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## Enhanced persistent properties of $\text{Mn}^{2+}$ activated $\text{CaZnOS}^\dagger$

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The red emitting  $\text{CaZnOS}:\text{Mn}^{2+}$  long persistent phosphor was successfully prepared via a conventional high-temperature solid-state reaction method. The persistent luminescence emission of  $\text{Mn}^{2+}$ -doped  $\text{CaZnOS}$  phosphor was investigated for the first time. Rare-earth trivalent ions were used as co-dopants to obtain a brighter red afterglow and longer persistence time. The optical properties of the as-prepared samples have been investigated systematically by employing photoluminescence spectroscopy, persistent luminescence spectroscopy, afterglow decay curves and thermoluminescence curves. In light of the experimental results of this study, a possible mechanism of persistent luminescence was illustrated and discussed in detail. These investigations provide a new and efficient long persistent phosphor.

## Introduction

The phenomenon of luminescence has fascinated people since the earliest times. Persistent luminescent is a particular luminescence type for which the emission is delayed by minutes, hours or even days after the excitation has stopped.<sup>1</sup> The luminescent materials with persistent luminescence properties are put into the context of emerging applications such as in imaging storage, optical memory and *in vivo* imaging.<sup>2,3</sup> Since the efficient persistent phosphor  $\text{SrAl}_2\text{O}_4:\text{Eu}^{2+}$ ,  $\text{Dy}^{3+}$  has been reported,<sup>4</sup> numerous novel persistent materials have been gaining attention among many researchers.<sup>5–10</sup> More recently, materials with emission in the deep red or in the near-infrared are desired for surveillance applications and *in vivo* medical imaging.<sup>11</sup> However, many researchers and publications on these persistent luminescent materials have focused on divalent europium as the doping ion. The broadband emission of  $\text{Eu}^{2+}$  is strongly dependent on the host material, more exactly, on the centroid shift and the strength of the crystal field acting on the ions.<sup>12</sup> This mixture of effects leads to the so-called red shift, and the value depends strongly on the structure of the host compound and the local coordination of the europium dopant ion. It is quite common to obtain a blue or green afterglow using oxide host compounds, but it is much more challenging to find a suitable host solid with appropriate red shift, in order to obtain orange or red persistent luminescence.

Moreover, red emitting  $\text{Eu}^{2+}$  doped persistent phosphors, such as  $\text{CaS}:\text{Eu}^{13}$  and  $(\text{Ca}, \text{Sr})_2\text{Si}_5\text{N}_8:\text{Eu}$ ,<sup>14,15</sup> the choice is limited and the host lattices are chemically unsteady or hard to prepare. At present, great efforts have been made to develop other doping ions in order to obtain efficient red radiating persistent materials.

Calcium zinc oxysulfide ( $\text{CaZnOS}$ ) is an intermediate production recovery of zinc from its sulfide by a carbothermal reduction in the presence of lime. This novel quaternary oxysulfide was first reported.<sup>16</sup> Later on, Igiehon and coworkers<sup>17,18</sup> tried to explore the products of carbothermal reduction of iron and zinc sulfides and to determine the crystal structures of  $\text{CaZnOS}$  and  $\text{CaFeOS}$ . In 2003, Petrova *et al.* refined the lattice parameters and determined the space group of this zinc calcium oxysulfide using the Rietveld profile analysis method. Subsequently, its synthesis, structure, and electrical properties were investigated in detail.<sup>19,20</sup> Recently, the photoluminescence properties of  $\text{CaZnOS}:\text{Mn}^{2+}$  phosphor were reported.<sup>21</sup>  $\text{Mn}^{2+}$ -activated  $\text{CaZnOS}$  shows a single symmetric narrow red emission band in the wavelength range of 550–700 nm with the peak centered at 614 nm. Very recently, the multi-stress sensitive elastico-mechanoluminescence characteristics of  $\text{CaZnOS}:\text{Mn}^{2+}$  phosphor were for the first time discovered.<sup>22</sup> It can simultaneously sense and image different mechanical stimuli including ultrasonic vibration, impact, friction, and compression with an intense emission, indicating the practical prospect on stress sensors. Upon reviewing the literature, it is clear that there is no research on the afterglow properties of  $\text{Mn}^{2+}$  doped  $\text{CaZnOS}$  phosphors. Thus, the investigation of the afterglow properties of  $\text{CaZnOS}:\text{Mn}^{2+}$  is a very interesting work.

In this article,  $\text{Mn}^{2+}$  doped  $\text{CaZnOS}$  phosphors were successfully prepared by solid-state reaction in an argon atmosphere. We report some new and preliminary phenomena

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of  $\text{CaZnOS:Mn}^{2+}$  phosphors. The red long afterglow emission enhancement of  $\text{CaZnOS:Mn}^{2+}$  by co-doping  $\text{Ce}^{3+}/\text{Pr}^{3+}$  into the phosphor is also reported. The photoluminescence (PL), afterglow and thermoluminescence (TL) properties of  $\text{CaZnOS:Mn}^{2+}$  and  $\text{CaZnOS:Mn}^{2+}$ ,  $\text{RE}^{3+}$  ( $\text{RE} = \text{Ce}^{3+}$ ,  $\text{Pr}^{3+}$ ) phosphors were investigated in detail. It was found that the additive  $\text{Ce}^{3+}$  ions extended the persistent time much longer. This result is of course an exciting phenomenon in the investigation field of the red afterglow phosphors.

