

# Photoluminescence properties of $\gamma\text{-Ca}_3(\text{PO}_4)_2\text{:Sm}^{3+}$ prepared under high-pressure and high-temperature conditions



Weihong Xue<sup>a,b,\*</sup>, Shuangmeng Zhai<sup>a</sup>, Shiqing Xu<sup>c</sup>

<sup>a</sup> Key Laboratory of High-Temperature and High-Pressure Study of the Earth's Interior, Institute of Geochemistry, Chinese Academy of Sciences, Guiyang 550002, China

<sup>b</sup> Key Laboratory of Orogenic Belts and Crustal Evolution, MOE; School of Earth and Space Sciences, Peking University, Beijing 100871, China

<sup>c</sup> College of Materials Science and Engineering, China Jiliang University, Hangzhou 310018, China

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## ABSTRACT

$\gamma\text{-Ca}_3(\text{PO}_4)_2\text{:Sm}^{3+}$  samples were synthesized under high-pressure and high-temperature conditions. The samples were characterized by X-ray diffraction measurements. The excitation and emission spectra and fluorescence decay curves of synthesized samples were collected to study the photoluminescence properties. The excitation spectra of  $\gamma\text{-Ca}_3(\text{PO}_4)_2\text{:Sm}^{3+}$  exhibit a series of narrow peaks attributed to the typical f–f transition of  $\text{Sm}^{3+}$ . The emission spectra of  $\gamma\text{-Ca}_3(\text{PO}_4)_2\text{:Sm}^{3+}$  compose of four bands attributed to transitions from  $^4\text{G}_{5/2}$  excited state to  $^6\text{H}_{J/2}$  ( $J = 5, 7, 9$  and  $11$ ) ground states of  $\text{Sm}^{3+}$ . The average decay lifetimes were obtained by fitting the decay curves for  $^6\text{H}_{7/2}$  level of  $\text{Sm}^{3+}$  emission using a double exponential function. The photoluminescent spectra of  $\gamma\text{-Ca}_3(\text{PO}_4)_2\text{:Sm}^{3+}$  show that the samples belong to a red-emitting phosphor and can be pumped in the near ultraviolet light region, which can easily be applied for NUV LED chips and thus in WLEDs. There exists a concentration quenching of  $\text{Sm}^{3+}$  in  $\gamma\text{-Ca}_3(\text{PO}_4)_2\text{:Sm}^{3+}$  samples. The critical concentration of  $\text{Sm}^{3+}$  in  $\gamma\text{-Ca}_3(\text{PO}_4)_2$  is about 1.0 mol%.

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## 1. Introduction

Phosphates were regarded as important hosts for phosphor and luminescent materials, and many phosphates doped with different lanthanides were investigated [1–16]. Among those phosphates,  $\beta$ -tricalcium phosphate ( $\beta\text{-TCP}$ ,  $\beta\text{-Ca}_3(\text{PO}_4)_2$ ) is one of the important calcium phosphates. Previous study shows that  $\beta\text{-TCP}$  is not stable under high-pressure and high-temperature conditions. At 4 GPa and 950 °C,  $\beta\text{-TCP}$  transforms into  $\gamma$ -tricalcium phosphate ( $\gamma\text{-TCP}$ ,  $\gamma\text{-Ca}_3(\text{PO}_4)_2$ ) [17]. There is a significant difference in the crystal structures between  $\beta\text{-TCP}$  and  $\gamma\text{-TCP}$ . In  $\beta\text{-TCP}$ , five kinds of Ca exist complicatedly with coordinated number of 3, 6, 7, 8 and 8, respectively [18]. However, only two kinds of Ca exist with coordinated number of 10 and 12 in  $\gamma\text{-TCP}$  [19]. Due to the crystal structure,  $\gamma\text{-TCP}$  is regarded as an important host for rare earth elements (REE) [19]. Actually, REE-bearing  $\gamma\text{-TCP}$  was obtained in high-pressure and high-temperature experiments [20].

