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# Effect of Cr<sup>3+</sup> doping on structural and optical properties of Eu<sup>3+</sup> doped LaVO<sub>4</sub> phosphor

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In this work, the Eu<sup>3+</sup>, Cr<sup>3+</sup> doped and co-doped LaVO<sub>4</sub> phosphors have been prepared through a high temperature solid-state reaction method. The powder XRD patterns of phosphors are very sharp and intense, which reflects a highly crystalline nature of phosphors. The XRD data were also refined by a Rietveld refinement method. The particle size of the phosphor samples lies in the sub-micron to micron range. The existence of La, Eu, Cr, V and O elements was verified by EDS spectra. The FTIR spectra show various absorption bands due to different vibrating groups. The optical band gap of the phosphor decreases on increasing concentration of Cr<sup>3+</sup> ion. The photoluminescence excitation spectra of Eu<sup>3+</sup>, Cr<sup>3+</sup> co-doped LaVO<sub>4</sub> phosphor exhibit bands due to Eu<sup>3+</sup> and Cr<sup>3+</sup> ions. The Eu<sup>3+</sup> doped LaVO<sub>4</sub> phosphor exciting at 393 and 316 nm wavelengths gives intense red color at 614 nm due to the <sup>5</sup>D<sub>0</sub> → <sup>7</sup>F<sub>2</sub> transition of the Eu<sup>3+</sup> ion. When the Cr<sup>3+</sup> ion is co-doped in the Eu<sup>3+</sup> doped LaVO<sub>4</sub> phosphor the emission spectra contain emission bands due to Eu<sup>3+</sup> and Cr<sup>3+</sup> ions. The emission intensity of Eu<sup>3+</sup> doped phosphor reduces due to energy transfer from Eu<sup>3+</sup> to Cr<sup>3+</sup> ions in presence of Cr<sup>3+</sup> ions upon 393 and 386 nm excitations. The lifetime of the <sup>5</sup>D<sub>0</sub> level of Eu<sup>3+</sup> ions decreases in the Eu<sup>3+</sup>, Cr<sup>3+</sup> co-doped LaVO<sub>4</sub> phosphor, which also reflects the energy transfer. The Eu<sup>3+</sup>, Cr<sup>3+</sup> co-doped LaVO<sub>4</sub> phosphor also produces a large amount of heat upon 980 nm excitation. Thus, the Eu<sup>3+</sup>, Cr<sup>3+</sup> co-doped LaVO<sub>4</sub> phosphors may be used for LEDs, solid state lighting and heat generating devices.

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## 1. Introduction

The rare earth doped phosphor materials have been a subject of extensive investigations for several decades due to their wide applications in different fields in day-to-day life.<sup>1–7</sup> They give intense sharp emissions in UV-vis-NIR regions and often lead to lasing, and Nd:YAG laser lasing at 1.06 μm is a good example of this.<sup>8</sup> The intensity of this laser in NIR region is large enough so that the frequency doubled, tripled and quadrupled laser emissions could be obtained at 0.532, 0.355 and 0.266 μm, respectively and they are used for various applications. The rare earth ions attached with biomolecules are used as biosensors.<sup>9,10</sup> The optical energy absorbed by biomolecules is transferred to the rare earth ion, which gives intense emission and thus, serves as a biosensor. The photoluminescence intensity of the rare earth ion in the host materials can be enhanced by co-

doping of another neutral/ionized species acting as a sensitizer/modifier for the rare earth ions. In several cases, the sensitizer plays a dominant role.<sup>11–15</sup> These materials have promising applications in different fields, such as display devices, light emitting diodes (LEDs), white LEDs, tunable color sources, optoelectronic devices, luminescent security ink, solar energy converters, optical heating, temperature sensing, bio-imaging, etc.<sup>1–15</sup>

The Eu<sup>3+</sup> ion is a well-studied rare earth ion. It emits almost pure red color and often used in display devices, color televisions and computer screens. It also gives laser emission in the far-infrared region. The Eu<sup>3+</sup> ion has been doped in large number of hosts including glass, micro- and nano-phosphors and also in the bio-mediums.<sup>7,16–25</sup> The ground state of Eu<sup>3+</sup> ion is <sup>7</sup>F<sub>J</sub> with <sup>7</sup>F<sub>0</sub> as the lowest one. The other component of it forms the low-lying excited states. The next low-lying excited state is regular <sup>5</sup>D<sub>J</sub> with <sup>5</sup>D<sub>0</sub> as the lowest one. The bands in visible region in it arise due to <sup>5</sup>D<sub>0,1</sub> → <sup>7</sup>F<sub>J</sub> transitions. The Eu<sup>3+</sup> ion emits very intense red band in almost all hosts in the range of 610–615 nm. The emitted radiation is mixed red because in many cases, it also emits intense orange at 590 nm and weak deep red near 650 nm and deep red at 700 nm. Ofcourse, the intensity of the red band is much larger than the orange band. Therefore, the Eu<sup>3+</sup> doped materials are sometimes treated as pure red source of light for many practical purposes. It is also

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used to generate white light (RGB) with other rare earth/earths emitting green and blue colors.<sup>1–6,16–25</sup> The  $^5D_0 \rightarrow ^7F_1$  transition lies in orange region whereas  $^5D_0 \rightarrow ^7F_3$  and  $^5D_0 \rightarrow ^7F_4$  in deep red (NIR) regions. However, these bands are relatively weak. The band  $^5D_0 \rightarrow ^7F_0$  appears with very small intensity and it is not seen in many cases. The  $^5D_0 \rightarrow ^7F_2$  transition is due to change in electric dipole moment whereas the  $^5D_0 \rightarrow ^7F_1$  transition occurs due to change in magnetic dipole moment. The ratio of the emission intensity of these bands *i.e.*,  $\frac{I(^5D_0 \rightarrow ^7F_2)}{I(^5D_0 \rightarrow ^7F_1)}$  gives the asymmetric nature of the host.<sup>16–21</sup> The  $\text{Eu}^{3+}$  ion has been studied in different phosphor host matrices (ref. 26–31 and references therein) in which some of them are found to be self-activated.<sup>26–29</sup>

The  $\text{LaVO}_4$  as a self-activated host absorbs UV light efficiently. It emits blue color in the range of 350–450 nm with its maximum  $\sim 430$  nm. The  $\text{Eu}^{3+}$  ion absorbs this radiation and emits photoluminescence in the range of 420–750 nm. This induced us to select this as a host matrix.<sup>32–34</sup>  $\text{LaVO}_4$  exists in monoclinic (m- $\text{LaVO}_4$ ) as well as in tetragonal (t- $\text{LaVO}_4$ ) phases. The formation of monoclinic or tetragonal phase depends on the temperature used for its preparation and also on the environment in which the samples have been prepared (in air or vacuum). Sometimes, it also depends on the doping of activator/sensitizer ions in the  $\text{LaVO}_4$  matrix.<sup>25,27,28</sup> We have used 1473 K temperature for the preparation of phosphor samples. The XRD measurements clearly show that the phosphor samples exist in monoclinic phase at this temperature.<sup>25,27,28,32–34</sup>

The triply ionized  $\text{Cr}^{3+}$  ion with configuration  $3d^3$  has been used as sensitizer as well as activators in different host matrices.<sup>35–38</sup> The  $^2E$  state of  $\text{Cr}^{3+}$  ion is a long-lived state, which gives long lasting phosphorescence.<sup>35,37</sup> It gives lasing at several wavelengths in IR and NIR regions in which the NIR emission at 695 nm is due to forbidden  $^2E \rightarrow ^4A_2$  transition and has been extensively studied in recent years.<sup>38</sup> The  $\text{Cr}^{3+}$  ion also gives the blue ( $^4T_1(^4P) \rightarrow ^4A_2$ )/( $^4T_1(^4F) \rightarrow ^4A_2$ ), green ( $^4T_2(^4P) \rightarrow ^4A_2$ ) and red ( $^4T_2(^4F) \rightarrow ^4A_2$ ) emissions. It will be interesting to see that the energy level  $^4T_2(^4F)$  of  $\text{Cr}^{3+}$  ion lies in the red region at  $15\,773\text{ cm}^{-1}$  while that of  $^4T_1(^4F)$  lies in the green region at  $17\,211\text{ cm}^{-1}$ . On the other hand, the  $^5D_0$  level of  $\text{Eu}^{3+}$  ion lies at  $17\,267\text{ cm}^{-1}$ . Thus, when  $\text{Eu}^{3+}$  and  $\text{Cr}^{3+}$  ions are present in the host the  $^4T_1(^4F)$  of  $\text{Cr}^{3+}$  ion lies close but slightly below to the  $^5D_0$  level of  $\text{Eu}^{3+}$  ion. Thus, when  $\text{Eu}^{3+}$  ions are excited they transfer a small amount of their energy to  $\text{Cr}^{3+}$  ions and therefore, the emission intensity of  $\text{Eu}^{3+}$  ion as well as its lifetime are decreased. The presence of excess  $\text{Cr}^{3+}$  ion may also quench its own emission intensity.

