

Cite this: *RSC Adv.*, 2018, 8, 2963

Improvement of O₂ adsorption for α -MnO₂ as an oxygen reduction catalyst by Zr⁴⁺ doping

Yicheng Wang,^a Yaozong Li,^a Zhenghang Lu^b and Wei Wang^{*a}

Zr⁴⁺ doped α -MnO₂ nanowires were successfully synthesized by a hydrothermal method. XRD, SEM, TEM and XPS analyses indicated that Mn³⁺ ions, Mn⁴⁺ ions, Mn^{4+ δ} ions and Zr⁴⁺ ions co-existed in the crystal structure of synthesized Zr⁴⁺ doped α -MnO₂ nanowires. Zr⁴⁺ ions occupied the positions originally belonging to elemental manganese in the crystal structure and resulted in a mutual action between Zr⁴⁺ ions and Mn³⁺ ions. The mutual action made Mn³⁺ ions tend to lose their electrons and Zr⁴⁺ ions tend to get electrons. Cathodic polarization analyses showed that the electrocatalytic activity of α -MnO₂ for oxygen reduction reaction (ORR) was remarkably improved by Zr⁴⁺ doping and the Zr/Mn molar ratio notably affected the ORR performance of the air electrodes prepared by Zr⁴⁺ doped α -MnO₂ nanowires. The highest ORR current density of the air electrodes prepared by Zr⁴⁺ doped α -MnO₂ nanowires in alkaline solution appeared at Zr/Mn molar ratio of 1 : 110, which was 23% higher than those prepared by α -MnO₂ nanowires. EIS analyses indicated that the adsorption process of O₂ molecules on the surface of the air electrodes prepared by Zr⁴⁺ doped α -MnO₂ nanowires was the rate-controlling step for ORR. The DFT calculations revealed that the mutual action between Zr⁴⁺ and Mn³⁺ in Zr⁴⁺ doped α -MnO₂ nanowires enhanced the adsorption process of O₂ molecules.

Received 10th September 2017

Accepted 23rd December 2017

DOI: 10.1039/c7ra10079e

rsc.li/rsc-advances

1. Introduction

Oxygen reduction reaction (ORR), as the main kinetically limiting cathode reaction in clean energy related technologies, is the primary cause of voltage loss in fuel cells and metal–air batteries.^{1,2} Developing catalysts for ORR is of significant importance. At present, the most extensively used ORR catalysts are still Pt-based materials due to their relatively high activity and stability.^{3,4} However, the low abundance of Pt in the earth's crust together with its high price limits its wide and practical applications in fuel cells. In recent years, various efforts have been made to explore non-Pt-based catalysts for ORR, such as Ag-based catalysts,^{5,6} Pd-based materials,^{7,8} N-doped carbon,^{9–11} M–N₄-macrocycles,¹² Fe–N–C hybrids^{13–15} as well as some transition metal oxides.^{16–19} Nevertheless, exploiting inexpensive and highly active electrocatalysts for ORR remains a great challenge.^{20,21}

Manganese dioxide (MnO₂) has been utilized as an electrocatalyst in metal–air batteries owing to its unique electrochemical performance, low cost and environmentally friendly properties.^{22–26} MnO₂ is a polymorphic material with three different crystal phases including α , β , and γ phases.²⁷ The catalytic activity of MnO₂ depends strongly on the

crystallographic structure, following an order of $\alpha > \beta > \gamma$ as reported in the literature.²⁸ The structural variations of manganese dioxides result from the different bonding ways of the basic [MnO₆] octahedral units. Among them, α -MnO₂ possesses both (2 × 2) and (1 × 1) tunnels surrounded by double binding octahedral chains, β -MnO₂ consists of only (1 × 1) tunnels separated by single chains, and γ -MnO₂ displays (1 × 1) and (1 × 2) tunnels enveloped in double chains.²⁹ To the best of our knowledge, morphology is another important influential factor for the electrochemical properties. It was reported that α -MnO₂ nanowires possess enhanced electrocatalytic activity compared to the other two shapes namely nanotubes and nanoparticles despite the nanotubes showing a much higher specific surface area.³⁰

Previous research on manganese dioxide as the catalysts toward the ORR mainly focused on the effect of the morphology, crystallographic structure and chemical composition on the catalytic activity.^{31–34} Recently, most research has been concentrated on further improving the ORR activity by hybridizing or doping manganese oxides with other transition metals, such as Ag,^{35–37} Pd³⁸ and Ni.^{39,40} Zr⁴⁺ doped cathode materials for metal–ion batteries could improve electronic structure and electrochemical performance.^{41,42} Up to now, related studies on Zr⁴⁺ doped α -MnO₂ have not been reported yet.

In the present work, Zr⁴⁺ doped α -MnO₂ nanowires were prepared via an *in situ* hydrothermal method. The structures and properties were characterized using X-ray diffraction (XRD), scanning electron microscopy (SEM), transmission electron

^aDepartment of Applied Chemistry, School of Chemical Engineering and Technology, Tianjin University, Tianjin 300072, PR China. E-mail: wangweipaper@163.com; Fax: +86-22-27403389; Tel: +86-13512958953

^bKimball Union Academy, NH 03770, USA

microscopy (TEM) and X-ray photoelectron spectroscopy (XPS). The electrocatalytic activity of Zr^{4+} doped $\alpha\text{-MnO}_2$ nanowires for ORR was evaluated by cathodic polarization curve (CPC) and electrochemical impedance spectroscopy (EIS) measurements. The enhancement mechanism of Zr^{4+} doping was further explored by density functional theory (DFT) calculations.

2. Experimental

2.1. Catalyst synthesis

Manganese dioxide ($\alpha\text{-MnO}_2$) nanowires were synthesized *via* a hydrothermal method as reported in the literatures.^{43,44} Typically, 30 mL of 0.1 M KMnO_4 aqueous solution was dropped into 30 mL of the MnSO_4 solution (0.1 M) at room temperature with vigorous stirring for 30 min. Then the mixture was transferred into a 100 mL Teflon-lined autoclave reactor to perform a hydrothermal reaction at 140 °C for 24 h in a hot air oven. After that, the reactor was cooled naturally to room temperature. The obtained precipitate was washed several times with distilled water and ethanol. Finally, the product was dried overnight in an oven at 60 °C and used for further characterization.

