


Cite this: *RSC Adv.*, 2020, 10, 21049

Received 6th April 2020

Accepted 12th May 2020

DOI: 10.1039/d0ra03094e

rsc.li/rsc-advances

Photoluminescence, TGA/DSC and photocatalytic activity studies of Dy³⁺ doped SrY₂O₄ nanophosphors

Santosh P. Ghorpade,^a Nagaraju Kottam,^b Raveendra Melavanki^{ce} and N. R. Patil^{de}

Dy³⁺:SrY₂O₄ nanophosphors were prepared *via* a solution combustion method using glycine as an organic fuel. The structural, optical, and thermal properties of the nanophosphors were studied. Strain and crystal size were calculated *via* W–H analysis. The direct energy band gap is nearly 4.9 eV and photocatalytic studies reveal that Rh-B degradation of almost 50% can be achieved.

1. Introduction

In the last few decades, Dy³⁺-doped nanophosphors have attracted the attention of researchers because of their vast potential for use in a modern technological capacity. Dy³⁺ is one of the most efficient rare-earth elements for use as an activator. Dy³⁺ usually has two visible emission bands (blue and yellow). The blue emission is due to the ⁴F_{9/2} → ⁶H_{15/2} transition and the yellow emission band is due to the ⁴F_{9/2} → ⁶H_{13/2} transition.^{1–4} Trivalent rare earth (RE³⁺) doped inorganic nanophosphors may be broadly applied in display devices, fluorescent tubes, optical sensors, white light emitting diodes (w-LEDs), biological imaging, and scintillators due to their superior thermal and chemical stabilities and unique optical properties.^{5–15} Among these uses, w-LEDs are regarded as future light devices that may replace present traditional sources of light because of their low energy consumption, better consistency, higher brightness, longer operating times, and eco-friendly properties.^{14–19} W-light can be obtained *via* three possible means: firstly, *via* coalescing three basic coloured LEDs, like red, green and blue, even though this can be rather expensive and complicated, since each coloured LED is destroyed after a dissimilar time interval;^{20–25} secondly, *via* stimulating red, green and blue emitting nanophosphors with ultra-violet LED excitation, although the strong reverse absorption of blue light by red and green nanophosphors can

considerably decrease the efficacy of the device;¹⁴ and finally, *via* yellow-emitting Ce³⁺:Y₃Al₅O₁₂ under GaN-based blue LED excitation, which suffers from chromatic aberration and low w-light emitting performance after some period of functioning because of distinct degradation and the deficiency of the red constituent.¹³ Hence, further exciting research is required to examine powerful w-light producing nanophosphors in the presence of UV excitation.^{13–19} Therefore, it is a matter of interest to analyse the structural, morphological, optical, thermal, and, in particular, photoluminescence properties of Dy³⁺-doped SrY₂O₄ nanophosphors designed for warm w-LED uses.

In this article, a series of SrY₂O₄:Dy³⁺ nanophosphors was synthesized efficaciously with variable dopant molar concentrations (0.01 to 0.11%) *via* a solution combustion method (SCM). The obtained nanophosphors were calcinated and analysed *via* powder X-ray diffraction (XRD), high-resolution transmission electron microscopy (HRTEM) and field emission scanning electron microscopy (FESEM) techniques to obtain the crystal sizes, particle sizes and morphologies. Photoluminescence (PL) spectroscopy was used for the analysis of the excitation and emission spectra. FTIR and Raman spectroscopy studies were carried out to study the presence of different types of bonds in SrY₂O₄:Dy³⁺. UV-visible diffuse reflectance spectroscopy (UV-vis DRS) was used to determine the direct energy bandgaps. Thermogravimetric (TG/DSC) analysis was used to study the thermal solidities of the synthesized nanophosphors.

2. Experimental

2.1. Materials

Extremely pure strontium nitrate [Sr(NO₃)₂], yttrium nitrate hexahydrate [Y(NO₃)₃·6(H₂O)], dysprosium nitrate hexahydrate [Dy(NO₃)₃·6(H₂O)], glycine [C₂H₅NO₂], and rhodamine blue [C₂₈H₃₁ClN₂O₃] were used as materials in this study. All these constituents were purchased from Sigma-Aldrich.

^aDept. of Physics, P. C. Jabin Science College, Vidyanagar, Hubli-580031, Karnataka, India

^bDept. of Chemistry, MS Ramaiah Institute of Technology (Affiliated to Visvesvaraya Technological University, Belagavi), Bangalore-560054, Karnataka, India

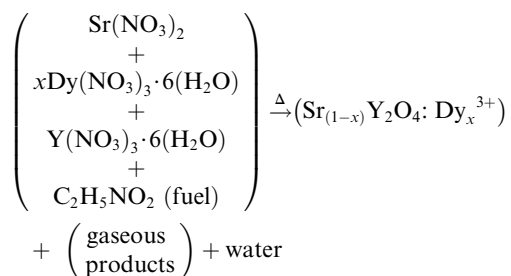
^cDept. of Physics, MSRIT, Bangalore-560054, Karnataka, India

^dDept. of Physics, B. V. B. College of Engg. & Tech., Vidyanagar, Hubli-580031, Karnataka, India. E-mail: patilnr23@gmail.com

^eDept. of Physics, Visvesvaraya Technological University, Belagavi-590018, Karnataka, India


2.2. Synthesis

Dy³⁺-activated SrY₂O₄ nanophosphors were successfully synthesized through a novel SCM with glycine as the fuel. The above-mentioned constituents were weighed according to the calculated compositions for different Sr_(1-x)Y₂O₄:Dy_x³⁺ ($x = 0.01, 0.03, 0.05, 0.07, 0.09, \text{ and } 0.11 \text{ mol\%}$) compounds and then added into a Borosil glass beaker, then 20 ml of double-distilled water was added into the beaker dropwise. Further, to achieve a homogenous solution, the mixture was stirred briskly using a magnetic stirrer for 10 min. Immediately the glass beaker was shifted into a muffle furnace, which was pre-heated to $500 \pm 10^\circ\text{C}$. After 5 min, oxidation occurs, resulting in the evaporation of N₂, CO₂ and water vapour. After the mixture was allowed to naturally cool, a large volume of white nanophosphors was obtained.



