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Novel antimony-based mixed halides exhibiting an excellent SHG response and a broad transmission range†

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In this study, we successfully synthesized two novel non-centrosymmetric (NCS) antimony halide compounds, **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**. These compounds were formed by combining Sb^{3+} cations, which possess stereochemically active lone pairs (SCALP), with halogen atoms (fluorine and chlorine). The introduction of mixed halogen atoms enables Sb atoms to adopt multiple coordination modes, which in turn promotes the development of both compounds towards NCS structures. Notably, **$Rb_3Sb_4F_{14}Cl$** demonstrated a large second harmonic generation (SHG) response, approximately three times greater than that of KH_2PO_4 (KDP), along with a wide transparency range from 0.27 to 13.3 μm and a high laser damage threshold (LDT) of 223 MW cm^{-2}. These exceptional properties indicate that **$Rb_3Sb_4F_{14}Cl$** is a highly promising nonlinear optical (NLO) crystal with excellent overall performance, making it suitable for potential applications spanning the short-wave ultraviolet (UV) to mid-infrared (IR) spectral regions. In contrast, **$RbSbF_3Cl$** exhibits a large birefringence of 0.31 at 546 nm as a uniaxial crystal, suggesting its practicability as a birefringent material.

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Introduction

Nonlinear optical (NLO) materials have significant academic and technological value in extending the wavelength range of solid-state lasers from deep ultraviolet (UV) (<200 nm) to infrared (IR) (~20 μm) through simple frequency conversion.[1–6] From a commercial perspective, an ideal NLO crystal must possess several key characteristics, one of which is a broad transparent window, which determines the application range of the crystal.[7–10] Over the past few decades, several well-known NLO crystals, including LiB_3O_5 (LBO),[11] β-BaB_2O_4 (β-BBO),[12] and $KTiOPO_4$ (KTP),[13] have found commercial applications in the UV and visible regions. However, in the IR region, the selection is limited to a few materials, such as $AgGaS_2$,[14] $AgGaSe_2$,[15] and $ZnGeP_2$,[16] which are suitable for the mid-IR range (3–12 μm). These materials are hampered by low laser damage thresholds (LDT), significantly restricting their utility in laser communication and high-power applications. Compounds that simultaneously exhibit strong second harmonic generation (SHG) effects and broad spectral transmission, from the UV region to the IR region, are exceptionally rare. This rarity stems from the inherent trade-offs between optical properties, such as the band gap and NLO response. Typically, a large SHG response is coupled with a narrow transmission range and low LDT. Thus, balancing the competing demands for a broad transmission range and superior overall optical properties remains a substantial challenge in the development of novel NLO materials.

It is well established that the primary requirement for NLO crystal materials is a non-centrosymmetric (NCS) structure.[17,18] Thus, designing and synthesizing crystals with NCS structures is of great significance. Recently, metal halides have demonstrated strong potential as NLO materials, largely due to their tendency to form NCS structures, as observed in compounds like **Na_2CeF_6**[19] and **$SrCl_2$**·$6H_2O$.[20] Research studies have indicated that the properties of metal halides are predominantly influenced by the choice of the central metal cation.[21–26] For instance, selecting metals with stereochemically active lone pairs (SCALP) (*e.g.*, Sn^{2+}, Sb^{3+}, Bi^{3+}, and Pb^{2+}) often results in compounds with exceptional NLO properties, as evidenced by $Pb_3Mg_3TeP_2O_{14}$ (13.5 × KDP),[27] $Bi(IO_3)F_2$ (11.5 × KDP),[28] and $Sn(IO_3)_2F_2$ (3 × KDP).[29] In this study, we focus on antimony

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halides, building on our recent work with antimony oxysalt NLO crystals. Sb-based cations possess SCALP, which are easily polarizable and can favorably contribute to NLO performance. Additionally, antimony is a relatively robust metal, and its compounds typically exhibit wide band gaps, resulting in high LDT values.[30] Compounds such as **$K_2Sb(P_2O_7)F$** (4.0 × KDP, 4.74 eV),[31] **$CsSbF_2SO_4$** (3.0 × KDP, 4.76 eV),[32] and **$Rb_2SbFP_2O_7$** (5.1 × KDP, 4.76 eV)[33] have been found to possess a strong SHG response and a broad transmittance range.

