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Elucidation of the surface structure–selectivity relationship in ethanol electro-oxidation over platinum by density functional theory†

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We have successfully built a general framework to comprehend the structure–selectivity relationship in ethanol electrooxidation on platinum by density functional theory calculations. Based on the reaction mechanisms on three basal planes and five stepped surfaces, it was found that only (110) and $n(111) \times (110)$ sites can enhance CO₂ selectivity but other non-selective step sites are more beneficial to activity.

The use of direct ethanol fuel cells (DEFCs) is a hopeful future energy solution for directly converting chemical energy into electricity to replace the usage of fossil fuels. Liquid ethanol, which can be sustainably produced from biomass, has a high energy density and is stored and transported easily compared to hydrogen gas. However, there are still challenges hindering DEFCs' widespread applications commercially, in particular, the slow ethanol oxidative kinetics and formation of some poisoning intermediates. Platinum catalysts are the most common in fuel cell systems operating under strongly acidic conditions and thus, understanding of the catalytic processes on platinum catalysts is fundamentally important.^{1–7} The complete oxidation of ethanol to CO₂ is the ideal anodic reaction in DEFCs, with 12 electron transfer ($\text{C}_2\text{H}_5\text{OH} + 3\text{H}_2\text{O} \rightarrow 2\text{CO}_2 + 12\text{H}^+ + 12\text{e}^-$), but in reality, acetaldehyde ($\text{C}_2\text{H}_5\text{OH} \rightarrow \text{CH}_3\text{CHO} + 2\text{H}^+ + 2\text{e}^-$) and acetic acid ($\text{C}_2\text{H}_5\text{OH} + \text{H}_2\text{O} \rightarrow \text{CH}_3\text{COOH} + 4\text{H}^+ + 4\text{e}^-$) predominate the final products, with only 2 and 4 electron transfer, respectively.^{1–7} The optimal catalyst for ethanol oxidation is inevitably linked with CO₂ selectivity which is expected to be highly active towards cleaving the C–C bond at low potentials. Unfortunately, the production of CO₂ is identified in the region of 0.5–7.5% on platinum catalysts in real systems,² which is far below the desirable selectivity for implementation of the technology.

Understanding the structure–selectivity relationship at atomic levels is one of the fundamental scientific challenges in surface science and catalysis, which is of great significance for rationally designing catalysts. For practical catalysts with a high complexity of surface structures, what sites may actually govern the product distribution is largely unresolved due to the current technological limitation in revealing the detailed selectivity at different surface sites. Surface structural effects on ethanol electrooxidation have been studied by using single-crystal planes or nanoparticles enclosed by unique facets, and the low-coordination sites on the surface have been identified to facilitate the reaction rate.^{8–16} However, in terms of selectivity, the low-coordination sites do not always promote CO₂ selectivity. Experiments have shown that the formation of acetic acid was also enhanced by the presence of low-coordination sites.^{8,11} The formation of CO₂ is sensitive to the surface structure, and surfaces with terraces of (111) symmetry separated by monoatomic (110) step sites were found to be very active for C–C bond breaking since the amounts of CO/CO₂ produced increase with the step density.^{8–12,15,16}

The Brønsted–Evans–Polanyi (BEP) relationship reveals that the thermodynamics of any catalytic reaction controls the kinetics in general.^{17–20} Since CO₂ is considerably more stable than acetaldehyde and acetic acid as the major products in ethanol electrooxidation, the thermodynamically favored CO₂ production is expected to be faster kinetically than acetic acid production, but in fact the CO₂ selectivity is rather low. The underlying reason behind the low CO₂ selectivity is believed to be the kinetics, which may significantly determine the reaction route. Therefore, a fundamental understanding of ethanol selective oxidation reactions is significant for elucidating the structure–selectivity relationship at the atomic level, also shedding light on the selectivity concerned in many other electrocatalytic systems.

