

RESEARCH ARTICLE

View Article Online
View Journal | View IssueCite this: *Inorg. Chem. Front.*, 2022,
9, 5820

Interesting dimensional transition through changing cations as the trigger in multinary thioarsenates displaying variable photocurrent response and optical anisotropy†

Chao Zhang,^a Sheng-Hua Zhou,^{b,c,d} Yu Xiao,^a Hua Lin ^{*b,c,d} and Yi Liu ^{*a}

Due to the intriguing component variability and structure–property flexibility, lone-pair cation-based chalcogenides have garnered substantial interest in recent years. Herein, two new multinary thioarsenates, Cs₂ZnAs₄S₈ and [(NH₄)Cs]CdAs₄S₈, were successfully discovered via a surfactant–thermal reaction. Both of them possess identical stoichiometry 2–1–4–8, but they exhibit surprisingly different structural features. Cs₂ZnAs₄S₈ demonstrates a three-dimensional (3D) [ZnAs₄S₈]^{2–} framework made from the corner-sharing [ZnS₄] tetrahedra and one-dimensional (1D) [As₄S₈]^{4–} chains, whereas [(NH₄)Cs]CdAs₄S₈ exhibits a two-dimensional (2D) [CdAs₄S₈]^{2–} layer constructed from the corner-sharing [CdS₄] tetrahedra and tetranuclear [As₄S₈] clusters. Photoelectric measurements display that Cs₂ZnAs₄S₈ has higher photogenerated electron–hole pair separation efficiency than [(NH₄)Cs]CdAs₄S₈ under visible light irradiation. Moreover, both of them show large optical anisotropy ($\Delta n > 0.17$ at 1064 and 2050 nm), while the low dimensional structure is more conducive to enhancing the optical anisotropy based on the theoretical calculations. These findings will provide inspiration for the exploration of multifunctional chalcogenides.

Received 4th August 2022,
Accepted 19th September 2022

DOI: 10.1039/d2qi01694j

rsc.li/frontiers-inorganic

Introduction

Over the past few decades, inorganic chalcogenides containing [M^{III}Q_n] (M^{III} = As, Sb, Bi) polyhedra have attracted increasing attention because of their intriguing structural and compositional diversity.^{1–15} These M^{III} cations with stereochemically active lone-pair electrons are favorable to form different asymmetric coordination modes, which will further influence the electronic structures and physical properties of the resultant chalcogenides. For example, M₂As₂Q₅ (M = Ba, Pb; Q = S, Se) exhibit excellent overall infrared nonlinear optical (IR-NLO) performance thanks to their zero-dimensional (0D) discrete arsenate anions.¹² Pentanary thioantimonate Rb₂Ba₃Cu₂Sb₂S₁₀

adopts a one-dimensional (1D) chain structure and displays a wide-bandgap and an intriguing photocurrent response.⁹ The narrow gap semiconductor RbBi_{11/3}Te₆, which consists of a two-dimensional (2D) infinite Bi₂Te₃-like layer, exhibits a sharp superconducting transition at ~3.2 K.⁴ The three-dimensional (3D) frameworks A₃Mn₂Sb₃S₈ (A = K and Rb) not only display IR-NLO performances but also possess temperature-dependent paramagnetism and photocurrent responses.¹³ On the other hand, transition-metal-based chalcogenides with d¹⁰ electronic configurations (e.g., Zn²⁺, Cd²⁺, Hg²⁺) have also been extensively investigated as these cations tend to show flexible coordination geometries, and hence engender an efficient route to design functional materials.¹⁶ For instance, Na₆Zn₃M₂Q₉ (M = Ga, In; Q = S, Se), 2D layered chalcogenides made of unprecedented T₃-supertetrahedra, exhibit desirable photoluminescence performances.¹⁷ KCd₃Ga₅S₁₁ adopts a diamond-like framework structure and achieves a strong second-harmonic-generation intensity (1.7 × AgGaS₂).¹⁸ Ternary IR-NLO BaHgSe₂ shows large susceptibility and physicochemical stability activated by the trigonal planar [HgSe₃]^{4–} motif.¹⁹

A tremendous amount of multinary lone-pair cation-based chalcogenides have been discovered so far, however, the systematic investigation of the A/TM/As/Q system (A = alkali metals; TM = group 12 metals; Q = chalcogen) is rarely reported. To the best of our knowledge, only 3 compounds in this system are known, namely NaCdAsS₃,²⁰ Rb₄CdAs₂S₉,²¹ and

^aInstitute for Composites Science Innovation (InCSI), School of Materials Science and Engineering, Zhejiang University, Hangzhou 310027, China.

