

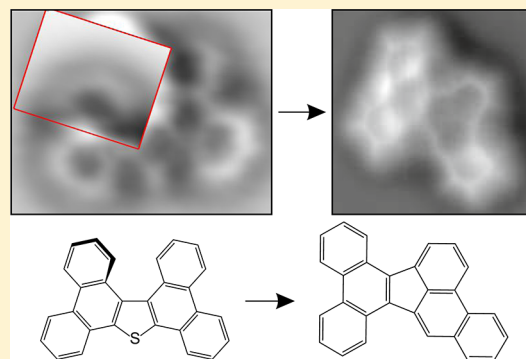
# Characterization of a Surface Reaction by Means of Atomic Force Microscopy

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**ABSTRACT:** We study a thermally activated on-surface planarization reaction by a detailed analysis of the reactant and reaction products from atomically resolved atomic force microscopy (AFM) images and spectroscopy. The three-dimensional (3D) structure of the reactant, a helical diphenanthrene derivative, requires going beyond constant-height imaging. The characterization in three dimensions is enabled by acquisition and analysis of the AFM signal in a 3D data set. This way, the structure and geometry of nonplanar molecules as well as their reaction products on terraces and at step edges can be determined.



## INTRODUCTION

By introducing CO-functionalized tips in frequency-modulated atomic force microscopy (FM-AFM), Gross and co-workers directly resolved the chemical structure of individual planar molecules<sup>1</sup> in real space. Inspired by this novel possibility, the technique has been applied to identify the structure of an unknown molecule,<sup>2</sup> to resolve the structure of reaction products in single molecule chemistry<sup>3,4</sup> as well as in thermally induced on-surface chemical reactions.<sup>5,6</sup> To avoid additional complexity in these demanding experiments, mostly planar molecules were studied, so that the structure determination could be achieved in the basic imaging mode, in which the tip is scanned in constant height over the sample. For select cases of nonplanar structures, the molecule was still flat enough to resolve its geometry.<sup>7–10</sup> To quantify slight deviations from a flat configuration, as, for example, an overall small tilt of the molecular plane with respect to the surface it is adsorbed on, FM-AFM spectroscopy was employed more recently.<sup>11</sup>

Most chemical reactions, however, involve three-dimensional (3D) molecular structures. In addition, the reaction pathway is critical for the reaction product. Although many catalytically activated reactions involve the presence of surfaces, on-surface chemistry and its reaction pathways are still poorly understood. In particular, the presence of a surface will strongly affect the possible reaction pathways, at least by restricting the molecular geometry.

Here, we study a thermally activated on-surface reaction by a detailed analysis of reactant and reaction products from atomically resolved FM-AFM images and 3D spectroscopy. To analyze the on-surface reaction in three dimensions, we extend the constant-height imaging used so far by analyzing the frequency shift along a nonplanar surface of a 3D data set. This way, the orientation and tilt angle of individual carbon rings in

the reactant can be determined. Subsequently, these molecules are submitted to an on-surface planarization reaction that is activated by annealing. The reaction products are also atomically resolved in real space from FM-AFM imaging. The comparison of the structures of reaction products and the 3D reactant suggests that the pathway of the on-surface reaction could involve an out-of-surface rotation of one phenanthrene moiety to enable a planarization of the molecule.

