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## First-principles study of a new structure and oxidation mechanism of Pt<sub>3</sub>Zr

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Zirconia (ZrO<sub>2</sub>)–metal interfaces are interesting for solid oxide fuel cells. Although the oxidation of Pt<sub>3</sub>Zr provides a new route for the formation of ZrO<sub>2</sub>–Pt interfaces, the crystal structure of Pt<sub>3</sub>Zr remains controversial and the oxidation mechanism of Pt<sub>3</sub>Zr is unclear. To solve these problems, we use first-principles calculations to explore the crystal structure of Pt<sub>3</sub>Zr. We demonstrate a stable structure of Pt<sub>3</sub>Zr based on phonon dispersion. Importantly, two new Pt<sub>3</sub>Zr structures, Ti<sub>3</sub>Pt-type (*Pm**m*) and Fe<sub>3</sub>Al-type (*Fm**m*), are predicted. To study the oxidation mechanism, two possible doped models are considered. The calculated results show that the O atom prefers to occupy the tetrahedral interstitial site (Tl) in comparison to the octahedral interstitial site (Ol). We find that the oxidizing capacity of the Fe<sub>3</sub>Al-type cubic structure is stronger than that of other structures. In particular, we predict that Pt<sub>3</sub>Zr exhibits better oxidation capacity in comparison to other metals because of the strong localized hybridization between Zr and O.

## 1. Introduction

Zirconia (ZrO<sub>2</sub>)–metal interfaces have widely used in solid oxide fuel cells (SOFCs), heterogeneous catalysis, gas sensors, *etc.*<sup>1–9</sup> Although ZrO<sub>2</sub> shows excellent chemical and thermal stabilities, the poor electrical conductivity hinders the application because of the requirement of sensitive probing techniques.<sup>10,11</sup> To solve this problem, an effective method is to oxidize Zr on an appropriate metal substrate. However, Zr deposited on a metal substrate should meet three conditions: oxygen reduction reaction (ORR), growth of the oxide layer and better catalytic activity of the metal substrate. In particular, the ORR clearly confirms the degree of oxidation between Zr and the metal substrate. Therefore, high ORR catalytic activity strongly depends on the d-state of the transition metal.<sup>12–14</sup>

Platinum-group-metals are important catalysts because of their excellent chemical and physical properties.<sup>15–18</sup> In particular, Pt is a fascinating catalyst because it effectively promotes the process of ORR.<sup>19–21</sup> Recently, the oxidation behavior of Pt<sub>3</sub>Zr (0001) surface was studied by the experiment and first-principles calculations.<sup>22</sup> They found that the weak localized hybridization between Pt and Zr is beneficial to oxidation of Zr metal. Note that the metal addition will accelerate the growth of ZrO<sub>2</sub> film on the Pt<sub>3</sub>Zr (0001) surface.<sup>23</sup> Therefore, structural configuration plays a crucial role in ORR and the formation of ZrO<sub>2</sub>–Pt interface. Unfortunately, the crystal structure of Pt<sub>3</sub>Zr remains controversial. Earlier work suggested that Pt<sub>3</sub>Zr is a Au<sub>3</sub>Cu-type cubic structure.<sup>24,25</sup> On the contrary, Stalick *et al.*

found that Pt<sub>3</sub>Zr belongs to a Ni<sub>3</sub>Ti-type hexagonal structure,<sup>26</sup> which is different from the previous viewpoint. As a result, the nature of oxidation mechanism of Pt<sub>3</sub>Zr is unclear.

To explore the catalytic activity of Pt<sub>3</sub>Zr and improve the formation of ZrO<sub>2</sub>–Pt interface, in our works, we investigate the crystal structure and oxidation mechanism of Pt<sub>3</sub>Zr by using the first-principles calculations. According to the structural feature, we predict two possible cubic structures. Importantly, the structural stability of Pt<sub>3</sub>Zr is estimated by the formation enthalpy and phonon dispersion. To examine the oxidation mechanism and oxidation capacity of Pt<sub>3</sub>Zr, we calculate and compare the oxygen doped formation energy between O-doped Pt<sub>3</sub>Zr and many metals. In particular, we examine the possible adsorption site of Pt<sub>3</sub>Zr. Finally, we predict that Pt<sub>3</sub>Zr shows better oxidation capacity in comparison to many metals.

## 2. Model and methods

To explore the oxidation mechanism, we firstly study the crystal structure of Pt<sub>3</sub>Zr. To our knowledge, one is that Pt<sub>3</sub>Zr is a Ni<sub>3</sub>Ti-type hexagonal structure with space group of *P6*<sub>3</sub>/*mmc*. The lattice parameters of hexagonal structure are *a* = 5.653 Å and *c* = 9.347 Å, respectively.<sup>24</sup> Another is that Pt<sub>3</sub>Zr belongs to the Au<sub>3</sub>Cu-type cubic structure with space group of *Pm**m*, and the lattice parameter is *a* = 4.051 Å.<sup>26</sup> In comparison to hexagonal structure, we suggest that Pt<sub>3</sub>Zr with cubic structure can promote the oxidation of Pt<sub>3</sub>Zr because of the localized hybridization between oxygen and Pt<sub>3</sub>Zr. Therefore, we further design two possible structures: Ti<sub>3</sub>Pt-type with cubic structure and Fe<sub>3</sub>Al-type with cubic structure, respectively. The structural models of Pt<sub>3</sub>Zr are shown in Fig. 1.

