

## UV-Light Absorption and Photocatalytic Properties of Zn-doped CeO<sub>2</sub> Nanopowders Prepared by Ultrasound Irradiation

Aula Fitra Efendi<sup>1,a</sup> and Iis Nurhasanah<sup>1</sup>

<sup>1</sup>Department of Physics, Faculty of Science and Mathematics, Diponegoro University, Semarang, Central Java, Indonesia

<sup>a</sup>efendi\_aula@st.fisika.undip.ac.id (Corresponding author)

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**Abstract.** Ceria (CeO<sub>2</sub>) nanopowders doped with various Zinc (Zn) compositions were synthesized from solution by irradiating ultrasound waves. Ultrasound waves were irradiated to aqueous/isopropanol solution of cerium nitrate and zinc nitrate mixtures. Aqueous solution of ammonium hydroxide was dropped into that solution until pH becomes 10. Dried precipitates were calcined at 100°C to form CeO<sub>2</sub> nanopowders. X-ray Diffraction (XRD) analysis shows the CeO<sub>2</sub> nanopowder possess fluorite cubic structure. Ultrasound irradiation resulted in nano-metric powder of CeO<sub>2</sub> with spherical in shape. The addition of Zn into CeO<sub>2</sub> reduces the particle size and shows strong absorbance in the ultra-violet (UV) region. Moreover, the addition of 20 mol% Zn is inhibiting photocatalytic activity of CeO<sub>2</sub> under sunlight irradiation. These results suggest that Zn-doped CeO<sub>2</sub> is more promising for UV radiation protection with no presence photocatalytic activity.

### Introduction

Ceria (CeO<sub>2</sub>) nanoparticles are extensively studied through the last few decades due to effective for many applications such as absorbent, UV filter and blocker [1-2]. This material become one of the most attractive nanomaterials for research objectives, one of which is because of its interesting properties for UV radiation protection. For these application, CeO<sub>2</sub> with low photocatalytic ability is needed. CeO<sub>2</sub> nanoparticles are oxide material with fluorite cubic structure. The properties of that CeO<sub>2</sub> can be changed by doping Zn. The inserting Zn into CeO<sub>2</sub> lattice can reduce that particle size. As consequence, it reduce catalyst and photocatalyst activity, the evolution of molecular oxygen and preventing degradation of organic materials [3]. In recent years, many effort have been made to synthesize CeO<sub>2</sub> nanoparticles through different approaches. These methods include sol-gel, precipitation, and sonochemical method [4-6] etc. Sonochemical method has many advantages including easy of operation, being fast, low cost, high efficiency, convenient, time saving, and enviromental friendly [6]. Chemical reaction of the starting materials occurred in the presence of an applied high-frequency ultrasonic waves. These method has been employed for several purpose, in various organic and inorganic reactions and fabrication of nanostructured materials [7]. The chemical effect of ultrasonic irradiation arise from acoustic cavitation, in other words, the formation, growth and impulsive collapse of bubbles in a liquid medium, which result in an instantaneously high temperature and pressure pulse [8,9]. These special conditions of high temperature, pressure and local intense micromixing attained during acoustic cavitation lead to many unique properties in the irradiated solution [10].

Zn-doped CeO<sub>2</sub> nanopowders were synthesized from solution assisted by ultrasound irradiation. Generally, the irradiation performs in solution with water solvent. In this paper, we use mixed solvent system (aqua/isopropanol) to obtain cerium and zinc precursor solution. It is expected to produce the fine particle and strong UV absorbance. Photocatalytic of Zn-doped CeO<sub>2</sub> nanopowders were examined to methylene blue (MB) degradation under sunlight irradiation. The Zn-doped CeO<sub>2</sub> nanopowders show excellent UV absorption and lower photocatalytic activity than TiO<sub>2</sub>. This result indicates that Zn-doped CeO<sub>2</sub> nanopowders are promising as an alternative UV radiation protection materials.

## Experimental Procedure

CeO<sub>2</sub> and Zn-doped CeO<sub>2</sub> nanopowders were synthesized from solution of Cerium (III) nitrate (0.07 M) and Zinc nitrate in a mixture of aqua DM and isopropanol solvent with volume ratio of 1:6. Ammonium hydroxide (3M) was dropped into that solution until pH 10. Ultrasound wave was irradiated to the solution with frequency of 40 kHz for 60 minutes. The result of irradiated solution was then washed and calcined at 100°C for 3 hours.

