Facile synthesis of the desired red phosphor $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6\text{:Eu}^{2+}$ for high CRI white LEDs and plant growth LED device

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Abstract

The red emission with suitable peak wavelength and narrow band is acutely required for

high color rendering index (CRI) white LEDs without at the cost of the luminous efficacy.

Herein, the Li₂Ca₂Mg₂Si₂N₆:Eu²⁺ red phosphor was prepared with facile solid-state method using

Ca₃N₂, Mg₃N₂, Si₃N₄, Li₃N, and Eu₂O₃ as the safety raw materials under atmospheric pressure

for the first time, which shows red emission peaking at 638 nm with full width at half maximum

(FWHM) of 62 nm under blue light irradiation and becomes the desired red phosphor to realize

the balance between luminous efficacy and high CRI in white LEDs. The morphology, structure,

luminescence properties, thermal quenching behavior, and chromaticity stability of the

Li₂Ca₂Mg₂Si₂N₆:Eu²⁺ phosphor are investigated in detail. Concentration quenching occurs when

the Eu²⁺ content exceeds 1.0 mol%, whereas high temperature photoluminescent measurements

show a 32% drop from the room temperature efficiency at 423 K. In view of the excellent

luminescence performances of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$: Eu^{2+} phosphor, a white LEDs with CRI of 91 as

a proof-of-concept was fabricated by coating the title phosphor with Y₃Al₅O₁₂:Ce³⁺ on blue LED

chip. In addition, the potential application of the title phosphor in plant growth LED device was

also demonstrated. All the results indicated that the Li₂Ca₂Mg₂Si₂N₆:Eu²⁺ is a promising

red-emitting phosphor for blue-LED-based high CRI white LEDs and plant growth lighting

sources.

Keywords: Li₂Ca₂Mg₂Si₂N₆:Eu²⁺; phosphor; high CRI white LEDs; plant growth LED device

Introduction

To pursue high luminous efficacy and reduce energy consumption caused by low eye sensitivity in the infrared region, red phosphors with suitable peak wavelength and narrow band are urgently needed for high color rendering index (CRI) white LEDs. ^{1,2} There is a tradeoff between luminous efficacy and high CRI in all types of white LEDs, which means that the improvement in one usually brings about the reduction of the other. The emission peak should be at ~630 nm with relatively narrow band for the optimum balance. ^{3,4}

Over the past, nitridosilicate phosphors with excellent luminescence properties and stability as red components were widely utilized in high CRI white LEDs, such as Sr₂Si₅N₈:Eu²⁺, (Ca.Sr)AlSiN₃:Eu^{2+,5,6} However, their broad emissions produce infrared emissions falling out the sensitivity range of the human eye, which leads to the wasted power input and thereby limits the $maximum \ achievable \ luminous \ efficacy. \ The \ nitridolithoaluminate \ SrLiAl_3N_4: Eu^{2+} \ phosphor$ debuted with peak at ~650 nm and narrow band emission (FWHM = 50 nm) as well as excellent thermal stability, which shows luminous efficacy improvement by 14% compared to commercial phosphor-converted white LEDs (pc-white LEDs).⁷⁻⁹ Unfortunately, slight red-shift of the emission maximum of SrLiAl₃N₄:Eu²⁺ phosphor means a considerable part in spectral regions with low human eye sensitivity not fulfilling requirements of both high CRI and luminous efficacy. 10,11 Exploration of UCr₄C₄ structure system contributes to the discovery of novel red emitting nitride phosphor. 12-15 CaBaLi₂Al₆N₈:Eu²⁺ derived from SrLiAl₃N₄ structure with heavier homolog Ba substituting for Sr has been demonstrated to the blue-shift of the peak wavelength, it exhibits red emission peaking at 636~639 nm with FWHM of 48~57 nm, as well as another emission peak at 790 nm with FWHM of 89 nm, ¹⁶ while the latter emission hinders the enhancement in luminous efficacy. Nitridomagnesosilicates SrMg₃SiN₄:Eu²⁺ phosphor gives a red emission peaking at 615 nm and shows the narrowest emission band (FWHM of 43 nm) up to date, but it undergoes serious thermal quenching,¹⁷ hampering its possible applications in pc-white LEDs.

