Zinc-bearing dust derived non-toxic mixed iron oxides as magnetically recyclable photo-Fenton catalyst for degradation of dye
Jian-ming Gao, ShuJia Ma, Zongyuan Du, Fangqin Cheng and Peng Li

ABSTRACT
In this paper, comprehensive utilization of hazardous zinc-bearing dust for preparation of non-toxic mixed iron oxides as a magnetically recyclable photo-Fenton catalyst for degradation of dye by a facile solid state reaction process was proposed. The as-prepared samples were characterized by X-ray diffraction (XRD), Raman spectra, ultraviolet and visible (UV-Vis) spectra and Physical Property Measurement System (PPMS), and the degradation performance of as-prepared catalysts was also tested and analyzed. The results show that spinel ferrite coexisting with or without Fe₂O₃ was the predominant phase in the as-prepared samples, which were confirmed by Raman analysis. The as-prepared samples presented high degradation efficiency (about 90%) of rhodamine B (RhB) in the presence of hydrogen peroxide (H₂O₂) with visible light irradiation, owing to the synergistic effect of photocatalyst reaction and Fenton-like catalyst reaction during the degradation process. The mixed iron oxides also presented stable structure and exhibited excellent reusability with a degradation efficiency of 87% after the fifth cycle of reuse. Importantly, the heavy metals in the zinc-bearing dust could be fixed in the stable spinel structure. This paper could provide a simple approach for comprehensive utilization of zinc-bearing dust to synthesize non-toxic mixed iron oxides as a magnetically recyclable photo-Fenton catalyst for degradation of dye.

Key words | dye degradation, magnetically recyclable, mixed iron oxide, photo-Fenton catalyst, zinc-bearing dust

HIGHLIGHTS
● Non-toxic mixed iron oxides were prepared from hazardous zinc-bearing dust.
● Mixed iron oxides could be used as a magnetically recyclable photo-Fenton catalyst.
● The catalyst exhibited degradation efficiency of 85% after the fifth cycle of reuse.
● Heavy metals in the zinc-bearing dust could be fixed in the stable spinel structure.

INTRODUCTION
The contamination caused by organic dyes has been of great concern due to the toxicity to humans, and various physical, biological and chemical methods including adsorption (Jiang et al. 2019; Zhou et al. 2019), membrane separation (Ramlow et al. 2017), and oxidation (Liu et al. 2021) have been developed for the treatment of dye wastewater. Among these methods, advanced oxidation processes (AOPs) based on the technology of Fenton reaction are potentially applied in industrial wastewater treatment to degrade recalcitrant organic pollutants, which are cost effective and environmentally friendly (Neyens & Baeyens 2003; Zapata et al. 2009; Gligorovski et al. 2015). The technology is based on the generation of powerful hydroxyl radicals, -OH, which can oxidize and mineralize organic pollutants in wastewater. The generation process involves simple
Fe$_3$Fe$_{78}$Si$_9$B$_{13}$ alloy etc. have been synthesized and catalysts, such as nano-flake Fe-SC hybrid, Cu-impregnated zeolite Y, sulfur-modified iron oxide, and amorphous Fe$_{78}$Si$_9$B$_{13}$ alloy etc. have been synthesized and confirmed the Fenton-like reaction on the solid-liquid interface (Du et al. 2016; Jia et al. 2016; Kong et al. 2016).