## EXPERIMENTAL SECTION

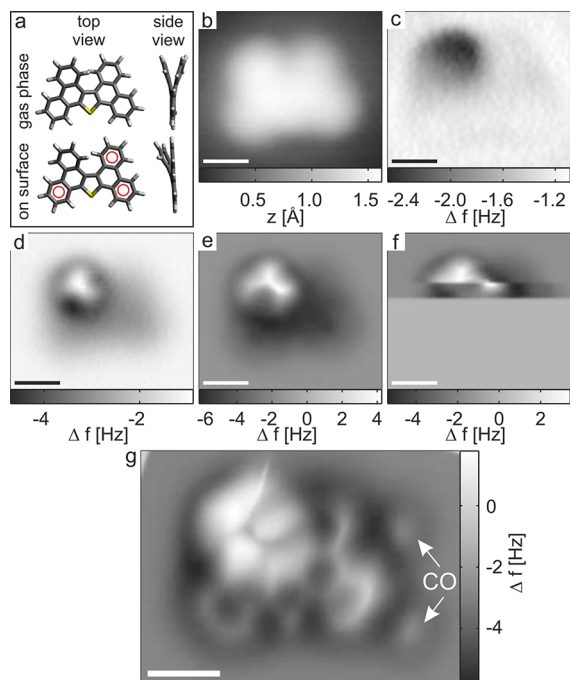
The experiments were carried out with a home-built combined scanning tunneling microscope/atomic force microscope, operating in ultrahigh vacuum at a temperature of 5 K. The atomic force microscope's qPlus tuning fork<sup>12</sup> was operated in the frequency modulation mode.<sup>13</sup> Sub-ångstrom oscillation amplitudes were used to maximize the lateral resolution.<sup>14</sup> Bias voltages refer to the sample with respect to the tip. The Cu(111) single-crystal sample was cleaned by several sputtering and annealing cycles. A small amount of CO was dosed onto the surface for tip functionalization.<sup>1</sup> NaCl islands were grown to facilitate the pick-up of CO molecules to the tip apex from these islands. All AFM data was obtained with a CO-terminated tip apex and away from the NaCl islands. Individual molecules were sublimed onto the sample being located in the microscope at a temperature of  $T \cong 5$  K. Density functional theory (DFT) calculations<sup>15</sup> of the free monomers were performed to obtain the molecular geometry. To confirm the structural assignment from AFM, matrix-assisted laser desorption and ionization–time-of-flight (MALDI-TOF) based mass spectrometry, nuclear-magnetic resonance (NMR) spectroscopy, and elemental analysis were performed.<sup>16</sup>

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## RESULTS AND DISCUSSION

Diphenanthro[9,10-*b*:9',10'-*d*]thiophene (DPAT) molecules were synthesized from a solid-state reaction under CO<sub>2</sub> atmosphere between phenanthrene and elemental sulfur.<sup>16,18</sup> The molecule in the gas phase is nonplanar because of steric hindrance. The two phenanthrene moieties are bent out-of-plane at the side opposing the sulfur atom, thus giving rise to a helical shape of the molecule. Its structure as calculated from DFT<sup>15</sup> is displayed in Figure 1a. Provided that DPAT remains



**Figure 1.** Characterization of DPAT molecules by means of scanning probe microscopy in standard modes of operation: (a) Stick models of the molecule's geometry as calculated for the free molecule after full relaxation (top) and with three benzene rings (marked in red) forced to one plane (bottom) to mimic the on-surface geometry, respectively. Carbon, hydrogen, and sulfur atoms are represented in black, white, and yellow, respectively.<sup>17</sup> (b, c) Simultaneously recorded topography (b) and frequency shift (c) acquired with active current feedback (constant-current mode,  $I = 2$  pA,  $V = 0.05$  V). (d–f) Set of  $\Delta f$ -images ( $\Delta z = 0.3, 0.5,$  and  $0.7$  Å with respect to the STM set point ( $I = 2$  pA,  $V = 0.1$  V)) of one DPAT molecule on clean copper. (g)  $\Delta f$ -image ( $\Delta z = 2.0$  Å with respect to the STM set point ( $I = 1$  pA,  $V = 0.1$  V)) of a DPAT molecule pinned to two CO molecules (scale bars 5 Å).

intact, adsorption onto the Cu(111) surface will break the symmetry between the two phenanthrene moieties since only one of them can directly face the surface. Interestingly, this asymmetry is not apparent in the topography of a constant current Scanning Tunneling Microscopy (STM) image, as depicted in Figure 1b. However, the simultaneously recorded frequency shift channel (Figure 1c) shows a clear asymmetry between the two phenanthrene moieties.<sup>19</sup> Figure 1d–g shows several constant-height AFM images of the molecule at decreasing tip–sample distances. Note that in Figure 1g the two CO molecules adsorbed next to the DPAT molecule stabilized the molecule against lateral displacement. Similar imaging attempts without CO molecules next to the DPAT molecule resulted in lateral displacement of the molecule in such a way that it could not be stably imaged (Figure 1f). At the upper left part of Figure 1d, the repulsive interaction sets in at

