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# Structural diversity dependent cation incorporation into magnetic Cr–Se nanocrystals†

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Chromium selenide (Cr–Se)-based low-dimensional materials have attracted significant attention in spintronics because of their diverse structure- and composition-dependent magnetic properties. While significant progress has been made in fabricating the Cr–Se family of nanomaterials through techniques like chemical vapor deposition, the synthesis of Cr–Se nanocrystals (NCs) *via* colloidal methods remains underexplored. In this work, we demonstrate the robust colloidal synthesis approach for producing Cr<sub>2</sub>Se<sub>3</sub> and Cr<sub>3</sub>Se<sub>4</sub> NCs with distinct morphologies by varying the Cr precursors and ligands. Cr–Se NCs can serve as templates for cation exchange (CE) reactions involving monovalent Cu<sup>+</sup> and Ag<sup>+</sup>, divalent Zn<sup>2+</sup> and Cd<sup>2+</sup>, and trivalent In<sup>3+</sup>, facilitating the creation of a diverse library of metal selenide NCs and nanoheterostructures. Our findings highlight how the outcomes of CE reactions are influenced by the structure of various Cr–Se phases. Furthermore, the magnetic properties of the as-synthesized Cr–Se NCs and their derivative CuCrSe<sub>2</sub> were investigated. Our work provides a robust synthesis route for the Cr–Se class of magnetic nanomaterials and a platform for creating a diverse range of functional metal selenide NCs *via* CE reactions.

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## Introduction

The Cr–Se family of materials has garnered significant attention in the exploration of two-dimensional magnets due to their high Curie temperatures and air-stability, providing opportunities to investigate a wide range of magnetic properties in the 2D realm. Various stable and metastable stoichiometries within the Cr–Se material class, such as CrSe,<sup>1</sup> Cr<sub>2</sub>Se<sub>3</sub>,<sup>2</sup> Cr<sub>3</sub>Se<sub>4</sub>,<sup>2</sup> Cr<sub>5</sub>Se<sub>8</sub>,<sup>3</sup> Cr<sub>9</sub>Se<sub>13</sub>,<sup>4</sup> and CrSe<sub>2</sub>,<sup>5,6</sup> have been synthesized. For instance, He's group demonstrated the presence of noticeable ferromagnetic properties in 2D CrSe crystals grown *via* chemical vapor deposition (CVD) at temperatures below 280 K.<sup>1</sup> In another study, Duan *et al.* grew air-stable CrSe<sub>2</sub> nanosheets on WSe<sub>2</sub> substrates using a CVD approach, showcasing ferromagnetic behavior.<sup>5</sup> Despite the predominance of previous research on Cr–Se nanomaterials relying on CVD and exfoliation techniques, which suffer from low productivity, hindering large-scale practical applications, colloidal synthesis emerges as a promising method for crafting 2D materials with a tailored morphology, thickness, and composition.<sup>7,8</sup> More importantly, it enables the production of freestanding 2D materials amenable for doping *via* the chemical method<sup>9</sup> and integration into thin-film devices through solution processing.<sup>10,11</sup> Despite the recent progress in colloidal Cr-based NCs,<sup>12–16</sup> to the best of our knowledge, the colloidal synthesis of Cr–Se NCs remains unexplored.

In colloidal synthesis, the CE reaction, which describes the scenario that the cations in the host lattice are replaced by

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materials for electrocatalysis and energy storage devices, as well as the discovery of novel magnetic two-dimensional materials.



foreign cations without largely modifying the anionic sublattice, presents a distinctive avenue for generating a diverse array of metal chalcogenide NC libraries.<sup>17,18</sup> The copper chalcogenide class of nanomaterials, *i.e.*, Cu<sub>2-x</sub>S, Cu<sub>2-x</sub>Se, and Cu<sub>2-x</sub>Te with  $0 \leq x \leq 1$ , serves as the robust template for CE reactions due to the varied crystal structures and morphologies achievable through colloidal synthesis.<sup>19,20</sup> Furthermore, the Cu<sup>+</sup> ions in the degenerately doped structure possess fluid-like ion mobility at elevated temperature.<sup>21</sup> In the metal selenide regime, CdSe NCs can serve as the template for CE with Cu<sup>+</sup> to produce Cu<sub>2</sub>Se and Cu<sub>2-x</sub>Se NCs.<sup>22</sup> Besides these advancements, there is a need for additional metal selenide templates with a tunable morphology and crystal structures for not only understanding the mechanism of CE reactions on metal selenides, but also fostering the development of previously unprecedented metal selenide-based nanostructures.

In this work, we demonstrate that Cr-Se NCs can serve as templates for CE reactions, facilitating the creation of a diverse range of metal selenide NCs. By employing Cr(acac)<sub>3</sub> or CrCl<sub>2</sub> precursors, we synthesised Cr-Se NCs with varying crystal structures and morphologies, resulting in the production of distinct Cr<sub>2</sub>Se<sub>3</sub> and Cr<sub>3</sub>Se<sub>4</sub> NCs. The subsequent CE reactions with monovalent Cu<sup>+</sup> and Ag<sup>+</sup>, as well as trivalent In<sup>3+</sup>, led to distinct reaction products when using the two Cr-Se templates, while the outcomes were similar when divalent Zn<sup>2+</sup> and Cd<sup>2+</sup> were applied. For example, ultrathin InSe nanosheets with lateral dimensions exceeding a micron can be synthesized *via* In<sup>3+</sup> exchange with Cr<sub>2</sub>Se<sub>3</sub> nanoplatelets. The valence selectivity of CE reactions originated from the structural disparities between more symmetric Cr<sub>2</sub>Se<sub>3</sub> unit cells and lattice-disordered Cr<sub>3</sub>Se<sub>4</sub>. Additionally, we investigated the magnetic properties of Cr-Se NCs and their derivative CuCrSe<sub>2</sub>. Our work provides valuable insights into the relationship between structural dependent cation incorporation and magnetic properties of Cr-Se based low-dimensional nanomaterials.

## Experimental

### Chemicals

Oleylamine (OAm, 80–90%) and cadmium(II) chloride (CdCl<sub>2</sub>, 99.99%) were purchased from Aladdin. Trioctylphosphine (TOP, 97%), chromium(III) acetylacetonate (Cr(acac)<sub>3</sub>, 97%), copper(II) acetylacetonate (Cu(acac)<sub>2</sub>, 97%), zinc(II) chloride (ZnCl<sub>2</sub>, ≥98%), hexamethyldisilazane (HMDS, 99.9%) and oleic acid (OA, 90%) were purchased from Sigma-Aldrich. Tri-*n*-octylamine (TOA, 97%) was purchased from TCI Chemicals. Chromium(II) chloride (CrCl<sub>2</sub>, 99.99%), chromium(0) hexacarbonyl (Cr(CO)<sub>6</sub>, 99%), selenium (Se, 99.999%, 200 mesh), and indium(III) chloride (InCl<sub>3</sub>, 99.99%) were purchased from Alfa Aesar. Silver(I) acetate (Ag(OAc), 99%) was purchased from Adamas-beta. All chemicals were used as received.

### Synthesis of Cr-Se NCs

All NC syntheses were carried out using standard Schlenk techniques under a nitrogen atmosphere. The detailed synthetic

parameters and product NCs are provided in Table S1.† In a typical synthesis of Cr-Se NCs, 1.0 mmol CrCl<sub>2</sub>, 1 mL OA, 1 mL HMDS, and 6 mL of TOA were loaded into a three-neck flask. Under a nitrogen flow, the mixture was maintained at room temperature for 30 min and then heated at 100 °C for another 30 min to form a clear solution. In a separate vial, the TOP-Se precursor was prepared by dissolving 2.0 mmol of Se powder into 2.0 mL of TOP *via* stirring in a nitrogen-filled glove box at 100 °C. The reaction mixture was heated to 300 °C before injection of the TOP-Se precursor. The reaction was kept at the reaction temperature for 3 hours before being air-cooled to room temperature. The Cr-Se NCs were isolated by 10 mL ethanol precipitation and centrifugation at 4000 rpm for 5 min. The NCs were redispersed in 10 mL of hexane and precipitated with 10 mL of ethanol before being centrifuged at 4000 rpm for 3 min. After another cycle of precipitation and centrifugation, Cr-Se NCs were dissolved in hexane for further use. The synthesis of Cr-Se nanoplatelets and nanoflowers follows the same synthetic procedure except for using different metal precursors, ligands, and synthetic parameters.

