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1. Introduction

Due to their exceptional optical properties, rare-earth-doped inorganic phosphors have been extensively researched and developed for a wide range of applications, including white light-emitting diodes (WLEDs), display screens, indoor plant cultivation, and biomedical uses.¹ Commercial WLEDs are generally fabricated by using a blue LED chip (InGaN) coated by a yellow YAG:Ce³⁺ phosphor;^{1,2} however, they only exhibit a low quality of white light (a poor color rendering index, CRI, <75 and high correlated color temperature, CCT, >4500 K) due to lack of red emission.² In addition, compared to incandescence and sunlight, the emission spectra of WLEDs lack a cyan emission

Highly efficient and thermally stable cyan-emitting ZnS/ZnO phosphors for full-visible-spectrum LED lighting⁺

Do Quang Trung, ^(b) *^a Manh Trung Tran, ^(b) *^b Nguyen Van Du, ^a Nguyen Duy Hung,^c Tong Thi Hao Tam, ^d Nguyen Tu, ^(b) ^a Pham Minh Tri, ^b Hoang Gia Chuc, ^b Nguyen Duc Trung Kien, ^e Nguyen Van Quang^f and Pham Thanh Huy ^(b) ^b

Near-ultraviolet (NUV)-pumped white light-emitting-diodes (WLEDs) often suffer from poor color rendering in the 480–520 nm range, highlighting the need for an efficient cyan phosphor with strong absorption at 370–420 nm. This study presents the successful synthesis of cyan-emitting ZnS/ZnO phosphors using a high-energy planetary ball milling method followed by post-annealing. The fabricated phosphors, with particle sizes ranging from 1 to 3 μ m, exhibit strong cyan emission with CIE chromaticity coordinates of (0.2302, 0.3759) and excellent thermal stability with an activation energy of 0.26 eV. A prototype near-ultraviolet (NUV)-pumped cyan LED was developed, achieving chromatic coordinates of (0.2769, 0.4380) and a quantum efficiency of 77% by coating an NUV chip at 370 nm with the synthesized phosphor. These results demonstrate the potential of ZnS/ZnO-based materials as efficient, non-toxic alternatives to rare-earth phosphors, paving the way for advancements in full-spectrum white LEDs for solid-state lighting.

(470-500 nm) between the primary blue LED chip and the YAG:Ce³⁺ phosphor emission.^{3,4} Recently, many efforts have been made to enhance the CRI values and adjust the CCT of YAG:Ce³⁺-based WLEDs by adding red phosphors, and a high CRI of around 90 has been achieved.⁵ On the contrary, little attention has been paid to fulfilling the cyan gap by discovering cyan-emitting phosphors well excited by blue light for ultrahigh-CRI (>95) pc-WLEDs.6 Another way to make full-spectrum WLEDs with high-CRI could be based on a combination of a near-ultraviolet (NUV) LED chip with red, green, and blue (RGB) or cyan and red phosphor.7,8 Many studies have recently been made to find efficient phosphors for NUV-pumped WLEDs. Among them, the research on new phosphors such as high-efficiency cyan emitting phosphors excitable by NUV light is rapidly increasing and has become one of the hot research topics.^{2,9} High-efficiency cyan emitting phosphors generally were synthesized by using rare-earth ions activated by several host lattices, such as Ba₃Y₂B₆O₁₅:Ce³⁺, Tb³⁺,¹ RbBa₂(PO₃)₅: $Eu^{2+,7}$ $Ba_9Y_2Si_6O_{24}:Eu^{2+,10}$ $RbNa(Li_3SiO_4)_2:Eu^{2+,11}$ $Ba_9(Lu_{2-x-y})_2$ Al_x)Si₆O₂₄:yCe³⁺, Ca₃SiO₄(Cl,Br)₂:Eu²⁺, BaZrSi₃O₉:Eu²⁺, *etc.* However, these phosphors contain rare-earth and complex components, making them expensive and difficult to synthesize.

As widely known, wide bandgap semiconductors such as zinc oxide (ZnO) and zinc sulfide (ZnS), featuring substantial bandgaps of approximately 3.72–3.77 eV for ZnS and around 3.37 eV for ZnO, hold significant promise as efficient and non-toxic alternatives to rare-earth-based phosphors in applications such as

^aFaculty of Fundamental Sciences, Phenikaa University, Yen Nghia, Ha-Dong District, Hanoi 10000, Vietnam. E-mail: trung.doquang@phenikaa-uni.edu.vn

^bFaculty of Materials Science and Engineering, Phenikaa University, Yen Nghia, Ha-Dong District, Hanoi 10000, Vietnam. E-mail: trung.tranmanh@phenikaa-uni.edu.vn ^cFaculty of Electronic Materials and Components, School of Materials Science and Engineering, Hanoi University of Science and Technology (HUST), No. 01 Dai Co Viet, Hanoi 10000, Vietnam

^dFaculty of Information Technology, College of Technology, National Economics University (NEU), 207 Giai Phong Street, Hanoi 10000, Vietnam

^eFaculty of Electrical and Electronic Engineering, Phenikaa University, Yen Nghia, Ha-Dong District, Hanoi 10000, Vietnam

^fDepartment of Chemistry, Hanoi Pedagogical University 2, Phuc Yen, Vinh Phuc, Vietnam

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fluorescent and solid-state lighting.¹³⁻¹⁶ Both ZnO and ZnS, when incorporating intrinsic defects, exhibit strong emission across the entire visible spectrum, contingent upon the specific defect type formed. Undoped ZnO usually emits green light associated with oxygen vacancies (V_O) or blue light related to the zinc interstitials (Zn_i) defect.^{17,18} In parallel, ZnS exhibits robust emission in the blue spectrum, primarily through the sulfur vacancies defect (V_S).¹⁹⁻²¹ Recent advancements in ZnS and ZnO-based materials include the successful development of warm-white light-emitting diodes (WLEDs),^{22,23} highlighting the potential of cyan-emitting ZnS/ZnO phosphors. Although, ZnS/ZnO nanocomposite phosphor was fabricated,²⁴ microsized ZnS/ZnO phosphors for WLED applications have not been studied yet.

High-energy planetary ball milling has recently emerged as an effective method for synthesizing phosphors at lower calcination temperatures than conventional solid-state reactions, while also introducing defect-related states that create electron/ hole traps within the band gap.²⁵ These new defect states can result in additional emission peaks in the photoluminescence spectra. Consequently, high-energy planetary ball milling shows great promise for synthesizing cyan-emitting ZnS/ZnO-based phosphors by creating these new defect states.

Herein, cyan-emitting ZnS/ZnO phosphors were synthesized using the high-energy planetary ball milling method coupled with post-annealing. The crystal structure and optical performances of the obtained phosphors were characterized in detail. Furthermore, a cyan LED coated with ZnS/ZnO phosphor prototype was successfully fabricated to demonstrate the potential for full-spectrum white LED applications.

2. Experimental

Zinc sulfide (ZnS, 99.99%) and zinc oxide (ZnO, 99.99%) powders from Sigma-Aldrich were used as received. ZnS, ZnO, and their 1:1 weight mixture were ball-milled at 500 rpm for 2 hours using a Fritsch Pulverisette7 (Germany), followed by annealing in an argon atmosphere at 1000 °C for 2 hours to evaluate their characteristics.

