

First Synthesis of Continuous Mesoporous Copper Films with Uniformly Sized Pores by Electrochemical Soft Templating

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Abstract: Although mesoporous metals have been synthesized by electrochemical methods, the possible compositions have been limited to noble metals (e.g., palladium, platinum, gold) and their alloys. Herein we describe the first fabrication of continuously mesoporous Cu films using polymeric micelles as soft templates to control the growth of Cu under sophisticated electrochemical conditions. Uniformly sized mesopores are evenly distributed over the entire film, and the pore walls are composed of highly crystallized Cu.

Creating extended metallic 3D architectures is important because such kind of well-patterned networks improves material utilization efficiency and catalytic activity.^[1] The nanoscale structure of metals have been manipulated through various physical and synthetic routes. For example, top down nanofabrication can generate precisely defined structures down to the nanometer level, but making complex 3D structures is challenging.^[2] Likewise, free-standing metal nanoparticles can be assembled into large-scale superstructures, but the process tends to be rather slow.^[3] Especially, mesoporous metals have sparked great interest due to their large surface areas and uniformly-sized pores, opening up a wide range of applications.^[4] Growing metals in sacrificial templates has been extensively studied by developing new kinds of hard and soft templates that can guide the deposition of metals and then be removed to reveal the mesoporous metallic structures.^[4b,c,5] The lyotropic liquid crystal (LLC) method^[6] has been designed to work in combination of electrochemical processes, and has been further extended for the synthesis of various mesoporous metals. To our knowledge, however, the compositions of mesoporous/nanoporous metals have been mostly limited in noble metals, such as Pt,^[7] Au,^[8] Pd,^[9] and their mixtures,^[10] despite the diversity of the morphologies and the wide range of usage. To extend the

utility of the mesoporous materials and make them ubiquitous, we must focus on non-noble metals that are far more Earth abundant than noble metals. The preparation of porous non-noble metals, however, is still very challenging because of their low reduction potentials, high susceptibility upon exposure to solution and air, and uncontrollable deposition behaviors.^[11]

Copper (Cu) is a good candidate for mesoporous synthesis because it is a relatively abundant metal and has been well known to exhibit size-dependent properties in various application fields (e.g., sensing, conductivity, anti-bacteria, selective conversion of CO₂ into fuels).^[12] In particular, the high surface area and fast diffusion through the porous network makes these surfaces ideal for sensing applications. For example, a good non-enzymatic glucose-detection performance has been realized by using a Cu-foam-based sensor because of the unimpeded mobility of glucose and its reaction products offered by the robust hierarchical porous architectures.^[13] Synthesis of such kind of Cu foams relied on evolved hydrogen bubbles as templates, exclusively resulting in micron-sized pores.^[14] We hypothesize that introducing uniformly spaced, nanometer-sized pores in Cu will greatly improve its sensing capability because of increased surface area and more uniform network for the diffusion of analytes. Nevertheless, it has been extremely difficult to create uniformly mesoporous Cu in part because Cu has a strong preference even at small length scales, favoring polyhedral particles and wires instead of extended networks.^[12a,15]

Herein we describe the synthesis of continuously mesoporous Cu films using block copolymers as soft templates. An external reducing potential was applied to guide the growth of Cu, ultimately forming a diaphanous slab of mesoporous Cu (Scheme 1). The pore walls of these films are highly crystalline throughout the film—from the bottom to the top surface. Finally, we show that these mesoporous Cu films can efficiently and selectively detect glucose, even in actual serum samples.

