

12-21-2013

Asymmetric Supercapacitors With Dominant Pseudocapacitance Based on Manganese Oxide Nanoflowers in a Neutral Aqueous Electrolyte

Qiang Li

The University of Texas Rio Grande Valley

Xing Sun

The University of Texas Rio Grande Valley

Karen Lozano

The University of Texas Rio Grande Valley, karen.lozano@utrgv.edu

Yuanbing Mao

The University of Texas Rio Grande Valley, yuanbing.mao@utrgv.edu

Follow this and additional works at: https://scholarworks.utrgv.edu/chem_fac

 Part of the [Chemistry Commons](#)

Recommended Citation

Li, Qiang; Sun, Xing; Lozano, Karen; and Mao, Yuanbing, "Asymmetric Supercapacitors With Dominant Pseudocapacitance Based on Manganese Oxide Nanoflowers in a Neutral Aqueous Electrolyte" (2013). *Chemistry Faculty Publications and Presentations*. 56.
https://scholarworks.utrgv.edu/chem_fac/56

This Article is brought to you for free and open access by the College of Sciences at ScholarWorks @ UTRGV. It has been accepted for inclusion in Chemistry Faculty Publications and Presentations by an authorized administrator of ScholarWorks @ UTRGV. For more information, please contact justin.white@utrgv.edu, william.flores01@utrgv.edu.

Electronic Supplementary Information

Asymmetric Supercapacitors with Dominant Pseudocapacitance Based on Manganese Oxide Nanoflowers in Neutral Aqueous Electrolyte

Qiang Li,^{a,b} Xing Sun,^a Karen Lozano^b and Yuanbing Mao^{a,*}

^a Department of Chemistry, University of Texas - Pan American, Edinburg, TX 78539 USA

^b Department of Mechanical Engineering, University of Texas - Pan American, Edinburg, TX 78539 USA

*To whom correspondence should be addressed: Phone: +1 956 665 2417; Fax: +1 956 665 5006;
E-mail: maoy@utpa.edu

Experimental Section

Synthesis of manganese oxide nanoflowers (MnO₂-NFs): In a typical synthesis, 0.8 mmol of KMnO₄ was dissolved into 36 mL of deionized water under vigorous magnetic stirring for 20 min. Then 4.385 mL of N-Methyl-2-pyrrolidone (NMP) was added into the above solution and the mixture was stirred for another 5 min before kept still at 25 – 60 °C for 6 – 20 h. Subsequently, the brown-black precipitate was collected and repeatedly washed with deionized water and ethanol, and then dried in a vacuum oven overnight.

Functionalization of carbon nanotubes (CNTs): 0.1 g of pristine MWCNTs (Sigma Aldrich, product No. 636517) were immersed into 150 mL concentrated nitric acid and refluxed at 70 °C for 3 h to remove metal catalyst impurities and yield oxygenated functional groups on the surface. The CNTs were filtered with a 0.45 μm PTFE membrane and washed with deionized water until the pH value of the supernatant was close to 7. The resultant CNTs were redispersed into ethanol for further utilization.

Material characterization: The MnO₂-NFs were characterized by field emission scanning electron microscopy (SEM, Carl Zeiss Sigma VP, equipped with scanning transmission electron microscopy (STEM)), energy dispersive X-ray spectroscopy (EDS), transmission electron microscopy (TEM, Zeiss LEO 900), X-ray powder diffraction (XRD, Rigaku Miniflex II with Cu K α radiation), Raman spectroscopy (Bruker SENTERRA, 785 nm laser as excitation source), porosimetry analyzer (Micromeritics ASAP 2020 Analyzer, nitrogen adsorption at 77K), and Fourier transform infrared spectroscopy (FTIR, Thermal Nicolet Nexus 470 spectrometer).