The crystalline structure was characterized by X-ray diffraction (XRD, Rigaku Smartlab, Japan D2 Phaser A26-X1, Bruker, Germany) with a Cu K α ($\lambda = 0.154$ nm) performing at 40 mA tube current. The surface morphologies and compositions were observed by a field emission scanning electron microscopy system (FESEM-JEOL/JSM-7600F) equipped with energydispersive X-ray spectroscopy (EDX). The Raman spectra were obtained by a Raman microscopy - XploRA Plus instrument (Horiba, France) using the wavelength of 532 nm of the laser source. The XPS spectra were investigated by an X-ray photoelectron Spectroscope (XPS, VG Scientific, ESCALAB250) radiated by a monochromatic Al Ka source. The UV-vis absorption spectra were obtained by a single-monochromator Jasco V-750 and spectroscopy. Photoluminescence (PL) photoluminescence excitation (PLE) spectra were studied by a spectrofluorometer excited by a 450 W xenon discharge lamp



Fig. 1 (a) XRD pattern refinement of ZnS/ZnO phosphor following grinding and post-annealing at 1000 °C for 2 h in Ar. (b) Visualization of the ZnO and ZnS crystal structure within the resulting phosphor, created using VESTA software.

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 Table 1
 Comparison of the average crystallite sizes of ZnO, ZnS, and ZnO/ZnS samples determined using the Scherrer equation in this study and previously published works

Sample	Fabrication method	Crystallite size of ZnS (nm)	Crystallite size of ZnO (nm)	Ref.
ZnS/ZnO nanocomposite phosphor	Combustion synthesis	11	10	24
ZnS/ZnO nanocomposites	Solid-state method	46-50	45-54	28
ZnO–ZnS core–shell nanostructure	Solution method	6.95	22.65	29
ZnS-ZnO nanocomposites	Hydrothermal routes	1.58-1.97	3.91-4.40	30
ZnS/ZnO phosphor	High-energy planetary ball milling method and annealing	16.71	17.37	This work

 Table 2
 The obtained parameters from Rietveld's refinement of the XRD data for the ZnSO sample

					Fractiona co-ordina	al position ates				
Phases	Lattice parameters (Å)	Lattice volume (Å ³)	Phase fraction (%)	Atoms	x	у	z	Wyckoff positions	Occupancy	U
ZnS phase	a = 3.82394	$\textbf{79.234} \pm \textbf{0.019}$	73.84 ± 0.52	Zn	0.33333	0.66667	0.00000	2b	1.000	0.104
				S	0.33333	0.66667	0.37480	2b	0.907	0.059
	c = 6.25692			0	0.33333	0.66667	0.37480	2b	0.067	0.059
ZnO phase		47.642 ± 0.008	26.16 ± 0.22	Zn	0.33333	0.66667	0.00000	2b	1.000	0.076
	a = 3.25075			0	0.33333	0.66667	0.38200	2b	0.944	-0.020
	c = 5.20587			S	0.33333	0.66667	0.38200	2b	0.015	0.000
"R" values		$R_{\rm p} = 9.42\%$		$R_{\rm exp} = 8.73\%$				$\chi^2 = 2.45$		

(Nanolog, HORIBA Jobin Yvon). The optical characteristics of LED devices were measured by a spectroradiometer (RadOMA, Gamma Scientific GS-1290).

Finally, a 4:1 mixture of polydimethylsiloxane (PDMS, Dow Corning OE-6370) and ZnS/ZnO phosphor was prepared to fabricate WLED prototypes. The mixture was coated onto



Fig. 2 (a) FESEM image, (b) EDS spectrum, and (c) Raman spectrum of ZnS/ZnO phosphor.

a 370 nm UV chip, and then cured in an oven (ASONE AVO-310SB-D) at 150 °C for 4 hours.

3. Results and discussion

Fig. 1a presents XRD pattern refinement of ZnS/ZnO phosphors received after grinding by a high-energy planetary ball milling and post-annealing at 1000 $^{\circ}$ C in Ar for 2 h.

Both ZnS and ZnO phases are evident in the XRD pattern (with ZnS, ZnO, and ZnS/ZnO annealed under the same conditions, as shown in Fig. S1[†]). The distinct diffraction peaks at $2\theta = 26.9^{\circ}$, 28.6°, and 30.6° correspond to the (100), (002), and (101) planes of the hexagonal ZnS phase (wurtzite-2H, space P63mc(186), JCPDS Card No. 01-1450).²⁶ Additionally, the peaks at $2\theta = 31.8^{\circ}$, 34.5° , and 36.3° are attributed to the (100), (002), and (101) planes of hexagonal zincite phase ZnO (JCPDS Card No. 36-1451).26,27 The crystallite sizes of the synthesized powders were calculated using the Scherrer equation, $d_{\text{Scherrer}} = 0.9\lambda/(\beta \times \cos \theta)$,²⁴ where λ is the X-ray wavelength ($\lambda = 1.54$ Å), θ is the diffraction angle, and β is the full width at half-maximum (FWHM, in radians) of a diffraction peak. Table 1 compares the average crystallite sizes of ZnS and ZnO phases synthesized by various methods, highlighting the significant impact of fabrication techniques on structural properties. Hydrothermal synthesis produces the smallest crystallite sizes (1.58-1.97 nm for ZnS and 3.91-4.40 nm for ZnO), making it ideal for ultrafine material application.³⁰ In contrast, combustion synthesis yields moderate sizes (10-11 nm),²⁴ while the solid-state method results in much larger crystallites, up to 54 nm,28 indicating limited refinement. The solution method offers intermediate crystallite sizes (6.95 nm for ZnS and 22.65 nm for ZnO), striking a balance between refinement and structural control.29

Thanks to the advantage of the high-energy planetary ball milling, this study achieves relatively small sizes (16.71 nm for ZnS and 17.37 for ZnO), combining efficient synthesis with potential defect-related enhancements. These results highlight the importance of selecting appropriate synthesis methods to optimize material performance.