X-ray Diffraction (XRD) measurement was used to analyze structure of nanopowders. Optical properties of nanopowders were measured using ultraviolet-visible (UV-vis) spectroscopy. Photodegradation of 32 ppm MB solution under sunlight irradiation for 120 minutes was carried out to evaluate photocatalytic activity of nanopowders. Particle morphology of the synthesized nanopowders were observed by scanning electron microscopy (SEM).

## Result and Discussion

**Structural Analysis.** The diffraction patterns of undoped CeO<sub>2</sub> and Zn-doped CeO<sub>2</sub> nanopowders are shown in Fig.1. Various of diffraction peaks were observed in all the powders indicating the polycrystalline structure. All the diffraction peaks of the samples correspond to a pure fluorite cubic structure of CeO<sub>2</sub> (JPDS 43-1002). The (111), (200), (220), (311) dominant diffraction peaks could be seen in all of the samples. The sharp of those peaks is indicating that CeO<sub>2</sub> well crystallized. Moreover, the intensity of all the diffraction peaks is increased with increasing Zn content. The peaks become more broader with increasing Zn content than undoped CeO<sub>2</sub>. The broadening of peak is expected due to decreasing in the particle size. It indicates that Zn-doped CeO<sub>2</sub> nanopowders are consist of nanometric size particle. The other peaks of (222), (400), (311), (420), and (422) with small intensity were also observed in the pure CeO<sub>2</sub> diffraction pattern and that peaks is not appearing in the Zn-doped CeO<sub>2</sub>. The diffraction patterns of both pure CeO<sub>2</sub> and Zn-doped CeO<sub>2</sub> do not show the peaks correspond to others phase, so that Zn is successful substituted into CeO<sub>2</sub>.

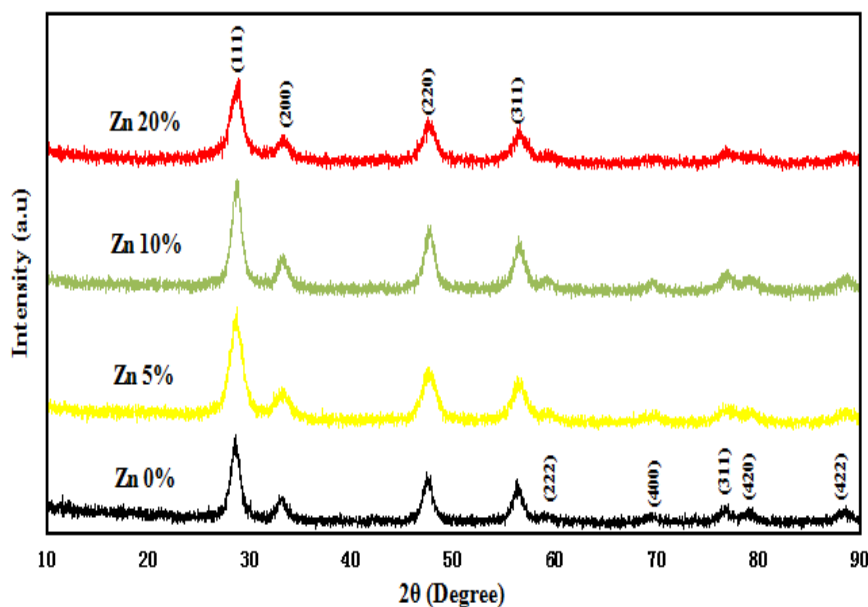


Fig. 1. X-ray diffraction patterns of Zn-doped CeO<sub>2</sub> with Zn content of 0%, 5%, 10%, 20%

The lattice constant of CeO<sub>2</sub> nanopowders can be determined by using Equation (Eq.) 1.