## Experimental

Undoped and doped  $\text{CaZnOS}$  powder samples were prepared by the conventional high-temperature solid-state reaction method. The 4 N pure  $\text{CaCO}_3$ ,  $\text{ZnS}$ ,  $\text{MnCO}_3$ ,  $\text{CeF}_3$  and  $\text{Pr}_2\text{O}_3$  were employed as the starting materials. Phosphor with molar compositions of  $\text{Ca}_{0.99}\text{Zn}_{0.99}\text{OS:0.01Mn}^{2+}, 0.01\text{RE}^{3+}$  ( $\text{RE} = \text{Ce}^{3+}$ ,  $\text{Pr}^{3+}$ ) were thoroughly mixed and ground in an agate mortar, and subsequently sintered in an alumina crucible at 1273 K for 4 h in a horizontal tube furnace under a flowing argon atmosphere. After sintering, these samples were gradually cooled down to room temperature in the heating system. Finally, the samples crystallization behavior were examined by powder X-ray diffraction (XRD, Bruker D8) with  $\text{Cu K}\alpha$  radiation ( $\lambda = 1.54056 \text{ \AA}$ ) operated at 36 kV and 30 mA in the  $2\theta$  range from  $10^\circ$  to  $80^\circ$  with a step size of  $0.02^\circ$  for a scanning rate of  $2^\circ \text{ min}^{-1}$ . The microstructure and morphological information of samples were measured using a scanning electron microscope (SEM, XL-30, Phillips) coupled with energy dispersive spectroscopy (EDS). X-ray photoelectron spectroscopy (XPS) measurements were performed by X-ray photoelectron spectrometer (Escalab 250Xi, Thermo Scientific) equipped with an  $\text{Al K}\alpha$  line within the 0–1300 eV energy range. The photoluminescence and persistent luminescence spectra of the as-synthesized materials were recorded using a fluorescence spectrometer (F-7000, Hitachi) equipped with a 150 W Xe lamp as the radiation source. Afterglow decay curves were measured by long afterglow phosphor tester after 15 min excitation with an unfiltered Xe arc lamp at 1000 lux (PR-305, Zhejiang Tri-color instrument Co. Ltd., Zhejiang, China). The thermoluminescence (TL) behaviors were investigated by a microcomputer controlled TL dosimeters (FJ427-AL; Beijing Nuclear Instrument Factory, Beijing, China) with a fixed heating ratio  $1 \text{ K s}^{-1}$ . Before measuring the TL, the samples were pre-irradiated by 365 nm UV light for 10 min. All measurements were carried out at room temperature except for the TL curves.

## Results and discussion

After prepared through solid-state reaction, the samples were ground. The size and shape of the particles are important when constructing of the optical device. The morphology and the size of a typical  $\text{Mn}^{2+}$ -dope  $\text{CaZnOS}$  sample were examined by SEM, as shown in Fig. 1. It can be clearly seen that most of the particles are distributed in the crystal size range of 1–3  $\mu\text{m}$ , together with some larger particles up to 5  $\mu\text{m}$ . And the phosphors are composed of small, roughened spherical grains.

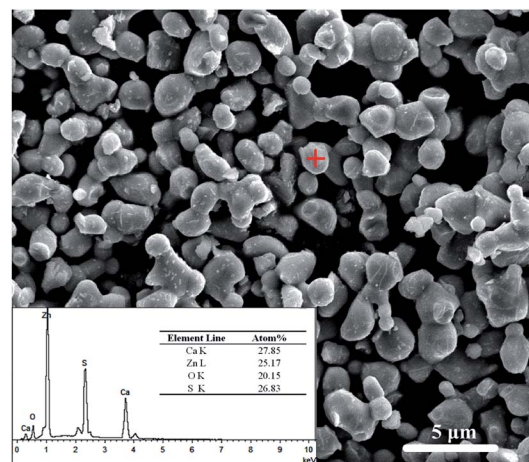


Fig. 1 SEM image of the  $\text{CaZnOS:Mn}^{2+}$  sample. Inset: EDS spectrum of the sample.

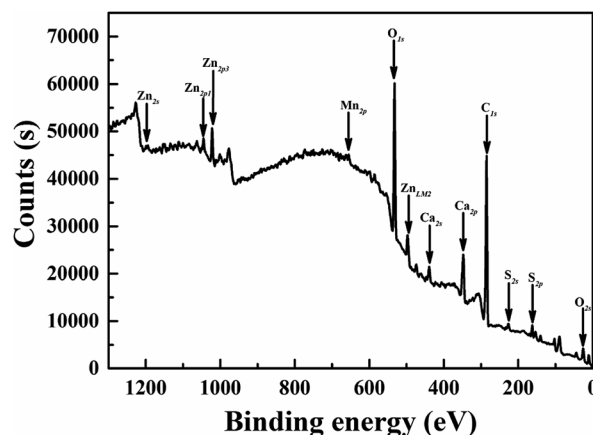


Fig. 2 XPS survey spectrum of  $\text{CaZnOS:Mn}^{2+}$  sample.

Composition and quantity of all elements in a spot were analyzed by using Energy Dispersive X-ray Spectrometer (EDS), as shown in the inset of Fig. 1. The EDS spectrum of the  $\text{CaZnOS:Mn}$  sample revealed that it is composed of Ca, Zn, O and S. However, Mn element was not detected due to its low-concentration doping. The EPR spectra of the samples in the present study were measured and illustrated as Fig. S1.† The well resolved signature of  $\text{Mn}^{2+}$  ions observed in the EPR spectra confirms the existence of  $\text{Mn}^{2+}$  ions in samples. In order to confirm the constituent elements of the synthesized  $\text{CaZnOS}$  phosphor, X-ray photoelectron spectroscopy (XPS) measurement was also carried out, as shown in Fig. 2. The XPS spectrum contains Ca, Zn, O, C, S and Mn signal.<sup>23</sup> As in the previous EDS testing result, we could not accurately determine the amount of Mn incorporated in the present XPS measurements because of the low sensitivity of the Mn signal.

