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## 2D self-assembly of phenylene-vinylene tectons at the liquid-HOPG interface: from the concentration and alkyl chains length effects to host-guest systems

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**Abstract.** Here we report the synthesis and characterization of a series of new phenylenevinylene tectons. The study by Scanning Tunneling Microscopy (STM) of their supramolecular self-assembly at the interface between a phenyloctane solution and Highly Oriented Pyrolytic Graphite (HOPG) demonstrates that variation of concentration and alkyl chains length led to the formation of different networks, a compact one and a nanoporous one, with a fine control of the lattice parameters. The study of host-guest properties of the nanoporous network revealed a selectivity toward guest compounds according to their shape and size.

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#### 1. Introduction

Surface noncovalent nanostructuration by supramolecular self-assembly of organic building blocks (so-called "tectons") has attracted much interest for several years [1], in view of new potential applications in nanoscience such as molecular electronics [2], photonics or nanomechanical devices [3]. In particular, the formation of a two-dimensional (2D) self-assembled monolayer at the interface between an atomically-flat substrate and a solution of molecular building blocks offers a lot of flexibility and ease of fabrication, together with the possibility of matter exchanges with the liquid environment when working on such surface-confined nanomaterials [4]. In addition to the control of molecule positioning leading to complex nanopatterns, the control of surface properties has also been addressed more recently [5]. In that respect, nanoporous networks with specific properties and presenting selective adsorption and/or surface diffusion of guest molecules were developed [6]. Owing to the host properties of these networks towards organic guests, they allowed for addressing various interesting aspects such as molecular dynamics, selection, recognition, and immobilization, hence leading to functional surfaces [7]. Moreover, these 2D surface-confined nanoporous molecular networks systems can thereby act as templates for the construction of multicomponent 2D hierarchical assemblies [8], most of them giving an isotropic network as obtained from trifunctional molecular building blocks. Among them, we previously developed a strategy to form "on demand" non-covalent self-assemblies with predetermined 2D topologies on sp2-hybridized carbon-based substrates at the liquid-solid interface at room temperature. This method dealt with 2D self-assembly of building blocks designed so as to enable intermolecular interdigitation of alkyl side-chains thus forming noncovalent bonding units (so-called "molecular clips") [9]. Following this principle, we first designed a family of trifunctional molecules based on a symmetrical tristilbene core substituted by three molecular clips leading consequently to the formation of isotropic honeycomb-like nanoporous monolayers. We demonstrated its interest as a molecular sieve [10] and determined the thermodynamic conditions of stability of such a nanoporous phase [11,12].

More recently, taking advantage of this novel functional unit, we obtained 2D patterns of well-defined supramolecular polymer-like chains, formed by a bifunctional tecton, which can self-organize into anisotropic nanoporous networks [13]. However, the precise study and description of this latter tecton self-assembly on HOPG was not investigated so far. In particular, its ability to trap or not guest molecules and the influence of parameters such as concentration on the experimental procedure has not been reported yet.

Here, we first report the synthesis of a series of tectons **BS-C**<sub>n</sub> varying in alkyl chains length (figure 1, n = 10, 12 and 14) (**BS-C**<sub>n</sub>). We then describe the influence of the concentration on the self-assembly of the tecton **BS-C**<sub>10</sub>, in order to find the optimal experimental parameters to favor the formation of an anisotropic nanoporous network. We demonstrate that by varying the alkyl chain length it is possible to finely tune the anisotropic network lattice parameters. Finally, preliminary results regarding the ability of this nanoporous network to host third-party molecules are presented.

#### 2. Results and discussion

#### 2.1. Synthesis of "bone-shape" tectons $BS-C_n$

All the **BS-C**<sub>n</sub> tectons were prepared as shown in figure 1, by adapting a procedure described in the literature [15]. First, compounds **Clip-C**<sub>n</sub> were synthesized with various alkyl chain length (n = 10, 12, and 14). Then, two equivalents of **Clip-C**<sub>n</sub> underwent a Wittig-Horner reaction with one equivalent of 2,5-dimethylterephthalaldehyde (dialdehyde on figure 1) obtained itself from the commercially available 1,4-bis(chloromethyl)-2,5-dimethylbenzene.



Figure 1. Synthesis of the BS-C<sub>n</sub> compounds, differing from the alkyl side chains -C<sub>n</sub>H<sub>2n+1</sub>.

The structures of target tectons  $BS-C_n$  were confirmed by proton and carbon NMR spectroscopies, elemental analysis, and mass spectrometry (see appendix A).

