



Mechanistic insight into the adsorption and photocatalytic activity of a magnetically separable γ -Fe₂O₃/Montmorillonite nanocomposite for rhodamine B removal

Is Fatimah^{a,*}, Gani Purwiandono^a, Arif Hidayat^b, Suresh Sagadevan^c, Azlan Kamari^d

^a Department of Chemistry, Faculty of Mathematics and Natural Sciences, Universitas Islam Indonesia, Kampus Terpadu UII, Jl. Kaliurang Km 14, Sleman, Yogyakarta, Indonesia

^b Department of Chemical Engineering, Faculty of Industrial Technology, Universitas Islam Indonesia, Kampus Terpadu UII, Jl. Kaliurang Km 14, Sleman, Yogyakarta, Indonesia

^c Nanotechnology and Catalysis Research Center, University Malaya, Malaysia

^d Chemistry Department, Universiti Pendidikan Sultan Idris Perak, Perak 35900, Malaysia

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ABSTRACT

The magnetically separable nanocomposite of γ -Fe₂O₃/montmorillonite (γ -Fe₂O₃/Mt) has been synthesized by a hydrothermal method. The as-obtained nanocomposite is composed of a single phase of γ -Fe₂O₃ nanoparticles homogeneously dispersed in a montmorillonite structure with high photocatalytic activity for rhodamine B photodegradation. Physicochemical characterization revealed particle sizes ranging from 10 to 30 nm with a band-gap energy of 2.3 eV. This study reveals a significant contribution of the adsorption mechanism to the degradation mechanism. The nanocomposite could be easily separated and reused by magnetic separation owing to its magnetism of 22 emu/g. The nanocomposite is also reusable without any chemical change until five cycles.

1. Introduction

Organic compound-containing wastewater is a serious problem in many industries that require emerging technology for removal. The persistence properties of organic compounds bring potential damage to aquatic environments and cause the accumulation of vast environmental problems, which must raise concerns before allowing their discharge to the environment [1,2]. Organic compounds classified as dyes, especially azo dyes, are considered serious and harmful contaminants characterized as biologically toxic (carcinogenic, mutagenic, and teratogenic), irritants, and infectious. The textile, printing, paint, and pharmaceutical industries highly consume these dye compounds; unfortunately, more than 15% of the consumed amount is discharged into wastewater [3]. Evidently, these dyes are the major pollutants from many industries and are clearly major hazards to groundwater. Rhodamine B (RhB) is a significant and well-known basic xanthene dye that is a major ingredient in the above-mentioned industries. The discharge of RhB-containing wastewater is a serious threat to the environment, even at low concentrations in water; thus, it is a representative target organic pollutant for photocatalytic efficiency examination.

Heterogeneous photocatalytic oxidation, as part of the advanced oxidation process (AOPs), has proven to be a promising technology for organic-containing water. Its removal mechanism, which includes oxidation, for the target compound is advantageous over the use of adsorption technologies [4,5]. The method depends on the photocatalytic activity of semiconductor photocatalysts, and its reusability and recyclability are value-added characteristics of the technology.

Adsorption capability, stability, easy separation, and high photocatalytic activity in the visible range are the significant properties required for materials to be used as photocatalysts [6]. Recent studies showed that adsorption capability of the photocatalyst is the important function in photocatalysis mechanism, and based on this, semiconductor immobilized onto porous material is ideal for the mechanism due to the surface area availability [7,8]. The development of nanomaterials with magnetic photocatalysts involves a combination of these properties and has been described as an efficient method for improving the effectiveness of the photocatalytic oxidation process [9–11]. Iron oxide-based magnetic nanoparticles are well known magnetically separable materials, and their combination with supportive solids with high porosity and chemical stability has received great attention. α -Fe₂O₃, γ -Fe₂O₃, and Fe₃O₄

* Corresponding author.

E-mail address: isfatimah@uii.ac.id (I. Fatimah).

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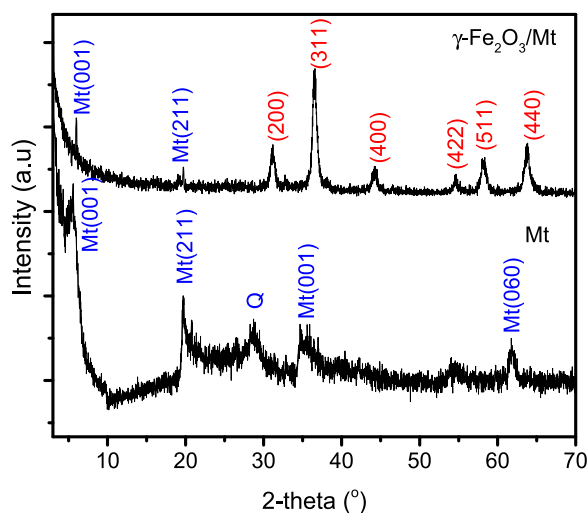


Fig. 1. XRD pattern of materials.

have been widely studied as magnetic active nanoparticles; within this scheme, nanocomposite development using silica-, carbon-, and clay-based materials has been attempted [12]. The swelling capability and modifiable properties of their chemical structures cause clay materials to produce homogeneously distributed magnetic nanoparticles. Surface properties potentially contribute to this activity via an adsorption mechanism [13–15]. In this work, dispersed $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles in montmorillonite (Mt) clay support were prepared. The aim of the present work was to determine the characteristics of materials and the photocatalytic features for the photocatalytic degradation of RhB in aqueous solutions as a model pollutant. In addition, this study provides mechanistic insight into adsorption and photocatalytic properties, especially with magnetically separable characteristics.

2. Materials and methods

2.1. Materials

The Mt used in this work was obtained from Pacitan, East Java, Indonesia. The material was previously washed with water, dried in an oven, and ground to a size of 200 mesh before use. The cation exchange capacity of the material was 69 meq/g. Chemicals consisting of FeCl_3 , RhB, NaOH, ethylene diamine tetra acetate (EDTA), isopropanol, and H_2O_2 were purchased from Merck (Darmstadt, Germany).

2.2. Preparation of nanocomposites

First, dispersed $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles in Mt ($\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$) was prepared using a Mt suspension, by dissolving Mt powder in water at a concentration of 10% wt. An iron oxide precursor was prepared, referring to the previous syntheses of $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles ($\gamma\text{-Fe}_2\text{O}_3$ NPs) by dissolving $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ and $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ with a $\text{Fe}^{3+}/\text{Fe}^{2+}$ molar ratio of 2:1 in double distilled water [16]. The precursor solution was continuously stirred before it was mixed with the Mt suspension at a theoretical percentage of Fe of 10% wt, followed by stirring for 1 h. Into the mixture, the NaOH solution was added slowly and then poured into a Teflon-lined autoclave and kept at 150 °C overnight. The obtained precipitate was dried in an oven at 100 °C and further calcined at 200 °C for 4 h to obtain powder samples, which were designated as $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$.

For comparison, $\gamma\text{-Fe}_2\text{O}_3$ was also synthesized using the same precursor, hydrothermal treatment condition, and calcination temperature with the synthesis of $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$, but without dispersion into a clay suspension.

2.3. Material characterization

The physicochemical properties of the samples were characterized using field emission-scanning electron microscopy (FE-SEM), transmission electron microscopy (TEM), X-ray diffraction (XRD), X-ray photoelectron spectroscopy (XPS), gas sorption analysis, ultraviolet diffuse reflectance spectrophotometry (UV-DRS), and Fourier transform infrared. A Phenom X instrument (US) was used for SEM analysis, while TEM images were taken using a JEOL TEM 2010 transition electron microscope operated at an applied voltage of 200 kV. The XRD of XPERT-PRO with a Ni-filtered $\text{Cu K}\alpha$ radiation ($\lambda = 0.154060$ nm) was used for analysis at a range of 2–70°. The Thermo Nicolet Avatar system (Madison, USA) was used to examine the changes in the surface functional groups of the materials, while the XPS was performed on a V.G. Scientific ESKALAB MKII instrument. Monochromatic $\text{Al K}\alpha$ radiation with a photon energy of 1486.6 ± 0.2 eV was used for analyses. Before analysis, approximately 0.2 mg of sample was slightly pressed into a small pellet of 15 mm in diameter and then mounted on the sample holder for degassing for 4 h to achieve a dynamic vacuum of $< 10^{-8}$ Pa. A vibration sample magnetometer (VSM)-BHV-5 (Japan) was used for the magnetism of the composite.

2.4. Adsorption and photocatalytic experiments

The adsorption experiments were conducted in a batch adsorption system using a horizontal shaker. The experimental setup for the photocatalytic activity measurements included a 40-W xenon lamp-

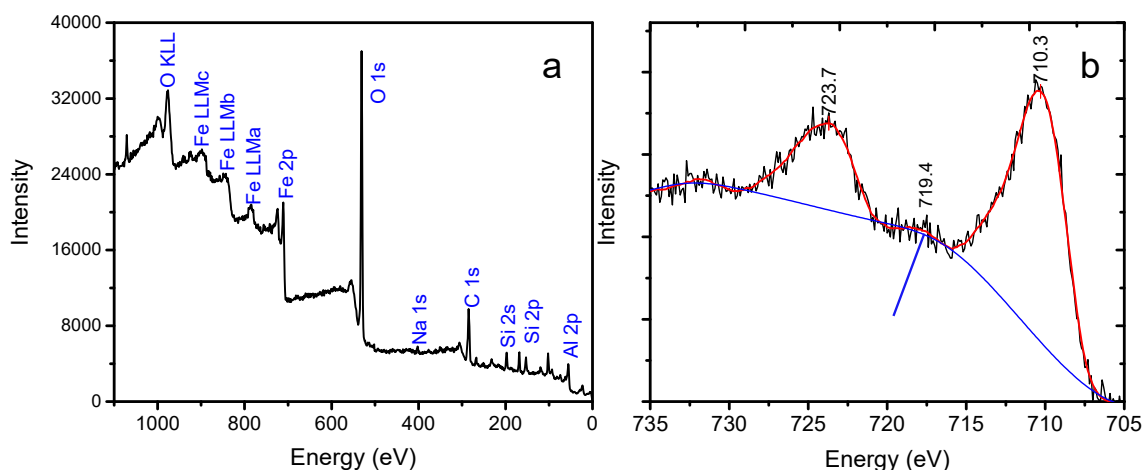


Fig. 2. XPS spectra of $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ (a) survey scan (b) Fe 2p spectrum.