For the synthesis of Zr^{4+} doped $\alpha\text{-MnO}_2$ nanowire catalysts, a certain amount of $\text{Zr}(\text{NO}_3)_4 \cdot 5\text{H}_2\text{O}$ was added into 30 mL of 0.1 M MnSO_4 solution under vigorous stirring until the formation of a homogeneous solution. The dosage of $\text{Zr}(\text{NO}_3)_4 \cdot 5\text{H}_2\text{O}$ changed depending on the Zr/Mn molar ratio desired 1 : 70, 1 : 90, 1 : 110, 1 : 140 and 1 : 150. The other preparation processes were exactly the same with the preparation of $\alpha\text{-MnO}_2$ nanowires.

2.2. Physical characterization

The X-ray diffraction (XRD) patterns of the prepared $\alpha\text{-MnO}_2$ and Zr^{4+} doped $\alpha\text{-MnO}_2$ catalysts were recorded with a Rigaku MiniFlex II X-ray diffractometer using Cu K α radiation with a Ni filter. The tube current was 30 mA with a tube voltage of 40 kV. The 2θ angular regions between 20° and 75° were explored at a scan rate of 4° min⁻¹. The morphologies and surfaces of the prepared catalysts were investigated using a Nanosem 430 field-emission scanning electron microscope (FE-SEM, FEI Company) with an acceleration voltage range of 10–30 kV and a Philips Tecnai G2 F20 field-emission transmission electron microscope (TEM, FEI Company) operated at 200 kV. The binding energy of the elements in the catalysts was analyzed by X-ray photoelectron spectroscopy (XPS, PHI1600 ESCA System, PERKIN ELMER, USA) with Al K α radiation ($h\nu = 1486.6$ eV).

2.3. Preparation of the air electrode

Briefly, the air electrode was prepared by dispensing 2.2 g of the synthesized catalyst, 1.1 g of acetylene black and 2.1 g of polytetrafluoroethylene (PTFE, 60 wt%) into 20 mL of ethanol. The mixture was stirred at room temperature for 1 hour and then stirred continually at 60 °C for 15 minutes until the mixture getting together. Afterwards, the mixture was rolled to form a film with thickness 0.6 mm. Finally, the film was coated onto a stainless steel net with area 2 cm × 2 cm.

2.4. Electrochemical measurements

The electrochemical measurements were carried out in a three-electrode and two-circuit electrochemical cell. One side of the working electrode was immersed in a 3 M KOH solution and the other side was exposed to the air. A ruthenium–titanium mesh was used as a counter electrode. An Hg/HgO (3 M KOH) reference electrode was used through a salt bridge, and all the potentials in this work were referred to Hg/HgO (3 M KOH) reference electrode. The cathodic polarization curves were recorded from the open circuit potentials and the scan rate of the potential was 0.5 mV s⁻¹. Electrochemical impedance spectroscopy (EIS) measurements were performed with an applied potential amplitude ± 5 mV in a frequency range from 10 kHz to 10 mHz. All the electrochemical measurements were conducted in 3 M KOH aqueous solution at 25 ± 0.1 °C. Distilled water and analytical grade reagents were used to prepare the solutions.

The electrochemical tests for charge transfer and durability analyses were carried out on a CHI660D electrochemical workstation (CH Instruments, China) with a rotation disk electrode (Pine instrument), using a conventional three-electrode electrochemical cell at 25 ± 0.1 °C. A glassy carbon electrode (GCE) with an area of 0.196 cm² was used as the substrate for the working electrode, a ruthenium–titanium mesh and Hg/HgO (1 M KOH) were employed as the counter and reference electrodes, respectively. The catalyst ink was fabricated as follows: 2.67 mg as-prepared Zr^{4+} doped $\alpha\text{-MnO}_2$ and 1.33 mg acetylene black was ultrasonically dispersed in a 1.0 mL solution (0.9 mL of ethanol and 0.1 mL of 5 wt% Nafion solution) for 30 min. Then 10 μL of the ink was pipetted onto the GCE surface, dried in air naturally. Linear sweep voltammetry (LSV) curves were obtained in O₂-saturated 0.1 M KOH solution at different electrode rotated speeds with a potential scan rate of 5 mV s⁻¹. The current–time durability measurement at -0.4 V (vs. Hg/HgO) was conducted by a rotating disk electrode (RDE) in O₂-saturated 0.1 M KOH solution with a rotation rate of 1600 rpm. Koutecky–Levich plots (K–L plots) was fitted by a line and the slopes was used to analyse the number of electrons transferred (n) during the ORR on the basis of Koutecky–Levich equation (K–L equation)⁴⁵

$$\frac{1}{J} = \frac{1}{J_k} + \frac{1}{B\omega^{1/2}} \quad (1)$$

$$B = 0.2nFD^{2/3}\nu^{-1/6}C_{\text{O}_2} \quad (2)$$

where J is the measured current density, J_k the kinetic current density, B the Levich constant, ω the angular velocity of the rotating electrode, F the Faraday constant (96 485 C mol⁻¹), D the diffusion coefficient of oxygen (1.9×10^{-5} cm² s⁻¹), ν the kinetic viscosity of the electrolyte (0.01 cm² s⁻¹), C_{O_2} the bulk concentration of oxygen in electrolyte (1.2×10^{-6} mol cm⁻³), and n the number of electron exchanged during ORR process. The constant is adopted to be 0.2 when ω is expressed in rpm.⁴⁶

2.5. Density functional theory(DFT) calculations

CASTEP^{47,48} based on the Density Functional Theory (DFT) was used to investigate the O adsorption of $\alpha\text{-MnO}_2$ and Zr^{4+} doped



α -MnO₂ by solving the Kohn–Sham equation of a many-body system with an iterative approach. The system was under the periodic boundary condition and computation was done in a reciprocal space. Plane-wave basis sets with a cutoff energy of 340 eV. The General Gradient Approximation (GGA)⁴⁹ of Perdew, Burke and Ernzerhof (PBE)^{50–52} was used as the exchange–correlational functional to describe the interactions among electrons. The Brillouin zone integration was done on a grid of $3 \times 2 \times 1$ with a Gaussian smearing parameter δ of 0.1 eV and a Self-Consistence-Field (SCF) convergence criterion of 1×10^{-5} eV.⁵³ Initial structures were optimized by allowing the atoms positions to be relaxed while the iterative algorithm was updated by the conjugate gradient approach with convergence criteria of 0.05 eV Å^{−1}. The crystal plane (211) of α -MnO₂ surface was modeled using three-layer periodic slabs repeating in a (2×2) surface structure. Periodic boundary conditions were applied in all three dimensions with a vacuum region of 12 Å along the z -axis. The atoms in the top two layers of the slab were relaxed, whereas the atoms in the third layer were frozen at their bulk positions. Oxygen was placed on one side of the slab. Excluding these species and the top layer of the slab that were allowed to relax in their positions to reach the most stable configuration, all other atoms were frozen at their ideal bulk positions to simulate the bulk and reduce the computational cost.