A series of trivalent REE cations have photoluminescence and have been widely used in cathodoluminescence, display phosphor screens, lasers, and lamps due to their f–f or d–f transitions [21–24]. It is interesting to investigate the luminescent properties

of  $\gamma\text{-TCP}$  doped with REE. In our previous study,  $\text{Eu}^{3+}$ -doped  $\gamma\text{-TCP}$  was investigated [25]. As we known, trivalent samarium ( $\text{Sm}^{3+}$ ) is another most important luminescence center.  $\text{Sm}^{3+}$  has a fundamental level  $^6\text{H}_{5/2}$  and three main emitting levels  $^4\text{G}_{5/2}$ ,  $^4\text{F}_{3/2}$  and  $^4\text{G}_{7/2}$ , which gives visible reddish-orange emission and exhibits relatively high quantum efficiency forming a practical view point for lighting and displays [26,27].  $\text{Sm}^{3+}$ -doped  $\beta\text{-TCP}$  was recently prepared and the luminescent properties were characterized [28]. In this study,  $\text{Sm}^{3+}$ -doped  $\gamma\text{-TCP}$  samples were first synthesized in high-pressure and high-temperature experiments and the luminescence properties of the samples were investigated.

## 2. Experimental

### 2.1. Sample preparation

The  $\text{Sm}^{3+}$ -doped  $\gamma\text{-TCP}$  samples were synthesized under high-pressure and high-temperature conditions by using the mixture of  $\beta\text{-Ca}_3(\text{PO}_4)_2$  and  $\text{Sm}_2\text{O}_3$  (99.99%) as starting material. The experimental procedure is similar to that described in our previous study [25]. The  $\text{Sm}^{3+}$  doping concentrations were 0.25%, 0.5%, 1.0%, 1.5% and 2.0% (in molar ratio), respectively. The experiments were carried out using a 1000-tonne Kawai-type apparatus (USSA-1000) installed at Institute for Study of the Earth's Interior, Okayama

\* Corresponding author at: Institute of Geochemistry, Chinese Academy of Sciences, Guiyang, Guizhou 550002, China. Tel.: +86 851 5895692.

E-mail address: [xuweihong@vip.gyig.ac.cn](mailto:xuweihong@vip.gyig.ac.cn) (W. Xue).

University, Japan. An 18/11 sample assembly is adopted, as illustrated in Fig. 1. The thermal insulators were ZrO<sub>2</sub> tube and rods. The heater was graphite. Two platinum capsules enclosed different mixtures of Sm<sub>2</sub>O<sub>3</sub> (99.99%) and β-TCP were put in one experiment. Between the graphite heater and Pt capsules, MgO sleeve was inserted. The temperature was monitored by a thin W<sub>97</sub>Re<sub>3</sub>/W<sub>75</sub>Re<sub>25</sub> thermocouple whose junction was set at the center of the sample assembly. Quenched method was used to synthesize Sm<sup>3+</sup>-doped γ-TCP samples. The experimental conditions were selected as 8 GPa and 1000 °C for 10 h based on the sample assembly and the phase relationship of Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub> [29]. After pressure was applied, temperature was raised. The samples were kept at desired pressure and temperature conditions for 10 h, and then they were quenched by turning off the electric power supply. Finally the samples were decomposed to the ambient pressure.

## 2.2. Characterization

The synthesized samples were ground into fine powder and checked by powder X-ray diffractometer (Rigaku's Smartlab) using rotating anode Cu Kα irradiation. The acceleration voltage and the current were 40 kV and 30 mA, respectively. The XRD patterns were collected in the 2θ range from 25° to 80° with step size of 0.01°. The photoluminescence excitation and emission spectra and fluorescence decay curves of the synthesized samples were recorded with a Jobin-Yvon FL3-211-P spectrofluorometer, using a 370-nm pulsed spectral-LED and Xe lamp as an excitation source for the samples.

