Grafting of Benzylic Amide Macrocycles onto Acid-Terminated Self-Assembled Monolayers Studied by XPS, RAIRS, and Contact Angle Measurements

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The grafting of benzylic amide macrocycles, the basic units of more complex mechanically interlocked architectures such as catenanes and rotaxanes, was performed via the functionalization of an acid-terminated self-assembled monolayer (SAM) of 11-mercaptoundecanoic acid (11-MUA). Both chemical and physical adsorption were investigated using macrocycles containing either a reactive hydroxyl functionality or an exo-pyridyl moiety, with characterization of the systems using X-ray photoelectron spectroscopy (XPS), contact angle measurements, and reflection absorption infrared spectroscopy (RAIRS). By comparing theoretical values with experimental data, it was found that up to 40% of the acid groups on the SAM surface were modified by the macrocycles and that the macrocycle coverage can be controlled by varying the reaction time. Stability tests demonstrated that the prepared films are stable in air, under ultrasound treatment and against chemical substitution. RAIRS measurements suggested a tilted orientation of the macrocycle with respect to the plane of the SAM surface.

1. Introduction

Considerable interest has been focused on the preparation and characterization of functionalized, switchable, ordered arrays of mechanically interlocked molecules for the purpose of developing new and promising multifunctional materials. Catenanes and rotaxanes, two classes of interlocked molecules, have shown great potential as components for nanoscale devices in the form of molecular shuttles, switches, and information storage systems. The key property of catenanes (two or more mechanically interlocked macrocycles) and rotaxanes (one or more macrocycles locked onto a linear “thread” by two bulky terminal stoppers) is that the mechanical bond holding the components of the molecule together allows large-amplitude relative movements of the components and can be used to modify the molecular properties by external stimuli. Ideally, the incorporation of these molecules into workable devices involves the formation of ordered bidimensional arrays while retaining their dynamic properties.

Functionalization of alkanethiol SAMs is a method often used to anchor macromolecular units such as proteins or enzymes onto a solid surface (see, for example, refs 13−24). The success of this approach is due to the simplicity of SAM film preparation, its reproducibility, and the possibility to create a wide range of surfaces via the incorporation of different groups at the end of the alkyl chains at the gas−solid interface. For example, it is possible to functionalize SAMs via covalent bonding reactions with experimental data, it was found that up to 40% of the acid groups on the SAM surface were modified by the macrocycles and that the macrocycle coverage can be controlled by varying the reaction time. Stability tests demonstrated that the prepared films are stable in air, under ultrasound treatment and against chemical substitution. RAIRS measurements suggested a tilted orientation of the macrocycle with respect to the plane of the SAM surface.

2. Experimental Section

Materials. Macrocycles 1 and 2 (Figure 1a and b) were synthesized using methods analogous to those previously...
described in the literature. To graft macrocycle 1, the carboxylic acid-terminated SAMs were immersed in 1 mM solutions (using dichloromethane as the solvent) of EDCI and 1 for 7 to 118 h. The grafting of macrocycle 2 took place over the same immersion times using a 1 mM dichloromethane solution of 2. Films of 11-MUA derivatized with EDCI alone were prepared by immersion in a 1 mM aqueous solution for 51 h. The modified surfaces were each rinsed and sonicated for 30 s in the pure solvent (dichloromethane or water) and dried under a stream of argon prior to analysis by XPS, RAIRS and contact angle measurements.

**Stability Tests.** The stability of the films was investigated upon exposure to ultrasound treatment for 30 s to 20 min. Their stability against aging under atmospheric conditions was verified by exposing the films to air for 3–20 days. To avoid light-induced aging effects, the films were kept in the dark during these stability tests. The SAM film functionalized with macrocycle 2 was also checked against chemical substitution by exposing it to molecules that can also physisorb onto the surface and therefore potentially displace the macrocycle. The films were exposed to either a 15 mM aqueous solution of ammonium chloride, a 1 mM methanol solution of aniline, or a 10 mM methanol solution of aniline for different immersion times and then rinsed in the pure solvent before analysis. Before and after each treatment, the surfaces were characterized by XPS and contact angle measurements.

