Morphology Controlled Synthesis of Composition Related Plasmonic CuCdS Alloy Nanocrystals

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Cu-based ternary alloy nanocrystals have emerged for extensive applications in solar cells, light-emitting devices (LEDs), and photoelectric detectors because of their low-toxicity, tunable band gaps, and large absorption coefficients. It is still an enormous challenge that regulating optical and electrical properties through changing their compositions and shapes in alloy nanocrystals. Herein, we present a facile method to synthesize CuCdS alloy nanocrystals (NCs) with tunable compositions and shapes at relatively low temperature. Different morphologies of monodisperse CuCdS nanocrystals are tailored successfully by simply adjusting the reaction temperature and Cu:Cd precursor molar ratio. The as-synthesized nanocrystals are of homogeneous alloy structures with uniform obvious lattice fringes throughout the whole particles rather than heterojunction structures. The localized surface plasmon resonance (LSPR) absorption peaks of CuCdS NCs are clearly observed and can be precisely tuned by varying the Cu:Cd molar ratio. Moreover, current–voltage (I–V) behaviors of different shaped CuCdS nanocrystals show certain rectification characteristics. The alloy CuCdS NCs with tunable shape, band gap, and composition possess a potential application in optoelectronic devices.

Keywords: morphology, composition, plasmonic, CuCdS, alloy nanocrystals

INTRODUCTION

Cu-based ternary semiconductor nanocrystals are of great scientific interest for applications in solar cells (Rivest et al., 2011), light-emitting devices (LEDs) (Yoon et al., 2019; Li et al., 2020), biological imaging (Meng et al., 2017), photo-catalytic (Zhao et al., 2016), photovoltaic detectors because of their low-toxicity, tunable band gaps, and high absorption coefficients (Pejova et al., 2020). Especially, ternary nanocrystals based on Cu1.94S, with localized surface plasmon resonance due to the large number of copper vacancies, have been widely demonstrated for photoelectrochemical water reduction and solar cells due to their high concentration of free charge carriers (Liu et al., 2017; Bai et al., 2020; Bera et al., 2020; Zhu et al., 2020). Developing the size-, shape-, and composition-controlled composite nanocrystals is a significant strategy to regulate their energy level position, carrier concentration, and photoluminescence quantum yield for the improvement of device performance.
Semiconductor–semiconductor hetero-nanostructures have been widely synthesized by hot-injection method and seed growth method (Yi et al., 2010; Shen H. et al., 2012; Li et al., 2013, 2014a). Antecedently, researches on transition-metal and II-VI composite nanocrystals mainly focused on semiconductor–semiconductor heterostructures such as Ag2S-ZnS, Ag2S-CdS, Ag2Se-ZnS, CuS-ZnS, Cu1.94S-CdS, Cu1.94S-Zn4Cd1.94S, which have been applied in related optoelectronic devices (Regulacio et al., 2011; Shen et al., 2011a; Zhang et al., 2011; Han et al., 2012; Xu et al., 2013; Wang et al., 2014). And the Cu1–xSx-CdS has been particularly investigated by researchers. For example, it has been reported in 1980 that the design and fabrication of thin-film CdS/Cu2S heterojunction solar cells with energy conversion efficiencies in sunlight of up to 9.15% (Devaney et al., 1979). Wang et al. employed n-CdS/p-Cu2S coaxial nanowire to fabricate photovoltaic devices (Pan et al., 2012). However, alloyed composite semiconductors of transition-metal and II-VI elements are extremely rare. It tends to evolve into heterostructures and is difficult to form alloyed nanostructures (Regulacio et al., 2011; Shen et al., 2011a; Wang et al., 2014; Xu et al., 2020). However, alloy nanomaterials have shown excellent characteristics, for instance, alloy quantum dots of II–VI and I–III–VI group show exceptional photovoltaic properties and stability of fluorescence (Swafford et al., 2006; Jun and Jang, 2012; Panthani et al., 2013; Feng et al., 2020). Recently, Cu-based ternary nanostructures such as Cu2Bi2S3 and CuMnS NGS (M: In and Ga) were prepared through a cation exchange process. Interestingly, these ternary nanocrystals possessed the alloyed nanostructure instead of a heterostructure. And they revealed enhanced photocatalytic activities toward photoelectrochemical water reduction, owing to their tailored optical band gaps, energy level alignments, and shape affecting (Bera et al., 2020; Zhu et al., 2020). Therefore, further exploiting new methods to synthesize highly crystallized transition-metal and II–VI alloyed nanostructures still worth systematical study.