To date, a general framework for describing the structure–selectivity relationship in ethanol electrooxidation is still lacking, and many puzzles regarding the catalytic selectivity of different sites have not been rationalized yet. In particular, the following questions remain to be solved: What is the key factor determining the complete oxidative (to form CO₂) or partial oxidative

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In contrast, in the β -dehydrogenation pathway for the formation of CH_2CO^* , which does not involve the formation of CH_3CO^* , the barrier is much reduced. The comparison of the overall barriers in the two pathways is presented in Fig. 3. It is apparent that the barriers in the β -dehydrogenation pathway (the red columns) are all lower than those in the α -dehydrogenation

Fig. 2 Reaction mechanism for ethanol selective dehydrogenation via the α -dehydrogenation pathway (blue) or the β -dehydrogenation pathway (red). The key intermediate structures on Pt(111) are displayed.

Fig. 3 Comparison of the overall barriers (in eV) for CH_3CHOH^* dehydrogenation to CH_2CO^* via the α -dehydrogenation (blue) and β -dehydrogenation (red) pathways on a series of Pt surfaces.

pathway (the blue columns). The smallest overall barrier difference between the two pathways is obtained on Pt(111), *i.e.*, 0.02 eV via the β -dehydrogenation pathway and 0.89 eV via the α -dehydrogenation pathway, indicating that the reaction rates for the formation of CH_2CO^* via the two pathways are rather close. But on the other surfaces, the advantage of the β -dehydrogenation pathway in kinetics is considerable owing to a lower barrier of 0.16–0.35 eV, which would result in increasing the rate at about two to four orders of magnitude compared with that via the α -dehydrogenation pathway at room temperature. It is therefore suggested that the β -dehydrogenation pathway is more favoured kinetically for C–C bond breaking.

Whether CH_2CO^* or CHCO^* is a reasonable precursor for C–C bond splitting is found to be dependent on the surface structure.³⁰ We have thus examined the two possible pathways: (i) $\text{CH}_2\text{CO}^* \rightarrow \text{CHCO}^* \rightarrow \text{CH}^* + \text{CO}^*$ and (ii) $\text{CH}_2\text{CO}^* \rightarrow \text{CH}_2^* + \text{CO}^*$, respectively. If the dehydrogenation barrier is lower than the C–C bond breaking barrier, it would suggest the dehydrogenation to CHCO^* could occur first. The comparison barriers between paths (i) and (ii) are displayed in Fig. S4 (ESI†). From our results, on Pt(111), Pt(100) and Pt(310), the mechanism follows path (i), while on other five surfaces, the C–C bond breaking in CH_2CO^* may occur directly without involving the formation of CHCO^* . According to the favoured pathways, the C–C bond breaking barriers on different surfaces are presented in Fig. S5 (ESI†). It can be seen that all the barriers are 0.48–0.77 eV, indicating that the C–C bond breaking rate is reasonable kinetically. Once CH_2^* is produced from CH_2CO^* , it would be further decomposed into CH^* which is stable on the surface since the further C^* formation is prohibited by thermodynamics or kinetics. As a result of C–C bond breaking, one would expect the final C1 products to be CH^* and CO^* on major surfaces.

From the above kinetic analyses, it is shown that the selectivity towards CO^* formation *versus* that towards CH_3CO^* formation on platinum is determined by the competition between the elementary step of $\text{CH}_3\text{CHOH}^* \rightarrow \text{CH}_3\text{COH}^*$ and the step of $\text{CH}_3\text{CHOH}^* \rightarrow \text{CH}_2\text{CHOH}^*$. The barriers in C–C bond breaking steps being presented are reasonably low, such that the origin of low CO_2 selectivity should date back to the selective dehydrogenation

in CH_3CHOH^* . Only when the β -dehydrogenation pathway is preferred kinetically than the α -dehydrogenation pathway in CH_3CHOH^* , ethanol C–C bond breaking can occur. We thus defined the selectivity as the difference between the barrier of α -dehydrogenation ($E_{\alpha,\alpha\text{-CH}}$) and that of β -dehydrogenation ($E_{\alpha,\beta\text{-CH}}$) as follows:

$$\Delta E_a = E_{\alpha,\alpha\text{-CH}} - E_{\alpha,\beta\text{-CH}}$$

Quantitatively, the higher ΔE_a indicates the more favoured β -dehydrogenation and the higher possibility of forming C1 products.²³ The comparison of the selectivities obtained on a series of Pt surfaces is shown in Fig. 4, based on the above definition. It is found that the stepped Pt(110) surface possesses the highest selectivity with ΔE_a being 0.13 eV, indicating that the predominant products (>99%, estimated by the Boltzmann distribution) on the surface would be CH^* and CO^* at low potentials. Upon increasing the width of the (111) terrace, (110) step sites could still hold the selectivity. On Pt(331), the selectivity is reduced to 82%, lower than that on Pt(110). Although having the same local structure of (110) sites, the presence of (111) terrace lowers the selectivity, indicating that the width of the terrace may affect the intrinsic properties of the step sites.