### CE reaction with Cr-Se NCs

The detailed synthetic parameters and product NCs are provided in Table S2.† In a typical synthesis of InSe nanosheets, 0.05 mmol InCl<sub>3</sub> and 6 mL OAm were loaded into a three-neck flask. Under a nitrogen flow, the mixture was maintained at room temperature for 30 min and then heated at 100 °C for another 30 min to form a clear solution. In a separate vial, Cr-Se nanoplatelet solution was prepared by dissolving 13.1 mg dry Cr-Se nanoplatelets into 3.0 mL OAm and 1.0 mL TOP by sonication. After injection of the Cr-Se solution, the reaction mixture was heated to 340 °C and kept at this temperature for 2 hours before being air-cooled to room temperature. The InSe nanosheets were isolated by 10 mL ethanol precipitation and centrifugation at 4000 rpm for 5 min. The nanosheets were redispersed in 10 mL of hexane and precipitated with 10 mL of ethanol before being centrifuged at 4000 rpm for 3 min. After another cycle of precipitation and centrifugation, InSe nanosheets were dissolved in hexane for further use. The synthesis of InSe nanosheets using Cr-Se nanoplates and nanoflowers follows the same synthetic procedure except for using different synthetic parameters.

### Characterization

Transmission electron microscopy (TEM) images, high-resolution TEM (HRTEM) images, and STEM-energy-dispersive X-ray spectroscopy (STEM-EDS) maps were obtained with a JEM-F200 operating at an accelerating voltage of 200 kV. TEM grids were prepared by placing a dilute NC dispersion onto a carbon-coated copper grid and allowing the sample to air dry. Powder X-ray diffraction (XRD) measurements were performed using a Rigaku Ultima IV diffractometer equipped with a Cu K $\alpha$  X-ray source. Samples were prepared by drop-casting concentrated NC dispersions onto glass slides. X-ray photoelectron spectroscopy (XPS) spectra were acquired using an ESCALAB 250Xi with an Al K $\alpha$  X-ray source (1486.6 eV) and a nominal spot diameter of 400  $\mu$ m. The analysis chamber pressure was maintained below  $5 \times 10^{-8}$

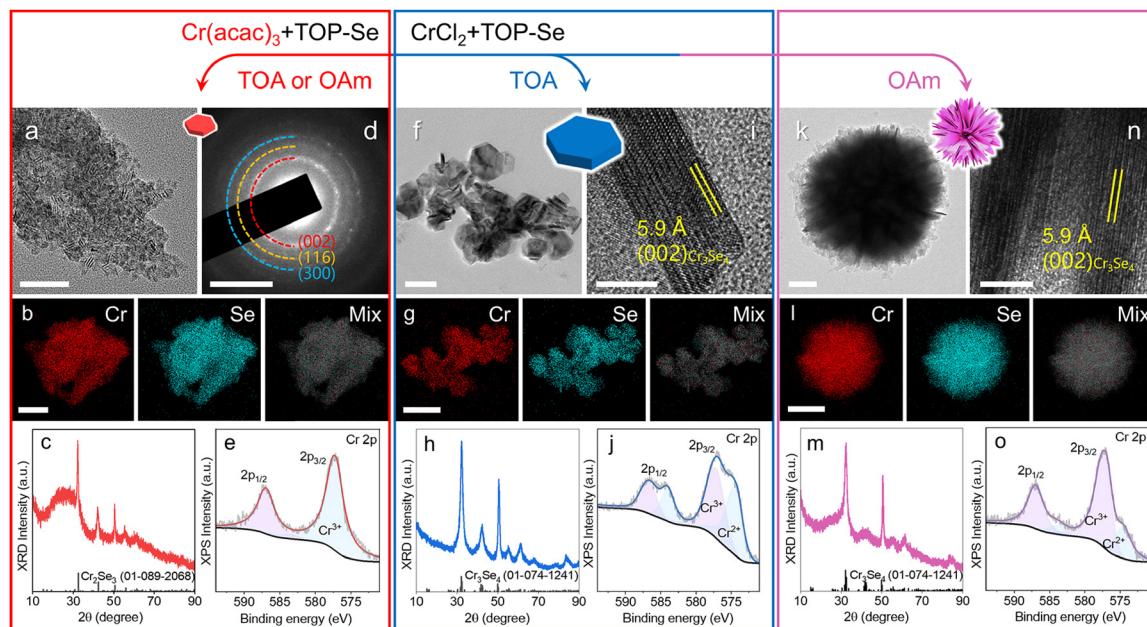


mbar. Magnetic hysteresis ( $M$ - $H$ ) loops, along with field-cooling (FC) and zero-field cooling (ZFC) curves, were measured using a Physical Property Measurement System (PPMS).

## Results and discussion

We synthesized Cr-Se NCs with different crystal structures and morphologies *via* a colloidal method. In general, the Se precursor was introduced into the Cr precursor complex *via* a hot-injection approach. When  $\text{Cr}(\text{acac})_3$  was used as the Cr precursor, the synthesis yielded small nanoplatelets, with an average diameter of  $13.2 \pm 0.1$  nm, as demonstrated by the TEM image in Fig. 1a. Size distribution histograms for this and the following samples are provided in Fig. S1 and S9.† The STEM-EDS elemental maps (Fig. 1b) demonstrate uniform distribution of Cr and Se signals across the nanoplatelets with a Cr : Se ratio of 42.6% : 57.4%, indicating the presence of Se vacancies. The corresponding EDS spectra and quantitative results of the elemental maps in this work are collectively provided in Fig. S2–S3 and Table S3.† The XRD pattern and selective area electron diffraction (SAED) pattern (Fig. 1c and d) suggest the formation of the  $\text{Cr}_2\text{Se}_3$  phase, with diffraction peaks/rings that correspond to the (002), (116), and (300) planes within the  $\text{Cr}_2\text{Se}_3$  unit cell. As shown in Fig. 1e, the XPS spectrum in the Cr 2p region reveals the peaks at 587.0 eV and 577.2 eV, suggesting that Cr is exclusively present as  $\text{Cr}^{3+}$ . The peaks observed in the XPS spectrum in the Se 3d region correspond to  $\text{Se}^{2-}$  (Fig. S4a†). The XPS results matched well with the previous observations on  $\text{Cr}_2\text{Se}_3$ .<sup>2,23</sup>

In contrast, the synthesis of Cr-Se NCs involving the  $\text{CrCl}_2$  precursor led to the formation of  $\text{Cr}_3\text{Se}_4$  NCs with a larger size in nanoplate and nanoflower morphologies, featuring the average size of  $45.5 \pm 0.2$  nm and  $444.0 \pm 6.6$  nm, respectively, with TOA and OAm ligands (Fig. 1f and k, and Fig. S1†). The STEM-EDS elemental maps illustrate the co-localization of Cr and Se signals on the individual nanoplate and nanoflower with Cr : Se ratios of 43.1% : 56.9% and 43.2% : 56.8%, respectively (Fig. 1g and l), consistent with their XRD patterns that matched well with the  $\text{Cr}_3\text{Se}_4$  phase (Fig. 1h and m). Note that the  $\text{Cr}_2\text{Se}_3$  nanoplatelets and  $\text{Cr}_3\text{Se}_4$  nanoplates exhibited varying full width at half maxima (FWHM) in XRD peaks, reflecting differences in crystalline size and anisotropy. Furthermore, the HRTEM images reveal an observed lattice spacing of 5.9 Å on the side-view of the nanoplate, corresponding to the (002) planes in the  $\text{Cr}_3\text{Se}_4$  unit cell (Fig. 1i), highlighting that the basal planes of the nanoplates are (002) planes in  $\text{Cr}_3\text{Se}_4$ . The XPS spectra of Cr (Fig. 1j and o) and Se (Fig. S4b and c†) for nanoplates and nanoflowers indicated that Cr is present as a mixture of  $\text{Cr}^{2+}$  and  $\text{Cr}^{3+}$ , while Se is present as  $\text{Se}^{2-}$ , consistent with a previously report on  $\text{Cr}_3\text{Se}_4$ .<sup>2</sup> The formation of the plate-like morphology is attributed to the layered atomic arrangement of the unit cell (Fig. S5†), as supported by the side-view HRTEM image. A similar lattice structure was observed on the nanoflowers, indicating that the nanoflowers were composed of assembled nanoplates (Fig. 1n). Ligand-assisted phase engineering and morphology engineering have been employed as powerful tools for creating diverse NCs with a controlled morphology, composition and crystal structure.<sup>12,24</sup> TOA, considered a weakly coordinating



**Fig. 1** Synthesis of Cr-Se NCs. (a) TEM image, (b) STEM-EDS elemental maps of Cr and Se, (c) XRD pattern, (d) SAED pattern, and (e) XPS spectrum of Cr for  $\text{Cr}_2\text{Se}_3$  nanoplatelets. (f) TEM image, (g) STEM-EDS elemental maps of Cr and Se, (h) XRD pattern, (i) HRTEM image, and (j) XPS spectrum of Cr for  $\text{Cr}_3\text{Se}_4$  nanoplates. (k) TEM image, (l) STEM-EDS elemental maps of Cr and Se, (m) XRD pattern, (n) HRTEM image, and (o) XPS spectrum of Cr for  $\text{Cr}_3\text{Se}_4$  nanoflowers. Scale bars: (a) 50 nm, (b, g and l) 200 nm, (f and k) 100 nm, (i and n) 5 nm, and (d)  $5 \text{ nm}^{-1}$ .