Herein we report the formation of CuCdS ternary alloy semiconductors via a facile one-pot colloidal route. CuCdS nanocrystals with adjustable shapes such as fusiform, rod, hexagon, and rhombus were synthesized by heating CuI and Cadmium diethyldithiocarbamate (Cd(DDTC)2), DDTC=diethyldithiocarbamate (Cd(DDTC)2), DDTC=diethyldithiocarbamate) as precursors at low temperature. These shapes were tuned by controlling the reaction temperature and Cu/Cd precursor molar ratio. High resolution transmission electron microscopy (HRTEM) and X-ray diffraction (XRD) results indicated the homogeneous alloy structure and high crystallinity. The localized surface plasmon resonance (LSPR) absorption spectrum and current–voltage (I–V) behaviors of different shaped CuCdS nanocrystals have also been studied.

**MATERIALS AND METHODS**

**Materials**

Copper iodide (CuI, 98%), Sodium N-diethyldithiocarbamate [Na(DDTC), 99%], and Cd(NO3)2·4H2O (95%) were purchased from XiYa Reagent Company (P. R. China). Oleylamine (OAM, 70%) was purchased from Sigma-Aldrich. Hexanes (analytical grade) and acetone (analytical grade) were purchased from Tianjin Chemical Reagents Co. Ltd. All chemicals were used as received without any further purification.

**Synthesis of Cd(DDTC)2 Precursors**

The preparation of Cd(DDTC)2 was referred to a previously published procedure (Shen S. et al., 2012). Typically, Na(DDTC) (18.02 g, 80 mmol) and Cd(NO3)2·4H2O (12.34 g, 40 mmol) were, respectively, dissolved in 200 mL of deionized water by ultrasonication. Subsequently, Cd(NO3)2 solution was dropwise added into Na(DDTC) solution and kept stirring for 30 min. The resulting white precipitate was generated immediately when the reaction started. Then, Cd(DDTC)2 was formed and washed for several times by deionized water and ethanol, and finally was dried in air at room temperature.

**Synthesis of CuCdS**

In a typical procedure, CuI (0.4 mmol) and Cd(DDTC)2 (0.1 mmol) were mixed with 8 mL of OAM in a 25 mL three-necked flask at room temperature. The mixture was heated to 120°C at a rate of 15°C/min under magnetic stirring and nitrogen flow for 10 min, and then cooled to room temperature. The brown product was collected by centrifugation (4,000 rpm, 10 min), and washed several times by acetone for further characterization. By changing the molar ratio of Cu/Cd and reaction temperature, different shaped nanostructures could be obtained.

**Device Fabrication**

A layered structure of Al/CuCdS/ITO was fabricated for conductivity test. Fixed amount of CuCdS nanocrystals solution (30 mg/mL, dispersed in hexanes) was applied directly to pre-cleaned ITO-coated glass substrates (≤40 Ω/sq.) by spin-casting. And uniform nanocrystal thin films were formed. Finally, Al electrodes were deposited by RF magnetron sputtering through a metal mask to form a sandwich structured device.

**Measurements and Characterization**

Thermal gravimetric analysis (TGA) was obtained by a TG/DTA 6200 thermogravimetric analyzer (SII Inc.) under nitrogen flow (200 cm3/min) with a heating rate of 10°C/min. Transmission electron microscopy (TEM) and high resolution transmission electron microscopy (HRTEM) images were taken by a JEOL JEM-2010 microscope with an accelerating voltage of 200 kV. X-Ray diffraction (XRD) results of nanocrystals were obtained employing an X-ray diffractometer (Philips X’ Pert Pro) using Cu-Kα radiation (λ = 1.54 Å) with a step size of 0.02° and a scan rate of 4°/min at a voltage/current of 40 kV/40 mA. Energy-dispersive X-ray spectroscopy data (EDS) was presented on an FEI Quanta 200 scanning electron microscope equipped with a Thermo NORAN System SIX energy dispersive X-ray spectrometer. UV–vis absorption was gained using UV–vis spectrometry (PE Lambda 950). I–V measurements were acquired using Keithley 4200 source meter.
RESULTS AND DISCUSSION

