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Interface catalytic regulation *via* electron rearrangement and hydroxyl radicals triggered by oxygen vacancies and heavy metal ions†

Xiang-Yu Xiao,^{ab} Zong-Yin Song,^{ab} Chong-Chong Zhang,^c Yong-Huan Zhao,^{ab} Zhi-Wei Gao,^{ab} Shi-Hua Chen,^a Pei-Hua Li,^a Yu-Feng Sun,^c Meng Yang^{*a} and Xing-Jiu Huang ^{ab}

Although the enhanced intrinsic activities of some nano-metal oxides are obtained by manufacturing oxygen vacancies (OVs), the effect of multiple roles of OVs is ambiguous. Herein, an interface catalytic regulation *via* electron rearrangement and hydroxyl radicals ($\cdot\text{OH}$) was proposed with the designed ZrO_2 hollow sphere rich in OVs (V_\circ -rich ZrO_2). Surprisingly, it was shown that the catalytic ability of V_\circ -rich ZrO_2 was 9.9 times higher than that of ZrO_2 with little OVs in electrochemical catalytic reduction of $\text{Pb}(\text{II})$. It was found that the generation of Zr^{2+} and Zr^{3+} caused by OVs results in the rearrangement of abundant free electrons to facilitate the catalytic reaction rates. The longer bond length between V_\circ -rich ZrO_2 and reactants, and the lower adsorption energy are beneficial for reactants to desorb, improving the conversion rates. Besides, the produced $\cdot\text{OH}$ were captured which were induced by OVs and trace divalent heavy metal ions in *in situ* electron paramagnetic resonance (EPR) experiments, contributing to lowering the energy barriers. This study not only revealed the enhanced interface catalytic effect of electron rearrangement and generated $\cdot\text{OH}$ triggered by OVs, but also provided unique insights into interface catalytic regulation on nano-metal oxides simulated by OVs.

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1 Introduction

Nano-metal oxide catalysts offering intriguing properties and stability, and low price are the subject of intense investigation, particularly in the energy and environment-related catalysis field.^{1–4} However, some nano-metal oxides, like TiO_2 , CeO_2 , and ZrO_2 , suffer from poor catalytic efficiency due to their low electron transfer rates and limited active sites, greatly restricting their application.^{5–8} To this end, manufacturing oxygen vacancies (OVs) on them is a simple and convenient way to enhance their intrinsic catalytic properties, which aroused tremendous interest.^{9–11} For example, Pei *et al.*¹² used TiO_2 , which possesses low conductivity and poor reactivity, to fabricate reductive TiO_{2-x} nanocrystals by self-doping with abundant OVs and achieve excellent catalytic activity in the oxygen reduction reaction (ORR). Sun *et al.*⁶ reported that CeO_2 with

rich OVs exhibited excellent conductivity and adsorption capacity, leading to satisfactory catalytic performance in the electrochemical reduction reaction.

It has been reported that the created OVs could play a significant role in catalysis. For instance, OVs serve as active sites for chemically absorbing the reactants, improving the number of reactants. Also, OVs could capture surrounding electrons to interact with intermediates with strong affinity and transfer electrons to them, boosting the redox reaction rates. Besides, the introduction of OVs changes the chemical environment and electronic structure of nano-metal oxides, modulating the band structure.¹³ However, the detailed interface catalytic mechanisms between the oxygen-defective nano-metal oxides and reactants induced by OVs were still unclear and insufficient. Therefore, taking advantage of OVs for interface catalytic regulation and revealing their roles in catalytic reactions are necessary and important.

What's more, it has been reported that OVs are beneficial for the generation of some reactive oxygen species (ROS), and the participation of these ROS in the catalytic process remarkably improved the catalytic efficiency.¹⁴ For example, Lyu *et al.* found that O_2 could be directly activated into singlet oxygen (${}^1\text{O}_2$) or superoxide radicals ($\cdot\text{O}_2^-$) spontaneously on TiO_{2-x} , exhibiting excellent performance in dye degradation, ORR, and cell viability.¹⁵ Yuan *et al.*¹⁶ presented BiO_{2-x} nanosheets rich in OVs to modulate ROS, catalyzing the production of $\cdot\text{O}_2^-$ and $\cdot\text{OH}$ in

^aKey Laboratory of Environmental Optics and Technology, Environmental Materials and Pollution Control Laboratory, Institute of Solid State Physics, HFSIPS, Chinese Academy of Sciences, Hefei 230031, P. R. China. E-mail: myang@iim.ac.cn; xingjiuhuang@iim.ac.cn

^bDepartment of Materials Science and Engineering, University of Science and Technology of China, Hefei 230026, P. R. China

^cCollege of Mechanical and Automotive Engineering, Anhui Polytechnic University, Wuhu, Anhui 241000, PR China

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cancer cells. Also, it was found that OVs are well suited to ROS catalytic kinetics. Li *et al.*¹⁷ found that the fabricated OVs on the surface of BiOCl are conducive to the formation of $\cdot\text{O}_2^-$, achieving complete visible light NO oxidation with excellent efficiency. However, the research on ROS induced by OVs participating in catalytic reactions is mainly focused on biocatalysis, photocatalysis, *etc.* Relevant studies on electrocatalysts are few, although there is a great possibility that ROS would be produced and exist during the process.¹⁸

As a p-type semiconductor with a wide band gap (about 5 eV), the low catalytic activity of ZrO_2 in the electrochemical process has been reported.^{19,20} Also, it can be used as a raw material for ceramic glaze because of its strong chemical inertness, improving the chemical stability, and acid and alkali resistance of glaze.^{21,22} Therefore, fabricating OVs in ZrO_2 also has a great possibility for improving its intrinsic properties, broadening its application as a catalyst. Besides, ZrO_2 is a kind of transition metal oxide with variable valence states of metal elements, which is quite similar to the above-mentioned TiO_2 and CeO_2 . However, the introduction of OVs induces the generation of Ti^{3+} and Ce^{3+} in TiO_2 and CeO_2 , respectively, participating in the catalytic reaction; while not only Zr^{3+} but also Zr^{2+} would be probably produced in oxygen-defective ZrO_2 , which is worth exploring. What's more, the possible generated ROS should also be studied during the electrochemical catalytic experiments.

In this work, ZrO_2 nanomaterials with mesoporous hollow sphere structures were fabricated, and different oxygen vacancy concentrations in the pristine ZrO_2 were modulated through calcining them in an oxidizing or reducing gas environment at high temperature, denoted as V_o -poor ZrO_2 and V_o -rich ZrO_2 . The difference in oxygen vacancy concentrations among these ZrO_2 nanomaterials was confirmed by electron paramagnetic resonance (EPR), X-ray photoelectron spectroscopy (XPS), and extended X-ray absorption fine structure (EXAFS). Besides, the effect of electron rearrangement induced by the abundant OVs is also verified to promote the catalytic reaction. To investigate the relationship between the catalytic ability of ZrO_2 materials and their oxygen vacancy concentrations, electrochemical reduction experiments were implemented. Surprisingly, the ZrO_2 possessing the highest concentration of OVs (V_o -rich ZrO_2) exhibited superior catalytic capacity. During electrochemical catalytic reduction of $\text{Pb}(\text{II})$, the current density of 1.0 μM $\text{Pb}(\text{II})$ signals with V_o -rich ZrO_2 is nearly 10 times higher than that of ZrO_2 with the lowest oxygen vacancy concentrations (V_o -poor ZrO_2). At the same time, *in situ* EPR spectra and DFT calculations were employed to prove the generation of $\cdot\text{OH}$ triggered by OVs and trace divalent heavy metal ions, such as $\text{Pb}(\text{II})$, boosting the catalytic efficiency. Moreover, the multiple roles of OVs in interface catalytic regulation for enhanced catalytic capacity of V_o -rich ZrO_2 were also explored.