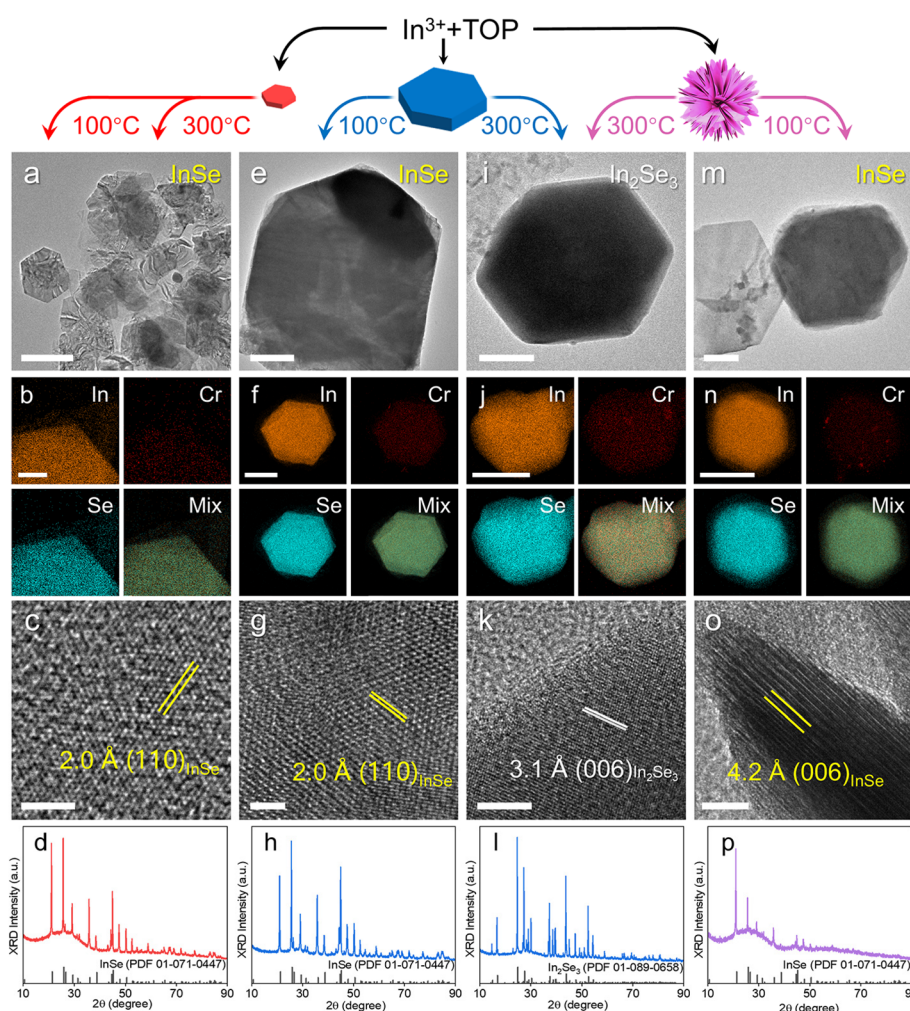




ligand compared to OAm due to its larger steric bulk and lower ligand surface packing, likely contributes to a more disordered and thicker ligand shell during nanoplate growth, preventing the formation of the assembled nanoplate architectures. Compared with small nanoplatelets, the formation of nanoplates and nanoflowers can be attributed to the utilization of an inorganic Cr precursor ( $\text{CrCl}_2$ ), whose reactivity is higher than that of  $\text{Cr}(\text{acac})_3$ . This can be also supported by the increased Cr:Se ratio from the stoichiometry between two phases and the increased size of the NCs. Furthermore, the synthesis of Cr–Se NCs using  $\text{Cr}(\text{CO})_6$  leads to the formation of  $\text{Cr}_2\text{Se}_3$  NCs, regardless of the ligands (OAm or TOA) involved in the reaction (Fig. S6†). The obtained NCs exhibited a nanoplatelet morphology, indicating that the  $\text{Cr}(\text{CO})_6$  precursor may exhibit a similar reactivity to  $\text{Cr}(\text{acac})_3$ .

CE reactions have emerged as a versatile method for tailoring the composition and crystal structure of metal selenide

NCs.<sup>17,25,26</sup> A template with diverse crystal structures and morphologies is pivotal for expanding the range of accessible metal selenide nanostructures. While copper selenide templates have commonly been used in CE reactions to synthesize a diverse range of metal selenide NCs, we demonstrate that the Cr–Se NCs can also serve as templates during CE reactions. We first explored the CE reactions on Cr–Se NCs with  $\text{In}^{3+}$ . The In–Se class of materials is known for their unique properties such as a large band gap, high mobility, and highly anisotropic electrical and thermal properties, making them promising for applications in optoelectronics, thermoelectric devices and photocatalysis.<sup>27–30</sup> The controlled syntheses in the previous reports yielded distinct In–Se phases in either  $\text{InSe}^{29}$  or  $\text{In}_2\text{Se}_3$ ,<sup>28</sup> while the CE reaction with Cr–Se NCs in our work led to the formation of both phases, depending on the Cr–Se template used and the injection temperature. As shown in Fig. 2a, at an injection temperature of 300 °C, monodisperse



**Fig. 2** Synthesis of In–Se nanosheets *via* CE reactions. TEM image, STEM-EDS elemental maps of In, Cr and Se, HRTEM image, XRD pattern of (a–d) InSe nanosheets obtained using  $\text{In}^{3+}$  exchange with  $\text{Cr}_2\text{Se}_3$  nanoplatelets at 300 °C. (e–h) InSe nanosheets obtained using  $\text{In}^{3+}$  exchange with  $\text{Cr}_3\text{Se}_4$  nanoplates at 100 °C and (i–l)  $\text{In}_2\text{Se}_3$  nanosheets obtained using  $\text{In}^{3+}$  exchange with  $\text{Cr}_3\text{Se}_4$  nanoplates at 300 °C. (m–p) InSe nanosheets obtained using  $\text{In}^{3+}$  exchange with  $\text{Cr}_3\text{Se}_4$  nanoflowers at 100 °C. Scale bars: (a, e and f) 1  $\mu\text{m}$ , (c and g) 2 nm, (k and o) 5 nm, (b and i) 100 nm, (j and n) 500 nm, and (m) 200 nm.



nanosheets with a lateral size approaching a micron can be produced *via* CE reactions employing small  $\text{Cr}_2\text{Se}_3$  nanoplatelet templates. The formation of larger nanosheets using  $\sim 13$  nm nanoplatelets suggested a ripening process occurring *via* the merging of smaller nanoplatelets during the CE reaction. STEM-EDS elemental maps (Fig. 2b) confirmed the uniformly distributed In and Se signals with an expected  $\sim 1:1$  In:Se ratio, along with minimal residual Cr signal presence. The side-view HRTEM image in Fig. 2c shows the lattice spacing of  $2.0 \text{ \AA}$ , corresponding to the (110) planes in InSe from XRD analysis (Fig. 2d). Similar InSe nanosheets were obtained by injecting  $\text{Cr}_2\text{Se}_3$  nanoplatelets into an In-complex solution at  $100^\circ\text{C}$  (Fig. S7†), indicating the negligible impact of injection temperature on the selective synthesis of In–Se compounds when using  $\text{Cr}_2\text{Se}_3$  templates.