Mixed iron oxides, spinel ferrites MFe$_2$O$_4$ (M = Ni, Mn, Co, Mg, Cu, Zn, etc.), have been applied as heterogeneous catalysts in the Fenton-like processes, which exhibit unique structural, magnetic properties and catalytic performance (Ahmed & Ahmaruzzaman 2015; Kefeni et al. 2017; Vinosha et al. 2017). Liu et al. (2012) successfully synthesized magnetic NiFe$_2$O$_4$ by the hydrothermal method, and as-prepared nickel ferrite exhibited photo-Fenton catalytic features for organic pollutants in the presence of oxalic acid. Seven cyclic tests for rhodamine B degradation demonstrated that the catalyst is stable and highly active with a degradation efficiency of above 90%. Highly ordered mesoporous CuFe$_2$O$_4$ was successfully obtained and proposed as a heterogeneous Fenton-like catalyst (Zhang et al. 2014). The CuFe$_2$O$_4$ catalyst presented low metal leaching (<1 ppm) even in acidic conditions and retained high catalytic activity (3.75% decrease) after six cycles of reuse. Besides, many studies demonstrated that spinel ferrites presented excellent catalytic activity in Fenton-like reaction systems, such as ZnFe$_2$O$_4$, CoFe$_2$O$_4$, MgFe$_2$O$_4$, and Zn$_x$Co$_{1-x}$Fe$_2$O$_4$ (Su et al. 2012; Feng et al. 2013; Shahid et al. 2013; Cai et al. 2016). Moreover, it is confirmed that for some metal elements, such as Cr, Mn, Ti or V, substitution into the spinel structure can be facilitated to improve the catalytic activity of the spinel ferrite-based Fenton-like catalysts (Zhong et al. 2014). Furthermore, they are all inverse spinels with excellent stability, and toxic ions could be stably fixed in spinel ferrites (Chen et al. 2011; Tu et al. 2012), demonstrating that multi-metal doped spinel ferrites could be used as magnetically recyclable Fenton-like catalysts for organic pollutant degradation.

Zinc-bearing dust, generated during the iron and steel production process, is classified as hazardous solid waste that contains various heavy metals, such as Zn, Mn, Fe, Pb, Cr etc. Currently, much effort was focused on the extraction and recycling of valuable metals from zinc-bearing dust using various complex pyrometallurgical and/or hydrometallurgical processes, leading to not only environmental pollution problems but also low recovery of valuable metals (Kukurugya et al. 2015; Lin et al. 2017; Zhang et al. 2017a). Unfortunately, few works have studied comprehensive utilization of valuable metals in zinc-bearing dust. Notably, Zn, Mn, Fe, Cr etc. in the zinc-bearing dust are the main chemical compositions of spinel ferrites (Gao & Cheng 2018a, 2018b), suggesting that zinc-bearing dust could be used as raw material for preparation of spinel ferrites as magnetically recyclable heterogeneous Fenton-like catalysts. Besides, the gap vacancy and lattice defects in spinel ferrites could provide structural conditions for metal substitution into the spinel structure, implying that it is reasonable to synthesize multi-metal doped spinel ferrites from zinc-bearing dust. Furthermore, heavy metal ions easily dissolved in water from zinc-bearing dust could be stably fixed in the stable spinel structure, which might be facilitated to improve the catalytic activity of the spinel ferrite-based Fenton-like catalysts. In short, comprehensive utilization of hazardous zinc-bearing dust for preparation of non-toxic mixed iron oxides as magnetically recyclable photo-Fenton catalyst for degradation of dye meets the requirements of a resource-saving and environment-friendly society.

