Synthesis and photoluminescence characteristics of high color purity \( \text{Ba}_3\text{Y}_4\text{O}_9: \text{Eu}^{3+} \) red-emitting phosphors with excellent thermal stability for warm W-LED application

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Single phase \( \text{Eu}^{3+} \)-activated \( \text{Ba}_3\text{Y}_4\text{O}_9 \) (\( \text{Ba}_3(\text{Y}_{1-x}\text{Eu}_x)_4\text{O}_9 \)) red-emitting phosphors with different \( \text{Eu}^{3+} \) doping concentrations were synthesized by a high temperature solid-state reaction method. The phase purity, crystal structure, photoluminescence properties, internal quantum efficiency, decay lifetimes, and thermal stability were investigated. Upon excitation at 396 nm near-ultraviolet light and 469 nm blue light, the \( \text{Ba}_3(\text{Y}_{1-x}\text{Eu}_x)_4\text{O}_9 \) phosphors exhibited a strong red emission at 614 nm due to the \( ^5\text{D}_0 \rightarrow ^7\text{F}_2 \) transition of \( \text{Eu}^{3+} \) ions. The optimal doping concentration of \( \text{Eu}^{3+} \) ions in \( \text{Ba}_3(\text{Y}_{1-x}\text{Eu}_x)_4\text{O}_9 \) was found to be \( x = 0.25 \). Furthermore, the critical distance was calculated to be 12.78 A and the energy transfer mechanism for the concentration quenching effect was determined to be quadrupole–quadrupole interaction. In addition, the Commission Internationale de l’Eclairage (CIE) chromaticity coordinates of \( \text{Ba}_3(\text{Y}_{0.75}\text{Eu}_{0.25})_4\text{O}_9 \) phosphors were measured to be (0.6695, 0.3302) which located at the red region, and significantly, the high color purity was about 97.9%. The as-synthesized phosphors also possessed excellent thermal stability and the activation energy was determined to be 0.29 eV. Therefore, the investigated results indicated that the \( \text{Ba}_3\text{Y}_4\text{O}_9: \text{Eu}^{3+} \) phosphor may be a suitable candidate as a red phosphor for white light-emitting diodes under effective excitation at near-ultraviolet and blue light.

1. Introduction

During the past decades, frequency conversion using luminescent materials, which can convert incident radiation into desired wavelengths, has been widely applied in various fields, such as lighting, displays, bioimaging, sensors, and solar cells.\(^1\)–\(^6\) In recent years, white light-emitting diodes (W-LEDs) have gradually replaced traditional incandescent and fluorescent lamps owing to their obvious advantages, such as high efficiency, low energy cost, environmental friendliness, and long lifetime.\(^7\)–\(^15\) They are widely applied in residential lighting, automotive lighting, displays, and so on.\(^16\)–\(^17\) For indoor illumination, the typical commercial W-LED is a combination of a blue InGaN LED chip with a yellow-emitting \( \text{Y}_3\text{Al}_5\text{O}_{12}:\text{Ce}^{3+} \) phosphor. But the deficiency in the red region brings about a low color rendering index (CRI) and high correlated color temperature (CCT).\(^18\)–\(^19\) In order to overcome the above defects, alternative W-LEDs have been developed using tricolor (red, green, and blue) phosphors pumped by near-ultraviolet (near-UV) LED chips.\(^20\)–\(^22\) However, the current red phosphors like sulfides and oxysulfides (\( \text{SrY}_2\text{S}_4: \text{Eu}^{2+} \) (ref. 22) and \( \text{Y}_2\text{O}_2\text{S}:\text{Eu}^{3+} \) (ref. 23)) have poor chemical stability and are moisture insensitive. Recently, \( \text{Eu}^{2+} \)-activated nitrides compounds (\( \text{Ba}_2\text{Si}_5\text{N}_8: \text{Eu}^{2+} \) (ref. 24) and \( \text{CaAlSiN}_3: \text{Eu}^{2+} \) (ref. 25)) have been synthesized as red-emitting phosphors to overcome the above-mentioned problems,\(^26\) but the rigorous synthesis conditions such as high pressure, certain atmosphere, and high temperature restrict the practical applications. Therefore, it is significant to search novel red phosphors with excellent chemical stability, high-efficiency, good color purity, and especially high absorption in near-UV or blue wavelength region for W-LED application.

