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## ARTICLE

# Unveiling the nature of supramolecular crown ether- $C_{60}$ interactions

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A series of exTTF-(crown ether)<sub>2</sub> receptors, designed to host  $C_{60}$ , has been prepared. The size of the crown ether and the nature of the heteroatoms have been systematically changed to fine tune the association constants, which were determined by a number of complementary spectroscopic techniques. Electrochemical measurements and transient absorption spectroscopy assisted in corroborating a charge transfer in the ground state and in the excited state leading to the formation of radical ion pairs featuring lifetimes in the range from 12 to 21 ps. To rationalize the nature of the exTTF-(crown ether)<sub>2</sub>• $C_{60}$  stabilizing interactions, theoretical calculations have been carried out, prompting to a synergetic interplay of donor-acceptor,  $\pi$ - $\pi$ , n- $\pi$  and CH $\cdots\pi$  interactions, which is the basis for the affinity of our novel receptors towards  $C_{60}$ .

## Introduction

The construction of non-covalent electron donor–acceptor (D–A) assemblies is a rational way for the creation of new and sophisticated electroactive materials impossible to obtain by covalent means. As representative examples, a variety of simple models for the study of electron and energy transfer processes such as those found in the photosynthetic centers of plants and bacteria have been prepared.<sup>1,2</sup> To this end, fullerenes have been widely used as electron acceptors given their spherical geometry,<sup>3</sup> small reorganization energy in electron transfer reactions,<sup>4</sup> low reduction potential,<sup>5,6</sup> appreciable absorption cross section throughout a wide range of the solar spectrum,<sup>7</sup> and efficient generation of long-lived charge-separated states following photoexcitation.<sup>8–10</sup> In this regard, their supramolecular chemistry is largely based on the use of fullerene derivatives giving rise to metal-ligand interactions,  $\pi$ - $\pi$  stacking, hydrogen bonding, electrostatic interactions, or mechanical bonds.<sup>11–16</sup> This approach involves, however, saturating at least one of their double bonds altering thus their electronic conjugation. In contrast, complexation of pristine fullerene enables maintaining their singular electronic properties. This is typically achieved by using host molecules endowed with large dispersion forces, namely  $\pi$ - $\pi$  and van der Waals, such as cyclodextrins,<sup>17–19</sup> calixarenes,<sup>20–22</sup> cyclotrimerarylenes,<sup>23–25</sup> porphyrins,<sup>26–30</sup> subphthalocyanines,<sup>31–33</sup> or 9,10-di(1,3-dithiol-2-ylidene)-9,10-dihydroanthracenes (exTTF).<sup>34–37</sup>

Interestingly, although an aza-crown ether decorated with lipophilic fragments was the first system ever reported to complex fullerenes in solution,<sup>38</sup> crown ethers have scarcely been explored as hosts for fullerenes. Indeed, despite the importance of both kind of molecules, whose discoverers were each awarded a Nobel Prize,<sup>39,40</sup>

to the best of our knowledge, a detailed experimental and theoretical study on the supramolecular interactions occurring between them has not been properly addressed thus far and many open questions still remain unanswered. A notable exception includes the work by Mukherjee and co-workers,<sup>41, 42</sup> who observed that the overall stability of the resulting complexes increased as a function of the cavity size of the crown ether. To this end, an interplay between different energy terms, such as solvation effects, electron donor-acceptor interactions, etc., was hypothesized to explain the complexation. Following the latter, Liu et al. evaluated the impact of introducing Se atoms in the crown ethers, which led to a better stabilization.<sup>43</sup> As in the previous example, a relation between the cavity size of the crown ether and the binding constants with  $C_{60}$  was noted. Another remarkable example is the porphyrin designed by D'Souza et al. bearing four benzo[18]crown-6 ethers in the *meso* positions and exhibiting a moderate but tunable affinity towards  $C_{60}$  depending on the presence or the absence of  $K^+$  ions.<sup>44</sup> More recently, our group developed a novel receptor for  $C_{60}$ , which was based on an exTTF derivative appended with two benzo[18]crown-6 ethers, exTTF-(crown ether)<sub>2</sub>. This receptor featured extraordinarily high binding constants ( $K_a$ ) for  $C_{60}$  and  $C_{70}$  with  $\log K_a = 6.7$  and 7.4 in benzonitrile at room temperature, respectively. As a matter of fact, it became the sole example of  $C_{60}$  complexation by a single exTTF molecule.<sup>45</sup>

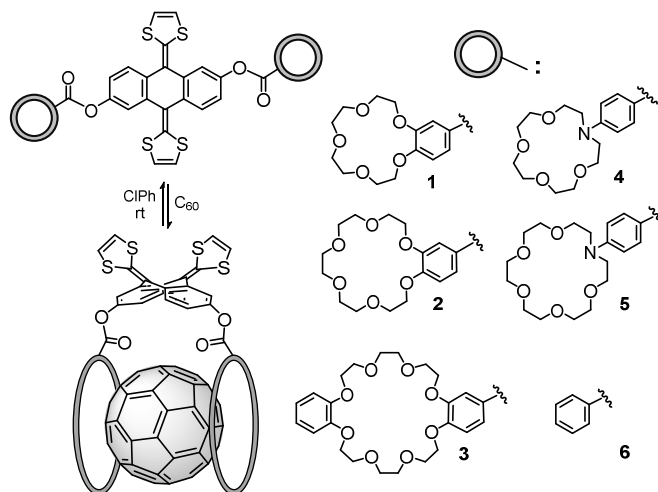
The impact of crown ethers on the receptor properties led us to focus on the study of the crown ether• $C_{60}$  interaction in order to finally unveil its nature. To do so, a series of exTTF-crown ether derivatives has been prepared, in which the cavity size and the nature of the heteroatoms have been systematically modified. Their complexation with  $C_{60}$  has been complementarily investigated by

both spectroscopic and electrochemical means. In addition, theoretical calculations have been carried out to draw conclusions about the key factors influencing the resulting binding constants.