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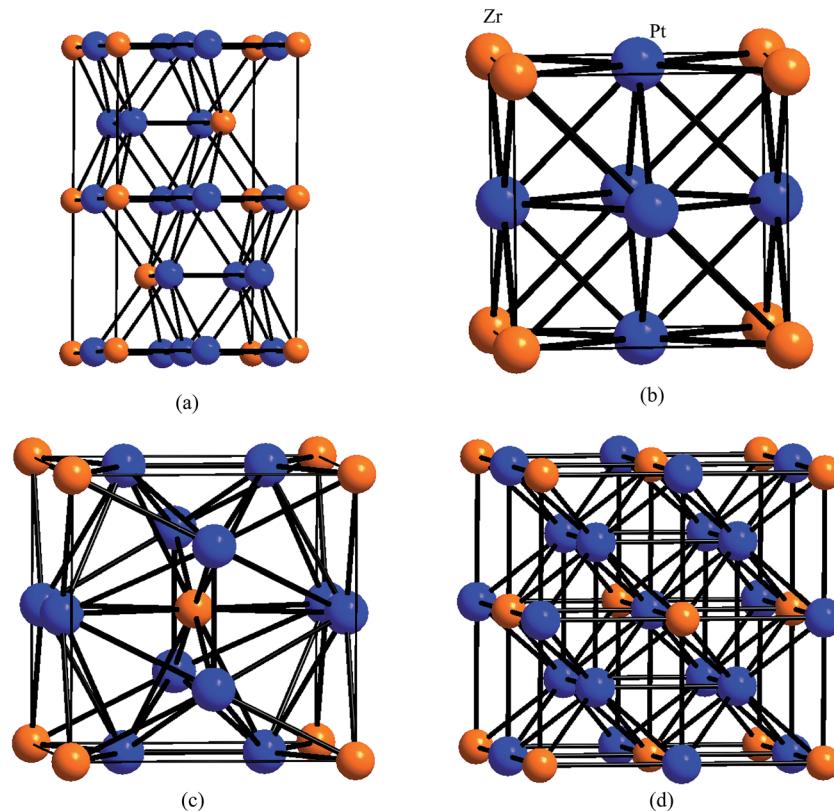


Fig. 1 Structural model of  $\text{Pt}_3\text{Zr}$ , (a)  $\text{Ni}_3\text{Ti}$ -type with hexagonal structure, (b)  $\text{Au}_3\text{Cu}$ -type with cubic structure, (c)  $\text{Ti}_3\text{Pt}$ -type with cubic structure and (d)  $\text{Fe}_3\text{Al}$ -type with cubic structure, respectively.

In this paper, the total energy, structural information, electronic structure and chemical bonding of  $\text{Pt}_3\text{Zr}$  and O-doped  $\text{Pt}_3\text{Zr}$  were calculated by using the first-principles calculations, as implemented in the CASTEP code.<sup>27</sup> The exchange-correlation-functional was calculated by using the generalized gradient approximation (GGA) within PW91 functional.<sup>28,29</sup> To treat the interaction between electrons and the ions, the atomic configurations of O, Pt and Zr were  $2s^22p^4$ ,  $5p^65d^96s^1$  and  $4p^64d^25s^2$ , respectively. Based on the convergence test, the plane-wave basis set for electron wave function with cutoff energy of 400 eV was used. The  $k$ -point grids of  $10 \times 10 \times 5$  for  $\text{Ni}_3\text{Ti}$ -type structure,  $12 \times 12 \times 12$  for  $\text{Au}_3\text{Cu}$ -type structure,  $10 \times 10 \times 10$  for  $\text{Ti}_3\text{Pt}$ -type structure,  $11 \times 11 \times 11$  for  $\text{Fe}_3\text{Al}$ -type structure,  $14 \times 14 \times 8$  for Zr and  $17 \times 17 \times 17$  for Pt were treated, respectively. To examine the dynamically stable, the phonon calculation was carried out by using the supercell method within the PHONON code.<sup>30</sup>

### 3. Results and discussions

#### 3.1 Structural prediction

The structural stability of  $\text{Pt}_3\text{Zr}$  is measured by the formation enthalpy and phonon dispersion, respectively. The equation of formation enthalpy ( $\Delta H$ ) is given by:

$$\Delta H = E_{\text{Pt}_3\text{Zr}} - 3E_{\text{Pt}} - E_{\text{Zr}} \quad (1)$$

where  $E_{\text{Pt}_3\text{Zr}}$ ,  $E_{\text{Pt}}$  and  $E_{\text{Zr}}$  are the total energy of  $\text{Pt}_3\text{Zr}$ , isolated Pt atom and isolated Zr atom, respectively.

Table 1 lists the calculated lattice parameters, density, and formation enthalpy of  $\text{Pt}_3\text{Zr}$  with four structures. It can be seen that these structures are thermodynamically stable at the ground state because the calculated formation enthalpy of these structures is smaller than zero. Our predicted two structures ( $\text{Ti}_3\text{Pt}$ -type and  $\text{Fe}_3\text{Al}$ -type) are also thermodynamically stable. Importantly, the calculated formation enthalpy of  $\text{Ni}_3\text{Ti}$ -type structure is  $-8.188$  eV per atom, which is smaller than that of  $\text{Au}_3\text{Cu}$ -type structure. The slight difference implies that external condition is easy to result in phase transition from  $\text{Ni}_3\text{Ti}$ -type structure to  $\text{Au}_3\text{Cu}$ -type structure. This result is similar to the Fairbank's viewpoint.<sup>24</sup>

Table 1 Calculated equilibrium lattice parameters,  $a$ -axis and  $c$ -axis (Å), density,  $\rho$  (g cm<sup>-3</sup>) and formation enthalpy,  $\Delta H$  (eV per atom) of  $\text{Pt}_3\text{Zr}$

