

Microwave Synthesis and Photocatalytic Properties of CeVO₄/FeVO₄ Nanocomposites

Su Chen, Shuibin Yang, Dingjin Fan, Xuehong Liao*

Hubei Key Laboratory for Processing and Application of Catalytic Materials, College of Chemical Engineering, Huanggang Normal University, Huanggang, China
Email: liaoxuehong@sohu.com

Received 13 April 2016; accepted 24 May 2016; published 27 May 2016

Copyright © 2016 by authors and Scientific Research Publishing Inc.

This work is licensed under the Creative Commons Attribution International License (CC BY).

<http://creativecommons.org/licenses/by/4.0/>



Open Access

Abstract

In this paper, CeVO₄/FeVO₄ nanocomposites were prepared by direct feeding microwave synthesis method with nine water iron nitrate (Fe(NO₃)₃·9H₂O), cerium nitrate hexahydrate (Ce(NO₃)₃·6H₂O) and ammonium metavanadate (NH₄VO₃) as raw material and Sodium Dodecyl Sulfate (SDS) as surfactant. Then X-Ray Diffractometer (XRD) and Scanning Electron Microscopy (SEM) were used to observe the CeVO₄/FeVO₄ nanocomposites. SEM image showed that the as-prepared CeVO₄/FeVO₄ nanocomposites calcined at 773 K is formed of small particles aggregation irregular sheet structure. We studied the photocatalytic activity of the as-prepared samples by using degradation of methyl orange in visible light. The results showed that the photocatalytic activity of CeVO₄/FeVO₄ nanocomposites were very well. It found that when the catalyst calcined at 773 K was 0.10 g, and 0.5 mL hydrogen peroxide joined as well as, pH was 2.0, the degradation ratio of catalyst for methylene orange of 100 mL 5 mg/L reached 98.63% in 40 min.

Keywords

CeVO₄/FeVO₄ Nanocomposites, Microwave Synthesis, Photocatalytic, Methyl Orange

1. Introduction

Environmental pollution, energy crisis and population growth are the three major challenges facing humanity in the 21st century. In recent years, the photocatalytic applications in environmental protection have been paid more and more people's attention.

Many researchers are already visible in the synthesis and catalytic vanadate complex aspects of exploration. These photocatalysts due to a relatively narrow band gap, and visible light photocatalytic activity are widespread

*Corresponding author.

concern. Monoclinic phase vanadate semiconductor photocatalyst material with a narrow band gap in the visible region can exhibit high photochemical activity, and is a promising class of applications highly active photocatalyst material [1]-[8].

In this article, CeVO₄/FeVO₄ complexes are prepared by the method of liquid-phase microwave synthesis, small particle size uniform, the catalytic effect is good. The results show that in 100 mL, 5 mg/L of methyl orange solution, adding CeVO₄/FeVO₄ 0.1 g nanocomposites by 773 K thermostatic heat 2 h after adding 0.5 mL H₂O₂, adjust to pH 2.0, at this time, the degradation rate reaches 98.63%. So, the as-prepared photocatalyst has good photocatalytic performance.

2. The Experiment

2.1. Instruments and Reagents

Microwave oven with 650 W (Sanle general electric corp. Nanjing, China) with refluxing system was used. Powder X-Ray Diffraction (XRD) was used to characterize the sample. Data were collected on a Shimadzu XRD-6100 X-ray diffractometer (Cu K α radiation, $\lambda = 0.15418$ nm). The morphology and size were determined by SEM. The SEM images were recorded on a Quanta 200 FEG field emission scanning electron microscope. Ultraviolet-visible diffuse reflectance spectrum was carried out on a UV-2600 UV—visible spectrophotometer. Lambda10 UV-vis spectrometer (Perkin-Elme Corp, USA) was used for monitoring the absorption spectra of photo-degradation of methyl orange.

All the reagents used were of analytical purity. Doubly distilled water was used throughout the experiments.

