Direct observation of titanium-centered octahedra in titanium–antimony–tellurium phase-change material

Feng Rao, Zhitang Song, Yan Cheng, Xiaosong Liu, Mengjiao Xia, Wei Li, Keyuan Ding, Xuefei Feng, Min Zhu & Songlin Feng

Phase-change memory based on Ti0.4Sb2Te3 material has one order of magnitude faster Set speed and as low as one-fifth of the Reset energy compared with the conventional Ge2Sb2Te5 based device. However, the phase-transition mechanism of the Ti0.4Sb2Te3 material remains inconclusive due to the lack of direct experimental evidence. Here we report a direct atom-by-atom chemical identification of titanium-centered octahedra in crystalline Ti0.4Sb2Te3 material with a state-of-the-art atomic mapping technology. Further, by using soft X-ray absorption spectroscopy and density function theory simulations, we identify in amorphous Ti0.4Sb2Te3 the titanium atoms preferably maintain the octahedral configuration. Our work may pave the way to more thorough understanding and tailoring of the nature of the Ti–Sb–Te material, for promoting the development of dynamic random access memory-like phase-change memory as an emerging storage-class memory to reform current memory hierarchy.
The performance gap between processor and memory has been widened in recent years since massively parallel computing with multicore processors being utilized. An emerging storage-class memory (SCM) technology was introduced to mitigate the bottleneck, which concerns the entire computing system. To reform the conventional memory hierarchy, the SCM technology needs to combine the attributes of memory and storage; that is to say, the performances of SCM are hoped to match commodity dynamic random access memory (DRAM), while the capacity and cost of SCM shall be able to compete with NAND Flash memory. Phase-change memory (PCM) has been regarded as the most promising candidate to replace incumbent NAND Flash memory as well as DRAM for SCM application. In terms of the speed, power, and endurance, the fact is PCM already has overall better bit performances than those of the NAND. Obviously, multi-level resistance drift limitation in single cell will need a lot more work.

Nevertheless, the key challenges to realize DRAM-like PCM still lie in how to further decrease Reset operation energy and further increase Set operation speed, because DRAM requires rigorously $10^{-11} \sim 10^{-12}$ J per bit program energy and <10 ns access time. The Reset operation of PCM, generally referring to the melt-quenching procedure, transforms the crystalline (c-) chalcogenide alloy, for example, Ge$_2$Sb$_2$Te$_5$ (GST), into amorphous (a-) phase. Conversely, the Set operation is to heat the a-phase to a temperature between crystallization temperature ($T_c$) and melting point ($T_m$) to obtain the c-phase. Due to the slow crystallization speed ($\sim 30$ ns), and high $T_m$ ($\sim 616^\circ C$) of GST-based materials, many efforts, such as diminish the grain size, promote the nucleation, reduce the active volume and enhance the thermal efficiency, have been made to try to lower the power and expedite the response.

Our previous work proposed an alternative and simpler approach to develop DRAM-like PCM via merely replacing the switching material from GST to Ti$_0.4$Sb$_2$Te$_3$ (TST) alloy. We demonstrated at least one order of magnitude faster Set speed and as low as one-fifth of the Reset energy on TST-based PCM cell compared with those of GST cell with the same size. Nevertheless, the origin of such performance improvements yet is controversial and not fully understood because of the lack of solid experimental evidences. In the present work, we apply Cs-corrected scanning transmission electron microscopy (STEM) and soft X-ray absorption spectroscopy (sXAS) combined with density functional theory (DFT) simulations to explore the phase transition mechanism of TST material in a more accurate way.

We report that the Ti-centered octahedra (TCOs) not only disperse inside of the hexagonal (HEX) Sb$_2$Te$_3$ (ST) lattice, but also form nano-lamellae separated outside, respectively acting as lattice distortion centers and nano-thermal insulators to induce low-energy amorphization of the restricted ST crystal grains. We also identify that in a-TST, the Ti atoms prefer to maintain the octahedral configuration. Such robust TCOs or nano-lamellae thus may serve as intrinsic nucleation centers or templates for surrounding Sb-centered atomic motifs to swiftly align into long-range order to achieve high-speed recrystallization. An understanding of the phase-transition mechanism of TST from the atomic and electronic structure points of view is essential for further material optimization to meet the requirements of future DRAM-like PCM application.

