

Facile Oxygen Reduction on a Three-Dimensionally Ordered Macroporous Graphitic C₃N₄/Carbon Composite Electrocatalyst**

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One of the greatest stumbling blocks hindering broad applications of fuel cells is the high cost and vulnerability of the platinum catalyst as well as the sluggish oxygen reduction reaction (ORR) on the cathode.^[1] Although Pt alloys or non-noble metals have been developed as substitute catalysts for the ORR,^[2,3] they still suffer from multiple disadvantages, such as low stability under fuel cell conditions, vulnerability to fuel crossover, and harmfulness to the environment.^[4] Thus, the ongoing search for metal-free catalysts for the ORR has attracted much attention. In this regard, nitrogen-containing carbon materials are of particular interest owing to their catalytic activity towards the ORR brought about by nitrogen incorporation, which has been confirmed by both experimental studies^[5–7] and quantum-mechanical calculations.^[8] Among these materials, graphitic carbon nitride (g-C₃N₄) is especially promising because of its high nitrogen content, low cost, and easily tailorable structure,^[9] which makes it potentially suitable for oxygen reduction as well as other applications, such as photocatalysis^[10] or hydrogen storage, under mild conditions.^[11]

However, the ORR electrocatalytic activity of g-C₃N₄ alone is still inferior to Pt catalysts.^[12,13] This is because of the extremely low electrical conductivity (ca. 10^{−9} S m^{−1})^[10,14] of g-C₃N₄ as indicated by our recent theoretical calculations.^[15] With the aim to better utilize this nitrogen-rich but poorly conducting material, a variety of carbon materials have been recently introduced into g-C₃N₄ by different routes,

including mechanical mixing of g-C₃N₄ with carbon black^[16] or in situ immobilization of g-C₃N₄ onto carbon black,^[17] graphene,^[13,18] or mesoporous carbon.^[15] Compared to pristine g-C₃N₄, the aforementioned composites show better catalytic performance for ORR (that is, higher cathodic reaction current or lower ORR starting potential in a half-cell test). However, the effect of the structure of these composites on the catalytic properties is still unknown.

The porosity of catalyst in a fuel cell is critical for mass transport and access of the proton exchange ionomers.^[19,20] However, the pores of previously investigated g-C₃N₄/carbon materials (g-C₃N₄/C) arise from pyrolysis of carbon precursors,^[16,17] random stacking of graphene layers,^[13,18] or mesoporous carbon with small pores.^[15] The pore sizes of these composites are hard to control and often too narrow for efficient transport and access.^[19,20] Furthermore, the potential prospects of g-C₃N₄/C as a substitute for Pt cannot be forecast on the basis of the current research because the ORR catalytic activity of the existing g-C₃N₄/C is hardly comparable to Pt^[16–18] and/or they are not stable enough in the fuel cell environments.^[13]

From this point of view, a more uniform structure with suitable pore size of the carbon substrate could be preferable for hybrid g-C₃N₄/C fuel cell catalysts. More specifically, a well-defined and continuous porous structure would facilitate reactant transport inside the pores, assure better contact between catalyst and ionomer, and better stability, as well as create an opportunity for an accurate evaluation of the structural effect of the added carbon on the ORR catalytic properties of g-C₃N₄/C. On this basis, a hard template method using silica microspheres as templates can be especially feasible and advantageous for preparing the carbon supports for g-C₃N₄ in several aspects. Preparation of g-C₃N₄/C with tunable pore size from meso to macropores could be easily achieved using this method.^[21–27] Besides, the carbon materials thus synthesized feature uniform cage-like interconnected pores, which not only facilitate the mass transport but also avoid possible confusion in assessing the effect of pore sizes due to structural differences. Furthermore, this three-dimensionally interconnected structure would be stronger than the microscopically separated carbon materials, and the silica spheres are either commercially available or easy to fabricate, making the resulting g-C₃N₄/C potentially favorable for commercialization at low cost.

