders - Issued.

Read: File No.CCE-AC/GEN/81/2019-Academic Cell dated 12.10.2019

The Commissioner of Collegiate Education has identified an imbalance in the teaching faculty in the colleges based on the workload calculation. In view of this, surplus regular faculty identified in some Government Degree Colleges are hereby shifted along with the posts to the required colleges based on the workload calculation for optimum utilization of regular teaching staff under administrative grounds.

In the circumstances above, the Commissioner of Collegiate Education, Telangana State, Hyderabad is hereby shift the Lecturer along with the post as shown hereunder:-

SI. No.	Name of the Lecturer	Subject	Presently working college	Shifting of the post along with the Person to the College
18	Jyothi Sardar	Hindi	Government Degree College (W), Sangareddy	

The Principals of Government Degree Colleges are requested to relieve and admit to duty immediately and report compliance.

File No.CCE-SER2/TRF/4/2019-SER2

The Principals of Government Degree Colleges concerned are requested to make necessary entries in the respective staff register and scale register and report compliance immediately.

The receipt of these Proceedings shall be acknowledged.

(Orders issued with the approval of the Commissioner of Collegiate Education, Telangana State, Hyderabad)

Signature Not Verified Digitally signed by Dr C Manjulatha Date: 2019.10.12 18:25:57 IST Reason: Approved

For Commissioner of Collegiate Education

То

The Individual Concerned

(through Principal of Government Degree College concerned)

Copy to the Principal, Government Degree College concerned.

File No.CCE-AC/ODWS/4/2019-ACADEMIC CELL

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI.NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education-Tara GDC, Sangareddy-A.Vishweshwara Sharma, Assistant Professor of English-Served as a **Resource person** for a day on **11.10.2019** in three day Orientation Programme organized by TTWREIS for newly recruited TS Residential Degree College Lecturers-Sanction of **On Duty**-Orders-Regarding.
- **Ref:** Letter received from Principal, Tara GDC, Sangareddy, dated: 13.10.2019.

Vide references read, the Commissioner of Collegiate Education has accorded On Duty to **A.Vishweshwara Sharma**, Assistant Professor of English, Tara GDC, Sangareddy served as a Resource person on **11.10.2019** in a three day Orientation Programme organized by Telangana Tribal Welfare Residential Educational Institutions Society (TTWREIS) for newly recruited TW residential Degree College Lecturers at Rajendranagar, Hyderabad.

The absence of the above said Assistant Professor shall be treated as **ON DUTY** on above said date. In view of the above, the Principal, Tara GDC, Sangareddy is informed to consider his absence in the college as **On DUTY**.

(Orders of the CCE have been obtained in the note file)



For Commissioner of Collegiate Education

To The Principal of GDC concerned.

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION TELANGANA::HYDERABAD

PRESENT: Sri Navin Mittal, I.A.S.,

- Sub: TSCES Dr. Ch. Jyothsna, Assistant Professor of Political Science, Government Degree College, Sadasivpet, Sangareddy Dist. - Permission to work at Government Degree College(A), Sangareddy on "On Duty" basis– Orders - Issued.
- Read: As per the Note orders of the CCE in Note File bearing No. File No. CCE-AC/GEN/6/2019-ACADEMIC CELL (Computer No. 72006).

In the circumstances stated in the reference read above, the Commissioner of Collegiate Education, Telangana, Hyderabad is hereby accorded permission to Dr. Ch. Jyothsna, Assistant Professor of Political Science, Government Degree College, Sadasivpet, Sangareddy Dist. to work at Government Degree College (A), Sangareddy on "on duty" for the Academic year 2019-2020.

The Principal, Government Degree College, Sadasivpet, Sangareddy Dist. is requested to relieve the incumbent of the college immediately so as to enable her to work at Government Degree College(A), Sangareddy on "On Duty "basis. She will be claiming her salary at her parent institution on production of attendance certificate at the end of every month from the Principal, Government Degree College (A), Sangareddy.

The Principal, Government Degree College (A), Sangareddy is requested to admit the incumbent to duty and then submit the date of relief and joining to the Commissioner of Collegiate Education promptly.

The receipt of these proceedings should be acknowledged by return of post.

(Orders of the Commissioner of Collegiate Education have been obtained in the note file)

Signature Not Verified Digitally signed by Dr C Manjulatha Date: 2019.07.15 16:26:14 IST Reason: Approved

For COMMISSIONER OF COLLEGIATE EDUCATION

То

- Dr. Ch. Jyothsna, Assistant Professor of Political Science, Government Degree College, Sadasivpet, Sangareddy Dsit. (through the Principal of the College).
- Copy to the Principal, Government Degree College, Sadasivpet, Sangareddy Dist.

Copy to the Principal, Government Degree College (A), Sangareddy.

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION TELANGANA::HYDERABAD PRESENT: Sri Navin Mittal, I.A.S.,

- Sub: TSCES –Smt. M. Praveena, Assistant Professor of English, Government Degree College(A),Sangareddy - Permission to work at Government Degree College, Patancheru, Sangareddy Dist. on "On Duty" basis– Orders - Issued.
- Read: As per the Note orders of the CCE in Note File bearing No. File No. CCE-AC/GEN/6/2019-ACADEMIC CELL (Computer No. 72006).

In the circumstances stated in the reference read above, the Commissioner of Collegiate Education, Telangana, Hyderabad is hereby accorded permission to Smt. M.Praveena, Assistant Professor of English, Government Degree College(A),Sangareddy to work at Government Degree College, Patancheru, Sangareddy Dist. on "on duty" for the Academic year 2019-2020.

The Principal, Government Degree College(A), Sangareddy is requested to relieve the incumbent of the college immediately so as to enable her to work at Government Degree College, Patancheru, Sangareddy Dist. on "On Duty "basis. She will be claiming her salary at her parent institution on production of attendance certificate at the end of every month from the Principal, Government Degree College, Patancheru, Sangareddy Dist.

The Principal, Government Degree College, Patancheru, Sangareddy Dist. is requested to admit the incumbent to duty and then submit the date of relief and joining to the Commissioner of Collegiate Education promptly.

The receipt of these proceedings should be acknowledged by return of post.

(Orders of the Commissioner of Collegiate Education have been obtained in the note file)

Signature Not Verified Digitally signed by Dr C Manjulatha Date: 2019.07.15 8:28:48 IST Reason: Approved For COMMISSIONER OF COLLEGIATE EDUCATION

То

Smt. M. Praveena, Assistant Professor of English, Government Degree College(A), Sangareddy (through the Principal of the College).

Copy to the Principal, Government Degree College (A),Sangareddy. Copy to the Principal, Government Degree College, Patancheru, Sangareddy Dist.

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION TELANGANA::HYDERABAD

PRESENT: Sri Navin Mittal, I.A.S.,

- Sub: TSCES Sri Rapolu Srinivas, Assistant Professor of Telugu, Government Degree College(A),Sangareddy – Permission to work at Government Degree College for Women, Nalgonda on "On Duty" basis– Orders - Issued.
- Read: As per the Note orders of the CCE in Note File bearing No. File No.CCE-AC/GEN/6/2019-ACADEMIC CELL(Computer No. 72006).

In the circumstances stated in the reference read above, the Commissioner of Collegiate Education, Telangana, Hyderabad is hereby accorded permission to Sri Rapolu Srinivas, Assistant Professor of Telugu, Government Degree College (A), Sangareddy to work at Government Degree College for Women, Nalgonda on "on duty" for the Academic year 2019-2020.

The Principal, Government Degree College(A), Sangareddy is requested to relieve the incumbent of the college immediately so as to enable him to work at Government Degree College for Women, Nalgonda on "On Duty "basis. He will be claiming his salary at his parent institution on production of attendance certificate at the end of every month from the Principal, Government Degree College for Women, Nalgonda.

The Principal, Government Degree College for Women, Nalgonda is requested to admit the incumbent to duty and then submit the date of relief and joining to the Commissioner of Collegiate Education promptly.

1

The receipt of these proceedings should be acknowledged by return of post.

(Orders of the Commissioner of Collegiate Education have been obtained in the note file)

Signature Not Verified Digitally signed by Dr C Manjulatha Date: 2019.07.15 17:52:26 IST For COMMISSIONER OF COLLEGIATE EDUCATION

То

Sri Rapolu Srinivas, Assistant Professor of Telugu, Government Degree College(A), Sangareddy (through the Principal of the College).

Copy to the Principal, Government Degree College (A), Sangareddy. Copy to the Principal, Government Degree College for Women, Nalgonda.

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION TELANGANA::HYDERABAD

PRESENT: Sri Navin Mittal, I.A.S.,

- Sub: TSCES Sri D. Srinivas, Assistant Professor ofHistory, Government Degree College, Sathupally, KhammamDist-Permission to work at Government Degree College, Sangareddy on "On Duty" basis– Orders - Issued.
- Read: As per the orders and approved listof the Commissioner of Collegiate Education, Nampally, Hyderabad in note file No. CCE-AC/GEN/6/2019, Dated 20.06.2019.

In the circumstances stated in the reference read above, the Commissioner of Collegiate Education, Telangana, Hyderabad is hereby accorded permission to Sri D. Srinivas, Assistant Professor of History, Government Degree College, Sathupally, Khammam Dist. to work at Government Degree College, Sangareddy on "on duty" basis for the Academic year 2019-20.

The Principal, Government Degree College, Sathupally, Khammam Dist. is requested to relieve Sri D. Srinivas, Assistant Professor of History of the college immediately so as to enable him to work at Government Degree College, Sangareddy on "On Duty" basis. He will be claiming his salary at his parent institution on submission of attendance certificate at the end of every month from the Principal, Government Degree College, Sangareddy Dist.

The Principal, Government Degree College, Sangareddy is requested to admit Sri D. Srinivas, Assistant Professor of History to duty and then submit the date of relief and joining to the Commissioner of Collegiate Education promptly.

The receipt of these proceedings should be acknowledged by return of post.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by BHANAVATH SWAMY Date: 2019.06.26 16:13:29 IST Reason: Approved

То

Sri D. Srinivas, Assistant Professor of History, Government Degree College, Sathupally, Khammam Dist. (through the Principal of the College).

Copy to the Principal, Government Degree College, Sathupally, Khammam Dist.

Copy to the Principal, Government Degree College, Sangareddy Dist. Copy to Services-I,II,IV Sections of this office.

TARA GOVERNMENT COLLEGE, SANGAREDDY - 502 001 (AUTONOMOUS)

BONAFIDE CERTIFICATE

Certified that the project report "Antimicrobial Silver Nanoparticle coating on Currency notes and Mobile phones using Eco-friendly Tollens process for prevention of infectious diseases"

S.No.	Name of the Student	Roll Number	Group	Year
1	A.Bhanupraksh	17016058445001	B.Sc.(BZC)	Ш
2	J.Manjulatha	17016058457025	B.Sc.(MZC)	III
3	A.Pranaya	17016058481001	B.Sc.(BCCA)	III
4	P. Pooja	17016058445028	B.Sc.(BZC)	III
5	J.Swetha	17016058457026	B.Sc.(MZC)	III
6	G.Sai Ganesh	17016058457023	B.Sc.(MZC)	Ш

15	the	bonaj	id	ewori	201	

who carried out the project work under my supervision in the academic year of 2019-20.

PROJECT SUPERVISOR

AD.

PRINCIPAL PRINCIPAL **FARA GOVT COLLEGE** (AUTONOMOUS) SANGAREDDY-502 001

DEPARTMENT OF CHEMISTRY Tara Govt. College, Sangareddy (A) Dist. Sangareddy - 502 001

Antimicrobial Silver Nanoparticle coating on Paper currency notes and Mobile phones using Eco-friendly Tollens process for prevention of infectious diseases

INTRODUCTION:

Contaminated Paper currency notes may cause a public health risk by spreading nosocomial (Hospital acquired infections) infections when simultaneous handling of food and in addition to this, also cause normal sort of contaminations in persons with immunodeficiency. Especially when banknotes recovered from hospitals may be highly contaminated by *Staphylococcus aureus*, *Salmonella* species and *Escherichia coli*. Laboratory studies revealed that methicillin-resistant *S. aureus* can easily survive on paper currency notes, whereas *E. coli*, *Salmonella* species and viruses, including human influenza virus, Norovirus, Rhinovirus, hepatitis A virus and Rotavirus, which can be transmitted through hand contact. Large-scale, 16S rRNA, metagenomic studies and culturomics have the capacity to dramatically expand the known diversity of bacteria and viruses on money and fomites [1].



Figure 1: Depiction microbes on paper currency notes (**Courtesy:** *Future Microbiol.* (2014) 9(2), 249–261).

Similarly, constant handling of the Mobile phone by different users exposes it to an array of microorganisms, and makes it a good carrier for microbes, especially those associated with the skin resulting in the spread of different microorganisms from user to user. Because of the multifaceted benefits of the mobile phones, it is easy to overlook its hazard to health; this is against the background that many users may have no regard for personal hygiene, and the number of people who may use the same phone. Many research studies has shown that the mobile phone could be a health hazard with tens of thousands of microbes living on each square inch of the phone [2,3].

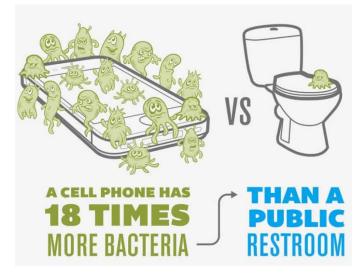


Figure 2: Depiction microbes on Mobile phones.

(Courtesy: https://www.phonesoap.eu/medical-professional)

However, Nosocomial infections caused by microorganisms which usually originated from hospital environments and cross-contamination due to the incorrect use of medical equipment can be prevented or reduced by replacing normal equipments with Silver nanoparticles (AgNPs) coated medical devices [4,5]. Many medical studies have revealed that silver is effective against more than 650 pathogens, having a broad spectrum of activity. Further its use in the form of Nanoparticles enhances this property up to great extent and allows its use in a wide range of applications [6, 7]. Therefore, in recent years Nano-silver is considered as one of the most viable

alternatives to antibiotics because it seems to have high potential to solve the problem of multidrug resistance, which is often observed in several bacterial strains [8-10].

Nanoparticles are usually a clusters of atoms, with sizes ranging between 1 and 100 nm, whereas the word "Nano" is used to indicate one billionth of a meter [11-13]. Because of the variation in the size of AgNPs, they exhibited variety of physical and chemical characteristics to that of metallic silver [14, 15].

The silver is well known for its antimicrobial activity. In Nano-metric form, silver has shown accentuated antimicrobial characteristics. Due to their nano scale size, AgNPs can enter in to cells and inhibit enzymatic systems in the respiratory chain of some bacteria and thereby alter their DNA synthesis. AgNPs, their use can be recommended as a good alternative for the control of microorganisms, with less risk of toxicity to human cells [16].

Various studies have revealed the effectiveness of AgNPs as dressings for covering burns to surgical devices and bone prostheses, and are incorporated into clothing – always with the aim of producing antimicrobial effect [17-20].

In this context, we have been used silver nanoparticles (AgNP) to coat the surfaces of both Paper currency notes and mobile phones for the prevention of microbial contamination.

RESEARCH PROBLEM:

In recent time, healthcare-associated infections are one of the most serious patient safety issues in healthcare today [21]. Most microbes are able to survive on surfaces and Paper currency note's surfaces can act as sources of pathogen transmission if no disinfection is performed. In addition, the survival of nosocomial bacterial strains, including methicillin-resistant *Staphylococcus aureus* (MRSA), in the environment is of great interest to infection control professionals [22]. Moreover, workers who are working with food and edible products have been implicated in several outbreaks of food-borne diseases who were frequently handling contaminated paper currency notes and human occupational activities could introduce the risk of food contamination [23]. Pathogens that can infect food workers have multiple sources among them paper currency notes occupy considerable portion and contaminated workers in turn become potential sources of contamination in food processing and preparation facilities [24].

In addition to the Paper currency notes, mobile phones also act as potential non-vector causative factors for microbial contamination due to its constant contact with humans. According to many microbial studies conducted by Microbiologists reveal the combination of constant handling with the heat generated by the mobile phones creates a prime breeding platform for many microorganisms that are normally found on the skin. Staphylococci species, particularly S. epidermidis are which belongs to the normal flora of the human skin, respiratory and gastrointestinal tracts. 20-50% of human beings contain S. aureus in their nasal carriage. are also found regularly on Clothes, bed linen, and other human environments usually contains Staphylococci species [25]. Staphylococcus aureus, a common bacterium found on the skin and in the nasal fluids of up to 25% of healthy people and animals can cause illnesses from pimples and boils to pneumonia and meningitis, and is a close relative of methicillin Resistant Staphylococcus aureus (MRSA). Human hand acts as main reservoir of S. aureus from where it is introduced into food during preparation [26]. The hands also serve as a major vehicle of transmission of various pathogens including the enteric species [27]. Proteus mirabilis is one of the most common Gram-negative pathogens found in clinical specimens. It can cause a variety of community or hospital-acquired infections, including those of the urinary tract, respiratory tract, wounds and burns, bacteraemia, neonatal meningoencephalitis, empyema and osteomyelitis [28]. After Escherichia coli, P. mirabilis is the member of the Enterobacteriaceae most often isolated in European clinical microbiology laboratories [29] and accounting for ~3% of nosocomial infections in the United States [30]. Pseudomonas aeruginosa is a metabolically versatile γ -Proteobacterium, which inhabits terrestrial, aquatic, animal, human, and plant-host-associated environments [31].

To address the above issues the common utility objects like Paper currency notes and mobile phone surfaces should be disinfected regularly with broad spectrum antibiotics. These antibiotics are chemically unstable under the normal handling conditions and their efficacy will reduced significantly with time. Hence these antibiotics should be used regularly used for disinfection, which leads to evolution of multi-drug resisted microbial strains. This further ruins the public health care with diseases which will not be controlled by normal dosage of antibiotics.

To answer the above research problem innovative approach of coating Silver nanoparticles on surfaces of Paper currency notes and mobile phones has been adopted. The most applied method for AgNPs preparation is by the reduction of Ag^+ in aqueous solution. For this purpose we have used simple Tollens process with slight modifications.

Owing to their peculiar properties Nanoparticles attracts great interest for applicative methods in many disciplines [32], among them the most advance application is in the field of biology and medicine [33]. Many metals like silver, copper, gold, magnesium etc. have been exhibited potential antimicrobial property in the form of Nano-particles and among these silver was the most efficient [34]. The antimicrobial activity of Silver nanoparticles (AgNPs) against both pathogenic fungi and bacterial strains is attracting researcher's attention in multidisciplinary applications of health care. Many bacterial strains have great intrinsic antimicrobial resistance limiting the number of effective antibiotics. Thus, metallic antimicrobial agents such as silver nanoparticles (AgNPs) are considered as potential agents to help manage and prevent infections. AgNPs can be used in several applications against bacteria which are resistant to common antibiotics or even multiresistant bacteria such as *P. aeruginosa*.

OBJECTIVES:

- The major objective of the project is to develop a protective strategy for prevention of microbial infections caused by contaminated Paper currency notes and Mobile phones using concepts of nanotechnology.
- To Design the novel synthetic strategy for coating silver nanoparticles (AgNP) on Paper currency notes and mobile phone screen using simple tollens reagent and evaluate its antimicrobial efficacy against both bacteria and fungi by microbial screening methods.

REVIEW OF LITERATURE:

Nosocomial infections caused by contaminated Paper currency notes have a significant impact on public health in recent time. According to the studies conducted by *Emmanouil Angelakis et al.*[1] stated that that contaminated money and coins are a public health risk when associated with the simultaneous handling of food, and Paper currency may spread nosocomial infections. We have highlighted the potential for banknotes and coins to carry bacteria and fungi, as well as their potential capacity to

spread infectious agents. In addition, banknotes and monetary coinage can act as potential reservoirs for antibiotic-resistant bacteria, such as MRSA.

- Similarly mobile phones also act as non vector factors for cross-contamination. Saeed Banawas et al.[35] reported that the cell phones of healthcare workers can be contaminated by a wide range of bacteria including multidrug resistance bacteria. Bacteria may be readily able to adhere to the surface of mobile phones, and the heat emitted by the cell phone enhances bacterial growth and these bacteria can then be transferred to one person to another. Another study conducted by Raghavendra Rao Morubagal et al.[36] also revealed the presence of pathogenic bacteria on Mobile phones which are capable of causing infections when dealt with health care associates.
- **R. Salomoni et al.** [16] in their study explained the antibacterial activity of AgNPs especially multidrug resistant strains of *Pseudomonas aeruginosa* which are common pathogens in nosocomial infections.
- According to study conducted by K. M. Alananbeh et al.,[37] AgNP have been possess
 potential antifungal activity against various fungal strains like *Aspergillus* sp. i.e. *A. niger* and *A. terreus*. The gradual growth reduction was clear in both *Aspergillus* species
 with the increase in concentration of the AgNP.
- *Yadong Yin et al.* [38] reported a simple and convenient procedure based on the Tollens process for the preparation of silver nanoparticles with a relatively narrow distribution in size in the range of 20–50 nm. These silver particles could be easily prepared either as stable aqueous dispersions or as decorative coatings on microspheres and surfaces.
- *Gayatri Dhulappanavar et al.* [39] reported an eco-friendly synthesis of AgNP using Lemon fruit juice (*Citrus limon L.*) as a reducing and stabilizing agent.
- According to study conducted by **Padma S Vankar** and **Dhara Shukla** [40] showed that Antimicrobial finish on fabric provided durable textile finish on cotton and silk fabric. The Preparation of silver nanoparticles (AgNP) have been carried out biosynthetically using aqueous extract of Lemon leaves (*Citrus limon*) which acts as reducing agent and encapsulating cage for AgNP.
- Polymer nanocomposites containing metal nanoparticles have attracted a great interest due to their unique chemical and physical properties. "Green" chemistry promotes application of natural fibers in such structures, among them cellulose is one of the most

frequently used. However, cellulose fabrics have ability to absorb moisture, so under certain conditions of humidity and temperature they can be subjected to microbial attack. One of the most popular and best known antibacterial agents is silver, which serves as a potential antibacterial material acting against an exceptionally broad spectrum of bacteria including activity against antibiotic-resistant bacteria. **Dagmara K. Chmielewska et al.**[41] revealed in their studies that Silver nanoparticles (Ag NPs) were grown at the cellulose fibers surface by direct reduction of AgNO₃ with electron beam (EB) application.

• E. Smiechowicz et al. [42] reported the enhanced antibacterial activity of nanocomposite cellulose fibers of Lyocell type modified with nanosilver particles and nanosilica.

RESEARCH METHODOLOGY:

Nanotechnology deals with various structural aspects of matter having dimensions of the order of a billionth of a meter. Based on the size, Nano-materials are usually intermediate between macroscopic solid materials and of atomic and molecular systems. Specific physical, chemical and biological properties of Nano-materials make them dissimilar from the macroscopic bulk materials. These properties of Nano-particles provide us the scope of multiple applications in advance research to day to day life.

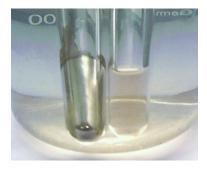
Based on the size, morphology, physical and chemical properties NPs (Nano-particles) were categorized into different types namely ceramic nanoparticles, carbon-based nanoparticles, metal nanoparticles etc. Among these Metal based Nps have shown multifaceted applications. Metal nanoparticles are prepared from metal precursors. These nanoparticles can be synthesized by chemical, electrochemical, or photochemical methods. In chemical methods, the metal nanoparticles are obtained by reducing the metal-ion precursors in solution by chemical reducing agents. These have the ability to adsorb small molecules and have high surface energy. In the present study we use nanotechnology to answer the research problem, i.e. to design antimicrobial protective layer on Paper currency notes and mobile phone surfaces for prevention of microbial contamination. For this we have chosen AgNP because of its significant antimicrobial activity,

AgNP have several merit over normal antibiotics which includes long period of effectiveness and will not initiate drug resistance among the microbial strains.

In the present study we have used very simple Tollens process which is used to identify aldehyde functional group in the organic chemistry.

CONCEPT:

Tollens' reagent is usually used to determine the presence of aldehyde functional group on aliphatic, aromatic and carbohydrate (reducing sugars) moieties, it will also gie positive test with some alpha-hydroxy ketones which can tautomerize into aldehydes. The reagent is prepared from aq.silver nitrate solution, ammonia and some sodium hydroxide (to maintain a basic pH of the reagent solution). It was named after its discoverer, the German chemist Bernhard Tollens [43]. A positive test with Tollens' reagent is indicated by the precipitation of elemental silver, often producing a characteristic "silver mirror" on the inner surface of the reaction vessel.



This reagent is freshly prepared in the laboratory for immediate use due to its short shelf life and easily decomposes hence it is not commercially available. Commonly the preparation involves two steps. First a few drops of dilute sodium hydroxide are added to some aqueous 0.1 M silver nitrate. The HO⁻ ions convert the silver aquo complex form into silver oxide, Ag₂O, which precipitate from the solution as a brown solid:

$$2 \text{ AgNO}_3 + 2 \text{ NaOH} \rightarrow \text{Ag}_2\text{O}(\text{s}) + 2 \text{ NaNO}_3 + \text{H}_2\text{O}$$

In the next step, sufficient aqueous ammonia is added to dissolve the brown silver(I) oxide. The resulting solution contains the $[Ag(NH_3)_2]^+$ complexes in the mixture, which is the main component of Tollens' reagent. Sodium hydroxide is reformed:

$$Ag_2O(s) + 4 NH_3 + 2 NaNO_3 + H_2O \rightarrow 2 [Ag(NH_3)_2]NO_3 + 2 NaOH$$

Aldehydes/Reducing sugars are easily oxidized by mild oxidizing agents such as Ag^+ . The silver mirror test is the reaction of a sample with a solution containing silver-ammonia complex ions. When this reagent oxidizes the aldehyde, the silver ions are reduced to metallic silver, which forms a black precipitate, and if the test tube is clean, a silver mirror on the test tube. The overall reaction is as follows:

 $\begin{array}{c} O\\ =\\ 2Ag(NH_3)_2^+ + 2OH^- + R - \overset{\parallel}{C} - H \longrightarrow 2Ag_{(s)} \text{ (silver mirror)} + R-COO^- + NH4^+ + 3NH_3 + H_2O \end{array}$

MATERIALS:

Silver nitrate (AgNO3; > 99.5% purity), Sodium hydroxide, D-Glucose anhydrous were purchased from SD fine (India). All chemicals were of analytical reagent grade and were used without further purification. Other chemicals used in this project extracted from natural sources.

METHODS:

Preparation of silver nanoparticles (AgNP):

A mixture of 5ml 0.001M AgNO₃ and 5ml 0.001M NaOH solution mixed to form turbid precipitated solution. To this, aqueous ammonia was slowly added to make clear solution of Tollens reagent, followed by the addition of 5ml 0.001M glucose solution. The mixture is taken in 10ml vials and immersed in a sonication bath for one hour for synthesis of AgNP. These aqueous dispersions of silver nanoparticles were also very stable and no sedimentation formed even after being stored for a longer periods. The solution of AgNP was then diluted to 2, 3, 4 folds with de-ionized water.

Coating of AgNP on Paper currency notes:

The Paper currency notes initially cleaned with cotton wool to remove dust and dirt particles and then AgNP solution sprayed on it uniformly an swapped with cotton wool dipped in dilute Lemon fruit juice (*Citrus limon L.*) as it stabilized the AgNP on Paper currency notes and also reduces any traces of Ag^+ ions left over [40]. The AgNP plausibly deposited on cellulose fibers of Paper currency notes which provide firm binding.

Coating of AgNP on Mobile phone Screen:

Coating of AgNP on mobile phone screen (tempered glass) is difficult as Polyethylene terephthalate (PET) or Thermoplastic polyurethane (TPU) which provided smooth surface over which adsorption or binding of AgNP is not possible. Therefore we opted a innovative approach, in which Bio-film of orange peel extract was initially coated over mobile phone screen. The bio-film cross-linked with polymeric material of screen and provides a transparent texture. Upon this layer AgNP solution sprayed uniformly and wiped with cotton wool dipped in dilute Lemon fruit juice (*Citrus limon L.*) for stabilization of nanoparticles.

Characterization of AgNP:

The AgNP samples collected by scratching AgNP treated Paper currency notes were characterized by using Scanning electron microscope (SEM). Morphology of the samples was studied using a scanning electron microscope with a detector of back-scattered electrons (SEM-BSE) equipped with an energy dispersive spectrometer (EDS) which allows to determine precisely elemental composition of materials. Samples for SEM-BSE were prepared according to the standard procedure, fixed with conductive glue and covered with a thin gold layer. The thermal investigation of cellulose fibers with silver particles was carried out with Q 500 TGA (T. A. Instruments) from 30 to 600°C at a heating rate of 10°C per min, under a constant flow (60 ml/min) of nitrogen gas.

Antimicrobial Screening:

The efficacy of AgNP was tested by Antimicrobial Screening on both bacterial (Gram positive and Gram negative strains) and fungal species.

Antibacterial activity:

Inoculums of bacterial strains were prepared by using Nutrient broth (pH 7.2) and for antibacterial screening the agar medium was sterilized by autoclaving at 120°C for 15 min. The Petri plates and pipettes were sterilized by dry heat in a hot- air oven at 150°C for 1 hr. About 20 mL of the molten agar medium was poured in each of sterilized Petri plates. The microorganisms employed in this study were one gram positive bacteria; Staphylococcus aureus (MTCC - 96) and one gram negative bacteria, Pseudomonas aeruginosa (MTCC - 424). The inoculum was standardized at 1*106 CFU/ml comparing with turbidity standard (0.5 MacFarland tube). The AgNP solutions with different concentrations (standard, 2, 3, 4 fold dilutions) were spray uniformly over sterilized whatman filter paper followed by wiping with cotton wool dipped in dilute Lemon fruit juice. This paper was cut into 5 mm discs and were screened in vitro for their antibacterial activity by the cup-plate agar diffusion method [44]. The auto calved Nutrient broth media Inoculation of Pseudomonas aeruginosa (gram-negative) and Staphylococcus aureus (gram-positive) were Incubate over night at 37° C in shaker for Bacterial growth. From this 0.3ml of bacterial culture was taken and inoculated by using spreader on freshly prepared auto calved agar plates. After drying of plate prepared AgNP 5 mm sample discs were kept on microbial plate along with positive controls Norfloxacin for Staphylococcus and Pseudomonas strains. After overnight incubation at 37° C in BOD incubator zone of inhibition is measured by measuring scale. The zone of inhibition (in mm) was compared with standard drugs.

Antifungal activity:

Sclerotium rolfsii inocculam was inoculated to the freshly prepared sterilized Potato Dextrose Broth and allowed for fungal growth. After the growth of fungus, inoculum was added to the sterilized PDA plates for anti fungal Activity. Further, 5 mm AgNP discs were prepared by using whatman filter paper as discussed above. The Inoculation of *Sclerotium rolfsii* fungal strain which were obtained from MTCC in autoclaved PDB media and incubate for 3-4 days at 30° C in shaker for fungal growth. From that 20 μ l of Fungal culture was taken and inoculated by inoculation loop on freshly prepared autoclaved agar plates. Different 5 mm AgNP discs were kept on microbial plate along with antifungal agent as a control i.e. Ketoconazole. These plates were incubated for 5-6 days at 30° C in BOD incubator and zone of inhibition (in mm) is measured by measuring scale.

RESULTS AND DISCUSSIONS:

(Data Analysis-Findings)

It is evident from SEM-BSE micrographs of AgNP-cellulose composites (**Fig-3**) obtained for 0.001M concentrations of the applied AgNO₃ solution that the silver nanoparticles present on the surface of cellulose fibers of Paper currency notes. The size and shape of the silver particles varies from 20nm to 50nm. Whereas the scratched material of mobile phone screens also confirmed the presence of AgNP with size and shape of the silver particles varies from 15nm to 50nm. (**Fig-4**)

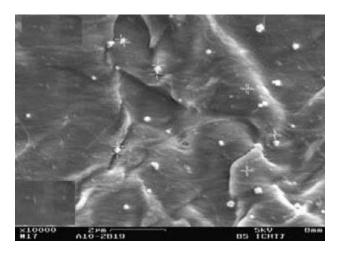
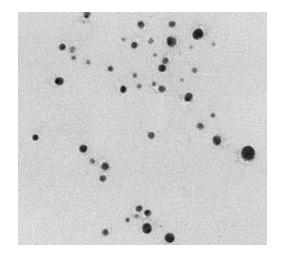
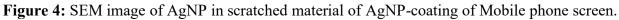


Figure 3: SEM image of AgNP distributed on cellulose fibers of Paper currency notes.





From the anti-microbial studies it is evident that the AgNP treatment on Paper currency notes and cell phone screens provide potential antimicrobial barrier against both bacteria and fungi. The *in vitro* antimicrobial (anti-fungal & anti-bacterial) results are tabulated in Table 1.

Entry	Sample	P. aeruginosa	S. aureus	S.rolfsii
1	Standard	7	8	9
2	2-fold dilution	7	6	8
3	3-fold dilution	6	5	2
4	4-fold dilution	2.5	2.5	2
5	5-fold dilution	2.5	2.5	2
6	Control ^[a]	11	15	5

Table 1. Antimicrobial activity (Zone of inhibition in mm) of different samples (Different dilutions)

[a] Controls: Norfloxacin for S. aureus, and P. aeruginosa and Ketoconazol for S. rolfsii.

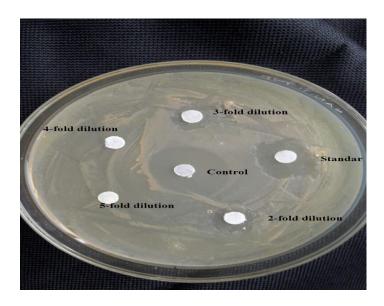


Figure 5. Antimicrobial activity (Zone of inhibition in mm) against *P.aeruginosa*.

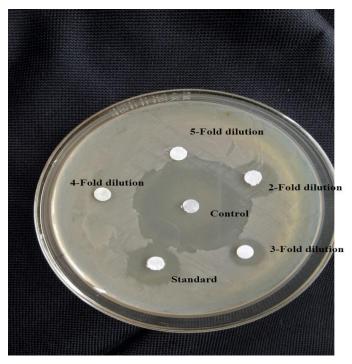


Figure 6. Antimicrobial activity (Zone of inhibition in mm) against S. aureus.

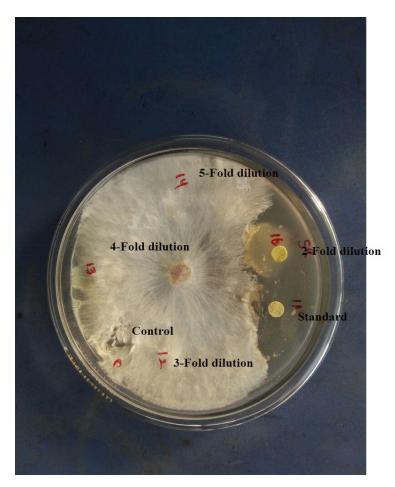


Figure 7: Antimicrobial activity (Zone of inhibition in mm) against S. rolfsii.

The presence of definite zone of inhibition surrounding the disc indicated antimicrobial activity. All the samples have shown Excellent to good antibacterial activity against both Gram positive bacteria and Gram negative bacteria compared to the standard antibiotic Norfloxacin. The samples up to 3-fold dilutions have shown excellent activity but beyond that (4 and 5-fold dilutions) antibacterial activity drastically decreased. Whereas, anti-fungal activity of the AgNP samples showed more potentiality compared to standard antifungal agent Ketoconazol especially for standard and 2-fold diluted AgNP samples.

This benign process is completely eco-friendly as the chemicals used are non-hazardous and extracted from natural sources. moreover the procedure of coating AgNP is very economical as each 500 Rs Paper currency note or Mobile phone screen needed 20 pisa only.

CONCLUSION:

From the current project it is clear that AgNP coating on Paper currency notes and Mobile phone screens by using innovative modified Tollens process provide an excellent protection against microbial contamination which will be fatal in nosocomial infections caused by microorganisms in hospitals when contaminated Paper currency notes and Mobile phones handled by healthcare associates. The problem of cross contamination will be serious in patients with immunodeficiency disorders like HIV/AIDS or patients who were underwent organ transplantation to whom usually immune-depressant drugs were given. The outlook of the projects also opens new avenue for public health care aspects especially in situations where multidrug bacterial strains like *mycobacterium tuberculi* evolved by excessive use of antibiotics. The process is very convenient, ecofriendly and economical to scale up for bulk usage.

SUGGESTIONS:

Further studies should investigate the combination of AgNPs and antibiotics against resistant hospital strains for the development of new materials and substances for medical application. As "*prevention is better than cure*" the following suggestions were made basd on this project:

- The paper currency notes should be printed with inks impregnated with the AgNP.
- Automatic Teller Machines (ATM) should be reloaded with AgNP suspension for spraying on used Paper currency notes.
- The interiors of hospitals to be painted with AgNp impregnated paints to prevent Nosocomial infections and to keep the intensive care units (ICU) hygiene and sterile.
- To prevent contagious diseases like swine flu, the public transport utilities like buses and metros to be coated with AgNP impregnated paints.
- The mobile phone screen guards to be manufactured with AgNP incorporated polymers.

ACKNOWLEDGEMENTS:

We are thankful to Ciencia Life sciences, Hyderabad, India for providing bacterial and fungal strains for antimicrobial screening.

ABBREVIATIONS

rRNA	Ribosomal ribonucleic acid
AgNP	Silver Nano-particles
DNA	Deoxyribonucleic acid
MRSA	Methicillin-resistant Staphylococcus aureus
EB	Electron beam
SEM	Scanning electron microscope
BSE	Back-scattered electrons
EDS	Energy dispersive spectrometer
MTCC	Microbial Type Culture Collection
PDA	Potato Dextrose Agar
PDB	Potato Dextrose broth
BOD	Biochemical oxygen demand
HIV	Human immunodeficiency virus
AIDS	Acquired immune deficiency syndrome

REFERENCES:

- Angelakis, E., Azhar, E.I., Bibi, F., Yasir, M., Al-Ghamdi, A.K., Ashshi, A.M., Elshemi, A.G., Raoult, D. (2014) Paper money and coins as potential vectors of transmissible disease. *Future Microbiol.* 9(2), 249–261. https://doi.org/10.2217/fmb.13.161
- Amira H. A. Al-Abdalall (2010) Isolation and identification of microbes associated with mobile phones in Dammam in eastern Saudi Arabia. *J Family Community Med.* 17(1), 11–14. doi: 10.4103/1319-1683.68783
- Ekrakene T, Igeleke C.L. (2007) Micro-organisms associated with public mobile phones along Benin-sapele Express Way, Benin City, Edo State of Nigeria. J Appl Sci Res. 3, 2009–12.
- Gales, A.C., Torres, P.L., Vilarinho, D.S., Melo, R.S., Silva, C.F., Cereda, R.F.(2004) Carbapenem resistant *Pseudomonas aeruginosa* outbreak in an intensive care unit of a teaching hospital. *Braz J Infect Dis.*, 8(4), 267–271. doi:10.1590/s1413-86702004000400001
- Menezes, E.A., Silveira, L.A., Cunha, F.A., et al. (2003) Perfil de resistencia aos antimicrobianos de *Pseudomonas* isoladas no Hospital Geral de Fortaleza [Antimicrobials profile resistance from isolated pseudomonas at the Fortleza's General Hospital]. *Rev Bras Anál Clín.*, 35(4), 177-180. Portugese.
- Dastjerdi, R., Montazer, M. (2010) A review on the application of inorganic nanostructured materials in the modification of textiles: focus on antimicrobial properties. *Colloids Surf B Biointerfaces.*, 79(1), 5–18. doi:10.1016/j.colsurfb.2010.03.029
- Yoon, K., Hoon Byeon, J., Park, J.H., Hwang, J. (2007) Susceptibility constants of Escherichia coli and Bacillus subtilis to silver and copper nanoparticles. Sci. Total. Environ., 373(2–3):572–575. doi:10.1016/j.scitotenv.2006.11.007
- Rai, M.K., Deshmukh, S.D., Ingle, A.P., Gade, A.K. (2012) Silver nanoparticles: the powerful nanoweapon against multidrug-resistant bacteria. *J. Appl. Microbiol.*, 112(5), 841–852. doi:10.1111/j.1365-2672.2012.05253.x

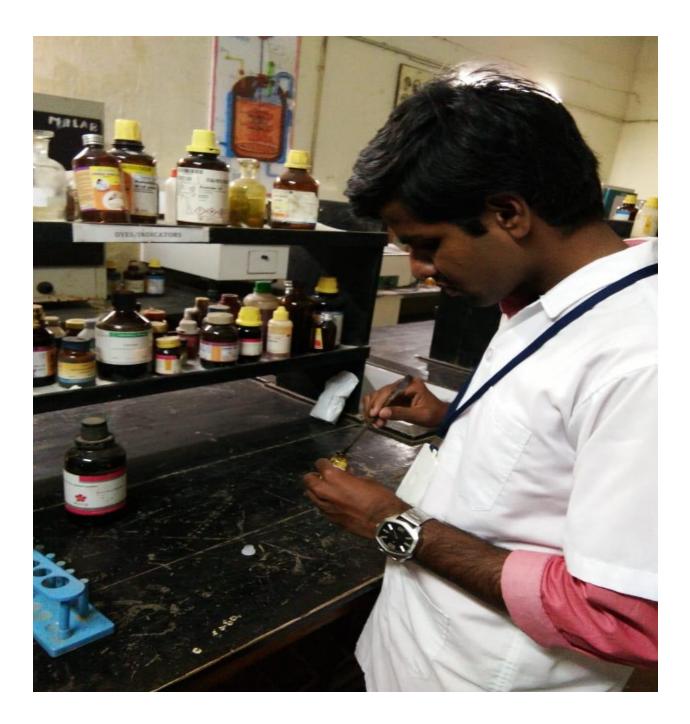
- 9. Franci, G., Falanga, A., Galdiero, S., et al. (2015) Silver nanoparticles as potential antibacterial agents. *Molecules.*, 20(5), 8856–8874.doi:10.3390/molecules20058856
- Salomoni, R., Leo, P., Rodrigues, M.F.A. (2015) Antibacterial activity of silver nanoparticles (AgNPs) in *Staphylococcus aureus* and cytotoxicity effect in mammalian cells. *Formatex Microbiol.*, 5, 851–857.
- Brigger, I., Dubernet, C., Couvreur, P. (2002) Nanoparticles in cancer therapy and diagnosis. *Adv. Drug. Deliv. Rev.*, 54(5), 631–651. doi:https://doi.org/10.1016/j.addr.2012.09.006
- Rai, M., Yadav, A., Gade, A. (2009) Silver nanoparticles as a new generation of antimicrobials. *Biotechnol. Adv.*, 27(1), 76–83. doi: https://doi.org/10.1016/j.biotechadv.2008.09.002
- Sudarenkov, V. (2013) Nanotechnology: balancing benefits and risks to public health and the environment. Strasbourg: Council of Europe, Committee on Social Affairs, Health and Sustainable Development, 2013.
- 14. Nordberg, G.F., Fowler, B.A., Nordberg, M., Friberg, L.T. (2007) Handbook on the Toxicology of Metals. 3rd ed. San Diego: Elsevier.
- 15. Marcone, G.P.(2011) Avaliacao da ecotoxicidade de nanoparticulas de dioxide de titanio e prata [Doutorado] [Assessment of ecotoxicity of nanoparticles of titatinium dioxide and silver]. Campinas: UNICAMP; 2011. Portuguese.
- Salomoni, R., Léo, P., Montemor, A.F., Rinaldi, B.G., Rodrigues, M.F.A.(2017) Antibacterial effect of silver nanoparticles in *Pseudomonas aeruginosa, Nanotechnol. Sci. Appl.*, 10, 115–121. doi: 10.2147/NSA.S133415
- Lansdown, A.B. (2006) Silver in health care: antimicrobial effects and safety in use. *Curr. Probl. Dermatol.*, 33,17–34. doi:10.1159/000093928
- Chen, J., Han, C.M., Lin, X.W., Tang, Z.J., Su, S.J. (2006) Effect of silver nanoparticle dressing on second degree burn wound. *Zhonghua Wai Ke Za Zhi*. 44(1), 50–52.
- Lee, H.Y., Park, H.K., Lee, Y.M., Kim, K., Park, S.B. (2007) A practical procedure for producing silver nanocoated fabric and its antibacterial evaluation for biomedical applications. *Chemic Communic (Camb).*, 28, 2959–2961. doi: 10.1039/B703034G

- Cohen, M.S., Stern, J.M., Vanni, A.J., et al. (2007) *In vitro* analysis of a nanocrystalline silver-coated surgical mesh. *Surg Infect (Larchmt)*. 8(3), 397–403. doi:https://doi.org/10.1089/sur.2006.032
- Pittet, D., Allegranzi, B., Sax, H. *et al.*(2006) Evidence-based model for hand transmission during patient care and the role of improved practices. *Lancet Infect. Dis.* 6(10), 641–652. doi:https://doi.org/10.1016/S1473-3099(06)70600-4
- 22. Tolba, O., Loughrey, A., Goldsmith, C.E., Millar, B.C., Rooney, P.J., Moore, J.E. (2007) Survival of epidemic strains of nosocomial- and community-acquired methicillinresistant *Staphylococcus aureus* on coins. *Am. J. Infect. Control* 35(5), 342–346. doi: https://doi.org/10.1016/j.ajic.2006.10.015
- 23. Todd, E.C., Greig, J.D., Bartleson, C.A., Michaels, B.S.(2009) Outbreaks where food workers have been implicated in the spread of foodborne disease. Part 6. Transmission and survival of pathogens in the food processing and preparation environment. J. Food Prot. 72(1), 202–219. doi:10.4315/0362-028x-72.1.202
- 24. Todd, E.C., Greig, J.D., Bartleson, C.A., Michaels, B.S. (2008) Outbreaks where food workers have been implicated in the spread of foodborne disease. Part 4. Infective doses and pathogen carriage. J. Food Prot., 71(11), 2339–2373. doi:10.4315/0362-028x-71.11.2339
- Melnick, J., Edward, A. Medical Microbiology. 23th ed. New York: McGraw-Hill Professional; 2004.
- 26. Hui Y.H., Sattar, S.A., Murrell, K.D., Nip, W.K., Stanfield, P.S., Food borne disease handbook. Viruses, parasites, pathogens and HACCP. (2nd ed) 2001;2
- Brande, A.I., Davis, C.E., Fraver, J. (1981) Food borne microbiology infections diseases. *Philadelphia: W.B. Sanders Company;* 1981. p. 1860.
- O'Hara, C.M., Brenner, F.W., Miller, J.M. (2000) Classification, identification, and clinical significance of Proteus, Providencia, and Morganella. *Clin Microbiol Rev.*, 13, 534–46. [PMCID: PMC88947] [PubMed: 11023955]
- Liu, P.Y., Gur, D., Hall, L.M. (1992) Survey of the prevalence of β-lactamases amongst 1000 gram-negative bacilli isolated consecutively at the Royal London Hospital. J Antimicrob. Chemother., 30, 429–47. [PubMed: 1490917]

- Centers for Disease Control and Prevention. National Nosocomial Infections Surveillance (NNIS) report, data summary October 1986– April 1996, issued May 1996. A report from the National Nosocomial Infections Surveillance (NNIS) System. Am. J. Infect. Control., 1996;24:380–8. [PubMed: 8902113]
- Ramos JL, editor. Pseudomonas. New York: Kluwer Academic/Plenum Publishers; 2004.
 p. 2132.
- Ollis, D.F., El-Akabi, H.(1993) Photocatalytic purification and treatment of water and air. Amsterdam: Elsevier Science Ltd., 957-61. ISBN-13: 978-0444898555
- Gong P, Li H, He X, Wang K, Hu J, Tan W. (2007) Preparation and antibacterial activity of Fe₃O₄ Ag nanoparticles. Nanotechnology, 18, 604-11.
- 34. Ahmad, Z., Pandey, R., Sharma, S., Khuller, G.K. (2005) Alginate nanoparticles as antituberculosis drug carriers: formulation development, pharmacokinetics and therapeutic potential. Indian J. Chest. Dis. Allied. Sci., 48, 219-26.
- 35. Banawas, S., Abdel-Hadi, A., Alaidarous, M., Alshehri, B., Dukhyil, A.Z.B., Alsaweed, M., Aboamer, M. (2018) Multidrug-Resistant Bacteria Associated with Cell Phones of Healthcare Professionals in Selected Hospitals in Saudi Arabia. *Canadian Journal of Infectious Diseases and Medical Microbiology*, Volume 2018, Article ID 6598918, 1-7. doi: https://doi.org/10.1155/2018/6598918
- 36. Morubagal, R.R., Shivappa, S.G., Mahale, R.P., Neelambike, S.M. (2017) Study of bacterial flora associated with mobile phones of healthcare workers and non-healthcare workers. *Iran J. Microbiol.*, 9(3), 143-151.
- 37. Alananbeh, K.M., Al-refaee, W.J. Al-qodah, Z. (2017) Antifungal Effect of Silver Nanoparticles on Selected Fungi Isolated from Raw and Waste Water. *Indian J Pharm Sci.*, 79(4), 559-567. doi: 10.4172/pharmaceutical-sciences.1000263
- Yin, Y., Li, Z.Y., Zhong, Z., Gates, B., Xia Y., Venkateswaran, S. (2002) Synthesis and characterization of stable aqueous dispersions of silver nanoparticles through the Tollens process. *J. Mater. Chem.*, 12, 522–527. doi:10.1039/B107469E
- Dhulappanavar, G., Hungund, B., Ayachit, N. (2011) Characterization of Silver Nanoparticles Biosynthesized Using Lemon Juice. *Proceedings of the ICONSET- 2011, IGCAR*, Kalpakkam, 28-30 November, 2011. doi: 10.1109/ICONSET.2011.6167936

- 40. Vankar, P.S., Shukla, D. (2012) Biosynthesis of silver nanoparticles using lemon leaves extract and its application for antimicrobial finish on fabric. *Appl Nanosci* 2, 163–168 doi:10.1007/s13204-011-0051-y
- Dagmara K. Chmielewska, D.K., Bożena Sartowska, B., Wojciech Starosta, W., Marta Walo, M. (2010) Radiation synthesis of silver nano and microparticles in cellulose fibers. *NUKLEONIKA*, 55(3), 345–349.
- Smiechowicz, E., Niekraszewicz, B., Kulpinski, P., Dzitko, K. (2018) Antibacterial composite cellulose fibers modified with silver nanoparticles and nanosilica. *Cellulose*, 25(6), 3499-3517. doi:10.1007/s10570-018-1796-1
- Tollens, B. (1882). "Ueber ammon-alkalische Silberlösung als Reagens auf Aldehyd" [On an ammonical alkaline silver solution as a reagent for aldehydes]. Berichte der Deutschen Chemischen Gesellschaft (in German). 15(2), 1635–1639. doi:10.1002/cber.18820150243.
- (a) Singh, H., Dhar, L., Yadav, S., Shukla, K.N., Dwivedi, R. (1990) J. Agric. Food. Chem. 1962-1964, 38. (b) D. Eugene & Weinberg, Burger's Medicinal Chemistry & Drug Diseases, 1997, Fifth Edition, 2, Therapeutic agents, 637. (c) Bennet, J.E., (1996) In Goodman Gilman's the Pharmacological Basis of Therapeutics, Ninth Edition, 1175.
 (d) Gomes, A.T., Smânia Jr, A., Seidel, C., Smânia, E.F.A., Honda, N.K., Roese, F.M. Muzzi, R.M. (2003) Braz. J. Microbiol. 34, 194-196. (e) Hadacek, F., Greger, H., (2000) Phytochem. Anal., 11, 137-147. (f) Hamburger, M.O., Cordell, G.A., (1987) J. Nat. Prod., 50, 19-22. (g) Heneine, I.F. (2000) Biofisica Básica, Atheneu, São Paulo. (h) Hostettman, K., Wolfender, J.L., Rodriguez, S. (1997) Planta Med., 63, 2-10. (i) US 3592932, Ciba Ltd.; Microbiology Abstr., 1974, Vol. 9, No. 2, 9A, 977. (j) US 3705903, Lilly Ind. Ltd.; Microbiology Abstr., 1975, Vol. 10, No. 3, 10A, 1636. (k) US 3702363, Ciba-Geigy AG.; Microbiology Abstr., 1975, Vol. 10, No. 3, 10A, 1771.
- 45. https://en.wikipedia.org

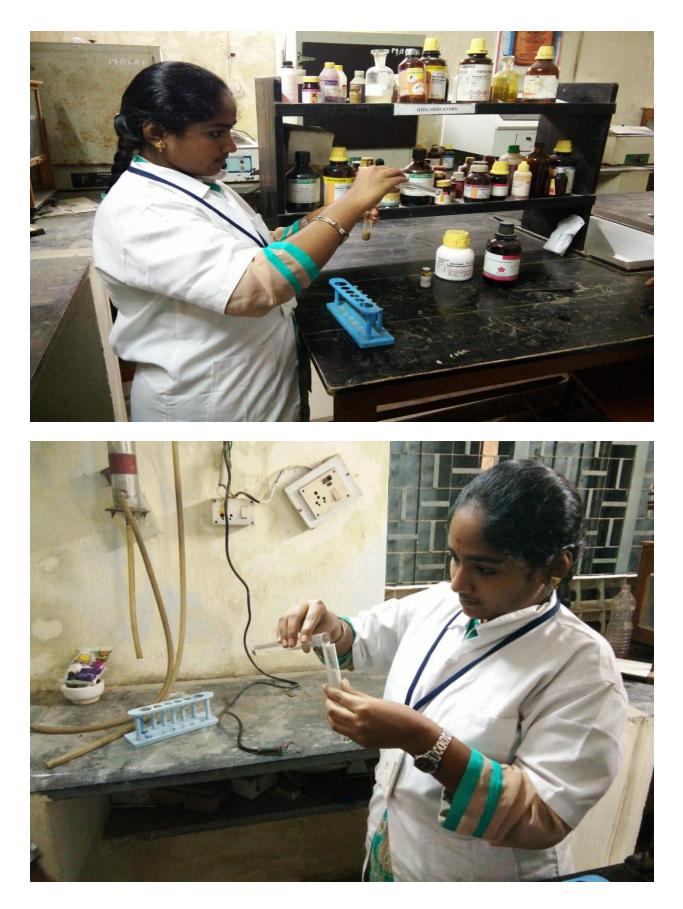
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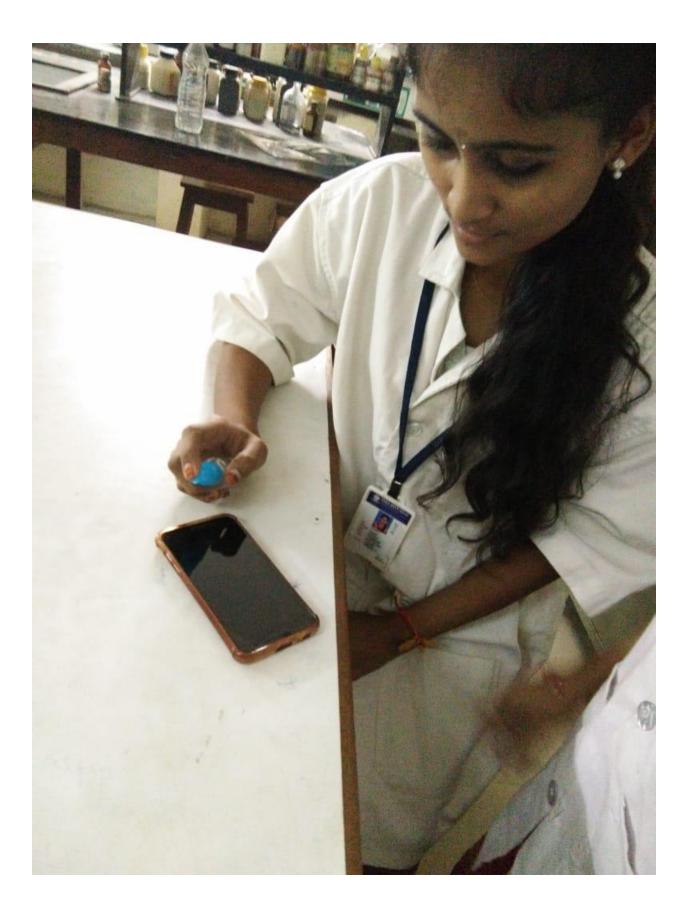


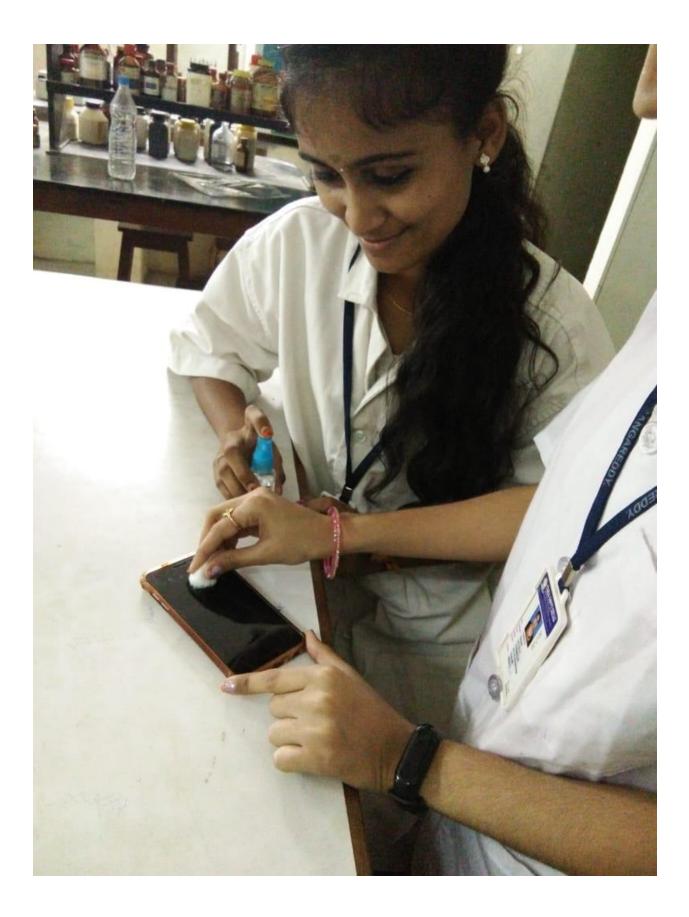


















For Academic and Scientific Cooperation Memorandum of Understanding (MoU) Ciencia Life Sciences, Hyderabad





TARA GOVERNMENT COLLEGE, Department of Chemistry, Sangareddy.

Managing Director Ciencia Life Sciences, Hyderahad,

Tara Govt. College(A), Sangareddy. ~

Department of Chemistry, Tara Govt. College (A), Sangareddy Head of Department,



For Academic and Scientific Cooperation **Memorandum of Understanding (MoU) Ciencia** Life Sciences, Hyderabad



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TARA GOVERNMENT COLLEGE, Department of Chemistry, (AUTONOMOUS) Sangareddy.

Ciencia Life Sciences, Hyderabad. Managing Director



Head of Department,

Tara Govt. College (A), Sangareddy Department of Chemistry,

File No.CCE-AC/QLTY/NAAC/2/2018-ACADEMIC CELL

PROCEEDINGS OF COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA PRESENT: SRI.NAVIN MITTAL, IAS.

Sub:- Accreditation of GDCs-New system of Assessment by NAAC- Status of preparation for NAAC Reaccreditation-Visits to certain GDCs-Reg. Ref:- <u>CCE-AC/QLTY/NAAC/2/2018-ACADEMIC CELL</u>

Vide reference read, certain GDCs due for NAAC reaccreditation have submitted the status of preparation to the Commissioner of Collegiate Education. In this connection, GDC (W) Siddipet has submitted IIQA and is preparing SSR for fresh accreditation.

In view of the above, Principal, GDC (W) Siddipet has requested Commissioner, Collegiate Education for the guidance of Sri.A.Vishweswara Sharma, Asst. Professor of English, GDC (A) Sangaredy, in the submission of data on College website from 28th to 31st January 2020.

In this connection, Principal, GDC (A) Sangareddy is informed that the absence of above mentioned faculty shall be considered as ON DUTY.

The faculty visiting the college have to submit the report on the actual status of preparation in respective GDC on or before 03.02.2020.

Signature Not Verified Digitally signed by PODILA BALA BHASKAR Date: 2020.01.31 10:49:57 Reason: Approved For Commissioner of Collegiate Education

То

The Principals of GDCs concerned

File No.CCE-AC/QLTY/NAAC/2/2018-ACADEMIC CELL



(Affiliated to Osmania University & Accredited by NAAC with 'B' grade, 2.75 CGPA)

SANGAREDDY-502 001, SANGAREDDY (Dist.), T.S.

E-mail: prl-gdc-srd-ce@telangana.gov.in Website: www.taragc.ac.in

Dr. Chandra Mukherji, M.A., M.Phil., Ph.D. Principal Mobile: 07032960368 E-mail: tara.sangareddy@gmail.com

Lr.Rc.No:190/TGC-SRD/2019-20

06 January 2020

TO The Commissioner of Collegiate Education, Telangana State, Nampally, Hyderabad-500001.

Esteemed sir,

- Sub: Joining of Smt.T.Navya, Contract Lecturer in Commerce, Govt. Degree College (W), Sangareddy- Identified the Surplus Contract lecturers due to Relocation of regular faculty- Submission of joining report -**Reg**.
- Ref: 1.CCE-SER/2/CLEC/9/2019-SER@,Dt:31.12.2019
 - 2. Relieving letter from GDC(W), Sangareddy
 - 3. Applicant Joining Letter, Dt:03.01.2020

In obedience to the orders of CCE reference cited, I submit that Smt.T.Navya , Contract Lecturer in Commerce, Govt. Degree college(W),Sangareddy.The Hon'ble CCE Identified the surplus Contract Lecturers due to relocation of regular faculty she has been transferred to Tara Govt. College(A),Sangareddy. She has reported to duty on 03.01.2020 AN at Tara Govt. College(A), Sangareddy.

This is for your kind information.

Thanking you

Yours faithfully

TARA AUTONO

SANGAREDDY-502 001

Date 03/01/2020

TO The Poincipal, Taxa Gort. Degree College(A), Sangareddy.

- Sub: Contract lecturers working in Gout Degree Colleges in the state - Identified the surplus contract lecturers - due to relocation of regular faculty - Joining Report.
- Ref: File NO. CCE-SER2 | CLEC/9 | 2019 SER2 Govt · memo NO. 2859 | CE | AI | 2015, Dt. 08.06.2019. Counselling Conducted on 13.12-2019 and optimy given by the Individuals.

3/01/2020

Respected Madam,

N/a immediat

(nerroe

" wowely

I am T. Navya, contract lecturer in Commune here with submitting my Joining Report in the Tara Govt. Degree college(A), to day i.e on 03/01/2020.

Thanking You madam.

Your's Faithhully Du T-Navya

GOVERNMENT OF TELANGANA COLLEGIATE EDUCATION DEPARTMENT

From: Dr.Anita,M.A., M.Phil.,Ph.D., Principal, Govt. Degree College for Women, Sangareddy, Sangareddy Dist.

То

The Principal, Govt. Degree College (A), <u>SANGAREDDY</u>

Lr.Rc.No.A1/356 /GDCWS/2020

Date: 02-01-2020.

Respected Madam,

 Sub: - Contract Faculty- Relocation of Contract Lecturers Working in Government Degree Colleges- Relieving of Smt.T.Navya, Contract Lecturer in Commerce due to posting at Tara Government College(A), Sangareddy - Reg.
 Ref:- Proc. File No. CCE-SER2/CLEC/9/2019-SER2 Dt. 31-12-2019 of C.C.E., T.S., Hyd.

-X-X-X-

With reference to the subject cited above, I am herewith relieving Smt. T. Navya,

Contract Faculty in Commerce of Govt. Degree College for Women, Sangareddy as per the

Orders of the C.C.E., T.S., Hyderabad vide reference cited above on 03-01-2020 F.N.

Smt. T. Navya, C.F. in Commerce of this college is hereby instructed to report at her new station i.e., Tara Government College(A), Sangareddy, Sangareddy District.

Copy to Smt.T.Navya, Contract Lecturer in Commerce The Commissioner of Collegiate Education, Hyderabad The pay Bills Spare PRINCIPAL Govt. Degree College (W) Sangareddy.

File No.CCE-SER2/CLEC/9/2019-SER2

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION TELANGANA STATE :: HYDERABAD

PRESENT: NAVIN MITTAL, I.A.S.,

- Sub:- Contract faculty- Contract Lecturers working in Government Degree Colleges in the State – Identified the surplus Contract Lecturers due to relocation of regular faculty - Relocation of Contract Lecturers Working in Government Degree Colleges - Orders -Issued.
- Read:- 1. Govt.Memo.No.2859/CE/A1/2015, dated 08.06.2019
 2. File NoCCE-AC/GEN/98-Academic Cell (Computer No.157641) dated 19.12.2019.
 3. Counselling conducted on 13.12.2019 and Options given by the individuals.

The Commissioner of Collegiate Education, Telangana State, Hyderabad have identified the surplus Contract Lecturers due to relocation of regular faculty. In reference 2nd read above a counselling was held on 13.12.2019 at IPGDC(W), Nampally and they opted the places as per their choice.

In the circumstances reported in references 1 to 3rd read above, the Commissioner of Collegiate Education, Telangana State, Hyderabad is hereby Relocate the following Contract Lecturer working in Government Degree College as mentioned against his/her name as shown hereunder:-

	Name of the Contract Lecturer	Subject	Presently working	Relocated at
79	T. Navya	Commerce	GDC (W) Sangareddy	GDC (A) Sangareddy

File No.CCE-SER2/CLEC/9/2019-SER2

Further, the Principals of Government Degree College concerned are requested to report compliance in the matter promptly.

The receipt of these proceedings shall be acknowledged.

(Orders issued with the approval of the Commissioner of Collegiate Education, Telangana State, Hyderabad)

Signature Not Verified Digitally signed by N VIJAYA BHASKAR RAO Date: 2019.12.31 16:55:26 IST Reason: Approved

For COMMISSIONER OF COLLEGIATE EDUCATION

То

The Principal, Government Degree Colleges as mentioned above. Copy to the Individual (through Principal of the College) Copy to the DTO / STO concerned Copy to the GDC-CS concerned

File No.CCE-SER2/DPOD/11/2020-SER2

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION TELANGANA, HYDERABAD

Present: Sri Navin Mittal, I.A.S.

- Sub:- TSCES Government Degree Colleges Permission to work on "On Duty" basis Orders Issued.
- Read: U.O Note. File No. CCE-AC/GEN/41/2019-ACADEMIC CELL, Dated.21.10.2020.

In the circumstances stated in the reference read above, the Commissioner of Collegiate Education, Telangana, Hyderabad is hereby directed to work on on "On Duty" basis for the Academic year 2020-21 to the following teaching staff as follows.

SI.	Zone	Name of the	Subject	Present	On Duty
No.		faculty		place of	at
				working	
1	VI	Dr.Rapolu	Telugu	GDC (A),	GDC (W),
		Srinivas		Sangareddy	Nalgonda

The Principal, Government Degree College(A), Sangareddy is requested to claim his salary from the parent institution on production of attendance certificate every month issued by the Principal, Government Degree College(W), Nalgonda.

Both the Principals of Government Degree Colleges are requested to report compliance immediately

The receipt of these proceedings should be acknowledged by return of post.

File No.CCE-SER2/DPOD/11/2020-SER2

(Orders of the Commissioner of Collegiate Education have been obtained in the note file bearing No.CCE-AC/GEN/41/2019-ACADEMIC CELL)

Signature Not Verified Digitally signed by YADAGIRI GOSTKA Date: 2020.11.23 13:00:20 IST Reason: Approved For COMMISSIONER OF COLLEGIATE EDUCATION

То

1) Dr.Rapolu Srinivas, Assistant Professor of Telugu,

Government Degree College(A), Sangareddy

(through the Principal of the College).

- 2) The Principal, Government Degree College(A), Sangareddy.
- 3) The Principal, Government Degree College(W), Nalgonda.

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI.NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education-Department of Telugu, Osmania University, Hyderabad- Preliminary meeting for revision of UG and PG syllabus in Telugu subject at Head & Chairperson BoS of Telugu Chamber, Department of Telugu, University College of Arts & Social Science, Osmania University, Hyderabad on 25th February 2020-Permission accorded to the list of faculty from various GDCs of Telangana State to attend the meeting-Orders-Regarding.
- **Ref:** Letter received from the Head, Department of Telugu, Osmania University, Hyderabad, dated:24.02.2020.

Vide reference read, the Commissioner of Collegiate Education has accorded permission to the list of following faculty members from various GDCs of Telangana State to participate in **preliminary meeting for revision of UG and PG syllabus in Telugu subject** at Head & Chairperson BoS of Telugu Chamber, Department of Telugu, University College of Arts & Social Science, Osmania University, Hyderabad on **25th February 2020**.

S.No.	Name of the Lecturers	College Name
1	Dr. J. Bharathi	Vivekananda Degree College, Hyderabad.
2	Dr.Uma Shashi, Principal	Govt. Women's Degree College, Gajwel.
3	Dr. Koyi Koteswar Rao	Govt. Degree college for womens, Nalgonda.
4	Dr. K.Narayana Murthy	BJR Degree college, Narayanaguda.
5	Dr. Veledanda Sridhar	Govt. Degree College, Hayathnagar.
6	Dr. R. Sumathi Devi	Tara Degree College, Sanga Reddy.

The absence of the above said faculty members from GDCS shall be treated as **ON DUTY** for the said date and they are not eligible to claim TA/DA from the Colleges. In view of the above, the Principals of concerned GDCs are informed to relieve them to attend the above said Seminar.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by PODILA BALA BHASKAR Date: 2020.02.25 06:29:36 IST Reason: Approved

For Commissioner of Collegiate Education

The Principals of GDCs concerned.

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI.NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education-Tara GDC, Sangareddy-Permission accorded to Dr.Chandra Mukherji, Principal, Tara GDC, Sangareddy to attend International Seminar to be held at Department of Hindi, The English and Foreign Languages University(EFLU), Hyderabad on 06.02.2020 & 07.02.2020 as Resource Person–Orders-Regarding.
- **Ref:** Email received from the Principal, Tara GDC, Sangareddy, dated: 05.02.2020.

Vide references read, the Commissioner of Collegiate Education has accorded permission to **Dr.Chandra Mukherji**, Principal, Tara GDC, Sangareddy to attend **International Seminar** to be held at Department of Hindi, The English and Foreign Languages University(EFLU), Hyderabad on **06.02.2020** & **07.02.2020** as **Resource Person**.

The absence of the above said Assistant Professor in the college shall be treated as **ON DUTY** for the said period. In view of the above, the Principal, Tara GDC, Sangareddy is informed to relieve and attend the above said Seminar.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by PODILA BALA BHASKAR Date: 2020.02.05 18:43:36 HST Reason: Approved

For Commissioner of Collegiate Education

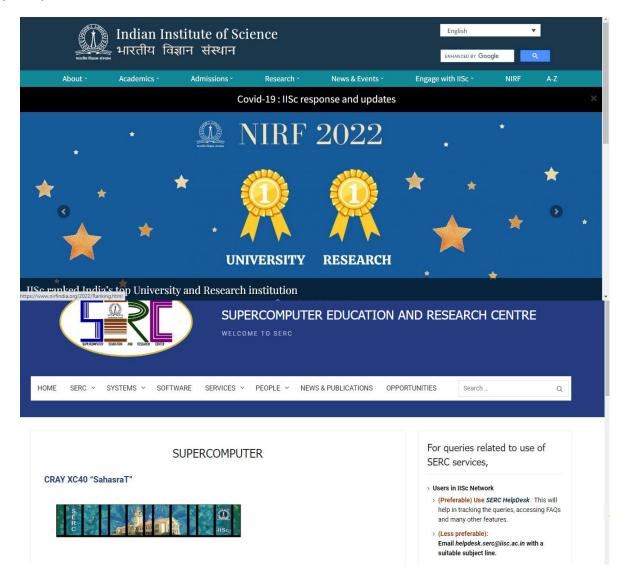
To The Principal of GDC concerned.

Collaboration with Indian Institute of Science (I.I.Sc.)

Bangalore.

The faculty members of the college are collaborating with the country's premier research institutions for the usage of laboratories/computing facilities for their own research projects.

One of the faculty members of our college, V Satya Prakash, Asst Prof of Physics, has collaborated with India's premier research institution IISc, which is India's No 1 as per NIRF 2022 rankings. This collaboration is reached in the year 2019-20 to do his research work on conductivity of electrolyte materials. As a part of this collaboration, he accessed Sahasra T Cray XC 40, country's first Peta Flops Super Computer, at Super Computing Education and Research Centre(SERC), Indian Institute of Science (IISc), Bangalore. This super computer used by him is the fourth fastest super computer in India and it ranks 327th position in the world (2019).



On Wed, 1 Jan 2020 at 15:26, Aditya Krishna Swamy adityaks@iisc.ac.in> wrote:

Dear Prof. Satyaprakash,

Greetings and best wishes for the new year 2020!

We have activated your account on SahasraT. Please find the instructions to access SERC network and SahasraT attached with this email. Also attached is a template job script. For all technical support, you may write to <u>supercomputing.serc@iisc.ac.in</u> with a copy to my email address.

Thanks, Aditya

On Wed, 4 Dec 2019 at 12:51, Aditya Krishna Swamy adityaks@iisc.ac.in> wrote:

Dear Prof. Satya Prakash,

I'm glad to update you that SERC has approved your proposal subject to the following requirements: 1. Existing software version (6.1) maybe used. If a version update is made in future, we will inform and provide you access to the same.

2. Jobs need to use a minimum of 256 cores on SahasraT CPU.

Kindly let me know if your proposed jobs will be able to meet these requirements. If yes, you may go ahead and send us the signed Agreement form (link below) and DD and we will inform you on how to access our systems.

http://www.serc.iisc.ac.in/serc_web_new/wp-content/uploads/2019/12/UsageAgreementForm-1.pdf

Let me know if you have any questions.

Thanks, Aditya

On Fri, 8 Nov 2019 at 15:40, Satyaprakash Vpet <<u>satyaprakashvpet@yahoo.in</u>> wrote:

Sir,

I am very grateful to you for your quick response.

The following are the details of using Sahasra T Cray XC 40 CPU:

1) Quantum Espresso 6.3 version I am using currently. If installing this version is possible, it is OK. Otherwise Espresso 6.1 is also OK.

2)Total core hours required are **15x45x24 =16200 core hours**(15 jobs each of taking 24 cores and 45 hours)

3)Number of **parallel runs** is **2** and typical **size of each run** is **24 cores per job** Thanking you Sir

Yours faithfully

V Satya Prakash Asst Prof of Physics Tara Govt Degree&PG

Sangareddy, Telangana, India-502001.

College(Autonomous)



OSMANIA UNIVERSITY HYDERABAD - 500 007

No. 702 /BOS/I/ Acad-1/2021

Date: 30 -08-2021

NOTIFICATION

Sub: BOARD OF STUDIES – Reconstitution of Board of Studies in Political Science (UG) under the Faculty of Social Sciences, OU -Notification - Issued.

In exercise of the powers conferred by the Ordinance No. II (as amended in the Ordinance XVI and XXX), sanction is accorded for the reconstitution of the Board of Studies in Political Science (UG) under the Faculty of Social Sciences, O.U. with the following Members:-

- 1. The Chairperson, Board of Studies in Political Science, UC A& SS, Osmana University.
- 2.* The Head, Department of Political Science, UCA&SS,O.U.
- 3. Prof. A.V. Satish Chandra, Dept. of Political Science, UCA&SS, O.U.
- 4. Dr. M. Krishna Kumar, Dept. of Political Science, UCA&SS, O.U.
- 5. Dr. Ch. Venkateshwarlu, Dept. of Political Science, UCA&SS, O.U.
- 6. Dr. Sabavath Ravi, Dept. of Political Science, UCA&SS, O.U.
 - 7. Dr. P. Venkata Ramana, Dept. of Political Science, Govt. College, Chanchalguda, Hyderabad.
- 8. Dr. Ch. Jyotsna, Dept. of Political Science, Tara Degree College, Sangareddy.
 - 9. Dr. K. Bhaskar, Dept. of Political Science, Govt. City (A) College, Nayapool, Hyderabad.
 - 10. Dr. Hussain, Dept-of Political Science, Govt. Degree College, Gajwel.
- 11. Dr. S. Sharwani, Dept. of Political Science, Govt. Degree College, Falaknuma, Hyderabad.
- 12. Dr. K. Prabhu, Dept. of Political Science, Govt. Degree College, Chevella.

The term of the Office of the Board of Studies in Political Science (UG) shall be for a period of Two (2) years with effect from the date of issue of this notification. However, any Member of the Board shall cease to be a Member when he/she vacates the Office when he/she was holding at the time of his/her nomination, unless otherwise permitted by the Vice-Chancellor.

The Membership is purely honorary. The TA and DA will be paid to the External Members as per the rules of the University in force whenever they attend the meeting of the Board of Studies.

Girraj Govt. College (A), Nizamabad

Re-Accredited by NAAC with "B"

Date: 06-11-2021

From:

The Principal, Girraj Govt. College(A), Nizamabad. To:

Dr. A. Venkatesham Head, Dept. of Economics Tara Govt. College (Autonomous), Sangareddy

sir,

Sub: GG College (A), Nzb- BOS Meeting in Dept. of Economics on 9-11-2021-Request to attend-Reg.

Ref: Proc. File No-GDC/NZB-GEN/261/2021-O/o PRINCIPAL-GDC-NZB-CE Dt: 01-11-2021

With the reference subject cited above, the BOS Meeting of Economics Department is to be held on 09-11-2021 at 3.00 PM in the Department of Economics in the College with the following Agenda.

- 1. Finalize and approval of Final Year V and VI Semester (UG) Syllabus, Question paper pattern etc., of CBCS common core syllabus as proposed by TSCHE.
- 2. Review on Results of I, III, V and II, IV, VI Semester of UG in the academic year 2020-21
- 3. Review of academic activities of the Dept for the academic year 2021-22.
- 4. Any other item with the permission of the chair.

Hence, I request you to make it convenient to attend the BOS meeting. Your cooperation is highly solicited.

Thanking you

PRINCIPAL

Encloses

1. Copy of BOS Members in Economics

Signed by Dr.e Laxmi Narayana Date: 08-11-2021 18:37:49 Reason: Approved

File No.GDCNZB-GEN/261/2021-O/o PRINCIPAL-GDC-NZB-CE

Proceedings of the Principal (FAC), Girraj Govt. College (A), Nizamabad.

(Re-Accredited by NAAC with "B")

Present: Dr. E. Laxminarayana, M.A, Ph.D Principal (FAC)

Sub: Reconstitution of Board of Studies Members for Dept. of Economics for the tenure of Three Years from the academic year 2021-22 – Orders – Issued.

ORDER:

Sanction is accorded for reconstitution of Board of Studies Members for Dept. of Economics with the following members for the tenure of Three Years from the academic year 2021-22.

1. Dr. Y. Venu Prasad	Chairperson, Board of Studies
Head, Dept of Economics, GG College (A), Nzb	
2. Dr. P. Nagaraj	University Nominee
Associate Professor of Economics,	•
BOS Chairperson, Telangana University. Nzb.	
3. Dr.K. Krishna Reddy	Subject Expert
Head, Dept. of Economics	
Dr. B. R Ambedkar Open University, Telangana.	
4. Dr. A. Venkatesham	Subject Expert
Head, Dept. of Economics	
Tara Govt. College (Autonomous), Sangareddy	
Osmania University, Hyd.	
5. Sri Bachu Venkatesh	Expert from Industry
Kamadhenu Food Processing Industries	
Rudrur, Nizamabad,T.S503188	
6. Dr. V. Mutyam Reddy	Member
Asst.Prof of Eco, GG College (A),Nzb	
7. Dr. D. Adeppa	Member
Asst.Prof of Eco, GG College (A),Nzb	
8. Dr. G. Sriram	Member
Asst.Prof of Eco, GG College (A),Nzb	
9. Smt. Tabassum Shaeen.	Member
Asst.Prof of Eco, GG College (A),Nzb	
10. Kum. B. Bhargavi.	P.G Meritorious Student Member
BA EPP (2017-20)	
MA (Economics), Arts College, OU, Hyd.	
	DRINCIDAL

PRINCIPAL

Copy submi	tted to:	Signed by Dr.e Laxmi
1.	The Registrar, TU, Nizamabad.	Narayana
Copy to:		Date: 01-11-2021 17:15:23
2.	The Coordinator Autonomy & UGC, GGC (A) Nizamabad.	Reason: Approved
3.	The Controller of Examinations, GGC (A), Nizamabad.	
4.	All the members of BOS.	





CERTIFICATE OF COLLABORATION

between Department of Chemistry, Girraj Govt. College(A), Nizamabad & Department of Chemistry, TARA Govt. College, Sangareddy(A)

Department of Chemistry, Girraj Govt. College(A), Nizamabad and **Department of Chemistry, TARA Govt. College, Sangareddy(A)** agree on engaging in collaboration for quality enhancement for the mutual progression of both the institutions. The subject to mutual consent, the area of cooperation will include the following aspects;

- 1. Participation in seminars, workshops and academic meetings.
- 2. Extension Lectures and Training.
- 3. Joint Research activities and publications.
- 4. Designing and development of teaching and learning modules.
- 5. Certificate programs and projects for skill development.
- 6. Quality enhancement initiatives.

TERMS OF IMPLEMENTATION:

1. Details of the implementation of any particular exchange resulting from this Collaboration shall be negotiated between the two organizations.

2. This Collaboration becomes effective on the day it is signed and remains valid for three years.

- 3. This Collaboration will be renewed after three years upon the consent of both organizations.
- 4. Any amendment or modification to the present text shall be submitted for review to the competent

authorities, and shall not binding unless reduced to writing and signed by both the organizations.

5. This Collaboration does not bind either of the two parties legally or financially. Its aim is to promote relations that will mutually benefit each organization, this being the primary aim of academic collaboration.

In witness whereof, the organizations hereto have offered their signatures.

Head of Department, Department of Chemistry Girraj Govt. College(A), Nizamabd Date: 1/2/21 Principa Girraj Govt. College(A), Nizamabd Date: 114/2021 FRINCIPA

Head of Department, Department of Chemistry Tara Govt. College, Sangareddy(A) Date: 01.04.21.

Principal, Tara Govt. College, Sangareddy(A) Date: <u>0 | 0 4 2</u>. **PRINCIPAL TARA GOVT. COLLEGE** (AUTONOMOUS) **SANGAREDDY**-502 001





Memorandum of Understanding (MoU) between MART Specialities Lab LLP, Hyderabad

&

Department of Chemistry, TARA Govt. College, Sangareddy (Autonomous)

MART Specialities Lab LLP, Hyderabad and Department of Chemistry, TARA Govt. College,

Sangareddy(A) agree on the importance and the usefulness of establishing scientific and academic links, in order to assert and to consolidate the ties of friendship between two institutions. The subject to mutual consent, the area of cooperation will include the following aspects;

1. Design and developments of Student research projects

- 2. Participation in Research and academic meetings.
- 2. Student Training Programmes.
- 3. Joint Research activities and publications.
- 4. Providing Analytical services to the research projects.
- 5. Support in Patent Filing procedures.
- 6. Quality enhancement initiatives.

TERMS OF IMPLEMENTATION:

1. Details of the implementation of any particular exchange resulting from this MoU shall be negotiated between the two organizations.

2. This MoU becomes effective on the day it is signed and remains valid for THREE years.

3. This MoU will be renewed after THREE years upon the consent of both organizations.

4. Any amendment or modification to the present text shall be submitted for review to the competent

authorities, and shall not binding unless reduced to writing and signed by both the organizations. 5. This MoU does not bind either of the two parties legally or financially. Its aim is to promote relations that will mutually benefit each organization, this being the primary aim of academic collaboration.

In witness whereof, the organizations hereto have offered their signatures.

X.7-j--

Managing Partner MART Specialities Lab LLP, Hyderabad, Hyderabad.

Date: 31.12.2021

Head of Department, Department of Chemistry Tara Govt. College, Sangareddy(A) Date: <u>3</u>].12 · 2].

Vuced

PRINCIPAL TARA GOVT. COLLEGE, SANGAREDDY(A) Date: 31.12.2

PRINCIPAL TARA GOVT. COLLEGE AUTONOMOUS SANGAREDDY-502001



PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education– On Duty to faculty from GDCs –TSBIE-ERTW-II(TB)- **Revision of Intermediate II year Telugu Syllabus** - Subject Committee Meeting– Orders Issued-Reg.
- Ref: Letter received from The Secretary, Telangana State Board of Intermediate Education, dated: 28.07.2021.

Vide reference read above, the Commissioner of Collegiate Education has accorded permission to the following faculty members to attend the meeting for Revision of syllabi of Telugu textbook for Intermediate II year as **Subject Committee members** at **Telangana State Board of Intermediate Education, Nampally**, Hyderabad on **29.07.2021.**

SI.No.	Name and Designation	Subject	Contact No. & Email.I.D	Date of Meeting
1.	Dr. J.Neeraja Assistant Professor, Govt. City College (A), Nayapul,Hyderabad.	Telugu	9848370018 nj.ccets@gmail.com	
2.	Dr. K.Koteshwara Rao Assistant Professor, Govt. City College (A), Nayapul, Hyderabad.	Telugu	9440480274 koyikoti@gmail.com officegdcw.nlg@gmail.c om	29.07.2021
3.	Sri. B.Venkateshwarlu, Assistant Professor, Govt.Degree College, Husnabad, Siddipet	Telugu	9491598040 Venkateshwarluboorla 500@gmail.com.	
4.	Dr.P.Bhaskara Yogi, Assistant Professor, Govt.Degree College Sangareddy	Telugu	9912070125 Bhaskarayogi.p@gmail. com	

The absence of the above said faculty members in the college shall be treated as **On Duty** on the days they attend the meeting and they are not eligible to claim TA/DA from the colleges. In view of the above, the concerned Principals are informed to relieve them to attend the above said meeting on the above said dates.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by Ghanshyam Date: 2021.07.28 13:18:00 IST Reason: Approved

For Commissioner of Collegiate Education

Copy To Secretary, TSBIE, Hyderabad Principal, Govt. City College, Nayapul Principal, GDC, Husnabad Principal, GDC, Sangareddy

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education– On Duty to faculty from GDCs –TSBIE-ERTW-II(TB)- **Revision of Intermediate II year Telugu Syllabus** - Subject Committee Meeting– Orders Issued-Reg.
- Ref: Letter received from The Secretary, Telangana State Board of Intermediate Education, dated: 20.07.2021.

Vide reference read above, the Commissioner of Collegiate Education has accorded permission to the following faculty members to attend the meeting for Revision of syllabi of Telugu textbook for Intermediate II year as **Subject Committee members** at **Telangana State Board of Intermediate Education, Nampally**, Hyderabad on **22.07.2021**, **23.07.2021** and **24.07.2021**.

SI.No.	Name and Designation	Subject	Contact No. & Email.I.D	Date of Meeting
1.	Dr. J.Neeraja Assistant Professor, Govt. City College (A), Nayapul,Hyderabad.	Telugu	9848370018 nj.ccets@gmail.com	
2.	Dr. K.Koteshwara Rao Assistant Professor, Govt. City College (A), Nayapul, Hyderabad.	Telugu	9440480274 koyikoti@gmail.com officegdcw.nlg@gmail.c om	22.07.2021 23.07.2021
3.	Sri. B.Venkateshwarlu, Assistant Professor, Govt.Degree College, Husnabad, Siddipet	Telugu	9491598040 Venkateshwarluboorla 500@gmail.com.	24.07.2021
4.	Dr.P.Bhaskara Yogi, Assistant Professor, Govt.Degree College Sangareddy	Telugu	9912070125 Bhaskarayogi.p@gmail. com	

The absence of the above said faculty members in the college shall be treated as **On Duty** on the days they attend the meeting and they are not eligible to claim TA/DA from the colleges. In view of the above, the concerned Principals are informed to relieve them to attend the above said meeting on the above said dates.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by Ghans yam Date: 2021.07.20 16:47,45 IST Reason: Approved

For Commissioner of Collegiate Education

Copy To Secretary, TSBIE, Hyderabad Principal, Govt. City College, Nayapul Principal, GDC, Husnabad Principal, GDC, Sangareddy

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education– On Duty to faculty from GDCs –TSBIE-ERTW-II(TB)- **Revision of Intermediate II year Telugu Syllabus** - Subject Committee Meeting– Orders Issued-Reg.
- Ref: Letter received from The Secretary, Telangana State Board of Intermediate Education, dated: 16.07.2021.

Vide reference read above, the Commissioner of Collegiate Education has accorded permission to the following faculty members to attend the meeting for Revision of syllabi of Telugu textbook for Intermediate II year as **Subject Committee members** at **Telangana State Board of Intermediate Education, Nampally**, Hyderabad on **17.07.2021**.

SI.No.	Name and Designation	Subject	Contact No. & Email.I.D	Date of Meeting
1.	Dr. J.Neeraja Assistant Professor, Govt. City College (A), Nayapul,Hyderabad.	Telugu	9848370018 nj.ccets@gmail.com	
2.	Dr. K.Koteshwara Rao Govt.Degree College for Women, Nalgonda.	Telugu	9440480274 koyikoti@gmail.com officegdcw.nlg@gmail.c om	17.07.2021
3.	Sri. B.Venkateshwarlu Govt.Degree College, Husnabad, Siddipet	Telugu	9491598040 Venkateshwarluboorla 500@gmail.com.	
4.	Dr.P.Bhaskara Yogi Govt.Degree College Sangareddy	Telugu	9912070125 Bhaskarayogi.p@gmail. com	

The absence of the above said faculty members in the college shall be treated as **On Duty** on the days they attend the meeting and they are not eligible to claim TA/DA from the colleges. In view of the above, the concerned Principals are informed to relieve them to attend the above said meeting on the above said dates.

(Orders of the CCE have been obtained in the note file)

Signature Not Ve	erified
Digitally signed by G Date: 2021.07.16 16 Reason: Approved	hanshyam : <mark>31:1</mark> 0 IST

For Commissioner of Collegiate Education

Copy To Secretary, TSBIE, Hyderabad Principal, Govt. City College, Nayapul Principal, GDC W, Nalgonda Principal, GDC, Husnabad Principal, GDC, Sangareddy

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education– On Duty to faculty from GDCs –TSBIE-ERTW-II(TB)- **Revision of Intermediate II year Telugu Syllabus** - Subject Committee Meeting– Orders Issued-Reg.
- Ref: Letter received from The Secretary, Telangana State Board of Intermediate Education, dated: 12.07.2021.

Vide reference read above, the Commissioner of Collegiate Education has accorded permission to the following faculty members to attend the meeting for Revision of syllabi of Telugu textbook for Intermediate II year as **Subject Committee members** at **Telangana State Board of Intermediate Education, Nampally**, Hyderabad on **13.07.2021**.

SI.No.	Name and Designation	Subject	Contact No. & Email.I.D	Date of Meeting
1.	Dr. J.Neeraja Assistant Professor, Govt. City College (A), Nayapul,Hyderabad.	Telugu	9848370018 nj.ccets@gmail.com	
2.	Dr. K.Koteshwara Rao Govt.Degree College for Women, Nalgonda.	Telugu	9440480274 koyikoti@gmail.com officegdcw.nlg@gmail.c om	13.07.2021
3.	Sri. B.Venkateshwarlu Govt.Degree College, Husnabad, Siddipet	Telugu	9491598040 Venkateshwarluboorla 500@gmail.com.	
4.	Dr.P.Bhaskara Yogi Govt.Degree College Sangareddy	Telugu	9912070125 Bhaskarayogi.p@gmail. com	

The absence of the above said faculty members in the college shall be treated as **On Duty** on the days they attend the meeting and they are not eligible to claim TA/DA from the colleges. In view of the above, the concerned Principals are informed to relieve them to attend the above said meeting on the above said dates.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by Ghanehyam Date: 2021.07.12 14:16:16 IST Reason: Approved

For Commissioner of Collegiate Education

Copy To Secretary, TSBIE, Hyderabad Principal, Govt. City College, Nayapul Principal, GDC W, Nalgonda Principal, GDC, Husnabad Principal, GDC, Sangareddy

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education– On Duty to faculty from GDCs –TSBIE-ERTW-II(TB)- **Revision of Intermediate II year Telugu Syllabus** - Subject Committee Meeting– Orders Issued-Reg.
- Ref: Letter received from The Secretary, Telangana State Board of Intermediate Education, dated: 07.07.2021.

