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Photoluminescence properties of γ-$Ca_3(PO_4)_2$:Sm^{3+} prepared under high-pressure and high-temperature conditions

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ABSTRACT

γ-$Ca_3(PO_4)_2$: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 γ-$Ca_3(PO_4)_2$:Sm^{3+} exhibit a series of narrow peaks attributed to the typical f–f transition of Sm^{3+}. The emission spectra of γ-$Ca_3(PO_4)_2$:Sm^{3+} compose of four bands attributed to transitions from $^4G_{5/2}$ excited state to $^6H_{J/2}$ (J = 5, 7, 9 and 11) ground states of Sm^{3+}. The average decay lifetimes were obtained by fitting the decay curves for $^6H_{7/2}$ level of Sm^{3+} emission using a double exponential function. The photoluminescent spectra of γ-$Ca_3(PO_4)_2$: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 Sm^{3+} in γ-$Ca_3(PO_4)_2$:Sm^{3+} samples. The critical concentration of Sm^{3+} in γ-$Ca_3(PO_4)_2$ is about 1.0 mol%.

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, β-tricalcium phosphate (β-TCP, β-$Ca_3(PO_4)_2$) is one of the important calcium phosphates. Previous study shows that β-TCP is not stable under high-pressure and high-temperature conditions. At 4 GPa and 950 °C, β-TCP transforms into γ-tricalcium phosphate (γ-TCP, γ-$Ca_3(PO_4)_2$) [17]. There is a significant difference in the crystal structures between β-TCP and γ-TCP. In β-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 γ-TCP [19]. Due to the crystal structure, γ-TCP is regarded as an important host for rare earth elements (REE) [19]. Actually, REE-bearing γ-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 γ-TCP doped with REE. In our previous study, Eu^{3+}-doped γ-TCP was investigated [25]. As we known, trivalent samarium (Sm^{3+}) is another most important luminescence center. Sm^{3+} has a fundamental level $^6H_{5/2}$ and three main emitting levels $^4G_{5/2}$, $^4F_{3/2}$ and $^4G_{7/2}$, which gives visible reddish-orange emission and exhibits relatively high quantum efficiency forming a practical view point for lighting and displays [26,27]. Sm^{3+}-doped β-TCP was recently prepared and the luminescent properties were characterized [28]. In this study, Sm^{3+}-doped γ-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 Sm^{3+}-doped γ-TCP samples were synthesized under high-pressure and high-temperature conditions by using the mixture of β-$Ca_3(PO_4)_2$ and Sm_2O_3 (99.99%) as starting material. The experimental procedure is similar to that described in our previous study [25]. The 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