The adsorption energies (ΔE_{ads}) were defined relative to the isolated substrate and the molecules in the gas phase, as shown in eqn (3). A negative ΔE_{ads} value indicates favorable (exothermic) adsorption.

$$\Delta E_{\text{ads}} = E_{(\text{adsorbate/catalyst})} - E_{\text{catalyst}} - E_{\text{adsorbate}} \quad (3)$$

3. Results and discussions

3.1. Characterization

Fig. 1 shows the XRD patterns of the synthesized α -MnO₂ and Zr⁴⁺ doped α -MnO₂ catalysts with different Zr/Mn molar ratios. It can be seen from Fig. 1a that all the patterns can be indexed to body-centered tetragonal α -MnO₂ (JCPDS NO. 44-0141, space group $I4/m$, $a = b = 9.784$ Å, $c = 2.863$ Å), indicating that the crystal structures of the synthesized catalysts are the same with α -MnO₂. It can also be observed from Fig. 1b that the positions of the two characteristic peaks at $2\theta = 28.9$ and 37.6

corresponding to crystal planes (310) and (211) remain the same as that of α -MnO₂ when the Zr/Mn molar ratios in the catalysts are 1 : 140 and 1 : 150. With increasing Zr/Mn molar ratio, a shift in the positions of the two characteristic peaks in Fig. 1b appears at Zr/Mn molar ratio 1 : 110 and the shift gets bigger at Zr/Mn molar ratio 1 : 90 and 1 : 70. The shift in the peak positions means that a lattice distortion exists. The lattice parameters were calculated by the data in Fig. 1 and shown in Table 1. The lattice parameters a , b and c of Zr⁴⁺ doped α -MnO₂ become smaller compared with that of α -MnO₂. The changes in the lattice parameters also show a distortion in the lattice cell. The results above indicate that α -MnO₂ is successfully synthesized and element zirconium is doped into the lattice structure of α -MnO₂.

Fig. 2 shows the FE-SEM morphologies of α -MnO₂ and Zr⁴⁺ doped α -MnO₂ synthesized in this work. The two kinds of synthesized products all possess a wire shape with diameter less than 100 nm. It can also be found that the surface of α -MnO₂ nanowires (Fig. 2a) is smoother than that of Zr⁴⁺ doped α -MnO₂ nanowires (Fig. 2b).

The high resolution TEM images of synthesized α -MnO₂ and Zr⁴⁺ doped α -MnO₂ nanowires (Zr/Mn molar ratio of 1 : 110) are shown in Fig. 3. The nanowires are straight in Fig. 3a and c, indicating that the crystal growth is along a preferred orientation. The diameter of synthesized α -MnO₂ nanowires is in the range of 10 to 50 nm and that of synthesized Zr⁴⁺ doped α -MnO₂ nanowires is in the range of 30 to 80 nm. The surfaces of α -MnO₂ nanowires are clean and smooth (Fig. 3a and b). As for Zr⁴⁺ doped α -MnO₂ nanowires, some attachments exist on the nanowires (Fig. 3c and d). The attachments possess an irregular morphology and its crystal structure is different from the nanowires. The different crystal structure between the attachments and nanowires illustrates that the attachments and nanowires are different materials. A semi-coherent interfacial relationship can also be found between the nanowires and the attachments as shown in Fig. 3d by the arrow, demonstrating that the attachments grow from the surface of the nanowires. The results above reveal that Zr⁴⁺ doped α -MnO₂ nanowires have been successfully synthesized and few attachments differing from α -MnO₂ in structure are formed at high Zr/Mn molar ratios.

XPS was used to probe the composition and chemical state of the synthesized nanowires, and the results are shown in Fig. 4. Compared with α -MnO₂ nanowires, a signal related to element zirconium is detected in Zr⁴⁺ doped α -MnO₂ nanowires as

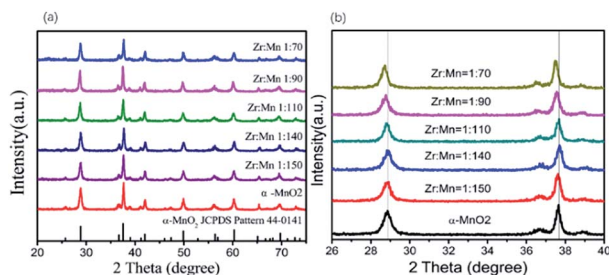


Fig. 1 XRD patterns of Zr⁴⁺ doped α -MnO₂ with different Zr/Mn molar ratios.

Table 1 Lattice parameters calculated by the data in Fig. 1

Sample	Lattice parameter		
	a (Å)	b (Å)	c (Å)
α -MnO ₂	9.815	9.815	2.847
Zr/Mn1 : 150	9.7998	9.7998	2.845
Zr/Mn1 : 140	9.7976	9.7976	2.842
Zr/Mn1 : 110	9.7969	9.7969	2.845
Zr/Mn1 : 90	9.7960	9.7960	2.844
Zr/Mn1 : 70	9.7952	9.7952	2.845



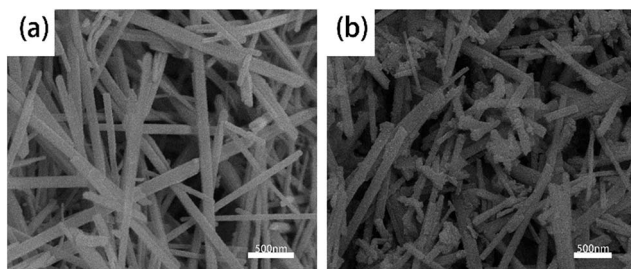


Fig. 2 FE-SEM images of (a) α -MnO₂ nanowires and (b) Zr⁴⁺ doped α -MnO₂ nanowires with Zr/Mn molar ratio of 1 : 110.

