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Anharmonic stabilization and band gap renormalization in the perovskite CsSnI₃

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Amongst the X(Sn,Pb)Y₃ perovskites currently under scrutiny for their photovoltaic applications, the cubic B-α phase of CsSnI₃ is arguably the best characterized experimentally. Yet, according to the standard harmonic theory of phonons, this deceptively simple phase should not exist at all due to rotational instabilities of the SnI₆ octahedra. Here, employing self-consistent phonon theory, we show that these soft modes are stabilized at experimental conditions through anharmonic phonon-phonon interactions between the Cs ions and their iodine cages. We further calculate the renormalization of the electronic energies due to vibrations and find an unusual opening of the band gap, estimated as 0.24 and 0.11 eV at 500 and 300 K, which we attribute to the stretching of Sn-I bonds. Our work demonstrates the important role of temperature in accurately describing these materials.

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Four decades after its identification as an unusual phase-change material [1], the inorganic perovskite CsSnI₃ has experienced a revival of interest in its technological applications. After being used as a hole transporter in solid-state photovoltaics [2], the subsequent explosion in activity surrounding perovskite solar cells [3] has seen CsSnI₃ incorporated into new devices as a lead-free light absorber [4] with favorable optical properties [5–8]. Like many perovskites [9] CsSnI₃ has a rich phase diagram, driven by low-energy rotations and tilts of the SnI₆ octahedra [10–12]. In addition CsSnI₃ has an unusual electronic structure, with a nondegenerate and highly-dispersive valence band [13] and an intra-atomic band gap strongly coupled to external strain [13–15].

In a wider context, CsSnI₃ is the gateway to understanding the basic physics of the family of X(Sn,Pb)Y₃ perovskites (X = cation, Y = halogen). Unlike its famous cousin MAPbI₃ (MA = methylammonium), CsSnI₃ has (i) no permanent cationic dipole moment [16], (ii) reduced spin-orbit coupling due to the lighter mass of Sn [17], and (iii) a high-symmetry cubic (B-α) phase characterized by many studies [10,12,18]. However, theoretical investigations [11,19–21] consistently find the B-α phase to be unstable against spontaneous rotation of the SnI₆ octahedra, so on energetic grounds this phase should not exist at all. The answer to this puzzle must partly lie in the fact that the B-α phase is stable at high temperature [12], where both energetic and entropic contributions determine the free energy $F$. Unfortunately the most widely-used approach of calculating $F$ from first principles, the quasiharmonic approximation [22], cannot be straightforwardly applied [21] due to the presence of the unstable (imaginary) phonon modes (Fig. 1).

In this Rapid Communication, we demonstrate the critical role played by anharmonicity in stabilizing the experimentally-observed cubic and tetragonal (B-α and B-β) phases of CsSnI₃. We perform our ab initio investigation using a stochastic implementation of self-consistent phonon theory [24–27]. We show that the SnI₆ octahedra are stabilized against tilts and rotations by interacting with the renormalized vibrations of the Cs ions. Unexpectedly our calculations also reveal a temperature-induced opening of the band gap, with a magnitude of 0.24 and 0.11 eV at 500 and 300 K, respectively. The significant size of these corrections (36 and 11% of the uncorrected gaps) places temperature effects at a similar level of importance as spin-orbit coupling for determining the band gap in these materials [17], yet usually they are not included in ab initio studies. We further find that the gap opening is not consistent with a harmonic theory of band gap renormalization [28] but can be understood in terms of an increase in average length of the Sn-I bonds.

All total energy and force calculations in this paper were performed within a generalized-gradient approximation to density-functional theory (the PBEsol functional [29]), expanding the wave functions in a plane-wave basis set [30] and treating the interactions between electrons and ion cores within the projector-augmented wave formalism [31] as implemented in the GPAW code [32].

In Fig. 1 we show the phonon band structure obtained for the B-α phase calculated in the harmonic approximation using the finite displacement method [33,34]. Instabilities corresponding to tilts and rotations of the SnI₆ octahedra are found at the M and R points of the Brillouin zone [11]. One of the triply-degenerate soft modes at M is shown in Fig. 2, together with a frozen-phonon calculation of the potential energy surface (PES) with respect to a distortion $x_{rot}$ along this mode. The cubic structure ($x_{rot} = 0$) is metastable, and the system can lower its potential energy through an octahedral rotation to a new structure with tetragonal symmetry. These soft modes have been observed under a number of different computational setups and theoretical approximations [11,19–21] and cannot be stabilized for example by the application of a strain. In fact increasing the lattice constant above its 0 K equilibrium value yields further soft modes, identified as ferroelectric instabilities in Ref. [21].

