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# New green synthesized reduced graphene oxide–ZrO<sub>2</sub> composite as high performance photocatalyst under sunlight

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We report a green route method for the effective reduction of graphene oxide using Cinnamon extract. ZrO<sub>2</sub> nanomaterial synthesized *via* facile green induced method and zirconia–reduced graphene oxide nanocomposite (NC) synthesized by reflux method. The structural properties of the prepared NC had been characterized using PXRD, SEM, TEM, Raman, PL and XPS analysis. Further, the excellent photocatalytic properties of NC for the decomposition of reactive blue 4 compared to commercial P25 can be attributed to its excellent electrical properties and well connected arrangement derived from the structural reward of the ZrO<sub>2</sub> and rGO support. This results in significant electron-acceptor/transport of graphene materials that effectively decrease the photo-generated electron–hole recombination thereby enhancing the absorption of light. It opens new window to use this eco-friendly, simple and low cost method to synthesize reduced graphene-based composite materials.

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## 1. Introduction

Recently, the main concerns for the environmentalists are water soluble and colored organic compounds such as dyes. The aquatic system present in the water are severely affected by the dye manufacturing, coloring and textile industrial residues which are highly toxic, carcinogenic to human beings.<sup>1</sup> The major source for water pollution, eutrophication and perturbation of aquatic ecosystem is due to release of dyes into the water. In the recent years, tremendous work involved in the rectification of these environmental issues mainly highlights the photocatalytic process which is the best candidate for the efficient dye remediation under light illumination. The study of photocatalytic technology can't be separated with photocatalytic materials. Environmental friendly ZrO<sub>2</sub> is one among the fundamental transition metal oxide with less toxicity and high chemical inertness.<sup>2</sup> In addition, it is biocompatible, cost-

effective with good thermal stability and electrochemical nature. It serves as an excellent material as an active electrode for the electro catalysis of critical analyses because of its excellent electro catalytic properties.<sup>3</sup>

Currently, majority of the materials used in photocatalysis are semiconductor materials, such as TiO<sub>2</sub>, ZnO and ZrO<sub>2</sub>, *etc.*, as the core body of the catalyst. Out of these, ZrO<sub>2</sub> is the only metallic oxide which has both oxidizability and reducibility due to its acidic and basic nature. Since ZrO<sub>2</sub> is a kind of p-type semiconductor,<sup>4</sup> it easily produces oxygen holes, as the carrier of the catalyst, hence generating interaction with active components. Thus compared to other materials it has more excellent performance as a photocatalyst. Although as a single semiconductor, zirconia has high carrier recombination rate and low quantum efficiency which delimit its practical application to some extent. Due to wide band gap, zirconia can only absorb UV light which constitutes less than 4% of the solar spectrum. Thus, its large scale applications get limited. At present, most of the researches focus on the modification of ZrO<sub>2</sub> by doping, compounding carbon nano tubes and several other techniques to enhance its photocatalytic activity.<sup>5</sup> The last approach is the best way and these composites have been prepared by many researchers to enhance their photocatalytic activity under visible light.

About relevant literature of ZrO<sub>2</sub>, the photodegradation of textile dyes like reactive yellow, rhodamine B, 2,2,4-diphenoxy-acetic under UV light has been reported earlier.<sup>6–9</sup> Also photocatalytic activity for the degradation of ethylene and 4-chlorophenol under UV light by TiO<sub>2</sub>–ZrO<sub>2</sub> binary oxides has been studied.<sup>10–12</sup> The increased activity under UV light in the above cases is ascribed to either stability of catalyst/increase in

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surface area or transfer of charges from  $\text{ZrO}_2$  to  $\text{TiO}_2$  hence preventing the electron–hole recombination. Recently, the photodegradation of phenol under UV light by mixed oxide  $\text{TiO}_2$ – $\text{ZrO}_2$  catalyst has been proved to be very efficient compared to P25. Here primarily fast transfer of photo-generated electrons from the conduction band of  $\text{ZrO}_2$  to  $\text{TiO}_2$  takes place effectively under UV light which is mainly due to its small particle size, high surface area and abundant surface OH groups.<sup>13</sup>

In the past few years, many researchers have been attracted by the improved photocatalytic activity of carbon-based NCs. Hence in order to prepare these composites, graphene, an extremely attractive component has been particularly chosen<sup>14,15</sup> because of its large specific surface area and an extensive two-dimensional  $\pi$ – $\pi$  conjugation structure that has excellent electron conductivity.<sup>16</sup> Carriers in pristine graphene sheets have been already studied to behave as massless Dirac fermions.<sup>17</sup> When combining  $\text{TiO}_2$  nanocrystals with graphene, excited electrons of  $\text{TiO}_2$  could transfer from the conduction band (CB) to graphene through a percolation mechanism.<sup>18</sup> The heterojunction formed at the interface (termed Schottky barrier) separates the photoinduced electron–hole pairs, thereby inhibiting charge recombination.<sup>16</sup> First time, Kamat and co-workers in 2010 have demonstrated the enhanced photocatalytic activity of graphene-based semiconductor–metal composites.<sup>18</sup> After which, Zhang *et al.*,<sup>19</sup> Shen *et al.*,<sup>20</sup> and Zhou *et al.*,<sup>21</sup> performed an one-step hydrothermal methods to prepare graphene– $\text{TiO}_2$  hybrid materials and showed that the composites exhibited enhanced photoactivity towards organic degradation over bare  $\text{TiO}_2$ . Fan *et al.*,<sup>22</sup> fabricated P25–graphene composites by three different preparation methods, *i.e.*, UV-assisted photocatalytic reduction, hydrazine reduction and hydrothermal method which had better photocatalytic performance for  $\text{H}_2$  evolution from methanol aqueous solution as compared to pure P25. Graphene (GR)– $\text{TiO}_2$  mesocrystal composites were synthesized by a facile template-free process based on the sol–gel and solvothermal combination method for the enhancement in visible light photocatalytic performance.<sup>23</sup> Bearing these results in mind, the study on the use of  $\text{ZrO}_2$ –reduced graphene composites on the photodecomposition of Reactive Blue 4 (RB 4) is still in its infancy to the best of our knowledge. This study is intended to examine the role of reduced graphene in the composite towards the photodecomposition of RB 4 under sunlight irradiation.

Our group already demonstrated green assisted solution combustion synthesis.<sup>24–26</sup> Since the plant based synthetic routes are environmental friendly, cost effective, energy saving and less time consumption. Cinnamon (*Cinnamomum zeylanicum*), a common spice used in various countries is well known for its various activities such as antiparasitic, antioxidant, antimicrobial and it was found to reduce blood glucose, blood pressure and serum cholesterol.<sup>27,28</sup> In traditional medicine for treating various ailments in many countries, the bark of Cinnamon has been extensively used. Presence of these health benefit compounds such as cinnamaldehyde, eugenol and their derivatives<sup>29</sup> are mainly responsible for the medicinal properties of Cinnamon and help them to act as a potential

antioxidant. Reduction of GO to rGO is attained by these anti-oxidant which acts as a very good reducing agent. Therefore, an attempt has been made to assess the effectiveness of Cinnamon extract towards the GO reduction.

In the present investigation, GO is reduced to rGO using green induced method. Further, a simple reflux method has been adopted to prepare  $\text{ZrO}_2$ –reduced graphene oxide ( $\text{ZrO}_2$ /rGO) NC. The Cyclic Voltammetry (CV) and Electrochemical Impedance Studies (EIS) were carried out for  $\text{ZrO}_2$  NM and  $\text{ZrO}_2$ /rGO NC to find the capacitance and charge transfer resistance. The results obtained from the electrochemical studies were correlated with photocatalytic and photoluminescence studies. The photoactivity of  $\text{ZrO}_2$ /rGO NC was compared to commercial P25 and studied by evaluating its performance in the photodecomposition of RB 4 dye under sunlight illumination. The  $\text{ZrO}_2$ /rGO NC was shown to exhibit excellent photocatalytic activity as compared to  $\text{ZrO}_2$  NMs and commercial P25.

## 2. Experimental

All the used chemicals were procured from S. D. Fine Chemicals Company and graphite flakes were received from Sigma Aldrich Company. Cinnamon was purchased from local market.

### 2.1 Preparation of Cinnamon extract

In a round bottomed flask, powdered Cinnamon bark and double distilled water were added in the ratio of 1 : 10. It was refluxed for 5 hours with constant stirring at a temperature of 100 °C. The obtained extract was filtered, concentrated and dried.

### 2.2 Preparation of GO

GO was prepared in the laboratory by adding 5 g of graphite and 2.5 g of sodium nitrate in 120 mL of 95% sulfuric acid.<sup>30</sup> This solution was placed in an ice bath and stirred for half an hour followed by the addition of 15 g of potassium permanganate with vigorous stirring at temperature below 20 °C. By adding 150 mL of distilled water slowly, the stirring was continued overnight. The temperature was increased rapidly from 20 to 98 °C and 50 mL of 30%  $\text{H}_2\text{O}_2$  was added. The product obtained was washed with 5% HCl and then with deionized water.<sup>31</sup> Finally, the obtained samples were collected after drying.

