


Cite this: *J. Mater. Chem. C*, 2022, **10**, 12266Enhancing the luminescence performance of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$ by substitution of Sr^{2+} for Ca^{2+} †Junlong Cao, Songsong Ding, Yunpeng Zhou, Xilin Ma and Yuhua Wang *

As reusable materials, persistent luminescence (PersL) materials have a wide range of applications in emergency response, information encryption and other fields. However, there is still a lack of commercial PersL materials with a yellow afterglow in current studies. Based on this, a series of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$ samples were synthesized by a high temperature solid-state method, and then their crystal structure, optical performance and trap properties are discussed in this work. First, EPR and XPS showed that Sr^{2+} substitution is beneficial for the generation of oxygen vacancy traps, which would improve the luminescence performance of the samples from 12 h to 46 h. Next, through PL and XANES it was found that the characteristic emission of oxygen vacancies and Eu^{2+} emission presented color-adjustable PL properties, and also produced single matrix white light, which solved the stroboscopic problem of the blue chip in LEDs. In addition, the emission of oxygen vacancies can transfer energy to Eu^{2+} excitation, which marks the first report on the energy transfer of oxygen vacancies. Finally, the ML properties of the samples were investigated, providing a new method for enhancing the luminescence properties of samples.

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Introduction

As science and technology develop by leaps and bounds, energy and environmental issues have posed an increasingly serious challenge to human life, so researchers in relevant fields are committed to seeking environmentally friendly and reusable materials. In this context, as a reusable energy storage material, long persistent luminescence materials have attracted much attention.^{1–3} Persistent luminescence (PersL) materials have unique properties for storing energy, with afterglow properties of different materials ranging from a few seconds to several days.^{4,5} Applied in coatings, inks and glass ceramics at first, PersL materials are now being used in fields including

emergency escape, defence and military, information encryption and light storage as research develops.^{6–8} In the current research studies, among PersL materials, which mainly emit short wavelength visible light, $\text{SrAl}_2\text{O}_4:\text{Eu}^{2+}, \text{Dy}^{3+}$ (green, >24 h),⁹ $\text{CaAl}_2\text{O}_4:\text{Eu}^{2+}, \text{Nd}^{3+}$ (blue, >19 h)¹⁰ and $\text{Y}_2\text{O}_2\text{S}:\text{Eu}^{3+}, \text{Mg}^{2+}, \text{Ti}^{4+}$ (orange-red, >5 h)¹¹ are more mature. These three commercial phosphors can cause damage to human eyes because most of them are cool-coloured phosphors. In fact, compared with other commercial PersL materials, most yellow PersL samples such as $\text{SrSi}_3\text{O}_5:\text{Eu}^{2+}, \text{Dy}^{3+12}$ and $\text{Ca}_2\text{ZnSi}_2\text{O}_7:\text{Eu}^{2+}, \text{Dy}^{3+13}$ have poor chemical stability, low initial brightness and short afterglow time. There is still a lack of yellow PersL phosphor materials due to inadequate research studies on the afterglow luminescence mechanism, and no practical application of yellow PersL phosphors has been reported. In this context, the research and development of novel warm-coloured yellow PersL phosphors is of great importance.

PersL fluorescent materials generally consist of a matrix and an activator, with the latter acting as a luminescence center in the material, and the same activator demonstrates different PersL properties in different substrates.^{14,15} Eu^{2+} can exhibit emission from violet to red and even near-infrared wavelengths because its 5d electron orbitals are naked and highly influenced by the crystal field.¹⁶ The PersL performance of samples singly doped with Eu^{2+} is poor, and other trivalent rare earth

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ions such as Dy³⁺ and Nd³⁺ ions need to be co-doped to improve the PersL properties of the sample. Therefore, unequal substitution of Dy³⁺ and Nd³⁺ ions would generate a large number of traps, which is beneficial to the improvement of afterglow performance.^{17,18} For example, Matsuzawa revealed that the afterglow properties of SrAl₂O₄:Eu²⁺ (green) could be improved from 10 h to 40 h by co-doping with Eu²⁺ and Dy³⁺;⁹ Wang discovered that the afterglow properties of Li₂SrSiO₄:Eu²⁺, Dy³⁺ (orange-yellow), can be promoted significantly from several minutes to 15 h by co-doping with Eu²⁺ and Dy³⁺;¹⁹ Wang *et al.* found that the afterglow properties of Sr₃SiO₅:Eu²⁺ could reach 14 h through co-doping with Eu²⁺ and Dy³⁺ (yellow).¹² However, in the present research studies, the performance of the yellow PersL materials is still not comparable to that of commercial phosphors including SrAl₂O₄:Eu²⁺, Dy³⁺ and Sr₂MgSi₂O₇:Eu²⁺, Dy³⁺, so there is an urgent need to find a warm PersL material with excellent performance.

In recent years, the design and modulation of phosphors by replacing anions or ionic groups with solid solution have become one of the research hotspots. Denalt *et al.* found that the luminescence performance of Ba_{2-x}Sr_xSiO₄:Eu²⁺ samples could be substantially improved by using the substitution of the same main group ion Sr²⁺ in Ba₂SiO₄:Eu²⁺, which is because Sr²⁺ substitution increased the thermal stability of phosphors and produced a more rigid crystal structure;²⁰ Xia pointed out that the the fluorescence performance of Ca₂(Al_{1-x}Mg_x)(Al_{1-x}Si_{1+x})O₇:Eu²⁺ could see significant improvements by replacing the [Al³⁺-Al³⁺] structural unit with [Mg²⁺-Si⁴⁺] to facilitate the electron-phonon coupling effect;²¹ Lin *et al.* substituted CaYAlO₄:Eu²⁺, Eu³⁺ phosphors with [Si⁴⁺-Al³⁺], [Ca²⁺-Y³⁺] to make the octahedral lattice of AlO₆ shrink and the polyhedral lattice of CaO₉ expand, so that some of the Eu³⁺ ions are reduced to Eu²⁺ and the luminous color of the samples is adjustable;²² Zhou noted that the near-infrared spectrum coverage of BaMgGa_xAl_{10-x}O₁₇:Cr³⁺ was greatly enhanced through the solid solution with Ga³⁺ and Al³⁺ and the half-peak width increased from 100 nm to 300 nm, and the substitution of Ga³⁺ for Al³⁺ led to an increase in CrO bond stress, a decrease in the bond length and an increase in energy level cleavage, resulting in a significant improvement in the optical properties of the sample;²³ Ma *et al.* investigated a 2000% increase in initial afterglow luminance and 350% increase in afterglow time in the BaGa_{2-x}Zn_xO₄:Pr³⁺ series of samples;²⁴ Xie found that the trap depth of Y₃Al_{5-x}Ga_xO₁₂:Ce³⁺, V³⁺ (x = 0–3) could be adjusted by solid solution of Al³⁺ by Ga³⁺, resulting in the change of the crystal field intensity. Meanwhile, the solid solution of Ga³⁺ led to the adjustment of the luminescence color of the sample from green to yellow, which has a good application prospect for optical storage.²⁵

