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Self-sorting of crown ether/secondary ammonium ion hetero-[c2]daisy chain pseudorotaxanes†

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Four monomeric building blocks equipped with one crown ether and one secondary ammonium ion are synthesized and studied with respect to their ability to form daisy chain dimers. Two crown ethers with different cavity sizes – *i.e.* [21]crown-7 and [24]crown-8 – and two ammonium ions substituted with either a thin alkyl group or a more bulky benzyl group are used as the binding motifs. Self-sorting behaviour can be expected as the [21]crown-7/alkyl ammonium and [24]crown-8/benzyl ammonium binding motifs are orthogonal. Three homodimers are characterized by NMR, X-ray crystallography and ESI mass spectrometry. They are recognizable by the presence of signals for diastereotopic protons in the ¹H NMR spectra. The formation of hetero-[c2]daisy chain dimers can be monitored by NMR spectroscopy and ESI mass spectrometry and show the expected self-sorting behaviour.

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Introduction

Nature efficiently and successfully assembles intricate and highly complex architectures,¹ in which high-fidelity self-sorting² is ubiquitous as the basis of the building blocks' cooperative functional interplay. Many supramolecular chemists have recently constructed synthetic self-sorting systems with orthogonal binding motifs fabricated on the basis of hydrogen bonding,³ metal–ligand interactions,⁴ π - π -stacking⁵ or solvophobic effects.⁶ A number of different factors, among them the orthogonality of the recognition motifs, size, shape, thermodynamic and kinetic binding parameters, stoichiometry and concentration define the molecular codes and are responsible for self-recognition (narcissistic self-sorting) or self-discrimination (social self-sorting).^{2,5a,7}

Recently, we have applied the concepts of social self-sorting to pseudorotaxane⁸ assemblies and supramolecular pseudorotaxane polymers based on the crown ether/secondary ammonium ion binding motifs.⁹ Benzo[21]crown-7 **C7** and dibenzo[24]crown-8 **C8** (Fig. 1) have two different cavity sizes and their pseudorotaxane binding motifs with secondary ammonium ions can be made orthogonal by using either

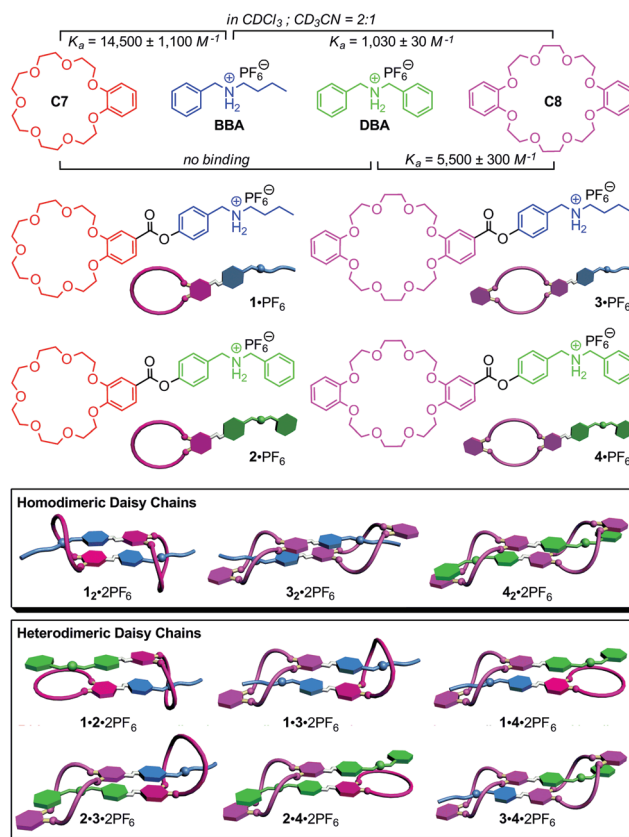


Fig. 1 Top: Structures and binding data of building blocks **C7**, **C8**, **BBA** and **DBA**. Centre: Monomers **1-PF₆**–**4-PF₆**. Bottom: Cartoons representing all possible homo- and heterodimeric daisy chains. Note that dimer **2-2-2PF₆** does not form and **1-2-2PF₆** and **1-4-2PF₆** cannot ring-close.

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benzyl-*n*-butyl ammonium **BBA** with one thin alkyl ammonium or dibenzyl ammonium **DBA** with two more bulky benzyl ammonium stopper groups. As the benzyl stopper is too large to penetrate **C7**, only three host-guest complexes **BBA-C7**, **BBA-C8** and **DBA-C8** can form. Fig. 1 summarizes the binding constants for these pairs in a CDCl_3 - CH_3CN (2:1) solvent mixture. When the feed ratio is controlled and all four components are mixed in equimolar amounts, self-sorting is obtained, because the **DBA** guest cation consumes all **C8** host molecules while only **C7** binds **BBA**. But even, when feed ratio is not perfectly matched, this self-sorting behaviour is expected to occur, because the **BBA-C7** and **DBA-C8** pairs exhibit the two highest binding constants of 14500 M^{-1} and 5500 M^{-1} , respectively, among the three possible pseudorotaxanes.¹⁰ Consequently, the resulting self-sorting is driven kinetically (stopper sizes) as well as thermodynamically (binding constants) into the same direction.

[c2]Daisy chains are cyclic interwoven oligo- or even polymers¹¹ that are usually constructed from AB-type pleiotopic monomers equipped with two complementary binding units A (host) and B (guest). Usually, small, often dimeric macrocyclic assemblies result, because they are enthalpically (all binding sites involved in binding) as well as entropically (high particle number) favourable. Such self-complementary building blocks – when equipped with two switchable stations – are also excellent building blocks for molecular muscle-type supramolecular polymers.¹² The daisy chains known so far are all constructed from identical monomers. This limits the structural diversity that can be obtained by self-assembly. In contrast, progress towards functional complexes containing different cooperating building blocks to realize high-level function would require the ability to build well-defined architectures, in which each subunit occupies a precisely controlled position. It is therefore highly desirable to provide strategies for programming different monomers into heteromeric assemblies. Self-sorting is one potential strategy in this direction.

Here, we apply the principles of social self-sorting to the construction of heterodimeric [c2]daisy chains. The aim of the present study is to investigate their self-sorting behaviour and to extend the scope of the self-assembly of hierarchical heteromeric assemblies. The investigation of heteromeric assemblies helps understand self-sorting better and will contribute to paving a way to build well-defined assemblies that carry cooperating groups for implementing function. Four different, yet structurally quite similar monomers **1-PF₆**–**4-PF₆** were designed (Fig. 1, centre) based on the two different crown ethers **C7** and **C8** and two different secondary ammonium ions **BBA** and **DBA**. In order to obtain these four AB-type monomers, the two binding sites – one crown ether and one ammonium ion – are connected through ester linkages as reported earlier for **1-PF₆**.¹³

Based on the self-sorting behaviour of **C7**, **C8**, **BBA** and **DBA**, one would arrive at the following expectations (Fig. 1, bottom). As **2-PF₆** is not self-complementary due to the combination of the larger benzyl stopper with smaller crown, this monomer is predicted not to dimerize. In solutions of the

pure monomers, three homodimers should therefore form under the appropriate conditions: **1₂-2PF₆**, **3₂-2PF₆** and **4₂-2PF₆**. A 1:1 mixture of **2-PF₆** and **3-PF₆** should convert fully into the heterodimer **2-3-2PF₆** (perfect self-sorting), because the two building blocks are matching each other. Binding can only involve all binding sites, when the heterodimeric daisy chain forms exclusively. Otherwise, monomeric **2-PF₆** would remain with unsaturated binding sites along with the **3₂-2PF₆** homodimer. For a 1:1 mixture of **1-PF₆** and **3-PF₆**, one predicts a statistical 1:2:1 mixture of **1₂-2PF₆**, **1-3-2PF₆** and **3₂-2PF₆** as the final result of the assembly, since both homodimers as well as the heterodimer can easily form and their equilibration should be more or less thermoneutral. For all other combinations, no significant heterodimer formation is expected, as the heterodimers **1-2-2PF₆**, **1-4-2PF₆** and **2-4-2PF₆** can merely form singly threaded heterodimeric daisy chains. The formation of heterodimers would thus lead to unsaturated binding sites and is then less favourable as the mixture of both homodimers.

