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Donor-Acceptor Properties of a Single-Molecule Altered by On-Surface Complex Formation

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Abstract

Electron donor-acceptor molecules are of outstanding interest in molecular electron-
ics and organic solar cells for their intramolecular charge transfer controlled *via* electri-
cal or optical excitation. The preservation of their electronic character in the ground
state upon adsorption on a surface is cardinal for their implementation in such single-
molecule devices. Here, we investigate by atomic force microscopy and scanning tunnel-
ing microscopy a prototypical system consisting of a π -conjugated tetrathiafulvalene-

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2
3 fused dipyridophenazine molecule adsorbed on thin NaCl films on Cu(111). Depending
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5 on the adsorption site, the molecule is either found in a nearly undisturbed, free state
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7 or in a bound state. In the latter case the molecule adopts a specific adsorption site
8
9 leading to the formation of a chelate complex with a single Na^+ alkali cation pulled out
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11 from the insulating film. Although expected to be electronically decoupled, the charge
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13 distribution of the complex is drastically modified leading to the loss of the intrinsic
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15 donor-acceptor character. The chelate complex formation is reversible with respect to
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17 lateral manipulations enabling tunable donor-acceptor molecular switches activated by
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19 on-surface coordination.
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22 **Keywords:** atomic force microscopy, scanning tunneling microscopy, fused donor-
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24 acceptor molecules, local contact potential difference, single-molecule, force spectroscopy,
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26 coordination.
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31 Electron Donor-Acceptor (DA) molecules are highly regarded in nanoscience for their
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33 properties at the base of a emerging class of single molecule based electronic devices.¹⁻⁵
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35 DA molecules with spatially separated electron donating and accepting moieties allow a
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37 controlled internal charge transfer.⁶ The spatial separation of the highest occupied molec-
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39 ular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) allows excited
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41 electron-hole pairs to be split upon photon absorption, thus making DA molecules useful for
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43 the conversion of light into electrical current leading to applications in organic solar cells.⁷
44
45 Furthermore, electrons can be tuned to resonantly tunnel in and out of the molecule by
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47 applying suitable bias voltages. Since there is a preferential direction for electron tunnel-
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49 ing through the molecule, DA molecules can also rectify the current and act as molecular
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51 diodes.^{8,9} The spatial separation of the electronic orbitals as well as the charge distribution
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53 in the ground state determine the properties of DA molecules upon excitation. The question
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55 whether these particular properties of DA molecules are still preserved upon adsorption on
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57 a surface is cardinal for their integration in molecular electronics.
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At the molecular level, subtle molecule-surface interactions may induce several stable adsorption configurations which depend on the alignment of the molecular structure with respect to the underlying substrate. Using high spatial resolution of non-contact atomic force microscopy (AFM) and scanning tunneling microscopy (STM), the interplay between structural and electronic properties of molecules as well as the visualization of on-surface chemical processes can be elucidated down to the atomic scale.^{10–17} Recently, the small fused DA molecule, TTF-PYZ₂, was reported to adopt two conformational states on thin NaCl films on Cu(111).¹⁰ By mapping the local contact potential difference (LCPD), the charge distribution of single molecules^{10,18} was found to be similar to the gas phase, thus demonstrating electronic decoupling from the metallic substrate.^{11,19–21} Pioneering studies with individual atoms have further demonstrated the capability of charging on NaCl films.^{22–24} The charge state was shown to be related with the adsorption position on the NaCl lattice which also induced the structural relaxation of the underlying atomic lattice.^{24,25} In contrast, site-dependent charge states were not reported for molecules on thin insulating films since the molecules are usually weakly adsorbed through van der Waals interactions. However, the intrinsic DA character of the molecules implies the coexistence of electron excess and depletion regions at the intramolecular level. Such particular charge distribution may locally interact with the alternating charges of the NaCl film like in the charged adatom case if those regions are sufficiently localized.