The flow chart of the SCM is shown in Fig. 1. The equivalent chemical reaction of the process is also given. The obtained nanophosphors were annealed using an alumina crucible at 1300°C for three hours in a muffle furnace.

2.3. Characterization

The phase pureness of Dy³⁺-doped SrY₂O₄ powder was analysed with a Rigaku Ultima IV powder X-ray diffractometer using Cu-K α radiation ($\lambda = 1.5406 \text{ \AA}$). The superficial morphology and particle size were examined using FESEM (JEOL JSM-7100F) at an accelerating voltage of 30 kV and HRTEM (a JEOL JEM 2100 microscope operating at 200 kV; source: LaB6). Photoluminescence spectroscopy studies were carried out using a Shimadzu RF-5301-PC fluorescence spectrophotometer. Fourier transform infrared (FTIR) spectra were recorded with a NICOLET 6700 spectrometer (USA) using KBr as a reference. UV-vis diffuse reflectance spectroscopy (DRS) data was obtained with the help of a UV 3092 UV-vis spectrophotometer. The DSC/TGA (differential scanning calorimetry/thermal gravimetric analysis) studies were accomplished using a NETZSCH STA

449F3 thermogravimeter instrument. Room temperature Raman spectroscopic studies were done using LabRM HR revolution HORIBA apparatus. The photocatalytic activity in rhodamine B (Rh-B) solution was inspected using a SPECORD-210 Plus (Germany) UV-visible spectrophotometer in the range from 200 to 800 nm.

2.4. Photocatalytic activity

Rh-B was used to estimate the photocatalytic activity of undoped and Dy³⁺ (0.03, 0.07, 0.11 mol%) doped SrY₂O₄ nanophosphors. The photocatalytic investigations were conducted at room temperature using a round glass container with a surface area of 176 cm^2 and an average-pressure mercury vapour lamp as the ultra-violet light source with a wavelength of 365 nm at 125 W. The light flux is around 8 mW cm^{-2} according to a ferrioxalate actinometric approach.^{26,27} The UV light was made to focus directly onto the reaction mixture. Experiments were conducted using double distilled water. Usually, an experiment comprises 50 mg of photocatalyst distributed in 100 ml of 10 ppm Rh-B solution. The reaction mixture was stirred vigorously for 30 min prior to radiation treatment to guarantee the development of adsorption/desorption equilibrium, and magnetic stirring was continued throughout the entire experiment. The amount of adsorption is given by eqn (1):

$$Q = \frac{(C_0 - C)V}{W} \quad (1)$$

where Q is the degree of adsorption, C_0 and C are the initial concentration (at time = 0 min) and concentration after adsorption (at time = t min), V is the volume of the reaction mixture, and W is the quantity of catalyst in grams; the unit of Q is ppm ml mg^{-1} . About 5 ml of solution was taken from the suspension after a certain time intervals (say 15 min), centrifuged, and then subjected to spectrophotometric analysis; the maximum absorption was observed at a wavelength of 555 nm and the residual concentration of Rh-B was estimated. The dye degradation rate was calculated from the following eqn (2):²⁸

$$\text{Rate of photodegradation} = \frac{(C_0 - C)}{C_0} \times 100 \quad (2)$$

3. Results and discussion

3.1. Powder X-ray diffraction analysis

X-ray diffraction (XRD) spectra were recorded in the range of $15^\circ \leq 2\theta \leq 80^\circ$ using Cu-K α radiation ($\lambda = 1.5406 \text{ \AA}$). As we know,

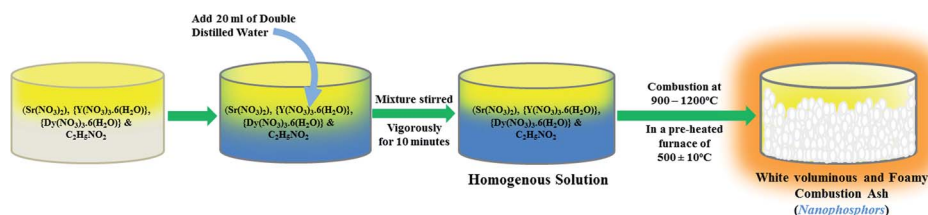


Fig. 1 A flow chart showing the SCM to produce SrY₂O₄:Dy³⁺ nanophosphors.



SrY₂O₄ fits the orthorhombic CaFe₂O₄ structure with the *Pnam* space group; it has a volume of 408.18 Å³, and cell parameters of *a* = 10.07 Å, *b* = 11.91 Å, and *c* = 3.41 Å.²³ The XRD patterns of annealed SrY₂O₄ nanophosphors doped with Dy³⁺ (0.01, 0.03, 0.05, 0.07, 0.09, and 0.11 at%) at different molar concentrations are shown in Fig. 2(a), and the corresponding peaks can be indexed to the orthorhombic structure and match the typical SrY₂O₄ standard indicated in JCPDS Card No. 32-1272. Fig. 2(b) shows the crystal structure of the host SrY₂O₄:Dy³⁺ nanophosphors created using the VESTA program.²⁹

Typically, the average crystallite size of phosphors can be evaluated *via* the Debye Scherrer formula (3):

$$D = \frac{0.9\lambda}{\beta \cos \theta} \quad (3)$$

where *D* is the typical nanocrystalline size, λ is the wavelength of the X-rays (1.5406 Å), β is the full width at half maximum (FWHM) in terms of radians, and θ is the Bragg diffraction angle of a measured peak in terms of radians. The strongest diffraction heights were considered for computing the crystallite sizes, and the data are tabulated in Table 1. With an increase in the Dy³⁺ concentration, the crystallite size also varies moderately, and it is found that the average crystallite size is about 38 nm.