The narrow band red-emitting nitridomagnesosilicate $\text{Li}_2(\text{Ca}_{1-x}\text{Sr}_x)_2\text{Mg}_2\text{Si}_2\text{N}_6$: Eu^{2+} ($x=0\sim0.06$) was reported by Wolfgang Schnick group in 2017 based on the work of describing the structure of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$ compound. 18 $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$: Eu^{2+} shows red emission at 638 nm as well as FWHM of 62 nm under blue light irradiation, which is desirable for high CRI white LEDs. 19 However, the preparation of this phosphor needs the rigorous preparation conditions, such as the sealed tantalum ampules into a silica glass tube, long reaction time more than 90 h, dangerous raw materials including $\text{Si}(\text{NH})_2$ and LiN_3 . Also, its luminescent properties, thermal quenching behavior, and potential application in high CRI white LEDs lacks of systematic research.

In this paper, narrow band red emitting nitride $Li_2Ca_2Mg_2Si_2N_6$: Eu^{2+} phosphor was prepared using Ca_3N_2 , Mg_3N_2 , Si_3N_4 , Li_3N , and Eu_2O_3 as the safety raw materials under atmospheric pressure for the first time. The morphology, structure, luminescence properties, optimal doped concentration, thermal quenching behavior, and chromaticity stability of the title phosphor are investigated in detail. Employing the title phosphor, commercial $Y_3Al_5O_{12}$: Ce^{3+} phosphor, and InGaN-based blue LED chip, a proof-of-concept white LEDs with a CRI of 90 and R9 = 90 is achieved.

Compared with the artificial plant growth LEDs through integrating of red and blue LED chips, the phosphor-converted plant growth LED device exhibits several merits including low

cost, facile fabrication, and spectral diversification. However, searching a red phosphor with suitable peak wavelength and narrow band matching well with the absorption spectrum of chlorophyl is acutely required. Thanks to the excellent luminescence performances of Li₂Ca₂Mg₂Si₂N₆:Eu²⁺, in this paper, we also investigated the potential application of the title red phosphor in plant growth LED device.

Experimental

Preparation of Li₂Ca₂Mg₂Si₂N₆:Eu²⁺ phosphor

The phosphor powders of $\text{Li}_2\text{Ca}_{2(1-x)}\text{Mg}_2\text{Si}_2\text{N}_6$: $x\text{Eu}^{2+}$ (x=0.5%-4.0%) were prepared with the conventional high-temperature solid-state reaction. The raw materials were Ca_3N_2 (99%, purchased from Desunmet Ceramic Material Co., Ltd.), Mg_3N_2 (99.9%, purchased from Desunmet Ceramic Material Co., Ltd.), Si_3N_4 (Aladdin, 99.5%), Li_3N (Alfa Assad, 99.5%), and Eu_2O_3 (Aladdin, 99.99%). The elemental precursors were weighted stoichiometrically and grounded thoroughly using agate mortar and pestle in a nitrogen-filled glovebox ($\text{H}_2\text{O} < 1$ ppm, $\text{O}_2 < 1$ ppm). Then, the mixtures were transferred into molybdenum crucibles and fired at 1000 °C for 12 hours in a tube furnace under a reducing atmosphere (95% $\text{N}_2/5\%$ H₂).

Characterizations

The X-ray powder diffraction were acquired on Rigaku (Cu K α irradiation, 40 kV, 20 mA, $\lambda = 1.5406$ Å). The Rietveld refinement was conducted by TOPAS software. Photoluminescence excitation (PLE) and photoluminescence (PL) spectra were measured with a Hitachi F-7000 fluorescence spectrometer equipped with a 450 W Xenon lamp.