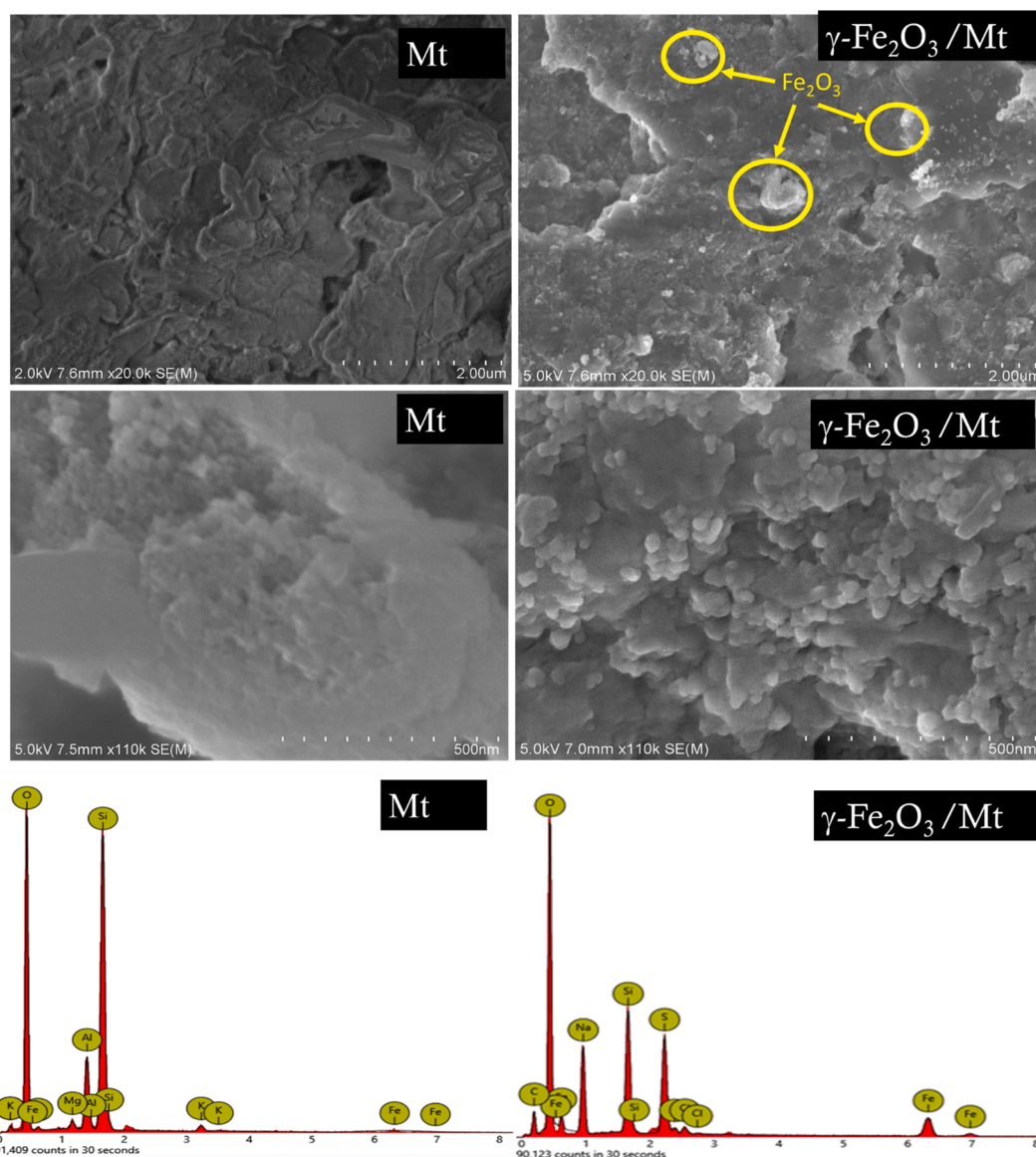


Fig. 3. SEM-EDX of materials.

equipped batch photoreactor. The entire photoreactor system was maintained at room temperature using a water cooler. For each photocatalytic activity test, the photocatalyst powder was mixed with the dye solution with stirring and light exposure. Samples from the adsorption and photocatalytic activity tests were collected at a certain time, and the RhB concentration was determined using the colorimetric method with a U-2010 UV-visible spectrophotometer (Hitachi, Singapore). The degradation efficiency (DE; %) in the photocatalytic activity test was calculated using Eq. (1).

$$DE(\%) = \frac{C_0 - C_t}{C_0} \times 100 \quad (1)$$

where C_0 and C_t are the initial RhB concentration and concentration of RhB at the sampling time t .

To ensure the degradation mechanism, experiments on effect of scavenger to the kinetics of RhB degradation was also performed. High performance liquid chromatography (HPLC) on Agilent 1260 (Singapore) was employed for analysis. Analysis was performed on the condition: analytical C81 column as stationary phase, mobile phase of acetonitrile:phosphate buffer pH 3.5 (1:1), ambient temperature (28–32 °C), flow rate 1.2 mL/min.

3. Results and discussion

3.1. Material characterization

Physically, $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ was prepared in the form of black powder and easily attracted by the magnet field. Fig. 1 shows the XRD profile of the materials. The Mt sample demonstrated peaks associated with the structure at 5.59° , 19.69° , and 34.67° , which corresponded to the (001), (211), and (004) planes, respectively, referring to JCPDS 43-0688 [17]. $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ represents some peaks attributed to (220), (311), (400), (422), (511), and (440), which are in agreement with the values of the standard data (JCPDS no. 4-755) [18–21]. In addition, the (001) and (211) planes are still maintained in the same position, reflecting no such increase or decrease in the basal spacing d_{001} of the interlayer structure. The peak associated with quartz as an impurity disappeared. By using the Scherer formula for the (311) plane, the particle size of $\gamma\text{-Fe}_2\text{O}_3$ was estimated at approximately 23.6 nm [22].

The obtained single phase of $\gamma\text{-Fe}_2\text{O}_3$ in the composite was mainly related to the easy dispersion of Fe(III) ions within the interlayer space of Mt due to the cation exchange capacity of Mt. We can also conclude that the optimum sintering temperature for producing the $\gamma\text{-Fe}_2\text{O}_3$ phase

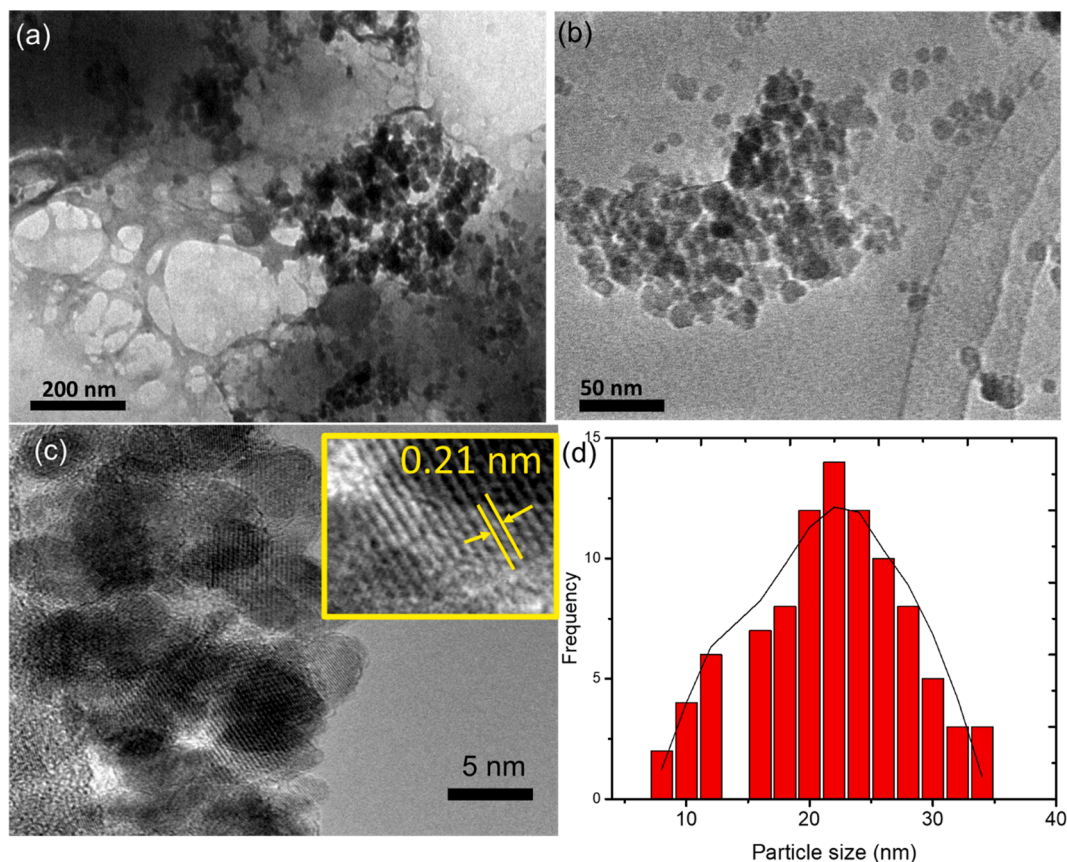


Fig. 4. (a-c) TEM image of γ - Fe_2O_3 /Mt (d) particle size distribution of γ - Fe_2O_3 in the composite.

