

Research Article

Influence of Europium Doping Amount on Properties of Antimony Doped Tin Oxide

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Abstract: In order to improve the performance of ATO nanometer powder, using $\text{SnCl}_4 \cdot 5\text{H}_2\text{O}$, SbCl_3 and Eu_2O_3 as raw materials, europium doped ATO non-powder was prepared by co-precipitation method. The precursor was examined by differential thermal gravimetric analyzer and the Eu^{3+} doped ATO nano-powder was examined by XRD and SEM to characterize the crystal structure, particle size and morphology. Comparison was carried out on the ATO nano-powder with or without Eu^{3+} doped. The results found that an appropriate Eu^{3+} doping amount can slightly increase the powder resistance, but to reduce the particle size of the powder and the agglomeration of the powder. Comparing with ATO nano-powder, Eu^{3+} doped ATO nano-powder had better performance.

Keywords: Antimony doped tin oxide, conductive nano-powder, Eu^{3+} doping amount, preparation

INTRODUCTION

Antimony-doped Tin Oxide (ATO) is an essential member of transparent conductive oxides (Muller *et al.*, 2009; Li *et al.*, 2010). SnO_2 is the first transparent conductor to attract many attentions to commercial (Varol and Hinsch, 1996). ATO has the advantage of chemically inert, mechanical hardness and heat-resistant (Burgurd *et al.*, 1998; Kim *et al.*, 2001; Crrijakorel *et al.*, 1992). Due to its excellent performances, ATO nano-powder was widely used in the fields of antistatic plastic (Al-Dahoudi *et al.*, 2001), coating (Zhang *et al.*, 2006), fibers (Wang *et al.*, 2004), gas sensors (Chatterjee *et al.*, 2003) and electrode materials (Wang *et al.*, 2006). ATO nano-particles have been synthesized by solid state reaction (Wang *et al.*, 2009), co-precipitation (Kim *et al.*, 2006) and hydrothermal methods (Jianrong and Gao, 2004). The catalytic and optical properties of the rare earth elements have been confirmed (Malpass *et al.*, 2007; Feng and Li, 2003; Feng *et al.*, 2009; Balaji *et al.*, 2007). Research of Eu^{3+} doped ATO nano-powder is few. It is important for improving characters of conductive ATO nano-powder. In this study, Eu^{3+} doped ATO nano-powder was prepared by chemical co-precipitation method. Influence of Eu^{3+} doping amount on ATO nano-powders performance was studied.

METHODOLOGY

In this study, co-precipitation method was used to prepare Eu^{3+} doped ATO precursor by controlling the molar ratio of Eu^{3+} ion, the liquid phase reaction

temperature and titration end point pH value. Then the precursor was calcined at 600°C to produce ATO nano-powder.

The preparation process is as follows: the molar ratio of $\text{SnCl}_4 \cdot 5\text{H}_2\text{O}$, SbCl_3 and Eu_2O_3 were dissolved in a certain amount of hydrochloric acid solution, forming a transparent acidic solution. In a thermostatic water bath, acidic mixed solution were stirred by adding aqueous ammonia until the solution reach and maintain a certain pH value for 0.5 h to form a precipitate. The mother liquor containing precipitate was stood and aged at room temperature for 2 h, then filtered. Filter cake obtained was washed successively with distilled water and alcohol several times and then dried at 80°C to gain a dry powder. Finally, the dry powder was sintered at 600°C for 2 h to produce Eu^{3+} doped ATO nano-powder.

Weigh 3 g Eu^{3+} doped ATO nano-powder, then put them into a Φ 16.5 mm stainless steel mould. The tablets were prepared by a direct compression method under 183 Mpa static pressure. The resistances of the tablets were measured using DC bridges. The resistivity of Eu^{3+} doped ATO nano-powder was calculated by following formula:

$$\rho = \frac{\pi R D^2}{4 L} \quad (1)$$

where,

ρ = Denotes resistivity ($\Omega \cdot \text{cm}$)

R = Resistance (Ω)

D = Tablet's diameter (cm)