So far, some mixed iron oxides as heterogeneous Fenton-like catalysts have been successfully synthesized from solid waste or natural minerals. Cao et al. (2017) reported that zinc ferrite catalysts for efficient degradation of organic dye were fabricated by the calcination of electroplating sludge, and the as-prepared catalysts exhibited excellent MB decolorization efficiency (about 85%) in a UV/H$_2$O$_2$ system. Zhang et al. (2017b) provided a novel method for the reuse of iron-containing Fenton sludge and nickel ferrite was synthesized as an efficient catalyst in the heterogeneous Fenton process. A phenol degradation efficiency of about 95% could be obtained in the presence of both nickel ferrite and H$_2$O$_2$. Furthermore, multi-metal co-doped magnesium ferrite was successfully synthesized from saprolite laterite ore by Diao et al. (2017), and used as heterogeneous photo-Fenton like catalyst for dye degradation. The as-prepared ferrite exhibited excellent catalytic activity with a degradation efficiency of 96.8%, which is more competitive compared to the catalysts prepared from chemical reagents. That is to say, it is feasible to obtain excellent heterogeneous Fenton-like catalysts from solid waste or liquid waste.
In this paper, multi-metal doped spinel ferrites were synthesized from zinc-bearing dust by the facile solid state reaction method. The structure and magnetic properties of as-prepared mixed iron oxides were characterized and discussed. Then the as-prepared ferrites were used for the degradation of organic pollutants by photo-Fenton reaction, and the effects of degradation conditions and Zn/Fe molar ratio on the degradation efficiency of rhodamine B (RhB) were investigated. Simultaneously, the dissolution characteristics of heavy metals in the zinc-bearing dust and as-prepared spinel ferrites were also tested to verify the environmental security of zinc-bearing dust-derived mixed iron oxide photo-Fenton catalysts. Finally, the possible catalytic mechanism for degradation of RhB using zinc-bearing dust-derived mixed iron oxides as photo-Fenton catalysts was proposed. This paper might provide a simple approach for comprehensive utilization of zinc-bearing dust in a green pollution-free method.

**EXPERIMENTAL PROCEDURE**

**Materials and reagents**

The zinc-bearing dust used in this study was collected from a stainless steel plant in China, and the main chemical compositions of the zinc-bearing dust are summarized in Table 1. In the zinc-bearing dust, spinel ferrites including ZnFe$_2$O$_4$ and Fe$_3$O$_4$, calcite (CaCO$_3$) and quartz (SiO$_2$) are the main phases as reported in the literature (Gao & Cheng 2011a). All the reagents used in this study are of analytical grade and used without any treatment.

**Sample preparation**

Using the facile solid state reaction method, the mixed iron oxides were prepared by the following procedure. Firstly, zinc-bearing dust was dried at 105°C for 12 h and ground to a particle size smaller than 75 μm. Then, 5.0 g zinc-bearing dust was mixed with a certain amount of ZnO reagent, and the mixture was ground adequately in an agate mortar to form a uniform powder. Finally, the uniform powder was transferred into a corundum crucible, and calcinated at 900°C for 60 min with a heating rate of 10°C/min in a muffle furnace to obtain mixed iron oxide. In this paper, to investigate the effect of Zn/Fe molar ratio on zinc ferrite (ZnFe$_2$O$_4$) based photo-Fenton catalysts synthesized from zinc-bearing dust, the mass ratios of 2:0.4, 2:0.6 and 2:0.8 were chose, and the molar ratio of Zn to Fe in the dust and the mixture of zinc-bearing dust and ZnO addition are calculated and listed in Table 1. The as-prepared mixed iron oxides with mass ratios of 2:0.4, 2:0.6 and 2:0.8 were labeled as ZF1, ZF2 and ZF3, respectively.

**Degradation experiments**

The as-prepared mixed iron oxides were used as a Fenton-like catalyst for the degradation of RhB at a constant temperature of 25°C. A 500 W Xe lamp with 420 nm cutoff filter was used as the visible light source. In the degradation experiment, 0.2 g as-prepared mixed iron oxides was first dispersed in 200 mL of the RhB aqueous solution (10 mg/L) in a dark environment for 30 min to ensure that the RhB reached an adsorption equilibrium on the surface of the photocatalysts at a pH value of 4.0. Then a solution containing H$_2$O$_2$ (2.0 mmol/L) was added in the aqueous solution, and reacted with magnetic stirring. During the irradiation process, 5 mL of the suspension was collected every 30 min and centrifuged for subsequent RhB absorbance analysis. The remaining RhB dye concentration was detected by ultraviolet-visible (UV-Vis) spectrophotometer at a wavelength of 554 nm (Bhargava et al. 2016). The degradation efficiency $\eta$ was calculated according to Equation (3).

