

## RESEARCH ARTICLE

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# Deep-ultraviolet nonlinear-optical crystals LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> synthesized by ionic potential modulation towards uniform arrangement of PO<sub>4</sub> groups†

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The uniform arrangement of functional groups is a key factor in improving nonlinear properties in nonlinear-optical (NLO) materials, but currently there is no feasible and guiding strategy to modulate the uniform arrangement. Herein, we first apply the ionic potential concept to deep-ultraviolet (DUV,  $\lambda < 200$  nm) NLO phosphates for a uniform arrangement of PO<sub>4</sub> tetrahedral functional groups. Adopting Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub> with a non-uniform arrangement of PO<sub>4</sub> as a structural model, by removing low ionic potential Cs<sup>+</sup> and Li<sup>+</sup> successively, two DUV NLO crystals LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were synthesized. LiBePO<sub>4</sub> features a [Be<sub>3</sub>P<sub>3</sub>O<sub>18</sub>] six-membered ring constructed by alternate connection of BeO<sub>4</sub> and PO<sub>4</sub>, while BeP<sub>2</sub>O<sub>6</sub> exhibits two kinds of [PO<sub>3</sub>]<sub>∞</sub> helical chains bridged by BeO<sub>4</sub>. Remarkably, the arrangement of the PO<sub>4</sub> in LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> exhibits uniform evolution. As a result, LiBePO<sub>4</sub> exhibits an enhanced second-harmonic-generation (SHG) effect up to 4.3 times that of Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub>, while BeP<sub>2</sub>O<sub>6</sub> shows an even more enhanced SHG effect, reaching 7.0 that of Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub> (2.1 × KDP). Moreover, BeP<sub>2</sub>O<sub>6</sub> exhibits a short DUV absorption edge below 175 nm and the shortest SHG phase-matching output wavelength down to 211 nm. The universality of the new ionic potential modulation strategy is supported through analyzing known NLO materials containing alkali/alkaline-earth metal cations.

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## Introduction

The exploration of deep-ultraviolet (DUV,  $\lambda < 200$  nm or band-gap  $> 6.2$  eV) nonlinear-optical (NLO) crystals with large second-harmonic-generation (SHG) effects is of tremendous academic interest,<sup>1,2</sup> as a large SHG effect is beneficial for high laser frequency-conversion efficiency, which can be widely applied in a series of optoelectronic fields, *e.g.*, material micromachining, generation of entangled photon pairs, photo-

electron spectroscopy, *etc.*<sup>3–6</sup> In particular, those crystals with both a large SHG effect and short SHG phase-matching output wavelength (preferably in the short-wave UV ( $\lambda \leq 266$  nm), and even DUV region) are popular research areas because of essential applications in frontier technology. However, a large SHG effect and wide band-gap are naturally incompatible due to the fact that the SHG effect is inversely proportional to the band-gap in principle. Over the past few decades, a common strategy for achieving a large SHG effect has been to introduce SHG-active  $\pi$ -conjugated groups such as [BO<sub>3</sub>] and [CO<sub>3</sub>] to construct DUV NLO crystals, and this strategy has produced many crystals, such as KBe<sub>2</sub>BO<sub>3</sub>F<sub>2</sub> (KBBF),<sup>7</sup> NH<sub>4</sub>Be<sub>2</sub>BO<sub>3</sub>F<sub>2</sub>,<sup>8</sup>  $\beta$ -Rb<sub>2</sub>Al<sub>2</sub>B<sub>2</sub>O<sub>7</sub>,<sup>9</sup> NH<sub>4</sub>B<sub>4</sub>O<sub>6</sub>F,<sup>10</sup> and LiZn(OH)CO<sub>3</sub>.<sup>11</sup>

Recent studies show that non- $\pi$ -conjugated tetrahedral groups such as [SiO<sub>4</sub>], [PO<sub>4</sub>], and [SO<sub>4</sub>] with large HOMO–LUMO gaps can achieve high transmittance in the DUV region, and a series of tetrahedral DUV NLO crystals were thus synthesized, *e.g.* Li<sub>2</sub>SrSiO<sub>4</sub>,<sup>12</sup> Rb<sub>6</sub>Si<sub>10</sub>O<sub>23</sub>,<sup>13</sup> Ba<sub>3</sub>P<sub>3</sub>O<sub>10</sub>Cl,<sup>14</sup> Cs<sub>6</sub>Mg<sub>6</sub>(PO<sub>3</sub>)<sub>18</sub>,<sup>15</sup> RbNaMgP<sub>2</sub>O<sub>7</sub>,<sup>16</sup> Ba<sub>5</sub>P<sub>6</sub>O<sub>20</sub>,<sup>17</sup> (NH<sub>4</sub>)<sub>2</sub>Na<sub>3</sub>Li<sub>9</sub>(SO<sub>4</sub>)<sub>7</sub>,<sup>18</sup> *etc.* However, most of these DUV NLO crystals exhibit a small SHG effect because the constituent tetrahedral groups show nearly nonpolar high symmetry and therefore exhibit weak SHG activity. In order to improve the

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†Electronic supplementary information (ESI) available: Crystallographic data, XPS spectra, powder XRD, TG and DSC curves, IR spectra. CCDC 2384142 and 2384141 for LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub>. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d5qi00779h>

SHG performance, some strategies have been proposed, mainly including the introduction of (i) second-order Jahn–Teller effect  $d^0$  cations (e.g.  $\text{Mo}^{6+}$ ,  $\text{Ti}^{4+}$ ) or  $ns^2$  lone-pair cations (e.g.  $\text{Pb}^{2+}$ ,  $\text{Sn}^{2+}$ ),<sup>19–22</sup> (ii) anisotropic tetrahedral groups involving partially substituted oxygen atoms (e.g.  $[\text{BO}_{4-x}\text{F}_x]$ ,  $[\text{PO}_{4-x}\text{F}_x]$ ,  $[\text{SO}_{4-x}(\text{NH}_2)_x]$ ),<sup>23–26</sup> and (iii)  $\pi$ -conjugated groups (e.g.  $[\text{BO}_3]$ ).<sup>27,28</sup> However, the strategies all face seemingly insurmountable difficulties, such as an evident red-shift of the optical absorption edge, low thermal stability or returning to  $\pi$ -conjugated-characteristic NLO materials. Notably, that alkali/alkaline-earth metal cations with large HOMO–LUMO gaps have been only thought of as counterions for charge balance in these acentric structures, and their electrostatic interactions with surrounding tetrahedral groups are always overlooked. However, the interaction could affect the arrangement of tetrahedral groups, while a uniform arrangement can drive the generation of a large SHG effect according to anionic group theory.

