Strongly enhanced luminescence of Sr₄Al₁₄O₂₅:Mn⁴⁺ phosphor by co-doping B³⁺ and Na⁺ ions with red emission for plant growth LEDs

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Development of a more cost-effective radiation source for use in plant-growing facilities would be of significant benefit for commercial crop production applications. A series of co-doped B³⁺ and Na⁺ ions Sr₄Al₁₄O₂₅:Mn⁴⁺ inorganic luminescence materials which can be used for plant growth were successfully synthesized through a conventional high-temperature solid-state reaction. Powder X-ray diffraction was used to confirm the crystal structure and phase purity of the obtained samples. Then scanning electron microscopy elemental mapping was undertaken to characterize the distribution of the doped ions. Detail investigations on the photoluminescence emission and excitation spectra revealed that emission intensity of tetravalent manganese ions can be well enhanced by monovalent sodium ions and trivalent boron ions under near-ultraviolet and blue excitation. Additionally, crystal field parameters and energies of states are calculated and discussed in detail. Particularly we achieve a photoluminescence internal quantum yield as high as 60.8% under 450 nm blue light excitation for Sr₄Al₁₄O₂₅:Mn⁴⁺, Na⁺, B³⁺. Therefore, satisfactory luminescence properties make these phosphors available to LEDs for plant growth.

Introduction

It is known that green plants can be grown only using red and blue monochromatic light, because chlorophyll has its second distinct absorption peak in the vicinity of 450 nm (the blue light region) and its first peak in the vicinity of 660 nm (red light region). The blue light is indispensable to the morphologically healthy plant growth, and the red light contributes to the plant leaf photosynthesis. The first successful plant growth experiment using only blue and red LEDs were achieved in June 1994 by Okamoto and Yanagi. With LEDs price reduced, LEDs are gradually used as plant lighting source, because of its high light efficiency, energy saving and other characteristics. As the driving currents of the blue chip and the red chip are inconsistent, the design of the driver circuit of LEDs for plant growth would be very complex, resulting in increased cost. Therefore, the plant-grown red lamp that is currently in widespread use in the marketplace is composed of a red phosphor rather than a red chip. The most commonly used red phosphors are nitride phosphor (e.g., Sr₂Si₃N₄:Eu²⁺, CaAlSiN₃:Eu²⁺) because of high luminous efficiency. But their high temperature (≈1600 °C), high pressure (1–10 MPa) preparation conditions and expensive raw materials containing rare earth elements lead to high cost.

In recent years, Mn⁴⁺-activated fluoride compounds, as an alternative to commercial (oxy)nitride phosphors, are emerging as a new class of non-rare-earth red phosphors for high-efficacy warm white LEDs compared to Mn⁴⁺-activated fluoride phosphor, the emission wavelength of Mn⁴⁺-activated oxide phosphors, such as Sr₄Al₁₄O₂₅:Mn⁴⁺, Y₃Al₅O₁₂:Mn⁴⁺, etc., is much longer which is very suitable for plant lighting, because the absorption peak of plant chlorophyll is near 660 nm. The following diagram shows spectroscopic range of Mn⁴⁺ ions in various crystals. We can see from Fig. 1 that the emission peak of the Mn⁴⁺-activated fluoride phosphor and the absorption peak of chlorophyll rarely overlap, while the emission peak of the Mn⁴⁺-activated oxide phosphors, such as Sr₄Al₁₄O₂₅:Mn⁴⁺, overlap with the absorption peaks of chlorophyll-a and chlorophyll-b.