## 3. Results and discussion

### 3.1. Powder X-ray diffraction

γ-TCP crystallizes in a rhombohedral structure with the space group R-3 m (No. 166) [19,30]. Phosphorus atom is tetrahedrally coordinated by oxygen atoms. Calcium atoms are located at two crystallographic sites of Ca(1) and Ca(2) with coordination numbers of 12 and 10, respectively. Along the *c* axis, the structure can be expressed by a translationally interconnected polyhedral sequence PO<sub>4</sub>-Ca(2)O<sub>10</sub>-Ca(1)O<sub>12</sub>-Ca(2)O<sub>10</sub>-PO<sub>4</sub>. The powder XRD patterns of 0.25–2.0 mol% Sm<sup>3+</sup>-doped γ-TCP samples are shown in Fig. 2. The XRD patterns of synthesized β-TCP and γ-TCP are also illustrated. The XRD results indicate that Sm<sup>3+</sup>-doped γ-TCP samples show an identical structure with γ-TCP since all peaks can be indexed to the γ-Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub> phase according to PDF 32-176, and no obvious impurity phase was detected when Sm<sup>3+</sup> ions were doped into the host lattice. The unit cell parameters of synthesized samples were listed in Table 1. With increasing of Sm concentration, the lattice parameters slightly increase. It is well-known that the radius of Sm<sup>3+</sup> (0.96 Å) is similar to that of Ca<sup>2+</sup> (1.00 Å). Therefore, the substitution of Ca<sup>2+</sup> by Sm<sup>3+</sup> is easy in γ-TCP. A substituted mechanism involving a vacancy at Ca sites, i.e., 2Sm<sup>3+</sup> + □ = 3Ca<sup>2+</sup>, is responsible for the present study. Indeed, such substituted mechanism involving vacancy was confirmed in previous studies on other rare earth elements in apatites [31,32]. There are two kinds of Ca<sup>2+</sup> in γ-TCP, but we cannot determine the occupancy of Sm<sup>3+</sup> in γ-TCP. According to mass balance, CaO should appear in the sample. However, due to the relative low intensity of peaks and overlapping with those of γ-TCP, it is difficult to index peaks of CaO in XRD patterns.

### 3.2. Photoluminescence spectra

The luminescent properties of Sm<sup>3+</sup> (0.25–2.0 mol%) doped γ-TCP samples were investigated at room temperature. Fig. 3

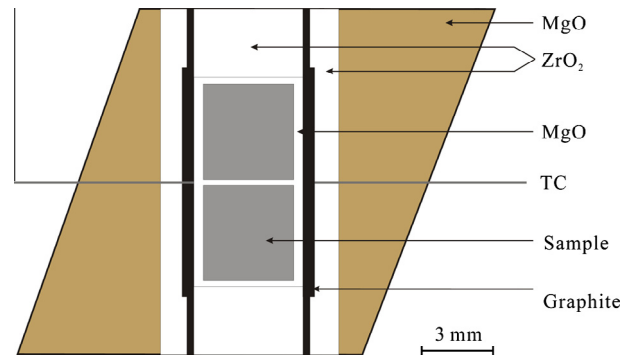


Fig. 1. Schematic drawing of the 18/11 sample assembly.

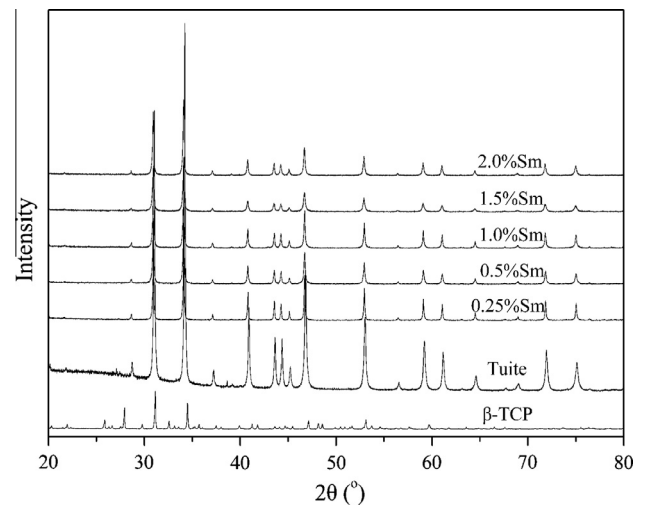


Fig. 2. X-ray diffraction patterns of β-TCP, γ-TCP and Sm<sup>3+</sup> (0.25–2.0 mol%) doped γ-Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub> samples.

Table 1  
Unit cell parameters, average life time (τ) and CIE of γ-Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub>:Sm<sup>3+</sup> samples.

