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Formation of Highly Ordered Self-Assembled Monolayers of Alkynes on Au(111) Substrate

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Supporting Information Placeholder

ABSTRACT: Self-assembled monolayers (SAMs), prepared by reaction of terminal *n*-alkynes ($\text{HC}\equiv\text{C}(\text{CH}_2)_n\text{CH}_3$, $n = 5, 7, 9$, and 11) with Au(111) at 60°C were characterized using scanning tunneling microscopy (STM), infra-red reflection absorption spectroscopy (IRRAS), X-ray photoelectron spectroscopy (XPS), and contact angles of water. In contrast to previous spectroscopic studies of this type of SAMs, these combined microscopic and spectroscopic experiments confirm formation of highly-ordered SAMs having packing densities and molecular chain orientations very similar to those of alkanethiolates on Au(111). Physical properties—hydrophobicity, high surface order, and packing density—also suggest that SAMs of alkynes are similar to SAMs of alkanethiols. The formation of high-quality SAMs from alkynes requires careful preparation and manipulation of reactants in an oxygen-free environment: trace quantities of oxygen lead to oxidized contaminants and disordered surface films. The oxidation process occurs during formation of the SAM; via oxidation of the $-\text{C}\equiv\text{C}-$ group (most likely catalyzed by the gold substrate in the presence of O_2).

Thin organic films based on self-assembled monolayers (SAMs)¹ are ubiquitous in surface science. The reaction of organic thiols (RSH) with group Ib metals (Au and Ag) to generate SAMs with composition Au/AgSR is the reaction most commonly used to prepare SAMs,¹ although reactions that generate organosilanes on silicon² (SiR) and organic carboxylates on silver³ (AgO_2CR) have attractive properties, and a number of other precursors have been surveyed. There have also been scattered descriptions of SAMs formed on gold from solutions of alkynes⁴ ($\text{HC}\equiv\text{C}(\text{CH}_2)_n\text{CH}_3$, $n = 3, 5, 7, 9, 11$, and 13), ethynylbenzene⁵ ($\text{HC}\equiv\text{CC}_6\text{H}_5$) or *n*-alkylmercury(II) tosylates⁶ ($\text{CH}_3(\text{CH}_2)_n\text{HgOTs}$, $n = 4$ and 18) on Au(111). Although the potential interest of SAMs having Metal–C $\equiv$ CR bonds is high, since they offer a new type of metal-organic bond, most of these studies have used preparations analogous to those employed with *n*-alkanethiols, and have generated SAMs that have not seemed to be highly ordered, and are thus, perhaps, unsuitable for detailed studies of the physical chemistry of the surface. In particular, there are no procedures that describe the formation of SAMs that are highly ordered in two dimensions—a key requirement for high-quality surface science. The most recent analyses of *n*-alkyl-based SAMs on Au(111) indicate a “liquid-like” structure of the monolayer,⁶ and XPS analyses of SAMs formed from alkynes^{4,5} suggest that these

SAMs are sensitive to oxidation at an undefined point in their formation; that is, oxidation occurs either during or after SAM formation (for example, by reaction of the $\text{AuC}\equiv\text{CR}$ bond with O_2). Contact angle analyses of increasing lengths of alkynes ($\text{HC}\equiv\text{C}(\text{CH}_2)_n\text{CH}_3$, $n = 5, 7, 9$ and 11) also suggest⁴ that the quality of these SAMs is lower than those based on *n*-alkanethiols.

Although SAMs have enabled studies of wetting,^{7,8} adhesion,^{9,10} and charge transport^{3,11–13} (inter alia), most of this work has focused on the terminal part of the SAM that is exposed to air, and there is relatively little work devoted to understanding the contribution of the anchoring groups of the SAM (as opposed to the terminal group, the thickness, or the electronic structure). In the current work, we characterize the SAMs formed by reactions of *n*-alkynes ($\text{HC}\equiv\text{C}(\text{CH}_2)_n\text{CH}_3$, $n = 5, 7, 9$ and 11) with Au(111). We believe that the $\text{AuC}\equiv\text{CR}$ group is particularly interesting as the basis for SAMs on Au for two reasons: i) the acetylene group connects with gold atoms on the surface by a strong σ -bond;¹⁴ ii) the orbital structure of the acetylene group, and the existence of a variety of stable organometallic compounds containing the $\text{AuC}\equiv\text{CR}$ group, suggests that the interface between the metal, Au, and the saturated organic component of the SAM ($\text{R} = (\text{CH}_2)_n\text{CH}_3$), might be especially informative in studies in which the interface connecting the SAM to the metallic substrate might contribute to its properties.