The fact that experiments observe only the cubic phase at temperatures above 440 K [10,12,18] indicates that this phase corresponds to a minimum of the free energy $F$. In the quasiharmonic approximation [22], $F$ is replaced with $\tilde{F}(\omega,T)$, the free energy of an ensemble of oscillators of temperature $T$ with frequencies $\omega = \{\omega_1, \omega_2, ..., \omega_\nu\}$:

$$\tilde{F}(\omega,T) = V_0 + \sum_\nu \left[ \frac{\hbar \omega_\nu}{2} - k_B T \ln[1 + n_B(\omega_\nu,T)] \right].$$

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\( V_0 \) is the energy of the ions in their equilibrium positions, \( \hbar \) and \( k_B \) the Planck and Boltzmann constants, and \( n_B \) the Bose-Einstein distribution function.

A quasiharmonic treatment of the B-\( \alpha \) phase would replace \( \omega \) with the phonon eigenfrequencies shown in Fig. 1, but there are two difficulties: First, Eq. (1) is defined only for real phonon frequencies, so the contribution to \( F \) from the soft modes cannot be included. Second, from the harmonic phonon frequencies and eigenvectors of Fig. 1 we calculate that Cs atoms would undergo typical oscillations with a root mean-square displacement of 0.8 Å at 500 K, corresponding to 18% of the distance to their iodine neighbors at equilibrium [35]. Such large displacements are unlikely to be well described within the harmonic approximation.

Determining \( F \) for the B-\( \alpha \) phase therefore requires moving beyond the (quasi)harmonic approximation. Different approaches to this problem have been developed, including methods based on parametrizations and perturbative expansions of the PES [36–38], molecular dynamics [39,40], and self-consistent phonons [24–27,41]. Here we follow the self-consistent phonon approach and calculate a fictitious free energy \( F(\omega, T) \) as

\[
F(\omega, T) = \tilde{F}(\omega, T) + \langle V \rangle_T - \langle \tilde{V} \rangle_T. \tag{2}
\]

Here \( \tilde{V} \) is a harmonic approximation to the true PES \( V \), and \( \langle A \rangle_T \) is a thermal average of a quantity \( A \) with respect to the fictitious harmonic system, whose exact value is obtained via Mehler’s formula [42,43] as

\[
\langle A \rangle_T = \prod_{\nu} \frac{1}{\sqrt{2\pi \langle \chi^2_\nu \rangle_T}} \int dx_\nu e^{-x^2_\nu/(2\langle \chi^2_\nu \rangle_T)} A(x). \tag{3}
\]

\( x \) gives the amplitudes along each phonon mode \( \nu \), with the mean-square amplitude at temperature \( T \) given as \( \langle \chi^2_\nu \rangle_T \). \( F \) is the free energy of the real system evaluated on the thermal equilibrium state of the fictitious system and is a rigorous upper bound to the true free energy \( F \) [27]. The self-consistent set of frequencies \( \omega \) are chosen as those which minimize \( F \).

A fully self-consistent phonon theory (e.g., Refs. [26,27]) also minimizes \( F \) with respect to phonon eigenvectors and equilibrium ionic positions, but in the current study we keep these quantities fixed at their harmonic values. The reasons for performing this simplification are (i) for the high-symmetry B-\( \alpha \) phase, many of the phonon eigenvectors (including the soft modes) are fixed by the crystal symmetry, and (ii) the large unit cells and low symmetry of the B-\( \beta \) and B-\( \gamma \) phases render a full minimization of \( F \) impractical [44], even after performing the symmetrization techniques of Ref. [27]. Then, as has been done previously for calculating free energies [26,27], absorption spectra [45], and magnetic spectroscopies [46], we evaluate the thermal averages of Eq. (3) stochastically from an ensemble of configurations with ionic displacements distributed according to \( \prod_{\nu} \exp[-x^2_\nu/(2\langle \chi^2_\nu \rangle_T)] \). We label the current scheme SCCo.

