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Methanol electro-oxidation on platinum modified tungsten carbides in direct methanol fuel cells: a DFT study

Tian Sheng,^{ab} Xiao Lin,^{†a} Zhao-Yang Chen,^c P. Hu,^a Shi-Gang Sun,^b You-Qun Chu,^c Chun-An Ma^{ac} and Wen-Feng Lin^{*ad}

In exploration of low-cost electrocatalysts for direct methanol fuel cells (DMFCs), Pt modified tungsten carbide (WC) materials are found to be great potential candidates for decreasing Pt usage whilst exhibiting satisfactory reactivity. In this work, the mechanisms, onset potentials and activity for electrooxidation of methanol were studied on a series of Pt-modified WC catalysts where the bare W-terminated WC(0001) substrate was employed. In the surface energy calculations of a series of Pt-modified WC models, we found that the feasible structures are mono- and bi-layer Pt-modified WCs. The tri-layer Pt-modified WC model is not thermodynamically stable where the top layer Pt atoms tend to accumulate and form particles or clusters rather than being dispersed as a layer. We further calculated the mechanisms of methanol oxidation on the feasible models *via* methanol dehydrogenation to CO involving C–H and O–H bonds dissociating subsequently, and further CO oxidation with the C–O bond association. The onset potentials for the oxidation reactions over the Pt-modified WC catalysts were determined thermodynamically by water dissociation to surface OH* species. The activities of these Pt-modified WC catalysts were estimated from the calculated kinetic data. It has been found that the bi-layer Pt-modified WC catalysts may provide a good reactivity and an onset oxidation potential comparable to pure Pt and serve as promising electrocatalysts for DMFCs with a significant decrease in Pt usage.

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1. Introduction

Fuel cells *via* converting the chemical energy directly into electricity are a promising clean energy solution to replace traditional fossil fuel technologies.^{1–5} Liquid methanol as a fuel has some advantages over gaseous hydrogen in handling, transportation and storage. In direct methanol fuel cells (DMFCs), the ideal anodic reaction is the complete oxidation of methanol to CO₂ with the release of 6 electrons per methanol molecule (CH₃OH + H₂O → CO₂ + 6H⁺ + 6e[−]).^{1–11} Since the fuel

cell devices have to operate in strongly basic or acidic electrolyte media, the corrosion of the electrode materials is problematic, resulting in the inevitable usage of novel metals such as Pt and Pd.¹² However, the estimated global reserves for Pt-group metals are extremely low, and they are remarkably expensive for widespread application commercially.^{1–12} To overcome this challenge, the alternative low-cost catalytic materials are strongly desired to reduce the Pt loading in order to make fuel cells more commercially viable.

In the exploration of low-cost catalyst materials, transition metal carbides are found to have great potential for reducing Pt usage. Noticeably, tungsten carbide (WC) which has a similar electronic structure in the region of the Fermi level has attracted widespread attention.¹³ However, the surface states of tungsten carbide are considerably different from those of the Pt surface, and therefore the catalytic properties could not be compared directly with Pt.¹³ To date, surface modification of low-cost tungsten carbide by active metals has been extensively investigated.^{14–32} Many groups have prepared varieties of platinum modified tungsten carbide materials, which exhibit a higher reactivity than pure platinum, in hydrogen evolution reaction (HER), oxygen reduction reaction (ORR) and methanol oxidation reaction (MOR).^{14–32}

^a Centre for the Theory and Application of Catalysis (CenTACat), School of Chemistry and Chemical Engineering, Queen's University of Belfast, Belfast BT9 5AG, UK. E-mail: w.lin@qub.ac.uk

^b Collaborative Innovation Center of Chemistry for Energy Materials, State Key Laboratory of Physical Chemistry of Solid Surfaces, Department of Chemistry, College of Chemistry and Chemical Engineering, Xiamen University, Xiamen, 361005, China

^c International Sci. & Tech. Cooperation Base of Energy Materials and Application, College of Chemical Engineering and Materials Science, Zhejiang University of Technology, Hangzhou 310032, China

^d Department of Chemical Engineering, Loughborough University, Loughborough, Leicestershire, LE11 3TU, UK. E-mail: w.lin@lboro.ac.uk

[†] Current address: Peterhouse, Trumpington Street, University of Cambridge, Cambridge CB2 1RD, UK.



To understand the mechanism of electrooxidation of methanol on platinum electrodes, a lot of effort has been made in experiments. Sun *et al.* and Herrero *et al.* studied methanol electrooxidation on a series of Pt(111), Pt(110) and Pt(100) facets in acidic electrolytes.^{33,34} These studies revealed the influence of the surface structures and the anions on the catalytic decomposition of methanol. Recently, density functional theory (DFT) calculations have been widely used to understand electrochemical catalytic reactions at the atomic level.³⁵ Theoretical studies of the electrooxidation of methanol have been performed extensively.^{36–50} Zhang *et al.* showed that the decomposition of methanol could occur *via* both C–H bond and O–H bond dissociations on the closed-pack flat (111) surface.³⁶ Greeley *et al.* carried out DFT calculations by investigating the reaction energy and activation barriers of the elementary steps for methanol decomposition to CO on Pt(111).^{38,39} Ferrin *et al.* showed the surface structure sensitivity of methanol electrooxidation on transition metals.⁴⁰ Cao *et al.* studied the methanol decomposition on three well-defined low index platinum single crystal planes by combining the experimental and theoretical methods.⁴¹ The decomposition pathways in methanol oxidation were also calculated over some bimetallic surfaces such as PtAu, PtRu and PdIn.^{42–44} Kramer *et al.* presented a model of the surface kinetics of methanol dehydrogenation on transition metals.⁴⁵ Stottleyer *et al.* calculated the methanol activation on the Pt-modified WC(0001) surface *via* C–H and O–H bonds.⁴⁸

In this work, in order to understand the catalytic performance of the Pt-modified WC catalysts in a DMFC, two key issues have been considered: (i) What are the reasonable effective theoretical models to describe the Pt-modified WC structures in reality? (ii) What are the possible pathways of methanol dehydrogenation and oxidation on the Pt-modified WC surfaces and what are the onset potentials and activities for the surface reactions? With the above questions in mind, we illustrated the surface energies of a series of Pt-modified WC models to determine the thermodynamic stability with an increase in the number of surface Pt atoms. The calculated results, including the mechanisms of methanol oxidation and C–H bond and O–H bond dissociations, are discussed in detail.

2 Theoretical methods

All the electronic structure calculations were performed using the Vienna Ab-initio Simulation Package (VASP) with the exchange–correlation functional of Perdew–Burke–Ernzerhof (PBE). The projector-augmented-wave (PAW) pseudopotentials were utilized to describe the core electron interaction. Geometry optimization was carried out using the BFGS algorithm.^{51–59} The cut-off energy was 400 eV and a $4 \times 4 \times 1$ Monkhorst–Pack k -point sampling was used. The transition states were located using a constrained optimization approach with the force converge criterion below $0.05 \text{ eV } \text{Å}^{-1}$ in modified VASP.^{60–62} Since the WC(0001) surface has been confirmed to be W-terminated which is more stable than C-terminated,⁴⁸ three layers of W-terminated WC(0001) as the $p(3 \times 3)$ substrate including 27 W and 27 C atoms was used

in the calculations with the bottom two layers being fixed while the top layer was allowed to relax during calculations to interact with Pt. The vacuum region was $\sim 12 \text{ Å}$ to ensure that there is little interaction between slabs. For modeling the Pt-modified WC models, mono-, bi- and tri-layer Pt atoms were placed epitaxially on the W-terminated WC(0001) substrate with 9, 18 and 27 Pt atoms, respectively. All the Pt atoms were allowed to relax in the calculations. For each adsorbate, different binding sites (top, bridge, and hollow) were calculated to determine the most stable bonding configuration.