**X-ray Photoelectron Spectroscopy (XPS) Analysis.** High-resolution XPS measurements were performed using an SSX-100 (Surface Science Instruments) photoelectron spectrometer with a monochromatic Al Kα X-ray source ($h\nu = 1486.6$ eV). The energy resolution was set to 0.92 eV, and the photoelectron takeoff angle (TOA) was 90°. All binding energies were referenced to the Au 4f\(^{7}/2\) core level. The base pressure in the spectrometer was in the low 10\(^{-10}\) Torr range.

Spectral analysis included a linear background subtraction and peak separation using mixed Gaussian–Lorentzian functions in a least-squares curve-fitting program (Winspec) developed in our laboratory. The photoemission peak areas of each element, used to estimate the amount of each species on the surface, have been normalized by the sensitivity factors of each element tabulated for the spectrometer used.

Three different points of each sample were analyzed to check for homogeneity. Within the error bars quoted, we found the same atomic proportions for all points of the same sample and therefore conclude that all samples can be considered homogeneous.

**Contact Angle Measurements.** Static contact angles of deionized (milli-Q) water deposited on the samples were measured in air using a video contact angle system VCA-2500XE (AST Products). All quoted angles are subject to an error of ±3°.

**Reflection Absorption Infrared Spectroscopy (RAIRS).** The RAIRS experiments were performed at room temperature in the sample compartment of a Biorad FTS-60A FT-IR spectrometer. P-polarized infrared radiation was reflected from the sample compartment of a Biorad FTS-60A FT-IR spectrometer with a monochromatic Al Kα X-ray source ($h\nu = 1486.6$ eV).
spectra are displayed as the ratio of the adsorbed macrocycle single beam to the single beam of 11-MUA on gold. In addition to the RAIRS measurements, infrared transmission spectra of solid-state macrocycle 1 were recorded using 150-mg pellets, 1 wt % of the macrocycle in KBr. Data collection conditions were similar to those used for the RAIRS experiments but with the spectrometer in transmission mode.

3. Results and Discussion

XPS Analysis. a. SAM Functionalization with Macrocyle 1. Characterization of the Modified Surfaces. Figure 3 shows the carbon 1s core-level photoemission spectra for a SAM of 11-MUA (bottom panel) and for a SAM functionalized with macrocycle 1 (top panel). Because no chlorine could be detected in the photoemission spectra of these samples (not shown), we can be sure that no solvent molecules (CH$_2$Cl$_2$) are incorporated in the photoemission spectra of these samples (not shown), we can be sure that no solvent molecules (CH$_2$Cl$_2$) are incorporated in the photoemission spectra of these samples. The most convincing evidence that macrocycle 1 is present on the surface comes from the analysis of the photoemission spectra of the nitrogen 1s core-level region. In the spectra recorded for the monolayer functionalized by the macrocycle, shown in Figure 4 (top panel), the single peak at 400.1 eV binding energy with a full width at half-maximum (fwhm) of 1.4 eV corresponds to the amide nitrogen. This agrees with the fact that no CI$^-$ remains in solution. Even in the N 1s spectra recorded for samples corresponding to shorter immersion times (not shown) we find exactly the same line shape as that shown in Figure 4 (top panel), whereas if we expose the EDCI film to a solution containing only the macrocycle, we find afterward an N 1s spectrum (not...
shown) very similar to the EDCI N signal, which can be reproduced by adding to the spectrum of Figure 4 (bottom panel) a tiny amount of the spectrum Figure 4 (top panel). This indicates that when the entire surface is covered by EDCI very few active sites on the SAM are accessible to the macrocycle.

When instead EDCI and the macrocycle are present in the solution to which the SAM is exposed, the activation of a site by the EDCI is followed by its functionalization until all of the available sites are functionalized, and this explains why no EDCI signal is present in the N 1 s spectrum of film functionalized with 1 (Figure 4, top panel).

Figure 5 presents the S 2p photoemission lines of the SAM before (bottom panel) and after (top panel) functionalization with the macrocycle. The latter signal is weaker (hence showing more noise), as expected for a thicker film, indirectly confirming the macrocycle’s presence. However, it is still possible to identify, as for the SAM alone (Figure 5, bottom panel), two components. For the first, the S 2p\(\frac{3}{2}\) peak is situated at 161.9 eV, and we assign it to sulfur bonded to gold. The other component with the S 2p\(\frac{1}{2}\) peak at 163.5 eV could be due to a small number of alkanethiols not covalently bonded to the substrate but only intercalated between the 11-MUA molecules bound to Au or physisorbed as a double layer. Alternatively, the second component might be derived from disulfides formed under the influence of X-rays during spectra acquisition. However, the aging behavior of the films supports the disulfide attribution. (See Stability of the Films below.)