As shown in Fig. 2, the emission spectrum of Sr₄Al₁₄O₂₅:Mn⁴⁺ red phosphor has overlap more efficiently with the absorption spectrum of chlorophyll compared with Sr₂Si₃N₄:Eu²⁺ red commercial phosphors. However, most Mn⁴⁺-activated oxide phosphors cannot be excited effectively by blue light, which limited its application on blue chip-based LEDs. There are some methods to improve the luminous efficiency and luminous intensity of Mn⁴⁺-activated oxide phosphors, such as impurity doping with Mg²⁺ ions, impurity doping with Bi³⁺ ions, and impurity doping with Na⁺ ions. The objective of this work is to develop a red emitting phosphor which can match with blue chip for possible application in plant growth. In this article, the red phosphors Sr₄Al₁₄O₂₅:Mn⁴⁺, Na⁺, B³⁺ with strong absorption at blue region were synthesized. The difference with
Sr₄Al₁₄O₂₅:Mn⁴⁺, Na⁺ phosphors reported by Lili Meng was that the excitation intensity of the phosphor we prepared was significantly improved, especially in the blue light region. We also found that the doping amount of B³⁺ ions was very crucial. Specific doping ratio of B³⁺ ions and Na⁺ ions make the luminous performance of the phosphor significantly improved. The advancements of current work include significant improvement of luminescent efficiency of Sr₄Al₁₄O₂₅:Mn⁴⁺ by doping B³⁺ and Na⁺ ions. Notably, doping of B³⁺ and Na⁺ ions can improve its visible light excitation efficiency in the spectral range of 400–500 nm so that it can be incorporated as a red component into blue chip-based LED applications for plant growth.

Experimental section

Materials and synthesis
Polycrystalline phosphors with composition of Sr₄Al₁₄O₂₅:Mn⁴⁺, Na⁺, B³⁺ were prepared with a high-temperature solid-state reaction. Briefly, the constituent raw materials SrCO₃ (A. R., 99.9%), Al₂O₃ (A. R., 99.9%), Na₂CO₃ (A. R., 99%), H₃BO₃ (A. R., 99%) and MnO₂ (A. R., 99.9%) were weighed according to the stoichiometric ratio. Individual batches of 10 g were weighed according to the designed stoichiometry and mixed homogeneously with the same mass of absolute ethyl alcohol as the dispersant. After planetary ball-milling process, the obtained homogeneous slurry was placed in a Petri dish and dried in an oven. Then, the dried mixtures were put into a crucible with a lid and heated in a tubular furnace at 1400 °C for 6 hours in the air. When cooled down to room temperature, the prepared phosphors were crushed and ground for subsequent measurements.

Characterization
All crystal structure compositions were checked for phase formation by using powder X-ray diffraction (XRD) analysis with a Rigaku X-ray diffractometer (Tokyo, Japan) with a graphite
Results and discussion

Microstructure

The compositions of the typical Sr₄Al₁₄O₂₅:0.014Mn⁴⁺, xB³⁺, 0Na⁺, (0 ≤ x ≤ 1.6) samples were displayed in Fig. 3a. When no flux is added, the compound Sr₄Al₁₄O₂₅ is not formed, and instead, the two phases SrAl₂O₄ (JCPDS-No. 34-0379) and SrAl₁₂O₁₉ (JCPDS-No. 80-1195) appear. When boric acid is added, for instance at x = 0.2, the sample turns into a single phase of Sr₄Al₁₄O₂₅. With increase of x from 0.2 to 1.2, XRD patterns of phosphors agree well with that of standard Sr₄Al₁₄O₂₅ (JCPDS-No. 52-1876). Further increment of boric acid content will induce an extra impurity phase of SrAl₂O₄. Because boric acid has lower melting temperature, it (a B³⁺ dopant content in Sr₄Al₁₄O₂₅:Mn⁴⁺, xB³⁺, 0Na⁺ (0 ≤ x ≤ 1.6) phosphors. (b) XRD patterns of Sr₄Al₁₄O₂₅:0.014Mn⁴⁺, 0.8B³⁺, yNa⁺ (0 ≤ y ≤ 3). The reference spectra of Sr₄Al₁₄O₂₅ (JCPDS-No. 52-1876), SrAl₁₂O₁₉ (JCPDS-No. 80-1195), SrAl₂O₄ (JCPDS-No. 34-0379) standard patterns were shown at the bottom.