Electrochemical measurements: All electrochemical measurements were conducted using a Gamry reference 600 Potentiostat/Galvanostat/ZRA workstation. For three-electrode testing, platinum gauze and Ag/AgCl were used as the counter and reference electrodes, respectively. The working electrodes of MnO₂-NFs or CNTs were fabricated as below: The as-prepared

MnO₂-NFs (70 wt.%) were evenly mixed and grinded with carbon black (20 wt.%) and polyvinylidene fluoride (PVDF, 10 wt.%) in ethanol and the formed slurry was pasted onto a graphite paper disk (19 mm in diameter). The CNT electrode was prepared by pasting a mixture of CNTs and PVDF (90:10 in weight ratio) onto graphite fiber paper disk. They were pre-dried in a vacuum oven at 70 °C overnight. The graphite fiber papers are chosen as current collectors to prevent capacitance fading from delamination issue and an undesirable electrochemical corrosion towards traditional metallic foils. The porous characteristic of them allows heavy depositions of electroactive materials without penalizing the rate capacity of ASCs to deliver sufficient power for practical applications. The specific capacitance in this work is obtained after excluding the slight contribution of graphite fiber paper.

Based on CV measurements, the charges (q) stored on both electrodes were balanced according to $q = C_s \times \Delta V \times m$, where C_s is the specific capacitance, ΔV is the potential window, and m is the mass of active electrode material.

To build our ASCs, the optimal mass ratio between the MnO₂-NFs and CNTs was calculated to be 0.48 based on the relationship of $q^+ = q^-$ with respective C_s of MnO₂-NFs and CNTs as 145.4 and 70.3 F g⁻¹, respectively, at 20 mV s⁻¹. The actual loading amount for positive and negative materials on graphite current collector is 0.62 mg and 1.02 mg including the additives, respectively. The MnO₂-NFs//CNTs ASCs separated by an electrolyte-cast glassy fiber separator (Millipore Co., 0.45 μm pore size, 19 mm in diameter) were then assembled into a stainless steel split flat cell (MTI Co.) in an Ar-filled glove box. The 2 M KCl aqueous solution was purged with high purity argon for 30 min for oxygen removal and used as the electrolyte. Chloride electrolyte was chosen over conventional sulfate electrolytes deliberately to improve the mobility of cations due to the relatively small ionic radius of chloride anions (ionic radii 1.67 Å of Cl⁻ vs 2.30 Å of SO₄²⁻) in order to further reduce the kinetic limitations. The MnO₂-NFs//MnO₂-NFs and CNTs//CNTs symmetric supercapacitors were constructed following the same procedure.

The specific capacitance calculated from the CV curves can be derived from:

$$C_s = \frac{\int_{V_2}^{V_1} i(V) dV}{2 \times (V_2 - V_1) \times m \times \nu}$$

where C_s is the specific capacitance from evaluated samples; V_1 and V_2 are the starting and ending points of potential window, respectively; $i(V)$ is the instantaneous current as the function of potential; m is the mass of active electrode materials, and is the total mass of both electrode materials for ACS device and two symmetric devices; and ν is the scan rate in mV s⁻¹. The mass is measured using A&D HR-202i analytical balance with readability of 0.01 mg.

The specific capacitance calculated from the galvanostatic charge/discharge measurements can be derived from:

$$C_s = \frac{I \times t}{\Delta V \times m}$$

I and t are the discharge current and the discharge time, respectively in the potential range ΔV excluding the IR drop.

The energy density E at different power density P of the MnO_2 -NFs//CNTs asymmetric supercapacitor, MnO_2 -NFs// MnO_2 -NFs and CNTs//CNTs symmetric supercapacitors is calculated from the CV curves at different scan rates according to the equation below:

$$E = \frac{C_s \times V^2}{2} = \frac{500 \times C_s \times V^2}{3600} \text{ (Wh / kg)}$$

and power density P is calculated following:

$$P = \frac{E}{t} = \frac{3600 \times E}{t} \text{ (W / kg)}$$

where C_s is the specific capacitance in F g^{-1} of devices at a corresponding scan rate; V is the cell potential and t is the discharge time period in seconds from V to 0 V.

