### Shetkari Shikshan Prasarak Mandal's

## Krishna Mahavidyalaya, Rethare Bk

IQAC 2019-2020

## ACTIVITY REPORT PHYSICS DEPARTMENT

#### IQAC ACTIVITY No:

DATE	FACULTY	DEPARTMENT/COMMITTEE	COORDINATOR NAME		
Throughout the academic year	Science	Physics	Dr. Dhanaji S. Dalavi		
TIME	VENUE	NUMBER OF PARTICIPANTS	NATURE: Outdoor/Indoor		
	Physics laboratory	04 students+01 teacher	Indoor		
SUPPORT/ASSISTANCE: Nil					
BRIEF INFORMATION A	BOUT THE ACT	IVITY (CRITERION NO ):			

TOPIC/SUBJECT OF THE ACTIVITY	"Lead college Activity-Research Project: "Synthesis and characterization of WO3 thin film by sol-gel method for smart window application."
OBJECTIVES	to analyze a scientific occurrence with an investigation or to solve a problem with an invention.
METHODOLOGY	Experimental work
OUTCOMES	students become active, engaged learners. It also helped students to develop independent critical thinking skills.

## PROOFS & DOCUMENTS ATTACHED (Tick mark the proofs attached):

Notice &     Letters	2. Student list of	3. Activity report	4. Photos	5. Feedback form
6. Feedback analysis	7. News clip with details	8. Certificate	9. Any other	10.

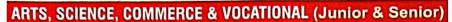
IQAC CELL ACTIV	VITY NUMBER:		
NAME OF TEACHER & SIGNATURE	NAME OF HEAD/ COMMITTEE INCHARGE & SIGNATURE	PRINCIPALS SIGNATURE	IQAC COORDINATOR (SEAL & SIGNATURE)
Dr. Dhanaji S.	Dr. Dhanaji S. Dalavi		
Dalavi <u>Falan'</u> HE	(Salan)	Produ	IQAC, Coordinator,
DEPARTMENT KRISHNA MAN	OF PHYSICS	Principal	KRISHNA MAHAVIDYALAYA, Rethare Bk; Shivnagar - 415108
Rethare Bk; Shi	TAVIDYALAYA, Krishna Mah Vnagar - 415108 Tal, Ka	avidya aya, Rethard rad : 415 108 (MS)	Bk, Tal. Karad, Dist. Satara
		1	

रयत शिक्षण संस्था,सातारा १९१९-२०१

## SADGURU GADAGE MAHARAJ COLLEGE, KARAD

**VIDYANAGAR**, Pin - 415 124, Dist. Satara (M.S.) INDIA P.O. Box No.3 **Ph.** Office: (02164) 271346 Resi. (02164) 271794 **Fax.** (02164) 271346

Website: www.sgm.edu.in E-mail: sgmkarad@yahoo.com



• Principal : Dr. Mohan Rajmane м.Sc., Ph.D.

Ref. No. 1689 19-20

Date:

01/11/2019

To

The Principal, Krishna Mahavidyalaya Rethare (Bk.) Tal.Karad, Dist. Satara.

Sub.: Submission of Research Project under Research Promotion Activity ofr College Students.

I am pleased to inform you that the Research Project under the Research Promotion Activity of the Shivaji University, Kolhapur is received with following details.

Name of Project Advisor	Name of Students	Title of Project	Amount Estimate Rs.
Dr. D. S. Dalavi	1) Lad Divya Dilip	Synthesis,	
	2) Mohini Rajendra Harale	Characterization of WO3 thin film by sol-gel route for electrochromic smart	10,000/-
	3) Ashlesha Arun Patil	windows application	
	4) Aniket Anil Dmame	e e	

Financial assistance under this scheme is subject to final approval and directions of the university.

Thank you,

Yours faithfully,

Principal Lead College,

Sadguru Gadage Maharaj College,

KARAD

De Dalari D.S.

Krishna Mahavidualaua. Shivnadar Haward No. 946, ाचार्य :

ॉ. मोहन राजमाने

एम.एस्सी., पीएच्.डी.

''स्वावलंबी शिक्षण हेच आमचे ब्रीद''-कर्पवीर

रयत शिक्षण संस्थेचे,

## महाराज कालंज, कराड

रवायत्त महाविद्यालय – शिवाजी विद्यापीठ, कोल्हापूर संलग्नित

विद्यानगर, कराड पिन - ४१५ १२४ जि. सातारा (महाराष्ट्र) पो. ऑ. बॉ. नं. ३ फोन: कार्यालय: (०२१६४) २७१३४६ फॅक्स: (०२१६४) २७१३४६ Website: www.sgm.edu.in E-mail: sgmkarad@yahoo.com

कला, विज्ञान, वाणिज्य व व्यवसाय शिक्षण (कनिष्ठ व वरिष्ठ))

नॅक मानांकन : A<sup>+</sup>CGPA 3.63 ● आयएसओ प्रमाणित कॉलेज : 9001:2015

Jr. College No. j.21.02.003

संदर्भ क्र.: 2269/19-20

REGISTERED AD

दिनांक :

30/05/5050

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डॉ. कर्मवीर माऊराव पाटील

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प्रति.

मा.प्राचार्य. कृष्णा महाविद्यालय, रेठरे बु ॥ जि.सातारा

> विषय: अग्रणी महाविद्यालय योजनेअंतर्गत रिसर्च प्रोजेक्टसाठी द्यावयाच्या ॲडव्हान्सबाबत.