Temperature-dependent emission was collected on Hitachi F-7000 fluorescence spectrometer equipped with an Oxford-instruments OptistatDN. Diffuse reflection spectra were measured by UV-Vis-NIR spectrophotometer (SHIMADZU, UV-3600). SEM images and EDS were obtained by an ultrahigh resolution field-emission scanning electron microscope (FEI, Nova Nano SEM 430). ²⁹Si solid-state NMR spectrum was obtained on Bruker Avance III NMR spectrometer. The optical microscope photographs were required on Zeiss Axio Imager 2. Fluorescence decay curve was recorded on Hamamatsu Quantaurus-Tau C11367 fluorescence spectrophotometer. The electroluminescence (EL) spectra were measured using an integrating sphere (Labsphere) equipped with an Ocean optics spectrograph under an operating forward current.

Plant cultivation

Pak-choi cultivation and light expose experiments were conducted under different illumination conditions including present as-prepared plant lamps and the commercial white LEDs lamp. All Pak-choi seedlings were firstly grown under sunlight treatment for 14 days in the outdoor greenhouse, and then the above-mentioned two light sources were used to illuminate the plant for 10 days where the light intensity is 110 μmol m⁻²s⁻¹ photosynthetic photon flux density (PPFD). The illuminate time are from 7:00-9:00 at the morning, and 19:00-21:00 at night. The cultivation method was hydroponics using Hoagland solution. The Pak-choi without supplementary light treatment was chosen as the reference.

Fabrication of White LEDs and plant growth LED device

Prototype white LEDs devices were fabricated using a mixture of epoxy resin,

commercial $Y_3Al_5O_{12}$: Ce^{3+} phosphor, with/without the as-prepared $Li_2Ca_2Mg_2Si_2N_6$: Eu^{2+} phosphor coated on a blue ($\lambda=460$ nm) LED chip. The plant growth LED device was fabricated using $Li_2Ca_2Mg_2Si_2N_6$: Eu^{2+} phosphor and a blue LED InGaN chip ($\lambda=455$ nm).

Results and discussions

The XRD pattern of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ sample is shown in **Fig. 1a** and such a pattern is consistent with the standard card apart from several weak peaks of unknown impurity phase. To confirm the structure and estimate the content of impurity, the Rietveld structure refinement for Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ was performed and shown in **Fig. S1**, the amount of impurity phase is negligible considering the intensities. The refined structure parameters are also demonstrated in **Table S1**. The obtained reliability factors R_{wp} , R_p , and R_b demonstrate a good fitting quality. The crystal structure of Li₂Ca₂Mg₂Si₂N₆ is shown in **Fig. 1b**, the network is made up of bow-tie units $[Si_2N_6]^{10}$ which consists of edge sharing double SiN₄ connected by chains of edge sharing MgN₄. The Ca²⁺ site is in a distorted octahedral coordination surrounded by 6 nitrogen depicted in Fig. 1c. The SEM image in Fig. 1d shows aggregation and distribution randomly of phosphor particles with size less than 10 µm. It should be pointed that the phosphor faces the problem of poor water resistance, thus the oxygen content is overestimated. The atomic ratio of phosphor particle is also illustrated in Fig. 1e, it reveals an atomic ratio Ca/Mg/Si = 1.03:1:1.04 which agrees well with the formula of Li₂Ca₂Mg₂Si₂N₆. The ²⁹Si solid-state MAS nuclear magnetic resonance spectrum of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ was measured and presented in **Fig. S2**. Mostly, the chemical shift of ²⁹Si is ranging from -40 ppm to -60 ppm in nitrides. ^{20,21} There is only one sharp peak observed at -55 ppm, in line with one crystallographic Si site in the

structure of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$. Considering the crystal structure of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$, the peak at -55 ppm could be assigned to the bow-tie units $\left[\text{Si}_2\text{N}_6\right]^{10}$, which also demonstrate that the purity of the sample and no Si based impurity phase was generated.