Herein we present the design and preparation of macroporous g-C₃N₄/C with three-dimensionally ordered interconnected structures using silica microspheres as hard templates. The main objective is to develop a high performance metal-free fuel cell catalyst, to evaluate the structural effect of the

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resulting $\text{g-C}_3\text{N}_4/\text{C}$ on the electrocatalytic activity as well as to reveal its potential suitability as a substitute for Pt/C used in fuel cells. The catalytic activity of $\text{g-C}_3\text{N}_4/\text{C}$ for ORR has been measured in comparison with mesoporous $\text{g-C}_3\text{N}_4/\text{C}$ and commercial Pt/C. The $\text{g-C}_3\text{N}_4/\text{C}$ catalysts with ordered pores of about 150 nm showed highest ORR performance, which was comparable with Pt/C in both the reaction current and the initial potential. Moreover, unlike the vulnerable commercial Pt/C, our material had extremely high tolerance against fuel crossover. Furthermore, this macroporous $\text{g-C}_3\text{N}_4/\text{C}$ catalyst also possessed an excellent durability compared to both commercial Pt/C and previously reported $\text{g-C}_3\text{N}_4/\text{C}$ in alkaline media.^[13,18] To our knowledge, all of these properties have not been achieved in other metal-free catalysts, which make the current macroporous $\text{g-C}_3\text{N}_4/\text{C}$ catalyst promising for the next generation of alkaline fuel cells based on organic fuels.

Macroporous $\text{g-C}_3\text{N}_4/\text{C}$ was prepared as shown in Figure 1 a. Silica spheres with different sizes were synthesized by

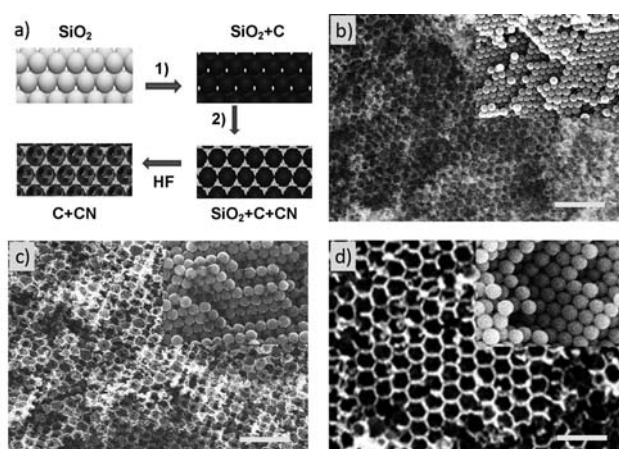


Figure 1. a) The synthesis of macroporous $\text{g-C}_3\text{N}_4/\text{C}$ as a reversal of the SiO_2 sphere structure. Conditions: 1) Sucrose, 900°C , N_2 ; 2) cyanamide, 550°C , N_2 . b)–d) Scanning electron microscopy (SEM) images of 150-C/CN, 230-C/CN, and 400-C/CN (and the corresponding silica spheres of each size (insets)). Scale bars: $1\ \mu\text{m}$.

Stober's method^[28] and packed up by gravitational sedimentation or solvent vaporization, forming an opal monolith. The monolith was then dried and calcined at 550°C to obtain interconnected silica spheres. In the next step, a thin carbon shell was coated on the silica spheres (C@SiO_2) through sucrose infiltration followed by carbonization at 900°C in nitrogen (Supporting Information, Figure S1). Afterward, melted cyanamide, as $\text{g-C}_3\text{N}_4$ precursor, was impregnated into the intersphere spaces of C@SiO_2 , followed by heating at 550°C in nitrogen to form $\text{g-C}_3\text{N}_4/\text{C@SiO}_2$. Finally, the silica template was removed with hydrofluoric acid after $\text{g-C}_3\text{N}_4$ synthesis, rather than in earlier stages, which successfully prevented the possible structural deterioration and pore blockage. Noticeably, as-synthesized $\text{g-C}_3\text{N}_4/\text{C}$ (denoted as X-C/CN, where X is the pore size) has well-defined interconnected ordered macropores with the same size as the silica spheres (Figure 1).