Sm (mol%)	<i>a</i> (Å)	<i>c</i> (Å)	<i>V</i> (Å <sup>3</sup> )	τ (ms)	<i>x</i>	<i>y</i>
0	5.2524(4)	18.681(3)	446.30(5)	–	–	–
0.25	5.2531(3)	18.688(2)	446.60(6)	0.61	0.55	0.45
0.5	5.2536(5)	18.690(2)	446.74(8)	0.62	0.57	0.43
1.0	5.2547(4)	18.694(2)	447.02(7)	1.50	0.58	0.42
1.5	5.2557(4)	18.698(2)	447.29(7)	0.86	0.58	0.42
2.0	5.2570(5)	18.703(3)	447.63(2)	0.80	0.58	0.42

shows the excitation spectra ( $\lambda_{em} = 601$  nm) of Sm<sup>3+</sup>-doped γ-TCP samples with various concentrations. Sm<sup>3+</sup>-doped γ-TCP samples exhibit a series of narrow peaks attributed to the typical f–f transition of Sm<sup>3+</sup> ions. The results show the bands corresponding to transitions: <sup>6</sup>H<sub>5/2</sub> → <sup>4</sup>K, <sup>4</sup>L<sub>17/2</sub> (340 nm), <sup>6</sup>H<sub>5/2</sub> → <sup>4</sup>D, <sup>6</sup>P<sub>15/2</sub> (357 nm), <sup>6</sup>H<sub>5/2</sub> → <sup>4</sup>L<sub>17/2</sub> (370 nm), <sup>6</sup>H<sub>5/2</sub> → <sup>4</sup>K<sub>11/2</sub> (399 nm), <sup>6</sup>H<sub>5/2</sub> → <sup>6</sup>P<sub>5/2</sub> + <sup>4</sup>M<sub>19/2</sub> (413 nm), <sup>6</sup>H<sub>5/2</sub> → <sup>4</sup>G<sub>9/2</sub> + <sup>4</sup>I<sub>15/2</sub> (435 nm), and <sup>6</sup>H<sub>5/2</sub> → <sup>4</sup>F<sub>5/2</sub> + <sup>4</sup>I<sub>13/2</sub> (470 nm), respectively [13]. It is noted that the strongest excitation sharp line corresponding to the <sup>6</sup>H<sub>5/2</sub> → <sup>4</sup>K<sub>11/2</sub> transition of Sm<sup>3+</sup> located at 399 nm, which indicates that the samples can be pumped in the NUV light region, and can easily be applied for NUV LED chips and thus in WLEDs [33].

The emission spectra ( $\lambda_{exc} = 399$  nm) of Sm<sup>3+</sup>-doped γ-TCP samples were illustrated in Fig. 4. There are four main sharp emission peaks at 563, 597, 646 and 704 nm, among which the most intense peaks are centered at 597 nm. The emissions are caused by the f–f forbidden transitions of the 4f electrons of Sm<sup>3+</sup>, corresponding to

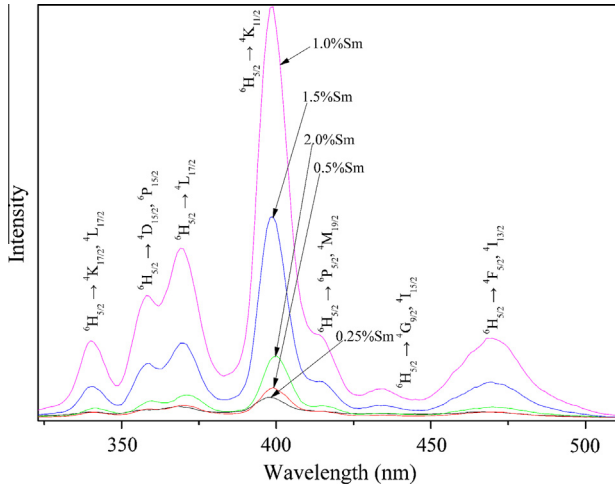


Fig. 3. The excitation spectra of Sm<sup>3+</sup> (0.25–2.0 mol%) doped  $\gamma$ -Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub> samples.

