Cyclophane with eclipsed pyrene units enables construction of spin interfaces with chemical accuracy†

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Advanced functionality in molecular electronics and spintronics is orchestrated by exact molecular arrangements at metal surfaces, but the strategies for constructing such arrangements remain limited. Here, we report the synthesis and surface hybridization of a cyclophane that comprises two pyrene groups fastened together by two ferrocene pillars. Crystallographic structure analysis revealed pyrene planes separated by ~352 pm and stacked in an eclipsed geometry that approximates the rare configuration of AA-stacked bilayer graphene. We deposited this cyclophane onto surfaces of Cu(111) and Co(111) at submonolayer coverage and studied the resulting hybrid entities with scanning tunnelling microscopy (STM). We found distinct characteristics of this cyclophane on each metal surface: on non-magnetic Cu(111), physisorption occurred and the two pyrene groups remained electronically coupled to each other; on ferromagnetic Co(111) nanolands, chemisorption occurred and the two pyrene groups became electronically decoupled. Spin-polarized STM measurements revealed that the ferrocene groups had spin polarization opposite to that of the surrounding Co metal, while the pyrene stack had no spin polarization. Comparisons to the non-stacked analogue comprising only one pyrene group bolster our interpretation of the cyclophane’s STM features. The design strategy presented herein can be extended to realize versatile, three-dimensional platforms in single-molecule electronics and spintronics.

Introduction

To design the electronic properties of nanoscale interfaces demands the precise arrangement of their structures. Magnetoresistance arises at interfaces between ferromagnetic electrodes and organic compounds if chemisorbed proximal aromatic group would form a hybrid molecular magnet, while has been observed only in limited cases in which it could be characterized sparingly.17,20–26 We hypothesized that such an AA-packed graphene bilayer would exhibit interfacial magnetoresistance when placed in contact with a ferromagnetic electrode, as has been found at boundaries between zinc methyl phenalenyl molecules and cobalt.2 Chemisorption of the lower aromatic group would form a hybrid molecular magnet, while the upper ring would act as a spin analyser because it retains sharp states with spin-polarization induced by \( \pi-\pi \) interactions.24,27 Such spin interface effects have yet to be demonstrated at the level of a single molecule.

graphene sheets if one is twisted by a “magic angle” (~1.1°) with respect to the other.14,15 These uncommon electronic properties arise because atoms at interfaces have been arranged in precise configurations that are challenging to achieve.

It is difficult to arrange two polyaromatic hydrocarbon (PAH) groups or continuous graphene sheets at their van der Waals (vdW) distance such that all carbon atoms of the upper sheet lie atop those of the bottom one. In such a fully eclipsed arrangement (AA-stacked), the frontier orbitals of the sheets overlap constructively and electronic repulsion is maximized (Fig. 1).10,11 The other limiting case is the half-eclipsed (AB-stacked) arrangement that is native to graphite.16 AA-stacked bilayer graphene has been a subject of theoretical interest,17–19 but it has been observed only in limited cases in which it could be characterized sparingly.17,20–26 We hypothesized that such an AA-packed graphene bilayer would exhibit interfacial magnetoresistance when placed in contact with a ferromagnetic electrode, as has been found at boundaries between zinc methyl phenalenyl molecules and cobalt.2 Chemisorption of the lower aromatic group would form a hybrid molecular magnet, while the upper ring would act as a spin analyser because it retains sharp states with spin-polarization induced by \( \pi-\pi \) interactions.24,27 Such spin interface effects have yet to be demonstrated at the level of a single molecule.
Results and discussion

Cyclophane design, synthesis, and structure

Our cyclophane design uses pyrene as an aromatic core that balances size, chemical processability, and vapor pressure.\textsuperscript{39–42} Pyrene chemisorbs onto metal surfaces with a face-on geometry,\textsuperscript{42} and its electronic states are influenced minimally by substituents at the 2- and 7-positions because the relevant frontier orbitals are bisected by a nodal plane (Fig. 1 and S13\textsuperscript{†}).\textsuperscript{43,44}