In this work, the adsorption energy was defined as

$$E_{\text{ad}} = E_{\text{adsorbate/slab}} - E_{\text{adsorbate}} - E_{\text{slab}} \quad (1)$$

where $E_{\text{adsorbate/slab}}$, $E_{\text{adsorbate}}$, and E_{slab} are the total energies of the adsorbate binding with the surface, the gaseous adsorbate and the clean surface, respectively.

For the calculation of the OH* formation potential,^{60–63} the reaction free energy change (ΔG) in the formation of OH* according to the water dissociation reaction of $\text{H}_2\text{O} \rightarrow \text{OH}^* + \text{H}^+ + \text{e}^-$ was calculated. The free energy was obtained using $G = E + \text{ZPE} + \text{TS}$, in which E is the total energy of species, S is the entropy and ZPE is the zero point energy at room temperature. Therefore, the free energy change of the reaction $\text{H}_2\text{O} \rightarrow \text{OH}^* + \text{H}^+ + \text{e}^-$ was derived as $\Delta G = G(\text{OH}^*) + G(\text{H}^+ + \text{e}^-) - G(\text{H}_2\text{O})$. When the electrode potential is 0 V, pH = 0 ($[\text{H}^+] = 1 \text{ M}$), at 298 K, due to the equilibrium of $\text{H}^+ + \text{e}^- \rightarrow \frac{1}{2}\text{H}_2$, we can use the free energy of gaseous $\frac{1}{2}\text{H}_2$ to replace that of $\text{H}^+ + \text{e}^-$. Two correction terms were introduced: the pH of the electrolyte ($-\text{pH}kT \ln 10$) and the electrode potential (eU) referring to the standard hydrogen electrode (SHE), resulting in $G(\text{H}^+ + \text{e}^-) = G(\frac{1}{2}\text{H}_2) - \text{pH}kT \ln 10 + \text{eU}$. When $\Delta G = 0$, the reaction $\text{H}_2\text{O} \rightarrow \text{OH}^* + \text{H}^+ + \text{e}^-$ is in equilibrium and the OH* formation potential can be obtained.

For the calculation of the surface energy, the definition of eqn (2), proposed by Boettger *et al.*,⁶⁴ was employed in this work:

$$E_{\text{Pt,surf}} = 1/n(E_{\text{Pt/substrate}} - E_{\text{substrate}} - nE_{\text{Pt,bulk}}) \quad (2)$$

where $E_{\text{Pt/substrate}}$ is the total energy of the slab, $E_{\text{substrate}}$ is the total energy of the substrate, n is the number of Pt atoms, and $E_{\text{Pt,bulk}}$ is the bulk energy per Pt atom obtained from an independent bulk calculation. In the calculation of the surface energy, every possible position of Pt atoms was tested and the most stable structure was used.

3. Results

3.1 Surface energies of Pt-modified WC(0001)

In this work, the models containing a three-layer W-terminated WC(0001) slab as the core and layers of Pt atoms as the shell were employed in the calculations. To quantify the thermodynamic stability of a series of Pt-modified WC(0001) surfaces, surface energies of a series of structures were calculated using eqn (2), with an increase of Pt loadings from mono-layer to tri-layers. Through comparing the calculated energy per Pt atom in the shell with an equivalent energy per Pt atom in the bulk as



the reference, the stability of Pt atoms adsorbed on the substrate could be assessed effectively. Specifically, if the surface energy is lower than zero, the dispersed Pt atoms on WC(0001) would be favoured. On the other hand, if the surface energy is higher than 0, the dispersed Pt atoms would prefer to accumulate to form clusters. The assessments can be summarized as follows:

$$E_{\text{Pt,surf}} < 0, \text{ Pt atoms} \rightarrow \text{Pt layer}$$

$$E_{\text{Pt,surf}} > 0, \text{ Pt atoms} \rightarrow \text{Pt cluster}$$

The calculated surface energies and the structures of each model with the top and side views at different Pt coverages are presented in Fig. 1. On mono-layer Pt-modified WC, the Pt_{1ML}/WC(0001) surface, the Pt atoms favour to occupy the hcp sites rather than the fcc sites, and the calculated surface energy is -1.25 eV per atom, indicating that Pt atoms are favoured to be dispersed over W-terminated WC(0001) since the chemical bonds between the Pt atoms and W atoms are considerably strong. On the bi-layer Pt-modified WC surface, Pt_{2ML}/WC(0001), Pt atoms could alternatively occupy the fcc or hcp sites on Pt_{1ML}/WC(0001) with the same surface energy of -0.01 eV, which is close to zero, implying that both surface structures are likely to exist. With an increase of Pt atoms on tri-layers, the most stable structure would be such that the top layer Pt atoms occupy the fcc sites on Pt_{2ML,hcp}/WC(0001). However, as shown in Fig. 1a, the calculated surface energy is noticeably higher than zero,

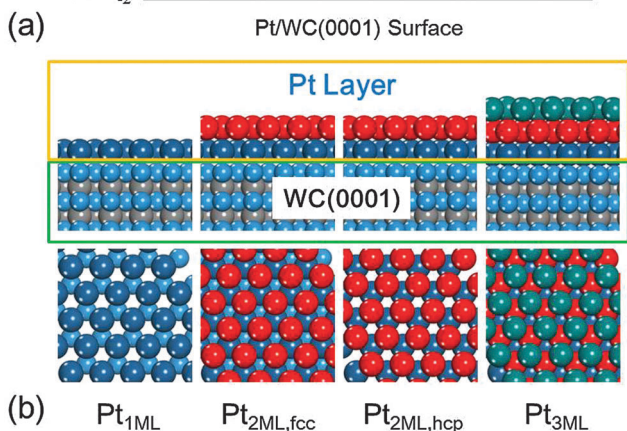
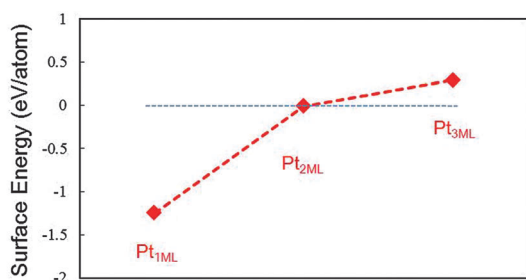


Fig. 1 (a) The calculated surface energies (in eV per atom) of a series of Pt-modified WC(0001) surfaces. (b) Top and side views of a series of Pt-modified WC(0001) surfaces with the increase of Pt atoms. Cyan: W; grey: C; blue: first layer Pt; red: second layer Pt; green: third layer Pt. (The same colors were used throughout this work.)

indicating that the Pt atoms of the third layers cannot be effectively dispersed but tend to accumulate forming Pt clusters or particles on Pt_{2ML}/WC(0001). According to these calculated results, we concluded that in reality mono- and bi-layer Pt-modified WC surfaces should commonly exist, depending on the amount of Pt loaded. Therefore, Pt_{1ML}/WC(0001) and Pt_{2ML}/WC(0001) as shown in Fig. 1b were used for the calculations in the study of the electrooxidation of methanol.

3.2 Methanol dehydrogenation

In the electrooxidation of methanol in DMFCs, the reactions could be divided into methanol dehydrogenation to surface CO adsorbates and the further CO oxidation in the presence of surface oxidants produced from water dissociation. Since in the dehydrogenation of CH₃OH* to CO*, four C-H bonds and one O-H bond need to be cleaved subsequently, two main pathways are considered: C-H and O-H pathways. To highlight the dehydrogenation mechanisms, we mapped out an overall reaction network systematically as shown in Fig. 2, including all the possible paths in the methanol dehydrogenation involving 10 elementary reactions and 8 intermediates: methanol (CH₃OH*), hydroxymethyl (CH₂OH*), hydroxymethylene (CHOH*), hydroxymethylidyne (COH*), methoxy (CH₃O*), formaldehyde (CH₂O*), formyl (CHO*) and carbon monoxide (CO*). In this section, since there are two structures of bi-layer Pt-modified WC surfaces, the calculations in the methanol dehydrogenation were performed on both Pt_{2ML,fcc}/WC(0001) and Pt_{2ML,hcp}/WC(0001). In order to compare with the methanol dehydrogenation on pure Pt, Pt(111) was also employed. All the calculated data for the Pt_{1ML}/WC(0001), Pt_{2ML,fcc}/WC(0001), Pt_{2ML,hcp}/WC(0001) and Pt(111), including kinetic barriers (E_a) and reaction energies (ΔE), are listed in Table 1. The most favoured reaction paths are highlighted in Fig. 2 with the kinetic data listed. The energy profiles for methanol dehydrogenation are presented in Fig. 3 whilst the optimized structures of intermediates and transition states involved in the reaction network are shown in Fig. 4.