FullProf Suite analysis reveals that the ZnS and ZnO phases comprise 73.84% and 26.16%, respectively. Their Rietveld refinement parameters, including the reliability factors (χ^2 , R_p , and R_{exp}), lattice constants and unit cell volume are listed in Table 2. It is worthwhile noting that the calculated patterns are consistent with the measured data, with low R_{p} and R_{exp} values confirming the reliability of the structural models used for the refinements. Based on the lattice parameters obtained from the Rietveld refinement, a schematic of the ZnS and ZnO unit cells with wurtzite structures was generated using VESTA software, as shown in Fig. 1b. Table 2 clearly presents detailed information on lattice parameters, phase fractions, atomic positions, and occupancy values for the ZnS and ZnO phases, allowing for easy comparison. The phase fractions are accurate, summing to 100%. Lattice parameters are precise, though indicating the symmetry type (e.g., cubic or hexagonal) would provide additional context. The fractional positions and Wyckoff positions are well-reported, but the rationale behind the sulfur occupancy in ZnS (0.907) could be clarified. Occupancy and atomic displacement (U) values are useful, though further explanation

31 32 33 34 35 35 36 37 38 39 40 71his work Ref. $E_2(high)/LO + TA$ 138.5 (ZnO) 115.2 (ZnS) 434 437 450 433 137 29 (DL) 382.0 382.0 \mathbf{A}_1 $A_1/E_1(LO)$ _____351 350 350 347.7 350 339 346.8351.2 330 333.2 S (OL E 2 (DTO) A_1/E_1 251.1 69.2 Raman peak position (cm^{-1}) 262 269 269 267 267 267 275 275 271 Comparison on Raman modes of ZnS/ZnO phosphor and previous reports ΓA $E_2(low)$ 100.6 High-energy planetary ball milling and post-annealing Chemical colloidal process Microwave irradiation Hydrothermal method Hydrothermal method Magnetron sputtering Hydrothermal method Thermal evaporation Thermal evaporation Solution combustion Synthesis method Sol-gel method Sol-gel method ZnS micro-nanostructures ZnS/ZnO nanostructure InO:N nanostructures ZnS-ZnO composites znO:Al nanoparticles znS nanostructures znO nanoparticles ZnS:Al³⁺ nanorods znS nanoribbons ZnO:Al thin film znS nanowires Materials ZnS/ZnO Table 3

of negative or low *U* values, especially for sulfur in ZnO, would be beneficial.

The refinement quality ($R_p = 9.42\%$, $R_{exp} = 8.73\%$, $\chi^2 = 2.45$) suggests a reasonable fit, but a brief explanation of these values would improve clarity. The low sulfur occupancy in ZnO (0.015) may indicate impurities, which should be addressed. Additionally, the table could benefit from details on the XRD data range and refinement software used for a more thorough assessment.

Fig. 2a presents the FESEM image of ZnS/ZnO phosphor, with particle sizes ranging from 0.5 to 3 μ m and clear grain boundaries. Fig. 2b shows the EDS spectrum of ZnS/ZnO

phosphor powder, revealing atomic proportions of Zn (47.7%), O (27.6%), and S (24.7%). The lower S content than O may be due to the partial oxidation of ZnS into ZnO during annealing, which is consistent with the XRD phase composition results. The morphology and elemental composition of the annealed ZnS and ZnO are detailed in Fig. S2 and Table S1.[†]

The Raman spectrum of the ZnS/ZnO phosphor, excited by a 532 nm laser, is shown in Fig. 2c, with the resonance modes for both ZnS and ZnO summarized in Table 3. This table compares the Raman peak positions of ZnS/ZnO phosphors synthesized through various methods, highlighting differences



Fig. 3 XPS spectra of ZnS/ZnO phosphor powder: (a) complete survey, and their high-resolution binding energy spectra of (b) Zn 2p (c) S 2p, (d), and O 1s.



Fig. 4 (a) UV-vis spectroscopy analysis of ZnS/ZnO sample and (b) Tauc plot.



Fig. 5 (a) PLE and (b) PL spectrum of ZnS/ZnO phosphor excited by the 325 nm wavelength of Xe lamp; (c) CIE chromaticity diagram of the ZnS/ZnO phosphors ($\lambda_{ex} = 325$ nm) and (d) proposed mechanism for emitting cyan emission from ZnS/ZnO phosphor.

across multiple samples. The distinctive peaks at 216.6, 269.2, 281.5, 333.2, 351.2, and 415.2 cm⁻¹ correspond to the longitudinal acoustic (LA), transverse optical [A₁/E₁(TO)], translational oscillation [E₂(TO)], surface optical (SO), longitudinal optical $[A_1/E_1(LO)]$, and translational oscillation $E_2(high)$ resonance modes of ZnS, respectively.^{20,31} Additionally, the peaks at 100.6, 382.0, and 438.5 cm⁻¹, corresponds to the E₂(low), A₁(TO), and $E_2(high)$ resonance modes of ZnO.^{40,42} Notably, the sharp peaks at 351.2 cm⁻¹ (A₁/E₁(LO) mode of ZnS) and 438.5 cm⁻¹ (E₂(high) mode of ZnO) confirm the presence of wurtzite ZnS and ZnO phases.32 These results demonstrate the effectiveness of this synthesis method in capturing a broad range of Raman modes, providing valuable insights into the material's vibrational properties. The absence of certain peaks in some references (indicated by dashes) may suggest variations in synthesis conditions or material quality.

3.1. Binding energy analysis

Fig. 3a shows the XPS survey spectrum of ZnS/ZnO phosphor powder, with clear peaks corresponding to Zn, O, S, and C. The XPS spectra are normalized according to the C 1s binding energy peak at 284.65 eV.^{43,44} In Fig. 3b, the high-resolution binding energy spectrum of Zn 2p exhibits a well-defined Zn $2p_{3/2}$ peak at approximately 1021.4 eV, aligning with the binding energy of Zn²⁺ in ZnO and ZnS.^{45,46} The deconvolution of the S 2p XPS spectra (Fig. 3c) shows two peaks at 161.5 eV and 162.6 eV, corresponding to the S $2p_{3/2}$ and S $2p_{1/2}$ electronic states, with the S $2p_{3/2}$ binding energy indicating Zn–S bonding.^{43,47–49}

The analysis of the O 1s XPS spectra, using multiple Gaussian fitting, reveals three peaks at 530.3, 531.4, and 532.1 eV (Fig. 3d). The peak at 530.3 eV can be attributed to the lattice oxygen in ZnO,^{39,50} while the peak at 531.4 eV is assigned to the O^{2-} ions in oxygen-deficient regions within ZnO matrix, indicating the presence of oxygen vacancy (V_O).^{39,51,52} The peak at 532.1 eV may be associated with H₂O or O₂ absorbed on the surface.^{41,50} These findings suggest a significant presence of defects related to oxygen vacancies within the host lattice of ZnS/ZnO phosphor.

3.2. UV-vis spectroscopy analysis and photoluminescence study

DRS UV-visible analysis was used to examine the optical properties of the synthesized ZnS–ZnO sample (Fig. 4), focusing on reflectance calculated using the Kubelka–Munk equation. Two



Fig. 6 (a) Temperature-dependent emission spectra excited by the wavelength of 235 nm and (b) the dependence of normalized PL emission intensity of spectra of ZnS/ZnO phosphor and temperature. The insets illustrate the relationship between $ln[(l_0/l) - 1]$ and 1/kT.

distinct optical bands were observed, corresponding to ZnS (3.46 eV) and ZnO (3.17 eV). The bandgap of ZnS/ZnO was compared to that of ZnS and ZnO annealed under the same conditions (see Fig. S3 and Table S2†). The reduced bandgap in

ZnS/ZnO may be attributed to surface defects or diffusion of S into ZnO and O into ZnS. 29,53

Fig. 5a presents the PLE spectrum of ZnS/ZnO phosphor monitored at 495 nm. The spectrum shows a characteristic absorption band of ZnS peaking at 341 nm (\sim 3.63 eV) and a weaker ZnO absorption band centered at 378 nm (\sim 3.28 eV). This effect may be due to the high-energy planetary ball milling and annealing at elevated temperatures, which likely induce intrinsic defects in the ZnS and ZnO host lattices²⁵ or result in the formation of ZnS/ZnO heterostructures.²⁸ These findings are consistent with Raman and XPS analyses.