#### 2.2. Photophysical properties in solution of tectons $BS-C_n$

Absorption and emission spectra in solution of compounds **Clip-C**<sub>n</sub>, **triPPV**, and **BS-C**<sub>n</sub> have been recorded. In absorption, all the spectra of tectons **BS-C**<sub>n</sub> have the same behavior and a typical UV-visible spectrum is reported in figure 2 (compound **BS-C**<sub>10</sub>). It is almost the superposition of the spectra of the corresponding **Clip-C**<sub>10</sub> ( $\lambda_{max} = 308$  nm) and the central stilbenes model compound **TriPPV** ( $\lambda_{max} = 353$  nm) (figure 2).



Figure 2. Absorption spectra of BS-C<sub>10</sub>, Clip-C<sub>10</sub> and TriPPV compounds.

In addition, molar extinction coefficients ( $\epsilon$ ) have been determined by varying the solution concentrations (table 1).

Compound	Molar extinction coefficient at 324 nm
$BS-C_{10}$	111545 L.mol <sup>-1</sup> .cm <sup>-1</sup>
Clip-C <sub>10</sub>	39284 L.mol <sup>-1</sup> .cm <sup>-1</sup>
TriPPV	26792 L.mol <sup>-1</sup> .cm <sup>-1</sup>

Table 1. Molar extinction coefficients (ε) of BS-C<sub>10</sub>, Clip-C<sub>10</sub> and TriPPV compounds.

The sum of **TriPPV** molar extinction coefficient and twice that of the **Clip-C**<sub>10</sub> gives  $\varepsilon = 105360 \text{ L.mol}^{-1} \cdot \text{cm}^{-1}$ , a value very close to the **BS-C**<sub>10</sub> molar extinction coefficient:  $\varepsilon = 111545 \text{ L.mol}^{-1} \cdot \text{cm}^{-1}$ . The difference between these two values can be explained by an increased conjugation in the **BS-C**<sub>10</sub> compound, compared to **Clip-C**<sub>10</sub> and **TriPPV** compounds. This result proves that **BS-C**<sub>10</sub> absorption is due to the absorption of both **Clip-C**<sub>10</sub> units and the central core.

Similar results have been obtained for **BS-C**<sub>12</sub> and **BS-C**<sub>14</sub> molecules demonstrating that there is no influence of the side alkyl chains length on the spectral shape. Only the molar extinction coefficient at 324 nm increases with the side alkyl chain length, to reach around  $\varepsilon = 123918$  L.mol<sup>-1</sup>.cm<sup>-1</sup> and  $\varepsilon = 135064$  L.mol<sup>-1</sup>.cm<sup>-1</sup> respectively for **BS-C**<sub>12</sub> and **BS-C**<sub>14</sub>. Such small increase with longer alkyl chains could be due to the increased mass of the alkyl chains, which could dump the skeleton fluctuations a little more efficiently.

Emission spectra have also been recorded, by exciting the compounds at their absorption maximum, e.g. 324 nm for **BS-C**<sub>n</sub> molecule, 308 nm for corresponding **Clip-C**<sub>n</sub>, and 353 nm for **TriPPV** molecule. All the spectra are very similar, whatever the side alkyl chain length. As an example with **BS-C**<sub>10</sub> (figure 3), emission maxima at 419 nm and 442 nm are observed. This structure is typical of central **TriPPV** emission, even though the spectra are switched toward higher wavelengths, due to an increased conjugation.



Figure 3. Emission spectra for BS-C<sub>10</sub>, Clip-C<sub>10</sub> and TriPPV compounds.

Excitation spectra have also been recorded for each compound at their emission maxima: in all cases, they were similar to the absorption spectra of the corresponding molecules. Therefore, in each case, there is only one specie at the excited state. To conclude, side alkyl chains variation does not affect photophysical properties of **BS-C**<sub>n</sub> compounds.

#### 2.3. Self-assembly properties of $BS-C_{10}$ at the liquid-solid interface: effect of the concentration

First, the self-assembly properties of compound  $BS-C_{10}$  were studied by STM at the solution-solid interface at different concentrations. The solutions of  $BS-C_{10}$  were deposited just after cleaving the substrate and were then imaged by STM at room temperature. Through this procedure, the growth is controlled by the kinetics rather than the thermodynamics and it thus leads to the formation of various types of networks and not only that which is thermodynamically the most stable. This coexistence is explained by an initial nucleation-growth phenomenon, followed by a kinetic blockade [12]. The

hypothesis is that during nucleation, molecules nucleate on the HOPG surface according to the "molecular clip" process and it is at this moment that the configuration is determined. Actually, for  $3.10^{-4}$  mol.L<sup>-1</sup> solutions, many different networks were obtained with surface densities ranging from 0.13 molecule/nm<sup>2</sup> (nanoporous network) to 0.21 molecule/nm<sup>2</sup> (most compact network). For a concentration reduced to  $1.10^{-4}$  mol.L<sup>-1</sup> solutions the majority network is the nanoporous one (figure 4 (a) and (b)).