2 Results and discussion

2.1 Structural and surface characterization

For obtaining ZrO_2 samples with different oxygen vacancy concentrations, pristine ZrO_2 , possessing a small quantity of intrinsic OVs, was first synthesized through the hydrothermal

method. Then, V_o -poor ZrO_2 and V_o -rich ZrO_2 were acquired through calcining pristine ZrO_2 in air or Ar/H_2 , respectively. As pristine ZrO_2 was partially reduced with the treatment of Ar/H_2 , a decreased number of oxygen atoms in V_o -rich ZrO_2 were obtained. In contrast, the oxidizing gas (air) caused pristine ZrO_2 to gain oxygen atoms, leading to the formation of V_o -poor ZrO_2 . The calcination treatment in reducing gas is also applicable for many other nano-metal oxides to create OVs,^{23,24} which is quite convenient and effective. The corresponding morphological characterization results of V_o -poor ZrO_2 , pristine ZrO_2 , and V_o -rich ZrO_2 are shown in Fig. 1a–c and S1, S2†. The transmission electron microscopy (TEM) images of pristine ZrO_2 and V_o -rich ZrO_2 showed distinct hollow sphere structures with a diameter of about 150 nm, and the hollow volumes of V_o -rich ZrO_2 particles significantly increased compared to those in pristine ZrO_2 particles. However, the morphology of V_o -poor ZrO_2 changed obviously, and severe aggregation was observed. The lattice orientations of (101) and (−111) in V_o -rich ZrO_2 particles were marked in high-resolution transmission electron microscopy (HRTEM) images (Fig. 1d–f). It is clear to see that the lattice fringes of V_o -poor ZrO_2 in Fig. 1d were clearer and neater than that of the others as the OVs were occupied by the oxygen atoms when being calcined in air. The energy-dispersive spectrometer (EDS) images in Fig. S2† show that the order of the percentage of oxygen atoms was V_o -poor ZrO_2 > pristine ZrO_2 > V_o -rich ZrO_2 , which was consistent with their corresponding oxygen vacancy concentrations. N_2 adsorption–desorption isotherm curves in Fig. S3† exhibited the types of hysteresis loops in the range of 0.3–1.0 (P/P_0), showing that all of them possessed mesoporous structures, and V_o -rich ZrO_2 had the largest surface areas compared to the others. The X-ray diffraction (XRD) spectra of these ZrO_2 particles are displayed in Fig. 1g. The peaks in V_o -rich ZrO_2 were in good agreement with the standard card of JCPDS NO. 80-2155 (tetragonal phase), while the monoclinic ZrO_2 characteristic peaks (JCPDS NO. 86-1449) were coincident with peaks in V_o -poor ZrO_2 which only exhibited a small proportion of tetragonal phase characteristics. Besides, the pristine ZrO_2 particles were also dominated by the tetragonal phase, showing a small intensity of characteristic peaks of monoclinic ZrO_2 .²⁵ A strong intensity of monoclinic ZrO_2 characteristic peak was observed in V_o -poor ZrO_2 as pure stoichiometric ZrO_2 crystals are the monoclinic phase which is highly stable at room temperature, and it would be converted to the tetragonal phase above 1400 K while to the cubic phase above 2600 K. It was found that the tetragonal phase could exist at normal temperature by introducing defects, and V_o -poor ZrO_2 possesses more monoclinic phases due to the decrease of the oxygen vacancy concentrations.²⁶ More details of crystalline phase transition were revealed by Raman spectra (Fig. S4†). The comparison of EPR signals provided evidence of different oxygen vacancy concentrations among V_o -poor ZrO_2 , pristine ZrO_2 , and V_o -rich ZrO_2 , as shown in Fig. 1h. The signals generated by the unpaired electrons at the OVs ($g = 2.00$) weakened sharply after heat treatment in air while strengthened in Ar/H_2 , which verified that V_o -rich ZrO_2 possessed the highest oxygen vacancy concentration while V_o -poor ZrO_2 had the lowest.^{27,28} The Fourier transform infrared (FT-IR) spectra



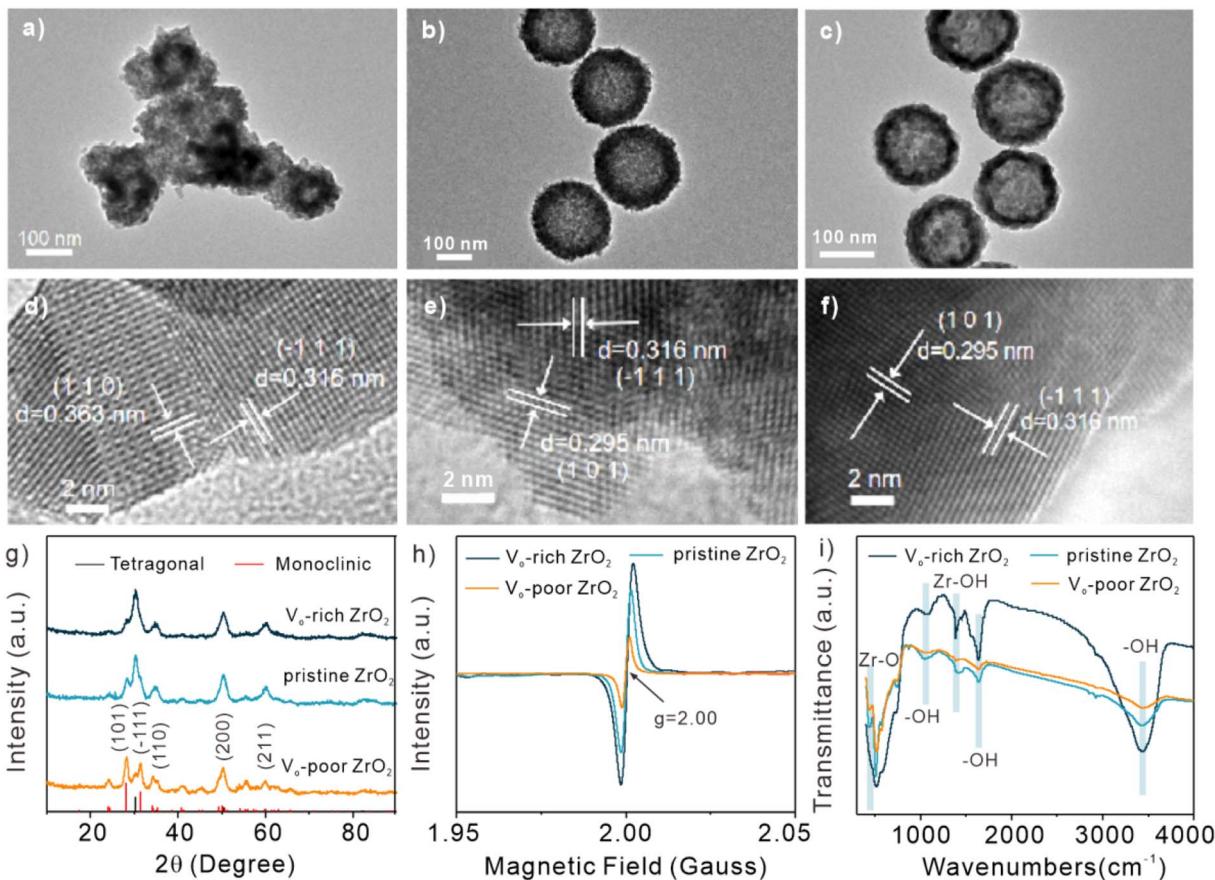


Fig. 1 (a–c) TEM and (d–f) HRTEM images of V_o -poor ZrO_2 , pristine ZrO_2 , and V_o -rich ZrO_2 , respectively. The specific details of the (101), (−111), and (110) planes are marked in the images. Comparison of (g) XRD spectra, (h) EPR spectra, and (i) FT-IR spectra among V_o -poor ZrO_2 , pristine ZrO_2 , and V_o -rich ZrO_2 .

