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Elucidation of active sites for the reaction of ethanol on TiO$_2$/Au(111)

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Elucidation of Active Sites for the Reaction of Ethanol on TiO$_2$/Au(111)

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An Honors College Project Presented to

the Faculty of the Undergraduate

College of Science and Mathematics

James Madison University

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by David Thomas Boyle

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Accepted by the faculty of the Department of Chemistry and Biochemistry, James Madison University, in partial fulfillment of the requirements for the Honors College.

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PUBLIC PRESENTATION

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Abstract

Obtaining a molecular-level understanding of the reaction of alcohols with heterogeneous model catalysts is critical for improving industrial catalytic processes, such as the production of H₂ from alcohols. Gold has been shown to be an excellent oxidation catalyst once oxygen is added to it. The use of reducible oxides provides a source of oxygen on Au(111) for the reaction of ethanol, which is easily regenerated in the presence of an oxygen background. In this work, ethanol operates as a probe molecule to investigate the role of Au(111), TiO₂ nanoparticles, and TiO₂/Au interfacial surface sites on the catalytic properties of TiO₂/Au(111). Ultrahigh vacuum temperature programmed desorption (TPD) studies with ethanol/Au(111) elucidate previously unreported adsorption sites for ethanol. Ethanol molecularly adsorbs to Au terrace sites, step edges, and undercoordinated kink sites with adsorption energies of -51.7, -55.8, and -65.1 kJ/mol, respectively. A TPD coverage study of ethanol on TiO₂/Au(111) indicates ethanol undergoes dissociative adsorption to form H*(a) and CH₃CH₂O*(a) on the inverse model catalyst surface. The desorption temperature of low coverages of ethanol from TiO₂/Au(111) (T_{des} = ~235 K) is at an intermediate temperature between the desorption temperatures from bulk Au(111) and TiO₂(110), indicating both Au and TiO₂ play a role in the adsorption of ethanol. Both low-temperature adsorption and high-temperature reactions are studied and indicate that ethanol-derived products—such as acetaldehyde and ethylene—desorb from TiO₂/Au(111) at ~500 K.

All the work reported herein is published in the Journal of Physical Chemistry C:
Introduction

Heterogeneous Au/TiO$_2$ model catalysts have been extensively studied due to their remarkable catalytic activity toward the water-gas shift reaction,$^{1-4}$ production of H$_2$ from alcohols,$^{5-8}$ oxidation of CO to CO$_2$,$^{4,9-11}$ and several other applications.$^1$ It has been proposed that the interface between Au nanoparticles and TiO$_2$ is the catalytically active site of Au/TiO$_2$ heterogeneous catalysts for the conversion of small alcohols to their respective carbonyls and alkenes, water-gas shift reaction, and H$_2$ dissociation.$^{4,9,10,12,13}$ In order to determine the role of distinct Au, TiO$_2$, and Au/TiO$_2$ interfacial sites in Au/TiO$_2$ heterogeneous catalysts, inverse model catalysts have become a material of interest. Using inverse model catalysts in which TiO$_2$ nanoparticles are deposited onto a Au(111) substrate allows for the use of surface science techniques to investigate the role of distinct surface sites on the overall reactivity of the TiO$_2$/Au(111) catalyst.

