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Selective binding in different adsorption sites of a 2D covalent organic framework†

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This study shows that surface-supported two-dimensional (2D) porous covalent organic frameworks (COFs) can selectively bind different molecules at specific sites *via* different types of interactions. Scanning tunneling microscopy (STM) images collected at the liquid/solid interface reveal the adsorption of 1,2,4-trichlorobenzene (TCB) in the hexagonal pore of a COF-1 template. A well-defined loop boundary formed by a chain of pentagonal and heptagonal pores allowed the investigation of the effect of pore shape and size on TCB adsorption, suggesting that both geometrical and size effects are important in binding TCB. When both C₆₀ and TCB are present at the solution/solid interface, the TCB molecules are selectively trapped in the pore-site, whereas fullerenes are adsorbed on the top-site of COF-1. While the former structure is stabilized by Cl⋯H hydrogen bonds, the latter is controlled by van der Waals interactions. These results suggest that COFs may offer a powerful platform for the recognition and patterning of guest molecules.

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Introduction

Host/guest (H/G) chemistry in two-dimensional (2D) supramolecular networks has been extensively studied as a means to immobilize a variety of guest molecules, with potential applications in separation technology, molecular recognition, sensing, catalysis and nanoscale patterning.^{1,2} In general, H/G networks rely on hierarchical interactions: the interactions binding the host together are stronger than the interactions binding the guest to the host. The host networks are typically sustained by hydrogen bonding,^{3–6} van der Waals (vdW) forces,^{7,8} or metal–ligand interactions,⁹ whereas the adsorption of guest molecules mainly occurs *via* weaker interactions, typically dispersive interactions with the host as well as with the underlying substrate. However, moving away from self-assembled templates may offer some new opportunities for the stabilization of guest molecules. 2D covalent organic frameworks (COFs) have recently been explored as host systems,^{10–12} opening the possibility of using stronger interac-

tions to stabilize molecules into pores. COF-1, a well ordered 2D porous material that can be synthesized through the on-surface polycondensation of benzenediboronic acid (BDDBA),¹³ is a promising candidate for this application.

The mutually specific recognition between the host network and the guest molecules, *i.e.*, a form of molecular recognition, is at the core of host–guest chemistry.^{14,15} Design strategies for selectivity in guest bonding have often relied on tuning the host geometry, such as pore size,^{16,17} shape¹⁸ and chirality.¹⁹ Much less is known about design based on the interaction between a host and a guest.²⁰ This is mainly due to limitations of hierarchical interaction strength imposed by self-assembled host templates. This aspect can be addressed by using a covalently-bonded template, such as COF-1, which enables a route to molecular recognition through a range of different interactions, including hydrogen bonding, halogen bonding or vdW interactions.

Here, we show that a template of COF-1 can host 1,2,4-trichlorobenzene (TCB) guest molecules that are stabilized through X⋯H hydrogen bonding. The adsorption is enabled through a combination of factors, including the host pore size and shape, and the specific stereochemical arrangement of binding sites. By introducing a solution of C₆₀ in TCB onto the COF-1 template, we observe that selective adsorption of C₆₀ on top of the COF-1 template is obtained simultaneously with trapping of TCB within the pores. This work demonstrates a strategy to separate and recognize molecules through different interactions within a covalent porous 2D template.

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Experimental section

Monolayer COF-1 on highly oriented pyrolytic graphite (HOPG) was formed through a modification of procedures described previously, where heptanoic acid was used as a solvent.^{13,21} In the present experiments, 1.8 mg of 1,4-benzenediboronic acid (Tokyo Chemical Industry Co. Ltd) was added to 1.2 ml of 1,2,4-trichlorobenzene (99%, Sigma-Aldrich), and then sonicated for approximately 30 min. This produced a whitish suspension. 15 μl of this BDBA suspension was dropped onto freshly cleaved HOPG (Structure Probe International, grade SPI-2) and placed into a reactor with a volume of ~ 16 ml. 130 μl of deionized water was added to the bottom of the reactor, and a valve to atmosphere was left slightly open to maintain an open system. The entire reactor was placed in an oven preheated to 125 $^{\circ}\text{C}$ and left for 60 min. After the thermal treatment, the reactor was taken out of the oven and allowed to cool down for at least 20 min before the samples were removed.