Rare-earth ions activated inorganic materials have been investigated extensively on account of their narrow emission bands and excellent luminescent properties.\(^27\) Among them, the trivalent europium (\( \text{Eu}^{3+} \)) ion is a significant activator. In many inorganic compounds, \( \text{Eu}^{3+} \) ion mainly shows an efficient bright red emission around 615 nm attributing to intra 4f–4f transitions.\(^28\)–\(^30\) Meanwhile, the oxide phosphors are more stable in comparison with sulfide phosphors. In the past few years, oxide based phosphors have been widely investigated as potential candidates for W-LEDs applications.\(^31\) Li et al. have studied the color-tunable \( \text{Bi}^{3+}/\text{Eu}^{3+} \) co-doped \( \text{Ba}_3\text{Y}_4\text{O}_9 \) phosphors; however, to our best knowledge, the \( \text{Eu}^{3+} \) single-doped \( \text{BYO} \) phosphors have not been reported and they may have better luminescence properties for further
research. In this paper, we reported Ba₃Y₂O₆:Eu³⁺ (BYO:Eu³⁺) red-emitting phosphors with superior thermal stability, internal quantum efficiency (IQE), and high color purity, which satisfied the potential application in fabrication of W-LEDs based on the near-UV/blue LED chip.

2. Experimental

2.1 Sample synthesis

The Ba₃(Y₁₋ₓEuₓ)₄O₉ (abbreviated as BYO:xEu³⁺; x = 0.10, 0.15, 0.20, 0.25, 0.30, 0.35, and 0.40) phosphors with different Eu³⁺ concentrations were synthesized via a conventional solid-state reaction route in an air atmosphere. Starting materials of BaCO₃ (analytical reagent), Y₂O₃ (99.99%), and Eu₂O₃ (99.99%) were weighed according to stoichiometric ratio and then mixed thoroughly in an agate mortar. Subsequently, the obtained mixtures were transferred into the alumina crucibles and sintered at 1400 °C for 5 h in a furnace. Finally, when the samples cooled down slowly to room temperature, the obtained products were ground again into powders for further characterizations.

2.2 Characterization

The X-ray diffraction (XRD) patterns of the as-prepared phosphors were recorded on Bruker D8 Advance diffractometer with Cu-Kα radiation (λ = 1.54056 Å). The morphology was analyzed by using a field-emission scanning electron microscope (FE-SEM; TESCAN MAIA3). The photoluminescence (PL) and photoluminescence excitation (PLE) spectra were examined by the Edinburgh FS5 spectrometer equipped with a 150 W continuous-wave xenon lamp. The IQE, decay curves, and temperature-dependent emission spectra of the samples were also measured by using the Edinburgh FS5 spectrometer equipped with an integrating sphere coated with barium sulfate, a pulsed xenon lamp, and a temperature controller.

2.3 Fabrication of the W-LED device

The commercial blue phosphor BaMgAl₁₁O₁₉:Eu²⁺ (BAM:Eu²⁺), commercial green phosphor (Ba,Sr)₂SiO₄:Eu²⁺ and our red phosphor BYO:0.25Eu³⁺ were mixed thoroughly with silicone coated on the surface of a 395 nm LED chip to fabricate a W-LED device. The fabricated device was driven by 3 V voltage and 60 mA current and the photoelectric properties of the device were measured by using an integrating sphere spectroradiometer system (HAAS2000, Everfine).

3. Results and discussion

The XRD patterns of BYO:xEu³⁺ (x = 0.10, 0.15, 0.20, 0.25, 0.30, 0.35, and 0.40) phosphors were displayed in Fig. 1. It can be observed that all of the diffraction peaks were well-matched with the standard PDF#89-5891 card of Ba₃Y₂O₆, and there was no other impurity phase. In addition, due to the similar ionic radii between Y³⁺ (r = 0.900 Å, CN = 6; CN is the coordination number) and Eu³⁺ (r = 0.947 Å, CN = 6) ions, there was no obvious shift in the diffraction peaks. The radius percentage difference between the dopants (Eu³⁺) and substituted ions (Y³⁺) can be calculated by the following equation:

\[
D_r = \frac{R_c(CN) - R_d(CN)}{R_c(CN)} \times 100\%
\]

where \(D_r\) is the radius percentage difference; \(R_c\) and \(R_d\) represent the ionic radii of host ion and doped ion. In this case, the value of \(D_r\) was determined to be 5.2%, which was far smaller than 30%. The result further manifested that the doped ions entered into the host lattice smoothly without any significant influence on the crystal structure.