To enhance the likelihood of obtaining NCS antimony halide crystals, a common strategy is to modify their structures and properties by introducing mixed halogens. Sb^{3+} cations readily coordinate with halogen atoms (F/Cl/Br/I), forming a variety of irregular geometrical configurations, including SbX_3 triangular pyramids, SbX_4 seesaws, and SbX_5 tetragonal pyramids. By integrating these configurations with different combinations of the four halogen elements, a diverse array of halide compounds with varied structural features can be synthesized. This approach enables the discovery of candidates with excellent comprehensive properties, including significant NLO effects, high LDTs, and broad transparency windows, such as **$Cs_2Hg_2Br_2I_4$**·H_2O (6 × KDP),[34] **$Rb_2CdBr_2I_2$** (4 × KDP),[35] and **Hg_2BrI_3** (1.2 × KTP).[36]

Building on the halogen mixing strategy, two novel antimony-based mixed halide crystals, **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**, have been successfully synthesized by combining Sb^{3+} cations with fluorine and chlorine atoms. The two compounds feature NCS structures composed of different Sb-based polyhedra. Remarkably, **$Rb_3Sb_4F_{14}Cl$** exhibits a large SHG response, achieving a value three times higher than that of KDP. It also possesses a broad transparency range of 0.27–13.3 μm and a high LDT of 223 MW cm^{-2}. Meanwhile, **$RbSbF_3Cl$**, as a uniaxial crystal, features a high birefringence of 0.31 at 546 nm.

Experimental section

Synthesis of $RbSbF_3Cl$ and $Rb_3Sb_4F_{14}Cl$

The reaction reagents SbF_3 (≥99.8%), Rb_2CO_3 (≥99.0%), and HCl (AR grade) were obtained from Aladdin and used without further purification.

The single crystals of both the compounds, **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**, were synthesized using an aqueous solution method. The same raw materials of Rb_2CO_3 (0.231 g, 1 mmol) and SbF_3 (0.358 g, 2 mmol) were dissolved in 1 mL of distilled water, to which 0.3 mL and 0.1 mL of HCl (12 mol L^{-1}) were added to obtain **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**, respectively. The resulting mixtures were stirred at room temperature for 15 to 20 minutes and then filtered. The filtrates were allowed to evaporate slowly in a refrigerator at 5 °C. After two days, colorless and transparent block crystals of **$RbSbF_3Cl$** and flake-like crystals of **$Rb_3Sb_4F_{14}Cl$** were collected from the bottom of the plastic beakers (Fig. S1†). The yields of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** were approximately 38% and 42%, respectively, based on antimony.

Results and discussion

Crystal structure description

$RbSbF_3Cl$ crystallizes in the tetragonal crystal system, belonging to the NCS space group $I\bar{4}2m$ (No. 121) (Tables S1, S2 and S4†).[37,38] The asymmetric unit comprises one Sb atom, one Rb atom, one Cl atom, and three F atoms. The Sb atom adopts a tetragonal pyramidal geometry as $[SbF_3Cl_2]^{2-}$ (**I**), coordinating with three F atoms and two Cl atoms (Fig. 1a). The Sb–F bond lengths range from 1.917 to 1.962 Å, while the Sb–Cl bond length measures 2.889 Å. In this compound, four $[SbF_3Cl_2]^{2-}$ (**I**) units interconnect by sharing four Cl atoms, forming a square-shaped $[Sb_4F_{12}Cl_4]^{4-}$ (**II**) structure with four-membered rings (4-MR) arranged orderly within the *a*–*b* plane (Fig. 1b). Additionally, Rb^+ ions occupy the cavities of the 4-MR, serving as charge balancers and exhibiting two distinct coordination modes in **$RbSbF_3Cl$**. Both Rb1 and Rb2 atoms coordinate with four Cl atoms, while they bind with six and eight F atoms, respectively, resulting in the anionic complexes $[RbF_6Cl_4]^{9-}$ and $[RbF_8Cl_4]^{11-}$. The Rb–F bond lengths range from 2.974 to 3.562 Å, while the Rb1–Cl and Rb2–Cl bond lengths are 3.544 Å and 3.424 Å, respectively (Fig. S2a†).

