

Structure and luminescence property of Eu^{2+} , Dy^{3+} -doped SrO-MgO-SiO_2 glass

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Abstract. Eu^{2+} , Dy^{3+} -doped SrO-MgO-SiO_2 glass was firstly synthesized by sol-gel method, its structure features and optical properties have been investigated by X-ray diffraction and luminescence spectra. X-ray diffraction pattern indicates the amorphous structure of glass phase. According to luminescence spectra, Strong red emission which centered at 587nm which are attributed to the energy level transition of $5D_i(i=0,1) \rightarrow 7F_j(j=0\sim4)$ from Eu^{2+} .

Introduction

With the development of the science and technology, the luminescent materials has attracted attention over the recent years^[1-4]. The kinds of the luminescent materials have from polycrystalline to monocrystalline particale, thin-film, ceramic, glass and so on^[5-9]. But the worse stability and comfection of crystals limited their further application. As the glass has the characters of homogeneous and transparent, and it can be mixed at higher concentration of doped luminescent ions, So many scientists have increased interest in glass, which has great potential for applications in optical information processing^[10-12].

In this paper, Eu^{2+} , Dy^{3+} -doped SrO-MgO-SiO_2 glass was synthesized by sol-gel preparing process.and their structure and luminescent properties were also studied.

Expriment

Synthesis. Eu^{2+} , Dy^{3+} -doped SrO-MgO-SiO_2 glass was synthesized by sol-gel preparing process. $\text{Mg}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, $\text{Sr}(\text{NO}_3)_2$ were dissolved to water; $\text{Si}(\text{OC}_2\text{H}_5)_4$ was dissolved to ethanol. The two kinds of solution above were mixed under stirring. Molar ratio of $\text{SrO} : \text{MgO} : \text{SiO}_2 : \text{Eu}_2\text{O}_3 : \text{Dy}_2\text{O}_3$ was 40: 20: 40: 0.1:0.2. When the solution was stirred for 0.5h, a mount of citric acid (CA) that was used as the chelating was added to the above solution, and subsequently pH was tuned to 1.0 by nitric acid. After stirring for 2h, the solution was aged for 24h at 80°C to form gel in a water bath. The dried gel was sintered at a higher temperature at 1000°C for 1h, with a heating rate of 5°C/min. All heat treatments were performed in an atmosphere of air.

Measurements. Fluorescence spectra was measured with a Hitachi F-7000 fluorescence spectrophotometer. Differential thermogravimetric analysis(DTG) and differential thermal analysis (DTA) of the powdered gel precursors was carried out on a HCT-2 microcomputer differential thermal Balance. X-ray diffraction (XRD) analysis was performed on a BDX3200 diffractometer using CuK_α radiation.

Results and discussion

DTA-TG analysis.