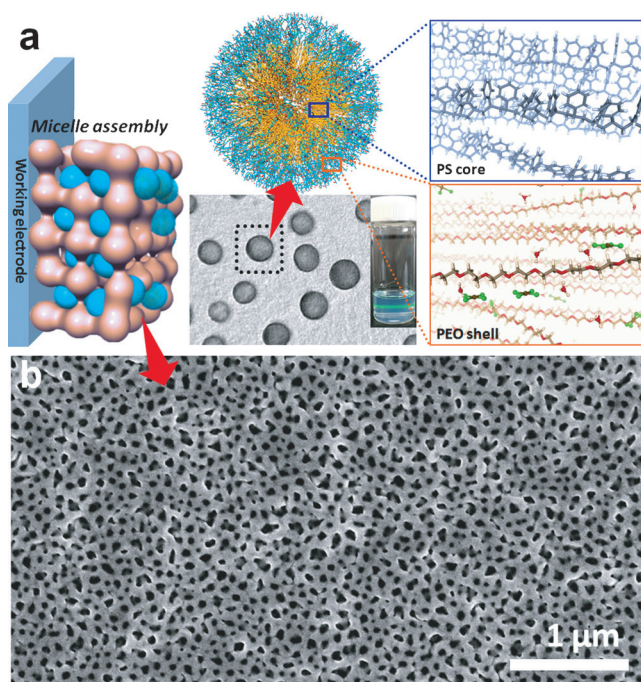
In a typical preparation, 10 mg of polystyrene-*b*-poly(oxyethylene) (PS₆₃₀₀₀-*b*-PEO₂₆₀₀₀, abbreviated as PS-*b*-PEO) was dissolved in 3 mL of tetrahydrofuran (THF) at 50 °C, then 1.5 mL of ethanol was added to the solution. After the solution was thoroughly mixed, 1 mL of aqueous CuSO₄ (80 mM) was added, resulting in the formation of spherical micelles. To increase the ionic conductivity of the electrolyte, 2.5 mL of H₂SO₄ (500 mM) was added slowly to this mixture. Gentle stirring for 5 h ensured that the dissolved Cu species were well incorporated in the exterior PEO region of the micelles, resulting in a transparent light-green colored

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Scheme 1. a) Schematic illustration for the preparation process, and b) the typical SEM image of the mesoporous Cu film.

solution that was used for electrodeposition (see Supporting Information for additional details).

Using this solution as the electrolyte, the mesoporous Cu films were electrodeposited on a conductive Au-coated Si(Au-Si) wafer at a constant voltage of -0.4 V at room temperature and without stirring (see Supporting Information for additional details on the electrochemistry setup). The PS-*b*-PEO was completely removed by ultraviolet ozone treatment. Scanning electron microscopy (SEM) micrographs show that the top side of the film is composed of uniformly distributed mesopores (Figure 1a,b and Figure S1). The entire area of the film is homogenous and free of any defects such as perforations or bumps. According to SEM, the average pore size is about 48 nm, with a wall thickness of approximately 65 nm (Figure 1b and Figure S1). Longer deposition times led to thicker films. But surprisingly no variation in the size and uniformity of the pores regardless of thickness was observed. This approach shows a fairly high repeatability in the porous constructions no matter the variations of the electrolyte batches, making it a robust method for large-area mesoporous coating, perhaps extending to meter length scales (Figure S2).

A cross-sectional specimen sampled from the resulting mesoporous Cu film was further investigated by using transmission electron microscopy (TEM). Even in the earliest stages of deposition, the mesoporous structures formed on the Au-Si substrate (Figure 1e and Figure S3a). High-resolution TEM (HRTEM) images show that the Cu frameworks are highly crystalline without any amorphous domains, and clear lattice fringes extended unabridged from bottom to top of the film (Figure 1c and Figure S3b–c). The lattice fringes with a constant d spacing of 0.21 nm can be well ascribed to the (111) lattice planes of face-centered cubic (fcc) Cu crystals