Following the confirmation of monolayer COF-1 on the HOPG surface using scanning tunneling microscopy (STM), 15 μl of 1,2,4-TCB (99%, Sigma-Aldrich) was applied onto the substrate. Subsequently, the samples were characterized by STM at the solvent/solid interface. To investigate the adsorption of fullerenes, 15 μl of a supernatant solution of C_{60} (99.5%, Sigma-Aldrich) in 1,2,4-TCB was applied onto a prepared COF-1 template.

STM was performed at room temperature at the liquid–solid interface, using a Digital Instruments STM equipped with a Nanoscope IIIa controller. Tips were cut from a $\text{Pt}_{0.8}\text{Ir}_{0.2}$ wire (Nanoscience Instruments). Bias voltages are reported with respect to the STM tip. STM images were calibrated with the COF-1 lattice parameter (1.476 nm) using free WSxM software.²²

Gas-phase density functional theory (DFT) calculations were carried out using Gaussian09.²³ In previous work, we compared the performance of a number of functionals and basis sets for

calculations of self-assembled systems comprising $\text{X}\cdots\text{H}$ and $\text{X}\cdots\text{X}$ interactions,^{24,25} and found that the combination of M06-2X²⁶/LANL2DZ provides an accurate description of energies for halogen-bonded systems. To simulate surface adsorption, all geometries were constrained to remain planar.

Results and discussion

Using TCB as a solvent, the synthesis of COF-1 on HOPG produces a high density of small domains across the surface, as shown in Fig. S1a.† The obtained surface-confined porous network is qualitatively identical to the one formed using heptanoic acid as a solvent.¹⁰ In both cases, large COF sheets appear to comprise multiple domains that have grown together, leaving disordered regions evident at the domain boundaries. However, when using TCB, we also observed ordered domain boundaries, as described below.

As shown in Fig. 1a, the COF-1 polycrystalline film synthesized in TCB is composed of two different domain orientations tilted at an angle $\alpha = 30 \pm 1^{\circ}$ to each other and separated by a grain boundary (GB). In our analysis of the epitaxial relationship of COF-1 and HOPG (Fig. 1b), we attribute the dominant domain as phase I, *i.e.*, the COF-1 lattice aligned with HOPG with a 6×6 epitaxial unit cell, which corresponds to a lattice parameter of $a_1 = 1.476$ nm, as reported in previous work.¹⁰ The lattice of phase I is represented by small blue hexagons in Fig. 1b. The smaller domain is denoted as phase II. The orientation of this domain, along the HOPG armchair direction, is nearly commensurate to a 10×10 superstructure (corresponding to 3×3 unit cells). We previously calculated that the phase II epitaxial orientation of COF-1 should be energetically disfavoured,¹⁰ consistent with our infrequent experimental observation of the phase. Phase II has also occasionally been observed for COF-1 synthesized from heptanoic acid. (See Fig. S1†).

