



# Activated Charcoal Supported Cadmium Doped TiO<sub>2</sub> for Photocatalytic and Antibacterial Applications

S.Prabha, V. L. Chandraboss, J.Kamalakaran and S. Senthilvelan\*

Department of Chemistry, Annamalai University, Annamalai Nagar 608002, India.

## ARTICLE INFO

### Article history:

Received: 15 November 2014;

Received in revised form:

19 April 2015;

Accepted: 28 April 2015;

### Keywords

Nanocomposites,  
Photocatalytic,  
Raman Spectroscopy,  
X-Ray diffraction,  
Antibacterial activity.

## ABSTRACT

The photocatalytic activity was studied under UV light using AC-Cd/TiO<sub>2</sub> prepared via precipitation method and characterized by powder X-ray diffraction (XRD), high resolution scanning electron micrographs (HR-SEM) with energy dispersive X-ray analysis (EDX), photoluminescence (PL) and Fourier transform Raman analysis (FT-RAMAN). The enhanced photocatalytic activity of the AC-Cd/TiO<sub>2</sub> is demonstrated through photodegradation of methylene blue under UV light radiation at 365 nm. The mechanism of photocatalytic effect of AC-Cd/TiO<sub>2</sub> Nano composite material has been discussed Further its antibacterial activity against two gram positive and two gram negative bacterial strain is studied.

© 2015 Elixir All rights reserved.

## Introduction

Textile industries produce extensive volume of coloured effluents. Among the different types of dyes used in industries, 65-75% belongs to azo compounds. The release of these coloured dyes in the ecosystem without treatment creates extreme environmental pollution problem by releasing toxic and potential carcinogenic substances into the aqueous phase and damaging to the aesthetic nature of the environment [1]. Many traditional techniques such as precipitation, adsorption, air stripping, reverse osmosis, ultra filtration and coagulation have been utilized so far the waste water treatment but they lost their vitality due to usually non destructive nature towards target molecules and they generate auxiliary pollutants of transferring contaminants from water to another phase[2]

Hence it is important to develop treatment methods that are more viable in destructing dyes from waste water. The removal of azo-dyes by advanced oxidation processes (AOPs) has been the subject of several recent studies. The mechanism of dye destruction in AOPs is based on the formation of a extremely reactive hydroxyl radical (•OH), that, with an oxidation potential of 2.80V can oxidize a broad range of organic compounds [3]. Photocatalysis, as a “green” technique, offers incredible potential in ecological remediation such as the photodegradation of the organic pollutants in the wastewater Semiconductor photocatalysts such as TiO<sub>2</sub> and ZnO nano particles have pulled much attention in recent years due to their various applications to the photocatalytic degradation of organic pollutants in water and air and dye sensitized photovoltaic solar cell.[4]

The interest is much more focused nowadays on the synthesis of photocatalyst materials to beat some limitations of the use of TiO<sub>2</sub>. Some of the important limitations are: TiO<sub>2</sub> powder is hard to reuse, easy to agglomerate, and causes a issue of separation from the solution[5]. Many efforts have been paid to enhance the photocatalytic activity of semiconductor TiO<sub>2</sub>, such as semiconductor combination transition metal doping and noble metal deposition. Among these techniques doping with

transition metals has been proved as an effective strategies to enhance photocatalytic activity[1].

In industrial wastewaters brought about in the textile industry, the heavy metals (cations or complexes), even in traces will adsorb on the photocatalyst surface. Cadmium is one of the presumable heavy metals in these wastewaters. It can be adsorbed on the active sites in processes of chemisorption or it can be integrated in the TiO<sub>2</sub> lattice acting as a doping agent and immobilizing TiO<sub>2</sub> on an inert and suitable supporting materials. For this reason several porous media used as far as possible, to provide a tremendous pore structure for dispersing TiO<sub>2</sub> photocatalyst such as zeolite, alumina, silica, glass, carbon nano tube and activated charcoal.[6-10]

Among these materials, activated charcoal is widely utilized as a support it gas and water remediation due to its good adsorption properties and seems to be an attractive support for TiO<sub>2</sub>. The increment in photocatalytic activity by the addition of charcoal has been well established.

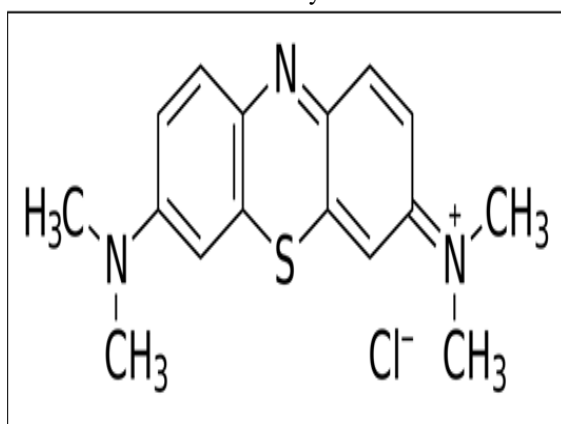
In this work, we report the preparation, characterization, photocatalytic and antibacterial activity of AC-supported Cd doped TiO<sub>2</sub> nano composite materials. The obtained material was characterized by HR-SEM, with EDX, XRD, FT-RAMAN and PL Analysis. In the application part, we have studied their photocatalytic activity towards methylene blue in aqueous solution under UV light irradiation. The reason for choosing methylene blue is due to its wider applications that include colouring paper, temporary hair colourant, dyeing cottons, wools and coating for paper stock and thus acute exposure to MB will cause health problems such as increased heart rate, vomiting, shock, Heinz body formation, cyanosis and tissue neurosis in humans.[11]. AC-Cd/TiO<sub>2</sub> reveals enhanced photocatalytic activity as compared to TiO<sub>2</sub> and Cd/TiO<sub>2</sub> for the degradation of MB under UV light irradiation for 0-60 minutes. The enhanced photocatalytic activity of AC supported Cd/TiO<sub>2</sub> is caused by the reduction of the recombination of electron hole pairs, the extension of the light adsorption range increase of absorption of

light intensity and enhancement of surface active sites. In addition, we report antibacterial activity of prepared AC-Cd/TiO<sub>2</sub> nano composite materials using three gram positive and two gram negative bacteria strains and ciprofloxacin used as a standard. The results shown AC-Cd/TiO<sub>2</sub> have higher bacterial activity than bare TiO<sub>2</sub> and Cd doped TiO<sub>2</sub> nano material.