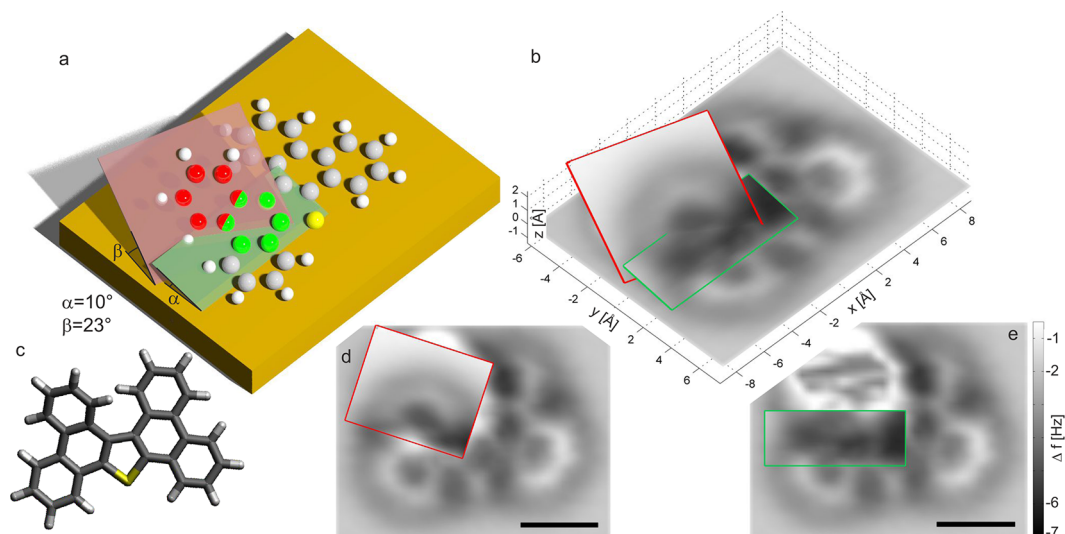
considerably larger tip height than over the rest of the molecule. Hence, this part of the molecule has to be pointing out toward the tip. Otherwise, the image acquired at closest tip–sample distance carries most of the information: four of the benzene rings are readily identified and seem to be oriented roughly parallel to the sample surface—all at a comparable adsorption height. This can be deduced from the intramolecular contrast being relatively homogeneous over these four benzene rings. This suggests that the phenanthrene moiety facing the surface (to the right in Figure 1g) is flattened out upon adsorption, whereas the other one bends even further due to steric hindrance. There is one atom exhibiting a bond angle slightly smaller than 90°, whereas all other atoms show approximately the 120° bond angles expected for sp<sup>2</sup>-hybridized carbon atoms. This suggests that the atom with the smaller bond angle is sulfur. Whereas one can guess from the image that the sulfur atom is part of a five-membered ring structure, the part of the molecule that sticks out of the surface plane cannot be identified, and it only leads to the strongly distorted part of the AFM image.

To also resolve the structure for this part of the molecule, we acquired a three-dimensional (3D) data set of frequency shift  $\Delta f$  as a function of all coordinates  $x, y$  and  $z$ , by recording  $\Delta f(z)$  versus distance  $z$  at a closely spaced grid in the surface plane  $x, y$ . To avoid too strong tip–sample interaction at close distances  $z$ , each of the  $\Delta f(z)$  spectra was aborted upon reaching a predefined  $\Delta f$  value as proposed and described by Mohn et al.<sup>20</sup> With this data set, we can extract and display the data for any arbitrary plane within the 3D data set. The analysis of such 3D imaging is shown in Figure 2. By a variation of the orientation of such planes and looking at the resulting images, one can unambiguously resolve the remaining structure of the molecule. Furthermore, as the orientations of the imaging planes are set, this technique allows us to extract the out-of-plane distortion angles of the molecule within to a few degrees uncertainty. Based on this analysis, we can identify the molecular structure itself as well as its 3D geometry after adsorption, despite the out-of-plane rotation of one of the benzene rings being as large as  $\approx 23^\circ$ . This possibility to resolve the 3D geometry of molecules is not at all self-evident, because at some even larger tilt angle one may expect that the CO bending at the tip apex<sup>21</sup> prevents a structural analysis.