When the CE of  $\text{In}^{3+}$  was performed with  $\text{Cr}_3\text{Se}_4$  nanoplates and nanoflowers at  $100^\circ\text{C}$ , comparable nanosheets were produced (Fig. 2e and m). The formation of the InSe phase was validated by the XRD patterns (Fig. 2h and p), the observed lattice spacings  $2.0 \text{ \AA}$  and  $4.2 \text{ \AA}$  for the (110) and (006) planes (Fig. 2g and o), and the homogeneously distributed In and Se signals in STEM-EDS elemental maps (Fig. 2f and n). In contrast to the use of small  $\text{Cr}_2\text{Se}_3$  templates, elevating the injection temperature with  $\text{Cr}_3\text{Se}_4$  nanoplates resulted in  $\text{In}_2\text{Se}_3$  nanosheets, as evidenced by the TEM image, HRTEM image, STEM-EDS elemental maps, and XRD patterns (Fig. 2i–l). Similarly, employing  $\text{Cr}_3\text{Se}_4$  nanoflowers as templates in CE reactions yielded predominantly the  $\text{In}_2\text{Se}_3$  phase, with a minor presence of InSe as a secondary phase (Fig. S8†). As shown in Fig. S9†, the results demonstrate that InSe nanosheets synthesized *via* CE reactions on  $\text{Cr}_3\text{Se}_4$  nanoplates exhibit the largest lateral dimensions and thickness ( $3.8 \mu\text{m}$  and  $68.6 \text{ nm}$ , respectively) compared to those grown on  $\text{Cr}_2\text{Se}_3$  nanoplatelets and  $\text{Cr}_3\text{Se}_4$  nanoflowers. This observation can be attributed to the highly faceted morphology of  $\text{Cr}_3\text{Se}_4$  nanoplates, which contrasts with the less defined structures of the other Cr–Se templates. The CE reaction, which is strongly dependent on cation diffusion, proceeds more efficiently on the well-defined facets of anisotropic NCs. In contrast,  $\text{Cr}_2\text{Se}_3$  nanoplatelets and  $\text{Cr}_3\text{Se}_4$  nanoflowers require significant morphological evolution to form large, faceted InSe nanosheets. Furthermore,  $\text{In}_2\text{Se}_3$  nanosheets are generally smaller but thicker than InSe nanosheets, a difference that can be explained by the non-layered structure of the  $\text{In}_2\text{Se}_3$  unit cell compared to the van der Waals structure of InSe (Fig. S10†).

Furthermore, the general CE reactions with Cr–Se NCs can be extended to various cations including  $\text{Cu}^+$ ,  $\text{Ag}^+$ ,  $\text{Zn}^{2+}$ , and  $\text{Cd}^{2+}$  (Fig. 3a). Fig. 3c1 shows a typical TEM image of the products obtained *via* the CE of  $\text{Cu}^+$  on small  $\text{Cr}_2\text{Se}_3$  nanoplatelets, revealing a lattice spacing of  $2.0 \text{ \AA}$  corresponding to the (220) planes of  $\text{Cu}_2\text{Se}$  (Fig. 3c2). The XRD pattern and the STEM-EDS elemental maps further confirm the formation of the  $\text{Cu}_2\text{Se}$  nanoparticles (Fig. 3b and c3), with the Cr content largely removed, indicating a tendency for  $\text{Cu}^+$  ions to completely replace the  $\text{Cr}^{3+}$  ions within the  $\text{Cr}_2\text{Se}_3$  lattice. By contrast, the CE reactions of  $\text{Cu}^+$  on  $\text{Cr}_3\text{Se}_4$  nanoplates and nanoflowers

lead to the ternary  $\text{CuCrSe}_2$  phase, retaining the morphology of the respective templates (Fig. 3d1 and e1). The formation of the ternary  $\text{CuCrSe}_2$  phase is confirmed by the XRD patterns and HRTEM images (Fig. 3b, d2, and e2). This indicates the potential of CE reactions in synthesizing 2D magnets in addition to conventional methods like CVD.<sup>31–33</sup> The STEM-EDS elemental maps confirmed the colocation of Cu, Cr, and Se with an atomic ratio of  $24.6\%:27.4\%:48.0\%$  and  $26.4\%:23.5\%:50.2\%$  (Fig. 3d3 and e3). These observations suggest that  $\text{Cu}^+$  ions tended to coexist with  $\text{Cr}^{3+}$  in the  $\text{Cr}_3\text{Se}_4$  lattice, further highlighting the significant influence of the crystal phase and morphology on the outcomes of CE reactions.

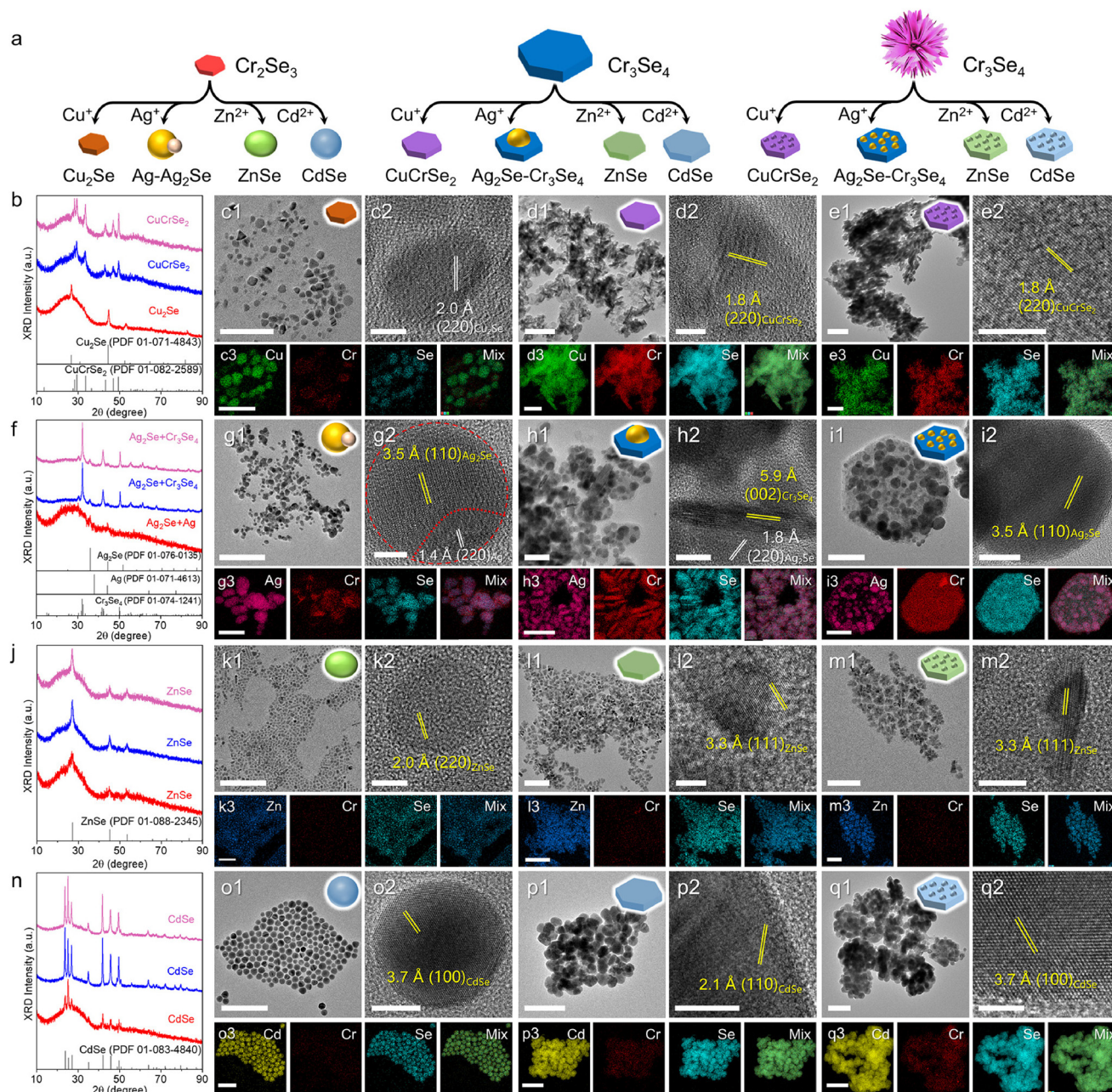
The CE reactions of  $\text{Ag}^+$  on Cr–Se NCs yielded results akin to those obtained with  $\text{Cu}^+$ . The reaction on  $\text{Cr}_3\text{Se}_4$  nanoplates and nanoflowers produced  $\text{Cr}_3\text{Se}_4\text{--Ag}_2\text{Se}$  heterostructures (Fig. 3f), suggesting that  $\text{Ag}^+$  partially occupied the surface of  $\text{Cr}_3\text{Se}_4$  NCs, followed by regioselective CE reactions (Fig. 3h1 and i1). The formation of these heterostructures was confirmed by the HRTEM image and separate signals of Cr and Ag (Fig. 3h2, h3 and i2, i3) as well as the clear heterointerface along the two domains in the HRTEM image (Fig. 3g1 and g2). In contrast, CE reactions on the small  $\text{Cr}_2\text{Se}_3$  nanoplatelets resulted in more complete reactions, forming  $\text{Cr}_2\text{Se}_3\text{--Ag}_2\text{Se--Ag}$  heterostructures, with the remaining Cr signals less pronounced compared to the Ag signals (Fig. 3g3). Notably, the lattice fringes were obviously different in the two areas of a single nanoparticle, consistent with our previous observation that the excess  $\text{Ag}^+$  could deposit on the  $\text{Ag}_2\text{S}$  surface as Ag metallic nanoparticles when the CE reaction was complete.<sup>34</sup>