*Estimation of the Functionalization Yield.* Figure 6a shows a plot of the ratios between the N 1 s and S 2p (top panel), N 1 s and Au 4f (center panel), and S 2p and Au 4f (bottom panel) photoemission peak areas as a function of the reaction time. As expected, the S 2p to Au 4f ratio remains reasonably constant for all immersion times, indicating that the monolayer of 11-MUA is stable under these conditions. The N 1 s to Au 4f and N 1 s to S 2p ratios increase with the immersion time, indicating macrocycle adsorption. We also compared the experimentally determined atomic percentages, obtained from the photoemission peak areas, with those calculated for the functionalization of 12, 30, and 50% of the acid groups of the SAM. In the calculation, we considered a model surface of 100 thiol chains and computed the atomic percentages for N, C, O, and S (excluding hydrogen, which cannot be detected by XPS) for a coverage of 12, 30, or 50 macrocycle molecules. The results are summarized in Table 1: after 15 h of reaction, 12% of the surface acid groups are functionalized by 1, while after 118 h a yield of 30% is reached. Comparing the size of an acid group to the dimensions of a macrocycle (average diameter 10 Å) and considering that the orientation of the macrocycle with respect to the surface may be random, we can conclude that nearly all of the self-assembled monolayer area is covered by macrocycle 1.

The error in the photoemission peak areas was estimated depending on the signal-to-noise ratio in the spectrum for each element: the carbon and oxygen signals are better defined because the errors were found to be 2% for C, 5% for O, 10% for S, and 15% for N. The error in the atomic percentages was estimated to be 2% for C, 5% for O, 10% for S, and 15% for N. The error in the photoemission peak areas was estimated to be 2% for C, 5% for O, 10% for S, and 15% for N.
Grafting Amide Macrocycles onto Monolayers


TABLE 2: Comparison between Experimental Atomic Percentages Derived from the Photoemission Peak Areas of an 11-MUA Film Functionalized with Macrocycle 2 Prepared with Different Increasing Immersion Times and Theoretical Values Calculated for Different Coveragesa

<table>
<thead>
<tr>
<th>Immersion Time</th>
<th>Experimental Atomic Percentages</th>
<th>Theoretical Atomic Percentages</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>%C exp</td>
<td>%O exp</td>
</tr>
<tr>
<td>7 h</td>
<td>76.7 ± 1.5</td>
<td>77.4 ± 1.5</td>
</tr>
<tr>
<td>24 h</td>
<td>17.1 ± 0.9</td>
<td>15.9 ± 0.8</td>
</tr>
<tr>
<td>103 h</td>
<td>5.2 ± 0.5</td>
<td>3.6 ± 0.4</td>
</tr>
<tr>
<td>10%</td>
<td>1.0 ± 0.2</td>
<td>3.2 ± 0.5</td>
</tr>
<tr>
<td>30%</td>
<td>0.2 ± 0.0</td>
<td>0.9 ± 0.2</td>
</tr>
<tr>
<td>40%</td>
<td>0.0 ± 0.0</td>
<td>0.0 ± 0.0</td>
</tr>
</tbody>
</table>

a The error in the experimental atomic percentages was estimated to be 2% for C, 5% for O, 10% for S, and 15% for N.

b. SAM Functionalization with Macrocycle 2. Characterization of the Modified Surfaces. As for macrocycle 1, the changes induced by the functionalization of the 11-MUA SAM with macrocycle 2 was followed by XPS. Figure 7 shows the N 1s and C 1s core-level regions in detail as well as their components deduced from the spectral analysis. First, the appearance of a photoemission signal in the N 1s core-level region (Figure 7, top panel) confirms the presence of the macrocycle on the surface of the acid-terminated SAM. The large peak at 400.2 eV (fwhm = 1.6 eV) is due to amide nitrogen atoms and to the pyridine nitrogen in the macrocycle. The C 1s core-level spectrum for the functionalized monolayer (Figure 7, bottom panel) shows a peak at 284.6 eV due to aromatic carbons and one at 285.4 eV corresponding to the aliphatic chain and to aromatic carbon atoms bound to electronegative groups. In addition, the shoulder at 286.4 eV comprises the C 1s signal of aliphatic carbon atoms bound to carboxylic or amide groups and to pyridine nitrogen. At 287.3 eV, we find the signature of amide carbon, at 288.7 eV, that of ester carbon, and at 289.6 eV, that of carboxylate carbon. The last feature at 291.8 eV is the shake-up associated with the aromatic rings.

