Preparation and Photocatalytic Performance of ZnO/Sepiolite Composite Materials

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Photocatalytic technology is a widely used water treatment method, whose efficiency can be increased by developing a suitable photocatalyst fabrication procedure. In this study, five different synthesis methods were utilised for the preparation of novel ZnO/sepiolite photolytic composites, namely, sol-gel method, hydrothermal reduction, hydrolytic precipitation, powder sintering, and impregnation-reduction. The obtained photocatalysts were characterised by scanning electron microscopy, infrared spectroscopy, and X-ray diffraction. The differences between the applied photocatalyst preparation methods and the reasons for these differences were discussed, and the photocatalytic activities of the prepared composite materials were compared. The obtained results revealed that the physical structure, chemical properties, and photocatalytic performance of the composite produced by the sol-gel method were superior to those of the materials fabricated by the other four methods. Moreover, this material also exhibited high photocatalytic stability, while its photocatalytic degradation of methylene blue dye proceeded via a quasi-first-order reaction. The prepared composites have broad application prospects in photocatalysis and can be potentially used for treating environmental pollutants.

1. Introduction

In 1977, Frank and Bard used TiO2 photocatalytic technology to treat CN− and SO3−2 plasmas, after which photocatalytic methods were successfully applied to control environmental pollution [1]. Photocatalytic materials typically include semiconductors such as metal oxide sulphides. Among various photocatalysts and their composites, TiO2 exhibits high and stable photocatalytic activity and, therefore, is commonly utilised in photocatalytic reactions. Some researchers studied [2–5] the photocatalytic performance of TiO2 for the treatment of aromatic hydrocarbon pollutants and methyl blue dye followed by its successful application in the degradation of landfill leachate. However, photocatalytic TiO2 pollution treatment technology has not been widely used in practical applications for the following reasons. (1) The adsorption capacity of TiO2 is relatively small, and the adsorbed species are difficult to separate in a liquid phase [6, 7]. (2) TiO2 has a large bandgap, which can absorb not more than 4–5% of the ultraviolet (UV) light from the solar spectrum, while its spectral response range is narrow [8, 9]. (3) TiO2 particles easily agglomerate, which reduces their practicability [10, 11]. Therefore, exploring other types of photocatalysts has recently become a hot research topic.

ZnO is a semiconductor with a forbidden band width of 3.37 eV and exciton binding energy of 60 MeV at room temperature, which is widely applied for the degradation of organic pollutants due to its excellent chemical stability, high photocatalytic activity, low fabrication cost, and other advantages [12–16]. The results of ZnO-related studies revealed that the photocatalytic activity of zinc oxide was sufficiently high to decompose various pollutants in the environment [17]. For example, it can photocatalyse the degradation of harmful compounds [18, 19] such as aniline [20] under low UV light irradiation. In addition, ZnO may be doped by various compounds such as lanthanum and graphene [21, 22] that effectively photocatalyse the degradation of methyl orange and methylene blue (MB) dyes. However, the usability of ZnO in photocatalytic degradation applications has several limitations, including a wide bandgap [23, 24], high recombination rate of photogenerated electrons with holes, and low quantum efficiency [25, 26]. To increase the photocatalytic efficiency of...
ZnO and expand the range of its spectral response to visible light. ZnO modification must be performed via metal/non-
metal ion doping and/or surface complexation reactions, which require the preparation of multiple composite materials.

2. Literature Review

The composite materials produced by various methods have different crystal structures, morphologies, optical properties, and photocatalytic efficiencies [27, 28]. Several researchers fabricated nanosized TiO₂ powder [29] and Sn-Al co-doped ZnO photocatalyst by sol-gel method [30]. The photocatalytic properties of the Cu/ZnO catalysts prepared by hydrolysis method [31] and graphene-ZnO nanorod composites [32] were strongly influenced by the proportions of the raw materials; furthermore, the catalysts fabricated by different methods possessed different particle shapes and specific surface areas. For example, the Al-doped ZnO nanoparticles [33], ZnO nanoparticles [34, 35], and CuO/ ZnO catalysts [36] prepared by the precipitation method showed that nanoparticle sizes could vary from several nanometres to several tens of nanometres under different experimental conditions and even reach hundreds of nanometres in the case of needle-like particles with various morphologies and degrees of cleanliness. The nanometre-sized zinc oxide powder synthesised by this method had pseudospherical particle shapes. In addition, a ZnO/ZnSe composite with a maximum degradation rate of 91% was fabricated by a one-pot method [37], while ZnO nanoparticles were produced by a microwave method [38], and their crystal morphology was enhanced by optimising the solvent properties and crystal growth process.