Fig. 5b illustrates the PL spectrum of ZnS/ZnO phosphor, exhibiting a broad visible emission (400–700 nm) with a peak at 495 nm under 325 nm Xe lamp excitation. Gaussian deconvolution reveals two prominent peaks at 473 nm (blue) and 527 nm (green), with green emission dominating the PL spectrum. The blue peak at 473 nm is attributed to intrinsic defects, such as sulfur vacancies (V_S) in ZnS^{54,55} or zinc interstitials (Zn_i) in ZnO.^{43,56,57}

The green emission at 527 nm is likely due to oxygen vacancies $(V_{\rm O})$ in $ZnO^{55,58}$ or the ZnO/ZnS interface.^59 Using



Fig. 7 (a) EL spectrum of NUV chip (370 nm, 3 W, 3.4 V); (b) EL spectra of NUV-pumped ZnS/ZnO phosphor; (c) the CIE 1931 chromaticity diagram of the fabricated cyan-LED and NUV-chip.

OSRAM Sylvania's ColorCalculator software, the CIE 1931 color coordinates of the ZnS/ZnO phosphor were calculated to be (0.2302, 0.3759), placing it in the cyan emission region (Fig. 5c). Fig. 5d depicts the energy level diagram, illustrating the combination of blue and green emissions to produce cyan light.

The thermal stability of phosphors is critical for LED applications, as operational temperatures can exceed 150 °C.²² Fig. 6a presents the PL spectra of ZnS/ZnO phosphor at various temperatures, excited at 325 nm.

While the spectra maintain their shape in the visible region, the intensity decreases as the temperature rises from 303 to 443 K. At 423 K, approximately 51% of the initial intensity is retained. For comparison, the Ca₃Y(PO₄)₃:Eu²⁺, Mn²⁺ phosphor retains 52% under similar conditions,⁶⁰ while the Ba₃Y₂B₆O₁₅:-Ce³⁺, Tb³⁺ phosphor retains only 21.9%.¹ The thermal quenching effect is linked to the activation energy (ΔE), calculated using the modified Arrhenius eqn (1):²²

$$I_{\rm T} = I_0 [1 + A \exp(-\Delta E/kT)]^{-1}$$
(1)

where I_0 and I_T are the initial PL intensity and the PL intensity at a specific temperature, respectively. A is a constant, and $k = 8.62 \times 10^{-5}$ eV K⁻¹ is Bolzmann's constant. By fitting the plot of ln $[I_0/I_T - 1]$ versus 1/kT, ΔE was determined as 0.26 eV (Fig. 6b). This value could be compared with some phosphors using for WLED such as Ca₃Y(PO₄)₃:Eu²⁺, Mn²⁺ (0.166 eV),⁶⁰ Ca₄ZrGe₃-O₁₂:Bi³⁺ (0.228 eV),² Ba₃Bi(PO₄)₃: Sm³⁺, Eu³⁺ (0.21 eV),⁶¹ and NaGd₉(SiO₄)₆O₂:Bi³⁺ (0.296 eV).⁶²

The electroluminescence (EL) spectrum of the NUV 370 nm chip driven at 500 mA in Fig. S5,[†] highlighting a strong NUV emission peak at 370 nm. The phosphor-converted cyan LEDs exhibit a broad visible emission region and a weaker NUV peak. The visible emissions are produced by the phosphor layer excited by the 370 nm NUV chip, while the residual NUV emission in the EL spectrum originates from unabsorbed light from the NUV chip. Quantum efficiency (QE) is a critical parameter for assessing the suitability of phosphors for LED applications.⁶³ The QE of the ZnS/ZnO phosphor, when coated onto a NUV chip, can be calculated using the following eqn (2):²²

$$QE = \frac{I_{em}}{I_{abs}} = \frac{\int I_{em, vis-LED} - \int I_{em, vis-chip}}{\int I_{em, UV-chip} - \int I_{em, UV-LED}} \times 100$$
(2)

where, $\int I_{\text{em,vis-LED}}$ and $\int I_{\text{em,vis-chip}}$ correspond to the integrated visible emission of the LED with and without the phosphor layer, respectively.

Using this equation, the QE of the ZnS/ZnO phosphor was estimated to be around 77%. This QE value surpasses those of several other reported materials, including ZnO nanowires (20%),⁶⁴ Ba₃Lu₂B₆O₁₅:Ce³⁺,Tb³⁺ phosphor (51%),⁶⁵ Mn-doped ZnS/ZnO nanobelts (61%),⁶⁶ RbNa_{1-y}(Li₃SiO₄)₂:yEu²⁺ phosphor (62.9%),¹¹ and it is comparable to Ca₂GdHf₂(AlO₄)₃:Ce³⁺ (80%).⁶⁷

The digital image and electroluminescence (EL) parameters of the NUV chip are presented in the inset of Fig. 7a. The EL spectrum of the NUV chip, driven by a 500 mA current, reveals strong NUV emission with chromaticity coordinates (x, y) of (0.2970, 0.2719), spanning the 350-410 nm range and peaking at approximately 377 nm. Fig. 7b shows the EL spectrum of the

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Phosphor powder	Synthesis method and experimental conditions	Strongest emission peak (nm)	FWHM (nm)	Highest absorption peaks (nm)	Color-coordinates	QE (%)	Lifetime (µs)	Activation energy, $E_{\rm a}$ (eV)	Ref.
ZnS-ZnO	Hydrothermal, 120 °C, 24 b. amhiont	~ 470	Ι	316-319	I	I	I	Ι	30
ZnS:KBr,Mn ²⁺	24 II, annorm Solid-state, 750–1100 °C, 2 h. CO ambiant	468-476	I	344	I		$1.35{ extrm{-}2.47 imes10^3}$	I	68
ZnO:Er ³⁺ /Ho ³⁺	solid-state, 500 °C, 2 h	377	I	502			8.47-27.5	I	69
ZnO:Eu ³⁺ ,Dy ³⁺	Precipitation at 80 °C,	482		390	(0.514, 0.395)				70
	calcination for 2 h at 700 °C								
CaZnOS:Bi ³⁺ ,Mn ²⁺	Solid-state, 1100 °C, 3 h	485 - 491		375	(0.350, 0.323)		I	Ι	71
ZnO	Hydrothermal, 200 °C, 4 h	520	Ι	355	(0.302, 0.488)	Ι	Ι	Ι	72
ZnO:Ce ³⁺	Solid-state, 1000 °C, 6 h	499	Ι	375	(0.23, 0.33)	Ι	Ι	Ι	73
ZnS	Solid-state, 1050 °C, N ₂ /H ₂	460	I	342		I	Ι	Ι	74
ZnO:Al ³⁺	ambient								22
ZnS/ZnO	High-energy planetary ball milling, 1000 °C, 2 h	495		341, 378	(0.277, 0.438)	77		0.26	This work

ZnS/ZnO phosphor-coated NUV chip, accompanied by a digital image of the phosphor-coated NUV chip under 500 mA current and no current.