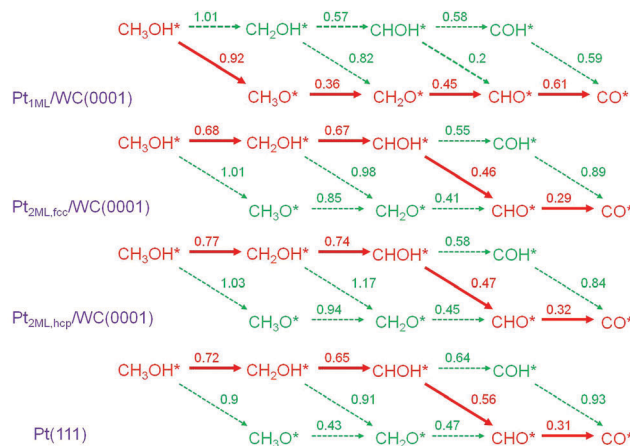


Fig. 2 Reaction networks and the calculated reaction barriers (in eV) for methanol dehydrogenation from CH₃OH* to CO* + 4H* over Pt_{1ML}/WC(0001), Pt_{2ML,fcc}/WC(0001), Pt_{2ML,hcp}/WC(0001) and Pt(111) surfaces. The paths in red color present the favored pathways in methanol dehydrogenation with the intermediates involved and the barriers shown.



Table 1 Calculated reaction barriers (E_a , in eV) and reaction energies (ΔE , in eV) of the elementary steps in methanol dehydrogenation to surface CO* and the further CO* oxidation over Pt_{1ML}/WC(0001), Pt_{2ML,fcc}/WC(0001), Pt_{2ML,hcp}/WC(0001) and Pt(111) surfaces

Surface reactions	Pt _{1ML} /WC		Pt _{2ML,fcc} /WC		Pt _{2ML,hcp} /WC		Pt	
	E_a	ΔE	E_a	ΔE	E_a	ΔE	E_a	ΔE
CH ₃ OH* → CH ₂ OH* + H*	1.01	0.29	0.68	-0.44	0.77	-0.18	0.72	-0.21
CH ₃ OH* → CH ₃ O* + H*	0.92	0.22	1.01	0.08	1.03	0.42	0.9	0.63
CH ₂ OH* → CHOH* + H*	0.57	0.43	0.67	-0.57	0.74	-0.2	0.65	-0.17
CH ₂ OH* → CH ₂ O* + H*	0.82	0.07	0.98	0.29	1.17	0.53	0.91	0.51
CH ₃ O* → CH ₂ O* + H*	0.36	0.13	0.85	-0.23	0.94	-0.07	0.43	-0.33
CHOH* → COH* + H*	0.58	0.14	0.55	-0.95	0.58	-0.72	0.64	-0.51
CHOH* → CHO* + H*	0.2	-0.61	0.46	-0.23	0.47	-0.09	0.56	-0.11
CH ₂ O* → CHO* + H*	0.45	-0.26	0.41	-1.1	0.45	-0.82	0.47	-0.79
CHO* → CO* + H*	0.61	-0.68	0.29	-1.4	0.32	-1.04	0.31	-0.73
COH* → CO* + H*	0.59	-1.43	0.89	-0.68	0.84	-0.42	0.93	-1.26
CO* + OH* → COOH*	0.49	0.04	0.58	-0.24	0.59	-0.24	0.43	-0.34

On Pt_{1ML}/WC(0001), methanol initially adsorbs on the surface *via* O–Pt bonding at the top site with an adsorption energy of -0.28 eV. Then, the adsorbed methanol CH₃OH* prefers to break the O–H bond forming CH₃O* with an activation energy of 0.92 eV; the O–H bond length is 1.50 Å at the transition state. On the other hand, C–H bond of CH₃OH* is harder to be activated to form CH₂OH* as a higher barrier of 1.01 eV is required with the C–H bond length of 1.64 Å at the transition state. The formation of both CH₃O* and CH₂OH* are endothermic by 0.22 eV and 0.29 eV, respectively. All in all, the initial dehydrogenation of CH₃OH* *via* the O–H bond is favourable than *via* the C–H bond, both kinetically and thermodynamically.

Therefore, the CH₃O* adsorbed at the top site was identified to be a reactive intermediate for producing CH₂O* with a low C–H bond dissociation barrier of 0.36 eV; the C–H bond length is 1.86 Å at the transition state. Once CH₂O* is formed, it readily overcomes the 0.45 eV barrier to further dehydrogenate to CHO* with the C–H bond length being 1.53 Å at the transition state. The final dehydrogenation step of CHO* to CO* takes place with a barrier of 0.61 eV and the distance of the C–H bond being 1.41 Å at the transition state. Overall, the pathway for methanol dehydrogenation on Pt_{1ML}/WC(0001) is as follows:



For the dehydrogenation of methanol on Pt_{2ML,fcc}/WC(0001) and Pt_{2ML,hcp}/WC(0001), the reaction mechanisms are generally the same over the two surfaces, but are different from those over Pt_{1ML}/WC(0001). Herein, we take methanol dehydrogenation on Pt_{2ML,fcc}/WC(0001) as an example. In contrast to the initial activation of methanol *via* the O–H bond on Pt_{1ML}/WC(0001), CH₃OH* is more easily dehydrogenated to CH₂OH* *via* the C–H bond here with a reaction barrier of 0.68 eV instead of *via* the O–H bond to CH₃O* for which the barrier would be much higher as 1.01 eV. At the transition state, the C–H bond length is 1.48 Å. The formation of CH₂OH* is exothermic by -0.44 eV, whilst the formation of CH₃O* is slightly endothermic by 0.08 eV. For the further dehydrogenation of CH₂OH* adsorbed at the top site, the C–H bond pathway was identified to be favoured with a reaction barrier of 0.67 eV and the C–H bond length is 1.43 Å at the transition state. The barrier is much lower than that of 0.98 eV *via* the O–H bond pathway, where the length of the O–H bond is 1.90 Å at the transition state. The so formed CHOH* adsorbs at the bridge site and undergoes further dehydrogenation *via* its O–H bond dissociation to yield CHO* with a lower barrier of 0.46 eV. If the C–H bond dissociation occurs to yield COH*, a higher barrier of 0.55 eV must be overcome; the formation of COH* is exothermic by -0.95 eV, whilst the formation of CHO* by -0.23 eV. The produced CHO* favours the top site and readily decomposes to CO* overcoming a small barrier of 0.29 eV with the C–H bond length of 1.39 Å at the transition state. On the other hand, if COH* is formed, it is considerably stable at the fcc site resulting in the difficulty in activating its O–H bond, which needs to overcome

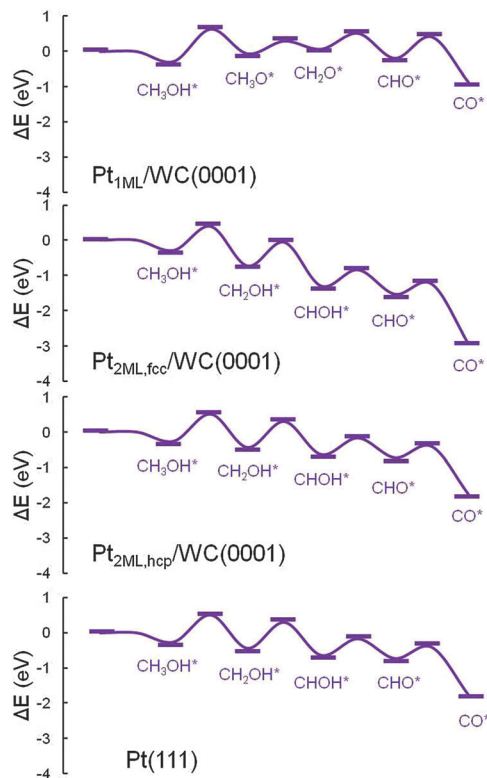


Fig. 3 Energy profiles for the dehydrogenation of methanol from CH₃OH* to CO* + 4H* over Pt_{1ML}/WC(0001), Pt_{2ML,fcc}/WC(0001), Pt_{2ML,hcp}/WC(0001) and Pt(111) surfaces.