$$a = \frac{\lambda}{2 \sin \theta} \sqrt{h^2 + k^2 + l^2} \quad (1)$$

where  $a$  is the lattice constant,  $\lambda = 1,54056 \text{ \AA}$ ,  $h$ ,  $k$ ,  $l$  is miller index and  $\theta$  is the diffraction angle. The values of lattice constant were calculated by Eq. 1 ranging from  $5.4006 \text{ \AA}$  to  $5.4211 \text{ \AA}$ . These

values still indicates Zn-doped CeO<sub>2</sub> nanoparticles have fluorite cubic structure. The value of lattice constant decreases as the diffraction angle shifts to the right. The relationship between Zn content with lattice constant of CeO<sub>2</sub> nanopowders was shown in Fig. 2. It can be seen that lattice constant decreases with increasing Zn content in CeO<sub>2</sub>. This decreasing shows that the volume of the CeO<sub>2</sub> cell has decreased due to the Zn<sup>2+</sup> effective ionic radius (0.74 Å) which is smaller than Ce<sup>4+</sup> ionic radius (0.97 Å) [11].

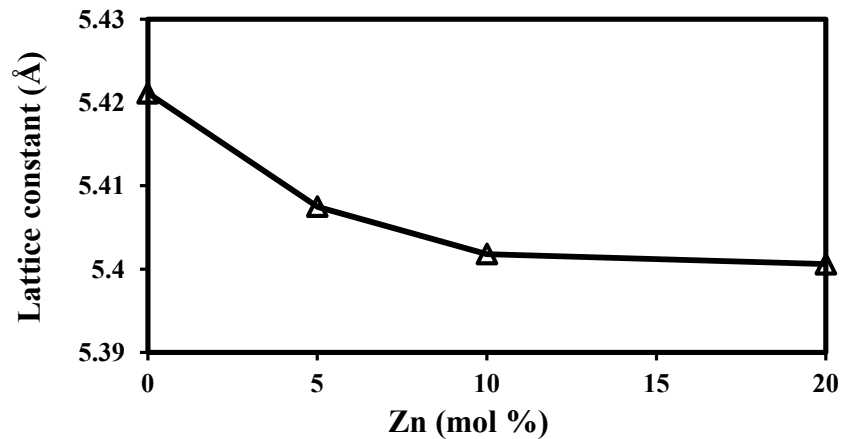


Fig. 2. Lattice constant of Zn-doped CeO<sub>2</sub> nanopowders vs Zn content.

**Particle Morphology.** Fig. 3 shows SEM image of undoped and Zn-doped CeO<sub>2</sub> nanopowders. The surface morphology of all that nanopowders appears to be rough and look spherical in shape. In this study, the estimation of average grain size are 37 nm for undoped CeO<sub>2</sub> and 28 nm for the 20 mol% Zn-doped CeO<sub>2</sub>. The addition of Zn content 5% and 10% was not estimated because the grain boundaries couldnot be clear seen. The average of grain sizes is reduced due to Zn doping.