UV-Vis diffuse reflectance spectra of host and $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6:1.0\%\text{Eu}^{2+}$ sample are presented in **Fig. 2**. The doped sample shows a broad band absorption in the blue to yellow region compared with the undoped sample, which is resulted from the $4f \rightarrow 5d$ transitions of Eu^{2+} ions. The inset presents the digital photograph of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6:1.0\%\text{Eu}^{2+}$ sample under daylight, it shows light orange body color consistent with its absorption in blue to yellow region. The host and the doped sample show identical absorption band at around $240 \sim 280$ nm, indicating it is ascribed to the host lattice absorption. The calculated bad gap is 4.26 eV (insert of **Fig. 2**) close to the value of 4.6 eV in the literature 18 , illustrating that this compound is an appropriate host for rare earth luminescence.

In the preparation of nitride phosphors, Li₃N usually plays a role as flux to improve the crystallinity.²² The effect of additional Li₃N concentration on phase and luminescence intensity are shown in **Fig. S3**, which indicates that the excess Li₃N contributes to the crystallinity of the phosphor and the optimal concentration is 120%. The photoluminescence excitation (PLE) and photoluminescence emission (PL) spectra of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ phosphor as well as SrLiAl₃N₄:Eu²⁺ (as a reference) are shown in **Fig. 3a**. The Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ phosphor shows a broad absorption band in blue region, which is consistent with diffuse reflectance spectrum (as demonstrated in **Fig. 2**), indicating it can be efficiently pumped by blue LED chip. Notably, a shoulder band at 530 nm can also be observed. To understand the origin of shoulder band, the PL and PLE spectra at 77 K were collected and presented in **Fig. S4**. The PLE

spectrum is obviously different from that at room temperature, which may be due to the Under line.²³ phonon-assisted transitions of zero phonon 460 excitation, Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ phosphor exhibits the emission peak at 638 nm with FWHM of 62 nm. The powder gives a bright red light under 365 nm UV lamp irradiation (Fig. 3a). The emission spectra of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ under different wavelength excitation are also shown in Fig. S5. The emission profile is nearly unchanged apart from luminescence intensity due to the difference of absorption value, indicating only one Eu²⁺ crystallographic site exists in Li₂Ca₂Mg₂Si₂N₆. SrLiAl₃N₄:Eu²⁺ phosphor exhibits an emission band peaking at 650 nm with FWHM of 50 nm under 460 nm excitation (Fig. 3a). As above mentioned, the blue shift of narrow band red-emitting phosphor makes a great impact on luminous efficacy of white LEDs device. The most advantage of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ phosphor lies in position of emission maximum. Compared to SrLiAl₃N₄:Eu²⁺ phosphor, Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ gives a short wavelength emission at 638 nm, resulting in less part of emission outside the human eye sensitivity (green dotted line). Differencing from Eu²⁺ ions in a highly symmetric cuboidlike environment of SrLiAl₃N₄ or in highly unusual EuN₉ coordination polyhedron of β-SiAlON,²⁴ this compound with distorted octahedral coordination for Eu²⁺ still shows relative narrow band. The possible reason is that the lattice ions like Li, Mg, and Si occupy distinct lattice sites instead of the same lattice site with a statistical distribution, inducing small Stokes shifts and restricting inhomogeneous emission band broadening, thus narrow band occurs. 25 The room temperature decay curve of Li₂Ca₂Mg₂Si₂N₆:1.0% Eu²⁺ phosphor under excitation at 460 nm, monitored at the peak of 648 nm is shown in **Fig. 3b**. The lifetime of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ is calculated to be 0.7 µs, which is in line with those determined for other Eu²⁺-doped nitridosilicate red

phosphors, such as $(Ca/Sr)AlSiN_3:Eu^{2+}$ ($\tau=0.6\sim0.8~\mu s$), $SrLiAl_3N_4:Eu^{2+}$ ($\tau=0.6~\mu s$). The images of phosphor particles were also checked by optical microscope and presented in **Fig. 3c-d**. These particles with size of 5-10 μm show red emission upon 530 nm green illumination, consistent with the results of emission spectra and SEM images.