In the present work, we have doped different concentrations of  $\text{Cr}^{3+}$  ions in the  $\text{LaVO}_4$ :1 mol%  $\text{Eu}^{3+}$  phosphor. Initially, the  $\text{Eu}^{3+}$  doped  $\text{LaVO}_4$  has been studied and its concentration was optimized for optimum emission intensity. It has been found that the emission intensity is optimum at 1 mol% concentration of  $\text{Eu}^{3+}$  ion. Further, the samples were prepared using  $\text{LaVO}_4$ :1 mol%  $\text{Eu}^{3+}$  phosphor with different concentrations of  $\text{Cr}^{3+}$  ions (where  $x = 0.01, 0.05, 0.1, 0.5, 1.0$  mol%). We have studied the effect of  $\text{Cr}^{3+}$  doping on the absorption and

photoluminescence emission intensity of  $\text{LaVO}_4$ :1 mol%  $\text{Eu}^{3+}$  phosphor. It has been found that the emission intensity of  $\text{LaVO}_4$ :1 mol%  $\text{Eu}^{3+}$  phosphor decreases even with small concentration of  $\text{Cr}^{3+}$  ions. We have also measured the lifetime of  $^5D_0$  level of  $\text{Eu}^{3+}$  ion corresponding to  $^5D_0 \rightarrow ^7F_2$  band at 614 nm. The lifetime of  $^5D_0$  level initially increases and then decreased with the increase in the concentration of  $\text{Eu}^{3+}$  ion. Further, the lifetime of  $^5D_0$  level decreases on increasing the concentration of  $\text{Cr}^{3+}$  ion, which is due to energy transfer from  $\text{Eu}^{3+}$  to  $\text{Cr}^{3+}$  ions. This supports that the  $\text{Cr}^{3+}$  ion behaves as a quenching center. We have also measured the heat produced in the  $\text{Eu}^{3+}$ ,  $\text{Cr}^{3+}$  co-doped  $\text{LaVO}_4$  phosphor on excitation with 980 nm diode laser.

## 2. Experimental methods

### 2.1. Synthesis and characterization of samples

The  $\text{La}_{(1-x)}\text{Eu}_x\text{VO}_4$  (where  $x = 0, 0.5, 1.0, 1.5$  and  $2$  mol%) and  $\text{La}_{(1-x-y)}\text{Eu}_x\text{Cr}_y\text{VO}_4$  (where  $x$  is fixed at 1 mol% and  $y$  was varied as 0.01, 0.05 and 0.1 mol%) phosphor samples were synthesized by a high temperature solid-state reaction method at 1473 K. The raw materials *viz.*  $\text{La}_2\text{O}_3$  (99.9% Himedia),  $\text{V}_2\text{O}_5$  (99.9% Alpha Aesar),  $\text{Eu}_2\text{O}_3$  (99.99% Merck) and  $\text{Cr}_2\text{O}_3$  (99.99% alpha Aesar) were used as the starting materials. The calculated amounts of all the raw materials were weighed and mixed homogeneously with the help of an agate mortar and pestle for one hour each using acetone as mixing agent. The prepared mixtures were put in alumina crucibles and calcined in a high temperature furnace at 1473 K for 5 h. After that, the furnace temperature was reduced till it reaches to the room temperature and products were taken out from different crucibles. The phosphor samples were then crushed to convert them into fine powder form. The structural and optical measurements of these phosphor samples were carried out separately and compared with each other.

The XRD measurements of  $\text{Eu}^{3+}$ ,  $\text{Cr}^{3+}$  doped and co-doped  $\text{LaVO}_4$  phosphors monitored in the  $2\theta$  range of  $10^\circ$ – $100^\circ$  were carried out using MiniFlex 600 (Rigaku Japan) X-ray diffractometer at  $2^\circ$  per min scan speed with  $\text{CuK}_\alpha$  X-ray radiation source ( $\lambda = 1.5406\text{ \AA}$ ). The Rietveld refinements of all the XRD data were made using FullProf Suite.<sup>32–34</sup> In order to measure the morphology and the energy dispersive X-ray spectroscopic (EDS) spectra of all the samples, a Zeiss EVO 18 Research unit of the scanning electron microscope (SEM) system was used. The absorption spectra of samples were recorded in the range in 200–800 nm using Lambda-750 PerkinElmer UV-vis-NIR spectrophotometer unit. The Fourier transform infrared (FTIR) spectra of samples were also measured with the help of Frontier-1 spectrometer, PerkinElmer unit. The photoluminescence excitation and emission spectra of the samples were monitored using Fluorolog-3 (Horiba) attached with 450 W Xenon lamp and photomultiplier tube (PMT). The lifetime of  $^5D_0$  level of the  $\text{Eu}^{3+}$  ion at 614 nm was also measured using the same unit in phosphorescence mode using 25 W pulsed Xenon lamp. The heat generated in all the samples were also measured using a thermo-couple set up at different power densities of 980 nm diode laser.



### 3. Results and discussion

#### 3.1. Crystal structure and rietveld analysis

For the structural analysis of  $\text{LaVO}_4:1\text{Eu}^{3+}:y\text{Cr}^{3+}$  phosphors with  $y = 0.01, 0.05$  and  $0.1$  mol%, the powder X-ray diffraction (XRD) patterns of these phosphor samples were monitored at room-temperature and are presented in Fig. 1. The powder XRD patterns of these phosphors are very sharp and highly intense, which reveal that the crystallinity of the  $\text{LaVO}_4:1\text{Eu}^{3+}:y\text{Cr}^{3+}$  phosphors are very high.

The diffraction peaks in the powder XRD patterns of all the compounds, well match with the standard XRD data of monoclinic  $\text{LaVO}_4$  compound belonging to  $P2_1/n$  space group with JCPDF File No. 50-0367 alongwith lattice parameters  $a = 7.0433(1)$  Å,  $b = 7.2806(2)$  Å,  $c = 6.7219(1)$  Å and  $\beta = 104.834(2)^\circ$  (see Fig. 1). This implies that all  $\text{Eu}^{3+}$  and  $\text{Cr}^{3+}$  co-doped  $\text{LaVO}_4$  phosphors crystallize into monoclinic crystal structure belonging to  $P2_1/n$  space group.<sup>32–34</sup> No traces of any secondary phase in the XRD profile of these compounds are seen. The XRD patterns shift towards higher  $2\theta$  value with the increase of concentration of  $\text{Cr}^{3+}$  ion.

In order to find out different crystal structure parameters, Rietveld structural analysis of the XRD patterns of all the phosphors have been carried out by the use of FullProf Suite.<sup>39,40</sup> In the course of Rietveld structural analysis, we had supposed that the  $\text{Eu}^{3+}$  and/or  $\text{Cr}^{3+}$  ions are substituting at La-site in the  $\text{LaVO}_4$  host. In the monoclinic crystal structure having  $P2_1/n$  space group,  $\text{V}^{5+}$ ,  $\text{La}^{3+}/\text{Eu}^{3+}/\text{Cr}^{3+}$ ,  $\text{O}^{2-}(\text{i})$ ,  $\text{O}^{2-}(\text{ii})$ ,  $\text{O}^{2-}(\text{iii})$  and  $\text{O}^{2-}(\text{iv})$  ions substitute at 4e site ( $\delta x$ ,  $\delta y$ ,  $\delta z$ ). Further, for Rietveld structural analysis, we have used atomic coordinates for constituent atoms as reported earlier.<sup>32–34</sup> A good fit between the calculated and the observed XRD patterns was observed. Rietveld plots for the  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphors are displayed in Fig. 2a–d, respectively. In Fig. 2, solid red dots designate the observed XRD

patterns and the continuous curve over the observed XRD patterns indicates the calculated patterns. The vertical bars in Rietveld fits demonstrate the positions of Bragg's peak and the lower most continuous curve signifies the difference among the observed and the calculated XRD patterns.

The obtained values of goodness of fit ( $\chi^2$ ), lattice parameters ( $a$ ,  $b$ ,  $c$ ,  $\alpha$ ,  $\beta$  and  $\gamma$ ) and the unit cell volume ( $V$ ) for all the doped and co-doped phosphor samples are given in Table 1. It has been observed that the value of unit cell volume decreases as the concentrations of  $\text{Cr}^{3+}$  ions are enhanced. Reduction in the unit cell volume of the  $\text{Cr}^{3+}$  doped  $\text{LaVO}_4:1\text{Eu}^{3+}$  phosphors may be due to the smaller ionic radius of  $\text{Cr}^{3+}$  ion (0.615 Å) as compared to the ionic radius of  $\text{La}^{3+}$  ion (1.216 Å) and  $\text{Eu}^{3+}$  ion (1.120 Å).<sup>34</sup> The  $\text{Cr}^{3+}$  ion would prefer to go at the La-site to maintain the charge balance in the  $\text{LaVO}_4$  phosphor.