Using the advances that have been made in Suzuki–Miyaura cross-couplings, we avoided the multistep approaches that have often been used to synthesize cyclophanes.\textsuperscript{39} 2,7-Bis[4,4,5,5-tetramethyl-1,3,2-dioxaborol-2-yl]pyrene ([Bpin]\textsubscript{2}pyrene), readily produced via direct Ir-catalyzed borylation,\textsuperscript{45,46} was cross-coupled with 1,1\texttextsuperscript{′}-diiodoferrocene\textsuperscript{47} utilizing [1,1\texttextsuperscript{′}-bis(di-tert-butylphosphino)ferrocene]dichloropalladium(II)\textsuperscript{48} as catalyst in N,N-dimethylformamide (DMF) (Scheme 1).

We obtained the target cyclophane 2 in <1% yield after purifying it by gradient sublimation (Fig. S1). Several oligomers and larger macrocycles were also obtained. We synthesized 2,7-diferrocenylpyrene 1 under similar catalytic conditions with a yield of 65%, compared with the 2% yield obtained by Preuß \textit{et al.} using Negishi coupling, illustrating the advantage of our optimized procedure.\textsuperscript{39} We confirmed the identities of 1 and 2 by matrix-assisted laser desorption/ionization mass spectrometry (MALDI-MS, Fig. S3 and S4\textsuperscript{†}), elemental analysis, nuclear magnetic resonance (NMR, Fig. S2 (1\textsuperscript{†})) and Fourier-transform infrared (FT-IR) spectroscopy (Fig. S6\textsuperscript{†}).

X-ray diffraction of a single crystal of 2 that we obtained from gradient sublimation confirms the resemblance of the pyrene groups to AA-stacked graphene bilayers (Fig. 2). The space group of cyclophane 2 is the same (P2\textsubscript{1}/c) as reported for 1,\textsuperscript{49} with intermolecular C–H⋯π contacts that resemble the dimeric herringbone packing of pyrene (Fig. S7 and S8\textsuperscript{†}).\textsuperscript{49} The intramolecular spacing between the consensus planes of the pyrene groups is 351.64(15) pm, and these planes have bonding angles\textsuperscript{49} of just 3.8(4)°. The pyrene groups are offset by only ~50 pm from perfectly eclipsed. The vdW contact distance between two AB-stacked graphene sheets is ~335 pm,\textsuperscript{16,50} the spacing of the AA arrangement has not previously been measured with the accuracy of our X-ray diffraction, but a distance of 355 pm has
been estimated.21 The distance of ~352 pm between pyrene groups in 2 appears to be the closest that two PAHs can be forced with the AA arrangement without distortion. The distance between cyclopentadienyl centroids in 2 is 330.10(18) pm, only marginally larger than the 329.6 pm of ferrocene;21 the pyrene groups are neither stretching nor contracting these pillars and are thus at the vdW distance for the AA configuration.

In other PAH-containing cyclophanes,28 the conflict between shorter bridge groups and repulsive interactions bends the aromatic planes dramatically (e.g. by ~37.9° in ethyl28,30 or ~16.9° in propyl29,30,35 bridged cyclophanes). For discrete aromatic bilayers linked by longer bridges (e.g. butylene29 or –CH₂CH₂OCH₂CH₃30) lesser bending and large lateral offset values were reported that are similar to the values observed in natural graphite (AB-stacking). The two Fc bridges in 2 gently force the two pyrene planes into an eclipsed arrangement at a threshold distance that prevents both bending and relaxation into an AB-stacked arrangement.

STM measurements of cyclophane 2 and reference compound 1

We performed STM measurements at ~4 K of 1 and 2 that were deposited at submonolayer coverages atop nanoscale Co(111) islands on a Cu(111) surface. Triangle-shaped, double-layered Co(111) islands (5–25 nm across) on Cu(111) have out-of-plane magnetization,29–33 and the Cu(111) regions are nonmagnetic. The interactions of aromatic adsorbates with transition metal surfaces have been the subject of extensive theoretical and experimental investigations.1–9 Co has 3d bands that are narrow, spin-split, close to the Fermi level, and partly filled, leading to strong chemisorption. This may be understood to result from Co d-orbitals that protrude at the positions of the Co surface atoms normal to the surface and interact with π-orbitals of pyrene. The broad 3d bands of Cu are fully occupied with energies well below the Fermi level, thus fostering weaker molecule–metal interactions than Co, i.e. physisorption.60