$Rb_3Sb_4F_{14}Cl$ forms crystals in the orthorhombic system's NCS space group $Pmn2_1$ (No. 31) (Tables S1, S3 and S5†). The asymmetric unit comprises four Sb atoms, three Rb atoms, one Cl atom, and fourteen F atoms. In this compound, the three distinct Sb atoms exhibit different coordination environments; $[SbF_3Cl_2]^{2-}$ (**I**) and $[SbF_5]^{2-}$ (**IV**) form tetragonal pyramidal configurations, while $[SbF_4]^-$ (**III**) adopts a seesaw configuration, with varying numbers of coordinating F and Cl atoms. The Sb–Cl bond length is 2.894 Å, whereas the Sb–F bond lengths vary from 1.913 to 2.450 Å. These polyhedra further assemble into structurally distinct clusters. For instance, two $[SbF_4]^-$ (**III**) polyhedra share an F atom to form a $[Sb_2F_7]^-$ (**V**) cluster, while a $[Sb_3F_{11}]^{2-}$ (**VI**) cluster is formed by sharing two F atoms among two $[SbF_4]^-$ (**III**) units and one $[SbF_5]^{2-}$ (**IV**) unit (Fig. 1c). These four clusters are arranged sequentially along the *a*-axis in the order $[SbF_3Cl_2]^{2-}$ (**I**), $[Sb_2F_7]^-$ (**V**), $[Sb_3F_{11}]^{2-}$ (**VI**), and $[Sb_2F_7]^-$ (**V**), repeating this pattern between adjacent layers (Fig. 1d). Additionally, Rb^+ cations serve as charge balancers and are systematically positioned within the cavities between the Sb-polyhedra, displaying three distinct coordination modes in **$Rb_3Sb_4F_{14}Cl$** (Fig. S2b†). Specifically, Rb1 coordinates with one Cl atom and nine F atoms to form the $[RbF_9Cl]^{9-}$ polyhedron, Rb2 coordinates with one Cl atom and ten F atoms to form $[RbF_{10}Cl]^{10-}$, and Rb3 coordinates with nine F atoms to form the $[RbF_9]^{8-}$ polyhedron. The Rb–F bond lengths range from 2.769 to 3.454 Å, while the Rb1–Cl and Rb2–Cl bond lengths are 3.356 Å and 3.393 Å, respectively.

By calculating the bond valence sum (BVS), the validity of the structures of both compounds was confirmed.[39] The atomic oxidation states of Rb^+, Sb^{3+}, F^-, and Cl^- in **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** were determined to be 0.93–1.15, 2.90–3.20, 0.90–1.29, and 0.89–1.09, respectively (Tables S2 and S3†).