Results and discussion

Monomer synthesis and characterisation

The synthesis of **2-PF₆**–**4-PF₆** follows the previously reported preparation of **1-PF₆** (Fig. 2).^{13,14} Briefly, the acid derivatives of the two crown ethers, **C7-COOH** and **C8-COOH**, were coupled with Boc-protected derivatives **5** and **6** of **DBA** and **BBA**, respectively. After de-protection and ion exchange, the desired target compounds **1-PF₆**–**4-PF₆** were obtained.

In order to characterize the samples as monomers, NMR spectra were recorded in $\text{DMSO-}d_6$ (ESI[†]), in which only one set of signals is observed. Although the substitution of the crown ethers' benzo moieties with the ester group renders all methylene groups of the crowns non-equivalent, they are somewhat broadened as they are pairwise almost isochronous and thus closely overlapping mostly depending on the distance to the aromatic ring.

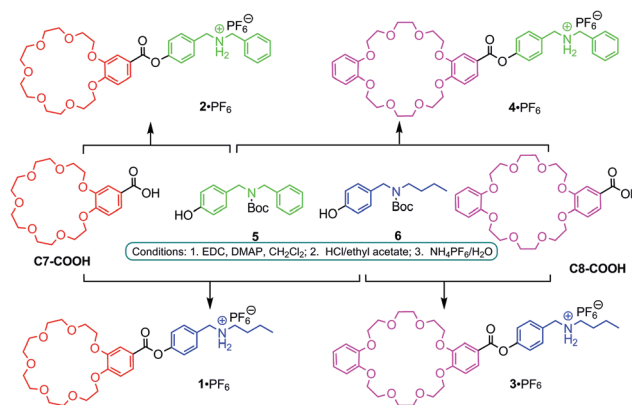


Fig. 2 Synthesis of monomers **1-PF₆**–**4-PF₆**. (EDC = 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride; DMAP = 4-dimethylaminopyridine).



Homodimers $1_2\cdot 2PF_6$, $3_2\cdot 2PF_6$ and $4_2\cdot 2PF_6$

Before discussing the formation of homodimers, a brief comment on the chirality features of the daisy chains may be indicated. All four monomers are achiral compounds. Upon threading an unsymmetrical axle through each of the crown ethers, chiral complexes are possible. The ester groups define directionality around the crown ether; the axle defines a direction perpendicular to the crown ether plane. Consequently, such a complex exhibits planar chirality. This is certainly true for all heterodimeric [c2]daisy chains, which may thus form four different stereoisomers that are in equilibrium with each other. However, one should be aware of the fact that homodimeric [c2]daisy chains may exist as a pair of enantiomers plus one achiral *meso*-diastereomer bearing a centre of inversion so that only three stereoisomers can exist in equilibrium here.

In less competitive solvents, three of the monomers, $1\cdot PF_6$,¹³ $3\cdot PF_6$ and $4\cdot PF_6$, are expected to form homodimers. Indeed, the 1H NMR spectra of these three compounds in a 2 : 1 mixture of chloroform and acetonitrile (Fig. 3) are much more complex than those in DMSO, while $2\cdot PF_6$ retains its simplicity also in this less competitive solvent mixture. The increased complexity is caused by two effects: threading on one hand leads to more significant differences in the environments of those crown ether methylene groups that overlap in the spectra of the monomers. In addition, the two faces of the crown ethers become different upon threading of the axle. Thus, at least some of the methylene groups give rise to separate signals for the two protons. Even with the help of 2D spectra, a complete assignment of all signals is difficult as many of them still strongly overlap (ESI†). The best signal separation is found for $4_2\cdot 2PF_6$ (Fig. 3) so that we can clearly identify two pairs of diastereomers (one from the phenyl-methyl proton and one from the crown methylene) and other diastereotopic signals of crown CH_2 groups here (ESI†). Also, typical complexation-induced shifts indicate the formation of daisy chain dimers: In particular, the aromatic protons H^a and H^c of $1_2\cdot PF_6$, $3_2\cdot PF_6$ and $4_2\cdot PF_6$ are shifted up-field relative to

those of the free monomer in DMSO- d_6 , the monomer form of $2\cdot PF_6$ in $CDCl_3$ - CD_3CN (2 : 1) or the potassium complex (*vide infra*), with the most significant shift observed for H^a . This indicates – irrespective of the reference system chosen – the up-field shift to be due to the anisotropy effect of stacked aromatic rings which are only present in the daisy chain dimers, but do not exist in the monomers. Together with the observation of intense signals for the homodimers in the ESI mass spectra, the drastic changes in the 1H NMR spectra for $1\cdot PF_6$, $3\cdot PF_6$ and $4\cdot PF_6$ and the absence of similar changes for $2\cdot PF_6$ clearly indicate the formation of the expected homodimers. Even though thermodynamically controlled macrocyclisations can give rise to larger cyclic oligomers,¹⁵ no larger complexes than dimers were detected in the mass spectra so that we rule out significant contributions from larger oligomers to the equilibrium situation.

Another piece of evidence for the formation of homodimers is their switchability with a competitive guest. When KPF_6 is added to the NMR solutions of the three homodimers (ESI†), the complex NMR spectra return into much simpler ones that again show only one set of signals. These signals of the potassium complex are shifted only very slightly relative to the uncomplexed monomer as demonstrated for example for $2\cdot PF_6$ (ESI†).

Although all attempts to crystallize the heterodimers under study failed unfortunately, single crystals of homodimers $3_2\cdot 2PF_6$ and $4_2\cdot 2PF_6$ were obtained. These solid state structures are shown in Fig. 4a, c and e together with the previously reported crystal structure of $1_2\cdot 2PF_6$,¹³ which is added to the figure to facilitate direct comparison. It is clear that in the crystal structures all three homodimers are doubly-threaded head-to-tail daisy chains. The ammonium ions are positioned inside the cavities of the crown ethers by multiple N-H...O and C-H...O hydrogen bonds. Table 1 summarizes the hydrogen bonding distances and angles for N-H...O and C-H...O hydrogen bonds which are in the ranges expected for crown ether/ammonium binding motifs. Apparently, the formation of dimers is also supported by face-to-face π -stacking interaction

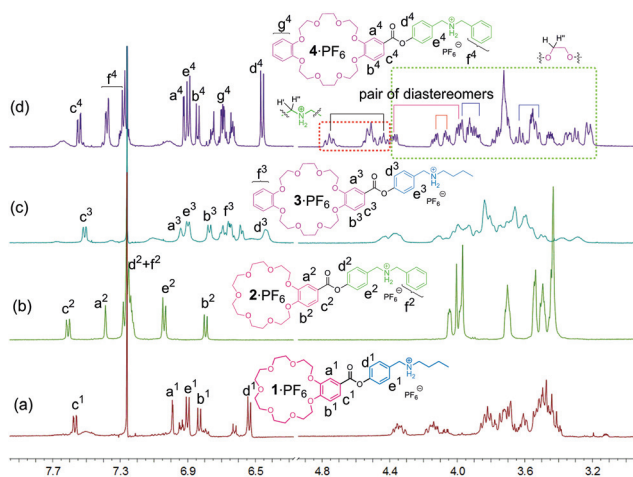


Fig. 3 Partial 1H NMR spectra (500 MHz, $CDCl_3$ - CD_3CN (2 : 1) 10.0 mM, 298 K) of (a) $1_2\cdot 2PF_6$, (b) $2\cdot PF_6$, (c) $3_2\cdot 2PF_6$ and (d) $4_2\cdot 2PF_6$.