## Experimental section

### Chemicals

Cadmium acetate dihydrate, titanium tetra isopropyl orthotitanate, activated charcoal, hydrochloric acid, ethanol and methylene blue were the guaranteed reagents of sigma Aldrich and used as such. Deionized water was used for all the reactions and treatment processes. All glassware's were cleaned with chromic acid followed by thoroughly washing with distilled water. Chemical structure of methylene blue is shown in Fig : 1



**Figure 1. Chemical structure of Methylene Blue**

### Synthesis of Ac-Supported Cd- Doped TiO<sub>2</sub> Nano Composite Materials

AC-Supported Cd doped TiO<sub>2</sub> nanocomposite material was synthesized by precipitation method. Initially cadmium acetate dihydrate (0.2 M) and Activated charcoal were dissolved in anhydrous ethanol (solution A). 2 ml of titanium tetraisopropyl ortho titanate (TTIP) is dissolved in ethanol is taken as another solution B. The solution B is added to Solution A and stirred well. To this 2-3 drops of conc. HNO<sub>3</sub> and 4 ml of conductivity water was added at room temperature under vigorous stirring until the precipitate formed. The obtained precipitate was washed with water and ethanol. Then the precipitate was collected and dried in oven at 100°C for 12 hrs. The resulting powder was finally calcinated at 300°C for 2 hrs.

### Photocatalytic Activity

The photocatalytic activity of the prepared photocatalysts was evaluated by the photodegradation of MB by using multilamp photoreactor. The light source was a UV lamp (365nm). The reaction was maintained at ambient temperature (303 K). In a typical experiment, aqueous suspensions of dye (45 mL,  $1 \times 10^{-4}$  mol L<sup>-1</sup>) and 0.06 g of the photocatalyst was loaded in the reaction tube. Prior to irradiation, the suspension was magnetically stirred in the dark to ensure the establishment of an adsorption/desorption equilibrium. The suspension was kept under constant air-equilibrated condition. At the intervals of given irradiation time. The reaction solution was measured spectro photometrically (660 nm) by diluting it five times to keep the absorptions within the Beer–Lambert law limit.

### Antibacterial Activity Study

Antibacterial activities of the synthesized TiO<sub>2</sub>, Cd/TiO<sub>2</sub> and AC-Cd/TiO<sub>2</sub> nanoparticles were determined in comparison with Ciprofloxacin, using the disc diffusion assay method [12-14], is a means of measuring the effect of an antimicrobial agent against bacteria grown in culture. The bacteria in question are

swabbed uniformly across a culture plate. A filter-paper disk, impregnated with the compound to be tested, is then placed on the surface of the agar. The compound diffuses from the filter paper into the agar. The concentration of the compound will be highest next to the disk, and will decrease as distance from the disk increases. If the compound is effective against bacteria at a certain concentration, no colonies will grow where the concentration in the agar is greater than or equal to the effective concentration. This is the zone of inhibition. Thus, the size of the zone of inhibition is a measure of the compound's effectiveness: the larger the clear area around the filter disk, the more effective the compound. Approximately (20 mL) of molten and cooled media (Nutrient agar) was poured in sterilized Petri dishes.

The plates were left overnight at room temperature to check for any contamination to appear. The bacterial test organisms were grown in nutrient broth for 24 h. A (100 mL) nutrient broth culture of each bacterial organism ( $1 \times 10^8$  cfu/mL) was used to prepare bacterial lawns (cfu=number of colony forming unite). Antibiotic disks of 5 mm in diameter were impregnated with TiO<sub>2</sub>, Cd/TiO<sub>2</sub> and AC-Cd/TiO<sub>2</sub> nanoparticles solution (60 µg/mL). The plates were incubated at 37°C and were examined for evidence of zones of inhibition, which appeared as a clear area around the wells. The diameter of such zones of inhibition was measured using a meter ruler and the mean value for each organism was recorded and expressed in millimeter.

### Analytical Methods

High resolution Scanning Electron Microscopy (HRSEM) and Elementary Dispersive X-ray (EDX) analysis experiments were carried out on a FEI Quanta FEG 200 instrument with EDX analyzer facility at 25 °C. XRD spectra was recorded on the X'PERT PRO model X-ray diffractometer from Pan Analytical instruments operated at a voltage of 40 kV and a current of 30 mA with Cu Kα radiation.

FT-RAMAN spectra were recorded with an integral microscope Raman system RFS27 spectrometer equipped with 1024 - 256 pixels liquefied nitrogen-cooled germanium detector. The 1064 nm line of the Nd:YAG laser (red laser) was used to excite. To avoid intensive heating of the sample, the laser power at the sample was not higher than 15 mW.