$$\eta = \frac{C_0V_o - C_eV_e}{C_0V_o} \times 100\%$$

(3)

**Table 1** Main chemical compositions and molar ratio of Zn to Fe of zinc-bearing dust and mixtures with different mass ratios used in this study wt.%

<table>
<thead>
<tr>
<th>Chemical compositions</th>
<th>Fe$_2$O$_3$</th>
<th>ZnO</th>
<th>MnO$_2$</th>
<th>MgO</th>
<th>CuO</th>
<th>PbO</th>
<th>Cr$_2$O$_3$</th>
<th>CaO</th>
<th>SiO$_2$</th>
<th>Molar ratio of Zn to Fe</th>
</tr>
</thead>
<tbody>
<tr>
<td>Zinc-bearing dust</td>
<td>64.1</td>
<td>10.4</td>
<td>0.8</td>
<td>1.7</td>
<td>0.3</td>
<td>1.3</td>
<td>0.5</td>
<td>1.0</td>
<td>5.3</td>
<td>0.32:2</td>
</tr>
<tr>
<td>2:0.4</td>
<td>64.1</td>
<td>30.4</td>
<td>0.8</td>
<td>1.7</td>
<td>0.3</td>
<td>1.3</td>
<td>0.5</td>
<td>1.0</td>
<td>5.3</td>
<td>0.93:2</td>
</tr>
<tr>
<td>2:0.6</td>
<td>64.1</td>
<td>40.4</td>
<td>0.8</td>
<td>1.7</td>
<td>0.3</td>
<td>1.3</td>
<td>0.5</td>
<td>1.0</td>
<td>5.3</td>
<td>1.25:2</td>
</tr>
<tr>
<td>2:0.8</td>
<td>64.1</td>
<td>50.4</td>
<td>0.8</td>
<td>1.7</td>
<td>0.3</td>
<td>1.3</td>
<td>0.5</td>
<td>1.0</td>
<td>5.3</td>
<td>1.54:2</td>
</tr>
</tbody>
</table>
where $C_o$ and $C_e$ represent the concentration of RhB dye in the initial solution and after the degradation process, respectively, g·mL$^{-1}$; $V_o$ and $V_e$ are the volumes of the initial RhB dye solution and after the degradation process, respectively, mL.

**Toxicity leaching test**

To evaluate the chemical stability and ensure the stability toward metal ion leaching from as-prepared mixed iron oxides to the solution under the reaction conditions, the Toxicity Characteristic Leaching Procedure (TCLP) (1311) formulated by the US Environmental Protection Agency (US EPA) (Sebag et al. 2009) was adopted and distilled water with a pH value of 2.88 was chosen as the leaching agent. The test procedures were as follows: firstly, 1.0 g zinc-bearing dust or as-prepared mixed iron oxide and 20 mL leaching agent were mixed in a centrifuge tube, and the mixture was shaken at a speed of 30 rpm at 25°C for 24 h, and then filtered to obtain the filtrate. The concentration of metal ions in the filtrate were determined by inductively coupled plasma optical emission spectrometer (ICP-OES).

**Analysis and characterization**

The mineral phase compositions of as-prepared samples were recorded by X-ray diffractometer (XRD, Rigaku, Cu Kα radiation, $\lambda = 0.15406$ nm) at a scanning rate of 0.02 deg·s$^{-1}$ at the diffraction angle ($2\theta$) 10–90° with a voltage of 40 kV and 40 mA. Room temperature Raman spectra of as-prepared samples were tested using a Raman spectrometer equipped with an Ar$^+$ laser (532 nm, 10 mW) excitation source and a CCD detector. The UV-Vis spectra of as-prepared samples were measured with an UV-Vis spectrophotometer (Hitachi U-3010). Physical Property Measurement System (PPMS, America, 9 T (EC-II)) was explored to test the magnetic properties of as-prepared samples with the applied magnetic field varying from −10,000 to 10,000 Oe.