Figure 4. STM images and molecular model of adsorption on HOPG at the HOPG-phenyloctane interface. a),b) STM images of BS-C<sub>10</sub> self-assembly for a deposition at a) 3.10<sup>-4</sup> mol.L<sup>-1</sup> (Circle: the most compact network, Rectangle: nanoporous network) (V = -930 mV, I = 40 pA, 100x100 nm<sup>2</sup>), b) 1.10<sup>-4</sup> mol.L<sup>-1</sup> (nanoporous network only) (V = -870 mV, I = 30 pA, 20x20 nm<sup>2</sup>), c) Molecular models of one unit cell of the monolayer adsorbed on HOPG for the nanoporous network.

On the STM images the conjugated moieties appear much brighter than the aliphatic parts. The lattice parameters of the nanoporous network shown on figure 4 are a = 3.9 nm, b = 2.1 nm and  $\alpha = 65^{\circ}$  and the density is 7.4 nm<sup>2</sup> per molecule, which is higher than the density of the compact network (5.6 nm<sup>2</sup> per molecule). Moreover, previous work has led to a reproducible procedure leading to the nanoporous network [12], therefore we will focus on the study of this network only. Similar concentration-dependent self-assemblies have been reported in literature [14], and an expression of the critical concentration and a procedure leading to the thermodynamically stable lattice have been derived for the self-assembly of trifunctional tristilbenes (TSB) [12].

#### 2.4. Self-assembly properties of BS-C<sub>n</sub> at the liquid-solid interface: influence of alkyl chain length

In the case of the nanoporous network, the exclusive formation of which is well-controlled [12], we investigated the possibility to finely tune the distance between cavities, as it constitutes a prerequisite toward the control and exploitation of such anisotropic network. To achieve that, first we developed and synthesized several building blocks with various peripheral alkyl chain lengths, more precisely bifunctionnal "bone-shape tectons" **BS-C**<sub>10</sub>, **BS-C**<sub>12</sub> and **BS-C**<sub>14</sub> which are presented on figure 5. Second, we managed to control and realize a series of nanoporous networks by self-assembly of these tectons on HOPG.



Figure 5. STM images and molecular model of adsorption on HOPG at the HOPG-phenyloctane interface. a) Large-scale STM image of a monolayer domain of compound BS-C<sub>10</sub> (V = -650 mV, I = 16 pA, 255x255 nm<sup>2</sup>) ; b) Close-up STM image of BS-C<sub>10</sub> self-assembly (V = -870 mV, I = 30 pA, 20x20 nm<sup>2</sup>), c) Molecular model associated to compound BS-C<sub>10</sub>, d) Large-scale STM image of a monolayer domain of compound BS-C<sub>12</sub> (V = -1150 mV, I = 11 pA, 70x70 nm<sup>2</sup>) ; e) Close-up STM image of BS-C<sub>12</sub> self-assembly (V = -1150 mV, I = 12 pA, 22x22 nm<sup>2</sup>), f) Molecular model associated to compound BS-C<sub>12</sub>, g) Large-scale STM image of a monolayer domain of compound BS-C<sub>14</sub> (V = -650 mV, I = 11 pA, 80x80 nm<sup>2</sup>) ; h) Close-up STM image of BS-C<sub>14</sub> self-assembly (V = -680 mV, I = 12 pA, 15x15 nm<sup>2</sup>), i) Molecular model associated to compound BS-C<sub>14</sub>.

In these conditions, the self-assemblies obtained are stable and self-replicated on distances of at least 100 nm. Moreover, this nanoporous network is obtained with the same self-assembly pattern for each alkyl chain length, as seen on models. Only the lattice parameter "a", *i.e.* the longitudinal distance between cavities, increases with the alkyl chain length, whereas the cavity size remains the same, as evidenced with the lattice parameters reported in table 2. As a consequence, obtaining the nanoporous network remains as easy for longer alkyl side chains, in contrast with TSB35 for which the cavity size increases with alkyl chains which restrains the stability domain for the nanoporous network. Therefore, the "molecular clip" function is effective for alkyl chain length ranging at least from ten to fourteen carbons. Note that, since the vector **a** is nearly parallel to the alkyl chains (i.e. the 110 direction of HOPG), the variation of the angle  $\alpha$  is not significant and the decrease of the angle predicted by the theory is lower than the uncertainty on the measurements.