The CE reactions of divalent  $\text{Zn}^{2+}$  and  $\text{Cd}^{2+}$  with Cr–Se NCs follow a similar trend, producing ZnSe and CdSe NCs, respectively, as demonstrated by the XRD patterns and homogeneous distribution of Zn/Cd and Se signals across the entire NCs (Fig. 3j–q). Generally, the morphologies of the products were preserved from the particle, plate, and flower templates, with the size of the products obtained with  $\text{Cd}^{2+}$  being consistently larger than those with  $\text{Zn}^{2+}$  (Fig. S1†). This size difference arises from the larger ionic radius of  $\text{Cd}^{2+}$  ( $78 \text{ pm}$ ) compared to  $\text{Zn}^{2+}$  ( $60 \text{ pm}$ ), leading to lattice expansion during the CE process. This is supported by the observed larger lattice spacing in the HRTEM images,  $3.3 \text{ \AA}$  for ZnSe (Fig. 3m2) and  $3.7 \text{ \AA}$  for CdSe (Fig. 3q2).

The above observations highlight the significant role of the crystal structure in dictating the outcomes of CE reactions. The products obtained by the CE reaction involving monovalent cations ( $\text{Cu}^+$  and  $\text{Ag}^+$ ) and the trivalent cation ( $\text{In}^{3+}$ ) on  $\text{Cr}_2\text{Se}_3$  nanoplatelets are generally different from those obtained with  $\text{Cr}_3\text{Se}_4$  nanoplates and nanoflowers, while similar results are observed when divalent cations ( $\text{Zn}^{2+}$  and  $\text{Cd}^{2+}$ ) are used. We speculate that the discrepancy originated from the structural difference between  $\text{Cr}_2\text{Se}_3$  and  $\text{Cr}_3\text{Se}_4$  during CE reactions. The  $\text{Cr}_2\text{Se}_3$  unit cell comprises a hexagonal lattice, while  $\text{Cr}_3\text{Se}_4$  adopts a monoclinic structure. In the  $\text{Cr}_3\text{Se}_4$  unit cell, there are two types of Cr cations ( $\text{Cr}^{2+}$  and  $\text{Cr}^{3+}$ ) with the coexistence of both ordered and disordered octa-







**Fig. 3** CE reactions with  $\text{Cu}^+$ ,  $\text{Zn}^{2+}$ ,  $\text{Ag}^+$ , and  $\text{Cd}^{2+}$ . (a) Schematic illustration of the valence-dependent CE reaction on Cr–Se templates. (b) XRD patterns of  $\text{Cu}^+$  exchange with Cr–Se NCs. TEM image, STEM-EDS elemental maps, and HRTEM image of  $\text{Cu}^+$  exchange with (c)  $\text{Cr}_2\text{Se}_3$  nanoplatelets, (d)  $\text{Cr}_3\text{Se}_4$  nanoplates, and (e)  $\text{Cr}_3\text{Se}_4$  nanoflowers. (f) XRD patterns of  $\text{Ag}^+$  exchange with Cr–Se NCs. TEM image, STEM-EDS elemental maps, and HRTEM image of  $\text{Ag}^+$  exchange with (g)  $\text{Cr}_2\text{Se}_3$  nanoplatelets, (h)  $\text{Cr}_3\text{Se}_4$  nanoplates, and (i)  $\text{Cr}_3\text{Se}_4$  nanoflowers. (j) XRD patterns of  $\text{Zn}^{2+}$  exchange with Cr–Se NCs. TEM image, STEM-EDS elemental maps, and HRTEM image of  $\text{Zn}^{2+}$  exchange with (k)  $\text{Cr}_2\text{Se}_3$  nanoplatelets, (l)  $\text{Cr}_3\text{Se}_4$  nanoplates, and (m)  $\text{Cr}_3\text{Se}_4$  nanoflowers. (n) XRD patterns of  $\text{Cd}^{2+}$  exchange with Cr–Se NCs. TEM image, STEM-EDS elemental maps, and HRTEM image of  $\text{Cd}^{2+}$  exchange with (o)  $\text{Cr}_2\text{Se}_3$  nanoplatelets, (p)  $\text{Cr}_3\text{Se}_4$  nanoplates, and (q)  $\text{Cr}_3\text{Se}_4$  nanoflowers. Scale bars: (c1, l1, m1, i1, m3, h3, i3, k3, l3 and o3) 100 nm, (d1, e1, g1, k1, o1, p1, q1, e3, p3 and q3) 200 nm, (c3, d3, g3 and h1) 50 nm, (m2, h2, o2 and p2) 10 nm, and (c2, d2, e2, g2, i2, l2, k2 and q2) 5 nm.

hedron-coordinated centers, leading to a higher degree of lattice disorder in  $\text{Cr}_3\text{Se}_4$ . These features allow the  $\text{Cr}_3\text{Se}_4$  structure to provide more available sites for incoming cations to replace, potentially lowering the energy barrier for Cr ions to coexist with incoming cations, thereby fostering greater

structural diversity in the products post-CE reactions. Furthermore, the coexistence of  $\text{Cr}^{3+}$  and  $\text{Cu}^+$  in the  $\text{CuCrSe}_2$  unit cell, the stoichiometrically decreased In : Se ratio in  $\text{In}_2\text{Se}_3$  with a disordered lattice (Fig. S10†), and the predominant formation of the  $\text{Cr}_3\text{Se}_4$ – $\text{Ag}_2\text{Se}$  heterostructures after CE reactions

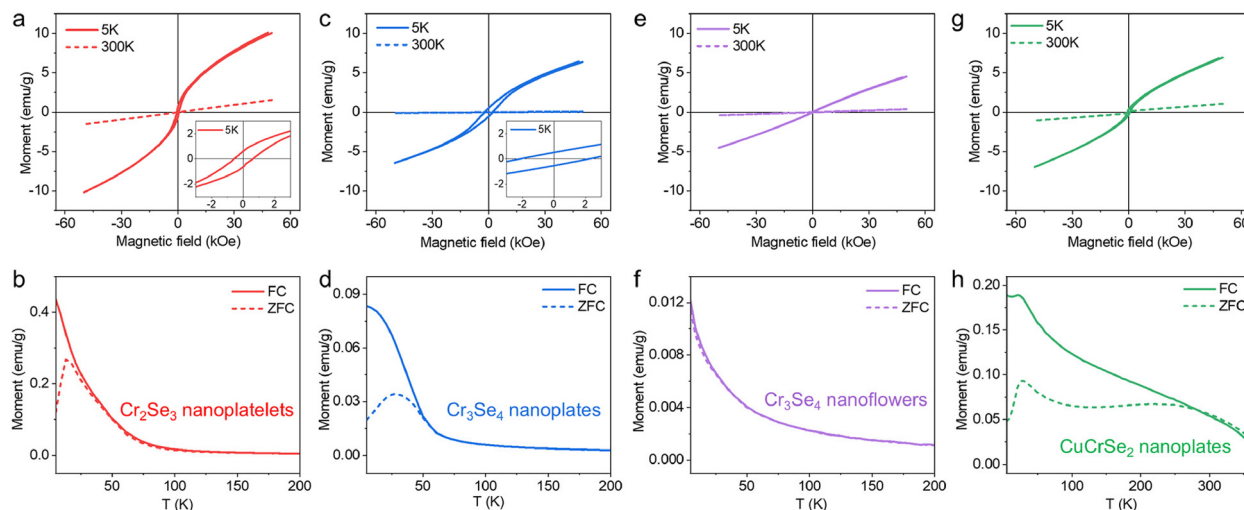


on  $\text{Cr}_3\text{Se}_4$  indicate a tendency towards relatively incomplete CE processes. By contrast, the more symmetric  $\text{Cr}_2\text{Se}_3$  unit cell facilitates more complete CE reactions, forming  $\text{Cu}_2\text{Se}$ ,  $\text{InSe}$ , and  $\text{Ag}_2\text{Se}$ - $\text{Ag}$  heterostructures when  $\text{Cu}^+$ ,  $\text{In}^{3+}$  and  $\text{Ag}^+$  ions are used, respectively. For divalent cations, the CE reactions produced binary  $\text{ZnSe}$  and  $\text{CdSe}$  indiscriminately, as other stoichiometric  $\text{Zn-Se}$  or  $\text{Cd-Se}$  phases may not be stabilized under ambient conditions. These results underscore that CE reactions can be used to identify the crystal phase and structures of the templates.<sup>35–37</sup>