Therefore, in this work a high temperature solid-phase method was used to successfully prepare the Ca_{2-x}Sr_xBO₃:Cl:Eu²⁺, Dy³⁺ solid solution borate system by replacing Ca²⁺ with Sr²⁺ ions of the same main group. The cationic solid solution substitution was adopted to improve the PersL performance of the phosphor so that its PersL performance could be increased from 12 h to 46 h. Finally, samples with different Sr²⁺

solid solution concentrations were investigated and some explanations were given for the mechanical luminescence properties produced by the samples after solid solution.

Materials and methods

Materials and synthesis

A series of samples were prepared by using the traditional high temperature solid phase method. Raw materials including CaCO₃ (A. R.), SrCO₃ (A. R.), H₃BO₃ (A. R.), CaCl₂ (A. R.), Eu₂O₃ (A. R.) and Dy₂O₃ (A. R.) were taken on the basis of the stoichiometric ratio, and then they were put into an agate mortar to mix evenly after adding a certain amount of alcohol. Next, the mixture was placed in a crucible under the reducing atmosphere of a high-temperature furnace. The samples were cooled to room temperature in the high temperature furnace after heating to 950 °C for 6 hours. Finally, the samples were collected for further research.

Measurements and characterization

The samples were subjected to analysis on a Rigaku D/Max-2400 X-ray diffractometer (XRD) with Cu K α radiation running from 10° to 80° and the working state of 40 kV and 60 mA. The structure of the experimental product was analyzed using the general structure analysis system (GSAS). HRTEM was performed on an FEI TecnaiF30 TEM operating at 300 kV. EDX and HAADF were used in conjunction to measure the elemental composition. The diffuse reflectance spectrum (DRS) using BaSO₄ as a reference substance was recorded on a spectrophotometer. An FS5 spectrofluorometer was adopted to measure photoluminescence (PL), photoluminescence excitation (PLE), PersL spectra and luminescence decay curves of samples. The samples were irradiated under an ultraviolet lamp at 254 nm for 10 min to record their PersL emission spectra, PersL decay curves, TL spectra and PersL images of the samples. The 3D thermoluminescence (TL) spectrum of the sample was obtained using a LTTL-3DS 3D TL spectrometer. The electron paramagnetic resonance (EPR) test was carried out using a JES-FA300 instrument. Analysis of the characteristic emission of Eu ions from samples was performed using the BL02U1 in the Shanghai Synchrotron Radiation Facility. It is worth noting that all samples were measured in a stable environment at room temperature.

Results and discussion

Structure and morphology

Fig. 1a shows the XRD patterns of Ca_{2-x}Sr_xBO₃Cl (0 ≤ x ≤ 30%) under different Sr²⁺ substitution concentrations. As the substitution concentration of Sr²⁺ increases, an impurity phase appears in the XRD patterns. The XRD patterns of the samples are pure single-phases except when the substitution concentration of Sr²⁺ is higher than 20%. The SrBO₃ impurity phase occurs when the substitution concentration of Sr²⁺ is above 20% (the impurity phase SrBO₃ is marked with “*”). It is worth

