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Theoretical study on the ferroelectric and light absorption properties of $\text{Li}_2\text{SbBiO}_6$ for harvesting visible light

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Ferroelectric oxides with large bandgaps have restricted applications in photovoltaic and photocatalytic fields. Based on recent experiments with the ferroelectric compound, LiSbO_3 , the stability and optoelectronic properties of a new ferroelectric compound, namely $\text{Li}_2\text{SbBiO}_6$, are investigated in this study. The calculated results demonstrate that $\text{Li}_2\text{SbBiO}_6$ satisfies the stability conditions of the elastic coefficients and phonon dynamics. $\text{Li}_2\text{SbBiO}_6$ maintains the ferroelectric polarization strength of LiSbO_3 and significantly reduces the bandgap, and thus has been explored for applications in photovoltaic and photocatalytic fields. $\text{Li}_2\text{SbBiO}_6$ is a new potential ferroelectric oxide for harvesting visible light owing to its suitable bandgap and a large hole–electron effective mass ratio.

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1. Introduction

Polar materials with non-centrosymmetric structures have pyroelectric, piezoelectric, ferroelectric, and nonlinear optical properties and have been found to exhibit unique applications in diverse fields.^{1–4} The conventional ferroelectric materials, LiNbO_3 and LiTaO_3 , belong to the noncentrosymmetric $R3c$ space group and show excellent ferroelectricity and unique features in the photovoltaic and photocatalytic decomposition of water.^{5–7} LiNbO_3 and LiTaO_3 are difficult to be promoted in photovoltaic and photocatalytic applications owing to their wide bandgaps. Some rare-earth ferroelectric oxides such as LuMnO_3 , YMnO_3 , and TbMnO_3 do not have very strong ferroelectric polarization strengths as LiNbO_3 and BiFeO_3 , but they exhibit excellent band gaps and high absorption coefficients, leading to significant interest in the ferroelectric photovoltaic field.^{8,9} The ferroelectric photovoltaic material, BiFeO_3 , can effectively maintain a high ferroelectric polarization strength and reduce the bandgap by doping to achieve high photovoltaic performance.¹⁰ The bandgap values of ferroelectric oxides with an LiNbO_3 structure are usually larger than 3 eV and poorly absorb visible light.^{11–15} LiSbO_3 with the LiNbO_3 structure recently synthesized by Inaguma *et al.* showed potential ferroelectricity and a high ferroelectric Curie temperature.¹⁶

Rich structural phase transitions can appear in LiSbO_3 at different pressures with large static dielectric tensor and insulating properties.¹⁷ LiSbO_3 with a $R3c$ structure has a large bandgap and mainly absorbs ultraviolet light according to our recent study.¹⁸ LiBiO_3 with the $Pccn$ space group can be prepared *via* the hydrothermal treatment method.¹⁹ Young *et al.* theoretically predicted that the energy of LiBiO_3 with a $R3c$ structure is relatively close to that of the $Pccn$ space group and is a potentially stable ferroelectric photovoltaic material with a strong bulk photovoltaic effect.²⁰ Bi-based materials have attracted attention for splitting water to produce hydrogen under visible light.^{21,22} In Bi-based compounds such as BiFeO_3 , BiVO_4 , Bi_2O_3 , KBiO_3 , LiBiO_3 , and NaBiO_3 excited by visible light, the bandgap can be reduced to below 3.0 eV due to hybridization between the O 2p and Bi 6s orbitals.^{23,24} The overlapping of O 2p and Bi 6s orbitals in Bi-based photocatalysts is beneficial for decreasing the bandgap to capture visible light.²⁵ The introduction of Cr ions into BiFeO_3 to form the $\text{Bi}_2\text{FeCrO}_6$ double perovskite with a $R3$ structure can also significantly reduce the bandgap and maintain a high ferroelectric polarization strength, effectively improving photovoltaic efficiency.²⁶ Considering Sb and Bi elements in the VA group and the unique 6s orbital of the Bi element and double perovskite $\text{Bi}_2\text{FeCrO}_6$ with the $R3$ structure, the introduction of Bi elements to form $\text{Li}_2\text{SbBiO}_6$ in the ferroelectric material LiSbO_3 can aid ferroelectric materials with the absorption of visible light. In this study, the stability and optoelectronic properties of $\text{Li}_2\text{SbBiO}_6$ are investigated using first-principles calculations. The calculation results show that $\text{Li}_2\text{SbBiO}_6$ satisfies stability conditions and maintains the ferroelectric

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strength of LiSbO_3 , while significantly reducing the bandgap to absorb visible light.