The formation of $\text{g-C}_3\text{N}_4$ in the composite is confirmed by X-ray diffraction (XRD) on the basis of the peaks at 2θ located at 30° and 52° and corresponding to the (002) and the (101) face (Figure 2 a).^[9] The existence of $\text{g-C}_3\text{N}_4$ on carbon is

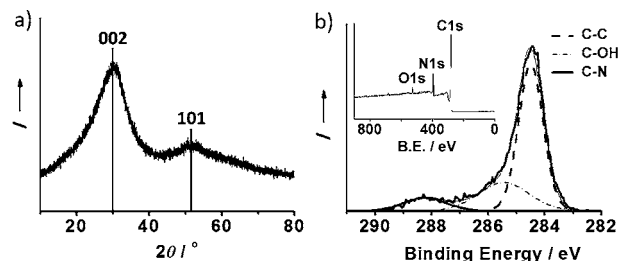


Figure 2. a) Typical XRD pattern obtained by $\text{Co K}\alpha$ X-rays; b) typical high-resolution C1s XPS spectrum. Inset: corresponding survey scan of $\text{g-C}_3\text{N}_4/\text{C}$.

also evidenced by FTIR spectra on the basis of characteristic $\text{g-C}_3\text{N}_4$ absorbance peaks (Supporting Information, Figure S2). X-ray photoelectron spectroscopy (XPS) is another method to probe the integration of $\text{g-C}_3\text{N}_4$ with carbon. The survey scan spectrum of $\text{g-C}_3\text{N}_4/\text{C}$ shows the presence of carbon, nitrogen, and oxygen (inset in Figure 2 b). High resolution C1s spectra reveal the carbon status in the material (Figure 2 b). The emergence of the peak at $288.3\ \text{eV}$ corresponding to C–N coordination and the enhancement of the peak at $284.5\ \text{eV}$ corresponding to C–C coordination, compared to pristine carbon and $\text{g-C}_3\text{N}_4$ (Supporting Information, Figure S3), confirm the successful incorporation of $\text{g-C}_3\text{N}_4$ onto carbon.^[13,16] Another peak at $285.7\ \text{eV}$ corresponds to C–OH, which is from the sucrose precursor and is commonly observed on carbon materials.^[29] High-resolution scans of N1s in $\text{g-C}_3\text{N}_4/\text{C}$ and $\text{g-C}_3\text{N}_4$ show similar nitrogen species, indicating comparable nitrogen status in both materials (Supporting Information, Figure S3). The porosity of these materials was investigated by nitrogen sorption measurements. As a result, all of the macroporous samples possess similar Brunauer–Emmett–Teller (BET) specific surface areas in the range between 50 and $100\ \text{m}^2\text{g}^{-1}$; the surface area tends to decrease with increasing pore size (Supporting Information, Figure S5). No significant presence of mesopores or micropores was observed.

To further evaluate the pore size effect on the ORR catalytic activity of $\text{g-C}_3\text{N}_4/\text{C}$, mesoporous $\text{g-C}_3\text{N}_4/\text{C}$ was also prepared for the purpose of comparison. The fabrication was firstly carried out by synthesizing mesoporous carbon using $12\ \text{nm}$ colloidal silica spheres as a hard template as previously reported.^[24] Cyanamide was then cast into the mesoporous carbon followed by heating at 550°C , forming $\text{g-C}_3\text{N}_4$ inside mesoporous carbon (denoted as 12-C/CN). Both the 12-C/CN and the mesoporous carbon have the similar isotherm hysteresis loop confirming the presence of mesopores, but slight pore shrinkage is observed in 12-C/CN as compared to the parent carbon material (Supporting Information, Figure S6). These results indicate that both mesoporous materials possess similar cage like pores and $\text{g-C}_3\text{N}_4$ impregnation into mesoporous carbon does not significantly change its pore

structure. Transmission electron microscopy (TEM) observation also confirms the similar structures of mesoporous g-C₃N₄/C and carbon (Supporting Information, Figure S6). The uniform mesoporous structure of 12-C/CN is extremely suitable for comparison with its macroporous counterparts under study.

