New green synthesized reduced graphene oxide–
ZrO$_2$ composite as high performance photocatalyst under sunlight

K. Gurushantha,$^a$ K. S. Anantharaju,$^{*bc}$ L. Renuka,$^{ac}$ S. C. Sharma,$^{dc}$
H. P. Nagaswarupa,$^a$ S. C. Prashantha,$^a$ Y. S. Vidy*$^e$ and H. Nagabhushana*$^f$

We report a green route method for the effective reduction of graphene oxide using Cinnamon extract.
ZrO$_2$ 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$_2$ 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.

effective with good thermal stability and electrochemical nature. It serves as an excellent material as an electroactive for the electro catalysis of critical analyses because of its excellent electro catalytic properties.$^3$

Currently, majority of the materials used in photocatalysis are
semiconductor materials, such as TiO$_2$, ZnO and ZrO$_2$, etc., as the
core body of the catalyst. Out of these, ZrO$_2$ is the only metallic oxide which has both oxidizability and reducibility due to its
acidic and basic nature. Since ZrO$_2$ is a kind of p-type semi-conductor,$^4$ 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$_2$ by doping, compounding carbon nano tubes and several other techniques to enhance its photocatalytic activity.$^5$ 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$_2$, the photodegradation of
textile dyes like reactive yellow, rhodamine B, 2,2,4-diphenoxo-
acetic under UV light has been reported earlier.$^6$–$^9$ Also photo-
catalytic activity for the degradation of ethylene and 4-
chlorophenol under UV light by TiO$_2$–ZrO$_2$ binary oxides has
been studied.$^{10}$–$^{12}$ The increased activity under UV light in the
above cases is ascribed to either stability of catalyst/increase in

$^a$Department of Science, Research Center, East West Institute of Technology, Bengaluru – 560091, India
$^b$Department of Chemistry, Dayananda Sagar College of Engineering, Shavige Malleshwara Hills, Kumarswamy Layout, Bengaluru – 560078, India. E-mail: iananothkurupalya@gmail.com
$^c$Dr. D. Premachandra Sagar Centre for Advanced Materials, DSCE, Bangalore – 560078, India
$^d$Department of Mechanical Engineering, Jain University, Advisor, Jain Group of Institutions, Bengaluru – 560092, India
$^e$Department of Physics, Lal Bahadur Shastri Government First Grade College, Bengaluru – 560032, India. E-mail: vidyps.phy@gmail.com
$^f$Prof. C.N.R. Rao Centre for Advanced Materials, Tumkur University, Tumkur – 572103, India. E-mail: bhushanlcf@gmail.com

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.$^1$ 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$_2$ is one among the fundamental transition metal oxide with less toxicity and high chemical inertness.$^2$ In addition, it is biocompatible, cost-effective with good thermal stability and electrochemical nature. It serves as an excellent material as an electroactive for the electro catalysis of critical analyses because of its excellent electro catalytic properties.$^3$
surface area or transfer of charges from ZrO₂ to TiO₂ hence preventing the electron–hole recombination. Recently, the photodegradation of phenol under UV light by mixed oxide TiO₂–ZrO₂ catalyst has been proved to be very efficient compared to P25. Here primarily fast transfer of photo-generated electrons from the conduction band of ZrO₂ to TiO₂ takes place effectively under UV light which is mainly due to its small particle size, high surface area and abundant surface OH groups.33