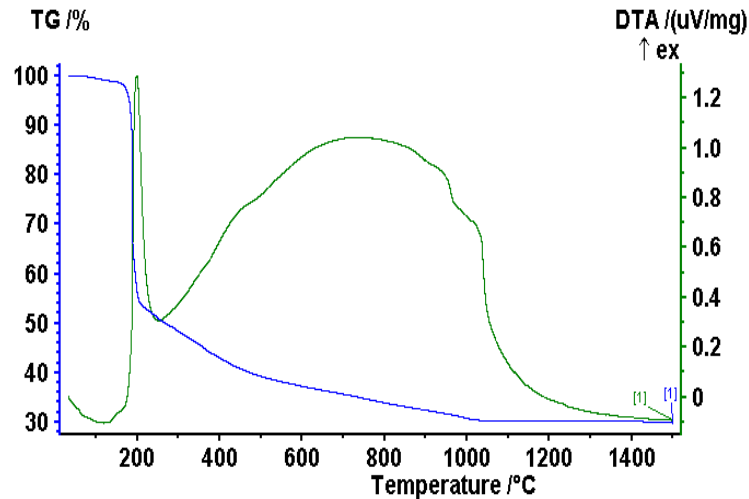


Fig.1 DTA-TG curves of the xerogel

Fig.1 shows the DTA-TG curves of xerogel compositing of $40\text{SrO}-20\text{MgO}-40\text{SiO}_2: 0.1\text{Eu}^{2+}, 0.2\text{Dy}^{3+}$, which has been dried at 60°C for 24 hours. TG curve indicates that the mainly weight loss appears between 150°C and 1050°C , which is 68.23%. The first exothermic peak occurs at $180\sim 200^\circ\text{C}$, accompanied by mass loss of 46.03%, which could be attributed to decomposing removal of citric acid in xerogel. The second endothermic peak near 260°C accompanied by mass loss of 17.67%, which could be caused by decomposing removal of nitric acid in xerogel. The third exothermic peak from 500°C to 1000°C , accompanied by mass loss of 23.07%, is due to pyrolysis of the residual citrate, residual nitrate and TEOS in the sample. According to the DTA-TG analysis, the heat processing treatment system of $\text{Eu}^{2+}, \text{Dy}^{3+}$ -doped SrO-MgO-SiO₂ glass was determined.

X-ray diffraction pattern. Fig.2 shows the XRD pattern of the $40\text{SrO}-20\text{MgO}-40\text{SiO}_2: 0.1\text{Eu}^{2+}, 0.2\text{Dy}^{3+}$ glass. It could be found that there is no diffraction peaks in the XRD pattern, but two typical dispersion peaks of glass, which means there is no crystalline phase, and the sample is in a state of non-crystalline structure.

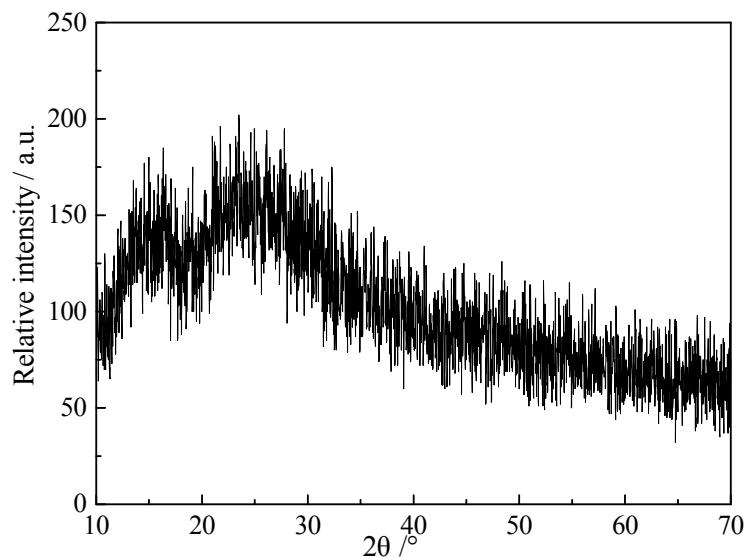


Fig.2 XRD pattern of the sample

Fluorescence spectra .