The objective of this work was to address the question of the order of *n*-alkyne-derived SAMs on gold through a combination of microscopic, spectroscopic and contact angle measurements. Our results show that the disorder and mixed organic functionality implied by previous work^{4–6} are artefacts reflecting oxidation of the terminal acetylene by oxygen in solution during formation of the SAM (perhaps in a reaction catalyzed by gold),¹⁵ and that using appropriate experimental conditions (e.g., rigorous exclusion of oxygen; a slightly elevated temperature of 60°C during formation of the SAM) results in well-organized SAMs of alkynes that have qualities similar to that of alkanethiols on Au(111). These studies thus establish that SAMs of the surface composition $\text{AuC}\equiv\text{CR}$ provide a new type of SAM for use in physical- and physical-organic studies having an interface to gold ($\text{AuC}\equiv\text{CR}$), and are complementary to the well-understood SAMs of alkanethiols (AuSR).

SAMs of alkynes were prepared on gold by submerging freshly evaporated Au(111) substrates in a 1mM ethanolic solution of *n*-alkyne ($\text{HC}\equiv\text{C}(\text{CH}_2)_n\text{CH}_3$, $n = 5, 7, 9$, and 11) for 15 hours at 60°C . Importantly, the preparation of the SAMs was performed in

Figure 1. In (a-c) STM data for decyne/Au(111) SAMs. Panels assigned as A, B and C show cross-sections A, B and C indicated in (a-c), respectively. The inset in (b) shows FFT spectrum of image (b) with indicated six-fold symmetry pattern. Yellow rectangle in (b) marks an area which corresponds to the image shown in (c) obtained after FFT filtering of (b). In (d) scheme of decyne adsorption in the $(\sqrt{3} \times \sqrt{3})R30^\circ$ structure with arbitrary taken adsorption seats.

an oxygen-free environment to avoid oxidation of the acetylene group. The supporting information (SI) provides a more detailed description of the experimental procedure, as well as additional details of measurements.

Figure 1 summarizes the results obtained by scanning tunneling microscopy (STM) for decyne ($n = 7$) chemisorbed on Au(111). Analysis on a larger scale (Figure 1a) shows formation of depressions on the substrate with depths compatible with substrate lattice steps due to a single atomic layer of gold (as indicated by the respective cross-section, marked by A). Such depressions are also characteristic of formation of both thiol-¹ and selenol-¹⁶ based SAMs on Au(111), and probably result mainly from lifting the Au(111) herringbone reconstruction upon chemisorption of these

Figure 2. Overview of IRRAS data for $n = 5, 7, 9$ and 11 alkynes on Au(111) together with the corresponding DDT/Au(111) in the characteristic C-H stretching range.

molecules.¹⁷ The detection of similar features for alkynes suggests the formation of densely packed chemisorbed structures on Au(111).

This inference of a dense monolayer is confirmed by high-resolution STM, accompanied by Fourier analysis (using Fast Fourier Transform, FFT), which shows a hexagonal lattice. The data obtained after FFT filtering (Figure 1c) suggest a hexagonal structure with a period of ca. $5 \text{ \AA}$, as indicated by the line scans along the axes. The structure inferred from this analysis (shown schematically in Figure 1d) is consistent with the $(\sqrt{3} \times \sqrt{3})R30^\circ$ lattice (which characterizes the structure formed by alkanethiols on Au(111)¹⁸ and leads to an area per molecule of $21.5 \text{ \AA}^2$. While the quality of our data does not permit a more detailed description of the structure (as was possible for alkanethiols, which exhibit a $c(4 \times 2)$ superlattice),¹⁸ it certainly shows that alkynes on Au(111) form well-ordered structures with packing densities similar to those of alkanethiols (area per molecule $21.5 \text{ \AA}^2$).