Coulombic efficiency is estimated according to:

$$\eta = \frac{t_D}{t_C} \times 100\%$$

where η is coulombic efficiency, t_D and t_C is the galvanostatic charge and discharge time, respectively.

The K^+ Faradic intercalation/deintercalation reaction of the as-synthesized MnO_2 -NFs is represented below:

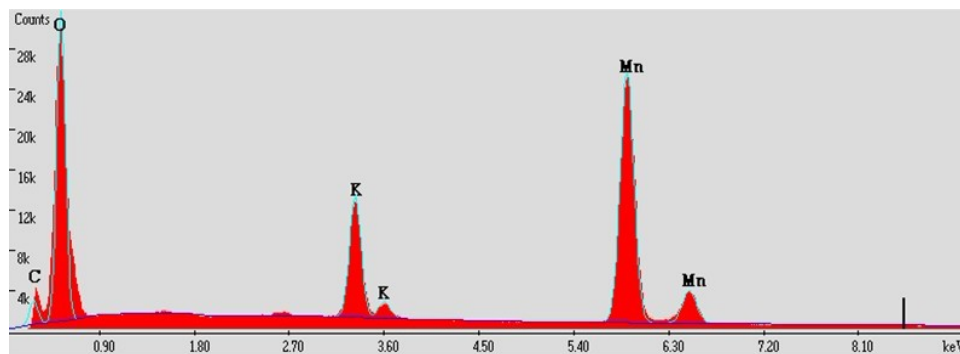
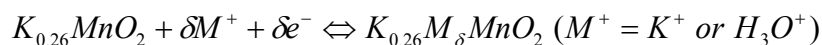


Fig. S1 Energy-dispersive X-ray spectroscopy (EDS) data of as-synthesized MnO_2 -NFs.

In virtue of the strong acidity and oxidizing property of KMnO_4 , we hypothesize that the pyrrolidone ring-opening reaction of the NMP molecules would yield refined MnO_2 nanostructures. In our synthesis, overabundant NMP is used to react with strong oxidizing agent KMnO_4 to synthesize homogeneously distributed MnO_2 nanostructures at relatively low temperature. The high bond strength of C-N (292 kJ mol^{-1}) would give rise to a high reaction activation energy and accordingly brings a significant effect on MnO_2 nucleation and growth rates. Infrared spectra (Fig. S2) were recorded to insightfully investigate synthesis process. The resultant precipitation was first deliberately filtrated and rinsed with water to exclude the influence of residual NMP to the effect that the superficial product compounds on MnO_2 could be detected by IR spectrum and this sample is denoted as filtrated MnO_2 (Fig. S2b).

As shown in Fig. S2, most interesting observation in the IR curve from the filtrated MnO_2 is that the N-C₃ stretching vibration bands, located at 1504 cm^{-1} , of N-C=O in the pyrrolidone ring disappears, whereas the N-C₁ at $\sim 1264 \text{ cm}^{-1}$ and N-C₂ bonds at $\sim 1301 \text{ cm}^{-1}$ have similar chemical environments in comparison with IR spectrum of pure NMP. The other great change is an apparent absorption peak at $\sim 1567 \text{ cm}^{-1}$ corresponding to N-H bending vibration of amines emerges in filtrated MnO_2 specimen. Additionally, the relative intensity difference among the C-H vibrational bands is noticeable, which was regarded as evidence of capping ligands chemisorbing on the MnO_2 nanocrystals. Meanwhile, a remarkable broad band centered at 500 cm^{-1} can be ascribed to the Mn-O vibrations in MnO_6 octahedra. The results demonstrate that the NMP undergoes the N-C₃ breaking of pyrrolidone rings in the presence of KMnO_4 and generate MnO_2 in spite of not eliminating the possibility that the partial NMP might be accompanied with their intrinsic hydrolysis in acidic atmosphere. However, whether the hypothetical by-product, namely 4-(Methylamino)butyric acid, is able to continue being oxidized by KMnO_4 through the breaking of C-N bonds is still under investigation and the likelihood of the occurrence of oxidative azo coupling reaction among the C-NH-C bonds is ruled out due to the absence of acidic catalyst and overabundance of NMP addition.