In this study, sepiolite was used as a support for loaded ZnO particles. Because sepiolite is a hydrated magnesium silicate clay mineral with the largest specific surface area among nonmetallic minerals (up to 900 m²/g), unique channel structure, high surface adsorption capacity, low weight, and high chemical stability, it can serve as the supporting material for the active photocatalyst components. To identify the best ZnO/sepiolite preparation method, the sepiolite support was modified by a thermal acid treatment, after which ZnO/sepiolite composite materials were fabricated by the sol-gel, hydrothermal reduction, hydrolytic precipitation, powder sintering, and impregnation-reduction methods. The obtained composites were characterised by various techniques, and the loading degrees and morphologies of ZnO particles were determined.

3. Experimental

3.1. Materials and Instruments

3.1.1. Materials. The following materials were used: sepiolite raw ore, hydrochloric acid (analytical grade), sodium hexametaphosphate, distilled water, sodium hydroxide, zinc chloride, ethylene glycol, zinc acetate, anhydrous ethanol, ammonia solution, ice water, ammonium sulphate, silver nitrate, zinc oxide, zinc sulphate, hydrazine hydrate, and MB.

3.1.2. Instruments. The following instruments were used: vacuum filter extractor, 60 mesh sieve, 80 mesh sieve, constant temperature drying oven, muffle oven, X-ray powder diffractometer, scanning electron microscope, water bath, magnetic stirrer, vacuum filter extractor, hydrothermal reaction kettle, electronic scale, pH meter, 50 W high-pressure mercury lamp, TGA-DSC/DTA synchronous thermal analyzer, UV-visible spectrophotometer.

3.2. Sepiolite Modification Procedure

3.2.1. Sepiolite Pretreatment. The raw sepiolite ore contained many impurities, including sepiolite, talc, calcite, and kaolinite, which required pretreatment. Briefly, 500 g of sepiolite was soaked in distilled water at a solid-liquid ratio of 1:10. After 24 h, 0.5 g of sodium hexametaphosphate was added to the obtained mixture, followed by settlement. The upper suspended matter was removed; it was then filtered under vacuum and placed into a drying oven at 105°C. After drying for 24 h, a sepiolite concentrate was obtained.

3.2.2. Thermal Acid Modification. In the next step, sepiolite was modified by a thermal acid treatment. After pretreatment, sepiolite was screened with 60 mesh and 80 mesh sieves, and the intermediate fraction was soaked for 6 h and filtered under vacuum. The obtained solid particles were repeatedly washed with distilled water to the neutral pH, filtered, and placed into a drying oven at 105°C. The produced sepiolite was annealed at 250°C in a muffle oven for 3 h and cooled to room temperature.

3.3. Preparation of ZnO/Sepiolite Composites. ZnO/sepiolite composite materials were prepared by the sol-gel, hydrothermal reduction, hydrolytic precipitation, powder sintering, and impregnation-reduction methods. The obtained composites were characterised by various techniques, and the loading degrees and morphologies of ZnO particles were determined.

3.3.1. Sol-Gel Method. Initially, 16 g of sodium hydroxide was added to a specified amount of glycol solution to achieve a final volume of 200 mL. The resulting solution was placed on a magnetic stirrer and stirred until all sodium hydroxide particles were completely dissolved to obtain a 2 mol/L sodium hydroxide solution in ethylene glycol (solution A). Afterwards, 27.26 g of zinc chloride was added to the ethylene glycol solution to achieve a final volume of 200 mL. The resulting mixture was placed on a magnetic stirrer and stirred until all zinc chloride particles were completely dissolved to obtain a 1 mol/L zinc chloride solution in ethylene glycol (solution B). After mixing solutions A and B, the produced mixture was stirred for 30 min to obtain a precursor of zinc oxide sol, which was placed in a water bath with a temperature of 95°C for 3 h to produce zinc oxide sol. Finally, 16.26 g of the activated and modified sepiolite was added to the sol, stirred at room temperature for 1 h, heated in a water bath at 80°C until it became almost dry, cooled,
and dried in a drying oven at 80°C. The obtained product was placed into a muffle furnace and heated to 450°C. After continuous annealing for 4 h, the solid was taken out of the furnace and cooled to room temperature to obtain a ZnO/sepiolite composite material.