Here, we report the investigation of the charge states of a prototypical DA molecule, the tetrathiafulvalene-fused dipyridophenazine (TTF-dppz) (Fig. 1a),¹ with respect to its adsorption site on a NaCl thin film. Using combined STM and high-resolution AFM measurements, we show that in the favored configuration oriented along the apolar axis of NaCl ([001]) the intramolecular charge distribution changes compared to the other adsorption configurations. This electronic peculiarity results from the local formation of a chelate complex between the dipyridophenazine (dppz) end of the molecule and a single Na⁺ cation pulled out from the NaCl film. As a result of the chelate complex formation, the intramolecular

charge distribution is dramatically altered, as revealed by force spectroscopy and density functional theory calculations, which leads to the loss of the intrinsic DA character defined by the spatial distribution of the HOMO localized on a donor and the LUMO on an acceptor in the ground state.¹⁰ Through lateral molecule-manipulations with the tip, complex formation can be suppressed by varying the adsorption position, which leads to the intrinsic DA character of the molecule. These results underline the fundamental role of atomic-scale coordination to control the electronic properties of donor-acceptor molecules at surfaces.

RESULTS AND DISCUSSION

To determine the adsorption geometries of the deposited molecules, we conducted high-resolution AFM measurements at 5 K with functionalized CO tips of the TTF-dppz as well as of the NaCl thin film.²⁶ The constant height AFM images reveal two main molecule appearances (Figs. 1b and c) which we name *bound* and *free* molecules, respectively. Statistically, we observed a ratio of approximately 2:1 between *bound* and *free* molecules upon adsorption at low temperature (Fig. S1). For the *bound* configuration, the two pyridyl rings are slightly bent downwards as shown by the darker contrast in this area. Between the TTF and the adjacent phenyl ring, the more positive contrast in the frequency shift (Fig. 1b) shows that the molecule is at this position slightly closer to the tip than above the pyridyl units. At the onset of the TTF unit, a bright stripe and two bright spots corresponding to stronger repulsive forces are visible which coincide with the sulfur atom position of the TTF (Fig. 1b). The TTF-dppz molecule image suggests a downwards bending of the dppz unit towards the surface with a maximum height at the edge of the TTF unit resulting in a bridge-shape geometry. Comparing the *bound* (Fig. 1b) to the *free* configuration (Fig. 1c), two main differences are observed. First, the contrast along the molecular axis shows a rather flat profile compared to the bridge-shape of the *bound* configuration. Moreover, the two last

Figure 1: **Structure and AFM images of TTF-dppz.** **a**, Chemical structure of TTF-dppz. **b,c**, Constant height Δf images of TTF-dppz of *bound* type at $z = 1.27 \text{ \AA}$ (**b**) and *free* type at $z = 0.65 \text{ \AA}$ (**c**). The heights are referenced to the STM setpoint on the NaCl layer at a sample bias of $V_b = 0.3 \text{ V}$, $I_t = 3 \text{ pA}$. The AFM images were recorded on NaCl(2ML)/Cu(111) at 5 K with a CO terminated tip at $V_b = 0 \text{ V}$. At the top left in **b** a CO molecule is adsorbed nearby. **d,e**, Corresponding simulated AFM images calculated with the probe-particle model.

sulfur atoms of the TTF are slightly tilted for the *free* molecule. Figures 1d and e show simulated AFM images using the probe-particle model described in ref.²⁷. In this model a probe-particle, which represents a CO molecule, is connected to a tip with a spring and is used to determine the interaction between tip and sample in AFM measurements, assuming a specific molecular adsorption geometry calculated by Density Functional Theory (DFT) beforehand. The simulation of the *bound* configuration (Fig. 1d) reproduces well the bridge-shape of the AFM image (Fig. 1b) as well as the characteristic bright contrast at the TTF moiety. However, the deformation of the phenyl ring and the adjacent part of the TTF is not visible. In the simulated AFM image the molecule stays in its relaxed adsorption geometry and does not have any flexibility which, presumably, plays a role in the contrast of the AFM experiments. The *free* configuration simulation (Fig. 1e) resembles the AFM features of Fig. 1c such as the relative flat geometry of the molecule and the slight bending of the dppz unit. The distortion of TTF sulfur atoms is however less pronounced compared to the AFM image.