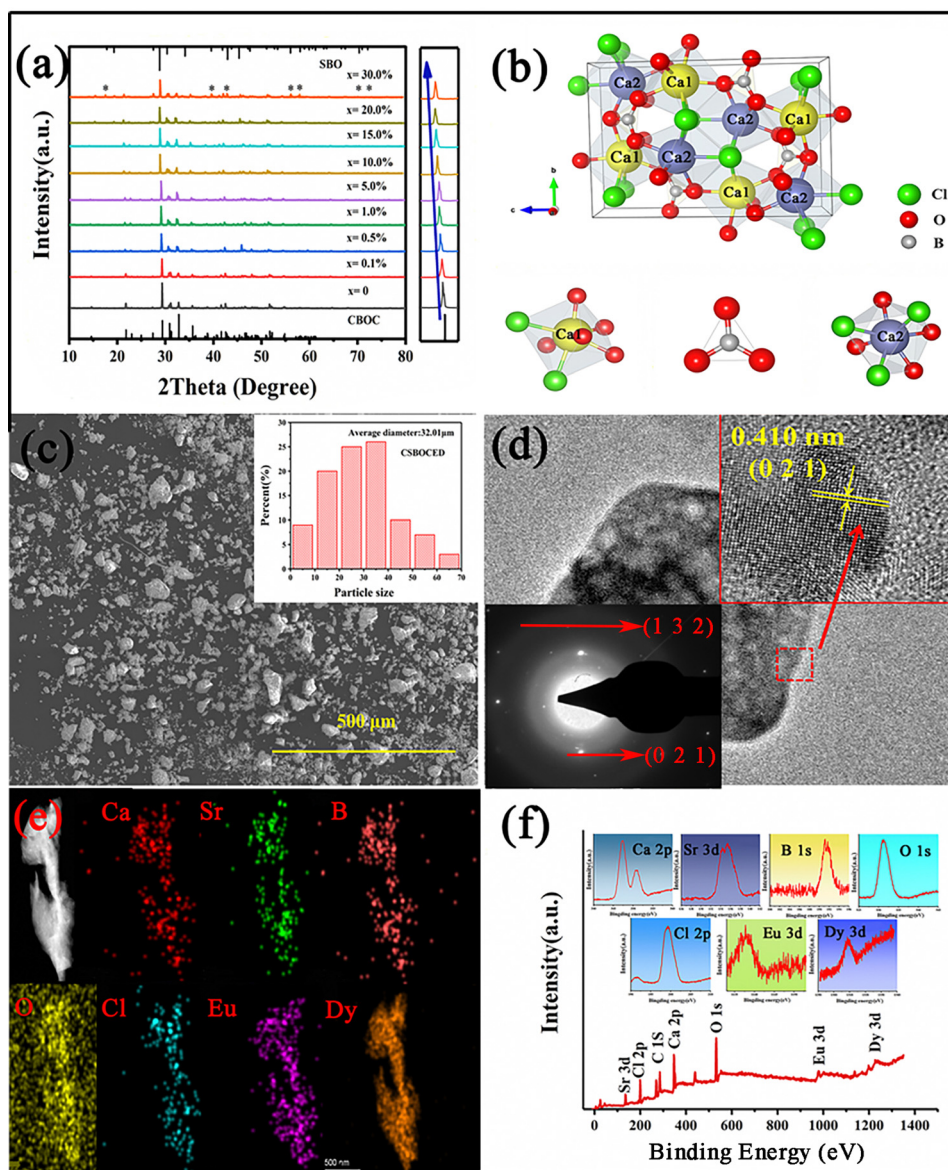


Fig. 1 (a) The XRD patterns of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}$ ($0 \leq x \leq 30\%$); (b) the structural model of $\text{Ca}_2\text{BO}_3\text{Cl}$ and the two Ca^{2+} site positions in the crystal structure; (c) the SEM image of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$ (the inset depicts the particle size distribution of the sample); (d) the TEM of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$ (the inset shows the HR-TEM particle size distribution of the sample); (e) the HAADF-STEM and elemental analysis of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$; and (f) XPS full spectrum and fine spectrum of each element in $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$.

noting that the peaks in some positions of the XRD patterns appear because the main phase $\text{Ca}_2\text{BO}_3\text{Cl}$ and impurity phase SrBO_3 overlap to some extent. The presence of the impurity phase will affect the luminescence performance of the main phase, so its luminescence performance will be no longer investigated when the substitution concentration of Sr^{2+} exceeds 20%. It can be seen that the diffraction peaks in the XRD pattern move in the small angle direction with the increase of Sr^{2+} doping (as shown in the illustration on the right), which is because Sr^{2+} (CN = 7, 1.21 Å) can replace the lattice Ca^{2+} ions when doping because as two elements of the same main group, the ionic radii of Sr^{2+} and Ca^{2+} are similar in size.²⁶ In order to obtain the changes in the crystal

structure of the samples after replacing Ca^{2+} with Sr^{2+} of different concentrations, the structure of the $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}$ ($x = 0, 10\%, 20\%$) matrix was optimized by using GSAS, as exhibited in Fig. S1 (ESI[†]). The three parameters χ^2 , R_{wp} and R_p obtained by refining and optimizing the structure of the main phase can demonstrate that the structure optimization result of the main phase has good reliability. The crystal structure of the main phase obtained by structural optimization is shown in Table S1 (ESI[†]). At the same time, the XRD patterns of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$ ($0 \leq x \leq 20\%$) with different Sr^{2+} concentrations are presented in Fig. S2 (ESI[†]). It was found that there was no impurity phase in the series of samples, so their optical properties were studied subsequently.

According to the above refined structure, the crystal structure of $\text{Ca}_2\text{BO}_3\text{Cl}$ was explored. Fig. 1b shows the crystal structure of the $\text{Ca}_2\text{BO}_3\text{Cl}$ - $\text{Ca}_2\text{BO}_3\text{Cl}$ crystal belonging to the $P_{121/C1}(14)$ space group with the monoclinic system. As can be seen, there are two Ca positions in $\text{Ca}_2\text{BO}_3\text{Cl}$, namely Ca1 and Ca2. The B^{3+} ion is connected to three O^{2-} , which is a typical sp^2 hybrid orbital. At the same time, Ca1 connects 5 O^{2-} and 2 Cl^- while Ca2 connects 4 O^{2-} and 3 Cl^- . Eu^{2+} (CN = 7, $r = 1.2 \text{ \AA}$) and Dy^{3+} (CN = 7, $r = 0.97 \text{ \AA}$) parameters are similar to those of Ca^{2+} (CN = 7, 1.06 \AA), so Eu^{2+} and Dy^{3+} can replace the position of Ca^{2+} when doping. But the Ca^{2+} ion has two lattice places, which means that Eu^{2+} has two possibilities for substitution. Lattice occupation will be discussed for the photoluminescence spectra below.²⁷

A scanning electron microscope (SEM) and a transmission electron microscope (TEM) were used to characterize the morphology of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$, as shown in Fig. 1c and d. It can be seen from the SEM image that the particle morphology of the sample is irregular, but its particle size is relatively uniform. The distribution diagram of particle size in the inset demonstrates that the particle size of the sample roughly ranges from 20 μm to 40 μm , with an average size of 32.01 μm , which is suitable for producing excellent afterglow properties. Fig. 1d displays the TEM image of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$, and the inset is the HR-TEM image and SAED pattern of the sample. According to the HR-TEM image, the crystal lattice of the sample can be clearly observed and the sample has good crystallinity.²⁸ After a certain measurement, the plane spacing of the sample is 0.410 nm, which belongs to the (0 2 1) crystal plane of the sample. In Fig. 1e and f, high-angle annular dark field (HAADF-STEM) and X-ray photoelectron spectroscopy (XPS) were used to analyze the elemental

composition and valence state of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$. From the HAADF, it can be seen that the sample does not have any impurity elements. The illustration depicts the mapping of each element in the sample. XPS showed that all the elements in $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ are in the expected valence state, which will be discussed in detail in the thermoluminescence section.