#### 3.2. Microstructural analysis

The microstructure of all the phosphor samples has been studied with the help of scanning electron microscopy (SEM) images, at a magnification of 10 K $\times$  and a scale of 2  $\mu\text{m}$ . Fig. 3a–d displays SEM micrographs for the  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor powder samples, respectively. The SEM images of the samples reveal that the particles size of the phosphor samples decreases as the doping concentration of the  $\text{Cr}^{3+}$  ions is increased. The particles of these phosphors are somewhat aggregated and most of the particles are spherical in shape (see Fig. 3). The particles size of the phosphor samples in general lies in the sub-micron to micron range; however, the average particles size in the present case is in the micron range. Thus, the average particles size of the phosphor samples has been calculated using ImageJ software. The mean value of the particles size was found to be 4.05, 3.47, 3.18 and 2.89  $\mu\text{m}$  for the  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphors, respectively. This clearly shows that the particles size reduces on doping/co-doping of  $\text{Eu}^{3+}$  and  $\text{Cr}^{3+}$  ions.

#### 3.3. Compositional analysis

Qualitative compositional analyses of the  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples have been carried out by monitoring energy dispersive X-ray spectroscopy (EDS) spectra of the phosphor samples at the time during the SEM micrographs measurements. The EDS spectra of the  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphors are presented in Fig. 4a–d, respectively. The occurrence of the characteristic peaks corresponding to the component atoms, *i.e.*, La, Eu/Cr, V and O in the EDS spectra exhibits the presence of the component elements. As the EDS method is not suitable for the quantitative analysis of the light elements only qualitative analysis was performed.

Fig. 5 displays the selected area used for elemental mapping for the elements La, Eu, Cr, V and O present in the  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor. The elemental mapping clearly demonstrates the homogeneous distribution of the elements (*i.e.*, La, Eu, Cr, V and O) present in the phosphor sample.

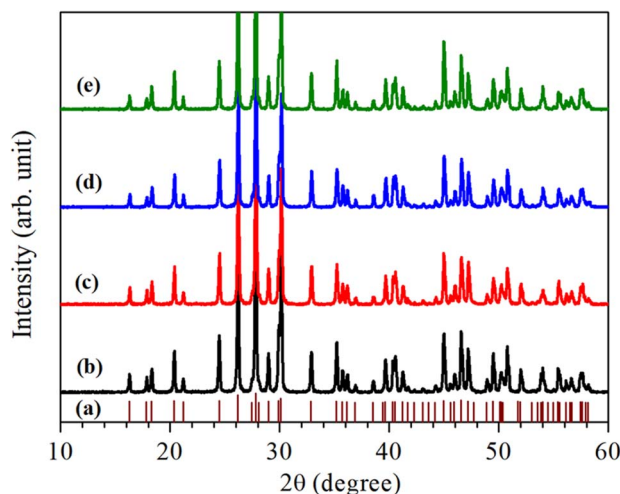


Fig. 1 Room-temperature XRD patterns of  $\text{Eu}^{3+}$  and/or  $\text{Cr}^{3+}$  co-doped  $\text{LaVO}_4$  phosphors: (a) Standard XRD pattern of  $\text{LaVO}_4$  (JCPDF File No. 50-0367), (b)  $\text{LaVO}_4:1\text{Eu}^{3+}$ , (c)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ , (d)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and (e)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphors.



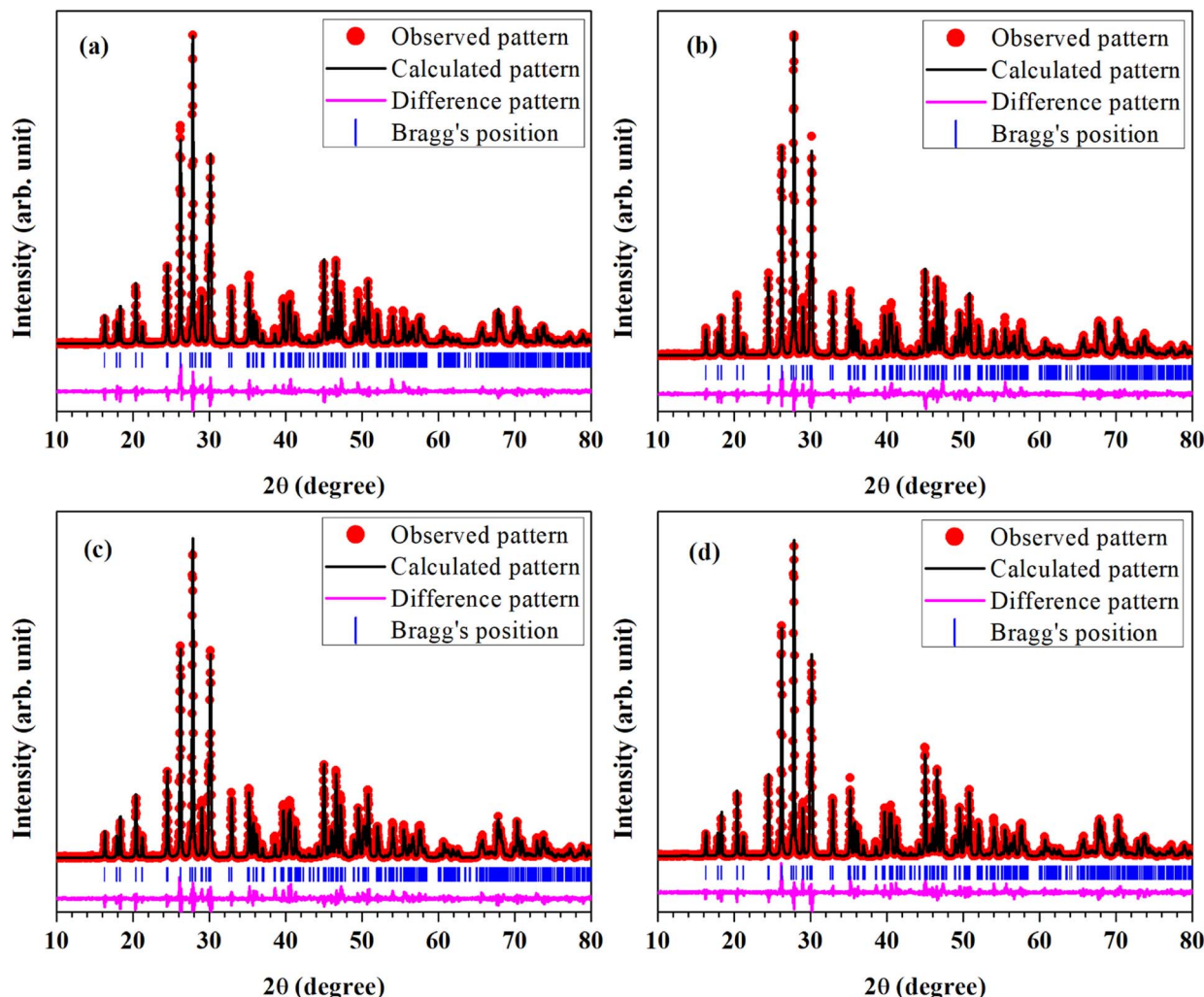


Fig. 2 Rietveld fits of XRD patterns for  $\text{Eu}^{3+}$  and/or  $\text{Cr}^{3+}$  co-doped  $\text{LaVO}_4$  phosphors: (a)  $\text{LaVO}_4:1\text{Eu}^{3+}$ , (b)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ , (c)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and (d)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphors.

Table 1 Refined value of goodness of fit ( $\chi^2$ ), lattice parameters ( $a$ ,  $b$ ,  $c$ ,  $\alpha$ ,  $\beta$  and  $\gamma$ ) and the unit cell volume ( $V$ ) of the monoclinic structure for  $\text{Eu}^{3+}$  and  $\text{Eu}^{3+} + \text{Cr}^{3+}$  doped and co-doped  $\text{LaVO}_4$  phosphors

Compound	Lattice parameters ( $\alpha = \gamma = 90^\circ$ )				Unit cell volume $V (\text{\AA}^3)$	$\chi^2$
	$a (\text{\AA})$	$b (\text{\AA})$	$c (\text{\AA})$	$\beta (^\circ)$		
$\text{LaVO}_4:1\text{Eu}^{3+}$	7.0388(2)	7.2771(2)	6.7200(2)	104.831(2)	332.75(2)	1.90
$\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$	7.0393(1)	7.2766(1)	6.7194(1)	104.837(2)	332.71(1)	1.76
$\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$	7.0389(2)	7.2762(2)	6.7195(2)	104.839(2)	332.67(2)	1.78
$\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$	7.0390(2)	7.2760(2)	6.7191(2)	104.837(2)	332.65(2)	1.63

### 3.4. Optical measurements

**3.4.1 Fourier transform infrared (FTIR) spectra.** The FTIR spectra of the  $\text{LaVO}_4$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples have been recorded in the spectral range of 400–4000  $\text{cm}^{-1}$  and they are shown in Fig. 6. There is a variation in intensity of different absorption bands due to variation of charge distribution of  $\text{Cr}^{3+}$  and  $\text{Eu}^{3+}$  ions in the host. All the

samples show three clear peaks near 431, 755 and 804  $\text{cm}^{-1}$ . These peaks are due to La–O stretching vibration; symmetric and asymmetric stretching vibrations of V–O bonds in  $\text{VO}_4^{3-}$  group, respectively. In fact, there are large number of weak peaks near 804  $\text{cm}^{-1}$  band due to  $\text{VO}_4^{3-}$  group.<sup>32–34</sup>

