

Red shift properties, crystal field theory and nephelauxetic effect on Mn⁴⁺-doped SrMgAl_{10-y}Ga_yO₁₇ red phosphor for plant growth LED light

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Abstract:

The discovery of novel Mn⁴⁺-doped oxide red phosphor with suitable spectrum for plant growth is a hot issue in the recent years due to the characteristic red photoluminescence of ²E_g → ⁴A₂ transition in Mn⁴⁺ ions. Generally, the emission position of Mn⁴⁺ is hard to tune because of specific crystal field in most phosphors. In this work, tunable luminescence property with obvious red shift in the spectra is observed in the Mn⁴⁺-doped SrMgAl_{10-y}Ga_yO₁₇ red phosphor via simple substitution of Ga³⁺ for Al³⁺, and crystal field theory and nephelauxetic effect are employed to explain this phenomenon. Meanwhile, the Ga³⁺ dopant changes the shape of the spectra because Ga³⁺ dopant guides the replacement site of Mn⁴⁺ and changes the luminescence center. Improved emission intensity is obtained because appropriate Ga³⁺ doping leads to larger band gap and reduces non-radiative transitions. Phosphor-converted LED (pc-LED) devices fabricated with blue chip (470 nm) and the as-obtained SrMgAl_{10-y}Ga_yO₁₇:1.0%Mn⁴⁺ phosphors emit bright blue and red light, which fit the absorption regions of plant pigments well, thus SrMgAl_{10-y}Ga_yO₁₇:Mn⁴⁺ phosphor can be a candidate for plant growth LED light.

Keywords: red shift; crystal field theory; nephelauxetic effect; $\text{SrMgAl}_{10-y}\text{Ga}_y\text{O}_{17}:\text{Mn}^{4+}$; plant growth LED light.

1. Introduction

Plant cultivation plays an important role in agricultural production. Conventional agriculture suffers from harsh environment such as frost, cloudy weather, droughts and rainstorms, resulting in a reduced yield, which cannot satisfy people's needs. In recent years, indoor plant cultivation (IPC) has caused a lot of concern for constructing a suitable and stable environment for plant growth. Light source is an essential condition in the growth process of all plants including branching, flowering, and fruiting [1-3]. Light energy can be converted into chemical energy by photosynthesis of plant pigments. Chlorophyll A, chlorophyll B, phytochrome P_R and phytochrome P_{FR} are four main plant pigments which mainly absorb blue (400-500 nm), red (600-700 nm) and far-red (700-780 nm) lights. The traditional light sources for indoor plant cultivation are incandescent lamps, high voltage halogen lamps and xenon lamps, which suffer from many disadvantages such as high energy consumption, short lifetime and spectral mismatch. Therefore, benefit from characteristic merits of energy saving, long lifetime, spectral match and environment friendliness, phosphor-converted light-emitting-diode (pc-LED) has gradually been widely used in IPC. As a light-conversion material, phosphors play an indispensable role in LED devices which directly determine the photoluminescence properties, so it is important to design and synthesize the phosphors with proper spectral emission and bright photoluminescence [4-6].

Based on this fact, many phosphors have been investigated and discussed to fulfil the light requirements of plant growth. Red phosphors are the ideal material for plant growth LED light because red emission is beneficial in promoting plant growth and biomass accumulation. Meanwhile, red phosphors also have wide application in white light-emitting diodes [7-10]. Nowadays, the commercial phosphor $\text{Y}_3\text{Al}_5\text{O}_{12}:\text{Ce}^{3+}$ widespread used in white LEDs is not suitable for plant growth LEDs because of the mismatched spectral bands [11, 12]. The main red phosphor Eu^{2+} -doped nitrides likes $(\text{Ca},\text{Sr})\text{AlSiN}_3:\text{Eu}^{2+}$ [13-15] and $(\text{Ca},\text{Sr})_2\text{Si}_5\text{N}_8:\text{Eu}^{2+}$ [16, 17] suffer from severe synthetic conditions with high temperature ($>1800^\circ\text{C}$) and high pressure in reductive atmosphere. The critical preparation requirement increases the cost of the product and limits their large-scale use in agricultural industry. Another red phosphor is Mn^{4+} -doped fluoride, such as $\text{K}_2\text{TiF}_6:\text{Mn}^{4+}$ [18] and $\text{K}_2\text{SiF}_6:\text{Mn}^{4+}$ [19]. Preparation of these materials often need HF while it is harmful to the environment, and they are unstable that easily decompose in moist environment because of the existence of $[\text{MnF}_6]^{2-}$ groups. As an alternative, Mn^{4+} -doped oxides have received extensive attention for series of advantages such as attractive photoluminescence properties, high stability, low cost and eco-friendliness [20, 21]. New types of Mn^{4+} -doped oxides such as SrLaAlO_4 [22], $\text{La}_2\text{LiSbO}_6$ [23], $\text{Gd}_2\text{ZnTiO}_6$ [24], $\text{Ca}_{14}(\text{Ga}/\text{Al})_{10}\text{Zn}_6\text{O}_{35}$ [25, 26] have been reported for plant cultivation in recent years. Mn^{4+} -activated oxide phosphors can usually be excited by commercial blue and near ultraviolet chips because they have large photoluminescence excitation spectra ranging from 220 to 580 nm, and the emission band of these phosphors is located at

the red and far-red regions from the characteristic ${}^2E_g \rightarrow {}^4A_2$ transitions of Mn^{4+} . Nonetheless, the emission position of Mn^{4+} in these matrixes is hard to tune to match the absorption of plant pigments better, so it still needs a novel matrix or method to develop red phosphors with more suitable spectrum for plant growth LED light.

Impurity doping plays a key role on the discovery of Mn^{4+} -doped oxides as red phosphors. It can be constructed easily through the substitution between elements in one main group, because these ions usually have similar ionic radius, analogical structure and electronegativity alike. Zhao *et al.* [27] and Zhou *et al.* [28] reported that the lattice site can be adjusted by replacing Ga^{3+} with Al^{3+} in $Ca_{14}Ga_{10-x}Al_xZn_6O_{35}$ phosphor, lead to an enhancement on photoluminescence intensity and quantum efficiency of the samples. Analogously, Qiao *et al.* reported a single-phase $(Ca_{9-x}Sr_x)MgK(PO_4)_7:Eu^{2+}$ phosphor with white light by adjusting the compositions and multiple activator sites, which show a significant shift and improvement on the photoluminescence spectra via Sr^{2+} replace Ca^{2+} [29]. Similar enhanced and tunable luminescence performances in $Lu_{3-x}Y_xAl_5O_{12}:Mn^{4+}$ red phosphor can also be explained in details due to the substitution of Y^{3+} for Lu^{3+} [30]. Taking the inspiration from these literatures, we assumed Ga^{3+} to replace Al^{3+} in $SrMgAl_{10}O_{17}:Mn^{4+}$ phosphor to improve luminescence properties. As far as we know, no similar literature has been reported.