Figure 1 shows the SCCo-calculated phonon band structure obtained at 500 K. There are three points to note. First, the soft modes at the \( M \) and \( R \) points are stabilized to positive energies of 2.3 meV. Second, the vibrational energies of the Cs atoms appearing at 1–3 meV in the harmonic approximation [21] are renormalized by more than a factor of two in SCCo. As a result, ferroelectric instabilities involving Cs atoms that appear at a strained lattice vanish at high temperatures [35]. Finally, the lattice constant which minimizes \( F \) is calculated to be 6.21 Å, which compared to experiment (6.206 Å [12]) is a significant improvement over the values of 6.131 Å found from a quasiharmonic analysis ignoring the soft modes [21].

The significant renormalization of the Cs vibrations points to the mechanism by which the soft modes are stabilized in SCCo. The SCCo potential calculated for octahedral rotations is far steeper than that expected from a one-dimensional analysis of a quartic potential, which yields a parabola wide enough for the system to sample the two minima [35]. Instead one must consider phonon-phonon interactions between the octahedral rotations and the vibrations of the Cs atoms. In Fig. 2(c) we show the PES obtained by simultaneously displacing the Cs atoms along an \( M \)-point phonon while rotating the SnI\( _6 \) octahedra. Harmonically for each Cs mode amplitude...
Given the interest in the optoelectronic properties of CsSnI$_3$, it is desirable to quantify the effects of phonons on the electronic band gap $E_g$. There is increasing evidence that semi-local exchange-correlation functionals find a weaker electron-phonon coupling strength compared to more sophisticated theories of electronic excitations, e.g., the GW approximation [47,48]. For this reason we perform electronic structure calculations using the derivative discontinuity-corrected GLLB-SC functional of Ref. [49], which has been found to improve the PBEsol description of the band gap for a range of materials [14,50]. We calculate a gap deformation potential of 7.20 eV with GLLB-SC, close to the value of 7.35 eV found from the quasiparticle self-consistent GW (QSGW) calculations of Ref. [13] and steeper than the values of 4.73 found with PBEsol or 4.65 eV from the local-density approximation [13]. The derivative discontinuity is responsible for this difference [35].

Expanding the lattice constant from 6.131 Å (harmonic, $T = 0$ K) to 6.21 Å (SCCo, $T = 500$ K) already accounts for an increase of the gap $E_g$ from 0.40 eV to 0.66 eV. However in addition there is a constant-volume renormalization of the gap due to phonons [51], which in the adiabatic approximation of Ref. [28] is calculated as $\Delta E_g = (E_g)_{\text{har}} - E_g^0$, where $E_g^0$ is the gap calculated with the ions in their equilibrium positions [43]. We use Eq. (3) to evaluate $(E_g)_{\text{har}}$ from the SCco frequencies at the experimental volume at 500 K. The band gaps calculated for 300 configurations is shown in Fig. 3. The calculated $\Delta E_g$ is remarkable for being both large and positive, i.e., the electron-phonon interaction increases the gap. Although the latter behavior has been observed experimentally for materials like copper halides [51,52], ab initio calculations of electron-phonon renormalization have so far focused on semiconductors like diamond and Si where the gap is reduced by temperature [47,53–55].

We have also studied the technologically-relevant low temperature B-β and B-γ phases at 380 and 300 K, respectively. Owing to the close agreement of the SCco B-α lattice constant with experiment, we used the experimental lattice constants reported in Ref. [12] for the other phases. We show the SCco band structures in the supplemental information [35]. For the B-β phase, the SCco calculations remove the unstable modes and renormalize the frequencies of the Cs modes, whilst for the B-γ phase some small changes in phonon frequencies occur across the spectrum [35]. The calculated corrections to the band gap are again large, yielding $+0.70$ eV (B-α phase, 500 K), $+0.45$ eV (B-β phase, 380 K), and $+0.31$ eV (B-γ phase, 300 K). However as discussed below these values are likely to be overestimates due to finite size effects in our supercell calculations.

To further investigate this band gap renormalization we first consider the harmonic theory of Ref. [28], where

$$\Delta E_g \approx \Delta E_g^{(2)} = \sum_{\nu} \frac{\partial E_g}{\partial n_\nu} \left[ n_B(\omega_\nu, T) + \frac{1}{2} \right].$$

(4)