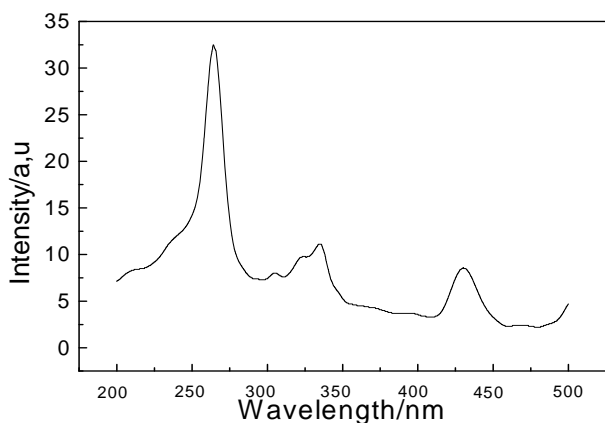


Fig.3 Emission spectra of the sample

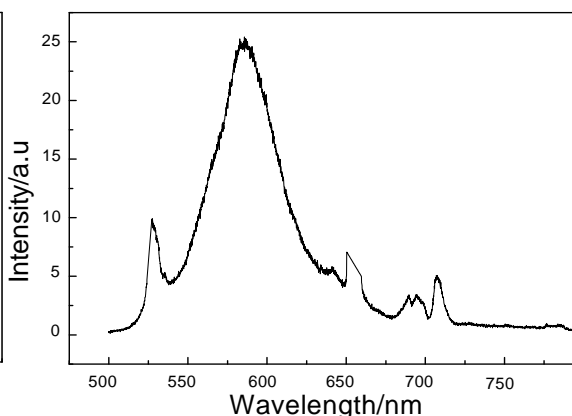


Fig.4 Excitation spectra of the sample

Fig.3 shows emission spectra of 40SrO-20MgO-40SiO₂: 0.1Eu²⁺, 0.2Dy³⁺ glass. The excitation spectra revealed three excitation bands such as 230~300nm, 350~400nm, 400~450nm. stronger broadband absorption occurs at 270nm, which belongs to the electron charge excitation stage transition of Dy³⁺→O²⁻. The other bands are due to the transition of Eu²⁺. Fig.4 shows Excitation spectra. The spectra excited at 530nm, 587nm, 654nm, 704nm and 718nm which ascribed to the ⁵D_i(i=0,1)→⁷F_j(j=0~4) transition of Eu³⁺ ion.

Conclusions

In conclusion, SrO-MgO-SiO₂:Eu²⁺, Dy³⁺ glass was prepared by sol-gel method. Heat treating process, luminescence properties and glass structure were attributed and analysed by luminescence spectra, XRD pattern, and DTA-TG and so on, which indicate that this glass with outstanding luminescence property has a huge potentiality in the information processing area.

Acknowledgements

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References

- [1] Gurkalenko Y A, Distanov V B, and Berdanova V F, et al. Application of luminescent materials to encoding in authorized access systems Dyes and Pigments[J].1999, 43(2):147-151.
- [2] Bomm J, Büchtemann A, and Amanda J, et al. Fabrication and full characterization of state-of-the-art quantum dot luminescent solar concentrators [J]. Solar Energy Materials and Solar Cells, 2011, 95(8):2087-2094.
- [3] Zhang H J, Fan R Q, and Zhou G P, et al. Syntheses, structures, and luminescent properties of three novel two-dimensional lanthanide coordination polymers with mixed aromatic carboxylate ligands[J]. Inorganic Chemistry Communications, 2012, 16: 100-103.
- [4] Tan C L, and Wang Q M. Photophysical studies of novel lanthanide (Eu³⁺ and Tb³⁺) luminescent hydrogels Inorganic Chemistry Communications, 2011,14(4): 515-518.
- [5] Yang Q H, Fang L, and Zheng F G, et al. Structural, electrical, luminescent, and magnetic properties of Ba_{0.77}Ca_{0.23}TiO₃:Eu ceramics [J]. Materials Chemistry and Physics, 2009, 118(2-3): 484-489.
- [6] Yamazaki M., Yamamoto Y., and Nagabama S., Long luminescent glass Tb³⁺-activated ZnO-B2O3-SiO₂ glass [J]. Non-Cryst Solids, 1998, 241(1): 71-73.

- [7] Tomozawa M, Hong J W, Ryu S R. Infrared (IR) investigation of the structural changes of silica glasses with fictive temperature [J]. *Journal of Non-Crystalline Solids*, 2005, 351: 1054-1060.
- [8] Pisarska J, Żur L, and Goryczka T, et al. Local structure and luminescent properties of lead phosphate glasses containing rare earth ions [J]. *Journal of Rare Earths*, 2011, 29(12):1157-1160
- [9] Narayana R C, Sreekanth R P. Elastic properties and spectroscopic studies of fast ion conducting $\text{Li}_2\text{O}-\text{ZnO}-\text{B}_2\text{O}_3$ glass system [J]. *Materials Research Bulletin*, 2007, 42: 1337-1347.
- [10] Bomm J, Büchtemann A, and Amanda J. Fabrication and full characterization of state-of-the-art quantum dot luminescent solar concentrators [J]. *Solar Energy Materials and Solar Cells*, 2011, 95(8): 2087-2094.
- [11] Boulon G. Fifty years of advances in solid-state laser materials[J]. *Optical Materials*, 2012, 34(3): 499-512.
- [12] Reddy C. M, Reddy B S, and Dillip G R, et al. FT-IR, FT-Raman and fluorescence studies of Tb^{3+} ions activated lead containing sodium fluoroborate glasses[J]. *Journal of Molecular Structure*, 2012, 1019(18):166-173.