

Synthesis of Controlled Size GdPO₄·H₂O Nanorods

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Abstract

Due to the wide nanoparticles application, the synthesis of nanoparticles and the study of their properties are of great interest to scientists. Special attention is paid not only to particle size but also to their morphology, since the physical properties depend on the morphology of the nanoparticles. The Eu doped GdPO₄ has luminescent properties. These particles can be applicable in biomedicine, for example, in biological labels, biological images and drug delivery [1,2].

In this study Eu doped GdPO₄·H₂O samples were prepared by hydrothermal method. Our goal is to investigate the dependence of morphology and size of GdPO₄·H₂O nanoparticles on the synthesis conditions. One parameter was changed during each synthesis. The synthesis of Gd_{0.85}Eu_{0.15}PO₄·H₂O was performed by changing the solvent amount (H₂O) in reactor of hydrothermal autoclave, the amount of reagents was kept the same. The pH of the solution adjusted with adding nitric acid. Samples were analysed by X-ray diffraction analysis and scanning electron microscopy. The luminescence properties also have been measured.

Experimental

The starting materials were Gd(NO₃)₃·6H₂O, (99.9%, Glentham), Eu(NO₃)₃·6H₂O (99.9%, Alfa Aesar), C₄H₆O₆ (99.5%), NH₄H₂PO₄. All materials were dissolving separately in distilled water. Gd(NO₃)₃·6H₂O, Eu(NO₃)₃·6H₂O and C₄H₆O₆ solutions were mixed together for 30 min. at 50 °C. NH₄H₂PO₄ solution were added slowly to prepared Gd(NO₃)₃·6H₂O, Eu(NO₃)₃·6H₂O and C₄H₆O₆ solution and mixed for 30 min. at 50 °C. The pH was adjusted by adding NH₄OH or HNO₃.

The hydrothermal syntheses were performed in Berghof High-pressure reactor BR, temperature controller BTC-3000, heating equipment BMH Heating jacket and IKA RH Digital mixer.

The temperature was slowly increased to 180 °C and maintained throughout the syntheses.

Obtained mixture was centrifuged at 7000 rpm, and washed with distilled water. This procedure was repeated 4 times, and 2 times this procedure was repeated using C₂H₅OH for washing. The precipitate was left to dry in air at 70 °C.

X-ray diffraction (XRD) analysis was performed using a Rigaku MiniFlex II. For particles morphology and size analyses the Hitachi SU-70 scanning electron microscope (SEM) was used.

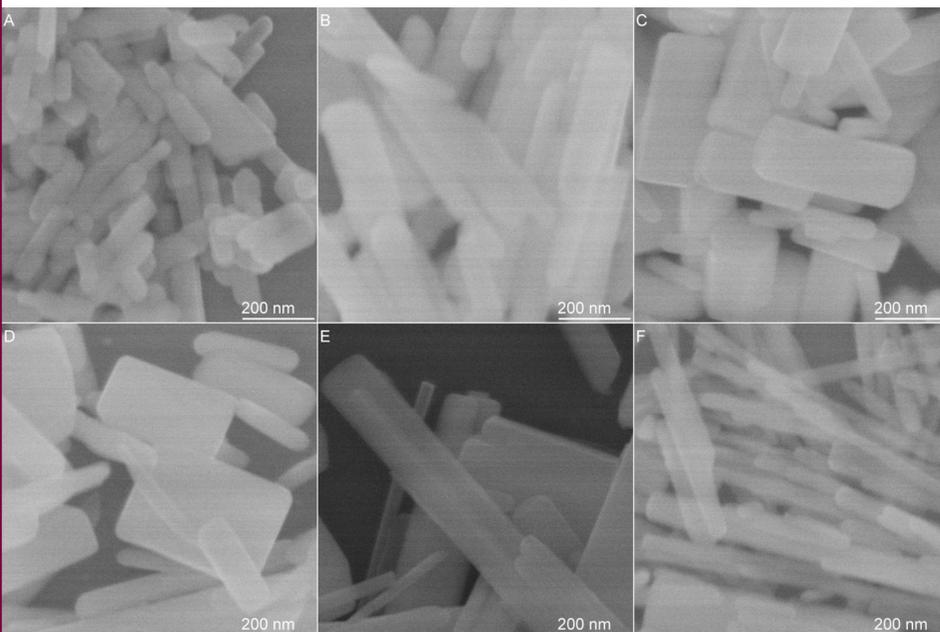


Fig. 1. SEM images of GdPO₄·H₂O synthesized using different amount of solvent (H₂O) in reactor of hydrothermal autoclave: (A) V=10ml, (B) V=20ml, (C) V=30ml, (D) V=40ml, (E) V=50ml, (F) V=60ml.