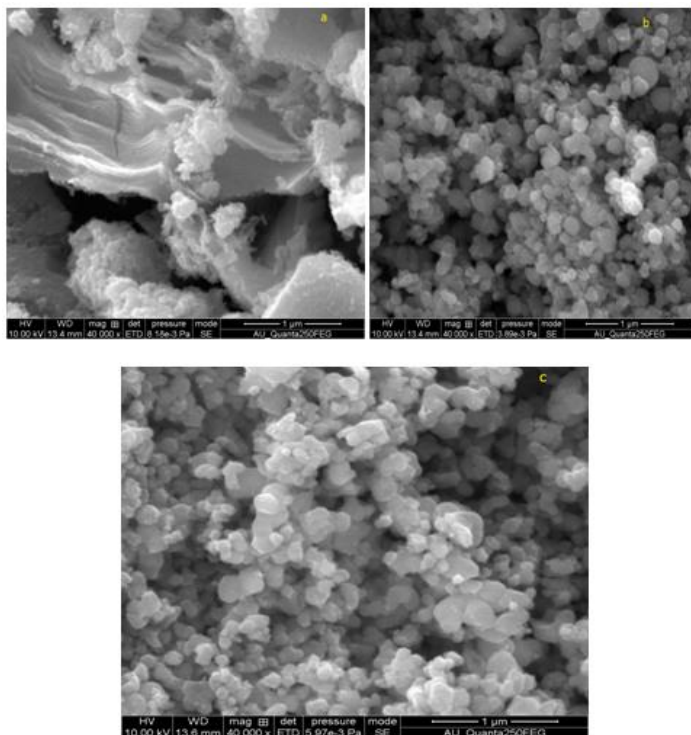
Each spectrum was recorded with an acquisition time of 18 s. UV–vis (ultraviolet and visible light) absorbance spectra were measured over a range of 200–800 nm with a Shimadzu UV-1650PC recording spectrometer using a quartz cell with 10 mm of optical path length. Photoluminescence spectra at room temperature were recorded using a perkin-Elmer LS 55 fluorescence spectrometer. Nano particles were dispersed in chloroform and excited using light of wavelength 300nm.

### Results and discussion

#### Hr - Sem micrograph analysis

Figure (2a), (2b) and (2c) shows the HR-SEM images of the as prepared TiO<sub>2</sub>, Cd/TiO<sub>2</sub> and AC-Cd/TiO<sub>2</sub>. The surface morphology of TiO<sub>2</sub>, Cd/TiO<sub>2</sub> and AC-Cd/TiO<sub>2</sub> has been studied using High Resolution Scanning Electron Microscope. The synthesized TiO<sub>2</sub> consisted of aggregated nanocrystalline anatase.

Due to its very small size of anatase, nanoscale each particle had high driving force to agglomerate, thus the morphology shown by HR-SEM was secondary particles of agglomerated nano sized anatase. The Cd/TiO<sub>2</sub> particles exhibited irregular morphology due to the agglomeration of primary particles and with an average diameter of ~90 nm and AC-Cd/TiO<sub>2</sub> particles exhibited average diameter of ~85 nm.



**Figure 2.** HR-SEM images of (a)  $\text{TiO}_2$  (b)  $\text{Cd/TiO}_2$  (c)  $\text{AC-Cd/TiO}_2$

The observation particle size of the  $\text{Cd/TiO}_2$  was greater than that of  $\text{AC-Cd/TiO}_2$  reveals that absence of AC and also AC matrix controlled the growth of  $\text{Cd/TiO}_2$  particles and prevents their agglomeration. Moreover, the decrease in the particle size can be correlated with the observed increase of the surface area[15]. Figure 3(a), 3(b) and 3(c) shows the EDX images of the as prepared  $\text{TiO}_2$ ,  $\text{Cd/TiO}_2$  and  $\text{AC-Cd/TiO}_2$ .

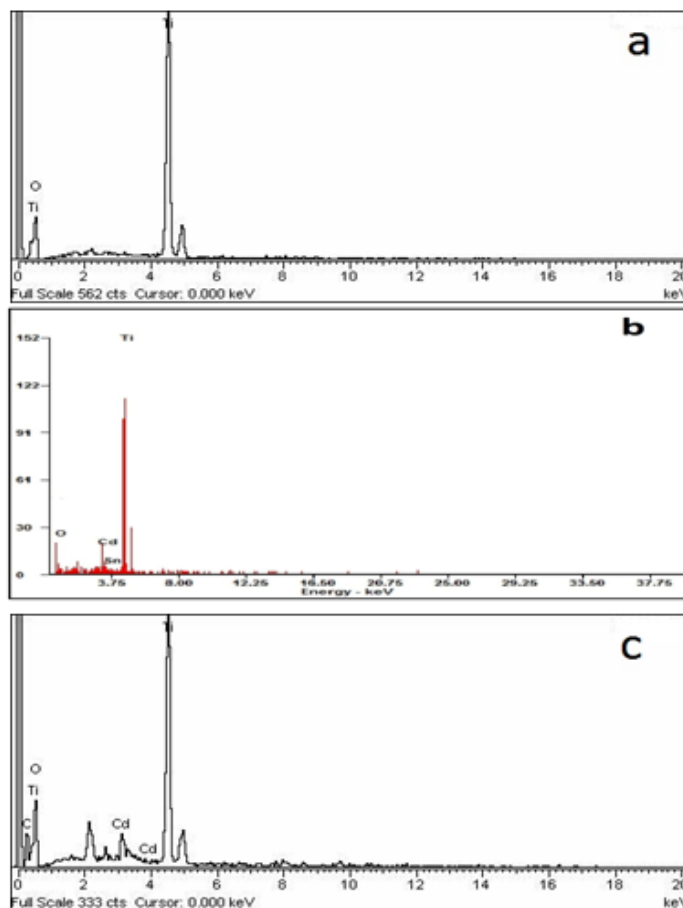
EDX analysis confirms only Titanium, cadmium and oxygen present  $\text{Cd/TiO}_2$  (Figure 3b), where as Titanium, cadmium, oxygen and carbon present in  $\text{AC-Cd/TiO}_2$  composite material (Figure 3c). EDX analysis of the composite sample confirms a doping of the Cd and C in the  $\text{TiO}_2$  material.