In this study, a novel Mn^{4+} -doped $SrMgAl_{10-y}Ga_yO_{17}$ red phosphor with tunable and improved luminescence properties is discovered. The samples were prepared through high-temperature solid-state method in atmospheric environment. The phase

structure, photoluminescence (PL) and photoluminescence excitation (PLE) spectra, absorption spectra, quantum efficiency, lifetime decay curves and electroluminescence properties are investigated in detail. Finally, series of LED devices combined with blue chips and the as-obtained phosphors show bright blue and red emissions which match the plant absorption spectrum well, indicates this phosphor can be a candidate for indoor plant growth light.

2. Experimental section

The raw materials were SrCO₃ (99.99%), MgO (99.99%), Al₂O₃ (99.99%), Ga₂O₃ (99.99%) and MnCO₃ (99.99%), which were bought from Aladdin without further purification. Different contents of H₃BO₃ (AR) (1, 2, 3, 4, 5% wt) acted as flux in the reaction process. All reagents were weighed following the stoichiometric ratio and put into an agate mortar, dropped with a certain amount of absolute ethyl alcohol. These powders were ground for 30 minutes to mix up uniformly then transferred into corundum crucibles. They were put into a tube furnace preheated at 800 °C for 2 hours and sintered at 1500 °C in ambient atmosphere for 5 hours with a heating rate of 5 °C/min in the whole process. All samples were cooled down to room temperature naturally and then reground to fine powders. Finally, the SrMgAl₁₀O₁₇:xMn⁴⁺ ($x = 0.1\%, 0.3\%, 0.5\%, 1.0\%, 1.5\%, 2.0\%$) and SrMgAl_{10-y}Ga_yO₁₇:1.0%Mn⁴⁺ ($y = 1, 2, 3, 4, 5, 6$) samples were obtained.

The X-ray powder diffraction (XRD) patterns were measured ranging from 10° to 80° by a diffractometer (D/SHIMADZU-6000, Japan) which equipped with Cu-Kα radiation ($\lambda=1.5406 \text{ \AA}$). The scanning rate was 6°/min and operating voltage and

current were 40 kV and 40 mA. F-4700 fluorescence spectrophotometer (Hitachi, Japan) was used to obtain the photoluminescence excitation (PLE) and photoluminescence (PL) spectra with a 150W Xe lamp as the excitation lamp. The UV-vis absorption spectra were tested on U-3310 spectrophotometer (Hitachi, Japan). Temperature-dependent PL spectra were measured using F-7000 Spectro-photometer (Hitachi, Japan) by changing the testing temperature from 298 K to 473 K. FLS 1000 fluorescence spectrometer (Edinburgh, UK) was used to get the lifetime curves and quantum efficiency. The photographs of sample were taken by Sony A6400 camera.

The LED devices were fabricated using two-component epoxy resin (Kraft K-9761) to encapsulate the as-obtained phosphors on the blue chip. The transparent resin A and B was mixed with the weight rate of 2:1, then the well-mixed resin and phosphor further mixed with the weight ration changed from 1:1 to 1:3. An ATA-500 measurement system (Everfine, China) was used to carry out the luminescence spectra of these LED devices.

3. Results and discussion

3.1. Structure and phase characterization

First of all, H_3BO_3 was chosen as flux in the synthesis process, as shown in [Fig.S1 \(a\)](#). The XRD pattern of samples exhibit impurity phase of SrAl_2O_4 before the content of H_3BO_3 is below 2 wt %, and after that the XRD represent pure phase of $\text{SrMgAl}_{10}\text{O}_{17}$ with the standard card of PDF#26-0879. Considering the photoluminescence excitation (PLE) and emission (PL) spectra of samples with different content of flux, the emission intensity come to the maximum when 2 wt %

H₃BO₃ participate in the synthesis reaction, and thus this condition is selected in the following section.

The SrMgAl₁₀O₁₇ crystal belongs to *P63/mmc* space group with hexagonal structure as shown in Fig.1 (a). It is worth noting that the asymmetric part of the unit cell contains four independent sites of Al³⁺ which can be written as two [AlO₆] and two [AlO₄]. One type of Al³⁺ in tetrahedron [AlO₄] shares the position with Mg²⁺ randomly. In general, activator ion Mn⁴⁺ prefers to replace Al³⁺ in octahedron [AlO₆] position due to the similar ionic radius between Mn⁴⁺ ($r = 0.530 \text{ \AA}$, CN = 6) and Al³⁺ ($r = 0.535 \text{ \AA}$, CN = 6), and Mn⁴⁺ is more stable in octahedral environment with stronger ligand-field stabilization energy [31]. When a small quantity of Mn⁴⁺ replace Al³⁺ in the SrMgAl₁₀O₁₇ matrix, the XRD patterns keep the original position without any shift, the experimental results are shown in Fig. S2. Rietveld refinement analysis is performed in SrMgAl₁₀O₁₇:1.0%Mn⁴⁺ sample, and there are still two impurities phase SrAl₂O₄ and MgAl₂O₄, the contents of them are determined to be 6.38% and 7.50%. The cell parameters of this sample are listed as $a = b = 5.62 \text{ \AA}$, $c = 22.41 \text{ \AA}$ and $V = 614.25 \text{ \AA}^3$, while these parameters in pure SrMgAl₁₀O₁₇ are $a = b = 5.63 \text{ \AA}$, $c = 22.47 \text{ \AA}$ and $V = 616.80 \text{ \AA}^3$, indicates that Mn⁴⁺ is doped into the unit cell successfully. With more Ga³⁺ doped into the phosphor, the XRD curves exhibit complex phase gradually, as seen in Fig.1 (b). The sample $y = 0$ match well with the standard card of SrMgAl₁₀O₁₇ at first, then MgAl₂O₄ appear and its content gradually increase. XRD pattern of the sample $y = 3$ shows two phases: SrMgAl₁₀O₁₇ and MgAl₂O₄, which are noted as ● and ◆ respectively. According to literature research,

there is no report about Mn^{4+} luminescence property in single MgAl_2O_4 matrix. Wu et al [32] found that $\text{MgAl}_2\text{O}_4:\text{Mn}^{4+}$ sample show no photoluminescence emission but MgAl_2O_4 phase have advantages on the luminescence performance of the $\text{MgAl}_2\text{O}_4/\text{CaAl}_2\text{O}_9:\text{Mn}^{4+}$ composite sample. Accordingly, combined with the experiment results, it can expect that a small amount of MgAl_2O_4 phase can enhance the luminescence properties of sample. With further increasing of Ga^{3+} dopant, the phase of samples are mainly composed of $\text{SrMgAl}_{10}\text{O}_{17}$ host, XAl_2O_4 ($\text{X}=\text{Mg}^{2+}$, Mn^{2+}) and Ga_2O_3 , and it should be noticed that the latter three compounds have similar XRD pattern with almost the same peak position. Considering there is no octahedral coordination in these phases as suitable luminescence centers for Mn^{4+} , thus they have little effect on the luminescence properties of sample theoretically.

