



Cite this: *RSC Adv.*, 2019, 9, 9244

Enhanced photoluminescence and energy transfer performance of $Y_3Al_4GaO_{12}:Mn^{4+}, Dy^{3+}$ phosphors for plant growth LED lights†

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Plant growth LEDs have attracted broad attention in modern society, desperate for specific phosphors with a characteristic emission band. A novel Mn^{4+} and Dy^{3+} co-doped $Y_3Al_4GaO_{12}$ phosphor were successfully prepared through a conventional high-temperature solid-state reaction. A three band emission including red (625–700 nm), orange (550–607 nm) and blue (462–490 nm) is observed in these phosphors when excited under a near-UV lamp, which is ascribed to ${}^2E \rightarrow {}^4A_2$ of Mn^{4+} , ${}^4F_{9/2} \rightarrow {}^6H_{15/2}$ and ${}^4F_{9/2} \rightarrow {}^6H_{13/2}$ of Dy^{3+} , respectively. The three emissions match the absorption spectra of chlorophyll A and chlorophyll B well. Meanwhile, the energy transfer from Dy^{3+} to Mn^{4+} was confirmed by the luminescence spectra and lifetime analysis. Finally, an LED device was fabricated that consisted of a 365 nm ultraviolet chip and the $Y_3Al_4GaO_{12}:Mn^{4+}, Dy^{3+}$ phosphor. The excellent properties indicate that the synthesized phosphor has a promising application in the optical agricultural industry.

Received 26th January 2019

Accepted 12th March 2019

DOI: 10.1039/c9ra00700h

rsc.li/rsc-advances

1. Introduction

Optical agriculture has been brought to the forefront in the recent years, for providing plants with necessary light. As we know, plant growth needs four main conditions including light, atmosphere, water and nutrients, and light plays crucial roles among the four factors during all the process of plant growth.¹ According to the previous experiments of plant cultivation, the effective emission ranges are orange-red (550–700 nm) and blue (400–500 nm) bands. Plants absorb orange-red and blue light by the photosynthesis pigments of chlorophyll and phytochrome.^{2,3} The latest way to promote plant growth is using LEDs to supply light in optical agricultural. Generally speaking, there are two main methods to obtain plant growth LEDs: “blue LED chip + red LED chip” and “phosphor + LED chip”. For the first technology, many shortcomings such as high cost, complex control circuit, high light failure and color drift limit its further application. The main problem is that two different monochromatic semiconductor chips are used, which require different drive circuits to control the emitting light ratio of blue

and red dynamically. The “phosphor + LED chip” technology can be divided into two forms for different types of chip. The first method is combined a blue-emitting InGaN chip with red emission phosphors such as $(Sr,Ca)AlSiN_3:Eu^{2+}$ ⁴ and $K_2TiF_6:Mn^{4+}$.⁵ This technology has the advantages of simple operate and low cost. However, the inconsistency of light wane and color drift exist between blue chip and red phosphor. On the one hand, chromaticity shifts in light-emitting diode (LED) devices arise from multiple mechanisms that produce chemical changes in the materials used to construct the LED devices. The chromaticity shifts in LED devices usually start with a fast-acting component which quickly reaches its maximum value, followed by one or more slower acting component. The fast-acting and slower acting component can be modeled with a bounded exponential function and a generalized logistic function.⁶ On the other hand, light wane because of the luminescence decay of phosphors in severe conditions such as high temperature, and LED device always reaches high temperature in long time work. What's more, it is difficult to guarantee the combination between the chip and phosphors, which weaken the light without effective excitation. The second approach is combined near-ultraviolet (n-UV) chips with red and blue phosphors or single-matrix red and blue dual emission phosphor, but this strategy depends more on the characteristics of phosphors and put forward higher requirements on phosphors. However, most traditional phosphors cannot match the aimed emission band, thus more accurate light sources are needed to provide the special light in the plant growth process.^{7–9}

The current primary red-emitting phosphors are Eu^{2+} -doped nitrides, such as $M_2Si_3N_8:Eu^{2+}$ ^{10,11} and $MAISiN_3:Eu^{2+}$ ($M = Ca,$

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9ra00700h

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Sr, Ba).^{4,12,13} They usually are synthesized in critical requirements (0.9 MPa N₂ pressure and 1800 °C), the high cost limits their utility in agricultural industry. As an alternative, Mn⁴⁺-doped oxides^{14–17} catch the public's attention due to its attractive luminescence properties of high-efficacy, low cost, high stability and eco-friendliness. Many strategies have been tried out to enhance the photoluminescence emission of Mn⁴⁺-doped phosphors, such as matrix solid solution modification,^{18–20} co-doped charge compensator^{21,22} or fluxing agent,^{23,24} and energy transfer from sensitizer to activator.^{25–27}

Solid solution treatment is a common method in phosphor modification with many advantages such as increase lattice distortion and lattice rigidity. Some materials change phase accompanied by a large volume contraction at high temperature reaction which have damage to their luminescence properties. This phenomenon can be reduced by the addition of solid solution ions. There is no crystal transformation in the formation of solid solution, but volume effect was reduced which can enhance the resistance of phosphors to high temperature. After the formation of solid solution, there are some distortions on the lattice structure which is in a high energy activated state, and it is conducive to chemical reaction. Therefore, solid solution treatment can improve the luminescence performance and thermal stability.^{28–30} There are many conditions for solid solution formation, such as similar ion radius size, alike crystal structure or chemical formula, similarity of ion electronegativity, and the same of ion value. For example, Ca₁₄Al₁₀Zn₆O₃₅³¹ and Ca₁₄Ga₁₀Zn₆O₃₅³² has the similar structure, Mn⁴⁺-doped Ca₁₄Al₁₀Zn₆O₃₅ phosphors show far-red emitting, and the luminescence properties can be improved by substituting Ga³⁺ for Al³⁺ ion.³³ The similar result was found in Mn⁴⁺ or Ce³⁺ doped (Y,Lu)₃(Al,Ga)₅O₁₂ system.^{22,34,35}

Another important way to enhance luminescence properties is co-doped charge compensator in same matrix. For instance, the luminescence performance of Y₃Al₅O₁₂:Mn⁴⁺ red phosphor can be enhanced by co-doping Ge⁴⁺, Sc³⁺, Ca²⁺, Mg²⁺, Sr²⁺, Li⁺, Na⁺ ions.^{21,33,36} In addition, aluminate phosphors are mainly compounded *via* high-temperature solid-state reaction in traditional production, which requires high sintering temperature. In order to reduce the reaction temperature and surface defects of the product, and increase the crystallinity of the product, fluxing agent often needs to add in the raw material such as BaF₂, NaF, NH₄Cl and H₃BO₃.³⁵ The third method to improve luminescence performances is to use energy transfer effect, such as Ca₁₄Al₁₀Zn₆O₃₅:Ti⁴⁺, Mn⁴⁺,³¹ Ca₃Al₄ZnO₁₀:Bi³⁺, Mn⁴⁺,²⁶ Gd₂ZnTiO₆:Mn⁴⁺, Yb³⁺,³⁷ and so on.