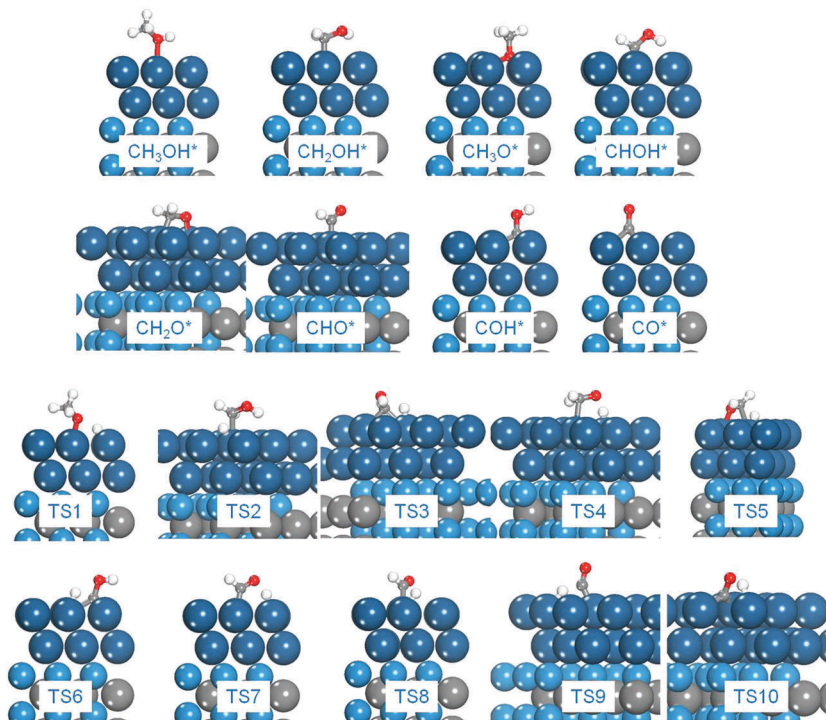
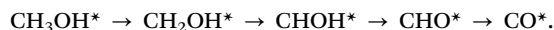


Fig. 4 Optimized structures of intermediates and transition states in methanol electro-oxidation on the $\text{Pt}_{2\text{MLfcc}}/\text{WC}(0001)$ surface as an example. TS1: $\text{CH}_3\text{OH}^* \rightarrow \text{CH}_3\text{O}^* + \text{H}^*$; TS2: $\text{CH}_2\text{OH}^* \rightarrow \text{CH}_2\text{OH}^* + \text{H}^*$; TS3: $\text{CH}_2\text{OH}^* \rightarrow \text{CHOH}^* + \text{H}^*$; TS4: $\text{CH}_2\text{OH}^* \rightarrow \text{CH}_2\text{O}^* + \text{H}^*$; TS5: $\text{CH}_3\text{O}^* \rightarrow \text{CH}_2\text{O}^* + \text{H}^*$; TS6: $\text{CHOH}^* \rightarrow \text{COH}^* + \text{H}^*$; TS7: $\text{CHOH}^* \rightarrow \text{CHO}^* + \text{H}^*$; TS8: $\text{CH}_2\text{O}^* \rightarrow \text{CHO}^* + \text{H}^*$; TS9: $\text{CHO}^* \rightarrow \text{CO}^* + \text{H}^*$; TS10: $\text{COH}^* \rightarrow \text{CO}^* + \text{H}^*$; TS11: $\text{CO}^* + \text{OH}^* \rightarrow \text{COOH}^*$. Cyan: W; grey: C; blue: Pt; white: H; red: O. (The same colors were used throughout this work.)

a barrier of 0.89 eV in forming CO^* with the O–H bond length of 1.25 Å at the transition state. Therefore, the overall methanol dehydrogenation pathway on $\text{Pt}_{2\text{MLfcc}}/\text{WC}(0001)$ follows these steps:



Comparing the calculated data for methanol dehydrogenation over the $\text{Pt}_{2\text{MLhcp}}/\text{WC}(0001)$ and $\text{Pt}_{2\text{MLfcc}}/\text{WC}(0001)$ surfaces, the dehydrogenation barriers calculated on the hcp surface are only slightly higher than those on the fcc surface in the range 0.01–0.09 eV. For instance, the initial C–H bond dissociation barrier in CH_3OH^* is 0.77 eV on $\text{Pt}_{2\text{MLhcp}}/\text{WC}(0001)$ and 0.68 eV on $\text{Pt}_{2\text{MLfcc}}/\text{WC}(0001)$; the O–H bond breaking barrier in CHOH^* is 0.47 eV on $\text{Pt}_{2\text{MLhcp}}/\text{WC}(0001)$ and 0.46 eV on $\text{Pt}_{2\text{MLfcc}}/\text{WC}(0001)$. Therefore, the activity of methanol dehydrogenation on $\text{Pt}_{2\text{MLfcc}}/\text{WC}(0001)$ is only slightly higher than that on $\text{Pt}_{2\text{MLhcp}}/\text{WC}(0001)$.

It is worth pointing out that we have also calculated the reaction energies and barriers for the dehydrogenation of methanol on Pt(111) as a reference to compare with the calculated data on Pt-modified WC. On Pt(111), the reaction pathways are the same to those on bi-layer Pt-modified WC(0001) surfaces as $\text{CH}_3\text{OH}^* \rightarrow \text{CH}_2\text{OH}^* \rightarrow \text{CHOH}^* \rightarrow \text{CHO}^* \rightarrow \text{CO}^*$. The initial activation of CH_3OH^* to form CH_2OH^* is endothermic by -0.21 eV with a reaction barrier of 0.72 eV. In the subsequent dehydrogenation processes, CH_2OH^* overcomes a barrier of 0.65 eV to produce CHOH^* at the bridge site and then the so produced CHOH^* continues to break the O–H bond with a barrier of 0.56 eV to yield CHO^* rather than to break C–H bond with a barrier of 0.64 eV to

yield COH^* . The final dehydrogenation step of CHO^* to CO^* requires to overcome a small barrier of 0.31 eV to break the C–H bond of CHO^* and it is exothermic by -0.73 eV.