## Results and discussion

In order to carry out this study, we have designed and synthesized a series of novel exTTF-based receptors endowed with two crown ethers **1-5** (Scheme 1). Non-commercially available crown ethers were obtained through a Buchwald-Hartwig cross coupling reaction or a Williamson ether synthesis (see ESI† for further details). Then, 9,10-bis(1,3-dithiol-2-ylidene)-9,10-dihydroanthracene-2,6-diol<sup>46</sup> was esterified with the corresponding crown ether ether appended carboxylic acids either *via* the acyl chloride or activation with EDC. Compound **6** lacking the crown ether moieties was also prepared as reference by condensation of the 2,6-dihydroxylated exTTF with benzoic acid.

Unambiguous characterization of all compounds was carried out employing standard spectroscopic and analytical techniques (see ESI† for details on the synthesis and characterization). Successful esterification was evidenced by the maintenance of the characteristic 1,3-thiol signal at around 6 ppm and the appearance of the crown ether signals in the 3-4 ppm region. UV-vis spectroscopy profile of all exTTFs exhibited the characteristic band at around 435 nm without significant shifts among the derivatives.

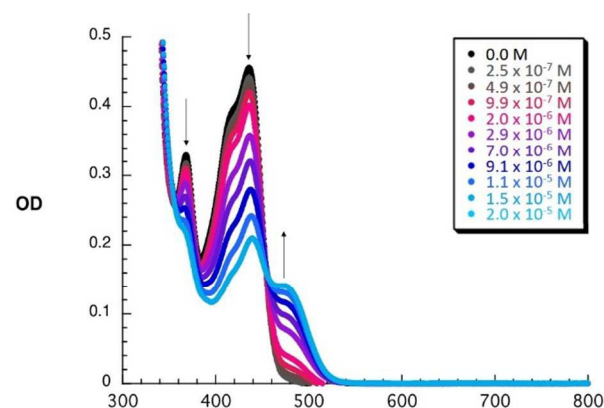


**Scheme 1.** Complexes obtained from exTTF-based **1-6** and  $C_{60}$ .

## Titration Experiments

To shed light onto the ground state interactions between **1-6** and  $C_{60}$ , absorption titrations were performed in PhCl at room temperature (see ESI† for further details). When adding  $C_{60}$  to constant concentrations of **1-6**, an increase of the typical absorption features of  $C_{60}$ , namely a strong absorption band at wavelengths <350 nm, a sharp band at 407 nm, and a broad absorption between 470 and 650 nm evolve. Simultaneously, the intrinsic exTTF features, which are noted between 350 and 450 nm, gradually decrease in the presence of  $C_{60}$ . All of these changes are assigned to a successful complexation of  $C_{60}$  by **1-6**. Additional support for this notion comes from a newly developing absorption band, which features a  $C_{60}^{\delta-}/\text{exTTF}^{\delta+}$  charge transfer character, between 455 and 530 nm – peaking at 475 nm – in PhCl. When going to the more polar PhCN, a

shift of the charge transfer band is observed (455-550 nm, peaking at 485 nm). This spectral shift is rationalized on grounds of a better stabilization of  $C_{60}^{\delta-}/\text{exTTF}^{\delta+}$  (see Figure S2). This charge transfer interaction has also been observed in other donor- $C_{60}$  systems as a relatively significant contribution to the overall stability of exTTF based fullerene receptors.<sup>47</sup> As a representative example, the spectroscopic changes observed for **3** upon titration with  $C_{60}$  are depicted in Figure 1. These spectral changes are a clear signature for the association of **3** with  $C_{60}$  and were also observed for all other receptors (Figures S1 and S2 in the ESI†).



**Figure 1.** Absorption spectra of dilute PhCl solutions of **3** ( $1.5 \times 10^{-5}$  M) with variable concentrations of  $C_{60}$  upon subtraction of the fullerene absorption profile to highlight the absorption changes and the isosbestic point.

The binding constants of **1-6** with  $C_{60}$  were obtained with non-linear curve fitting analyses of the UV-vis absorption titration experiments. For all the studied systems, the best fits to the experimental data were obtained when assuming a 1:1 stoichiometry, which was also observed by MS-MALDI experiments (Figure S4). As documented in Table 1, the nature and size of the crown ether have a clear impact on the affinity towards  $C_{60}$ , leading to  $K_a$  values that vary by as much as three orders of magnitude. A clear trend between the size of the crown ether and the magnitude of the binding constant can be gathered in PhCl. Thus, the highest constant of the series is obtained for **3**, bearing the largest crown ether. The lowest binding constant is obtained for reference **6**, lacking the crown ether subunits. In between, both bis-aza-crown ether derivatives **4** and **5** exhibit significantly smaller  $K_a$  values when compared to their corresponding oxygen-bearing analogues **1** and **2**, respectively. This experimental finding could be accounted for by their less appropriate geometry to accommodate  $C_{60}$  as will be further discussed in the Computational Studies (see below).

**Table 1.** Calculated binding constants ( $K_a$ ) for exTTF molecular tweezers **1-6** towards  $C_{60}$  in PhCl at 298 K.