3.3.2. Hydrothermal Reduction Method. First, 4.51 g of zinc acetate was mixed with 20 mL of distilled water and 60 mL of anhydrous ethanol. The resulting mixture was stirred until all zinc acetate particles were completely dissolved followed by the addition of 2 g of modified sepiolite and continuous stirring for 30 min. After that, 8 mL of ammonia solution was diluted with 20 mL of anhydrous ethanol, and the obtained product was added dropwise to the mixture. The latter was stirred for 1 h, transferred to a 200 mL hydrothermal reaction kettle, and heated to 120°C for 5 h. Subsequently, the reaction mixture was taken out of the kettle and placed into a drying oven with a temperature of 105°C. The dry product was heated in a muffle furnace at 450°C for 4 h and then cooled down to room temperature to obtain a ZnO/sepiolite composite.

3.3.3. Hydrolytic Precipitation Method. First, ice was crushed with an ice breaker. Then, 200 mL of distilled water was added to 20 g of sepiolite to achieve a solid-to-liquid mass ratio of 1:10. After mixing, the solution was placed into an ice water bath followed by the addition of 1 mL of pure hydrochloric acid. In the next step, 25.215 g of zinc chloride was weighted and dissolved in an appropriate amount of distilled water and then transferred to a volumetric flask with a constant volume of 50 mL to obtain a 3.7 mol/L zinc chloride solution. Subsequently, 19.8 g of ammonium sulphate was dissolved in an appropriate amount of distilled water and transferred to a volumetric flask with a constant volume of 100 mL to obtain a 1.5 mol/L ammonium sulphate solution. The prepared zinc chloride solution was added to the first solution. After 10 min, 10 mL of the ammonium sulphate solution was added to the reaction mixture, which was then stirred for 10 min and heated to 80°C for 1 h in a water bath. The mixture was the neutralised with ammonia solution to pH = 6, stored for 1 h, washed with distilled water, and filtered. Finally, 3.397 g of silver nitrate was dissolved in an appropriate amount of distilled water and transferred to a 200 mL volumetric flask to obtain a 0.1 mol/L silver nitrate solution. After detecting free chloride ions in the reaction mixture with this solution, the fabricated sample was dried in a drying oven at 105°C, heated to 450°C for 2.5 h in a muffle furnace, and cooled down to room temperature to obtain a ZnO/sepiolite composite.

3.3.4. Powder Sintering Method. After mixing 5 g of zinc oxide with 5 g of sepiolite, a certain amount of anhydrous ethanol was added to the obtained mixture to fully wet the solid particles, which were subsequently ground for 30 min. The resulting mixture was placed into a muffle furnace and continuously annealed at 300°C for 6 h. After cooling to room temperature, the sample was ground until its particle sizes became uniform to obtain a ZnO/sepiolite composite material.

3.3.5. Impregnation-Reduction Method. First, 12 g of sodium hydroxide was dissolved in an appropriate amount of distilled water and then transferred to a volumetric flask with a constant volume of 50 mL to obtain a 6 mol/L sodium hydroxide solution. Next, 100 mL of the hydrazine hydrate solution was diluted to 514.3 mL to obtain a 2 mol/L hydrazine solution. Subsequently, 16.144 g of zinc sulphate was added to 400 mL of deionized water. After all zinc sulphate species were completely dissolved, 8.138 g of sepiolite was added to the mixture, which was then stirred for 30 min using a magnetic stirrer. During this process, 5 mL of the 6 mol/L sodium hydroxide solution was added to the reaction system. After continuous stirring for 30 min, 3 mL of the 2 mol/L hydrazine solution was added dropwise, and after 10 minutes of reaction, 0.015 mol of ammonia (2.25 mL of the concentrated ammonia solution) was added to the mixture. The subsequent reaction was performed at room temperature for 3 h. A ZnO/sepiolite composite was obtained by washing the resulting solid with deionized water three times and then with anhydrous ethanol three times. The composite was annealed in a muffle furnace at 450°C for 3 h and then cooled down to room temperature.

