Crystal structure evolution and luminescence properties of color tunable solid solution phosphors Ca_{2+x}La_{8-x}(SiO_4)_{6-x}(PO_4)_xO_2:Eu^{2+} 

Yuefei Xia, Jian Chen, Yan-gai Liu, Maxim S. Molokeev, Ming Guan, Zhaohui Huang and Minghao Fang

A series of apatite solid solution phosphors Ca_{2+x}La_{8-x}(SiO_4)_{6-x}(PO_4)_xO_2:Eu^{2+} (x = 0, 2, 4, 6) were synthesized by a conventional high-temperature solid-state reaction. The phase purity was examined using XRD, XPS and XRF. The crystal structure information, such as the concentration, cell parameters and occupation rate, was analyzed using a Rietveld refinement, demonstrating that the Eu^{2+} activated the Ca_{2+x}La_{8-x}(SiO_4)_{6-x}(PO_4)_xO_2 and Ca_{8-x}La_{2+}(PO_4)_xO_2 to form continuous solid solution phosphors. Different behaviors of luminescence evolution in response to structural variation were verified among the series of phosphors. Two kinds of Eu^{2+} ion sites were proved using low temperature PL spectra (8k) and room temperature decay curves. The substitution of large La^{3+} ions by small Ca^{2+} ions induced a decreased crystal field splitting of the Eu^{2+} ions, which caused an increase in emission energy from the 5d excited state to the 4f ground state and a resultant blue-shift from 508 nm to 460 nm. Therefore, with the crystal structure evolution, the emitted color of the series of phosphors could be tuned from green to blue by adjusting the ratio of Ca/La.

1. Introduction

Compared to conventional incandescent or fluorescent lamps, LED-based lighting devices fabricated by coating the yellow-emitting phosphor YAG:Ce^{3+} on blue InGaN LED chips have incited a revolution in the lamp industry due to their many advantages, such as significant power reduction, longer lifetime, higher luminous efficiency, environmental friendliness and brightness.\(^3\) However, this combination exhibits a poor color rendering index (CRI \(\approx 70\)–80) and a high correlated color temperature (CCT \(\approx 7750\) K) attributed to the insufficient red emission.\(^4\) \(^5\) One solution is to assemble LEDs by pumping a near-ultraviolet (n-UV) emitting chip with a mixture of blue, green and red emitting phosphors, which can then emit high quality white light with a smoother spectral distribution over the whole visible range.\(^6\) \(^7\)

The crystal field strength and coordination environment have great influences on the outermost electron transition of the Eu^{2+} ion, the most frequently used activator in phosphors, because the active electronic level is not shielded against the surrounding ligands,\(^8\) \(^9\) \(^10\) indicating that Eu^{2+} ions can emit light from the ultraviolet to the infrared with broadband emitting fluorescence.\(^11\) \(^12\) It is well known that apatite structure compounds (space group \(P6_3/m\)), with a general chemical formula of the form \(A_{10}(XO_4)_6Z_2\) (\(A = Ca^{2+},Ba^{2+},Ce^{3+},La^{3+},Y^{3+}\), etc., \(X = P^{5+},As^{5+},Si^{4+}\), etc., and \(Z = O^{2-},F^{-},Cl^{-},OH^{-}\), etc.),\(^13\) contain two kinds of cation sites: the 9-fold coordinated 4f sites with \(C_3\) point symmetry and the 7-fold coordinated 6 h sites with a \(C_6\) point symmetry, which are suitable for the substitution of various rare-earth-metal ions.\(^14\) \(^15\) \(^16\) Consequently, due to their adjustable structures, and excellent thermal and physicochemical stabilities, apatite compounds have become highly efficient host materials for the luminescence of various rare earth ions and have aroused widespread attention.\(^17\) \(^18\) \(^19\) As discussed above, the coordination environment of the Eu^{2+} site is anticipated to be changed via chemical composition variation among solid solution phosphors.\(^20\) Thus, the emitted color can be controlled by doping Eu^{2+} ions into a series of apatite solid solution hosts which are expected to display adjustable emission spectra in a wide range to meet the requirements of multi-color phosphors.

