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LiMg(IO₃)₃: an excellent SHG material designed by single-site aliovalent substitution†

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An excellent second harmonic generation (SHG) material, LiMg(IO₃)₃ (LMIO), has been elaborately designed from Li₂M^{IV}(IO₃)₆ (M^{IV} = Ti, Sn, and Ge) by aliovalent substitution of the central M^{IV} cation followed by Wyckoff position exchange. The new structure sustains the ideal-alignment of (IO₃)⁻ groups. Importantly, LMIO exhibits an extremely strong SHG effect of roughly 24 × KH₂PO₄ (KDP) under 1064 nm laser radiation or 1.5 × AgGaS₂ (AGS) under 2.05 μ m laser radiation, which is larger than that of α -LiIO₃ (18 × KDP). The replacement of M^{IV} with Mg²⁺ without d-d electronic transitions induces an obviously larger band gap (4.34 eV) with a short absorption edge (285 nm). This study shows that single-site aliovalent substitution provides a new synthetic route for designing SHG materials.

Introduction

Noncentrosymmetric (NCS) or polar crystals have attracted scientific endeavors since they are potential candidates as SHG materials. 1,2 Metal iodates with polar units, $(IO_3)^-$ or $(IO_4)^{3-}$, are a class of important compounds which can exhibit excellent SHG performance.³ Numerous synthetic efforts have been made and afforded a number of iodates with outstanding SHG properties, 3-8 such as BiO(IO₃) (12.5 \times KDP)7a and GdI₅O₁₄ (15 \times KDP).8d However, generally, the larger SHG effects of iodates are usually accompanied by smaller band gaps. Recently, aliovalent substitution has become an effective approach for exploring novel SHG crystals.9,10 Aliovalent substitution of BiO(IO3) gave BiFSeO₃, which has the highest SHG effect among selenites reported.94 The novel route of aliovalent substitution involving three atom sites has expended d⁰-TM cations to post main group cations, such as from V⁵⁺ to Ga³⁺. 10 In this way, isostructural compounds sustained both large polarizability and a wide band gap. ¹⁰ For example, α - and β -Ba₂[GaF₄(IO₃)₂(IO₃)] (6 × KDP; 4.61 and 4.31 eV)10a have clearly larger band gaps and slightly weaker SHG signals than α - and β -Ba₂[VO₂F₄(IO₃)₂(IO₃)] $(9 \times \text{KDP}; 2.59 \text{ and } 2.55 \text{ eV}).^{9d}$

Recently, a serial of isostructural SHG iodates, namely ${\rm Li_2}$ - ${\rm M^{IV}(IO_3)_6}$ (${\rm M^{IV}}={\rm Ti}$, Sn, and Ge), were reported, whose structures feature well-arranged (${\rm IO_3}$) $^-$ units. The regulation of ${\rm Ti^{4+}}$,

Sn⁴⁺ and Ge⁴⁺ cations induces SHG effects shifting from 17×, $15\times$, to $32\times$ KDP and band gaps changing from 3.0, 3.9 to 3.89 eV for Ti, Sn and Ge compounds, respectively.11 Interestingly, in these structures, both Li⁺ and M⁴⁺ cations are octahedrally coordinated but they occupy different Wyckoff positions. The Li⁺ cation site on 2b and the LiO₆ octahedra are bridged by iodate groups into a 3D [Li(IO₃)₆]⁵⁻ framework. M⁴⁺ cations located at 2a are only 50% occupied and 0D [M^{IV}(IO₃)₆]²⁻ polyanions were formed. The full occupation of M^{IV} would be not consistent with the colorless and insulating crystals, which had been discussed in detail by P. Shiv Halasyamani. 11b We consider that aliovalent substitution of the M⁴⁺ cation with two divalent metal ions may result in LiMII(IO3)3 with a completely ordered structure, and this expands the family of this class of metal iodates with excellent SHG effects. Additionally, the use of alkaline-earth cations can avoid the unfavorable d-d electronic transitions for the transition metal ions and favors the blueshift of the absorption edge. By using such a single-site aliovalent substitution method, LiMg(IO₃)₃ (LMIO) was designed from Li₂M^{IV}(IO₃)₆ via replacing the defect-containing M⁴⁺ cation with an ordered Mg2+ cation followed by Wyckoff position exchange. In LMIO, a 3D $[Mg(IO_3)_6]^{4-}$ anionic framework is formed instead of 0D $[M^{IV}(IO_3)_6]^{2-}$ anions. LMIO displays an extremely large SHG response (24 \times KDP under 1064 nm laser radiation or 1.5 \times AGS under 2.05 μ m laser radiation), large band gap (4.34 eV), high LDT (46 \times AGS) and good thermal stability (>500 °C).