3.4. Photocatalytic Performance Testing. To investigate the photocatalytic properties of the produced composites, the photocatalytic degradation of MB was performed at room temperature under simulated UV light irradiation with a power of 50 W using a high-pressure mercury lamp. In selected reactions, 20 mg of the photocatalyst was dispersed in 100 mL of the MB aqueous solution (0.015 g/L) under magnetic agitation. In all photodegradation experiments, the distance between the experimental device and the light source was 7 cm. Before the photodegradation experiment, the catalyst-dye mixture was stirred in the dark for 1 h to reach an adsorption-desorption equilibrium. Afterwards, the mixture was exposed to the light generated by the high-pressure mercury lamp and continuously stirred. During the reaction, 3 mL of the mixture was taken out every 10 min, after which the catalyst was removed by centrifugation, and the change in absorbance was measured by the UV-Vis spectrophotometer. The reusability and photostability of the composite with the best photocatalytic performance were evaluated during the first five MB degradation cycles. Finally, the photocatalytic mechanism of the composite was analyzed, and the methods to improve the photocatalytic effect were proposed.

4. Results and Discussion

4.1. Morphological Analysis. Each analyzed sample was attached to a piece of copper sheet with conductive tape, sprayed with gold, and placed on the sample holder for SEM testing. Figures 1 and 2 display the SEM images of sepiolite obtained before and after thermal acid modification.
The figures show that the modified sepiolite has a higher specific surface area, larger surface pores, and looser structure as compared with those of the nonmodified sepiolite. Moreover, it likely exhibits higher catalytic activity and higher degree of immobilisation of ZnO particles.

The SEM image of the ZnO/sepiolite composite prepared by the sol-gel method is shown in Figure 3. In contrast to Figure 2, the composite surface is completely covered by the layer of a gel-like material consisting of the loaded zinc oxide gel obtained from the sol precursor. These results indicate that zinc oxide has successfully adhered to the sepiolite surface.

The SEM image of the composite prepared by the hydrothermal reduction method is presented in Figure 4. Similar to Figure 3, the sepiolite surface is covered with some material. In the hydrothermal reduction method, a zinc salt is crystallised on sepiolite under hydrothermal conditions and then decomposed into a metal oxide at a high temperature. As a result, zinc ions are loaded on the sepiolite surface in the form of crystals and then oxidised to produce zinc oxide. Therefore, the material deposited on the sepiolite surface consists of zinc oxide; however, its particle distribution is not uniform.

The SEM image of the ZnO/sepiolite composite fabricated by the hydrolytic precipitation method is displayed in Figure 5. It shows a layer of fine particles with a relatively large mass attached to the sepiolite surface; however, their distribution is not uniform.

The SEM image of the composite prepared by the powder sintering method is shown in Figure 6. In this image, the sepiolite surface is coated by the fine ZnO particles with an average loading; however, their distribution is more uniform than that of the particles prepared by the hydrolytic precipitation method.

The SEM image of the ZnO/sepiolite composite fabricated by the impregnation-reduction method is presented in Figure 7. It shows that the coated particles have a large load and uniform distribution. Here, zinc oxide is attached to the
sepiolite surface in the form of small particles with a large specific surface area, containing many catalytically active sites. The composite prepared by this method retained the large specific surface area; however, a significant fraction of zinc oxide particles precipitated and aggregated on the sepiolite surface.

4.2. Spectral Analysis. A sample was evenly mixed with potassium bromide and pressed into tablets. Its infrared spectrum was recorded by an infrared spectrometer. Figure 8 shows the infrared spectrum of sepiolite, which contains the characteristic peaks of its various groups. In the high-frequency region, the absorption band at 2900–3700 cm\(^{-1}\) is generated by the stretching vibration of the OH group; the absorption band at 3676.32 cm\(^{-1}\) is produced by the stretching vibration of the Mg-OH group; and the characteristic peak of crystal water is centred at 3564.45 cm\(^{-1}\). This peak belongs to the H-O-H bending deformation vibration zone located between 1400 and 1700 cm\(^{-1}\) and has a relatively high intensity. In the range of 900–1300 cm\(^{-1}\), there is a strong absorption band caused by the stretching vibration of the Si-O bond. In addition, the characteristic calcite peak is located between 600 and 800 cm\(^{-1}\). In the 400–600 cm\(^{-1}\) region, a strong bending deformation of the Si-O bond is observed.