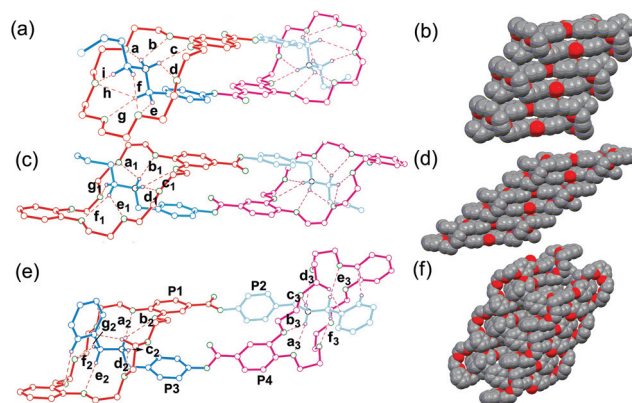


Fig. 4 Left: Ball-stick views of the X-ray structures of (a) $1_2\cdot 2PF_6$, (c) $3_2\cdot 2PF_6$ and (e) $4_2\cdot 2PF_6$. PF_6^- counterions and hydrogen atoms except the ones involved in hydrogen bonding are omitted for clarity. Right: Space-filling representations of the packing of (b) $1_2\cdot 2PF_6$, (d) $3_2\cdot 2PF_6$ and (f) $4_2\cdot 2PF_6$.



Table 1 C–H...O and N–H...O hydrogen-bonding parameters: H...O, C...O, and N...O distances are given in Å, C–H...O and N–H...O angles in degrees; for comparison, the previously published crystal structure data of $1_2\cdot 2PF_6$,¹³ is included. Labels a_n-i_n refer to the bond labels in Fig. 4

		$1_2\cdot 2PF_6$ (a-i)	$3_2\cdot 2PF_6$ (a ₁ -g ₁)	$4_2\cdot 2PF_6$ (a ₂ -g ₂)	$4_2\cdot 2PF_6$ (a ₃ -f ₃)
a	$d_{H...O}$	2.46(3)	2.49(5)	2.32(4)	2.40(8)
	$d_{N...O}$	3.12(5)	3.20(2)	3.14(5)	3.32(1)
	Angle	130.6(4)	129.4(6)	139.7(6)	170.7(2)
b	$d_{H...O}$	2.16(6)	2.29(0)	2.56(5)	2.58(7)
	$d_{N...O}$	2.98(3)	3.14(0)	3.47(1)	3.11(1)
	Angle	150.5(7)	153.3(8)	168.4(4)	116.6(7)
c	$d_{H...O}$	2.25(3)	2.40(2)	2.46(9)	2.05(3)
	$d_{N...O}$	2.91(2)	3.04(0)	3.13(1)	2.96(2)
	Angle	129.7(6)	126.5(4)	129.0(0)	169.6(7)
d	$d_{H...O}$	2.11(2)	1.99(6)	2.05(0)	2.56(6)
	$d_{N...O}$	2.91(6)	2.86(5)	2.91(7)	3.11(8)
	Angle	148.2(0)	157.0(0)	156.9(2)	119.0(4)
e	$d_{H...O}$	2.61(4)	2.64(5)	2.33(8)	2.49(9)
	$d_{C...O}$	3.39(7)	3.45(6)	2.97(5)	3.48(1)
	Angle	138.0(0)	139.0(5)	126.2(6)	171.0(6)
f	$d_{H...O}$	2.29(5)	2.56(0)	2.51(5)	2.35(6)
	$d_{C...O}$	3.18(8)	3.47(1)	3.48(7)	3.13(7)
	Angle	152.7(0)	152.8(6)	167.0(6)	135.1(1)
g	$d_{H...O}$	2.71(1)	2.62(4)	2.55(1)	
	$d_{C...O}$	3.15(9)	3.59(8)	3.43(0)	
	Angle	108.7(2)	167.7(6)	147.9(3)	
h	$d_{H...O}$	2.64(2)			
	$d_{C...O}$	3.49(9)			
	Angle	147.4(9)			
i	$d_{H...O}$	2.40(5)			
	$d_{C...O}$	3.28(0)			
	Angle	149.7(6)			

between the aromatic ring of the host moiety and the phenyl ring of the ammonium salt unit. The centroid-centroid distances are 3.64 Å in $1_2\cdot 2PF_6$ and 3.68 Å in $3_2\cdot 2PF_6$. The structure of $4_2\cdot 2PF_6$ differs in three respects from the other two: (i) although both monomers in the homodimer are identical, its crystal structure reveals two different binding motifs with different hydrogen bonding parameters realized – an effect that is likely due to packing effects. (ii) The aromatic rings are not aligned in a face-to-face fashion, but are twisted into arrangements with inclinations of 70.1° and 82.2° between the two planes of adjacent aromatic rings. (iii) Both ester carbonyl groups point in the same direction, while the other two homodimers have ester groups that point in opposite directions. Consequently, $1_2\cdot 2PF_6$ and $3_2\cdot 2PF_6$ bear a centre of inversion and realize the above-mentioned *meso*-forms, while no centre of inversion exists in the crystal of $4_2\cdot 2PF_6$, which thus is present as a chiral complex. The daisy chain homodimers pack in the crystal to yield infinite supramolecular tapes driven by face-to-face π -stacking interactions (Fig. 4b, d and f). In the crystal structure of $4_2\cdot 2PF_6$, homochiral tapes each containing only one of the two enantiomeric assemblies alternate.

Mass spectrometry as a method to detect self-sorting

As discussed above, already the NMR spectra of the homodimers are rather complicated and difficult to interpret. If one attempts to analyse the generation of heterodimers, even more

complicated spectra are expected to be observed. A detailed understanding, in particular a quantification of the contributions of homo- and heterodimers in solution will therefore not be straightforward by NMR spectroscopy. Crystallography neither provides a picture of the solution situation, even if suitable crystals could be grown, as solution equilibria may be shifted significantly upon crystallization. Therefore, ESI mass spectrometry appears to be a favourable tool to at least semi-quantitatively analyse the self-sorting behaviour of the daisy chains under study. A number of earlier studies on the assembly of other self-sorting pseudorotaxanes support this.^{3d,e,6b,9b,c,e,f}

Four points, however, may limit the scope of the analysis and should be considered in the interpretation of the data: (i) doubly charged homodimers will appear at the same *m/z* as singly charged monomers. They can however be easily distinguished as the peak spacing in the isotope patterns is 0.5 for the dicationic dimer and 1 for the singly charged monomer. Isotope pattern deconvolution has therefore been performed for all samples under study here. (ii) The sample solution concentrations used in ESI mass spectrometry are considerably lower than NMR concentrations. This may lead to a shift of the monomer-dimer equilibria towards the monomers so that the monomers may appear more prominently than expected from solution experiments. Furthermore, the monomers may form from the dimers, if the ions dissociate in part during ionization. Consequently, some contribution from monomeric species is expected to be observed. In the experiments reported here, sample concentrations were 100 μ M in each monomer to reduce the concentration difference of NMR and MS samples as much as possible. (iii) Ion intensities do not necessarily directly reflect solution concentrations. The efficiencies with which the ions form during the electrospray process depend significantly on the desolvation energies¹⁶ which may vary for different structures. Although the differences in the so-called ESI response factors are often not very large for structurally similar ions, exceptions exist. (iv) Non-specific binding often occurs in ESI mass spectrometry. If it is prominent, it may blur the analysis of self-sorting, because non-specific complexes will not exhibit self-sorting and superimpose the true distribution of self-sorted complexes. In our study, we chose ionization conditions such that non-specific binding is minimal.