For In exchange, when  $\text{Cr}_2\text{Se}_3$  nanoplatelets were used as a template, the reaction yielded  $\text{InSe}$  nanosheets.  $\text{InSe}$  exhibits promising physical properties including higher carrier mobilities and thickness-dependent bandgaps, which enhance its potential for application in field-effect transistors (FETs) with high on/off current ratios at room temperature.<sup>38–40</sup> In contrast, when  $\text{Cr}_3\text{Se}_4$  nanoplates and nanoflowers were used as templates for cation exchange, both  $\text{InSe}$  and  $\text{In}_2\text{Se}_3$  nanosheets were obtained.  $\text{In}_2\text{Se}_3$  has become one of the leading candidates for visible photodetection, owing to its narrow bandgap, direct bandgap characteristics, high absorption coefficient within the visible spectrum, and exceptional sensitivity.<sup>28,41–43</sup> These results highlight the potential of the template in regulating the properties of cation exchange products. For monovalent  $\text{Cu}^+$  exchange, the complete cation exchange product  $\text{Cu}_2\text{Se}$ , which was obtained using the  $\text{Cr}_2\text{Se}_3$  template, features a p-type semiconductor with a tunable bandgap (1.1–1.5 eV) and a direct bandgap (2.0–2.3 eV). These characteristics allow  $\text{Cu}_2\text{Se}$  to play significant roles in energy conversion and storage applications, including solar cells, lithium-ion batteries, supercapacitors, and thermoelectric devices.<sup>44–46</sup> By contrast, the incorporation of Cu into  $\text{Cr}_3\text{Se}_4$  leads to the formation of  $\text{CuCrSe}_2$  NCs, which exhibit antiferromagnetism. Very recently, multiferroicity has been observed in thin layer  $\text{CuCrSe}_2$ ,<sup>32,33</sup> highlighting the influence of the

template on the properties of products after cation exchange reactions.  $\text{Ag}_2\text{Se}$  is a typical n-type thermoelectric material near room temperature with high carrier mobility and low thermal conductivity.<sup>47,48</sup> Integration with metallic Ag nanoparticles has the potential to enhance the thermoelectric performance of the  $\text{Ag}_2\text{Se}$  domain by improving electron transfer.  $\text{ZnSe}$  and  $\text{CdSe}$  are well-known for their applications in light-emitting diodes (LEDs) and optoelectronic devices.<sup>49,50</sup> Although their crystal phase and stoichiometry are difficult to tune *via* the choice of Cr-Se templates for cation exchange, the resulting  $\text{ZnSe}$  and  $\text{CdSe}$  NCs inherit the size and morphology of their Cr-Se templates. The size- and structure-dependent optical and electronic properties allow for diverse applications.

The metal selenide NCs discussed above hold significant potential for a wide array of applications, such as catalysis, energy storage, and optoelectronics. As a proof-of-concept demonstration, we aimed to leverage the magnetic properties of  $\text{Cr}_2\text{Se}_3$ ,  $\text{Cr}_3\text{Se}_4$ , and  $\text{CuCrSe}_2$ . These colloidal Cr-Se-based nanomaterials offer two immediate advantages. Firstly, the colloidal approach provides a pathway for the scalable synthesis of magnetic materials for solution-processed thin film devices, with carrier mobility comparable to bulk materials achievable through ligand exchange and annealing processes. Secondly, these NCs with their standalone nature hold promise as templates for incorporating foreign cations/anions to alter magnetic ordering or enhance the Curie temperature. To explore these advantages, the magnetic properties of the synthesized Cr-Se and  $\text{CuCrSe}_2$  NCs with different morphologies were investigated by PPMS measurements. Fig. 4a and b display the hysteresis loops and ZFC and FC magnetization results for  $\text{Cr}_2\text{Se}_3$  nanoplatelets, indicating a ferromagnetic behavior with a  $T_C$  of  $\sim 80$  K and exhibiting paramagnetic ordering at higher temperature, consistent with the previous reports.<sup>2,51,52</sup> Compared to  $\text{Cr}_2\text{Se}_3$ , the  $\text{Cr}_3\text{Se}_4$  nanoplates displayed a slightly lower  $T_C$  of  $\sim 50$  K but a higher coercivity (Fig. 4c and d). The



**Fig. 4** Magnetic measurements of Cr-Se and  $\text{CuCrSe}_2$  NCs.  $M-H$  and  $M-T$  curves of (a and b)  $\text{Cr}_2\text{Se}_3$  nanoplatelets, (c and d)  $\text{Cr}_3\text{Se}_4$  nanoplates, (e and f)  $\text{Cr}_3\text{Se}_4$  nanoflowers, and (g and h)  $\text{CuCrSe}_2$  nanoplates obtained through CE with  $\text{Cr}_3\text{Se}_4$  nanoplates.





magnetic behavior is different from the recently reported  $\text{Cr}_3\text{Se}_4$  flakes, possibly due to the size effect of the sample in this work.<sup>2</sup> In single-domain particles, coercivity increases with size due to magnetic anisotropy scaling with volume. However, multi-domain transitions at larger sizes reduce coercivity *via* domain wall-mediated reversal.<sup>53–55</sup>  $\text{Cr}_3\text{Se}_4$  nanoplates likely remain single-domain, favoring coherent magnetization reversal, while smaller  $\text{Cr}_2\text{Se}_3$  may approach pseudo-single-domain states with lower energy barriers for switching. Surface pinning also dominates coercivity in nanostructures.  $\text{Cr}_3\text{Se}_4$ 's nanoplate geometry (high aspect ratio, distinct facets) enhances surface anisotropy, stabilizing moments against reversal. In contrast,  $\text{Cr}_2\text{Se}_3$ 's compact platelets exhibit weaker pinning, reducing coercivity. This is consistent with the previous reports that shape-dependent surface anisotropy aligns with studies on anisotropic NCs.<sup>56</sup> Moreover,  $\text{Cr}_3\text{Se}_4$ 's monoclinic structure likely hosts stronger magnetocrystalline anisotropy than trigonal  $\text{Cr}_2\text{Se}_3$ , amplified by stoichiometry-driven exchange interactions.

For  $\text{Cr}_3\text{Se}_4$  nanoflowers, the linear relationship in  $M$ - $H$  loops at both room and low temperatures (Fig. 4e) as well as almost zero coercivity revealed their paramagnetic nature. The pronounced increase in magnetization at low temperatures can be elucidated by dipolar interactions prevailing over thermal energy, thereby promoting moment alignment and consequently augmenting magnetization (Fig. 4f). The discrepancies may arise from the distinct exchange interactions that lead to long-range magnetic order,<sup>2</sup> potentially attributed to the uniformity in sample thickness and the presence of defects during synthesis. While  $\text{Cr}_3\text{Se}_4$  exhibits paramagnetic ordering overall, the ZFC/FC splitting in nanoplates ( $\sim 50$  K) *versus* overlapping curves in nanoflowers stems from morphology-dependent spin dynamics. In nanoplates, the large planar geometry enhances surface spin density and disorder.<sup>57</sup> At low temperatures, these surface spins experience frustration due to competing interactions,<sup>58</sup> leading to a glass-like state that manifests by the irreversibility in ZFC/FC. Conversely, nanoflowers' 3D branched structure promotes stronger interparticle coupling, which suppresses surface spin effects and stabilizes faster spin relaxation, resulting in reversible paramagnetic behavior.<sup>59</sup> The magnetic properties of Cr-Se NCs are intricately linked to the valence states of Cr cations and their corresponding electron spin configurations.<sup>60</sup> In  $\text{Cr}_2\text{Se}_3$  and  $\text{Cr}_3\text{Se}_4$ , the valence states of Cr cations ( $\text{Cr}^{2+}$ ,  $\text{Cr}^{3+}$ ) play a critical role in determining spin-orbit coupling, which influences the magnetic ordering. Specifically,  $\text{Cr}^{3+}$  ( $d^3$  configuration) typically exhibits a high-spin state, favoring ferromagnetic coupling *via* superexchange/double-exchange interactions.<sup>2</sup> In contrast,  $\text{Cr}^{2+}$  ( $d^4$  configuration) can exhibit varying spin states depending on the crystal field splitting, which may lead to complex magnetic behaviors such as antiferromagnetism or spin-glass states.<sup>51,61</sup> In our study, the observed ferromagnetic behavior in  $\text{Cr}_2\text{Se}_3$  nanoplatelets ( $T_C \sim 80$  K) and  $\text{Cr}_3\text{Se}_4$  nanoplates ( $T_C \sim 50$  K) can be attributed to the dominance of  $\text{Cr}^{3+}$  cations, which promotes strong ferromagnetic coupling. The higher coercivity in  $\text{Cr}_3\text{Se}_4$  nanoplates