Fig. 3a shows the X-ray powder diffraction patterns of as-prepared samples and the standard ICSD card of  $\text{CaZnOS}$  for comparison. It can be seen from Fig. 3a that all the diffraction peaks are in good agreement with the standard data of  $\text{CaZnOS}$  (ICSD#245309). It indicates that the obtained samples are well



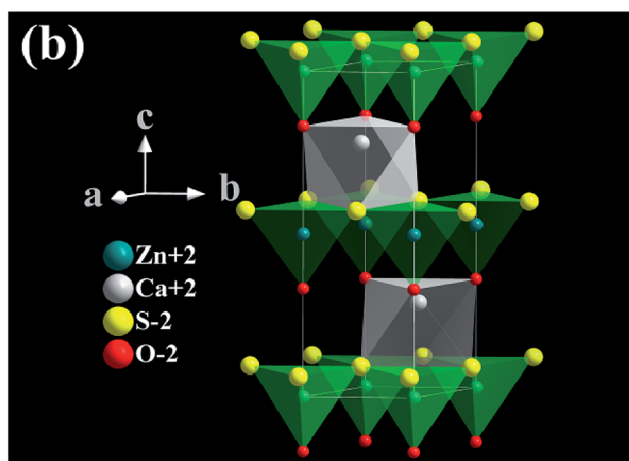
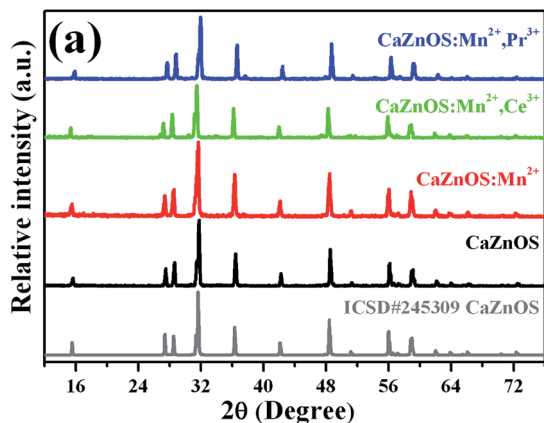


Fig. 3 (a) X-ray diffraction patterns of the as-prepared samples and the pattern of ICSD#245309 for CaZnOS; (b) crystal structure of CaZnOS.

crystallized and the incorporation of  $\text{Mn}^{2+}/\text{Ce}^{3+}/\text{Pr}^{3+}$  ions almost has no significant influence on the host lattice. The structure of CaZnOS is investigated in detail by Clarke *et al.*<sup>20</sup> Based on the above article, the oxysulfide compound crystallizes in the hexagonal space group  $P6_3mc$  with  $a = 3.75726(3)$  Å,  $c = 11.4013(1)$  Å,  $V = 139.388(2)$  Å<sup>3</sup> and  $Z = 2$ . The crystal structure of CaZnOS and the coordination environments of the  $\text{Ca}^{2+}/\text{Zn}^{2+}$  sites are plotted in Fig. 3b. It can be seen that the structure framework of CaZnOS host lattice is composed of layers of  $\text{ZnS}_3\text{O}$  tetrahedral and  $\text{CaS}_3\text{O}_3$  octahedral. Obviously,  $\text{ZnS}_3\text{O}$  tetrahedra all aligned with Zn–O bond vectors parallel and directed parallel to  $c$  axis and linked at all their S-containing vertices, yielding  $[\text{ZnS}_{3/3}\text{O}]^{2-}$  layers.  $\text{Ca}^{2+}$  ions occupy distorted octahedral interstices between these  $[\text{ZnS}_{3/3}\text{O}]^{2-}$ .

To further examine the crystal structure and sites in this compound, Rietveld structure refinements of these typical samples have been performed using the GSAS program, and the final refinement patterns of the typical  $\text{CaZnOS:Mn}^{2+}$  sample are shown in Fig. 4. Correspondingly, these refinement parameters are shown in Table 1. The good fitness of  $\chi^2$ ,  $R_{\text{wp}}$ , and  $R_p$  suggests that the refined results are reliable.

The effective ionic radii for the given coordination number (CN) are listed in Table 2. In view of similar ionic radii and

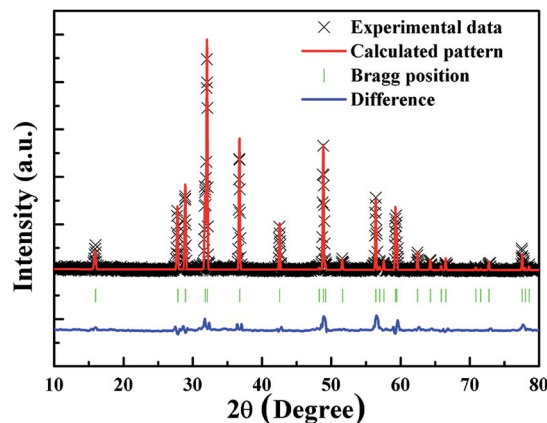


Fig. 4 Observed (crossed) and calculated (red line) XRD patterns and the difference profile (blue line) of the Rietveld refinement of  $\text{CaZnOS:Mn}^{2+}$ . Bragg reflections are shown as green vertical bars.

Table 1 Refinement data for CaZnOS and  $\text{CaZnOS:Mn}^{2+}$

Formula	CaZnOS	$\text{CaZnOS:Mn}^{2+}$
Space group	$P6_3mc$	$P6_3mc$
$a/\text{\AA}$	3.75610(3)	3.75611(2)
$b/\text{\AA}$	3.75610(3)	3.75611(2)
$c/\text{\AA}$	11.4011(2)	11.4012(7)
$V/\text{\AA}^3$	139.30	139.31
$\alpha = \beta/\text{\AA}^\circ$	90	90
$\gamma/\text{\AA}^\circ$	120	120
$Z$	2	2
$R_{\text{wp}}/\%$	3.99	3.76
$R_p/\%$	2.55	2.71
$\chi^2$	6.58	6.36

valence state, the  $\text{Mn}^{2+}$  ions can be dissolved into the host lattice completely by the substitution for  $\text{Zn}^{2+}$  ions. The ionic radii of  $\text{Ce}^{3+}$  and  $\text{Pr}^{3+}$  are similar to that of  $\text{Ca}^{2+}$ , but much larger than that of  $\text{Zn}^{2+}$ . As a logical consequence, the  $\text{Ca}^{2+}$  sites will be substituted by  $\text{Ce}^{3+}/\text{Pr}^{3+}$ .