In the $[01\bar{1}]$ zone, ΔE_a is –0.22 eV on Pt(111) and –0.19 eV on Pt(100), respectively, suggesting that hardly any C1 product can form on the surface. The monoatomic (100) step on the (111) terrace, as the most common defect on surfaces, presents a moderate ability to break the C–C bond, although it is not comparable with the selectivities on Pt(110) and Pt(331); about 32% ethanol on the Pt(211) step would decompose into CH^* and CO^* finally. However, the monoatomic (111) step on the (100) terrace cannot promote CO_2 selectivity. The selectivity obtained on Pt(511) is almost the same as that on Pt(100). Herein, in the $[01\bar{1}]$ zone, only upon increasing the density of $n(111) \times (100)$ step sites, the selectivity can be enhanced. In the $[001]$ zone, despite the selectivities on Pt(320) and Pt(310) being better than that on Pt(100), the former two surfaces still cannot provide any C1 product, indicating that the increase of step sites in this zone is not beneficial to CO_2 production.

The overall barriers in their kinetically favoured pathways can depict appropriately the activity in ethanol dehydrogenation on different surfaces. It can be seen from Table 1 that Pt(111) is the least active for ethanol dehydrogenation with the highest barrier

Fig. 4 Comparison of the selectivity (ΔE_a , in eV) on a series of Pt surfaces.

Table 1 Calculated activity (E_a , in eV) and selectivity (ΔE_a , in eV) on a series of Pt surfaces

	(111)	(211)	(511)	(100)	(310)	(320)	(110)	(331)
E_a	0.78	0.54	0.51	0.64	0.54	0.52	0.68	0.70
ΔE_a	-0.22	-0.02	-0.18	-0.19	-0.17	-0.12	0.14	0.04

of 0.78 eV and Pt(100) is more active with a lower barrier of 0.64 eV. Pt(211), Pt(511), Pt(310) and Pt(320) steps are the most active where the barriers are between 0.51 and 0.54 eV. The activities on Pt(110) and Pt(331) are similar with the overall barriers being 0.68 and 0.70 eV, which are determined by the C–C bond breaking barriers. The activities are in the following order:

$$(111) < (110) \approx (331) < (100) < (310) \approx (211) \approx (320) \approx (511)$$

From these results, it is understood that the presence of (110) or $n(111) \times (110)$ step sites may give rise to a high CO_2 selectivity on practical catalysts, successfully explaining the experimental observations.^{8–12,15,16} However, the activities of such sites are less than other typical step sites, on which the partial oxidation to acetic acid is preferred. The reactive selectivity is strongly dependent on the local structure of the step sites and the low-coordination sites are not always active for C–C bond breaking. These findings highlight the need for control of the surface structure that will allow facile C–C bond breaking but hardly provide a high activity at the same time. It is reasonable to infer that how to balance the selectivity and activity will be more crucial for making platinum catalysts more efficient. It is hardly possible that pure platinum catalysts are sufficiently active for complete ethanol electrooxidation to CO_2 without acetic acid as the partial oxidation product in DEFCs.

In summary, we have successfully built a general framework to comprehend the surface structure–selectivity relationship in ethanol electrooxidation on platinum catalysts. The reasonable formation mechanisms for the C1 products are pointed out *via* the β -dehydrogenation pathway from CH_3CHOH^* . The selectivity is defined as the barrier difference based on the kinetic analyses. It is found that only (110) and $n(111) \times (110)$ step sites may enhance CO_2 selectivity but other non-selective step sites are more beneficial to the activity towards partial oxidation. This work could help gain more insights into the catalytic processes on practical catalysts with various surface sites, which is essential for the search for new highly active and selective catalysts.

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