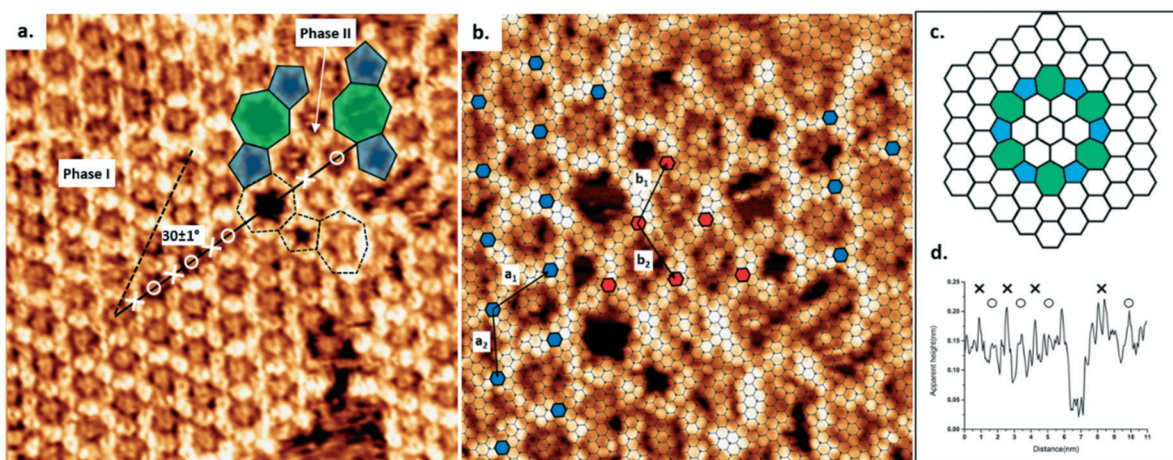


Fig. 1 (a) COF-1 STM image showing the grain boundary consisting of pentagon and heptagon rings, connecting two domains rotated by $30 \pm 1^{\circ}$ with respect to one another, as shown by the dashed and solid black lines. The image was collected at the interface of TCB and HOPG. Image width: 15 nm. Scanning conditions: $V = -1000$ mV, $I = 100$ pA. (b) The epitaxy of COF-1 and HOPG at the grain boundary region. Phase I: $a_1 = a_2 = 1.476$ nm; phase II: $b_1 = b_2 = 1.42$ nm. Image width: 9.6 nm. (c) Schematic structure of the rotational grain boundary identified in (b), *i.e.*, a loop defect with C_6 symmetry. Five- and sevenfold rings are shaded by blue and green, respectively. (d) Line profile of the measured height along the black solid line in (a). The positions of benzene rings in COF-1 are marked as \times and the positions of adsorbed TCB molecules are marked as \circ .



The GB region in Fig. 1 consists of elementary topological defects, pentagons (shaded blue) and heptagons (shaded green). These defects represent some of the basic building blocks of the disordered COF-1 network that have been observed on oriented noble metal surfaces under ultra-high vacuum conditions (UHV),^{27,28} where the random distribution of these topological defects can be attributed to kinetic trapping during synthesis, since the water-driven self-correction mechanism is not available under UHV conditions. However, the ordered pentagonal/heptagonal defects that we observe in this work are directly analogous to defects formed in another single layer covalent system, *i.e.*, single layer graphene.^{29–31} According to the atomic model proposed by Cockayne *et al.*,³² the chain of alternating pentagon and heptagon defects is consistent with a loop GB of graphene, as shown in Fig. 1c. Compared with the linear GB, reported by Xu *et al.* in COFs formed by benzene-1,3,5-tricarbaldehyde (BTA) and *p*-phenylenediamine (PDA),³³ the loop defect boundary has the lowest energy per dislocation core, suggesting that this defect is likely to form under conditions where mobile dislocations exist. This is consistent with our COF-1 synthesis method, since defect correction within the COF lattice leads to improved structural order under humid synthesis conditions.¹³ In our case this translates to the minimization of the energy at domain boundaries through the inclusion of well-defined Stone–Wales type defects.²⁹

The presence of the loop boundary defect enables the identification of the adsorption of TCB in the COF-1 pore. Fig. 1d shows the apparent height profile along the zigzag direction of the phase I COF-1 lattice (the black solid line in Fig. 1a). The adsorption of TCB in hexagonal rings can be clearly distinguished by comparison with the apparent height of the heptagonal ring.

High resolution STM images allow the elucidation of the host–guest structure formed by the hexagonal pore of COF-1 and a TCB molecule, as shown in Fig. 2a. The six phenyl rings in the backbone of COF-1 can be easily distinguished, as are submolecular features associated with the adsorbed TCB molecule. These features are consistent with the benzene ring and the chlorine atoms, and their presence suggests the stable adsorption of the TCB molecule (*i.e.* that the molecule is not rotating).³⁴ Our gas-phase DFT calculations suggest that the TCB adsorption is stabilized through Cl⋯H hydrogen bonding interactions, as indicated in Fig. 2b; consistent with previous work, we identify these interactions through Cl⋯H distances that are shorter than the sum of the Cl and H vdW radii, as shown in Table S1 in the ESI.^{24,35†} The electron density associated with the chlorine atoms in the TCB molecule is anisotropically distributed, as shown in Fig. S2.†³⁶ The nucleophilic belt orthogonal to the covalent bond, with higher electron density and showing negative electrostatic potential, forms X⋯H hydrogen bonds with neighboring hydrogen atoms on the COF-1 backbone (dashed blue line).³⁷ Although the σ -hole of the chlorine atoms points toward the nucleophilic oxygen atom of the COF-1 (dotted black line in Fig. 2b), the interaction between the chlorine and the oxygen is screened by