In considering the interaction between the SAM and macrocycle 2, pyridine N 1s photoemission is expected between 399.0 and 399.5 eV depending on the chemical environment of the pyridine rings. However, theoretical calculations on rotaxanes investigating the effects of intra- and intermolecular interactions on the binding energy showed that hydrogen bonding interactions can produce shifts to higher binding energy of up to 1 eV. An electrostatic interaction between a protonated pyridine nitrogen and an anionic carboxylate function following the exchange of the proton of the carboxylic acid group should give rise to a new component in the N 1s core-level region between 399.0 and 402.0 eV, which is not seen. We can therefore rationalize the fact that we do not distinguish two separate components for amide and pyridine nitrogen in the N 1s core level (Figure 7, top panel) by assuming that the interaction between the pyridine moiety of the macrocycle and the acid group of the SAM does not occur through proton exchange but through hydrogen bonding, at least for the large majority of molecules.

Estimation of the Functionalization Yield. Similar to the procedure detailed above for 1, we followed the evolution of the functionalized surface as a function of the immersion time by XPS. Figure 6b shows the ratios between the photoemission peak areas of N 1s and S 2p (top panel), N 1s and Au 4f (center panel), and S 2p and Au 4f (bottom panel) versus time. The S 2p to Au 4f ratio remains constant for all immersion times, confirming the stability of the SAMs, as observed for 1. The N 1s to Au 4f and N 1s to S 2p ratios increase with the immersion time from 7 to 103 h, indicating a gradual increase in the functionalized surface area. Moreover, a comparison of the experimental atomic percentages obtained from the peak area with the theoretical values calculated as explained previously for 1 and shown in Table 2 indicates that after 7 h of immersion 10% of the acid groups of the SAM are functionalized by 2 and after 103 h of immersion the functionalization yield reaches 40%. As for 1, we note that on the basis of the dimensions of both the acid groups and 2 this corresponds to a value close to the maximum possible surface coverage.

We note that the functionalization yields for 1 and 2, determined as detailed above, rely on the implicit assumption that the SAM is compact and no free gold surface is available for direct adsorption. Deducing reliable information on the compactness and the thickness of both the SAM of 11-MUA and the SAM functionalized with either 1 or 2 from the XPS data is extremely difficult because precise values for the electron escape depth are not available. We therefore decided to approach the problem through a completely independent series of electrochemical experiments. The results of this work are too copious to be included here, and we are currently preparing a separate manuscript detailing them. However, we found evidence of a highly ordered and compact SAM (no holes or defects).

c. Stability of the Films. More detailed information on the type of bonding interactions between the macrocycles and the 11-MUA SAM would be of great interest and is contained within the C 1s photoemission signal. However, because of the complexity of this core-level region consisting of multiple components, it was not possible to prove the exact nature of the interaction between the macrocycle and the acid-terminated SAM surface by XPS. Nevertheless, photoemission can be used to investigate the stability of the macrocycle-functionalized SAMs under different external conditions such as ultrasound treatment and aging in air. Contact angle measurements were performed to distinguish the different surfaces by their polarity, and XPS allowed us to monitor possible chemical changes.

The contact angle (CA) obtained for the 11-MUA SAM immediately after deposition of the drop is 29°; however, during...
the following minute, the drop wets the surface, and the contact angle drops below 10°, as expected for such a highly hydrophilic surface. The reaction of EDCI with the acid groups of the SAM according to the scheme in Figure 2 increases the CA to 76°, indicating that a more hydrophobic surface has been produced. After reaction of the SAM activated by EDCI with I, a CA of 55° confirms the presence of the macrocycle, whose amide and ester groups are responsible for the more hydrophilic character of the surface. Because I and 2 are very similar, it is not surprising that we find a CA of 57° after functionalization of the SAM with the latter. Macrocyle films subjected to ultrasound treatment showed the same contact angle values even after 20 min of treatment. Figure 8a shows the N 1s to S 2p, N 1s to Au 4f, and S 2p to Au 4f ratios calculated from the XPS peak areas for the same films of I and 2 after sonication; these ratios are constant for all sonication times in agreement with the contact angle data, confirming that macrocycle-functionalized SAMs are stable with respect to ultrasound treatment.