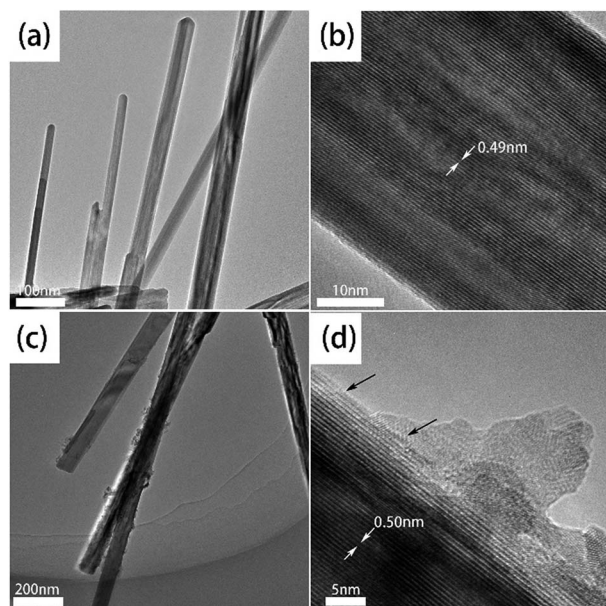


Fig. 3 TEM images of α -MnO₂ nanowires (a and b) and Zr⁴⁺ doped α -MnO₂ nanowires (c and d) with Zr/Mn molar ratio of 1 : 110.

shown in Fig. 4a. Fig. 4b is the XPS spectrums of O 1s for α -MnO₂ nanowires and Zr⁴⁺ doped α -MnO₂ nanowires. Curve simulations were conducted and the simulated curves are also shown in Fig. 4b. The fitted curve is in accord with the measured data, demonstrating the simulated results are reasonable. The binding energy values at the peak positions 529.4 eV and 531.7 eV in the O 1s spectrum of α -MnO₂ nanowires correspond to the lattice oxygen and the adsorbed oxygen, respectively.^{54,55} It can also be seen from Fig. 4b that the peak position corresponding to lattice oxygen shifts to 529.2 eV while the peak position corresponding to adsorbed oxygen keeps the same after Zr⁴⁺ doping, which means that the chemical state of lattice oxygen in Zr⁴⁺ doped α -MnO₂ nanowires is altered compared with that in α -MnO₂ nanowires.

In order to obtain the detail information about the presence of element manganese, the XPS spectra of Mn 2p for α -MnO₂ nanowires were simulated with three groups of curves colored by pink, dark yellow and dark blue, respectively (Fig. 4c). The fitted curve is in accord with the measured data, demonstrating the simulated results are reasonable. The three groups of curves

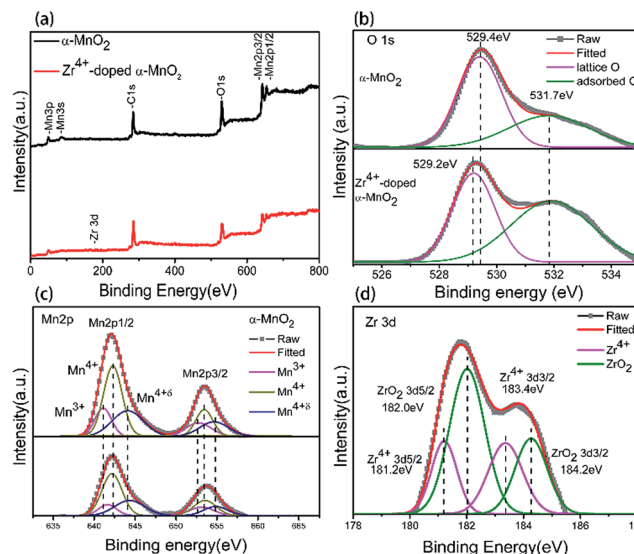


Fig. 4 XPS spectrums of α -MnO₂ and Zr⁴⁺ doped α -MnO₂ with Zr/Mn molar ratio of 1 : 110. (a) Full spectrums; (b) O 1s; (c) Mn 2p; (d) Zr 3d.

correspond well to the XPS spectra of Mn³⁺ (pink), Mn⁴⁺ (dark yellow) and Mn^{4+δ} (dark blue) in α -MnO₂,^{56,57} indicating that there are three kinds of element manganese with different valences in α -MnO₂ nanowires. Comparing the XPS spectra of α -MnO₂ nanowires and Zr⁴⁺ doped α -MnO₂ nanowires in Fig. 4c, it can be found that the peak position corresponding to Mn³⁺ shifts positively with 0.5 eV while the peak position corresponding to Mn⁴⁺ and Mn^{4+δ} remains the same after Zr⁴⁺ doping. The positive shift of Mn³⁺ binding energies with a value less than 1 eV means that the chemical state of Mn³⁺ ions in Zr⁴⁺ doped α -MnO₂ has been changed and Mn³⁺ ions tend to lose their electrons by Zr⁴⁺ doping.

Fig. 4d is Zr3d XPS spectrum of Zr⁴⁺ doped α -MnO₂ nanowires. The measured data were simulated with two group curves colored with pink and green. The fitted curve is consistent with the measured data, demonstrating the simulated results are reasonable. The binding energy values of green peaks in Fig. 4d are separately 182.0 eV and 184.2 eV, which is in consistent with that of ZrO₂.⁵⁸ The result manifests the existence of ZrO₂. The binding energy values of pink peaks in Fig. 4d are separately 181.2 eV (Zr3d5/2) and 183.4 eV (Zr3d3/2), which are 0.8 eV negative shifts compared with that of green peaks in Fig. 4d. The negative shift in Zr3d binding energies with a value less than 1 eV means that some zirconium ions in Zr⁴⁺ doped α -MnO₂ tend to get electrons. Comprehensively considering the XPS results in Fig. 4, it can be concluded that Mn³⁺ ions, Mn⁴⁺ ions, Mn^{4+δ} ions and Zr⁴⁺ ions co-existed in the crystal structure of synthesized Zr⁴⁺ doped α -MnO₂ nanowires. Zr⁴⁺ ions occupied the positions originally belonging to element manganese in the crystal structure and resulted in a mutual action between Zr⁴⁺ ions and Mn³⁺ ions. The mutual action made Mn³⁺ ions tend to lose their electrons and Zr⁴⁺ ions tend to get electrons. The existence of Zr⁴⁺ ions in the crystal structure also changed the chemical state of the lattice oxygen. XPS analyses in Fig. 4d also indicate that there are some ZrO₂ in Zr⁴⁺ doped α -MnO₂,



which can be considered to be the attachments shown in Fig. 3d.