The Williamson–Hall (W–H) fitting method was applied to evaluate the strain (ϵ) and crystallite size values of the samples.^{30,31} Induced strain in nanophosphors may be caused by virtue of crystal faults and defects. Induced strain is estimated *via* eqn (4):

$$\epsilon = \frac{\beta}{4 \tan \theta} \quad (4)$$

If the crystal size and strain contribution to enlargement are independent of one another and both possess a Cauchy-like profile,³¹ then the measured line breadth is merely the sum of eqn (3) and (4):

$$\beta = \frac{0.9\lambda}{D \cos \theta} + 4\epsilon \tan \theta. \quad (5)$$

Further, rearranging eqn (5), one can get:

$$\beta \cos \theta = \frac{0.9\lambda}{D \cos \theta} + 4\epsilon \sin \theta. \quad (6)$$

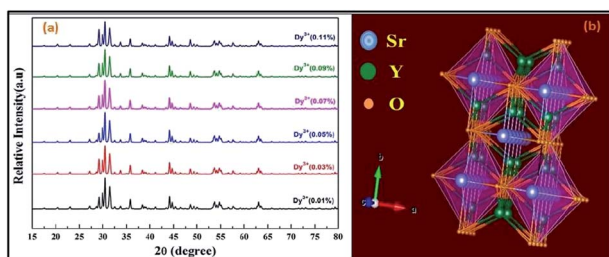


Fig. 2 (a) X-ray diffraction spectra of SrY₂O₄ nanophosphors doped with Dy³⁺ at different molar concentrations and annealed at a temperature of 1300 °C and (b) the crystal structure of the host SrY₂O₄ nanophosphors.

Table 1 The crystallite sizes of various concentrations of Dy³⁺-doped SrY₂O₄ phosphors annealed at a temperature of 1300 °C

Phosphor	Debye Scherrer method	W–H method	
	Crystal size (nm)	Lattice strain	Crystal size (nm)
Sr _{0.99} Y ₂ O ₄ :Dy _{0.01} ³⁺	38	0.00109	67
Sr _{0.97} Y ₂ O ₄ :Dy _{0.03} ³⁺	40	0.00142	63
Sr _{0.95} Y ₂ O ₄ :Dy _{0.05} ³⁺	40	0.00158	50
Sr _{0.93} Y ₂ O ₄ :Dy _{0.07} ³⁺	39	0.00110	58
Sr _{0.91} Y ₂ O ₄ :Dy _{0.09} ³⁺	38	0.00147	76
Sr _{0.89} Y ₂ O ₄ :Dy _{0.11} ³⁺	37	0.00129	69

From eqn (6), the slope of the linear fit of a plot of $\beta \cos \theta$ versus $4 \sin \theta$ is equal to the strain ϵ and linear fits of the graphs of 0.01 to 0.11 at% Dy³⁺-doped SrY₂O₄ are shown in Fig. 3. The induced strain values are in the range of 0.00109 to 0.00158, and the positive slopes specify the existence of tensile strain in the arrangement. Similar behaviour has been indicated for Eu³⁺-doped SrWO₄ nanophosphors³² and a Gd³⁺-co-doped CaMoO₄:Eu³⁺ system.³³ Further, the W–H fitting method was used to compute the crystal size by using the intercept $\left(\frac{0.9\lambda}{D \cos \theta}\right)$ of the line with the y-axis; crystallite size (*D*) values were calculated and they average 64 nm, as tabulated in Table 1.

3.2. FESEM and HRTEM analysis

With the help of HRTEM, the morphologies of the formed nanophosphors were analysed, showing that the particles remain nearly spherical with their usual size.^{34–36} The average particle size is about 49 nm, as shown in Fig. 4. FESEM morphological images of SrY₂O₄:Dy³⁺ nanophosphors at different magnifications are shown in Fig. 5. From the FE-SEM image, it is clear that the SrY₂O₄:Dy³⁺ nanophosphors are spherical in nature with quite identical sizes and closely crowded with each other.^{34–37} Also from Fig. 5 it is found that the average phosphor particle size is around 55 nm; these very crowded phosphor particles avoid the scattering of light,

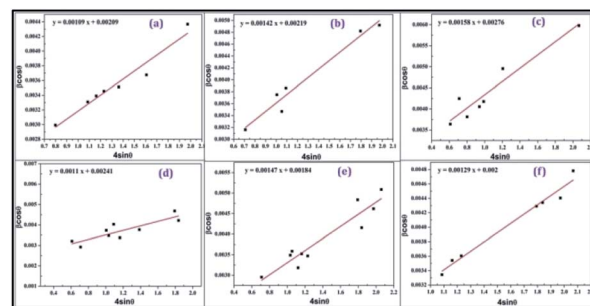


Fig. 3 W–H plots of $\beta \cos \theta$ against $4 \sin \theta$ for SrY₂O₄ doped with Dy³⁺ at different molar concentrations and annealed at 1300 °C: (a) 0.01%, (b) 0.03%, (c) 0.05%, (d) 0.07%, (e) 0.09% and (f) 0.11%.



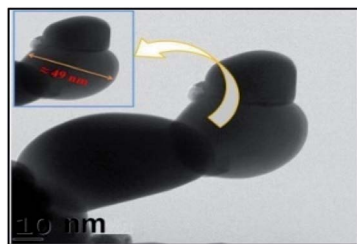


Fig. 4 A HR-TEM image of 0.11 at% Dy^{3+} -doped SrY_2O_4 annealed at 1300°C (the inset indicates an extended view of a single particle having an average diameter of 49 nm).

creating strong light production and acting as a suitable material for potential applications in the area of lighting.

3.3. Photoluminescence study

Room temperature photoluminescence excitation and emission spectra were recorded. Fig. 6(a) shows the excitation spectra of the $\text{SrY}_2\text{O}_4:\text{Dy}^{3+}$ nanophosphors, and it is observed that prominent excitation peaks appear at wavelengths of 327 nm, 354 nm and 367 nm. Of the three peaks, the peak at 354 nm has the highest intensity and so we excited the $\text{SrY}_2\text{O}_4:\text{Dy}^{3+}$ sample at this wavelength.