(Fig. 1i) in the range of 400 to 4000 cm^{-1} demonstrated that they all have five characteristic peaks and there is no significant difference in peak position. The small peaks at 475, 1080, and 1385 cm^{-1} could be attributed to Zr–O vibration, bending vibration of hydroxyl groups, and Zr–OH vibration, respectively. The peaks at 1600 and 3430 cm^{-1} were assigned to the –OH flexural vibration mode and tensile mode.²⁹ It was clearly shown that the content of –OH is also positively correlated with the concentration of OVs and the –OH content in V_o -rich ZrO_2 was much higher than that in pristine ZrO_2 and V_o -poor ZrO_2 .

2.2 Valence states and electronic structure study

For figuring out the valence states of Zr in the three samples with different concentrations of OVs, the normalized Zr K-edge X-ray absorption near edge structure (XANES) spectra of Zr foil, $ZrCl_4$, pristine ZrO_2 , V_o -poor ZrO_2 , and V_o -rich ZrO_2 were analyzed, as shown in Fig. 2a. It is clear to see that the position of pre-edge peak tendency and the white line peak of Zr K-edge XANES in V_o -rich ZrO_2 was in lower energy compared with that in pristine ZrO_2 and V_o -poor ZrO_2 , indicating that the valence states of Zr in V_o -rich ZrO_2 were lower than those of the others, and all of them were lower than that in $ZrCl_4$ (Zr^{4+}). Besides, Fig. 2b demonstrated the fitting results of Zr K-edge EXAFS spectra of V_o -rich

ZrO_2 , which showed that V_o -rich ZrO_2 had Zr–O and Zr–Zr coordination peaks. More information is shown in Fig. S5, S6, and Table S1.† It was found that after calcining in Ar/H₂, the coordination numbers of Zr–O in V_o -rich ZrO_2 (3.1) decreased apparently compared to that in pristine ZrO_2 (3.6) and V_o -poor ZrO_2 (3.9), further proving the highest oxygen vacancy concentration in V_o -rich ZrO_2 . There are three kinds of charge-characteristic OVs possibly generated in hypoxic ZrO_2 due to Madelung potential, which are neutral, single, and double charge, respectively.²⁶ The corresponding schematic diagrams are shown in Fig. 2c. The oxygen atom escaped with two electrons remaining, resulting in the formation of neutral OVs; while single-charged OVs retain one electron, and the other one is transferred to the adjacent Zr^{4+} to generate Zr^{3+} . Regarding the double-charged OVs, Zr^{4+} is reduced to Zr^{3+} or Zr^{2+} depending on the number of electrons transferred.²⁵ The high-resolution X-ray photoelectron spectroscopy (XPS) spectra of Zr 3d (Fig. 2d) and O 1s (Fig. 2e) of the three ZrO_2 samples were also analysed for exploring their different OV species. As shown in Fig. 2d, the peak at 182.1 eV was in the majority of Zr 3d in pristine ZrO_2 (56.14%) and V_o -poor ZrO_2 (61.04%), indicating that the Zr^{4+} was the main state present in these two ZrO_2 samples. Besides, the peaks coloured with cyan and orange were the non-stoichiometric ZrO_2 caused by OVs and assigned as ZrO_{x1} ($1.5 < x_1 < 2$) and ZrO_{x2}



($1 < x_2 \leq 1.5$), respectively. In Table S2,[†] we summarized the proportions of ZrO_{x1} , ZrO_{x2} , and their sums in the total area of O 1s, as well as that in the total area of Zr 3d. It was obvious that the sum proportion of ZrO_{x1} and ZrO_{x2} in V_o -rich ZrO_2 was much higher than that in pristine ZrO_2 and V_o -poor ZrO_2 , indicating that more single and double-charged OVs were produced in V_o -rich ZrO_2 . Similarly, ZrO_{x1} and ZrO_{x2} can be allocated to the position of 531.2 and 532.2 eV in the O 1s region in pristine ZrO_2 and V_o -poor ZrO_2 ,^{25,30} as shown in Fig. 2e. It is worth noting that the peaks of ZrO_{x1} and ZrO_{x2} in V_o -rich ZrO_2 were much obvious than that of the other two, which all verified the highest concentration of Zr^{2+} and Zr^{3+} in V_o -rich ZrO_2 . And the peak at 529.9 eV in pristine ZrO_2 , which was attributed to Zr^{4+} , shifted toward higher binding energy (530.2 eV) after fabricating

abundant OVs while shifted toward lower binding energy (529.6 eV) in V_o -poor ZrO_2 , indicating that more dramatic changes in the oxygen environment occurred.

To investigate the different electrochemical properties of these samples caused by OVs, the cyclic voltammetry (CV) and electrochemical impedance spectroscopy (EIS) experiments on pristine ZrO_2 , V_o -poor ZrO_2 , and V_o -rich ZrO_2 were implemented. It was found that the conductivities of ZrO_2 samples were positively correlated with the concentration of OVs, as exhibited in Fig. S7.[†] Besides, the UV-vis absorption spectra of the three ZrO_2 particles (Fig. 2f) were collected and converted to a Tauc plot (Fig. 2g) as there is a close relationship between band gap and electrical conductivity. The energy band gap was