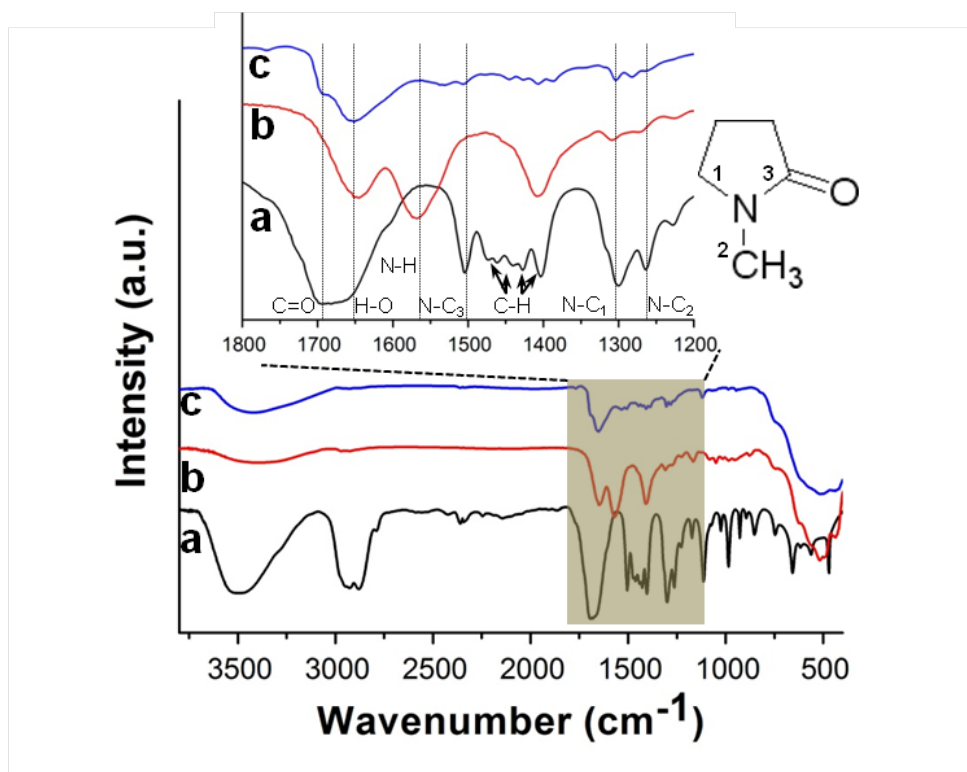


Fig. S2 FTIR spectra of (a) pure NMP, (b) filtrated MnO₂-NFs after DI water rinse, and (c) thoroughly rinsed MnO₂-NFs with DI water and ethanol.

Therefore, the plausible formation mechanism of these monodisperse sub-30 nm MnO₂-NFs is as follows: N-Methyl-2-pyrrolidone (NMP) reduces KMnO₄ to generate MnO₂. The initially formed MnO₂ nuclei experience a rapid growth and fuse into amorphous aggregations. As the reaction proceeds, an Ostwald ripening mechanism takes over the growth. The ultrathin nanoflakes form according to the lamellar crystal structures of birnessite-type MnO₂ and/or anisotropic growth caused by the selective chemisorption of 4-(Methylamino)butyric acid on certain crystal planes, and subsequently self-assemble into flower-like nanostructures. The further growth of these nanoflowers stops after depletion of KMnO₄ in the solution. The possible surface capping by 4-(Methylamino)butyric acid and/or its protection networks by hydrogen bonds between its nitrogen donors may prevent MnO₂-NFs from growing. Comparing to previous studies, the high activation energy for the pyrrolidone ring-opening of NMP is capable of effectively downsizing the formed MnO₂-NFs.