The emitted light is biased toward cyan, demonstrating the ZnS/ZnO phosphor's ability to convert NUV to visible cyan light. The measured chromaticity coordinates (x, y) and color rendering index (CRI) of the cyan-LED are (0.2769, 0.4380) and 58, respectively. Fig. 7c displays the CIE chromaticity coordinates for both the NUV chip and the ZnS/ZnO phosphor-coated NUV chip. The color coordinate discrepancy between the phosphor-coated NUV-LED and the phosphor powder is likely due to visible emission contributions from the NUV chip. Table 4 provides a comprehensive comparison of several cyanemitting phosphors.

This table details their synthesis methods, excitation sources, emission peaks, color coordinates, quantum efficiency (QE), and activation energy. The data highlights a variety of synthesis techniques and their impact on the material's optical properties. Notably, the ZnS/ZnO phosphor in this work exhibits a strong emission peak at 495 nm, a quantum efficiency of 77%, and an activation energy of 0.26 eV, which demonstrates excellent performance compared to other reported materials. The inclusion of these metrics strengthens the understanding of the relationship between synthesis conditions and phosphor properties.

4. Conclusions

In summary, the cyan-emitting phosphor derived from ZnS/ZnO has been successfully synthesized using a high-energy planetary ball milling method combined with post-annealing. The ZnS/ZnO phosphor, with particle sizes ranging from 1 to 3 μ m, exhibits cyan emission with CIE chromaticity coordinates of (*x*, *y*) = (0.2302, 0.3759). A prototype near-ultraviolet (NUV)-pumped cyan LED was also developed, showing chromatic coordinates of (0.2769, 0.4380) and a quantum efficiency of 77%, achieved by coating the NUV chip at 370 nm with ZnS/ZnO phosphor. The phosphor exhibits strong thermal stability with an activation energy of 0.26 eV. These findings open new avenues for cyan LED development using ZnS/ZnO phosphor, offering substantial promise for phosphor-converted white LEDs (pc-WLEDs) with full-color capability.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

The authors declare no conflict of interest.

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References

- B. Li, J. Liang, L. Sun, S. Wang, Q. Sun, B. Devakumar, G. Annadurai, D. Chen, X. Huang and Y. Wu, Cyanemitting Ba₃Y₂B₆O₁₅:Ce³⁺,Tb³⁺ phosphor: A potential color converter for near-UV-excited white LEDs, *J. Lumin.*, 2019, 211, 388–393, DOI: 10.1016/j.jlumin.2019.04.001.
- 2 Z. Jiang, J. Gou, Y. Min, C. Huang, W. Lv, X. Yu, X. Su and L. Duan, Crystal structure and luminescence properties of a novel non-rare-earth activated blue-emitting garnet phosphor $Ca_4ZrGe_3O_{12}:Bi^{3+}$ for n-UV pumped lightemitting diodes, *J. Alloys Compd.*, 2017, 727, 63–68, DOI: 10.1016/j.jallcom.2017.08.109.
- 3 X. Huang, Cyan phosphors for full-visible-spectrum lighting: shining new light on high-CRI white pc-LEDs, *Sci. Bull.*, 2019, 64, 1649–1651, DOI: 10.1016/j.scib.2019.09.008.
- 4 L. Wang, C. Yuan, P. Hu, R. Li, Y. Liu, P. Sun, R. Dong, X. Qi,
 J. Jiang and H. Jiang, A low cost and high efficient Ba₉(Lu_{2-x-y}Al_x)Si₆O₂₄:yCe³⁺ cyan-emitting phosphor, *Ceram. Int.*, 2020,
 46, 11466–11473, DOI: 10.1016/j.ceramint.2020.01.172.
- 5 H. Zhu, C. C. Lin, W. Luo, S. Shu, Z. Liu, Y. Liu, J. Kong, E. Ma, Y. Cao, R. Liu and X. Chen, Highly efficient nonrare-earth red emitting phosphor for warm white lightemitting diodes, *Nat. Commun.*, 2014, 5, 4312, DOI: 10.1038/ncomms5312.
- 6 M. Zhao, H. Liao, M. S. Molokeev, Y. Zhou, Q. Zhang, Q. Liu and Z. Xia, Emerging ultra-narrow-band cyan-emitting phosphor for white LEDs with enhanced color rendition, *Light Sci. Appl.*, 2019, **8**, 38, DOI: **10.1038/s41377-019-0148-8**.
- 7 Q. Zhang, X. Wang and Y. Wang, Design of a broadband cyan-emitting phosphor with robust thermal stability for high-power WLED application, *J. Alloys Compd.*, 2021, **886**, 161217, DOI: **10.1016/j.jallcom.2021.161217**.
- 8 J. Y. Park, K. S. Shim, J. S. Yu and H. K. Yang, Cyan-emitting BaZrSi₃O₉:Eu²⁺ phosphors for near-UV based white lightemitting diodes, *Mater. Lett.*, 2016, **173**, 68–71, DOI: **10.1016/j.matlet.2016.03.011**.
- 9 C. K. Shilpa, S. V. Jasira, V. P. Veena, B. Abhishek and K. M. Nissamudeen, Photometric study on excitation tailored cyan to white color conversion in SrCeO₃:Dy³⁺ for wLEDs, latent fingerprint, and security ink applications, *Mater. Today Commun.*, 2024, **38**, 107954, DOI: **10.1016**/ **j.mtcomm.2023.107954**.
- 10 C. Hu, B. Peng, K. X. Song, B. Liu, D. Wang and I. M. Reaney, The cyan-green luminescent behaviour of nitrided Ba₉Y₂Si₆O₂₄:Eu²⁺ phosphors for W-LED, *Ceram. Int.*, 2018, 44, S2–S6, DOI: 10.1016/j.ceramint.2018.10.166.
- 11 M. Liao, Z. Mu, Q. Wang, X. Zhang, H. Dong, M. Wen and F. Wu, Understanding the cyan-emitting phosphor RbNa(Li₃SiO₄)₂:Eu²⁺ by providing Rb ion vacancies, *J. Alloys Compd.*, 2020, 837, 155084, DOI: 10.1016/ j.jallcom.2020.155084.
- 12 Y. Zhu, L. Fu, D. Wu, J. Peng, F. Du, X. Ye, L. Chen and W. Zhuang, A cyan-emitting phosphor Ca₃SiO₄(Cl,Br)₂:Eu²⁺ with high efficiency and good thermal stability for full-

visible-spectrum LED lighting, *J. Lumin.*, 2021, **232**, 117854, DOI: **10.1016/j.jlumin.2020.117854**.