In 2020, we reported two isostructural DUV NLO crystals  $\text{M}_4\text{LiBe}_4\text{P}_7\text{O}_{24}$  ( $\text{M} = \text{Cs}, \text{Rb}$ ) by combining alkali and alkaline-earth metal cations. Unfortunately, both crystals exhibit a weak SHG effect ( $\sim 0.3 \times \text{KDP}$ ).<sup>29</sup> Structurally, the weak SHG effect is mainly caused by a non-uniform arrangement of  $\text{PO}_4$  groups (Fig. S1†). Additionally, it is found that the characteristic differences between the three metal cations  $\text{Cs}^+$ ,  $\text{Li}^+$  and  $\text{Be}^{2+}$  in  $\text{Cs}_4\text{LiBe}_4\text{P}_7\text{O}_{24}$  (as an example) are size and charge number. The influence of the size and charge number on the electrostatic interactions of the respective ions is consistent with the ionic potential concept. In this regard, we take  $\text{Cs}_4\text{LiBe}_4\text{P}_7\text{O}_{24}$  as a structural model and put forward an ionic potential modulation strategy towards uniform arrangement of  $\text{PO}_4$ . By removing low ionic potential  $\text{Cs}^+$  and  $\text{Li}^+$ , two NLO-active crystals  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  were synthesized successively, and the arrangement of the  $\text{PO}_4$  exhibits uniform evolution. An increased SHG effect from  $\text{Cs}_4\text{LiBe}_4\text{P}_7\text{O}_{24}$  to  $\text{LiBePO}_4$  to  $\text{BeP}_2\text{O}_6$  is thus displayed. Notably,  $\text{BeP}_2\text{O}_6$  exhibits a large SHG effect of  $7.0 \times \text{Cs}_4\text{LiBe}_4\text{P}_7\text{O}_{24}$  ( $2.1 \times \text{KDP}$ ) and the shortest SHG phase-matching output down to the short-wave UV region of 211 nm. Furthermore, the universality of the ionic potential modulation strategy is supported through investigating known alkali/alkaline-earth metal NLO structures.

## Experimental

### Reagents

$\text{BeO}$  (99.99%),  $\text{NH}_4\text{H}_2\text{PO}_4$  (99.9%),  $\text{Li}_2\text{CO}_3$  (99.99%),  $\text{MoO}_3$  (99.9%) and  $\text{Li}_2\text{MoO}_4$  (99.9%) were commercially available and used as received. Caution: due to the high toxicity of  $\text{BeO}$  upon inhalation, all experiments were performed with sufficient ventilation.

### Synthesis

A polycrystalline sample of  $\text{LiBePO}_4$  was synthesized by a solid-state reaction method. Stoichiometric amounts of  $\text{Li}_2\text{CO}_3$ ,  $\text{BeO}$ , and  $\text{NH}_4\text{H}_2\text{PO}_4$  were mixed, ground completely, put in a platinum

crucible, and heated in a muffle furnace at 350 °C for 10 h. Then, the material was reground thoroughly, pressed into a pellet, and heated at 850 °C for 3d with several intermittent grindings.

### Crystal growth

We used a high-temperature solution method to grow the crystals of  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$ . For  $\text{LiBePO}_4$ , a mixture of  $\text{LiBePO}_4$  polycrystalline powder (0.5 mmol, 0.055 g),  $\text{MoO}_3$  (4 mmol, 0.576 g), and  $\text{Li}_2\text{MoO}_4$  (3 mmol, 0.522 g) was melted at 850 °C, then quickly cooled to 750 °C and held for 10 h. Then, the melt was cooled to 550 °C at a rate of 3 °C  $\text{h}^{-1}$  and finally cooled to 350 °C at a rate of 20 °C  $\text{h}^{-1}$ , and we switched off the furnace. Colorless millimeter-crystals were obtained after dissolving the material in water.

For  $\text{BeP}_2\text{O}_6$ , a mixture of  $\text{Li}_2\text{CO}_3$  (0.5 mmol, 0.037 g),  $\text{BeO}$  (1 mmol, 0.025 g) and  $\text{NH}_4\text{H}_2\text{PO}_4$  (3 mmol, 0.345 g) was melted at 850 °C and held for 12 h. Then, the melt was cooled down to 400 °C at a rate of 1 °C  $\text{h}^{-1}$ . Finally, the melt was cooled down to 350 °C at a rate of 10 °C  $\text{h}^{-1}$ , and we switched off the furnace. Colorless millimeter-crystals were obtained after dissolving the material in water.

### Single-crystal X-ray diffraction (XRD)

Single-crystal XRD data for  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  were collected at 293 K on a Bruker D8 Venture diffractometer equipped with a graphite monochromator using  $\text{Mo K}\alpha$  radiation ( $\lambda = 0.71073 \text{ \AA}$ ). Empirical absorption correction using spherical harmonics was performed using the SCALE3 ABSPACK scaling algorithm. The structure was solved by direct methods with the program SHELXS and refined by the full-matrix least-squares program SHELXL.<sup>30</sup> The structure was carefully checked using the program PLATON and no higher symmetries were found.<sup>31</sup> The details of relevant crystallographic data are summarized in Table 1. The atomic coordinates and equi-

