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Introduction

Nonlinear optical (NLO) materials, as the core components of all-solid-state lasers, play a crucial role in current scientific and technological applications.^{1–8} Based on these NLO crystals, ultraviolet (UV) and deep-ultraviolet (DUV; $\lambda < 200$ nm) coherent radiation can be directly obtained through cascading second harmonic generation (SHG), which has a variety of applications in laser micromachining, semiconductor manufacturing, photolithography, and high-resolution spectroscopy.^{9–15}

During the past few decades, borates have been considered a major part of the exploration of UV or DUV NLO crystals because of their excellent structural units, *i.e.*, $[\text{BO}_3]$ or $[\text{BO}_4]$, and their combinations subsequently result in the formation of a wide variety of structural topologies,^{16–18} such as $[\text{BO}_3]$ in $\text{KBe}_2\text{BO}_3\text{F}_2$ (KBBF),¹⁹ $[\text{B}_3\text{O}_6]$ in $\beta\text{-BaB}_2\text{O}_4$ ($\beta\text{-BBO}$),²⁰ $[\text{B}_3\text{O}_7]$ in LiB_3O_5 (LBO),²¹ and $[\text{B}_3\text{O}_7]$ in $\text{CsLiB}_6\text{O}_{10}$ (CLBO).²² These functional units make borates possess not only a short UV cutoff edge, but also a large SHG response in non-centrosymmetric (NCS) structures through their excellent combination and arrangement. Among them, in KBBF, $[\text{BO}_3]$ groups with a coplanar configuration and an aligned arrangement are linked by $[\text{BeO}_3\text{F}]$ to build unique ${}^2\infty[\text{Be}_2\text{BO}_3\text{F}_2]$ layers, which

result in the balance of its short cutoff edge, suitable SHG response and birefringence. These superior optical properties make KBBF the sole NLO crystal to generate DUV coherent radiation light by SHG. However, its strong layering tendency due to the weak interlayer bonding and the high toxicity of the contained beryllium limits the wide applications of KBBF in the DUV region.¹⁹ To overcome the layer habit of KBBF, an effective strategy is to develop borates with reinforced interlayer bonding by introducing stronger B–O/Be–F bonds or developing single layers into double layers.²³ The successful examples include $\text{NaSr}_3\text{Be}_3\text{B}_3\text{O}_9\text{F}_4$ (NSBBF),²⁴ $\text{Na}_2\text{CsBe}_6\text{B}_5\text{O}_{15}$ (NCBBO),²⁵ $\text{Sr}_2\text{Be}_2\text{B}_2\text{O}_7$ (SBBO),²⁶ *etc.* However, the toxicity of beryllium still hinders their applications.²⁷

Aluminoborates have been considered attractive candidates for replacing Be-borates or Be-borate fluorides in UV NLO materials since the tetrahedral coordination environment of the Al^{3+} cation is similar to that of the Be^{2+} cation, and aluminum has lower toxicity than beryllium.²⁸ Based on this, a series of aluminoborates have been synthesized, such as $\text{K}_2\text{Al}_2\text{B}_2\text{O}_7$,^{29,30} $\beta\text{-Rb}_2\text{Al}_2\text{B}_2\text{O}_7$,²⁸ $\text{Cs}_2\text{Al}_2\text{B}_2\text{O}_{13}$,³¹ $\text{K}_3\text{Ba}_3\text{Li}_2\text{Al}_4\text{B}_6\text{O}_{20}\text{F}$,³² $\text{Rb}_3\text{Ba}_3\text{Li}_2\text{Al}_4\text{B}_6\text{O}_{20}\text{F}$,³³ *etc.* Among them, $\text{K}_2\text{Al}_2\text{B}_2\text{O}_7$ has a transparent window into the DUV region (180–3600 nm) and an appropriate birefringence of $\Delta n = 0.07$ at 589 nm. The shortest Type I phase-matching (PM) wavelength is evaluated to be 232.5 nm, which can be used for the fourth harmonic generation (FHG) of the Nd:YAG laser (1064 nm).³⁴ The results demonstrate the UV NLO application potential of $\text{K}_2\text{Al}_2\text{B}_2\text{O}_7$. As such, we will

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continue to explore new UV or DUV NLO crystals in aluminoborates.