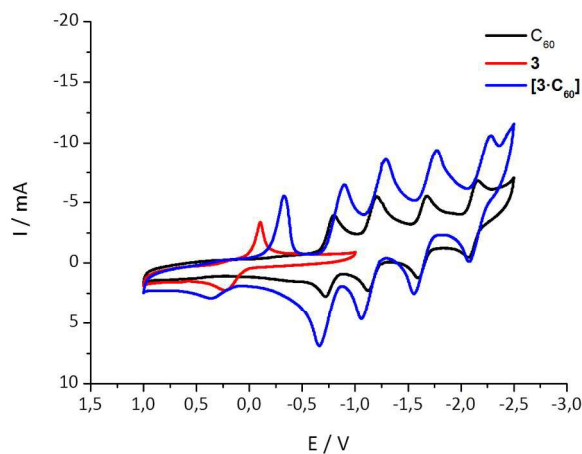
<b>1</b> • $C_{60}$	<b>2</b> • $C_{60}$	<b>3</b> • $C_{60}$	<b>4</b> • $C_{60}$	<b>5</b> • $C_{60}$	<b>6</b> • $C_{60}$
$4.8 \pm 0.9$	$6.7 \pm 0.2^{45}$	$6.9 \pm 0.2$	$3.8 \pm 0.6$	$5.1 \pm 0.1$	$3.3 \pm 0.4$

Insights into excited state interactions between **1-6** and  $C_{60}$  in either PhCl or PhCN came from emission studies exciting at 350, 400, and 450 nm (see ESI†). Upon addition of  $C_{60}$  to a solution of **1-**

6, a new and broad emission at around 530 and 550 nm in PhCl and PhCN, respectively, grows in at the expense of the intrinsic exTTF emission centered around 460 nm. The substantial 530 to 550 nm shift is due to the underlying intermolecular charge transfer character. Like in the ground state, more polar PhCN facilitates the stabilization of the  $(C_{60}^{\delta-} / \text{exTTF}^{\delta+})^*$  excited state, when compared to PhCl (see Figure S3). Importantly, the underlying energetics are, on one hand, comparable to those found for other  $C_{60}/\text{exTTF}$  systems, and, on the other hand, appreciably higher than what is typically found in  $C_{60}/\text{porphyrins}$  with values of 2.2 and 1.5 eV, respectively.<sup>48-53</sup>

### Electrochemical Studies

Further insights into the interactions in the ground state came from cyclic voltammetry (CV) investigations with equimolar mixtures of **1-6** and  $C_{60}$ . Notably, the lower stability of the azacrown ether complexes limited the electrochemical measurements to the crown ethers as summarized in Table S1. As an example, Figure 2 shows the CVs for  $C_{60}$ , **3**, and a 1:1 mixture of **3** and  $C_{60}$  (see ESI† for more details). Because of complexation, the first, second, third, and fourth quasireversible reductions of  $C_{60}$  are clearly shifted to more cathodic potentials (see Table S1 and Figure S5). The smallest shift for the first reduction ( $\sim 30$  mV) is found for  $6 \cdot C_{60}$  and it increases to  $\sim 50$  mV for both  $1 \cdot C_{60}$  and  $2 \cdot C_{60}$ , and to  $\sim 100$  mV for  $3 \cdot C_{60}$ . At this point, we postulate that the magnitude of the reduction potential shift, as seen upon complexation, relates to the binding strength. Electronic interactions between the electron-donating exTTF host and the electron-accepting  $C_{60}$  guest in the ground state are the basis for this trend and scale with the size of the crown ether. A shift is also found in the exTTF centered oxidation, moving towards more positive values (see Figure 2 and Figure S5). These shifts, however, do not fully correlate with the experimentally determined binding constants most likely due to adsorption phenomena upon oxidation.



**Figure 2.** Cyclic voltammetry of **3**,  $C_{60}$ , and  $3 \cdot C_{60}$  in a 4/1 v/v solvent mixture of PhCl and MeCN with 0.1 M  $n\text{-Bu}_4\text{NPF}_6$  and at 0.1  $\text{V}\cdot\text{s}^{-1}$ . Potentials are given versus  $\text{Ag}/\text{Ag}^+$ .

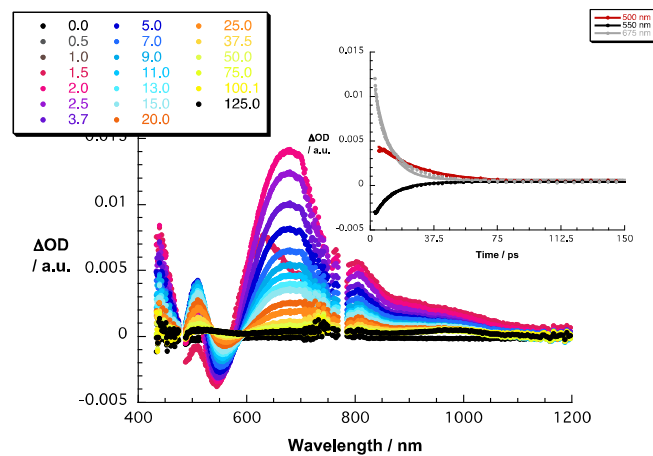
### Transient Absorption Studies

In transient absorption measurements with **1-5** in PhCl only a single transient evolves. The excited state transient, which is exTTF-centered, appears simultaneously with the conclusion of the 387 nm laser excitation. Transient maxima arise at 465, 605, and 910 nm,

while ground state bleaching is observed at wavelengths  $< 450$  nm. Kinetic analyses of the transients reveal short-lived excited states with lifetimes in the range from 0.9 to 1.1 ps. Such short lifetimes are rationalized on the basis of strong second-order vibronic spin-orbit coupling, which originate from the sulfur atoms.

With respect to  $C_{60}$ , upon excitation at 480 nm in PhCl, the characteristic singlet excited state transient emerges in the near infrared around 980 nm. This singlet excited state deactivates via intersystem crossing within 1.2 ns and produces the corresponding triplet excited state with a transient maximum at 750 nm and a lifetime of about 45  $\mu\text{s}$ .