A topographic constant-current STM plot (Fig. 3a) imaged intact molecules of 2 as protrusions of length and symmetry consistent with our crystallographic measurements on both Cu (dashed red ellipse) and Co (dashed green ellipse). We noted other fragments (dashed blue ellipse) that appeared after surface deposition. In pristine areas of Co(111), cyclophane 2 adsorbed with the longitudinal axis perpendicular to one of the island edges, which run along the crystallographic [010], [110] or [011] axis. Wang et al. previously showed that pyrene adsorbed to Cu(111) with its four ring centroids occupying bridge sites upon the metal,65 since the spacings of these centroids (~244 pm) are only slightly mismatched from the Cu lattice constant (aCu = 256 pm). A similar geometry of adsorption for 2 on Co(111) (aCo = 251 pm)65 is consistent with our data; we propose such an arrangement in Fig. 4. This excellent match of the benzoid repeat units with the Co(111) surface lattice favours strong hybridization of out-of-plane d-orbitals to molecular π-orbitals.

Cyclophane 2 adsorbed to Co(111) more strongly than to Cu(111). On Cu(111), the profile of the apparent height traced along the main axis of 2 (red plot in Fig. 3e) is flat and featureless. The corresponding profile of 2 on Co(111) (green plot in Fig. 3e) clearly differs, with a pronounced dip at the centre. The ~300 pm maxima of both of these apparent height profiles are similar to previous reports of Fc adsorbed with a face-on configuration,65 and the symmetries and lengths of the profiles agree with the adsorption scheme in Fig. 4. Cyclophane 2 tended to form aggregates on Cu(111) (white dashed ellipse in Fig. 3a), and was often found at the edges of islands, indications that its interactions with this metal were weak and physisorptive in character. On the Co islands, we found many isolated molecules of 2, suggesting that lateral diffusion was restricted and that 2 chemisorbs to Co(111).

Reference compound 1 preferentially adsorbed on both metals with a face-on geometry, and bound to Cu(111) even more weakly than did 2 (Fig. 3b). We observed no aggregates, and few isolated molecules of 1 that were static on Cu(111). In fact, measuring the profile of the apparent height for 1 atop Cu(111) proved difficult, and the only plots we obtained (red in Fig. 3d) have lower maxima than we expect for intact molecules. We do not know the reason for this discrepancy but suspect that the only molecules of 1 that remained sufficiently static for this measurement were pinned at defect sites of the substrate or had been chemically modified in a way analogous to that which we induced using tip manipulation (vide infra). There are no indications that 1 diffused across Co(111), which corroborates its chemisorption to this metal, and its Fc groups gave apparent heights of the expected ~300 pm.65

Hybridization of cyclophane 2 to Co(111) suppresses conductance through the pyrene stack

We investigated the role of the upper pyrene group on the electronic structure of cyclophane 2 by comparing the profile of its apparent height with that of 1 under similar imaging conditions. The apparent height at the centre of the profile of 2 is ~245 pm, while for 1 it is ~205 pm (Fig. 3). We might have expected the difference between these values to be larger based upon the crystallographic structures of 1 and 2, but the apparent height measured by STM in the constant-current mode convolutes the local topography with the local conductance between the surface and the tip. The lack of infinite sharpness of the STM tip hindered our plumbing of the depth of the crevice between Fc groups in 1, but the apparent height at

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the centre may be inflated by the strong chemisorption of pyr-ene to Co(111), providing a channel of conducting states between the tip and the substrate.

The apparent dip of ~55 pm at the centre of cyclophane 2 reflects the lesser tunnelling conductance of the stacked pyrene groups when compared with the Fc groups, a signal that the upper pyrene group of 2 is electronically decoupled from the lower one after it has chemisorbed to Co(111). This dip is unlikely to reflect a topographic depression because the upper pyrene group is suspended at a level that is even with the upper cyclopentadienyl rings. In contrast, on Cu(111) the upper pyrene of cyclophane 2 has an apparent height equal to those of the Fc groups.