Through systematic investigations, a new beryllium-free SBBO-like aluminoborate, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$, was successfully synthesized. In $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$, the two-dimensional (2D) $^2\infty[\text{B}_3\text{AlO}_6]$ single layers are derived from the $^2\infty[\text{Be}(\text{BO}_3)\text{O}]$ single layers of SBBO, in which the $[\text{BeO}_4]$ tetrahedra are replaced by $[\text{AlO}_4]$ tetrahedra and the $[\text{BO}_3]$ triangles are substituted by $[\text{B}_3\text{O}_7]$ units. The single layers are linked by the one-dimensional (1D) $^1\infty[\text{B}_6\text{O}_{13}]$ chains, forming $^2\infty[\text{B}_{12}\text{Al}_2\text{O}_{28}]$ double layers with tunnels, which are filled with the Ba^{2+} cations. Meanwhile, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ exhibits a large SHG response ($2.7 \times \text{KH}_2\text{PO}_4$ (KDP)), a short absorption edge (<190 nm), and suitable birefringence. Compared to other reported aluminoborates, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ achieves an optimal balance between SHG efficiency and DUV transparency. Notably, congruent melting thermal properties are also observed, and a single crystal with $9 \times 5 \times 2$ mm³ has been grown by a top-seeded-solution-growth (TSSG) technique. These results demonstrate that $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ is a promising UV NLO material. In this report, we present the synthesis, crystal structure, thermal behavior, NLO properties, structure–property relationships, and the first-principles calculations of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$.

Results and discussion

Pure polycrystalline $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ was synthesized through a high-temperature solid-state reaction. The experimental powder X-ray diffraction (PXRD) patterns exhibit good agreement with the calculated results derived from the structural data (Fig. S1). To investigate the thermal behavior of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$, thermogravimetry-differential scanning calorimetry (TG-DSC) analysis was performed on the pure polycrystalline sample (Fig. S2). Clearly, there is only one endothermic peak at

840 °C on its DSC curve, and no weight loss is observed on the TG curve. To further confirm its thermal behavior, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ was heated to 850 °C, and the polycrystalline powder sample completely melted. Then the melt was slowly cooled to room temperature. The PXRD patterns of the sample after melting were consistent with those of the pure polycrystalline sample (Fig. S3). These results demonstrate that $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ melts congruently. Owing to its congruent-melting behavior, a single crystal of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ with a size of $9 \times 5 \times 2$ mm³ has also been successfully grown from its stoichiometric melt (Fig. S4).

The structure of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ was determined by single-crystal X-ray diffraction (Table S1), which shows that $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ crystallizes in the monoclinic space group of *Cc* (No. 9). The asymmetric unit of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ contains three unique Ba, two unique Al, twelve unique B, and twenty-four unique O atoms (Table S2). All of the Al atoms are four-coordinated, forming $[\text{AlO}_4]$ tetrahedra. The B atoms exhibit two different coordination environments: $[\text{BO}_3]$ triangles and $[\text{BO}_4]$ tetrahedra, which are further connected *via* corner-sharing to form a $[\text{B}_2\text{O}_5]$ dimer, a $[\text{B}_4\text{O}_9]$ group, and two types of $[\text{B}_3\text{O}_7]$ groups with different symmetries (Fig. 1a). The $[\text{B}_2\text{O}_5]$ dimers alternately connect with the $[\text{B}_4\text{O}_9]$ groups to form a $^1\infty[\text{B}_6\text{O}_{13}]$ chain (Fig. 1b). The $[\text{B}_3\text{O}_7]$ rings are linked with the $[\text{AlO}_4]$ tetrahedra to create $^2\infty[\text{B}_3\text{AlO}_6]$ layers, namely A, B, A', and B' (Fig. 1c). The arrangement of the $^2\infty[\text{B}_3\text{AlO}_6]$ single layers in $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ is ABA'B'ABA'B'..., in which the A' (B') layers and the A (B) layers have a mirror image relationship along the *c*-axis. Furthermore, the A (A') and B (B') layers are connected by $^1\infty[\text{B}_6\text{O}_{13}]$ chains to generate $^2\infty[\text{B}_{12}\text{Al}_2\text{O}_{28}]$ double layers (Fig. 1d). These layers are separated by the Ba^{2+} cations. Meanwhile, Ba^{2+} cations also occupy the void space within the double layer. The Ba atoms are in sixfold and tenfold coordination environments, $[\text{Ba}(1)\text{O}_6]$, $[\text{Ba}(2)\text{O}_{10}]$, and $[\text{Ba}(3)\text{O}_{10}]$ polyhedra. In the structure, the B–O distances fall in the range of 1.336(16) to 1.538(18) Å. The Al–O