The $1_2\cdot 2PF_6/4_2\cdot 2PF_6$ (1 : 1) and $2\cdot PF_6/4_2\cdot 2PF_6$ (2 : 1) mixtures

When $1_2\cdot 2PF_6$ and $4_2\cdot 2PF_6$ are mixed in equimolar amounts, no significant change is expected to occur. Both compounds form homodimers already in the stock solutions before mixing so that all binding sites are saturated. The heterodimer instead must remain singly threaded if it forms at all as the benzyl stopper in $4\cdot PF_6$ is unable to thread through the [21]-crown-7 ether in $1\cdot PF_6$. As this situation is energetically less favourable, there is consequently no driving force for heterodimer formation.

Fig. 5a shows the corresponding ESI mass spectrum recorded after 15 min of equilibration. Clearly, the two homodimers are the dominating species; 1_2^{2+} appears with *ca.* 35%,



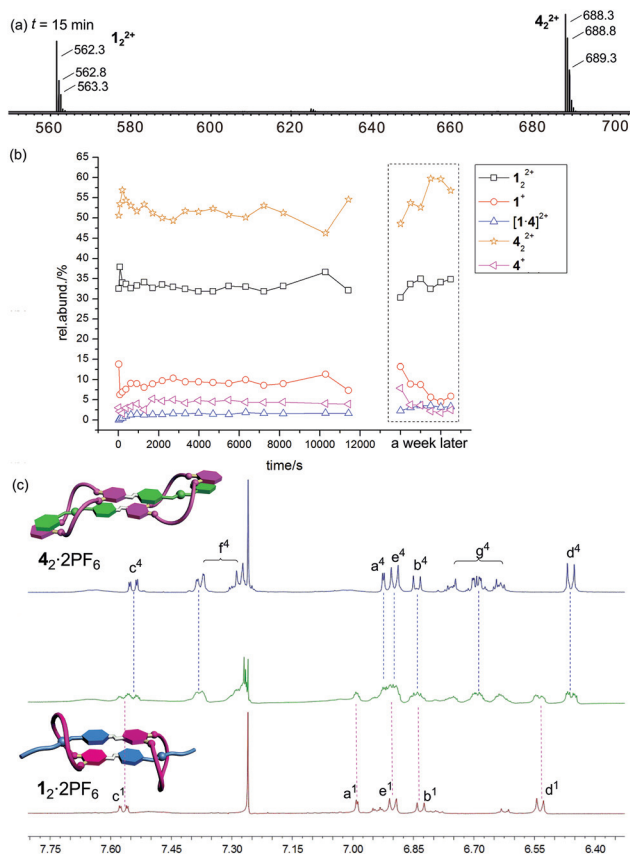


Fig. 5 (a) Partial ESI mass spectrum of a 1:1 mixture of 1_2-PF_6 and 4_2-PF_6 (100 μM for each monomer) in $\text{CHCl}_3\text{-CH}_3\text{CN}$ (2:1) recorded after 15 min of equilibration time. (b) More spectra have been recorded after different time intervals. No change is observed even after a week. (c) Partial ^1H NMR spectra (500 MHz, $\text{CDCl}_3\text{-CD}_3\text{CN}$ = 2:1, 10.0 mM, 298 K) of 1_2-PF_6 (bottom row), an equimolar mixture of 1_2-PF_6 and 4_2-PF_6 (centre), and 4_2-PF_6 (top row).

4_2^{2+} with *ca.* 55% intensity. The heterodimer $[1\cdot 4]^{2+}$ is always below 3%. No changes are found when the sample was left to equilibrate for periods as long as a week. This result first of all confirms that heterodimer formation is unfavourable as expected. The low intensity of the heterodimer also indicates that non-specific dimerization does not play a significant role under the experimental conditions applied. The differences of the intensities of the two homodimers point to some differences in the ESI response factors. Nevertheless, a semiquantitative picture is obtained.

The mass spectrometric results are in agreement with the ^1H NMR spectra in Fig. 5c. Although the spectrum of the mixture is somewhat less well-resolved, it is quite clear that this spectrum is a superposition of the two spectra of the homodimers. Full spectra and $^1\text{H}, ^1\text{H}$ COSY data are given in the ESI†. In conclusion, both methods are in agreement with the hypothesis that 1_2-PF_6 and 4_2-PF_6 do not form heterodimers.

A similar situation is observed for the 2:1 mixture of 2-PF_6 and 4_2-PF_6 that is equimolar in monomer concentration. The heterodimer appears in the mass spectra (ESI†) below 5%

intensity accompanied mainly by 2^+ and 4_2^{2+} . Due to the larger structural differences between a monomer (2^+) and a dimer (4_2^{2+}), larger differences in the ESI response factors can be expected. Indeed, the 2^+ ion is much lower in intensity than the 4_2^{2+} dimer. When judging the intensities, it must however be taken into account that they are proportional to the charge state in Fourier-transform ion-cyclotron-resonance (FTICR) mass spectrometry. Consequently, the 4_2^{2+} dication intensity is two times higher than that of a singly charged cation of the same abundance. The NMR spectra (ESI†) are again complex, but can be rationalized by a superposition of 2-PF_6 and 4_2-PF_6 .

Upon closer inspection, this result might be surprising. As 2-PF_6 does not form homodimers, an exchange of 2-PF_6 against a monomer in 4_2-PF_6 to yield the heterodimer would result in the same number of satisfied binding sites and should thus be thermoneutral. Consequently, one might have expected more singly threaded heterodimer and an exchange equilibrium in which 2-PF_6 , 4_2-PF_6 and $2\cdot 4\text{-PF}_6$ coexist. However, a doubly threaded, divalent dimer 4_2-PF_6 may well benefit from chelate cooperativity¹⁷ as observed recently for other divalent crown ammonium pseudorotaxanes.^{9h} The open, singly threaded dimer $2\cdot 4\text{-PF}_6$ does of course not profit from such a chelate cooperativity effect.

The $1_2\text{-PF}_6/3_2\text{-PF}_6$ (1:1) mixture

Mixing 1_2-PF_6 and 3_2-PF_6 in equimolar amounts leads indeed to the expected statistical mixture of 1_2-PF_6 , $1\cdot 3\text{-PF}_6$, and 3_2-PF_6 as indicated by the appearance of 1_2^{2+} , $[1\cdot 3]^{2+}$ and 3_2^{2+} in the ESI mass spectrum of the mixture (Fig. 6a). As anticipated, the homodimer dications are superimposed by signals of monomers 1^+ and 3^+ . The intensity ratio of the three dimeric dications is about 3:8:6 ($1_2^{2+}:[1\cdot 3]^{2+}:3_2^{2+}$). If one takes into account the likely differing desolvation energies, this ratio is more or less in agreement with a statistical formation of the three dimeric daisy chains.

