

# Chemical Science

Volume 13  
Number 35  
21 September 2022  
Pages 10163–10584

rsc.li/chemical-science



ISSN 2041-6539

## EDGE ARTICLE

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Cite this: *Chem. Sci.*, 2022, 13, 10260

All publication charges for this article have been paid for by the Royal Society of Chemistry

# Wide bandgaps and strong SHG responses of hetero-oxyfluorides by dual-fluorination-directed bandgap engineering†

Yilei Hu,<sup>‡a</sup> Xingxing Jiang,<sup>‡b</sup> Tianhui Wu,<sup>‡a</sup> Yanyan Xue,<sup>‡a</sup> Chao Wu,<sup>ID \*a</sup> Zhipeng Huang,<sup>ID a</sup> Zheshuai Lin,<sup>ID b</sup> Jun Xu,<sup>a</sup> Mark G. Humphrey,<sup>ID c</sup> and Chi Zhang,<sup>ID \*a</sup>

A wide bandgap is an essential requirement for a nonlinear optical (NLO) material. However, it is very challenging to simultaneously engineer a wide bandgap and a strong second-harmonic generation (SHG) response, particularly in NLO materials containing second-order Jahn–Teller (SOJT) distorted units. Herein, we employ a bandgap engineering strategy that involves the dual fluorination of two different types of SOJT distorted units to realize remarkably wide bandgaps in the first examples of 5d<sup>0</sup>-transition metal (TM) fluoriodates. Crystalline A<sub>2</sub>WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>) (A = Rb (RWOFl) and Cs (CWOFl)) exhibit the largest bandgaps yet observed in d<sup>0</sup>-TM iodates (4.42 (RWOFl) and 4.29 eV (CWOFl)), strong phase-matching SHG responses of 3.8 (RWOFl) and 3.5 (CWOFl) × KH<sub>2</sub>PO<sub>4</sub>, and wide optical transparency windows. Computational studies have shown that the excellent optical responses result from synergism involving the two fluorinated SOJT distorted units ([WO<sub>3</sub>F<sub>3</sub>]<sup>3−</sup> and [IO<sub>2</sub>F<sub>2</sub>]<sup>−</sup>). This work provides not only an efficient strategy for bandgap modulation of NLO materials, but also affords insight into the relationship between the electronic structure of the various fluorinated SOJT distorted units and the optical properties of crystalline materials.

Received 14th April 2022  
Accepted 25th April 2022

DOI: 10.1039/d2sc02137d

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## Introduction

Nonlinear optical (NLO) materials, exemplified by commercial β-BaB<sub>2</sub>O<sub>4</sub> (β-BBO), LiB<sub>3</sub>O<sub>5</sub> (LBO), KH<sub>2</sub>PO<sub>4</sub> (KDP), KTiOPO<sub>4</sub> (KTP) and AgGaS<sub>2</sub> (AGS),<sup>1</sup> can effectively expand the spectral range of lasers, and may therefore serve as key materials for the all-solid-state laser devices that are widely applied in modern laser technologies such as photolithography, spectral analysis, tissue imaging, and environmental monitoring.<sup>2</sup> A high-performance NLO material should not only possess non-centrosymmetry, but it should also exhibit a large second-harmonic generation (SHG) coefficient, a wide bandgap, sufficient birefringence, and good physicochemical stability.<sup>3</sup> Unfortunately, two of the key optical properties, a wide bandgap and a strong SHG coefficient, are two often-competing optical parameters and difficult to

engineer simultaneously in a material, owing to their contrasting microstructural requirements.<sup>4</sup>

Modulation of SHG responses can be achieved by incorporating second-order Jahn–Teller (SOJT) distorted cations,<sup>5–7</sup> such as octahedrally coordinated d<sup>0</sup>-transition metal (TM) cations (*e.g.*, Ti<sup>4+</sup>, V<sup>5+</sup>, Nb<sup>5+</sup>, Mo<sup>6+</sup>, *etc.*)<sup>5</sup> or stereochemically-active lone-pair cations (*e.g.*, Se<sup>4+</sup>, Te<sup>4+</sup>, and I<sup>5+</sup>, *etc.*)<sup>6</sup> although, in general, these microstructural building units promote a red-shift in the absorption edge. A structurally related strategy involves fluorination of the TM cation-centered oxyanions<sup>8</sup> in order to blue-shift the absorption edge, but the SHG responses of the resultant materials are usually limited (<1.0 KDP) owing to the often-antiparallel arrangements of the fluorinated TM oxyanions. A complementary approach relies on combination of two types of oxyanions containing SOJT distorted cations; this has been demonstrated by the successful syntheses of a variety of hetero-oxyanion materials with strong SHG responses,<sup>9,10</sup> such as ASe<sub>2</sub>V<sub>3</sub>O<sub>12</sub> (A = Rb, Tl),<sup>9a</sup> A<sub>2</sub>-(MoO<sub>3</sub>)<sub>3</sub>(SeO<sub>3</sub>)<sub>2</sub> (A = Rb, Tl),<sup>9b</sup> MgTeMoO<sub>6</sub>,<sup>9c</sup> AMoO<sub>3</sub>(IO<sub>3</sub>) (A = Rb, Cs, NH<sub>4</sub>) (A = Rb, Cs, NH<sub>4</sub>),<sup>9d,e</sup> K(VO)<sub>2</sub>O<sub>2</sub>(IO<sub>3</sub>)<sub>3</sub>,<sup>9f</sup> and [C(NH<sub>2</sub>)<sub>3</sub>]<sub>2</sub>Mo<sub>2</sub>O<sub>5</sub>(IO<sub>3</sub>)<sub>4</sub>·2H<sub>2</sub>O.<sup>9g</sup> However, the bandgaps of these materials are relatively narrow owing to the existence of d-orbital electrons in the TM cations, which significantly restricts the optical transparency range in the short-wavelength region and reduces the laser damage threshold.

<sup>a</sup>China–Australia Joint Research Center for Functional Molecular Materials, School of Chemical Science and Engineering, Tongji University, Shanghai 200092, China. E-mail: chizhang@tongji.edu.cn; wuc@tongji.edu.cn

<sup>b</sup>Technical Institute of Physics and Chemistry, Chinese Academy of Sciences, Beijing 100190, China

<sup>c</sup>Research School of Chemistry, Australian National University, Canberra, ACT 2601, Australia

† Electronic supplementary information (ESI) available. CCDC 2164004 and 2164005. For ESI and crystallographic data in CIF or other electronic format see <https://doi.org/10.1039/d2sc02137d>

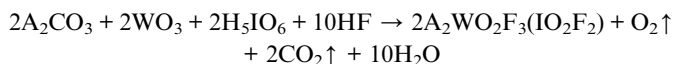
‡ Y. L. Hu, X. X. Jiang, T. H. Wu and Y. Y. Xue contributed equally to the work.