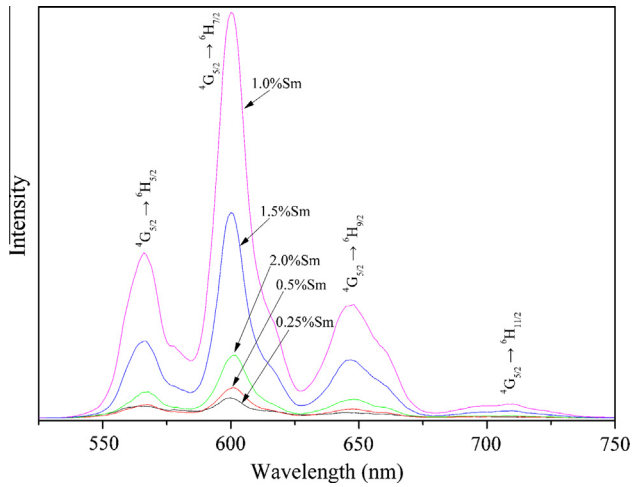


Fig. 4. The emission spectra of Sm<sup>3+</sup> (0.25–2.0 mol%) doped  $\gamma$ -Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub> samples.

${}^4G_{5/2} \rightarrow {}^6H_{5/2}$  (563 nm),  ${}^4G_{5/2} \rightarrow {}^6H_{7/2}$  (597 nm),  ${}^4G_{5/2} \rightarrow {}^6H_{9/2}$  (646 nm) and  ${}^4G_{5/2} \rightarrow {}^6H_{11/2}$  (704 nm), respectively [13].

It is obviously observed that the intensity of peaks in excitation and emission spectra change with the concentration of Sm<sup>3+</sup> while the peak locations remain practically unchanged. The excitation and emission intensity initially increases with the Sm<sup>3+</sup> concentration increasing from 0.25 to 1.0 mol%, and then decreases as the Sm<sup>3+</sup> concentration increases from 1.0 to 2.0 mol%, showing energy migration between Sm<sup>3+</sup> at different sites in the lattice. The excitation band with the highest intensity was the 399 nm one, corresponding to the  ${}^6H_{5/2} \rightarrow {}^4K_{11/2}$  transition for Sm<sup>3+</sup>. And the emission band with the highest intensity was the 597 nm one, corresponding to the  ${}^4G_{5/2} \rightarrow {}^6H_{7/2}$  transition for Sm<sup>3+</sup>. Fig. 5 shows the relative intensity of excitation and emission bands with different concentration of Sm<sup>3+</sup> ions. It should be pointed out that the intensity was normalized to highest intensity of each transition of Sm<sup>3+</sup> ions. The relative intensity of each transition of Sm<sup>3+</sup> ions shows similar trends for both excitation and emission bands. As the Sm<sup>3+</sup> concentration increases, the excitation and emission intensities of each transition gradually increase and reach a maximum at 1.0 mol%. The concentration quenching happens since the intensity decreases when the concentration increases from 1.0 to 2.0 mol%.

The causes of concentration quenching may be attributed to the facts that the activated Sm<sup>3+</sup> concentration is higher, the distance

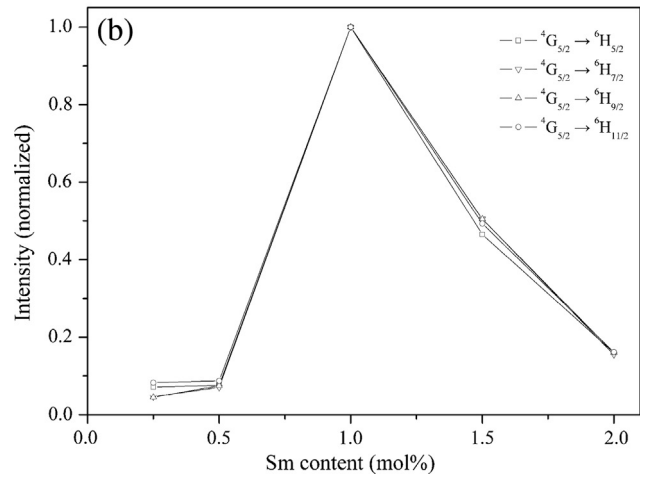
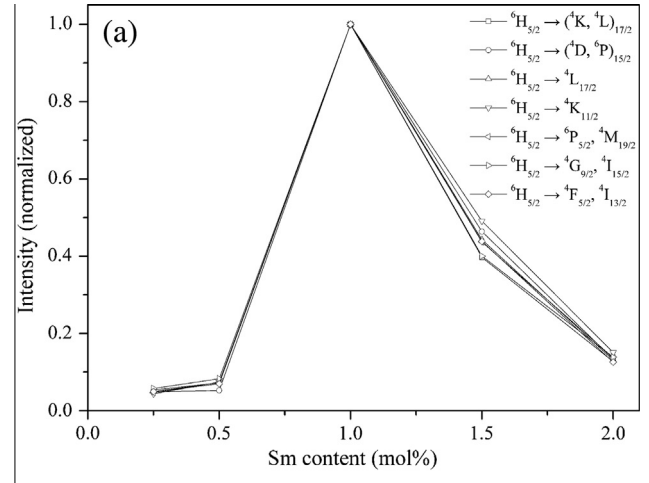