At an excitation wavelength of 354 nm, the Dy^{3+} -doped SrY_2O_4 nanophosphors show broad blue and yellow colour emission at wavelengths of 491 nm ($^4\text{F}_{9/2} \rightarrow ^6\text{H}_{15/2}$) and 581 nm ($^4\text{F}_{9/2} \rightarrow ^6\text{H}_{13/2}$) with the highest intensity,¹⁻⁴ as shown in Fig. 6(b). The yellow $^4\text{F}_{9/2} \rightarrow ^6\text{H}_{13/2}$ emission is associated with a forced electric dipole type transition, possible only at low symmetry *via* a non-inversion centre, and the peak intensity is strongly dependent on the crystal-field environment. The blue emission is related to the $^4\text{F}_{9/2} \rightarrow ^6\text{H}_{15/2}$ magnetic dipole transition.²⁶ Also, it is observed that with a rise in the Dy^{3+} concentration, there is an increase in the intensity of the emission spectrum. In addition to these peaks, there is the presence of small peaks; the source of these peaks may be magnetic dipoles or electric dipoles, after emitting to the completion levels, which is dependent on the sites of Dy^{3+} in the lattice of SrY_2O_4 , and the type of transfer is determined by selection rules.³⁴⁻³⁷ The observed photoluminescence mechanism is described schematically through the energy level diagram in Fig. 7. From the PL emission spectra, it is noticed that the dopant Dy^{3+} emits blue and yellow colour light

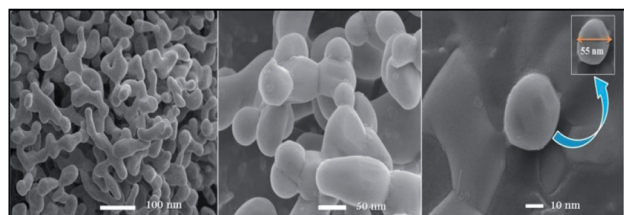


Fig. 5 FE-SEM micrographs of $\text{SrY}_2\text{O}_4:\text{Dy}^{3+}$ (0.11 at%) nanophosphors at various resolutions.

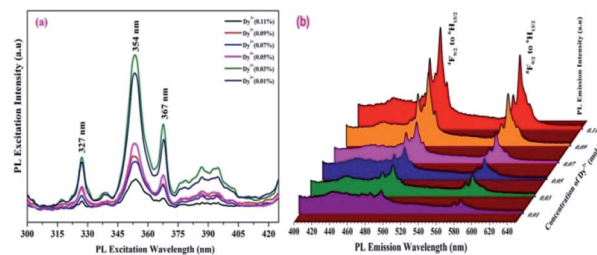


Fig. 6 Photoluminescence (a) excitation and (b) emission spectra of Dy^{3+} (0.01 to 0.11 mol%)-doped SrY_2O_4 nanophosphors.

successfully, which is helpful for many types of lighting equipment.

Fig. 8 shows the Commission Internationale de l'Eclairage (CIE) chromaticity graph of $\text{SrY}_2\text{O}_4:\text{Dy}^{3+}$ (0.01, 0.03, 0.05, 0.07, 0.09, and 0.11 at%) nanophosphors. The calculated CIE 'x' and 'y' colour co-ordinates of all samples are presented in Table 2. At an excitation wavelength of 354 nm, blue and yellow light emission is perceived.¹⁻⁴ It is found that, as the concentration of Dy^{3+} increases, the CIE coordinates initially are positioned in the blue region, further moving to the area that is very close to standard w-light illumination. These observed x and y coordinate values affirm that the colour is tuneable from blue-yellow to white in the visible spectrum region suitable for w-LED applications.³⁴⁻³⁷

3.4. FTIR studies

FTIR spectra of the Dy^{3+} -doped SrY_2O_4 nanophosphors over wavenumbers ranging from 400 to 4000 cm^{-1} are shown in Fig. 9. In the FTIR spectra, the broad intense absorption peak at 3462 cm^{-1} implies O-H stretching vibrations, which is due to the existence of water molecules present superficially on the nanophosphors; two close weak peaks are detected at 2922 cm^{-1} and 2845 cm^{-1} , which correspond to C-H symmetric and asymmetric stretching; and the weak extensive absorption peak observed at 1750 cm^{-1} can be allocated to C=C symmetric

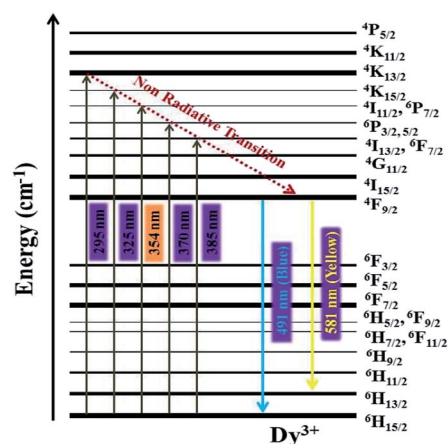


Fig. 7 The energy level diagram of Dy^{3+} -doped SrY_2O_4 nanophosphors at $\lambda_{\text{Excitation}} = 354\text{ nm}$.

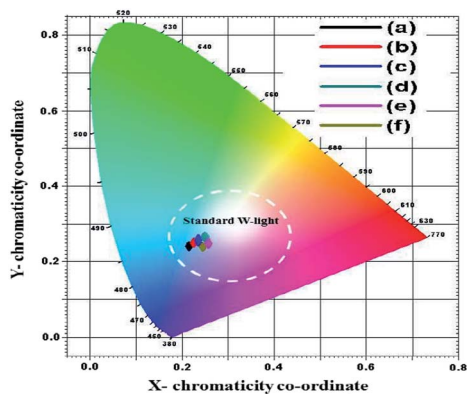


Fig. 8 The Commission Internationale de l'Eclairage (CIE) chromaticity coordinates of $\text{SrY}_2\text{O}_4:\text{Dy}^{3+}$ nanophosphors with varied Dy^{3+} ion molar concentrations: (a) 0.01, (b) 0.03, (c) 0.05, (d) 0.07, (e) 0.09, and (f) 0.11 at%.