Table 1  Comparison of B³⁺ dopant content in Sr₄Al₁₄O₂₅:Mn⁴⁺, xB³⁺

| Range of x   | Optimum value of x | Ref. |
|-------------|--------------------|-----|
| ---          | 0.7                | 15  |
| 0–1.6        | 0.8                |     |
| 0–3.5        | 0.7                | 15  |
| ---          | 0.54               | 14  |
| ---          | 0.4                | 12  |

Fig. 4 shows a schematic of the Sr₄Al₁₄O₂₅ crystal structure. The space group of Sr₄Al₁₄O₂₅ is Pmma and orthorhombic. The networks of orthorhombic Sr₄Al₁₄O₂₅ are built by one layer of the octahedral anion groups (AlO₆) and several layers of tetrahedral anion groups (AlO₄) alternatively. There are three types of AlO₆ octahedrons [Al(1)O₆, Al(5)O₆ and Al(6)O₆], and three types of AlO₄ tetrahedron [Al(1)O₄, Al(2)O₄ and Al(3)O₄]. Compared with the strong covalence effect of the AlO₄ tetrahedron, a little weak polarization field of the AlO₆ octahedron is more suitable for Mn⁴⁺ incorporating. In addition, the Mn⁴⁺ ion always experiences a strong CF due to its high effective positive charge with the result that the emission spectrum is always dominated by the sharp emission line corresponding with the spin-forbidden 2E_g → 4A_2g transition.

Luminescence property

Fig. 5a show the PLE spectrum of Sr₄Al₁₄O₂₅:0.8B³⁺, 2Na⁺, 2Mn⁴⁺ (0.05% ≤ z ≤ 0.5%) phosphor monitored at 652 nm. Fig. 5b show the PL spectrum of Sr₄Al₁₄O₂₅:0.8B³⁺, 2Na⁺, 2Mn⁴⁺ (0.05% ≤ z ≤ 0.5%) phosphor monitored at 652 nm. Fig. 5c show the PL spectrum of Sr₄Al₁₄O₂₅:0.8B³⁺, 2Na⁺, 2Mn⁴⁺ (0.05% ≤ z ≤ 0.5%) phosphor monitored at 652 nm. Fig. 5d show the PL spectrum of Sr₄Al₁₄O₂₅:0.8B³⁺, 2Na⁺, 2Mn⁴⁺ (0.05% ≤ z ≤ 0.5%) phosphor monitored at 652 nm. Fig. 5e show the PL spectrum of Sr₄Al₁₄O₂₅:0.8B³⁺, 2Na⁺, 2Mn⁴⁺ (0.05% ≤ z ≤ 0.5%) phosphor monitored at 652 nm.
$z \leq 0.5\%$) phosphor excited at 450 nm. Fig. 5c and d visually show the excitation and emission intensity of $\text{Mn}^{4+}$ on dopant concentration for $\text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.8\text{B}^{3+}, 2\text{Na}^+$, $z\text{Mn}^{4+}$ phosphor, respectively. As we can see, the optimum molar concentration of $\text{Mn}^{4+}$ in $\text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.8\text{B}^{3+}, 2\text{Na}^+$, $z\text{Mn}^{4+}$ in this work was $0.1\%$. The photoluminescence emission spectrum of the phosphor presented a double-peak structure between 600 and 700 nm with two strong bands at about 654 and 664 nm, which were attributed to the $^1\text{E} \rightarrow ^3\text{A}_2$ transition of $\text{Mn}^{4+}$ ions and a phonon sideband transition, respectively.$^{13}$

After obtained optimum $\text{Mn}^{4+}$ doping concentration, we adjusted the amount of $\text{B}^{3+}$ and $\text{Na}^+$. Fig. 6a and b show the photographs of $\text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, x\text{B}^{3+}, y\text{Na}^+ (0 \leq x \leq 1.6, \ 0 \leq y \leq 3)$ under natural sunlight and 365 nm UV light, respectively. Along with Na$^+$ ions and B$^{3+}$ doping concentrations increasing, the color of $\text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, x\text{B}^{3+}, y\text{Na}^+$ phosphors under natural sunlight change from light yellow to bright yellow. This might be ascribed to the Na$^+$ and B$^{3+}$ ions enhanced absorption band (220–500 nm), especially in the region of 400–500 nm, as illustrated in Fig. 7a.