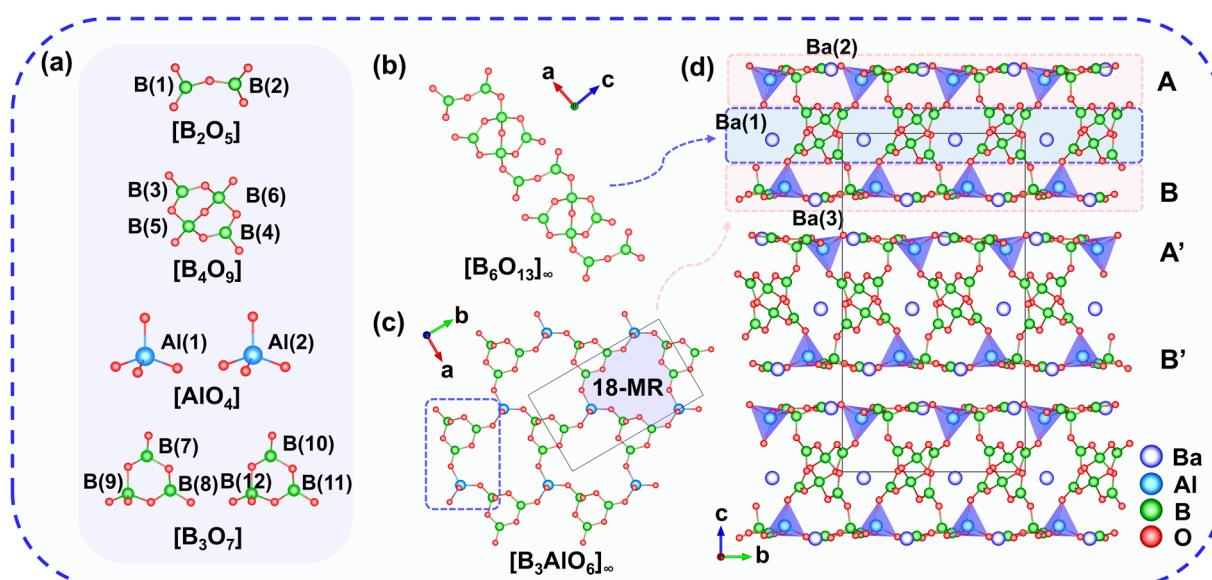


Fig. 1 (a) Basic building units (BBUs) of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$: $[\text{B}_2\text{O}_5]$, $[\text{B}_4\text{O}_9]$, $[\text{AlO}_4]$, and $[\text{B}_3\text{O}_7]$ units; (b) $^1\infty[\text{B}_6\text{O}_{13}]$ chain of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$; (c) $^2\infty[\text{B}_3\text{AlO}_6]$ layer of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$; (d) 2D double-layered structure of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$.



distances range from 1.728(9) to 1.780(9) Å. The Ba–O distances have a range from 2.593(9) to 3.133(9) Å (Table S3). The bond valence sum (BVS) analysis shows that the BVS values of Ba, Al, B, and O are 1.81–2.27, 3.08–3.13, 2.90–3.10, and 1.81–2.17, respectively (Table S2),³⁵ which are in agreement with their ideal oxidation states for each atom.