Table 2 The CIE x and y co-ordinates of $\text{Sr}_{(1-x)}\text{Y}_2\text{O}_4:\text{Dy}_x^{3+}$ with variable molar concentrations

Nanophosphor	CIE co-ordinates	
	x	y
$\text{Sr}_{0.99}\text{Y}_2\text{O}_4:\text{Dy}_{0.01}^{3+}$	0.2607	0.2720
$\text{Sr}_{0.97}\text{Y}_2\text{O}_4:\text{Dy}_{0.03}^{3+}$	0.2616	0.2761
$\text{Sr}_{0.95}\text{Y}_2\text{O}_4:\text{Dy}_{0.05}^{3+}$	0.2534	0.2580
$\text{Sr}_{0.93}\text{Y}_2\text{O}_4:\text{Dy}_{0.07}^{3+}$	0.2562	0.2611
$\text{Sr}_{0.91}\text{Y}_2\text{O}_4:\text{Dy}_{0.09}^{3+}$	0.2404	0.2415
$\text{Sr}_{0.89}\text{Y}_2\text{O}_4:\text{Dy}_{0.11}^{3+}$	0.2395	0.2409

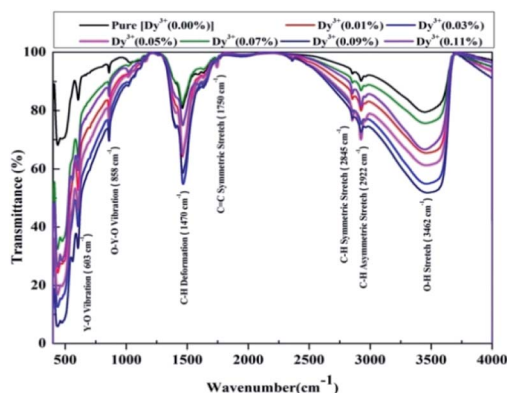


Fig. 9 FT-IR spectra of SrY_2O_4 nanophosphors doped with Dy^{3+} at different concentrations and annealed at 1300°C .

stretching vibrations. The sharp strong peak at 1470 cm^{-1} be allocated to the bending vibrations of C-H deformation. The medium sharp peak at 858 cm^{-1} corresponds to O-Y-O deformation, owing to the enlargement of the lattice of SrY_2O_4 after calcination at 1300°C . The presence of the O-Y-O vibration peak is seen at 603 cm^{-1} , and Y-O vibration is observed at 490 cm^{-1} .^{36,37} From Fig. 9, it is clear that the peaks do not shift with a rise in the Dy^{3+} concentration, but the percentage

absorption rises as the concentration is increased, clearly indicating the stability of the structure. Thus, the CaFe_2O_4 structure of the Dy^{3+} -doped SrY_2O_4 nanophosphors is verified via FTIR analysis.

3.5. Raman studies

Fig. 10 shows Raman spectra of $\text{SrY}_2\text{O}_4:\text{Dy}^{3+}$ at a wavelength of 532 nm in the frequency range of 100 to 600 cm^{-1} . It is observed that for 0.01%, 0.03%, 0.05%, 0.07%, and 0.09% Dy^{3+} , no prominent peaks are found, but for 0.11% Dy^{3+} , noticeable Raman bands are perceived at 104 , 144 , 299 , 380 and 419 cm^{-1} , which are related to the SrY_2O_4 orthorhombic structure. In Fig. 10, the peak situated at 483 cm^{-1} can be ascribed to the symmetric stretching of Y-O, whereas the peak at 219 cm^{-1} is allocated to the symmetric bending of O-Y-O. The peaks at 419 and 380 cm^{-1} are ascribed to O-Y-O antisymmetric stretching, and the peaks at 144 and 109 cm^{-1} are ascribed to Y-O anti-symmetric bending.³⁸

3.6. DSC/TGA analysis

Fig. 11 illustrates the synchronized DSC/TGA curves along with the DTA curves of SrY_2O_4 doped with Dy^{3+} at a concentration of 0.11%. Under a nitrogen atmosphere, samples were measured in the temperature range of 28 to 1400°C at a heating rate of $10^\circ\text{C min}^{-1}$.

The TG study (Fig. 11) shows weight loss of 11.6% between 28°C and 300°C , 2.6% in the range of 300 – 750°C , 1.8% in the range of 750 – 825°C , and insignificant loss beyond 825°C for 0.11 at% Dy^{3+} -doped SrY_2O_4 . The mass loss up to 300°C is assigned to dehydration of the sample, while mass loss up to 600°C is because of the evaporation of organic materials like glycine and some impurities.³⁸ In the DTA curve, there are four exothermic peaks in the locality of 100°C , 260°C , 450°C and 800°C , which relate to water molecule evaporation, organic molecule decay, and the phase creation of $(0.11\%)\text{Dy}^{3+}:\text{SrY}_2\text{O}_4$.

To analyse the heat flow against temperature under a nitrogen gas atmosphere related to the transitions in $\text{SrY}_2\text{O}_4:(0.11\%)\text{Dy}^{3+}$, DSC was performed, and both exothermic and endothermic peaks were found. The peak at around 166°C

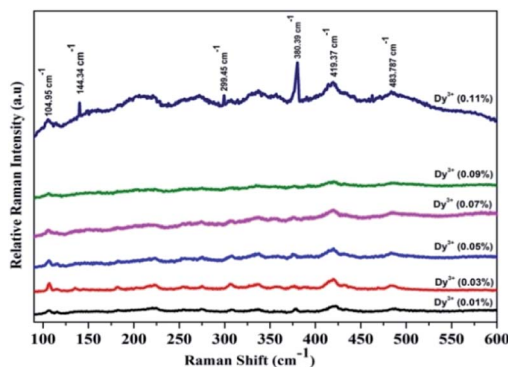


Fig. 10 Room temperature Raman spectra of $\text{SrY}_2\text{O}_4:\text{Dy}^{3+}$ nanophosphors with varied Dy^{3+} concentrations at an excitation wavelength of 532 nm after being treated at 1300°C for 3 h.