E-mail: liuyimse@zju.edu.cn

^bState Key Laboratory of Structural Chemistry, Fujian Institute of Research on the Structure of Matter, Chinese Academy of Sciences, Fuzhou, Fujian 350002, China.
E-mail: linhua@fjirsm.ac.cn

^cFujian Science & Technology Innovation Laboratory for Optoelectronic Information of China, Fuzhou, Fujian 350108, China

^dUniversity of Chinese Academy of Sciences, Beijing 100049, China

† Electronic supplementary information (ESI) available: Additional experimental and theory results, together with additional tables and figures. CCDC 2167064 and 2167065. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d2qi01694j>

CsHgAsS₃.²² The first two are obtained by the solid-state method, while the last one is prepared by solvothermal synthesis. Recently, a surfactant–thermal reaction has been proved to be a facile method for exploring novel chalcogenides.^{23–31} In this study, our continuous explorations based on the surfactant–thermal method have led to the discovery of two new thioarsenates, Cs₂ZnAs₄S₈ and [(NH₄)Cs]CdAs₄S₈. Although these two compounds have identical stoichiometry 2–1–4–8, they are not isostructural and undergo an interesting dimensional transition from a 3D framework to a 2D layered structure. Herein, the crystal structure, physical properties, and corresponding theoretical studies are systematically investigated.

Results and discussion

Single-crystal X-ray diffraction (SXRD) test results show that Cs₂ZnAs₄S₈ and [(NH₄)Cs]CdAs₄S₈ belong to different space

Table 1 Crystallographic data and refinement details of Cs₂ZnAs₄S₈ and [(NH₄)Cs]CdAs₄S₈

Formula	Cs ₂ ZnAs ₄ S ₈	[(NH ₄)Cs]CdAs ₄ S ₈
FW	887.35	819.51
Crystal system	Tetragonal	Tetragonal
Temperature (K)	Yellow	Orange
Crystal color	<i>I</i> ₄ /a (no. 88)	<i>P</i> ₄ / <i>n</i> (no. 85)
Space group	11.0828(5)	10.3879(2)
<i>a</i> (Å)	11.0828(5)	10.3879(2)
<i>b</i> (Å)	13.2303(8)	7.7691(2)
<i>c</i> (Å)	90	90
α (°)	90	90
β (°)	90	90
γ (°)	1625.06(18)	838.35(4)
<i>V</i> (Å ³)	4	2
<i>Z</i>	3.627	3.246
<i>D</i> _c (g cm ⁻³)	15.004	12.243
μ (mm ⁻¹)	1.039	1.071
GOOF on <i>F</i> ²	0.0265, 0.0603	0.0152, 0.0300
<i>R</i> ₁ , <i>wR</i> ₂ (<i>I</i> > 2 σ (<i>I</i>)) ^a	0.0369, 0.0645	0.0189, 0.0316
<i>R</i> ₁ , <i>wR</i> ₂ (all data)	1.826/–1.271	0.818/–0.552
Largest diff. peak/hole (e Å ⁻³)	887.35	819.51

$$^a R_1 = \sum ||F_o| - |F_c|| / \sum |F_o|, wR_2 = [\sum w(F_o^2 - F_c^2)^2 / \sum w(F_o^2)^2]^{1/2}.$$

groups, *i.e.*, tetragonal *I*₄/a (no. 88) for Cs₂ZnAs₄S₈ with cell parameters of *a* = *b* = 11.0828(5), *c* = 13.2303(8) Å, and *Z* = 4 and tetragonal *P*₄/*n* (no. 85) for [(NH₄)Cs]CdAs₄S₈ with cell parameters of *a* = *b* = 10.3879(2), *c* = 7.7691(2) Å, and *Z* = 2. The detailed crystallographic information is listed in Table 1. There are 1 unique Cs atom (Wyckoff site: 8*e*), 1 unique As atom (Wyckoff site: 16*f*), 1 unique Zn atom (Wyckoff site: 4*a*), and 2 unique S atoms (Wyckoff sites: 16*f* and 16*f*) in its asymmetric unit (Table 2). As illustrated in Fig. 1a, the three-dimensional (3D) crystal structure of Cs₂ZnAs₄S₈ is constructed from charge-balanced Cs⁺ cations, regular [ZnS₄] tetrahedra, (Fig. 1b) and one-dimensional (1D) anionic chains [As₄S₈]⁴⁻ made of the corner-sharing [As₄S₉] groups (Fig. 1c). The important bond distances of Cs₂ZnAs₄S₈ are listed in Table S1.† The As atom adopts a common coordination with 3 S atoms in the bond distance range of 2.2143(5)–2.3267(5) Å to build a [AsS₃]³⁻ triangular pyramid. The Zn–S bond distance in the [ZnS₄] tetrahedron is 2.3309(5) Å. The S–As–S angles in Cs₂ZnAs₄S₈ range from 98.07(2)° to 103.28(2)°, while the S–Zn–S angles vary from 105.79(2)° to 117.12(2)°. These bond distances and angles in Cs₂ZnAs₄S₈ are normal and can also be comparable to the reported Cs-based chalcogenides.^{32–34}

