

Electronic Supplementary Information

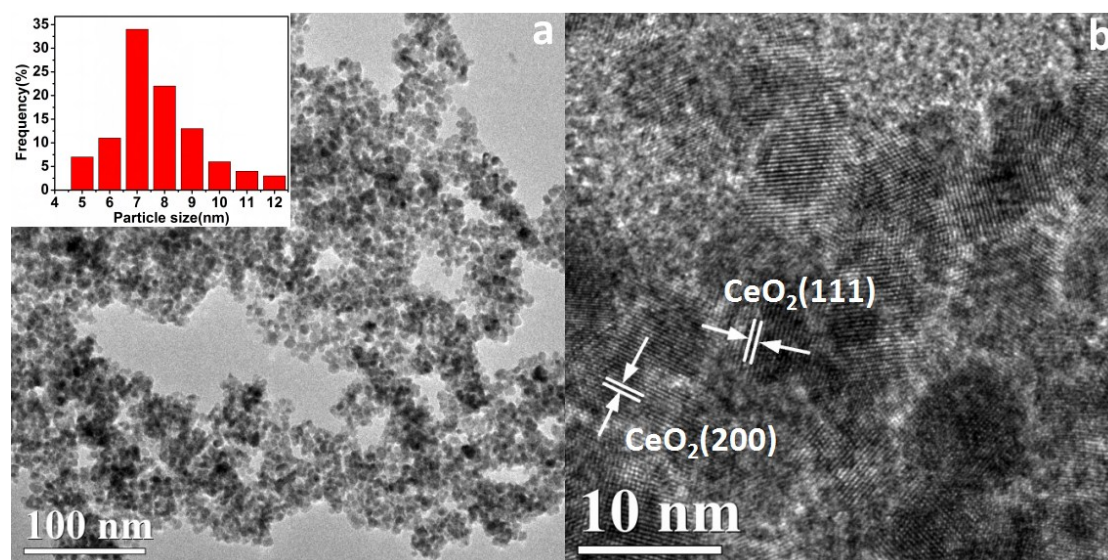


Fig. S1 (a) TEM and (b) HRTEM images of CeO₂ nanoparticles by calcining the CeO₂·xH₂O nanoparticles at 400 °C. Inset in FS1(a) is particle size distribution of the CeO₂ nanoparticles.