Fig. 2(a) represents the Rietveld XRD refinement of Eu³⁺-activated BYO phosphors and the refined crystallographic parameters were summarized in Table 1. As shown, the experimental and calculated spectra were accordant with each other, indicating that the BYO compound belonged to trigonal system with R3 space group. Meanwhile, the cell parameters were \(a = b = 6.12099 \text{ Å}, c = 25.2800(8) \text{ Å}, \text{ and } V = 820.258(35) \text{ Å}^3\). Fig. 2(b) shows the crystal structure of BYO, it can be seen that there were four types of Y atom sites, named as Yn \((n = 1, 2, 3, 4)\). All of the Y atoms were surrounded by 6 oxygen atoms to form octahedrons. In addition, there were three types of Ba atom sites in this crystal structure, defined as Ba1, Ba2, and Ba3, which were coordinated with 6, 6, and 3 oxygen atoms, respectively.

Considering the same valence state, Eu³⁺ ions preferred to occupy Y³⁺ sites rather than Ba²⁺ sites, which were consistent with the previous results.

The representative FE-SEM image of BYO:0.25Eu³⁺ phosphors was shown in Fig. 3(a). The obtained compounds consisted of aggregated and irregular particles with the size of several micrometers. Moreover, as displayed in Fig. 3(b), elements of O, Y, Ba, and Eu were observed, and the elemental mapping results implied that the elements in BYO were evenly distributed over the whole particles.

Since the optimal doping concentration was 0.25 mol, the representative PLE and PL spectra of BYO:0.25Eu³⁺ phosphors were shown in Fig. 4. The PLE spectrum contained a broad band
and several sharp peaks when monitored at 614 nm. The broad band ranging from 250 to 300 nm was attributed to $O^{2-} \rightarrow \text{Eu}^{3+}$ charge transfer band, which represented the electronic transition from 2p orbital of $O^{2-}$ to empty 4f orbital of Eu$^{3+}$. Meanwhile, several sharp excitation peaks located at 325, 366, 387, 396, 407, and 469 nm were assigned to $^5D_0 \rightarrow ^7H_{6}$, $^7F_0 \rightarrow ^5D_1$, $^7F_0 \rightarrow ^5G_2$, $^7F_0 \rightarrow ^5H_{6}$, $^7F_0 \rightarrow ^5D_3$, and $^7F_0 \rightarrow ^5D_2$ transitions, respectively.\textsuperscript{52–37} Particularly, the excitation intensities of $^7F_0 \rightarrow ^5L_6$ (396 nm) and $^7F_0 \rightarrow ^5D_2$ (469 nm) transitions were almost equivalent. Because the energy of near-UV light was larger than that of blue light, under the same situation, the PL intensity of BYO:0.25Eu$^{3+}$ phosphors upon 396 nm excitation was stronger than that upon 469 nm, as shown in the PL spectra. The emission bands peaking at 594, 614, 656, and 704 nm were ascribed to $^5D_0 \rightarrow ^7F_j$ ($j = 1, 2, 3, \text{and } 4$) transitions of Eu$^{3+}$ ions.\textsuperscript{38–40} Besides, the shape profiles of peaks had no variation except a slight change in PL intensity under the excitation of 396 and 469 nm, and also the strongest peak at 614 nm was ascribed to the $^5D_0 \rightarrow ^7F_1$ transition.

The magnetic dipole (MD) $^5D_0 \rightarrow ^7F_1$ transition (594 nm) is insensitive to the distortion of the inversion symmetry, and inversely the electric dipole (ED) $^5D_0 \rightarrow ^7F_2$ transition (614 nm) is hypersensitive.\textsuperscript{41} To evaluate the site symmetry of Eu$^{3+}$ ions, the integrated PL intensity ratio (asymmetry ratio) $(^5D_0 \rightarrow ^7F_2/ ^5D_0 \rightarrow ^7F_1)$ suggests the degree of distortion.\textsuperscript{42} In our case, the ED (614 nm) $^5D_0 \rightarrow ^7F_2$ transition was stronger than the MD (594 nm) $^5D_0 \rightarrow ^7F_1$ transition. The asymmetry ratio value of BYO:0.25Eu$^{3+}$ was calculated to be 3.54, and others with different doping concentrations were listed in Table 2. This result revealed that the Eu$^{3+}$ ions primarily occupied the non-inversion symmetry lattice sites in BYO host material.