#### XRD Patterns

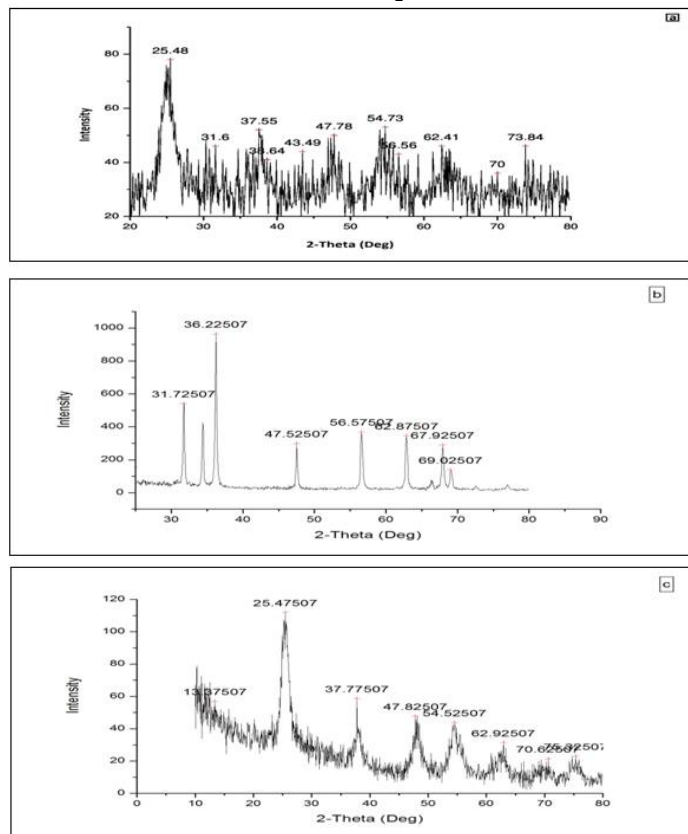
The obtained XRD of the  $\text{TiO}_2$ ,  $\text{Cd/TiO}_2$  and  $\text{AC-Cd/TiO}_2$  nanoparticle are shown in Fig. 4a, 4b and 4c. The  $\text{TiO}_2$  peaks at 25.4, 37.5, 47.8, 54.7 and 62.4, are the diffractions of the  $\text{TiO}_2$  (1 0 1), (0 0 4), (2 0 0), (1 0 5) and (2 0 4) crystal planes. The experimental XRD pattern agrees with the JCPDS card no. 21-1272 (anatase  $\text{TiO}_2$ ) and the XRD pattern of  $\text{TiO}_2$  nanoparticles other literature [16]. The  $2\theta$  at peak  $25.4^\circ$  confirms the  $\text{TiO}_2$  anatase structure [17].  $\text{Cd/TiO}_2$  peaks at 25.4, 34.2, 37.7, 47.8, 54.5, 62.8, and 69.1 are the diffractions of the  $\text{TiO}_2$  (101), Cd (111), (JC PDS File No. 05-0640)  $\text{TiO}_2$  (004),  $\text{TiO}_2$  (200),  $\text{TiO}_2$  (105),  $\text{TiO}_2$  (204), Cd(222).

$\text{AC-Cd/TiO}_2$  peaks at 13, 25.4, 29.3, 33.4, 37.7, 47.8, 54.5, 62.8, and 70.6 are the diffractions of the C (002),  $\text{TiO}_2$  (101), C(002), Cd (111), (JC PDS File No. 05-0640)  $\text{TiO}_2$  (004),  $\text{TiO}_2$  (200),  $\text{TiO}_2$  (105),  $\text{TiO}_2$  (204), Cd(222). Two peaks with a value of  $2\theta$  around 13, 29.3 (JCPDS card no.50 0927) are assigned as graphite oxide and graphite carbon.

The average crystalline size (L) of the  $\text{TiO}_2$  and  $\text{AC-Cd/TiO}_2$  particles can be calculated from the Debye-Scherrer formula [15],  $L = 0.89/\lambda \cos \theta$  where L is the crystalline size (in nm),  $\lambda$  is the wavelength (in nm),  $\theta$  is the Bragg diffraction angle.



**Figure 3a.** EDX Spectrum of  $\text{TiO}_2$  (b)  $\text{Cd/TiO}_2$  (c)  $\text{AC-Cd/TiO}_2$

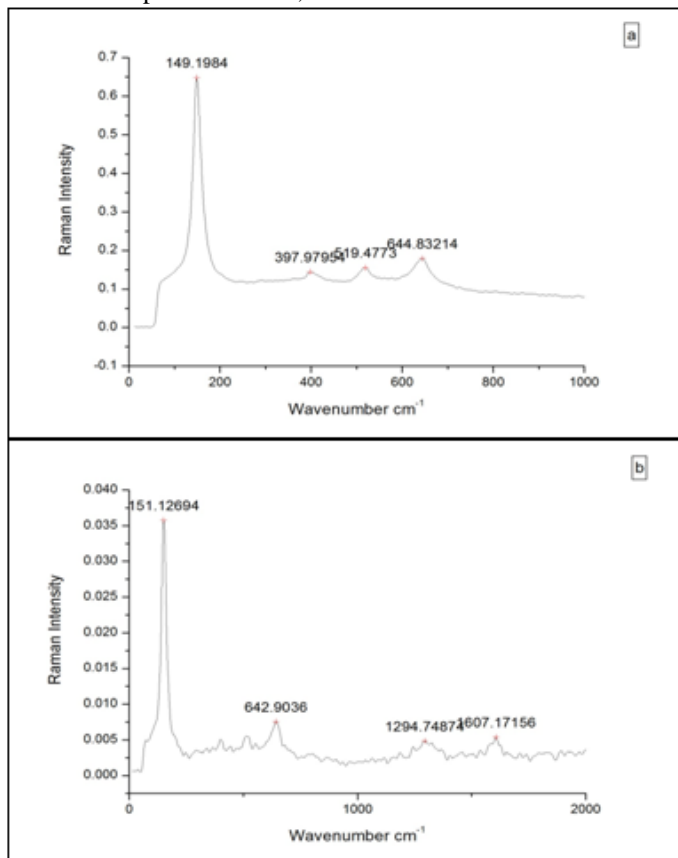


**Figure 4.** XRD patterns of (a)  $\text{TiO}_2$  Nano material, (b)  $\text{Cd/TiO}_2$  Nanocomposite materials and (c)  $\text{AC-Cd/TiO}_2$  Nanocomposite material