2.2. Direct Feeding Microwave Synthesis of CeVO₄/FeVO₄ Nanocomposites

At a molar ratio of 1:1, weighed by calculating the required 2.17 g Ce(NO₃)₃·6H₂O, 1.01 g Fe(NO₃)₃·9H₂O, was dissolved in 50 mL 1 mol/L HNO₃ solution, dispersed and dissolved with ultrasonic waves, mixed uniform for A solution. Weigh 1.17 g NH₄VO₃ was dissolved in 10 mL containing 1 mol/L of NH₃·H₂O, add deionized water to 50 mL, 1.0 g SDS was added to the solution to the dispersion, the dispersion was dissolved by ultrasonic mixing, for B solution. The A, B two solutions were mixed rapidly transferred into 250 mL round bottom flask, fitted with reflux apparatus, with 40% power (total power constant, 30s work cycle, the work 12 s, stop 18 s) microwave irradiation 20 min, allowed to cool to room temperature, pour the supernatant liquid by centrifugation, washed three times with distilled water, dehydrated with acetone once, 60°C vacuum drying 4 h, and then placed in a muffle furnace and heated treated 2 h at different temperatures. Products collected for characterization and photocatalytic experiments.

2.3. Photocatalytic Experiment of CeVO₄/FeVO₄ Nanocomposites

Preparation of 100 mL concentration 5 mg/L of methyl orange solution with dilute nitric acid pH was adjusted to 2.0 by adding a certain amount of CeVO₄/FeVO₄ nanocomposites and 0.5 mL of hydrogen peroxide, and uniformly dispersed ultrasonically in full under dark conditions adsorption, placed under visible light photocatalytic degradation experiments carried out. Sampling every 10 min, the supernatant after centrifugation taken by UV—Vis absorbance spectroscopy and observe the color change of the solution until the solution is completely faded or color does not change until the absorbance. Finally, according to the change in absorbance to calculate the degradation rate of the solution, the calculation formula is as follows:

$$D_t\% = \frac{(A_0 - A_t)}{A_0} \times 100\%$$

In this equation: D_t for methyl orange solution in the presence of catalyst through degradation under visible light irradiation time rate after t , A_0 as the catalyst without the absorbance of methyl orange solution when irradiated with visible light, A_t as catalyst by the absorbance of the methyl orange solution under visible light irradiation time after t .

3. Results and Discussion

Figure 1 is a XRD pattern of the sample. As can be seen, the larger the heat treatment temperature on the sample crystal. When the calcination temperature is increased to 773 k when, XRD diffraction peaks of the sample

sharp, crystal has a very complete explanation.

Figure 2 is a scanning electron micrograph of the sample prepared, it can be seen, most of the catalyst prepared in irregular sheet structure, these sheets are formed by a number of fine particles aggregated.

Figure 3 is a sample of UV—visible diffuse reflectance spectroscopy, can clearly be seen, $\text{CeVO}_4/\text{FeVO}_4$ nanocomposites in the visible range has a strong absorption, is a good light-responsive photocatalyst.

Figure 4 shows the photocatalytic degradation of methyl orange curve. After adding 0.1 g at 773 k fired $\text{CeVO}_4/\text{FeVO}_4$ nanocomposites in 100 mL 5 mg/L methyl orange solution, placed in sunlight degrades, it can be seen only in the 40 min, methyl orange solution the degradation rate of 98.63%, good degradation effect. Well, the photocatalytic performance specifications prepared catalyst.

We also studied the influence factors of the photocatalytic performance, such as heat treatment temperature of the catalyst, the dosage of the catalyst. The results showed the best photocatalytic effect, when the heat treatment temperature is 773 K, the dosage of the catalyst is 1.0 g/L.

4. Conclusions

In this paper, through direct feeding microwave prepared $\text{CeVO}_4/\text{FeVO}_4$ nanocomposites, SEM display 773 K

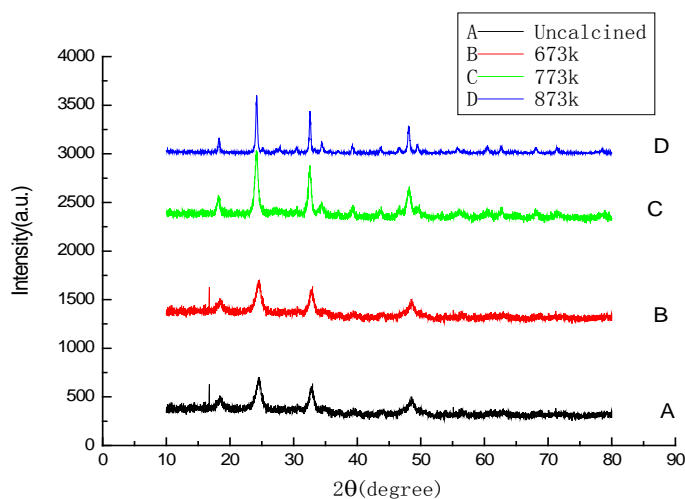


Figure 1. XRD pattern of as-prepared sample.