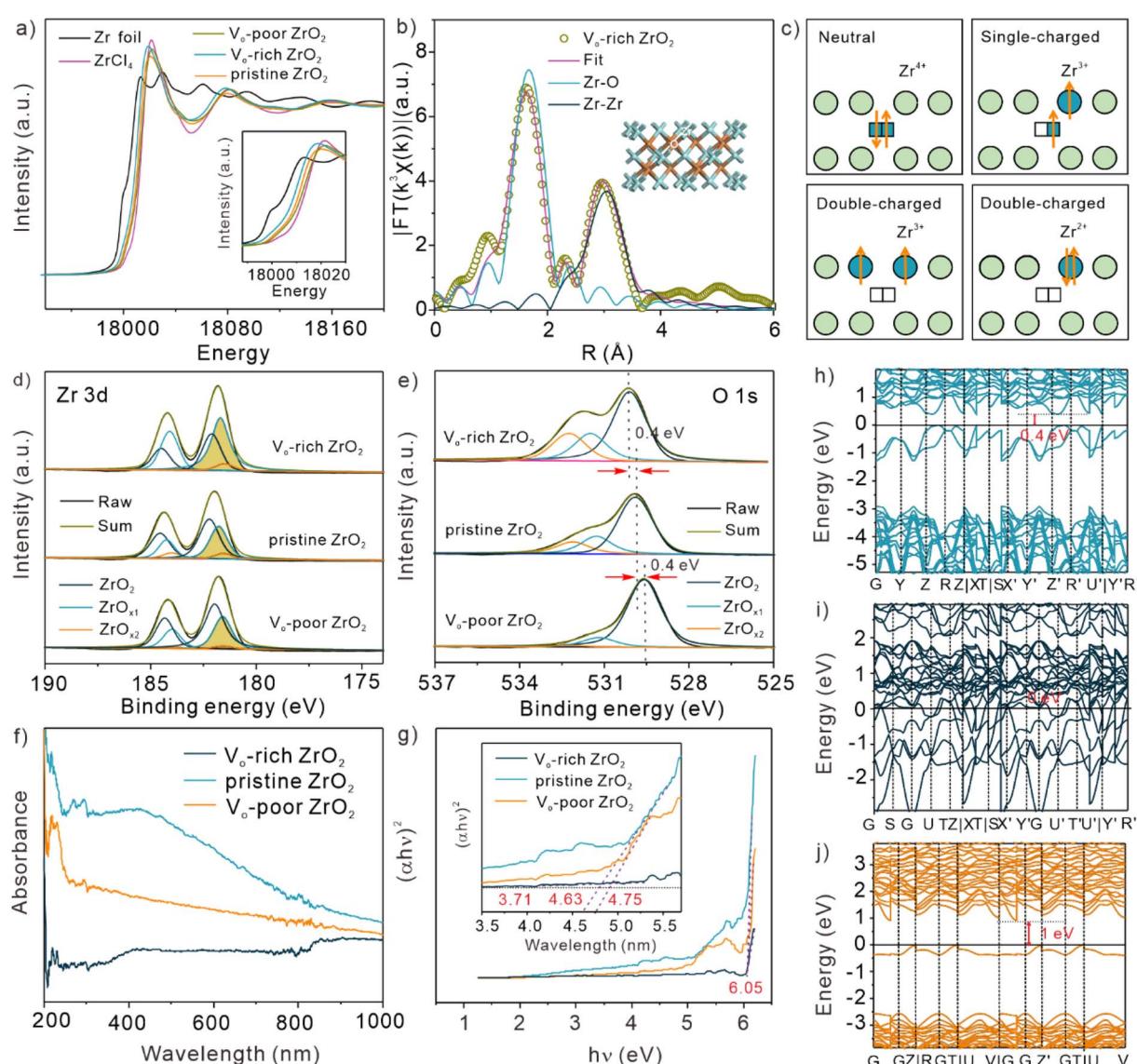


Fig. 2 (a) Normalized Zr K-edge XANES spectra of Zr foil, ZrCl_4 , pristine ZrO_2 , V_o -poor ZrO_2 , and V_o -rich ZrO_2 . (b) Zr K-edge EXAFS spectra (R space, k^3 -weighted) of V_o -rich ZrO_2 and corresponding fitting curves. The inset image of (b) is coordination structure diagrams of V_o -rich ZrO_2 . (c) Schematic diagram of four different forms of oxygen vacancies (electrons with oxygen vacancies indicated by up and down orange arrows). Comparison of high-resolution XPS spectra of (d) Zr 3d and (e) O 1s among pristine ZrO_2 , V_o -poor ZrO_2 , and V_o -rich ZrO_2 . (f) UV-vis DRS spectra, (g) Tauc plot obtained from the UV-vis data of pristine ZrO_2 , V_o -poor ZrO_2 , and V_o -rich ZrO_2 . Band gap diagrams of (h) pristine ZrO_2 , (i) V_o -rich ZrO_2 , and (j) V_o -poor ZrO_2 .

calculated by converting the data according to the following formula:

$$(\alpha h\nu)n = k(h\nu - E_g) \quad (1)$$

where α represents the absorption coefficient and $h\nu$ is the photoelectron energy, and the value of k depends on the material. The characteristic value of a transition in a semiconductor is represented by two fixed values, denoted as n . For the direct transition, the value is 2 and for the indirect transition it is 1/2.^{31,32} Among them, ZrO_2 belongs to the direct transition type and the intercept value was obtained by extending the straight part to the abscissa, which was the band gap energy (E_g) of the semiconductor. The main band gap energy of all three samples was 6.05 eV, while the new band gap energies introduced by OVs were 4.78, 4.92, and 3.71 eV for pristine ZrO_2 , V_o -poor ZrO_2 and V_o -rich ZrO_2 , respectively. This showed that it was OVs that caused a sharp change in the energy band gap of the material which enhanced the transfer of electrons.^{27,33,34} DFT calculations were also carried out to uncover the enhanced conductivity. After optimizing the configurations of ZrO_2 samples based on their different crystalline phases (Fig. S8†), a comparison of the total density of states (TDOS) and the corresponding band gap diagrams of pristine ZrO_2 , V_o -rich ZrO_2 , V_o -poor ZrO_2 , and ZrO_2 with no OVs showed that the band gaps reduced obviously due to abundant OVs existing in V_o -rich ZrO_2 , which were 0.4, 0, 1, and 3.6 eV, respectively, as shown in Fig. 2h–j, S9, and S10.† Therefore, a conclusion could be drawn that abundant OVs are key factors in the enhanced conductivity and electrochemical activity of V_o -rich ZrO_2 .

2.3 Remarkable catalytic ability of V_o -rich ZrO_2 triggered by OVs

To explore the relationship between OV concentration and catalytic capacity, a divalent lead solution was adsorbed on the three ZrO_2 samples, revealing the interface catalytic effect between V_o -rich ZrO_2 and lead ions. XAFS spectra of the three samples after adsorption of $\text{Pb}(\text{II})$ were first studied, which were denoted as pristine ZrO_2/Pb , V_o -poor ZrO_2/Pb , and V_o -rich ZrO_2/Pb , respectively. Fig. 3a shows the normalized Pb L_3 -edge XANES spectra of Pb foil, PbO , pristine ZrO_2/Pb , V_o -poor ZrO_2/Pb , and V_o -rich ZrO_2/Pb . It has been reported that the height of the white line peak is related to the electronic structure and usually a higher white line peak indicates a higher valence state. Surprisingly, the height of the white line peak of Pb in V_o -rich ZrO_2/Pb was similar to that in Pb foil, while that of pristine ZrO_2/Pb and V_o -poor ZrO_2/Pb was between Pb foil and PbO , indicating that almost all $\text{Pb}(\text{II})$ was catalytically reduced to $\text{Pb}(0)$ in V_o -rich ZrO_2 while part of $\text{Pb}(\text{II})$ in pristine ZrO_2 and V_o -poor ZrO_2 . It is worth mentioning that $\text{Pb}(\text{II})$ could be effectively reduced without an electric field, showing that V_o -rich ZrO_2 had superb catalytic ability. Comparison of Pb 4f XPS spectra of V_o -rich $\text{ZrO}_2/\text{Pb}(\text{II})$ and $\text{Pb}(\text{NO}_3)_2$ is shown in Fig. 3b. It was observed that the two peaks of Pb $4f_{7/2}$ and Pb $4f_{5/2}$ at 138.7 eV and 143.3 eV in V_o -rich ZrO_2 , respectively, were a little lower than those in $\text{Pb}(\text{NO}_3)_2$ (139.1 eV and 144.4 eV),³⁵ indicating that $\text{Pb}(\text{II})$ possibly obtained some electrons from V_o -rich ZrO_2 ,

which was in good agreement with the above XANES results. Besides, a comparison of XPS on different samples before and after the adsorption of $\text{Pb}(\text{II})$ was also performed. It is clear to see that the peak intensity of Pb 4f followed the trend as V_o -rich ZrO_2 > pristine ZrO_2 > V_o -poor ZrO_2 , as shown in Fig. S11,† indicating that V_o -rich ZrO_2 had the largest amount of the adsorbed lead. At the same time, the surface energies of (101) and (−111) with different OVs concentrations on V_o -rich ZrO_2 have been calculated and the detailed information is shown in Table S2,† which indicated that with the increased concentration of OVs, the surface energies of both facets increase, showing better adsorption abilities.³⁶ As there was a positive correlation between OV concentration and adsorption capacity, it could be deduced that the produced OVs could increase the number of active sites on the surface of ZrO_2 to adsorb more $\text{Pb}(\text{II})$. Besides, the zeta potential test results in HAc-NaAc buffer solution at pH 5.0 are displayed in Fig. S12.† Clearly, as the concentration of OVs increased, more free electrons covered the surface of ZrO_2 , resulting in a stronger electronegativity. Combined with the FT-IR results in Fig. 1i, perhaps more $\text{Pb}(\text{II})$ will be electrostatically adsorbed on V_o -rich ZrO_2 as $-\text{OH}$ could chemically bond and interact with $\text{Pb}(\text{II})$, resulting in better adsorption capacity toward $\text{Pb}(\text{II})$.^{29,37} In addition, the changes of XPS Zr 3d and O 1s in V_o -rich ZrO_2 were also investigated, as displayed in Fig. 3c, d and S13, S14.† Through the comparison, it was found that the total ratio of $(\text{ZrO}_{x1} + \text{ZrO}_{x2})$ decreased significantly after the adsorption of $\text{Pb}(\text{II})$ compared with that of pristine $\text{ZrO}_2/\text{Pb}(\text{II})$ and V_o -poor $\text{ZrO}_2/\text{Pb}(\text{II})$, as shown in Table S3,† which indicated that Zr^{3+} and Zr^{2+} directly participated in boosting the catalytic reduction of $\text{Pb}(\text{II})$. For investigating the catalytic mechanisms of V_o -rich ZrO_2 toward $\text{Pb}(\text{II})$ at the atomic level, the EXAFS spectra of Pb L_3 -edge in R space of pristine ZrO_2/Pb , V_o -poor ZrO_2/Pb , and V_o -rich ZrO_2/Pb are exhibited in Fig. 3e, f and S15.† More detailed fitting results are shown in Table S4.† It was found that the first-shell coordination peak of V_o -rich ZrO_2/Pb was at $R_{\text{Pb}-\text{O}} = 2.30 \text{ \AA}$ with 1.2 O atoms around,³⁸ where the second-shell coordination peak of V_o -rich ZrO_2/Pb was at $R_{\text{Pb}-\text{Zr}} = 3.04 \text{ \AA}$ with 1.0 Zr atom around. Compared with pristine ZrO_2/Pb and V_o -poor ZrO_2/Pb , the bond length of Pb-Zr in V_o -rich ZrO_2/Pb was much longer than that of them, which was 3.01 \AA and 3.00 \AA , respectively. It suggested that Pb ions adsorbed on V_o -rich ZrO_2 were easier to desorb than that adsorbed on pristine ZrO_2 and V_o -poor ZrO_2 , contributing to improving the interface catalytic efficiency.