Figure 2 shows the relaxed molecule geometries with respect to their adsorption sites obtained by DFT calculations and in agreement with the determined adsorption geometries by AFM (Fig. S2).²⁶ The *bound* molecule is always adsorbed along the apolar axis of NaCl (Fig. 2a) whereas three nearly energetically equivalent configurations of the *free* molecule are found less frequently (*free1*, *free2*, *free3*). These diverse adsorption configurations also explain the slight discrepancy between calculated and experimental AFM images (Figs. 1c and e). The *bound* molecule (Fig. 2b) has however a characteristic geometry with a height maximum at the TTF onset with a height difference of ≈ 27 pm to the dppz unit pointing towards the surface (Figs. 1b and Fig. 5c). Interestingly, the side-view of Fig. 2b shows a Na⁺ cation pulled out from the NaCl substrate located below the two nitrogen atoms of the dppz unit. Note, that the Na⁺ cation is part of the NaCl layer and relatively far away from the tip, so that it cannot be directly detected in the AFM measurements without disturbing the molecule. This adsorption peculiarity also accounts for a significant decrease of the

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3 molecule adsorption energy of ≈ 200 meV with respect to all *free* configurations (Fig. 2d).
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5 This energetically most favorable configuration is indeed the most frequently observed in
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7 the experiments (approximately 70% of the observed molecules, Fig. S1). In the case of the
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9 *free* configurations (Fig. 2a), the molecule is always slightly rotated compared to the *bound*
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11 configuration. As a result, the adsorption geometry is flatter (Fig. 2c) than the bridge-shape
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13 geometry (Fig. 2b) and no pulled out Na^+ cation is observed. Further rotated adsorption
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15 configurations (*free2* and *free3*, Fig. 2a) aligned along the polar Cl axis are also found and
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17 show similar geometries as the *free1* one. All orientations misaligned with respect to the
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19 apolar NaCl axis, differ from the *bound* molecule by the absence of a Na^+ below the two
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21 nitrogens of the dppz unit and are equivalent in terms of geometries and adsorption energies
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23 (Fig. 2d) as the *free1* configuration. The sulfur atoms are preferentially located close to Na^+
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25 cations. In the case of *free1*, one of the sulfur atoms is above a Cl^- which leads to a higher
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27 adsorption height at this position. This observation also holds for the two sulfur atoms closer
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29 to the center in the bound configuration. There, the bright stripe occurs because the sulfur
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31 atoms stick out. For *free2*, no tilt or sticking out of the sulfur atoms is observed (Fig. S2e)
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33 since the sulfur atoms are symmetrically aligned with the Na^+ cations underneath. *Free3*
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35 shows again a slight tilt because of one sulfur atom lying above a Cl^- . As a conclusion we
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37 see also a contribution of the sulfur atoms to the geometrical adaptations of the molecules
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39 with respect to the adsorption site.
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42 To study the impact of those configurations on the intrinsic DA character of the TTF-
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44 dppz, we performed scanning tunneling spectroscopy (STS) combined with STM (Fig. 3)
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46 and resolved the corresponding spatial contributions of HOMO and LUMO of the adsorbed
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48 molecule.²⁸ The *bound* molecule shows resonant tunneling at -2.3 V (Fig. 3a) where the spa-
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50 tial distribution of the HOMO dominates the STM image (Fig. 3c). The HOMO is clearly
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52 located at the donor TTF and only a small tail is observed at the acceptor site. Moreover,
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54 the measured HOMO image matches reasonably the density distribution of the HOMO of
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56 the gas phase molecule calculated by DFT (Fig. S3c). The LUMO resonance of the *bound*
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Figure 2: **Simulated adsorption sites and geometries.** **a**, Adsorption sites of TTF-dppz calculated by DFT on 2 ML of NaCl and observed in experiments. The experimentally determined adsorption sites are illustrated in Fig. S2. **b,c**, Side views of the relaxed *bound* molecule showing the pulling out of the Na^+ cation and of the *free* molecule depicted in Figure 1c; the insets are orthogonal back views. **d**, Calculated adsorption energies at different adsorption sites. For the representation of the NaCl layer the atomic radii are used. Color coding: grey C, blue N, yellow S, pink Na, green Cl, white H.

