Gd₃B(W,Mo)O₉: Eu³⁺ red phosphor: From structure design to photoluminescence behavior and near-UV white-LEDs performance

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Abstract

The photoluminescence behavior of a high efficient red phosphor Gd₃B(W,Mo)O₉:Eu³⁺ designed according to crystal structure and electronic structure is presented in detail. The substitution of isolated WO₆ group by MoO₆ makes the absorption band edge extend from 360 nm to 440 nm, as designed to fit the excitation of near ultraviolet (NUV) LED chip, which could be interpreted by the density functional theory calculations. The extremely low symmetry of Eu³⁺ site enables Eu³⁺ ion to show dominant electronic dipole transition 2D₀ → 2F₂ (~616 nm) with five apparent splitting peaks according to group theory, which is almost 10 times stronger than that of commercial Y₂O₃:S:Eu³⁺ upon ~385 nm excitation. The outstanding thermal stability of photoluminescence of this phosphor up to 523 K may probably be owed to the energy transfer from isolated MoO₆ groups to the well-dispersed Eu³⁺ ion separated by BO₆ groups. The performance of NUV white-LEDs package fabricated with a blue, green and red phosphor is also demonstrated. It shows high color stability of the white light (CIE coordinate (0.320, 0.349), CCT 6031K, at 20 mA) under the DC drive current varied from 20 mA to 350 mA, verifying the potential application of the red phosphor in tri-color phosphors coated white-LEDs.

1. Introduction

White LEDs (w-LEDs), considered as the next generation for general lighting, have attracted much attention because of their unique advantages, for instance, high color rendering index, long lifetime, low power consumption, and environmental friendliness [1–4]. Remarkable progress has been made in the development of w-LEDs with blue LED chip coated with yellow phosphor, while for that assembled using a near ultraviolet (NUV) LED chip with tri-color phosphors, which is viewed as a more promising approach to have a better white light performance, is less investigated [3–5]. Some phosphors are selected for NUV w-LEDs application, for instance, BaMgAl₁₂O₁₉:Eu³⁺ as blue phosphor, ZnS:Cu⁺Al³⁺ as green phosphor, and Y₂O₂S:Eu³⁺ as red phosphor [6]. However, the red phosphor Y₂O₂S:Eu³⁺ could not efficiently absorb NUV (370–400 nm) light emitted by NUV LED chip, resulting in low emission efficiency of this red phosphor and blocking the development of NUV w-LEDs [7,8]. Therefore, it is very urgent to develop new efficient red phosphors for NUV w-LEDs [9–11]. Red phosphors with efficient absorption of the NUV light and emitting light at 610 nm are considered the best choice for a light source with respect to luminous efficiency and color rendering [8,12,13]. Recent theoretical analysis has shown that four narrow emitting sources at 459 nm, 535 nm, 573 nm, and 614 nm, respectively, are needed to obtain high luminous efficiency, high color rendering index and low correlated color temperature [14]. Particularly the red phosphor emitting 610–620 nm with FWHM less than 20 nm is highly desirable [14,15]. For this, Eu³⁺ ion is the perfect and unique choice. Moreover, the temperature dependence of photoluminescence is also significant since it has great influence on the light output. Meanwhile, color rendering index of phosphor-coated w-LEDs is also influenced by temperature, which is a significant issue for device application since the w-LEDs devices radiate much heat [16].

Compounds with the MO₆ group (M is a high valence ion), such as vanadates, molybdates, tungstates, are widely used as hosts for Eu³⁺-doped red phosphors. These hosts could efficiently absorb UV light through the charge transfer state (CTS) of MO₆ group and easily transfer the absorbed energy to Eu³⁺ for red emission. Tungstates widely attract people’s interests due to its high melting point, excellent thermal stability and excellent...
photoluminescence properties [17–21]. It is reported that Eu4+BWO3 bortungstate exhibits attractive red emission performance [7]. While it shows sharp line-shape excitation characteristic in the NUV region, which is not desirable for those NUV LED chips produced by different corporation or different manufacturing batches with different emitting wavelengths. The charge transfer band (CTB) of isolated WO6 group in Eu4+BWO3 locates at about 270–330 nm, and the substitution of WO6 group by MoO6 group would tune the CTB of the bortungstate to the desired NUV region as demonstrated in other cases [8,15,22–25]. To avoid the use of expensive and highly concentrated Eu3+ ions, Gd3+BWO3 is utilized as phosphor host. The well-defined single site of the cations (Gd3+Eu3+4Mo6+,0.06Eu3+) [7,26] makes it easier to discuss the relationship between the structure and photoluminescence properties, which would open a perspective to design novel red phosphors. This work will give insight into the crystal structure-electronic structure-photoluminescence behavior of Gd4+Bi(W,Mo)O12:Eu3+ and its performance in the fabricated w-LEDs application.