Vide reference read above, the Commissioner of Collegiate Education has accorded permission to the following faculty members to attend the meeting for Revision of syllabi of Telugu textbook for Intermediate II year as **Subject Committee members** at **Telangana State Board of Intermediate Education, Nampally**, Hyderabad on **09.07.2021**.

SI.No.	Name and Designation	Subject	Contact No. & Email.I.D	Date of Meeting
1.	Dr. J.Neeraja Assistant Professor, Govt. City College (A), Nayapul,Hyderabad.	Telugu	9848370018 nj.ccets@gmail.com	
2.	Dr. K.Koteshwara Rao Govt.Degree College for Women, Nalgonda.	Telugu	9440480274 koyikoti@gmail.com officegdcw.nlg@gmail.c om	09.07.2021
3.	Sri. B.Venkateshwarlu Govt.Degree College, Husnabad, Siddipet	Telugu	9491598040 Venkateshwarluboorla 500@gmail.com.	
4.	Dr.P.Bhaskara Yogi Govt.Degree College Sangareddy	Telugu	9912070125 Bhaskarayogi.p@gmail. com	

The absence of the above said faculty members in the college shall be treated as **On Duty** on the days they attend the meeting and they are not eligible to claim TA/DA from the colleges. In view of the above, the concerned Principals are informed to relieve them to attend the above said meeting on the above said dates.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by Ghanstyam Date: 2021.07.08 13:48/32 IST Reason: Approved

For Commissioner of Collegiate Education

Copy To Secretary, TSBIE, Hyderabad Principal, Govt. City College, Nayapul Principal, GDC W, Nalgonda Principal, GDC, Husnabad Principal, GDC, Sangareddy

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education– On Duty to faculty from GDCs –TSBIE-ERTW-II(TB)- **Revision of Intermediate II year Telugu Syllabus** - Subject Committee Meeting– Orders Issued-Reg.
- Ref: Letter received from The Secretary, Telangana State Board of Intermediate Education, dated: 01.07.2021.

Vide reference read above; the Commissioner of Collegiate Education has accorded permission to the following faculty members to attend the meeting for Revision of syllabi of Telugu textbook for Intermediate II year as **Subject Committee members** at **Telangana State Board of Intermediate Education, Nampally**, Hyderabad on **06.07.2021**.

SI.No.	Name and Designation	Subject	Contact No. & Email.I.D	Date of Meeting
1.	Dr. J.Neeraja Assistant Professor, Govt. City College (A), Nayapul,Hyderabad.	Telugu	9848370018 nj.ccets@gmail.com	
2.	Dr. K.Koteshwara Rao Govt.Degree College for Women, Nalgonda.	Telugu	9440480274 koyikoti@gmail.com officegdcw.nlg@gmail.c om	06.07.2021
3.	Sri. B.Venkateshwarlu Govt.Degree College, Husnabad, Siddipet	Telugu	9491598040 Venkateshwarluboorla 500@gmail.com.	
4.	Dr.P.Bhaskara Yogi Govt.Degree College Sangareddy	Telugu	9912070125 Bhaskarayogi.p@gmail. com	

The absence of the above said faculty members in the college shall be treated as **On Duty** on the days they attend the meeting and they are not eligible to claim TA/DA from the colleges. In view of the above, the concerned Principals are informed to relieve them to attend the above said meeting on the above said dates.

(Orders of the CCE have been obtained in the note file)

Signature Not Verified Digitally signed by Ghanshyam Date: 2021.07.01 18:41:03 IST Reason: Approved

For Commissioner of Collegiate Education

Copy To Principal, Govt. City College, Nayapul Principal, GDC W, Nalgonda Principal, GDC, Husnabad Principal, GDC, Sangareddy

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD PRESENT: SRI NAVIN MITTAL, I.A.S.

- Sub: Collegiate Education Government Degree College, Zaheerabad, Sangareddy Dist. – Deputation of faculty members on On Duty basis for assisting in NAAC Peer Team visit – 29.11.2021 to 02.12.2021- On Duty Orders issued - Reg.
- Ref: Letter received from Principal, Government Degree College, Zaheerabad, Sangareddy Dist. Dated: 27.11.2021.

In pursuance of the circumstances informed in the reference read above, the Commissioner of Collegiate Education, Hyderabad, hereby accorded permission to the following faculty members from various GDCs to work in Government Degree College, Zaheerabad, Sangareddy Dist. from 29.11.2021 to 02.12.2021 for providing assistance in NAAC related work.

SI. No.	Name of the Faculty member	Subject	Name of the College
1	Md. Jaleel	English	GDC A, Sangareddy
2	Dr. L. Raji Reddy	Commerce	GDC Patancheru

In view of the above, the concerned Principals are instructed to relieve the above mentioned faculty members on the above mentioned dates and their absence in the college on the dates mentioned above may be treated as On Duty. The faculty members are not eligible to claim TA/DA from the college.

(Orders of the CCE have been obtained in the note file)

Signed by Ghanshyam Date: 27-11-2021 16:52:58 Reason: Approved For Commissioner of Collegiate Education Copy To Principal, GDC Sangareddy Principal, GDC Patancheru Principal, GDC Zaheerabad



Tara Government College Sangareddy (Autonomous) An ISO 9001: 2015 certified college **Department of Botany**



TRAINING ON SOIL TESTING

The MoU has been done between the department of Botany and Soil Testing Lab, Sangareddy, for two years where Smt. Haritha Madam, Asistant Director of Agriculture has signed the agreement and agreed on the importance and the usefulness of establishing scientific and academic links.

The objective of this MoU is to train the students and to fecilitate the testing of the samples collected randomly by the students. Nearly 50-60 students have been given the training in the month of December- 2021 in two batches. The analysis was done for the samples collected by the students and the report was given by the lab authorites. Based on the report the levels of Macro and Micro nutrients have been identified. The students visited the farmers and recommended the use of Bioferilisers prepared by the Department.

A certificate course has been introduced in soil testing, the classes were taken by the faculty in the Botany Dept. and an examination has also been conducted and the certificate has been given to the students.



Memorandum of Understanding (MoU) Between DEPT OF BOTANY, TARA GOVERNMENT COLLEGE (A), SANGAREDY & SOIL TESTING LAB, SANGAREDDY



Soil Testing Lab, Sangareddy and Department of Botany, Tara Government College, Sangaredy agree on the importance and the usefulness of establishing scientific and academic links, in order to assert and to consolidate the ties of friendship hetween two departments. The subject to mutual consent, the area of cooperation will include the following aspects;

- 1. The objective of this MoU is to facilitate the testing of samples collected by the students for macro and micro nutrient analysis.
- 2. Participation in seminars and workshops.
- 3. Extension Lectures and Training.
- 4. Certificate programs and projects for skill development.
- Coordination in creating awareness among the farmers for the significance f soil testing to have a better yield.
- 6. Quality enhancement initiatives.

TERMS OF IMPLEMENTATION:

1. Details of the implementation of any particular exchange resulting from this MoU shall be negotiated between the two organizations.

2. This MoU becomes effective on the day it is signed and remains valid for two years.

3. This MoU will be renewed after two years upon the consent of both organizations.

4. Any amendment or modification to the present text shall be submitted for review to the competent authorities, and shall not binding unless reduced to writing and signed by both the organizations.

5. This MoU does not bind either of the two parties legally or financially. Its aim is to promote relations that will mutually benefit each organization, this being the primary aim of academic collaboration.

In witness where of, the organizations here to have offered their signatures.

PRINFINALPAL iovi College LEGE **HAMOUS** 02001

Head of Advantment, Dopathonah v Bolan Xy Tara Government, Collano (A) Sangareddy Sez con Telano

Asst Director of Agriculture Soil Testing Labor of Agriculture Soil Testing Laboratory Sauthedd SANGAREDDY-502 001



Asst.Director-Ms. Haritha Madam

Training the Students



Training the Students



Soil Testing Lab, Sangareddy



Principal Ms. M.Praveena Madam addressing the Students



Certificates given to the students



		PG.NO
S.NO	CONTENTS	
1	INTRODUCTION SOIL HEALTH MANAGEMENT IMPORTANCE AND OBJECTIVES	
2	IMPORTANCE OF SOIL TESTING	
3	SOIL SAMPLING METHODS AND PROCEDURES	
4	SOIL PARAMETERES	
5	DETERMINATION OF SOIL COLOUR AND TENTURE BY #112, D-METHOD FEEL	
0	DETERMINATION OF SOIL REACTION	
7	DETERMINATION OF ELECTONIC CONDUCTIVITY	
s	DETERMINATION OF ORGANIC CARBON	
9	DETERMINATION OF AVAILABLE NITROGEN	
10	DETERMINATION OF AVAILABLE PHOSPHORUS	
11	DETERMINATION OF AVAILABLE POTASSIUM	
12	DETERMINATION OF BORON	
13	DETERMINATION OF SULPHUR	
14	DETERMINATION OF MICRO NUTRIENTS	
15	INTERPRETATION OF SOIL TEST $^{\underline{S}HH}_{A}$: Ranges & Reclamations	

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TARA GOVERNMENT COLLEGE (A), SANGAREDDY

SOIL TESTING-EXAMINATION

25 MARKS-1hr

1. Write about the soil sampling and also about the steps in soil testing? 5 marks

Short Answers= 2marks each

1

2. Define labeling?

3. At what time you have to collect the soil samples?

4. Write a short note on soil texture?

5. What is the pH for acidic and basic soils?

6. State the principle of estimation of Gypsum required for alkali soil?

7. Write a short note on a wet digestion method?

8. What is the major nutrient deficiencies in soil and what are the symptoms seen on plants?

9. What is the percentage of Nitrogen in the atmosphere? How does the fixation takes place?

10. Define the soil? VI. Doms a diagram of Soil pufile

4

TARA GOVERNMENT COLLEGE, SANGAREDDY

(ISO 9001:2015 certified) (District Identified College & District Resource Centre) (Affiliated to Osmania University & Accredited by NAAC with 'B' grade, 2.75 CGPA) SANGAREDDY (Dist.), Telangana.



E-mail: prl-gdc-srd-ce@telangana.gov.in

Website: https://gdcts.cgg.gov.in

TASK DATA

TASK-HYSEA	Dear Sir/Madam,	14.09.2022
WINTalk series		
on AI/ML		
	Creatings for the day!	
	Greetings for the day!	
	This mail is regarding WINtalk Series session on AI/ML	
	which has been rescheduled on 16th September 2022 from	
	3:00 PM to 4:00 PM.	
	The panelists for the session are Mr. Srinivas Atreya, Chief	
	Data Scientist, Round Sqr, Mr. Kishore Seshagiri, EVP,	
	Brodridge Financial services.	
	Moderator for the session Mr.Shrikant Sinha, CEO-TASK.	
	Joining link for the session:	
	https://task.radiusedutech.com/	
	Attached is the flyer for the WINtalk Series session	

	Request you to share with faculty and Students and encourage them to make use of this opportunity.	
	Dear Sir/Madam,	13.09.2022
TASK -		
INNOHAT		
System -	TASK in association with InnoHat System is conducting an	
Training on RPA	Instructor-led virtual Training Program on Robotic process automation (RPA). This program is offered to all TASK registered students and Faculty. Below are the complete details and attached is the EDM.	
	Training Program:-RPA	
	No of Days:-5 Days	
	Training Start Date:19th Sep 2022	
	Training End Date: 23rd Sep 2022	
	Timings: 6.00 PM to 7.00 PM	
	Note: Students/Faculty who attend all five days will get the participation certificate from the InnoHat system.	
	Registration link: - https://forms.gle/J8XoHtvRgyLK64vp9	
	Faculty and students from Engineering/Degree can attend this training program.	
	Last Date to register: 16th Sept 22.	

ASK - College Registration/Ren ewal	Dear Sir/Madam,	13.09.2022
Notification for the Academic Year 2022-23	Greetings from TASK!	
	Telangana Academy for Skill and Knowledge (TASK) is established under Department of ITE & C, Government of Telangana to enhance the employability quotient of youth in Telangana state.	
	TASK releases a notification for College Registration/Renewal for the Academic Year 2022-23. I would like to request you to please complete the registration process online.	
	Website: https://task.telangana.gov.in/	
	The last date is on 30th September 2022, for detailed notification please find the attached document.	
	Below are the important dates:	
	College Registration/Renewal	
	Start Date & Time	
	End Date & Time	
	Without late Fee	

	01–09–2022	
	10 AM onwards	
	30-09-2022	
	till 11.59 PM	
	With a late fee of Rs.1000/- + Applicable Taxes	
	01–10–2022	
	10 AM onwards	
	10-10-2022	
	till 11.59 PM	
TASK - HYSEA Drona 12 weeks online faculty	Dear Sir/Madam,	29.08.2022
immersive learning program	Greetings from TASK!	
	TASK – HYSEA would like to conduct a 12 weeks online faculty immersive learning program for IT industry needs.	
	Kindly nominate your faculty in below link on or before 5th September 2022	
	https://docs.google.com/forms/d/e/1FAIpQLScBUvs7m- 23qmsYMxPJefoqtvayaUCG3S3REBRblBNAJPpBVA/viewf orm	
	Program Design:	

3 emerging technology tracks (Full stack, AI/ML/ Digital transformation)
Self-learning on Infosys Springboard and help at frequent intervals by Mentors via masterclass.
Experiential learning via capstone & presentation that will be evaluated for understanding of the code & business sense.
One / Two days Faculty immersive experience in corporate to understand project management process and various roles & responsibilities.
Learning Objectives
By the end of the program faculty will be able to:
1. Analyse various emerging technologies
2. Apply the concepts to solve problems
3. Understand the program management process
4. Demonstrate understanding of various roles in IT industry
What's in it for the Faculty?
Get hands on with what Industry needs

Emerging Technology – hands on ability to code
Business Sense & Customer Centricity
Program Management with focus on Professional Skills Development
Exposure to nuances of IT industry
What is the program execution plan?
Faculty enrols into a technology track. Learn the concepts in an online mode. A pool of faculty for each track will be available at stipulated hours for doubt clarification. Applies the concepts in a capstone program. The faculty makes a presentation of the capstone. Industry emerging technology practicing experts evaluate the capstone & offer feedback.
Certification – 4 weeks each track – 12 weeks program
Completion of 1 track – Eligible for 1 track course completion certificate
Completion of 2 tracks – Eligible for faculty immersive experience + 2 track course completion certificate
Completion of 3 tracks – Award Program Completion certificate
Eligibility:
Faculty in teaching Computer Science, IT or related subjects in Engineering, BCA, MCA, Degree, and Polytechnic institutions.

	Proposed Date	
	Program launch – 2:00 PM to 3:30PM, 5th Sept 22	
	Please reach out to me for any further clarifications.	
	Dear Sir/Madam,	16.08.2022
TASK-		
Quantum		
Science and	Greetings of the Day!	
Technology		
Hackathon 2022		
	TASK in association Quantum Ecosystems and Technology Council of India is organising Quantum Science and Technology Hackathon 2022.	
	The Real Value of Quantum Computers lies in the problems that they can solve for Humanity, which are either difficult or impossible to solve purely with Classical Computers.	
	We are announcing the Quantum Science and Technology Hackathon 2022 !	
	Registration and Idea Submission starts : Registration closes on 15th September 6:29 pm and the registered participants continue Hacking till end of October.	
	Eligibilty :	
	To tackle challenges in applying quantum technologies, we need not only people with skills in computer science,	

physics, and math, but also people with expertise in the application domains of this hackathon: financial services, life sciences, quantum security, and quantum sensors/communications, and even applications you can propose yourself!
Anybody starting students studying for undergraduate degrees
The detailed Schedule is attached.
Link to Register Online Hackathon HackerEarth developer event Quantum Science and Technology Hackathon 2022
Through this Hackathon, we want to :
· Enable hackers to come up with solutions using a Quantum Stack/Technology of their choice
 Collaborate across Geographies and get access to mentors from around the world to help them solve the problems. Provide access to Incubation and mentorship support for people who want to create a startup from their project
• Provide an opportunity for students and practitioners to learn and be part of a larger community, providing them access to Leading Researchers and experts who can help them with their careers.
• Some of the Partners and Sponsor Companies will offer Internships and other job opportunities to hackers who they find suitable for their work. Completion of a project in the hackathon will go a long way in building confidence in the Industry.
 Pick up some of the winning projects to be part of the Open Source Projects that we want the Ecosystem to focus

 on.	
The Hackathon themes are around Financial Services, Life Sciences (Pharmaceutical, Healthcare, Biosciences, and	
Biotech), Quantum Security, Quantum sensors, and Quantum Communications. Apart from this, there is a Theme Others under which people can hack on any problem using and leveraging Quantum Science, technology and concepts.	
There are some really exciting PRIZES	
- Cash Prizes totaling to Rs 10 Lacs	
- Winners in each Category – Top 3 get access to demo day with investors and incubators	
- All winners get Winner Certificates and QETCI Membership opportunities	
- Winning Projects get Credits for access to quantum hardware	
- Two Projects will get an opportunity to present their projects at the Quantum. Tech APAC Conference in Singapore	
- Project teams that want to register their startups, will get support from empanelled Companies in the registration process, as well as any patent filing process.	
So what are you waiting for - Start Hacking	
Online Hackathon HackerEarth developer event	

	Quantum Science and Technology Herberther 2002	
	Quantum Science and Technology Hackathon 2022	
	I request you to share with the students and encourage	
	them to make use of this opportunity.	
	Dear Sir/Madam,	11.08.2022
TASK-Celonis		
Academic		
Alliance	Greetings from TASK!	
registration for		
FDP	Thank you for Joining the awareness session . I am sharing more details about the program and the next steps for faculty/institution.	
	Celonis is the worldwide leader in Process Mining and Execution Management that empowers people to analyse any gap while creating, building and sharing their innovations. Our technology is changing the nature of work and the way we live. Our solutions deliver a brave new model of technology that is fluid, instant and more secure. Customers can innovate faster by rapidly developing, automatically delivering and more safely consuming any application.	
	As part of efforts to empower academia, want to discuss on Celonis Academic Alliance Program and work jointly on the following:	
	• Offer a high quality curriculum that combines in-depth technical training and professional skills development.	
	• Choose courses to complement your academic program and prepare students for certification.	
	• Leverage our learning platform to track student	

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	success and connect with educators and Process Mining experts.	
	• Local student chapter including annual project competitions, workforce development programs and others.	
	• Point students to career building resources created with employers for our alumni.	
	We are planning to conduct a FDP for the colleges. Please share this information with all colleges and and ask them to nominate the faculty using the below link by 12th August 2022.	
	https://forms.gle/w1M2kiYUNUD4mV2d8	
	Details regarding the session will be shared shortly.	
	For more information sharing the program URL for your reference: https://www.celonis.com/academic-alliance/ and attached are the PDF's for your reference.	
	Kindly reach out to me in case of any questions.	
TASK - Series of Speed Mentoring	Dear Sir/Madam,	11.08.2022
Sessions by Amazon	Greetings from TASK	

Telangana Academy for Skill and Knowledge(TASK) in collaboration with Mentor Together and Amazon is inviting your students to be part of the mentoring series on placement readiness.	
Inform all your students to join this mentorship series to get the chance to interact with experts from Amazon who will help them to create a winning CV, prepare for an interview, and build their professional brand on LinkedIn.	
Here is a google form and students will receive mentoring session invitations directly to their email.	
Mentoring series details:	
Session 01: Preparing a winning CV on 18 Aug 2022, at 4.15 pm to 5.45 pm	
Session 02: Building a professional brand on LinkedIn on / 25 Aug 2022, at 4.15 pm to 5.45 pm	
Session 03: Preparing for an interview on / 30 Aug 2022, at 4.15 pm to 5.45 pm	
Session 04: Preparing for an interview on / 05 Sep 2022, at 4.15 pm to 5.45 pm	
Session 05: Preparing for an interview on / 06 Sep 2022, at 4.15 pm to 5.45 pm	
Looking forward to having your students in TASK - MT mentoring series!!	

	Register now to interact with industry professionals https://forms.gle/AMZCdyXNvgkfZU72A	
	Get updates on your WhatsApp by joining this group https://chat.whatsapp.com/ErEguQjhfMkCz00UZoGPIH	
TASK - HYSEA WINtalk series on 'UI/UX' on 5th August 2022	Dear Sir/Madam,	04.08.2022
	Greetings from TASK!	
	TASK - WINtalk series session on 'UI/UX' on 5th August 2022 from 3:00 pm to 4:00 pm.	
	The panelists for the session are Mr.Ameya Naik,Sr.Manager Product Design,ServiceNow and Mr.Prasad Kantamneni,Founding Partner,UXReactor.	
	Moderator for the session Mr. R. Srinivas Rao, Chief Operating Officer, HYSEA.	
	Joining link for the session: https://task.radiusedutech.com/	
	Attached is the flyer for WINtalk series session.	
	Request you to share with all faculty and students and encourage them to make use of this opportunity.	

TASK-Celonis Academic Alliance	Dear Sir/Madam,	27.07.2022
Partnership - Online Meeting on 1st August 22	TASK have planned Celonis Academic Alliance Partnership - Meeting on 1st August 22.	
	Joining Link: https://celonis.zoom.us/j/98920600021?pwd=SlpsNXIxb2d1S 2hZQWNjVWw5RFZJZz09#success	
	ID: 98920600021 passcode: j75S3EkZ	
	Timings: 03:00 PM to 04:00 PM	
	Kindly join the meeting on time.	
	Partnership Details:	
	TASK in association with" Celonis Academic Alliance" is giving opportunity to all colleges to become a member of " Celonis Academic Alliance Program" to avail opportunities designed exclusively for Institutions, Faculties and their Students.	
	About Celonis Academic Alliance Program:	

Celonis is the worldwide leader in Process Mining and Execution Management that empowers people to analyze any gap while creating, building and sharing their innovations. Our technology is changing the nature of work and the way we live. Our solutions deliver a brave new model of technology that is fluid, instant and more secure. Customers can innovate faster by rapidly developing, automatically delivering and more safely consuming any application.

Celonis collaborates with TASK as a Regional Academic Alliance Training Partner to enhance students and educators on Process Mining and EMS tool and make them ready to deploy resources at no cost.

As a part of efforts to empower academia, TASK and Celonis Academic Alliance Program work jointly on the following:

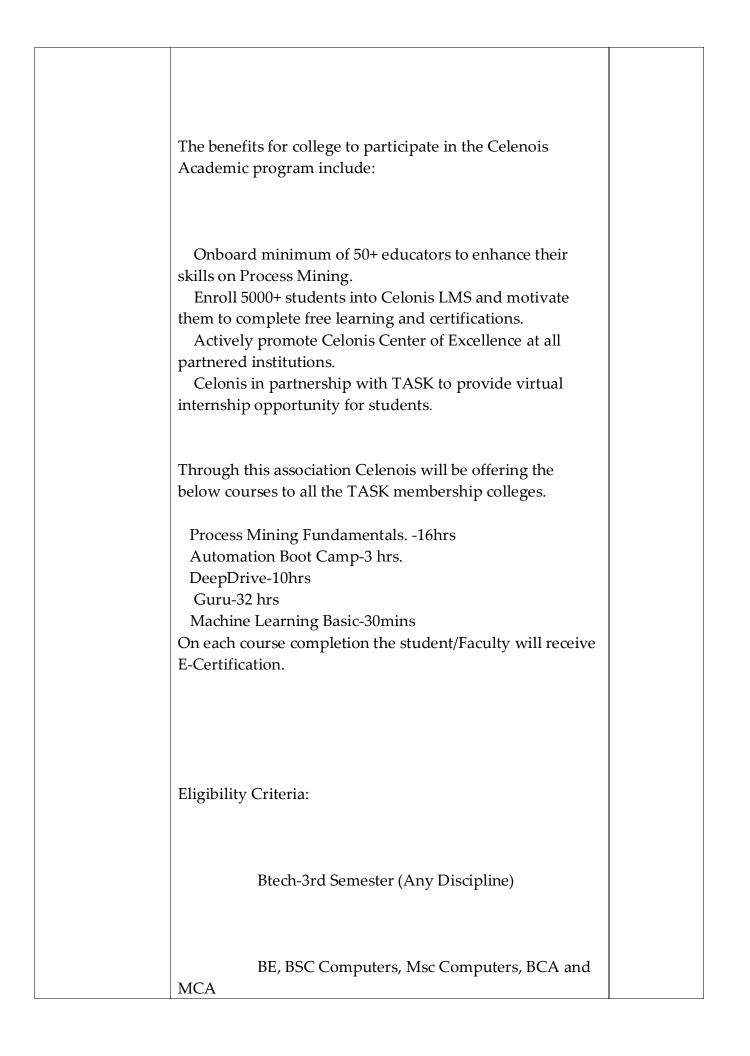
• Offer a high quality curriculum that combines in-depth technical training and professional skills development.

• Choose courses to complement your academic program and prepare students for certification.

• Leverage our learning platform to track student success and connect with educators and Process Mining experts.

• Local student chapter including annual project competitions, workforce development programs and others.

• Point students to career building resources created with employers for our alumni.



	For more information about the program URL for your reference: https://www.celonis.com/academic-alliance/	
	Kindly reach out to me in case of any questions.	
TASK-Celonis Academic Alliance Partnership -	ear Sir/Madam,	27.07.2022
Online Meeting on 1st August 22	TASK have planned Celonis Academic Alliance Partnership - Meeting on 1st August 22.	
	Joining Link: https://celonis.zoom.us/j/98920600021?pwd=SlpsNXIxb2d1S 2hZQWNjVWw5RFZJZz09#success	
	ID: 98920600021 passcode: j75S3EkZ	
	Timings: 03:00 PM to 04:00 PM	
	Kindly join the meeting on time.	
	Partnership Details:	

TASK in association with" Celonis Academic Alliance" is giving opportunity to all colleges to become a member of " Celonis Academic Alliance Program" to avail opportunities designed exclusively for Institutions, Faculties and their Students.
About Celonis Academic Alliance Program:
Celonis is the worldwide leader in Process Mining and Execution Management that empowers people to analyze any gap while creating, building and sharing their innovations. Our technology is changing the nature of work and the way we live. Our solutions deliver a brave new model of technology that is fluid, instant and more secure. Customers can innovate faster by rapidly developing, automatically delivering and more safely consuming any application.
Celonis collaborates with TASK as a Regional Academic Alliance Training Partner to enhance students and educators on Process Mining and EMS tool and make them ready to deploy resources at no cost.
As a part of efforts to empower academia, TASK and Celonis Academic Alliance Program work jointly on the following:
• Offer a high quality curriculum that combines in-depth technical training and professional skills development.
Choose courses to complement your academic

program and prepare students for certification.
• Leverage our learning platform to track student success and connect with educators and Process Mining experts.
 Local student chapter including annual project competitions, workforce development programs and others.
• Point students to career building resources created with employers for our alumni.
The benefits for college to participate in the Celenois
Academic program include:
Onboard minimum of 50+ educators to enhance their skills on Process Mining. Enroll 5000+ students into Celonis LMS and motivate them to complete free learning and certifications. Actively promote Celonis Center of Excellence at all
partnered institutions. Celonis in partnership with TASK to provide virtual internship opportunity for students.
Through this association Celenois will be offering the below courses to all the TASK membership colleges.
Process Mining Fundamentals16hrs Automation Boot Camp-3 hrs. DeepDrive-10hrs Guru-32 hrs Machine Learning Basic-30mins
On each course completion the student/Faculty will receive E-Certification.

	Eligibility Criteria:	
	Btech-3rd Semester (Any Discipline)	
	BE, BSC Computers, Msc Computers, BCA and MCA	
	For more information about the program URL for your reference: https://www.celonis.com/academic-alliance/	
TASK - HYSEA	Kindly reach out to me in case of any questions. Dear Sir/Madam,	06.07.2022
WINtalk series on Cloud Computing	Greetings from TASK!	
	TASK would like to conduct WINtalk series session on 'Cloud Computing' on 8th July 2022 from 3:00 pm to 4:00 pm.	
	The panelists for the session are Mr.Arun Michael,Cloud Delivery Head - EMEA Region, Tech Mahindra Ltd and Mr.Ramesh Swaminathan,Principal Product Manager, Informatica.	

	Moderator for the session Mr.Shrikant Sinha, CEO -TASK.	
	Joining link for the session https://task.radiusedutech.com/	
	Attached is the flyer for WINtalk series session.	
TASK CEO - Principal meet -	Request you to share with all faculty and students and encourage them to make use of this opportunity. Dear Sir/Madam,	29.06.2022
3PM,30th June 22(Tomorrow) at TARA Government	Greetings from TASK!	
College, Sangareddy	TASK would like to invite you to meet TASK - CEO at 3:00 PM, 30th June 2022(tomorrow) at Tara Government College, Sangareddy.	
	Request you to please attended the meeting, He would like to meet all the Principals & TASK Coordinators. Dear Madam,	29.06.2022
TASK-Oracle		
FDP on		
Artificial	Greetings from TASK!	
Intelligence with		
Machine		
learning in	TASK in association with Oracle Academy is planning to conduct FDP on Artificial Intelligence in Machine learning	
JAVA		

with JAVA August 2022.
with jr vr i rugust 2022.
As I informed earlier, Oracle Academy FDP module has
been upgraded to Supported Self Study (SSS) in which,
now the training will be completed in 9 weeks.
In SSS, the training will be setup in Member Hub and the
trainer from Oracle will take 2 sessions – orientation session
on first day of training (90 minutes) and final session on last day of training (90 minutes).
ady of training (50 minutes).
Details of scheduled FDP are given below:
Topic: Artificial Intelligence with Machine Learning in Java
Training type: Virtual
Duration: 9 weeks
Start Date (Orientation): 22nd August Monday 2022
End Date (Final Session): 24th October Monday 2022
Target Audience: Degree and Engineering Faculty.
No of Faculty: Minimum 35 and Maximum 55
Link to nominate for FDP:
https://forms.gle/oD5mg1R6tbAkbezQ7
Last date for nomination: 4th July 2022
Faculty has to complete the course on their own in 9 weeks
or less. In case, they need support from trainer then they

	can write a mail or contact the trainer directly and accordingly a separate zoom call will be setup by the trainer for resolving those queries.	
	Points to be noted:	
	• College must be a Oracle Academy member	
	• Faculty must have college domain email id to register for the FDP	
	• Faculty should register in Oracle Academy to enroll for the FDP(Deatils will be shared after giving nomination in above link)	
	• Faculty who will complete all the quizzes and exams will be received certificate directly from Oracle Academy	
	• Faculty members have requested to deliver the same course to their students after successful completion of FDP.	
	• Post FDP instructions will be shared with all certified faculty once the training is completed.	
	Please share this information with interested faculty members to nominate in above link for the FDP with their domain email id by 4th July 2022.	
TASK - Skill Next 360Digi TMG Training on Cloud	Dear Sir/Madam,	22.06.2022
Computing from 4th to 7th July 2022	Greetings from TASK!	
	TASK in collaboration with 360Digi TMG is conducting	

training on Cloud Computing from 4th of July 2022.	
Training details are as followed:	
Title of the Module : Cloud computing Duration : 5 Days (1.5 Hrs / Day; Total 7.5Hrs)	
Start Day / Date : Monday, 4th July 2022	
End Day / Date : Thursday, 8th July 2022	
Timings : 7.30 PM – 9 PM	
Resource Person : Mrs Srija	
Student Eligibility : Any Graduate / Post Graduate	
End Date of registration :- 1st July 2022	
Registration link:- https://docs.google.com/forms/d/e/1FAIpQLSeYewzF4IYGx nVc44HbsNOF_nrgy6NACbx65a37hY6sDNz1rA/viewform ?usp=sf_link	
Or	

	https://forms.gle/RcTsFhGrPXF8sR8s7	
	Request you to circulate to All student.	
TASK - Skill	Dear Sir,	21.06.2022
Next 360Digi		
TMG Trainings		
For the Week	Greetings from TASK!	
from 4th july to		
7th july 2022		
	TASK in collaboration with 360Digi TMG is conducting trainings on "Social media marketing" from 4th July 2022.	
	Training details are as followed:	
	Title of the Module : Social media marketing	
	Duration : 4 Days (1.5 Hrs / Day; Total 6Hrs)	
	Duration . 4 Days (1.5 mis / Day, 10tal 01115)	
	Start Day / Date : Monday, 4th July 2022	
	Fred Days / Data , Three days 7th Isla 2022	
	End Day / Date : Thursday, 7th July 2022	
	Timings : 7.30 PM – 9 PM	
	Resource Person : Mr. Enoch Joy	
	Charlent Elizibilitza Arra Craduata / D. J. C. J. J.	
	Student Eligibility : Any Graduate / Post Graduate	

	End Date of registration :- 1st July 2022	
	Registration link:-	
	https://docs.google.com/forms/d/e/1FAIpQLSdSguEVSxUX	
	sOn00ccaQC9- MZIiTHqgJE9HPVzokgCahjbBWA/viewform?usp=sf_link	
	Request you to circulate to the students.	
TASK-HYSEA WINtalk Series	Dear Sir/Madam,	18.06.2022
on Fullstack		
	Greetings from TASK!	
	This mail is regarding WINtalk series session on 'Fullstack'	
	on 24th June 2022 from 3:00 pm to 4:00 pm.	
	The panelist for the session Mr.Paparao Kapuganti,Senior	
	Principal Architect,Infosys Ltd and Ms.Sirisha Peyyeti,Head of Consulting,Roundsqr.	
	Moderator for the session Mr.Shrikant Sinha, CEO -TASK.	
	Joining link for the session https://task.radiusedutech.com/	
	Attached is the flyer for WINtalk series session.	
	Request you to share with students and encourage them to	

	make use of this opportunity.	
AWS Deep Racer Student League	Dear Sir/Madam,	16.06.2022
	Greetings for the day!	
	AWS has launched the Deep Racer Student League-India 2022 Program in the presence of the Honorable Minister of Education and Minister of Skill Development and Entrepreneurship, Shri Dharmendra Pradhan ji, AWS DeepRacer Student league aims to skill students on emerging technologies via a game based racing stimulator.	
	In continuation with the successful AWS DeepRacer Women's league in 2021, this year's event is open to all students who are 18+ in age and enrolled in a recognized academic institutes. Throughout the course of the event, there will be different self-learning and enablement activities that will help students to get started on Machine learning using AWS Educate and DeepRacer content.	
	This year's AWS DeepRacer Student league, brings exciting employment opportunities for top 100 winners from AWS partners and an opportunity to win the AWS Artificial Intelligence (AI) and Machine Learning (ML) Scholarship program, in collaboration with Intel and Udacity.	
	To know more kindly visit AWS DeepRacer Student league -India 2022.	
	Awards and Recoginition for participants:	

1st Prize :	
- Amazon Vocher worth 1,00,000	
- Amazon Branded Swag Kit	
- Direct entry to AWS Global DeepRacer Student Championship Cup	
2nd Prize :	
- Amazon Vocher worth 75,000	
- Amazon branded Swag Kit	
3rd Prize:	
- Amazon Vocher worth 50,000	
- Amazon branded Swag Kit	
- Top 10 will be awarded with Amazon Vocher worth 5,000 and Amazon branded Swag Kit.	
- Top 100 will be awarded with Digital certificate recognizing them as Semi-finalist.	
- All Participants will be awarded with Digital Certificate.	
Click the below link to register for the program:	
https://tinyurl.com/2p8k5z2c	

	Last Date for Registration 6th July 2022.	
	Request you to share with the students and encourage them to register where they can make use of this opportunity. Dear Sir/Madam,	16.06.2022
CDAC FDP on		
Advance IOT	Greetings from TASK,	
	I am glad to inform you that TASK in collaboration with CDAC and NASSCOM initiative, "Future Skills PRIME" is conducting a Faculty Development program on Advance IoT(online).	
	The training program will be held in a blended model	
	Program Details are as follows: -	
	Course Title: - Advanced IoT.	
	Dates: - 20th June to 11th July 2022	
	Mode of Delivery: - Blended model(both online & offline)	
	AGENDA OF THE TRAINING: - 20TH JUNE (9 AM-4PM) 23RD JUNE (9 AM-4PM)	
	27TH JUNE (9 AM- 4PM)	

	30TH JUNE (9 AM- 4PM)	
	4TH JULY (9 AM- 4PM)	
	7TH JULY (9 AM- 4PM)	
	11TH JULY (9 AM- 4PM)	
	Certificate will be provided after the assessment test.	
	Those, who will enroll, will also get a chance to attend 4- day guided practical session at our E-City campus on mutually convenient date. Food and lodging will be provided to participants free of cost during practical sessions.	
	So request you to give your faculty nominations in below link by 17th June 2022(by EOD).	
	https://docs.google.com/forms/d/e/1FAIpQLSfvF8U- VK5AuK4hw6pAqzqyLCykJqUAtw9is3MNwRPu71FMBw /viewform	
	Please find attachment for execution plan of the program and IOT syllabus.	
	Feel free for any queries.	
TASK-TechGig Code Competition	Dear Praveena Garu,	23.05.2022

Greetings from TASK!	
TASK in association with TechGig is planning for Code competition 2022.	
About TechGig:	
TechGig is a division of Times Internet Limited, India's largest digital products company. We are India's largest and fastest growing developer community of 4.2 million software professionals. We are an innovative and enthusiastic technology community of super-active developers who love to compete and showcase their skills, learn new technologies, and keep up with the latest technology news to grow in their career.	
We curate technology news, deep insights on latest technologies, host exciting skill contests and help developers' upskill on new-age technologies. Our coding events, Code Gladiators and Geek Goddess are well-known in the technology industry. We bring to our readers the latest software updates, deep insights on programming languages, gadget previews and a quirky take on all things technology in our podcasts and videos. We help developers learn, compete and grow in their career.	
Registrations Link for the competition:	
https://www.techgig.com/codegladiators/signup?sourcetyp e=campus_TASK	

	Last Date for registration 6th June 2022. Participation of students in this competition will enable	
	them to: 1. Win cash prizes worth INR 50 Lakh*	
	2. Create developer profiles and get closer to their dream job.	
	3. Job Opportunities with top IT companies	
	Eligibility: Any student above 18 years and interested in coding is eligible to apply irrespective of his academic year and stream	
	Request you to kindly share with your respective colleges and students and encourage them to participate in the competition.	
	Thanks,	
TASK - HYSEA WINtalk series on Deveops	Dear Sir/Madam,	12.05.2022
	Greetings for the day!	

	This mail is regarding WINtalk series session on 'DevOps' on 13th May 2022 from 3:00 pm to 4:00 pm.	
	The panelists for the session are Ms. Rajita Singh,Head HR,Broadridge and Mr.Satyen Sharma,Head of Delivery,Technology,State Street.	
	Moderator for the session Mr.Shrikant Sinha, CEO -TASK.	
	Joining link for the session https://task.radiusedutech.com/	
	Attached is the flyer for WINtalk series session.	
	Request you to share with colleges and students and encourage them to make use of this opportunity.	
Salesforce	Dear Sir/Madam,	12.05.2022
Fundamentals Free Virtual Program for 3 weeks- Starting 19th May 2022	Greetings from Telangana Academy for Skill and Knowledge !!	
	Join Salesforce Fundamentals, a FREE virtual learning journey to get on the path to earning your first Salesforce credential, and discover your career in the cloud.	
	It's a three-week course, each week is 2 hours of self-guided	

	learning and 1 hour of live teaching. Weekly sessions are every Thursday at 9:30am IST starting May 19th 2022.	
	From Marketing, Sales, and Consultant roles, to, Developer, and Administrator ones, there's plenty to gain for anyone who clicks or codes.	
	Eligibility: TASK Registered 1st and 2nd year students of B.E/Btech, BCA, B.Sc, BCom streams. TASK Registered 1st year students of MBA, MCA streams.	
	Students Registration Link: https://forms.gle/cVCHNUcgeqcXkEji9	
	Last Date to register: 17th May 2022	
	Check out what students will learn:	
	The basics of Salesforce, our platforms, and the Salesforce ecosystem How Salesforce skills lead to in-demand and high-paying	
	careers How to navigate landing your first role in the ecosystem	
TASK - HYSEA WINtalk series on **TESTING**	Dear Sir/Madam,	28.04.2022
for students and faculty	Greetings from TASK!	
	TASK - WINtalk series session on ""TESTING" on 29th	

	April 2022 from 3:00 pm to 4:00 pm.	
	The panellists for the session are:	
	Mr. Hemchand Srigiriraju AVP – Delivery & Global Head of Testing, ZenQ	
	Mr. Prashanth Achanta EVP & CTO, firstsource	
	Mr. Sesha M. Rao Board Member, Insideview Technologies	
	Joining link for the session https://task.radiusedutech.com/	
	Attached is the flyer for WINtalk series session.	
	Request you to share with all faculty and students to make use of this opportunity.	
Salesforce Virtual Internship	Dear Sir/Madam,	09.04.2022
Program 2022 Launch- 11th April 2022- 11	Greetings from Telangana Academy For Skill and Knowledge.	

AM till 12 PM		
	TASK in association with Smartbridge is launching the Virtual Internship Program supported by Salesforce on 11th April 2022, in partnership with AICTE, NASSCOM FutureSkills, Monster India & GitHub.	
	We have attached the Launch Program agenda, Salesforce Virtual Internship details.	
	Virtual Internship Program is a 8 week project based learning program delivered through smartinternz platform. The program consists of live hands-on training sessions, self-paced learning on the salesforce trailhead platform, Ask a mentor sessions, project development, soft skills training, and followed by a career fair for eligible students.	
	Benefits to student:	
	100+ hrs of experiential learning on salesforce technologies Work on a Guided project with Mentor support Acquire super badges and build salesforce trailhead trailblazer profile Virtual internship completion certificate Access to career fair organized by hiring partners only for final year & passed-out students	
	Virtual Internship Roles:	
	Virtual internship program has two learning tracks as listed below	

	Salesforce Administrator Track : This learning track is suitable for the learner with limited coding skills. Salesforce Developer Track: This learning track is suitable for the learners with good coding skills. The learners enrolling for this track will be routed through Nasscom FutureSkills Platform for NOS (National Occupation Standard) aligned course.	
	Eligibility: Students currently pursuing BE, B.Tech,MCA, MBA, B.Sc, etc belonging to 2022 and 2023 passout batch. Students who graduated in 2020 & 2021 can also apply. Students must be available virtually for a period of two months or 8 weeks.	
	Target Audience for the launch: Principals, TPOs, HODs, Faculties of Engineering, Degree and MBA colleges.	
	Please register for the launch using the following link : https://us06web.zoom.us/meeting/register/tZcrf- 6qpzIqE9JsaO-mnJ619RpbwIMflQop	
	Program timings: 11 AM till 12 PM	
	Please register in advance for this meeting and after registering, you will receive a confirmation email containing information about joining the meeting.	
TASK-HYSEA Virtual Job Fair	Dear Sir/Madam,	21.03.2022
	TASK in association with HYSEA (Hyderabad Software	

Enterprises Association) is taking great pleasure in announcing the 3rd edition of HYSEA JOB FAIR(Virtual) targeting Engineering (B.E/B.Tech) and Non-Engineering (MCA, MBA, BCA, B.Sc. Comp Science, BBA, B.A, B.Sc. & B.com) pass outs of 2020, 2021 and 2022. This year HYSEA is planning to target Tier-3 towns and colleges in Telangana. In addition to the tech jobs, the Job Fair will also assist in hiring for ITES/BPO & other jobs.	
A unique feature of this year's job fair includes focusing on the hiring of Women from the Economically Weaker Section of the Society who are trained by our NGO Partner NIRMAAN. For this category graduates from the year, 2019 will also be considered.	
All eligible candidates interested in participating in this VIRTUAL JOB FAIR will be required to Register Online at https://hyseajobfair2022.in/	
The registrations open on 1st March 2022 and the last date to register is 28th March 2022.	
Students/Candidates are advised to read the details of the Job Fair at Webpage: https://hyseajobfair2022.in/ before registering.	
Registration Link: https://hyseajobfair2022.in/student- registration.php	
Key points to be noted by Candidates/Students:	
Each candidate/student participating in the job fair will	

	 choose the top 10 preferences from the participating companies. Shortlisted candidates' details will be shared with participating companies based on the number of open positions, skill set required & candidate preference. Once details of the candidates are shared with companies, they are free to follow their hiring process of further tests/interviews with shortlisted candidates. It is possible that a candidate may receive multiple offers from different companies. HYSEA reserves the right to share all candidates' data with all registered companies at a later stage. It is entirely up to the participating companies to shortlist or make a job offer. HYSEA is just a facilitator. 	
	For any help/info with the registration process, please feel free reach me out.	
	Dear Sir/Madam,	12.03.2022
TASK Oracle 5-		
Day Boot Camp		
and Build-a- Thon Application	TASK in association with Oracle Academy is planning to conduct 5-Day Boot Camp and Build-a-Thon Application Development using Oracle APEX	
Development using Oracle APEX	About the event:	
	TASK in association with Oracle Academy is launching a program exclusively for students to provide Hands-On Experience on building projects. Exposure to live projects will help students to gain technical skills and also gain Hands-On experience on Oracle APEX	

Eligibility :	
Students from 3rd & Final year CSE, IT and ECE	
backgrounds	
Students from Final Year Degree which support computers	
and Information technology.	
Students who will complete the course Database	
Programming with SQL Learner English Course in Oracle Academy will have have an opportunity to attend 5 - Day	
Boot Camp.	
Registration link for the Boot camp will be only sent to	
students who have certified on Database Programming	
with SQL Learner English Course in Oracle Academy.	
Students who will attend 5- Day Boot camp only can	
participate in Project Build-A-Thon.	
Benefits of the Program:	
Students will be trained on Oracle Application Express – Application Development Foundations content during 5	
Days.	
Students will have an understanding on how to make use	
of Oracle APEX platform.	
Exposure to live Projects will help students in improving	
technical skills.	
Students will be presented with awards based on their	
performance.	
Mentoring support from trainers.	
Tentative Dates:	
7th April 2022 to 7th June 2022	
Program Agenda:	
Event launch	
E learning- Database Programming with SQL Learner	

	English in Oracle Academy 5- Day Boot camp Project Build-a-Thon Project Reviews & Announcing Top 15 projects Final Jury Round & Winner Announcement	
	Students would need faculty assistance to complete E learning on Database Programming with SQL Learner English Course in Oracle Academy	
	So, Request you to kindly take faculty nominations (Domain Email Id) from Oracle registered colleges(attached OA registered list for your reference) in below link by 15th March 2022.	
	https://forms.gle/18CF9WzC1Sb2ifwy5	
	Onboarding students into Oracle Academy process will be shared with Faculty members once we get enough nominations.	
	Please try to push more college participation	
	Let me know in case of any queries.	
SCOST NIPER Hyderabad 6- month Pharmaceutical EDP with Scholarship – reg	Dear Sir/Madam, Greetings from TASK!!	12.03.2022

We are happy to inform you that NIPER, Hyderabad has been awarded a grant from DBT, GoI through Telangana State Council of Science and Technology (TSCOST) to conduct a Skill Vigyan Program in the Pharmaceutical Sector. A Six-month Entrepreneurship Development Program (EDP) is being organized by the Department of Pharmaceutical Management, National Institute of Pharmaceutical Education and Research - Hyderabad, during March – September 2022.

This program is designed to enhance the understanding of the participants on the Entrepreneurship Journey, Various Opportunities & Challenges encountered in the Journey and the support available from the Government for Entrepreneurs.

It is a 6-month full time program where the participants will be given classroom/laboratory training for 2 months at NIPER Hyderabad and industrial training for 4 months. The participants will receive a stipend of Rs. 10,000 per month for six months.

Applications are called from interested candidates of all the Degree, Degree & PG, Pharmacy, and Engineering colleges registered with TASK. Last date for applications is 15th March 2022.

As Telangana Academy for Skill and Knowledge is known for its industry academia connect to help the youth acquire the required skillsets for future, we request your kind support to circulate the Entrepreneurship Development Program (EDP) details (Brochure and application form enclosed) to the passed-out pharmacy/related science discipline students in colleges registereded with TASK.

	Students with entrepreneurial interests from these colleges may kindly be encouraged to apply for the program.	
	We look forward to your support in reaching the entrepreneurial aspirants from the TASK registered colleges.	
	For any queries, please write to edp.niperhyd@gmail.com	
TSCOST NIPER Hyderabad 6- month Pharmaceutical EDP with Scholarship – reg	Dear Sir/Madam, Greetings from TASK!!	08.03.2022
	We are happy to inform you that NIPER, Hyderabad has been awarded a grant from DBT, GoI through Telangana State Council of Science and Technology (TSCOST) to conduct a Skill Vigyan Program in the Pharmaceutical Sector. A Six-month Entrepreneurship Development Program (EDP) is being organized by the Department of Pharmaceutical Management, National Institute of Pharmaceutical Education and Research - Hyderabad, during March – September 2022.	
	This program is designed to enhance the understanding of the participants on the Entrepreneurship Journey, Various Opportunities & Challenges encountered in the Journey and the support available from the Government for Entrepreneurs.	

	It is a 6-month full time program where the participants will be given classroom/laboratory training for 2 months at NIPER Hyderabad and industrial training for 4 months. The participants will receive a stipend of Rs. 10,000 per month for six months.	
	Applications are called from interested candidates of all the Degree, Degree & PG, Pharmacy, and Engineering colleges registered with TASK. Last date for applications is 15th March 2022.	
	As Telangana Academy for Skill and Knowledge is known for its industry academia connect to help the youth acquire the required skillsets for future, we request your kind support to circulate the Entrepreneurship Development Program (EDP) details (Brochure and application form enclosed) to the passed-out pharmacy/related science discipline students in colleges registereded with TASK.	
	Students with entrepreneurial interests from these colleges may kindly be encouraged to apply for the program.	
	We look forward to your support in reaching the entrepreneurial aspirants from the TASK registered colleges.	
	For any queries, please write to edp.niperhyd@gmail.com	
TASK - 'JobFair@JNTUH ' 15th & 16th March 2022	Dear Sir/Madam,	04.03.2022
	Greetings from TASK!	

	Announcing 'JobFair@JNTUH', a multi-industry and multi- profile job fair organised by Solvix and supported by TASK and DEET as part of JNTU's Golden Jubilee celebrations on 15th & 16th March 2022.	
	With 20,000+ vacancies and over 100 companies participating in this job fair, we will help you connect with some of the biggest employers and a huge array of jobs to choose from. But hurry as registrations are filling in fast.	
	Kindly share with student.	
	Last Date to Register 11th March 2022	
	Registration link: https://forms.gle/BfP88d8JB3jUSeCj8. Please note that there is no registration fee.	
TASK-HYSEA Virtual Job Fair	Dear Sir/Madam,	04.03.2022
	TASK in association with HYSEA (Hyderabad Software Enterprises Association) is taking great pleasure in announcing the 3rd edition of HYSEA JOB FAIR(Virtual) targeting Engineering (B.E/B.Tech) and Non-Engineering (MCA, MBA, BCA, B.Sc. Comp Science, BBA, B.A, B.Sc. & B.com) graduates of 2020, 2021 and 2022.	
	A unique feature of this year's job fair includes focusing on the hiring of Women from the Economically Weaker	

Section of the Society who are trained by our NGO Partner NIRMAAN. For this category graduates from the year, 2019 will also be considered.	
All eligible candidates interested in participating in this VIRTUAL JOB FAIR will be required to Register Online at https://hyseajobfair2022.in/	
The registrations open on 1st March 2022 and the last date to register is 15th March 2022.	
Students/Candidates are advised to read the details of the Job Fair at Webpage: https://hyseajobfair2022.in/ before registering.	
Registration Link: https://hyseajobfair2022.in/student- registration.php	
Key points to be noted by Candidates/Students:	
Each candidate/student participating in the job fair will choose the top 10 preferences from the participating companies. Shortlisted candidates' details will be shared with participating companies based on the number of open positions, skill set required & candidate preference. Once details of the candidates are shared with companies, they are free to follow their hiring process of further tests/interviews with shortlisted candidates. It is possible that a candidate may receive multiple offers from different companies. HYSEA reserves the right to share all candidates' data with all registered companies at a later stage	
all registered companies at a later stage. It is entirely up to the participating companies to shortlist	

	or make a job offer.	
Verisk Applied students data	Dear Sir/Madam,	07.02.2022
	Greetings from TASK!	
	Please find the attachment of the applied students for Verisk. Students will receive communication mail from the company for the interview.	
TASK Oracle FDP on Artificial Intelligence in	Dear Madam,	25.01.2022
Machine Learning with Java March 2022	Greetings from TASK!	
	TASK in association with Oracle Academy is planning to conduct FDP on Artificial Intelligence in Machine Learning with Java (Duration : 9 weeks) in the month of March 2022.	
	Oracle Academy FDP module has been upgraded to Supported Self Study (SSS) in which, now the training will be completed in 9 weeks.	
	In SSS, the training will be setup in Member Hub and the trainer from Oracle will take 2 sessions – orientation session on first day of training (90 minutes) and final session on last day of training (90 minutes).	
	Details of scheduled FDP are given below:	

Topic: Artificial Intelligence in Machine Learning with Java	
Training type: Virtual	
Duration: 9 weeks	
Start Date (Orientation): 11th March 2022	
End Date (Final Session): 13th May 2022	
Target Audience: Degree and Engineering Faculty.	
Link to nominate for FDP: https://forms.gle/UYdkBWJZbDu3oEj69	
Last date for nomination: 30th Jan 2022	
Faculty has to complete the course on their own in 9 weeks	
or less than 9 weeks. In case, they need support from trainer then they can write a mail or contact the trainer directly and accordingly a separate zoom call will be setup by the trainer for resolving those queries.	
Points to be noted:	
· College must be a Oracle Academy member	
• Faculty must have college domain email id to register for the FDP	
• Faculty should register in Oracle Academy to enroll for the FDP(Deatils will be shared after giving nomination in above link)	
• Faculty who will complete all the quizzes and exams will be received certificate directly from Oracle Academy	
• Faculty members have requested to deliver the	

	 same course to their students after successful completion of FDP. Post FDP instructions will be shared with all certified faculty once the training is completed. Please nominate faculty to attend the FDP with their domain email id by 30th Jan 2022 	
TALLY Training	Dear Madam / Sir,	27.12.2021
Schedule; Date-	Greetings from TASK!	
28th to 30th, December-2021 & Time: 9:30 AM to 4:30 PM	We are confirming the Tally training schedule at your college for the proposed period with the following details. We request you to follow the Covid norms.	
	TALLY Training Schedule:	
	28th to 30th December, 2021 Trainer details	
	1). Mr. Ramakrishna - 91007 98186	
	Following are the requirements from our end for TALLY Training Sessions	
	As a part of our ongoing TASK skilling initiatives, we have scheduled TALLY training sessions at your college on TALLY with GST for all years TASK registered commerce students. This is for six hours each day, ideally between 9:30 AM and 4:30 PM including 60 minutes for lunch. Each class to have 40 to 50 students max.	

[
	In this connection we would like you to provide us the following:	
	1. Neat and Tidy Class Room with seating capacity of 50.	
	2. Projector with Sound System.	
	3. White Board & Markers.	
	4. Lunch and Tea for trainer	
	5. Collar mikes or normal mikes	
	Also a request regarding help with accommodation. Since the trainer will be travelling from Hyderabad it would be great if your college can provide the same as it would also be safer.	
	Also a request regarding help with local transport, it would be great if your college can provide a cab for pick and drop of the trainer, as many of them are not familiar about the college locations at long distances.	
	If at all any cancellation regarding this schedule please let us know immediately	
	Please find the attachment of attendance. We will soon share the feedback form, assessment and request to handover all the documents to the trainer after the sessions.	

		· · · · · · · · · · · · · · · · · · ·
	We would request you to take the print out of the same attachment which we are sending for attendance.	
	Thanks in advance for your support.	
	Suiderri Cuedume Project Managore 0020021(24	
	Sridevi Guduru, Project Manager - 9989931684	
	Telangana Academy for Skill and Knowledge	
	Begumpet	
TASK- BLUEPRISM	Dear Sir/Madam,	23.12.2021
ACADEMIA	Please find the recordings of Day 4 below.	
FDP DETAILS	Day 4 - <u>https://youtu.be/EP2fiIVnCn8</u>	
TASK - ExcelR - Training on Data Science	Dear Sir/Madam,	21.12.2021
	Greetings from TASK!	
	TASK in association with ExcelR is conducting a Instructor led virtual Training Program on Data science. This program is offered to all the TASK registered Degree colleges. Below are the complete details and attached is the EDM.	
	Training Program :-Data science	
	Training Start Date :04-01-2022	

	Training End Date : 2-02-2022	
	Note:- (we will not have training on Saturday & Sunday)	
	Timings : 6.00 PM to 8.00 PM	
	Note:-13th and 14th of jan also we will not have the training as they are festival holidays.	
	Registration link : https://docs.google.com/forms/d/e/1FAIpQLSdtiOoSNlqUv KgvmA1alMXUwamNUWypp8XGet_epCDNgEfwQg/vie wform?usp=sf_link	
	OR	
	https://forms.gle/RU6327Ct4jQddLsE6	
	Faculty from Degree can attend this training program. Topics such as Data Science Project Lifecycle , Data Mining Unsupervised , Data Mining Unsupervised , will be covered in this training program.	
	Request you all to take this forward to your respective Degree colleges and take the nominations .	
	Request you all to take the nomination before 30th december.	
TCS BPS Fresher Hiring for 2022 Graduates	Dear Sir/Madam,	20.12.2021

	Creating of from TACK!	
	Greetings from TASK!	
	We thank you for your continuous support in identifying quality talent and greatly value your contribution towards TCS BPS Fresher Hiring.	
	With pleasure, we would like to invite your students for "TCS BPS Fresher Hiring" for YOP 2022 Arts, Commerce & Science Graduates.	
	For more details and to register : Pls. refer to the below poster or click on this link <u>https://on.tcs.com/3D8qBkQ</u>	
	Dear Educators,	20.12.2021
CII-EDU		
SUMMIT on 21		
December 2021	CII-EDU SUMMIT	
at HICC	(A cademic Excellence in Higher Education through	
Novotel:	'Academic Excellence in Higher Education through Industry Collaborations'	
Madhapur:		
Hyderabad	1000-1630 hrs: 21 December 2021; Hotel HICC Novotel, Madhapur, Hyderabad	
	I am pleased to inform you that CII Telangana in collaboration Department of Collegiate & Technical Education, Government of Telangana is organizing EDU Summit with the theme 'Academic Excellence in Higher Education through Industry Collaborations' between 1000- 1630 hrs on 21 December 2021 at Hotel HICC Novotel, Madhapur, Hyderabad. (Program registration starts at 0915 hrs).	
	The objective of the Edu Summit is to deliberate on the	

	Need for Strengthening the Core Courses, Accreditation & Ranking Processes, Importance of Multi-Disciplinary Education, and Developing the Academic Leaders. We are writing to cordially invite you and colleagues to join the conference.	
	Kindly note that there is no registration fee however prior registration is mandatory and seats at the venue will be allocated on a first-come-first-serve basis. Kindly note that COVID protocols will be followed at the program venue.	
	TASK is collaborating Partner for EDU Summit. We request you to use the attached Registration form for nominations of TPO/Faculty from your college. Please share the filled up Registration form with aruna.mantena@cii.in	
	Registration Link for the program at Hotel HICC Novotel: https://bit.ly/CIIedusummit2021-HICCNovotel	
	Program flyer copies are attached for reference.	
	We look forward to your support and confirmation from your college.	
Infosys Spring board Career Compass - Journey from Me to We December 21 - 23, 2021	Dear Sir/Madam, Greetings from TASK!	18.12.2021
	TASK – Infosys Spring board would like to conduct Career	

	Compass from 21st to 23rd Dec 2021 from 3:30 PM to 5:30 PM.	
	Kindly send the attachments to all students .	
TASK- BLUEPRISM ACADEMIA	Dear Sir/Madam, Greetings for the day!!	17.12.2021
FDP DETAILS	Please find the details of scheduled Blue Prism academia program FDP :	
	Topic : Foundation Training Training Type : Virtual Duration: 3 hrs per day Start Date : 20-12-2021 End Date : 23-12-2021	
	Trainer : Smart Bridge trainer Time:10 am to 1:00 pm Session Link - https://zoom.us/meeting/register/tJ0rcOitqT8oGND6KSoaN 1dnIbX8PsrAtinJ	
	Benefits:	
	Participation Certificates would be provided.	
Blue Prism TASK FDP- Faculty Nominations	Request all the faculty to join the session on time. Dear Madam,	15.12.2021
ivonimations	Greetings from TASK!	
	TASK – Blue Prism would like to conduct a Faculty Development Program(FDP) from 20th to 23rd Dec 2021.	
	Request you to please nominate the faculty in below link by	

	[]
17-12-2021 end of the day.	
https://forms.gle/AP8TqsMsHKtEu0a28	
Details of scheduled Blue Prism academia program FDP are given below:	
Topic : Foundation Training	
Training Type : Virtual	
Duration: 3 hrs per day	
Start Date : 20-12-2021	
End Date : 23-12-2021	
Trainer : Smart Bridge trainer	
Time:10 am to 1:00 pm	
Benefits:	
Participation Certificates would be provided.	
Eligibility Criteria:	

		1
	Faculty of Engineering. Faculty of Degree(B.com computers and B.sc and BCA).	
	Kindly let me know if you need any additional information from my end.	
Task-BluePrism	Dear Sir/Madam,	10.12.2021
Academia Program partnership Certificate	Please find the attachment of Blueprism academia program partnership certificate.	
Recruitment -	Dear Sir/Madam,	09.12.2021
TCS BPS Hiring 2021 & 2020 Graduates II BCom, BA, BBA/BBM/BMS, BSc	Request you to share the link to apply with students. Students who were unable to take the online assessment previously due to exams etc also can apply(Students who have applied and are not qualified are not eligible to	
(CS/IT)/General	apply).	
	Attached are the documents for your reference.	
	Link to apply: https://forms.gle/wexydtSwg5uE4vdL9	
	Last date to apply : 12th December 2021	
	Students would receive communication from TCS only.	
	We thank you for your continuous support in identifying quality talent and greatly value your contribution towards TCS Campus Hiring Program. With pleasure, we would like to invite you for 'TCS BPS Off Campus Recruitment'.	
	We are currently hiring 2021 & 2020 BCom, BA, BBA/BBM/BMS, BSc (IT &CS), BCA candidates for	

immediate requirements.	
Interview process - Online Test> HR> TR> MR. Interviews to be conducted virtually	
Job Description- is herewith attached for your kind perusal.	
As the requirement is urgent, joining of selected candidates will be immediate.	
Both, Clearing Online Test & subsequent rounds of Virtual Online Interviews are mandatory for selection and receiving an offer.	
IMPORTANT:	
Looking at the pandemic scenario, the selected candidates shall work from home, once the situation normalizes candidate needs to start reporting to nearest assigned TCS Operations Centre. Candidate may / may not have location preference, decision on deployment will be purely business in nature.	
Please encourage interested candidates to fill in their respective details in the below format and share it with you. (Same excel format attached) Request you to share collated information in one single file and share it with us so that candidates can receive Online Test & Online Interviews related information. DT No. is not mandatory while updating the student details to be considered for the Test. However, you may inform the candidates to register their profile in TCS Next Step portal under BPS section and submit the complete form as DT No. will be required later to process the candidature of selected candidates.	
Candidates who have already registered in TCS NextStep Portal under BPS Section and have their DT No. can share the same no and need not create new profile.	
Important Points Regarding TCS NQT Test:	
 Aptitude test will be remote proctored (online). 	

	 Students should login 15 minutes prior for the test; Eg If test timing is 12pm, students should login by 11:30 or 11:45am. 	
	• If students login after test time – system will not allow them to attempt test.	
	 Test details will be shared on their registered Email ID (as mentioned by students on NextStep Portal). 	
	• Students can appear for Online Remote Proctored Assessment using a laptop/desktop, Mobile devices are not supported for this assessment.	
	Please ensure students are equipped with Windows/Ubuntu OS, Google Chrome browser, Minimum 2 MBPS Internet connectivity & A working webcam as it will be a proctored examination.	
	Dear Professor,	03.12.2021
Campus Connect 2.0	Warm Greetings from Campus Connect Team!	
Launch and Invite for Principals' Meet Wednesday -	Digital Literacy is essential for every individual. Infosys has always benefitted from the association with educational institutions such as yours and the corporate experience helps us anticipate the needs of the industry.	
15 December,	Launching Campus Connect 2.0	
2021	We are pleased to launch Campus Connect 2.0 for our partnering colleges across the country. The offerings are now powered by Infosys Springboard, our online digital platform for learning.	
	We invite you to participate in the Principals' Meet on 15th December 2021 from 10:00 AM - 1:30 PM and join the discussion to strengthen our Industry - Academia partnership.	
	The tentative agenda for the Principals' Meet is as follows:	
	• Infosys Digital literacy mission	

 Campus Connect 2.0 framework and partnership model
o Internship
o Microcredit Courses
o Industry ready new age roles
o Masterclasses
o Research Collaboration Opportunities
o Partnership scorecard model
· Interaction with eminent speakers
· Rewards and Recognition
 Strengthening Industry academia partnership – A Discussion
The Principals' Meet details are given below:
Event Date
Wednesday, 15th December 2021
Time
10:00 AM to 1:30 PM
Venue
Virtual event
Nomination for the event
The success of this event depends on your valuable presence, and hence we would urge you to attend the same along with your Head of Institute (Principal, Chairman, Director), Head of department(max 3 nominations from an

institution).
Nomination Link
https://exp.infosysmeridian.com/register/PMMeet2021
Rewards and recognition
We would like to thank you for your support towards onboarding your students to the platform, leveraging the content and the opportunities offered by Infosys. We are also celebrating educational institutions who have used Infosys Springboard to their advantage through our rewards and recognition awards for the following categories:
Best Ambassadors
Extra Milers
Ace Partners
Institutions that have onboarded all their students onto Infosys Springboard
Institutions that have led in terms of learning time on Infosys Springboard
SPOC's of colleges who have collaborated with Campus Connect
Digital Leaders
Learning Trailblazers
Star Achievers
Institutions who have created and used microsites effectively
Institutions wherein students complete numerous courses
Colleges that complete maximum no. of certifications

	Your institution could stand a chance to win these awards. Please onboard all your students, encourage them to learn and complete the courses & certifications available on the platform. Also create your own microsites and use them in your teaching-learning process. Testimonials If you wish to share a testimonial video or message with us to publish during the event, please share it with us through an email to Springboard-support@infosys.com with the subject line "Principals' Meet – 2021: Testimonials" For any queries or clarification(s), please write to Springboard-support@infosys.com with the subject line "Principals' Meet – 2021: Queries"	
	We look forward to your active participation.	
iNFOSYS GBFS TRAINING PROGRAM- VIRTUAL MODE- STUDENTS LIST	Dear Madam, Thank you for sharing the students data.	01.12.2021
	Kindly share the data in attached format.	
Capgemini Campus drive for BCA and BSc graduates - 2022	Dear Sir/Madam,	01.12.2021
Batch! Registration	Greetings from TASK!	
	Capgemini is a global leader in partnering with companies to transform and manage their business by harnessing the power of technology. The Group is guided everyday by its purpose of unleashing human energy through technology	

	for an inclusive and sustainable future. It is a responsible and diverse organization of over 300,000 team members in nearly 50 countries. With its strong 50-year heritage and deep industry expertise, Capgemini is trusted by its clients to address the entire breadth of their business needs, from strategy and design to operations, fueled by the fast evolving and innovative world of cloud, data, AI, connectivity, software, digital engineering and platforms. The Group reported in 2020 global revenues of €16 billion.	
	Capgemini in India comprises over 150,000 team members working across 13 locations: Bangalore, Bhubaneswar, Chennai, Coimbatore, Gandhinagar, Gurugram, Hyderabad, Kolkata, Mumbai, Noida, Pune, Salem and Tiruchirappalli.	
	We are excited to announce the launch of our campus drive for BCA and BSc graduates of 2022 batch. We cordially invite all the eligible students of your institute to register and apply.	
	Please find below the details of the eligibility criteria for your reference, last date for the registration is 12th December 2021.	
TALLY Training Schedule (Date: 6th Dec to 10th	Dear Sir/Madam,	01.12.2021
Dec 2021 and Time: 2:00 PM to 4:00 PM	Greetings from TASK!	
	TASK is offering TALLY online training to the TASK registered students of B.com (1st, 2nd, 3rd years), M.com and MBA (1st, 2nd years). It is a five-day program and free of cost for the TASK registered students.	

We request you to please inform to the students.	
TALLY Training details:	
Date: 6th, December-2021 to 10th, December – 2021	
Time: 2:00 PM to 4:30 PM	
Training is on Octa platform.	
Student Registration Link:	
https://forms.gle/W4ptpokn72i5RWuc7	
Last date for closing the registrations : 4th, December - 2021 by 11:00 AM	
Requesting to share the above link with the students.	

	Note:	
	We are providing 2 certificates based on the following eligibility criteria	
	1). Participation Certificate - Students must attend all 5 days of training	
	2). Assessment Certificate - Students must attend all 5 days of training and also have to complete the assessment with a minimum of 60% marks. Assessment will be on the last day of the training.	
Virtual event	Dear Sir/Madam,	30.11.2021
hosted by Brookings on		
Digital Mentoring - 2nd Dec 21, 6.30 PM	We hope you're all doing well. We are really excited to share about a virtual event upcoming at Brookings THURS 2nd DEC, 6.30 PM IST, where Mentor Together policy brief on a digital mentoring policy and practice ecosystem for India, will be launched.	
	As you all know that, TASK Partnered with Mentor Together with the objective of Speed Mentoring is to support students to learn skills that are essential for them to get into the workforce. For the details CLICK HERE.	
	The workshop will also have as panelists Hana Brixi, Global Director for Gender, at the World Bank, and Jayesh Ranjan, Principal Secretary, Government of Telangana for Department of Information Technology, Electronics and Communications (ITE&C), and Industries and Commerce (I&C).	

	The event followed by the policy brief builds on evidence from the Mentor To Go program, and has as a central thesis that a digital mentoring policy and practice ecosystem could tackle at scale the triple threat of a skills deficit, a network gap, and restrictive gender norms, which constrain young women in India from equitable labor force participation rate.	
	We would love to have you are all attending if this time and date aligns. Link to sign up here: bit.ly/2dec21.	
	The role of the college is central in our recommendations, so I think it would be really great to have TPOs attend and hear the global perspective on mentoring.	
Student	The overall model of TASK has been a key idea in the digital mentoring ecosystem Dear Praveena ma'am,	30.11.2021
registration form		
	We have closed the pre-placement talk and there is an update on the CTC being revised to 2.5Lacs PA	
	Awaiting registration data to conduct further assessment of the applicants	
	Do call me if you have any further questions	
Taragc Sangareddy - TASK- Naandi	Dear Madam,	26.11.2021

foundation- Mahindra Pride Classroom Training Programme- Batch V- Reg	We are pleased to confirm the Mahindra Pride classroom training program from 29th Nov 2021 onwards. Below are the details of the trainer.	
	Ms. Anitha – Mobile Number: 9030886639	
	Kindly arrange all the necessary arrangements.	
TASK-Esri India: ArcGIS Platform Launch Event: Reimagine	Dear Sir/Madam,	22.11.2021
Location in Your Apps	Greetings from TASK!	
	TASK in collaboration with Esri INDIA planning to launch a new ArcGIS Platform - A location-focused platform-as-a- service (PaaS) offering for the developers in India on 26th November 2021.	
	Request you to share the below information to the faculty and students to enroll for this program.	
TASK - Hackmania is launching global hackathon series	Dear Sir/Madam,	17.11.2021
	Greetings from TASK! TASK in Association with Hackmania is launching a big global hackathon series starting with a bilateral hackathon	
	amongst India and the United States of America.	

Don't forget to secure your spot for DAOHACK – INDIA- USA BILATERAL EDITION, a virtual online hackathon with USD 25,000 as prizes for the top 15 teams.	
Registration Link: https://hackathon.hackmania.com/hackathons/daohack	
Last Date for registrations: 20th November 2021, 6:00 PM IST	
DAOHACK is being organised to find and fund the best solutions built on the Decentralized Autonomous Organisation tech stack.	
Certificate :	
All participants get a blockchain authenticated/verified participation certificate with the details mentioned.	
All winners get a blockchain authenticated/verified winning certificate with the details mentioned.	
(EVERY PARTICIPANT GETS A PARTICIPATION CERTIFICATE authenticated on the blockchain with a unique ID for every single student who registers and participates. This is being presented by our challenge sponsors DOCK.) ALL winners get winning certificates.	

Prizes by USDAO:	
Winner: USD 2,500 in Ethereum. Runner-Up: USD 1,500 in Ethereum. Conditional Prize (for top 10 teams): USD 2,500 in USDAO tokens, only for continuing to work on solutions post hackathon.	
Prizes by DOCK	
Winner: USD 2,500 in DOCK tokens. Runner-Up: USD 1,500 in DOCK tokens. 2nd Runner-Up (for top 10 teams): USD 1,000 in DOCK tokens.	
NOTE: Request you to Nominate a SPOC from your college for Hackathon and share their details	
Attached is the hackathon ppt for a better understanding. We do have mentors, but they will only be mentoring virtually on blockchain technologies.	
Respected Sir/Madam,	16.11.2021
Greetings from TASK!!	
	 Winner: USD 2,500 in Ethereum. Runner-Up: USD 1,500 in Ethereum. Gnditional Prize (for top 10 teams): USD 2,500 in USDAO tokens, only for continuing to work on solutions post hackathon. Prizes by DOCK Winner: USD 2,500 in DOCK tokens. Runner-Up: USD 1,500 in DOCK tokens. Autanner-Up (for top 10 teams): USD 1,000 in DOCK tokens. NOTE: Request you to Nominate a SPOC from your college for Hackathon and share their details Attached is the hackathon ppt for a better understanding. We do have mentors, but they will only be mentoring virtually on blockchain technologies. Respected Sir/Madam,

	I take this opportunity to thank you for your continuous support to TASK. It has been striving to enhance the employability quotient of the youth and enabled them choose multiple avenues as they graduate.	
	We are glad to inform you that TASK is opened student registrations across the state for the Academic year 2021-22 for the students who are pursuing B.E / B.Tech / B.Pharmacy / BA / B.Sc / B.Com / BCA / BBM / BBA / MBA / MCA / MA / M.Com / M.Sc / Polytechnic from 15th Nov 2021 to 31st Dec 2021.	
	Kindly communicate to the students and encourage them to register in below link:	
	Link to register: https://tasklms.telangana.gov.in/lms/login/signup_process. php	
	Note: Attached Student Registration Notification FY 2021- 22.	
TASK-Oracle Cloud Infrastructure(O CI) 7th Dec 2021	Dear Sir/Mam,	11.11.2021
	TASK in association with Oracle Academy is coming up with Oracle Cloud Infrastructure(OCI) event is on 7th December 2021 at 11 AM to 12:30 PM.	
	Oracle is accepting only 60 faculty nominations from 20 colleges and the faculty who have Oracle MEMBER HUB access only can attend this event.	

	About the Event:	
	Oracle Cloud Infrastructure (OCI) leads cloud computing with a deep and broad platform of cloud services that enables customers to build and run a wide range of applications in a scalable, secure, highly available and high- performance environment.	
	 Throughout the course, learners gain an understanding of: the core infrastructure of cloud how it works with databases cloud security, administration, monitoring, and management 	
	who are eligible?	
	Faculty members who have the access of Oracle MEMBER HUB Faculty who are having institution domain id Request you to please inform the faculty members who have member hub access to give their nomination in below link on or before 15th November 2021.	
	https://forms.gle/j9rZ25FVaW9wacdv6	
	Please reach out to me for any queries.	
TASK - NRSC Industry Visit	Dear Sir/Madam,	28.10.2021

Greetings From TASK!	
TASK in Collaboration with National Remote Sensing Centre (NRSC) Indian Space Research Organisation (ISRO) offering Industry Visit to all the TASK registered Colleges.	
About NRSC:	
National Remote Sensing Centre (NRSC) is one of the primary centres of Indian Space Research Organisation (ISRO), Department of Space (DOS). NRSC has the mandate for the establishment of ground stations for receiving satellite data, generation of data products, dissemination to the users, development of techniques for remote sensing applications including disaster management support, geospatial services for good governance and capacity building for professionals, faculty and students.	
Who Can Visit: Any Students from all Streams/Branch of Engineering/Degree &PG / MBA / MCA	
When: Every Friday except holidays	
Timings: 10:00 AM to 12:00 PM	
No. of Students: 200 - 250 maximum per visit	

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	Note: Students coming for ISRO (Industry Visit) Should be wearing formals & follow the covid norms (mask, sanitiser)	
	kindly share the student's data a week prior so that we can share it with NRSC Team as we prefer first come first serve basis.	
5G Empower for	Dear Sir,	21.10.2021
the youth of Telangana!		21.10.2021
	Greetings from TASK!	
	STL Academy is ready to train the students on 5G training. Please find below instructions.	
	I would like to request each college please download the template and enter the details, who are interested to enroll for 5G Training.	
	To upload bulk students please follow the following steps:	
	Step 1 Visit the website https://5g.stlacademy.tech/ and click on Upload button in the top right corner of the menu.	
	Step 2 Download the CSV file template by clicking on the Click here link. Fill the student database as per the template format.	
L		

	Step 3 On the same page input your institute / college name, contact person name, contact number and upload the file template.	
	Dear Sir,	19.10.2021
Infosys Off- Campus Recruitment Program: Invitation to participate in BCA and BSc graduates hiring	Please find below email from Infosys regarding Infosys off Campus Recruitment Program for BCA and BSc students from 2019,2020 and 2021 batches. Attached is the eligibility criteria for your reference.	
	Eligible students who wish to participate in the program can do so by filling out the Infosys Off-Campus Recruitment Program Application Form latest by Sunday, November 7, 2021.	
	Please refer to the email from Infosys reg Online test Pattern and other information.	
	Infosys Campus Recruitment Program will be conducted online in a phased manner comprising of Infosys online test and virtual interview. Candidates who clear Infosys online test will progress to the interview round.	
	Please be aware that we will not have any visibility regarding the applied students details, shortlisted student details , Interview schedule , selected student details or DOJ As the students would apply using link provided by Infosys	

	Academic Year 2021-22 for your reference.	
Infosys Campus	For any queries please feel free to contact me. Dear Sir/Madam,	01.10.2021
Recruitment		
Program:		
Operations		
Executive role	Infosys online test for Operation Executive is scheduled at 3PM on Sunday, October 3.	
	Candidates must carefully go through the attached Infosys	
	online test guidelines and rules to adhere to the mandatory instructions. If any malpractice or violation of instructions is detected during the test, candidate's application will be cancelled.	
	Students can access the sample test paper here to prepare for the online test.	
	Candidates who clear Infosys online test will proceed to the virtual interview round.	
	Request you to share the information with students.	
TASK - Infosys	Dear Sir/Madam,	01.10.2021
BPM Online		
FDP from 8th to		
19th Nov	Greetings from TASK!	
2021(Batch2)		

We are happy to initiate the 2nd batch of Infosys BPM online Faculty Development Program from 8th to 19th November 2021.

Infosys will deliver Global Business Foundation Skills curriculum which is aimed at bridging this gap between course-curriculum at graduate schools and the industry requirements. Our target audience is faculty members in English and Mathematics, who handle undergraduate students from streams like BA, BCom, BSc, BBA, BCA. To ensure that we have a multiplier effect we follow the train the trainer methodology. We use the adult learning methodology for training the faculty members from streams like BA, BCom. BSc, BBA, BCA. The Broad categories of the curriculum that would be covered includes – Facilitation Skills, Language Enhancement/Communication (Spoken and Written), Email Etiquette, Interview Skills, Analytical Skills (Creative thinking, introduction to management tools), Quantitative Aptitude, Corporate Etiquette etc. This training is conducted free of cost for the teachers under the banner of Infosys BPM -Corporate Social Responsibility. Our expectation is that each teacher will further train minimum 100 students.

Request you to share faculty details in attached format on or before 7th Oct 2021. The nominated faculty should be committed to make a difference in the student community.

Below are the details of the three modules that are covered as part of the program.

Track

	Participants with 100% attendance only will be given participatory certificates. No exceptions will be allowed	
	Looking forward to a successful Batch!	
	Please feel free to reach out to me for any clarifications and support.	
Infosys Springboard	Dear Sir/Madam,	27.09.2021
	Thanks for your interest for Springboard, Please share the students list in attached format.	
	Sharing an overview of Infosys new initiative – Infosys Springboard:	
	Why Infosys Springboard?	
	Infosys as part of our ESG (Environment, Social and Governance) vision has launched a new "Digital literacy Initiative", enabling Digital Talent at scale, to empower people, communities, and society. Through this initiative, Infosys plans to enable over 10 million people with digital and life skills by 2025. Towards this, we have launched the digital learning platform, Infosys Springboard in March 2021.	
	Who can leverage Infosys Springboard?	
	Students across India in the age group of 10 - 22 years as well as lifelong learners who are from government, Semi government or private institutions can benefit from Infosys Springboard. It caters to students from all streams not just Engineering and science. We already have ~400,000 students registered on Infosys Springboard.	

What content is available on Infosys Springboard?	
Infosys Springboard includes learning content developed by Infosys and leading content providers, spanning across digital and emerging technologies and life skills which are needed for students at all levels and streams. For a holistic learning experience, the platform has technology and soft skills playgrounds, programming challenges, and social learning features.	
300+ schools/colleges have started using this content for their students.	
Salient Features:	
Infosys Springboard is available free of cost for all learners in India including Govt and Private schools and NGOs. All the cost is taken by Infosys as part of our CSR initiative. Infosys Springboard is powered by Infosys Wingspan®, an integrated digital learning and collaboration platform which is a cloud hosted multitenant platform made available on Mobiles, tablets and Laptop/Desktop computers. Learners can learn relevant content on device of their choice anytime and anywhere.	
· Cloud hosted App – Wider accessibility for anywhere	
• Android and IOS enabled – Access Anytime, Anywhere, Any Device	
• Authoring Access – Institutes/schools can create content of their choice/assessments on the platform	
• Telemetry for all levels - Realtime updates on student learning data	
• Playgrounds – Ready to use technical and soft skills playground for practice, no pre-installation required	
· Assessments and Certifications – Video proctored	

certifications	
• Microsites –Customized microsites for educational institutions	
• Cohorts based learning – Learning with each other, discussion forums for learners	
• Multiple language – Already available in three languages – English, Hindi and Marathi. Planned to be made available in 13 major Indian languages by end of year. Content is also being made available in local languages.	
· Infosys Masterclass and student sessions - On demand as well as per planned calendar	
· Sessions with experts to discuss industry trends	
· Free conferencing app	
• Opportunity to participate in hackathon	
What is in for educational institutions:	
Customized Microsite for you	
Ability to create courseware (and learning path) combining content available on portal and your own customized	
content Playground for hands on practice for students: no need to worry about licenses, hardware, downloads, configuration etc	
Zero cost hosting of content	
Telemetry Access for your students	
Authoring Access for Faculty members – Faculty members can create their own content and assessments and host it on Springboard	
Masterclasses by Infosys SMEs – Live and interactive sessions for students. No additional overhead for teachers	
Link to platform	

	Infosys Springboard can be accessed at	
	https://infyspringboard.onwingspan.com/	
	Institutes can share details for bulk registration(details	
	attached)	
TASK-"C2C -	Dear Sir,	21.09.2021
Campus to		
Career"		
	Greetings from Telangana Academy of Skills and	
	Knowledge (TASK)	
	We are happy to announce, yet another key initiative	
	launched by our Industry Partner CSS Corp – which is	
	branded as "C2C - Campus to Career" for the benefit of	
	TASK registered institutions across Telangana.	
	Through this Academic – Industry partnership, academic	
	institutions are welcome to explore opportunities to work	
	very closely with CSS Corp in the areas of Virtual	
	Internship, Curriculum Integration, Setting up Incubation	
	Center & Creating employment opportunities for your	
	students.	
	Please click on the below link to know more about CSS	
	Corp's "C2C – Campus to Career	
	https://www.task.telangana.gov.in/Announcements/TASK_	
	CSS	
	Interested college please do confirm your interest by	
	writing mail to Campus.career@csscorp.com on or before	
	21st September 2021.	
		I

	We invite your esteemed institution to be a part of this new initiative and make Campus to Career a reality for your students.	
Infosys Springboard - New Digital Literacy Initiative	Dear Sir/Madam, Sharing an overview of Infosys new initiative – Infosys Springboard:	20.09.2021
	Why Infosys Springboard? Infosys as part of our ESG (Environment, Social and Governance) vision has launched a new "Digital literacy Initiative", enabling Digital Talent at scale, to empower people, communities, and society. Through this initiative, Infosys plans to enable over 10 million people with digital and life skills by 2025. Towards this, we have launched the digital learning platform, Infosys Springboard in March 2021.	
	Who can leverage Infosys Springboard? Students across India in the age group of 10 - 22 years as well as lifelong learners who are from government, Semi government or private institutions can benefit from Infosys Springboard. It caters to students from all streams not just Engineering and science. We already have ~400,000 students registered on Infosys Springboard.	
	What content is available on Infosys Springboard? Infosys Springboard includes learning content developed	

by Infosys and leading content providers, spanning across digital and emerging technologies and life skills which are needed for students at all levels and streams. For a holistic learning experience, the platform has technology and soft skills playgrounds, programming challenges, and social learning features. 300+ schools/colleges have started using this content for their students.	
Salient Features:	
Infosys Springboard is available free of cost for all learners in India including Govt and Private schools and NGOs. All the cost is taken by Infosys as part of our CSR initiative. Infosys Springboard is powered by Infosys Wingspan®, an integrated digital learning and collaboration platform which is a cloud hosted multitenant platform made available on Mobiles, tablets and Laptop/Desktop computers. Learners can learn relevant content on device of their choice anytime and anywhere.	
· Cloud hosted App – Wider accessibility for anywhere	
 Android and IOS enabled – Access Anytime, Anywhere, Any Device 	
• Authoring Access – Institutes/schools can create content of their choice/assessments on the platform	
• Telemetry for all levels - Realtime updates on student learning data	
 Playgrounds – Ready to use technical and soft skills playground for practice, no pre-installation required 	
• Assessments and Certifications – Video proctored certifications	
• Microsites –Customized microsites for educational institutions	

• Cohorts based learning – Learning with each other, discussion forums for learners	
• Multiple language – Already available in three languages – English, Hindi and Marathi. Planned to be made available in 13 major Indian languages by end of year. Content is also being made available in local languages.	
 Infosys Masterclass and student sessions - On demand as well as per planned calendar 	
· Sessions with experts to discuss industry trends	
· Free conferencing app	
• Opportunity to participate in hackathon	
What is in for educational institutions:	
Customized Microsite for you Ability to create courseware (and learning path) combining content available on portal and your own customized content	
Playground for hands on practice for students: no need to worry about licenses, hardware, downloads, configuration etc	
Zero cost hosting of content Telemetry Access for your students	
Authoring Access for Faculty members – Faculty members can create their own content and assessments and host it on Springboard	
Masterclasses by Infosys SMEs – Live and interactive sessions for students. No additional overhead for teachers	
Link to platform	
Infosys Springboard can be accessed at https://infyspringboard.onwingspan.com/	
Institutes can share details for bulk registration(details	

	attached)	
TASK - Amazon Job Requirement for 2020 and 2021 graduates	Dear Sir/Madam,	16.09.2021
	Greetings from TASK!	
	Amazon JD is published on TASK portal.	
	B.E/B.Tech, B. Sc, B.Com, BBA, BBM, B.A, BCA (any discipline) 2020/2021 graduates with 55% aggregate throughout the academics and no backlogs are eligible to apply	
	Graduates from 2021 batch can apply using below link	
	https://www.task.telangana.gov.in/Placements/Amazon_Development_Centre_India_Pvt_Ltd_2021	
	Graduates from 2020 batch can apply using below link	
	https://www.task.telangana.gov.in/Placements/Amazon_Development_Centre_India_Pvt_Ltd_2020	
	As its urgent requirement last date to apply is 17th September 2021.	
	Applied students would receive communication from amazon regarding the online assessment post last date to apply.	

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	Applied students are expected to check their emails from Amazon for the assessment link and also need to check in spam.	
	Request you to encourage eligible and interested students to apply on TASK portal.	
Infosys Campus Recruitment Program(Job Opening): Invitation to participate in	Dear Sir/Madam, Greetings from TASK!	16.09.2021
BCA and BSc graduates hiring	Please find below email from Infosys regarding Infosys Campus Recruitment Program 2021-22. BCA and BSc students from 2020, 2021 and 2022 batches.	
	Attached is the eligibility criteria for your reference.	
	To apply Interested and eligible students need to fill in their details in the link provided in the below email. Please refer to the email from Infosys reg Online test Pattern and other information Last Date to apply Sunday, September 19, 2021.	
TASK-"C2C - Campus to Career"	Dear Sir / Madam,	16.09.2021
	Greetings from Telangana Academy of Skills and Knowledge (TASK)	

	We are happy to announce, yet another key initiative launched by our Industry Partner CSS Corp – which is branded as "C2C - Campus to Career" for the benefit of TASK registered institutions across Telangana. Through this Academic – Industry partnership, academic institutions are welcome to explore opportunities to work very closely with CSS Corp in the areas of Virtual Internship, Curriculum Integration, Setting up Incubation Center & Creating employment opportunities for your students. Please click on the below link to know more about CSS Corp's "C2C – Campus to Career	
	https://www.task.telangana.gov.in/Announcements/TASK_ CSS	
	Interested college please do confirm your interest by writing mail to Campus.career@csscorp.com on or before 24th September 2021.	
	We invite your esteemed institution to be a part of this new initiative and make Campus to Career a reality for your students.	
TASK - Mentor Together - TPO Meeting Invite	Dear Sir,	16.09.2021

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	<u>TASK</u> in collaboration with <u>Mentor Together</u> , India's first ar focused mentorship non-profit organization is providing a r - Mentor To Go to all students in Telangana. (<u>https://www.task.telangana.gov.in/Partners/Mentor_Togeth</u>	nobile mento	
	In this regard, TASK is inviting all the TPOs from its partner webinar to learn about 'Mentor To Go' and extend the progr	U 1	-
	Mentor To Go is an android and web-bas students improve their work readiness th sessions, and personalised mentorship pr professionals from a range of careers. Me provide career mentorship to young peop across the length and breadth of India.	rough self-lea ovided by ind ntor to Go's v	arning dustry vision is t
Infosys Campus Recruitment Program(Job	Dear Sir/Madam,	15.09.2021	
Opening): Invitation to participate in BCA and BSc	Greetings from TASK!		
graduates hiring	Please find below email from Infosys regarding Infosys Campus Recruitment Program 2021-22. BCA and BSc students from 2020, 2021 and 2022 batches.		
	Attached is the eligibility criteria for your reference.		
	To apply Interested and eligible students need to fill in their details in the link provided in the below email. Please refer to the email from Infosys reg Online test Pattern and other information Last Date to apply Sunday, September 19, 2021.		
Student to Entrepreneurs "Angadi" - Registrations	Dear Sir/Madam,	14.09.2021	

Greetings from TASK - i4TS - Mandi.	
i4TS Concept was initiated by TASK under the key mission of Entrepreneurship, and its inception i4TS exploring social entrepreneurial mindset development in the state of Telangana.	
We are introducing a unique concept "Angadi" A field marketing event in the state as a tribute to Bharat Ratna Shri Mokshagundam Visvesvaraya's birth anniversary on 15th September 2021, and closing on Mahatma Gandhi birth anniversary.	
Why this program?	
With a motto to develop an entrepreneurial mind-set, enterprise development among graduating students, and youth of Telangana state.	
What will be done POST to the field experiment?	
Post to the field experiment, students will be made to reflect on their field experience so as to reconfigure the field learnings. Experience sharing, learning possibilities. Students may be asked to make presentations, videos of their experiences. Learning presentations will be appreciated and provided with certificates or prizes.	
How the Program will work out?	
TASK Will procure the engineering merchandise in bulk & fixes a price to the profit to the student and distributed it to the CMs/RMs as per their targets and handover to the colleges to sell to their students.	

