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Investigation of CuAl-layered double hydroxides as efficient electrocatalysts of CO₂ reduction

<u>TARHINI Ali</u>^{(a)*}, GUYOT Mélanie^(b), COSTENTIN Cyrille^(b), CHARDON-NOBLAT Sylvie^(b), MOUSTY Christine^(a), PREVOT Vanessa^(a)

^(a) Institut de Chimie de Clermont-Ferrand, Clermont-Ferrand (France) ^(b) Département de Chimie Moléculaire, Grenoble (France) * ali.tarhini@etu.uca.fr

The emission of CO_2 by industrial action has a serious impact on our environment which leads us to look for ways to decrease the emitted CO_2 to the atmosphere. The electrochemical conversion of CO_2 (CO_2ER) into other molecules such as CO, HCOOH, CH_3COOH , and CH_4 is one of the many ways to achieve this goal. Not only it will help to reduce pollution, but it will also convert CO_2 to valuable products to use as a feedstock.^[1] Recently, efficient copper based electro-catalysts for CO_2ER were prepared from layered double hydroxides (LDH) ^[2]. Using CuAl-NO₃ LDH prepared by the coprecipitation method and thermal treatments, a morphology control allowed to improve the catalytic activity by tailoring the structure of active sites and increasing the surface area/number of active sites. The resulting catalyst promoted the electrochemical conversion, in galvanostatic conditions, of CO_2 to CO and HCOOH with a faradaic efficiency (FY) up to 42% and 22%, respectively.

In the present work, the influence of the chemical composition of Cu based LDH of both the layer and the interlayer domain on the CO₂ER was studied. Several copper based LDH were synthesized by varying the synthetic method and taking into account the Cu/Al ratio and the interlayer anion $(CO_3^{2^-})$, Acetate Ac and dodecylsulfate SDS). The electrochemical behavior of these phases was first studied in cyclic voltammetry in different electrolytes under CO₂ saturated atmosphere in order to put in light their electrocatalytic behavior towards CO₂ER. The experimental parameters were optimized to perform the macroscale electrolyses. Among the different phases tested, Cu₂Al-CO₃ LDH phase prepared by coprecipitation showed the highest faradic yield of HCOOH in 0.5 M KCl, reaching 48%.

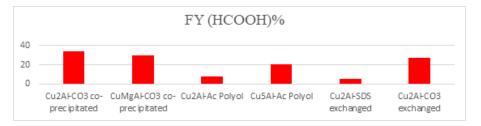


Figure.1: Faradic yields of HCOOH formation by copper based LDH synthesized by different

methods in 0.45 M KHCO3

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