2. Experiment

2.1. Sample preparation

Nominal Gd3+BW1−xOxMo6+,0.06Eu3+ (x = 0–0.4) and Gd3+BW0.80Mo0.20O9:0.20Mo6+,0.06Eu3+ (y = 0.06–1.20) powder samples were synthesized through a high-temperature solid-state reaction technique. The starting materials were MoO3 (analytical reagent, A.R.), Eu2O3(99.99%), Gd2O3(99.99%), WO3(A.R.) and H2BO3(A.R.). The stoichiometric amounts of the starting materials according to nominal formulae Gd3+BW1−xOxMo6+,0.06Eu3+ (x = 0–0.40) and Gd3+BW0.80Mo0.20O9:0.20Mo6+,0.06Eu3+ (y = 0.06–1.20) were mixed and ground thoroughly in an agate mortar. The mixture was firstly preheated at 500 °C for 5 h, then at 1100 °C for 10 h to eliminate the evaporation of MoO3, and finally fired at 1200 °C for 10 h to form the single-phased Gd3+BW1−xOxMo6+,0.06Eu3+ in a muffle furnace with some intermediate grindings. In here 3 mol% excess of H2BO3 was added as flux to accelerate the reaction.

2.2. Characterization

Phase identification of the obtained products were analyzed by means of a Philips PW1830 X-ray powder diffractometer (XRD) using graphite monochromator and Cu Kα (λ = 1.54056 Å) radiation at 40 kV and 40 mA. XRD data for refinement were collected on a Bruker D8 ADVANCE X-ray diffractometer with Cu target and Ni filter. The step-scans mode was used with setup of 40 kV × 40 mA, step 0.02°, 0.2 s/step. Rietveld refinement was conducted by TOPAS Academic program. Diffuse reflection spectra were measured on Hitachi U-3010 spectrophotometer with BaSO4 as a reference. Theoretical calculations based on density functional theory (DFT) were conducted by the Vienna ab initio simulation package [27] with the frozen-core projector-augment-wave (PAW) method [28]. The generalized gradient approximation [29] and Pw(Pw) = 1 × 1 × 1 k points were employed in the calculations. The energy cutoff of the plane wave basis set was set to 450 eV. The convergence criterion for the electronic energy was 10−4 eV and the structures were relaxed until the Hellmann–Feynman forces were less than 0.02 eV/Å. The excitation and photoluminescence (PL) spectra were recorded on the Jobin Yvon TRIXAX320 spectrometer equipped with a 450-W xenon lamp as the excitation source and a Hamamatsu R928 photomultiplier tube as detector. The temperature-dependent PL spectra were measured on the same spectrophotometer equipped with an additional TAP-02 high-temperature fluorescence instrument (Tian Jin Orient–KJII instrument Co., Ltd.). Decay curves were recorded on an Edinburgh FLS920 spectrophotometer with a µFlash 300 flash lamp as the excitation resource and a thermo-electron cooled red sensitive Hamamatsu R928 photomultiplier tube as detector. NUV w-LEDs were fabricated by coating proper amounts of red phosphor of Gd3+Bi(W,Mo)O12:Eu3+; a green phosphor of BaMgAl11O19:Eu2+,Mn2+ (Lab-prepared) [30] and a blue phosphor of alkaline earth chloroborate (Lab-prepared) [31] on an NUV LED chip (λ = 385 nm, EP-U4543C-A3) as the primary light source. The commercial Y2O3:S:Eu phosphor (Red-1, Dalian Luminglight Co., Ltd., China) is used for the comparison with the as-prepared Gd3+Bi(W,Mo)O12:Eu3+ phosphor. The optical properties of w-LEDs were performed by YF1000 lamp complete analysis system under a forward-bias current of 20–350 mA at room temperature (RT).