Photoluminescence properties

Fig. 2a shows the emission spectra of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ excited at 338 nm. Just as illustrated in the figure, the emission spectra of the series of samples consist of two peaks, among which the emission peak at 585 nm is due to an electron transition of Eu^{2+} from $4f^7$ to $4f^65d^1$.²⁹ With the increase of the Sr^{2+} substitution concentration, the full width at half-maximum increases from 20 nm to 41 nm; the emission peak at 470–750 nm is attributed to Eu^{2+} characteristic emission, and full width at half-maximum decreases from 132 nm to 103 nm. And the IQE of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ was 38.06%. The X-ray absorption near-edge spectroscopy (XANES) spectra of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+}$ are shown in Fig. 2b. By comparing with the XANES spectra of the $\text{BAM}:\text{Eu}^{2+},\text{Eu}_2\text{O}_3$ standard sample, it is found that the peak value of the sample is 6974 eV, which is completely consistent with the peak value of the standard sample, indicating that the characteristic emission peak of the sample is attributed to Eu^{2+} .³⁰ The source of the emission peak at 420 nm is worth exploring. According to the current literature, it is mainly caused by Eu^{2+} occupying different Ca^{2+} positions, or by the emission peak of the oxygen vacancies generated in the solid solution process of Ca^{2+} by Sr^{2+} . To further prove the above two possibilities, a series of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}$ samples were prepared under a vacuum environment

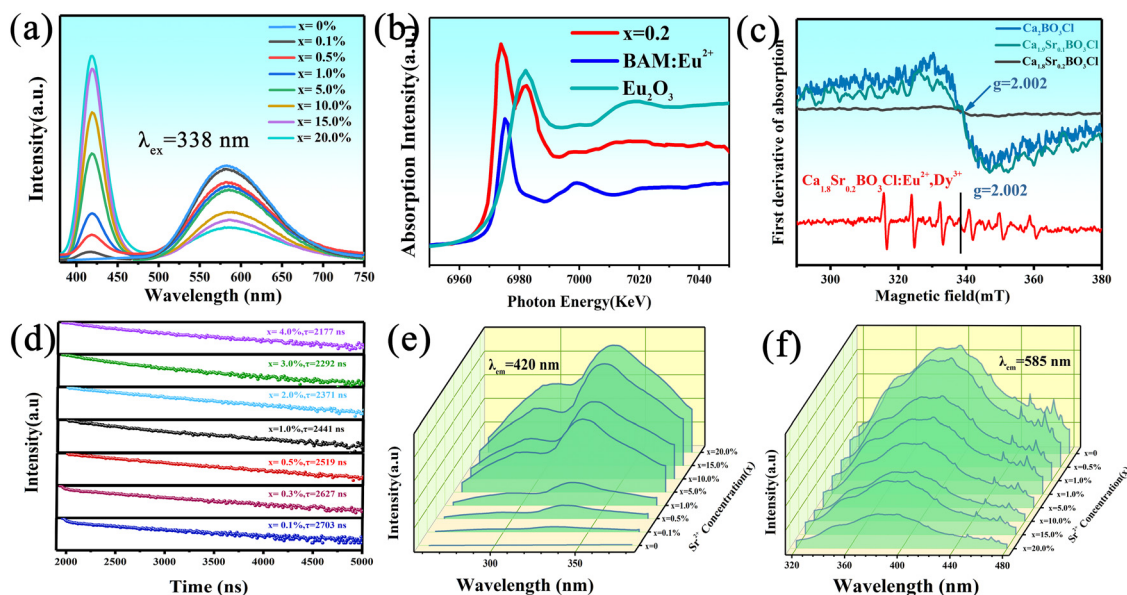


Fig. 2 (a) Photoluminescence emission spectra of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$; (b) XANES spectra of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$; (c) EPR spectra of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}$ ($x = 0, 10\%, 20\%$) and $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$; (d) the decay curves of Eu^{2+} in $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+}$ ($0.1\% \leq x \leq 4\%$); (e) the excitation spectra of oxygen vacancies of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$; and (f) the excitation spectra of Eu^{2+} in $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$.

and reducing atmosphere, respectively, and then their luminescence properties were tested. As shown in Fig. S3 (ESI[†]), solid line in reducing atmosphere, dotted line in vacuum atmosphere. The series of samples without Eu²⁺ doping also have an emission peak at 420 nm, and the emission peak intensity of the samples prepared under a reducing atmosphere is significantly higher than that under a vacuum atmosphere. Therefore, the first possibility is ruled out and the second possibility must be the reason. Electron paramagnetic resonance (EPR) was performed on the Ca_{2-x}Sr_xBO₃Cl series of samples prepared under a reducing atmosphere, and the test results are displayed in Fig. 2c. According to EPR spectra, the resonance signal of the oxygen vacancies in the Ca_{2-x}Sr_xBO₃Cl series of samples was $g = 2.002$.³¹ As the amount of Sr²⁺ solid solution concentration increases, the number of oxygen vacancies increases because Sr²⁺ generates a certain amount of oxygen vacancy traps during solid solution, which is caused by oxygen ions entering the lattice gap and generating traps in the lattice. Moreover, the EPR test was carried out on the Ca_{1.8}Sr_{0.2}BO₃Cl:Eu²⁺,Dy³⁺ sample, which presented the resonance signal not only for oxygen vacancies but also for Eu²⁺, supporting the presence of Eu²⁺ in the sample. The electronic structure of Eu²⁺ is 4f⁷ (the ground state of 8S_{7/2}), and the spin quantum number of Eu²⁺ is $S = 7/2$, which makes the EPR spectrum of the sample present 7 peaks. Meanwhile, it can be seen that the 7 spectral lines of the sample are asymmetrical due to anisotropy caused by the splitting of the crystal field of the sample.^{32,33}