The average crystalline size of Cd/TiO<sub>2</sub> product was figured out to be about 88 nm. The average crystalline size of AC-Cd/TiO<sub>2</sub> particles are almost 82 nm. As can be understood, these values are very close to the SEM value.

#### FT-Raman Analysis

Fig. 5a & 5b presents the Raman spectrum of the Cd/TiO<sub>2</sub> and AC-Cd/TiO<sub>2</sub>. The Raman peaks observed at 197.1, 398.4, 518.7 and 644.0 cm<sup>-1</sup> in Fig. 5 are assigned as the Eg, B1g, A1g + B1g, and Eg modes[18] of Cd/TiO<sub>2</sub>. AC-Supported Cd/TiO<sub>2</sub> consists of peaks at 151.1,



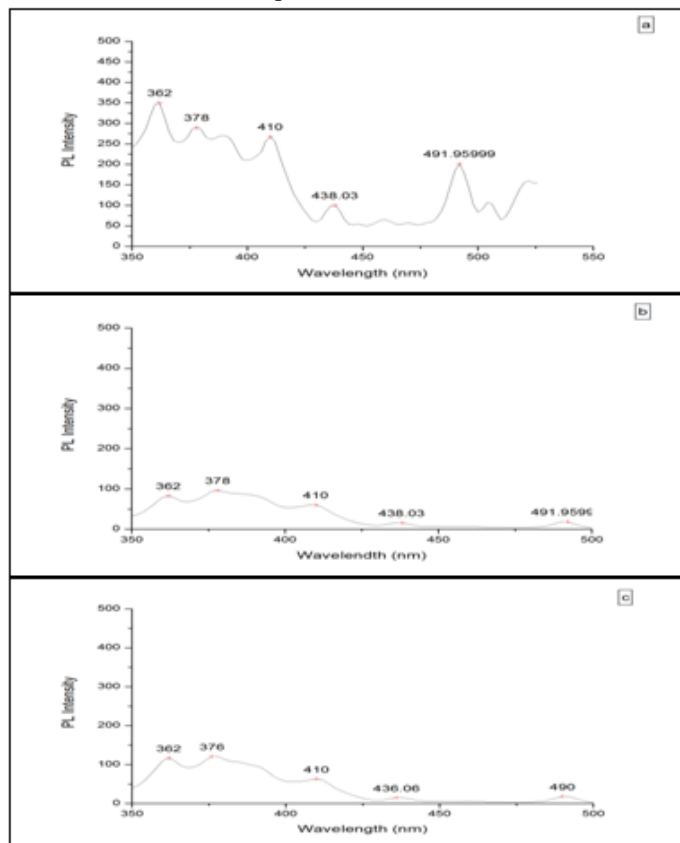
**Figure 5, FT-RAMAN Spectra of (a) Cd/TiO<sub>2</sub> Nanocomposites and (b) AC-CD/TiO<sub>2</sub> Nano composite material**

399.67, 642.28, 511.5 and 1606. The high intensity of the 1325cm<sup>-1</sup> and 1606 cm<sup>-1</sup>, attributed to the D- and G-bands of Carbon, respectively. Which correspond to carbon present in the Cd/TiO<sub>2</sub> material[19]

#### PL Analysis

Photoluminescence spectra of TiO<sub>2</sub>, Cd/TiO<sub>2</sub> and AC-Cd/TiO<sub>2</sub> are shown in figure 6a,6b and 6c respectively. As the photoluminescence occurs due to electron hole recombination, its intensity is directly proportional to the rate of electron-hole recombination. TiO<sub>2</sub> gave emissions at 362, 378, 390, 410, 438, 460, 491 The Photoluminescence spectrum of anatase phase nano TiO<sub>2</sub> resulted from three origins: self trapped excitons [20,21] surface states [22] and oxygen vacancies. The peak situated at 431 nm corresponds to the emission band of self-trapped excitons of the TiO<sub>2</sub> nanoparticles. As it has been reported by Liu et al. [23], a strong emission at 491 nm could be attributed to the indirect recombination via defect states with the interaction of phonons in TiO<sub>2</sub> lattice Oxygen vacancy peaks normally observed at 391nm and 409 nm. the peak with the center at 371 nm could be attributed to the intrinsic emission of the TiO<sub>2</sub>. The PL Spectrum of Cd/TiO<sub>2</sub> nanoComposites show as in the case of undoped TiO<sub>2</sub> and there is no significant shift in the position of the undoped TiO<sub>2</sub> on Cd substitution. AC-

Cd/TiO<sub>2</sub> gave five emissions at 362, 376, 410, 436 and 490 nm. Cd doping TiO<sub>2</sub> and AC Supported Cd/TiO<sub>2</sub> slightly shift the emission of Undoped TiO<sub>2</sub>. The intensity of AC-Cd/TiO<sub>2</sub> . PL emission is less when compared to TiO<sub>2</sub> and Cd/TiO<sub>2</sub> nano composite materials. A weaker PL emission signal is commonly indicative of higher photocatalytic activity[24,25]. The AC-Cd/TiO<sub>2</sub> presented very low emission intensities, indicating that the residual carbon species significantly reduced the recombination rate of the photo-induced electrons and holes[26]



