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First-principle investigation of pressure and temperature influence on structural, mechanical and thermodynamic properties of Ti_3AC_2 (A = Al and Si)

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ABSTRACT

The mechanical and thermodynamic properties of Ti_3AlC_2 and Ti_3SiC_2 have been investigated by the first principles. The structure is briefly discussed, the compressibility of Ti_3SiC_2 is better than Ti_3AlC_2. Its mechanical stability is confirmed by the elastic constant, and the results show that the pressure can enhance the resistance to the deformation. In addition, the ductility is also improved with the pressure increasing. The elastic anisotropy is enhanced with increasing the pressure. Finally, the temperature and pressure dependences of thermodynamic properties of Ti_3AlC_2 and Ti_3SiC_2 are evaluated by the quasi-harmonic Debye approximation theory, and analyzed the effect of pressure and temperature on the Debye temperature, bulk modulus, heat capacity and the thermal expansion coefficient. The Debye temperature and bulk modulus decrease with increasing the temperature but increase with increasing the pressure. The temperature and pressure have opposite influences on the heat capacity, moreover, the ability of Ti_3AlC_2 to absorb or release heat is stronger than that of Ti_3SiC_2.

1. Introduction

The $M_{n+1}AX_n$ phases, where M is an early transition metal, A is an A-group element, and X is either C or N [1–5], have attracted more and more attention due to their uniquely combined properties of both ceramics and metals. Similar to ceramics, they are elastically rigid, lightweight and maintain their strengths to high temperature. Like metals, they are thermally plastic at high temperature and most readily machinable, which makes them potentially interesting for technological applications [6–8], including high-temperature structural applications [8], electrode brush materials, chemical anticorrosive materials and high-temperature heating materials [9–11].

Over the last few years, a concerted effort has been made to investigate the properties of the MAX phase. For example, Whittle et al. [9] studied the radiation tolerance of Ti_3AlC_2 and Ti_3SiC_2. Drouelle et al. [10] concluded the deformation mechanisms of Ti_3AlC_2 by the high temperature tensile creep experiments. The electronic and structural properties of the layered ternary carbide Ti_3AlC_2 were evaluated by Zhou et al. [11], they analyzed the band structure and Fermi level and revealed an electronic conductor of the Ti_3AlC_2. Wang et al. [12] examined the properties of vacancies in Ti_3AlC_2 and Ti_3SiC_2 and got the effective results of, they found that an A-group element mono-vacancy was more easily formed in Ti_3AlC_2, the vacancies tended to disperse in Ti_3SiC_2 but aggregate in Ti_3AlC_2. P. Finkel et al. [13] measured the low temperature dependence of the elastic properties including Young's shear modulus and they investigated the Debye temperature. Low temperature heat capacity of Ti_3SiC_2 was calculated by Ho et al. [14,15]. However there are little research on the effects of the temperature and pressure on the Ti_3AlC_2 and Ti_3SiC_2.

In this paper, we concentrated on exploring the effects of Ti_3AlC_2 and Ti_3SiC_2 under the pressure ranging from 0 to 50 GPa based on the first principle and calculating the bulk modulus, Debye temperature, heat capacity and the thermal expansion coefficient at temperature ranging from 0 to 1000 K to investigate the thermodynamic properties by the quasi-harmonic Debye approximation theory.

2. Computational methods

The first principle calculations have been performed based on the density functional theory (DFT) [16] and the Cambridge Serial Total

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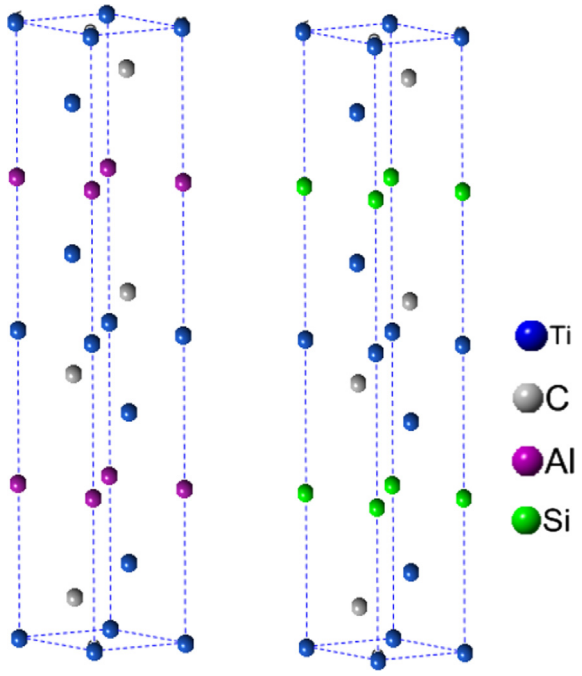

Fig. 1. The crystal structure of Ti_3AlC_2 and Ti_3SiC_2.

Energy Package (CASTEP) [17], in which a plane wave ultra-soft pseudopotential [18] is used. The generalized gradient approximation (GGA) with Perdew-Burke-Ernzerhof (PBE) approach [19] is performed to represent the exchange-correlation function. By convergence, the plane-wave basis set cutoff was set as 440 eV and the *k*-point mesh to 13 × 13 × 2 for all cases, the energy was converged to better than 5.0×10^{-6} eV/atom, whereas the maximum force and stress had been determined as 0.01 eV/Å and 0.02 GPa, respectively. The maximum displacement was set as 5.0×10^{-4} Å.