molecule observed at $V_b = 1.5$ V exhibits two new lobes located at the nitrogen positions of the phenazine subunit (Fig. 3e) compared to the gap image (Fig. 3d). They can be attributed to the acceptor site and shows that a part of the LUMO is located at the dppz unit. However, there is a appreciable LUMO weight on the TTF side which clearly indicates a qualitative deviation from the calculated LUMO distribution of the gas phase molecule (Fig. S3d). This calculated LUMO distribution, centered on the dppz moiety, is in better agreement with the STM image at the LUMO resonance of the *free* molecule (Fig. 3h). Comparing the *bound* and *free* molecules, the location and shape of the LUMO resonances are quite different, as well as their energetic positions in the STS spectra. The onset of the LUMO resonance varies from 1 V for the *bound* to 1.5 V for the *free* molecule which also indicates that the electronic properties of the *free* molecules on the NaCl film are closer to

those of the gas-phase molecules than for the *bound* ones. In addition, at negative bias the observed energetic shifts of the resonances indicate a different influence on the HOMO levels between *bound* and *free* adsorption sites. Since the HOMO is strongly localized on the TTF moiety, in a first approximation, the different adsorption geometries of the TTF unit are expected to be the dominant contribution for this feature.

The electronic properties are determined by the charge distribution in the molecule. Using force spectroscopy, *i.e.* measuring $\Delta f(V)$ which is proportional to the vertical force gradient, to resolve the local contact potential difference (LCPD) between molecule and tip, the intramolecular charge distribution is accessible in real-space as previously demonstrated with donor-acceptor molecules as well as other species.^{10,18,29,30} The LCPD contrast is known to vary with the tip-sample distance, particularly at close tip-sample distance, where the CO bending can produce complex contrasts.¹⁰ In our measurements, we used copper tips to characterize the LCPD contrasts which prevent probing at close-distance regimes compared to CO-terminated tips. In that case, *i.e.* at medium distance regime, compared to the molecule dimensions, the LCPD contrast represents the partial charge distribution of the DA molecules. Figure 4 shows the LCPD maps at constant tip height of the *bound* and *free* DA molecule, respectively. The LCPD corresponds to the voltage V^* at the maximum frequency shift (Δf^*) of the parabola fitted to the quadratic dependence of the frequency shift Δf on the bias voltage V .¹⁰ The $\Delta f^*(x, y)$ maps reveal striking differences between the *bound* and *free* configurations (Fig. 4a,b). For *bound* molecules (Fig. 4a) Δf^* exhibits a minimum at the edge of the TTF donor whereas, for *free* molecules, the Δf^* drop extends over the whole molecule with an absolute minimum at the dppz location. This trend is also visible in the $\Delta f^*(x)$ profiles (Fig. 4c) extracted along the longitudinal axis of each molecule. Interestingly, the LCPD map of the *bound* molecule shows one region with a less negative LCPD compared to the surrounding NaCl layer which is attributed to local negative charges within the molecule (Fig. 4d). These negative charges are located between the phenazine subunit of the acceptor and the edge of the TTF donor. The most negative Δf^* is detected at the same

Figure 3: Energetic position and spatial distribution of the TTF-dppz' molecular orbitals. **a,b**, dI/dV spectra of *bound* and *free* molecules, recorded on the donor (TTF) and acceptor (dppz) sides. **c**, STM constant current image of the *bound* molecule at the first resonance of negative sample bias ($I_t = 1$ pA, $V_b = -2.3$ V), **d**, in the gap between the first negative and positive resonance ($I_t = 3$ pA, $V_b = 0.3$ V) and **e**, at the first resonance of positive sample bias ($I_t = 1$ pA, $V_b = 1.5$ V). **f**, STM constant current image of the *free* molecule at $I_t = 1$ pA, $V_b = -2.5$ V, **g**, at $I_t = 1$ pA, $V_b = 0.3$ V and **h**, at $I_t = 1$ pA, $V_b = 1.8$ V. The scale bars in **c-h** correspond to 1 nm. All STM/STS measurements were performed at 5 K. See Fig. S3 for a comparison with the calculated distribution of the molecular orbitals in the gas phase.