3. Results and discussion

3.1. XRD patterns and crystal structure

Some typical XRD patterns of Gd2.94BW1−xOxMo6+,0.06Eu3+ (x = 0, 0.10, 0.20, 0.40) powder samples are demonstrated in Fig. 1. The standard card of Gd3+BWO3 (JCPDS 50-1859) is also given for comparison. The XRD pattern of Gd2.94BWO3:0.06Eu3+ sample (red curve) agrees well with the standard card of Gd3+BWO3 (black curve) in Fig. 1, indicating that the samples is single-phased and the crystal structure is not significantly changed when the dopant ions incorporate into the host. Owing to the similar radii and the same electrovalence for Eu3+ and Gd3+, we deduce that Eu3+ ion substitutes for Gd3+ ion in the host. The XRD patterns of Gd2.94BW1−xOxMo6+,0.06Eu3+ (x = 0.10, 0.20) (magenta curve, cyan curve) powder samples are also consistent with the standard card of Gd3+BWO3 (black curve). While for the sample with high Mo content x = 0.4, additional peaks of impurity phase (Gd3+MoO6, JCPDS file 26-0656) appear. The result suggests that the W-Mo solid solution range x in this phosphor is no larger than 0.40. To investigate the crystal structure, the Rietveld refinement of XRD pattern of Gd2.94BWO3:0.06Eu3,0.20Mo6+,0.06Eu3+ was performed (see Fig. 2), and the results are listed in Table 1. This compound has a space group of P63/m, with lattice parameters of a = b = 8.5571 Å, c = 5.4079 Å, and cell volume of 342.93 Å3. Every cation has only one crystallographic site, and the Gd/Eu site has an extremely low symmetry C1. The isolated (W/Mo)O6 groups and the isolated BO3 groups with dispersive Eu3+ ions are arranged along C6 rotation axis (c axis), as depicted in Fig. 3.

3.2. Diffuse reflection spectra, electronic structure and photoluminescence behavior

Diffuse reflection spectra of Gd2.94BW1−xOxMo6+,0.06Eu3+ (x = 0, 0.05, 0.10, 0.20, 0.25) phosphors are shown in Fig. 4. It can be seen that Gd2.94BWO3:0.06Eu3+ has a strong absorption band in the range of 240–360 nm, which is mainly ascribed to the CTS of WO6 group [8,15,22–25]. With the increase of Mo content x, the absorption band gradually shifts to the long wavelength, reaching a maximum value of 440 nm when x = 0.20. Apparently, the redshift is owing to the CTS of MoO6 group [8,15,22–25], and the gradual redshift implies there is interaction among MoO6 groups [24]. To make better understand of this redshift, DFT calculations of electronic structure of Gd3+BWO3 and Gd3+BW0.80Mo0.20O9 were conducted and their band structures are depicted in Fig. 5(a) and (b), respectively. It is clear that some defect levels are formed in the band gap of Gd3+BW0.80Mo0.20O9 compared to that of Gd3+BWO3, which could be responsible to the redshift shown in Fig. 4. The partial density states, as shown in Fig. 5(c), indicate that the defect bands are attributed to the incorporation of Mo, in line with the
corresponding experimental observation. Meanwhile, we found that the valence bands are dominated by oxygen while the conduction bands are mainly contributed by W/Mo (Fig. 5(c)), in line with the earlier reports that the gap is correlated to the CTS of WO$_6$ and MoO$_6$ groups [8,15,22–25].

Typical PL spectra of selected Gd$_{2.94}$BW$_{0.80}$O$_9$:0.20Mo$^{6+}$,0.06-Eu$^{3+}$ upon excitation of 385 nm light (red curve) and Gd$_{2.94}$BWO$_9$:0.06-Eu$^{3+}$ upon excitation of 340 nm light (black curve) are described in Fig. 6(a). Emission peaks at 578, 588–598, 610–622, 650, 701–704 nm are assigned to $5D_0 \rightarrow 7F_J$ ($J = 0–4$) transition of the Eu$^{3+}$ ion, respectively. The dominant electronic dipole transition of $5D_0 \rightarrow 7F_2$ suggests that Eu$^{3+}$ ion locate at the site without a symmetric center, which is coincident with the crystallographic data in Table 1. According to group theory and character tables, the $C_1$ symmetry of Eu$^{3+}$ site (see the seven-coordinated polyhedron in the insert of Fig. 6(a)) would result in five splitting peaks for $5D_0 \rightarrow 7F_2$ electronic dipole transition. There are three irreducible representations $\langle A + A + A \rangle$ for electric dipole transition Hamiltonian, five irreducible representations $\langle A + A + A + A + A \rangle$ for $7F_2$ level and one irreducible representation for $5D_0$ level when Eu$^{3+}$ ion locates at a site with $C_1$ symmetry. Therefore, the direct product of initial state wavefunction and final state wavefunction $\langle \psi(5D_0) | \psi(7F_2) \rangle$ has five irreducible representations $\langle A + A + A + A + A \rangle$, each of which could be contained in the three irreducible

