Photoluminescence in Li2basio4: Eu 2+ and Ce3+

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Abstract: Phosphor Li2BaSiO4: Eu2+ and Li2BaSiO4 :Ce are prepared through solid state reaction method at 700 C for 12 to 14 hrs. The photoluminescence of given phosphors are studied. PL emission of Li2BaSiO4: Eu2+ is observed in bluish green region. Emission is due to Eu2+ transition between the ground state 8S7/2(4f7) and the excited states of 4f65d1configuration. Weak long after glow emission is also observed. In Li2BaSiO4:Ce, emission peak is observed at 392 nm at excitation of 330 nm. Keywords: Photoluminescence, Eu2+and Ce3+ transition.

I. Introduction

Nowadays, the studies on red/green/blue tricolor phosphors suitable for UV excitation have attracted more and more attention because of their significant applications in solid state lighting (SSL), Plasma display panel, Liquid crystal display etc.. Up to now, many phosphors suitable for near-ultraviolet (UV) excitation have been reported, including orthosilicates, akermanites, aluminates, sulfides, molybdates, oxynitrides/nitrides, and so on[1].

Optical transitions of divalent europium 4f7 have been investigated in many silicate phosphors. The spectra of Eu2+ doped compounds are due to electric dipole 4f7 - 4f65d transitions which are parity-allowed so that they occur with high transition probabilities. The Eu2+ emission color varies from UV to red depending on the host lattice, covalency, the size of the cation, and the strength of the crystal field. Researchers are taking interest in development of phosphors like Li2CaSiO4, Li2BaSiO4, and Li2MgSiO4.

II. Experimental

In present work Li2BaSiO4 : Eu2+ and Li2BaSiO4 : Ce3+ phosphors are prepared through solid state reaction.Initially, BaSiO3:Eu (2%)/Ce(1%)was precipitated from aqueous solution of Barium nitrate by slowly adding Na2SiO3 solution. The precipitate formed was filtered, dried and then annealed in air at 700 C for 1 hour. Li2BaSiO4 was prepared by heating BaSiO3 with Li2CO3 at 700 C for 12 hours. The obtained phosphor is reduced at 800 C for 1 hr.

III. Results And Discussion

Fig1. shows the XRD pattern of Li2BaSiO4, which excellently matched with that reported by Kulshreshtha et al [2] and the data calculated by He et al [3], indicating that the compound crystallizes in the hexagonal structure (P63cm, a = 8.1 Å, c = 10.6, Z = 6). Fig.2, shows the hexagonal crystal structure of Li2BaSiO4.



Photoluminescence spectra are studied. In photoluminescence spectra of Li2BaSiO4:Eu2+ emission spectrum is obtained at 495 nm at excitation 335 nm (Fig.3). The intense bluish green emission is obtained. Both

the emission

and excitation bands can be assigned to the well-known Eu2+ transitions between the ground state 8S7/2 (4f7) and the excited states of 4f65d1 configuration. As the position of the band corresponding to 4f65d configuration is strongly influenced by the host, the emission can be anywhere from 365 nm to 650 nm.Baginskiy et al [4]. Weak long after glow emission is obtained, which is not measured. Hence given phosphor may possess the characteristics of thermoluminescence. This phosphor maybe used in light emitting diode



Figure 3: Photoluminescence (PL) spectra ofLi2BaSiO4.(a) Eu2+ emission for 335 nm excitation(b) Eu2+ excitation for 495 nm emission

The photoluminescence spectra of Li2BaSiO4:Ceis also studied.Fig.4,shows the PL spectra of Li2BaSiO4: Ce. The peak of emission spectrum is observed at 392 nm at excitation of 330 nm.PL spectra of Ce3+ can be conveniently discussed in terms of the scheme used by Dorenbos[5,6,7]. The Ce3+ ion has, from a spectroscopic point of view, a very simple electronconfiguration in the ground and excited state: 4f1 and 5d1, respectively. The 4f1 state is split by spin-orbit coupling into a doublet (2F5/2, 2F7/2) with an energy difference of 2000 cm-1. The 5d1 state is split by the crystal field into several components with an averaged total splitting of some 10000 cm -1. The emission consists of a transition from the lowest crystal-field component of the 5d1 state. The emission band has two maxima due to the spin-orbit splitting of the ground state. The spin–orbit split 2D3/2 and 2D5/2 states of the 5d configuration of free Ce3+ are located at 49700 and 52100 cm-1 above the 4f1 (2F5/2) ground state of Ce3+. When Ce3+ is introduced in a compound the average energy of the 5d configuration is lowered and the 2D3/2 and 2D5/2 states are further split by the crystal field. Depending on the site symmetry at most five distinct 5d states may form.



Figure 4: Photoluminescence (PL) spectra of Li2BaSiO4: Ce: (a) Ce emission for 392 nm excitation (b) Ce excitation for 330 nm emission

The energy difference between the maxima of the highest and lowest 5d band in spectra is defined as the total crystal field splitting (cfs). The energy shift of the average of the 5d configuration is defined as the centroid shift (c).



Figure 5: Transition in Ce³

IV. Conclusion

Photoluminescence spectra are studied. In $Li_2BaSiO_4:Eu^{2+}$, emission peak is observed at 495 nm at excitation 335 nm. The intense bluish green emission is obtained. Weak long after glow emission is obtained, which is not measured. Hence given phosphor may possess the characteristics of thermoluminescence. This phosphor maybe used in light emitting diode. In $Li_2BaSiO_4:Ce^{3+}$ emission peak of spectrum is observed at 392 nm at excitation of 330

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