3.2. Photoluminescence properties of $\text{SrMgAl}_{10-y}\text{Ga}_y\text{O}_4:x\text{Mn}^{4+}$ phosphors

Fig. 2(a) and (c) exhibit the PLE ($\lambda_{\text{em}} = 662 \text{ nm}$) and PL ($\lambda_{\text{ex}} = 320 \text{ nm}$ and $\lambda_{\text{ex}} = 467 \text{ nm}$) spectra of $\text{SrMgAl}_{10}\text{O}_{17}:x\text{Mn}^{4+}$ phosphors. It is obvious that the excitation and emission intensities are increasing with a rising concentration of Mn^{4+} at first, both of them reach maximum when $x = 1.0\%$ and then decrease with more Mn^{4+} doping. For the purpose of researching the concentration quenching behavior of $\text{SrMgAl}_{10}\text{O}_{17}:x\text{Mn}^{4+}$ phosphor, the critical distance R_c is taken into account, whose value can be determined by the equation below [33, 34]:

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

where V stands for the cell volume, x_c is the critical concentration of Mn^{4+} and N refers to the number of available sites occupied by Mn^{4+} , respectively. For

SrMgAl₁₀O₁₇:1.0%Mn⁴⁺ phosphor, $V = 614.25 \text{ \AA}^3$, $x_c = 0.010$ and $N = 2$, and the R_c is calculated to be 38.86 \AA . This value is much larger than 5 \AA which corresponding to the interaction between activator ions, therefore, the concentration quenching of SrMgAl₁₀O₁₇: x Mn⁴⁺ belongs to the multipolar-multipolar interaction which can be divided to be dipole–dipole (d–d), dipole–quadrupole (d–q), and quadrupole–quadrupole (q–q) interactions, respectively. They refer to different values of θ in the following formula, which is often used to reflect the interactions based on the Dexter's theory [35, 36]:

$$\frac{I}{x} = k[1 + \beta(x)^{\theta/3}]^{-1} \quad (2)$$

where I and x mean the emission intensity and concentration of Mn⁴⁺ dopant, k and β refer to a certain excitation condition and specific matrix crystal, and $\theta = 6, 8, 10$ refer to d–d, d–q, and q–q interactions, respectively.

The specific value of θ is usually obtained by the slope of the dependence of $\log(I/x)$ versus $\log(x)$. From the inset in Fig. 2(a), there are two fitting lines corresponding to 320 nm and 467 nm for the excitation wavelength, two slopes are -1.029 and -1.046, respectively. The θ_1 and θ_2 are calculated to be 3.087 and 3.138, respectively, both of them are close to 6 which indicate the concentration quenching mechanism is dipole–dipole (d–d) interaction in SrMgAl₁₀O₁₇: x Mn⁴⁺ phosphor.

According to Fig.2 (b), the PLE spectrum of SrMgAl₁₀O₁₇:1.0%Mn⁴⁺ is fitted into three Gaussian peaks located at 32468, 27472 and 22026 cm⁻¹, they are related to $^4A_2 \rightarrow ^4T_1$, $^4A_2 \rightarrow ^2T_2$ and $^4A_2 \rightarrow ^4T_2$ transitions of Mn⁴⁺, respectively. The PLE spectra of SrMgAl₁₀O₁₇: x Mn⁴⁺ phosphors have a large range from 220 to 580 nm,

indicates that these phosphors can be easily excited by commercial blue and near-UV chips. Fig. 3(a) and (b) exhibit the photoluminescence excitation (PLE) and photoluminescence (PL) spectra of SrMgAl_{10-y}Ga_yO₁₇:1.0%Mn⁴⁺ samples. The intensity of PLE and PL spectra both increase with an incremental content of Ga³⁺ doping, and come to the maximum when $y = 3$. It is worth noting that the shape of PL spectrum changed significantly upon Ga³⁺ doping which related to a shift of the luminescence center. In the sample of SrMgAl₁₀O₁₇:1.0%Mn⁴⁺ ($y = 0$), the emission come from the Al center absolutely. When Ga³⁺ replace Al³⁺ in samples from $y = 1$ to $y = 6$, new luminescence center of Ga site is generated, thus resulted in the change of spectra. Another interesting phenomenon is that the normalized PLE spectra exhibit obvious red shift as shown in Fig.3 (c), that the ⁴A₂→²T₂ transitions show a great enhancement and red shift with the increase of Ga³⁺ doping. Meanwhile, PL spectra excited at different wavelength all show slight red shift in Fig. 3 (d) and Fig. S4.

Based on this situation, crystal field theory and nephelauxetic effect are employed to explain these phenomena. There are three parameters to assess the effect of crystal field on the luminescent properties of Mn⁴⁺ that are described as the crystal-field parameter (D_q), Racah parameters (B) and (C), respectively. They are directly related to the PLE and PL spectra, and the values of D_q , B and C can be calculated from following equations [30, 37]:

$$D_q = \frac{E(^4A_2 - ^4T_2)}{10} \quad (3)$$

$$\frac{D_q}{B} = \frac{15(x-8)}{(x^2-10x)} \quad (4)$$

$$x = \frac{E(^4A_2 - ^4T_1) - E(^4A_2 - ^4T_2)}{D_q} \quad (5)$$

$$\frac{E(^2E_g \rightarrow ^4A_2)}{B} = \frac{3.05C}{B} + 7.9 - \frac{1.8B}{D_q} \quad (6)$$