**3.4.2 UV-vis absorption spectra of phosphor samples.** The UV-vis absorption spectra of  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples have been monitored in diffuse reflectance



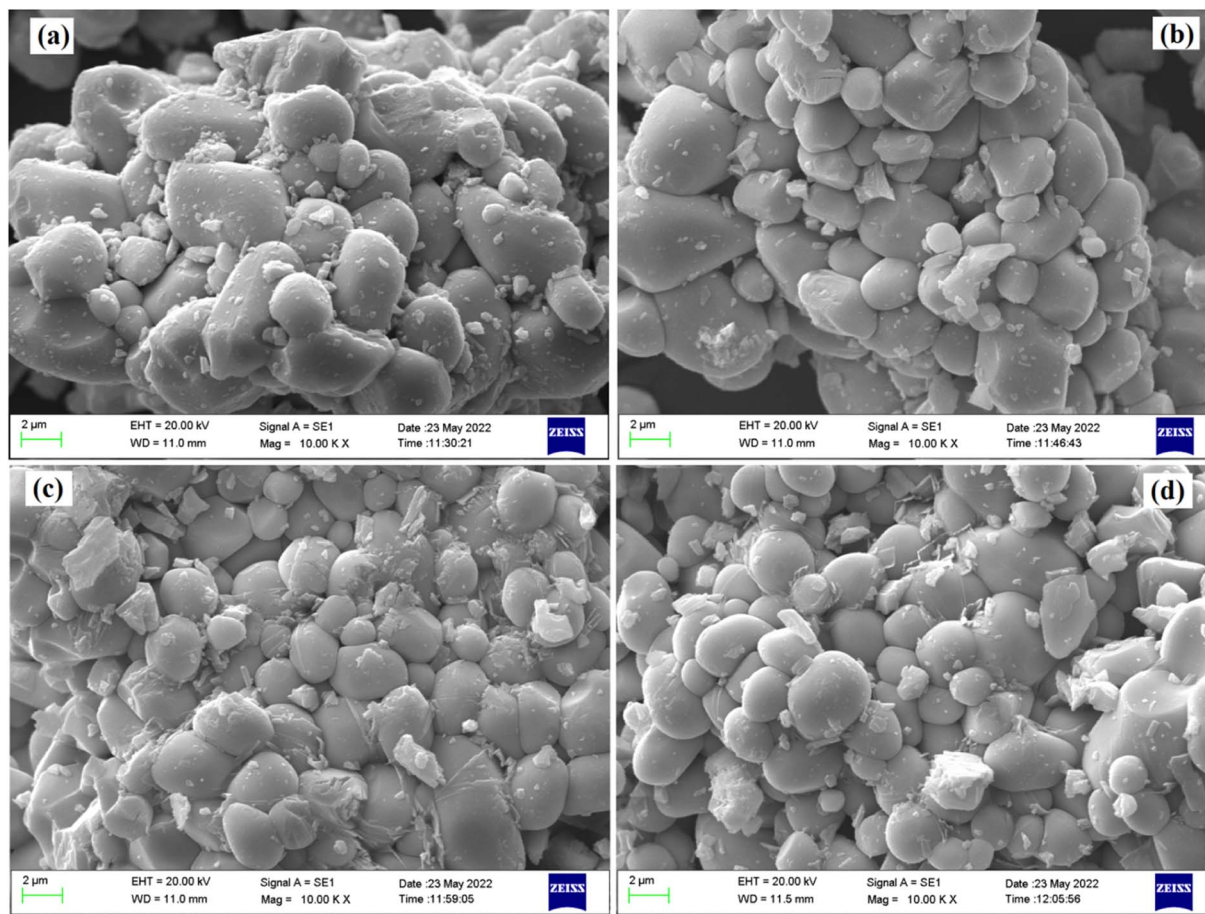


Fig. 3 Scanning electron microscopic images for the  $\text{Eu}^{3+}$  and  $\text{Eu}^{3+} + \text{Cr}^{3+}$  co-doped  $\text{LaVO}_4$  phosphors; (a)  $\text{LaVO}_4:1\text{Eu}^{3+}$ , (b)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ , (c)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and (d)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphors.

mode in the spectral region 200–400 nm and the resulting spectra are shown in Fig. 7. The spectra contain two intense broad bands at 254 and 316 nm. The intensity of these bands varies with  $\text{Cr}^{3+}$  concentrations; however, the peak positions remain unchanged. These bands arise due to charge transfer transition of  $\text{O}^{2-} \rightarrow \text{V}^{5+}$  ions and from  $\text{O}^{2-} \rightarrow \text{Eu}^{3+}$  ions in the  $\text{LaVO}_4$  phosphors. The doping of  $\text{Cr}^{3+}$  ion in the  $\text{Eu}^{3+}$  doped  $\text{LaVO}_4$  phosphor causes a field effect on the absorption intensity of  $\text{O}^{2-} \rightarrow \text{V}^{5+}$  transition. The ground state of  $(\text{VO}_4)^{3-}$  ion is  ${}^1\text{A}_2({}^1\text{T}_1)$ . The observed bands arise due to transition from ground state to the close-lying electronic states  ${}^1\text{A}_1({}^1\text{E})$  and  ${}^1\text{E}({}^1\text{T}_1)$ . The charge transfer transition (CTS) of  $\text{O}^{2-} \rightarrow \text{Eu}^{3+}$  also lies in 270–325 nm range and overlaps on  $(\text{VO}_4)^{3-}$  bands.<sup>17–19,32–34</sup>

We have also monitored the absorption spectra of the samples in Kubelka–Munk (KM) mode to determine the optical band gap of  $\text{LaVO}_4$  and its dependence on different doping concentrations of  $\text{Cr}^{3+}$  ions. The Kubelka–Munk (KM) function is given as:<sup>39–42</sup>

$$F(R) = \frac{(1 - R_\infty)^2}{2R_\infty} \quad (\text{i})$$

where,  $R_\infty$  represents the diffuse reflectance intensity of the samples normalized to the non-absorbing standard samples. The KM function  $F(R)$  is related to the band gap ( $E_g$ ) as:

$$F(R) = \frac{B(h\nu - E_g)^m}{h\nu} \quad (\text{ii})$$

This relation is also called as Wood and Tauc formula (41) where  $E_g$ ,  $h\nu$  and  $B$  specify the optical band gap, energy of incident photon and the band tailoring parameter, respectively.

The value of  $m$  is taken  $\frac{1}{2}$  supposing the case to be of direct optical band gap.<sup>39–42</sup> Therefore,

$$\frac{[F(R) \cdot h\nu]}{B} = (h\nu - E_g)^{\frac{1}{2}} \quad (\text{iii})$$

or

$$\left[ \frac{F(R) \cdot h\nu}{B} \right]^2 = (h\nu - E_g) \quad (\text{iv})$$

If we assume  $\frac{F(R)}{B} = \alpha$ , then  $(\alpha h\nu)^2 = (h\nu - E_g)$

A graph between  $[\alpha h\nu]^2$  versus  $h\nu$  also called as Wood and Tauc plot for different phosphor samples are shown in Fig. 8. The intercepts of the tangents on the curves on X-axis directly give the band gap. The band gap values for different concentrations of  $\text{Cr}^{3+}$  ions are given in the figure. This clearly shows