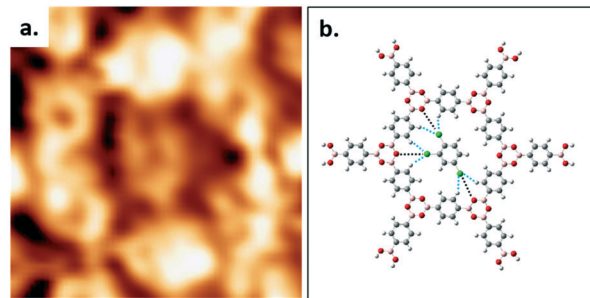


Fig. 2 (a) The detailed STM image of one TCB molecule adsorbing into the hexagonal ring formed by COF-1. Image width: 2.1 nm. Scanning conditions: $V = -1000$ mV, $I = 100$ pA. (b) DFT-calculated structure of one TCB molecule in the hexagonal pore (M06-2X/LANL2DZ). Dashed blue and black lines represent the interactions of Cl⋯H and Cl⋯O, respectively.

the hydrogens on the phenyl rings, as shown in Fig. S2.† Our DFT calculation confirms that the chlorine–oxygen distance is too large (~ 4.38 Å) to allow a significant interaction.

The presence of the pentagonal/heptagonal defects at the GB allows the exploration of the topological and pore size dependence of the guest molecule adsorption. In previous work, fullerene guest molecules were shown to preferentially adsorb in the irregularly-shaped pores present at domain boundaries in a nanoporous hydrogen-bonded network of trimesic acid (TMA).³⁸ However, in contrast to the TMA network where the domain boundary dynamically evolves,³⁹ the loop defects in COF-1 are stable during STM scanning and present a well-defined molecular template, as shown in Fig. 3. The diameters of pentagonal rings (blue dashed line) and heptagonal rings (green dashed line) are 1.381 nm and 1.959 nm respectively, compared to the hexagonal pore size (red dashed line) of 1.758 nm.²⁷ Pores of different sizes can adsorb different numbers of guest molecules, as shown in Fig. 3a–c. There is no adsorption of TCB in pentagonal rings; basic geometric considerations reveal that a TCB molecule cannot be included in this pore without creating interference between hydrogen atoms on the TCB and the pentagonal ring (multiple H⋯H contacts with distances smaller than twice the hydrogen vdW radius), as shown in Fig. S3.†

Two TCB molecules can be adsorbed simultaneously in the heptagonal pore, as shown in Fig. 3. The adsorption of a different number of guest molecules within different pore

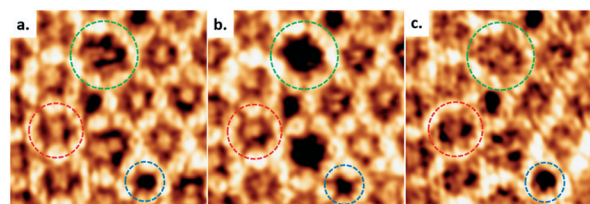


Fig. 3 (a–c) The sequential collection of STM images showing the evolution of adsorption of TCB in the grain boundary defects. Different geometries are marked by different color dashed circles: heptagonal ring (green circle); hexagonal ring (red circle); and pentagonal ring (blue circle). Image width: 5.8 nm. Scanning conditions: $V = -1000$ mV, $I = 100$ pA.



sizes is driven by the commensurability of host and guest molecules. Networks with sufficiently large cavities can host clusters of molecules.^{4,9,40} The adsorption of TCB in porous self-assembled networks has been reported previously for large-pore self-assembled templates,^{17,41} including instances where TCB was trapped in close-packed domains within the pore.^{41,42} In the present study, DFT calculations suggest that both inter-TCB Cl⋯Cl interactions and TCB-COF Cl⋯H interactions stabilize the adsorption of the TCB pair within the heptagonal pore (see the ESI†).