![Fig. 2.](image1.png)  
**Fig. 2.** The experimental (blue dotted line) and calculated (red solid line) XRD patterns and their difference (gray solid line). (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

<table>
<thead>
<tr>
<th>Compounds</th>
<th>Space group</th>
<th>Wyckoff position</th>
<th>$x$</th>
<th>$y$</th>
<th>$z$</th>
<th>Site symmetry</th>
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<tr>
<td>Gd/Eu</td>
<td>P$_6_3$</td>
<td>6c</td>
<td>0.3579(9)</td>
<td>0.0828(8)</td>
<td>0.211B(8)</td>
<td>C$_1$</td>
</tr>
<tr>
<td>W/Mo</td>
<td></td>
<td>2b</td>
<td>0.3333(3)</td>
<td>0.6666(7)</td>
<td>0.2500(0)</td>
<td>C$_1$</td>
</tr>
<tr>
<td>B</td>
<td></td>
<td>3a</td>
<td>0.0000(0)</td>
<td>0.0000(0)</td>
<td>0.7500(0)</td>
<td>C$_1$</td>
</tr>
<tr>
<td>O1</td>
<td></td>
<td>6c</td>
<td>0.1887(1)</td>
<td>0.3613(8)</td>
<td>0.7567(8)</td>
<td>C$_1$</td>
</tr>
<tr>
<td>O2</td>
<td></td>
<td>6c</td>
<td>0.2757(4)</td>
<td>0.3613(8)</td>
<td>0.7567(8)</td>
<td>C$_1$</td>
</tr>
<tr>
<td>O3</td>
<td></td>
<td>6c</td>
<td>0.1675(5)</td>
<td>0.4952(4)</td>
<td>0.4697(2)</td>
<td>C$_1$</td>
</tr>
</tbody>
</table>

![Fig. 3.](image2.png)  
**Fig. 3.** The sketch of Gd$_3$B(W,Mo)O$_9$ crystal structure along $c$ axis.

![Fig. 4.](image3.png)  
**Fig. 4.** Diffuse reflection spectra of Gd$_{2.94}$BW$_{1-x}$O$_9$:0.06Eu$^{3+}$ ($x = 0, 0.05, 0.10, 0.20, 0.25$) phosphors.
representations \( A + A + A \) of electric dipole transition Hamiltonian and thus all five splitting transition peaks are deduced theoretically. This is evidenced in Fig. 6(a) and in accordance with previous work [26]. The electronic dipole transition of \( ^5D_0 \to ^7F_2 \) at ~616 nm is desirable for tri-color phosphors coated NUV w-LEDs. The excitation spectra monitoring 616 nm emission of \( \text{Gd}_2\text{BWO}_9:x\text{Mo}^{6+},0.06\text{Eu}^{3+} \) \((x = 0, 0.05, 0.10, 0.20, 0.25)\) phosphors in Fig. 6(b), are composed of a broad band and some sharp lines. For \( \text{Gd}_2\text{BWO}_9:0.06\text{Eu}^{3+} \) sample, broad excitation band locates at 240–360 nm. With the increase of Mo content \( x \), the broad excitation band gradually extends to the long wavelength due to the CTB of \( \text{MoO}_6 \) groups [9], which is consistent with the diffuse reflection spectra. The broad bands originate from CTB of \( \text{WO}_6 \) groups for \( \text{Gd}_2\text{BWO}_9 \) \((x = 0)\) sample and from \( \text{MoO}_6 \) groups for \( \text{Gd}_2\text{BWO}_9:x\text{Mo}^{6+},0.06\text{Eu}^{3+} \) \((x = 0.05, 0.10, 0.20, 0.25)\) samples, manifesting that energy transfer process take place from \( \text{WO}_6/\text{MoO}_6 \) groups to \( \text{Eu}^{3+} \) ions. The strong sharp lines are the characteristic intra-configurational \( 4f-4f \) transitions of \( \text{Eu}^{3+} \): 395 nm \((^7F_0 \to ^5L_6)\) and 465 nm \((^7F_0 \to ^3D_2)\), indicating that the low symmetry of \( \text{Eu}^{3+} \) site would break down the parity of \( f \) orbital function and result in large increase of the parity-forbidden \( 4f-4f \) transition rate. The redshift is beneficial to NUV w-LEDs application because it matches well with the emitted wavelength of NUV LED chips [8]. Thus the optimum concentration for Mo in
phosphor Gd$_{2.94}$BW$_{1-x}$O$_{9}$:0.06Eu$^{3+}$ is 0.20. Fig. 6(c) presents the luminescence decay curves by monitoring emission at 616 nm for Gd$_{2.94}$BW$_{1-x}$O$_{9}$:0.06Eu$^{3+}$ ($x = 0.10, 0.20$) (blue curve, magenta curve) upon 385 nm excitation and for Gd$_{2.94}$BWO$_{9}$:0.06-Eu$^{3+}$ under 340 nm excitation (cyan curve). The luminescence lifetime of Gd$_{2.94}$BWO$_{9}$:0.06Eu$^{3+}$ phosphor is about 740 μs, which decreases slightly with the introduction of Mo. It may be caused by some defects introduced by Mo in the lattice, which act as non-radiative relaxation centers.

