

Microwave Synthesis and Photoluminescence Properties of $\text{CaMoO}_4 : \text{Eu}_{0.1}^{3+}$ Nanocomposites

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How to cite this paper: Li, Q.C., Zeng, X.M., Yang, S.B. and Liao, X.H. (2017) Microwave Synthesis and Photoluminescence Properties of $\text{CaMoO}_4 : \text{Eu}_{0.1}^{3+}$ Nanocomposites. *Open Journal of Composite Materials*, 7, 99-104.

<https://doi.org/10.4236/ojcm.2017.72006>

Received: February 27, 2017

Accepted: April 27, 2017

Published: April 30, 2017

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Abstract

In this paper, using calcium chloride and sodium molybdate as raw material, polyethylene glycol (PEG2000) as surfactant, the nanocomposites of $\text{CaMoO}_4 : \text{Eu}^{3+}$ were prepared by a direct feeding microwave synthesis method. The as-prepared sample was characterized by X-ray diffraction (XRD), scanning electron micrograph (SEM) and photoluminescence spectrum (PL). The XRD Pattern showed that the samples are scheelite structure of CaMoO_4 . The SEM image showed that the majority of as-prepared sample is a relatively flake structure, and some fine particles attached to it. PL spectra showed that as-prepared samples have strong luminescence properties; it had purity red emission at 615 nm. The effects of different Eu^{3+} ions doping amount and surface active agent on the photoluminescence properties were studied. The results showed that when the molar ratio of Eu^{3+} was 0.10, PEG2000 as surfactant, the luminescence intensity of as-prepared sample was maximum.

Keywords

Calcium Molybdate, Eu^{3+} Ion, Doping, Nanocomposite, Microwave Synthesis, Photoluminescence

1. Introduction

Molybdate has superior optical, electrical, magnetic properties, in terms of scintillator, light soldering, sensors and catalysts has a wide application prospect. Metal molybdate as luminescent materials in important family, due to its excellent luminescence has been widely attention. Molybdate system because of it's in the near UV region has wide and strong charge transfer absorption band. After UV excitation energy can be through non radiative transition is passed to the activator ion. So we often used molybdate as matrix material doped with rare earth

ions prepared in near ultraviolet excitation efficient red phosphor, used for white light emitting diode to arouse people's great interest [1]-[11]. At present, several techniques were used to synthesize $\text{CaMoO}_4:\text{Eu}^{3+}$ red phosphors, such as high temperature solid-state, hydrothermal, sol-gel, chemical co-precipitation, combustion, microwave radiation method and so on.

In this study, we report on a direct feeding microwave synthesis method to synthesize $\text{CaMoO}_4:\text{Eu}^{3+}$ red phosphors. The calcium molybdate as matrix, using Eu^{3+} ion as activator, by changing the concentration of Eu^{3+} ion, the choice of different surfactant, to seek the strongest luminescence.

2. The Experiment

2.1. Synthesis of $\text{CaMoO}_4:\text{Eu}_{0.1}^{3+}$ Nanocomposite

All chemicals were analytical grade and used without further purification. nanocomposites of $\text{CaMoO}_4:\text{Eu}^{3+}$ were prepared by a direct feeding microwave synthesis method. In a typical procedure, at a molar ratio of Ca:Mo:Eu of 1:1:0.1, 1.1 g of CaCl_2 was dissolved in 50 ml of 2% PEG aqueous solution, dispersed and dissolved with ultrasonic waves, add 1 mmol of Eu^{3+} reserve liquid, mixed uniform for A solution. 2.42 g of $\text{Na}_2\text{MoO}_4 \cdot 2\text{H}_2\text{O}$ was dissolved in 50 mL of 2% PEG aqueous solution, the dispersion was dissolved by ultrasonic mixing, for B solution. The A, B solutions were mixed rapidly transferred into 250 ml of flask, then the mixed solution was placed in a microwave refluxing system to react for 20 min with a power microwave radiation of 40% and cool down naturally to the room temperature. Then the precipitate was centrifuged, washed with the deionized water for several times and dried at 60°C in the vacuum for 8 h, The final product was collected for the characterization.