The exchange of monomers between the two homodimers to yield the heterodimeric daisy chain is quite slow and can be followed by ESI mass spectrometry (Fig. 6b). If one takes the development of the heterodimer ion intensity as an example, one arrives at an estimate for the half-life of roughly 230 seconds. The exchange reaction thus proceeds on a minute time scale.

The ^1H NMR spectra are very complicated and difficult to interpret – in particular when it comes to quantification of the different dimers present in solution. However, quite clearly, the signals for H^b and H^c of the two compounds in the aromatic region provide evidence that the spectrum of the mixture is more than a mere superposition of the two homodimer spectra. Additional signals are observed, which together with the ESI mass spectrometric results at least provide qualitative evidence for the presence of the homo- and heterodimers in the sample solution.

The $2\text{-PF}_6/3_2\text{-PF}_6$ (2:1) mixture

Finally, the mixture of $2\text{-PF}_6/3_2\text{-PF}_6$ (equimolar in monomer concentration) is expected to self-sort into the heterodimer



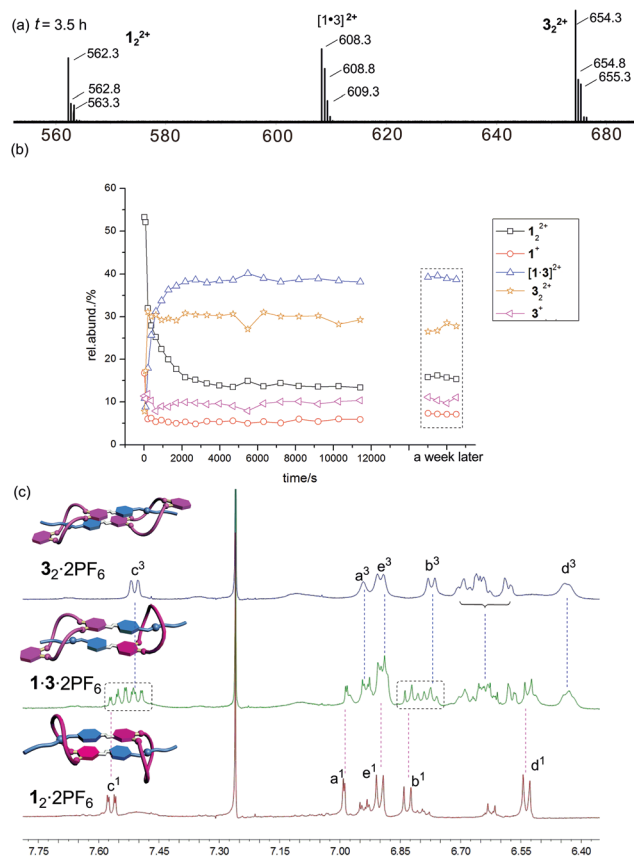


Fig. 6 (a) Partial ESI mass spectrum of a 1:1 mixture of $1_2:2PF_6$ and $3_2:2PF_6$ (100 μ M for each monomer) in $CHCl_3-CH_3CN$ (2:1) recorded after 3.5 h of equilibration time. (b) The equilibration can be followed over time and occurs on a minute time scale. (c) Partial 1H NMR spectra (500 MHz, $CDCl_3-CD_3CN = 2:1$, 10.0 mM, 298 K) of $1_2:2PF_6$ (bottom row), an equimolar mixture of $1_2:2PF_6$ and $3_2:2PF_6$ (centre), and $3_2:2PF_6$ (top row).

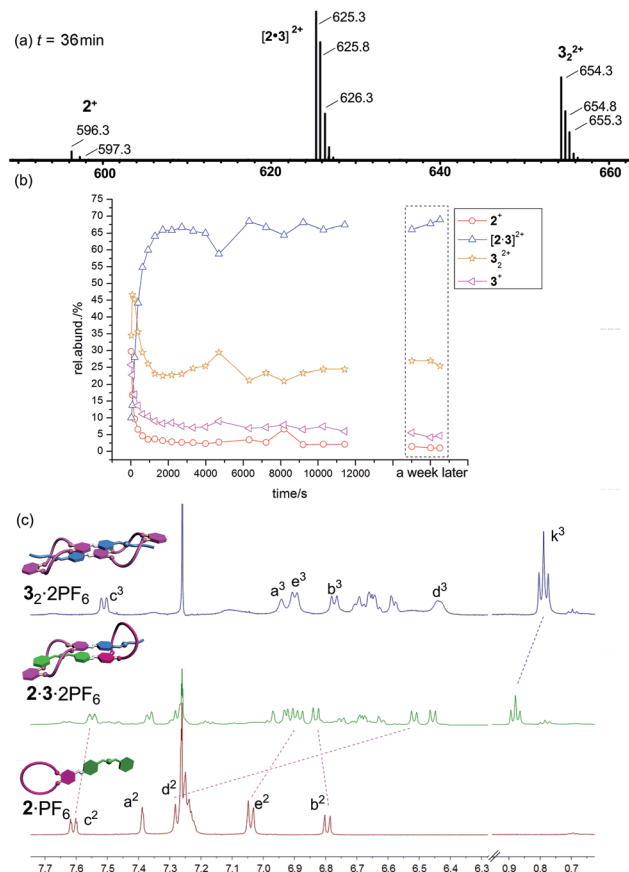


Fig. 7 (a) Partial ESI mass spectrum of a 2:1 mixture of $2-PF_6$ and $3_2:2PF_6$ (100 μ M for each monomer) in $CHCl_3-CH_3CN$ (2:1) recorded after 36 min of equilibration time. (b) The self-sorting can be followed over time and occurs on a minute time scale. (c) Partial 1H NMR spectra (500 MHz, $CDCl_3-CD_3CN = 2:1$, 10.0 mM, 298 K) of $2-PF_6$ (bottom row), a 2:1 mixture of $2-PF_6$ and $3_2:2PF_6$ (centre), and $3_2:2PF_6$ (top row).

exclusively. A look at the corresponding ESI mass spectrum (Fig. 7a) confirms the heterodimer to be the most prominent species in the sample solution which again forms on a minute time scale (Fig. 7b). As time passed by, we could see the decrease of the signal of 2^+ from 50% to less than 6% and the gradual increase of the intensity of $[2\cdot3]^{2+}$ (ESI $^+$). Similar ESI mass spectrometric behaviour (higher intensity) were found for 3_2^{2+} in Fig. 6 and 7, maybe due to its much different desolvation energy from those of other species. Nevertheless, some contribution from $2-PF_6$ and $3_2:2PF_6$ is still observed and one would arrive at the conclusion that self-sorting is imperfect.

A closer inspection of the NMR spectra of the individual compounds and the mixture of both, however, reveals that self-sorting is almost complete. The 1H NMR spectrum of the mixture exhibits new sets of signals that are not present in the individual parent compounds. In turn, the signals for $2-PF_6$ and $3_2:2PF_6$ are missing in the spectrum of the mixture as seen, for example, for c^3 , k^3 and e^2 (Fig. 7c). Such a divergence of the ESI mass spectrometric and NMR spectroscopic results has been observed earlier for other supramolecular self-sorting complexes.^{3d,e} As the NMR (10.0 mM) results reflect the

solution situation more directly than ESI mass spectrometry (100 μ M), we can therefore conclude that self-sorting quite nicely works for the $2-PF_6/3_2:2PF_6$ combination.