The diffuse reflectance spectra of nondoped and doped CaZnOS are comparatively demonstrated in Fig. 5. It is noticed that the CaZnOS host shows high reflection in the range of 450–800 nm and then starts to decline from 450 nm to 250 nm due to the host absorption. For  $\text{CaZnOS:Mn}^{2+}$  and  $\text{CaZnOS:Mn}^{2+},\text{Pr}^{3+}$ , the similar reflection are observed in the range of 400–800 nm, and then they decrease obviously to 200 nm. For  $\text{Mn}^{2+}$ ,  $\text{Ce}^{3+}$  co-

Table 2 Ionic radii (Å) for given coordination number (CN) of positive ions

Ion	CN	Ionic radius (Å)
$\text{Zn}^{2+}$	4	0.60
$\text{Ca}^{2+}$	6	1.00
$\text{Mn}^{2+}$	4 (HS)	0.66
$\text{Mn}^{2+}$	6 (LS)	0.67
$\text{Mn}^{2+}$	6 (HS)	0.83
$\text{Ce}^{3+}$	6	1.01
$\text{Pr}^{3+}$	6	0.99



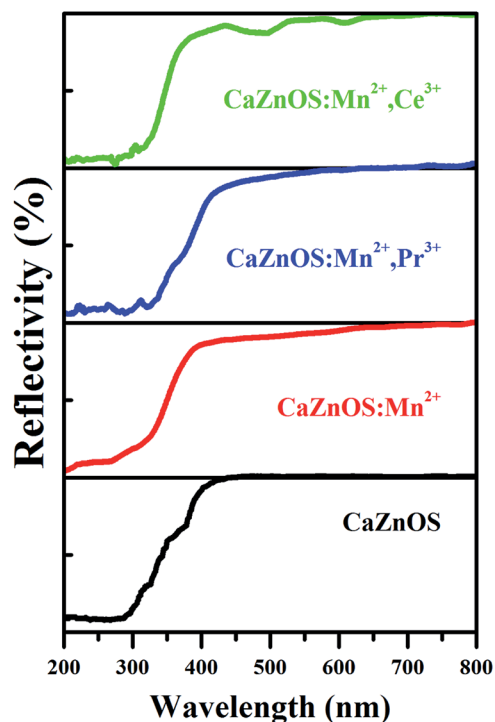


Fig. 5 Diffuse reflectance spectrum of CaZnOS; CaZnOS:Mn<sup>2+</sup>; CaZnOS:Mn<sup>2+</sup>,Ce<sup>3+</sup>; CaZnOS:Mn<sup>2+</sup>,Pr<sup>3+</sup>.

doped CaZnOS, the high reflection ends up until 450 nm, and then declines quickly to 320 nm. Two absorption bands can be observed (at about 500 and 600 nm), which should be related to the absorption of doped Ce<sup>3+</sup>.<sup>24,25</sup>

According to a previous report, the optical band gap ( $E_g$ ) of the sample can be determined by the following equation provided by Tauc:<sup>26</sup>

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

The  $\alpha$  in eqn (1) is absorption coefficient  $h$  is Planck's constant,  $\nu$  is the light frequency,  $A$  is the absorption constant,  $n$  is a constant exponent that determines the type of optical transitions.  $n = 3$  represents indirect forbidden transition,  $n = 2$  represents indirectly allowed transition,  $n = 3/2$  represents direct forbidden transition and  $n = 1/2$  represents direct allowed transition. In the CaZnOS host,  $n = 1/2$ .<sup>20,24</sup>

Absorption coefficient  $\alpha$  of the sample at different wavelengths is proportional to  $F(R)$ . Based on Kubelka–Munk model,<sup>27</sup>  $F(R)$  can be expressed by the following equation:<sup>28</sup>

$$F(R) = (1 - R)^2/2R \quad (2)$$

where  $R$  is the reflectance of the material and can be estimated from the experimental reflectance in our CaZnOS sample. Fig. 6 presents the plot of  $[F(R)h\nu]^2$  vs.  $h\nu$  according to eqn (1). We can get that the  $E_g$  value from the intercepts of the fitted straight lines using the linear extrapolations of  $[F(R)h\nu]^2 = 0$ , as shown in Fig. 6. For undoped CaZnOS, its optical band gap is calculated to be about 3.91 eV, which is in fair agreement with the

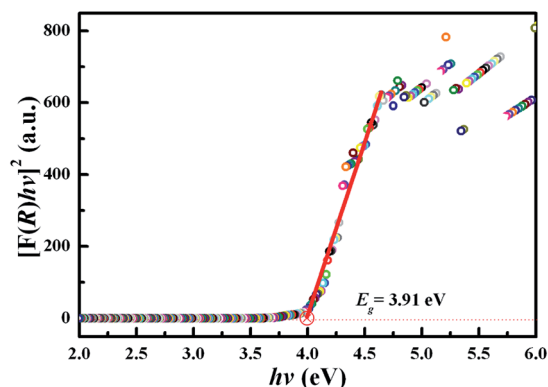


Fig. 6 The  $[F(R)h\nu]^2$  variation versus photon energy  $h\nu$  and the  $E_g$  of CaZnOS host.

Table 3 Summary of optical band gap of CaZnOS in different articles

Optical band gap of CaZnOS						
Value (eV)	3.71	3.88	3.90	4.00	4.16	3.91
Ref.	20	29	30	21	22	This work

value of the previously reported.<sup>20–22,29,30</sup> Table 3 shows the optical band gap of CaZnOS in previous articles.