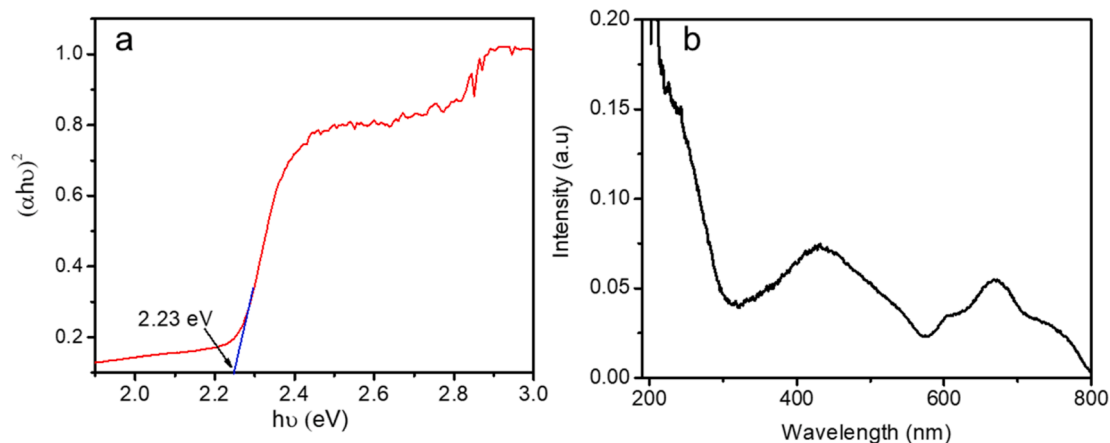
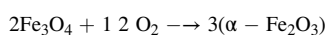
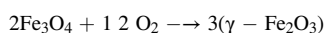
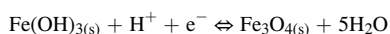
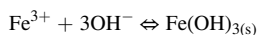


Fig. 5. (a) UV-DRS, and (b) PL spectrum spectrum of γ - Fe_2O_3 /Mt.

is 400 °C. Meanwhile, other possible phases such as Fe_3O_4 , α - Fe_2O_3 , and ϵ - Fe_2O_3 can be formed via the following reaction steps [23]:



The surface composition of γ - Fe_2O_3 /Mt was further demonstrated using XPS with the spectra presented in Fig. 2. The survey scan represents some peaks attributed to Si, Al, O, and Fe, indicating the major composition of the composite. In addition, the low-intensity peak at 400

eV is the characteristic peak of Na 1 s, which is consistent with the naturally occurring Mt structure. The deconvolution of the Fe 2p spectra exhibits two appearing peaks at 723.7 and 710.3 eV, respectively, which are attributed to Fe 2p_{1/2} and Fe 2p_{3/2}, respectively [24–27]. No signal or shoulder representing the presence of the Fe²⁺ ion, which usually exists at approximately 708 eV, was observed. This confirmed that the addition of the shakeup satellite peak occurred at 719.4 eV when no Fe²⁺ ion was found, consistent with the XRD pattern.

The effect of the γ - Fe_2O_3 nanoparticle dispersion on the morphology was investigated using FE-SEM. The SEM images in Fig. 3c depict different surface patterns in which γ - Fe_2O_3 /Mt shows dispersed irregular forms of γ - Fe_2O_3 on the surface (as marked in the yellow dashed circles). With higher magnification, the rounded cubic and irregular

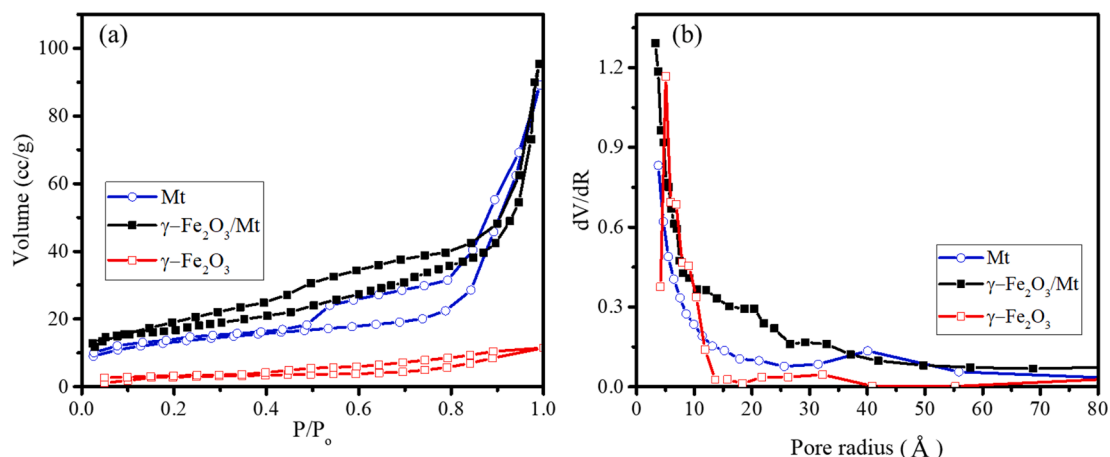


Fig. 6. a. Adsorption/desorption profile (b) Pore size distribution of $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ in comparison with Mt and $\gamma\text{-Fe}_2\text{O}_3$.

spherical forms are confirmed. Further structural identification using TEM and high-resolution (HRTEM) revealed that the particle size ranged from 10 to 30 nm (Fig. 4d). These sizes are consistent with the calculated particle size from the XRD measurement, which was close to 23.6 nm on average. Furthermore, the HRTEM images of the irregular forms exhibited the distances of the adjacent lattice fringes, with an average of 0.21 nm, indicating agreement with the interplanar distance of the $\gamma\text{-Fe}_2\text{O}_3$ (311) space, in reference to JCPDS, Ref. Code: 024-081 [25,28].

With respect to their photocatalytic application, the optical properties of the composites were identified using UV-DRS and photoluminescence spectrum measurements. The Tauc plot of photon energy ($h\nu$) against the $(\alpha h\nu)^2$ curve based on the Kubelka-Munk theory of materials is presented in Fig. 5. Conclusively, the plots show that the E_g of $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ is 2.23 eV. This value is higher than the band-gap values

of $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles that were reported in previous articles, in which the synthesized $\gamma\text{-Fe}_2\text{O}_3$ expressed a band gap of 2.03 eV [2228]. The increasing band-gap energy is related to the smaller particle size due to the quantum size effect and/or crystallite defect. A similar phenomenon was found in the dispersion of $\gamma\text{-Fe}_2\text{O}_3$ into porous Ni and in the composite of $\gamma\text{-Fe}_2\text{O}_3/\text{b-TiO}_2$ [29,30]. The narrow band-gap energy extends the photoresponse to visible-light and near-infrared regions, which is the benefit of the low-cost photocatalytic system [31,32].

Furthermore, the PL spectrum for $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ ranged from 350 to 700 nm (Fig. 5b). The excitation peaked at around 426 and 659 nm, and the peak at 659 nm was attributed to the $6A_1 \rightarrow 4T_1(4G)$ ligand field transition of Fe^{3+} . The data represent the condition that leads to the capability of composites to interact with photons in the UV and visible regions [33,34].

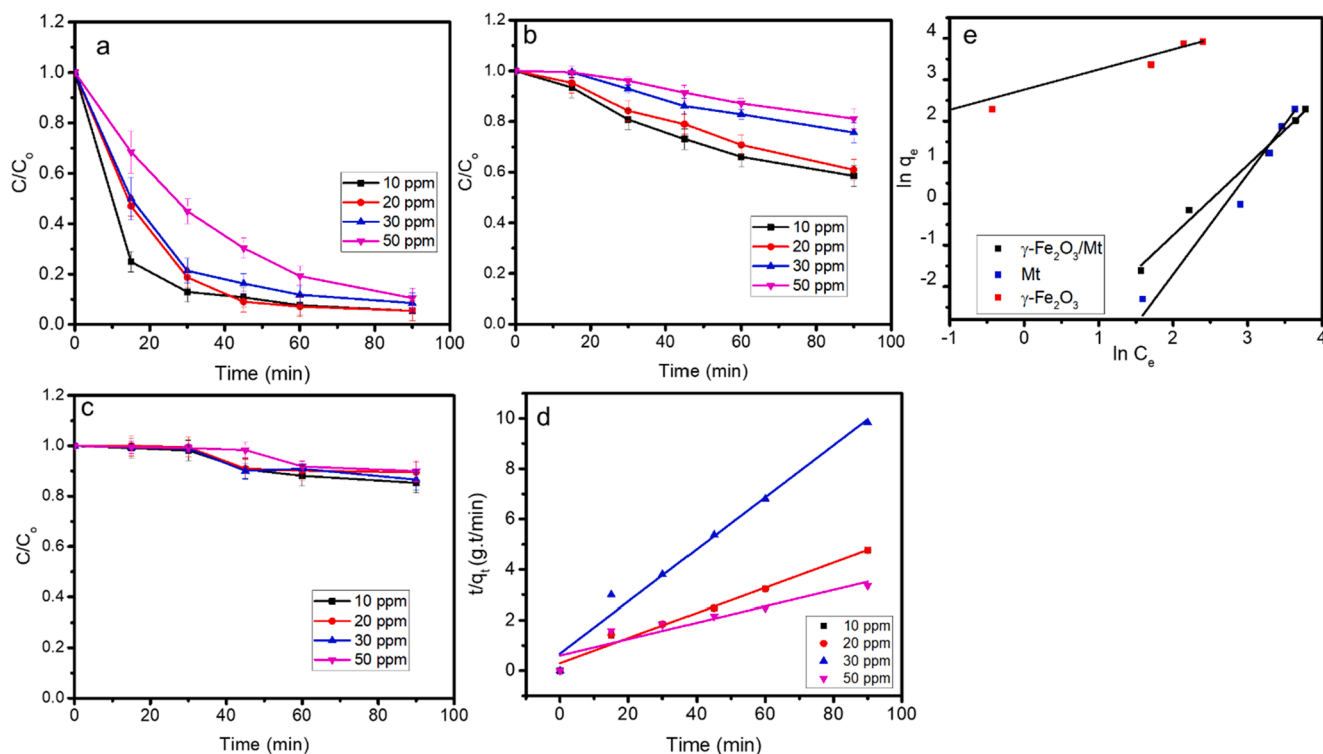


Fig. 7. (a). Kinetics of RhB adsorption using $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ (b). Kinetics of RhB adsorption using Mt (c). Kinetics of RhB adsorption using $\gamma\text{-Fe}_2\text{O}_3$ (d) Pseudo-second order plots of RhB adsorption using $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ (e). Freundlich isotherm plots of RhB adsorption using $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ and Mt.

Table 1
calculated parameters from kinetics studies.

Pseudo-first-order			
	Kinetics parameter		
	R ²	k	Q _e (mg/g)
γ-Fe ₂ O ₃	0.897	1.06 × 10 ⁻³	5.55
γ-Fe ₂ O ₃ /Mt	0.944	2.55 × 10 ⁻²	48.8
Mt	0.997	2.46 × 10 ⁻³	15.4
Pseudo-second-order			
	R ²	k	Q _e
γ-Fe ₂ O ₃	0.977	1.18 × 10 ⁻³	5.68
γ-Fe ₂ O ₃ /Mt	0.997	2.08 × 10 ⁻²	47.64
Mt	0.704	3.06 × 10 ⁻³	9.86
Intra-particle diffusion			
	R ²	k _i	C
γ-Fe ₂ O ₃	0.706	0.23	-0.48
γ-Fe ₂ O ₃ /Mt	0.991	3.00	-0.35
Mt	0.900	0.60	-1.11

Table 2
Calculated isotherm parameters.