**RESULTS AND DISCUSSION**

**Characterization of mixed iron oxides**

The XRD patterns of as-prepared samples with different mass ratios of zinc-bearing dust to ZnO addition of 2:0.4, 2:0.6, and 2:0.8 is shown in Figure 1. According to the PDF card, peaks appear at 29.9°, 35.3°, 42.8°, 56.6° and 62.2° match well with PDF card 22–1022 (Franklinite, ZnFe$_2$O$_4$) (National Bureau of Standards 1971), which correspond to the crystal face (220), (311), (400), (511) and (440) of franklinite, respectively. Peaks appear 24.1°, 33.2°, 40.9° and 49.5° match well with PDF card 33–0664 (Hematite, Fe$_2$O$_3$) (National Bureau of Standards 1981), which correspond to the crystal face (012), (104), (113) and (024) of hematite, respectively. As can be seen from Figure 1, when the mass ratio is controlled at 2:0.4, the diffraction peaks of both spinel ferrite ZnFe$_2$O$_4$ and the rhombohedral structure of Fe$_2$O$_3$ were detected. With the mass ratio decreasing to 2:0.6 and 2:0.8, the intense diffraction peaks were well indexed to spinel ferrite ZnFe$_2$O$_4$, and the diffraction peaks of Fe$_2$O$_3$ disappeared, indicating the formation of single-phase spinel ferrite. Furthermore, with the mass ratios increasing, the diffraction peaks increased in intensity, implying that the purity and crystallinity of as-prepared samples might be improved.

In the Raman spectra, there are five optical Raman-active modes for spinel space group (Fd3 m), A1g + Eg + 3T2g. The notations A, E and T are one, two and three dimensional representations, respectively, and g represents the symmetry regarding the center of inversion. In spinel ferrites, the assignment of the Raman-active modes and the regions where they are located are given as follows:

(i) The T2g(1) active mode located at 213–243 cm$^{-1}$ are associated to translational movements of the whole MO$_4$ tetrahedral. M and O represent Metal and Oxygen, respectively.

(ii) The Eg active mode located at 334–352 cm$^{-1}$ are related to symmetric bending vibrations of O atoms with respect to M in tetrahedral (A) sites.
(iii) The T2 g(2) active mode appearing at 478–488 cm⁻¹ is ascribed to asymmetric stretching vibrations of M–O in tetrahedral sites.

(iv) The T2 g(3) active mode appearing at 540–571 cm⁻¹ is assigned to asymmetric bending vibration of O atoms with respect to M in tetrahedral sites.

(v) The A1g symmetry active mode located in the range 600–710 cm⁻¹ is related to symmetric stretching vibrations of M–O in tetrahedral (A) sites. For this mode, it has been confirmed that the Raman spectrum of normal spinel ferrites is located in the 600–620 cm⁻¹ region, while that of inverse spinel ferrite occurs in the 670–710 cm⁻¹ region.

The as-prepared mixed iron oxides are also characterized by Raman spectra, as presented in Figure 2. It can be observed that the peaks appearing around 205, 280, 335, 475, 645 and 700 cm⁻¹ were assigned to the A1g, Eg, and 3T2g Raman active modes of spinel ferrite ZnFe2O4 (Yan et al. 2015; Aakash et al. 2019), further confirming the formation of spinel ferrite. Furthermore, the A1g(1) mode (around 700 cm⁻¹) and A1g(2) mode (around 640 cm⁻¹) are the symmetric stretch of O along the Fe-O bonds and Zn-O bonds at the tetrahedral sites, respectively (Thota et al. 2015). With the mass ratios of zinc-bearing dust to ZnO addition decreasing from 2:0.4 to 2:0.8, the intensity of A1g(1) mode decreased while that of the A1g(2) mode increased, implying that Zn²⁺ ion substitution into spinel ferrites could cause cation redistribution. When Zn was substituted in the as-prepared spinel ferrites, Zn²⁺ ions preferred to occupy the tetrahedral (A) sites, and forced the same amount of Fe³⁺ ions to transfer to the octahedral (B) sites.