Ethanol is the simplest molecule containing C—H, C—O, C—C, and O—H bonds, which makes it an excellent probe molecule to elucidate active sites on TiO$_2$/Au(111) inverse model catalyst surfaces. Additionally, ethanol is particularly interesting to the catalytic community, as it is used as an alternative fuel and chemical feedstock for important industrial reactions like the production of H$_2$ and acetic acid.$^{14}$ Ethanol has been utilized to probe the catalytic activity of several Au/TiO$_2$-based surfaces including: O/Au(111),$^{15,16}$ TiO$_2$(110),$^{17-20}$ and Au/TiO$_2$(110)$^1$. It is well understood that ethanol undergoes molecular adsorption on clean Au(111) and dissociative adsorption on O/Au(111) and bulk TiO$_2$(110) to form ethoxy and hydroxyl adsorbates.$^{15-20}$ The reaction of ethanol on bulk rutile-TiO$_2$(110) single crystals showed that both acetaldehyde and ethylene were produced at ~620 K.$^{20}$ The formation of a carbonyl group versus an alkene was dependent on the number of defects in the crystal surface, as an increased concentration of oxygen vacancies promoted the formation of ethylene.$^{20}$
Interestingly, low temperature ethanol adsorption and ethanol chemistry on TiO₂/Au(111) surfaces has not yet been reported. Studies with other simple alcohols methanol²¹ and 2-propanol²² on TiO₂/Au(111) inverse model catalysts showed that the alcohols underwent an elimination or oxidation reaction depending on the method of surface preparation. Both studies proposed that the formation of the oxidized products—formaldehyde and acetone—were the result of excess oxygen on the surface. The formation of elimination products—methane and propene—were the result of undercoordinated Ti cations within the TiO₂ nanoparticles supported on Au(111). While TiO₂/Au(111) showed catalytically valuable chemistry toward methanol and 2-propanol, the role of distinct Au, TiO₂, and Au/TiO₂ interfacial sites on the overall reactivity of the surface was not reported.

In this work, a comprehensive study of low temperature ethanol adsorption and chemistry over Au(111) and TiO₂/Au(111) is reported across a wide temperature range to determine the catalytically active sites of TiO₂/Au(111). The role of distinct Au, TiO₂, and Au/TiO₂ interfacial sites is investigated through the use of ultrahigh vacuum temperature-programmed desorption (UHV-TPD), atomic force microscopy (AFM), and X-ray photoelectron spectroscopy (XPS). Low-temperature desorption relays information about binding energies and adsorption sites of ethanol to Au(111) and TiO₂/Au(111), and high-temperature desorption is used to investigate the chemistry of the ethanol over the TiO₂/Au(111) surface. The results shown here demonstrate the TiO₂–Au interface as the active catalytic site on inverse model catalysts for the reaction of ethanol.
Methods

Experiments were performed in a UHV-TPD chamber. The UHV chamber is pumped by turbo and titanium sublimation pumps and maintained a base pressure of ~5 x 10⁻¹⁰ Torr during experiments. The chamber contains a quadrupole mass spectrometer (Hiden HAL3F-RC) for TPD and residual gas analysis. The heating rate for TPD experiments was 1.05 K/s. Collimated high precision leak valves were used for controlling gas doses into the chamber.

A Au(111) single crystal (Princeton Scientific) was mounted on a sample holder and held vertically by tungsten wires (Goodfellow, 99.95%, 0.25 mm diameter) between two copper blocks in the center of the chamber, which allowed the sample to be resistively heated and cooled with liquid nitrogen. Temperature measurements were recorded with a k-type thermocouple that was directly attached to a hole in the side of the Au(111) crystal. The Au(111) surface was prepared via cycles of 1 keV Ar⁺ sputtering and annealing in UHV at ~850 K. The cleanliness of the surface was determined via Auger electron spectroscopy (AES), TPD of probe molecules, ex situ AFM, and XPS. TiO₂ was deposited on the Au(111) single crystal by the physical vapor deposition of Ti at an oxygen background pressure of 2 x 10⁻⁷ Torr at room temperature, and then annealed in O₂ to 600 K. For the physical vapor deposition, a homemade Ti filament made with Ti wire (0.125 mm diameter, 99.6 %, Goodfellow) was used. Monolayer coverages of TiO₂ were controlled by monitoring the power input and deposition time and determined by ex situ AFM and XPS. Ex situ AFM images were recorded by a Digital Instruments MultiMode AFM in tapping mode under ambient conditions. The saturation coverage of ethanol is defined as 1 monolayer (ML) and was calibrated using TPD. XPS was performed on a PerkinElmer 5400 X-ray Photoelectron Spectrometer equipped with a monochromatized Mg(Kα) radiation source (1253.6 eV) and a position sensitive, multichannel
plate detector. Measurements were taken at a pressure of $\sim 10^{-7}$ mbar and with a takeoff angle of 15° with respect to the surface normal. Binding energies are referenced to Au 4f$_{7/2}$ at 84.0 eV.
**Results and Discussion**