Adsorbent	Freundlich Isotherm parameters			Langmuir Isotherm Parameters			
	K _F (L/g)	1/n	R ²	q _m (mg/g)	K _L (L/mg)	R _L	R ²
γ-Fe ₂ O ₃	0.65	0.67	0.977	7.89	4.1 × 10 ⁻³	0.971	0.974
γ-Fe ₂ O ₃ /Mt	1153.70	0.39	0.995	45.08	7.8 × 10 ⁻²	0.999	0.965
Mt	418.29	0.45	0.991	26.23	3.8 × 10 ⁻²	0.991	0.973

Fig. 6 shows the adsorption–desorption profile of γ-Fe₂O₃/Mt in comparison with Mt and the compared pore distributions. The plots demonstrate the increasing adsorption capability of Mt after modification by γ-Fe₂O₃ dispersion. The calculated parameters based on the adsorption/desorption profile provided a specific surface area of 89.9 m²/g in γ-Fe₂O₃/Mt, which is higher than that of Mt (79.9 m²/g) and γ-Fe₂O₃ (24.6 m²/g). The specific surface area of γ-Fe₂O₃ is similarly with was reported in other previous works [35,36]. In this case, Mt plays a role in supporting a specific surface area and porosity, which theoretically influences the adsorption mechanism. Similar phenomenon was also occurred in the dispersion of γ-Fe₂O₃ onto kaolinite [35]. The increasing surface area parameter is consistent with the profile in that the pore distribution also increased, mainly at the microporous structure. Based on the pore distribution pattern, which implied no change in dominant pores, the average diameters of the pores were not significantly different, 11.3 Å in Mt and 11.8 Å in γ-Fe₂O₃/Mt.

A similar pattern was reported in a previous work on the dispersion of Fe₃O₄ into Mt and the dispersion of SnO₂ nanoparticles [17,37]. The increasing specific surface area in the composite generally represents the homogeneous distribution of the dispersed materials.

3.2. Adsorption experiment

The adsorption experiments of RhB using γ-Fe₂O₃, γ-Fe₂O₃/Mt, and Mt were performed in a horizontal shaker at room temperature. The kinetics of RhB removal using γ-Fe₂O₃/Mt, Mt, and γ-Fe₂O₃ are presented in Fig. 7a–7c. The plots indicate the significant improvement in RhB removal by γ-Fe₂O₃/Mt compared with Mt and γ-Fe₂O₃. Basically, the increasing adsorption was related to the specific surface area enlarged by the composite formation. This was confirmed by the lower

adsorption rate and capacity over γ-Fe₂O₃. The data showed that the kinetics of the adsorption was evaluated in accordance with Lagergren's pseudo-first order, Ho and McKay's pseudo-second order, and Weber and Morris' intraparticle diffusion models based on Eqs. (2–4).

$$\ln(q_e - q_t) = \ln q_e - kt \quad (2)$$

$$\frac{t}{q_t} = \frac{1}{k_2 q_e^2} + \frac{t}{q_e} \quad (3)$$

$$q_t = k_i t^{0.5} + C \quad (4)$$

where q_t (mg/g) is the amount of adsorbed metal ions at time t , q_e (mg/g) is the adsorption capacity, k (min⁻¹) is the first-order rate constant, k_2 (g/mg min) is the second-order rate constant of adsorption (min⁻¹), and k_i (mg.min^{1/2}/g) and C are the kinetic constant and constant of the intra-particle diffusion model, respectively [38,39].

The R² values of the calculated parameters listed in Table 1 show that the pseudo-second-order kinetic model is the best fit for adsorption by γ-Fe₂O₃/Mt and γ-Fe₂O₃. Meanwhile, adsorption by Mt follows the pseudo-first-order kinetics. From the Q_e data from all adsorbents, it can also be seen that adsorption capacity is in following order: γ-Fe₂O₃/Mt > Mt > γ-Fe₂O₃, suggesting the provided surface active for adsorption by Mt as support. A synergistic effect of the increasing adsorption capacity is in consistence with the increasing specific surface area. The plots of pseudo-second-order from the adsorption using γ-Fe₂O₃/Mt at varied concentrations are presented in Fig. 7d. These data suggest that adsorption by γ-Fe₂O₃/Mt adsorbents is largely controlled by the chemisorption mechanism rather than mass transport. This is in contrast to the adsorption by Mt, for which the best fit is pseudo-first-order kinetics. Moreover, the adsorption capacity was enhanced either by the pseudo-first- or pseudo-second-order kinetic model. Based on these kinetic parameters, the chemical interaction caused by the γ-Fe₂O₃ dispersion in the Mt surface can be described.

The details of the adsorbate-adsorbent interaction were evaluated by determining the adsorption isotherm using the Freundlich and Langmuir isotherms. The equations used for the Freundlich and Langmuir models were as follows (Eqs. 4 and 5):

$$q_e = K_F C_e^{1/n} \quad (4)$$

$$q_e = \frac{q_m K_L C_e}{1 + K_L C_e} \quad (5)$$

where q_e and q_m (mg/g) are the adsorption capacities of the adsorbent at the equilibrium and maximum values, respectively; C_e (mg/L) is the concentration of the adsorbate in equilibrium; K_F and n are the Freundlich constants related to the adsorption–desorption equilibrium, K_L (L/mg) is the Langmuir constant related to the energy of adsorption and adsorption intensity [40,41].

The R_L parameter from the Langmuir isotherm is represented as follows (Eq. 6):

$$R_L = \frac{1}{1 + K_L q_m} \quad (6)$$

Table 2 lists the calculated isotherm parameters, and the fittest plot is presented in Fig. 7e.

The R² values suggest that the Freundlich isotherm model is more feasible than the Langmuir isotherm for adsorption by all tested materials. However, for RhB adsorption by γ-Fe₂O₃/Mt, the R² value was higher than that for the Langmuir isotherm. By contrast, adsorption by Mt showed a better fit for the Langmuir isotherm. The K_F and K_L values showed significant improvements in adsorption capability. Both constants from γ-Fe₂O₃/Mt were twice the values from Mt, while the

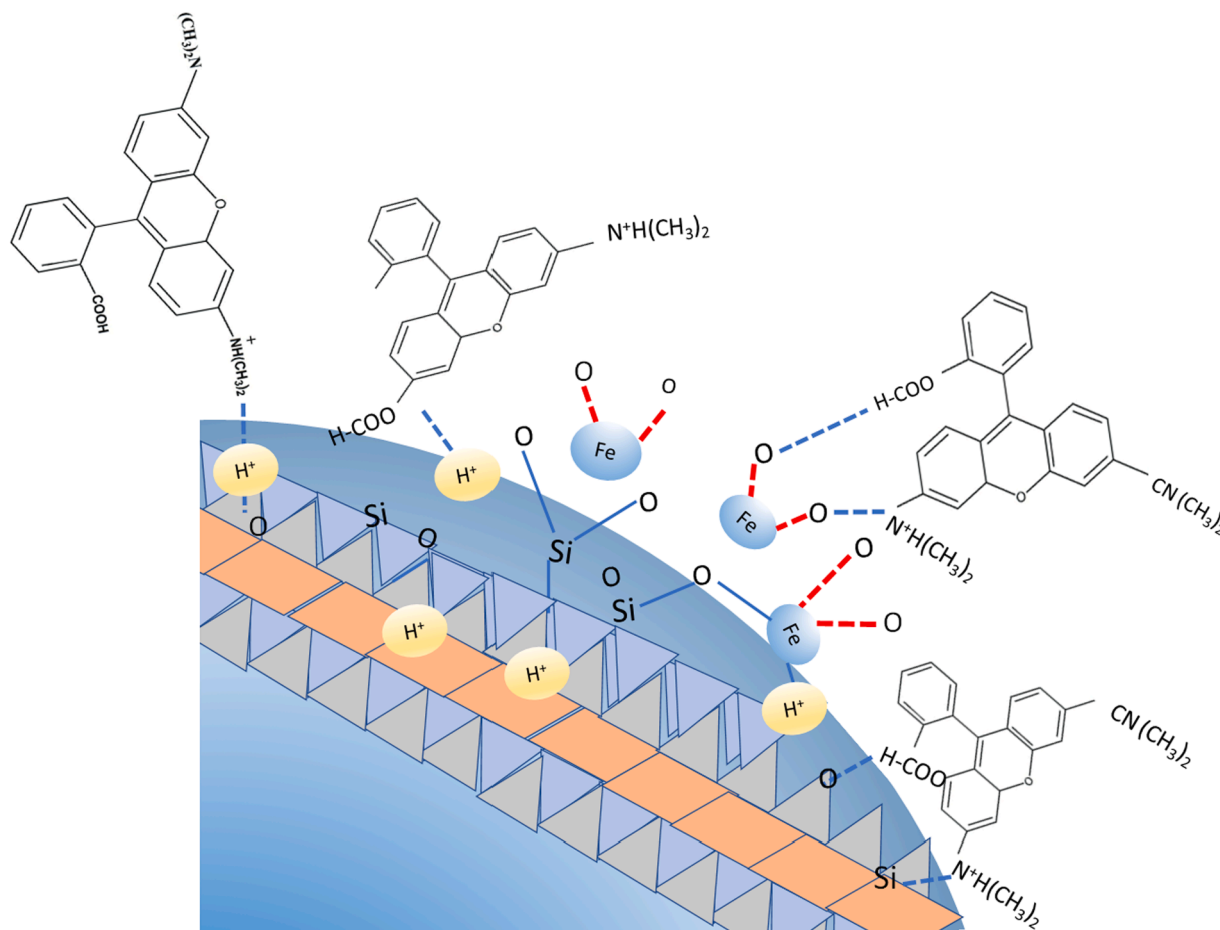


Fig. 8. Schematic diagram of RhB adsorption mechanism by $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$.

Brunauer-Emmett-Teller (BET) surface area of $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ was not higher than twice the BET surface area value. This can be ascribed to the role of dispersed nanoparticles in adsorption-active sites [42]. The enhanced hydrophilicity, along with the presence of abundant negative functional surface groups ($-\text{O}$ and $-\text{OH}$), is the most possible explanation for the rapid adsorption of RhB. This assumption is justified by the low zeta potential of $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ (pH 5.8) compared with that of Mt (pH 6.8). A schematic representation of the mechanism can be seen in Fig. 8.