Fig. 4a-b show the emission spectra and the emission intensity variation of $\text{Li}_2\text{Ca}_{2(1-x)}\text{Mg}_2\text{Si}_2\text{N}_6$: $x\text{Eu}^{2+}$ (x=0.5%-4.0%) samples under 460 nm excitation. The luminescent intensity is very sensitive to the concentration of Eu^{2+} , determining the optimal Eu^{2+} doping concentration is 1.0%, namely the critical concentration. As the concentration of Eu^{2+} increases, the peak wavelength stabilizes in range of 637-639 nm and the FWHM stabilizes in range of 62-67 nm. **Fig. 4d** presents the relationship of $\log(x)$ and $\log(I/x)$. The slope is calculated to be -1.8, indicating the value of θ is 5.4 close to 6. According to Dexted theory, ²⁶ the concentration quenching in $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$: Eu^{2+} phosphors most likely comes from dipole–dipole interactions.

Thermal stability is of great importance affecting the luminescence performance, it is indispensable to evaluate the thermal quenching behavior and chromaticity stability of phosphor.^{27–33} The temperature-dependent PL spectra of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ from room temperature (RT) to 473 K are depicted in **Fig. 5a-b**. With the temperature increasing, the emission intensity gradually decreases attributed to thermal quenching effect and the peak wavelength shows a slight blue-shift. **Fig. 5c** illustrates the peak and integrated intensities of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ with the increase in temperature, respectively. The integrated intensity drops to 87% at 373 K and 68% at 423 K of that initially at RT, respectively. The corresponding peak intensities were 86% and 64%, respectively. The chromaticity shift of

 $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6:1.0\%\text{Eu}^{2+}$ phosphor with different temperature is presented in **Fig. 5d**. Results indicate that the $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6:1.0\%\text{Eu}^{2+}$ has a good chromaticity stability below 373 K but exhibits an obvious chromaticity shift at high temperature.

The desirable red emission spectra of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ is an attractive candidate for high CRI white LEDs. To demonstrate the potential application of Li₂Ca₂Mg₂Si₂N₆:1.0% Eu²⁺, the white LEDs (2) device was fabricated by combining the commercial yellow phosphor (Y₃Al₅O₁₂:Ce³⁺) with the as-prepared red phosphor on a blue LED chip. And the same time, a commercial white LEDs (1) was also fabricated using a blue LED chip combined with commercial vellow Y₃Al₅O₁₂:Ce³⁺ phosphor for comparison. The electroluminescence (EL) spectra of the devices under a current of 50 mA current are presented in Fig. 6a. The conventional white LEDs (1) encounters high CCT of 7650 K and low CRI of 76, while the CRI of the white LEDs (2) can be improved to 91 and the corresponding CCT decreases to 4226 K with the addition of Li₂Ca₂Mg₂Si₂N₆: Eu²⁺ phosphor. Interestingly, the value of R9 was greatly enhanced to 90, indicating that the red spectral part was enhanced obviously. The color coordinates and digital photographs of the white LEDs devices are shown in Fig. 6b. The light changes from cold white (0.2951, 0.3208) to warm white (0.3585, 0.3159), indicating that Li₂Ca₂Mg₂Si₂N₆:1.0% Eu²⁺ is a promising candidate material for use in the fabrication of high CRI white LEDs.

Given the suitable peak wavelength and narrow band of $Li_2Ca_2Mg_2Si_2N_6$: Eu^{2+} , it can be a good candidate for plant growth LED device. Hence, red-emitting $Li_2Ca_2Mg_2Si_2N_6$: Eu^{2+} was coated onto blue LED chips (λ = 455 nm) to fabricate a plant growth LED device, which gives the emission matching well with the absorption range of chlorophyl (a or b)

(**Fig. 7c**). After being irradiated for 10 days under the same conditions using different light sources (daylight; white LEDs; plant growth LED device), the pak-choi treated by plant growth LED device shows better quality than that illuminated by daylight and white LEDs (**Fig. 7a-b**). The fresh weight, total chlorophyl content, soluble protein content, and total soluble sugar of the pak-choi were analyzed and these results are list in **Table 1**. ^{34,35} All the results indicate that Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺-based plant growth LED device is a suitable light source for enhancing the growth, increasing the accumulation of nutrients, and promoting photosynthesis in plants.

