

Observation of electronic metal support interaction: investigation of Au/Ceria catalysts by HERFD-XANES

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CO oxidation on a supported gold nanocatalyst has been a paradigmatic and pivotal problem in catalysis in the last two decades¹. The scientific debate is focused on the relevance of particle size, valence state of gold and interaction with the support, however, despite the huge amount of experimental data and model simulations reported in the literature, an ultimate analysis of the key-factors determining the impressive low-temperature activity of nanosized gold towards CO oxidation is still lacking. Actually, these factors are difficult to isolate because they are closely intertwined.¹ However, the beneficial effect of the metal support interaction remains one of more accredited phenomenon able to explain the high catalytic performance of gold based metal catalysts. To address this crucial issue, a detailed investigation, by both experimental and computational approach, of the metal-support interface was undertaken.

According to theoretical prediction, the most important information arises from the interface Au atoms, which determine band modification both for the metal cluster and the support.

In this paper we report the results of a recent in situ HERFD-XANES at RT experiment carried out at the ID26 beam line of ESRF. It is evidence that a cyclic modification occurs and this is reversible during the redox treatments. Interestingly, Au cluster mediates the oxygen uptake in the ceria support (see fig below).