3.3. Photocatalytic activity study

The photocatalytic activity of $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ is presented by the kinetics of RhB removal at different initial concentrations (Fig. 9a). For each experiment, a mixture of dye solution and photocatalyst powder was first kept in the dark for 15 min with constant stirring. The stirring in the dark was aimed to get adsorption equilibrium prior photocatalysis. The plot shows that the degradation efficiency reached greater than 96% at 30 min and 99% at 60 min at all concentrations. The comparison of DE values by photocatalysis over $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ and $\gamma\text{-Fe}_2\text{O}_3$ presented in Fig. 9b suggest the similar DE at low RhB concentration (2–10 ppm), but at increasing concentration $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ was superior respect to $\gamma\text{-Fe}_2\text{O}_3$. The higher degradation is attributed to the contributing adsorption mechanism which is more dominant in $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$. The contribution adsorption in photocatalytic mechanism was also reported in the use of

BiOBr, $\text{Fe}_3\text{O}_4@\text{MoS}_2$, and Fe(III)/montmorillonite [43–45]. The DE values indicated that the $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ had high efficiency, as also demonstrated in the comparison with the DE values obtained from other similar works listed in Table 3. The DE value obtained in this work was higher than that of $\gamma\text{-Fe}_2\text{O}_3/\text{SiO}_2$, which was 95% at the same duration of 30 min, and that of Ag–AgI/ $\text{Fe}_3\text{O}_4@\text{SiO}_2$, which was 95% at 1 h of treatment [46–51].

3.4. Degradation mechanism

The identification of the degradation mechanism in photocatalysis in comparison with the adsorption mechanism was performed using UV–visible spectrophotometry and liquid chromatography coupled to mass spectrometry (LCMS) analysis. The UV–visible spectra in Fig. 9b and 9c represent a significant difference among the spectral changes obtained by adsorption and the photocatalysis mechanism. Adsorption produced reduced absorbance spectra along with prolonged treatment time. From the photocatalysis treatment, not only reduced absorbance but also the red shift of the characteristic peak of 556 nm was found. The red shift is characteristic of deethylation and decarboxylation. This is also confirmed by the LCMS analysis presented in Fig. 10. The LC analysis of the treated solution revealed some other peaks as proof of the degradation products (Fig. 10a). Besides the other peaks found in the treated solution from the LC analysis (Fig. 10b), the treated solutions exhibited fractions with m/z values of 359, 331, 181, 168, 146, and 128

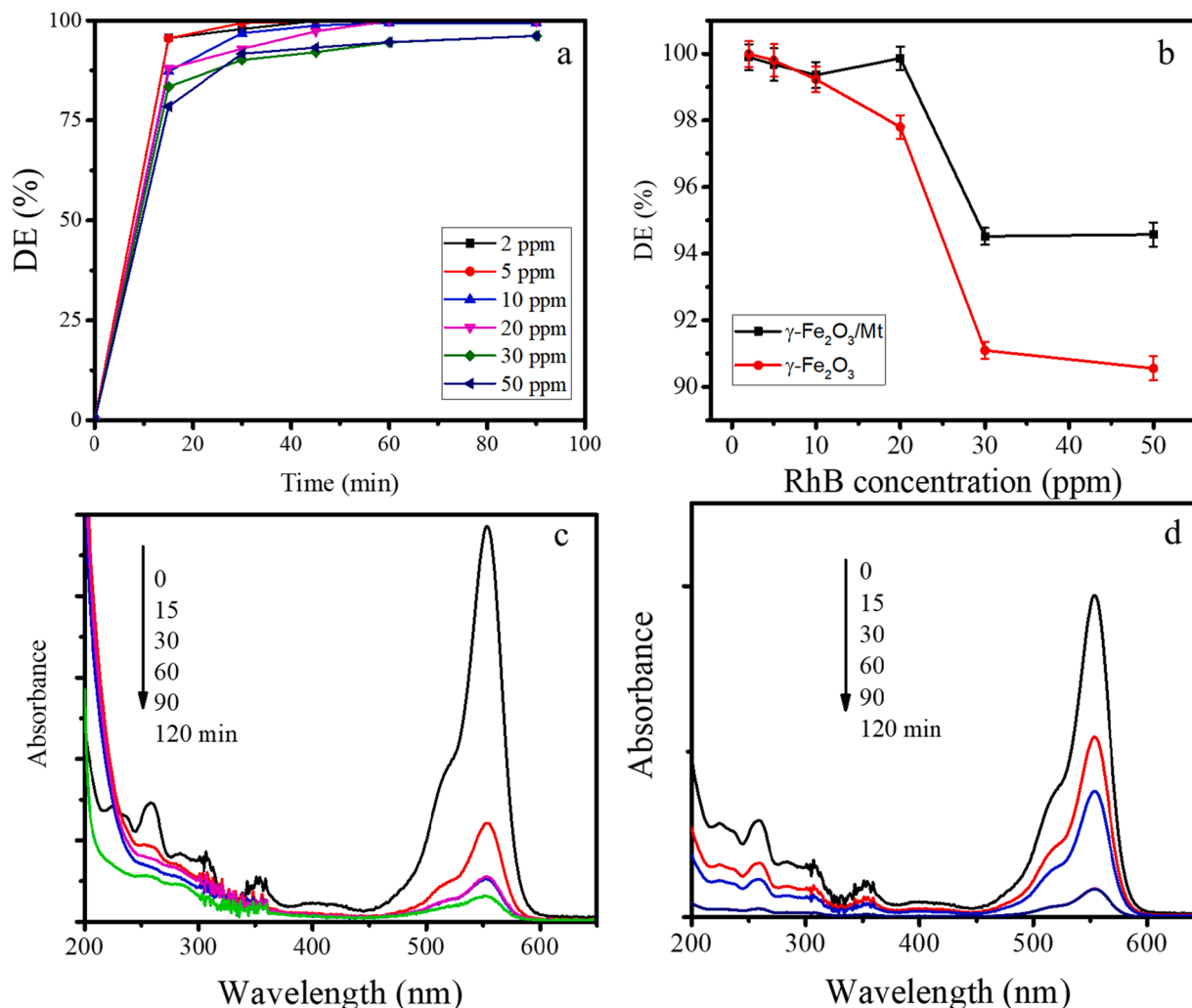


Fig. 9. (a) DE of photocatalytic degradation using $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ at varied concentration (b) Comparison on DE by photocatalytic degradation using $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ and $\gamma\text{-Fe}_2\text{O}_3$ (c) UV-Visible spectral change by photocatalytic mechanism (d) UV-Visible spectral change by adsorption mechanism.

Table 3
RhB removal by various materials.

Material	DE (%)	Remark	Reference
$\alpha\text{Fe}_2\text{O}_3$ NPs	73 at 1 h	Particle size of $\alpha\text{Fe}_2\text{O}_3$ is ranging at 48–55 nm with the band gap energy of 2.14 eV	[39]
$\gamma\text{-Fe}_2\text{O}_3/\text{SiO}_2$	95 at 30 min	The DE achieved by the photocatalyst dose of 0.8 g/L, silica was derived from rice husk ash	[40]
$\text{BaTiO}_3/\text{MnO}_2$	70 at 3 h	photodegradation of RhB in the presence of MnO_2 -modified BTO ceramics followed a pseudo-first order model a	[41]
$\text{Ag-AgI}/\text{Fe}_3\text{O}_4/\text{SiO}_2$	95 at 1 h	The light source was a 250 W lamp, photocatalyst dosage of 0.1 g/L	[42]
$\text{Cu-Fe}_3\text{O}_4$ magnetic nanoparticles	92.8 at 1 h	The optimum degradation was achieved at pH \sim 7 and 25 °C and the rate constant of about 0.0256 min^{-1} .	[43]
NiMn_2O_4 nanoparticles	98 at 80 min	The photocatalyst has the band gap energy of 3.0 eV	[44]

as degradation products. These spectra elucidate that the possible degradation mechanism can be described using the following scheme (Fig. 11):

These analyses are similar to the identification of RhB degradation over other TiO_2 -supported activated carbons and SnO_2/Mt [17].

To further determine how reactive species induce the degradation mechanism, hole and hydroxy radical scavengers were added to the photocatalytic system. To this, isopropanol (IPR) was added as the hydroxy radical scavenger, while the EDTA was added as a hole scavenger. The effect of the addition to the degradation rate is depicted in the kinetic plots and the calculated initial rates presented in Fig. 12a and Fig. 12b show that the photodegradation rate was increased with the addition of EDTA and decreased with the addition of IPR. In order to reduce the interpretation on decreasing concentration caused by adsorption pathway, the test on scavenger addition under light illumination was performed by previously treated by adsorption under dark condition. The proof that the degradation mechanism occurred by varied treatment, HPLC analysis was performed with the chromatogram presented in Fig. 13. The chromatogram of initial RhB shows the single dominant peak at 6.8 min as identity of RhB molecule. After was treated