Conclusions

In summary, the desired red-emitting Li₂Ca₂Mg₂Si₂N₆:Eu²⁺ phosphors were successfully synthesized by the conventional solid-state method under atmospheric pressure. With the XRD, SEM, PLE and PL, and NMR measurements, the phase, morphology, luminescence properties, and the structure of phosphor were systematically studied. The Li₂Ca₂Mg₂Si₂N₆:Eu²⁺ phosphor with a broad absorption in blue to yellow region shows red emission peaking around 638 nm with FWHM of 62 nm, which is desirable pursuit for the balance of luminous efficacy and CRI. The aid-sintering additive of Li₃N contributes to the crystallinity and enhance luminescence intensity. The title phosphor shows good thermal quenching behavior that the integrated intensity can remain 68% of the pristine even heating up to 423 K and the quenching of emission intensity occurs when the doping concentration of Eu²⁺ exceeds 1.0%. Moreover, the warm white with low CCT (4224 K) and high CRI (91) could be garnered through combining the title

phosphor and yellow-emitting $Y_3Al_5O_{12}$: Ce^{3+} in blue LED chip. These results demonstrate that $Li_2Ca_2Mg_2Si_2N_6$: $1.0\%Eu^{2+}$ has a great potential to serve as the desired red-light component applied in high CRI white LEDs. The further study in indoor pak-choi cultivation experiment also indicates that $Li_2Ca_2Mg_2Si_2N_6$: $1.0\%Eu^{2+}$ red phosphor is a promising material applied in plant growth.

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Figure captions

Fig. 1. (a) XRD patterns of standard Li₂Ca₂Mg₂Si₂N₆ and Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ sample. (b) Crystal structure of Li₂Ca₂Mg₂Si₂N₆. The yellow spheres denote Ca; the pink spheres signify Li; the blue tetrahedrals represent SiN₄; the purple tetrahedrals are MgN₄. (c) Perspective view of the six-fold coordination of Ca²⁺ by six nitrogen. (d) SEM image of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ sample. (e) EDS results (atom%) on the Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ phosphor particle.

Fig. 2. Diffuse reflectance spectra of host and $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$:1.0% Eu^{2+} sample; the inset show the calculated band gap of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$:1.0% Eu^{2+} sample and the digital photograph of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$:1.0% Eu^{2+} powder under daylight.

Fig. 3. (a) PLE and PL spectra of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$:1.0% Eu^{2+} and $\text{SrLiAl}_3\text{N}_4$: Eu^{2+} phosphors; the dotted green curve indicates the upper limit of sensitivity of the human eye; The inset shows the digital photograph of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$:1.0% Eu^{2+} phosphor under 365 nm UV lamp; (b) The decay curve of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$:1.0% Eu^{2+} under excitation at 460 nm and monitored at 638 nm; Optical microscope images of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$:1.0% Eu^{2+} particles in the (c) daylight and (d) upon 530 nm green illumination.

Fig. 4. (a) PL spectra of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$: $x\text{Eu}^{2^+}$ ($x = 0.5\% \sim 4.0\%$); (b) The dependence of peak intensity on Eu^{2^+} concentration. (c) The variations of peak wavelengths and FWHM on Eu^{2^+} concentration. (d) Relationship between the $\log(\text{I/}x)$ and $\log(x)$ of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$: Eu^{2^+} phosphor.

Fig. 5. (a,b) Temperature-dependent PL spectra of Li₂Ca₂Mg₂Si₂N₆:1.0%Eu²⁺ under 460 nm excitation. (c) Temperature-dependent normalized integrated PL intensities, normalized peak (638 nm) intensities. (d) Chromaticity shift of Li₂Ca₂Mg₂Si₂N₆:Eu²⁺ with different temperature.

Fig. 6. (a) Electroluminescence spectra of the white LEDs fabricated by a 460 nm blue LED chip, commercial $Y_3Al_5O_{12}$: Ce^{3+} phosphor, and as-prepared $Li_2Ca_2Mg_2Si_2N_6$: $1\%Eu^{2+}$ absent (1) and present (2); (b) Chromaticity coordinates of the two white LEDs, the inset shows photographs of the two white LEDs (1,2).