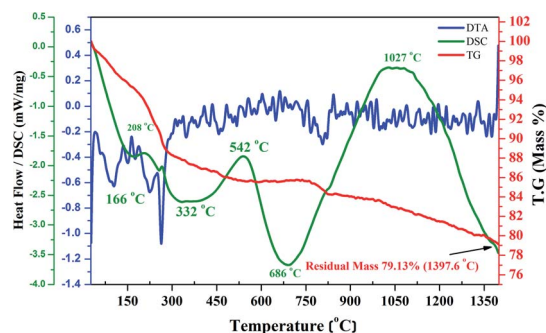


Fig. 11 Simultaneous DSC/TG analysis of SrY_2O_4 doped with Dy^{3+} at a concentration of 0.11% concentration along DTA.

signifies mass loss owing to the vaporization of water and glycine. The exothermic peaks at around 332 and 686 °C imply the evaporation of organic compounds. The big and sharp endothermic DSC peaks at about 542 and 1027 °C suggest the phase establishment of $\text{SrY}_2\text{O}_4:(0.11\%)\text{Dy}^{3+}$. These outcomes indicate that the $\text{SrY}_2\text{O}_4:(0.11\%)\text{Dy}^{3+}$ nanophosphors are stable with respect to rising temperature, and they may be useful in the areas of lighting and display-related devices.³⁸

3.7. UV-visible diffuse reflectance study

From the UV-vis diffuse reflectance spectroscopic data, the direct energy bandgaps of the un-doped and Dy^{3+} -doped SrY_2O_4 nanophosphors were evaluated by plotting graphs of $[F(R)h\nu]^2$ against the energy of the incident photons ($h\nu$).^{39,40} The Kubelka–Munk (K–M) approximation is one suitable method used for the estimation of the direct energy bandgap of pure SrY_2O_4 and Dy^{3+} -doped nanophosphors and the respective K–M plots are shown in Fig. 12.

According to the K–M approximation function, the correlation between the optical absorption coefficient $[F(R)h\nu]^2$ and the energy bandgap ($h\nu$) is given in eqn (7):

$$[F(R)h\nu]^2 = A(h\nu - E_g) \quad (7)$$

where $F(R)$, h , ν , E_g , and A are the K–M function, Planck's constant, the frequency, the energy band gap and a proportionality constant, respectively.^{41,42}

The energy bandgap is projected by extrapolating the rectilinear region of the plot of $[F(R)h\nu]^2$ versus $h\nu$ to the photon energy axis, as shown in Fig. 12. The corresponding energy bandgap values are estimated for pure SrY_2O_4 and $\text{SrY}_2\text{O}_4:(0.01\text{--}0.11\%)\text{Dy}^{3+}$ nanophosphors, as listed in Table 3.

A study of Table 3 specifies that un-doped SrY_2O_4 has a direct energy band gap of 5.04 eV, but the occurrence of dopant ions reduces the energy bandgap; this is due to the substitution of Sr^{2+} by Dy^{3+} in the crystalline arrangement, which is able to generate additional energy levels. The creation of a charge transfer band in between the valence band (VB) and conduction band (CB) of un-doped SrY_2O_4 is accountable for the reduction in the energy bandgaps of the doped samples. In addition, the reduction of the energy bandgap is attributed to an increase in the superficial defect concentrations of the doped samples.⁴³

3.8. Photocatalytic activity

The photocatalytic performances of un-doped and Dy^{3+} (0.03, 0.07 and 0.11 at%)-doped SrY_2O_4 nanophosphors were analysed via the decolourization of Rh-B in aqueous solution. Graphs of

Table 3 The energy band gaps for various concentrations of Dy^{3+} -doped SrY_2O_4 nanophosphors annealed at a temperature of 1300 °C

Nanophosphor	Direct energy band gap (eV)
$\text{Sr}_{0.00}\text{Y}_2\text{O}_4:\text{Dy}_{0.00}^{3+}$	5.04
$\text{Sr}_{0.99}\text{Y}_2\text{O}_4:\text{Dy}_{0.01}^{3+}$	4.89
$\text{Sr}_{0.97}\text{Y}_2\text{O}_4:\text{Dy}_{0.03}^{3+}$	4.91
$\text{Sr}_{0.95}\text{Y}_2\text{O}_4:\text{Dy}_{0.05}^{3+}$	4.83
$\text{Sr}_{0.93}\text{Y}_2\text{O}_4:\text{Dy}_{0.07}^{3+}$	4.90
$\text{Sr}_{0.91}\text{Y}_2\text{O}_4:\text{Dy}_{0.09}^{3+}$	4.93
$\text{Sr}_{0.89}\text{Y}_2\text{O}_4:\text{Dy}_{0.11}^{3+}$	4.89

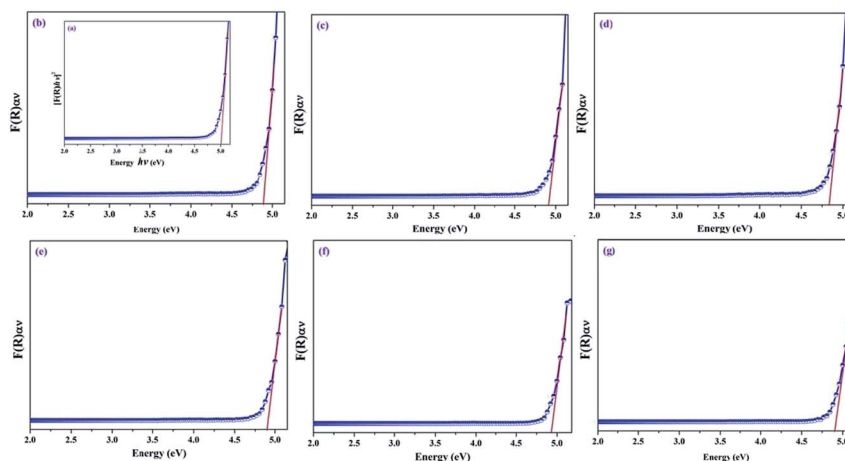


Fig. 12 Kubelka–Munk approximation plots for the direct band transition of $\text{SrY}_2\text{O}_4:\text{Dy}^{3+}$ nanophosphors with varying Dy^{3+} molar concentrations: (a) un-doped (inset), (b) 0.01, (c) 0.03, (d) 0.05, (e) 0.07, (f) 0.09, and (g) 0.11 at%.