3.3 Onset potential for surface oxidant formation and CO oxidation

It is well known that electrooxidation of surface CO^* at high potentials requires the dissociation of water to form the active surface oxidant OH^* to turn over CO^* to CO_2 . Water dissociation involves the separation of a proton and an electron ($\text{H}_2\text{O} \rightarrow \text{OH}^* + \text{H}^+ + \text{e}^-$). Therefore, the onset potential of OH^* formation usually coincides with the onset potential of surface oxidation reaction.^{44,62} Herein, in order to estimate the onset potentials for the CO electrooxidation on pure Pt(111) and Pt-modified WC surfaces, we have calculated the OH^* formation potentials on these surfaces. On Pt(111), the calculated OH^* formation potential is 0.64 V (*vs.* SHE) at pH = 0. On $\text{Pt}_{1\text{ML}}/\text{WC}(0001)$, OH^* forms at a very low onset potential of -0.26 V (*vs.* SHE), which implies that this surface is easily covered by OH^* . On $\text{Pt}_{2\text{MLfcc}}/\text{WC}(0001)$ and $\text{Pt}_{2\text{MLhcp}}/\text{WC}(0001)$ surfaces, the calculated onset potentials for OH^* formation are 0.67 V (*vs.* SHE) and 0.66 V (*vs.* SHE), respectively, which are very close to the potential (0.64 V) obtained on Pt(111), indicating that bi-layer Pt-modified WC surfaces have a similar ability to facilitate water dissociation as that of pure Pt(111). Table 2 lists the calculated onset potentials for OH^* formation (U_{OH^*} , in V *vs.* SHE) on $\text{Pt}_{1\text{ML}}/\text{WC}(0001)$, $\text{Pt}_{2\text{MLfcc}}/\text{WC}(0001)$, $\text{Pt}_{2\text{MLhcp}}/\text{WC}(0001)$ and Pt(111) surfaces for comparison.



Table 2 The calculated one-set potentials for OH* formation (U_{OH^*} , in V vs. SHE), the barriers ($E_{\text{a},1}$, in eV) of the rate-determining step in methanol dehydrogenation and the barriers ($E_{\text{a},2}$, in eV) in CO oxidation on Pt_{1ML}/WC(0001), Pt_{2ML,fcc}/WC(0001), Pt_{2ML,hcp}/WC(0001) and Pt(111) surfaces, respectively

	Pt _{1ML} /WC	Pt _{2ML,fcc} /WC	Pt _{2ML,hcp} /WC	Pt
U_{OH^*}	-0.26	0.67	0.66	0.64
$E_{\text{a},1}$	0.92	0.68	0.77	0.72
$E_{\text{a},2}$	0.49	0.58	0.59	0.4

For the surface oxidative reaction processes of CO* + OH*, CO* adsorbs at the fcc hollow site and OH* stays at the top site initially. At the transition state, CO* moves towards the top site to couple with OH* for producing COOH*, which is readily dehydrogenated to CO₂ in the presence of OH*. The overall reaction barrier on Pt(111) is 0.43 eV, which is relatively low for the reaction to occur at room temperature. For mono- and bi-(fcc or hcp) layer Pt-modified WC surfaces, the calculated barriers are 0.49 eV on Pt_{1ML}/WC(0001), 0.58 eV on Pt_{2ML,fcc}/WC(0001) and 0.59 eV on Pt_{2ML,hcp}/WC(0001); all of these are slightly higher than those on Pt(111) (0.40 eV) implying that CO* oxidation to CO₂ still readily occurs in the presence of OH*.

We found that the surface CO* adsorption energy is -1.96 eV at the fcc site on Pt_{2ML,fcc}/WC(0001) which is stronger than -1.76 eV obtained on Pt_{2ML,hcp}/WC(0001), but a similar barrier is obtained on the fcc and hcp sites, *i.e.*, 0.58 eV and 0.59 eV, respectively. According to a previous work, the geometry effect plays an important role in the total barriers.⁵⁹ From the changes in the structures between the initial and transition states, it was evidenced that CO* diffused from the fcc to top sites for coupling but OH* still remained at the top sites. Therefore, we examined the CO* adsorption energy at the top sites and found that the adsorption energy is -1.61 eV on Pt_{2ML,fcc}/WC(0001) and -1.43 eV on Pt_{2ML,hcp}/WC(0001). The corresponding energy barriers for CO* diffusion over these two surfaces are 0.35 eV and 0.33 eV, respectively, which indicates that there is a similar CO* geometry effect on the two surfaces, the latter in turn results in similar CO* + OH* barriers on both surfaces. In addition, the reaction energy for the formation of COOH* on both surfaces is the same, -0.24 eV. According to the BEP relationship for bridging the kinetics and thermodynamics,⁵⁸ two reactions with the same reaction energy usually encounter a similar kinetic barrier. Once COOH* forms, it can be readily transferred to CO₂ in the presence of OH* *via* the proton transfer.⁶⁷ Therefore, the CO adsorption energy hardly affects the CO oxidation rate on the Pt_{2ML}/WC(0001) surface at higher potentials where the DMFC operates.

As listed in Table 2, compared with the barriers of 0.68 eV–0.92 eV in methanol dehydrogenation to produce CO*, the oxidative removal of CO* with the lower barriers of 0.4–0.59 eV should be much faster than the methanol dehydrogenation; therefore, in the overall methanol electrooxidation reaction, the rate-determining step should be the C–H bond activation of methanol but not CO oxidation. The calculated onset potentials (U_{OH^*}), reaction barriers ($E_{\text{a},1}$) of the rate-determining

step in methanol dehydrogenation and the barriers ($E_{\text{a},2}$) in CO oxidation on Pt_{1ML}/WC(0001), Pt_{2ML,fcc}/WC(0001), Pt_{2ML,hcp}/WC(0001) and Pt(111) surfaces are listed together in Table 2 for a clear comparison among the four surfaces.

3.4 CO oxidation at the interface

It is worth mentioning that, for understanding and screening improved Pt-based catalysts for methanol electrooxidation, the bi-functional mechanism is widely considered in a binary catalyst in which the surface Pt site catalyses the methanol dehydrogenation and the second element activates water to provide oxidants for surface CO* oxidation.^{62,65,66} Both W-terminated WC(0001) and Pt_{1ML}/WC(0001) surfaces are easily oxidized and we therefore examined another possibility for the CO* oxidation at the interface between Pt and WC, in which WC(0001) or Pt_{1ML}/WC(0001) provides surface oxidants (O*/OH* formed from water oxidation) and the edge of the Pt island adsorbs the CO* formed from methanol dehydrogenation. In this case, CO* oxidation occurs at the interface instead of the flat surface. For modelling the Pt–WC(0001) interface comprising the WC(0001) and Pt island, we expanded the Pt_{1ML}/WC(0001) unit cell from $p(3 \times 3)$ to $p(3 \times 6)$ and removed half of the surface Pt atoms to create the interface. The same approach was used for building the Pt–Pt_{1ML}/WC(0001) interface, and we found that the second Pt atom prefers to occupy the fcc sites at the interface. The two interface structures are shown in Fig. 5. At the Pt–WC(0001) interface, the bare W-terminated WC(0001) surface was identified to be occupied extremely easily by O* and the O binding energy is around -4.0 eV larger than that on pure Pt(111), indicating that once the clean WC(0001) is exposed to water or air, it would be oxidized immediately. The formation of CO₂ from the CO*/Pt-edge and O*/WC(0001) is endothermic by 2.55 eV, mostly due to the overly strong adsorption of O*, which indicates that the adsorbed O* on WC(0001) is not reactive towards CO* oxidation. Therefore, this path is not energetically favourable. On the other hand, at the Pt–Pt_{1ML}/WC(0001) interface, OH* is formed at a potential above -0.26 V (*vs.* SHE), also implying that the OH* binding energy is much stronger in comparison with that on Pt_{2ML}/WC(0001) (where the onset potential for OH* formation is 0.66–0.67 V). Furthermore, CO*

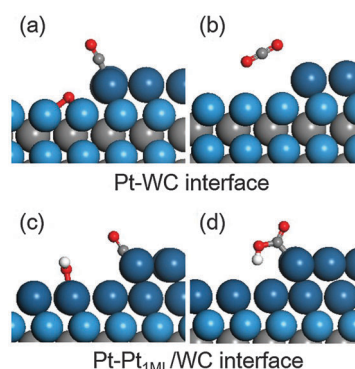


Fig. 5 Optimized structures for surface CO* oxidation at the Pt–WC(0001) and Pt–Pt_{1ML}/WC(0001) interfaces: (a) CO*/Pt and O*/WC; (b) CO₂ product; (c) CO*/Pt-edge and OH*/Pt_{1ML}/WC; (d) COOH*/Pt-edge.



adsorption at the edge site is even stronger with an adsorption energy of -2.22 eV in comparison with -1.96 eV at the flat $\text{Pt}_{2\text{ML},\text{fcc}}/\text{WC}(0001)$. Herein, CO^* oxidation *via* the coupling of CO^* and OH^* is endothermic by 0.69 eV, indicating that CO^* oxidation can hardly occur at this interface. The structures of the initial and final states at the two interfaces are shown in Fig. 5. Therefore, on the Pt modified WC catalyst, the bi-functional aspect for methanol electrooxidation is not feasible. We would suggest that the further modification on the $\text{Pt}_{2\text{ML}}/\text{WC}(0001)$ surface with an additional element like Ru or Sn may enable the bi-functional mechanism and further enhance the reactivity.