### 2.3 Preparation of rGO

The above prepared GO (80 mg) was added to 50 mL of distilled water and sonicated for about 40 min. Cinnamon extract was mixed with the suspension and refluxed for 45 min, hence reducing GO to rGO.<sup>32</sup> The final product was washed with distilled water, dried and stored.

### 2.4 Preparation of $\text{ZrO}_2$ nanomaterial

Solution combustion synthesis was carried out by using zirconyl nitrate ( $\text{ZrO}(\text{NO}_3)_2$ ) as oxidizer and Cinnamon extract as fuel. Stoichiometric ratio of oxidizer and fuel were taken in a cylindrical crucible. This reaction mixture was placed in a muffle furnace preheated at 450 °C. Initially, the mixture boils and



undergoes dehydration followed by liberation of large amount of gases with formation of nanomaterials. The obtained  $\text{ZrO}_2$  nanomaterial (NM) was collected.

## 2.5 Preparation of $\text{ZrO}_2/\text{rGO}$ NC

100 mg of prepared  $\text{ZrO}_2$  was taken in 100 mL of water and added into a round bottom flask along with refluxing for about half an hour. 20 mg of prepared rGO was dispersed in 2 litre of water and stirred for half an hour. rGO solution was added dropwise to  $\text{ZrO}_2$  solution with constant stirring and mixture was stirred for about 24 hours to attain  $\text{ZrO}_2/\text{rGO}$  NC. The schematic diagram for synthesis of  $\text{ZrO}_2/\text{rGO}$  is shown in Fig. 1.

## 2.6 Characterization

SEM (Scanning Electron Microscopy), TEM (Transmission Electron Microscopy) and high resolution TEM were performed to find the morphology of the synthesized samples using SEM, Hitachi-3000 and TEM, TECNAIF-30 respectively. UV-Vis absorption spectra (SL 159 ELICO UV-Visible spectrophotometer) determined the energy band gap ( $E_g$ ). Using  $\text{CuK}\alpha$  (1.541 Å) radiation at a scan rate of  $2^\circ \text{ min}^{-1}$  with nickel filter in the  $2\theta$  range  $20\text{--}70^\circ$ , the crystalline size and phase present in the synthesized samples were analyzed by using room temperature Shimadzu Powder X-ray diffractometer (PXRD). With the help of Horiba Fluorolog Spectrofluorometer, PL emission spectra of the samples were recorded at room temperature. Raman spectrophotometer (Model: Lab RAM HR) had been utilized to analyze

Raman spectra with an exposure time of about 5 s with number of repetitions as three times (spot size: 100 (11); long working distance: 50 (21); power: 3 mW without filter). Using AXIS ULTRA from AXIS 165, XPS (X-ray photoelectron spectroscopy) analysis were performed with Kratos patented magnetic immersion lens, spherical mirror and analyzer charge neutralization system. A total organic carbon (TOC) analyzer (Shimadzu) was used to determine the TOC. Fourier transform infrared spectroscopy (FTIR) studies for the synthesized samples were performed with a Perkin Elmer FTIR spectrometer (Spectrum-1000).

## 2.7 Photocatalytic procedure

Photocatalytic experiments were carried out under UV light irradiation in the laboratory and also under direct sunlight. The experiment was carried out in a photocatalytic reactor with surface area  $176.6 \text{ cm}^2$ . In the photocatalytic reactor, dye solution of known ppm was mixed with required amount of synthesized catalyst ( $\text{ZrO}_2/\text{rGO}$ ). Before starting the experiment, the mixture was stirred for half an hour continuously in order to attain adsorption/desorption equilibrium. Once experiment starts, 5 mL of suspension was withdrawn from photocatalytic reactor at regular intervals. All the experiments were conducted between 11 a.m. to 2 p.m. in the month of May when the solar intensity fluctuations were minimal. The average solar intensity in a day was found to be  $0.753 \text{ kW m}^{-2}$ . In order to avoid the error arising due to changes in the solar intensities, the experiments were carried out simultaneously to ensure similar exposure to sunlight irradiation and their activities were

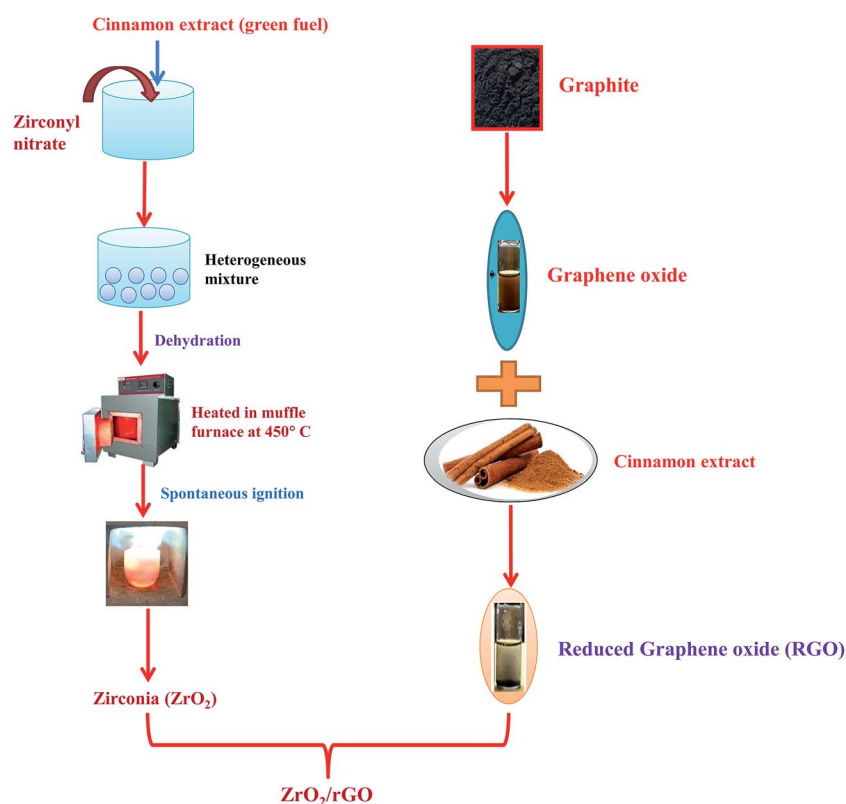


Fig. 1 Schematic diagram for synthesis of  $\text{ZrO}_2/\text{rGO}$  NC.





compared. Finally the suspension was centrifuged and analyzed in UV-Visible spectrophotometer. The amount of dye adsorbed was calculated from the difference in the absorbance before and after the decomposition and tabulated.

### 3. Results and discussions

#### 3.1 Morphological studies

Fig. 2a–c shows the SEM images of  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC. It could be observed that the crystals are irregular in shape/size and slight agglomeration is observed (Fig. 2a). Here, rGO helps  $\text{ZrO}_2$  NM to disperse more smoothly in  $\text{ZrO}_2/\text{rGO}$  NC. Thus, the separation between the agglomerated NPs slightly increases (Fig. 2b and c at different magnification). In these spaces, the rGO sheets exist in the form of flakes. The material was found to be porous with black appearance, which helps in the increased photocatalytic performance. This is analogous with the study carried out by Mao *et al.*, demonstrated black  $\text{TiO}_2$  can more efficiently harvest solar light for photocatalysis.<sup>33</sup>

The microstructure of  $\text{ZrO}_2/\text{rGO}$  NC is studied by TEM (Fig. 3a and b). Abundant  $\text{ZrO}_2$  nanocrystals are densely deposited on the wrinkled graphene sheets due to preferential heterogeneous nucleation and interfacial interactions. As shown in Fig. 3a and b, the wrinkled paper like structure of the 2D graphene sheet can be clearly observed. The HRTEM image (Fig. 3d) of the wrinkled part obtained from the area marked by the red rectangle in Fig. 3a depicts that there were approximately 18 graphene layers.

The graphene plays a vital role in supporting  $\text{ZrO}_2$  nanoparticles dispersion on its surface while  $\text{ZrO}_2$  facilitates to avoid the agglomeration of rGO. The crystallite sizes of the deposited  $\text{ZrO}_2$  nanocrystals are normally 14–18 nm, which is in good agreement with the PXRD results. The intimate interfacial contact between rGO and  $\text{ZrO}_2$  NC can be observed in HRTEM image (Fig. 3c and e) which shows the lattice of both graphene and  $\text{ZrO}_2$ . Since that the transfer process of charge carriers in rGO– $\text{ZrO}_2$  NC is related to the interfacial interaction between graphene and  $\text{ZrO}_2$  NC. This leads to the expectation for improved photocatalysis process due to good charge transfer property. The growth direction of the  $\text{ZrO}_2$  NC is parallel to the lattice fringes and the  $d$  spacing of the (–111) plane is about

0.3368 nm which is close to the lattice spacing of the monoclinic  $\text{ZrO}_2$ . Considering our PXRD study and close observation of the SAED images (Fig. 3f), it can be concluded that the diffraction rings signify the monoclinic  $\text{ZrO}_2$  structure. The SAED pattern of the NC demonstrates the crystalline nature.

