

Synthesis of Fe-doped CeO₂ Nanoparticles Prepared by Solgel Method

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Abstract

Nanomaterials have achieved remarkable technological advances in bulk materials due to their excellent physical, chemical and biological properties. cerium oxide (CeO₂) nanostructured doped with Fe ions is attractive due to improvement in redox properties, transport property and surface-to-volume ratio. In this research, Fe-doped CeO₂ nanoparticles (NPs) were prepared by simple solgel method. The as-synthesized and annealed samples were studied by transmission electron microscopy (TEM), field emission scanning electron microscopy (FESEM) and x-ray diffraction (XRD). The XRD result revealed the cubic crystal structure of Fe-doped CeO₂ NPs. The FESEM images showed that the uniformity of the NPs increase with increasing calcination temperature. The TEM studies demonstrated the 20 nm uniform NPs.

Keywords: Fe/CeO₂ nanoparticles; Crystal structure; Solgel; Chemical synthesis.

Introduction

Metal oxide semiconductor (MOS) NPs have greatly applications such as magnetic recording media, nanocatalysis and biology sciences [1–28]. Nanostructured materials are one of the most popular materials in science and engineering and will have many applications in the coming technologies [29-36].

Due to excellent physical and chemical properties such as fuel cells, CeO₂ has recently received much attention [37]. Cerium oxide is one of the most important rare earth metal oxides and has many applications in the industry, including in solid oxide catalysts and fuel cells. Cerium oxide, as a relatively high dielectric constant, has attracted researchers' attention in applications such as dielectric capacitors available in dynamically accessible memory and oxygen

sensors. Another important application is magnetic oxides diluted at a temperature higher than room temperature, creating a new classification for the study of magnetic properties. This kind of oxide has a potential electrolyte for medium-temperature solid oxide fuel cells and has recently become a promising material for second-generation spintronics due to the ferromagnetic properties observed at room temperature. Nanostructured materials based on CeO₂ have achieved significant impact in practical applications due to the overall improvement of catalytic, electrical and optical properties. Solid oxide fuel cell technology is currently the most suitable field in which CeO₂ applications can be found. CeO₂ is a semiconductor with UV (388nm, E_g= 3.2eV) and visible light absorption.

Defects of CeO₂ include rapid formation and

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elimination, and formation in the concentrated or decentralized 4f electron layer. The doping element needs to be biocompatible. In that context, Fe^{3+} seems to be a good candidate for cosmetic application, which has not yet been studied for UV filtration applications. Indeed, this biocompatible element has a lower valence state than Ce^{4+} , as well as being cheap and abundant. Furthermore, it can bring a brown color to the CeO_2 phase, which is naturally pale yellow in colour, and thus give a brown tinge to the final product [38]. High oxygen storage capacity is one kind of applications of Fe-doped CeO_2 .

Fe^{3+} -doped CeO_2 nanocrystals have been prepared by various methods such as sol-gel [39], hydrothermal [40], coprecipitation method [41] and ball milling process [42]. Among these methods, the sol gel method has been widely used for easy reaction control and synthesis of single-size nanoparticles [17].

Materials and Methods

Fe-doped CeO_2 NPs were fabricated by a simple solgel synthesis. Firstly, 3g $\text{CeCl}_2 \cdot 5\text{H}_2\text{O}$ was dissolved in 100 mL pure water and then 3g $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ was dissolved to the solution and then stirred at room temperature for 10 min and finally synthesis temperature was increased to 85°C . The pH=4 was adjusted during the fabrication. The product were synthesized for 3 hours and annealed at 500°C for 3 hours. The crystal structure and morphology of the as-

prepared and calcined NPs were done by XRD, FESEM and TEM analyses.

Results and Discussion

Crystal structure and crystalline phases was identified by XRD analysis. Figure 1 shows the XRD spectra of the sample after calcination at 500°C for 3 hours. The peaks, which were formed at 2θ angles of 28.67° , 33.25° , 47.59° , 56.45° , 69.53° , 76.80° and 79.16° , were indexed to the x-ray reflections of (111), (200), (220), (311), (400), (331) and (420) respectively, representing the cubic structure (according JCPDS Card no. 21-1272). The average size of the NPs has been determined by Sherrer formula [43].

The crystallite size for as-synthesized sample was determined in the range size of 15-25 nm from the Sherrer formula. The results showed that iron impurities ions penetrate the crystalline network of cerium oxide nanoparticles and decrease their size in the boundaries. In addition, the increase in iron ions dopant increases the depletion in the Ce network through the oxygen present in the crystalline network, thereby reducing the size of the nanoparticles. In fact, doped ions of iron are dispersed in the CeO_2 lattice and replaced with cerium atoms. In addition, decrease in the size of the particles is due to adding Fe atoms on the surface of the nanoparticles.