4. Discussion

4.1 Activity of the Pt-modified WC surfaces towards methanol oxidation

To understand the theoretical activity of these Pt-modified WC surfaces towards methanol oxidation under realistic conditions, we calculated the turnover frequency (TOF, in s^{-1}) using the Arrhenius equation, $\text{TOF} = k_{\text{B}}T/h \exp(-E_{\text{a}}/RT)$, to roughly estimate the catalyst/surface performance in the DMFC, where E_{a} is the activation energy for methanol oxidation to CO_2 , in which the initial activation barrier of methanol dehydrogenation is the rate determining step as illustrated above in the reaction pathways. Since the surface energies of $\text{Pt}_{2\text{ML},\text{fcc}}/\text{WC}(0001)$ and $\text{Pt}_{2\text{ML},\text{hcp}}/\text{WC}(0001)$ are the same, the positions of the second layer Pt atoms have the same possibility to occupy the fcc and hcp sites on $\text{Pt}_{1\text{ML}}/\text{WC}(0001)$. The TOF of methanol oxidation on bi- (fcc or hcp) layer Pt-modified WC(0001) is therefore calculated using the equation $\text{TOF}_{2\text{ML}} = \frac{1}{2}\text{TOF}_{2\text{ML},\text{fcc}} + \frac{1}{2}\text{TOF}_{2\text{ML},\text{hcp}}$, considering the same contributions from the fcc and hcp surfaces to the total activity. The theoretical activity on mono- and bi-layer Pt-modified WC(0001) surfaces and Pt(111) are presented in Fig. 6. It can be seen from the figure that the activity on $\text{Pt}_{1\text{ML}}/\text{WC}(0001)$ is extremely low, having only 0.2% of the Pt(111) activity. It could be understood based on the fact that, due to the significantly strong chemical interactions between the Pt atoms and WC(0001) on $\text{Pt}_{1\text{ML}}/\text{WC}(0001)$, inferred from the lowest surface energy, the

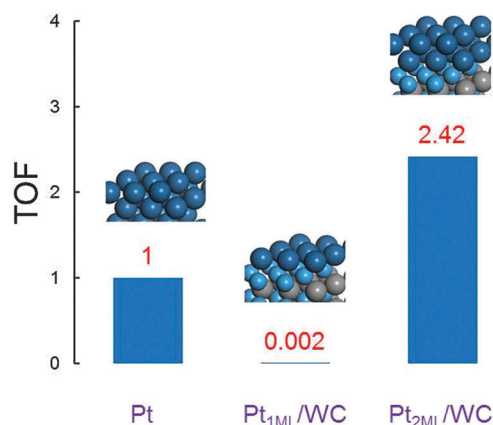


Fig. 6 The calculated normalized turnover frequency TOF in the electro-oxidation of methanol on Pt(111), $\text{Pt}_{1\text{ML}}/\text{WC}(0001)$ and $\text{Pt}_{2\text{ML}}/\text{WC}(0001)$ surfaces, respectively.

Pt atoms here are too stable to be active for methanol dehydrogenation. However, on bi- (fcc or hcp) layer Pt-modified WC(0001) surfaces, the activity is increased significantly to about 2.42 times that of Pt(111). The latter data suggest that the bi-layer Pt-modified WC materials could provide a better performance in the electrooxidation of methanol than pure Pt. These results are in fact supported by the recent experimental data which show that the onset potential is hardly changed, but the activity is increased significantly for the Pt/WC catalysts compared to that of pure Pt catalysts.^{26,32}

4.2 The effect of potential

In methanol electrocatalysis, there are two elementary reactions involved in the first dehydrogenation process: (i) $\text{CH}_3\text{OH}^* \rightarrow \text{CH}_2\text{OH}^* + \text{H}^*$ and (ii) $\text{H}^* \rightarrow \text{H}^+ + \text{e}^-$.^{68–70} It can be seen that the surface H^* electrooxidative stripping in (ii) enables the removal of H^* to refresh the surface with the current created. Liu's group has reported that the first reaction (i) is hardly affected by the potential, but the second step (ii) with the proton and electron transfer is indeed sensitive to the potential.^{68–70} The electrooxidative removal of H^* in the electrolyte at potentials where the DMFC operates (*e.g.* above 0.20 V vs. SHE) is relatively quicker and the rate-determining step is (i), supported by the evidence observed in the isotope experiments.⁷¹ Therefore, we only compare the dehydrogenation barriers in (i) between the Pt modified WC and pure Pt for the methanol dehydrogenation.

4.3 Water effect

Since the methanol surface reactions occur in aqueous solution, we further investigated the effect of water on the initial dehydrogenation of methanol, which is the rate-determining step. The simulation of water structures is still a huge challenge in electrocatalysis, the main reason being that the water structure is always fluctuating at the dynamics scale. Usually, the choice of a statistic water structure is a good approximation for generally understanding the role of water.^{60,72–74} Here we compare the adsorption energy and dehydrogenation barrier through the $[\text{CH}_3\text{OH} \cdots \text{H}_2\text{O}]$ complex structure. All the calculated data, including the adsorption energy, the bond distance and the dehydrogenation barrier, are listed in Table 3. The complex is trapped on the $\text{Pt}_{2\text{ML},\text{fcc}}/\text{WC}(0001)$ surface at the top site with an adsorption energy of -0.82 eV, and the optimized structure with a H-down configuration of H_2O is shown in Fig. 7. It has been found that the O–Pt bond is shortened, from 2.46 Å without H_2O to 2.29 Å due to the water effect, indicating that water increases

Table 3 The calculated adsorption energy (E_{ads} , in eV), the distance of the O–Pt bond (in Å), the reaction barrier (E_{a} , in eV) and the distance of the C–H bond at the transition state of the $[\text{CH}_3\text{OH} \cdots \text{H}_2\text{O}]$ complex on $\text{Pt}_{2\text{ML},\text{fcc}}/\text{WC}(0001)$, $\text{Pt}_{2\text{ML},\text{hcp}}/\text{WC}(0001)$ and Pt(111) surfaces, respectively. The values in parentheses are the data obtained without the H_2O effect

	$\text{Pt}_{2\text{ML},\text{fcc}}/\text{WC}$	$\text{Pt}_{2\text{ML},\text{hcp}}/\text{WC}$	Pt
E_{ads}	-0.82 (-0.3)	-0.76 (-0.27)	-0.75 (-0.24)
$d(\text{O–Pt})$	2.29 (2.46)	2.30 (2.48)	2.28 (2.38)
E_{a}	0.72 (0.68)	0.79 (0.77)	0.75 (0.72)
$d(\text{C–H})$	1.44 (1.47)	1.46 (1.49)	1.45 (1.47)



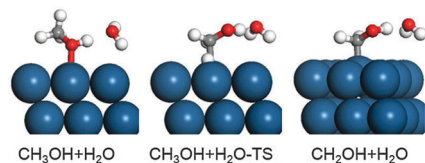


Fig. 7 Optimized structures in the dehydrogenation of the $[\text{CH}_3\text{OH}\cdots\text{H}_2\text{O}]$ complex.

the interaction between methanol and the surface Pt site. On Pt(111), this phenomenon is also observed where the O–Pt bond, which is originally 2.38 Å in the absence of water, decreases to 2.28 Å in the presence of H_2O , yielding a significant adsorption energy of -0.75 eV. Looking at all the three surfaces studied, the adsorption energy of the $[\text{CH}_3\text{OH}\cdots\text{H}_2\text{O}]$ complex has increased by 0.49–0.52 eV in comparison with CH_3OH adsorption in the absence of water, indicating that the water effect on the adsorption of methanol on the three surfaces studied is very similar.