Fig. 5(a) and (b) present the PL spectra of BYO:xEu$^{3+}$ phosphors with different doping concentrations under the excitation of 396 and 469 nm, respectively. It can be seen that the contours of the PL spectra hardly varied with the increase of the Eu$^{3+}$ concentration. The normalized PL intensities of BYO:xEu$^{3+}$ phosphors as functions of Eu$^{3+}$ concentrations under the excitation of 396 and 469 nm were displayed in Fig. 5(c). It was clearly found that the overall intensities under 469 nm excitation were slightly weaker than that under 396 nm excitation. These results indicated that the BYO:xEu$^{3+}$ phosphors can be effectively pumped with near-UV and blue LED chips. Moreover, the PL intensity gradually enhanced when Eu$^{3+}$ ions concentration increased, then it reached a maximum at $x = 0.25$. Subsequently, when $x$ was further increased, the PL intensity decreased due to the concentration quenching effect, which was triggered by the nonradiative energy transfer among the nearest Eu$^{3+}$ ions.\textsuperscript{43} According to the theory proposed by Blass,\textsuperscript{44} the concentration quenching mechanism can be determined by the critical distance ($R_c$) of Eu$^{3+}$ ions. If $R_c$ is larger than 5 Å, the electric multipolar interaction will lead to the concentration quenching phenomenon, otherwise it will be the exchange interaction. Therefore, it’s necessary to calculate $R_c$ via the following formula:\textsuperscript{44}

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

where $V$ is the volume of the unit cell, $x_c$ is the critical concentration and $N$ refers to the number of formula units per unit cell. In this present paper, $V = 820.258 \, \text{Å}^3$, $N = 3$, and $x_c = 0.25$. According to the above equation, the value of $R_c$ was determined to be 12.78 Å and it was much larger than 5 Å. In BYO:xEu$^{3+}$ phosphors, the energy transfer among Eu$^{3+}$ ions took place in the manner of electric multipolar interaction. In addition, to further research the interaction between Eu$^{3+}$ ions in BYO:Eu$^{3+}$ phosphors, Dexter’s formula of multipolar interaction was used as expressed below:\textsuperscript{45}

$$I / x = K [1 + \beta(x)^j]^{-1} \quad (3)$$
where $I$ is the emission intensity, $x$ is the activator concentration, $K$ and $\beta$ are the constants for the same excitation condition, while $\theta = 6, 8, \text{ and } 10$ correspond to the dipole–dipole (d–d), dipole–quadrupole (d–q), and quadrupole–quadrupole (q–q) interactions, respectively. The slope of the fitting line between $\log(I/x)$ and $\log(x)$ was $-3.7$, as shown in Fig. 5(d). Thus the value of $\theta$ was calculated to be $11.1$, which was close to $10$, indicating that the energy transfer among Eu$^{3+}$ ions can be attributed to q–q interaction in BYO:Eu$^{3+}$ phosphors.

The energy level diagram of Eu$^{3+}$ ions was presented in Fig. 6(a) to demonstrate the energy transfer process. Upon the excitation of near-UV (396 nm) and blue (469 nm) light, the electrons at ground state ($^7F_0$) transited to the higher excited states $^5D_0$ and $^5D_1$, then relaxed to lower excited state ($^5D_0$) level through nonradiative transition. Subsequently, the lower excited state electrons returned to $^7F_J$ ($J = 1, 2, 3, \text{ and } 4$) levels, resulting in four characteristic emissions due to the $^5D_0$/$^7F_J$ transitions, as shown in Fig. 6(a).

Fig. 6(b) shows the room-temperature decay curves of $614$ nm red emission in BYO:xEu$^{3+}$ phosphors with different Eu$^{3+}$ concentrations under the excitation of 396 nm. It’s necessary to calculate the decay lifetimes and all the decay curves can be well fitted with the double-exponential equation:

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

(4)

where $I(t)$ is the PL intensity at time $t$; $\tau_1$ and $\tau_2$ represent the lifetimes for exponential components; $A_1$ and $A_2$ are constants. The average lifetime $\tau_s$ of Eu$^{3+}$ can be calculated by the following function:

$$\tau_s = (A_1\tau_1^2 + A_2\tau_2^2)/(A_1\tau_1 + A_2\tau_2)$$

(5)

the $\tau_s$ were found to be $0.695, 0.650, 0.613, 0.521, 0.438, 0.231, \text{ and } 0.187$ ms for BYO:xEu$^{3+}$ phosphors with Eu$^{3+}$ concentrations $x = 0.10, 0.15, 0.20, 0.25, 0.30, 0.35, \text{ and } 0.40$, respectively. From Fig. 6(b), it was observed that $\tau_s$ dropped slowly at the beginning, when the Eu$^{3+}$ concentration reached $0.25$ mol, the decay lifetimes started to decline quickly, revealing the existence of the nonradiative energy transfer between the Eu$^{3+}$ ions.