3.2. Electrochemical activity of synthesized catalysts of ORR

The cathodic polarization curves of the air electrodes prepared by α -MnO₂ nanowires and Zr⁴⁺ doped α -MnO₂ nanowires with different Zr/Mn molar ratios were measured and the results are shown in Fig. 5a. As the potential moves negatively, the current density of the electrodes increases slowly in the potential range from 0.3 V to 0.1 V. Then the current density begins to rise rapidly with the potential shifting more negative than 0.1 V. According to the linear sweep voltammetry curves in Fig. 5a, the onset potentials of the electrodes were plotted in Fig. 5b. The onset potential of the electrodes prepared by pure α -MnO₂ is 0.02 V (vs. Hg/HgO) while that of the electrodes prepared by Zr⁴⁺ doped α -MnO₂ catalyst with molar ratio Zr : Mn = 1 : 90 shows the highest onset potential (0.082 V). It can be observed that the onset potential increases with the rise of Zr : Mn ratio and drops after reaching the peak value. To clearly show the effect of Zr/Mn molar ratios in the catalysts on the current density, the current densities at potential -0.2 V in Fig. 5a were plotted against Zr/Mn molar ratios and the curve is shown in Fig. 5c. The current density in Fig. 5c alters in a volcano shape with Zr/Mn molar ratio increasing. The largest current density (60 mA cm⁻²) emerges at Zr/Mn molar ratio of 1 : 110, which is 23% larger than that of the electrode prepared with α -MnO₂ nanowires (49 mA cm⁻²). The current during the cathodic polarization process is caused by ORR on the surface of the electrodes. The larger the current density during the cathodic polarization process is, the more efficiently the catalyst promotes ORR. The results indicate that the electrocatalytic activity of α -MnO₂ on ORR can be remarkably improved by Zr⁴⁺ doping. After Zr/Mn molar ratios exceeding 1 : 110, excessive zirconium forms ZrO₂ crystals attached on the surfaces of Zr⁴⁺ doped α -MnO₂ nanowires as shown in Fig. 3d (the attachments). The existence of ZrO₂ attachments reduces the catalytical area of Zr⁴⁺ doped α -MnO₂ nanowires, resulting in the decrease of the current density as shown in Fig. 5b.

The electrochemical impedance spectroscopies (EIS) of the air electrodes prepared by α -MnO₂ nanowires and Zr⁴⁺ doped α -MnO₂ nanowires with different Zr/Mn molar ratios as catalysts were measured and shown in Fig. 6. All the EIS plots in Fig. 6a

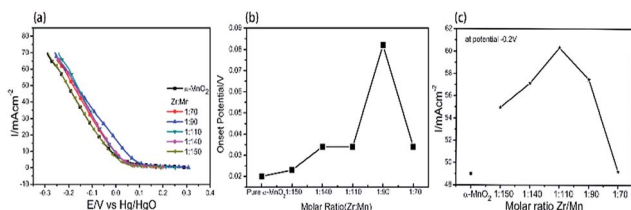


Fig. 5 (a) Cathodic polarization curves of the air electrodes prepared with α -MnO₂ nanowires and Zr⁴⁺ doped α -MnO₂ nanowires with different Zr/Mn molar ratios as catalysts. (b) The onset potentials of the electrodes prepared by Zr⁴⁺ doped α -MnO₂ catalyst with different Zr/Mn ratios (c) the relations between Zr/Mn molar ratios and the current densities at potential -0.2 V.

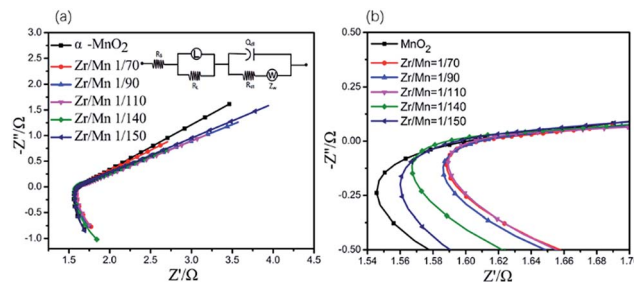
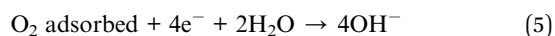


Fig. 6 (a) EIS of the air electrodes prepared by α -MnO₂ nanowires and Zr⁴⁺ doped α -MnO₂ nanowires with different Zr/Mn molar ratios as catalysts. Upper inset is the equivalent circuit. (b) The amplified part of the EIS. Dot: measured data; solid line: simulated results.

exhibit the same shape, which are composed of a large arc in the high frequency range in the fourth quadrant linked with a very small semi-circle in the middle frequency range in the first quadrant followed by a long straight line in the low frequency range in the first quadrant.

The reduction of oxygen molecules on the surface of the air electrodes can be considered to proceed by following two steps.⁵⁹ In the first step, oxygen molecules from the air diffuse through the air electrode and absorb on the surface of the air electrode near the solution side, which can be described in eqn (4). In the second step, absorbed oxygen molecules are electrochemically reduced and forms OH⁻, which can be described in eqn (5).



The very small semi-circle in the EIS (Fig. 6) means that the electrochemical reduction of absorbed oxygen molecules described in eqn (5) is easy to proceed. The large arc in the EIS (Fig. 6) indicates that the adsorption of oxygen molecules described in eqn (4) is difficult to proceed. This suggests that the adsorption of oxygen molecules on the surface of the air electrodes is the rate-controlling step for ORR.