To monitor changes due to aging in the dark, XPS spectra of films of SAMs functionalized with I and 2 were collected after exposure to air for 3, 6, and 20 days. The atomic percentages calculated from these spectra for I are shown in Table 3; an increase in the amount of oxygen on the surface is clearly seen after only 3 days of exposure. After 6 days, the oxygen contamination is still the same, but after 20 days, there is still more oxygen present.

Figure 9 presents the S 2p core-level region measured for the SAM surface. Ammonium chloride (NH₄⁺Cl⁻) and aniline molecules are still intact on the SAM surface. Films of 11-MUA and macrocycle 1 freshly prepared and after Aging in Air  

<table>
<thead>
<tr>
<th></th>
<th>freshly prepared</th>
<th>3 days</th>
<th>6 days</th>
<th>20 days</th>
</tr>
</thead>
<tbody>
<tr>
<td>%C</td>
<td>80.5 ± 1.6</td>
<td>74.9 ± 1.5</td>
<td>75.4 ± 1.5</td>
<td>74.7 ± 1.5</td>
</tr>
<tr>
<td>%O</td>
<td>13.1 ± 0.7</td>
<td>18.2 ± 0.9</td>
<td>17.9 ± 0.9</td>
<td>19.4 ± 1.0</td>
</tr>
<tr>
<td>%S</td>
<td>3.3 ± 0.3</td>
<td>3.8 ± 0.4</td>
<td>3.4 ± 0.3</td>
<td>2.9 ± 0.3</td>
</tr>
<tr>
<td>%N</td>
<td>3.1 ± 0.5</td>
<td>3.2 ± 0.5</td>
<td>3.0 ± 0.5</td>
<td>3.0 ± 0.5</td>
</tr>
<tr>
<td>N/S</td>
<td>0.9 ± 0.2</td>
<td>0.8 ± 0.1</td>
<td>0.8 ± 0.1</td>
<td>1.0 ± 0.2</td>
</tr>
</tbody>
</table>

The error in the experimental atomic percentages was estimated to be 2% for C, 5% for O, 10% for S, and 15% for N.

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Figure 9 presents the S 2p core-level region measured for these films of I. Whereas the 3- and 6-day-old films show spectral features identical to the freshly prepared film (Figure 5), the 20-day-old film has at least one new spectral component at 168.5 eV binding energy that can be attributed to a sulfonite species (SO₃⁻). We can therefore conclude that the additional oxygen in the films exposed for 3 or 6 days is due to atmospheric contamination physiosorbed onto the films and that after 20 days of aging the amount of oxygen still increases because of the oxidation of sulfur atoms. Nevertheless, the consistency in the N1s to S 2p, N1s to Au 4f, and S 2p to Au 4f ratios with time (Figure 8b) confirms that the macrocycle molecules are still intact on the SAM surface. Films of macrocycle 2 that were subjected to the same aging treatment showed a more rapid oxidation of sulfur than films of macrocycle 1: in Figure 10, we show the S 2p core-level spectra recorded for the same film at different aging times. We can observe a weak peak at 168.5 eV indicating sulfur oxidation after 6 days of exposure to air. After 20 days of aging, the same peak represents a significant proportion of the total sulfur content of the film. The difference in the oxidation rate of the two types of macrocycle films can be explained on the basis of the difference in interaction with the SAM for chemically bonded I and physically adsorbed 2: a different orientation of the macrocycles in the two cases or different homogeneities of these surfaces could allow for different channels for the oxygen penetration into the monolayer chains to reach the sulfur atoms. It is important to note that the stability of our functionalized surfaces against oxidation is in line with what is reported in the literature for CH₃-terminated SAMs with the same chain length as that of the alkanethiols used here. Moreover, we observe that it is the component corresponding to the sulfur bonded to gold that decreases as the sulfonite species increases. The component with the S 2p{3/2} peak at 163.5 eV remains unchanged, and this supports the attribution of the latter to disulfides. In fact, if noncovalently bound alkanethiols were present on the surface, then one would expect them to oxidize easier than the sulfur bonded to gold.

