High surface area, micro/mesoporous carbon particles with selectable 3-D biogenic morphologies for tailored catalysis, filtration, or adsorption†

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Biogenic 3-D SiO2 micro-assemblies (diatom microshells) of low specific surface area (SSA, <2 m2g−1) have been transformed into 3-D replicas of high SSA C (>1300 m2g−1) and Pt-loaded C (>600 m2g−1) by a shape-preserving serial reaction process. Such high SSA micro-assemblies may be generated in a wide variety of selectable 3-D shapes for tailored catalysis, filtration, or adsorption for energy storage/production, water purification, and biofuel separation.

Highly-porous carbon assemblies have attracted appreciable interest as catalyst supports in fuel cells or sensors, as electrodes for supercapacitors or batteries, and as adsorbants for hydrogen or natural gas storage, carbon dioxide capture, and (bio)chemical or water purification.1 Templating approaches have been used to generate very high surface area (>1,000 m2g−1) porous carbon structures with simple shapes (e.g., spheres, tubes, films) or with complex but uncontrolled irregular shapes (e.g., non-uniform powders).2 Here we demonstrate a shape-preserving serial reaction process for converting complex-shaped, three-dimensional (3-D) biogenic SiO2 micro-assemblies (i.e., the microshells, or frustules, of single-celled aquatic algae known as diatoms) of low specific surface area (SSA <2 m2g−1) into positive 3-D replicas comprised of micro/mesoporous C of very high SSA (>1,300 m2g−1). To our knowledge, this is the first report of a method capable of yielding such high-SSA carbon assemblies in a wide variety of complex 3-D microscale morphologies available from biogenic3 or synthetic4 templates (e.g., each of the tens of thousands of extant diatom species forms a silica frustule with a unique 3-D morphology5). While diatom frustules have been used as passive (non-reacting) scaffolds for the infiltration of C-bearing precursors (sucrose, furfuryl alcohol6), subsequent carbonization heat treatment and SiO2 dissolution has yielded negative C replicas with much lower values of SSA (169–312 m2g−1) than observed for the reaction-induced positive replicas of the present work. The ability to synthesize microscale high-SSA carbon assemblies with selectable 3-D morphologies can provide for enhanced control of adsorption, filtration, or catalysis. Indeed, upon loading with Pt nanoparticles, the 3-D high-SSA C frustule replicas of the present work exhibited significantly higher electrocatalytic activity for oxygen reduction (under conditions relevant for polymer electrolyte fuel cells) than irregular particles of Pt-bearing commercial C black and SiC-derived C of comparable SSA.

† Electronic supplementary information (ESI) available: Experimental details on the syntheses and characterization of C replicas, along with Figures S1–S3. See DOI: 10.1039/c1ee02102h
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Broader context

The electrical, chemical, and mechanical characteristics of carbon have made this material attractive for energy storage/harvesting, environmental remediation, and (bio)chemical separation (e.g., as electrodes for batteries or supercapacitors; catalyst supports in fuel cells or sensors; adsorbants for hydrogen or natural gas storage, carbon sequestration, water purification, and biofuel separation). The performance of carbon is strongly dependent upon the carbon morphology, specific surface area (SSA), and pore size distribution. To date, high SSA (>1,000 m2g−1) carbon has been produced in simple regular shapes (spheres, tubes, films) or uncontrolled irregular shapes (non-uniform powders). Herein, we demonstrate, for the first time, a bio-enabled process for synthesizing micro/mesoporous carbon of very high SSA (>1,300 m2g−1) and with complex, but controlled three-dimensional (3-D) morphologies inherited from the silica microshells of diatoms (unicellular algae). The low SSA (<2 m2g−1) silica microshells are transformed into 3-D carbon replicas of very high SSA (>1,300 m2g−1) by a series of shape-preserving reactions. Upon loading with Pt nanoparticles, the 3-D high-SSA C microshell replicas exhibited significantly higher oxygen reduction activity (e.g., for fuel cells) than irregular Pt-bearing C particles of comparable SSA. High-SSA carbon assemblies with a wide range of 3-D architectures selected from biogenic silica templates may be generated for tailored catalysis, filtration, intercalation, or adsorption in environmental or energy-related applications.
A series of shape-preserving chemical reactions was used to convert 3-D microscale silica diatom frustules of low SSA into positive 3-D carbon replicas of very high SSA for subsequent platinum loading. The starting *Aulacoseira* diatom frustules (Fig. 1a) possessed a hollow cylindrical shape, side walls with fine channels and rows of macropores (102 nm dia.), and end faces with a protruding rim and a circular hole. Nitrogen adsorption analyses indicated that these SiO2 frustules possessed a SSA of only 1.65 m2g⁻¹, a specific mesopore volume of <0.005 cm3g⁻¹, and no detectable microporosity. The following magnesiothermic reaction was used to convert *Aulacoseira* frustules into MgO/Si replicas:

\[
2\text{Mg(g)} + \text{SiO}_2(s) \rightarrow 2\text{MgO(s)} + \text{Si(s)}
\]  

The magnesia product of this reaction was removed by selective acid dissolution to yield polycrystalline silicon, as confirmed by X-ray diffraction (XRD) analysis (Fig. 2). Exposure of these porous silicon replicas to flowing methane at 950 °C for 2.5 h, followed by thermal treatment at 1200 °C for 12 h in flowing argon, resulted in conversion of the silicon into silicon carbide (β-SiC), as revealed by XRD, transmission electron microscopy (TEM), and selected area electron diffraction (SAED) analyses (Fig. 2, S1a†). The SiC structures were then exposed to Cl2 gas at 950 °C for 2 h to allow for selective Si removal and amorphous carbon formation, as confirmed by XRD, TEM/SAED, and energy-dispersive X-ray (EDX) analyses (Fig. 2, S1b†). The C replicas were then impregnated with Pt nanoparticles via use of platinum dicarboxyl chloride (Pt(CO)2Cl2) vapor (230 °C, 0.5 h), EDX and HRTEM analyses (Fig. 3a, 3b, S1c†) confirmed the presence of Pt nanoparticles within the porous walls of the C replicas. Thermogravimetric analysis (Fig. S2a†) indicated that the Pt-bearing C frustule replicas contained 38.3 wt% platinum.

The 3-D cylindrical morphology and fine features (rows of fine pores and channels) of the starting SiO2 *Aulacoseira* frustules were retained upon reactive conversion into Si, β-SiC, and amorphous C (Fig. 1b–d). However, the porosity within the walls of these frustules replicas was altered by the reaction-induced changes in solid volume.

**Fig. 1** Secondary electron images of: a) a starting SiO2 *Aulacoseira* diatom frustule, b) a Si replica, c) a SiC replica, and d) a C replica.

**Fig. 2** XRD patterns obtained from Si, SiC, and C replicas of *Aulacoseira* diatom frustules.

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Selective dissolution of the MgO product of reaction (1) yielded a porous, but interconnected Si network with enhanced SSA (52.6 m²g⁻¹) and mesopore volume (0.448 cm³g⁻¹) relative to the starting frustules. Subsequent conversion of the porous Si into β-SiC resulted in modest reductions in SSA and mesopore volume (to 48.4 m²g⁻¹ and 0.370 cm³g⁻¹, respectively), which was consistent with the modest difference in the volumes of Si (12.05 cm³mol⁻¹) and β-SiC (12.47 cm³mol⁻¹).18 However, amorphous C possesses a significantly smaller molar volume (~5.2–6.0 cm³mol⁻¹) than β-SiC.19 Hence, since the Cl₂-induced selective removal of Si from the β-SiC frustules replicas yielded amorphous C replicas of similar size (i.e., the β-SiC and C frustule replicas possessed diameters of ~10–11 μm, see Fig. 1C, 1D, and 5), the large reduction in solid volume associated with the conversion of β-SiC into C was accommodated by dramatic enhancements in the values of SSA (1370 m²g⁻¹), mesopore volume (1.53 cm³g⁻¹), and micropore volume (0.282 cm³g⁻¹) for the C frustule replicas (Table 1).