From the above structural analysis, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ possesses a similar layered structure to KBBF (single-layer structure) and SBBO (double-layer structures). Therefore, we conducted their structure comparison (Fig. 2). All these crystals feature layered structural units composed of coplanar $[\text{BO}_3]$ triangles ($[\text{B}_3\text{O}_7]$ for $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$) and tetrahedra ($[\text{BeO}_3\text{F}]$ for KBBF, $[\text{BeO}_4]$ for SBBO, and $[\text{AlO}_4]$ for $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$) (Fig. 2a–c). First, the $[\text{BeO}_4]$ tetrahedra were used to substitute the $[\text{BeO}_3\text{F}]$ tetrahedra in $^2\infty[\text{Be}_2\text{BO}_3\text{F}]$ single layers of KBBF, generating $^2\infty[\text{Be}(\text{BO}_3)\text{O}]$ single layers in SBBO. Furthermore, the $[\text{BeO}_4]$ tetrahedra and the $[\text{BO}_3]$ units in the $^2\infty[\text{Be}(\text{BO}_3)\text{O}]$ single layers of SBBO were replaced by $[\text{AlO}_4]$ tetrahedra and the $[\text{B}_3\text{O}_7]$ units to form $^2\infty[\text{B}_3\text{AlO}_6]$ single layers of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$. In SBBO, the $^2\infty[\text{Be}_2(\text{BO}_3)_2\text{O}]$ double layers are formed by the $^2\infty[\text{Be}(\text{BO}_3)\text{O}]$ single layers through bridged O atoms. Notably, the bridged O atoms between $^2\infty[\text{Be}(\text{BO}_3)\text{O}]$ single layers are replaced by $^1\infty[\text{B}_6\text{O}_{13}]$ chains in the $^2\infty[\text{B}_{12}\text{Al}_2\text{O}_{28}]$ double layers of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$. Meanwhile, compared with KBBF, the weak interlayer bonding ($\text{K}^+\text{–F}^-$ ionic bonds) is evidently reinforced in the other two compounds (Sr–O bonds for SBBO and Ba–O bonds for $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$). In particular, the interlayer spacing greatly decreases from 6.25 Å in KBBF to 3.43 Å in SBBO and 3.00 Å in $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ (Fig. 2d–f).^{25,36} The reduced interlayer space in $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ is expected to reinforce the bonding force and will

be conducive to overcoming the layering growth tendency. In $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$, the use of highly toxic beryllium oxide is effectively avoided by the substitution of the $[\text{BeO}_4]$ tetrahedra with the $[\text{AlO}_4]$ tetrahedra. Meanwhile, the structure contains planar $[\text{B}_3\text{O}_7]$ groups with greater microscopic polarizability than the $[\text{BO}_3]$ groups, which can be conducive to generating large SHG response and optical anisotropy.

For $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$, a $9 \times 5 \times 2$ mm³ crystal has been successfully obtained. The crystal can be further used to evaluate its optical properties. We measured its transparent windows from the UV to IR regions by using a polished $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ crystal wafer with 2 mm thickness, which demonstrates that $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ has a wide transparent spectral region from 190 to 3520 nm (Fig. 3a). This indicates that $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ is DUV transparent, and its UV absorption edge is comparable with those of other aluminoborate materials, such as $\text{K}_2\text{Al}_2\text{B}_2\text{O}_7$ (180 nm),³⁰ $\beta\text{-Rb}_2\text{Al}_2\text{B}_2\text{O}_7$ (<200 nm),²⁸ $\text{Cs}_2\text{Al}_2\text{B}_6\text{O}_{13}$ (185 nm),³¹ $\text{K}_3\text{Ba}_3\text{Li}_2\text{Al}_4\text{B}_6\text{O}_{20}\text{F}$ (190 nm),³² $\text{Rb}_3\text{Ba}_3\text{Li}_2\text{Al}_4\text{B}_6\text{O}_{20}\text{F}$ (198 nm),³³ etc. As $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ crystallizes in the NCS space group, powder SHG was measured using the Kurtz–Perry method. SHG intensity *versus* particle size revealed that $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ is phase-matchable at 1064 nm and has a large SHG intensity, $\sim 2.7 \times$ KDP (Fig. 3b and c). Furthermore, we investigated 11 reported NCS aluminoborate compounds containing alkali or alkaline earth metals (Table S4). When compared with other aluminoborates, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ exhibited the largest SHG response among these reported aluminoborates (Fig. 3d). As expected, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ demonstrates a good balance between SHG response and DUV transparency in aluminoborates (Fig. 3e).