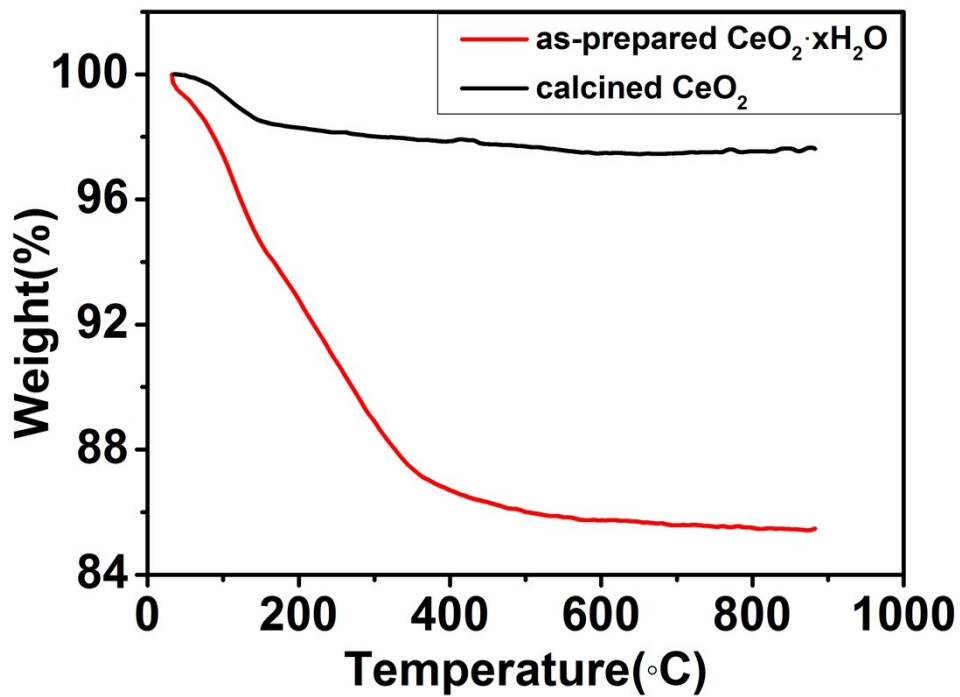


Fig. S2 TGA curves of as-prepared CeO₂·xH₂O nanoparticles and calcined CeO₂ nanoparticles.

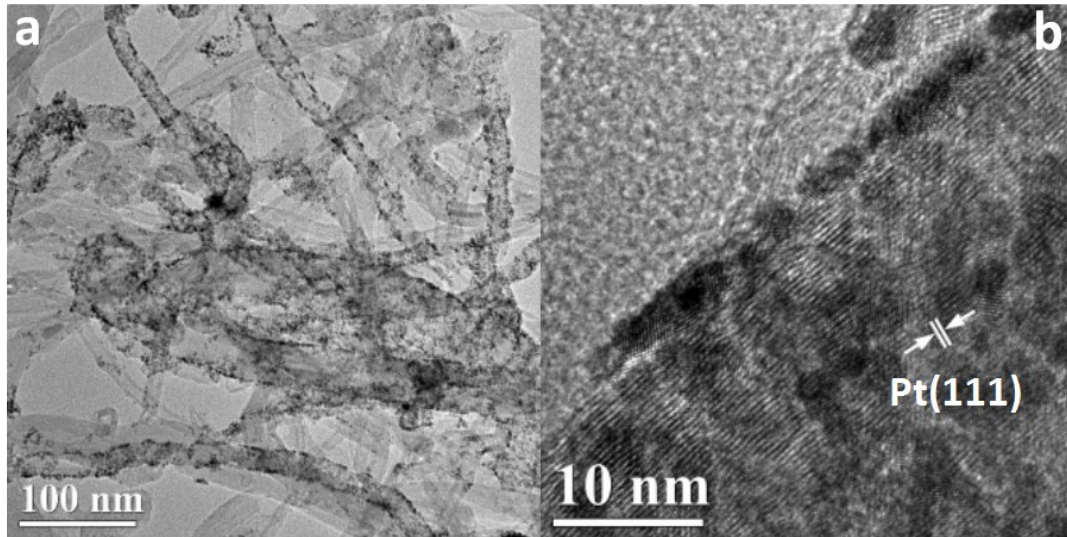


Fig. S3 The TEM and HRTEM images of Pt/CNTs catalyst.

Table S1 Comparison of methanol oxidation behavior in alkaline electrolytes on the various electrocatalysts.