In order to evaluate the utility of these hollow 3-D high-SSA carbon frustule replicas as catalyst templates, the replicas (referred to herein as CF microparticles) were impregnated with platinum nanoparticles. These Pt-bearing CF (Pt/CF) microparticles possessed a SSA of 662 m²g⁻¹, along with appreciable retained mesoporosity (0.689 cm³g⁻¹) and reduced microporosity (0.0584 cm³g⁻¹) relative to the CF microparticles (Table 1). For comparison of catalytic activity, two other types of carbon microparticles were impregnated with platinum: i) porous SiC-derived C (referred to as CS microparticles) generated via exposure of β-SiC powder (8.7 μm ave. dia.) to Cl₂(g) at 950 °C for 2 h, and ii) C black (4.9 μm ave. dia., Vulcan XC-72R, referred to as CV microparticles). The CS microparticles (Fig. S3a†) were selected for comparison to the CF microparticles, as both types of microparticles possessed similar specific values of surface area, mesopore volume, and micropore volume (i.e., these values agreed to within 11%, Table 1) and comparable particle diameters (~8.7 vs. ~10–11 μm). Although the corresponding values of specific surface area, specific micropore volume, and specific mesopore volume for the CV powder (Fig. S3b†) were much smaller, the selection of these CV microparticles was based on the common use of this catalyst support in PEM fuel cells.22 After Pt deposition with Pt(CO)₂Cl₂ vapor (230 °C, 0.5 h), the Pt-bearing CS (Pt/CS) and CV (Pt/CV) microparticles possessed 41.4 wt% and 14.2 wt% Pt, respectively (Figs. S2b, S2c†). That is, after Pt deposition with Pt(CO)₂Cl₂ vapor under identical conditions, a similar amount of Pt (38.3 and 41.4 wt%) was deposited onto the CF microparticles (which possessed similarly high values of SSA), whereas less Pt (14.2 wt%) was deposited onto the modest SSA CV microparticles. The resulting Pt/CF and Pt/CS microparticles also possessed higher specific values of surface area, mesopore volume, and micropore volume than did the C/PTV microparticles (Table 1). Comparison of the micropore volumes before and after Pt deposition indicated a larger reduction in micropore volume for CF microparticles (from 0.282 to 0.0584 cm³g⁻¹ = 79% reduction) than for CS (from 0.286 to 0.170 cm³g⁻¹ = 41% reduction) or CV (from 0.0232 to 0.0215 cm³g⁻¹ = 7.3% reduction) microparticles, which was consistent with a relatively high population of very fine (≤2 nm) Pt nanoparticles in the Pt/CF specimens (Fig. 3b).

The electrocatalytic activities of Pt/CF, Pt/CS, and Pt/CV microparticles for the oxygen reduction reaction (ORR) in an oxygen-saturate 0.5 M H₂SO₄ solution were evaluated via the rotating disk electrode (RDE) method. A suspension of each type of platinum-bearing microparticle in a perfluorosulfonic acid-bearing (Nafion®) solution was deposited onto a glassy carbon working electrode and dried. The microparticle content deposited onto the working electrode was adjusted to achieve a similar amount of total platinum loading (0.1 mgcm⁻²) for each type of microparticle. Polarization curves are shown in Fig. 4a. At potentials ≥0.9 V (where the overall ORR rate was dominated by charge-transfer kinetics), the Pt/CV-bearing working electrode exhibited substantially greater catalytic activity (carried a higher current density) than for the Pt/CS-bearing or Pt/CF-bearing working electrodes. Over the potential range of 0.90 to 1.05 V, the ORR activity of the Pt/CF-bearing electrode (as revealed by the current density) was higher than for the Pt/CS-bearing and Pt/CV-bearing electrodes by factors of 4.5–4.9 and 7.7–9.2, respectively. Electrochemical impedance spectroscopic (EIS) analyses obtained at 0.8 V (i.e., comparable to operating voltages used for PEM fuel cells44) are shown in Fig. 4b. While similar values of Ohmic resistance (given by the high frequency intercepts of the curves in Fig. 4b with the real axis) were seen for the various Pt/C-bearing electrodes, the value of interfacial charge transfer resistance (given by the difference between the high frequency and low frequency intercepts of these curves with the real axis) was appreciably lower for the Pt/CS-bearing electrode than for the Pt/CS-bearing and Pt/CV-bearing electrodes. Chronoamperometric analysis (Fig. 4c) at 0.8 V indicated that the steady-state current for the Pt/CF-bearing electrode was also significantly higher than for the Pt/CS-bearing and Pt/CV-bearing electrodes. It is worth noting that, although the Pt/CS microparticles possessed higher specific micropore and mesopore volumes than did the Pt/CF microparticles (Table 1), the latter microparticles exhibited significantly higher catalytic activity for ORR.

