

CO₂ Adsorption on Copper Catalyst as Enhanced by Ion Liquid

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Electroreduction of CO₂ represents a promising solution toward addressing global challenges in energy and sustainability [1-3]. The technology success depends on developing efficient electrocatalysts capable of selectively reducing CO₂ to valuable hydrocarbon products at low overpotentials (ethylene, methanol, CO etc.). So far, the best candidate is nanostructured copper and for efficient catalyst reaction, the CO₂ should adsorb from gaseous phase on the catalyst surface. CO₂ capture with ionic liquids is well known. The distinct properties of ionic liquids such as negligible vapor pressure and their affinity to capture the CO₂ molecules make them a feasible alternative for currently available solvents including, different amines. [4]. In this work, we investigated the adsorption process stimulated by ionic liquid. CO₂ adsorption on Cu surface was studied by ATR-FTIR spectroscopy and electrochemical processes by voltammetry.

Spectra measured using FTIR spectrometer Vertex 80v (Bruker, Germany) together with ATR accessory: VeeMAX III (PIKE Technology Inc., USA). The surface of the ATR crystal from germanium was coated with 20 nm thick copper. This surface was blown by a stream of CO₂. In situ measurements showed a significant increase in the adsorption of CO₂ on the surface of copper in the presence of an ionic liquid.

Electrochemical measurements were performed using Autolab potentiostat/ galvanostat in electrochemical cell with KHCO₃ electrolyte (1M and 2.5M) at a room temperature. Three-electrode configuration was applied with a copper on ATR crystal as a working electrode and black platinum as a counter electrode.

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