480 nm excitation of equimolar mixtures of **1-5** and  $C_{60}$  [ $(C_{60}/\text{exTTF}) \approx 10^{-5}$  M] into the charge transfer band results in the instantaneous formation of photoexcited  $C_{60}^{\delta-} / \text{exTTF}^{\delta+}$  as seen in Figure 3 for  $1 \cdot C_{60}$  and in Figure S6 for the remaining complexes. This excited charge transfer state features maxima at 507 and 673 nm as well as a broad band in the near infrared around 950 nm. Additionally, transient bleaching is observed at around 550 nm. The latter relates, however, to stimulated charge transfer emission - *vide supra*. In terms of kinetics, the transients transform to the fully charge-separated state, that is,  $C_{60}^{\cdot-} / \text{exTTF}^{\cdot+}$ , on a time scale ranging from 1.6 to 2.3 ps (Table 2). In terms of spectroscopy, the presence of the characteristic transient absorption due to the one-electron oxidized exTTF in the visible at around 680 nm confirms our hypothesis.<sup>54-56</sup> Importantly, the latter is complemented by the feature of the one-electron reduced  $C_{60}$ , which maximizes in the near-infrared at around 1100 nm.<sup>57</sup> These radical ion pair states recombine in any of the probed systems within 12 to 21 ps into lower lying excited states of  $C_{60}$ , that is, the singlet and triplet excited states with maxima at 750 and 980 nm, respectively (Table 2). In general, stronger binding causes acceleration of the charge recombination – **3** vs. **2** and **5** vs. **4** – due to tighter interactions.



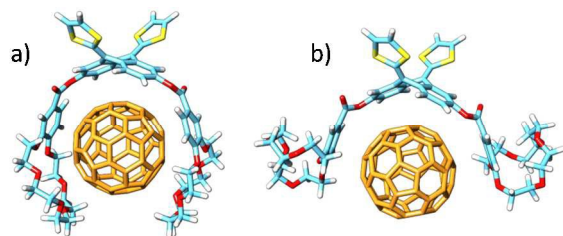
**Figure 3.** Differential absorption spectra (visible and near-infrared) obtained upon femtosecond flash photolysis (480 nm) of  $1 \cdot C_{60}$  (1:1) in PhCl [ $(C_{60}/\text{exTTF}) \approx 10^{-5}$  M] with several time delays between 0 and 125 ps at room temperature. Inset: time-absorption profiles of the spectra at 500, 550, and 675 nm monitoring the charge separation / charge recombination.

**Table 2.** Charge separation (CS) and charge recombination (CR) dynamics obtained upon laser flash photolysis at 480 nm of equimolar mixtures of **1-5** with  $C_{60}$  in argon-saturated PhCl.

Complex	CS (s <sup>-1</sup> )	CR (s <sup>-1</sup> )
1•C <sub>60</sub>	6.3 × 10 <sup>11</sup>	8.3 × 10 <sup>10</sup>
2•C <sub>60</sub>	4.3 × 10 <sup>11</sup>	4.8 × 10 <sup>10</sup>
3•C <sub>60</sub>	4.8 × 10 <sup>11</sup>	5.7 × 10 <sup>10</sup>
4•C <sub>60</sub>	4.3 × 10 <sup>11</sup>	7.6 × 10 <sup>10</sup>
5•C <sub>60</sub>	4.8 × 10 <sup>11</sup>	7.9 × 10 <sup>10</sup>

### Computational Studies

The different conformations that 1–6 may adopt when complexing C<sub>60</sub> were initially explored by using semiempirical PM7 calculations. Only 1:1 stoichiometric ratios were computed according to the experimental evidences. Figure 4a surveys the minimum-energy optimized structure calculated for 2•C<sub>60</sub> as a representative example (see Figure S7 for all the associates). In any of the complexes between 1–6 and C<sub>60</sub>, the latter interacts with the anthracene concave region of exTTF and, at the same time, the crown ether-based arms embrace C<sub>60</sub> in a pinzer-like shape. Non-embraced host–guest arrangements, in which the crown ethers fold themselves away from C<sub>60</sub>, were also optimized for 1•C<sub>60</sub>, 2•C<sub>60</sub> and 3•C<sub>60</sub> (see Figure 4b for 2•C<sub>60</sub>) to assess the stabilization due to the embracing movement. PM7 predicts association energies of –68.12, –72.43 and –88.75 kcal/mol for the embraced conformations of 1•C<sub>60</sub>, 2•C<sub>60</sub> and 3•C<sub>60</sub>, respectively, whereas the values for their non-embraced homologues were computed to be –50.49, –51.20 and –51.56 kcal/mol. Calculations therefore suggest that the embraced conformations are favored by an increase of the total binding energy that grows up with the size of the crown ether. Intermediate structures in which C<sub>60</sub> is embraced by only one arm of the exTTF-(crown ether)<sub>2</sub> receptor were also calculated for complexes 2•C<sub>60</sub> and 3•C<sub>60</sub>. For 2•C<sub>60</sub>, the two crown ether arms stabilize the complex by a similar energy amount of about –10.5 kcal/mol (Figure S8a). In contrast, the first arm of 3•C<sub>60</sub> stabilizes the complex by –24.9 kcal/mol due to the larger size of the crown ether and to the additional interaction with the terminal benzene ring, whereas the second arm leads to a significantly lower stabilization of –12.3 kcal/mol due to the steric hindrance between the two crown ether arms (Figure S8b).