**Table 1** Crystal data and structure refinements for  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$

Empirical formula	$\text{LiBePO}_4$	$\text{BeP}_2\text{O}_6$
Formula weight	221.84	166.95
Temperature/K	293(2) K	293(2) K
Crystal system	Monoclinic	Monoclinic
Space group	$Cc$	$P2_1$
$a/\text{\AA}$	16.331(1)	7.084(2)
$b/\text{\AA}$	9.151(1)	8.586(3)
$c/\text{\AA}$	15.965(1)	14.092(7)
$\alpha/^\circ$	90	90
$\beta/^\circ$	111.62(1)	94.06(3)
$\gamma/^\circ$	90	90
Volume/ $\text{\AA}^3$	2218.34(9)	855.0(6)
$Z$	16	8
$\rho_{\text{calc}}/\text{g cm}^{-3}$	2.117	2.594
$\mu/\text{mm}^{-1}$	0.788	0.953
$F(000)$	1728	656
Goodness-of-fit on $F^2$	1.03	1.036
$R_1, wR_2 [I \geq 2 \sigma(I)]^a$	0.0261, 0.0674	0.0328, 0.0793
$R_1, wR_2$ (all data) <sup>a</sup>	0.0316, 0.0717	0.0381, 0.0823
Flack parameter	0.02(3)	−0.01(5)

$$^a R_1 = \sum ||F_o| - |F_c|| / \sum |F_o|, \text{ and } wR_2 = [w(F_o^2 - F_c^2)^2 / w(F_o^2)^2]^{1/2}.$$

valent isotropic displacement parameters are listed in Table S1.† The selected bond distances and angles are shown in Table S2.†

### Powder X-ray diffraction (XRD)

Powder XRD data of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were collected using a Rigaku MiniFlex600 powder diffractometer equipped with Cu K<sub>α</sub> radiation ( $\lambda = 1.5418 \text{ \AA}$ ) in the  $2\theta$  angular range of 5–80° with a scan step width of 0.01° at 293 K. These experimental XRD patterns were in good agreement with the calculated patterns (Fig. S2 and S3†).

### Element analysis

Element analysis was performed by using a PerkinElmer Optima 8000 DV inductively coupled plasma optical emission spectrometer (ICP-OES). Two crystal samples were dissolved in perchloric acid at the boiling point for 0.5 h. ICP-OES measurements give molar ratios of 1.08(Li):0.95(Be):1.00(P) for LiBePO<sub>4</sub> and 1.00(Be):2.11(P) for BeP<sub>2</sub>O<sub>6</sub>, which are consistent with their formulae.

### X-ray photoelectron spectroscopy (XPS)

The chemical states and surface compositions of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were analyzed by a Thermo Scientific K-Alpha<sup>+</sup> X-ray photoelectron spectrometer with a monochromatic Al K<sub>α</sub> X-ray source (1486.6 eV) operating at 72 W (12 kV, 6 mA). All of the XPS spectra underwent background subtraction and were fitted using mixed Gaussian-Lorentzian peak shapes.

As seen in Fig. S4,† XPS spectra of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> demonstrate the presence of Li (for LiBePO<sub>4</sub>), Be, P, and O (Fig. S4a and f†). For LiBePO<sub>4</sub>, a peak is observed at 55.38 eV for Li 1s, ascribed to the Li(I) oxidation state (Fig. S4b†). The Be 1s XPS spectrum (Fig. S4c†) shows a peak at 113.50 eV, which is attributed to the Be(II) oxidation state. The peak at 133.46 eV (Fig. S4d†) corresponds to the P 2p binding energy for the +5 oxidation state. The O 1s XPS spectrum exhibits a peak at 531.54 eV (O<sup>2-</sup>) (Fig. S4e†). For BeP<sub>2</sub>O<sub>6</sub>, the Be 1s XPS spectrum (Fig. S4g†) shows a peak at 114.20 eV, which is attributed to the Be(II) oxidation state. The peak at 133.56 eV (Fig. S4h†) corresponds to the P 2p binding energy for the +5 oxidation state. The O 1s XPS spectrum exhibits a peak at 531.74 eV (O<sup>2-</sup>) (Fig. S4i†).

### Thermal behavior

Thermogravimetry (TG) and differential scanning calorimetry (DSC) analysis of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were carried out on a NETZSCH STA 449 F5 Jupiter simultaneous analysis system under flowing N<sub>2</sub>. About 15 mg of polycrystalline samples of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were separately placed in Pt crucibles, and heated from room temperature to 1300 °C (for LiBePO<sub>4</sub>) or 1100 °C (for BeP<sub>2</sub>O<sub>6</sub>) at a rate of 10 °C min<sup>-1</sup> under a flowing N<sub>2</sub> atmosphere.

TG and DSC curves of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> (Fig. S5†) show that weight losses in the TG curves were not obviously observed up to 1300 °C for LiBePO<sub>4</sub> and 1100 °C for BeP<sub>2</sub>O<sub>6</sub>. An endothermic peak at 1263 °C in the DSC curve for LiBePO<sub>4</sub>

was detected, while the endothermic peak of BeP<sub>2</sub>O<sub>6</sub> was not observed until 1068 °C, indicating the excellent thermostability. Further, polycrystalline samples of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were separately calcined at 1300 °C and 1100 °C for 0.5 h under an air atmosphere. Powder XRD of the residues (Fig. S2 and S3†) suggests that LiBePO<sub>4</sub> samples before and after melting are consistent, showing that it is a congruently melting compound, while BeP<sub>2</sub>O<sub>6</sub> samples before and after melting are inconsistent. According to the powder XRD pattern of the residue, BeP<sub>2</sub>O<sub>6</sub> after melting is transformed into another known polymorph BeP<sub>2</sub>O<sub>6</sub> and an unknown phase (Fig. S3b†), demonstrating that BeP<sub>2</sub>O<sub>6</sub> is an incongruently melting compound.

### Infrared (IR) spectroscopy

Fourier transform infrared (FTIR) spectra of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were recorded from KBr pellets in the range 4000–400 cm<sup>-1</sup> on a Nicolet Model 5DX spectrometer. As shown in Fig. S6,† the IR spectrum of LiBePO<sub>4</sub> indicates that the peak at 1055 cm<sup>-1</sup> is attributed to the P–O asymmetric stretching vibration of PO<sub>4</sub>. The peaks at 789, 715, and 611 cm<sup>-1</sup> are attributed to the P–O symmetric stretching vibrations of PO<sub>4</sub>. The peaks at 577 and 508 cm<sup>-1</sup> are attributed to the P–O bending vibrations of PO<sub>4</sub>. The IR spectrum of BeP<sub>2</sub>O<sub>6</sub> indicates that the peaks at 1287, 1174, 1109, 1030, and 995 cm<sup>-1</sup> are attributed to the P–O asymmetric stretching vibrations of PO<sub>4</sub>. The peak at 735 cm<sup>-1</sup> is attributed to the P–O symmetric stretching vibration of PO<sub>4</sub>. The peaks at 518 and 438 cm<sup>-1</sup> are attributed to the P–O bending vibrations of PO<sub>4</sub>. The IR spectra further indicate the presence of P–O units of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub>.