- 13 J. G. Simmons, J. V. Foreman, J. Liu and H. O. Everitt, The dependence of ZnO photoluminescence efficiency on excitation conditions and defect densities, *Appl. Phys. Lett.*, 2013, 103, 2–5, DOI: 10.1063/1.4829745.
- 14 V. Lahariya and S. J. Dhoble, Development and advancement of undoped and doped zinc sulfide for phosphor application, *Displays*, 2022, **74**, 102186, DOI: **10.1016**/ **j.displa.2022.102186**.
- 15 X. Fang, Y. Bando, U. K. Gautam, T. Zhai, H. Zeng, X. Xu, M. Liao and D. Golberg, ZnO and ZnS nanostructures: Ultraviolet-light emitters, lasers, and sensors, *Crit. Rev. Solid State Mater. Sci.*, 2009, 34(3–4), 190–223, DOI: 10.1080/10408430903245393.
- 16 A. Tiwari and S. J. Dhoble, Critical analysis of phase evolution, morphological control, growth mechanism and photophysical applications of zns nanostructures (zerodimensional to three-dimensional): A review, *Cryst. Growth Des.*, 2017, 17, 381–407, DOI: 10.1021/acs.cgd.6b01463.
- 17 J. Wang, R. Chen, L. Xiang and S. Komarneni, Synthesis, properties and applications of ZnO nanomaterials with oxygen vacancies: A review, *Ceram. Int.*, 2018, 44, 7357–7377, DOI: 10.1016/j.ceramint.2018.02.013.
- 18 S. Pramanik, S. Mondal, A. C. Mandal, S. Mukherjee, S. Das, T. Ghosh, R. Nath, M. Ghosh and P. K. Kuiri, Role of oxygen vacancies on the green photoluminescence of microwaveassisted grown ZnO nanorods, *J. Alloys Compd.*, 2020, 849, 156684, DOI: 10.1016/j.jallcom.2020.156684.
- 19 M. A. Shakil, S. Das, M. A. Rahman, U. S. Akther, M. K. Hassan and M. K. Rahman, A Review on Zinc Sulphide Thin Film Fabrication for Various Applications Based on Doping Elements, *Mater. Sci. Appl.*, 2018, 09, 751–778, DOI: 10.4236/msa.2018.99055.
- 20 X. Wang, J. Shi, Z. Feng, M. Li and C. Li, Visible emission characteristics from different defects of ZnS nanocrystals, *Phys. Chem. Chem. Phys.*, 2011, 13, 4715–4723, DOI: 10.1039/c0cp01620a.
- 21 M. Joy, A. P. Mohamed, K. G. K. Warrier and U. S. Hareesh, Visible-light-driven photocatalytic properties of binary MoS₂/ZnS heterostructured nanojunctions synthesized via one-step hydrothermal route, *New J. Chem.*, 2017, **41**, 3432– 3442, DOI: **10.1039/c6nj03727e**.
- 22 D. Q. Trung, N. V. Quang, M. T. Tran, N. V. Du, N. Tu, N. D. Hung, D. X. Viet, D. D. Anh and P. T. Huy, Singlecomposition Al³⁺-singly doped ZnO phosphors for UVpumped warm white light-emitting diode applications, *Dalton Trans.*, 2021, **50**, 9037–9050, DOI: **10.1039**/ **D1DT00971K**.
- 23 N. Karar, H. Chander and S. M. Shivaprasad, Enhancement of luminescent properties of ZnS:Mn nanophosphors by controlled ZnO capping, *Appl. Phys. Lett.*, 2004, **85**, 5058– 5060, DOI: **10.1063/1.1815059**.
- 24 M. Zahiri, M. Shafiee Afarani and A. M. Arabi, Combustion Synthesis of ZnO/ZnS Nanocomposite Phosphors, *J. Fluoresc.*, 2019, **29**, 1227–1239, DOI: **10.1007/s10895-019-02434-9**.

- 25 N. Tu, H. Van Bui, D. Q. Trung, A. T. Duong, D. M. Thuy, D. H. Nguyen, K. T. Nguyen and P. T. Huy, Surface oxygen vacancies of ZnO: A facile fabrication method and their contribution to the photoluminescence, *J. Alloys Compd.*, 2019, **791**, 722–729, DOI: **10.1016/j.jallcom.2019.03.395**.
- 26 X. F. Liu, X. R. Cui, X. B. Chen, N. Yang and R. H. Yu, Shellenhanced photoluminescence and ferromagnetism of Co:ZnS/Co:ZnO core-shell nanostructure, *Mater. Res. Bull.*, 2014, 50, 113–117, DOI: 10.1016/j.materresbull.2013.09.048.
- 27 S. Kar, P. Dutta, T. Pal and S. Ghosh, Simple solvothermal route to synthesize S-doped ZnO nanonails and ZnS/ZnO core/shell nanorods, *Chem. Phys. Lett.*, 2009, **473**, 102–107, DOI: **10.1016/j.cplett.2009.03.027**.
- 28 A. Oskenbay, D. Salikhov, O. Rofman, I. Rakhimbek,
 Z. Shalabayev, N. Khan, B. Soltabayev, A. Mentbayeva,
 M. Baláž and B. Tatykayev, Solid-state synthesis of ZnS/
 ZnO nanocomposites and their decoration with NiS
 cocatalyst for photocatalytic hydrogen production, *Ceram. Int.*, 2023, 49, 32246–32260, DOI: 10.1016/
 j.ceramint.2023.07.200.
- 29 M. Sookhakian, Y. M. Amin, W. J. Basirun, M. T. Tajabadi and N. Kamarulzaman, Synthesis, structural, and optical properties of type-II ZnO-ZnS core-shell nanostructure, *J. Lumin.*, 2014, 145, 244–252, DOI: 10.1016/ j.jlumin.2013.07.032.
- 30 S. Ali, S. Saleem, M. Salman and M. Khan, Synthesis, structural and optical properties of ZnS–ZnO nanocomposites, *Mater. Chem. Phys.*, 2020, **248**, 122900, DOI: **10.1016/j.matchemphys.2020.122900**.
- 31 U. P. Gawai, U. P. Deshpande and B. N. Dole, A study on the synthesis, longitudinal optical phonon-plasmon coupling and electronic structure of Al doped ZnS nanorods, *RSC Adv.*, 2017, 7, 12382–12390, DOI: 10.1039/c6ra28180j.
- 32 P. Devaraji, M. Mapa, H. M. A. Hakkeem, V. Sudhakar, K. Krishnamoorthy and C. S. Gopinath, ZnO – ZnS Heterojunctions : A Potential Candidate for Optoelectronics Applications and Mineralization of Endocrine Disruptors in Direct Sunlight, ACS Omega, 2017, 6768–6781, DOI: 10.1021/acsomega.7b01172.
- 33 Y. C. Cheng, C. Q. Jin, F. Gao, X. L. Wu, W. Zhong, S. H. Li and P. K. Chu, Raman scattering study of zinc blende and wurtzite ZnS, *J. Appl. Phys.*, 2009, **106**(12), 123505, DOI: **10.1063/1.3270401**.
- 34 M. Hafeez, S. Rehman, U. Manzoor, M. A. Khan and A. S. Bhatti, Catalyst driven optical properties of the selfassembled ZnS nanostructures, *Phys. Chem. Chem. Phys.*, 2013, 15, 9726–9734, DOI: 10.1039/c3cp50534k.
- 35 N. Prasad and B. Karthikeyan, Resonant and Off-Resonant Phonon Properties of Wurtzite ZnS: Effect of Morphology on Fröhlich Coupling and Phonon Lifetime, *J. Phys. Chem. C*, 2018, **122**, 18117–18123, DOI: **10.1021/acs.jpcc.8b05164**.
- 36 M. Hafeez, T. Zhai, A. S. Bhatti, Y. Bando and D. Golberg, Enhanced field emission and optical properties of controlled tapered ZnS nanostructures, *J. Phys. Chem. C*, 2012, **116**, 8297–8304, DOI: **10.1021/jp3010635**.
- 37 S. S. Lo, Y. T. Hung and D. J. Jan, Stress induction, UV emission variation and efficiency variation in dye-

sensitized solar cells of hollow ZnS/ZnO/ZnS nanostructures, *J. Mater. Chem. A*, 2013, **1**, 10274–10280, DOI: **10.1039**/c3ta12213a.