| Catalyst | Mass / specific activity | ESCA(m ² /g) | Scan rate (mV/s) | Electrolyte | Reference |
|--|--------------------------|-------------------------|------------------|----------------------------------|-----------|
| Pt/CNTs + CeO ₂ ·xH ₂ O | 2304mA/mg | | 20mV/s | 1M KOH + 1M CH ₃ OH | This work |
| Pt/Ni(OH) ₂ /rGO | 1236mA/mg | 64.1 | 20mV/s | 1M KOH + 1M CH ₃ OH | 1 |
| Pt/CeO ₂ /rGO | 1140mA/mg | | 50mV/s | 1M KOH + 1M CH ₃ OH | 2 |
| Pt/CeO ₂ /C | 36mA/cm ² | | 50mV/s | 1M KOH + 1M CH ₃ OH | 3 |
| Pt-Mn ₃ O ₄ -MWCNT | 970.2mA/mg | 64.8m ² /g | 50mV/s | | 4 |
| Pt-TiO ₂ /ITO | 1.94mA/cm ² | | 50mV/s | 1M KOH + 1M CH ₃ OH | 5 |
| Pt-TiO ₂ /RGO | 1.36mA/cm ² | | 50mV/s | 0.5M KOH + 1M CH ₃ OH | 6 |
| Pd/MnO ₂ /graphene | 838mA/mg | 82.6m ² /g | 50mV/s | 0.5M KOH + 1M CH ₃ OH | 7 |
| Pd-CeO ₂ /C | 36.6mA/cm ² | | 50mV/s | 1M KOH + 1M CH ₃ OH | 8 |
| Pt/In _{0.1} SnO ₂ | 2320mA/mg | 43m ² /g | 20mV/s | 0.5M KOH + 2M CH ₃ OH | 9 |
| Pt _{3.5} Pb | 2840mA/mg | | 50mV/s | 0.5M KOH + 1M CH ₃ OH | 10 |
| porous Pt ₁ Cu ₃ /1-aminopyrene-graphene | 2750mA/mg | 76.58m ² /g | 50mV/s | 1M KOH + 0.5M CH ₃ OH | 11 |
| Mesoporous Co-Pt films | 4.3 mA/cm ² | | 50mV/s | 1M NaOH + 1M CH ₃ OH | 12 |
| PtNi/C | 1200mA/mg | | 50mV/s | 1M NaOH + 2M CH ₃ OH | 13 |
| PtAg/C | 3mA/mg | 56m ² /g | 20mV/s | 0.5M KOH + 2M CH ₃ OH | 14 |

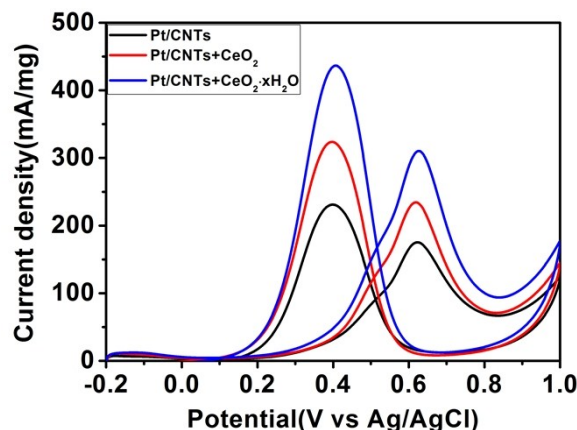


Fig. S4 Cyclic voltammograms curves of Pt/CNTs + CeO₂·xH₂O, Pt/CNTs + CeO₂ and Pt/CNTs catalysts in 0.5M H₂SO₄ and 0.5M CH₃OH with a scan rate of 20 mV/s.

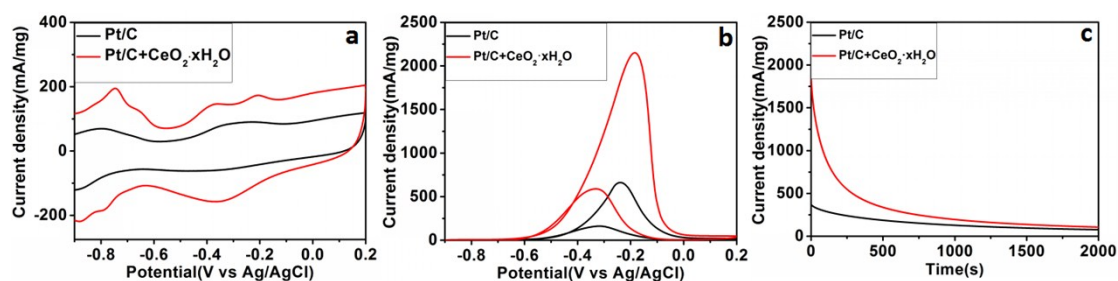


Fig. S5 (a) Cyclic voltammograms curves of methanol electro-oxidation for Pt/C and Pt/C+CeO₂·xH₂O in 1M KOH with a scan rate of 200 mV/s. (b) Cycle voltammograms curves of all the catalysts in 1M KOH and 1M CH₃OH in the potential of -0.9V to 0.2V with a scan rate of 20 mV/s. (c) Chronoamperometric curves of all the catalysts in 1M KOH and 1M CH₃OH.