Nevertheless, of the possible chemical systems, assembly of hetero-oxyanions may still offer an ideal platform for optical property modulation because of their multiple tunable micro-structural building units.<sup>10</sup> The two mutually exclusive optical properties of bandgap and SHG response are intimately related to the electronic band structure of the material, which can be micro-controlled by the intrinsic properties and spatial arrangements of the building units.<sup>11</sup> In this study, we propose a general bandgap engineering approach based on fluorinating two types of oxyanions, both containing SOJT cations. In this approach, a judicious assembly of fluorine-rich units containing heavy cations addresses the problematic competition between bandgap and SHG response in hetero-oxyanion materials. To implement this strategy, heavy 5d<sup>0</sup>-TM octahedra (e.g., Hf<sup>4+</sup>, Ta<sup>5+</sup> and W<sup>6+</sup>) are preferred to conventional 3d<sup>0</sup>/4d<sup>0</sup>-TM octahedra of the same family because heavy 5d<sup>0</sup>-TM octahedra are anticipated to form less-covalent bonds with oxygen/fluorine owing to their lower effective electronegativity (referring to the ease of oxygen-to-metal charge transfer for the heavy 5d<sup>0</sup>-TM octahedra);<sup>12</sup> this will increase the bandgap of the resultant materials. The simultaneous fluorination of two different types of oxyanions in the one structure (i.e. d<sup>0</sup>-TM-centered octahedra and non-metal-centered polyhedra) can induce further significant differences in the band structures and increase the bandgap owing to the highly electronegative fluorine, an outcome observed in recently reported fluorooxoborates<sup>13</sup> and fluorophosphates.<sup>14</sup> Furthermore, fluorine acting as chemical “scissors” can effectively reduce the dimensionality of the structure, which may be highly beneficial for the enhancement of bandgap.<sup>15</sup> Finally, large macroscopic polarization (corresponding to a large SHG effect) may be achieved by stacking the oxyfluorine anions in an additive mode. To demonstrate the effectiveness of our proposed strategy, we disclose the successful synthesis of the first examples of 5d<sup>0</sup>-TM fluoroiodates A<sub>2</sub>WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>) (A = Rb (RWOFI), Cs (CWOFI)). Their unique lambda (Λ)-shaped [WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>)]<sup>2-</sup> hetero-oxyfluorine anions not only drive the formation of the polar structures that exhibit strong phase-matching SHG responses of 3.8 (RWOFI) and 3.5 (CWOFI) times that of KDP, sufficient birefringences, and wide optical transparency windows from the ultraviolet to the mid-infrared, but more importantly induce very wide bandgaps of 4.42 eV (RWOFI) and 4.29 eV (CWOFI) – both among the largest bandgaps for d<sup>0</sup>-TM iodates. Herein we report their syntheses, crystal structures, electronic structures, and optical properties, as well as structure–property correlations employing a combination of experiments and first-principles calculations.

## Results and discussion

Although considerable research pursuing non-centrosymmetric iodates containing 3d<sup>0</sup>/4d<sup>0</sup>-TM cations susceptible to the SOJT effect has been undertaken, there has thus far been only one 5d<sup>0</sup>-TM iodate reported (K<sub>5</sub>(W<sub>3</sub>O<sub>9</sub>F<sub>4</sub>)(IO<sub>3</sub>),<sup>16</sup> containing the W<sup>6+</sup> cation). The synthetic difficulty lies in the fact that tungsten is a refractory metal, and is extremely difficult to dissolve in common solvents. Unlike the previous synthetic method, in this

study hydrofluoric acid was used not only as the solvent but also as a fluorine source for the syntheses of hetero-oxyfluorides A<sub>2</sub>WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>) based on the following reaction equation:



Our efforts to synthesize homologues with other alkali metal cations (e.g., Na<sup>+</sup>, K<sup>+</sup>) were unsuccessful, possibly due to the large differences in ionic radii among these alkali metal cations (0.102, 0.138, 0.152, and 0.167 nm for Na<sup>+</sup>, K<sup>+</sup>, Rb<sup>+</sup>, and Cs<sup>+</sup>, respectively).<sup>17a</sup> The purities of the two hetero-oxyfluorides were confirmed by comparison of their experimental PXRD patterns with simulated patterns derived from single-crystal X-ray diffraction data (Fig. S1†). The energy dispersive X-ray spectroscopy analyses reveal element distribution maps for RWOFI and CWOFI with molar ratios of 9.38 : 5.01 : 4.02 : 26.55 and 8.75 : 4.78 : 4.54 : 25.11 for Rb/Cs, W, I and F, respectively, which are consistent with the molar ratios determined from single-crystal X-ray diffraction (Fig. S2†).

A<sub>2</sub>WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>) (A = Rb, Cs) are the first examples of 5d<sup>0</sup>-TM fluoroiodates. They are isostructural and crystallize in the same non-centrosymmetric and polar orthorhombic space group, *Cmc*2<sub>1</sub> (no. 36), so only the structure of CWOFI is described in detail here (Tables S1–S5†). Its structure is composed of zero-dimensional (0D) [WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>)]<sup>2-</sup> hetero-oxyfluorine anions and Cs<sup>+</sup> cations (Fig. 1b). In the [WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>)]<sup>2-</sup> hetero-oxyfluorine anion, the unique I<sup>5+</sup> cation is coordinated by two O atoms and two F atoms, forming an unsymmetrical [IO<sub>2</sub>F<sub>2</sub>] tetrahedron (Fig. 1a). The two I–F bond lengths (1.974(8) Å) are longer than the two I–O lengths (1.790(17) and 1.759(14) Å), and the F–I–F bond angle is close to linear (177.8(6)°). Each W<sup>6+</sup> cation is octahedrally coordinated with two terminal O atoms, three terminal F atoms, and one bridging O atom from an [IO<sub>2</sub>F<sub>2</sub>]<sup>-</sup> group bound in a unidentate fashion (Fig. 1a). The W–F and W–O bond distances lie in the ranges 1.733(16)–1.953(8) and 1.800(10)–2.263(17) Å, respectively. One [WO<sub>2</sub>F<sub>3</sub>] octahedron further connects with one [IO<sub>2</sub>F<sub>2</sub>] tetrahedron by sharing O atoms, forming the Λ-shaped [WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>)]<sup>2-</sup> hetero-oxyfluorine anion (Fig. 1a) with a W–O–I angle of 140.78°; this differs from other Λ-shaped iodate-containing species such as *cis*-[ZrF<sub>6</sub>(IO<sub>3</sub>)<sub>2</sub>],<sup>17a</sup> *cis*-[VO<sub>2</sub>F<sub>2</sub>(IO<sub>3</sub>)<sub>2</sub>],<sup>17b</sup> and [Ga(IO<sub>3</sub>)<sub>2</sub>F<sub>4</sub>].<sup>17c</sup> The [WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>)]<sup>2-</sup> hetero-oxyfluorine anion is the first example of a TM iodate containing two different types of heavy atom-containing SOJT distorted cations. The Cs<sup>+</sup> cations connect the Λ-shaped [WO<sub>2</sub>F<sub>3</sub>(IO<sub>2</sub>F<sub>2</sub>)]<sup>2-</sup> hetero-oxyfluorine anions, forming a 3D layer-piled structure along the *b*-axis (Fig. 1b).