### Results

**Identification of triple-layered TiTe$_2$ lamellae.** From the STEM bright-field (BF) image of c-TST (Fig. 1a), several atomically resolved triple-layered lamellae can be clearly observed. They locate adjacent to the quintuple layers (QLs). Figure 1b shows the high-angle annular dark-field (HAADF) STEM image of the triple-layered lamellae zooming into the field marked by dashed line in Fig. 1a. The energy dispersive spectrometer (EDS) mappings of the Ti, Sb and Te elements for the corresponding zooming area in Fig. 1b are given by Fig. 1c–e, respectively. The triple-layered lamellae are determined to be composed with a stacking sequence of $-v$-Te-Ti-v-Te-Ti-v-Te-Ti-v-Te-Ti-v-Te-Ti-v-Te-Ti-v-Te- (Fig. 1b and Supplementary Fig. 1), where v denotes the van der Waals interaction gap. Such stacking sequence is the typical HEX lattice structure of TiTe$_2$ (TT) with lattice parameter $a = 3.77$ Å (ref. 19). While the adjacent QLs are believed to be $-v$-Te-Sb-Te-Sb-Te-v- stacking structure of HEX- ST with a little larger lattice parameter $a = 4.25$ Å (ref. 20), which can be confirmed in Fig. 2. Although TT and ST were commonly used to construct super-lattice-like materials, the $\sim 11\%$ mismatch of $a$ between TT and ST blocks still tends to result in the phase segregation. Differently magnified STEM-HAADF images and the corresponding EDS mappings of other observing regions in the same c-TST sample also confirm this segregation phenomenon that the ST crystal

![Figure 1 | Structural and chemical identifications of TiTe2 lamellae in crystalline Ti0.4Sb2Te3.](image-url)
grains are shrunk or even cut into pieces by the TT lamellae (TTL) (Supplementary Figs 2 and 3). In contrast, the \( a \) values of GeTe (GT) and ST constructing the flagship GST alloy only deviate slightly (\( \sim 2.2\% \) for HEX lattice and \( \sim 1.0\% \) for rocksalt lattice)\(^{20} \). Thus, unlike the incompatibility between TT and ST, it would be rare to find severe GT segregation out of ST in c-GST phase. Nevertheless, the smaller ST crystal grains defined by TT in c-TST may be helpful to boost the speed and power performances of PCM\(^{14} \).

Identification of Ti-centered octahedra in Sb\(_2\)Te\(_3\) quintuple layers. The HAADF image (Fig. 2a) shows the clear ST-like HEX lattice structure of the QLs in c-TST, which is atom-by-atom chemically identified as -v-Te\(_{(2)}\)-Sb-Te\(_{(1)}\)-Sb-Te\(_{(2)}\)-v- stacking sequence (Fig. 2b), where subscripts (1) and (2), respectively, refer to 3-coordinated and 6-coordinated Te atoms. To be noted that there are non-ignorable Ti signals simultaneously detected in some Sb layers (Fig. 2c), which indicates that dopant Ti atoms can occupy Sb lattice sites in the HEX-ST lattice\(^{17} \). It is also worthy of pointing out that not all the QLs of ST in c-TST accommodate the Ti dopants (Supplementary Fig. 4).