As we know, the Mn⁴⁺ doped Y₃(Al,Ga)₅O₁₂ system show a wide range of 225–575 nm excitation spectrum. The Dy³⁺ can exhibit blue 480 nm (⁴F_{9/2}–⁶H_{15/2}) sharp band and orange 585 nm (⁴F_{9/2}–⁶H_{13/2}) luminescence performance in Y₃(Al,Ga)₅O₁₂ matrix for n-UV excitation.^{38–41} Based on the resonance-type energy transfer theory, we assume that Dy³⁺ could improve the emission intensity of Mn⁴⁺ in Y₃(Al,Ga)₅O₁₂ matrix. Energy transfer from Dy³⁺ to Mn⁴⁺ has been reported in several works. Chen *et al.*⁴¹ and Zhou *et al.*³² verified energy transfer from Dy³⁺ to Mn⁴⁺ in Ca₁₄Al₁₀Zn₆O₃₅ and Ca₁₄Zn₆Ga₁₀O₃₅ phosphors respectively, which indicate that it is

possible for the occurrence of similar energy transfer in Y₃(Al,Ga)₅O₁₂ matrix. Herein, we intend to synthesize a novel Dy³⁺ and Mn⁴⁺ co-doped Y₃(Al,Ga)₅O₁₂ phosphor which show both blue and red luminescence for plant growth LEDs.

In this study, a novel Y₃Al₄GaO₁₂:Mn⁴⁺,Dy³⁺ (YAGO:Mn⁴⁺,Dy³⁺) phosphor was prepared *via* conventional solid-state method at high-temperature in air. Samples as-obtained show bright red, orange and blue light emission, which match the absorption of plant pigments in a large extent. The crystal structure, optical properties and luminescence lifetime are researched in detail. All measurements indicated that the YAGO:Mn⁴⁺,Dy³⁺ phosphors is promising to applicate in plant growth LEDs.

2. Experimental work

2.1 Preparation of samples and device

A series of Y₃Al₄GaO₁₂:Mn⁴⁺ (YAGO:Mn⁴⁺), Y₃Al₄GaO₁₂:Dy³⁺ (YAGO:Dy³⁺) and Y₃Al₄GaO₁₂:Mn⁴⁺,Dy³⁺ (YAGO:Mn⁴⁺,Dy³⁺) phosphors were synthesized *via* high-temperature solid-state method. The chemical reagents were Y₂O₃ (99.999%), Ga₂O₃ (99.999%), Al₂O₃ (99.999%), Dy₂O₃ (99.999%), Mn(NO₃)₂ (AR), MgO (AR), H₃BO₃ (AR) and LiF (99.99%), all raw materials bought from Aladdin without further purification. Simultaneously, 5 w% H₃BO₃ (AR) and LiF acted as a flux. First of all, stoichiometric amounts of the raw material were mixed up adequately with a moderate amount of alcohol and ground thoroughly for 30 min. Then put the powder into alumina crucibles, fired at 1500 °C for heating rate at 5 °C min⁻¹ in ambient atmosphere and last for 4 hours. Finally, the products were cooled to room temperature naturally and then ground for further used. The LED devices were fabricated by using the Y₃-Al₄GaO₁₂:Mn⁴⁺,Dy³⁺ phosphors and 365 nm near-UV chips.

2.2 Characterization

X-ray powder diffraction (XRD) analysis was tested through a Rigaku D/SHIMADZU-6000 X-ray diffractometer with Cu-K α radiation ($\lambda = 1.5406 \text{ \AA}$), operating at 40 kV and 40 mA, setting the 2θ from 10° to 80°. The photoluminescence excitation (PLE) and emission (PL) spectra were measured by F-7000 Spectrophotometer (Hitachi, Japan) equipped with a 150 W Xe lamp. For the measurement of UV-vis absorption spectra, U-3310 spectrophotometer (Hitachi, Japan) was used. The decay curves and photoluminescence quantum efficiency were measured under excitation at 366 nm *via* FLS980 fluorescence spectrometer (Edinburgh, UK) using a millisecond flash lamp. All the characterizations obtained at room temperature except the temperature-dependent PL spectra.

3. Result and discussion

3.1 Structure characterization and phase identification

The phase composition characterized by X-ray diffraction (XRD). Fig. 1(a) presents the XRD patterns of the Y₃Al_{5–m}Ga_mO₁₂:0.004-Mn⁴⁺ ($m = 0–5$). With the doping concentration of Ga³⁺ increased, the curve changed from Y₃Al₅O₁₂ (PDF#72-1315) to