### SHG measurement

Powder SHG measurements of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were performed by the Kurtz–Perry method at 298 K. The measurement was carried out using a Q-switched Nd:YAG solid-state laser at 1064 nm and 532 nm, and the intensity of the frequency-doubled output emitted from the samples was measured using a photomultiplier tube. Polycrystalline samples LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were sieved into a series of distinct size ranges: 20–40 μm, 40–60 μm, 60–80 μm, 80–120 μm, 120–150 μm, 150–200 μm, and 200–300 μm. Polycrystalline KH<sub>2</sub>PO<sub>4</sub> (KDP) and β-BaB<sub>2</sub>O<sub>4</sub> (BBO) with the same particle size ranges were used as references. The SHG signal intensity of the LiBePO<sub>4</sub>/KDP and BeP<sub>2</sub>O<sub>6</sub>/KDP (or BBO) ratio was identified based on the same size (150–200 μm).

### Optical transmission spectrum

Optical transmission spectra of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were measured using a Shimadzu SolidSpec-3700 DUV spectrometer in the 175–800 nm region. Two polished single-crystals with sizes of 2.0 × 1.2 × 0.3 mm<sup>3</sup> for LiBePO<sub>4</sub> and 1.8 × 1.1 × 0.3 mm<sup>3</sup> for BeP<sub>2</sub>O<sub>6</sub> were separately used to perform the measurements.

### Birefringence measurement

Birefringences of LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were measured on a CNOPTec cross-polarizing microscope BK-POLR with an LED

light filter. The LED light source has an optical wavelength of  $590 \pm 3$  nm. The birefringence was calculated by using the following equation:  $R = d \times |N_e - N_o| = d \times \Delta n$ , where  $R$ ,  $d$ , and  $\Delta n$  represent the optical retardation, thickness of the crystal, and birefringence, respectively.

Interference colors of the selected  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  single-crystal samples are second-order yellow and second-order green based on a Michel–Levy diagram, and complete extinction can be achieved (Fig. S7a and S7b†). The tested crystals have retardations ( $R$ ) of 810 and 710 nm, with thicknesses ( $d$ ) of 45.13 and 25.23  $\mu\text{m}$ , respectively. Accordingly, the measured birefringences are 0.018 and 0.028 for  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$ , respectively.

### Bond-valence-sum (BVS) calculation

BVS calculations for  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  were performed to further validate the structural model, and the bond valence sums were calculated using the formula  $V_i = \sum S_{ij} = \sum \exp[(r_0 - r_{ij})/B]$ , where  $S_{ij}$  is the bond valence associated with the bond length  $r_{ij}$ , and  $r_0$  and  $B$  (usually 0.37) are empirically determined parameters.

### Computational method

Single-crystal structural data of  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  were used for the theoretical calculations. The electronic structures and optical properties were calculated using a plane-wave pseudo-potential method within density functional theory (DFT) implemented in the total energy code CASTEP.<sup>32,33</sup> Generalized gradient approximation (GGA) Perdew–Burke–Ernzerhof (PBE) was used as the exchange and correlation functional.<sup>34</sup> The norm-conserving pseudopotential was applied to present the core–electron interactions.<sup>35</sup> Li  $2s^1$ , Be  $2s^2$ , O  $2s^2 2p^4$ , and p  $3s^2 2p^3$  orbital electrons were considered as valence electrons. A cutoff energy of 850 eV determined the number of plane waves included in the basis sets. Monkhorst–Pack  $k$ -point sampling of  $2 \times 1 \times 4$  was used to perform the numerical integration of the Brillouin zone. During the optical property calculations, about 540 empty bands were set to ensure the convergence of SHG coefficients. Birefringence is calculated by subtracting the minimum refractive index from the maximum refractive index. The theoretical birefringences of  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  are 0.016 and 0.024, respectively, at 590 nm, which are close to the experimental results. The calculations of second-order NLO susceptibilities were based on length-gauge formalism within the independent particle approximation.<sup>36,37</sup>

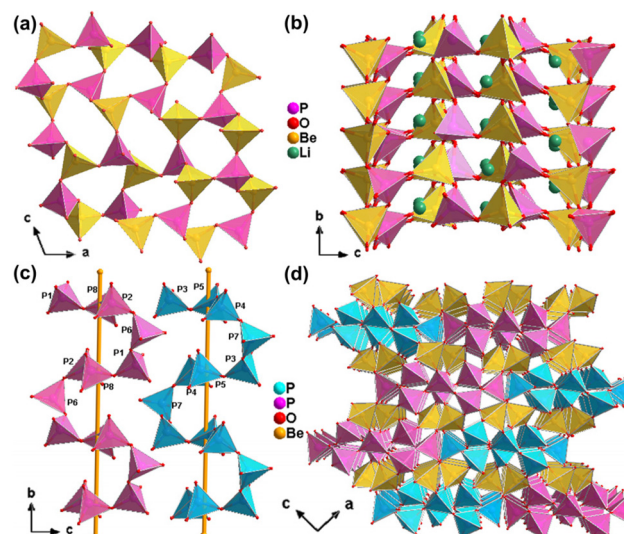
## Results and discussion

### Crystal structures

$\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  single-crystals were synthesized through high-temperature molten reactions, and both crystal structures were characterized by single-crystal XRD.  $\text{LiBePO}_4$  crystallizes in acentric and polar monoclinic space group  $Cc$  (no. 9), and contains eight crystallographically independent Li atoms,

