

# LUMINESCENCE AND CHARGE TRANSFER BANDS OF THE Sm(3+) AND Eu(3+) IN Mg<sub>3</sub>BO<sub>3</sub>F<sub>3</sub>

Xingren LIU, Yinglan ZHANG, Zhihua WANG and Shuhua LU

Changchun Institute of Physics, Academia Sinica, Changchun, China

The excitation and emission spectra and the charge transfer bands of the Sm(3+) and Eu(3+) in Mg<sub>3</sub>BO<sub>3</sub>F<sub>3</sub> have been investigated under UV radiation and CR excitation. The charge transfer bands of Sm(3+) and Eu(3+) which center around at 43.5 and 34.2 X10<sup>3</sup>cm<sup>-1</sup>, respectively are observed at room temperature

## 1. INTRODUCTION

The magnesium fluoroborate material studied has composition Mg<sub>3</sub>BO<sub>3</sub>F<sub>3</sub>. The luminescence of the rare earth activators in the system Mg<sub>3</sub>BO<sub>3</sub>F<sub>3</sub> has not been reported. In the past investigation of charge transfer band was mainly made on oxides doped with Eu(3+) ion<sup>1,2</sup>.

In this paper we report the luminescence properties of the host lattice, Sm(3+) and Eu(3+) and charge transfer bands presenting in the excitation spectra of the Sm(3+) and Eu(3+) emissions in the magnesium fluoroborate.

## 2. EXPERIMENTAL

Samples discussed were prepared by solid-state reaction. An appropriate amount of MgO, MgF<sub>2</sub>, B<sub>2</sub>O<sub>3</sub> (or H<sub>3</sub>BO<sub>3</sub>), and Sm<sub>2</sub>O<sub>3</sub> or Eu<sub>2</sub>O<sub>3</sub> compounds were mixed and fired at 1100°C for 2 hours. The white powder samples obtained were checked by x-ray powder diffraction analysis and have the hexagonal structure of r-Mg<sub>3</sub>BO<sub>3</sub>F<sub>3</sub>. The luminescence properties of samples were measured by the YF-2 cathode-ray measurement system and MPF-4 type fluorospectrophotometer.

## 3. RESULTS AND DISCUSSION

It is found that a strong emission band peaking at 388 nm is the emission of the host lattice under the excitation of the

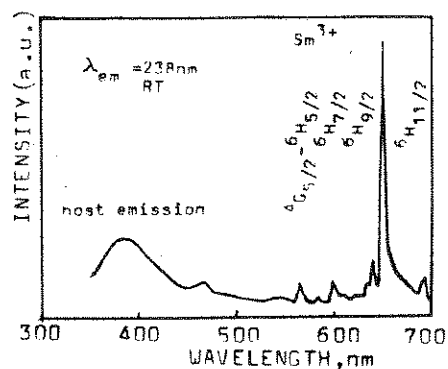


FIGURE 1  
Emission spectrum of Mg<sub>3</sub>BO<sub>3</sub>F<sub>3</sub>:Sm<sup>3+</sup> phosphor

short wavelength UV radiation. Main emissions of the Sm(3+) and Eu(3+) ions in Mg<sub>3</sub>BO<sub>3</sub>F<sub>3</sub> are respectively located at 651 nm of the Sm(3+) <sup>4</sup>G<sub>5/2</sub>-<sup>6</sup>H<sub>9/2</sub> transition and at 613 nm corresponding to the <sup>5</sup>D<sub>0</sub>-<sup>7</sup>F<sub>2</sub> transition of the Eu(3+) under UV radiation and cathode-ray excitation. The decay time to 10% initial brightness value is about 80 ms for the Sm(3+) and Eu(3+) doped samples under CR excitation.