Fig. 7. (a) Indoor pak-choi cultivation irradiated by different lighting sources; (b) Digital photographs of pak-choi cultivation irradiated by different lighting sources; (c) Absorption spectra of Chlorophyl (a and b) and the emission spectrum of $\text{Li}_2\text{Ca}_2\text{Mg}_2\text{Si}_2\text{N}_6$:1.0% Eu²⁺-based plant growth LED device.

Table 1. Effect of different light sources on the indicators of pak-choi (a-c represent daylight, white LEDs, and plant growth LED device, respectively)

	Fresh weight (g plant ⁻¹)	Total chlorophyl content (mg g ⁻¹)	Soluble protein content (mg g ⁻¹)	Total soluble sugar (mg g ⁻¹)
a	32.09	764.2	4.80	1.29
b	35.74	836.8	7.69	1.95
c	38.22	1073.7	6.82	2.49

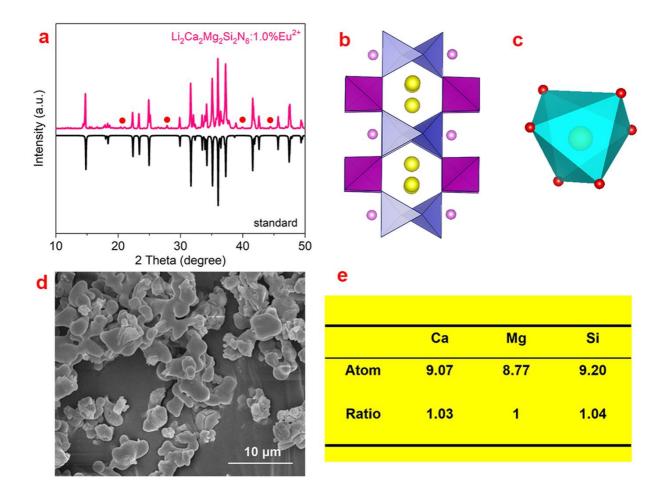


Fig. 1

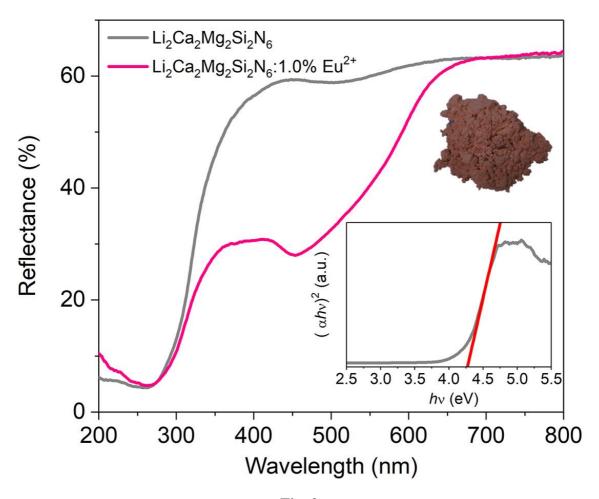


Fig. 2

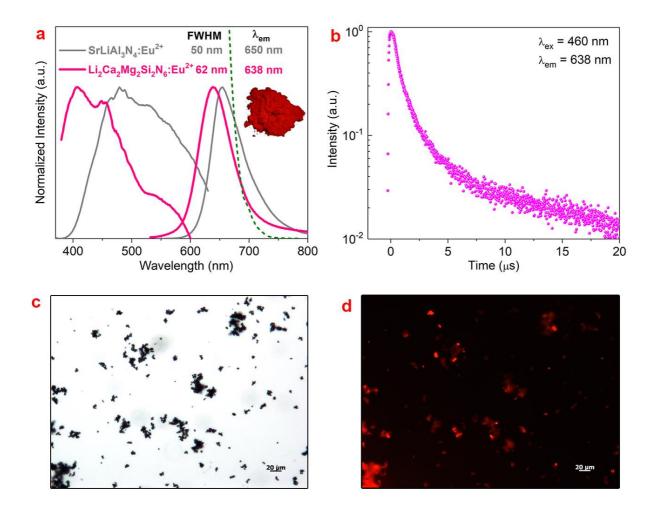


Fig. 3

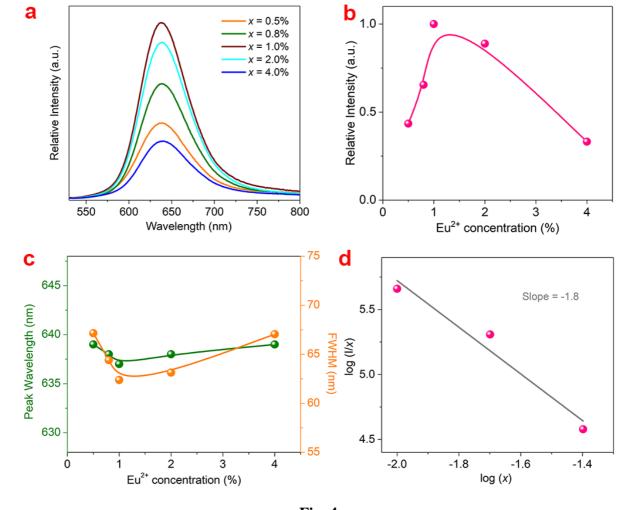


Fig. 4

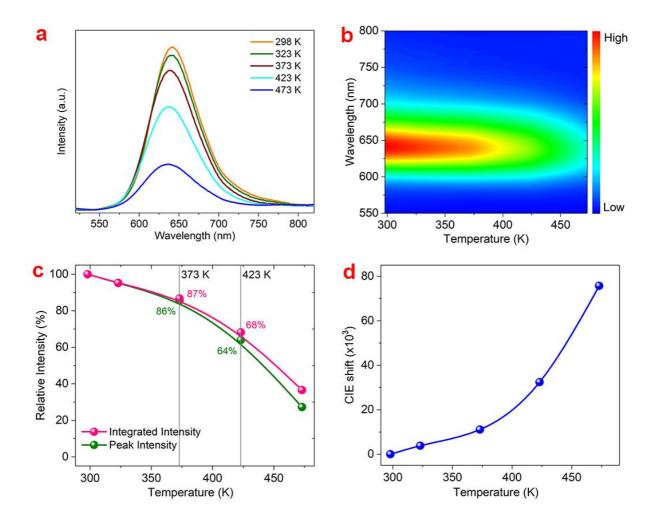


Fig. 5

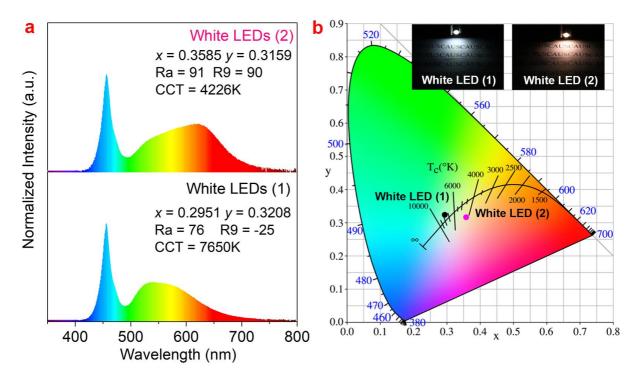


Fig. 6

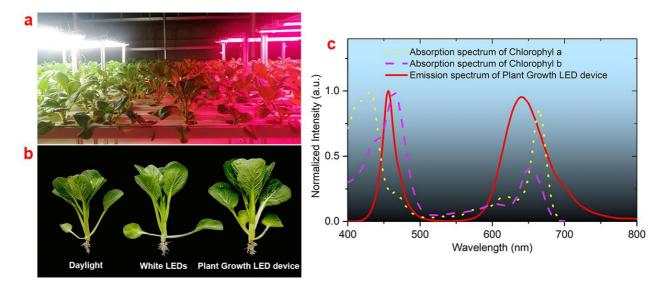


Fig. 7