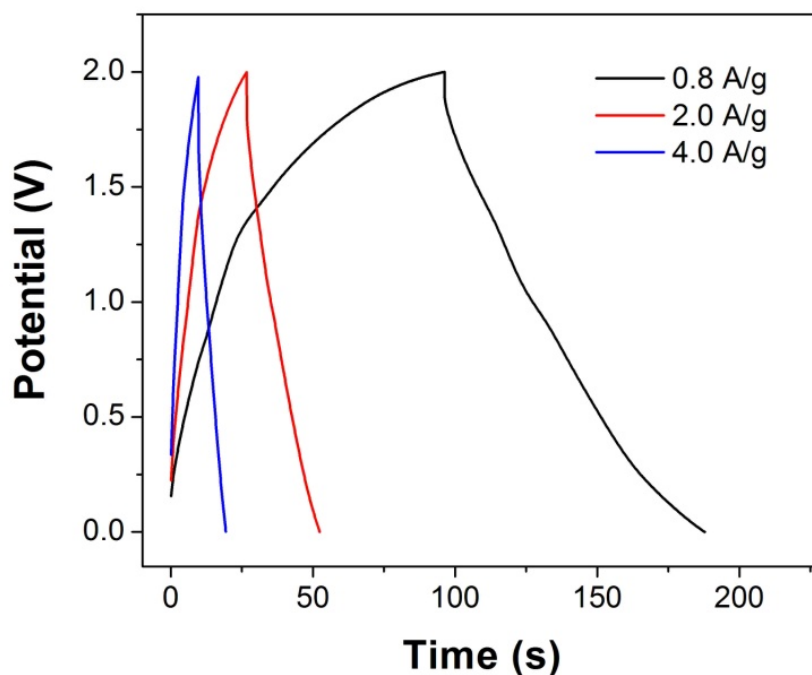


Fig. S3 Galvanostatic charge/discharge curves at different current densities of 0.8, 1.0 and 2.0 A g⁻¹ of the as-assembled ASCs.

Table S1: Summary of Electrochemical Data of MnO₂-based Supercapacitors

Cathode material	Cathode material morphology/size	Anode material	Electrolyte solution	Potential window (V)	Maximum capacitance (F/g)	Energy density (Wh/kg) at P power density (W/kg)	Capacitance retention after n cycles
MnO ₂ ¹	-	AC	KCl	2.0	52	28.8 (P = 500)	(n = 100)
MnO ₂ ²	20-500 nm grains	AC	K ₂ SO ₄	1.5	-	~7.0 (P = 400)	76%(n = 23,000)
MnO ₂ ³	~ 200 nm grains	AC	K ₂ SO ₄	2.2	31	17.3 (P = 605)	~ 87% (n = 5,000)
MnO ₂ ⁴	-	AC	K ₂ SO ₄	2.0	21±2	11.7 (-)	87.5% (n = 195,000)
δ-K _{0.27} MnO ₂ ⁵	50 – 200 nm aggregated particles	AC	K ₂ SO ₄	1.8	57.7	25.3 (P = 140)	≥98% (n = 10,000)
NaMnO ₂ ⁶	50 μm aggregated spherical particles	AC	Na ₂ SO ₄	1.9	38.9	19.5 (P = 130)	≥97% (n = 10,000)
δ-MnO ₂ ⁷	400-600 nm corona structures	AC	Na ₂ SO ₄	2.0	30.8	17.1 (P = 100)	-
α-MnO ₂ ⁸	3D mesoporous frameworks	AC	Na ₂ SO ₄	1.8	23.1	10.4 (-)	- (n = 1,200)
ε-MnO ₂ ⁹	Aligned nanoplates on nickel foam	Graphene hydrogel	Na ₂ SO ₄	2.0	26.8	23.2 (P = 1,000)	83.4% (n = 5,000)
δ-MnO ₂ ¹⁰	≤10 nm in diameter nanorods	AC	K ₂ SO ₄	1.8	53.7	17 (P = 2,000)	≥94% (n = 23,000)
MnO ₂ ¹¹	~ 50 nm nanoparticles	Graphene	Na ₂ SO ₄	2.0	37	25.2 (P = 100)	96% (n = 500)