Results and discussion

Polar LMIO was prepared by hydrothermal reactions of LiCl, MgCl₂, I₂O₅ and 3% HCl solution at 230 °C for 3 days. Its purity

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was checked by PXRD study and EDS elemental analyses (Fig. S1 and $S2^{\dagger}$).

LMIO crystallizes in the NCS and polar space group P6₃ (no. 173). The asymmetric unit of LMIO contains one Li, one Mg, one I, and three O atoms. Both Li and Mg atoms occupy sites with 3-fold axis symmetry whereas the remaining atoms are located at the normal sites. Both Li⁺ and Mg²⁺ cations connect with six O atoms to form LiO6 and MgO6 octahedra, and each octahedron links with six (IO₃) groups which are ideally aligned along the c direction, leading to very large polarizability. The lengths of the Li-O bond (2.075(6) and 2.088(5) Å) and Mg-O bond (2.076(2) and 2.111(2) Å) are pretty close indicating that the distortions of LiO₆ and MgO₆ are very small. The I⁵⁺ atom is three coordinated in a triangular-pyramidal geometry with I-O bond lengths of 1.7966(19), 1.809(2) and 1.815(2) Å. BVS calculations exhibit the values of 1.14 (Li), 2.03 (Mg) and 5.06 (I), verifying that these cations are in oxidation states of +1, +2 and +5, respectively.

Neighboring LiO $_6$ octahedra are inter-linked into a 1D chain along the c-axis via face-sharing (Fig. 1a) whereas neighboring MgO $_6$ octahedra are bridged by iodate groups into a 3D [Mg(IO $_3$) $_6$] $^{4-}$ anionic framework with 1D tunnels of Mg $_6$ I $_6$ 12-membered rings (MRs) (Fig. 1b). The overall structure of LMIO can be viewed as the 1D chains of face-sharing LiO $_6$ octahedra inserted into the 1D tunnels of the 3D [Mg(IO $_3$) $_6$] $^{4-}$ anionic framework (Fig. 1c). Because that a process of Wyckoff position exchange occurred as the structural evolution from Li $_2$ M IV (IO $_3$) $_6$ to LMIO and different occupancy of the cation at the 2a position (fully occupied Li $^+$ in LMIO but 50% occupied M $^{4+}$ in Li $_2$ -M IV (IO $_3$) $_6$), it is interesting to note that there is in the structures of Li $_2$ M IV (IO $_3$) $_6$ (M = Ti, Sn, Ge), neighboring LiO $_6$ octahedra or [M IV (IO $_3$) $_6$] $^{2-}$ anions are not interconnected, that is they are isolated. 11

The ionic radii of the six-coordinated Li⁺, Mg²⁺, Ti⁴⁺, Sn⁴⁺, and Ge⁴⁺ are 0.076, 0.072, 0.0605, 0.069, and 0.053 nm, respectively.¹² Obviously, the ionic radius of Mg²⁺ is much closer to that of Li⁺ than that of M⁴⁺. Thus, Wyckoff position exchange occurred during the aliovalent substitution of M⁴⁺ with Mg²⁺ cations. In addition, our calculations of total energy show that LMIO (–12 816.63 eV) is more energy-favorable than the one adopting the Li₂M^{IV}(IO₃)₆ (M = Ti, Sn, Ge) structural type (–12814.03 eV). Attempts to replace Li⁺ in LMIO with Na⁺ to get an isostructural sodium phase were not successful, probably owing to the much larger size of Na⁺ (0.102 nm). We consider that a disordered M³⁺ cation, such as Al³⁺ and Ga³⁺ with 67% occupancy, may replace the M⁴⁺ cation in Li₂M^{IV}(IO₃)₆ (M = Ti, Sn, Ge).