According to Fig. 2, the energy levels of 4T_1 , 4T_2 and 2E_g in $\text{SrMgAl}_{10}\text{O}_{17}:1.0\%\text{Mn}^{4+}$ are 32468, 22026 and 15110 cm^{-1} , and its D_q , B and C are calculated to be 2203, 1123 and 2423 cm^{-1} , respectively. The value of D_q/B is evaluated to be 2.1, means that Mn^{4+} is exposed in a weak crystal field in $\text{SrMgAl}_{10}\text{O}_{17}$ matrix. Meanwhile, with the concentration of Ga^{3+} dopant increasing from 10% to 60%, the values of D_q/B are determined to be 2.22, 2.23, 2.24, 2.35, 2.43 and 2.49 which reveal a strong crystal field [38]. However, the change of crystal field is not the main reason in the red shift of Mn^{4+} emission due to its characteristic parity-forbidden d-d transition, and the nephelauxetic effect takes the predominant role in the luminescence of Mn^{4+} in most oxide phosphors[39, 40]. The nephelauxetic effect has an influence on the energy of parity-forbidden $^2E_g \rightarrow ^4A_2$ transition of Mn^{4+} , which can be described by the following formula [41, 42]:

$$\beta_1 = \sqrt{\left(\frac{B}{B_0}\right)^2 + \left(\frac{C}{C_0}\right)^2} \quad (7)$$

where B_0 and C_0 are the Racah parameters for free ions, and the values of them are $B_0 = 1160 \text{ cm}^{-1}$ and $C_0 = 4303 \text{ cm}^{-1}$ for Mn^{4+} , respectively. The details of these parameters about crystal field and nephelauxetic effect of $\text{SrMgAl}_{10-y}\text{Ga}_y\text{O}_{17}:1.0\%\text{Mn}^{4+}$ phosphors are listed in Table 1.

According to the Tanabe-Sugano energy-level diagram of Mn^{4+} as shown in Fig. 4, the energy of $^2E_g \rightarrow ^4A_2$ transition of Mn^{4+} is mainly determined by the Racah parameter B which relates to chemical bonding between the Mn^{4+} and the ligand. In general, a weaken nephelauxetic effect in bonding has smaller Racah parameter B ,

and thus shift down energy scale of 2E level [43, 44], means it has lower energy. Meanwhile, wavelength is inversely proportional to energy that can ascribe to $\lambda=1240/E$, thus lower energy E results in longer wavelength, corresponding to red-shift phenomenon of the emission position. To sum up, a smaller value of the Racah parameter B gives the lower energy of ${}^2E_g \rightarrow {}^4A_2$ transition, corresponding to the red-shift phenomenon in the luminescence emission, and experimental results are in agreement with the theory.

Fig. 5(a) shows the diffuse reflection spectra of these powder samples, it is obvious that the reflection rate are decreasing with more content of Ga^{3+} , which has the same trend as a change of the PLE spectra. All diffuse reflection curves can be divided into four parts corresponding to the absorption of matrix and ${}^4A_2 \rightarrow {}^4T_1$, ${}^4A_2 \rightarrow {}^2T_2$, ${}^4A_2 \rightarrow {}^4T_2$ transitions of Mn^{4+} , respectively, which is consistent with the Gaussian fitting of PLE spectrum mentioned above. The diffuse reflection spectra can be translated into UV-vis absorption spectra, and get the band gap energy further through the following equation [45, 46]:

$$(\alpha hv)^{1/2} = A(hv - E_g) \quad (8)$$

where α and A stand for the absorption parameter and proportional constant, hv and E_g refer to the photon energy and the band gap, respectively. The band gap energy of this series samples increases from 5.25 eV to 5.44 eV and then drops down to 5.21 eV, as shown in Fig.5 (b).

For the purpose of investigating the luminescence kinetics in $SrMgAl_{10-y}Ga_yO_{17}:Mn^{4+}$ phosphors, the lifetime of these samples are measured and

displayed in Fig.6 (a). All of these curves fit well with the double-exponential decay model, see the following formula [47]:

$$I(t) = I_0 + C_1 \exp(-t/\tau_1) + C_2 \exp(-t/\tau_2) \quad (9)$$

And the average lifetime $\tau_{\text{ave.}}$ can be further calculated as [48]:

$$\tau_{\text{ave.}} = \frac{C_1\tau_1^2 + C_2\tau_2^2}{C_1\tau_1 + C_2\tau_2} \quad (10)$$

where I and I_0 stand for the photoluminescence intensity at time t and 0, C_1 and C_2 refer to pre-exponential factors, τ_1 and τ_2 are the lifetime components, respectively. The average lifetimes of $\text{SrMgAl}_{10-y}\text{Ga}_y\text{O}_{17}:\text{Mn}^{4+}$ are calculated to be 0.446, 0.476, 0.383, 0.320, 0.266, 0.207 and 0.126 ms when the Ga^{3+} dopant increased from 0 to 60%. Details about fitting information of the decay times are shown in the Table S1. The double-exponential decay model indicates that Mn^{4+} should have two luminescence centers in $\text{SrMgAl}_{10-y}\text{Ga}_y\text{O}_{17}$ matrix, which is consistent with the emission spectra.

Quantum efficiency (QE) is an important parameter to evaluate luminescence property of phosphors, the internal quantum efficiency (IQE) of $\text{SrMgAl}_7\text{Ga}_3\text{O}_{17}:\text{Mn}^{4+}$ and $\text{SrMgAl}_{10}\text{O}_{17}:\text{Mn}^{4+}$ samples are shown in Fig.6 (b) and Fig.S5, which can be calculated via the following formula [49, 50]:

$$\eta = \frac{\int L_S}{\int E_R - \int E_S} \quad (11)$$

herein, η is internal quantum efficiency, L_S stands for the luminescence spectra of the sample, E_R and E_S refer to the excitation line of BaSO_4 reference and sample, respectively. The IQE of $\text{SrMgAl}_7\text{Ga}_3\text{O}_{17}:\text{Mn}^{4+}$ and $\text{SrMgAl}_{10}\text{O}_{17}:\text{Mn}^{4+}$ are determined to be 29.67% and 51.40%. These values of the samples are at a moderate

level compared to other red or deep red phosphors that have been reported such as SrLaScO₄:Mn⁴⁺ (IQE: 12.2%) [51], NaGdCa₄W₂O₁₂:Mn⁴⁺ (IQE: 21.0%) [52], SrAl₃BO₇:Mn⁴⁺ (IQE: 26.0%) [53], Ba₂YNbO₆:Mn⁴⁺ (IQE: 29.2%) [54], Ca₂LaSbO₆:Mn⁴⁺ (IQE: 52.2%) [55]. Nevertheless, the photoluminescence properties of SrMgAl_{10-y}Ga_yO₁₇:Mn⁴⁺ phosphors are expected to improve via further modification process likes charge compensation or energy transfer with co-doping sensitizer ion.

Fig.7 (a) and (b) are the temperature-dependence luminescence spectra to evaluate the thermal stability of phosphor. The emission intensity decreases rapidly as the temperature increasing in both of the two samples, and the decline trend is more serious in SrMgAl₇Ga₃O₁₇:Mn⁴⁺ phosphor. It because that Ga³⁺ doping destroys the stability of matrix structure to some extent and leads to lower activation energy, promotes the loss of energy with non-radiative processes and results in worse thermal stability. Therefore, thermal stability of this phosphor needs to be further improved through modifications such as coating and synthesis optimization to fit further applications.