Figure 9 displays the infrared spectrum of the modified sepiolite. In this sample, magnesium ions leached due to their substitution with hydrogen ions. As compared with Figure 8, the absorption peak centred at 3564.45 cm\(^{-1}\) is enhanced, and the peak at 3676.32 cm\(^{-1}\) is significantly weakened. Because calcite reacts with hydrochloric acid to produce carbon dioxide, the three absorption peaks in the 600–800 cm\(^{-1}\) region were transformed into two peaks, and the calcite characteristic peaks disappeared. Owing to the high-temperature annealing and activation processes, water was desorbed from the sepiolite surface; as a result, the characteristic peak at 1425.40 cm\(^{-1}\) disappeared. Meanwhile, the peak intensities in the low-frequency region did not change significantly after modification. Hence, the thermal acid treatment removed some magnesium ions, carbonate impurities, and adsorbed water from the sepiolite surface. Figure 10 shows the infrared spectrum of zinc oxide.

According to literature data, the absorption peak at 3427.51 cm\(^{-1}\) is the characteristic peak of the O-H bonds in physisorbed water molecules. The characteristic absorption peak of the nanosized zinc oxide is located between 400 and 600 cm\(^{-1}\) [39].

Figures 11–15 display the infrared spectra of the ZnO/sepiolite composites prepared by the sol-gel, hydrothermal reduction, hydrolytic precipitation, powder sintering, and impregnation-reduction methods, respectively.

The figures show that the wavelength range of the zinc oxide characteristic peak overlaps with that of sepiolite, which has a certain influence on the conclusion of the loading condition of zinc oxide on sepiolite. However, the zinc oxide loading on the sepiolite surface can be determined from the intensity variations of the characteristic absorption peaks located between 400 and 600 cm\(^{-1}\). The infrared spectrum of the composite sample obtained by the hydrothermal reduction method has the largest variation range of these peaks in the specified region followed by the spectra of the ZnO/sepiolite composites prepared by powder sintering, impregnation-reduction, and sol-gel methods.

4.3. Structural Analysis. In this paper, 100 mesh sepiolite from Xiangtan sepiolite Technology Co., Ltd., was selected, and the chemical formula was \(\text{Mg}_8\left(\text{H}_2\text{O}\right)_4\left(\text{Si}_6\text{O}_{10}\right)_2\left(\text{OH}_2\right)_4\cdot8\text{H}_2\text{O}\). The theoretical chemical composition is MgO = 21%–25%; SiO\(_2\) = 54%–60%. There is also a small number of exchange cations, the charge for the tetrahedral weight of the ion exchange produced. Si\(^{4+}\) in tetrahedron can be replaced by Fe\(^{3+}\) and Al\(^{3+}\), the Mg\(^{2+}\) in octahedron can be replaced by Mn\(^{2+}\), Fe\(^{2+}\) or Fe\(^{3+}\). XRD analysis shows that SiO\(_2\) content is high, containing Fe, K, Ca and other impurities. The spectral analysis results of sepiolite are shown in Table 1.

In Figure 16, the XRD analysis of sepiolite ore samples, the value of \(2\theta\) at 7.4° is the diffraction peak of sepiolite on plane 110, the \(2\theta\) values at 26.7° and 9.3° are characteristic diffraction peaks of quartz and talc, respectively, the diffraction peak of impurity calcite is \(2\theta\) at 29.59°, and the main
Figure 8: Infrared spectrum of sepiolite.

Figure 9: Infrared spectrum of the modified sepiolite.

Figure 10: Infrared spectrum of ZnO.
Figure 11: Infrared spectrum of the ZnO/sepiolite composite prepared by the sol-gel method.

Figure 12: Infrared spectrum of the ZnO/sepiolite composite prepared by the hydrothermal reduction method.

Figure 13: Infrared spectrum of the ZnO/sepiolite composite prepared by the hydrolytic precipitation method.
components of crystal are sepiolite, quartz, talc, and calcite. Calculate according to XRD software, 41.9% of sepiolite, 31.1% of quartz, 12.8% of talc, 14.1% of calcite; it indicates that the impurities in the original sepiolite ore used in this study are mainly quartz, calcite, and a small amount of talc.

The density of quartz is 2.65 g/cm³, that of sepiolite is 2.03 g/cm³, after purification by precipitation, and the diffraction peaks of sepiolite are enhanced, while those of quartz and talc are weakened. In Figure 17, calculated by XRD, the sepiolite content increased from 41.9% to 72%, quartz content decreased from 31.1% to about 10%, talc from 12.8% to 5%; it shows that the impurities are basically removed after purification and the purpose of purification is achieved.