Prior to studying the low-temperature adsorption and chemistry of ethanol on TiO$_2$/Au(111), a comprehensive understanding of ethanol adsorption on clean Au(111) is critical for interpreting the role of each individual material on the reactivity of the composite inverse model catalyst. In this work, the coverage-dependent TPD studies of ethanol/Au(111) elucidate previously unreported adsorption sites for ethanol on Au(111). Figure 1 shows TPD results from ethanol coverages below 0.20 ML, where a m/z ratio of 31 was followed, as it is the most intense fragment for ethanol. At ethanol coverages below 0.20 ML, the TPD spectra show two desorption features with desorption temperatures of $T_{des}=172$ and 185 K. The desorption feature at $T_{des}=172$ K is related to ethanol desorbing from the Au(111) terrace, and is in agreement with previous work on ethanol/Au(111). The consistency in $T_{des}=172$ K as the monolayer coverage of ethanol increases indicates first-order kinetics of desorption, which confirms that ethanol adsorbs and desorbs molecularly from the Au(111) terrace. Interestingly, TPD spectra of low ethanol coverages on Au(111) show a desorption feature at $T_{des}=185$ K. At ethanol coverages as low as 0.01 ML, the desorption feature at $T_{des}=185$ K predominates with a slight low-temperature shoulder, indicating that the adsorption site corresponding to $T_{des}=185$ K is preferentially populated prior to Au(111) terrace sites. Upon increasing the coverage of ethanol above 0.01 ML, the desorption feature at $T_{des}=185$ K saturates, and the desorption feature at $T_{des}=172$ K, representing ethanol desorption from Au(111) terrace sites, continues to grow in intensity up to a coverage of 0.20 ML of ethanol.
Figure 1. TPD spectra of low coverages of ethanol (m/z = 31) from Au(111). The peak at $T_{\text{des}}=172$ K represents submonolayer coverages of the molecular desorption of ethanol from Au terraces, and the peak at $T_{\text{des}}=185$ K is assigned to the molecular desorption of ethanol from step edges.

The observation of two distinct desorption features in the TPD spectra indicates that there are two distinct adsorption sites for ethanol on Au(111). Previous studies of CO/Au(111) demonstrated that a sputtered Au(111) surface enhanced the adsorption of CO by exposing more undercoordinated Au step edges and kink sites.\textsuperscript{23–28} CO had stronger binding energies to undercoordinated Au sites, which resulted in a higher $T_{\text{des}}$.\textsuperscript{27} In order to investigate the desorption...
of ethanol from undercoordinated Au sites, the clean Au(111) surface was sputtered by bombarding the surface with 1 keV Ar$^+$ ions at 300 K. The morphology of a sputtered Au(111) single crystal was imaged using \textit{ex situ} AFM, which showed the presence of large pits with roughened step edges (Figure S1). The sputtered surface results in an ethanol TPD spectrum with noticeable differences from the clean Au(111) surface (Figure 2). Upon sputtering the surface, the desorption feature at $T_{\text{des}}=185$ K grows in intensity, whereas the desorption feature at $T_{\text{des}}=172$ K decreases in intensity. Consequently, the peak at $T_{\text{des}}=185$ K is assigned to the desorption of ethanol from undercoordinated Au sites. Annealing Au(111) to 400 K and above decreases the number of undercoordinated Au sites by smoothing the sputtered surface.$^{27,29}$ After annealing the sputtered Au(111) surface to 400 K, the desorption feature at $T_{\text{des}}=172$ K increases in intensity, while the desorption feature for undercoordinated Au sites ($T_{\text{des}}=185$ K) decreases, indicating that the 400 K anneal results in a smoother Au(111) surface. While sputtering the surface results in an increased number of undercoordinated sites of both step edges and kink sites,$^{23,25-28}$ the peak at 185 K is assigned to Au step edges, as the concentration of step edges is expected to be much greater than that of kink sites.
Figure 2. TPD spectra of ethanol on clean and sputtered Au(111) surfaces (m/z = 31). Annealing the surface to 400 K partially smooths the surface, which results in a decrease in ethanol desorption at $T_{\text{des}}=185$ K (step edges) and an increase in ethanol desorption from the terraces at $T_{\text{des}}=172$ K.