Our observations suggest that the electronic coupling between the pyrene groups of 2 is stronger when the cyclophane is physisorbed to Cu(111) than when it is chemisorbed to Co(111). The coupling between group orbitals is proportional to their spatial overlap and inversely proportional to the difference in their energies when the groups are isolated. The reduced coupling between pyrene groups in 2 after chemisorption on Co(111) is likely to be caused by a reduced $\pi-\pi$ overlap or a shift of the energies of the lower pyrene orbitals induced by hybridization with the metal. On Cu(111), hybridization of the lower pyrene group to the metal is weaker, this energetic shift as well as the impact on the orbital overlap are smaller, and the pyrene groups remain strongly coupled. Due to the wide HOMO–LUMO gap of pyrene (Fig. S14†), the isolated pyrene group of 2 contributes little to the density of states between this bias voltage of 500 mV and the Fermi level. Similar observations were made for naphthalenediimide (NDI) cyclophanes with...
large interplanar distances adsorbed to Au(111) or Ag(111). The NDIs are electronically decoupled, and the photophysical properties of the upper NDIs are partially preserved and could be exploited in photonics.66,72

Our efforts to directly probe the states of the upper pyrene group of cyclophane 2 by STM were hindered because the molecules were not stable to measurements at larger bias voltages. This instability manifests itself in the occurrence of molecules with asymmetric appearance (black ellipse in Fig. 3a). Molecules of 2 were readily modified by placing the STM tip over a pristine area of a Co island, applying a bias voltage \( V_b = 1.5 \text{ V} \) at a setpoint current \( I_s = 50 \text{ pA} \), and moving the tip over a molecule (Fig. S9†). The tip retracted by \( \sim 350 \text{ pm} \), the apparent height of 2 at \( V_b = 1.5 \text{ V} \), and then irreversibly advanced by more than 100 pm after \( \sim 3 \text{ s} \). We subsequently collected topographic maps over the area in constant-current mode to find that the Fc portion of the molecule subjected to this treatment had been truncated (Fig. S9b†). This is consistent with other reports of the partial decomposition of Fc groups into cyclopentadienyl (Cp) and FeCp fragments on Au,73 Pt74,75 and Cu76,77 surfaces, and with the reported modification by an STM tip of a tetracetylene-bridged di-ferrocene molecule on Cu(110).78 We also applied this modification procedure to the reference molecule 1, with results that were generally similar (Fig. S10†). For a more detailed discussion see ESL†

**Spin-polarized STM of cyclophane 2 on Co(111) islands**

We mapped the spin-polarized conductance of cyclophane 2 on Co(111) islands using an antiferromagnetic Cr tip at a low bias voltage of \( V_b = -0.5 \text{ V} \) \( (I_s = 1.0 \text{ nA}, \pm 1 \text{ T}) \). Under these conditions, the molecules were sufficiently stable for us to measure SP-STM conductance maps with acquisition times of up to 6 h. Spin asymmetry maps were obtained from the difference of conductance maps measured at the same bias voltage \( V_b \) but for parallel and antiparallel alignment of tip and sample magnetizations. Spin asymmetry maps reflect spatial variations of the spin polarization of the sample’s density of states at the energy \( |e|V_b \). Further details on spin asymmetry and our acquisition methods can be found in the ESL†

We found positive spin asymmetry over the pristine Co(111) and inverted spin asymmetry over the Fc groups, while the spin asymmetry returned to zero as the tip passed over the pyrene stack (see traces along the lengths of two molecules of cyclophane 2 in Fig. 5). Theoretical calculations9,27 indicate that the upper plane of a discrete aromatic bilayer structure that is chemisorbed to a ferromagnetic metal retains sharp but spin-polarized electronic states. This polarization is induced by coupling between aromatic layers and may lead to resistive spin filter functionality. We did not observe spin polarization at the pyrene site of 2 after chemisorption to Co. One possible explanation is that the bias voltage we applied may have been too low to observe spin asymmetry at this site, since the upper pyrene
A cyclophane hybrid entity with ferromagnetic coupling between individual Fe and Fc molecule atop a Co adatom on Cu(111) to give a magnetic ferromagnetic metal surface with chemisorbed molecules, an effect that has been encountered in films (e.g. aluminium-tris(8-hydroxyquinolin) (Alq3))85 or individual molecules (e.g. phthalocyanine86 or 2,4,6-triphenyl-1,3,5-triazine (TPT)87), and theoretically explained in detail.9,45,64,81,82