Fig. 8 shows the mechanism diagram in SrMgAl₉GaO₁₇:Mn⁴⁺ and SrMgAl₄Ga₆O₁₇:Mn⁴⁺ for clearly describing the energy transition process. There is only Mn⁴⁺ ion transition emission in these phosphors, electron is excited from ⁴A₂ energy level to ⁴T₁, ⁴T₂ and even conduction band, and then relaxes to ²E_g level through non-radiative transition process, finally transfers to ⁴A₂ level with bright red emission. According to the results calculated above, the sample with more Ga³⁺

dopant has lower energy in the 4T_1 , 4T_2 and 2E_g states, and the energy of ${}^2E_g \rightarrow {}^4A_2$ transition directly decide the position of red emission. Therefore, there is an obvious red shift on the PLE and PL spectra with more content of Ga^{3+} in $SrMgAl_{10-y}Ga_yO_{17}:Mn^{4+}$ phosphors.

3.3. The emission property of as-obtained LED device

In order to search for the potential application of $SrMgAl_{10-y}Ga_yO_{17}:Mn^{4+}$ phosphor, LED devices are fabricated with 470 nm blue chips and different content of the as-obtained $SrMgAl_7Ga_3O_{17}:Mn^{4+}$ sample, the emission spectra are shown in [Fig.9 \(a\) and \(b\)](#). The red emission continues rising with an increasing content of phosphor, indicates that the ratio of blue and red light can be easily adjusted to meet the needs of different plants growth. The CIE chromaticity coordinates of these device change from (0.1456, 0.0365) to (0.5776, 0.2291), that from blue light to red light. Other four chromaticity coordinates with different content of the phosphor are (0.1852, 0.0603), (0.2149, 0.0754), (0.3391, 0.1256), and (0.4578, 0.1782), respectively. The emission spectrum of the fabricated LED device with appropriate phosphor is mainly composed of the blue band (420-500 nm) and red band (650-750 nm) with the peak at 470 nm and 663 nm, which can fit the absorption spectra of plant pigment including chlorophyll *A*, *B* and phytochrome P_R , P_{FR} , indicates that the $SrMgAl_{10-y}Ga_yO_{17}:Mn^{4+}$ phosphors can be a candidate for plant growth LED light.

4. Conclusions

In this work, a series of novel $SrMgAl_{10-y}Ga_yO_{17}:xMn^{4+}$ red phosphors with enhanced and tunable luminescence properties were synthesized through high

temperature approach. An appropriate amount of Ga^{3+} dopant improves the luminescent intensity to 163% in $\text{SrMgAl}_7\text{Ga}_3\text{O}_{17}:\text{Mn}^{4+}$ sample compared to the original $\text{SrMgAl}_{10}\text{O}_{17}:\text{Mn}^{4+}$ phosphor and causes the red shift in photoluminescence excitation (PLE) and photoluminescence (PL) spectra. The red shift observed in the spectra is explained by crystal field theory and nephelauxetic effect. All samples have a broad band in the PLE spectra ranging from 220 to 580 nm indicates they can be excited by both near ultraviolet and blue chips. The Ga^{3+} dopant increases the band gap of $\text{SrMgAl}_7\text{Ga}_3\text{O}_{17}:\text{Mn}^{4+}$ sample to reduce non-radiative transitions and improve the emission intensity but leads to worse thermal stability for its low activation energy. Devices assembled with blue chip and the as-obtained $\text{SrMgAl}_7\text{Ga}_3\text{O}_{17}:\text{Mn}^{4+}$ phosphor emits bright blue and red light which match the plant absorption spectra well, thus the $\text{SrMgAl}_{10-y}\text{Ga}_y\text{O}_{17}:\text{Mn}^{4+}$ phosphor have potential application on plant growth LED light.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://www.sciencedirect.com/>

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Table 1 Energy States, crystal field parameters and calculated values of β_1 in $\text{SrMgAl}_{10-y}\text{Ga}_y\text{O}_{17}:1.0\%\text{Mn}^{4+}$ ($y=0, 1, 2, 3, 4, 5, 6$) phosphors.

Sample	${}^4\text{A}_2 \rightarrow {}^4\text{T}_1$ (cm^{-1})	${}^4\text{A}_2 \rightarrow {}^4\text{T}_2$ (cm^{-1})	${}^2\text{E}_g \rightarrow {}^4\text{A}_2$ (cm^{-1})	D_q	B	C	D_q/B	β_1
y=0	32468	22026	15110	2203	1123	2423	1.96	1.120
y=1	30303	21097	15133	2110	951	2797	2.22	1.046
y=2	29940	20877	15129	2088	935	2833	2.23	1.040
y=3	29674	20704	15115	2070	924	2852	2.24	1.036
y=4	29240	20619	15106	2062	876	2952	2.35	1.020
y=5	28986	20576	15088	2058	847	3008	2.43	1.011
y=6	28653	20450	15047	2045	821	3051	2.49	1.002

Figures Captions

Fig.1. (a) The crystal structure of SrMgAl₁₀O₁₇ host, the coordination environment of the octahedron [AlO₆] and tetrahedron [AlO₄]; (b) The XRD patterns of SrMgAl_{10-y}Ga_yO₁₇:1.0%Mn⁴⁺ ($y = 0, 1, 2, 3, 4, 5, 6$) phosphors.

Fig.2. (a) The photoluminescence excitation (PLE) spectra of SrMgAl₁₀O₁₇:xMn⁴⁺ ($x=0.1\%, 0.3\%, 0.5\%, 1.0\%, 1.5\%, 2.0\%$), and the inset is the dependence of $\log(I/x)$ versus $\log(x)$; (b) Gaussian fitting of SrMgAl₁₀O₁₇:1.0%Mn⁴⁺ sample; (c) The photoluminescence (PL) spectra of samples under the excitation of 320 nm and 467 nm respectively, and the inset is the tendency of emission intensity varies with Mn⁴⁺ doping concentration.

Fig.3. (a) Photoluminescence excitation and (b) photoluminescence spectra of SrMgAl_{10-y}Ga_yO₁₇:1.0%Mn⁴⁺ ($y=0, 1, 2, 3, 4, 5, 6$) phosphors; the normalized PLE spectra (c) monitored at 663 nm; the normalized PL spectra under (d) 340 nm excitation, and the insets are enlarged part of the peaks in the wavelength ranging from 658 to 668 nm.