The adsorption energies of ethanol on Au(111) terrace sites and Au step edges can be calculated using the Redhead approximation for first-order desorption

$$E_{\text{des}} = RT_{\text{des}} \left[ \ln \left( \frac{T_{\text{des}}}{\beta} \right) - 3.46 \right]$$

where $\beta$ is the heating rate of $1.05 \pm 0.07$ K/s and $\nu$ is the pre-exponential factor. A pre-exponential factor of $10^{15}$ was used to calculate adsorption energies of -0.54 eV (-51.7 kJ/mol) and -0.58 eV (-55.8 kJ/mol) for ethanol adsorbed to terrace Au(111) sites and Au step edges,
respectively. The calculated adsorption energy on Au(111) terraces of -0.54 eV is in agreement with previous DFT studies, which resulted in an adsorption energy of -0.536 eV.\textsuperscript{30} The same DFT calculations, which included a van der Waals correction, predict that ethanol adsorbs to the Au(111) surface with the C–C bond parallel to the surface.\textsuperscript{30}

To further investigate the desorption of ethanol from Au(111), TPD studies of ethanol coverages above 1.0 ML were performed on clean Au(111) (Figure 3). As the coverage of ethanol increases, the Au terrace peak ($T_{des}=172$ K) increases in intensity and saturates at 1.0 ML of ethanol. It is important to note that the peak previously observed for step edges (185 K) is now obscured by the terrace desorption peak. Additionally, upon increasing the ethanol coverage above 1.0 ML, low-temperature peaks corresponding to multilayered ethanol appear ($T_{des} < 155$ K). The desorption feature corresponding to multilayered ethanol increases to a maximum desorption temperature of $T_{des}=150$ K, until a low-temperature shoulder appears at $T_{des}=145$ K above 1.8 ML. The desorption of multilayered ethanol at $T_{des}=150$ K and ethanol from Au(111) terraces at $T_{des}=172$ K is consistent with previous reports.\textsuperscript{15} The desorption peaks at $T_{des}=150$ and $T_{des}=145$ K are assigned as multilayers of ethanol. Previous work with methanol exhibited more than one multilayer peak on Au(111),\textsuperscript{31} Ag(111),\textsuperscript{32} and Pt(111),\textsuperscript{33} which were labeled as amorphous and crystalline phases due to the early work of Ehlers using UV photoemission and IR reflection to differentiate between multilayer structures.\textsuperscript{33} However, ethanol multilayer studies on highly oriented pyrolytic graphite (HOPG) resulted in two desorption peaks ($T_{des}=145$ K and $T_{des}=142$ K), labeled as a bilayer and multilayer of ethanol, respectively.\textsuperscript{34} We cannot rule out one assignment over another in this case to differentiate the peaks; therefore, we label desorption peaks below $T_{des}=155$ K as multilayers of ethanol.
Figure 3. TPD spectra of a series of ethanol coverages (m/z = 31) at saturation coverage (1.0 ML) and above on Au(111). Desorption peaks for ethanol from multilayers \((T_{des} < 155 \text{ K})\), Au terrace sites \((T_{des}=172 \text{ K})\), and undercoordinated Au kinks \((T_{des}=215 \text{ K})\) are present.

In addition to desorption peaks for multilayers of ethanol and ethanol adsorbed to Au(111) terraces, a small peak at \(T_{des}=215 \text{ K}\) appears with coverages of ethanol above 1 ML, as seen in Figures 3 and 4. The adsorption energy for ethanol desorbing at 215 K, calculated using the Redhead approximation described above, is -0.67 eV (-65.1 kJ/mol). The difference in adsorption energy between Au step edges and the higher temperature desorption peak \((T_{des}=215 \text{ K})\) is -9.3 kJ/mol (-0.096 eV), which is in excellent agreement with the calculated difference in adsorption energy between step edges and kink sites on Au(111) for CO. Therefore, the
desorption peak at $T_{des}=215$ K is labeled as ethanol desorption from undercoordinated kink sites on Au(111) step edges.