Structure	Method	$a$	$c$	$\rho$	$\Delta H$
$\text{Ni}_3\text{Ti}$ -type	GGA	5.742	9.398	16.75	-8.188
	Exp <sup>24</sup>	5.653	9.347		
$\text{Au}_3\text{Cu}$ -type	GGA	4.061		17.77	-8.178
	Theo <sup>25</sup>	3.980			
	Exp <sup>26</sup>	4.051			
$\text{Ti}_3\text{Pt}$ -type	GGA	5.198		16.00	-7.918
$\text{Fe}_3\text{Al}$ -type	GGA	6.513		16.26	-7.871



In addition to thermodynamically stable, the dynamically stable of  $\text{Pt}_3\text{Zr}$  should examine follow. To study the dynamically stable, Fig. 2 displays the calculated phonon dispersion curves of  $\text{Pt}_3\text{Zr}$  with four structures. It is clear that  $\text{Pt}_3\text{Zr}$  with  $\text{Ni}_3\text{Ti}$ -type structure is a dynamically unstable because the imaginary phonon frequency is observed in this structure. However, we find that there is no imaginary phonon frequency in  $\text{Au}_3\text{Cu}$ -type structure, indicating that this structure is a dynamically stable at the ground state. Based on the first-principles calculations, we demonstrate the Stalick's viewpoint. Importantly, we predict two new  $\text{Pt}_3\text{Zr}$  structures ( $\text{Ti}_3\text{Pt}$ -type and  $\text{Fe}_3\text{Al}$ -type) because no imaginary phonon frequencies are found in these structures.

To further insight into the nature of dynamically stable, Fig. 3 shows the calculated phonon density of state (PhDOS) of  $\text{Pt}_3\text{Zr}$  with four structures. It can be seen that negative frequencies are found in  $\text{Ni}_3\text{Ti}$ -type structure, indicating that this structure is a mechanically unstable at the ground state. The calculated PhDOS profile reveals that the mechanically unstable of  $\text{Ni}_3\text{Ti}$ -type is attributed to the vibration of Pt atom at low frequency region. However,  $\text{Au}_3\text{Cu}$ -type,  $\text{Ti}_3\text{Pt}$ -type and  $\text{Fe}_3\text{Al}$ -type structures are mechanically stable because no negative frequencies are observed in these structures. In particular, the whole low frequency model of  $\text{Au}_3\text{Cu}$ -type structure derives from the vibration of Pt atom and Zr atom. With increasing frequency, Zr's vibration plays an important role in thermodynamic properties. Therefore, it is concluded that the structural stability of  $\text{Pt}_3\text{Zr}$  is markedly influenced by Pt-Zr bond.

To further reveal the structural stability, the structural information of  $\text{Pt}_3\text{Zr}$  is discussed. As listed in Table 1, the calculated lattice parameters of  $\text{Ni}_3\text{Ti}$ -type structure (space group:  $P6_3/mmc$ , no: 194) are  $a = 5.742 \text{ \AA}$  and  $c = 9.398 \text{ \AA}$ , which are in good agreement with the experimental data.<sup>24</sup> In this structure (see

Fig. 1(a)), the alternative Pt layer and Pt-Zr layer can be viewed along the  $c$ -axis. In particular, each Zr atom is surrounded by 4 Pt atoms at Pt-Zr layer. Therefore, Pt-Zr bond ( $2.601 \text{ \AA}$ ) can improve the structural stability of  $\text{Pt}_3\text{Zr}$  with  $\text{Ni}_3\text{Ti}$ -type structure. However, the cohesive force between layers is determined by the bond strength of Pt-Pt bond and part of Pt-Zr bond. In other words, Pt-Pt and Pt-Zr bonds play a key role in structural stability. The calculated bond length of Pt-Pt and Pt-Zr bonds is  $2.924 \text{ \AA}$  and  $2.968 \text{ \AA}$ , respectively.

For  $\text{Au}_3\text{Cu}$ -type structure (see Fig. 1(b)), the calculated lattice parameter is  $a = 4.061 \text{ \AA}$ , which is in excellent agreement with the experimental data and theoretical results.<sup>25,26</sup> Note that the symmetrical Pt-Zr bonds effectively improve its structural stability. The calculated bond length of Pt-Zr bond is  $2.871 \text{ \AA}$ , which is similar to the previous theoretical result.<sup>31</sup> Note that the bond length of Pt-Zr bond of  $\text{Au}_3\text{Cu}$ -type structure is slightly smaller than the corresponding bond for  $\text{Ni}_3\text{Ti}$ -type structure, indicating that the structural feature of the former can obviously improve the structural stability in comparison to the latter. Thus, we should consider the cubic structure to oxidize Zr.

The calculated lattice parameter of  $\text{Ti}_3\text{Pt}$ -type and  $\text{Fe}_3\text{Al}$ -type structures is  $a = 5.198 \text{ \AA}$  and  $a = 6.513 \text{ \AA}$ , respectively. From Fig. 1,  $\text{Ti}_3\text{Pt}$ -type structure is similar to  $\text{Nb}_3\text{Si}$ -type structure. In comparison to  $\text{Au}_3\text{Cu}$ -type structure, Pt atom in  $\text{Ti}_3\text{Pt}$ -type structure occurs migration from  $(0.50, 0, 0.50)$  site to  $(0.25, 0, 0.50)$  site. As a result, the variation of atomic position changes the localized hybridization between Pt and Zr, which forms two different Pt-Zr bonds. The calculated bond length of Pt-Zr bond is  $2.599 \text{ \AA}$  and  $2.906 \text{ \AA}$ , respectively. In particular, the atomic configuration can provide a large number of interstitial sites to adsorb oxygen. In addition, we find that the calculated lattice parameter of  $\text{Ti}_3\text{Pt}$ -type and  $\text{Fe}_3\text{Al}$ -type structures is larger than that of  $\text{Au}_3\text{Cu}$ -type structure.