#### 3.2 X-ray photoelectron spectroscopy

To investigate the elemental composition of  $\text{ZrO}_2/\text{rGO}$  NC, XPS measurements have been carried out (Fig. 4). The existence of carbon and  $\text{ZrO}_2$  was confirmed by XPS analysis of the  $\text{ZrO}_2/\text{rGO}$  composite as shown in Fig. 4a. The presence of Zr, oxygen and carbon elements in the sample were indicated by Zr 3d, O 1s and C 1s characteristic peaks in the wide scan XPS spectrum of  $\text{ZrO}_2/\text{rGO}$  NC. The existence of graphene sheets were confirmed by the signal for C 1s obtained at the O–C=O (carboxyl groups, 296.4 eV), C=O (carbonyl groups, 288.1 eV), C–OH/C–O–C (hydroxyl and epoxy groups, 285.3 eV) and C=C/C–C (284.4 eV) (see Fig. 4b).

Furthermore, the spin–orbit splitting of the Zr 3d components, Zr 3d<sub>3/2</sub>, Zr 3d<sub>5/2</sub> with a peak separation of 2.4 eV results in the peaks located at 179.8 and 189.3 eV (see Fig. 4c). The peak of the Zr 3d of the nanocomposite showed peaks centered at 179.8 eV is attributed to the formation of a Zr–C bond in the composite.<sup>34</sup> Additionally, the presence of multi-component oxygen species in the surface is understood by the presence of oxygen vacancies of O1s which are asymmetric (Fig. 4d). The curve was deconvoluted into two separate peaks located at 532.29 and 536.01 eV. In the spectrum, there arises a high binding energy component (HBEC) peak located at 536.01 eV and low binding energy component (LBEC) peak located at 532.29 eV due to two different types of oxygen species. The chemisorbed oxygen and lattice oxygen were two different types of oxygen species present in HBEC and LBEC that was originated by surface chemisorbed species such as OH<sup>–</sup> and H<sub>2</sub>O. Thus, from the above explanations it was clear that  $\text{ZrO}_2$  had been incorporated into the rGO layers during synthesis.<sup>35,36</sup>

#### 3.3 Raman studies

Raman spectroscopy is a widely used tool for the characterization of sp<sup>2</sup> and sp<sup>3</sup> hybridized carbon atoms in graphene based materials to differentiate the order and disorder/defects.<sup>37,38</sup> The

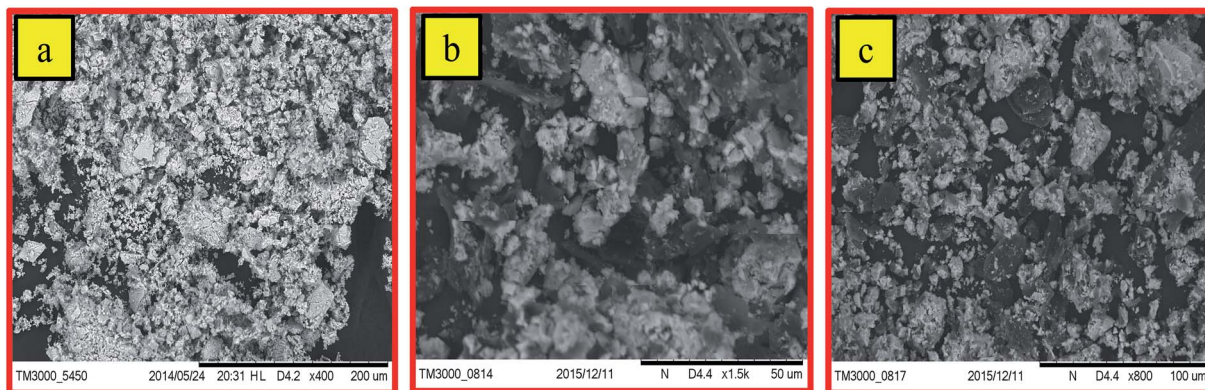


Fig. 2 Scanning electron microscopy of (a)  $\text{ZrO}_2$  NM (b & c)  $\text{ZrO}_2/\text{rGO}$  NC.



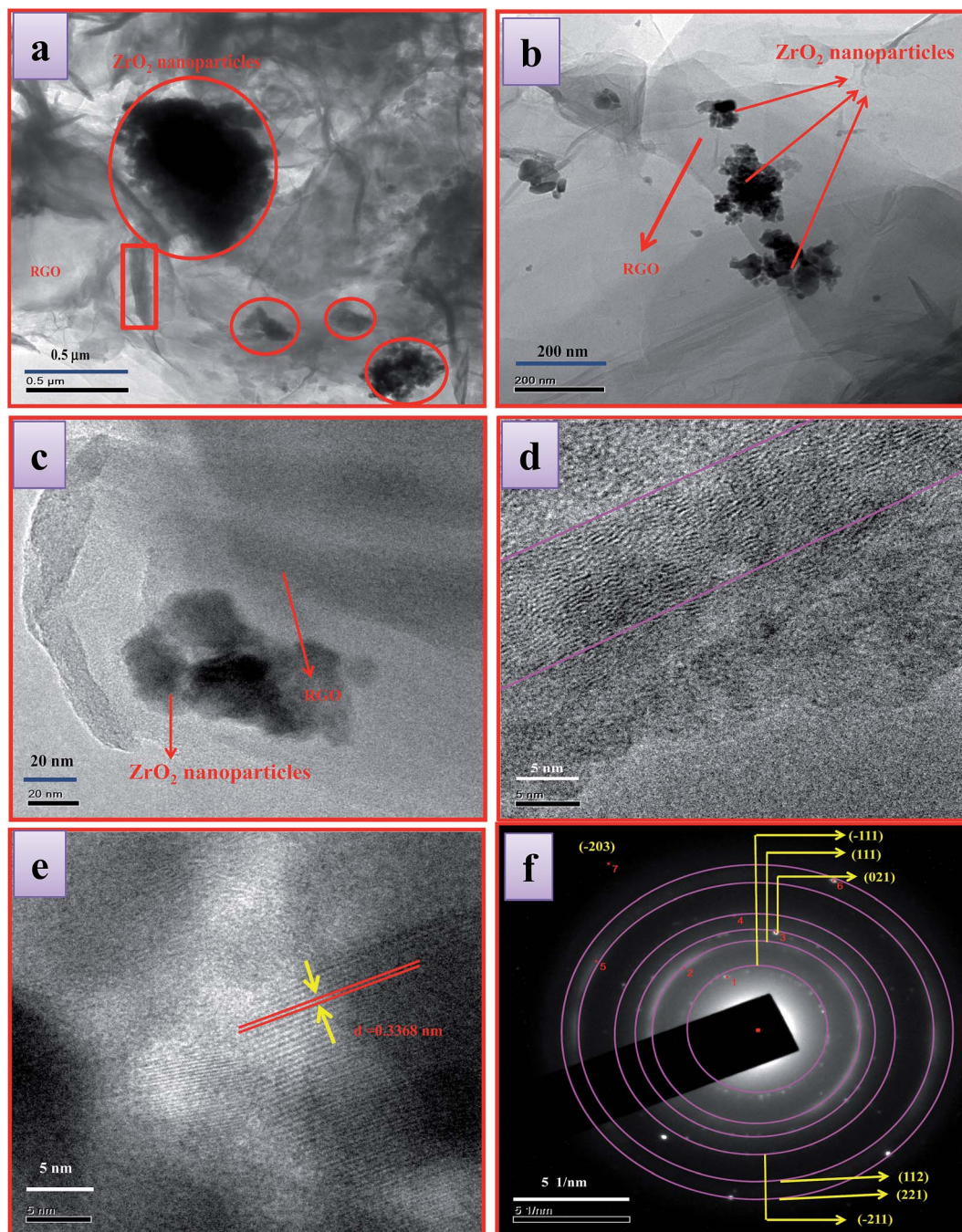


Fig. 3 TEM (a & b), HRTEM (c & d), and SAED pattern (e) of  $\text{ZrO}_2/\text{rGO}$  NC.

Raman signature of GO, rGO and  $\text{ZrO}_2/\text{rGO}$  are depicted in the Fig. 5. The Raman spectrum of GO can be characterized by two main features, D band (the symmetric  $A_{1g}$  mode) at  $1353\text{ cm}^{-1}$  and G band (the  $E_{2g}$  mode of the  $sp^2$  carbon atoms) at  $1581\text{ cm}^{-1}$ .<sup>39</sup> The G band related to the first-order scattering of the  $E_{2g}$  mode noticed for  $sp^2$  carbon domains, while the D band is ascribed to a breathing mode of point phonons of  $A_{1g}$  symmetry, which is a familiar feature of  $sp^3$  defects in carbon and usually can be allied with the structural defects, amorphous carbon, or edges that split the symmetry and selection rule.<sup>38</sup> As a result, the intensity ratio of the D to the G band is usually

a gauge of the disorder/defects in graphene, and a smaller intensity of  $I_D/I_G$  ratio can be consigned to fewer  $sp^3$  defects/disorders and better average size (or less amount) of the in-plane graphitic crystallite  $sp^2$  domains.<sup>38,40</sup> The GO (0.442) shows lesser  $I_D/I_G$  ratio compared to rGO (0.452), signifying that the rGO include more defects.<sup>41</sup> For the  $\text{ZrO}_2$ -rGO composite, the  $I_D/I_G$  ratio is further enhanced to 0.657, signifying the formation of more  $sp^3$  defects in carbon. The probable reason for the increased  $sp^3$  defects can be credited to the strong interaction like Zr–O–C bond between the interface of  $\text{ZrO}_2$  and rGO.