Fig. 7 exhibits the excitation and emission spectra of as-prepared samples. Fig. 7a shows the excitation and emission spectra of host material CaZnOS. In the spectra, CaZnOS shows a very broad green light emission band between 400 and 650 nm with a dominated peak at about 500 nm. An asymmetrical excitation band centered at 376 nm is observed when monitoring host emission at 500 nm. According to a previous report,<sup>31</sup> the green emission can be assigned to the recombination of donors ( $V_O$ ) and acceptors ( $V_{Zn}$ ) which are associated with native defects. The photoluminescence of the Mn<sup>2+</sup> doped CaZnOS exhibits a relatively broad orange-red emission band caused by  ${}^4T_1-{}^6A_1$  transition of Mn<sup>2+</sup> (Fig. 7b). Normally, the tetrahedral Mn<sup>2+</sup> will give out the green emission,<sup>32,33</sup> this red emission observed is ascribed to further splitting of the two level system of Mn<sup>2+</sup> ion into more levels in a distorted tetrahedral coordination environment resulting in the first excited-state shifting to lower energy. In addition to the strong  ${}^4T_1-{}^6A_1$  emission from Mn<sup>2+</sup>, there is a weak emission band located at 530 nm. This band is most likely due to the Mn<sup>2+</sup>-related trap levels introduced into host lattice CaZnOS.<sup>26</sup> Interestingly, it is completely disappeared by codoping Pr<sup>3+</sup> or Ce<sup>3+</sup> into CaZnOS:Mn<sup>2+</sup> (Fig. 7d and c). We will examine this situation in more detail below. The excitation spectra of CaZnOS:Mn<sup>2+</sup> extended a broad range of wavelength, which could be ascribed to the host lattice absorption and a small part of weak d–d transition of Mn<sup>2+</sup>. Similarly, the excitation spectra of CaZnOS:Mn<sup>2+</sup>,M (M = Pr<sup>3+</sup>, Ce<sup>3+</sup>) also consist of two parts, one is the excitation band of host lattice (200–325 nm) and the other one is d–d transition of Mn<sup>2+</sup> (325–550 nm). The former is much stronger than the latter. In addition, if compared with





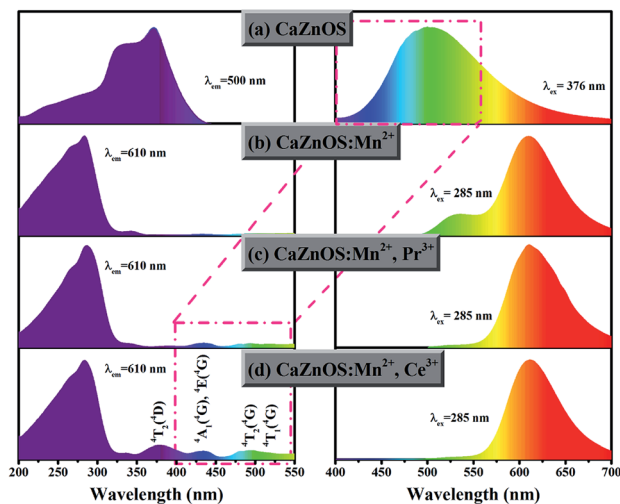


Fig. 7 Comparison of the excitation and emission spectra from (a) CaZnOS, (b) CaZnOS:Mn<sup>2+</sup>, (c) CaZnOS:Mn<sup>2+</sup>, Pr<sup>3+</sup>, (d) CaZnOS:Mn<sup>2+</sup>, Ce<sup>3+</sup>.

CaZnOS:Mn<sup>2+</sup>, the intensities of Mn<sup>2+</sup> (d-d transition) absorption peaks of CaZnOS:Mn<sup>2+</sup>, Ce<sup>3+</sup> increased obviously. These peaks at approximately 380 nm, 436 nm and 495 nm can be attributed to the transitions of <sup>6</sup>A<sub>1</sub> (<sup>6</sup>S) to <sup>4</sup>T<sub>2</sub> (<sup>4</sup>D), [<sup>4</sup>E (<sup>4</sup>G), <sup>4</sup>A<sub>1</sub> (<sup>4</sup>G)] and <sup>4</sup>T<sub>2</sub> (<sup>4</sup>G), respectively. According to Dexter theory,<sup>34</sup> an efficient energy transfer requires a spectral overlap between the donor emission and the acceptor excitation. Fig. 7 (the red bounding box) demonstrates a significant spectral overlap between the emission bands of CaZnOS host and the excitation peaks of the Pr<sup>3+</sup>/Ce<sup>3+</sup> codoped CaZnOS:Mn<sup>2+</sup> sample. As a result, there could exist efficient energy transfer from host to Mn<sup>2+</sup> in CaZnOS:Mn<sup>2+</sup>, M (M = Pr<sup>3+</sup>, Ce<sup>3+</sup>). These results give strong evidence for why the emission intensities at about 530 nm in CaZnOS:Mn<sup>2+</sup>, M (M = Pr<sup>3+</sup>, Ce<sup>3+</sup>) decreased obviously. By codoping with Pr<sup>3+</sup> or Ce<sup>3+</sup> can significantly improve the intensity of Mn<sup>2+</sup> emission peak.

Fig. 8 displays persistent luminescence emission spectra recorded at different times after the 285 nm excitation source is switched off. As can be seen in Fig. 8a–c, the persistent luminescence intensity of CaZnOS:Mn<sup>2+</sup>, M (M = Pr<sup>3+</sup>, Ce<sup>3+</sup>) are remarkably stronger than that of CaZnOS:Mn<sup>2+</sup> in the same test conditions. It is found that the emission peaks at about 610 nm and the spectral features in the persistent luminescence spectra remain unchanged and are consistent with the photoluminescence emission. Moreover, no other typical emission which belongs to the Pr<sup>3+</sup> or Ce<sup>3+</sup> are observed in the persistent luminescence emission spectra (Fig. 8b and c). These give evidence that the long afterglow characteristics in all cases are from the same <sup>4</sup>T<sub>1</sub>–<sup>6</sup>A<sub>1</sub> transition of Mn<sup>2+</sup>. It should be noted that the CaZnOS host also shows weak persistent luminescence emission. However, the persistent luminescence emission centered at 530 nm from the host in the CaZnOS:Mn<sup>2+</sup> system is very weak (Fig. 8a).