	After completion of the selling, students will be asked to fill a questionnaire based on the learning outcomes.	
	a questionnaire based on the learning outcomes.	
	The purchase amount will be collected back from the colleges and handover to the TASK.	
	Based on the learning outcomes, the responses will be evaluated and awarded by TASK.	
	Edibility Criteria: Youth between 18-25 years (Engineering/Degree/Diploma) belonging to Telangana State.	
	Last date to apply: 20th September 2021.	
	Registration link: t.ly/rziDq	
	Please let me know if you have any questions or need additional information. Dear Sir/Madam,	07.09.2021
i4TS TASK - Mandi Webinar-		07.07. L ULI
3 on 8th September 2021	Telangana Academy for Skill and Knowledge (TASK) is a not-for-profit organization initiated by the government of Telangana in 2014 to build synergy between academia,	
from 4 PM to 6 PM	industry, and government.	
	The organization has three fundamental focus areas:1.	

Education, 2. Employability, & 3. Entrepreneurship.	
i4TS (Innovation for Telangana State) is a flagship program under the Entrepreneurships of TASK.	
We are glad to inform you that our initiative i4TS (Innovating for Telangana State) - Samajika AavishkaranaluSaralamaina Jeevithaniki (సామజిక ఆవిష్కరణలుసరళమైన జీవితానికి) is progressing well.	
Under the program of i4TS, TASK planned a webinar called i4TS TASK - Mandi on 8th September 2021, 4 PM to 6 PM.	
Speaker-1:	
Dr. Prasad Teegalapelly, Faculty Adviser, Center for Student Enterprises & Associate Professor of Organizational Behavior, National Institute of Industrial Engineering (NITIE) Mumbai, India.	
Over the last 15 years, he has managed numerous competitively funded basic research and Institute-industry collaborative projects. In the latter capacity, he has worked in collaboration with a number of Indian companies and government departments to assist in organizational redesign and improvements, and management development. These organizations include Reliance, BEL, ITI Bangalore, Zensar, Satyam, Kribhco, etc.	
About Speaker success story: 1. https://www.youtube.com/watch?v=dl8X7QD0mIo; 2. https://youtu.be/sUkLx0sd1h0; 3. https://youtu.be/dl8X7QD0mIo;	

		,
	joining link: https://task.radiusedutech.com Time: 04.00 AM to 6.00 PM.	
	Topic: The art of selling (అమ్మడమనే కళ) Who can attend: Youth between 18 to 25 years.	
	Kindly share with all students.	
i4TS TASK - Mandi Webinar- 2 on 1st	Dear Sir/Madam,	31.08.2021
September 2021 from 4 PM to 6 PM.	Greetings from Telangana Academy for Skill and Knowledge (TASK)!	
	TASK is a not-for-profit organization initiated by the government of Telangana in 2014 to build synergy between academia, industry, and government.	
	The vision and mission of TASK are to empower youth so that they transform into quality assets to the industry.	
	The organization has three fundamental focus areas:1. Education, 2. Employability, & 3. Entrepreneurship.	
	i4TS (Innovation for Telangana State) is a flagship program under the Entrepreneurships of TASK.	
	We are glad to inform you that our initiative i4TS	

(Innovating for Telangana State) - Samajika AavishkaranaluSaralamaina Jeevithaniki is progressing well.	
Under the program of i4TS, TASK planned a webinar called i4TS TASK - Mandi on 1st September 2021, 4 PM to 6 PM.	
Speaker-1:	
Dr. Prasad Teegalapelly, Faculty Adviser, Center for Student Enterprises & Associate Professor of Organizational Behavior, National Institute of Industrial Engineering (NITIE) Mumbai, India.	
Over the last 15 years, he has managed numerous competitively funded basic research and Institute-industry collaborative projects. In the latter capacity, he has worked in collaboration with a number of Indian companies and government departments to assist in organizational redesign and improvements, and management development. These organizations include Reliance, BEL, ITI Bangalore, Zensar, Satyam, Kribhco, etc.	
About Speaker success story: 1. https://www.youtube.com/watch?v=dl8X7QD0mIo; 2. https://youtu.be/sUkLx0sd1h0; 3. https://youtu.be/dl8X7QD0mIo;	
joining link: https://task.radiusedutech.com	
Time: 04.00 AM to 6.00 PM.	
Topic: Registering a Company (for entrepreneurs)	

	Who can attend: Youth between 18 to 25 years.	
Google -	Dear Sir/Madam,	25.08.2021
Android Development	Creatings from TASKI	
with Kotlin TTT	Greetings from TASK!	
Nomination		
Form		
	We bring you our new 'Android Development with Kotlin'	
	course which will empower and enable your faculty to	
	teach Android development in Kotlin to their students in a classroom/online lecture format and through hands-on	
	practical labs.	
	We are offering an online 'Train The Trainer' session to faculty.	
	faculty.	
	Training Model	
	No. of Days	
	Duration	
	No. of Participants	
	Online ILT Model	
	3	
	7 hours	
	30 (Per Batch)	
	For now, you can share your nominations on the basis of	
	the below-mentioned prerequisites and specifications	
	required for the course.	

	Technical Know-How: Participants taking the course must have Object-oriented programming experience (eg Java, .NET, etc).	
	Machine Hardware Specifications required for this course (If college or university can also provide this specification for faculty in their labs would be appreciated) :	
	Processor: i3 or Higher	
	RAM: 8GB or higher	
	Virtualization Technology - enabled	
	Operating System: Windows 10/ Macintosh/ Linux	
	Internet: Good Internet speed (recommended 10 MBPS at least)	
	Kindly share the below link with all the interested faculty members of your institution for joining the session,	
	Click here for nomination: Google - Android Development with Kotlin TTT Nomination Form	
	We appreciate your involvement and look forward to active participation and hope for more responses from your side. Attached Program PDF for ready reference.	
Batch - 11 TALLY Training from 1st Sept to	Dear Sir/Madam.	25.08.2021
6th Sept 21 and Time: 2:00 PM to 4:30 PM	Greetings from TASK!	

We are offering the 11th batch of TALLY online training to the TASK registered students of B.com (1st, 2nd, 3rd years), M.com and MBA (1st, 2nd years). It is a five-day program and free of cost for the TASK registered students.	
TALLY Training details:	
Date: 1st, September-2021 to 6th, September-2021 Time: 2:00 PM to 4:30 PM	
Student Registration Link:	
https://forms.gle/ERBkjcDXKgTSmCoc9	
Last date for closing the registrations is 27th, August - 2021 by 4:30 PM	
Requesting to share the above link with the students.	
Note:	

	We are providing 2 certificates based on the following eligibility criteria 1). Participation Certificate - Students must attend all 5 days of training	
	2). Assessment Certificate - Students must attend all 5 days of training and also have to complete the assessment with a minimum of 60% marks. Assessment will be on the last day of the training.	
Crux Services -	Dear Sir/Madam,	25.08.2021
New Job Requirement for 2020 and 2021 graduates.	Please find the link of new requirement for 2020 and 2021 graduates.	
	Company name - Crux Services	
	JD Link for 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/Crux_servic es_2020	
	JD Link for 2021 pass out students:	
	https://www.task.telangana.gov.in/Placements/Crux_servic es_2021	

	Request you to encourage the eligible students to apply for the above job posting.	
KIMS Hospitals - New Job Requirement for 2020 and 2021 graduates	Dear Sir/Madam, Please find the link of new requirement for 2020 and 2021 graduates.	23.08.2021
	Company name - KIMS Hospitals	
	JD Link for 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/KIMS_Hosp itals_Trainee	
	JD Link for 2021 pass out students:	
	https://www.task.telangana.gov.in/Placements/KIMS_Hosp itals_Trainee_2021	

the above job posting.	
Dear Sir/Madam,	23.08.2021
Greetings from TASK!	
Deloitte has come up with requirement for Degree students (Non Engineering) from 2021 batch. As a pilot students from Hyderabad, Rangareddy, Medchal and Sangareddy are eligible to apply	
Profile: Associate Analyst – Deloitte Consulting	
· Eligibility Criteria: BSc- (Mathematics, Statistics, Physics, Electronics) with Computer Science	
· BCA	
· 2021 Graduates only	
• No active backlogs	
• Aggregate 60% or CGPA of 6.5	
 Please refer to the attachment for the details of the profile being offered. The compensation set for the requisite role is Rs.3,57,500 Please ensure all the students have: Laptop/desktop with functional web-cam capability Stable and secure internet connectivity 	
) ((1 (1 (1) (1) (1) (1) () (Dear Sir/Madam, Greetings from TASK! Deloitte has come up with requirement for Degree students (Non Engineering) from 2021 batch. As a pilot students from Hyderabad, Rangareddy,Medchal and Sangareddy are eligible to apply Profile: Associate Analyst – Deloitte Consulting Eligibility Criteria: BSc- (Mathematics, Statistics, Physics, Electronics) with Computer Science BCA 2021 Graduates only No active backlogs Aggregate 60% or CGPA of 6.5 Please refer to the attachment for the details of the profile being offered. The compensation set for the requisite role is Rs.3,57,500 Please ensure all the students have: 1. Laptop/desktop with functional web-cam capability

	link https://forms.gle/wGEFUKcxBwucifAf7	
	Last date to apply is 11:00 am 28th August 2021.	
	Please Note : All the applied student details would be shared with Deloitte and Deloitte based on their discretion would shortlist the students.	
	All shortlisted students only would receive online test Invite from Deloitte assessment vendor.	
[24]7.ai - New Job Requirement	Dear Sir/Madam,	19.08.2021
for 2021 graduates.	Please find the link of new requirement for 2021 graduates.	
	Company name - [24]7.ai	
	JD Link for 2021 pass out students:	
	https://www.task.telangana.gov.in/Placements/24_7ai_dig_ adv	
	Request you to encourage the eligible students to apply for the above job posting.	

TekFriday Processing Solutions Pvt	Dear Sir/Madam,	17.08.2021
Ltd - Job requirement for 2020 graduates	Greetings from TASK!	
	Please find the link of new requirement for 2020 graduates.	
	Company name - TekFriday Processing Solutions Pvt Ltd	
	JD Link for 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/TekFriday_ Processing_Solutions_Pvt_Ltd	
	Request you to encourage the eligible students to apply for the above job posting.	
TASK Unschool - A Series of Free	Dear Sir/Madam,	17.08.2021
Webinars	Greetings from TASK	
	TASK in association with Unschool is coming up with free webinars for TASK registered Students.	

Please find below the list of webinars that we wish to conduct in the month of August:	
Date	
Timing	
Category	
Proposed Topics	
18.08.2021	
18:00 - 19:00	
Personal Development	
Preparing yourself for an interview	
21.08.2021	
18:00 - 19:00	
Coding & Development	
NLP- Build your own Chabot	
22.08.2021	
18:00 - 19:00	
Software & Technology	
Cybersecurity as a Career	
28.08.2021	
18:00 - 19:00	
Software & Technology	
Machine Learning using Python	

	Registration link to register for above webinars:	
	http://unschool-7019118.hs-sites.com/webx	
	Kindly share it with TASK students to avail this opportunity.	
	Registration is open till 18th August 2021 afternoon for the first webinar " Preparing yourself for an interview".	
Batch - 10 TALLY Training Schedule : 24th, August-2021 to 28th, August- 2021 and Time: 2:00 PM to 4:30	Dear Sir/Madam, Greetings from TASK!	17.08.2021
PM	We would like to offer the 10th batch of TALLY online training to the TASK registered students of B.com (1st, 2nd, 3rd years), M.com and MBA (1st, 2nd years). It is a five- day program and free of cost for the TASK registered students.	
	TALLY Training details:	
	Date: 24th, August-2021 to 28th, August-2021	

Time: 2:00 PM to 4:30 PM	
Student Registration Link:	
https://forms.gle/3Rv7nwUDWTkZFSBDA	
Last date for closing the registrations is 20th, August - 2021 by 2:00 PM	
Requesting to share the above link with the students.	
For any further clarifications, please feel free to contact me.	
Note:	
We are providing 2 certificates based on the following eligibility criteria	
1). Participation Certificate - Students must attend all 5	

	days of training 2). Assessment Certificate - Students must attend all 5 days	
	of training and also have to complete the assessment with a minimum of 60% marks. Assessment will be on the last day of the training.	
First American (India) Pvt Ltd Job Requirement for 2020	Dear Sir/Madam,	16.08.2021
graduates.	Greetings from TASK!	
	Please find the link of new requirement for 2020 graduates.	
	Company name - First American (India) Pvt Ltd	
	JD Link for 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/First_Ameri can_India_Pvt_Ltd_2020	
	Request you to encourage the eligible students to apply for the above job posting.	
New Job Requirement for	Dear Sir/Madam,	16.08.2021

2020 graduates.		
	Greetings from TASK!	
	Please find the link of new job requirement for 2020 graduates.	
	Company name - Genpact	
	JD Link for 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/Genpact_A pprentice	
	Request you to encourage the eligible students to apply for the above job posting.	
NASSCOM//Cyb er security Centre of	Dear Sir/Madam,	12.08.2021
Excellence//MO NTH-LONG PROGRAM on "APP SECURITY	Greetings from TASK!	
CHAMPION"	Please find the below mailer about "APP SECURITY CHAMPION"	
	Students being in exams, we did a soft (low-key) kickoff	

with a few students mainly to record the session which explains how to use the platform, so that the subsequent batches can watch the instruction video. We'll have a grand finale!	
MONTH-LONG PROGRAM as be extended their students registration dates, Please find the details below Dear Madam.	12.08.2021
	1_1001_0_1
Greetings from TASK!	
Deloitte has come up with requirement for students from 2019,2020,2021 batches.	
Designation : Intern Analyst	
B.Tech/ B.E. – Any branch and MCA - 2019, 2020, 2021 Graduates only	
Students should be certified or trained in any of the technologies.	
No active backlogs	
Aggregate 60% or CGPA of 6.5	
	explains how to use the platform, so that the subsequent batches can watch the instruction video. We'll have a grand finale! MONTH-LONG PROGRAM as be extended their students registration dates, Please find the details below Dear Madam, Greetings from TASK! Deloitte has come up with requirement for students from 2019,2020,2021 batches. Designation : Intern Analyst B.Tech/ B.E. – Any branch and MCA - 2019, 2020, 2021 Graduates only Students should be certified or trained in any of the technologies. No active backlogs

	Probation Period: 1 year	
	The stipend for the requisite role INR 2.5 L per annum and INR 70,000 joining bonus	
	Please Note: Students from Colleges Where Deloitte has already visited for campus 2021 are not be eligible to apply.	
	Request you to share the link to the colleges encourage TASK registered students to apply using the link https://forms.gle/iDxLh9BFCpjDgsr5A	
	Last Date to apply 11:00 am 17th August 2021	
AIR Worldwide Corporation Job	Dear Sir/Madam,	07.08.2021
Requirement for 2021 graduates	Please find the link of new requirement for 2021 graduates.	
	Company name - AIR Worldwide Corporation (AIR)	
	JD Link for 2021 pass out students:	
	https://www.task.telangana.gov.in/Placements/AIR_World wide_Corporation_2021	

	Request you to encourage the eligible students to apply for the above job posting.	
Launching of TASK – UNICEF - YuWaab, Young	Dear Sir/Madam,	07.08.2021
YuWaah_Young Warrior Movement to Combat COVID- 19	Greetings from TASK!	
	TASK has served Telangana state in many ways since its inception and you are all the front-line warriors to all its services.	
	Now we are in a position to take up a real challenge of "TASK - Young Warrior" A youth movement to combat COVID-19.	
	The overwhelming of the world about Corona Virus outreach with various variants and waves. Coronavirus is here for the long haul and scientists predict for the next 10 to 20 years the situation remains the same.	
	Keeping this adventure, we have to stay safe and informed by a trusted source and trained volunteers to prevent this situation.	
	We are excited to announce that, a collaborative initiative of Youth-led Movement to Combat COVID-19 between TASK – UNICEF - YuWaah.	
	For launch of this partnership event in the presence of Shri Jayesh Ranjan, Principal Secretary, ITE&C Dept. Telangana, TASK and Officials from Government Departments, Senior	

Officials from UNICEF/YuWaah, TASK Alumni and participants from institutions including faculty and students.	
 Target Audience: Officials from Government Departments Faculty, Students and TASK Alumni 	
Date & Time of Launching the program: Will be informed after successful registration of the participants.	
Modalities of launching the program: - Virtual event and live broadcast on social media	
For more details about Young Warrior please go through the link -	
https://www.task.telangana.gov.in/Announcements/TASK_ Young_WarriorYouth- led_Movement_to_Combat_COVID-19	
Communicate the same with all the faculty to register for this unique program in below link in a massive way.	
Registration link - https://forms.gle/p3aLQvaZVYJqe5X37	

Invitation: Inaugural of HYSEA Internship Fair 2021 - Saturday, 7th August 2021 at 3.30pm	Dear Sir/Madam, Greetings from TASK!	06.08.2021
	You are cordially invited to the Inaugural of the first-ever HYSEA Internship Fair 2021 being organized in association with Tech Partner CONDUIRA and outreach partners JNTUH and TASK. Mr. V. Laxmikanth, MD, Broadridge India has kindly consented to be the Chief Guest on the occasion.	
	An important part of the Inaugural event will be a panel discussion by eminent Industry and Academic Leaders on 'Are we bridging the gap right?' in the context of Industry- Academia efforts towards creating Industry ready students. The panel will be moderated by Mr. Sesha M Rao, Board Member, Insideview Technologies & Leader – Industry- Academia Forum – HYSEA and will consist of eminent personalities from the Industry and the Academia.	
	The panel members include: Mr. Vinay Agrawal, Global Head - Business HR, Tech Mahindra Ms. Manisha Saboo, AVP & Delivery Head and Head of Pocharam, Hyd and Indore Campuses, Infosys Dr. A. Govardhan, Professor & Rector, JNTUH Mr. Shrikant Sinha, CEO, TASK Mr. Prakash Yalla Veera, Technology Transfer Officer & Head - Product Labs, IIIT-H	

		[]
	Day & Date: Saturday, 7th August 2021 Time: 3.30pm to 5pm Zoom Registration Link: https://us02web.zoom.us/webinar/register/WN_D3f_kaXqQ LqND9w4SPGViw	
	Look forward to your participation.	
TASK- Times Jobs (The Times Group) free aptitude test //A gateway for your first job - TimesJobs Pro	Dear Sir/Ma'am, Greetings from TASK!	06.08.2021
	Here is an exclusive opportunity, TASK in association with Times Jobs (Times Group) for your students from Arts & Science Branches. We are conducting a free aptitude test for your students, the hiring companies associated with us will get the pre-assessed data of your students along with their resumes for the recruitment process. We will be sharing the details with the best companies in the market.	
	Aptitude Round:	
	 25 - 30 Aptitude Questions. Timing will be around 40-45 Mins. 	
	3. Students will have 2 attempts.	

	 Process: o Students are requested to register in the link below. o Once they are registered, we will share the link for Aptitude Assessment. o After the completion of the assessment, we will share the results with you. 	
	o We will share the assessment results along with their resumes with the companies as per their requirement.	
	Request you to circulate the link to the candidates once they register, we will share the test link with them	
	https://www.timesjobs.com/candidate/register.html?outSou rce=Camptask	
Invitation TASK- Oracle Academy	Dear Sir/Mam,	29.07.2021
Virtual Faculty Day (10th Aug, 2021)	TASK in association with Oracle Academy is coming up with a virtual event on 10th Aug, 2021 at 11:00 AM IST to celebrate Oracle Academy Faculty Day. Oracle Academy Institutional members will present how they are using free resources and aligning them with their existing curriculum. They also will share how these resources have impacted their teaching and their students learning. Learn how Oracle Academy Institutional members are	
	using Oracle Academy resources for the benefit of their students. Key highlights of the event are: Event is open for faculty/management of the institutes	

	Certificate of attendance will be provided to attendees of the event who will attend at least 75% of the total duration of live event. Timing for Event: 11 AM IST to 1 PM IST Event is open for registration to Non- OA members as well.	
	Webinar prerequisites:	
	You must have access to a laptop/tablet compatible with Zoom.	
	Contact Oracle Academy for any issues prior to the start of the workshop.	
	Registration Link for Event -	
	https://go.oracle.com/LP=113938?elqCampaignId=298531&s rc1=:pp:ptr::::TASK	
	Other details regarding the event can be found on the registration page. Faculty members need to register for the event via above mentioned link.	
	Kindly share this information with TASK registered colleges and inform the faculty members to register for the event.	
Principals' and TPOs/TASK Coordinators Meet on 30th	Dear Sir/ Madam,	27.07.2021
July 2021 from 2:30 PM to 4:00 PM	Greetings from Telangana Academy for Skill and knowledge (TASK)!	
	Trust you and your family are healthy and staying safe.	

We are delighted to invite you to join us for the Principals' Meet 2021 (Virtual) scheduled on 30th July 2021 from 02:30 PM to 04:00 PM	
The virtual event will be presided over by Sri Navin Mittal, IAS, Commissioner of Collegiate Education	
The agenda of the meet includes talks on skilling initiatives, hiring scenario, skilling plans for the next academic year and an interactive session. The objective is to share and understand each other's points of view so that we can work together more cohesively for the benefit of the youth.	
Your feedback and review are essential for us to constantly improve our services and make them more relevant to your requirements.	
Kindly confirm your participation in advance by filling the below form on or before 28th July 2021 (5:00 PM)	
https://forms.gle/DThUYj4X9QRsvPF56	
After registering, you will receive a confirmation email containing the meeting joining link	
We look forward for your participation on the 30th July 2021	

TASK-NRSC- ISRO: Quiz on 75 years of India and ISRO	Dear Sir/Madam, Greetings from TASK!	25.07.2021
	We are glad to inform you that, TASK in association with NRSC-ISRO is organizing online Quiz competition on "Indian Independence and ISRO" based on pre Independence struggle, post independence development and major development in ISRO over these years.	
	Please find the attached flyer requesting you to share the information with all students. (It is an open quiz for TASK registered and non TASK registered students)	
	Following are the details:	
	Quiz Name: Azadi ka Amrut Mahotsav	
	Date of the Quiz: 29th July (Thursday)	
	Time of the Quiz: 5.00 PM	
	Eligibility: All branches of Engineering, Degree & P.G Colleges(pursuing)	
	Registration link and Mock test link: https://bhuvan- app1.nrsc.gov.in/quiz/home/index.php	

Start Date of Registration: 23.07.2021	
End date and time of Registration: 28.07.2021,2:00 pm	
Attractive prizes and certificates to the winners (conditions apply).	
How to register:	
• For all details and updates visit https://bhuvan- app1.nrsc.gov.in/quiz/home/index.php	
• Registration is open till 28th July (2:00 PM).	
• Registration can be done using the "REGISTER" link on the home page.	
• Note down the unique ID generated after successful registration and use the same for the actual quiz.	
• There is no fee for registration.	
Quiz Details:	
• A mock practice paper will be available on the Quiz home page "MOCK QUIZ"	
• No negative marking. Recommended using a desktop browser.	
• Quiz language - English	

	 Click on a question number to go to the question location on the Bhuvan map/image. The question can also be seen by clicking on the location tag on the map. Click on a location to get the question and options. Choose the correct option. There is only one correct option for each question. The button will remain highlighted. Do not press submit button after each answer. You can zoom/de-zoom the Bhuvan map/image to know nearby details. You can view Satellite data by selecting the 'Layer stack' icon on the map and selecting the Satellite option to know more details. Press "Finish Quiz" after completion of the quiz before the stipulated time ends. Attempt all the questions but a partial quiz can also be submitted. Only one submission per unique ID is allowed. Quiz time is from 17:00 Hrs to 17:20 Hrs. to attempt 25 questions. 	
	Dear Sir/Madam,	25.07.2021
KIA - Job Requirement for 2019 and 2020 graduates.	Please find the link of new requirement for 2019 and 2020 graduates.	
	Company name - CAR KIA	
	JD Link for 2019 pass out students:	

	https://www.task.telangana.gov.in/Placements/CAR_KIA_2 019	
	JD Link for 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/CAR_KIA_2 020	
	Request you to encourage the eligible students to apply for the above job posting.	
TASK - New Job Requirements	Dear Sir,	22.07.2021
for 2020 graduates	Please find the link of new requirements for 2020 graduates.	
	Company name - Matrimony.com Ltd	
	JD Link for 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/Matrimony. com_Ltd_2020_telesales_exe	
	Company name - IKEA India Pvt Ltd	

	JD Link for 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/Ikea_CSC_ Gen	
	Request you to encourage the eligible students to apply for the above job posting.	
Bharti Airtel - New Job	Dear Sir/Madam,	16.07.2021
Requirement for 2019 and 2020 graduates.	Greetings from TASK!	
	Please find the link of new requirement for 2019 and 2020 graduates.	
	Company name - Bharti Airtel	
	JD's Links:	
	For 2019 pass out students	
	https://www.task.telangana.gov.in/Placements/Bharti_Airte 1_2019	

	For 2020 pass out students	
	https://www.task.telangana.gov.in/Placements/Bharti_Airte 1_2020	
	Request you to encourage the eligible students to apply for the above job posting.	
TASK - Job requirement for 2019,2020 and	Dear Sir,	12.07.2021
2021 graduates at Tutoroot Technologies Pvt. Ltd	Please find the link of new requirement for 2019,2020 and 2021 graduates.	
	Company name - Tutoroot Technologies Pvt Ltd	
	JD Links:	
	For 2019 pass out students:	
	https://www.task.telangana.gov.in/Placements/Tutoroot_Te chnologies_Pvt_Ltd_2019	
	For 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/Tutoroot_Te chnologies_Pvt_Ltd_2020	
	For 2021 pass out students:	
	https://www.task.telangana.gov.in/Placements/Tutoroot_Te chnologies_Pvt_Ltd_2021	
	Request you to encourage the eligible students to apply for	

	the above job posting.	
TASK - Job Requirement for	Dear Sir/Madam,	12.07.2021
2019,2020 and 2021 graduates at Ninjacart	Greetings from TASK!	
	Please find the link of new requirement for 2019 and 2020 graduates.	
	Company name - NINJACART	
	JD Links:	
	For 2019 pass out students:	
	https://www.task.telangana.gov.in/Placements/NINJACAR T_sales_exe_2019	
	For 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/NINJACAR T_2020	
	Request you to encourage the eligible students to apply for the above job posting.	
New Jobs -	Dear Sir/Madam,	09.07.2021
Justdial &		
RACEWIN	Creetings from TASK!	
Tradelinks for	Greetings from TASK!	

students		
graduated in		
2020	Please find the link of new requirement for 2020 graduates.	
	Company name - Justdial	
	ID Link For 2020 page out students:	
	JD Link For 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/Justdial_202	
	Company name - RACEWIN TRADELINKS	
	JD Link For 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/RACEWIN_	
	TRADELINKS	
	Request you to encourage the eligible students to apply for	
	the above job posting.	
TASK - Job	Dear Sir/Madam.	07.07.2021
Requirement for		
students		
graduated 2020		
in Intralinks .	Greetings from TASK!	
	Please find the below Job requirement for 2020 graduates.	

Company name – Intralinks	
Domain: IT	
Designation: Associate Backoffice Specialist	
CTC Per Annum : 3,00,000 Per Annum	
Job Description:	
Responsible for completion of projects on Customer Success team.	
Complete Projects in a timely manner and with high quality.	
Confirm client requirements, manage internal customer expectations, and liaise with other internal teams as needed	
Eligibility Criteria:	
B.Sc,B.Com, BCA ,BA (any discipline) with minimum 65% aggregate throughout the academics graduated in 2020 and with no backlogs, need to apply.	
Bond Period : NA	

	Interview Procedure:	
	Excel Test	
	Technical rounds - 2	
	Manager round	
	HR Round	
	Should be willing to join immediately.	
	For more details please visit:	
	https://www.task.telangana.gov.in/Placements/Intralinks_2 020	
	Request you to encourage the eligible students to apply for the above job posting.	
FDP on Machine	Dear Sir,	06.07.2021
Learning on Cloud from 19th to 24th July 2021	Greetings from TASK!	
	TASK in collaboration with 360Digi TMG & Innodatatics is conducting a 5 Day FDP on Machine Learning on Cloud(Advanced and trending concept). Below are the complete details of the same.	

Duration : 5 Days	
Start Date : 19th July 2021	
End Date : 24th July 2021	
Time : 2 PM to 3:30 PM	
Resource Person : Mr.Bharani Kumar	
Director, Innodatatics Inc.	
Alumnus of ISB & IIT	
Topics covered : Introduction to Auto ML	
Overview of Auto ML techniques	
Practical Application of opensource Auto ML Algorithm - TPOT	
Cloud based Automated ML	
Advanced Concepts of ML on Cloud.	
Interested Faculty from any branch can register.	
Nomination Start Date : 6th July 2021	
Nomination End Date : 14th July 2021	
Link for submitting Nominations: https://forms.gle/sVKmy5AXXMYYaLv17	
Attached is the flyer of the program.	

	A participation certificate will be issued to all the	
	participants who attended all 5 days of the training . Hence request the faculty to submit their certificate names in the nomination form itself. The same names will be considered and corrections in names will not be encouraged.	
	Request you to ensure maximum registrations for the program form your faculty.	
Mentor session	Dear Sir/Madam,	02.07.2021
on How to build		
successful career	Greetings for the day!	
in AI		
	TASK in collaboration with Yes mentor is planning to organize a session on Building a successful career in Cyber Security	
	Below are the details of the session:	
	Mentoring Topic Name	
	Target Audience	
	Duration	
	Proposed Date and Time	
	Mentor Name	
	Introduction to Artificial Intelligence and How to build successful career in AI	

	Undergraduate or Master students from Engineering, Science	
	120 mins	
	03rd July, 4:00 PM to 6:00 PM	
	Mr. Surrendra	
	Tipparaju	
	Students can join the session from link below :	
	https://ciscolearning.webex.com/ciscolearning/k2/j.php?MT ID=t549adb1bca6d06875a6847f8d873a643	
	Attached is the profile of the Speaker and flyer.	
	Please communicate the same with students and ensure the	
	maximum participation.	
TASK - Crux Management	Dear Sir/Madam,	29.06.2021
Services Job requirement for		
2019 and 2020 graduates	Please find the link of new requirement for 2019 and 2020 graduates.	
	Company name - Crux Management Services Pvt Ltd	
	JD Links:	

	For 2019 pass out students: https://www.task.telangana.gov.in/Placements/Crux_Mana gement_Services_Pvt_Ltd_Exe_data_mgmt_2019	
	For 2020 pass out students: https://www.task.telangana.gov.in/Placements/Crux_Mana gement_Services_pvt_Ltd_Exe_data_mgmt_2020	
	Request you to encourage the eligible students to apply for the above job posting. Dear Sir/Madam,	28.06.2021
Medvarsity Job Requirement for 2020 graduates	Please find the link of new requirement for 2020 graduates.	
	Company name - Medvarsity Online Ltd JD Links:	
	For 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/Medvarsity _Online_Ltd_2020	
	Request you to encourage the eligible students to apply for the above job posting.	

TASK - TALLY	Dear Sir/Madam,	24.06.2021
5day online training - (Date:		
29th, June-2021	Creating as from TASK!	
to 3rd, July-2021) Time: 2:00 PM to	Greetings from TASK!	
4:30 PM		
	We are offering TALLY online training to the TASK	
	registered students of B.com (2nd, 3rd years), M.com and MBA (1st, 2nd years). It is a five-day program and free of	
	cost for the TASK registered students.	
	We request you to please communicate with the students to	
	attend the training.	
	TALLY Training details:	
	8	
	Date: 29th, June-2021 to 3rd, July-2021	
	Time: 2:00 PM to 4:30 PM	
	Students Production Link:	
	Students Registration Link:	
	https://forms.gle/LtxMAz1JkAez2y5h7	

	Last date for closing the registrations is 26th, June-2021 by 3:00 PM	
	Note: Students who attend all 5 days will get a participation certificate (Digital).	
	For any further clarifications, please feel free to contact me.	
NASSCOM//Cyb er security	Dear Sir/Madam,	23.06.2021
Centre of Excellence//MO NTH-LONG PROGRAM on "APP SECURITY CHAMPION"	Greetings from Telangana Academy for Skill and Knowledge(TASK)!	
	Hope you are doing well! Our Cybersecurity Centre of Excellence, in partnership with our parent body, NASSCOM, is delighted to bring you this excellent opportunity for your students this year in the booming field of cybersecurity, which is rich with career prospects.	
	This programme, "APP SECURITY CHAMPION", is completely free of cost for the participating students and colleges and has several rewards and certificates waiting to be taken!	
	ABOUT US:	
	The Cybersecurity Centre of Excellence (CCoE) is a joint initiative of Data Security Council of India (DSCI) and the Government of Telangana. It is created to accelerate the cybersecurity and privacy momentum and create a conducive CYBERSECURITY and	

PRIVACY ecosystem which nurtures innovation, entrepreneurship, and capability building. CCoE has played a pivotal role in the Security domain with their SKILL BUILDING INITIATIVES and is bringing you this initiative. BACKGROUND:	
There are over 2 MILLION APPS EACH on Android and iOS. Yet, 55% do not make it to the platform as they are not security and privacy compliant! Security is not limited to a few experts! Knowledge in security and privacy is very important for app developers and infrastructure specialists of the future. The Career Opportunities in Application Security are IMMENSE.	
THE EVENT:	
This MONTH-LONG PROGRAM called "APP SECURITY CHAMPION" will run from 1-JULY-2021-15-AUGUST-2021. It contains THREE PARTS. Registration Last Date : 25 JUNE 2021 THE FIRST PART: A Course on "VULNERABILITY ASSESSMENT FOR APP SEC"- with self-paced classes (51.5 hours of learning, including 13.5 hours of theory and 38 hours of hands-on exercises) and assessments. Live sessions throughout the month with industry experts and veterans from the Industry, Academia and Research. THE GREAT APPSEC HACKATHON in August-2021.	
ELIGIBILITY CRITERIA FOR PARTICIPATION AND	

	BENEFITS:	
	All students in Telangana that are currently pursuing a graduation / post-graduation degree [B.Tech/M.Tech/Diploma-All Streams, B.Sc/M.Sc- (Computers, Electronics, Electricals, Computers),BCA/MCA] and are interested in coding. Multiple exciting cash prizes, certificates, and awards to be won at both individual and college levels.	
	HOW TO PARTICIPATE:	
	Students can register themselves by filling the form given in the link below: https://zfrmz.in/4ajmXJdjba6hMjqLuLy4	
	To know more about this event visit- https://ccoe.dsci.in/ccoe-app-security-champion-students/	
TASK - New Job Requirement for 2020 graduates(2 Job Roles)	Dear Sir/Madam,	21.06.2021
	Greetings from TASK!	
	Please find the link of new requirements for 2020 graduates(2 job roles).	

	Company name - MOL Information Processing Services India Pvt. Ltd	
	JD Links:	
	For 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/MOL_Infor mation_Processing_Services_India_PvtLtd_Doc	
	https://www.task.telangana.gov.in/Placements/MOL_Infor mation_Processing_Services_India_PvtLtd_Acct	
TASK New	Request you to encourage the eligible students to apply for the above job posting. Dear Sir/Madam,	18.06.2021
Job Requirement for 2020 graduates.	Greetings from TASK!	10.00.2021
	Please find the link of new requirement for 2020 graduates.	
	Company name - Teleperformance	

	JD Links: For 2020 pass out students: https://www.task.telangana.gov.in/Placements/Teleperform ance_2020	
	Request you to encourage the eligible students to apply for the above job posting.	
TASK - New Job Requirement for 2019,2020,2021 Graduates	Dear Sir/Madam, Please find the link of new requirement for 2019,2020,2021 Graduates.	04.06.2021
	Company name - Tata Consultancy Services	
	JD Links:	
	For 2021 pass out:	
	https://www.task.telangana.gov.in/Placements/TCS_BPS_2 021	
	For 2020 Pass out:	
	https://www.task.telangana.gov.in/Placements/TCS_BPS_2 020	
	For 2019 Pass out:	
	https://www.task.telangana.gov.in/Placements/TCS_BPS_2	

	019	
	Request you to encourage the eligible students to apply for the above job posting.	
TASK - New Job	Greetings from TASK!	04.06.2021
requirement for 2019 and 2020	Please find the link of new requirement for 2019 and 2020 graduates.	
graduates.	Company name - Adecco India Private Limited	
	JD Links:	
	For 2020 pass out students:	
	https://www.task.telangana.gov.in/Placements/Adecco_Ind ia_Private_Limited_2020	
	For 2019 pass out students: https://www.task.telangana.gov.in/Placements/Adecco_Ind	
	ia_Private_Limited_2019	
	Request you to encourage the eligible students to apply for the above job posting.	
TASK New	Greetings from TASK!	29.05.2021
Requirement for 2020 Pass out students.	Please find the link of new requirement for 2020 Graduated students.	
	Company name - 4AT Consulting	
	JD Link:	
	For 2020 Pass out:	
	https://www.task.telangana.gov.in/Placements/4AT_Consu lting_2021	
	Request you to encourage the eligible students to apply for the above job posting.	
TASK - New Job	Greetings from TASK!	17.05.2021
Requirement for 2019 and 2020 Graduated	Please find the link of new requirement for 2019,2020 students.	

students		
	Company name - Gland Pharma	
	JD Links:	
	For 2019 Pass out: https://www.task.telangana.gov.in/Placements/Gland_Pha rma_2019	
	For 2020 Pass out: https://www.task.telangana.gov.in/Placements/Gland_Pha rma_2020	
	Request you to encourage the eligible students to apply for the above job posting.	
TASK-Oracle FDP on Database Programming	Thank you for attending 5 Day Virtual Faculty Development Program on Database Programming with SQL.	11.05.2021
Programming with SQL April 2021	As part of your training program you are requested to deliver the Oracle Official Curriculum to your students which you have gained through FDP	
	In order to train the students you need to Register the student in your Member Hub for your training program initially and assign the Curriculum to students	
	In this email I am attaching a file contains How to add students and How to assign Course to students Process.	
	Kindly register Minimum 30 Students and Maximum 90 students using your account and inform the students to go through the course and attempt practice tests on their own.	
	Students are eligible to appear for final Certification Exam Online Upon successful completion of all practice tests available in students accounts (This training process should be completed With in 3 months after attending FDP).	
	Student once completed with final certification exam with a pass percentage of 60% Minimum will be awarded with a	

	Certificate from Oracle & TASK	
	Please find the attachment of student registration process.	
	Please feel free to contact me in case of any assistance.	
TASK - New Job Requirement for 2020 & 2021	Please find the link of new requirement for 2020/2021 graduates.	07.05.2021
Graduates	Company name - AXIS BANK JD Link: https://www.task.telangana.gov.in/Placements/AXIS_ BANK2	
	Request you to encourage the eligible students to apply for the above job posting.	
Infosys Campus	Greetings from TASK!	23.04.2021
recruitment Program for NEG students	Please find below the requirement from Infosys for NEG students from 2019, 2020, 2021 batches.	
from 2019, 2020, 2021 batches	This a good oppurtunity for students who have missed out to apply / participate for the previous recruitment drive.	
	Eligible and Interested students can click on Infosys Campus Recruitment Program Form to apply on Infosys portal.	
	Last Date to Apply: Sunday, May 2 <u>, 2021</u>	
	Domain: Infrastructure Services Designation: Operations Executive	
	 Eligibility Criteria: BCA or B.Sc. (Computer Science / Electronics / Mathematics / Physics / Statistics / Information Technology / Information Science only) students from 2019, 2020, 2021 batches 	
	Students who possess aggregate of 60% or 6 CGPA in Class X, XII & Graduation without any active/standing backlogs only need to apply.	
	CTC per Annum: INR 2,22,000	

Additional benefits include a health insurance of INR 4 lakhs per annum and a life cover of INR 30 lakhs per annum				
Bond Pe	eriod: NA			
skills. © M	equired: andidates should have lust be proficient in wo dividual.			
1st roun 2nd rou	n Process: d: Online Test nd: Technical and HR st duration – 100 minut			
ection	Skill tested	No. of questions	Time allocated	
ection I	Reasoning Ability	15	30 minutes	
ection II	Technical Ability	10	30 minute	
ection I	Verbal Ability	20	25 minute	
ection √	Numerical Puzzle Ability	4	15 minutes	
Limited BPM) se Candida environ	ates should not have pa and/or Infosys Group election process in the l ates should be willing t ment. <u>Note: Post</u> the cutoff dat	Company (such a ast 6 months. o relocate and wo	ns – Infosys ork in a 24x7	
discretion will analyze the data of all students who have applied and would shortlist the students based on the criteria specified and shall send out invite letters to the eligible students for the Virtual placement drive.				
All the c	eligible students for the Virtual placement drive. All the details regarding online test schedule, Interview details (for online test shortlist) would be directly communicate (via email) by Infosys only to the students. Selected students would receive communication from			
commu				

	As the data would be captured by Infosys we would not have the visibility of applied students or selected students. Request you to share the requirement with the eligible students and colleges.	
TASK - UNYCC Conclave on 1st May 2021: 21st Century Workplace Skills & Attitude	Greetings from TASK! TASK and UNYCC would like to invite the students of your prestigious institution to participate in a dialogue on "21st century workplace skills". The use of technology and innovative business models is transforming hiring criteria of employers across sectors. In addition to academic and technical skills, companies prefer hiring candidates who are trained in essential new age soft skills such as entrepreneurship, creative thinking, critical thinking and problem solving. In-fact institutions such as the World Economic Forum rank these amongst the top 10 must have skills for entry level candidates. Therefore it is essential that students are made aware of what they need to do in order to build sustained career trajectories in this new world of work. Our eminent line up of speakers include Deloitte's Managing Director Mr Vikas Gupta, Dell India's Chief of Staff Mr Vijaya Mohan Reddy, Novartis Biome India Head RenikaBodla and other such accomplished CEOs & HR Heads. Mr PullelaGopichand (Chief Coach, Indian National Badminton Team) and Mr JayeshRanjan (Principal Secretary, Industries & Commerce and IT, Telangana Government) will deliver the keynote address. Date: May 1st 2021 Time: 3pm to 5pm Medium: Digital All participants will also receive a certificate which will be of immense value during the placement process.	22.04.2021
	name, Email ID & Mobile Number or students can directly	

	1	
	enroll on this link: UNYCC TASK - EVENT	
	REGISTRATION (google.com) or http://bit.ly/unycc_task.	
	For any further information please feel free to call me.	
	Look forward to a positive response.	
TASK-Oracle FDP on	Greetings from TASK,	22.04.2021
Database	Thank you for your interest towards scheduled Faculty	
Programming with SQL April	Development Program on Database Programming with SQL.	
2021	Please find the training details below:	
	Course Title: Database Programming with SQL	
	Training Type: Virtual Training	
	Start Date: 26th April 2021	
	End Date: 30th April 2021	
	Timings: 10 am to 5 pm(one hour break for lunch) Platform: Zoom	
	Trainer Name: JitendraJha (Oracle Master trainer)	
	Tranter Tvarite. Jiteriorajita (Oracle Waster tranter)	
	Single click URL to join	
	Zoom: https://oracle.zoom.us/j/6757609387?pwd=QTQ3OG	
	NEY2xSWVVnaGs1S3RraTRHQT09.	
	Request you to please attend 5 day session by using above link.	
	Kindly login and check your oracle accounts before joining	
	the session in academy.oracle.com->Sign into member hub.	
	Please reach out to me in case of any login problems or any queries.	
Workshop for the foundation	Greetings from TASK!	19.04.2021
program on UX Design by	This is to bring to your kind notice that, TASK, Adobe and NASSCOM Future Skills are providing the UX / UI course	
TASK, Adobe	for the TASK registered students of Engineerig,	
and NASSCOM Future Skills	Polytechnic, Degree, PG and MBA of all years.	

TASK registered students need to regist through the below mentioned google fo	1 0
Link for the Student Registration : https://forms.gle/aYL6sweavieCE9sU9)
After the registrations, students receive from NASCCOM Future Skills. Students registered for this program car	
sessions mentioned below. Course com will be issued, once the student complet course	pletion certificate
You can find the attachment regarding t	he course brochure.
The Calendar for the Workshop:	
Activity	Description
Registration Opening date	
Registration Closing date	
Mail communication from NASSCOM	
Future Skills to the Student	
Orientation Workshop for Students	Course details step t
Mentoring Session Live interaction with Industry experts o Technology 14 th May-2021 3:00 PM to 4:00 PM	n UI/UX
Career Orientation for Students Interaction with Industy experts about t UI/UX domain 28 th May-2021	he Job prospects in
3:00 PM to 4:00 PM Validatory Function	
1 st June-2021 3:00 PM to 4:00 PM	

	with all your associated colleges & students, Here is an exclusive link for TASK to register for the world's biggest coding competition - Code Gladiators 2021.	
	PFB the exclusive link for your students to register & participate: We are excited to share this with you. Kindly share this	
	title of Code Gladiators up for grabs, the competition sees enthusiastic participation and has grown from strength to strength with each passing year. This is open to the students from all the departments and anyone who wishes to participate can give their solutions to solve healthcare challenges faced in the country.	
	Code Gladiators is an annual coding competition by TechGig presented by Cognizant, that draws the best and the brightest coding talent from all parts of India. With multiple contests in emerging technologies and the coveted	
Times Group) CODE GLADIATORS 2021	CODE GLADIATORS 2021Calling the world's best coders to the world's biggest coding arena	
TASK - TECHGIG (registered students and ensure maximum participation. We are excited to inform you that one of our partner TECHGIG is conducting a competition	16.04.2021

Impact by faculties in educating the next generation developers and engineers is unparalleled, and we want to enable them with the best curriculum on Android development and programs, The focus of the summit would help you understand the importance of Android development in today's age, modern programming paradigms on Android and introduce you to our curriculum and programs which can help you deliver high-quality sessions to your students. The summit will be attended by educationalists, scholars, influencers from academia, and faculties from universities. This will also be a chance for you to connect with other faculties and experts interested in this domain. Registrations to the summit are Open: (Click here) Google Faculty Summit 2021 We have opened up the registrations for the Faculty summit on our summit website. Feel free to check out the agenda and Panelist details. We request you to kindly fill the registrations from your colleges/universities. It is an open event for all the faculties/College Management. Digital format: Please feel free to share a Digital invitation to all your faculties for registering for the event. Attached with the email.

	Let's make it a Big Show! and I am really hoping we will get maximum participation from colleges.	
TASK - TALLY 5 day online training - (Date: 19th to 24th, April-2021) Time: 2:00 PM to 4:30 PM	Greetings from TASK! We are offering TALLY online training to the TASK registered students of B.com (2nd, 3rd years), M.com and MBA (1st, 2nd years). It is a five day program and free of cost for the TASK registered students. TALLY Training details: Date: 19th to 24th, April-2021 (21st, April-2021 is holiday) Time: 2:00 PM to 4:30 PM Students Registration Link: https://forms.gle/iocsmwwQRHtmdifX8 Last date for closing the registrations is 16th, April-2021 by 3:00 PM Request to share the above link to the students. For any further clarifications, please feel free to contact me. Note: Students who attends all 5 days will get participation certificate (Digital).	13.04.2021
TASK-Oracle	Greetings from TASK!	09.04.2021
Academy Virtual Girls in ICT Day April 2021	TASK in association with Oracle Academy is coming up with an event <u>Oracle Academy Virtual Girls in ICT Day</u> <u>2021, Japan & Asia Pacific</u> on 22 nd April, 2021 at 10:00 AM to celebrate Girls in ICT Day.	
	International Girls in ICT Day seeks to inspire young women to study and pursue technology. This day highlights female role models in technology as a way to	

	increase interest in information and communication technology and encourage girls to explore careers in the technology sector.	
	Key highlights of the event are:	
	 Event is open for faculty as well as students. Faculties and students attending the event on 22nd April 2021 will be eligible for free course on Data Visualization from Oracle Academy. Participants will be given 15-20 days to complete the course on Data Visualization Attendees completing the course Data Visualization Attendees completing the course Data Visualization will be awarded with OA Badge and course completion certificate. Certificate of attendance will be provided to all attendees of the event. Registration Link for the event: https://go.oracle.com/LP=109493?elqCampaignId=28 7417&src1=:pp:ptr::::TASK Kindly share this registration link with all students and faculty members to register for the event. Students and Faculty members from Degree / Engineering / Polytechnic and PG colleges(Oracle and Non Oracle registered colleges both can apply)are eligible for the event. 	
FDP on Data Analyst by Excel R	Greetings from TASK! TAS would like to mentioned below dates and time for the	06.04.2021
	proposed FDP has been finalised. We have about 300 registrations from social welfare colleges and we request you to register your respective degree and polytechnic college faculty also. The registrations will be closed by 8th April 2021	
	<mark>Data Analyst</mark> / Business <mark>Analyst</mark>	

	No of Days : 15 Days Start Date : 15-04-2021 End Date : 01-05-2021 No. of Hours per Day : 2 Hours/Day Proposed Timings : 1.00PM to 4.00PM Topics Covered: Basic and advanced concepts of Microsoft Excel, Introduction to the tool TABLEAU and My SQL. Registration link : https://forms.gle/PEbjD8yGmKERernW9	
Infosys Headstart - New Digital Learning Initiative : Reg.	 Greetings from TASK! TASK – Infosys would like to introduce a new free learning portal, Infosys Headstart https://infosysheadstart.onwingspan.com/. We have enabled bulk registrations from the backend. We have attached a registration template and Data Processing Agreement template with this mailer for the same. Please check the files and submit student information for bulk registration. Reference Links: Link to platform - https://infosysheadstart.onwingspan.com/ Link to platform - https://infosysheadstart.onwingspan.com/ Link to "Headstart initiative video" which can be shown to students – https://youtu.be/3OVQxtjj6dk Next Steps: Share the list of students from 1st year to final year to be registered in the excel format enclosed(Headstart_Student_List.xlsx) Note : If the student in your list is already registered on the platform, the platform will take care not to register them again with the same email id. Please share the required documents with rm6-task@telangana.gov.in latest by 2nd Apr 2021. We are sharing this information with you as part of a beta launch, while the formal launch is expected in the first / second week of April 2021. We request you not to post this on social media. 	01.04.2021

TCS requirement for 2020 graduating NEG students	TCS has come up with requirement for 2020 NEG students. Please click on : https://www.task.telangana.gov.in/Placements/TCS to view the JD. Request you to encourage students to apply.	01.04.2021
Virtual Meeting to brief TASK UI / UX Course for TASK registered Degree and PG Faculty	Greetings from TASK! TASK would like to conduct a virtual meeting with the TASK registered college Principals, HOD's, Faculties for an overview of TASK-Adobe NASCCOM future skills program. Attached Brochure for your reference. Virtual Meeting Date : 1st, April-2021 Time: 12:00 PM to 1:00 PM We request you to please register in Below link to participate in the meeting. https://forms.gle/zSuDPhRwTRsEVavq6 Last date to close the registration is 29th, March-2021 by 5:00 PM Details of the program are attached. How does UX benefit students of Management stream	27.03.2021
	Stream Potential Job roles Details	

			Product research	
	Management	UX strategist UX product manager UX researcher Data analyst UX erchitect	The foundational stage of Depends on understandin research, the designer can Excellent channels for con interviews with users and channel.	ig customer needs, go: make informed decis ducting research inclu
	Marketing / Advertising)	UX architect UX analyst UX content strategist UX copywriter	Content strategy Content strategy plans for content. Consequently, a c needs. He/ she then works bridges the gap between h profitability and usability writing, and editing it, fol	content strategist unde s on planning, develop pusiness goals and use of content via a conte
		C benefit students of		
	Stream Arts, Design	Potential Job roles Visual designer UI designer Digital designer UI artist 	Details Visual designing · Visual Designers prov websites. Their duties incl visual concepts for approv their graphic designs. The	
Salesforce Fundamentals Women in Tech Initiative- Program details	program calle Initiative. Many of Sales looking forwa organisations	ed Salesforce Fundar sforce hiring partner ard to hire talented v as part of their Dive	· · · · · · · · · · · · · · · · · · ·	27.03.2021

	students.	
	The enrolled students will go through a series of curated trailmixes and virtual trainings taken by instructors over the weekends.	
	Please find the attachment for the complete details about the program.	
	Kindly note that this program is only for TASK registered girl students.	
	Eligible students for this initiative is as below :	
	Degree Colleges - 2nd year Girl Students of Bcom, BSc, BBA	
	Please see below the key highlights.	
	 <u>Key Highlights:</u> Launch date - April, 2021 Duration: 12 weeks Time commitment from students: 4-6 hours a week Format: Self paced with guided help from instructors Mentorship and Job Connections I request to forward to the colleges and get the Colleges, 	
	College SPOCS and faculty nominations by 29th March, 2021:	
	https://forms.gle/8TafFW44f8zUtTe6A	
i4TS online Conference - III on 27th March '2021	We are glad to inform you that our initiative i4TS (Innovating for Telangana State) - SamajikaAavishkaranaluSaralamainaJeevithaniki is progressing as per schedule. So far, around 200 youth from different parts of Telangana are going through the 3-month Entrepreneurship Development Program.	26.03.2021
	Conferences are an integral part of the i4TS program. The first and second conferences were held on December 19th, 2020 and February 13th, 2021 respectively.	

Saturday, March 22		
Time: 10.00 AM to	1.30 PM	
The Agenda:		
Time	Session	About the session
1000 – 1015 hrs	I4TS program recap	
1015 – 1115 hrs	Panel – I	Evolution of im
	Investing for Impact	do investors loc
		Measurement o
		• Ms.Padm
		Indian A
		Mr.Jayes
		Protein VMs.Ansh
		Ms.AnsnMr. Ravi
		• IVII. Kavi
		Session N
		Member,
		wiember,
		What is the pers
		leading change
		and rural entrep
		and challenges?
		about a social cl
		• P. Balaji -
		Officer, V
	Panel - II	Mr.Prade
1115 - 1215 hrs	CEO's Panel	Group
	Leading the Change	• Mr.Saura
		• Dr. Gane
		Honeyw
		Director
		Internatio
		• Dr. Krish
		Biotech I
		Session M
1215 - 1220 hrs	l	

	Dorthographin (TPC)	
	Partnership (TBC) Announcement of AIM partnering with TASK	
	Mr.RamananRamanathan, Mission Director Atal	
	Innovation Mission	
	1220 - 1235 hrs	
	Key Note	
	Sri JayeshRanjan, IAS. Principal Secretary, Industries	
	& Commerce & Information Technology, Electronics	
	& Communications, Government of Telangana	
	1235 – 1315 hrs	
	Fireside chat	
	Dr.KiranKarnik , Director, Central Board of	
	Directors, RBI	
	1315 - 1320 hrs	
	Vote of Thanks	
	Ms.Shruti, Sr.Program Manager, TASK	
	Please find the attached eDM	
	Please find the attached eDM	
	Looking forward to your support and presence,	
	Looking forward to your support and presence,	
TASK-UIPATH	This is an introductory mail to one of our Industry Partner	23.03.2021
Academic	"UIPATH Academic Alliance."	20.00.2021
Alliance	on min matter marke.	
Program	About UIPATH Academic Alliance:	
riogram		
	UIPATH provide individuals with free courses for UIPATH	
	Academy and started teaching RPA(Robotic Process	
	Automation) to over 35,000 now certified developers,	
	propelling our rocket ship growth. UIPATH is the fastest-	
	growing enterprise software company in history. The	
	employee base grew to over 2,900 employees today, across	
	25+ offices. In 2019 UIPATH was named a leader in the	
	Gartner Magic Quadrant for Robotic Process Automation	
	Software.	
	Institutionalizing RPA (Robotic Process	
	Automation) as a discipline in academia.	
	• Preparing students and educators with in-demand	
	to automation skills.	
	• Creating a differentiation for universities.	
	• Connecting students and educators to a community	
	of practitioners.	
	TASK & UIPATH program has signed for partnership to	

help students & faculty of partner institutes gain i expertise in latest industry technologies.	insight &
Through this association we will be offering three courses to all the TASK membership colleges.	different
Robotic Process Automation (RPA) E Development v1.0- Lecture (40 hours), Hands-On hours).	e
• Step in to RPA(Robotic Process Autom	naton) - 4
hours. • Introduction to RPA (Robotic Automaton) -14 hours, can be a 1 credit elective or course.	
At the end of each course, students will earn a part certificate and Learners earn an UIPATH Juni Developer Certification after successfully passi certification exam.	ior RPA
As part of UIPATH and TASK collaboration would like to open-up academic membership colleges under TASK who can get the direct adva UIPATH Certification.	o to the
The benefits for an engineering college to participa UIPATH Academic program include:	ate in the
1. FDP is organized on Robotic Process Aut (RPA) Design & Development v1.0- Lecture (40) Hands-On Lab (20 hours) to the faculty of the p colleges,Instructor logins, course conten given.	0 hours), partnered nt etc. are
2. Industry aligned global cert 3. Faculty to attend FDP and roll out 4. Dedicated slots (at least 2 hrs per week) in a	Courses. academic
curriculum for students to go through a p 5. Monthly monitoring of program.	program.
Who can take:	
Students of 3.1 & 4.1sem year of Engineering. Students of 2.1 & 3.1sem year of MCA and De	

	Kindly confirm your Interest to be part of this partnership on and before 27-03-2021.Attached PDFs will give all the additional information you might need.For more details regarding this program, please feel free to contact me for any assistance	
TASK-AWS DeepRacer Women's League	 Greetings from TASK! Amazon Web Services(AWS) is hosting a DeepRacer League exclusively for female students. Hosting a DeepRacer league is a conscious attempt to foster community learning and mark a headway towards the early adoption of AI, ML technologies for a future, gender diversified workforce. 	23.03.2021
	The AWS Deepracer Women's League will help foster community learning and help with the early adoption of artificial intelligence and machine learning technology for a more diverse workforce of the future. The initiative aims to bring together women students to learn collaboratively and get hands on with machine learning through a cloud-based virtual racing simulator. During the course of the competition, participants get the opportunity to build their ML models and compete on tracks virtually while driving deep into advanced reinforcement learning and upskilling with ML.	
	India has a significant stake in the AI revolution of the world. The Govt. of India, at the helm, is driving AI innovation & adoption with multiple initiatives like #AIForAll, Digital India, Future Skill Prime etc. This is a clear opportunity to build an AI Skilled workforce for the future. AWS DeepRacer Women's League is designed to promote up-skilling in machine learning (ML), as well as inclusion and diversity in the technology sector. This is a women centric league to attract diverse talent in this fiercely competitive talent market place and upskill them on emerging technologies like AWS machine learning.	

	Eligibility : All streams(Any year) - Engineering and Non Engineering Women who are at least 18 years of age currently enrolled with an academic institution. Registration Link : (https://awsdeepracerleague.in/womens- league-2021/) Last Date for Registration - April 4th This event is to mark the occasion of International Women's day and will begin with regional elimination round comprising of AWS DeepRacer community raceacross four regions in India (North, East, West, South), followed by a national elimination community race and then the grand finale <u>on 21* April</u> , where participants will upload their models and run on the virtual track which will be streamed live on twitch.tv. The event is open to all women students currently enrolled with higher education academic institutions in India. They would register on AWS Educate, complete the AWS machine learning and AWS DeepRacer course modules as well as attend the enablement sessions planned during the event. You are requested to promote the event amongst your female student audience and ensure maximum participation.	
TASK-NRSC- ISRO : Water Quiz on 23rd March (Tuesday)	Greetings from TASK! We are glad to inform you that on the occasion of world water day 2021 (i.e, 22.03.2021), TASK and NRSC-ISRO are conducting the water quiz for all the students. please, find attached the document. The details are: Date of the Quiz: 23rd March (Tuesday) Time: 5.00 PM	22.03.2021

	Registration link: https://bit.ly/3f8JQml	
	Kindly share the information with all our students.	
TASK - Tableau DataViz	Greetings from TASK!	19.03.2021
Challenge - 2021 by SmartBridge in Collaboration	TASK in association with Smart Bridge organizing a boot camp for faculty and student.Below are the details	
with Tableau	About the Program:	
	SmartBridge in partnership with Tableau is hosting India's Biggest DataViz Challenge, a unique program for both students & educators.	
	This exclusive academic initiative program is to build practical and analytical data skills in students & educators. The DataViz Challenge will provide hands-on exposure to the data analytics powered by Tableau E-Learning and Tableau Software.	
	Event Name: Tableau DataViz Challenge - 2021 Registration Link: <u>https://smartinternz.com/tableau- dataviz-challenge-2021</u> Event Format: 5 Days Bootcamp + 10 Days DataViz Challenge	
	Participants Eligible: <mark>Faculty & Students from Engineering, MBA, BBA, Degree & Diploma</mark> Last Date of Registration: 31st March 2021	
	This program is completely FREE of cost for the participants as it is sponsored by Tableau.	
	 Program Benefits : Free Tableau License for Students & Educators Access to Tableau's E-Learning Technical Bootcamp on Tableau Software Mentoring Support from Data Experts Exciting prizes for the Top 3 Winners Special Jury Awards for Best Vizzes Internship Opportunities for the Winners 	

	Bootcamp Bootcamp	& Challenge Particip	ation Certificates.	
	Program Agenda : 5 Days Bootcamp on Data Visualization powered by Tableau Timings : 5:30PM – 8:30PM IST Mode of Training: Instructor-led (Online via Zoom)			
	Description	Start Date	End Date	
	Boot Camp	01st April 2021	06th April 2021	
	Data Viz Challenge	07th April 2021	17th April 2021	
	Winner Announcement	301	h April	
	We request you to circ colleges Faculty & Stu from this wonderful In Please let us know if y	dents to leverage ma nitiative.	aximum benefits	
TASK - New Job Requirement updated on TASK portal	Greetings from TASK Requirement from Foo graduates is published Students can click here https://www.task formance-3 to view JD Request you encourag	cussoft and Teleperfo d on TASK portal. c.telangana.gov.in/Pl d and apply for Telep	acements/Teleper	19.03.2021
TASK-Yes Mentor Partnership	Greetings for the day! This is introductory m Mentor'. About Yes Mentor:		ustry partner 'Yes	17.03.2021

connect college stud industry subject ma mentoring session th and at the same time skilled resource poor Mentor talk session	GO for providing mentorship session to dents and entry-level professionals with atter experts through focused hereby helping with reduced skill-gap be contributing increased and competent of towards self reliantTelangana. I will be done on both generalized and elow are the topics and Target
Category	То
	Introduction to information Techno ir
	Introduction to Cyber Security and Sec
Specialized Mento	oring Introduction to Artificial Intelligen AI and Da
	Building a successful career in
	Building a successful
	Building a successful
	How to build Future S
	Developing and exercising Cog
	Analytical Thinking skills, Critica
Generalized Mento	oring Skills, and Probl
	Developing and exercising Emd
	Advanced communications Skills, F
	Empathy and add Developing and Practi
	Becoming a Socio-Cultu
pm to 5:00 pm and o Session request show that respective week	will be typically on Friday from 1:00 or on Saturday 9:00 am to 5:00 pm IST. uld be given on or before Tuesday of

	https://forms.gle/UYX8AxJYobTyzpSXA	
	We can deliver the session for individual college or for 2 or 3 colleges together.	
TASK - FDP on Data Analyst by Excel R	Greetings from TASK! TASK in collaboration with ExcelR is planning to conduct a Faculty Development Program on Data Analysis to the faculty of Degree,Govt.Polytechnic and Social Welfare colleges.Below are the details of the same.Attached is the proposal. The faculty registration link is also mentioned below. Data Analyst/ Business Analyst No of Days : 15 Days No. of Hours per Day : 2 Hours/Day Proposed Timings : 11AM - 1PM / 2PM to 4PM / 6PM - 8PM Topics Covered: Basic and advanced concepts of Microsoft Excel, Introduction to the tool TABLEAU and My SQL. Kindly communicate the same to your faculty to register on below link for the program. Registration link : https://forms.gle/PEbjD8yGmKERernW9 Please note that the execution date of the training program will be communicated soon.	17.03.2021
TASK - TALLY 5 day online training - (Date: 15th to 19th, March-2021) Time: 2:00 PM to	Greetings from TASK! We are offering 4th batch of TALLY online training to the TASK registered students of B.com, M.com and MBA . It is a five day program and free of cost for the TASK registered students.	09.03.2021

	TALLY Training details:	
	Date: 15th to 19th, March -2021 Time: 2:00 PM to 4:30 PM	
	Students Registration Link:	
	https://forms.gle/dXxFFszbRwfijUEy7	
	Last date for closing the registrations is 12th, March-2021 by 3:00 PM	
	Request to share the above link to the students.	
	Training is through Octa platform and after the registrations, I will coordinate with the octa team for further process.	
	For any further clarifications, please feel free to contact me.	
TASK - Boot camp and Build- A-Thon on Machine Learning with	TASK in association with Oracle Academy is planning to conduct Boot camp and Build-A-Thon on Machine Learning with Java.	03.03.2021
Java	About the event:	
	TASK in association with Oracle Academy is launching a program exclusively for students to provide Hands-On Experience on building projects. Exposure to live projects will help students to gain technical skills and also gain Hands-On experience on Machine Learning with Java.	
	 Eligibility : Students from 3rd & Final year CSE, IT and ECE backgrounds Students from Final Year Degree which support computers and Information technology. Students who will complete the course Artificial Intelligence with Machine Learning in Java in Oracle Academy will have have an opportunity to attend 3 - Day Boot Camp. 	

 Registration link for the Boot camp will be only sent to students who have certified on Artificial Intelligence with Machine Learning in Java in Oracle Academy. Students who will attend 3- Day Boot camp only can participate in Project Build-A-Thon. 	
 Benefits of the Program: E Participation Certificate (participant should Build a project to Receive the Participation Certificate) Top three projects submitted by students will be presented with awards. Mentoring support from trainers. 	
 Program Agenda: Event launch-1st April 2021 E learning- Artificial Intelligence with Machine Learning in Java in Oracle Academy From 2nd April 2021 to 15th April 2021 3- Day Boot camp is from 23rd April to 25th April 2021 Project Build-a-Thon is from 26th April to 8th May 2021 Project Reviews & Announcing Top 15 projects - 13th May 2021 Final Jury Round & Winner Announcement 14th May 2021 to 16th May 2021 Students would need faculty assistance to complete E learning on Artificial Intelligence with Machine Learning in Java in Oracle Academy From 2nd April 2021 to 15th April 2021 	
So, Request you to kindly take faculty nominations from Oracle registered colleges(attached OA registered list for your reference) in below link by 10th March 2021(any mail id's will be accepted).	
https://forms.gle/QUfRLsm6CdDKt6PC6 Onboarding students into Oracle Academy process will be shared with Faculty members once we get enough	

	nominations.	
	Please reach out to me in case of any queries.	
	Greetings!	02.03.2021
TASK - Infosys Soft Skills Webinar from	TASK in association with Infosys planned Soft Skills webinar from 3 rd to 5 th March 2021 for all students. <u>Modules:</u>	
3rd to 5th March 2021	 Body Language for Successful Interviews Elevating your Virtual Presence Interactive Skills 	
	4. Professional Etiquette Kindly share the below link with all Colleges and student groups.	
	Days/Date of Session: 3 rd to 5 th March 2021(Wednesday to Friday)	
	Time: 3PM to 4:30PM	
	Attached poster for your reference.	
	Kindly click the below link to join the session	
	Link: https://infosys.webex.com/meet/haripriya.chauhan Resource Person: Ms. HariPriya Chauhan(Infosys)	
TASK UI / UX Course for Degree and Other PG Students	Greetings from TASK!	28.02.2021
	We are glad to inform you that TASK is launching UI / UX course in partnership with Adobe and NASSCOM Future Skills.	
	Details of the program are attached.	
	How does UX benefit students of Degree and Other PG stream:	

Stream	Potential Job roles
Arts, Design	 Visual designer UI designer Digital designer UI artist
Implementation Process 1). Student need to regis google form link mention	ster for the program through the
https://forms.gle/SPoz5B	
2). Students will receive Skills team 2 to 3 days af regarding platform detai link etc. A separate comm	a mail from NASSCOM Future fter the last date of registration ils, webinar and project submission munity of students who register rogram will be created for focussed
3). The course starts with workshop on 6th March	h a 2-hour Virtual Instructor led
Batch size: Workshop date: Time:	Up to 2000 students 6th, March - 2021 4:00 PM to 6:00 PM
Duration: Requirement's: Internet facility	2 hours Computer / laptop with
	I Future Skills will host 4 live th and 27th of March (all Saturdays
Students registered once	can attend all the 4 sessions

Virtual Workshop: Session on Introduction to Oracle Academy Primavera P6 Teaching Resources	 4). After the first workshop, student can access the self-learning modules to practice working with UX. The course contains 38 hours of self-learning content. They can complete the course and submit a relevant project to achieve the certificate by 31st March 2021. 5). Projects submitted will be evaluated and a course completion certificate co-branded by TASK, Adobe and NASCCOM Future Skills will be given to students within 20 days of project submission. The program is free of cost. This is a reminder mail regarding Virtual workshop on Introduction to Oracle Academy Primavera P6 Teaching Resources, scheduled on 25th February 2021, Thursday at 5.30 p.m. Kindly remind OA members who are teaching project management and civil engineering courses to register for the session in below link: https://go.oracle.com/LP=96468?elqCampaignId=258014&csr c1=:pp:ptr:::TASK 	23.02.2021
TASK TALLY 5	c1=:pp:ptr::::TASK Greetings from TASK!	23.02.2021
day online training - (Date: 1st to 5th, March-2021), Time: 2:00 PM to 4:30 PM)	We are offering TALLY - ACE online training to the TASK registered students of B.com (2nd & 3rd yrs), M.com (2nd yrs) and MBA (2nd yrs). It is a five day program and free of cost for the TASK registered students.	
	We request you to please ask the students to register for the Program. <u>TALLY Training details:</u>	

	Date: 1st to 5th, March -2021 Time: 2:00 PM to 4:30 PM	
	Students Registration Link:	
	https://forms.gle/gUTHw9LtPmSvNWmv7	
	Last date for closing the registrations is 25th, February-2021 by 4:00 PM	
	Request to share the above link to the students	
	Training is through octa platform and after the registrations, I will coordinate with the octa team for further process.	
	For any further clarifications, please feel free to contact me.	
TASK-Unschool Partnership	TASK in association with Unschool is offering free online courses for 2nd, Final year Degree and 2nd Year MBA students.	23.02.2021
	Unschool is an online learning market-place for everyone who has the knowledge to share or a skill to teach, to coach inspired individuals and ensure employability with the right skill set. Unschool believe that teaching is not meant only for conventional teachers or institutions, but in the idea that anyone and everyone can teach and learn.	
	The Courses Unschool offer on, 1. Technology 2. Management 3. Humanities Benefits for Students:	
	 Courses which are taught by industry experts. Unschool will provide free access to 10,000 students under the partnership with TASK for a validity duration of 60 days (two months) from the date of enrolment Unschool provides 24/7 support and doubt clarification 	

	 Unschool platform is accessible on all devices with limited internet as well Learner can learn the way they want to. So, videos, PDFs and voice notes. Upon completion of the course, if the student requests for a course completion certificate then Unschool shall charge INR 2500/- from the learner. The Students who get a course completion certificate from Unschool would be eligible to apply for the internship opportunities available at Unschool at no extra cost. Kindly share this email with TASK associated Degree and MBA colleges and inform interested colleges to register for the courses in below link: https://forms.gle/56fPk7fHoYZonnDEA Note: Student should complete one course in Unschool platform and Colleges should give commitment that 50% of the students will complete the course. Please let me know in case of any queries. 	
New Job requirements for 2020 & 2021 Graduates	Please find the link of new requirements from the below companies for 2020/2021 graduates. VMax e- Solutions : https://www.task.telangana.gov.in/Placements/Vmax_e- Solutions_India_Pvt_Ltd MRF Tyres 2021: https://www.task.telangana.gov.in/Placements/MRF_ Limited2021_N Request you to encourage the eligible students to apply for the above job posting.	17.02.2021
Ramky Jobs for	Greetings from TASK!	16.02.2021

2021 graduating students for TASK Registered /Non Engginering MBA/MSc students	We have a requirement for MBA(Marketing)/MSc(Chemistry) for RamkyEnviro Engineers Ltd. We request you to encourage all students to apply for Ramky (Both TASK Registed and Non TASK Registered) can apply Using below link. Eligibility: 2021 Graduating students with minimum 60% aggregate throughout the academics without any backlogs and should be open to work across site locations in India can apply through https://forms.gle/ngDjcXtMsLWyDMTaA	
TASK Infosys Soft Skills FEP from 22nd to 24th Feb 2021	Greetings from TASK! TASK in association with Infosys is planned to conduct a Soft Skills FEP for Faculty of English and H&S department. Request you to please share the email to nominate the faculty in below link. Please note that we can only allow 3 Faculty from each Institution and also on FCFS basis. Please Share the nominations in below link as soon as possible. https://docs.google.com/forms/d/e/1FAIpQLSd3x7iF7cows VNOOnUDRwq7loZBnrnrg7b3shIp9C21CC9jVw/viewform	16.02.2021
TASK - New Job Requirements for 2019/2020 graduates	Greetings from TASK! Please find the link of new requirements from the below companies for 2019/ 2020 graduates. Waterleaf Group: https://www.task.telangana.gov.in/Placements/WA TERLEAF_CONSULTANTS_PVT_LTD MOL (Accounting) - https://www.task.telangana.gov.in/Placements/MOL_Infor mation_Processing_Services_India_PvtLtdACC	13.02.2021

	MOL (Documentation) - https://www.task.telangana.gov.in/Placements/MOL_Infor mation_Processing_Services_India_PvtLtdDoc_	
TASK-Oracle FDP on Database Programming	TASK would like to inform you that we have scheduled Oracle Academy 5 Day FDP program. Please find below updated schedule:	10.02.2021
Programming with SQL April 2021	Course Title: Database Programming with SQL Training Type: Virtual Training Start Date: 26th April 2021 End Date: 30th April 2021 Timings: 10 am to 5 pm(one hour break for lunch) Batch Size: 32	
	Please forward the updated schedule to all oracle registered colleges and inform the faculty to register for the FDP in below link: https://forms.gle/ExhfxJpMmf637eVQA	
TASK - Batch-2	Kindly let me know in case of any questions. Greetings from TASK!	09.02.2021
TALLY 5 day online training - (Date: 15th to 19th, Feb 2021	TASK would like to offer the TALLY - ACE online training to the TASK registered students of B.com (2nd & 3rd yrs), M.com (2nd yrs) and MBA (2nd yrs). It is a five day program and free of cost for the TASK registered students. Batch size is minimum 150.	
	<u>TALLY Training details:</u> Date: 15th to 19th, February -2021 Time: 2:00 PM to 4:30 PM	

	Students Registration Link:	
	https://forms.gle/hqYGXshHYQxoXzjo6	
	Last date for closing the registrations is 11th, February-2021 by 4:00 PM	
	Request to share the above link to the students	
	Training is through octa platform and after the registrations, I will coordinate with the octa team for further process.	
	For any further clarifications, please feel free to contact me.	
TASK - New Job	Greetings from TASK!	04.02.2021
requirements for 2020 Graduates	Please find the link of new requirement from the below companies for 2020 graduates.	
	VITA Technologies: https://www.task.telangana.gov.in/Placemen ts/Vita_technologies_(VITATECH_Solutions_Private_Limit ed)	
	Request you to encourage the eligible students to apply for the above job postings.	
TASK - New Job Requirements from Urban Tech Company	Please find the link of new requirement from the UrbanTech company for 2019/2020 graduates for different positions.	04.02.2021
	Urban Tech:	
	1. https://www.task.telangana.gov.in/Placements/URBANT ECH_SERVICES_PHP	

	 2. https://www.task.telangana.gov.in/Placements/URBANT ECH_SERVICES_app_dev 3. https://www.task.telangana.gov.in/Placements/URBANT ECH_SERVICES_JrAndroid_ Request you to encourage the eligible students to apply for the above job postings. 	
TASK - New Job Requirements for 2021 graduates	Please find the links of new requirement from the below companies for 2021 graduates. MRF Tyres: https://www.task.telangana.gov.in/Placements/MRF _Limited2021	30.01.2021
Oracle Academy I-Learning Upgrade details	This is to inform you that Oracle Academy i-Learning is upgraded to Member Hub. Please find the attachments for the new steps included as part of Academy Adding Faculty Account, Create Channel and Adding Students, How to generate Student Reports and Login to Student Account Procedure. Please forward it to all Oracle registered colleges and get them to use Oracle Academy Member Hub effectively and also inform them to reach out to me for any questions or doubts regarding Member Hub. In case of existing faculty accounts, Please inform the	28.01.2021
	 In case of existing faculty accounts, i fease inform the faculty to follow below steps: Navigate to academy.oracle.com -> Sign into Member Hub -> Username is email id -> Click on forgot password and reset the password. Please let me know in case of any additional information from my end. 	
IBM Open PTECH Platform	This is an introductory mail to one of our Industry Partner "IBM Open PTECH."	23.01.2021

WALK		
THROUGH	About "IBM Open PTECH":	
	IBM Open PTECH is a free online learning platform that has industry courses on emerging technologies and professional development. IBM is bringing learning content that is relevant for students who are getting ready to join the workplaces where more than 90% of the jobs will require relevant understanding of new technologies. Emerging technology learning is relevant to every student whether they are on an IT pathway or not. Employers today across industries are looking for well- balanced people with transferable skills like collaboration and critical thinking, in combination with technical skills and workplace readiness. Open PTECH provides self- paced, self-learning courses for students to equip themselves with relevant skills, that are essential for current and future workplaces.	
	 Open PTECH has 80+ hours of learning content from IBM and another 40+ hours from IBM industry partners. Data Science, Artificial Intelligence, Cloud computing, Cyber security, Block chain, Coding and Computational thinking, Quantum Computing, machine learning, Professional skills (collaboration and team building, presentation skill, critical thinking, inter-personal skills, work delivery skills and Agile methodologies) and, Workplace skills like Design thinking, Project Management and others IBM and its partners would support the colleges/education institutions associated with TASK, to onboard students through custom links created for the colleges, train and hand-hold colleges SPOCs and faculty members to navigate the Open PTECH platform to learn (and guide students) as well as to track learning of their students through dashboards and (downloadable) reports, and organize webinars/online seminars on topics relevant to technology and careers to be conducted by IBM leaders and industry experts to additionally support students leaning. 	
	In the first phase we are planning to roll out the program in 45 colleges. We are looking for 25 engineering, 15 Degree	

	and 10 Polytechnic colleges to take up the course.	
	Please nominate the college details here on or before 27th Jan 2021.Attached PDF will give all the additional information you might need.For more details regarding this program, please feel free to contact me for any assistance	
TASK - RPA Build-A-Thon by SmartBridge in collaboration with Blue Prism University	 TASK in association with Smart Bridge organizing an Faculty Development Program on RPA Build-a-thon.Below are the details About the program: Smart Bridge in collaboration with Blue Prism invites all faculty members to participate 3 day Hands on Training on Robotic Process Automation and 3 Day Mentor guided Project Development in RPA Build-A-Thon program. RPA Build-A-Thon is an exclusive skill building initiative by SmartBridge& Blue Prism University to Enable Educator with Robotic Process Automation skills. This program is a learning by doing event packed with Hands-on training,Mentoring sessions and Digital workforce development powered by SmartInternz& Blue Prism University. Benefits for the Participants: Free access to Blue Prism Courses Free access to Blue Prism Software Tech Boot camps on Robotic Process Automation Mentoring support by RPA Experts Develop RPA applications for Real time user case Boot camp and Build a thon Participation Certificate (Must submit project to receive certificate) 	19.01.2021
	Eligibilty - <mark>Engineering/MCA/Sciences/Polytechnics</mark> Program Agenda: 3-Days Hands-on Training; Virtual Program via WebEx	

	Date	Content of 3-Day Han	
		Introduction to Robotic Process Au	
		Introduction to Blue Prism Tool	
		What is Process Studio	
		What is Object Studio	
		Difference between Process and Ob	
	01-02-2021 to 03-02-2021	Setting-up the Blue Prism Tool	
		Work Queues	
		Exception Management Overview	
		Email Automation	
		Building Web Application Automa	
	04-02-2021 to 06-02-2021		
	To know more details abo	out the program Click here	
	Click here to register for t	he program	
	Last date for registration	- <mark>30th January 2021</mark>	
	Request you to please con colleges,so that the faculty opportunity.	nmunicate the same with y can make maximum use of this	
Visionary RCM Applied student list- 2021 campus	5	ew application for Visionary RCM 2021. Applied student details your referance.	19.01.2021
recruitment.	We will extend the last	date to apply till 22nd January.	
	Eligiblity	criteria:	
	Biochemistry, Biotechr Anatomy, Bioinformatics Nano-technology, Food Dietetics, Botany) // Graduating students w	Science Graduate in (Microbiology, nology, Biology, Bio-Medical, , Physiology, Zoology, Bioscience, d technology, Nutrition and B.Pharma /M Pharma 2021 with minimum 50% aggregate nics without any backlogs only apply.	

	CTC	per	An	num	:	
	Doct 2	monthe	Training	INID	1 56 000	
	Post 2	months	Training:	INR	1,56,000	
		andidate needs	0	-	0	
	-	ost training and ndidate would				
	of		NR		1,56,000)	
	Dlasa				-1: -1.	
	Please on https://	www.task.telar	igana.gov.in/P	lacements	click Visionar	
	-	<u>/led_code_2021</u>	0 0	the	JD.	
	Request	you to encou	irage eligible	TASK	registered	
	students	jou to cheot	to	111010	apply.	
TASK - Unicorn	Greetings	from TASK!				18.12.2020
Technologies Job		1 1 • 11.		1	1 1	
openings, LLC Requirements -	Portal.	echnologies LL	c requirement	s are uplo	aded on	
on TASK Portal						
	Below are	the links :				
	BE/B.Tech,	,MBA graduate	s from 2020 ba	tch can c	lick	
	-	www.task.telar				
	recimolog	ies_LLC2 to ap	piy ioi ii iecit	iner posit	1011.	
		,BCom,MBA gr				
	-	www.task.telar ies_LLC to app	· ·			
	position.		iy for ear eer	ner nepre	bentative	
	DE/D To ale	PCom MDA or	a decato a fuerra 1	010 h a t al		
	-	,BCom,MBA gr www.task.telar				
	Technolog	ies_LLC1 to ap	0 0			
	position.					
	BE/B.Tech,	,MBA graduate	s from 2019 ba	tch can cl	ick	
	-	www.task.telar	0 0			
	recuniting	ies_LLC3 to ap	PIY IOI II Keci	uner pos		

	Greetings from TASK!!	16.12.2020
NOTIFICATION FOR EXTENSION OF STUDENT REGISTRATION FOR THE ACADEMIC YEAR 2020-21	 TASK would like to inform you that, TASK has decided to extend the dates for Students Registrations till 30th Jan 2021 for the academic year 2020-21. Please note that, this will the final extension and no further extension request will be entertained. Late fee charges is Rs.100.00 + 18% GST till 13th Feb 2021. Note: ATTACHED NOTIFICATION FOR EXTENSION OF STUDENT REGISTRATION FOR THE ACADEMIC YEAR 2020-21. 	
Infosys-TASK Special Sessions for Girl Students 11th Dec 2020	Greetings from TASK! Infosys in conducting a session for Girl students from Degree Colleges, Kindly ask Girl students to attend the session. We have only 250 students for each session. Link: https://infosys.webex.com/meet/hina.azad Date: 11th Dec 2020 Time: 9AM to 11AM	10.12.2020
Infosys NEG Campus Recruitment Program 2021 : Guidelines for Infosys Online Test and FAQ's	Time: 9AM to 11AM Greetings from TASK! All Infosys shortlisted students would be given opportunity to participate in the Infosys recruitment drive irrespective whether they have registered in InfyTQ. Students who have registered in InfyTQ would be able to take online test as per their preferred slot. Online test slots would be allocated by Infosys for candidates who have not registered in InfyTQ / could not select their slots. Student would receive series of emails from Infosys regarding the online test details. Pls ask students to check	08.12.2020

	their emails regularly.	
	Shortlised student should be ready with PC /laptop as per the guidelines from Infosys.Students can also take support for their respective college to take the test.	
	Request you to share this Communication with students and be ready with necessary arrangements, and encourage maximum students to take online test.	
	As you are aware that TASK has scheduled " EXCELLERATE !" A 5 day program for Infosys shortlisted students from 6th - 10th December 2020.	
	These are very useful sessions would help students to prepare for the online test / Interviews, Hence pls encourage students to take leverage.	
	Attached is the list of shortlisted candidates for your reference,	
RE: TASK - Confirmation for	Dear Madam,	05.12.2020
Jio Drive -200 nos Home Sales	Jio HR has confirmed the date as on 14 th Dec 2020.	
Officers in Jio !!!	Kindly arrange all necessary requirements as pet the date i.e 14 th Dec 2020.	
	Thanks,	
	Regards, kumar	
TASK - Confirmation for	Greetings from TASK!	04.12.2020
Jio Drive -200 nos Home Sales Officers in Jio !!!	As discussed, Jio is ready to conduct a drive in Sangareddy, Kindly confirm your willingness to host the drive in your college.	
	Jio is having 200+ positions opened in JioFiber Business and attached the JD and all positions are for Hyd city (enclosed for your reference) -Its an <u>ONROLL</u> <u>OPPORTUNITY IN JIO</u>	

	to hav to SAI	uld request necessary l e SMART Phone Must .ES minimum 6 month <mark>)T OFFERSSPOT O</mark>	and candidates incli s - <mark>SPOT OFFERS</mark>		
Infosys NEG Campus Recruitment Program 2021 : Guidelines for Infosys Online Test and FAQ's	can Pro Afte regi nex The whi on I Info sche can is S One can fror date logi	e first stage of the prog didates have registered cess on InfyTQ. er the student has suce stering on InfyTQ only tstage else not eligible next stage of the prog ch registered candidat infyTQ. osys has sent an email a eduling Infosys online didates. The last date f aturday, December 5, 2 ee the slot has been suce didates will receive an n Talent.Acquisition@i e and time of their onli n details.	d for Infosys Recruit ssfully completed y then will move on t to take the online Te ram is Infosys online es need to book a tes about the process of test to all the register or booking a test slot 2020. eccessfully booked, email nfosys.com confirmi ne test along with a l	ment to the st. e test for st slot red t ng the link and	04.12.2020
	tion	Skill tested	Number of questions	Time allocate	
	tion I	Reasoning Ability	15	30 minut	
	tion II	Mathematical Ability	10	30 minut	
	tion	Verbal Ability	20	25 minut	
	tion	Numerical Puzzle Ability	4	15 minut	
	The	al duration of the test i re are some mandator uirements for students	y system and location		

	are the guidelines for Infosys Online Test	
	For more questions related to Infosys online test that you might have, attached are the FAQs.	
	If any malpractice or violation of instructions is detected during the test, Infosys will not move ahead with the application of that candidate.	
	A sample test will be available on the InfyTQ Infosys Recruitment Process homepage of the candidates who successfully scheduled their online test.	
	Candidates who clear the Infosys online test will qualify for the virtual interview round.	
	Incase if the student doesn't have PC / Laptop the student can take test in their respective colleges with the approval of their colleges as per the system and location requirements provided in the Guidelines.	
	Request you to share this Communication with students.	
New Job	Greetings from TASK!	02.12.2020
Requirement for students Graduated in	Please find the requirement from below companies for 2020 graduates.	
2020	Broadridge: https://www.task.telangana.gov.in/Placements/Broadridge_ Financial_Solutions_(India)_Private_Limitedn	
	Selectys https://www.task.telangana.gov.in/Placements/Selectsys- _2020	
	Greetings from TASK!	26.11.2020

entrepreneurshi p program with TASK	The i4TS program for entrepreneurship development is all set to launch on 2nd December 2020. We request you to circulate the registration link among the student community. Program details: i4TS is a 4 to 5 months complete entrepreneurship training program to promote social innovation and rural entrepreneurship in 5 thematic areas - Rural Healthcare, Rural Education, Rural Logistics and Access, Green Economy, Agriculture. The program involves the following stages: Stage 1: 25-hour Entrepreneurship Mindset Development Program, starting on December 2nd (2.5 hrs live session on all the 5 Wednesdays in December and 10 hours of home assignments) for 3000 to 5000 youth Stage 2: Ideathon, mentoring by experts and Demo Day to present start-up ideas Stage 3: 3-months Entrepreneurship Development Program for the best 300 ideas Stage 4: Best 100 ideas to be provided Incubation support with the best incubators in the state Eligibility: All youth between 18 and 25 years of age Focus: Social Innovation and Rural Entrepreneurship Registration form: https://forms.gle/XmsLdatGWgYgYe5GA Last time and date for registration: 2.00 PM, 30th November 2020 The dates on Ideathon and any change, in case, in the Entrepreneurship Development Program will be informed later.	
Infosys Campus Recruitment Program -B.Sc	Greetings from TASK! Infosys has come up with requirement for B.Sc,BCA students graduating in 2021 Please find eligibility as below	20.11.2020

and BCA	Role: Operations Executive
students graduating in	Compensation Offered: INR 2,22,000 per annum
2021	Eligibility Criteria:
	 BCA or B.Sc. (Computer Science / Electronics / Mathematics / Physics / Statistics / Information Technology / Information Science only) Candidates must be graduating from the 2021 batch Candidates should not have any active/standing backlogs Simple average aggregate of 60% or CGPA 6 throughout Class X, XII & Graduation Candidates should not have participated in the Infosys Ltd and/or Infosys Group Company (such as – Infosys BPO) selection process in the last 6 months. Candidates should have excellent communication skills. Candidates should be willing to relocate and work in a 24x7 environment. Important, please read carefully: Please note the minimum academic criteria enables the students only to apply for the job and does not guarantee confirmation to participate or get shortlisted for the placement drive. Post the cutoff date as specified on this page, Infosys at their own discretion will analyze the data of all students who have applied and decide on the final academic percentage /GPA cut-off which may be higher or equivalent to minimum academic criteria specified and shall send out invite letters for the placement drive. The details of all students shortlisted by Infosys will also be put up on this page.
	JD is Published on TASK portal and Students can click on https://forms.gle/yhoLoa5WZGCinU7D7 to apply

(1	
	Last Date to Apply: 11:00 am , 23rd November 2020	
	Request you to encourage all TASK registered eligible students to apply	
Job openings -	Greetings from TASK!	13.10.2020
New requirements for 2020 graduates	Please find the links of new Job openings from the below companies for 2020 graduates.	
	Doyen Solutions - https://www.task.telangana.gov.in/Placements/Doyen_Sol utions2020	
	AIR Worldwide(Degree) - https://www.task.telangana.gov.in/Placements/AIR_World wide2020_DSA	
	AIR Worldwide(Engineering) - https://www.task.telangana.gov.in/Placements/AIR_World wide2020_CAT	
	Please note that the last date to apply for AIR worldwide is tomorrow, 14th Oct.	
	Greetings from TASK!	09.10.2020
TASK - Online	As you avone that TCC has an accurate 1 Company II'm of	
Training Program for TCS National Qualifier Test	As you aware that TCS has announced Campus Hiring for YoP 2021 students. In this regard TASK is planning to start the training program for TASK registered students to attend for the TCS National Qualifier Test (NQT) is scheduled on 24th/25th/26th October 2020.The NQT Scored secured by a student would be considered for TCS Hiring Process.	