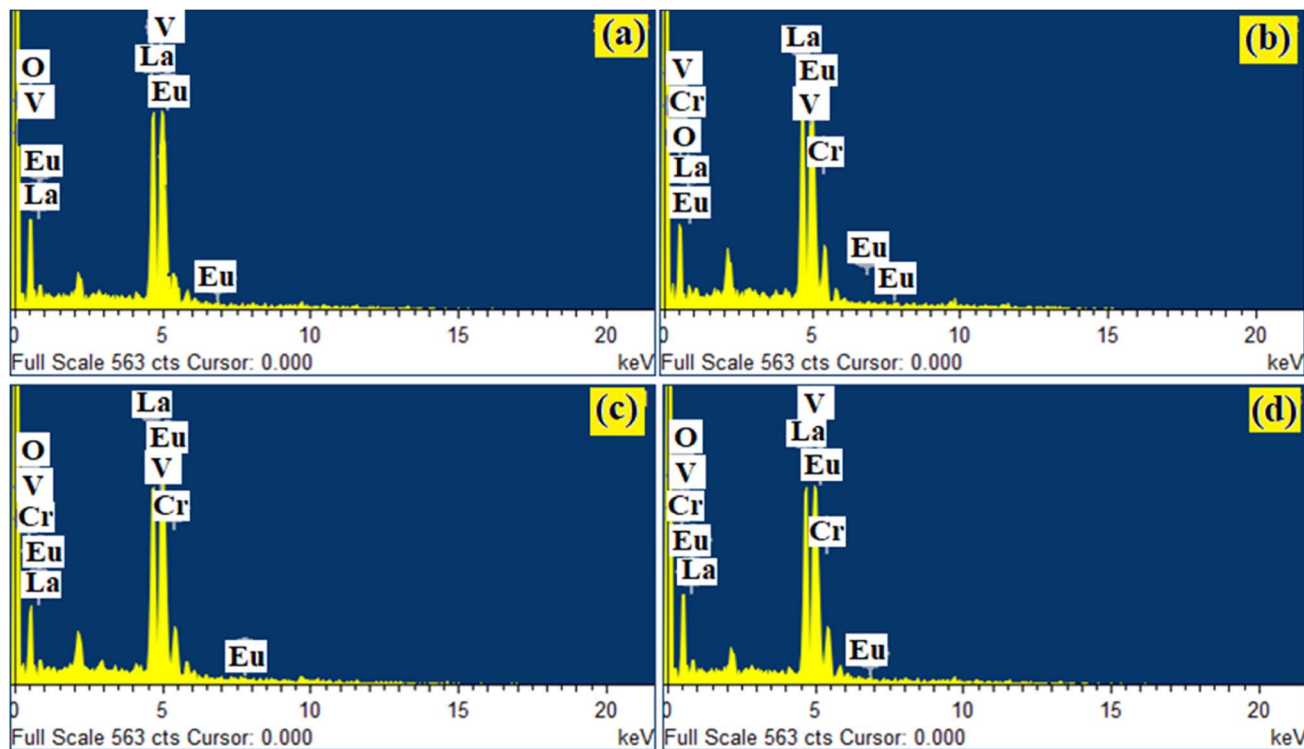


Fig. 4 EDS spectra for  $\text{Eu}^{3+}$  and  $\text{Eu}^{3+} + \text{Cr}^{3+}$  co-doped  $\text{LaVO}_4$  phosphors *i.e.* (a)  $\text{LaVO}_4:1\text{Eu}^{3+}$ , (b)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ , (c)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and (d)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphors.

that the optical band gap decreases with the increase in the concentration of  $\text{Cr}^{3+}$  ion, which is due to metallic nature of phosphor samples.<sup>43,44</sup>

### 3.4.3 Photoluminescence excitation and emission spectra.

The photoluminescence excitation (PLE) spectra of  $\text{LaVO}_4$  doped with different concentrations of  $\text{Eu}^{3+}$  ions ( $x = 0.5, 1.0, 1.5, 2.0$  mol%) have been monitored in the spectral region of 200–575 nm with  $\lambda_{\text{em}}$  fixed at 614 nm. The PLE spectra thus obtained are shown in Fig. 9. A broad band is observed between 230–350 nm with its maximum at 316 nm. The maxima are slightly shifted towards higher wavelength side from 316 to 323 nm on increasing concentration of  $\text{Eu}^{3+}$  ion. The corresponding broad band is due to overlapping between two charge transfer bands (CTBs) *i.e.* ( $\text{O}^{2-} \rightarrow \text{V}^{5+}$ ) and ( $\text{O}^{2-} \rightarrow \text{Eu}^{3+}$ ). The spectra also contain many sharp weak peaks at 362, 376, 381, 393, 413 and 464 nm, which are due to  ${}^7\text{F}_0 \rightarrow {}^5\text{D}_4$ ,  ${}^7\text{F}_0 \rightarrow {}^5\text{L}_8$ ,  ${}^7\text{F}_0 \rightarrow {}^5\text{L}_7$ ,  ${}^7\text{F}_0 \rightarrow {}^5\text{L}_6$ ,  ${}^7\text{F}_0 \rightarrow {}^5\text{D}_3$  and  ${}^7\text{F}_0 \rightarrow {}^5\text{D}_2$  transitions of  $\text{Eu}^{3+}$  ion, respectively.<sup>11–23,32,33</sup> The intensity of these peaks follows the trend  $\lambda_{393} > \lambda_{464} > \lambda_{362}$ . The peak at 393 nm lies in the region of n-UV LED; therefore, it is very important. The intensity of charge transfer band (CTB) is much larger than the intensity of discrete bands due to 4f–4f transitions of  $\text{Eu}^{3+}$  ion. The intensity of excitation band is optimum for 1 mol% concentration of  $\text{Eu}^{3+}$  ion.

We have monitored the photoluminescence (PL) emission spectra of  $\text{LaVO}_4:x\text{Eu}^{3+}$  (where  $x = 0.5, 1.0, 1.5$  and  $2.0$  mol%  $\text{Eu}^{3+}$ ) exciting at 316 and 393 nm wavelengths in the range 500–800 nm. In all the cases, we have observed intense bands in

between 520–700 nm regions due to  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_j$  transitions.<sup>11–23,32,33</sup> The intensity of emission bands on excitation with CTB wavelength is nearly 10 times larger than on excitation with discrete bands of  $\text{Eu}^{3+}$  ion *i.e.* at 393 nm. The photoluminescence emission spectra on excitation with these wavelengths are shown in Fig. 10a and b.

As mentioned earlier, the bands observed between 570–700 nm are due to  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_j$  transitions of  $\text{Eu}^{3+}$  ion in which the band at 614 nm due to  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition is very intense. The other observed bands are due to  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_0$  (579 nm),  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_1$  (582 and 591 nm),  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_3$  (644 nm) and  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_4$  (700 nm) transitions.<sup>11–23,32,33,45,46</sup> The intensity of the bands follows the  ${}^7\text{F}_2 > {}^7\text{F}_1 > {}^7\text{F}_4 > {}^7\text{F}_3 > {}^7\text{F}_0$  trends. Though the excitation wavelengths (316 and 393 nm) populate high lying levels of  $\text{Eu}^{3+}$  ion; however, due to closeness of the energy levels, the excited  $\text{Eu}^{3+}$  ions relax rapidly to low lying  ${}^5\text{D}_1$  and  ${}^5\text{D}_0$  excited states. It is due to this reason that the observed emission bands are arising from these two lower lying levels. The weak emission bands in the green region are due to  ${}^5\text{D}_1 \rightarrow {}^7\text{F}_0$  (525 nm),  ${}^5\text{D}_1 \rightarrow {}^7\text{F}_1$  (537 and 531 nm) and  ${}^5\text{D}_1 \rightarrow {}^7\text{F}_2$  (565 nm) transitions, which are reported only in few host materials.<sup>46,47</sup> The emission bands from  ${}^5\text{D}_1$  level is observed only on excitation with 316 nm wavelength.

The  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition occurs due to change in electric dipole moment. Its intensity depends on the external conditions. However, the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_1$  transition is magnetic dipole allowed transition and it is independent of the local environment round the  $\text{Eu}^{3+}$  ions. The  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_3$  has an electric and



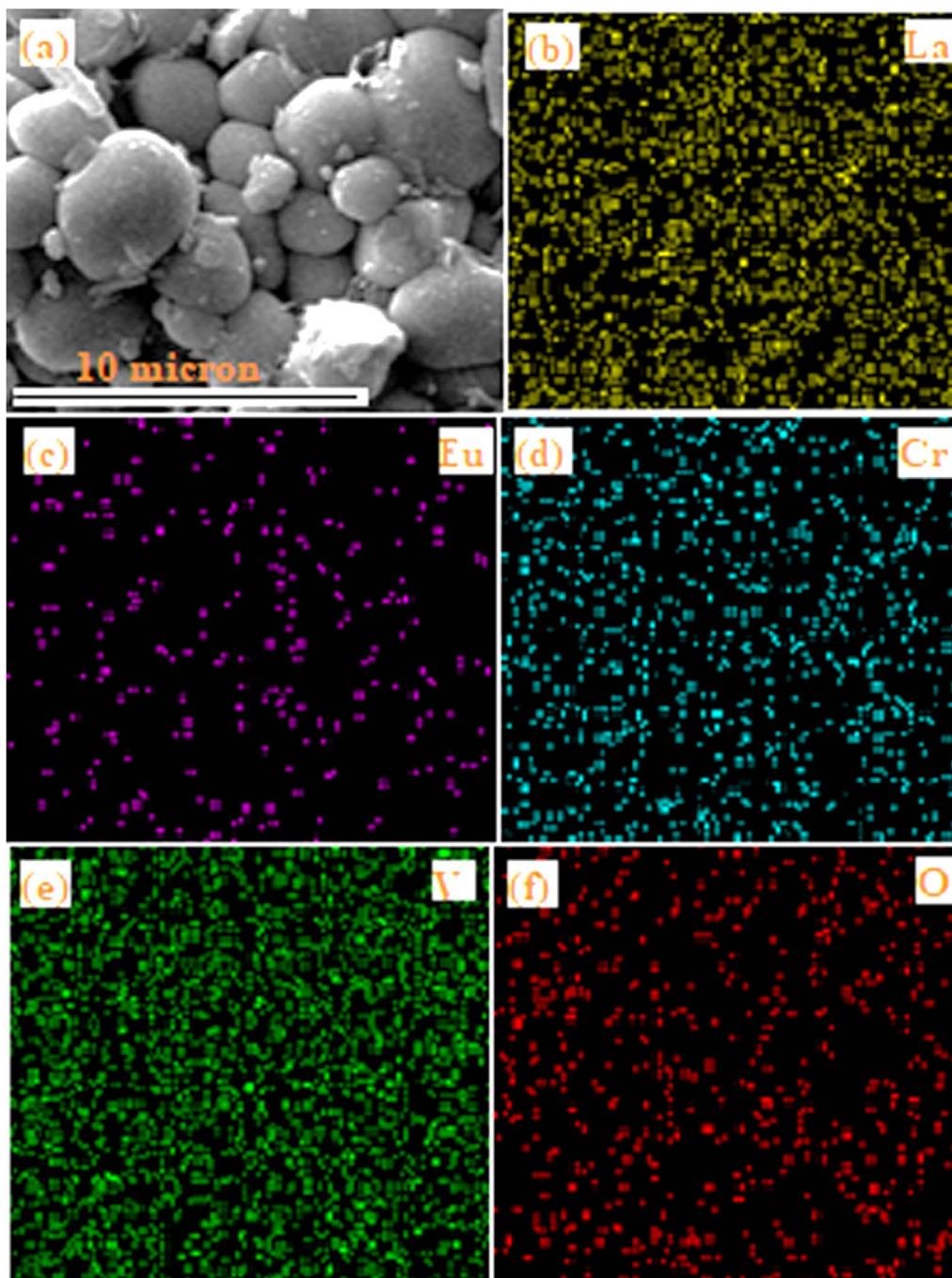


Fig. 5 Elemental mapping and distribution of the component elements present in the  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor: (a) selected area used for elemental mappings, (b) La, (c) Eu, (d) Cr, (e) V and (f) O elements.