2.4 The observed hydroxyl radicals induced by divalent heavy metal ions

The *in situ* EPR experiments were carried out to study the possibly produced ROS species during the electrochemical catalytic process with V_o -rich ZrO_2 . Fig. 4a and b display the EPR spectra of observed $\cdot\text{OH}$ under the specific experimental conditions, and 5,5-dimethyl-1-oxaporphyrin-N-oxide (DMPO) was adopted as a spin-trapping agent. Obviously, it is clear to see that no matter whether $\text{Pb}(\text{II})$ and DMPO are added or not, there are no peak signals when without the effect of negative potential and the catalyst of V_o -rich ZrO_2 . After applying the negative potential, no



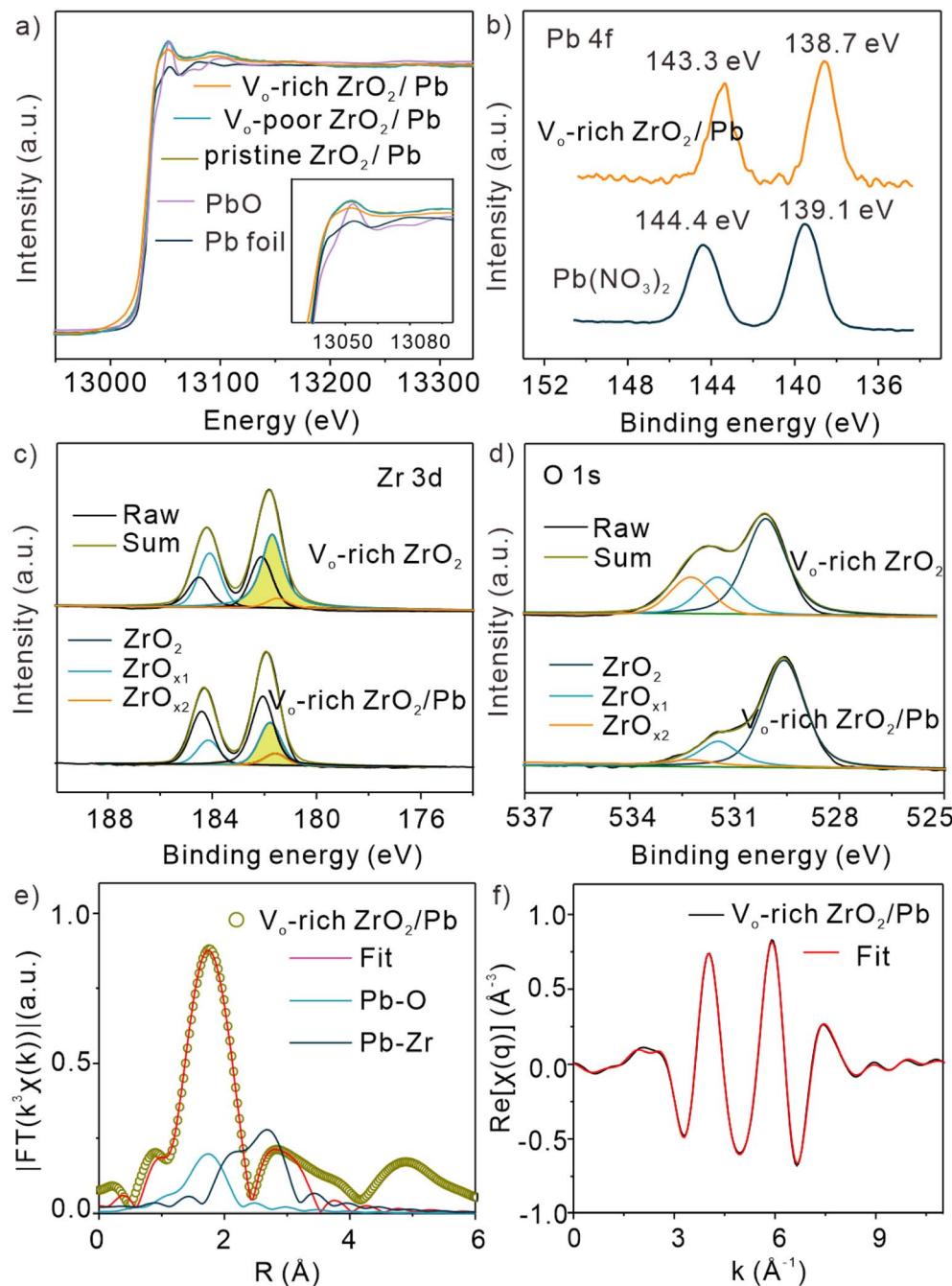


Fig. 3 (a) Normalized Pb L_3 -edge XANES spectra of Pb foil, PbO , pristine ZrO_2/Pb , $\text{V}_{\text{o}}\text{-poor } \text{ZrO}_2/\text{Pb}$, and $\text{V}_{\text{o}}\text{-rich } \text{ZrO}_2/\text{Pb}$. (b) Comparison of high-resolution Pb 4f XPS between $\text{V}_{\text{o}}\text{-rich } \text{ZrO}_2/\text{Pb}$ and $\text{Pb}(\text{NO}_3)_2$. (c) High-resolution XPS of Zr 3d , (d) O 1s results of $\text{V}_{\text{o}}\text{-rich } \text{ZrO}_2$ and $\text{V}_{\text{o}}\text{-rich } \text{ZrO}_2/\text{Pb}$. (e) Pb L_3 -edge EXAFS spectra (R space, k^3 -weighted) of $\text{V}_{\text{o}}\text{-rich } \text{ZrO}_2/\text{Pb}$ and corresponding fitting curves. (f) The corresponding $\text{k}^3(\text{k})$ oscillations of $\text{V}_{\text{o}}\text{-rich } \text{ZrO}_2/\text{Pb}$.

EPR signals could be detected when there is no $\text{V}_{\text{o}}\text{-rich } \text{ZrO}_2$ existing even though $\text{Pb}(\text{II})$ and DMPO were added. As shown in Fig. 4b, with the effect of negative potential and $\text{V}_{\text{o}}\text{-rich } \text{ZrO}_2$, several peaks appeared in the presence of $\text{Pb}(\text{II})$ and DMPO, while there were no peaks observed without the addition of $\text{Pb}(\text{II})$. Interestingly, with the increasing concentration of $\text{Pb}(\text{II})$, the typical peaks of DMPO-·OH (1:2:2:1) became more evident, indicating that a growing number of ·OH was produced during this electrochemical catalytic process. It is worth mentioning that

2,2,6,6-tetramethyl-4-piperidine (TEMP), another type of spin-trapping agent, was also solely added for excluding the possibility of the generation of other ROS species. In brief, it could be deduced that ·OH would be produced during the electrochemical catalytic reaction of $\text{V}_{\text{o}}\text{-rich } \text{ZrO}_2$, where $\text{Pb}(\text{II})$ is necessary and act as a radical initiator.