In terms of the dehydrogenation barrier of $[\text{CH}_3\text{OH}\cdots\text{H}_2\text{O}]^*$ to $[\text{CH}_2\text{OH}\cdots\text{H}_2\text{O}]^*$, on $\text{Pt}_{2\text{ML},\text{fcc}}/\text{WC}(0001)$, the barrier increased from 0.68 eV to 0.72 eV and the C–H bond distance at the transition state elongated from 1.44 Å to 1.47 Å due to the presence of water; on $\text{Pt}_{2\text{ML},\text{hcp}}/\text{WC}(0001)$, the barrier also increased slightly from 0.77 eV to 0.79 eV; on Pt(111), the barrier increased from 0.72 eV to 0.75 eV. From the data listed in Table 3, it can be seen clearly that the barrier only increased by around 0.02–0.04 eV, showing that water only slightly slows down the C–H bond dissociation rate. Because the C–H bond is relatively hydrophobic and the water which forms hydrogen bonds with the OH group hardly affects the C–H bond dissociation barriers, the slight influence of water is thought not to be unreasonable. More importantly, it provides evidence that water could equally affect the adsorption and kinetic barrier for the methanol electrooxidation on Pt(111) and Pt/WC(0001) surfaces. All in all, our data suggest that the role of water is indeed significant in determining the absolute adsorption and kinetic values in electrocatalysis, but it does not significantly affect the assessment of the general trend in the inherent activity of catalysts.

5. Conclusions

In this work, the electrooxidation of methanol in a DMFC has been investigated on a series of Pt-modified WC(0001) model surfaces. Firstly, through the calculation of surface energies, we found that the most stable Pt coverage is the mono- or bi- (fcc or hcp) layer but not the tri-layer Pt-modified WC(0001) which is considerably unstable thermodynamically. Then, on the three possible surfaces of $\text{Pt}_{1\text{ML}}/\text{WC}(0001)$, $\text{Pt}_{2\text{ML},\text{fcc}}/\text{WC}(0001)$ and $\text{Pt}_{2\text{ML},\text{hcp}}/\text{WC}(0001)$, we calculated the reaction network in the methanol dehydrogenation involving the intermediates CH_3OH^* , CH_2OH^* , CHOH^* , COH^* , CH_3O^* , CH_2O^* ; CHO^* and CO^* via the four C–H and one O–H bond dissociations subsequently. It has been found that the favourable methanol dehydrogenation paths on $\text{Pt}_{1\text{ML}}/\text{WC}(0001)$ are $\text{CH}_3\text{OH}^* \rightarrow \text{CH}_3\text{O}^* \rightarrow \text{CH}_2\text{O}^* \rightarrow \text{CHO}^* \rightarrow \text{CO}^*$ but those on $\text{Pt}_{2\text{ML}}/\text{WC}(0001)$ and Pt(111) are

$\text{CH}_3\text{OH}^* \rightarrow \text{CH}_2\text{OH}^* \rightarrow \text{CHOH}^* \rightarrow \text{CHO}^* \rightarrow \text{CO}^*$. The initial activation of methanol was found to be the rate-determining step. Subsequently CO^* can be readily oxidized to CO_2 in the presence of surface OH^* at high potentials where the DMFC operates. The on-set potentials for the surface OH^* formation through water dissociation were calculated for the Pt-modified WC surfaces and it has been found that bi- (fcc or hcp) layer Pt-modified WC catalysts have similar onset potentials compared with that on pure Pt(111), but exhibit up to 2.4 times higher reactivity compared to that of pure Pt. Our theoretical data were supported by recent experimental results and suggests that bi-layer Pt-modified WC catalysts could be a promising low cost high performance electrocatalysts for DMFC application with the benefit of reducing Pt usage.

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References

- 1 M. Baldauf and W. Preidel, *J. Power Sources*, 1999, **84**, 161.
- 2 C. Lamy, A. Lima, V. LeRhun, F. Delime, C. Coutanceau and J. M. Leger, *J. Power Sources*, 2002, **105**, 283.
- 3 Z. G. Shao, W. F. Lin, P. A. Christensen and H. M. Zhang, *Int. J. Hydrogen Energy*, 2006, **31**, 1914.
- 4 Z. G. Shao, F. Y. Zhu, W. F. Lin, P. A. Christensen and H. M. Zhang, *J. Power Sources*, 2006, **161**, 813.
- 5 U. B. Demirci, *J. Power Sources*, 2007, **169**, 239.
- 6 Z. G. Shao, F. Y. Zhu, W. F. Lin, P. A. Christensen, H. M. Zhang and B. L. Yi, *J. Electrochem. Soc.*, 2006, **153**, A1575.
- 7 Z. G. Shao, W. F. Lin, F. Y. Zhu, P. A. Christensen, H. M. Zhang and B. L. Yi, *J. Power Sources*, 2006, **160**, 1003.
- 8 Z. G. Shao, W. F. Lin, F. Y. Zhu, P. A. Christensen and H. M. Zhang, *Phys. Chem. Chem. Phys.*, 2006, **8**, 2720.
- 9 H. X. Liu, N. Tian, M. P. Brandon, J. Pei, Z. Huangfu, C. Zhan, Z. Y. Zhou, C. Hardacre, W. F. Lin and S. G. Sun, *Phys. Chem. Chem. Phys.*, 2012, **14**, 16415.
- 10 W. F. Lin, P. A. Christensen and A. Hamnett, *Phys. Chem. Chem. Phys.*, 2001, **3**, 3312.
- 11 M. Winter and R. J. Brodd, *Chem. Rev.*, 2004, **104**, 4245.
- 12 S. A. Grigoriev, P. Millet and V. N. Fateev, *J. Power Sources*, 2008, **177**, 281.
- 13 R. B. Levy and M. Boudart, *Science*, 1973, **181**, 547.
- 14 D. V. Esposito and J. G. Chen, *Energy Environ. Sci.*, 2011, **4**, 3900.
- 15 H. H. Hwu and J. G. Chen, *Chem. Rev.*, 2005, **105**, 185.
- 16 M. B. Zellner and J. G. Chen, *Surf. Sci.*, 2004, **569**, 89.
- 17 D. V. Esposito, S. T. Hunt, A. L. Stottlemeyer, K. D. Dobson, B. E. McCandless, R. W. Birkmire and J. G. Chen, *Angew. Chem., Int. Ed.*, 2010, **49**, 9859.