The calculations of bond strain index (BSI) and global instability index (GII) values gave small values of 0.030 (BSI) and 0.079 (GII) for LMIO, which indicated that the structure of LMIO is unstrained and stable. The corresponding BSI and GII values of $\text{Li}_2\text{M}^{\text{IV}}(\text{IO}_3)_6$ (M = Ti, Sn, Ge) are much larger than those of LMIO (Table S3†), which are consistent with the disorder in $\text{Li}_2\text{M}^{\text{IV}}(\text{IO}_3)_6$ (M = Ti, Sn, Ge).

Thermogravimetric analysis (TGA) reveals the high thermal stability (>500 °C) of LMIO (Fig. 2). It shows two steps of weight loss in 500–650 °C (sharp) and 750–910 °C (smooth), and both are related to the release of $\rm I_2$ and $\rm O_2$. The DSC curves exhibit

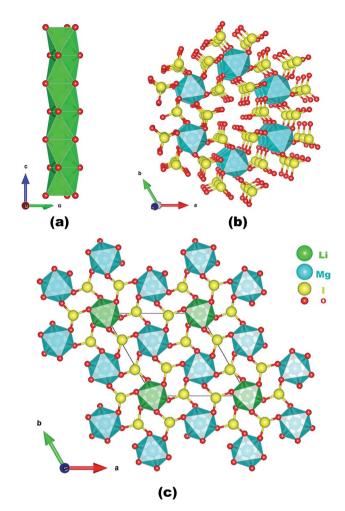


Fig. 1 A 1D chain of face-sharing LiO_6 octahedra along the c-axis (a); the 3D $[Mg(IO_3)_6]^{4-}$ anionic framework along the c-axis (b); and the view of the structure of LMIO along the c-axis (c).

three endothermic peaks at 610, 647 and 903 °C and one exothermic peak at 626 °C. Compared to $\text{Li}_2\text{M}^{\text{IV}}(\text{IO}_3)_6$ (M = Ti, Sn, Ge) (\sim 400 °C), LMIO has higher thermal stability.¹¹

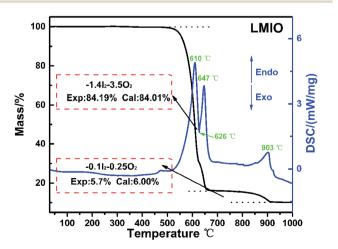


Fig. 2 TG and DSC analyses of LMIO.

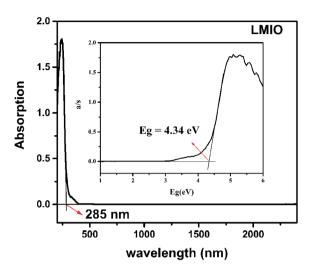


Fig. 3 UV-Vis-IR analysis of LMIO.

The IR spectrum shows that LMIO is transparent in the wavelength region of 2.5–11.2 μm (4000–890 cm $^{-1}$) (Fig. S3†). The UV-Vis-IR study revealed the absorption edge of 285 nm for LMIO (Fig. 3). Hence, the transparent range (0.28–11.2 μm) of LMIO covers the UV, Visible and mid-IR regions. The band gap of LMIO (4.34 eV) is significantly wider than those of Li₂- $M^{IV}(IO_3)_6$ (M = Ti, Sn, Ge) (3.0, 3.9 and 3.86 eV for Ti, Sn and Ge compounds, respectively). 11