2. The calculation method and structure

First-principles calculations are based on the plane-wave basis set of the VASP software.^{27,28} The generalized gradient approximation and the plane pseudopotential wave method are used to deal with the interactions between electrons and ions.²⁹ The Perdew–Burke–Ernzerhof (PBE) functional is used as the exchange–correlation function.³⁰ The cutoff energy is chosen to be 520 eV, and the K -point sample including the Γ point is selected to be $6 \times 6 \times 6$. The lattice parameters and atomic positions are completely relaxed in the calculations. The electron self-consistency accuracy is 10^{-6} eV, and the atomic force is less than $0.01 \text{ eV } \text{\AA}^{-1}$. The Heyd–Scuseria–Ernzerhof (HSE) hybrid functional is employed to calculate the bandgap and complex frequency-dependent dielectric matrix in random phase approximation (RPA) schemes with an $8 \times 8 \times 8$ K -point sample.³¹ The elastic constants (C_{ij}) are calculated using the strain–stress method in the VASP package with the $8 \times 8 \times 8$ K -point sample.³² The standard berry phase method was applied to calculate ferroelectric polarization.^{33,34} A $2 \times 2 \times 2$ supercell containing 80 atoms was used to calculate the phonon frequency for $\text{Li}_2\text{SbBiO}_6$, and a $4 \times 4 \times 4$ K -point sample was used for this supercell. LiSbO_3 exhibits the $R3c$ symmetry based on experimental reports,¹⁶ while $\text{Li}_2\text{SbBiO}_6$ shows the $R3$ symmetry by Bi substitution for Sb similar to the $\text{Bi}_2\text{FeCrO}_6$ double perovskite, as shown in Fig. 1. The calculated lattice parameters are shown in Table 1, where the lattice parameters

Table 1 The calculated lattice parameters (lattice constant and angle α)

	LiSbO_3	LiSbO_3	LiBiO_3	LiBiO_3	$\text{Li}_2\text{SbBiO}_6$
a (Å)	5.39 (ref. 16)	5.46	5.67 (ref. 20)	5.71	5.60
α (°)	56.4 (ref. 16)	56.6	56 (ref. 20)	56.1	56.3

of $R3c$ - LiSbO_3 and $R3c$ - LiBiO_3 are in good agreement with the experimental¹⁶ and theoretical values,²⁰ respectively.

3. The stability of $\text{Li}_2\text{SbBiO}_6$

$R3c$ - LiSbO_3 has six independent elasticity coefficients, while $R3$ - $\text{Li}_2\text{SbBiO}_6$ has seven independent elasticity coefficients. The calculated elastic coefficients are shown in Table 2. The elastic coefficients of LiSbO_3 satisfy the inequalities of the stability conditions of the $R3c$ structure:³⁵

$$C_{11} - C_{12} > 0,$$

$$C_{13}^2 < 0.5 \times C_{33} \times (C_{11} + C_{12}),$$

$$C_{14}^2 < 0.5 \times C_{44} \times (C_{11} - C_{12}),$$

$$C_{44} > 0.$$

The seven independent elastic coefficients of $\text{Li}_2\text{SbBiO}_6$ also satisfy the stability conditions of the $R3$ space group:³⁵