महोदय,

शिवाजी विद्यापीठाच्या Research Promotion Activity for students of the affiliated Colleges या योजनेअंतर्गत आपल्या विद्यार्थ्यांच्या "Synthesis Characterization of WO3 thin film by solgel route for electrochromic smart windows application" या रिसर्च प्रपोजलसाठी विद्यापीठाने रू. १००००/- मंजूर केले असून सदर ॲडव्हान्स रकमेचा चेक नं. १०००७१३ दि. १ (10२/२०२० ने सोबत पाठविला आहे. कृपया सदर रकमेची पोहोच पावती त्वरीत पाठवून द्यावी.

वरील रिसर्च प्रोजेक्टसाठी मंजूर रक्कम रू.१०,०००/- खर्च करून त्याचा हिशोव व प्रोजेक्ट रिपोर्ट या अग्रणी महाविद्यालयाकडे त्वरीत साद्धर करावा.

KARA

कळावे, ही विनंती.

आपला विश्वास

अग्रणी महाविद्यालय सद्गुरू गाडगे महाराज कॉलेज, कराड

सोबत : वरीलप्रमाणे.

Kitshiia menavidyalaya. Shivpagar

Inward No.- 1357

25/02/20 Date-



## संहती कार्यसाधिका । शिलं परं भूपणम्

Shetkari Shikshan Prasarak Mandal's

## KRISHNA MAHAVIDYALAYA, RETHARE BK.

Shivnagar, Tal. Karad, Dist. Satara, 415108 (M.S.)

Email: kmr\_sspm@yahoo.co.in

Website: www.krishnamahavidyalaya.com

NAAC 'B+' Grade (CGPA 2.65)

Principal (I/C): Dr. Salunkhe C. B., M.Sc., Ph. D.

Ref. No. : KMR/313/2019-2020

Founder: Hon. Jaywantrao Bhosale

Date: 03-03-2020.



प्रति. मा समन्वयक अग्रणी महाविद्यालय सदगुरू गाडगे महाराज कॉलेज कराड जि सातारा

विषय: - अग्रणी महाविद्यालया अंतर्गत राबवलेल्या रिसर्च प्रोजेक्टच्या खर्चाचा अहवःस.

महोदय.

शिवाजी विद्यापीठाच्या Research Promotion activity for students of the affiliated collages यात्राजनेअंतर्गत यात्राहाविद्यालयातील विद्यार्थ्यांच्या "Synthesis, Characterization of WO3 thin film by sol-gel route for electrochromic smart windows application" यारिसर्च प्रोपोसलं साठी विद्यापीठाने रु. १०,०००/- मलंगू क्षेत्रहे हो सदर रिसर्च प्रोपोसलं साठी आपणकाड्यू रु. १०,०००/- चाएँडव्हान्स रकमधेा मके नां 10७३१३ दि. 18/02/2020 राज्रे मिळाला असस् त्यानुसार वरील रिसर्च प्रोजेक्टसाठी मलंगू रकमेच्या खर्चाचा तपशकी व अहवका आपणकाडके कारण विदेश साहे

खर्चाचा तपशील खालीलप्रमाणे.

अ नं	तपशील	साहित्य	खर्च
٧.	Shri. Samarth Trading Company	Chemicals	7500.00
٦,	Bhadi and Company	Stationary	154.00
3.	Chingale Surgicals	Stationary	1030.00
٧.	Dhiraj Stationers	Stationary	684.00
4.	Anushka Stationers	Stationary	654.00
٤.	Uma Xerox	Xerox	25.00
		Total	10,047=00

समन्वयक

Received pohol 12/3/20

Principal

Cisma Mahavdyaaya, Rathere Bit

Tel - Cared - 4TE(10) (MS)

## FORMAT FOR STATEMENT OF EXPENDITURE

- 1. SUK file No: SGM/1689/2019-2020 dated 01/11/2019
- 2. Title of the Lead college Project: "Synthesis, Characterization of WO<sub>3</sub> thin film by sol-gel route for electrochromic smart windows application".
- 3. Name of the Project Advisor: Dr. Dhanaji S. Dalavi, Physics.
- 4. Number of students participated in the project: 04
- 5. Duration of the Activity: July 2019 to March 2020.

Sanction No. and	Grant	Details of expenditure in	curred item	Amount	
date	sanctioned	wise		, 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1	
SGM/1689/2019-	10,000/-	Shri. Samarth Trading	Chemicals	7500.00	
2020 dated	and the second	Company			
01/11/2019	Baldagian.	Bhadi and Company	Glass	154.00	
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&		Chingale Surgicals	Stationary	1030.00	
		Dhiraj Stationers	Stationary	684.00	
SGM/2289/2019-		Anushka Stationers	Stationary	654.00	
2020 dated	Let for Lin	Uma Xerox	Xerox	25.0	
17/02/2020			 al Expenditure	10,047.0	
Advance cheque No. 107313.	THE STREET		Grant Received	10,000.0	

It is certified that the sanctioned amount of Rs. 10,000/- (Rupees Ten thousand only) out of the total grant of Rs. 10,000/- (Rupees Ten Thousand) has been received for the project work to the Department of Physics, Krishna Mahavidyalaya, Rethare (Bk) by the Shivaji University, Kolhapur. its Letter No. SGM/2289/2019-2020, **cheque No. 107313**. **dated 17/02/2020** has been utilized for the purpose for which it was sanctioned and in accordance with the terms and conditions laid down by the Shivaji University, Kolhapur.

**Project Advisor** 

Dr. Dhanaji S. Dalavi T. Dhanaji S. Dalavi

insor (Physics) thevidyalay: k.) Tot. Karos

#### **UTILIZATION CERTIFICATE**

Certified that the Rs.10,000/- (Ten Thousand Only) has been sanctioned by the Shivaji University, Kolhapur to couduct the lead college Project entitled "Synthesis, Characterization of WO<sub>3</sub> thin film by sol-gel route for electrochromic smart windows application". on 2019-2020 has been incurred by the observing scrupulously all the rules and as per rates prescribed by the Shivaji University, Kolhapur.