The CIE chromaticity coordinates are crucial factors to evaluate the luminescent properties of phosphors. Fig. 7(a) illustrates the CIE chromaticity diagram of BYO:0.25Eu$^{3+}$ phosphors. According to the PL spectra, the CIE chromaticity coordinates of BYO:xEu$^{3+}$ phosphors ($\lambda_{ex} = 396$ nm) with various Eu$^{3+}$ concentrations were summarized in Table 2. The green and blue asterisks in Fig. 7(a) represented the locations of CIE chromaticity.

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**Table 2.** The asymmetry ratio, CIE chromaticity coordinates, and color purity of BYO:Eu$^{3+}$ phosphors with different Eu$^{3+}$ concentrations

<table>
<thead>
<tr>
<th>Eu$^{3+}$ concentration</th>
<th>Asymmetry ratio</th>
<th>CIE coordinates ($x$, $y$)</th>
<th>Color purity</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.10</td>
<td>3.36</td>
<td>(0.6662, 0.3335)</td>
<td>97.0%</td>
</tr>
<tr>
<td>0.15</td>
<td>3.50</td>
<td>(0.6682, 0.3316)</td>
<td>97.5%</td>
</tr>
<tr>
<td>0.20</td>
<td>3.48</td>
<td>(0.6689, 0.3308)</td>
<td>97.7%</td>
</tr>
<tr>
<td>0.25</td>
<td>3.54</td>
<td>(0.6695, 0.3302)</td>
<td>97.9%</td>
</tr>
<tr>
<td>0.30</td>
<td>3.41</td>
<td>(0.6685, 0.3312)</td>
<td>97.6%</td>
</tr>
<tr>
<td>0.35</td>
<td>3.40</td>
<td>(0.6678, 0.3320)</td>
<td>97.4%</td>
</tr>
<tr>
<td>0.40</td>
<td>3.24</td>
<td>(0.6662, 0.3335)</td>
<td>97.0%</td>
</tr>
</tbody>
</table>
coordinates of BYO:0.25Eu³⁺ phosphors under the excitation of 396 and 469 nm, respectively. The corresponding chromaticity coordinates were calculated to be (0.6695, 0.3302) and (0.6692, 0.3305), which were very close to that of the standard red light (0.6700, 0.3300). Furthermore, it can be clearly seen from Fig. 7(a) that the both points basically coincided with each other, indicating that the phosphors can be excited efficiently by both near-UV and blue light from another point of view. The color purity was further investigated as follows:

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**Fig. 5** (a) PL spectra of BYO:xEu³⁺ phosphors under 396 nm excitation. (b) PL spectra of BYO:xEu³⁺ phosphors under 469 nm excitation. (c) Normalized PL intensity of BYO:xEu³⁺ phosphors as a function of Eu³⁺ concentration. (d) Plot of log(I/I₀) vs. log(x) of Eu³⁺ ions in BYO:xEu³⁺ phosphors excited at 396 nm and monitored at 614 nm.

**Fig. 6** (a) Energy level diagram of Eu³⁺ ions. (b) Decay curves of BYO:xEu³⁺ phosphors excited at 396 nm and monitored at 614 nm.
Color purity = \frac{\sqrt{(x - x_d)^2 + (y - y_d)^2}}{\sqrt{(x - x_i)^2 + (y - y_i)^2}} \times 100\% \quad (6)

where \((x, y)\), \((x_i, y_i)\), and \((x_d, y_d)\) present the CIE chromaticity coordinates of the sample, white illumination, and dominated wavelength, respectively. For BYO:0.25Eu³⁺ phosphors under 396 nm excitation, \((x, y) = (0.6695, 0.3302)\), \((x_i, y_i) = (0.310, 0.316)\), and \((x_d, y_d) = (0.6775, 0.3224)\). In this work, the value of color purity reached up to 97.9%, which was higher than that of Ca₂SiO₄:Eu³⁺ (91.2%), NaBiF₄:Eu³⁺ (90.4%), and CaWO₄:Eu³⁺ (91.3%), meaning that the high quality red-emitting BYO:0.25Eu³⁺ phosphors were obtained. The color purity of BYO:xEu³⁺ samples with different Eu³⁺ doping concentrations were listed in Table 2.