$$C_{11} > |C_{12}|,$$

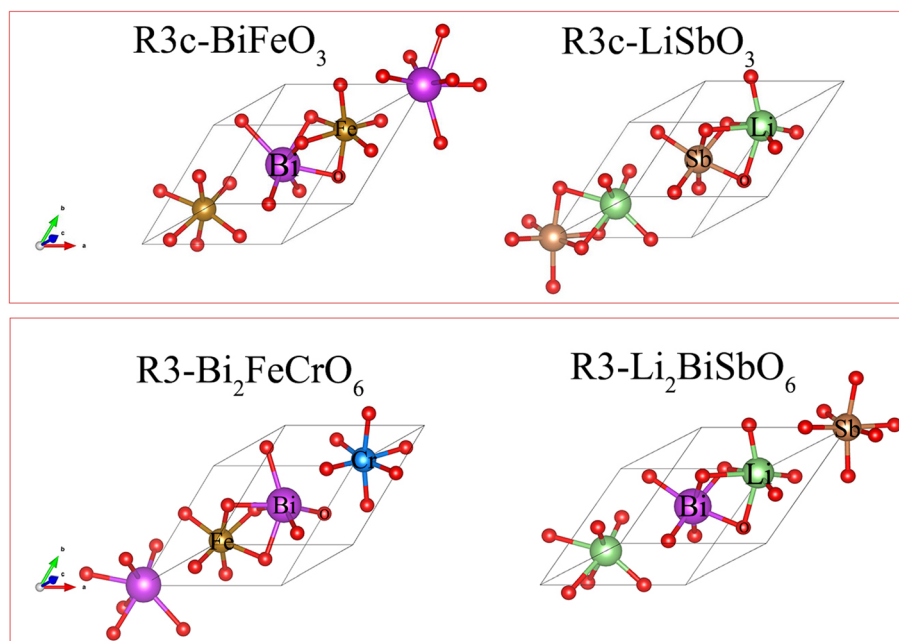


Fig. 1 The structures of LiSbO_3 and $\text{Li}_2\text{SbBiO}_6$.



Table 2 The calculated elastic coefficients (GPa)

	C_{11}	C_{12}	C_{13}	C_{14}	C_{33}	C_{44}	C_{66}
LiSbO ₃ (present)	245.3	52.0	86.4	−23.4	258.6	112.9	—
LiSbO ₃ (ref. 18)	283.6	56.2	104.0	−28.8	308.7	135.0	—
LiSbO ₃ (10 GPa) (ref. 18)	322.3	77.8	130.3	−32.2	336.4	150.8	—
LiSbO ₃ (10 GPa) (ref. 17)	289.6	75.0	119.8	−28.5	304.1	135.9	—
LiSbO ₃ (12 GPa) (ref. 17)	295.8	79.3	124.6	−28.9	309.8	138.3	—
Li ₂ BiSbO ₆	196.7	50.2	68.7	−16.2	200.1	84.3	73.2

Table 3 The calculated eigenvalue matrices of elastic coefficients (GPa)

	λ_1	λ_2	λ_3	λ_4	λ_5	λ_6
LiSbO ₃	79.7	100.7	131.1	156.2	207.6	396.7
Li ₂ BiSbO ₆	61.6	78.3	101.0	124.2	159.1	318.7

$$C_{13}^2 < 0.5 \times C_{33} \times (C_{11} + C_{12}),$$

$$C_{14}^2 + C_{15}^2 < 0.5 \times C_{44} \times (C_{11} - C_{12}),$$

$$C_{44} > 0.$$

The elasticity coefficient of *R3c*-LiSbO₃ calculated using the PBE function is smaller than that of the SCAN function (Table 2),¹⁸ and the elasticity coefficient becomes larger under high pressure conditions according to reports in the literature.^{17,18} All elasticity coefficients of *R3c*-LiSbO₃ and *R3*-Li₂SbBiO₆ in Table 2 meet the stability conditions. The eigenvalue matrices of the elastic coefficients of LiSbO₃ and Li₂SbBiO₆ in Table 3 are greater than 0 and satisfy the mechanical stability condition. The phonon frequencies and phonon density of states were calculated to verify the stability of LiSbO₃ and Li₂SbBiO₆ (Fig. 2). The results show that they are potentially stable structures with no imaginary frequencies throughout the Brillouin zone. Li₂SbBiO₆ has a smaller elasticity coefficient than LiSbO₃, which implies that mechanical properties, such as Young's modulus, bulk modulus, and hardness, may be modulated by the mutual