There are 7 unique crystallographic atoms in the asymmetric unit of the structure of [(NH₄)Cs]CdAs₄S₈, including 1 Cs (Wyckoff site: 2*c*), 1 Cd (Wyckoff site: 2*a*), 1 As (Wyckoff site: 8*g*), 2 S (Wyckoff sites: 8*g* and 8*g*), 1 N (Wyckoff site: 2*b*) and 1 H (Wyckoff site: 8*g*) (Tables 1 and 2). As given in Fig. 1d, the crystal structure of [(NH₄)Cs]CdAs₄S₈ is composed of regular [CdS₄] tetrahedra with *d*_(Cd–S) = 2.5481(5) Å and tetranuclear [As₄S₈] clusters with *d*_(As–S) = 2.2676(5)–2.3008(5) Å, which interconnect with each other by sharing vertexes to form a two-dimensional (2D) [CdAs₄S₈]²⁻ layer filling the dispersed (NH₄)⁺ and Cs⁺ cations.

The interesting structural evolution between Cs₂ZnAs₄S₈ and [(NH₄)Cs]CdAs₄S₈ is illustrated in Fig. 1. Both of them possess an identical stoichiometry of 2–1–4–8 and belong to the same tetragonal system, but they are not isostructural and have some significantly different structural features: (i) pyramidal [AsS₃] units in Cs₂ZnAs₄S₈ forms a 1D infinite [As₄S₈]⁴⁻

Table 2 Atomic coordinates and equivalent isotropic displacement parameters of Cs₂ZnAs₄S₈ and [(NH₄)Cs]CdAs₄S₈

Atom	Wyckoff	<i>x</i>	<i>y</i>	<i>z</i>	<i>U</i> _(eq) ^a	Occu.
Cs₂ZnAs₄S₈						
Cs	8 <i>e</i>	0	0.25	0.45521(2)	0.02597(6)	1.0
As	16 <i>f</i>	0.16243(2)	0.01798(2)	0.00487(2)	0.01245(5)	1.0
Zn	4 <i>a</i>	0	0.25	0.125	0.01320(8)	1.0
S2	16 <i>f</i>	0.17589(5)	0.21445(4)	0.03310(3)	0.01605(9)	1.0
S1	16 <i>f</i>	0.31599(5)	0.51563(5)	0.10909(3)	0.01892(10)	1.0
[(NH₄)Cs]CdAs₄S₈						
Cs	2 <i>c</i>	0.25	0.25	0.68028(3)	0.02210(7)	1.0
Cd	2 <i>a</i>	0.25	0.75	0	0.01361(7)	1.0
As	8 <i>g</i>	0.04860(2)	0.15875(2)	0.14479(2)	0.01204(6)	1.0
S2	8 <i>g</i>	0.16315(4)	0.02562(4)	0.31813(6)	0.01556(10)	1.0
S1	8 <i>g</i>	0.07233(4)	0.64420(4)	0.17643(6)	0.01402(9)	1.0
N	2 <i>b</i>	0.25	0.75	0.5	0.0082(5)	1.0
H	8 <i>g</i>	0.2013(19)	0.724(2)	0.446(3)	0.014(6)	1.0

^a *U*_(eq) is defined as one-third of the trace of the orthogonalized *U*_{*ij*} tensor.