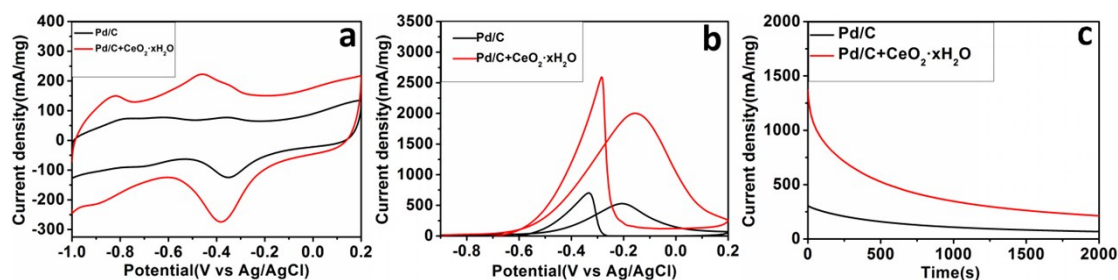


Fig. S6 (a) Cyclic voltammograms curves of methanol electro-oxidation for Pd/C and Pd/C+CeO₂·xH₂O on the all catalysts in 1M KOH with a scan rate of 200mV/s. (b) Cycle voltammograms curves of all the catalysts in 1M KOH and 1M CH₃CH₂OH in the potential of -0.9V to 0.2V with a scan rate of 20mV/s. (c) Chronoamperometric curves of all the catalysts in 1M KOH and 1M CH₃CH₂OH.

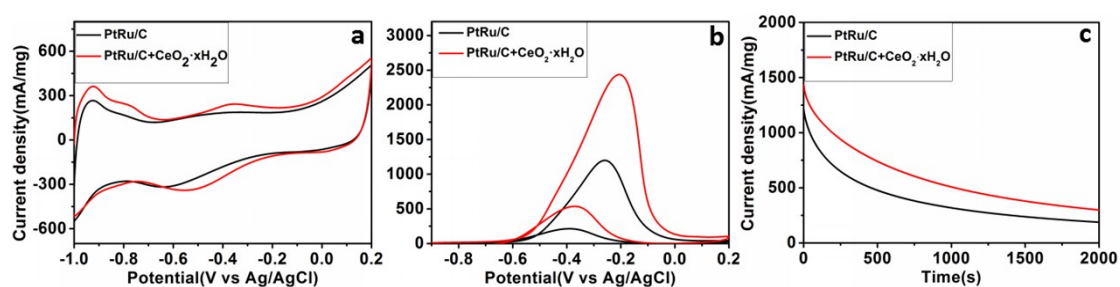


Fig. S7 (a) Cyclic voltammograms of methanol electro-oxidation for PtRu/C and PtRu/C+CeO₂·xH₂O catalysts in 1M KOH with a scan rate of 200mV/s. (b) Cycle voltammograms of all the catalysts in 1M KOH and 1M CH₃OH in the potential of -0.9V to 0.2V with a scan rate of 20mV/s. (c) Chronoamperometric curves of all the catalysts in 1M KOH and 1M CH₃OH.