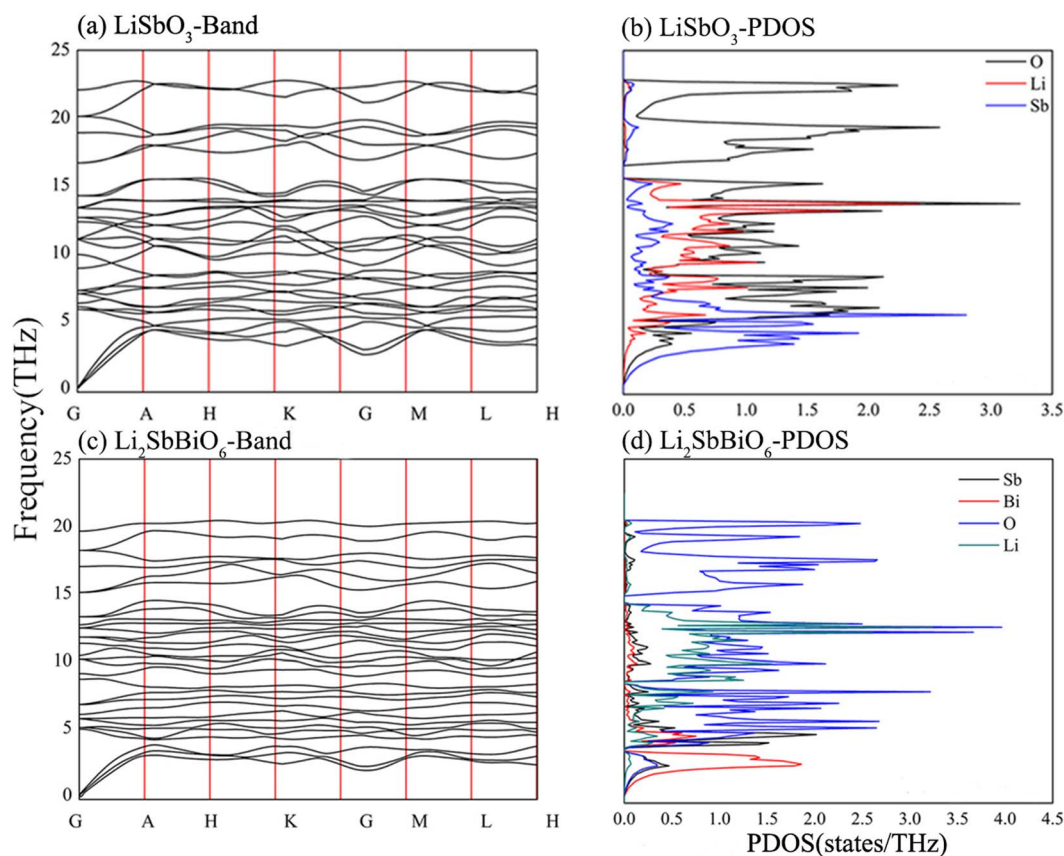


Fig. 2 The calculated phonon frequency and phonon density of states.



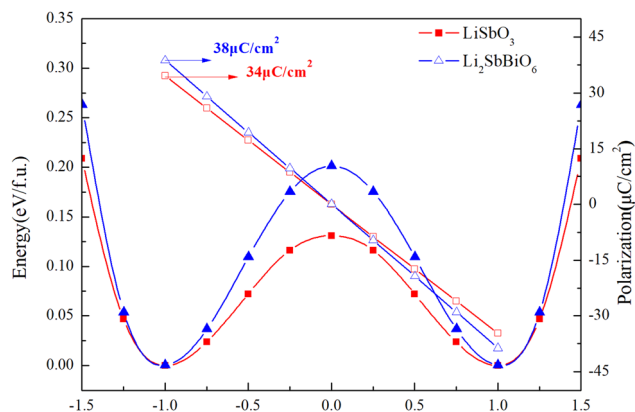


Fig. 3 The calculated energy and polarization as the position of the atom changes.

doping of Sb and Bi. The calculated phonon density of state shows that the phonon frequency of $\text{Li}_2\text{SbBiO}_6$ shifts toward lower frequencies as compared to that of LiSbO_3 , which is in line with the trend of the reduced elasticity coefficient.