Discussion

Given above, the representative lattice structure of HEX-TST should be depicted as the model presented in Fig. 3 from which, in addition to the pure ST QLs, one may also find the separated TTL as well as the TCOs scattered in ST QLs. The charge density difference\(^{22} \) maps indicate that in the Sb-centered octahedron, there are three strong and three weak bonds\(^{23} \); while among Ti and its six Te ligands in the TCO, there are six equally stronger covalent-like bonds formed by overlapping Ti-3d and Te-5p orbitals, which are quite similar with the electronic configuration of pure HEX-TT\(^{24,25} \). Due to the stronger Ti-Te\(_{(2)}\) bonds, the Te\(_{(2)}\)-Sb bonds in the adjacent layer without the Ti are significantly elongated and weakened, as indicated in Fig. 3 by dashed lines. The rigid TCOs possibly bring tensile stress to the ST QLs, and those fragile Te\(_{(2)}\)-Sb bonds have to be deformed to accommodate such lattice distortion. Thus, with more Ti atoms occupying in the same Sb layer, the interlayer distance (\( d_{1} \approx 1.9 \text{ Å} \)) between Sb(Ti) and Te\(_{(2)}\) layers is found to be a little smaller than that (\( d_{2} \approx 2.1 \text{ Å} \)) between the Te\(_{(2)}\) and Sb layers (Fig. 3). This subtle interplane expansion is also pointed out by black arrows in Fig. 2a,d. In comparison, there should be no such

---

**Figure 2** | Structural and chemical identifications of quintuple layers in crystalline Ti\(_{0.4}\)Sb\(_{2}\)Te\(_{3}\). (a) Scanning transmission electron microscopy (STEM) high-angle annular dark-field (HAADF) image of quintuple layers, projected along \(<100>\) direction. (b) Energy dispersive spectrometer (EDS) mappings of Sb and Te. (c) EDS mappings of Ti and Te. (d) STEM-HAADF image zooming into the field marked in a. Compared with the interlayer distance (\( d_{1} \)) between Sb(Ti) and Te\(_{(2)}\) layers, the wider gap (\( d_{2} \)) between Te\(_{(2)}\) and Sb layers is pointed out with black arrows. The Te···Te Van der Waals interaction gaps are marked by double-line. Scale bar, (a,d) \( 1 \) nm.

**Figure 3** | Lattice structure and bonding chemistry of crystalline Ti\(_{0.4}\)Sb\(_{2}\)Te\(_{3}\). Hexagonal lattice of Ti\(_{0.4}\)Sb\(_{2}\)Te\(_{3}\) with charge-density difference. The differences are calculated with respect to that of a superposition of isolated atoms. The isosurface (transparent yellow area) shows electron pileup mostly at the bonds. The isosurface value is fixed at \( \pm 0.005 \ e \ a_{0}^{-3} \) (\( a_{0} = \) bohr). The Sb-centered octahedron has three strong bonds and three weak bonds, while the Ti-centered octahedron has six strong bonds and no weak bond. In quintuple layers (QLs) with the Ti dopants, the formation of strong Ti-Te\(_{(2)}\) bonds weakens some of the adjacent Te\(_{(2)}\)-Sb bonds as marked by dashed lines, leading to a larger interlayer distance (\( d_{2} \)) between Te\(_{(2)}\) and Sb layers compared with that (\( d_{1} \)) between Sb(Ti) and Te\(_{(2)}\) layers. While in pure Sb\(_{2}\)Te\(_{3}\) QLs, the interlayer distances (\( d_{3} \)) among Sb-Te\(_{(2)}\)-Sb layers are almost the same.
inter-distance variation for Sb–Te(2)-Sb layers ($d_3 \approx 2.0$ Å) in pure ST QLs (Fig. 3). During amorphization the fragile Te(2)-Sb bonds will be easily ruptured to destroy the long-range order, leading to a low-energy Reset operation. By contrast, the TTL with all rigid Ti-Te bonds may tend to keep its ordered configuration in amorphization, and in reverse effectively restrict the growth of ST grain. Since the quasi-two-dimensional c-TT is semimetallic23, and also has quite lower thermal conductivity (0.12 W mK$^{-1}$)26 than those of c-GST (0.41 W mK$^{-1}$)27 and c-ST (0.78 W mK$^{-1}$)28, electric current shall be more conveniently conducted to the small ST grains to generate Joule heat that is insulated by adjacent TTL to accomplish a highly efficient Reset operation.