Experimental section

General

All reagents were commercially available and used as supplied without further purification. Benzo[21]crown-7 carboxylic acid **C7-COOH**, dibenzo[24]crown-8 carboxylic acid **C8-COOH**, monomer **1-PF₆**, and axle precursors **5** and **6** were prepared according to literature procedures.^{13,14} 1H NMR spectra were collected on a temperature-controlled 500 MHz Bruker AVANCE DMX-500 spectrometer with the deuterated solvent as the lock and the residual undeuterated solvent or TMS as the internal reference. Chemical shifts are reported in ppm and coupling constants in hertz (Hz). ^{13}C NMR spectra were recorded on a Bruker AVANCE DMX-500 spectrometer. The crystal structure data were collected on an Oxford Diffraction Xcalibur Gemini Ultra diffractometer with an Atlas detector.



The electrospray-ionization Fourier-transform ion-cyclotron-resonance (ESI-FTICR) mass spectrometric experiments were performed with a Varian/IonSpec QFT-7 FTICR mass spectrometer equipped with a superconducting 7 Tesla magnet and a Micromass Z-spray ESI ion source utilizing a stainless steel capillary with a 0.65 mm inner diameter. The solutions of samples (100 μM in $\text{CHCl}_3\text{-CH}_3\text{CN}$ 2 : 1) were introduced into the source with a syringe pump (Harvard Apparatus) at a flow rate of approximately $2\text{-}3\ \mu\text{L}\ \text{min}^{-1}$. The ions were collected in the instrument's accumulation hexapole long enough to obtain useful signal-to-noise ratios, introduced into the FTICR analyzer cell, which was operated at pressures below 10^{-9} mbar, and detected by a standard excitation and detection sequence. The ionization voltage was set to 3000–3500 V, source and probe temperature were 40 °C. The sample cone voltage was set to 25–30 V.

Syntheses

Monomer 2-PF₆. A solution of **C7-COOH** (170 mg, 0.42 mmol), **5** (133 mg, 0.42 mmol), EDC (178 mg, 0.90 mmol) and DMAP (24 mg, 0.20 mmol) in dichloromethane (10 mL) was stirred for 24 h at room temperature. The solvent was removed to give a crude product, which was subjected to column chromatography (ethyl acetate) to obtain the pure product. The product was dissolved in 10% HCl–ethyl acetate (10 mL) and stirred overnight. The white solid was filtered, washed with ethyl acetate thoroughly, and dissolved in warm deionized water (20 mL). A saturated aqueous solution of NH_4PF_6 was added to afford a white precipitate, which was filtered off and washed with deionized water to yield monomer 2-PF₆ as a white solid (188 mg, 60%), mp 117–120 °C. ¹H NMR (500 MHz, DMSO-*d*₆, 295 K) δ (ppm): 7.76 (d, *J* = 8.5 Hz, 1H), 7.60–7.52 (m, 3H), 7.51–7.37 (m, 5H), 7.33 (d, *J* = 8.3 Hz, 2H), 7.17 (d, *J* = 8.3 Hz, 1H), 4.24–4.20 (s, 2H), 4.20–4.15 (s, 2H), 4.15–4.07 (m, 4H), 3.82–3.75 (m, 4H), 3.65–3.59 (m, 4H), 3.59–3.54 (m, 4H), 3.54–3.48 (m, 8H). ¹³C NMR (125 MHz, acetone-*d*₆, 295 K) δ (ppm): 164.11, 153.05, 152.08, 148.06, 131.69, 131.48, 130.05, 129.45, 129.24, 129.01, 124.60, 122.52, 121.82, 113.59, 112.06, 70.07, 69.82, 69.65, 69.33, 69.19, 67.96, 51.93, 51.29. HRESIMS: *m/z* calcd for $[\text{M} - \text{PF}_6]^+$ C₃₀H₄₄NO₉, 562.3011; found 562.3032, error: 3.7 ppm.

Monomer 3-PF₆. A solution of **C8-COOH** (243 mg, 0.49 mmol), **6** (138 mg, 0.49 mmol), EDC (192 mg, 1.00 mmol) and DMAP (24 mg, 0.20 mmol) in dichloromethane (20 mL) was stirred for 24 h at room temperature. After workup as described above for 2-PF₆, monomer 3-PF₆ was obtained as a white solid (350 mg, 89%), mp 120–122 °C. ¹H NMR (500 MHz, DMSO-*d*₆, 295 K) δ (ppm): 8.75–8.63 (br, 2H), 7.75 (d, *J* = 8.6 Hz, 1H), 7.60–7.54 (m, 3H), 7.35 (d, *J* = 8.1 Hz, 2H), 7.15 (d, *J* = 8.6 Hz, 1H), 6.98–6.83 (m, 4H), 4.23–4.13 (m, 6H), 4.09–4.03 (m, 4H), 3.84–3.74 (m, 8H), 3.71–3.64 (m, 8H), 2.99–2.90 (m, 2H), 1.65–1.54 (m, 2H), 1.39–1.29 (m, 2H), 0.90 (t, *J* = 7.3 Hz, 3H). ¹³C NMR (125 MHz, DMSO-*d*₆, 295 K) δ (ppm): 164.63, 153.82, 151.63, 148.87, 148.45, 131.73, 130.07, 124.86, 122.82, 121.58, 121.18, 114.45, 113.01, 71.02, 70.95, 70.91, 70.88, 69.63, 69.60, 69.47, 69.31, 69.14, 49.95, 46.95,

27.88, 19.72, 13.94. HRESIMS: *m/z* calcd for $[\text{M} - \text{PF}_6]^+$ C₃₆H₄₈NO₁₀, 654.3273; found 654.3243, error 4.6 ppm.

Monomer 4-PF₆. A solution of **C8-COOH** (247 mg, 0.50 mmol), **5** (157 mg, 0.50 mmol), EDC (192 mg, 1.00 mmol) and DMAP (24 mg, 0.20 mmol) in dichloromethane (20 mL) was stirred for 36 h at room temperature. After the workup as described above, monomer 4-PF₆ was obtained as a white solid (300 mg, 72%), mp 103–105 °C. ¹H NMR (500 MHz, DMSO-*d*₆, 295 K) δ (ppm): 9.27–9.14 (br, 2H), 7.74 (d, *J* = 8.5 Hz, 1H), 7.59–7.53 (m, 3H), 7.53–7.39 (m, 5H), 7.34 (d, *J* = 8.0 Hz, 2H), 7.13 (d, *J* = 7.9 Hz, 1H), 6.96–6.82 (m, 4H), 4.27–4.09 (m, 8H), 4.08–4.00 (m, 4H), 3.84–3.72 (m, 8H), 3.71–3.60 (m, 8H). ¹³C NMR (125 MHz, DMSO-*d*₆, 295 K) δ (ppm): 164.66, 153.84, 151.68, 148.89, 148.47, 132.31, 131.89, 130.46, 129.92, 129.55, 129.21, 124.88, 122.80, 121.61, 121.21, 114.46, 113.03, 71.05, 70.98, 70.94, 70.91, 69.66, 69.63, 69.49, 69.34, 69.15, 50.73, 50.11. HRESIMS: *m/z* calcd for $[\text{M} - \text{PF}_6]^+$ C₃₉H₄₆NO₁₀, 688.3116; found 688.3108, error 1.2 ppm.