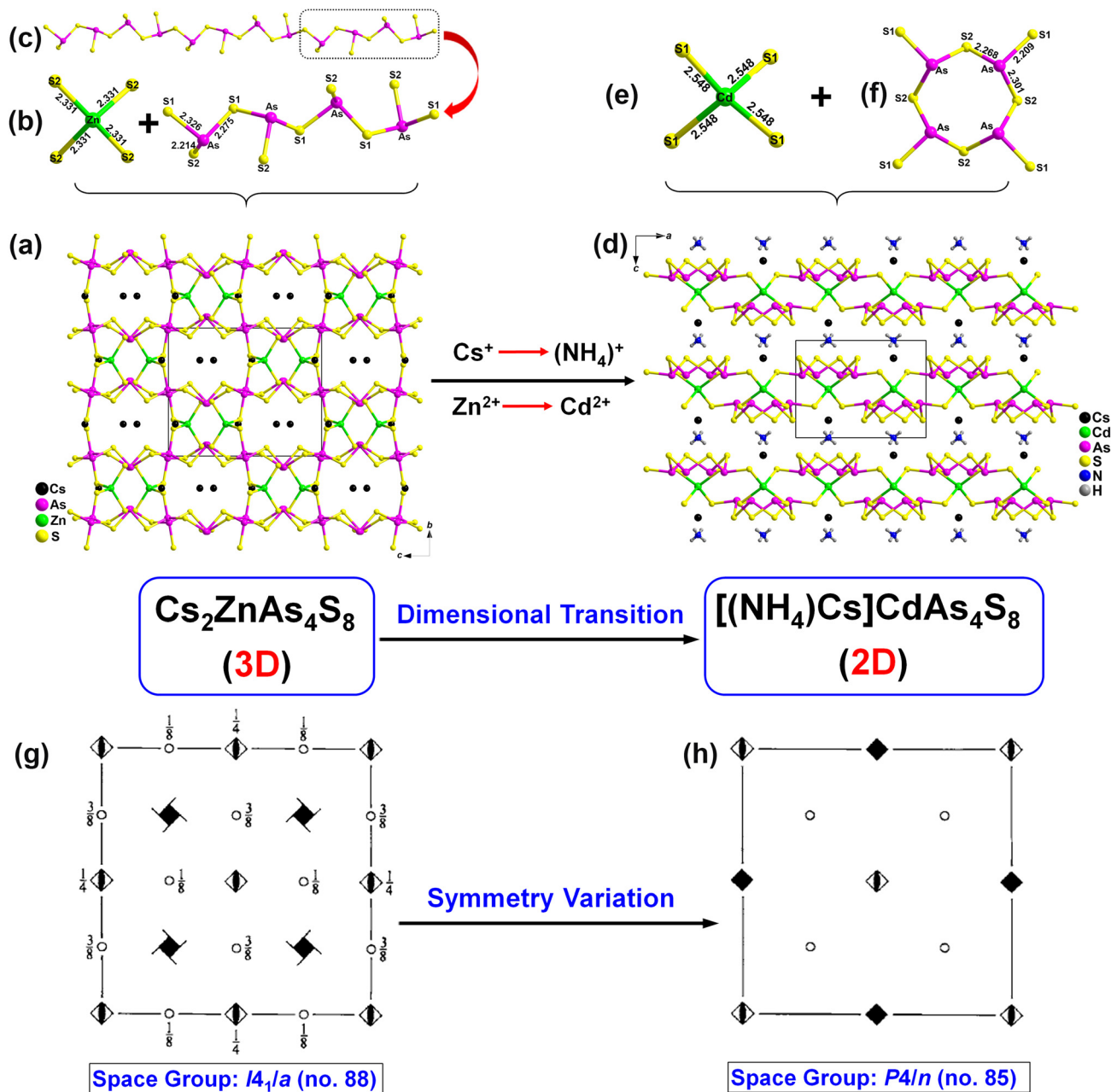


Fig. 1 Dimensional transition from 3D $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ to 2D $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ based on the isovalent cation substitution. (a) Crystal structure of 3D $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ viewed down the a -direction; (b) 1D $[\text{As}_4\text{S}_8]$ chain with the coordination environment of the $[\text{As}_4\text{S}_9]$ unit marked; (c) the coordination environment of the tetrahedral $[\text{ZnS}_4]$ unit; (d) crystal structure of 2D $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ viewed down the a -direction; (e, f) the coordination environment of the tetrahedral $[\text{ZnS}_4]$ unit and tetra-nuclear $[\text{As}_4\text{S}_8]$ cluster; (g, h) spatial symmetry operation from $I4_1/a$ (no. 88) to $P4/n$ (no. 85).