In this report, CuI and Cd(DDTC)$_2$ which acted as the reactants were directly dispersed in oleylamine. Among them, Cd(DDTC)$_2$ as single-source precursor can simultaneously provide sulfur source and cadmium source (Supplementary Figure 1). The as-prepared Cd(DDTC)$_2$ was air-stable and low-toxic. The TGA results illustrated that the decomposition temperature of Cd(DDTC)$_2$ in its pure solid form was between about 250 and 350°C in the nitrogen gas environment (Supplementary Figure 2). Owing to the presence of oleylamine, various morphologies of CuCdS alloy nanostructures were synthesized at relatively low temperature. The influence of the reaction conditions on the architectures of CuCdS nanocrystals was mainly researched by changing molar ratio of Cu:Cd and reaction temperature. TEM was used to observe the morphology of as-prepared CuCdS nanocrystals. Figure 1 shows four different shapes of CuCdS alloyed nanostructures, which were obtained by changing the molar ratio of Cu:Cd at 120°C with reaction time of 10 min. CuCdS crystals, which possess different surface energies in different directions, allow the shape of the nanoparticles to be controlled by increasing and/or decreasing the growth rate in different directions (Han et al., 2008). We changed the molar ratio of Cu:Cd from 1:1 to 8:1 and kept the other parameters the same. Low Cu ratio (Cu:Cd = 1:1) led to the formation of fusiform morphology, as shown in Figure 1a. However, further changing the Cu:Cd ratio to 2:1 led to the formation of rod shape morphology with average length of 44 nm and width of 24 nm (Figure 1b). Contrastively, when the Cu/Cd ratios were increased to 4:1 and 8:1, rod-shaped hexagon and regular hexagonal nanocrystals were finally obtained, respectively (Figures 1c,d).

From the XRD results (Figure 1e), we can find that the diffraction patterns of as-synthesized CuCdS nanocrystals could be regarded as the combination of diffraction patterns of monoclinic Cu$_{1.94}$S (JCPDS No. 23-0959) and wurtzite CdS (JCPDS No. 80-0006). With the increase of Cu content in the reaction, the intensity of Cu$_{1.94}$S peak gradually increased and the intensity of CdS gradually decreased. The evolution of XRD patterns showed that different composed CuCdS nanocrystallines were successfully synthesized without impurities.

The corresponding high-resolution TEM (HRTEM) images of CuCdS nanocrystals synthesized at 120°C with reaction time of 10 min using different Cu/Cd molar ratios are shown in Figure 2. All the HRTEM images of CuCdS nanocrystals show good crystallinity with uniform lattice fringes throughout the whole particle, indicating the homogeneous CuCdS alloy structures, rather than the Cu$_{1.94}$S-CdS heterostructure. The analysis of the different light and dark lattice spacings reveals $d_{\text{light}} = 0.268$ nm and $d_{\text{dark}} = 0.341$ nm, which fit the interplanar spacing of monoclinic Cu$_{1.94}$S (1000) and wurtzite CdS (002) planes, respectively. EDS analysis in Table 1 quantifies the elemental compositions of CuCdS alloy nanocrystals. Cu, Cd, and S signals are all detected, indicating that the alloy nanostructures are composed mainly of Cu, Cd, and S. All the molar ratio of Cu:Cd in products were greater than that of raw materials, declaring that the activity of Cu was higher than that of Cd.