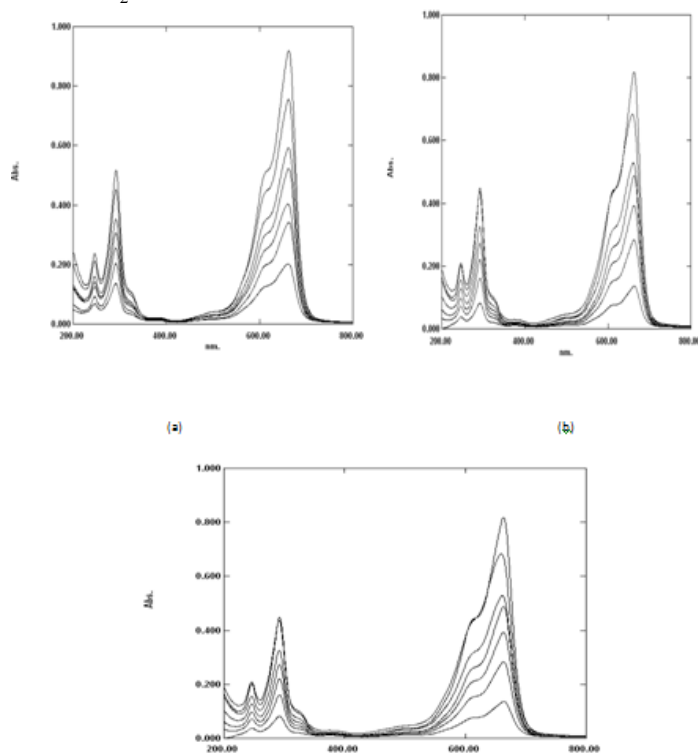
**Figure 6. PL Spectra of (a) TiO<sub>2</sub> Nanomaterial (b) CD/TiO<sub>2</sub> Nano composite material and (c) AC-CD/TiO<sub>2</sub> Nanocomposite material**

#### Photodegradation and Decolourization of Mb Under UV Light Irradiation

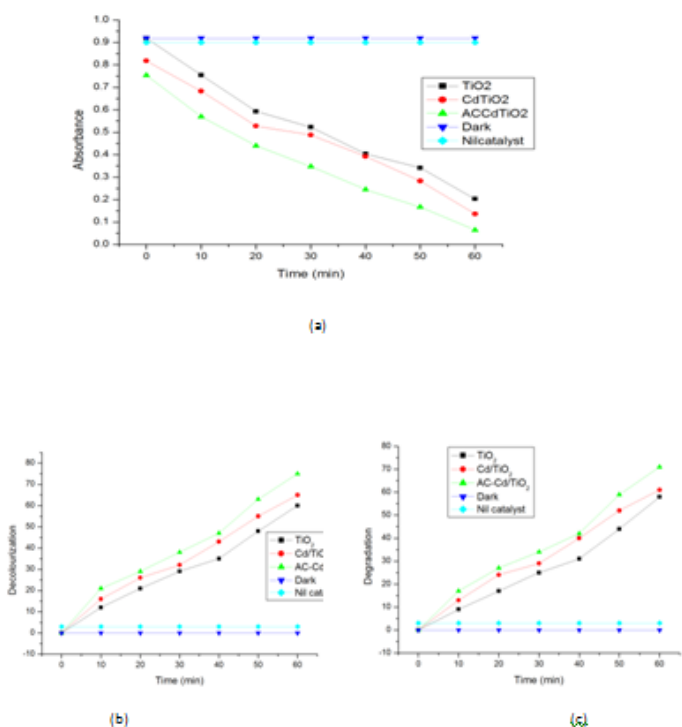
The textile industries daily discharge million litres of untreated effluents in the form of wastewater into public drains that eventually empty into rivers. This alters the pH, increases the biochemical oxygen demand, chemical oxygen demand (COD) and gives the rivers intense colouration. The use of these water resources is limited and the ecosystem is affected. The removal of the polluting dyes is an important problem[27]. So We have chosen photodegradation of MB as model dye with UV irradiation to evaluate the photocatalytic activity of the proposed photocatalyst. Fig. 6 shows the time course of decrease in the concentration of MB under UV irradiation. The photodegradation process of MB, the UV-vis spectra and the colours of MB aqueous solution as a function of under UV light irradiation time in the presence of photocatalysts(TiO<sub>2</sub>, Cd/TiO<sub>2</sub> and AC-Cd/TiO<sub>2</sub> ) are illustrated in Figures 7a , 7b, and 7c. It can be seen from UV-visible spectra changes , while in the case of MB two peaks were initially observed at 663 and 612nm, which merged and the maximum absorbance was finally observed at around 636nm, the strong adsorption peak of MB solution at 636 nm steadily decreased and degraded with increasing the light irradiation time, and the initial blue colour of the solution gradually turned to light – coloured. The comparative study of



photocatalytic activity of  $\text{TiO}_2$ ,  $\text{Cd/TiO}_2$  and  $\text{AC-Cd/TiO}_2$  photocatalysts for photodegradation of MB was shown in figure 7d.  $\text{AC-Cd/TiO}_2$  exhibited excellent photocatalytic activity for MB under UV light irradiation when compared to that of  $\text{TiO}_2$  &  $\text{Cd/TiO}_2$ .