The most critical structural feature of CWOFI, however, is the tungsten-centered oxyfluorine anion, which plays a key role in construction of the polar fluoroiodate. The W<sup>6+</sup> cation undergoes intra-octahedral distortion toward the terminal fluorine atom, a corner (C4) distortion, with one short [1.733(16) Å], one long [2.263(17) Å], and four normal [1.800(10)–1.953(8) Å] W–O/F bonds (Fig. S3†). The magnitude of the out-of-center distortion ( $\Delta d$ ) is 0.84, which corresponds to strong distortion ( $\Delta d > 0.8$ ).<sup>18</sup> The C4 distorted [WO<sub>3</sub>F<sub>3</sub>] octahedron is favorable





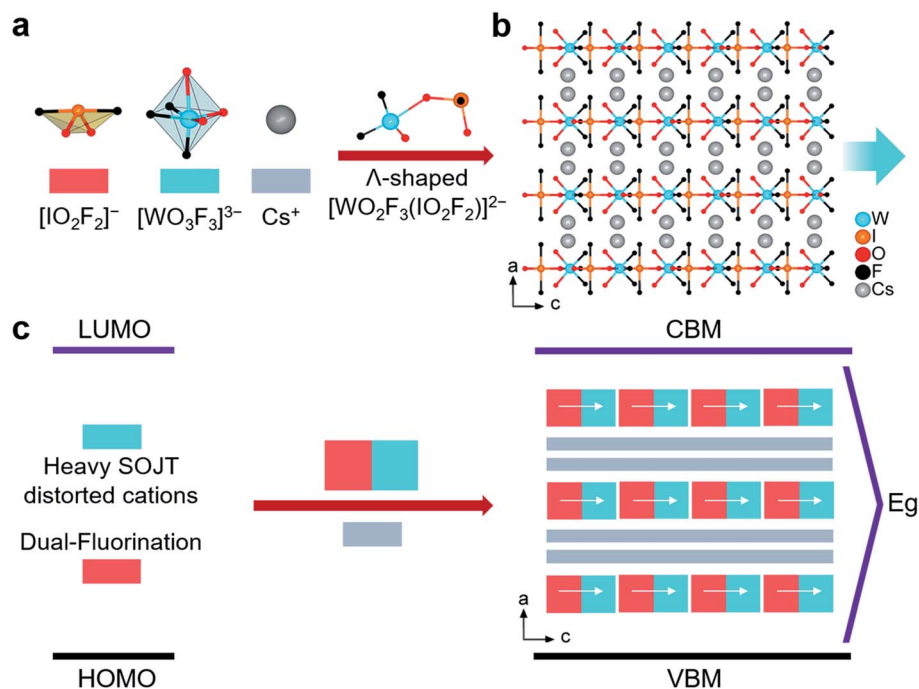


Fig. 1 (a) Structural building units in CWOFI. (b) Crystal structure of CWOFI viewed along the  $b$ -axis. The blue arrow in the panel represents the net dipole moment of the structure. (c) Structural building unit description of the arrangement of 0D  $[\text{WO}_2\text{F}_3(\text{IO}_2\text{F}_2)]^{2-}$  units and  $\text{Cs}^+$  cations. The Cs–O/F bonds are omitted for clarity.

for creating  $\Lambda$ -shaped units with a fixed polarization orientation. The bridging O(2) with one long bond has the greater residual negative charge, based on BVS calculations (Table S5<sup>†</sup>), and is therefore predicted to bond in preference to the other O/F terminal ligands; it consequently links to the lone-pair cation  $\text{I}^{5+}$  to form the  $\Lambda$ -shaped  $[\text{WO}_2\text{F}_3(\text{IO}_2\text{F}_2)]^{2-}$  hetero-oxyfluorine anion. The  $\text{W}^{6+}$  cations are blocked from distorting toward a corner, owing to the unsymmetrical  $[\text{IO}_2\text{F}_2]^-$  oxyfluorine anions, the lone-pair cation  $\text{I}^{5+}$  serving to reinforce the direction of the intra-octahedral distortion. The polar orientation of the  $\Lambda$ -shaped  $[\text{WO}_2\text{F}_3(\text{IO}_2\text{F}_2)]^{2-}$  hetero-oxyfluorine anion is fixed and it cannot rotate due to significant steric effects (Fig. 1a). As a result, the  $[\text{WO}_2\text{F}_3(\text{IO}_2\text{F}_2)]^{2-}$  hetero-oxyfluorine anions pack in the crystal structure with their polar orientations arranged in an additive fashion.

Our proposed bandgap engineering strategy involving dual fluorination of two different types of SOJT distorted units has afforded  $\text{A}_2\text{WO}_2\text{F}_3(\text{IO}_2\text{F}_2)$ , the structure of which bears a remarkable resemblance to those of conventional  $\text{d}^0$ -TM iodates. Their structures are constructed by judicious assembly of two different types of SOJT distorted units (Fig. 1), except for three following crucial differences. Firstly,  $3\text{d}^0/4\text{d}^0$ -TM cations (*e.g.*,  $\text{V}^{5+}$ ,  $\text{Ti}^{4+}$ ,  $\text{Nb}^{5+}$ , and  $\text{Mo}^{6+}$ ) are responsible for the bandgaps of the  $\text{d}^0$ -TM iodates, and these bandgaps are generally less than 3.8 eV<sup>19</sup> owing to the presence of d–d transitions. By contrast, the bandgaps of the hetero-oxyfluorides RWOFI and CWOFI are expected to blue-shift because the effective electronegativity<sup>12</sup> of the heavy  $5\text{d}^0$  tungsten cation is lower than that of the  $3\text{d}^0/4\text{d}^0$ -TM cations. Secondly, in stark contrast to conventional  $\text{d}^0$ -TM iodates, the dual fluorination of two different types of oxyanions

containing SOJT cations may afford structure-building oxyfluorine anions ( $[\text{WO}_3\text{F}_3]^{3-}$  and  $[\text{IO}_2\text{F}_2]^-$ ) with relatively wide HOMO–LUMO gaps,<sup>20</sup> and thereby potentially induce a wide bandgap in RWOFI and CWOFI. Thirdly, the abundant fluorine, acting as chemical “scissors”, induces formation of the 0D  $\Lambda$ -shaped  $[\text{WO}_2\text{F}_3(\text{IO}_2\text{F}_2)]^{2-}$  hetero-oxyfluorine anion, which favors the formation of polar structures, and is beneficial for strong SHG responses, owing to the crystal packing along the  $c$ -axis.