The macrocycle 2 films were subjected to other types of stability tests to examine whether it was possible to substitute the macrocycle at the interaction site with the SAM with molecules containing functionality that can also interact with the SAM surface.
The substitution tests indicate that the bond of macrocycle 2 with the SAM is stable and indirectly confirm our result for the functionalization yield in that it is only when the SAM surface is completely covered by 2 that the access of aniline or ammonium chloride molecules to the remaining free-acid terminal groups is sterically hindered.

d. RAIRS Measurements on Macrocycle 1. Because the grafting of macrocycle 1 seems to yield the most stable films with respect to aging in air, we decided to extend the analysis of this system using RAIRS to derive information on the orientation of the macrocycles with respect to the SAM surface plane.

The RAIRS spectrum obtained for the adsorbed macrocycle in the C–H stretching group frequency region is shown in Figure 11 (top panel). Although the absorption bands are inherently weak, it is possible to discern four features in the spectrum. The assignment of these bands is aided by comparison with the solid-state infrared transmission spectrum of 1 wt % macrocycle 1 in KBr (bottom panel).

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### Table 5: Assignments of Vibrational Modes for an 11-MUA Film Functionalized with Macrocycle 1 and the Solid-State Spectrum of 1 wt % Macrocycle 1 in KBr

<table>
<thead>
<tr>
<th>Mode</th>
<th>Solid State (cm$^{-1}$)</th>
<th>RAIRS (cm$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\nu$(NH)</td>
<td>3314</td>
<td>3314</td>
</tr>
<tr>
<td>$\nu$(OH)</td>
<td>3221</td>
<td>3221</td>
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<tr>
<td>$\nu$(CH)</td>
<td>3073</td>
<td>3073</td>
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<tr>
<td>$\nu$(CO)</td>
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<td>1756</td>
</tr>
<tr>
<td>$\nu_{as}$(CH$_3$)</td>
<td>2928</td>
<td>2928</td>
</tr>
<tr>
<td>$\nu_{as}$(CH$_3$)</td>
<td>2855</td>
<td>2855</td>
</tr>
<tr>
<td>$\nu$(CC)</td>
<td>1641</td>
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<tr>
<td>$\nu$(CC)</td>
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</tr>
<tr>
<td>$\nu$(CC)</td>
<td>1463</td>
<td>1463</td>
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</tbody>
</table>

*The error in the experimental atomic percentages was estimated to be 2% for C, 5% for O, 10% for S, and 15% for N.*
accompanied by a weaker CH$_2$ symmetric stretch at lower frequency,$^{43}$ and this band is observed in the spectrum at 2853 cm$^{-1}$.

The highest-frequency band in the C–H stretching region of the infrared spectrum occurs at 3018 cm$^{-1}$. The position of this band agrees with that expected for the C–H in-plane stretching mode of a phenyl ring,$^{44}$ and because macrocycle 1 contains phenyl groups, this is consistent with the presence of the macrocycle on the surface. The vibrational frequency for this mode is shifted down from the solid-phase frequency of 3041 cm$^{-1}$ possibly because of the phenyl rings experiencing an interaction with the surface resulting in weakening of the carbon–hydrogen bonds in the phenyl rings. The final band in the spectrum observed at approximately 2960 cm$^{-1}$ is an instrumental artifact due to instrumental instability and is not related to the macrocycle 1/11-MUA system. Unfortunately, it was not possible to determine if the amide nitrogen–hydrogen stretch (expected between 3270 and 3330 cm$^{-1}$ for the trans species$^{45}$) is present in the RAIRS spectra because a broad absorption band due to ice condensing on the detector window obscures the region above 3100 cm$^{-1}$.