In this study, the coordination environment variation of the Eu^{2+} ion has been realized by the replacement of Ca^{2+} ions with La^{3+} ions and resulted in crystal splitting decreases of the Eu^{2+} ion, a series of color tunable solid solution phosphors...
Ca$_{2+}$La$_{x}$(SiO$_4$)$_6$(PO$_4$)$_4$O$_2$:Eu$^{2+}$ ($x = 0,2,4,6$) were successfully prepared by a high-temperature solid-state reaction. Moreover, it’s worth noting that the replacement of [PO$_4$]$^{3-}$ with the [SiO$_4$]$^{4-}$ tetrahedron was introduced into the solid solution to realize the charge compensation because of the different valences between Ca$^{2+}$–La$^{3+}$ and also that the Ca$_{4}$La$_{6}$SiO$_{4}$O$_2$:Eu$^{2+}$ and Ca$_{4}$La$_{6}$SiO$_{4}$O$_2$:Eu$^{2+}$ phosphors were synthesized for the first time.

The phase purity was demonstrated by XRD, XPS and XRF, and the crystal structure information was analyzed base on Rietveld refinement results. In addition, the relationship between the crystal structure evolutions, the PLE and PL spectra at normal and low temperatures, the lifetimes and the temperature dependence spectra have been discussed in detail.

2. Experimental

2.1. Materials and synthesis

A series of apatite solid solution phosphors Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_4$O$_2$:0.02Eu$^{2+}$ ($x = 0,2,4,6$) were synthesized according to a conventional high-temperature solid-state method. CaCO$_3$ (analytical reagent (A. R.)), La$_2$O$_3$ (A. R.), SiO$_2$ (A. R.), NH$_4$H$_2$PO$_4$ (A. R.), and Eu$_2$O$_3$ (A. R.) were used as the raw materials. After the raw materials were well ground in an agate mortar for 10 min according to the stoichiometric amounts of reactants, all the materials were preheated for 1 h at 1000°C in a corundum crucible to eliminate H$_2$O and CO$_2$. Then, the mixed powders were sintered at 1525°C in a reductive atmosphere (H$_2$ 10%, N$_2$ 90%) for 6 h in a horizontal tube furnace. Finally, the sintered products were well ground after cooling down to room temperature.

2.2. Characterization

The X-ray diffraction (XRD) data of the powder samples were collected on an X-ray powder diffractometer (D/max-IIIA, Rigaku, Japan) with a step-wise scanning mode over the $2\theta$ range of 10°–110° using Cu K$_\alpha$ radiation (1.5406 Å) under an operating voltage (40 kV) and current (100 mA). The XRD patterns, which were submitted for Rietveld refinement, were acquired at a step size of 0.02° with a counting time of 2 s per step.

X-ray photoelectron spectroscopy (XPS) measurements were collected using a Kratos Axis Ultra DLD, employing an MCP stack & delay-line photoelectron detector with scanned & snapshot spectroscopy modes. X-ray Fluorescence (XRF) measurements were measured by utilizing a Rigaku ZSX Primus II X-ray fluorescence spectrophotograph. The photoluminescence emission (PL) and the photoluminescence excitation (PLE) spectra at 298k and 8k were measured using a Hitachi F-4600 fluorescence spectrophotometer (Japan) equipped with a 150 W Xe lamp as the excitation source. The temperature-dependent luminescence properties were measured on the same spectrophotometer which was assembled with a computer-controlled electric furnace and a self-made heating attachment. The morphology was observed using high-resolution transmission electron microscopy (HRTEM; JEM-21000, JEOL, Japan). The room-temperature luminescence decay curves were obtained from a spectrofluorimeter (Horiba, Jobin Yvon TBXPS) using a tunable pulse laser radiation (nano-LED) for the excitation.

3. Results and discussion

3.1. Phase formation and structural characteristics

The XRD patterns of Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_4$O$_2$:0.02Eu$^{2+}$ ($x = 0,2,4,6$) and the standard PDF card (JCPDF 29-0337) of Ca$_3$La$_8$(SiO$_4$)$_6$O$_2$ are displayed in Fig. 1a. All the diffraction peaks of these samples matched well with the JCPDF 29-0337 card, indicating that the series of solid solution phosphors had been prepared successfully and that the introduction of the Eu$^{2+}$ ion did not cause any impurity phase. In addition, the diffraction peaks of the last three samples ($x = 2, 4, 6$) have obviously shifted towards the high degrees with the increase of the ratio.

