

Electro-catalyst systems for energy storage through coupled water oxidation and CO₂ reduction

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Project Publications

1. K. Klingan, T. Kottakkat, Z.P. Jovanov, S. Jiang, C. Pasquini, F. Scholten, P. Kubella, A. Bergmann, B.R. Cuenya, C. Roth, H. Dau, Cavity versus Surface Catalysis on Electrodeposited Cu Foams for Electrochemical CO₂ Reduction, in preparation.
2. T. Kottakkat, K. Klingan, Z.P. Jovanov, H. Dau, C. Roth, Electrodeposited nanostructured Ag-Cu catalysts for enhanced reduction of CO₂ to CO, in preparation.

Project Summary

The steadily rising atmospheric content of CO₂ due to burning of fossil fuels generates much concern with respect to changing global climate. Considering the ambient operating conditions and the possibility of direct integration with renewable resources, electrochemical conversion of CO₂ to fuels and useful chemicals opens door to a sustainable carbon neutral fuel cycle. However, the main challenge is to find a suitable catalyst to overcome the high activation barrier for the reduction of thermodynamically stable CO₂. In addition, when **operated in an aqueous medium (pH ~ 7)**, the selected

catalyst material has to impede the hydrogen evolution reaction (HER) initiated already at much lower overpotentials.

Cu is in the focus of electrochemical reduction of CO₂ owing to its ability to catalyze CO₂ reduction towards energy-dense hydrocarbon products. However, the electrochemical reduction of CO₂ on Cu is challenged by insufficient selectivity and a wide spectrum of products formed. The surface of the catalyst plays a key role in affecting the intermediate adsorption and by controlling the number of active sites for the reactants. The performance of Cu-based catalysts can be improved by tuning the structure, morphology and composition of the catalyst.

Evaluation of Cu-based nanostructured materials with controlled morphology for CO₂ reduction

By utilizing the **hydrogen bubble templated electrodeposition method**, the morphology and the pore size can be easily controlled by changing the deposition current and time. During the electrodeposition of Cu at high current densities, the hydrogen bubbles originating from the cathodic reaction create a continuous path from the electrode surface to the electrolyte air interphase, and these bubbles act as a dynamic negative template for the Cu deposition (Fig. 1).

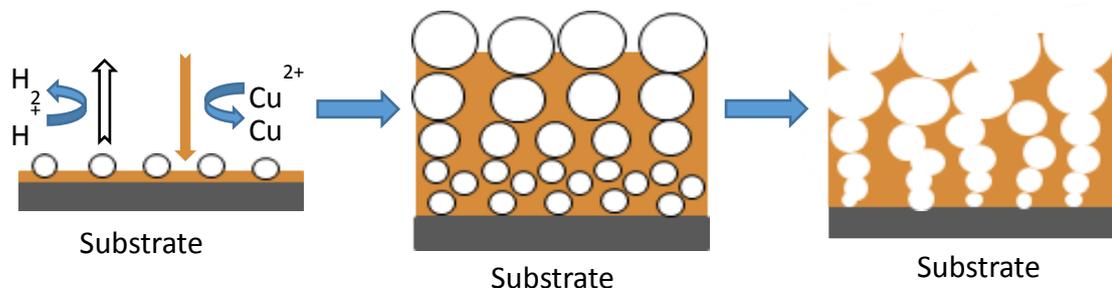


Figure 1. Hydrogen bubble templated electrodeposition of Cu

Apart from the Cu only structures, we also develop bimetallic electrocatalysts for CO₂ reduction, namely **AgCu** and **CuZn** by electrochemical synthesis. Fig. 2 shows the porous Cu and bimetallic AgCu synthesized by hydrogen bubble templated electrodeposition method.

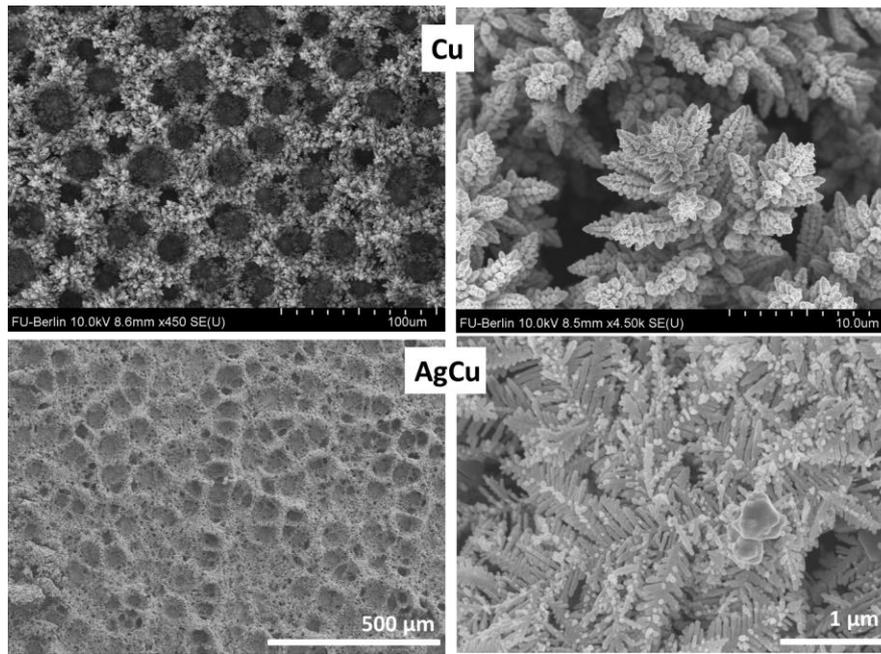


Figure 2. SEM images of Cu and Ag-Cu structures electrodeposited using the hydrogen bubble templated electrodeposition method.

The synthesized materials are initially characterized for the electrochemical CO₂ reduction by means of linear sweep and cyclic voltammetry and chronoamperometry. Determination of the product distribution of the electrochemical CO₂ reduction is carried out by an online gas analysis using a gas chromatograph.