$K_{0.15}MnO_2$ /PEDOT ¹²	Nanosheets	PEDOT	Na_2SO_4	1.2	77.2	15.4 (-)	87.8% (n = 1,000)
α - MnO_2 /CNTs mixture ¹³	-	AC	KNO_3	2.0	140	21 (-)	88% (n = 1,000)
α - MnO_2 /CNTs mixture ¹⁴	-	PANI/CNTs composite	KNO_3	1.2	-	5.86 (-)	-
α - MnO_2 /CNTs mixture ¹⁴	-	PPy/CNTs composite	H_2SO_4	1.4	-	7.37 (-)	-
α - MnO_2 /CNTs mixture ¹⁴	-	PEDOT /CNTs	KNO_3	1.8	-	13.5 (-)	91% (n = 500)
α - MnO_2 /graphene ¹⁵	10-20 nm in diameter nanowires on graphene	Graphene	Na_2SO_4	2.0	31	30.4 (P = ~ 100)	~ 79% (n = 1,000)
MnO_2 /GHCS ¹⁶	20-50 nm in length nanofibers on GHCS	GHCS	Na_2SO_4	2.0	24.5	22.1 (P = 7,000)	99% (n = 1,000)
MnO_2 /graphene ¹⁷	Nanorods on graphene	Graphene/Ag	Na_2SO_4	1.8	112.8	50.8 (P = 101.5)	-
MnO_2 /graphene ¹⁸	Nanoparticles on graphene	ACN	Na_2SO_4	1.8	113.5	51.1 (P = 198)	97% (n = 1,000)
$K_{0.26}MnO_2$ (this work)	~ 25 nm nanoflake assemblies	CNTs	KCl	2.0	81.7	45.5 (P = 163.5)	No definite loss (n = 20,000)

Abbreviations: activated carbon (AC), polyaniline (PANI), polypyrrole (Ppy), poly(3,4-ethylenedioxythiophene) (PEDOT), stainless steel (SS), carbon nanotubes (CNTs), graphitic hollow carbon spheres (GHCS). To identify easily, the monocomponent MnO_2 -based ASCs are red-labeled and the hybrid MnO_2 -based ASCs are blue-labeled.