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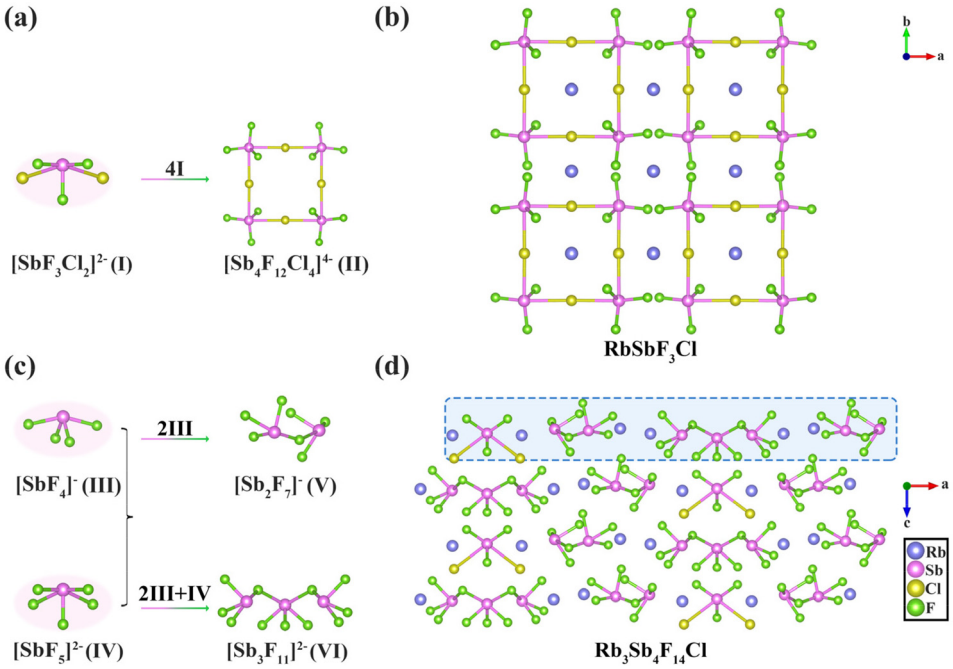

Fig. 1 (a and c) Various coordination patterns and linkage modes of Sb^{3+}. (b) Arrangement of 4-MR and Rb^{+} cations in **$RbSbF_3Cl$** along the *a*–*b* plane. (d) Arrangement of different Sb polyhedra and Rb^{+} cations in **$Rb_3Sb_4F_{14}Cl$**.

Powder X-ray diffraction

The phase purity of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** was verified through powder X-ray diffraction analysis. The experimental data, as presented in Fig. S3,† are consistent with the patterns derived from single-crystal X-ray diffraction, confirming that the powder samples are indeed pure phases.

Thermal properties

Thermogravimetric analysis (TGA) was conducted to evaluate the thermal stability of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**. As depicted in Fig. S4,† the TGA curves indicate that both the compounds are stable up to approximately 200 °C.

Optical properties

The IR spectra of the two compounds are shown in Fig. 2a, with characteristic absorption bands near 568/513/471 cm^{-1} and 751/572/528/479 cm^{-1}, corresponding to the stretching vibrations of the Sb–F and Sb–Cl bonds. These vibrational modes align with those reported in the literature.[40,41] The UV-vis-NIR diffuse reflectance spectra of both **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**, presented in Fig. 2b and Fig. S5,† indicate band gaps of 4.49 and 4.60 eV, corresponding to UV cutoff edges at 276 and 269 nm.[42] Based on the data from the IR and UV-vis-NIR spectra, the two compounds are transparent across 0.28 to 17.6 and 0.27 to 13.3 μm, respectively, covering short-wave UV to mid-IR regions.

To develop broadband NLO materials, the main group Sb^{3+} cations and F^{-} ions were combined to shift the cutoff edge towards shorter wavelengths. The distortion of Sb^{3+} upon coordination promotes asymmetric structures, which may also enhance SHG response. Additionally, the incorporation of Rb^{+}, which lacks d–d and f–f transitions, minimizes electronic absorption in the visible and UV regions, potentially lowering the UV cutoff and extending the transparency range. As a result, the title compounds exhibit short UV cutoffs at 276 and 269 nm. Fig. 2c compares the UV cutoff edges of the title compounds with various metal halides,[30,35,36,43–54] such as **$K_2SbF_2Cl_3$**,[30] **$Cs_2HgI_2Cl_2$**,[43] and **$CsSbF_3Cl$**,[44] which have cutoffs at 309 nm, 394 nm, and 405 nm, respectively, and are transparent in the UV range. Unlike most reported metal halides, the two title compounds presented here exhibit broad transmission ranges (0.28–17.6 μm and 0.27–13.3 μm), covering the short-wave UV to mid-IR regions, highlighting their potential as broadband optical materials.

