



# Tetragonal high-pressure phase of InI predicted from first principles

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## ARTICLE INFO

### Article history:

Received 23 September 2011

Accepted 3 November 2011

Available online 9 November 2011

### Keywords:

InI

Swarm optimization algorithm

High pressure transition

First principles

## ABSTRACT

*Ab initio* particle swarm optimization algorithm for crystal structural prediction was employed to uncover the high-pressure crystal structure of indium iodide (InI). We have predicted one tetragonal high-pressure phase for InI with P4/nmm symmetry, which is energetically much superior to the previously proposed CsCl-type structure. The P4/nmm-InI possesses alternative stacking of double I and In layers. The arrangement of adjacent I and In layers of P4/nmm-InI is similar to that of the CsCl-type structure. The calculated electronic density of states supports a metallic character for this tetragonal phase that is similar to the high-pressure behavior of IIA–VIB families. Furthermore, the phase transition path from the ambient pressure TII–InI → P4/nmm-InI has been discussed.

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## 1. Introduction

Indium halides have attracted much attention due to their unique physical and chemical properties in fundamental science and technological applications, such as high volatility at reasonably low temperatures, fast recombination rates fairly low dissociation energies, etc. [1–4]. One of indium halides, InI, is a promising semiconductor for radiation detector. The high atomic number of In and I, along with high density, give the InI detector a higher photon stopping power (attenuation coefficient) compared to Ge and comparable to that of CdTe. The resistivity of InI was reported to be very high,  $\sim 10^{11} \Omega \text{ cm}$ , at room temperature with a mobility lifetime product for electron  $\sim 7 \times 10^{-5} \text{ cm}^2/\text{V}$ . Another major advantage of InI is that both In and I are less toxic compared to other compound semiconductors used for fabrication of solid state detector such as CdTe (Cd), HgI<sub>2</sub> (Hg), and GaAs (As) [5–7].

Materials under high pressure exhibit rich transition behavior, which gives us broadened views into the essence of atomic binding in solids while greatly challenging our instinctive understanding. At ambient pressure, InI crystallizes in the base-center orthorhombic structure of TII-type (space group Cmcm) [8], which differs from that of the well investigated cubic thallium halides. However, the high-pressure behaviors of InI are rarely studied, and there is lack of confirmed reports on the existence of pressure-induced phase transitions. Recently, Becker and Beck [9,10] have investigated the high-pressure phase transitions of

TII-type compounds including InI by density functional theory (DFT) calculations. They proposed a however CsCl-type phase as the candidate structure at high pressure on account of its low Madelung energy, which becomes dominant in determining structures at small volumes. They determined this new high-pressure structure by energy calculations of different known possible structures. This procedure may miss some unexpected yet more stable structures. Accordingly, this question is open until the convincing proof is provided. The peculiarity and the absence of characterized high-pressure phase of InI prompted our endeavor to investigate its structural behavior at high pressure.

Here, we extensively explored the crystal structures of InI over a wide range of pressures (0–60 GPa) using a specifically developed particle swarm optimization (PSO) algorithm for crystal structure prediction [11,12]. This methodology has been successful in correctly predicting crystal structures for various systems including elements and compounds at high pressure [13–16], unbiased by any known information. We found one layer-like tetragonal P4/nmm structure, which is energetically more preferable than the previously proposed CsCl-type structure. The phase transition path from the TII-type → P4/nmm structure has also been discussed.

## 2. Computational details

Our PSO methodology on crystal structural prediction has been implemented in CALYPSO (Crystal structure Analysis by Particle Swarm Optimization) code [12] at 0 GPa, 25 GPa, and 60 GPa with 1–4 formula units (f.u.) per simulation cell. The underlying

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*ab initio* structural relaxations and electronic calculations were carried out using DFT within the Perdew–Burke–Ernzerhof exchange–correlation as implemented in the VASP code [17–19]. The all-electron projector augmented wave method [20] was adopted with  $5s^25p^5$  and  $4d^{10}5s^25p^1$  treated as valence electrons for I and In, respectively. The energy cutoff 400 eV and appropriate Monkhorst–Pack  $k$  meshes [21] were chosen to ensure that enthalpy calculations are well converged to better than 1 meV/atom. The phonon calculations were carried out using a supercell approach as implemented in the PHONOPY code [22].

### 3. Results and discussion

At 0 GPa, our simulations with the only input of chemical composition of In:I=1:1 predicted the most stable structure to be orthorhombic TII-type phase, in complete agreement with experiments. The TII–InI contains four f.u. per unit cell, and the I ions are fivefold coordinated and the In ions are sevenfold coordinated. Fig. 1a presents the crystal structure of the orthorhombic modification of TII projected on the  $bc$ -plane. Along the  $c$ -direction one can recognize zigzag chains of In–I sticks resembling the structure of trans-polyacetylene with In and I ions at the sites of C and H, respectively. Moreover, in order to provide some insight into the pressure behavior of TII–InI according to Ref. [9], the pressure dependences of the TII–InI lattice constants are plotted in Fig. 2, along with the experimental data and other theoretical results [9]. Strikingly, the calculated results are in agreement with these experimental data and theoretical values, indicating the reliability of our calculations.