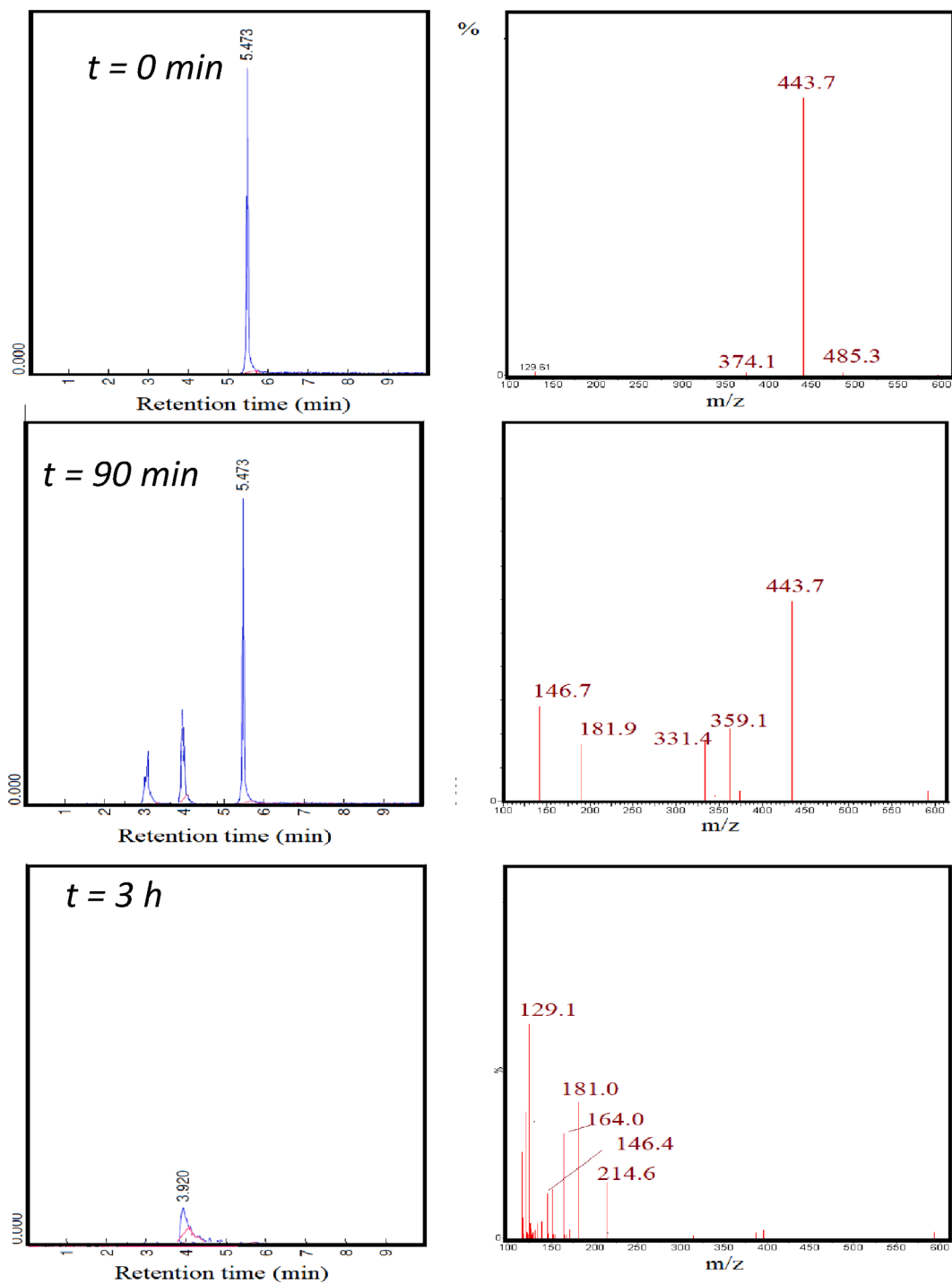


Fig. 10. LCMS profile of initial and treated solutions.

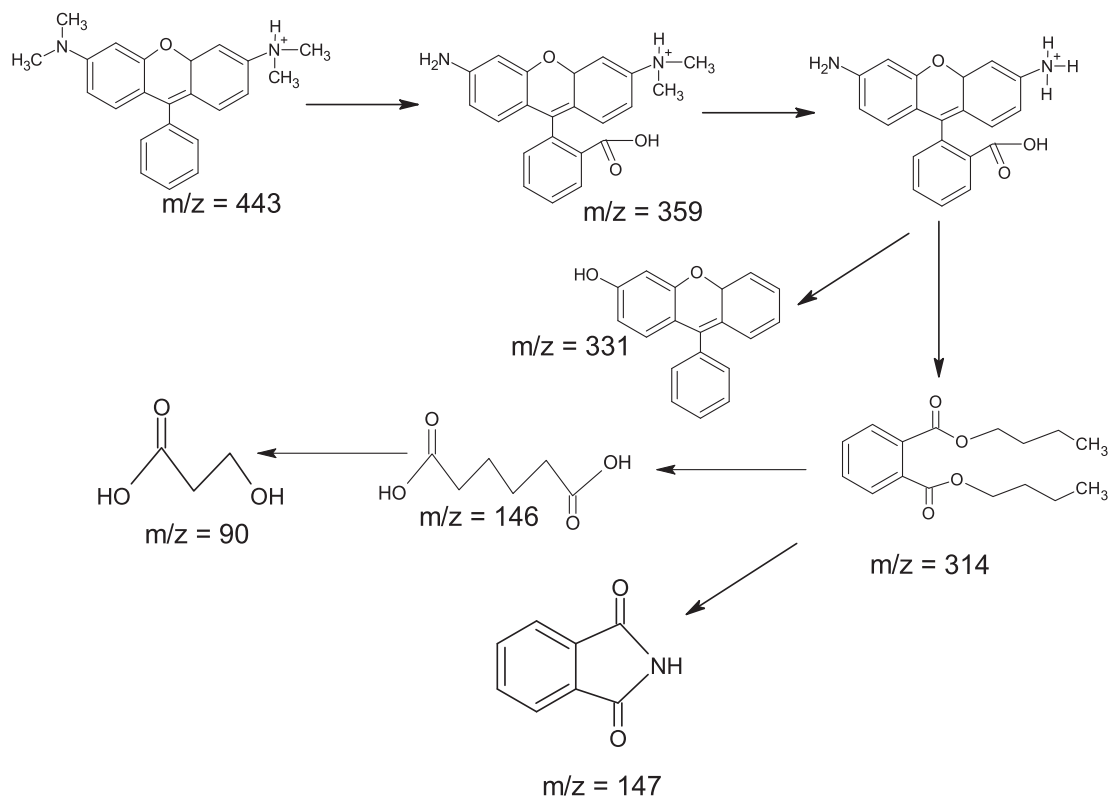


Fig. 11. Proposed mechanism of RhB degradation by photooxidation using $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$.

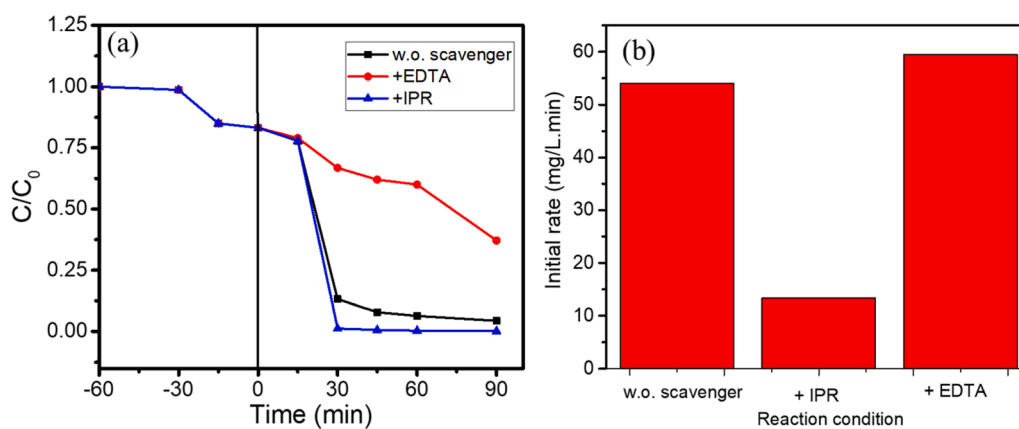


Fig. 12. (a). DE RhB (b) Initial rate of degradation using $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ with and without scavenger.

by varied condition, the chromatograms exhibit other peaks at less retention time, associated with smallest molecular weight. It suggest that there was degradation of RhB structure as shown by other peaks in all varied treated solutions for 30 min. The slower degradation rate by IPR addition was confirmed by the higher concentration of rest RhB in the solution. Contrarily, the faster degradation by EDTA addition was expressed by less intense RhB peak, along with some more intensive peaks of degradation products. This phenomenon revealed the role of the promoted separation of electron-hole pairs related directly to the increased lifetime of electron-hole pairs. In addition, the trapping hydroxy radicals by IPR inhibited the propagation reaction owing to their consumption in further oxidation process toward RhB. These results imply that the radicals were the predominant reactive species for the photocatalytic degradation of RhB. A similar effect was reported for the photocatalytic activity of Fe_2O_3 nanoparticles [52] and BiFeO_3 nanostructures [53].

3.5. Reusability study

Examination of reusability is an important evaluation for a photocatalyst to be applied on an industrial scale. The experiments were conducted in five cycles based on the initial rate and DE achieved at 30 min. The photocatalyst was recycled by filtering, washing using n-hexane, and calcined for 200 °C after the completion of each cycle. The easily separable property of the material is related to the magnetism feature measured using the VSM analysis presented in Fig. 14a. The magnetism plot shows that the $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ demonstrates specific magnetic saturation magnitudes of 22.7 emu/g. This value is less than the magnetism of $\gamma\text{-Fe}_2\text{O}_3$ NPs prepared from various experimental methods, which ranged from 60 to 70 emu/g, representing the influence of the non-magnetic properties of clay support. However, magnetism is still applicable for magnetic attraction for easy separation with high recoverability. The bar chart depicted in Fig. 14b shows that the