2.2. Characterization

The crystal structure of nanocomposites of $\text{CaMoO}_4:\text{Eu}^{3+}$ was measured by XRD on a Shimadzu XRD-6100 X-ray diffractometer (Cu $K\alpha$ radiation, $\lambda = 0.15418$ nm). The morphology and size of products were determined by SEM. The SEM images were recorded on a Quanta 200 FEG field emission scanning electron microscope. The optical property was obtained by Cary Eclipse fluorescence spectrometer (USA Varian Company).

3. Results and Discussion

3.1. XRD and SEM Analysis

Figure 1 shows the XRD pattern graph of as-prepared sample. The XRD pattern (peak 2θ 18.75, 28.82, 31.36, 34.39, 47.13, 49.37, 54.14, 58.11) showed that the product is the Tetragonal system of scheelite structure of CaMoO_4 (JCPDS File No. 29-0351). The diffraction peak is strong and sharp, which indicates that the sample has a high degree of crystallinity.

Figure 2 shows the SEM image of as-prepared sample. It shows that the majority of the catalyst is a relatively flake structure, parts overlap each other, and some particles attached to it. The size of most of the flakes is 50 to 500 nm.

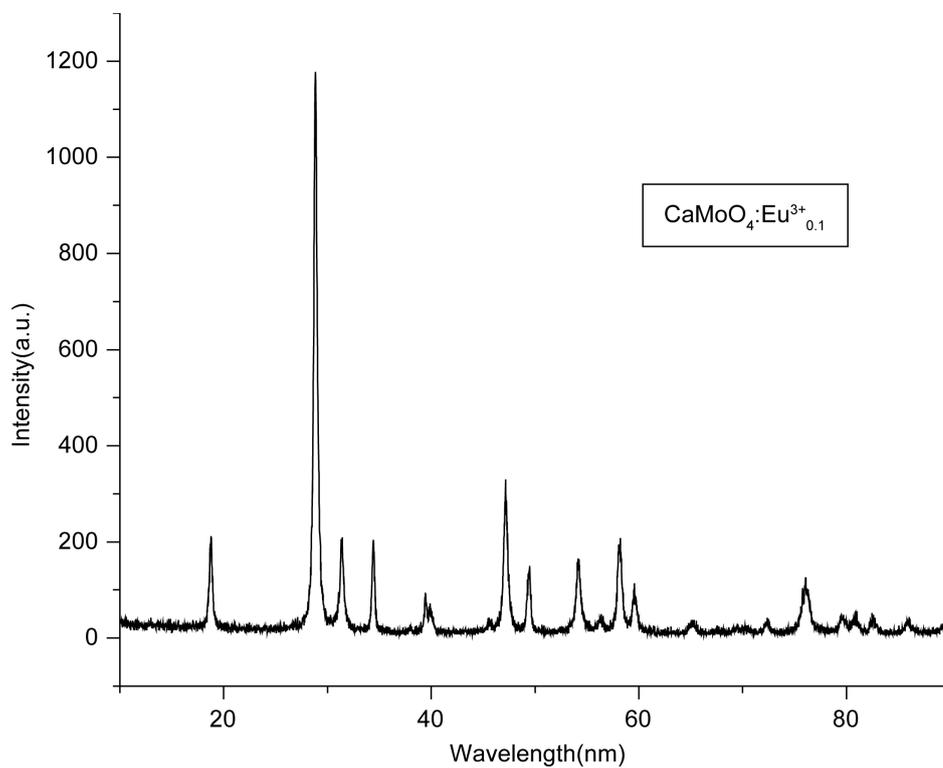


Figure 1. X-ray diffraction pattern of as-prepared samples.