The optical transmission spectra of RWOFI and CWOFI single crystals suggest that both compounds possess broad optical transparency windows of 0.28–5.28 and 0.289–5.33  $\mu\text{m}$  (Fig. 2a, b, S4a, and b<sup>†</sup>) that cover the IR atmospheric window (3–5  $\mu\text{m}$ ). The optical bandgaps of RWOFI and CWOFI from their UV cutoff wavelengths are 4.42 and 4.29 eV. In contrast to  $\text{K}_5(\text{W}_3\text{O}_9\text{F}_4)(\text{IO}_3)$  (3.83 eV) with a similar composition, the bandgaps of RWOFI and CWOFI are significantly blue-shifted, which can be attributed to the combination of two different types of SOJT-distorted oxyfluorine anions as well as the reduced 0D structures. The bandgaps of RWOFI and CWOFI are the largest for  $\text{d}^0$ -TM iodates reported to date. IR absorption below  $1000\text{ cm}^{-1}$  was measured on crystalline powder samples of the two materials (Fig. S4c and d<sup>†</sup>). The absorption bands ranging from 460 to  $861\text{ cm}^{-1}$  can be assigned to the stretching and bending vibrations of the  $[\text{IO}_2\text{F}_2]$  tetrahedra, while the strong peaks between 950 and  $900\text{ cm}^{-1}$  belong to the vibration frequencies of the  $[\text{WO}_3\text{F}_3]$  octahedra. From the two-phonon approximation, the IR edges of RWOFI and CWOFI should be larger than 5  $\mu\text{m}$ , with the maximum absorption bands around



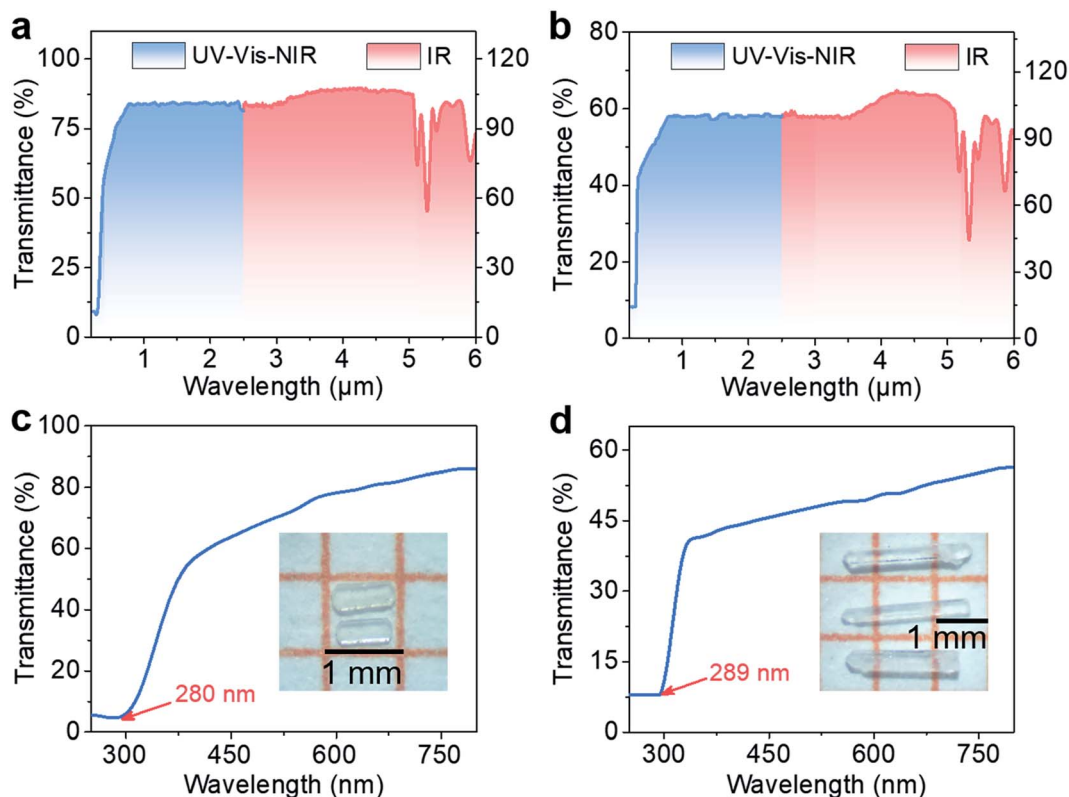


Fig. 2 Optical transparency windows (top) and UV-Vis-NIR transmission spectra (bottom) of RWO (a and c) and CWO (b and d). The insets show the crystals of RWO (c) and CWO (d) used for measurements.

977  $\text{cm}^{-1}$  and 966  $\text{cm}^{-1}$ , respectively, indicating that RWO and CWO are mid-IR transparent NLO materials.

The thermogravimetric analysis (Fig. S5†) reveals that RWO and CWO are stable at temperatures up to 288 °C, beyond which a two-step decomposition is observed. The first step, a sharp drop between 288–543 °C, corresponds to a weight loss of 28.20% and 25.31%, assigned to the loss of 0.5  $\text{I}_2$  and 1.5  $\text{F}_2$  molecules (28.72% and 25.28% for calculated values). The second step between 543 °C and 700 °C can be assigned to the loss of 1  $\text{F}_2$  molecule, with experimental weight losses of 5.53% and 4.93% (5.93% and 5.17% for calculated values).