Figures 16 and 18 display the XRD patterns of the nonmodified and modified sepiolite, respectively. They show that the spectrum of modified sepiolite lacks the calcite peak; in addition, the peaks corresponding to the other impurities either decreased in intensity or disappeared completely, while the intensity of the sepiolite peak increased from

| Ingredient | SiO₂ | Al₂O₃ | Fe₂O₃ | MgO | CaO | Na₂O | K₂O | MnO | TiO₂ | P₂O₅ | Loss on ignition |
|------------|------|-------|-------|-----|-----|------|-----|-----|------|------|-----------------|
| Content    | 64.45| 7.32  | 2.50  | 12.10| 4.55| 0.15 | 0.42| 0.05| 0.21 | 0.05| 8.20            |

Figure 14: Infrared spectrum of the ZnO/sepiolite composite prepared by the powder sintering method.

Figure 15: Infrared spectrum of the ZnO/sepiolite composite prepared by the impregnation-reduction method.
approximately 800 to about 950 after modification because hydrogen ions replaced magnesium ions in the sepiolite structure after the acid treatment, and the grain shape was good. After modification, the sepiolite characteristic peak shifted to a lower angle, indicating that both the pore size and specific surface area of sepiolite increased. The TGA/DTA test results of sepiolite are shown in Figure 19.

The experimental results show that the structure of sepiolite does not change from room temperature to 300°C, which is mainly due to the process of water desorption. The structure of sepiolite changes at 300–800°C, which is a process of decrystallising water, the sepiolite loses crystal water and forms anhydrous sepiolite phase, and its structure is locally adjusted. At 800–1000°C stage, the mineral composition of sepiolite is recombined and its structure is completely destroyed; the main process of dehydroxy water resulted in the formation of new mineral facies calcite and enstatite. Therefore, the thermal stability of sepiolite is better than other clay minerals, and good thermal stability is the premise of sepiolite as raw material for the preparation of composite materials.

Figure 20 depicts the XRD pattern of zinc oxide, which is consistent with the hexagonal wurtzite ZnO structure. No impurities were detected in the zinc oxide sample. The observed XRD peak is sharp and exhibits a relatively high intensity, which indicates a high degree of crystallinity. However, some widening was observed in the lower part of the diffraction peak, suggesting that ZnO grains were refined, which was consistent with the particle characteristics.

Figures 21–25 show the XRD patterns of the ZnO/sepiolite composites prepared by the sol-gel, hydrothermal reduction, hydrolytic precipitation, and impregnation-reduction methods, respectively. All these patterns exhibit smaller characteristic sepiolite peaks (as compared with those in the pure sepiolite spectra) as well as the characteristic peaks of anatase and zinc oxide. These phenomena indicate that the material deposited on the sepiolite surface consists of zinc oxide. Furthermore, the zinc oxide diffraction peaks of the composites prepared by the sol-gel method, hydrothermal reduction, and powder sintering methods are relatively strong and exhibit good shapes, whereas the ZnO peaks of the materials prepared by the other methods are relatively weak. The main ZnO/sepiolite composite characteristics are summarised in Table 2.

4.4. Photocatalytic Performance. To evaluate the photocatalytic performance of the fabricated composites, the ZnO/sepiolite composites and ZnO were applied to degrade organic pollutant MB. First, the catalytic material and MB were continuously stirred in the dark for 1 h to reach an adsorption-desorption equilibrium. The absorbance of the obtained solution was measured after centrifugation under UV irradiation at different intervals using the high-pressure mercury lamp. The observed variation of MB concentration with exposure time is shown in Figure 26. The absorption intensity of the raw MB (without ZnO or ZnO/sepiolite composites) exposed to UV-Vis light decreased only slightly, indicating that very little MB amount degraded under UV light irradiation in the absence of a catalyst (the corresponding degradation rate was only 6.5%). With the addition of ZnO or the ZnO/sepiolite composite materials, the MB concentration decreased with an increase in the irradiation time, suggesting that all these materials exhibited high photocatalytic activity for the MB degradation. The ZnO/sepiolite composite prepared by the sol-gel method degraded approximately 93.5% of the initial MB amount within 90 min, while the degradation rate of MB over ZnO catalyst was only 81%. Therefore, the catalytic activity of the ZnO/sepiolite composite for MB photodegradation was superior to that of ZnO catalyst.