![Temperature-Programmed Desorption (TPD) spectrum](image)

**Figure 4.** TPD, m/z = 31. Zoomed in perspective of desorption features corresponding to terrace Au sites (172 K) and Au kink sites (215 K). At ethanol coverages >1 ML, desorption from kink sites increases and desorption from terrace sites decreases.

It is expected that undercoordinated Au sites, including step edges and kinks, would act as preferential binding sites due to their increased adsorption energy and desorption temperature compared to terrace Au sites. Interestingly, Figures 3 and 4 show an opposite trend for the population of kink sites ($T_{des}=215$ K) and terrace sites ($T_{des}=172$ K) as the coverage of ethanol sequentially increases. At an ethanol coverage of 1.0 ML, the only observed desorption peak is
172 K. As the coverage of ethanol increases, the desorption peak at 172 K decreases in intensity and shifts to lower temperatures while the kink site desorption peak appears ($T_{des} = 215$ K) and continually increases (Figure 4). We hypothesize that the shift to lower temperatures of the 172 K desorption peak arises from the depletion of step edge desorption sites ($T_{des} = 185$ K) to form kink desorption sites. These results indicate that the presence of high coverages of ethanol roughen Au(111) step edges below room temperature, increasing the concentration of undercoordinated kink sites ($T_{des} = 215$ K). In a separate set of experiments, the coverage of ethanol was increased to >1 ML to produce the kink site desorption feature. Subsequently, a 1 ML ethanol dose was exposed to the surface, and the kink site desorption peak was not observed, indicating that the higher coverages of ethanol lead to the roughening of Au(111) step edges. While future studies will be required to understand the mechanism behind the formation of kinks, it is speculated that the interaction between ethanol molecules and surface Au atoms will decrease the Au–Au bond energy, which would thereby decrease the barrier for Au diffusion. While we do not suspect the interaction between ethanol and Au is strong enough to lift the herringbone reconstruction, it is plausible that Au step edge atoms, which exhibit mobility at room temperature on clean Au(111), will have an enhanced mobility in the presence of ethanol, resulting in the rearrangement of step edge atoms to form kinks. Furthermore, the high coverage of ethanol required to visualize the kink site desorption feature indicates that intermolecular repulsion of ethanol molecules may also play a role in the destabilization of Au–Au bonds. It was previously reported that intermolecular repulsion influenced the molecular packing of methanol on Au(111) even at submonolayer coverages.\textsuperscript{35}

The rearrangement and restructuring of Au surface atoms has been observed at low pressures and temperatures after exposing small molecules to Au(111).\textsuperscript{36–38} It is expected that
ethanol roughens the surface of reducible oxides. However, the oxide is reduced by ethanol via a reaction with the lattice oxygen and acidic hydrogens from ethanol to form hydroxyl groups, which recombine to form water and leave oxygen vacancies in the surface.\textsuperscript{39–41} The TPD spectra in Figures 3 and 4 therefore demonstrate the first reported roughening of the Au(111) step edges by ethanol at low temperatures. Future imaging studies will allow us to investigate the morphology of the Au(111) before, during, and after the desorption of ethanol, to investigate the reversibility of the kink formation. Additionally, imaging studies will be conducted to visualize the change in surface morphology due to ethanol adsorption at high coverages, as well as to identify a mechanism for atomic rearrangement at the step edges.