eight Be atoms and eight  $\text{PO}_4$  groups, all of which are localized on crystallographically general positions with site occupancies of 1. Basic tetrahedral  $\text{BeO}_4$  and  $\text{PO}_4$  show typical bond distances of 1.606(3)–1.645(3) Å for Be–O and 1.525(1)–1.542(1) Å for P–O bonds. The material features a  $[\text{Be}_3\text{P}_3\text{O}_{18}]$  six-membered ring constructed by alternate connection of  $\text{BeO}_4$  and  $\text{PO}_4$ . The six-membered rings are fused by shared oxygen to generate  $[\text{BePO}_4]_\infty$  two-dimensional (2D) pseudo layers in the  $ac$  plane (Fig. 1a). The pseudo layers are interconnected through Be–O–P bonds to form a 3D structure with pores (Fig. 1b). It should be noted that  $c$  glide planes in the structure cause polarization cancellation along the  $b$ -axis and net polarization in the  $ac$  plane. Small four-coordinated  $\text{Li}^+$  fills in pores to balance charge (Fig. 1b). In comparison, in  $\text{Cs}_4\text{LiBe}_4\text{P}_7\text{O}_{24}$ , large high-coordinated  $\text{Cs}^+$  requires large coordination space that induces four-coordinated  $\text{Li}^+$  on large  $[\text{LiBe}_4\text{P}_7\text{O}_{24}]$  twelve-membered rings of framework layers (Fig. S1†).

$\text{BeP}_2\text{O}_6$  crystallizes in acentric and polar monoclinic space group  $P2_1$  (no. 4). There are four crystallographically independent Be atoms and eight  $\text{PO}_4$  groups, all of which are localized on crystallographically general positions with Wyckoff letter  $2a$ . These  $\text{PO}_4$  by corner-sharing generate two kinds of  $[\text{PO}_3]_\infty$  helical chains with a  $2_1$  axis along the  $b$ -axis (Fig. 1c). Adjacent  $[\text{PO}_3]_\infty$  chains are bridged by  $\text{BeO}_4$  to produce a 3D structure. Net polarization of the structure is along the  $b$ -axis, as deduced from its  $P2_1$  space group. Viewed along the  $b$ -axis (Fig. 1d), two  $[\text{PO}_3]_\infty$  chains both exhibit almost uniform stacking. In all  $\text{BeO}_4$  and  $\text{PO}_4$  units, Be–O (1.579(6)–1.633(6) Å) and P–O (1.434(4)–1.603(3) Å) bond lengths both vary in reasonable ranges. Validity of both structure models is checked through BVS calculations and XPS measurements, revealing their con-



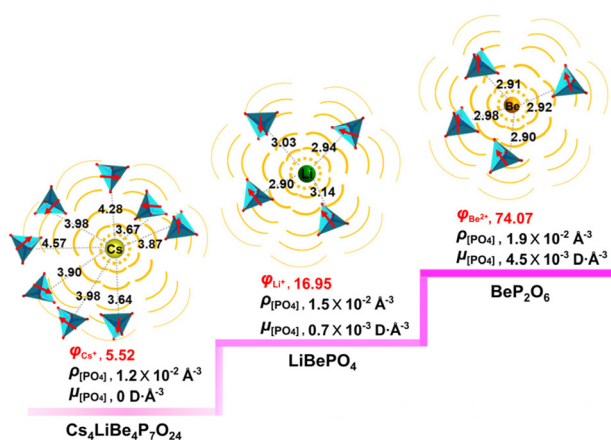
**Fig. 1** Crystal structures of  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$ .  $\text{LiBePO}_4$ : (a)  $[\text{BePO}_4]_\infty$  2D pseudo layer formed by  $[\text{Be}_3\text{P}_3\text{O}_{18}]$  six-membered rings. (b) 3D stacking of  $[\text{BePO}_4]_\infty$  pseudo layers viewed along the  $a$ -axis.  $\text{BeP}_2\text{O}_6$ : (c) two kinds of  $[\text{PO}_3]_\infty$  chains with  $2_1$  axes along the  $b$ -axis. (d) 3D stacking of  $[\text{PO}_3]_\infty$  chains bridged by  $\text{BeO}_4$  viewed along the  $b$ -axis.



stituent elements in rational oxide states (Table S1 and Fig. S4†).

### Ionic potential modulation model of PO<sub>4</sub> arrangement

Arrangement changes of the PO<sub>4</sub> from Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub> to LiBePO<sub>4</sub> to BeP<sub>2</sub>O<sub>6</sub> are related to the respective counter-balance cations Cs<sup>+</sup>, Li<sup>+</sup> and Be<sup>2+</sup>. The Pauling electronegativity of these elements increases in turn from Cs (0.79) to Li (0.98) to Be (1.57). In Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub>, Cs<sup>+</sup> has the strongest ionic bonding character and thus is regarded as the counter-balance cation of the structure, and the remaining ions Li<sup>+</sup> and Be<sup>2+</sup> mainly act as local charge balancing cations in the structure. Similarly, in LiBePO<sub>4</sub>, Li<sup>+</sup> is counter-balance cation of the structure and Be<sup>2+</sup> mainly acts as a local charge balancing cation. The size and charge number are the main differences between these cations. Their effect on PO<sub>4</sub> arrangement is studied using an ionic potential modulation model. According to the equation  $\varphi = z/r$  ( $\varphi$ , ionic potential;  $z$ , ionic charge number;  $r$ , ionic radius),  $z$  values of Cs<sup>+</sup>, Li<sup>+</sup> and Be<sup>2+</sup> are 1, 1 and 2, respectively, and  $r$  values (effective ionic radii) of these cations in Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub>, LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> are 0.18, 0.059 and 0.027 nm, respectively.  $\varphi_{\text{Li}^+}$  and  $\varphi_{\text{Be}^{2+}}$  are calculated to be 16.95 and 74.07, 3.1 and 13.4 times greater, respectively, than 5.52 for  $\varphi_{\text{Cs}^+}$ . The impacts of the alkali/alkaline-earth metal cations on the respective PO<sub>4</sub> groups are depicted in the radar maps in Fig. 2. As shown in Fig. 2, with increasing  $\varphi$  from  $\varphi_{\text{Cs}^+}$  to  $\varphi_{\text{Li}^+}$  to  $\varphi_{\text{Be}^{2+}}$ , the arrangement of these PO<sub>4</sub> groups around the cations exhibits uniform evolution. The uniform evolution can bring the following advantages: (i) the number of PO<sub>4</sub> groups per unit volume increases, which is confirmed by calculating the packing density ( $\rho$ ) of PO<sub>4</sub>, showing that  $\rho_{[\text{PO}_4]}$  increases from  $1.2 \times 10^{-2}$  to  $1.5 \times 10^{-2}$  to  $1.9 \times 10^{-2}$  per Å<sup>3</sup>; (ii) favorable superposition of microscopic hyperpolarizability of PO<sub>4</sub> that is beneficial for increasing the overall SHG effect is generated, and it is verified by calculating the dipole moment ( $\mu$ ) of PO<sub>4</sub>, revealing that  $\mu_{[\text{PO}_4]}$  increases from 0,  $0.7 \times 10^{-3}$  to  $4.5 \times 10^{-3}$  Debye (D) Å<sup>-3</sup>.