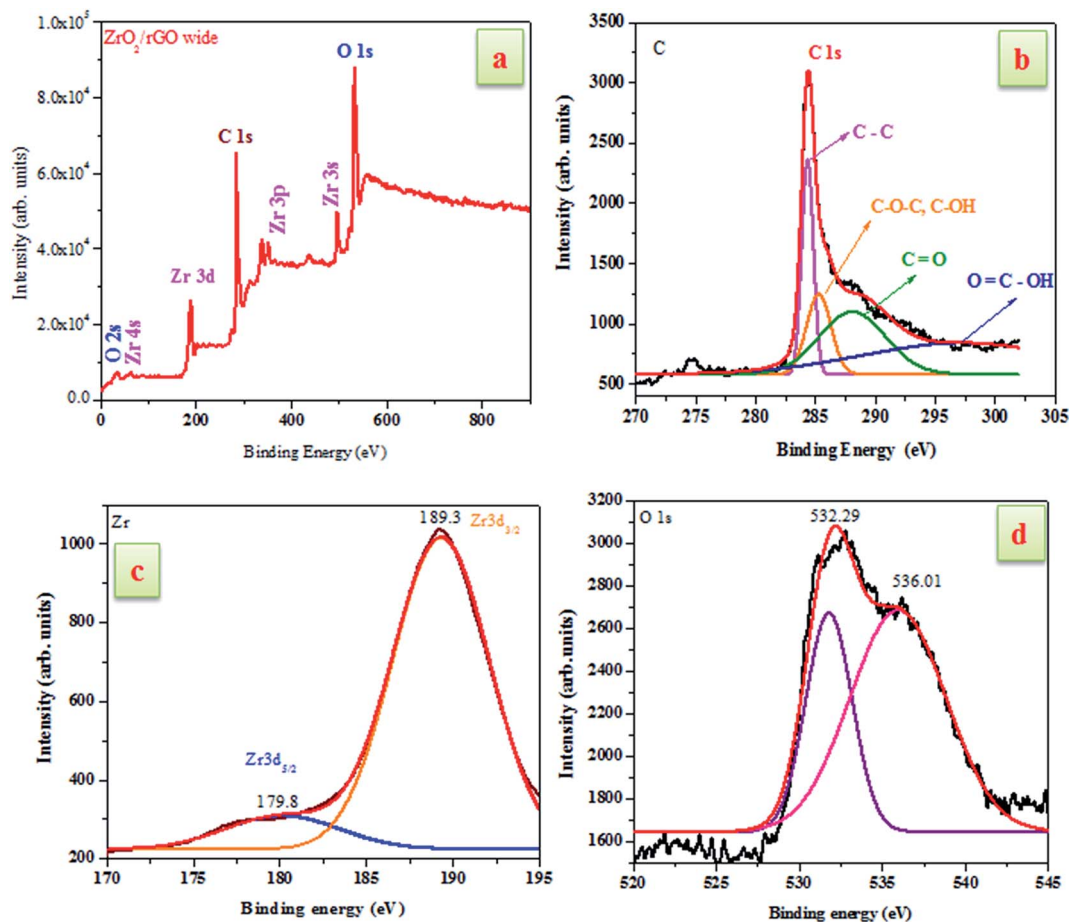


Fig. 4 XPS spectra of  $\text{ZrO}_2/\text{rGO}$  NC (a) survey spectrum (b) C 1s (c) Zr 3d (d) O 1s peaks.

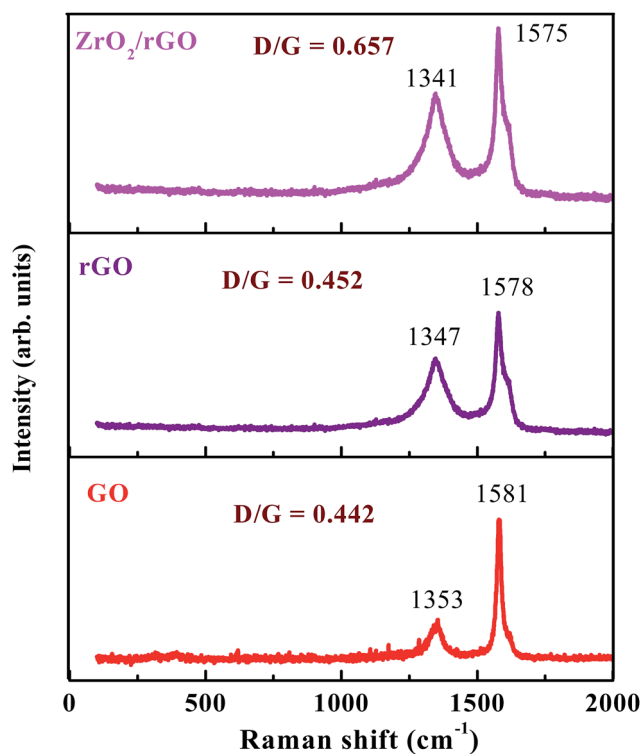


Fig. 5 Raman spectra of GO, rGO and  $\text{ZrO}_2/\text{rGO}$  NC.

### 3.4 PXRD analysis

Fig. 6 shows the PXRD patterns of  $\text{ZrO}_2$ , GO, rGO and  $\text{ZrO}_2/\text{rGO}$  NC. The diffraction peaks of synthesized  $\text{ZrO}_2$  NM show the existence of both monoclinic and tetragonal mixed phase, in good agreement with the standard cards (JCPDS PDF no. 37-1484 and JCPDS PDF no. 49-1642 respectively). But the maximum crystallization takes place along the  $(-111)$  plane of monoclinic phase. The peak at  $\sim 10^\circ$  due to  $(001)$  plane for GO corresponds to the interlayer distance of 0.76 nm (Fig. 6). The fusing of many oxygenated functional groups between the carbon layers increases the interlayer distance. The synthesized GO was subjected to reduction using Cinnamon extract resulting in the shift of the diffraction from  $\sim 10$  to  $\sim 25.6^\circ$ . Thus rGO shows a major  $(002)$  peak at about  $25.6^\circ$ , which is ascribed to the reduction of GO sheets and restacking into an ordered crystalline structure.<sup>42</sup>

While, the PXRD pattern of  $\text{ZrO}_2/\text{rGO}$  NC is similar to diffraction peaks of host matrix except the variation in intensity. The  $(002)$  diffraction peak of graphene was not properly resolved, as the same peak at  $\sim 25^\circ$  overlapped with the  $(110)$  diffraction peak of the monoclinic  $\text{ZrO}_2$ . The broadening of the diffraction peaks observed in  $\text{ZrO}_2/\text{rGO}$  NC indicated the formation of  $\text{ZrO}_2$ -rGO nanocrystallites. The primary size of the



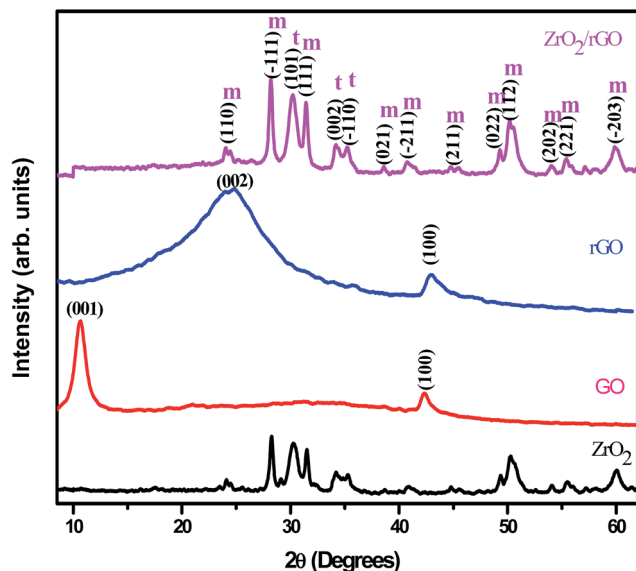


Fig. 6 PXRD patterns of (a)  $\text{ZrO}_2$  NM (b) GO (c) rGO (d)  $\text{ZrO}_2/\text{rGO}$  NC.

$\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC was estimated to be  $\sim 14$  and  $\sim 15$  nm from the Debye Scherrer's equation.

