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Optimizing the luminescence efficiency of an europium (Eu³⁺) doped SrY₂O₄ phosphor for flexible display and lighting applications

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This research paper reports the synthesis and luminescence study of an Eu³⁺ activated SrY₂O₄ phosphor prepared by a modified solid-state reaction method with varying concentrations of Eu³⁺ ions (0.1–2.5 mol%). X-ray diffraction (XRD) revealed the orthorhombic structure and Fourier transform infrared spectroscopy (FTIR) methods were used to analyse the produced phosphors. Photoluminescence emission and excitation spectra were recorded for varying concentrations of Eu³⁺ ions, and an optimum concentration of 2.0 mol% was found to produce the highest intensity. Under 254 nm excitation the emission peaks were found to be at 580 nm, 590 nm, 611 nm and 619 nm, corresponding to transitions at ⁵D₀ → ⁷F₀, ⁵D₀ → ⁷F₁, and ⁵D₀ → ⁷F₂ respectively. Because of Eu³⁺ inherent luminosity, these emission peaks indicate radiative transitions between excited states of ions, making them useful for developing white light-emitting phosphors for optoelectronic and flexible display applications. The 1931 CIE (x, y) chromaticity coordinates were calculated from the photoluminescence emission spectra and found to be near white light emission, indicating the potential application of the prepared phosphor for light emitting diodes (white component). TL glow curve analysis was also performed for various concentrations of doping ions and UV exposure times, and a single broad peak was observed at 187 °C. Using the computerised glow curve deconvolution (CGCD) method, kinetic parameters were computed.

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1. Introduction

The rosy fantasy of a life that can be controlled with the press of a button and more automation would make the daily lives of humans easier. The current trend of innovation and research has made it easier for people to explore new areas of science and technology. Smart materials are one of the aspects of this new frontier. Smart materials are any substances that are capable of detecting or responding to outside stimuli. There are many other types of stimuli that may be used, including environmental, optical, chemical, electrical, thermal, biological, physical, *etc.*^{1–4} Since the reduction in the size of display systems, luminescent materials have attracted a lot of research attention.^{5–7} Applications for smart luminescent materials are

widespread, ranging from polymeric optical fibers and LEDs to displays, self-emitting devices, and light sensors.^{8–12} Achieving durability and adaptability of the electronics and materials used in flexible electronic devices without compromising performance is a critical challenge for smart materials.^{13–18} This obstacle is readily overcome with the use of nanotechnology. The usefulness of several micro- and nanoparticles as smart materials has been reported.^{19–22} Because of this, the intrinsic luminescence feature is the main focus of this study. This property has the potential to be employed in an intelligent (smart) material regarding applications of flexible display.

Europium (Eu³⁺) doped SrY₂O₄ is a well-known luminescence material because it exhibits strong and long-lasting red luminescence under ultraviolet or blue light excitation. The luminescence properties of this material are due to the presence of Eu³⁺ ions, which are known for their unique electronic transitions. There are several reasons why Eu³⁺ doped SrY₂O₄ is a good luminescence material:

(1) Quantum yield: the quantum yield refers to the efficiency of a phosphor in converting absorbed energy into emitted light. Eu³⁺-doped SrY₂O₄ phosphor has a relatively high quantum yield, typically ranging from 60% to 80%. However, it's important to note that the quantum yield can vary depending on the specific manufacturing process and conditions.

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(2) Color purity:

- Eu^{3+} -doped SrY_2O_4 : this phosphor typically exhibits a red emission providing good color purity.
- $\text{Y}_2\text{O}_3:\text{Eu}$: this phosphor is known for its red emission, offering high color purity.
- $\text{Y}_2\text{O}_2\text{S}:\text{Eu}$: this phosphor generally produces a yellowish-red emission, which also has good color purity.

(3) Cost: the cost of phosphor materials can vary based on several factors, including the availability of raw materials and the complexity of the manufacturing process. Eu^{3+} -doped SrY_2O_4 phosphor is relatively more expensive compared to some other commonly used phosphors such as cerium-doped yttrium aluminum garnet (YAG:Ce) or various sulfide-based phosphors. The higher cost can be attributed to the specific composition and synthesis methods required to produce $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphor.

(4) Stability and lifetime: phosphor stability and lifetime are crucial factors to consider in practical applications. Eu^{3+} -doped SrY_2O_4 phosphor exhibits good thermal and chemical stability, allowing it to withstand the harsh conditions typically encountered in solid-state lighting devices. Additionally, it offers a long operational lifetime, ensuring sustained performance over extended periods.

(5) Excitation wavelength: another important aspect is the excitation wavelength required to activate the phosphor material. Eu^{3+} -doped SrY_2O_4 phosphor typically requires ultraviolet (UV) or near-UV excitation in the range from 250 to 350 nm. This wavelength range can be achieved using various light sources, including UV LEDs and mercury lamps.

(6) Luminescent properties:

- Eu^{3+} -doped SrY_2O_4 : this phosphor has a relatively high luminescence efficiency, long decay time, and good thermal stability, making it suitable for various applications such as lighting and displays.
- $\text{Y}_2\text{O}_3:\text{Eu}$: phosphor is known for its high luminescence efficiency and good thermal stability, making it commonly used in red-emitting devices.
- $\text{Y}_2\text{O}_2\text{S}:\text{Eu}$: phosphor also has high luminescence efficiency and good thermal stability. It is often used in yellowish-red-emitting devices.

(7) Application specifics:

- Eu^{3+} -doped SrY_2O_4 : due to its blue-green emission, this phosphor is often used in applications requiring a specific color range, such as white LEDs and plasma displays.
- $\text{Y}_2\text{O}_3:\text{Eu}$: phosphor finds extensive use in red-emitting devices like fluorescent lamps, plasma displays, and cathode ray tubes (CRTs).
- $\text{Y}_2\text{O}_2\text{S}:\text{Eu}$: phosphor is commonly employed in yellowish-red-emitting devices like white LEDs, plasma displays, and CRTs.

Metal oxides with rare earth ion doping have mostly been used for their luminous characteristics. Rare earth ion-doped alkali earth metallic yttrates (MY_2O_4 , $\text{M} = \text{Ba}$, Ca , and Sr) have been seen to exhibit prominent optical emission. According to reports, $\text{SrY}_2\text{O}_4:\text{RE}^{3+}$ ($\text{RE} = \text{Dy}$, Eu , Tb , Gd , Sm , Ce , and Er) may be made using a variety of techniques, including solution combustion, sol-gel, citrate sol-gel, polyol, oil emulsion, aldo-

keto gel, and co-precipitation.²³ The red optical emission of europium makes it one of the intriguing and flexible lanthanides, making it a prime candidate for luminescence applications. $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ (SYO:Eu) has a structure similar to CaFe_2O_4 and exhibits magnificent down-conversion luminescence. At a wavelength of 611 nm, the very powerful emission line produced by the Eu^{3+} long-lived transition $^5\text{D}_0 \rightarrow ^7\text{F}_2$ may be seen. With the excitation of 254 nm (UV region), europium ions are useful for light emitting phosphor and flexible display applications.^{24–36} In this paper, we describe how to easily make new Eu^{3+} doped SrY_2O_4 phosphors, which may be utilized to improve the attractive performance of intelligent (smart) materials for flexible displaying applications. SYO:Eu phosphors emit red light when exposed to UV radiation.

2. Experimental

The preparation of the SrY_2O_4 doped with Eu^{3+} specimen involves a solid-state reaction method. The necessary starting materials, including SrCO_3 , Y_2O_3 , Eu_2O_3 , and H_3BO_3 , Here are the steps for synthesis and characterization of Eu^{3+} doped SrY_2O_4 phosphor:

- Preparation of starting materials: SrCO_3 , Y_2O_3 , and Eu_2O_3 are weighed and mixed in stoichiometric amounts to obtain the desired composition of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$.
- Mixing and grinding: the mixture of powders is then ground together in an agate mortar and pestle for 2 hours to ensure a homogenous mixture.
- Calcination: once the mixing was complete, the resulting powder was placed into a furnace and heated to a temperature of 1000 °C for duration of 1 hour.
- Sintering: after calcination the mixture of powders grind for 1 hour in an agate mortar and pestle and the homogenized powder mixture is then calcined in air at a temperature of 1250 °C for 3 hours to form the desired $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphor. Higher temperature was needed to ensure that the reaction was complete and that the product was fully formed.

The resulting specimen, SrY_2O_4 doped with Eu^{3+} , was then allowed to cool to room temperature. The specimen was characterized using various techniques to confirm the desired properties, such as its crystal structure, morphology, and luminescent properties.