In order to specify the morphology of the samples, FESEM analysis was used for as-prepared and annealed

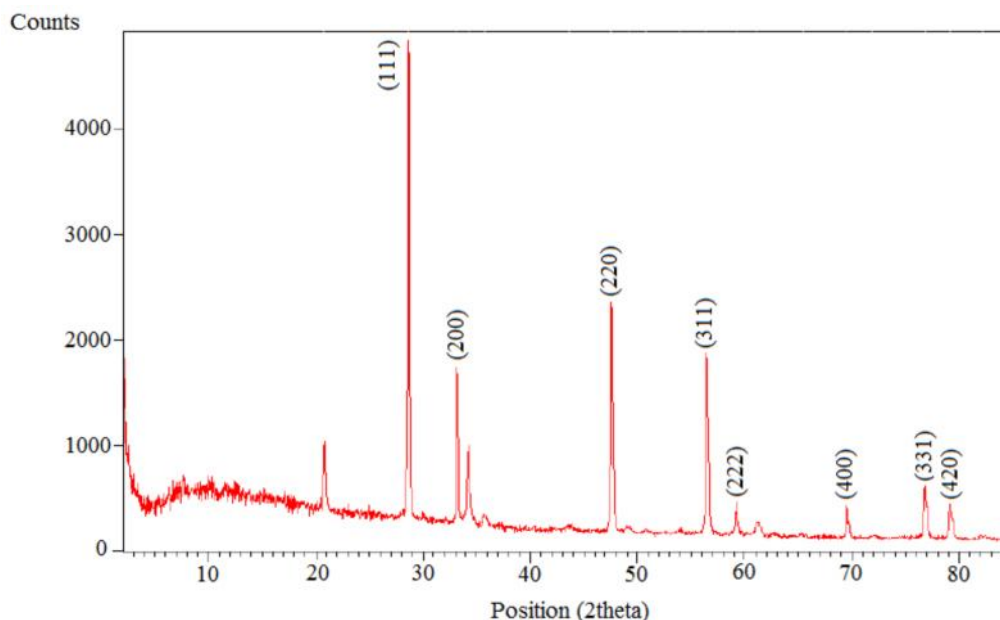


Figure 1. XRD pattern of annealed Fe-doped CeO_2 sample

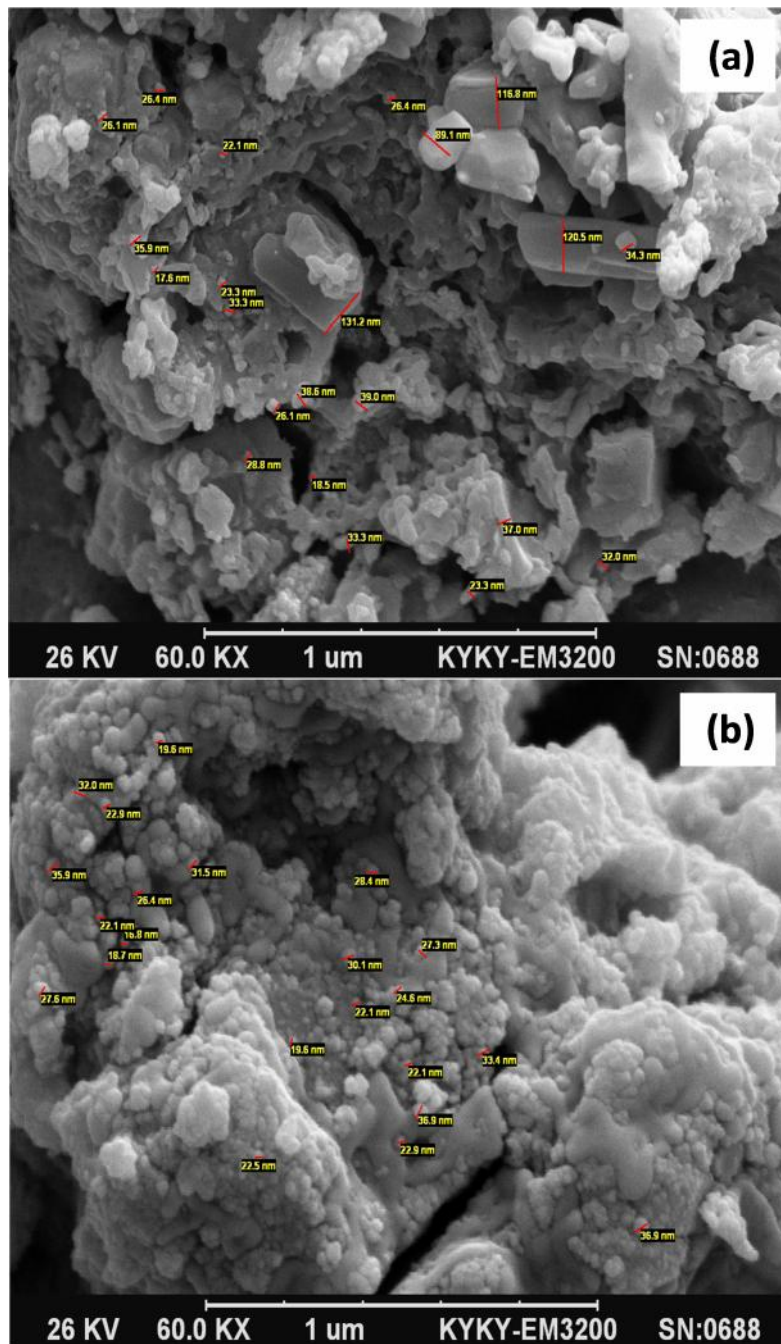


Figure 2. FESEM images of the (a) as-prepared (b) annealed Fe-doped CeO₂ NPs at 500 °C

NPs. Figure 2(a) reveals the FESEM image of the as-synthesized Fe-doped CeO₂ NPs. Figure 2(b) demonstrates the FESEM image of the heated FeCe NPs at 500°C for 3 hours. The FESEM analysis shows that the uniformity of the particles increase with increasing annealing temperature. In fact, the interatomic forces

increase with decreasing size of the nanoparticles, causing the nanoparticles to become closer to each other, resulting in their aggregation. The results is in correspondence with the average size of the nanoparticles measured by XRD analysis.