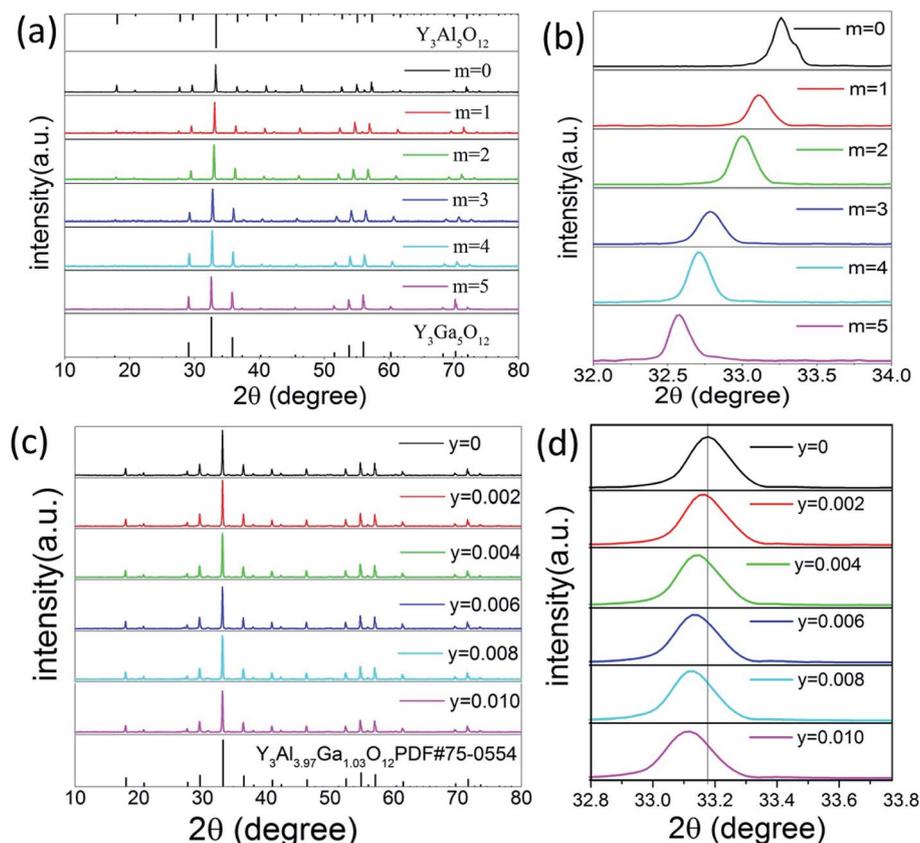


Fig. 1 (a) The XRD patterns of $Y_3Al_{5-m}Ga_mO_{12}:0.004Mn^{4+}$ ($m = 0-5$); (b) peak position change between 32.0 and 34.0° ; (c) XRD curves of $Y_3Al_4GaO_{12}:0.004Mn^{4+},yDy^{3+}$ ($y = 0-0.01$); (d) the enlarge main peak between 32.6 and 33.8° .

$Y_3Ga_5O_{12}$ (PDF#43-0512) gradually. Fig. 1(b) shows a broad peak ranging from 32° to 34° , indicating the Al site in the matrix was replaced by Ga significantly. This substitution induced the diffraction peak of these samples shifting towards low-angle direction, because Ga^{3+} has large ionic radius ($r = 0.47 \text{ \AA}$, CN = 4; $r = 0.62 \text{ \AA}$, CN = 6) than Al^{3+} ($r = 0.39 \text{ \AA}$, CN = 4; $r = 0.54 \text{ \AA}$, CN = 6).²⁶ The situation was confirmed by the following analysis of PL spectra excited at different wavelengths. The XRD patterns of $Y_3Al_4GaO_{12}:Mn^{4+},yDy^{3+}$ ($y = 0-0.010$) phosphors are shown in Fig. 1(c), which well indexed to standard card PDF#75-0554 of $Y_3Al_{3.97}Ga_{1.03}O_{12}$ phase. This result indicates $Y_3Al_4GaO_{12}:Mn^{4+},yDy^{3+}$ phosphors keep its original crystal structure without significant variation. The main peak between 32.6 and 33.8° was found to be slightly offset to low-angle in Fig. 1(d), for Dy^{3+} has larger ionic radius ($r = 1.03 \text{ \AA}$, CN = 8) than Y^{3+} ion ($r = 1.02 \text{ \AA}$, CN = 8) and Mn^{4+} ion has similar ionic radius ($r = 0.39 \text{ \AA}$, CN = 4; $r = 0.53 \text{ \AA}$, CN = 6) compared to Al^{3+} ion ($r = 0.39 \text{ \AA}$, CN = 4; $r = 0.53 \text{ \AA}$, CN = 6).⁴² This result is owing to the Dy^{3+} ion replaced Y^{3+} and Mn^{4+} replaced Al^{3+} ion in the $[YO_8]$, $[AlO_4]$ and $[AlO_6]$ site of the matrix effectively which is similar to the literature review reported by Liudmyla M. *et al.*⁴³ and Chen *et al.*³² Fig. S1(a)† presents the XRD patterns of Dy^{3+} -doped $Y_3Al_4GaO_{12}$ phosphors, and they match the standard card of $Y_3Al_{3.97}Ga_{1.03}O_{12}$ well and have the same variation tendency like Fig. 1(c).

Fig. 2 shows the unit-cell structure of $Y_3Al_{3.97}Ga_{1.03}O_{12}$ crystal (abbreviated as YAGO) and details about atomic in the

cubic octahedral structure. The unit-cell is composed of three main sites. They are tetrahedral $[AlO_4]$ octahedral $[AlO_6]$, and dodecahedral $[YO_8]$. Generally, Mn^{4+} activator likes to replace the Al^{3+} in $[AlO_6]$ site because Mn^{4+} has the similar ionic radii to Al^{3+} and octahedron are unstable compared to tetrahedron. Dy^{3+} should occupy Y^{3+} ion site in the host according to the principle of ionic radii matching,⁴⁴ which has been elucidated in the previous section. The substitution mechanism

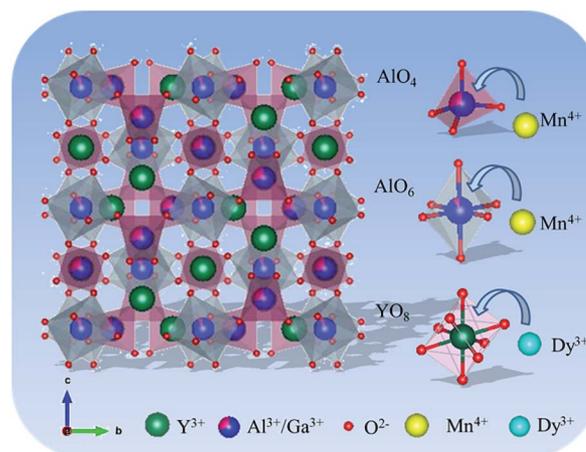


Fig. 2 Unit-cell structure of $Y_3Al_{3.97}Ga_{1.03}O_{12}$ crystal and substitution mechanism among different ion in the phosphors.



demonstrates that Dy^{3+} replaced Y^{3+} on dodecahedral site $[\text{YO}_8]$ and Mn^{4+} prefer to substitute Al^{3+} on octahedral $[\text{AlO}_6]$ compared to $[\text{AlO}_4]$ in the unit-cell structure of YAGO crystal.