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University, Japan. An 18/11 sample assembly is adopted, as illustrated in Fig. 1. The thermal insulators were ZrO_2 tube and rods. The heater was graphite. Two platinum capsules enclosed different mixtures of Sm_2O_3 (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 $W_{97}Re_3$/$W_{75}Re_{25}$ thermocouple whose junction was set at the center of the sample assembly. Quenched method was used to synthesize Sm^{3+}-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 $Ca_3(PO_4)_2$ [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_4–$Ca(2)O_{10}$–$Ca(1)O_{12}$–$Ca(2)O_{10}$–PO_4. The powder XRD patterns of 0.25–2.0 mol% Sm^{3+}-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^{3+}-doped γ-TCP samples show an identical structure with γ-TCP since all peaks can be indexed to the γ-$Ca_3(PO_4)_2$ phase according to PDF 32-176, and no obvious impurity phase was detected when Sm^{3+} 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^{3+} (0.96 Å) is similar to that of Ca^{2+} (1.00 Å). Therefore, the substitution of Ca^{2+} by Sm^{3+} is easy in γ-TCP. A substituted mechanism involving a vacancy at Ca sites, i.e., $2Sm^{3+} + \square = 3Ca^{2+}$, 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^{2+} in γ-TCP, but we cannot determine the occupancy of Sm^{3+} 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^{3+} (0.25–2.0 mol%) doped γ-TCP samples were investigated at room temperature. Fig. 3 shows the excitation spectra (λ_{em} = 601 nm) of Sm^{3+}-doped γ-TCP samples with various concentrations. Sm^{3+}-doped γ-TCP samples exhibit a series of narrow peaks attributed to the typical f–f transition of Sm^{3+} ions. The results show the bands corresponding to transitions: $^6H_{5/2} \rightarrow (^4K, {}^4L)_{17/2}$ (340 nm), $^6H_{5/2} \rightarrow (^4D, {}^6P)_{15/2}$ (357 nm), $^6H_{5/2} \rightarrow {}^4L_{17/2}$ (370 nm), $^6H_{5/2} \rightarrow {}^4K_{11/2}$ (399 nm), $^6H_{5/2} \rightarrow {}^6P_{5/2} + {}^4M_{19/2}$ (413 nm), $^6H_{5/2} \rightarrow {}^4G_{9/2} + {}^4I_{15/2}$ (435 nm), and $^6H_{5/2} \rightarrow {}^4F_{5/2} + {}^4I_{13/2}$ (470 nm), respectively [13]. It is noted that the strongest excitation sharp line corresponding to the $^6H_{5/2} \rightarrow {}^4K_{11/2}$ transition of Sm^{3+} 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 (λ_{exc} = 399 nm) of Sm^{3+}-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^{3+}, corresponding to

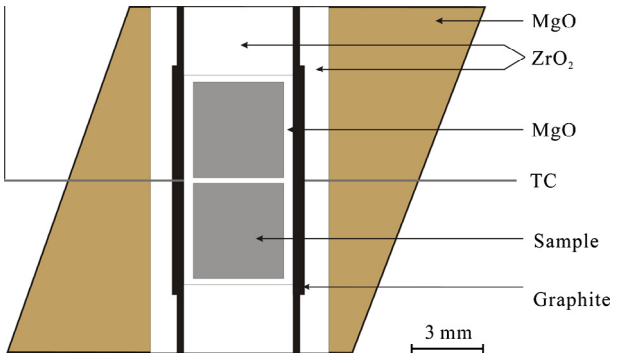

Fig. 1. Schematic drawing of the 18/11 sample assembly.

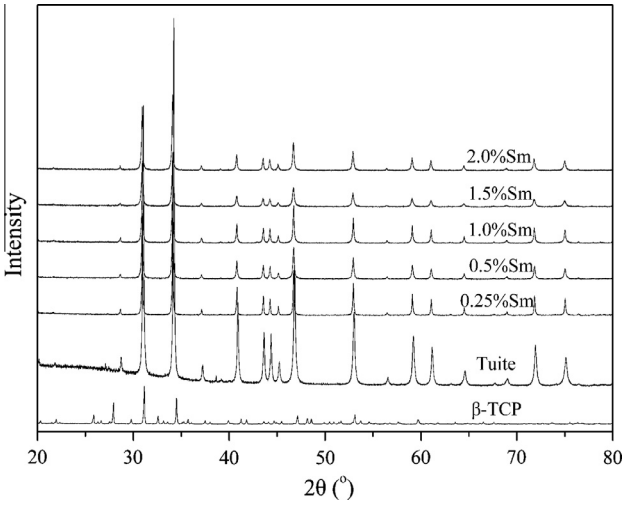

Fig. 2. X-ray diffraction patterns of β-TCP, γ-TCP and Sm^{3+} (0.25–2.0 mol%) doped γ-$Ca_3(PO_4)_2$ samples.

Table 1
Unit cell parameters, average life time (τ) and CIE of γ-$Ca_3(PO_4)_2$:Sm^{3+} samples.

Sm (mol%)	*a* (Å)	*c* (Å)	*V* (Å^3)	τ (ms)	*x*	*y*
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