chain by vertex-sharing S1 atoms, while in $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, the tetra-nuclear $[\text{As}_4\text{S}_8]$ cluster is constructed from the same $[\text{AsS}_3]$ units; (ii) the coordination number (CN) of the crystallographically independent Cs atom in $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ is 10 and the Cs–S bond distances are in the range of 3.617 (5)–4.116 (6) Å, which are different from those of $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ [CN = 8, $d_{(\text{Cs}-\text{S})} = 3.696(5)\text{--}3.76(5)$ Å] (Fig. S1†). Such differences between them can be attributed to the different sizes of cations, *i.e.*, the 3D $[\text{ZnAs}_4\text{S}_8]^{2-}$ framework in $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ can be viewed as $[\text{ZnS}_4]$ tetrahedra connected by 1D infinite $[\text{As}_4\text{S}_8]^{4-}$

chains by sharing S atoms and relatively large Cs^+ cations occupied the space of the framework. However, when one of the “large” radius Cs^+ cations was replaced by smaller radius $(\text{NH}_4)^+$ cations, the linkages between the $[\text{As}_4\text{S}_8]^{4-}$ chains in $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ are broken, and the 3D $[\text{ZnAs}_4\text{S}_8]^{2-}$ framework in $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ transforms into the 0D $[\text{As}_4\text{S}_8]$ clusters in $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$. In addition, the above-mentioned structural evolution is ultimately reflected by their space groups, from $I4_1/a$ (for $\text{Cs}_2\text{ZnAs}_4\text{S}_8$) to $P4/n$ (for $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$). The detailed symmetric operation change based on isovalent cation substitution is shown in Fig. 1g and h.

Polycrystalline samples of $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ were prepared by a facile surfactant-thermal method at 413 K for 7 days with $\text{CsOH}\cdot\text{H}_2\text{O}$, Zn (or Cd), As_2S_3 , S, oleic acid, hydrazine monohydrate (98%) and PEG-400 as starting materials, in a yield of approximately 80–90% based on Zn (or Cd) (further experimental details see the ESI†). As displayed in Fig. S2 and S3,† semi-quantitative energy-dispersive X-ray (EDX) elemental analysis provides the average atomic ratios of 2.0/1.1(2)/3.9(8)/8.2(1) and 1.0/1.1(4)/4.0(5)/7.9(7) for $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, respectively. In addition, the purity of the polycrystalline samples was confirmed by powder X-ray diffraction (XRD) analysis (Fig. S4 and S5†). Moreover, both of them exhibit desirable thermal stability (up to 700 K) under N_2 conditions, as shown in Fig. S6 and S7.† Based on the different calculated formulas for direct semiconductors ($(\alpha h\nu)^2$ vs. energy) or indirect semiconductors ($(\alpha h\nu)^{1/2}$ vs. energy),³⁵ the experimental energy gaps (E_g) of $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ are about 2.24 and 2.36 eV, respectively (Fig. 2). These values are compared to those of other reported quaternary thioarsenates, such as $\text{RbCu}_4\text{AsS}_4$ ($E_g = 2.15$ eV),³⁶ $\text{Cs}_3\text{CuAs}_4\text{S}_8$

($E_g = 2.26$ eV),³⁰ $\text{Rb}_8\text{Cu}_6\text{As}_8\text{S}_{19}$ ($E_g = 2.29$ eV),³⁷ and $\text{CsCu}_2\text{AsS}_3$ ($E_g = 2.30$ eV).²⁷

In addition, inspired by recent reports that most of the lone-pair-based chalcogenides display intriguing photocatalytic properties,^{38–43} the photoelectrochemical experiment was performed through a standard three-electrode system using simulated solar light illumination to study the photoelectric properties of $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$. As illustrated in Fig. 3, the photocurrent-time curves exhibit a rapid and consistent photocurrent response in a multiple 20 s switching period. Clearly, $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ shows a remarkable transient photocurrent response, which is about 4 times that of $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, that is, $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ possesses higher photogenerated electron-hole pair separation efficiency than $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ under visible light irradiation. Meanwhile, these repeatable anodic photocurrent responses suggest that $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ belong to n-type semiconductors. It is worth mentioning that these values of photocurrent densities (*ca.* 3.0 and 0.75 $\mu\text{A cm}^{-2}$ for $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, respectively) are much higher than those of most recently reported chalcogenides, such as $\text{Rb}_2\text{Ba}_3\text{Cu}_2\text{Sb}_2\text{S}_{10}$ (*ca.* 6 nA cm^{-2}),⁹ BaCuSbSe_3 (*ca.* 30 nA cm^{-2}),³⁸ and BaCuSbS_3 (*ca.* 55 nA cm^{-2}).³⁸