NLO and birefringence properties

The SHG properties of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** were evaluated using KDP as a reference since they crystallize in the NCS space groups $I\bar{4}2m$ and $Pmn2_1$, respectively.[55] However, the results of the frequency doubling performance of the two compounds vary a lot, with compound **$RbSbF_3Cl$** being undetectable due to the small value of the frequency doubling response signal, while **$Rb_3Sb_4F_{14}Cl$** indicates a large SHG response, which is approximately 3 times that of KDP, as depicted in Fig. 3a. Additionally, the SHG signal increases with the increased particle size of the **$Rb_3Sb_4F_{14}Cl$** crystals, suggesting that compound **$Rb_3Sb_4F_{14}Cl$** exhibits type I phase-matching properties (Fig. 3a). The difference in the SHG response of the two compounds was explored, which may be affected by the orientations of the lone pair of electrons of Sb in the unit cell.

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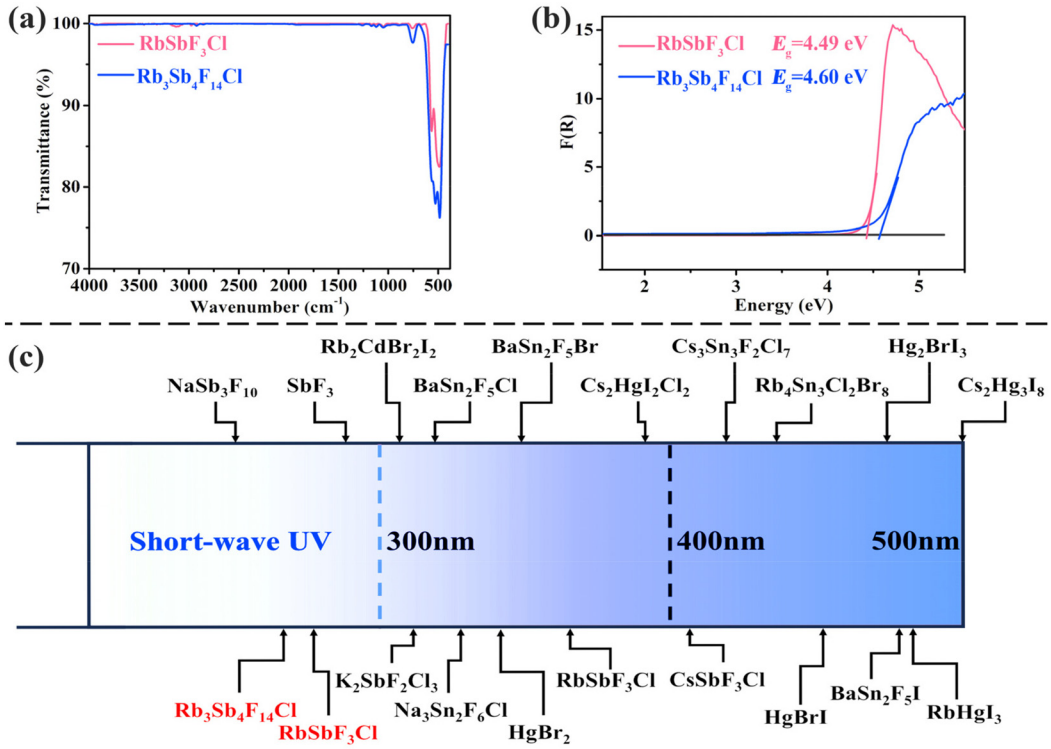

Fig. 2 (a) IR spectra of compounds **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**. (b) Band gaps for **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**. (c) Comparison of the cutoff edges of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** with those various reported metal halides.