The occupation states of the pentagonal and hexagonal pores appear unchanged with time under STM scanning: the pentagonal pores remained empty, whereas the hexagonal pores retain a single TCB molecule. Conversely, the adsorption of TCB molecules in heptagonal rings exhibits dynamical change. The adsorption-desorption-readsorption process of two TCB molecules can be clearly distinguished in the STM images, as marked by green circles in Fig. 3a-c. Although both empty pores and double-occupied pores were observed, we did not detect the intermediate state, *i.e.*, single TCB adsorbed in a heptagonal ring. The most likely explanation for this is the relatively weak stabilization for an individual TCB molecule in a heptagonal pore. Unlike the matched symmetry between the TCB and the hexagonal pore, the reduced symmetry of the heptagonal pore is a poor match for the TCB molecule, reducing the number of possible Cl⋯H contacts for a single molecule, and rendering single-molecule adsorption unfavourable.

Previous work showed that C₆₀ introduced from a solution containing heptanoic acid as a solvent is adsorbed on the COF-1 template in two different sites, the top-site and pore-site.¹⁰ The top-site geometry corresponds to the adsorption of a fullerene molecule on the boroxine ring and the pore-site corresponds to the adsorption of a fullerene molecule in the hexagonal pore. Our present experiments reveal that C₆₀ introduced from a solution in TCB exhibits a different behaviour, as shown in Fig. 4a. Our interpretation of the STM data is shown in Fig. 4b, which is consistent with the results obtained in the absence of C₆₀. Therefore, we attribute the tightly-packed hexagonal pattern to TCB molecules adsorbed in the COF-1 pores (indicated as shaded blue circles in Fig. 4a and b), and the high-contrast, larger-periodicity hexagonal domain to C₆₀ molecules adsorbed on top-sites. The top-site adsorption of C₆₀ is consistent with the calculated geometry, where the architecture is stabilized by vdW interactions.¹⁰

The growing body of literature on H/G architectures based on COF-1 suggests that the solvent used for C₆₀ deposition plays an important role in the adsorption of the guest molecules. Past work shows that C₆₀ molecules grow layer-by-layer sequentially from a COF-1 template dispersed in 1-phenyloctane, suggesting that the solvent does not compete with C₆₀ for adsorption in the pore site.¹¹ Using heptanoic acid as a solvent, we anecdotally observed a preference for top-site adsorption of fullerenes, suggesting that heptanoic acid may compete for adsorption in the pore-site. Combined with the present results, this suggests that the solvent can be used to select the initial adsorption site for fullerenes, that is TCB and 1-phenyloctane will favor top-site

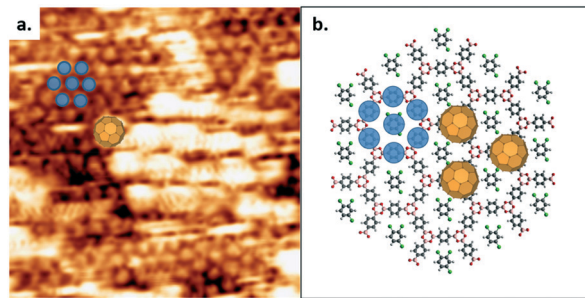


Fig. 4 (a) The top-site selective formation in the C₆₀/COF-1 system. Image width: 12 nm. Scanning conditions: $V = -800$ mV, $I = 100$ pA. (b) Schematic image of top-site selective formation in the C₆₀/COF-1 system. The shaded blue circles over benzene rings represent the closed packed background in STM image. Top-site C₆₀ molecules are placed over boroxine rings, corresponding to the large bright spots in (a).

and pore-site adsorption, respectively, whereas heptanoic acid cannot predictably favor either adsorption site.

Conclusions and perspectives

A 2D porous COF-1 layer was synthesized on HOPG from solution in TCB. Two different epitaxial orientations of COF-1 were observed, with a well-defined loop boundary defect formed by a chain of pentagonal rings and heptagonal rings separating the two domain orientations. STM images collected at the TCB/solid interface reveal the adsorption of TCB within the hexagonal pores of the COF-1 template. Our DFT calculations show that this host/guest structure is stabilized by Cl⋯H hydrogen bonding. The presence of the loop boundary defect permitted the investigation of the effect of pore shape and size on TCB adsorption: no TCB was adsorbed in the pentagonal pores, but the larger heptagonal pores accommodated two TCB molecules, stabilized through a combination of Cl⋯H and Cl⋯Cl bonding. Including C₆₀ in the TCB solution allowed the investigation of the molecular recognition of the template. When both C₆₀ and TCB are present at the solution/solid interface, TCB molecules are selectively trapped in the pores of the COF-1 template, whereas fullerenes are adsorbed on top of the COF-1 in sites identified as top sites. The selective adsorption of TCB and C₆₀ in COF-1 suggests that the different adsorption sites in the COF-1 lattice can be used to separate guest molecules by different interactions, opening opportunities for applications in molecular patterning and recognition.