The elucidation of ethanol adsorption and desorption from distinct Au(111) surface sites allows for the subsequent study of low-temperature ethanol adsorption on the TiO\textsubscript{2}/Au(111) inverse model catalyst. Several preparations of Au(111)–supported TiO\textsubscript{2} nanoparticles are reported in the literature.\textsuperscript{42–46} For the experiments reported herein, TiO\textsubscript{2} nanoparticles were deposited onto Au(111) via physical vapor deposition in an O\textsubscript{2} background (2x10\textsuperscript{-7} torr) at 300 K and annealed to 600 K in O\textsubscript{2} prior to experiments. \textit{Ex situ} AFM and \textit{ex situ} XPS confirmed the successful deposition of TiO\textsubscript{2} nanoparticles on the Au(111) surface as compared to clean Au(111) (Figure 5, S2). Figure 5A shows an \textit{ex situ} AFM image of the Au(111) surface prior to the deposition of TiO\textsubscript{2} nanoparticles, with Au step edges running diagonally across the image. Figures 5B and 5C show TiO\textsubscript{2} nanoparticles well-dispersed across the Au terrace and step edges. Analysis of the XPS data and the Au 4f peak attenuation confirmed a TiO\textsubscript{2} coverage of ~0.6 ML on the Au(111) surface. The average diameter of the TiO\textsubscript{2} nanoparticles is 10.1 ± 2.2 nm, and a particle distribution histogram is shown in Figure S3.
**Figure 5.** *Ex situ* AFM images of (A) Au(111) and (B, C) ~0.6 ML TiO$_2$/Au(111). TiO$_2$ nanoparticles are highly dispersed across the Au(111) surface, residing on Au terraces and step edges.

TPD experiments of ethanol on TiO$_2$/Au(111) showed significantly different results from those of the clean Au(111) surface. A coverage dependent study of ethanol on ~0.7 ML TiO$_2$/Au(111) indicated several differences from clean Au(111) TPD spectra (Figure 6). First,
the $T_{\text{des}}$ of ethanol from the TiO$_2$/Au(111) surface trends toward lower $T_{\text{des}}$ with increasing coverages of ethanol. Ethanol desorption from the TiO$_2$/Au(111) surface is $T_{\text{des}}=\sim 235$ K after only 0.3 L of ethanol exposure, whereas ethanol desorbs from the same TiO$_2$/Au(111) surface at $T_{\text{des}}=\sim 195$ K after 2.0 L of ethanol exposure. This trend suggests that ethanol undergoes dissociative adsorption on the surface into an adsorbed H and ethoxy group. The dissociative adsorption of ethanol was expected on the TiO$_2$/Au(111) surface as previous work with aliphatic alcohols on bulk TiO$_2$(110) showed that ethanol dissociates into H and ethoxy on the surface.$^{15,20}$

$$\text{CH}_3\text{CH}_2\text{OH}(g) \rightarrow \text{H}^*(a) + \text{CH}_3\text{CH}_2\text{O}^*(a)$$

The dissociative adsorption of ethanol on TiO$_2$/Au(111) indicates the potential for the addition of other reactant molecules to react with the activated H and ethoxy species on the surface.
Figure 6. Coverage-dependent study of ethanol/TiO$_2$/Au(111) TPD, m/z = 31. The trend in TPD spectra indicates dissociative adsorption of ethanol to H and ethoxy species.

The $T_{des} \approx 235$ K of ethanol from TiO$_2$/Au(111) is at an intermediate temperature between those of ethanol desorption from clean Au(111) (172 K, Figure 3) and bulk TiO$_2$(110) at 300-400 K. The intermediate $T_{des}$ of ethanol from the TiO$_2$/Au(111) surface indicates that both Au and TiO$_2$ sites may contribute to the adsorption of ethanol. As the size of TiO$_2$ decreases from bulk to nanoparticles, the number of undercoordinated atoms increases as the surface area increases, and it is expected that the desorption temperature at the undercoordinated atoms of TiO$_2$ would be higher than that of bulk TiO$_2$. Instead, in Figure 6, the desorption temperature related to TiO$_2$ sites is lower than that of bulk TiO$_2$, indicating that the ethanol adsorbs to
interfacial sites between the TiO₂ and Au. Previous work with ethanol/CeO₃/Au(111) proposed
that the oxide/metal interface played a strong role in the catalytic activity of the material toward
ethanol using XPS. The intermediate T_{des} of ethanol from TiO₂/Au(111) suggests that the
interface between the TiO₂ nanoparticles and the Au(111) substrate plays a role in the conversion
of ethanol to ethoxy.