3.2 Photoluminescence properties of $\text{Y}_3\text{Al}_5\text{O}_{12}:\text{xMn}^{4+}$ ($x = 0.002\text{--}0.010$) and $\text{Y}_3\text{Al}_{5-m}\text{Ga}_m\text{O}_{12}:0.004\text{Mn}^{4+}$ ($m = 0\text{--}5$) phosphors

Photoluminescence property is an important part to evaluate phosphors, which should be investigated in details. Fig. 3(a) and (b) show the luminescence excitation and emission spectra of $\text{Y}_3\text{Al}_5\text{O}_{12}:0.004\text{Mn}^{4+}$ phosphors. When monitoring at 674 nm, there were two broad bands of each excitation spectrum that ranged from 220 to 430 nm (labeled A) and 430 to 600 nm (labeled B), and their peak located at 366 nm and 472 nm respectively. The PLE spectrum can be fitted into five Gaussian peaks, of which the 327 nm and 370 nm peaks are contributed to the ${}^4\text{A}_2 \rightarrow {}^4\text{T}_1$ transitions of Mn^{4+} , other peaks located at 407 nm, 469 nm and 479 nm all originated from the ${}^4\text{A}_2 \rightarrow {}^4\text{T}_2$ transitions of Mn^{4+} .²⁷ Upon excitation at 366 and 472 nm, the as-synthesized phosphors show red emission ranging from 600 to 720 nm with two peaks at 648 nm and 674 nm which can be ascribed to the spin-forbidden ${}^2\text{E} \rightarrow {}^4\text{A}_2$ transition of Mn^{4+} . The emission intensity came to the maximum when the concentration of Mn^{4+} was 0.004 no matter excited at 366 nm or 472 nm, as presented in the inset of Fig. 3(b). The details about the emission spectra of $\text{Y}_3\text{Al}_5\text{O}_{12}:\text{xMn}^{4+}$ ($x = 0.002\text{--}0.010$) are shown in Fig. S3.†

Single ion doped phosphor concentration quenching model is shown in the inset of Fig. 3(a). The type of multipolar-multipolar interaction can be reflected by the formula⁴⁵ based on the Dexter's theory:

$$\frac{I}{x} = k \left[1 + \beta(x)^{\theta/3} \right]^{-1} \quad (1)$$

where k , β and θ refer to the constants for the same excitation condition, host crystal and indication of multipolar character, I and x stand for luminescence intensity and the activator concentration, respectively. Three types of dipole-dipole (d-d), dipole-quadrupole (d-q), and quadrupole-quadrupole (q-q) interactions can be speculated when θ value to 6, 8 and 10 respectively. The fitting slope is -1.4229 and the θ was calculated to be 4.2687 which is close to 6, indicating that the main quenching mechanism of $\text{YAGO}:\text{Mn}^{4+}$ phosphor is d-d interactions.^{22,29,30,32}

As shown in Fig. 3(c), it is obvious that the substitution of Ga^{3+} enhances the intensity of excitation spectra, which indicates these phosphors can be excited easily. With concentration of Ga^{3+} increasing, the intensity of photoluminescence excitation and emission spectra reached maximum when $m = 1$ then decreased. So we chose $\text{Y}_3\text{Al}_4\text{GaO}_{12}$ (abbreviated as YAGO) as matrix in the follow-up experiment.

3.3 Energy transfer in $\text{YAGO}:\text{Mn}^{4+}, \text{Dy}^{3+}$ phosphors

The emission of Mn^{4+} ion can be further improved *via* the addition of Dy^{3+} ion, as shown in Fig. 4(a). Two characteristic

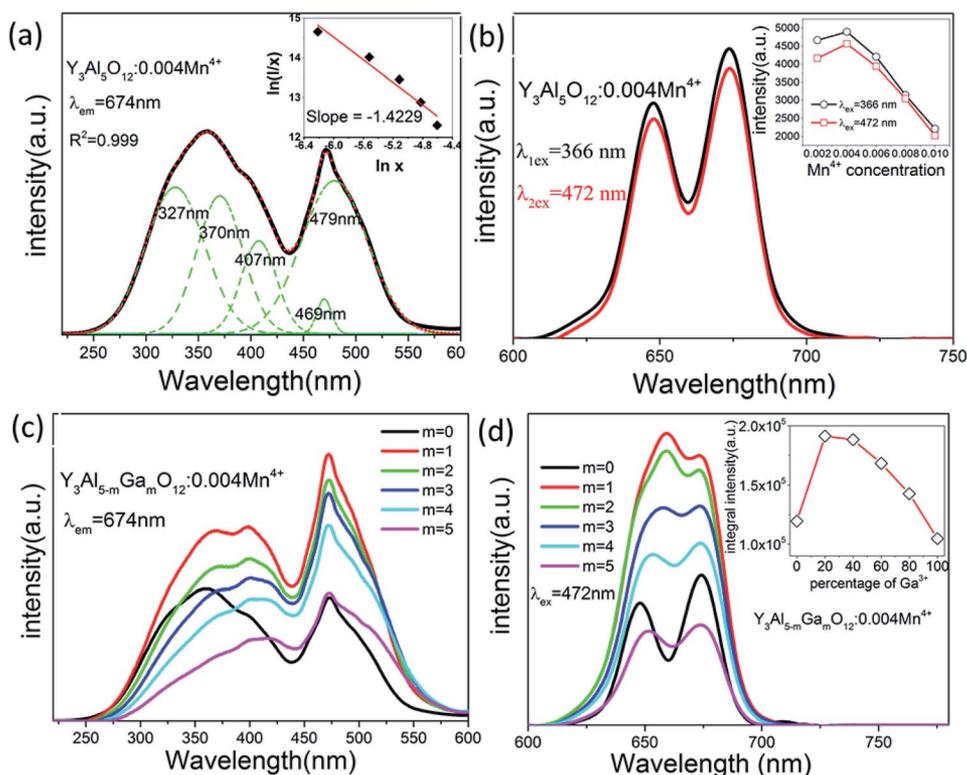


Fig. 3 (a) PLE spectrum of $\text{Y}_3\text{Al}_5\text{O}_{12}:0.004\text{Mn}^{4+}$ and fitting into five Gaussian curves, inset is the dependence of $\log(I/x)$ versus $\log(x)$; (b) PL spectra of these samples under excitation at 366 nm and 472 nm respectively, inset is the relationship between intensity and concentration of Mn^{4+} ; (c) excitation and (d) emission spectra of $\text{Y}_3\text{Al}_{5-m}\text{Ga}_m\text{O}_{12}:0.004\text{Mn}^{4+}$ ($m = 0\text{--}5$) phosphors.