- 38 M. S. Ghamsari, S. Alamdari, D. Razzaghi and M. Arshadi Pirlar, ZnO nanocrystals with narrow-band blue emission, *J. Lumin.*, 2019, 205, 508–518, DOI: 10.1016/ j.jlumin.2018.09.064.
- 39 J. Ghosh, R. Ghosh and P. K. Giri, Tuning the visible photoluminescence in Al doped ZnO thin film and its application in label-free glucose detection, *Sens. Actuators, B*, 2018, **254**, 681–689, DOI: **10.1016/j.snb.2017.07.110**.
- 40 F. Ajala, A. Hamrouni, A. Houas, H. Lachheb, B. Megna, L. Palmisano and F. Parrino, The influence of Al doping on the photocatalytic activity of nanostructured ZnO: The role of adsorbed water, *Appl. Surf. Sci.*, 2018, 445, 376–382, DOI: 10.1016/j.apsusc.2018.03.141.
- 41 N. P. Herring, L. S. Panchakarla and M. S. El-Shall, P-Type nitrogen-Doped ZnO nanostructures with controlled shape and doping level by facile microwave synthesis, *Langmuir*, 2014, **30**, 2230–2240, DOI: **10.1021/la404593w**.
- 42 Ü. Özgür, Y. I. Alivov, C. Liu, A. Teke, M. A. Reshchikov, S. Doğan, V. Avrutin, S. J. Cho and H. Morkoç, A comprehensive review of ZnO materials and devices, *J. Appl. Phys.*, 2005, **98**, 1–103, DOI: **10.1063/1.1992666**.
- 43 Y. Cheng, R. Chen, H. Feng, W. Hao, H. Xu, Y. Wang, J. Li, F. Li, S. Zhang and H. Sun, Variation of the coordination environment and its effect on the white light emission properties in a Mn-doped ZnO?ZnS complex structure, *Phys. Chem. Chem. Phys.*, 2014, 16, 4544, DOI: 10.1039/ c3cp54830a.
- 44 X. Liu, X. Chen, X. Cui and R. Yu, The structure and multifunctional behaviors of Mn-ZnO/Mn-ZnS nanocomposites, *Ceram. Int.*, 2014, **40**, 13847–13854, DOI: **10.1016/j.ceramint.2014.05.102**.
- 45 D. Zhu, W. Li, L. Ma, Y. Lei, H. Qu, Y. G. Sun, B. H. Dong, M. G. Bawendi and Y. Masumoto, Glutathionefunctionalized Mn:ZnS/ZnO core/shell quantum dots as potential time-resolved FRET bioprobes, *RSC Adv.*, 2014, 4, 9372–9378, DOI: 10.1039/c3ra45491f.
- 46 N. V. Nghia, N. D. Dung, P. T. Huy and D. H. Nguyen, Lasing from ZnO Nanocrystals in ZnO-ZnS Microbelts, *J. Electron. Mater.*, 2017, 46, 3295–3300, DOI: 10.1007/s11664-017-5420-1.
- 47 J. Jiang, H. Xu, L. Zhu, W. Niu, Y. Guo, Y. Li, L. Hu, H. He and Z. Ye, Structural and optical properties of ZnSO alloy thin films with different S contents grown by pulsed laser deposition, *J. Alloys Compd.*, 2014, 582, 535–539, DOI: 10.1016/j.jallcom.2013.07.015.
- 48 X. Gu, S. Zhang, Y. Zhao and Y. Qiang, Band alignment of ZnO/ZnS heterojunction prepared through magnetron sputtering and measured by X-ray photoelectron spectroscopy, *Vacuum*, 2015, **122**, 6–11, DOI: **10.1016**/ j.vacuum.2015.09.005.
- 49 R. R. Thankalekshmi and A. C. Rastogi, Structure and optical band gap of ZnO_{1-x}S_x thin films synthesized by chemical spray pyrolysis for application in solar cells, *J. Appl. Phys.*, 2012, **112**(6), 063708, DOI: **10.1063/1.4754014**.

- 50 Y. Lu, Y. Lin, T. Xie, S. Shi, H. Fan and D. Wang, Enhancement of visible-light-driven photoresponse of Mn/ ZnO system: Photogenerated charge transfer properties and photocatalytic activity, *Nanoscale*, 2012, **4**, 6393–6400, DOI: **10.1039/c2nr31671d**.
- 51 Y. Quan, J. Yao, S. Yang, L. Chen, J. Li, Y. Liu, J. Lang, H. Shen, Y. Wang, Y. Wang, J. Yang and M. Gao, ZnO nanoparticles on MoS₂ microflowers for ultrasensitive SERS detection of bisphenol A, *Microchim. Acta*, 2019, **186**, 4–11, DOI: **10.1007/s00604-019-3702-4**.
- 52 M. A. Hassan, A. Waseem, M. A. Johar, I. V. Bagal, J. S. Ha and S. W. Ryu, Single-step fabrication of 3D hierarchical ZnO/ZnS heterojunction branched nanowires by MOCVD for enhanced photoelectrochemical water splitting, *J. Mater. Chem. A*, 2020, **8**, 8300–8312, DOI: **10.1039**/ **c9ta13714a**.
- 53 M. M. Hasan Farooqi and R. K. Srivastava, Structural, optical and photoconductivity study of ZnO nanoparticles synthesized by annealing of ZnS nanoparticles, *J. Alloys Compd.*, 2017, **691**, 275–286, DOI: **10.1016**/ j.jallcom.2016.08.245.
- 54 A. Kanti Kole, C. Sekhar Tiwary and P. Kumbhakar, Morphology controlled synthesis of wurtzite ZnS nanostructures through simple hydrothermal method and observation of white light emission from ZnO obtained by annealing the synthesized ZnS nanostructures, *J. Mater. Chem. C*, 2014, **2**, 4338–4346, DOI: **10.1039/c4tc00091a**.
- 55 P. C. Patel, S. Ghosh and P. C. Srivastava, Structural, magnetic and optical properties of ZnO nanostructures converted from ZnS nanoparticles, *Mater. Res. Bull.*, 2016, 81, 85–92, DOI: 10.1016/j.materresbull.2016.05.005.
- 56 J. Li, G. J. Fang, C. Li, L. Y. Yuan, L. Ai, N. S. Liu, D. S. Zhao, K. Ding, G. H. Li and X. Z. Zhao, Synthesis and photoluminescence, field emission properties of stalactitelike ZnS-ZnO composite nanostructures, *Appl. Phys. A: Mater. Sci. Process.*, 2008, **90**, 759–763, DOI: **10.1007**/ **s00339-007-4352-7**.
- 57 Q. Shi, Z. Wang, Y. Liu, B. Yang, G. Wang, W. Wang and J. Zhang, Single-phased emission-tunable Mg-doped ZnO phosphors for white LEDs, *J. Alloys Compd.*, 2013, 553, 172–176, DOI: 10.1016/j.jallcom.2012.11.135.
- 58 W. Tian, C. Zhang, T. Zhai, S. L. Li, X. Wang, J. Liu, X. Jie, D. Liu, M. Liao, Y. Koide, D. Golberg and Y. Bando, Flexible ultraviolet photodetectors with broad photoresponse based on branched ZnS-ZnO heterostructure nanofilms, *Adv. Mater.*, 2014, 26, 3088– 3093, DOI: 10.1002/adma.201305457.
- 59 M. Ahmad, X. Yan and J. Zhu, Controlled synthesis, structural evolution, and photoluminescence properties of nanoscale one-dimensional hierarchical ZnO/ZnS heterostructures, *J. Phys. Chem. C*, 2011, **115**, 1831–1837, DOI: **10.1021/jp112047f**.
- 60 Y. Tian, N. Feng, M. Wierzbicka-Wieczorek, P. Huang,
 L. Wang, Q. Shi and C. Cui, Energy transfer-induced
 tunable emission color and thermal quenching of Ca₃Y(PO₄)₃:Eu²⁺,Mn²⁺ phosphor for NUV-pumped white