Fig.4. Tanabe-Sugano energy-level diagram of Mn⁴⁺ in SrMgAl_{10-y}Ga_yO₁₇:1.0%Mn⁴⁺ phosphors.

Fig.5. (a) The diffuse reflection spectra of SrMgAl_{10-y}Ga_yO₁₇:1.0%Mn⁴⁺ phosphors and one PLE spectrum of SrMgAl₇Ga₃O₁₇:1.0%Mn⁴⁺ sample; (b) the corresponding band gap energy fitting.

Fig.6. (a) The lifetime decay curves of SrMgAl_{10-y}Ga_yO₁₇:1.0%Mn⁴⁺ phosphors monitored at 661 nm; the measurement of quantum efficiency of (b) SrMgAl₇Ga₃O₁₇:1.0%Mn⁴⁺, and the insets are the enlarged pattern ranging from 600 nm to 780 nm.

Fig.7. The temperature-dependence luminescence spectrum of (a) SrMgAl₁₀O₁₇:1.0%Mn⁴⁺ and (b) SrMgAl₇Ga₃O₁₇:1.0%Mn⁴⁺; (c) configurational coordinate diagram for Mn⁴⁺ ions in SrMgAl_{10-y}Ga_yO₁₇ phosphor.

Fig.8. Mechanism diagram of luminescence in SrMgAl₆GaO₁₇:1.0%Mn⁴⁺ and SrMgAl₄Ga₆O₁₇:1.0%Mn⁴⁺ phosphors.

Fig.9. (a) (b) Emission spectra of the fabricated LEDs combined with 470 nm blue chip and different contents of SrMgAl₇Ga₃O₁₇:1.0%Mn⁴⁺ sample, and the insets are the photos of the device; (c) CIE chromaticity diagram of LED devices; (d) the comprehensive comparison between emission spectra of device and the absorption curves of the plant pigments of Chlorophyll *A*, *B* and phytochrome *P_R*, *P_{FR}*.

