The contribution of Eu$^{3+}$ doping concentration on the modulation of morphology and luminescence properties of InVO$_4$:Eu$^{3+}$

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Nanostructures of InVO$_4$ and InVO$_4$:Eu$^{3+}$ were synthesized through a hydrothermal method with a post-annealing process. The morphology and luminescence properties of InVO$_4$:Eu$^{3+}$ can be modulated by Eu$^{3+}$ ion doping concentration. SEM and (HR)TEM studies indicated that different sizes of nanomaterials were obtained when the Eu$^{3+}$ ion concentration ranged from 2 mol% to 30 mol%, and middle-concave nanodisks or nanoparticles with different sizes were obtained with the Eu$^{3+}$ ion concentration ranging from 35 mol% to 45 mol%. Luminescence property studies indicated that the photoluminescence emission originated both from VO$_4$$^{3-}$ and Eu$^{3+}$, and the emission of VO$_4$$^{3-}$ was blue shifted, affected by the doped Eu$^{3+}$ ions. The CIE chromaticity diagram of Eu$^{3+}$-doped indium vanadate exhibited green, blue-green, red, giving especially white light emission.

1 Introduction

Nowadays, materials on the mesoscopic scale are being paid more and more attention. Nano-size materials with uniform morphology usually have different special characteristics, such as their magnetism, optics, conductivity and so on, due to small size effects and the specific effect of large surface area. As we all know, InVO$_4$ is a semiconductor with a narrow band gap. The photoluminescence properties of InVO$_4$ have previously been studied. Since the morphology of nanomaterials may have some important effects on their optical properties, it is important to investigate various morphologies of InVO$_4$ in detail to obtain outstanding luminescent nanomaterials. However, nano-size InVO$_4$ has been rarely investigated. Ai et al. synthesized nanocrystalline InVO$_4$ hollow microspheres with an aerosol flow synthetic method, and other nano-sized indium vanadates have been synthesized by the sol-gel method and hydrothermal processes. In spite of these methods for synthesis of indium vanadate nano-/micro-materials, the size distribution and regularity of these materials can still be improved. The diversity of regular and dispersed nanostructures for indium vanadates still needs to be expanded to meet their desired potential applications.

In addition, white-LEDs have attracted much attention in recent decades because of their properties of high brightness, long fluorescence lifetimes and environmental friendliness. Lanthanide ions are usually codoped with each other or with transition metals in a host lattice to obtain white light materials. It is very important to find a suitable host for lanthanide luminescent centers to give white emission. In this work, we found that different emission colors including white light could be achieved from different concentrations of Eu$^{3+}$-doped InVO$_4$ nanomaterials. First, Eu$^{3+}$-doped indium vanadates were synthesized through a hydrothermal process with sodium citrate as a coordination agent. Nanostructures of the as-obtained InVO$_4$:Eu$^{3+}$ with different europium ion doping concentrations were investigated. The morphology and size of the nano-indium vanadates varied regularly with the variation of the Eu$^{3+}$ doping concentration. Different sizes of nanoparticles, middle-concave nanoparticles and middle-concave nanodisks of InVO$_4$:Eu$^{3+}$ were obtained. Second, the photoluminescence properties of InVO$_4$:Eu$^{3+}$ were studied to reveal the different emission spectra with different Eu$^{3+}$ doping concentration. The CIE chromaticity diagram of Eu$^{3+}$-doped indium vanadate exhibited green, blue-green, red, giving especially white light emission, making this material a suitable candidate for optical LEDs.

2 Experimental section

2.1 Characterization

X-ray diffraction (XRD) patterns were collected on an X-ray diffractometer (Bruker Axs D2 PHASER diffractometer) with Cu Kα radiation ($\lambda = 1.5405$ Å). SEM images were recorded on a scanning electron microscope (SEM, JSM-5601). TEM images were recorded on a transmission electron microscope (TEM JSM 2100). The photoluminescence spectra were recorded on a PL3-211-P spectrometer (HORIBA Jobin Yvon, America) and a 450W xenon lamp was used as the excitation source. The
luminescence quantum efficiency (Φ) was obtained by using BaSO₄ as a reference.

2.2 Synthesis

General method for synthesis of InVO₄: 10 mL 0.1 mol L⁻¹ NH₄VO₃ aqueous solution was introduced dropwise to a vigorously stirred 10 mL aqueous solution of 1 mmol In(NO₃)₃·5H₂O. After addition, the mixture was stirred for 10 min, and a certain amount of sodium citrate was added to the suspension. The as-obtained yellow colloidal precipitate was transferred to a 50 mL autoclave, sealed, and heated at 150 °C for 15 h. After completion of the reaction time, the autoclave was then cooled to room temperature naturally. The products were washed three times with distilled water and centrifuged for 3 min at 10 000 rpm. The washed powders were dried under vacuum at 60 °C. The final products were obtained by annealing at 900 °C in air for 6 h and used for further investigation. For different doping concentrations in InVO₄ₓEu³⁺, the same method was used with starting materials containing different proportions of In(NO₃)₃·5H₂O and Eu(NO₃)₃·6H₂O.

