



 Cite this: *RSC Adv.*, 2021, 11, 3673

Controllable synthesis of non-layered two-dimensional plate-like CuGaSe₂ materials for optoelectronic devices†

 Wenling Feng, Yutong Zhao, Di Zhao, Wenjian Wang, Zenghao Xia, Xiaoxia Zheng, Xu Wang, Weihua Wang and Wenliang Wang *

CuGaSe₂ semiconductor materials, as an important member of the I–III–VI₂ family, have sparked tremendous attention due to their fascinating structure-related properties and promising applications in solar energy storage and conversion. Nevertheless, the controllable preparation of two-dimensional (2D) CuGaSe₂ structures is still a daunting challenge owing to the intrinsic non-layered crystal structure and inaccessible reactivity-matching of multiple reaction precursors, which will seriously impede the much deeper research progress on their properties and applications. Herein, non-layered 2D CuGaSe₂ plates possessing high crystallinity, and uniform size and morphology have been first synthesized by a feasible cation exchange strategy. Because the fabrication of 2D CuGaSe₂ crystals is rarely reported, a particular highlight is laid on the compositional analysis, structural characterization, and formation mechanism. Furthermore, the optical absorption and optoelectronic measurements reveal that the as-synthesized CuGaSe₂ plates exhibit high light harvesting capacity and excellent photoelectric performance. This study opens up a new avenue for the feasible fabrication of non-layered CuGaSe₂ plates possessing a high-quality crystalline structure and provides a promising candidate for the development of novel solar energy conversion and storage devices.

 Received 11th October 2020
 Accepted 11th January 2021

DOI: 10.1039/d0ra08662b

rsc.li/rsc-advances

1. Introduction

Since the first successful synthesis of graphene in 2004,¹ two-dimensional (2D) materials have aroused the great interest and enthusiasm of researchers owing to their high specific surface area, unique electronic structure and excellent physical and chemical properties.^{2–4} Generally, 2D materials can be divided into layered and non-layered types according to their crystal structures. Despite the fact that previous work mainly focused on 2D materials with a layered crystal structure, the research on 2D non-layered materials has gradually attracted extensive attention.^{5–7} Importantly, the emergence and development of 2D non-layered semiconductor structures plays a vital role in broadening and deepening the fundamental investigation on 2D materials, which also exhibit intriguing structure- and composition-dependent physicochemical properties and advanced applications.^{8–10} Compared with layered materials, non-layered 2D crystals feature abundant dangling bonds on the surface and intrinsic crystal distortion that render them specially promising applications in solar energy

conversion and storage, such as photovoltaic cells, optoelectronic detection, and photocatalytic fields.^{6,11–13} Solar energy, featuring its universality, inexhaustibility, environmental benignity and high capacity, is identified as the most promising option for renewable energy. Accordingly, developing advanced materials to efficiently convert solar energy into chemical and/or electronic energy is attracting great research interest. In particular, 2D semiconductor materials originating from their unique physical, electronic, and structural properties have been intensively investigated for application in electronic and optoelectronic devices.⁵ To date, considerable efforts have been made to achieve the anisotropic growth on non-layered materials.⁶ However, the most studies of these 2D structures have largely limited to binary compounds, such as CdX (X = S, Se, Te)^{14–16} and PbX (X = S, Se).¹⁷ In addition, further applications of these materials are severely restricted by the inherent toxicity of Cd and Pb. Strikingly, as the potential alternatives to the CdX or PbX crystals, copper-based chalcogenide compounds have spawned growing interest and research owing to their promising applications in optoelectronic devices.^{18–20} Therefore, rational design a facile approach for the controllable fabrication of 2D non-layered semiconductor materials with appropriate bandgap energy, low toxicity and high light absorption coefficient is of great significance for improving conversion of solar energy into chemical and/or electronic energy.

School of Chemistry and Chemical Engineering, Qufu Normal University, Qufu 273165, Shandong, P. R. China. E-mail: wliwang@qfnu.edu.cn; Tel: +86-1565-023-5536

† Electronic supplementary information (ESI) available: SEM and HRTEM of the Cu_{2–x}Se plates. SEM, EDX spectrum, ICP-AES, UV-vis-NIR absorption spectra, and TGA of the as-synthesized CuGaSe₂ plates. See DOI: 10.1039/d0ra08662b



Currently, accumulating evidences demonstrate that copper-based ternary I-III-VI₂ (I = Cu; III = Ga, In; VI = S, Se, Te) chalcogenide semiconductor materials have received a great deal of attention due to their distinct composition and structure-tunable properties and extensive applications.^{21–31} For instance, Wang *et al.* reported a hot injection method for the preparation of monodispersed wurtzite structure CuInSe₂ nanocrystals with uniform hexagonal shape as well as their high performance in optoelectronic application.³² Xia *et al.* demonstrated a solvothermal synthesis technique to prepare CuInS₂ quantum dots and studied their size-dependent optical properties.³³ Among these I-III-VI₂ materials, CuGaSe₂ as a typical p-type semiconductor possesses an appropriate bandgap and high light absorption coefficient, which makes it as an ideal effective light-absorbing material in optoelectronic and photovoltaic applications.^{18,34–36} Over the past several years, tremendous endeavors have been devoted to develop synthetic strategies for controllable preparation of CuGaSe₂ materials, and significant advances have been achieved. For example, Tang *et al.* prepared CuGaSe₂ nanocrystals using a hot injection method.³⁴ Feng *et al.* developed a solid-state reaction to synthesize CuGaSe₂ nanocrystals.¹⁸ Nevertheless, the morphology of the as-prepared CuGaSe₂ materials is mostly restricted in the form of nanoparticles, which may heavily hamper their potential applications in optoelectronic devices. In sharp contrast to the synthesis of binary chalcogenides, there is still a big challenge to control the composition, shape, and crystal phase of ternary nanocrystals due to the difficult to balance the reactivity of three different precursors to favor the formation of ternary compound and restrain binary by-products.^{29,37} To date, the controllable preparation of 2D CuGaSe₂ crystals through wet-chemical methods is still in its infancy. Especially, tetragonal phase 2D CuGaSe₂ structures with high crystalline quality and uniform size have rarely been reported.¹⁸ Consequently, exploring and developing a feasible synthesis method for controlled preparation of tetragonal phase 2D CuGaSe₂ materials is urgently needed, which will further deepen the understanding of the growth mechanism and promote comprehensive applications of 2D non-layered materials.