The ORR electrocatalytic activity of g-C₃N₄/C was firstly tested through conventional three-electrode cyclic voltammetry (CV) in O₂ or N₂ saturated 0.1 M KOH aqueous solutions (Figure 3a). A voltammogram without any signifi-

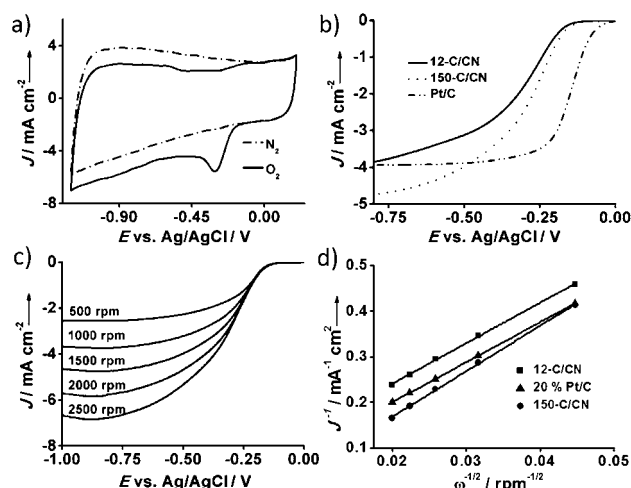


Figure 3. Catalytic activity towards electrochemical reduction of oxygen in 0.1 M KOH aqueous solution at room temperature. a) Cyclic voltammetry performed for 150-C/CN in O₂ and N₂; b) onset potential of mesoporous (12-C/CN), macroporous (150-C/CN) g-C₃N₄/C, and Pt/C obtained with a rotating disk electrode (RDE) at 1500 rpm and 5 mV s⁻¹; c) linear sweep voltammograms (LSVs) obtained for macroporous g-C₃N₄/C (150-C/CN) at various speeds; d) Koutecky-Levich plot for g-C₃N₄/C and Pt/C obtained from LSVs in (c) and the Supporting Information, Figure S7 at -0.6 V.

cant peak was obtained in the absence of oxygen. The quasi-rectangular shape is due to a capacitor effect.^[6,7] On the contrary, when oxygen was introduced, a characteristic ORR peak starting at -0.177 V was observed, showing the electrochemical reduction of oxygen initiated on g-C₃N₄/C. Compared to previous reports on nitrogen-doped carbon materials, the starting potential has increased positively,^[6,30,31] indicating an easier ORR process on g-C₃N₄/C. At the same time, pure g-C₃N₄ with similar structure was also tested under the same conditions, showing the negligible ORR catalytic activity. The results indicate that the performance enhancement of g-C₃N₄/C was brought about by the introduction of carbon (Supporting Information, Figure S7).

To further investigate the ORR catalytic activity of g-C₃N₄/C and the effect of pore size, these materials and commercial Pt/C (Vulcan, 20 wt %) were tested using a rotating disk electrode (RDE) in O₂-saturated 0.1 M KOH solution. The g-C₃N₄ amounts in each sample on the electrode in the RDE tests were the same by carefully controlling the catalyst ink concentration according to the elemental and thermogravimetric analysis results (Supporting Information, Fig-

ure S8 and Table S2). The ORR onset potentials of each sample were acquired from RDE linear sweep at 1500 rpm (Figure 3b). Remarkably, the macroporous g-C₃N₄/C showed obviously better performance than the mesoporous material in both onset potential (ca. -0.14 V) and reaction current density, which is comparable to Pt/C and indicates better performance brought about by larger pores.

Linear sweep voltammograms (LSV) were also recorded from 500 to 2500 rpm. LSVs for all g-C₃N₄/Cs show typical increasing current with higher rotations speeds (Figure 3c; Supporting Information, Figure S9). This result can be explained by shortened diffusion distance at high speeds, which is in accordance with other studies.^[6,7] Noticeably, 12-C/CN showed a simultaneously increasing current density along with overpotential at all rotation speeds (Supporting Information, Figure S8), indicating a surface dominant reaction without a diffusion limited current, which is possibly due to the difficulties in reactant transfer within narrow 12 nm mesopores. The slight decrease in the current density at high rotation (2500 rpm) and high overpotential (> 0.9 V) might be caused by the fast O₂ consumption in the test cell.