compared to  $\text{Cr}_2\text{Se}_3$  suggests enhanced spin alignment and magnetic anisotropy, likely due to differences in the  $\text{Cr}^{2+}/\text{Cr}^{3+}$  ratio and the resulting electron spin configurations. This is consistent with previous reports that highlight the role of valence states in modulating magnetic properties.<sup>2,61</sup> The paramagnetic nature of  $\text{Cr}_3\text{Se}_4$  nanoflowers, as evidenced by the linear  $M$ - $H$  loops and negligible coercivity, further supports the influence of valence states. However, the absence of long-range magnetic order in  $\text{Cr}_3\text{Se}_4$  nanoflowers may arise from the presence of  $\text{Cr}^{2+}$  cations, which introduce competing exchange interactions and potential spin frustration.

Upon incorporation with  $\text{Cu}^+$ ,  $\text{CuCrSe}_2$  exhibited the anti-ferromagnetic behavior with a Néel temperature ( $T_N$ ) of  $\sim 45$  K (Fig. 4g and h), consistent with the previously reported magnetic properties of  $\text{CuCrSe}_2$ .<sup>62,63</sup> This arises from the antiferromagnetically coupled  $\text{CrSe}_2$  layers between interlayers in  $\text{CuCrSe}_2$ , resulting in an antiferromagnetic ground state. Recent findings indicated the observation of ferromagnetism in  $\text{CuCrSe}_2$  when the thickness is reduced to a few layers. Work is currently underway to explore the novel magnetic properties of Cr-Se nanoplates with precisely controlled thickness.

## Conclusions

In summary, we successfully synthesized a series of colloidal Cr-Se NCs with different crystal phases and morphologies through manipulating the Cr precursor and reaction temperature, producing  $\text{Cr}_2\text{Se}_3$  nanoplatelets and  $\text{Cr}_3\text{Se}_4$  nanoplates and nanoflowers. We show that, in addition to the commonly used copper selenide, the Cr-Se class of nanomaterials can serve as the template for the CE reactions to create diverse metal selenide NCs. By reacting the Cr-Se templates with different cations ( $\text{Cu}^+$ ,  $\text{Ag}^+$ ,  $\text{Zn}^{2+}$ ,  $\text{Cd}^{2+}$ , and  $\text{In}^{3+}$ ), we demonstrated that the outcomes of the CE reactions were strongly influenced by the crystal structure of the templates, which was primarily attributed to the lattice ordering in the Cr-Se template NCs, wherein monodisperse ultrathin InSe nanosheets can be produced. We further explored the magnetic properties of Cr-Se and  $\text{CuCrSe}_2$  NCs. These findings underscore the significant potential of Cr-Se based NCs in creating novel metal selenide derivatives and exploring their unique properties.

## Data availability

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

## Conflicts of interest

There are no conflicts to declare.





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## References

- 1 Y. Zhang, J. Chu, L. Yin, T. A. Shifa, Z. Cheng, R. Cheng, F. Wang, Y. Wen, X. Zhan, Z. Wang and J. He, *Adv. Mater.*, 2019, **31**, 1900056.
- 2 F. Cui, K. He, S. Wu, H. Zhang, Y. Lu, Z. Li, J. Hu, S. Pan, L. Zhu, Y. Huan, B. Li, X. Duan, Q. Ji, X. Zhao and Y. Zhang, *ACS Nano*, 2024, **18**, 6276–6285.
- 3 J. Zhang, Y. Xiao, K. Li, Y. Chen, S. Liu, W. Luo, X. Liu, S. Liu, Y. Wang, S. Y. Li and A. Pan, *Nanoscale*, 2024, **16**, 8028–8035.
- 4 S. Li, J. Tan, Y. Sun, J. Liu, H. Nong, L. He, Y. Zhang, J. Wang and B. Liu, *Adv. Funct. Mater.*, 2024, **34**, 2403453.
- 5 B. Li, Z. Wan, C. Wang, P. Chen, B. Huang, X. Cheng, Q. Qian, J. Li, Z. Zhang, G. Sun, B. Zhao, H. Ma, R. Wu, Z. Wei, Y. Liu, L. Liao, Y. Ye, Y. Huang, X. Xu, X. Duan, W. Ji and X. Duan, *Nat. Mater.*, 2021, **20**, 818–825.
- 6 M. Liu, J. Gou, Z. Liu, Z. Chen, Y. Ye, J. Xu, X. Xu, D. Zhong, G. Eda and A. T. S. Wee, *Nat. Commun.*, 2024, **15**, 1765.
- 7 M. Nasilowski, B. Mahler, E. Lhuillier, S. Ithurria and B. Dubertret, *Chem. Rev.*, 2016, **116**, 10934–10982.
- 8 J. H. Han, M. Kwak, Y. Kim and J. Cheon, *Chem. Rev.*, 2018, **118**, 6151–6188.
- 9 H. Chen, Z. Chen, B. Ge, Z. Chi, H. Chen, H. Wu, C. Cao and X. Duan, *Chem. Mater.*, 2017, **29**, 10019–10026.
- 10 L. Hu, R. Zhang and Q. Chen, *Nanoscale*, 2014, **6**, 14064–14105.
- 11 F. Wang and X. Wang, *Nanoscale*, 2014, **6**, 6398–6414.
- 12 J. Wang, B. Wang, Y. Wang, R. Yang, L. Wang, F. Wang, X. Xu and Y. Liu, *J. Mater. Chem. C*, 2024, **12**, 7725–7731.
- 13 F. Wang, H. Yang, H. Zhang, J. Zhou, J. Wang, L. Hu, D. J. Xue and X. Xu, *Nano Lett.*, 2021, **21**, 7684–7690.
- 14 H. Yang, H. Zhang, L. Guo, W. Yang, Y. Wu, J. Wang, X. Li, H. Du, B. Peng, Q. Liu, F. Wang, D. J. Xue and X. Xu, *Nano Lett.*, 2024, **24**, 10519–10526.
- 15 H. Yang, F. Wang, H. Zhang, L. Guo, L. Hu, L. Wang, D. J. Xue and X. Xu, *J. Am. Chem. Soc.*, 2020, **142**, 4438–4444.
- 16 Q. Tu, Q. Fu, C. Tang, Y. Xiao, B. Cheng and S. Lei, *Cryst. Growth Des.*, 2022, **22**, 4277–4287.
- 17 L. T. De and L. Manna, *Chem. Rev.*, 2016, **116**, 10852–10887.
- 18 C. Coughlan, M. Ibáñez, O. Dobrozhan, A. Singh, A. Cabot and K. M. Ryan, *Chem. Rev.*, 2017, **117**, 5865–6109.
- 19 Y. Liu, M. Liu and M. T. Swihart, *J. Phys. Chem. C*, 2017, **121**, 13435–13447.
- 20 M. Liu, Y. Liu, B. Gu, X. Wei, G. Xu, X. Wang, M. T. Swihart and K. T. Yong, *Chem. Soc. Rev.*, 2019, **48**, 4950–4965.
- 21 T. T. Zhuang, F. J. Fan, M. Gong and S. H. Yu, *Chem. Commun.*, 2012, **48**, 9762–9764.
- 22 H. Li, M. Zanella, A. Genovese, M. Povia, A. Falqui, C. Giannini and L. Manna, *Nano Lett.*, 2011, **11**, 4964–4970.
- 23 C. H. Ho and X. R. Lai, *ACS Appl. Electron. Mater.*, 2019, **1**, 370–378.
- 24 Q. Yun, Y. Ge, B. Huang, Q. Wa and H. Zhang, *Acc. Chem. Res.*, 2023, **56**, 1780–1790.
- 25 W. Li, R. Zamani, M. Ibáñez, D. Cadavid, A. Shavel, J. R. Morante, J. Arbiol and A. Cabot, *J. Am. Chem. Soc.*, 2013, **135**, 4664–4667.
- 26 X. Y. Gan, R. Sen and J. E. Millstone, *J. Am. Chem. Soc.*, 2021, **143**, 8137–8144.
- 27 W. Huang, L. Gan, H. Li, Y. Ma and T. Zhai, *Chem. – Eur. J.*, 2018, **24**, 15678–15684.
- 28 G. Almeida, S. Dogan, G. Bertoni, C. Giannini, R. Gaspari, S. Perissinotto, R. Krahne, S. Ghosh and L. Manna, *J. Am. Chem. Soc.*, 2017, **139**, 3005–3011.
- 29 K. H. Park, K. Jang, S. Kim, H. J. Kim and S. U. Son, *J. Am. Chem. Soc.*, 2006, **128**, 14780–14781.
- 30 J. Lauth, F. E. S. Gorris, M. K. Samadi, T. Chassé, W. Friedrich, V. Lebedeva, A. Meyer, C. Klinke, A. Kornowski, M. Scheele and H. Weller, *Chem. Mater.*, 2016, **28**, 1728–1736.
- 31 J. Peng, Y. Su, H. Lv, J. Wu, Y. Liu, M. Wang, J. Zhao, Y. Guo, X. Wu, C. Wu and Y. Xie, *Adv. Mater.*, 2023, **35**, 2209365.
- 32 P. Wang, Y. Zhao, R. Na, W. Dong, J. Duan, Y. Cheng, B. Xu, D. Kong, J. Liu, S. Du, C. Zhao, Y. Yang, L. Lv, Q. Hu, H. Ai, Y. Xiong, V. S. Stolyarov, S. Zheng, Y. Zhou, F. Deng and J. Zhou, *Adv. Mater.*, 2024, **36**, 2400655.
- 33 Z. Sun, Y. Su, A. Zhi, Z. Gao, X. Han, K. Wu, L. Bao, Y. Huang, Y. Shi, X. Bai, P. Cheng, L. Chen, K. Wu, X. Tian, C. Wu and B. Feng, *Nat. Commun.*, 2024, **15**, 4252.
- 34 Y. Liu, D. Yin and M. T. Swihart, *Chem. Mater.*, 2018, **30**, 8089–8098.
- 35 Y. Liu, D. Yin and M. T. Swihart, *Chem. Mater.*, 2018, **30**, 1399–1407.
- 36 Y. Liu, M. Liu and M. T. Swihart, *J. Am. Chem. Soc.*, 2017, **139**, 18598–18606.
- 37 Y. Liu, M. Liu, D. Yin, L. Qiao, Z. Fu and M. T. Swihart, *ACS Nano*, 2018, **12**, 7803–7811.
- 38 W. Feng, W. Zheng, W. Cao and P. Hu, *Adv. Mater.*, 2014, **26**, 6587–6593.
- 39 S. Sucharitakul, N. J. Goble, U. R. Kumar, R. Sankar, Z. A. Bogorad, F. C. Chou, Y. T. Chen and X. P. A. Gao, *Nano Lett.*, 2015, **15**, 3815–3819.
- 40 D. A. Bandurin, A. V. Tyurnina, G. L. Yu, A. Mishchenko, V. Zólyomi, S. V. Morozov, R. K. Kumar, R. V. Gorbachev, Z. R. Kudrynskyi, S. Pezzini, Z. D. Kovalyuk, U. Zeitler, K. S. Novoselov, A. Patané, L. Eaves, I. V. Grigorieva, V. I. Fal'ko, A. K. Geim and Y. Cao, *Nat. Nanotechnol.*, 2017, **12**, 223–227.