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 chosen34,35 because of its large specific surface area and an extensive two-dimensional π–π conjugation structure that has excellent electron conductivity.36 Carriers in pristine graphene sheets have been already studied to behave as massless Dirac fermions.37 When combining TiO₂ nanocrystals with graphene, excited electrons of TiO₂ could transfer from the conduction band (CB) to graphene through a percolation mechanism.18 The heterojunction formed at the interface (termed Schottky barrier) separates the photoinduced electron–hole pairs, thereby inhibiting charge recombination.48 First time, Katam and co-workers in 2010 have demonstrated the enhanced photocatalytic activity of graphene-based semiconductor–metal composites.19 After which, Zhang et al.,19 Shen et al.,20 and Zhou et al.,21 performed an one-step hydrothermal methods to prepare graphene–TiO₂ hybrid materials and showed that the composites exhibited enhanced photoactivity towards organic degradation over bare TiO₂. Fan et al.,22 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 H₂ evolution from methanol aqueous solution as compared to pure P25. Graphene (GR)–TiO₂ 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.23 Bearing these results in mind, the study on the use of ZrO₂–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.24–26 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.27,28 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 derivatives29 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 antioxidant 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 ZrO₂–reduced graphene oxide (ZrO₂/rGO) NC. The Cyclic Voltammetry (CV) and Electrochemical Impedance Studies (EIS) were carried out for ZrO₂ NM and ZrO₂/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 ZrO₂/rGO NC was compared to commercial P25 and studied by evaluating its performance in the photodecomposition of RB 4 dye under sunlight illumination. The ZrO₂/rGO NC was shown to exhibit excellent photocatalytic activity as compared to ZrO₂ 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.40 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% H₂O₂ was added. The product obtained was washed with 5% HCl and then with deionized water.31 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.32 The final product was washed with distilled water, dried and stored.

2.4 Preparation of ZrO₂ nanomaterial

Solution combustion synthesis was carried out by using zirconyl nitrate [ZrO(NO₃)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 ZrO₂ nanomaterial (NM) was collected.

### 2.5 Preparation of ZrO₂/rGO NC

100 mg of prepared ZrO₂ 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 ZrO₂ solution with constant stirring and mixture was stirred for about 24 hours to attain ZrO₂/rGO NC. The schematic diagram for synthesis of ZrO₂/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 CuKα (1.541 Å) radiation at a scan rate of 2°/min with nickel filter in the 2θ range 20–70°, 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 Flurolog 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 (1l); long working distance: 50 (2l); 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 cm². In the photocatalytic reactor, dye solution of known ppm was mixed with required amount of synthesized catalyst (ZrO₂/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 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 ZrO₂/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 ZrO$_2$ NM and ZrO$_2$/rGO NC. It could be observed that the crystals are irregular in shape/size and slight agglomeration is observed (Fig. 2a). Here, rGO helps ZrO$_2$ NM to disperse more smoothly in ZrO$_2$/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 TiO$_2$ can more efficiently harvest solar light for photocatalysis.\(^{33}\)

The microstructure of ZrO$_2$/rGO NC is studied by TEM (Fig. 3a and b). Abundant 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 ZrO$_2$ nanoparticles dispersion on its surface while ZrO$_2$ facilitates to avoid the agglomeration of rGO. The crystallite sizes of the deposited ZrO$_2$ nanocrystals are normally 14–18 nm, which is in good agreement with the PXRD results. The intimate interfacial contact between rGO and ZrO$_2$ NC can be observed in HRTEM image (Fig. 3c and e) which shows the lattice of both graphene and ZrO$_2$. Since that the transfer process of charge carriers in rGO-ZrO$_2$ NC is related to the interfacial interaction between graphene and ZrO$_2$ NC. This leads to the expectation for improved photocatalysis process due to good charge transfer property. The growth direction of the 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 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 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 ZrO$_2$/rGO NC, XPS measurements have been carried out (Fig. 4). The existence of carbon and ZrO$_2$ was confirmed by XPS analysis of the ZrO$_2$/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 ZrO$_2$/rGO NC. The existence of graphene sheets were confirmed by the signal for C 1s obtained at the O–C=O (carbonyl 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$_{3/2}$, Zr 3d$_{5/2}$ 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.\(^{34}\) 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$^\cdot$ and H$_2$O. Thus, from the above explanations it was clear that ZrO$_2$ had been incorporated into the rGO layers during synthesis.\(^{35,36}\)

3.3 Raman studies

Raman spectroscopy is a widely used tool for the characterization of $sp^2$ and $sp^3$ hybridized carbon atoms in graphene based materials to differentiate the order and disorder/defects.\(^{37,38}\)
Raman signature of GO, rGO and ZrO$_2$/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 cm$^{-1}$ and G band (the E$_{2g}$ mode of the sp$^2$ carbon atoms) at 1581 cm$^{-1}$. 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. 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. The GO (0.442) shows lesser I$_D$/I$_G$ ratio compared to rGO (0.452), signifying that the rGO include more defects. For the 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 ZrO$_2$ and rGO.