One of the most important result of the present work is that we have observed obvious enhancement and longer phosphorescence when RE<sup>3+</sup> (Pr<sup>3+</sup>, Ce<sup>3+</sup>) co-doped into CaZnOS:Mn<sup>2+</sup>. Accordingly, the afterglow decay curves of as-prepared samples after irradiation by an artificial light have been conducted in Fig. 9a. Their initial intensity is normalized for comparison. The afterglow intensity of all samples exhibits a rapid decay and subsequential long-lasting phosphorescence. Clearly, the RE<sup>3+</sup> (Pr<sup>3+</sup>, Ce<sup>3+</sup>) co-dopants can increase the intrinsic positive defects and result in longer the duration time. In particular, it is very exciting to note that the afterglow intensity of the CaZnOS:Mn<sup>2+</sup>, Ce<sup>3+</sup> sample can be recorded for about 1790 s by definition of 0.32 mcd m<sup>−2</sup> after the 1000 lux artificial light promoted (the inset of Fig. 9a). As shown in Fig. 9b and c, the afterglow decay curves of the RE<sup>3+</sup> co-doped CaZnOS:Mn<sup>2+</sup> samples are plotted as a function of the reciprocal afterglow intensity (*I*<sup>−1</sup>) versus time (*t*). Both of the curves can be well fitted by the linear equations. The linear dependence of *I*<sup>−1</sup> vs. *t* indicates that the afterglow of the samples are probably caused

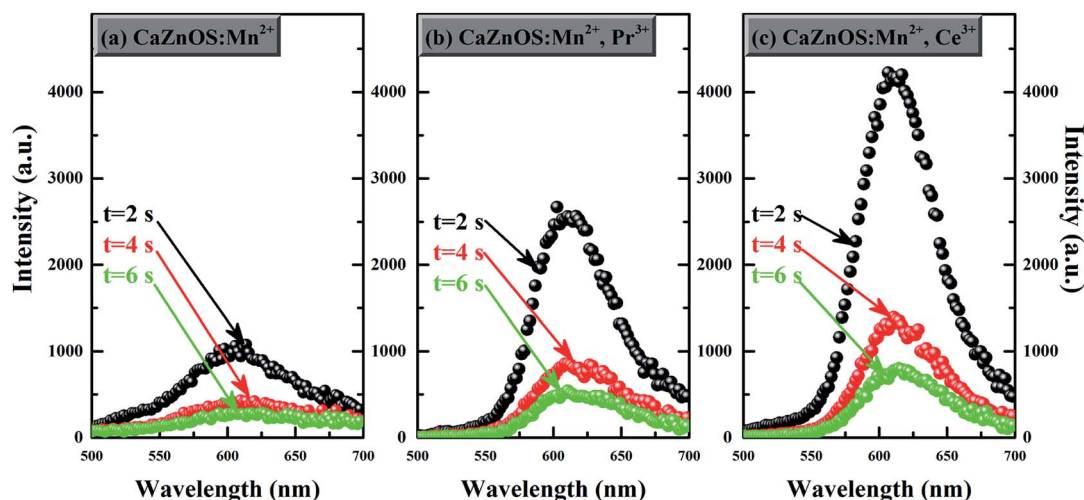


Fig. 8 Persistent luminescence emission spectra of (a) CaZnOS:Mn<sup>2+</sup>; (b) CaZnOS:Mn<sup>2+</sup>, Pr<sup>3+</sup>; (c) CaZnOS:Mn<sup>2+</sup>, Ce<sup>3+</sup> at different times after removal of the excitation source.



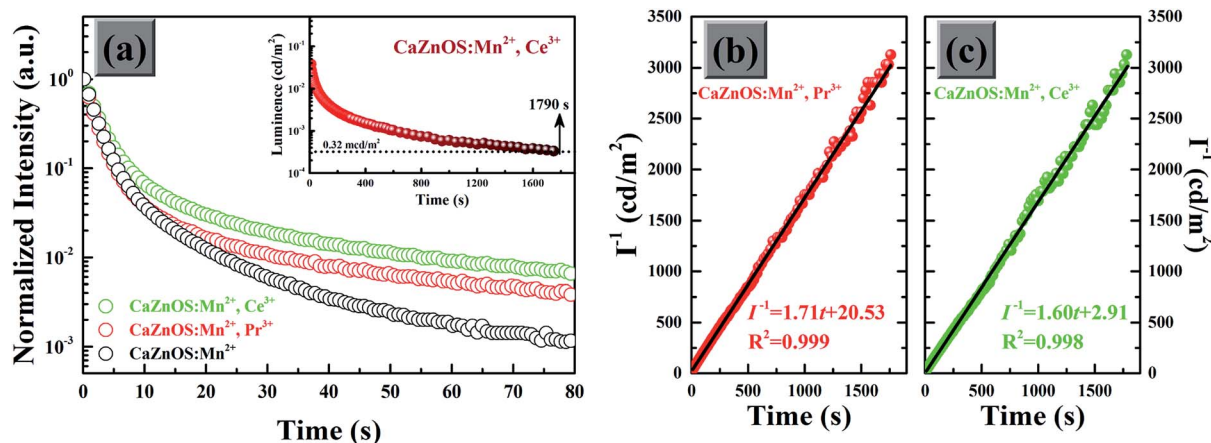


Fig. 9 (a) Afterglow decay curves of  $\text{Mn}^{2+}$ -doped and  $\text{Mn}^{2+}$ ,  $\text{RE}^{3+}$  co-doped  $\text{CaZnOS}$  samples with their initial intensity normalized for comparison. Inset: The afterglow decay curve of  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$  by definition of  $0.32 \text{ mcd m}^{-2}$ . Function of reciprocal afterglow intensity versus time of (b)  $\text{CaZnOS:Mn}^{2+}, \text{Pr}^{3+}$  and (c)  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$ .

by only one type effective trap center.<sup>35</sup> The electron traps created by  $\text{Pr}^{3+}$  and  $\text{Ce}^{3+}$  seemed to have similar nature. This result demonstrates that the incorporation of  $\text{RE}^{3+}$  ( $\text{Pr}^{3+}$ ,  $\text{Ce}^{3+}$ ) does not influence the type of traps, which are predominantly responsible for the afterglow.