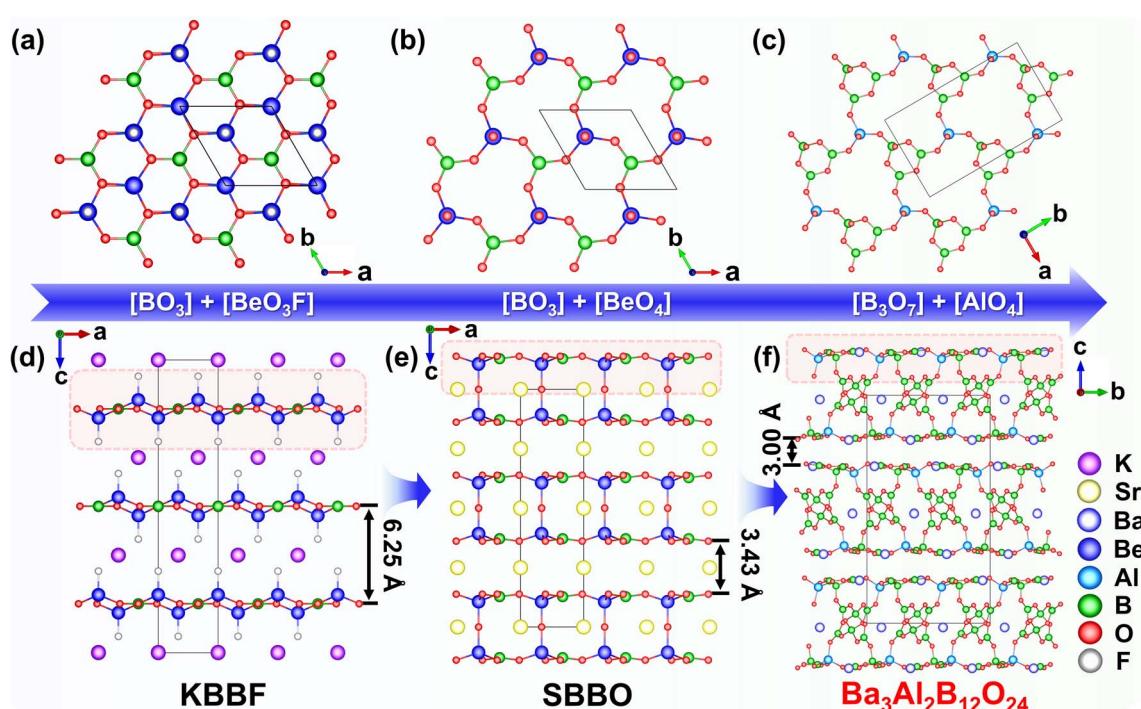


Fig. 2 (a)–(c) Layers in KBBF, SBBO, and $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$, respectively; (d)–(f) the structural evolution from the single-layer structure of KBBF to the double-layer structures of SBBO and $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$.