![Fig. 1](image-url)
of Ca/La. This can be explained as the substitution of the large La<sup>3+</sup> ion by the small Ca<sup>2+</sup> ion and the large [SiO<sub>4</sub>]<sup>−</sup> tetrahedron by the small [PO<sub>4</sub>]<sup>3−</sup> tetrahedron causes a decrease in the lattice constant of the Ca<sub>2</sub>xLa<sub>8−x</sub>[SiO<sub>4</sub>]<sub>6</sub>[PO<sub>4</sub>]<sub>6</sub>O<sub>2</sub> hosts, which is also evidence demonstrating the existence of a solid solution in the whole range (x = 0,2,4,6). The photoelectron survey spectrum of the Ca<sub>2</sub>xLa<sub>8−x</sub>[SiO<sub>4</sub>]<sub>6</sub>[PO<sub>4</sub>]<sub>6</sub>O<sub>2</sub>:0.02Eu<sup>2+</sup> sample is plotted in Fig. 1b, depicting photoelectron peaks corresponding to O 1s, Si 2p, P 2p, La 4p<sub>3/2</sub>, Ca 2p, La 2s, O 1s, La 3d<sub>5/2</sub> and Eu 3d<sub>5/2</sub> and 3d<sub>3/2</sub> emission. The inset shows the high-resolution XPS spectrum at the Eu 3d<sub>5/2</sub> position, and the peak at 1126.28 ev agrees well with the signal of Eu<sup>2+</sup> 3d<sub>5/2</sub>, demonstrating the existence of the Eu<sup>2+</sup> ion. Furthermore, the elemental content of the Ca<sub>2</sub>xLa<sub>8−x</sub>[SiO<sub>4</sub>]<sub>6</sub>[PO<sub>4</sub>]<sub>6</sub>O<sub>2</sub>:0.02Eu<sup>2+</sup> sample was further examined and calculated by XRF. As shown in Table 1, the proportions of all components match well with the suggested formula.

Fig. 2a displays the crystal structure of the 2 × 2 × 2 unit cells of Ca<sub>2</sub>xLa<sub>8−x</sub>[SiO<sub>4</sub>]<sub>6</sub>[PO<sub>4</sub>]<sub>6</sub>O<sub>2</sub>, which is chosen as the representative. Obviously, Ca<sub>2</sub>xLa<sub>8−x</sub>[SiO<sub>4</sub>]<sub>6</sub>[PO<sub>4</sub>]<sub>6</sub>O<sub>2</sub> has a layered structure and contains two kinds of cation sites: the inner-cation site of Ca<sup>2+</sup> or La<sup>3+</sup> and the inter-laminar site labeled M(i) with the local symmetry C<sub>3</sub> and the inter-laminar site labeled M(ii) with the local symmetry C<sub>6</sub>. The two kinds of different coordination environment of the cation sites are displayed in Fig. 2b and c. The M(i) site at the 4f site (C<sub>4</sub>) was surrounded by 9 oxygen atoms to form a monocapped square antiprism, which was connected with tetrahedral PO<sub>4</sub>/SiO<sub>4</sub> groups, and the M(ii) site at the 6 h site (C<sub>6</sub>) formed a pentagonal bipyramid with the surrounding 7 oxygen atoms and these bipyramids were connected with each other through a vertex. Theoretically, both the Ca<sup>2+</sup> ion and the La<sup>3+</sup> ion are uniformly distributed in the two kinds of cationic sites, illustrating that the ratio of Ca/La is easily changed by adjusting the proportion of raw materials.