To well understand the origin of difference of electronic structures and structure-activity relationships, theoretical calculations of $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ have been systematically carried out based on DFT methods. As shown in Fig. 4a and b, the valence band maximum (VBM) and the conduction band minimum (CBM) located at the same high-symmetry points (*i.e.*, Z|R) indicate that $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ is a direct-band-gap ($E_g = 1.92$ eV) semiconductor, while $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ is an indirect-band-gap ($E_g = 2.13$ eV) semiconductor since the VBM and the CBM are located at different high-symmetry points (*i.e.*, Z|R and V, respectively). These calculated values are slightly smaller than the experimental observations

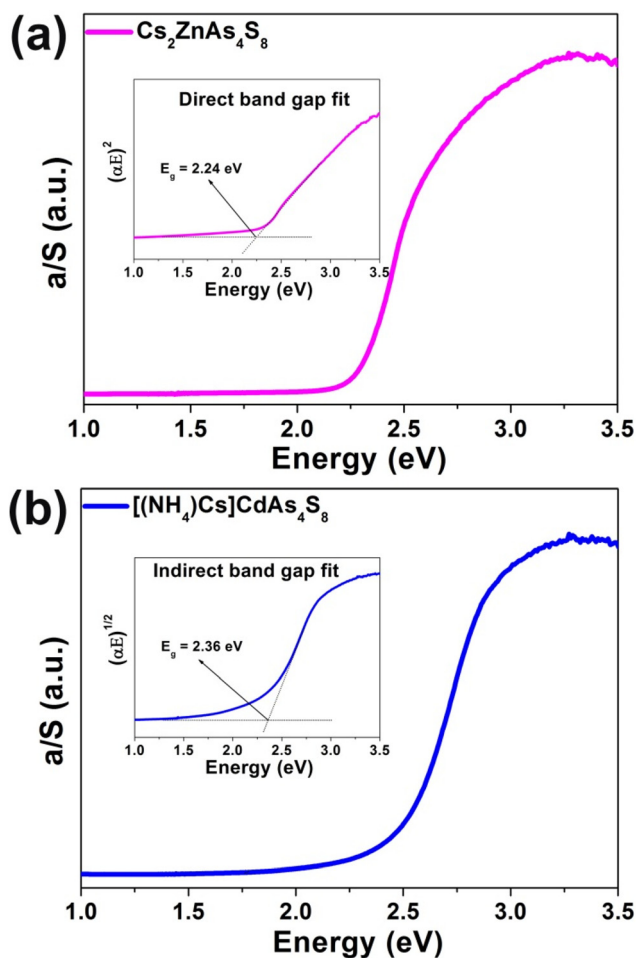


Fig. 2 UV-vis-NIR diffuse reflectance spectra of (a) $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and (b) $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ (inset: direct or indirect band gap fit based on the DFT results).



Fig. 3 Photocurrent response curves of $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$.