The TPD spectra of higher coverages of ethanol on TiO₂/Au(111) indicate that the
presence of TiO₂ increases the total adsorption of ethanol (Figure 7). Ethanol desorption from
clean Au(111) has a sharp desorption peak in the TPD spectrum that occurs at T_{des}=172 K.
Interestingly, the presence of TiO₂ nanoparticles results in a broad higher temperature peak
centered around T_{des}≈190 K with a long high temperature tail. We hypothesize that the increase
in total adsorbed ethanol which is demonstrated by increased area and broadening of the
desorption peak, could be the result of several factors: (1) The presence of TiO₂ could increase
the total number of adsorption sites on the surface. Figure 5 C shows that the Au(111) substrate
has several step edges across the surface and TiO₂ nanoparticles could pin the step edges during
annealing cycles. TiO₂ pins the Au step edges and results in a roughened surface with an
increased number of ethanol adsorption sites on Au in addition to the increased number of
adsorption sites on TiO₂ nanoparticles. Figure S4 shows the peak fitting for the
ethanol/TiO₂/Au(111) spectrum in Figure 7. The peak fitting suggests that the low-temperature
desorption features occur at temperatures that correspond well with our assignments for ethanol
adsorbed on terrace Au (T_{des}= 175 K), step edges (T_{des}= 195 K), kink sites (T_{des}= 217 K), and a
new peak (T_{des} = 265 K) corresponding to TiO₂ (Figure S4). (2) The presence of TiO₂
nanoparticles potentially disrupts the ordered packing of ethanol on the exposed Au(111)
surface. Previous work with methanol on Au(111) showed that H-bonding significantly
influenced the packing structure of methanol on the surface. The presence of TiO$_2$ nanoparticles on the surface causes the dissociative adsorption of ethanol, which could disrupt H-bonding between ethanol molecules on the surface and allow for the enhanced total adsorption of ethanol by tightly packing ethanol on the surface. Furthermore, the stronger adsorption energy between TiO$_2$ and ethanol could influence the density of the molecules on the nanoparticles, as compared to on the Au surface, where the H-bonded networks will dominate the packing density.

![TPD spectra](image)

**Figure 7.** TPD spectra of ethanol desorption from clean Au(111) (black line) and nanoparticle TiO$_2$/Au(111) (red line). The increased area under the red curve suggests an increase in the distribution of adsorption sites as well as the concentration of surface-bound molecules.
Temperature-programmed reaction spectroscopy (TPRS) experiments of ethanol/TiO$_2$/Au(111) show that the nanoparticle TiO$_2$/Au(111) surface is active for the transformation of ethanol into both ethylene and acetaldehyde (Figure 8). Previous work with methanol and 2-propanol demonstrated that both alcohols underwent redox chemistry that depended on the specific method of surface preparation and the oxidation state of TiO$_2$.\textsuperscript{21,22} The preparation of the Au(111) supported TiO$_2$ nanoparticles was consistent for the TPRS experiments described herein. Prior to TPRS experiments, the TiO$_2$/Au(111) surface was oxidized at 600 K with an O$_2$ background of 2 x 10$^{-7}$ torr. Several m/z ratios were monitored during experiments to elucidate different products, as several m/z ratios for potential reaction products overlap. The main species monitored were ethanol (m/z=31, 46, 45), ethylene, (m/z=28, 27, 26), and acetaldehyde (m/z=29, 44, 43). Coupling products as well as smaller molecules resulting from the cleavage of C—C bonds were not detected. Six representative m/z ratios are shown to represent ethanol (m/z=46), acetaldehyde (m/z=29, 43, 44), ethylene (m/z=27), and water (m/z=18). The left panel displays the full temperature range of the TPR spectrum, which expectedly shows the desorption of ethanol over the lower temperature range between 170 and 250 K.
Figure 8. TPR spectrum shows the ethanol–derived products formed over TiO$_2$ nanoparticles supported on Au(111). The left panel shows low-temperature ethanol desorption, and the right panel highlights the product formation at $T_{des}$ $\approx$ 500 K.