The four XRD patterns were analyzed using the Rietveld refinement. The observed (x), calculated (red) and difference (gray) XRD profiles for the refinements of Ca<sub>2</sub>xLa<sub>8−x</sub>[SiO<sub>4</sub>]<sub>6</sub>[PO<sub>4</sub>]<sub>6</sub>O<sub>2</sub>:0.02Eu<sup>2+</sup> (x = 0,2,4,6) are shown in Fig. 3a-d. The main refinement parameters of the processing and refinement results are presented in Table 2. The results of the refinement further demonstrate that the series of solid solution phosphors are single phase without any impurity or secondary phases. On the other hand, the ranges of the weighted profile R-factor (Rwp) and the R-Bragg factor (RBr) are 10.038%–11.012% and 1.792%–2.777%, respectively, indicating that the crystal structures of these phosphors match well with the starting model (Ca<sub>2.87(2)</sub>La<sub>7.13(2)</sub>[SiO<sub>4</sub>]<sub>6</sub>O<sub>2</sub>) after the refinement and that the results are believable and publishable. Meanwhile, we also refined the concentration of these samples while obtaining the high-quality XRD data. According to these refinement results, the calculated formula are Ca<sub>2</sub,xLa<sub>8−x</sub>[SiO<sub>4</sub>]<sub>6</sub>[PO<sub>4</sub>]<sub>6</sub>O<sub>2</sub>, Ca<sub>4</sub>,47(2)[La<sub>5.53(2)</sub>[SiO<sub>4</sub>]<sub>3.53(2)</sub>[PO<sub>4</sub>]<sub>3</sub>O<sub>2</sub>, Ca<sub>5.52(2)</sub>[La<sub>1.48(2)</sub>[SiO<sub>4</sub>]<sub>1.48(2)</sub>[PO<sub>4</sub>]<sub>4.52(2)</sub>O<sub>2</sub> and Ca<sub>8.28(2)[La<sub>1.72(2)</sub>[SiO<sub>4</sub>]<sub>1.72(2)</sub>[PO<sub>4</sub>]<sub>6</sub>O<sub>2</sub>, demonstrating that the refined concentrations are highly consistent with the suggested formula and that the coordination environment variation is mainly caused by the replacement of a Ca<sup>2+</sup> with a La<sup>3+</sup> ion. Moreover, according to the XRD data and the Rietveld refinement, no additive superstructure peaks were detected, this feature proves that the Ca<sup>2+</sup> ions and the La<sup>3+</sup> ions were randomly mixed at the atomic level. In addition, the unit lattice parameters and the unit cell volumes of the as-prepared phosphors are given in Fig. 4. The high linear fitting coefficients (0.9934–0.99796) proved the crystal structure evolution of this continuous solid solution. The lattice parameters and unit cell volumes show linear decreases and are proportional to the value of x, which is attributed to the substitution of the large La<sup>3+</sup> ions by small Ca<sup>2+</sup> ions, suggesting that the coordination environment of the cations become more unconsolidated as x increases.

As shown in Fig. 5a-d, the linear change in the crystal structure of these solid solution phosphors was further verified by HRTEM and fast Fourier transform (FFT) images. Both the HRTEM and FFT images illustrate that no significant structural defects appeared in the selected areas of these single-phase samples and that good crystallinity was obtained. Moreover, the lattice fringe measurements with the d spacings of 0.315 nm, 0.837 nm, 0.354 nm, and 0.353 nm could be assigned to the planes (210), (010), (021) and (021) for

**Table 1.** The main parameters of the XRF measurements

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**Fig. 2.** (a) The crystal structure of the 2 × 2 × 2 unit cells of Ca<sub>2</sub>xLa<sub>8−x</sub>[SiO<sub>4</sub>]<sub>6</sub>[PO<sub>4</sub>]<sub>6</sub>O<sub>2</sub>:0.02Eu<sup>2+</sup>; (b) the coordination environment of the M(i) site with the local symmetry C<sub>3</sub> and (c) the M(ii) site with the local symmetry C<sub>6</sub>.
Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_{6-x}$(PO$_4$)$_x$O$_2$:0.02Eu$^{2+}$ ($x = 0, 2, 4, 6$) samples with the corresponding Rietveld refinement (red) and residuals (gray).

Table 2 The main refinement parameters of Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_{6-x}$(PO$_4$)$_x$O$_2$ ($x = 0, 2, 4, 6$) crystal structure. Therefore, four kinds of emitting blocks will be included in the host: Ca(I), Ca(II), La(I), and La(II). The occupation rates of Ca(I), Ca(II), La(I), and La(II), reflecting the distribution of Ca$^{2+}$ ions and La$^{3+}$ ions in the two cationic sites, have been refined and displayed in Table 3. When the content of Ca$^{2+}$ ions increases, the occupation rate of La(II) is decreased faster than that of La(I) and the M(II) site is totally occupied by Ca$^{2+}$ ions while the chemical formula is Ca$_8$La$_2$(PO$_4$)$_6$O$_2$ ($x = 6$), which shows similar results to the previous studies. The phenomenon can be explained as follows: the Eu$^{2+}$ ion located at the M(II) site that is connected with a free oxygen ion, results in a very short Eu–O distance due to the small sum of the electrostatic bond strength of the cations toward the free oxygen ion. Thus, it must be very unfavorable for the (6 h) sites to be occupied by a cation with a large radius in these compounds.