Powder SHG measurements with 1064 nm laser radiation reveal that RWO and CWO are phase-matchable and display SHG signals of  $3.8\times$  and  $3.5\times$  KDP, respectively, in the particle size range 105–150 nm (Fig. 3). The measured SHG signals are significantly larger than those of previously reported crystals with single-fluorinated SOJT distorted units, such as  $\alpha$ - $\text{BaMoO}_2\text{F}_4$  ( $0.7\times$   $\alpha$ - $\text{SiO}_2$ ),<sup>8b</sup>  $\text{NaVOF}_4(\text{H}_2\text{O})$  ( $1\times$   $\alpha$ - $\text{SiO}_2$ ),<sup>8c</sup>  $\text{Rb}_2\text{-VO}(\text{O}_2)_2\text{F}$  ( $0.8\times$  KDP),<sup>8d</sup>  $\text{KNaNbOF}_5$  ( $3\times$  KDP),<sup>8a</sup>  $\text{KWO}_3\text{F}$  ( $3\times$  KDP),<sup>8e</sup> and  $\text{CsIO}_2\text{F}_2$  ( $3\times$  KDP).<sup>21</sup> Dipole moment calculations were performed on RWO and CWO by the bond valence method,<sup>22</sup> the results being summarized in Table S6.† The calculated net dipole moments of their unit cells are similar, consistent with their similar measured SHG results. The dipole moments of the  $[\text{WO}_3\text{F}_3]^{3-}$  and  $[\text{IO}_2\text{F}_2]^-$  oxyfluorine anions in the *a* and *b* direction nearly cancel, the vector sum of the dipole moments of both oxyanions effectively pointing along the *c*-axis.

The dipole moment of the  $[\text{IO}_2\text{F}_2]^-$  tetrahedron is larger than that of the  $[\text{WO}_3\text{F}_3]^-$  octahedron, indicating that the  $[\text{IO}_2\text{F}_2]^-$  oxyfluorine anions make the major contribution to the SHG responses.

Electronic structure calculations of RWO and CWO were undertaken to shed light on the origin of the linear and nonlinear optical properties. The band structures demonstrate that the two compounds are direct bandgap materials with a high efficiency of light utilization. The calculated bandgaps of RWO and CWO obtained from the band structures are 3.14 and 3.09 eV (Fig. S6†), respectively, smaller than the experimental results of 4.42 and 4.29 eV, a difference that can be ascribed to the discontinuity of exchange–correlation energy in DFT calculations. Examination of the partial and total density of states show that the I 5p and W 5d orbitals overlap with the O 2p and F 2p orbitals over the entire energy range (Fig. 4a and S7a†). The Rb 4p nonbonding orbitals in RWO are far from the Fermi level compared to the Cs 5p nonbonding orbitals in CWO. Thus, RWO shows a slightly larger bandgap, consistent with the trend in experimental results. The contribution to the valence band (VB) close to the Fermi level is essentially O 2p and F 2p in nature, with the latter located at a lower energy because of its larger electronegativity. Both I 5p and O 2p orbitals play key roles in the lower energy regions of the conduction band (CB), consistent with strong I–O/F and W–O/F bonds interactions.



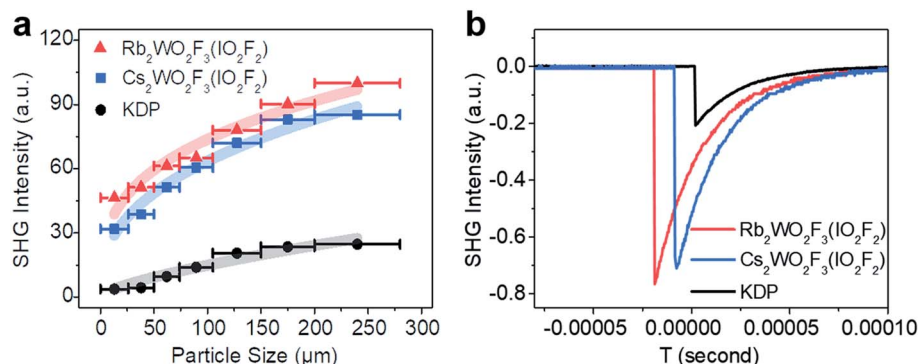


Fig. 3 (a) Phase-matching curves of RWOFI and CWOFI with 1064 nm laser radiation. The solid curves are guides for the eye and not fits to the data. (b) Oscilloscope traces of the SHG signals (105–150 μm) for powders of RWOFI and CWOFI at  $\lambda = 1064$  nm. KDP samples serve as the references.

To clearly demonstrate the influences of the two heavy oxyfluorine anions on the bandgaps, we firstly compare the electronic structures<sup>23</sup> of RWOFI and CWOFI with  $K_5(W_3O_9F_4)(IO_3)$ , which possess similar compositions but exhibit different configurations near the Fermi level.  $K_5(W_3O_9F_4)(IO_3)$  has isolated  $[IO_3]^-$  oxyanions with three terminal oxygen atoms that manifest as non-bonding O 2p orbitals in the electronic structure, while most oxygen atoms in the  $[IO_2F_2]^-$  oxyfluorine anion of RWOFI and CWOFI are either further linked to  $[WO_3F_3]^{3-}$  oxyfluorine anions or replaced by highly electronegative  $F^-$ . The contributions of non-bonding O 2p orbitals in RWOFI and CWOFI are consequently much less important than those in  $K_5(W_3O_9F_4)(IO_3)$ . Secondly, in contrast to the interconnected

tungsten-centered octahedra in  $K_5(W_3O_9F_4)(IO_3)$ , the  $[WO_3F_3]$  octahedra in RWOFI and CWOFI are isolated from each other, which leads to a narrow conduction band width and a significant increase in the bandgap.<sup>15</sup> Thirdly, the bottom of the CB in conventional  $3d^0/4d^0$ -TM iodates mainly consists of  $d^0$ -TM empty d orbitals and O 2p orbitals; in contrast, the heavy  $5d^0$ - $W^{6+}$  cation has a weaker orbital overlap with  $O^{2-}/F^-$  anions in RWOFI and CWOFI due to its relatively low effective electronegativity, which results in an upward shift of the bottom of the CB.<sup>12</sup> The wide bandgaps observed in RWOFI and CWOFI can clearly be attributed to synergism between the  $[WO_3F_3]^{3-}$  and  $[IO_2F_2]^-$  oxyfluorine anions.