The other structural parameters such as dislocation density ( $\delta$ ), stacking fault were calculated using the following equation and mentioned in Table 1.

$$\delta = \frac{1}{D^2} \quad (1)$$

$$\text{SF} = \left[ \frac{2\pi^2}{45(3 \tan \theta)^{1/2}} \right] \quad (2)$$

### 3.5 FTIR studies

The presence of functional groups in the synthesized samples was analyzed by FTIR. The analysis was carried out by exposing the samples to infrared radiations in the range of 400 to 4000  $\text{cm}^{-1}$ . The FTIR spectra of  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC are represented in Fig. 7. The peaks at 3443  $\text{cm}^{-1}$  correspond to hydroxyl stretching and 1635  $\text{cm}^{-1}$  was attributed to vibration modes from the moisture absorbed on the samples. As reported by several researchers, these two peaks are almost present in all the FTIR spectra (Kacurakova *et al.*, 2000; Nejjatzadeh-Barandozi & Enferadi, 2012). The broad absorption peaks at around 2930  $\text{cm}^{-1}$  are due to the characteristic stretching vibration of  $\text{CH}_2$ . The peak at 1387  $\text{cm}^{-1}$  is due to the presence of OH deformations in the C–OH groups and at 1656  $\text{cm}^{-1}$  is due to the presence of  $\text{sp}^2$  bond of graphite in GO. The band at 1060  $\text{cm}^{-1}$

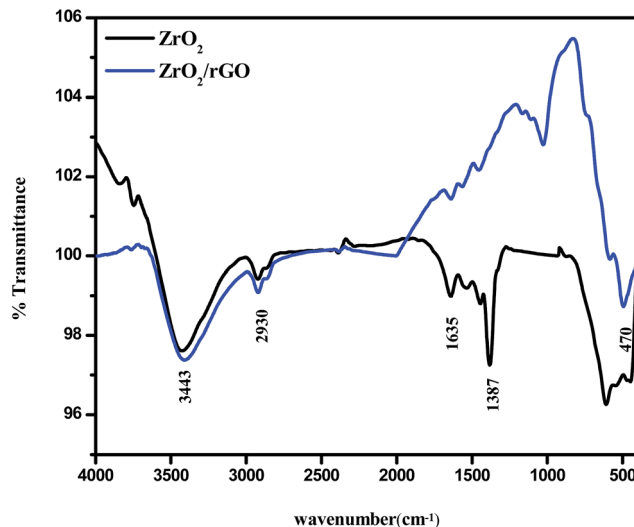


Fig. 7 FTIR spectra of  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC.

is assigned to C–O (epoxy) group which is consistent with XPS studies while the band at 1100  $\text{cm}^{-1}$  is usually contributed to alkoxy group.<sup>43–45</sup>

The absence of peaks at 1060, 1210 and 1656  $\text{cm}^{-1}$  in the  $\text{ZrO}_2/\text{rGO}$  confirms that GO has been reduced to rGO. The peak at 470  $\text{cm}^{-1}$  can be assigned to the Zr–O vibration. These observations may signify that most oxygen functionalities in the GO were removed and  $\text{ZrO}_2/\text{rGO}$  has been formed. The intensity of all moieties containing oxygen present in GO has been reduced suggesting efficient and successful reduction of GO to rGO.

### 3.6 Optical characterization

The difference between the valence band ( $E_{\text{VB}}$ ) and the conduction band ( $E_{\text{CB}}$ ) gives energy band gap ( $E_{\text{g}}$ ), from which the electronic structure of  $\text{ZrO}_2/\text{rGO}$  NC was characterized. In the photocatalytic mechanism, reactive oxygen species has been induced by metal oxide nanoparticles. The production of these reactive oxygen species such as  $\cdot\text{OH}$  and  $\text{O}_2\cdot$  by  $\text{ZrO}_2/\text{rGO}$  were related to the energy band gap and also to redox potential of different reactive species.<sup>46–48</sup> So calculation of energy band gap for the synthesized NC is very important and it is determined by the Wood–Tauc relationship. As shown in Fig. 8, an extrapolation of the linear region of a plot of  $F(R)^2$  on the Y-axis versus energy (eV) on the X-axis gives the value of the  $E_{\text{g}}$ . The  $E_{\text{g}}$  value of  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC was found to be 4.6 and 3.25 eV respectively. This  $E_{\text{g}}$  value for  $\text{ZrO}_2/\text{rGO}$  was very close to the  $E_{\text{g}}$  value reported by Braj Raj Singh and his co-workers.<sup>49</sup> The very

Table 1 Estimated crystallite size, strain, stacking fault and dislocation density of  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC

Samples	FWHM (radians)	Crystallite size (nm) by Scherrer's method	Micro strain ( $\epsilon \times 10^{-3}$ )	SF	$\delta$ ( $10^{15}$ lines per $\text{m}^2$ )
$\text{ZrO}_2$	0.653	14.7	2.68	0.438	4.5
$\text{ZrO}_2/\text{rGO}$	0.627	15.2	2.56	0.460	4.3



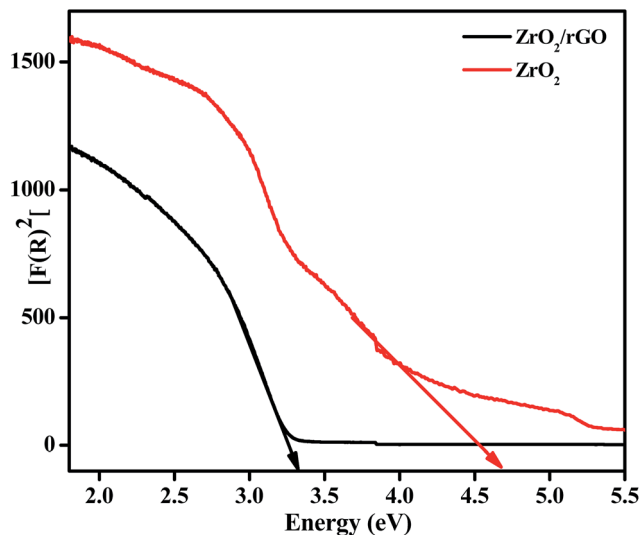


Fig. 8 Plots of  $F(R)^2$  vs. energy (eV) for  $ZrO_2$  NM and  $ZrO_2/rGO$  NC to find energy band gap.

less value of  $ZrO_2/rGO$  NC compared to  $ZrO_2$  NM clearly depicts that the NC has good electron transport and hence can act as a very good photocatalyst with enhanced photocatalytic activity under sunlight irradiation.

The decrease in the  $E_g$  value might be due to high crystallinity or less oxygen vacancies of  $ZrO_2/rGO$  NC as confirmed by PL. The narrowing of band gap could be ascribed to chemical bonding between  $ZrO_2$  and specific sites of carbon during the preparation, which is similar to the case of carbon nanotube– $TiO_2$  composite materials.<sup>50,51</sup> The value of  $E_g$  is decreased to 3.25 eV after anchoring of  $ZrO_2$  onto the enhanced rGO sheet, signifying that the  $ZrO_2/rGO$  can absorb the visible light irradiation efficiently. The narrowing in the band gap can be attributed to the formation of Zr–C and Zr–O–C bond between  $ZrO_2$  and rGO.<sup>52</sup> Compared to  $ZrO_2$  NM,  $ZrO_2/rGO$  NC has favorable energy band gap which agrees with the black

appearance of the NC consistent with SEM analysis. These results suggest that  $ZrO_2/rGO$  NC could exhibit an enhanced photocatalytic activity.

### 3.7 Photoluminescence studies

Fig. 9a shows the excitation spectrum of  $ZrO_2$  NM by monitoring the emission wavelength at 482 nm. The excitation spectrum shows two sharp emission peaks centred at 308 and 428 nm and a less intense peak at 234 nm. These peaks result from electron transfer from valence band to conduction band of the  $ZrO_2$  host. Due to trapping of electrons on the surface states, broad emission peaks arise at 482 nm. This may be also due to the interstitial oxygen lattice defects and oxygen vacancies of the  $ZrO_2$  to valence band of  $ZrO_2$  nanoparticles. The other low intense emission peak centered at 587 nm is owing to the band edge emission of  $ZrO_2$  NM. Compared to  $ZrO_2$  NM, the PL emission peak of  $ZrO_2/rGO$  NC for both broad emission and low intense peak was found to be quenched. Moreover, the broad peak at 482 nm has been slightly shifted towards lower wavelength (467 nm) in  $ZrO_2/rGO$  NC. The decrease in PL intensity of the composite stems from the band alignment at the interface of  $ZrO_2$  and rGO takes place as shown in inset of Fig. 9b.