The unspent balance of Rs. **Nil** is refunded to the Shivaji University, Kolhapur wide challan dated------ in the bank ------ in A/C No-----receipt No.-----dated-----

The excess expenditure of Rs. Nil over the advance is receivable from Shivaji University, Kolhapur.

The expenditure incurred of Rs. 10, 047 out of which Rs. 10,000/- has been received wide cheque No. 107313. dated 17/02/2020 and remaining Rs.Nil is receivable from Shivaji University, Kolhapur.

The penal interest of Rs. **Nil** is credited to the University, vide receipt No.-----dated----- under the budget head A.4.R.2.

Certified that the original vouchers, bills and stamped receipt for the above mentioned of A/C are retained in this office and will be made available as and when required.

Place: Shivnagar

Date:

Project Advisor

Dr. Dhanaji S. Dalavi

Krim Libraridyalaya, Kerner BKN Tal. Karan

Dr. Dhanaji S. Dalavi, Assistant Professor Department of Physics, Krishna Mahavidyalaya, Rethare (Bk). Date: 27/02/2020

To,
The Principal,
Krishna Mahavidyalaya, Rethare (Bk)
Shivnagar-415108

Subject: Submission of Bills toward purchase of chemicals and Glassware under lead college Activity research project.

Respected Sir,

With reference to above mentioned subject, we have purchased chemicals and glassware through Research Promotion activity for students of the affiliated college initiated by Shivaji University, Kolhapur for the research project entitled "Synthesis, Characterization of WO<sub>3</sub> thin film by sol-gel route for electrochromic smart windows application".

Herewith I kindly request you to issue check of Rs. 7,500/- in favour of **Shri. Samarth Trading Company, Islampur** toward the purchase of chemicals and glassware's.

Thanking You

Yours Faithfully

Project Advisor Research Promotion Activity

Or. Dhanaji : Dalavi
Assistant Profer (Physics)
Krishna Mah (Idyalaya,
Rethare (Bk.), Fal. Karad

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TAX INVOICE



Peth Sangli Road, Opp. Ganesh Servicing Center, Shripadnagar, ISLAMPUR - 415 409 Tal. Walwa, Dist. Sangli. Tel. (02342) 225394, 225773 Mob. 9822256373 E-mail: samarth.trading@yahoo.com samarth4092@gmail.com

Original Buyer's Copy

oice No.:- CO/411/2019-20 oice date:-19/01/2020

me: The Principal, Krishna Mahavidyalaya

verse Charge (Y/N): te: Maharashtra

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::- Satara IN:-

PO No .:-

Bill to Party

Date :-15/01/2020

Code

Transport Mode:-Vehicle number:-Date of Supply:-

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Ship to Party

Authorised signatory

Name:-Address:-

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Absolute Ethanol 99.9 % AR 500ml	China	18%	2	450.00	900.00	405.00	495.00	9	44.55	9	44.55	584.
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Common Seal

Dr. Dhanaji S. Dalavi, Assistant Professor Department of Physics, Krishna Mahavidyalaya, Rethare (Bk). Date: 27/02/2020.

To, The Principal, Krishna Mahavidyalaya, Rethare (Bk) Shivnagar-415108

Subject: Submission of Bills toward purchase of contingency under lead college Activity research project.

Respected Sir,

With reference to above mentioned subject, we have purchased contingent items of Rs.2,547/- through Research Promotion activity for students of the affiliated college initiated by Shivaji University, Kolhapur for the research project entitled "Synthesis, Characterization of  $WO_3$  thin film by sol-gel route for electrochromic smart windows application". For the purchase of said items I have paid Rs. 2,547/-

Herewith I kindly request you to issue check of Rs. 2,547/- in favour of Dr. Dhanaji Suresh Dalavi toward the purchase of contingent items.

Thanking You

Yours Faithfully

Project Advisor Research Promotion Activity Dr. Dhanaji S. Dalavi

Ausiatant Professor (Physics) Krichna Mahavidyalaya, Rethard Bk.), Tal. Karad

## GST INVOICE

## BHADI

# BHADI & COMPANY

• Hardware Merchant •

H.O.: 31/32, Shaniwar Peth, KARAD - 415110 ☎ 222821

M/s. प्राचारी कृष्णा अस्तिक्यालिश			INVOICE No. 18	6
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#### BILL OF SUPPLY Cash Sale

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are by : e.Sense InfoSolutions, Karad (9921946393)

## रिज स्टेशनर्स

स्रोमवाव पेठ, कवाउ, जि. सातावा

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BHIVNAGAR 415108

Authorised sang.

## **UMA STORES & XEROX**

Near Krishna Mahavidyalaya, Julewadi.Tal.- Karad, Dist. - Satara.