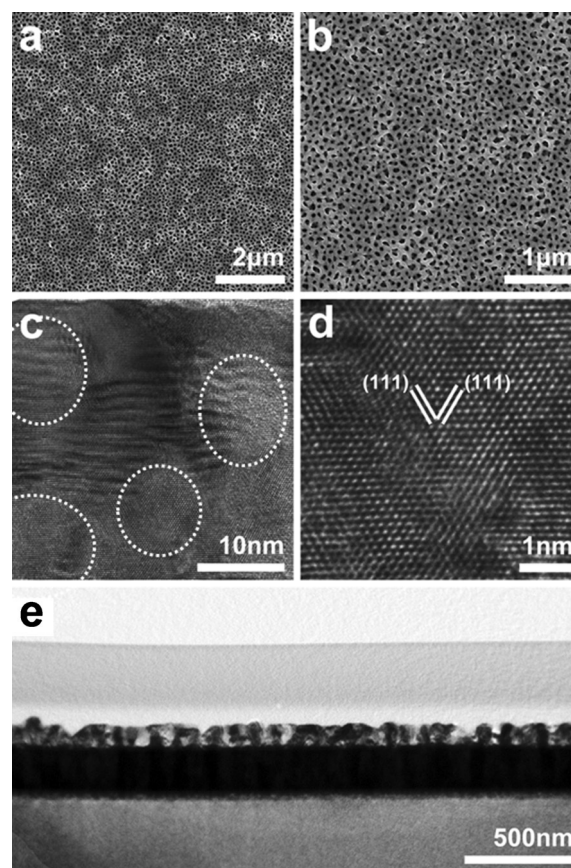


Figure 1. a),b) SEM images of the top-surface of the mesoporous Cu films. c),d) Cross-sectional HRTEM images of the mesoporous Cu films. The pores are indicated by circles in panel (c). e) Low-magnification TEM image of the cross-section of the mesoporous Cu film.

(Figure 1d). The crystallinity of the obtained Cu film was then studied by wide-angle X-ray diffraction (XRD) carried out in in-plane mode with various incident angles, which is sensitive to the lattice planes that are perpendicular to the film surface. With an incident angle in the range of 0.5 – 1.5° , the (220) diffraction peak ascribed to Cu fcc crystal is clearly observed, although a tiny intensity of other diffraction planes are observed (Figure S4). A deeper scanning to the Cu films at a higher incident angle ($>5.0^\circ$) showed the diffraction peaks from the Au substrate. Thus, the in-plane XRD data shows the Cu crystals have a preferential orientation according to the (110) plane vertical to the substrate. In the case of one-dimensional (1D) Cu nanowires, previous reports have demonstrated a crystal growth along the $\langle 110 \rangle$ direction.^[16]

The polymeric micelles play the central role in the fabrication of uniform mesoporous Cu films. Therefore, an investigation to the micellization process was conducted. In pure THF, the diblock copolymer, PS-*b*-PEO, is completely dissolved to be a clear solution without any micelles or aggregates (Figure 2a-i). The addition of ethanol and aqueous solution (CuSO_4 and H_2SO_4) led to a cloudy solution indicating the polymeric micelles, which was further evidenced by the Tyndall effect in the electrolyte (Figure 2a-ii). It should be noted that the addition of CuSO_4 , caused the