**Figure 7. Absorption spectral changes of Methylene blue solution ( $1 \times 10^{-4}$  M, 45 ml) in the presence of Photocatalyst (a)  $\text{TiO}_2$  (b)  $\text{Cd/TiO}_2$  (c)  $\text{AC-Cd/TiO}_2$  under UV light irradiation at 10 min interval.**



**Figure 8a. Comparative studies of Photocatalytic activity (Absorbance Vs time (min) of (a)  $\text{TiO}_2$  (b)  $\text{Cd/TiO}_2$  (c)  $\text{AC-Cd/TiO}_2$**

Photocatalyst for photodegradation and decolourization of MB under UV light irradiation at 10 minute interval. (b) Comparison of photocatalytic activity (dye decolourization (%))

Vs time (min) of decolourization of MB dye by (a)  $\text{TiO}_2$  (b)  $\text{Cd/TiO}_2$  (c)  $\text{AC-Cd/TiO}_2$  under UV light irradiation at 10 min interval. (c) comparative studies of photocatalytic activity (degradation (%)) Vs time (min) of degradation of MB dye by (a)  $\text{TiO}_2$  (b)  $\text{Cd/TiO}_2$  (c)  $\text{AC-Cd/TiO}_2$  under UV light irradiation at 10 minute interval.

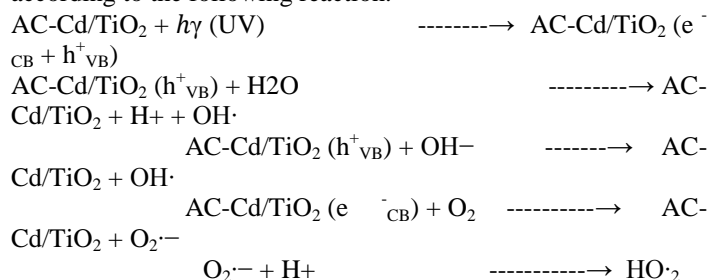
The effect of time on the degradation of MB was examined under UV light irradiation in the reaction time ranging from 0 to 60 minutes. Samples were withdrawn at different time intervals after photodegradation and centrifuged immediately and percentage mineralisation was studied.

Results show that as time increases the percentage decolourization increases. It has been observed that the decolourization of the compound proceeds much more rapidly in the presence of  $\text{AC-Cd/TiO}_2$  doped catalyst in UV light. The reason for better photoactivity could be attributed to the fact that the catalyst is composed of nanocrystalline form. The presence of dopant prevents the recombination of photogenerated electrons and holes leading to better photocatalytic activity.

### Mechanism

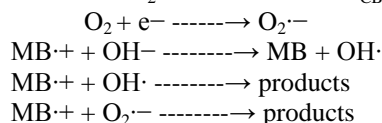
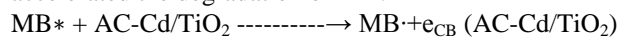
Based on previous reports, the photocatalytic degradation of MB under UV or visible light irradiation is mainly related to the generation of reactive hydroxy and hydroperoxy radicals [28–30]. The hydroxy and hydroperoxy radicals produced by the action of photocatalyst are capable of oxidizing MB molecules at the surface layer of photocatalyst. MB molecules tend to be adsorbed by the surface of the catalyst[31].

In water, UV-irradiated  $\text{Cd-TiO}_2$  is able to completely destroy organic contaminants through the  $\text{TiO}_2$  activation, according to the following reaction.



The electron and the hole are generated in the conduction band and in the valence band of  $\text{TiO}_2$  by UV irradiation, respectively [32]. The positive hole can oxidize hydroxide ions (or water molecule) adsorbed on the surface of  $\text{TiO}_2$  particles to produce hydroxyl radical [33], the electrons of conduction band can react with the oxygen to produce superoxide radical anions [34]. The superoxide radical anion reacts with a proton to form hydroperoxyl radical.

The hydroxyl radical existing on the surface of catalyst accelerated the degradation of MB.



For the mechanism of photosensitized oxidation, in the presence of catalysts the excited state of MB injects an electron into the conduction band. The MB dye is then converted to a cationic dye radical that undergoes degradation to yield products.

### Antibacterial Activity

Antibacterial activity of  $\text{TiO}_2$ ,  $\text{Cd/TiO}_2$  and  $\text{AC-Cd/TiO}_2$  nano composite materials are shown in Table 1. The antimicrobial properties of  $\text{TiO}_2$ ,  $\text{Cd/TiO}_2$  and  $\text{AC-Cd/TiO}_2$  nano composite materials were studied against Gram positive bacteria

namely *Bacillus subtilis*, *Staphylococcus aureus* and Gram negative bacteria *Escherichia coli*, *Pseudomonas aeruginosa* with a size dependent effect.

**Table 1. Antibacterial activity [disc diffusion method]**

S. No.	Bacteria	Standard Antibiotic Disk*	Zone of inhibition (mm)			
			Ti O <sub>2</sub>	Cd/Ti O <sub>2</sub>	AC-Cd/Ti O <sub>2</sub>	Control (DMSO)
1	<i>Bacillus subtilis</i>	22	14	16	17	-
2	<i>Escherichia coli</i>	24	10	13	16	-
3	<i>Pseudomonas aeruginosa</i>	24	09	14	14	-
4	<i>Staphylococcus aureus</i>	22	11	13	16	-

\*ciprofloxacin

17 mm zone of inhibition was developed against *B. subtilis* which was the biggest zone developed against this strain by AC-Cd/TiO<sub>2</sub> nano composite materials but it show 16 mm against the other gram-positive strain *Staphylococcus aureus*. The improved activity was because of its small size compared with TiO<sub>2</sub> and Cd/TiO<sub>2</sub>

AC-Cd/TiO<sub>2</sub> nano composite materials showed activity against both gram-positive strains and also gram-negative strains. 16mm and 14mm inhibition zone was developed against *Escherichia coli* and *Pseudomonas aeruginosa*. AC-Cd/TiO<sub>2</sub> nano composite materials were found good against gram-positive bacteria rather than gram-negative. This was due to the differences in the cell wall composition of these two bacteria.