In detail, Fig. 7a shows PLE spectrum of $\text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, x\text{B}^{3+}, y\text{Na}^+$ phosphor monitored at 652 nm. In order to compare the relative changes of the excitation bands, the PLE intensity is normalized. When no Na$^+$ is added, with $x$ of B$^{3+}$
increases from 0.2 to 1.6, the relative magnitude of excitation bands varies little. Furthermore, the optimum $x$ of $B^{3+}$ is 0.4, not 0.8, according to Fig. 7b. When Na$^+$ is added, for instance at $y = 1$, the intensity of the blue excitation band (400–500 nm) has been dramatically increased, as shown in Fig. 7a. In Fig. 7b, we can see that the optimum $x$ of $B^{3+}$ turns to 0.8 from 0.4. When $x$ of $B^{3+}$ is fixed at 0.8, the emission intensity increased first, reached the maximum ($y = 2$), and then decreased with the increase of Na$^+$ content. Cross experiment can clearly found that the optimal combination is $x = 0.8, y = 2$. To illustrate the difference between our experimental results and Meng’s,$^{12}$ the detail differences are listed in Table 2. As we can see the amount of doping Na$^+$ and B$^{3+}$ is observably different. As Fig. 7a shown, when $x = 0.4$, the addition of Na$^+$ does not significantly affect the relative intensity between the two excitation bands. This is why Meng didn’t report the asynchronous increase of excitation bands. This asynchronous increase of excitation bands, however, allows the phosphor to be more easily excited by blue light. As Fig. 7b shows, the PL emission intensity excited by 450 nm blue light sharply increased 3 times when $x$ of $B^{3+}$ turns to 0.8 from 0.4 (when $y = 2$). Therefore, addition of $B^{3+}$ and Na$^+$ ions in Sr$_4$Al$_{14}$O$_{25}$:0.014Mn$^{4+}$, xB$^{3+}$, Na$^+$ can significantly improve its visible light excitation efficiency in the spectral range of 400–500 nm so that it can be incorporated as a red component into blue chip-based LED applications for plant growth.

Fig. 8a and b show the photoluminescent excitation and emission spectra of the Sr$_4$Al$_{14}$O$_{25}$:0.014Mn$^{4+}$, 0.8B$^{3+}$ based phosphors with or without co-incorporating Na$^+$. The fluorescent intensities of the phosphors excited at 450 nm reached a maximum at $x = 0.8$, $y = 2$ and $z = 0.014$, and the strongest emission intensity of Sr$_4$Al$_{14}$O$_{25}$:0.8B$^{3+}$, 2Na$^+$, 0.014Mn$^{4+}$ sample were increased by 700% compared with Sr$_4$Al$_{14}$O$_{25}$:0.8B$^{3+}$, 2Na$^+$, 0.014Mn$^{4+}$ without Na$^+$ co-doping. The photographs of the Sr$_4$Al$_{14}$O$_{25}$:0.014Mn$^{4+}$, 0.8B$^{3+}$, 2Na$^+$ sample and Sr$_4$Al$_{14}$O$_{25}$:0.014Mn$^{4+}$, 0.8B$^{3+}$ sample exposed to 450 nm blue light and 365 nm UV light are shown in the insert of the Fig. 8a and b. After the incorporating of sodium ions, the brightness of phosphor becomes larger. We can find that the excitation spectra are consists of three conjoint bands by multi-peaks fitting, which located from near UV region to visible blue region. These three bands are corresponding to $^4A_2 \rightarrow ^2T_2$, $^4A_2 \rightarrow ^2T_2$ and $^4A_2 \rightarrow ^4T_1$ transition, respectively. Though according to the spin selection rule of $\Delta S = 0$, the transitions between $^4T_2$, $^4T_1$ and ground $^4A_2$ levels are spin-allowed, the spin-forbidden transition $^4A_2 \rightarrow ^2T_2$ is still be found in our result and other Mn-incorporated phosphors, such as CaAl$_{12}$O$_{20}$:Mn$^{2+}$,$^{17}$ SrMgAl$_{10}$O$_{17}$:Mn$^{2+}$,$^{18}$ Mg$_2$TiO$_4$:Mn$^{2+}$,$^{19}$ Li$_2$Mg$_2$SnO$_6$:Mn$^{2+}$,$^{20}$ Ba$_2$TiGe$_2$O$_8$:Mn$^{2+}$ and Na$_2$MgAl$_{12}$O$_{27}$:Mn$^{2+}$.$^{21}$ By fitting the peaks of the excitation spectra, we found that all the excitation peaks have a red shift as the sodium incorporated.