The Koutecky-Levich plots (j^{-1} vs. $\omega^{-1/2}$) of each catalyst are obtained from LSVs according to the current at -0.6 V, and all plots show good linearity at various rotation speeds (Figure 3d; Supporting Information, Figure S8). The macroporous samples exhibit higher current density than the mesoporous sample, and the 150-C/CN possesses the highest current density among all of the samples, which is even higher than that obtained for Pt/C at all rotation speeds (Figure 3b,d; Supporting Information, Figure S9), revealing the outstanding catalytic activity of this macroporous g-C₃N₄/C. The electron transfer numbers of ORR on different g-C₃N₄/C samples can be obtained from the slope of Koutecky-Levich plots (see details in the Supporting Information; these numbers are listed in Table S1).^[32] All of the g-C₃N₄/C samples possess a similar value of about 3, indicating a reasonably similar oxygen reduction process with combined two-electron and four-electron reaction pathways.^[32]

To obtain insight into the ORR catalytic kinetic differences on g-C₃N₄/C with different pore sizes, the Tafel slopes, representing the overall resistance in the ORR process, were obtained according to the linear plots of LSVs at 1500 rpm for all g-C₃N₄/Cs (Figure 4a; Supporting Information, Figure S10). All of the plots show typical two-stage linear regions at low overpotential (> -0.2 V) and high overpotential (< -0.25 V), and the slope values are listed in Figure 4b.

In the low-overpotential region, where the overall ORR speed is determined by the surface reaction rate on the catalyst,^[33] all of the Tafel slopes were between 51.1 and 72.6 mV dec⁻¹. The comparable values of the Tafel slopes for g-C₃N₄/C are also in agreement with their similar electron transfer numbers. This is in accordance with the fact that all g-C₃N₄/C can electrochemically reduce oxygen by a similar oxygen reduction process mechanism combining two- and four-electron reaction pathways. In the high overpotential region, where the overall ORR rate is dependent on the oxygen diffusion, the Tafel slope values for macroporous g-C₃N₄/Cs are between 150 and 190 mV dec⁻¹, which are much smaller than that of 12-C/CN (308 mV dec⁻¹), confirming

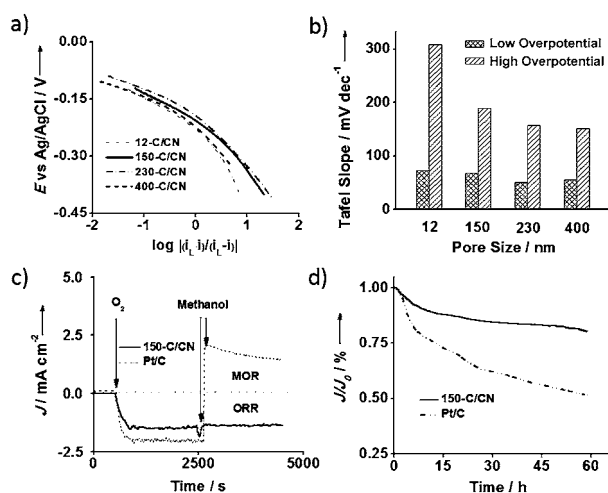


Figure 4. a) Tafel plots obtained from the RDE measurements on g-C₃N₄/C at 1500 rpm and b) the corresponding slope values of low and high overpotential regions; c) current–time chronoamperometric response of 150-C/CN and Pt/C in 0.1 M KOH solution. The arrow indicates the introduction of O₂ and 10 vol % methanol; d) current–time chronoamperometric response of 150-C/CN and Pt/C in O₂-saturated 0.1 M KOH solution.

significantly smoother reactant diffusion in the macropores than that in mesopores. As all of the samples show similar ORR mechanism at low overpotential, it is reasonable to assume that the better ORR activity of macroporous g-C₃N₄/C, compared to the mesoporous sample, is caused by the facile mass transfer in macropores. The superior performance of 150-C/CN could be the combination of both the positive effect of macropores on diffusion and the sufficient amount of active sites linked to the highest BET surface area among all macroporous samples.