Cation exchange reactions, in which involve replacing cations of a parent crystalline structure with diverse metal ions.^{38,39} Cation exchange method has been demonstrated to be an extremely efficient strategy in the colloid synthesis of semiconductor materials with unique composition, morphology, and crystal structure that are difficult to be achieved by simple direct growth methods.^{40–44} For example, Berends *et al.* synthesized CuInSe₂ quantum dots, nanoplatelets, and nanosheets by partial indium-cation exchange into Cu_{2–x}Se nanocrystals.⁴⁵ Wang *et al.* reported the successful conversion of layered SnS₂ into nonlayered 2D Cu₂SnS₃ nanosheets *via* cation exchange.⁴⁶ Ramirez *et al.* demonstrated a morphology transformation from Cu_{2–x}Se nanoparticles to Cu₃SbSe₃ nanoplates by cation exchange reactions.²⁰ Such convincing evidences indicate that cation exchange reaction may offer an effective approach for the preparation of tetragonal phase 2D CuGaSe₂ structures.

In this work, we demonstrate for the first time a partial cation exchange strategy to prepare tetragonal phase 2D CuGaSe₂ plates. Firstly, Cu_{2–x}Se plates are synthesized *via* a hot-injection method. Subsequently, Ga³⁺ ions are incorporated into the Cu_{2–x}Se crystal lattice *via* the partial replacement of Cu⁺ by Ga³⁺. The shape and size of the parent Cu_{2–x}Se without obvious change is retained in resulting CuGaSe₂ crystals. The phase purity of the synthesized CuGaSe₂ plates can be regulated by varying the initial precursor Ga : Cu ratio. On the basis of systematic analysis by several characterization techniques, a possible formation mechanism of CuGaSe₂ plates has been proposed. Moreover, optoelectronic measurements confirm that a photodetector based on the CuGaSe₂ plates exhibits excellent sensitivity and stability.

2. Experimental section

2.1 Materials

Gallium (iii) acetylacetonate (Ga(acac)₃, 99.99%) and diphenyl diselenide (Ph₂Se₂, 98%) were purchased from Alfa Aesar. Octadecene (ODE, 90%) and oleylamine (OAm, 70%) were ordered from Sigma-Aldrich. Oleic acid (OA), anhydrous copper chloride (CuCl), absolute ethanol, and toluene were purchased from Sinopharm Chemical Reagent Ltd., China. All chemicals were used directly without further purification.

2.2 Preparation of Se–ODE and Ga–OAM precursors

The Se–ODE precursor solution was prepared by addition of 0.0624 g of Ph₂Se₂ (0.2 mmol) to 1.0 mL of ODE, followed by heating to 70 °C, yielding a transparent orange liquid.⁴⁷ The Ga–OAM precursor solution was prepared in a nitrogen atmosphere, by dissolving 0.3670 g of Ga(acac)₃ (1.0 mmol) in 10.0 mL OAm in a three-neck flask under magnetic stirring, followed by heating to 120 °C for 60 min until the Ga(acac)₃ absolutely dissolved.⁴⁸

2.3 Synthesis of the two-dimensional CuGaSe₂ plates

CuGaSe₂ plates were prepared according to our previously reported method with a slight modification.⁴⁹ In a typical synthesis procedure, 0.0099 g of CuCl (0.10 mmol) together with 0.2 mL of OAm, 0.2 mL of OA and 6.0 mL of ODE were brought into a three-neck round-bottom flask (100 mL) at room temperature. In order to eliminate dissolved oxygen, adventitious water and other low-boiling-point impurities, the mixture was first heated to 130 °C and maintained for 30 min under nitrogen atmosphere and constant stirring. After that, the reaction mixture was further heated to 230 °C. At this point, the Se–ODE precursor solution was transferred into a syringe equipped with a large needle and injected quickly into the flask at 230 °C and kept at the temperature for 10 min. Subsequently, 2.0 mL Ga–OAM precursor solution was immediately injected into the above solution and the reaction mixture was further raised to 255 °C and maintained for 90 min with continuous stirring. After the reaction was complete, the resulting solution was allowed to cool to room temperature naturally. The as-synthesized product was collected by centrifugation

(9000 rpm, 3 min) and washed several times with absolute ethanol and toluene.

2.4 Characterization

The as-prepared samples were characterized by X-ray diffraction (XRD) performed on a Philips X'pert PRO X-ray diffractometer (Cu $K\alpha$, $\lambda = 1.54182 \text{ \AA}$). The morphologies of the samples were examined using a Scanning Electron Microscopy (SEM, JSM-6700F). The high resolution transmission electron microscopy (HRTEM), selected area electron diffraction (SAED) patterns, high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM), and corresponding energy-dispersive X-ray spectroscopy (EDX) mapping analyses were performed on a JEOL JEM-ARF200F TEM/STEM with a spherical aberration corrector. Inductively Coupled Plasma Atomic Emission Spectroscopy (ICP-AES) analysis was performed on a PerkinElmer Model Optima 3000DV to quantify the copper, gallium, and selenium content of the resulting crystals. X-ray Photoelectron Spectroscopy (XPS) was performed using an ESCALAB MK II with Mg $K\alpha$ as the excitation source. The ultraviolet-visible-near infrared (UV-vis-NIR) absorption spectra were recorded on a spectrophotometer (Shimadzu Solidspec-3700DUV) at room temperature. The samples were prepared by dispersing the plates in toluene. Thermogravimetric analysis (TGA) was performed on a STA-449-F3 thermal analyzer (NETZSCH Corporation) with a heating rate of $10 \text{ }^\circ\text{C min}^{-1}$ in flowing argon. Current–voltage (I – V) characteristics of the devices were recorded with a two-probe method using an electrochemical station (CHI660e) and xenon lamp (PLS-SXE300) was selected as a white light source.