4. Ferroelectricity analysis

The ferroelectric polarization strengths of LiSbO_3 and $\text{Li}_2\text{SbBiO}_6$ are calculated using the berry phase method. As shown in Fig. 3, the energy curves and ferroelectric polarization strengths are calculated by the linear interpolation of the atomic positions between the centrosymmetric and polarized structures. The centrosymmetric and polarized structures for LiSbO_3 are $R\bar{3}c$ and $R3c$, respectively, while $R\bar{3}$ and $R3$ structures are chosen for $\text{Li}_2\text{SbBiO}_6$. Displacement-type ferroelectric materials can form energy double potential well curves between the ferroelectric (polar) and paraelectric (centrosymmetric) states. The calculated energies of LiSbO_3 and $\text{Li}_2\text{SbBiO}_6$ show significant ferroelectric double potential wells. Interestingly, the calculated ferroelectric polarization values of LiSbO_3 and $\text{Li}_2\text{SbBiO}_6$ are $34 \mu\text{C cm}^{-2}$ and $39 \mu\text{C cm}^{-2}$, respectively. Thus, the ferroelectric polarization strength of $\text{Li}_2\text{SbBiO}_6$ is slightly greater than that of

LiSbO_3 . According to our previous study, Li–O atomic interactions play a significant role in the ferroelectricity of LiSbO_3 .¹⁸ We further calculated the effect of the ionic movement on the ferroelectricity of $\text{Li}_2\text{SbBiO}_6$ (Fig. 4). The movement of Li–O atoms induces the double potential well shift and promotes the ferroelectric stability in $\text{Li}_2\text{SbBiO}_6$. This is consistent with the results of the phonon density of states calculations (Fig. 2), where evidently the Li–O phonon coupling effect is stronger in the high phonon frequency area.

5. Discussion of photoelectric properties

As shown in Fig. 5, LiSbO_3 and $\text{Li}_2\text{SbBiO}_6$ present the characteristics of an indirect bandgap material with 3.2 and 1.9 eV bandgap values, respectively. The calculated bandgap of LiSbO_3 using the HSE06 functional is close to the value (3.4 eV) of the MBJ functional as-obtained in our earlier report.¹⁸ The energy band of $\text{Li}_2\text{SbBiO}_6$ has been calculated using the HSE06 + SOC (spin orbit coupling) approach due to the heavy Bi element. The calculation results from Fig. 6 show that the energy band structure of $\text{Li}_2\text{SbBiO}_6$ remains basically unchanged by the SOC effect, and the bandgap increases slightly to 2.1 eV. Considering the calculation of HSE06 + SOC is relatively expensive and HSE06 calculations can usually give accurate results in general, the HSE06 approach is mainly used to calculate the optical properties and electronic density of states of $\text{Li}_2\text{SbBiO}_6$. The bandgap of $\text{Li}_2\text{SbBiO}_6$ maintains the nature of the indirect bandgap of LiSbO_3 , which is unfavorable for photovoltaic performance, but the introduction of Bi ions can significantly reduce the bandgap to absorb visible light. From the density of states calculations (Fig. 5), the valence charges of both LiSbO_3 and $\text{Li}_2\text{SbBiO}_6$ mainly comprise O-2p orbitals. It is also found that the antibonding states formed by O-2p and Bi-6s orbitals significantly reduce the bandgap of $\text{Li}_2\text{SbBiO}_6$. The absorption coefficients ($\alpha(\omega)$) of $\text{Li}_2\text{SbBiO}_6$ were acquired using the frequency-dependent dielectric matrix:

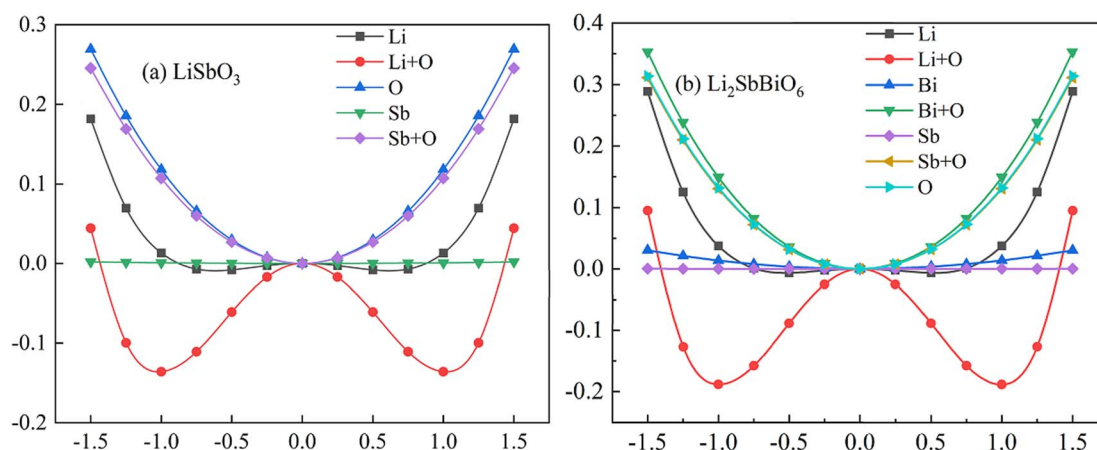


Fig. 4 The calculated energy as the position changes with the movement of different atoms.