The EIS data were simulated using an equivalent circuit shown in Fig. 6b, and the simulated results are also shown in Fig. 6a in solid lines. Therein, R_s represents the solution resistance, R_{ct} the charge transfer resistance incurred during the electrochemical reactions taking place on the electrodes, Q the constant phase element modeling the non-faradaic processes, Z_w the Warburg element representing the diffusion control process, L the inductance element representing the adsorption of oxygen molecules on the electrode, R_L the adsorption resistance. The solid lines in Fig. 6a fit well with the measured data, demonstrating the proposed equivalent circuit are reasonable. Some of the parameters in the equivalent circuit are listed in Table 2. It is well known that the reaction rate of the electrochemical reaction step is inversely proportional to R_{ct} .⁶⁰ All the values of R_{ct} in Table 2 are small, illustrating that the reaction described in eqn (5) is easy to proceed. The electrode prepared with Zr⁴⁺ doped α -MnO₂ nanowires with Zr/Mn molar ratio of 1 : 110 shows the smallest



Table 2 Equivalent circuit elements and their values

Sample	L (H)	R_L (Ω)	R_s (Ω)	Z_W (Ω)	R_{ct} (Ω)
α -MnO ₂	1.422	4.685	1.804	1.57	29.81
1 : 70	1.367	3.221	1.565	1.84	27.34
1 : 90	1.336	3.195	1.524	2.68	24.42
1 : 110	1.391	3.141	1.308	2.46	20.64
1 : 140	1.886	3.721	1.679	1.67	29.34
1 : 150	1.53	4.849	1.9745	1.34	37.88

value of R_L . The smaller R_L value is, the more easily the adsorption process described in eqn (4) proceeds. Considering the adsorption of oxygen molecules is the rate-controlling step of ORR on the air electrode, the lowest R_L value reasonably explains why the catalyst with Zr/Mn molar ratio 1 : 110 exhibits the highest electrocatalytical activity as shown in Fig. 6.

To further elucidate the pathway and kinetics of ORR, LSV curve of the electrode prepared by Zr⁴⁺ doped α -MnO₂ on RDE were recorded at different rotation rates (400–1600 rpm) in O₂-saturated 0.1 M KOH solution and the result is shown in Fig. 7a. The derived K-L plot of J^{-1} vs. $\omega^{-1/2}$ at a potential of -0.4 V is shown in Fig. 7b. It can be found that a good linear relationship between J^{-1} vs. $\omega^{-1/2}$. The electron transfer number n was calculated by eqn (2) based on the slope (1/B) of the K-L plot. The electron transfer number for Zr⁴⁺ doped α -MnO₂ is 3.8, which clearly indicate that the electrode prepared by Zr⁴⁺ doped α -MnO₂ follows a desirable four-electron pathway for ORR.

The stability of catalysts is very important issue for their practical application in various energy technologies. Stability was examined during the ORR process. The durability of the electrode prepared by Zr⁴⁺ doped α -MnO₂ and Pt/C as the catalyst for ORR was evaluated by chronoamperometric technique at potential -0.4 V for 7200 s. It can be seen from Fig. 7c that the ORR current density of the Zr⁴⁺ doped α -MnO₂ electrode decreased by only 16.3%, but the Pt/C electrode exhibited a 30.1% decrease in current density under the same conditions. The result reveals that the performance of Zr⁴⁺ doped α -MnO₂ catalyst is more stable than that of the commercial Pt/C catalyst.

3.3. Optimized configuration and O₂ adsorption

To understand the influence of Zr⁴⁺ doping on the adsorption of oxygen molecules, the DFT calculations were carried out aiming

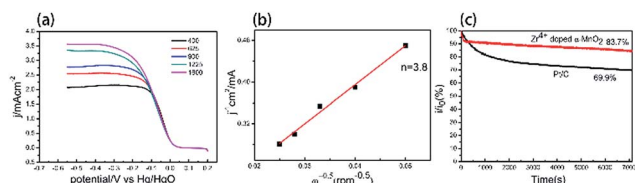


Fig. 7 (a) LSV curves of the electrodes prepared by Zr⁴⁺ doped α -MnO₂ catalyst at different rotation speed (400–1600 rpm) in oxygen-saturated 0.1 M KOH solution. (b) Koutecky–Levich (K–L) plots. (c) Chronoamperometric curves of the electrode prepared by Zr⁴⁺ doped α -MnO₂ and Pt/C measured in oxygen-saturated 0.1 M KOH solution at a rotating rate 1600 rpm.

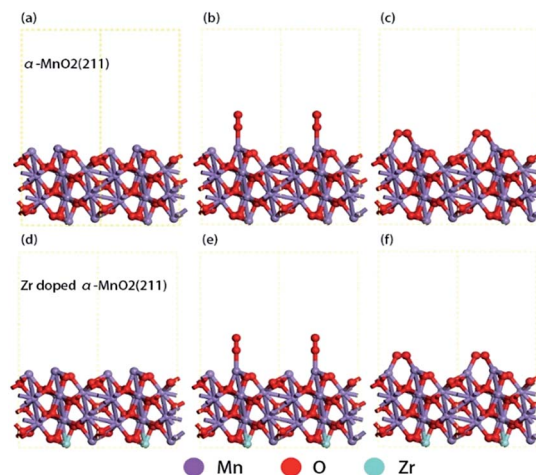


Fig. 8 Optimized adsorption structures. (a) Side view of the optimized α -MnO₂ (211) surface; (b) O₂ top adsorption on α -MnO₂ (211) surface; (c) O₂ bridge adsorption on α -MnO₂ (211) surface; (d) side view of the optimized Zr⁴⁺ doped α -MnO₂ (211) surface; (e) O₂ top adsorption on Zr⁴⁺ doped α -MnO₂ (211); (f) O₂ bridge adsorption on Zr⁴⁺ doped α -MnO₂.

at the surface of crystal plane (211) of α -MnO₂ (Fig. 8a and d). Different adsorption configurations of O₂ molecules were optimized by the computation methods mentioned in 2.4. The optimized adsorption structures on the surfaces are shown in Fig. 8b, c, e and f. The direct adsorption of O₂ molecules on the surfaces includes the following two modes: top adsorption (Fig. 7b and e) and bridge adsorption (Fig. 7c and f). The adsorption energy of O₂ molecules was calculated by eqn (3) and the results were listed in Table 3.

As shown in Table 3, the bridge adsorption energy is more negative than top adsorption energy, indicating that bridge adsorption is more favorable for O₂ molecules. Comparing with α -MnO₂, the bridge adsorption energy of O₂ molecules on Zr⁴⁺ doped α -MnO₂ decreases by 0.26 eV. The more negative adsorption energy indicates an easier adsorption process and a more stable adsorption structure.^{61,62} The results in Table 3 reveal that the mutual action between Zr⁴⁺ and Mn³⁺ in Zr⁴⁺ doped α -MnO₂ nanowires enhances the adsorption process of O₂ molecules. The bridge adsorption configuration is supposed to be the preferential adsorbing manner for O₂ molecules on the surface of Zr⁴⁺ doped α -MnO₂. According to EIS analyses, the adsorption process of O₂ molecules is the rate-controlling step for ORR. The results in Table 3 provide a reasonable explanation about the improvement in the performance of ORR by Zr⁴⁺ doping.