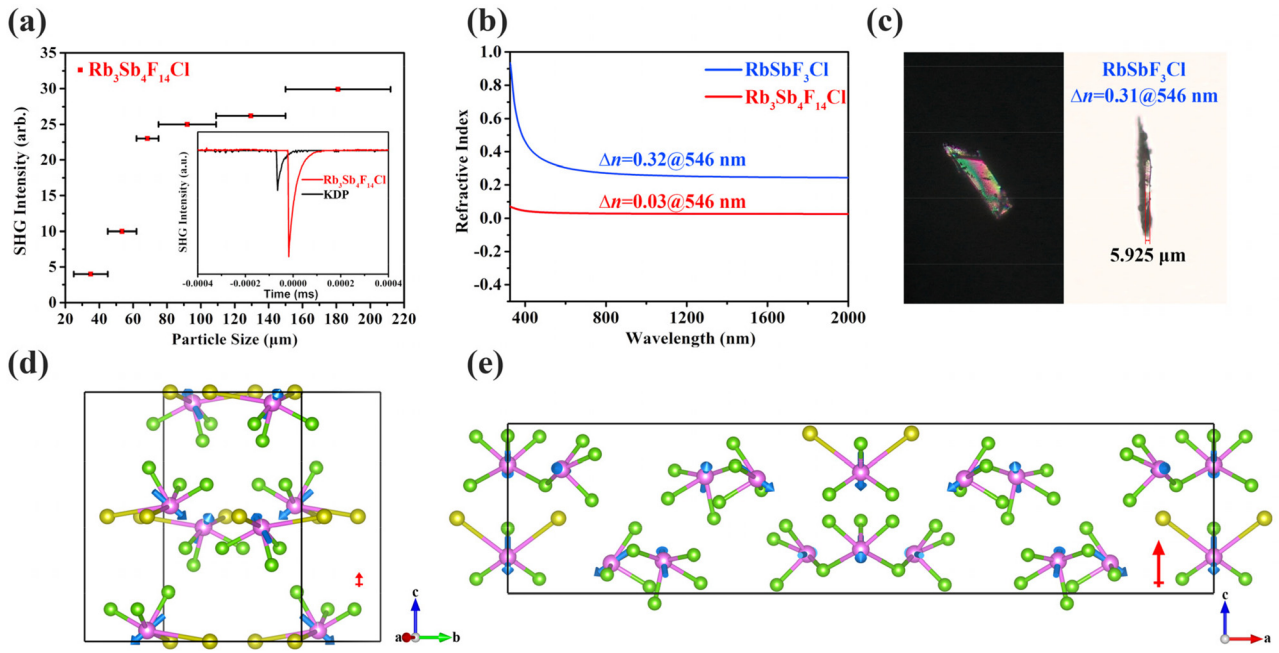

Fig. 3 (a) The phase-matching curve for **$Rb_3Sb_4F_{14}Cl$**. Inset: SHG intensity with KDP as the reference (150–212 μm). (b) Calculated birefringence for **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**. (c) The experimental birefringence of **$RbSbF_3Cl$**. (d) The orientations of the lone pair of electrons in the unit cell of **$RbSbF_3Cl$**. (e) The orientations of the lone pair of electrons on Sb^{3+} within a single cell in **$Rb_3Sb_4F_{14}Cl$**. The direction of the overall dipole moments is highlighted by a red arrow.

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The orientations of the lone pair of electrons in compound **$RbSbF_3Cl$** are demonstrated in Fig. 3d. By roughly estimating the total dipole moment, due to the nearly opposite orientations of the effective dipole moments of Sb^{3+} ions along the *c*-axis, the contribution of Sb^{3+} cations to the SHG effect is almost entirely canceled out. As a result, the SHG effect of this compound is extremely weak and cannot be detected by instrumentation. However, this arrangement is conducive to the superposition of birefringence, because birefringence is scarcely affected or canceled by the opposite directions of the lone pair of electrons based on current research studies.[56,57] Hence, **$RbSbF_3Cl$** exhibits a large experimental birefringence of 0.31@546 nm ascribed to the superposition of the lone pair of electrons of Sb^{3+} along the *c*-axis direction (Fig. 3c).[58] For the other compound **$Rb_3Sb_4F_{14}Cl$**, it exhibits a significantly larger SHG effect, approximately three times that of KDP, compared to **$RbSbF_3Cl$**, which can be attributed to the higher density of Sb^{3+} in the unit cell (Table S6†), despite the relatively disordered orientation of the lone pair of electrons of the central Sb^{3+} cation (Fig. 3e). However, the disordered orientation results in a much smaller calculated birefringence of 0.03 at 546 nm (Fig. 3b). Compared to some SCALP-containing oxysalts,[59] such as $BiOIO_3$[60] and $LiHgPO_4$,[61] the title compound **$Rb_3Sb_4F_{14}Cl$** shows a relatively weaker SHG response but features a significantly shorter UV cutoff edge. This confirms that the mixed halide antimony compounds have potential as broadband second-harmonic generation materials.