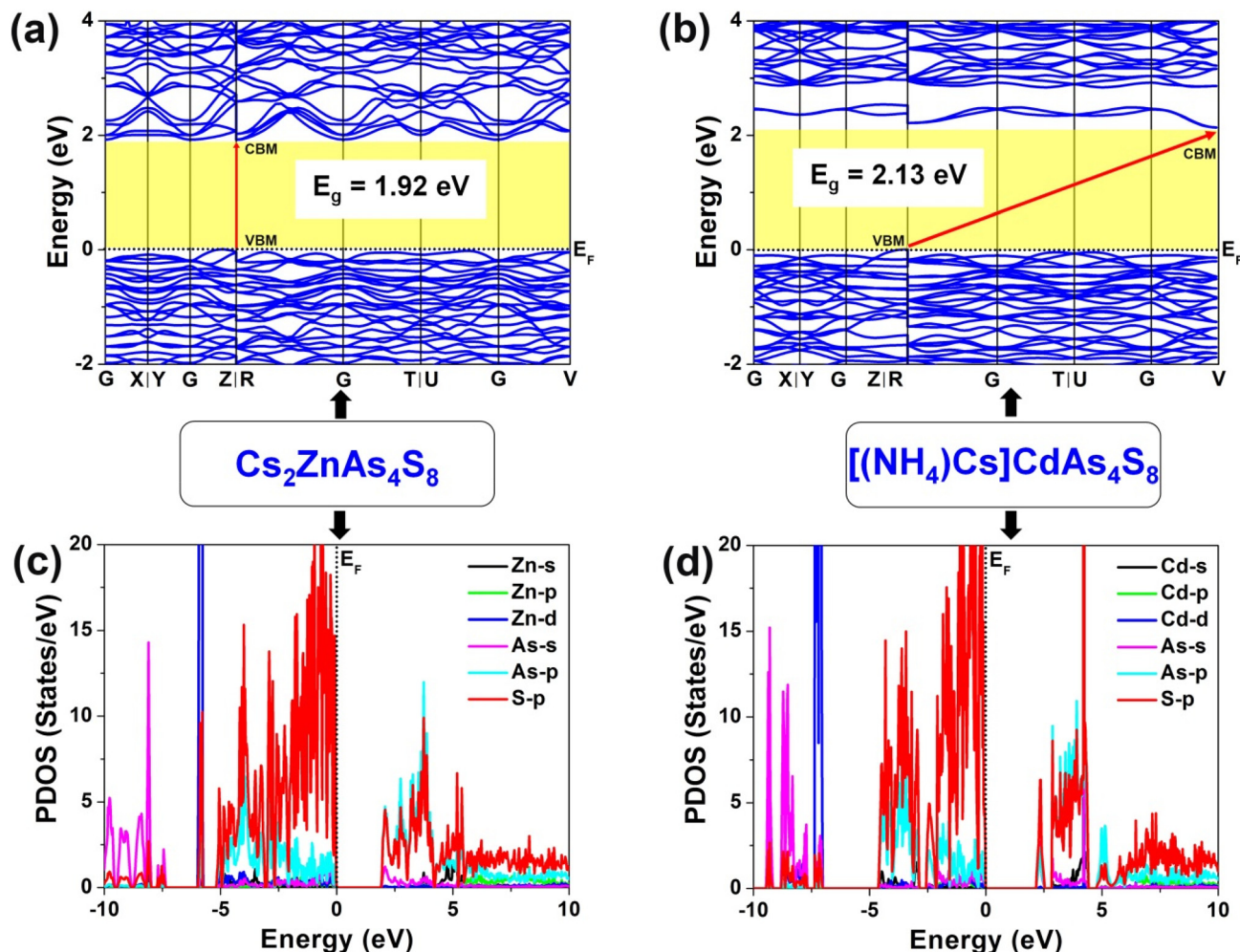


Fig. 4 Theoretical calculated results of $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$: (a, b) electronic band structures; (c, d) PDOSs (states with fewer contributions are omitted for a better view). The Fermi level E_F is set at 0.0 eV.

(2.24 eV for $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and 2.36 eV for $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, as given in Fig. 2), which is a well-known phenomenon for the local approximations to DFT.⁴⁴ It is widely known that the photocatalytic activity mainly depends on the separation and diffusion rate of the photogenerated charge carriers, which can be judged by calculating the relative effective masses of electrons and holes based upon the electronic structures of the VBM and CBM in the title compounds.³⁸ Through comparison, we can clearly see that the electronic bands of $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ are steeper than those of $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$. Namely, $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ is more conducive to improving the efficiency of photocatalysis, which is basically consistent with the experimental observation (Fig. 3). In addition, the projected density of states (PDOSs) with major contributions of title compounds is illustrated in Fig. 4c and d. $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ exhibit similar results: the band from -5 eV to the Fermi level (E_F), S-3p as well as As-4p and Zn-3d (or Cd-4d) states makes the main contribution, while the bottom of the CB is mainly composed of electronic hybridization of S-3p, As-4p and As-4s

states. Consequently, the optical E_g values of $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ mainly come from the charge transfer of Zn (or Cd)-S and As-S units, in which charge-balancing cations (e.g., Cs^+ , $(\text{NH}_4)^+$) show negligible contributions for the DOS. On the basis of their electron structures, we also calculated the birefringence (Δn) of $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, respectively. As displayed in Fig. 5, their Δn values in both the important wavelengths can be calculated as 0.17@1064 nm and 0.18@2050 nm for $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and 0.29@1064 nm and 0.31@2050 nm for $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, which are comparable with some recently reported chalcogenides, indicating that they have potential as UV-vis or IR birefringent crystals. As known, the Δn value will mainly depend on the anisotropy of the anionic substructure, whereas the contribution of charge-balancing cations can be neglected.^{45–50} In other words, compared with the 3D anionic $[\text{CdAs}_4\text{S}_8]^{2-}$ framework in $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, the anisotropy of the 2D anionic $[\text{ZnAs}_4\text{S}_8]^{2-}$ layer in $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ is more obvious, which is more beneficial for producing larger Δn values.