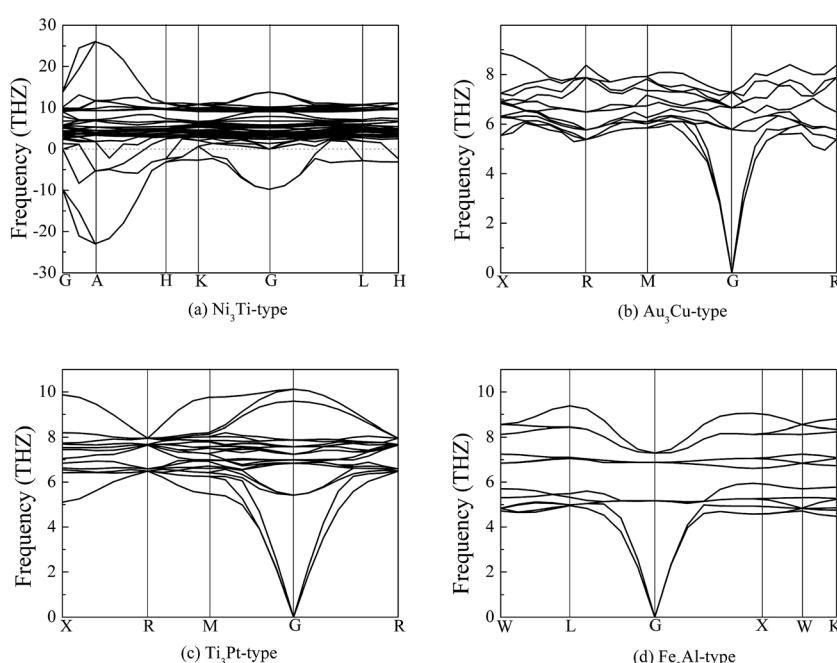


Fig. 2 Calculated phonon dispersion curves of  $\text{Pt}_3\text{Zr}$ , (a)  $\text{Ni}_3\text{Ti}$ -type structure, (b)  $\text{Au}_3\text{Cu}$ -type structure, (c)  $\text{Ti}_3\text{Pt}$ -type structure and (d)  $\text{Fe}_3\text{Al}$ -type structure, respectively.

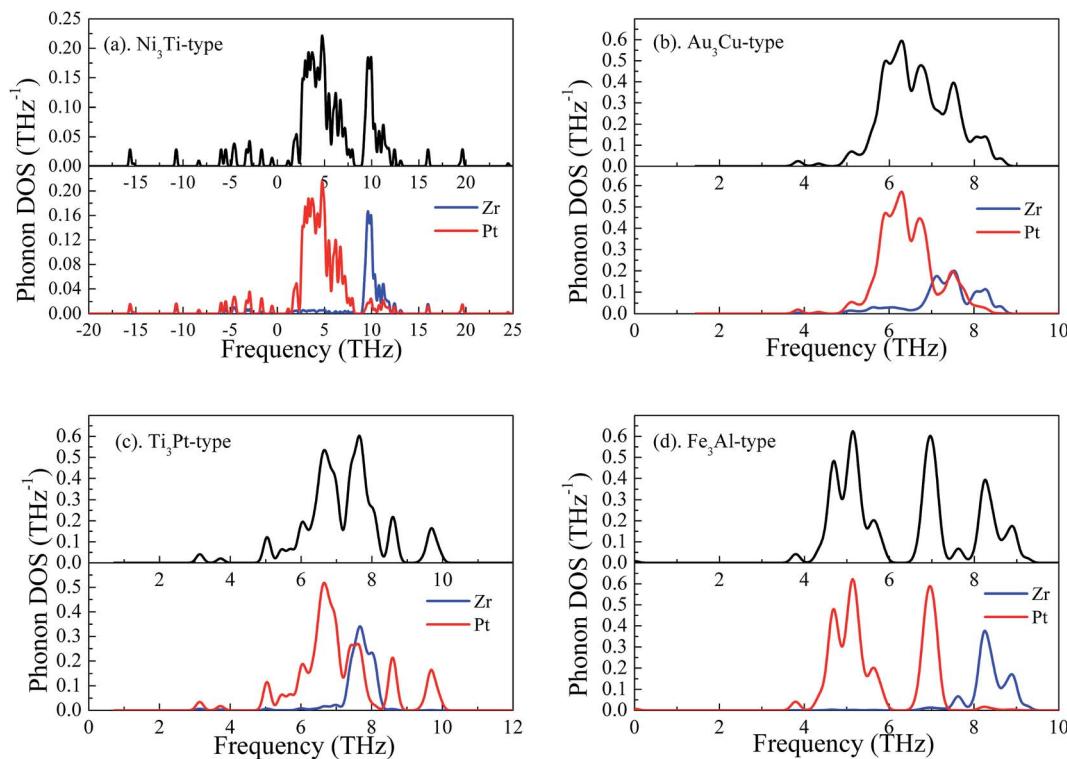


Fig. 3 Phonon density of state of  $\text{Pt}_3\text{Zr}$ , (a)  $\text{Ni}_3\text{Ti}$ -type structure, (b)  $\text{Au}_3\text{Cu}$ -type structure, (c)  $\text{Ti}_3\text{Pt}$ -type structure and (d)  $\text{Fe}_3\text{Al}$ -type structure, respectively.

For  $\text{Fe}_3\text{Al}$ -type structure, Pt atom migrates from  $(0.50, 0, 0.50)$  site to  $(0.25, 0.25, 0.25)$  site. Each Pt atom is surrounded by 4 Zr atoms and 8 Pt atoms. The calculated bond length of Pt-Zr bond is  $2.820 \text{ \AA}$ . Importantly, the network Pt-Zr bonds can improve the structural stability of  $\text{Pt}_3\text{Zr}$ .