X-ray crystallographic data

Solid-state structure of 3₂-2PF₆. Block, colourless, C₃₆H₄₈F₆NO₁₀P, FW 799.73, triclinic, space group $P\bar{1}$, *a* = 11.9860(5), *b* = 12.2786(7), *c* = 16.2211(9) Å, α = 103.393(5)°, β = 110.555(5)°, γ = 92.801(4)°, *V* = 2152.18(19) Å³, *Z* = 2, *D_c* = 1.234 g cm⁻³, *T* = 140(2) K, μ = 0.140 mm⁻¹, 7859 measured reflections, 3825 independent reflections, 609 parameters, 1729 restraints, *F*(000) = 840, *R*₁ = 0.1665, *wR*₂ = 0.3278 (all data), *R*₁ = 0.0970, *wR*₂ = 0.2891 [*I* > 2 σ (*I*)], max. residual density 0.386 e Å⁻³, and goodness-of-fit (*F*²) = 1.090. CCDC number: 952285.

Solid-state structure of 4₂-2PF₆. Block, colourless, C₃₉H₄₆F₆NO₁₀P, FW 833.73, triclinic, space group $P2_1/n$, *a* = 10.4663(3), *b* = 42.4866(16), *c* = 25.4339(9) Å, α = 90.00°, β = 113.902(3)°, γ = 90.00°, *V* = 10 339.9(6) Å³, *Z* = 4, *D_c* = 1.070 g cm⁻³, *T* = 140(2) K, μ = 1.054 mm⁻¹, 13 857 measured reflections, 8431 independent reflections, 1040 parameters, 126 restraints, *F*(000) = 3480, *R*₁ = 0.1859, *wR*₂ = 0.4125 (all data), *R*₁ = 0.1501, *wR*₂ = 0.3841 [*I* > 2 σ (*I*)], max. residual density 1.172 e Å⁻³, and goodness-of-fit (*F*²) = 1.342. CCDC number: 981643.

Conclusions

The results obtained in this study confirm the expected self-sorting behaviour of daisy chain monomers 1-PF₆–4-PF₆. While the pairs 1₂-2PF₆/4₂-2PF₆ and 2-PF₆/4₂-2PF₆ do not form heterodimeric daisy chains, 1₂-2PF₆ and 3₂-2PF₆ yield a more or less statistical mixture. The 2-PF₆/3₂-2PF₆ pair instead forms almost exclusively the heterodimeric daisy chain. The present study thus provides evidence that hetero-[c₂]daisy chain pseudo-rotaxanes can be programmed based on the appropriate choice of orthogonal binding motifs. Both motifs used here are very similar and the self-sorting is based on the size complementarity of crown ether size and ammonium axle width.



It is, however, this similarity of the two binding motifs which also poses difficulties in the analysis of the solution situation and points to the limits of current routine characterization methods. While X-ray crystallography and NMR experiments are quite straightforward, when it comes to the characterization of the monomers and homodimers, the determination of solution concentrations in mixtures of them is not trivial – even though the molecules under study are structurally not very complicated.

Finally, a combination of ESI mass spectrometry and NMR spectroscopy provides semi-quantitative insight into the self-sorting behaviour as well as a rough estimate on the equilibration time scale. Nevertheless, to know the self-sorting behaviour is a key issue for future studies. For example, 2-PF₆ and 3-PF₆ can now be equipped with two different functional groups. They can then be brought into proximity to each other, when the heterodimer daisy chain forms – thus providing complexes with two functionalities which might be capable of cooperating with each other *e.g.* in organocatalysis. The two point attachment between the two monomers provides the basis for better geometrical control over the complex formed than a single-site recognition between the two partners. Furthermore, the exclusive formation of the heterodimer maximizes the concentration of complexes that bear both functional groups and therefore maximizes complex activity. The use of the crown ether/ammonium ion binding motifs renders the complexes switchable by (de)protonation. Consequently, the cooperation of the two functional groups could be switched on and off.

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Notes and references

- G. M. Whitesides, J. P. Mathias and C. T. Seto, *Science*, 1991, **254**, 1312–1319.
- For reviews, see: (a) M. M. Safont-Sempere, G. Fernández and F. Würthner, *Chem. Rev.*, 2011, **111**, 5784–5814; (b) M. L. Saha and M. Schmittel, *Org. Biomol. Chem.*, 2012, **10**, 4651–4684.
- Selected examples: (a) Y. Ma, S. V. Kolotuchin and S. C. Zimmerman, *J. Am. Chem. Soc.*, 2002, **124**, 13757–13769; (b) A. Wu and L. Isaacs, *J. Am. Chem. Soc.*, 2003, **125**, 4831–4835; (c) E. S. Barrett, T. J. Dale and J. Rebek Jr., *J. Am. Chem. Soc.*, 2008, **130**, 2344–2350; (d) Y. Rudzevich, V. Rudzevich, F. Klautzsch, C. A. Schalley and V. Böhmer, *Angew. Chem., Int. Ed.*, 2009, **48**, 3867–3871; (e) C. Talotta, C. Gaeta, Z. Qi, C. A. Schalley and P. Neri, *Angew. Chem., Int. Ed.*, 2013, **52**, 7437–7441.
- Selected examples: (a) P. N. Taylor and H. L. Anderson, *J. Am. Chem. Soc.*, 1999, **121**, 11538–11545; (b) J.-M. Lehn, *Science*, 2002, **295**, 2400–2403; (c) I.-W. Hwang, T. Kamada, T. K. Ahn, D. M. Ko, T. Nakamura, A. Tsuda, A. Osuka and D. Kim, *J. Am. Chem. Soc.*, 2004, **126**, 16187–16198; (d) K.-W. Chi, C. Addicott, A. M. Arif and P. J. Stang, *J. Am. Chem. Soc.*, 2004, **126**, 16569–16574; (e) B. H. Northrop, Y.-R. Zheng, K.-W. Chi and P. J. Stang, *Acc. Chem. Res.*, 2009, **42**, 1554–1563; (f) Y. R. Zheng, H. B. Yang, K. Ghosh, L. Zhao and P. J. Stang, *Chem. – Eur. J.*, 2009, **15**, 7203–7214; (g) S. Ulrich and J.-M. Lehn, *J. Am. Chem. Soc.*, 2009, **131**, 5546–5549; (h) S. Ulrich and J.-M. Lehn, *Chem. – Eur. J.*, 2009, **15**, 5640–5645; (i) K. Mahata and M. Schmittel, *J. Am. Chem. Soc.*, 2009, **131**, 16544–16554; (j) T. Murase, S. Horiuchi and M. Fujita, *J. Am. Chem. Soc.*, 2010, **132**, 7864–7865; (k) Y. Yamauchi, M. Yoshizawa, M. Akita and M. Fujita, *J. Am. Chem. Soc.*, 2010, **132**, 960–966; (l) K. Mahata, M. L. Saha and M. Schmittel, *J. Am. Chem. Soc.*, 2010, **132**, 15933–15935; (m) M. Schmittel and K. Mahata, *Chem. Commun.*, 2010, **46**, 4163–4165; (n) M. M. J. Smulders, A. Jiménez and J. R. Nitschke, *Angew. Chem., Int. Ed.*, 2012, **51**, 6681–6685; (o) A. S. Singh and S.-S. Sun, *Chem. Commun.*, 2012, **48**, 7392–7394; (p) M. L. Saha, S. Pramanik and M. Schmittel, *Chem. Commun.*, 2012, **48**, 9459–9461.
- Selected examples: (a) A. D. Shaller, W. Wang, H. Gan and A. D. Q. Li, *Angew. Chem., Int. Ed.*, 2008, **47**, 7705–7709; (b) S. Ghosh, A. Wu, J. C. Fettinger, P. Y. Zavalij and L. Isaacs, *J. Org. Chem.*, 2008, **73**, 5915–5925; (c) S. Ghosh, X.-Q. Li, V. Stepanenko and F. Würthner, *Chem. – Eur. J.*, 2008, **14**, 11343–11357; (d) M. M. Safont-Sempere, P. Osswald, M. Stolte, M. Grüne, M. Renz, M. Kaupp, K. Radacki, H. Braunschweig and F. Würthner, *J. Am. Chem. Soc.*, 2011, **133**, 9580–9591.
- Selected examples: (a) N. Tomimasu, A. Kanaya, Y. Takashima, H. Yamaguchi and A. Harada, *J. Am. Chem. Soc.*, 2009, **131**, 12339–12343; (b) W. Jiang, Q. Wang, I. Linder, F. Klautzsch and C. A. Schalley, *Chem. – Eur. J.*, 2011, **17**, 2344–2348.
- (a) P. Mukhopadhyay, P. Y. Zavalij and L. Isaacs, *J. Am. Chem. Soc.*, 2006, **128**, 14093–14102; (b) P. Mukhopadhyay, A. Wu and L. Isaacs, *J. Org. Chem.*, 2004, **69**, 6157–6164.
- (a) P. R. Ashton, E. J. T. Chrystal, P. T. Glink, S. Menzer, C. Schiavo, N. Spencer, J. F. Stoddart, P. A. Tasker, A. J. P. White and D. J. Williams, *Chem. – Eur. J.*, 1996, **2**, 709–728; (b) N. Yamaguchi, L. M. Hamilton and H. W. Gibson, *Angew. Chem., Int. Ed.*, 1998, **37**, 3275–3279; (c) P. R. Ashton, I. Baxter, M. C. T. Fyfe, F. M. Raymo, N. Spencer, J. F. Stoddart, A. J. P. White and D. J. Williams, *J. Am. Chem. Soc.*, 1998, **120**, 2297–2307; (d) T. Clifford, A. Abushamleh and D. H. Busch, *Proc. Natl. Acad. Sci. U. S. A.*, 2002, **99**, 4830–4836; (e) F. Huang, K. A. Switek, L. N. Zakharov, F. R. Fronczek, C. Sleboznick, M. Lam,