Fig. 3 TEM (a & b), HRTEM (c & d), and SAED pattern (e) of ZrO$_2$/rGO NC.
3.4 PXRD analysis

Fig. 6 shows the PXRD patterns of ZrO$_2$, GO, rGO and ZrO$_2$/rGO NC. The diffraction peaks of synthesized 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 ($\frac{111}{2}$) plane of monoclinic phase. The peak at $\frac{24}{10}$° 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 diffraction from $\sim 10^{°}$ to $\sim 25.6^{°}$. Thus rGO shows a major (002) peak at about 25.6°, which is ascribed to the reduction of GO sheets and restacking into an ordered crystalline structure.$^{42}$

While, the PXRD pattern of ZrO$_2$/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 $\sim 25^{°}$ overlapped with the (110) diffraction peak of the monoclinic ZrO$_2$. The broadening of the diffraction peaks observed in ZrO$_2$/rGO NC indicated the formation of ZrO$_2$–rGO nanocrystallites. The primary size of the
ZrO$_2$ NM and ZrO$_2$/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.

$$\delta = \frac{1}{D^2}$$

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

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 cm$^{-1}$. The FTIR spectra of ZrO$_2$ NM and ZrO$_2$/rGO NC are represented in Fig. 7. The peaks at 3443 cm$^{-1}$ correspond to hydroxyl stretching and 1635 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; Nejatzadeh-Barandozi & Enferadi, 2012). The broad absorption peaks at around 2930 cm$^{-1}$ are due to the characteristic stretching vibration of CH$_2$. The peak at 1387 cm$^{-1}$ is due to the presence of OH deformations in the C–OH groups and at 1656 cm$^{-1}$ is due to the presence of sp$^2$ bond of graphite in GO. The band at 1060 cm$^{-1}$ is assigned to C–O (epoxy) group which is consistent with XPS studies while the band at 1100 cm$^{-1}$ is usually contributed to alkoxy group.$^{44-45}$

The absence of peaks at 1060, 1210 and 1656 cm$^{-1}$ in the ZrO$_2$/rGO confirms that GO has been reduced to rGO. The peak at 470 cm$^{-1}$ can be assigned to the Zr–O vibration. These observations may signify that most oxygen functionalities in the GO were removed and ZrO$_2$/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 ($E_{VB}$) and the conduction band ($E_{CB}$) gives energy band gap ($E_g$), from which the electronic structure of ZrO$_2$/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 ‘OH and O$_2$$^•$ by ZrO$_2$/rGO were related to the energy band gap and also to redox potential of different reactive species.$^{46-48}$ 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 the plot of $F(R)^2$ on the Y-axis versus energy (eV) on the X-axis gives the value of the $E_g$. The $E_g$ value of ZrO$_2$ NM and ZrO$_2$/rGO NC was found to be 4.6 and 3.25 eV respectively. This $E_g$ value for ZrO$_2$/rGO was very close to the $E_g$ value reported by Braj Raj Singh and his co-workers.$^{49}$ The very