Fig. 9b depicts PL emission spectra of  $ZrO_2$  NM and  $ZrO_2/rGO$  NC excited at 308 nm. When  $ZrO_2$  NM are attached to the rGO, it is clear from the figure that the electron affinity for  $ZrO_2$  is 2.5 eV (ref. 53) and the Fermi level of the graphene is known to be 4.42 eV.<sup>54</sup> This is due to the direct electron transfer from the conduction band of photo excited  $ZrO_2$  NM to the rGO which is energetically constructive, resulting in the formation of interfaces between the  $ZrO_2$  and rGO. This results in the efficient charge separation of the photo-excited charge carriers leading to inhibition of the recombination of photo-excited electron–hole pairs.<sup>55–58</sup> Hence due to the interaction of  $ZrO_2$  surface with rGO sheets, PL quenching appears with decrease in intensity.<sup>57</sup> Further, there is considerable PL quenching observed in  $ZrO_2$ –rGO compared to  $ZrO_2$  NM, suggesting additional pathways for the disappearance of charge carriers. Obviously,  $ZrO_2$  NM shows

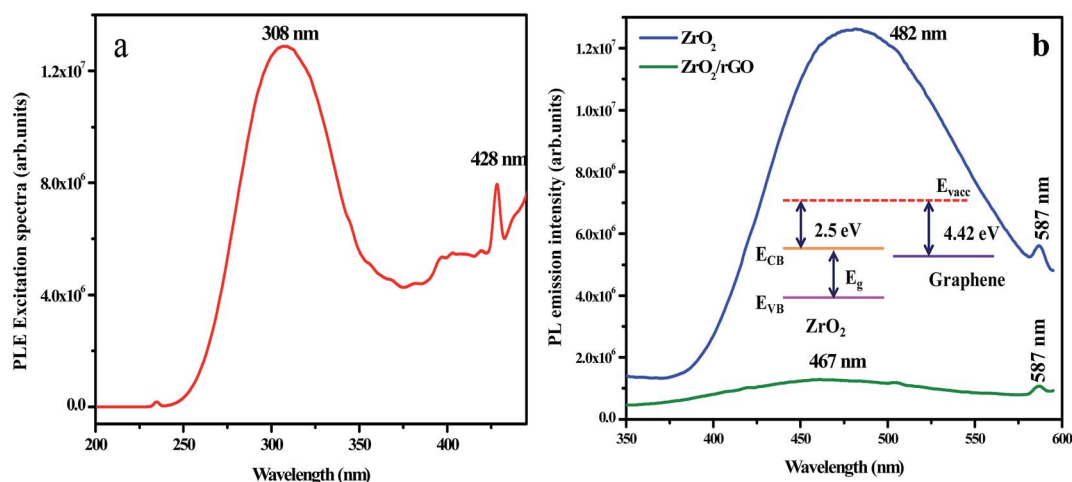


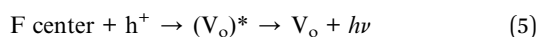
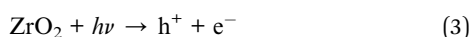
Fig. 9 PL excitation spectrum of  $ZrO_2$  NM excited at 308 nm (b) PL emission spectrum of  $ZrO_2$  NM and  $ZrO_2/rGO$  NC (inset: decrease in PL intensity of the composite stems from the band alignment at the interface of  $ZrO_2$  and rGO).





higher PL intensity than ZrO<sub>2</sub>/rGO NC, indicating ZrO<sub>2</sub> has the highest optical combination rate that deteriorated the photodegradation.

PL spectra mainly depend on the size of the particle, defects and impurities found in the materials. The PL properties of ZrO<sub>2</sub> nanomaterial have been widely investigated by many researchers<sup>59–61</sup> however, the role of particle size and the crystal phases of ZrO<sub>2</sub> nanomaterial has not yet been analyzed in a systematical manner. Harrison *et al.*<sup>62</sup> proved that the luminescent centers constituted by Zr<sup>4+</sup> ions in an asymmetric site bounded by the oxygen ions contribute to the emission peak. The presence of defects and impurities in the systems could be the major reason for the luminescent properties of ZrO<sub>2</sub> nanomaterials. The broad emission band arises due to the singly ionized oxygen vacancies defects in ZrO<sub>2</sub> nanomaterials. The radiative recombination of a photo generated hole with an electron occupying the oxygen vacancy results in an emission band. The electron–hole pairs were obtained when the excitation energy exceeds to the band gap of ZrO<sub>2</sub> nanomaterials. The electrons were trapped rapidly by the oxygen vacancies (VO) thereby producing F centers. The Zr<sup>4+</sup> ions adjacent to the bulk oxygen vacancies had retained the electrons leading to the formation of Zr<sup>3+</sup> ions.<sup>63,64</sup> The holes recombine with the F centers forming the excited states of the emitter. These excited emitters undergo radiative transitions to the ground state. The recombination phenomenon could be summarized by the following reactions:<sup>65</sup>



The mechanism possibly involved in the emission phenomenon of the sample ZrO<sub>2</sub> has been demonstrated with the help of schematic diagram as shown in Fig. 10. The major source of defects centers in ZrO<sub>2</sub> sample was oxygen vacancies/interstitial, Zr vacancies/interstitials (Zr<sup>3+</sup> interstitials).

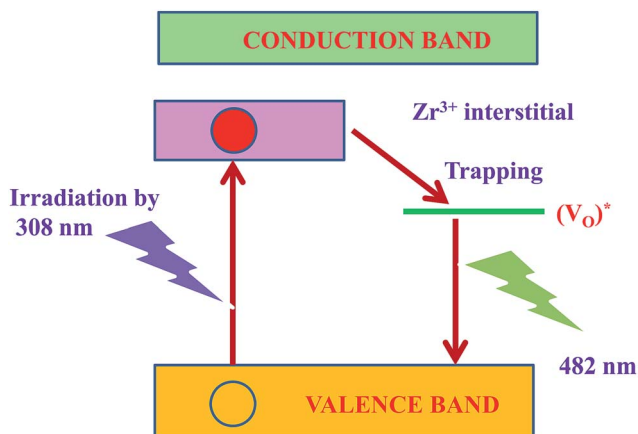


Fig. 10 Energy level diagram for emission mechanism in ZrO<sub>2</sub> nanophosphor.

Moreover, the broad peak at 482 nm has been slightly shifted towards lower wavelength (467 nm) in ZrO<sub>2</sub>/rGO NC. The decrease in PL intensity of the composite stems from the band alignment at the interface of ZrO<sub>2</sub> and rGO takes place as shown in inset Fig. 9b. This blue shifting of the emission band could have originated from the modification of stress/strain in the samples (Table 1) is mainly because of lattice distortions. The decrease in PL intensity is due to the decrease in density of oxygen vacancies. The increase in the particle size could be the possible reason for the decrease in the density of oxygen vacancies.

### 3.8 Electrochemical properties

The Cyclic Voltammetry (CV) data were detected within controlled potential region (−1.2 to 1.2 V) to set a collection of the reproducible current–potential dependencies in 0.1 M sodium nitrate (NaNO<sub>3</sub>) solution using three electrode cell arrangement. Fig. 11a shows the capacitance variations of the ZrO<sub>2</sub>/rGO electrodes at various scanning speed from 10 to 50 mV s<sup>−1</sup>. The capacitance increases with increase in the scanning speeds and predominantly became more obvious at the high scanning speed. For comparison purpose, CV plot of ZrO<sub>2</sub> NM and ZrO<sub>2</sub>/rGO NC were run at 30 mV s<sup>−1</sup> and depicted in Fig. 11b. It is seen from the figure that ZrO<sub>2</sub> NM found to have a very broad semi-rectangular plot while ZrO<sub>2</sub>/rGO NC has less broad semi rectangular shape. This indicated that the capacitance value for ZrO<sub>2</sub> NM is more compared to ZrO<sub>2</sub>/rGO.

Electrochemical impedance spectroscopy (EIS) were recorded by the application of an AC voltage with 10 mV amplitude in the frequency range of 10<sup>5</sup> to 10<sup>−2</sup> Hz in 0.1 M sodium nitrate (NaNO<sub>3</sub>) using three electrode cell that was connected to an electrochemical station, where the prepared samples served as the working electrode, a platinum foil as the counter electrode and the silver as the reference electrode. As an extensively used electrochemical method, EIS is very effective to demonstrate the properties of the electron-transfer phenomenon across the ZrO<sub>2</sub>–electrolyte interfaces.<sup>66,67</sup> Fig. 11c illustrates the Nyquist plots of the electrodes with X-axis as real part Z' of impedance and Y-axis as the inverse of imaginary part −Z''. This semi-circular curve in high frequency region and inclined straight lines in low frequency region represents a typical AC impedance curve for super capacitor.

The diameter of the semicircle in high frequency region represents the charge transfer resistance (R<sub>ct</sub>) value in terms of ohms, which generally associated with the pore structure of electrode materials and faraday pseudo capacitive reaction.<sup>68</sup> The less impedance arc radius in Nyquist plots indicates the faster electron transfer rate.<sup>69</sup> The diameter for ZrO<sub>2</sub> NM and ZrO<sub>2</sub>/rGO NC were found to be 14 and 8 Ω respectively. The diameter for ZrO<sub>2</sub>/rGO was found to be less indicating less charge transfer resistance and hence enhanced photocatalytic activity and decrease in photoluminescence intensity.