Besides the molar ratio of the reactants, we found that the reaction temperature could also significantly influence the reaction kinetics and the morphology of the obtained CuCdS nanocrystals. For example, when the molar ratio
of Cu:Cd was kept of 1:1, and changing the reaction temperature to 100 and 150°C (Figures 3a,e), the shape of CuCdS ternary semiconductors remained fusiform with different length. At the 2:1 Cu/Cd molar ratio, with the reaction temperature decreased to 100°C, CuCdS nanocrystals had some agglomeration with inhomogeneous rectangular and hexagonal morphology (Figure 3b); while, uniform elliptical CuCdS materials were synthesized at 150°C (Figure 3f). The nanocrystals synthesized with Cu:Cd = 4:1 showed rod-shaped hexagon without obvious corner angle at 100°C and unified rhombus at 150°C (Figures 3c,g). For the products with a Cu:Cd molar ratio of 8:1, rising or falling of the temperature similarly caused the change of morphology to nearly rotundity or homogeneous jujube pit (Figures 3d,h). Based on experimental data and literatures, Generally, Cu ions combined with Cd(DDTC)2 forming CuDDTC. Then, the CuDDTC decomposed to obtain Cu$_{1.94}$S crystals with a dominating (100) plane (Han et al., 2012). CdDDTC decomposed to form Cds nanorods with a prominent (002) plane in OAM solution (Shen et al., 2011b). Simultaneously, the evolution of the reaction process includes a cation exchanging between Cu ions and Cd ions to form CuCdS crystals. In summary, the Cu ions dominated the (100) planes of CuCdS crystals and the $\text{S}^-$ ions were strongly combined with Cu in the precursor. Moreover, Cd ions dominated the (002) planes of CuCdS and were only weakly bonded by the DDTC ligands, the difference of reactivity on different crystal planes resulted in the various shapes. From the results in Figures 1, 3, we found that with the decrease of Cd content at different reaction temperatures, the length to width ratio of CuCdS nanocrystals was reduced.

To give an in-depth understanding of the evolution of CuCdS NCs, the effect of reaction time was studied. The size and morphology evolution of the NCs were monitored by TEM observation as shown in Figure 4. Nanocrystals prepared with the reaction time of 1 min and 10 min were almost the same appearance, which indicates that the reaction occurred in the extremely instantaneous time. In the rest of the reaction time, Ostwald ripening dominated the growth process, and the CuCdS nanocrystallines appeared tiny increase in size and tended to be more uniform and dispersive.

The LSPR related absorption spectra of CuCdS nanostructures with different morphologies by reaction at 120°C are shown in Figure 5. The CuCdS NCs exhibited a NIR-LSPR peak which had a broad band. And the plasmonic absorption band exhibits a red-shift from 960 to 1,067 nm with the decreasing of Cu content in the nanocrystals. The strong LSPR absorption peak was clearly detected at 960 nm for products using 8:1 Cu:Cd molar ratio. With the decreasing of Cu content to Cu:Cd = 4:1, the LSPR absorption peak red-shifted to 1,010 nm. Further decrease of the amount of Cu caused the LSPR peak further red-shifted to 1,043 nm for the NCs prepared using 2:1 and 1:1 Cu:Cd molar ratios. According to previous reports, the LSPR related absorption is dependent on the free carrier density, high frequency dielectric constant, and crystalline anisotropy (Zhou et al., 2016; Zhu et al., 2016). Based on the literatures, the dominant influence of crystalline anisotropy usually results in the adjustable LSPR absorption, which has been demonstrated experimentally and theoretically (Kim et al., 2016). The effect of crystalline anisotropy includes the size, shape and aspect ratio of nanocrystal. The LSPR absorption peak shows a continuous red shift with the decrease of Cu content in these nanocrystals, which is attributed to the decrease of the carrier density (Zhou et al., 2016).

The composition and valence states of nanocrystals synthesized with 4:1 Cu/Cd molar ratio and 120°C reaction temperature were further identified by X-ray photoelectron spectroscopy (XPS), as shown in Figure 6. XPS survey spectrum presents the presence of C 1s, Cu 2p, Cd 3d, and S 2p core level peaks. In the high-resolution XPS spectrum of the Cu 2p core level, the 2p3/2 and 2p1/2 signals are, respectively, located at 931.6 eV and 951.1 eV, revealing that the binding energies are well-matched with that of Cu$^+$ according to previous reports (Tang et al., 2010). The XPS peaks of Cd 3d is centered at the binding energies of 404.5 eV (Cd 3d5/2) and 411.2 eV (Cd 3d3/2), which are in agreement with the Cd 3d core levels from Cds (Zhu et al., 2014). The featured peaks of S 2p were located at the binding energies of 160.8 eV and 161.9 eV, which indicated

![HRTEM images of CuCdS nanostructures prepared at 120°C with reaction time of 10 min using different Cu/Cd molar ratios: (a) 1:1, (b) 2:1, (c) 4:1, and (d) 8:1.](image-url)
the S 2p binding energy with $-2$ oxidation state on the basis of previous reports (Li et al., 2014b).