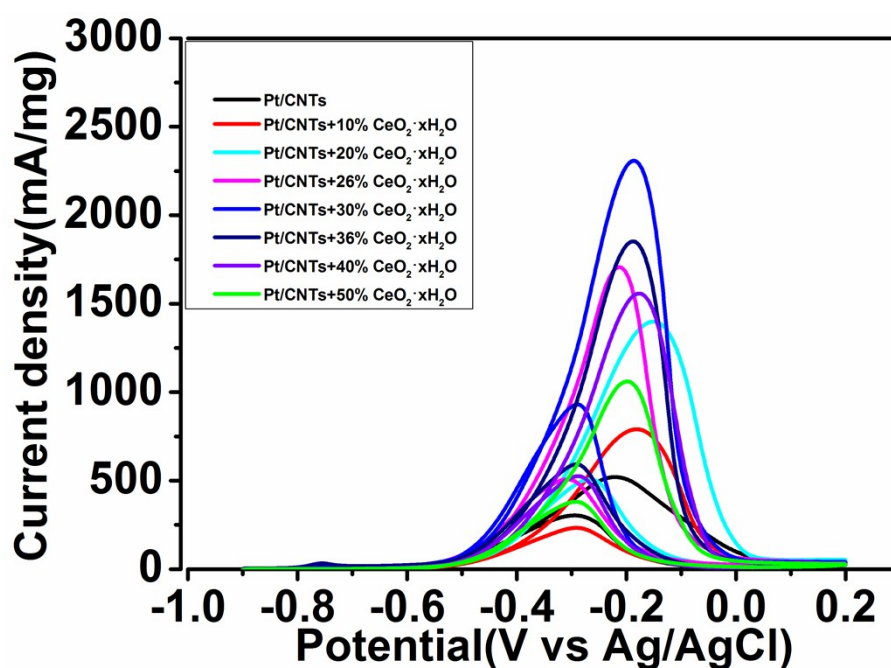


Fig. S8 CVs of methanol electrooxidation in a N₂-saturated solution of 1.0 M KOH and 1.0 M CH₃OH on the Pt/CNTs and Pt/CNTs + CeO₂·xH₂O catalysts prepared with different ratio of CeO₂·xH₂O. The scan rate is 20 mV/s.

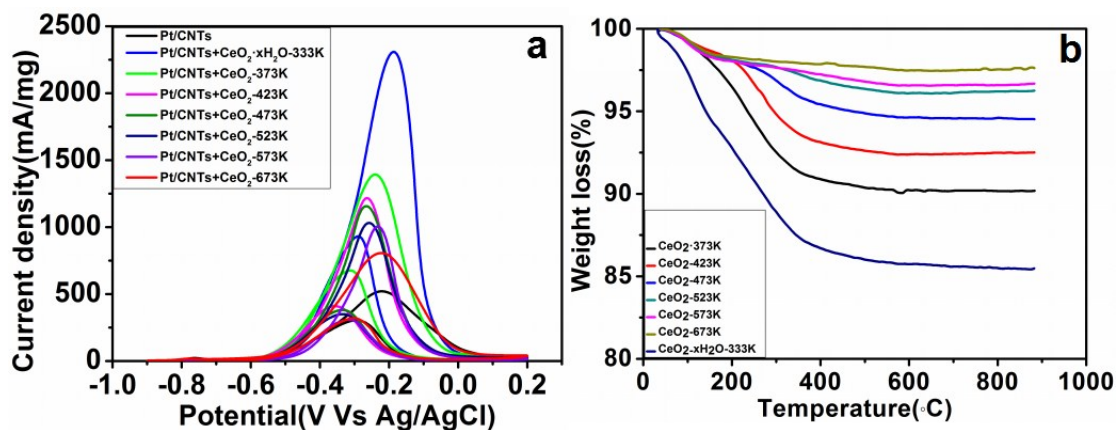


Fig. S9 (a) CVs of Pt/CNTs + $\text{CeO}_2 \cdot x\text{H}_2\text{O}$ catalysts prepared with distinct $\text{CeO}_2 \cdot x\text{H}_2\text{O}$ calcined at different temperature. (b) TGA curves of the distinct $\text{CeO}_2 \cdot x\text{H}_2\text{O}$ nanoparticles.