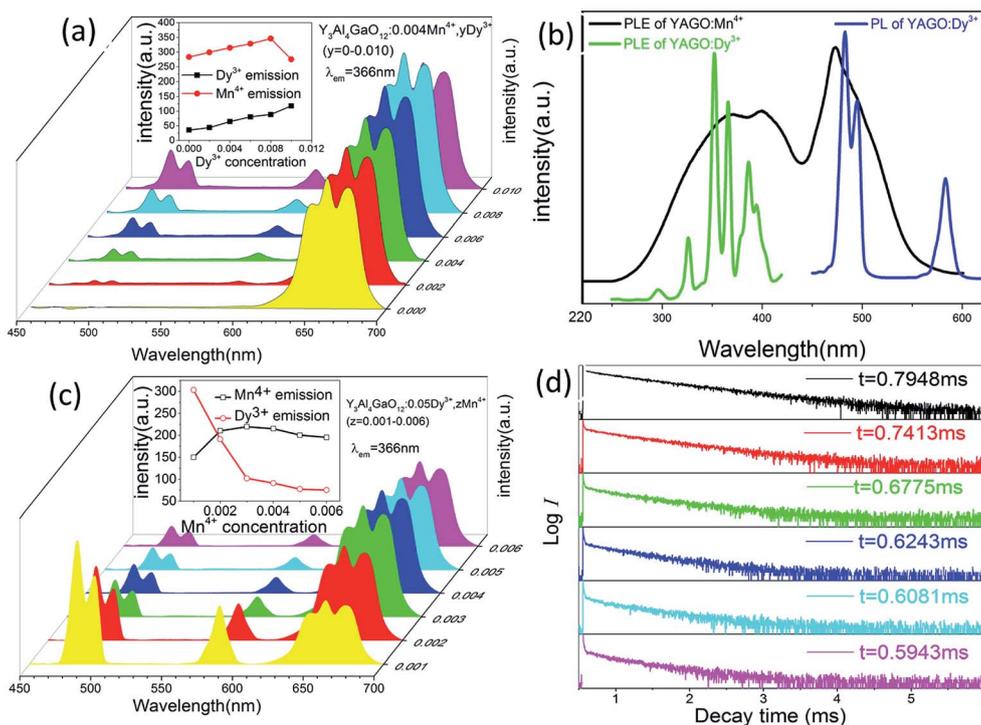


Fig. 4 (a) Photoluminescence spectra of YAGO:0.004Mn⁴⁺,yDy³⁺ (y = 0, 0.002, 0.004, 0.006, 0.008, 0.010), inset is the change of emission intensity of two ions with different Dy³⁺ concentration; (b) the excitation spectrum of YAGO:Mn⁴⁺ (black line), the PLE (green line) and PL spectra (blue line) of YAGO:Dy³⁺; (c) PL spectra of YAGO:0.05Dy³⁺,zMn⁴⁺ (z = 0.001–0.006) under excitation at 366 nm, inset is the change of emission intensity of two ions with different Mn⁴⁺ concentration; (d) decay curves of YAGO:0.05Dy³⁺,zMn⁴⁺ (z = 0.001–0.006) monitored at 483 nm under excitation at 366 nm.

emission bands can be detected at 462–490 nm and 554–607 nm in YAGO:Mn⁴⁺,Dy³⁺ phosphor under excitation at 366 nm, which corresponding to $^4F_{9/2} \rightarrow ^6H_{15/2}$ and $^4F_{9/2} \rightarrow ^6H_{13/2}$ transitions in the 4f levels of Dy³⁺ ions, respectively.⁴⁶ And red emission ranging from 600 to 720 nm should be ascribed to the $^2E \rightarrow ^4A_2$ transition of Mn⁴⁺. What's more, Fig. S2(a) and (b)† show the photoluminescence excitation and emission spectra of YAGO:Dy³⁺, the optimal Dy³⁺ doping concentration is 0.05. The excitation peaks located at 326.5, 352.5, 366.5 and 387.0 nm attribute to the intrinsic f–f transitions of Dy³⁺ from the ground state $^6H_{15/2}$ to the excited state $^4L_{19/2}$, $^6P_{7/2}$, $^6P_{5/2}$ and $^4I_{13/2}$, respectively.⁴⁶

Fig. 4(b) presents a large overlapping part between the PLE spectrum of YAGO:Mn⁴⁺ and the PL spectrum of YAGO:Dy³⁺. It is one of the indispensable factors for effective energy transfer. Therefore, it is possible for the occurrence of energy transfer from Dy³⁺ to Mn⁴⁺ ions in YAGO matrix. In order to confirm the energy transfer further, the content of Dy³⁺ ion was fixed at 0.05 and the concentration of Mn⁴⁺ was increased from 0.001 to 0.006. According to Fig. 4(c), with the concentration of Mn⁴⁺ increasing, the emission of Dy³⁺ decreased gradually while the emission of Mn⁴⁺ increased until Mn⁴⁺ concentration exceed 0.003 and then decreased due to the concentration quenching effect. Such experimental results are consistent with the prediction that there is energy transfer from Dy³⁺ to Mn⁴⁺ in YAGO host.

As mentioned in Fig. 4(c), the emission intensity of Mn⁴⁺ decrease when the content of Mn⁴⁺ exceeds the critical

concentration because of the concentration quenching phenomenon which is related to energy transfer. In generally, the critical distance (R_c) is a key parameter to evaluate the performance of sensitizer and activator, which is usually calculated by the following formula:^{47,48}

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

where V , N and x_c refer to the volume of the unit cell, the critical concentration of Mn⁴⁺ and the number of lattice sites in the unit cell that can be occupied by activators, respectively. Herein, V is 1754.05 Å³, x_c is 0.0025 and N is 8 for YAGO:Mn⁴⁺,Dy³⁺ phosphors, then the critical distance R_c was calculated to be 55.13 Å.