position where the topographic maximum of the *bound* geometry is observed in AFM images and DFT calculations (Fig. 1b,d). For the *free* molecule (Fig. 4e), the LCPD also reveals a negative charge at this location. In contrast to the *bound* molecule, the least negative LCPD is clearly observed at the dppz moiety. This indicates that the acceptor side of the *free* molecule is more negatively charged compared to the surrounding NaCl whereas, for the *bound* molecule, the acceptor has the same charge relative to the surface. Furthermore, the LCPD at the TTF moiety of the *free* molecule is slightly below the background LCPD which suggests a positive charge at the donor (Fig. 4e). All the features are reproduced in

Figure 4: **LCPD measurements of *bound* and *free* molecules.** **a, d,** Δf^* and corresponding LCPD map of a *bound* molecule on NaCl(2 ML)/Cu(111). **b, e,** Δf^* and LCPD maps of a *free* molecule adsorbed on NaCl(3 ML)/Cu(111). **c, f,** LCPD line scans along the molecular axis of *bound* and *free* molecules on NaCl(3 ML)/Cu(111), which are shown in the STM topography inset ($I_t = 3$ pA, $V_b = 0.3$ V) and measured with exactly the same Cu tip. For all LCPD measurements a Cu tip was used and the sample was kept at 5 K.

the V^* line scans (Fig. 4f) measured along the longitudinal axis of both molecules. The negative charge region is laterally shifted within the molecule. These observations demonstrate that *bound* and *free* molecules have different charge distributions which only depend on the adsorption site.

When the TTF-dppz molecule is aligned along the apolar axis of NaCl, the bending of the dppz unit towards the surface is triggered by the coordination of the N atoms of the pyridyl units in dppz to the positively charged Na^+ cation in order to favor the formation of a chelate complex. Dppz is indeed known for its metal ion chelating property, mainly with ruthenium(II) and iron(II) ions^{31,32} but also with rare earth ions.^{33,34} Compared to transition metal ions, alkali metal ions exhibit weaker complexing abilities and chelation is usually essential to complex formation.³⁵

A common notion is that the sodium ion and its higher congeners are not prone to

Figure 5: **Switching between *bound* and *free* molecules.** **a,b**, STM topography of *bound* and *free* molecule at $I_t = 3 \text{ pA}$, $V_b = 0.3 \text{ V}$. Between the images the voltage was ramped until a jump in the current occurred and the adsorption site changed. These voltage values are given next to the arrows. The tip was positioned above the topographic maximum of the molecule. The switching of the adsorption site could be done in both directions. The scale bars are 500 pm. **c,d**, Corresponding height profiles of the two molecules. *Bound* molecules show one distinct topographic maximum which forms the bridge shape geometry, whereas *free* molecules exhibit two local maxima.

generate coordination compounds and only the introduction of crown ethers and cryptands opened up different avenues for this field. From the viewpoint of Hard-Soft-Acid-Base theory, the coordination of sodium ions to ligands with oxygen donors clearly dominates over the coordination with nitrogen (soft donor) containing ligands, since Na^+ qualifies as a hard acid and oxygen as a hard donor. Consequently, there are only few reports on complexation of sodium ions with N^N chelating ligands.^{36–38} Interestingly, the study of Qian *et al.* shows mono- and bidentate complexation modes for the coordination of sodium ions

with phenanthroline-based ligands.³⁷ For this rare case of a monodentate linkage, the Na-N coordination distances are in the range of 2.66 Å to 2.73 Å, and the uncoordinated Na...N distances are larger than 3.19 Å. For the chelating N^N coordination, the Na-N bond lengths range from 2.44 Å to 2.46 Å, and the chelating N-Na-N bond angle amounts to 67.7°. Not unexpected, the above-mentioned bond distances are larger than those for a corresponding O^O chelation mode of sodium ions, which typically shows values around 2.32 Å.³⁹ From the DFT simulations (Fig.2), we extract the bond length between the Na and the pyridyl N atoms of ~ 2.63 Å. Therefore, our measurements suggest the formation of a complex between Na⁺ and N which is in the range of the referred Na-N coordination distance. The smaller bond length resulting from the bidentate complexation mode is likely not reached in our system since the Na⁺ pulled out from the ionic film is still bound to the lattice.