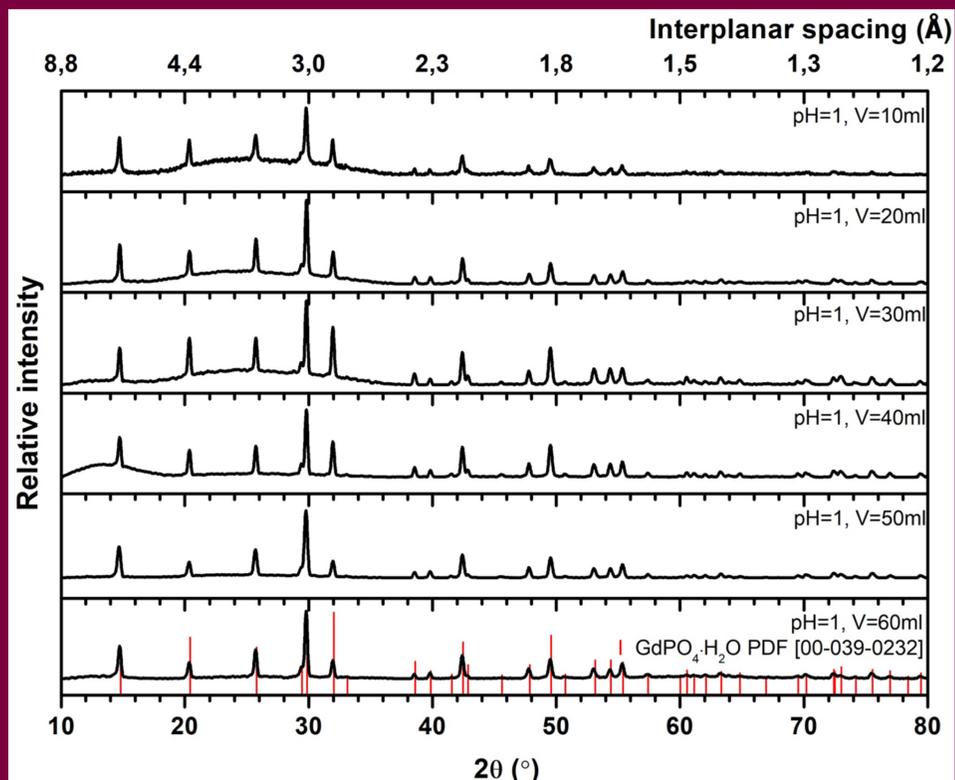


Fig. 2. XRD patterns of GdPO₄·H₂O samples synthesized using different amount of solvent (H₂O) in reactor of hydrothermal autoclave.

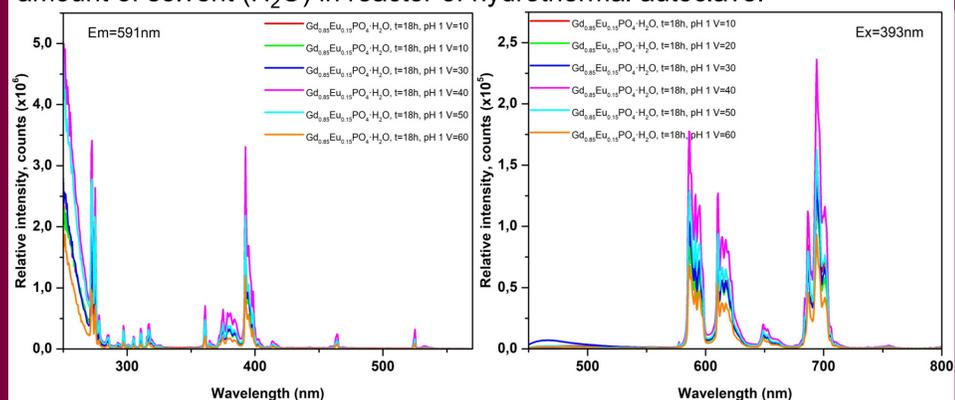


Fig. 3. Excitation (λ_{em} =591 nm) and emission (λ_{ex} =393 nm) spectra of GdPO₄·H₂O samples synthesized using different amount of solvent (H₂O).

Conclusions

Eu doped GdPO₄·H₂O nanoparticles synthesized using different amount of solvent (H₂O) were nanorods shape and not uniform in size. With increasing amount of solvent, the longer nanorods were obtained. The diameter of the nanorods increased by increasing the volume of the solvent, but with the volume of the solvent 50ml and 60ml the diameter does not increase.

References

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2. H. Song, L. Zhou, L. Li, F. Hong, X. Luo. Hydrothermal synthesis, characterization and luminescent properties of GdPO₄·H₂O:Tb³⁺ nanorods and nanobundles. *Materials Research Bulletin* Vol. 48, 2013, pp 5013–5018.