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Au/TiO₂ Catalyst for Conversion of CO₂ in Ethanol

Tomaz Neves Garcia,^{1,*} Adriano Braga¹, Liane Marcia Rossi¹

¹Instituto de Química – Universidade de São Paulo, São Paulo, Brazil

*Corresponding author: tomaz@iq.usp.br

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Global warming is one of the most serious problems humanity is facing, changing fauna and flora over the years and putting at risk the existence of life as we know it. There is an urgent need to reduce the emission of greenhouse gases, because the emission of these types of gases may one day be no longer optional. Aligning environmental concerns with the need for the formation of the valuable C-C bonds from a non-fossil resource, widely available and renewable, the valorization of CO₂, as well as its use as a source of carbon in fine chemicals synthesis, receives great relevance today.^{1,2} However, because it is a thermodynamically stable and kinetically inert molecule, employing CO₂ as a building block in fine chemical synthesis is very challenging. Among the various types of CO₂ utilization reactions, its direct application to olefin carbonylation has attracted great interest. The classical process of hydroformylation, carbonylation of olefins with synthesis gas (CO and H₂), which employs Rh and Co catalysts, is considered a leader in the industrial production of aldehydes. The direct use of CO₂, possibly with the formation of CO in situ, requires new catalysts or CO₂ activators.

In the present work gold nanoparticles supported in TiO₂ (anatase), has been successfully synthesized by deposition–precipitation with urea (DPU), where H₂AuCl₄ is vigorously stirred at 80°C for 20h with a little excess of urea in deionized water. Afterward the solid is washed with acetone and water and dried under reduced pressure. The catalyst has been characterized by transmission electron microscopy, **Figure 1**, showing the formation of 1 nm averaged gold nanoparticles well distributed in the TiO₂ (anatase) surface. Atomic absorption spectrometry analysis demonstrated that 91% of the H₂AuCl₄ gold was impregnated in the TiO₂ (anatase) surface, resulting in 0.97 wt.% Au.

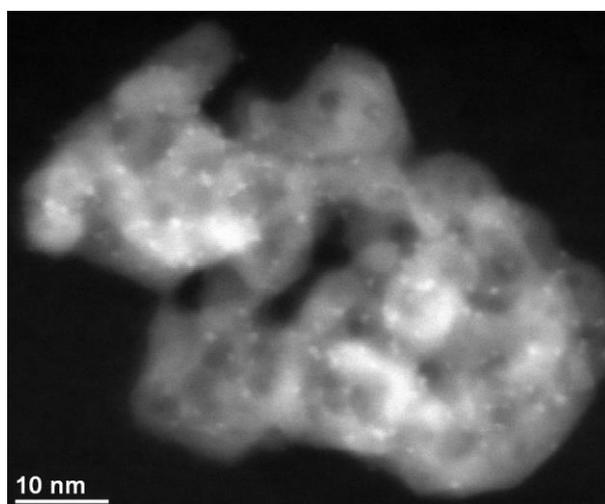


Figure 1. Au/TiO₂ catalyst

The catalyst successfully converts CO₂ and H₂ giving ethanol as the only product. The reaction was performed at 40 bars of CO₂/H₂ 3:1 (at room temperature) and heated to 200°C with DMF or Acetonitrile as solvent for 10 hours. Currently the work is focusing on studying the effect of solvent, additives and catalyst doping in the yield/selectivity of the reaction. The obtained results exhibit the potential of the Au/TiO₂ system to activate small molecules like CO₂ and H₂, as already demonstrated by our group,³ having the potential to catalyze other types of CO₂ conversion, like olefin carbonylation.

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