Alkyl chain length	Reconstruction	Lattice parameter a	Lattice parameter b	Angle α
		Theory (measure)	Theory (measure)	Theory (measure)
10 carbons	$\begin{pmatrix} 0 & 8\\ 17 & -3 \end{pmatrix}$	4.0 nm (3.9)	2.0 nm (2.1)	70° (65)
12 carbons	$\begin{pmatrix} 0 & 8\\ 18 & -3 \end{pmatrix}$	4.2 nm (4.1)	2.0 nm (2.1)	69° (69)
14 carbons	$\begin{pmatrix} 0 & 8\\ 19 & -3 \end{pmatrix}$	4.4 nm (4.3)	2.0 nm (2.1)	68° (70)

 Table 2. Analysis of molecular lattice evolution with increasing alkyl chain lengths from C10H21 to C14H29. The reconstruction with respect to HOPG is given.

In addition, a closer look on some STM images reveals a slight variant in alkyl chain packings. More specifically two configurations can be discerned. The first one can be called "regular" (figure 6 (a)), which is the template generally proposed, and a second one corresponding to a reconstruction  $\begin{pmatrix} 0 & 8\\ 12 + n/2 & +1 \end{pmatrix}$  with respect to HOPG (*n* is the alkyl chain length) is named "gauche" (figure 6 (b)), which is encountered sometimes. It corresponds to an inversion of the clips, that is to a shift by a vector (0 4) of an entire lamella. This shift being parallel to a lattice vector (b = (0 8)) this variant preserves the unit cell area and the surface molecular density. The most visible change is the decrease of the lattice angle  $\alpha$ , *e.g.* from 70° (gauche) to 57° (regular) in the case n = 10.



Figure 6. Molecular model of nanoporous network in both configurations: a) Regular, b) Gauche.

On most of the STM images, the regular configuration is observed,. Experimentally the gauche configuration appears as a defect. More precisely, it appears at the boundaries of domains with translational mismatch accidentally fitting the gauche configuration. At places where such a domain boundary crosses a lamella, a local breaking of the periodicity along the lamella is formed as shown on figure 7.



Figure 7. STM image of compound BS-C<sub>14</sub> with both configurations observed: gauche (left rectangle), regular (right rectangle) and local defect resulting from a domain boundary crossing the lamella (white circle).

#### 2.5. Co-assembly structures of $BS-C_{10}$ and guest molecules on HOPG surface

Nanoporous networks are likely to host third-party compounds in their cavities. We investigated the ability of a **BS-C**<sub>10</sub> self-assembly to trap guest molecules. Three guest candidates have been chosen, benzo[*rst*]pentaphene (BPP), pentacene and hexabenzocoronene (HBC) (figure 8), the dimensions of which fitting the cavity size. For BPP and pentacene, even one hour after their addition, only few successful insertions have been observed by STM (figures 8 (a) and (b)): this could be explained by the fact that they are either too small to be adsorbed inside the cavities or too small again to give a very easy STM signature enabling its differentiation from the host self-assembly (their motion inside the cavity leading to a blurred poorly contrasted STM response). Indeed, the low interaction surface between the molecules and the substrate, which induce a low adsorption enthalpy, and their high entropy in the solution could increase their desorption, compared to bigger molecules. The same deposition protocol has been carried out at various concentrations and led to the same conclusion. Oppositely, HBC remained sequestered inside the network and all the cavities appeared filled after a waiting time of one hour, as seen on figure 8 (c). This can be explained by the fact that the shape and size of the cavities.



Figure 8. a) Molecular structures of the guest molecules BPP, pentacene and HBC, b-d) STM images of the self-assembled monolayer of the matrix formed through self-assembly of BS-C<sub>10</sub> on HOPG, after deposition of b) BPP (V = -970 mV, I = 25 pA, 24x24 nm<sup>2</sup>), c) pentacene (V = -1150 mV, I = 40 pA, 40x40 nm<sup>2</sup>), or d) HBC (V = -1425 mV, I = 25 pA, 79x79 nm<sup>2</sup>). Red circles highlight guest insertion.

At lower concentrations these molecules can still be trapped again in the cavities, but not all of the latter are filled (sub-stoichiometry conditions). Then, guest molecules dynamics appears as a blinking of the corresponding spots in successive images. In most cases the appearance of a spot in one cavity is concomitant with the disappearance of another one in its neighborhood, as illustrated on figure 9. The circle shows a disappearing HBC molecule and the arrow shows uniformly distributed stripes parallel to the fast-scan direction. Those stripes correspond to occupation changes occurring in the course of cavity imaging and evidence that the guest molecules are moving when working in such substoichiometry conditions.