magnetic dipole mixed character. On the other hand,  $^5\text{D}_0 \rightarrow ^7\text{F}_4$  is forced electric dipole transition. Thus, the electric dipole allowed transitions have larger intensity compared to the magnetic dipole allowed transitions.<sup>11–23,32,33</sup> Actually, the doping of  $\text{Eu}^{3+}$  ion in the  $\text{LaVO}_4$  host replaces  $\text{La}^{3+}$  ion and forms  $\text{EuO}_9$  instead of  $\text{LaO}_9$  in the case of monoclinic structure of  $\text{LaVO}_4$  host. The  $\text{EuO}_9$  in  $\text{LaVO}_4$  host forms a highly asymmetric structure, which is the basic reason for large PL intensity.

**3.4.4 Effect of  $\text{Cr}^{3+}$  doping on PL intensity.** The PLE spectra of  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples have been monitored by fixing  $\lambda_{\text{emi}} = 614$  and  $700$  nm. The  $\text{Cr}^{3+}$  ion gives an intense emission at  $700$  nm due to forbidden  $^2\text{E} \rightarrow ^4\text{A}_2$  transition. There is also a weak  $^5\text{D}_0 \rightarrow ^7\text{F}_4$  transition due to  $\text{Eu}^{3+}$  ion at the same wavelength. We have monitored the PLE spectra taking  $\lambda_{\text{emi}} = 700$  nm just to see, is there any other excitation band due to  $\text{Cr}^{3+}$  ion in this region. The PLE spectra of the



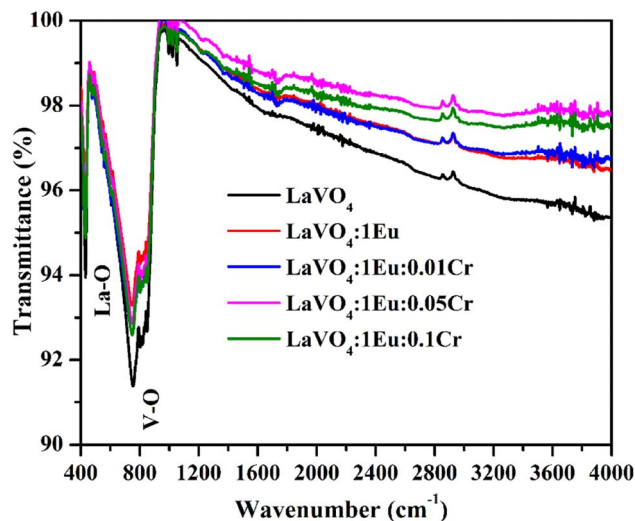


Fig. 6 FTIR spectra of  $\text{LaVO}_4$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples.

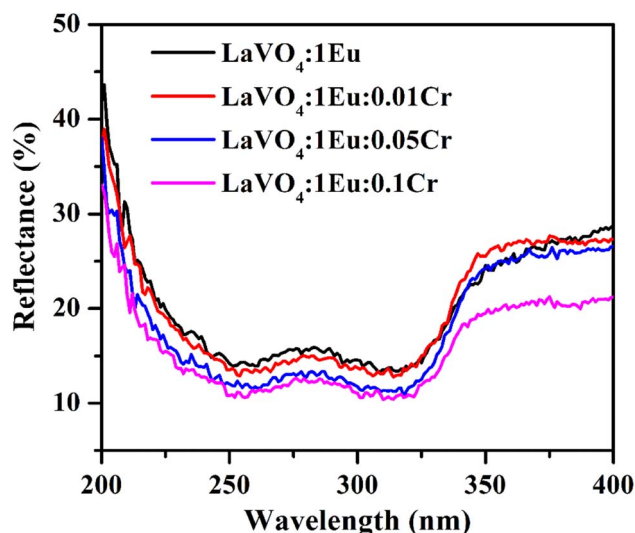


Fig. 7 The UV-vis absorption spectra of  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples.

samples for  $\lambda_{\text{emi}} = 614$  and  $700$  nm are shown in Fig. 11a and b. The PLE spectra appear almost identical in the two cases though there is a decrease in the PLE intensity for  $\lambda_{\text{emi}} = 700$  nm. An additional peak also appears near  $386$  nm, which have been taken due to  $\text{Cr}^{3+}$  ion as there is a band due to  $\text{Cr}^{3+}$  ion at this wavelength. The position of peaks remains unchanged; however, the intensity of the peaks varies with the concentration of  $\text{Cr}^{3+}$  ion.

The PL spectra of  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples were monitored on excitation with  $393$  and  $386$  nm and they are shown in Fig. 12a and b. The PL spectra of these phosphor samples in the range of  $420$ – $750$  nm show similar emission bands on excitation with  $393$  nm (see Fig. 10b).

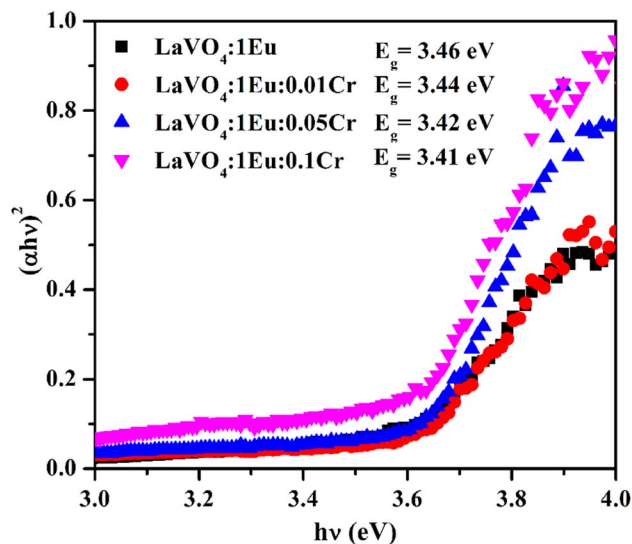


Fig. 8 Optical band gap of  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples.

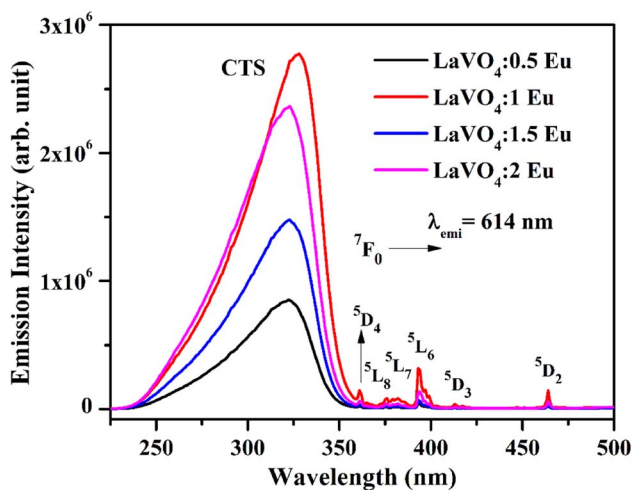


Fig. 9 Photoluminescence excitation (PLE) spectra of  $\text{LaVO}_4:x\text{Eu}^{3+}$  (where  $x = 0.5, 1.0, 1.5, 2.0$  mol%) phosphors monitored at  $\lambda_{\text{emi}} = 614$  nm.