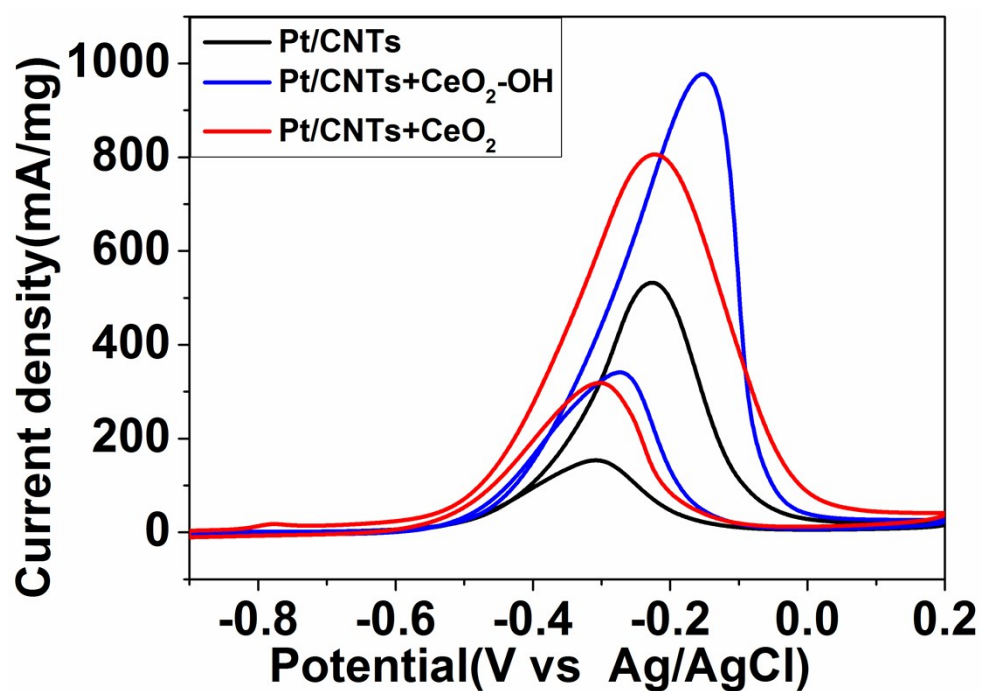


Fig. S10 CVs of Pt/CNTs, Pt/CNTs + CeO_2 , Pt/CNTs + $\text{CeO}_2\text{-OH}$ in 1.0 M KOH + 1.0 M CH_3OH with a scan rate of 20 mV/s. $\text{CeO}_2\text{-OH}$ was obtained by stirring CeO_2 nanoparticles in 1.0 M NaOH solution for 1 h.