Figure 9. Two successive STM images which show an occupation state change during the imaging (arrow) and the disappearance of a HBC molecule (circle). V = -1000 mV, I = 30 pA, 25x25 nm<sup>2</sup>.

In addition, this behavior was observed whatever the length of the peripheral alkoxy chains, which means that in this case (contrary to our precedent studies with trifunctional TSB entities) the chain length parameter don't lead to a modification of the cavities and thus of the molecular sieve properties: increasing the chain length is just a way we can play with in order to tune in a control way the distance between these cavities.

#### 3. Conclusion

In summary, a series of new phenylene-vinylene tectons substituted with peripheral alkyl chains with various lengths have been synthesized. The supramolecular self-assembly properties at liquid/solid interface have been investigated by STM. Changes in concentration led to the formation of various networks, several compact ones and two nanoporous ones. Fine tuning of the peripheral alkyl chain length led as well to the variation of the distance between cavities on one axis only. This distance varies step-by-step, while preserving the cavity size and the detailed features of the HOPG-induced alkyl chain interdigitation. Finally, the nanoporous network can be used as a host-guest system with selectivity toward guest molecules depending on their sizes and shapes, therefore it could be used as a molecular sieve.

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#### Appendix A

Tetrahydrofurane (THF), Toluene and Dichloromethane (CH<sub>2</sub>Cl<sub>2</sub>) used in synthesis come from a fountain. All other solvents were purchased from commercial suppliers and used without further purification. Chromatographic separations were carried out with a flash chromatographic instrument (Grace, Reveleris) and with silica gel (40  $\mu$ m, Grace). Structural information of materials was obtained by <sup>1</sup>H NMR and <sup>13</sup>C spectroscopy. Spectra were recorded at 200 MHz, proton chemical shifts ( $\delta$ ) are reported in ppm and referenced to the residual solvent signal: CDCl<sub>3</sub> (7.26). <sup>13</sup>C NMR spectra were recorded at 200 MHz and  $\delta$  referenced to the residual solvent signal: CDCl<sub>3</sub> (77.16).

The **BS-C**<sub>n</sub> tectons have been synthesized and purified by adapting a general procedure for the Wittig-Hörner reaction described in the literature [15], using the corresponding "molecular clip" compounds **Clip-C**<sub>n</sub>. To a solution of **Clip-C**<sub>n</sub> (2.2eq) in dry THF under argon atmosphere was added tBuOK (1.5eq). The mixture was stirred for 15mn at 0°C then the dialdehyde (1eq) was added. The reaction was stirred for 2h at 0°C. The resulting mixture was quenched with water and concentrated to remove the THF, then extracted with CH<sub>2</sub>Cl<sub>2</sub> and washed with brine. The combined organic layers were dried over MgSO<sub>4</sub>, filtered and concentrated under vacuum. The residue was purified by flash chromatography (1:1 CH<sub>2</sub>Cl<sub>2</sub>:PE) to yield the pure target material **BS-C**<sub>n</sub>.

# $3,3',3'',3'''-((1E,1'E,1''E,1''E)-(((1E,1'E)-(2,5-dimethyl-1,4-phenylene)bis(ethene-2,1-diyl))bis(benzene-5,3,1-triyl))tetrakis(ethene-2,1-diyl))tetrakis((decyloxy)benzene) (BS-C_{10})$