	Details of the Training (TCS NQT Pattern): -	
	 6 days @ 3 Hours/day (12th to 16th and 19th October 2020, 2.00 PM to 5.00PM) During the training period the following topics will be covered. 1. Verbal Ability 2. Reasoning Ability 3. Numerical Ability 4. Programming Logic 5. Coding 	
	I request you to send the following link to TASK registered students and ask them to register for this training program.	
	https://docs.google.com/forms/d/e/1FAIpQLSeGuPsIzau2w OCbBy6ceMNjMUjIQzihdFTnJaUdyX1zlJo_Gw/viewform	
	Greetings from TASK!	09.10.2020
Placement - Job Requirements for 2020	Please find the links of new requirements from the below companies for 2020 graduates. Kindly ask the students to apply the jobs.	
graduates	Raam Group - https://www.task.telangana.gov.in/Placements/Raam_Gro upRM_2020_new	
	Copart - https://www.task.telangana.gov.in/Placements/Copart_Ind ia_Pvt_Ltd2020	
	Zensar - https://www.task.telangana.gov.in/Placements/Zensar _2020	

TASK - Extension of	Greetings from TASK!	10.09.2020
College Registration notification for the academic year 20-21	Attached TASK - Extension of College Registration notification for the academic year 20-21. Kindly complete the renewal process.	
TASK - College Renewal Notification released for the academic Year 2020-21	 Greetings from Telangana Academy for Skill and Knowledge (TASK). We hope you and your family are doing well and taking all necessary precautions under this Pandemic Period. Thank you for being with TASK in skilling the youth of Telangana. We have successfully completed the academic year 2019-20. Now college renewal notification released for the year 2020-21. APPLY Online with e-Payment through: https://task.telangana.gov.in/College-Registration Note: Attached College Registration Notification For Academic Year 2020-21 for your reference. For any queries please feel free to contact me. 	10.08.2020
TASK - Session on Expectations of the Infrastructure Sector Post COVID19 on 16th July 2020	In the times of COVID19, when employment opportunities are scare and competition is more, how should students prepare themselves to face the job market? To bring clarity on the employment scenario in the Infrastructure Sector, TASK brings you a Webinar by industry experts titled "Expectations of the Infrastructure Sector Post COVID19" Please block your date and time for this enlightening webinar. Date: 16 th July 2020 Time: 3.00 PM to 5.00 PM The joining link will be shared shortly.	13.07.2020

TASK -	Greetings from TASK!	01.07.2020
Communication		
skills - Beginner	Course: Communication skills - Beginner.	
from 6th to 8th	This course is open to all the registered students of all	
July 2020 for	years.	
Student	Date: 6th to 8th July 2020.	
Registration	Duration: 90 minutes	
	Number of students per batch: 300	
	Timings for Slot : 12.30 PM to 2.00 PM	
	Platform: Octa	
	Here is the google form link for the slot students to apply.	
	Kindly share the link to the students.	
	Please inform the students to check the confirmation email	
	to their inbox one hour before the scheduled time for login	
	credentials.	
	https://forms.gle/j2WWb57cNWT9Z9Gd9	
	The link shall be closed by 3rd July at 4:00 PM	
	Please feel free to contact me if you need	
	any further information.	
	Greetings from Telangana Academy for Skill and	26.06.2020
Request to fill	Knowledge (TASK)!	
the Feedback	We hope you and your family are doing well and taking all	
Form	necessary precautions under this Pandemic Period.	
	TASK would like to know your feedback and suggestions	
	after the post COVID-19. Request you to please fill the	
	below link.	
	https://docs.google.com/forms/d/e/1FAIpQLSd0c6QdRIVM	
	<u>I2NGz9UdUhUnovyn5ZHpFyz5JVosLgbhGqxCzQ/viewfor</u> <u>m?usp=sf_link</u>	
TASK - Schedule	Greetings from TASK.	19.06.2020
inter oureaute		17.00.2020

for Essential Skills Training	TASK would like to conducting online sessions on Essential
from 22nd to	Skills for our TASK registered students with interactive
24th June 2020	methodologies. With jobs being scarce, experts say that
for Slot-1 Reg-	essential skills will play an important role in the job
U	market.
	We have redesigned the entire essential skills content and
	structure to suit online course delivery. Twelve skill-
	specific modules have been identified as below:
	1. Communication skills
	2. Learning to learn
	3. Problem-solving and decision making
	4. Interpersonal skills
	5. Networking skills
	6. Self-empowerment
	7. Performance management
	8. Presentation skills
	9. Teamwork
	10. Leadership skills
	11. Interview skills
	12. Effective writing skills
	Each of these modules, except Interview Skills, will be
	offered in 3 levels - Beginner, Intermediate, and Intensive.
	In this quarter (July to August) only Beginner level will be
	offered. From the next semester, Intermediate and Intensive
	will also be added. Each course will be scheduled multiple
	times in the year so that the maximum number of students
	can get a chance to attend each course.
	These courses are open to all students of all branches of all
	years. Students have to complete the levels one after
	another, starting from the Beginner level.
	Interview skills is only for the final year students or III year
	second-semester students. This is offered in two levels.
	Starting from 22nd June, we are going to offer 4 courses
	every week (each course is for 3 days with 90 minutes
	session per day). We are starting with the Beginner Level.
	Courses will be conducted in two time slots each day, as

	mentioned in the calendar.	
	Please see the attached calendar to know the weekly allotm ent of courses. Students can choose any one of the courses and apply for it through the Google form link given. Right now, we are taking applications only for the courses offered from 22nd to 24th June.	
	Courses will be offered through Octa Work platform. Students can log in through their hall ticket numbers. A standard common password will be shared with all the students for login access. Once they log in, they can change their password and use the same login and password for all future courses.	
	Course-1: Communication skills - Beginner Course-2: Interview skills Level 1	
	Date: 22 nd to 24 th June 2020 Duration: 90 minutes	
	Slot Timings: 10.30 AM to 12.00 PM	
	Number of students per batch: 300	
	Here is the google form link for the slot-1 students to apply. It asks for basic details of the students including the choice of anyone course.	
	https://forms.gle/j2WWb57cNWT9Z9Gd9	
	For further clarifications please feel to call back.	
Just Dial recruitment for	Greetings from TASK!	10.03.2020
B.E/ B.Tech, B.Com, B.Sc, B.A, MBA (any	Just dial has recruitment for B.E/ B.Tech, B.Com, B.Sc, B.A, MBA (any discipline) graduated in 2019 & 2020 at Master's Degree College, Sangareddy	
discipline) graduated in 2019 & 2020	Website: http://www.justdial.com	
	Domain: Sales & Marketing	

Designation: Business Development Executives
 Eligibility Criteria: B.E/ B.Tech, B.Com, B.Sc, B.A, MBA (any discipline) 2020 graduates with 50% throughout the academics and no back logs only need to apply. Only Male Candidates need to apply
CTC per Annum: INR 2,59 ,000 (for Remote cities) - INR 2,90,000 (for Hyderabad) + Incentives + Insurance Coverage (Personal accidental insurance +Medical Insurance) and other benefits. Only Freshers are eligible for Salary revision Post 6 months by Re 2000 & an antillement of Repuse amount after and of 6
by Rs 3000 & an entitlement of Bonus amount after end of 6 and 12 months which is linked to the performance.
 Job Description: Need to approach Business Establishments of an area and capture all their data. Convincing the Customer for listing in Just Dial (Sales) Meeting daily/ weekly/ monthly targets. Giving presentation to the clients about the company and other services. Explaining the concepts and benefits of listing in Just dial and need to get Contracts (Business). Following up with the client for Sales closure. Meeting daily/ weekly/ monthly targets
 Bond Period: NA Skills Required: Excellent communication skills (Written and verbal), ability to communicate correctly and clearly with all customers. Basic knowledge of methods involved in promoting and selling products or services. Sales strategies and tactics, product demonstration and sales techniques. Fluency in English, Telugu, and Hindi. Work successfully in a team environment as well as independently, ability to successfully adapt to changes in their environment.

	 Computer knowledge/ Skills, Ability to use a desktop computer system. Demonstrable conflict resolution, negotiating, and de-escalation skills Ability to determine customer needs and provide appropriate solutions Ability to approach problems logically & rationally, action-oriented, organized and self-disciplined Qualified candidates will be comfortable in a multitasking, high-energy environment. They will be a creative and analytical problem solvers with a passion for excellent customer service. Selection Process: 1st round: Online Test 2nd round: HR interview 4th round: Managerial Intervieww 	
	Please carry an updated Resume, 2 Passport size photographs and Government-approved (original & Photocopy) identity proof.	
	For More details:	
	Please click on https://www.task.telangana.gov.in/Placements/Justdial- _Business_Dev_Ex_2020	
Confirmation - Introduction to Python - Tara	The below training schedule is confirmed and the same updated on the training calendar.	07.02.2020
Govt. Degree	Introduction to Python	
College, Sangareddy	Duration : 2 Days	
	Tara Govt. Degree College, Sangareddy Dates : 13th and 14th Feb 2020	
	Instructor : Mr.Jaffer Pasha Contact No : 8886220324	
	Attached are the attendance and feedback forms. Kindly	

	Submit the same.	
	Pre- Requisites : 1.Computer Lab with High Speed Internet facility 2.Each student should be allocated one single system.	
Deadline Approaching to apply for 500 Scholarship from Facebook for students	 February 12th is the last date for students to apply for the 500 scholarships from Facebook's School of Innovation programme to learn Augmented Reality from industry experts. You may please share this information with students from your institute so that the interested students may apply and benefit from this industry led learning experience. 	06.02.2020
	Last date to apply: February 12th 2020. To learn more, visit www.sv.co/fb <u>/apply</u> Thanks,	
Requirement from ELICO for 2020 Graduates	Greetings from TASK!! We would like to inform you that TASK received a requirement from ELICO company for 2020 graduating students. Kindly ask the student ti apply on below links. Please find the below links: https://www.task.telangana.gov.in/Placements/ELICO- _AR_Calling_2020 https://www.task.telangana.gov.in/Placements/Elico- _Medical_Coder_2020	08.01.2020

MPC Sessions	Greetings from TASK!	06.01.2020
from 6th Jan	TACK would like to inform you that we have asheduled	
2020	TASK would like to inform you that we have scheduled Mahindra Pride Training Session from 6th Jan 2020 with	
	below trainer, Kindly coordinate the program.	
	Tara Government College - Mr.Thomas	
Bulk Student	Greetings from TASK!!	27.11.2019
Registration		
	We have successfully updated Students data in TASK website. Please inform to the students to update their	
	profiles within next 6 working days i.e (03-Dec-2019).	
	Student login details are:	
	Username :Hall Ticket Number (without special characters)	
	Password : Task@2019 (common for all students)	
Excel	Dear Ramana,	20.11.2019
	Kindly fill the format.	
Proceedings-List	Respected Sir/ Madam, Please find the attachment-Proceedings-List of nominated	19.11.2019
of nominated	candidates for Infosys Project Genesis-Faculty	
candidates for	Development Program on GBFS between 28th November to	
Infosys Project	6th December 2019 to be held at Infosys SEZ, Pocharam	
Genesis-Faculty	Campus, Hyderabad.	
Development		
Program on		
GBFS between		
28th November		
to 6th December		
2019 to be held		
at Infosys SEZ,		
Pocharam		
Campus,		

Hyderabad.		
Students Registration Notification 2019-2020	Greetings from TASK!! TASK has been striving to enhance the employability quotient of the youth and enabled them choose multiple avenues as they graduate. We have added new services for the academic year 2019-20 to enrich the students in technical skills and Soft Skills. We are glad to inform you that TASK has opened the student registrations across the state for the Academic year 2019-20 from 1st July 2019 to 31st August 2019. (Late fee of Rs.100/- +Tax is applicable from 1st September - 09th September 2019). The services offered through the Telangana Academy for Skill and Knowledge (TASK) has been detailed out in the website (Course Catalogue). For more information, please visit TASK websitewww.task.telangana.gov.in. Note: Attached Student Registration Notification.	22.08.2019
Proceedings-OD Approval-List of Faculty Faculty from GDCs-To attend FDP - TASK-Infosis from 5th to 13th August, 2019	Respected Madam/Sir, Please find the attachment of Proceedings-OD Approval- List of Faculty Faculty from GDCs-To attend FDP -TASK- Infosis from 5th to 13th August, 2019	03.08.2019

PROCEEDINGS OF THE COMMISSIONER OF COLLEGIATE EDUCATION GOVERNMENT OF TELANGANA, HYDERABAD Present: Sri. Navin Mittal, IAS.

- Sub: Collegiate Education Government Degree College, Narayankhed, Sangareddy Dist. - Permission granted to certain faculty members from various GDCs to work in GDC Banswada, in view of the NAAC Peer Team visit to GDC Narayankhed - On Duty Orders issued - Reg.
- Ref: Letter received from Principal, Government Degree College, Narayankhed. Dated: 07.09.2022.

In pursuance of the circumstances informed in the reference read above, the Commissioner of Collegiate Education, Hyderabad, has accorded permission to the following faculty members to work in GDC Narayankhed from **07.09.2022** to **13.09.2022** in view of the NAAC Peer Team visit to GDC Narayankhed, scheduled on the dates 12.09.2022 to 13.09.2022.

S.No	Name of the Faculty member	Subject	Present working place
1.	A. Rajendhar	Mathematics	GDC Kamareddy
2.	Md. Sharif Miya	English	GDC A Sangareddy
3.	S. Ramulu	Economics	GDC Zaheerabad
4.	Dr. Mithun Kumar Rathod	Zoology	GDC Sangareddy
5.	K. Krishna Priya	History	GDCW Sangareddy
6.	Giridhar Gopal	Librarian	GDC A Sangareddy
7.	P. Supriya	Physical Director	GDC Jogipet

The absence of the above said faculty members in the colleges on the dates mentioned above shall be treated as On Duty and they are not eligible to claim TA/ DA from the colleges. Further, in view of the above, the concerned Principals are informed to relieve the above mentioned faculty members to attend the above said duty on the above said dates.

(Orders of the CCE have been obtained in the Note file)

Signed by D Thiruvengala Chary Date: 08-09-2022 13:45:28 Reason: Approved For Commissioner of Collegiate Education

File No.CCE-AC/GEN/53/2021-ACADEMIC CELL

To Principal GDC Narayankhed Copy To Principal GDC Kamareddy Principal GDC A Sangareddy Principal GDC Zaheerabad Principal GDC W Sangareddy Principal GDC Jogipet

File No.GDCNKD-ESTT/75/2022-O/o PRINCIPAL-GDC-NKD-CE

PROCEEDINGS OF THE PRINCIPAL, GOVT. DEGREE COLLEGE, NARAYANKHED

PRESENT: Sri.K.Venkateshwarlu, M.Sc., B.Ed.,

- Sub: Collegiate Education GDC, Narayankhed, District Sangareddy Certain faculty members to work in GDC, Narayankhed from various GDCs on OD basis in view of the NAAC Peer Team visit scheduled from 12.09.2022 & 13.09.2022 – Relieving Orders – Issued.
- Read: CCETS, Hyderabad, Proc. File No.CCE-AC/GEN/53/2021-ACADEMIC CELL, Dated: 08.09.2022.

0 0 0

<u>O R D E R:-</u>

In pursuance of the orders issued by the Commissioner of Collegiate Education, TS, Hyderabad under reference read above, the following faculty members has been attended their duties at Government Degree College, Narayankhed, District Sangareddy **from 07.09.2022 to 13.09.2022** in view of the NAAC Peer Team visit scheduled from 12.09.2022 & 13.09.2022 as **ON DUTY** basis.

SI. No.	Name of the Employee	Subject	Name of the College
1	Sri. Md. Sharif Miya	English	Tara GDC, Sangareddy
2	Dr. Mithun Kumar Rathod	Zoology	Tara GDC, Sangareddy
3	Sri. Giridher Gopal	Librarian	Tara GDC, Sangareddy
4	Sri. K. Krishna Priya	History	GDCW, Sangareddy
5	Sri. S. Ramulu	Economics	GDC, Zaheerabad
6	Smt. P. Supriya	Physical Director	GDC, Jogipet
7	Sri. A. Rajendhar	Mathematics	GDC, Kamareddy

Therefore, the above faculty members are hereby relieved from their duties on **13.09.2022 A/N**.

Signed by Venkateshwarlu Kandukuri Daterin(1941-, 909, NRA/ANRHED 9:15:22 Reason: Approved

То

Principal, Tara GDC (A), Sangareddy

Principal, GDCW, Sangareddy

Principal, GDC, Zaheerabad

Principal, GDC, Jogipet

Principal, GDC, Kamareddy.





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D-122, Phase-III, IDA., Jeedimetla, Hyderabad - 500 055. Telangana., India

Phone : +91-40-23090147, Fax : +91-40-23091549, e-mail : basrtech@basrtech.com, www.basrtech.com

This is to certify that following students of Tara Govt. College, Sangareddy (A) have successfully completed the One-Week internship at our commercial laboratory facility on "CHEMICAL INDUSTRIAL SAFETY MANAGEMENT".

S.No.	Name of the Student	Class/Group	Roll Number
1	Aaluri Madappa	B.Sc.BtBC-III	6058-19-572-001
2	Agnikula Kshatriya Vikranth Singh	B.Sc.BtBC-III	6058-19-572-002
3	Bagili Pavan Kumar	B.Sc.BtBC-III	6058-19-572-003
4	Errolla Vivek Teja	B.Sc.BtBC-III	6058-19-572-004
5	Golla Sheshikanth	B.Sc.BtBC-III	6058-19-572-005
6	Kallem Sai Prasanna	B.Sc.BtBC-III	6058-19-572-006
7	Kurma Anil Kumar	B.Sc.BtBC-III	6058-19-572-007
8	Maldoddi Nikitha	B.Sc.BtBC-III	6058-19-572-008
9	Naikoti Sai Teja	B.Sc.BtBC-III	6058-19-572-010
10	Rathod Meenakshi	B.Sc.BtBC-III	6058-19-572-012
11	Sanadhi Laxman	B.Sc.BtBC-III	6058-19-572-013
12	U Navaneetha	B.Sc.BtBC-III	6058-19-572-014
13	Uppari Vamsi	B.Sc.BtBC-III	6058-19-572-015

BATCH-I

ВАТСН-П

S.No.	Name of the Student	Class/Group	Roll Number
1	Solkampally Niharika	B.Sc.MZC-III	1701-6058-457-046
2	Amrin	B.Sc.MZC-III	6058-19-457-001
3	Angoth Sangeetha	B.Sc.MZC-III	6058-19-457-002

4	Chakali Yathish Chandra	B.Sc.MZC-III	6058-19-457-004
5	Dhannaram Sudhakar	B.Sc.MZC-III	6058-19-457-006
6	Erigipally Shivaleela	B.Sc.MZC-III	6058-19-457-007
7	Kunadoddi Shalini	B.Sc.MZC-III	6058-19-457-009
8	M Pragna	B.Sc.MZC-III	6058-19-457-011
9	Mudgula Ruchitha	B.Sc.MZC-III	6058-19-457-012
10	Nallolla Ranjitha	B.Sc.MZC-III	6058-19-457-013
11	Neeradi Anitha	B.Sc.MZC-III	6058-19-457-014
12	Nelapati Vijaya	B.Sc.MZC-III	6058-19-457-016
13	Pathri Kiran Kumar	B.Sc.MZC-III	6058-19-457-017
14	Patnam Prashanth Kumar	B.Sc.MZC-III	6058-19-457-018
15	Peddasayeb Praveen Kumar	B.Sc.MZC-III	6058-19-457-019
16	Perka Sai Charan	B.Sc.MZC-III	6058-19-457-020
17	Ponna Srija	B.Sc.MZC-III	6058-19-457-021
18	Puramola Madhavi	B.Sc.MZC-III	6058-19-457-022
19	Ravula Abhishekasumalatha	B.Sc.MZC-III	6058-19-457-023
20	Regoti Vamshi Priya	B.Sc.MZC-III	6058-19-457-024
21	Sangeetham Vyshnavi	B.Sc.MZC-III	6058-19-457-025
22	Shaik Arshad	B.Sc.MZC-III	6058-19-457-026
23	Thammali Eshwar	B.Sc.MZC-III	6058-19-457-027
24	Upthalla Kavya Sri	B.Sc.MZC-III	6058-19-457-028
25	Uroojunnisa	B.Sc.MZC-III	6058-19-457-029
26	Vadla Keerthana	B.Sc.MZC-III	6058-19-457-030

ВАТСН-Ш

S.No.	Name of the Student	Class/Group	Roll Number
1	Aarpula Jeremiah	B.Sc.MCCs-III	6058-19-578-001
2	Aluri Lavanya	B.Sc.MCCs-III	6058-19-578-002
3	Begari Mahesh Kumar	B.Sc.MCCs-III	6058-19-578-004

4	Begari Nikhilesh Babu	B.Sc.MCCs-III	6058-19-578-005
5	Bisa Sravani	B.Sc.MCCs-III	6058-19-578-006
6	Chembeti Srinivas	B.Sc.MCCs-III	6058-19-578-008
7	Dande Anil	B.Sc.MCCs-III	6058-19-578-009
8	Dandu Sai Kiran	B.Sc.MCCs-III	6058-19-578-010
9	Gangeri Laxmikanth	B.Sc.MCCs-III	6058-19-578-012
10	Gudala Omkar	B.Sc.MCCs-III	6058-19-578-014
11	Kammari Manasa	B.Sc.MCCs-III	6058-19-578-015
12	Kemsaram Narasimha	B.Sc.MCCs-III	6058-19-578-016
13	Kolipaka Dheeraj Kumar	B.Sc.MCCs-III	6058-19-578-017
14	Kummari Prashanth Reddy	B.Sc.MCCs-III	6058-19-578-018
15	Kunadoddi Rahul	B.Sc.MCCs-III	6058-19-578-019
16	Lagula Sai Ram Goud	B.Sc.MCCs-III	6058-19-578-020
17	Meedinti Arjun	B.Sc.MCCs-III	6058-19-578-021
18	Mukthikanth Rout	B.Sc.MCCs-III	6058-19-578-022
19	Munugala Binesh	B.Sc.MCCs-III	6058-19-578-023
20	Munugala Madhav	B.Sc.MCCs-III	6058-19-578-024
21	Nethi Shiva Shankar	B.Sc.MCCs-III	6058-19-578-025
22	Nirudi Saimahesh	B.Sc.MCCs-III	6058-19-578-026
23	Pocharam Sujil Kumar	B.Sc.MCCs-III	6058-19-578-027
24	Shankarampet Alekhya	B.Sc.MCCs-III	6058-19-578-028
25	Singitham Shirisha	B.Sc.MCCs-III	6058-19-578-029
26	Sura Raja Ramesh	B.Sc.MCCs-III	6058-19-578-030
27	Talelma Prasannalaxmi	B.Sc.MCCs-III	6058-19-578-031
28	Thammali Chandana	B.Sc.MCCs-III	6058-19-578-032
29	Thammalithota Pranay Kumar	B.Sc.MCCs-III	6058-19-578-033
30	Tudumu Shiva Charan	B.Sc.MCCs-III	6058-19-578-034
31	Turpati Raghavendra	B.Sc.MCCs-III	6058-19-578-035
32	Vadla Prashanth	B.Sc.MCCs-III	6058-19-578-036
33	Vislavath Aravind	B.Sc.MCCs-III	6058-19-578-037

34	Yellayolla Avinash Reddy	B.Sc.MCCs-III	6058-19-578-038
35	Yerrolla Shashi Pranay	B.Sc.MCCs-III	6058-19-578-039
			Ltd.
		For B	ASR FARE CHEMISPIP
		For D	ASR File Chemisals Put. Ltd.

ADSORPTION STUDIES OF Fe[III] ON CELITE IN THE PRESENCE OF D-RIBOSE AS CHELATING AGENT

Dissertation submitted in Partial fulfillment for the requirements for the award of degree of

Bachelor of Science ín CHEMISTRY

By

1.S.	Shirisha,	B.Sc(Chemistry),	III	Year
2.A.	Lavanya,	B.Sc(Chemistry),	III	Year
З.Т.	Prasanna Laxmi,	B.Sc(Chemistry),	III	Year
4.S.	Alekhya,	B.Sc(Chemistry),	III	Year
5.К.	Manasa,	B.Sc(Chemistry),	III	Year
6.В.	Sravani,	B.Sc(Chemistry),	III	Year

Under the Guídance:

Dr. K.ABHIJIT *HEAD* DEPARTMENT OF CHEMISTRY



TARA GOVERNMENT COLLEGE, SANGAREDDY(A)

<u>CERTIFICATE</u>

This is to certify that the project work entitled "ADSORPTION STUDIES OF Fe[III] ON CELITE IN THE PRESENCE OF D-RIBOSE AS CHELATING AGENT" is presented by B.Sc (CHEMISTRY) students in partial fulfillment of the requirements for the degree of Bachelor of Science in Chemistry by the Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

The results embodied in this report have not been to any other University or Institution for the award of any degree.

(Dr. K. ABHIJIT)

Project Supervisor & Head, Department of Chemistry Tara Government College, Sangareddy(A)

EXTERNAL EVALUATOR

ACKNOWLEDGEMENTS

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We wish to express my gratitude to Smt. M.Praveena, Principal, Tara Govt. College, Sangareddy(A)-502001, INDIA for her constant support, cooperation and suggestions during the research work. We wish to express my sincere thanks to K.Sreedhar, Asst. Professor, Department of Chemistry for providing me facilities, help and support for the entire research work.

We express our special thanks to **MART Specialities Lab, Hyderabad** for providing technical assistance in Atomic absorption spectrometric analysis.

Finally, we take this opportunity to thank one and all that has directly or indirectly helped me in completing the task.

1.S. Shirisha, B.Sc(Chemistry), III Year 2.A. Lavanya, B.Sc(Chemistry), III Year 3.T. Prasanna Laxmi, B.Sc(Chemistry), III Year 4.S. Alekhya, B.Sc(Chemistry), III Year 5.K. Manasa, B.Sc(Chemistry), III Year 6.B. Sravani, B.Sc(Chemistry), III Year

<u>DECLARATION</u>

We hereby declare that the project report entitled "ADSORPTION STUDIES OF Fe[III] ON CELITE IN THE PRESENCE OF D-RIBOSE AS CHELATING AGENT" is the work done by us in the campus at Department of Chemistry, Tara Government College, Sangareddy(A) during the academic year 2021-2022 and is submitted in partial fulfillment of the requirements for the degree of Bachelor of Science by Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

S.No.	Name of the Student	Roll Number	Group	Year	Signature
1	S.SHIRISHA	605819578029	B.Sc (MCCs)	III	
2	A.LAVANYA	605819578002	B.Sc (MCCs)	III	
3	T.PRASANNA LAXMI	605819578031	B.Sc (MCCs)	III	
4	S.ALEKHYA	605819578028	B.Sc (MCCs)	III	
5	K.MANASA	605819578015	B.Sc (MCCs)	III	
6	B.SRAVANI	605819578006	B.Sc (MCCs)	III	

TARA GOVERNMENT COLLEGE, SANGAREDDY – 502 001 (AUTONOMOUS)

BONAFIDE CERTIFICATE

Certified that the project report" ADSORPTION STUDIES OF Fe[III]

ON CELITE IN THE PRESENCE OF D-RIBOSE AS CHELATING AGENT" is

the bonafide work of

1.S.	Shirisha,	B.Sc(Chemistry),	III Year
2.A.	Lavanya,	B.Sc(Chemistry),	III Year
З.Т.	Prasanna Laxmi,	B.Sc(Chemistry),	III Year
4.S.	Alekhya,	B.Sc(Chemistry),	III Year
5.К.	Manasa,	B.Sc(Chemistry),	III Year
6.В.	Sravani,	B.Sc(Chemistry),	III Year

who carried out the project work under my supervision.

Dr.K.ABHIJIT **PROJECT SUPERVISOR** Smt. M.PRAVEENA PRINCIPAL

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Chapter-I

INTRODUCTION

Adsorption

Adsorption is a process in which solids come into contact with liquids or gases, and the mass transfer occurs from liquids to solids. Desorption is the reversal of this action. Adsorption operations take advantage of a solid's capacity to concentrate certain chemicals from a fluid on to its surface. Adsorbate refers to the adsorbed substance, while adsorbent refers to the solid substance. The following are some examples of solid-liquid and solid-gas applications:

- Removing dissolved moisture from gasoline.
- Decolorization of sugar solutions and petroleum products.
- Removing noxious odors and tastes from water. Dehumidification of air and gases is one of the solid-gas activities.
- To fractionate mixtures of hydrocarbon gases such as methane, ethane, and propane.
- To remove undesirable odors and contaminants from gases.
- To recover valuable solvent vapors from dilute gas mixtures.

Nature of adsorbents

Adsorbents are typically granular in nature, ranging in size from 0.5 mm to 12 mm. They can't have a lot of pressure decrease or get swept away by a fast-moving stream. During handling, they must maintain their shape and size. They'll need a lot of pores and a lot of surface area per unit mass.

Sl. No.	Adsorbent	Source	Application
1.	Fuller's earth	Naturally occurring clay is heated and dried to get a porous structure.	De-colorizing, drying of lubricating oils, kerosene and engine oils.
2.	Activated charcoal	Bentonite or other activated clay which are activated by treatment with sulfuric acid and further washing, drying and crushing.	Used for de-colorizing petroleum products.
3.	Bauxite	A naturally occurring hydrated alumina, activated by heating at 230-815	Used for de-colorizing petroleum products and for drying gases.

Some of the commonly used adsorbents, their sources and applications are given below:

4.	Alumina	A hard hydrated aluminium oxide, which is activated by heating to drive off the moisture and then crushed to desired size.	Used as desiccant.
5.	Bone-char	Obtained by destructive distillation of crushed bones at 600-900	Used for refining sugar and can be reused after washing and burning.
6.	Silica gel	A hard granular and porous product obtained from sodium silicate solution after treatment with acid. Normally has 4 to 7% water in the product.	Dehydration of gases and liquids, and separation of gas-liquid hydrocarbon mixture.

7.	Activated carbon	 (1) Vegetable matter is mixed with calcium chloride, carbonized and finally the inorganic compounds are leached away. (2) Organic matter is mixed with porous pumice stones and then heated and carbonized to deposit the carbonaceous matter throughout the porous particle. (3) Carbonizing substances like wood, sawdust, coconut shells, fruit pits, coal, lignite and subsequent activation with hot air steam. It is available in granular or pellated form. 	De-colorizing of sugar solutions, chemicals, drugs, water purification, refining of vegetable and animal oils, recovery of gold and silver from cyanide ore-leach solution, recovery of solvent vapour from gas-mixtures, collection of gasoline hydro- carbons from natural gas, fractionation of hydrocarbon gases.
8.	Molecular sieves	These are porous synthetic zeolite crystals, metal alumino-silicates.	Dehydration of gases and liquids, and separation of gas-liquid hydrocarbon mixture.

Significance of Research problem

Adsorption of heavy metals is an important strategy to develop newer remediation technologies for the sustainable environmental protection. But the efficacy of adsorption of heavy metals under the normal conditions using suitable adsorbent depends on several factors which need to be finely tuned to get efficient adsorption process. In the presence of proper facilitating agents, the adsorption of heavy metals enhanced which will certainly improves the existing heavy metal techniques.

Objective of the Project

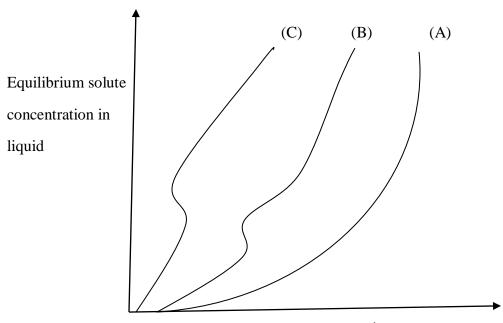
To evaluate the impact of D-Ribose as chelating agent in the adsorption of Fe (III) from aqueous solution by Celite as an adsorbent to develop efficient remediation technology using concept of Coordination chemistry.

Chapter-II

ADSORPTION OF SOLUTE FROM DILUTE SOLUTIONS

Both the solvent and the solute are adsorbed whenever a mixture of solute and solvent is adsorbed using an adsorbent. As a result, only relative or apparent solute adsorption can be determined. As a result, treating a known volume of solution of original concentration Cwith a known weight of adsorbent is standard procedure. Let C* be the solution's final equilibrium solute concentration. If v is the volume of solution per unit mass of adsorbent (cc/g), and C and C^* are the starting and equilibrium concentrations (g/cc) of the solute, then the apparent adsorption of the solute per unit mass of adsorbent is v(C -C*), (g/g), neglecting any volume change. This statement is most useful in the case of dilute solutions. The C* value is determined by the temperature, nature, and properties of the adsorbent when the proportion of the original solvent that can be adsorbed is tiny. The Freundlich adsorption isotherm, $C^*=K[v(C - C^*)]n$, represents the adsorption phenomena in dilute fluids across a small concentration range. The Freundlich adsorption equation is especially useful in situations where the identification of the solute is unknown, such as the removal of colouring substances from sugar solutions, oils, and other liquids. A spectrophotometer or colorimeter can quickly determine the colour composition of the solute. In worked example 2, the interpretation of this data is demonstrated. Adsorption is good if the value of n is high, say 2 to 10. If it's between 1 and 2, it's relatively challenging, and if it's less than 1, it's easy and it indicates poor adsorption characteristics. Freundlich adsorption equation is also useful in such a case where the actual identification of solute is not known, e.g. removal of colouring substance from sugar solutions, oils etc. A typical adsorption isothermal for

the adsorption of various adsorbents A, B and C in dilute solution at the same temperature for the same adsorbent is represented in a graph.



Kg solute apparently adsorbed /kg adsorbent

Adsorption isotherms for various adsorbents

Metal Complexation of A D- Ribose- Based Ligand

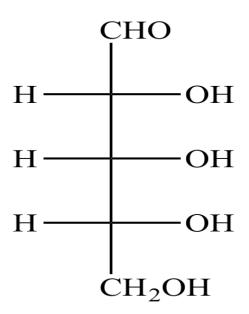
The complexation characteristics of methyl 2,3,4-tri-O-(2-picolyl)—D-ribopyranoside, a novel sugar-derived hexadentate ligand, were investigated using a combination of experimental and theoretical approaches. In the complexes with Mn^{II}, Co^{II}, Ni^{II}, and Zn^{II}, the coordination bond lengths reveal significant departures from perfect octahedral, with deformation towards trigonal-prismatic geometries, indicating a conformationally constrained ligand. The metal-cation–ligand interactions for D-Ribose and its acyclic analogue ligand [1, 2, 3-tri-O-(2-picolyl)-1,2,3-propanetriol] were investigated using spectroscopic techniques and isothermal calorimetric

titrations for the series Mn["], Co["], Ni["], Zn["], and Cu["]. Depending on the nature of the metal, the results show that the complexes produced with D-Ribose are more stable than those obtained with According to molecular modelling studies; the presence of the sugar moiety greatly favours conformations that are compatible with metal binding. In many ways, a close comparision can be sketched between metal complexation and other host-guest interactions. To describe them both, several molecular variables have to be taken into consideration. One of the most complex variables is the relation between the conformational flexibility of the ligand and the binding strength of the metal cation. For ligands with a reduced number of low-energy conformations, relatively straightforward estimations of the binding same wavelength of metal cations as well as a direct interaction between structural and thermodynamic data are available. However, for ligands with a huge number of degrees of freedom, the correct orientation of the chelating groups becomes one of the most important issues in metal-ion identification processes. For coordination chemists, the prediction and understanding about coordination properties of new ligands is one the most challenging tasks. We decided to investigate how entropic and enthalpic variables can impact on metal binding and selectivity for systems with a substantial number of degrees of freedom.

Ferric complexes of sugar-type ligand

Fe(111) complexes formed with sugar-type ligands such as aldoses, ketoses, polyalcohols, sugar acids, di and trisaccharides was prepared. These complexes composition was determined by standard analytical methods. Deprotonated alcoholic hydroxy groups participate in the complex formation which results in the formation of polynuclear species. Mossbauer spectra reflected the presence of high spin iron(111) central atoms.EPR spectra showed antiferromagnetic

interactions between the iron(111) centres in the complexes indicating dimeric or oligomeric complex structures. Depending upon the nature of a ligand as well as on the preparation mode of the complexes the ratio of interacting and isolated iron(111) is calculated. Formation of polynuclear iron(111) complexes of D-fructose, sorbose, lactobionic acid, glucose, galactose, mannose, and lactose were shown by the analysis and structural study by the sugar type ligand with ferric system. Ferric complexes of reducing sugars also contained ferrous species in some cases. The combination of Mossbauer spectroscopic measurements with ERP or magnetic susceptibility studies has led valuable information concerning the intramolecular and electronic structure on which the system has been developed.



Structure of D-Ribose

Celite

Celite is a naturally occurring, soft, siliceous sedimentary rock that can be crumbled into a fine white to off-white powder. It has a particle size ranging from more than 3 μ m to less than 1 mm, but typically 10 to 200 μ m. Depending on the granularity, this powder can have an abrasive feel, similar to pumice powder, and has a low density as a result of its high porosity. The typical chemical composition of oven-dried celite is 80–90% silica, with 2–4% alumina (attributed mostly to clay minerals), and 0.5–2% iron oxide.

Celite consists of fossilized remains of diatoms, a type of hard-shelled protist. It is used as a filtration aid, mild abrasive in products including metal polishes and toothpaste, mechanical insecticide, absorbent for liquids, matting agent for coatings, reinforcing filler in plastics and rubber, anti-block in plastic films, porous support for chemical catalysts, litter boxes, activator in coagulationstudies, a stabilizing component of dynamite, a thermal insulator, and a soil for potted plants and trees like bonsai.

Usages

Filtration: The Celle engineer, Wilhelm Berkefeld, recognized the ability of the celite to filter and developed tubular filters (known as filter candles) fired from celite. During the cholera epidemic in Hamburg in 1892, these Berkefeld filters were used successfully. One form of celite is used as a filter medium, especially for swimming pools. It has a high porosity because it is composed of microscopically small, hollow particles. Celite is used in chemistry as a filtration aid, to increase flow rate, and filter very fine particles that would otherwise pass through or clog filter paper. It is also used to filter water, particularly in the drinking water treatment process and in fish tanks, and other liquids, such as beer and wine. It can also filter syrups, sugar, and honey without removing or altering their color, taste, or nutritional properties.

Catalyst support: Celite also finds some use as a support for catalysts, generally serving to maximize a catalyst's surface area and activity.For example, nickel can be supported on the material—the combination is called Ni–Kieselguhr—to improve its activity as a hydrogenation catalyst.

Adsorbent: Celite is also used as an adsorbent in the process of adsorption.



Powdered Celite

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Chapter-III

ATOMIC ABSORPTION SPECTROMETRY

Atomic absorption spectrometry (AAS) is an analytical technique that measures the concentrations of elements. Atomic absorption is so sensitive that it can measure down parts per billion of a gram (μ g dm⁻³) in a sample. The technique makes use of the wavelengths of light specifically absorbed by an element. They correspond to the energies needed to promote electrons from one energy level to another, higher, energy level.

Atomic absorption spectrometry has many uses in different areas of chemistry.

Clinical analysis: Analysing metals in biological fluids such as blood and urine.

Environmental analysis: Monitoring our environment- eg finding out the levels of various elements in rivers, seawater, drinking water, air, petrol and drinks such as wine, beer and fruit drinks.

Pharmaceuticals: In some pharmaceutical manufacturing processes, minute quantities of a catalyst used in the process (usually a metal) are sometimes present in the final product. By using AAS the amount of catalyst present can be determined.

Industry: Many raw materials are examined and AAS is widely used to check that the major elements are present and that toxic impurities are lower than specified- eg in concrete, where calcium is a major constituent, the lead level should be low because it is toxic.

Mining: By using AAS the amount of metals such as gold in rocks can be determined to see whether it is worth mining the rocks to extract the gold.

How it works

Atoms of different elements absorb characteristic wavelengths of light. Analysing a sample to see if it contains a particular element means using light from that element. For example with lead, a lamp containing lead emits light from excited lead atoms that produce the right mix of wavelengths to be absorbed by any lead atoms from the sample. In AAS, the sample is atomized- i.e. converted into ground state free atoms in the vapour state- and a beam of electromagnetic radiation emitted from excited lead atoms is passed through the vaporized sample. Some of the radiation is absorbed by the lead atoms in the sample. The greater the number of atoms there is in the vapour, the more radiation is absorbed. The amount of light absorbed is proportional to the number of lead atoms. A calibration curve is constructed by running several samples of known lead concentration under the same conditions as the unknown. The amount the standard absorbs is compared with the calibration curve and this enables the calculation of the lead concentration in the unknown sample.

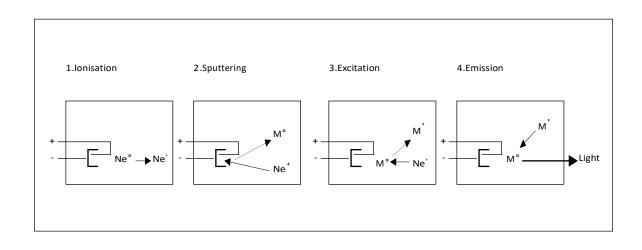
Consequently an atomic absorption spectrometer needs the following three components: a light source; a sample cell to produce gaseous atoms; and a means of measuring the specific light absorbed.

The light source

The common source of light is a 'hollow cathode lamp'. This contains a tungsten anode and a cylindrical hollow cathode made of the element to be determined. These are sealed in a glass tube filled with an inert gas- e.g neon or argon- at a pressure of between 1 Nm⁻² and 5 Nm⁻².

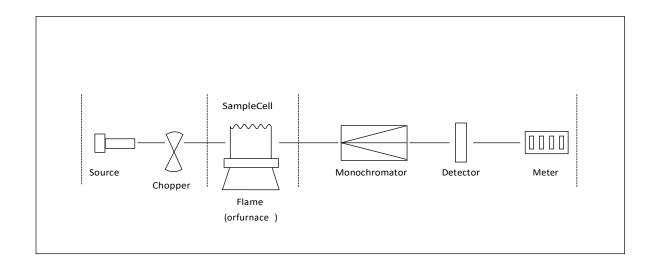


The ionization of some gas atoms occurs by applying a potential difference of about 300-400V between the anode and the cathode. These gaseous ions bombard the cathode and eject metal atoms from the cathode in a process called sputtering. Some sputtered atoms are in excited states and emit radiation characteristic of the metal as they fall back to the ground state $-egPb^* \rightarrow Pb + hv$. The shape of the cathode concentrates the radiation into a beam which passes through a quartz window, and the shape of the lamp is such that most of the sputtered atoms are redeposited on the cathode. A typical atomic absorption instrument holds several lamps each for a different element. The lamps are housed in a rotating turret so that the correct lamp can be quickly selected.



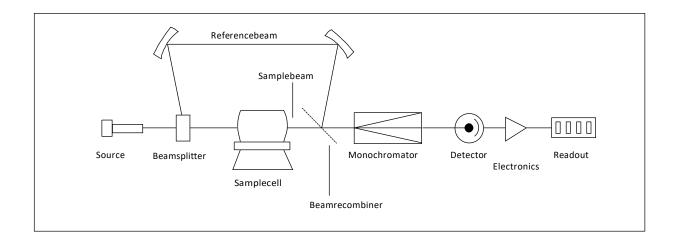
The optical system and detector

A monochromator is used to select the specific wavelength of light –ie spectral line – which is absorbed by the sample, and to exclude other wavelengths. The selection of the specific light allows the determination of the selected element in the presence of others. The light selected by the monochromator is directed onto a detector that is typically a photomultiplier tube. This produces an electrical signal proportional to the light intensity



Double beam spectrometers

Modern spectrometers incorporate a beam splitter so that one part of the beam passes through the sample cell and the other is the reference. The intensity of the light source may not stay constant during an analysis. If only a single beam is used to pass through the atom cell, a blank reading containing no analyte (substance to be analysed) would have to be taken first, setting the absorbance at zero. If the intensity of the source changes by the time the sample is put in place, the measurement will be inaccurate. In the double beam instrument there is a constant monitoring between the reference beam and the light source. To ensure that the spectrum does not suffer from loss of sensitivity, the beam splitter is designed so that as high a proportion as possible of the energy of the lamp beam passes through the sample.



Atomisation of the sample

Two systems are commonly used to produce atoms from the sample. Aspiration involves sucking asolution of the sample into a flame; and electrothermal atomisation is where a drop of sample is placed into a graphite tube that is then heated electrically.

Some instruments have both atomisation systems but share one set of lamps. Once the appropriate lamp has been selected, it is pointed towards one or other atomisation system.

Flame aspiration

Ethyne/air (giving a flame with a temperature of 2200–2400°C) or ethyne/dinitrogen oxide (2600– 2800°C) are often used. A flexible capillary tube connects the solution to the nebuliser. At the tip of the capillary, the solution is 'nebulised' –ie broken into small drops. The larger drops fall out and drain off while smaller ones vaporise in the flame. Only ca 1% of the sample is nebulised.

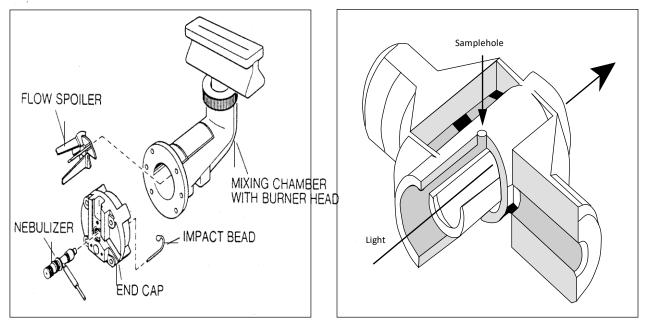


Figure 1

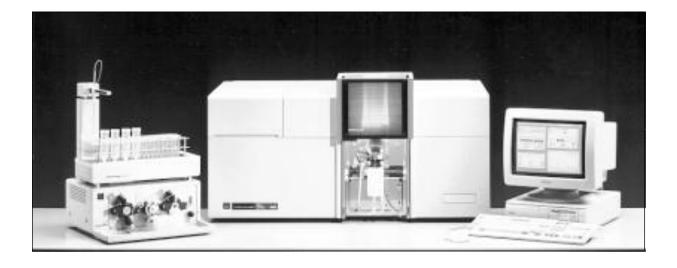
Figure 2

Electro-thermal atomization

25 µl of sample (ca 1/100th of a raindrop) is placed through the sample hole and onto the platform from an automated micropipette and sample changer. The tube is heated electrically by passing a current through it in a pre-programmed series of steps. The details will vary with the sample but typically they might be 30–40 seconds at 150°C to evaporate the solvent, 30 seconds at 600°C to drive off any volatile organic material and char the sample to ash, and with a very fast heating rate (ca 1500 °C s⁻¹) to 2000-2500°C for 5–10 seconds to vaporise and atomise elements (including the element being analysed). Finally heating the tube to a still higher temperature –ca 2700°C – cleans it ready for the next sample. During this heating cycle the graphite tube is flushed with argon gas to prevent the tube burning away. In electrothermal atomisation almost 100% of the sample is atomised. This makes the technique much more sensitive than flame AAS.

Sample preparation

Sample preparation is often simple, and the chemical form of the element is usually unimportant. This is because atomisation converts the sample into free atoms irrespective of its initial state. The sample is weighed and made into a solution by suitable dilution. Elements in biological fluids such as urine and blood are often measured simply after a dilution of the original sample.



When making reference solutions of the element under analysis, for calibration, the chemical environment of the sample should be matched as closely as possible –i.e. the analyte should be in the same compound and the same solvent. Teflon containers may be used when analyzing very dilute aolutions because elements such as lead are sometimes leached out of glass vessels and can affect the results

Background absorption

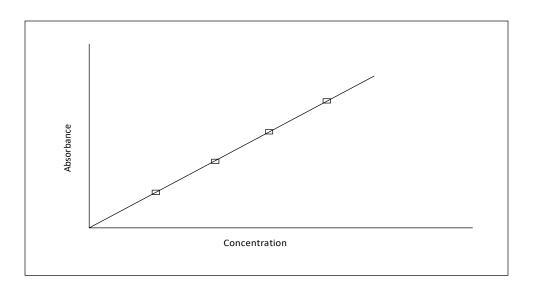
It is possible that other atoms or molecules apart from those of the element being determined will absorb or scatter some radiation from the light source. These species could include unvaporised solvent droplets, or compounds of the matrix (chemical species, such as anions, that tend to accompany the metals being analysed) that are not removed completely. This means that there is a background absorption as well as that of the sample.

One way of measuring and correcting this background absorption is to use two light sources, one of which is the hollow cathode lamp appropriate to the element being measured. The second light source is a deuterium lamp.

The deuterium lamp produces broad band radiation, not specific spectral lines as with a hollow cathode lamp. By alternating the measurements of the two light sources – generally at 50 -100 Hz – the total absorption (absorption due to analyte atoms plus background) is measured with the specific light from the hollow cathode lamp and the background absorption is measured with the light from the deuterium lamp. Subtracting the background from the total absorption gives the absorption arising from only analyte atoms.

Calibration

A calibration curve is used to determine the unknown concentration of an element -eg lead – in a solution. The instrument is calibrated using several solutions of known concentrations. A calibration curve is produced which is continually rescaled as more concentrated solutions are used – the more concentrated solutions absorb more radiation up to a certain absorbance. The calibration curve shows the concentration against the amount of radiation absorbed in the given figure.(a) The sample solution is fed into the instrument and the unknown concentration of the element-e.g. lead- is then displayed on the calibration curve given in the below figure.(b)





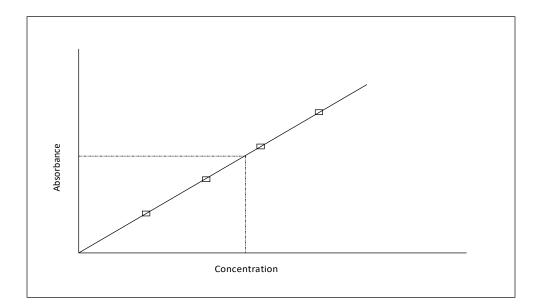


Figure (b)

Interferences and matrix modification

Other chemicals that are present in the sample may affect the atomisation process. For example, in flame atomic absorption, phosphate ions may react with calcium ions to form calcium pyrophosphate. This does not dissociate in the flame and therefore results in a low reading for calcium. This problem is avoided by adding different reagents to the sample that may react with the phosphate to give a more volatile compound that is dissociated easily. Lanthanum nitrate solution is added to samples containing calcium to tie up the phosphate and to allow the calcium to be atomised, making the calcium absorbance independent of the amount of phosphate. With electrothermal atomisation, chemical modifiers can be added which react with an interfering substance in the sample to make it more volatile than the analyte compound. This volatile component vaporises at a relatively low temperature and is removed during the low and medium temperature stages of electrothermal atomisation.

Chapter-IV

METHODOLOGY

MATERIALS REQUIRED

- Ferric chloride Hexahydrate
- D- Ribose
- Celite
- Volumetric flask
- Digital Weighing Machine
- Watch glasses
- Hot air oven
- Ultra pure water (Demineralized)
- 2% of Nitric acid

PROCEDURE

- Using a 250ml Volumetric flask prepare a metal ligand solution by adding 100ppm of Ferric chloride Hexahydrate (MOLYCHEM MCR-11580) and 200ppm of D-Ribose (Avra N2200133). Prepare 250ml solution by adding Ultra pure water and then keep this system aside for a few hours.
- Weigh 5 grams of Celite(Avra N2001901) using a Digital weighing machine (Citizen Scales(I) PVT LTD CTG302-300) and take this into a beaker.

- **3.** Now add 100ml of the above prepared metal ligand solution into the beaker and stir the mixture well for 10 minutes using a glass rod.
- **4.** Keep this mixture aside for 48 hours without disturbing it as at this step Ferric is going to be adsorbed on Celite in the presence of D-Ribose which acts as a chelating agent.
- **5.** After completion of 48 hours take the mixture and filter it off using Whatman Grade 1 filter paper and a funnel.
- **6.** After filtration of the mixture again add Ultra pure water for 3 times and then filtrate it to obtain pure concentration of Ferric which is get adsorbed on Celite.
- 7. Collect the filtered Celite powder and place it on a watch glass and keep this in a Hot air owen at 60 °C for 10 hours to get rid off moisture present in it.
- 8. Now weigh each 1 gm of Celite in glass vials.
- 9. Now take a beaker and rinse it with ultra pure water then followed by Nitric acid.
- **10.** Take 0.5 grams of Celite sample in the beaker and add 2% of Nitric acid and stir the mixture well for 10-15 minutes.
- **11.** Filter the mixture using Whatman Grade 1 filter paper and again 3 times by using Ultra pure water to obtain pure concentration of Ferric present in the mixture prepared using the sample.
- **12.** Take this collected sample solution and keep this system under AAS (Thermo Scientific iCE 3300)
- **13.** Calculate the concentration of Ferric adsorbed on Celite at different ppm levels. Observe the graph obtained and note down the readings of the result we obtained.
- **14.** Same Experiment carried out without the interference/addition of Ligand i.e. D-Ribose for Control Experiment.



Ferric chloride Hexahydrate, D-Ribose, Celite



Preparing 250 ml of metal ligand solution using volumetric flask



Samples with metal ligand solution



Metal ligand solution and celite solution



Addition of Celite to Metal-Ligand Solution





Measuring Flask



Watch glass



Digital Weighing Machine



Hot air Oven



Measuring 100mL of Metal Ligand solution



Celite added to Metal Ligand solution is prepared





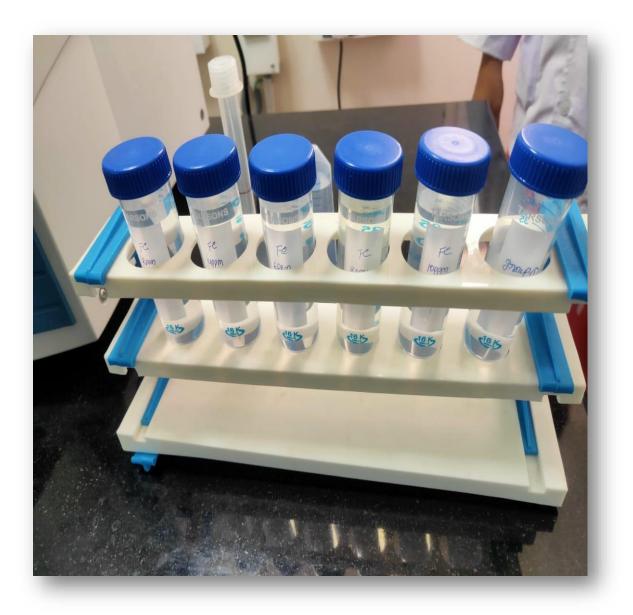
Collecting Celite sample into glass vials



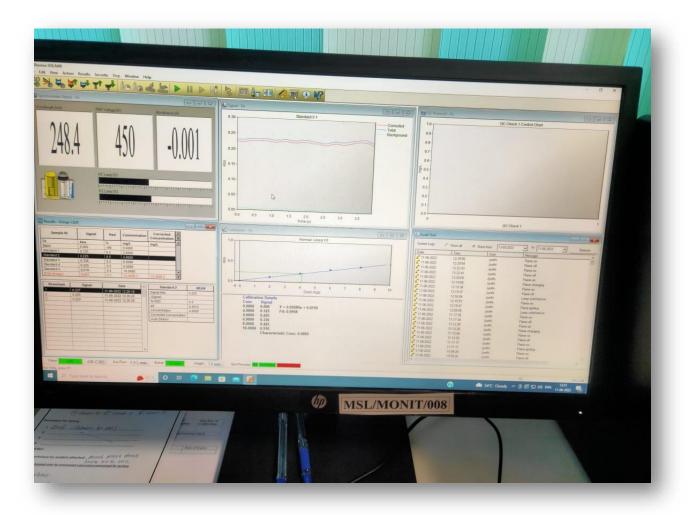
Collected Samples



Atomic absorption spectrometer



Sample added to 2% of Nitric acid

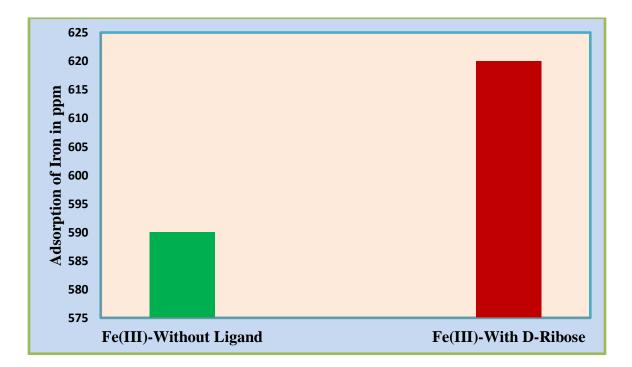


Results obtained on the Computer System under AAS Method

Chapter-V

RESULTS AND DISCUSSION

Celite adsorbs **620** ppm of Iron metal from aqueous solution of Fe(III)-D-Ribose metal ligand solution. Whereas, Celite adsorbs only **590** ppm when D-Ribose is absent. It is evident from the AAS results, ligand involvement enhanced the metal adsorption by initiating potential chemical interactions between adsorbate and adsorbent. D-Ribose firmly interacts with Fe(III) to form a stable complex in aqueous condition. The complex coordination sphere in the resulted complex facilitates strong interactions with the polar points of the adsorbent, Celite. From the AAS results, it is conclusive that **5.08** % of adsorbtion increased in the presence of D-Ribose as chelating agent.



Impact of D-Ribose on Adsorption of Fe (III) ions from aqueous solution by Celite

Spectrometer Parameters-Fe

Element: Fe	Measurement Mode:	Lamp Current: 75%
Wavelength: 248.3nm	Absorbance	Optimise Spectrometer Parameters:
Background Correction:	Bandpass:0.2nm	No
D2	High Resolution: Off	Number Of Resamples: 3
Signal Type: Continuous Measurement Time: 4.0secs Use RSD Test : No	Resamples: Fast Flier Mode: No	

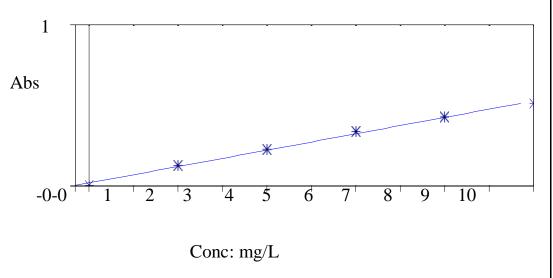
Flame Parameters-Fe				
Flame Type: Air-C2H2 Nebuliser Uptake: 4secs	Fuel Flow: 0.9 L/min Burner Stabilisation :0mins	Auxilary Oxidant: Off Optimise Fuel Flow: No		
Burner Height: 7.0mm	Optimise Burner Height: No	Optimise Fuel Flow. No		

Calibration Parameters	-Fe		
Calibration Mode: Normal	Line Fit: Linear	Use Stored Calibration: No	
Concentration Units: mg/L	Scaled Units: mg/L	Scaling Factor: 1.0000	
Acceptable Fit: 0.990	Rescale Limit : 10.0%	Failure Action: Flag and Continue	
Standard 1 2.0000		Standard 4 8.0000	
Standard 2 4.0000		Standard 5 10.0000	
Standard 3 6.0000			

Solution Results – Fe

Y=0.05089x+ 0.0159 Fit:0.9958

CharacteristicConc:0.0865Normal: Linear Fit



Sample ID	Signal	Rsd	Conc
	Abs	%	mg/L
Fe Blank	0.000	>99	0.0000
1	0.000	Background:	0.000
2	-0.000	Background:	0.000
3	-0.000	Background	0.000
Fe Standard 1	0.125	0.3	2.0000
1	0.125	Background:	0.006
2	0.125	Background:	0.006
3	0.126	Background:	0.006
Fe Standard 2	0.225	0.5	4.0000
1	0.227	Background:	0.007
2	0.225	Background:	0.007
3	0.224	Background:	0.007
Fe Standard 3	0.336	0.2	6.0000
1	0.336	Background :	0.008
2	0.336	Background :	0.008
3	0.335	Background:	0.008
Fe Standard 4	0.425	0.2	8.0000
1	0.426	Background :	0.009
2	0.425	Background:	0.008

3	0.425	Background :	0.009
Fe Standard 5	0.510	0.4	10.0000
1	0.510	Background:	0.009
2	0.509	Background:	0.009
3	0.513	Background:	0.009
Fe B.No.Group-I	0.647	0.3	12.4006C
1	0.647	Background:	0.006
2	0.649	Background:	0.006
3	0.645	Background:	0.006

Celite-Control	0.647	0.3	11.8005C
1	0.614	Background:	0.008
2	0.615	Background:	0.008
3	0.615	Background:	0.008

TEST RESULTS

S.No.	Test Parameter	Sample	Result
01.	Iron by AAS Analysis: (ppm)	Fe(III)-D-Ribose- Celite Sample	620ppm
01.	Iron by AAS Analysis: (ppm)	Fe(III)- Celite Sample (Control)	590ppm

CONCLUSION

From the current project it is clear that D-Ribose as a Chelating agent has played a vital role in adsorption of Fe (III) ions from aqueous solution and increases the adsorption up to **5.08%**. This aspect will be useful in designing the newer strategies of Heavy metal Remediation techniques using Natural Chelating Ligands as Facilitating agents in Metal Adsorption processes.

REFERENCES

- [1] A.Tzschach, K.Ponicke, L.Koreczand K.Burger, J. Organomet. Chem., 59, 199(1973).
- [2] A. D. Buckingham, Can. J. Chem. 1960, 38, 300-307
- [3] A.F.CottonandG.Wilkinson,AdvancedInorganicChemistry,6thedition,JohnWiley&Sons,1999,p.
- [4] A. Martinez, J. Suarez, T. Shand, R. S. Magliozzo and R. A. Sanchez Delgado, J. Inorg. Biochem. 2011, 105, 39-45; S. S. Hosseini, M. Bhadbhade, R. J. Clarke, P. J. Rutledge and L. M. Rendina, Dalton Trans. 2011, 40,506-513.
- [5] A. Bentz, P. Comba, R. J. Deeth, M. Kerscher, B. Seibold and H. Wadepohl, Inorg. Chem. 2008,47,95189527;C.Diedrichand,R.J.Deeth,Inorg.Chem.2008,47,24942506;L.Bonniard,A. delaLande, S. Ulmer, J.-P. Piquemal, O. Parisel and H. Gérard, Catalysis Today 2011, 177, 79-86; Y. D. M. Champouret, J.-D. Maréchal, I. Dadhiwala, J. Fawcett, D. Palmer, K. Singh and G. A. Solan, Dalton Trans. 2006, 2350-2361.
- [6] A. D. Becke, J. Chem. Phys. **1993**, 98,5648-5652.
- [7] B.Lakatos, I.Meisel, A.Rockenbauer, P.SimonandL.Korecz, Jnorg.Chim.Acta, 79, 269(198).
- [8] B.Chiari, O.Piovesana, T.Tarantelliand P.F.Zanazzi, Jnorg. Chem., 22, 2781 (1983), and refs. Therein.
- [9] B. P. Hay and R. D. Hancock, Coord. Chem. Rev. 2001, 212,61-78.
- [10]C.Gessa, M.L.DeCherchi, A.Dessi, S.Deianaand G. Micera, Inorg. Chim. Acta, 80, L53 (1983).
- [11] C. F. Macrae, P. R. Edgington, P. McCabe, E. Pidcock, G. P. Shields, R. Taylor, M. Towler and J. v. d. Streek, J. Appl. Cryst. 2006, 39,453-457
- [12] C. A. G. Haasnoot, F. A. A. M. de Leeuw and C. Altona, Tetrahedron1980, 36,2783-1792.
- [13] C. A. Podlasek, J. Wu, W. A. Stripe, P. B. Bondo and A. S. Serianni, J. Am. Chem. Soc. 1995, 117, 8635-8644.
- [14]C. Lee, W. Yang and R. G. Parr, Phys. Rev. B 1988, 37,785-789.
- [15] D. J. Cram and J. M. Cram, Container Molecules and their Guests, The Royal Society of Chemistry, Cambridge, 1994, p; J. B. Wittenberg and L. Isaacs in Complementarity and Preorganization, (Ed. J. W. S. Philip A. Gale), John Wiley and Sons, 2012.
- [16] D. Cremer and J. A. Pople, J. Am. Chem. Soc. 1975, 97,1354-1358.
- [17] D. E. Wilcox, Inorg. Chim. Acta 2008, 361, 857-867; A. Trapaidze, C. Hureau, W. Bal, M.

Winterhalter and P. Faller, J. Biol. Inorg. Chem. 2012, 17,37-47.

- [18] D. W. Johnson and K. N. Raymond, Supramol. Chem. 2001, 13,639-659.
- [19] E. Chekmeneva, R. Prohens, J. M. Díaz-Cruza, C. Ariñoa and M. Esteban, Anal. Biochem. 2008, 375, 82-89.
- [20] F.W.Lichtenthaler, Acc.Chem.Res.2002,35,728-737; M.J.Climent, A.CormaandS.Iborra, Green Chem. 2011, 13, 520-540.
- [21] F. Bellot, R. Hardré, G. Pelosi, M. Thérisod and C. Policar, Chem. Commun. 2005,5414-5416.
- [22]F. Cisnetti, R. Guillot, G. Pelosi, M. Desmadril and C. Policar, Dalton Trans. 2007,1473-1476.
- [23]F. Cisnetti, R. Guillot, M. Thérisod and C. Policar, Acta Cryst. 2007, C63,m201-m203.
- [24] F.Cisnetti, R.Guillot, N.Ibrahim, F.Lambert, M.Thérisodand C.Policar, Carbohydr. Res. 2008, 343, 530-535.
- [25] F. Cisnetti, R. Guillot, M. Thérisod, M. Desmadril and C. Policar, Inorg. Chem. 2008, 47,2243-2245.
- [26] F. E. Jorge, J. Autschbach and T. Ziegler, J. Am. Chem. Soc. 2005, 127, 975-985; E. I. Stiefel and G. F. Brown, Inorg. Chem. 1972, 11,434-436.
- [27] F. Mohamadi, N. G. J. Richard, W. C. Guida, L. R., M. Lipton, C. Caufield, G. Chang, T. Hendrickson and W. C. Still, J. Comput. Chem. 1990, 11,440-467.
- [28]G.Micera, S.Deiana, C.Gessaand M.Petrera, Jnorg. Chim. Acta, 56, 109 (1981).
- [29]G. Charron, F. Bellot, F. Cisnetti, G. Pelosi, J.-N. Rebilly, E. Rivière, A.-L. Barra, T. Mallahand C. Policar, Chem. Eur. J. 2007, 13,2774-2782.
- [30] G. Camci-Unal and N. L. B. Pohl, Carbohydr. Polym. 2010, 81, 8-13; S. Striegler and M. G. Gichinga, Chem. Commun. 2008,5930-5932.
- [31] G. Sheldrick, SHELXS-97 Program for Crystal Structure Refinement, 1997.
- [32] G. Sheldrick, SHELXL-97 Program for Crystal Structure Refinement, 1997.
- [33] G. L. Strati, J. L. Willett and F. A. Momany, Carbohydr. Res. 2002, 337,1833-1849.
- [34] H. M. Irving and R. J. P. Williams, J. Chem. Soc. 1953,3192-3210.
- [35] H. D. Flack, Acta Cryst. 1983, A39, 876-881.
- [36]I.Zay, A. Vertes, G. Takacsi Nagy, M. Subaand K. Burger, Radioanal. Nuc/. Chem., 8,343(1985)
- [37]I. Berger, M. Hanif, A. A. Nazarov, C. G. Hartinger, R. O. John, M. L. Kuznetsov, M.

Groessl, F. Schmitt, O. Zava, F. Biba, V. B. Arion, M. Galanski, M. A. Jakupec, L. Juillerat-Jeanneret, P. J. Dyson and B. K. Keppler, Chem. Eur. J. 2008, 14, 9046-9057; M. Gottschaldt, U. S. Schubert, S. Rau, S. Yano, J.G. Vos, T. Kroll, J. Clement and I. Hilger, ChemBioChem2010, 11, 649-652; M. Hanif, S. M. Meier, W. Kandioller, A. Bytzek, M. Hejl, C. G. Hartinger, A. A. Nazarov, V. B. Arion, M. A. Jakupec, P. J. Dyson and B. K. Keppler, J. Inorg. Biochem. 2011, 105, 224-231; M. Hanif, A. A. Nazarov, C. G. Hartinger, W. Kandioller, M. A. Jakupec, V. B. Arion, P. J. Dyson and B. K. Keppler, Dalton Trans. **2010**, 39, 7345- 7352; J. K. H. Hui, Z. Yu, T. Mirfakhrai and M. J. MacLachlan, Chem. Eur. J. 2009, 15, 13456-13465; R. Kikkeri, D. Grünstein and P. H. Seeberger, J.2 Am. Chem. Soc. 2010, 132, 10230-10232; M.-W. Louie, H.-W. Liu, M. H.-C. Lam, Y.-W. Lam and K. K.-W. Lo, Chem. Eur. J. 2011, 17, 8304-8308; Y. Mikata, S. Fujii, M. Naemura, K.Takahashi and Y. Noguchi, Dalton Trans. 2009, 10305-10310; T. Shibata, H.Hashimoto, I. Kinoshita, S. Yano and T. Nishioka, Dalton Trans. 2011, 40, 4826-4829; T. Storr, L. E. Scott, M. L. Bowen, D. E. Green, K. H. Thompson, H. J. Schugar and C. Orvig, Dalton Trans. 2009, 3034-3043; C.-C. Yang, P.-S. Lin, F.-C. Liu, I. J. B. Lin, G.-H. Lee and S.-M. Peng, Organometallics 2010, 29, 5959-5971; E. C. Constable, C. E. Housecroft, M. Neuburger and P. Rosel, Chem. Commun. 2010, 46, 1628-1630.

- [38] In this Paper, solvation issues are not discussed, however as we consider two ligands with the same coordinating sites and their interactions with metal cations possessing the same charge, solvation effects could most confidently be neglected while discussing the metal selectivities in this study. ,Vol.
- [39]Ju.A.Komkov,M.A.Ivanov,G.A.ShagisultanovaandJu.B.Jakovlev,ZhNeorg.Khim.,29,15 202(1984).
- [40]J.-H. Liao, C.-S. Chen, C.-C. Hu, W.-T. Chen, S.-P. Wang, I. L. Lin, Y.-H. Huang, M.-H. Tsai, T.-H. Wu, F.-Y. Huang and S.-H. Wu, Inorg. Chem. 2011, 50, 365-377; P. G. Devi, S. Pal, R. Banerjee and D. Dasgupta, J. Inorg. Biochem. 2007, 101,127-137.
- [41]J. Ali-Torres, J.-D. Maréchal, L. Rodriguez-Santiago and M. Sodupe, J. Am. Chem. Soc. 2011, 133, 15008-15014; V. Muñoz Robles, E. Ortega-Carrasco, E. G. Fuentes, A. Lledos and J.-D. Maréchal, Faraday Discuss. 2011, 148,137-159.
- [42] K.Burger, I.Zayand G. Takacsi Nagy, Jnorg. Chim. Acta, 80, 231 (1983).
- [43]K. Born, P. Comba, R. Ferrari, G. A. Lawrance and H. Wadepohl, Inorg. Chem. 2007,

46,458-464.

- [44] L. Garcia, S. Maisonneuve, J. Xie, R. Guillot, P. Dorlet, E. Rivière, M. Desmadril, F.Lambert and C. Policar, Inorg. Chem. 2010, 49,7282-7288.
- [45] L. Garcia, S. Maisonneuve, J. Oudinet-Sin Marcu, R. Guillot, F. Lambert, J. Xie and C. Policar, Inorg. Chem. 2011, 50,11353–11362.
- [46]L. J. Farrugia, J. Appl. Cryst. 1999, 32, 837-838.M.Tonkovic, S.Music, I.Hadzija, J.Nagy-Czak6and A.Vertes, ActaChim.Hung., 110, 197(1982).
- [47]M. M. K. Boysen, Chem. Eur. J. 2007, 13, 8648-8659.
- [48] M.GottschaldtandU.S.Schubert, Chem.Eur.J.2009,15,1548-1557; M.L.BowenandC.Orvig, Chem. Commun. 2008, 5077-5091.
- [49] Chem. Eur. J. 2010, 16, 10202-10213; A. H. Younes, L. Zhang, R. J. Clark and L. Zhu, M. Ostermeier, M.-A. Berlin, R. M. Meudtner, S. Demeshko, F. Meyer, C. Limberg and S. Hecht, J. Org. Chem. 2009,74,8761-8772;L.ZhangandL.Zhu,J.Org.Chem.2008,73,8321-8330; N.K.Dalley,U.Olsher,J. C. Lee, M. D. Eley, J. Wang and R. A. Bartsch, Tetrahedron 2007, 63, 10576-10580; L. Mikulášek, B. Grüner, C. Dordea, V. Rudzevich, V. Böhmer, J. Haddaoui, V. Hubscher-Bruder, F. Arnaud-Neu, J. Čáslavský and P. Selucký, European J. Inorg. Chem. 2007, 2007, 4772-4783; P. Comba, N. Dovalil, G. Haberhauer, G. Hanson, Y. Kato and T. Taura, J. Biol. Inorg. Chem. 2010, 15, 1129-1135.
- [50] M. Miguirditchian, D. Guillaneux, D. Guillaumont, P. Moisy, C. Madic, M. P. Jensen and K. L.Nash, Inorg. Chem. 2005, 44, 1404-1412. M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, J.J. A. Montgomery, T. Vreven, K. N. Kudin, J. C. Burant, J. M. Millam, S. S. Iyengar, J. Tomasi, V. Barone, B. Mennucci, M. Cossi, G. Scalmani, N. Rega, G. A. Petersson, H. Nakatsuji, M. Hada, M. Ehara, K.Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, M. Klene, X. Li, J. E. Knox, H. P. Hratchian, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, P. Y. Ayala, K. Morokuma, G. A. Voth, P. Salvador, J. J. Dannenberg, V. G. Zakrzewski, S. Dapprich, A. D. Daniels, M. C. Strain, O. Farkas, D. K. Malick, A. D. Rabuck, K. Raghavachari, J. B. Foresman, J. V. Ortiz, Q. Cui, A. G. Baboul, S. Clifford, J. Cioslowski, B. B. Stefanov, G. Liu, A. Liashenko, P. Piskorz, I. Komaromi, R. L. Martin, D. J. Fox, T. Keith, M. A. Al-Laham, C. Y. Peng, A. Nanayakkara, M. Challacombe, P. M. W. Gill, B. Johnson, W. Chen,

M. W. Wong, C. Gonzalez and J. A. Pople, Gaussian 03, Gaussian, Inc., Wallingford CT, 2004.

- [51] N. Ghaschghaie, T. Hoffmann, M. Steinborn and P. Klufers, *DaltonTrans.* 2010, 39, 5535-5543; A. L. Petrou, *Coord. Chem. Rev.* 2002, 228,153-162.
- [52]P.Saltman,].Chem.Educ.,42,682(1965).
- [53] P.J.Charley, B. Sarkar, C. F. Stitt and P. Saltman, Bio-chim. Biophys. Acta, 69, 313 (1963).
- [54]P. Comba, Coord. Chem. Rev. 1999, 182, 343-371; P. Comba and W. Schiek, Coord.Chem.
 Rev.2003, 238-239, 21-29.
- [55] P. Comba, T. W. Hambley and B. Martin, Molecular Modeling of inorganic compounds, 3rd edition, Wiley VCH Weinheim, 2010, p.
- [56]P.Comba,Coord.Chem.Rev.1999,185186,898;P.CombaandM.Kerscher,Coord.Chem.Rev.20 09, 253, 564-574.
- [57]P. J. Hay and W. R. Wadt, J. Chem. Phys. 1985, 82, 299-310.
- [58] R. D. Hancock, D. L. Melton, J. M. Harrington, F. C. McDonald, R. T. Gephart, L.L.Boone, S. B. Jones,
- [59] N. E. Dean, J. R. Whitehead and G. M. Cockrell, Coord. Chem. Rev. 2007, 251,1678-1689.
- [60]R. Deeth and C. Diedrich, J. Biol. Inorg. Chem. 2009, 15, 117-129'N.Gresh, G. A. Cisneros,
 T. A. Darden and J. P. Piquemal, J. Chem. Theory Comput. 2007, 3, 1960-1986.
- [61] R. J. Deeth, A. Anastasi, C. Diedrich and K. Randell, Coord. Chem. Rev. 2009, 253,795-816.
- [62] R. D. Hancock and K. Hegetschweiler, J. Chem. Soc., Dalton Trans. 1993,2137-2140.
- [63] R. Jeanloz, H. G. Fletcher and C. S. Hudson, J. Am. Chem. Soc. 1948, 70,4052-4054.
- [64] R. D. Shannon, Acta Cryst. 1976, A32, 751-767.
- [65] S.WolowiecandK. Drabent, J.Radioanal.Nucl. Chem. Lett., 95, 1 (1985).
- [66] S.Dhungana, J.M.Harrington, P.Gebhardt, U.Möllmannand A.L.Crumbliss, Inorg. Chem. 2007, 46, 8362-8371.
- [67] S. Woodward, M. Diéguez and O. Pàmies, Coord. Chem. Rev. 2010, 254,2007-2030.
- [68] S. J. Angyal, Tetrahedron 1974, 30, 1695-1702; S. J. Angyal, Chem. Soc. Rev. 1980, 9, 415-428; S. J.Angyal, Aust. J. Chem. 2000, 53, 567-570; B. Gyurcsick and L. Nagy, Coord. Chem. Rev. 2000, 203, 81- 149; N. Nagy and A. Szorcsik, J. Inorg. Biochem. 2002, 89, 1-

12; J.-F. Verchèreand S. Chapelle, Progr. Inorg. Chem. **1998**, 47, 837-945; D. M. Whitfield, S. Stojkovski and B. Sarkar, Coord. Chem. Rev. **1993**, 122, 171-225.

[69] S. J. Angyal, Adv. Carbohydr. Chem. Biochem. 1989, 47,1-43.

- [70] S. Bjelić and I. Jelesarov, J. Mol. Recognit. 2009, 21, 289-312; R. J. Falconer and B. M. Collins, J. Mol. Recognit. 2011, 24, 1-16; R. J. Falconer, A. Penkova, I. Jelesarov and B. M. Collins, J. Mol. Recognit. 2010, 23, 395-413; O. Okhrimenko and I. Jelesarov, J. Mol. Recognit. 2008, 21,1-19.
- [71] Schrodinger, Inc., Portland, Oregon, 2000,
- [72]T.K.Lindhorst, Essentials of Carbohydrate Chemistry and Biochemistry, Wiley-VCH, 2003, p
- [73] T.Marino, N.Russo and M.Toscano, J.Inorg. Biochem. 2000, 79, 179-185; J. Bertran,L.Rodriguez-Santiago and M. Sodupe, J. Phys. Chem. B 1999, 103,2310-2317.
- [74] The use of unprotected sugar polyols or polyolate as ligands, while not discussed here indetail, is conceptually related to urwork. For a review see: T.Allscher, P.Klüfers,
- [75]P.Mayerin:B.Fraser-Reid,K. Tatsuta, J. Thiem (eds.) in Glycoscience, Springer-Verlag, Berlin, 2008, 1079–1139., Vol.
- [76] V. J. James, J. D. Stevens and F. H. Moore, Acta Crist. 1978, B34,188-193.
- [77] W. O. Gillum, R. A. D. Wentworth and R. F. Childers, Inorg. Chem. 1970, 9,1825-1832.
- [78] W. Jorgensen, D. Maxwell and J. Tirado-Rives, J. Am. Chem. Soc. 1996, 118, 11225-11236.
- [79] W.J.Hehre, R.Ditchfieldand J.A.Pople, J.Chem. Phys. 1972, 56, 2257-2261; P.C.Hariharanand
 J. A. Pople, Theor. Chim. Acta 1973, 28, 213-222.
- [80]Z.Jingkun,L.Rongehuan,W.Shuxin,X.YuanfuandQ.Shouren,KexeuTongbao,1091(1984).

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Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

General Parameters

Method : Iron (Fe) Autosampler : None Use SFI: No Operator : Jyothi

Instrument Mode: Flame Dilution: None

mA Hours

n/a

Report Date: 11-06-2022 17:40:20

Valid Method Signatures

11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Method Audit Trail

11-06-2022 17:22:18 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC

Signed with Reason : Approved by:

Analysis Details

Spectrometer: iCE 3000 AA01204906 v1.30

Analysis Name: Iron (Fe) 11-06-2022 Operator Name: Jyothi

n/a

Lamp Information Serial Number

Element(s) Fe

. . .

Deuterium Lamp Hours: 68.34

Sequence Table

Shared Standards: Yes	
Action	Fe
Calibration	√
B.No.Group-II	√
B.No.Group-III	√
B.No.Group-IV	√
B.No.Group-V	√
B.No.Group-VI	✓

Sample Details

		Nominal Mass: 1.000	00
No.	Sample Id	Sample Mass	Dilution Ratio
1	B.No.Group-II	1.0000	1.0000
2	B.No.Group-III	1.0000	1.0000
3	B.No.Group-IV	1.0000	1.0000
4	B.No.Group-V	1.0000	1.0000
5	B.No.Group-VI	1.0000	1.0000

Valid Analysis Signatures

11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Analysis Audit Trail

- 11-06-2022 17:30:16 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
- 11-06-2022 17:38:27 Jyothi(M. Jyothi):DESKTOP-39TDEGC Error MD147 - Activity manually aborted by user.
- 11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
- 11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

MART SPECIALITIES LAB LLP.

Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Spectrometer Parameters - Fe

Element: Fe Wavelength: 248.3nm Background Correction: D2 Signal Type: Continuous Measurement Time: 4.0secs Use RSD Test: No

Flame Type: Air-C2H2 Nebuliser Uptake: 4secs Burner Height: 7.0mm

Sampling: None

Calibration Mode: Normal	
Concentration Units: mg/L	
Acceptable Fit: 0.990	
Standard 1	2.0000
Standard 2	4.0000
Standard 3	6.0000

No changes are recorded for this element

Measurement Mode: Absorbance Bandpass: 0.2nm High Resolution: Off Resamples: Fast Flier Mode: No

Lamp Current: 75% Optimise Spectrometer Parameters: No Number Of Resamples: 3

Flame Parameters - Fe

Fuel Flow: 0.9L/min Burner Stabilisation: 0mins Optimise Burner Height: No Auxilary Oxidant: Off Optimise Fuel Flow: No

Sampling Parameters - Fe

Calibration Parameters - Fe

Line Fit: Linear Scaled Units: mg/L Rescale Limit: 10.0% Standard 4 Standard 5 Use Stored Calibration: No Scaling Factor: 1.0000 Failure Action: Flag and Continue 8.0000 10.0000

Element Audit Trail - Fe

Solution Results - Fe

Y = 0.04924x + 0.0141 Fit: 0.9965 Characteristic Conc: 0.0894

> 1 2 3

> 1 2 3

> 1 2 3

> 1 2 3

1 2

3

Sample ID

Fe Blank

Fe Standard 1

Fe Standard 2

Fe Standard 3

Fe Standard 4

	4				Norma	I: Linea	ar Fit				
	Abs		*		*		*		-*-		*
	0 * -0-0	1	2	3	4	5	6	7	8	9	10
	-0-0		2	0		nc:mg/		,	0	0	10
Signal	Rsd	Conc				cted C					
Abs	%	mg/L			mg/L						
0.001	35.6	0.000)		Ũ						
0.001	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:26	i		
0.001	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:30)		
0.000	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:35			
0.120	0.3	2.0000)								
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:05			
0.119	-	ound: 0.00						7:32:09			
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:13			
0.215	0.3	4.0000)								
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:40)		
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:45			
0.216	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:49)		
0.322	0.2	6.0000)								
0.323	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:19)		
0.322	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:23			
0.322	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:27			
0.411	0.3	8.0000)								
0.410	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:59)		
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:03			
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:07	•		

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Operator Name: Jyothi

Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Solution Results - Fe

Sample ID	Signal	Rsd	Conc	Corrected Conc
-	Abs	%	mg/L	mg/L
Fe Standard 5	0.493	0.1	10.0000	
1	0.494	Background: 0.005		11-06-2022 17:34:38
2	0.494	Backg	ground: 0.005	11-06-2022 17:34:42
3	0.493	Backg	ground: 0.006	11-06-2022 17:34:46
Fe B.No.Group-II	0.834	0.1	16.6490 C	16.6490 C
1	0.834	Backg	ground: 0.005	11-06-2022 17:35:14
2	0.833	Backg	ground: 0.005	11-06-2022 17:35:19
3	0.834	Backg	ground: 0.005	11-06-2022 17:35:23
Fe B.No.Group-III	0.759	0.1	15.1286 C	15.1286 C
1	0.758	Backg	ground: 0.003	11-06-2022 17:35:52
2	0.760	Backg	ground: 0.003	11-06-2022 17:35:56
3	0.758	Backg	ground: 0.004	11-06-2022 17:36:00
Fe B.No.Group-IV	0.645	0.3	12.8189 C	12.8189 C
1	0.647	Backg	ground: 0.003	11-06-2022 17:36:31
2	0.644	Background: 0.003		11-06-2022 17:36:35
3	0.645	Backg	ground: 0.003	11-06-2022 17:36:40
Fe B.No.Group-V	0.809	0.2	16.1488 C	16.1488 C
1	0.808	Backg	ground: 0.005	11-06-2022 17:37:11
2	0.809	Backg	ground: 0.005	11-06-2022 17:37:15
3	0.811	Backg	ground: 0.005	11-06-2022 17:37:20
Fe B.No.Group-VI	0.711	0.3	14.1597 C	14.1597 C
1	0.709	Backg	ground: 0.005	11-06-2022 17:37:55
2	0.713	Backg	ground: 0.005	11-06-2022 17:37:59
3	0.712	Backg	ground: 0.004	11-06-2022 17:38:03

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-1 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
I	DETAILS OF THE SAMPLE
Sample Name : NA	
Name of the Manufacturer : NA	
Storage condition To b	up-I Mfg. : NA Exp. : NA e stored at room date date
	Derature Batch : NA C±3°C size
	by AAS Analysis.
	6/2022
Analysis Completion Date : 11/0	6/2022
Mfg. License No. : Not	provided
A.R.NO : NA	The Andrew Rest

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	620ppm

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



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	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00		
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-2 Sample Received Date : 11/06/2022 Report Date : 13/06/2022		
Q	DETAILS OF THE SAMPLE		
Sample Name : N	A		
Name of the Manufacturer : N	A		
Storage condition : To	roup-II Mfg. : NA Exp. : NA be stored at room date date		
Room Temperature : 25	Temperature 25°C±3°C 4gm		
Tests Required:IreMethod:NAnalysis Starting Date:11	on by AAS Analysis.		
	ot provided		
A.R.NO : N	A		
	Sample analyzed as received		

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	832.5ppm

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



OCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinaga Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Sample Received Date : 11/06/2022
a.y.	DETAILS OF THE SAMPLE
Sample Name : N	JA
Name of the Manufacturer : N	JA
Storage condition	Group-III Mfg. : NA Exp. : NA To be stored at room date date Cemperature Batch : NA
	5°C±3°C size
	gm /
Method : I Analysis Startin <mark>g Date : I</mark>	ron by AAS Analysis. NA 1/06/2022 1/06/2022
and the second se	Not provided
A.R.NO	NA
Remark : S	Sample analyzed as received

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	756.5ppm

×

Authorized Signatory

(Dr.R.Marayya)

Office: 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile: 98481 98350, 98481 25019 Mail: rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College	
Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000	Sample Received Date: 11/06/2022Report Date: 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name : N	A
Name of the Manufacturer : N	A
Storage condition : T	roup-IV Mfg. : NA Exp. : NA o be stored at room date date emperature Batch : NA
	5°C±3°C size
Method : N Analysis Starting Date : 1	on by AAS Analysis. A 1/06/2022 1/06/2022
Mfg. License <mark>No. :</mark> N	ot provided
A.R.NO : N	A
Remark : Sa	ample analyzed as received

Test Results

Harris I.		pristance of	
S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	641ppm	

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/	G
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Custome	r: Reference / Report No. : MSL/2022/JUNE/TARA/0064-5
Tara Government College	Sample Received Date : 11/06/2022
Prashanth Nagar Colony, Balajinag	ar Sumpto Received 2 mil
Sangareddy Telangana. 502000	Report Date : 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name :	NA
Name of the Manufacturer :	NA
Batch no : Storage condition	Group-VMfg. : NAExp. : NATo be stored at roomdatedateTemperatureBatch : NA
Room Temperature :	25°C±3°C size
Quantity Received	4gm
Tests Required:Method:Analysis Starting Date:	Iron by AAS Analysis. NA 11/06/2022
Analysis Completion Date :	11/06/2022
Mfg. License No.	Not provided NA
A.R.NO Remark	Sample analyzed as received

Test Results

S.No.	Test Parameter	Result
01. Ir	on by AAS Analysis: (ppm)	807.5ppm

>

Authorized Signatory

(Dr.R.Marayya)

Office : 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile : 98481 98350, 98481 25019 Mail : rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-6 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
1	DETAILS OF THE SAMPLE
Storage condition	up-V1 Mfg. : NA Exp. : NA be stored at room date date perature Batch : NA C±3°C size
Tests Required:IronMethod:NAAnalysis Starting Date:11/0Analysis Completion Date:11/0Mfg. License No.:NotA.R.NO:NA	by AAS Analysis. 6/2022 6/2022 provided

Test Results

S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	708ppm	

*

Authorized Signatory

(Dr.R.Marayya)

ADSORPTION STUDIES OF Fe[III] ON ACTIVATED CHARCOAL IN THE PRESENCE OF LACTIC ACID AS CHELATING AGENT

Dissertation submitted in Partial fulfillment for the requirements for the award of degree of

Bachelor of Science in CHEMISTRY

By

1. MukthiKanth Rout,	B.Sc(Chemistry),	III	Year
2. S.Raja Ramesh,	B.Sc(Chemistry),	III	Year
3. P.Sujil Kumar,	B.Sc(Chemistry),	III	Year
4. Ch.srinivas,	B.Sc(Chemistry),	III	Year
5. B.Mahesh Kumar,	B.Sc(Chemistry),	III	Year
6. B.NikhileshBabu,	B.Sc(Chemistry),	III Y	Year

Under the Guídance:

Dr. K.ABHIJIT *HEAD* DEPARTMENT OF CHEMISTRY



TARA GOVERNMENT COLLEGE, SANGAREDDY (A) 2021-22

<u>CERTIFICATE</u>

This is to certify that the project work entitled "ADSORPTION STUDIES OF FE[III] ON ACTIVATED CHARCOAL IN THE PRESENCE OF LACTIC ACID AS CHELATING AGENT " is presented by B.Sc (CHEMISTRY) students in partial fulfillment of the requirements for the degree of Bachelor of Science in Chemistry by the Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

The results embodied in this report have not been to any other University or Institution for the award of any degree.

(Dr. K. ABHIJIT)

Project Supervisor& Head, Department of Chemistry Tara Government College, Sangareddy(A)

EXTERNAL EVALUATOR

ACKNOWLEDGEMENTS

We express my deep gratitude to my research supervisor **Dr.K.Abhíjít**, Head, Department of Chemistry, Tara Govt. College, Sangareddy(A)-502001, INDIA for his inspiring guidance during the course of the Project work. The continuous encouragement extended by him propelled me to update my research skills and pedigree to engage in fruitful research.

We wish to express my gratitude to Smt. M.Praveena, Principal, Tara Govt. College, Sangareddy(A)-502001, INDIA for her constant support, cooperation and suggestions during the research work. We wish to express my sincere thanks to K.Sreedhar, Asst. Professor, and Department of Chemistry for providing me facilities, help and support for the entire research work.

We express our special thanks to **MART Specialities Lab**, **Hyderabad** for providing technical assistance in Atomic absorption *spectrometric analysis*.

Finally, we take this opportunity to thank one and all that has directly or indirectly helped me in completing the task.

1. MukthiKanth Rout,	B.Sc(Chemistry),	III Year	
2. S.Raja Ramesh,	B.Sc(Chemistry),	III Year	
3. P.Sujil Kumar,	B.Sc(Chemistry),	III Year	
4. Ch.srinivas,	B.Sc(Chemistry),	III Year	
5. B.Mahesh Kumar,	B.Sc(Chemistry),	III Year	
6. B.NikhileshBabu,	B.Sc(Chemistry),	III Year	

<u>DECLARATION</u>

We hereby declare that the project report entitled "ADSORPTION STUDIES OF Fe[III] ON ACTIVATED CHARCOAL IN THE PRESENCE OF LACTIC ACID AS CHELATING AGENT" is the work done by us in the campus at **Department of Chemistry,** Tara Government College, Sangareddy(A) during the academic year 2021-2022 and is submitted in partial fulfillment of the requirements for the degree of **Bachelor of Science** by Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

S.No.	Name of the Student	Roll Number	Group	Year	Signature
1	MukthiKanth Rout	6058-19-578-022	B.sc - MCCS	III	
2	S. Raja Ramesh	6058-19-578-030	B.sc - MCCS	III	
3	P. Sujil Kumar	6058-19-578-027	B.sc - MCCS	III	
4	Ch. Srinivas	6058-19-578-008	B.sc - MCCS	III	
5	B. Mahesh Kumar	6058-19-578-004	B.sc - MCCS	III	
6	B. NikhileshBabu	6058-19-578-005	B.sc - MCCS	III	

TARA GOVERNMENT COLLEGE, SANGAREDDY – 502 001 (AUTONOMOUS)

BONAFIDE CERTIFICATE

Certified that the project report "ADSORPTION STUDIES OF Fe[III] ON

ACTIVATED CHARCOAL IN THE PRESENCE OF LACTIC ACID AS CHELATING AGENT \H is

the bonafide work of

1. MukthiKanth Rout,	B.Sc(Chemistry),	III Year
2. S.Raja Ramesh,	B.Sc(Chemistry),	III Year
3. P.Sujil Kumar,	B.Sc(Chemistry),	III Year
4. Ch.srinivas,	B.Sc(Chemistry),	III Year
5. B.Mahesh Kumar,	B.Sc(Chemistry),	III Year
6. B.NikhileshBabu,	B.Sc(Chemistry),	III Year

who carried out the project work under my supervision.