3. Results and discussion

3.1 Characterizations of the CuGaSe₂ plates

The typical CuGaSe₂ plates have been synthesized by means of a cation exchange method. The method consists of a sequential two-step strategy, *i.e.*, starting from the as-synthesized Cu_{2–x}Se plates followed by the incorporation of Ga³⁺ *via* an *in situ* cation exchange reaction, leading to homogeneous CuGaSe₂ plates. The structure, morphology, and composition of the as-obtained CuGaSe₂ plates are comprehensively characterized by various techniques. As shown in Fig. 1a, the crystal structure of the as-prepared CuGaSe₂ plates is characterized by the powder XRD. The major specific diffraction peaks at 27.7° , 45.7° , 46.1° , 54.2° , and 55.0° can be respectively indexed to the (112), (220), (204), (312), and (116) planes of the tetragonal chalcopyrite phase of CuGaSe₂ (JCPDS no. 31-0456).³⁶ The none-existence of any observable impurity peak and peak splitting can confirm the absence of the chance of the phase separation during the synthesis of CuGaSe₂ plates. Meanwhile, the presence of strong and sharp diffraction peaks verifies the highly crystalline nature of the resulting CuGaSe₂ plates. The morphology of CuGaSe₂ crystals is examined by SEM measurement. As displayed in Fig. 1b, it can be seen that the as-synthesized CuGaSe₂ crystals are roughly hexagonal structure. The thickness and edge length of the CuGaSe₂ plates are around $45.3 \pm 2.3 \text{ nm}$ and $328.1 \pm$

17.6 nm (Fig. S1 of the ESI†), respectively. HRTEM, SAED, and EDX are performed to investigate the crystal microstructure and composition of the resultant CuGaSe₂ plates. As shown in Fig. 1c, the lattice fringes can be clearly observed, uncovering the well-defined single-crystalline structure. Moreover, the detected interplanar distances of the CuGaSe₂ plates are 0.32 and 0.20 nm, which can be assigned to the (112) and (2–20) lattice planes of tetragonal phase CuGaSe₂,^{18,50} respectively. For the corresponding SAED pattern depicted in Fig. 1d, the observed regular symmetric diffractions can also demonstrate that the as-prepared CuGaSe₂ plates possess good crystallinity. In addition, the elemental composition of CuGaSe₂ plates is determined by EDX. As detected in Fig. S2,† the corresponding Cu : Ga : Se atomic ratio is close to 1 : 1 : 2, which is consistent with the ICP-AES analysis (Table S1 of the ESI†). All of the above measurement results confirm that as-prepared CuGaSe₂ plates are of high purity and good crystalline structure.

In order to further verify the structure and composition of the CuGaSe₂ plates, STEM-EDX analysis was conducted. As shown in Fig. 2a, STEM image demonstrates a good consistent with the SEM observation, in which the prepared CuGaSe₂ plates are of approximately hexagonal morphology. As shown in Fig. S2 of the ESI,† the EDX measurement confirms that the resulting products are composed of Cu, Ga, and Se elements. The atomic ratio of Cu : Ga : Se is close to 1 : 1 : 2, which is well consistent with the stoichiometric ratio of CuGaSe₂. In addition, as shown in Fig. 2b–d, STEM-EDX elemental mapping of the as-synthesized CuGaSe₂ plates further reveals that three elements are homogeneously distributed in the plates.

X-ray photoelectron spectroscopy (XPS) has been performed to investigate the chemical state and elemental composition of the synthesized CuGaSe₂ plates. As illustrated in Fig. 3a, the XPS survey spectrum uncovers the existence of Cu, Ga, Se, C, and O in the resulting CuGaSe₂ plates. As shown in Fig. 3b the copper high-resolution XPS spectrum, two characteristic peaks of Cu 2p_{3/2} and Cu 2p_{1/2} locating at 931.5 and 951.3 eV with a peak splitting of 19.8 eV demonstrate the presence of Cu(I) in the CuGaSe₂ plates.⁵¹ The gallium high-resolution XPS spectrum in Fig. 3c exhibits two peaks at 1143.6 and 1116.7 eV corresponding to Ga 2p_{1/2} and Ga 2p_{3/2},^{52,53} respectively, suggesting that the chemical state of the Ga in the composite is +3. In Fig. 3d the high-resolution XPS spectrum of Se 3d, the two peaks of 53.2 and 54.1 eV are assigned to Se 3d_{5/2} and Se 3d_{3/2}, respectively, indicating the –2 oxidation state of Se in the as-prepared CuGaSe₂ plates.⁵⁴ Consequently, XPS results further verify that CuGaSe₂ plates without any other impurities have been successfully prepared.

3.2 Formation mechanism of the CuGaSe₂ plates

To gain a further insight into the formation mechanism of the as-prepared CuGaSe₂ plates, the products of different reaction stages before and after gallium precursor injection have been carefully investigated. Firstly, we will analyze the results of reaction system before injecting gallium precursor. In the first step, the crystal structure, chemical composition, and morphology of the resulting crystals after injecting selenium

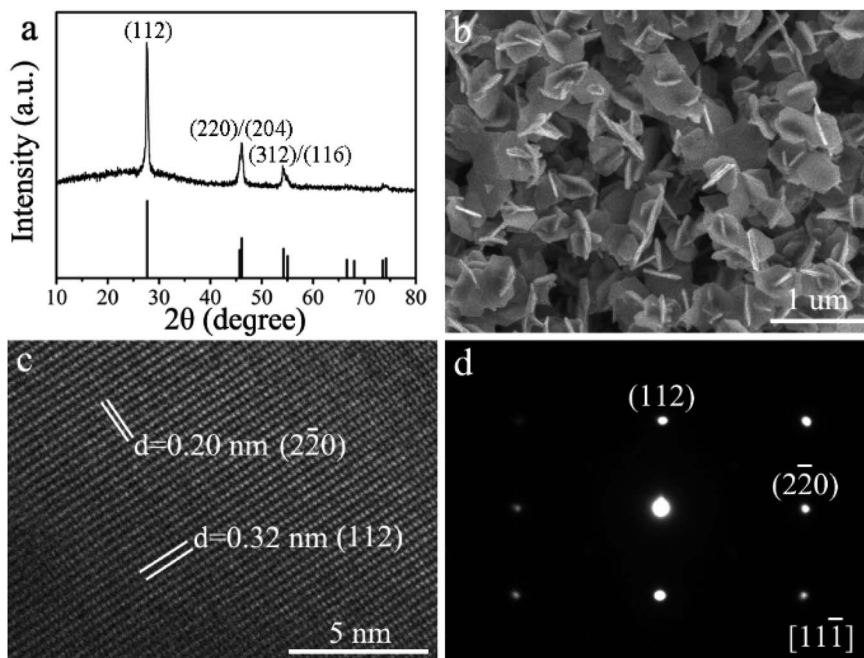


Fig. 1 (a) XRD pattern along with standard JCPDS no. 31-0456 for reference and (b) SEM image of the as-synthesized CuGaSe_2 plates. (c) HRTEM image and (d) the corresponding SAED pattern of the CuGaSe_2 plates.