As already known, the density of traps generally plays a key role in the afterglow emission of optical materials. Defects with suitable structure could capture the carrier more effectively and deliver them more slowly to the emission centers. The thermoluminescence technique was carried out because of its sensitivity to the trapping property of defects. Fig. 10 represents the TL spectra of  $\text{CaZnOS:Mn}^{2+}$  and  $\text{CaZnOS:Mn}^{2+}, \text{RE}^{3+}$  ( $\text{Pr}^{3+}$ ,  $\text{Ce}^{3+}$ ) samples. One very weak peak is observed above room temperature at around 350 K in  $\text{CaZnOS:Mn}^{2+}$  sample, which is formed by native defect in  $\text{CaZnOS}$  (Fig. 10a). As show in Fig. 10b and c, the TL intensity of  $\text{CaZnOS:Mn}^{2+}, \text{RE}^{3+}$  ( $\text{Pr}^{3+}$ ,  $\text{Ce}^{3+}$ ) samples are more higher than the  $\text{CaZnOS:Mn}^{2+}$ , which

indicating that more traps are created when  $\text{RE}^{3+}$  ( $\text{Pr}^{3+}$ ,  $\text{Ce}^{3+}$ ) co-doped in  $\text{CaZnOS:Mn}^{2+}$ . Because of the nonequivalent substitution, an excess of positive charge in the lattice must be compensated. There are two possible ways to fulfill the charge compensation of the  $\text{RE}^{3+}$  co-doped in  $\text{CaZnOS:Mn}^{2+}$  phosphor. One possible way is that two  $\text{RE}^{3+}$  ions replace three  $\text{Ca}^{2+}$  ions to balance the charge of the phosphor, which creates two  $\text{RE}'_{\text{Ca}}$  positive defects and one  $\text{V}''_{\text{Ca}}$  negative defect. The other possibility of the charge compensation, the vacancies of  $\text{V}''_{\text{Zn}}$  created during the synthesis process, is also feasible because of the relatively high vapor pressure of the  $\text{Zn}^{2+}$  component. By adsorbing and integrating the previous research results,<sup>36,37</sup> it is considered that the mainly defects in Fig. 10 belong to  $\text{V}''_{\text{Zn}}$  created. This negative defect is responsible for the improvement of long afterglow of  $\text{Mn}^{2+}$ . According to Chen's method,<sup>38</sup> the trap depth and density of TL peak can be estimated by the following equations:<sup>39,40</sup>

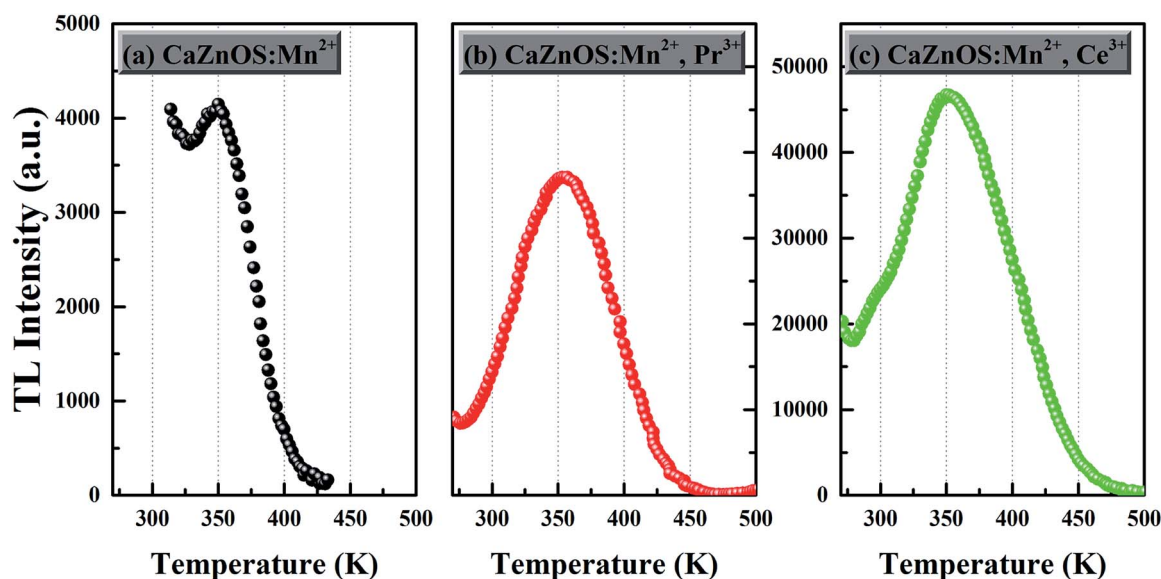


Fig. 10 Thermoluminescence curves of (a)  $\text{CaZnOS:Mn}^{2+}$ , (b)  $\text{CaZnOS:Mn}^{2+}, \text{Pr}^{3+}$  and (c)  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$ .