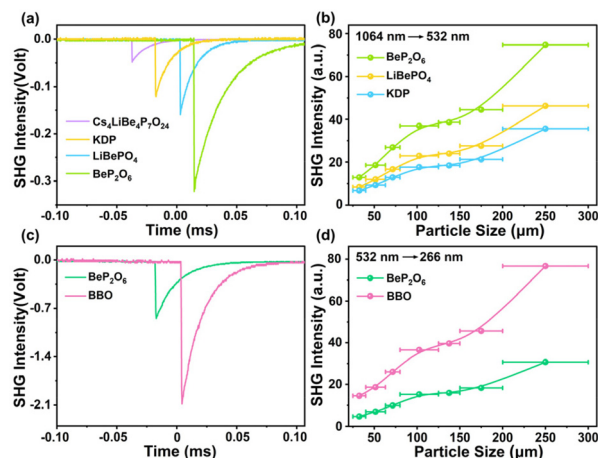


**Fig. 2** Ionic potential ( $\varphi$ ) modulation model of PO<sub>4</sub> arrangement. In the radar maps, the numbers and arrows represent the distance (Å) between the cations and dipole moment direction of PO<sub>4</sub>. Only PO<sub>4</sub> tetrahedra are displayed for clarity.

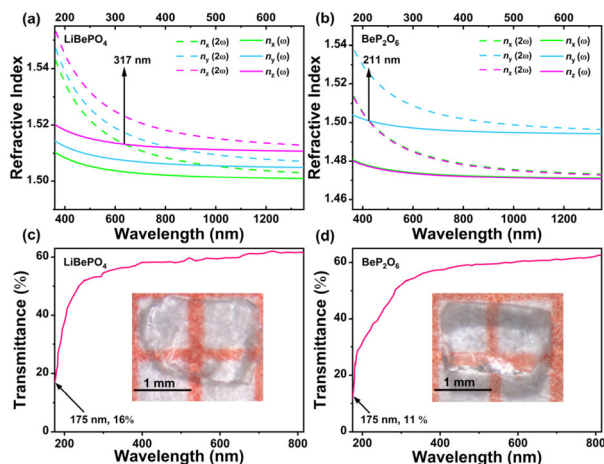
Essentially, ionic potential modulation is mainly due to electrostatic interactions between alkali/alkaline-earth metal cations and surrounding ions. The electrostatic interaction can be estimated using Coulomb's law based on a simple point-charge model, revealing that the electrostatic interactions between Li<sup>+</sup> and P<sup>5+</sup> in LiBePO<sub>4</sub> and between Be<sup>2+</sup> and P<sup>5+</sup> in BeP<sub>2</sub>O<sub>6</sub> are 1.8 and 3.7 times that between Cs<sup>+</sup> and P<sup>5+</sup> in Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub> (average distances of Cs and P, Li and P, and Be and P are 3.99, 3.00 and 2.93 Å according to Fig. 2). In addition, within the same structure, the ionic potential concept remains effective and applicable. In Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub> and LiBePO<sub>4</sub>, the arrangement uniformity of the PO<sub>4</sub> surrounding Cs<sup>+</sup>/Li<sup>+</sup>/Be<sup>2+</sup> and Li<sup>+</sup>/Be<sup>2+</sup> is improved in turn (Fig. S8 and S9†).

### SHG effect analysis

SHG effects based on the Kurtz–Perry method were measured at laser wavelengths 1064 and 532 nm, showing that the SHG effect of LiBePO<sub>4</sub> is  $1.3 \times$  benchmark KDP (KH<sub>2</sub>PO<sub>4</sub>) at 1064 nm (Fig. 3a), and 4.3 times that of Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub>. The SHG effect of BeP<sub>2</sub>O<sub>6</sub> further increases to  $2.1 \times$  KDP (Fig. 3a), reaching 7.0 times that of Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub>, and meanwhile, BeP<sub>2</sub>O<sub>6</sub> exhibits an SHG effect of  $0.4 \times$  BBO ( $\beta$ -BaB<sub>2</sub>O<sub>4</sub>) at 532 nm (Fig. 3c). The large SHG effect of BeP<sub>2</sub>O<sub>6</sub> is compared not only with reported non- $\pi$ -conjugated tetrahedral DUV NLO materials, such as Ba<sub>2</sub>NaClP<sub>2</sub>O<sub>7</sub> ( $0.9 \times$  KDP),<sup>38</sup> KMg(H<sub>2</sub>O)PO<sub>4</sub> ( $1.14 \times$  KDP),<sup>39</sup> BPO<sub>4</sub> ( $2.0 \times$  KDP),<sup>40</sup> Li<sub>2</sub>SrSiO<sub>4</sub> ( $0.3 \times$  KDP),<sup>13</sup> Li<sub>3</sub>AlSiO<sub>5</sub> ( $0.8 \times$  KDP),<sup>41</sup> Li<sub>2</sub>NaNH<sub>4</sub>(SO<sub>4</sub>)<sub>2</sub> ( $1.1 \times$  KDP),<sup>18</sup> and LiKSO<sub>4</sub> ( $1.2 \times$  KDP),<sup>42</sup> but also with well-known  $\pi$ -conjugated NLO materials, such as KBBF ( $1.21 \times$  KDP),<sup>7</sup> LiB<sub>3</sub>O<sub>5</sub> (LBO,  $3.0 \times$  KDP),<sup>43</sup> CsB<sub>3</sub>O<sub>5</sub> (CBO,  $2.7 \times$  KDP),<sup>44</sup> and CsLiBe<sub>6</sub>O<sub>10</sub> (CLBO,  $2.2 \times$  KDP).<sup>45</sup> Furthermore, LiBePO<sub>4</sub> reveals phase-matchable behavior in the visible region, whereas BeP<sub>2</sub>O<sub>6</sub> exhibits phase-matchable behavior in both visible and ultraviolet (UV) regions, as shown by the increasing SHG intensities with increasing particle size (Fig. 3b and d).