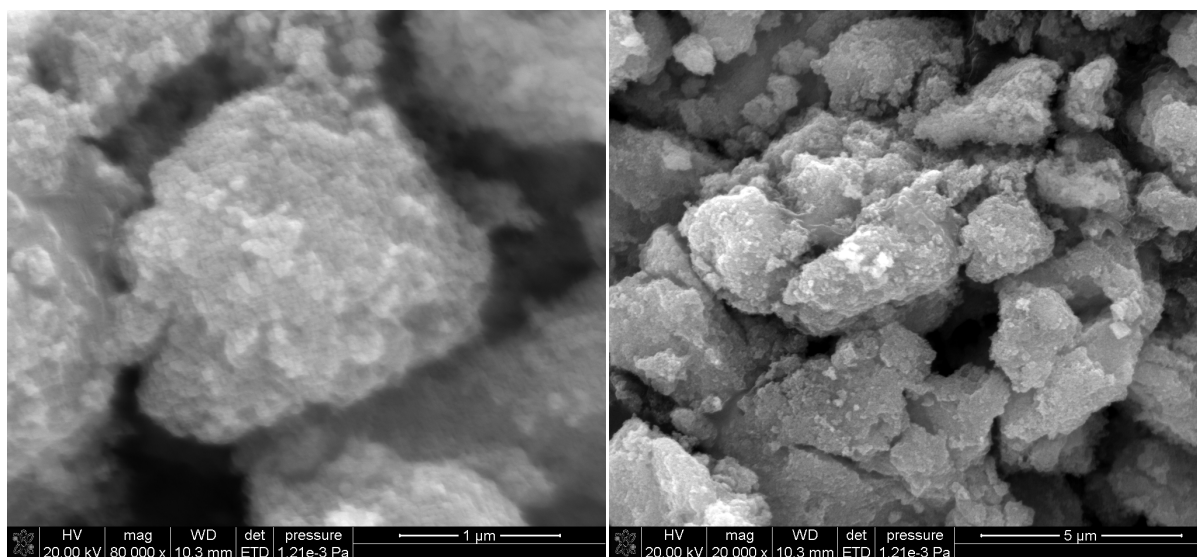


Figure 2. SEM image of as-prepared sample.

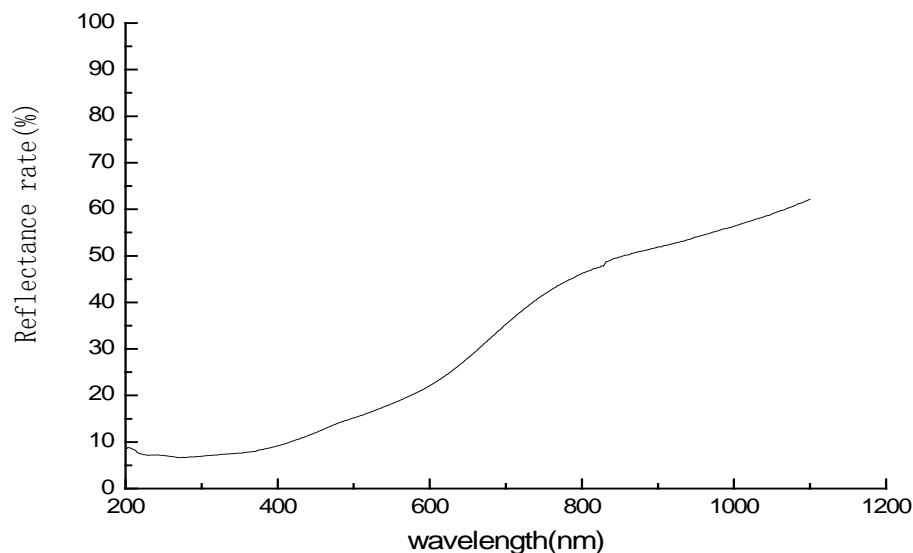


Figure 3. UV-Vis diffuse reflectance spectrum of CeVO₄/FeVO₄ nanocomposite.