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The obtained experimental data were fitted with the following equation derived for the first-order reaction kinetic model:

\[
\frac{-dC}{dt} = k \times C, \\
\ln \left( \frac{C}{C_0} \right) = k \times t, \quad (1)
\]

Here, \(C_0\) is the initial MB concentration, \(C\) is the MB concentration measured after a 10-minute interval, \(t\) is the irradiation time, and \(K\) (min\(^{-1}\)) is the first-order rate constant. Figure 27 shows the linear relationship between \(\ln(C/C_0)\) and degradation time \((t)\) under UV irradiation. After the addition of the ZnO/sepiolite composite prepared by the sol-gel method, the first-order rate constant of MB degradation was \(K = 0.03722 \text{ min}^{-1}\). Without the addition of a photocatalyst, its value was \(k = 0.0061 \text{ min}^{-1}\). Finally, after the ZnO addition, the first-order rate constant of MB degradation was \(K = 0.013889 \text{ min}^{-1}\). The obtained \(R^2\) values exceeded 0.9; hence, the photocatalytic degradation of MB was a quasi-first-order reaction.

The reusability and photostability of the ZnO/sepiolite composite, containing 100 mg, prepared by the sol-gel method were studied as well. In this experiment, five MB degradation cycles were performed. After each cycle, the utilised composite was collected and washed for the next degradation cycle. The obtained powder was dried at 60°C for 12 h. The photodegradation efficiencies of the ZnO/sepiolite composite measured during the five degradation cycles are shown in Figure 28 (their values are equal to 93.5%, 90.9%, 88.7%, 86.1%, and 85.8%). These results indicate that the ZnO/sepiolite composite prepared by the sol-gel method exhibits excellent reusability and photostability properties, which can be attributed to its high chemical stability and photoelectric performance.

The phase structure of the composites after recycling was analysed by XRD to observe whether the material structure changes after photocatalytic reaction.

As shown in Figure 29, the XRD pattern of the catalyst after repeated use was basically the same as the composition of the original compound. It shows that ZnO is well loaded on the sepiolite surface and the bond is firm. Therefore, ZnO/sepiolite composite is a kind of photocatalytic composite material with stable chemical properties, easy to be separated, and easy to be recycled.
Figure 16: XRD pattern of sepiolite.

Figure 17: XRD phase analysis of sepiolite and purified sepiolite (1: raw ore; 2: purification of sepiolite; S: sepiolite; T: talc; Q: quartz; C: calcite).

Figure 18: XRD pattern of the modified sepiolite.
4.5. Reaction Mechanism. When zinc oxide, as a semiconductor photocatalyst, is exposed to ultraviolet light, the valence band electrons of zinc oxide are excited to transition to the conduction band; thus, holes ($h_{VB}^+$) are generated in the valence band and photogenerated electron ($e_{CB}^-$) pairs are generated in the conduction band. In the solution, $h_{VB}^+$ and $e_{CB}^-$ can react with water and oxygen in a series of reactions, generates hydroxyl radical ($\cdot$OH) and other strong oxidation radicals, which oxidizes and degrades organic matter thoroughly and mineralizes into inorganic small molecules. The reaction steps are as follows:

![Test result](image1)

**Figure 19: Test result.**

![XRD pattern of ZnO](image2)

**Figure 20: XRD pattern of ZnO.**

![XRD pattern of the ZnO/sepiolite composite prepared by the sol-gel method](image3)

**Figure 21: XRD pattern of the ZnO/sepiolite composite prepared by the sol-gel method.**
Figure 22: XRD pattern of the ZnO/sepiolite composite prepared by the hydrothermal reduction method.

Figure 23: XRD pattern of the ZnO/sepiolite composite prepared by the hydrolytic precipitation method.

Figure 24: XRD pattern of the ZnO/sepiolite composite prepared by the powder sintering method.
\[
\begin{align*}
\text{ZnO} + h\nu & \rightarrow e_{CB}^- + h_{VB}^+ \quad (2) \\
O_2 + e_{CB}^- & \rightarrow O_2^- \quad (3) \\
(H_2O \rightarrow H^+ + OH^-) + h_{VB}^- & \rightarrow \cdot OH + H^+ \quad (4) \\
O_2^- + H^+ & \rightarrow HO_2 \quad (5) \\
2HO_2 & \rightarrow H_2O_2 + O_2 \quad (6)
\end{align*}
\]

\[
\text{H}_2\text{O}_2 + e^- \rightarrow \cdot \text{OH} + \text{OH}^- \quad (7)
\]

\[
\text{MB} + \cdot \text{OH} \rightarrow \text{degradation product} + \text{H}_2\text{O} \quad (8)
\]

- \cdot \text{OH} is the most important oxidant for MB degradation, and the production of \cdot \text{OH} is directly related to the production of \text{h}_{VB}^-.