---

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

| Samples     | FWHM (radians) | Crystallite size (nm) by Scherrer’s method | Micro strain ($\varepsilon$) $\times 10^{-3}$ | SF     | $\delta$ $(10^{15}$ lines per m$^2$) |
|-------------|----------------|------------------------------------------|------------------|--------|-----------------------------|
| ZrO$_2$     | 0.653          | 14.7                                     | 2.68             | 0.438  | 4.5                         |
| ZrO$_2$/rGO | 0.627          | 15.2                                     | 2.56             | 0.460  | 4.3                         |
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.\textsuperscript{50,51} 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.\textsuperscript{52} 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.\textsuperscript{54} 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.\textsuperscript{55–58} Hence due to the interaction of ZrO$_2$ surface with rGO sheets, PL quenching appears with decrease in intensity.\textsuperscript{57} 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
higher PL intensity than ZrO2/rGO NC, indicating ZrO2 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 ZrO2 nanomaterial have been widely investigated by many researchers, however, the role of particle size and the crystal phases of ZrO2 nanomaterial has not yet been analyzed in a systematical manner. Harrison et al. proved that the luminescent centers constituted by Zr4+ 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 ZrO2 nanomaterials. The broad emission band arises due to the singly ionized oxygen vacancies defects in ZrO2 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 ZrO2 nanomaterials. The electrons were trapped rapidly by the oxygen vacancies (VO) thereby producing F centers. The Zr4+ ions adjacent to the bulk oxygen vacancies had retained the electrons leading to the formation of Zr3+ ions. 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:

\[ \text{ZrO}_2 + h\nu \rightarrow h^+ + e^- \]  
\[ V_o + e^- \rightarrow F \text{ center} \]  
\[ F \text{ center} + h^+ \rightarrow (V_o)^* \rightarrow V_o + h\nu \]

The mechanism possibly involved in the emission phenomenon of the sample ZrO2 has been demonstrated with the help of schematic diagram as shown in Fig. 10. The major source of defects centers in ZrO2 sample was oxygen vacancies/interstitial, Zr vacancies/interstitials (Zr4+ interstitials).

Moreover, the broad peak at 482 nm has been slightly shifted towards lower wavelength (467 nm) in ZrO2/rGO NC. The decrease in PL intensity of the composite stems from the band alignment at the interface of ZrO2 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 (NaNO3) solution using three electrode cell arrangement. Fig. 11a shows the capacitance variations of the ZrO2/rGO electrodes at various scanning speed from 10 to 50 mV s⁻¹. 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 ZrO2 NM and ZrO2/rGO NC were run at 30 mV s⁻¹ and depicted in Fig. 11b. It is seen from the figure that ZrO2 NM found to have a very broad semi-rectangular plot while ZrO2/rGO NC has less broad semi rectangular shape. This indicated that the capacitance value for ZrO2 NM is more compared to ZrO2/rGO.

Electrochemical impedance spectroscopy (EIS) were recorded by the application of an AC voltage with 10 mV amplitude in the frequency range of 10⁵ to 10⁻² Hz in 0.1 M sodium nitrate (NaNO3) 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 ZrO2–electrolyte interfaces. Inset 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 (Rct) value in terms of ohms, which generally associated with the pore structure of electrode materials and faraday pseudo capacitive reaction. The less impedance arc radius in Nyquist plots indicates the faster electron transfer rate. The diameter for ZrO2 NM and ZrO2/rGO NC were found to be 14 and 8 Ω respectively. The diameter for ZrO2/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 NaNO3 solution is depicted as element Rш. Element Rт corresponds to electrolyte resistance present in porosity of the
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. UV-Vis absorption spectra of the ZrO$_2$ NM and ZrO$_2$/rGO NC are shown in Fig. 12a. In the present work, photocatalytic activity is studied for ZrO$_2$ NM and ZrO$_2$/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 ZrO$_2$/rGO NC than 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 ZrO$_2$ NM and ZrO$_2$/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 ZrO$_2$ NM and ZrO$_2$/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 ZrO$_2$/rGO NC has got higher photocatalytic activity compared to commercial P25. It was evident that the absorbent capacity of ZrO$_2$/rGO NC.
towards RB 4 was stronger than that of ZrO$_2$ NM, which was because rGO presented large surface area to bind dye molecules through the π–π 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$_2$ (−1.16) > rGO (−4.4), that shows conduction band for ZrO$_2$ NM is less negative than compared to rGO.$^{73}$ 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
CBs of ZrO2 to rGO, leaving h+ behind in the ZrO2. The rGO acts as an e− reservoir with ZrO2 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/C0 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 ZrO2 NM and ZrO2/rGO NC under sunlight irradiation was found to be 0.0014 and 0.0203 min−1 and under UV light irradiation was found to be 0.0083 and 0.018 min−1 respectively.