Fig. 5. Effect of Sm contents on the excitation (a) and emission (b) intensities of typical transitions of Sm<sup>3+</sup> ions in  $\gamma$ -Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub> host.

between the ions is shorter, ionic bonding interaction enhances and the non-radiative energy transfer Sm ( ${}^4G_{5/2}$ ) + Sm ( ${}^6H_{5/2}$ )  $\rightarrow$  Sm ( ${}^6F_{9/2}$ ) happens [22,24]. It is dependent on the critical distance  $R_c$  as the shortest average distance between the nearest activator Sm<sup>3+</sup> ions at a critical concentration  $x_c$ . Previous study pointed out that if the activator is introduced solely on Z ion sites, the critical transfer distance  $R_c$  is approximately equal to twice the radius of a sphere with this volume, expressed as the following equation [34]:

$$R_c \approx 2 \left( \frac{3V}{4\pi x_c N} \right)^{1/3} \quad (1)$$

where  $V$  is the unit cell volume,  $x_c$  is critical concentration, and  $N$  is the number of Z cations in the unit cell. By using the available values of  $V$ ,  $x_c$  and  $N$  (445.57 Å<sup>3</sup>, 0.01 and 6, respectively), the critical distance is calculated to be about 24.21 Å.

In order to obtain additional information on the luminescence properties of Sm<sup>3+</sup> ions in  $\gamma$ -Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub> host, the decay curves of the Sm<sup>3+</sup> emission at 601 nm corresponding to the  ${}^4G_{5/2} \rightarrow {}^6H_{7/2}$  emission line have been measured and given in Fig. 6. As shown in Fig. 6, the decay curves for  ${}^6H_{7/2}$  level of Sm<sup>3+</sup> emission can be well fitted into a double exponential function as follows:

$$I(t) = I_0 + A_1 \exp(-t/\tau_1) + A_2 \exp(-t/\tau_2) \quad (2)$$

where  $I$  and  $I_0$  are the luminescence intensity at time  $t$  and 0,  $A_1$  and  $A_2$  are constants,  $t$  is the time, and  $\tau_1$  and  $\tau_2$  are the decay times for

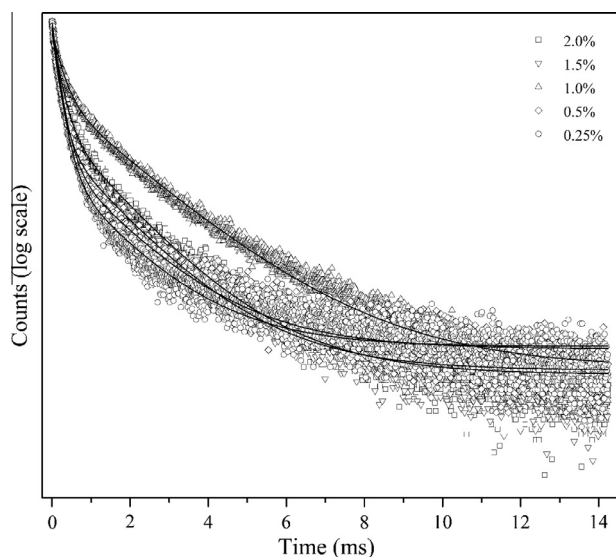


Fig. 6. Decay curves of the  $\text{Sm}^{3+}$  emission at 601 nm corresponding to the  $^4\text{G}_{5/2} \rightarrow ^6\text{H}_{7/2}$  emission line for  $\text{Sm}^{3+}$  (0.25–2.0 mol%) doped  $\gamma\text{-Ca}_3(\text{PO}_4)_2$  samples. The solid curves represent fittings by using a double exponential function.