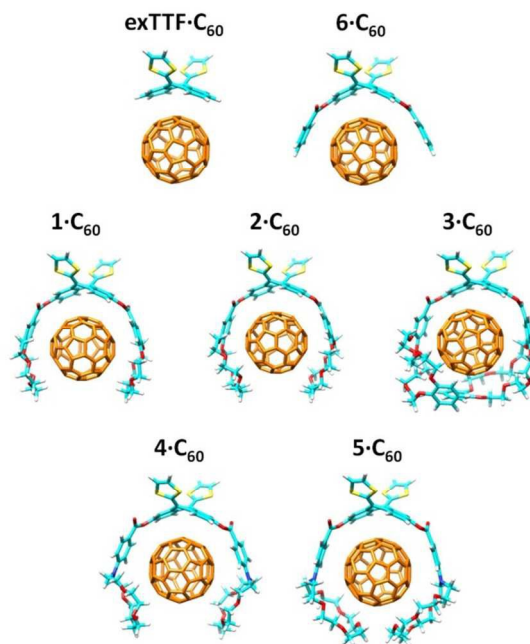


**Figure 4.** Minimum-energy embraced (a) and non-embraced (b) conformation calculated at the PM7 level for the 2•C<sub>60</sub> associate.

The more stable embraced conformations were subsequently optimized using the dispersion-corrected B97-D functional and the cc-pVDZ basis set (Figure 5). The exTTF•C<sub>60</sub> complex, which is not observed experimentally, was also calculated as a reference. Geometry optimizations were performed under C<sub>2</sub> symmetry restrictions but for 3•C<sub>60</sub>. For the latter, no symmetry was assumed

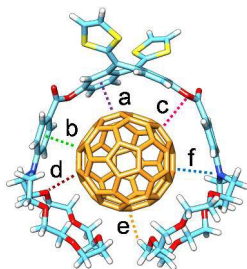
because the terminal benzene rings of the crown ethers disturb each other when complexing C<sub>60</sub>, resulting in a C<sub>1</sub> symmetry.

The B97-D/cc-pVDZ-optimized geometries reveal intermolecular contacts of different nature along the host-guest interface. Table 3 summarizes the shortest distances computed for the intermolecular contacts determining the stabilization of the complexes between 1–6 and C<sub>60</sub>. To estimate the binding energies, single-point energy calculations were performed on the B97-D/cc-pVDZ-optimized structures using the revPBE0-D3 functional and the more extended triple- $\zeta$  cc-pVTZ basis set. Table 3 collects the binding energies computed for the resulting complexes. A binding energy of –10.24 kcal/mol is predicted for exTTF•C<sub>60</sub> due to the  $\pi$ – $\pi$  interactions between the lateral benzene rings of exTTF and the benzene rings of C<sub>60</sub> with centroid–centroid distances of 3.42 Å (a in Table 3). Since exTTF•C<sub>60</sub> is not detected experimentally, entropic and solvent effects are expected to provide a positive contribution that cancels out the stabilizing interaction. In 6•C<sub>60</sub>, two additional interactions originating from the presence of the benzoates are found:  $\pi$ – $\pi$  interactions at 3.25 Å between the benzene rings of the benzoate moiety and C<sub>60</sub> (b in Table 3) and n– $\pi$  interactions due to short O(host)···C(guest) intermolecular distances (3.16 Å, c in Table 3). The positive effect of these interactions is evidenced by the folding angle of the anthracene in exTTF, which becomes sharper in passing from exTTF•C<sub>60</sub> (142.5°) to 6•C<sub>60</sub> (137.0°). The association energy computed for 6•C<sub>60</sub> amounts to –22.85 kcal/mol, which is more than twice of the binding energy found for exTTF•C<sub>60</sub>, and, in turn, is high enough to experimentally detect the complex in solution (Table 1).



**Figure 5.** B97-D/cc-pVDZ minimum-energy geometries calculated for the exTTF•C<sub>60</sub> and 1–6•C<sub>60</sub> complexes.

**Table 3.** Intermolecular distances (a–f, in Å) and binding energies (E<sub>bind</sub>, in kcal/mol) calculated at the B97-D/cc-pVDZ and revPBE0-D3/cc-pVTZ levels, respectively, for the exTTF•C<sub>60</sub> and 1–6•C<sub>60</sub> complexes.<sup>a</sup>



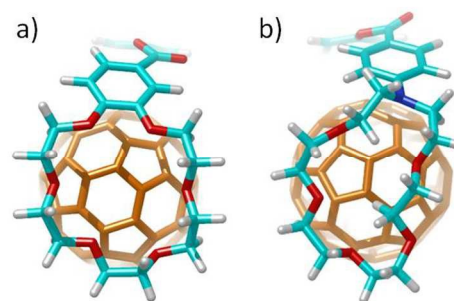
Complex	a	b	c	d	e	f	$E_{\text{bind}}$ (kcal/mol)
exTTF•C <sub>60</sub>	3.42	–	–	–	–	–	–10.24
<b>1</b> •C <sub>60</sub>	3.46	2.99	3.30	3.19	2.61	–	–39.69
<b>2</b> •C <sub>60</sub>	3.45	2.95	3.42	2.79	2.69	–	–44.76
<b>3</b> •C <sub>60</sub> <sup>b</sup>	3.49 <sup>c</sup>	2.98 <sup>c</sup>	3.44 <sup>c</sup>	2.85	2.50	–	–54.36
<b>4</b> •C <sub>60</sub>	3.37	3.41	3.25	3.56	2.57	4.14	–36.77
<b>5</b> •C <sub>60</sub>	3.37	3.06	3.14	3.14	2.54	3.50	–43.33
<b>6</b> •C <sub>60</sub>	3.45	3.25	3.16	–	–	–	–22.85

<sup>a</sup> **a** is the distance between the centroid of the lateral benzene rings of exTTF and that of the closest benzene rings of C<sub>60</sub>. **b** is the distance between the centroid of the benzene ring of the benzoate moiety and the center of the closest C<sub>60</sub> 6:6 double bond. **c** is the distance between the benzoate *sp*<sup>3</sup> oxygen and the closest carbon atom of C<sub>60</sub>. **d** and **e** are the shortest O...C<sub>60</sub> and H...C<sub>60</sub> distances, respectively, between the crown ether and C<sub>60</sub>. **f** is the distance between the nitrogen atom of the aza-crown ether and the closest carbon atom of C<sub>60</sub>. <sup>b</sup> Two additional  $\pi$ - $\pi$  interactions between the outer benzene rings of the crown ethers and C<sub>60</sub> are computed at 3.13 and 3.68 Å. <sup>c</sup> Average values.