**Fig. 3** (a) SHG signals and (b) SHG signal intensity vs. particle size for LiBePO<sub>4</sub>, BeP<sub>2</sub>O<sub>6</sub>, and benchmarks Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub> and KDP at 1064 nm. (c) SHG signals and (d) SHG signal intensity vs. particle size for BeP<sub>2</sub>O<sub>6</sub> and benchmark BBO at 532 nm.



**Fig. 4** Refractive index dispersions of (a)  $\text{LiBePO}_4$  and (b)  $\text{BeP}_2\text{O}_6$ . Single-crystal transmission spectra of (c)  $\text{LiBePO}_4$  and (d)  $\text{BeP}_2\text{O}_6$ , with single-crystal photographs in the insets.

The phase-matching capacities of both structures are studied through experimental and theoretical methods. Calculated refractive index dispersions (Fig. 4a and b) reveal that the shortest SHG phase-matching output wavelengths are 317 and 211 nm for  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$ , respectively, consistent with the corresponding experimental observations of phase-matching in the 1064-to-532 nm and 532-to-266 nm second-harmonic processes (Fig. 3b and d), confirming their phase-matching capacity. In addition, calculated birefringences of  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  are 0.016 and 0.024, respectively, at 590 nm, and experimental birefringences determined using a cross-polarizing microscope at  $590 \pm 3$  nm through an optical retardation method show that the tested birefringences are 0.018 for  $\text{LiBePO}_4$  and 0.028 for  $\text{BeP}_2\text{O}_6$  (Fig. S7<sup>†</sup>), in general agreement with the calculated results. Remarkably, the shortest SHG phase-matching output of  $\text{BeP}_2\text{O}_6$  is down to the short-wave UV region of 211 nm, much better than non-phase-matching  $\text{BPO}_4$ , 450 nm of  $\text{LiCs}_2\text{PO}_4$ ,<sup>46</sup> and well-known borates LBO (277 nm), CBO (273 nm) and CLBO (236 nm) (Table 2).  $\text{BeP}_2\text{O}_6$  may be applied to frequency quadruple and quintuple the Nd:YAG laser output, producing 266 nm and 213 nm coherent light. In addition, optical transmission spectra show that  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  both have short DUV absorption edges below 175 nm (Fig. 4c and d), comparable to that of  $\text{Cs}_4\text{LiBe}_4\text{P}_7\text{O}_{24}$  (<190 nm).<sup>29</sup>

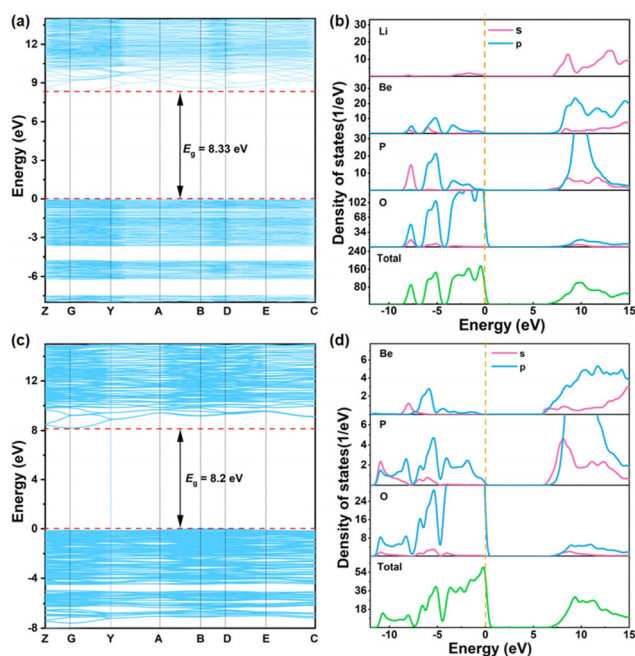
**Table 2** Comparison of DUV absorption edges ( $\lambda_{\text{DUV}}$ ), SHG effects at 1064 nm, and SHG phase-matching cutoff wavelengths ( $\lambda_{\text{PM}}$ ) of well-known crystals

Material	$\lambda_{\text{DUV}}$ (nm)	SHG effect ( $\times$ KDP)	$\lambda_{\text{PM}}$ (nm)
$\text{BPO}_4$	134	2.0	Nonphase-matching
$\text{LiCs}_2\text{PO}_4$	174	2.6	450
LBO	158	3.0	277
CBO	167	2.7	273
CLBO	180	2.2	236
$\text{BeP}_2\text{O}_6$	<175	2.1	211