To verify our structural assignment, we performed MALDI-TOF and NMR<sup>16</sup> measurements of the molecules. Both mass and NMR spectra fully support our structural analysis. Note that, without expecting the reaction product that was identified from AFM, the NMR spectrum is not easily converted into a structural model. To further analyze the geometry of the molecule, we mimicked the flattening of one phenanthrene moiety upon adsorption in the geometry calculations by forcing three of the carbon rings to be planar, and by letting the rest of the molecule relax. The resulting geometry fits very well to the geometry deduced from AFM data, as can be seen in Figure 2, including the out-of-surface tilt angles of  $\approx 10^\circ$  and  $\approx 23^\circ$ , respectively.

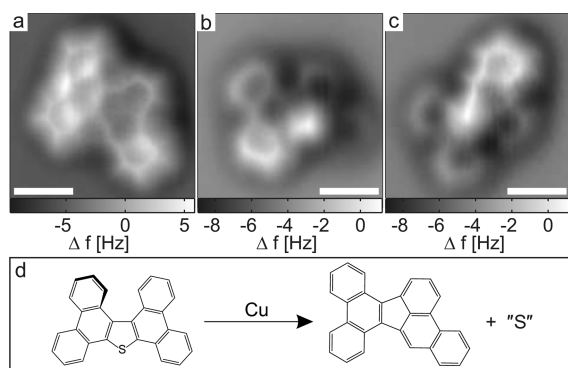
The molecular structure and the strong out-of-plane distortion suggest that the molecule can be planarized by annealing. In analogy to other on-surface reactions,<sup>22,23</sup> one might expect a ring closure by a C–C bond formation at the position of steric hindrance resulting in a C<sub>2v</sub>-symmetric molecule<sup>18</sup> (having C<sub>s</sub>-symmetry when adsorbed).

To activate such an on-surface-reaction, we annealed the sample to 200 °C. After annealing, we analyzed 32 individual



**Figure 2.** Molecular structure identification from 3D data set: To resolve the molecular structure from  $\Delta f$  imaging, the 3D data set is displayed along three different planar cross sections. While one of them is parallel to the surface, the other two are aligned with respect to the two carbon rings that are tilted with respect to the surface (green and red). (a) 3D representation of molecular structure including the cross sections, along which the experimental data is displayed in (b). (c) Top view of the geometric structure for reference.<sup>17</sup> (d, e) Top views of 3D data with only one of the tilted cross sections in each panel (scale bars 5 Å).

molecules from atomically resolved AFM images. Further molecules were identified to be of the same species as some of the atomically resolved ones from their appearance in STM images (not shown). Via the atomically resolved AFM images we found four different species, which together account for 30 out of the 32 individual molecules. The remaining two individual molecules must have formed from more than just a single reactant and are disregarded here. One of the four species is the nonreacted DPAT molecule. This observation confirms that the anneal temperature is just at the verge of initiating the reaction. From the annealing time of 2 min and the observation that 25 out of 30 molecules have reacted, we estimate an activation energy of  $\approx 1.4$  eV. The constant-height AFM images of the three remaining species are shown in Figure 3a–c. One of the latter is completely planar, so that its structure can be readily identified from the AFM image (see



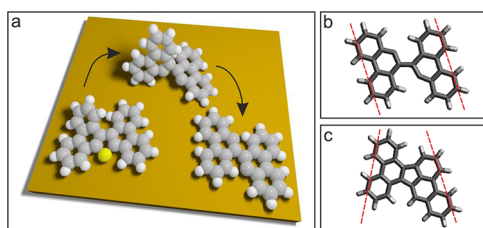
**Figure 3.** Reaction products: (a–c) Constant height  $\Delta f$ -images of different reaction products. All products are planar (a) or almost planar (b, c), so that the two phenanthrene moieties can be clearly identified. The latter are attached to each other in a nonsymmetric way and are clearly *not* aligned (anti-) parallel. Imaging parameters:  $\Delta z = 3.2$  Å,  $\Delta z = 2.55$  Å, and  $\Delta z = 2.75$  Å with respect to the STM set point ( $I = 1$  pA,  $V = 0.1$  V) (scale bars 5 Å). (d) Reaction scheme for planar product.