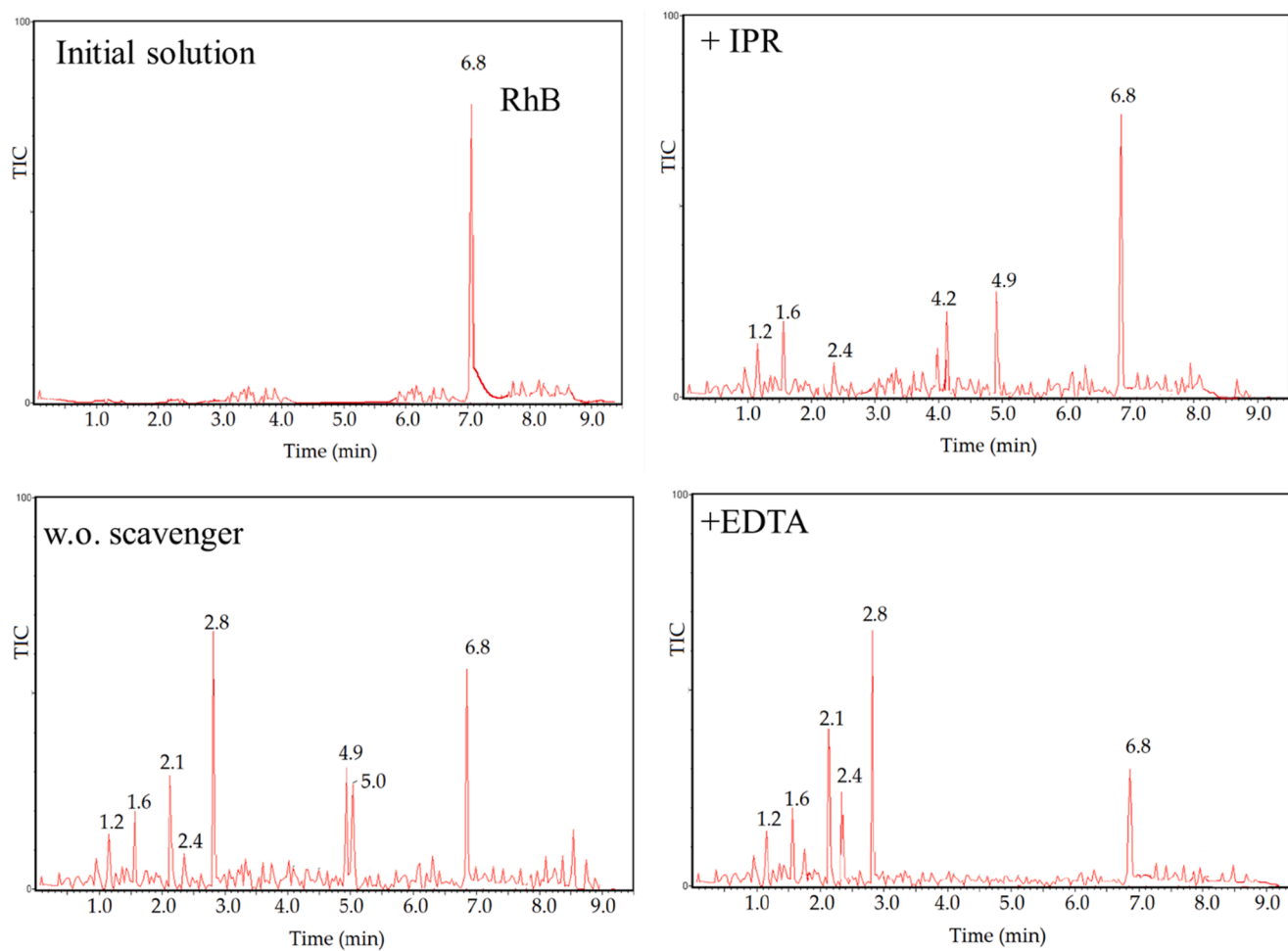


Fig. 13. Chromatograms of initial RhB solution and varied treated solutions for 30 min.

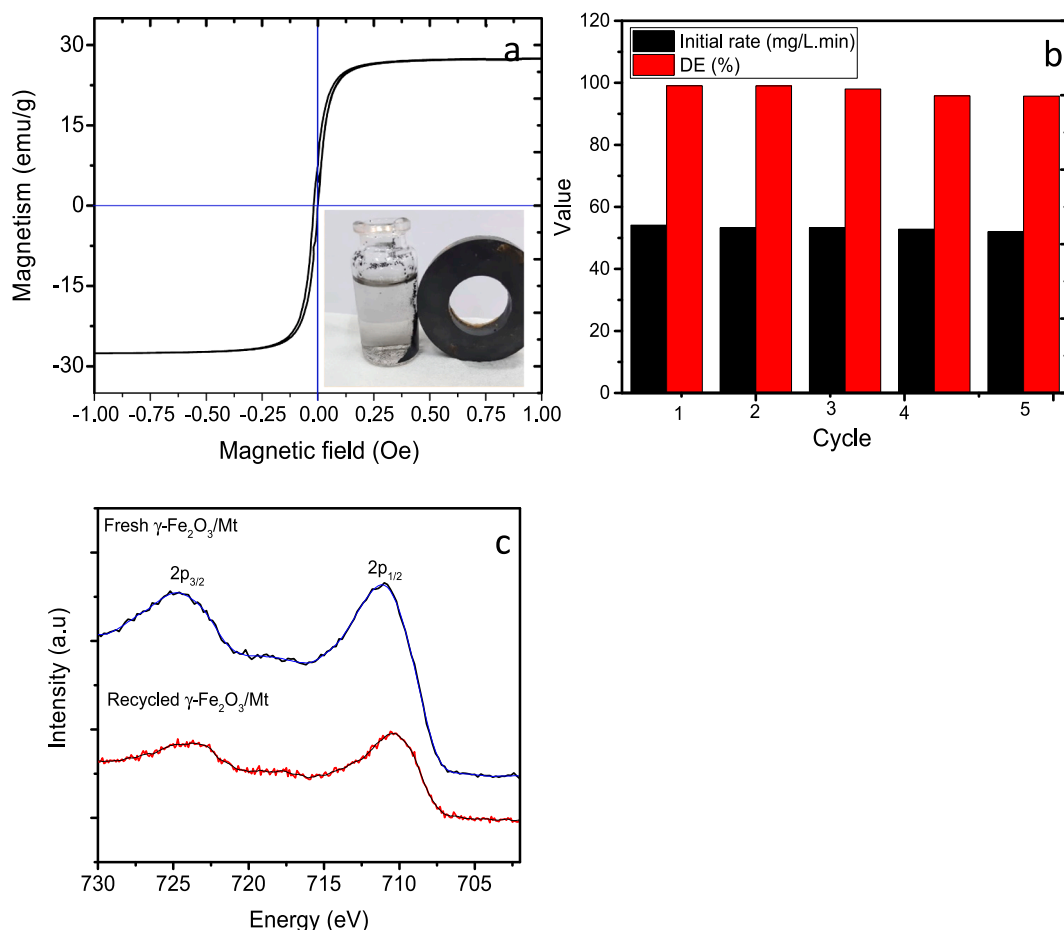


Fig. 14. (a). Magnetism curve of $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ (b) Initial rate and DE RhB degradation using $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ at various cycles (c). Comparison on XPS Fe 2p spectrum fresh and recycled $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$.

photocatalytic activity achieved was quite stable, as confirmed by the maintained DE at approximately 97–99% and the reduction of the initial rate to $\leq 10\%$. In addition, the stability of the photocatalyst was also monitored by XRD measurement and XPS of the reused photocatalyst. The spectra of Fe 2p presented in Fig. 14 c suggest the insignificant change of the material. No other reflection was identified in the recycled photocatalyst, and no other peaks in the XPS profile were observed, confirming that the single-ion species of Fe^{3+} were maintained.

Finally, the $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ nanocomposite exhibited excellent properties in terms of visible-light activity, easy magnetic separation, stability, and reusability, making it practically applicable in dye photo-oxidation. Mt is an abundant, low-cost, naturally occurring material that will contribute to the economic value of nanocomposites to be further developed.

4. Conclusion

A nanocomposite of $\gamma\text{-Fe}_2\text{O}_3/\text{Mt}$ was successfully prepared using a hydrothermal method. The nanocomposite demonstrated excellent features as an adsorbent and photocatalyst for RhB removal. The homogeneous dispersion of the single-phase $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles in the Mt structure contributed to the enhanced dye-surface interaction, as shown by the significant improvement in adsorption capability. The narrow band-gap energy of 3.2 eV is fit visible light and attractivity to magnetic force provide the benefits of a low-cost and easy handling photocatalytic system. In addition, the nanocomposite showed a magnetically separable performance for complete recovery after use. The stability of the nanocomposite, proven by the relatively unchanged phase after use for many cycles, greatly promotes its potential practical application for the

removal of dye pollutants from wastewater.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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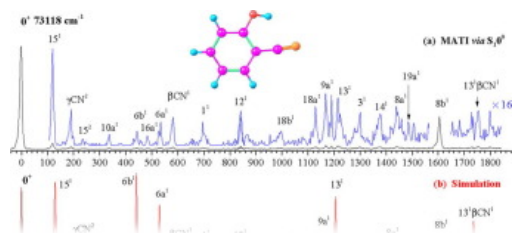
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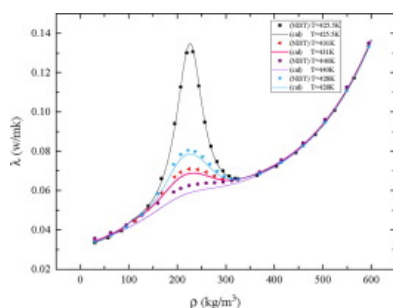
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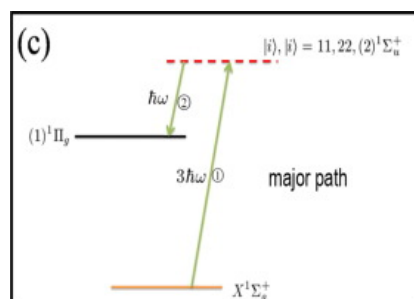
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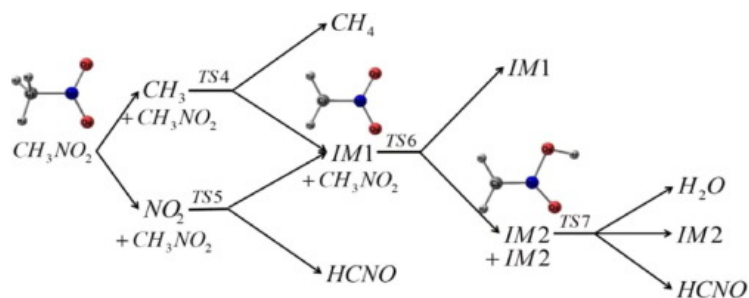
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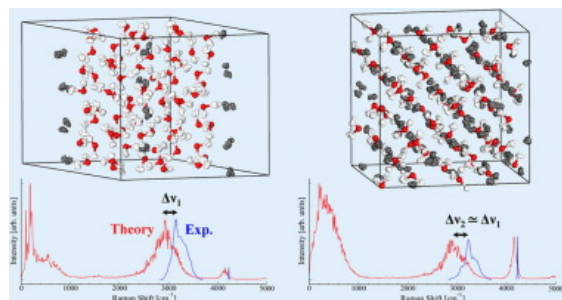
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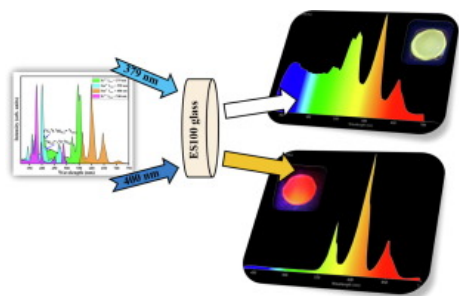
Excitation dependent luminescence of Er³⁺/Sm³⁺ co-activated multicomponent borosilicate glasses via energy transfer analysis

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Nanostructures and Materials

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Ecofriendly synthesis and white light-emitting properties of $\text{BaLa}_2\text{ZnO}_5:\text{Dy}^{3+}$ nanomaterials for lighting application in NUV-WLEDs and solar cells

Priyanka Sehrawat, R.K. Malik, R. Punia, Sanjeev Maken, Neelam Kumari

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Abstract

Abstract

Herein, we report a new set of white-emitters i.e., $\text{BaLa}_2\text{ZnO}_5:\text{Dy}^{3+}$ nanomaterials, fabricated via an inexpensive and energy-saving urea-aided-combustion route. Tetragonal phase-transparency had been authorized via X-ray powder diffraction-facilitated-Rietveld refinement analysis. Under 355 nm energization, target samples manifested white-luminance via two bands, namely bluish and golden (485 & 578 nm). Fine photometric characteristics of optimal nanopowder had been reflected in white coordinates (0.2693, 0.3278) with high purity of white-light (27.61×10^{-2}), quantum-efficiency (74.95%), and appropriate color-temperature (9290 K). Briefly, the aforementioned results suggested their promising employment for generating cool-white-light in high-performance cool-wLEDs (white-light emanating diodes), lasers, indoor-cultivation, sensors, and photovoltaic-cells.