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 ZrO2 NM has a monoclinic and tetragonal mixed phase. The increase in intensity of ZrO2/rGO peak in PXRD confirms the incorporation of ZrO2 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 ZrO2 and RGO in ZrO2/rGO NC. The interaction between the surface of the ZrO2 NM and rGO sheets leads to PL quenching and therefore the intensity of PL gets decreased. The ZrO2/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.

Acknowledgements

The authors S. C. Sharma and K. S. Anantharaju thank VGST, Govt. of Karnataka, India for sanctioning the research project (VGST/CESEM/2015-16).

References

1 A. Umar, M. S. Akhtar, A. Al-Hajry, M. S. Al-Assiri, G. N. Dar and M. Saif Islam, Chem. Eng. J., 2015, 262, 588.
2 B. Liu, J. Hu and J. S. Foord, Electrochem. Solid-State Lett., 2011, 14, D22.
3 A. T. Ezhil Vilian, S.-M. Chen, M. Ajmal Alib and F. M. A. Al-Hemaidib, RSC Adv., 2014, 4, 30358.
4 K. Samson, M. S. liwa, R. P. Socha, K. Gora-Marek, D. Mucha, D. Rutkowska-Zbik, J. F. Paul, M. Ruggiero-Mikolajczyzk, R. Grabowski and J. Sloczynski, ACS Catal., 2014, 4, 3730.
5 Y. L. Yu, P. Zhang, Y. J. Kuang, Y. H. Ding, J. H. Yao, J. J. Xu and Y. A. Cao, J. Phys. Chem. C, 2014, 118, 20982.
6 C. Wu, X. Zhao, Y. Ren, Y. Yue, W. Hua, Y. Cao, Y. Tang and Z. Gao, J. Mol. Catal. A: Chem., 2005, 229, 233.
7 H. Zheng, K. Liu, H. Cao and X. Zhang, J. Phys. Chem. C, 2009, 113, 18259.
8 M. Alvarez, T. Lopez, J. A. Odrozolab, M. A. Centenoc, M. I. Dominguezb, M. Montesb, P. Quintanab, D. H. Aguilarb and R. D. Gonzalez, Appl. Catal., B, 2007, 73, 34.
9 B. Neppolian, H. Choi, S. Sathivel, V. M. B. Arabindoo, B. Neppolian and V. Murugesan, J. Hazard. Mater., 2002, 89, 303.
10 B. M. Reddy and A. Khan, Catal. Rev., 2007, 47, 257.
11 A. Burri, N. Jiang and S. E. Park, Catal. Sci. Technol., 2012, 2, 514.
12 B. Neppolian, Q. Wang, H. Yamashita and H. Choi, Appl. Catal., A, 2007, 333, 264.
13 A. Kambur, G. S. Pozan and I. Boz, Appl. Catal., B, 2012, 115–116, 149.
14 L. L. Tan, S. P. Chai and A. R. Mohamed, ChemSusChem, 2012, 5(10), 1868.
15 Q. Xiang, J. Yu and M. Jaroniec, Chem. Soc. Rev., 2012, 41(2), 782.
16 Q. Xiang, J. Yu and M. Jaroniec, Nanoscale, 2011, 3(9), 3670.
17 K. S. Novoselov, A. K. Geim, S. V. Morozov, D. Jiang, M. I. Katsnelson, I. V. Grigorieva, S. V. Dubonos and A. A. Firsov, Nature, 2005, 438(7065), 197.
18 G. Williams, B. Seger and P. V. Kamat, ACS Nano, 2008, 2(7), 1487.
19 H. Zhang, X. Lv, Y. Li, Y. Wang and J. Li, ACS Nano, 2009, 4(1), 380.
20 J. Shen, B. Yan, M. Shi, H. Ma, N. Li and M. Ye, J. Mater. Chem., 2011, 21(10), 3415.