Yellow solid. Yield: 200mg, 27%.  $\mathbf{R}_f = 0.59$  (1:1 CH<sub>2</sub>Cl<sub>2</sub>:PE). <sup>1</sup>H NMR (**250** MHz, CD<sub>3</sub>Cl)  $\delta$ : 0.87 (t, J = 6.7 Hz, 12H, H<sub>27</sub>), 1.28 (m, 56H, H<sub>20-26</sub>), 1.78 (m, 8H, H<sub>19</sub>), 2.49 (s, 6H, H<sub>28</sub>), 3.98 (t, J = 6.5 Hz, 8H, H<sub>18</sub>), 6.81 (m, 4H, H<sub>15</sub>), 7.10 (d, J = 16.0 Hz, 2H, H<sub>5</sub>), 7.11 (m, 8H, H<sub>10,11</sub>), 7.17 (s, 8H, H<sub>13,17</sub>), 7.27 (t, J = 7.7 Hz, 4H, H<sub>14</sub>), 7.42 (d, J = 16.0 Hz, 2H, H<sub>4</sub>), 7.50 (s, 2H, H<sub>2</sub>), 7.56 (m, 6H, H<sub>7,9</sub>). <sup>13</sup>C NMR (63 MHz, CDCl<sub>3</sub>)  $\delta$ : 14.3 (C<sub>27</sub>), 19.8 (C<sub>28</sub>), 22.8, 26.3, 29.5, 29.6, 29.7, 32.1 (C<sub>19-26</sub>), 68.1 (C<sub>18</sub>), 112.5, 114.2 (C<sub>15,17</sub>), 119.3, 124.3, 126.7, 127.4, 128.7, 129.4, 129.8 (C<sub>4-5,7,9-11,13-14</sub>), 133.8, 135.6, 138.1, 138.5, 138.7 (C<sub>1,3,6,12</sub>), 159.6 (C<sub>16</sub>). UV maximum absorption in CH<sub>3</sub>Cl at 20°C  $\lambda_{abs} = 324$ nm. UV molar absorption coefficient at 324nm  $\varepsilon = 111545$  L.mol<sup>-1</sup>.cm<sup>-1</sup>. Fluorescence spectroscopy emission in CH<sub>3</sub>Cl at 20°C  $\lambda_{em} = 419$ nm, excitation  $\lambda_{exc} = 324$ nm. Anal. calcd for C<sub>104</sub>H<sub>142</sub>O<sub>4</sub>: C, 85.79; H, 9.45; O, 4.76; found: C, 85.69; H, 9.49; O, 4.81.

3,3',3'',3'''-((1*E*,1'*E*,1''*E*,1'''*E*)-(((1*E*,1'*E*)-(2,5-dimethyl-1,4-phenylene)bis(ethene-2,1-diyl))bis(benzene-5,3,1-triyl))tetrakis(ethene-2,1-diyl))tetrakis((dodecyloxy)benzene) (BS-C<sub>12</sub>)



Yellow solid. Yield: 59mg, 18%. **R**<sub>f</sub> = 0.60 (1:1 CH<sub>2</sub>Cl<sub>2</sub>:PE). <sup>1</sup>**H** NMR (200MHz, CDCl<sub>3</sub>)  $\delta$ : 0.87 (t, J = 6.7 Hz, 12H, H<sub>29</sub>), 1.28 (m, 72H, H<sub>20-28</sub>), 1.81 (m, 8H, H<sub>19</sub>), 2.50 (s, 6H, H<sub>30</sub>), 4.00 (t, J = 6.5 Hz, 8H, H<sub>18</sub>), 6.83 (m, 4H, H<sub>15</sub>), 7.11 (d, J = 16.0 Hz, 2H, H<sub>5</sub>), 7.15 (m, 8H, H<sub>10-11</sub>), 7.18 (s, 8H, H<sub>13,17</sub>), 7.28 (t, J = 7.7 Hz, 4H, H<sub>14</sub>), 7.44 (d, J = 16.0 Hz, 2H, H<sub>4</sub>), 7.48 (s, 2H, H<sub>2</sub>), 7.55 (m, 6H, H<sub>79</sub>). <sup>13</sup>C NMR (200MHz, CDCl<sub>3</sub>)  $\delta$ : 14.2 (C<sub>29</sub>), 19.7 (C<sub>30</sub>), 22.7, 26.1, 29.4, 29.5, 29.6, 29.6, 29.7, 29.7, 32.2, 31.9 (C<sub>19-28</sub>), 68.0 (C<sub>18</sub>), 112.4, 114.1 (C<sub>15,17</sub>), 119.2, 123.8, 124.3, 126.7, 127.4, 128.7, 129.4, 129.8 (C<sub>4-5,7,9-11,13-14</sub>), 133.8, 135.6, 138.1, 138.5, 138.7 (C<sub>1,3,6,8,12</sub>), 159.6 (C<sub>16</sub>). UV maximum absorption in CH<sub>3</sub>Cl at 20°C  $\lambda_{abs} = 324$ nm. UV molar absorption coefficient at 324nm  $\varepsilon = 123918$  L.mol<sup>-1</sup>.cm<sup>-1</sup>. Fluorescence spectroscopy emission in CH<sub>3</sub>Cl at 20°C  $\lambda_{em} = 419$ nm, excitation  $\lambda_{exc} = 324$ nm. Anal. calcd for C<sub>104</sub>H<sub>142</sub>O<sub>4</sub>: C, 85.78; H, 9.83; O, 4.39; found: C, 85.63; H, 9.79; O, 4.28.