Shree Principle Krishnu muhawal

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HEAD

PARTMENT OF PHYSICA

RISHEA MANAVIDYALAYA

SHIVNAGAR 415108

कॅश-मेमो अनुष्का स्टेशनरी ॲन्ड जनरल स्टीअर्स कार्वेनाका, कराड. मो. ९७६७२७१९७२ श्री. कुछा। महाविद्यालय रेठरे दिनांक ०३ / ०२ /२०५० तपशील किमंत रुपये प्लास्थिक डबा 801-40 व्हाइरुनर .20 801-टियू पेपर 25 501-कटर -401-20 क्लीप 351-35 फेळीकॉल 45 451-फाईल 601-10 फेव्ही क्वीक 30 601-सार्कर c.D 8. 641-फोम टेप ALL Page No 1000 20 601-फॉइल पेपर 801-80 Order No. 6541 अभूष्का शालेय स्टेशनस्कूण -ॲन्ड जनस्त स्टोअर्स Amount of bill Rs... 654 = 70 कार्वे नाका, कराड 🗯 :९८८ १५७६८७३ सही/शिक्वा Deduction if any Rs.... Bill passed for . Rs. 454

HEAD
DEPARTMENT OF PHYSICA
KRISHNA MAHAVIDYALAYA
SHIVNAGAR 415108

# UNIVERSITY, KOLHAPUR RESEARCH SENSITIZATION SCHEME FOR COLLEGE STUDENTS

UNDER LEAD COLLEGE, ACTIVITY

## SUBMISSION OF PROJECT REPORT

TITLE OF THE PROJECT: SYNTHESIS,
CHARACTERIZATION OF WO<sub>3</sub> THIN FILM BY SOL-GEL
ROUTE FOR ELECTROCHROMIC SMART WINDOWS
APPLICATION.

BY

MISS. LAD DIVYA DILIP,
MISS. MOHINI HARALE RAJENDRA,
MISS. PATIL ASHLESHA ARUN,
MR. DAMAME ANIKET ANIL

UNDER THE GUIDANCE OF

DR. DHANAJI S. DALAVI

ASSISTANT PROFESSOR,

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# DECLARATION

We hereby declare that, the project report entitled "Synthesis, Characterization of  $WO_3$  thin film by sol-gel route for electrochromic smart windows application" submitted by us has been completed and written by us, has not previously formed and published in any other University in India or any other country or examining body to the best of  $\ensuremath{\text{our}}$ knowledge.

Place: Shivnagar

Date: 29/02/2019

## Name of project students.

1. Lad Divya Dilip.

Patil Ashlesha Arun
 A · A · Patel
 Mohini Harale Rajendra.

4. Damame Aniket Anil.

## CERTIFICATE

This is to certify that thesis entitled "Synthesis, Characterization of WO<sub>3</sub> thin film by sol-gel route for electrochromic smart windows application" which is being submitted herewith is the result of original research work completed by Miss. Lad Divya Dilip, Miss. Patil Arun Ashlesha, Miss. Mohini Harale Rajendra, Mr. Damame Aniket Anil under my supervision and guidance and to the best of my knowledge and belief, the work embodied in this project work has not formed earlier.

Place: Shivnagar

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Project Advisor

Salan.

(Dr. Dhanaji S. Dalavi)

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#### 1. Introduction:

The term chromogenic is referred to as the change in optical properties of the compound when it is subjected to change in their environment. So, depending upon the chosen environment chromogenic devices are divided into thermochromic device, photochromic device, electrochromic device, phase dispersed liquid crystal device, gasochromic device and suspended particle liquid crystal device. The history of electrochromism started in 1704, when Diesbach discovered the Prussian blue, an excellent dye which had also electrochromic properties. This material changes its color from dark blue to transparent when a voltage is applied across it. In 1815 the electrochromism of WO<sub>3</sub> was discovered, by Berzelius [1]. In fact, it was showed that pure WO<sub>3</sub> changed color on reduction when warmed under a flow of dry hydrogen gas. Later in 1824 Wohler effected a similar chemical reduction with sodium metal. Kobosew and Nekrasso in 1830, recorded that WO<sub>3</sub> powders could acquire the color blue by electrochemical reduction in an acidic solution. The first step towards an electrochromic device was taken in 1942 by Talmey, in studies on the coloration associated with electrolytic reduction of artificially produced particulate molybdenum and tungsten oxide layers. In 1953 Kraus made a very clear description of electrochromism in tungsten oxide films. As none of these studies attracted much attention, probably most current investigators attribute the first widely accepted suggestion of an electrochromic device to Deb, in 1969, with the tungsten oxide films, and after this point, there was a visible increase of the interest in electrochromism. In spite of the innovation on Deb's first electrochromic device it wasn't able to keep up with the fast development of liquid crystal devices [1, 2].

In 1971, Blanc and Staebler produced an electrochromic effect superior to most of the previously published. They applied electrodes to the opposing faces of doped, crystalline SrTiO3 (Strontium Titanium Trioxide) and observed an electrochromic color move into the crystal from the two electrodes. In 1972, Beegle developed a display having identical counter and working electrodes as the one from Blanc and Staebler, but made of WO<sub>3</sub> [1, 2]. Nowadays, Deb's paper form 1973 is quoted as the work responsible for the true birth of electrochromic technology. Faughan et al. [3] in 1975 accomplished a significant progress in developing the electrochromic display device. This was followed by an increase in

electrochromic devices developed for display applications. Nevertheless, electrochromism has remained an active area for basic and applied research, with large possibilities for applications in emerging technologies.

The interest was boosted in the mid-1980s with the awareness that electrochromism was of much interest as a mean to achieve energy efficiency in buildings, using smart windows [4]. The smart windows and other electrochromic systems consist of two electrodes and an electrolyte. When applied voltage with appropriate polarity, charge in the cell drives in and out of the electrochromic material and an electrochemical redox reaction causes a corresponding color change. Therefore, electrochromic materials are currently attracting much interest in industry for their commercial applications [5]. In the recent year various attempts has been made to prepare nanostructured electrochromic devices with the help of various sophisticated physical and chemical techniques and tools. Physical techniques require sophisticated instruments which are of high cost which impact on the end product to be delivered in the market. Sol-gel route is the simple and inexpensive technique which is capable to produce highly transparent nanostructured WO<sub>3</sub> thin film.

Therefore, in the present project an emphasis has been given to synthesize WO3 thin films by simple, low cost sol-gel dip coating technique.