3. Results and discussion

3.1 X-ray analysis

XRD is a widely used technique that involves shining a beam of X-rays onto a sample and measuring the angle and intensity of the diffracted X-rays. The sample's crystal structure and content may be determined from the ensuing XRD pattern. The XRD pattern of Eu^{3+} doped SrY_2O_4 phosphors typically shows characteristic peaks corresponding to the crystalline phases present in the sample. The main phase is SrY_2O_4 , which exhibits orthorhombic crystal structure. The XRD peaks of SrY_2O_4 are sharp and well-defined, indicating a high degree of crystallinity and phase purity. The positions and intensities of the peaks can



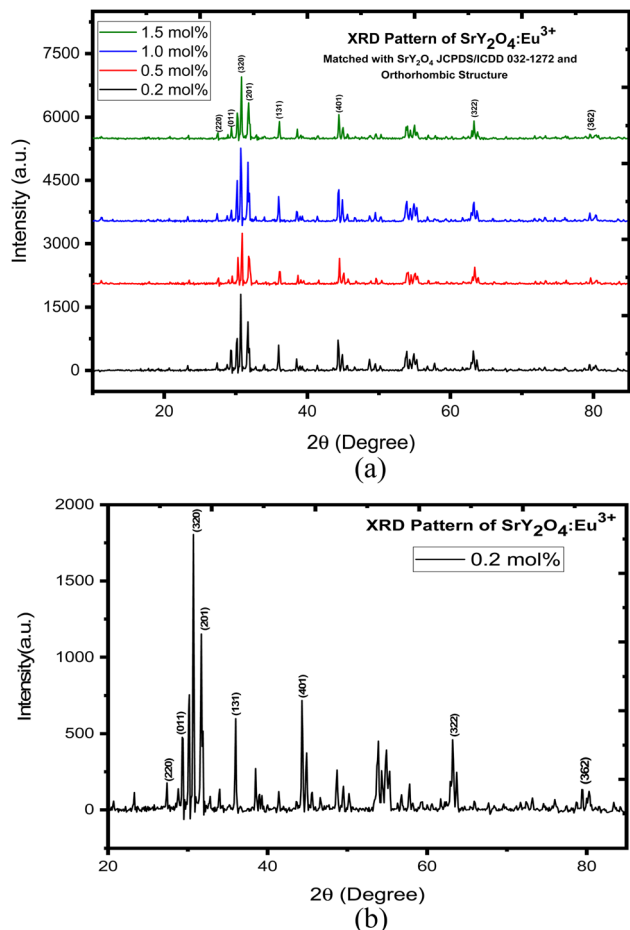


Fig. 1 (a) XRD pattern of SrY₂O₄:Eu³⁺ phosphor. (b) XRD pattern of SrY₂O₄:Eu³⁺ phosphor for 0.2 mol%.

be used to calculate the lattice parameters and crystal structure of the material.

The XRD pattern of the sample is shown in Fig. 1 where aorthorhombic structure is displayed. These values match those of the International Centre for Diffraction Data (JCPDS) card No. 032-1272. The crystallite size of the phosphor was calculated using the Scherer equation. The average crystallite size of the phosphor was found to be 34.068 nm (Table 1).

Table 1 Crystallite size for 0.2 mol% of SrY₂O₄:Eu³⁺ phosphor

<i>hkl</i>	Peak position (2θ)	FWHM	Crystallite size <i>D</i> (nm)	Average size <i>D</i> (nm)
220	23.21705	1.38028	5.876894136	34.06808
*011	27.24806	0.13642	59.93173053	
320	30.67829	0.2346	35.12083109	
201	31.62791	0.2148	38.44694105	
131	35.83333	0.13504	61.83951978	
401	44.26357	0.19296	44.45405375	
322	63.17829	0.52107	17.90162841	
362	79.24419	1.14963	8.973048289	

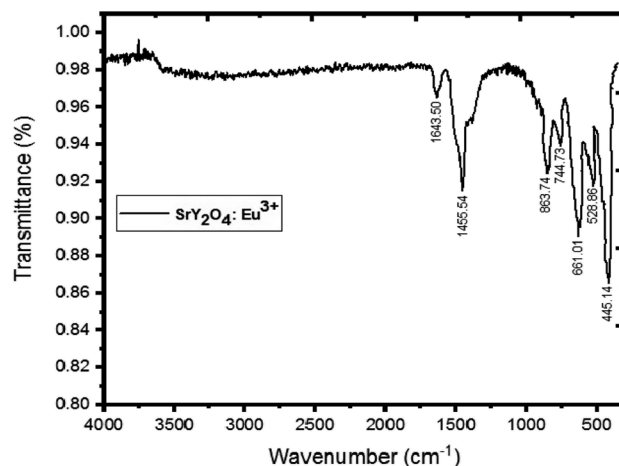


Fig. 2 FTIR spectra of SrY₂O₄:Eu³⁺ phosphor (2.0 mol%).

3.2 FTIR analysis

Fig. 2 displays the SrY₂O₄:Eu³⁺ (2%) phosphor's FTIR spectrum. Strong, sharp peaks may be seen in this spectrum between 445–863 cm⁻¹, which are indicative of Y–O vibrations. An IR peak with a wavelength of 1455–1643 cm⁻¹ is produced when Sr–O is present.³⁷ The production of SrY₂O₄ phosphor is confirmed by all of these observed peaks taken together (Table 2).

3.3 Scanning electron microscopy (SEM)

Scanning Electron Microscopy (SEM) is a powerful characterization tool that is used to examine the morphology and surface features of europium (Eu³⁺) doped SrY₂O₄ phosphors. The

Table 2 Crystallite size for 0.5 mol% of SrY₂O₄:Eu³⁺ phosphor

<i>hkl</i>	Peak position (2θ)	FWHM	Crystallite size <i>D</i> (nm)	Average size <i>D</i> (nm)
220	23.34254	0.9755	8.317365003	33.4503
*011	27.31909	0.13849	59.04481715	
320	30.8561	0.29096	28.32989183	
201	31.86547	0.26985	30.62172567	
131	36.14804	0.15282	54.69350767	
401	44.49721	0.18065	47.52278414	
322	63.39878	0.3074	30.38083641	
362	79.58315	1.1897	8.692154104	

Table 3 Crystallite size for 1.0 mol% of SrY₂O₄:Eu³⁺ phosphor

<i>hkl</i>	Peak position (2θ)	FWHM	Crystallite size <i>D</i> (nm)	Average size <i>D</i> (nm)
220	23.33911	0.6755	12.01116121	34.7837
*011	27.39439	0.1938	42.20033311	
320	30.7273	0.30141	27.33922054	
201	31.70426	0.28279	29.2088247	
131	36.01264	0.17439	47.91012036	
401	44.35684	0.17304	49.58794225	
322	63.23297	0.58242	16.02063895	
362	79.48164	0.19139	53.99151117	

Table 4 Crystallite size for 1.5 mol% of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphor

<i>hkl</i>	Peak position (2θ)	FWHM	Crystallite size <i>D</i> (nm)	Average size <i>D</i> (nm)
220	23.35613	0.38574	21.03434538	32.0282
*011	27.56954	0.1934	42.30342242	
320	30.80483	0.28707	28.71023965	
201	31.80803	0.25283	32.67844571	
131	36.17191	0.17999	46.4405259	
401	44.47332	0.19451	44.13273883	
322	63.33333	0.42113	22.16839805	
362	79.5098	0.551	18.75779935	

technique involves focusing a beam of electrons onto the surface of the sample, causing it to emit secondary electrons that are then collected and analyzed by detectors (Table 4).

When SEM is used to study Eu^{3+} doped SrY_2O_4 phosphors, it provides high-resolution images of the particle size, shape, and distribution (Fig. 3). The technique can also be used to observe any morphological changes that occur as a result of the synthesis method or thermal treatment. SEM imaging can reveal the surface features of the phosphor particles, such as cracks, pores, or other defects, which can affect the luminescence properties of the material (Table 5).

3.4 Energy dispersive X-ray spectroscopy (EDX)

Energy Dispersive X-ray Spectroscopy (EDX) analysis is a valuable tool for studying the elemental composition of Eu^{3+} doped SrY_2O_4 phosphors and can provide important insights into the materials' luminescence properties. By optimizing the elemental composition and concentration of the phosphor, using this technique we can enhance the luminescence

Table 5 Average crystallite size for concentration (0.2–1.5) mol% of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphor

S. no	Concentration (mol%)	Average crystallite size <i>D</i> (nm)
1	0.2	34.068
2	0.5	33.4503
3	1	34.7837
4	1.5	32.0282

efficiency and stability of the material, which can have important implications for applications such as lighting and sensing.

