Interface Structures of Gold Nanoparticles on TiO$_2$ (110)

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Scanning transmission electron microscopy and density functional theory are used to characterize atomic structures of nanoscale heterointerfaces between gold nanoparticles and a TiO$_2$ (110) surface. It is found that when the gold nanoparticle size is smaller than a few nanometers, gold atoms preferentially attach to specific sites on the TiO$_2$ surface and thus form an epitaxial and coherent heterointerface. Conversely, as the gold size becomes larger, the gold-TiO$_2$ interface loses lattice coherency in order to accommodate the large lattice mismatch between the two dissimilar crystals.

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Gold nanoparticles dispersed on metal-oxide supports are active catalysts for a variety of chemical reactions [1–4]. It has been demonstrated that the unique catalytic activity strongly depends on the size of Au nanoparticles and the type of metal-oxide support [2,5], suggesting the importance of Au-support interfacial interactions on a nanometer scale. However, a detailed description of nanoscale heterointerface formation has not been established, especially its relation to the Au nanoparticle size, and the size-dependent catalytic activity of supported Au is still a matter of conjecture. It is thus essential to characterize the atomic structure of nanosized Au-support interfaces in order to truly understand the origin of the remarkable catalytic activity with strong particle size dependency.

Atomic-resolution scanning transmission electron microscopy (STEM) is a powerful method for characterizing metal cluster catalysts at subnanometer dimensions [6,7]. Under the high-angle annular dark-field (HAADF) imaging mode, the STEM image can be considered as an incoherent image, and the image intensity strongly depends on the atomic number ($Z$) of the constituent atoms [8]. In this study, Au/TiO$_2$ (110) model catalysts were fabricated and atomic-resolution plan-view STEM imaging was carried out to directly observe the atomic structures of Au nanoparticles on the TiO$_2$ (110) surface. The observed structures are interpreted using first-principles density functional theory (DFT) calculations.

A commercially available rutile TiO$_2$ (110) substrate was thinned down by mechanical polishing followed by ion bombardment to obtain electron transparent TEM samples. The TEM samples were annealed in air at 973 K for 30 min to produce atomically flat (110) surfaces, in accordance with previous reports [9–11]. High-purity gold (99.95%) was deposited on the TEM sample surface by vacuum evaporation at room temperature with a base pressure of about $1 \times 10^{-3}$ Pa. Atomic-resolution HAADF STEM images were taken with a 200-kV JEM-2100F TEM-STEM electron microscope (JEOL Ltd., Tokyo, Japan) equipped with an aberration corrector (CEOS GmbH, Heidelberg, Germany). A HAADF detector with an inner angle greater than 70 mrad was used for the STEM observations.

Figure 1(a) shows a HAADF STEM image of a rutile TiO$_2$ crystal observed in the $\{110\}$ projection. There are two types of bright spots with different image intensities as indicated by the arrows. These bright spots correspond to the two different Ti-containing columns as shown schematically in Fig. 1(b). The brighter spots correspond to the atomic columns with both Ti and O atoms (Ti-O columns), whereas the duller spots correspond to the Ti-only columns. Figure 1(c) shows a HAADF STEM image of Au nanoparticles deposited on a TiO$_2$ (110) surface. Inset shows the image intensity profile across a single Au atom, showing strong image intensity on the Ti-O columns.
while the darker spots correspond to the atomic columns with body-centered Ti atoms only (Ti-only columns), as confirmed by HAADF STEM simulation [12]. The signal from the oxygen columns is extremely weak, and thus the troughs in between the Ti-containing columns appear dark. Figure 1(c) shows a typical HAADF STEM image of Au nanoparticles deposited on the TiO$_2$ (110) surface. The Au nanoparticles on the TiO$_2$ surface are clearly visible due to the strong Z-dependent image contrast (Z: Au = 79, Ti = 22, O = 8). The projected Au nanoparticle sizes are in the range 1–5 nm. Figure 1(d) shows single Au atoms attached on the TiO$_2$ (110) surface as indicated by the arrows. The strong image intensity of the Au single atoms is preferentially found on top of Ti-O columns. This site corresponds to the on-top sites of bridging O sites ([13–18]. If the surface is reduced and vacancies are introduced, Au atoms are most stable on the vacancy sites [13–16,18]. These two possibilities agree well with our STEM images. On the other hand, if the surface is oxidized or hydrated, the stable attachment sites become inconsistent with our images [14,18]. The observed Au single atoms are thus considered to be either on the fivefold coordinated Ti sites or the O$_{br}$ vacancy sites on the TiO$_2$ (110) surface. In the present experiment, however, there should be a certain amount of surface oxygen vacancies because the Au deposition was done in vacuum condition. Thus, Au atoms observed here are more likely to attach to the vacancy sites.

Figures 2(a)–2(e) show typical atomic-resolution HAADF STEM plan-view images of Au nanoparticles on a TiO$_2$ (110) surface, arranged in order of projected particle size. It is seen that the atomic structures of both Au and TiO$_2$ can be simultaneously resolved. In the case of very small Au islands (<~2 nm), as in Figs. 2(a) and 2(b), the strong image intensities corresponding to Au atoms are found on the Ti-O columns and on the O columns along the trough of the Ti-containing columns [along the arrows in the magnified image in Fig. 2(f)]. The strong preference of Au atom attachments on the Ti-O columns is consistent with the Au single atom cases. The present results suggest that Au atoms in the small Au islands also preferentially attach to specific sites on the TiO$_2$ surface, and therefore form an epitaxial coherent-type heterointerface [19]. These epitaxial Au islands are not only small in size, but are also very thin (a few atomic layers) as estimated from the STEM image intensity profiles. If the Au atoms on the Ti-O columns are in the first Au atomic layer, Au atoms on the trough of Ti-containing columns (on top of O columns) can be considered as second layer atoms, because nearest Au-Au distances would be too close to each other (less than 80% of the stable Au-Au interatomic distance in bulk Au) if these Au atoms were on the same first atomic layer. This proposed Au bilayer stacking sequence is similar to the (110) stacking ([110]$_{Au}$ || [110]$_{TiO_2}$, [100]$_{Au}$ || [110]$_{TiO_2}$) or (100) stacking ([100]$_{Au}$ || [110]$_{TiO_2}$, [110]$_{Au}$ || [110]$_{TiO_2}$) of fcc Au on TiO$_2$ (110), despite the extremely large lattice mismatch (>20%) in one direction.