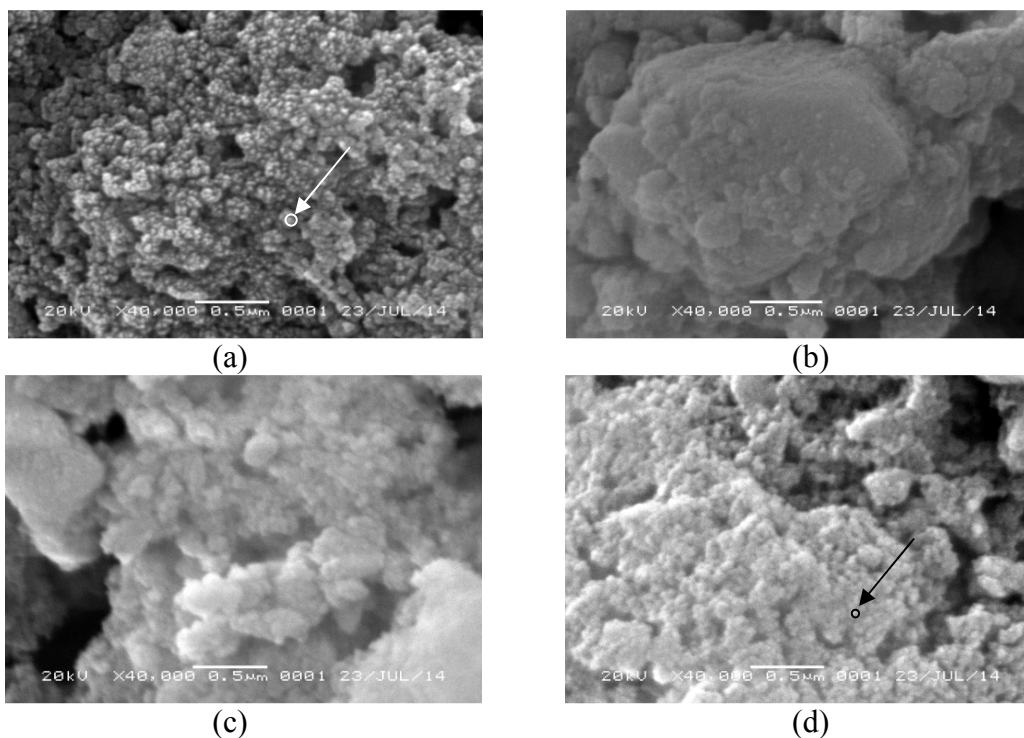


Fig. 3. SEM images of (a) undoped CeO<sub>2</sub> and Zn-doped CeO<sub>2</sub> nanopowders with Zn content (b) 5%, (c) 10%, (d) 20%

**Optical Properties.** The UV absorption spectra of undoped and Zn-doped CeO<sub>2</sub> nanopowders is shown in Fig. 4. CeO<sub>2</sub> nanopowders have strong absorption properties in the ultraviolet region. The undoped and Zn-doped CeO<sub>2</sub> increase slightly in the UV absorption. The change in optical properties of CeO<sub>2</sub> due to presence Zn dopant in the lattice CeO<sub>2</sub> are related to the particle size and the aggregation condition of particle [4]. The smoother particle surface is better light scattered than the rougher particle surface. It can be correlated by the results of SEM image. The surface morphology of undoped and the 20 mol% Zn-doped CeO<sub>2</sub> are more smoother than the 5 and 10 mol% Zn-doped CeO<sub>2</sub>. It causes that the UV absorption of undoped CeO<sub>2</sub> is more little stronger than the others. It is caused by 5 and 10 mol% Zn-doped CeO<sub>2</sub> nanopowders have a little agglomeration as can be seen in SEM image.

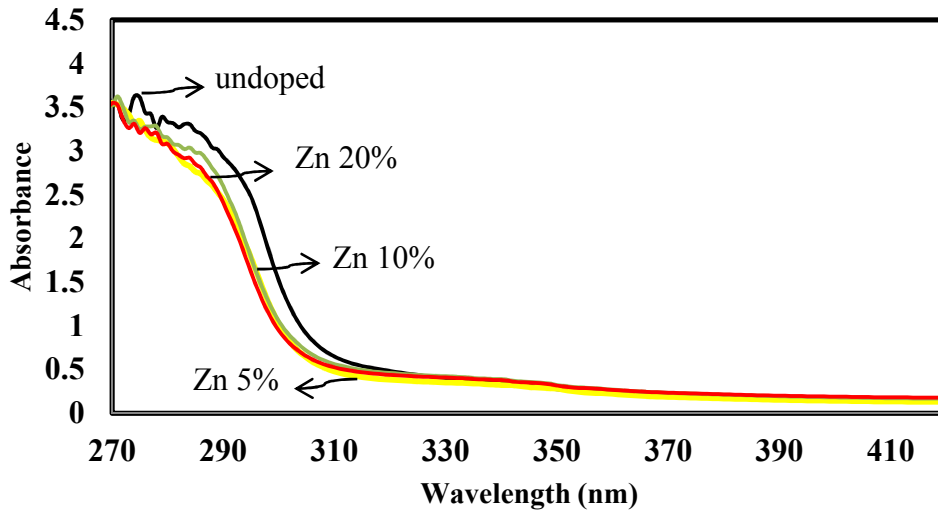


Fig. 4. UV absorbance spectra of CeO<sub>2</sub> nanopowders with Zn content of 0% (undoped), 5%, 10% and 20%.