It is an analysis of the components of produced phosphor. Fig. 4a displays quantitative representations of the sample's Y, Sr, O, and Eu elements. The data supports the conclusion that $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphor can be synthesized. The qualitative analysis of the process of creating the components is shown in Fig. 4b.

3.5 Photoluminescence (PL) observation

Doping with Eu^{3+} ions improved the emission spectra of the host SrY_2O_4 phosphor, which has no luminescence.²³ The SrY_2O_4 phosphor doped with Eu^{3+} PL excitation spectra are shown in Fig. 5. Excitation spectra were captured at an emission wavelength of 613 nm. It has a wide spectrum between 190 and 254 nanometers. Eu^{3+} doped SrY_2O_4 phosphor's PL emission spectrum, measured at 254 nm, shows the typical emissions at 580, 590, 611, and 619 nm. Fig. 6. The emission spectra of Eu^{3+} are produced by transitions between $^5\text{D}_0 \rightarrow ^7\text{F}_j$ ($j = 0, 1, 2$).³⁸ The brightest emission band can be seen at 611 nm, which is associated with the electric dipole transition $^5\text{D}_0 \rightarrow ^7\text{F}_2$ (ref. 23–25). The $^5\text{D}_0 \rightarrow ^7\text{F}_0$ and $^5\text{D}_0 \rightarrow ^7\text{F}_1$ magnetic dipole

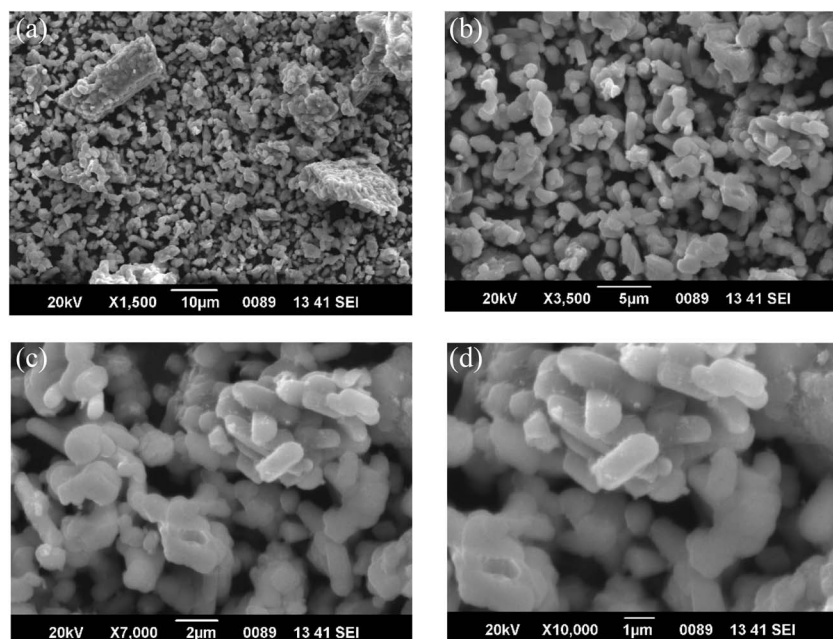
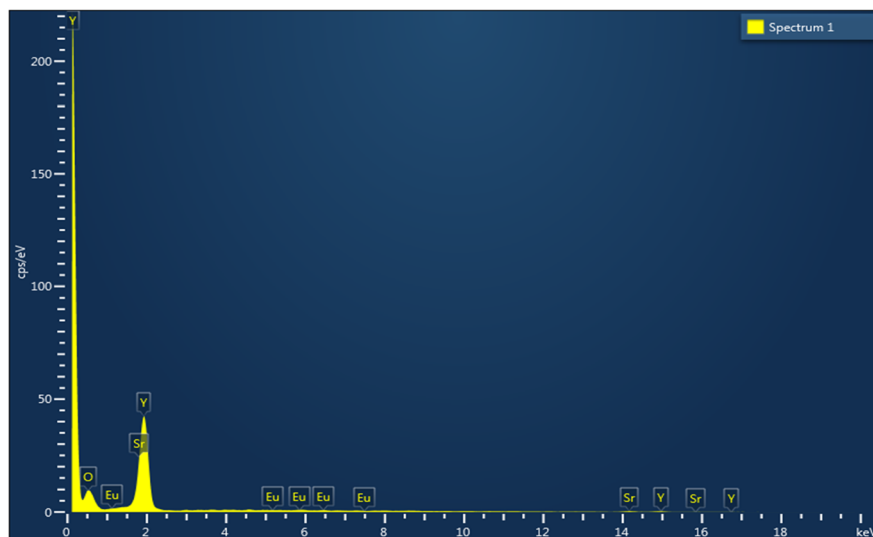
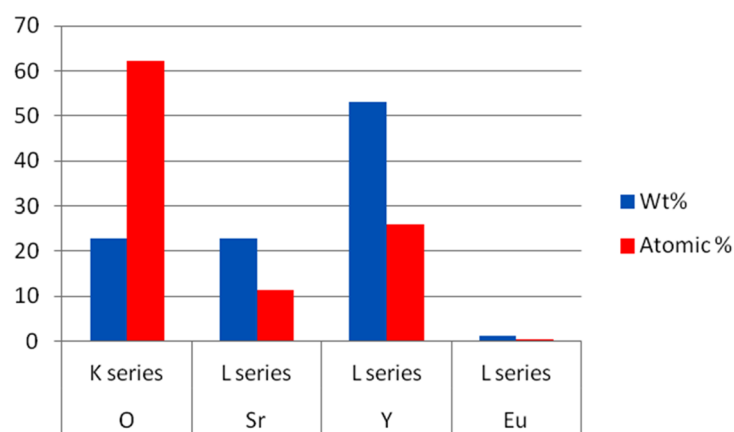


Fig. 3 SEM images of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ (2.0 mol%) (a) $\times 1.5\text{k}$ 10 μm (b) $\times 3.5\text{k}$ 5 μm (c) $\times 7\text{k}$ 2 μm (d) $\times 10\text{k}$ 1 μm .





(a)



(b)

Fig. 4 (a) EDX pattern of prepared phosphor Eu^{3+} (2.0 mol%) doped SrY_2O_4 . (b) Quantitative EDX of Eu^{3+} doped SrY_2O_4 phosphor (2.0 mol%).

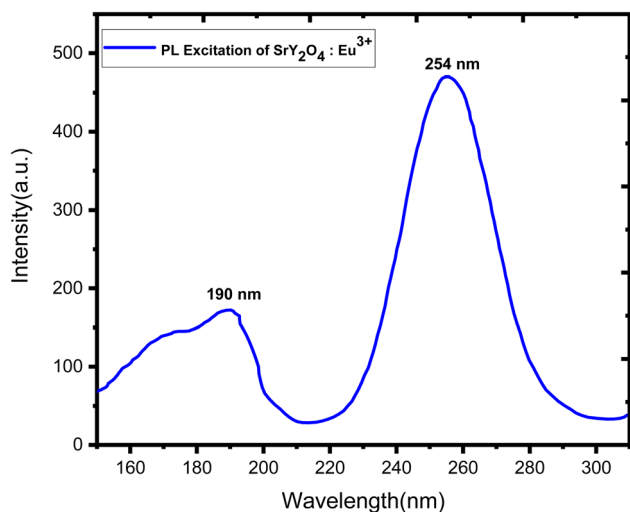


Fig. 5 PL excitation spectra of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ (2.0 mol%) phosphor.

transitions⁶⁰ give rise to the 580 nm and 590 nm bands, respectively. Due to similar ionic radii, the emergence of the $^5\text{D}_0 \rightarrow ^7\text{F}_1$ transition supports the substitution of Eu^{3+} ions at Y^{3+} sites.³⁹ The minor separation of the $^5\text{D}_0 \rightarrow ^7\text{F}_1$ and $^5\text{D}_0 \rightarrow ^7\text{F}_2$ transition lines in the emission spectra of the $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphor amply demonstrates the considerable influence of the host composition, crystal structure, and coordination environment on luminescence qualities.⁴⁰ Fig. 6 shows how the concentration of the dopant affects the intensity of the emission. The intensity continues to rise as the Eu^{3+} ion concentration rises from 0.1 to 2.0 mol%. However, the concentration quenching event causes the intensity to dramatically falldown for 2.5 mol% of the time. Intensity is reduced by quenching, which happens when Eu^{3+} ions move in closer proximity to one another and interact with one another to transfer charge. Eu^{3+} doped SrY_2O_4 phosphors using 2.0 mol% had the highest

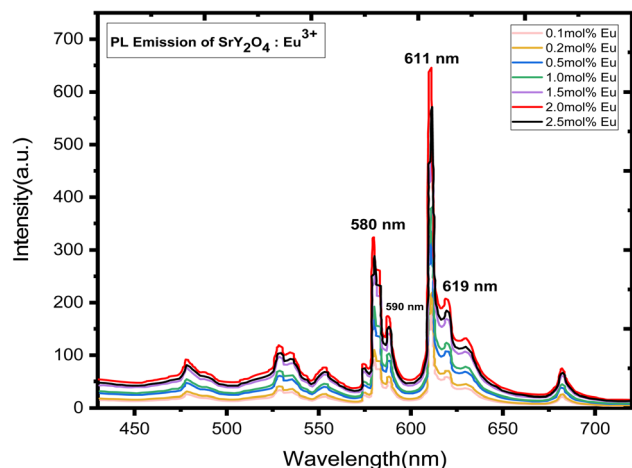


Fig. 6 PL emission spectra of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ (0.1–2.5 mol%) phosphor.

emission intensity, making it suitable for various light-emitting applications (Table 6).