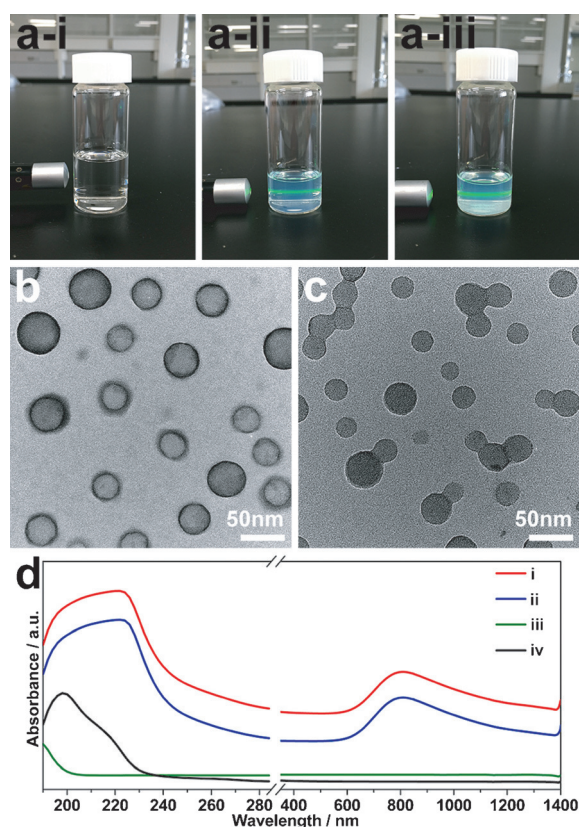


Figure 2. a) Optical micrographs demonstrating the emergence of the Tyndall effect in the electrolyte solution with different compositions: a-i) PS-*b*-PEO dissolved in THF, a-ii) PS-*b*-PEO + CuSO₄ + H₂SO₄, and a-iii) PS-*b*-PEO + CuSO₄. Typical TEM images of the polymeric micelles b) before and c) after the addition of CuSO₄ precursor. The samples are stained with 1.0 wt% phosphotungstic acid to highlight the micelles. d) UV/Vis-NIR spectra of different solutions: i) PS-*b*-PEO micelles + CuSO₄ + H₂SO₄, ii) CuSO₄, iii) H₂SO₄, and iv) PS-*b*-PEO micelles, respectively. (Note: to highlight the interactions among different compositions, UV/Vis-NIR spectra of low- and high-wavelengths were obtained at different composition concentrations: low concentration for low-wavelength, and high concentration for high-wavelength.)

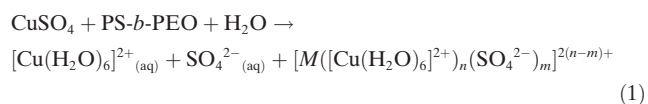
micelles to aggregate slightly (Figure 2b,c and Figure S5) and decrease in diameter (from 36 nm to 28 nm; Figure S5). These observations indicate that the CuSO₄ stimulated the interaction between the micelles.^[5b]

Retaining the micelle structure during Cu deposition, which depends on the interaction between the micelles and the metal precursors, was vital to the templating of uniform mesopores. To understand the formation mechanism of mesoporous Cu, ultraviolet-visible-near infrared (UV/Vis-NIR) absorption measurements were carried out for the electrolyte solutions under our working conditions (Figure 2d). Cu²⁺ ions are readily complexed with water and the peak at around 800 nm corresponds to a d–d transition of [Cu(H₂O)₆]²⁺ ion (Figure 2d).^[17] The other bands in the UV region of the spectrum originate from various moieties of the surfactants, such as the benzene rings of the phenyl group of the surfactant. Cu²⁺ ions in an octahedral field display a single d–d transition from a ²E_g ground state to a ²T_{2g} excited state.^[17] The tail on the low energy site is due to Jahn–Teller

distortion.^[17] Although it is difficult to predict whether the coordination sphere is water molecule interacting with the ethylene oxide moiety of the micelles through hydrogen bonding or surfactant oxygen of the ethylene oxide units, the dissolved Cu species should be 2⁺ charged. The d–d band completely disappears upon electrochemical reduction process.

In the micelle solution, it is likely that the CuSO₄ dissolved as [Cu(H₂O)₆]²⁺ and SO₄²⁻ ions and interact with the ethylene oxide shell domains of the micelles.^[18] We consider that the Cu ions can be, in the solution, as free ions as well as in the hydrophilic domains of the micelles. The micelles with Cu species could be neutral, negatively or positively charged depending on the Cu²⁺/SO₄²⁻ ratios in the micelles. Since reduction takes place at the cathode, the Cu²⁺ rich micelles (overall positively charged ones) and free [Cu(H₂O)₆]²⁺ ions can be selectively directed to the cathode surface, where the Cu²⁺ ions are reduced to the metallic copper. The above formation mechanism satisfactorily explains the formation of mesopores in our self-assembly-reduction process.

The following chemical equilibrium [Eq. (1)] summarizes the assembly process, where “*M*” stands for micelle, (aq) species are free ions, and (*n*–*m*) can be negative or positive depending on the Cu²⁺/SO₄²⁻ ratios.