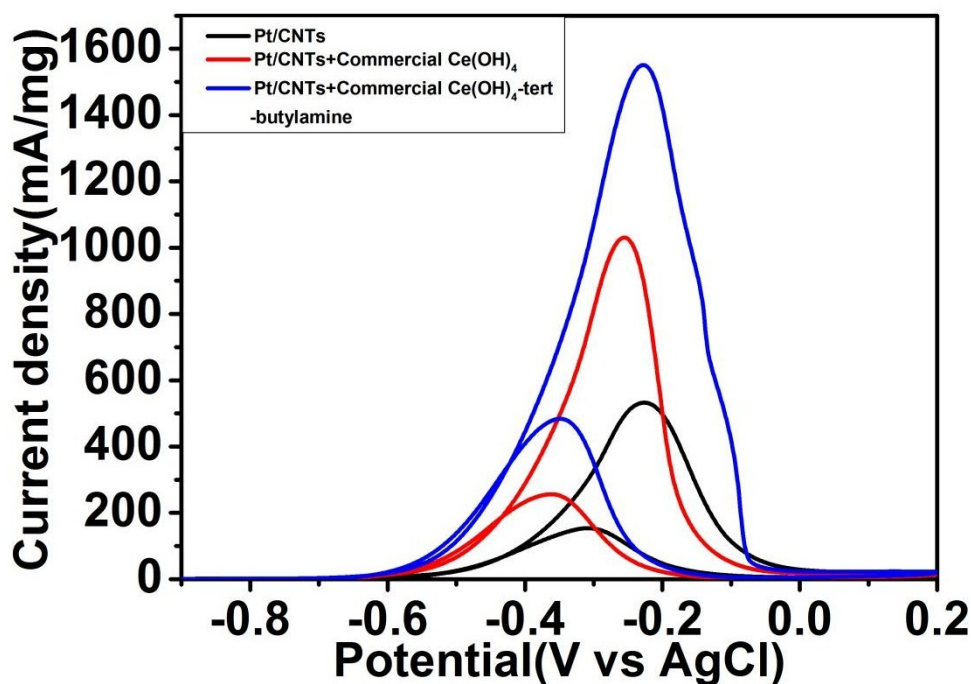


Fig. S11 CVs of Pt/CNTs, Pt/CNTs + commercial $\text{Ce}(\text{OH})_4$ and Pt/CNTs + $\text{Ce}(\text{OH})_4$ -*tert*-butylamine in 1.0 M KOH + 1.0 CH_3OH with a scan rate of 20 mV/s.

Reference

1. W. Huang, H. Wang, J. Zhou, J. Wang, P. N. Duchesne, D. Muir, P. Zhang, N. Han, F. Zhao, M. Zeng, J. Zhong, C. Jin, Y. Li, S. T. Lee and H. Dai, *Nat. Commun.*, 2015, **6**, 10035.
2. Q. He, Y. Shen, K. Xiao, J. Xi and X. Qiu, *Int. J. Hydrogen Energy*, 2016, **41**, 20709-20719.
3. C. Xu and P. K. Shen, *Chem. Commun.*, 2004, 2238-2239.
4. X. Yang, X. Wang, G. Zhang, J. Zheng, T. Wang, X. Liu, C. Shu, L. Jiang and C. Wang, *Int. J. Hydrogen Energy*, 2012, **37**, 11167-11175.
5. H. Zhang, W. Zhou, Y. Du, P. Yang, C. Wang and J. Xu, *Int. J. Hydrogen Energy*, 2010, **35**, 13290-13297.
6. C. Zhai, M. Zhu, D. Bin, H. Wang, Y. Du, C. Wang and P. Yang, *ACS Appl. Mater. Interfaces*, 2014, **6**, 17753-17761.
7. H. Huang and X. Wang, *Phys. Chem. Chem. Phys.*, 2013, **15**, 10367-10375.
8. K.-H. Ye, S.-A. Zhou, X.-C. Zhu, C.-W. Xu and P. K. Shen, *Electrochim. Acta*, 2013, **90**, 108-111.
9. Y.-Y. Feng, W.-Q. Kong, Q.-Y. Yin, L.-X. Du, Y.-T. Zheng and D.-S. Kong, *J. Power Sources*, 2014, **252**, 156-163.
10. L. Huang, Y. Han, X. Zhang, Y. Fang and S. Dong, *Nanoscale*, 2017, **9**, 201-207.
11. G. Zhang, Z. Yang, W. Zhang and Y. Wang, *J. Mater. Chem. A*, 2016, **4**,

- 3316-3323.
12. A. Serrà, E. Gómez, I. V. Golosovsky, J. Nogués and E. Vallés, *J. Mater. Chem. A*, 2016, **4**, 7805-7814.
 13. Q. Jiang, L. Jiang, S. Wang, J. Qi and G. Sun, *Catal. Commun.*, 2010, **12**, 67-70.
 14. Y.-Y. Feng, L.-X. Bi, Z.-H. Liu, D.-S. Kong and Z.-Y. Yu, *J. Catal.*, 2012, **290**, 18-25.