Usually the coupling coefficient $\partial E_g/\partial n_\nu$ is defined as $l_\nu^2 \partial^2 E_g/\partial x_\nu^2$ with $l_\nu$ being the characteristic length of the normal mode [35], but with this definition Eq. (4) gives a too large gap renormalization of 0.95 eV at 500 K, demonstrating the failure of a harmonic expansion of $E_g$ with respect to $x_\nu$. However Fig. 3 reveals a correlation between the calculated gap and $D_{\text{SnI}}$, the average Sn-I bond length, which accounts for the gap increase both from the electron-phonon interaction and between the unperturbed $\alpha$, $\beta$, and $\gamma$ phases (orange stars). Following Ref. [13] we attribute this sensitivity to a weakened Sn-Sn-I-$\rho$ antibonding interaction as the bond length increases, narrowing the valence band and widening the band gap. This correlation motivates a resummation and new coupling constant definition:

$$\frac{\partial E_g}{\partial n_\nu} = l_\nu^2 \frac{d}{dD_{\text{SnI}}} \frac{\partial^2 D_{\text{SnI}}}{\partial x_\nu^2},$$

(5)

where $dE_g/dD_{\text{SnI}}$ is the gradient of the straight line in Fig. 3. In Fig. 4 we plot the spectral functions [56] $g^2 F(\omega) = \sum_\nu \partial E_g/\partial n_\nu \delta(\omega - \omega_\nu)$ for the two definitions of $\partial E_g/\partial n_\nu$, showing that (a) the harmonic expansion of $E_g$ predicts much larger contributions from polar modes at 6 meV, and (b) both expansions yield an important contribution to the gap renormalization from the octahedral rotations at 2.3 meV, which can only be described with an anharmonic treatment of the ground state.

Equation (5) yields a gap renormalization of 0.70 eV for the B-α phase at 500 K, exactly reproducing the ensemble average of Fig. 3. Noting that the B-β and B-γ phases display a similar correlation of $E_g$ with $D_{\text{SnI}}$ [35], we combined $dE_g/dD_{\text{SnI}}$ from Fig. 3 with $\partial^2 D_{\text{SnI}}/\partial x_\nu^2$ obtained from the phonon eigenvectors of these phases and found renormalizations of

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**FIG. 3.** (Color online) GLLB-SC band gaps calculated with/without (green squares/gray circles) derivative discontinuity contribution for an ensemble of 300 configurations vs average Sn-I bond length $D_{\text{SnI}}$. A linear fit to the data is shown. $E_g$ is taken as the difference between the highest occupied state and the average of the three lowest unoccupied states at $R$ [35]. We also show the gaps of the unperturbed B-α, B-β, and B-γ structures (orange stars). Note that these calculations were performed in a $2 \times 2 \times 2$ supercell and are subject to the finite size effects discussed in the text.
to a significant reduction of $dE_g/dS_{\text{Sat}}$ by 62%, thus giving revised estimates of the gap renormalization from Eqs. (4) and (5) of +0.24, +0.16, and +0.11 eV for CsSnI$_3$ at 500, 380, and 300 K, or corrected GLLB-SC gaps of 0.90, 1.04, and 1.17 eV. Future work is required to study the nature and origin of this slow size convergence.

Connecting our work to experimental studies, we note that Ref. [8] found the peak photoluminescence (PL) to increase in energy by 0.09 eV from 9 to 300 K. Although this data appears to agree with our calculated shift of +0.11 eV, we note that (a) the latter value does not include thermal expansion effects, and (b) it is unclear whether the PL corresponds to band-band transitions or defects [12,60]. At higher temperatures, our calculations indicate that the band gap will reduce, e.g., by 0.14 eV between 380 and 500 K. The measurement of the absorption spectrum of CsSnI$_3$ over the 0–500 K temperature range would be highly useful to further explore these effects.

Finally, we note that while anharmonicity has been demonstrated to play a crucial role for materials at very high temperatures or pressures [27,41], the conditions simulated here are relevant to the expected operating conditions for solar cells [61]. It is notable that the 0.24 eV shift obtained for the cubic phase is of similar magnitude to the spin-orbit correction [13], with opposite sign. Our paper thus illustrates the importance of anharmonic temperature effects to the realistic modeling of the $X$(Sn,Pb)Y$_3$ perovskites.

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We used plane-wave cutoffs of 1000 and 600 eV for the phonon and band gap calculations. We used supercells and (Γ-centered) k-point grids of $2 \times 2 \times 2 / 6 \times 6 \times 6$ (B-α), $1 \times 1 \times 2 / 6 \times 6 \times 4$ (B-β), and $1 \times 1 \times 1 / 6 \times 4 \times 6$ (B-γ).

As an example, a $1 \times 1 \times 1$ sampling for the B-γ phase would require minimizing $F$ with respect to 861 basis functions, compared to the 25 realized in Ref. [27].