For a new ORR electrocatalyst for fuel cells, a high catalytic selectivity for cathode reactions against fuel oxidation is important, especially when using small-molecule organic fuels, such as methanol, which could cross over through the polymer electrolyte membrane from anode to cathode, seriously compromising the whole cell performance.^[13,31] The methanol crossover effect was evaluated on both macroporous g-C₃N₄/C and commercial Pt/C (Figure 4c). A negative current appeared when oxygen was introduced to N₂-saturated 0.1 M KOH aqueous solution at about 500 s, indicating ORR occurred on both g-C₃N₄/C and Pt/C. After adding 10 vol % methanol to the electrolyte, g-C₃N₄/C retained stable current response whereas the current in the Pt/C system instantaneously jumped to positive values owing to the methanol oxidation reaction (MOR) on Pt/C. These results clearly show the excellent catalytic selectivity of g-C₃N₄/C and its better suitability as a cathode catalyst for direct methanol fuel cells than the vulnerable Pt/C.

The durability of g-C₃N₄/C and Pt/C was also assessed through chronoamperometric measurements at −0.3 V; at this potential the current was the largest, indicating the highest reaction rate (Figure 4d). The 100 h test only caused slight activity loss (reaction current decrease) on g-C₃N₄/C, whereas Pt/C lost nearly 50 % of its initial activity in just 60 h,

confirming a much better stability of active reaction sites on the g-C₃N₄/C materials than that on commercial Pt/C in alkaline environment, which is favored for the next generation alkaline fuel cells.

Taking into account the similar composition of the g-C₃N₄/C catalysts in this study and previous reports,^[13] the observed enhancement in the ORR durability on the macroporous g-C₃N₄/C can be attributed to their uniquely porous structure. The composite catalyst synthesized in our study have a three-dimensionally interconnected porous structure, which effectively prevent the structural collapse or catalyst detachment that happens on microscopically separated carbon supported Pt or g-C₃N₄ catalysts, and thus help to keep our g-C₃N₄/C far more stable than other g-C₃N₄/C materials or commercial Pt/C.^[13]

In summary, we have successfully designed and constructed a three-dimensionally ordered macroporous g-C₃N₄/C catalyst with outstanding ORR performance for fuel cells. This novel catalyst possesses prominent ORR catalytic activity, which is comparable with commercial Pt/C in both reaction current density and onset potential. As a metal-free catalyst, the macroporous g-C₃N₄/C showed much better fuel crossover resistance and long-term durability than the commercial Pt/C in alkaline medium. Furthermore, this material was synthesized through a simple procedure using inexpensive cyanamide as a precursor and easily fabricated silica microspheres as a template, which gives a great promise for large scale production. The pore size of g-C₃N₄/C can be easily tailored by using different size silica spheres as templates. All these features make macroporous g-C₃N₄/C a potentially promising and suitable substitute for the expensive noble metal catalysts in the next generation alkaline and direct methanol fuel cells.

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- [1] B. C. H. Steele, A. Heinzl, *Nature* **2001**, 414, 345.
- [2] R. Bashyam, P. Zelenay, *Nature* **2006**, 443, 63.
- [3] H. Gasteiger, N. Markov, *Science* **2009**, 324, 48.
- [4] D. S. Yu, E. Nagelli, F. Du, L. M. Dai, *J. Phys. Chem. Lett.* **2010**, 1, 2165.
- [5] K. Gong, F. Du, Z. Xia, M. Durstock, L. Dai, *Science* **2009**, 323, 760.
- [6] W. Yang, T.-P. Fellingner, M. Antonietti, *J. Am. Chem. Soc.* **2011**, 133, 206.
- [7] R. Liu, D. Wu, X. Feng, K. Müllen, *Angew. Chem.* **2010**, 122, 2619; *Angew. Chem. Int. Ed.* **2010**, 49, 2565.
- [8] S. Maldonado, K. J. Stevenson, *J. Phys. Chem. B* **2005**, 109, 4707.
- [9] S. Hwang, S. Lee, J.-S. Yu, *Appl. Surf. Sci.* **2007**, 253, 5656.
- [10] X. Wang, K. Maeda, A. Thomas, K. Takanabe, G. Xin, J. M. Carlsson, K. Domen, M. Antonietti, *Nat. Mater.* **2009**, 8, 76.
- [11] S. S. Park, S. W. Chu, C. Xue, D. Zhao, C. S. Ha, *J. Mater. Chem.* **2011**, 21, 10801.
- [12] K. Kwon, Y. J. Sa, J. Y. Cheon, S. H. Joo, *Langmuir* **2012**, 28, 991.