The Ti_3AlC_2 and Ti_3SiC_2 are hexagonal structures, as shown in Fig. 1. Which can be regarded as alternating stacks of two layers, the space group is $P6_3/mmc$. The Al or Si atoms are located in the (0, 0, 1/4) positions, the Ti atoms are located in at (0, 0, 0) and (2/3, 1/3, z_{Ti}) and the C atoms are located at (1/3, 2/3, z_c) [20,21].

3. Results and discussions

3.1. Structural properties

The relative changes of equilibrium volume and lattice parameters at various pressures ranging from 0 to 50 GPa with a step of 10 GPa are shown in Fig. 2, where V_0 and a_0 are the zero pressure equilibrium structural parameters. It is easy to see that the ratio of V/V_0 of Ti_3AC_2 (X = Al and Si) was reduced by 18.7% and 16.9%, respectively. Therefore, the compressibility of the system is strong. The volume change ratio with pressure gradually decreases as the order of Ti_3AlC_2 > Ti_3SiC_2. As for the Ti_3AC_2, the volume was reduced with increasing the internal pressure. At high pressure, the V/V_0 curves become gentle, since the distance change in the atoms gets smaller and thus the mutual repulsion of the atoms gets stronger, which leads to the difficulty of compression in crystal.

3.2. Mechanical properties

The elastic constants (C_{ij}), bulk modulus (B) and shear modulus (G) were calculated and listed in Table 1. The available theoretical date under different pressures was included for comparison. Because the Ti_3AC_2 is a hexagonal crystal, which has six different independent elastic constants (C_{11}, C_{12}, C_{13}, C_{33}, C_{44} and C_{66}), but only five of them

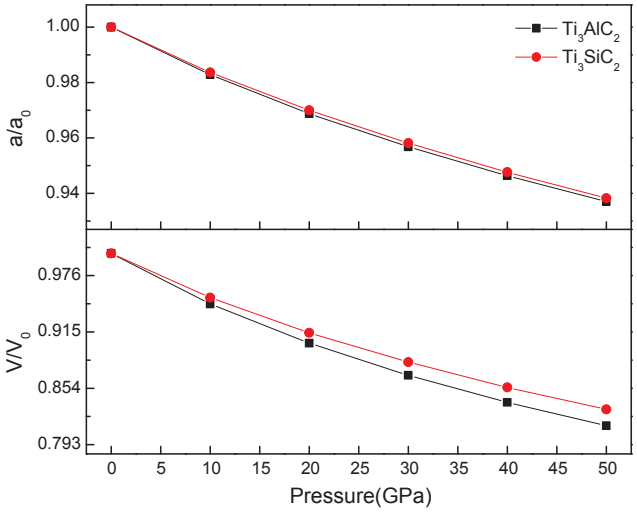

Fig. 2. The normalized lattice parameters a/a_0 and volume V/V_0 of Ti_3AlC_2 and Ti_3SiC_2 at various pressures.

are independent since $C_{66} = (C_{11} - C_{12})/2$ [22], thus only five independent elastic constants were listed. The mechanical stability of system can be judged by the Born stability criteria [23]:

$$C_{11} > 0,\ C_{11} - C_{12} > 0,\ C_{44} > 0,\ C_{66} > 0,\ (C_{11} + C_{12})C_{33} - 2C_{13}^2 > 0 \quad (1)$$

It is obviously to note that the elastic constants in the pressure range 0–50 GPa satisfy Eq. (1), which indicates that they are mechanically stable under compression and the value of C_{ij} goes up linearly with increasing the pressure, besides the C_{11} and C_{33} increase more rapidly than others.

As for the bulk modulus and shear modulus, these quantities are given by Eqs. (2)(7) [24–26]:

$$B_V = \frac{2(C_{11} + C_{12}) + C_{33} + 4C_{13}}{9} \quad (2)$$

$$B_R = \frac{(C_{11} + C_{12})C_{33} - 2C_{13}^2}{C_{11} + C_{12} + 2C_{33} - 4C_{13}} \quad (3)$$

$$B_H = \frac{B_V + B_R}{2} \quad (4)$$

$$G_V = \frac{C_{11} + C_{12} + 2C_{33} - 4C_{13} + 12C_{55} + 12C_{66}}{30} \quad (5)$$

$$G_R = \frac{5}{2} \frac{[(C_{11} + C_{12})C_{33} - 2C_{13}^2]C_{55}C_{66}}{3B_V C_{55}C_{66} + [(C_{11} + C_{12})C_{33} - 2C_{13}^2](C_{55} + C_{66})} \quad (6)$$

$$G_H = \frac{G_V + G_R}{2} \quad (7)$$

where bulk modulus B_B, B_R and B_H represent the maximum, minimum and average value, respectively, and the shear modulus G_B, G_R and G_H also represent the maximum, minimum and average value.

The values of B and G play an important role on the mechanical properties of materials. The value of the B and G increases linearly at various pressures ranging from 0 to 50 GPa with a step of 10 GPa, the bulk modulus is usually assumed to be a measure of resistance to volume change by applied pressure, which implies that pressure can improve the resistance to volume deformation. As for shear modulus and Young's modulus, the trend is similarly to the bulk modulus. The Young's modulus is used to provide a measure of the capability of resisting the tension and pressure in the range of elastic deformation [27]. Generally speaking, the larger the Young's modulus is, the harder it is to deform the material. The value of B, G and E of Ti_3SiC_2 is found to be higher than that of Ti_3AlC_2, the effect of resisting the deformation for the Ti_3SiC_2 is better than Ti_3AlC_2. Therefore, the change of external