Fig. 7(a) illustrates the Eu$^{3+}$ concentration dependence of 616 nm emission intensity of Gd$_{3-y}$BW$_{0.80}$O$_{9}$:0.20Mo$^{6+}$, $y$Eu$^{3+}$ under excitation at 385 nm. With the increase of Eu$^{3+}$ concentration ($y$), the 616 nm emission intensity increases firstly and reaches a maximum value around $y = 0.80$, then decreases due to the concentration quenching [32], as seen in Fig. 7(a). Thus the optimum concentration $y$ for Eu$^{3+}$ ion in phosphor Gd$_{3-y}$BW$_{0.80}$O$_{9}$:0.20Mo$^{6+}$, $y$Eu$^{3+}$ is 0.80. The emission intensity of optimal phosphor is almost 10 times stronger than that of commercial Y$_2$O$_2$S:Eu$^{3+}$ upon the excitation of ~385 nm NUV light, as seen in Fig. 7(b). The thermal performance of phosphor is also a significant technological parameter for phosphors applied in w-LEDs [33], especially in high power devices [34]. The junction temperatures of typical LEDs can be higher than 373 K. Normally, if there is thermal quenching of a phosphor at $T > 373$ K, emission color would shift [35]. Fig. 7(c) demonstrates the dependence of 616 nm emission intensity of Gd$_{2.20}$BW$_{0.80}$O$_{9}$:0.20Mo$^{6+}$,0.80Eu$^{3+}$ on the temperature upon 465 nm (red curve) and 385 nm (blue curve) excitation. For 465 nm excitation corresponding to the $f$–$f$ transition of Eu$^{3+}$, the intensity almost keeps constant up to 423 K, and then decreases dramatically with the increase of temperature to 300 °C (decrease to about 60% of the initial value at 300 K when the temperature reaches 523 K). The sharp decrease at high temperature is due to the temperature quenching effect of Eu$^{3+}$ luminescence caused by energy migration and transfer to nonradiative traps in the host lattice [7]. While under the 385 nm excitation corresponding to the CTS of MoO$_6$ group, the intensity could maintain constant up to 523 K. There exists energy transfer from isolated MoO$_6$ group to the well-dispersed Eu$^{3+}$ upon excitation at 385 nm. The energy transfer would alleviate the decrease caused by temperature quenching effect of Eu$^{3+}$ itself. The outstanding thermal performance of the phosphor is highly desirable for NUV w-LEDs application.

3.3. w-LEDs performance

To evaluate the application of the red phosphor Gd$_{3-y}$B(W,Mo)O$_{9}$:Eu$^{3+}$ in NUV w-LEDs, we fabricated the w-LEDs device consisting of the as-prepared red phosphor Gd$_{3-y}$B(W,Mo)O$_{9}$:Eu$^{3+}$, a chloroborate blue phosphor [31], an aluminate green phosphor [30] and a NUV LED chip (emission wavelength 385 nm). The photos of the above mentioned phosphors under a 365 nm UV lamp are illustrated in Fig. 8(a). Intense red, blue and green light are observed. Images of fabricated w-LEDs lamp without and with a 20 mA DC drive current are shown in Fig. 8(b) and (c), respectively. Bright white light from the LED driven by a 20 mA direct current (DC) is observed by naked eyes and the emission spectrum of the w-LEDs lamp is demonstrated in Fig. 8(d). The Commission Internationale de l’Eclairage (CIE) chromaticity coordinate of the lamp operated at 20 mA is calculated to be (0.320, 0.349), locating at the white region as depicted in Fig. 8(e). The correlated color temperature (CCT) is 6031 K, very close to that of the daylight [12].