The right panel highlights the high-temperature component of the TPR spectrum. This data indicates the formation of acetaldehyde (m/z=29, 44, 43), ethylene (m/z=27), and water (m/z=18) from ethanol (m/z=46) over the TiO$_2$/Au(111) surface. While m/z=29 corresponds to both ethanol and acetaldehyde, the combination of high $T_{des}$ peaks for m/z=29, 44, 43 with the lack of a high $T_{des}$ peak for m/z=46 shows that the desorption features are not the result of ethanol fragmentation but rather that acetaldehyde is in fact a high $T_{des}$ product. Similarly, a m/z ratio of 27 is observed for both ethanol and ethylene, but the lack of high-temperature desorption...
of m/z=46 indicates that the desorption of m/z=27 corresponds to ethylene. The desorption temperatures of acetaldehyde and ethylene are significantly lower on the TiO$_2$/Au(111) surface ($T_{\text{des}} = \sim 500$ K) as compared to those on the TiO$_2$(110) surface ($T_{\text{des}} = \sim 620$ K). Sequential ethanol TPR experiments did not result in a change in a distribution in the ethanol–derived products, i.e., after multiple experiments the ratio between oxidized (acetaldehyde) and reduced (ethylene) products did not change, indicating that there was not a significant oxidation or reduction of the surface due to ethanol reaction. We hypothesize that ethanol deprotonates at the interface to form ethoxy and adsorbed H at low temperatures. Future work will extend to isotopic studies, as well as altering coverages and surface preparations of TiO$_2$ to discern the effect of TiO$_2$ nanoparticle size and surface condition on the reaction pathway for the reaction of ethanol.
Conclusions

The desorption of ethanol from multilayers, Au terraces, Au step edges, and Au kink sites were differentiated via coverage-dependent TPD studies. Upon the deposition of TiO$_2$ nanoparticles on Au(111), a broadening of the low-temperature ethanol desorption peak is observed with a maximum desorption temperature of $T_{des}$ $\approx$ 235 K. The desorption of ethanol from TiO$_2$/Au(111) occurs at an intermediate temperature between bulk Au(111) and TiO$_2$(110). This intermediate desorption temperature indicates that ethanol dissociatively adsorbs into ethoxy and adsorbed hydrogen at the interface between TiO$_2$ nanoparticles and Au(111) prior to forming acetaldehyde or ethylene.
Ex situ AFM images show the morphology of the clean (left panel) and roughened (right panel) Au(111) surface. The roughened AFM image has roughly 50 kinks in the 1000 nm square image on the right. While this is not a true count of the total number of kinks in the surface, the shape of the edges allows for the rough estimation of the low end number of kinks per area.

**Figure S1.** *Ex situ* AFM images of Au(111) (left) and roughened Au(111) (right). The roughened Au(111) was sputtered and annealed to 600 K.
Figure S2. *Ex situ* XPS: Clean Au(111) (Blue) and 0.6 ML TiO$_2$/Au(111) (Orange).
Figure S3. *Ex situ* AFM Analysis: Histogram of particle diameters for nanoparticle TiO$_2$ coverage (0.6 ML) on a Au(111) surface. The average diameter of the TiO$_2$ nanoparticles is 10.1 ± 2.2 nm.
Peak analysis indicates that the broad low $T_{\text{des}}$ is the result of several desorption peaks for ethanol, which make up the overall peak (Figure 7). The formation of several new peaks for ethanol with a higher $T_{\text{des}}$ indicates that TiO$_2$ forms new adsorption sites with higher binding energies to ethanol. Analysis of the underlying peaks shows that a portion of the broad low temperature peak is made up of ethanol desorption from the Au(111) terrace (175 K), Au step edges (195 K), Au kink sites (217 K) and TiO$_2$ nanoparticle sites (265 K) (Figure S4).

**Figure S4.** Peak fitting of ethanol desorption from ~0.3 ML TiO$_2$/Au(111). The peaks correspond to ethanol adsorbed to the Au(111) terrace (175 K), Au step edges (195 K), Au kink site (217 K), and TiO$_2$ site (265 K).
References


(9) Grunwaldt, J.-D.; Baiker, A. Gold/Titania Interfaces and Their Role in Carbon Monoxide


(18) Kim, Y. K.; Kay, B. D.; White, J. M.; Dohnálek, Z. Alcohol Chemistry on Rutile


15384–15385.