Upon going from **6**•C<sub>60</sub> to **1**•C<sub>60</sub>, **2**•C<sub>60</sub>, and **3**•C<sub>60</sub> new n- $\pi$  (**d**) and CH... $\pi$  (**e**) interactions with intermolecular distances of 3.4 and 2.8 Å (averaged over all the O...C<sub>60</sub> and C-H...C<sub>60</sub> interactions shorter than 3.8 and 3.2 Å, respectively, in **1–3**•C<sub>60</sub> associates) contribute to the complex stabilization due to the inclusion of the crown ethers in the host system (Table 3). Calculations predict that the binding energies of the complexes rise as the size of the crown ether increases passing from –39.69 kcal/mol for **1**•C<sub>60</sub>, to –44.76 kcal/mol for **2**•C<sub>60</sub>, and to –54.36 kcal/mol for **3**•C<sub>60</sub>. This trend is in good agreement with the increase of the  $K_a$  value estimated experimentally (Table 1). It has to be attributed to increasing contributions from n- $\pi$  and CH... $\pi$  interactions, which are associated with the increasing size of the crown ethers when going from **1**•C<sub>60</sub> to **3**•C<sub>60</sub>. The crown ether arms wrap C<sub>60</sub> and lead to more compact complexes, in which the benzene rings of the benzoate moiety are closer (by 0.2 Å) to C<sub>60</sub> as compared, for example, with **6**•C<sub>60</sub> (distance **b** in Table 3). This gain in compactness underpins the positive effect that noncovalent interactions between C<sub>60</sub> and the crown ethers exert on the complex stability.

Finally, nitrogens, which bridge the crown ether and the benzoate in **4** and **5**, confer additional flexibility to the aza-crown ethers in **4**•C<sub>60</sub> and **5**•C<sub>60</sub>. The latter features structures that are more folded than their oxygenated analogues **1**•C<sub>60</sub> and **2**•C<sub>60</sub> (Figure 6). These structures are less appropriate to accommodate C<sub>60</sub> and, as a consequence, they lead to less efficient host-guest interactions. For instance, in **4**•C<sub>60</sub>, the intermolecular contacts defined by parameters **b** and **d** are found at significantly larger distances relative to **1**•C<sub>60</sub> (Table 3). The binding energies computed for **4**•C<sub>60</sub> (–36.77 kcal/mol) and **5**•C<sub>60</sub> (–43.33 kcal/mol) are indeed smaller than those computed for the oxygenated complexes **1**•C<sub>60</sub> and **2**•C<sub>60</sub> (–39.69

and –44.76 kcal/mol, respectively). The lower affinity in terms of interacting with C<sub>60</sub> for the azacrown ethers is in agreement with the experimentally determined binding constants (Table 1) and is ascribed to an overall weakening of the host-guest interactions provoked by the less efficiently oriented aza-crown ether arms. Calculations therefore suggest that the ability of the exTTF-based molecular tweezers to bind C<sub>60</sub> arises from an interplay of different  $\pi$ - $\pi$ , n- $\pi$  and CH... $\pi$  interactions, and that the size and nature of the crown ether are key factors for the relative stabilization of the resulting complexes between **1–6** and C<sub>60</sub>.



**Figure 6.** Side view of the B97-D/cc-pVDZ-optimized geometries calculated for complexes **2**•C<sub>60</sub> (left) and **5**•C<sub>60</sub> (right) showing the different spatial arrangement of the crown and aza-crown ethers, respectively, along the C<sub>60</sub> guest.

Theoretical calculations predict a charge transfer from exTTF to C<sub>60</sub> for all the complexes in the ground state. For **6**•C<sub>60</sub>, a small charge-transfer of 0.06e is computed that accounts for the negative shift of ~30 mV observed in the first reduction of **6**•C<sub>60</sub> when compared to C<sub>60</sub> (Table S1). **1–3** interact stronger with C<sub>60</sub> and a noticeable increase in the charge transferred to C<sub>60</sub> is obtained along the series: **1**•C<sub>60</sub> (0.14e), **2**•C<sub>60</sub> (0.15e), and **3**•C<sub>60</sub> (0.18e). Such an increase justifies the larger cathodic shifts measured for the first reductions in **1**•C<sub>60</sub> and **2**•C<sub>60</sub> (~50 mV) as well as in **3**•C<sub>60</sub> (~100 mV) when compared to C<sub>60</sub> (Table S1). In the excited state, electron promotion from the HOMO to the LUMO, which are respectively localized on exTTF and C<sub>60</sub> (Figure S9), leads to a fully charge-separated C<sub>60</sub><sup>•–</sup>/exTTF<sup>•+</sup> associate.

## Conclusions

In summary, we have obtained a series of new exTTF-(crown ether)<sub>2</sub> receptors featuring two crown ethers of different size and composition. These receptors bind C<sub>60</sub> in PhCl with moderate to high efficiencies. The interactions between **1–6** and C<sub>60</sub> have been further studied both in the ground state by CV, pointing to the presence of electronic interactions, and in the excited state by transient absorption studies. Importantly, the latter corroborate the formation of radical ion pair states, which feature lifetimes in the 12–21 ps range. Complementary computational investigations have further documented the stabilization energy associated with the embraced conformation (25–30 %) and have provided critical insights into each of the interactions involved in the process. The nature of the supramolecular exTTF-(crown ether)<sub>2</sub>•C<sub>60</sub> affinity interactions arises, thus, from an interplay of  $\pi$ - $\pi$ , n- $\pi$  and CH... $\pi$  forces whose intensity depends on the size and nature of the crown ether.