3.2. Photoluminescence characteristics

The PLE spectra (monitored in the range of 460 nm–508 nm) and the PL spectra (under 365 nm excitation) of the Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_{6-x}$(PO$_4$)$_x$O$_2$:0.02Eu$^{2+}$ ($x = 0, 2, 4, 6$) phosphors at room temperature are displayed in Fig. 6a and b, respectively. The PLE spectra depict a series of broad excitation bands between 250 nm and 450 nm, which are attributed to the 4f$^7$ → 4f$^6$5d$^1$ transition of the Eu$^{2+}$ ion, and the emission spectra show a series of broad emission bands from 400 nm to 650 nm, which are attributed to the 5d$^1$ → 4f$^7$ transition of the Eu$^{2+}$ ion. As shown in Fig. 6b, a strong asymmetric broad band with a maximum wavelength peak at 508 nm was detected in the PL spectrum of Ca$_2$La$_8$(PO$_4$)$_6$O$_2$:0.02Eu$^{2+}$, which was
The refinement of Ca2La8(PO4)6O2:0.02Eu2+ (x = 0, 2, 4, 6) shows that the emission band can be decomposed into two Gaussian components with peaks at 526 and 576 nm, and the PL spectrum of Ca2La8(PO4)6O2:0.02Eu2+ can be separated into two components with peaks at 526 nm and 576 nm, and the PL spectrum of Ca2La8(PO4)6O2:0.02Eu2+ obviously contains two components with peaks at 460 nm and 625 nm. The results further demonstrate that there are two kinds of cation sites included in the series of apatite phosphors which could be occupied by the Eu2+ ion.

Actually, considering the valence state and the previous results,16,29 the blue emission band observed in the PL spectrum of Ca2La8(PO4)6O2:0.02Eu2+ is attributed to the substitution of a Ca2+ ion by a Eu2+ ion. According to the report by Van Uitert, to further demonstrate the relationship between emission peaks and emission centers, the possible crystallographic site may be investigated theoretically by the following equation:30

$$E \left( \text{cm}^{-1} \right) = Q^* \left[ 1 - \left( \frac{V}{4} \right)^{1/4} \times 10^{-\left(6eVr/60\right)} \right],$$  

where $E$ represents the energy position of the d-band edge for the rare-earth ion (cm$^{-1}$), $Q^*$ is the energy position for the lower d-band edge for the free ion (34 000 cm$^{-1}$ for Eu$^{2+}$), $V$ is the valence of the activator (Eu$^{2+}$) ion ($V = 2$), $n$ is the number of anions in the immediate shell around the Eu$^{2+}$ ion, $r$ is the radius of the host cation replaced by the Eu$^{2+}$ ion (in Å), and $E_a$ is the electron affinity of the atoms that form anions (in eV) depending on the anion complex type. Here, $E_a$ was approximately determined as 1.60 for the [SiO$_4$]$^{4-}$ in the oxide host$^{31}$ and the $r$ values of Ca$i$ and Ca$ii$ were calculated to be 118 pm and 106 pm, respectively. Consequently, the $E_{calcd}$ values of Ca$i$ and Ca$ii$ were calculated to be 19 693 cm$^{-1}$ and 18 546 cm$^{-1}$, respectively. The results indicated that the emission band centered at 504 nm was attributed to a Eu$^{2+}$ ion occupying the Ca(i) site with $C_1$ symmetry and nine-coordination and that the emission band peaked at 542 nm was attributed to a Eu$^{2+}$ ion occupying the Ca(ii) site with $C_3$ symmetry and seven-coordination.

The emission wavelengths of the Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_2$O$_2$:0.02Eu$^{2+}$ ($x = 0, 2, 4, 6$) PL spectra show a wide blue shift from 508 nm to 460 nm with the increase of the $x$ value which is dependent on the crystal field strength variation. The structural model of the cation substitution around the Eu$^{2+}$ ion sites is depicted in Fig. 7. In this regard, the crystal field splitting of the Eu$^{2+}$ ions can be determined as obeying,$^{17,32,33}$

$$D_n = \frac{Ze^2r^4}{6R^6},$$  

where $D_n$ is a measure of energy level separation, $R$ represents the distance from the central ion to its ligands, $z$ stands for the charge or valence of the anion, $r$ is the radius of the d wave

**Table 3** Atomic occupation rates in Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_2$O$_2$ with different $x$ values

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**Fig. 4** The refined unit cell parameters (a, c) and cell volume (v) of Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_2$:0.02Eu$^{2+}$ ($x = 0, 2, 4, 6$).