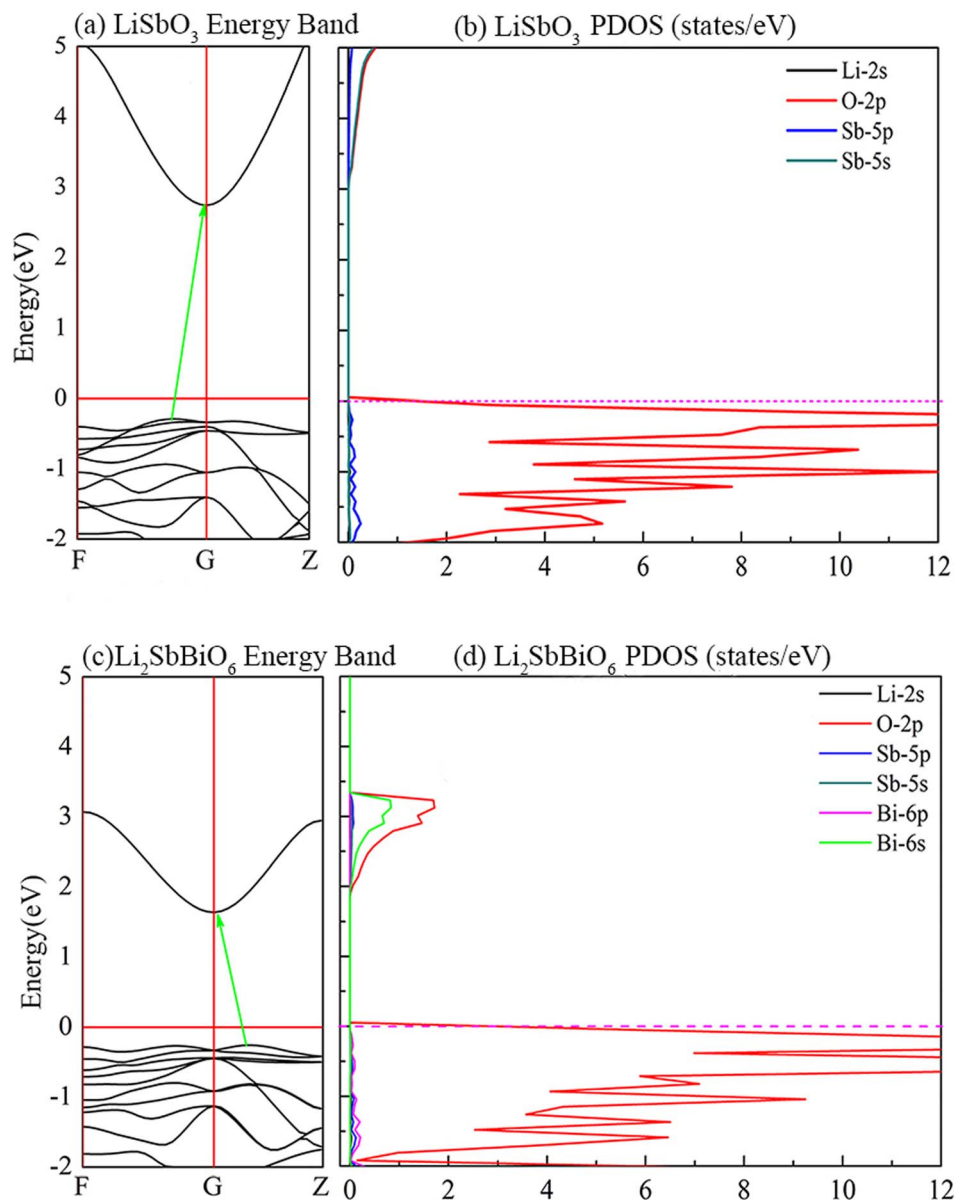


Fig. 5 The calculated energy band structure and density of states.

$$\alpha(\omega) = \frac{2\omega}{c} \left(\frac{\sqrt{\varepsilon_i(\omega)^2 + \varepsilon_r(\omega)^2} - \varepsilon_r(\omega)}{2} \right)^{\frac{1}{2}}$$

where ω is the incident light frequency, and c is the speed of light in a vacuum. The imaginary and real parts of the dielectric function are denoted by ε_i and ε_r , respectively. The calculated absorption coefficient of $\text{Li}_2\text{SbBiO}_6$ is shown in Fig. 7. LiSbO_3 does not absorb visible light because of its large bandgap. $\text{Li}_2\text{SbBiO}_6$ can significantly absorb visible light and is a potential ferroelectric photovoltaic material. The bandgap of $\text{Li}_2\text{SbBiO}_6$ using the HSE06 calculation is 1.9 eV, which is close to that of the photocatalytic materials BiOI ³⁶ and LiBiO_3 (ref. 37) with

applications in photocatalytic water splitting. The bandgap value of $\text{Li}_2\text{SbBiO}_6$ is not as favorable as that of rare-earth ferroelectric oxides for absorbing visible light,^{8,9} but the ferroelectric polarization strength of $\text{Li}_2\text{SbBiO}_6$ is relatively large, which could be favorable for driving carrier separation by the depolarization field. A ferroelectric material with an internal electric field can reduce the probability of carrier recombination and improve the efficiency of photogenerated carriers. We calculated the effective mass to investigate the photoelectric properties. The expression for effective mass is $m^*(k) = \pm \hbar^2 [\partial^2 E(k) / \partial^2 k]^{-1}$. The calculation results of the effective mass and ferroelectric polarization strength are shown in Table 4. The calculated electron effective mass of LiSbO_3 and $\text{Li}_2\text{SbBiO}_6$



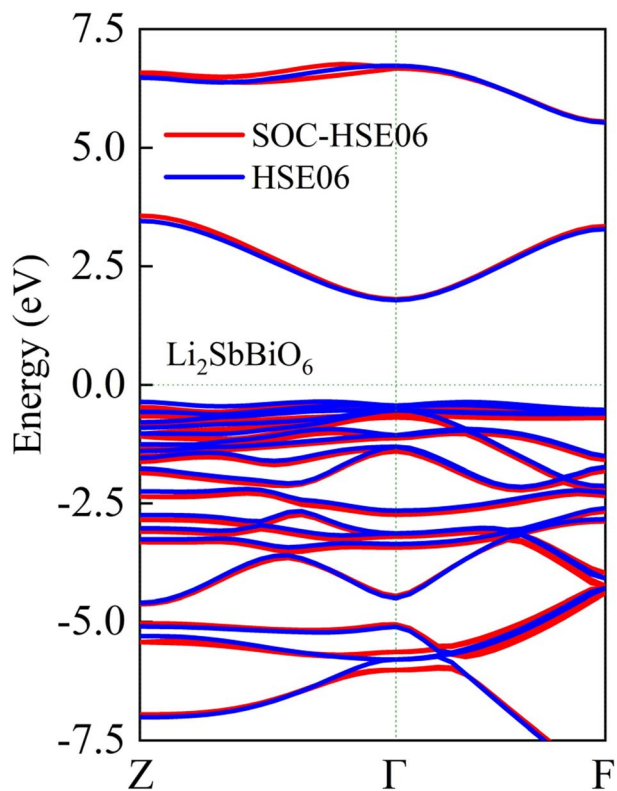


Fig. 6 The calculated energy band of $\text{Li}_2\text{SbBiO}_6$ with and without the SOC effect.