The PL decay curves were investigated to calculate the lifetime and then to verify the energy transfer from Dy³⁺ to Mn⁴⁺. Monitor at 483 nm for YAGO:Mn⁴⁺,Dy³⁺ phosphors under the excitation of 366 nm, the decay curves are illustrated in Fig. 4(d). These data fitted a typical single-exponential function well:²⁵

$$I = I_0 \times \exp\left(\frac{-t}{\tau}\right) \quad (3)$$

where t refer to the time and I is the luminescence intensity at that moment, I_0 and τ stand for the initial emission intensity at time 0 and the luminescence lifetime, respectively. For YAGO:0.05Dy³⁺,zMn⁴⁺ (z = 0.001–0.006) phosphors, the lifetime of Dy³⁺ decreased distinctly with the content of Mn⁴⁺ increasing. They were 0.795, 0.741, 0.678, 0.624, 0.608 and 0.594 ms when



the content of Mn^{4+} changed from 0.001 to 0.006, respectively. This result prove that energy transfer from Dy^{3+} to Mn^{4+} exist in these samples, similar conclusion in $\text{Ca}_{14}(\text{Al,Ga})_{10}\text{Zn}_6\text{O}_{35}$ matrix was reported by Zhou *et al.*³¹

Fig. 5(a) presents the UV-vis absorption spectra of YAGO, YAGO: Mn^{4+} , YAGO: Dy^{3+} and YAGO: $\text{Mn}^{4+},\text{Dy}^{3+}$. Two characteristic absorption bands were found in all four samples that name Band A (220–260 nm) and Band B (260 to 430 nm). Band C (430–550 nm) existed only in Mn^{4+} -doped phosphor, which should be attributed to the ${}^4\text{A}_2 \rightarrow {}^4\text{T}_1$ transition of Mn^{4+} . The absorption intensity enhanced by adding Dy^{3+} ion, and further enhanced by co-doping Mn^{4+} and Dy^{3+} ions.

To understand the effect of the addition of Mn^{4+} and Dy^{3+} ions on YAGO matrix, the band gap of four materials were researched, which can be calculated by the following equation according to the UV-vis absorption plots:⁴⁹

$$(\alpha h\nu)^2 = A(h\nu - E_g) \quad (4)$$

where α , h , ν and A are the absorption coefficient, the Planck constant, the frequency and constant; and E_g is the band gap energy. According to the equation and absorption dates, the relationship between the plot of $(\alpha h\nu)^2$ and $h\nu$ based on the direct gap are presented in Fig. 5(b). The value of the band gap E_g for YAGO, YAGO: Mn^{4+} , YAGO: Dy^{3+} and YAGO: $\text{Mn}^{4+},\text{Dy}^{3+}$ were calculated to be 4.12 eV, 4.10 eV, 3.15 eV and 3.06 eV, respectively. There is a great change of band-gap after Mn^{4+} -doping (from 4.12 to 3.15 eV), which can be explained by the electro-negativity of ions. In the one hand, it requires less energy for electrons to get rid of the limiting valance bonds when Mn^{4+} ion replace Al^{3+} and Ga^{3+} ion because Mn^{4+} ion has lower electro-negativity than Al^{3+} and Ga^{3+} ion, which resulting in a reduce in energy band gap.²⁶ Meanwhile, Dy^{3+} dopant could also reduce the band gap slightly for its small electro-negativity. In the other hand, the intimate addition of E_g on Mn^{4+} ions concentration indicates that the Mn^{4+} orbitals are probably involved in the conduction band besides of the band gap of YAGO.

According to Reisfeld's approximation and Dexter's energy transfer formula of multipolar interaction, the equation can evaluate interaction mechanism:^{50,51}

$$\frac{\eta_0}{\eta} = C^n \quad (5)$$

herein, C is the sum of the concentration of Mn^{4+} and Dy^{3+} , η and η_0 stand for the luminescence quantum efficiency of Dy^{3+} in

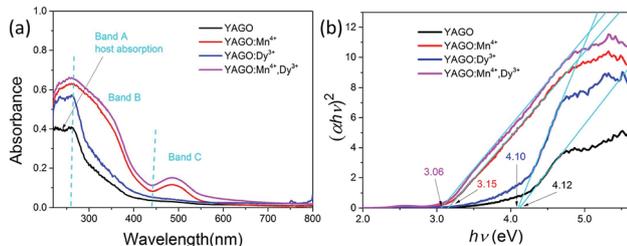


Fig. 5 (a) Comparison the UV-vis absorption spectra of YAGO phosphors with different dopants; (b) the plot of $(\alpha h\nu)^2$ versus $h\nu$ based on the direct gap of different samples.

the presence and absence of Mn^{4+} , respectively. The η_0/η is usually calculated through the ratio of initial luminescence intensity of YAGO: Dy^{3+} and YAGO: $\text{Dy}^{3+},\text{Mn}^{4+}$, and the equation can be written as $I_{\text{so}}/I_s \propto C^{n/3}$. Based on the formula, three types of interaction named dipole-dipole (d-d), dipole-quadrupole (d-p) and quadrupole-quadrupole (p-p) can be expected while $n = 6, 8$ and 10 respectively. Fig. 6 presents the relationship between I_{so}/I_s and C . When $n = 6$, linear fitting come to optimum with $R^2 = 0.998$, implying that it is dipole-dipole interaction mechanism for the energy transfer from Dy^{3+} to Mn^{4+} in YAGO phosphor.

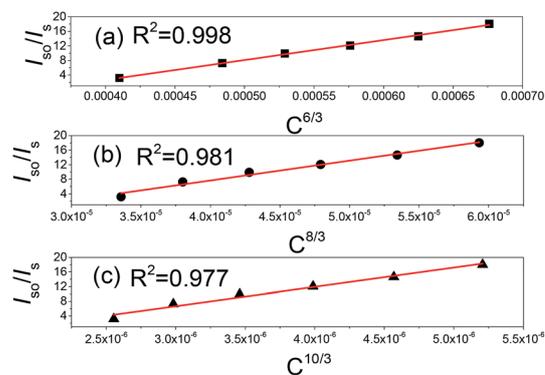


Fig. 6 Linear relationship between I_{so}/I_s and $C^{n/3}$ (a) $n = 6$, (b) $n = 8$, (c) $n = 10$.