Comparing with our previous differential scanning calorimetry (DSC) result of TST material27, the remeasured DSC curve (Supplementary Fig. 5) expands the measuring temperature from $\sim 600$ to $\sim 650$ °C. Here we only focus on the melting procedures. The onsets of two endothermic peaks at $\sim 557$ and $\sim 603$ °C are taken as the first and the second $T_m$s of TST material, respectively. Note that most of the heat is consumed by the second melting process which has a lower $T_m$ than that of pure ST ($\sim 618$ °C)12,29. Chemically, we can rewrite the Ti$_{0.4}$Sb$_2$Te$_3$ stoichiometry into (TiTe$_2$)$_x$Ti$_{0.4}$Sb$_2$Te$_3-2x$, ($0 < x < 0.4$), where (TiTe$_2$)$_x$ part accounts for the TTL located outside of the ST QLs, then the rest Ti$_{0.4}$-Sb$_2$Te$_3-2x$, from Ti$_{0.4}$-Sb$_2$Te$_3-2x$ part should be considered as the TCOs inside some ST QLs. Note that the Ti$_{0.4}$-Sb$_2$Te$_3-2x$ part ($0 < x < 0.4$) resembles the Sb-rich Sb-Te compound (2/3 < Sb at%/Te at% < 2/2)22 whose widely accepted microstructural fingerprint is the Sb$_2$ bilayers stacking adjacent to some ST QLs. Indeed, we also observed such structure of minority from individual crystal grains in c-TST (marked by black arrows in Supplementary Fig. 6). No doubt, this Sb-rich composition can naturally help to result in a faster crystallization speed than that of a-GST30. However, without accurately quantifying the value of $x$, we cannot determine the repeated stacking manner of Sb$_2$ and ST blocks for Ti$_{0.4}$-Sb$_2$Te$_3-2x$ part, thus we prefer not to include the Sb$_2$ bilayer in the representative lattice model of HEX-TST shown in Fig. 3. Furthermore, because pure TT material has quite larger $T_m$ (> 1,200 °C) than those of the Sb–Te compounds30,32, it is rational to deduce the TTL and TCOs could be unmelted during c-TST amorphization. Actually, our in situ X-ray diffraction results of pure TT film on Si substrate verify that even at 700 °C (> $T_m$ of TST) (Supplementary Fig. 7), TT film still maintains good crystallinity through a typical HEX lattice structure22. Therefore, we believe the Ti$_{0.4}$-Sb$_2$Te$_3-2x$ part in TTL play the major role in causing the two consequent melting processes resembling the case of pure Sb-rich Sb–Te compounds29, especially the robust TCOs highly distort the HEX-ST lattice, promoting the significant decrease of the second $T_m$.

On the other hand, in recrystallization, we speculate that the long-range ordered TTL and the structurally similar TCOs may serve as the heterogeneous nucleation templates or centers for neighbouring distorted or destructive Sb-centered octahedra to easily and swiftly align into long-range order13,14,33,34. This ordering mechanism is qualitatively different with the one of a-GST that involves a reconfiguration of considerable tetrahedrally bonded Ge atoms to octahedrally bonded ones35,36 or a cavity-supported reorientation of four-membered ring structures37,38. In this regard, we also note that in Si doped ST material, Si and ST exhibit nano-scale phase segregation morphology resembling that of TST; namely, dopant Si atoms keep in the a-phase outside of the ST-rich areas and do not participate in the phase transition39. Nevertheless, the a-Si phase cannot play the role of inducing swift recrystallization of the a-ST phase because of the complete lattice mismatch between a-Si and c-ST, leading to a sluggish Set operation39.

