Carbon Foam Decorated with Silver Nanoparticles for Electrochemical CO₂ Conversion


Electrochemistry is a promising method to recycle CO₂ into useful carbon feedstock and for storing intermittent renewable energy. To date, Au and Ag nanoparticles are the most active catalysts for electrochemical conversion of CO₂ to CO. However, agglomeration reduces the activity and the high cost slows widespread commercialization. Suitable support materials are thus needed to improve catalyst utilization. We explore carbon foam (CF) as a catalyst support. Compared with carbon black or graphene nanoplatelets, CF has higher surface area, larger pores, and more defects, resulting in improved uniformity of Ag nanoparticle distribution as well as higher activity and efficiency for CO₂ conversion to CO.

Atmospheric CO₂ levels have increased significantly, and this has been linked to climate change, erratic weather patterns, increasing temperatures, and rising sea levels. Renewable energy sources such as wind and solar are being implemented at large scale to lower carbon emissions.[1] However, the intermittency of such technologies hinders their widespread adoption as they are not always available when demand is high. Electroreduction of CO₂ to CO using off-peak renewable energy has potential to directly reduce CO₂ emissions, create fuels or chemical precursors, and act as an energy storage medium to smooth out supply and demand.[2] Noble metals such as Ag and Au are highly active for the electrochemical conversion of CO₂ to CO.[2a,3] However, suitable catalyst support materials with high electrical conductivity, suitable porosity, and good stability are necessary to improve catalyst utilization and reduce cost.[4] We previously developed carbon foam (CF) as a Pt catalyst support for fuel cells.[5] Large surface area and good nanoparticle (NP) distribution resulted in high electrochemical surface area and oxygen reduction reaction activity. Such CF may thus also be suitable as an Ag support for the electroreduction of CO₂. Here, we report the synthesis of silver-decorated CF for electroreduction of CO₂ to CO.

Scanning electron microscope (SEM) images (Figure 1) show that CF has a highly porous carbon network with μm-scale voids and thin, graphene-like cell walls. In contrast, commercial graphene nanoplatelets (CG) comprise μm-scale agglomerates of restacked graphene sheets whilst carbon black (CB) forms ≈5–20 μm agglomerates with a primary particle size of ≈40 nm. Brunauer–Emmett–Teller (BET) nitrogen adsorption reveals surface areas of 904, 652, and 237 m²/g for CF, CG, and CB, respectively. CF has by far the highest surface area. CB and CG have broad pore-size distribution (Figure 2a) whereas CF mainly has pore radii of <2 nm in addition to the micrometer-scale pores observed by SEM.

Raman spectra for CF, CG, and CB are shown in Figure 2b. The G peaks at 1592, 1574, and 1582 cm⁻¹; the G-peak full-width-at-half-maximum (FWHM) of 85.8, 63.4, and 88.9 cm⁻¹; and the I₆/I₅ ratios of 0.99, 0.95, and 1.04, respectively, indicate that CF has a high degree of disorder similar to CB whereas CG is a more crystalline material.[6,7] The morphology and distribution of Ag nanoparticles formed on the support surface was investigated using trans-

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mission electron microscopy (TEM, Figure 3). The sheet-like structure of both CF and CG is clearly observed whereas CB is formed of spheroidal carbon NPs with \( \approx 40 \text{ nm} \) diameter. Ag NPs with a diameter of \( \approx 5 \text{ nm} \) are formed and distributed uniformly across the surface of Ag/CF (several larger particles are also observed). In contrast, the NPs are much larger (\( > 20 \text{ nm} \)) and have poor distribution in both Ag/CG and Ag/CB. The difference in Ag distribution is attributed to the larger surface area of CF and the higher degree of disorder providing numerous anchoring sites for NP nucleation at edges and defects.

The catalytic activity towards electrochemical CO\(_2\) reduction was measured in a flow reactor.\(^{[4a,8]}\) Figure 4a shows partial current densities for CO production (\( j_{\text{CO}} \)). Ag/CF has by far the highest (\( \approx 120 \text{ mA cm}^{-2} \) at \(-2.06 \text{ V}_{\text{Ag/AgCl}}\)), followed by Ag/CG (\( \approx 58 \text{ mA cm}^{-2} \)), and Ag/CB (\( \approx 30 \text{ mA cm}^{-2} \)). The differences are mainly driven by the Faradaic efficiency (FE, Figure 4b). Ag/CF has the highest FE (84% at \(-1.82 \text{ V}_{\text{Ag/AgCl}}\)), but this drops at more negative potentials possibly due to Ag agglomeration during operation for high loadings.\(^{[4a]}\) The improved performance of Ag/CF is attributed to the porosity, large surface area, and improved NP distribution, leading to increased mass transport and larger triple-phase boundary.\(^{[9]}\) Ag/CG performs better than Ag/CB due to its higher electrical conductivity and surface area. Additionally, some Ag NPs may be trapped in the deep pores of CB.\(^{[4a]}\) Cyclic voltammetry was performed from 0 to \(-1.8 \text{ V} \) vs. Ag/AgCl in a standard three-electrode cell with 0.5 M KHCO\(_3\) electrolyte (data not shown). This showed that Ag/CF has