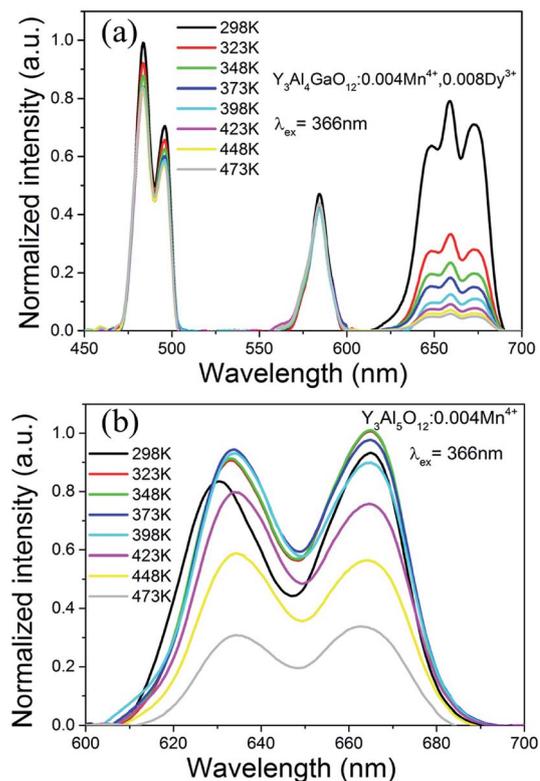


Fig. 7 Temperature-dependent emission spectra of (a) $\text{Y}_3\text{Al}_4\text{GaO}_{12}:\text{Mn}^{4+},\text{Dy}^{3+}$ and (b) $\text{Y}_3\text{Al}_5\text{O}_{12}:\text{Mn}^{4+}$ from 298 K to 473 K under excitation at 366 nm.



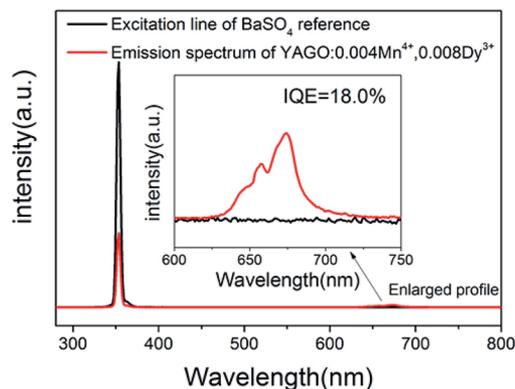


Fig. 8 Excitation line of BaSO_4 and the PL spectrum of $\text{Y}_3\text{Al}_4\text{GaO}_{12}:\text{0.004Mn}^{4+},\text{0.008Dy}^{3+}$ phosphor collected using an integrating sphere.

We tested the temperature-dependent emission spectra of $\text{Y}_3\text{Al}_4\text{GaO}_{12}:\text{0.004Mn}^{4+},\text{0.008Dy}^{3+}$ and $\text{Y}_3\text{Al}_5\text{O}_{12}:\text{0.004Mn}^{4+}$ from 298 K to 473 K in details, as shown in Fig. 7. The integral intensity of the $\text{YAGO}:\text{Mn}^{4+},\text{Dy}^{3+}$ sample kept 59.61% when the temperature came to 423 K while $\text{YAG}:\text{Mn}^{4+}$ maintained 90.10%. Though it seems decreased in the value of thermal stability, other effective emission bands for plant growth lighting were induced and keep its thermal stability well. In addition, it is possible that its thermal stability can be improved through a series of modifications in the future.

The quantum yield of $\text{Y}_3\text{Al}_4\text{GaO}_{12}:\text{0.004Mn}^{4+},\text{0.008Dy}^{3+}$ phosphor was measured under 366 nm excitation which as shown in Fig. 8, the value of IQE was calculated to be 18.0% through the following equation:⁵²

$$\eta = \frac{\int L_S}{\int E_R - \int E_S} \quad (6)$$

where η and L_S are IQE and the emission spectrum of the sample, E_S and E_R are the spectra of excitation light with sample and only with BaSO_4 reference, respectively. This IQE is at a low level likes $\text{SrLaScO}_4:\text{Mn}^{4+}$ (IQE: 12.2%)⁵³ and $\text{Ca}_{14}\text{Al}_{10}\text{Zn}_6\text{O}_{35}:\text{Mn}^{4+}$ (IQE: 19.4%)⁵⁴ which have been reported, and the property would be increased further.

For clearly describing the energy change in $\text{YAGO}:\text{Mn}^{4+},\text{Dy}^{3+}$ phosphor, the excitation, emission and energy transfer processes of the sample were shown in Fig. 9. Dy^{3+} ions were excited from ${}^6\text{H}_{15/2}$ energy level to high excited state level such as ${}^6\text{P}_{7/2}$, ${}^4\text{I}_{13/2}$, ${}^4\text{F}_{9/2}$ or even the conduction band under irradiation of the n-UV light, then relaxed to ${}^6\text{H}_{13/2}$ and ${}^6\text{H}_{15/2}$ from ${}^4\text{F}_{9/2}$ with blue and orange emission. Meanwhile, Mn^{4+} ion can be excited to ${}^4\text{T}_1$, ${}^4\text{T}_2$ level and conduction band from ${}^4\text{A}_2$ under n-UV radiation, the excited Mn^{4+} dropped to the ${}^2\text{E}$ energy level then relaxed to ${}^4\text{A}_2$ with red light emission. In this process, the energy transfer occurs from excited state level ${}^4\text{F}_{9/2}$ of Dy^{3+} ions to the excited state level ${}^4\text{T}_2$ of Mn^{4+} ions, then relaxed to the lowest excited state level ${}^2\text{E}$ of Mn^{4+} ions through non-radiative relaxation, finally came back to the ground state in the form of red emission.