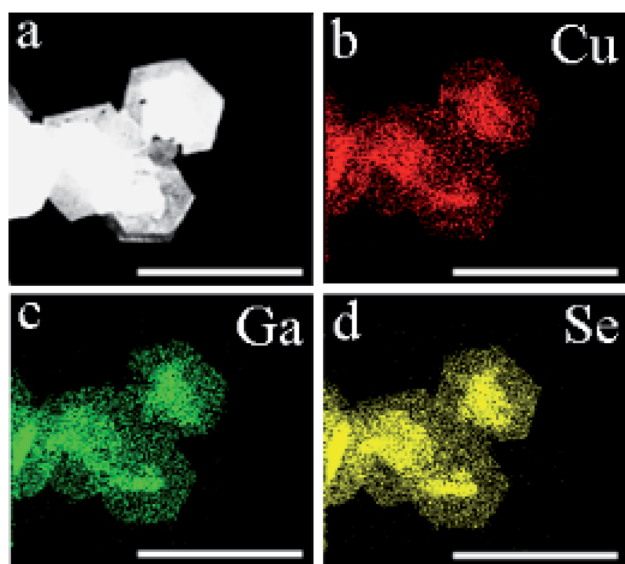


Fig. 2 (a) HAADF-STEM image of the as-prepared CuGaSe_2 plates and the corresponding STEM-EDX elemental mappings for (b) Cu, (c) Ga and (d) Se, respectively. The scale bar is 1 μm .

precursor at 230 °C for 10 min (Experimental section for details) are characterized by several techniques. The SEM image in Fig. 4 depicts the synthesized crystals are of fairly uniform plates. The thickness and edge length of the plates are around 43.8 ± 2.9 nm and 320.5 ± 15.8 nm (Fig. S3[†]), respectively. As seen in Fig. S4,[†] the interplanar distance of 0.33 nm can be well indexed to the (111) lattice space of the cubic Cu_{2-x}Se .⁵⁵ The crystal phase of the resulting product is characterized by XRD.

As shown in Fig. 5a, the emerged diffraction peaks at 27.0°, 31.3°, 44.8°, 53.1°, 65.2°, and 71.9° can be assigned to the (111), (200), (220), (311), (400), and (331) planes of the cubic phase of Cu_{2-x}Se (JCPDS no. 06-0680),⁵⁵ respectively. Furthermore, EDX analysis in Fig. 5b demonstrates that as-prepared Cu_{2-x}Se crystals have Cu deficient stoichiometry with the Cu : Se ratio of 1.85 : 1. Strikingly, slightly Cu-deficient composition of Cu_{2-x}Se crystals is favorable for cation exchange reactions.⁵⁶ XPS analysis is conducted to probe surface chemistry of prepared Cu_{2-x}Se plates. For the high-resolution XPS spectrum of Cu 2p in Fig. 5c, the peaks at 932.0 ($2p_{3/2}$) and 951.8 eV ($2p_{1/2}$) demonstrate the presence of Cu^+ .⁵⁷ Fig. 5d displays the high-resolution XPS spectrum of Se 3d, where the peaks at 53.8 ($3d_{5/2}$) and 54.3 ($3d_{3/2}$) eV indicate the (−2) oxidation state of Se.⁵⁸ Overall, these experimental results can verify the formation of cubic Cu_{2-x}Se plates prior to the injection of gallium precursor.

The synthesized Cu_{2-x}Se plates remained in the crude solution for the following cation exchange reaction toward CuGaSe_2 plates. Generally, high temperature is favorable to impel the proceeding of cation exchange reactions for the synthesis of Ga-containing nanomaterials.²³ In addition, it is worth noting that molar ratio of precursors is judiciously controlled since it has a significant effect on the phase purity of the resulting product. We raise the reaction temperature to 255 °C after injecting gallium precursor. The XRD characterizations have been employed to monitor the phase evolution of the resulting products obtained from various Ga : Cu precursor molar ratios during the process of cation exchange reactions. As shown in Fig. 6a and b, a group of XRD patterns reveal that the initial cubic phase Cu_{2-x}Se has partially transformed into