The addition of H₂SO₄ takes a substantial role in obtaining continuous mesoporous Cu films. In the absence of H₂SO₄, only isolated Cu nanoparticles are confirmed on the substrate (Figure S6). By gradual increase the concentration of H₂SO₄, the porous structures of the continuous Cu film become apparent. Further optimization of the technique led us to settle on a H₂SO₄ concentration of 156.25 mM (Figure S7). Our method is distinctly different from previous reports that use H₂SO₄ to create H₂ bubbles that direct the formation of large-sized pores in the films.^[14,19] In our method, H₂SO₄ has two essential effects: 1) the addition of H₂SO₄ solution can prevent a formation of hydroxides of Cu precursors,^[20] which can be well supported by the transparent solution with H₂SO₄ solution (Figure 2a-ii), while, in the absence of H₂SO₄, some precipitates are formed at the bottom of the electrolyte (Figure 2a-iii); and 2) H₂SO₄ increases the ionic conductivity of the electrolyte, favoring a high deposition rate of Cu and the formation of continuous films. When the used acids were changed to other types, the quality of mesoporous structures was decreased (Figure S8).

The impact of H₂SO₄ on ionic conductivity can be observed in the electrodeposition traces before and after addition of acid (Figure S9a). Without H₂SO₄, the cyclic voltammetric (CV) curves show a very small and linear current response, suggesting the huge resistance of the electrolyte. Upon addition of H₂SO₄ solution, the CV curve showed two reduction processes that correspond to: Cu²⁺ → Cu⁺ and Cu⁺ → Cu⁰ (Figure S9a). According to the Pourbaix diagram, direct reduction of Cu²⁺ to Cu⁰ is possible by

manipulating the reduction potential at such an acidic condition in the present study.^[21] This excludes the possibility for the formation of other Cu phases (e.g., Cu₂O). Additionally, the charge transfer resistance was further investigated at two different electrolytes. Without any doubt, the charge transfer resistance decreases dramatically upon the addition of H₂SO₄ (Figure S9b,c). We believe that this is the key to preparing continuous Cu films with high reproducibility.

The reduction rate of Cu species is also extremely important for the final mesoporous structures.^[6a,8a,22] Electrochemical deposition enables the precise regulation of the crystal growth manner of the Cu walls that form around the micelles. Variations in the structure of the mesopores could be observed by using different constant potentials between -300 mV and -700 mV (Figure S10). From the top surface structures of the obtained films, it is obvious that the mesoporous structures are greatly changed, and optimized at particular potentials (i.e., -300 mV and -400 mV). A high reduction rate of Cu at relatively low potentials (e.g., -600 mV and -700 mV in the present work), affords extremely large sized Cu crystals at the initial stage, which then grow rapidly to exceed the size of the hydrophilic volume of the micelles, thus leading to films with inadequate porous structures (Figure S10d,e).^[6a,9b]

The films with good mesoporous structures, that were prepared at three different potentials (i.e., -300 mV, -400 mV and -500 mV), were studied by cyclic voltammetry to sign light on how the accessible surface area depends on the applied constant potentials. The mesoporous Cu film prepared at -400 mV had the highest current density compared to other films (Figure S11), indicating that the great advantage of perfect mesoporous structures. A high deposition potential (-300 mV) resulted in a relatively slow reduction speed, and small nanocrystals are obtained on the surface of the films, as shown in the inset of Figure S11a. The formation of small nanocrystals caused by the slow deposition rate finally decreased the surface area of the porous film.

Non-enzymatic glucose sensors based on nanostructured Cu have been intensively studied owing to the material's good conductivity, low cost, and superior performance. As a result of its self-supported mesoporous structures, mesoporous Cu film is a good candidate for the direct detection of glucose to satisfy the requirements of diverse applications by maximizing the sensitivity and achieving favorable selectivity. A highly improved activity and selectivity toward the glucose detection was realized by using the nanoporous Cu film, indicating an attractive application for the daily use. The details are given in Figure S12 and Table S1.

In summary, we have, for the first time, demonstrated the preparation of large-scale mesoporous Cu films based on a novel and simple approach to precisely balance the growth manner of the Cu frameworks and the micelle structures regardless the substrate variations. The present success not only indicates the capability of creating mesoporous structures with highly crystalized metals, but also paves a new way for preparation of porous Earth-abundant metals for diverse applications and at low cost.

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