Figure 3a and d) as phenanthro(9,10-*e*)acephenanthrylene.<sup>24</sup> None of the bond angles is particularly small, which may indicate that the sulfur was eliminated from the molecule. The molecule does *not* have an out-of-plane mirror symmetry that would be expected for the simple reaction pathway for planarization discussed above.<sup>18</sup> Instead, the two phenanthrene moieties are now connected to each other at the center of the molecule with different orientations.

Apart from the planar reaction product shown in Figure 3a, there are two more species, displayed in Figure 3b and c, that are not completely planar. While the entire structure is not directly apparent from their AFM images, these do allow for an identification of the orientation of the two phenanthrene moieties. In both cases, the orientation agrees with the one of the planar reaction product. As the central part of the molecule cannot be imaged so clearly, it is at first glance unclear whether a five-membered ring has formed at their center in analogy to the planar reaction product. However, the five-membered ring enforces an orientation of the two phenanthrene moieties in an angle of  $150^\circ$ , whereas a single bond between the two would result in an angle of  $180^\circ$  (see Figure 4b and c). As the observed angle in the AFM images agrees with the former, we conclude that the five-membered ring has to be present. Most probably, the sulfur atom is still present in these molecular structures, giving rise to the additional feature at the center of the target molecule.

Hence, from the AFM images one can assign the backbone of all three products as phenanthro(9,10-*e*)acephenanthrylene—a molecule that has been synthesized from phenanthrene, using wet chemistry.<sup>25</sup>

We now turn to the discussion of a potential pathway of the observed reaction. By comparing the reactant and the product (see Figure 3d) and their respective symmetries, one realizes that the two phenanthrene moieties are interconnected quite differently. As illustrated in Figure 4a, a reaction pathway involving the sulfur atom elimination, an (almost)  $180^\circ$  out-of-plane rotation of one of them, and, finally, a ring closure yield the observed molecular structure. Note that, after a bond



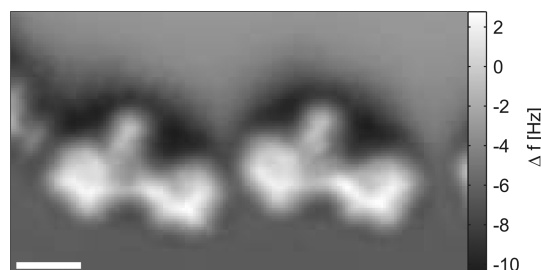
**Figure 4.** Potential reaction pathway: (a) 3D representation of the proposed reaction pathway starting with DPAT on the left-hand side. The experimentally observed nonsymmetric arrangement of the two phenanthrene moieties can be explained by a reaction pathway involving an out-of-plane rotation of one of the phenanthrene moieties as depicted by the models from left to right. (b, c) Stick models of planar reaction products without (b) and with (c) pentagonal ring closure, respectively. The dashed red lines highlight the different alignment of the phenanthrene moieties in both cases. In comparison, all the experimentally observed structures indicate a ring closure in their center. Molecular models in panels (b) and (c) were structurally optimized and displayed using Avogadro.<sup>17</sup>

cleavage at the sulfur atom, the two phenanthrene moieties are connected by a single  $\sigma$ -bond only, facilitating the rotation. Without such an out-of-plane rotation, a reorganization of *both* bonds that connect the two phenanthrene moieties would be required. Such a pathway would not be easily reconciled with the observation of almost all reaction products still consisting of two phenanthrenes—though we cannot exclude such bond reorganization from our data. We therefore propose C–S bond cleavage, the out-of-plane rotation of one phenanthrene moiety, and a ring closure as the reaction pathway. As has been observed previously,<sup>26,27</sup> the surface can coordinate atoms with open bonds that occur from e. g. the sulfur elimination and thereby stabilize intermediate structures, such as the one displayed in Figure 4b. It may well be that the hydrogen that is released upon pentagonal ring closure will saturate the open bond in the final product as shown in Figure 4c. While the planar structure of the reaction product may indicate the saturation of all bonds by hydrogen atoms, we cannot exclude that instead the respective atoms are coordinated to the substrate.