3.4 LED device fabricated with $\text{YAGO}:\text{Mn}^{4+},\text{Dy}^{3+}$ and its electro-luminescent property

In order to investigate the application of the synthetic $\text{YAGO}:\text{Mn}^{4+},\text{Dy}^{3+}$ in indoor plant growth, LED devices were fabricated with $\text{YAGO}:\text{Mn}^{4+},\text{Dy}^{3+}$ phosphor and 365 nm near-UV chip. As shown in Fig. 10(a), the phosphor powder was white in daylight and presented purplish red under the ultraviolet lamp; it appeared milky white in the LED device and sent bright purplish red emission under constant current of 20 mA. The electroluminescence spectrum was mainly consisted of red, blue and little orange emission from $\text{YAGO}:\text{Mn}^{4+},\text{Dy}^{3+}$ phosphor and near ultraviolet emission from the chip, which

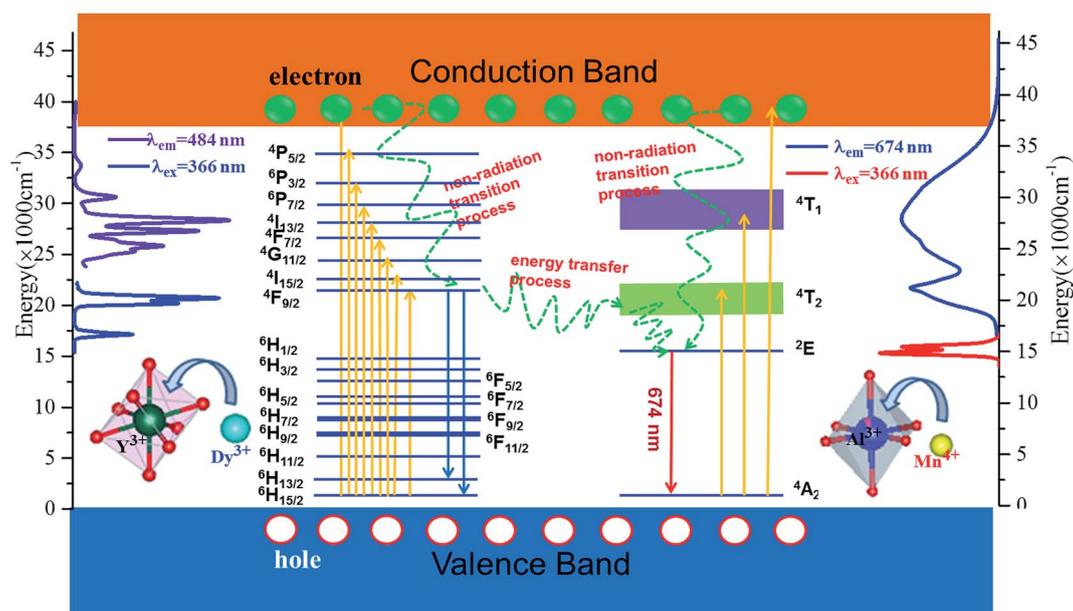


Fig. 9 Energy level, electron transitions and energy transfer schematic diagram of Dy^{3+} and Mn^{4+} in YAGO matrix.



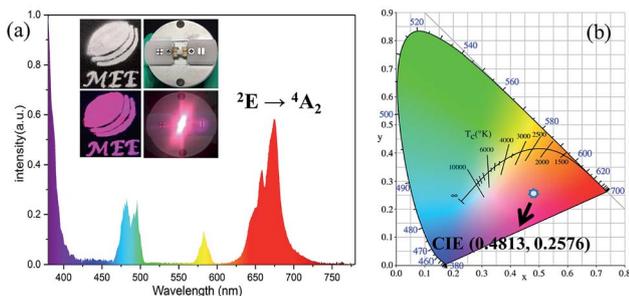


Fig. 10 (a) Electroluminescence spectrum of the as-fabricated LED device by using YAGO:0.004Mn⁴⁺,0.008Dy³⁺ phosphors and a 365 nm near-UV LED chip under a current of constant current of 20 mA. Insets are the images of LED device and phosphor powder in daylight and under a 365 nm UV lamp. (b) CIE chromaticity coordinates of the sample.

consistent with luminescence spectrum of the sample. Fig. 10(b) shows the resultant CIE coordinate value of this LED device was (0.4813, 0.2576). Other CIE coordinate values of devices that combined with different concentration of Mn⁴⁺ and Dy³⁺ co-doped YAGO phosphor are exhibited in Fig. S4,[†] they changed from purplish red to white light gradually with the increasing content of Dy³⁺. The result suggests that the novel YAGO Mn⁴⁺,Dy³⁺ phosphor have potential application in plant growth lighting and white LED lighting for their unique optical property.

4. Conclusions

To sum up, Mn⁴⁺ and Dy³⁺ co-doped YAGO phosphors were successfully synthesized *via* conventional high-temperature solid-state method. These phosphors are derived from the crystal structure of Y₃Al₅O₁₂ and changed to Y₃Al₄GaO₁₂ host, activators Mn⁴⁺ replacing Al³⁺ site while Dy³⁺ occupying Y³⁺ site in the matrix. The Y₃Al₄GaO₁₂:Mn⁴⁺,Dy³⁺ phosphors show blue, orange and red emission located at 462–490 nm, 554–607 nm and 625–700 nm respectively, which match the absorption spectrum of plant pigment well. And, these three emission bands can be ascribed to the ⁴F_{9/2} → ⁶H_{15/2} and ⁴F_{9/2} → ⁶H_{13/2} transition of Dy³⁺ and the ²E → ⁴A₂ transition of Mn⁴⁺, respectively. When the doping concentration of Mn⁴⁺ was 0.4% and Dy³⁺ was 0.8%, optimal property of these phosphors was obtained. There are many evidences for energy transfer from Dy³⁺ to Mn⁴⁺ in the as-obtained Y₃Al₄GaO₁₂:Mn⁴⁺,Dy³⁺ phosphors. Moreover, LED devices made from the phosphor have good properties. It is indicate that the as-obtained phosphors have potential applications both in optical agriculture and white LED lighting.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors would like to gratefully acknowledge funds from National Natural Science Foundation of China (Grant No.

21706060, 51703061), Natural Sciences Foundation of Hunan Province, China (Grant No. 2017JJ3103), Scientific Research Foundation of Hunan Provincial Education Department (Grant No. 17B118) and Hunan Provincial Engineering Technology Research Center for Optical Agriculture (Grant No. 2018TP2003).

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