- [13] S. Yang, X. Feng, X. Wang, K. Müllen, *Angew. Chem.* **2011**, *123*, 5451; *Angew. Chem. Int. Ed.* **2011**, *50*, 5339.
- [14] Y. Zhang, T. Mori, J. Ye, M. Antonietti, *J. Am. Chem. Soc.* **2010**, *132*, 6294.
- [15] Y. Zheng, Y. Jiao, J. Chen, J. Liu, J. Liang, A. Du, W. Zhang, Z. Zhu, S. Smith, M. Jaroniec, G. Q. M. Lu, S. Z. Qiao, *J. Am. Chem. Soc.* **2011**, *133*, 20116.
- [16] S. M. Lyth, Y. Nabae, S. Moriya, S. Kuroki, M.-a. Kakimoto, J.-i. Ozaki, S. Miyata, *J. Phys. Chem. C* **2009**, *113*, 20148.
- [17] S. M. Lyth, Y. Nabae, N. M. Islam, S. Kuroki, M. Kakimoto, S. Miyata, *J. Electroanal. Chem.* **2011**, *158*, B194.
- [18] Y. Sun, C. Li, Y. Xu, H. Bai, Z. Yao, G. Shi, *Chem. Commun.* **2010**, *46*, 4740.
- [19] E. Antolini, *Appl. Catal. B* **2009**, *88*, 1.
- [20] H. Chang, S. H. Joo, C. Pak, *J. Mater. Chem.* **2007**, *17*, 3078.
- [21] S. H. Joo, S. J. Choi, I. Oh, J. Kwak, Z. Liu, O. Terasaki, R. Ryoo, *Nature* **2001**, *412*, 169.
- [22] A. Lu, A. Kiefer, W. Schmidt, F. Schuth, *Chem. Mater.* **2004**, *16*, 100.
- [23] J. Fan, C. Yu, L. Wang, B. Tu, D. Zhao, Y. Sakamoto, O. Terasaki, *J. Am. Chem. Soc.* **2001**, *123*, 12113.
- [24] H. I. Lee, G. D. Stucky, J. H. Kim, C. Pak, H. Chang, J. M. Kim, *Adv. Mater.* **2011**, *23*, 2357.
- [25] Y. Meng, D. Gu, F. Zhang, Y. Shi, H. Yang, Z. Li, C. Yu, B. Tu, D. Zhao, *Angew. Chem.* **2005**, *117*, 7215; *Angew. Chem. Int. Ed.* **2005**, *44*, 7053.
- [26] Y. Deng, T. Yu, Y. Wan, Y. Shi, Y. Meng, D. Gu, L. Zhang, Y. Huang, C. Liu, X. Wu, D. Zhao, *J. Am. Chem. Soc.* **2007**, *129*, 1690.
- [27] C. Liang, S. Dai, *J. Am. Chem. Soc.* **2006**, *128*, 5316.
- [28] W. Stober, A. Fink, E. Bohn, *J. Colloid Interface Sci.* **1968**, *26*, 62.
- [29] H. Darmstadt, C. Roy, S. Kaliaguine, S. J. Choi, R. Ryoo, *Carbon* **2002**, *40*, 2673.
- [30] S. H. Liu, J. R. Wu, *Int. J. Hydrogen Energy* **2011**, *36*, 87.
- [31] L. Qu, Y. Liu, J. B. Baek, L. Dai, *ACS Nano* **2010**, *4*, 1321.
- [32] H. S. Wroblowa, P. Yen Chi, G. Razumney, *J. Electroanal. Chem. Interfacial Electrochem.* **1976**, *69*, 195.
- [33] M. T. de Groot, M. Merkx, A. H. Wonders, M. T. M. Koper, *J. Am. Chem. Soc.* **2005**, *127*, 7579.