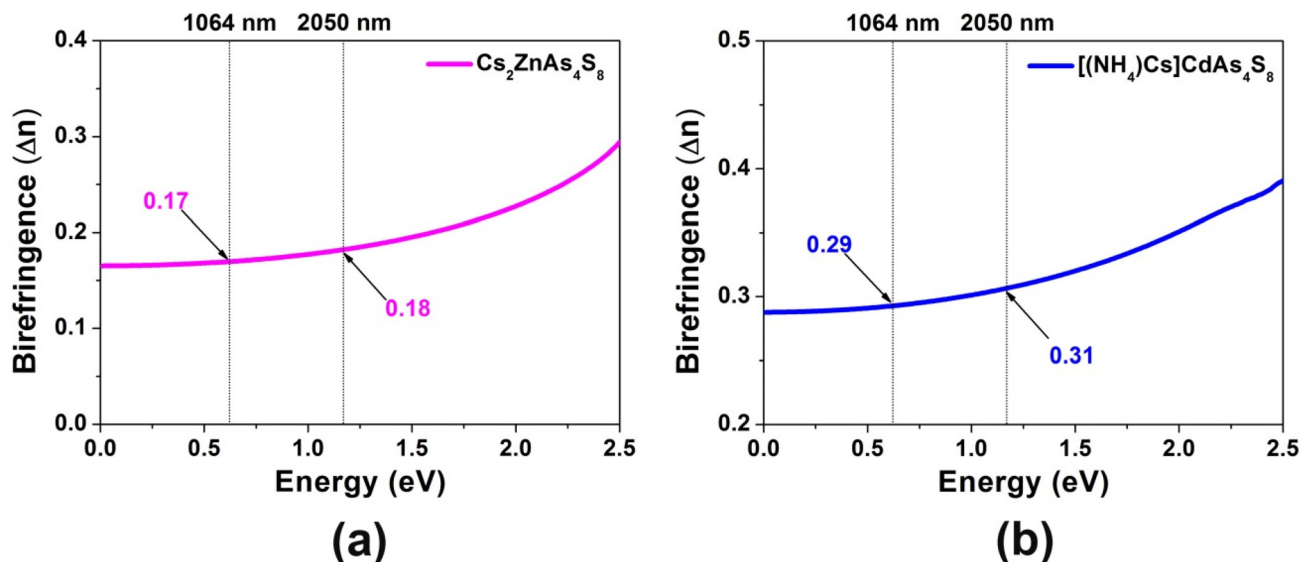


Fig. 5 Curves of the calculated birefringence (Δn) as a function of energy (eV) for (a) $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and (b) $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$.

Conclusions

In conclusion, two new members of the multinary X-TM-As-S family, $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ and $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, have been prepared by a simple surfactant-thermal method. Although they have similar chemical stoichiometry 2-1-4-8, they undergo an intriguing dimensional transition from a 3D framework to a 2D layered structure. The optical absorption spectra and photoelectric results confirm that both the thioarsenates are wide-bandgap semiconductors and $\text{Cs}_2\text{ZnAs}_4\text{S}_8$ exhibits a better photocurrent response than $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$ under the same test conditions. Meanwhile they also show large birefringence. In particular for $[(\text{NH}_4)\text{Cs}]\text{CdAs}_4\text{S}_8$, the calculated birefringence values are 0.29@1064 nm and 0.31@2050 nm, respectively, suggesting its potential for application as a dual-waveband birefringent crystal. The analysis results of the structure-activity relationship show that the low dimensional structure of this family will be favorable for the generation of large optical anisotropy, that is, to obtain high birefringence. These results provide new insights into the exploration of novel functional chalcogenides and further research on the other physical properties of the title compounds is ongoing.

Author contributions

C. Zhang prepared the samples, and designed and carried out the experiments. S. H. Zhou carried out the theoretical calculations. Y. Xiao measured the optical properties. H. Lin and Y. Liu conceived the experiments, analyzed the results and wrote and edited the manuscript. All the authors have approved the final version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (No. 52171277, 22175175 and 21771179), the Natural Science Foundation of Fujian Province (No. 2019J01133), the Fujian Science & Technology Innovation Laboratory for Optoelectronic Information of China (2021ZR118) and the Shanxi-Zheda Institute of Advanced Materials and Chemical Engineering (2022SZ-TD006).

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