For figuring out whether other kinds of divalent heavy metal ions, except $\text{Pb}(\text{II})$, could trigger the production of ·OH, 1.6 μM $\text{Cd}(\text{II})$ and $\text{Hg}(\text{II})$ were added respectively into the buffer solution

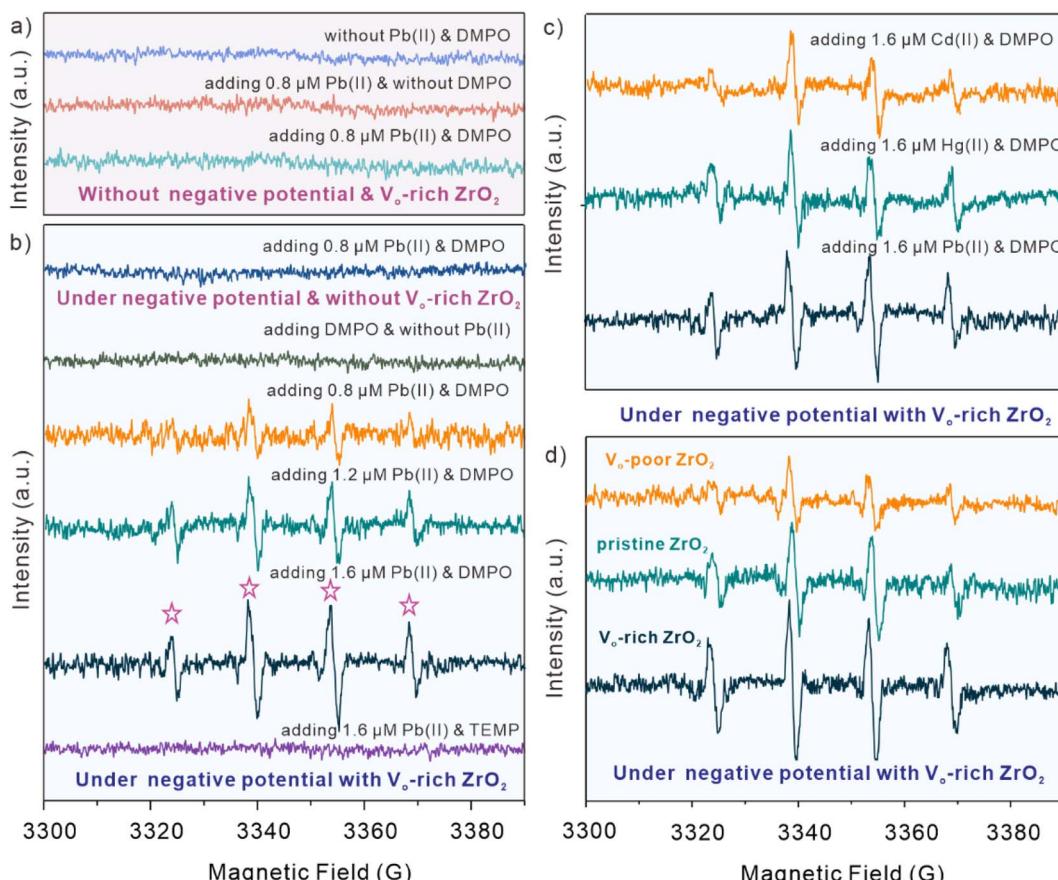


Fig. 4 (a) The obtained EPR spectra without negative potential and V_o -rich ZrO_2 . (b) The obtained EPR spectra under negative potential. (c) Comparison of EPR spectra of V_o -rich ZrO_2 in the presence of DMPO and 1.6 μM different kinds of divalent heavy metal ions, including $Cd(II)$, $Hg(II)$, and $Pb(II)$. (d) Comparison of EPR spectra of V_o -rich ZrO_2 , pristine ZrO_2 , and V_o -poor ZrO_2 in the presence of DMPO and 2.0 μM $Pb(II)$.

under the same experimental conditions. Surprisingly, as displayed in Fig. 4c, the peaks of DMPO-·OH were also detected when adding $Cd(II)$ and $Hg(II)$, while their intensities seemed a bit lower than that of adding $Pb(II)$. This indicated that divalent heavy metal ions, not just confined to $Pb(II)$, could induce the generation of ·OH with the electrochemical catalytic effect of V_o -rich ZrO_2 . What's more, pristine ZrO_2 and V_o -poor ZrO_2 were also used to investigate the relationship between OV concentrations and the generated ·OH. Fig. 4c shows the comparison of EPR spectra of V_o -rich ZrO_2 , pristine ZrO_2 , and V_o -poor ZrO_2 for electrochemical catalysis in the presence of DMPO and 2.0 μM $Pb(II)$. It was found that V_o -rich ZrO_2 had the largest amount of produced ·OH during the reaction, and the intensities of DMPO-·OH peak signals were positively correlated with the concentration of OVs. It could be concluded that the captured different intensities of DMPO-·OH peak signals of these three samples were ascribed to their different concentration of OVs. And the generated ·OH perhaps have a specific impact on boosting the catalytic capacities of V_o -rich ZrO_2 .

2.5 The investigation of changed electronic structure and energy barrier

DFT calculations were employed to further understand the inner mechanisms for obviously enhanced catalytic capacity of

V_o -rich ZrO_2 due to the generated abundant OVs. The electronic structure of V_o -rich ZrO_2 after adsorption of $Pb(II)$ was studied first. The optimized structures of $Pb(II)$ adsorbed on the surface of the three ZrO_2 samples are exhibited in Fig. S16.† It showed that the adsorption energies of $Pb(II)$ on pristine ZrO_2 , V_o -rich ZrO_2 , and V_o -poor ZrO_2 were -1.542 , -0.196 and -1.695 eV. Apparently, V_o -rich ZrO_2 /Pb had the lowest adsorption energy. The changes in the partial DOS of Zr, O, and Pb atoms in V_o -rich ZrO_2 before and after the adsorption are shown in Fig. 5a. The degree of overlap between the orbitals of Pb 6p and Zr 4d near the Fermi level followed the trend as V_o -rich ZrO_2 /Pb < pristine ZrO_2 /Pb < V_o -poor ZrO_2 /Pb (Fig. S17†), reflecting the opposite strength on desorption capacities. This evidence was in good agreement with the EXAFS results. Besides, it is clear to see that the electron density of Zr 3d and O 1s close to the Fermi surface increased a lot in V_o -rich ZrO_2 /Pb, indicating that the intrinsic material becomes active when interacting with Pb atoms. A prominent newborn orbital hybrid peak (filled with purple) emerged in the partial DOS of O 1s of all three absorptive configurations, which proved the speculation that the adsorbed Pb atoms might take up the positions of OVs. Besides, the charge density difference images of the three adsorption structures showed that the electron cloud density (yellow area) around the Pb atoms in V_o -rich ZrO_2 /Pb was the least while that