To reveal the nature of chemical bonding, Fig. 4 shows the calculated density of state (DOS) of  $\text{Pt}_3\text{Zr}$  with four structures. We can see that the DOS profile of  $\text{Pt}_3\text{Zr}$  is composed of Pt-5d state and Zr-4d state. The strong localized hybridization between Pt and Zr forms the Pt-Zr bond. It is worth noticing

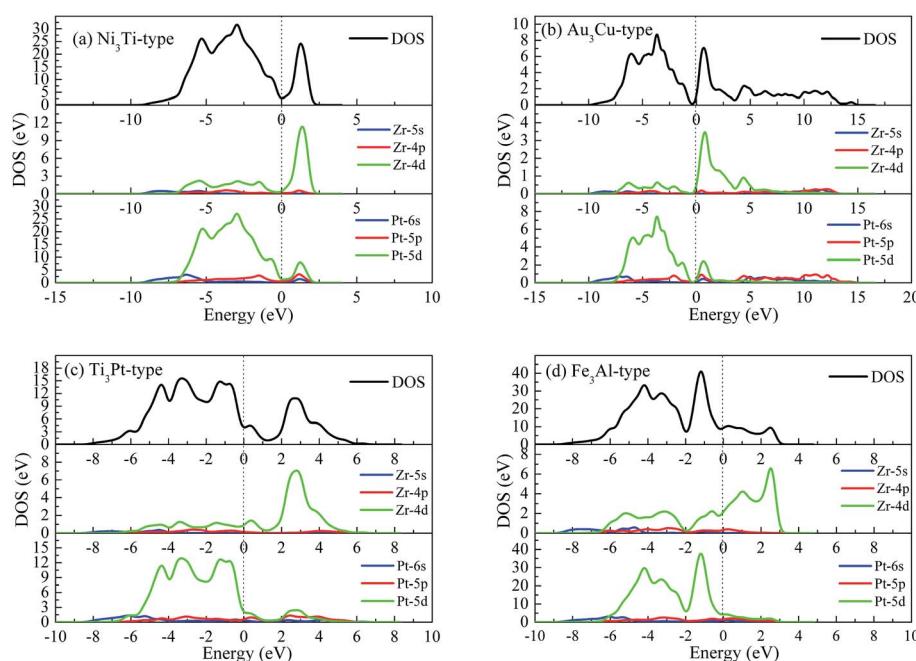


Fig. 4 Total and partial density of state (DOS) of  $\text{Pt}_3\text{Zr}$ , (a)  $\text{Ni}_3\text{Ti}$ -type structure, (b)  $\text{Au}_3\text{Cu}$ -type structure, (c)  $\text{Ti}_3\text{Pt}$ -type structure and (d)  $\text{Fe}_3\text{Al}$ -type structure, respectively.

that the DOS profile of  $\text{Ni}_3\text{Ti}$ -type structure is similar to  $\text{Au}_3\text{Cu}$ -type structure. From Fig. 4, there is a deep valley near Fermi level ( $E_F$ ), which separates the bonding state and antibonding state. For  $\text{Ti}_3\text{Pt}$ -type and  $\text{Fe}_3\text{Al}$ -type structures, however, the Zr-PDOS profile right shifts from  $E_F$  to high energy region. That is to say, Pt's migration weakens the localized hybridization between Pt and Zr. As a result, Pt-Pt bond plays an important role in structural stability, particularly for catalytic properties.

### 3.2 Oxidation mechanism

We suggest that the formation of  $\text{ZrO}_2$ -Pt interface strongly depends on the interaction between  $\text{Pt}_3\text{Zr}$  and O atom. To examine the formation of  $\text{ZrO}_2$ -Pt interface, we calculate the oxygen doped formation energy of O-doped  $\text{Pt}_3\text{Zr}$ . In particular, we discuss and analyze the chemical bonding of O-doped  $\text{Pt}_3\text{Zr}$  to reveal the formation of  $\text{ZrO}_2$ -Pt interface. It must be mentioned that the formation of  $\text{ZrO}_2$  film is related to the interstice radius of  $\text{Pt}_3\text{Zr}$ , which is determined by the atomic configuration of  $\text{Pt}_3\text{Zr}$ . Therefore, it is necessary to insight into the oxidation mechanism of  $\text{Pt}_3\text{Zr}$ .

According to the first-principles calculations, we consider the oxidation behavior of three stable  $\text{Pt}_3\text{Zr}$  structures:  $\text{Au}_3\text{Cu}$ -type,  $\text{Ti}_3\text{Pt}$ -type and  $\text{Fe}_3\text{Al}$ -type, respectively. To reveal the oxidation mechanism, we design the possible interstice models: tetrahedral interstice site (TI) and octahedral interstice site (OI), respectively. Oxygen mechanism of  $\text{Pt}_3\text{Zr}$  is measured by the oxygen doped formation energy ( $E_{\text{ad}}$ ), which is given by:

$$E_{\text{ad}} = E_{\text{Pt}_3\text{Zr}}^{\text{O}} - E_{\text{Pt}_3\text{Zr}} - \mu_{\text{O}} \quad (2)$$

where  $E_{\text{Pt}_3\text{Zr}}^{\text{O}}$  and  $E_{\text{Pt}_3\text{Zr}}$  are the total energy of O-doped  $\text{Pt}_3\text{Zr}$  and parent  $\text{Pt}_3\text{Zr}$  at the ground state, respectively.  $\mu_{\text{O}}$  is the chemical potential of O atom. Generally, the negative oxygen doped formation energy indicates the thermodynamically stable at the ground state.