3,3',3'',3'''-((1*E*,1'*E*,1''*E*,1'''*E*)-(((1*E*,1'*E*)-(2,5-dimethyl-1,4-phenylene)bis(ethene-2,1-diyl))bis(benzene-5,3,1-triyl))tetrakis(ethene-2,1-diyl))tetrakis((tetradecyloxy)benzene) (BS-C<sub>14</sub>)



Yellow solid. Yield: 135mg, 22%. **R**<sub>f</sub> = 0.57 (1:1 CH<sub>2</sub>Cl<sub>2</sub>:PE). <sup>1</sup>**H NMR (200MHz, CDCl<sub>3</sub>) \delta:** 0.88 (t, J = 6.7 Hz, 12H, H<sub>31</sub>), 1.27 (m, 88H, H<sub>20-30</sub>), 1.82 (m, 8H, H<sub>19</sub>), 2.52 (s, 6H, H<sub>32</sub>), 4.02 (t, J = 6.5 Hz, 8H, H<sub>18</sub>), 6.84 (m, 4H, H<sub>15</sub>), 7.11 (d, J = 16.0 Hz, 2H, H<sub>5</sub>), 7.15 (m, 8H, H<sub>10,11</sub>), 7.18 (s, 8H, H<sub>13,17</sub>), 7.28 (t, J = 7.7 Hz, 4H, H<sub>14</sub>), 7.44 (d, J = 16.0 Hz, 2H, H<sub>4</sub>), 7.48 (s, 2H, H<sub>2</sub>), 7.56 (m, 6H, H<sub>79</sub>). <sup>13</sup>C **NMR (200MHz, CDCl<sub>3</sub>) \delta:** 14.2 (C<sub>31</sub>), 19.7 (C<sub>32</sub>), 22.7, 26.1, 29.4, 29.5, 29.6, 29.6, 29.7, 29.7, 32.2, 31.9 (C<sub>19-30</sub>), 68.0 (C<sub>18</sub>), 112.4, 114.1 (C<sub>15,17</sub>), 119.2, 123.8, 124.3, 126.7, 127.4, 128.7, 129.4, 129.8 (C<sub>4,5,7,9,10,11,13,14</sub>), 133.8, 135.6, 138.1, 138.5, 138.7 (C<sub>1,3,6,8,12</sub>), 159.6 (C<sub>16</sub>). **UV maximum absorption in CH<sub>3</sub>Cl at 20°C**  $\lambda_{abs} = 324$ nm. **UV molar absorption coefficient at 324nm**  $\varepsilon = 135064$  L.mol<sup>-1</sup>.cm<sup>-1</sup>. **Fluorescence spectroscopy emission in CH<sub>3</sub>Cl at 20°C**  $\lambda_{em} = 419$ nm, excitation  $\lambda_{exc} = 324$ nm. **Anal. calcd for C<sub>112</sub>H<sub>158</sub>O<sub>4</sub>:** C, 85.77; H, 10.15; O, 4.08; found: C, 85.63; H, 10.19; O, 4.18.

#### Appendix B

UV-visible spectra were recorded with a ThermoFisher evolution UV/Vis spectrophotometer coupled with a thermal controller. Cells were in quartz with two faces (thickness = 1cm).

UV-Visible spectra have been recorded in  $CHCl_3$  at room temperature, for concentrations between  $10^{-6}$  mol.L<sup>-1</sup> and  $10^{-5}$  mol.L<sup>-1</sup>. Molar extinction coefficients have been calculated thanks to Beer-Lambert law: A = 1. $\epsilon$ .c. This law is applicable in the concentration range used, so we can consider that there are neither dimerisations nor oligomerisations of the molecules.

Emission and excitation spectra have been recorded in  $CHCl_3$  at room temperature, for concentrations ranging from  $10^{-6}$  mol.L<sup>-1</sup> to  $10^{-5}$  mol.L<sup>-1</sup>. Emission spectra have been obtained by exciting the compounds at their absorption maximum, e.g. 324 nm for **BS-C**<sub>n</sub> molecules, 308 nm for **Clip-C**<sub>n</sub> molecules, and 353 nm for **TriPPV** molecule.

#### Appendix C

STM images were acquired at room temperature with a homemade digital system. The fast-scan axis was kept perpendicular to the sample slope. All images were obtained in the height mode, i.e. with real-time current regulation. Images acquired simultaneously in both fast scan directions were systematically recorded and compared. All images were corrected for the drift of the instrument, by combining two successive images with downward and upward slow-scan directions, using especially developed image cross-correlation software called Imago. The solvent for the self-assembly was 1-phenyloctane (98%, Aldrich), which is well suited for *in situ* scanning tunneling microscopy (STM) because of its low conductivity, slow volatility and high dielectric rigidity and also avoids the coadsorption often observed with linear alkanes. The substrate was a freshly cleaved HOPG (SPI, grade 2) and the tips were mechanically formed from a 250mm Pt/Ir wire (Pt80Ir20, Goodfellow). The freshly-cleaved sample and tip quality was systematically checked by STM observation of HOPG atomic network prior to molecular deposition. The monolayers were formed by immersing the STM junction in a droplet (ca. 5  $\mu$ L) of a solution immediately after observation of HOPG atomic network. Imaging was then carried out in situ at the liquid-solid interface.