### 2. Synthesis of Nanostructured WO<sub>3</sub> thin film:

The precursor solution used for the deposition of WO<sub>3</sub> thin films was prepared by dissolving 7.48 g of tungsten metal powder (99% pure, Sigma Aldrich) in 80 ml of H<sub>2</sub>O<sub>2</sub> (30%) [6]. The reaction mixture that was kept for 48 h with constant stirring yielded a deep yellow-coloured PTA sol. As reaction being exothermic, it was conducted between 0 and 10°C in an ice bath. After completion of reaction the reaction mixture was filtered with whatman filter paper and heated at 55 °C in order to remove excess peroxide. As-prepared sol was used as a starting precursor for the deposition of WO<sub>3</sub> thin films and kept for to form gel. The WO<sub>3</sub> thin film of desired thickness was deposited by using sol-gel dip coating method.

#### 3. Characterization:

The structural properties of the films were studied by X-ray diffraction (XRD) patterns recorded using X-ray diffractometer (Bruker AXS Analytical Instruments Pvt. Ltd., Germany), D2 phaser model with Cu-Ka radiation (k = 1.5418 A°). The scanning rate of 10°/min was applied to record the patterns in the range of 10° -80°. The infrared (IR) spectrum of powder collected from all NiO samples were recorded using Perkin-Elmer IR spectrophotometer (model-100) in the spectral range of 400–4,000 cm<sup>-1</sup>. The pellets were prepared by mixing KBr with WO<sub>3</sub> powder collected by scratching film from glass substrates, in the ratio 300:1 and then pressing the powder between two pieces of polished steel. The surface morphology of the films was examined by scanning electron microscopy (SEM; Model JEOL-JSM-6360, Japan, operated at 20 kV) with a thin layer of gold sputter coated prior to analyses. The optical transmittance spectra of fully colored and fully bleached states were measured over the range of 350-1,100 nm using an UV-vis spectrophotometer (Shimadzu, model: UV-1800, Japan). All the electrochromic measurements were performed in an electrolyte (1 M LiClO<sub>4</sub>+Propylene carbonate) in a conventional three-electrode arrangement comprising platinum wire as the counter electrode and SCE serving as the reference electrode using electrochemical quartz crystal measurements (model-CHI-400A) made by CH Instruments, USA. Colorimetric determinations were done with the help of Shimadzu color analysis software by analyzing the transmittance spectra of color/bleach state to evaluate the L\*a\*b\* and Yxy coordinate values. These obtained values were used as reference data in order to get the observed color in reduced and oxidized state for all samples from online color analysis software with 1931 2° observer and D-65 illuminant proposed by CIE Yxy and L\*a\*b\* coordinate.

#### 4. Results and Discussion:

#### **4.1 X-Ray Diffraction:**

The X-ray diffraction (XRD) pattern of nanostructured WO<sub>3</sub> thin film deposited on ITO/glass substrate is shown in Fig.1. It was observed that, the XRD exhibits a broad hump in the low  $2\theta$  region for nanostructured WO<sub>3</sub> thin film typically of an amorphous in nature. Normally, amorphous WO<sub>3</sub> film is more suitable than crystalline WO<sub>3</sub> film for

electrochromic applications. A crystallized structure is less favorable for ions to diffuse through because of the densely packed atomic structure and due to this lithium ion movement through the film is obstructed by the dense structure leading to a lower response time. X-ray diffraction patterns realized on these film allowed to confirmed that WO<sub>3</sub> film is totally amorphous. Such characteristics, typical of amorphous materials combined with a nanostructured WO<sub>3</sub>, are favorable for a fast-electrochromic response.

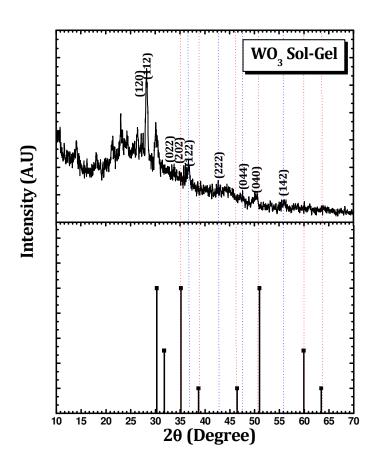


Figure.1 X-ray diffraction pattern of sol-gel deposited WO<sub>3</sub> thin film

### 4.2 FT-IR Analysis:

Fig 2. Shows FT-IR spectra of sol gel deposited nanostructured WO<sub>3</sub> thin film. A single absorption band observed at 950 cm<sup>-1</sup> is characteristic for the terminal W=0 stretching vibration in tungsten trioxide [7].

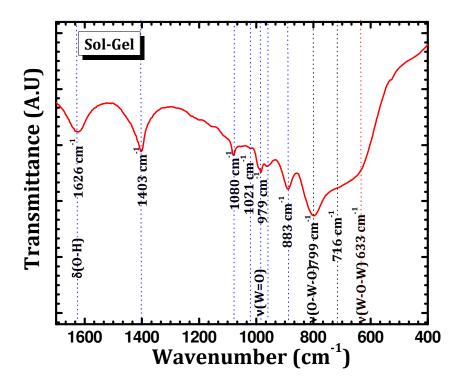


Figure.2 FT-IR spectrum of sol-gel deposited WO<sub>3</sub> thin film.

A well-defined bands seen at 799 and 883 cm<sup>-1</sup>, in the spectra of nanostructured WO<sub>3</sub> thin film is due to the (O–W–O) inter bridging stretching mode in WO<sub>3</sub> [8]. However, the band at 633 cm<sup>-1</sup> is due to W–O–W stretching vibrations. A band centered at 1626 cm<sup>-1</sup> ascribable to the  $\delta$ (H–O–H ) deformation mode is also observed in spectrum of the NPs-WO<sub>3</sub> thin film and a band due to the W–OH...OH<sub>2</sub> stretching mode of hydroxyl groups linked to tungsten on one side and hydrogen bonded with water molecules, on the other, is produced at 1403 cm<sup>-1</sup>[9].