The current–voltage ($I$–$V$) characteristics of four different shaped CuCdS nanocrystals which were synthesized with different Cu:Cd molar ratios are presented in Figure 7. The bottom-right inset of Figure 7 depicts a typical schematic structure of devices, containing ITO anode, CuCdS nanocrystals film, and Al cathode. To measure these devices, we used copper conducting resin to form the contact pads between the probes and the Al electrodes and obtained rectifying $I$–$V$ curves. $I$–$V$ curves of the device were measured for a 5 V bias range at room temperature in air environment. We can gain four asymmetric rectifying $I$–$V$ curves, which running through the positive and the negative bias voltages. For these four different shaped samples, the values of the current show a gradually increasing trend with increasing the molar ratio of Cu:Cd from 1:1 to
8:1. The non-linearity of current illustrates that the nanocrystals formed Schottky junction with Al. Additionally, the change of conductivity may be related to the decrease in the percentage of CdS semiconductor. As a high resistivity semiconductor, CdS may reduce the current values on account of the high series resistance in the diodes. These rectification characteristics implied that the CuCdS alloy nanostructures would have potential occupancy in photovoltaic devices.

**CONCLUSION**

In summary, we report the high-quality CuCdS alloy nanocrystals with tunable compositions and various shapes synthesized by using CuI and Cd(DDTC)2 as precursors. The shapes of this product are controlled by reaction temperatures.
(100–150°C) and Cu: Cd precursor molar ratios, which includes fusiform, rod, hexagon, rhombus and so on. The CuCdS alloy nanocrystals exhibit tunable LSPR absorption spectra dependent on the free carrier density, high frequency dielectric constant, and crystalline anisotropy. In addition, the CuCdS nanocrystals reveal certain rectification characteristics. These interesting properties make the CuCdS alloy nanocrystals applicable in photovoltaic and photoelectric researches.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Materials, further inquiries can be directed to the corresponding author/s.

**AUTHOR CONTRIBUTIONS**

YG conducted the experiments and wrote the results. LW wrote the manuscript. GT, SZ, and HW characterized the samples. JN and LL revised the manuscript. All authors contributed to the article and approved the submitted version.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2020.628536/full#supplementary-material

**REFERENCES**

Bai, X., Finn, P.-M., and Gun’ko, Y. K. (2020). Near-infrared-emitting CIZSe/CIZS/ZnS colloidal heteronanomaterial structures. *Nanoscale* 12, 15295–15303. doi: 10.1039/D0NR02777D

Bera, S., Shyamal, S., Sen, S., and Pradhan, N. (2020). Insights of crystal growth, nucleation density, and shape modulations in the formation of I-VI ternary semiconductor nanoplatelet photoelectrocatalysts. *J. Phys. Chem. C* 124, 15607–15615. doi: 10.1021/acs.jpcc.0c03947

Devaney, W. E., Barnett, A. M., Storti, G. M., and Meakin, J. D. (1979). The design and fabrication of CdS/CuS Cells of 8.5-percent conversion efficiency. *IEEE, Trans. Electron Devices* 26, 205–210. doi: 10.1109/TE-D.1979.19405

Feng, H., Song, J., Song, B., Lin, Q., Shen, H., Li, L. S., et al. (2020). Highly efficient near-infrared light-emitting diodes based on chloride treated CdTe/CdSe type-II quantum dots. *Front. Chem.* 8:266. doi: 10.3389/fchem.2020.00266

Han, S.-K., Gong, M., Yao, H.-B., Wang, Z.-M., and Yu, S.-H. (2012). One-pot controlled synthesis of hexagonal-prismatic CuInS2-ZnS-CuInS2-CuS nanocrystals and CuInS2-ZnS-CuInS2-ZnS-CuInS2 heteronanostuctures. *Angew. Chem. Int. Ed.* 51, 6365–6368. doi: 10.1002/anie.201202128

Han, W., Yi, L., Zhao, N., Tang, A., Gao, M., and Tang, Z. (2008). Synthesis and shape-tailoring of copper sulfide/indium sulfide-based nanocrystals. *J. Am. Chem. Soc.* 130, 13152–13161. doi: 10.1021/ja8046393

Jun, S., and Jang, E. (2012). Bright and stable alloy core/multishell quantum dots. *Angew. Chem. Int. Ed.* 52, 679–682. doi: 10.1002/anie.201206333

Kim, I., Agrawal, A., Krieg, F., Bergerud, A., and Milliron, D. J. (2016). The interplay of shape and crystalline anisotropies in plasmonic semiconductor nanocrystals. *Nano Lett.* 16, 3879–3884. doi: 10.1021/acs.nanolett.6b01390