Table 3 Adsorption energy of O₂ molecules with different configurations

Surface	Top adsorption/eV	Bridge adsorption/eV
Crystal plane (211) of α -MnO ₂	−1.2242	−2.8777
Crystal plane (211) of Zr doped α -MnO ₂	−1.3313	−3.1317



4. Conclusions

Zr⁴⁺ doped α -MnO₂ nanowires were successfully synthesized by a hydrothermal method. XRD, SEM, TEM and XPS analyses indicated that Mn³⁺ ions, Mn⁴⁺ ions, Mn^{4+ δ} ions and Zr⁴⁺ ions co-existed in the crystal structure of synthesized Zr⁴⁺ doped α -MnO₂ nanowires. Zr⁴⁺ ions occupied the positions originally belonging to element manganese in the crystal structure and resulted in a mutual action between Zr⁴⁺ ions and Mn³⁺ ions. The mutual action made Mn³⁺ ions tend to lose their electrons and Zr⁴⁺ ions tend to get electrons. The existence of Zr⁴⁺ ions in the crystal structure also changed the chemical state of the lattice oxygen. Cathodic polarization analyses showed that Zr/Mn molar ratio notably affects the ORR performance of the air electrodes prepared by Zr⁴⁺ doped α -MnO₂ nanowires. The highest ORR current density of the air electrodes prepared by Zr⁴⁺ doped α -MnO₂ nanowires in alkaline solution appeared at Zr/Mn molar ratio of 1 : 110, which was 23% higher than that prepared by α -MnO₂ nanowires. The results indicated that the electrocatalytic activity of α -MnO₂ for ORR remarkably improved by Zr⁴⁺ doping. EIS analyses indicated that the adsorption process of O₂ molecules on the surface of the air electrodes prepared by Zr⁴⁺ doped α -MnO₂ nanowires was the rate-controlling step for ORR. The calculation results based on density functional theory (DFT) revealed that the mutual action between Zr⁴⁺ and Mn³⁺ in Zr⁴⁺ doped α -MnO₂ nanowires enhanced the adsorption process of O₂ molecules.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This research is supported by the Chinese Natural Science Foundation (No. 21276181). We are grateful for the support.

Notes and references

- X. Huang, Z. Zhao, L. Cao, Y. Chen, E. Zhu, Z. Lin, M. Li, A. Yan, A. Zettl, Y. M. Wang, X. Duan, T. Mueller and Y. Huang, *Science*, 2015, **348**, 1230.
- C. Chen, Y. Kang, Z. Huo, Z. Zhu, W. Huang, H. L. Xin, J. D. Snyder, D. Li, J. A. Herron, M. Mavrikakis, M. Chi, M. N. Markovic, G. A. Somorjai, P. Yang and V. R. Stamenkovic, *Science*, 2014, **343**, 1339.
- V. R. Stamenkovic, B. Fowler, B. S. Mun, G. F. Wang, P. N. Ross, C. A. Lucas and N. M. Markovic, *Science*, 2007, **315**, 493–497.
- V. Komanicky, H. Iddir, K. C. Chang, A. Menzel, G. Karapetrov, D. Hennessy, P. Zapol and H. You, *J. Am. Chem. Soc.*, 2009, **131**, 5732–5733.
- C. L. Lee, C. M. Syu, H. P. Chiou, C. H. Chen and H. L. Yang, *Int. J. Hydrogen Energy*, 2011, **36**, 10502–10512.
- L. Tammeveski, H. Erikon, A. Sarapuu, J. Kozlova, P. Ritslaid, V. Sammelselg and K. Tammeveski, *Electrochem. Commun.*, 2012, **20**, 15–18.
- B. P. Vinayan, K. Sethupathi and S. Ramaprabhu, *Int. J. Hydrogen Energy*, 2013, **38**, 2240–2250.
- G. F. Alvarez, M. Mamlouk and S. M. Senthil Kumar, *J. Appl. Electrochem.*, 2011, **41**, 925–937.
- P. Fu, L. H. Zhou, L. H. Sun, B. H. Huang and Y. Yuan, *RSC Adv.*, 2017, **7**, 13383–13389.
- Y. Z. Xu, C. L. Chen, M. Zhou, G. Y. Fu, Y. Y. Zhao and Y. H. Chen, *RSC Adv.*, 2017, **7**, 26722–26728.
- W. X. Wang, Y. Shi, M. C. Li, Z. Y. Wang, S. F. Wu, F. C. Lyu, C. Q. Shang and Z. G. Lu, *RSC Adv.*, 2016, **6**, 110758–110764.
- R. Bashyam and P. Zelenay, *Nature*, 2006, **443**, 63–66.
- X. Y. Xu, C. X. Shi, Q. Li, R. Chen and T. H. Chen, *RSC Adv.*, 2017, **7**, 14382–14388.
- G. Panomsuwan, N. Saito and T. Ishizaki, *RSC Adv.*, 2016, **6**, 114553–114559.
- X. W. Bai, E. J. Zhao, W. C. Wang, Y. Wang, K. Li, L. Lin, J. C. Yang, H. Sun and Z. J. Wu, *RSC Adv.*, 2017, **7**, 23812–23819.
- S. Jin, A. G. Hubert, Y. Naoaki, N. Haruyuki, B. G. John and S. H. Yang, *Nat. Chem.*, 2011, **3**, 546–550.
- Y. Y. Liang, H. L. Wang, J. G. Zhou, Y. G. Li, J. Wang, T. Regier and H. J. Dai, *J. Am. Chem. Soc.*, 2012, **134**, 3517–3523.
- J. Xu, P. Gao and T. S. Zhao, *Energy Environ. Sci.*, 2012, **5**, 5333–5339.
- Z. S. Wu, S. Yang, Y. Sun, K. Parvea, X. L. Feng and K. Mullen, *J. Am. Chem. Soc.*, 2012, **134**, 9082–9085.
- N. S. Lewis and D. G. Nocera, *Proc. Natl. Acad. Sci. U. S. A., Early Ed.*, 2006, **103**, 15729–15735.
- A. J. Bard and M. A. Fox, *Acc. Chem. Res.*, 1995, **28**, 141–145.
- I. Roche, E. Chainet and M. Chatenet, *J. Phys. Chem. C*, 2007, **111**, 1434–1443.
- M. L. Calegario, F. H. B. Lima and E. A. Ticianelli, *J. Power Sources*, 2006, **158**, 735–739.
- Y. L. Cao, H. X. Yang, X. P. Ai and L. F. Xiao, *J. Electroanal. Chem.*, 2003, **557**, 127–134.
- M. Lanqun, T. Sotomura, K. Nakatsu, N. Koshiba, Z. Dun and T. Ohsaka, *J. Electrochem. Soc.*, 2002, **149**, A504–A507.
- F. H. B. Lima, M. L. Calegario and E. A. Ticianelli, *J. Electroanal. Chem.*, 2006, **590**, 152–160.
- S. Devaraj and N. Munichandraiah, *J. Phys. Chem. C*, 2008, **112**, 4406–4417.
- F. Cheng, Y. Su, J. Liang, Z. Tao and J. Chen, *Chem. Mater.*, 2009, **22**, 898–905.
- Y. J. Zhang, C. S. Sun, P. Lu, K. Y. Li, S. Y. Song and D. F. Xue, *CrystEngComm*, 2012, **14**, 5892–5897.
- K. Selvakumar, S. M. S. Kumar, R. Thangamuthu, G. Kruthika and P. Murugan, *Int. J. Hydrogen Energy*, 2014, **39**, 21024–21036.
- F. Cheng, J. Shen, W. Ji, Z. Tao and J. Chen, *ACS Appl. Mater. Interfaces*, 2009, **1**, 460–466.
- W. Xiao, D. Wang and X. W. Lou, *J. Phys. Chem. C*, 2010, **114**, 1694–1700.
- L. Mao, D. Zhang, T. Sotomura, K. Nakatsu, N. Koshiba and T. Ohsaka, *Electrochim. Acta*, 2003, **48**, 1015–1023.
- F. Cheng, T. Zhang, Y. Zhang, J. Du, X. Han and J. Chen, *Angew. Chem., Int. Ed.*, 2013, **52**, 2474–2477.