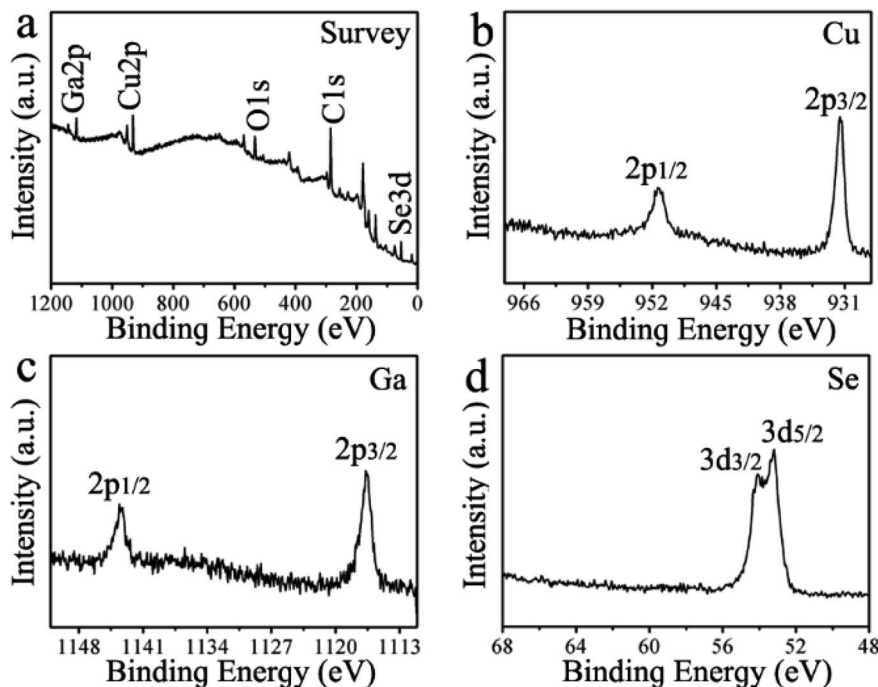


Fig. 3 XPS spectra of the as-prepared CuGaSe₂ plates. (a) Survey spectrum, (b) Cu 2p core level spectrum, (c) Ga 2p core level spectrum, and (d) Se 3d core level spectrum.

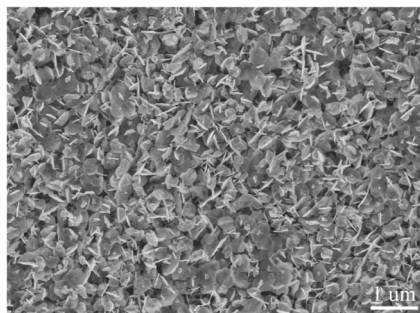


Fig. 4 SEM image of the sample synthesized at 230 °C for 10 min after Se-precursor injection.

tetragonal phase CuGaSe₂ when the Ga : Cu ratio changes from 0 : 1 to 1 : 1. Further increase of the Ga : Cu ratio up to 1.5 : 1 in Fig. 6c results in the enhancement of the representative diffraction peaks at 27.7°, 46.1°, and 54.2°, which can be indexed to the (112), (220)/(204), and (312)/(116) planes of the tetragonal chalcopyrite phase of CuGaSe₂. Meanwhile the fading of the main characteristic peaks at 27.0°, 44.8°, and 53.1° matched with the (111), (220), and (311) planes of cubic Cu_{2-x}Se. Finally, as present in Fig. 6d, a pure tetragonal structure of CuGaSe₂ plates with nearly hexagonal shape is successfully achieved when the Ga : Cu ratio increases to 2 : 1. It is demonstrated that CuGaSe₂ plates with desired stoichiometric combination can be obtained by suitable adjustment of molar precursor ratios.

3.3 Optical and optoelectronic properties of the CuGaSe₂ plates

To prove the ability of the optoelectronic device application of the CuGaSe₂ plates, UV-vis-NIR absorption spectra are measured to investigate the optical properties. As depicted in Fig. 7a, an obvious comparison in absorption curves can be observed between Cu_{2-x}Se and CuGaSe₂ crystals. The spectral response curve of initial Cu_{2-x}Se plates exhibits two stages. For the ranges within 800 nm, they should be induced by excitonic or interband transitions. However, the range beyond 800 nm, they should be caused by localized surface plasmon resonance (LSPR). Remarkably, a distinct LSPR of Cu_{2-x}Se plates in the NIR region arising from the copper-defect-induced free carriers is thoroughly restrained after the partial replacement of Cu⁺ with Ga³⁺ *via* cation exchange reactions. Such results of UV-vis-NIR absorption demonstrate another evidence of the successful transformation from Cu_{2-x}Se to CuGaSe₂. Meanwhile, the as-synthesized CuGaSe₂ plates demonstrate a stronger absorption in visible light region. On the base of absorption spectra, the direct optical bandgap energy of CuGaSe₂ plates in Fig. 7b is determined by extrapolation of the linear region of the Tauc plot to the photon energy. As a result, the calculated bandgap is 1.61 ± 0.003 eV (Fig. S5[†]), which is in well consistent with the literature report.⁵⁹ Moreover, we have investigated the thermal stability of the resulting CuGaSe₂ plates. According to TG analysis in Fig. S6,[†] the small amount of weight loss over the temperature range from 100 to 380 °C may be attributed to the removal of absorbed water molecules and some organic compounds on the materials' surface. With the increase of the temperature, up to about 530 °C, the CuGaSe₂ plates will be

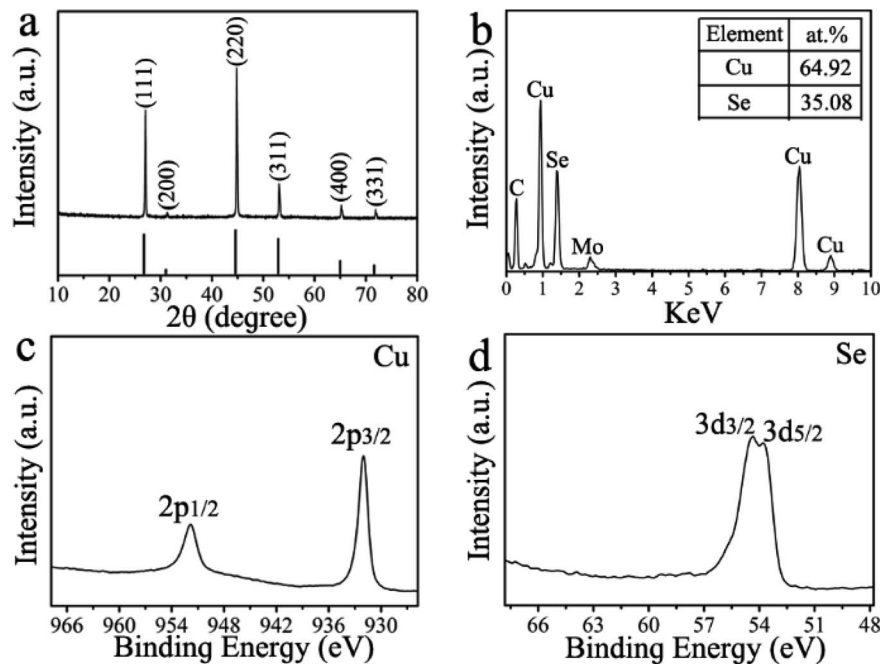


Fig. 5 (a) XRD pattern, (b) EDX spectrum, XPS spectra of (c) Cu 2p core level and (d) Se 3d core level of the product synthesized at 230 °C for 10 min after Se-precursor injection.