- 18 D. V. Esposito, S. T. Hunt, Y. C. Kimmel and J. G. Chen, *J. Am. Chem. Soc.*, 2012, **134**, 3025.
- 19 C. Ma, J. Sheng, N. Brandon, C. Zhang and G. Li, *Int. J. Hydrogen Energy*, 2007, **32**, 2824.
- 20 M. Wu, P. K. Shen, Z. D. Wei, S. Q. Song and M. Nie, *J. Power Sources*, 2007, **166**, 310.
- 21 D. J. Ham, R. Ganesan and J. S. Lee, *Int. J. Hydrogen Energy*, 2008, **33**, 6865.
- 22 Y. Hara, N. Minami, H. Matsumoto and H. Itagaki, *Appl. Catal., A*, 2007, **332**, 289.
- 23 M. K. Jeon, K. R. Lee, W. S. Lee, H. Daimon, A. Nakahara and S. I. Woo, *J. Power Sources*, 2008, **185**, 927.
- 24 D. J. Ham, Y. K. Kim, S. H. Han and J. S. Lee, *Catal. Today*, 2008, **132**, 117.
- 25 M. Nie, P. K. Shen, M. Wu, Z. Wei and H. Meng, *J. Power Sources*, 2006, **162**, 173.
- 26 R. Wang, Y. Xie, K. Shi, J. Wang, C. Tian, P. Shen and H. Fu, *Chem. – Eur. J.*, 2012, **18**, 7443.
- 27 M. D. Obradovic, B. M. Babic, V. R. Radmilovic, N. V. Krstajic and S. L. Gojkovic, *Int. J. Hydrogen Energy*, 2012, **37**, 10671.
- 28 Z. J. Mellinger, T. G. Kelly and J. G. Chen, *ACS Catal.*, 2012, **2**, 751.
- 29 C. K. Poh, S. H. Lim, Z. Tian, L. Lai, Y. P. Feng, Z. Shen and J. Lin, *Nano Energy*, 2013, **2**, 28.
- 30 H. Meng and P. K. Shen, *Chem. Commun.*, 2005, 4408.
- 31 C. Ma, W. Liu, M. Shi, X. Lang, Y. Chu, Z. Chen, D. Zhao, W. Lin and C. Hardacre, *Electrochim. Acta*, 2013, **114**, 133.
- 32 Z. Y. Chen, C. A. Ma, Y. Q. Chu, J. M. Jin, X. Lin, C. Hardacre and W. F. Lin, *Chem. Commun.*, 2013, **49**, 11677.
- 33 S. G. Sun and J. Clavilier, *J. Electroanal. Chem.*, 1987, **236**, 95.
- 34 E. Herrero, K. Franaszczuk and A. Wieckowski, *J. Phys. Chem.*, 1994, **98**, 5074.
- 35 B. Braunschweig, D. Hibbitts, M. Neurock and A. Wieckowski, *Catal. Today*, 2013, **202**, 197.
- 36 C. J. Zhang and P. Hu, *J. Chem. Phys.*, 2001, **115**, 7182.
- 37 S. K. Desai, M. Neurock and K. Kourtakis, *J. Phys. Chem. B*, 2002, **106**, 2559.
- 38 J. Greeley and M. Mavrikakis, *J. Am. Chem. Soc.*, 2002, **124**, 7193.
- 39 J. Greeley and M. Mavrikakis, *J. Am. Chem. Soc.*, 2004, **126**, 3910.
- 40 P. Ferrin and M. Mavrikakis, *J. Am. Chem. Soc.*, 2009, **131**, 14381.
- 41 D. Cao, G. Q. Lu, A. Wieckowski, S. A. Wasileski and M. Neurock, *J. Phys. Chem. B*, 2005, **109**, 11622.
- 42 D. W. Yuan, X. G. Gong and R. Q. Wu, *J. Chem. Phys.*, 2008, **128**, 64706.
- 43 J. Ye, C. Liu and Q. Ge, *Phys. Chem. Chem. Phys.*, 2012, **14**, 16660.
- 44 B. Y. Liu, J. M. Jin, X. Lin, C. Hardacre, P. Hu, C. A. Ma and W. F. Lin, *Catal. Today*, 2015, **242**, 230.
- 45 Z. C. Kramer, X. K. Gu, D. Y. Zhou, W. X. Li and R. T. Skodje, *J. Phys. Chem. C*, 2014, **118**, 12364.
- 46 R. Jiang, W. Guo, M. Li, D. Fu and H. Shan, *J. Phys. Chem. C*, 2009, **113**, 4188.
- 47 A. Morgan, R. Kavanagh, W. F. Lin, C. Hardacre and P. Hu, *Phys. Chem. Chem. Phys.*, 2013, **15**, 20170.
- 48 A. L. Stottlemeyer, P. Liu and J. G. Chen, *J. Chem. Phys.*, 2010, **133**, 104702.
- 49 C. Ma, T. Liu and L. Chen, *Appl. Surf. Sci.*, 2010, **256**, 7400.
- 50 D. D. Vasic, I. A. Pasti and S. V. Mentus, *Int. J. Hydrogen Energy*, 2013, **38**, 5009.
- 51 G. Kresse and J. Hafner, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1993, **48**, 13115.
- 52 G. Kresse and J. Furthmuller, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1996, **54**, 11169.
- 53 G. Kresse and J. Hafner, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1993, **47**, 558.
- 54 G. Kresse and J. Hafner, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **49**, 14251.
- 55 G. Kresse and J. Furthmuller, *Comput. Mater. Sci.*, 1996, **6**, 15.
- 56 P. E. Blochl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**, 17953.
- 57 (a) G. Kresse and D. Joubert, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, **59**, 1758; (b) A. Alavi, P. Hu, T. Deutsch, P. L. Silvestrelli and J. Hutter, *Phys. Rev. Lett.*, 1998, **80**, 3650.
- 58 (a) J. P. Pedrew and K. Burke, *Phys. Rev. Lett.*, 1996, **77**, 3865; (b) A. Michadelides, Z. P. Liu, C. J. Zhang, A. Alavi, D. A. King and P. Hu, *J. Am. Chem. Soc.*, 2003, **125**, 3704.
- 59 (a) J. P. Pedrew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1997, **78**, 1396; (b) Z. P. Liu and P. Hu, *J. Am. Chem. Soc.*, 2003, **125**, 1958.
- 60 J. Rossmeisl, J. K. Norskov, C. D. Taylor, M. J. Janik and M. Neurock, *J. Phys. Chem. B*, 2006, **110**, 21833.
- 61 J. K. Norskov, J. Rossmeisl, A. Logadottir, L. Lindqvist, J. R. Kitchin, T. Bligaard and H. Jonsson, *J. Phys. Chem. B*, 2004, **108**, 1520.
- 62 J. M. Jin, T. Sheng, X. Lin, R. Kavanagh, P. Hamer, P. Hu, C. Hardacre, A. Martinez-Bonastre, J. Sharman, D. Thompsett and W. F. Lin, *Phys. Chem. Chem. Phys.*, 2014, **16**, 9432.
- 63 T. Sheng, W. F. Lin, C. Hardacre and P. Hu, *Phys. Chem. Chem. Phys.*, 2014, **16**, 13248.
- 64 J. C. Boettger, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **16**, 798.
- 65 P. Ferrin, A. U. Nilekar, J. Greeley, M. Mavrikakis and J. Rossmeisl, *Surf. Sci.*, 2008, **602**, 3424.
- 66 J. Rossmeisl, P. Ferrin, G. A. Tritsarlis, A. U. Nilekar, S. Koh, S. E. Bae, S. R. Brankovic, P. Strasser and M. Mavrikakis, *Energy Environ. Sci.*, 2012, **5**, 8335.
- 67 X. Q. Gong, P. Hu and R. Raval, *J. Chem. Phys.*, 2003, **119**, 6324.
- 68 Y. H. Fang and Z. P. Liu, *J. Am. Chem. Soc.*, 2010, **132**, 18214.
- 69 Y. H. Fang and Z. P. Liu, *Surf. Sci.*, 2015, **631**, 42.
- 70 Y. H. Fang, G. F. Wei and Z. P. Liu, *J. Phys. Chem. C*, 2013, **117**, 669.
- 71 J. Ren, Y. Y. Yang, B. W. Zhang, N. Tian, W. B. Cai, Z. Y. Zhou and S. G. Sun, *Electrochem. Commun.*, 2013, **37**, 49.
- 72 Y. H. Fang, G. F. Wei and Z. P. Liu, *Catal. Today*, 2013, **202**, 98.
- 73 X. Sun, X. Cao and P. Hu, *Sci. China: Chem.*, 2015, **58**, 55.
- 74 T. Sheng, W. F. Lin, C. Hardacre and P. Hu, *J. Phys. Chem. B*, 2014, **118**, 5762.