Further evidence for the presence of the macrocycle on the surface is provided by the RAIRS spectrum in the lower-frequency region as shown in Figure 12 (top panel), which we again compare to the solid-state infrared transmission spectrum of macrocycle 1 (Figure 12, bottom panel). The spectrum is rather complicated and contains a considerable number of absorption bands, but the most intense bands can be discerned. In the carbonyl stretching region, two infrared absorption bands are observed that provide strong evidence for the presence of amide functional groups on the surface: the most dominant peak at 1662 cm$^{-1}$ corresponds to the amide I band (i.e., primarily a carbonyl stretch but also an in-phase bending of N–H and stretching of C–N of the amide functional group$^{46}$). This band is accompanied by the amide II band (a C–N stretch combined with an N–H bend), which is observed as a weak feature at 1533 cm$^{-1}$. The carbonyl band at 1762 cm$^{-1}$ is assigned to the carbonyl stretch of the ester functional group contained within the macrocycle.$^{47}$ Weak bands are observed at 1611 and 1511 cm$^{-1}$, and a very weak, broad feature is centered around 1452 cm$^{-1}$. The carbon–carbon stretching frequencies for 1,4-substituted phenyl rings normally occur in the regions of 1620–1585, 1590–1565 (weak), 1525–1480, and 1400–1420 cm$^{-1}$ $^{43}$ and close to 1612 and 1460 cm$^{-1}$ for 1,3,5-substituted phenyls.$^{47}$ Therefore, we deduce that the bands observed in the spectrum are due to both 1,4- and 1,3,5-substituted phenyl groups at the surface, as expected for the macrocycle grafted on the surface.

If the RAIRS spectra agree with the XPS data concerning the presence of I on the surface, the question still remains as to the nature of the interaction of the macrocycle with the self-assembled monolayer of 11-MUA. Because macrocycle 1 is expected to react with 11-MUA upon activation with EDCI to form a chemical attachment via an ester function, it should be possible to observe the ester in the RAIRS spectra. Clearly, we do observe an absorption band indicative of an ester group at 1762 cm$^{-1}$ in the RAIRS spectrum, but this is due not only to a possible ester group between the macrocycle and 11-MUA but also to the ester functions within the macrocycle itself because both ester functions are in an identical chemical environment. Consequently, it is impossible to differentiate them using vibrational spectroscopy. An alternate approach to determine whether covalent attachment occurs between the macrocycle and 11-MUA is via monitoring the intensity of the carbonyl stretch of the carboxylic acid. Reaction between the macrocycle hydroxyl group and the acid group of 11-MUA effectively reduces the number of free acid groups at the surface. Because the RAIRS spectra are the ratio of the single-beam spectrum of the adsorbed macrocycle to that of 11-MUA on gold, we would therefore expect to see an anti-absorption band at the vibrational excitational frequency of the acid group. The most intense absorption band of the acid function is the carbonyl stretching mode of 11-MUA, which occurs at 1718 cm$^{-1}$ for 11-MUA adsorbed on polycrystalline gold as shown in Figure 13, in agreement with ref 48. An examination of the spectrum reveals no evidence of an anti-absorption band in this region. However, this may be obscured by the absorption band due to the ester functional group occurring at 1762 cm$^{-1}$.

Additional interesting information concerning the adsorption geometry of the macrocycle can be obtained from vibrational spectroscopy. A comparison of the RAIRS spectra to the solid-state spectrum reveals that there is a marked difference in the relative intensity of the absorption bands. For a given poly-
crystalline solid sample, intensities of absorption bands depend only on the change in dipole moment associated with the vibrational mode. RAIRS absorption band intensity for adsorbates on a metal surface, however, is strongly influenced by the orientation and magnitude of the change in dipole moment of the vibration by consideration of the metal surface selection rule.\(^{49}\) Vibrational modes that undergo a change in dipole moment with a large component parallel to the underlying metal will have weaker absorption bands than those with a large component perpendicular to the metal. Whereas the intensities of the amide I band (1662 cm\(^{-1}\)) and the carbonyl stretch of the ester (1762 cm\(^{-1}\)) are similar in the solid-state spectrum (Figure 12, bottom panel), the intensity of the ester is considerably weaker than that of the amide absorption band in the RAIRS spectrum (Figure 12, top panel). Consequently, the carbonyl groups of the esters (those within the macrocyle and, if they exist, those formed between the macrocycle and the surface) must be oriented such that the change in dipole moment associated with the carbonyl stretch has a smaller component perpendicular to the plane of the underlying gold surface. Therefore, on average, the ester groups must be angularly aligned so that the carbonyl groups are closer to the plane of the gold than the amide carbonyl groups.