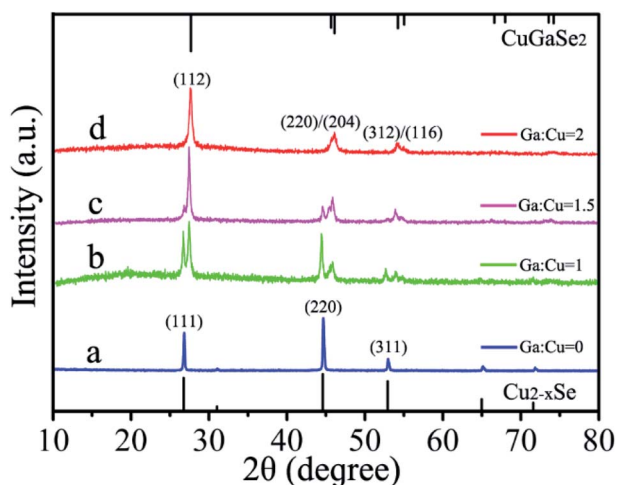


Fig. 6 XRD patterns of the samples synthesized with different precursor molar ratios: (a) Ga : Cu = 0, (b) Ga : Cu = 1, (c) Ga : Cu = 1.5, (d) Ga : Cu = 2.

further decomposed. Based on the above results of TGA, we suggest that the synthesized CuGaSe_2 plates have good thermal stability.

Based on the above results of absorption spectra, the synthesized CuGaSe_2 plates are expected as promising light-harvesting materials for optoelectronic applications. To further evaluate the potential application of CuGaSe_2 plates in optoelectronics, a photodetector is constructed by drop-casting a concentrated toluene solution of CuGaSe_2 plates onto glass substrate and employed two adjacent ITO film as conductive electrodes.⁵¹ A xenon lamp is utilized as a white light source by

equipped with a cutoff filter of 400 nm, which can afford visible light ranging from 400 to 780 nm.

Fig. 7c shows the current–voltage (I – V) curves of the CuGaSe_2 plates photodetector measured in the dark and under light illumination at a bias of 1.0 V, respectively. The linear curves demonstrate good ohmic contacts between the ITO electrodes and CuGaSe_2 plates. In addition, it is obviously observed that the current under light illumination is higher than that in the case of dark. The stability and responsivity are also identified as key factors to assess the property of a photodetector. For purpose of exploring the stability and responsivity of the CuGaSe_2 -based photodetector, the time-dependent current response has been conducted under illumination with light intensity of 6.5 mW cm^{-2} at a voltage of 1.0 V. Fig. 7d exhibits the photocurrents of the device during repetitive switching of light illumination, or on/off switching. The photocurrent increases and decreases as a response to the periodically switching on and off operation for a duration of 50 s, exhibiting high sensitivity. Furthermore, the photocurrent can still be changed by periodically turning on and off the light even though after a number of cycles. Such results indicate that the photodetector based on the CuGaSe_2 plates has an excellent stability and repeatability.

4. Conclusions

In summary, we have first demonstrated a cation exchange method to prepare non-layered tetragonal phase 2D CuGaSe_2 plates. Starting from the synthesis of Cu_{2-x}Se plates, a subsequent cation exchange with Ga^{3+} results in the formation of CuGaSe_2 plates while retaining their original shape and size.

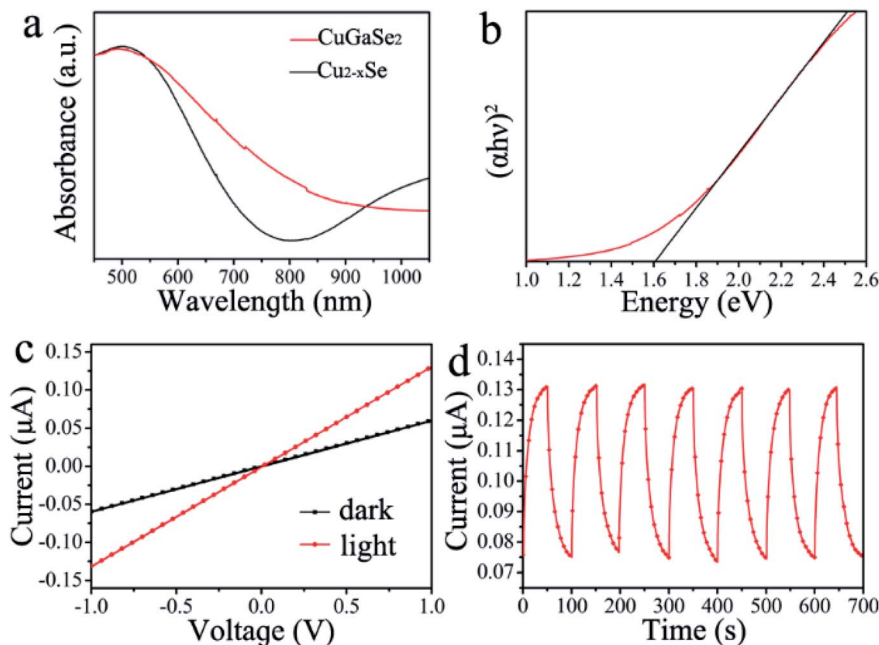


Fig. 7 (a) UV-vis-NIR absorption spectra of Cu_2-xSe and CuGaSe_2 plates dispersed in toluene. (b) The linear extrapolations of plots of $(\alpha hv)^2$ vs. photon energy. (c) Current versus voltage curves of the CuGaSe_2 plates measured in dark and under xenon lamp illumination with a power intensity of 6.5 mW cm^{-2} , and (d) current versus time curves during ON and OFF cycles.

The phase purity of CuGaSe_2 plates can be controlled by regulating the initial precursor ratio. Furthermore, we also reveal that CuGaSe_2 plates have promising potential application in photoelectric conversion. Expectedly, the present synthetic methodology of CuGaSe_2 plates can also be extended for the preparation of other 2D materials from non-layered structures to further explore the physicochemical properties and advanced applications.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (21805163), the National College Students Innovation and Entrepreneurship Training Program (201910446030), and the Qufu Normal University Innovation and Entrepreneurship Training Program for College Students (2019A004).