Fig.1

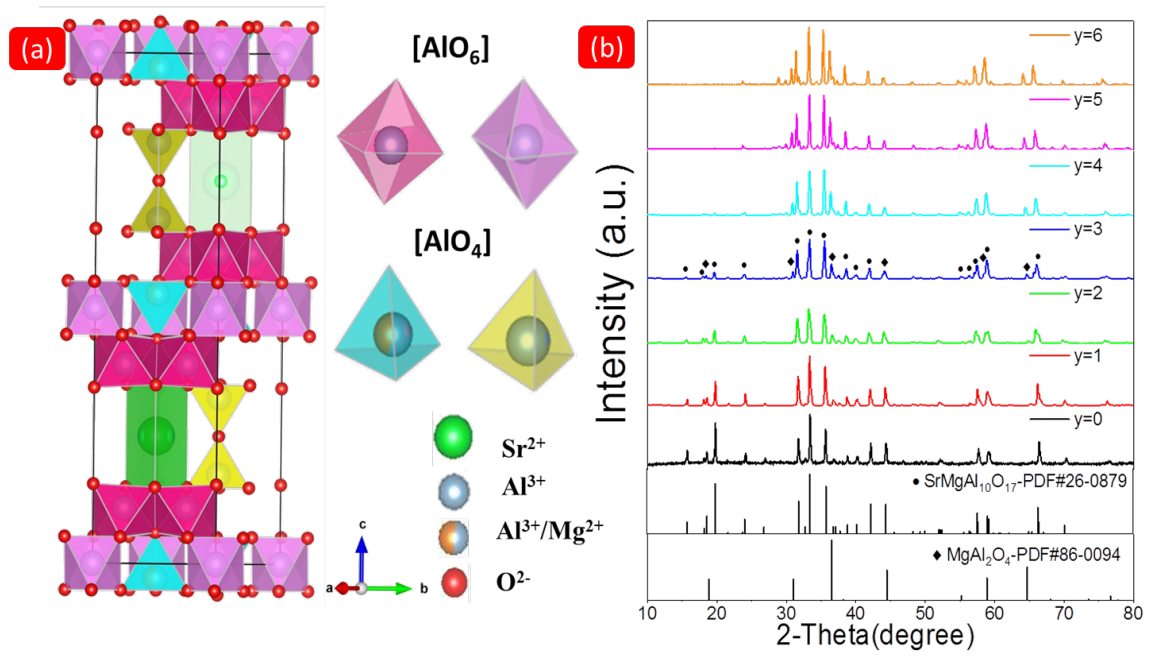


Fig.2

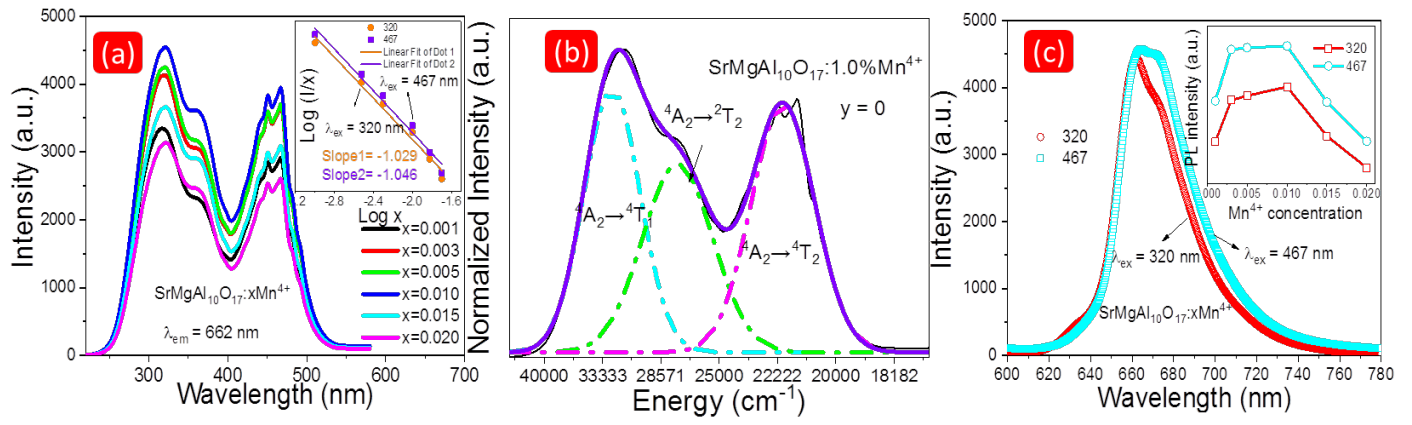


Fig.3

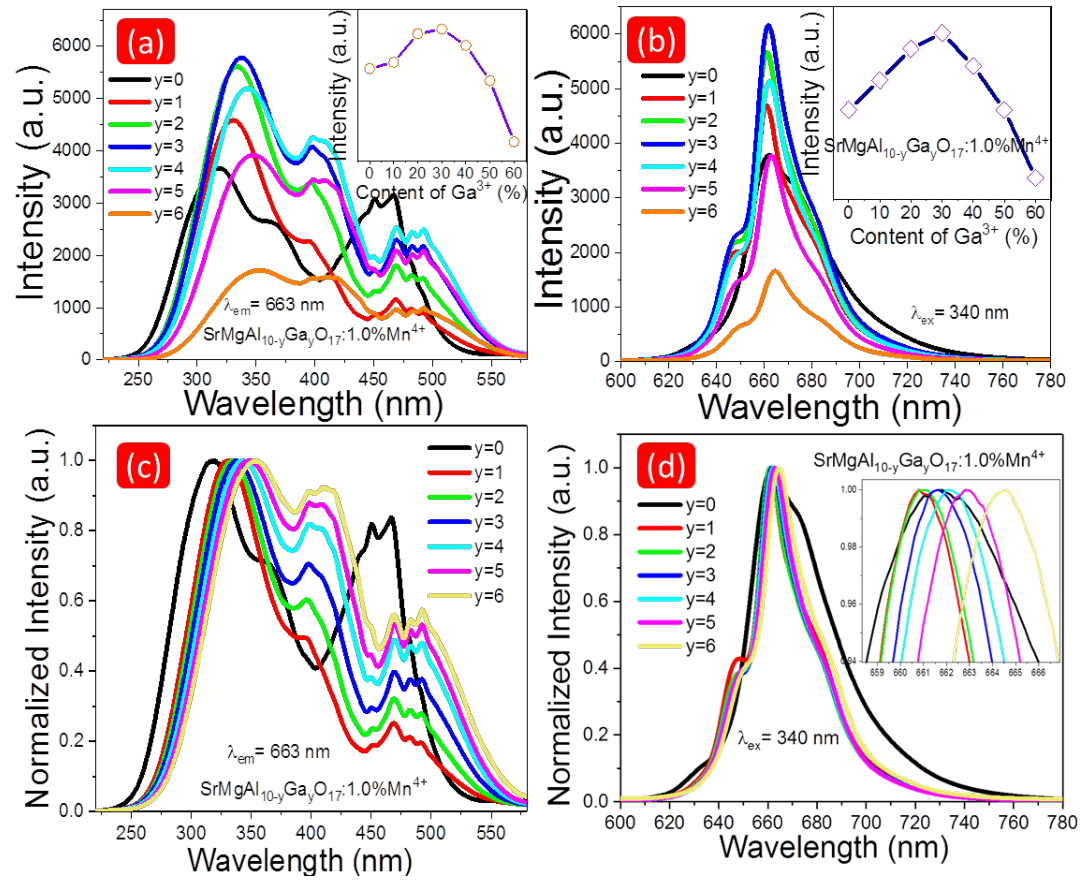


Fig.4

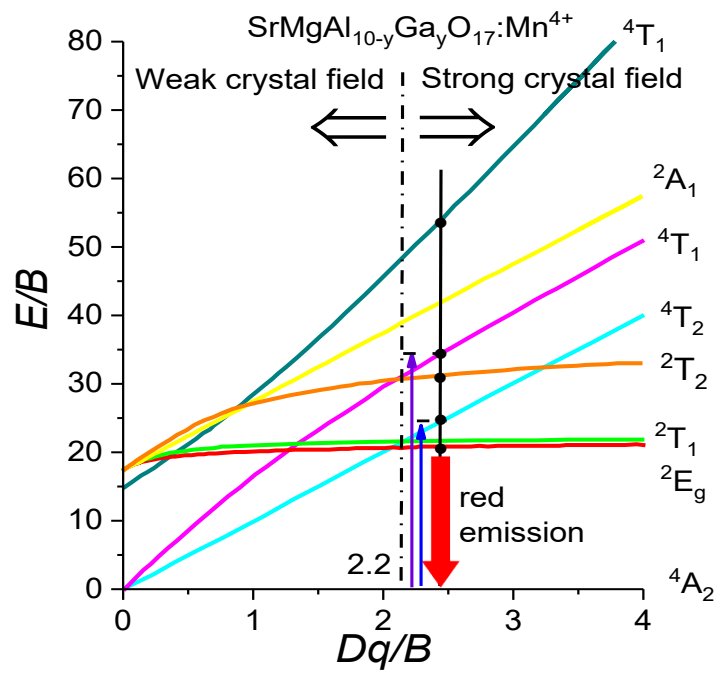