One may still inquire whether the Ti atoms could keep their octahedral configurations even during high-temperature amorphization process. We thereby used sXAS method to verify the existence of TCOs in a-TST. The spot size of the incident X-ray beam for taking sXAS is about 1.5 mm$^2$. In contrast to the STEM for microstructural probe, the sXAS technique can provide the overall information in macroscopic scale. For 3$d$ transition-metal (TM) compounds, sXAS involves the excitation of 2$p$ core electrons to empty 3$d$ orbitals (2$p^3\rightarrow3d^0$ to $2p^3\rightarrow3d^+$) through dipole selection rules40,41. Thus, sXAS directly probes the TM 3$d$ unoccupied electronic structures that is sensitive to its oxidation states, spin states, and local environment such as chemical bonding, symmetry and metal–ligand distance. Figure 4a shows the Ti L-edge absorption spectra of TST film with 1 K min$^{-1}$ in situ heating rate. Note that the as deposited TST film is amorphous, and will begin to be crystallized when the in situ heating temperature reaches the $T_c$ ~ 451 K (Supplementary Fig. 8). The absorption spectra are divided into $L_2$ and $L_3$ regions, corresponding to 2$p_{3/2}$ and 2$p_{1/2}$, levels resulting from the 2$p$ core-hole spin-orbital coupling, and further split to double-peak structure (labelled as A and B in Fig. 4a) by the crystal field generated by the Te atoms surrounding each Ti atom. To understand the effect of this crystal field on the electronic structure of TST, we perform the DFT calculation to analyse the electron density of states. Here we use HEX-TT (space group of P3m1 and point group of $D_{3h}$)32 as a reference to analyse the sXAS spectrum taken at 513 K corresponding to the c-TST phase. In contrast to the well-studied analogue TiO$_2$ with point group of $D_{4h}$ (refs 42,43), where the slightly distorted octahedron of O atoms leads the splitting of degeneracy Ti-3$d$ orbitals into $t_2$ and $e_g$ orbitals with an energy separation 10 $D_q$, the fivefold degenerate orbitals of HEX-TT are split into three parts: the upper two degenerate orbitals of $d_{xy}$, $d_{yz}$, and $d_{zx}$, the middle orbital of $d_{z^2}$, and the rest two degenerate orbitals of $d_{x^2-y^2}$ and $d_{xy}$, as shown in Fig. 4b. The simulated Ti-$L_3$ sXAS from electron-hole approximation is displayed in Fig. 4c, which shows the double peaks with energy separation about 1.6 eV. This energy scale quantitatively agrees with that of the experimental sXAS spectra for c-TST (Fig. 4a). Although the ratio of double-peak intensities is not well reproduced that may due to the lack of considering the electron-hole interaction44, this distinction is trivial for our study that focuses on the effect of crystal field. Because the temperature-dependent sXAS spectra, in particular the crystal-field separation, display almost the identical features, the Ti electronic configurations should be nearly unchanged in both a- and c-TST phases. This is the evidence, from the electronic structure point of view, proving that the Ti atoms maintain in the octahedral configuration in a-phase. Furthermore, our ab initio molecular dynamics simulation on melting TST alloy (data not shown here) also indicates that the ST QLs can be adequately disordered while the TTL basically preserve the ordered triple-layered structures. In this regard, we speculate that the separated TTL are able to survive after amorphization if the Reset operation of the TST PCM cell can be carefully manipulated. It has great potentials to design a suitable heating profile (< $T_m$ of pure TT) to preserve the ordered TTL inside a disordered a-ST network. Higher-quality STEM and EDS evidences of ordered TTL formed in Reset state of the TST PCM cell still need more work.

In conclusion, STEM and sXAS studies of the TST material have revealed its essential microstructure feature that Ti atoms do exist in the octahedral configuration in both a- and c-phases. We have identified the dispersed TCOs in the HEX-ST QLs as well as the externally separated TTL. The former ones highly distort the ST lattice, while the latter ones effectively aggregate the heat in
the small ST grains, both promoting the low-energy amorphization of c-TST. The octahedral configuration centered by Ti atom is proved to be prevailed in a-TST, which should originate from the strong covalent-like characteristic of Ti-Te bonds. The precise roles of the TTT and TCOs in triggering the swift recrystallization are not yet clear. We conjecture the long-range ordered TTT and the structurally similar TCOs may probably act as the medium for inducing the reorder of surrounding Sb-centered atomic motifs. The quantification of Ti contents in TTT and TCOs requires more study, and will be a great help to determining the lattice structure, understanding the phase transition mechanism and further optimizing the material composition for DRAM-like high-performance PCM application. In addition, the finding of intrinsic quasi-two-dimensional c-TT lamella in the polycrystalline TST alloy may open up the possibility of searching the topological insulating or superconducting properties from a lattice configuration well-controlled Ti–Sb–Te alloy45,46.