Li, H., Brescia, R., Povia, M., Prato, M., Bertoni, G., Manna, L., et al. (2013). Synthesis of uniform disk-shaped copper telluride nanocrystals and cation exchange to cadmium telluride quantum disks with stable red emission. *J. Am. Chem. Soc.* 135, 12270–12278. doi: 10.1021/ja40494k

Li, H., Jiang, X., Wang, A., and Du, Z. (2020). Simple synthesis of CuInS2/ZnS core/shell quantum dots for white light-emitting diodes. *Front. Chem.* 8:669. doi: 10.3389/fchem.2020.00669

Li, J., Bloemen, M., Parsi, J., and Kolny-Olesiak, J. (2014a). Role of copper sulfide seeds in the growth process of CuInS2 nanorods and networks. *ACS Appl. Mater. Interfaces* 6, 20535–20543. doi: 10.1021/am506145Y

Li, J., Shen, H., Zhou, C., Li, N., Wang, H., and Li, L. (2014b). Inorganic Sn–X complex ligands capped CuInS2 nanocrystals with high electron mobility. *J. Nanopart. Res.* 16:2802. doi: 10.1007/s11051-014-2802-7

Liu, Y., Liu, M., and Swihart, M. T. (2017). Plasmonic copper sulfide-based materials: a brief introduction to their synthesis, doping, alloying, and applications. *J. Phys. Chem. C* 121, 13435–13447. doi: 10.1021/acs.jpcc.7b06894

Meng, X., Liu, Z., Cao, Y., Dai, W., Zhang, K., Dong, H., et al. (2017). Fabricating aptamer-conjugated pegylated-MoS2/CuInS2 theranostic nanoplatform for multiplexed imaging diagnosis and chemo-photothermal therapy of cancer. *Adv. Funct. Mater.* 27:160592. doi: 10.1002/adfm.201605592

Pan, C., Niu, S., Ding, Y., Dong, L., Yu, R., Liu, Y., et al. (2012). Enhanced Cu2S/CdS coaxial nanowire solar cells by piezo-phototronic effect. *Nano Lett.* 12, 3302–3307. doi: 10.1021/nl304082

Pantani, M. G., Stolle, C. J., Reid, D. K., Rhee, D. J., Harvey, T. B., Akhavan, V. A., et al. (2013). CuInSe2 quantum dot solar cells with high open-circuit voltage. *J. Phys. Chem. Lett.* 4, 2030–2034. doi: 10.1021/jl4010015

Pejova, B., Sherif, E., and Minde, M. W. (2020). Sonochemically synthesized quantum nanocrystals of cubic CuInS2: evidence for multilatrf surfapce morphology, size-dependent structure, and particle size distribution. *J. Phys. Chem. C* 124, 20240–20255. doi: 10.1021/acs.jpcc.0c06070

Regulacio, M. D., Ye, C., Lim, S. H., Bosman, M., Polavarapu, L., Koh, W. L., et al. (2011). One-pot synthesis of CuInS2-CdS and CuInS2-ZnS-CdS nanodisk heterostructures. *J. Am. Chem. Soc.* 133, 2052–2055. doi: 10.1021/ja1090589

Rivest, J. F., Swisher, S. L., Fong, L., Zheng, H., and Alivisatos, A. P. (2011). Assembled monolayer nanorod heterojunctions. *ACS Nano* 5, 3811–3816. doi: 10.1021/nn2001454

Shen, H., Niu, J., Li, X., Wang, H., Xing, M., Chen, X., et al. (2012). Fabrication of “strong” columnar Cu2Se superstructures assisted by inorganic ligands. *Nanoscale* 4, 2741–2747. doi: 10.1039/c2nr00057a

Shen, S., Zhang, Y., Liu, Y., Peng, L., Chen, X., and Wang, Q. (2012). Manganese-Doped Ag2S-ZnS Heteronanostuctures. *Chem. Mater.* 24, 2407–2413. doi: 10.1021/cm301342z

Shen, S., Zhang, Y., Peng, L., Du, Y., and Wang, Q. (2011a). Matchstick-shaped Ag2S-ZnS heteronanostuctures preserving both uv/blue and near-infrared photoluminescence. *Angew. Chem. Int. Ed.* 2011, 50, 7115–7118. doi: 10.1002/anie.201101084