To explore the catalytic activity of  $\text{Pt}_3\text{Zr}$ , we calculate and compare the capacity of oxygen between  $\text{Pt}_3\text{Zr}$  and many metals. Firstly, we explore the oxidation mechanism of  $\text{Pt}_3\text{Zr}$ . From Fig. 1,  $\text{Au}_3\text{Cu}$ -type is a typical cubic structure. Therefore, we design two possible occupied sites: TI site and OI site, respectively. However, the structural feature of  $\text{Ti}_3\text{Pt}$ -type structure is more complex than that of  $\text{Au}_3\text{Cu}$ -type structure. According to the structural feature, we design two different OI sites and one TI site. Although the doped model of  $\text{Fe}_3\text{Al}$ -type structure is similar to  $\text{Au}_3\text{Cu}$ -type structure, the atomic interaction of the former is stronger than the latter. That is to say,  $\text{Fe}_3\text{Al}$ -type structure is easily to adsorb a large number of oxygen.

Fig. 5 shows the calculated oxygen doped formation energy of O-doped  $\text{Pt}_3\text{Zr}$  and many metals. We can conclude that TI model is more thermodynamically stable than that of OI model because the calculated oxygen doped formation energy of the former is lower than the latter. This discrepancy is attributed to the localized hybridization between O and Zr. For  $\text{Pt}_3\text{Zr}$ , TI site can effectively improve the charge interaction between Zr and O, which forms the Zr-O bond. In particular, O atom prefers to occupy the TI site of  $\text{Fe}_3\text{Al}$ -type structure in comparison to other structures. Therefore, we can adjust the crystal structure of

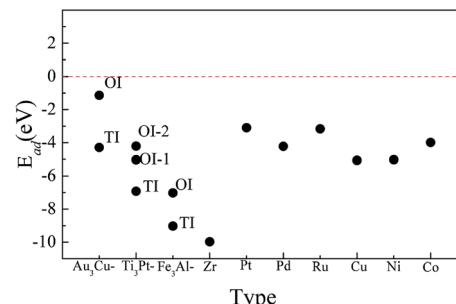


Fig. 5 Calculated oxygen adsorption energy of O-doped  $\text{Pt}_3\text{Zr}$  at tetrahedral interstice site (TI) and octahedral interstice site (OI), together with many metals.

$\text{Pt}_3\text{Zr}$  to promote the formation of  $\text{ZrO}_2$ . In particular, the calculated oxygen doped formation energy of  $\text{Fe}_3\text{Al}$ -type with TI site is about  $-9.029$  eV, which is smaller than that of other structures. This result indicates that  $\text{Pt}_3\text{Zr}$  with  $\text{Fe}_3\text{Al}$ -type structure shows the better catalytic activity in comparison to other structures. This reason is attributed to the atomic configuration of  $\text{Pt}_3\text{Zr}$ .

As we know, many metals such as Pt, Cu, Ni and Co *etc.* are important parts of solid oxide fuel cells.<sup>32-35</sup> To estimate the catalytic activity of  $\text{Pt}_3\text{Zr}$ , we compare the oxygen doped formation energy between  $\text{Pt}_3\text{Zr}$  and many metals. As shown in Fig. 5, the calculated oxygen doped formation energy of metal Zr is  $-10.068$  eV, which is smaller than that of  $\text{Pt}_3\text{Zr}$ . This result indicates that metal Zr shows better oxidation capacity in comparison to  $\text{Pt}_3\text{Zr}$ . This is why  $\text{ZrO}_2$  has been widely investigated over the last years. Importantly, although the oxygen doped formation energy of  $\text{Pt}_3\text{Zr}$  is larger than that of metal Zr, the oxygen doped formation energy of  $\text{Pt}_3\text{Zr}$  is smaller than that of other metals (see Fig. 5). The calculated electronic structure shows that the better oxidation capacity of  $\text{Pt}_3\text{Zr}$  derives from the strong localized hybridization between O and metal Zr. As mentioned above, we can predict that  $\text{Pt}_3\text{Zr}$  is expected to have better catalytic activity in comparison to other metals.

To reveal the nature of the oxidation mechanism, we further analyze the chemical bonding of O-doped  $\text{Pt}_3\text{Zr}$ . The first-principle calculations show that the bond length of Zr-O bond for  $\text{Au}_3\text{Cu}$ -type structure and  $\text{Ti}_3\text{Pt}$ -type structure is  $2.026$  Å and  $2.014$  Å, respectively, which are smaller than the corresponding Zr-O bond ( $2.05$  Å) of  $\text{ZrO}_2$ .<sup>36,37</sup> However, the calculated bond length of Zr-O bond of  $\text{Fe}_3\text{Al}$ -type structure is  $2.076$  Å, which is close to the bond length of Zr-O bond for  $\text{ZrO}_2$ . On the other hand, the calculated electronic localization density shows that there is a strong localized hybridization between Pt and O for  $\text{Au}_3\text{Cu}$ -type structure. The calculated bond length of Pt-O bond is  $2.117$  Å. However, Pt-O antibonding state in  $\text{Ti}_3\text{Pt}$ -type and  $\text{Fe}_3\text{Al}$ -type structures is found. In other words,  $\text{Ti}_3\text{Pt}$ -type and  $\text{Fe}_3\text{Al}$ -type structures are beneficial to the formation of  $\text{ZrO}_2$ . The oxidation mechanism of  $\text{Pt}_3\text{Zr}$  with three structures is shown in Fig. 6. As mentioned above, we predict that  $\text{Pt}_3\text{Zr}$  with  $\text{Fe}_3\text{Al}$ -type structure is beneficial to the formation of  $\text{ZrO}_2$ .