Dr.K.ABHIJIT **PROJECT SUPERVISOR** Smt. M.PRAVEENA PRINCIPAL

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Chapter-I INTRODUCTION

Adsorption

Adsorption operation involves the touch of solids with both liquids and gases in which the mass switch is towards solids. The reverse of this operation is known as "desorption." Adsorption operations exploit the potential of positive solids to concentrate precise materials from fluid on to their surfaces. The adsorbed substance is known as an adsorbate, and the solid substance is known as an adsorbent. Typical packages of this stable-liquid operation are as follows:

- Gas: Elimination of moisture dissolved in gas
- de-colorization of petroleum merchandize and sugar answers
- water for the removal of objectionable flavours and odours.

The stable-gasoline operations consist of:

- Dehumidification of air and gases
- Gas purification is the process of removing noxious odours and impurities from gases.
- The recovery of precious solvent vapours from dilute gasoline combinations
- fractionate combinations of hydrocarbon gases, inclusive of methane, ethane, and propane.

NATURE OF ADSORBENTS

Adsorbents are commonly in granular form, with their size starting from zero to five to twelve millimetres. They must neither provide a high strain drop nor get overly excited by the flowing stream. They should now not lose their shape and size at the same time as managing. They must have a large surface area per unit mass and a lot of pores.

S.NO	Adsorbent	Source	Application
1.	Fuller's earth	Naturally occurring clay is	De-colorizing,
		heated and dried to get a	drying of
		porous structure.	lubricating oils,
			kerosene and
			engine oils.
2.	Activated clay	Bentonite or other activated	Used for de-
		clay which are activated by	colorizing
		treatment with sulfuric acid	petroleum
		and further washing, drying	products.
		and crushing	
3.	Bauxite	A naturally occurring	Used for de-
		hydrated alumina, activated	colorizing
		by heating at 230-815 ⁰ C	petroleum
			products and for

			drying gases.
4.	Alumina	Hard hydrated aluminum	Used as a
		oxide is activated by heating	desiccant.
		to drive off the moisture and	
		then crushed to the desired	
		size.	
5.	Bone – Char	Obtained by destructive	Used for refining
5.	Bone – Chai		_
		distillation of crushed bones at	sugar and can be
		600-9000C.	reused after
			washing and
			burning.
6.	Activated Carbon	(i) Vegetable matter	De-colorizing of
		is mixed with	sugar solutions,
		calcium chloride,	chemicals, drugs,
		carbonized and	water purification,
		finally, the	refining of
		inorganic	vegetable and
		compounds are	animal oils,
		leached away.	recovery of gold
		(ii) Organic matter is	and silver from
		mixed with	cyanide ore-leach

					1	
			porous p	umice	solution,	recovery
			stones and	then	of solvent	vapour
			heated	and	from	gas-
			carbonized	to	mixtures,	
			deposit	the	collection	of
			carbonaceous		gasoline	hydro-
			matter		carbons	from
			throughout	the	natural	gas,
			porous parti	cle.	fractionatio	on of
		(iii)	Carbonizing	5	hydrocarbon	
			substances	like	gases.	
			wood, sav	vdust,		
			coconut s	shells,		
			fruits pits, li	gnite,		
				quent		
			activation	with		
			the hot air s	steam.		
			it is availat	ole in		
			granular	or		
			pelleted forr			
			peneceu ion			
7.	Silice col	A hard or	anular and no	r 0116	Used fo	or de-
/.	Silica gel		anular and po			
		product		from	hydration	
		sodium silicate solution after			and other	
		treatment	with a	acid.	fractionation	on of

		Normally has 4 to 7% water	hydrocarbons.
		in the product.	
8.	Molecular sieves	These are porous synthetic	Dehydration of
		zeolite crystals, metal	gases and liquids,
		aluminosilicates.	and separation of
			gas— liquid
			hydrocarbon
			mixture.

<u>Heat of adsorption</u>

The differential heat of adsorption (—H) is described as the heat liberated at a regular temperature whilst a unit quantity of vapour is adsorbed on a large quantity of stable already containing adsorbate. The solid so used is in such a huge quantity that the adsorbate concentration stays unchanged.

The indispensable warmness of adsorption (AH) at any concentration X is described as the enthalpy of the adsorbate—adsorbent mixture minus the sum of the enthalpies of unit weight of natural stable adsorbent and sufficient pure adsorbed substance (before adsorption) to offer the specified concentration X at an equal temperature.

The differential warmness of adsorption and the necessary heat of adsorption are capabilities of temperature and adsorbate concentration.

<u>Effect of temperature</u>

An increase in temperature at steady stress decreases the quantity of solute adsorbed from a combination. However, the generalisation of the end result isn't smooth.

Effect of pressure

Generally, reducing stress reduces the quantity of adsorbate adsorbed upon the adsorbent. However, the relative adsorption of paraffin hydrocarbons on carbon decreases at high pressures.

<u>Liquids</u>

The pollutants are available both at low and high fixations in fluids. These are typically eliminated by an adsorption procedure. The attributes of adsorption of low and high fixation debasements are unique.

Adsorption of solute from Dilute solutions

Whenever a combination of solute and dissolvable is adsorbed, utilising an adsorbent, both the dissolvable and the solute are adsorbed. Because of this, the main family member or evident adsorption of solute cannot be entirely set in stone.

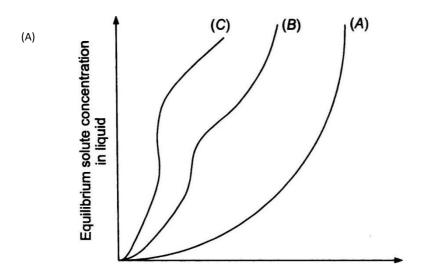
Thus, it is a typical practice to treat a known volume of an arrangement of unique focus (Co) with a known load of adsorbent. Allow C* to be the last harmony grouping of solfege in the arrangement.

Assuming that v is the volume of arrangement per unit mass of adsorbent (cc/g) and Co and C* are the underlying and balance focuses (g/cc) of the solute, then the evident adsorption of solute per unit mass of adsorbent, dismissing any adjustment of volume, is v (Co-C*), (g/g).

This articulation is, for the most part, pertinent to weakening arrangements. At the point when the negligible part of the first dissolvable which can be adsorbed is small, the C^* esteem relies upon the temperature, nature, and properties of the adsorbent.

On account of weakening arrangements and over a small focus range, the Freundlich adsorption isotherm portrays the adsorption peculiarities.

The Freundlich adsorption condition is additionally very helpful in situations where the real character of the solute isn't known, for example, the expulsion of shading substances from sugar arrangements, oils, and so on. The arrangement's variety content can be easily estimated by utilising a spectrophotometer or colorimeter. The translation of this information is shown in working model 2. Assuming the value of n is high, i.e., 2 to 10, adsorption is great. Assuming it lies between I and 2, tolerably troublesome and short of what I demonstrate, I demonstrate unfortunate adsorption attributes. A normal adsorption isothermal for the adsorption of different adsorbates A, B, and C in a weakened arrangement at a similar temperature for a similar adsorbent is displayed in Fig.



Kg solute apparently adsorbed / kg adsorbent

Fig. Adsorption isotherms for various adsorbents.

Significance of Research problem

Adsorption of heavy metals is a key development approach for more advanced remediation technologies for the preservation of the environment. However, the effectiveness of heavy metal adsorption under typical conditions using a good adsorbent depends on a number of variables that must be precisely adjusted to get an effective adsorption process. The adsorption of heavy metals was boosted in the presence of suitable facilitating agents, which undoubtedly improved the heavy metal procedures already in use.

Objective of the Project

To evaluate the impact of Lactic acid as chelating agent in the adsorption of Fe (III) from aqueous solution by activated charcoal as an adsorbent to develop efficient remediation technology using Metal-ligand interactions.

Chapter-II

ADSORPTION OF HEAVY METALS USING ACTIVATED CHARCOAL

Nowadays, various heavy metals have been released into the waters, causing serious pollution of water resources and endangering human health. Therefore, it is very important to study the removal of heavy metals from water. In this work, the adsorption of heavy metal ions including Pb(II), Cu(II), Zn(II), Cd(II) and Cr(VI) onto activated carbon (AC) from aqueous solutions was investigated in both single and mixed systems. The batch adsorption experiment for heavy metal ions on AC in single system was studied at different conditions including contact time, initial concentration, initial pH and adsorbent dosage. The results indicated that the adsorption kinetics and isotherms followed the pseudo-first order model and the Langmuir isotherm model, respectively. The final removal efficiencies and maximum adsorption capacity followed the order of Pb(II)> Cr(VI) Cd(II) > Cu(II) > Zn(II) at the pH 5.0 and 25°C in single system. The adsorption of heavy metals was mainly determined by the pH of the solution and the surface properties of the adsorbent, and it was found that the initial pH of 3 was optimal for the removal of Cr(VI), which was different from the optimum initial pH of 7.0 for Pb(II), Cu(II), Zn(II) and Cd(II). The results demonstrated that the electrostatic interaction between the surface of AC and heavy metal ions played an important role in the adsorption of heavy metal ions. In addition, compared with the results in single system, the results in multi-component mixed system showed that the adsorption of Cu(II) and Cr(VI) was promoted, but the adsorption of Zn(II) and Cd(II) was inhibited. The factors affecting the adsorption impetuses are interactive involving electrostatic interaction, promotion or inhibition between heavy metal ions, chemical reaction and so on.

Adsorption of Iron complex using Activated Charcoal

Among pollutants, iron have attracted our attention because; they are the most released by industries. To curb this kind of pollution, a local alternative must be developed by these countries to ensure the health of their inhabitants.

There are several methods of treatment geared towards removing these pollutants from water, such as: filtration, ozonation, clarification, photocatalysis, adsorption, membrane processes, electro coagulation, and chemical processes. Although, the above-mentioned methods are efficient in treating high concentration of heavy metal ions, nevertheless these techniques also have disadvantages including incomplete metal removal, high consumption of reagent and very high cost incurred in the process. For the lower concentration of heavy metal ions, adsorption is a much preferable technique and activated carbon has been widely applied for treating industrial wastewater.

Activated carbons as adsorbents from plant origin have been developed from agricultural by-products, called biosorbent. Crude olive stone was used for adsorption of iron present in industrial wastewater. Adsorption on activated carbon remains one of the mainly used methods for these countries.

The aim of this work is to investigate the removal of iron from synthetic solution and industrial wastewater with commercial activated carbon from local wood. The adsorption of metal showed two important parameters: initial concentration and pH of solution. These parameters allows for maximizing the amount of adsorbed metals. However, the problems associated with these adsorbents are the regeneration and recovery of the useful materials, which makes them unattractive for wider commercial applications. The carboxyl group as a part of many compounds (pharmaceuticals, supplements) is a "universal" O-donor ligand which can form complexes of different coordination, and can act as a monodentate or a bidentate ligand.

 α -Hydroxycarboxylic acids are constitutive components of many biochemical processes in the living world. They are often used in pharmaceutical preparations. Lactic acid, 2hydroxypropanoic acid, (LA) is formed by fermentation of sugars and other carbohydrates under the action of anaerobic bacteria and it can be ingested with food into the body. In medicine it is used as a component of the Ringer's and Hartman's solutions. Lactic acid is a suitable molecule for model systems for the study of interaction of biometals with O-donor ligands in biosystems. With M(II) metal ions from solutions of molar concentrations, depending on the conditions of complexation, properties and synergistic effect of the metal ions, LA can form different types of complex associates With M(II) dmetal ions LA can form complexes of the type [M(LA)₂R], where it behaves as a bidentate ligand, and R is molecule of solvent . Cu(II) ion can form mononuclear and binuclear complexes with aliphatic a-hydroxycarboxylic acids at pH values.

Previous studies have shown that there is a mobility of metal ions from biological hard mineral tissue of teeth that had been exposed to the impact of various media (lactic acid, acetic acid, etc.) as a result of the interaction of metal ions with the test media. In the literature there are no enough data about interactions of d-metals and LA under approximately physiological conditions, and at the level of micro-molar concentration at which they are present in human body.

Iron - Lactic acid complex in aqueous solutions

The photochemistry of a I: I Fe [lll]—lactic acid complex Fe(Lact)]⁺, in aqueous solutions was studied by stationary photolysis, nanosecond laser flash photolysis (355 nm, 6 ns), and

femtosecond pump-probe spectroscopy (400 nm, 200 fs). The quantum yield of photolysis of Fe(Lact)]⁺, upon excitation at 355 nm is 0.4 and 0.22 in the deoxygenated and airsaturated solutions, respectively. Weak transient absorption in the range of 500-750 nm was observed in the experiments. It nanosecond was assigned to а Fe^{II}...⁻O-CH(Me)-COO[•]]⁺ radical complex. The spectral properties of the ligandto-metal charge transfer excited state and the characteristic time of formation of the radical complex (1.5 ps) were determined in the femtosecond spectroscopy experiments. A reaction mechanism was proposed, which involves innersphere electron transfer in the excited complex with the formation of a radical complex Fe^{II}...⁻O-CH(Me)-COO[•]]⁺ and its subsequent transformation to the end product of the photochemical reaction.

The photochemistry of Fe [111] carboxylate complexes with natural carboxylic acids (tartaric, pyruvic, lactic, succinic, as well as humic and fulvic acids) is of considerable interest from the standpoint of environmental photochemistry and photocatalysis because photolysis of these compounds is accompanied by the formation of reactive H_2O_2)¹⁻⁵ oxygen species (•OH, H02•), which leads to mineralization of organic matter and to CO and CO2 generation in natural water.

A traditional mechanism of photolysis of Fe [III] carboxylate complexes implies that the primary photochemical event involves inner-sphere electron transfer accompanied by reduction of Fe [III] to Fe [II] and escape of a free radical into the solvent bulk, followed by fast decarboxylation of the radical. A secondary radical generated as a result of decarboxylation reacts with Fe [III] complexes and dissolved oxygen to give reactive oxygen species.

This mechanism is based on the analysis of photochemical reaction end products, and the generation of organic radicals in the primary photochemical process was frequently not confirmed experimentally. Simultaneously, studies on the photochemistry of iron(ll) complexes with oxalic acid 13 and ethylene diamine tetra acetic acid 12'14 using pulsed methods enabled the formulation of an alternative photolysis mechanism based on the production of a long-lived radical complex [Fe^{ll}s • OOC—R] ²+ in the primary photochemical event.

In this work, we studied primary photochemical processes, taking a Fe^{lli}complex with lactic acid as an example. The emphasis is placed on the detection of corresponding radical complex and its precursor, i.e., the excited state, and on the determination of their spectral and kinetic properties.

Chapter-III

ATOMIC ABSORPTION SPECTROMETRY

Atomic absorption spectrometry (AAS) is an analytical technique that measures the concentrations of elements. Atomic absorption is so sensitive that it can measure down to parts per billion of a gram (μ g dm⁻³) in a sample. The technique makes use of the wavelengths of light specifically absorbed by an element. They correspond to the energies needed to promote electrons from one energy level to another, higher, energy level.

Atomic absorption spectrometry has many uses in different areas of chemistry.

Clinical analysis, analysing metals in biological fluids such as blood and urine. **Environmental analysis** monitoring our environment -eg finding out the levels of various elements in rivers, seawater, drinking water, air, petrol and drinks such as wine, beer and fruit drinks.

Pharmaceuticals. In some pharmaceutical manufacturing processes, minute quantities of a catalyst used in the process (usually a metal) are sometimes present in the final product. By using AAS the amount of catalyst present can be determined.

Industry. Many raw materials are examined and AAS is widely used to check that the major elements are present and that toxic impurities are lower than specified -eg in concrete, where calcium is a major constituent, the lead level should be low because it is toxic.

Mining. By using AAS the amount of metals such as gold in rocks can be determined to see whether it is worth mining the rocks to extract the gold.

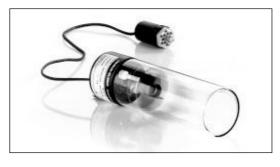
How it works

Atoms of different elements absorb characteristic wavelengths of light. Analysing a sample to see if it contains a particular element means using light from that element. For example with lead, a lamp containing lead emits light from excited lead atoms that produce the right mix of wavelengths to be absorbed by any lead atoms from the sample. In AAS, the sample is atomised *-ie* converted into ground state free atoms in the vapour state – and a beam of electromagnetic radiation emitted from excited lead atoms is passed through the vaporised sample. Some of the radiation is absorbed by the lead atoms in the sample. The greater the number of atoms there is in the vapour, the more radiation is absorbed. The amount of light absorbed is proportional to the number of lead atoms. A calibration curve is constructed by running several samples of known lead concentration under the same conditions as the unknown. The amount the standard absorbs is compared with the calibration curve and this enables the calculation of the lead concentration in the unknown sample.

Consequently an atomic absorption spectrometer needs the following three components: a light source; a sample cell to produce gaseous atoms; and a means of measuring the specific light absorbed.

The light source

The common source of light is a 'hollow cathode lamp' (Fig. 1).





This contains a tungsten anode and a cylindrical hollow cathode made of the element to be determined. These are sealed in a glass tube filled with an inert gas -eg neon or argon – at a pressure of between 1 Nm⁻² and 5 Nm⁻². The ionisation of some gas atoms occurs by applying a potential difference of about 300–400 V between the anode and the cathode. These gaseous ions bombard the cathode and eject metal atoms from the cathode in a process

called sputtering. Some sputtered atoms are in excited states and emit radiation characteristic of the metal as they fall back to the ground state $-egPb^* \rightarrow Pb + h \vee (Fig. 2)$.

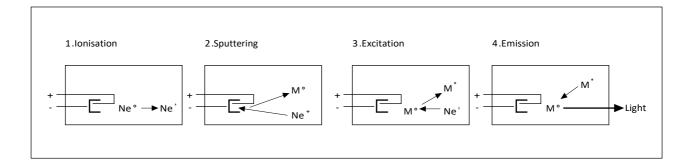
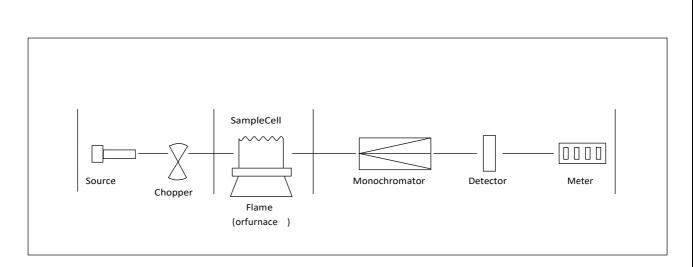


Figure. 2

The shape of the cathode concentrates the radiation into a beam which passes through a quartz window, and the shape of the lamp is such that most of the sputtered atoms are redeposited on the cathode.

A typical atomic absorption instrument holds several lamps each for a different element. The lamps are housed in a rotating turret so that the correct lamp can be quickly selected.

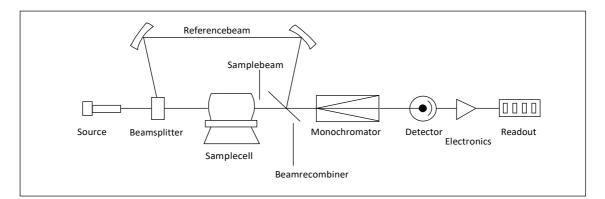
The optical system and detectorA mono chromator is used to select the specific wavelength of light – ie spectral line – which is absorbed by the sample, and to exclude other wavelengths. The selection of the specific light allows the determination of the selected element in the presence of others. The light selected by the monochromator is directed onto a detector that is typically a photomultiplier tube. This produces an electrical signal proportional to the light intensity (Fig. 3)





Double beam spectrometers

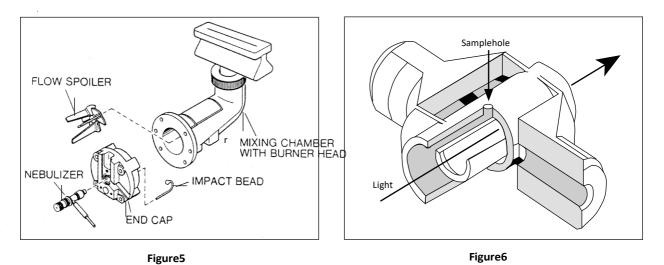
Modern spectrometers incorporate a beam splitter so that one part of the beam passes through the sample cell and the other is the reference (*Fig. 4*). The intensity of the light source may not stay constant during an analysis. If only a single beam is used to pass through the atom cell, a blank reading containing no analyte (substance to be analysed) would have to be taken first, setting the absorbance at zero. If the intensity of the source changes by the time the sample is put in place, the measurement will be inaccurate. In the double beam instrument there is a constant monitoring between the reference beam and the light source. To ensure that the spectrum does not suffer from loss of sensitivity, the beam splitter is designed so that as high a proportion as possible of the energy of the lamp beam passes through the sample.





Atomisation of the sample Two systems are commonly used to produce atoms from the sample. Aspiration involves sucking a solution of the sample into a flame; and electrothermal atomisation is where a drop of sample is placed into a graphite tube that is then heated electrically.

Some instruments have both atomisation systems but share one set of lamps. Once the



appropriate lamp has been selected, it is pointed towards one or other atomisation system.

Flame aspiration

Figure 5 shows a typical burner and spray chamber. Ethyne/air (giving a flame with a temperature of 2200–2400°C) or ethyne/dinitrogen oxide (2600– 2800°C) is often used. A flexible capillary tube connects the solution to the nebuliser. At the tip of the capillary, the solution is 'nebulised' -i.e. broken into small drops. The larger drops fall out and drain off while smaller ones vaporise in the flame. Only *ca* 1% of the sample is nebulised.

Electro thermal atomisation of the original sample *Figure 7* shows a flame atomic *Figure 6* shows a hollow graphite tube with a platform. Absorption spectrometer with an auto sampler and 25 μ l of sample (*ca* 1/100th of a raindrop) is placed through the sample hole and onto the platform from an automated micropipette and sample changer. The tube is heated electrically

by passing a current through it in a pre-programmed series of steps. The details will vary with the sample but typically they might be 30–40 seconds at 150°C to evaporate the solvent, 30 seconds at 600°C to drive off any volatile organic material and char the sample to ash, and with a very fast heating rate (*ca* 1500 °C s⁻¹) to 2000– 2500°C for 5–10 seconds to vaporise and atomise elements (including the element being analysed). Finally heating the tube to a still higher temperature -ca 2700°C – cleans it ready for the next sample. During this heating cycle the graphite tube is flushed with argon gas to prevent the tube burning away. In electro thermal atomisation almost 100% of the sample is atomised. This makes the technique much more sensitive than flame AAS.

Sample preparation

Sample preparation is often simple, and the chemical form of the element is usually unimportant. This is because atomisation converts the sample into free atoms irrespective of its initial state. The sample is weighed and made into a solution by suitable dilution. Elements in biological fluids such as urine and blood are often measured simply after a dilution flow injection accessory.

When making reference solutions of the element under analysis, for calibration, the chemical environment of the sample should be matched as closely as possible – *i.e.* the analyte should be in the same compound and the same solvent. Teflon containers may be used when analysing very dilute solutions because elements such as lead are sometimes leached out of glass vessels and can affect the results

Background absorption

It is possible that other atoms or molecules apart from those of the element being determined will absorb or scatter some radiation from the light source. These species could include un vaporised solvent droplets, or compounds of the matrix (chemical species, such as anions, that tend to accompany the metals being analysed) that are not removed completely. This means that there is background absorption as well as that of the sample.

One way of measuring and correcting this background absorption is to use two light sources, one of which is the hollow cathode lamp appropriate to the element being measured. The second light source is a deuterium lamp.

The deuterium lamp produces broad band radiation, not specific spectral lines as with a hollow cathode lamp. By alternating the measurements of the two light sources – generally at 50 -100 Hz – the total absorption (absorption due to analyte atoms plus background) is measured with the specific light from the hollow cathode lamp and the background absorption is measured with the light from the deuterium lamp. Subtracting the background from the total absorption gives the absorption arising from only analyte atoms.

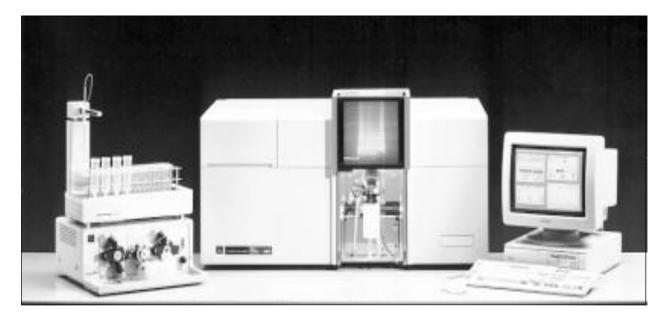


Figure 7

Calibration

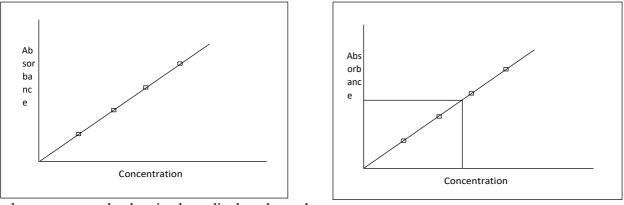
A calibration curve is used to determine the unknown concentration of an element -eg lead -

in a solution.

20 | P a g e

The instrument is calibrated using several solutions of known concentrations. A calibration curve is produced which is continually rescaled as more concentrated solutions are used – the more concentrated solutions absorb more radiation up to a certain absorbance. The calibration curve shows the concentration against the amount of radiation absorbed (*Fig.* 8(a)).

The sample solution is fed into the instrument and the unknown concentration of the



element -e.g. lead - is then displayed on the

calibration curve (*Fig.* 8(*b*)).

Figure 8(a) and Figure 8(b)

Interferences and matrix modification other chemicals that are present in the sample may affect the atomisation process. For example, in flame atomic absorption, phosphate ions may react with calcium ions to form calcium pyrophosphate. This does not dissociate in the flame and therefore results in a low reading for calcium. This problem is avoided by adding different reagents to the sample that may react with the phosphate to give a more volatile compound that is dissociated easily. Lanthanum nitrate solution is added to samples containing calcium to tie up the phosphate and to allow the calcium to be atomised, making the calcium absorbance independent of the amount of phosphate. With electrothermal atomisation, chemical modifiers can be added which react with an interfering substance in the sample to make it more volatile than the analyte compound. This volatile component vaporises at a relatively low temperature and is removed during the low and medium temperature stages of electrothermal atomisation.

Chapter-IV

METHODOLOGY

Required materials:

- FeCl3.6H2O (MOLYCHEM)
- Lactic acid (AVRA)
- Activated charcoal (SDFCL)
- Ultra-pure water (demineralized)
- Distilled water
- 2% nitric acid
- Volumetric flask 250 mL (Borosilicate glass)
- Watch glass
- Hot air oven (BIO-TECHNICS INDIA)
- Digital weighing machine (CITIZEN)
- AAS (ICE 3300)

Procedure:

- Using a 250ml volumetric flask, prepare 100 ppm of Ferric chloride hexahydrate metal solution.
- For the preparation of 100 ppm of Ferric chloride hexahydrate metal solution we have to calculate the amount of Ferric chloride hexahydrate required for the solution.
- Now add 3 drops of lactic acid in the Ferric chloride hexahydrate metal solution using a dropper, then we obtain the metal ligand solution which will be in 1:2 ratio.
- Keep the solution for 24 hrs. for better results.
- Now take 100ml of metal ligand solution in a 500ml beaker and add 5 grams of Activated Charcoal to it.
- Keep this solution aside for 48 hours. To obtain a metal ligand solution absorbed charcoal, filter the solution using a filter paper and rinse it 3 times using distilled water.
- Take the metals adsorbed activated Charcoal on a watch glass and keep it in the oven

at 60°C for 10 hours to get rid of moisture.

- After getting rid of moisture, we get fine powder of metal adsorbed activated Charcoal.
- Later shift the compound to a glass valve.
- Now send the adsorbent sample to the AAS for further tests.
- Here in AAS an Atomic Absorption Spectrometer is used for the tests, which consists of different bulbs that have wavelengths of different elements.
- The spectrometer cannot intake the solid particles, it can only intake the liquid.
- So the given adsorbent sample is mixed with the nitric acid solution.
- Now the adsorbent sample that is mixed with nitric acid is heated in the fume hood at 100°C for 15 minutes.
- The test sample is reduced to slurry liquid.
- Not all of the adsorbent samples given at the AAS are mixed, but 0.5 gm. of the adsorbent sample is diluted in 25mL of nitric acid solution (in which the nitric acid is present 2% in the ultra-pure water).
- Now filter the slurry liquid in thick filter paper. The ultra-pure water is used in rinsing of the adsorbent sample to 25mL, as the sample that should be used in the spectrometer has reduced because of the heating.
- Rinsing the slurry liquid with the ultra-pure water also helps the metal ions present in the adsorbent sample, flow with the nitric acid present in the water for the test purpose.
- Now 5 test standards are prepared using 1ppm, 2ppm, 3ppm, 4ppm and 5ppm "Fe" in it.
- Firstly the spectrometer is checked using only ultra-pure water with 2% nitric acid

present in it.

- After that it is tested with the 5 different prepared Fe standards.
- Now our test sample is tested after the 5 Fe standards are tested.
- The bulb present in the spectrometer is set at the same wavelengths as of the "Fe".
- When the test sample is being tested the bulb that consists of the "Fe" wavelength automatically transmits the same wavelengths through the sample.
- When the test sample is being tested in the spectrometer the Fe ions present with nitric acid are burned with the burner present in the spectrometer.
- As the concentration of the Fe increases the temperature also increases and the color gradually turns from orange to red and red to crimson and so on... The average temperature is around 1500°C.
- The waste present in the test sample other than Fe and nitric acid after the completion of tests are collected in the waste collector present below.
- The signal from the spectrometer can be seen in the monitor, which is already connected to the spectrometer.
- The signal from the spectrometer can be recorded and calculated from the graphs formed and numeric values shown in the monitor.
- Same Experiment carried out without the interference/addition of Ligand i.e. Lactic acid for Control Experiment.



AA spectrometer (a)



AA spectrometer (b)



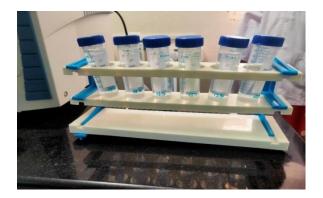
<u>Model no. ICE 3300</u>



Fumehood (a)



<u>Fume hood (b)</u>



Fe standards with test sample



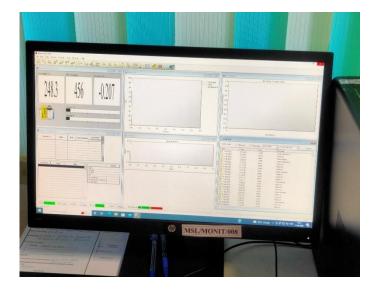
Fe standards with test sample



Burning of Nitric acid solution in AA Spectometer



Burning of Test sample in AA Spectometer



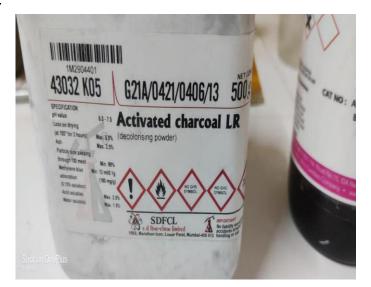
Values before intake of test samples

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Values after intake of test samples



Lactic acid



Activated charcoal



Metal ligand solution (a)



<u>Glass plate</u>



<u>Spatula</u>



Measuring Tube



<u>Hot air Oven</u>



5gm of Activated charcoal in metal ligand solution



Metal adsorbed charcoal





Test Samples















| P a g e







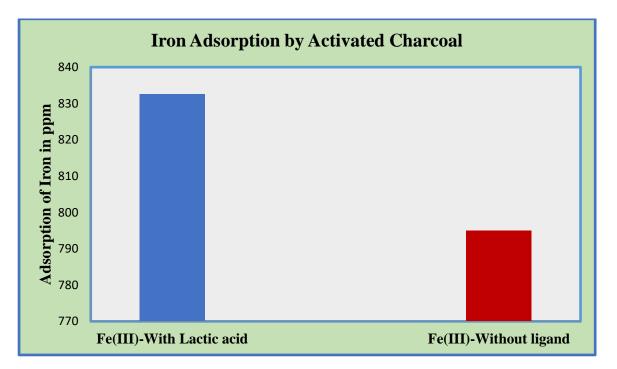
| P a g e



Chapter-V

RESULTS AND DISCUSSION

Activated charcoal adsorbed **832.5** ppm of Iron metal from aqueous solution of Fe(III)-Lactic acid metal ligand solution. Whereas, Activated charcoal adsorbed only **795.0** ppm when Lactic acid is absent. It is evident from the AAS results, ligand involvement enhanced the metal adsorption by initiating potential chemical interactions between adsorbate and adsorbent. Lactic acid strongly form coordination bonding with Fe(III) to form a soluble complex in aqueous solution. The ligand capped Fe(III) complex strongly adsorbed to Activated charcoal by establishing chemical bonding. From the AAS results, it is conclusive that **4.71** % of adsorption increased in the presence of Lactic acid as chelating agent.



Impact of Lactic acid on Adsorption of Fe (III) ions from aqueous solution by Activated Charcoal

Spectrometer Parameters – **<u>Fe</u>:**

Element : Fe	Measurement mode : Absorbance	
Wavelength : 248.3nm	Band pass : 0.2nm	Lamp current : 75%
Background correction : D2	High Resolution : Off	Optimise Spectrometer Parameters : No
Signal type : continuous	Resamples : Fast	Number of resamples : 3
Measurement time : 4.0secs	Flier mode : No	
Use RSD Test : No		

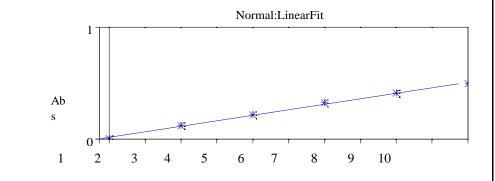
<u>Flame Parameters – Fe:</u>

Flame type : Air – C2H2	Fuel Flow : 0.9L/min	Auxiliary Oxidant : Off
Nebuliser Uptake : 4secs	Bunsen Stabilisation : Omins	Optimise Fuel Flow : No
Burner Height :7.0mm	Optimise Burner fuel : No	

<u>Calibration Parameters-Fe</u>:

Calibration mode : Normal	Line fit : Linear	Use stored calibration : No
Concentration units : mg/L	Scales units : mg/L	Scaling factor : 1.0000
Acceptable fit : 0.990	Rescale Limits : 10.0%	Failure Action : Flag and Continue
Standard 1 - 2.0000	Standard 4 - 8.0000	
Standard 2 - 4.0000	Standard 5 - 10.0000	
Standard 3 - 6.0000		

Solutions Results-Fe:



Y = 0.04924x + 0.0141 Fit: 0.9965 Characteristic Conc.: 0.0894

Sample ID	Signal	RSD	Conc.
	Abs	%	Mg/L
Fe Blank	0.001	35.6	0.0000
1	0.001	Back	ground: -0.003
2	0.001	Background: -0.003	
3	0.000	Back	ground: -0.003
Fe Standard 1	0.120	0.3	2.0000
1	0.120	Back	ground: 0.003
2	0.119	Back	ground: 0.003
3	0.120	Backs	ground: 0.003
Fe Standard 2	0.215	0.3	4.0000
1	0.215	Backs	ground: 0.004
2	0.215	Backs	ground:0.004
3	0.216	Backs	ground:0.004
Fe Standard 3	0.322	0.2	6.0000
1	0.323	Backs	ground: 0.005
2	0.322	Background: 0.005	
3	0.322	Background: 0.005	
Fe Standard 4	0.411	0.3	8.0000
1	0.410	Backş	ground: 0.005
2	0.412	Backs	ground: 0.005
3	0.412	Background: 0.005	
Fe Standard 5	0.493	0.1	10.0000
1	0.494	Background: 0.005	
2	0.494	Backs	ground: 0.005
3	0.494	Backy	ground: 0.006
Fe(III)-Lactic acid- Activated Charcoal	0.834	0.1	16.6490 C
1	0.834	Back	ground: 0.005
2	0.833	Back	ground: 0.005
3	0.834	Backs	ground: 0.005

Fe(III)-Activated Charcoal (Without Ligand)	0.796	0.1 15.8990 C
1	0.795	Background: 0.005
2	0.797	Background: 0.005
3	0.796	Background: 0.005

Test Results:

S.no	Test Parameters	Sample	Results
01.	Iron by AAS	Fe(III)-Lactic acid –	832.5ppm
	Analysis: (ppm)	Activated charcoal	
		Sample	
02.	Iron by AAS	Fe(III)–Activated	795.0 PPM
	Analysis: (ppm)	charcoal (control)	

CONCLUSION:

From the current project it is clear that Lactic acid as a Chelating agent has played a vital role in adsorption of Fe (III) ions from aqueous solution by Activated charcoal and increases the adsorption up to **4.716**%. This aspect will be useful in designing the newer strategies of Heavy metal Remediation techniques using Natural Chelating Ligands as Facilitating agents in Metal Adsorption processes.

<u>References</u>:

- 1. A. A. El-Asmy, E. M. Saad, M. S. El-Shahawi, Transit. Metal Chem., 19, 406 (1994).
- 2. A. P. B. Lever, Inorganic electronic spectroscopy 1, 2, Mir, Moskva, 1987.
- 3. A. Cotton, G. Wilkinson, Advanced inorganic chemistry, 3rd ed., John Wiley and Sons, New YorkLondon-Sydney, 1976.
- A. Alahabadi, H. Malvandi, Contamination and ecological risk assessment of heavy metals and metalloids in surface sediments of the Tajan River, Iran. Mar. Pollut. Bull., 133 (2018) 741–749.
- A. Kolbasov, S. Sinha-Ray, A.L. Yarin, B. Pourdeyhimi, Heavy metal adsorption on solution-blown biopolymer nanofiber membranes, J. Membr. Sci., 530 (2017) 250– 263.
- A.R. Kaveeshwar, P.S. Kumar, E.D. Revellame, D.D. Gang, M.E. Zappi, R. Subramaniam, Adsorption properties and mechanism of barium (II) and strontium (II) removal from fracking wastewater using pecan shell based activated carbon, J. Clean. Prod., 193 (2018) 1–13.
- A. Ozcan, A.S. Ozcan, S. Tunali, T. Akar, I. Kiran, Determination of the equilibrium, kinetic and thermodynamic parameters of adsorption of copper (II) ions onto seeds of *Capsicum annuum*, J. Hazard. Mater., 124 (2005) 200–208.
- A. Naghizadeh, R. Nabizadeh, Removal of reactive blue 29 dye by adsorption on modified chitosan in the presence of hydrogen peroxide[J], Environ. Protect. Eng., 42 (2016) 149–168.
- 9. A. Mohseni-Bandpi, D.J. Elliott, M.A. Zazouli, Biological nitrate removal processes from drinking water supply-a review[J], J. Environ. Sci. Eng., 11 (2013) 35–35.
- 10. A. Vicek, Jr., Coord. Chem. Rev., 2000, 200-202, 933.
- 11. A. Pigliucci, G. Duvanel, M. L. Lawson Daku, E. Vauthey, J. Phys. Chem. A, 2007, 111, 6135.
- 12. B. S. Faust, R. G. zepp, Environ. Sci. Technol., 1993, 27, 2517.
- B. Abei, J. Assmann, M. Buback, C. Grimm, M. Kling, S. Schmatz, J. Schroeder, T. Witte, J. Phys. Chem. A, 2003, 107, 9499.
- 14. Voelker, F. M. M. Morel, B. Sulzberger, Environ. Sci. Technol., 1997, 31, 1004.
- 15. B. Sarkar, S. Mandal, Y.F. Tsang, P. Kumar, K.H. Kim, Y.S. Ok, Designer carbon nanotubes for contaminant removal in water and wastewater: a critical review, Sci. Total Environ., 612 (2018) 561–581.
- 16. B. Kaličanin, R. Nikolić, in: Wide spectra of quality control, Akyr, Isin (eds), Croatia: InTech, Rijeka, 2011, p. 211.
- 17. C.M. Park, D. Wang, J. Han, J. Heo, C. Su, Evaluation of the colloidal stability and adsorption performance of reduced graphene oxide-elemental silver/magnetite

nanohybrids for selected toxic heavy metals in aqueous solutions, Appl. Surf. Sci., 471 (2019) 8–17.

- 18. C.G. Lee, M.K. Song, J.C. Ryu, C. Park, J.W. Choi, S.H. Lee, Application of carbon foam for heavy metal removal from industrial plating wastewater and toxicity evaluation of the adsorbent, Chemosphere, 153 (2016) 1–9.
- C.G. Lee, S. Lee, J.A. Park, C. Park, S.J. Lee, S.B. Kim, B. An, S.T. Yun, S.H. Lee, J. W. Choi, Removal of copper, nickel and chromium mixtures from metal plating wastewater by adsorption with modified carbon foam, Chemosphere, 166 (2017) 203–211.
- C. Chen, X. Yang, J. Wei, X. Tan, X. Wang, Eu(III) uptake on rectorite in the presence of humic acid: a macroscopic and spectroscopic study, J. Colloid Interface Sci., 393 (2013) 249–256.
- 21. C.B. Godiya, X. Cheng, D. Li, Z. Chen, X. Lu, Carboxymethyl cellulose/polyacrylamide composite hydrogel for cascaded treatment/reuse of heavy metal ions in wastewater, J. Hazard. Mater., 364 (2019) 28–38.
- C. Dendrinou-Samara, G. Tsotsou, L. V.Ekateriniadou, A. H. Kortsaris, C. P. Raptopoulou, A. Terzis, D. A. Kyriakidis, D. P. Kessissoglou, *J. Inorg. Biochem.*, 71, 171 (1998).
- 23. C. E. Mortimer, Chemistry, 5th ed. Litton Educational Publishing Inc., Belmont, California 94002, 1983.
- 24. C. J. Miles, P. L. Brezonik, Environ. Sci. Technol., 1981, 15, 1089.
- 25. D. E. Falvey, G. B. Schuster, J. Am. Chem. Soc., 1986, 108, 7419.
- 26. D. Jensen, A. Ledin, T. Christensen, W.A. Wenzel, Speciation of heavy metals in landfill-leachate polluted groundwater, Water Res., 33 (1999) 2642–2650.
- 27. D. Eeshwarasinghe, P. Loganathan, S. Vigneswaran, Simultaneous removal of polycyclic aromatic hydrocarbons and heavy metals from water using granular activated carbon, Chemosphere, 223 (2019) 616–627.
- 28. D. H. Anđelković, R. S. Nikolić, D. Z. Marković, T. D. Anđelković, G. M. Kocić, Z. B. Todorović, A. Lj. Bojić, J. Serb. Chem. Soc., 78(1), 137 (2012).
- 29. E. M. Glebov, 1. P. Pozdnyakov, V. P. Grivin, V. F. Plyusnin, X. Zhang, F. Wu, N. Deng, Photochem. Photobiol. Sci., 2011, 10, 425.
- 30. E. Mentasti, Inorg. Chem., 1979, 18, 1512.
- 31. E. A. Juban, J. K. McCusker, J. Am. Chem. Soc., 2005, 127, 6857.
- 32. E. Da'na, A. Awad, Regeneration of spent activated carbon obtained from home filtration system and applying it for heavy metals adsorption, J. Environ. Chem. Eng., 5 (2017) 3091–3099.
- 33. E.S. Dragan, D. Humelnicu, M.V. Dinu, R.I. Olariu, Kinetics, equilibrium modeling, and thermodynamics on removal of Cr (VI) ions from aqueous solution using novel composites with strong base anion exchanger microspheres embedded into chitosan/poly(vinyl amine) cryogels, Chem. Eng. J., 330 (2017) 675–691.
- 34. E.E. ElSayed, Natural diatomite as an effective adsorbent for heavy metals in water and wastewater treatment (a batch study), Water Sci., 32 (2019) 32–43.

- 35. F. Lian, G. Cui, Z. Liu, L. Duo, G. Zhang, B. Xing, One-step synthesis of a novel Ndoped microporous biochar derived from crop straws with high dye adsorption capacity, J. Environ. Manage., 176 (2016) 61–68.
- 36. F. Cao, C. Lian, J. Yu, H. Yang, S. Lin, Study on the adsorption performance and competitive mechanism for heavy metal contaminants removal using novel multipore activated carbons derived from recyclable long-root *Eichhorniacrassipes*, Bioresour. Technol., 276 (2019) 211–218.
- 37. F. Weng, D. Nansheng, Chemosphere, 2000, 41, 1137.
- 38. F. wu, N. Deng, E. M. Glebov, 1. P. Pozdnyakov, V. P. Grivin, V. F. Plyusnin, N. M. Bazhin, Russ. Chem. Bull. (Int. Ed.), 2007, 56, 900 1/o. Akad. Nauk, Ser. Khim., 2007, 8661.
- F. Dimiza, F. Perdih, V. Tangoulis, I. Turel, D. Kessissoglou, G. Psomas, J. Inorg. Biochem., 105, 476 (2011).
- 40. G. G. Duka, D. G. Batyr, L. S. Romanchuk, A. Ya. Sychev, Koordinats. Khim., 1990, 16, 93 [Sov. Coord. Chem. (Engl. Transl.), 1990, 161.
- 41. G. Xiao, Y. Wang, S. Xu, P. Li, C. Yang, Y. Jin, Q. Sun, H. Su, Superior adsorption performance of graphitic carbon nitride nanosheets for both cationic and anionic heavy metals from wastewater, J. Chem. Eng., 27 (2019) 305–313.
- 42. G. Xiao, X. Zhang, H. Su, T. Tan, Plate column biosorption of Cu(II) on membranetype biosorbent (MBS) of Penicillium biomass: optimization using statistical design methods, Bioresour. Technol., 143 (2013) 490–498.
- 43. H. B. Abrahamson, A. B. Rezvani, J. G. Brushmiller, InorgChim. Acta, 1994, 226, 117.
- 44. H. Gao, R. G. zepp, Environ. Sci. Technol., 1998, 32, 2940.
- 45. H. B. Abrahamson, A B. Rezvani, J. G. Brushmiller, Inorg. Chim. Acta, 1994, 226, 117.
- 46. H.-J. Benkelberg, P. Warneck, J. Phys. Chem., 1995, 99, 5214.
- 47. H. Guedidi, L. Reinert, J. M. Lévêque, Y. Soneda, N. Bellakhal, L. Duclaux, The effects of the surface oxidation of activated carbon, the solution pH and the temperature on adsorption of ibuprofen, Carbon, 54 (2013) 432–443.
- 48. H.A. Hegazi, Removal of heavy metals from wastewater using agricultural and industrial wastes as adsorbents, HBRC J., 9 (2013) 65–76.
- 49. H. Xu, H. Yuan, J. Yu, S. Lin, Study on the competitive adsorption and correlational mechanism for heavy metal ions using the carboxylated magnetic iron oxide nanoparticles (MNPs-COOH) as efficient adsorbents, Appl. Surf. Sci., 473 (2019) 960–966.
- 50. I.A. Aguayo-Villarreal, A. Bonilla-Petriciolet, R. Muñiz-Valencia, Preparation of activated carbons from pecan nutshell and their application in the antagonistic adsorption of heavy metal ions, J. Mol. Liq., 230 (2017) 686–695.
- 51. I. P. Pozdnyakov, O. V. Kel, V. F. Plyusnin, V. P. Grivin, N. M. Bazhin, Phys. Chem. n, 2008, 112, 8316.

- 52. I. P. Pozdnyakov, E. M. Glebov, V. F. Plyusnin, V. P. Grivin, E. Bunduki, N. V. Goryacheva, V. Gladkii, G. G. Duka, Khim. Vys. Energii, 2009, 43, 461 [High Energy Chem. (Engl. Thansl.), 2009, 43, 4061.
- 53. I. P. Pozdnyakov, A. V. Kolomeets, V. F. Plyusnin, A A. Melnikov, V. O. Kompanets, S. V. Chekalin, N. Tkachenko, H. Lemmetyinen, Chem. Phys. Lett., 2012, 530, 45.
- 54. I. P. Pozdnyakov, V. F. Plyusnin, V. P. Grivin, D. Yu. Vorobyev, N. M. Bazhin, E. Vauthey, J. Photochem. Photobiol., A: Chem., 2006, 182, 75.
- 55. J. G. Calvert, Jr., J. N. Pitts, Photochemistry, Wiley, New York, 1966, 782 pp.
- 56. J. W. Hilborn, J. A. Pincock, J. Am. Chem. Soc., 1991, 113, 2683.
- 57. J. Shima, J. Makanova, Coord. Chem. Rev., 1997, 160, 161.
- 58. J. Jortnor, G. J. Stein, Phys. Chem., 1962, 66, 1264.
- 59. J.B. Dima, C. Sequeiros, N.E. Zaritzky, Hexavalent chromium removal in contaminated water using reticulated chitosan micro/ nanoparticles from seafood processing wastes, Chemosphere, 141 (2015) 100–111.
- 60. J. Shen, G. Huang, C. An, X. Xin, C. Huang, S. Rosendahl, Removal of Tetrabromobisphenol A by adsorption on pineconederived activated charcoals: synchrotron FTIR, kinetics and surface functionality analyses, Bioresour. Technol., 247 (2018) 812–820.
- 61. J. Valentin-Reyes, R.B. Garcia-Reyes, A. Garcia-Gonzalez, E. Soto-Regalado, F. Cerino-Cordova, Adsorption mechanisms of hexavalent chromium from aqueous solutions on modified activated carbons, J. Environ. Manage., 236 (2019) 815–822.
- J. Sun, M. Li, Z. Zhang, J. Guo, Unravelling the adsorption disparity mechanism of heavy-metal ions on the biomass derived hierarchically porous carbon, Appl. Surf. Sci., 471 (2019) 615–620.
- 63. J. Sun, Z. Zhang, J. Ji, M. Dou, F. Wang, Removal of Cr⁶⁺ from wastewater via adsorption with high-specific-surface-area nitrogen-doped hierarchical porous carbon derived from silkworm cocoon, Appl. Surf. Sci., 405 (2017) 372–379.
- 64. J. Piispanen, L. H. J. Lajunen, Acta Chem. Scand., 49, 241 (1995).
- 65. J. M. Jovanović, R. S. Nikolić, G. M. Kocić, N. S. Krstić, M. M. Krsmanović, J. Serb. Chem. Soc., **78(2)**, 197 (2013).
- 66. K. Nakamoto, Infrared and Raman spectra of inorganic and coordination compounds, Part A and B, 6th ed., John Wiley and Sons Inc., New Jersey, 2009.
- 67. L. Patterson, R. Small, J. Scaiano, Radiat. Res., 1977, 72, 218.
- 68. L. Vafajoo, R. Cheraghi, R. Dabbagh, G. McKay, Removal of cobalt (II) ions from aqueous solutions utilizing the pre-treated 2-Hypnea Valentiae algae: equilibrium, thermodynamic, and dynamic studies, Chem. Eng. J., 331 (2018) 39–47.
- 69. L. Wang, Y. Wang, F. Ma, V. Tankpa, S. Bai, X. Guo, X. Wang, Mechanisms and reutilization of modified biochar used for removal of heavy metals from wastewater: a review, Sci. Total Environ., 668 (2019) 1298–1309.
- L.Y. Li, X. Gong, O. Abida, Waste-to-resources: exploratory surface modification of sludge-based activated carbon by nitric acid for heavy metal adsorption, Waste Manage., 87 (2019) 375–386.

- 71. L. Li, F. Zhu, Y. Lu, J. Guan, Synthesis, adsorption and selectivity of inverse emulsion Cd (II) imprinted polymers, J. Chem. Eng., 26 (2018) 494–500.
- 72. M.G. Kiran, K. Pakshirajan, G. Das, Heavy metal removal from multicomponent system by sulfate reducing bacteria: mechanism and cell surface characterization, J. Hazard. Mater., 324 (2017) 62–70.
- 73. M. Bilal, J.A. Shah, T. Ashfaq, S.M. Gardazi, A.A. Tahir, A. Pervez, H. Haroon, Q. Mahmood, Waste biomass adsorbents for copper removal from industrial wastewater-a review, J. Hazard. Mater., 263 (2013) 322–333.
- 74. M. N. Schuchmann, C. von Sonntag, J. Am. Chem. Soc., 1988, 110, 5698
- 75. M. Tuzen, M. Soylak, Multi-element coprecipitation for separation and enrichment of heavy metal ions for their flame atomic absorption spectrometric determinations, J. Hazard. Mater., 162 (2009) 724–729.
- 76. M.R. Awual, M.A. Shenashen, T. Yaita, H. Shiwaku, A. Jyo, Efficient arsenic (V) removal from water by ligand exchange fibrous adsorbent, Water Res., 46 (2012) 5541–5550.
- 77. M.M. Rao, A. Ramesh, G.P. Rao, K. Seshaiah, Removal of copper and cadmium from the aqueous solutions by activated carbon derived from *Ceibapentandra* hulls, J. Hazard. Mater., 129 (2006) 123–129.
- M. Bansal, D. Singh, V.K. Garg, A comparative study for the removal of hexavalent chromium from aqueous solution by agriculture wastes' carbons, J. Hazard. Mater., 171 (2009) 83–92.
- 79. M.F.N. Secondes, V. Naddeo, V. Belgiorno, F. Ballesteros Jr., Removal of emerging contaminants by simultaneous application of membrane ultrafiltration, activated carbon adsorption, and ultrasound irradiation, J. Hazard. Mater., 264 (2014) 342–349.
- 80. M. Obradović, D. Veselinović, P. Đurđević, Physicochemical methods of testing the balance of the complexing environments, Faculty of Philosophy, Niš, Faculty of Physical Chemistry, Belgrade, 1996. (in Serbian)
- 81. M. A. Agotegaray, M. Dennehy, M. A. Boeris, M. A. Grela, R. A. Burrow, O. V. Quinzani, *Polyhedron*, **34**, 74 (2012).
- 82. N. A. El-Ragehy, M. Abdelkawy, A. El-Bayoumy, Anal. Lett., 27(11), 2127 (1994).
- 83. N. Krstić, R. Nikolić, M. Krstić, Hem. preg., 55(6), 154 (2014) (in Serbian).
- 84. N. S. Krstić, R. S. Nikolić, M. N. Stanković, N. G. Nikolić, D. M. Đorđević, *Trop. J. Pharm. Res.*, **14(2)**, 337 (2014).
- 85. N. Deng, F. wu, F. Luo, M. Xiao, Chemosphere, 1998, 36, 3101.
- 86. P. Natarajan, J. F. Endicott, J. Phys. Chem., 1973, 77, 2049.
- P. Kocot, K. Szacilowski, Z. J. Stasicka, Photochem. Photobiol. A: Chem. 2007, 188, 128.
- 88. P. Tan, J. Sun, Y. Hu, Z. Fang, Q. Bi, Y. Chen, J. Cheng, Adsorption of Cu²⁺, Cd²⁺ and Ni²⁺ from aqueous single metal solutions on graphene oxide membranes, J. Hazard. Mater., 297 (2015) 251–260.

- 89. P. Tan, Y. Hu, Q. Bi, Competitive adsorption of Cu²⁺, Cd²⁺ and Ni²⁺ from an aqueous solution on graphene oxide membranes, Colloids Surf. A, 509 (2016) 56–64.
- 90. R. Tovar-Gómez, M.D.R. Moreno-Virgen, J. Moreno-Pérez, A. Bonilla-Petriciolet, V. Hernández-Montoya, C.J. DuránValle, Analysis of synergistic and antagonistic adsorption of heavy metals and acid blue 25 on activated carbon from ternary systems, Chem. Eng. Res. Des., 93 (2015) 755–772.
- 91. R. Carballo, B. Covelo, S. Balboa, A. Castiñeiras, J. Niclós, Z. Anorg. Allg. Chem., 627, 948 (2001).
- 92. R. Battino, T. R. Rettich, T. Tominaga, J. Phys. Chem. Ref. Data, 1983, 12, 163.
- 93. R. Carballo, A. Castiñeiras, S. Balboa, B. Covelo, J. Niclós, *Polyhedron*, **21**, 2811 (2002).
- 94. R. P. Sharma, S. Singh, A. Singh, V. Ferreti, J. Mol. Struct., 918, 188 (2009).
- 95. R. R. Crichton, Biological Inorganic Chemistry An Introduction, Elsevier, 2008.
- 96. R. Nikolić, B. Kaličanin, N. Krstić, Connect. Tissue Res., 53(3), 229 (2012).
- R. S. Nikolić, N. V. Radosavljević-Stevanović, T. D.Anđelković, M. N. Stanković, N. S. Krstić, *J. Serb. Chem. Soc.*, **79**, 1395 (2014).
- 98. R. S. Nikolić, N. S. Krstić, G. M. Nikolić, G. M. Kocić, M. D. Cakić, D. H. Anđelković, *Polyhedron*, **80**, 223 (2014).
- 99. R. Drago, Physical methods in chemistry 1, 2, Mir, Moskva, 1981.
- 100. S. Spasić, Z. Jelić-Ivanović, V. Spasojević-Kalimanovska, Fundamentals of biochemistry, Belgrade, 2000. (in Serbian)
- S. S. Mitić, G. Z. Miletić, A. N. Pavlović, B. B Arsić, V. V. Živanović, J. Serb. Chem. Soc., 73(8-9), 879 (2008).
- 102. S. V. Chekalin, Usp. Piz. Nauk, 2006, 176, 657 IPhys.-Usp. (Engl. Transl.), 2006, 49, 6341.
- 103. S. Goldstein, J. Rabani, J. Photochem. Photobiol. A: Chem., 2008, 193, 50.
- 104. T. M. Bockman, S. M. Hubig, J. K. Kochi, J. Org. Chem., 1997, 62, 2210.
- 105. T.A. Kurniawan, G.Y.S. Chan, W. H. Lo, S. Babel, Physicochemical treatment techniques for wastewater laden with heavy metals, Chem. Eng. J., 118 (2006) 83–98.
- 106. T.A.H. Nguyen, H.H. Ngo, W.S. Guo, J. Zhang, S. Liang, Q.Y. Yue, Q. Li, T.V. Nguyen, Applicability of agricultural waste and by-products for adsorptive removal of heavy metals from wastewater, Bioresour. Technol., 148 (2013) 574–585.
- V. Balzani, V. Carassiti, Photochemistry ofCoordination Compounds, Academic Press, London—New York, 1970, 432 pp.
- 108. V. Nadtochenko, J. Kiwi, J. Photochem. Photobiol. A: Chem., 1996, 99, 145.
- 109. V. Nejadshafiee, M.R. Islami, Adsorption capacity of heavy metal ions using sultone-modified magnetic activated carbon as a bio-adsorbent, Mater. Sci. Eng. C, 101 (2019) 42–52.

- 110. W. Zhang, H. Feng, J. Chang, J. Qu, H. Xie, L. Yu, Heavy metal contamination in surface sediments of Yangtze River intertidal zone: an assessment from different indexes, Environ. Pollut., 157 (2009) 1533–1543.
- X. Deng, L. Lu, H. Li, F. Luo, The adsorption properties of Pb (II) and Cd (II) on functionalized graphene prepared by electrolysis method, J. Hazard. Mater., 183 (2010) 923–930.
- X. Zhang, Y. Gong, F. wu, N. Deng, 1. P. Pozdnyakov, E. M. Glebov, V. P. Grivin, V. F. Plyusnin, N. M. Bazhin, Russ. Chem. Bull. (Int. Ed.), 2009, 58, 1828 1/zv. Akad. Nauk, Ser. Khim., 2009, 17711.
- X. Luo, Z. Zhang, P. Zhou, Y. Liu, G. Ma, Z. Lei, Synergic adsorption of acid blue 80 and heavy metal ions (Cu²⁺/Ni²⁺) onto activated carbon and its mechanisms, J. Ind. Eng. Chem., 27 (2015) 164–174.
- 114. X. Cai, J. He, L. Chen, K. Chen, Y. Li, K. Zhang, Z. Jin, J. Liu, C. Wang, X. Wang, A 2D-g-C3N4 nanosheet as an eco-friendly adsorbent for various environmental pollutants in water, Chemosphere, 171 (2017) 192–201.
- 115. Y. Wu, H. Pang, Y. Liu, X. Wang, S. Yu, D. Fu, J. Chen, X. Wang, Environmental remediation of heavy metal ions by novelnanomaterials: a review, Environ. Pollut., 246 (2019) 608–620.
- 116. Y. Cui, Q. Ge, X.Y. Liu, T.S. Chung, Novel forward osmosis process to effectively remove heavy metal ions, J. Membr. Sci., 467 (2014) 188–194.
- 117. Y. Zuo, J. Hoigne, Afrnospher. Environ., 1994, 28, No. 7, 1231.
- 118. Y. Ahmadi, S.E. Eshraghi, P. Bahrami, M. Hasanbeygi, Y. Kazemzadeh, A. Vahedian, Comprehensive Water-AlternatingGas (WAG) injection study to evaluate the most effective method based on heavy oil recovery and asphaltene precipitation tests, J. Pet. Sci. Eng., 133 (2015) 123–129.

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Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

General Parameters

Method : Iron (Fe) Autosampler : None Use SFI: No Operator : Jyothi

Instrument Mode: Flame Dilution: None

mA Hours

n/a

Report Date: 11-06-2022 17:40:20

Valid Method Signatures

11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Method Audit Trail

11-06-2022 17:22:18 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC

Signed with Reason : Approved by:

Analysis Details

Spectrometer: iCE 3000 AA01204906 v1.30

Analysis Name: Iron (Fe) 11-06-2022 Operator Name: Jyothi

n/a

Lamp Information Serial Number

Element(s) Fe

. . .

Deuterium Lamp Hours: 68.34

Sequence Table

Shared Standards: Yes	
Action	Fe
Calibration	√
B.No.Group-II	√
B.No.Group-III	√
B.No.Group-IV	√
B.No.Group-V	√
B.No.Group-VI	✓

Sample Details

		Nominal Mass: 1.0000		
No.	Sample Id	Sample Mass	Dilution Ratio	
1	B.No.Group-II	1.0000	1.0000	
2	B.No.Group-III	1.0000	1.0000	
3	B.No.Group-IV	1.0000	1.0000	
4	B.No.Group-V	1.0000	1.0000	
5	B.No.Group-VI	1.0000	1.0000	

Valid Analysis Signatures

11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Analysis Audit Trail

- 11-06-2022 17:30:16 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
- 11-06-2022 17:38:27 Jyothi(M. Jyothi):DESKTOP-39TDEGC Error MD147 - Activity manually aborted by user.
- 11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
- 11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

MART SPECIALITIES LAB LLP.

Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Spectrometer Parameters - Fe

Element: Fe Wavelength: 248.3nm Background Correction: D2 Signal Type: Continuous Measurement Time: 4.0secs Use RSD Test: No

Flame Type: Air-C2H2 Nebuliser Uptake: 4secs Burner Height: 7.0mm

Sampling: None

Calibration Mode: Normal	
Concentration Units: mg/L	
Acceptable Fit: 0.990	
Standard 1	2.0000
Standard 2	4.0000
Standard 3	6.0000

No changes are recorded for this element

Measurement Mode: Absorbance Bandpass: 0.2nm High Resolution: Off Resamples: Fast Flier Mode: No

Lamp Current: 75% Optimise Spectrometer Parameters: No Number Of Resamples: 3

Flame Parameters - Fe

Fuel Flow: 0.9L/min Burner Stabilisation: 0mins Optimise Burner Height: No Auxilary Oxidant: Off Optimise Fuel Flow: No

Sampling Parameters - Fe

Calibration Parameters - Fe

Line Fit: Linear Scaled Units: mg/L Rescale Limit: 10.0% Standard 4 Standard 5 Use Stored Calibration: No Scaling Factor: 1.0000 Failure Action: Flag and Continue 8.0000 10.0000

Element Audit Trail - Fe

Solution Results - Fe

Y = 0.04924x + 0.0141 Fit: 0.9965 Characteristic Conc: 0.0894

> 1 2 3

> 1 2 3

> 1 2 3

> 1 2 3

1 2

3

Sample ID

Fe Blank

Fe Standard 1

Fe Standard 2

Fe Standard 3

Fe Standard 4

	Normal: Linear Fit										
	Abs		*		*		*		-*-		*
	0 * -0-0	1	2	3	4	5	6	7	8	9	10
	-0-0		2	0		nc:mg/		,	0	0	10
Signal	Rsd	Conc				cted C					
Abs	%	mg/L			mg/L						
0.001	35.6	0.000)		Ũ						
0.001	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:26	i		
0.001	Background: -0.003				11-06	-2022 1	7:31:30)			
0.000	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:35			
0.120	0.3	2.0000)								
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:05			
0.119	-	ound: 0.00						7:32:09			
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:13			
0.215	0.3	4.0000)								
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:40)		
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:45			
0.216	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:49)		
0.322	0.2	6.0000)								
0.323	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:19)		
0.322	Background: 0.005				11-06	-2022 1	7:33:23				
0.322	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:27			
0.411	0.3	8.0000)								
0.410	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:59)		
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:03			
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:07			

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Operator Name: Jyothi

Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Solution Results - Fe

Sample ID	Signal	Rsd	Conc	Corrected Conc
-	Abs	%	mg/L	mg/L
Fe Standard 5	0.493	0.1	10.0000	
1	0.494	Backg	ground: 0.005	11-06-2022 17:34:38
2	0.494	Backg	ground: 0.005	11-06-2022 17:34:42
3	0.493	Backg	ground: 0.006	11-06-2022 17:34:46
Fe B.No.Group-II	0.834	0.1	16.6490 C	16.6490 C
1	0.834	Backg	ground: 0.005	11-06-2022 17:35:14
2	0.833	Backg	ground: 0.005	11-06-2022 17:35:19
3	0.834	Backg	ground: 0.005	11-06-2022 17:35:23
Fe B.No.Group-III	0.759	0.1	15.1286 C	15.1286 C
1	0.758	Backg	ground: 0.003	11-06-2022 17:35:52
2	0.760	Backg	ground: 0.003	11-06-2022 17:35:56
3	0.758	Backg	ground: 0.004	11-06-2022 17:36:00
Fe B.No.Group-IV	0.645	0.3	12.8189 C	12.8189 C
1	0.647	Backg	ground: 0.003	11-06-2022 17:36:31
2	0.644	Backg	ground: 0.003	11-06-2022 17:36:35
3	0.645	Backg	ground: 0.003	11-06-2022 17:36:40
Fe B.No.Group-V	0.809	0.2	16.1488 C	16.1488 C
1	0.808	Backg	ground: 0.005	11-06-2022 17:37:11
2	0.809	Backg	ground: 0.005	11-06-2022 17:37:15
3	0.811	Backg	ground: 0.005	11-06-2022 17:37:20
Fe B.No.Group-VI	0.711	0.3	14.1597 C	14.1597 C
1	0.709	Backg	ground: 0.005	11-06-2022 17:37:55
2	0.713	Backg	ground: 0.005	11-06-2022 17:37:59
3	0.712	Backg	ground: 0.004	11-06-2022 17:38:03

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-1 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
I	DETAILS OF THE SAMPLE
Sample Name : NA	
Name of the Manufacturer : NA	
Storage condition To b	up-I Mfg. : NA Exp. : NA e stored at room date date
	Derature Batch : NA C±3°C size
	by AAS Analysis.
	6/2022
Analysis Completion Date : 11/0	6/2022
Mfg. License No. : Not	provided
A.R.NO : NA	The Andrew Rest

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	620ppm

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



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Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-2 Sample Received Date : 11/06/2022 Report Date : 13/06/2022	
Q	DETAILS OF THE SAMPLE	
Sample Name : N	A	
Name of the Manufacturer : N	A	
Storage condition : To	roup-II Mfg. : NA Exp. : NA be stored at room date date	
Room Temperature : 25	Batch : NA size	
Tests Required:IreMethod:NAnalysis Starting Date:11	on by AAS Analysis.	
	Not provided	
A.R.NO : N	A	

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	832.5ppm

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



OCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00	
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinaga Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Sample Received Date : 11/06/2022	
a.y.	DETAILS OF THE SAMPLE	
Sample Name : N	JA	
Name of the Manufacturer : N	JA	
Storage condition	Group-III Mfg. : NA Exp. : NA To be stored at room date date Cemperature Batch : NA	
	5°C±3°C size	
	gm /	
Method : I Analysis Startin <mark>g Date : I</mark>	: NA tte : 11/06/2022	
and the second se	Not provided	
A.R.NO	NA	
Remark : S	Sample analyzed as received	

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	756.5ppm

×

Authorized Signatory

(Dr.R.Marayya)

Office: 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile: 98481 98350, 98481 25019 Mail: rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College	
Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000	Sample Received Date: 11/06/2022Report Date: 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name : N	A
Name of the Manufacturer : N	A
Storage condition : T	roup-IV Mfg. : NA Exp. : NA o be stored at room date date emperature Batch : NA
	5°C±3°C size
Method : N Analysis Starting Date : 1	on by AAS Analysis. A 1/06/2022 1/06/2022
Mfg. License <mark>No. :</mark> N	ot provided
A.R.NO : N	A
Remark : Sa	ample analyzed as received

Test Results

	and the second se	AN SECTION	
S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	641ppm	

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/	G
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Custome	r: Reference / Report No. : MSL/2022/JUNE/TARA/0064-5
Tara Government College	Sample Received Date : 11/06/2022
Prashanth Nagar Colony, Balajinag	ar Sumpto Received 2 mil
Sangareddy Telangana. 502000	Report Date : 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name :	NA
Name of the Manufacturer :	NA
Batch no : Storage condition	Group-VMfg. : NAExp. : NATo be stored at roomdatedateTemperatureBatch : NA
Room Temperature :	25°C±3°C size
Quantity Received : 4gm	
Tests Required:Iron by AAS Analysis.Method:NAAnalysis Starting Date:11/06/2022	
Analysis Completion Date : 11/06/2022	
Mfg. License No.	Not provided NA
A.R.NO Remark	Sample analyzed as received

Test Results

S.No.	Test Parameter	Result
01. Ir	on by AAS Analysis: (ppm)	807.5ppm

>

Authorized Signatory

(Dr.R.Marayya)

Office : 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile : 98481 98350, 98481 25019 Mail : rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-6 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
1	DETAILS OF THE SAMPLE
Storage condition	up-V1 Mfg. : NA Exp. : NA be stored at room date date perature Batch : NA C±3°C size
Tests Required:IronMethod:NAAnalysis Starting Date:11/0Analysis Completion Date:11/0Mfg. License No.:NotA.R.NO:NA	by AAS Analysis. 6/2022 6/2022 provided

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	708ppm

*

Authorized Signatory

(Dr.R.Marayya)

ADSORPTION OF Fe[III] ON CELLULOSE USING D-VALINE AS FACILITATING AGENT

Dissertation submitted in Partial fulfillment for the requirements for the award of degree of

Bachelor of Science ín CHEMISTRY

By

1.	K.Dheeraj Kumar,	B.Sc(Chemistry),	III	Year
2.	G.Laxmi kanth,	B.Sc(Chemistry),	III	Year
3.	K.Narasimha,	B.Sc(Chemistry),	III	Year
4.	G.Omkar,	B.Sc(Chemistry),	III	Year
5.	D.Anil,	B.Sc(Chemistry),	III	Year
6.	A.Jeremiah,	B.Sc(Chemistry),	III	Year

Under the Guídance:

Dr. K.ABHIJIT *HEAD* DEPARTMENT OF CHEMISTRY



Tara Government College, Sangareddy(A)

2021-22

<u>CERTIFICATE</u>

This is to certify that the project work entitled "ADSORPTION OF *Fe*[*III*] ON CELLULOSE USING D-VALINE AS FACILITATING AGENT" is presented by B.Sc (CHEMISTRY) students in partial fulfillment of the requirements for the degree of Bachelor of Science in Chemistry by the Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

The results embodied in this report have not been to any other University or Institution for the award of any degree.

(Dr.K.ABHIJIT)

Project Supervisor& Head, Department of Chemistry Tara Government College, Sangareddy(A)

EXTERNAL EVALUATOR

ACKNOWLEDGEMENTS

We express my deep gratitude to my research supervisor **Dr.K.Abhíjít**, Head, Department of Chemistry, Tara Govt. College, Sangareddy(A)-502001, INDIA for his inspiring guidance during the course of the Project work. The continuous encouragement extended by him propelled me to update my research skills and pedigree to engage in fruitful research.

We wish to express our gratitude to Smt. M.Praveena, Principal, Tara Govt. College, Sangareddy(A)-502001, INDIA for her constant support, cooperation and suggestions during the research work. We wish to express my sincere thanks to K.Sreedhar, Asst. Professor, Department of Chemistry for providing me facilities, help and support for the entire research work.

We express our special thanks to **MART Specialities Lab, Hyderabad** for providing technical assistance in Atomic absorption spectrometric analysis.

Finally, we take this opportunity to thank one and all that has directly or indirectly helped me in completing the task.

K.Dheeraj Kumar, B.Sc(Chemistry), III Year
 G.Laxmi kanth, B.Sc(Chemistry), III Year
 K.Narasimha, B.Sc(Chemistry), III Year
 G.Omkar, B.Sc(Chemistry), III Year
 D.Anil, B.Sc(Chemistry), III Year
 A.Jeremiah, B.Sc(Chemistry), III Year

<u>DECLARATION</u>

We hereby declare that the project report entitled "ADSORPTION OF Fe[III] ON CELLULOSE USING D-VALINE AS FACILITATING AGENT" is the work done by us in the campus at Department of Chemistry, Tara Government College, Sangareddy(A)during the academic year 2021-2022 and is submitted in partial fulfillment of the requirements for the degree of Bachelor of Science by Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

S.No.	Name of the Student	Roll Number	Group	Year	Signature
1	K.Dheeraj Kumar	605819578017	BSc Mccs	III	
2	G.Laxmi kanth	605819578012	BSc Mccs	III	
3	K.Narasimha	605819578016	BSc Mccs	III	
4	G.Omkar	605819578014	BSc Mccs	III	
5	D.Anil	605819578009	BSc Mccs	III	
6	A.Jeremiah	605819578001	BSc Mccs	III	

TARA GOVERNMENT COLLEGE, SANGAREDDY – 502 001 (AUTONOMOUS)

BONAFIDE CERTIFICATE

Certified that the project report "ADSORPTION OF Fe[III] ON

CELLULOSE USING D-VALINE AS FACILITATING AGENT" is the

bonafidework of

1.K.Dheeraj Kumar,	B.Sc(Chemistry),	III	Year
2.G.Laxmi kanth,	B.Sc(Chemistry),	III	Year
3.K.Narasimha,	B.Sc(Chemistry),	III	Year
4.G.Omkar,	B.Sc(Chemistry),	III	Year
5.D.Anil,	B.Sc(Chemistry),	III	Year
6.A.Jeremiah,	B.Sc(Chemistry),	III	Year

who carried out the project work under my supervision.

Dr.K.ABHIJIT **PROJECT SUPERVISOR** Smt. M.PRAVEENA PRINCIPAL

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Chapter-I

INTRODUCTION

ADSORPTION

Adsorption is a process in which solids come into contact with liquids or gases, and the mass transfer occurs from liquids to solids. Desorption is the reversal of this action. Adsorption operations take advantage of a solid's capacity to concentrate certain chemicals from a fluid on to its surface. Adsorbate refers to the adsorbed substance, while adsorbent refers to the solid substance. The following are some examples of solid-liquid and solid-gas applications:

- Removing dissolved moisture from gasoline.
- Decolorization of sugar solutions and petroleum products.
- Removing noxious odors and tastes from water. Dehumidification of air and gases is one of the solid-gas activities.
- To fractionate mixtures of hydrocarbon gases such as methane, ethane, and propane.
- To remove undesirable odors and contaminants from gases.
- To recover valuable solvent vapors from dilute gas mixtures.

NATURE OF ADSORBENTS

Adsorbents are typically granular in nature, ranging in size from 0.5 mm to 12 mm. They can't have a lot of pressure decrease or get swept away by a fast-moving stream. During handling, they must maintain their shape and size. They'll need a lot of pores and a lot of surface area per unit mass.

Some of the commonly used adsorbents, their sources and applications are given below:

S. No.	Adsorbent	Source	Application
1.	Fuller's earth	Naturally occurring clay is heated and dried to get a porous structure.	De-colorizing, drying of lubricating oils, kerosene and engine oils.
2.	Activated charcoal	Bentonite or other activated clay which are activated by treatment with sulfuric acid and further washing, drying and crushing.	Used for de-colorizing petroleum products.
3.	Bauxite	A naturally occurring hydrated alumina, activated by heating at 230-815	Used for de-colorizing petroleum products and for drying gases.

4.	Alumina	A hard hydrated aluminium oxide, which is activated by heating to drive off the moisture and then crushed to desired size.	Used as desiccant.
5.	Bone-char	Obtained by destructive distillation of crushed bones at 600-900	Used for refining sugar and can be reused after washing and burning.
6.	Silica gel	A hard granular and porous product obtained from sodium silicate solution after treatment with acid. Normally has 4 to 7% water in the product.	Dehydration of gases and liquids, and separation of gas-liquid hydrocarbon mixture.

7.	Activated carbon	 (1) Vegetable matter is mixed with calcium chloride, carbonized and finally the inorganic compounds are leached away. (2) Organic matter is mixed with porous pumice stones and then heated and carbonized to deposit the carbonaceous matter throughout the porous particle. (3) Carbonizing substances like wood, sawdust, coconut shells, fruit pits, coal, lignite and subsequent activation with hot air steam. It is available in granular or pellated form. 	De-colorizing of sugar solutions, chemicals, drugs, water purification, refining of vegetable and animal oils, recovery of gold and silver from cyanide ore-leach solution, recovery of solvent vapour from gas-mixtures, collection of gasoline hydro-carbons from natural gas, fractionation of hydrocarbon gases.
8.	Molecular sieves	These are porous synthetic zeolite crystals, metal alumino-silicates.	Dehydration of gases and liquids, and separation of gas- liquid hydrocarbon mixture.

SIGNIFICANCE OF RESEARCH PROBLEM

Cellulose can be used for adsorption of heavy metals owing to its polyhydric functional groups which will be a fetching tool to develop eco-friendly remediation strategy. Mean while, there is a need to improve the adsorption efficacy of cellulose by the incorporation of facilitating agents to design and develop standardized heavy metal remediation technology. For this purpose organic ligands will be useful to bring the metal-adsorbent linkages by stabilizing the metal complexes which will bind to the adsorbent more covalently.

OBJECTIVE OF THE PROJECT

To evaluate the impact of D-Valine as facilitating agent in the adsorption of Fe (III) from aqueous solution by Cellulose as an adsorbent to develop efficient remediation technology using concept of Coordination chemistry.

Chapter-II

ADSORPTION OF HEAVY METALS FROM DILUTE SOLUTIONS

The phrase "heavy metals" refers to a collection of metals and metalloids that have an atomic density more than 6 g cm—3. Although it is a broad phrase, it is typically used to refer to elements like Cd (cadmium), Cr (chromium), Cu (copper), Hg (mercury), Ni (nickel), Pb (lead), and Zn (zinc) that are commonly related with pollution and toxicity problems. Heavy metals, unlike most organic contaminants, occur naturally.

Each of these elements has a range of normal background concentrations in soils, sediments, waterways, and living creatures because they occur naturally in rock-forming and ore minerals. Significant amounts of various heavy metals are produced each year from the mining of their respective ores. Approximately 14,500 103 tonnes of copper were produced in 2004. (US Geological Survey, 2004). Soils, sediments, waterways, and live species all play a role in the environment. Significant amounts of various heavy metals are produced each year from the mining of their respective ores.

Metals used in industry and other domestic processes (e.g., burning fossil fuels, incineration of wastes, automobile exhausts, smelting processes, and the use of sewage sludge as a landfill material and fertiliser) have released large amounts of potentially toxic heavy metals into the atmosphere, as well as into aquatic and terrestrial environments. Cd, Pb, and Zn are common hazardous metals released into the environment.Cu, Ni, Cr, Co, Zn, and Pb are some of the most common metals (Babich et al., 1985). Table 1 shows the industrial sources of a variety of metals emitted, as well as the potential for contamination.

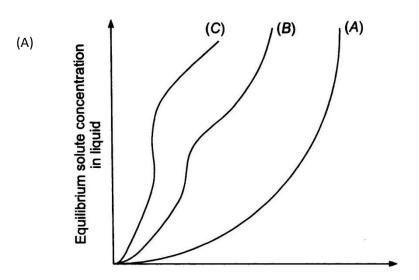
While many heavy metals are required by plants as micronutrients, larger quantities have been linked to a variety of harmful effects. Lead causes encephalopathy, cognitive impairment, behavioural problems, renal damage, anaemia, and reproductive system toxicity at high exposure levels (Pagliuca and Mufti, 1990). In its hexavalent state, chromium is well acknowledged to be hazardous (Rowbotham et al., 2000). Exposure to Cr (VI) compounds has been linked to a greater incidence of respiratory malignancies in humans (IARC, 1990). Cadmium has been linked to nephrotoxic effects, especially at high levels of exposure; long-term exposure may also induce bone dam- age (Friberg, 1985). Neurobehavioural problems and developmental abnormalities, such as dyslexia, attention deficit hyperactivity disorder, and intellectual retardation, can be caused by high mercury levels (Weiss and Landrigan, 2000). Copper poisoning can cause weakness, tiredness, and anorexia, as well as damage to the stomach and intestines (Theophanides and Anastassopoulou, 2002). Nordberg et al. go into great length about the toxicity of nickel and other heavy metals (2007)

Industry	Metals	Pollution Arising	Reference
Metalliferous mining	Cd,Cu,Ni,Cr,Co,Zn	Acid mine drainage,	Babich et al.(1985)
		tailings, slag heaps	Aswathanarayana(2003)
Agriculture materials	Cd,Cr,Mo,Pb,U,V,Zn	Run-off, surface and	Nicholson et al.(2003)
Fertilisers		groundwater	Otero et al.(2005)
		contamination, plant	
		bioaccumulation	
Manures sewage	Zn,Cu,Ni,Pb,Cd,Cr,As,Hg	Landspreading threat to	Nicholson et al.(2003)
sludge		ground and surface water	Cheung and Wong
			(1983)
			Walter et al.(2006)

 Table : Significant anthropogenic sources of metals in the environment.

Metallurgical	Pb,Mo,Ni.Cu,Cd,As,Te,U,	Manufacture, disposaland	Alloway and Ayres
industries Specialist	Zn	recycling of metals,	(1993)
alloys and steels		Tailings and slag heaps	Rule et al (2006)
			Cheng (2003)
Waste disposal	Zn,Cu,Cd,Pb,Ni,Cr,Hg	Landfill leachate,	Kjeldson et al.(2002)
Landfill leachate		contamination of ground	Fernandez et al.(2005)
		and surface	
Electronics	Pb,Cd,Hg,Pt,Au,Cr,As,Ni,	Aqueous and solid	Veglio et al. (2003)
	Mn	metallic waste food	
		manufacturing and	
		recycling process	
Metal finishing	Cr,Ni,Zn,Cu	Liquid effluents from	Castelblanque and
industry		plating processes	salimbeni(2004)
Electroplating			Zhao et al.(1999)
			Alvarez-Ayuso et
			al.(2003)
Miscellaneous	Pb,Sb,Zn,Cd,Ni,Hg	Waste battery fluid,	EU Directorate general
sources Batteries		contamination of soil and	of the Environment
		groundwater	(2004)
Paints and pigments	Pb,Cr,As,Ti,Ba,Zn	Aqueous waste from	Davis and Burns (1999)
		manufacture, old paint	Barnes and Davis (1996)
		deterioration and soil	Monken(2000)
		pollution	

Both the solvent and the solute are adsorbed whenever a mixture of solute and solvent is adsorbed using an adsorbent. As a result, only relative or apparent solute adsorption can be determined. As a result, treating a known volume of solution of original concentration C with a known weight of adsorbent is standard procedure. Let C* be the solution's final equilibrium solute concentration. If v is the volume of solution per unit mass of adsorbent (cc/g), and C and C* are the starting and equilibrium concentrations (g/cc) of the solute, then the apparent adsorption of the solute per unit mass of adsorbent is v(C -C*), (g/g), neglecting any volume change. This statement is most useful in the case of dilute solutions. The C* value is determined by the temperature, nature, and properties of the adsorbent when the proportion of the original solvent that can be adsorbed is tiny. The Freundlich adsorption isotherm, $C^*=K[v(C - C^*)]n$, represents the adsorption phenomena in dilute fluids across a small concentration range. The Freundlich adsorption equation is especially useful in situations where the identification of the solute is unknown, such as the removal of colouring substances from sugar solutions, oils, and other liquids. A spectrophotometer or colorimeter can quickly determine the colour composition of the solute. In worked example 2, the interpretation of this data is demonstrated. Adsorption is good if the value of n is high, say 2 to 10. If it's between 1 and 2, it's relatively challenging, and if it's less than 1, it's easy and it indicates poor adsorption characteristics. Freundlich adsorption equation is also useful in such a case where the actual identification of solute is not known, e.g. removal of colouring substance from sugar solutions, oils etc. A typical adsorption isothermal for the adsorption of various adsorbents A, B and C in dilute solution at the same temperature for the same adsorbent is represented in a graph.



Kg solute apparently adsorbed / kg adsorbent

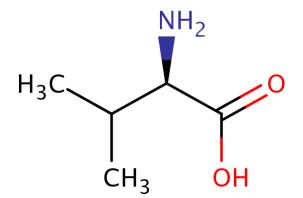
METAL COMPLEXES OF AMINO ACIDS

Transition metal amino acid complexes are a large family of coordination complexes containing the conjugate bases of the amino acids, the 2-aminocarboxylates. Amino acids are prevalent in nature, and all of them function as ligands toward the transition metals. Not included in this article are complexes of the amides (including peptide) and ester derivatives of amino acids. Also excluded are the polyamino acids including the chelating agents EDTA and NTA.

Most commonly, amino acids coordinate to metal ions as N,O bidentate ligands, utilizing the amino group and the carboxylate. They are "L-X" ligands. A five-membered chelate ring is formed. The chelate ring is only slightly ruffled at the sp³-hybridized carbon and nitrogen centers. For those amino acids containing coordinating substituents, the resulting complexes are more structurally diverse since these substituents can coordinate. Histidine, aspartic acid, methionine, and cysteine sometimes form

tridentate N,N,O, N,O,O, S,N,O, and S,N,O complexes, respectively. Using kinetically inert metal ions, complexes containing monodentate amino acids have been characterized. These complexes exist in either the N or the O linkage isomers. It can be assumed that such monodentate complexes exist transiently for many kinetically labile metal ions (e.g. Zn^{2+}). Mixing simple metal salts with solutions of amino acids near neutral or elevated pH often affords bis- or tris complexes. For metal ions that prefer octahedral coordination, these complexes often adopt the stoichiometry M(aa)₃ (aa = amino carboxylate, such as glycinate, H₂NCH₂CO₂⁻). Complexes of the 3:1 stoichiometry have the formula is [M(O₂CC(R)HNH₂)₃]^{*r*}. Such complexes adopt octahedral coordination geometry. These complexes can exist in facial and meridional isomers, both of which are chiral. The stereochemical possibilities increase when the amino acid ligands are not homochiral. Complexes with the 2:1 stoichiometry are illustrated by copper(II) glycinate [Cu(O₂CC(R)HNH₂)₂], which exists both in anhydrous and pentacoordinate geometries. When the metal is square planar, these complexes can exist as cis and trans isomers. The stereochemical possibilities increase when the amino acid ligands are not homochiral is tridentate amino acid are not homochiral. Homoleptic complexes are also known where the amino carboxylate is tridentate amino acids. One such complex is Ni(κ^3 -histidinate)₂.

D-Valine can easily involved in M-L complex formation due to its functional flexibility. Moreover its water solubility encourages the metal-ligand complex formation under aqueous conditions.



D-Valine

CELLULOSE AS ADSORBENT

Cellulose is the most plentiful and renewable polymer material available on the planet. According to estimates, deterioration and soil pollution Photosynthesis produces 10^{11} – 10^{12} tonnes of cellulose every year in a relatively pure form, such as in the seed hairs of cotton plants, but it is more commonly mixed with lignin and other polysaccharides (so-called hemicelluloses) in the cell walls of woody plants (Klemm et al., 2002). For thousands of years, cellulose has been utilised as an energy source, a building material, and a clothing material in the form of wood and cotton.

As a carbohydrate polymer, cellulose has a molecular structure that is made up of repeating b-Dglucopyranose units that are covalently linked by acetal functionalities between the OH groups of the C4 and C1 carbon atoms (b-1,4-glucan). Cellulose is a long, linear-chain polymer with a lot of hydroxyl groups (three per anhydroglucose (AGU) unit) and the 4C1 conformation. Every second AGU unit is rotated 180 degrees in the plane to accommodate the preferred bond angles. The number of constituent AGU units (degree of polymerization, DP) determines the length of the polymeric cellulose chain, which varies depending on the origin and treatment of the cellulose raw material (Klemm et al., 2002).

Cellulose has a ribbon form that allows it to twist and bend in directions other than the plane, making it a fairly flexible molecule. Due to the presence of hydroxyl (–OH) groups that protrude from the chain and create intermolecular hydrogen bonds, there is a relatively strong interaction between neighbouring cellulose molecules in dry fibres. Each chain of cellulose regenerated fibres contains 250– 500 repeating units (Klemm et al., 2002). The hydrophilicity, chirality, and degradability features of cellulose are due to this molecular structure. Chemical reactivity is mostly determined by the OH groups' high donor reactivity.

Chapter-III

ATOMIC ABSORPTION SPECTROMETRY

Atomic absorption spectrometry (AAS) is an analytical technique that measures the concentrations of elements. Atomic absorption is so sensitive that it can measure down parts per billion of a gram (µg dm⁻³) in a sample. The technique makes use of the wavelengths of light specifically absorbed by an element. They correspond to the energies needed to promote electrons from one energy level to another, higher, energy level.

Atomic absorption spectrometry has many uses in different areas of chemistry.

Clinical analysis: Analysing metals in biological fluids such as blood and urine.

Environmental analysis: Monitoring our environment- eg finding out the levels of various elements in rivers, seawater, drinking water, air, petrol and drinks such as wine, beer and fruit drinks.

Pharmaceuticals: In some pharmaceutical manufacturing processes, minute quantities of a catalyst used in the process (usually a metal) are sometimes present in the final product. By using AAS the amount of catalyst present can be determined.

Industry: Many raw materials are examined and AAS is widely used to check that the major elements are present and that toxic impurities are lower than specified- eg in concrete, where calcium is a major constituent, the lead level should be low because it is toxic.

Mining: By using AAS the amount of metals such as gold in rocks can be determined to see whether it is worth mining the rocks to extract the gold.

HOW IT WORKS

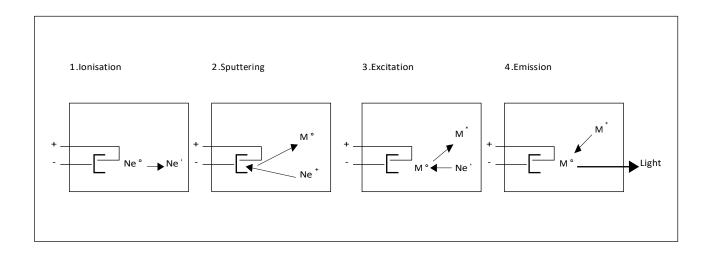
Atoms of different elements absorb characteristic wavelengths of light. Analysing a sample to see if it contains a particular element means using light from that element. For example with lead, a lamp containing lead emits light from excited lead atoms that produce the right mix of wavelengths to be absorbed by any lead atoms from the sample. In AAS, the sample is atomized- i.e. converted into ground state free atoms in the vapour state- and a beam of electromagnetic radiation emitted from excited lead atoms is passed through the vaporized sample. Some of the radiation is absorbed by the lead atoms in the sample. The greater the number of atoms there is in the vapour, the more radiation is absorbed. The amount of light absorbed is proportional to the number of lead atoms. A calibration curve is constructed by running several samples of known lead concentration under the same conditions as the unknown. The amount the standard absorbs is compared with the calibration curve and this enables the calculation of the lead concentration in the unknown sample. Consequently an atomic absorption spectrometer needs the following three components: a light source; a sample cell to produce gaseous atoms; and a means of measuring the specific light absorbed.