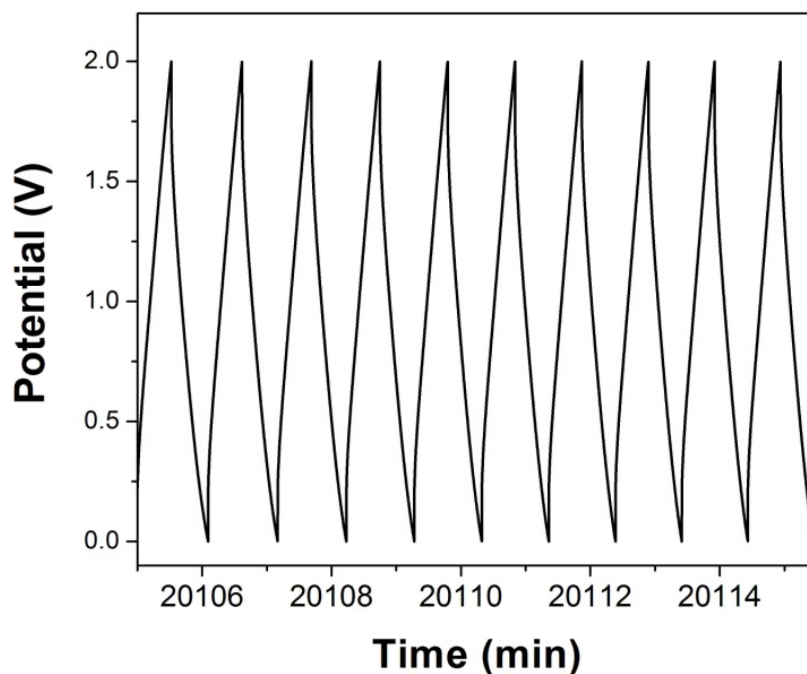


Fig. S4 Galvanostatic charge/discharge curves of the last 10 cycles out of 20,000 cycles at a current density of 2.0 A g^{-1} of the as-assembled ASCs.

Reference

1. M. S. Hong, S. H. Lee and S. W. Kim, *Electrochem. Solid. St.*, 2002, **5**, A227-A230.
2. T. Brousse, M. Toupin and D. Belanger, *J. Electrochem. Soc.*, 2004, **151**, A614-A622.
3. T. Cottineau, M. Toupin, T. Delahaye, T. Brousse and D. Belanger, *Appl. Phys. a-Mater.*, 2006, **82**, 599-606.
4. T. Brousse, P. L. Taberna, O. Crosnier, R. Dugas, P. Guillemet, Y. Scudeller, Y. Zhou, F. Favier, D. Belanger and P. Simon, *J. Power. Sources*, 2007, **173**, 633-641.
5. Q. T. Qu, L. Li, S. Tian, W. L. Guo, Y. P. Wu and R. Holze, *J. Power Sources*, 2010, **195**, 2789-2794.
6. Q. T. Qu, Y. Shi, S. Tian, Y. H. Chen, Y. P. Wu and R. Holze, *J. Power Sources*, 2009, **194**, 1222-1225.
7. X. Zhang, P. Yu, H. T. Zhang, D. C. Zhang, X. Z. Sun and Y. W. Ma, *Electrochim. Acta*, 2013, **89**, 523-529.
8. Y. T. Wang, A. H. Lu, H. L. Zhang and W. C. Li, *J. Phys. Chem. C*, 2011, **115**, 5413-5421.
9. H. C. Gao, F. Xiao, C. B. Ching and H. W. Duan, *ACS Appl. Mater. Inter.*, 2012, **4**, 2801-2810.
10. Q. T. Qu, P. Zhang, B. Wang, Y. H. Chen, S. Tian, Y. P. Wu and R. Holze, *J. Phys. Chem. C*, 2009, **113**, 14020-14027.
11. J. Cao, Y. Wang, Y. Zhou, J.-H. Ouyang, D. Jia and L. Guo, *J. Electroanal. Chem.*, 2013, **689**, 201-206.
12. M. Yeager, W. X. Du, R. Si, D. Su, N. Marinkovic and X. W. Teng, *J. Phys. Chem. C*, 2012, **116**, 20173-20181.
13. V. Khomenko, E. Raymundo-Pinero and F. Beguin, *J. Power Sources*, 2006, **153**, 183-190.
14. V. Khomenko, E. Raymundo-Pinero, E. Frackowiak and F. Beguin, *Appl. Phys. A-Mater.*, 2006, **82**, 567-573.
15. Z. S. Wu, W. C. Ren, D. W. Wang, F. Li, B. L. Liu and H. M. Cheng, *ACS Nano*, 2010, **4**, 5835-5842.
16. Z. B. Lei, J. T. Zhang and X. S. Zhao, *J. Mater. Chem.*, 2012, **22**, 153-160.
17. Y. L. Shao, H. Z. Wang, Q. H. Zhang and Y. G. Li, *J. Mater. Chem. C*, 2013, **1**, 1245-1251.
18. Z. J. Fan, J. Yan, T. Wei, L. J. Zhi, G. Q. Ning, T. Y. Li and F. Wei, *Adv. Funct. Mater.*, 2011, **21**, 2366-2375.