The fluorescence quantum yield (ϕ_F) is a measure of the efficiency of fluorescence, representing the ratio of absorbed photons to emitted photons through fluorescence. It quantifies the likelihood of the excited state being deactivated through fluorescence rather than non-radiative mechanisms. The

comparative method developed by Williams *et al.* is considered the most reliable approach for determining (ϕ_F). This method involves utilizing well-characterized standard samples with known (ϕ_F) values, enabling accurate measurement and comparison of fluorescence efficiencies.⁶²

$$\phi_F = \eta \frac{\text{Number of emitted photons}}{\text{Number of absorbed photons}}$$

where the η denote the refractive index of the sample.

The fluorescence quantum yield (ϕ_F) is a parameter that measures the efficiency of fluorescence by comparing the number of absorbed photons to the number of emitted photons. To determine ϕ_F , one common approach involves analyzing the integrated areas of the absorption and emission spectra. The integrated area represents the number of absorbed photons and the number of emitted photons, respectively. By comparing these values, the fluorescence quantum yield can be accurately calculated (Table 7).

3.6 Commission international de l'Eclairage (CIE) coordinates

The Commission International de l'Eclairage (CIE) coordinates are a standard way to describe the colour of light emitted by these phosphors. The CIE colour space is a three-dimensional space that describes all possible colours based on three

Table 6 Quantum yield values for different concentrations (mol%) of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$

S. no	Concentration (mol%)	Absorption of photons	Emission of photons	Quantum yield
1	0.1	22 988.91862	5870.677682	0.242601398
2	0.2		7114.49341	0.29400116
3	0.5		10 769.65826	0.445048135
4	1		12 347.99237	0.51027162
5	1.5		16 981.28813	0.701739129
6	2		23 090.70313	0.954206169
7	2.5		18 523.81395	0.765482863

Table 7 Table of standard materials and their literature quantum yield values

Compound	Solvent	Literature quantum yield	Emission range/nm	Reference ^{63–74}
Cresyl violet	Methanol	0.54	600–650	<i>J. Phys. Chem.</i> , 1979, 83 , 696
Rhodamine 101	Ethanol + 0.01% HCl	1.00	600–650	<i>J. Phys. Chem.</i> , 1980, 84 , 1871
Quinine sulfate	0.1 M H_2SO_4	0.54	400–600	<i>J. Phys. Chem.</i> , 1961, 65 , 229
Fluorescein	0.1 M NaOH	0.79	500–600	<i>J. Am. Chem. Soc.</i> , 1945, 1099
Norharmane	0.1 M H_2SO_4	0.58	400–550	<i>J. Lumin.</i> , 1992, 51 , 269–74
Harmane	0.1 M H_2SO_4	0.83	400–550	<i>J. Lumin.</i> , 1992, 51 , 269–74
Harmine	0.1 M H_2SO_4	0.45	400–550	<i>J. Lumin.</i> , 1992, 51 , 269–74
2-Methylharmane	0.1 M H_2SO_4	0.45	400–550	<i>J. Lumin.</i> , 1992, 51 , 269–74
Chlorophyll A	Ether	0.32	600–750	<i>Trans. Faraday Soc.</i> , 1957, 53 , 646–55
Zinc phthalocyanine	1% pyridine in toluene	0.30	660–750	<i>J. Chem. Phys.</i> , 1971, 55 , 4131
Benzene	Cyclohexane	0.05	270–300	<i>J. Phys. Chem.</i> , 1968, 72 , 325
Tryptophan	Water, pH 7.2, 25C	0.14	300–380	<i>J. Phys. Chem.</i> , 1970, 74 , 4480
2-Aminopyridine	0.1 M H_2SO_4	0.60	315–480	<i>J. Phys. Chem.</i> , 1968, 72 , 2680
Anthracene	Ethanol	0.27	360–480	<i>J. Phys. Chem.</i> , 1961, 65 , 229
9,10-Diphenyl anthracene	Cyclohexane	0.90	400–500	<i>J. Phys. Chem.</i> , 1983, 87 , 83



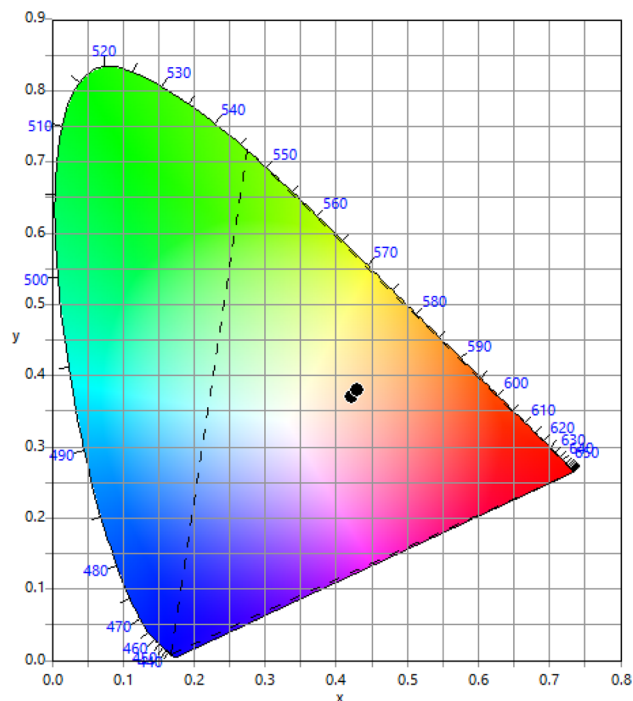


Fig. 7 CIE coordinate for $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphor (0.1–2.5 mol%).

parameters: X, Y, and Z. The X, Y, and Z coordinates represent the relative amounts of red, green, and blue light needed to create a specific colour. The CIE colour space is a useful tool for describing the colour of light emitted by luminescent materials like Eu^{3+} doped SrY_2O_4 phosphors. The CIE coordinates of Eu^{3+} doped SrY_2O_4 phosphors can be determined using photoluminescence spectroscopy. The emitted light is collected and passed through a monochromator to separate the light into its component wavelengths. The intensity of the light at each wavelength is measured, and the CIE coordinates are calculated based on the spectral distribution of the emitted light. Eu^{3+} doped SrY_2O_4 phosphors typically exhibit a near white region luminescence (Fig. 7), with CIE coordinates of according to Table 8. The exact CIE coordinates can vary depending on the specific composition and synthesis conditions of the phosphor.

3.7 TL glow curve analysis

Europium (Eu^{3+}) doped SrY_2O_4 phosphors exhibit thermoluminescence (TL) properties, which make them useful for

Table 8 CIE coordinates of Eu^{3+} doped SrY_2O_4 phosphor with different concentration

Concentration	x	y	u'	v'	CCT	CRI
0.1	0.4301	0.3801	0.2568	0.5105	2912	85
0.2	0.43	0.3789	0.2572	0.51	2904	85
0.5	0.4297	0.3797	0.2567	0.5103	2916	85
1	0.4306	0.3802	0.257	0.5106	2906	85
1.5	0.4294	0.3793	0.2566	0.51	2919	85
2	0.4226	0.3692	0.2567	0.5046	2951	85
2.5	0.4295	0.3791	0.2568	0.51	2915	85

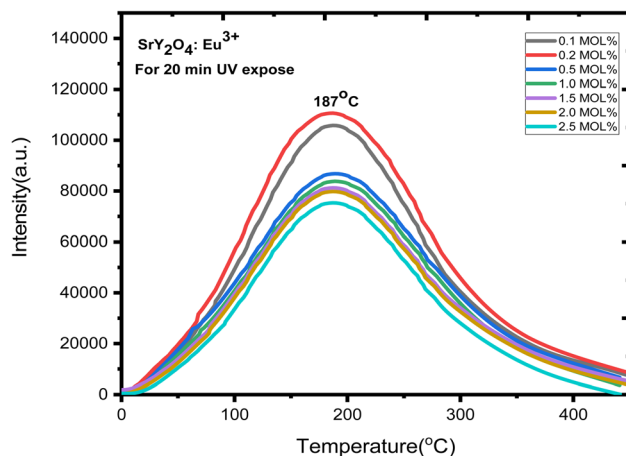


Fig. 8 TL glow curve analysis of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$.