As a photo-Fenton catalyst, it is essential to understand the optical properties of as-prepared samples, and Figure 3 presents the UV-Vis spectra of as-prepared ZF1, ZF2 and ZF3 with wavelengths ranging from 200 to 800 nm. All three samples exhibit intense absorption in a wide wavelength varying from UV to visible light with an absorption tail extending into the infrared region. The band gap (Eg) can be calculated by the Tauc’s equation as follows:

\[ (\alpha h\nu)^n = A(h\nu - E_g) \]  

where \( \alpha \) is the absorption coefficient, \( h \) is the Planck’s constant, \( \nu \) is the frequency of light and \( A \) is a constant. The bandgap energy could be evaluated for direct (\( n = 2 \)) and indirect (\( n = 1/2 \)) electronic transitions determined by extrapolation of the linear regions of the Tauc’s plots to zero absorption. The bandgap energy is obtained by extrapolating the tangent of the curve to x-axis as plotted in the inset. The bandgaps for ZF1, ZF2 and ZF3 are 1.92 eV, 1.94 eV and 1.96 eV, respectively, which are close to that for zinc ferrite.

The magnetic properties of as-prepared samples were measured by PPMS, and room temperature hysteresis loops are illustrated in Figure 4. All the as-prepared samples exhibit ferrimagnetism behaviors. With Zn²⁺ ion substitution content increasing (mass ratios from 2:0.4 to 2:0.8), the saturation magnetization values decreased from 33.8 emu g⁻¹ to 20.3 emu g⁻¹. From the inset of Figure 4, it can be observed that the as-prepared sample could be easily separated from the reaction system by using a magnet, providing a practical approach to recycle and reuse the catalyst.
Degradation of dye

The RhB degradation experiments using as-prepared ZF1, ZF2 and ZF3 as catalysts were carried out as shown in Figures 5 and 6. Figure 5 shows the degradation efficiency of RhB using ZF2 as the catalyst under different reaction conditions. Clearly, as shown in Figure 5, the self-degradation of RhB is very weak under visible light irradiation in the absence of mixed iron oxides and H2O2. It can be also observed that the degradation efficiency is only 2% using H2O2 as the catalyst in the dark after reaction for 210 min. Under the visible light irradiation, the degradation efficiency could be improved (about 50%) using H2O2 as the catalyst in the dark after reaction for 210 min. Using ZF2 as catalyst, the degradation efficiency is increased to 50% with the addition of H2O2 in the dark. However, using ZF2 as photocatalyst without the addition of H2O2, the degradation efficiency is about 20%, implying that ZF2 is not an efficient photocatalyst for the degradation of RhB. In the presence of H2O2 and with visible light irradiation, ZF2 catalyst could reach 91% RhB degradation. Such results demonstrated that zinc-bearing dust-derived mixed iron oxide ZF2 is an effective heterogeneous photo-Fenton-like catalyst for RhB degradation. Besides, the key roles of H2O2 addition and visible light irradiation are also indispensable in the photo-Fenton reaction system.

As observed from Figure 6, the degradation of RhB for ZF1, ZF2 and ZF3 catalysts slightly increases from 91% to 95% after reaction for 210 min, which might be ascribed to the active catalytic sites increasing when more Zn2⁺ ions occupied the tetrahedral (A) sites and forced more Fe3⁺ ions to occupy the octahedral (B) sites, as discussed in Figure 2. Moreover, the required reaction time for the similar degradation of dye using as-synthesized mixed iron oxide catalysts is longer than that of some reported literatures (Cao et al. 2011; Diao et al. 2011), which might be ascribed to the relatively low catalytic activity of as-synthesized mixed iron oxide catalysts resulting from high synthesis temperature as analyzed in the literature (Cao et al. 2011; Ismael 2021).