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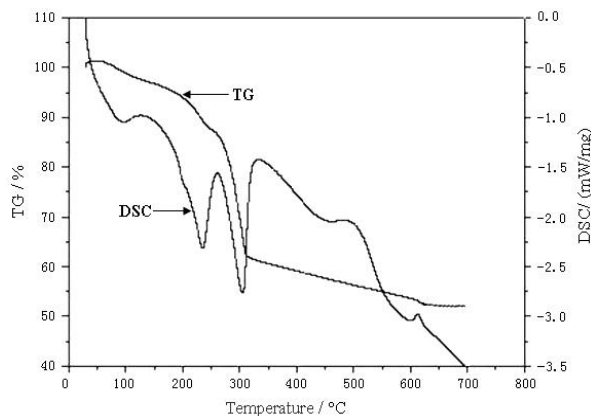
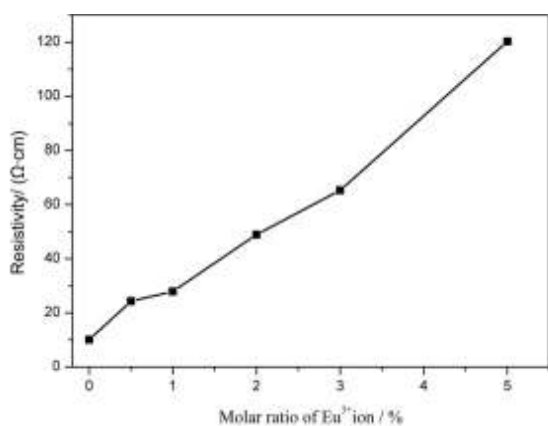


Fig. 1: DSC/TG curves of precursor

Fig. 2: Resistivity of ATO nano-powder with different molar ratio of Eu^{3+} ion

L = Tablet's thickness (cm)

The thermal decomposition behavior of the precursor obtained was investigated by DSC/TG (TA, SDT-2960 and USA). XRD analysis of the Eu^{3+} doped ATO nano-powder was performed on a XRD diffractometry (Rigaku, D/max-rA10, Japan). Morphology of Eu^{3+} doped ATO nano-powder was characterized on a Scanning Electron Microscope (SEM) (JEOL, JSM 5600LV, Japan).

RESULTS AND DISCUSSION

Thermogravimetric analysis of the precursor: Thermal decomposition result of the precursor under optimum conditions from the ambient temperature to 800°C in nitrogen atmosphere was given in Fig. 1. The DSC/TG curves indicate that a little, endothermic effect, one DSC peak is observed about 90°C , which can be considered to the evolution of water in the precursor. The two endothermic peaks with 22% weight loss at about 220°C and 320°C are observed for the decomposition of $\text{Sn}(\text{OH})_4$ and $\text{Sb}(\text{OH})_3$ with reactions $\text{Sn}(\text{OH})_4 = \text{SnO}_2 + 2\text{H}_2\text{O}$ and $2\text{Sb}(\text{OH})_3 = \text{Sb}_2\text{O}_3 + 3\text{H}_2\text{O}$, respectively.

Effect of Eu^{3+} doping amount on ATO resistivity:

Figure 2 shows the relationship of ATO resistivity with different molar ratio of Eu^{3+} ion under the conditions of $\text{pH} = 9$, the reaction temperature = 60°C and heat treatment temperature = 600°C . It indicated that the ATO resistivity increased significantly with the increase of molar ratio of Eu^{3+} ion. SnO_2 is a wide band gap n-type semiconductor, whose resistivity is larger. Research (Jeon *et al.*, 2005) found that the composition of the SnO_2 powder particles was deviated from the stoichiometric ratio of SnO_2 . Doping with other elements can be taken to improve the conductive properties of the SnO_2 powder. After heat treatment, Sb^{5+} and Eu^{3+} ions doped dissolved in the SnO_2 lattice, Sb^{5+} and Eu^{3+} ions easily became substitutional ions to occupy Sn^{4+} position. When Sb^{5+} ion occupied Sn^{4+} position in the SnO_2 lattice, a positively charged ion and an excess of valence electron were formed. The valence electron easily broke free of the shackles to be a conduction electron. Therefore, Sb^{5+} ion doped into SnO_2 can increase net conduction electron and increase the ATO conductivity. Rare earth europium represented +3 valence and Eu^{3+} ion was acceptor level for SnO_2 . So presence of Eu^{3+} ion will increase the resistivity of the ATO powder. When Eu^{3+} doping amount was lower, the increase of ATO nano-powder resistivity was relatively small. Nonetheless, continuing to increase the molar ratio of Eu^{3+} ion, the resistivity of the ATO nano-powder will increase rapidly.

The ionic radius of Sn^{4+} is 0.071 nm and the ionic radius of Eu^{3+} (0.095 nm) is much larger than that of Sn^{4+} . When Eu^{3+} ion substitute Sn^{4+} ion in the SnO_2 lattice, SnO_2 lattice will be deformed and movement of conductive carrier will be hindered, reducing ATO nano-powder conductivity. When Eu^{3+} doping amount was very low, distortion of SnO_2 lattice was also insignificant and had no apparent effect on ATO resistivity. Distortion rate of SnO_2 lattice was increased with the increase of the molar ratio of Eu^{3+} ion, resulting in an increase of ATO resistivity. According to the effect of Eu^{3+} doping amount on ATO nano-powder resistivity, molar ratio of Eu^{3+} ion was selected as 1%.