- 41 Z. Q. Zheng, J. D. Yao and G. W. Yang, *J. Mater. Chem. C*, 2016, **4**, 8094–8103.
- 42 S. Chen, X. Liu, X. Qiao, X. Wan, K. Shehzad, X. Zhang, Y. Xu and X. Fan, *Small*, 2017, **13**, 1604033.
- 43 W. Feng, W. Zheng, F. Gao, X. Chen, G. Liu, T. Hasan, W. Cao and P. Hu, *Chem. Mater.*, 2016, **28**, 4278–4283.
- 44 H. Liu, X. Shi, F. Xu, L. Zhang, W. Zhang, L. Chen, Q. Li, C. Uher, T. Day and G. J. Snyder, *Nat. Mater.*, 2012, **11**, 422–425.
- 45 J. Zhu, Q. Li, L. Bai, Y. Sun, M. Zhou and Y. Xie, *Chem. – Eur. J.*, 2012, **18**, 13213–13221.
- 46 Y. Su, G. Li, Z. Guo, Y. Y. Li, Y. X. Li, X. J. Huang and J. H. Liu, *ACS Appl. Nano Mater.*, 2018, **1**, 245–253.
- 47 D. Li, B. L. Zhang, H. W. Ming, L. Wang, Y. Zu and X. Y. Qin, *ACS Appl. Mater. Interfaces*, 2021, **13**, 34543–34549.
- 48 S. Huang, T. R. Wei, H. Chen, J. Xiao, M. Zhu, K. Zhao and X. Shi, *ACS Appl. Mater. Interfaces*, 2021, **13**, 60192–60199.
- 49 Y. Wang, Y. Luo, X. Kong, T. Wu, Y. Lin, Z. Chen and S. Wang, *Nanoscale*, 2025, **17**, 1764–1789.
- 50 W. Zhu, Z. Lin, X. Zhang, W. Wang and Y. Li, *Nanoscale*, 2022, **14**, 11210–11217.
- 51 S. Luo, X. Zhu, H. Liu, S. Song, Y. Chen, C. Liu, W. Zhou, C. Tang, G. Shao, Y. Jin, J. Guan, V. C. Tung, H. Li, X. Chen, F. Ouyang and S. Liu, *Chem. Mater.*, 2022, **34**, 2342–2351.
- 52 X. Zhu, H. Liu, L. Liu, L. Ren, W. Li, L. Fang, X. Chen, L. Xie, Y. Jing, J. Chen, S. Liu, F. Ouyang, Y. Zhou and X. Xiong, *Chem. Mater.*, 2021, **33**, 3851–3858.
- 53 A. G. Kolhatkar, A. C. Jamison, D. Litvinov, R. C. Willson and T. R. Lee, *Int. J. Mol. Sci.*, 2013, **14**, 15977–16009.
- 54 P. Guardia, A. Labarta and X. Batlle, *J. Phys. Chem. C*, 2011, **115**, 390–396.
- 55 J. Msomi and T. Moyo, *J. Magn. Magn. Mater.*, 2009, **321**, 1246–1250.
- 56 Q. Song and Z. J. Zhang, *J. Am. Chem. Soc.*, 2004, **126**, 6164–6168.
- 57 W. Li, Y. Wang, X. Y. Cui, S. Yu, Y. Li, Y. Hu, M. Zhu, R. Zheng and S. P. Ringer, *ACS Appl. Mater. Interfaces*, 2018, **10**, 19235–19247.
- 58 R. Fuglsby, P. Kharel, W. Zhang, S. Valloppilly, Y. Huh and D. J. Sellmyer, *J. Appl. Phys.*, 2015, **117**, 17D115.
- 59 J. Jang, P. K. Srivastava, M. Joe, S. G. Jung, T. Park, Y. Kim and C. Lee, *ACS Nano*, 2025, **19**, 999–1006.
- 60 L. Lin, P. Su, Y. Han, Y. Xu, Q. Ni, X. Zhang, P. Xiong, Z. Sun, G. Sun and X. Chen, *eScience*, 2025, **5**, 100264.
- 61 X. Liu, Y. Gebredingle, X. Guo, F. Zhang and N. Kim, *Adv. Funct. Mater.*, 2024, **34**, 2316834.
- 62 G. C. Tewari, T. S. Tripathi, H. Yamauchi and M. Karppinen, *Mater. Chem. Phys.*, 2014, **145**, 156–161.
- 63 G. C. Tewari, M. Karppinen and A. K. Rastogi, *J. Solid State Chem.*, 2013, **198**, 108–113.