Inset Fig. 11c represents the circuit used for fitting the experimental data for impedance plot. Electrolyte resistance for 0.1 M NaNO<sub>3</sub> solution is depicted as element R<sub>s</sub>. Element R<sub>1</sub> corresponds to electrolyte resistance present in porosity of the



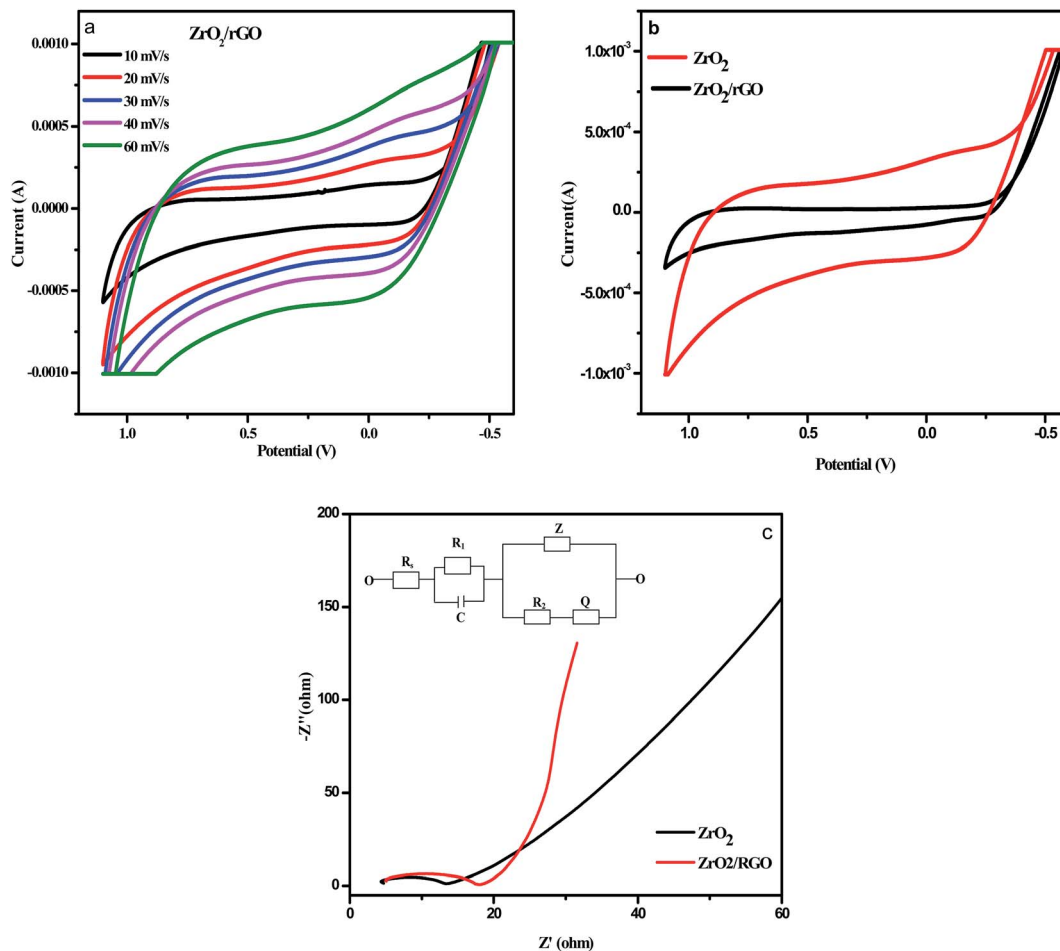


Fig. 11 (a) Comparison of results of cyclic voltammetry (CV) test on  $\text{ZrO}_2/\text{rGO}$  NC at different scan rates tested in 0.1 M  $\text{NaNO}_3$  (b) comparison of results of cyclic voltammetry (CV) test on  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC at different scan rates tested in 0.1 M  $\text{NaNO}_3$  with scan rate of  $30 \text{ mV s}^{-1}$  (c) impedance spectroscopy plot for  $\text{ZrO}_2$  NMs and  $\text{ZrO}_2/\text{rGO}$  NC (inset: circuit fitting program for  $\text{ZrO}_2/\text{rGO}$  NC).

material, while element  $R_2$  corresponds to charge transfer resistance present in phase interface and is inversely proportional to surface area. The element  $C$  is the double layer capacitance that formed when ions from the solution move towards the electrode surface.  $Z$  is Warburg impedance that characterized by having identical real and imaginary contributions, resulting in a phase angle of  $45^\circ$ , which generates a line in the impedance plot. This double layer capacitance ( $C$ ), Warburg impedance ( $Z$ ) and  $Q$  constitutes the Constant Phase Element (CPE).  $Q$  is included in the constant phase element to avoid the lack of homogeneity in the electrode.

### 3.9 Photocatalytic experiment

For evaluating the optical properties of photocatalyst, the performance of UV-Vis light absorption is an important factor.<sup>70</sup> UV-Vis absorption spectra of the  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC are shown in Fig. 12a. In the present work, photocatalytic activity is studied for  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC for degradation of Reactive Blue 4 (RB 4) dye under UV and sunlight irradiation for 120 min. The enhanced photocatalytic activity was observed in  $\text{ZrO}_2/\text{rGO}$  NC than  $\text{ZrO}_2$  NM due to high absorption of sunlight

and because of quenching in recombination of electron-hole pair in metal oxides with the existence of reduced graphene oxide. After achieving the complete adsorption of dye molecules on the composite material, the photocatalytic experiment was conducted.

It was noted that there was very low decomposition of dye under dark condition. But under UV and sunlight irradiation, in the absence of the photocatalyst, RB 4 was very stable. After 120 min of UV and sunlight irradiation, the photolysis efficiency of the dye was negligible. The degradation efficiencies of RB 4 by  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC under sunlight irradiation have been measured to be of 12% and 93% respectively as depicted in Fig. 12b. The photocatalytic decomposition of RB 4 under UV light irradiation for  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC has been carried out for comparison purpose and found to be 65% and 89% (Fig. 12c). Commercial P25 was tested for the photocatalytic activity under the same experimental condition and the degradation percentage was found to be 80% for sunlight and 65% for UV light irradiation. However, it is seen that  $\text{ZrO}_2/\text{rGO}$  NC has got higher photocatalytic activity compared to commercial P25. It was evident that the absorbent capacity of  $\text{ZrO}_2/\text{rGO}$  NC