It is worth noting that as the amount of Sr²⁺ increases, as shown in Fig. S4 (ESI[†]) CIE coordinates (Commission Internationale de l'Éclairage), the sample can move from yellow to white and thus a warm white phosphor could be obtained. In addition, with the increase of Sr²⁺ substitution concentration, emission peaks of oxygen vacancies gradually increase while the emission peak of Eu²⁺ decreases, which may be relevant to the change of crystal field caused by Sr²⁺ substitution. As the substitution concentration of Sr²⁺ increases, the crystal field symmetry of the matrix is reduced, which directly lowers the Eu²⁺ emission intensity. However, in view that the excitation peaks of Eu²⁺ and the emission peaks of the oxygen vacancies of the samples overlap to some extent, energy transfer occurs.^{34,35} Therefore, energy transfer was used to compensate for the reduction in the characteristic emission peak of Eu²⁺ caused by changes in the crystal field, thus decreasing the reduction rate of Eu²⁺ emission intensity. Fig. 2d shows the decay curves for the series of samples of Ca_{1.8}Sr_{0.2}BO₃Cl:xEu²⁺. The calculated average lifetime is marked in the decay curve. It can be seen that the average lifetime decreases as the concentration of Eu²⁺ increases because the oxygen vacancies will transfer energy to Eu²⁺.³⁶ The characteristic of the sample lies in that white light can be obtained by combining the sample and the UV chip instead of mixing with other phosphors. The electroluminescence spectra under excitation of the 340 nm UV chip are shown in Fig. S5 (ESI[†]), with the best emission peaks located at 420 nm and 585 nm, which is in line with the photoluminescence spectra. On the successive UV chip, the

emission peak at 420 nm in the electroluminescence (EL) spectrum of the sample decreases over time while the emission peak at 585 nm remains unchanged, indicating that there is energy transfer from the emission peak of the oxygen vacancies to the emission of Eu²⁺ and that the LED device has good stability.

Fig. 2e shows the excitation spectra of Ca_{2-x}Sr_xBO₃Cl:Eu²⁺,Dy³⁺ emission at 420 nm. The main peak at 338 nm is produced by the oxygen vacancies, and the emission intensity gradually increases as the concentration of Sr²⁺ solid solution increases. Fig. 2f shows the excitation peak of Eu²⁺, which belongs to the electron transition of Eu²⁺ from 4f⁶5d¹ to 4f⁷. Fig. S6 (ESI[†]) displays the normalized excitation spectra of oxygen vacancies and Eu²⁺, and it is interesting that the excitation spectra of the samples are adjustable after normalizing their excitation peak, which is beneficial for expanding the application fields of the phosphor.

As the concentration of Sr²⁺ solid solution increases, the emission is gradually blue-shifted because the substitution of Ca²⁺ by Sr²⁺ is a large ionic radius substitution, which gives birth to lattice expansion, weakening of the crystal field, an increase in the band gap and reduction in energy level cleavage.³⁷ At the same time, DRS and density functional theory were used to calculate the band gap of the series of samples. The calculated values from DRS and density functional theory (DFT) are exhibited in Fig. S1 and S6 (ESI[†]) respectively. DRS and DFT calculations demonstrate that with the increase of the Sr²⁺ substitution concentration, the band gap value of the samples increases from 4.95 eV to 5.05 eV. This increase in band gap not only affects the energy level difference between the conduction band of the matrix and the 5d excited state of Eu²⁺, but also influences the luminescence intensity of Eu²⁺, which is in accordance with the reduction of Eu²⁺ characteristic emission. However, the reduction rate of Eu²⁺ emission will be decreased by using energy transfer. The specific calculation formula is as follows:³⁸

$$(\alpha h\nu)^n = A(h\nu - E_g) \quad (1)$$

$$\alpha = \frac{(1 - R)^2}{2R} \quad (2)$$

where $h\nu$ represents the energy of the incident light and α represents the absorption coefficient. It can be concluded that the solid solution of Sr²⁺ widens the band gap, which is consistent with the results obtained from density functional theory.

Persistent luminescence (PersL) properties

In order to characterise the afterglow properties of the samples, a series of samples were tested for their afterglow properties. All samples were pre-irradiated with a 254 nm UV lamp for 10 min before being tested for afterglow decay, with a threshold value of 0.32 mcd m⁻².³⁹ Fig. 3a demonstrates the afterglow decay curve of the Ca_{2-x}Sr_xBO₃Cl:Eu²⁺,Dy³⁺ series of samples. It can be seen that the afterglow time of samples without solid solution only reaches more than 10 h, but their afterglow time