- 35 D. A. Slanac, A. Lie, J. A. Paulson, K. J. Stevenson and K. P. Johnston, *J. Phys. Chem. C*, 2012, **116**, 11032–11039.
- 36 Q. Tang, L. Jiang, J. Qi, O. Jiang, S. Wang and G. Sun, *Appl. Catal., B*, 2011, **104**, 337–345.
- 37 S. W. Liu and X. Qin, *RSC Adv.*, 2015, **5**, 15627–15633.
- 38 W. Sun, A. Hsu and R. Chen, *J. Power Sources*, 2011, **196**, 4491–4498.
- 39 I. Roche, E. Chainet and M. Vondrak, *J. Phys. Chem. C*, 2007, **111**, 1434–1443.
- 40 A. C. Garcia, A. D. Herrera, E. A. Ticianelli, M. Chatenet and C. Poinssignon, *J. Electrochem. Soc.*, 2011, **158**, B290–B296.
- 41 K. A. Walz, C. S. Johnson, J. Genthe, L. C. Stoiber, W. A. Zeltner, M. A. Anderson and M. M. Thackeray, *J. Power Sources*, 2010, **195**, 4943–4951.
- 42 J. Xu, G. Chen, H. J. Zhang, W. Zheng and Y. Li, *J. Appl. Electrochem.*, 2015, **45**, 123–130.
- 43 V. Subramanian, H. W. Zhu and B. Q. Wei, *J. Power Sources*, 2006, **159**, 361–364.
- 44 F. Hashemzadeh, M. M. K. Motlagh and A. Maghsoudipour, *J. Sol-Gel Sci. Technol.*, 2009, **51**, 169–174.
- 45 R. E. Davis, G. L. Horvath and C. W. Tobias, *Electrochim. Acta*, 1967, **12**, 287.
- 46 K. Chen, Y. J. Hao and M. R. Zhang, *RSC Adv.*, 2017, **7**, 5782.
- 47 M. C. Payne, D. C. Allan, T. A. Arias and J. D. Joannopoulos, *Rev. Mod. Phys.*, 1992, **64**, 1045.
- 48 V. Milman, B. Winkler, J. A. White, C. J. Pickard, M. C. Payne, E. V. Akhmataskaya and R. H. Nobes, *Int. J. Quantum Chem.*, 2000, **77**, 895.
- 49 J. A. White and D. M. Bird, *Phys. Rev. B*, 1976, **13**, 5188.
- 50 J. P. Perdrew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865–3868.
- 51 J. P. Perdrew, K. Burk and M. Ernzerhof, *Phys. Rev. Lett.*, 1997, **78**, 1396.
- 52 Y. H. Li, G. Y. Zhong and F. Peng, *Phys. Chem. Chem. Phys.*, 2015, **17**, 21950.
- 53 H. J. Monkhorst and J. Pack, *Phys. Rev. B*, 1976, **13**, 5188.
- 54 F. Wang, H. X. Dai, J. G. Deng, G. M. Bai, K. M. Ji and Y. X. Liu, *Environ. Sci. Technol.*, 2012, **46**, 4034–4041.
- 55 J. B. Jia, P. Y. Zhang and L. Chen, *Appl. Catal., B*, 2016, **189**, 210–218.
- 56 D. A. Pena, B. S. Uphade and P. G. Smirniotis, *J. Catal.*, 2004, **221**, 421–431.
- 57 E. Pargoletti, G. Cappelletti, A. Minguzzi, S. Rondinini, M. Leoni, M. Marelli and A. Vertova, *J. Power Sources*, 2016, **325**, 116–128.
- 58 C. D. Wagner, W. M. Riggs, L. E. Davis and J. F. Moulder, *Handbook Of X-Ray Photoelectron Spectroscopy*, ed. G. E. Mullenberg, Perkin-Elmer Corporation Press, Minnesota, 1979, vol. 2, p. 100.
- 59 Z. Shi, J. J. Zhang, Z. S. Liu, H. J. Huang and D. P. Wikinson, *Electrochim. Acta*, 2006, **51**, 1905–1916.
- 60 Z. He and F. Mansfeld, *Energy Environ. Sci.*, 2009, **2**, 215–219.
- 61 L. Li, Z. D. Wei, S. G. Chen, X. Q. Qi, W. Ding, M. R. Xia, R. Li, K. Xiong, Z. H. Deng and Y. Y. Gao, *Chem. Phys. Lett.*, 2012, **539**, 88–93.
- 62 M. H. Yu, Z. K. Wang and C. Hou, *Adv. Mater.*, 2017, **29**, 1602868.