#### 4.3 Morphological Study:

Fig. 3 (a and b) shows low- and high-resolution SEM images of nanostructured WO<sub>3</sub> thin film. Fig. 3 (a and b) revealed agglomerated nanoclusters with average nanoparticles size of 30-40 nm. The film is uniform in nature with high surface are which is beneficial for good electrochromic performance. The thickness of the deposited film is observed to be  $\sim$ 950 nm as depicted in Fig. 3(c). The existence of tungsten and oxygen in the prepared film

was confirmed by the EDS results as shown in Fig 3 (d). It is noted that the high oxygen content in the results was due to the influence of the ITO glass substrate, which was also confirmed by the existence of In and Sn in the EDS analysis.

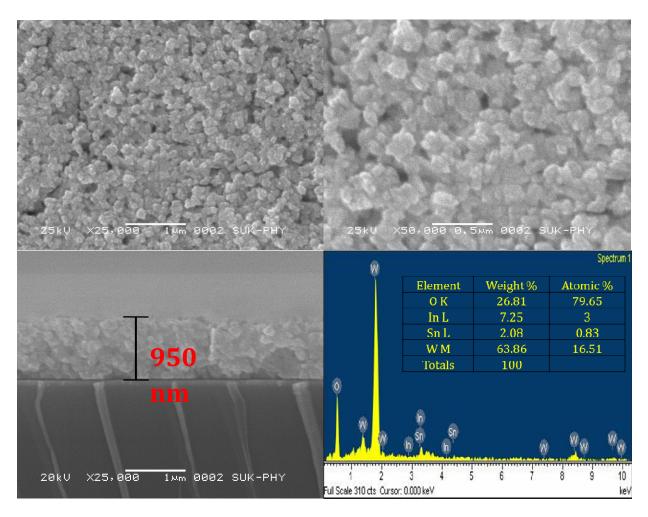


Figure.3 (a and) shows low- and high-resolution images (c) cross sectional image of and (d) EDS spectrum of nanostructured WO<sub>3</sub> thin film deposited on ITO coated conducting glass substrate.

#### 4.4 Electrochromic Study:

Cyclic voltammetry (CV) technique is employed to investigate the cathodic/anodic behavior of  $WO_3$  thin film. The CV was recorded at different scan rates carrying from 20 mV/Sec to 100 mV/sec in 1 M LiClO<sub>4</sub>-PC electrolyte with a potential window of +1.4 to -1.4 V. The shape of the curves is typical of electrochromism in nanostructured  $WO_3$  film. It is observed that the cathodic and anodic current densities for the nanostructured  $WO_3$  film

was  $4.5 \text{ mA/cm}^2$  (cathodic) and  $2.5 \text{ mA/cm}^2$  (anodic) at the scan rate of 20 mV/Sec and achieved a value as high as  $6 \text{ mA/cm}^2$  (cathodic) and  $3.9 \text{ mA/cm}^2$  (anodic) at 100 mV/Sec. The progressive increase in the cathodic and anodic current densities with respect to scan rate indicates the reduction of the  $W^{6+}$  ionic state to the  $W^{5+}$  state due to intercalation of  $Li^+$  ions towards extreme cathodic potentials as a result of  $\{WO_3 + Li^+ + e^- \rightarrow Li_xWO_3\}$  reaction and eventually responsible for blue coloration. Upon anodic polarization (+1.4V), oxidation of  $WO_3$  takes place with simultaneous deintercalation of  $Li^+$  ions and  $e^-$  from the film to acquire a transparent (bleached) state as a result of  $\{Li_xWO_3 \rightarrow WO_3 + Li^+ + e^-\}$  reaction. When the potentials swept from -1.4 V to +1.4 V the reduced W (i.e:  $W^{5+}$ ) gets converted into  $W^{6+}$  state. It is well known that the area under the curve is directly related to the amount of charge intercalated in the film. This confirms that the nanostructured  $WO_3$  thin film shows pronounced electrochromic properties.

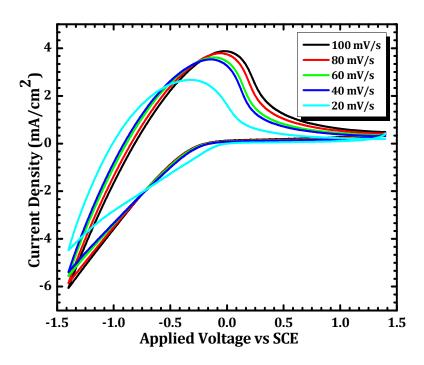


Figure.4 Cyclic voltammograms for the coloration and bleaching cycles of the nanostructured WO<sub>3</sub> thin film recorded in 1M LiClO<sub>4</sub>-PC electrolyte at different scan rates with a potential window from +1.4 to -1.4V versus SCE.

#### 4.5 X-Ray Photoelectron Spectroscopy Analysis:

X-ray photoelectron spectroscopic (XPS) analysis was carried out on the nanostructured WO<sub>3</sub> thin film to investigate the generation of reduced and oxidized tungsten species under the influence of cathodic and anodic potentials. The binding energies of the samples were corrected using a value of 284.6eV for the C 1s peak of carbon. There is no other contaminated element except C in the nanostructured WO<sub>3</sub> film. Fig. 5 (a) shows the XPS spectra of nanostructured WO<sub>3</sub> film under the action of anodic (+1.4 V) potential.