3 Results and discussion

3.1 Synthesis and structure

The nanostructures of indium vanadate or Eu³⁺-doped indium vanadate were synthesized by a hydrothermal method with a post annealing process. Indium(III) nitrate hydrate and ammonium vanadate were used as starting materials with sodium citrate as a complexing agent. The sodium citrate played an important role in the formation of the indium vanadate nanostructures. Control experiments without sodium citrate were carried out and irregularly shaped particles were obtained.

Fig. 1 shows the XRD patterns of the annealed hydrothermal products of InVO₄ and InVO₄ doped with various concentrations of Eu³⁺. It indicates that the XRD of InVO₄ shown in Fig. 1a can be easily indexed to orthorhombic InVO₄ JCPDS: 71-1689. As the Eu³⁺ doping concentration increased, the XRD patterns showed a slight change to indicate Eu³⁺ was well-doped in the crystal lattices of InVO₄, whose special obvious peaks are illustrated in Fig. 1. For lower doping concentrations, no obvious differences were found in the XRD patterns. When the doping-concentration of Eu³⁺ ions is higher than 10 mol% (Fig. 1b–d) the InVO₄ phase still remains, but the full width at half maximum of these peaks broadens.

The morphologies of InVO₄ and InVO₄xEu³⁺ were characterized by SEM and (HR)TEM. The results indicated that uniform and dispersed nanostructures were obtained through this method. The SEM micrograph of as-obtained InVO₄ is shown in Fig. 2a. Regular nanoparticles with a diameter around 200 nm were formed. Fig. 3 shows the (HR)TEM and EDX results of as-obtained InVO₄. The TEM image of InVO₄ shown in Fig. 3a displays the same uniform nanoparticles as those in the SEM image (Fig. 2a). The HRTEM image (Fig. 3b) of the nanoparticles shows lattice fringes with interplanar spacing of 0.271 nm that corresponds to the (1 1 2) plane of the orthorhombic crystal structure of InVO₄. The EDX in Fig. 3c clearly showed the composition of the InVO₄ nanoparticles.

Fig. 2b–g show the SEM images of the InVO₄ host doped with 2, 5, 10, 15, 20 and 30 mol% of Eu³⁺ ions. They are all uniform and dispersed nanoparticles. It is obvious that as the Eu³⁺ ion concentration increased from 2 mol% to 30 mol%, the InVO₄xEu³⁺ particle sizes decreased. The diameters of the
nanoparticles with different Eu$^{3+}$ ion concentrations are given in Table 1. Diameters of the particles decreased from 200 to 50 nm with increased Eu$^{3+}$ ion concentration. This nanoparticle diameter trend can be attributed to the large difference in the ion radius between indium(III) ions and europium(III) ions, which is 0.8 Å for In$^{3+}$ and 0.947 Å for Eu$^{3+}$.$^{18}$ The nanoparticle sizes were proven to be influenced by the nucleation and growth rates in the formation process of the nanostructures.$^{9}$ The presence of large ion radius Eu$^{3+}$ lowered the potential barrier for the formation of nuclei and slowed the growth rates which resulted in a decrease in nanoparticle size. This is in good accordance with the nanoparticle diameter trend seen in the SEM results. In addition, slight aggregation of the calcined nano-InVO$_4$ can also be observed (Fig. 2a), while the increased doping of Eu$^{3+}$ ions made it better and more dispersed nanoparticles were obtained in Fig. 2b–g.

Fig. 4 and 5 show the SEM and TEM images of the InVO$_4$ host doped with 35, 40 and 45 mol% of Eu$^{3+}$ ions. It is observed that when the Eu$^{3+}$ ion doping concentrations are higher than 35 mol%, the images of the InVO$_4$:Eu$^{3+}$ do not show nanoparticles any more. Both their size and morphology changed. The morphologies and diameters of the nanostructures with different Eu$^{3+}$ ion concentrations are collected in Table 1. InVO$_4$:35% Eu$^{3+}$ displays special nanodisks with diameters around 180 nm. Fig. 4a and b show the SEM and TEM images of