In contrast to our observation, in an earlier work<sup>18</sup> the symmetric reaction product dibenzo[2,3:10,11]perylene[1,12-*bcd*]thiophene was obtained from DPAT molecules in the presence of copper particles serving as a catalytic surface. It is well-known that at step edges the reactivity of the substrate is enhanced<sup>26,28,29</sup> which possibly enables alternative reaction pathways. Bearing this in mind, we imaged substrate defect step edges, to which molecules are attached, after annealing. Interestingly, in a corresponding AFM image displayed in Figure 5, we find planar molecules that can be readily identified as containing one of the two phenanthrene moieties only. Hence, step edges obviously facilitate the dissociation of the molecules separating the two phenanthrene moieties. This observation resolves the apparent contradiction of different reactions, as one might speculate that at even higher anneal temperatures the two phenanthrenes can reunite to form a symmetric reaction product.

## CONCLUSION

In summary, we analyzed the nonplanar adsorption geometry of DPAT molecules adsorbed on Cu(111) from atomically resolved 3D FM-AFM data. Upon annealing, the molecules



**Figure 5.** Reaction products at substrate step edge: AFM image of phenanthrene units attached to a copper step edge after sample annealing. The higher reactivity at step edges apparently facilitates dissociation of molecules. Imaging parameters:  $\Delta z = 1.55 \text{ \AA}$  with respect to the STM set point ( $I = 1 \text{ pA}$ ,  $V = 0.1 \text{ V}$ ) at the lower laying terrace in the top part of the image (scale bar  $5 \text{ \AA}$ ).

undergo an on-surface planarization reaction. The reaction pathway is proposed to involve (i) C–S bond cleavage, (ii) an (almost)  $180^\circ$  out-of-plane rotation of one phenanthrene moiety, and (iii) aromatization by dehydrogenation. Finally, the higher reactivity at step edges leads to a dissociation of the molecule separating the two phenanthrene moieties.

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### Notes

The authors declare no competing financial interest.

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## REFERENCES

- Gross, L.; Mohn, F.; Moll, N.; Liljeroth, P.; Meyer, G. *Science* **2009**, *325*, 1110.
- Gross, L.; Mohn, F.; Moll, N.; Meyer, G.; Ebel, R.; Abdel-Mageed, W. M.; Jaspars, M. *Nat. Chem.* **2010**, *2*, 821.
- Mohn, F.; Repp, J.; Gross, L.; Meyer, G.; Dyer, M. S.; Persson, M. *Phys. Rev. Lett.* **2010**, *105*, 266102.
- Albrecht, F.; Neu, M.; Quest, C.; Swart, I.; Repp, J. *J. Am. Chem. Soc.* **2013**, *135*, 9200.
- de Oteyza, D. G.; Gorman, P.; Chen, Y.-C.; Wickenburg, S.; Riss, A.; Mowbray, D. J.; Etkin, G.; Pedramrazi, Z.; Tsai, H.-Z.; Rubio, A.; Crommie, M. F.; Fischer, F. R. *Science* **2013**, *340*, 1434.
- Riss, A.; Wickenburg, S.; Gorman, P.; Tan, L. Z.; Tsai, H.-Z.; de Oteyza, D. G.; Chen, Y.-C.; Bradley, A. J.; Ugeda, M. M.; Etkin, G.; Louie, S. G.; Fischer, F. R.; Crommie, M. F. *Nano Lett.* **2014**, *14*, 2251.
- Pavliček, N.; Fleury, B.; Neu, M.; Niedenführ, J.; Herranz-Lancho, C.; Ruben, M.; Repp, J. *Phys. Rev. Lett.* **2012**, *108*, 086101.
- Hanssen, K.; Schuler, B.; Williams, A. J.; Demissie, T. B.; Hansen, E.; Andersen, J. H.; Svenson, J.; Blinov, K.; Repisky, M.; Mohn, F.; Meyer, G.; Svendsen, J.-S.; Ruud, K.; Elyashberg, M.; Gross, L.; Jaspars, M.; Isaksson, J. *Angew. Chem., Int. Ed.* **2012**, *51*, 12238.