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References

- J. Teyssandier, S. De Feyter and K. S. Mali, *Chem. Commun.*, 2016, **52**, 11465–11487.
- F. Cicoira, C. Santato and F. Rosei, in *Stm and Afm Studies on (bio) molecular systems: Unravelling the nanoworld*, Springer, 2008, pp. 203–267.
- S. J. Griessl, M. Lackinger, F. Jamitzky, T. Markert, M. Hietschold and W. M. Heckl, *J. Phys. Chem. B*, 2004, **108**, 11556–11560.
- J. A. Theobald, N. S. Oxtoby, M. A. Phillips, N. R. Champness and P. H. Beton, *Nature*, 2003, **424**, 1029–1031.
- J. M. MacLeod, O. Ivasenko, C. Fu, T. Taerum, F. Rosei and D. F. Perepichka, *J. Am. Chem. Soc.*, 2009, **131**, 16844–16850.
- O. Ivasenko, J. M. MacLeod, K. Y. Chernichenko, E. S. Balenkova, R. V. Shpanchenko, V. G. Nenajdenko, F. Rosei and D. F. Perepichka, *Chem. Commun.*, 2009, 1192–1194.
- E. Mena-Osteritz and P. Bäuerle, *Adv. Mater.*, 2006, **18**, 447–451.
- S. Lei, K. Tahara, X. Feng, S. Furukawa, F. C. De Schryver, K. Müllen, Y. Tobe and S. De Feyter, *J. Am. Chem. Soc.*, 2008, **130**, 7119–7129.
- S. Stepanow, M. Lingenfelder, A. Dmitriev, H. Spillmann, E. Delvigne, N. Lin, X. Deng, C. Cai, J. V. Barth and K. Kern, *Nat. Mater.*, 2004, **3**, 229–233.
- D. Cui, J. MacLeod, M. Ebrahimi, D. Perepichka and F. Rosei, *Chem. Commun.*, 2015, **51**, 16510–16513.
- J. Plas, O. Ivasenko, N. Martsinovich, M. Lackinger and S. De Feyter, *Chem. Commun.*, 2016, **52**, 68–71.
- J. Sun, X. Zhou and S. Lei, *Chem. Commun.*, 2016, **52**, 8691–8694.
- J. R. F. Dienstmaier, D. D. Medina, M. Dogru, P. Knochel, T. Bein, W. M. Heckl and M. Lackinger, *ACS Nano*, 2012, **6**, 7234–7242.
- D. Bonifazi, S. Mohnani and A. Llanes-Pallas, *Chem – Eur. J.*, 2009, **15**, 7004–7025.
- M. Li, Q. Zeng and C. Wang, *Sci. China: Phys., Mech. Astron.*, 2011, **54**, 1739–1748.
- M. Li, K. Deng, S. B. Lei, Y. L. Yang, T. S. Wang, Y. T. Shen, C. R. Wang, Q. D. Zeng and C. Wang, *Angew. Chem.*, 2008, **120**, 6819–6823.
- K. Tahara, K. Nakatani, K. Iritani, S. De Feyter and Y. Tobe, *ACS Nano*, 2016, **10**, 2113–2120.
- J. Adisoejoso, K. Tahara, S. Okuhata, S. Lei, Y. Tobe and S. De Feyter, *Angew. Chem.*, 2009, **121**, 7489–7493.
- E. Ghijssens, H. Cao, A. Noguchi, O. Ivasenko, Y. Fang, K. Tahara, Y. Tobe and S. De Feyter, *Chem. Commun.*, 2015, **51**, 4766–4769.
- K. Tahara, K. Katayama, M. O. Blunt, K. Iritani, S. De Feyter and Y. Tobe, *ACS Nano*, 2014, **8**(8), 8683–8694.
- J. R. F. Dienstmaier, A. M. Gigler, A. J. Goetz, P. Knochel, T. Bein, A. Lyapin, S. Reichlmaier, W. M. Heckl and M. Lackinger, *ACS Nano*, 2011, **5**, 9737–9745.
- I. Horcas, R. Fernández, J. Gomez-Rodriguez, J. Colchero, J. Gómez-Herrero and A. Baro, *Rev. Sci. Instrum.*, 2007, **78**, 013705.