**Methods**

**Film characterization.** The composition of TST film was measured by X-ray fluorescence spectroscopy using a Rigaku RIX 2100 system. TST film was deposited on silicon substrate at room temperature, and annealed at 250 °C for 2 min to get the c-phase37. Cross-sectional view TEM sample of TST film was prepared by using Ar ion milling method. The atomic identification was carried out by using JEOL ARM 300F and FEI Titan 300 with probe-corrected STEM mode. The element mapping was obtained through using a supercell calculation, where we removed a core electron on one atom in the supercell of 216 Ti, 65 Sb and 132 Te atoms. To reproduce the lattice structure of c-TST used the all-electron full potential linear augmented plane wave plus local orbitals method35 as implemented in the Wien2k code36. The exchange correlation potential was calculated also using the PBE functional. The absorption spectroscopy was obtained through using a supercell calculation, where we removed a core electron on one atom in the supercell of 2 × 2 × 2, and added such electron into the conduction band.

**Ab initio theoretical simulation.** The lattice model of c-TST was investigated by employing the DFT47. The Vienna Ab initio Simulations Package48 was used. The projector augmented wave pseudopotentials49 were used for electron-ion interactions. For the exchange-correlation energies between electrons, the Perdew-Burke-Ernzerhof (PBE) functional50 was used. A 216-atom supercell was built with 19 Ti, 65 Sb and 132 Te atoms. To reproduce the lattice structure of c-TST, a triple-layered TT was inserted between a Te···Te Van der Waals gap in HEX-ST and some Sb atoms were randomly replaced by dopant Ti atoms. The energy cutoff was set to 235 eV. The Γ point was chosen for relaxation and a 3 × 3 × 3 k-point mesh was used for electronic properties studies. In addition, xAS simulation of c-TT used the all-electron full potential plane wave plus local orbitals method35 as implemented in the Wien2k code36. The exchange correlation potential was calculated also using the PBE functional. The absorption spectroscopy was obtained through using a supercell calculation, where we removed a core electron on one atom in the supercell of 2 × 2 × 2, and added such electron into the conduction band.