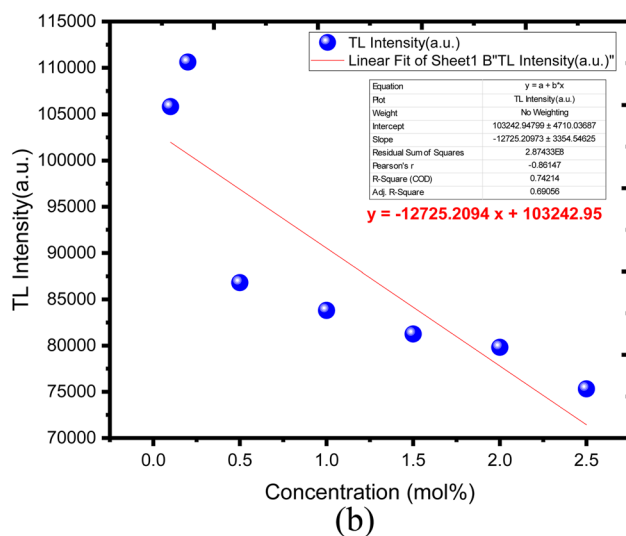
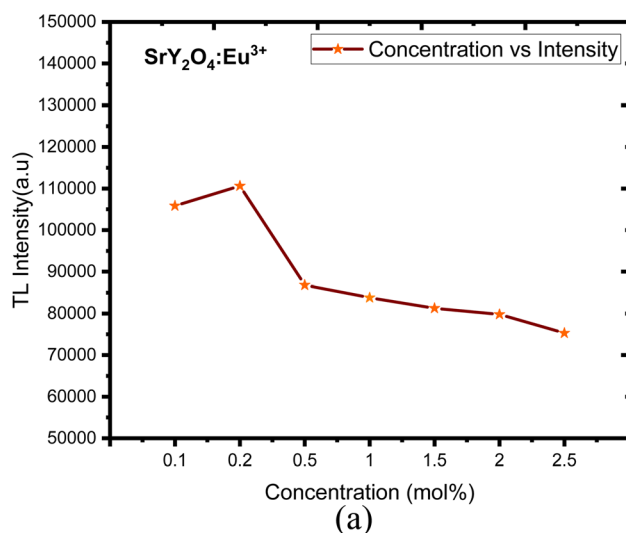


Fig. 9 (a) $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ concentration vs. intensity. (b) Linear fit with a mean error bar of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ for (0.1–2.5) mol%.

radiation dosimetry, medical radiation dosimetry, environmental monitoring, and space radiation dosimetry applications. TL refers to the phenomenon where a material emits light when heated after being exposed to ionizing radiation. The intensity and shape of the TL glow curve, which is the plot of the emitted light intensity as a function of temperature, can provide information about the energy and type of radiation absorbed by the material.

Europium-doped SrY_2O_4 phosphor has a linear response to a 20 minutes UV exposure, as seen by the TL glow curve (Fig. 8). Thermoluminescence (TL) intensity rises with increasing europium concentration (Fig. 8 inset), and its wide peak centers about 187°C (where the trapped electrons are released from the lattice defects), making it an excellent peak for a thermoluminescence dosimeter. Up to 0.2 mol%, the TL intensity rises; beyond that, it falls as a result of the concentration quenching phenomena (Fig. 9a). The TL glow curve for an optimum concentration of 0.2 mol% is shown in Fig. 10, which displays a linear wide peak with varying UV exposure durations at a fixed rate of heating of 2.5°C per s . The concentration quenching effect causes TL intensity to drop after the 25 minutes UV treatment for 0.2 mol% (Fig. 11a). Fitting the curve for the optimal concentration of 0.2 mol% using the CGCD method yields an excellent theoretical and experimental fit.

Fig. 9b showed the linear fit corresponding to its mean error with respect to concentration response. The obtained TL phosphor exhibits a linear fit for different concentration over from 0.1 to 2.5 mol%. TL intensity was decreased linearly when the dose was increased, obtaining a determination coefficient $R^2 = 0.74214$.⁷⁵

Fig. 11b showed the linear fit corresponding to its mean error with respect to dose response. The obtained TL phosphor exhibits a linear fit for different concentration over from 5 to 30 min UV dose exposed. TL intensity was increased linearly when the dose was increased, obtaining a determination coefficient $R^2 = 0.59899$.⁷⁵

3.7.1 CGCD analysis. TLD phosphor often shows many peaks upon charge carrier (hole or electron) emission. The

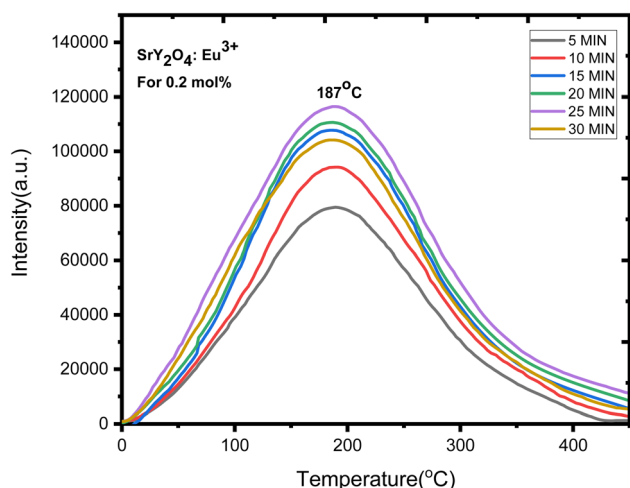


Fig. 10 TL glow curve analysis of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ for 0.2 mol%.

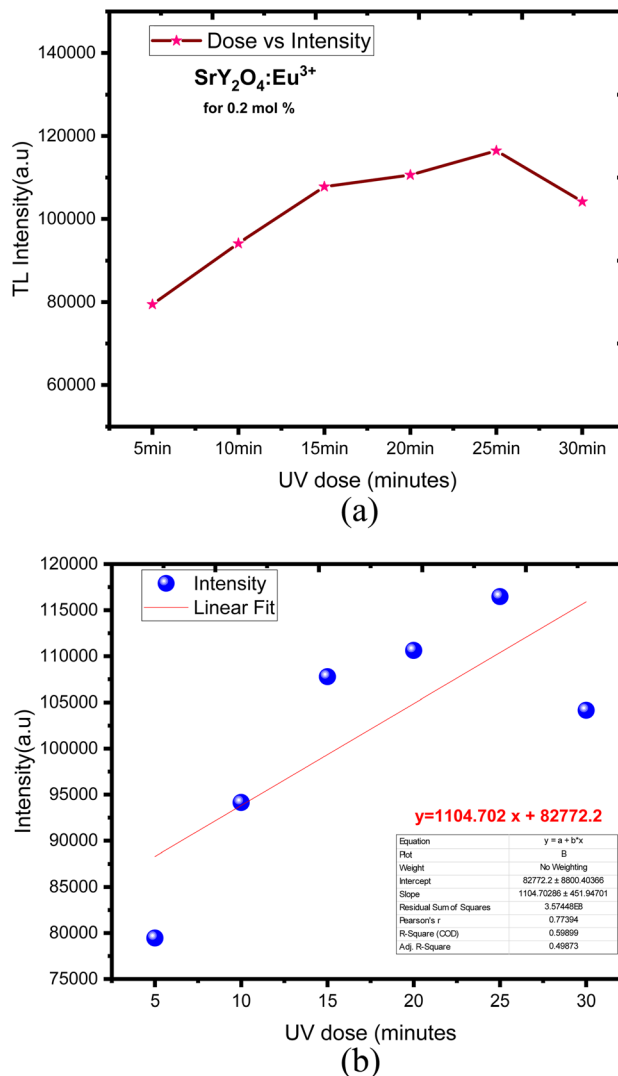


Fig. 11 (a) $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ dose vs. intensity. (b) Linear fit with a mean error bar of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ for (5–30 min) UV dose.

kinetic factors/parameters (E , b , and s) affect the dosimetric features of TL materials a lot. These parameters will provide significant information regarding the mechanism of phosphor emission. Understanding the kinetic parameters of a TLD phosphor is crucial for producing effective TLDs. An empirical set of equations^{41–51} developed by Chen's allows the estimation of these parameters. Using the Glow fit software, the glow curve is deconvoluted (Fig. 12) in order to implement the peak shape approach. In order to calculate the kinetic parameters of the TL materials' glow peaks, we employed the peak shape approach, also known as Chen's peak method.