Notes and references

- 1 K. S. Novoselov, A. K. Geim, S. V. Morozov, D. Jiang, Y. Zhang, S. V. Dubonos, I. V. Grigorieva and A. A. Firsov, *Science*, 2004, **306**, 666–669.
- 2 X. L. Li and L. J. Zhi, *Chem. Soc. Rev.*, 2018, **47**, 3189–3216.
- 3 Y. Chen, Z. X. Fan, Z. C. Zhang, W. X. Niu, C. L. Li, N. L. Yang, B. Chen and H. Zhang, *Chem. Rev.*, 2018, **118**, 6409–6455.

- 4 Y. Liu, N. O. Weiss, X. D. Duan, H. C. Cheng, Y. Huang and X. F. Duan, *Nat. Rev. Mater.*, 2016, **1**, 16042.
- 5 P. C. Tao, S. Yao, F. Y. Liu, B. Wang, F. Huang and M. Y. Wang, *J. Mater. Chem. A*, 2019, **7**, 23512–23536.
- 6 F. Wang, Z. X. Wang, T. A. Shifa, Y. Wen, F. M. Wang, X. Y. Zhan, Q. S. Wang, K. Xu, Y. Huang, L. Yin, C. Jiang and J. He, *Adv. Funct. Mater.*, 2017, **27**, 1603254.
- 7 Y. H. Dou, L. Zhang, X. Xu, Z. Q. Sun, T. Liao and S. X. Dou, *Chem. Soc. Rev.*, 2017, **46**, 7338–7373.
- 8 C. L. Tan and H. Zhang, *Nat. Commun.*, 2015, **6**, 7873.
- 9 S. L. Zhao, Y. Wang, Q. H. Zhang, Y. F. Li, L. Gu, Z. H. Dai, S. L. Liu, Y. Q. Lan, M. Han and J. C. Bao, *Inorg. Chem. Front.*, 2016, **3**, 1501–1509.
- 10 S. Liu, L. Xie, H. Qian, G. Y. Liu, H. Zhong and H. B. Zeng, *J. Mater. Chem. A*, 2019, **7**, 15411–15419.
- 11 D. Wang, W. W. Zhou, R. Zhang, J. J. Zeng, Y. Du, S. Qi, C. X. Cong, C. Y. Ding, X. X. Huang, G. W. Wen and T. Yu, *Adv. Mater.*, 2018, **30**, 1803569.
- 12 X. Y. Zhang, Z. Zhang, J. Liang, Y. G. Zhou, Y. C. Tong, Y. Wang and X. X. Wang, *J. Mater. Chem. A*, 2017, **5**, 9702–9708.
- 13 Y. Y. Zhan, Z. B. Shao, T. H. Jiang, J. Ye, X. F. Wu, B. C. Zhang, K. Ding, D. Wu and J. S. Jie, *J. Mater. Chem. A*, 2020, **8**, 789–796.
- 14 B. Jin, P. Huang, Q. Zhang, X. Zhou, X. W. Zhang, L. Li, J. W. Su, H. Q. Li and T. Y. Zhai, *Adv. Funct. Mater.*, 2018, **28**, 1800181.
- 15 D. D. Zhu, J. Xia, L. Wang, X. Z. Li, L. F. Tian and X. M. Meng, *Nanoscale*, 2016, **8**, 11375–11379.