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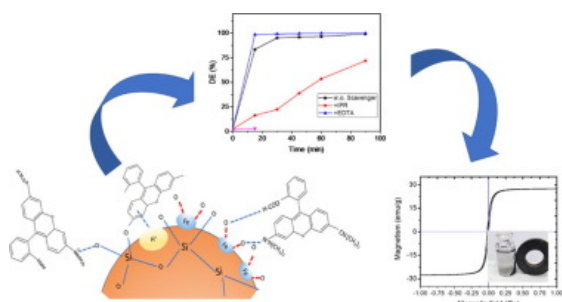
Is Fatimah, Gani Purwiandono, Arif Hidayat, Suresh Sagadevan, Azlan Kamari

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Exploring the mechanism of CdTe quantum dots as fluorescent probe to detect Hg(II) ion from the perspectives of fluorescence polarization and light scattering

Hongyu Zhou, Congling Yang, Mengxia Liao, Minglian Li, ... Jiayang Wang

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The mechanism of quantum dots (QDs) sensing metal ions remains controversial. Herein, we investigate the mechanism from the perspectives of fluorescence anisotropy and light scattering, where 3-mercaptopropionic acid (MPA) capped CdTe QDs are employed as a model fluorescent sensor and Hg^{2+} is used as a model quencher. We observe that when with the QDs is mixed with the Hg^{2+} , the fluorescence is quenched while the light scattering is enhanced and the fluorescence anisotropy is increased. These observations indicate that the Hg^{2+} destabilizes the QDs and drives them to agglomerate. Since the PL mechanism of QDs is closely depended on the quantum size effect, it implies that the agglomeration of QDs might be an important contributor to the fluorescence quenching. This work provides novel insights to explain the mechanism of QDs as fluorescent probe to detect metal ions.

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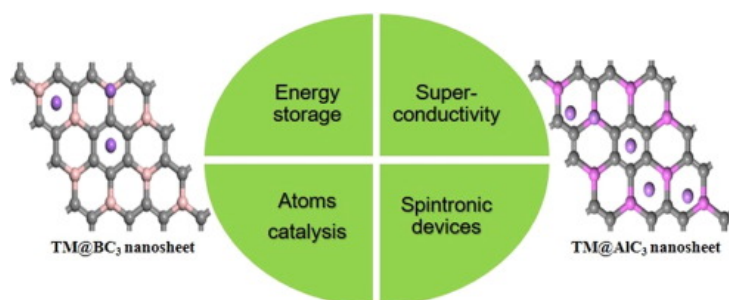
Jiawen Wang, Yingke Yang, Haili Liu, Huilong Dong, ... Youyong Li

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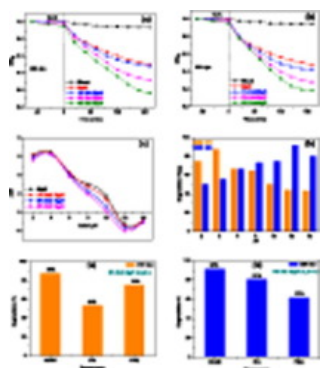
R. Vijaya Shanthi, R. Kayalvizhi, M. John Abel, K. Neyvasagam

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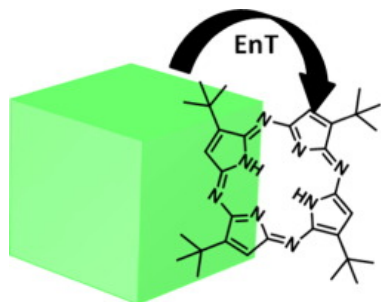
Arunkumar Kathiravan

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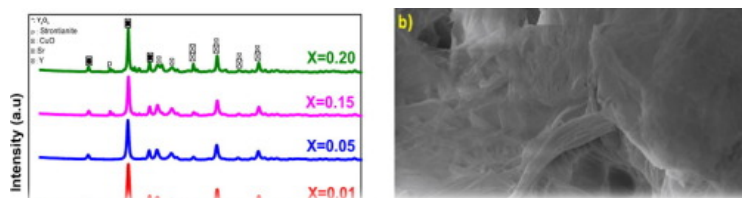
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Simple and fast detection of homocysteine by cucurbit[7]uril fluorescent probe based on competitive strategy

Chenrui Jiang, Deli Xiao, Ping Yang, Wenyao Tao, ... Hua He

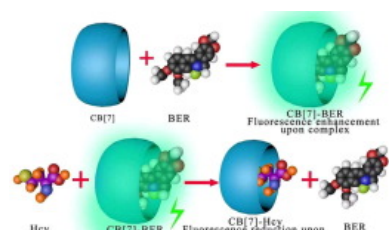
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The host–guest fluorescent probe based on competition mode was established to detect Hcy. CB[7] and BER can self-assemble to form a strong fluorescent host–guest inclusion complex. Due to the concentration advantage, the added analyte Hcy can compete to expel part of the BER molecules in the hydrophobic cavity of CB[7] macrocyclic host, resulting in a decrease in fluorescence.



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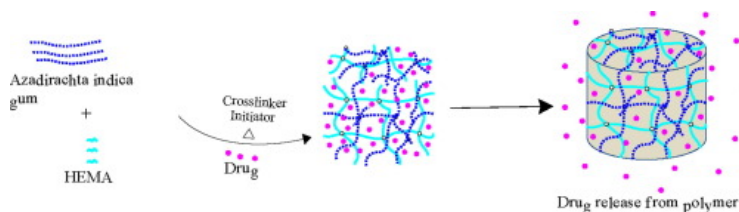
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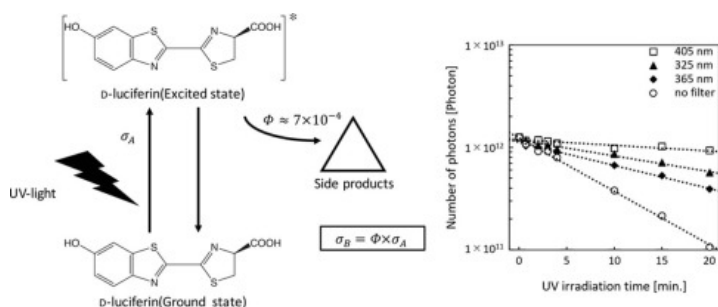
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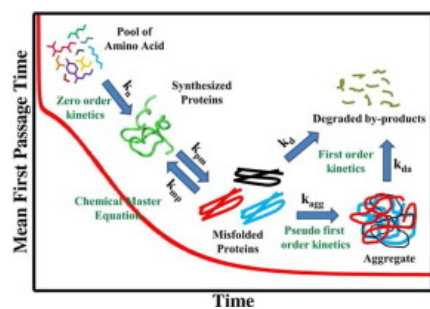
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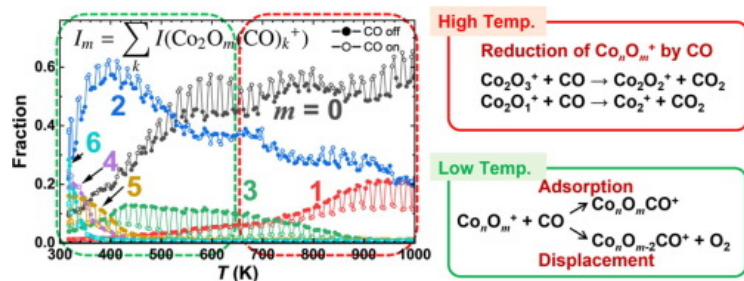
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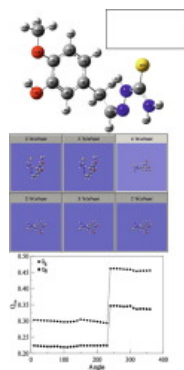
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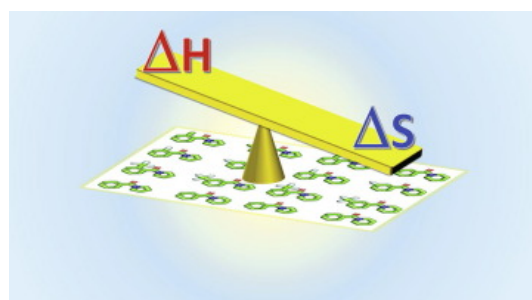
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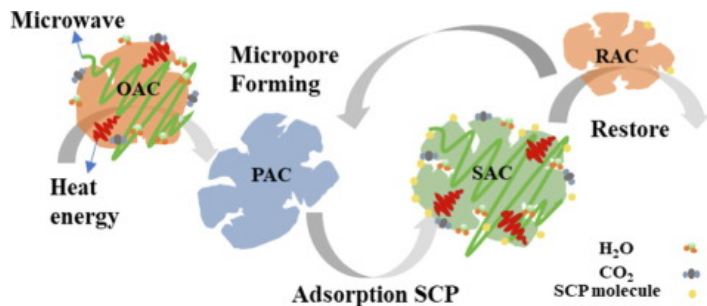
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