Shen, S., Zhang, Y., Peng, L., Xu, B., Du, Y., Deng, M., Xu, H., and Wang, Q. (2011b). Generalized synthesis of metal sulfide nanocrystals from single-source precursors: size, shape and chemical composition control and their properties. *CrystEngcomm* 13, 4572–4579. doi: 10.1039/c0ce00982b

Swafford, L. A., Weigand, L. A., Bowers, M. J., McBride, J. R., Rappaport, J. L., Watt, T. L., et al. (2006). Homogeneously alloyed CdS/Se$_x$ nanocrystals: synthesis, characterization, and composition/size-dependent band gap. *J. Am. Chem. Soc.* 128, 12299–12306. doi: 10.1021/ja063939e
Tang, A., Yi, L., Han, W., Teng, F., Wang, Y., Hou, Y., et al. (2010). Synthesis, optical properties, and superlattice structure of Cu(I)-doped CdS nanocrystals. *Appl. Phys. Lett.* 97:033112. doi: 10.1063/1.3466664

Wang, J., Feng, H., Chen, K., Fan, W., and Yang, Q. (2014). Solution-phase catalytic synthesis, characterization and growth kinetics of Ag$_2$S–CdS matchstick-like heteronanostructures. *Dalton Trans.* 43, 3990–3998. doi: 10.1039/C3DT52693C

Xu, Q., Li, X., Lin, Q., Shen, H., Wang, H., and Du, Z. (2020). Improved efficiency of all-inorganic quantum-dot light-emitting diodes via interface engineering. *Front. Chem.* 8:265. doi: 10.3389/fchem.2020.00265

Xu, W., Niu, J., Wang, H., Shen, H., and Li, L. S. (2013). Size, Shape-dependent growth of semiconductor heterostructures mediated by Ag$_2$Se nanocrystals as seeds. *ACS Appl. Mater. Interfaces* 5, 7537–7543. doi: 10.1021/am4018303

Yi, L., Tang, A., Niu, M., Han, W., Hou, Y., and Gao, M. (2010). Synthesis and self-assembly of Cu$_{1.94}$S–ZnS heterostructured nanorods. *CrystEngComm* 12, 4124–4130. doi: 10.1039/c0ce00026d

Yoon, S.-Y., Kim, J.-H., Jang, E.-P., Lee, S.-H., Jo, D.-Y., Kim, Y., et al. (2019). Systematic and extensive emission tuning of highly efficient Cu-In-S-based quantum dots from visible to near infrared. *Chem. Mat.* 31, 2627–2634. doi: 10.1021/acs.chemmater.9b00550

Zhang, J., Yu, J., Zhang, Y., Li, Q., and Gong, J. (2011). Visible light photocatalytic h$_2$-production activity of CuS/ZnS porous nanosheets based on photoinduced interfacial charge transfer. *Nano Lett.* 11, 4774–4779. doi: 10.1021/nl202587b

Zhao, M., Huang, F., Lin, H., Zhou, J., Xu, J., Wu, Q., et al. (2016). CuGa$_x$Zn$_{1-x}$S p-n nanoheterostructures: a promising visible light photocatalyst for water-splitting hydrogen production. *Nanoscale* 8, 16670–16676. doi: 10.1039/C6NR05002F

Zhou, D., Liu, D., Xu, W., Yin, Z., Chen, X., Zhou, P., et al. (2016). Observation of considerable upconversion enhancement induced by Cu$_{2−x}$S plasmon nanoparticles. *ACS Nano* 10, 5169–5179. doi: 10.1021/acsnano.6b00649

Zhu, D., Tang, A., Lan, P., Liu, Z., Yang, C., and Teng, F. (2016). Tuning the plasmonic resonance of Cu$_{2−x}$S nanocrystals: effects of the crystal phase, morphology and surface ligands. *J. Mater. Chem. C* 4, 4880–4888. doi: 10.1039/C6TC00980H

Zhu, G., Bao, C., Liu, Y., Shen, X., Xi, C., Xu, Z., et al. (2014). Self-regulated route to ternary hybrid nanocrystals of Ag–Ag$_2$S–CdS with near-infrared photoluminescence and enhanced photothermal conversion. *Nanoscale* 6, 11147–11156. doi: 10.1039/C4NR03001J

Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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