Evidently, there are eight wide peaks in the CGCD pattern (Fig. 12), and kinetic parameters are computed for each of those eight peaks in Table 3. After 20 minutes of UV irradiation at varying concentrations of europium ions, the recorded glow curves of the manufactured phosphors reveal a broad peak, suggesting that they are composite in nature. Therefore, the kinetic parameters were deconvoluted and calculated using the

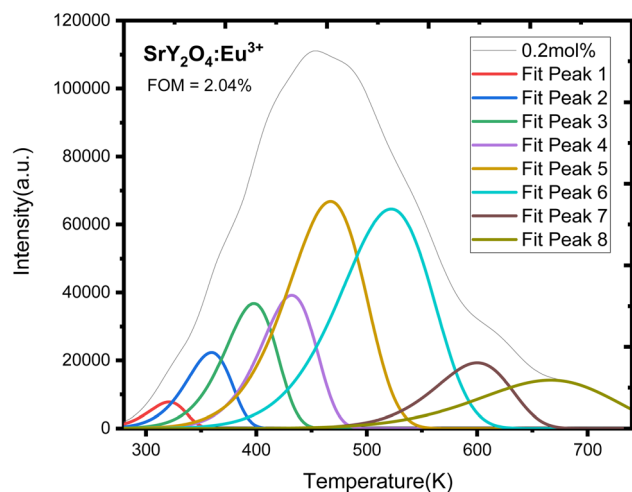


Fig. 12 CGCD pattern of UV induced $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ doped phosphor for optimized UV dose and 0.2 mol% concentration.

CGCD technique. The glow curves of the greatest intensity peak, 0.2 mol% europium ion irradiated with UV for 20 min, were deconvoluted using the CGCD technique using the Glow fit software. The Halperin and Barner formulas provide the mathematical foundation for this computer software. These equations explain the movement of charges between the different energy levels that occur during the process of trap emptying caused by thermal heating. This software calculated the trap level kinetic parameters for each deconvoluted peak. The experimental glow curves were used to fit the theoretically produced glow curves, and the accuracy of the fit was assessed by computing the figure of merit (FOM) associated with each fitting. When FOM values were less than 5%, it was determined that the fits were satisfactory. Current research puts FOM at 2.04%, confirming excellent agreement between theoretically derived and actually observed glow curves.^{52,53} Fig. 12 displays the fitted TL glow curves, and Table 3 provides a summary of the CGCD-method-calculated values for the frequency factors (s) and trap depths (E) of captured charges.

The location of trapping levels inside the forbidden gap is referred to as trap depth or activation energy (E), and it plays a significant role in the loss of dosimetry information that had been preserved in the materials after irradiation. The order of kinetics (b) is the process by which detrapped charge carriers

recombine with their equivalents. When the trap is modeled as a potential well, the number of times an electron collides with the wall multiplied by the wall reflection coefficient gives us the frequency factor (s). As a result, the trapping parameters of a thermoluminescent material are the foundation for an accurate dosimetry investigation.^{54–61}

Here, the information about the trap level obtained from the kinetic parameters (Table 9) shows that almost one electron is trapped there because of the value of (shape factor) 0.435 and it shows the general order kinetics (b) and the needed energy to escape one electron from the trap level is high (0.50–0.86 eV), *i.e.* activation energy (E), which shows the formation of traps is high and stable. The rate of escaping electrons per second, which falls between 2.12×10^9 and $3.66 \times 10^9 \text{ s}^{-1}$, is likewise fairly high.

3.8 Conclusion

The present study provides valuable insights into the properties of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphors, as summarized in the following conclusions:

- (1) XRD analysis confirmed the existence of a single orthorhombic crystalline phase in the newly synthesized $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphor, with nanocrystalline behavior.
- (2) The $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphors exhibited a fine surface morphology with a spherical shape which is confirmed by SEM imaging.
- (3) FTIR spectra revealed specific vibration modes of Y–O, Sr–O, and the production of SrY_2O_4 phosphor is confirmed by all of these observed peaks taken together.
- (4) Photoluminescence excitation spectra showed a broad excitation centered at 254 nm, with well-resolved emission peaks 580 nm, 590 nm, 611 nm and 619 nm corresponding to transitions at $^5\text{D}_0 \rightarrow ^7\text{F}_0$, $^5\text{D}_0 \rightarrow ^7\text{F}_1$, and $^5\text{D}_0 \rightarrow ^7\text{F}_2$, respectively.
- (5) The emission color of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphors was in the near white light region, as shown in the CIE 1931 graph, and exhibited good color tenability. These can act as a white light-emitting phosphor for optoelectronic and flexible display applications.
- (6) The TL glow curve demonstrated that the thermoluminescence intensity of $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ phosphors increased with increasing UV exposure time and peaked at 187 °C, indicating a good thermoluminescence dosimeter peak.
- (7) The CGCD methodology provided a successful theoretical and experimental fit, with kinetic parameters computed for

Table 9 Calculation of kinetic parameters using CGCD programme for UV induced $\text{SrY}_2\text{O}_4:\text{Eu}^{3+}$ doped phosphor

T_1 (K)	T_m (K)	T_2 (K)	τ	δ	ω	$\mu = \delta/\omega$	Activation energy (eV)	Frequency factor s^{-1}
298	321	338	23	17	40	0.425	0.5	2.12×10^9
332	359	380	27	21	48	0.4375	0.5175	2.26×10^9
366	398	422	32	24	56	0.4285714	0.5458	2.33×10^9
398	432	458	34	26	60	0.4333333	0.6118	2.65×10^9
420	467	504	47	37	84	0.4404762	0.5011	2.20×10^9
466	522	566	56	44	100	0.44	0.523	2.30×10^9
552	600	636	48	36	84	0.4285714	0.8559	3.66×10^9
587	666	731	79	65	144	0.4513889	0.5823	2.62×10^9



SrY₂O₄:Eu³⁺ phosphor and it shows general order kinetics (b) (Table 9).