LDT measurements

As described above, the band gaps of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** were found to be 4.49 and 4.60 eV, respectively. Generally, a larger band gap indicates a higher LDT. The LDTs of the two compounds were tested using a 1064 nm laser with a 10 ns pulse width, resulting in values of 163 and 223 MW cm^{-2}, respectively, five and seven times higher than that of $AgGaS_2$ (30 MW cm^{-2}) under the same conditions (Table S7†).[62]

Theoretical calculations

DFT calculations were conducted to investigate the relationship between the structures and optical properties of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**.[63] The birefringence of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** was calculated to be 0.32 and 0.03 at 546 nm (Fig. 3b), respectively. For that $n_0 > n_e$ in **$RbSbF_3Cl$**, indicating that it is a uniaxial crystal (Fig. S6†). Furthermore, the theoretical band gaps of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** were determined to be 3.79 and 4.04 eV, respectively, both of which are 0.70 and 0.56 eV lower than their experimental values (Fig. 4a and b).[64] These discrepancies align with the well-known tendency of the DFT-GGA approach to underestimate band gaps.[65] Additionally, the total densities of states (TDOS) and partial densities of states (PDOS) for **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** were computed, offering detailed insights into the contributions of individual atomic orbitals to the energy bands (Fig. 4c and d). For **$RbSbF_3Cl$**, the upper valence bands, spanning from −10 to 0 eV, are primarily composed of F–2p, Cl–3p, Sb–5p, and Sb–5s orbitals. In contrast, the lower conduction bands, within the energy range of 0 to 10 eV, predominantly consist of Sb–5p, Rb–5s, and Rb–5p orbitals. Similarly, in **$Rb_3Sb_4F_{14}Cl$**, the valence bands from −10 to 0 eV are primarily made up of F–2p, Rb–5p, Cl–3p, Sb–5s, and Sb–5p orbitals. Moreover, the conduction bands between 0 and 10 eV are largely dominated by Sb–5p and F–2p orbitals. The graphs of the density of states for **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$** show a significant overlap