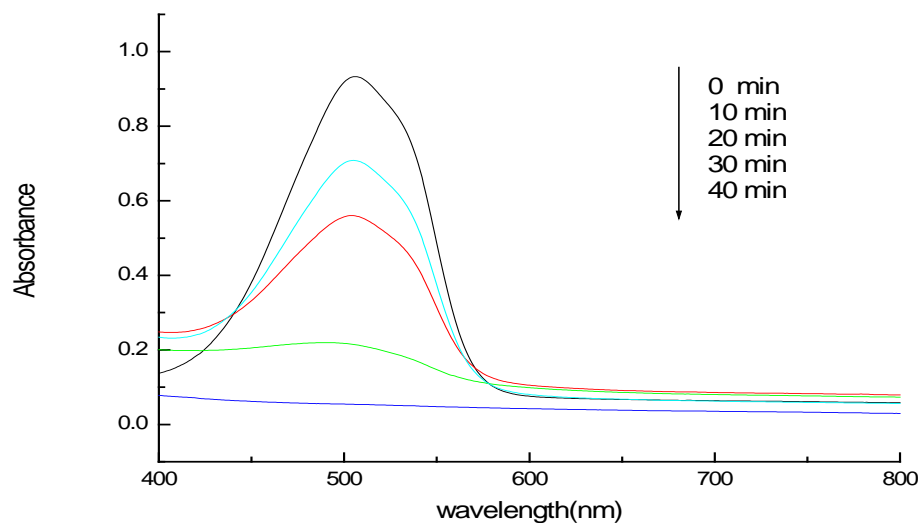


Figure 4. UV-Vis absorption spectra of photo-degradation of methyl orange.

heat treatment of 2 h samples showed irregular flake structure, these layers by some small particles aggregated to form.

The results showed that the CeVO₄/FeVO₄ nanocomposite microwave synthesis had excellent photocatalytic performance. In the 100 mL, 5 mg/L methyl orange solution, adding catalyst 0.1 g, adjusted pH to 2, with 0.5 mL H₂O₂, in the sunlight degradation, the degradation rate could reach 98.63% within 40 min.

References

- [1] da Silva, J.L.F., Ganduglia-Pirovano, M.V. and Sauer, J. (2007) Formation of the Cerium Orthovanadate CeVO₄:DFT + U Study. *Condensed Matter and Materials Physics*, **76**, 3398-3407.
- [2] Jiang, H.Q., Nagai, M. and Kobayashi, K. (2009) Enhanced Photocatalytic Activity for Degradation of Methylene Blue over V₂O₅/BiVO₄ Composite. *Journal of Alloys and Compounds*, **303**, 9-14.
- [3] Wang, Q., Li, H., Liu, J., *et al.* (2015) Preparation of the Photocatalyst Ag₃VO₄ by Hydrothermal Method and Evaluation of Its Visible-Light-Driven Photocatalytic Performance. *Chemical Research*, **26**, 519-523.
- [4] Turkovic, A., Orel, B., Lucic, M., *et al.* (2007) Gixsax Study of Temperature Evolution in Nanostructured CeVO₄ Films. *Solar Energy*, **91**, 1299-1304.

-
- [5] Opara, U., Krasove, B., Orel, A., *et al.* (1999) Structural and Spectroelectrochemical Investigations of Tetragonal CeVO₄ and Ce/V-Oxide Sol-Gel Derived ion-Storage Films. *Solid State Ionics*, **118**, 103-105.
- [6] Konta, R., Kato, H., Kobayashi, H., *et al.* (2003) Photophysical Properties and Photocatalytic Activities under Visible Light Irradiation of Silver Vandates. *Physical Chemistry Chemical Physics*, **5**, 3061-3065.
- [7] Wang, N., Chen, W., Li, L., *et al.* (2009) Synthesis and Magnetic Properties of CeVO₄ Nanorods. *Materials Review*, **23**, 23-26.
- [8] Watanabe, A. (2000) Highly Conductive Oxides, CeVO₄, Ce_{1-x}M_xVO_{4-0.5x}(M= Ca, Sr, Pb) and Ce_{1-y}Bi_yVO₄ with Zircon-Type Structure Prepared by Solid-State Reaction in Air. *Journal of Solid State Chemistry*, **153**, 174-179.
<http://dx.doi.org/10.1006/jssc.2000.8773>