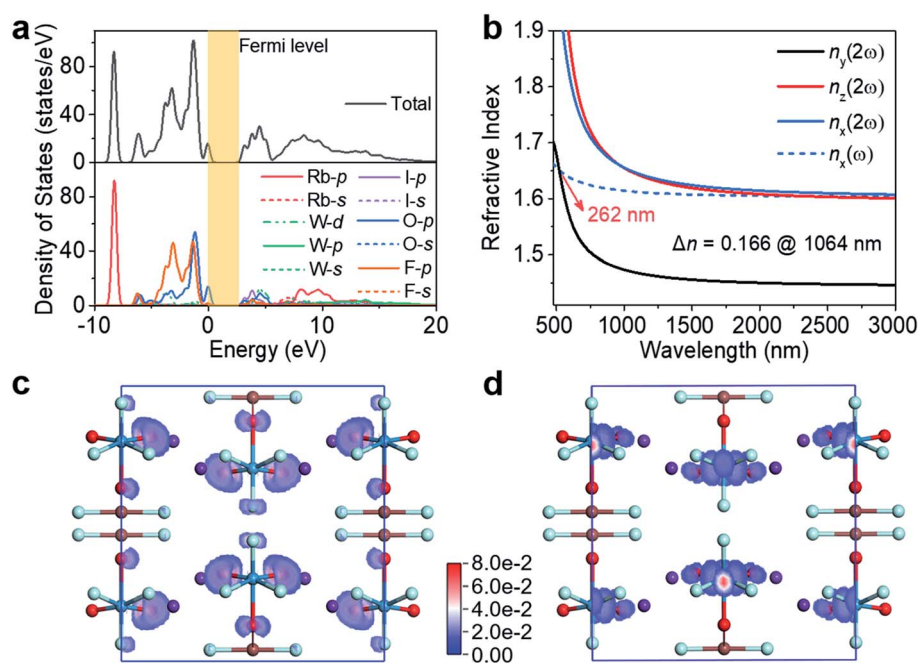


Fig. 4 (a) Total and partial density of states projected onto the constituent atoms in RWOFI. (b) Calculated refractive indexes in RWOFI. SHG-weighted densities for (c) occupied and (d) unoccupied electronic states in RWOFI. Color codes: I brown, W green, O red, F light green, Rb purple.



Based on the restriction of Kleinman's symmetry, there are three non-zero SHG tensors ( $d_{31}$ ,  $d_{32}$ ,  $d_{33}$ ) for RWOFI and CWOFI (listed in Table S7†). The absolute values of the largest tensor  $d_{31}$  are 1.59 and 1.85 pm/V for RWOFI and CWOFI, respectively, corresponding to approximately 4.1 and 4.7 times that of KDP ( $d_{36} = 0.39 \text{ pm V}^{-1}$ ), and therefore consistent with the experimental values. The birefringences are calculated to be 0.166 (RWOFI) and 0.137 (CWOFI) @ 1064 nm. The shortest phase-matchable wavelength of RWOFI is *ca.* 262 nm (Fig. 4b), which is shorter than its UV cut-off edge of 280 nm. In comparison, the shortest phase-matchable wavelength of CWOFI is red-shifted to 302 nm (Fig. S7b†), which is attributed to its relatively small birefringence and large chromatic dispersion. These results are consistent with the phase-matching capability of the two crystalline materials shown experimentally at 1064 nm. A real-space atom-cutting analysis<sup>24</sup> of the contributions of the constituent oxyfluorine anions  $[\text{IO}_2\text{F}_2]^-$  and  $[\text{WO}_3\text{F}_3]^{3-}$  to the NLO properties has been undertaken, the resultant data being listed in Table S7.† The  $[\text{IO}_2\text{F}_2]^-$  oxyfluorine anions make the major contributions to the coefficient  $d_{31}$  (66.5 and 70.2%), the  $[\text{WO}_3\text{F}_3]^{3-}$  oxyfluorine anions account for 27.1 and 27.3%, and the contributions from the alkali-metal cations (6.3 and 2.5%) are negligible, for RWOFI and CWOFI, respectively. In general, the SHG properties are closely related to the relevant virtual (electron and hole) excitations between the states near the Fermi level.<sup>25</sup> A wide bandgap usually increases the difficulty of the virtual excitation, resulting in a weak SHG response, making the simultaneous engineering of a wide bandgap and a strong SHG response a key problem in the development of efficient NLO materials.<sup>26</sup> SHG-weighted electron densities<sup>27</sup> illustrated in Fig. 4c, d, S7c, and d† further confirm the dominant contributions of the 0D  $[\text{WO}_2\text{F}_3(\text{IO}_2\text{F}_2)]^{2-}$  hetero-oxyfluorine anions electron clouds to the SHG responses. Since the SHG properties are closely related to the band structures, the excellent linear and nonlinear optical properties in RWOFI and CWOFI can be assigned to their unique band structures that result from the bandgap engineering by dual fluorination of the two types of SOJT distorted units.

## Conclusions

In summary, the first examples of 5d<sup>0</sup>-TM fluoroiodates,  $\text{A}_2\text{-WO}_2\text{F}_3(\text{IO}_2\text{F}_2)$ , have been successfully synthesized by employing a bandgap engineering strategy based on dual fluorination of two different oxyanions. Due to the optimized combination of two types of fluorinated SOJT distorted units,  $\text{A}_2\text{WO}_2\text{F}_3(\text{IO}_2\text{F}_2)$  exhibit wide bandgaps (4.42 eV (RWOFI), 4.29 eV (CWOFI)), the largest of d<sup>0</sup>-TM iodates to date. They additionally possess strong SHG responses of  $3.8 \times \text{KDP}$  (RWOFI) and  $3.5 \times \text{KDP}$  (CWOFI) at 1064 nm, sufficient birefringence (0.166 (RWOFI) and 0.137 (CWOFI) at 1064 nm), and broad optical transparency windows. Theoretical calculations have elucidated that the wide bandgaps of the two hetero-oxyfluorides originate from collaboration of the  $[\text{WO}_3\text{F}_3]^{3-}$  and  $[\text{IO}_2\text{F}_2]^-$  oxyfluorine anions. This study provides an efficient bandgap engineering strategy for the development of high-performance NLO materials with an optimized balance of linear and nonlinear optical properties.

## Data availability

All of the related experimental and computational data are provided in the ESI.†

## Author contributions

Y. L. H. synthesized the crystals, performed the experiments, and wrote the draft. X. X. J. performed the theoretical calculations. T. H. W. and Y. Y. X. collected the data. W. C. performed SHG measurements and edited the manuscript. Z. P. H., Z. S. L. and M. G. H. reviewed the manuscript. J. X. provided methods for crystal growth. C. Z. supervised the process and edited the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This research was financially supported by the National Natural Science Foundation of China (no. 51432006, 52002276), the Ministry of Education of China for the Changjiang Innovation Research Team (no. IRT14R23), the Ministry of Education and the State Administration of Foreign Experts Affairs for the 111 Project (no. B13025), and the Innovation Program of Shanghai Municipal Education Commission. M. G. H. thanks the Australian Research Council for support (DP170100411).