As a result, the Na⁺ alkali metal ion attracted by the lone pairs of the dppz N atoms forms a chelate complex which induces the intramolecular redistribution of charges. A reduction of negative charge at the acceptor side is indeed observed in the LCPD measurements of such complex (Fig. 4d,f) which suggests a partial charge transfer from the N atoms to the Na⁺ altering the acceptor charge state. This interpretation is further confirmed in the STS measurements of the LUMO resonances shifting from the calculated LUMO distribution (Fig. S3d) to lower energies (Fig. 3a) as shown after complex formation.³² In contrast, the *free* molecules do not bind to a Na⁺ cation and show a charge distribution (Fig. 4e) as predicted for DA molecules with a positive charge located at the donor region and a negative charge at the acceptor site. The simulated electrostatic potential of the *free* TTF-dppz (Fig. S3a) qualitatively agrees with the measured LCPD (Fig. 4e). For the *bound* type however, the variation of the electrostatic potential experimentally observed for the chelation is not as significant in the DFT calculations. Note that the charge around the Na⁺ cation cannot be investigated directly in LCPD measurements since it is located close to the substrate and hence too far away from the tip to detect a LCPD difference at the position between

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3 molecule and substrate.
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5 In order to demonstrate that only the adsorption sites affect the charge distribution and
6 exclude the influence of possible defects of the NaCl thin layer, we conducted single-molecule
7 manipulation experiments through inelastic electron tunneling (Fig. 5).⁴⁰ We displaced a
8 *bound* molecule to neighboring adsorption sites to induce the variation of electronic proper-
9 ties as described above. The investigation of the initial adsorption site confirmed that no
10 defects are present below the *bound* molecule. Since the displaced molecule shows the typical
11 STM topography of the *free* molecules, thus we further illustrate the interplay between the
12 adsorption site and its molecular charge state. Switching in the opposite direction meaning
13 from *free* to *bound* was also possible. These observations demonstrate that chelate complex
14 formation of the Na⁺-TTF-dppz can be controllably reversed by atomic manipulation.
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30 CONCLUSIONS

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32 We have demonstrated that interactions of a single DA molecule with the NaCl atomic lat-
33 tice on Cu(111) result in the formation of a chelate complex with a single Na⁺ cation. This
34 chemical process depends on the positioning of the molecule with respect to the underlying
35 substrate as characterized by combined high-resolution AFM measurements and DFT cal-
36 culations. As a consequence of the Na⁺-molecule bond formation, the local alteration of the
37 acceptor charge state results in the loss of the intrinsic DA molecule properties *in the ground*
38 *state*. These findings highlight the interplay between the DA character of the molecule and
39 the alternating charges of the insulating thin film at the atomic scale. Since the charge
40 distribution in DA molecules in the ground state is strongly relevant to the properties in the
41 excited state, our results are of great relevance for applications in optical energy conversion
42 and molecular electronics where the functionality of specifically designed molecules must be
43 maintained on surfaces. Interestingly, the complex formation can be triggered reversibly by
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voltage ramps thus enabling the design of a switchable donor-acceptor molecular compound controlled by coordination on an ionic thin film.

METHODS

STM/AFM experiments. The STM and AFM measurements were performed with an Omicron GmbH low temperature system equipped with a tuning fork in the Qplus configuration⁴¹ ($k \approx 1800$ N/m, $f_{\text{res}} = 25$ kHz) and operated with Nanonis controllers RC5. For all measurements the sample was kept at 5 K. The amplitude of the tuning fork oscillation was set to a value between 50 - 100 pm. Scanning tunneling spectroscopy (STS) data was acquired with the lock-in technique. The bias modulation amplitude was 10 mV at a frequency of 520 Hz. The bias voltage is given by the potential of the sample referred to the tip. The STM/AFM tip consists of a tungsten wire that is gently indented into the clean Cu(111) surface several times to obtain a Cu tip. All STM images were recorded with a Cu tip, unless a different termination is explicitly stated. CO tip terminations were done by picking up single CO molecules adsorbed on the NaCl film. For the LCPD mapping, a measurement grid was layed over the molecule. At each point of the grid a bias sweep was performed and the frequency shift was detected. Between the bias sweeps the position was adjusted by atom tracking.⁴²

Sample preparation. The Cu(111) surface was prepared in ultra high vacuum by sputtering and annealing cycles. Two to three monolayers of sodium chloride (NaCl, purchased from Sigma-Aldrich GmbH) were then deposited on the surface at room temperature. Tetrathiafulvalene-fused dipyrrophenazine (TTF-dppz) molecules were thermally evaporated onto the cold substrate ($T_{\text{sample}} < 10$ K) to obtain single-molecules.