To characterize the orientational order of alkyne-based SAMs further, we used infra-red reflection absorption spectroscopy (IRRAS). IRRAS measurements of the SAMs—formed by reaction of octyne ($n = 5$), decyne ($n = 7$), dodecyne ($n = 9$) and tetradecyne ($n = 11$) with Au(111)—showed vibrational bands in the C-H stretching range at 2965 cm^{-1} ($\nu_a\text{CH}_3$), 2938 cm^{-1} ($\nu_s\text{CH}_3$, FR), 2920 cm^{-1} ($\nu_a\text{CH}_2$), 2878 cm^{-1} ($\nu_s\text{CH}_3$, FR), and 2851 cm^{-1} ($\nu_s\text{CH}_2$) (Figure 2). For comparison, analogous IRRAS measurements were performed on SAMs of dodecanethiol (DDT) on Au(111). This comparison shows that the frequencies, relative intensities and the band width of the spectra for $\text{AuC}\equiv\text{CR}$ and AuSR are similar (for more details see Figure 1S in the SI).¹⁹ For the same orientation of n -alkynyl and alkanethiol on Au(111), however, we would expect the DDT ($n_{\text{CH}_2} = 11$) spectrum to be close to that of tetradecyne ($n_{\text{CH}_2} = 11$), which has a comparable aliphatic chain length $((\text{CH}_2)_n\text{CH}_3)$. Instead, the DDT spectrum intensity is similar to that of octyne ($n_{\text{CH}_2} = 5$). As a result of

Figure 3. IRRAS data analysis for tetradecyne/Au(111) with fitting individual bands. The inset shows linear increase in 2920 cm^{-1} peak intensity as a function of the CH_2 group number.

surface selection rules, IRRAS intensities are sensitive to the orientation of the molecules within the SAM (with respect to the metal substrate); the data in Figure 2 suggest slightly different twisting and/or tilting of the $(\text{CH}_2)_n\text{CH}_3$ chain for the n -alkynyl and alkanethiolate on Au(111). These differences are not surprising considering the different bonding geometries for each SAM. We also note that the observed differences in the IRRAS spectra for n -alkynyl and alkanethiol on Au(111) are much smaller than those reported previously¹⁹ for alkanethiolates on Au(111) and Ag(111), which have similar surface structures.

Figure 3 shows a more detailed analysis of the IRRAS spectra for tetradecyne, including a fitting for the C–H stretching modes. The inset in Figure 3 shows that an increase in the length of the aliphatic chain (from $n = 5$ up to $n = 11$) correlates linearly with the intensity of the CH_2 -related symmetric (2920 cm^{-1}) mode. This observation indicates that the conformation of n -alkynes on

Figure 4. XPS C1s and O1s data for $n = 5, 7, 9$ and 11 alkynes on Au(111) together with the corresponding DDT/Au(111).

Figure 5. Film thickness calculated in number of methylene groups $((\text{CH}_2)_n)$ from the XPS data (see text) for SAMs of $\text{HC}\equiv\text{C}(\text{CH}_2)_n\text{CH}_3$, ($n = 5, 7, 9$, and 11) on Au(111). The solid line indicates a linear fit.

Au(111) is preserved across the entire series of alkynes that we investigated; Nuzzo *et al.* reported a similar observation in earlier spectroscopic studies of alkanethiols on Au.¹⁹ In contrast to our observations, recent PM-IRRAS experiments by Scholz *et al.*⁶ (based on analysis of the C–H stretching range) performed for alkyl-based SAMs— n -butylmercury tosylate ($\text{C}_4\text{H}_9\text{HgOTs}$) and n -octadecylmercury tosylate ($\text{C}_{18}\text{H}_{37}\text{HgOTs}$) on gold—demonstrated fundamental differences in the relative intensities and band broadening between these SAMs and analogous alkanethiols: that is, the n -alkyls had a liquid-like structure (in contrast to the crystalline structure observed for n -alkanethiols). In contrast, the IRRAS data presented in Figures 2 and 3 demonstrates that it is possible to form alkynes that have order similar to that of alkanethiols on Au(111).