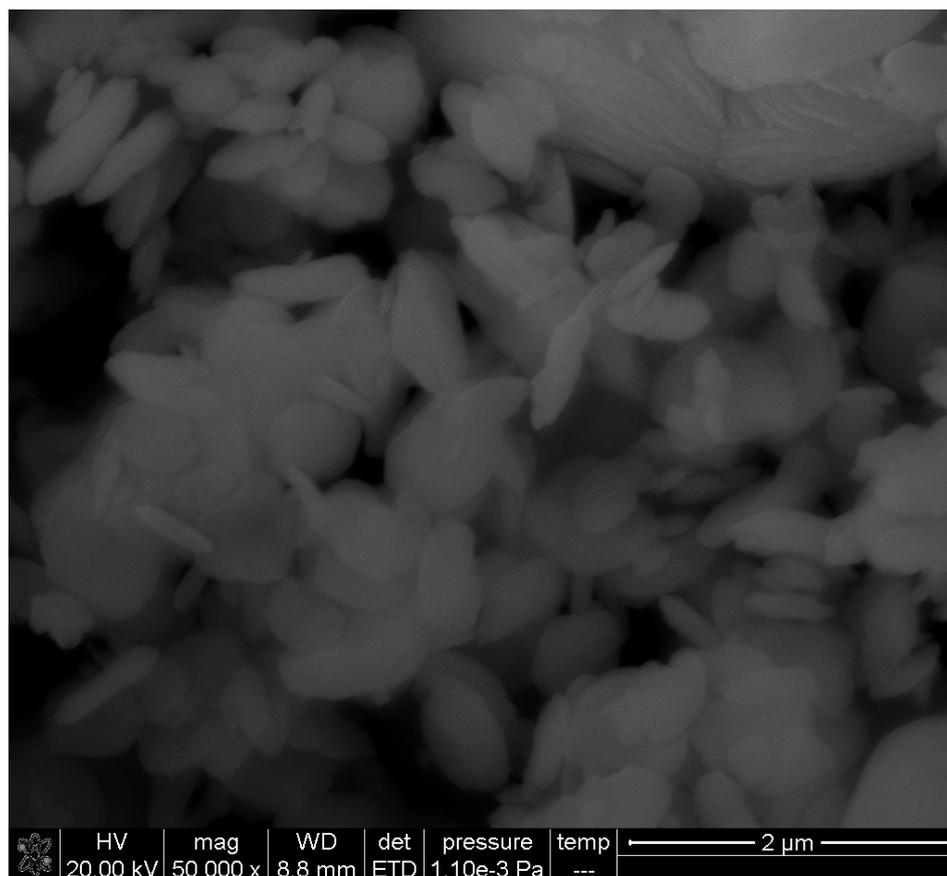


Figure 2. Scanning electron micrograph image of as-prepared sample.

3.2. Photoluminescence Properties of $\text{CaMoO}_4 : \text{Eu}_{0.1}^{3+}$ Nanocomposite

Figure 3 is photoluminescence spectrum of as-prepared sample. The excitation wavelength is 258 nm. It can be seen in the 430 - 450, 590 - 600, 610 - 620, 700 - 710 nm have a certain luminescence, which is the strongest at 615 nm. 430 - 450 nm belongs to calcium molybdate luminescence. The luminescence mechanism of calcium molybdate is due to transition induced by electrons in MoO_4^{2-} ion within the groups, belonging to the intrinsic emission. 516 nm has a very high peak, this is a frequency doubling peak of excitation light. At 590 - 600 nm, 610 - 620 nm and 700 - 710 nm, luminescence by Eu^{3+} (${}^5\text{D}_0 \rightarrow {}^7\text{F}_1$), (${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$), (${}^5\text{D}_0 \rightarrow {}^7\text{F}_4$) electronic energy level transition, respectively. Among them, 615 nm (${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$) electric dipole transition emission is the main, it has high emission efficiency, the strongest luminescence properties, and pure red light. So, as-prepared $\text{CaMoO}_4 : \text{Eu}_{0.1}^{3+}$ nanocomposite is an ideal red light-emitting material.