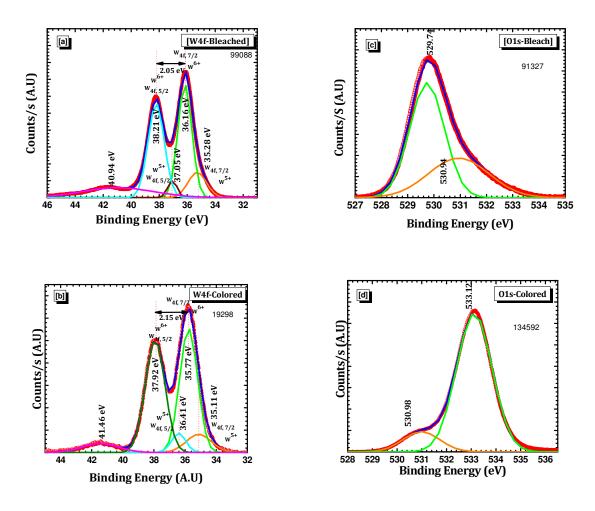


Figure 5. High resolution XPS spectra (a and b) of the W (4f) and (c and d) O (1s) core levels of the WO<sub>3</sub> thin film in bleached and colored state, respectively.

The analysis using XPS revealed a tungsten 4f spectrum in the bleached state composed of the  $W4f_{5/2}$  and  $W4f_{7/2}$  peaks, which may be deconvoluted into Gaussian peaks centered at

38.21 and 36.11 eV, with XPS W4f<sub>7/2</sub>–W4f<sub>5/2</sub> spin-orbit separation being 2.05 eV and the area ratio of the two peaks of each doublet being 0.96 corresponds to the tungsten in W<sup>6+</sup> valence state, which suggests that the film have nominal stoichiometry [10]. On the other hand, the other doublet at 35.28 and 37.05 eV corresponds to a typical W<sup>5+</sup> oxidation state of W. However, during cathodic (-1.4 V) potential (colored state) peaks corresponds to W4f<sub>7/2</sub> and W4f<sub>5/2</sub> shifts toward lower binding energies located at 35.77 eV, 35.11 eV and 37.92 eV, 36.41 eV respectively (Fig.5 (b)). This indicates that the redox reaction takes place between W<sup>6+</sup> and W<sup>5+</sup>. The XPS O1s spectrum in both bleached and colored state has been deconvoluted into two components (Fig.5 (c, d)). The binding energy of the first component situated at 529.74 eV (before coloration) above the W 4f<sub>7/2</sub> core level line, corresponds to the W=O bond in the oxide (Fig.5 (c)). However, there is increase in the intensity and shift in binding energy towards higher energy (533.12 eV) after coloration. The second component in both bleached and colored state observed at about 530.94 eV and 530.98 eV could be assigned to water bounded in the film structure or to water molecules adsorbed on the sample surface [11].

#### 4.6 Reversibility:

Chronocoulometry gives quantitative information about the number of protons/ions intercalated or deintercalated on the application of a potential double step for a known amount of time. The reversibility is then given by the ratio of charges deintercalated to the charges intercalated, i.e.

Reversibility = 
$$\frac{Q_{di}}{Q_i}$$

From the chronocoulometry studies (Fig. 6), the electrochromic reversibility of nanostructured  $WO_3$  film was found to be 75 %.

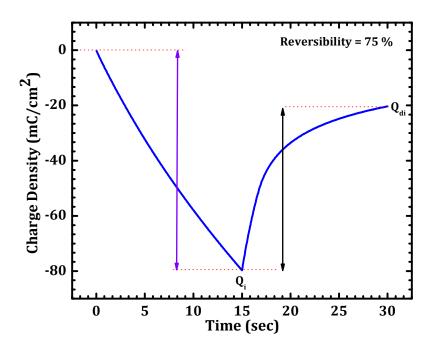


Figure. 6 Chronocoulometry curves of a nanostructured WO<sub>3</sub> thin films recorded in recorded in 1M LiClO<sub>4</sub>-PC electrolyte upon application of step potential of -1 to +1.4 V vs SCE.

#### 4.7 Optical Transmittance Study:

Fig.7 (a) shows the optical transmission spectra of WO<sub>3</sub> thin film at different applied potentials of -0.2, -0.6, -1.0 and -1.4 V respectively, in the wavelength range from 300 to 1100 nm. The optical transmittance of nanostructured WO<sub>3</sub> thin film in the bleached state (+1.4 V) was found to be 67 % at 555 nm and it changes immensely to 3 % as the potential switched to -1.4 V. Therefore, the optical transmittance modulation of nanostructured WO<sub>3</sub> thin film was 64 %. The enhancement in the transmittance modulation of nanostructured WO<sub>3</sub> thin film because of large surface area and increased textural boundaries where actual coloration/bleaching processes take place. The coloration efficiency (g) describes the optical density change ( $\Delta$ OD) at a specific wavelength as a function of the injected/ejected electronic charge (Qi), i.e., the amount of charge required to change the optical density, as shown in Eq. 2 [12].

$$\eta = \left(\frac{\Delta \text{OD}}{Q_i}\right)_{\lambda=550 \text{ nm}} = \left(\frac{\ln(T_b/T_c)}{Q_i}\right),$$
 (2)

where  $T_b$  is the bleached transmittance and  $T_c$  is the colored transmittance. The coloration efficiency of the nanostructured WO<sub>3</sub> thin films has been shown in the Table.1. The high coloration efficiency supports from the fact that nanostructured WO<sub>3</sub> with smaller dimensions prepared by sol-gel provide larger surface area for charge-transfer reactions. It makes the diffusion of ions easier among the materials.