<table>
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<tr>
<th>Compounds</th>
<th>Morphology</th>
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<tr>
<td>InVO$_4$</td>
<td>Nanoparticle</td>
<td>200</td>
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<tr>
<td>InVO$_4$:2% Eu$^{3+}$</td>
<td>Nanoparticle</td>
<td>180</td>
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<td>Middle-concave particle</td>
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<tr>
<td>InVO$_4$:45% Eu$^{3+}$</td>
<td>Middle-concave particle</td>
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the InVO₄:35% Eu³⁺ nanodisks, respectively. With careful observation we can find that these nanodisks are middle-concave, and the thickness of the edge of a nanodisk is about 50 nm. The morphologies of InVO₄ doped with 40 mol% (Fig. 5a and b) and 45 mol% Eu³⁺ (Fig. 5c and d) are novel middle-concave nanoparticles with diameters of 230 nm for 40 mol% Eu³⁺ and 260 nm 45 mol% Eu³⁺. The EDX shown in Fig. 4c clearly indicated the composition of the InVO₄: Eu³⁺ nanoparticles.

Generally, the growth process of nanocrystals includes two main steps: an initial nucleating stage and a crystal growth stage. In the first stage, the formation of seeds is crucial for further growth of the crystals (second stage). The morphologies of the final crystals are usually determined by the very early seed formation. So when the doping concentration of Eu³⁺ is lower than 30 mol%, In³⁺ dominated the initial stage of the crystal growth and different sizes of nanoparticles were obtained (as shown in Fig. 2b–g). When the doping concentration of Eu³⁺ is higher than 35 mol%, the Eu³⁺ probably had such a huge
impact on the initial stage that the morphology of the final crystals turned out to be middle-concave nanodisks or nanoparticles. As mentioned before, the ion radius of Eu$^{3+}$ is much bigger than that of In$^{3+}$, so the first nucleation stage would be changed by a higher doping concentration of bigger Eu$^{3+}$ ions even under the same experimental conditions. So it is understandable that InVO$_4$ with Eu$^{3+}$ doping concentrations higher than 35 mol% shows different morphologies. At the same time, we should notice that when the Eu$^{3+}$ doping concentration is higher than 35 mol%, the sizes of the nanostructures increase with the Eu$^{3+}$ doping concentration increasing. This is opposite to the Eu$^{3+}$ lower doping concentration, where nanoparticle size decreased with Eu$^{3+}$ concentration increasing (Fig. 2b–g). From the comparison of Fig. 4 and 5, we can see that InVO$_4$ with a doping concentration of Eu$^{3+}$ from 35 mol% to 45 mol% displays middle-concave nanostructures, and the size of the nanostructures increases from 180 nm to 260 nm. From this point we can speculate that the higher Eu$^{3+}$ doping concentrations (more than 35 mol%) in InVO$_4$ have an effect on both morphology and size of the nanostructures. When Eu$^{3+}$ doping concentration is higher than 50%, no regular nanostructures were obtained.