The success in the prediction of TII-type structure gives us confidence to further explore the high-pressure phases of InI. For higher pressure at 20 GPa and 60 GPa, a new stable tetragonal P4/nmm phase having 2 f.u./cell (Fig. 1b) was discovered for InI. At 20 GPa, the optimized lattice parameters of P4/nmm–InI are  $a=3.497$  Å and  $c=7.771$  Å, with I and In atoms occupying  $2c$  (0, 0, 0.1534) and  $2c$  (0, 0, 0.6177) positions, respectively. This P4/nmm structure possesses alternative stacking of double I and In layers, and the coordination number of I and In ions increased up to 13. Each I ion is surrounded by five In ions and eight I ions (Fig. 1d), and each In ion is surrounded by five I ions and eight In ions (Fig. 1e), resulting in a similar polyhedron structure. We have performed the calculations on the phonon dispersion curves of P4/nmm–InI and CsCl–InI at 60 and 30 GPa, respectively. As shown in Fig. 3a, no imaginary phonon frequency was detected in the whole Brillouin zone for P4/nmm–InI, indicating its dynamical stability. For CsCl–InI shown in Fig. 3b, however, it is clearly seen that the phonon spectrum shows a large imaginary frequency at M point, signaling its structural instability.

To determine the phase transition pressure, we have plotted out the enthalpy curves of the P4/nmm–InI relative to the ambient-pressure TII–InI (Cmcm–InI) in Fig. 4a, and the CsCl–InI was also considered for comparison. It is confirmed that the TII–InI transforms to the predicted P4/nmm structure above 17 GPa, and thus the CsCl–InI can be ruled out as existing. Meanwhile, the calculated volumes as a function of pressure for TII–InI and P4/nmm–InI are also shown in Fig. 4b. The results suggest that TII–InI  $\rightarrow$  P4/nmm–InI phase transition is first-order with clear volume drops of 5.9%, which can be easily detected in further experiment. Furthermore, the thermodynamic stability of P4/nmm–InI at high pressure with respect to decomposition is quantified in terms of the formation enthalpy,  $\Delta H_f = H_{\text{InI}} - H_{\text{I}} - H_{\text{In}}$ . The  $\Delta H_f$  is the formation enthalpy, the orthorhombic I (space group: Cmca) and tetragonal In (space group: I4/mmm) are chosen as the reference phases. The obtained results have demonstrated the structural stability of P4/nmm–InI against the decomposition into the mixture I and In up to 50 GPa.

Since Cmcm is a subgroup of P4/nmm, it is essential to investigate the relations between these two structures, which can help us to understand the mechanism of the Cmcm–InI  $\rightarrow$  P4/nmm–InI transition. In fact, as shown in Fig. 5(a) and (c), both Cmcm and P4/nmm are layer-like structures with the stacking order  $KLKL\ldots$  and  $NN'MM'NN'\ldots$ . The transition path can be understood qualitatively as follows: for Cmcm structure, the I and In atoms in  $K$  layers are frozen, while the I and In atoms in  $L$  layers moved as a whole along  $b$  direction in  $ab$  planes during

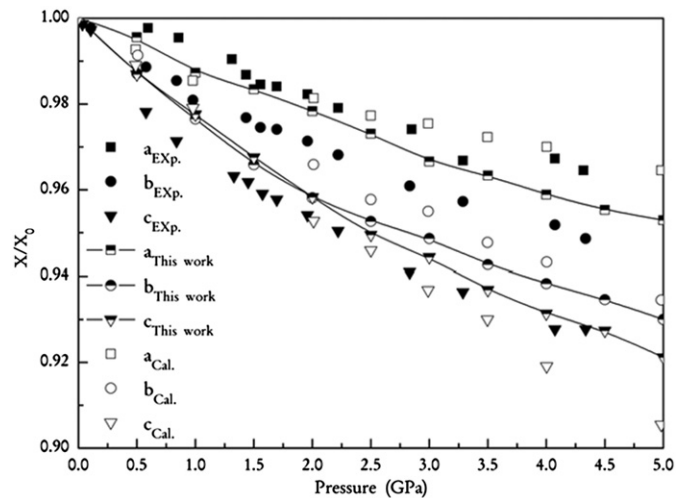


Fig. 2. Pressure dependences of lattice constants for TII–InI, together with the experimental data and other theoretical values.

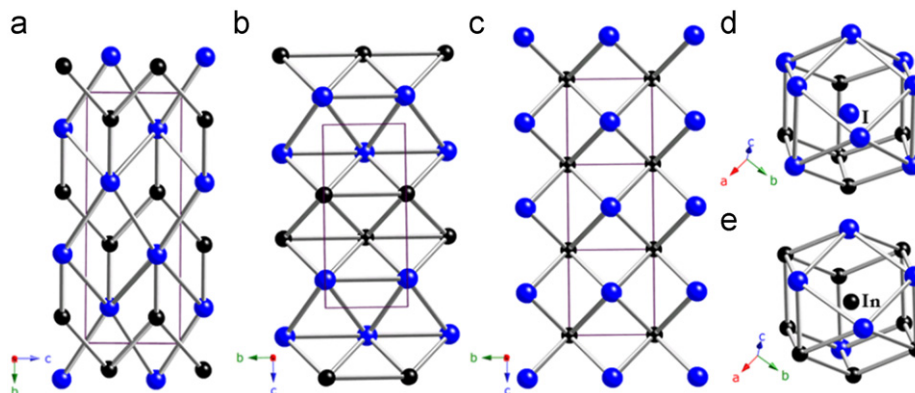


Fig. 1. Crystal structures of InI for TII-type (a), P4/nmm (b), and CsCl-type (c). The large and small spheres represent I and In atoms, respectively.

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