Paper

LEDs, *Dyes Pigm.*, 2016, **131**, 91–99, DOI: **10.1016**/ **j.dyepig.2016.04.010**.

- 61 H. Guo, Q. Shi, K. V. Ivanovskikh, L. Wang, C. Cui and P. Huang, A high color purity red-emission phosphor based on Sm³⁺ and Eu³⁺ co-doped Ba₃Bi(PO₄)₃, *Mater. Res. Bull.*, 2020, **126**, 110836, DOI: **10.1016**/ j.materresbull.2020.110836.
- 62 J. Zhou, Y. Lai, D. Wu, M. Chen, B. Wang, Z. Tang, L. Lin, M. Chen, J. Ding and Q. Wu, Site-selective excitation and photoluminescence properties of a cyan-emitting $NaGd_9(SiO_4)_6O_2$: Bi³⁺ phosphor for potential application in white light-emitting diodes, *Ceram. Int.*, 2021, 47, 13769– 13775, DOI: 10.1016/j.ceramint.2021.01.239.
- 63 Y. Lan, Z. Sun, Z. Lu, C. Liang and L. Zhou, Synthesis, luminescence property, and application of a novel redemitting, *Opt. Laser Technol.*, 2019, **119**, 105614, DOI: **10.1016/j.optlastec.2019.105614**.
- 64 D. J. Gargas, H. Gao, H. Wang and P. Yang, High quantum efficiency of band-edge emission from ZnO nanowires, *Nano Lett.*, 2011, **11**, 3792–3796, DOI: **10.1021/nl201850k**.
- 65 P. P. Lohe, D. V. Nandanwar, P. D. Belsare and S. V. Moharil, Cyan emitting $Ca_3Sc_2Si_{1.5}Ge_{1.5}O_{12}:Ce^{3+}$ phosphor with 10.4 ns lifetime, *J. Lumin.*, 2019, **216**, 116744, DOI: **10.1016**/ j.jlumin.2019.116744.
- 66 D. Q. Trung, M. T. Tran, N. D. Hung, Q. Nguyen Van, N. T. Huyen, N. Tu and H. Pham Thanh, Emission-tunable Mn-doped ZnS/ZnO heterostructure nanobelts for UVpump WLEDs, *Opt. Mater.*, 2021, **121**, 111587, DOI: **10.1016/j.optmat.2021.111587**.
- 67 Q. Sun, S. Wang, L. Sun, J. Liang, B. Devakumar and X. Huang, Achieving full-visible-spectrum LED lighting via employing an efficient Ce³⁺-activated cyan phosphor, *Mater. Today Energy*, 2020, 17, 100448, DOI: 10.1016/j.mtener.2020.100448.

- 68 X. Wang, Z. Qiu, Y. Liang, Z. Song, S. Li, J. Zhang and S. Lian, Achieving dynamic multicolor luminescence in ZnS:KBr,Mn²⁺ phosphor for anti-counterfeiting, *Chem. Eng. J.*, 2022, **429**, 132537, DOI: **10.1016/j.cej.2021.132537**.
- 69 N. Khichar, S. Bishnoi and S. Chawla, Introducing dual excitation and tunable dual emission in ZnO through selective lanthanide (Er³⁺/Ho³⁺) doping, *RSC Adv.*, 2014, 4, 18811–18817, DOI: **10.1039/c4ra01248h**.
- 70 A. S. Altowyan, M. B. Coban, U. H. Kaynar, E. A. Çin, M. Ayvacikli, J. Hakami and N. Can, Structural and photoluminescent analysis of novel Eu³⁺ and Dy³⁺ Codoped ZnO nanoparticles by incorporation of Li⁺ and K⁺ ions, *Ceram. Int.*, 2024, 50, 14529–14541, DOI: 10.1016/j.ceramint.2024.01.366.
- 71 Y.-L. Yang, X.-C. Yang, J.-Y. Yuan, T. Li, Y.-T. Fan, L. Wang, Z. Deng, Q.-L. Li, D.-Y. Wan, J.-T. Zhao and Z.-J. Zhang, Time-Resolved Bright Red to Cyan Color Tunable Mechanoluminescence from CaZnOS, *Adv. Opt. Mater.*, 2021, 9(16), 2100668, DOI: 10.1002/adom.202100668.
- 72 S. Senapati and R. Naik, Ratiometric optical temperature sensing based on greenish-white light emitting ZnO nanosheets and rhodamine 6G composite, *Mater. Res. Bull.*, 2022, **153**, 111904, DOI: **10.1016**/ **j.materresbull.2022.111904**.
- 73 M. L. A. Letswalo, L. Reddy, A. Balakrishna, H. C. Swart and O. M. Ntwaeaborwa, The role of sulfate ions on distinctive defect emissions in ZnO:Ce³⁺ nanophosphors A study on the application in color display systems, *J. Lumin.*, 2021, 240, 118462, DOI: 10.1016/j.jlumin.2021.118462.
- 74 H. Liu, Y. Zheng, S. Liu, J. Zhao, Z. Song, D. Peng and Q. Liu, Realizing Red Mechanoluminescence of ZnS:Mn²⁺ Through Ferromagnetic Coupling, *Adv. Funct. Mater.*, 2024, 34, 1–9, DOI: 10.1002/adfm.202314422.