The fluorescence spectrum of Mg<sub>3</sub>BO<sub>3</sub>F<sub>3</sub>:Sm(3+) involves a emission band of the host lattice situated in the blue-violet region and some characteristic emissions of Sm(3+) where the energy levels of Sm(3+) transition were marked, as in Fig. 1. The excitation spectrum of the Sm<sup>3+</sup> 615 nm emission in Mg<sub>3</sub>BO<sub>3</sub>F<sub>3</sub> is shown in Fig. 2. The excitation and emission

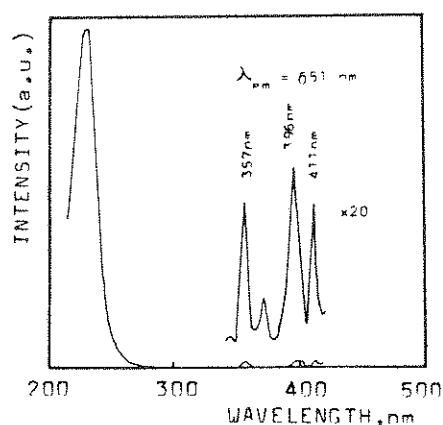


FIGURE 2  
Excitation spectrum of  $\text{Sm}^{3+}$  emission in  $\text{Mg}_3\text{BO}_3\text{F}_3$

spectra of the  $\text{Eu}^{3+}$  in the magnesium fluoroborate phosphor are shown in Fig. 3. All measurements were performed at room temperature.

Both the excitation spectra of the  $\text{Sm}(3+)$  and  $\text{Eu}(3+)$  emission in the magnesium fluoroborate contain an intense charge transfer band (CTB) and some narrow lines of the  $\text{Sm}(3+)$  and  $\text{Eu}(3+)$  4f-4f transitions. The energy positions of the charge transfer bands of the  $\text{Sm}(3+)$  and  $\text{Eu}(3+)$  ions in  $\text{Mg}_3\text{BO}_3\text{F}_3$  are located at 43.5 and 34.2 ( $\times 10^3 \text{ cm}^{-1}$ ), respectively. The difference value ( $9.3 \times 10^3 \text{ cm}^{-1}$ ) between the CTB positions of the  $\text{Sm}(3+)$  and  $\text{Eu}(3+)$  in magnesium fluoroborate very coincides with that one obtained by Jørgensen<sup>3</sup>

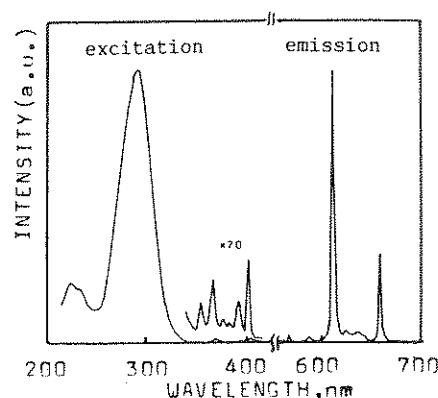


FIGURE 3  
Excitation(left,  $\lambda_{\text{em}} = 613 \text{ nm}$ ) and emission(right  $\lambda_{\text{ex}} = 295 \text{ nm}$ ) spectra of  $\text{Eu}(3+)$  in  $\text{Mg}_3\text{BO}_3\text{F}_3$ .

and Blasse<sup>4</sup> in other compounds.

#### ACKNOWLEDGEMENTS

The authors are grateful to Engineers Yu Baogui and Guan Zhongsu for measuring the optical spectra of samples.

#### REFERENCES

1. G. Blasse and A. Bril, J. Inorg. Nucl. Chem. 29 (1967) 2231.
2. G. Blasse, J. Chem. Phys. 45 (1966) 2356.
3. C. K. Jørgensen, Theoretical Chemistry of Rare Earths, in: Handbook on the Physics and Chemistry of Rare Earths, Vol. 3, eds. K. Gschneidner, Jr. and L. Eyring (North-Holland, Amsterdam, 1979) pp. 111-169.
4. G. Blasse and A. Bril, Phys. Letters 23 (1966) 4401.

SYNTHESIS  
Ho, Er,

Hongjie  
Changchun

Present  
Ca, Y, Sb, O  
The R<sup>3+</sup>  
yellow-  
discuss

#### 1. INTRODUCTION

Recently  
 $\text{Sb}_2\text{O}_3\text{-Eu}_2$   
have been  
on the lum  
timate i  
vestigate  
In this  
 $\text{Sm}^{3+}$ ,  $\text{Dy}^{3+}$   
(M=Ba, Ca  
of their  
our prece  
is to pres  
of the eff  
luminesce  
In add  
 $\text{Dy}^{3+}$  emis  
tion and  
concentra  
ratio of

#### 2. EXPERIMENTAL

The  
phosphors  
Their lat  
powder n  
structure  
monoclin

#### 3. RESULTS

0022-2313/  
(North-Holl