

Support Effect in High Activity Gold Catalysts for CO Oxidation

Massimiliano Comotti, Wen-Cui Li, Bernd Spliethoff, and Ferdi Schüth*

Contribution from the Max-Planck-Institut für Kohlenforschung, Kaiser-Wilhelm-Platz 1, D-45470 Mülheim a.d. Ruhr, Germany

Received September 7, 2005; E-mail: schueth@mpi-muelheim.mpg.de

Abstract: In this work, we present a detailed study concerning the evaluation of the metal–support interaction in high activity gold catalysts for CO oxidation. Using the colloidal deposition method, model catalysts were prepared, which allow the isolation of the effect of the support on the catalytic activity. Prefabricated gold particles were thus deposited on different support materials. Since the deposition process did not change the particle sizes of the gold particles, only the influence of the support could be studied. TiO₂, Al₂O₃, ZrO₂, and ZnO were used as support materials. Catalytic tests and high resolution transmission electron microscopy clearly show that the support contributes to the activity. However, our results are not in line with the distinction between active and passive supports based on the semiconducting properties of the oxidic material. The most active catalysts were obtained with TiO₂ and Al₂O₃, while ZnO and ZrO₂ gave substantially less active catalysts. Furthermore, the effect of other important parameters on the catalytic activity (i.e., particles size distribution, calcination temperature, and aging time for a Au/TiO₂ catalyst) has also been studied. Using this preparation route, the catalysts show high-temperature stability, size dependent activity, and a very good long-term stability.