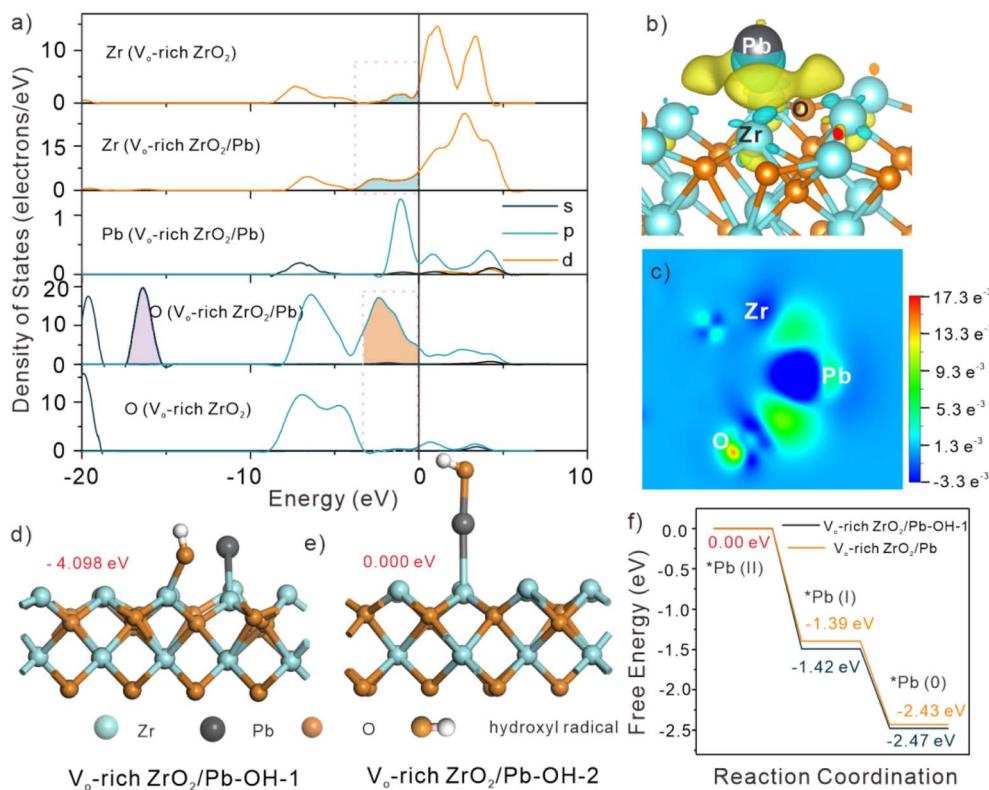


Fig. 5 (a) Comparison of DOS between V_o-rich ZrO₂/Pb and V_o-rich ZrO₂. (b) Charge density difference images of V_o-rich ZrO₂/Pb. The blue areas represent electron depletion, and the yellow areas represent electron accumulation. (c) Charge density difference image on the slice through Zr, O, and Pb atoms. The navy-blue areas represent electron depletion, and the green areas represent electron accumulation. (d and e) The two optimized configurations of V_o-rich ZrO₂/Pb when 'OH are participating in Pb(II) reduction. (f) Comparison of the free energy of reaction intermediates during the reduction process of Pb(II) on V_o-rich ZrO₂/Pb-OH-1 and V_o-rich ZrO₂/Pb.

in V_o-rich ZrO₂/Pb was the largest, which further proved the low absorptive ability of Pb(II) on V_o-rich ZrO₂, as shown in Fig. 5b and S18.† The charge density difference image on the slice through Pb, Zr, and O atoms (Fig. 5h) displayed the electron flow condition that Zr atoms lost electrons to Pb atoms.³⁹ In other words, the electron rearrangement in V_o-rich ZrO₂ caused by OVs is beneficial for the catalytic reduction of Pb(II). The Bader charge of Pb, Zr, and O atoms was computed, and the results are displayed in Fig. S19 and Table S5.† It was found that the adsorbed Pb could obtain 0.167e more in V_o-rich ZrO₂ than that in V_o-poor ZrO₂, which was conducive to boosting the reduction rates, showing the excellent catalytic ability of V_o-rich ZrO₂. The function of 'OH in catalysis of Pb(II) was also studied. According to different active sites of 'OH toward Pb(II), two different optimized configurations of V_o-rich ZrO₂/Pb when 'OH participate in Pb(II) reduction are exhibited in Fig. 5d and e, which were denoted as V_o-rich ZrO₂/Pb-OH-1 and V_o-rich ZrO₂/Pb-OH-2. Considering that ZrO₂/Pb-OH-1 had the larger absorptive energy, -4.098 eV, which had a more stable configuration, the energies for Pb(II) reduction on V_o-rich ZrO₂/Pb-OH-1 were simulated. The reduction process of Pb(II) to Pb(0) was in two steps, as shown in Fig. 5f. Apparently, every step of V_o-rich ZrO₂/Pb-OH-1 for Pb(II) reduction had lower energies than that of V_o-rich ZrO₂/Pb, indicating that the participation of 'OH was more conducive to the reaction.

2.6 Super-high catalytic capacity of V_o-rich ZrO₂ in electrochemical reduction

For verifying the catalytic capacity and practical application prospect of V_o-rich ZrO₂ in the electrochemical reduction

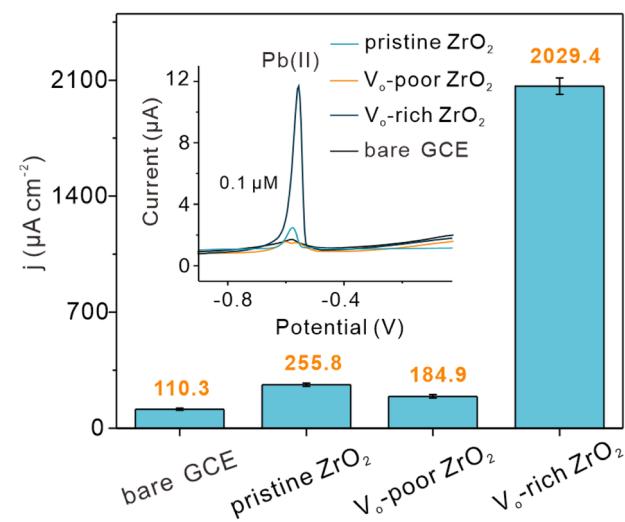


Fig. 6 Comparisons of the current densities and the SWASV curves of Pb(II) at the same concentration on the bare GCE and the three modified electrodes.

reaction, the three ZrO_2 samples were modified as catalysts on the same glassy carbon electrode (GCE) respectively for electrochemical catalysis of $\text{Pb}(\text{II})$ through square wave anode stripping voltammetry (SWASV). The bare GCE was also tested for comparison. The current responses of $\text{Pb}(\text{II})$ with increasing concentrations are shown in Fig. S20a-d.[†] Apparently, the current values of these modified GCEs toward $\text{Pb}(\text{II})$ at *ca.* -0.58 V all increased with the $\text{Pb}(\text{II})$ concentrations and an excellent linear relationship between $\text{Pb}(\text{II})$ concentration and peak current density was observed. The values of their current density per unit concentration were obtained by dividing the active area by the SWASV slope, and the active area of each modified electrode is shown in Fig. S21.[†] The current density per unit concentration of pristine ZrO_2 , V_o -poor ZrO_2 , and V_o -rich ZrO_2 modified electrodes toward $\text{Pb}(\text{II})$ was 255.8 , 184.9 , and 2029.4 $\mu\text{A cm}^{-2} \mu\text{M}^{-1}$, respectively, (Fig. 6). It was surprising that the catalytic ability of V_o -rich ZrO_2 was almost 6.9 times higher than that of pristine ZrO_2 and even 9.9 times higher than that of V_o -poor ZrO_2 , which indicated that the incredibly improved electrocatalytic ability of pristine ZrO_2 was obtained after creating OVs. Besides, the order of their catalytic ability (V_o -rich ZrO_2 $>$ pristine ZrO_2 $>$ V_o -poor ZrO_2) was consistent with their concentration of OVs, implying the vital role of OVs in the catalytic reduction of $\text{Pb}(\text{II})$. Besides, the corresponding experiments for characterizing catalytic cycle properties of V_o -rich ZrO_2 have also been carried out, showing excellent practical application prospects. More information is displayed in Fig. S22.[†]

Fig. 7 vividly depicts the interface catalytic regulation *via* significant roles of OVs in V_o -rich ZrO_2 for enhancing the catalytic efficiency in the electrochemical reduction of $\text{Pb}(\text{II})$.