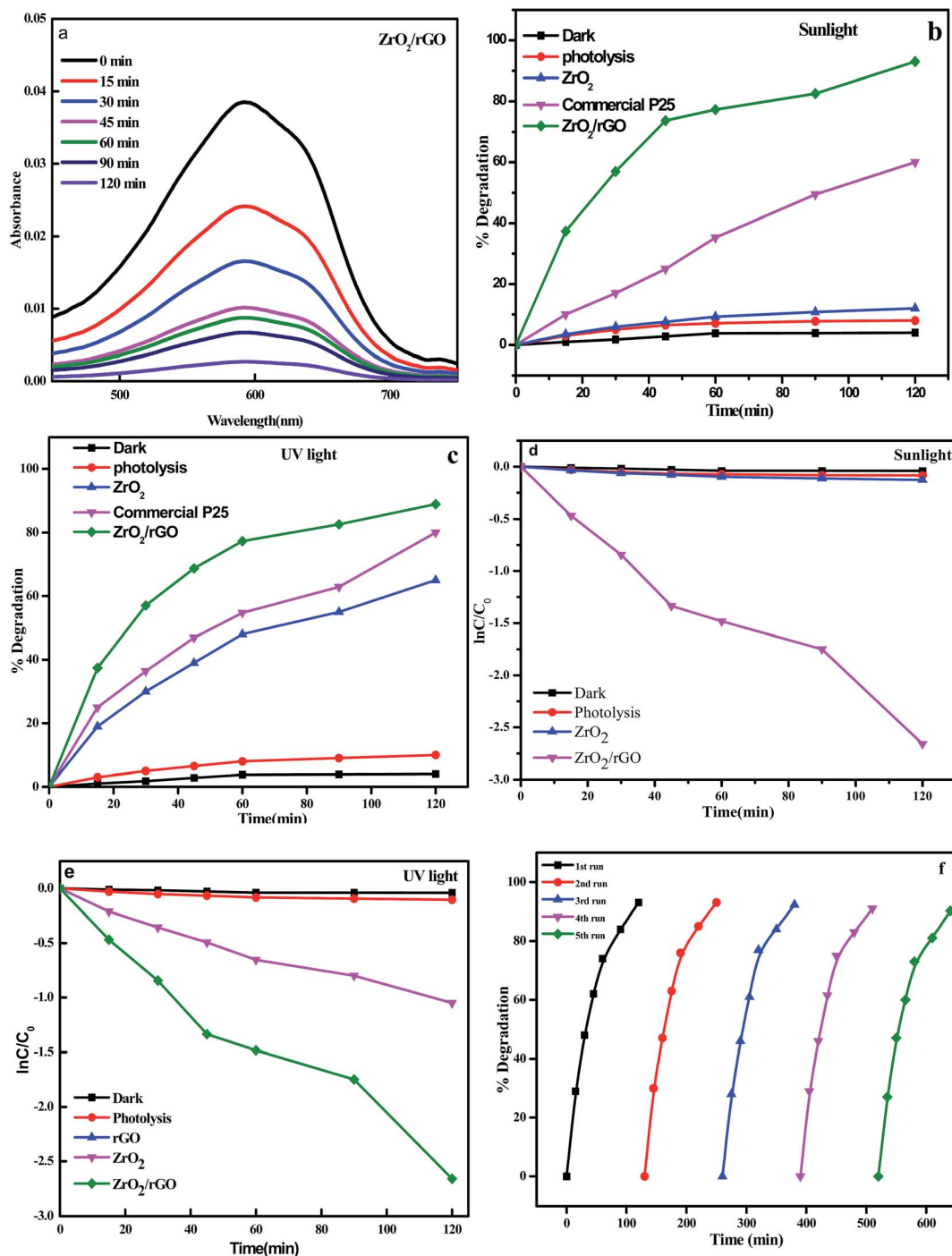


Fig. 12 (a) Spectral absorbance changes with ZrO<sub>2</sub>/rGO photocatalyst; (b) plot of % decomposition of RB 4 under sunlight irradiation; (c) plot of % decomposition versus irradiation time for the degradation of RB 4 under UV irradiation; (d) plot of  $\ln(C/C_0)$  versus irradiation time for the degradation of RB 4 under sunlight irradiation; (e) plot of  $\ln(C/C_0)$  versus irradiation time for the degradation of RB 4 under UV light irradiation; (f) reusability of the ZrO<sub>2</sub>/rGO photocatalyst for five consecutive recycle runs.

towards RB 4 was stronger than that of ZrO<sub>2</sub> NM, which was because rGO presented large surface area to bind dye molecules through the  $\pi$ - $\pi$  conjugation with offset face-to-face direction. This degradation is due to the stepwise energy level structure present in the NC. As per the earlier reported data, the order for the conduction band offset is ZrO<sub>2</sub> (-1.16) > rGO (-4.4), that

shows conduction band for ZrO<sub>2</sub> NM is less negative than compared to rGO.<sup>71</sup> Hence under the sunlight irradiation, system absorbs sufficient amount of energy from irradiated light.

As a result of lower Fermi level and greater electrical conductivity of rGO, the excited electrons will migrate from the





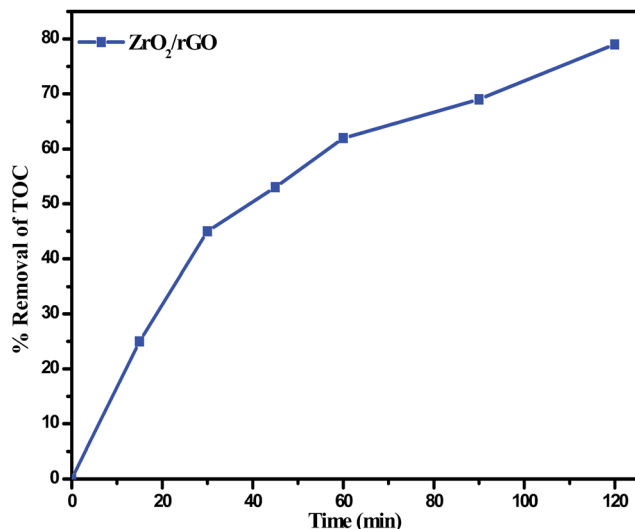


Fig. 13 Percentage TOC removal for the degradation of RB 4 under sunlight in presence of  $\text{ZrO}_2/\text{rGO}$  photo catalyst.

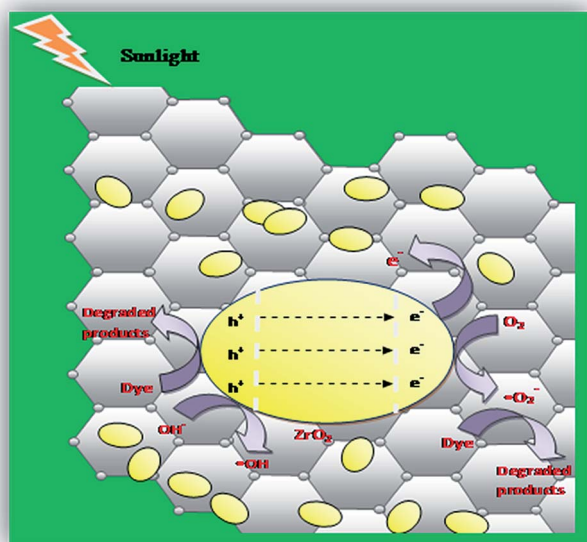


Fig. 14 Mechanism for the photocatalytic degradation of RB 4 over  $\text{ZrO}_2/\text{rGO}$  photocatalyst.

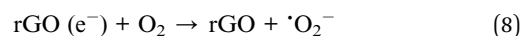
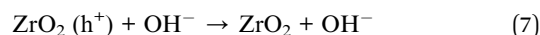
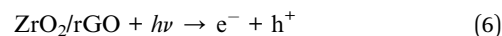
CBs of  $\text{ZrO}_2$  to rGO, leaving  $\text{h}^+$  behind in the  $\text{ZrO}_2$ . The rGO acts as an  $\text{e}^-$  reservoir with  $\text{ZrO}_2$  in this system by accepting the induced-photo electrons. This results in the disconnection of generated electron and hole pairs and increases life span of holes and photoelectrons, thus improving the activity of the photocatalyst dramatically. The obtained results were in consistent with photoluminescence and electrochemical impedance spectroscopic studies. Plots of  $\ln C/C_0$  vs. time for degradation of RB 4 under sunlight and UV light irradiation is represented in Fig. 12d and e. The determined rate constant for  $\text{ZrO}_2$  NM and  $\text{ZrO}_2/\text{rGO}$  NC under sunlight irradiation was found to be 0.0014 and 0.0203  $\text{min}^{-1}$  and under UV light irradiation was found to be 0.0083 and 0.018  $\text{min}^{-1}$  respectively.

The obtained results portrayed excellent photocatalytic performance in the presence of sunlight.

Recycling experiments were performed in order to study the photostability of the photocatalysts under sunlight (Fig. 12f). The recycling experiment is carried out for  $\text{ZrO}_2/\text{rGO}$  NC for five cycles by choosing the same sample and loading a fresh set of dye for degradation in each cyclic runs. The samples were washed for each cycle and dried before starting the next cycle. For four recycles, the percentage of degradation was found to be same and after the fourth cycle, the degradation was decreased which may be due to loss in the sample while washing. Hence it is confirmed that the  $\text{ZrO}_2/\text{rGO}$  NC shows a very good recycling efficiency and it is also stable.

Fig. 13 depicts the time dependency TOC data for degradation of RB 4 under sunlight irradiation. The degradation of RB 4 was characterized by choosing TOC concentration as mineralization index. As the reaction time increases, the TOC removal concentration decreased gradually. Most of the TOC was mineralized at 120 min of irradiation of RB 4 indicating the maximum removal at 78% which played a vital role for practical applications in order to avoid secondary pollution.

When the NC is exposed to sunlight, surface adsorbed oxygen molecules and hydroxyl ions of  $\text{ZrO}_2/\text{rGO}$  NC are the vital active species for the mechanism that reacts with excited electron and holes to convert it into  $\cdot\text{O}_2$  and  $\cdot\text{OH}$  radicals respectively (Fig. 14). These reactive oxygen species degrade the dye by breaking it into simple organics and furthermore into carbon dioxide and water. The main steps involved in the reaction process of photocatalytic degradation under sunlight irradiation is as follows:



## 4. Conclusions

The present work demonstrates an eco-friendly, green route based and simple approach for the rGO synthesis using Cinnamon extract. X-ray diffraction analysis ensured that the  $\text{ZrO}_2$  NM has a monoclinic and tetragonal mixed phase. The increase in intensity of  $\text{ZrO}_2/\text{rGO}$  peak in PXRD confirms the incorporation of  $\text{ZrO}_2$  into rGO sheets and also clearly evidenced from SEM and TEM morphological analysis. Shift of peaks in the Raman spectrum and formation of C–C bond and C–O bond in deconvoluted peak for C 1s XPS spectrum confirms the presence of  $\text{ZrO}_2$  and rGO in  $\text{ZrO}_2/\text{rGO}$  NC. The interaction between the surface of the  $\text{ZrO}_2$  NM and rGO sheets leads to PL quenching and therefore the intensity of PL gets decreased. The  $\text{ZrO}_2/\text{rGO}$  NC serves as a very good photocatalyst for the decomposition of RB 4 under sunlight. The NC shows enhanced photocatalytic activity compared to commercial P25 with 93%



degradation under sunlight. The photocatalyst was found to be stable even after five cyclic runs indicating its high reusability for several practical applications.

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