To gain insight the nature of oxidation behavior, Fig. 7 shows the calculated DOS of O-doped  $\text{Pt}_3\text{Zr}$  with three structures. We

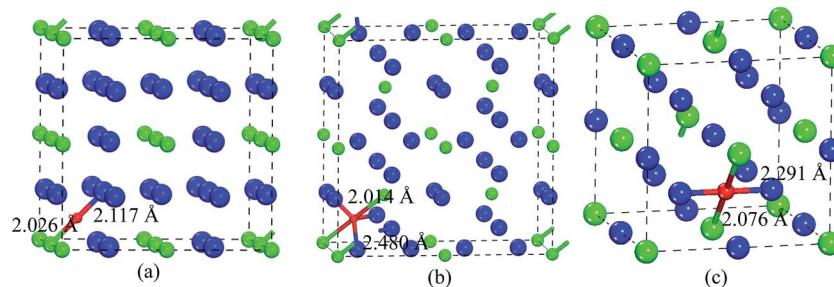


Fig. 6 Oxygen adsorption mechanism of  $\text{Pt}_3\text{Zr}$ , (a)  $\text{Au}_3\text{Cu}$ -type structure, (b)  $\text{Ti}_3\text{Pt}$ -type structure and (c)  $\text{Fe}_3\text{Al}$ -type structure, respectively.

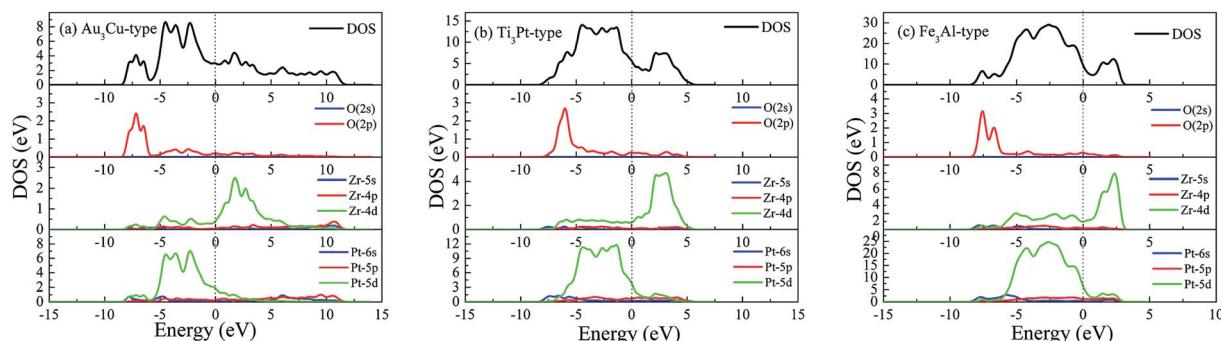


Fig. 7 Total and partial density of state (DOS) of O-doped  $\text{Pt}_3\text{Zr}$ , (a)  $\text{Au}_3\text{Cu}$ -type structure, (b)  $\text{Ti}_3\text{Pt}$ -type structure and (c)  $\text{Fe}_3\text{Al}$ -type structure, respectively.

observe that the DOS profile of  $\text{Pt}_3\text{Zr}$  is mainly contributed by Pt-5d state and Zr-4d state. The strong localized hybridization between Pt and Zr forms the Pt-Zr bond. However, oxygen addition can improve the charge equilibrium between Pt and Zr. As shown in Fig. 7, we can see that the charge interaction between O and Zr forms the Zr-O bond, which demonstrates the existence of Zr-O bond.

## 4. Conclusions

In summary, we apply first-principles calculations to study the crystal structure and oxidation mechanism of  $\text{Pt}_3\text{Zr}$ . To explore the stable structure, we calculate the formation enthalpy, phonon dispersion, lattice parameters and electronic structure of  $\text{Pt}_3\text{Zr}$ . In addition, we predict two possible new  $\text{Pt}_3\text{Zr}$  structures:  $\text{Ti}_3\text{Pt}$ -type (space group:  $\text{Pm}\bar{3}m$ , no. 223) and  $\text{Fe}_3\text{Al}$ -type (space group:  $\text{Fm}\bar{3}m$ , no. 225). To investigate the oxidation mechanism, we calculate the oxygen doped formation energy and chemical bonding of O-doped  $\text{Pt}_3\text{Zr}$ . In particular, we compare the oxygen doped formation energy of  $\text{Pt}_3\text{Zr}$  and many metals.

The calculated results show that although  $\text{Pt}_3\text{Zr}$  with hexagonal structure is more thermodynamically stable than that of cubic structure, hexagonal structure is a dynamically unstable at the ground state. The calculated oxygen doped formation energy of TI site is smaller than that of OI site. The calculated oxygen doped formation energy of  $\text{Pt}_3\text{Zr}$  with  $\text{Fe}_3\text{Al}$ -type structure is smaller than that of other structures. The calculated chemical bonding shows that  $\text{Pt}_3\text{Zr}$  with  $\text{Fe}_3\text{Al}$ -type

structure is easy to form the  $\text{ZrO}_2$  because of the formation of Zr-O bond. In particular, the calculated oxygen doped formation energy of  $\text{Fe}_3\text{Al}$ -type with TI site is smaller than that of other metals, indicating that  $\text{Pt}_3\text{Zr}$  shows the good catalytic activity in comparison to metals.

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 H. Y. T. Chen, S. Tosoni and G. Pacchioni, *ACS Catal.*, 2015, 5, 5486–5495.
- 2 L. M. Toscani, A. Raievich, M. C. Aantini, D. G. Lamas and S. A. Larrondo, *J. Phys. Chem. C*, 2016, **120**, 24165–24175.
- 3 A. P. Kulkarni, S. Giddey and S. P. S. Badwal, *J. Phys. Chem. C*, 2016, **120**, 15675–15683.
- 4 N. S. Yuzbasi, A. M. Kierzkowska, Q. Imtiaz, P. M. Abdalla, A. Kurlov, J. L. M. Rupp and C. R. Muller, *J. Phys. Chem. C*, 2016, **120**, 18977–18985.