Fig.5

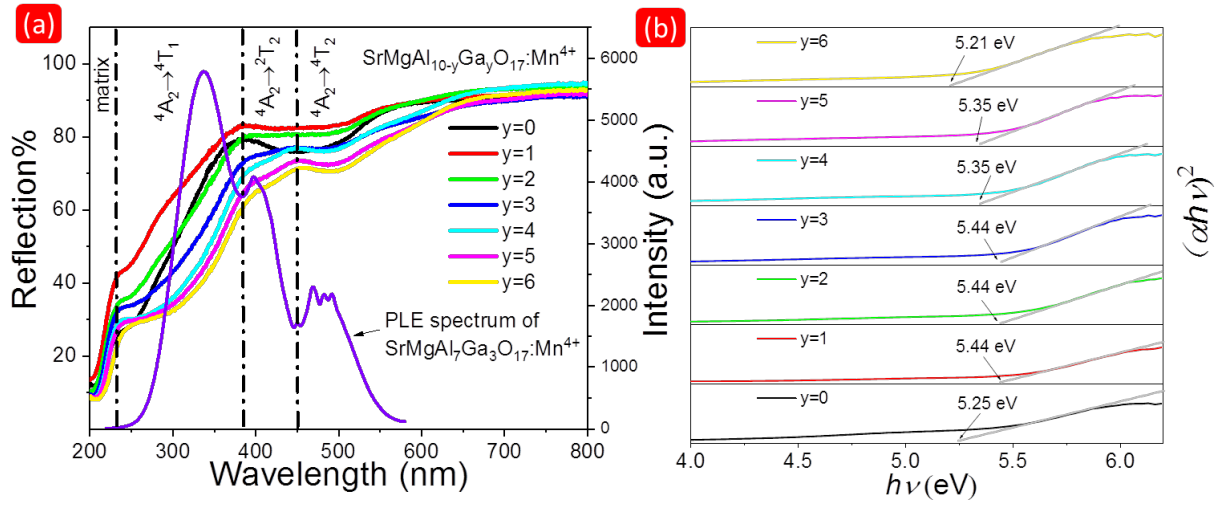


Fig.6

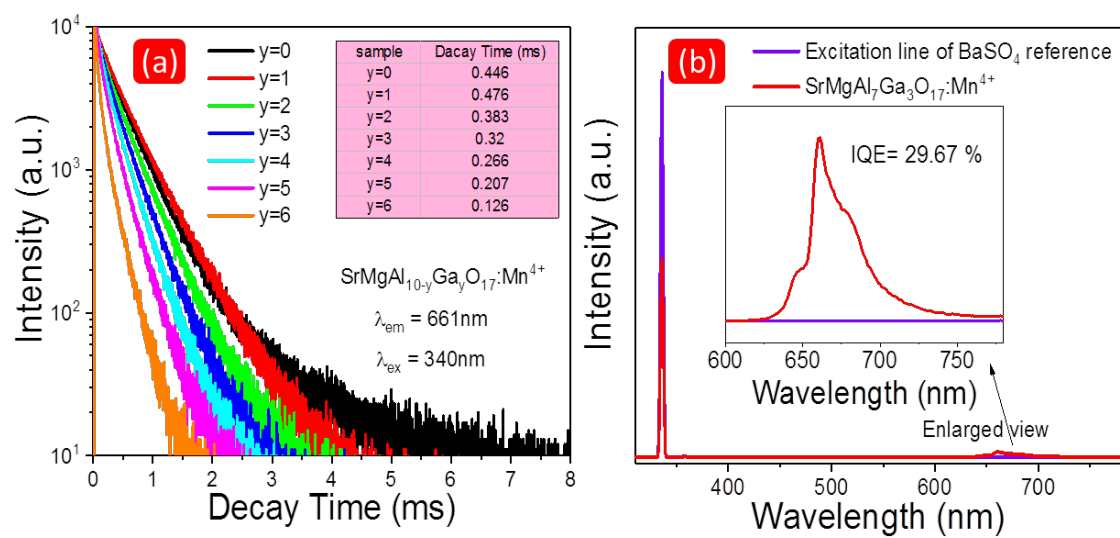


Fig.7

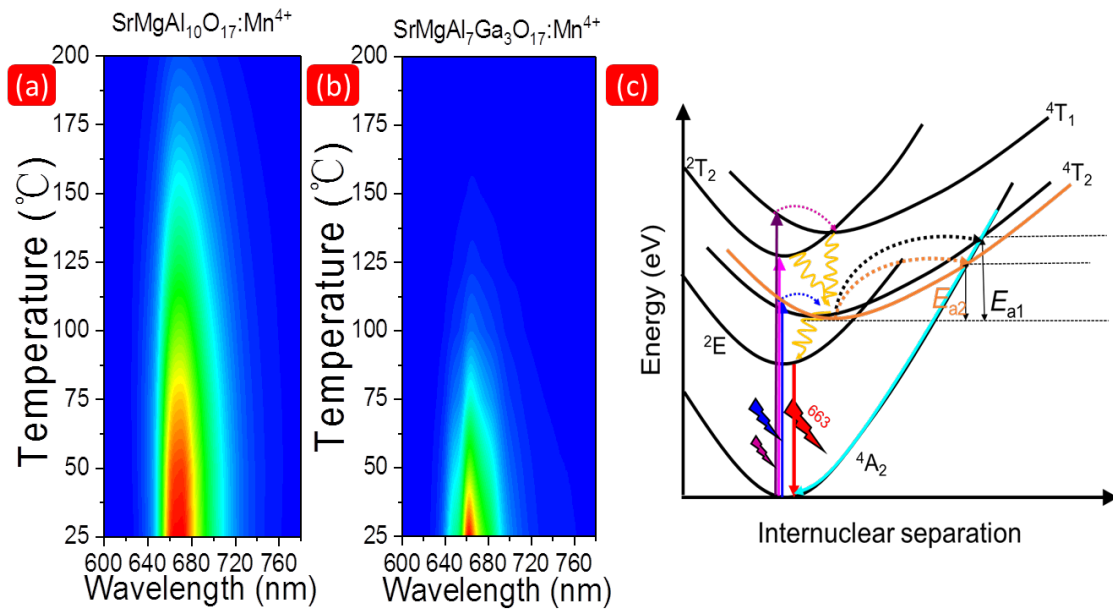


Fig.8

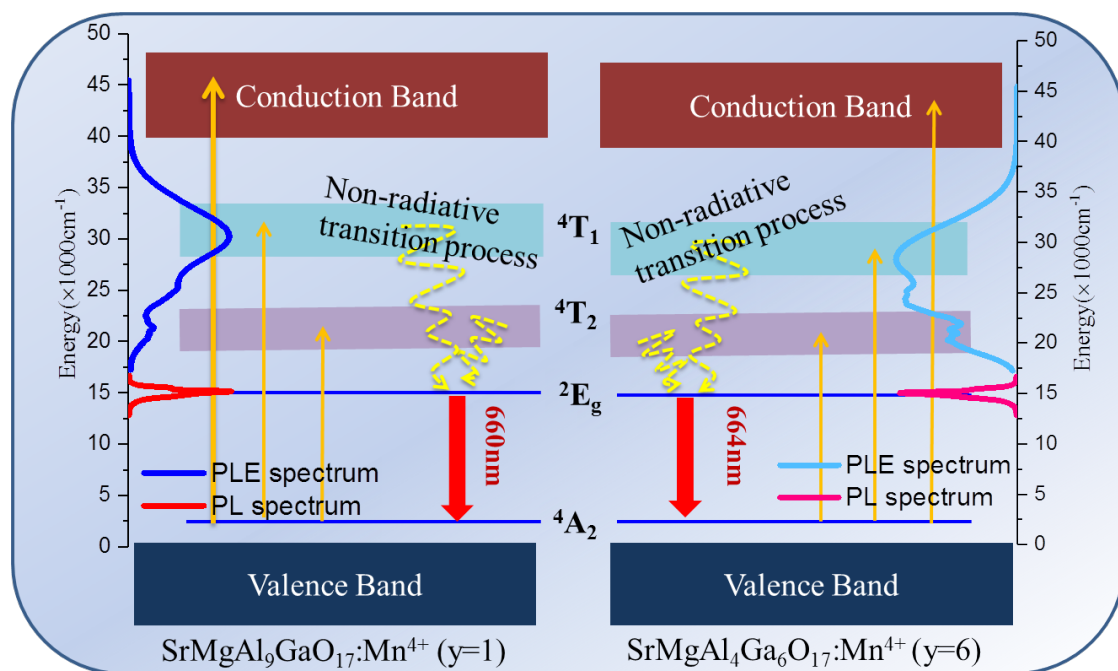


Fig.9