Eu³⁺ doped SrY₂O₄ is a good luminescence material due to its high quantum efficiency, narrow emission bands, long luminescence lifetime, and high stability, making it a versatile material for various applications in optoelectronics, such as solid-state lighting and flat panel displays.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 D. C. Celikel, Smart E-Textile Materials, in *Advanced functional materials*, ed. N. Tasaltin, P. S. Nnamchi and S. Saud, IntechOpen, London, 2020, pp. 1–16, DOI: [10.5772/intechopen.92439](#).
- 2 L. Liu, W. Xu, Y. Ding, S. Agarwal, A. Greiner and G. Duan, A review of smart electrospun fibers toward textiles, *Comput. Commun.*, 2020, **22**, 100506, DOI: [10.1016/j.coco.2020.100506](#).
- 3 M. Ghahremani Honarvar and M. Latifi, Overview of wearable electronics and smart textiles, *J. Text. Inst.*, 2017, **108**(4), 631–652, DOI: [10.1080/00405000.2016.1177870](#).
- 4 M. Chen, Y. Ma, J. Song, C. F. Lai and B. Hu, Smart clothing: connecting human with clouds and big data for sustainable health monitoring, *Mob. Netw. Appl.*, 2016, **21**(5), 825–845, DOI: [10.1007/s11036-016-0745-1](#).
- 5 A. Bratovic, Different applications of nanomaterials and their impact on the environment, *Int. J. Mater. Sci. Eng.*, 2019, **5**(1), 1–7, DOI: [10.14445/23948884/IJMSE-V5I1P101](#).
- 6 P. Kumar, S. Singh and B. K. Gupta, Future prospects of luminescent nanomaterial based security inks: from synthesis to anti-counterfeiting applications, *Nanoscale*, 2016, **8**(30), 14297–14340, DOI: [10.1039/C5NR06965C](#).
- 7 J. Yu, M. Luo, Z. Lv, S. Huang, H. H. Hsu, C. C. Kuo and Y. Zhou, Recent advances in optical and optoelectronic data storage based on luminescent nanomaterials, *Nanoscale*, 2020, **12**(46), 23391–23423, DOI: [10.1039/D0NR06719A](#).
- 8 S. Kwon, Y. H. Hwang, M. Nam, H. Chae, H. S. Lee, Y. Jeon and K. C. Choi, Recent progress of fiber shaped lighting devices for smart display applications—a fibertronic perspective, *Adv. Mater.*, 2020, **32**(5), 1903488, DOI: [10.1002/adma.201903488](#).
- 9 M. Cinquino, C. T. Prontera, M. Pugliese, R. Giannuzzi, D. Taurino, G. Gigli and V. Maiorano, Light-emitting textiles: device architectures, working principles, and applications, *Micromachines*, 2021, **12**(6), 652, DOI: [10.3390/mi12060652](#).
- 10 X. Shi, Y. Zuo, P. Zhai, J. Shen, Y. Yang, Z. Gao and H. Peng, Large-area display textiles integrated with functional systems, *Nature*, 2021, **591**(7849), 240–245, DOI: [10.1038/s41586-021-03295-8](#).
- 11 M. Haque, Nano fabrics in the 21st century: a review, *Asian J. Nanosci. Mater.*, 2019, **2**(2), 131–148, DOI: [10.26655/ajnanomat.2019.3.2](#).
- 12 Z. Gong, Z. Xiang, X. OuYang, J. Zhang, N. Lau, J. Zhou and C. C. Chan, Wearable fiber optic technology based on smart textile: a review, *Materials*, 2019, **12**(20), 3311, DOI: [10.3390/ma12203311](#).
- 13 C. W. Kan and Y. L. Lam, Future trend in wearable electronics in the textile industry, *Appl. Sci.*, 2021, **11**(9), 3914, DOI: [10.3390/app11093914](#).
- 14 A. Angelucci, M. Cavicchioli, I. A. Cintorino, G. Lauricella, C. Rossi, S. Strati and A. Aliverti, Smart textiles and sensorized garments for physiological monitoring: a review of available solutions and techniques, *Sensors*, 2021, **21**(3), 814, DOI: [10.3390/s21030814](#).
- 15 X. Hu, Y. Dou, J. Li and Z. Liu, Buckled structures: fabrication and applications in wearable electronics, *Small*, 2019, **15**(32), 1804805, DOI: [10.1002/sml.201804805](#).
- 16 Q. Zhang, Y. L. Wang, Y. Xia, P. F. Zhang, T. V. Kirk, X. D. Chen and X. D., Textile-only capacitive sensors for facile fabric integration without compromise of wearability, *Adv. Mater. Technol.*, 2019, **4**(10), 1900485, DOI: [10.1002/admt.201900485](#).
- 17 A. A. Simegnaw, B. Malengier, G. Rotich, M. G. Tadesse and L. Van Langenhove, Review on the integration of microelectronics for E-Textile, *Materials*, 2021, **14**(17), 5113, DOI: [10.3390/ma14175113](#).
- 18 E. A. Hassan, M. A. Ahmed, S. Zhu and O. K. Alebeid, Fibre-based wearable electronic technology for personal protective clothing, in *Protective textiles from natural resources*, Woodhead Publishing, 2022, pp. 511–547, DOI: [10.1016/B978-0-323-90477-3.00019-5](#).
- 19 A. A. Basheer, Advances in the smart materials applications in the aerospace industries, *Aircr. Eng. Aerosp. Technol.*, 2020, **92**(7), 1027–1035, DOI: [10.1108/AEAT-02-2020-0040](#).
- 20 A. Raza, U. Hayat, T. Rasheed, M. Bilal and H. M. Iqbal, “Smart” materials-based near-infrared light-responsive drug delivery systems for cancer treatment: a review, *J. Mater. Sci. Technol.*, 2019, **8**(1), 1497–1509, DOI: [10.1016/j.jmrt.2018.03.007](#).
- 21 C. Mendes-Felipe, J. Oliveira, I. Etxebarria, J. L. Vilas-Vilela and S. Lanceros-Mendez, State-of-the-art and future challenges of UV curable polymer-based smart materials for printing technologies, *Adv. Mater. Technol.*, 2019, **4**(3), 1800618, DOI: [10.1002/admt.201800618](#).
- 22 Z. Li, F. Yang and Y. Yin, Smart materials by nanoscale magnetic assembly, *Adv. Funct. Mater.*, 2020, **30**(2), 1903467, DOI: [10.1002/adfm.201903467](#).
- 23 R. Priya, S. Kaur, U. Sharma, O. P. Pandey and S. J. Dhoble, A review on recent progress in rare earth and transition metals activated SrY₂O₄ phosphors, *J. Mater. Sci.: Mater. Electron.*,