**References**

1. Lam, C. H. in Storage Class Memory. 10th IEEE International Conference on Solid-State and Integrated Circuit Technology (ICSICT). 1080–1083 (Shanghai, China, 2010).
2. Frietas, R. F. & Wilcke, W. W. Storage-class memory: the next storage system technology. IBM J. Res. Dev. 25, 439 (2008).
3. Cheng, H. Y. et al. in IEEE International Electron Devices Meeting (IEDM) 30.6.1–30.6.4 (Washington DC, USA, 2013).
4. Kim, I. S. et al. in Symposium on VLSI Technology (VLSIT) 203–204 (Honoolum, HI, USA, 2010).
5. Wong, H. S. P. et al. Phase change memory. Proc. IEEE 98, 2201–2227 (2010).
6. Kau, D. et al. in IEEE International Electron Devices Meeting (IEDM)1–4 (Baltimore, MD, USA, 2009).
7. Kinoshita, M. et al. in Symposium on VLSI Technology (VLSIT). 35–36 (Honolulu, HI, USA, 2012).
8. Hwang, Y. N. et al. in Symposium on VLSI Technology (VLSIT). 201–202 (Honolulu, HI, USA, 2010).
9. Oh, G. H. et al. in Symposium on VLSI Technology (VLSIT). 220–221 (Kyoto, Japan, 2009).
10. Weitem, M. Phase-change materials: Towards a universal memory? Nature 4, 265–266 (2005).
11. Atwood, G. Phase-change materials for electronic memories. Science 321, 210–211 (2008).
12. Yamada, N. et al. Rapid-phase transitions of GeTe-Sb2Te3, pseudobinary amorphous thin films for an optical disk memory. Nat. Phys. 65, 2849–2856 (1991).
13. Loke, D. et al. Breaking the speed limits of phase-change memory. Science 336, 1566–1569 (2012).
14. Wang, W. J. et al. in IEEE International Electron Devices Meeting (IEDM) 31.3.1–31.3.4 (San Francisco, CA, USA, 2012).
15. Xiong, F. et al. Low-power switching of phase-change materials with carbon nanotube electrodes. Science 332, 568–570 (2011).
16. Lee, S. et al. Highly scalable non-volatile and ultra-low power phase-change nanowire memory. Nat. Nanotechnol. 10, 626–630 (2007).
17. Zhu, M. et al. One order of magnitude faster phase change at reduced power in Ti-Sb-Te. Nat. Commun. 5, 4086 (2014).
18. Zhu, M. et al. Understanding the crystallographic behavior of as-deposited Ti-Sb alloy through real-time radial distribution function. Nanoscale 7, 9935–9944 (2015).
19. Greenaway, D. L. & Nitsche, R. Preparation and optical properties of group IV-VI chalcogenides having the CdI₂ structure. J. Phys. Chem. Solids 26, 1445–1458 (1965).
20. Da Silva, J. L. F. et al. Insights into the structure of the stable and metastable (GeTe)₅(Sb₂Te₃) compounds. Phys. Rev. B 78, 224111 (2008).
21. Schmid, B. et al. in ICT 2005. 24th International Conference on Thermoelectrics. 270–272 (Clemson, SC, USA, 2005).
22. Kolobov, A. et al. Vacancy-mediated three-center four-electron bonds in GeTe-Sb₂Te₃ phase-change memory alloys. Phys. Rev. B 87, 165206 (2013).
23. Kostylev, V. et al. Why phase-change media are fast and stable: a new approach to an old problem. Jpn J. Appl. Phys. 44, 3345–3349 (2005).
24. Kim, Y. et al. Electronic structures and chemical bonding of TiX₂ (X = S, Se, and Te). Jpn J. Appl. Phys. 37, 4878–4883 (1998).
25. Calessen, R. et al. Complete band-structure determination of the quasi-two-dimensional Fermi-liquid reference compound TiTe₂. Phys. Rev. B 54, 2453–2465 (1996).
26. Chiritecus, C. et al. Lower limit to the lattice thermal conductivity of nanostructured Bi₂Te₃-based materials. J. Appl. Phys. 106, 075303 (2009).
27. Ryu, S. et al. SiO₂ doped GeₓSb₁₋ₓTe films with high thermal efficiency for applications in phase change random access memory. Nanotechnology 22, 255005 (2011).
28. Chen, J. et al. Sb₂Te₃ nanoparticles with enhanced seebeck coefficient and low thermal conductivity. Chem. Mater. 22, 3086–3092 (2010).
29. Ghosh, G. The Sb-Te (antimony-tellurium) system. J. Phase Equilib. 15, 349–360 (1994).
30. Youm, M. et al. Effects of excess Sb on crystallization of β-phase Sb₂Te₅ binary thin films. Phys. Stat. Sol. A 205, 1636–1640 (2008).
31. Kifune, K. et al. Crystal structures of X-phase in the Sb–Te binary alloy system. Cryst. Res. Technol. 48, 1011–1021 (2013).
32. Cordes, H. & Schmid-Fetzer, R. Phase equilibria in the Ti-Sb-Te system. J. Alloy. Compd. 216, 197–206 (1994).
33. Caravati, S. et al. First-principles study of liquid and amorphous Sb₂Te₃. Phys. Rev. B 81, 205118 (2010).
34. Matsuanga, T. et al. From local structure to nanosecond recrystallization dynamics in Ag₈SnSbTe phase-change materials. Nat. Mater. 10, 129–134 (2011).
35. Kolobov, A. et al. Understanding the phase-change mechanism of rewritable optical media. Nat. Mater. 3, 703–708 (2004).
36. Krbal, M. et al. Intrinsic complexity of the melt-quenched amorphous GeₓSb₁₋ₓTe₃ memory alloy. Phys. Rev. B 83, 054203 (2011).
37. Akola, J. & Jones, R. O. Structural phase transitions on the nanoscale: the crucial pattern in the phase-change materials GeₓSb₁₋ₓTe₃ and GeTe. Phys. Rev. B 76, 235201 (2007).
38. Kalikka, J. et al. Nucleus-driven crystallization of amorphous GeₓSb₁₋ₓTe₃: A density functional study. Phys. Rev. B 86, 144113 (2012).
39. Rao, F. et al. Si–Sb–Te materials for phase change memory applications. Nanotechnology 22, 145702 (2011).
40. Groot, F. & Kotani, A. Core Level Spectroscopy of Solids (Taylor & Francis Ltd, 2008).
41. Stohr, J. NEXAFS Spectroscopy (Springer, 1992).
42. Crocombette, J. P. & Jollet, F. Ti 2p X-ray absorption in titanium dioxides (TiO₂): the influence of the cation site environment. J. Phys. Condens. Matter 6, 10811–10821 (1994).
43. Okada, K. & Kotani, A. Theory of core level X-ray photoemission and photoabsorption in Ti compounds. J. Electron. Spectrosc. Relat. Phenom. 62, 131–140 (1992).
44. Gallano, A. et al. A highly exciting: a full-potential all-electron package implementing density-functional theory and many-body perturbation theory. J. Phys. Condens. Matter 26, 363202 (2014).
45. Chhowalla, M. et al. The chemistry of two-dimensional layered transition metal dichalcogenide nanosheets. Nat. Chem. 5, 263–275 (2013).
46. Wray, L. et al. Observation of topological order in a superconducting doped topological insulator. Nat. Phys. 6, 855–859 (2010).
47. Hohenberg, P. & Kohn, W. Inhomogeneous electron gas. Phys. Rev. 136, B864–B871 (1964).
48. Kresse, G. & Hafner, J. Ab initio molecular dynamics for liquid metals. Phys. Rev. B 47, 558–561 (1993).
49. Bögli, P. E. Projector augmented-wave method. Phys. Rev. B 50, 17953–17979 (1994).
50. Perdew, J. P., Burke, K. & Ernzerhof, M. Generalized Gradient Approximation Made Simple. Phys. Rev. Lett. 77, 3865–3868 (1996).
51. Singh, D. J. & Nordstrom, L. Planewaves, Pseudopotentials, and the LAPW Method (Springer-Verlag, 2006).
52. Blaha, P. et al. WIEN2K: An Augmented PlaneWave + Local Orbitals Program for Calculating Crystal Properties (Technical University Wien, 2001).