XRD and average grain size of ATO nano-powder with different molar ratio of Eu^{3+} ion: XRD patterns of ATO nano-powder after heat treatment for 2 h with different molar ratio of Eu^{3+} ion were given in Fig. 3. As can be seen from Fig. 3, incorporation of Eu^{3+} does not form a phase containing europium and the crystal structure of the ATO nano-powder remains tetragonal rutile structure of SnO_2 . Nevertheless, incorporation of Eu^{3+} influences the orientation of the crystal and the average grain size. Diffraction peak intensity of ATO nano-powder becomes weaker and weaker with the increase of the molar ratio of Eu^{3+} and peak width is

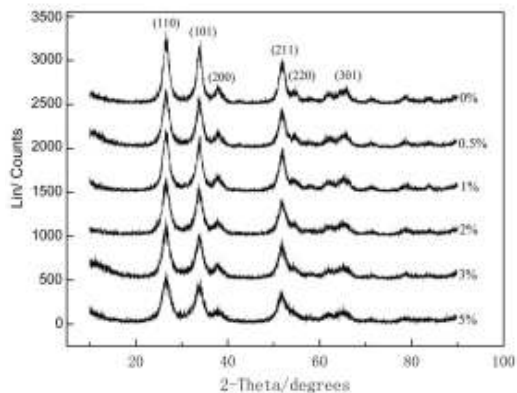


Fig. 3: XRD patterns of ATO nano-powder with different molar ratio of Eu^{3+} ion

increasing, indicating that crystallinity of the ATO powder becomes poor and the grain size becomes small.

Grain size of ATO nano-powder can be approximately estimated by Scherrer formula. The calculation of average grain size of ATO powder with different molar ratio of Eu^{3+} ion was given in Fig. 4. It is obvious that the grain size of ATO powder is smaller when Eu^{3+} doped in ATO powder. When the molar ratio rapidly, Doped Eu^{3+} ion has replaced Sn^{4+} into the SnO_2 of Eu^{3+} ion exceeds 1%, ATO powder grain size decreases lattice to form a solid solution. A large

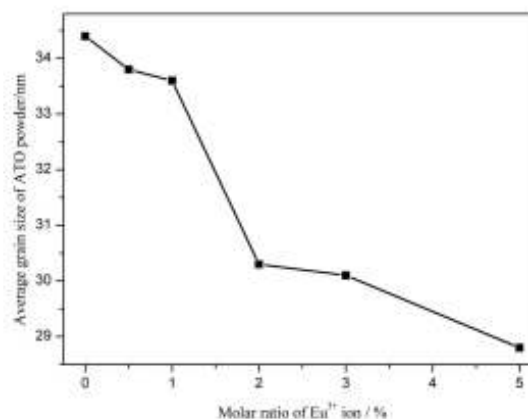


Fig. 4: Average grain size of ATO powder with different molar ratio of Eu^{3+} ion

number of lattice defects will be produced in the substitution process. It makes lower crystallinity degree of ATO powder and the lattice structure of SnO_2 is not complete. Simultaneously, the Eu^{3+} concentration in the crystal surface is relatively high, increasing the surface diffusion barrier. So that the grain boundary diffusion activation energy increases, hindering the diffusion movement of the grain boundary. Less Eu^{3+} doping amount has a small effect on SnO_2 lattice and surface diffusion barrier and the grain size of ATO powder changes little. With the increase of molar ratio