However, some additional emission bands are observed at  $431$  and  $536$  nm, which are assigned to arise due to  ${}^5\text{D}_3 \rightarrow {}^7\text{F}_2$  and  ${}^5\text{D}_1 \rightarrow {}^7\text{F}_1$  transitions of  $\text{Eu}^{3+}$  ion, respectively. The emission intensity of  $\text{Cr}^{3+}$  doped  $\text{LaVO}_4:1\text{Eu}^{3+}$  phosphor is slightly reduced on excitation with these wavelengths. It is also clear from Fig. 12a that the pure  $\text{LaVO}_4:1\text{Eu}^{3+}$  phosphor does not contain any emission band due to  $\text{Cr}^{3+}$  ion. When  $\text{Cr}^{3+}$  ion is doped in the  $\text{LaVO}_4:1\text{Eu}^{3+}$  phosphor; an additional broad emission band is observed in the range of  $700$ – $735$  nm, which occurs due to  ${}^2\text{E} \rightarrow {}^4\text{A}_2$  transition of  $\text{Cr}^{3+}$  ion.<sup>34,38</sup> This additional band alongwith the other emission bands are clearly observed by exciting the phosphor samples at  $386$  nm in the range of  $400$ – $750$  nm. In this case, the PL spectra contain additional broad emission bands at  $417$ ,  $518$  and ( $700$  and  $725$





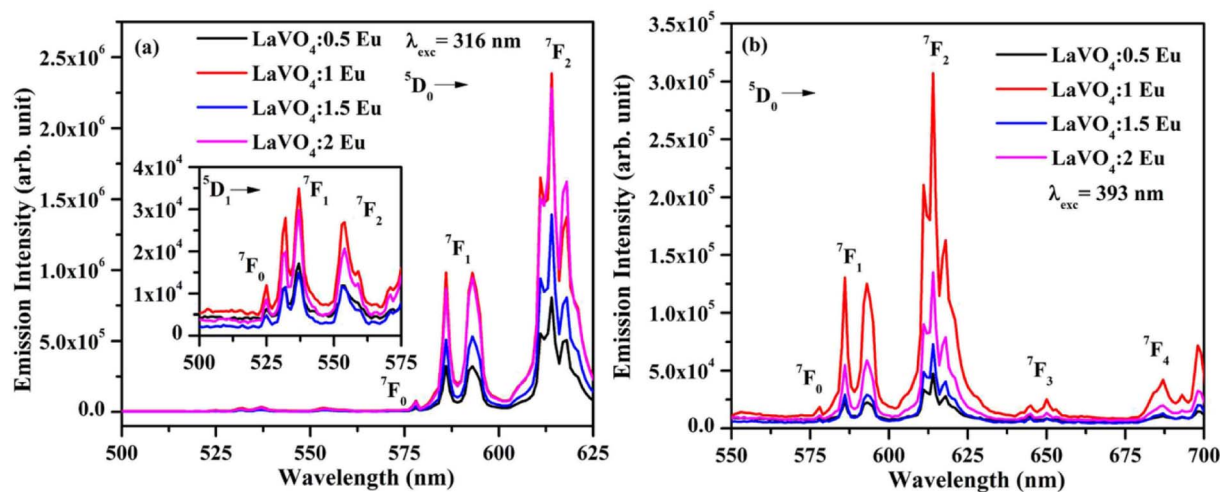


Fig. 10 Photoluminescence (PL) emission spectra of  $\text{LaVO}_4:x\text{Eu}^{3+}$  phosphors on excitation with (a) 316 nm and (b) 393 nm (where  $x = 0.5, 1.0, 1.5, 2.0$  mol%).

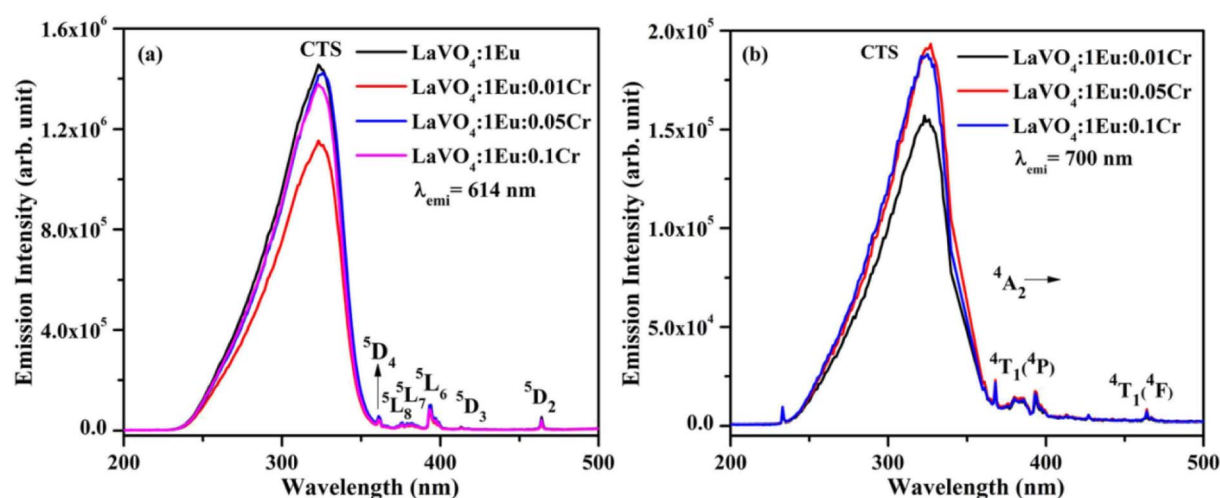


Fig. 11 Photoluminescence excitation (PLE) spectra of  $\text{LaVO}_4:1\text{Eu}^{3+}:y\text{Cr}^{3+}$  (where  $y = 0, 0.01, 0.05, 0.1$  mol%) phosphors monitored at  $\lambda_{\text{emi}} =$  (a) 614 nm and (b) 700 nm.

nm) clearly due to  $\text{Cr}^{3+}$  ion with weak PL intensity (see Fig. 12b) and they are assigned to arise due to  ${}^4\text{T}_1(4\text{P}) \rightarrow {}^4\text{A}_2$ ,  ${}^4\text{T}_2(4\text{P}) \rightarrow {}^4\text{A}_2$  and  ${}^2\text{E} \rightarrow {}^4\text{A}_2$  transitions of  $\text{Cr}^{3+}$  ion, respectively.<sup>34,38</sup>

The  $\text{LaVO}_4:1\text{Eu}^{3+}$  phosphor gives poor PL intensity due to the reason that 386 nm wavelength also excites the  $\text{Eu}^{3+}$  ion weakly. However, the PL intensity of 614 nm band is relatively larger than that of 700 nm band. The increasing concentration of  $\text{Cr}^{3+}$  ion reduces the PL intensity of  $\text{Eu}^{3+}$  as well as  $\text{Cr}^{3+}$  emission bands. This is due to energy transfer from  $\text{Eu}^{3+}$  to  $\text{Cr}^{3+}$  ions. Actually, the energy level  ${}^4\text{T}_2(4\text{F})$  of  $\text{Cr}^{3+}$  ion lies in the red region at  $15773\text{ cm}^{-1}$  while that of  ${}^4\text{T}_1(4\text{F})$  lies in the green region at  $17211\text{ cm}^{-1}$ . On the other hand, the  ${}^5\text{D}_0$  level of  $\text{Eu}^{3+}$  ion is found to lie at  $17267\text{ cm}^{-1}$ . The  ${}^4\text{T}_1(4\text{F})$  level of  $\text{Cr}^{3+}$  ion lies close but slightly below to the  ${}^5\text{D}_0$  level of  $\text{Eu}^{3+}$  ion. On excitation of  $\text{Eu}^{3+}$  ion, a small amount of energy is transferred to  $\text{Cr}^{3+}$  ion, which decreases the PL intensity of  $\text{Eu}^{3+}$  ion.<sup>47,48</sup> This clearly shows

that the  $\text{Cr}^{3+}$  ion works as intensity quencher even for smaller concentration of  $\text{Cr}^{3+}$  ion. This can be well reflected in the lifetime measurements as the lifetime of  ${}^5\text{D}_0$  level of  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition at 614 nm is expected to decrease with the concentration of  $\text{Cr}^{3+}$  ion.

**3.4.5 Lifetime measurements.** The lifetime of  ${}^5\text{D}_0$  level of  $\text{Eu}^{3+}$  ion corresponding to  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition at 614 nm has been measured in absence and presence of  $\text{Cr}^{3+}$  ion. The sample was excited by 393 nm radiation and the fluorescence decay of  $\text{LaVO}_4:1\text{Eu}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples are shown in Fig. 13a and b. We have tried to fit the decay curves using single exponential relation:<sup>32,33</sup>

$$I = I_0 e^{-\frac{t}{\tau}} \quad (\text{v})$$

where,  $I$  is the emission intensity at time  $t$  and  $I_0$  is the initial intensity, and  $\tau$  is the lifetime of  ${}^5\text{D}_0$  level. It was found that the



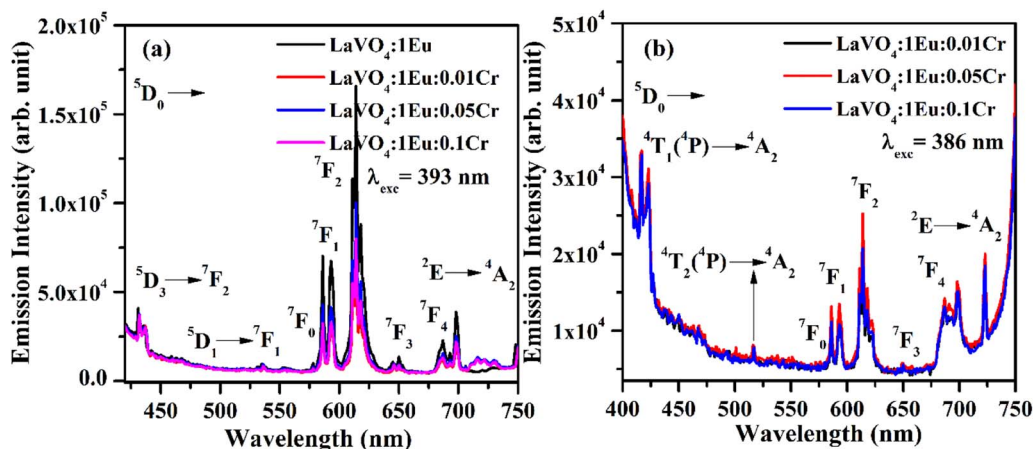


Fig. 12 (a) PL emission spectra of  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples on excitation with 393 nm and (b) PL emission spectra of  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples on excitation with 386 nm.