3.3. Effect of Eu^{3+} Doping on Photoluminescence Properties

The effect of Eu^{3+} ions doping on the luminescence properties was studied. **Figure 4** is photoluminescence spectra of samples with different Eu^{3+} doping. It can be seen that when the doping amount of Eu^{3+} ion is less than 0.10 (molar ratio), the luminescence intensity of $\text{CaMoO}_4 : \text{Eu}^{3+}$ composite increases with the increase of the amount of Eu^{3+} ions. When the doping amount of Eu^{3+} ion is 0.10, the luminescence properties are the best. When the doping amount of Eu^{3+} ion is more than 0.10, the amount of Eu^{3+} ions doping will continue to increase, the intensity of the luminescence will decrease obviously due to the concentration of the particle, the luminescence intensity of the $\text{CaMoO}_4 : \text{Eu}^{3+}$ nanocomposite decreases with the increase of Eu^{3+} doping amount.

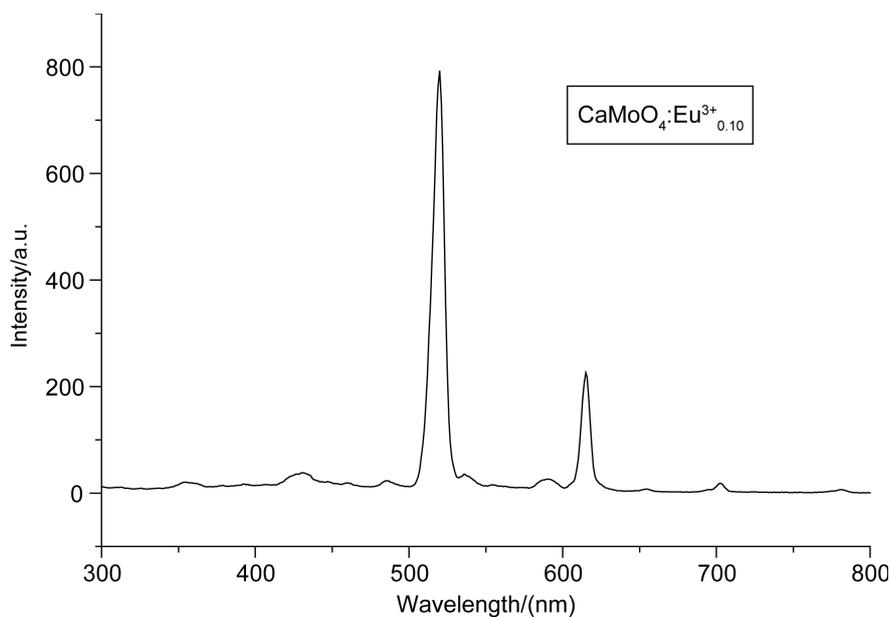


Figure 3. Photoluminescence spectrum of as-prepared sample.

3.4. Effect of Different Surfactants on Photoluminescence Properties

We also investigated the effect of different surfactants on the luminescence properties. **Figure 5** is photoluminescence spectra of samples with different surfactants. It can be seen that the nonionic surfactant PEG is the best, next is the anionic surfactant sodium dodecyl sulfate (SDS), the cationic surfactant cetyltrimethyl ammonium bromide (CTAB) is worst.