### Conclusion

AC Supported cadmium doped TiO<sub>2</sub> nanocomposite material was prepared by a precipitation method. This material was characterized by powder X-ray diffraction (XRD), high resolution scanning electron micrographs (HRSEM) with energy dispersive X-ray analysis, photoluminescence (PL) and Fourier transform Raman analysis (FT-RAMAN). These results confirmed the formation of AC-Cd/TiO<sub>2</sub> nanocomposite material. HR SEM and XRD analysis of Cd/TiO<sub>2</sub> showed the average size of as 90nm, AC-Cd/TiO<sub>2</sub> showed the average particle size of as 85nm. moreover the decrease in particle size can be correlated with increase of the surface area. The EDX shows the presence of C and Cd in the TiO<sub>2</sub> material. XRD reveal the all strong peaks can be indexed as anatase form of TiO<sub>2</sub>. PL Spectra explain the suppression of recombination of the photogenerated electron hole pairs by AC-Cd/TiO<sub>2</sub> nano material. AC-Cd/TiO<sub>2</sub> reveals enhanced photocatalytic activities as compared to TiO<sub>2</sub> and Cd/TiO<sub>2</sub> for the photodegradation and decolourization of MB under UV light irradiation for 0 to 60 minutes. The mechanism of photo catalytic effect of AC-Cd/TiO<sub>2</sub> nano composite material has been discussed.

### Reference

- Dongfang Zhang et al., J Mater Sci, 47 (2012) 2155-2161
- Adel A Ismail et al., chemical Engineering Journal., 229 (2013), 225-233
- Nadia Riaz et al., chemical Engineering Journal., 185-186(2012) 108-119
- Senthilvelan S et al., Material Research Bulletin, 48(2013) 3707-3712
- Housda Slimen et al., Journal of photochemistry and photobiology, 221(2011) 13-21
- M V Shankar et al., chemosphere, 63(2006) 1014-1021
- X Zhang et al., Applied catalysis A: General, 282(2005) 285-293

- T Yazawa et al., Ceramics International, 35(2009) 3321-3325
- Y Yu et al., Applied catalysis General, 289(2005), 186-196
- B Thyba., Applied catalysis B: Environmental, 41(2003) 427-433.
- Karakitsou K Eet al., J Phys.Chem, 97(1993) 1184.
1. In: Microbiological assays and tests, Indian Pharmacopoeia, Ministry of Health and Family Welfare, The Controller of Publications, New Delhi, A-100 (1996).
2. H.W. Seely and P. J. Van Demark., A Laboratory Manual of Microbiology, Taraporewala Sons and Co., Mumbai, 55 (1975).
3. A. L. Barry., The antimicrobial susceptibility test. Principle and Practice, Lea and Febiger, Philadelphia, 180 (1976).
- Mohd Athar., Adv. Mater. Rev. 1(1)( 2014) 25-37
- Antic Z, Krsmanovic RM, Nikolic MG, Cincovic MM, Mitric M, Polizzi S, Dramicanin MD., Multisite luminescence of rare earth doped TiO<sub>2</sub> anatase nanoparticles. Mat.Chem.Phys. 135(2012) 1064-1069.
- Ba-Abbad M, Kadhum AH, Mohamad A, Takriff MS, Sopian K, Synthesis and Catalytic Activity of TiO<sub>2</sub> Nanoparticles for Photochemical Oxidation of Concentrated Chlorophenols under Direct Solar Radiation Int.J.Electrochem.Sci. 7(2012) 4871-4888.
- Ohsaka, T. J. Phys. Soc. Jpn. 48(1980) 1661.
- B. W. Mwakikunga., Nanoscale Res Lett 3 (2008) 421-426
- L.V. Saraf, S.I. Patil, S.B. Ogalae, S.R. Sainkar, S.T. Kshirsager, Int. J. Mod. Phys. B12 (1998) 2635.
- H. Tang, H. Berger, P.E. Schmid, F. Levy, Solid State Commun. 87 (1993) 847.
- L. Fross, M. Schubnell, Appl. Phys. B: Photophys. Laser chem. 56 (1993) 363.
- Liu B., Zhao X., Zhao Q., He X., Feng., J Journal of Electron Spectroscopy and Related Phenomena 148(3)( 2005) 158-63.
- Chen, X.F., Wang, X.C., Hou, Y.D., Huang, J.H., Wu, L., Fu, X.Z., J. Catal. 255(1)(2008) 59-67.
- He, Z.L., Zhu, Z.F., Li, J.Q., Zhou, J.Q., Wei, N., J. Hazard. Mater. 190(1-3)(2011) 133-139.
- Yu, J.G., Ma, T.T., Liu, S.W., Phys. Chem. Chem. Phys. 13(8)(2011) 3491-3501.
- Darshan Marjadi., International Journal of Chemtech Applications 2(2)(2003) 126-136.
- Tschirch, J.; Dillert, R.; Bahnemann, D.; Proft, B.; Biedermann, A.; Goer, B. Res. Chem. Intermed. 34(2008) 381-392.
45. Fu, P.-F.; Zhao, Z.; Peng, P.; Dai, X.-G. Chin. J. Process Eng. 8(2008) 65-71.
46. Kim, J.; Choi, W.; Park, H. Res. Chem. Intermed. 36(2010) 127-140.
- Tursun Abdiryim et al., Materials 7(2014) 3786-3801
- H. A. Le, L. T. Linh, S. Chin, and J. Jurng, Powder Technology, 225(2012) 167-175,
- R.-J. Wu, C.-C. Chen, C.-S. Lu, P.-Y. Hsu, and M.-H. Chen, Desalination, 250(3)(2010) 869-875
- G. Wang, F. Wu, X. Zhang, M. Luo, and N. Deng, Journal of Hazardous Materials, vol. 133(1-3)(2006) 85-91.