We designed and synthesized a cyclophane in which ferrocene pillars hold two pyrene groups at their vdW distance in an eclipsed arrangement. The aromatic planes are virtually undistorted with a bending angle of just ~3.8° and are displaced laterally from a fully-eclipsed alignment by only ~50 pm. X-ray diffraction showed that the pyrene groups of this cyclophane are separated by 351.64(15) pm, the most precise measure to date of the vdW distance in AA-stacked graphene bilayers.

The organization of molecules at metal surfaces determines the characteristics of devices in molecular electronics and spintronics. The synthesis of 1 and cyclophane 2, compounds amenable to deposition by evaporation, allowed us to investigate and compare spin interfaces at single- and double-layered hybrid entities with chemical accuracy. Using STM, we determined that cyclophane 2 physorsors onto nonmagnetic Cu(111) with a face-on geometry, low barrier to lateral migration, and a maximum of the apparent height profile of ~300 pm. On Co(111), 2 chemisorbs with a face-on geometry, no evidence of lateral migration, and an apparent height of ~245 pm over the pyrene stack. We conclude that hybridization of the lower pyrene group to Co realigns the energy levels of the π-orbitals and reduces their overlap to weaken the electronic coupling between pyrene groups. This interlayer decoupling is consistent with our spin-polarized STM measurements that showed no spin polarization above the chemisorbed pyrene stacks. Further investigations at higher bias voltages were hindered by partial fragmentation of 2 that we confirmed with controlled manipulations using the STM tip.

We expect that the cyclophane design presented in this study can be extended to craft well-defined structures with variable aromatic stacking patterns and sizes. Over the Fe units of 2 on Co(111), we measured spin polarization of comparable strength, but opposite direction, to that of the pristine Co surface. This shows the efficiency with which Fe transmits spin polarization by the spin-dependent hybridization mechanism, and that covalent extension of Fe is an approach to enforce face-on hybridization of Fe groups to ferromagnetic surfaces. This opens a path to metalloocene nanopatterning and the construction of more complicated systems. The chemisorption of such structures to ferromagnetic metal surfaces is an important step toward engineering spin interfaces by precise orientation of their constituents in three dimensions.

Conclusions

We designed and synthesized a cyclophane in which ferrocene pillars hold two pyrene groups at their vdW distance in an eclipsed arrangement. The aromatic planes are virtually undistorted with a bending angle of just ~3.8° and are displaced laterally from a fully-eclipsed alignment by only ~50 pm. X-ray diffraction showed that the pyrene groups of this cyclophane are separated by 351.64(15) pm, the most precise measure to date of the vdW distance in AA-stacked graphene bilayers.

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Author contributions

D. E. B., C. M. S., and P. K. resources, funding acquisition, and project administration. D. E. B., J. R., and P. K. conceptualization and supervision of the study. S. S. (Schleicher), T. H., B. B. F. M., and D. E. B. STM investigation. M. M. (lead), S. S. (Sanz), J. R., and P. K. synthesis and chemical investigation. M. M., F. M., D. E. B., J. R., and P. K. writing – original manuscript. M. M., S. S., T. K., B. B., F. M., S. S. (Sanz), D. E. B., J. R., C.M.S., and P. K. formal analysis and writing – review & editing.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

M. M. and P. K. gratefully acknowledge financial support from the Deutsche Forschungsgemeinschaft (DFG, project 380626961). We thank Dr Nicolae Atodiresei (Research Center Jülich) for helpful discussions and Dr Natalya Izarova for crystallographic measurements (RWTH Aachen University). We acknowledge NMR, MS and EA measurements performed by the Central Institute for Engineering, Electronics and Analytics (ZEA-3) at Research Centre Jülich.

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