3.2 Optical properties

The room temperature photoluminescence excitation and emission spectra of InVO$_4$:30% Eu$^{3+}$ are shown in Fig. 6. The excitation spectrum of InVO$_4$:30% Eu$^{3+}$ monitored at 615 nm (Fig. 6a) exhibits an intense wide band centered at 316 nm, which is attributed to the charge transfer from the oxygen atoms to the central vanadium atom inside the VO$_4^{3-}$ anionic group. There are several groups of sharp peaks between 360 and 400 nm which belong to the Eu$^{3+}$ intra-4f transitions. The emission spectrum monitored at 316 nm (Fig. 6b) includes different lines of $^5D_0 \rightarrow ^7F_J$ ($J = 1, 2, 3, 4$) multiplet transitions, in which the very strong lines ($^5D_0 \rightarrow ^7F_2$) of hypersensitive transitions at 615 nm dominate the spectrum. Different transitions corresponding to the excitation and emission peaks are also illustrated in Fig. 6b. They are the typical well-known Eu$^{3+}$ ion emission, which will not be discussed in detail. The comparison of the emission spectra of different Eu$^{3+}$-doped InVO$_4$ is the important issue that needs to be noticed here. Fig. 7a depicts the photoluminescence emission spectrum of InVO$_4$ doped with different Eu$^{3+}$ ion concentrations monitored at 316 nm. As we all know, InVO$_4$ is a self-activating phosphor. When Eu$^{3+}$ is lower than 2 mol%, the spectra are dominated by InVO$_4$ emission at 524 nm, which originates from the VO$_4^{3-}$ charge transfer transition. The nanoparticles exhibited a blue-green emission under short UV irradiation. Meanwhile, in InVO$_4$:5% Eu$^{3+}$ and InVO$_4$:8% Eu$^{3+}$ emission spectra, the InVO$_4$ emission peak in 524 nm is reduced and a new peak at 434 nm appears, and at the same time the Eu$^{3+}$ emission enhances. So these two nanomaterials, especially InVO$_4$:8% Eu$^{3+}$, display obvious white light emission. The change of the InVO$_4$:Eu$^{3+}$ emission indicated that the Eu$^{3+}$ was well doped in the InVO$_4$ host. The increasing doping concentration of Eu$^{3+}$ made the emission of InVO$_4$ blue shift gradually (from 524 nm to 434 nm).
because the much larger ions of Eu$^{3+}$ occupied some of the crystal lattices instead of the smaller In$^{3+}$ ions. And in the InVO$_4$:10% Eu$^{3+}$ emission, the InVO$_4$ emission blue shifted to 434 nm completely, while the emission of Eu$^{3+}$ gradually enhanced and dominated the emission spectrum instead of InVO$_4$. Then the emission of InVO$_4$ disappeared gradually with Eu$^{3+}$ ion doping concentration increasing. The emission intensity of Eu$^{3+}$ reached a maximum at 30% doping concentration. The concentration quenching effect occurs at Eu$^{3+}$ concentrations higher than 30%, which is caused by the effect of energy exchange between Eu$^{3+}$ ions as the distance between them decreases with increasing Eu$^{3+}$ ion concentrations. The luminescence quantum efficiency ($\Phi$) which fundamentally determined the luminescence properties of the materials was studied. The experimental value ($\Phi$) of the InVO$_4$: 10% Eu$^{3+}$ is 3.04%, while the value ($\Phi$) of InVO$_4$: 30% Eu$^{3+}$ is 15.38% with the same experimental conditions. It can be seen that the luminescence increased about 5 times when Eu$^{3+}$ doping concentration increased from 10% to 30%.

It is interesting to find that the emission wavelength of InVO$_4$:5% Eu$^{3+}$ and InVO$_4$:8% Eu$^{3+}$ almost covers the whole visible region. The CIE chromaticity coordinates are $x = 3937$ $y = 3593$ for InVO$_4$:5% Eu$^{3+}$ and $x = 0.3195$ $y = 0.3181$ for InVO$_4$:8% Eu$^{3+}$ as shown in Fig. 7b, which could make these materials suitable candidates for white LEDs. Fig. 7b shows the CIE chromaticity diagram for InVO$_4$:Eu$^{3+}$ phosphors. The CIE chromaticity coordinates of InVO$_4$:xEu$^{3+}$ have been clearly marked with black spots in Fig. 7b corresponding to the emission of InVO$_4$:xEu$^{3+}$ ($x = 1$–40%) shown in Fig. 7a. The emission colors are green, blue green, white and red for different Eu$^{3+}$ ion concentrations. In this case, we can modulate the emission color region through InVO$_4$ doping with different concentrations of Eu$^{3+}$, which made the nanomaterials of InVO$_4$:Eu$^{3+}$ excellent new materials for optoelectronic applications.

4 Conclusion

Uniform nanostructures of InVO$_4$:xEu$^{3+}$ were obtained in this work. The InVO$_4$:xEu$^{3+}$ products are different sizes of nanoparticles when the Eu$^{3+}$ concentrations range from 0 mol% to 30 mol%, and middle-concave nanodisks or nanoparticles with different sizes when the Eu$^{3+}$ concentrations range from 35 mol% to 45 mol%. The morphology difference is related to the large difference in ion radius between In$^{3+}$ ions and Eu$^{3+}$ ions. Emission colors of InVO$_4$:xEu$^{3+}$ including green, blue green, white and red light were obtained by modulating the Eu$^{3+}$ ion doping concentration. When the Eu$^{3+}$ doping concentration is less than 20%, the emissions originate both from Eu$^{3+}$ and VO$_4^{3-}$, of which the emission of the InVO$_4$:5% Eu$^{3+}$ and InVO$_4$:8% Eu$^{3+}$ almost cover the whole visible region and exhibit white light. The emission intensity of Eu$^{3+}$ ions reached a maximum at 30% doping concentration and InVO$_4$:30% Eu$^{3+}$ also has the smallest size in morphology, which can be attributed to the quantum confinement effect of the matrix where the Eu$^{3+}$ ions were trapped. Research about micro/nanostructures of indium compounds is still in progress in our lab.

Conflicts of interest

There are no conflicts to declare.

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References

2. (a) J. Ge, J. D. Lei and R. N. Zare, Nat. Nanotechnol., 2012, 7, 428–432; (b) D. Kowalski, D. Kim and P. Schmuki, Nano Today, 2013, 8, 235–264.