- (9) Pavliček, N.; Herranz-Lancho, C.; Fleury, B.; Neu, M.; Niedenführ, J.; Ruben, M.; Repp, J. *Phys. Status Solidi B* **2013**, *250*, 2424.
- (10) Schuler, B.; Liu, S.-X.; Geng, Y.; Decurtins, S.; Meyer, G.; Gross, L. *Nano Lett.* **2014**, *14*, 3342.
- (11) Schuler, B.; Liu, W.; Tkatchenko, A.; Moll, N.; Meyer, G.; Mistry, A.; Fox, D.; Gross, L. *Phys. Rev. Lett.* **2013**, *111*, 106103.
- (12) Giessibl, F. J. *Appl. Phys. Lett.* **2000**, *76*, 1470.
- (13) Albrecht, T. R.; Grütter, P.; Horne, D.; Rugar, D. *J. Appl. Phys.* **1991**, *69*, 668.
- (14) Giessibl, F. J. *Rev. Mod. Phys.* **2003**, *75*, 949.
- (15) CPMD V3.15 Copyright IBM Corp 1990–2011, Copyright MPI fuer Festkoerperforschung Stuttgart 1997–2001.
- (16) Herranz-Lancho, C. Synthesis and Characterization of Molecules for Electronic Devices. Ph.D. Thesis, Université de Strasbourg, 2013.
- (17) Molecular models are displayed using Avogadro, an open-source molecular builder and visualization tool, version 1.1.0, <http://avogadro.openmolecules.net/>.
- (18) Zander, M.; Franke, W. H. *Chem. Ber.* **1973**, *106*, 2752.
- (19) In constant current imaging mode, the  $\Delta f$ -signal is influenced by the z-feedback. However, the  $\Delta f$ -image shows much more asymmetry between the two phenanthrene units than the topography.
- (20) Mohn, F.; Gross, L.; Meyer, G. *Appl. Phys. Lett.* **2011**, *99*, 53106.
- (21) Moll, N.; Gross, L.; Mohn, F.; Curioni, A.; Meyer, G. *New J. Phys.* **2010**, *12*, 125020.
- (22) Cai, J.; Ruffieux, P.; Jaafar, R.; Bieri, M.; Braun, T.; Blankenburg, S.; Muoth, M.; Seitsonen, A. P.; Saleh, M.; Feng, X.; Müllen, K.; Fasel, R. *Nature* **2010**, *466*, 470.
- (23) Treier, M.; Pignedoli, C. A.; Laino, T.; Rieger, R.; Müllen, K.; Passerone, D.; Fasel, R. *Nat. Chem.* **2011**, *3*, 61.
- (24) We cannot exclude partial dehydrogenation of the reaction product on the surface.
- (25) Franck, H.-G.; Buffleb, H. *Liebigs Ann. Chem.* **1967**, *701*, 53.
- (26) Hla, S.-W.; Bartels, L.; Meyer, G.; Rieder, K.-H. *Phys. Rev. Lett.* **2000**, *85*, 2777.
- (27) Zhao, A.; Li, Q.; Chen, L.; Xiang, H.; Wang, W.; Pan, S.; Wang, B.; Xiao, X.; Yang, J.; Hou, J. G.; Zhu, Q. *Science* **2005**, *309*, 1542.
- (28) Dahl, S.; Logadottir, A.; Egeberg, R. C.; Larsen, J. H.; Chorkendorff, I.; Törnqvist, E.; Nørskov, J. K. *Phys. Rev. Lett.* **1999**, *83*, 1814.
- (29) Vang, R. T.; Honkala, K.; Dahl, S.; Vestergaard, E. K.; Schnadt, J.; Lægsgaard, E.; Clausen, B. S.; Nørskov, J. K.; Besenbacher, F. *Nat. Mater.* **2005**, *4*, 160.