21 K. Zhou, Y. Zhu, X. Yang, X. Jiang and C. Li, New J. Chem., 2011, 35(2), 353.
22 W. Fan, Q. Lai, Q. Zhang and Y. Wang, J. Phys. Chem. C, 2011, 115(21), 10694.
23 X. Yang, J. Qin, Y. Li, R. Zhang and H. Tang, J. Hazard. Mater., 2013, 261, 342.
24 K. Gurushantha, K. S. Anantharaju, S. C. Sharma, H. P. Nagaswarupab, S. C. Prashanthac, K. R. Vishnud, L. Renukab, Y. S. Vidyad and H. Nagabhushana, J. Alloys Compd., 2016, 685, 761.
25 L. Renuka, K. S. Anantharaju, S. C. Sharma, H. P. Nagaswarupa, S. C. Prashantha, H. Nagabhushana and Y. S. Vidyaj, J. Alloys Compd., 2016, 672, 622.
26 P. Mishra, Y. P. Singh, H. P. Nagaswarupa, S. C. Sharma, Y. S. Vidyad, S. C. Prashantha, H. Nagabhushana, K. S. Anantharaju, S. Sharma and L. Renuka, J. Alloys Compd., 2016, 685, 656.
27 P. Ranasinghe, S. Pigera, G. A. S. Premakumuraa, P. Galappathgyb, G. R. Constantine and P. Katulandaa, BMC Complementary Altern. Med., 2013, 13, 275.
28 K. Sambaiah and K. Srinivasan, Nahrung, 1991, 35, 47.
29 G. Singh, S. Maurya, M. P. DeLampasona and C. A. Catala, Food Chem. Toxicol., 2007, 45, 1650.
30 W. S. Hummers and R. E. Offeman, J. Am. Chem. Soc., 1958, 80, 1339.
31 S. Gurunathan, J. W. Han, V. Eppakayalaa and J. Kim, Colloids Surf., B, 2013, 102, 772.
32 D. Suressb, Udayabhanuss, H. Nagabhushana and S. C. Sharma, Mat. Lett., 2015, 142, 4.
33 X. Chen, L. Liu, P. Y. Yu and S. S. Mao, Science, 2011, 331, 476.
34 S. Calderon, R. E. Galindo, N. Benito, C. Palico, A. Cavaleiro and S. Carvalho, J. Mater. Lett., 2013, 7065, 197.
35 H. Teymouriani, A. Salimi, S. Firoozia, A. Koranib, S. Soltanian, Electrochim. Acta, 2014, 143, 196.
36 H. L. Wang, J. T. Robinson, X. L. Li and H. J. Dai, J. Am. Chem. Soc., 2009, 131, 9910–9911.
37 J. Shen, T. Li, Y. Long, M. Shi, N. Li and M. Ye, Carbon, 2012, 50, 2134–2140.
38 D. C. Luo, G. X. Zhang, J. F. Liu and X. M. Sun, J. Phys. Chem. C, 2011, 115, 11327–11335.
39 H. Teymourian, A. Salimi, S. Firoozi, A. Korani and S. Soltanian, Electrochim. Acta, 2014, 143, 196.
40 H. L. Wang, J. T. Robinson, X. L. Li and H. J. Dai, J. Am. Chem. Soc., 2009, 131, 9910–9911.
41 P. Wanga, J. Wangb, X. Wangab, H. Yuab, Y. Juba, M. Lei and Y. Wang, Appl. Catal., B, 2013, 132–133, 452–459.
42 S. Kumar, J. G. Sharma, S. Majib and B. D. Malhotrab, Biosens. Bioelectron., 2016, 15, 497.
43 D. C. Marcanob, D. V. Kosynkinb, J. M. Berlin, A. Sinitskib, Z. Sun, A. Slesarevb, L. B. Alemanyb, W. Lu and J. M. Turn, ACS Nano, 2010, 4, 4806.
44 Y. Xu, H. Bai, G. Lu, C. Li and G. Shi, J. Am. Chem. Soc., 2008, 130, 5856.