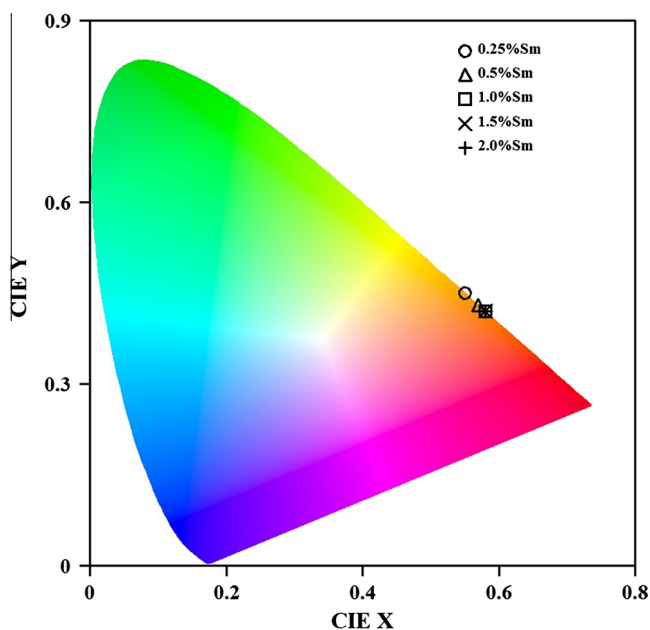


Fig. 7. CIE chromaticity coordinates of the  $\text{Sm}^{3+}$  (0.25–2.0 mol%) doped  $\gamma\text{-Ca}_3(\text{PO}_4)_2$  samples.

the exponential components, respectively. Further, the average decay lifetimes ( $\tau$ ) can be calculated as follows:

$$\tau = (A_1\tau_1^2 + A_2\tau_2^2)/(A_1\tau_1 + A_2\tau_2). \quad (3)$$

The analysis yields the average decay lifetime as 0.61, 0.62, 1.50, 0.86, and 0.80 ms for the Sm concentration of 0.25, 0.5, 1.0, 1.5, and 2.0 mol%, respectively, as listed in Table 1. Obviously, an effect of Sm concentration on the decay lifetime of the samples exists. The 1.0 mol% Sm doped sample is with the longest decay lifetime.

Color is usually communicated on the basis of the color coordinates established by the Commission Internationale de l'Éclairage (CIE) 1931, in a two-dimensional graphical representation of any color perceptible by the human eye on an  $x$ - $y$  plot. The obtained results were shown in Table 1 and Fig. 7. The CIE of 0.25 and

0.5 mol%  $\text{Sm}^{3+}$ -doped  $\gamma\text{-TCP}$  are about  $x=0.55$ ,  $y=0.45$ , and  $x=0.57$ ,  $y=0.43$ , as an open circle and open triangle in Fig. 7, and the CIEs of other three samples are merged due to the same  $x=0.58$  and  $y=0.42$ . The present results are slight different from that of  $\text{Sm}^{3+}$ -doped  $\beta\text{-Ca}_3(\text{PO}_4)_2$  (0.615,0.385) [28], but close in proximity to the Nicha corporation-developed amber LED NSPAR70BS (0.575,0.425) and locates in the red<sup>1</sup> region (0.57,0.41). The CIE chromaticity coordinates of present samples hardly change with the doped  $\text{Sm}^{3+}$  concentrations. The result is easily comprehensible because the location of the luminescent emission dose not shift and the dominant emission does not change, as shown in Fig. 4. The results are consistent with previous studies [28,33].

#### 4. Conclusions

Novel red phosphor  $\gamma\text{-Ca}_3(\text{PO}_4)_2:\text{Sm}^{3+}$  was synthesized at high-pressure and high-temperature conditions. The synthesized samples exhibited an intense emission at 597 nm. The relationship between the luminescent intensity and the concentration of activated ions  $\text{Sm}^{3+}$  shows that there is concentration quenching of  $\text{Sm}^{3+}$ -doped  $\gamma\text{-Ca}_3(\text{PO}_4)_2$ , and the critical concentration is 1.0 mol%. The CIE of  $\text{Sm}^{3+}$ -doped  $\gamma\text{-Ca}_3(\text{PO}_4)_2$  show that they are suitable to near-UV LEDs as the red phosphors.

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<sup>1</sup> For interpretation of color in Fig. 7, the reader is referred to the web version of this article.

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