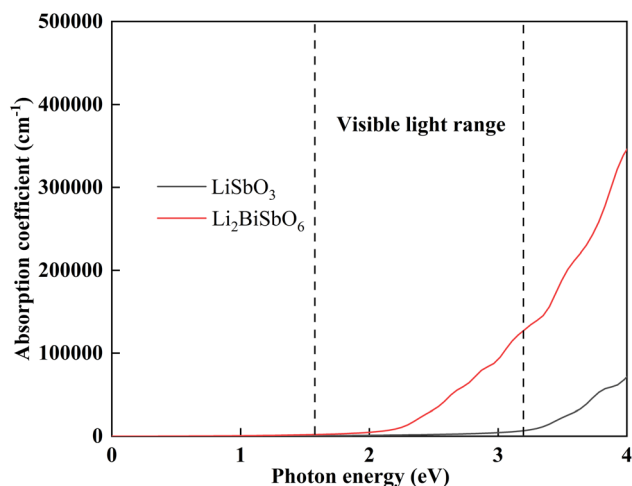


Fig. 7 The calculated light absorption coefficient.

is close to $0.5m_0$, and the electron carrier mobility could be larger due to the small effective mass. The ratio of m_h/m_e was calculated to further explore the response to photoelectric performance. A lower recombination rate and higher transfer rate of photogenerated carriers could be induced for a larger ratio of m_h/m_e . The m_h/m_e ratios of $\text{Li}_2\text{SbBiO}_6$ are 7.414 and 5.934 without and with the SOC effect, respectively, which are greater than those of LiSbO_3 polarized structures. The m_h/m_e ratio of the ferroelectric material BiFeO_3 is 4.589,³⁸ and the ferroelectric polarization strength is higher than that of $\text{Li}_2\text{SbBiO}_6$, but the bandgap of BiFeO_3 is relatively large and absorbs only a small amount of visible light for photocatalytic properties.^{39,40} The high degradation efficiency could arise from the strong visible light absorption, depolarization field of ferroelectrics, intrinsic small carrier effective mass, and large m_h/m_e ratio in $\text{Li}_2\text{SbBiO}_6$, which contribute to the photoinduced carrier separation in the photocatalytic process.

6. Conclusions

The stability and photoelectric properties of the $\text{Li}_2\text{SbBiO}_6$ ferroelectric compound have been investigated using density functional theory. The calculation results as well as the elastic coefficient and phonon dynamics analyses show that $\text{Li}_2\text{SbBiO}_6$ is a potentially stable ferroelectric material. The ferroelectric stability of $\text{Li}_2\text{SbBiO}_6$ and LiSbO_3 is related to the strong effect of the Li–O bond. $\text{Li}_2\text{SbBiO}_6$ slightly improves the ferroelectric polarization of LiSbO_3 and significantly reduces the bandgap to expand its applications in photovoltaic and photocatalysis with visible light. The larger m_h/m_e ratio of $\text{Li}_2\text{SbBiO}_6$ accompanied by a suitable bandgap and ferroelectric properties is more favorable for the separation of carriers and absorption of visible light.

Conflicts of interest

The authors declared that they have no conflicts of interest to this work. We declare that we do not have any commercial or associative interest that represents a conflict of interest in connection with the work submitted.

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Table 4 The calculated effective mass of holes (m_h) and electrons (m_e), the ratio of m_h/m_e , and ferroelectric polarization strength (P_s)

Compounds	$m_h (m_0)$	$m_e (m_0)$	m_h/m_e	$P_s (\mu\text{C cm}^{-2})$
LiSbO_3	1.484	0.433	3.424	34
LiBiO_3	1.651	0.504	3.272	47 (50 (ref. 20))
$\text{Li}_2\text{SbBiO}_6$	3.618(3.258-soc)	0.488(0.549-soc)	7.414(5.934-soc)	39
BiFeO_3	3.171 (ref. 38)	0.691 (ref. 38)	4.589	90



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