45 X. Li, Q. Wang, Y. Zhao, W. Wu, J. C. Feng and H. Meng, J. Colloid Interface Sci., 2013, 141, 69.
46 M. Shoeb, B. R. Singh, J. A. Khan, W. Khan, B. N. Singh, H. B. Singh and A. H. Naqvi, Adv. Nat. Sci.: Nanosci. Nanotechnol., 2013, 4, 035015.
47. J. A. Khan, M. Qasim, B. R. Singh, S. Singh, M. Shoeb, W. Khan, D. Das and A. H. Naqvi, *Spectrochim. Acta, Part A*, 2013, 109, 313.
48. C. D. Vecitis, K. R. Zodrow, S. Kang and M. Elimelech, *ACS Nano*, 2010, 4, 5471.
49. B. Raj Singh, M. Shoeb, W. Khan and A. H. Naqvi, *J. Alloys Compd.*, 2015, 651, 598.
50. K. Woan, G. Pyrgiotakis and W. Sigmund, *Adv. Mater.*, 2009, 21, 2233.
51. Y. Yu, J. C. Yu, C. Y. Chan, Y. K. Che, J. C. Zhao, L. Ding, W. K. Ge and P. K. Wong, *Appl. Catal., B*, 2005, 61, 1.
52. Y. L. Min, K. Zhang, W. Zhao, F. C. Zheng, Y. C. Chen and Y. G. Zhang, *Chem. Eng. J.*, 2012, 193–194, 203.
53. W. Zheng, K. H. Bowen, J. Li, I. Dabkowska and M. Gutowski, *J. Phys. Chem. A*, 2005, 109, 11521.
54. Y.-C. Yeo, T.-J. King and C. Hu, *J. Appl. Phys.*, 2002, 92, 7266.
55. Y. B. Shahar, F. Scotognella, N. Waiskopf, I. Kriegel, S. D. Conte, G. Cerullo and U. Banin, *Small*, 2015, 11, 462.
56. W. T. Chen, T. T. Yang and Y. J. Hsu, *Chem. Mater.*, 2008, 20, 7204.
57. Z. Wang, C. Yang, T. Lin, H. Yin, P. Chen, D. Wan, F. Xu, F. Huang, J. Lin, X. Xie and M. Jiang, *Adv. Funct. Mater.*, 2013, 23, 5444.
58. D. Zhang, B. Hua, D. Guan and Z. Luo, *Catal. Commun.*, 2016, 76, 7.
59. C. Lin, C. Zhang and J. Lin, *J. Phys. Chem. C*, 2007, 111, 3300.
60. A. Mondal, A. Zachariah, P. Nayak and B. B. Nayak, *J. Am. Ceram. Soc.*, 2010, 93, 387.
61. H. Cao, X. Qiu, B. Luo, Y. Liang, Y. Zhang, R. Tan, M. Zhao and Q. Zhu, *Adv. Funct. Mater.*, 2004, 14, 243.
62. D. E. Harrison, N. T. Melamed and E. C. Subbarao, *J. Electrochem. Soc.*, 1963, 110, 23.
63. H. Liu, L. Feng, X. Zhang and Q. Xue, *J. Phys. Chem.*, 1995, 99, 332.
64. S. Kumar and A. K. Ojha, *J. Alloys Compd.*, 2015, 644, 654.
65. N. G. Petrlik, D. P. Taylor and T. M. Orlando, *J. Appl. Phys.*, 1999, 85, 6770.
66. A. Zaban, A. Meier and B. A. Gregg, *J. Phys. Chem. B*, 1997, 101, 7985.
67. F. F. Santiago, G. G. Belmonte, J. Bisquert, A. Zaban and P. Salvador, *J. Phys. Chem. B*, 2002, 106, 334.
68. M. Toupina, D. Belanger, I. R. Hill and D. Quinn, *J. Power Sources*, 2005, 140, 203.
69. H. Liu, S. Cheng, M. Wu, H. Wu, J. Zhang, W. Li and C. Cao, *J. Phys. Chem. A*, 2000, 104, 7016.
70. L. S. Wang, M. W. Xiao, X. J. Huang and Y. D. Wu, *J. Hazard. Mater.*, 2009, 161, 49.
71. F. T. Johra and W. G. Jung, *Appl. Catal., A*, 2015, 491, 52.