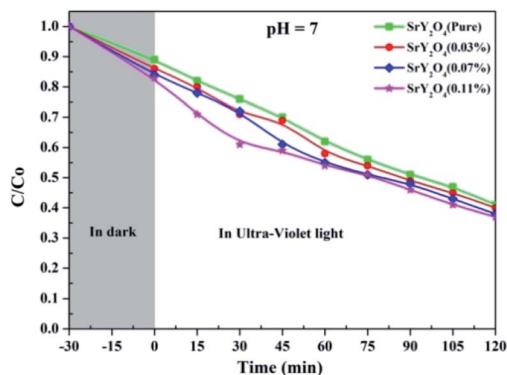


Fig. 13 The relative concentration variations against irradiation time for SrY_2O_4 nanophosphors doped with different molar concentrations of Dy^{3+} .

C/C_0 versus the UV light irradiation time for the decolourization of Rh-B at pH 7 were obtained, with the reaction was performed in the dark for 30 min followed by UV irradiation for 120 min (Fig. 13). The self-catalysis of Rh-B was ruled out, since only a small quantity of Rh-B decolorized in the absence of the photocatalyst; this verifies that the decolourization is due to photocatalysis. The Rh-B photo-decolourization efficiency gradually declines as the Dy^{3+} dopant concentration rises.

Photo-degradation arises as a result of Dy^{3+} stimulating charge separation, charge-carrier recombination and the amount of interfacial charge transfer; it can act as a mediator and affect quantum productivity.^{44,45} Dy^{3+} -doped SrY_2O_4 nanophosphors are treated with UV light; electrons present in the VB are transmitted to the CB, leading to the creation of some holes in the VB and commencing a sequence of reactions to create hydroperoxyl and hydroxyl radicals.

These hydroxyl radicals are strong oxidants that destroy dangerous impurities.⁴⁶ In the said method, the undesirable recombination of electrons and holes can arise too quickly, reducing the photocatalytic activity. Dy^{3+} might take electrons from the CB of SrY_2O_4 to form Dy^{2+} , which results in the relocation of an electron to dissolved O_2 to create a superoxide radical,⁴⁷ which prevents the recombination of light-generated electrons and holes. Therefore, Dy^{3+} on the surfaces of SrY_2O_4 nanophosphors could act as an electron hunter.

The use of photocatalysts and also the control of photocatalytic properties is of serious interest during the extended usage of photocatalysts, since the photocatalytic activity usually declines as an outcome of photo-corrosion due to frequent use.⁴⁸

4. Conclusions

Dy^{3+} -doped SrY_2O_4 nanophosphors were synthesized through an easy and low-cost SCM. Powder XRD data highlight the monophasic orthorhombic CaFe_2O_4 structure with a $Pn\text{am}$ space group. FESEM micrographs show the spherical nature, with quite identical size and a closely crowded morphology. The Debye-Scherrer and W-H methods are used to calculate the

particle sizes. The strain induced is in the range of 0.00109 to 0.00158. PL emission spectra reveal broad intense blue and yellow emission peaks at an excitation wavelength of 354 nm. In addition, the samples showed admirable CIE chromaticity coordinates (x , y), an outcome showing relative suitability for display and W-LED uses. The thermal stability was verified *via* DSC/TGA analysis for display applications. UV-vis DRS studies show that the energy band gap decreases upon Dy^{3+} doping. Photocatalytic activity studies highlight almost 50% Rh-B dye degradation under UV light treatment. Altogether, these results indicate that SrY_2O_4 is a likely contender for the improvement of blue and yellow nanophosphors in w-LEDs, and it could provide a suitable approach for developing efficient photocatalysts to destroy contaminants, particularly using UV-light.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors acknowledge Dr Vikas Dubey, Dept. of Physics, Bhilai Institute of Technology, Raipur, Chhattisgarh, India for providing the instrumental apparatus of a Shimadzu RF-5301-PC fluorescence spectrophotometer to measure the photoluminescence excitation and emission spectroscopy data.

References

- 1 P. Haritha, I. R. Martín, C. S. Dwaraka Viswanath, N. Vijaya, K. Venkata Krishnaiah, C. K. Jayasankar, D. Haranath, V. Lavín and V. Venkatramu, *Opt. Mat.*, 2017, **70**, 16.
- 2 P. Babu, K. H. Jang, E. S. Kim, L. Shi and H. J. Seo, *J. Korean Phys. Soc.*, 2009, **54**, 1488.
- 3 S. K. Gupta, M. Mohapatra, V. Natarajan and S. V. Godbole, *Int. J. Appl. Ceram. Technol.*, 2012, **10**, 593.
- 4 S. K. Gupta, M. Kumar, V. Natarajan and S. V. Godbole, *Opt. Mater.*, 2013, **35**, 2320.
- 5 Y. Huang, H. You, G. Jia, Y. Song, Y. Zheng, M. Yang, K. Liu and N. Guo, *J. Phys. Chem. C*, 2010, **114**, 18051.
- 6 H. Qiu, G. Chen, L. Sun, S. Hao, G. Han and C. Yang, *J. Mater. Chem.*, 2011, **21**, 17202.
- 7 C. Zhang, P. Ma, C. Li, G. Li, S. Huang, D. Yang, M. Shang, X. Kang and J. Lin, *J. Mater. Chem.*, 2011, **21**, 717.
- 8 Y. Lei, M. Pang, W. Fan, J. Feng, S. Song, S. Dang and H. Zhang, *Dalton Trans.*, 2011, **40**, 142.
- 9 X. Zhai, S. Liu, X. Liu, F. Wang, D. Zhang, G. Qin and W. Qin, *J. Mater. Chem. C*, 2013, **1**, 1525.
- 10 H. Liu, W. Lu, H. Wang, L. Rao, Z. Yi, S. Zeng and J. Hao, *Nanoscale*, 2013, **5**, 6023.
- 11 F. Wang and X. Liu, *Chem. Soc. Rev.*, 2009, **38**, 976.
- 12 K. Nini, N. P. Singh, L. P. Singh and S. S. Krishna, *Nanoscale Res. Lett.*, 2015, **10**, 347.
- 13 H. Guan, Y. Song, P. Ma, M. Chang, J. Chen, Y. Wang, B. Yuan and H. Zou, *RSC Adv.*, 2016, **6**, 53444.
- 14 P. Haritha, I. R. Martín, K. Linganna, V. Monteseuro, P. Babu, S. F. Le-on-Luis, C. K. Jayasankar,