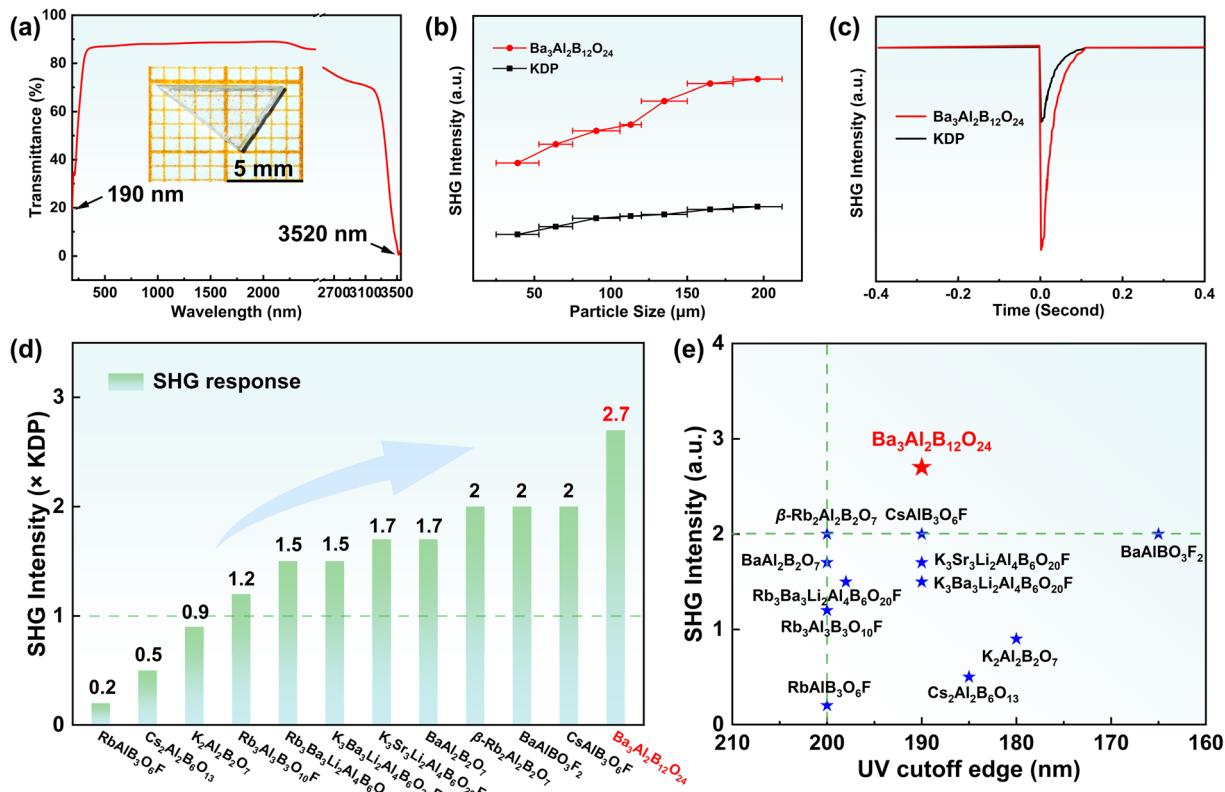


Fig. 3 (a) Ultraviolet-visible and infrared (UV-Vis-IR) transmission spectra of single crystals of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$; (b) and (c) powder SHG measurements at 1064 nm; (d) comparison of SHG responses of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ and reported aluminoborates; (e) scatter diagrams of SHG intensity and UV cutoff edges of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ and reported aluminoborates.

In order to investigate the origin of the large SHG response of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$, the dipole moment was calculated based on the BVS method to evaluate the contributions of individual groups to the SHG response (Table S5).³⁷ We found that the dipole moments of the B-O anionic groups in $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ are significantly larger than that of the Al-O anionic groups (Fig. 4a). This result indicates that the B-O anionic groups are important contributors to the large SHG response of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$. Meanwhile, the $[\text{B}_3\text{O}_7]$ units in the $^2\infty[\text{B}_3\text{AlO}_6]$ layers maintain an almost consistent coplanar arrangement, which is conducive to producing a large SHG response (Fig. 4b).

Besides these, birefringence is also important for NLO crystals to achieve a wide PM range. Therefore, the birefringence

value of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ was measured using a cross-polarizing microscope based on the formula $R = \Delta n \times d$, where R , Δn , and d are retardation, birefringence, and thickness of the crystal, respectively.³⁸ In the measurement, the crystal of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ with a thickness of 18.7 μm was used. And the I-order green interference color was observed using a cross-polarized light microscope (Fig. S5b). By comparing with the Michal-Levy chart, R is approximately 780 nm. Consequently, the birefringence of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ is about 0.042 in the visible region. The birefringence of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ was calculated based on the formulae $\varepsilon(\omega) = \varepsilon_1(\omega) + i\varepsilon_2(\omega)$ and $n^2(\omega) = \varepsilon(\omega)$. The calculated birefringence of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ at 1064 nm is 0.041, which is close to the measured birefringence (Fig. S5a).

To further explore the relationship between the structure and optical properties, the electronic structure was calculated via first-principles calculations based on the density functional theory (DFT).³⁹ $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ exhibits a direct band gap of 5.00 eV (Fig. 5a), which is smaller than the experimental result. This discrepancy can be attributed to the underestimation of the band gap (E_g) using DFT calculations.⁴⁰ Furthermore, we investigated the density of states (DOS) and partial density of states (PDOS) of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ (Fig. 5b). Since the optical properties of the compound are mainly determined by the states near the Fermi level,⁴¹ the top of the valence band (VB) and the bottom of the conduction band (CB) were analyzed. We found that the upper parts of the VB from -10 to 0 eV for $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$

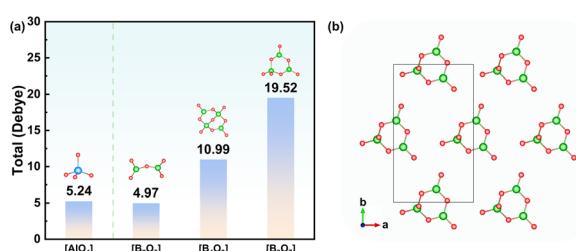


Fig. 4 (a) The sum of the dipole moments of all corresponding units $[\text{AlO}_4]$, $[\text{B}_2\text{O}_5]$, $[\text{B}_4\text{O}_9]$, and $[\text{B}_3\text{O}_7]$ in the unit cell of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$; (b) the arrangements of $[\text{B}_3\text{O}_7]$ groups in $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$.