Acknowledgements
This work was supported by Strategic Priority Research Program of Chinese Academy of Sciences (XDA09000402), National Key Basic Research Program of China (2011CB910201, 2011CB910202), National Natural Science Foundation of China (61076121, 61161622, 61216106001, 61376006, 21473235, 11272902), Science and Technology Council of Shanghai (13ZJ1442700, 13DJ2295700). X.L. also thank One Hundred Person Project of the Chinese Academy of Sciences, and Shanghai Pujiang Program (14PJ1404400). The supercomputer time was provided by National Supercomputer Center in Tianjin, and the calculations were performed on TianHe-1 (A). Many thanks for the strong support from JIOEL and FEI companies in STEM experiments, and Dr Jin-Ming Chen, Dr Jenn-Min Lee from National Synchrotron Radiation Research Center (NSRRC, Taiwan) and Prof. Cheng-hao Chuang from Tamkang University in xAS experiments. We also thank Dr W. Zhang from Xi’an Jiaotong University, Dr X.B. Li from Jilin University, and Prof. S.B. Zhang from Rensselaer Polytechnic Institute for useful discussions.

Additional information
Supplementary Information accompanies this paper at http://www.nature.com/naturecommunications
Competing financial interests: The authors declare no competing financial interests.
Reprints and permission information is available online at http://npg.nature.com/reprintsandpermissions/
How to cite this article: Rao, F. Direct observation of titanium-centered octahedra in titanium–antimony–tellurium phase-change material. Nat. Commun. 6, 10040; doi: 10.1038/ncomms10040 (2015).

This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/