![Figure 2](image1.png)  
*Figure 2.* (a) Pore-size distributions and (b) Raman spectra of the CF, CG, and CB samples.

![Figure 3](image2.png)  
*Figure 3.* TEM images of (a) Ag/CF, (b) Ag/CG, and (c) Ag/CB.

![Figure 4](image3.png)  
*Figure 4.* Flow-cell results for Ag/CF, Ag/CG, and Ag/CB: (a) partial current density for CO and (b) Faradaic efficiency.
the highest double-layer capacitance, which may contribute to enhanced desorption of key reaction intermediates, further promoting CO formation and resulting in the improved performance.

In summary, carbon foam (CF) was utilized as a high-performance Ag catalyst support for electrochemical CO₂ conversion to CO, far outperforming CG and CB, with high partial current density and Faradaic efficiency. The improved activity is attributed to the large surface area and porosity as well as to the improved catalyst nanoparticle distribution.

**Experimental Section**

CF was synthesized by thermal decomposition of sodium ethoxide (Wako) in air. [46,10] The resulting powder was crushed, sonicated, and washed in deionized water to remove NaOH, then pyrolysed at 600 °C for 2 h under N₂ flow. The citrate-protecting method was used for Ag decoration. [46,10,11] CF or CG (Graphene Supermarket, Grade 4; 80 mg) were mixed with AgNO₃ solution (188.6 mg AgNO₃ in 18.5 mL H₂O, AgNO₃ solid from Sigma–Aldrich), and stirred for 30 min. Sodium citrate solution (Fisher Chemicals, 131 mg, 18.5 mL aqueous solution) was added dropwise, then NaBH₄ solution (Sigma–Aldrich, 30 mg, 37.5 mL aqueous solution) was added while stirring vigorously in an ice-bath to reduce Ag⁺ to Ag. After gentle stirring overnight, the product was centrifuged, washed, and dried under vacuum at 80 °C for 4 h. Ag/CG was used as Ag catalyst (60 wt%, E-Tek). Characterisation was done by SEM (S-5200, Hitachi), TEM (JEOL 2100 CRYO, 200 kV), BET nitrogen adsorption (Belsorp mini II-VS, Bel Japan, Inc.), and Raman spectroscopy (DM2500M, Renishaw, UK, λ = 532 nm). AG loading was determined by inductively coupled plasma atomic emission spectroscopy (ICP-AES) (PerkinElmer-Optima 2000DV) to be 57.4% for CF and 53.3% for CG.

Cathode catalyst inks were prepared by sonicating Millipore water (200 mL), electrocatalyst powder (2 mg), Nafion® solution (2.6 μL, 5 wt%, Fuel Cell Earth), and isopropyl alcohol (200 μL) for 15 min. The ink was painted onto Sigracet 35BC gas diffusion layers (GDLs, Ion Power, 1 cm²) with a loading of 1 mg cm⁻². The anode loading was also 1 mg cm⁻² (IrO₂). [13] The flow reactor was operated under ambient conditions. [46,10] A potentiostat (Autolab PGSTAT-30, EcoChemie) operating in steady-state chronoamperometric mode was used to measure the current. [14,11] Seven cell potentials were applied, and the product gas was sampled after a steady state was reached (200 s). The current was then averaged over 180 s before changing the potential. Electrode potentials were measured with multiampere (AMPROBE 15XP-B) connected to each electrode and an Ag/AgCl reference electrode (RE-5B, BASi) downstream. CO₂ was introduced at 7 sccm, and 1 M KOH electrolyte was supplied at 0.5 mL min⁻¹. Gaseous products were extracted through the GDL by vacuum and quantitatively analyzed by gas chromatography (GC, Thermo Finnegan Trace, thermal conductivity detection module, He carrier gas). The only cathode products detected were CO and H₂. [14] Partial current densities were calculated by multiplying the total current density by the Faradaic efficiency.

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