The significantly enhanced electrocatalytic activity of the Pt/CF microparticles was consistent with the presence of a higher population of very fine (≤2 nm) Pt nanoparticles (as discussed above), and a reduced oxygen diffusion distance, for these microparticles than for the Pt/CS and Pt/CV microparticles. Because the Pt/CF microparticles retained the hollow morphology of the diatom frustules, the average oxygen diffusion distance through these porous microparticles was ~0.7 μm (half the wall thickness shown in Fig. 5a and 5b), as opposed to ~4.3 μm and ~2.5 μm (half the average particle diameters) for Pt/CS and Pt/CV microparticles, respectively. The hollow and thin-walled nature of the CF microparticles, coupled with the relatively high specific micropore volume of these particles, also appears to have enabled the deposition of a relatively high population of fine platinum nanoparticles within the CF micropores. Although the CF and CS microparticles possessed similar starting

### Table 1 Specific Surface Area (SSA), Specific Micropore Volume (SMiV), and Specific Mesopore Volume (SMeV)

<table>
<thead>
<tr>
<th>Specimena</th>
<th>SSA (m²g⁻¹)</th>
<th>SMiV (cm³g⁻¹)</th>
<th>SMeV (cm³g⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pt/CF</td>
<td>1370</td>
<td>0.282</td>
<td>1.53</td>
</tr>
<tr>
<td>Pt/CS</td>
<td>1230</td>
<td>0.286</td>
<td>1.40</td>
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<tr>
<td>Pt/CV</td>
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<td>0.0232</td>
<td>0.413</td>
</tr>
<tr>
<td>Pt/CS</td>
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<td>0.0584</td>
<td>0.689</td>
</tr>
<tr>
<td>Pt/CV</td>
<td>734</td>
<td>0.170</td>
<td>0.837</td>
</tr>
<tr>
<td>Pt/CV</td>
<td>141</td>
<td>0.0215</td>
<td>0.183</td>
</tr>
</tbody>
</table>

a CF = C frustule replicas, CS = C replicas of SiC powder; CV = C black (Vulcan XC-72R); Pt/CF = Pt-bearing C frustule replicas; Pt/CS = Pt-bearing SiC powder replicas; Pt/CV = Pt-bearing C black.

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specific micropore volumes (0.282 and 0.286 cm³/g, respectively), the shorter diffusion distance for the CF microparticles appears to have resulted in greater filling of the micropores, and the generation of a larger population of fine platinum nanoparticles, than for the CS microparticles (as indicated by the larger reduction in specific micropore volume after Pt deposition for the CF microparticles than for the CS microparticles). The relatively low population of micropores in the CV microparticles, and the small reduction in the specific micropore volume upon Pt loading of these microparticles, indicated that the population of fine Pt nanoparticles in these microparticles was low compared to the CF or CS microparticles.

This work provides the first demonstration of the use of a series of shape-preserving reactions to transform silica templates of complex 3-D micro/nanostructured morphology and modest SSA (<2 m²/g) into positive 3-D replicas comprised of micro/mesoporous carbon with very high SSA (i.e., >1,300 m²/g, comparable to commercial activated carbons). By allowing for the retention of the 3-D microscale shape and fine features of silica templates, this process enables the syntheses of high-SSA carbon assemblies with a wide variety of selectable architectures directly inherited from numerous available sustainable (inexpensive) biogenic silica (as well as synthetic silica) templates. Such control over 3-D morphology allows for the tailoring of fluid (gas or liquid) transport through, and nanoparticle dispersions within, highly porous carbon structures for enhanced catalysis, filtration, intercalation, or adsorption for numerous applications, such as in energy storage and harvesting, sensing, water purification, carbon sequestration, and (bio)chemical separation.

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Fig. 4 a) Polarization curves for ORR measured on a RDE in an O₂-saturated 0.5M H₂SO₄ solution at 2000 rpm with a voltage scan rate of 10 mV/sec, b) impedance spectra at 0.8 V under the influence of an ac voltage of 10 mV, and c) chronoamperometric curves for ORR at 0.8 V.

Fig. 5 Secondary electron images of an ion-milled cross-section of a carbon replica of an Aulacoseira diatom frustule.
References


10 Powder Diffraction File, Card No. 89-2955 for Si, Card No. 73-1708 for β-SiC, Card No. 75-0444 for C (International Center on Diffraction Data, Newtown Square, PA).