Crystal filed strength calculation

The values of $D_{Q1}$, $B$ and $C$ can be calculated based on experimentally determined energy levels using the following equations: $^3$
\[ D_q = \frac{E(4T_2g - 4A_2g)}{10} \]  
\[ \frac{D_q}{B} = \frac{15(x - 8)}{(x^2 - 10x)} \]  
\[ x = \frac{E(4A_2g \rightarrow 4T_1g) - E(4A_2g \rightarrow 4T_2g)}{D_q} \]  
\[ E(\frac{2T_1}{C_0} - \frac{2E_g}{C_0}) = \frac{66B^2}{(10D_q)} \]  
\[ E(\frac{2A_1}{C_0} - \frac{4A_2}{C_0}) = 10D_q + 4B + 3C \]  
\[ E(\frac{4T_1(P)}{C_0} - \frac{4A_2}{C_0}) = 15D_q + 7.5B - 0.5[100D_q^2 - 180D_qB + 225B^2]^{1/2} \]

Where \( D_q \) represents the crystal field strength and the parameter \( x \) is defined as:

\[ x = \frac{E(4A_2g \rightarrow 4T_1g) - E(4A_2g \rightarrow 4T_2g)}{D_q} \]

From Fig. 8a, the energy levels of \( 4T_2, 4T_1 \) and \( 2E_g \) in the \( \text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, 0.8\text{B}^{3+}, 2\text{Na}^+ \) host were determined at 22 124, 28 986 and 15 314 cm\(^{-1}\), respectively. From Fig. 8b, the energy levels of \( 4T_2, 4T_1 \) and \( 2E_g \) in the \( \text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, 0.8\text{B}^{3+} \) host were determined at 22 321, 30 769 and 26 667 cm\(^{-1}\), respectively. From Eqn (1)–(4), the crystal field parameters of \( D_q, B, C \) in the \( \text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, 0.8\text{B}^{3+} \) were calculated to be 2212, 644 and 3735 cm\(^{-1}\) respectively. The crystal field parameters of \( D_q, B, C \) in the \( \text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, 0.8\text{B}^{3+} \) were calculated to be 2232, 830 and 3540 cm\(^{-1}\) respectively. Once these parameters have been determined, the energies of all other states such as \( 2T_1, 2A_1 \) and \( 4T_1 \) can be theoretically predicted by:

\[ E(\frac{2T_1}{C_0} - \frac{2E_g}{C_0}) = \frac{66B^2}{(10D_q)} \]

\[ E(\frac{2A_1}{C_0} - \frac{4A_2}{C_0}) = 10D_q + 4B + 3C \]

\[ E(\frac{4T_1(P)}{C_0} - \frac{4A_2}{C_0}) = 15D_q + 7.5B - 0.5[100D_q^2 - 180D_qB + 225B^2]^{1/2} \]