To estimate the actual size and shape of the NPs

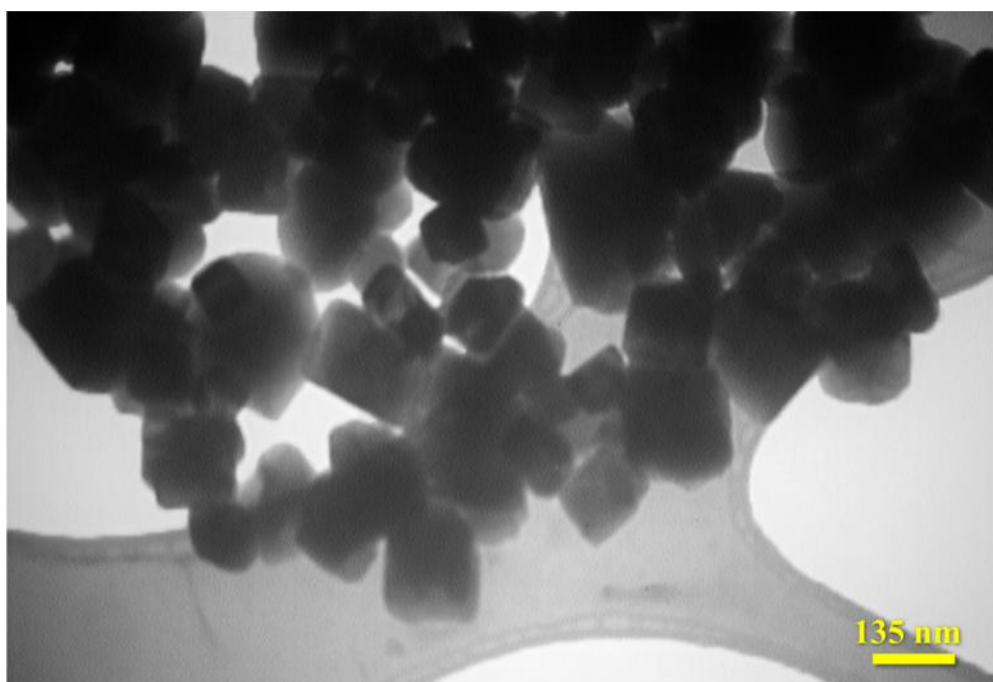


Figure 3. TEM image of the as-synthesized Fe-doped CeO₂ NPs

TEM analysis was carried out. Figure 3 shows the as-prepared TEM image of Fe/CeO₂ NPs with mean diameter of 20 nm prepared by solgel method. TEM also shows that the particles have been aggregated, which is due to the removal carboxylic group at 500°C.

Conclusions

Fe/CeO₂ NPs have been fabricated using simple solgel method. XRD spectrum shows single-crystal with face-centered cubic crystal cell of the CeO₂ structure and crystallizes in the cubic Fm-3m space group. FESEM images showed that the uniformity of the annealed NPs increased with less agglomeration. TEM image exhibits good uniformity of the as-prepared Fe-doped CeO₂ NPs with a mean diameter of 20 nm.

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