Fig. 7a shows that the extra electrons remaining in OVs transferred to the adjacent Zr^{4+} , leading to the generation of Zr^{3+} and Zr^{2+} . Then, more $\text{Pb}(\text{II})$ is adsorbed on the surface OVs of V_o -rich ZrO_2 due to the large surface area, occupying the position of OVs and resulting in the formation of $\text{Pb}-\text{O}$ and $\text{Pb}-\text{Zr}$ bonds, as shown in Fig. 7b. The accumulated electrons in Zr^{3+} and Zr^{2+} directly transferred to $\text{Pb}(\text{II})$, promoting the catalytic reduction of absorbed $\text{Pb}(\text{II})$ to $\text{Pb}(0)$. Besides, the longest $\text{Pb}-\text{Zr}$ bonds and the lowest adsorption energy in V_o -rich ZrO_2/Pb compared to those in pristine ZrO_2/Pb and V_o -poor ZrO_2/Pb are beneficial for $\text{Pb}(\text{II})$ to desorb, improving the rates of conversion. What's more, the generated $\cdot\text{OH}$ which are induced by OVs and $\text{Pb}(\text{II})$ also act as redox mediators to accelerate the electron transfer rates in electrochemical reduction reactions, boosting the interface catalytic effect.

3 Conclusions

In summary, ZrO_2 mesoporous hollow spheres rich in OVs were fabricated and the multiple roles of OVs in interface catalytic regulation were revealed. First, the introduction of OVs brings more catalytic reaction sites. After fabricating abundant OVs, the adsorption capacity is improved due to increased surface area, and more surface $\cdot\text{OH}$ is produced for chemical interaction with reactants. Unlike other catalytic systems where OVs are indirectly involved in catalytic processes, in this study, OVs themselves are catalytic rooms for the adsorbed reactants, enhancing the rates of conversion and the reaction. Second, OVs cause the rearrangement of electrons. The excess electrons in OVs transfer to the adjacent Zr^{4+} , promoting the formation of Zr^{2+} and Zr^{3+} , which in turn transfer electrons to the reactants. Zr^{2+} and Zr^{3+} serve as redox mediators, accelerating the electron transfer rates and boosting the catalytic reduction reaction. The changed electronic and energy band structures induced by OVs also enhanced the intrinsic conductivity and activity of pristine ZrO_2 . Besides, DFT calculations and EXAFS revealed that longer $\text{Pb}-\text{Zr}$ and lower absorptive energies are beneficial for desorption. Third, for the first time, $\cdot\text{OH}$ have been observed to be triggered by OVs and divalent heavy metal ions in *in situ* EPR experiments. It was shown that when adding divalent heavy metal ions, more produced $\cdot\text{OH}$ were captured in the electrochemical reduction reaction with V_o -rich ZrO_2 , which also act as a catalyst. In the experiments of electrochemical reduction of $\text{Pb}(\text{II})$, it was shown that the catalytic ability of V_o -rich ZrO_2 was almost 9.9 times higher than that of V_o -poor ZrO_2 , showing superior practical application prospects in the fields of chemistry, nanomaterials, catalysis, *etc.* This finding will provide enlightenment and guidance for wide communities including advanced *in situ* measurements, especially in the fields of chemistry, catalysis, nanomaterials, *etc.*, in boosting the catalytic effect through interface catalytic regulation.

4 Experimental section

4.1 Synthesis of ZrO_2 hollow spheres

The synthesis of ZrO_2 hollow spheres referred to the previous literature with revision.⁴⁰ Zirconium dichloride (0.3 g) is added

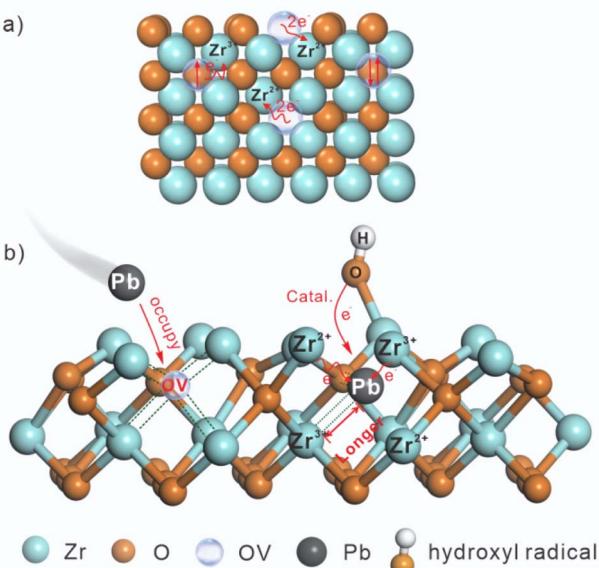


Fig. 7 The schematic diagram of (a) electron rearrangement on V_o -rich ZrO_2 triggered by OVs; (b) interface catalytic regulation *via* OVs and $\cdot\text{OH}$ on V_o -rich ZrO_2 .



to acetone (90 mL) and stirred magnetically for a few minutes to completely dissolve to form a colourless solution. The solution was immediately filled into a 100 mL Teflon-lined container, sealed, and placed in an oven at 200 °C for 12 h, and then naturally cooled down to room temperature. The resulting solution was centrifuged at 10 000 rpm for 3 minutes, then alternately washed twice with water, ethanol, and acetone. The obtained precipitate was freeze-dried all day, and the resulting powder was called pristine ZrO_2 .⁴⁰ The pristine ZrO_2 powder was calcined at 650 °C in a tube furnace with an average ramp rate of 3 °C min⁻¹ for 90 minutes, and the gas introduced was (Ar/H₂, air); the products were named V_o-rich ZrO_2 and V_o-poor ZrO_2 .

4.2 Computational methods

Spin-polarized DFT calculations were performed using the Vienna *ab initio* simulation package (VASP).^{41,42} The generalized gradient approximation proposed by Perdew, Burke, and Ernzerhof (GGA-PBE) is selected for the exchange-correlation potential.⁴³ The pseudo-potential was described by the projector-augmented-wave (PAW) method.⁴⁴ The geometry optimization is performed until the Hellmann–Feynman force on each atom is smaller than 0.02 eV·Å⁻¹. The energy criterion is set to 10⁻⁶ eV in the iterative solution of the Kohn–Sham equation. For DOS calculation, a 2 × 1 × 1 supercell adopted from tetragonal ZrO_2 was used for pristine ZrO_2 and V_o-rich ZrO_2 , and a 1 × 12 × 1 supercell adopted from monoclinic ZrO_2 was used for V_o-poor ZrO_2 . The supercell size for pristine ZrO_2 and V_o-rich ZrO_2 is 10.14 × 5.07 × 5.07 Å³, and it is 5.1454 × 10.4150 × 5.3107 Å³ for V_o-poor ZrO_2 . The k-mesh for structure optimization is 0.06 2π Å⁻¹. For adsorption calculation, a 2 × 2 supercell with 12 Å vacuum is applied. The size of V_o-rich ZrO_2 /Pb and pristine ZrO_2 is 10.14 × 10.14 × 16.8022 Å³. The size of V_o-poor ZrO_2 /Pb is 10.415 × 10.6214 × 17.0749 Å³. The k-mesh for structure optimization is 0.06 2π Å⁻¹. The hydroxyl radicals were calculated using PBE and their charge states are automatically assigned by the program.

Data availability

We have provided all the data in the manuscript and ESI.†

Author contributions

X. H., M. Y. and X. X. designed this project. X. X. and C. Z. implemented the experiments. Z. S. implemented the DFT calculations. X. X., Y. Z. and Z. G. completed the data analysis. X. X. and C. Z. drafted the manuscript. S. C., P. L. and Y. S. reviewed and edited the manuscript. All authors approved the final version.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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