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## Dedications

This work is dedicated to Professor José Barluenga on the occasion of his 75th birthday.

## Notes and references

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- S. Berardi, S. Drouet, L. Francas, C. Gimbert-Surinach, M. Guttentag, C. Richmond, T. Stoll and A. Llobet, *Chem. Soc. Rev.*, 2014, **43**, 7501-7519.
  - G. Bottari, O. Trukhina, M. Ince and T. Torres, *Coord. Chem. Rev.*, 2012, **256**, 2453-2477.
  - H. W. Kroto, J. R. Heath, S. C. O'Brien, R. F. Curl and R. E. Smalley, *Nature*, 1985, **318**, 162-163.
  - I. Hiroshi, H. Kiyoshi, A. Tsuyoshi, A. Masanori, T. Seiji, O. Tadashi, S. Masahiro and S. Yoshiteru, *Chem. Phys. Lett.*, 1996, **263**, 545-550.
  - Q. Xie, E. Pérez-Cordero and L. Echegoyen, *J. Am. Chem. Soc.*, 1992, **114**, 3978-3980.
  - L. Echegoyen and L. E. Echegoyen, *Acc. Chem. Res.*, 1998, **31**, 593-601.
  - H. Ajie, M. M. Alvarez, S. J. Anz, R. D. Beck, F. Diederich, K. Fostiropoulos, D. R. Huffman, W. Kraetschmer, Y. Rubin, K. E. Schriver, D. Sensharma and R. L. Whetten, *J. Phys. Chem.*, 1990, **94**, 8630-8633.
  - N. Martín, L. Sánchez, B. Illescas and I. Pérez, *Chem. Rev.*, 1998, **98**, 2527-2548.
  - D. M. Guldi and M. Prato, *Acc. Chem. Res.*, 2000, **33**, 695-703.
  - S. Castellanos, A. A. Vieira, B. M. Illescas, V. Sacchetti, C. Schubert, J. Moreno, D. M. Guldi, S. Hecht and N. Martín, *Angew. Chem. Int. Ed.*, 2013, **52**, 13985-13990.
  - F. D'Souza and O. Ito, *Chem. Commun.*, 2009, 4913-4928.
  - G. Bottari, G. de la Torre, D. M. Guldi and T. Torres, *Chem. Rev.*, 2010, **110**, 6768-6816.
  - A. Mateo-Alonso, D. M. Guldi, F. Paolucci and M. Prato, *Angew. Chem. Int. Ed.*, 2007, **46**, 8120-8126.
  - D. M. Guldi and N. Martín, *J. Mater. Chem.*, 2002, **12**, 1978-1992.
  - N. Martín and J.-F. Nierengarten, eds., *Supramolecular Chemistry of Fullerenes and Carbon Nanotubes*, Wiley-VCH, Weinheim, 2012.