Powder SHG experiments indicate that LMIO is a phase-matchable SHG material with remarkable SHG signals of 24 \times KDP under 1064 nm laser radiation or 1.5 \times AGS under 2.05 μ m laser radiation (Fig. 4). Compared with commercial α -LiIO $_3$ (18 \times KDP), LMIO possesses an obviously enhanced SHG signal. The SHG response of LMIO is also much larger than those of Li $_2$ Ti(IO $_3$) $_6$ (17 \times KDP) and Li $_2$ Sn(IO $_3$) $_6$ (15 \times KDP) but is slightly smaller than that of Li $_2$ Ge(IO $_3$) $_6$ (32 \times KDP). Eliminating the influence induced by the disorder effect, we adopt the Gaussian09 program to calculate the first-order

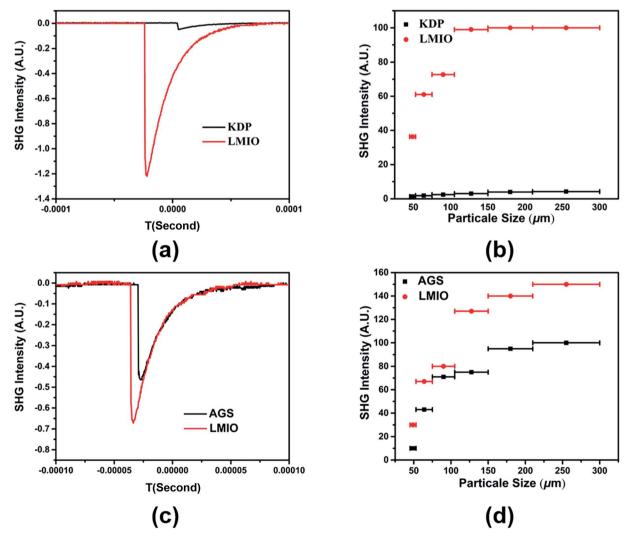


Fig. 4 Oscilloscope traces of the SHG signal (150–210 mm) of LMIO at (a) 1064 nm and (c) 2.05 μ m. Plots of measured SHG intensity vs. particle size of LMIO under laser radiation at (b) 1064 nm and (d) 2.05 μ m. KDP and AGS samples serve as the references for 1064 and 2.05 μ m laser radiation, respectively.

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hyperpolarizabilities of $[Mg(IO_3)_6]^{4-}$ and $[M^{IV}(IO_3)_6]^{2-}$ anions $(M=Ti, Sn \ and \ Ge)$, which are obtained to be 3047, 2982, 2602 and 1707 a.u. for Ge, Mg, Sn, and Ti, respectively, and such a trend is roughly consistent with results from the SHG measurements $(Li_2Ge(IO_3)_6 > LMIO > Li_2Sn(IO_3)_6 \approx Li_2Ti(IO_3)_6)$. Notably, LMIO is the only NLO material featuring excellent optical nonlinearity $(>20 \times KDP)$ and a near-UV absorption edge $(<300 \ nm)$.

LDT experiments revealed a large value of 101.86 MW cm⁻² for LMIO, which is about 46 times that of AGS (2.22 MW cm⁻²). This value is comparable to those of $\text{Li}_2\text{M}^{\text{IV}}(\text{IO}_3)_6$ (M = Ti, Sn, Ge) (49×, 41×, and 42 × AGS for Ti, Sn and Ge compounds, respectively) and $\alpha\text{-LiIO}_3$ (54 × AGS).^{11d}

Theoretical calculations based on DFT methods were applied for understanding the source of the outstanding SHG effect. Band structure calculations revealed that LMIO is an indirect band gap compound with a gap of 3.35 eV (Fig. S4†), which is much smaller than the measured value (4.34 eV). Hence scissor of 0.99 eV was applied during the subsequent optical property analyses.