Figure 4 shows X-ray photoelectron spectroscopy (XPS) analysis of the alkyne-based SAMs. In contrast to previously reported XPS data for increasing lengths of alkynes ($\text{HC}\equiv\text{C}(\text{CH}_2)_n\text{CH}_3$, $n = 5, 7, 9$, and 11),⁴ the C1s signal in Figure 4 shows a single symmetric peak at 285 eV, with no additional higher energy components (e.g., peaks at 287 eV, 289 eV). The C1s spectra are also consistent with the corresponding data obtained for DDT (Figure 4), as well as other literature data²⁰ for un-oxidized alkanethiols on Au(111). We calculated the film thickness using the C1s/Au4f intensity ratios (assuming an exponential attenuation of the photoelectron signal²¹ and using attenuation lengths reported earlier).²² The calculated values (summarized in Figure 5) show a linear increase in thickness as a result of an increase in the length of the chain (from $n = 5$ to $n = 11$). Importantly, the XPS data are consistent with the IRRAS data shown in Figure 3; they indicate a linear increase in the IR signal with an increasing number of methylene units (CH_2). Moreover, using the linear relation obtained from our XPS measurements data, we can extrapolate the thickness of the film for $n = 0$ to a value of 5.9 Å ; this value corresponds to the length expected for a $-\text{C}\equiv\text{CCH}_3$ fragment. This extrapolation is also consistent with the ca. 5.6 Å dimensions of an upright configuration of the $-\text{C}\equiv\text{CCH}_3$ molecule on Au(111), on the basis of the bonds lengths provided by previous DFT²³ calculations. Thus, the estimation of thickness using XPS is consistent with SAMs of $\text{HC}\equiv\text{C}(\text{CH}_2)_n\text{CH}_3$ on Au(111) having the $-\text{C}\equiv\text{C}-$ group perpendicular to the surface.¹⁴

Previously reported XPS data for SAMs derived from ethynylbenzene ($\text{HC}\equiv\text{CC}_6\text{H}_5$) on gold⁵ suggested that the SAM contained oxidized components, as is inferred from both a significant O1s signal and higher-energy contributors to the C1s peak (e.g., peaks at 287 eV, 289 eV). In contrast, data shown in Figure 4 show virtually no O1s signal for all SAMs analyzed, which were prepared carefully in an oxygen-free environment (the SI details the procedure). The oxidation of alkynes is indeed clearly visible for samples that were exposed to oxygen (ambient condi-

Figure 6. XPS region scans (C1s and O1s) for SAMs of $\text{HC}\equiv\text{C}(\text{CH}_2)_{11}\text{CH}_3$ on Au(111) prepared in an oxygen-free atmosphere (upper spectra) and an oxygen-containing atmosphere (lower spectra).

tions) during preparation of the SAMs (see Figure 6). In the XPS data (Figure 6), an intense O1s signal in this case is associated with significant changes in the C1s peak, and the appearance of higher energy components (289 eV, 287 eV) which cause significant asymmetry and broadening of the C1s peak, and are indicative of the oxidation of the $-\text{C}\equiv\text{C}-$ group as reported in ref. 5. Our data demonstrate that the oxidation of alkynes does not involve the $\text{C}-\text{Au}(111)$ bond formation (as had been suggested previously⁵), but instead involves oxidation of the alkynes in solution during formation of the SAM.

The interfacial free energy of n -alkyl SAMs on Au(111) was investigated using measurements of advancing contact angles of water ($\Theta_{\text{H}_2\text{O}}^a$). The data (see Table 1 in the SI file) are in a range ($\Theta_{\text{H}_2\text{O}}^a = 111^\circ$ – 115°) that is consistent with the values measured here for DDT/Au(111), as well as literature data²⁴ for SAMs of alkanethiols on Au(111). The values of $\Theta_{\text{H}_2\text{O}}^a$ presented here are significantly higher than those reported previously (89° – 98°) for alkynes on Au(111),⁴ and serve as additional evidence of a similar degree of order for SAMs of alkynes (prepared properly) and alkanethiols on Au(111).

In conclusion, our STM and IRRAS data show that alkynes on Au(111) form SAMs that have organization and structure similar to alkanethiols on Au(111). High surface ordering and packing density of alkynes is also consistent with high values of $\Theta_{\text{H}_2\text{O}}^a$. The XPS data show that the presence of oxygen marks contamination of alkyne-based SAMs through oxidation of the acetylene group. The oxidation of $-\text{C}\equiv\text{C}-$ (most likely catalyzed by the gold substrate in the presence of O_2) can be monitored by the appearance of high energy components and a broadening and asymmetry in the C1s signal. Alkynes appear to be more sensitive to oxidation than alkanethiols during formation of the SAM, and require the formation to be carried out carefully in an oxygen-free atmosphere.

ASSOCIATED CONTENT

Supporting Information

Experimental details are available free of charge via the Internet at <http://pubs.acs.org>

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Notes

The authors declare no competing financial interests.

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