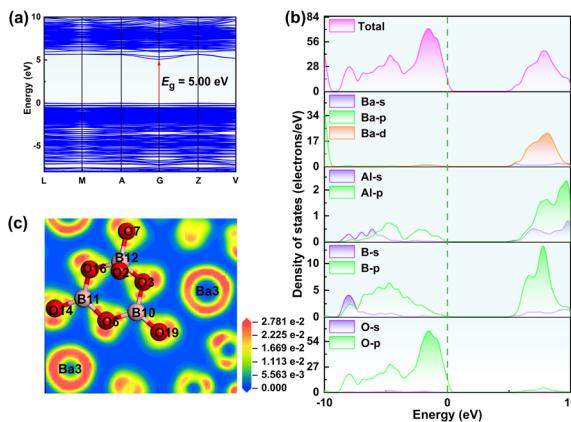


Fig. 5 (a) Electronic band structure of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$; (b) DOS and PDOS projected on the constitutional atoms of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$; (c) ELF map of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$.

mainly originate from the O 2p orbitals, which are mixed with a small part of B 2p orbitals and Al 3p orbitals. Meanwhile, the bottom of the CB from 0 to 10 eV primarily comprises B 2s2p orbitals with small amounts from the Ba 5d6s orbitals and Al 3s3p orbitals. From the above analysis, we can conclude that the large SHG response of $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ is mainly attributable to the B–O anionic groups. Based on the electron localization function (ELF) maps (Fig. 5c), we can intuitively observe that the clover-shaped asymmetric electron densities are arranged around the oxygen atoms to form the NLO-active $[\text{B}_3\text{O}_7]$ groups, which indicates that they have a positive contribution to the SHG response. Additionally, the nearby Ba cations form slightly distorted spherical asymmetric electron localized densities under the influence of the NLO-active $[\text{B}_3\text{O}_7]$ groups, which also contributes to the SHG response (Fig. S6). Similar situations have also been found in other compounds, such as $\text{Cs}_3\text{VO}(\text{O}_2)_2\text{CO}_3$ and $\text{Ba}_4\text{B}_{11}\text{O}_{20}\text{F}$.^{42,43} Therefore, the large SHG response in $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ originates mainly from B–O groups.

Conclusions

In summary, a new SBBO-like aluminoborate, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$, has been successfully synthesized by the chemical disubstitution strategy. $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ features $^2\infty[\text{B}_{12}\text{Al}_2\text{O}_{28}]$ double layers composed of $^2\infty[\text{B}_3\text{AlO}_6]$ single layers that are transformed from $^2\infty[\text{Be}(\text{BO}_3)\text{O}]$ single layers in SBBO through substituting $[\text{BeO}_4]$ tetrahedra and $[\text{BO}_3]$ triangles with $[\text{AlO}_4]$ tetrahedra and $[\text{B}_3\text{O}_7]$ units, respectively, which are further linked together *via* interlayer $^1\infty[\text{B}_6\text{O}_{13}]$ chains. This connection enables the reduced interlayer space in $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ to reinforce the bonding force and is conducive to overcoming the layering growth tendency. More importantly, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ satisfies the performance balance for NLO applications in the UV region among a large SHG response ($2.7 \times \text{KDP}$), a short UV absorption edge (<190 nm), and a suitable birefringence. Structural analyses and theoretical calculations indicate that the $[\text{B}_3\text{O}_7]$ units with coplanar configurations and well-ordered arrangements make a significant contribution to the excellent optical properties.

Meanwhile, $\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$ melts congruently, which is helpful for its crystal growth. These results demonstrate that site substitution based on the classical structure is an effective strategy for developing new UV and DUV NLO crystals.

Author contributions

X. R. L. performed the experiments, data analysis, theoretical calculations, and paper writing. H. W. Y. designed and supervised the experiments. H. P. W. provided major revisions of the manuscript. Z. G. H. supervised the optical experiments. J. Y. W. and Y. C. W. helped with the analyses of the crystallization process and the data. All the authors discussed the results and commented on the manuscript.

Conflicts of interest

There are no conflicts to declare.

Data availability

CCDC 2480204 ($\text{Ba}_3\text{Al}_2\text{B}_{12}\text{O}_{24}$) contains the supplementary crystallographic data for this paper.⁴⁴

The data supporting this article have been included as part of the SI. Supplementary information: Experimental section and additional tables and figures. See DOI: <https://doi.org/10.1039/d5sc03854e>.

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