- Under UV irradiation, electron-hole pairs are generated on the ZnO surface. At the same time, when the concentration of OH- ions in the solution increases, more OH- will be oxidised to \cdot \text{OH} by the holes generated on the ZnO surface.
Figure 26: MB concentration plotted as a function of the exposure time for the ZnO/sepiolite composites prepared by five different methods (1: sol-gel; 2: hydrothermal reduction; 3: hydrolytic precipitation; 4: powder sintering; 5: impregnation-reduction).

Figure 27: \( \ln(C_0/C) \) plotted as a function of the degradation time \( t \) for various samples subjected to UV light irradiation (1: ZnO/sepiolite composites).

Figure 28: MB photodegradation efficiencies measured during the five treatment cycles.
the photocatalyst surface. The increase of \( \cdot \)OH concentration is conducive to the improvement of the treatment effect. Therefore, the \( h_{VB}^+ \) increased yield or the addition of cocatalyst (strong oxidant) is beneficial to improve the processing capacity of photocatalytic materials. Meanwhile, in the process of degrading MB, MB can be used as a photosensitizer. Under ultraviolet light irradiation, the system is photosensitized, and the ground-state oxygen is irradiated to produce singlet oxygen \((^1O_2)\):

\[
\text{sensitizer} - \text{hv} - \text{sensitizer}^{T1} \tag{9}
\]

\[
\text{sensitizer}^{T1} + \text{O}_2 \text{(groundstate)} - \text{energy transfer} - \text{sensitizer} + ^1\text{O}_2 \tag{10}
\]

5. Conclusion

In this study, sepiolite was used as a catalytic substrate to synthesise novel ZnO/sepiolite composites by five different methods, and the catalytic efficiencies of the produced materials were determined by investigating the photocatalytic degradation of MB dye. The microstructures of these materials were observed, their photocatalytic properties and stabilities were evaluated, and their reaction kinetics was examined. The obtained conclusions were summarised as follows:

(1) After the thermal acid treatment, carbonate impurities were removed from the sepiolite surface, while the sizes of sepiolite pores and channels as well as the specific surface area and catalytic activity of sepiolite increased.

(2) The ZnO/sepiolite composites prepared by different methods had different loading forms and loads. When the sol-gel method was used, a zinc oxide layer was coated onto the sepiolite surface in the form of a gel with a relatively large load and uniform distribution. When the composite was prepared by the hydrothermal reduction method, zinc ions were loaded onto the sepiolite surface in the form of crystals and then oxidised to form unevenly distributed zinc oxide particles. When the composite material was prepared by the hydrolytic precipitation method, zinc ions precipitated into zinc hydroxide species attached to the sepiolite surface and then decomposed into zinc oxide species under high-temperature conditions. Therefore, the loaded particles exhibited microspherical shapes and a relatively large load with an uneven distribution. The powder sintering method resulted in a composite material with a uniform particle distribution using a simple technique. Finally, when the composite was prepared by the impregnation-reduction method, loose zinc oxide particles were uniformly distributed across the sepiolite surface and partially agglomerated. Therefore, the sol-gel method is superior to the hydrothermal reduction, hydrolytic precipitation, powder sintering, and impregnation-reduction methods.

(3) The photocatalytic degradation of MB by the ZnO/sepiolite composite prepared by the sol-gel method represented a quasi-first-order reaction. In addition, this material demonstrated excellent chemical stability and reusability.

The results of this study revealed that the sol-gel method could be used to prepare photocatalytic composite materials.
with excellent performance characteristics, which was consistent with their good photocatalytic properties observed during pollutant degradation. Thus, these findings can provide a basis for the development, preparation, and application of novel photocatalysts. In future works, the photocatalytic performance of composite materials will be enhanced by controlling the reaction conditions to expand the application range of ZnO/sepiolite composites.

**Data Availability**

The experimental data used to support the findings of this study are available from the corresponding author upon request.

**Conflicts of Interest**

The authors declare no conflicts of interest related to the publication of this study.

**Authors’ Contributions**

Chao Li was responsible for the writing of the original draft, validation, and visualization. Huijuan Li was responsible for the experiment and data curation. Guicheng He was responsible for the supervision. Wenyuan Wu was responsible for the cartography and documentation.

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