THE LIGHT SOURCE

The common source of light is a 'hollow cathode lamp'. This contains a tungsten anode and a cylindrical hollow cathode made of the element to be determined. These are sealed in a glass tube filled with an inert gas- e.g neon or argon- at a pressure of between 1 Nm⁻² and 5 Nm⁻².

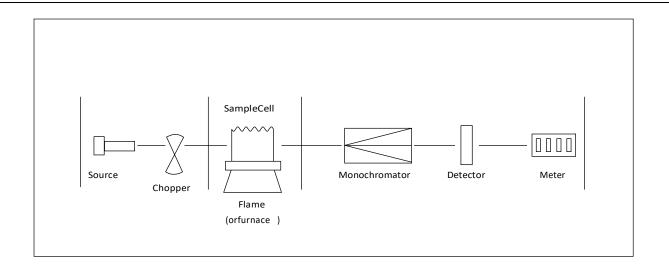


The ionization of some gas atoms occurs by applying a potential difference of about 300-400V between the anode and the cathode. These gaseous ions bombard the cathode and eject metal atoms from the cathode in a process called sputtering. Some sputtered atoms are in excited states and emit radiation characteristic of the metal as they fall back to the ground state $-egPb^* \rightarrow Pb + h v$. The shape of the cathode concentrates the radiation into a beam which passes through a quartz window, and the shape of the lamp is such that most of the sputtered atoms are redeposited on the cathode. A typical atomic absorption instrument holds several lamps each for a different element. The lamps are housed in a rotating turret so that the correct lamp can be quickly selected.



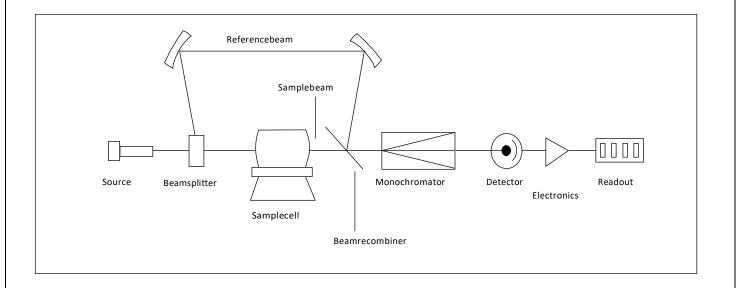
THE OPTICAL SYSTEM AND DETECTOR

A monochromator is used to select the specific wavelength of light –ie spectral line – which is absorbed by the sample, and to exclude other wavelengths. The selection of the specific light allows the determination of the selected element in the presence of others. The light selected by the monochromator is directed onto a detector that is typically a photomultiplier tube. This produces an electrical signal proportional to the light intensity



DOUBLE BEAM SPECTROMETERS

Modern spectrometers incorporate a beam splitter so that one part of the beam passes through the sample cell and the other is the reference. The intensity of the light source may not stay constant during an analysis. If only a single beam is used to pass through the atom cell, a blank reading containing no analyte (substance to be analysed) would have to be taken first, setting the absorbance at zero. If the intensity of the source changes by the time the sample is put in place, the measurement will be inaccurate. In the double beam instrument there is a constant monitoring between the reference beam and the light source. To ensure that the spectrum does not suffer from loss of sensitivity, the beam splitter is designed so that as high a proportion as possible of the energy of the lamp beam passes through the sample.



ATOMISATION OF THE SAMPLE

Two systems are commonly used to produce atoms from the sample. Aspiration involves sucking asolution of the sample into a flame; and electrothermal atomisation is where a drop of sample is placed into a graphite tube that is then heated electrically.

Some instruments have both atomisation systems but share one set of lamps. Once the appropriate lamp has been selected, it is pointed towards one or other atomisation system.

FLAME ASPIRATION

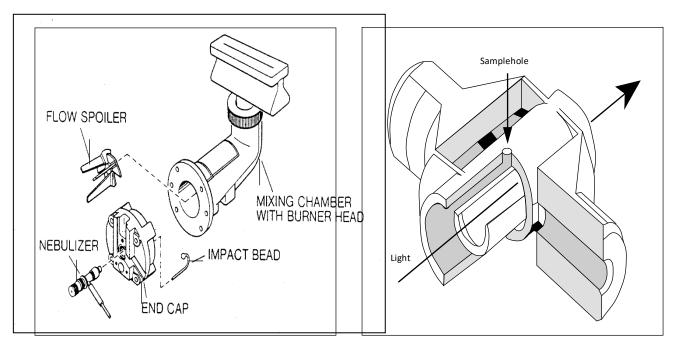




Figure 2

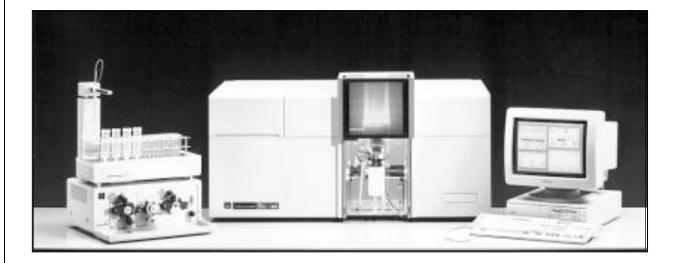
Ethyne/air (giving a flame with a temperature of 2200–2400°C) or ethyne/dinitrogen oxide (2600–2800°C) are often used. A flexible capillary tube connects the solution to the nebuliser. At the tip of the capillary, the solution is 'nebulised' –ie broken into small drops. The larger drops fall out and drain off while smaller ones vaporise in the flame. Only ca 1% of the sample is nebulised.

ELECTROTHERMAL ATOMIZATION

 $25 \ \mu$ l of sample (ca 1/100th of a raindrop) is placed through the sample hole and onto the platform from an automated micropipette and sample changer. The tube is heated electrically by passing a current through it in a pre-programmed series of steps. The details will vary with the sample but typically they might be 30–40 seconds at 150°C to evaporate the solvent, 30 seconds at 600°C to drive off any volatile organic material and char the sample to ash, and with a very fast heating rate (ca 1500 °C s⁻¹) to 2000-2500°C for 5–10 seconds to vaporise and atomise elements (including the element being analysed). Finally heating the tube to a still higher temperature –ca 2700°C – cleans it ready for the next sample. During this heating cycle the graphite tube is flushed with argon gas to prevent the tube burning away. In electrothermal atomisation almost 100% of the sample is atomised. This makes the technique much more sensitive than flame AAS.

SAMPLE PREPARATION

Sample preparation is often simple, and the chemical form of the element is usually unimportant. This is because atomisation converts the sample into free atoms irrespective of its initial state. The sample is weighed and made into a solution by suitable dilution. Elements in biological fluids such as urine and blood are often measured simply after a dilution of the original sample.



When making reference solutions of the element under analysis, for calibration, the chemical environment of the sample should be matched as closely as possible –i.e. the analyte should be in the same compound and the same solvent. Teflon containers may be used when analyzing very dilute aolutions because elements such as lead are sometimes leached out of glass vessels and can affect the results

BACKGROUND ABSORPTION

It is possible that other atoms or molecules apart from those of the element being determined will absorb or scatter some radiation from the light source. These species could include unvaporisedsolvent droplets, or compounds of the matrix (chemical species, such as anions, that tend to accompany the metals being analysed) that are not removed completely. This means that there is a background absorption as well as that of the sample.

One way of measuring and correcting this background absorption is to use two light sources, one of which is the hollow cathode lamp appropriate to the element being measured. The second light source is a deuterium lamp.

The deuterium lamp produces broad band radiation, not specific spectral lines as with a hollow cathode lamp. By alternating the measurements of the two light sources – generally at 50 -100 Hz – the total absorption (absorption due to analyte atoms plus background) is measured with the specific light from the hollow cathode lamp and the background absorption is measured with the light from the deuterium lamp. Subtracting the background from the total absorption gives the absorption arising from only analyte atoms.

CALIBRATION

A calibration curve is used to determine the unknown concentration of an element -eg lead – in a solution. The instrument is calibrated using several solutions of known concentrations. A calibration curve is produced which is continually rescaled as more concentrated solutions are used – the more concentrated solutions absorb more radiation up to a certain absorbance. The calibration curve shows the concentration

against the amount of radiation absorbed in the given figure.(a) The sample solution is fed into the instrument and the unknown concentration of the element-e.g. lead- is then displayed on the calibration curve given in the below figure.(b)

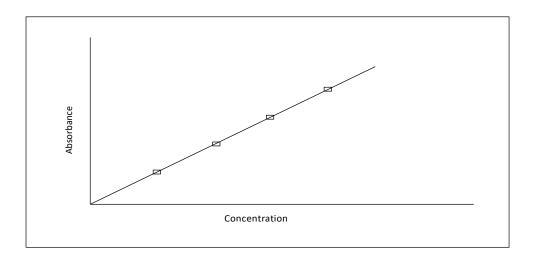


Figure (a)

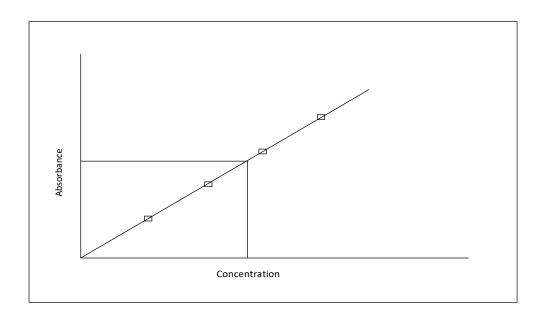


Figure (b)

Chapter-IV

METHODOLOGY

MATERIALS REQUIRED

- Ferric chloride pure Hexahydrate
- D-Valine
- Cellulose
- Volumetric flask
- Digital Weighing Machine
- Watch glasses
- Hot air oven
- Ultra pure water (Demineralized)
- 2% of Nitric acid

PROCEDURE

- Using a 250ml Volumetric flask prepare a metal ligand solution by adding 100ppm of Ferric chloride Hexahydrate (MOLYCHEM MCR-11580) and 200ppm of D-Valine (Avra N2200133).
 Prepare 250ml solution by adding Ultra pure water and then keep this system aside for a few hours.
- Weigh 5 grams of Cellulose (Avra N2001901) using a Digital weighing machine (Citizen Scales(I) PVT LTD CTG302-300) and take this into a beaker.
- **3.** Now add 100ml of the above prepared metal ligand solution into the beaker and stir the mixture well for 10 minutes using a glass rod.

- **4.** Keep this mixture aside for 48 hours without disturbing it as at this step Ferric is going to be adsorbed on Cellulose in the presence of D-Valine which acts as a chelating agent.
- **5.** After completion of 48 hours take the mixture and filter it off using Whatman Grade 1 filter paper and a funnel.
- **6.** After filtration of the mixture again add Ultra pure water for 3 times and then filtrate it to obtain pure concentration of Ferric which is get adsorbed on Cellulose.
- Collect the filtered Cellulose powder and place it on a watch glass and keep this in a Hot air owen at 60 °C for 10 hours to get rid off moisture present in it.
- 8. Now weigh each 1 gm of Cellulose in glass vials.
- 9. Now take a beaker and rinse it with ultra pure water then followed by Nitric acid.
- **10.** Take 0.5 grams of Cellulose sample in the beaker and add 2% of Nitric acid and stir the mixture well for 10-15 minutes.
- **11.** Filter the mixture using Whatman Grade 1 filter paper and again 3 times by using Ultra pure water to obtain pure concentration of Ferric present in the mixture prepared using the sample.
- Take this collected sample solution and keep this system under AAS (Thermo Scientific iCE 3300)
- **13.** Calculate the concentration of Ferric adsorbed on Cellulose at different ppm levels. Observe the graph obtained and note down the readings of the result we obtained.
- 14. Same Experiment carried out without the interference/addition of Ligand i.e. D-Valine for Control Experiment.

AA spectrometer

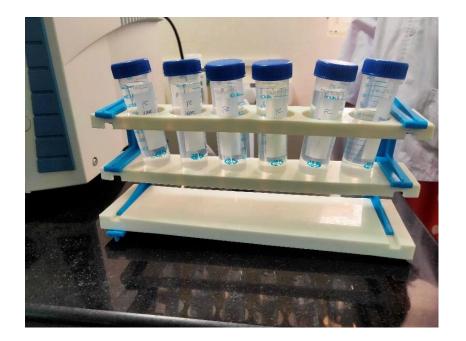


Model no. ICE 33 0

Fume hood



Fe standards with test sample





Sample is Taking in to the Machine

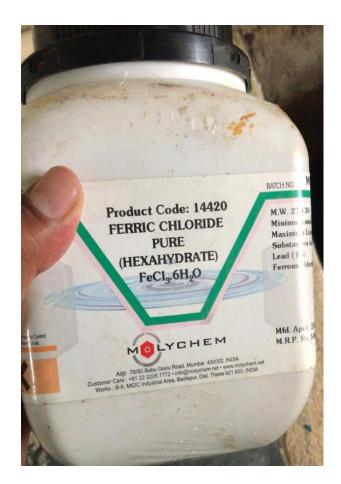
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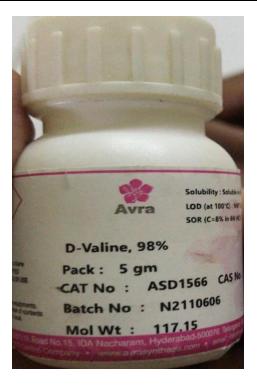
Burning of test sample in AASpectometer

Values after intake of test samples

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Ferric Chloride (FeCl₃6H₂O)



D-VALINE (Avra)



Solution of Fe(III)-D-Valine Solution

Other Apparatus Used



Watch glass



Spatula



Measuring Jar 50 ml



Digital weighing Machine







Filtering the solution



China Dish











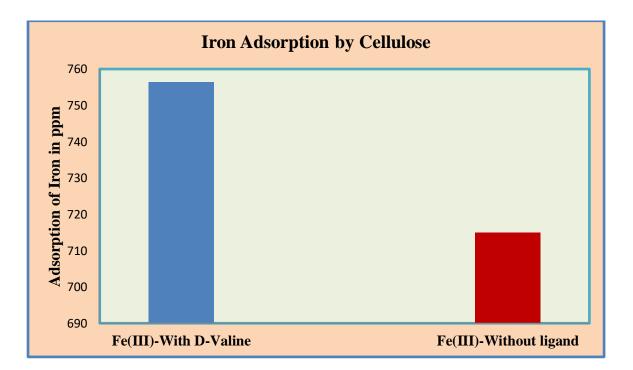
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Results obtained on the monitor under AAS Method

Chapter-V

RESULTS AND DISCUSSIONS

From an aqueous solution of the metal ligand solution Fe(III)-D-Valine, Cellulose absorbed **756.5**ppm of iron. When D-Valine is absent, Cellulose only absorbed **715.0**ppm. The AAS data clearly show that the addition of ligands improved the metal adsorption by triggering possible chemical interactions between the adsorbent and adsorbate. Strong coordination bonds between D-Valine and Fe(III) result in the formation of a soluble complex in aqueous solution. By creating chemical bonds, the ligand-capped Fe(III) complex firmly adhered to Cellulose. The AAS data clearly show that **5.804**% more adsorption occurred when D-Valine was included as a chelating agent.



Impact of D-Valine on Adsorption of Fe (III) ions from aqueous solution by Cellulose.

Spectrometer Parameters – Fe:

Element : Fe	Measurement mode : Absorbance	
Wavelength : 248.3nm	Band pass : 0.2nm	Lamp current : 75%
Background correction : D2	High Resolution : Off	Optimise Spectrometer Parameters : No
Signal type : continuous	Resamples : Fast	Number of resamples : 3
Measurement time : 4.0secs	Flier mode : No	
Use RSD Test : No		

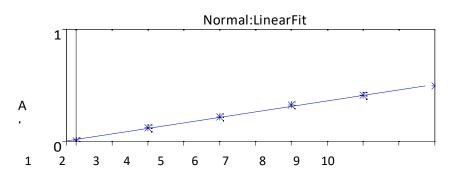
Flame Parameters – Fe:

Flame type : Air – C2H2	Fuel Flow : 0.9L/min	Auxiliary Oxidant : Off
Nebuliser Uptake : 4secs	Bunsen Stabilisation : Omins	Optimise Fuel Flow : No
Burner Height :7.0mm	Optimise Burner fuel : No	

Calibration Parameters-Fe:

Calibration mode : Normal	Line fit : Linear	Use stored calibration : No			
Concentration units : mg/L	Scales units : mg/L	Scaling factor : 1.0000			
Acceptable fit : 0.990	Rescale Limits : 10.0%	Failure Action : Flag and Continue			
Standard 1 - 2.0000	Standard 4 - 8.0000				
Standard 2 - 4.0000	Standard 5 - 10.0000				
Standard 3 - 6.0000					

Solutions Results-Fe:



Y = 0.04924x + 0.0141

Fit: 0.9965

Characteristic Conc.: 0.0894

Sample ID Signal		RSD	Conc.			
	Abs	%	Mg/L			
Fe Blank	0.001	35.6	0.0000			
1	0.001	Backg	round: -0.003			
2	0.001	Backg	round: -0.003			
3	0.000	Backg	round: -0.003			
Fe Standard 1	0.120	0.3	2.0000			
1	0.120	Background: 0.003				
2	0.119	Background: 0.003				
3	0.120	Background: 0.003				
Fe Standard 2	0.215	0.3 4.0000				
1	0.215	Background: 0.004				
2	0.215	Background:0.004				
3	0.216	Background:0.004				
Fe Standard 3	0.322	0.2	6.0000			

0.323	Background: 0.005					
0.322	Background: 0.005					
0.322	Background: 0.005					
0.411	0.3 8.0000					
0.410	Background: 0.005					
0.412	Background: 0.005					
0.412	Background: 0.005					
0.493	0.1 10.0000					
0.494	Background: 0.005					
0.494	Background: 0.005					
0.494	Background: 0.006					
0.759	0.1 15.1286					
0.758	Background: 0.003					
0.760	Background: 0.003					
0.758	Background: 0.004					
0.717	0.1 14.2986 C					
0.719	Background: 0.004					
0.716	Background: 0.003					
0.716	Background: 0.004					
	0.322 0.322 0.411 0.410 0.412 0.412 0.412 0.493 0.494 0.494 0.759 0.758 0.760 0.758 0.717 0.719 0.716					

Test Results:

S.no	Test Parameters	Sample	Results
01.	Iron by AAS Analysis:	Fe(III)-D-Valine –	756.5ppm
	(ppm)	Cellulose Sample	
02.	Iron by AAS Analysis:	Fe(III)– Cellulose	715.0 PPM
	(ppm)	(control)	

CONCLUSION

According to the results of the current experiment, D-Valine, which acts as a chelating agent, is crucial to the adsorption of Fe (III) ions from aqueous solution by Cellulose and enhances adsorption by up to **5.804**%. Designing the latest heavy metal remediation systems that employ natural chelating ligands as facilitation agents in metal adsorption processes would benefit from this feature.

REFERENCES

- 1 I. Nakagawa, R. J. Hooper, J. L. Walter and T. J. Lane, Spectrochim. Acta, 1966, 21, 1-14.
- 2 M. Tomassetti, E. Cardarelli, R. Curini and G. D'ascenzo, *Thermochim. Acta*, 1987, 113, 243–255.
- 3 M. M. Suvarna, V. S. Panse, U. P. Fulwadva and P. K. Jadhav, *Asian J. Chem.*, 1995, 7(3), 459–499.
- 4 J. G. Shao, Y. X. Yang, B. W. Li, L. P. Zhang, Y. R. Chen and X. L. Liu, *J. Therm. Anal. Calorim.*, 2009, 96(1), 277–285.
- 5 S. Shirazian and S. Gronert, J. Mol. Struct.: THEOCHEM, 1997, 397, 107-112.
- 6 T. Dudev and C. Lim, Annu. Rev. Biophys., 2008, 39, 97-116.7 J. J. R. Frausto da Silva and R. J.
- 7 P. Williams, The biological chemistry of the elements, Claredon Press, Oxford, 1991.
- 8. *Handbook on Metalloproteins*, ed. I. Bertini, A. Sigel and H.Sigel, Marcel Dekker, New York, 2001.
- 9. T. Dudev and C. Lim, Chem. Rev., 2003, 103, 773-787.

- 10D. W. Christianson and J. D. Cox, Annu. Rev. Biochem., 1999,68, 33-57.
- 11 P. M. Kroneck, V. Vortisch and P. Hemmerich, Eur. J. Biochem., 1980, 12, 603-612.
- 1212 B. M. Rode, Peptides, 1999, 20, 773-786.
- 13 B. M. Rode, J. Bujdak and A. H. Eder, Trends Inorg. Chem., 1993, 3, 45-62.
- 14 E. Constantino, A. Rimola, L. Rodriguez-Santiago and M. Sodupe, *New J. Chem.*, 2005, 29, 1585–1593.
- 15 W. F. DeGrado, C. M. Summa, V. Pavone, F. Nastri and A. Lombardi, Annu. Rev. Biochem., 1999, 68, 779–819.
- 16 Y. Lu, S. M. Berry and T. D. P ster, *Chem. Rev.*, 2001, 101, 3047–3080.
- 17 Y. Lu, Curr. Opin. Chem. Biol., 2005, 9, 118-126.
- 18 J. Xie, W. Liu and P. G. Schultz, Angew. Chem., Int. Ed., 2007, 46, 9239-9242.
- 19 A. L. Garay, A. Pichon and S. L. James, Chem. Soc. Rev., 2007, 36, 846-855.
- 20 X. Q. Xin and L. M. Zheng, J. Solid State Chem., 1993, 106, 451-456.
- 21 S. Mandal, G. Das and H. Askari, Struct. Chem., 2014, 25, 43-51.
- 22 H. Koshima, D. Matsushige, M. Miyauchi and J. Fujita, *Tetrahedron*, 2000, 56, 6845–6852.
- 23 T. Shimo, T. Uezono, T. Obata, M. Yasutake, T. Shinmyozuand K. Somekawa, *Tetrahedron*, 2002, 58, 6111–6116.
- 24 M. R. Wormald, A. J. Petrescu, Y. Pao, A. Glithero, T. Elliottand R. A. Dwek, *Chem. Rev.*, 2002, 102(2), 371–386.
- 25 N. Foloppe, B. Hartmann, L. Nilsson and A. D. MacKerell Jr, Biophys. J., 2002, 82, 1554–1569.
- 26 S. Miertus, E. Scrocco and J. Tomasi, Chem. Phys., 1981, 55(1), 117-129.
- 27 A. D. Becke, J. Chem. Phys., 1993, 98, 5648-5652.
- 28 C. Lee, W. Yang and R. G. Parr, Phys. Rev. B: Condens. Matter Mater. Phys., 1988, 37, 785-789.

- 29 M. J. Frisch, et al. Gaussian 03, Revision D.01, Gaussian Inc, Wallingford, CT, 2004.
- 30 M. J. Frisch, et al. Gaussian 09, Revision D02, Gaussian, Wallingford, CT, 2009.
- 31 I. R. Gould and I. H. Hillier, J. Chem. Soc., Chem. Commun., 1993, 951–952.
- 32 M. P. Andersson and P. Uvdal, J. Phys. Chem. A, 2005, 109, 2937–2941.
- 33 J. B. Foresman and A. Frisch, *Exploring chemistry with electronic structure methods*, 2nd edn, Gaussian, Inc., Pittsburgh, PA, 1996.
- 34 F. Freeman and K. T. Le, J. Phys. Chem. A, 2003, 107, 2908-2918.
- 35 R. Bauernschmitt and R. Ahlrichs, Chem. Phys. Lett., 1996, 256, 454-464.
- 36 M. E. Casida, C. Jamorski, K. C. Casida and D. R. J. Salahub, J. Chem. Phys., 1998, 108, 4439– 4449.
- 37 M. A. L. Marques and E. K. U. Gross, Annu. Rev. Phys. Chem., 2004, 55, 427-455.
- 38 E. Runge and E. K. U. Gross, *Phys. Rev. Lett.*, 1984, 52, 997–1000.
- 39 R. E. Stratmann, G. E. Scuseria and M. J. Frisch, J. Chem. Phys., 1998, 109, 8218-8224.
- 40 K. Morokuma, J. Chem. Phys., 1971, 55, 1236–1244.
- 41 F. Freeman and K. T. Le, J. Phys. Chem. A, 2003, 107, 2908–2918.
- 42 R. Bauernschmitt and R. Ahlrichs, Chem. Phys. Lett., 1996, 256, 454-464.
- 43 M. E. Casida, C. Jamorski, K. C. Casida and D. R. J. Salahub, *J. Chem. Phys.*, 1998, 108, 4439–4449.
- 44 M. A. L. Marques and E. K. U. Gross, Annu. Rev. Phys. Chem., 2004, 55, 427-455.
- 45 E. Runge and E. K. U. Gross, *Phys. Rev. Lett.*, 1984, 52, 997–1000.
- 46 R. E. Stratmann, G. E. Scuseria and M. J. Frisch, J. Chem. Phys., 1998, 109, 8218-8224.
- 44 S. A. Zabin and C. R. Jejurkar, Asian J. Chem., 1995, 7(3), 542–550.
- 45 S. Kapoor and M. S. Sastry, Proc. Indian Acad. Sci., Chem. Sci., 2000, 112, 459-463.
- 46 O. Zulbiye, Z. Huseyin and S. Mehmet, Turk. J. Chem., 2011, 35, 905–914.
- 47 S. V. Sanap and R. M. Patil, Res. J. Pharm. Sci., 2013, 2(1), 1-10.

43 | Page

- 48 K. Nakamoto, Y. Morimoto and A. E. Martell, J. Am. Chem. Soc., 1961, 83, 4528-4532.
- 49 P. R. Reddy, M. Radhika and P. Manjula, J. Chem. Sci., 2005, 117(3), 239–246.
- 50 M. M. Mashaly, Z. H. Abd-Elwahab and A. A. Faheim, J. Chin. Chem. Soc., 2004, 51, 901–915.
- 51 S. A. Zabin and C. R. Jejurkar, Asian J. Chem., 1995, 7(3), 542-550.
- 52 S. Kapoor and M. S. Sastry, Proc. Indian Acad. Sci., Chem. Sci., 2000, 112, 459-463.
- 53 O. Zulbiye, Z. Huseyin and S. Mehmet, Turk. J. Chem., 2011, 35, 905–914.
- 54 E. L. S. Gomes and O. L. Casagrande Jr, Thermochim. Acta, 1999, 331, 87-91.
- 55 J. M. Berg and D. L. Merkle, J. Am. Chem. Soc., 1989, 111,3759-3761.
- 56 N. C. Polfer, J. Oomens, D. T. Moore, G. von Helden, G. Meijer and R. C. Dunbar, J. *Am. Chem. Soc.*, 2006, 128,517–525.
- S. G. Stepanian, I. D. Reva, E. D. Radchenko, M. T. S. Rosado, M. L. T. S. Duarte, R. Fausto and
 L. Adamowicz, J. Phys. Chem. A, 1998, 102, 1041–1054.
- 58 K. Fukui, T. Yonezawa and H. Shingu, J. Chem. Phys., 1952, 20, 722-725.
- 59 M. T. Baei and S. Z. Sayyed-Alangi, Eur. J. Chem., 2012, 9, 1244–1250.
- 60. C. L. Brooks III, M. Karplus and B. M. Pettitt, *Proteins: a theoretical perspective of dynamics, structure, and thermodynamics*, Wiley, New York, 1988.
- 61 M. Remko, D. Fitz and B. M. Rode, J. Phys. Chem. A, 2008, 112, 7652-7661.
- 62 K. Wiberg, Tetrahedron, 1968, 24, 1093–1096.
- W. J. Hehre, L. Radom, P. v. R. Schleyer and J. A. Pople, *Abinitio molecular orbital theory*, Wiley, New York, 1986.
- 64 G. Das, J. Mol. Model., 2013, 19, 2981-2991.
- 65 S. Y. Lee and B. H. Boo, J. Phys. Chem., 1996, 100, 8782-8785.

66 K. Yokota, M. Hagimori, N. Mizuyama, Y. Nishimura, H. Fujito, Y. Shigemitsu and Y. **44** | P a g e

Tominaga, Beilstein J. Org. Chem., 2012, 8, 266–274.

- 67 Abdel-Aal, S.E., Gad, Y., Dessouki, A.M., 2006. The use of wood pulp and radiation modified starch in wastewater treatment. Journal of Applied Polymer Science 99, 2460–2469.
- 68 Abollino, O., Aceto, M., Malandrino, M., Sarzanini, C., Mentasti, E., 2003. Adsorption of heavy metals on Na-montmorillonite: effect of pH and organic substances. Water Research 37 (7), 1619–1627.
- 69. Aderhold, D., Williams, C.J., Edyvean, R.G.J., 1996. The removal of heavy-metal ions by seaweeds and their derivatives. Bioresource Technology 58 (1), 1–6.
- 70. Ajmal, M., Rao, R.A.K., Ahmad, R., Ahmad, J., 2000. Adsorption studies on Citrus reticulata (fruit peel of orange): removal and recovery of Ni(II) from electroplating wastewater. Journal of Hazardous Materials 79 (1–2), 117–131.
- 71. Altin, O., Ozbelge, H.O., Dogu, T., 1998. Use of general purpose adsorption isotherms for heavy metal–clay mineral interactions. Journal of Colloid Interface Science 198, 130–140.
- 72.Alloway, B.J., Ayres, D.C., 1993. Chemical Principles of Environmental Pollution, First ed. Chapman & Hall, London, pp. 190–216.
- 73.Alvarez-Ayuso, E., Garcia-Sanchez, A., Querol, X., 2003. Purification of metal electroplating waste waters using zeolites. Water Research 37 (20), 4855–4862.
- 74. Andini, S., Cioffi, R., Montagnaro, F., Pisciotta, F., Santoro, L., 2006. Simultaneous adsorption of chlorophenol and heavy metal ions on organophilic bentonite. Applied Clay Science 31 (1–2), 126–133.

- 75. Aoki,N.,Fukushima,K.,Kurakata,H.,Sakamoto,M.,Furuhata,K.,1999a.6-Deoxy-6mercaptocelluloseB and its S-substituted derivatives as sorbents for metal ions. Reactive and Functional Polymers 42 (3), 223–233.
- 76. Aoki, Y., Tanaka, K., Sakamoto, M., Furuhata, K., 1999b. Sorption of metal ions by bead cellulose grafted with amidoximated polyacrylo- nitrile. Sen'i Gakkaishi 55 (12), 569–575.
- 77. Aswathanarayana, U., 2003. Mineral Resources Management and the Environment. Routledge, Netherlands, pp. 223–256.
- Babich, H., Devanas, M.A., Stotzky, G., 1985. The mediation of mutagenicity and clastogenicity of heavy metals by physicochemical factors. Environmental Research 37 (2), 253–286.
- Bao-Xiu, Z., Peng, W., Tong, Z., Chun-yun, C., Jing, S., 2006. Preparation and adsorption performance of a cellulosic adsorbent resin for copper(II). Journal of Applied Polymer Science 99, 2951–2956.
- Barnes, G.L., Davis, A.P., 1996. Dissolution of lead paint in aqueous solution. Journal of Environmental Engineering ASCE 122, 663–666.
- 81. Bayramoglu, G., Bektas, S., Arica, M.Y., 2003. Biosorption of heavymetal ions on immobilized white-rot fungus Trametes versicolor. Journal of Hazardous Materials 101 (3), 285–300.
- Benke, N., Takács, E., Wojnárovits, L., Borsa, J., 2007. Pre-irradiation grafting of cellulose and slightly carboxymethylated cellulose (CMC) fibres. Radiation Physics and Chemistry 76 (8–9), 1355–1359.
- 83. Berber-Mendoza, M.S., Leyva-Ramos, R., Alonso-Davila, P., Mendoza- Barron, J., Diaz-Flores, P.E., 2006. Effect of pH and temperature on the ion-exchange isotherm of Cd(II) and
 46 | P a g e

Pb(II) on clinoptilolite. Journal of Chemical Technology and Biotechnology 301, 40–45.

- 84. Bhattacharya, A., Misra, B.N., 2004. Grafting: a versatile means to modify polymers: techniques, factors and applications. Progress in Polymer Science 29 (8), 767–814.
- 85. Bhattacharya, A., 2000. Radiation and industrial polymers. Progress in Polymer Science 25 (3), 371–401.
- 86. Bicak, N., Sherrington, D.C., Senkal, B.F., 1999. Graft copolymer of acrylamide onto cellulose as mercury selective sorbent. Reactive and Functional Polymers 41, 69–76.
- 87. Bicak, N., Senkel, S.F., 1998. Sorbitol-modified poly(*N*-glycidyl styrene sulfonamide) for removal of boron. Journal of Applied Polymer Science 68, 2113–2119.
- 88. Brown, P., Atly Jefcoat, I., Parrish, D., Gill, S., Graham, E., 2000. Evaluation of the adsorptive capacity of peanut hull pellets for heavy metals in solution. Advances in Environmental Research 4 (1), 19–29.
- 89. Castelblanque, J., Salimbeni, F., 2004. NF and RO membranes for the recovery and reuse of water and concentrated metallic salts from waste water produced in the electroplating process. Desalination 167, 65–73.
- 90.Cheng, S.P., 2003. Heavy metal pollution in China: origin, pattern and control. Environmental Science and Pollution Research 10 (3), 192–198.
- 91.Cheung, Y.H., Wong, M.H., 1983. Utilisation of animal manures and sewage sludges for growing vegetables. Agricultural Wastes 5 (2), 63–81.

- 92.Cochrane, E.L., Lu, S., Gibb, S.W., Villaescusa, I., 2006. A comparison of low cost biosorbents and commercial sorbents for the removal of copper from aqueous media. Journal of Hazardous Materials 137 (1), 198–206.
- 93. Crini, G., 2005. Recent developments in polysaccharide-based materials used as adsorbents in wastewater treatment. Progress in Polymer Science 30 (1), 38–70.
- 94.Dal Bosco, S.M., Jimenez, R.S., Carvalho, W.A., 2005. Removal of toxic metals from wastewater by Brazilian natural scolecite. Journal of Colloid and Interface Science 281 (2), 424–431.
- 95.Davis, A.P., Burns, M., 1999. Evaluation of lead concentration in runoff from painted structures. Water Research 33 (13), 2949–2958.
- 96. Demirbas, E., Kobya, M., Oncel, S., Sencan, S., 2002. Removal of Ni(II) from aqueous solution by adsorption onto hazelnut shell activated carbon: equilibrium studies. Bioresource Technology 84 (3), 291–293.
- 97.Dilek, F.B., Erbay, A., Yetis, U., 2002. Ni(II) biosorption by polyporous versicolor. Process Biochemistry 37 (7), 723–726.
- 98. Dimitrova, S.V., 1996. Metal sorption on blast-furnace slag. Water Research 30 (1), 228–232.
- 99. Feng, D., van Deventer, J.S.J., Aldrich, C., 2004. Removal of pollutants from acid mine wastewater using metallurgical by-product slags. Separation and Purification Technology 40 (1), 61–67.
- 100. Fernandez, Y., Maranon, E., Castrillon, L., Vazquez, I., 2005. Removal of Cd and Zn from inorganic industrial waste leachate by ion exchange. Journal of Hazardous Materials 126 (1–3), 169–175.

- 101. Freundlich, H., 1926. Adsorption. Journal of Physical Chemistry 7, 57-64.
- 102. Friberg, L.I., 1985. Rationale of biological monitoring of chemicals –with special reference to metals. American Industrial Hygiene Association Journal 46 (11), 633–642.
- 103.Gaballah, I., Kilbertus, G., 1998. Recovery of heavy metal ions through decontamination of synthetic solutions and industrial effluents using modified barks. Journal of Geochemical Exploration 62 (1–3), 241–286.
- 104. Gaey, M., Marchetti, V., Clement, A., Loubinoux, B., Gerardin, P., 2000. Decontamination of synthetic solutions containing heavy metals using chemically modified sawdusts bearing polyacrylic acid chains. Journal of Wood Science 46, 331–333.
- 105.Guclu, G., Gurdag, G., Ozgumus, S., 2003. Competitive removal of heavy metal ions by cellulose graft copolymers. Journal of Applied Polymer Science 90, 2034–2039.
- 106.Guibal, E., 2004. Interactions of metal ions with chitosan-based sorbents: a review. Separation and Purification Technology 38 (1), 43–74.

107. Gupta, K.C., Khandehar, K., 2002. Graft copolymerisation of acrylam- ide–methylacrylate comonomers onto cellulose using ceric ammonium nitrate. Journal of Applied Polymer Science 86, 2631–2642.

- 108. Gurdag, G., Yasar, M., Gurkaynak, M.A., 1997. Graft copolymerisation of acrylic acid on cellulose. Journal of Applied Polymer Science 66, 929–934.
- 109.Hancock, R.D., Martell, A.E., 1989. Ligand design for selective complex- ation of metal ions in aqueous solution. Chemical Reviews 89, 1875–1914.
- 110.Harris, J.A., Arthur, J.C., Carra, J.H., 1978. Photoinitiated polymerisa- tion of glycidyl methacrylate with cotton cellulose. Journal of Applied Polymer Science 22, 905–915.

- 111.Hashem, A., 2006. Amidoximated sunflower stalks (ASFS) as a new adsorbent for removal of Cu(II) from aqueous solution. Polymer– Plastics Technology and Engineering 45, 35–42.
- 112. Hilal, N., Al-Khatib, L., Atkin, B.P., Kochkodan, V., Potapchenko, N., 2003.
 Photochemical modification of membrane surfaces for (bio)foul- ing reduction: a nano-scale study using AFM. Desalination and the Environment: Fresh Water for All 158 (1–3), 65–72.
- 113.Hon, D.N.S., 1982. Graft Copolymerization of Lignocellulosic Fibers. American Chemical Society Symposium Series, 187, Washington DC, USA.
- 114. IARC, 1990. Monograph on the Evaluation of Carcinogenic Risk to Humans, vol.49, Chromium, Nickel and Welding. International Agency for Research on Cancer, Lyon.
- 115.Kamel, S., Hassan, E.M., El-Sakhawy, M., 2006. Preparation and application of acrylonitrilegrafted cyanoethyl cellulose for the removal of copper(II) ions. Journal of Applied Polymer Science 100, 329–334.
- 116. Langmuir, I., 1918. The adsorption of gases on plane surfaces of glass, mica and platinum.Journal of the American Chemical Society 40 (9), 1361–1402.
- 117. Lee, S.T., Mi, F.L., Shen, Y.J., Shyu, S.S., 2001. Equilibrium and kinetics studies of copper(II) ion uptake by chitosan-tripolyphosphate chelat- ing resin. Polymer 42, 1879–1892.
- 118. Theophanides, T., Anastassopoulou, J., 2002. Copper and carcinogenesis, critical reviews. Oncology/Haematology 42 (1), 57–64.
- 119.US Geological Survey, 2004. Minerals commodity summaries. Available from: http://minerals.usgs.gov./minerals/pubs/commodity>.

- 120. Vaughan, T., Seo, C.W., Marshall, W.E., 2001. Removal of selected metal ions from aqueous solution using modified corncobs. Bioresource Technology 78 (2), 133–139.
- 121.Veglio, F., Quaresima, R., Fornari, P., Ubaldini, S., 2003. Recovery of valuable metals from electronic and galvanic industrial wastes by leaching and electrowinning. Waste Management 23 (3), 245–252.
- 122.Villaescusa, I., Fiol, N., Martinez, M., Miralles, N., Poch, J., Serarols, J., 2004. Removal of copper and nickel ions from aqueous solutions by grape stalk wastes. Water Research 38 (4), 992–1002.
- 123.Walter, I., Martinez, F., Cala, V., 2006. Heavy metal speciation and phytotoxic effects of three representative sewage sludges for agricul- tural uses. Environmental Pollution 139 (3), 507–514.
- 124.Wang, L.K., Hung, N.K., Shammas, N.K., 2004. Physiochemical Treat- ment Processes, vol. 3. Human Press, New Jersey, pp. 103–138.
- 125.Wang, Y.H., Lin, S.H., Juang, R.S., 2003. Removal of heavy metal ions from aqueous solutions using various low-cost adsorbents. Journal of Hazardous Materials 102 (2–3), 291–302.
- 126.Waste Strategy, 2004. New energy goes into battery recycling. In: Environment for Europeans Magazine of the E.U. Directorate General of the Environment.
- 127.Weiss, B., Landrigan, P.J., 2000. The developing brain and the environ- ment: an introduction. Environmental Health Perspectives 108, 373–374.
- 128.Yavuz, O., Altunkaynak, Y., Guzel, F., 2003. Removal of copper, nickel, cobalt and manganese from aqueous solution by kaolinite. Water Research 37 (4), 948–952.

- 129.Zhao, M., Duncan, J.R., van Hille, R.P., 1999. Removal and recovery of zinc from solution and electroplating effluent using Azolla filiculoides. Water Research 33 (6), 1516–1522.
- 130.Zhou, D., Zhang, L., Zhou, J., Guo, S., 2004. Cellulose/Chitin beads for adsorption of heavy metals in aqueous solution. Water Research 38, 2643–2650.

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Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

General Parameters

Method : Iron (Fe) Autosampler : None Use SFI: No Operator : Jyothi

Instrument Mode: Flame *Dilution:* None

mA Hours

n/a

Report Date: 11-06-2022 17:40:20

Valid Method Signatures

11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Method Audit Trail

11-06-2022 17:22:18 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC

Signed with Reason : Approved by:

Analysis Details

Spectrometer: iCE 3000 AA01204906 v1.30

Analysis Name: Iron (Fe) 11-06-2022 Operator Name: Jyothi

n/a

Lamp Information Serial Number

Element(s) Fe

. . .

Deuterium Lamp Hours: 68.34

Sequence Table

Shared Standards: Yes	
Action	Fe
Calibration	√
B.No.Group-II	√
B.No.Group-III	√
B.No.Group-IV	√
B.No.Group-V	√
B.No.Group-VI	✓

Sample Details

		Nominal Mass: 1.0000					
No.	Sample Id	Sample Mass	Dilution Ratio				
1	B.No.Group-II	1.0000	1.0000				
2	B.No.Group-III	1.0000	1.0000				
3	B.No.Group-IV	1.0000	1.0000				
4	B.No.Group-V	1.0000	1.0000				
5	B.No.Group-VI	1.0000	1.0000				

Valid Analysis Signatures

11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Analysis Audit Trail

- 11-06-2022 17:30:16 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
- 11-06-2022 17:38:27 Jyothi(M. Jyothi):DESKTOP-39TDEGC Error MD147 - Activity manually aborted by user.
- 11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
- 11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

MART SPECIALITIES LAB LLP.

Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Spectrometer Parameters - Fe

Element: Fe Wavelength: 248.3nm Background Correction: D2 Signal Type: Continuous Measurement Time: 4.0secs Use RSD Test: No

Flame Type: Air-C2H2 Nebuliser Uptake: 4secs Burner Height: 7.0mm

Sampling: None

Calibration Mode: Normal	
Concentration Units: mg/L	
Acceptable Fit: 0.990	
Standard 1	2.0000
Standard 2	4.0000
Standard 3	6.0000

No changes are recorded for this element

Measurement Mode: Absorbance Bandpass: 0.2nm High Resolution: Off Resamples: Fast Flier Mode: No

Lamp Current: 75% Optimise Spectrometer Parameters: No Number Of Resamples: 3

Flame Parameters - Fe

Fuel Flow: 0.9L/min Burner Stabilisation: 0mins Optimise Burner Height: No Auxilary Oxidant: Off Optimise Fuel Flow: No

Sampling Parameters - Fe

Calibration Parameters - Fe

Line Fit: Linear Scaled Units: mg/L Rescale Limit: 10.0% Standard 4 Standard 5 Use Stored Calibration: No Scaling Factor: 1.0000 Failure Action: Flag and Continue 8.0000 10.0000

Element Audit Trail - Fe

Solution Results - Fe

Y = 0.04924x + 0.0141 Fit: 0.9965 Characteristic Conc: 0.0894

> 1 2 3

> 1 2 3

> 1 2 3

> 1 2 3

1 2

3

Sample ID

Fe Blank

Fe Standard 1

Fe Standard 2

Fe Standard 3

Fe Standard 4

	4				Norma	I: Linea	ar Fit				
	Abs		*		*		*		-*-		*
	0 * -0-0	1	2	3	4	5	6	7	8	9	10
	-0-0		2	0		nc:mg/		,	0	0	10
Signal	Rsd	Conc				cted C					
Abs	%	mg/L			mg/L						
0.001	35.6	0.000)		Ũ						
0.001	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:26	i		
0.001	Backgro	ound: -0.0	03	11-06-2022 17:31:30							
0.000	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:35			
0.120	0.3	2.0000)								
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:05			
0.119	-	ound: 0.00						7:32:09			
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:13			
0.215	0.3	4.0000)								
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:40)		
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:45			
0.216	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:49)		
0.322	0.2	6.0000)								
0.323	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:19)		
0.322	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:23			
0.322	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:27			
0.411	0.3	8.0000)								
0.410	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:59)		
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:03			
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:07	•		

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Operator Name: Jyothi

Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Solution Results - Fe

Sample ID	Signal	Rsd	Conc	Corrected Conc
-	Abs	%	mg/L	mg/L
Fe Standard 5	0.493	0.1	10.0000	
1	0.494	Backg	ground: 0.005	11-06-2022 17:34:38
2	0.494	Backg	ground: 0.005	11-06-2022 17:34:42
3	0.493	Backg	ground: 0.006	11-06-2022 17:34:46
Fe B.No.Group-II	0.834	0.1	16.6490 C	16.6490 C
1	0.834	Backg	ground: 0.005	11-06-2022 17:35:14
2	0.833	Backg	ground: 0.005	11-06-2022 17:35:19
3	0.834	Backg	ground: 0.005	11-06-2022 17:35:23
Fe B.No.Group-III	0.759	0.1	15.1286 C	15.1286 C
1	0.758	Backg	ground: 0.003	11-06-2022 17:35:52
2	0.760	Backg	ground: 0.003	11-06-2022 17:35:56
3	0.758	Backg	ground: 0.004	11-06-2022 17:36:00
Fe B.No.Group-IV	0.645	0.3	12.8189 C	12.8189 C
1	0.647	Backg	ground: 0.003	11-06-2022 17:36:31
2	0.644	Backg	ground: 0.003	11-06-2022 17:36:35
3	0.645	Backg	ground: 0.003	11-06-2022 17:36:40
Fe B.No.Group-V	0.809	0.2	16.1488 C	16.1488 C
1	0.808	Backg	ground: 0.005	11-06-2022 17:37:11
2	0.809	Backg	ground: 0.005	11-06-2022 17:37:15
3	0.811	Backg	ground: 0.005	11-06-2022 17:37:20
Fe B.No.Group-VI	0.711	0.3	14.1597 C	14.1597 C
1	0.709	Backg	ground: 0.005	11-06-2022 17:37:55
2	0.713	Backg	ground: 0.005	11-06-2022 17:37:59
3	0.712	Backg	ground: 0.004	11-06-2022 17:38:03

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DCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00	
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-1 Sample Received Date : 11/06/2022 Report Date : 13/06/2022	
I	DETAILS OF THE SAMPLE	
Sample Name : NA	NA	
Name of the Manufacturer : NA		
Storage condition To b	up-I Mfg. : NA Exp. : NA e stored at room date date	
	Derature Batch : NA C±3°C size	
	by AAS Analysis.	
	11/06/2022	
Analysis Completion Date : 11/0	6/2022	
Mfg. License No. : Not	provided	
A.R.NO : NA	The second se	

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	620ppm

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Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-2 Sample Received Date : 11/06/2022 Report Date : 13/06/2022	
Q	DETAILS OF THE SAMPLE	
Sample Name : N	NA	
Name of the Manufacturer : N	A	
Storage condition : To	roup-II Mfg. : NA Exp. : NA be stored at room date date	
Room Temperature : 25	Batch : NA size	
Tests Required:IreMethod:NAnalysis Starting Date:11	on by AAS Analysis.	
	Not provided	
A.R.NO : N	A	

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	832.5ppm

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Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinaga Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Sample Received Date : 11/06/2022	
a.d.	DETAILS OF THE SAMPLE	
Sample Name : N	JA	
Name of the Manufacturer : N	JA	
Storage condition	Group-III Mfg. : NA Exp. : NA To be stored at room date date Cemperature Batch : NA	
	5°C±3°C size	
	gm /	
Method : I Analysis Startin <mark>g Date : I</mark>	ron by AAS Analysis. NA 1/06/2022 1/06/2022	
and the second sec	Not provided	
A.R.NO	NA	
Remark : S	Sample analyzed as received	

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	756.5ppm

×

Authorized Signatory

(Dr.R.Marayya)

Office: 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile: 98481 98350, 98481 25019 Mail: rmarayya@yahoo.com

MART Specialities Lab. LLP

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College	
Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000	Sample Received Date: 11/06/2022Report Date: 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name : N	A
Name of the Manufacturer : N	A
Storage condition : T	roup-IV Mfg. : NA Exp. : NA o be stored at room date date emperature Batch : NA
	5°C±3°C size
Method : N Analysis Starting Date : 1	on by AAS Analysis. A 1/06/2022 1/06/2022
Mfg. License <mark>No. :</mark> N	ot provided
A.R.NO : N	A
Remark : Sa	ample analyzed as received

Test Results

Marine Marine Street Street		for second	
S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	641ppm	

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DCA Approval No: 05/ML/TS/2020/	G
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Custome	r: Reference / Report No. : MSL/2022/JUNE/TARA/0064-5
Tara Government College	Sample Received Date : 11/06/2022
Prashanth Nagar Colony, Balajinag	ar Sumpto Received 2 mil
Sangareddy Telangana. 502000	Report Date : 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name :	NA
Name of the Manufacturer :	NA
Batch no : Storage condition	Group-VMfg. : NAExp. : NATo be stored at roomdatedateTemperatureBatch : NA
Room Temperature :	25°C±3°C size
Quantity Received	4gm
Tests Required:Method:Analysis Starting Date:	Iron by AAS Analysis. NA 11/06/2022
Analysis Completion Date :	11/06/2022
Mfg. License No.	Not provided NA
A.R.NO Remark	Sample analyzed as received

Test Results

S.No.	Test Parameter	Result
01. Ir	on by AAS Analysis: (ppm)	807.5ppm

>

Authorized Signatory

(Dr.R.Marayya)

Office: 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile: 98481 98350, 98481 25019 Mail: rmarayya@yahoo.com

MART Specialities Lab. LLP

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-6 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
1	DETAILS OF THE SAMPLE
Storage condition	up-V1 Mfg. : NA Exp. : NA be stored at room date date perature Batch : NA C±3°C size
Tests Required:IronMethod:NAAnalysis Starting Date:11/0Analysis Completion Date:11/0Mfg. License No.:NotA.R.NO:NA	by AAS Analysis. 6/2022 6/2022 provided

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	708ppm

*

Authorized Signatory

(Dr.R.Marayya)

ANALYSIS OF ADSORPTION OF Fe[III] ON CELLULOSE USING D-RIBOSE AS A CHELATING AGENT

Dissertation submitted in Partial fulfillment for the requirements for the award of degree of

Bachelor of Science ín CHEMISTRY

By

N.Shiva shankar,
 K.Rahul,
 L.Sai ram goud,
 M.Arjun,
 M.Madhav,
 M.Binesh,

B.Sc(Chemistry),	III	Year
B.Sc(Chemistry),	III	Year

Under the Guídance:

Dr. K.ABHIJIT *HEAD* DEPARTMENT OF CHEMISTRY



Tara Government College, Sangareddy(A) 2021-22

<u>CERTIFICATE</u>

This is to certify that the project work entitled "ANALYSIS OF ADSORPTION OF Fe[III] ON CELLULOSE USING D-RIBOSE AS A CHELATING AGENT" is presented by B.Sc (CHEMISTRY) students in partial fulfillment of the requirements for the degree of Bachelor of Science in Chemistry by the Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

The results embodied in this report have not been to any other University or Institution for the award of any degree.

(Dr.K.ABHIJIT)

Project Supervisor& Head, Department of Chemistry Tara Government College, Sangareddy(A)

EXTERNAL EVALUATOR

ACKNOWLEDGEMENTS

We express my deep gratitude to my research supervisor **Dr.K.Abhijit**, Head, Department of Chemistry, Tara Govt. College, Sangareddy(A)-502001, INDIA for his inspiring guidance during the course of the Project work. The continuous encouragement extended by him propelled me to update my research skills and pedigree to engage in fruitful research.

We wish to express our gratitude to Smt. M.Praveena, Principal, Tara Govt. College, Sangareddy(A)-502001, INDIA for her constant support, cooperation and suggestions during the research work. We wish to express my sincere thanks to K.Sreedhar, Asst. Professor, Department of Chemistry for providing me facilities, help and support for the entire research work.

We express our special thanks to **MART Specialities Lab, Hyderabad** for providing technical assistance in Atomic absorption spectrometric analysis.

Finally, we take this opportunity to thank one and all that has directly or indirectly helped me in completing the task.

- 1.N.Shiva shankar,
- 2.K.Rahul,
- 3.L.Sai ram goud,
- 4.M.Arjun,
- 5.M.Madhav,
- 6.M.Binesh,

- B.Sc(Chemistry), III Year

<u>DECLARATION</u>

We hereby declare that the project report entitled "ANALYSIS OF ADSORPTION OF Fe[III] ON CELLULOSE USING D-RIBOSE AS A CHELATING AGENT" is the work done by us in the campus at Department of Chemistry, Tara Government College, Sangareddy(A)during the academic year 2021-2022 and is submitted in partial fulfillment of the requirements for the degree of Bachelor of Science by Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

S.No.	Name of the Student	Roll Number	Group	Year	Signature
1	N.Shiva shankar	6058-19-578-025	B.Sc (MCCS)	III	
2	K.Rahul	6058-19-578-019	B.Sc (MCCS)	III	
3	L.Sai ram goud	6058-19-578-020	B.Sc (MCCS)	III	
4	M.Arjun	6058-19-578-021	B.Sc (MCCS)	III	
5	M.Madhav	6058-19-578-024	B.Sc (MCCS)	III	
6	M.Binesh	6058-19-578-023	B.Sc (MCCS)	III	

TARA GOVERNMENT COLLEGE, SANGAREDDY – 502 001 (AUTONOMOUS)

BONAFIDE CERTIFICATE

Certified that the project ANALYSIS OF ADSORPTION OF Fe[III]

ON CELLULOSE USING D-RIBOSE AS A CHELATING AGENT"

is the bonafidework of

2.K.Rahul, 4.M.Arjun, 5.M.Madhav, 6.M.Binesh,

1.N.Shiva shankar, B.Sc(Chemistry), III Year B.Sc(Chemistry), III Year 3.L.Sai ram goud, B.Sc(Chemistry), III Year B.Sc(Chemistry), III Year B.Sc(Chemistry), III Year B.Sc(Chemistry), III Year

who carried out the project work under my supervision.

Dr.K.ABHIJIT PROJECT SUPERVISOR Smt. M.PRAVEENA PRINCIPAL

CONTENTS

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Chapter-I

INTRODUCTION

ADSORPTION

Adsorption is a process in which solids come into contact with liquids or gases, and the mass transfer occurs from liquids to solids. Desorption is the reversal of this action. Adsorption operations take advantage of a solid's capacity to concentrate certain chemicals from a fluid on to its surface. Adsorbate refers to the adsorbed substance, while adsorbent refers to the solid substance. The following are some examples of solid-liquid and solid-gas applications:

- Removing dissolved moisture from gasoline.
- Decolorization of sugar solutions and petroleum products.
- Removing noxious odors and tastes from water. Dehumidification of air and gases is one of the solid-gas activities.
- To fractionate mixtures of hydrocarbon gases such as methane, ethane, and propane.
- To remove undesirable odors and contaminants from gases.
- To recover valuable solvent vapors from dilute gas mixtures.

NATURE OF ADSORBENTS

Adsorbents are typically granular in nature, ranging in size from 0.5 mm to 12 mm. They can't have a lot of pressure decrease or get swept away by a fast-moving stream. During handling, they must maintain their shape and size. They'll need a lot of pores and a lot of surface area per unit mass.

Some of the commonly used adsorbents, their sources and applications are given below:

Sl. No.	Adsorbent	Source	Application
1.	Fuller's earth	Naturally occurring clay is heated and dried to get a porous structure.	De-colorizing, drying of lubricating oils, kerosene and engine oils.
2.	Activated charcoal	Bentonite or other activated clay which are activated by treatment with sulfuric acid and further washing, drying and crushing.	Used for de-colorizing petroleum products.
3.	Bauxite	A naturally occurring hydrated alumina, activated by heating at 230-815	Used for de-colorizing petroleum products and for drying gases.

4.	Alumina	A hard hydrated aluminium oxide, which is activated by heating to drive off the moisture and then crushed to desired size.	Used as desiccant.
5.	Bone-char	Obtained by destructive distillation of crushed bones at 600-900	Used for refining sugar and can be reused after washing and burning.
6.	Silica gel	A hard granular and porous product obtained from sodium silicate solution after treatment with acid. Normally has 4 to 7% water in the product.	Dehydration of gases and liquids, and separation of gas-liquid hydrocarbon mixture.

7.	Activated carbon	 (1) Vegetable matter is mixed with calcium chloride, carbonized and finally the inorganic compounds are leached away. (2) Organic matter is mixed with porous pumice stones and then heated and carbonized to deposit the carbonaceous matter throughout the porous particle. (3) Carbonizing substances like wood, sawdust, coconut shells, fruit pits, coal, lignite and subsequent activation with hot air steam. It is available in granular or pellated form. 	De-colorizing of sugar solutions, chemicals, drugs, water purification, refining of vegetable and animal oils, recovery of gold and silver from cyanide ore-leach solution, recovery of solvent vapour from gas-mixtures, collection of gasoline hydro- carbons from natural gas, fractionation of hydrocarbon gases.
8.	Molecular sieves	These are porous synthetic zeolite crystals, metal alumino-silicates.	Dehydration of gases and liquids, and separation of gas-liquid hydrocarbon mixture.

SIGNIFICANCE OF RESEARCH PROBLEM

Due to its polyhydric functional groups, cellulose may be utilised to adsorb heavy metals, making it a valuable tool for developing environmentally acceptable remediation methods. The adsorption efficiency of cellulose must be increased in the interim by the addition of facilitating agents in order to design and create standardised heavy metal remediation technology. By stabilising the metal complexes that will more strongly bind to the adsorbent, organic ligands will be helpful for this aim in bringing the metal-adsorbent connections.

OBJECTIVE OF THE PROJECT

To evaluate the impact of D-Ribose as chelating agent in the adsorption of Fe (III) from aqueous solution by Cellulose as an adsorbent to develop efficient remediation technology using concept of Coordination chemistry.

Chapter-II

ADSORPTION OF HEAVY METALS FROM DILUTE SOLUTIONS

The phrase "heavy metals" refers to a collection of metals and metalloids that have an atomic density more than 6 g cm⁻³. Although it is a broad phrase, it is typically used to refer to elements like Cd (cadmium), Cr (chromium), Cu (copper), Hg (mercury), Ni (nickel), Pb (lead), and Zn (zinc) that are commonly related with pollution and toxicity problems. Heavy metals, unlike most organic contaminants, occur naturally. Each of these elements has a range of normal background concentrations in soils, sediments, waterways, and living creatures because they occur naturally in rock-forming and ore minerals. Significant amounts of various heavy metals are produced each year from the mining of their respective ores. Soils, sediments, waterways, and live species all play a role in the environment. Significant amounts of various heavy metals are produced each year from the mining of their respective ores. Metals used in industry and other domestic processes (e.g., burning fossil fuels, incineration of wastes, automobile exhausts, smelting processes, and the use of sewage sludge as a landfill material and fertiliser) have released large amounts of potentially toxic heavy metals into the atmosphere, as well as into aquatic and terrestrial environments. Cd, Pb, and Zn are common hazardous metals released into the environment.Cu, Ni, Cr, Co, Zn, and Pb are some of the most common metals (Babich et al., 1985). Table 1 shows the industrial sources of a variety of metals emitted, as well as the potential for contamination. While many heavy metals are required by plants as micronutrients, larger quantities have been linked to a variety of harmful effects. Lead causes encephalopathy, cognitive impairment, behavioural problems, renal damage, anaemia, and reproductive system toxicity at high exposure levels (Pagliuca and Mufti, 1990). In its hexavalent state, chromium is

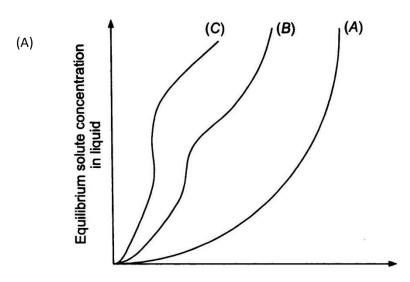
well acknowledged to be hazardous (Rowbotham et al., 2000). Exposure to Cr (VI) compounds has been linked to a greater incidence of respiratory malignancies in humans (IARC, 1990). Cadmium has been linked to nephrotoxic effects, especially at high levels of exposure; long-term exposure may also induce bone dam- age (Friberg, 1985). Neurobehavioural problems and developmental abnormalities, such as dyslexia, attention deficit hyperactivity disorder, and intellectual retardation, can be caused by high mercury levels (Weiss and Landrigan, 2000). Copper poisoning can cause weakness, tiredness, and anorexia, as well as damage to the stomach and intestines (Theophanides and Anastassopoulou, 2002). Nordberg et al. go into great length about the toxicity of nickel and other heavy metals (2007)

Industry	Metals	Pollution Arising	Reference
Metalliferous mining	Cd,Cu,Ni,Cr,Co,Zn	Acid mine drainage,	Babich et al.(1985)
		tailings, slag heaps	Aswathanarayana(2003)
Agriculture materials	Cd,Cr,Mo,Pb,U,V,Zn	Run-off, surface and	Nicholson et al.(2003)
Fertilisers		groundwater	Otero et al.(2005)
		contamination, plant	
		bioaccumulation	
Manures sewage	Zn,Cu,Ni,Pb,Cd,Cr,As,Hg	Landspreading threat to	Nicholson et al.(2003)
sludge		ground and surface water	Cheung and Wong
			(1983)
			Walter et al.(2006)
Metallurgical	Pb,Mo,Ni.Cu,Cd,As,Te,U,	Manufacture, disposaland	Alloway and Ayres

 Table 1 : Significant anthropogenic sources of metals in the environment.

industries Specialist	Zn	recycling of metals,	(1993)
alloys and steels		Tailings and slag heaps	Rule et al (2006)
			Cheng (2003)
Waste disposal	Zn,Cu,Cd,Pb,Ni,Cr,Hg	Landfill leachate,	Kjeldson et al.(2002)
Landfill leachate		contamination of ground	Fernandez et al.(2005)
		and surface	
Electronics	Pb,Cd,Hg,Pt,Au,Cr,As,Ni,	Aqueous and solid	Veglio et al. (2003)
	Mn	metallic waste food	
		manufacturing and	
		recycling process	
Metal finishing	Cr,Ni,Zn,Cu	Liquid effluents from	Castelblanque and
industry		plating processes	salimbeni(2004)
Electroplating			Zhao et al.(1999)
			Alvarez-Ayuso et
			al.(2003)
Miscellaneous	Pb,Sb,Zn,Cd,Ni,Hg	Waste battery fluid,	EU Directorate general
sources Batteries		contamination of soil and	of the Environment
		groundwater	(2004)
Paints and pigments	Pb,Cr,As,Ti,Ba,Zn	Aqueous waste from	Davis and Burns (1999)
		manufacture, old paint	Barnes and Davis (1996)
		deterioration and soil	Monken(2000)
		pollution	

Both the solvent and the solute are adsorbed whenever a mixture of solute and solvent is adsorbed using an adsorbent. As a result, only relative or apparent solute adsorption can be determined. As a result, treating a known volume of solution of original concentration C with a known weight of adsorbent is standard procedure. Let C* be the solution's final equilibrium solute concentration. If v is the volume of solution per unit mass of adsorbent (cc/g), and C and C* are the starting and equilibrium concentrations (g/cc) of the solute, then the apparent adsorption of the solute per unit mass of adsorbent is $v(C - C^*)$, (g/g), neglecting any volume change. This statement is most useful in the case of dilute solutions. The C* value is determined by the temperature, nature, and properties of the adsorbent when the proportion of the original solvent that can be adsorbed is tiny. The Freundlich adsorption isotherm, $C^*=K[v(C - C^*)]n$, represents the adsorption phenomena in dilute fluids across a small concentration range. The Freundlich adsorption equation is especially useful in situations where the identification of the solute is unknown, such as the removal of colouring substances from sugar solutions, oils, and other liquids. A spectrophotometer or colorimeter can quickly determine the colour composition of the solute. In worked example 2, the interpretation of this data is demonstrated. Adsorption is good if the value of n is high, say 2 to 10. If it's between 1 and 2, it's relatively challenging, and if it's less than 1, it's easy and it indicates poor adsorption characteristics. Freundlich adsorption equation is also useful in such a case where the actual identification of solute is not known, e.g. removal of colouring substance from sugar solutions, oils etc. A typical adsorption isothermal for the adsorption of various adsorbents A, B and C in dilute solution at the same temperature for the same adsorbent is represented in a graph.



Kg solute apparently adsorbed / kg adsorbent

METAL COMPLEXATION OF A D- RIBOSE- BASED LIGAND

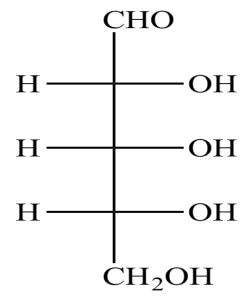
The complexation characteristics of methyl 2,3,4-tri-O-(2-picolyl)—D-ribopyranoside, a novel sugar-derived hexadentate ligand, were investigated using a combination of experimental and theoretical approaches. In the complexes with Mn[#], Co[#], Ni[#], and Zn[#], the coordination bond lengths reveal significant departures from perfect octahedral, with deformation towards trigonal-prismatic geometries, indicating a conformationally constrained ligand. The metal-cation–ligand interactions for D-Ribose and its acyclic analogue ligand [1, 2, 3-tri-O-(2-picolyl)-1,2,3-propanetriol] were investigated using spectroscopic techniques and isothermal calorimetric titrations for the seriesMn[#], Co[#], Ni[#], Zn[#], and Cu[#]. Depending on the nature of the metal, the results show that the complexes produced with D-Ribose are more stable than those obtained with According to molecular modelling studies; the presence of the sugar moiety greatly favours conformations that are compatible with metal binding. In many ways, a close comparision can be sketched between metal complexation and other host-guest interactions. To describe them both, several molecular variables have to be taken into consideration. One of the most complex

variables is the relation between the conformational flexibility of the ligand and the binding strength of the metal cation. For ligands with a reduced number of low-energy conformations, relatively straightforward estimations of the binding same wavelength of metal cations as well as a direct interaction between structural and thermodynamic data are available. However, for ligands with a huge number of degrees of freedom, the correct orientation of the chelating groups becomes one of the most important issues in metal-ion identification processes. For coordination chemists, the prediction and understanding about coordination properties of new ligands is one the most challenging tasks. We decided to investigate how entropic and enthalpic variables can impact on metal binding and selectivity for systems with a substantial number of degrees of freedom.

FERRIC COMPLEXES OF SUGAR-TYPE LIGAND

Fe(III) complexes formed with sugar-type ligands such as aldoses, ketoses, polyalcohols, sugar acids, di and trisaccharides was prepared. These complexes composition was determined by standard analytical methods. Deprotonated alcoholic hydroxy groups participate in the complex formation which results in the formation of polynuclear species. Mossbauer spectra reflected the presence of high spin iron(III) central atoms.EPR spectra showed antiferromagnetic interactions between the iron(III) centres in the complexes indicating dimeric or oligomeric complex structures. Depending upon the nature of a ligand as well as on the preparation mode of the complexes the ratio of interacting and isolated iron(III) is calculated. Formation of polynuclear iron(III) complexes of D-fructose, sorbose, lactobionic acid, glucose, galactose, mannose, and lactose were shown by the analysis and structural study by the sugar type ligand with ferric system. Ferric complexes of reducing sugars also contained ferrous species in some

cases. The combination of Mossbauer spectroscopic measurements with ERP or magnetic susceptibility studies has led valuable information concerning the intramolecular and electronic structure on which the system has been developed.



Structure of D-Ribose

CELLULOSE AS ADSORBENT

Cellulose is the most plentiful and renewable polymer material available on the planet. According to estimates, deterioration and soil pollution Photosynthesis produces 10^{11} – 10^{12} tonnes of cellulose every year in a relatively pure form, such as in the seed hairs of cotton plants, but it is more commonly mixed with lignin and other polysaccharides (so-called hemicelluloses) in the cell walls of woody plants (Klemm et al., 2002). For thousands of years, cellulose has been utilised as an energy source, a building material, and a clothing material in the form of wood and cotton.

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As a carbohydrate polymer, cellulose has a molecular structure that is made up of repeating b-D-glucopyranose units that are covalently linked by acetal functionalities between the OH groups of the C4 and C1 carbon atoms (b-1,4-glucan). Cellulose is a long, linear-chain polymer with a lot of hydroxyl groups (three per anhydroglucose (AGU) unit) and the 4C1 conformation. Every second AGU unit is rotated 180 degrees in the plane to accommodate the preferred bond angles. The number of constituent AGU units (degree of polymerization, DP) determines the length of the polymeric cellulose chain, which varies depending on the origin and treatment of the cellulose raw material (Klemm et al., 2002).

Cellulose has a ribbon form that allows it to twist and bend in directions other than the plane, making it a fairly flexible molecule. Due to the presence of hydroxyl (–OH) groups that protrude from the chain and create intermolecular hydrogen bonds, there is a relatively strong interaction between neighbouring cellulose molecules in dry fibres. Each chain of cellulose regenerated fibres contains 250–500 repeating units (Klemm et al., 2002). The hydrophilicity, chirality, and degradability features of cellulose are due to this molecular structure. Chemical reactivity is mostly determined by the OH groups' high donor reactivity.

Chapter-III

ATOMIC ABSORPTION SPECTROMETRY

Atomic absorption spectrometry (AAS) is an analytical technique that measures the concentrations of elements. Atomic absorption is so sensitive that it can measure down parts per billion of a gram (μ g dm⁻³) in a sample. The technique makes use of the wavelengths of light specifically absorbed by an element. They correspond to the energies needed to promote electrons from one energy level to another, higher, energy level.

Atomic absorption spectrometry has many uses in different areas of chemistry.

Clinical analysis: Analysing metals in biological fluids such as blood and urine.

Environmental analysis: Monitoring our environment- eg finding out the levels of various elements in rivers, seawater, drinking water, air, petrol and drinks such as wine, beer and fruit drinks.

Pharmaceuticals: In some pharmaceutical manufacturing processes, minute quantities of a catalyst used in the process (usually a metal) are sometimes present in the final product. By using AAS the amount of catalyst present can be determined.

Industry: Many raw materials are examined and AAS is widely used to check that the major elements are present and that toxic impurities are lower than specified- eg in concrete, where calcium is a major constituent, the lead level should be low because it is toxic.

Mining: By using AAS the amount of metals such as gold in rocks can be determined to see whether it is worth mining the rocks to extract the gold.

HOW IT WORKS

Atoms of different elements absorb characteristic wavelengths of light. Analysing a sample to see if it contains a particular element means using light from that element. For example with lead, a lamp containing lead emits light from excited lead atoms that produce the right mix of wavelengths to be absorbed by any lead atoms from the sample. In AAS, the sample is atomized- i.e. converted into ground state free atoms in the vapour state- and a beam of electromagnetic radiation emitted from excited lead atoms is passed through the vaporized sample. Some of the radiation is absorbed by the lead atoms in the sample. The greater the number of atoms there is in the vapour, the more radiation is absorbed. The amount of light absorbed is proportional to the number of lead atoms. A calibration curve is constructed by running several samples of known lead concentration under the same conditions as the unknown. The amount the standard absorbs is compared with the calibration curve and this enables the calculation of the lead concentration in the unknown sample.

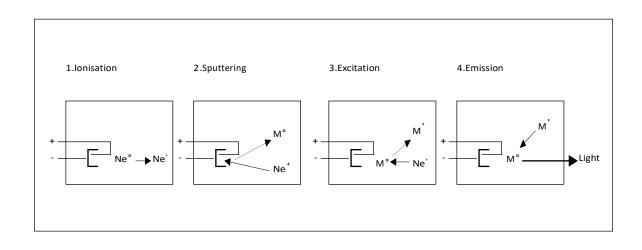
Consequently an atomic absorption spectrometer needs the following three components: a light source; a sample cell to produce gaseous atoms; and a means of measuring the specific light absorbed.

THE LIGHT SOURCE

The common source of light is a 'hollow cathode lamp'. This contains a tungsten anode and a cylindrical hollow cathode made of the element to be determined. These are sealed in a glass tube filled with an inert gas- e.g neon or argon- at a pressure of between 1 Nm⁻² and 5 Nm⁻².

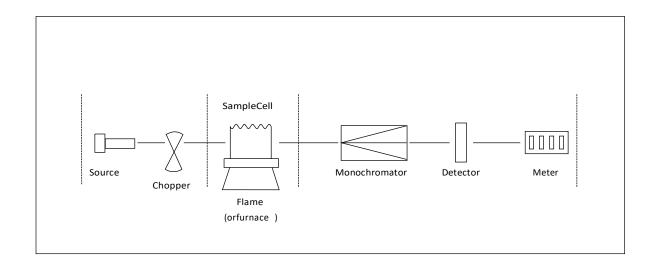


The ionization of some gas atoms occurs by applying a potential difference of about 300-400V between the anode and the cathode. These gaseous ions bombard the cathode and eject metal atoms from the cathode in a process called sputtering. Some sputtered atoms are in excited states and emit radiation characteristic of the metal as they fall back to the ground state $-egPb^* \rightarrow Pb + hv$. The shape of the cathode concentrates the radiation into a beam which passes through a quartz window, and the shape of the lamp is such that most of the sputtered atoms are redeposited on the cathode. A typical atomic absorption instrument holds several lamps each for a different element. The lamps are housed in a rotating turret so that the correct lamp can be quickly selected.



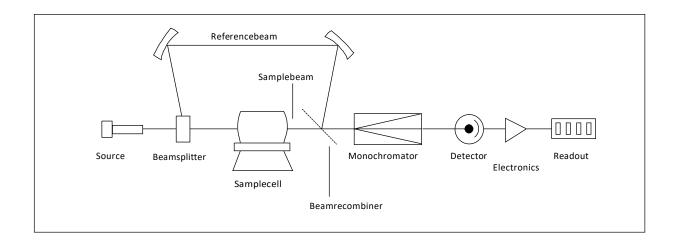
THE OPTICAL SYSTEM AND DETECTOR

A monochromator is used to select the specific wavelength of light –ie spectral line – which is absorbed by the sample, and to exclude other wavelengths. The selection of the specific light allows the determination of the selected element in the presence of others. The light selected by the monochromator is directed onto a detector that is typically a photomultiplier tube. This produces an electrical signal proportional to the light intensity



DOUBLE BEAM SPECTROMETERS

Modern spectrometers incorporate a beam splitter so that one part of the beam passes through the sample cell and the other is the reference. The intensity of the light source may not stay constant during an analysis. If only a single beam is used to pass through the atom cell, a blank reading containing no analyte (substance to be analysed) would have to be taken first, setting the absorbance at zero. If the intensity of the source changes by the time the sample is put in place, the measurement will be inaccurate. In the double beam instrument there is a constant monitoring between the reference beam and the light source. To ensure that the spectrum does not suffer from loss of sensitivity, the beam splitter is designed so that as high a proportion as possible of the energy of the lamp beam passes through the sample.



ATOMISATION OF THE SAMPLE

Two systems are commonly used to produce atoms from the sample. Aspiration involves sucking asolution of the sample into a flame; and electrothermal atomisation is where a drop of sample is placed into a graphite tube that is then heated electrically.

Some instruments have both atomisation systems but share one set of lamps. Once the appropriate lamp has been selected, it is pointed towards one or other atomisation system.

FLAME ASPIRATION

Ethyne/air (giving a flame with a temperature of 2200–2400°C) or ethyne/dinitrogen oxide (2600– 2800°C) are often used. A flexible capillary tube connects the solution to the nebuliser. At the tip of the capillary, the solution is 'nebulised' –ie broken into small drops. The larger drops fall out and drain off while smaller ones vaporise in the flame. Only ca 1% of the sample is nebulised.

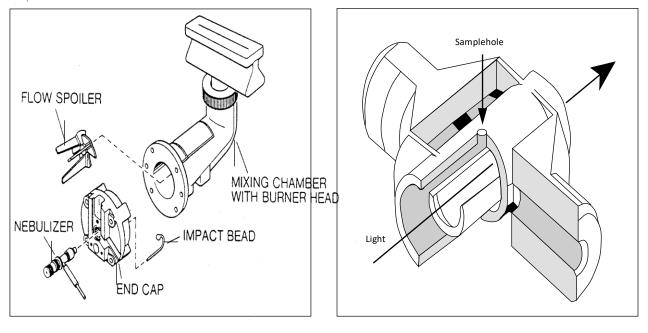


Figure 1

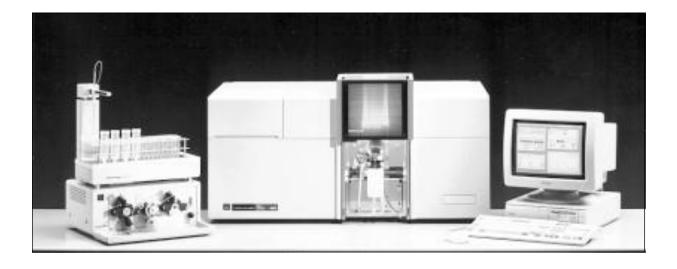
Figure 2

ELECTRO-THERMAL ATOMIZATION

25 µl of sample (ca 1/100th of a raindrop) is placed through the sample hole and onto the platform from an automated micropipette and sample changer. The tube is heated electrically by passing a current through it in a pre-programmed series of steps. The details will vary with the sample but typically they might be 30–40 seconds at 150°C to evaporate the solvent, 30 seconds at 600°C to drive off any volatile organic material and char the sample to ash, and with a very fast heating rate (ca 1500 °C s⁻¹) to 2000-2500°C for 5–10 seconds to vaporise and atomise elements (including the element being analysed). Finally heating the tube to a still higher temperature –ca 2700°C – cleans it ready for the next sample. During this heating cycle the graphite tube is flushed with argon gas to prevent the tube burning away. In electrothermal atomisation almost 100% of the sample is atomised. This makes the technique much more sensitive than flame AAS.

SAMPLE PREPARATION

Sample preparation is often simple, and the chemical form of the element is usually unimportant. This is because atomisation converts the sample into free atoms irrespective of its initial state. The sample is weighed and made into a solution by suitable dilution. Elements in biological fluids such as urine and blood are often measured simply after a dilution of the original sample.



When making reference solutions of the element under analysis, for calibration, the chemical environment of the sample should be matched as closely as possible –i.e. the analyte should be in the same compound and the same solvent. Teflon containers may be used when analyzing very dilute aolutions because elements such as lead are sometimes leached out of glass vessels and can affect the results

BACKGROUND ABSORPTION

It is possible that other atoms or molecules apart from those of the element being determined will absorb or scatter some radiation from the light source. These species could

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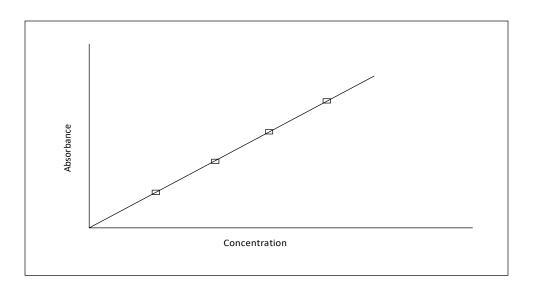
include unvaporised solvent droplets, or compounds of the matrix (chemical species, such as anions, that tend to accompany the metals being analysed) that are not removed completely. This means that there is a background absorption as well as that of the sample.

One way of measuring and correcting this background absorption is to use two light sources, one of which is the hollow cathode lamp appropriate to the element being measured. The second light source is a deuterium lamp.

The deuterium lamp produces broad band radiation, not specific spectral lines as with a hollow cathode lamp. By alternating the measurements of the two light sources – generally at 50 -100 Hz – the total absorption (absorption due to analyte atoms plus background) is measured with the specific light from the hollow cathode lamp and the background absorption is measured with the light from the deuterium lamp. Subtracting the background from the total absorption gives the absorption arising from only analyte atoms.

CALIBRATION

A calibration curve is used to determine the unknown concentration of an element -eglead – in a solution. The instrument is calibrated using several solutions of known concentrations. A calibration curve is produced which is continually rescaled as more concentrated solutions are used – the more concentrated solutions absorb more radiation up to a certain absorbance. The calibration curve shows the concentration against the amount of radiation absorbed in the given figure. (a) The sample solution is fed into the instrument and the unknown concentration of the element-e.g. lead- is then displayed on the calibration curve given in the below figure. (b)





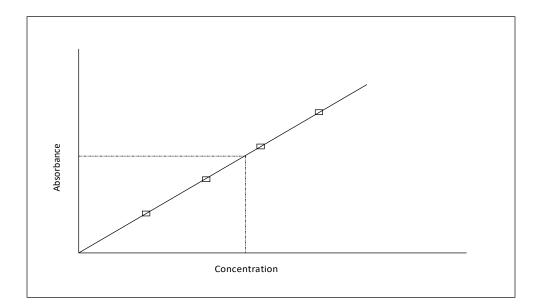


Figure (b)

INTERFERENCES AND MATRIX MODIFICATION

Other chemicals that are present in the sample may affect the atomisation process. For example, in flame atomic absorption, phosphate ions may react with calcium ions to form calcium pyrophosphate. This does not dissociate in the flame and therefore results in a low reading for calcium. This problem is avoided by adding different reagents to the sample that may react with the phosphate to give a more volatile compound that is dissociated easily. Lanthanum nitrate solution is added to samples containing calcium to tie up the phosphate and to allow the calcium to be atomised, making the calcium absorbance independent of the amount of phosphate. With electrothermal atomisation, chemical modifiers can be added which react with an interfering substance in the sample to make it more volatile than the analyte compound. This volatile component vaporises at a relatively low temperature and is removed during the low and medium temperature stages of electrothermal atomisation.

Chapter-IV

METHODOLOGY

MATERIALS REQUIRED

- Ferric chloride Hexahydrate
- D- Ribose
- Cellulose
- Volumetric flask
- Digital Weighing Machine
- Watch glasses
- Hot air oven
- Ultra pure water (Demineralized)
- 2% of Nitric acid

PROCEDURE

- Using a 250ml Volumetric flask prepare a metal ligand solution by adding 100ppm of Ferric chloride Hexahydrate (MOLYCHEM MCR-11580) and 200ppm of D-Ribose (Avra N2200133). Prepare 250ml solution by adding Ultra pure water and then keep this system aside for a few hours.
- Weigh 5 grams of Cellulose(Avra N2001901) using a Digital weighing machine (Citizen Scales(I) PVT LTD CTG302-300) and take this into a beaker.

- **3.** Now add 100ml of the above prepared metal ligand solution into the beaker and stir the mixture well for 10 minutes using a glass rod.
- **4.** Keep this mixture aside for 48 hours without disturbing it as at this step Ferric is going to be adsorbed on Cellulose in the presence of D-Ribose which acts as a chelating agent.
- **5.** After completion of 48 hours take the mixture and filter it off using Whatman Grade 1 filter paper and a funnel.
- **6.** After filtration of the mixture again add Ultra pure water for 3 times and then filtrate it to obtain pure concentration of Ferric which is get adsorbed on Cellulose.
- 7. Collect the filtered Cellulose powder and place it on a watch glass and keep this in a Hot air owen at 60 °C for 10 hours to get rid off moisture present in it.
- **8.** Now weigh each 1 gm of Cellulose in glass vials.
- 9. Now take a beaker and rinse it with ultra pure water then followed by Nitric acid.
- **10.** Take 0.5 grams of Cellulose sample in the beaker and add 2% of Nitric acid and stir the mixture well for 10-15 minutes.
- **11.** Filter the mixture using Whatman Grade 1 filter paper and again 3 times by using Ultra pure water to obtain pure concentration of Ferric present in the mixture prepared using the sample.
- **12.** Take this collected sample solution and keep this system under AAS (Thermo Scientific iCE 3300)
- **13.** Calculate the concentration of Ferric adsorbed on Cellulose at different ppm levels. Observe the graph obtained and note down the readings of the result we obtained.
- **14.** Same Experiment carried out without the interference/addition of Ligand i.e. D-Ribose for Control Experiment.



Ferric chloride Hexahydrate, D-Ribose



Preparing 250 ml of metal ligand solution using volumetric flask



5 grams of Cellulose added to 100ml of Metal Ligand solution



Metal Ligand solution



Measuring flask



Digital Weighing Machine



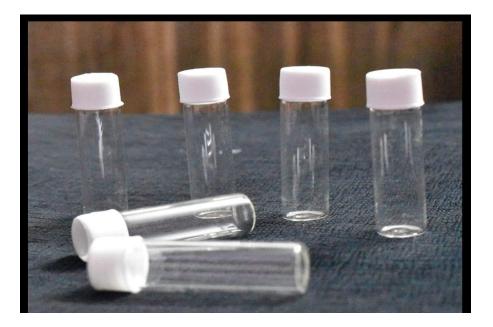
Hot air oven



Measuring 100ml of metal ligand prepared solution



Cellulose added to Metal Ligand solution is prepared



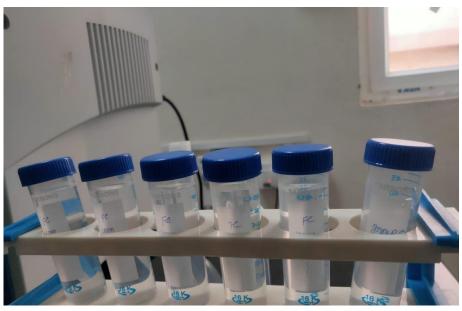
Glass vials



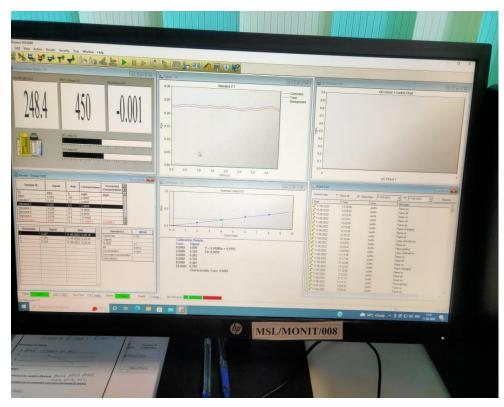
Sample collecting in glass vials



AA spectrometer



Sample solution added to 2% of nitric acid

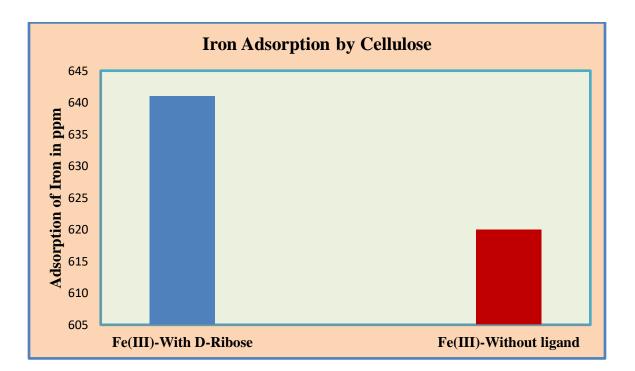


Results obtained on the monitor under AAS Method

Chapter-V

RESULTS AND DISCUSSIONS

Cellulose adsorbs **641** ppm of Iron metal from aqueous solution of Fe(III)-D-Ribose metal ligand solution. Whereas, Cellulose adsorbs only **620** ppm when D-Ribose is absent. It is evident from the AAS results, ligand involvement enhanced the metal adsorption by initiating potential chemical interactions between adsorbate and adsorbent. D-Ribose firmly interacts with Fe(III) to form a stable complex in aqueous condition. The complex coordination sphere in the resulted complex facilitates strong interactions with the polar hydroxyl functional groups of the adsorbent, Cellulose. From the AAS results, it is conclusive that **3.387** % of adsorbtion increased in the presence of D-Ribose as chelating agent.



Impact of D-Ribose on Adsorption of Fe (III) ions from aqueous solution by Cellulose.

Spectrometer Parameters – Fe:

Element : Fe	Measurement mode : Absorbance	
Wavelength : 248.3nm	Band pass : 0.2nm	Lamp current : 75%
Background correction : D2	High Resolution : Off	Optimise Spectrometer Parameters : No
Signal type : continuous	Resamples : Fast	Number of resamples : 3
Measurement time : 4.0secs	Flier mode : No	
Use RSD Test : No		

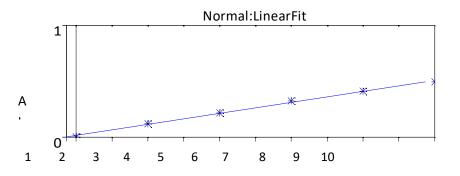
Flame Parameters – Fe:

Flame type : Air – C2H2	Fuel Flow : 0.9L/min	Auxiliary Oxidant : Off
Nebuliser Uptake : 4secs	Bunsen Stabilisation : Omins	Optimise Fuel Flow : No
Burner Height :7.0mm	Optimise Burner fuel : No	

Calibration Parameters-Fe:

Calibration mode : Normal	Line fit : Linear	Use stored calibration : No
Concentration units : mg/L	Scales units : mg/L	Scaling factor : 1.0000
Acceptable fit : 0.990	Rescale Limits : 10.0%	Failure Action : Flag and Continue
Standard 1 - 2.0000	Standard 4 - 8.0000	
Standard 2 - 4.0000	Standard 5 - 10.0000	
Standard 3 - 6.0000		

Solutions Results-Fe:



Y = 0.04924x + 0.0141

Fit: 0.9965

Characteristic Conc.: 0.0894

Sample ID	Signal	RSD	Conc.
	Abs	%	Mg/L
Fe Blank	0.001	35.6	0.0000
1	0.001	Backgr	ound: -0.003
2	0.001	Backgr	ound: -0.003
3	0.000	Backgr	ound: -0.003
Fe Standard 1	0.120	0.3	2.0000
1	0.120	Backgr	ound: 0.003
2	0.119	Backgr	ound: 0.003
3	0.120	Background: 0.003	
Fe Standard 2	0.215	0.3	4.0000
1	0.215	Background: 0.004	
2	0.215	Backgr	ound:0.004
3	0.216	Backgr	ound:0.004

Fe Standard 3	0.322	0.2	6.0000	
1	0.323	Background: 0.005		
2	0.322	Background: 0.005		
3	0.322	Backg	ground: 0.005	
Fe Standard 4	0.411	0.3	8.0000	
1	0.410	Backg	ground: 0.005	
2	0.412	Backg	ground: 0.005	
3	0.412	Backg	ground: 0.005	
Fe Standard 5	0.493	0.1	10.0000	
1	0.494	Background: 0.005		
2	0.494	Background: 0.005		
3	0.494	Background: 0.006		
Fe(III)-D-Ribose- Cellulose	0.645	0.3	12.8191 C	
1	0.647	Background: 0.003		
2	0.644	H	Background: 0.003	
3	0.645	Background: 0.003		
Fe(III)-Cellulose (Without Ligand)	0.624	0.3 12.3991 C		
1	0.622	Backg	ground: 0.003	
2	0.626	Backg	ground: 0.003	
3	0.624	Backg	ground: 0.003	

Test Results:

S.no	Test Parameters	Sample	Results
01.	Iron by AAS Analysis:	Fe(III)-D-Ribose –	641ppm
	(ppm)	Cellulose Sample	
02.	Iron by AAS Analysis:	Fe(III)– Cellulose	620 PPM
	(ppm)	(control)	

CONCLUSION

According to the results of the current experiment, D-Ribose, which acts as a chelating agent, is crucial to the adsorption of Fe (III) ions from aqueous solution by Cellulose and enhances adsorption by up to **3.387** %. Designing the latest heavy metal remediation systems that employ natural chelating ligands as facilitation agents in metal adsorption processes would benefit from this feature.