- U. R. Rodríguez-Mendoza, V. Lavín and V. Venkatramu, *J. Appl. Physiol.*, 2014, **116**, 174308.
- 15 M. Shang, C. Li and J. Lin, *Chem. Soc. Rev.*, 2014, **43**, 1372.
- 16 B. Liu, L. Kong and C. Shi, *J. Lumin.*, 2007, **122–123**, 121.
- 17 X. Huang, S. Han, W. Huang and X. Liu, *Chem. Soc. Rev.*, 2013, **42**, 173.
- 18 X. Liu, R. Deng, Y. Zhang, Y. Wang, H. Chang, L. Huang and X. Liu, *Chem. Soc. Rev.*, 2015, **44**, 1479.
- 19 F. Wang and X. Liu, *Chem. Soc. Rev.*, 2009, **38**, 976.
- 20 F. Auzel, *Chem. Rev.*, 2004, **104**, 139.
- 21 Q. Zhang, C. F. Wang, L. T. Ling and S. Chen, *J. Mater. Chem. C*, 2014, **22**, 4358.
- 22 R. Rajeswari, C. K. Jayasankar, D. Ramachari and S. Surendra Babu, *Ceram. Int.*, 2013, **39**, 7523.
- 23 Y. Shimizu, K. Sakano, Y. Noguchi and T. Moriguchi, *US Pat.*, 59998925, 1999.
- 24 Y. Hu, W. Zhuang, H. Ye, D. Wang, S. Zhang and X. Huang, *J. Alloys Compd.*, 2005, **390**, 226.
- 25 X. He, M. Guan, N. Lian, J. Sun and T. Shang, *J. Alloys Compd.*, 2010, **492**, 452.
- 26 C. A. Parker, A New Sensitive Chemical Actinometer. I., *Proc. R. Soc. London, Ser. A*, 1953, **220**, 104.
- 27 C. G. Hatchard and C. A. Parker, *Proc. R. Soc. London, Ser. A*, 1956, **235**, 518.
- 28 S. A. Khayyat, M. S. Akhtar and A. Umar, *Mater. Lett.*, 2012, **81**, 239.
- 29 K. Momma and F. Izumi, *J. Appl. Crystallogr.*, 2008, **41**, 653.
- 30 L. Robindro Singh, R. S. Ningthoujam, V. Sudarsan, I. Srivastava, S. Dorendrajit Singh, G. K. Dey and S. K. Kulshreshtha, *Nanotechnology*, 2008, **19**, 055201.
- 31 V. D. Mote, Y. Purushotham and B. N. Dole, *Journal of Theoretical and Applied Physics*, 2012, **6**, 6.
- 32 Maheshwary, B. P. Singh, J. Singh and R. A. Singh, *RSC Adv.*, 2014, **4**, 32605.
- 33 B. P. Singh, A. K. Parchur, R. S. Ningthoujam, A. A. Ansari, P. Singh and S. B. Rai, *Dalton Trans.*, 2014, **43**, 4770.
- 34 E. Pavitra, G. Seeta Rama Raju, J.-h. Oh and J. Su Yu, *New J. Chem.*, 2014, **38**, 3413.
- 35 V. Dubey, J. Kaur, S. Agrawal, N. S. Suryanarayana and K. V. R. Murthy, *Optik*, 2013, **124**, 5585.
- 36 V. Dubey, J. Kaur and S. Agrawal, *Mater. Sci. Semicond. Process.*, 2015, **31**, 27.
- 37 E. Pavitra, G. Seeta Rama Raju, Y. H. Ko and J. Su Yu, *Phys. Chem. Chem. Phys.*, 2012, **14**, 11296.
- 38 Maheshwary, B. P. Singh and R. A. Singh, *New J. Chem.*, 2015, **39**, 4494.
- 39 B. Murphy, *Sol. Energy Mater. Sol. Cells*, 2007, **91**, 1326.
- 40 S. P. Ghorpade, R. Hari Krishna, R. M. Melavanki, V. Dubey and N. R. Patil, *Optik*, 2020, **208**, 164533.
- 41 Z. Jiao, T. Chen, J. Xiong, T. Wang, G. Lu, J. Ye and Y. Bi, *Sci. Rep.*, 2013, **3**, 1.
- 42 T. Selvalakshmi, S. Sellaiyan, A. Uedonob and A. Chandra Bose, *RSC Adv.*, 2014, **4**, 34257.
- 43 A. G. Bispo Jr, D. A. Ceccato, S. A. M. Lima and A. M. Pires, *RSC Adv.*, 2017, **7**, 53752.
- 44 Y. Zong, Z. Li, X. Wang, J. Ma and Y. Men, *Ceram. Int.*, 2014, **40**, 10375.
- 45 S. Liu and Y. Chen, *Catal. Commun.*, 2009, **10**, 894.
- 46 Z. Zhang, W. Wang, E. Gao, M. Shang and J. Xu, *J. Hazard. Mater.*, 2011, **196**, 255.
- 47 A. S. Weber, A. M. Grady and R. T. Koodali, *Catal. Sci. Technol.*, 2012, **2**, 683.
- 48 P. Wang, J. Wang, X. Wang, H. Yu, J. Yu, M. Lei and Y. Wang, *Appl. Catal. B Environ.*, 2013, **132–133**, 452.