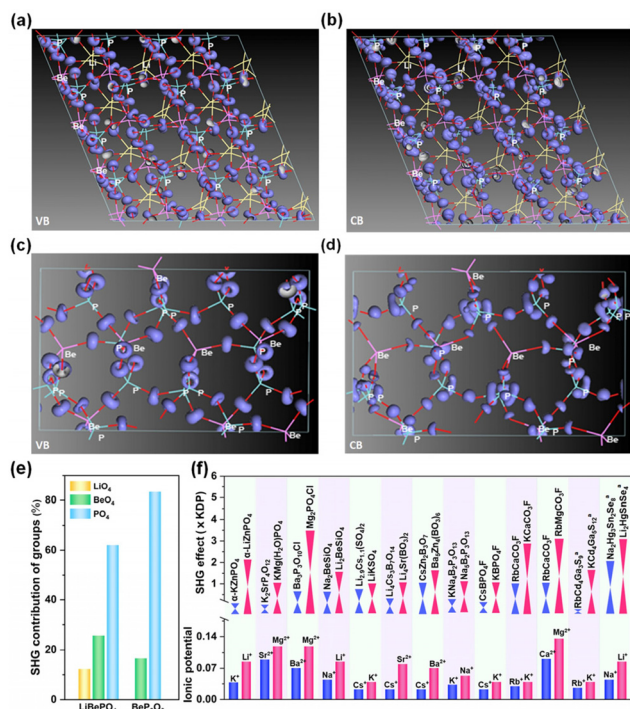
## Theoretical calculation

To gain deep insight into the origin of the good optical properties of  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$ , first-principles calculations were implemented. Band structures calculated using the hybrid exchange–correlation functional of HSE06 (Fig. 5a and c) indicate that  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$  have very wide band-gaps of 8.33 and 8.20 eV, corresponding to DUV absorption edges of 149 and 151 nm, respectively, in good support of the experimental results. Partial densities of states (Fig. 5b and d) show that the upper part of the valence-band consists mainly of P 3p, O 2p, and Be 2p electronic states, and the bottom part of the conduction-band is mainly composed of unoccupied P 3s 3p, O 2p, Be 2s 2p, and Li 2s (for  $\text{LiBePO}_4$ ) orbitals, revealing that all the chemical bonding interactions (including P–O, Be–O and Li–O) make contributions to the optical transitions of  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$ .

Calculated SHG coefficients show that the maximum tensors  $d_{15}$  for  $\text{LiBePO}_4$  and  $d_{23}$  for  $\text{BeP}_2\text{O}_6$  are 0.42 and 0.57  $\text{pm V}^{-1}$ , respectively, close to experimentally measured SHG effects. To intuitively determine the main SHG-contributed electronic orbitals, SHG-density analyses for both maximum SHG tensors were carried out. For  $\text{LiBePO}_4$  and  $\text{BeP}_2\text{O}_6$ , in the valence-band (Fig. 6a and c), the dominant SHG-contributed orbitals are O 2p nonbonding states, while in the conduction-band (Fig. 6b and d), unoccupied P 3s 3p and O 2p orbitals have a crucial impact on SHG effects, while the unoccupied Be 2p orbitals make a small contribution to the SHG effects. Furthermore, SHG contributions of different groups are summed based on the SHG-density analysis results shown in Fig. 6e, showing that  $\text{PO}_4$  is in both cases the leading group



**Fig. 5** Electronic band structures of (a)  $\text{LiBePO}_4$  and (c)  $\text{BeP}_2\text{O}_6$ . Density of states diagrams for (b)  $\text{LiBePO}_4$  and (d)  $\text{BeP}_2\text{O}_6$ .



**Fig. 6** SHG density maps of the valence band (VB) and conduction band (CB) for (a and b) LiBePO<sub>4</sub> and (c and d) BeP<sub>2</sub>O<sub>6</sub>. (e) SHG contribution of different groups in LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub>. (f) Applicability of the ionic potential modulation strategy in known alkali/alkaline-earth metal NLO materials. Blue represents cations with low ionic potential and their corresponding compounds, and red represents cations with high ionic potential and their corresponding compounds. <sup>a</sup> Represents the SHG effect based on benchmark AGS (AgGaS<sub>2</sub>).

producing SHG effects, accounting for 62.1% in LiBePO<sub>4</sub> and 83.4% in BeP<sub>2</sub>O<sub>6</sub>. The BeO<sub>4</sub> groups make small contributions to the SHG effects, accounting for 25.6% in LiBePO<sub>4</sub> and 16.6% in BeP<sub>2</sub>O<sub>6</sub>, respectively. The applicability of the ionic potential modulation strategy is investigated for known NLO structures containing alkali/alkaline-earth metal cations (Fig. 6f and Table S3<sup>†</sup>), and it is found that the strategy can be applied to these NLO materials well.

## Conclusions

An ionic potential modulation strategy is applied to tetrahedral DUV NLO phosphates that have long suffered from a weak SHG effect. Taking Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub> as a structural prototype, through continuous removal of low ionic potential Cs<sup>+</sup> and Li<sup>+</sup>, two DUV NLO crystals LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> were successfully synthesized. LiBePO<sub>4</sub> exhibits a 3D structure constructed from [Be<sub>3</sub>P<sub>3</sub>O<sub>18</sub>] six-membered rings composed of alternate BeO<sub>4</sub> and PO<sub>4</sub>, while BeP<sub>2</sub>O<sub>6</sub> features a 3D structure constructed from two kinds of [PO<sub>3</sub>]<sub>∞</sub> helical chains bridged by BeO<sub>4</sub>. Notably, the arrangement of PO<sub>4</sub> in LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub> exhibits uniform evolution. Thus, an increased SHG effect from Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub> to LiBePO<sub>4</sub> to BeP<sub>2</sub>O<sub>6</sub> is generated.

Furthermore, BeP<sub>2</sub>O<sub>6</sub> exhibits a large SHG effect of 7.0 × Cs<sub>4</sub>LiBe<sub>4</sub>P<sub>7</sub>O<sub>24</sub> (2.1 × KDP), the shortest SHG phase-matching output down to the short-wave UV region of 211 nm, and a short DUV absorption edge below 175 nm, showing that BeP<sub>2</sub>O<sub>6</sub> is a potential short-wave UV NLO crystal. Moreover, the new modulation strategy is universally supported through known alkali/alkaline-earth metal NLO materials. This work may not only be used to update the possible roles of alkali/alkaline-earth metal cations in NLO materials but also provide new opportunities for the precise design of material structures and properties.

## Author contributions

All authors have given approval to the final version of the manuscript.

## Data availability

The detailed crystallographic information and ESI figures and tables can be found in the ESI.<sup>†</sup> Crystallographic data are available *via* the Cambridge Crystallographic Data Centre (CCDC): 2384142 and 2384141 for LiBePO<sub>4</sub> and BeP<sub>2</sub>O<sub>6</sub>. The data supporting this article have been included as part of the ESI.<sup>†</sup>

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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