The partial density of states (PDOS) was studied for the energy band assignment and bond interaction (Fig. 5). In the whole energy region, full overlapping of the electronic states between I atoms and O atoms is observed, indicating strong interactions in I–O bonds. The band gap of LMIO is determined by IO₃ units because the nonbonding orbitals from I and O

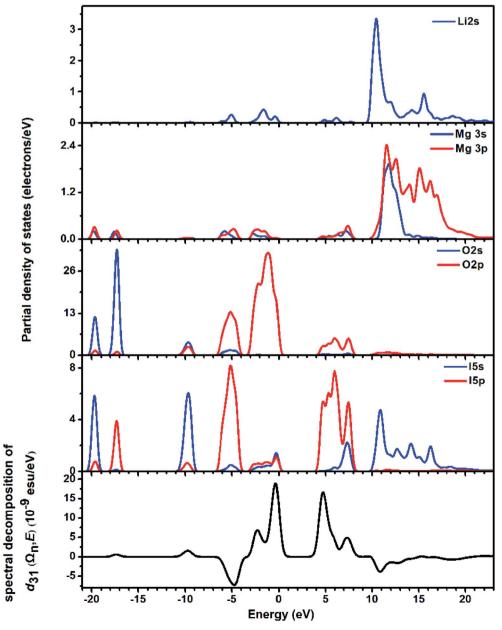


Fig. 5 The partial density of states (the upper four panels) and the spectral decomposition of d_{31} (the bottommost panel) for LMIO.

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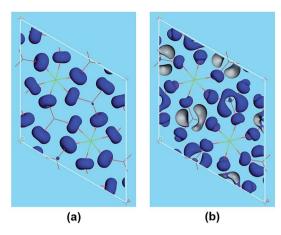


Fig. 6 The SHG density of d_{31} in the VB (a) and CB for LMIO (b).

atoms dominate the highest VB; meanwhile the empty orbitals from I and O atoms define the lowest CB.

Calculations of the SHG coefficient of LMIO revealed the largest SHG tensor d_{31} of 2.02×10^{-8} esu at 1064 nm. This result is consistent with our experimental value (24 × KDP). In addition, the refractive index curves of LMIO were recorded. It shows large anisotropy with large birefringence at 1064 nm (0.22), ensuring phase-matching abilities (Fig. S5†).

The SHG origin of LMIO has been further explored from spectral decomposition and SHG density. The spectral decomposition plot of d_{31} (the bottom panel in Fig. 5) indicates that the main SHG-contributed regions are located at the upper part of the VB and the lower part of the CB, corresponding to O-2p nonbonding states and unoccupied I-5p and O-2p states, respectively. There are small negative contributions from the VB regions of -6.3 to -3.6 eV and the CB region of 10-15 eV. Intuitively, for the SHG density (Fig. 6), the major SHGcontributors are the O-2p nonbonding orbitals in the VB and the unoccupied O-2p and I-5p orbitals in the CB. Accordingly, the IO₃ building units are the leading source of the large SHG coefficient. Numerically, the SHG-contributed percentages of Li⁺, MgO₆ and IO₃ groups are 0.43%, 8.94% and 90.6%, respectively. Hence, it is concluded that the very strong SHG response of LMIO originates predominantly from the idealalignment of (IO₃)⁻ groups.

Conclusions

In conclusion, via the aliovalent substitution of disordered M^{IV} (50% occupancy) with fully ordered Mg^{2^+} and the exchange of Wyckoff positions, a promising SHG material, LiMg(IO₃)₃ (LMIO), has been obtained. It displays a strong SHG effect (24 × KDP) with a phase matching ability and large band gap (4.34 eV). LMIO also has superior thermal stability (>500 °C) and LDT properties (46 × AGS). This novel approach of replacement of M^{IV} by M^{II} atoms further expands the family of iodate-based SHG materials. Our future research efforts will be devoted to aliovalent substitution of M^{IV} with defect containing M^{III} (66.7% occupancy) cations.

Conflicts of interest

The authors declare no competing financial interest.

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