On the other hand, if the Au size becomes larger (> ~ 3 nm), as in Figs. 2(c)–2(e), various orientation relationships exist between the Au nanoparticles and the TiO$_2$ substrate. Frequently observed moiré fringes [20] suggest that the lattices are gradually displaced or rotated from each other. These results indicate that the Au/TiO$_2$ (110) interface is not a coherent interface, but an
incoherent-type heterointerface [19], which is often found in large-mismatched metal-oxide heterointerface systems [21,22]. Figure 2(g) shows a histogram of the formation of coherent or incoherent interfaces as a function of the projected Au nanoparticle size. Although the thickness of the Au nanoparticles along the beam direction is difficult to precisely estimate, our systematic observations strongly suggest that there is a structural transition at the Au/TiO$_2$ (110) interface for Au particles around 2–3 nm in size.

To verify the structural transition at the Au/TiO$_2$ interfaces theoretically, first-principles DFT calculations were performed on model Au(110) || TiO$_2$(110) interfaces, whose interface orientation is based on the one estimated from the STEM images in Fig. 2. The projector augmented wave method, implemented in the VIENNA $ab$ initio simulation package, was used [23]. The electron exchange-correlation potential was described by the generalized gradient approximation in the form of Perdew, Burke, and Ernzerhof [24]. Spin polarization was included throughout the calculations. Brillouin-zone sampling using a $1 \times 8 \times 1$ Monkhorst-Pack mesh [25] (four irreducible $k$ points) was performed for the Au/TiO$_2$ supercells. It was confirmed that these conditions ensured a good total energy convergence of 1 meV/atom for the Au/TiO$_2$ supercells.

To estimate the effect of Au nanoparticle mass on the interface structure, two interface models with two (thin) and nine (thick) Au atomic layers stacked on the TiO$_2$ surfaces were simulated [26]. To allow for the large interfacial misfits in the simulation, the TiO$_2$ (110) and Au (110) surface units were extended by $1 \times 5$ and $1 \times 8$, respectively, to minimize the total lattice mismatch between the extended TiO$_2$ and Au slabs in the initial structure. According to our systematic DFT calculations, the stable termination of TiO$_2$ in the interface formation is strongly dependent on the external atmosphere and the reduced TiO$_2$ surface termination is more stable than the stoichiometric one in the reducing atmosphere [27]. Because our Au deposition was done in a reducing atmosphere, the reduced TiO$_2$ (110) termination model for the initial Au/TiO$_2$ interface models was used. In fact, the reduced interface model can reasonably reproduce the Au island structure observed by STEM. DFT calculations of the same two-layer Au on the stoichiometric TiO$_2$ (110) surface were also performed, but the results were inconsistent with our experimental observation.

Figures 3(a) and 3(b) show the relaxed structures of the respective interface models. In the case of the two-layer model, Au atoms significantly relax their positions: Au atoms in the first layer move to the on-top sites of the Ti-O columns (O$_{br}$ vacancy sites), whereas Au atoms in between these sites merge into the top surface layer. As a result, the Au structure shows the unique structural feature that the second Au layer is attached on the first Au layer rigidly anchored by the TiO$_2$ substrate. The Au atomic positions in the epitaxial islands are consistent with our experimental observations, which can also be confirmed from the STEM image simulation shown in the inset.

In the case of the nine-layer model, the relaxed structure shows no obvious lattice coherency at the Au/TiO$_2$ inter-
onto the atomic layers across the Au/TiO$_2$ interface. The interface loses lattice coherency in order to accommodate the top surface layer of the reconstructed Au structure. Moreover, electron transfer from the reduced TiO$_2$ surface to the Au layers is observed. The reconstructed electronic structure of the two-layer Au is thus completely different from that of bulk Au. In contrast, the electron density change in the Au structure in the nine-layer model is much smaller and limited to within a few atomic layers from the interface. Above those interface layers, the electron density of Au becomes similar to that of the bulk Au. These results suggest that the electronic structure of Au nanoparticles is strongly influenced by the structure of the nanoscale heterointerface.

In summary, the present study demonstrates that the atomic structure of Au nanoparticles is strongly influenced by the supporting TiO$_2$ surfaces. When the Au nanoparticles are very small and atomically thin, the strong templating behavior of the TiO$_2$ surface reconstructs both the atomic and the electronic structure of the Au nanoparticles. When the Au nanoparticle size becomes larger and thicker, the interface loses lattice coherency in order to accommodate large lattice mismatch, and the support effect rapidly decays across the interface. Our systematic observations thus clearly show that there is a structural transition at the Au/TiO$_2$ (110) interface depending on the Au particle size. We believe the strong interaction between nanosized Au and oxide support via unique nanoscale heterointerface structure will be an essential factor in understanding the origin of size-dependent catalytic activity of Au nanoparticles supported on metal oxides.

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