To further identify the main active oxidative species in the visible light/ZF2/H2O2 system, the degradation experiments with the addition of hydroxyl radicals -OH scavenger (tert-butanol (TBA)) and the holes scavenger (Ethylene Diamine Tetraacetic Acid (EDTA)) were carried out and the results are shown in Figure 7. As observed in Figure 7, with the addition of EDTA (1 mM), the degradation of RhB decreases from 91% to 60%, implying that
EDTA have some effects on the degradation of RhB. When TBA was added in the system, the degradation of RhB decreases from 91% to 48%, indicating that TBA (1 M) greatly inhibited the degradation reaction. As a result, the generated hydroxyl radicals \( \cdot \text{OH} \) played a key role in degradation of RhB and the generated electrons were of great importance for the degradation process.

Generally, in a Fenton reaction system, \( \text{Fe}^{2+} \) ions can be oxidized by \( \text{H}_2\text{O}_2 \) to generate powerful hydroxyl radicals \( \cdot \text{OH} \), which have a standard potential of 2.8 V (Kostedt et al. 2005). They can oxidize and mineralize organic pollutants in wastewater. In the \( \text{H}_2\text{O}_2 \) system, few hydroxyl radicals \( \cdot \text{OH} \) could be generated and resulted in low degradation efficiency. However, with visible light irradiation, hydroxyl radicals \( \cdot \text{OH} \) can be yielded in the \( \text{H}_2\text{O}_2 \) system, leading to the improvement of degradation efficiency. Using the mixed iron oxides as Fenton-like catalyst in the presence of \( \text{H}_2\text{O}_2 \), the plus two (+2) and three (+3) Fe ions could be oxidized by \( \text{H}_2\text{O}_2 \), and lots of hydroxyl radicals \( \cdot \text{OH} \) were produced (Munoz et al. 2015). As a result, the degradation efficiency could be obviously increased. Furthermore, in the presence of \( \text{H}_2\text{O}_2 \) with visible light irradiation, the as-prepared mixed iron oxides exhibited enhanced RhB degradation performance, implying that visible light irradiation is essential for effective dye degradation. The results might be ascribed to the generation of hydroxyl radicals \( \cdot \text{OH} \) being accelerated by visible light irradiation, and the synergistic effect of the photocatalyst reaction and Fenton-like catalyst reaction. Figure 8 shows the proposed mechanism for degradation of RhB by Zinc-bearing dust derived mixed iron oxides photo-Fenton catalysts.

**Catalyst stability and reusability**

As known, many heavy metals such as Zn, Mn, Cu, Pb, Cr are in the zinc-bearing dust. It is necessary to ensure security and stability towards hazardous metal leaching from as-prepared mixed iron oxides into the solutions under the reaction conditions. The hazardous metal leaching characteristics of zinc-bearing dust and as-prepared mixed iron oxides samples were tested according to the TCLP process, and the results are listed in Table 2. It can be found that generated hydroxyl radicals \( \cdot \text{OH} \) played a key role in degradation of RhB and the generated electrons were of great importance for the degradation process.

Table 2 | Metal ion concentrations in the solutions leached from zinc-bearing dust and as-prepared ZF1, ZF2 and ZF3