## Notes and references

- 1 D. N. Nikogosyan, *Nonlinear Optical Crystals: A Complete Survey*, Springer, Heidelberg, 2012.
- 2 (a) C. T. Chen, Y. Wang, B. Wu, K. Wu, W. Zeng and L. Yu, *Nature*, 1995, **373**, 322–324; (b) D. Cyranoski, *Nature*, 2009, **457**, 953–955; (c) L. Wu, S. Patankar, T. Morimoto, N. L. Nair, E. Thewalt, A. Little, J. G. Analytis, J. E. Moore and J. Orenstein, *Nat. Phys.*, 2017, **13**, 350–355; (d) C. Wu, G. Yang, M. G. Humphrey and C. Zhang, *Coord. Chem. Rev.*, 2018, **375**, 459–488.
- 3 (a) K. M. Ok, *Acc. Chem. Res.*, 2016, **49**, 2774–2785; (b) H. H. Cheng, F. M. Li, Z. H. Yang and S. L. Pan, *Angew. Chem., Int. Ed.*, 2022, **61**, e202115669; (c) G. Peng, N. Ye, Z. S. Lin, L. Kang, S. L. Pan, M. Zhang, C. S. Lin, X. F. Long, M. Luo, Y. Chen, Y. H. Tang, F. Xu and T. Yan, *Angew. Chem., Int. Ed.*, 2018, **57**, 8968–8972; (d) P. S. Halasyamani and J. M. Rondinelli, *Nat. Commun.*, 2018, **9**, 2972; (e) C. Wu, T. H. Wu, X. X. Jiang, Z. J. Wang, H. Y. Sha, L. Lin, Z. S. Lin, Z. P. Huang, X. F. Long, M. G. Humphrey and C. Zhang, *J. Am. Chem. Soc.*, 2021, **143**, 4138–4142.
- 4 (a) G. H. Zou and K. M. Ok, *Chem. Sci.*, 2020, **11**, 5404–5409; (b) Z. Y. Bai, L. H. Liu, D. M. Wang, C. L. Hu and Z. B. Lin, *Chem. Sci.*, 2021, **12**, 4014–4020; (c) H. P. Wu, B. B. Zhang, H. W. Yu, Z. G. Hu, J. Y. Wang, Y. C. Wu and P. S. Halasyamani, *Angew. Chem., Int. Ed.*, 2020, **59**, 8922–