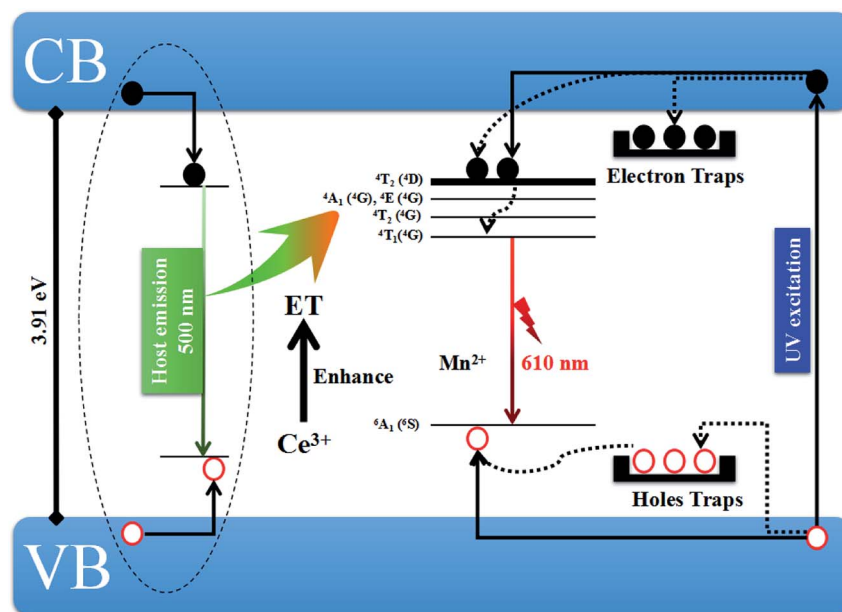


Fig. 11 Possible schematic diagram of host emission and the red afterglow emission in  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$ .

$$E = [2.52 + 10.2(\mu_g - 0.42)](k_B T_m^2 / \omega) - 2k_B T_m \quad (3)$$

$$n_0 = \omega I_m / [2.52\beta + 10.2\beta(\mu_g - 0.42)] \quad (4)$$

where  $E$  is the trap depth;  $\mu_g$  is the asymmetry parameter and defined as  $\mu_g = \delta/\omega$ ;  $\omega$  is the FWHM (full width of half maximum) and defined as  $\omega = \delta + \tau$ , with  $\delta$  being the high-temperature half width and  $\tau$  the low-temperature half-width;  $k_B$  is Boltzmann's constant;  $T_m$  is the temperature of the glow peak;  $n_0$  is the initial concentration of trapped charges;  $I_m$  is the TL intensity of glow peak and  $\beta$  is the heating rate ( $1 \text{ K s}^{-1}$ ). Therefore, the trap depths  $E$  corresponding to  $\text{CaZnOS:Mn}^{2+}, \text{Pr}^{3+}$  and  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$  phosphors are about 0.623 eV and 0.652 eV, respectively. The calculated  $n_0$  are about  $8.6 \times 10^5 \text{ (cm}^{-3})^{-1}$  and  $1.3 \times 10^6 \text{ (cm}^{-3})^{-1}$ , respectively. It is reported that a trap depth around 0.6–0.7 eV whose TL peaks located at 320–390 K is ideal for generating long afterglow at room temperature. On the basis of the TL peak position and the calculated trap depth, we believe that the  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$  phosphor have an opportune trap depth so that the sample could exhibit excellent afterglow at room temperature.

Fig. 11 displays the schematic diagram of the generation processes of host emission in  $\text{CaZnOS}$  and the after glow emission in  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$ . For undoped  $\text{CaZnOS}$ , under the 376 nm excitation, electrons can be pumped from valence band (VB) into conduction band (CB). The recombination of electrons and holes immediately generates the emission of host (500 nm). According to previous studies and the above results, we give the possible mechanism of afterglow in  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$  as following. After UV excitation, electrons are excited to CB and holes are generated in VB in  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$ . Some of the excited electrons will be transferred via the lattice directly to the luminescence centers  $\text{Mn}^{2+}$  and

finally to excited energy level of  $4T_1(4G)$  with non-radiative transitions. Afterwards, they will release to ground state and combines with holes accompanying with the 610 nm red emission. However, most of the excited electrons and the holes are trapped by lattice defects. Subsequently, with the thermal disturbance, these carriers will be released from the traps and transferred via the host to the excited state of  $\text{Mn}^{2+}$  ions. In addition, some of released electrons also will transfer gradually to the excited level of  $\text{Mn}^{2+}$  ions through tunneling. Owing to the slowly releasing of trapped carriers, the  $\text{Mn}^{2+}$  emission can last for a relatively long time after the removal of excitation.

## Conclusions

In conclusion, we extend the persistent luminescence into the red spectral region by developing a novel  $\text{Mn}^{2+}$ -doped  $\text{CaZnOS}$  persistent phosphor. The structure and luminescent properties of  $\text{CaZnOS}$  and  $\text{CaZnOS:Mn}^{2+}$  phosphors were investigated. Codoping with  $\text{RE}^{3+}$  ( $\text{Pr}^{3+}$ ,  $\text{Ce}^{3+}$ ) ions, the  $\text{CaZnOS:Mn}^{2+}, \text{RE}^{3+}$  ( $\text{Pr}^{3+}$ ,  $\text{Ce}^{3+}$ ) sample has more stronger afterglow intensity and longer duration of the afterglow. The afterglow intensity for the  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$  phosphor can be recorded for about 1790 s by definition of  $0.32 \text{ mcd m}^{-2}$  after the irradiation have been removed. From the TL curves, we can calculate that the trap depth of  $\text{CaZnOS:Mn}^{2+}, \text{Pr}^{3+}$  is 0.623 eV and  $\text{CaZnOS:Mn}^{2+}, \text{Ce}^{3+}$  is 0.652 eV. Moreover, a simplified schematic illustration was also constructed based on the photoluminescence and thermoluminescence results. The results suggest that  $\text{CaZnOS}$  may be an excellent host material for  $\text{Mn}^{2+}$ -based long afterglow. The further work can be focused on the improvement of the decay time and further enhancement of the afterglow brightness by adjusting the concentrations of  $\text{RE}^{3+}$  ( $\text{Pr}^{3+}$ ,  $\text{Ce}^{3+}$ ) and  $\text{Mn}^{2+}$ .



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