REFERENCES

1.Abdel-Aal, S.E., Gad, Y., Dessouki, A.M., 2006. The use of wood pulpand radiation modified starch in wastewater treatment. Journal of Applied Polymer Science 99, 2460–2469.

2.Abollino, O., Aceto, M., Malandrino, M., Sarzanini, C., Mentasti, E.,2003. Adsorption of heavy metals on Na-montmorillonite: effect of pH and organic substances. Water Research 37 (7), 1619–1627.

3.Aderhold, D., Williams, C.J., Edyvean, R.G.J., 1996. The removal ofheavy-metal ions by seaweeds and their derivatives. Bioresource Technology 58 (1), 1–6.

4.Ajmal, M., Rao, R.A.K., Ahmad, R., Ahmad, J., 2000. Adsorptionstudies on Citrus reticulata (fruit peel of orange): removal and recoveryof Ni (II) from electroplating wastewater. Journal of HazardousMaterials 79 (1–2), 117–131.

5.Altin, O., Ozbelge, H.O., Dogu, T., 1998. Use of general-purposeadsorption isotherms for heavy metal–clay mineral interactions.Journal of Colloid Interface Science 198, 130–140.

Alloway, B.J., Ayres, D.C., 1993. Chemical Principles of EnvironmentalPollution, First ed. Chapman & Hall, London, pp. 190–216.

6.Alvarez-Ayuso, E., Garcia-Sanchez, A., Querol, X., 2003. Purification ofmetal electroplating waste waters using zeolites. Water Research 37 (20), 4855–4862.

7.Andini, S., Cioffi, R., Montagnaro, F., Pisciotta, F., Santoro, L., 2006.Simultaneous adsorption of chlorophenol and heavy metal ions onorganophilic bentonite. Applied Clay Science 31 (1–2), 126–133.

8.Aoki, N., Fukushima, K., Kurakata, H., Sakamoto, M., Furuhata, K.,1999a. 6-Deoxy-6mercaptocellulose and its S-substituted derivativesas sorbents for metal ions. Reactive and Functional Polymers 42 (3),223–233.

9. Aoki, Y., Tanaka, K., Sakamoto, M., Furuhata, K., 1999b. Sorption ofmetal ions by bead cellulose grafted with amidoximated polyacrylonitrile. Sen'iGakkaishi 55 (12), 569–575.

10.Aswathanarayana, U., 2003. Mineral Resources Management and theEnvironment. Routledge, Netherlands, pp. 223–256.

11.Babich, H., Devanas, M.A., Stotzky, G., 1985. The mediation ofmutagenicity and clastogenicity of heavy metals by physicochemical factors. Environmental Research 37 (2), 253–286.

12.Bao-Xiu, Z., Peng, W., Tong, Z., Chun-yun, C., Jing, S., 2006. Preparationand adsorption performance of a cellulosic adsorbent resin forcopper (II). Journal of Applied Polymer Science 99, 2951–2956.

13.Barnes, G.L., Davis, A.P., 1996. Dissolution of lead paint in aqueoussolution. Journal of Environmental Engineering ASCE 122, 663–666.

14.Bayramoglu, G., Bektas, S., Arica, M.Y., 2003. Biosorption of heavymetal ions on immobilized white-rot fungus Trametes versicolor. Journal of Hazardous Materials 101 (3), 285–300.

15.Benke, N., Taka´cs, E., Wojna´rovits, L., Borsa, J., 2007. Pre-irradiationgrafting of cellulose and slightly carboxymethylated cellulose (CMC)fibres. Radiation Physics and Chemistry 76 (8–9), 1355–1359.

16.Berber-Mendoza, M.S., Leyva-Ramos, R., Alonso-Davila, P., Mendoza-Barron, J., Diaz-Flores, P.E., 2006. Effect of pH and temperature on the ion-exchange isotherm of Cd(II) and Pb(II) on clinoptilolite.Journal of Chemical Technology and Biotechnology 301,40–45.

17.Bhattacharya, A., Misra, B.N., 2004. Grafting: a versatile means tomodify polymers: techniques, factors and applications. Progress inPolymer Science 29 (8), 767–814.

18.Bhattacharya, A., 2000. Radiation and industrial polymers. Progress in Polymer Science 25 (3), 371–401.

19.Bicak, N., Sherrington, D.C., Senkal, B.F., 1999. Graft copolymer of acrylamide onto cellulose as mercury selective sorbent. Reactive and Functional Polymers 41, 69–76.

20.Bicak, N., Senkel, S.F., 1998. Sorbitol-modified poly(N-glycidyl styrenesulfonamide) for removal of boron. Journal of Applied PolymerScience 68, 2113–2119.

21.Brown, P., AtlyJefcoat, I., Parrish, D., Gill, S., Graham, E., 2000.Evaluation of the adsorptive capacity of peanut hull pellets for heavymetals in solution. Advances in Environmental Research 4 (1), 19–29.

22.Castelblanque, J., Salimbeni, F., 2004. NF and RO membranes for therecovery and reuse of water and concentrated metallic salts from wastewater produced in the electroplating process. Desalination 167, 65–73.

23.Cheng, S.P., 2003. Heavy metal pollution in China: origin, pattern and control. Environmental Science and Pollution Research 10 (3), 192–198.

24.Cheung, Y.H., Wong, M.H., 1983. Utilisation of animal manures and sewage sludges for growing vegetables. Agricultural Wastes 5 (2), 63–81.

25.Cochrane, E.L., Lu, S., Gibb, S.W., Villaescusa, I., 2006. A comparison oflow cost biosorbents and commercial sorbents for the removal ofcopper from aqueous media. Journal of Hazardous Materials 137 (1),198–206.

26.Crini, G., 2005. Recent developments in polysaccharide-based materialsused as adsorbents in wastewater treatment. Progress in PolymerScience 30 (1), 38–70.

27.Dal Bosco, S.M., Jimenez, R.S., Carvalho, W.A., 2005. Removal of toxicmetals from wastewater by Brazilian natural scolecite. Journal ofColloid and Interface Science 281 (2), 424–431.

28.Davis, A.P., Burns, M., 1999. Evaluation of lead concentration in runofffrom painted structures. Water Research 33 (13), 2949–2958.

29.Demirbas, E., Kobya, M., Oncel, S., Sencan, S., 2002. Removal of Ni (II)from aqueous solution by adsorption onto hazelnut shell activated carbon: equilibrium studies. Bioresource Technology 84 (3), 291–293.

30.Dilek, F.B., Erbay, A., Yetis, U., 2002. Ni (II) biosorption by polyporousversicolor. Process Biochemistry 37 (7), 723–726.

31.Dimitrova, S.V., 1996. Metal sorption on blast-furnace slag. WaterResearch 30 (1), 228–232.

Feng, D., van Deventer, J.S.J., Aldrich, C., 2004. Removal of pollutantsfrom acid mine wastewater using metallurgical by-product slags.Separation and Purification Technology 40 (1),61–67.

32.Fernandez, Y., Maranon, E., Castrillon, L., Vazquez, I., 2005. Removal ofCd and Zn from inorganic industrial waste leachate by ion exchange.Journal of HazardousMaterials 126 (1–3), 169–175.

33.Freundlich, H., 1926. Adsorption. Journal of Physical Chemistry 7, 57–64.Friberg, L.I., 1985. Rationale of biological monitoring of chemicals – withspecial reference to metals. American Industrial Hygiene AssociationJournal 46 (11), 633–642.

34.Gaballah, I., Kilbertus, G., 1998. Recovery of heavy metal ions throughdecontamination of synthetic solutions and industrial effluents usingmodified barks. Journal of Geochemical Exploration 62 (1–3), 241–286.

35.Gaey, M., Marchetti, V., Clement, A., Loubinoux, B., Gerardin, P., 2000.Decontamination of synthetic solutions containing heavy metals usingchemically modified sawdusts bearingpolyacrylic acid chains. Journalof Wood Science 46, 331–333.

36.Guclu, G., Gurdag, G., Ozgumus, S., 2003. Competitive removal of heavymetal ions by cellulose graft copolymers. Journal of Applied Polymer Science 90, 2034–2039.

37.Guibal, E., 2004. Interactions of metal ions with chitosan-based sorbents: a review. Separation and Purification Technology 38 (1), 43–74.

38.Gupta, K.C., Khandehar, K., 2002. Graft copolymerisation of acrylamide–methylacrylate comonomers onto cellulose using ceric ammoniumnitrate. Journal of Applied Polymer Science 86, 2631–2642.

39.Gurdag, G., Yasar, M., Gurkaynak, M.A., 1997. Graft copolymerization of acrylic acid on cellulose. Journal of Applied Polymer Science 66, 929–934.

40.Hancock, R.D., Martell, A.E., 1989. Ligand design for selective complexation f metal ions in aqueous solution. Chemical Reviews 89, 1875–1914.

41.Harris, J.A., Arthur, J.C., Carra, J.H., 1978. Photoinitiated polymerization of glycidyl methacrylate with cotton cellulose. Journal of Applied, Polymer Science 22, 905–915.

42.Hashem, A., 2006. Amidoximated sunflower stalks (ASFS) as a newadsorbent for removal of Cu(II) from aqueous solution. Polymer–Plastics Technology and Engineering 45, 35–42.

Hilal, N., Al-Khatib, L., Atkin, B.P., Kochkodan, V., Potapchenko, N.,2003. Photochemical modification of membrane surfaces for (bio)foulingreduction: a nano-scale study using AFM. Desalination and theEnvironment: Fresh Water for All 158 (1–3), 65–72.

43.Humans, vol. 49, Chromium, Nickel and Welding. InternationalAgency for Research on Cancer, Lyon.

44.Kamel, S., Hassan, E.M., El-Sakhawy, M., 2006. Preparation and application of acrylonitrilegrafted cyanoethyl cellulose for theremoval of copper(II) ions. Journal of Applied Polymer Science 100,329–334.

45.Kelleher, B., 2001. The development of alternative adsorbents for organiccompounds in aqueous environments. Ph.D. Thesis, University ofLimerick, Ireland, p. 194Khan, F., Ahmad, S.R., Kronfli, E., 2003. UV-radiation induced preirradiationgrafting of methyl methacrylate onto lignocellulose fibers inan aqueous medium and characterisation. Journal of Applied Polymer

Science 91 (3), 1667–1675.

46.Kjeldson, P., Barlaz, M.A., Rooker, A.P., Baun, A., Ledin, A., Christensen, T.H., 2002. Present and long term composition of MSW landfillleachate: a review. Critical Reviews in Environmental Science and Technology 32, 297–336

47.Klemm, D., Schmauder, H.P., Heinze, T., 2002. Cellulose. In: De Baets,S., Vandamme, E.J., Steinbuchel, A. (Eds.), Polysaccharides II.Polysaccharides from Eukaryotes, vol. 6. Wiley-VCH,Weinheim, pp.275–320.

48.Klemm, D., Heublein, B., Fink, H.F., Bohn, A., 2005. Cellulose:fascinating biopolymer and sustainable raw material. AngewandteChemie International Edition 44, 3358–3393.

49.Kondo, T., Ishizu, A., Nakano, J., 1989. Preparation of glycidyl cellulosesfrom completely allylated methylcellulose and Tri-O-allylcellulose.Journal of Applied Polymer Science 37, 3003–3009

50.Kongsricharoern, N., Polprasert, C., 1996. Chromium removal by abipolar electro-chemical precipitation process. Water Science andTechnology 34 (9), 109–116.

51.Kongsricharoern, N., Polprasert, C., 1995. Electrochemical precipitation of chromium Cr(VI) from an electroplating wastewater. Water Scienceand Technology 31 (9), 109–117.

52.Kubota, H., Suzuki, S., 1995. Comparative examinations of reactivity ofgrafted celluloses prepared by ultra violet and ceric salt-initiatedgraftings. European Polymer Journal 31 (8), 701–704.

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Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

General Parameters

Method : Iron (Fe) Autosampler : None Use SFI: No Operator : Jyothi

Instrument Mode: Flame Dilution: None

mA Hours

n/a

Report Date: 11-06-2022 17:40:20

Valid Method Signatures

11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Method Audit Trail

11-06-2022 17:22:18 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC

Signed with Reason : Approved by:

Analysis Details

Spectrometer: iCE 3000 AA01204906 v1.30

Analysis Name: Iron (Fe) 11-06-2022 Operator Name: Jyothi

n/a

Lamp Information Serial Number

Element(s) Fe

. . .

Deuterium Lamp Hours: 68.34

Sequence Table

Shared Standards: Yes	
Action	Fe
Calibration	√
B.No.Group-II	√
B.No.Group-III	√
B.No.Group-IV	√
B.No.Group-V	√
B.No.Group-VI	✓

Sample Details

		Nominal Mass: 1.0000			
No.	Sample Id	Sample Mass	Dilution Ratio		
1	B.No.Group-II	1.0000	1.0000		
2	B.No.Group-III	1.0000	1.0000		
3	B.No.Group-IV	1.0000	1.0000		
4	B.No.Group-V	1.0000	1.0000		
5	B.No.Group-VI	1.0000	1.0000		

Valid Analysis Signatures

11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Analysis Audit Trail

- 11-06-2022 17:30:16 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
- 11-06-2022 17:38:27 Jyothi(M. Jyothi):DESKTOP-39TDEGC Error MD147 - Activity manually aborted by user.
- 11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
- 11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

MART SPECIALITIES LAB LLP.

Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Spectrometer Parameters - Fe

Element: Fe Wavelength: 248.3nm Background Correction: D2 Signal Type: Continuous Measurement Time: 4.0secs Use RSD Test: No

Flame Type: Air-C2H2 Nebuliser Uptake: 4secs Burner Height: 7.0mm

Sampling: None

Calibration Mode: Normal	
Concentration Units: mg/L	
Acceptable Fit: 0.990	
Standard 1	2.0000
Standard 2	4.0000
Standard 3	6.0000

No changes are recorded for this element

Measurement Mode: Absorbance Bandpass: 0.2nm High Resolution: Off Resamples: Fast Flier Mode: No

Lamp Current: 75% Optimise Spectrometer Parameters: No Number Of Resamples: 3

Flame Parameters - Fe

Fuel Flow: 0.9L/min Burner Stabilisation: 0mins Optimise Burner Height: No Auxilary Oxidant: Off Optimise Fuel Flow: No

Sampling Parameters - Fe

Calibration Parameters - Fe

Line Fit: Linear Scaled Units: mg/L Rescale Limit: 10.0% Standard 4 Standard 5 Use Stored Calibration: No Scaling Factor: 1.0000 Failure Action: Flag and Continue 8.0000 10.0000

Element Audit Trail - Fe

Solution Results - Fe

Y = 0.04924x + 0.0141 Fit: 0.9965 Characteristic Conc: 0.0894

> 1 2 3

> 1 2 3

> 1 2 3

> 1 2 3

1 2

3

Sample ID

Fe Blank

Fe Standard 1

Fe Standard 2

Fe Standard 3

Fe Standard 4

	4				Norma	I: Linea	ar Fit				
	Abs		*		*		*		-*-		*
	0 * -0-0	1	2	3	4	5	6	7	8	9	10
	-0-0		2	0		nc:mg/		,	0	0	10
Signal	Rsd	Conc				cted C					
Abs	%	mg/L			mg/L						
0.001	35.6	0.000)		Ũ						
0.001	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:26	i		
0.001	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:30)		
0.000	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:35			
0.120	0.3	2.0000)								
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:05			
0.119	-	ound: 0.00						7:32:09			
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:13			
0.215	0.3	4.0000)								
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:40)		
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:45			
0.216	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:49)		
0.322	0.2	6.0000)								
0.323	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:19)		
0.322	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:23			
0.322	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:27			
0.411	0.3	8.0000)								
0.410	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:59)		
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:03			
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:07			

MART SPECIALITIES LAB LLP.

Operator Name: Jyothi

Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Solution Results - Fe

Sample ID	Signal	Rsd	Conc	Corrected Conc
-	Abs	%	mg/L	mg/L
Fe Standard 5	0.493	0.1	10.0000	
1	0.494	Backg	ground: 0.005	11-06-2022 17:34:38
2	0.494	Backg	ground: 0.005	11-06-2022 17:34:42
3	0.493	Backg	ground: 0.006	11-06-2022 17:34:46
Fe B.No.Group-II	0.834	0.1	16.6490 C	16.6490 C
1	0.834	Backg	ground: 0.005	11-06-2022 17:35:14
2	0.833	Backg	ground: 0.005	11-06-2022 17:35:19
3	0.834	Backg	ground: 0.005	11-06-2022 17:35:23
Fe B.No.Group-III	0.759	0.1	15.1286 C	15.1286 C
1	0.758	Backg	ground: 0.003	11-06-2022 17:35:52
2	0.760	Backg	ground: 0.003	11-06-2022 17:35:56
3	0.758	Backg	ground: 0.004	11-06-2022 17:36:00
Fe B.No.Group-IV	0.645	0.3	12.8189 C	12.8189 C
1	0.647	Backg	ground: 0.003	11-06-2022 17:36:31
2	0.644	Backg	ground: 0.003	11-06-2022 17:36:35
3	0.645	Backg	ground: 0.003	11-06-2022 17:36:40
Fe B.No.Group-V	0.809	0.2	16.1488 C	16.1488 C
1	0.808	Backg	ground: 0.005	11-06-2022 17:37:11
2	0.809	Backg	ground: 0.005	11-06-2022 17:37:15
3	0.811	Backg	ground: 0.005	11-06-2022 17:37:20
Fe B.No.Group-VI	0.711	0.3	14.1597 C	14.1597 C
1	0.709	Backg	ground: 0.005	11-06-2022 17:37:55
2	0.713	Backg	ground: 0.005	11-06-2022 17:37:59
3	0.712	Backg	ground: 0.004	11-06-2022 17:38:03

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-1 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
I	DETAILS OF THE SAMPLE
Sample Name : NA	
Name of the Manufacturer : NA	
Storage condition To b	up-I Mfg. : NA Exp. : NA e stored at room date date
	Derature Batch : NA C±3°C size
	by AAS Analysis.
	6/2022
Analysis Completion Date : 11/0	6/2022
Mfg. License No. : Not	provided
A.R.NO : NA	The American Company

Test Results

S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	620ppm	

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



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	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00	
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No.:MSL/2022/JUNE/TARA/0064-2Sample Received Date:11/06/2022Report Date:13/06/2022	
Q	DETAILS OF THE SAMPLE	
Sample Name : N	A	
Name of the Manufacturer : N	A	
Storage condition : To	roup-II Mfg. : NA Exp. : NA be stored at room date date	
Room Temperature : 25	Batch : NA size	
Tests Required:IreMethod:NAnalysis Starting Date:11	on by AAS Analysis.	
	ot provided	
A.R.NO : N	A	

Test Results

S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	832.5ppm	

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



OCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinaga Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Sample Received Date : 11/06/2022
a.y.	DETAILS OF THE SAMPLE
Sample Name : N	JA
Name of the Manufacturer : N	JA
Storage condition	Group-III Mfg. : NA Exp. : NA To be stored at room date date Cemperature Batch : NA
	5°C±3°C size
	gm /
Method : I Analysis Startin <mark>g Date : I</mark>	ron by AAS Analysis. NA 1/06/2022 1/06/2022
and the second sec	Not provided
A.R.NO	NA
Remark : S	Sample analyzed as received

Test Results

S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	756.5ppm	

×

Authorized Signatory

(Dr.R.Marayya)

Office: 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile: 98481 98350, 98481 25019 Mail: rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College	
Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000	Sample Received Date: 11/06/2022Report Date: 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name : N	A
Name of the Manufacturer : N	A
Storage condition : T	roup-IV Mfg. : NA Exp. : NA o be stored at room date date emperature Batch : NA
	5°C±3°C size
Method : N Analysis Starting Date : 1	on by AAS Analysis. A 1/06/2022 1/06/2022
Mfg. License No: N	ot provided
A.R.NO : N	A
Remark : Sa	ample analyzed as received

Test Results

and the second se		for some of	
S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	641ppm	

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G				
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00			
Name & Address of the Custome	r: Reference / Report No. : MSL/2022/JUNE/TARA/0064-5			
Tara Government College	Sample Received Date : 11/06/2022			
Prashanth Nagar Colony, Balajinag	ar Sumpto Received 2 mil			
Sangareddy Telangana. 502000	Report Date : 13/06/2022			
Contact Person: Dr. Abhijeet Contact Number :9502344392				
	DETAILS OF THE SAMPLE			
Sample Name :	NA			
Name of the Manufacturer :	NA			
Batch no : Storage condition	Group-VMfg. : NAExp. : NATo be stored at roomdatedateTemperatureBatch : NA			
Room Temperature :	25°C±3°C size			
Quantity Received	4gm			
Tests Required:Method:Analysis Starting Date:	Iron by AAS Analysis. NA 11/06/2022			
Analysis Completion Date :	11/06/2022			
Mfg. License No.	Not provided NA			
A.R.NO Remark	Sample analyzed as received			

Test Results

S.No.	Test Parameter	Result
01. Ir	on by AAS Analysis: (ppm)	807.5ppm

>

Authorized Signatory

(Dr.R.Marayya)

Office : 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile : 98481 98350, 98481 25019 Mail : rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-6 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
1	DETAILS OF THE SAMPLE
Storage condition	up-V1 Mfg. : NA Exp. : NA be stored at room date date perature Batch : NA C±3°C size
Tests Required:IronMethod:NAAnalysis Starting Date:11/0Analysis Completion Date:11/0Mfg. License No.:NotA.R.NO:NA	by AAS Analysis. 6/2022 6/2022 provided

Test Results

S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	708ppm	

*

Authorized Signatory

(Dr.R.Marayya)

ADSORPTION STUDIES OF Fe[III] ON BENTONITE IN THE PRESENCE OF D-RIBOSE AS CHELATING AGENT

Dissertation submitted in Partial fulfillment for the requirements for the award of degree of

Bachelor of Science ín CHEMISTRY

By

1.T. Chandana, B.Sc(Chemistry), III Year 2.T. Pranay kumar, B.Sc(Chemistry), III Year 3.T. Shiva charan, B.Sc(Chemistry), III Year 4.N. Sai mahesh, B.Sc(Chemistry), III Year 5.T. Raghavendra, B.Sc(Chemistry), III Year

Under the Guídance:

K.SREEDHAR Assistant Professor DEPARTMENT OF CHEMISTRY



Tara Government College, Sangareddy(A) 2021-22

<u>CERTIFICATE</u>

This is to certify that the project work entitled "ADSORPTION STUDIES OF Fe[III] ON BENTONITE IN THE PRESENCE OF D-RIBOSE AS CHELATING AGENT" is presented by B.Sc (CHEMISTRY) students in partial fulfillment of the requirements for the degree of Bachelor of Science in Chemistry by the Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

The results embodied in this report have not been to any other University or Institution for the award of any degree.

(K.SREEDHAR)

Asst. Professor, Department of Chemistry Tara Government College, Sangareddy (A)

EXTERNAL EVALUATOR

ACKNOWLEDGEMENTS

We express my deep gratitude to my research supervisor **K.Sreedhar**, Asst. Professor, Department of Chemistry, Tara Govt. College, Sangareddy(A)-502001, INDIA for his inspiring guidance during the course of the Project work. The continuous encouragement extended by him propelled me to update my research skills and pedigree to engage in fruitful research.

We wish to express our gratitude to Smt. M.Praveena, Principal, Tara Govt. College, Sangareddy(A)-502001, INDIA for her constant support, cooperation and suggestions during the research work. We wish to express our sincere thanks to **Dr.K.Abhíjít**, Head, Department of Chemistry, Tara Govt. College, Sangareddy(A)-502001, INDIA for providing me facilities, help and support for the entire research work.

We express our special thanks to **MART Specialities Lab, Hyderabad** for providing technical assistance in Atomic absorption spectrometric analysis.

Finally, we take this opportunity to thank one and all that has directly or indirectly helped me in completing the task.

- 1.T. Chandana,
- 2.T. Pranay kumar,
- 3.T. Shiva charan,
- 4.N. Sai mahesh,
- 5.T. Raghavendra,

B.Sc(Chemistry), III Year B.Sc(Chemistry), III Year B.Sc(Chemistry), III Year

- B.Sc(Chemistry), III Year
- B.Sc(Chemistry), III Year

<u>DECLARATION</u>

We hereby declare that the project report entitled "ADSORPTION STUDIES OF Fe[III] ON BENTONITE IN THE PRESENCE OF D-RIBOSE AS CHELATING AGENT" is the work done by us in the campus at Department of Chemistry, Tara Government College, Sangareddy(A) during the academic year 2021-2022 and is submitted in partial fulfillment of the requirements for the degree of Bachelor of Science by Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

S.No.	Name of the Student	Roll Number	Group	Year	Signature
1	T.CHANDANA	6058-19-578-032	B.Sc(MCCs)	III	
2	T.PRANAY KUMAR	6058-19-578-033	B.Sc(MCCs)	III	
3	T.SHIVA CHARAN	6058-19-578-034	B.Sc(MCCs)	III	
4	N.SAI MAHESH	6058-19-578-026	B.Sc(MCCs)	III	
5	T.RAGHAVENDRA	6058-19-578-035	B.Sc(MCCs)	III	

TARA GOVERNMENT COLLEGE, SANGAREDDY – 502 001 (AUTONOMOUS)

BONAFIDE CERTIFICATE

Certified that the project report" ADSORPTION STUDIES OF Fe[III] ON BENTONITE IN THE PRESENCE OF D-RIBOSE AS CHELATING AGENT" is the bonafide work of

1.Т.	Chandana,	B.Sc(Chemistry),	III	Year
2.т.	Pranay kumar,	B.Sc(Chemistry),	III	Year
З.Т.	Shiva charan,	B.Sc(Chemistry),	III	Year
4.N.	Sai mahesh,	B.Sc(Chemistry),	III	Year
5.т.	Raghavendra,	B.Sc(Chemistry),	III	Year

who carried out the project work under my supervision.

K.Sreedhar **PROJECT SUPERVISOR** Dr.K.ABHIJIT HEAD, DEPARTMENT OF CHEMISTRY

Smt. M.PRAVEENA PRINCIPAL

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Chapter-I

INTRODUCTION

ADSORPTION

Adsorption is a process in which solids come into contact with liquids or gases, and the mass transfer occurs from liquids to solids. Desorption is the reversal of this action. Adsorption operations take advantage of a solid's capacity to concentrate certain chemicals from a fluid on to its surface. Adsorbate refers to the adsorbed substance, while adsorbent refers to the solid substance. The following are some examples of solid-liquid and solid-gas applications:

- Removing dissolved moisture from gasoline.
- Decolorization of sugar solutions and petroleum products.
- Removing noxious odors and tastes from water. Dehumidification of air and gases is one of the solid-gas activities.
- To fractionate mixtures of hydrocarbon gases such as methane, ethane, and propane.
- To remove undesirable odors and contaminants from gases.
- To recover valuable solvent vapors from dilute gas mixtures.

NATURE OF ADSORBENTS

Adsorbents are typically granular in nature, ranging in size from 0.5 mm to 12 mm. They can't have a lot of pressure decrease or get swept away by a fast-moving stream. During handling, they must maintain their shape and size. They'll need a lot of pores and a lot of surface area per unit mass.

Some of the commonly used adsorbents, their sources and applications are given below:

Sl. No.	Adsorbent	Source	Application
1.	Fuller's earth	Naturally occurring clay is heated and dried to get a porous structure.	De-colorizing, drying of lubricating oils, kerosene and engine oils.
2.	Activated charcoal	Bentonite or other activated clay which are activated by treatment with sulfuric acid and further washing, drying and crushing.	Used for de-colorizing petroleum products.
3.	Bauxite	A naturally occurring hydrated alumina, activated by heating at 230-815	Used for de-colorizing petroleum products and for drying gases.

4.	Alumina	A hard hydrated aluminum oxide, which is activated by heating to drive off the moisture and then crushed to desired size.	Used as desiccant.
5.	Bone-char	Obtained by destructive distillation of crushed bones at 600-900	Used for refining sugar and can be reused after washing and burning.
6.	Silica gel	A hard granular and porous product obtained from sodium silicate solution after treatment with acid. Normally has 4 to 7% water in the product.	Dehydration of gases and liquids, and separation of gas-liquid hydrocarbon mixture.

7.	Activated carbon	 (1) Vegetable matter is mixed with calcium chloride, carbonized and finally the inorganic compounds are leached away. (2) Organic matter is mixed with porous pumice stones and then heated and carbonized to deposit the carbonaceous matter throughout the porous particle. (3) Carbonizing substances like wood, sawdust, coconut shells, fruit pits, coal, lignite and subsequent activation with hot air steam. It is available in granular or pelleted form. 	De-colorizing of sugar solutions, chemicals, drugs, water purification, refining of vegetable and animal oils, recovery of gold and silver from cyanide ore-leach solution, recovery of solvent vapor from gas-mixtures, collection of gasoline hydro- carbons from natural gas, fractionation of hydrocarbon gases.
8.	Molecular sieves	These are porous synthetic zeolite crystals, metal alumino-silicates.	Dehydration of gases and liquids, and separation of gas- liquid hydrocarbon mixture.

SIGNIFICANCE OF RESEARCH PROBLEM

Adsorption of heavy metals is an important strategy to develop newer remediation technologies for the sustainable environmental protection. But the efficacy of adsorption of heavy metals under the normal conditions using suitable adsorbent depends on several factors which need to be finely tuned to get efficient adsorption process. In the presence of proper facilitating agents, the adsorption of heavy metals enhanced which will certainly improves the existing heavy metal techniques.

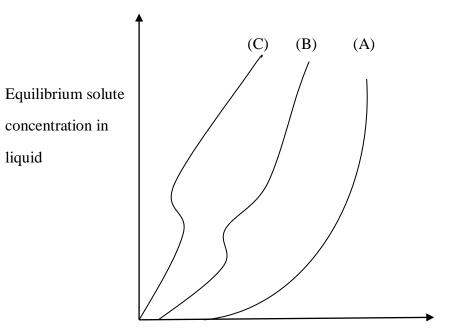
OBJECTIVE OF THE PROJECT

To evaluate the impact of D-Ribose as a chelating agent in the adsorption of Fe (III) from aqueous solution by Bentonite as an adsorbent to develop efficient remediation technology using concept of Coordination chemistry.

Chapter-II

ADSORPTION OF SOLUTES FROM DILUTE SOLUTIONS

Both the solvent and the solute are adsorbed whenever a mixture of solute and solvent is adsorbed using an adsorbent. As a result, only relative or apparent solute adsorption can be determined. As a result, treating a known volume of solution of original concentration Cwith a known weight of adsorbent is standard procedure. Let C* be the solution's final equilibrium solute concentration. If v is the volume of solution per unit mass of adsorbent (cc/g), and C and C* are the starting and equilibrium concentrations (g/cc) of the solute, then the apparent adsorption of the solute per unit mass of adsorbent is $v(C - C^*)$, (g/g), neglecting any volume change. This statement is most useful in the case of dilute solutions. The C* value is determined by the temperature, nature, and properties of the adsorbent when the proportion of the original solvent that can be adsorbed is tiny. The Freundlich adsorption isotherm, $C^*=K[v(C - C^*)]n$, represents the adsorption phenomena in dilute fluids across a small concentration range. The Freundlich adsorption equation is especially useful in situations where the identification of the solute is unknown, such as the removal of coloring substances from sugar solutions, oils, and other liquids. A spectrophotometer or colorimeter can quickly determine the color composition of the solute. In worked example 2, the interpretation of this data is demonstrated. Adsorption is good if the value of n is high, say 2 to 10. If it's between 1 and 2, it's relatively challenging, and if it's less than 1, it's easy and it indicates poor adsorption characteristics. Freundlich adsorption equation is also useful in such a case where the actual identification of solute is not known, e.g., removal of coloring substance from sugar solutions, oils etc. A typical adsorption isothermal for the adsorption of various adsorbents A, B and C in dilute solution at the same temperature for the same adsorbent is represented in a graph.



Kg solute apparently adsorbed /kg adsorbent

Adsorption isotherms for various adsorbents

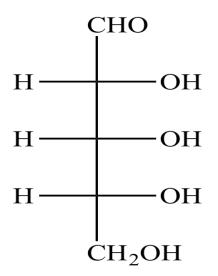
METAL COMPLEXATION OF A D- RIBOSE- BASED LIGAND

The complexation characteristics of methyl 2, 3, 4-tri-O-(2-picolyl)—D-ribopyranoside, a novel sugar-derived hexadentate ligand, were investigated using a combination of experimental and theoretical approaches. In the complexes with Mn^{II}, Co^{II}, Ni^{II}, and Zn^{II}, the coordination bond lengths reveal significant departures from perfect octahedral, with deformation towards trigonal-prismatic geometries, indicating a conformationally constrained ligand. The metal-cation–ligand interactions for D-Ribose and its acyclic analogue ligand [1, 2, 3-tri-O-(2-picolyl)-1, 2, 3-propanetriol] were investigated using spectroscopic techniques and isothermal calorimetric titrations for the series Mn^{II}, Co^{II}, Ni^{II}, Zn^{II}, and Cu^{II}. Depending on the nature of the metal, the results show that the complexes produced with D-Ribose are more stable than those obtained with According to molecular modelling studies; the presence of the sugar moiety greatly favours conformations that are compatible with metal binding. In many ways, a close comparision can be sketched between metal complexation and other host-guest interactions. To describe them both, several molecular variables have to be taken into consideration. One of the most

complex variables is the relation between the conformational flexibility of the ligand and the binding strength of the metal cation. For ligands with a reduced number of low-energy conformations, relatively straightforward estimations of the binding same wavelength of metal cations as well as a direct interaction between structural and thermodynamic data are available. However, for ligands with a huge number of degrees of freedom, the correct orientation of the chelating groups becomes one of the most important issues in metal-ion identification processes. For coordination chemists, the prediction and understanding about coordination properties of new ligands is one the most challenging tasks. We decided to investigate how entropic and enthalpic variables can impact on metal binding and selectivity for systems with a substantial number of degrees of freedom.

FERRIC COMPLEXES OF SUGAR-TYPE LIGAND

Fe (III) complexes formed with sugar-type ligands such as aldoses, ketoses, polyalcohols, sugar acids, di and trisaccharides was prepared. These complexes composition was determined by standard analytical methods. Deprotonated alcoholic hydroxy groups participate in the complex formation which results in the formation of polynuclear species. Mossbauer spectra reflected the presence of high spin iron (III) central atoms.EPR spectra showed antiferromagnetic interactions between the iron(III) centres in the complexes indicating dimeric or oligomeric complex structures. Depending upon the nature of a ligand as well as on the preparation mode of the complexes the ratio of interacting and isolated iron(III) is calculated. Formation of polynuclear iron(III) complexes of D-fructose, sorbose, lactobionic acid, glucose, galactose, mannose, and lactose were shown by the analysis and structural study by the sugar type ligand with ferric system. Ferric complexes of reducing sugars also contained ferrous species in some cases. The combination of Mossbauer spectroscopic measurements with ERP or magnetic susceptibility studies has led valuable information concerning the intramolecular and electronic structure on which the system has been developed.



Structure of D-Ribose

BENTONITE AS ADSORBENT

Because of heavy metals toxicity and non-biodegradable nature, the advent of heavy metals in water is turning into a critical environmental and public fitness concern. A variety of technology were evolved to do away with poisonous heavy metals from wastewater. The maximum crucial technology for the heavy steel ions elimination from wastewater consist of perception, ion exchange, adsorption, coagulation, evaporation and opposite osmosis. Adsorption on strong matrices has been proven to be an economically viable opportunity method (Abollino et al., 2003¹; Hoda et al., 2009² Kapoor and Vira Raghavan, 1998³). Cheaper Na-bentonite has been observed to be so beneficial for elimination of heavy steel ions from aqueous answers that it has attracted geologists and environmental engineers (Al-Quadbit et al., 2005⁴; Guo et al.,(2009).

The Na-bentonite from Gaomiaozi has been used as boundaries to save you infection of wastewater containing heavy metals. For this cause it's far crucial to take a look at the adsorption of metals of metals via way of means of Na-bentonite so one can offer crucial parameters and essential principle for the knowledge of adsorptive elimination of heavy metals via way of means of Na-bentonite from an aqueous environment. The bivalence ions of copper and nickel are not unusual

place observed withinside the commercial wastewater. The bivalence ions of copper and nickel are commonly found in the industrial wastewater. On a Na-montmorillonite turned into studied as feature of answer pH, dosage of Na-bentonite, temperature and make contact with and make contact with time. The aggressive isothermal Adsorption conduct of copper and nickel on Na-bentonite in singleaspect structures and binary-aspect structures has been investigated.



BENTONITE POWDER

Chapter-III

ATOMIC ABSORPTION SPECTROMETRY

Atomic absorption spectrometry (AAS) is an analytical technique that measures the concentrations of elements. Atomic absorption is so sensitive that it can measure down parts per billion of a gram (µg dm⁻³) in a sample. The technique makes use of the wavelengths of light specifically absorbed by an element. They correspond to the energies needed to promote electrons from one energy level to another, higher, energy level.

Atomic absorption spectrometry has many uses in different areas of chemistry.

Clinical analysis: Analysing metals in biological fluids such as blood and urine.

Environmental analysis: Monitoring our environment- eg finding out the levels of various elements in rivers, seawater, drinking water, air, petrol and drinks such as wine, beer and fruit drinks.

Pharmaceuticals: In some pharmaceutical manufacturing processes, minute quantities of a catalyst used in the process (usually a metal) are sometimes present in the final product. By using AAS the amount of catalyst present can be determined.

Industry: Many raw materials are examined and AAS is widely used to check that the major elements are present and that toxic impurities are lower than specified- eg in concrete, where calcium is a major constituent, the lead level should be low because it is toxic.

Mining: By using AAS the amount of metals such as gold in rocks can be determined to see whether it is worth mining the rocks to extract the gold.

HOW IT WORKS

Atoms of different elements absorb characteristic wavelengths of light. Analysing a sample to see if it contains a particular element means using light from that element. For example with lead, a lamp containing lead emits light from excited lead atoms that produce the right mix of wavelengths to be absorbed by any lead atoms from the sample. In AAS, the sample is atomized- i.e. converted into ground state free atoms in the vapour state- and a beam of electromagnetic radiation emitted from excited lead atoms is passed through the vaporized sample. Some of the radiation is absorbed by the lead atoms in the sample. The greater the number of atoms there is in the vapour, the more radiation is absorbed. The amount of light absorbed is proportional to the number of lead atoms. A calibration curve is constructed by running several samples of known lead concentration under the same conditions as the unknown. The amount the standard absorbs is compared with the calibration curve and this enables the calculation of the lead concentration in the unknown sample.

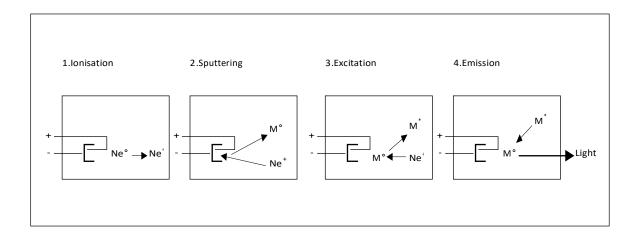
Consequently an atomic absorption spectrometer needs the following three components: a light source; a sample cell to produce gaseous atoms; and a means of measuring the specific light absorbed.

THE LIGHT SOURCE

The common source of light is a 'hollow cathode lamp'. This contains a tungsten anode and a cylindrical hollow cathode made of the element to be determined. These are sealed in a glass tube filled with an inert gas- e.g neon or argon- at a pressure of between 1 Nm⁻² and 5 Nm⁻².

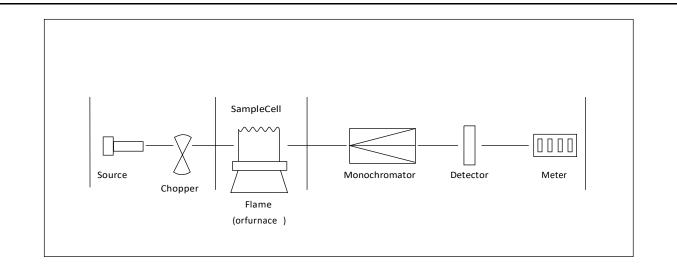


The ionization of some gas atoms occurs by applying a potential difference of about 300-400V between the anode and the cathode. These gaseous ions bombard the cathode and eject metal atoms from the cathode in a process called sputtering. Some sputtered atoms are in excited states and emit radiation characteristic of the metal as they fall back to the ground state $-egPb^* \rightarrow Pb + h v$. The shape of the cathode concentrates the radiation into a beam which passes through a quartz window, and the shape of the lamp is such that most of the sputtered atoms are redeposited on the cathode. A typical atomic absorption instrument holds several lamps each for a different element. The lamps are housed in a rotating turret so that the correct lamp can be quickly selected.



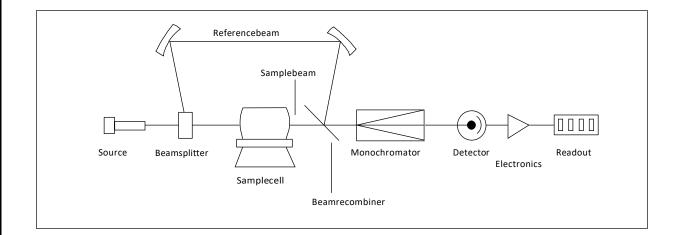
THE OPTICAL SYSTEM AND DETECTOR

A monochromator is used to select the specific wavelength of light –ie spectral line – which is absorbed by the sample, and to exclude other wavelengths. The selection of the specific light allows the determination of the selected element in the presence of others. The light selected by the monochromator is directed onto a detector that is typically a photomultiplier tube. This produces an electrical signal proportional to the light intensity



DOUBLE BEAM SPECTROMETERS

Modern spectrometers incorporate a beam splitter so that one part of the beam passes through the sample cell and the other is the reference. The intensity of the light source may not stay constant during an analysis. If only a single beam is used to pass through the atom cell, a blank reading containing no analyte (substance to be analysed) would have to be taken first, setting the absorbance at zero. If the intensity of the source changes by the time the sample is put in place, the measurement will be inaccurate. In the double beam instrument, there is a constant monitoring between the reference beam and the light source. To ensure that the spectrum does not suffer from loss of sensitivity, the beam splitter is designed so that as high a proportion as possible of the energy of the lamp beam passes through the sample.



ATOMIZATION OF THE SAMPLE

Two systems are commonly used to produce atoms from the sample. Aspiration involves sucking a solution of the sample into a flame; and electrothermal atomization is where a drop of sample is placed into a graphite tube that is then heated electrically.

Some instruments have both atomization systems but share one set of lamps. Once the appropriate lamp has been selected, it is pointed towards one or other atomization system.

FLAME ASPIRATION

Ethyne/air (giving a flame with a temperature of 2200–2400°C) or ethyne/dinitrogen oxide (2600–2800°C) are often used. A flexible capillary tube connects the solution to the nebulizer. At the tip of the capillary, the solution is 'nebulized' –ie broken into small drops. The larger drops fall out and drain off while smaller ones vaporize in the flame. Only ca 1% of the sample is nebulized.

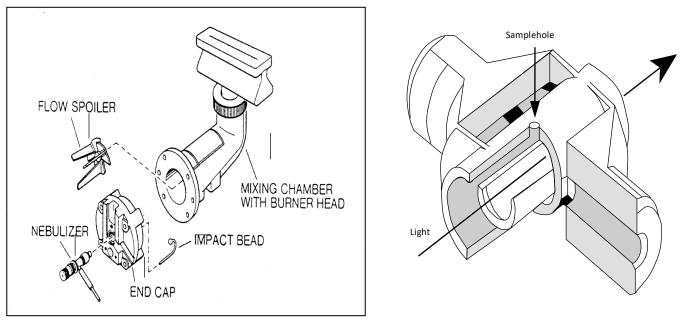


Figure 1

Figure 2

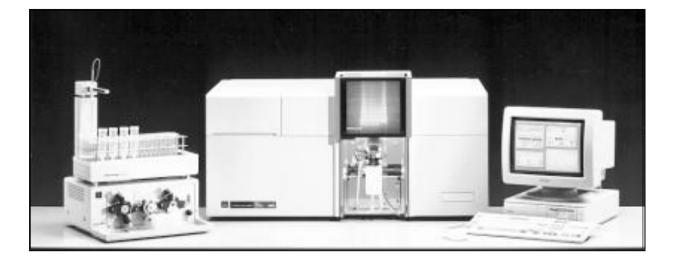
ELECTROTHERMAL ATOMIZATION

 $25 \ \mu$ l of sample (ca 1/100th of a raindrop) is placed through the sample hole and onto the platform from an automated micropipette and sample changer. The tube is heated electrically by passing a current through it in a pre-programmed series of steps. The details will vary with the sample but typically they

might be 30–40 seconds at 150°C to evaporate the solvent, 30 seconds at 600°C to drive off any volatile organic material and char the sample to ash, and with a very fast heating rate (ca 1500 °C s⁻¹) to 2000-2500°C for 5–10 seconds to vaporise and atomize elements (including the element being analysed). Finally heating the tube to a still higher temperature –ca 2700°C – cleans it ready for the next sample. During this heating cycle the graphite tube is flushed with argon gas to prevent the tube burning away. In electrothermal atomization almost 100% of the sample is atomised. This makes the technique much more sensitive than flame AAS.

SAMPLE PREPARATION

Sample preparation is often simple, and the chemical form of the element is usually unimportant. This is because atomization converts the sample into free atoms irrespective of its initial state. The sample is weighed and made into a solution by suitable dilution. Elements in biological fluids such as urine and blood are often measured simply after a dilution of the original sample.



When making reference solutions of the element under analysis, for calibration, the chemical environment of the sample should be matched as closely as possible –i.e., the analyte should be in the same compound and the same solvent. Teflon containers may be used when analysing very dilute Solutions because elements such as lead are sometimes leached out of glass vessels and can affect the results

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BACKGROUND ABSORPTION

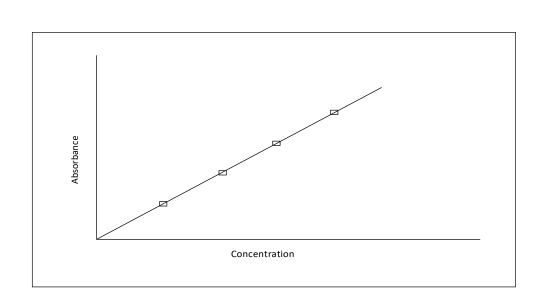
It is possible that other atoms or molecules apart from those of the element being determined will absorb or scatter some radiation from the light source. These species could include unvaporised solvent droplets, or compounds of the matrix (chemical species, such as anions, that tend to accompany the metals being analysed) that are not removed completely. This means that there is a background absorption as well as that of the sample.

One way of measuring and correcting this background absorption is to use two light sources, one of which is the hollow cathode lamp appropriate to the element being measured. The second light source is a deuterium lamp.

The deuterium lamp produces broad band radiation, not specific spectral lines as with a hollow cathode lamp. By alternating the measurements of the two light sources – generally at 50 - 100 Hz – the total absorption (absorption due to analyte atoms plus background) is measured with the specific light from the hollow cathode lamp and the background absorption is measured with the light from the deuterium lamp. Subtracting the background from the total absorption gives the absorption arising from only analyte atoms.

CALIBRATION

A calibration curve is used to determine the unknown concentration of an element -eg lead – in a solution. The instrument is calibrated using several solutions of known concentrations. A calibration curve is produced which is continually rescaled as more concentrated solutions are used – the more concentrated solutions absorb more radiation up to a certain absorbance. The calibration curve shows the concentration against the amount of radiation absorbed in the given figure. (a) The sample solution is fed into the instrument and the unknown concentration of the element-e.g., lead- is then displayed on the calibration curve given in the below figure. (b)





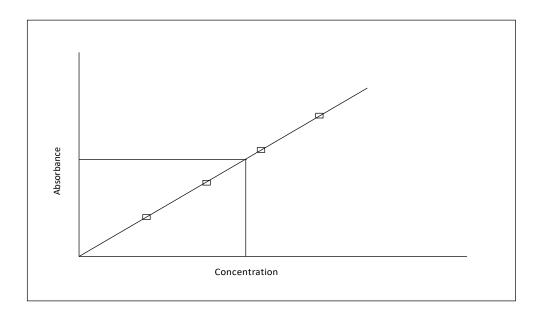


Figure (b)

Chapter-IV

METHODOLOGY

MATERIALS REQUIRED

- Ferric chloride Hexahydrate
- D-Ribose
- Bentonite
- Volumetric flask
- Digital Weighing Machine
- Watch glasses
- Hot air oven
- Ultra-pure water (Demineralized)
- 2% of Nitric acid

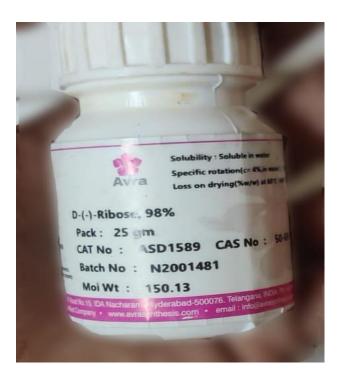
PROCEDURE

- Using a 250ml Volumetric flask prepare a metal ligand solution by adding 100ppm of Ferric chloride Hexahydrate (MOLYCHEM MCR-11580) and 200ppm of D-Ribose (AVRAN2200133).
 Prepare 250ml solution by adding Ultra-pure water and then keep this system aside for a few hours.
- Weigh 5 grams of Bentonite(AVRA N2101070) using a Digital weighing machine (Citizen Scales(I) PVT LTD CTG302-300) and take this into a beaker.
- **3.** Now add 100ml of the above prepared metal ligand solution into the beaker and stir the mixture well for 10 minutes using a glass rod.
- **4.** Keep this mixture aside for 48 hours without disturbing it as at this step Ferric is going to be adsorbed on Bentonite in the presence of D-Ribose which acts as a chelating agent.
- **5.** After completion of 48 hours take the mixture and filter it off using Whatman Grade 1 filter paper and a funnel.

- **6.** After filtration of the mixture again add Ultra-pure water for 3 times and then filtrate it to obtain pure concentration of Ferric which is get adsorbed on Bentonite.
- Collect the filtered Bentonite powder and place it on a watch glass and keep this in a Hot air Owen at 60 °C for 10 hours to get rid of moisture present in it.
- 8. Now weigh each 1 gm of Bentonite in glass vials.
- 9. Now take a beaker and rinse it with ultra-pure water then followed by Nitric acid.
- **10.** Take 0.5 grams of Bentonite sample in the beaker and add 2% of Nitric acid and stir the mixture well for 10-15 minutes.
- **11.** Filter the mixture using Whatman Grade 1 filter paper and again 3 times by using Ultra-pure water to obtain pure concentration of Ferric present in the mixture prepared using the sample.
- **12.** Take this collected sample solution and keep this system under AAS (Thermos Scientific iCE 3300)
- **13.** Calculate the concentration of Ferric adsorbed on Bentonite at different ppm levels. Observe the graph obtained and note down the readings of the result we obtained.
- 14. Same Experiment carried out without the interference/addition of Ligand i.e. D-Ribose for Control Experiment.



Bentonite Powder



D – Ribose



FERRIC CHLORIDE (HEXAHYDRATE) FeCL36H2O



Metal Ligand Solution



Watch Glass



Spatula



Digital Weighing Machine



Hot air Owen

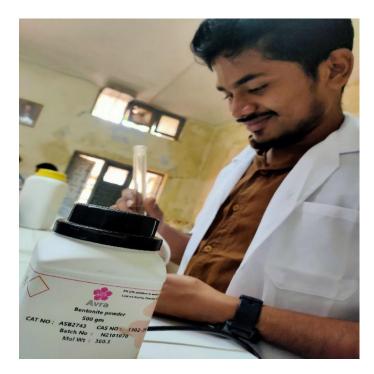










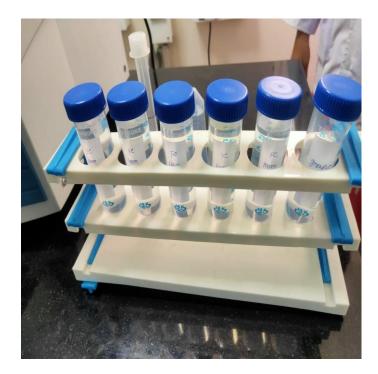








ATOMIC ASORPTION SPECTROSCOPY

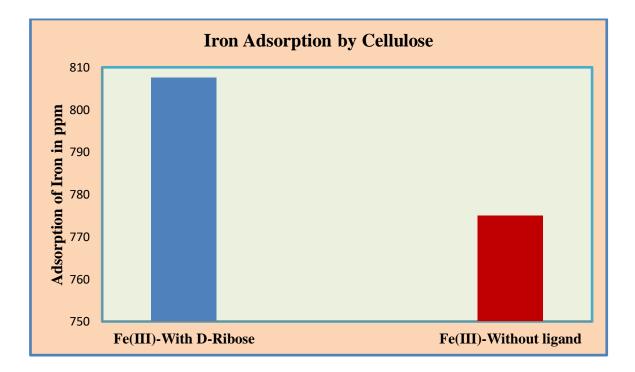


Fe standards with test sample

Chapter-V

RESULTS AND DISCUSSIONS

Bentonite adsorbs **807.5ppm** of Iron metal from aqueous solution of Fe(III)-D-Ribose metal ligand solution. Whereas, Bentonite adsorbs only **775.0ppm** when D-Ribose is absent. It is evident from the AAS results, ligand involvement enhanced the metal adsorption by initiating potential chemical interactions between adsorbate and adsorbent. D-Ribose firmly coordinates with Fe(III) to form a stable complex in aqueous condition. The complex coordination sphere in the resulted complex facilitates strong interactions with the polar points of Bentonite. From the AAS results, it is conclusive that **4.193%** of adsorbtion increased in the presence of D-Ribose as chelating agent.



Impact of D-Ribose on Adsorption of Fe (III) ions from aqueous solution by Bentonite.

<u>Spectrometer Parameters</u> – <u>Fe</u>:

Element : Fe	Measurement mode :	
	Absorbance	
Wavelength : 248.3nm	Band pass : 0.2nm	Lamp current : 75%
Background correction :	High Resolution : Off	Optimise Spectrometer
D2		Parameters : No
Signal type : continuous	Resamples : Fast	Number of resamples : 3
Measurement time :	Flier mode : No	
4.0secs		
Use RSD Test : No		

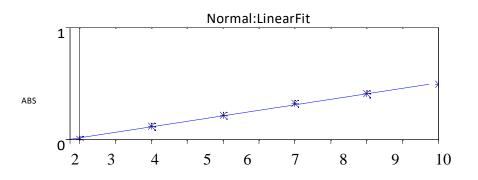
Flame parameters – Fe:

Flame type : Air – C2H2	Fuel Flow : 0.9L/min	Auxiliary Oxidant : Off
Nebuliser Uptake : 4secs	Bunsen Stabilisation : Omins	Optimise Fuel Flow : No
Burner Height :7.0mm	Optimise Burner fuel : No	

<u>Calibration Parameters</u> – <u>Fe</u>:

Calibration mode : Normal	Line fit : Linear	Use stored calibration : No
Concentration units : mg/L	Scales units : mg/L	Scaling factor : 1.0000
Acceptable fit : 0.990	Rescale Limits : 10.0%	Failure Action : Flag and Continue
Standard 1 - 2.0000	Standard 4 - 8.0000	
Standard 2 - 4.0000	Standard 5 - 10.0000	
Standard 3 - 6.0000		

Solutions Results-Fe:



Y = 0.04924x + 0.0141

Fit: 0.9965

Characteristic Conc.: 0.0894

Sample ID	Signal	RSD	Conc.	
	Abs	%	Mg/L	
Fe Blank	0	<99		0
	0		Background: 0.	000
	2 0		Background: 0.	000
	3 0		Background: 0.	000
Fe Standard 1	0.125		0.3	2
	0.125		Background: 0.	006
	2 0.125		Background: 0.	006
	3 0.126		Background: 0.	006
Fe Standard 2	0.225		0.5	4
	0.227		Background: 0.	007
	2 0.225		Background:0.0)07
	0.224		Background:0.0	007

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Fe Standard 3	0.336	0.2	6
1	0.336	Background: 0.008	
2	0.336	Background: 0.008 Background: 0.008	
3	0.335		
Fe Standard 4	0.425	0.2	
1	0.426	Background: 0.009	
2	0.425	Background: 0.008	
3	0.425	Background: 0.009	
Fe Standard 5	0.51	0.4	10
1	0.51	Background: 0.009	
2	0.509	Background: 0.009	
3	0.513	Background: 0.009	
Fe(III)-D-Ribose-Bentonite	0.809	0.2 16.1488 C	
1	0.808	Background: 0.005	
2	0.809	Background: 0.005	
3	0.811	Background: 0.005	
Fe(III)-Bentonite (Without Ligand)	0.776	0.2 15.4988C	
1	0.778	Background: 0.004	
2	0.775	Background: 0.004	
3	0.775	Background: 0.004	

TEST RESULTS:

S.No.	Test Parameters	Sample	Results
01.	Iron by AAS	Fe(III)- D-Ribose-	807.5ppm
	Analysis: (ppm)	Bentonite Sample	
02.	Iron by AAS	Fe(III)-Bentonite	775.0ppm
	Analysis: (ppm)	(control)	

CONCLUSION

From the current project it is clear that D-Ribose as a Chelating agent has played a vital role in adsorption of Fe(III) ions from aqueous solution and increases the adsorption up to **4.193** %. This aspect will be useful in designing the newer strategies of Heavy metal Remediation techniques using Natural Chelating Ligands as Facilitating agents in Metal Adsorption processes.

REFERENCES

- [1] A.Tzschach, K.Ponicke, L.Koreczand K.Burger, J. Organomet. Chem., 59, 199(1973).
- [2] A. D. Buckingham, Can. J. Chem. 1960, 38, 300-307
- [3] A.F.CottonandG.Wilkinson, AdvancedInorganicChemistry, 6thedition, JohnWiley&Sons, 1999, p.
 A. Martinez, J. Suarez, T. Shand, R. S. Magliozzo and R. A. Sanchez Delgado, J. Inorg. Biochem. 2011, 105, 39-45; S. S. Hosseini, M. Bhadbhade, R. J. Clarke, P. J. Rutledge and L. M. Rendina, Dalton Trans. 2011, 40, 506-513.
- [4] A. Bentz, P.Comba, R.J. Deeth, M. Kerscher, B. Seibold and H. Wadepohl, Inorg Chem. 2008,47,9518-9527;C.DiedrichandR.J.Deeth,Inorg.Chem.2008,47,2494-2506;L.Bonniard,A.delaLande, S. Ulmer, J.-P. Piquemal, O. Parisel and H. Gérard, Catalysis Today 2011, 177, 79-86; Y. D. M. Champouret, J.-D. Maréchal, I. Dadhiwala, J. Fawcett, D. Palmer, K. Singh and G. A. Solan, Dalton Trans. 2006, 2350-2361.
- [5] A. D. Becke, J. Chem. Phys. **1993**, 98,5648-5652.
- [6] B.Lakatos, I.Meisel, A.Rockenbauer, P.SimonandL.Korecz, Jnorg.Chim.Acta, 79, 269 (1983).
- B.Chiari, O.Piovesana, T.Tarantelliand P.F.Zanazzi, Jnorg. Chem., 22,2781(1983), and refs. There
 B. P. Hay and R. D. Hancock, Coord. Chem. Rev. 2001, 212,61-78.
- [8] C.Gessa, M.L.DeCherchi, A.Dessi, S.Deianaand G.Micera, Inorg. Chim. Acta, 80, L53(1983).
- [9] C. F. Macrae, P. R. Edgington, P. McCabe, E. Pidcock, G. P. Shields, R.Taylor, M. Towler and J. v. d. Streek, J. Appl. Cryst. 2006, 39,453-457
- [10]C. A. G. Haasnoot, F. A. A. M. de Leeuw and C. Altona, Tetrahedron **1980**, 36,2783-1792.C. A. Podlasek, J. Wu, W. A. Stripe, P. B. Bondo and A. S. Serianni, J. Am. Chem. Soc. **1995**, 117, 8635-8644.
- [11]C. Lee, W. Yang and R. G. Parr, Phys. Rev. B 1988, 37,785-789.

- [12] D. J. Cram and J. M. Cram, Container Molecules and their Guests, TheRoyal Society of Chemistry, Cambridge, 1994, p; J. B. Wittenberg and L. Isaacs in Complementarity and Preorganization, (Ed. J. W. S. Philip A. Gale), John Wiley and Sons, 2012.
- [13]D. Cremer and J. A. Pople, J. Am. Chem. Soc. 1975, 97,1354-1358.
- [14] D. E. Wilcox, Inorg. Chim. Acta 2008, 361, 857-867; A. Trapaidze, C. Hureau, W. Bal, M. Winterhalter and P. Faller, J. Biol. Inorg. Chem. 2012, 17,37-47.
- [15] D. W. Johnson and K. N. Raymond, Supramol. Chem. 2001, 13,639-659.
- [16] E. Chekmeneva, R. Prohens, J. M. Díaz-Cruza, C. Ariñoa and M. Esteban, Anal. Biochem. 2008, 375, 82-89.
- [17] F.W.Lichtenthaler, Acc.Chem.Res.2002,35,728-737; M.J.Climent, A.CormaandS.Iborra, Green Chem. 2011, 13, 520-540.
- [18] F. Bellot, R. Hardré, G. Pelosi, M. Thérisod and C. Policar, Chem. Commun. 2005,5414-5416.
- [19]F. Cisnetti, R. Guillot, G. Pelosi, M. Desmadril and C. Policar, Dalton Trans. 2007,1473-1476.
- [20]F. Cisnetti, R. Guillot, M. Thérisod and C. Policar, Acta Cryst. 2007, C63,m201-m203.
- [21] F.Cisnetti,R.Guillot,N.Ibrahim,F.Lambert,M.ThérisodandC.Policar,Carbohydr.Res.**2008**,343 ,530-535.
- [22] F. Cisnetti, R. Guillot, M. Thérisod, M. Desmadril and C. Policar, Inorg. Chem. 2008, 47,2243-2245.
- [23] F. E. Jorge, J. Autschbach and T. Ziegler, J. Am. Chem. Soc. 2005, 127, 975-985; E. I. Stiefel and G. F. Brown, Inorg. Chem. 1972, 11,434-436.
- [24] F. Mohamadi, N. G. J. Richard, W. C. Guida, L. R., M. Lipton, C. Caufield, G. Chang, T. Hendrickson and W. C. Still, J. Comput. Chem. 1990, 11,440-467.
- [25]G.Micera,S.Deiana,C.GessaandM.Petrera,Jnorg. Chim. Acta, 56, 109 (1981).
- [26]G. Charron, F. Bellot, F. Cisnetti, G. Pelosi, J.-N. Rebilly, E. Rivière, A.-L. Barra, T. Mallahand C. Policar, Chem. Eur. J. 2007, 13,2774-2782.
- [27] G. Camci-Unal and N. L. B. Pohl, Carbohydr. Polym. 2010, 81, 8-13; S. Striegler and M. G. Gichinga, Chem. Commun. 2008,5930-5932.
- [28] G. Sheldrick, SHELXS-97 Program for Crystal Structure Refinement, 1997.
- [29] G. Sheldrick, SHELXL-97 Program for Crystal Structure Refinement, 1997.
- [30]G. L. Strati, J. L. Willett and F. A. Momany, Carbohydr. Res. 2002, 337,1833-1849.
- [31] H. M. Irving and R. J. P. Williams, J. Chem. Soc. 1953,3192-3210.
- [32]H. D. Flack, Acta Cryst. 1983, A39, 876-881.
- [33]I.Zay, A.Vertes, G.TakacsiNagy, M.Subaand K.Burger, Radioanal. Nuc/. Chem., 88, 343(1985).
- [34]I. Berger, M. Hanif, A. A. Nazarov, C. G. Hartinger, R. O. John, M. L. Kuznetsov, M. Groessl, F.
 - Schmitt, O. Zava, F. Biba, V. B. Arion, M. Galanski, M. A. Jakupec, L. Juillerat-Jeanneret, P

J.Dyson and B. K. Keppler, Chem. Eur. J. 2008, 14, 9046-9057; M. Gottschaldt, U. S. Schubert, S.Rau, S. Yano, J.G. Vos, T. Kroll, J. Clement and I. Hilger, ChemBioChem2010, 11, 649-652; M. Hanif, S. M. Meier, W. Kandioller, A. Bytzek, M. Hejl, C. G. Hartinger, A. A. Nazarov, V. B. Arion, M. A. Jakupec, P. J. Dyson and B. K. Keppler, J. Inorg. Biochem. 2011, 105, 224-231; M. Hanif, A. A. Nazarov, C. G. Hartinger, W. Kandioller, M. A. Jakupec, V. B. Arion, P. J. Dyson and B. K. Keppler, Dalton Trans. 2010, 39, 7345-7352; J. K. H. Hui, Z. Yu, T. Mirfakhrai and M. J. MacLachlan, Chem. Eur. J. 2009, 15, 13456-13465; R. Kikkeri, D. Grünstein and P. H.Seeberger, J.2Am. Chem. Soc. 2010, 132, 10230-10232; M.-W. Louie,H.-W. Liu, M. H.-C. Lam, Y.-W. Lam and K. K.-W. Lo, Chem. Eur. J. 2011, 17, 8304-8308; Y. Mikata, S. Fujii, M. Naemura, K.Takahashi and Y. Noguchi, Dalton Trans. 2009, 10305-10310; T. Shibata,H.Hashimoto, I. Kinoshita, S. Yano and T. Nishioka, Dalton Trans. 2011, 40, 4826-4829; T. Storr, L. E. Scott, M. L. Bowen, D. E. Green, K. H. Thompson, H. J. Schugar and C. Orvig, Dalton Trans. 2010, 29, 5959-5971; E. C. Constable, C. E. Housecroft, M. Neuburger and P. Rosel, Chem. Commun. 2010, 46, 1628-1630.

- [35] In this Paper, solvation issues are not discussed, however as we consider two ligands with the same coordinating sites and their interactions with metal cations possessing the same charge, solvation effects could most confidently be neglected while discussing the metal selectivities in this study. ,Vol.
- [36]Ju.A.Komkov,M.A.Ivanov,G.A.ShagisultanovaandJu.B.Jakovlev,ZhNeorg.Khim.,2915202(1 984).
- [37]J.-H. Liao, C.-S. Chen, C.-C. Hu, W.-T. Chen, S.-P. Wang, I. L. Lin, Y.-H. Huang, M.-H. Tsai, T.-H. Wu, F.-Y. Huang and S.-H. Wu, Inorg. Chem. 2011, 50, 365-377; P. G. Devi, S. Pal, R. Banerjee and D. Dasgupta, J. Inorg. Biochem. 2007, 101,127-137.
- [38]J. Ali-Torres, J.-D. Maréchal, L. Rodriguez-Santiago and M. Sodupe, J. Am. Chem. Soc. 2011, 133, 15008-15014; V. Muñoz Robles, E. Ortega-Carrasco, E. G. Fuentes, A. Lledos and J.-D. Maréchal, Faraday Discuss. 2011, 148,137-159.
- [39] K.Burger, I.Zayand G. Takacsi Nagy, Jnorg. Chim. Acta, 80, 231 (1983).
- [40]K. Born, P. Comba, R. Ferrari, G. A. Lawrance and H. Wadepohl, Inorg. Chem. 2007, 46,458-464.
- [41] L. Garcia, S. Maisonneuve, J. Xie, R. Guillot, P. Dorlet, E. Rivière, M. Desmadril, F.Lambert and C. Policar, Inorg. Chem. 2010, 49,7282-7288.
- [42]L. Garcia, S. Maisonneuve, J. Oudinet-Sin Marcu, R. Guillot, F. Lambert, J. Xie and C. Policar, Inorg. Chem. 2011, 50,11353–11362.
- [43]L. J. Farrugia, J. Appl. Cryst. 1999, 32, 837-838.
- [44] M.Tonkovic, S.Music, I.Hadzija, J.Nagy-Czak6and A.Vertes, ActaChim.Hung., 110, 197(1982).

- [45]M. M. K. Boysen, Chem. Eur. J. 2007, 13, 8648-8659.
- [46] M.GottschaldtandU.S.Schubert, Chem.Eur.J.2009,15,1548-1557; M.L.BowenandC.Orvig, Chem. Commun. 2008, 5077-5091.
- [47]Chem. Eur. J. 2010, 16, 10202-10213; A. H. Younes, L. Zhang, R. J. Clark and L. Zhu, M. Ostermeier, M.-A. Berlin, R. M. Meudtner, S. Demeshko, F. Meyer, C. Limberg and S. Hecht, J. Org.Chem.2009,74,8761-8772;L.ZhangandL.Zhu,J.Org.Chem.2008,73,8321-8330;
 N.K.Dalley,U.Olsher, J. C. Lee, M. D. Eley, J. Wang and R. A. Bartsch, Tetrahedron 2007, 63, 10576-10580; L.Mikulášek,B.Grüner,C.Dordea,V.Rudzevich,V.Böhmer, J.Haddaoui,V. Hubscher-Bruder, F. Arnaud-Neu, J. Čáslavský and P. Selucký, European J. Inorg. Chem. 2007, 2007, 4772-4783; P. Comba, N. Dovalil, G. Haberhauer, G. Hanson, Y. Kato and T. Taura, J. Biol. Inorg. Chem. 2010, 15, 1129-1135.
- [48] M.Miguirditchian, D.Guillaneux, D.Guillaumont, P.Moisy, C.Madic, M.P.JensenandL.Nash, Inorg. Chem. 2005, 44, 1404-1412.M.J.Frisch, G.W.Trucks, H.B.Schlegel, G.E.Scuseria, M. A. Robb, J. R. Cheeseman, J.J. A. Montgomery, T. Vreven, K. N. Kudin, J. C. Burant, J. M. Millam, S. S. Iyengar, J. Tomasi, V. Barone, B. Mennucci, M. Cossi, G. Scalmani, N. Rega, G. A. Petersson, H. Nakatsuji, M. Hada, M. Ehara, K.Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, M. Klene, X. Li, J. E. Knox, H. P. Hratchian, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, P. Y. Ayala, K.Morokuma, G.A.Voth, P. Salvador, J. J. Dannenberg, V. G. Zakrzewski, S. Dapprich, A.D.Daniels, M.C.Strain O.Farkas, D. K. Malick, A. D. Rabuck, K. Raghavachari, J. B. Foresman, J.V.Ortiz, Q.Cui, A.G.Baboul, S. Clifford, J. Cioslowski, B. B. Stefanov, G. Liu, A. Liashenko, P. Piskorz, I.Komaromi R.L.Martin, D.J.Fox, T. Keith, M.A.Al-Laham, C.Y.Peng, A.NanayakkaraM.Challacombe, P.M.W.Gill, B.Johnson, W.Chen, M.W.Wong, C.G onzalez and J.A.Pople, Gaussian 03, Gaussian, Inc., Wallingford CT, 2004.
- [49] N. Ghaschghaie, T. Hoffmann, M. Steinborn and P. Klufers, *DaltonTrans.* 2010, 39, 5535-5543;
 A. L. Petrou, *Coord. Chem. Rev.* 2002, 228,153-162.
- [50]P.Saltman,].Chem.Educ.,42,682(1965).
- [51] P. J.Charley, B. Sarkar, C. F. Stitt and P. Saltman, Bio-chim. Biophys. Acta, 69, 313 (1963).
- [52]P. Comba, Coord. Chem. Rev. 1999, 182, 343-371; P. Comba and W. Schiek, Coord. Chem. Rev.2003, 238-239, 21-29.
- [53] P. Comba, T. W. Hambley and B. Martin, Molecular Modeling of inorganiccompounds, 3rd edition, Wiley VCH Weinheim, 2010, p.
- [54] P.Comba,Coord.Chem.Rev.1999,185186,8198;P.CombaandM.Kerscher,Coord.Chem.Rev.2009, 253, 564-574.
- [55]P. J. Hay and W. R. Wadt, J. Chem. Phys. 1985, 82, 299-310.

- [56] R. D. Hancock, D. L. Melton, J. M. Harrington, F. C. McDonald, R. T. Gephart, L.L.Boone, S.B. Jones,
- [57] N. E. Dean, J. R. Whitehead and G. M. Cockrell, Coord. Chem. Rev. 2007, 251,1678-1689.
- [58]R. Deeth and C. Diedrich, J. Biol. Inorg. Chem. 2009, 15, 117-129' N.Gresh G. A. Cisneros, T. A.

Darden and J. P. Piquemal, J. Chem. Theory Comput. 2007, 3, 1960-1986.

- [59] R. J. Deeth, A. Anastasi, C. Diedrich and K. Randell, Coord. Chem. Rev. 2009, 253,795-816.
- [60] R. D. Hancock and K. Hegetschweiler, J. Chem. Soc., Dalton Trans. 1993,2137-2140.
- [61] R. Jeanloz, H. G. Fletcher and C. S. Hudson, J. Am. Chem. Soc. 1948, 70,4052-4054.
- [62] R. D. Shannon, Acta Cryst. 1976, A32, 751-767.
- [63]S.WolowiecandK. Drabent, J.Radioanal.Nucl. Chem. Lett., 95, 1 (1985).
- [64] S.Dhungana, J.M.Harrington, P.Gebhardt, U.Möllmannand A.L.Crumbliss, Inorg. Chem. 2007, 46, 8362 -8371.
- [65] S. Woodward, M. Diéguez and O. Pàmies, Coord. Chem. Rev. 2010, 254,2007-2030.
- [66] S. J. Angyal, Tetrahedron 1974, 30, 1695-1702; S. J. Angyal, Chem. Soc. Rev. 1980, 9, 415-428;
 S. J.Angyal, Aust. J. Chem. 2000, 53, 567-570; B. Gyurcsick and L. Nagy, Coord. Chem. Rev. 2000, 203, 81- 149; N. Nagy and A. Szorcsik, J. Inorg. Biochem. 2002, 89, 1-12; J.-F. Verchèreand S. Chapelle, Progr. Inorg. Chem. 1998, 47, 837-945; D. M. Whitfield, S. Stojkovski and B. Sarkar, Coord. Chem. Rev. 1993, 122, 171-225.
- [67] S. J. Angyal, Adv. Carbohydr. Chem. Biochem. 1989, 47,1-43.
- [68] S. Bjelić and I. Jelesarov, J. Mol. Recognit. 2009, 21, 289-312; R. J. Falconer and B. M. Collins, J. Mol. Recognit. 2011, 24, 1-16; R. J. Falconer, A. Penkova, I. Jelesarov and B. M. Collins, J. Mol. Recognit. 2010, 23, 395-413; O. Okhrimenko and I. Jelesarov, J. Mol. Recognit. 2008, 21,1-19.
- [69] Schrodinger, Inc., Portland, Oregon, 2000,
- [70] T. K. Lindhorst, Essentials of Carbohydrate Chemistry and Biochemistry, Wiley-VCH, 2003, p.
- [71] T.Marino, N.Russo and M Toscano, J.Inorg. Biochem. 2000, 79, 179-185; J.Bertran, L. Rodriguez-Santiago and M. Sodupe, J. Phys. Chem. B 1999, 103, 2310-2317.
- [72] The use of unprotected sugar polyols or polyolate as ligands, while not discussed hereindetail, is conceptuallyrelatedtoourwork.Forareviewsee:T.Allscher,P.Klüfers,
- [73]P.Mayerin:B.Fraser-Reid,K. Tatsuta, J. Thiem(eds.) in Glycoscience,Spring Verlag, Berlin, 2008, 1079–1139., Vol.
- [74] V. J. James, J. D. Stevens and F. H. Moore, Acta Crist. 1978, B34,188-193.
- [75] W. O. Gillum, R. A. D. Wentworth and R. F. Childers, Inorg. Chem. 1970, 9,1825-1832.
- [76] W. Jorgensen, D. Maxwell and J. Tirado-Rives, J. Am. Chem. Soc. 1996, 118, 11225-11236.
- [77] W.J.Hehre, R.Ditchfield and J.A.Pople, J.Chem. Phys. 1972, 56, 2257-2261; P.C.Hariharan and
- [78] J. A. Pople, Theor. Chim. Acta 1973, 28, 213-222.

 $\label{eq:constraint} \ensuremath{\left[79\right]} Z. Jingkun, L. Rongehuan, W. Shuxin, X. Yuanfuand Q. Shouren, Kexeu Tongbao, 29, 1091 (1984).$

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Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

General Parameters

Method : Iron (Fe) Autosampler : None Use SFI: No Operator : Jyothi

Instrument Mode: Flame *Dilution:* None

mA Hours

n/a

Report Date: 11-06-2022 17:40:20

Valid Method Signatures

11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Method Audit Trail

11-06-2022 17:22:18 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC

Signed with Reason : Approved by:

Analysis Details

Spectrometer: iCE 3000 AA01204906 v1.30

Analysis Name: Iron (Fe) 11-06-2022 Operator Name: Jyothi

n/a

Lamp Information Serial Number

Element(s) Fe

. . .

Deuterium Lamp Hours: 68.34

Sequence Table

Shared Standards: Yes	
Action	Fe
Calibration	√
B.No.Group-II	√
B.No.Group-III	√
B.No.Group-IV	√
B.No.Group-V	√
B.No.Group-VI	✓

Sample Details

		Nominal Mass: 1.0000					
No.	Sample Id	Sample Mass	Dilution Ratio				
1	B.No.Group-II	1.0000	1.0000				
2	B.No.Group-III	1.0000	1.0000				
3	B.No.Group-IV	1.0000	1.0000				
4	B.No.Group-V	1.0000	1.0000				
5	B.No.Group-VI	1.0000	1.0000				

Valid Analysis Signatures

11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Analysis Audit Trail

- 11-06-2022 17:30:16 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
- 11-06-2022 17:38:27 Jyothi(M. Jyothi):DESKTOP-39TDEGC Error MD147 - Activity manually aborted by user.
- 11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
- 11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

MART SPECIALITIES LAB LLP.

Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Spectrometer Parameters - Fe

Element: Fe Wavelength: 248.3nm Background Correction: D2 Signal Type: Continuous Measurement Time: 4.0secs Use RSD Test: No

Flame Type: Air-C2H2 Nebuliser Uptake: 4secs Burner Height: 7.0mm

Sampling: None

Calibration Mode: Normal	
Concentration Units: mg/L	
Acceptable Fit: 0.990	
Standard 1	2.0000
Standard 2	4.0000
Standard 3	6.0000

No changes are recorded for this element

Measurement Mode: Absorbance Bandpass: 0.2nm High Resolution: Off Resamples: Fast Flier Mode: No

Lamp Current: 75% Optimise Spectrometer Parameters: No Number Of Resamples: 3

Flame Parameters - Fe

Fuel Flow: 0.9L/min Burner Stabilisation: 0mins Optimise Burner Height: No Auxilary Oxidant: Off Optimise Fuel Flow: No

Sampling Parameters - Fe

Calibration Parameters - Fe

Line Fit: Linear Scaled Units: mg/L Rescale Limit: 10.0% Standard 4 Standard 5 Use Stored Calibration: No Scaling Factor: 1.0000 Failure Action: Flag and Continue 8.0000 10.0000

Element Audit Trail - Fe

Solution Results - Fe

Y = 0.04924x + 0.0141 Fit: 0.9965 Characteristic Conc: 0.0894

> 1 2 3

> 1 2 3

> 1 2 3

> 1 2 3

1 2

3

Sample ID

Fe Blank

Fe Standard 1

Fe Standard 2

Fe Standard 3

Fe Standard 4

	4			Norma	I: Linea	ar Fit				
	Abs	*		*		*		*		*
	0 * -0-0	1 2	3	4	5	6	7	8	9	10
	-0-0	1 2	Ū		nc:mg/l		,	0	0	10
Signal	Rsd	Conc			cted Co					
Abs	%	mg/L		mg/L						
0.001	35.6	0.0000		0						
0.001	Backgro	ound: -0.003			11-06	-2022 1	7:31:26	;		
0.001	Backgro	ound: -0.003			11-06	-2022 1	7:31:30)		
0.000	Backgro	ound: -0.003			11-06	-2022 1	7:31:35	5		
0.120	0.3	2.0000								
0.120	Backgro	ound: 0.003			11-06	-2022 1	7:32:05	5		
0.119	-	ound: 0.003					7:32:09			
0.120	Backgro	ound: 0.003			11-06	-2022 1	7:32:13	5		
0.215	0.3	4.0000								
0.215	Backgro	ound: 0.004			11-06	-2022 1	7:32:40)		
0.215	Backgro	ound: 0.004			11-06	-2022 1	7:32:45	5		
0.216	Backgro	ound: 0.004			11-06	-2022 1	7:32:49)		
0.322	0.2	6.0000								
0.323	Backgro	ound: 0.005			11-06	-2022 1	7:33:19)		
0.322	Backgro	ound: 0.005			11-06	-2022 1	7:33:23	5		
0.322	Backgro	ound: 0.005			11-06	-2022 1	7:33:27	,		
0.411	0.3	8.0000								
0.410	Backgro	ound: 0.005			11-06	-2022 1	7:33:59)		
0.412	Backgro	ound: 0.005			11-06	-2022 1	7:34:03	5		
0.412	Backgro	ound: 0.005			11-06	-2022 1	7:34:07	,		

MART SPECIALITIES LAB LLP.

Operator Name: Jyothi

Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Solution Results - Fe

Sample ID	Signal	Rsd	Conc	Corrected Conc
-	Abs	%	mg/L	mg/L
Fe Standard 5	0.493	0.1	10.0000	
1	0.494	Backg	ground: 0.005	11-06-2022 17:34:38
2	0.494	Backg	ground: 0.005	11-06-2022 17:34:42
3	0.493	Backg	ground: 0.006	11-06-2022 17:34:46
Fe B.No.Group-II	0.834	0.1	16.6490 C	16.6490 C
1	0.834	Backg	ground: 0.005	11-06-2022 17:35:14
2	0.833	Backg	ground: 0.005	11-06-2022 17:35:19
3	0.834	Backg	ground: 0.005	11-06-2022 17:35:23
Fe B.No.Group-III	0.759	0.1	15.1286 C	15.1286 C
1	0.758	Backg	ground: 0.003	11-06-2022 17:35:52
2	0.760	Backg	ground: 0.003	11-06-2022 17:35:56
3	0.758	Backg	ground: 0.004	11-06-2022 17:36:00
Fe B.No.Group-IV	0.645	0.3	12.8189 C	12.8189 C
1	0.647	Backg	ground: 0.003	11-06-2022 17:36:31
2	0.644	Backg	ground: 0.003	11-06-2022 17:36:35
3	0.645	Backg	ground: 0.003	11-06-2022 17:36:40
Fe B.No.Group-V	0.809	0.2	16.1488 C	16.1488 C
1	0.808	Backg	ground: 0.005	11-06-2022 17:37:11
2	0.809	Backg	ground: 0.005	11-06-2022 17:37:15
3	0.811	Backg	ground: 0.005	11-06-2022 17:37:20
Fe B.No.Group-VI	0.711	0.3	14.1597 C	14.1597 C
1	0.709	Backg	ground: 0.005	11-06-2022 17:37:55
2	0.713	Backg	ground: 0.005	11-06-2022 17:37:59
3	0.712	Backg	ground: 0.004	11-06-2022 17:38:03

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-1 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
I	DETAILS OF THE SAMPLE
Sample Name : NA	
Name of the Manufacturer : NA	
Storage condition To b	up-I Mfg. : NA Exp. : NA e stored at room date date
	Derature Batch : NA C±3°C size
	by AAS Analysis.
	6/2022
Analysis Completion Date : 11/0	6/2022
Mfg. License No. : Not	provided
A.R.NO : NA	The Andrew Rest

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	620ppm

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



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Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-2 Sample Received Date : 11/06/2022 Report Date : 13/06/2022	
Q	DETAILS OF THE SAMPLE	
Sample Name : N	A	
Name of the Manufacturer : N	A	
Storage condition : To	roup-II Mfg. : NA Exp. : NA be stored at room date date	
Room Temperature : 25	Batch : NA size	
Tests Required:IreMethod:NAnalysis Starting Date:11	on by AAS Analysis.	
	Not provided	
A.R.NO : N	A	

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	832.5ppm

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



OCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00	
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinaga Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Sample Received Date : 11/06/2022	
a.u.	DETAILS OF THE SAMPLE	
Sample Name : N	JA	
Name of the Manufacturer : N	JA	
Storage condition	Group-III Mfg. : NA Exp. : NA To be stored at room date date Cemperature Batch : NA	
	5°C±3°C size	
	gm /	
Method : I Analysis Startin <mark>g Date : I</mark>	: NA : 11/06/2022	
and the second sec	: Not provided	
A.R.NO	NA	
Remark : S	Sample analyzed as received	

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	756.5ppm

×

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(Dr.R.Marayya)

Office: 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile: 98481 98350, 98481 25019 Mail: rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College	
Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000	Sample Received Date: 11/06/2022Report Date: 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name : N	A
Name of the Manufacturer : N	A
Storage condition : T	roup-IV Mfg. : NA Exp. : NA o be stored at room date date emperature Batch : NA
	5°C±3°C size
Method : N Analysis Starting Date : 1	on by AAS Analysis. A 1/06/2022 1/06/2022
Mfg. License <mark>No. :</mark> N	ot provided
A.R.NO : N	A
Remark : Sa	ample analyzed as received

Test Results

	and the second se	And the most of	
S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	641ppm	

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/	G
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Custome	r: Reference / Report No. : MSL/2022/JUNE/TARA/0064-5
Tara Government College	Sample Received Date : 11/06/2022
Prashanth Nagar Colony, Balajinag	ar Sumpto Received 2 mil
Sangareddy Telangana. 502000	Report Date : 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name :	NA
Name of the Manufacturer :	NA
Batch no : Storage condition	Group-VMfg. : NAExp. : NATo be stored at roomdatedateTemperatureBatch : NA
Room Temperature :	25°C±3°C size
Quantity Received	4gm
Tests Required:Iron by AAS Analysis.Method:NAAnalysis Starting Date:11/06/2022	
Analysis Completion Date : 11/06/2022	
Mfg. License No.	Not provided NA
A.R.NO Remark	Sample analyzed as received

Test Results

S.No.	Test Parameter	Result
01. Ir	on by AAS Analysis: (ppm)	807.5ppm

>

Authorized Signatory

(Dr.R.Marayya)

Office : 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile : 98481 98350, 98481 25019 Mail : rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G		
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00	
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-6 Sample Received Date : 11/06/2022 Report Date : 13/06/2022	
1	DETAILS OF THE SAMPLE	
Storage condition	up-V1 Mfg. : NA Exp. : NA be stored at room date date perature Batch : NA C±3°C size	
Tests Required:IronMethod:NAAnalysis Starting Date:11/0Analysis Completion Date:11/0Mfg. License No.:NotA.R.NO:NA	 Iron by AAS Analysis. NA 11/06/2022 	

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	708ppm

*

Authorized Signatory

(Dr.R.Marayya)

ADSORPTION STUDIES OF Fe[III] ON BENTONITE IN THE PRESENCE OF D-VALINE AS CHELATING AGENT

Dissertation submitted in Partial fulfillment for the requirements for the award of degree of

Bachelor of Science ín CHEMISTRY

By

1.Y. Avinash Reddy,	B.Sc(Chemistry),	III	Year
2.K.Prashanth Reddy,	B.Sc(Chemistry),	III	Year
3.V. Prashanth,	B.Sc(Chemistry),	III	Year
4.Y.Shashi Pranay,	B.Sc(Chemistry),	III	Year
5.D.Sai Kiran,	B.Sc(Chemistry),	III	Year
6.V.Aravind,	B.Sc(Chemistry),	III	Year

Under the Guídance:

K.SREEDHAR Assistant Professor DEPARTMENT OF CHEMISTRY



Tara Government College, Sangareddy(A) 2021-22

<u>CERTIFICATE</u>

This is to certify that the project work entitled "ADSORPTION STUDIES OF Fe[III] ON BENTONITE IN THE PRESENCE OF D-VALINE AS CHELATING AGENT " is presented by B.Sc (CHEMISTRY) students in partial fulfillment of the requirements for the degree of Bachelor of Science in Chemistry by the Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

The results embodied in this report have not been to any other University or Institution for the award of any degree.

(K.SREEDHAR)

Asst. Professor, Department of Chemistry Tara Government College, Sangareddy (A)

EXTERNAL EVALUATOR

ACKNOWLEDGEMENTS

We express my deep gratitude to my research supervisor **K.Sreedhar**, Asst. Professor, Department of Chemistry, Tara Govt. College, Sangareddy(A)-502001, INDIA for his inspiring guidance during the course of the Project work. The continuous encouragement extended by him propelled me to update my research skills and pedigree to engage in fruitful research.

We wish to express our gratitude to Smt. M.Praveena, Principal, Tara Govt. College, Sangareddy(A)-502001, INDIA for her constant support, cooperation and suggestions during the research work. We wish to express our sincere thanks to Dr.K.Abhijit, Head, Department of Chemistry, Tara Govt. College, Sangareddy(A)-502001, INDIA for providing me facilities, help and support for the entire research work.

We express our special thanks to **MART Specialities Lab, Hyderabad** for providing technical assistance in Atomic absorption spectrometric analysis.

Finally, we take this opportunity to thank one and all that has directly or indirectly helped me in completing the task.

1.Y. Avinash Reddy, B.Sc(Chemistry), III Year 2.K.Prashanth Reddy, B.Sc(Chemistry), III Year 3.V. Prashanth, B.Sc(Chemistry), III Year 4.Y.Shashi Pranay, B.Sc(Chemistry), III Year 5.D.Sai Kiran, B.Sc(Chemistry), III Year 6.V.Aravind, B.Sc(Chemistry), III Year

<u>DECLARATION</u>

We hereby declare that the project report entitled "ADSORPTION STUDIES OF Fe[III] ON BENTONITE IN THE PRESENCE OF D-VALINE AS CHELATING AGENT" is the work done by us in the campus at Department of Chemistry, Tara Government College, Sangareddy(A)during the academic year 2021-2022 and is submitted in partial fulfillment of the requirements for the degree of Bachelor of Science by Tara Govt. College, Sangareddy(A) (Affiliated to Osmania University, Hyderabad) during the academic year 2021-2022.

S.No.	Name of the Student	Roll Number	Group	Year	Signature
1	V. Prashanth	6058-19-578-036	B.Sc.(MCCs)	III	
2	V. Aravind	6058-19-578-037	B.Sc.(MCCs)	III	
3	Y. Avinash Reddy	6058-19-578-038	B.Sc.(MCCs)	III	
4	Y. Shashi Pranay	6058-19-578-039	B.Sc.(MCCs)	III	
5	D. Sai Kiran	6058-19-578-010	B.Sc.(MCCs)	III	
6	K.Prashanth Reddy	6058-19-578-018	B.Sc.(MCCs)	III	

TARA GOVERNMENT COLLEGE, SANGAREDDY – 502 001 (AUTONOMOUS)

BONAFIDE CERTIFICATE

Certified that the project report"ADSORPTION STUDIES OF Fe[III] ON

BENTONITE IN THE PRESENCE OF D-VALINE AS CHELATING AGENT"

is the bonafidework of

1.Y.Avinash Reddy, B.Sc(Chemistry), III Year 2.K.Prashanth Reddy, B.Sc(Chemistry), III Year 3.V.Prashanth, B.Sc(Chemistry), III Year 4.Y.Shashi Pranay, B.Sc(Chemistry), III Year 5.D.Sai Kiran, B.Sc(Chemistry), III Year 6.V.Aravind, B.Sc(Chemistry), III Year

who carried out the project work under my supervision.

K.Sreedhar **PROJECT SUPERVISOR**

DY.K.ABHIJIT HEAD, DEPARTMENT OF CHEMISTRY

Smt. M.PRAVEENA PRINCIPAL

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Chapter-I

INTRODUCTION

ADSORPTION

Adsorption is a process in which solids come into contact with liquids or gases, and the mass transfer occurs from liquids to solids. Desorption is the reversal of this action. Adsorption operations take advantage of a solid's capacity to concentrate certain chemicals from a fluid on to its surface. Adsorbate refers to the adsorbed substance, while adsorbent refers to the solid substance. The following are some examples of solid-liquid and solid-gas applications:

- Removing dissolved moisture from gasoline.
- Decolorization of sugar solutions and petroleum products.
- Removing noxious odors and tastes from water. Dehumidification of air and gases is one of the solid-gas activities.
- To fractionate mixtures of hydrocarbon gases such as methane, ethane, and propane.
- To remove undesirable odors and contaminants from gases.
- To recover valuable solvent vapors from dilute gas mixtures.