DFT Calculations Calculations were performed using the projector augmented plane-wave⁴³ method and the PBE functional⁴⁴ as implemented in the VASP 5.3 code.⁴⁵ Plane-wave cutoffs were set to 400 eV for wave-functions and 644 eV for potential. The rectangular

simulation box of x,y,z dimensions 28.2x22.56x20.00 Å corresponds to a 4x5 super-cell containing a NaCl (100) slab composed of 2 layers. Only the Γ point was used for k-space sampling. Convergence criterion was set to 10^{-6} eV/step for electronic and 10^{-5} eV/step for ionic relaxation.

Synthesis of 2-(1,3-dithiol-2-ylidene)-[1,3]benzodithiole-5,6-diamine (compound

2): Under N₂ protection, 6-(1,3-dithiol-2-ylidene)-[1,3]dithiolo[4,5-*f*]-2,1,3-benzothiadiazole⁴⁶ (156 mg, 0.5 mmol) was dissolved in dry THF (10 mL). Then Lithium aluminum hydride (1.25 mL, 1M in THF, 1.25 mmol) was added dropwise to the reaction mixture over a period of 30 min at 0°C. The reaction mixture was then stirred for 30 min under nitrogen. Afterwards it was slowly quenched by saturated aqueous NH₄Cl (1 mL) at 0°C. The resulting precipitate was collected and washed by water and Et₂O to give analytically pure product as a light yellow solid. Yield: 131 mg (91%); ¹H NMR (300 MHz, DMSO-*d*₆) δ = 6.72 (s, 2H), 6.54 (s, 2H), 4.68 (s, 4H); ¹³C NMR (75.5 MHz, THF-*d*₈): δ = 134.8, 121.6, 119.8, 111.5, 107.6, 107.5 ppm; HRMS (ESI): *m/z* calcd for C₁₀H₈N₂S₄: 283.9565; found 283.9556.

Synthesis of 2-(1,3-dithiol-2-ylidene)-[1,3]dithiolo[4,5-*i*]dipyrido[3,2-*a*:2,3-*c*]phenazine

(TTF-dppz): A solution of compound 2 (57 mg, 0.2 mmol) and 1,10-phenanthroline-5,6-dione (41 mg, 0.2 mmol) in ethanol (40 mL) was refluxed for 3 h under N₂. After filtration, the precipitate was washed by CH₂Cl₂ and purified by recrystallization in toluene to give analytically pure TTF-dppz as a deep blue powder (50 mg, 55%). ¹H NMR (300 MHz, DMSO-*d*₆) δ = 9.48 (d, *J* = 8.1, 2H), 9.21 (d, *J* = 4.4 Hz, 2H), 8.40 (s, 2H), 7.92 (dd, *J* = 8.1, 4.4 Hz, 2H), 6.82 (s, 2H); ¹³C NMR is not available due to its limited solubility. HRMS (ESI): *m/z* calcd for C₂₂H₁₀N₄S₄: 457.9861; found 457.9866. Further details on the synthesis of TTF-dppz can be found in the Supplementary Information.

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12 E.M., S.D., T.G. and R.P. planned the experiments. T.M. performed the STM/AFM ex-
13 periments, processed the data and wrote the manuscript. P.H. and P.J. performed the DFT
14 calculations. Y.G., X.L., S.X.L. and S.D. synthesized the molecular compounds. All co-
15 authors commented the data and contributed to the manuscript.
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ASSOCIATED CONTENT

Supporting Information Available

Further details concerning experiments, simulations and synthesis can be found in the Supplementary material section. This material is available free of charge via the Internet at <http://pubs.acs.org/>.

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Graphical TOC Entry