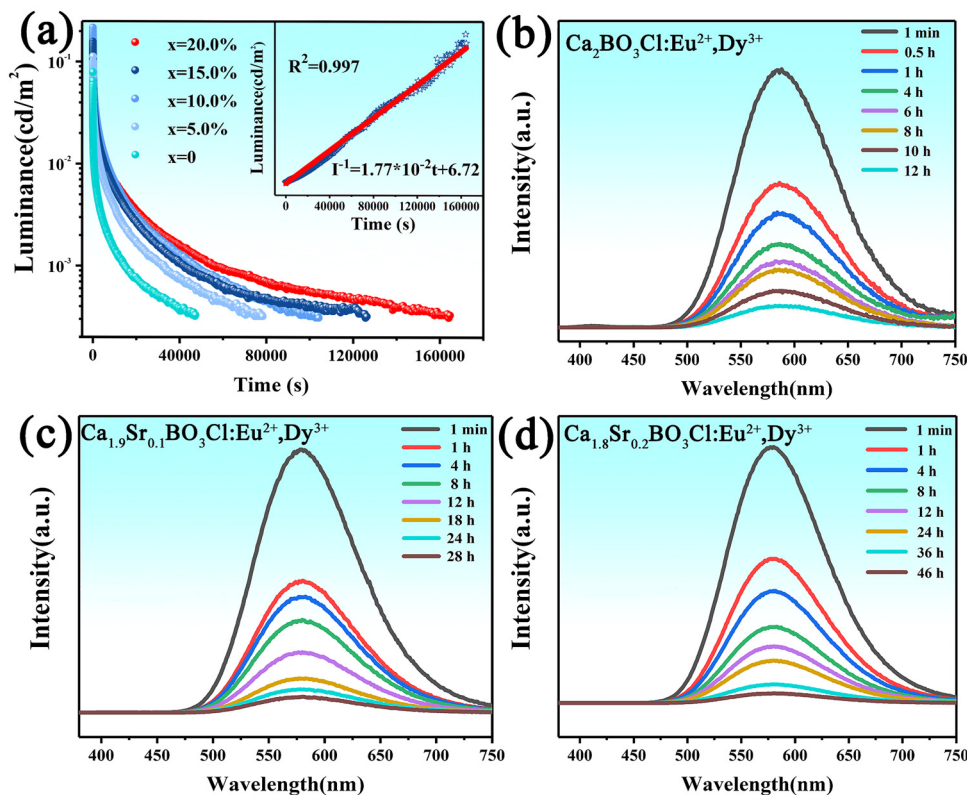


Fig. 3 (a) Afterglow decay curves of the $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ series of samples (the inset indicates the inverse of the afterglow intensity *versus* time for $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$) and (b–d) afterglow spectra of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ ($x = 0, 10\%, 20\%$).

can be increased to 46 h with the increase of Sr^{2+} solution concentration, which is improved by up to 400%. For the sample with the best afterglow time, a linear fitting of the inverse of the afterglow intensity *versus* time was performed, and the inset represents the inverse of the afterglow intensity (Γ^{-1}) *versus* the afterglow decay time (t) for the $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ sample, and $\Gamma^{-1}t$ can be well fitted by⁴⁰

$$\Gamma^{-1} = 1.77 \times 10^{-2}t + 6.26. \quad (3)$$

The fitting parameter is $R^2 = 0.997$, indicating that the fitting is reliable. As can be seen from the figure, the relationship appears to be linear, which in turn indicates that the traps acting on the afterglow are continuously distributed and that the release of the traps is a continuous and linear process.^{41,42} We will analyse this in the subsequent TL spectra.

The afterglow spectra of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ are exhibited in Fig. 3b–d. Also, in the previous work, the oxygen vacancies exhibited an emission peak at 420 nm, but there was no characteristic emission of oxygen vacancies in the afterglow spectrum. When the solid solution concentration of Sr^{2+} increases, the number of oxygen vacancies increases and the free electrons captured by traps strengthen, which in turn enhances the afterglow performance of the sample. As for the formation principle of oxygen vacancies, it will be mentioned in the following discussion.

Trap analysis

After performing tests on the afterglow properties of the series of samples, the present work has carried out some tests on the thermoluminescence (TL) spectra of the series of samples. In general, the afterglow phenomenon is inextricably linked to the type of trap in the sample. The type and number of traps have a strong influence on the PeSL performance, so we tested the three-dimensional TL spectra of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ samples and the results are presented in Fig. 4a. It can be seen that the samples have a wide trap distribution in the temperature range of 300–500 K and that the trap distribution of the samples is continuously distributed.⁴³

The trap depth (E) is calculated with the following equation⁴⁴

$$E = T_m/500 \quad (4)$$

where T_m represents the temperature peak in the TL spectrum. After calculation, the trap energy level of the $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ sample is mainly distributed in the range of 0.60–1.00 eV and such a trap depth is suitable for afterglow generation. Fig. 4b–d show the TL spectra of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ ($x = 0, 10\%, 20\%$) samples after 10 min of excitation and placed for different testing times. It is obvious that as time increases, the intensity of the shallow trap peak decreases and gradually approaches that of the deep trap peak. This result also indicates that the trap distribution of the samples is continuous. The TL spectra of the samples demonstrate that

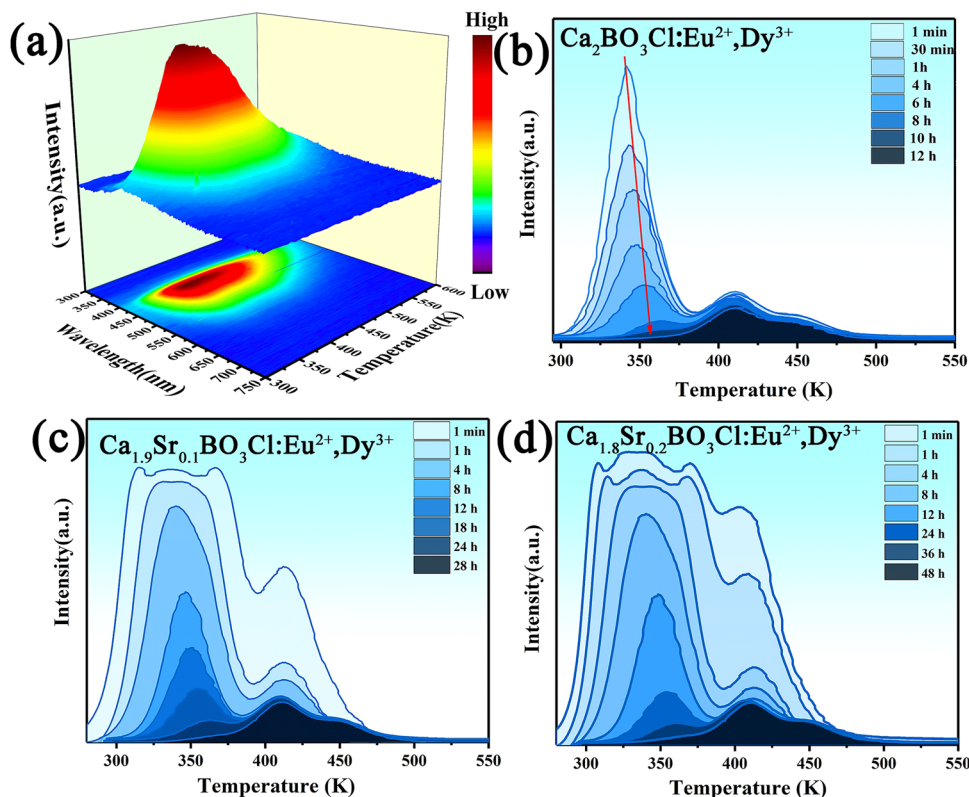


Fig. 4 (a) 3D-TL spectrum of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$; TL spectra of (b) $\text{Ca}_2\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$; (c) $\text{Ca}_{1.9}\text{Sr}_{0.1}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$; and (d) $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$.

there is no increase in the type of traps in the samples but a change in the number. In the series of samples, traps arise due to Ca^{2+} equivalent and unequal substitution of Eu^{2+} and Dy^{3+} , with Dy^{3+} replacing the Ca^{2+} in the lattice to form traps.