- 8926; (d) C. Wu, X. X. Jiang, Z. J. Wang, H. Y. Sha, Z. S. Lin, Z. P. Huang, X. F. Long, M. G. Humphrey and C. Zhang, *Angew. Chem., Int. Ed.*, 2021, **60**, 14806–14810; (e) S. F. Li, X. M. Jiang, Y. H. Fan, B. W. Liu, H. Y. Zeng and G. C. Guo, *Chem. Sci.*, 2018, **9**, 5700–5708.
- 5 (a) H. G. Kim, T. Tran, W. Choi, T. S. You, P. S. Halasyamani and K. M. Ok, *Chem. Mater.*, 2016, **28**, 2424–2432; (b) T. L. Chao, W. J. Chang, S. H. Wen, Y. Q. Lin, B. C. Chang and K. H. Lii, *J. Am. Chem. Soc.*, 2016, **128**, 9061–9064; (c) W. G. Zhang, H. W. Yu, J. Cantwell, H. P. Wu, K. R. Poeppelmeier and P. S. Halasyamani, *Chem. Mater.*, 2016, **28**, 4483–4491; (d) C. Wu, X. X. Jiang, L. Lin, Y. L. Hu, T. H. Wu, Z. S. Lin, Z. P. Huang, M. G. Humphrey and C. Zhang, *Angew. Chem., Int. Ed.*, 2021, **60**, 22447–22453.
- 6 (a) H. W. Yu, J. Young, H. P. Wu, W. G. Zhang, J. M. Rondinelli and P. S. Halasyamani, *J. Am. Chem. Soc.*, 2016, **138**, 4984–4989; (b) J. Chen, C. L. Hu, F. F. Mao, B. P. Yang, X. H. Zhang and J. G. Mao, *Angew. Chem., Int. Ed.*, 2019, **58**, 11666–11669; (c) F. G. You, F. Liang, Q. Huang, Z. G. Hu, Y. C. Wu and Z. S. Lin, *J. Am. Chem. Soc.*, 2019, **141**, 748–752; (d) J. Y. Chung, H. Jo, S. Yeon, H. R. Byun, T. S. You, J. I. Jang and K. M. Ok, *Chem. Mater.*, 2020, **32**, 7318–7326; (e) C. Wu, L. H. Li, L. Lin, Z. P. Huang, M. G. Humphrey and C. Zhang, *Chem. Mater.*, 2020, **32**, 3043–3053.
- 7 (a) P. S. Halasyamani and K. R. Poeppelmeier, *Chem. Mater.*, 1998, **10**, 2753–2769; (b) J. Chen, C. L. Hu, Y. L. Lin, Y. Chen, Q. Q. Chen and J. G. Mao, *Chem. Sci.*, 2022, **13**, 454–460; (c) H. X. Fan, C. S. Lin, K. C. Chen, G. Peng, B. X. Li, G. Zhang, X. F. Long and N. Ye, *Angew. Chem., Int. Ed.*, 2020, **59**, 5268–5272; (d) Y. N. Li, Z. X. Chen, W. D. Yao, R. L. Tang and S. P. Guo, *J. Mater. Chem. C*, 2021, **9**, 8659–8665; (e) R. B. Fu, Q. R. Shui, Z. J. Ma, Z. Q. Zhou, H. X. Tang and X. T. Wu, *Chem.–Eur. J.*, 2022, **28**, e202103687.
- 8 (a) F. H. Ding, K. J. Griffith, C. P. Kocer, R. J. Saballos, Y. R. Wang, C. Zhang, M. L. Nisbet, A. J. Morris, J. M. Rondinelli and K. R. Poeppelmeier, *J. Am. Chem. Soc.*, 2020, **142**, 12288–12298; (b) M. R. Marvel, J. Lesage, J. Baek, P. S. Halasyamani, C. L. Stern and K. R. Poeppelmeier, *J. Am. Chem. Soc.*, 2007, **129**, 13963–13969; (c) H. Jo, M. H. Lee and K. M. Ok, *Chem. Mater.*, 2021, **33**, 1875–1882; (d) G. Peng, Y. Yang, T. Yan, D. Zhao, B. X. Li, G. Zhang, Z. S. Lin and N. Ye, *Chem. Sci.*, 2020, **11**, 7396–7400; (e) S. Liu, X. M. Liu, S. G. Zhao, Y. C. Liu, L. N. Li, Q. R. Ding, Y. Q. Li, Z. S. Lin, J. H. Luo and M. C. Hong, *Angew. Chem., Int. Ed.*, 2020, **59**, 9414–9417; (f) H. X. Tang, R. B. Fu, Z. J. Ma and X. T. Wu, *Inorg. Chem.*, 2021, **60**, 17364–17370.
- 9 (a) H. Y. Chang, S. H. Kim, K. M. Ok and P. S. Halasyamani, *Chem. Mater.*, 2009, **21**, 1654–1662; (b) H. Y. Chang, S. W. Kim and P. S. Halasyamani, *Chem. Mater.*, 2010, **22**, 3241–3250; (c) J. J. Zhang, Z. H. Zhang, Y. X. Sun, C. Q. Zhang, S. J. Zhang, Y. Liu and X. T. Tao, *J. Mater. Chem.*, 2012, **22**, 9921–9927; (d) R. E. Sykora, K. M. Ok, P. S. Halasyamani and T. E. Albrecht-Schmitt, *J. Am. Chem. Soc.*, 2002, **124**, 1951–1957; (e) Y. H. Li, G. P. Han, H. W. Yu, H. Li, Z. H. Yang and S. L. Pan, *Chem. Mater.*, 2019, **31**, 2992–3000; (f) C. F. Sun, C. L. Hu, X. Xu, B. P. Yang and J. G. Mao, *J. Am. Chem. Soc.*, 2011, **133**, 5561–5572; (g) W. Zeng, X. H. Dong, Y. Tian, L. Huang, H. M. Zeng, Z. E. Lin and G. H. Zou, *Chem. Commun.*, 2022, **58**, 3350–3353.
- 10 (a) Y. L. Hu, C. Wu, X. X. Jiang, Z. J. Wang, Z. P. Huang, Z. S. Lin, X. F. Long, M. G. Humphrey and C. Zhang, *J. Am. Chem. Soc.*, 2021, **143**, 12455–12459; (b) C. Wu, X. X. Jiang, Z. J. Wang, L. Lin, Z. S. Lin, Z. P. Huang, X. F. Long, M. G. Humphrey and C. Zhang, *Angew. Chem., Int. Ed.*, 2021, **60**, 3464–3468; (c) J. Chen, C. L. Hu, X. H. Zhang, B. X. Li, B. P. Yang and J. G. Mao, *Angew. Chem., Int. Ed.*, 2020, **59**, 5381–5384.
- 11 L. Kang, F. Liang, X. X. Jiang, Z. S. Lin and C. T. Chen, *Acc. Chem. Res.*, 2020, **53**, 209–217.
- 12 H. W. Eng, P. W. Barnes, B. M. Auer and P. M. Woodward, *J. Solid State Chem.*, 2003, **175**, 94–109.
- 13 (a) M. Mutailipu, M. Zhang, Z. H. Yang and S. L. Pan, *Acc. Chem. Res.*, 2019, **52**, 791–801; (b) G. Q. Shi, Y. Wang, F. F. Zhang, B. B. Zhang, Z. H. Yang, X. L. Hou, S. L. Pan and K. R. Poeppelmeier, *J. Am. Chem. Soc.*, 2017, **139**, 10645–10648.
- 14 J. Lu, J. N. Yue, L. Xiong, W. K. Zhang, L. Chen and L. M. Wu, *J. Am. Chem. Soc.*, 2019, **141**, 8093–8097.
- 15 E. A. Axtell, Y. Park, K. Chondroudis and M. G. Kanatzidis, *J. Am. Chem. Soc.*, 1998, **120**, 124–136.
- 16 C. Wu, L. Lin, X. X. Jiang, Z. S. Lin, Z. P. Huang, M. G. Humphrey, P. S. Halasyamani and C. Zhang, *Chem. Mater.*, 2019, **31**, 10100–10108.
- 17 (a) L. Lin, X. X. Jiang, C. Wu, Z. S. Lin, Z. P. Huang, M. G. Humphrey and C. Zhang, *Chem. Mater.*, 2021, **33**, 5555–5562; (b) H. W. Yu, M. L. Nisbet and K. R. Poeppelmeier, *J. Am. Chem. Soc.*, 2018, **140**, 8868–8876; (c) J. Chen, C. L. Hu, F. F. Mao, J. H. Feng and J. G. Mao, *Angew. Chem., Int. Ed.*, 2019, **58**, 2098–2102.
- 18 P. S. Halasyamani, *Chem. Mater.*, 2004, **16**, 3586–3592.
- 19 J. Chen, C. L. Hu, F. Kong and J. G. Mao, *Acc. Chem. Res.*, 2021, **54**, 2775–2783.
- 20 M. Q. Gai, T. H. Tong, Y. Wang, Z. H. Yang and S. L. Pan, *Chem. Mater.*, 2020, **32**, 5723–5728.
- 21 Q. Wu, H. M. Liu, F. C. Jiang, L. Kang, L. Yang, Z. S. Lin, Z. G. Hu, X. G. Chen, X. G. Meng and J. G. Qin, *Chem. Mater.*, 2016, **28**, 1413–1418.
- 22 (a) P. A. Maggard, T. S. Nault, C. L. Stern and K. R. Poeppelmeier, *J. Solid State Chem.*, 2003, **175**, 27–33; (b) C. Wu, X. X. Jiang, L. Lin, W. Y. Dan, Z. S. Lin, Z. P. Huang, M. G. Humphrey and C. Zhang, *Angew. Chem., Int. Ed.*, 2021, **60**, 27151–27157.
- 23 Z. S. Lin, X. X. Jiang, L. Kang, P. F. Gong, S. Y. Luo and M. H. Lee, *J. Phys. D: Appl. Phys.*, 2014, **47**, 253001.
- 24 J. Lin, M. H. Lee, Z. P. Liu, C. T. Chen and C. J. Pickard, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, **60**, 13380–13389.
- 25 R. He, Z. S. Lin, M. H. Lee and C. T. Chen, *J. Appl. Phys.*, 2011, **109**, 103510.
- 26 K. Wu and S. L. Pan, *Coord. Chem. Rev.*, 2018, **377**, 191–208.
- 27 M. H. Lee, C. H. Yang and J. H. Jan, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2004, **70**, 235110.