NATURE OF ADSORBENTS

Adsorbents are typically granular in nature, ranging in size from 0.5 mm to 12 mm. They can't have a lot of pressure decrease or get swept away by a fast-moving stream. During handling, they must maintain their shape and size. They'll need a lot of pores and a lot of surface area per unit mass.

Some of the commonly used adsorbents, their sources and applications are given below:

Sl. No.	Adsorbent	Source	Application
1.	Fuller's earth	Naturally occurring clay is heated and dried to get a porous structure.	De-colorizing, drying of lubricating oils, kerosene and engine oils.
2.	Activated charcoal	Bentonite or other activated clay which are activated by treatment with sulfuric acid and further washing, drying and crushing.	Used for de-colorizing petroleum products.
3.	Bauxite	A naturally occurring hydrated alumina, activated by heating at 230-815	Used for de-colorizing petroleum products and for drying gases.

4.	Alumina	A hard hydrated aluminum oxide, which is activated by heating to drive off the moisture and then crushed to desired size.	Used as desiccant.
5.	Bone-char	Obtained by destructive distillation of crushed bones at 600-900	Used for refining sugar and can be reused after washing and burning.
6.	Silica gel	A hard granular and porous product obtained from sodium silicate solution after treatment with acid. Normally has 4 to 7% water in the product.	Dehydration of gases and liquids, and separation of gas-liquid hydrocarbon mixture.

7.	Activated carbon	 (1) Vegetable matter is mixed with calcium chloride, carbonized and finally the inorganic compounds are leached away. (2) Organic matter is mixed with porous pumice stones and then heated and carbonized to deposit the carbonaceous matter throughout the porous particle. (3) Carbonizing substances like wood, sawdust, coconut shells, fruit pits, coal, lignite and subsequent activation with hot air steam. It is available in granular or pelleted form. 	De-colorizing of sugar solutions, chemicals, drugs, water purification, refining of vegetable and animal oils, recovery of gold and silver from cyanide ore-leach solution, recovery of solvent vapor from gas-mixtures, collection of gasoline hydro- carbons from natural gas, fractionation of hydrocarbon gases.
8.	Molecular sieves	These are porous synthetic zeolite crystals, metal alumino-silicates.	Dehydration of gases and liquids, and separation of gas- liquid hydrocarbon mixture.

SIGNIFICANCE OF RESEARCH PROBLEM

Adsorption of heavy metals is an important strategy to develop newer remediation technologies for the sustainable environmental protection. But the efficacy of adsorption of heavy metals under the normal conditions using suitable adsorbent depends on several factors which need to be finely tuned to get efficient adsorption process. In the presence of proper facilitating agents, the adsorption of heavy metals enhanced which will certainly improves the existing heavy metal techniques.

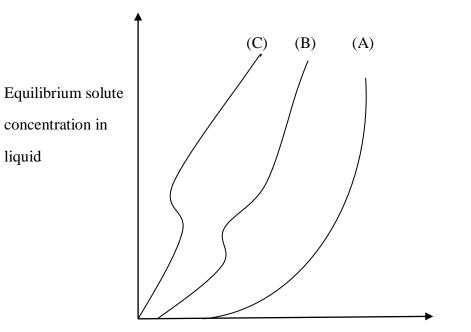
OBJECTIVE OF THE PROJECT

To evaluate the impact of D-Valine as a chelating agent in the adsorption of Fe (III) from aqueous solution by Bentonite as an adsorbent to develop efficient remediation technology using concept of Coordination chemistry.

Chapter-II

ADSORPTION OF SOLUTES FROM DILUTE SOLUTIONS

Both the solvent and the solute are adsorbed whenever a mixture of solute and solvent is adsorbed using an adsorbent. As a result, only relative or apparent solute adsorption can be determined. As a result, treating a known volume of solution of original concentration Cwith a known weight of adsorbent is standard procedure. Let C* be the solution's final equilibrium solute concentration. If v is the volume of solution per unit mass of adsorbent (cc/g), and C and C* are the starting and equilibrium concentrations (g/cc) of the solute, then the apparent adsorption of the solute per unit mass of adsorbent is $v(C - C^*)$, (g/g), neglecting any volume change. This statement is most useful in the case of dilute solutions. The C* value is determined by the temperature, nature, and properties of the adsorbent when the proportion of the original solvent that can be adsorbed is tiny. The Freundlich adsorption isotherm, $C^*=K[v(C - C^*)]n$, represents the adsorption phenomena in dilute fluids across a small concentration range. The Freundlich adsorption equation is especially useful in situations where the identification of the solute is unknown, such as the removal of coloring substances from sugar solutions, oils, and other liquids. A spectrophotometer or colorimeter can quickly determine the color composition of the solute. In worked example 2, the interpretation of this data is demonstrated. Adsorption is good if the value of n is high, say 2 to 10. If it's between 1 and 2, it's relatively challenging, and if it's less than 1, it's easy and it indicates poor adsorption characteristics. Freundlich adsorption equation is also useful in such a case where the actual identification of solute is not known, e.g., removal of coloring substance from sugar solutions, oils etc. A typical adsorption isothermal for the adsorption of various adsorbents A, B and C in dilute solution at the same temperature for the same adsorbent is represented in a graph.



Kg solute apparently adsorbed /kg adsorbent

Adsorption isotherms for various adsorbents

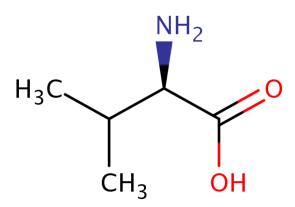
METAL COMPLEXES OF AMINO ACIDS

Transition metal amino acid complexes are a large family of coordination complexes containing the conjugate bases of the amino acids, the 2-aminocarboxylates. Amino acids are prevalent in nature, and all of them function as ligands toward the transition metals. Not included in this article are complexes of the amides (including peptide) and ester derivatives of amino acids. Also excluded are the polyamino acids including the chelating agents EDTA and NTA.

Most commonly, amino acids coordinate to metal ions as N,O bidentate ligands, utilizing the amino group and the carboxylate. They are "L-X" ligands. A five-membered chelate ring is formed. The chelate ring is only slightly ruffled at the sp³-hybridized carbon and nitrogen centers. For those amino acids containing coordinating substituents, the resulting complexes are more structurally diverse since these substituents can coordinate. Histidine, aspartic acid, methionine, and cysteine sometimes form tridentate N,N,O, N,O,O, S,N,O, and S,N,O complexes, respectively. Using kinetically inert metal ions,

complexes containing monodentate amino acids have been characterized. These complexes exist in either the N or the O linkage isomers. It can be assumed that such monodentate complexes exist transiently for many kinetically labile metal ions (e.g. Zn^{2+}). Mixing simple metal salts with solutions of amino acids near neutral or elevated pH often affords bis- or tris complexes. For metal ions that prefer octahedral coordination, these complexes often adopt the stoichiometry $M(aa)_3$ (aa = amino carboxylate, such as glycinate, $H_2NCH_2CO_2^{-}$). Complexes of the 3:1 stoichiometry have the formula is $[M(O_2CC(R)HNH_2)_3]^2$. Such complexes adopt octahedral coordination geometry. These complexes can exist in facial and meridional isomers, both of which are chiral. The stereochemical possibilities increase when the amino acid ligands are not homochiral. Complexes with the 2:1 stoichiometry are illustrated by copper(II) glycinate $[Cu(O_2CC(R)HNH_2)_2]$, which exists both in anhydrous and pentacoordinate geometries. When the metal is square planar, these complexes can exist as cis and trans isomers. The stereochemical possibilities increase when the amino acid ligands are not homochiral. Homoleptic complexes are also known where the amino carboxylate is tridentate amino acids. One such complex is $Ni(\kappa^3-histidinate)_2$.

D-Valine can easily involved in M-L complex formation due to its functional flexibility. Moreover its water solubility encourages the metal-ligand complex formation under aqueous conditions.



D-Valine

BENTONITE AS ADSORBENT

Because of heavy metals toxicity and non-biodegradable nature, the advent of heavy metals in water is turning into a critical environmental and public fitness concern. A variety of technology were evolved to do away with poisonous heavy metals from wastewater. The maximum crucial technology for the heavy steel ions elimination from wastewater consist of perception, ion exchange, adsorption, coagulation, evaporation and opposite osmosis. Adsorption on strong matrices has been proven to be an economically viable opportunity method (Abollino et al., 2003¹; Hoda et al., 2009² Kapoor and Vira Raghavan, 1998³). Cheaper Na-bentonite has been observed to be so beneficial for elimination of heavy steel ions from aqueous answers that it has attracted geologists and environmental engineers (Al-Quadbit et al., 2005⁴; Guo et al.,(2009).

The Na-bentonite from Gaomiaozi has been used as boundaries to save you infection of wastewater containing heavy metals. For this cause it's far crucial to take a look at the adsorption of metals of metals via way of means of Na-bentonite so one can offer crucial parameters and essential principle for the knowledge of adsorptive elimination of heavy metals via way of means of Na-bentonite from an aqueous environment. The bivalence ions of copper and nickel are not unusual place observed withinside the commercial wastewater. The bivalence ions of copper and nickel are commonly found in the industrial wastewater. On a Na-montmorillonite turned into studied as feature of answer pH, dosage of Na-bentonite, temperature and make contact with and make contact with time. The aggressive isothermal Adsorption conduct of copper and nickel on Na-bentonite in single-aspect structures and binary-aspect structures has been investigated.



Chapter-III

ATOMIC ABSORPTION SPECTROMETRY

Atomic absorption spectrometry (AAS) is an analytical technique that measures the concentrations of elements. Atomic absorption is so sensitive that it can measure down parts per billion of a gram (µg dm⁻³) in a sample. The technique makes use of the wavelengths of light specifically absorbed by an element. They correspond to the energies needed to promote electrons from one energy level to another, higher, energy level.

Atomic absorption spectrometry has many uses in different areas of chemistry.

Clinical analysis: Analysing metals in biological fluids such as blood and urine.

Environmental analysis: Monitoring our environment- eg finding out the levels of various elements in rivers, seawater, drinking water, air, petrol and drinks such as wine, beer and fruit drinks.

Pharmaceuticals: In some pharmaceutical manufacturing processes, minute quantities of a catalyst used in the process (usually a metal) are sometimes present in the final product. By using AAS the amount of catalyst present can be determined.

Industry: Many raw materials are examined and AAS is widely used to check that the major elements are present and that toxic impurities are lower than specified- eg in concrete, where calcium is a major constituent, the lead level should be low because it is toxic.

Mining: By using AAS the amount of metals such as gold in rocks can be determined to see whether it is worth mining the rocks to extract the gold.

HOW IT WORKS

Atoms of different elements absorb characteristic wavelengths of light. Analysing a sample to see if it contains a particular element means using light from that element. For example with lead, a lamp containing lead emits light from excited lead atoms that produce the right mix of wavelengths to be absorbed by any lead atoms from the sample. In AAS, the sample is atomized- i.e. converted into ground state free atoms in the vapour state- and a beam of electromagnetic radiation emitted from excited lead atoms is passed through the vaporized sample. Some of the radiation is absorbed by the lead atoms in the sample. The greater the number of atoms there is in the vapour, the more radiation is absorbed. The amount of light absorbed is proportional to the number of lead atoms. A calibration curve is constructed by running several samples of known lead concentration under the same conditions as the unknown. The amount the standard absorbs is compared with the calibration curve and this enables the calculation of the lead concentration in the unknown sample.

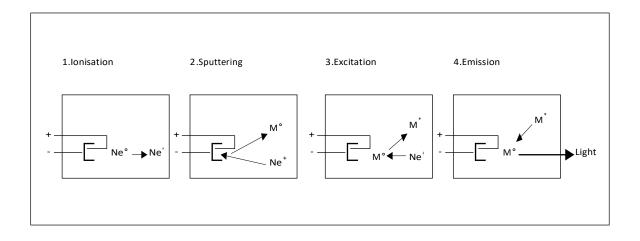
Consequently an atomic absorption spectrometer needs the following three components: a light source; a sample cell to produce gaseous atoms; and a means of measuring the specific light absorbed.

THE LIGHT SOURCE

The common source of light is a 'hollow cathode lamp'. This contains a tungsten anode and a cylindrical hollow cathode made of the element to be determined. These are sealed in a glass tube filled with an inert gas- e.g neon or argon- at a pressure of between 1 Nm⁻² and 5 Nm⁻².

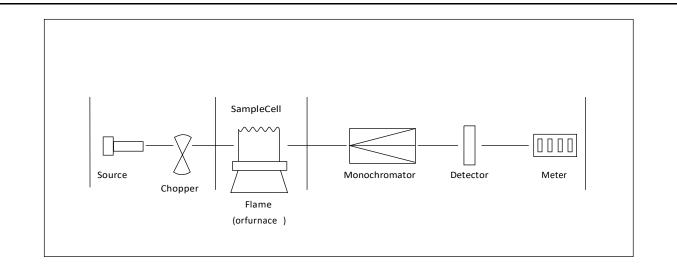


The ionization of some gas atoms occurs by applying a potential difference of about 300-400V between the anode and the cathode. These gaseous ions bombard the cathode and eject metal atoms from the cathode in a process called sputtering. Some sputtered atoms are in excited states and emit radiation characteristic of the metal as they fall back to the ground state $-egPb^* \rightarrow Pb + h v$. The shape of the cathode concentrates the radiation into a beam which passes through a quartz window, and the shape of the lamp is such that most of the sputtered atoms are redeposited on the cathode. A typical atomic absorption instrument holds several lamps each for a different element. The lamps are housed in a rotating turret so that the correct lamp can be quickly selected.



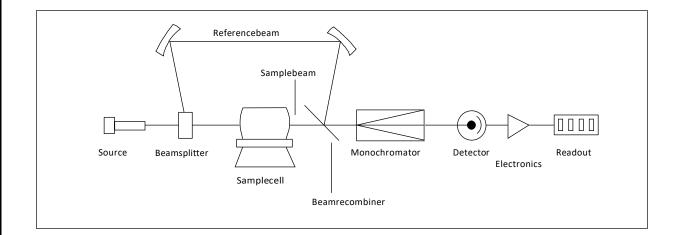
THE OPTICAL SYSTEM AND DETECTOR

A monochromator is used to select the specific wavelength of light –ie spectral line – which is absorbed by the sample, and to exclude other wavelengths. The selection of the specific light allows the determination of the selected element in the presence of others. The light selected by the monochromator is directed onto a detector that is typically a photomultiplier tube. This produces an electrical signal proportional to the light intensity



DOUBLE BEAM SPECTROMETERS

Modern spectrometers incorporate a beam splitter so that one part of the beam passes through the sample cell and the other is the reference. The intensity of the light source may not stay constant during an analysis. If only a single beam is used to pass through the atom cell, a blank reading containing no analyte (substance to be analysed) would have to be taken first, setting the absorbance at zero. If the intensity of the source changes by the time the sample is put in place, the measurement will be inaccurate. In the double beam instrument, there is a constant monitoring between the reference beam and the light source. To ensure that the spectrum does not suffer from loss of sensitivity, the beam splitter is designed so that as high a proportion as possible of the energy of the lamp beam passes through the sample.



ATOMIZATION OF THE SAMPLE

Two systems are commonly used to produce atoms from the sample. Aspiration involves sucking a solution of the sample into a flame; and electrothermal atomization is where a drop of sample is placed into a graphite tube that is then heated electrically.

Some instruments have both atomization systems but share one set of lamps. Once the appropriate lamp has been selected, it is pointed towards one or other atomization system.

FLAME ASPIRATION

Ethyne/air (giving a flame with a temperature of 2200–2400°C) or ethyne/dinitrogen oxide (2600–2800°C) are often used. A flexible capillary tube connects the solution to the nebulizer. At the tip of the capillary, the solution is 'nebulized' –ie broken into small drops. The larger drops fall out and drain off while smaller ones vaporize in the flame. Only ca 1% of the sample is nebulized.

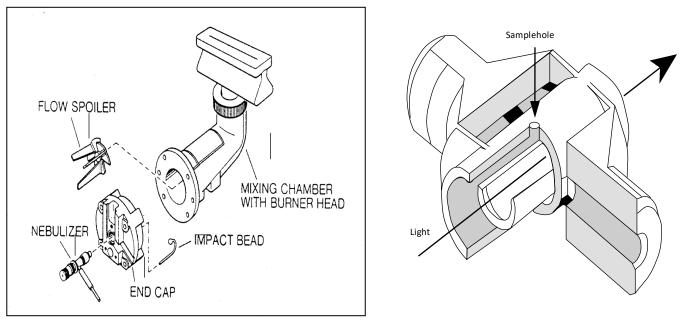


Figure 1

Figure 2

ELECTROTHERMAL ATOMIZATION

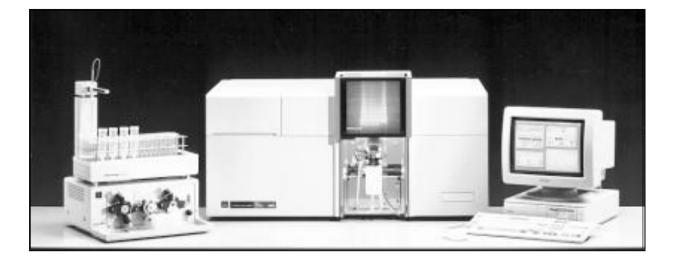
 $25 \ \mu$ l of sample (ca 1/100th of a raindrop) is placed through the sample hole and onto the platform from an automated micropipette and sample changer. The tube is heated electrically by passing a current through it in a pre-programmed series of steps. The details will vary with the sample but typically they

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might be 30–40 seconds at 150°C to evaporate the solvent, 30 seconds at 600°C to drive off any volatile organic material and char the sample to ash, and with a very fast heating rate (ca 1500 °C s⁻¹) to 2000-2500°C for 5–10 seconds to vaporise and atomize elements (including the element being analysed). Finally heating the tube to a still higher temperature –ca 2700°C – cleans it ready for the next sample. During this heating cycle the graphite tube is flushed with argon gas to prevent the tube burning away. In electrothermal atomization almost 100% of the sample is atomised. This makes the technique much more sensitive than flame AAS.

SAMPLE PREPARATION

Sample preparation is often simple, and the chemical form of the element is usually unimportant. This is because atomization converts the sample into free atoms irrespective of its initial state. The sample is weighed and made into a solution by suitable dilution. Elements in biological fluids such as urine and blood are often measured simply after a dilution of the original sample.



When making reference solutions of the element under analysis, for calibration, the chemical environment of the sample should be matched as closely as possible –i.e., the analyte should be in the same compound and the same solvent. Teflon containers may be used when analysing very dilute Solutions because elements such as lead are sometimes leached out of glass vessels and can affect the results

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BACKGROUND ABSORPTION

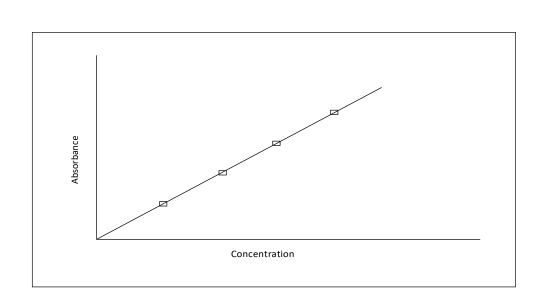
It is possible that other atoms or molecules apart from those of the element being determined will absorb or scatter some radiation from the light source. These species could include unvaporised solvent droplets, or compounds of the matrix (chemical species, such as anions, that tend to accompany the metals being analysed) that are not removed completely. This means that there is a background absorption as well as that of the sample.

One way of measuring and correcting this background absorption is to use two light sources, one of which is the hollow cathode lamp appropriate to the element being measured. The second light source is a deuterium lamp.

The deuterium lamp produces broad band radiation, not specific spectral lines as with a hollow cathode lamp. By alternating the measurements of the two light sources – generally at 50 - 100 Hz – the total absorption (absorption due to analyte atoms plus background) is measured with the specific light from the hollow cathode lamp and the background absorption is measured with the light from the deuterium lamp. Subtracting the background from the total absorption gives the absorption arising from only analyte atoms.

CALIBRATION

A calibration curve is used to determine the unknown concentration of an element -eg lead – in a solution. The instrument is calibrated using several solutions of known concentrations. A calibration curve is produced which is continually rescaled as more concentrated solutions are used – the more concentrated solutions absorb more radiation up to a certain absorbance. The calibration curve shows the concentration against the amount of radiation absorbed in the given figure. (a) The sample solution is fed into the instrument and the unknown concentration of the element-e.g., lead- is then displayed on the calibration curve given in the below figure. (b)





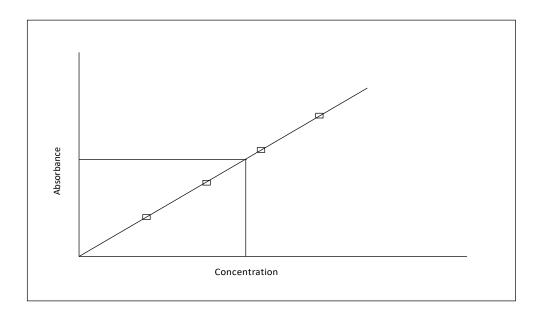


Figure (b)

Chapter-IV

METHODOLOGY

MATERIALS REQUIRED

- Ferric chloride Hexahydrate
- D-Valine
- Bentonite
- Volumetric flask
- Digital Weighing Machine
- Watch glasses
- Hot air oven
- Ultra-pure water (Demineralized)
- 2% of Nitric acid

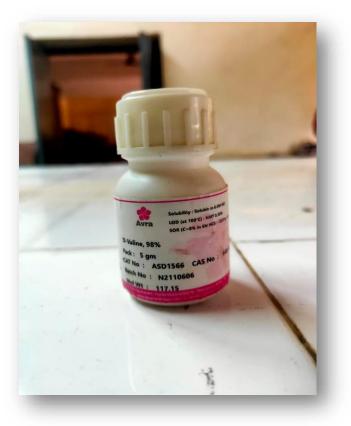
PROCEDURE

- Using a 250ml Volumetric flask prepare a metal ligand solution by adding 100ppm of Ferric chloride Hexahydrate (MOLYCHEM MCR-11580) and 200ppm of L-Valine (AVRAN2110606).
 Prepare 250ml solution by adding Ultra-pure water and then keep this system aside for a few hours.
- Weigh 5 grams of Bentonite (AVRA N2101070) using a Digital weighing machine (Citizen Scales(I) PVT LTD CTG302-300) and take this into a beaker.
- **3.** Now add 100ml of the above prepared metal ligand solution into the beaker and stir the mixture well for 10 minutes using a glass rod.
- **4.** Keep this mixture aside for 48 hours without disturbing it as at this step Ferric is going to be adsorbed on Bentonite in the presence of D-Valine which acts as a chelating agent.
- **5.** After completion of 48 hours take the mixture and filter it off using Whatman Grade 1 filter paper and a funnel.

- **6.** After filtration of the mixture again add Ultra-pure water for 3 times and then filtrate it to obtain pure concentration of Ferric solution which is get adsorbed on Bentonite.
- Collect the filtered Bentonite powder and place it on a watch glass and keep this in a Hot air Owen at 60 °C for 10 hours to get rid of moisture present in it.
- 8. Now weigh each 1 gm of Bentonite in glass vials.
- 9. Now take a beaker and rinse it with ultra-pure water then followed by Nitric acid.
- **10.** Take 0.5 grams of Bentonite sample in the beaker and add 2% of Nitric acid and stir the mixture well for 10-15 minutes.
- **11.** Filter the mixture using Whatman Grade 1 filter paper and again 3 times by using Ultra-pure water to obtain pure concentration of Ferric present in the mixture prepared using the sample.
- **12.** Take this collected sample solution and keep this system under AAS (Thermos Scientific iCE 3300)
- **13.** Calculate the concentration of Ferric adsorbed on Bentonite at different ppm levels. Observe the graph obtained and note down the readings of the result we obtained.
- 14. Same Experiment carried out without the interference/addition of Ligand i.e. D-Valine for Control Experiment.



Bentonite Powder



L-VALINE



FERRIC CHLORIDE (HEXAHYDRATE) FeCL₃6H₂O

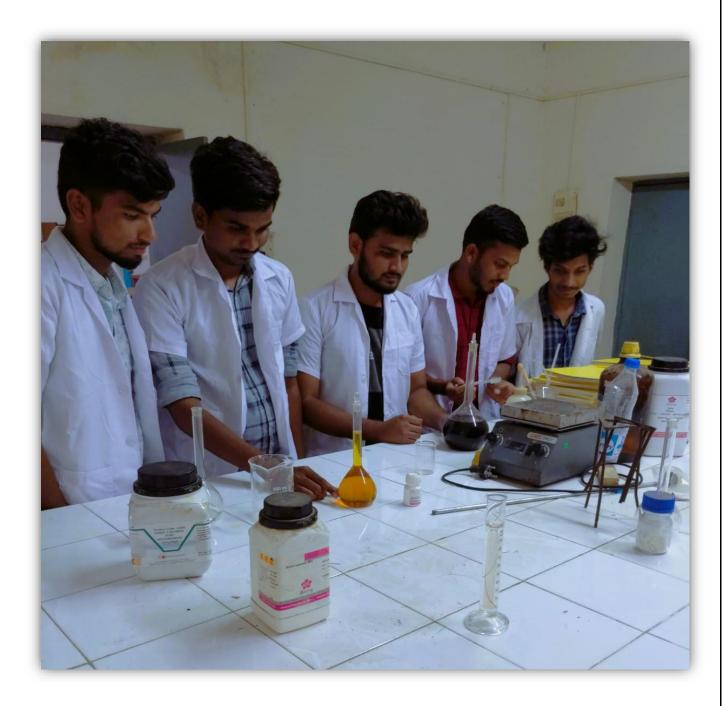


Measuring Flask





Preparing metal ligand solution





5 grams of bentonite added to 100ml of metal ligand solution



Filtering The Mixture using Whatman grade 1 filter paper and Again, three times by using Ultra-Pure water







Hot air oven



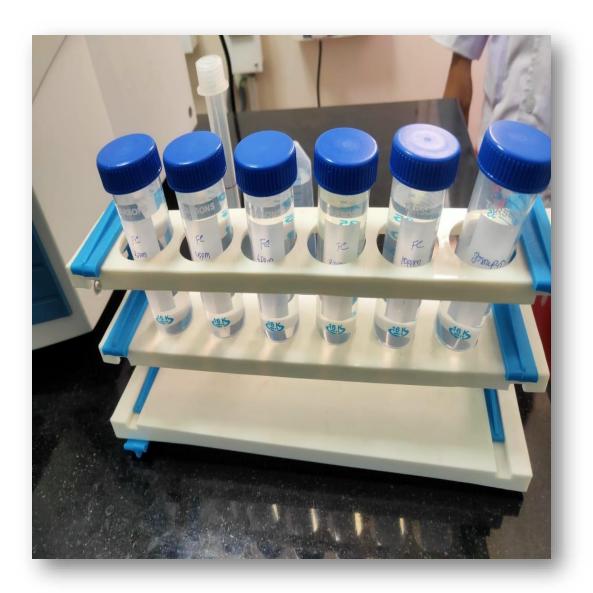
Glass vials



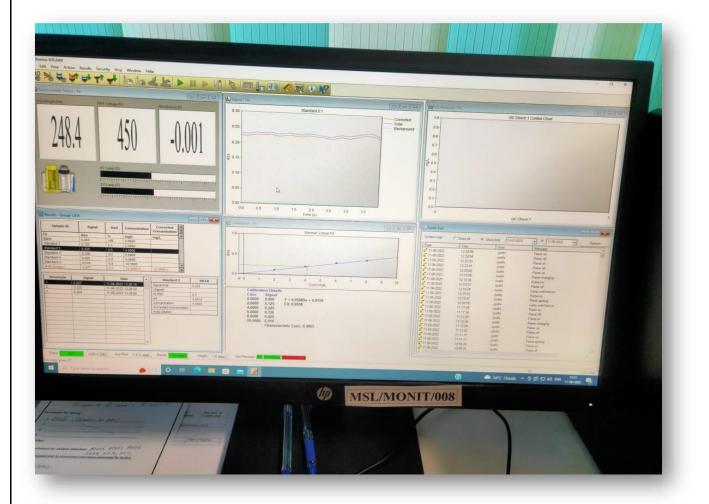
Collected samples



AAS (Atomic Absorption Spectrometry)



Sample solution added to 2% of nitric acid

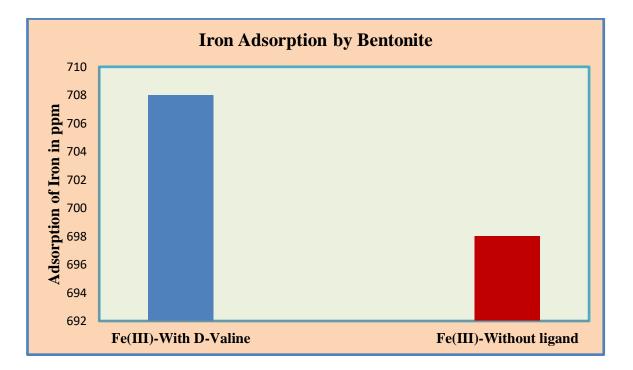


Results obtained on the monitor under AAS Method

Chapter-V

RESULTS AND DISCUSSIONS

Bentonite adsorbs **708.0ppm** of Iron metal from aqueous solution of Fe(III)-D-Valine metal ligand solution. Whereas, Bentonite adsorbs only **698.0ppm** when D-Valine is absent. It is evident from the AAS results, ligand involvement enhanced the metal adsorption by initiating potential chemical interactions between adsorbate and adsorbent. D-Valine firmly coordinates with Fe(III) to form a stable complex in aqueous condition. The complex coordination sphere in the resulted complex facilitates strong interactions with the polar points of Bentonite. From the AAS results, it is conclusive that **1.432%** of adsorbtion increased in the presence of D-Valine as chelating agent.



Impact of D-Valine on Adsorption of Fe (III) ions from aqueous solution by Bentonite.

<u>Spectrometer Parameters</u> – <u>Fe</u>:

Element : Fe	Measurement mode : Absorbance	
Wavelength : 248.3nm	Band pass : 0.2nm	Lamp current : 75%
Background correction :	High Resolution : Off	Optimise Spectrometer
D2		Parameters : No
Signal type : continuous	Resamples : Fast	Number of resamples : 3
Measurement time :	Flier mode : No	
4.0secs		
Use RSD Test : No		

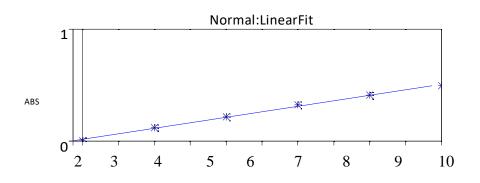
<u>Flame parameters – Fe:</u>

Flame type : Air – C2H2	Fuel Flow : 0.9L/min	Auxiliary Oxidant : Off
Nebuliser Uptake : 4secs	Bunsen Stabilisation : Omins	Optimise Fuel Flow : No
Burner Height :7.0mm	Optimise Burner fuel : No	

<u>Calibration Parameters</u> – <u>Fe</u>:

Calibration mode : Normal	Line fit : Linear	Use stored calibration : No
Concentration units : mg/L	Scales units : mg/L	Scaling factor : 1.0000
Acceptable fit : 0.990	Rescale Limits : 10.0%	Failure Action : Flag and Continue
Standard 1 - 2.0000	Standard 4 - 8.0000	
Standard 2 - 4.0000	Standard 5 - 10.0000	
Standard 3 - 6.0000		

Solutions Results-Fe:



Y = 0.04924x + 0.0141

Fit: 0.9965

Characteristic Conc.: 0.0894

Sample ID	Signal	RSD	Conc.
	Abs	%	Mg/L
Fe Blank	0	<99	0
1	0	В	ackground: 0.000
2	0	В	ackground: 0.000
3	0	В	ackground: 0.000
Fe Standard 1	0.125	0.	3 2
1	0.125	В	ackground: 0.006
2	0.125	В	ackground: 0.006
3	0.126	В	ackground: 0.006
Fe Standard 2	0.225	0.	5 4
1	0.227	В	ackground: 0.007
2	0.225	В	ackground:0.007

2	0.702		_	ground: 0.005 ground: 0.005	
1	0.700		_	ground: 0.005	
Fe(III)-Bentonite (Without Ligand)	0.701	0.1		16.4138 C	
3	0.712	Background: 0.004			
2	0.713	Background: 0.005			
1	0.709		Back	ground: 0.005	
Fe(III)-D-Valine-Bentinite-	0.711	0.1	1	16.6490 C	
3	0.513	3 Background: 0.009			
2	0.509	9 Background: 0.009			
1	0.51		Backg	ground: 0.009	
Fe Standard 5	0.51		0.4		10
3	0.425		Backg	ground: 0.009	
2	0.425		Backg	ground: 0.008	
1	0.426		Backg	ground: 0.009	
Fe Standard 4	0.425		0.2		8
3	0.335		Backg	ground: 0.008	
2	0.336		Backg	ground: 0.008	
1	0.336		Backg	ground: 0.008	
Fe Standard 3	0.336		0.2		6
3	0.224		Backg	ground:0.007	

TEST RESULTS:

S.No.	Test Parameters	Sample	Results
01.	Iron by AAS	Fe(III)- D-Valine-	708.0ppm
	Analysis: (ppm)	Bentonite Sample	
02.	Iron by AAS	Fe(III)-Bentonite	698.0ppm
	Analysis: (ppm)	(control)	

CONCLUSION

From the current project it is clear that D-Valine as a Chelating agent has played a vital role in adsorption of Fe(III) ions from aqueous solution and increases the adsorption up to **1.432** % using Bentonite. This aspect will be useful in designing the newer strategies of Heavy metal Remediation techniques using organic bifunctional Chelating Ligands as Facilitating agents in Metal Adsorption processes.

REFERENCES

- 1. Abollino, O., Aceto, M., Malandrino, M., Sarzanini, C. and Mentasti, E., 2003, Adsorption of metals on Na-montmorillonite. Effect of pH and organic substances. Water Res, 37: 1619-1627.
- Al-Qunaibit, M.H., Mekhemer, W.K. and Zaghloul, A.A., 2005, The adsorption of Cu (II) ions on bentonite—a kinetic study. Jcolloid interface sci, 283: 316-321.
- Adhikari T, Singh M V. Sorption characteristics of lead and cadmium in some soils of India. Geoderma, 2003(114): 81–92.
- mental Science Health, 1996(31): 2099–2109. [17] Altin O, Ozbelge O H, Dogu T. Effect of pH, flow rate and concentration on the sorption of Pb and Cd on montmorillonite. J Chem Tech Biotechnol, 1999(74): 1131–1138.
- Alvarez-Ayuso E, Garcia-Sanchez A (2003) Removal of heavy metals from waste waters by natural and Na-exchanged bentonites. Clays Clay Miner 51:475–480
- Argun ME, Dursum S (2008) A new approach to modification of natural adsorbent for heavy metal adsorption. Bioresource Technol 99:2516–2527.
- Ayari F, Srasra E, Trabelsi-Ayadi M (2005) Characterization of bentonitic clays and their use as adsorbent. Desalination 185:391–397.
- Bahranowski K. Structure and texture of copper-doped alumina-pillared montmorillonites. Mineralogia Polonica, 1998, 29(1): 55–65.

- Barakat MA (2011) New trends in removing heavy metals from industrial wastewater. Arab J Chem 4:361–377
- 10. Bereket G, Aroguz AZ, Ozel MZ (1997) Removal, Cu (II), and Ni (II) from aqueous solutions by adsorption on bentonite. J Colloid Intern Sci 187:338–343.
- 11. Bhattacharya AK, Mandal SN, Das SK (2006) Adsorption of cu (II), and Ni (II) from aqueous solution by using different adsorbents. Chem Eng J 123:43–51.
- 12. Bourliva A, Michailidis K, Sikalidis C, Filippidis A, Betsiou M (2012) nickel removal from aqueous solutions utilizing Greek natural bentonite and vermiculite. Fresen Environ Bull 21:2466–2471.
- Bourliva A, Michailidis K, Sikalidis C, Filippidis A, Betsiou M (2013) Lead removal from aqueous solutions by Na-bentonites. Clay Miner 48:771–787.
- Chen J, Du G C. The Production and Application of Environmentally Friendly Materials. Beijing: Chemical Industry Press, 2002. (In Chinese).
- 15. Cummings NE, Williams EM, Kasza I, Konon EN, Schaid MD, Schmidt BA, et al. (February 2018). . *The Journal of Physiology*. **596** (4): 623–645.
- CHRISTENSEN T.H., KJELDSEN P., BJERG P.L., JENSEN D.L., CHRISTENSEN J.B., BAUN
 A., ALBRECHTSEN H.J., HERN H. Biogeochemistry of landfill leachate plumes. Appl. Geochem.
 16, 659, 2001.
- 17. CHEN Y.G., YE W.M., YANG X.M., DENG F.Y., HE Y. Effect of contact time, pH, and ionic strength on Cu (II) adsorption from aqueous solution onto bentonite from Gaomiaozi, China. Environ. Earth Sci. 64, 329, 2011.
- CHAI J.C., MIURA N. Comparing the performance of landfill liner systems. J. Mater. Cycles. Waste Manage. 4, 135, 2002.
- CUERVO M., ASEDEGBEGANIETO E., DIAZ E., ORDONEZ S., VEGA A., DONGIL A. Modification of the adsorption properties of high surface area graphite's by oxygen functional groups. Carbon 46, 2096, 2008.
- 20. Chen YG, He Y, Ye WM, Lin CH, Zhang XF, Ye B (2012) Removal of cu(II) from aqueous solutions by adsorption on bentonite from Gaomiaozi, China. Environ Earth Sci 67:1261–1268.

- Dawson RM, Elliott DC, Elliott WH, Jones KM, eds. (1959). Data for Biochemical Research.
 Oxford: Clarendon Press.
- 22. Donat R, Akdogan A, Erdem E, Cetisli H (2005) Thermodynamics of cu2? and Ni2? adsorption onto natural bentonite from aqueous solutions. J Colloid Interf Sci 286:43–52.
- 23. Dong Q H, Cai R, Yang W F. Simulation of water-resistance of a clay layer during mining: analysis of a safe water head. Journal of China University of Mining & Technology, 2007, 17(3): 345–348.
- 24. DING S., SUN Y., YANG C., XU B. Removal of copper from aqueous solutions by bentonites and the factors affecting it. Min Sci Technol. 19, 489, 2009.
- 25. Etci O[°], Bektas, N, O[°] ncel MS (2010) Single and binary adsorption of lead and cadmium ions from aqueous solution using the clay mineral beidellite. Environ Earth Sci 61:231–240.
- 26. Evans RC (1966) An introduction to crystal chemistry. Cambridge University Press, London Freundlich H (1907) Ueber die Adsorption in Loesungen. Z Phys Chem 57:385–470.
- 27. Fu F, Wang Q (2011) Removal of heavy metal ions from wastewaters: a review. J Environ Manage 92:407–418.
- 28. Freundlich H (1907) Ueber die Adsorption in Loesungen. Z Phys Chem 57:385-470.
- 29. Fu F, Wang Q (2011) Removal of heavy metal ions from wastewaters: a review. J Environ Manage 92:407–418.
- 30. Fontana L, Cummings NE, Arriola Apelo SI, Neuman JC, Kasza I, Schmidt BA, et al. (July 2016). . Cell Reports. 16 (2): 520–530. .
- 31. FRANCISCA F.M., GLATSTEIN D.A. Long term hydraulic conductivity of compacted soils permeated with landfill leachate. Appl. Clay Sci. 49, 187, 2010.
- 32. FREUNDLICH H.M.F. Uber die adsorption in losangen Z. Phys.Chem. 57,385, 1906.
- Garci´a-delgado R A, Garci´a-herruzo F. Influence of soil carbonates in lead fixation. Journal of Environmental Science Health, 1996(31): 2099–2109.

- 34. Guo, Z.J., Xu, J., Shi, K.L., Tang, Y.P., Wu, W.S. and Tao, Z.Y., 2009, Eu(III) adsorption/desorption on Na-bentonite: experimental and modeling studies. Colloids Surf A: Physicochem Eng Aspects, 339: 126–133.
- 35. Giles CH, Smith D, Huitson A (1974) A general treatment and classification of the solute adsorption isotherm: I Theoretical. J Colloid Interf Sci 47:755–765.
- 36. Gupta SS, Bhattacharyya KG (2008) Immobilization of Pb(II), Cd(II) and Ni(II) ions on kaolinite and montmorillonite surfaces from aqueous medium. J Environ Manage 87:46–58.
- 37. GLATSTEIN D.A., FRANCISCA F.M. Influence of pH and ionic strength on Cd, Cu and Pb removal from water by adsorption in Na-bentonite. Appl. Clay Sci. 118, 61, 2015.
- 38. Hui X J. Environmental Toxicology. Beijing: Chemical Industry Press, 2003. (In Chinese).
- Hlavaty V. The montmorillonite interlayer space expansion control by meth acrylamide intercalation. Geologica, 1994, 38(2–4): 217–228.
- 40. Ho, Y.S. and McKay, G., 2000, The kinetics of sorption of divalent metal ions onto sphagnum moss peat. Water Res, 34: 735–742.
- 41. Hoda, O., Hassan, A. and Abdelhakium, D., 2009, Adsorption of 60Co radionuclides from aqueous solution by raw and modified bentonite. Appl Clay Sci, 44: 21–26.
- 42. Hamidpour M, Kalbasi M, Afyuni M, Shariatmadari H, Furrer G (2011) Sorption of lead on Iranian bentonite and zeolite: kinetics and isotherms. Environ Earth Sci 62:559–568.
- 43. Irving, H. and Williams, R.J.P., 1953, The stability of transition-metal complexes. J Chem Soc, 3192–3210.
- 44. Ijagbemi C, Baek MH, Kim DS (2009) Montmorillonite surface properties and sorption characteristics for heavy metal removal from aqueous solutions. J Hazard Mater 166:538–546.
- 45. (2002).. Dietary Reference Intakes for Energy, Carbohydrates, Fiber, Fat, Fatty Acids, Cholesterol, Protein, and Amino Acids. Washington, DC: The National Academies Press. pp. 589–768.
- 46. Jiang M, Jin X, Lu X, Chen Z (2010) Adsorption of Pb(II), Cd(II), Ni(II) and Cu(II) onto natural kaolinite clay. Desalination 252:33–39.

- 47. Jang C, Oh SF, Wada S, Rowe GC, Liu L, Chan MC, et al. (April 2016). <u>"</u>. Nature Medicine. 22 (4): 421–6
- 48. JIANG M., JIN X., LU X.Q., CHEN Z. Adsorption of Pb(II), Cd(II), Ni(II) and Cu(II) onto natural kaolinite clay. Desalination 252 (1-3), 33, 2010.
- 49. Kapoor, A. and Viraraghavan, T., 1998, Use of immobilized bentonite in removal of heavy metals from wastewater. J Environ Eng, 124: 1020–1024.
- 50. Karapinar N, Donat R (2009) Adsorption behavior of Cu2? and Cd2? onto natural bentonite. Desalination 249:123–129.
- 51. Kubilay S, Gu[¨]rkan R, Savran A, Sahan T (2007) Removal of Cu (II), Zn (II) and Co (II) ions from aqueous solutions by adsorption onto natural bentonite. Adsorption 13:41–51.
- 52. Kumric' KR, Đuckic' AB, Trtic'-Petrovic' TM, Vukelic' NS, Stojanovic' Z, Grbovic' Novakovic', Matovic' LL (2013) Simultaneous removal of divalent heavy metals from aqueous solutions using raw and mechanochemically treated interstratified montmorillonite/kaolinite clay. Ind Eng Chem Res 52:7930–7939.
- 53. KHALFA L., BAGANE M., CERVERA M.L., NAJJAR S. Competitive Adsorption of Heavy Metals onto Natural and Activated Clay: Equilibrium, Kinetics and Modeling. International Journal of Chemical and Molecular Engineering, 10 (5), 583, 2016.
- 54. Luo T A, Liu X D. The Cd (II) adsorption on modified bentonite properties. Environmental Engineering, 2005, 23(1): 80–81. (In Chinese).
- 55. Lu H J, Luan M T, Zhang J L, Yu Y X. Study on the adsorption of Cr (VI) onto landfill liners containing granular activated carbon or bentonite activated by acid. Journal of China University of Mining & Technology, 2008, 18(1): 125–130.
- 56. Lin, S.H. and Juang, R.S., 2002, Heavy metal removal from water by sorption using surfactantmodified montmorillonite. J Hazard Mater, 92: 315–326.
- 57. Liu, J.F., Cao, C.H. and Jiang, Y.C., 2000, Preliminary study on the dynamics of competitive adsorption of heavy metals in soil. Soil Fertile, 3: 10–15 (In Chinese).

- 58. Langmuir I (1918) The adsorption of gases on plane surfaces of glass, mica and platinum. J Am Chem Soc 40:1361–1403.
- 59. Liu Z, Zhou S (2010) Adsorption of copper and nickel on Na bentonite. Process Saf Environ 88:62–
 66.
- 60. Lehninger, Albert L.; Nelson, David L.; Cox, Michael M. (2000). Principles of Biochemistry (3rd ed.). New York: W. H. Freeman. ..
- 61. Lynch CJ, Adams SH (December 2014). . Nature Reviews. Endocrinology. 10 (12): 723-36.
- 62. LI B., LI L.Y., GRACE J.R. Adsorption and hydraulic conductivity of landfill-leachate perfluorinated compounds in bentonite barrier mixtures. J. Environ. Manage.156, 236, 2015.
- 63. LIU J.F., CAO C.H., JIANG Y.C. Preliminary study on the dynamics of competitive adsorption of heavy metals in soil. Soil Fertil. 3, 10, 2000.
- 64. LIU X., HICHER P., MURESAN B., SAIYOURI N., HICHER P.Y. Heavy metal retention properties of kaolin and bentonite in a wide range of concentration and different pH conditions.Appl. Clay Sci. 119, 365, 2016.
- 65. LIANG S., GUO X., FENG N., Q. TIAN. Isotherms kinetics and thermodynamic studies of adsorption of Cu (II) from aqueous solution by Mg/K type orange peel adsorbents. J Hazard Mat. 174, 756, 2010.
- 66. LIU Z., ZHOU S. Adsorption of copper and nickel on Na bentonite. Process Saf. Environ. 88, 62, 2010.
- 67. Mcbride M B. Environmental Chemistry of Soils. New York: Oxford University Press, 1994.
- 68. Ma, Q.Y., Traina, S.J. and Logan, T.J., 1994, Effects of aqueous Al, Cd, Cu, Fe (II), Ni, and Zn on Pb immobilization by hydroxyapatite. Environ Sci Technol, 28: 1219–1228.
- 69. Mohan, D. and Singh, K.P., 2002, Single-and multi-component adsorption of cadmium and zinc using activated carbon derived from bagasse—an agricultural waste. Water Res, 36: 2304–2318.
- 70. Malamis S, Katsou E (2013) A review on zinc and nickel adsorption on natural and modified zeolite, bentonite and vermiculite: examination of process parameters, kinetics and isotherms. J Hazard Mater 252:428–461.

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- 71. Mishra PC, Patel RK (2009) Removal of lead and zinc ions from water by low-cost adsorbents. J Hazard Mater 168:319–325.
- 72. Mohan D, Singh KP (2002) Single-and multi-component adsorption of cadmium and zinc using activated carbon derived from bagasse-an agricultural waste. Water Res 36:2304–2318.
- 73. Mathews CK (2000). Biochemistry. Van Holde, K. E., Ahern, Kevin G. (3rd ed.). San Francisco, Calif.: Benjamin Cummings. p. 776.
- 74. Marvel CS (1940). 20: 106.; Collective Volume, vol. 3, p. 848.
- 75. MISHRA A. K., OHTSUBO M., LI L., HIGASHI T. Controlling factors of the swelling of various bentonites and their correlations with the hydraulic conductivity of soil-bentonite mixtures. Appl. Clay Sci. 52, 78, 2011. 9.
- 76. MUSSO T.B., FRANCISCA F.M., PETTINARI G., ROEHL K.E. Suitability of a cretaceous natural Nab bentonite as construction material for landfill liners. Environ Eng Manage J. 15 (11), 2519, 2016.
 10.
- 77. MOHELLEBI F., LAKEL F. Adsorption of Zn2+ on Algerian untreated bentonite clay. Desalin. Water Treat. 57 (13), 6051, 2016.
- 78. MAVAKALA B.K., LE FAUCHEUR S., MULAJI C.K., LAFFITE A., DEVARAJAN N., BIEY E.M., GIULIANI G., OTAMONGA J.P., KABATUSUILA P., MPIANA P.T., POTÉ J. Leachates draining from controlled municipal solid waste landfill: Detailed geochemical characterization and toxicity tests. Waste Manage. 55, 238, 2016.
- 79. MUSSO T.B., ROEHL K.E., PETTINARI G., VALLÉS J.M. Assessment of smectite-rich claystones from Northpatagonia for their use as liner materials in landfills. Appl. Clay Sci. 48, 438, 2010.
- 80. MUSSO T.B., PETTINARI G., PAROLO M.E., MESQUÍN L. Smectitic clays from Northpatagonic region of Argentina as hydraulic barriers of landfills and heavy metals retention agents. Rev. Int. Contam. Ambie. 33 (1), 141, 2017.
- 81. MUSSO T.B., PAROLO M.E., PETTINARI G., FRANCISCA F.M. Cu (II) and Zn (II) adsorption capacity of three different clay liner materials. J. Environ. Manage. 146, 50, 2014.

- 82. MOHAN D., SINGH K.P. Single- and multi-component adsorption of cadmium and zinc using activated carbon derived from bagasse-an agricultural waste. Water Res. 36, 2304, 2002.
- 83. MELICHOVÁ Z., HROMADA L. Adsorption of Pb2+ and Cu2+ ions from aqueous solutions on natural bentonite. Pol. J. Environ. Stud. 22 (2), 457, 2013.
- 84. . IUPAC-IUB Joint Commission on Biochemical Nomenclature. 1983. from the original on 9 October 2008. Retrieved 5 March 2018.
- 85. NAVEEN B.P., DURGA MADHAB MAHAPATRA, SITHARAM T.G., SIVAPULLAIAH P.V., RAMACHANDRA T.V. Physico-chemical and biological characterization of urban municipal landfill leachate. Environ. Pollut. 220, 1, 2017.
- 86. PADILLA-ORTEGA E., LEYVA-RAMOS R., FLORESCANO J.V. Binary adsorption of heavy metals from aqueous solution onto natural clays. Chem. Eng. J. 225, 535, 2013.
- 87. PARE S., PERSSON I., GUEL B., LUNDBERG D., ZERBO L., KAM S., TRAORÉ K. Heavy metal removal from aqueous solutions by sorption using natural clays from Burkina Faso. Afr. J. Biotechnol. 11 (45), 10395, 2012.
- 88. & Technology, 2007, 17(3): 345–348. [13] Rawat J P, Ansari A A, Singh R P. Sorption equilibria of lead (II) on some Indian soils—the natural ion exchangers. Colloids and Surfaces, 1990(50): 207– 214.
- 89. ROBINSON T. Removal of toxic metals during biological treatment of landfill leachates. Waste Manage. 63, 299, 2017.
- 90. . REDDAD Z., GÉRENTE C., ANDRÈS Y., RALET M.C., THIBAULT J.F., LE CLOIREC P. Ni
 (II) and Cu (II) binding properties of native and modified sugar beet pulp. Carbohyd. Polym. 49 (1), 23, 2002.
- 91. Saha UK, Taniguchi S, Sakurai K (2002) Simultaneous adsorption of calcium, zinc, and lead on hydroxy aluminum- and hydroxy aluminosilicate-montmorillonite complexes. Soil Sci Soc Am J 66:117–128 Sen TK, Gomez D (2011) Adsorption of zinc (Zn2?) from aqueous solution on natural bentonite. Desalination 267:286–294.

- 92. Seo DC, Yu K, DeLaune RD (2008) Comparison of monometal and multimetal adsorption in Mississippi River alluvial wetland sediment: batch and column experiments. Chemosphere 73:1757– 1764 Shukla A, Zhang YH, Dubey P, Margrave JL,
- 93. Shukla SS (2002) The role of sawdust in the removal of unwanted materials from water. J Hazard Mater 95:137–152.
- 94. Srivastava P, Singh B, Angove M (2005) Competitive adsorption behavior of heavy metals on kaolinite. J Colloid Interf Sci 290:28–38.
- 95. Srivastava VC, Mall ID, Mishra IM (2008) Removal of cadmium (II) and zinc (II) metal ions from binary aqueous solution by rice husk ash. Colloid Surfaces A 312:172–184.
- 96. SDIRI A.T., HIGASHI T., JAMOUSSI F. Adsorption of copper and zinc onto natural clay in single and binary systems. Int. J. Environ. Sci. Technol. 11 (4), 1081, 2014.
- 97. Serrano S, Garrido F, Campbell C G. Competitive sorption of cadmium and lead in acid soils of central Spain. Geoderma, 2005(124): 91–104.
- 98. Saha U M, Iwasaki K, Sakurai K. Desorption behavior of Cd, Zn and Pb sorbed on hydroxy aluminum and hydroxy aluminosilicate-montmorillonite complexes. Clays and Clay Miner, 2003, 51(5): 481–492. [15].
- 99. Sheta A S, Falatah A M, Salam A S H. Sorption characteristic of zinc and iron by natural zeolite and bentonite. Micropor Mesopor Mater, 2003(61): 127–136.
- 100. TURAN N.G., GUMUSEL E.B. Utilizing natural liner materials for heavy metal removal in simulated landfill conditions. Clean, 41 (4), 403, 2013.
- 101. UDDIN M.K. A review on the adsorption of heavy metals by clay minerals, with special focus on the past decade. Chem. Eng. J. 308, 438, 2017.
- 102. VHAHANGWELE M., MUGERA G. W. The potential of ball-milled South African bentonite clay for attenuation of heavy metals from acidic wastewaters: Simultaneous sorption of Co2+, Cu2+, Ni2+, Pb2+, and Zn2+ ions. J. Environ. Chem. Eng. 3, 2416, 2015.
- Veli S, Alyuz B (2007) Adsorption of copper and zinc from aqueous solutions by using natural clay. J Hazard Mater 149:226–233.

45 | Page

- 104. Vieira MGA, Neto AFA, Gimenes ML, da Silva MGC (2010) Sorption kinetics and equilibrium for the removal of nickel ions from aqueous phase on calcined Bofe bentonite clay. J Hazard Mater 177:362–371.
- 105. Wang H K. Treatment of Cu2+-containing wastewater from copper smelter with bentonite. Nonferrous Metals, 2007, 59(1): 108–110. (In Chinese).
- 106. WU X.L., ZHAO D., YANG S.T. Impact of solution chemistry conditions on the sorption behavior of Cu (II) on Lin'an montmorillonite. Desalination 269, 84, 2011.
- 107. Xu, D., Shao, D.D., Chen, C.L., Ren, A.P. and Wang, X.K., 2006, Effect of pH and fulvic acid on sorption and complexation of cobalt onto bare and FA bound MX-80 bentonite. Radiochim Acta, 94: 97–102.
- 108. XU D., ZHOU X., WANG X. Adsorption and desorption of Ni2+ on Na-montmorillonite: Effect of pH, ionic strength, fulvic acid, humic acid and addition sequences. Appl. Clay Sci. 39, 133, 2008.
- 109. Yang S, Li J, Lu Y, Chen Y, Wang X (2009) Sorption of Ni (II) on GMZ bentonite: effects of pH, ionic strength, foreign ions, humic acid and temperature. Appl Radiat Isotopes 67:1600–1608.
- 110. Yang S, Zhao D, Zhang H, Lu S, Chen L, Yu X (2010) Impact of environmental conditions on the sorption behavior of Pb (II) in Na-bentonite suspensions. J Hazard Mater 183:632–640.
- 111. ZAKI A.A., AHMAD M.I., ABD EL-RAHMAN K.M. Sorption characteristics of a landfill clay soil as a retardation barrier of some heavy metals. Appl. Clay Sci. 135, 150, 2017.
- 112. Zhang H, Tong Z, Wei T, Tang Y (2011) Removal characteristics of Zn (II) from aqueous solution by alkaline Ca-bentonite. Desalination 276:103–10.
- 113. Zaitan, H., Bianchi, D., Achak, O. and Chafik, T., 2008, A comparative study of the adsorption and desorption of o-xylene onto bentonite clay and alumina. J Hazard Mater, 153: 852–859.
- Zhu, L.Z., Ren, X.G. and Yu, S.B., 1998, Use of cetyltrimethylammonium bromide-bentonite to remove organic contaminants of varying polar character from water. Environ Sci Technol, 32: 3374– 3378.

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Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

General Parameters

Method : Iron (Fe) Autosampler : None Use SFI: No Operator : Jyothi

Instrument Mode: Flame *Dilution:* None

mA Hours

n/a

Report Date: 11-06-2022 17:40:20

Valid Method Signatures

11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Method Audit Trail

11-06-2022 17:22:18 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
11-06-2022 17:22:33 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:23:24 parjanya(Parjanya):DESKTOP-39TDEGC

Signed with Reason : Approved by:

Analysis Details

Spectrometer: iCE 3000 AA01204906 v1.30

Analysis Name: Iron (Fe) 11-06-2022 Operator Name: Jyothi

n/a

Lamp Information Serial Number

Element(s) Fe

. . .

Deuterium Lamp Hours: 68.34

Sequence Table

Shared Standards: Yes	
Action	Fe
Calibration	√
B.No.Group-II	√
B.No.Group-III	√
B.No.Group-IV	√
B.No.Group-V	√
B.No.Group-VI	✓

Sample Details

		Nominal Mass: 1.0000				
No.	Sample Id	Sample Mass	Dilution Ratio			
1	B.No.Group-II	1.0000	1.0000			
2	B.No.Group-III	1.0000	1.0000			
3	B.No.Group-IV	1.0000	1.0000			
4	B.No.Group-V	1.0000	1.0000			
5	B.No.Group-VI	1.0000	1.0000			

Valid Analysis Signatures

11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

Analysis Audit Trail

- 11-06-2022 17:30:16 Jyothi(M. Jyothi):DESKTOP-39TDEGC Record created
- 11-06-2022 17:38:27 Jyothi(M. Jyothi):DESKTOP-39TDEGC Error MD147 - Activity manually aborted by user.
- 11-06-2022 17:38:41 jyothi(M. Jyothi):DESKTOP-39TDEGC Signed with Reason : Analysed by:
- 11-06-2022 17:39:33 parjanya(Parjanya):DESKTOP-39TDEGC Signed with Reason : Approved by:

MART SPECIALITIES LAB LLP.

Operator Name: Jyothi Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Spectrometer Parameters - Fe

Element: Fe Wavelength: 248.3nm Background Correction: D2 Signal Type: Continuous Measurement Time: 4.0secs Use RSD Test: No

Flame Type: Air-C2H2 Nebuliser Uptake: 4secs Burner Height: 7.0mm

Sampling: None

Calibration Mode: Normal	
Concentration Units: mg/L	
Acceptable Fit: 0.990	
Standard 1	2.0000
Standard 2	4.0000
Standard 3	6.0000

No changes are recorded for this element

Measurement Mode: Absorbance Bandpass: 0.2nm High Resolution: Off Resamples: Fast Flier Mode: No

Lamp Current: 75% Optimise Spectrometer Parameters: No Number Of Resamples: 3

Flame Parameters - Fe

Fuel Flow: 0.9L/min Burner Stabilisation: 0mins Optimise Burner Height: No Auxilary Oxidant: Off Optimise Fuel Flow: No

Sampling Parameters - Fe

Calibration Parameters - Fe

Line Fit: Linear Scaled Units: mg/L Rescale Limit: 10.0% Standard 4 Standard 5 Use Stored Calibration: No Scaling Factor: 1.0000 Failure Action: Flag and Continue 8.0000 10.0000

Element Audit Trail - Fe

Solution Results - Fe

Y = 0.04924x + 0.0141 Fit: 0.9965 Characteristic Conc: 0.0894

> 1 2 3

> 1 2 3

> 1 2 3

> 1 2 3

1 2

3

Sample ID

Fe Blank

Fe Standard 1

Fe Standard 2

Fe Standard 3

Fe Standard 4

	4				Norma	I: Linea	ar Fit				
	Abs		*		*		*		-*-		*
	0 * -0-0	1	2	3	4	5	6	7	8	9	10
	-0-0		2	0		nc:mg/		,	0	0	10
Signal	Rsd	Conc				cted C					
Abs	%	mg/L			mg/L						
0.001	35.6	0.000)		Ũ						
0.001	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:26	i		
0.001	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:30)		
0.000	Backgro	ound: -0.0	03			11-06	-2022 1	7:31:35			
0.120	0.3	2.0000)								
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:05			
0.119	-	ound: 0.00						7:32:09			
0.120	Backgro	ound: 0.00)3			11-06	-2022 1	7:32:13			
0.215	0.3	4.0000)								
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:40)		
0.215	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:45			
0.216	Backgro	ound: 0.00)4			11-06	-2022 1	7:32:49)		
0.322	0.2	6.0000)								
0.323	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:19)		
0.322	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:23			
0.322	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:27			
0.411	0.3	8.0000)								
0.410	Backgro	ound: 0.00)5			11-06	-2022 1	7:33:59)		
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:03			
0.412	Backgro	ound: 0.00)5			11-06	-2022 1	7:34:07	•		

MART SPECIALITIES LAB LLP.

Operator Name: Jyothi

Results File: E:\AAS System Data\2022\JUNE\11 06 2022\TARA 0064 2-6\Iron (Fe)1.SLR

Solution Results - Fe

Sample ID	Signal	Rsd	Conc	Corrected Conc
-	Abs	%	mg/L	mg/L
Fe Standard 5	0.493	0.1	10.0000	
1	0.494	Backg	ground: 0.005	11-06-2022 17:34:38
2	0.494	Backg	ground: 0.005	11-06-2022 17:34:42
3	0.493	Backg	ground: 0.006	11-06-2022 17:34:46
Fe B.No.Group-II	0.834	0.1	16.6490 C	16.6490 C
1	0.834	Backg	ground: 0.005	11-06-2022 17:35:14
2	0.833	Backg	ground: 0.005	11-06-2022 17:35:19
3	0.834	Backg	ground: 0.005	11-06-2022 17:35:23
Fe B.No.Group-III	0.759	0.1	15.1286 C	15.1286 C
1	0.758	Backg	ground: 0.003	11-06-2022 17:35:52
2	0.760	Backg	ground: 0.003	11-06-2022 17:35:56
3	0.758	Backg	ground: 0.004	11-06-2022 17:36:00
Fe B.No.Group-IV	0.645	0.3	12.8189 C	12.8189 C
1	0.647	Backg	ground: 0.003	11-06-2022 17:36:31
2	0.644	Backg	ground: 0.003	11-06-2022 17:36:35
3	0.645	Backg	ground: 0.003	11-06-2022 17:36:40
Fe B.No.Group-V	0.809	0.2	16.1488 C	16.1488 C
1	0.808	Backg	ground: 0.005	11-06-2022 17:37:11
2	0.809	Backg	ground: 0.005	11-06-2022 17:37:15
3	0.811	Backg	ground: 0.005	11-06-2022 17:37:20
Fe B.No.Group-VI	0.711	0.3	14.1597 C	14.1597 C
1	0.709	Backg	ground: 0.005	11-06-2022 17:37:55
2	0.713	Backg	ground: 0.005	11-06-2022 17:37:59
3	0.712	Backg	ground: 0.004	11-06-2022 17:38:03

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-1 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
I	DETAILS OF THE SAMPLE
Sample Name : NA	
Name of the Manufacturer : NA	
Storage condition To b	up-I Mfg. : NA Exp. : NA e stored at room date date
	Derature Batch : NA C±3°C size
	by AAS Analysis.
	6/2022
Analysis Completion Date : 11/0	6/2022
Mfg. License No. : Not	provided
A.R.NO : NA	The Andrew Rest

Test Results

S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	620ppm	

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



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Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-2 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
Q	DETAILS OF THE SAMPLE
Sample Name : N	A
Name of the Manufacturer : N	A
Storage condition : To	roup-II Mfg. : NA Exp. : NA be stored at room date date
Room Temperature : 25	Batch : NA size
Tests Required:IreMethod:NAnalysis Starting Date:11	on by AAS Analysis.
	ot provided
A.R.NO : N	A

Test Results

S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	832.5ppm	

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



OCA Approval No: 05/ML/TS/2020/G	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinaga Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Sample Received Date : 11/06/2022
a.u.	DETAILS OF THE SAMPLE
Sample Name : N	JA
Name of the Manufacturer : N	JA
Storage condition	Group-III Mfg. : NA Exp. : NA To be stored at room date date Cemperature Batch : NA
	5°C±3°C size
	gm /
Method : I Analysis Startin <mark>g Date : I</mark>	ron by AAS Analysis. NA 1/06/2022 1/06/2022
and the second sec	Not provided
A.R.NO	NA
Remark : S	Sample analyzed as received

Test Results

S.No.	Test Parameter	Result
01.	Iron by AAS Analysis: (ppm)	756.5ppm

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Authorized Signatory

(Dr.R.Marayya)

Office: 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile: 98481 98350, 98481 25019 Mail: rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College	
Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000	Sample Received Date: 11/06/2022Report Date: 13/06/2022
Contact Person: Dr. Abhijeet Contact Number :9502344392	
	DETAILS OF THE SAMPLE
Sample Name : N	A
Name of the Manufacturer : N	A
Storage condition : T	roup-IV Mfg. : NA Exp. : NA o be stored at room date date emperature Batch : NA
	5°C±3°C size
Method : N Analysis Starting Date : 1	on by AAS Analysis. A 1/06/2022 1/06/2022
Mfg. License <mark>No. :</mark> N	ot provided
A.R.NO : N	A
Remark : Sa	ample analyzed as received

Test Results

	and the second se	An alternation	
S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	641ppm	

Authorized Signatory

(Dr.R.Marayya)

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G				
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00			
Name & Address of the Custome	r: Reference / Report No. : MSL/2022/JUNE/TARA/0064-5			
Tara Government College	Sample Received Date : 11/06/2022			
Prashanth Nagar Colony, Balajinag	ar Sumpto Received 2 mil			
Sangareddy Telangana. 502000	Report Date : 13/06/2022			
Contact Person: Dr. Abhijeet Contact Number :9502344392				
	DETAILS OF THE SAMPLE			
Sample Name :	NA			
Name of the Manufacturer :	NA			
Batch no : Storage condition	Group-VMfg. : NAExp. : NATo be stored at roomdatedateTemperatureBatch : NA			
Room Temperature :	25°C±3°C size			
Quantity Received	4gm			
Tests Required:Method:Analysis Starting Date:	Iron by AAS Analysis. NA 11/06/2022			
Analysis Completion Date :	11/06/2022			
Mfg. License No.	Not provided NA			
A.R.NO Remark	Sample analyzed as received			

Test Results

S.No.	Test Parameter	Result
01. Ir	on by AAS Analysis: (ppm)	807.5ppm

>

Authorized Signatory

(Dr.R.Marayya)

Office: 13-6-438/A/83, Sri Satyanarayana Nagar, Mehdipatnam, Hyderabad - 500 028, Telangana State, India Mobile: 98481 98350, 98481 25019 Mail: rmarayya@yahoo.com

Work : Road No. : 03, Plot No.: 31, Ground Floor & 1st Floor, Gajularamaram, ALEAP Industrial Area, Pragathi Nagar, Medhcal, Medchal-Malkajgiri - 500 055, Telangana, India, **Tel.** : 85007 98350, 81423 98350 & 98481 98350 **Mail**:martspecialities@gmail.com



DCA Approval No: 05/ML/TS/2020/G	
	CERTIFICATE OF ANALYSIS MSL/QA/017-03/F07-00
Name & Address of the Customer: Tara Government College Prashanth Nagar Colony, Balajinagar Sangareddy Telangana. 502000 Contact Person: Dr. Abhijeet Contact Number :9502344392	Reference / Report No. : MSL/2022/JUNE/TARA/0064-6 Sample Received Date : 11/06/2022 Report Date : 13/06/2022
1	DETAILS OF THE SAMPLE
Storage condition	up-V1 Mfg. : NA Exp. : NA be stored at room date date perature Batch : NA C±3°C size
Tests Required:IronMethod:NAAnalysis Starting Date:11/0Analysis Completion Date:11/0Mfg. License No.:NotA.R.NO:NA	by AAS Analysis. 6/2022 6/2022 provided

Test Results

S.No.	Test Parameter	Result	
01.	Iron by AAS Analysis: (ppm)	708ppm	

*

Authorized Signatory

(Dr.R.Marayya)

Field Visit to Mart Specialities Lab Pvt. Ltd, Hyderabad For "Training and Hands-on Experience on Atomic Absorption Spectroscopy"



DEPARTMENT OF CHEMISTRY



TARA GOVERNMENT COLLEGE, SANGAREDDY (A) 2021-22



DEPARTMENT OF CHEMISTRY Tara Govt. College, Sangareddy(A)

Field Visit to Mart Specialities Lab Pvt. Ltd, Hyderabad For "Training and Hands-on Experience on Atomic Absorption Spectroscopy"

Department of chemistry has organized Field Visit to MART Specialities Lab Pvt. Ltd., Hyderabad for the final year students of B.Sc. Chemistry on 11.06.2022 to train the student in Atomic Absorption Spectroscopy (AAS) to assess the heavy metal content at ppm levels. Six (6) students of B.Sc. Chemistry Final year student were perfectly trained in this programme. Dr.Abhijit Kantankar, Head, Department of Chemistry acted as Coordinator of the Filed Visit.

Objective of the Programme:

To provide Scientific and Research knowledge to the students to enhance their research skills to carry out assigned research projects.

Context of the Programme:

Field Visit to Scientific and Research Institutes provides positive effect on both teachers and students in the cognitive and affective aspects. It is one of the most effective and the best way of teaching which provides the elements of scientific inquiry. Even the faculty will have the chance to develop their teaching abilities, including the capacity to make complex subjects understandable to people who aren't academics. Students have a stronger comprehension of the research elements in their field of study as a result of the procedure. Additionally, the positive experience of taking part in the process influenced their desire to take part in subsequent outreach initiatives. Implementing the Field visit provides a complex experience of learning.

In this regard, Department of chemistry has arranged a Field visit to **MART Specilaities Lab Pvt. Ltd, Hyderabad** for **Real time learning and hands-on training** on Atomic Absorption Spectroscopy with **THERMO Scientific iCE-3000** equipment which is essential to carry out their allotted Research Projects on Metal adsorption studies. This Field Visit was arranged as part of **POST-MoU activity** as MART Specilaities Lab Pvt. Ltd, Hyderabad and Department of Chemistry, Tara Govt. College, Sangareddy(A) reached MoU on 31.12.2021.

Students Trained in the Programme:

S.No.	Name of the Student	Roll Number	Group	Year
1	S.Shirisha	6058-19-578-029	B.Sc.(MCCs)	III
2	Mukthi Kanth Rout	6058-19-578-022	B.Sc.(MCCs)	III
3	N.Shiva Shankar	6058-19-578-025	B.Sc.(MCCs)	III
4	Y. Avinash Reddy	6058-19-578-038	B.Sc.(MCCs)	III
5	T.Chandana	6058-19-578-032	B.Sc.(MCCs)	III
6	K.Dheeraj Kumar	6058-19-578-017	B.Sc.(MCCs)	III

TRAINING MANUAL

Atomic Absorption Spectrometry

Atomic absorption spectrometry (AAS) is an analytical technique that measures the concentrations of elements. Atomic absorption is so sensitive that it can measure down parts per billion of a gram (μ g dm⁻³) in a sample. The technique makes use of the wavelengths of light specifically absorbed by an element. They correspond to the energies needed to promote electrons from one energy level to another, higher, energy level.

Atomic absorption spectrometry has many uses in different areas of chemistry.

Clinical analysis: Analysing metals in biological fluids such as blood and urine.

Environmental analysis: Monitoring our environment- eg finding out the levels of various elements in rivers, seawater, drinking water, air, petrol and drinks such as wine, beer and fruit drinks.

Pharmaceuticals: In some pharmaceutical manufacturing processes, minute quantities of a catalyst used in the process (usually a metal) are sometimes present in the final product. By using AAS the amount of catalyst present can be determined.

Industry: Many raw materials are examined and AAS is widely used to check that the major elements are present and that toxic impurities are lower than specified- eg in concrete, where calcium is a major constituent, the lead level should be low because it is toxic.

Mining: By using AAS the amount of metals such as gold in rocks can be determined to see whether it is worth mining the rocks to extract the gold.

How it works

Atoms of different elements absorb characteristic wavelengths of light. Analysing a sample to see if it contains a particular element means using light from that element. For example with lead, a lamp containing lead emits light from excited lead atoms that produce the right mix of wavelengths to be absorbed by any lead atoms from the sample. In AAS, the sample is atomized- i.e. converted into ground state free atoms in the vapour state- and a beam of electromagnetic radiation emitted from excited lead atoms is passed through the vaporized sample. Some of the radiation is absorbed by the lead atoms in the sample. The greater the number of atoms there is in the vapour, the more radiation is absorbed. The amount of light absorbed is proportional to the number of lead atoms. A calibration curve is constructed by running several samples of known lead concentration under the same conditions as the unknown. The amount the standard absorbs is compared with the calibration curve and this enables the calculation of the lead concentration in the unknown sample.

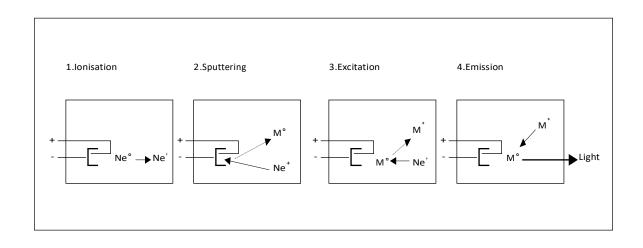
Consequently an atomic absorption spectrometer needs the following three components: a light source; a sample cell to produce gaseous atoms; and a means of measuring the specific light absorbed.

The light source

The common source of light is a 'hollow cathode lamp'. This contains a tungsten anode and a cylindrical hollow cathode made of the element to be determined. These are sealed in a glass tube filled with an inert gas- e.g neon or argon- at a pressure of between 1 Nm⁻² and 5 Nm⁻².

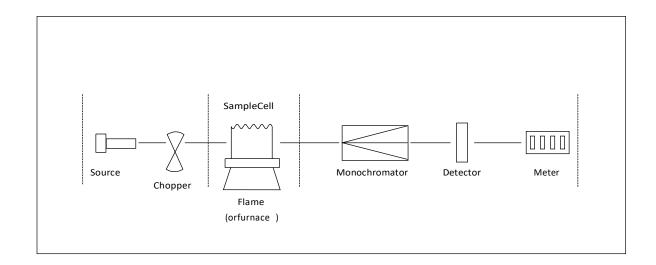


The ionization of some gas atoms occurs by applying a potential difference of about 300-400V between the anode and the cathode. These gaseous ions bombard the cathode and eject metal atoms from the cathode in a process called sputtering. Some sputtered atoms are in excited states and emit radiation characteristic of the metal as they fall back to the ground state $-egPb^* \rightarrow Pb + h v$. The shape of the cathode concentrates the radiation into a beam which passes through a quartz window, and the shape of the lamp is such that most of the sputtered atoms are redeposited on the cathode. A typical atomic absorption instrument holds several lamps each for a different element. The lamps are housed in a rotating turret so that the correct lamp can be quickly selected.



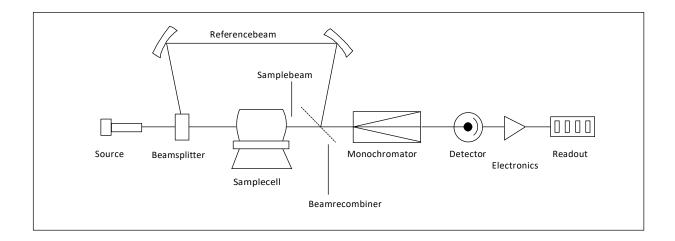
The optical system and detector

A monochromator is used to select the specific wavelength of light –ie spectral line – which is absorbed by the sample, and to exclude other wavelengths. The selection of the specific light allows the determination of the selected element in the presence of others. The light selected by the monochromator is directed onto a detector that is typically a photomultiplier tube. This produces an electrical signal proportional to the light intensity



Double beam spectrometers

Modern spectrometers incorporate a beam splitter so that one part of the beam passes through the sample cell and the other is the reference. The intensity of the light source may not stay constant during an analysis. If only a single beam is used to pass through the atom cell, a blank reading containing no analyte (substance to be analysed) would have to be taken first, setting the absorbance at zero. If the intensity of the source changes by the time the sample is put in place, the measurement will be inaccurate. In the double beam instrument there is a constant monitoring between the reference beam and the light source. To ensure that the spectrum does not suffer from loss of sensitivity, the beam splitter is designed so that as high a proportion as possible of the energy of the lamp beam passes through the sample.



Atomisation of the sample

Two systems are commonly used to produce atoms from the sample. Aspiration involves sucking asolution of the sample into a flame; and electrothermal atomisation is where a drop of sample is placed into a graphite tube that is then heated electrically.

Some instruments have both atomisation systems but share one set of lamps. Once the appropriate lamp has been selected, it is pointed towards one or other atomisation system.

Flame aspiration

Ethyne/air (giving a flame with a temperature of 2200–2400°C) or ethyne/dinitrogen oxide (2600– 2800°C) are often used. A flexible capillary tube connects the solution to the nebuliser. At the tip of the capillary, the solution is 'nebulised' –ie broken into small drops. The larger drops fall out and drain off while smaller ones vaporise in the flame. Only ca 1% of the sample is nebulised.

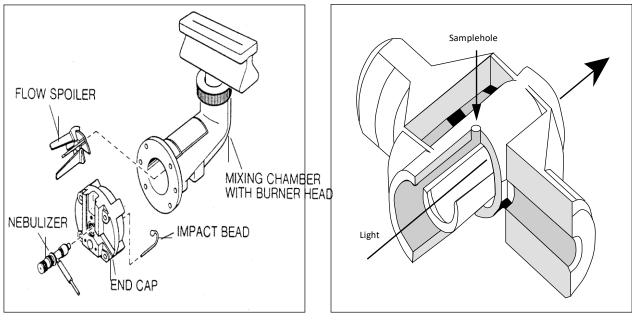


Figure 1

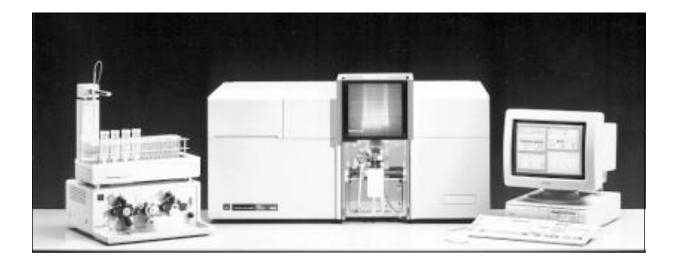
Figure 2

Electro-thermal atomization

 $25 \ \mu$ l of sample (ca 1/100th of a raindrop) is placed through the sample hole and onto the platform from an automated micropipette and sample changer. The tube is heated electrically by passing a current through it in a pre-programmed series of steps. The details will vary with the sample but typically they might be 30–40 seconds at 150°C to evaporate the solvent, 30 seconds at 600°C to drive off any volatile organic material and char the sample to ash, and with a very fast heating rate (ca 1500 °C s⁻¹) to 2000-2500°C for 5–10 seconds to vaporise and atomise elements (including the element being analysed). Finally heating the tube to a still higher temperature –ca 2700°C – cleans it ready for the next sample. During this heating cycle the graphite tube is flushed with argon gas to prevent the tube burning away. In electrothermal atomisation almost 100% of the sample is atomised. This makes the technique much more sensitive than flame AAS.

Sample preparation

Sample preparation is often simple, and the chemical form of the element is usually unimportant. This is because atomisation converts the sample into free atoms irrespective of its initial state. The sample is weighed and made into a solution by suitable dilution. Elements in biological fluids such as urine and blood are often measured simply after a dilution of the original sample.



When making reference solutions of the element under analysis, for calibration, the chemical environment of the sample should be matched as closely as possible –i.e. the analyte should be in the same compound and the same solvent. Teflon containers may be used when analyzing very dilute aolutions because elements such as lead are sometimes leached out of glass vessels and can affect the results

Background absorption

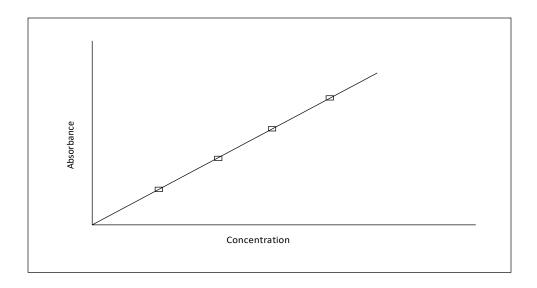
It is possible that other atoms or molecules apart from those of the element being determined will absorb or scatter some radiation from the light source. These species could include unvaporised solvent droplets, or compounds of the matrix (chemical species, such as anions, that tend to accompany the metals being analysed) that are not removed completely. This means that there is a background absorption as well as that of the sample.

One way of measuring and correcting this background absorption is to use two light sources, one of which is the hollow cathode lamp appropriate to the element being measured. The second light source is a deuterium lamp.

The deuterium lamp produces broad band radiation, not specific spectral lines as with a hollow cathode lamp. By alternating the measurements of the two light sources – generally at 50 -100 Hz – the total absorption (absorption due to analyte atoms plus background) is measured with the specific light from the hollow cathode lamp and the background absorption is measured with the light from the deuterium lamp. Subtracting the background from the total absorption gives the absorption arising from only analyte atoms.

Calibration

A calibration curve is used to determine the unknown concentration of an element -eg lead – in a solution. The instrument is calibrated using several solutions of known concentrations. A calibration curve is produced which is continually rescaled as more concentrated solutions are used – the more concentrated solutions absorb more radiation up to a certain absorbance. The calibration curve shows the concentration against the amount of radiation absorbed in the given figure.(a) The sample solution is fed into the instrument and the unknown concentration of the element-e.g. lead- is then displayed on the calibration curve given in the below figure.(b)





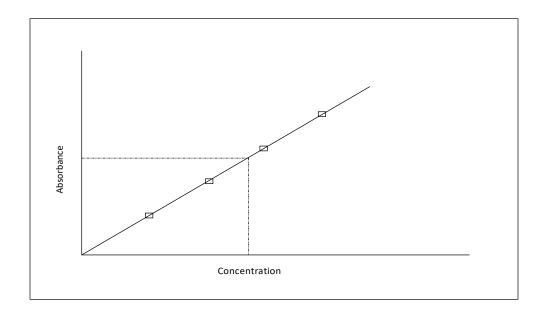


Figure (b)

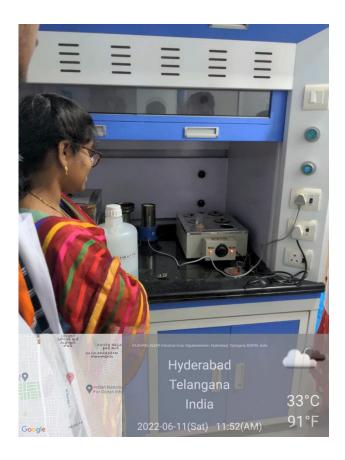
Interferences and matrix modification Other chemicals that are present in the sample may affect the atomisation process. For example, in flame atomic absorption, phosphate ions may react with calcium ions to form calcium pyrophosphate. This does not dissociate in the flame and therefore results in a low reading for calcium. This problem is avoided by adding different reagents to the sample that may react with the phosphate to give a more volatile compound that is dissociated easily. Lanthanum nitrate solution is added to samples containing calcium to tie up the phosphate and to allow the calcium to be atomised, making the calcium absorbance independent of the amount of phosphate. With electrothermal atomisation, chemical modifiers can be added which react with an interfering substance in the sample to make it more volatile than the analyte compound. This volatile component vaporises at a relatively low temperature and is removed during the low and medium temperature stages of electrothermal atomisation.

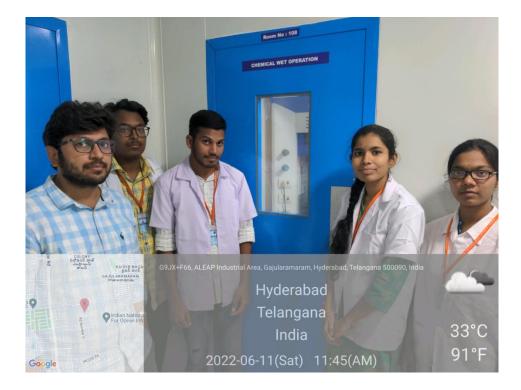
COURTESY: The Education Department, The Royal Society of Chemistry, Burlington House, Piccadilly, London W1J OBA. (The Royal Society of Chemistry Fine Chemicals and Medicinals Group).



Atomic absorption spectrometer (THERMO Scientific iCE-3000)

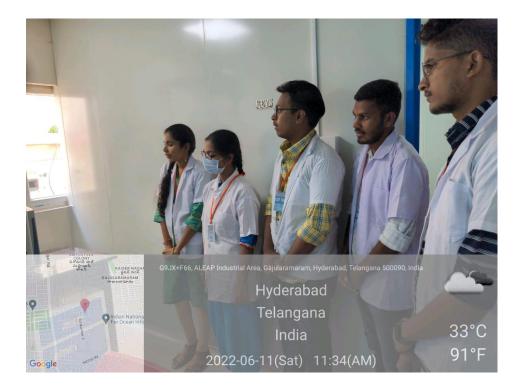




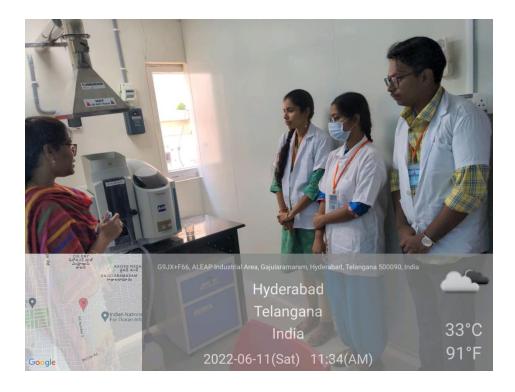
















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TARA GOVERNMENT DEGREE COLLEGE

SANGAREDDY

UROOJ UNNISA SANADHI LAXMAN AMREEN U. NAVNEETHA

SANADILAKSHMAN321@GMAIL.COM

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Executive Summary

RTC Buses are the most widely-used form of public transport in the India serving cities as well as suburban and rural areas. They are also the most cost-efficient and flexible form of public transport, When you consider that, most people use private transport whether it maybe Ola , uber ,Autorikshwa or every other private transport due to motives we don't realize when will the bus comes , loss of endurance or to avoid crowd adventure to conquer these problems we are able to inbuit GPS tracker in each bus to offer the precise place of required bus for passengers and additionally QR scanner based ticket purchase.

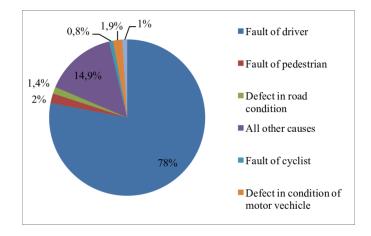
Business Description

To increase the revenues for public transport system ie. RTC buses by imparting beneficial services to passengers.

Many passengers shifting from public transport to private transport and self transport due to they don't realize when will the bus comes .To overcome the issues facing with the aid of passengers through waiting lengthy hours in bus stops for bus , with the help of Gps tracker on each bus to locate the proper location of bus correctly in the Smartphone itself. Additionally the organization provides Qr scanner based ticket purchasing and They might provide brief occupancy of bus on every stop on Google map app or Rtc app itself. No person will pick out non-public shipping in the event that they know while will bus comes.

Mission

Our motive is that passengers should be provided with most cost-efficient and flexible form of public transport with safety. Many accidents on avenue takes place because of self riding automobiles



Services Line

Service Offerings

QR SCANNER BASED TICKET PURCHASE -

The bus conductor will provided with QR scanner and passengers will scan and pay to buy ticket, and give offers and discounts to attract customers .it will be a great move to attract customers in these digital India . Our reason to introduce these QR scanner based ticket purchasing is to remedy the change issues while buying ticket hence many passengers are using digital currency usually. A few customers might not familiar with digital money therefore they are able to use traditional method to buy ticket .

OCCUPANCY OF BUS;

They might provide brief occupancy of bus on every stop on app itself, by that passenger could decide whether to board bus or not.

Market Analysis

Target Market

To draw the passengers who use personal or private transport in place of public transport via offering with low-cost ticket and services . And to create habitatual for each passengers to use public delivery in an effort to create extremely good effect on self riding fuel consumption and additionally controlling of accident rate due to self riding through preserve sources including petroleum sustainably.

Competitor Analysis

Ola, uber and a few different personal buses had been taking half of the percentage of transport market in India they're driven with huge marketplace shares due to the services and facilities they providing.

- Comparative Strength most cost-efficient and flexible form and safety.
- Comparative Weakness fail to provide attractive services to passengers.

Competitor Name	Strength(s)	Weakness(es)
Private buses	services	High ticket price
Taxis	Exact point pickup	High charges
Autorikshaw	Exact drop point	Not safety

Marketing Plan

The cost efficient ticket prices and protection transport itself is tremendous pass to attract passengers.

Financial Considerations

RTC revenue sees 1.5 cr per day according to Times of India ; march 14 2020 statistics .

Startup Costs

When you consider that in Telangana itself their are 9,384 TSRTC buses present on account that every bus have to be in-built with GPS tracking device to offer uninterrupted carrier for the passenger and one unit of GPS tracking device cost almost 1500 rupees.

Funding Requirements

To start with it is required 10,00,000 amount to experiment upon the 10 percentage of TSRTC buses to see the result.

Perhaps, the built in of GPS tracking device in every TSRTC bus is more money ingesting process subsequently, these will circuitously generate big revenues to TSRTC through their service to passengers indirectly by shifting private and self transport passengers to public transport. The total project fee is around 1 day revenue of TSRTC.

Appendix

• Floor plans



The above attached image depicts the similar interface of our service in google maps and TSRTC app to locate the precise place of required bus for passengers in smartphone .



CERTIFICATE OF APPRECIATION

This is to certify that K SREEDHAR of Tara Government College Sangareddy (A) had undergone training and facilitated in implementationing of the program, Youth for Social Impact - 2022 in their college which was organised across the state of Telangana by TSIC, Inqui-lab Foundation and UNICEF.

DR. SHANTA THOUTAM Chief Innovatin Officer Government of Telangana



VIVEK PIDDEMPALLY Co-founder Inqui-Lab Foundation



Maital Rusclia

MS. MEITAL RUSDIA Chief, UNICEF office for Andhra Pradesh, Karnataka and Telangana.





भारतीय प्रौद्योगिकी संस्थान मुंबई पवई, मुंबई-400 076, भारत

Indian Institute of Technology Bombay Powai, Mumbai-400 076, India

दूरभाष/Phone	;	(+91-22) 2572 2545
फैक्स/Fax	:	(+91-22) 2572 3480
वेबसाईट/Website	:	www.iitb.ac.in-



LETTER OF ASSOCIATION

Ref.No. STIITB/2022/060

Date - 21/09/2022

To, The Principal, Tara Government College (Autonomous), Sangareddy, Prashanth Nagar Colony, Sangareddy Dist. Telangana - 502001.

Your association is highly appreciable. We are happy to announce the continue ASSOCIATION of Knowledge Partner Spoken Tutorial Program, IIT Bombay with **Tara Government College** (Autonomous), Sangareddy. Your College is officially partnered as an <u>Academic Partner of IIT</u> <u>Bombay Spoken Tutorial (Academic year February 2022-23)</u>. We appreciate the efforts that College took and trained 494 students in the last semester.

The Program is a part of the National Mission on Education through I CT, <u>MoE, Govt. of India</u>, to spread IT Literacy all over India. We are promoting the learning and usage of Free & Open Source Software (FOSS), through an Audio-Video teaching tool, viz, 'Spoken Tutorial'.

Looking forward to see the same successful working relationship and keen to see **Tara Government College (Autonomous), Sangareddy.** as a model in the matter of introducing the ICT MOOCs, FOSS based training in **Telangana** via ST IIT Bombay.

For and On behalf of Spoken Tutorials, Indian Institute of Technology, Bombay



Mrs. Akanksha Saini National Coordinator Spoken Tutorial Project, IIT Bombay