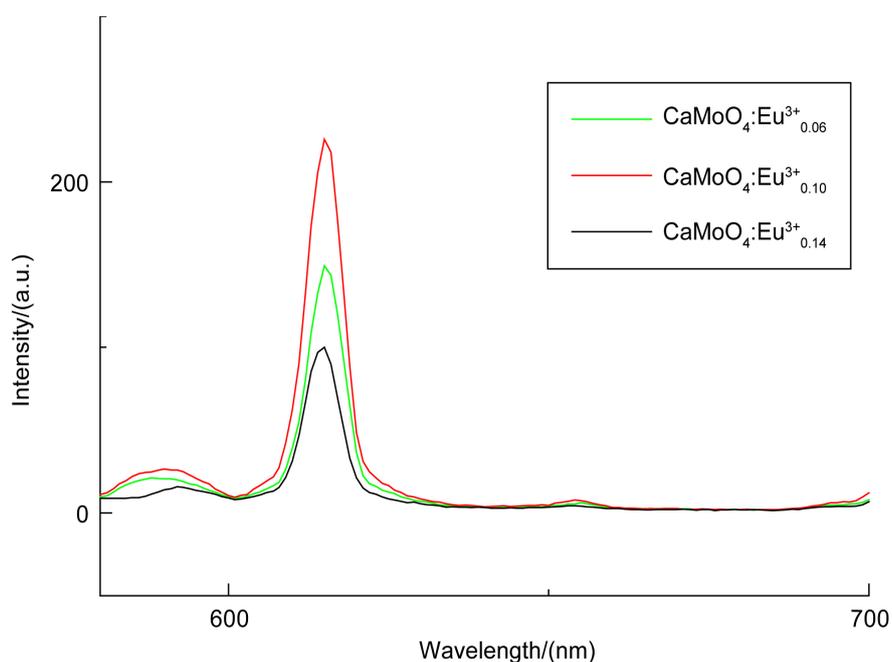


Figure 4. Effect of Eu^{3+} doping on photoluminescence properties.

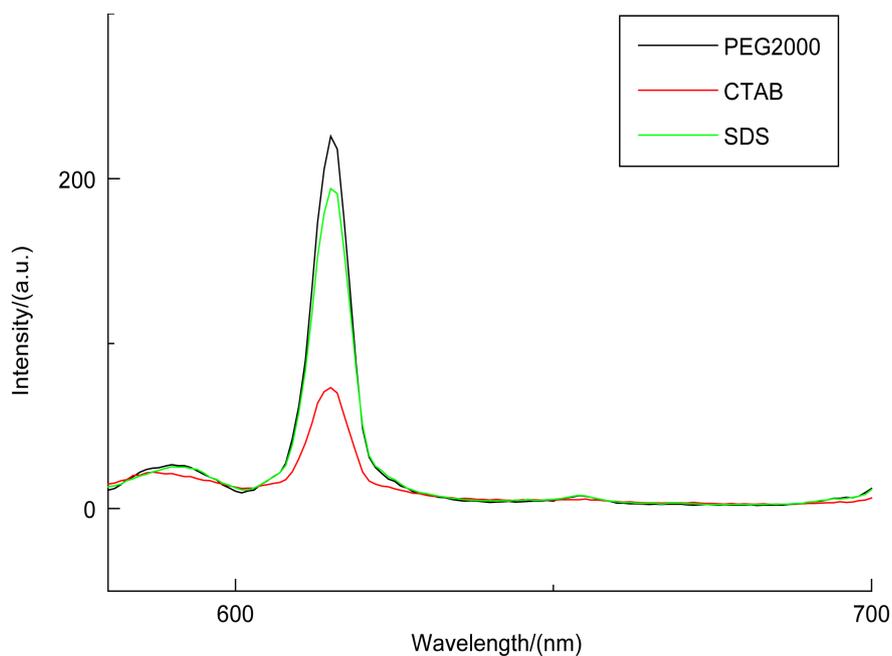


Figure 5. Effect of different surfactants on photoluminescence properties.

4. Conclusions

CaMoO₄:Eu³⁺ nanocomposites were successfully prepared by a direct feeding microwave synthesis method. This method is simple.

As-prepared samples have strong luminescence properties; it had purity red emission at 615 nm. When the molar ratio of Eu³⁺ was 0.10, PEG2000 as surfactant, the luminescence intensity of as-prepared sample was maximum.

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