The crystal field parameters and the energies of states in \( \text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, 0.8\text{B}^{3+}, 2\text{Na}^+ \) and \( \text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, 0.8\text{B}^{3+} \) crystal lattices are summarized in the Table 3. As shown in Fig. 8c, the dependence of energy levels of \( \text{Mn}^{4+} \) on crystal field strength can be illustrated by Tanabe–Sugano energy diagram. The \( 2E_g \) levels are almost parallel to the ground state \( 4A_2 \), which results that the location of the emission peak is difficult to be influenced by crystal field strength. While, the energy gap between \( 4T_1 (or 4T_2) \) levels and ground state \( 4A_2 \) can be changed by variation of the crystal field strength. The electron transition schematic diagrams are shown in the Fig. 8c with blue and green dot lines. The value of \( D_q/B \) increased to 3.43 from 2.69 as the \( \text{Na}^+ \) ions addition. There is an increase in the asynchronous increases of the near UV and visible absorption bands. The increase of the excitation

Table 2  Comparison of dopant content in \( \text{Sr}_4\text{Al}_{14}\text{O}_{25}:x\text{B}^{3+}, y\text{Na}^+, z\text{Mn}^{4+} \)

| Range of x | Optimum value of x | Range of y | Optimum value of y | Optimum value of z | Excitation bands | Ref. |
|-----------|-------------------|-----------|-------------------|-------------------|-----------------|------|
| 0.4       | —                 | 0.05      | 0–0.09            | 0.01              | Synchronous increase | 11   |
| 0.8       | 0–1.6             | 2         | 0–3               | 0.014             | Asynchronous increase | This work |

Fig. 8  (a) PL (EX = 450 nm) and PLE (EM = 652 nm) spectra of \( \text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, 2\text{Na}^+, 0.8\text{B}^{3+} \) and (b) \( \text{Sr}_4\text{Al}_{14}\text{O}_{25}:0.014\text{Mn}^{4+}, 0.8\text{B}^{3+} \) respectively; (c) Tanabe–Sugano energy diagram of a 3d\(^5\) system in an octahedral crystal field; (d) configurational coordinate diagram for \( \text{Mn}^{4+} \) ions in \( \text{Sr}_4\text{Al}_{14}\text{O}_{25} \) hosts.
intensity at visible blue region is much larger than at near UV region, even both excitation intensity at these two regions are almost equal. Na⁺ compounds are well known fluxes in solid state synthesis. However, the shapes of the excitation bands cannot be changed by fluxes and meanwhile the redshift of the excitation is a change on the luminescent mechanism instead of fluxes effect.

Fig. 8d shows a schematic diagram of a process of photo-luminescence. The 2Eg → 2T1, 2T2 and 4A2 levels are derived from the t2g electronic orbital, whereas the 4T1 and 4T2 levels are formed from another t2g orbital, resulting a displacement between the parabolas of ground state 4A2 and 4T1 (or 4T2). The electrons are excited from to ground state 4A2 to 4T1, 4T2 or 2T2 levels by radiation. Then, the excited electrons usually relax non-radiatively to 2Eg followed by the spin-forbidden 2Eg → 4A2 transition characterized by wide emission bands.

Conclusions

A series of Sr₄Al₄O₁₂:xB⁺, yNa⁺, zMn⁴⁺ red phosphors were synthesized by a high-temperature solid-state reaction method at 1400 °C for 6 hours in the air. The fluorescent intensities of the phosphors excited at 450 nm reached a maximum at x = 0.8, y = 2 and z = 0.014, and the strongest emission intensity of Sr₄Al₄O₁₂:0.014Mn⁴⁺, 0.88B⁺, 2Na⁺ sample was increased by 700% compared with Sr₄Al₄O₁₂:0.88B⁺, 0.014Mn⁴⁺ without Na⁺ co-doping. In comparison with Mn⁺⁺ single incorporated phosphor, Sr₄Al₄O₁₂:0.014Mn⁴⁺, 0.88B⁺, 2Na⁺ shows greater advantage of promising application incorporated as a red component into blue chip-based LED for plant growth because of much stronger absorption at blue light region and enhanced red emission. The prepared phosphors could be efficiently excited by both near-UW light and the commercially available blue light of LED chips at 450 nm.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was financially supported by the programs of the National Key Research and Development Program of China (2017YFB0403200).

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