- 16 R. Q. Cheng, Y. Wen, L. Yin, F. M. Wang, F. Wang, K. L. Liu, T. A. Shifa, J. Li, C. Jiang, Z. X. Wang and J. He, *Adv. Mater.*, 2017, **29**, 1703122.
- 17 Y. Wen, Q. S. Wang, L. Yin, Q. Liu, F. Wang, F. M. Wang, Z. X. Wang, K. L. Liu, K. Xu, Y. Huang, T. A. Shifa, C. Jiang, J. Xiong and J. He, *Adv. Mater.*, 2016, **28**, 8051–8057.
- 18 W. Feng, W. Zheng and P. Hu, *Phys. Chem. Chem. Phys.*, 2014, **16**, 19340–19344.
- 19 H. Liu, M. Yu, F. Qin, W. Feng and P. Hu, *ACS Appl. Nano Mater.*, 2018, **1**, 5414–5418.
- 20 O. Ramirez, P. Ramasamy, Y. C. Choi and J. S. Lee, *Chem. Mater.*, 2019, **31**, 268–276.
- 21 S. D. Deshmukh, R. G. Ellis, D. S. Sutandar, D. J. Rokke and R. Agrawal, *Chem. Mater.*, 2019, **31**, 9087–9097.
- 22 N. Oh, L. P. Keating, G. A. Drake and M. Shim, *Chem. Mater.*, 2019, **31**, 1973–1980.
- 23 S. O. Hinterding, A. C. Berends, M. Kurttepel, M.-E. Moret, J. D. Meeldijk, S. Bals, W. van der Stam and C. de Mello Donega, *ACS Nano*, 2019, **13**, 12880–12893.
- 24 H. L. Guthrey, A. G. Norman, J. Nishinaga, S. Niki, M. M. Al-Jassim and H. Shibata, *ACS Appl. Mater. Interfaces*, 2020, **12**, 3150–3160.
- 25 O. Yarema, M. Yarema and V. Wood, *Chem. Mater.*, 2018, **30**, 1446–1461.
- 26 B. Chen, N. Pradhan and H. Zhong, *J. Phys. Chem. Lett.*, 2018, **9**, 435–445.
- 27 A. C. Berends, M. J. Mangnus, C. Xia, F. T. Rabouw and C. de Mello Donega, *J. Phys. Chem. Lett.*, 2019, **10**, 1600–1616.
- 28 F.-J. Fan, L. Wu and S.-H. Yu, *Energy Environ. Sci.*, 2014, **7**, 190–208.
- 29 X. J. Wu, X. Huang, X. Qi, H. Li, B. Li and H. Zhang, *Angew. Chem., Int. Ed.*, 2014, **53**, 8929–8933.
- 30 T. Plirdpring, K. Kurosaki, A. Kosuga, T. Day, S. Firdosy, V. Ravi, G. J. Snyder, A. Harnwungmoung, T. Sugahara and Y. Ohishi, *Adv. Mater.*, 2012, **24**, 3622–3626.
- 31 H. L. Hwang, H. H. Chang, P. Sharma, A. J. Letha, L. Shao, Y. Zhang and B. H. Tseng, *Adv. Sci.*, 2016, **3**, 1500196.
- 32 J.-J. Wang, Y.-Q. Wang, F.-F. Cao, Y.-G. Guo and L.-J. Wan, *J. Am. Chem. Soc.*, 2010, **132**, 12218–12221.
- 33 C. Xia, W. Wu, T. Yu, X. Xie, C. Van Oversteeg, H. C. Gerritsen and C. de Mello Donega, *ACS Nano*, 2018, **12**, 8350–8361.
- 34 J. Tang, S. Hinds, S. O. Kelley and E. H. Sargent, *Chem. Mater.*, 2008, **20**, 6906–6910.
- 35 C.-R. Ho, M.-L. Tsai, H.-J. Jhuo, D.-H. Lien, C.-A. Lin, S.-H. Tsai, T.-C. Wei, K.-P. Huang, S.-A. Chen and J.-H. He, *Nanoscale*, 2013, **5**, 6350–6355.
- 36 M. G. Panthani, V. Akhavan, B. Goodfellow, J. P. Schmidtke, L. Dunn, A. Dodabalapur, P. F. Barbara and B. A. Korgel, *J. Am. Chem. Soc.*, 2008, **130**, 16770–16777.
- 37 J. F. L. Lox, Z. Y. Dang, M. L. Anh, E. Hollinger and V. Lesnyak, *Chem. Mater.*, 2019, **31**, 2873–2883.
- 38 J. B. Rivest and P. K. Jain, *Chem. Soc. Rev.*, 2013, **42**, 89–96.
- 39 B. C. Steimle, A. M. Fagan, A. G. Butterfield, R. W. Lord, C. R. McCormick, G. A. Di Domizio and R. E. Schaak, *Chem. Mater.*, 2020, **32**, 5461–5482.
- 40 S. Gupta, S. V. Kershaw and A. L. Rogach, *Adv. Mater.*, 2013, **25**, 6923–6944.
- 41 V. Lesnyak, C. George, A. Genovese, M. Prato, A. Casu, S. Ayyappan, A. Scarpellini and L. Manna, *ACS Nano*, 2014, **8**, 8407–8418.
- 42 J. M. Hodges, K. Kletetschka, J. L. Fenton, C. G. Read and R. E. Schaak, *Angew. Chem., Int. Ed.*, 2015, **54**, 8669–8672.
- 43 Y. Liu, M. Liu and M. T. Swihart, *J. Am. Chem. Soc.*, 2017, **139**, 18598–18606.
- 44 B. J. Beberwyck, Y. Surendranath and A. P. Alivisatos, *J. Phys. Chem. C*, 2013, **117**, 19759–19770.
- 45 A. C. Berends, W. van der Stam, Q. A. Akkerman, J. D. Meeldijk, J. van der Lit and C. D. Donega, *Chem. Mater.*, 2018, **30**, 3836–3846.
- 46 Y. X. Wang, Y. V. Morozov, M. Zhukovskiy, R. Chatterjee, S. Draguta, P. Tongying, B. Bryant, S. Rouvimov and M. Kuno, *ACS Energy Lett.*, 2016, **1**, 175–181.
- 47 R. L. Brutchey, *Acc. Chem. Res.*, 2015, **48**, 2918–2926.
- 48 T. Wang, S. S. Farvid, M. Abulikemu and P. V. Radovanovic, *J. Am. Chem. Soc.*, 2010, **132**, 9250–9252.
- 49 W. L. Wang, W. L. Feng, Q. Li, Y. T. Zhao, D. Zhao, Z. H. Xia, W. J. Wang, S. L. Zhang, X. X. Zheng and Z. H. Jing, *Cryst. Growth Des.*, 2019, **19**, 1226–1232.
- 50 J. Li, Z. G. Jin, T. Liu, J. Wang, D. L. Wang, J. Y. Lai, H. Y. Du and L. Cui, *CrystEngComm*, 2013, **15**, 7327–7338.
- 51 W. Wang, W. Feng, T. Ding and Q. Yang, *Chem. Mater.*, 2015, **27**, 6181–6184.
- 52 M. Bandi, V. Zade, S. Roy, A. N. Nair, S. Seacat, S. Sreenivasan, V. Shutthanandan, C. G. Van de Walle, H. Peelaers and C. V. Ramana, *Cryst. Growth Des.*, 2020, **20**, 1422–1433.
- 53 P. U. Londhe, A. B. Rohom, R. Fernandes, D. C. Kothari and N. B. Chaure, *ACS Sustainable Chem. Eng.*, 2018, **6**, 4987–4995.
- 54 W. L. Wang, J. Jiang, T. Ding, C. D. Wang, J. Zuo and Q. Yang, *ACS Appl. Mater. Interfaces*, 2015, **7**, 2235–2241.
- 55 Y. X. Wang, M. Zhukovskiy, P. Tongying, Y. Tian and M. Kuno, *J. Phys. Chem. Lett.*, 2014, **5**, 3608–3613.
- 56 V. Lesnyak, R. Brescia, G. C. Messina and L. Manna, *J. Am. Chem. Soc.*, 2015, **137**, 9315–9323.
- 57 W. L. Wang, T. Ding, G. H. Chen, L. Zhang, Y. Q. Yu and Q. Yang, *Nanoscale*, 2015, **7**, 15106–15110.
- 58 W. L. Wang, L. Zhang, G. H. Chen, J. Jiang, T. Ding, J. Zuo and Q. Yang, *CrystEngComm*, 2015, **17**, 1975–1981.
- 59 D. W. Houck, S. V. Nandu, T. D. Siegler and B. A. Korgel, *ACS Appl. Nano Mater.*, 2019, **2**, 4673–4680.