Another key factor to assess the application of the phosphors is IQE. Fig. 7(b) depicts the IQE of the BYO:0.25Eu³⁺ red-emitting phosphors and the value of IQE can be obtained by the following equation:  

\[ \eta = \frac{\int L_S}{\int E_R - \int E_S} \quad (7) \]

where \(L_S\) is the emission spectrum of the sample, \(E_S\) and \(E_R\) are the spectra of excitation light with and without sample, respectively. Under 396 nm excitation, the IQE of BYO:0.25Eu³⁺ phosphor was calculated to be 46%, which was slightly higher than that upon 469 nm excitation (IQE: 40%). In addition, the external QE of BYO:0.25Eu³⁺ phosphors excited by near-UV light was found to be 5%.

Fig. 8(a) shows the temperature-dependent PL spectra of BYO:0.25Eu³⁺ phosphors excited at 396 nm. As the temperature increased from 303 K to 503 K, the PL intensity decreased gradually due to the thermal quenching effect. The normalized PL intensity of BYO:0.25Eu³⁺ phosphors as a function of temperature was depicted in the inset of Fig. 8(a). As shown, the PL intensity at 423 K remained about 89% relative to its initial value at 303 K, which was higher than that of other reported Eu³⁺ doped phosphors, such as Ca₃Gd(AlO)₃(BO₃)₄:Eu³⁺ (84%).
and Na₃Sc₂(PO₄)₃:Eu³⁺ (73.4%), indicating that the phosphors had a splendid thermal stability. Moreover, another parameter, activation energy was used to evaluate the thermal property according to the Arrhenius equation:

\[
I(t) = \frac{I_0}{1 + c \exp\left(-\frac{E_a}{kT}\right)}
\]

where \(I_0\) is the PL intensity at room temperature, \(I(t)\) is the PL intensity at different given temperatures, \(E_a\) is the activation energy, \(c\) is a constant, and \(k\) is the Boltzmann constant (8.629 × 10⁻⁵ eV). Fig. 8(b) illustrates the linear relationship between \(\ln\left(\frac{[I_0/I]}{1}\right)\) and \(1/kT\), and the value of \(E_a\) was estimated to be 0.29 eV by the slope of the fitting line. Generally, the phosphors present superior thermal stability when \(E_a\) is high. The high \(E_a\) value suggested that the BYO:xEu³⁺ phosphors possessed good thermal stability and were suitable for W-LEDs applications.

For further testing the application of BYO phosphors in W-LEDs, a W-LED device was fabricated based on 395 nm near-UV chip, commercial (Ba,Sr)₂SiO₄:Eu²⁺ green-emitting phosphors, commercial BAM:Eu²⁺ blue-emitting phosphors, and as-prepared BYO:0.25Eu³⁺ red-emitting phosphors. As disclosed in Fig. 9, under 60 mA current and 3 V voltage, the device gave bright warm white light with high CRI (92) and low CCT (2850 K). The CIE chromaticity coordinates of the W-LED device were (0.4454, 0.4025), which located at warm white range. The above results further demonstrated that the Eu³⁺-activated BYO phosphors had potential application for warm W-LEDs.

4. Conclusion

In summary, the BYO:xEu³⁺ red phosphors were synthesized via a conventional solid-state reaction. The BYO:xEu³⁺ phosphors which can be effectively excited at 396 nm (near-UV) and 469 nm (blue) exhibited an intense red emission around 614 nm. The optimal concentration of Eu³⁺ was \(x = 0.25\). Furthermore, the CIE chromaticity coordinates of BYO:0.25Eu³⁺ phosphors were (0.6695, 0.3302) and the value of IQE reached up to 46%. Meanwhile, the BYO:0.25Eu³⁺ phosphors also exhibited high thermal stability due to the fact that their PL intensity at 423 K maintained 89% of its starting value at 303 K and the activation energy was up to 0.29 eV. In addition, a warm W-LED device with excellent properties was fabricated with the help of a 395 nm chip, as-prepared BYO:0.25Eu³⁺ red phosphors, commercial (Ba,Sr)₂SiO₄:Eu²⁺ green-emitting phosphors, and commercial BAM:Eu²⁺ blue-emitting phosphors. All of the obtained results suggested that our BYO:xEu³⁺ phosphors were promising candidates for warm W-LEDs application pumped by both near-UV and blue light.

**Conflicts of interest**

There are no conflicts to declare.

**Acknowledgements**

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