

Stabilising the Au₉ Clusters at the TiO₂ Film using Chromium Oxide Layer Depends on the Stability of TiO₂ Film

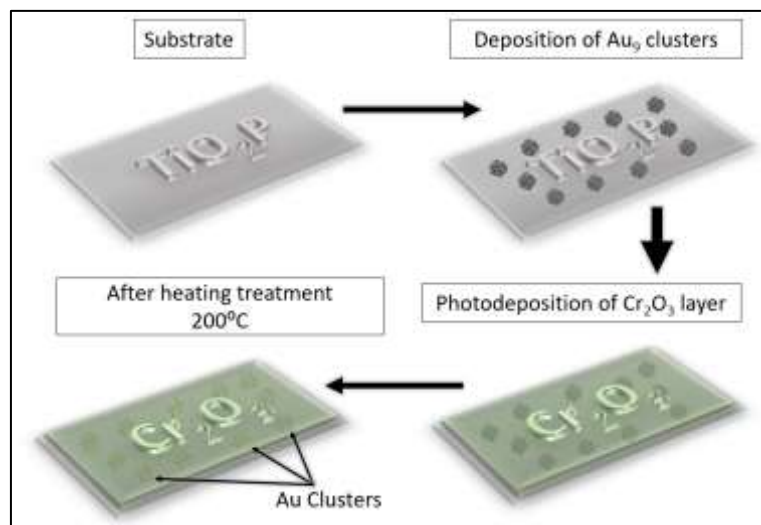
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Abstract

Hydrogen is considered as one of the future energy sources. Hydrogen is a storable commodity and can be used fuel for a large range of applications. One possibility to produce hydrogen from renewable energy resources is photocatalytic water splitting. This process requires a photocatalyst with a minimum bandgap, which is estimated to be around 1.6 eV (Wang and Domen 2020). The efficiency of photocatalysts can be enhanced by using co-catalysts (Kurashige, Kumazawa et al. 2018). Stabilising chemically synthesised gold clusters on a metal oxide surface without agglomeration is a major challenge (Metha, Andersson et al. 2019). We have investigated the stability of phosphine-stabilised clusters Au₉(PPh₃)₈(NO₃)₃ (Au₉) on TiO₂ films by depositing a Cr₂O₃ overlayer. Two different thicknesses of TiO₂ films were used in this study; TiO₂P and TiO₂G. Both were made using the radio-frequency sputtering technique. The TiO₂ film thickness for TiO₂P and TiO₂G is ~ 400 nm and 1100 nm. X-ray photoelectron spectroscopy was used to determine the chemical composition and electronic structure of the surface. The thickness of TiO₂ films was imaged by scanning electron microscopy. The crystallinity and the surface topography of TiO₂ films were characterised using x-ray diffraction and laser confocal microscopy. Au₉ clusters were successfully stabilised on the surface of the TiO₂P film; however, Au₉ clusters agglomerated at the surface of TiO₂G. The agglomeration of Au₉ clusters in the latter is due to the high mobility of TiO₂G film during the heating treatment to remove the ligands.

Metha, G. F., et al. (2017). Photocatalytic conversion of carbon dioxide and water into substituted or unsubstituted hydrocarbon(s) WO Patent App. 2016/051175.

Kurashige, W., et al. (2018). "Au₂₅-Loaded BaLa₄Ti₄O₁₅ Water-Splitting Photocatalyst with Enhanced Activity and Durability Produced Using New Chromium Oxide Shell Formation Method." The Journal of Physical Chemistry C **122**(25): 13669-13681.

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