<table>
<thead>
<tr>
<th>Samples</th>
<th>Fe (mg/L)</th>
<th>Zn (mg/L)</th>
<th>Mn (mg/L)</th>
<th>Cu (mg/L)</th>
<th>Pb (mg/L)</th>
<th>Cr (mg/L)</th>
<th>Mg (mg/L)</th>
<th>Ca (mg/L)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Raw dust</td>
<td>12.23 ± 0.5</td>
<td>30.05 ± 0.6</td>
<td>0.92 ± 0.2</td>
<td>0.33 ± 0.1</td>
<td>12.05 ± 0.3</td>
<td>10.11 ± 0.3</td>
<td>0.63 ± 0.1</td>
<td>0.54 ± 0.1</td>
</tr>
<tr>
<td>ZF1</td>
<td>0.12 ± 0.02</td>
<td>0.30 ± 0.08</td>
<td>0.18 ± 0.03</td>
<td>0.09 ± 0.02</td>
<td>0.27 ± 0.05</td>
<td>0.26 ± 0.04</td>
<td>0.26 ± 0.05</td>
<td>0.06 ± 0.02</td>
</tr>
<tr>
<td>ZF2</td>
<td>0.07 ± 0.02</td>
<td>0.26 ± 0.05</td>
<td>0.07 ± 0.02</td>
<td>0.08 ± 0.02</td>
<td>0.15 ± 0.02</td>
<td>0.15 ± 0.03</td>
<td>0.29 ± 0.05</td>
<td>0.09 ± 0.02</td>
</tr>
<tr>
<td>ZF3</td>
<td>0.06 ± 0.01</td>
<td>0.38 ± 0.06</td>
<td>0.24 ± 0.03</td>
<td>0.06 ± 0.02</td>
<td>0.11 ± 0.02</td>
<td>0.07 ± 0.02</td>
<td>0.23 ± 0.05</td>
<td>0.11 ± 0.02</td>
</tr>
<tr>
<td>Maximum limit (Wang et al. 2017)</td>
<td>4</td>
<td>2</td>
<td>2</td>
<td>0.5</td>
<td>0.5</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
</tbody>
</table>
heavy metals leaching from zinc-bearing dust to the solution is serious, and most of the heavy metal concentrations are above the maximum limit; however, the heavy metals leaching from as-prepared samples are very limited (far below the maximum limit), implying that heavy metals could be fixed and doped in the stable spinel structure during the facile solid state calcination process, as reported by Chen et al. (2011).

Recovery and reusability are important aspects of a resource-saving and environment-friendly catalyst for practical application in wastewater treatment. The reusability of as-prepared ZF2 was studied by a five-cycle degradation of RhB under identical reaction conditions as plotted in Figure 9. The degradation efficiency was slightly decreased from 91% to 87%, implying that zinc-bearing dust-derived mixed iron oxides were stable during the reaction and could be reusable catalysts for degradation of dye. In general, zinc-bearing dust-derived mixed iron oxides could be easily separated and reused as heterogeneous photo-Fenton-like catalysts.

**CONCLUSIONS**

In conclusion, non-toxic mixed iron oxides as the magnetically recyclable photo-Fenton catalyst for degradation of dye were synthesized from zinc-bearing dust using a facile solid state reaction method. In the as-prepared samples, spinel ferrite coexisting with or without Fe₂O₃ was the predominant phase. The as-prepared mixed iron oxides exhibited excellent degradation efficiency (above 90%) of RhB in the presence of 2.0 mmol/L H₂O₂ and with visible light irradiation for 210 min when the initial RhB concentration is 10 mg/L and the pH value is controlled at 4.0, owing to the synergistic effect of the photocatalyst reaction and Fenton-like catalyst reaction during the degradation process. When more Zn²⁺ ions occupied the tetrahedral (A) sites and forced more Fe³⁺ ions to occupy the octahedral (B) sites, the active catalytic sites were increased, and the degradation efficiency was increased accordingly. Moreover, heavy metals in the zinc-bearing dust were fixed stably in the spinel structure, and the as-prepared samples presented excellent reusability with a degradation efficiency of 87% after the fifth cycle of reuse. This work paves a green pathway towards the comprehensive utilization of hazardous solid waste to produce high value-added material product.

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**DATA AVAILABILITY STATEMENT**

All relevant data are included in the paper or its Supplementary Information.

**REFERENCES**


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