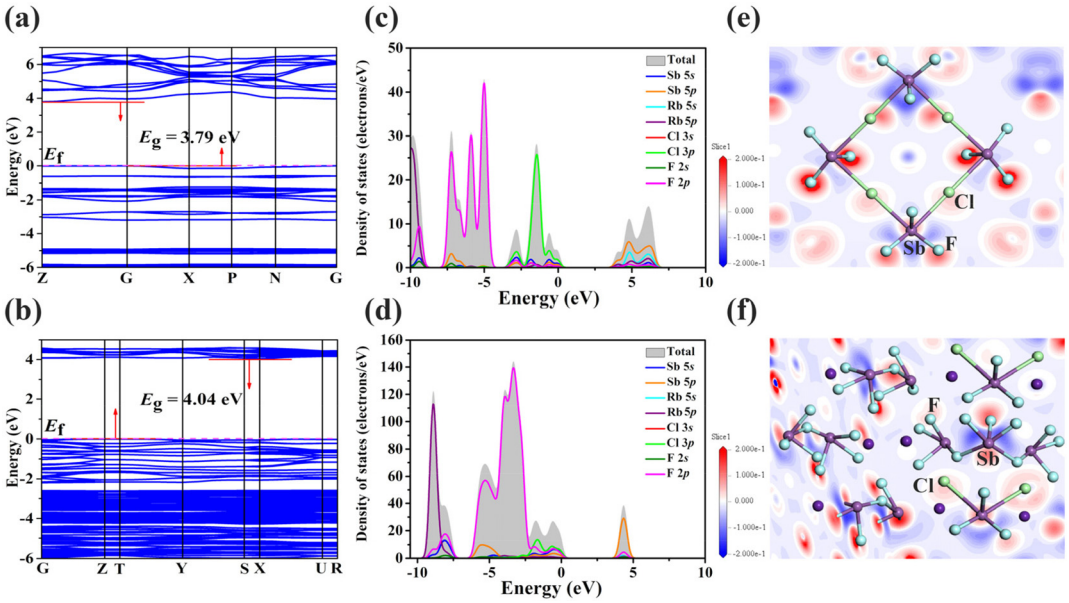

Fig. 4 (a and b) Calculated band structures for **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**. (c and b) The TDOS and PDOS of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**. (e and f) Electron density difference maps of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**.

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between the Sb–5s and Sb–5p orbitals and the F–2p and Cl–3p orbitals, indicating the presence of Sb–F and Sb–Cl bonds. It is well established that states near the Fermi level play a critical role in determining the NLO properties of a compound. In the case of **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**, the electronic orbitals around the Fermi level include Sb–5s, Sb–5p, F–2p, and Cl–3p, indicating that the optical properties of **$RbSbF_3Cl$** are primarily influenced by the $[SbF_3Cl_2]^{2-}$ groups, whereas those of **$Rb_3Sb_4F_{14}Cl$** are mainly influenced by the $[SbF_3Cl_2]^{2-}$, $[SbF_4]^-$, and $[SbF_5]^{2-}$ groups. To further validate this inference, dipole moment calculations were performed for both compounds. The results indicate that in **$RbSbF_3Cl$**, the calculated dipole moment of the $[SbF_3Cl_2]^{2-}$ groups is almost zero (Table S8†). For **$Rb_3Sb_4F_{14}Cl$**, the local dipole moments of the tetragonal pyramidal $[SbF_3Cl_2]^{2-}$ and $[SbF_5]^{2-}$ units and the seesaw $[SbF_4]^-$ unit were calculated to be 35.18 D (Debye), 45.64 D, and 39.71 D along the *z*-component, respectively, while the *x*- and *y*-components of all dipole moments in **$Rb_3Sb_4F_{14}Cl$** were close to zero (Table S9†). This indicates that all three polyhedra contribute collectively to the NLO performance of **$Rb_3Sb_4F_{14}Cl$**. Moreover, highly asymmetric lobes observed around the Sb^{3+} cations can be attributed to the presence of lone pairs (Fig. 4e and f).

Conclusions

In this study, we successfully synthesized two NCS compounds, **$RbSbF_3Cl$** and **$Rb_3Sb_4F_{14}Cl$**, using a halogen-mixing strategy in a Sb^{3+}-containing system. Among these, **$Rb_3Sb_4F_{14}Cl$** stands out due to its large SHG response, which is approximately 3 times greater than that of KDP. Additionally, it demonstrates high transparency across a wide spectral range, from short-wave UV to mid-IR (0.27 to 13.3 μm), and exhibits a high LDT of 223 MW cm^{-2}. These remarkable properties make **$Rb_3Sb_4F_{14}Cl$** a highly promising NLO crystal with superior overall optical performance. Furthermore, **$RbSbF_3Cl$**, a uniaxial crystal, displays a notable birefringence of approximately 0.31 at 546 nm. This work thus not only presents **$Rb_3Sb_4F_{14}Cl$** as a new benchmark in advanced NLO crystal design but also highlights the exceptional birefringent characteristics of **$RbSbF_3Cl$**, demonstrating a promising pathway for the development of next-generation nonlinear optical materials.

Data availability

The authors confirm that the data supporting the findings of this study are available within the article and its ESI.† The authors will supply the relevant data in response to reasonable requests.

Conflicts of interest

The authors declare no competing financial interest.

Acknowledgements

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