In addition, according to the principle of trap formation, there are many intrinsic traps in samples during the preparation by the high-temperature solid-phase method, as illustrated in Table 1. It is believed that the afterglow of the sample without solid solution is caused by many trap energy levels in the sample, such as oxygen vacancies and so on. XPS was used to analyze the chemical environment of each element in

$\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$, as illustrated in Fig. S8 (ESI[†]). Chemical environment analysis of oxygen elements in the samples discovered that the 1s electronic orbital of the oxygen atom can be fitted into three peaks, among which O_I is attributed to lattice oxygen, O_{II} to oxygen vacancy and O_{III} to adsorbed oxygen.⁴⁵ With the increase of the Sr^{2+} solid solution concentration, the number of oxygen vacancies increases dramatically, so the number of oxygen ions around Ca^{2+} decreases due to the reduction of the coordination number, leading to the change of the binding energy.⁴⁶ At the same time, the increase in oxygen vacancies leads to the greater energy difference between the trap energy level and the bottom of the host conduction band, as well as the greater trap energy level depth. When the trap depth increases, more electrons are captured, which also causes an increase in the number of electrons in the shallow traps. In addition, as the trap depth deepens, the synergistic effect of the increased number of electrons and the slower escape rate of electrons from the traps makes the sample have a stronger initial brightness and longer afterglow time. Therefore, the luminescence intensity and decay time of PersL materials can be improved by adjusting the distribution and depth of traps *via* the increase of Sr^{2+} substitution concentration.

After the substitution of Sr^{2+} for Ca^{2+} , the number of oxygen vacancy traps increased and the band gap of the sample widened from 4.95 eV to 5.05 eV, resulting in an increase in the afterglow properties of the $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$

Table 1 Point defect reactions for $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ LPL phosphors

Type (intrinsic)	Reaction	
Frenkel		$\text{Ca}_{\text{Ca}} \rightarrow \text{V}_{\text{Ca}}'' + \text{Ca}_i^{\bullet\bullet}$
		$\text{B}_{\text{B}} \rightarrow \text{V}_{\text{B}}''' + \text{B}_i^{\bullet\bullet\bullet}$
		$\text{O}_{\text{O}} \rightarrow \text{V}_{\text{O}}^{\bullet\bullet} + \text{O}_i''$
		$\text{Cl}_{\text{Cl}} \rightarrow \text{V}_{\text{Cl}}' + \text{Cl}_i'$
Schottky	$2\text{Ca}_{\text{Ca}} + \text{B}_{\text{B}} + 3\text{O}_{\text{O}} + \text{Cl}_{\text{Cl}} \rightarrow \text{Ca}_2\text{BO}_3\text{Cl} + \text{V}_{\text{Ca}}'' + \text{V}_{\text{B}}''' + \text{V}_{\text{O}}^{\bullet\bullet} + \text{V}_{\text{Cl}}'$	
Type (extrinsic)	Charge compensation	Reaction
$\text{Eu}^{2+} \rightarrow \text{Ca}^{2+}$	Interstitial	$\text{Eu}^{2+} + \text{Ca}^{2+} \rightarrow \text{Eu}_{\text{Ca}}$
$\text{Dy}^{3+} \rightarrow \text{Ca}^{2+}$	Interstitial	$2\text{Dy}^{3+} + 3\text{Ca}^{2+} \rightarrow 2\text{Dy}'_{\text{Ca}} + \text{V}_{\text{Ca}}^{\bullet\bullet}$
$\text{Sr}^{2+} \rightarrow \text{Ca}^{2+}$	Interstitial	$\text{Sr}^{2+} + \text{Ca}^{2+} \rightarrow \text{Sr}_{\text{Ca}}$

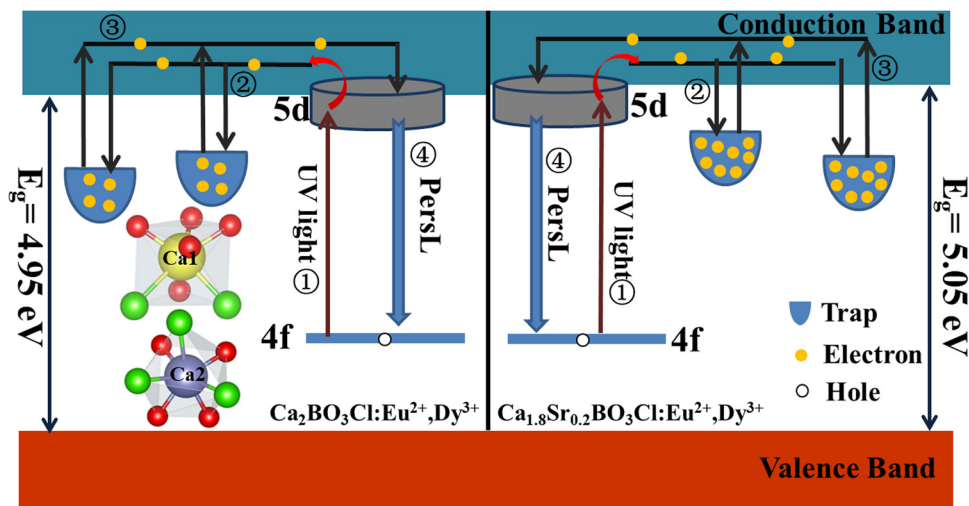


Fig. 5 The schematic diagram of the PersL mechanism of $\text{Ca}_2\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ and $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$.