The effect of Zn dopant to photocatalytic activity of CeO<sub>2</sub> was evaluated by the photodegradation of MB under sunlight irradiation for 120 minutes. The photocatalytic activity of Zn-doped CeO<sub>2</sub> nanopowder is also compared to the photocatalytic activity of TiO<sub>2</sub> commercial. The degradation of MB solution was calculated by Eq. 2

$$\text{Degradation (\%)} = \frac{A_0 - A_1}{A_0} \times 100\% \quad (2)$$

where  $A_0$  and  $A_1$  are absorbance of MB solution before irradiation ( $t = 0$  minutes) and after irradiation for  $t = 120$  minutes, respectively.

Degradation of MB solution under sunlight irradiation is shown in Fig. 5. By comparison to MB solution without photocatalyst material, the degradation of MB solution in the presence undoped and Zn-doped CeO<sub>2</sub> is almost equal. In the other hand, MB solution with TiO<sub>2</sub> possesses high degradation. These results show that the photocatalytic activity of undoped and Zn-doped CeO<sub>2</sub> is lower than TiO<sub>2</sub>. The lowest degradation is achieved for MB with 20 mol% Zn content in CeO<sub>2</sub>. It can be correlated to vacancies resulted from Zn<sup>2+</sup> incorporation into Ce<sup>4+</sup> lattice. That explanation can be shown from the decreasing of lattice constant. That vacancies act as traps for the charge [4]. The charge cannot reach the particle surface to induce free radical formation. This charge is responsible for MB degradation. The decreasing of charge on particle surface causes the inhibition of MB degradation.

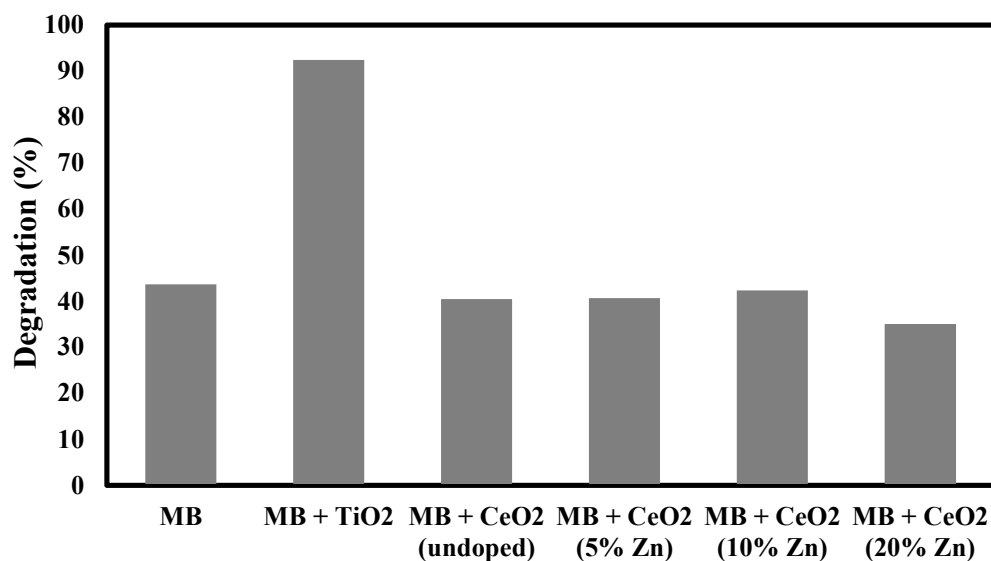


Fig. 5. The MB degradation under sunlight irradiation for 120 minutes.

### Summary

Undoped and doped with various Zn compositions (5%, 10%, 20%) of CeO<sub>2</sub> nanopowders were synthesized from solution by ultrasound irradiation. The effect of Zn doping on the structural, morphological and optical properties of that CeO<sub>2</sub> were investigated. The diffraction peaks in XRD pattern show that both undoped and Zn-doped CeO<sub>2</sub> were well crystalline and possess fluorite cubic structure. It is found that CeO<sub>2</sub> nanopowders consist of spherical particles. The addition of Zn into CeO<sub>2</sub> results in a decrease in particle size. The undoped and Zn-doped CeO<sub>2</sub> nanopowders have strong absorption properties in the ultraviolet region. These results show that Zn-doped CeO<sub>2</sub> is an excellent UV absorber and almost has no photocatalytic activity.

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