#### References

Bonifazi D, Mohnani S and Llanes-Pallas A 2009 Chem. Eur. J. 15 7004

Bilbao N, Destoop I, De Feyter S and Gonzalez-Rodriguez D 2016 Angew. Chem. Int. Ed. 55 659

Tahara K, Nakatyani K, Iritani K, De Feyter S and Tobe Y 2016 ACSNano 10 2113

Liu C, Zhang W, Zeng Q and Lei S 2016 Chem. Eur. J. 22 6768

[7] Pan G-B, Liu J-M, Zhang H-M, Wan L-J, Zheng Q-Y and Bai C-L 2003 Angew. Chem. Int. Ed. 42 2747

El Garah M, Ciesielski A, Marets N, Bulach V, Hosseini MW and Samorì P 2014 Chem. Commun. 50 12250

Palma C-A, Ciesielsski A, Oner MA, Schaeffer G, Lehn J-M, Barth JV and Samorì P 2015 Chem. Commun. **51** 1797

El Garah M, Marets N, Mauro M, Aliprandi A, Bonacchi S, De Cola L, Ciesielski A, Bulach V, Hosseini MW and Samorì P 2015 *J. Am. Chem. Soc.* **137** 8450

[8] Ciesielski A, Palma C-A, Bonini M and Samorì P 2010 Advanced Materials 22 3506

[9] Bléger D, Kreher D, Mathevet F, Attias A-J, Schull G, Huard A, Douillard L, Fiorini-Debuischert C and Charra F 2007 *Angew. Chem. Int. Ed.* **46** 7404

[10] Schull G, Douillard L, Fiorini-Debuisschert C, Charra F, Mathevet F, Kreher D and Attias A-J 2006 *Advanced Materials* **18** 2954

[11] Bellec A, Arrigoni C, Schull G, Douillard L, Fiorini-Debuisschert C, Mathevet F, Kreher D, Attias A-J and Charra F 2011 *The Journal of Chemical Physics* **134** 124702

[12] Arrigoni C, Schull G, Bléger D, Douillard L, Fiorini-Debuisschert C, Mathevet F, Kreher D, Attias A-J and Charra F 2010 *The Journal of Physical Chemistry Letters* **1** 190

[13] Bléger D, Kreher D, Mathevet F, Attias A-J, Arfaoui I, Metgé G, Douillard L, Fiorini-Debuisschert C and Charra F 2008 *Angewandte Chemie* **120** 8540

[14] Ciesielski A, Szabelski P J, Rżysko W, Cadeddu A, Cook T R, Stang P J, Samorì P 2013 *J. Am. Chem. Soc.* **135** 6942

Lee S-L, Adisoejoso J, Fang Y, Tahara K, Tobe Y, Mali K S and De Feyter S 2015 Nanoscale 7 5344

El Garah M, Dianat A, Cadeddu A, Gutierrez R, Cecchini M, Cook TR, Ciesielski A, Stang PJ, Cuniberti G and Samorì P 2016 *Small* **12** 343

<sup>[1]</sup> Barth J V, Costantini G and Kern K 2005 *Nature* **437** 671

<sup>[2]</sup> Wang Q H and Hersam M C 2009 Nature Chemistry 1 206

<sup>[3]</sup> Urdampilleta M, Klyatskaya S, Cleuziou J-P, Ruben M and Wernsdorfer W 2011 *Nature Materials* **10** 502

<sup>[4]</sup> Mazur U and Hipps K W 2015 Chem. Commun. 51 4737

<sup>[5]</sup> De Feyter S and De Schryver F C 2003 Chemical Society Reviews 32 139

Bellec A, Arrigoni C, Douillard L, Fiorini-Debuisschert C, Mathevet F, Kreher D, Attias A J and Charra F 2014 Nanotechnology **25** 435604

<sup>[6]</sup> Kudernac T, Lei S, Elemans J A A W and De Feyter S 2009 Chemical Society Reviews 38 402

Tahara K, Katayama K, Blunt MO, Iritani K, De Feyter S and Tobe Y 2014 ACSNano 8 8 8683

Lee S-L, Fang Y, Velpula G, Cometto F-P, Lingenfelder M, Mullen K, Mali K-S and De Feyter S 2015 ACSNano 9 11608