- 5 P. Li, X. Chen, X. Wang, J. Shao, G. Lin, H. Yang, Q. Yang and H. Chen, *Energy Fuels*, 2017, **31**, 3979–3986.
- 6 A. Ganesan, M. Narayanasamy, K. Shunmugavel and I. J. Chinnappa, *Int. J. Hydrogen Energy*, 2016, **41**, 8963–8977.
- 7 F. Menegazzo, M. Signoretto, D. Marchese, F. Pinna and M. Manzoli, *J. Catal.*, 2015, **326**, 1–8.
- 8 E. Cifturek, C. D. Mcmillen, K. Sabolsky and E. M. Sabolsky, *Sens. Actuators, B*, 2015, **207**, 206–215.
- 9 J. Ftouni, A. M. Murillo, A. Goryachev, J. P. Hofmann, E. J. M. Hensen, L. Lu, C. J. Kiely, P. C. A. Bruijnincx and B. M. Weckhuysen, *ACS Catal.*, 2016, **6**, 5462–5472.
- 10 F. Giannici, G. Canu, M. Gambino, A. Longo, M. Salome, M. Viviani and A. Martorana, *Chem. Mater.*, 2015, **27**, 2763–2766.
- 11 M. Kuzminska, T. V. Kovalchuk, R. Backov and R. M. Gaigneaux, *J. Catal.*, 2014, **320**, 1–8.
- 12 S. Grieshammer, *J. Phys. Chem. C*, 2017, **121**, 15078–15084.
- 13 H. H. Li, S. Y. Ma, Q. Q. Fu, X. J. Liu, L. Wu and S. H. Yu, *J. Am. Ceram. Soc.*, 2015, **137**, 7862–7868.
- 14 Y. Sha, T. H. Yu, B. V. Merinov and W. A. Goddard, *ACS Catal.*, 2014, **4**, 1189–1197.
- 15 L. A. Avakyan, N. A. Kolpacheva, E. V. Paramonova, J. Singh, U. Hartfelder, J. A. V. Bokhoven and L. A. Bugaev, *J. Phys. Chem. C*, 2016, **120**, 28057–28066.
- 16 H. Ostrom, H. Oberg, H. Xin, J. Larue, M. Beye, M. D. Angela and A. Nilsson, *Science*, 2015, **347**, 978–982.
- 17 K. Ding, A. Gulec, A. M. Johnson, N. M. Schweitzer, G. D. Stucky, L. D. Marks and P. C. Stair, *Science*, 2015, **350**, 189–192.
- 18 Y. Pan, C. Jin and P. Mao, *J. Electron. Mater.*, 2017, **46**, 6639–6645.
- 19 H. Sohn and U. Ozkan, *Energy Fuels*, 2016, **30**, 5309–5322.
- 20 J. Durst, M. L. Haro, L. Dubau, M. Chatenet, Y. S. Olivier, L. Guetaz, P. B. Guillemaud and F. Maillard, *J. Phys. Chem. Lett.*, 2014, **5**, 434–439.
- 21 C. Zhang, S. Y. Hwang, A. Trout and Z. Peng, *J. Am. Chem. Soc.*, 2014, **136**, 7805–7808.
- 22 M. Antlanger, W. M. Schmölzer, J. Pavelec, F. Mittendorfer and J. Redinger, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2012, **86**, 035451.
- 23 J. J. Choi, W. M. Schmolzer, I. Valenti, P. Luches, F. Mittendorfer, J. Redinger, U. Diebold and M. Schmid, *J. Phys. Chem. C*, 2016, **120**, 9920–9932.
- 24 G. B. Fairbank, C. J. Humphreys, A. Kelly and C. N. Jones, *Intermetallics*, 2000, **8**, 1091–1100.
- 25 Y. Pan, Y. Lin, X. Wang, S. Chen, L. Wang, C. Tong and Z. Cao, *J. Alloys Compd.*, 2015, **643**, 49–55.
- 26 J. K. Stalick and R. M. Waterstrat, *J. Alloys Compd.*, 2007, **430**, 123–131.
- 27 M. D. Segall, P. J. D. Lindan, M. J. Probert, C. J. Pickard, P. J. Hasnip, S. J. Clark and M. C. Payne, *J. Phys.: Condens. Matter*, 2002, **14**, 2717–2744.
- 28 J. P. Perdew and Y. Wang, *Phys. Rev. B: Condens. Matter Phys.*, 1992, **45**, 13244–13249.
- 29 G. Kresse and D. Joubert, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, **59**, 1758–1775.
- 30 S. Baroni, S. d. Gironcoli, A. D. Corso and P. Giannozzi, *Rev. Mod. Phys.*, 2001, **73**, 515.
- 31 Y. Pan, W. Guan, M. Wen, J. Zhang, C. Wang and Z. Tan, *J. Alloys Compd.*, 2014, **585**, 549–554.
- 32 Z. Zheng, C. Sun, R. Dai, S. Wang, X. Wu, X. An, Z. Wu and C. Xie, *Energy Fuels*, 2017, **31**, 3091–3100.
- 33 M. Deutsch, F. Horvath, C. Knoll, D. Lager, C. G. Mayer, P. Weinberger and F. Winter, *Energy Fuels*, 2017, **31**, 2324–2334.
- 34 K. Nakao, T. Ishimoto and M. Koyama, *J. Phys. Chem. C*, 2016, **120**, 16641–16648.
- 35 X. Yang, Y. Wang, M. Li, B. Sun, Y. Li and Y. Wang, *Energy Fuels*, 2016, **30**, 2198–2203.
- 36 W. Piskorz, J. Grybos, F. Zasada, S. Cristol, J. F. Paul, A. Adamski and Z. Sojka, *J. Phys. Chem. C*, 2011, **115**, 24274–24286.
- 37 M. Taddei, F. Costantino, V. Manuali and R. Vivani, *Inorg. Chem.*, 2011, **50**, 10835–10843.