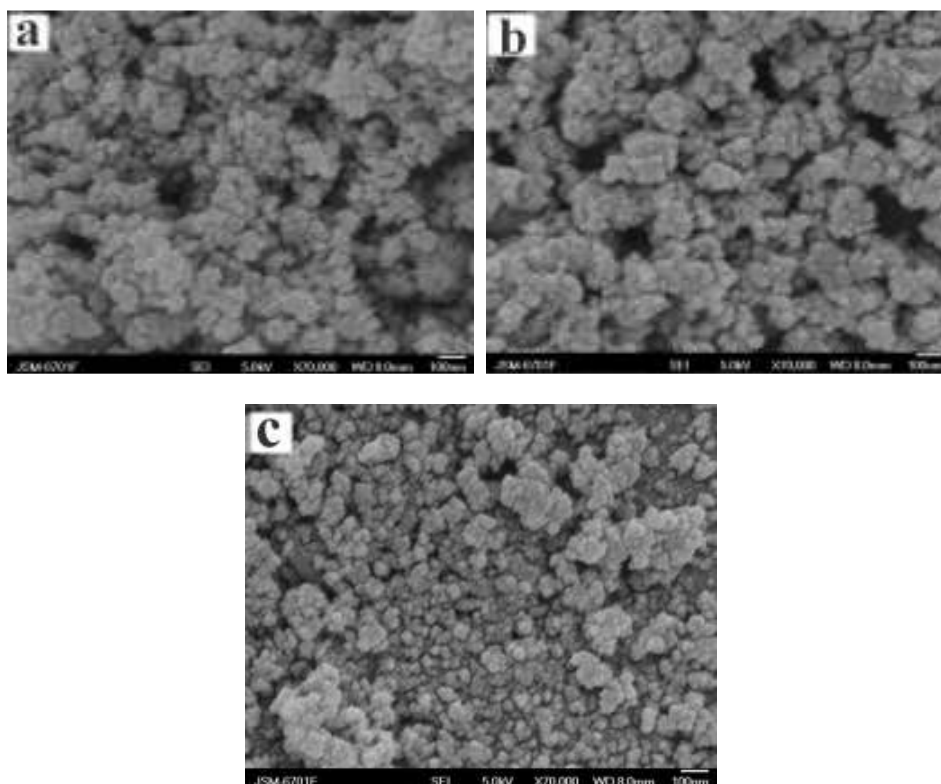


Fig. 5: SEM images of ATO powder with different molar ratio of Eu^{3+} ion (a) 0%, (b) 1%, (c) 5%

Table 1: Ti/SnO₂-Sb electrode electro-catalytic degradation efficiency and electrode life

Molar ratio of Eu ³⁺ ion/%	0%	1%
Electro-catalytic degradation efficiency of electrode/ %	48.3%	69.1%
Electrode life/hour	26.1	56.6

of Eu³⁺ ion, its impact on SnO₂ lattice and surface diffusion barrier will continue to increase, thus impeding the growth rate of the ATO powder grains and decreasing the grain size of the ATO powder. Considering the effect of Eu³⁺ doping amount on conductive properties, structure and grain size of ATO powder, molar ratio of 1% Eu³⁺ ion is more appropriate.

Morphology analysis of ATO nano-powder:

Figure 5a, b and c are SEM pictures of ATO nano-powder with the molar ratio of 0, 1 and 5% Eu³⁺ ion, respectively. As can be seen from Fig. 5, there are different degrees of agglomeration for three molar ratios of ATO powder. Agglomerations of un-doped (Fig. 5a) and 1% Eu³⁺ ion doped powder (Fig. 5b) are more serious, but 1% Eu³⁺ doped powder is smaller than that of un-doped powder. Comparing with the un-doped and 1% Eu³⁺ doped ATO powder, agglomeration for molar ratio of 5% Eu³⁺ doped ATO powder is significantly smaller and single irregular ellipsoidal ATO powder particles can be seen from Fig. 5c.

Different agglomeration of the above mentioned three molar ratios of Eu³⁺ doped ATO powder indicates that ATO powder doped with Eu³⁺ can reduce the powder agglomeration. Nano-powder prone to soft reunion due to its small particle has a great specific surface and the surface atoms of powder are extremely active for the presence of a large number of vacancy bonds, which make the van der Waals force, are very large. The presence of van der Waals force is easy to cause the powder agglomeration. Research (Qiang *et al.*, 1999) found that the powder surface existed potential, which was closely related to powder dispersion. Enrichment of Eu³⁺ ion in the powder surface is able to increase the surface potential of the powder and repulsive effects between the same kinds of particles. Thereby the action of the van der Waals force is offset and agglomeration of the powder is reduced.

Electro-catalytic degradation performance of Eu³⁺ doped Ti/SnO₂-Sb electrode prepared by ATO nano-powder:

Using p-nitrophenol as a target organic, electro-catalytic degradation performances and life of Eu³⁺ doped Ti/SnO₂-Sb electrode prepared by the molar ratio of 0% and 1% Eu³⁺ doped ATO nano-powder was carried out, respectively. The results were listed in Table 1. Eu³⁺ doped has an observable effect on electro-catalytic degradation performance and electrode life of Ti/SnO₂-Sb electrode. Therefore, an appropriate Eu³⁺ doped in ATO nano-powder can apparently increase powder electro-catalytic performance.

CONCLUSION

- Eu³⁺ doped ATO nano-powder was prepared by co-precipitation method. The effect of Eu³⁺ doping amount on properties of ATO nano-powder was investigated. The results found that the molar ratio of 1% Eu³⁺ doping amount can significantly improve the electro-catalytic performance of the ATO nano-powder. Electro-catalytic degradation efficiency of 1% Eu³⁺ doped electrode, which is greater than that of Eu³⁺ un-doped electrode, is up to 69.1%.
- Eu³⁺ doped in ATO nano-powder can reduce the grain size and agglomeration of powder, but will increase the resistivity of the powder. Eu³⁺ doped in ATO powder does not change the rutile structure of SnO₂.

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