**Fig. 5** HRTEM and FFT images for Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_2$:0.02Eu$^{2+}$ with different $x$ values, $x = 0$ (a), $x = 2$ (b), $x = 4$ (c) and $x = 6$ (d).
function, and $e$ is the charge of an electron. For the $d_{(5u-)}$ orbital, if $z$, $e$, and $r$ are equal, then $D_q$ is only a function of $1/R^5$. When the large La$^{3+}$ ion is substituted by the small Ca$^{2+}$ ion, the distance between Eu$^{2+}$ and O$^{2+}$ becomes longer and the magnitude of the crystal field strength decreases.\textsuperscript{34,35} Thus, the crystal field splitting of the Eu$^{2+}$ ion is reasonably decreased and this results in a gradual increase of the lowest 5d state.\textsuperscript{36} As a consequence, the emission wavelength shows a blue shift among the as-prepared phosphors and a color variation from green to blue.

In addition, as shown in Table 4, the stokes shifts were estimated to be 6135 cm$^{-1}$–5985 cm$^{-1}$ and the full width at half-maximum (FWHMs) of the PLE spectra decrease from 142 nm to 113 nm with the increase in $x$. The crystal-field splitting of the Eu$^{2+}$ ions was estimated to be 22 230 cm$^{-1}$–18 190 cm$^{-1}$, which is calculated by the gap between the first and the last component peaks of the PLE spectra. All the computations further demonstrate the decrease of the crystal field splitting.\textsuperscript{37}

Furthermore, Fig. 8 presents the room temperature decay curves of the Eu$^{2+}$ ion luminescence in the Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_x$O$_2$:0.02Eu$^{2+}$ ($x = 0, 2, 4, 6$) series. All of the decay curves can be well fitted with a second order exponential equation:

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

where $I(t)$ is the luminescence intensity, $A_1$ and $A_2$ represent constants, $\tau$ is the time, and $\tau_1$ and $\tau_2$ stand for rapid and

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Fig. 6 (a) The PLE monitored in the range of 460 nm–508 nm and (b) the PL spectra under 365 nm excitation of Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_x$O$_2$:0.02Eu$^{2+}$ ($x = 2, 4, 6$) at room temperature. (c) The PL and PLE spectra at low temperature (8k) of Ca$_2$La$_4$(SiO$_4$)$_6$O$_2$:0.02Eu$^{2+}$ and (d) the PL and PLE spectra at low temperature (8k) of Ca$_5$La$_6$(PO$_4$)$_x$O$_2$:0.02Eu$^{2+}$.

Fig. 7 Structural model for explaining the photoluminescence evolution observed in the Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_x$O$_2$:0.02Eu$^{2+}$ ($x = 0, 2, 4, 6$) phosphors: (a–d) represent Eu$^{2+}$ located in the M(I) site, (e–h) represent Eu$^{2+}$ located in the M(II) site.

Table 4 Excitation and emission bands, stokes shift, and the crystal field splitting of the Ca$_{2+x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_x$O$_2$:0.02Eu$^{2+}$ ($x = 0, 2, 4, 6$) phosphors.
slow lifetimes for exponential components, respectively. Moreover, the effective lifetime constant ($\tau^*$) can be calculated as:

$$\tau^*/C_3 = (A_1\tau_1^2 + A_2\tau_2^2) / (A_1\tau_1 + A_2\tau_2) \quad (4)$$

The effective decay times were calculated to be 745.34, 468.61, 384.15 and 353.33 ns with $x = 0$, 2, 4 and 6, respectively. One can see that the decay times decrease with the increasing Ca$^{2+}$ ion content. Due to the phase structures becoming more unconsolidated compared to the original Ca$_2$La$_8$(SiO$_4$)$_6$(PO$_4$)$_x$O$_{2-x}$:0.02Eu$^{2+}$ phase, the increasing possibility of energy transfer among the Eu$^{2+}$ ions increases the possible non-radiative transition and leads to the decreasing lifetime values.$^{32,38}$ Also the two decay components ($\tau_1$ and $\tau_2$) proved that the Eu$^{2+}$ occupied two different Ca$^{2+}$ sites.$^{13}$

The temperature dependence experiment proved that the thermal stability is consistent with the crystal structure evolution in the as-prepared series phosphors, and the peak emission intensity that normalized to 25 °C values was depicted in Fig. 9. As shown in Fig. 9, the emission intensity of all the samples decreases with the increase in temperature. Additionally, the thermal stability gradually decreases with the increasing $x$ values, this phenomenon can be explained by the neighboring-cation effect.$^{39,40}$ The replacement of large La$^{3+}$ ions by small Ca$^{2+}$ ions makes the distances between the Eu$^{2+}$ activator ion and the neighboring cations become smaller, as demonstrated by the refined cell parameters and cell volume, resulting in a larger coulombic force following the inverse-square law$^{41,42}$ and the decrease of the thermal quenching barrier height. Thus, the thermal stability becomes lower when the replacement occurs.