The intensity of the in-plane C–H stretching mode of the phenyl ring for the RAIRS spectrum in Figure 11 (top panel) was observed to diminish with respect to the CH\(_2\) stretching modes when compared to the solid-state spectrum (Figure 11, bottom panel). RAIRS studies of low coverages of benzene adsorbed on Cu(110) and Pt(111)\(^{50}\) demonstrate that benzene adsorbs with the plane of the ring parallel to the plane of the metal and that the C–H stretching mode is strongly infrared-inactive. An examination of Figure 11 reveals that for macrocycle 1 adsorbed on 11-MUA the in-plane stretch is not disallowed but is observed as an extremely weak band. Hence, we can conclude that all three phenyl rings of the macrocycle are not orientated parallel to the plane of the underlying gold surface but some or all must exhibit some amount of tilt with respect to the metal surface. The observation of the weak carbon–carbon stretching bands in Figure 12 supports this assertion. The \(\nu(C)\) modes in 1, 4- and 3, 5-substituted phenyl groups result in a change in dipole moment associated with the vibration that is in-plane. Therefore, there must be some tilt of the rings with respect to the surface for the vibrational modes to be infrared-active. We note that this geometry of 1 is different from what is found for a benzylic amide macrocycle directly adsorbed onto an Au surface.\(^{30}\)

4. Conclusions

We have established that macrocycles, the basic units of more complex mechanically interlocked architectures such as catenanes and rotaxanes, can be grafted onto acid-terminated SAMs by either chemical modification or physical adsorption, with the possibility to control the portion of functionalized surface area in both cases by varying the immersion time for the functionalization step. The two methods allow one to functionalize up to 30 or 40% of the acid groups on the SAM surface. This coverage appears quite important if we compare the dimensions of the acid groups with those of the macrocycles. If one wants to avoid intermolecular hydrogen bonding interactions that could hamper intramolecular motion, a functionalization of 40% represents the upper limit (i.e., the whole 11-MUA monolayer surface covered by 2) if one considers that the orientation of the macrocycles on the surface may well be random.

From the photoemission spectra, we obtained no clear evidence for the type of interaction between the macrocycles and the SAM because of the complexity of the spectral regions comprising details of carbon and oxygen atoms involved in the interaction. Therefore, we checked a posteriori the stability of the interaction of the prepared films under different chemical or physical conditions. The films of 1 prepared by chemical bonding were tested by ultrasound treatment and with respect to aging in air and were proven to be stable. The films of 2 prepared by physical adsorption were also subjected to ultrasound treatment and demonstrated high stability. They are, however, less resistant than 1 to aging in air because they show incipient sulfur oxidation after only 6 days. Physisorbed macrocycle 2 was also tested against chemical substitution and was found to be stable.

The RAIRS spectra also provided evidence for the grafting of macrocycle 1 intact onto the surface. Moreover, a consideration of the relative absorption band intensities revealed that the ester carbonyl groups are on average oriented closer to the surface plane than the amide carbonyl groups and that the phenyl rings must have some degree of tilt with respect to the surface.

The encouraging results obtained for these macrocycles provide the basis for the grafting of more complex catenanes and rotaxanes using a similar approach.

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References and Notes

(25) The SAM activation by the EDCI was performed both in water and in dichloromethane; for the same reaction times, the functionalization yield was more important in a water solution producing a stronger photoemission signal. For this reason, we choose to show the spectra recorded for a film prepared in water instead of the ones recorded for the film prepared in dichloromethane. However, the spectral analysis of the experimental data allowed the identification of the same components in both cases. We also note that because of the deprotonation of EDCI no high binding-energy peak characteristic of a protonated amine is found in either N 1s spectrum.
(32) We calculated the atomic percentages for many different coverages of the macrocycle but report here only the results for the functionalization of 12, 30, and 50% of the surface because these values correspond best to the experimental values.
(37) Bureau, Ch. Private communication.
(40) See, for example, Yan, L.; Marzolin, Ch.; Terfort, A.; Whitesides, G. M.; Langmuir 1997, 13, 6704.
(49) Only vibrations with a component of the dipole moment change normal to the surface may be observed.