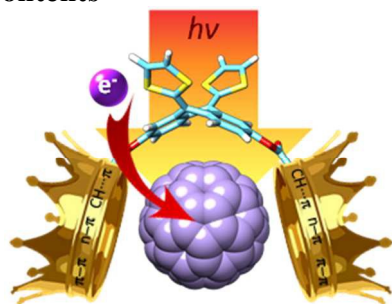
- L. Sánchez, M. Sierra, N. Martín, A. J. Myles, T. J. Dale, J. Rebek, W. Seitz and D. M. Guldi, *Angew. Chem. Int. Ed.*, 2006, **45**, 4637-4641.
- T. Andersson, K. Nilsson, M. Sundahl, G. Westman and O. Wennerstrom, *J. Chem. Soc., Chem. Commun.*, 1992, DOI: 10.1039/C39920000604, 604-606.
- Z.-i. Yoshida, H. Takekuma, S.-i. Takekuma and Y. Matsubara, *Angew. Chem. Int. Ed.*, 1994, **33**, 1597-1599.
- Y. Liu, G.-S. Chen, Y. Chen, N. Zhang, J. Chen and Y.-L. Zhao, *Nano Lett.*, 2006, **6**, 2196-2200.
- J. L. Atwood, G. A. Koutsantonis and C. L. Raston, *Nature*, 1994, **368**, 229-231.
- T. Suzuki, K. Nakashima and S. Shinkai, *Chem. Lett.*, 1994, **23**, 699-702.
- T. Hirao, M. Tosaka, S. Yamago and T. Haino, *Chem. Eur. J.*, 2014, DOI: 10.1002/chem.201404328, n/a-n/a.
- J. W. Steed, P. C. Junk, J. L. Atwood, M. J. Barnes, C. L. Raston and R. S. Burkharter, *J. Am. Chem. Soc.*, 1994, **116**, 10346-10347.
- J.-F. Nierengarten, L. Oswald, J.-F. Eckert, J.-F. Nicoud and N. Armaroli, *Tetrahedron Lett.*, 1999, **40**, 5681-5684.
- D. Felder, B. Heinrich, D. Guillon, J.-F. Nicoud and J.-F. Nierengarten, *Chem. Eur. J.*, 2000, **6**, 3501-3507.
- Y. Sun, T. Drovetskaya, R. D. Bolskar, R. Bau, P. D. W. Boyd and C. A. Reed, *J. Org. Chem.*, 1997, **62**, 3642-3649.
- K. Tashiro, T. Aida, J.-Y. Zheng, K. Kinbara, K. Saigo, S. Sakamoto and K. Yamaguchi, *J. Am. Chem. Soc.*, 1999, **121**, 9477-9478.
- D. Sun, F. S. Tham, C. A. Reed, L. Chaker and P. D. W. Boyd, *J. Am. Chem. Soc.*, 2002, **124**, 6604-6612.
- A. Hosseini, S. Taylor, G. Accorsi, N. Armaroli, C. A. Reed and P. D. W. Boyd, *J. Am. Chem. Soc.*, 2006, **128**, 15903-15913.
- J. Song, N. Aratani, H. Shinokubo and A. Osuka, *J. Am. Chem. Soc.*, 2010, **132**, 16356-16357.
- C. G. Claessens and T. Torres, *Chem. Commun.*, 2004, DOI: 10.1039/B401232A, 1298-1299.
- I. Sánchez-Molina, B. Grimm, R. M. Krick Calderon, C. G. Claessens, D. M. Guldi and T. Torres, *J. Am. Chem. Soc.*, 2013, **135**, 10503-10511.
- I. Sanchez-Molina, C. G. Claessens, B. Grimm, D. M. Guldi and T. Torres, *Chem. Sci.*, 2013, **4**, 1338-1344.
- E. M. Pérez, L. Sánchez, G. Fernández and N. Martín, *J. Am. Chem. Soc.*, 2006, **128**, 7172-7173.
- D. Canevet, M. Gallego, H. Isla, A. de Juan, E. M. Pérez and N. Martín, *J. Am. Chem. Soc.*, 2011, **133**, 3184-3190.
- G. Fernández, E. M. Pérez, L. Sánchez and N. Martín, *Angew. Chem. Int. Ed.*, 2008, **47**, 1094-1097.
- H. Isla, E. M. Pérez and N. Martín, *Angew. Chem. Int. Ed.*, 2014, **53**, 5629-5633.
- J. Effing, U. Jonas, L. Jullien, T. Plesniviy, H. Ringsdorf, F. Diederich, C. Thilgen and D. Weinstein, *Angew. Chem. Int. Ed.*, 1992, **31**, 1599-1602.
- C. J. Pedersen, *Angew. Chem. Int. Ed.*, 1988, **27**, 1021-1027.
- R. E. Smalley, *Angew. Chem. Int. Ed.*, 1997, **36**, 1594-1601.
- S. Bhattacharya, A. Sharma, S. K. Nayak, S. Chattopadhyay and A. K. Mukherjee, *J. Phys. Chem. B*, 2003, **107**, 4213-4217.
- A. Saha, S. K. Nayak, S. Chattopadhyay and A. K. Mukherjee, *J. Phys. Chem. B*, 2003, **107**, 11889-11892.
- Y. Liu, J.-R. Han, Y.-L. Zhao, H.-Y. Zhang and Z.-Y. Duan, *J. Inclusion Phenom. Macrocyclic Chem.*, 2005, **51**, 191-198.
- F. D'Souza, R. Chitta, S. Gadde, M. E. Zandler, A. L. McCarty, A. S. D. Sandanayaka, Y. Araki and O. Ito, *J. Phys. Chem. A*, 2006, **110**, 4338-4347.
- B. Grimm, J. Santos, B. M. Illescas, A. Muñoz, D. M. Guldi and N. Martín, *J. Am. Chem. Soc.*, 2010, **132**, 17387-17389.
- S. González, N. Martín and D. M. Guldi, *J. Org. Chem.*, 2002, **68**, 779-791.
- E. M. Pérez, A. L. Capodilupo, G. Fernández, L. Sánchez, P. M. Viruela, R. Viruela, E. Ortí, M. Bietti and N. Martín, *Chem. Commun.*, 2008, 4567-4569.
- S. S. Gayathri, M. Wielopolski, E. M. Pérez, G. Fernández, L. Sánchez, R. Viruela, E. Ortí, D. M. Guldi and N. Martín, *Angew. Chem. Int. Ed.*, 2009, **48**, 815-819.
- S. G. Radhakrishnan, D. M. Guldi, E. M. Pérez, I. Pérez, M. Bietti and N. Martín, *J. Coord. Chem.*, 2010, **63**, 2939-2948.
- B. Grimm, H. Isla, E. M. Perez, N. Martín and D. M. Guldi, *Chem. Commun.*, 2011, **47**, 7449-7451.

## Journal Name

51. V. Chukharev, N. V. Tkachenko, A. Efimov, D. M. Guldi, A. Hirsch, M. Scheloske and H. Lemmetyinen, *J. Phys. Chem. B*, 2004, **108**, 16377-16385.
52. J. L. Sessler, J. Jayawickramarajah, A. Gouloumis, T. Torres, D. M. Guldi, S. Maldonado and K. J. Stevenson, *Chem. Commun.*, 2005, DOI: 10.1039/B418345B, 1892-1894.
53. D. M. Guldi, C. Luo, M. Prato, E. Dietel and A. Hirsch, *Chem. Commun.*, 2000, DOI: 10.1039/A909657D, 373-374.
54. D. M. Guldi, L. Sánchez and N. Martín, *J. Phys. Chem. B*, 2001, **105**, 7139-7144.
55. A. E. Jones, C. A. Christensen, D. F. Perepichka, A. S. Batsanov, A. Beeby, P. J. Low, M. R. Bryce and A. W. Parker, *Chem. Eur. J.*, 2001, **7**, 973-978.
56. N. Martín, L. Sánchez, M. Á. Herranz, B. Illescas and D. M. Guldi, *Acc. Chem. Res.*, 2007, **40**, 1015-1024.
57. D. R. Lawson, D. L. Feldheim, C. A. Foss, P. K. Dorhout, C. M. Elliott, C. R. Martin and B. Parkinson, *J. Electrochem. Soc.*, 1992, **139**, L68-L71.



## Table of contents



A series of exTTF-(crown ether)<sub>2</sub> receptors, designed to host C<sub>60</sub>, has been prepared in order to understand the nature of the fullerene-crown ether interaction. The combination of experimental and in silico studies points to a synergetic interplay of donor-acceptor,  $\pi$ - $\pi$ , n- $\pi$  and CH $\cdots$  $\pi$  interactions as the basis of this affinity