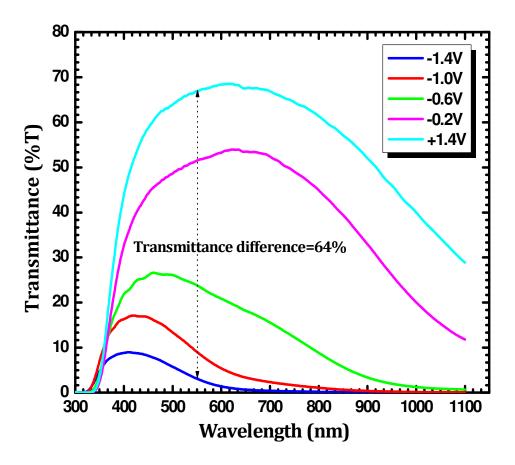


Figure 7. Optical transmission spectra showing colored and bleached states of WO<sub>3</sub> thin film with respect to applied potential.

Table.1 shows various preparative parameters evaluated from electrochromic and optical transmittance studies.

Applied	Photopic		Photopic	Optical	Coloration efficiency
Voltage	Transmittance T (%)		Transmittance	Density	$(cm^2/C)$
	(T <sub>P</sub> )	(T <sub>P</sub> )	difference	( $\Delta$ OD) $_{\lambda=555 \text{ nm}}$	λ=555 nm
	bleached	colored	ΔT <sub>Photopic</sub> (%)		
-0.2	67	51.5	15.5	0.2631	
-0.6	67	24	43	1.02	116
-1.0	67	9	58	2.007	
-1.4	67	3	64	3.10	

#### 4.8 Chromaticity Analysis:

A two-dimensional x-y representation known as the chromaticity diagram utilized to identify the colors of WO<sub>3</sub> thin film in its oxidized and reduced state as shown in Fig.8 (a-b). The shift in x-y co-ordinates occurs once the potential switched from oxidized to reduced state. Fig.4 (a) shows the CIE chromaticity curve of nanostructured WO<sub>3</sub> thin film at different applied potentials. Initially when nanostructured WO<sub>3</sub> thin film is in oxidized state, exhibits a transparent state and its position on the chromaticity curve is close to the white point. As cathodic potential increased from  $\pm 1.4$  to  $\pm 1.4$  V, color of the film immensely changes from transparent to dark blue state as seen by the shift in the position of the x-y coordinate on the chromaticity diagram. In the CIE 1931 Yxy color space, the tristimulus value Y is defined as a measure of the brightness or luminance of the color [13, 14]. Fig.8 (b) shows the relative luminous transmittance (% Y) with applied potential for nanostructured WO<sub>3</sub> thin film. As the potential is switched from  $\pm 1.4$  V to  $\pm 1.4$  V vs SCE, a large change in the xy coordinates occurs as the relative luminance changes from 67 % (bleached) to 3 % (colored) having luminous transmittance difference ( $\pm \Delta Y$ ) of 64 %.

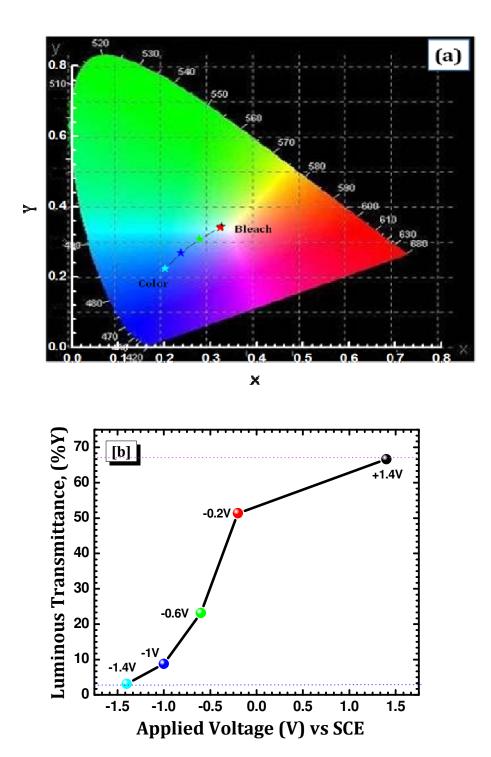


Figure 8. (a) CIE 1931Yxy chromaticity diagram and (a) Luminous transmittance for  $WO_3$  thin film at different applied voltage. The dotted horizontal lines (Fig.8 (b)) indicate difference of luminous transmittance in its colored and bleached state.

#### **Conclusions:**

Nanostructured WO<sub>3</sub> thin film has been deposited by sol-gel dip coating method for energy efficient electrochromic smart window application. The XRD pattern confirms the formation of nanocrystalline WO<sub>3</sub> thin film with amorphous background which is suitable for electrochromic window application. A well-defined bands observed at 799 and 883 cm<sup>-1</sup>, in the FT-IR spectra of nanostructured WO<sub>3</sub> thin film is due to the (O-W-O) inter bridging stretching mode in WO<sub>3</sub> confirms the formation of WO<sub>3</sub>. From electrochromic study it has been concluded that the sol-gel deposited nanostructured WO<sub>3</sub> thin films exhibits transmittance modulation of 64 % at 555 nm, reversibility of 75 % and coloration efficiency of about 116 cm<sup>2</sup>/C. The good transmittance modulation and excellent coloration efficiency of the WO<sub>3</sub> thin films is due to large surface area provided by nanostructured WO<sub>3</sub> thin film. From CIE system of colorimetric analysis and Luminous transmittance modulation it has been evidenced that the color of the WO<sub>3</sub> thin immensely changes from transparent to deep blue which confirms that the film deposited by sol-gel route are well suited for energy efficient electrochromic smart window application.

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