- Gaussian 09, Revision A.02*, M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery, J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, O. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski and D. J. Fox, Gaussian, Inc., Wallingford CT, 2016.
- R. Gatti, J. M. MacLeod, J. A. Lipton-Duffin, A. G. Moiseev, D. F. Perepichka and F. Rosei, *J. Phys. Chem. C*, 2014, **118**, 25505–25516.
- R. Gutzler, C. Fu, A. Dadvand, Y. Hua, J. M. MacLeod, F. Rosei and D. F. Perepichka, *Nanoscale*, 2012, **4**, 5965–5971.
- Y. Zhao and D. G. Truhlar, *Theor. Chem. Acc.*, 2008, **120**, 215–241.
- O. Ourdjini, R. Pawlak, M. Abel, S. Clair, L. Chen, N. Bergeon, M. Sassi, V. Oison, J.-M. Debierre and R. Coratger, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2011, **84**, 125421.
- N. A. Zwaneveld, R. Pawlak, M. Abel, D. Catalin, D. Gimes, D. Bertin and L. Porte, *J. Am. Chem. Soc.*, 2008, **130**, 6678–6679.
- J. Ma, D. Alfe, A. Michaelides and E. Wang, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2009, **80**, 033407.
- S. Kurasch, J. Kotakoski, O. Lehtinen, V. Skákalová, J. Smet, C. E. Krill III, A. V. Krashenninnikov and U. Kaiser, *Nano Lett.*, 2012, **12**, 3168–3173.
- J. MacLeod and F. Rosei, *Small*, 2014, **10**, 1038–1049.
- E. Cockayne, G. M. Rutter, N. P. Guisinger, J. N. Crain, P. N. First and J. A. Stroscio, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2011, **83**, 195425.
- L. Xu, X. Zhou, W. Q. Tian, T. Gao, Y. F. Zhang, S. Lei and Z. F. Liu, *Angew. Chem., Int. Ed.*, 2014, **53**, 9564–9568.
- S. J. Griessl, M. Lackinger, F. Jamitzky, T. Markert, M. Hietschold and W. M. Heckl, *Langmuir*, 2004, **20**, 9403–9407.
- R. Gutzler, O. Ivasenko, C. Fu, J. L. Brusso, F. Rosei and D. F. Perepichka, *Chem. Commun.*, 2011, **47**, 9453–9455.
- I. S. Gutiérrez, F.-Y. Lin, K. Vanommeslaeghe, J. A. Lemkul, K. A. Armacost, C. L. Brooks and A. D. MacKerell, *Med. Chem.*, 2016, **24**, 4812–4825.



- 37 G. Cavallo, P. Metrangolo, R. Milani, T. Pilati, A. Priimagi, G. Resnati and G. Terraneo, *Chem. Rev.*, 2016, **116**, 2478–2601.
- 38 D. den Boer, G. D. Han and T. M. Swager, *Langmuir*, 2014, **30**, 762–767.
- 39 M. Lackinger, S. Griessl, L. Kampschulte, F. Jamitzky and W. M. Heckl, *Small*, 2005, **1**, 532–539.
- 40 M. O. Blunt, J. C. Russell, N. R. Champness and P. H. Beton, *Chem. Commun.*, 2010, **46**, 7157–7159.
- 41 M. O. Blunt, J. Adisoejoso, K. Tahara, K. Katayama, M. Van der Auweraer, Y. Tobe and S. De Feyter, *J. Am. Chem. Soc.*, 2013, **135**, 12068–12075.
- 42 R. Gutzler, S. Lappe, K. Mahata, M. Schmittel, W. M. Heckl and M. Lackinger, *Chem. Commun.*, 2009, 680–682.