- 2020, **31**(16), 13011–13027, DOI: [10.1007/s10854-020-03930-6](https://doi.org/10.1007/s10854-020-03930-6).
- 24 S. P. Ghorpade, R. H. Krishna, R. M. Melavanki, V. Dubey and N. R. Patil, Effect of Eu^{3+} on optical and energy bandgap of SrY_2O_4 nanophosphors for FED applications, *Optik*, 2020, **208**, 164533, DOI: [10.1016/j.ijleo.2020.164533](https://doi.org/10.1016/j.ijleo.2020.164533).
- 25 D. R. Taikar, Synthesis and luminescence property of SrY_2O_4 : M ($\text{M} = \text{Eu}^{3+}$, Tb^{3+} , Sm^{3+} , Tb^{3+} , Sm^{3+} , Ce^{3+} , Bi^{3+}) phosphors, *J. Lumin.*, 2018, **204**, 24–29, DOI: [10.1016/j.jlumin.2018.07.040](https://doi.org/10.1016/j.jlumin.2018.07.040).
- 26 E. Pavitra, G. S. R. Raju, W. Park and J. S. Yu, Concentration and penetration depth dependent tunable emissions from Eu^{3+} co-doped SrY_2O_4 : Dy^{3+} nanocrystalline phosphor, *New J. Chem.*, 2014, **38**(1), 163–169, DOI: [10.1039/C3NJ00987D](https://doi.org/10.1039/C3NJ00987D).
- 27 R. K. Tamrakar and K. Upadhyay, Combustion synthesis and luminescence behaviour of the Tb^{3+} doped SrY_2O_4 phosphor, *J. Electron. Mater.*, 2018, **47**(1), 651–654, DOI: [10.1007/s11664-017-5825-x](https://doi.org/10.1007/s11664-017-5825-x).
- 28 S. P. Ghorpade, N. Kottam, R. Melavanki and N. R. Patil, Photoluminescence, TGA/DSC and photocatalytic activity studies of Dy^{3+} doped SrY_2O_4 nanophosphors, *RSC Adv.*, 2020, **10**(36), 21049–21056, DOI: [10.1039/D0RA03094E](https://doi.org/10.1039/D0RA03094E).
- 29 V. Singh, K. Swapna, S. Kaur, A. S. Rao and J. L. Rao, Narrow-band UVB-emitting Gddoped SrY_2O_4 phosphors, *J. Electron. Mater.*, 2020, **49**(5), 3025–3030, DOI: [10.1007/s11664-020-08032-x](https://doi.org/10.1007/s11664-020-08032-x).
- 30 R. Wei, Z. Zheng, Y. Shi, X. Peng, H. Wang, X. Tian and H. Guo, Tunable emission with excellent thermal stability in single-phased SrY_2O_4 : Bi^{3+} , Eu^{3+} phosphors for UV-LEDs, *J. Alloys Compd.*, 2018, **767**, 403–408, DOI: [10.1016/j.jallcom.2018.07.101](https://doi.org/10.1016/j.jallcom.2018.07.101).
- 31 S. Bhatia, R. Priya, D. Kumar, S. Upadhyay and O. P. Pandey, Structural and luminescent studies of titanium co-doped SrY_2O_4 : Eu phosphors, *Luminescence*, 2022, **37**(9), 1585–1596, DOI: [10.1002/bio.4335](https://doi.org/10.1002/bio.4335).
- 32 I. Suryawanshi, R. Kalia, R. K. Kunchala, S. L. Mudavath and B. S. Naidu, Detection of latent fingerprints using luminescent $\text{Gd}_{0.95}\text{Eu}_{0.05}\text{PO}_4$ nanorods, *J. Rare Earths*, 2022, **40**(4), 572–578, DOI: [10.1016/j.jre.2021.01.015](https://doi.org/10.1016/j.jre.2021.01.015).
- 33 S. P. Singh, S. Srinidhi, R. K. Kunchala, R. Kalia, S. N. Achary and B. S. Naidu, Structural and excitation-dependent photoluminescence properties of $\text{Bi}_{0.95x}\text{Gd}_x\text{Eu}_{0.05}\text{PO}_4$ ($0 \leq x \leq 0.95$) solid solutions and their anticounterfeiting applications, *Cryst. Growth Des.*, 2021, **21**(8), 4619–4631, DOI: [10.1021/acs.cgd.1c00467](https://doi.org/10.1021/acs.cgd.1c00467).
- 34 I. Suryawanshi, S. Srinidhi, S. Singh, R. Kalia, R. K. Kunchala, S. L. Mudavath and B. S. Naidu, Downshifting and upconversion dual mode emission from lanthanide doped GdPO_4 nanorods for unclonable anti-counterfeiting, *Mater. Today Commun.*, 2021, **26**, 102144, DOI: [10.1016/j.mtcomm.2021.102144](https://doi.org/10.1016/j.mtcomm.2021.102144).
- 35 I. Pushpendra, R. K. Kunchala, S. N. Achary and B. S. Naidu, $\text{NaBi}_{0.9}\text{Eu}_{0.1}(\text{MoO}_4)_2$ nanomaterials: tailoring the band gap and luminescence by La^{3+} substitution for light-emitting diodes, *ACS Appl. Nano Mater.*, 2019, **2**(9), 5527–5537, DOI: [10.1021/acsanm.9b01111](https://doi.org/10.1021/acsanm.9b01111).
- 36 I. Pushpendra, R. K. Kunchala, S. N. Achary, A. K. Tyagi and B. S. Naidu, Rapid, room temperature synthesis of Eu^{3+} Doped $\text{NaBi}(\text{MoO}_4)_2$ nanomaterials: structural, optical, and photoluminescence properties, *Cryst. Growth Des.*, 2019, **19**(6), 3379–3388, DOI: [10.1021/acs.cgd.9b00267](https://doi.org/10.1021/acs.cgd.9b00267).
- 37 M. A. Salim, R. Hussin, M. S. Abdullah, S. Abdullah, N. S. Alias, S. A. A. Fuzi, M. N. M. Yusuf and K. M. Mahbor, *Solid State Sci. Technol.*, 2009, **17**(2), 59–64.
- 38 J. Chen, J. Wang, F. Zhang, D. Yan, G. Zhang, R. Zhuo and P. Yan, Structure and photoluminescence property of Eu-doped SnO_2 nanocrystalline powders fabricated by sol-gel calcination process, *J. Phys. D*, 2008, **41**(10), 105306, DOI: [10.1088/0022-3727/41/10/105306](https://doi.org/10.1088/0022-3727/41/10/105306).
- 39 V. Dubey, J. Kaur and S. Agrawal, Effect of europium doping levels on photoluminescence and thermoluminescence of strontium yttrium oxide phosphor, *Mater. Sci. Semicond. Process.*, 2015, **31**, 27–37, DOI: [10.1016/j.mssp.2014.10.052](https://doi.org/10.1016/j.mssp.2014.10.052).
- 40 H. J. Song, L. Q. Zhou, Y. Huang, L. Li, T. Wang and L. Yang, Synthesis, characterization and luminescent properties of $\text{La}_2\text{Zr}_2\text{O}_7$: Eu^{3+} nanorods, *Chin. J. Chem. Phys.*, 2013, **26**(1), 83, DOI: [10.1063/1674-0068/26/01/83-87](https://doi.org/10.1063/1674-0068/26/01/83-87).
- 41 R. J. Chen, *Electrochem Soc.*, 1969, **116**, 1254; V. Pagonis and C. Michael, *Radiation Meas*, 1994, **23**, 131.
- 42 S. Doredrajit and S. Ingotombi, *J. Phys. D: Appl. Phys.*, 1995, **28**, 1509.
- 43 Y. Krish, P. D. Townsend and S. Soval, *Nuclear tracks and radiation measurements*, 1987, **13**, 115.
- 44 K. S. V. Nambi and S. Mitra, *Thermochim. Acta*, 1978, **27**, 61.
- 45 R. K. Gartia, S. J. Singh and P. S. Mazumdar, *Phys. Status Solidi A*, 1989, **114**, 407.
- 46 S. W. S. McKeever, *Thermoluminescence of solids*, Cambridge University Press, Cambridge, 1985, p. 390.
- 47 M. Martini and F. Meinardi, Thermally stimulated luminescence: new Perspectives in the study of defects in solids, *Riv. Nuovo Cimento*, 1997, **20–4**(8), 1–71.
- 48 R. Chen and S. W. S. McKeever, *Theory of Thermoluminescence and Related Phenomenon*, World Scientific, Singapore, New Jersey, London, Hong Kong, 1997, p. 559.
- 49 S. W. S. McKeever, M. Moscovitch and P. D. Townsend, *Thermoluminescence Dosimetry materials: Properties and Uses*, Nuclear Technology Publishing, Ashford, 1995, p. 221.
- 50 M. Martini, S. Paravisi and C. Iguori, A new high sensitivity spectrometer For 3D-thermoluminescence analysis, *Radiat. Prot. Dosim.*, 1996, **66**(4), 47–51.
- 51 G. kitis, E. C. Zoragoza and C. Furetta, *Appl. Radiat. Isot.*, 2005, **63**, 247–254.
- 52 K. S. Chung, H. S. Choe, J. I. Lee and S. Y. Chang, *Radiat. Prot. Dosimetry*, 2005, **115**, 136–143.
- 53 S. Som, M. Choedhury and S. K. Sharma, *Radiat. Phys. Chem.*, 2015, **110**, 51–58.
- 54 V. Dubey, N. S. Suryanarayana and J. Kaur, *J. Miner. Mater. Charact. Eng.*, 2010, **9**(12), 1101–1111.
- 55 R. Shrivastava, J. Kaur, V. Dubey and B. Jaykumar, *Bull. Mater. Sci.*, 2014, **37**(4), 925–929.
- 56 R. Shrivastava, J. Kaur, V. Dubey, N. S. Suryanarayana and B. Jaykumar, *Res. Chem. Intermed.*, 2014, **40**(2), 487–493.



- 57 V. Dubey, J. Kaur, N. S. Suryanarayana and K. V. R. Murthy, *Res. Chem. Intermed.*, 2015, **40**(2), 531–536.
- 58 J. Kaur, V. Dubey and N. S. Suryanarayana, *Res. Chem. Intermed.*, 2014, **39**(9), 4337–4349.
- 59 R. Tamrakar, V. Dubey, N. K. Swamy, R. Tiwari and S. V. N. Pammi, Research on Chemical, *Intermediates*, 2013, **39**(8), 3919–3923.
- 60 V. Dubey, J. Kaur, N. S. Suryanarayana and K. V. R. Murthy, *Res. Chem. Intermed.*, 2013, **39**(8), 3689–3697.
- 61 V. Dubey, *Thermoluminescence Study of Semaria Limestone of C.G.Basin*, LAP LAMBERT Academic Publishing, 2012, ISBN 978-3-8473-4210-6.
- 62 T. R. Williams, S. A. Winfield and J. N. Miller, Relative fluorescence quantum yields using a computer controlled luminescence spectrometer, *Analyst*, 1983, **108**, 1067.
- 63 D. Magde, J. H. Brannon, T. L. Cremers and J. Olmsted, *J. Phys. Chem.*, 1979, **83**(6), 696–699, DOI: [10.1021/j100469a012](#).
- 64 R. More and A. L. Capparelli, *J. Phys. Chem.*, 1980, **84**(14), 1870–1871, DOI: [10.1021/j100451a028](#).
- 65 W. H. Melhuish, *J. Phys. Chem.*, 1961, **65**(2), 229–235, DOI: [10.1021/j100820a009](#).
- 66 J. Q. Umberger and V. K. LaMer, *J. Am. Chem. Soc.*, 1945, **67**(7), 1099–1109.
- 67 A. Pardo, D. Reyman, J. M. L. Poyato and F. Medina, *J. Lumin.*, 1992, **51**(5), 269–274.
- 68 G. Weber and F. W. J. Teale, *Trans. Faraday Soc.*, 1957, **53**, 646–655.
- 69 P. S. Vincett, E. M. Voigt and K. E. Rieckhoff, *J. Chem. Phys.*, 1971, **55**, 4131–4140.
- 70 W. R. Dawson and M. W. Windsor, *J. Phys. Chem.*, 1968, **72**(9), 3251–3260.
- 71 E. P. Kirby and R. F. Steiner, *J. Phys. Chem.*, 1970, **74**(26), 4480–4490.
- 72 R. Rusakowicz and A. C. Testa, *J. Phys. Chem.*, 1968, **72**(7), 2680–2681.
- 73 W. H. Melhuish, *J. Phys. Chem.*, 1961, **65**(2), 229–235.
- 74 S. Hamai and F. Hirayama, *J. Phys. Chem.*, 1983, **87**(1), 83–89.
- 75 T. Swarnapriya, V. Hernandez, M. Ceron, P. Gomez, C. Wiechers, E. Carlos Montes, N.-M. Ricardo and S. Modesto, Thermoluminescence of Cu doped Li₂B₄O₇ + PTFE Annealed by Graphene exposed to X-rays and Gamma radiation, *J. Mol. Eng. Mater.*, 2020, **8**, 3–4, DOI: [10.1142/S2251237320500057](#).