**Fig. 7.** (a) The dependence of 616 nm emission intensity under excitation at 385 nm for Gd$_{3-y}$BW$_{0.80}$O$_{9}$:0.20Mo$^{6+}$, $y$Eu$^{3+}$ on the concentrations ($y$) of Eu$^{3+}$ ions. (b) PL spectra comparison between Gd$_{2.20}$BW$_{0.80}$O$_{9}$:0.20Mo$^{6+}$,0.80Eu$^{3+}$ (red curve) and commercial red phosphor Y$_2$O$_2$S:Eu$^{3+}$ (blue curve) under excitation at 385 nm. (c) The dependence of 616 nm emission intensity under 465 nm (red curve) and 385 nm (blue curve) excitation for Gd$_{2.20}$BW$_{0.80}$O$_{9}$:0.20Mo$^{6+}$,0.80Eu$^{3+}$ on the temperature (Both data are normalized at 300 K). (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)
Fig. 8. (a) The photos of red/blue/green phosphors under a 365 nm UV lamp. (b) Images of fabricated w-LEDs. (c) Images of fabricated w-LEDs driven by a 20 mA DC current. (d) The emission spectrum of the fabricated w-LEDs lamp operated at 20 mA. (e) The CIE chromaticity coordinates for the lamp operated at 20 mA. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

Fig. 9. (a) The dependence of CIE chromaticity coordinates value \((x, y)\) and color temperature on DC drive current. (b) The dependence of three primary colors intensities on DC drive current. (c) The variation of CIE chromaticity coordinates with different DC drive current for w-LEDs. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)
The color stability of the w-LEDs lamp under various DC drive current were performed and the results are depicted in Fig. 9. Both the chromaticity coordinate x and y decline gradually with a larger trend for y, and the CCT varies from 6031 K to 8726 K under the DC drive current varied from 20 mA to 350 mA (Fig. 9a). The integral emission intensity of the optimal red phosphor leads to the extension of the excitation band to the NUV region, which is ascribed to the CTS of MoO6 octahedra. The integral emission intensity of the optimal red phosphor is about 10 times higher than that of commercial red phosphors. The outstanding thermal stability of photo-luminescence of these phosphors is probably owed to the energy transfer from isolated MoO6 groups to the well-dispersed Eu3+ ion separated by BO3 groups. The white light performance under various DC drive current were performed and the results are depicted in Fig. 9. Both the chromaticity of the lamp is highly stable when the DC drive current varied from 20 mA to 350 mA (Fig. 9a). The integral emission intensity of the optimal red phosphor is about 10 times higher than that of commercial red phosphors. The outstanding thermal stability of photo-luminescence of these phosphors is probably owed to the energy transfer from isolated MoO6 groups to the well-dispersed Eu3+ ion separated by BO3 groups. The white light performance under various DC drive current were performed and the results are depicted in Fig. 9. Both the chromaticity of the lamp is highly stable when the DC drive current varied from 20 mA to 350 mA (Fig. 9a). The integral emission intensity of the optimal red phosphor is about 10 times higher than that of commercial red phosphors.

4. Conclusions

In conclusion, we have developed a novel red phosphor Gd3B(W,Mo)O9:Eu3+ for NUV w-LEDs. The incorporation of Mo to this phosphor leads to the extension of the excitation band to the NUV region, which is ascribed to the CTS of MoO6 octahedra and evidenced by the DFT calculations. The phosphors exhibit strong and dominant electronic dipole transition D0 → F2 of Eu3+ with five obvious splitting peaks, which could be corresponded to the C1 site symmetry in the crystallographic data and interpreted by group theory. The integral emission intensity of the optimal phosphor is about 10 times higher than that of commercial red phosphors Y2O3:S:Eu3+. The outstanding thermal stability of photo-luminescence of these phosphors is probably owed to the energy transfer from isolated MoO6 groups to the well-dispersed Eu3+ ion separated by BO3 groups. The white light performance under various DC drive current is investigated. The research verified the potential application of the red phosphors Gd3B(W,Mo)O9:Eu3+ in tri-color phosphor coated NUV w-LEDs.

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Reference