sample. Based on the above experimental data and speculation, the schematic diagram of afterglow generation was proposed, as shown in Fig. 5. Under the excitation of the UV light, the electrons in the ground state entered the excited state of the CB and become free electrons, as illustrated in ①. Next, these free electrons were caught by the traps, as displayed in ②. Then, when the excitation light source was removed, the electrons captured by the shallow traps and deep traps returned to the CB under the influence of thermal disturbance, as exhibited in ③. Finally, they returned to the ground state and produced yellow PersL, which is in line with the Dorenbos afterglow mechanism model. The applications of some samples in mechanoluminescence and life will be discussed below.²⁸

Mechanoluminescence

$\text{Ca}_2\text{BO}_3\text{Cl}$ is an asymmetric crystal structure. Asymmetric piezoelectric materials will produce piezoelectricity under pressure because the energy band of the material will be tilted and the deep trap will be released, which will compound with the luminescence center to produce mechanoluminescence (ML) properties.⁴⁷ The traps of the sample caused by Eu^{2+} and Dy^{3+} doping will lead to ML, and meanwhile the substitution of Ca^{2+} by Sr^{2+} at different concentrations will lead to an increase in the number of traps of the sample, which will improve its ML performance. To explore the ML properties of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$, the samples in this part were prepared by mixing a certain proportion of PDMS (polydimethylsiloxane) with a curing agent. Fig. 6a demonstrates the 120% stretching of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$, and the ML phenomenon after stretching was captured by the camera. Fig. 6b displays that when the force of 2 N was applied to the sample, there was yellow light emission in the applied area, and the sample could exhibit good ML properties. Fig. 6c exhibits the ML spectra of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ under different pressures. Within the test range (0.1–2 N), the luminescence intensity of the sample increases with the increase of pressure, and the peak position

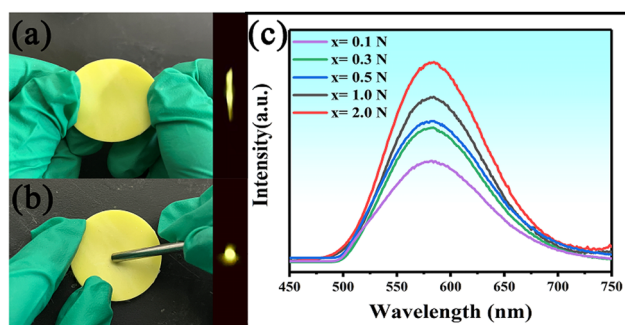


Fig. 6 (a) The image of a tensile test on $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$; (b) the image of the sample with an applied force of 2 N; (c) the ML spectra of $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ under different force.

of the ML spectra remains at 585 nm, which is attributed to the $4f^7-4f_6^5d^1$ electron transition of Eu^{2+} . The sample has good potential for engineering applications such as bridge fracture detection, tensile testing, biocompatibility and self-powered fields due to its ML properties.

Application of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$

White light emitting diodes (wLEDs) are generally a mixture of red, green and blue (RGB). The sample $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ has emission peaks at 420 nm and 585 nm under ultraviolet excitation, and then emits a yellow afterglow. On the basis of this performance, an LED device based on $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+},\text{Dy}^{3+}$ was successfully prepared using a 340 nm UV chip. In addition, the CIE of the pc-LED was tested and the CIE color coordinate values of pc-LED were 0.339, 0.307. Fig. 7a–c show the pictures of the LED without power and under working conditions, respectively. After 10 min of excitation, the device showed a yellow afterglow when the power was disconnected. Fig. 7d–g display the yellow afterglow of the device placed for different times after the power was disconnected. The results show that the device emits white light in the powered state and

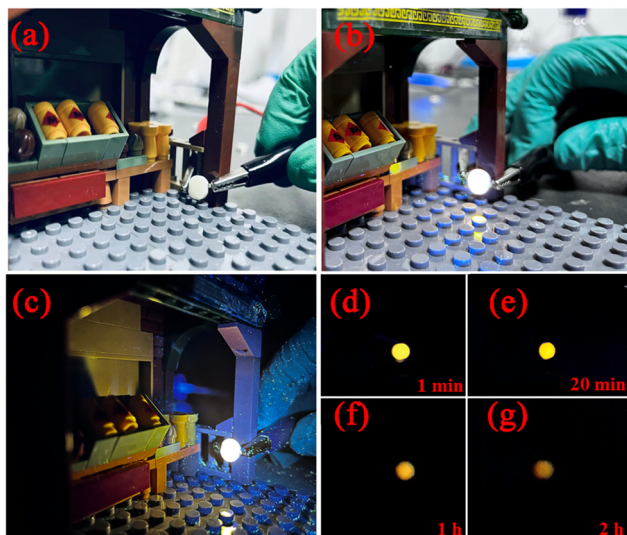


Fig. 7 (a and b) WLED without power supply and working under natural light; (c) wLED working in the night environment; and (d–g) photos of long afterglow placed for different times after power failure.

after disconnection it emits a yellow afterglow with a colour temperature of 3824 K, which conforms to the human sleep index. Thus it is concluded that $\text{Ca}_{1.8}\text{Sr}_{0.2}\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$ can be used as a sleeping lamp.

Conclusion

In conclusion, a new solid solution yellow PersL material $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$ was successfully designed and prepared by a high temperature solid phase method. PL and EPR spectra demonstrated that Sr^{2+} solid solution could increase the oxygen vacancies in samples, resulting in an emission peak at 420 nm in the series of samples. Eu^{2+} doping produced an emission peak at 585 nm, which interacted with the emission peak at 420 nm to produce white light. With the increase of the Sr^{2+} solid solution concentration, the trap depth and width of $\text{Ca}_{2-x}\text{Sr}_x\text{BO}_3\text{Cl}:\text{Eu}^{2+}, \text{Dy}^{3+}$ are improved significantly, leading to an enhancement in the afterglow performance from 12 h to 46 h and an improvement of ML properties of the samples. Therefore, the sample with the most excellent properties in the series of samples has large potential for application in engineering.

Author contributions

Junlong Cao: experimental design, data collection and interpretation, and writing the manuscript. Songsong Ding and Yunpeng Zhou: data curation and software. Xilin Ma: idea refinement and additional proposal. Yuhua Wang: funding acquisition, writing, review & editing, and supervision.

Conflicts of interest

The authors declare no conflicts of interest.

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