Luminescence efficiency is an important technological parameter for the application of phosphors. The internal quantum efficiency (QE) of the Ca$_{2-x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_x$O$_{2-x}$:0.02Eu$^{2+}$ ($x = 0,2,4,6$) phosphors was measured and calculated following:$^{12}$

$$\eta_{QE} = \frac{\int L_S}{\int E_R + \int E_S} \quad (5)$$

where $L_S$ is the luminescence emission spectrum of the sample, $E_R$ represents the spectrum of the excitation light from the empty integrated sphere (without the sample), and $E_S$ stands for the excitation spectrum for exciting the sample. As a result, under 365 nm excitation, the internal QE of the Ca$_{2-x}$La$_{8-x}$(SiO$_4$)$_6$(PO$_4$)$_x$O$_{2-x}$:0.02Eu$^{2+}$ ($x = 0,2,4,6$) phosphors was estimated to be about 45.44%, 46.39%, 44.12% and
46.56% for x = 0, 2, 4, and 6, respectively. The QE of the as-prepared phosphors can be further optimized by improving the preparation conditions because the QE depends closely on the prepared conditions, crystalline defects, particle sizes and the morphology of the phosphor.25,39

The CIE coordinates and the digital photos of the Ca$_2$$_x$La$_{8-x}$Si$_6$O$_{20}$:Eu$^{2+}$ ($x$ = 0, 2, 4, 6) phosphors under 365 nm excitation are displayed in Fig. 10. The calculated CIE coordinates are (0.2099, 0.4884) for $x$ = 0, (0.2167, 0.3585) for $x$ = 2, (0.1955, 0.2992) for $x$ = 4 and (0.1846, 0.1851) for $x$ = 6. Obviously, both the digital photos and the coordinates demonstrate that the emitted color of the solid solution phosphors Ca$_2$$_x$La$_{8-x}$Si$_6$O$_{20}$:Eu$^{2+}$ ($x$ = 0, 2, 4, 6) can be adjusted in the wide range from green to blue by changing the ratio of Ca/La.

4. Conclusions

In summary, a series of continuous solid solution phosphors Ca$_2$$_x$La$_{8-x}$Si$_6$O$_{20}$:Eu$^{2+}$ ($x$ = 0, 2, 4, 6) were successfully synthesized by a solid-state reaction method. The phase purity was examined by XRD, XPS and XRF, the crystal structure information was identified by both Rietveld refinement and HRTFM. The refined concentration, cell parameters and occupation rate matched well with the suggested formula and the variation tendency of Ca/La demonstrating that Eu$^{2+}$ activated Ca$_2$La$_{8-x}$Si$_6$O$_{20}$ and Ca$_2$La$_{8-x}$PO$_{20}$:Eu$^{2+}$ ($x$ = 0, 2, 4, 6) to form continuous solid solution phosphors. Different spectroscopic behavior properties based on structural variations were verified among the series of phosphors. The PL spectra at room temperature and low temperature (8K) were well separated into two component peaks, demonstrating that there are two kinds of Eu$^{2+}$ ion sites included in these apatite phosphors. Furthermore, the short wavelength emission peak is from the Eu$^{2+}$ ion located at the 4f site with nine coordination and the long wavelength emission peak is ascribed to the Eu$^{2+}$ ion occupying the 6h site with seven coordination. Under 365 nm excitation, the series of phosphors show a strong blue shift from 508 nm to 460 nm. It can be explained by the fact that the substitution of the large La$^{3+}$ ions by small Ca$^{2+}$ ions induced a decreased crystal field splitting of the Eu$^{2+}$ ions, which led to the emission energy from the 5d excited state to the 4f ground state increasing. Consequently, the emitted color of the series of phosphors could be tuned from green to blue by adjusting the ratio of Ca/La.

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References