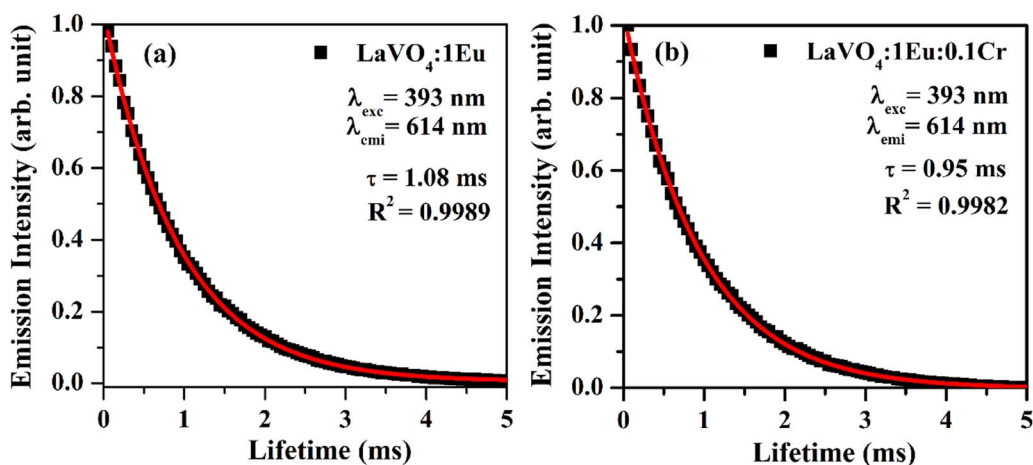


Fig. 13 Fluorescence decay curves of (a)  $\text{LaVO}_4:1\text{Eu}^{3+}$  and (b)  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples for  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition on excitation with 393 nm.

single exponential relation does not reproduce the decay curves correctly and the error limit was found large. We therefore fitted the decay curves using double exponential relation:<sup>34</sup>

$$I = A_1 e^{-t/\tau_1} - A_2 e^{-t/\tau_2} \quad (\text{vi})$$

As is clear from the Fig. 13, the double exponential fitting reproduced the decay curves in a better way. The average lifetime was calculated using the following relation:<sup>34</sup>

$$\tau_{\text{avg}} = \frac{A_1 \tau_1^2 + A_2 \tau_2^2}{A_1 \tau_1 + A_2 \tau_2} \quad (\text{vii})$$

We have measured the lifetime of  ${}^5\text{D}_0$  level of  $\text{Eu}^{3+}$  in the case of  $\text{LaVO}_4:1\text{Eu}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples. It was found that the lifetime of the  ${}^5\text{D}_0$  level decreases with the addition of  $\text{Cr}^{3+}$  ion. This also supports the energy transfer from  $\text{Eu}^{3+}$  to  $\text{Cr}^{3+}$  ions,<sup>48</sup> which quenches the PL intensity of the phosphor.

**3.4.6 Heating effect under 980 nm excitation.** We have also studied the heat generated in the  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples on excitation with different pump power densities of 980 nm diode laser for 5 minutes. The idea is to see the heat generated in the sample because of low phonon frequency of  $\text{LaVO}_4$  host. The samples were prepared in the form of discs of 2 mm diameter and of  $\sim 0.5$  mm thickness. The disc was kept on an aluminum foil. The laser beam was focused on the disc in such a way that its spot size is of  $\sim 1$  mm radius on the disc. The heat generated in the samples was measured at a distance of 2 mm from the edge of the disc. The laser power density was varied from 0–64  $\text{W cm}^{-2}$ . It is well known that the heat generated in the samples will be maximum at the point where the laser beam is falling on the phosphor sample.<sup>34</sup> However, it is difficult to measure it at that point due to sparking. If the heat generated is sufficiently large it can be used for localized thermal massage (physiotherapy) or controlled heating of the substances including water.



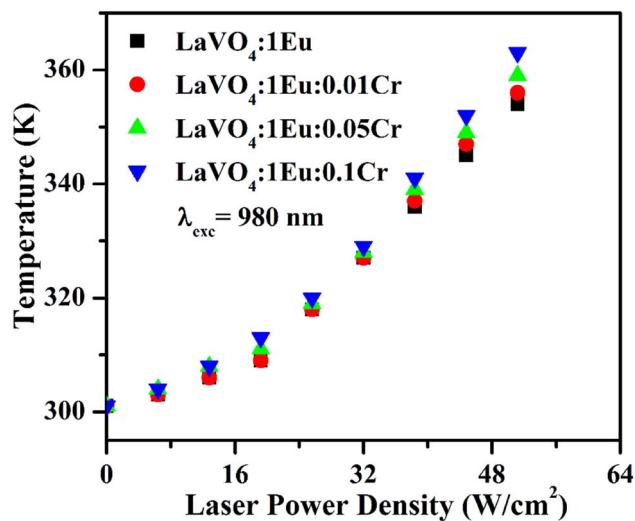


Fig. 14 A plot of generated temperature versus laser power density for the  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples on excitation with 980 nm.

The heat generated in the phosphor samples at different power densities of 980 nm is shown in Fig. 14. In the case of pure  $\text{LaVO}_4$  phosphor, the temperature generated is 336 K at  $64 \text{ W cm}^{-2}$  power density.<sup>34</sup> This value is increased to 354 K at  $51.2 \text{ W cm}^{-2}$  in the case of  $\text{LaVO}_4:1\text{Eu}^{3+}$  phosphor. Above  $51.2 \text{ W cm}^{-2}$ , the phosphor sample starts burning. The heat generated is increased further via  $\text{Cr}^{3+}$  doping and it increases with the increase of concentration of  $\text{Cr}^{3+}$  ion. We have measured the heat generated in the  $\text{LaVO}_4:1\text{Eu}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.01\text{Cr}^{3+}$ ,  $\text{LaVO}_4:1\text{Eu}^{3+}:0.05\text{Cr}^{3+}$  and  $\text{LaVO}_4:1\text{Eu}^{3+}:0.1\text{Cr}^{3+}$  phosphor samples on excitation at different power densities. The heat generated in the samples also depends on the NIR radiation absorbed by the samples. Since the  $\text{Eu}^{3+}$  and  $\text{Cr}^{3+}$  ions absorb the NIR radiation it may also be contributing to the heat generation. The brown color of these phosphor samples is an additional feature for generating large heat. A higher heat is generated in the colored phosphor samples, which is due to excitation of thermal phonons.

## 4. Conclusions

The  $\text{Eu}^{3+}$ ,  $\text{Cr}^{3+}$  doped and co-doped  $\text{LaVO}_4$  phosphors were synthesized by high temperature solid-state reaction method. The powder XRD patterns of phosphors are of crystalline nature. The particles of phosphor samples are spherical with their size in the sub-micron to micron range. The La, Eu, Cr, V and O elements are present in the  $\text{Eu}^{3+}$ ,  $\text{Cr}^{3+}$  co-doped  $\text{LaVO}_4$  phosphor. The FTIR spectra show the absorption bands due to La–O and V–O groups. The optical band gap of phosphor decreases on increasing concentration of  $\text{Cr}^{3+}$  ion. The  $\text{Eu}^{3+}$  doped  $\text{LaVO}_4$  phosphor was excited at 393 and 316 nm wavelengths, which gives intense red color at 614 nm due to  $^5\text{D}_0 \rightarrow ^7\text{F}_2$  transition of  $\text{Eu}^{3+}$  ion. When the  $\text{Cr}^{3+}$  ion is doped in the  $\text{Eu}^{3+}:\text{LaVO}_4$  phosphor; the emission intensity of phosphor is found to decrease due to energy transfer from  $\text{Eu}^{3+}$  to  $\text{Cr}^{3+}$  ions. The lifetime of  $^5\text{D}_0$

level of the  $\text{Eu}^{3+}$  ion also decreases in the  $\text{Eu}^{3+}$ ,  $\text{Cr}^{3+}$  co-doped  $\text{LaVO}_4$  phosphor due to energy transfer. The  $\text{Eu}^{3+}$ ,  $\text{Cr}^{3+}$  doped/co-doped  $\text{LaVO}_4$  phosphors also produces heat under 980 nm excitations. Therefore, the  $\text{Eu}^{3+}$ ,  $\text{Cr}^{3+}$  co-doped  $\text{LaVO}_4$  phosphor may be used for LEDs and heat generating devices.

## Conflicts of interest

There are no conflicts of interest to declare.

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