- J. A. Golen, W. S. Bryant, P. E. Mason, A. L. Rheingold, M. Ashraf-Khorassani and H. W. Gibson, *J. Org. Chem.*, 2005, **70**, 3231–3241; (f) Z. Niu, F. Huang and H. W. Gibson, *J. Am. Chem. Soc.*, 2011, **133**, 2836–2839.
- 9 (a) F. Wang, C. Han, C. He, Q. Zhou, J. Zhang, C. Wang, N. Li and F. Huang, *J. Am. Chem. Soc.*, 2008, **130**, 11254–11255; (b) W. Jiang, H. D. F. Winkler and C. A. Schalley, *J. Am. Chem. Soc.*, 2008, **130**, 13852–13853; (c) W. Jiang and C. A. Schalley, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 10425–10429; (d) F. Wang, B. Zheng, K. Zhu, Q. Zhou, C. Zhai, S. Li, N. Li and F. Huang, *Chem. Commun.*, 2009, 4375–4377; (e) W. Jiang, A. Schäfer, P. C. Mohr and C. A. Schalley, *J. Am. Chem. Soc.*, 2010, **132**, 2309–2320; (f) W. Jiang and C. A. Schalley, *J. Mass Spectrom.*, 2010, **45**, 788–798; (g) W. Jiang, D. Sattler, K. Rissanen and C. A. Schalley, *Org. Lett.*, 2011, **13**, 4502–4505; (h) W. Jiang, K. Nowosinski, N. L. Löw, E. V. Dzyuba, F. Klautzsch, A. Schäfer, J. Huuskonen, K. Rissanen and C. A. Schalley, *J. Am. Chem. Soc.*, 2012, **134**, 1860–1868; (i) S. Dong, B. Zheng, M. Zhang, X. Yan, X. Ding, Y. Yu and F. Huang, *Macromolecules*, 2012, **45**, 9070–9075; (j) S. Dong, X. Yan, B. Zheng, J. Chen, X. Ding, Y. Yu, D. Xu, M. Zhang and F. Huang, *Chem. – Eur. J.*, 2012, **18**, 4195–4199.
- 10 (a) P. R. Ashton, P. J. Campbell, P. T. Glink, D. Philp, N. Spencer, J. F. Stoddart, E. J. T. Chrystal, S. Menzer, D. J. Williams and P. A. Tasker, *Angew. Chem., Int. Ed. Engl.*, 1995, **34**, 1865–1869; (b) C. Zhang, S. Li, J. Zhang, K. Zhu, N. Li and F. Huang, *Org. Lett.*, 2007, **9**, 5553–5556.
- 11 Reviews: (a) L. Fang, M. A. Olson, D. Benitez, E. Tkatchouk, W. A. Goddard III and J. F. Stoddart, *Chem. Soc. Rev.*, 2010, **39**, 17–29; (b) J. Rotzler and M. Mayor, *Chem. Soc. Rev.*, 2013, **42**, 44–62; (c) B. Zheng, F. Wang, S. Dong and F. Huang, *Chem. Soc. Rev.*, 2012, **41**, 1621–1636. Selected examples: (d) P. R. Ashton, I. Baxter, S. J. Cantrill, M. C. T. Fyfe, P. T. Glink, J. F. Stoddart, A. J. P. White and D. J. Williams, *Angew. Chem., Int. Ed.*, 1998, **37**, 1294–1297; (e) S. J. Cantrill, G. J. Youn, J. F. Stoddart and D. J. Williams, *J. Org. Chem.*, 2001, **66**, 6857–6872; (f) D. G. Amirsakis, A. M. Elizarov, M. A. Garcia-Garibay, P. T. Glink, J. F. Stoddart, A. J. P. White and D. J. Williams, *Angew. Chem., Int. Ed.*, 2003, **42**, 1126–1132; (g) M. P. L. Werts, M. van den Boogaard, G. M. Tsiygoulis and G. Hadziioannou, *Macromolecules*, 2003, **36**, 7004–7013; (h) H. W. Gibson, N. Yamaguchi, Z. Niu, J. W. Jones, C. Slebodnick, A. L. Rheingold and L. N. Zakharov, *J. Polym. Sci., Part A: Polym. Chem.*, 2010, **48**, 975–985; (i) L. Cao and L. Isaacs, *Org. Lett.*, 2012, **14**, 3072–3075; (j) Z. Zhang, G. Yu, C. Han, J. Liu, X. Ding, Y. Yu and F. Huang, *Org. Lett.*, 2011, **13**, 4818–4821; (k) Z. Zhang, C. Han, G. Yu and F. Huang, *Chem. Sci.*, 2012, **3**, 3026–3031.
- 12 Review: (a) J.-P. Collin, C. Dietrich-Buchecker, P. Gaviña, M. C. Jimenez-Molero and J.-P. Sauvage, *Acc. Chem. Res.*, 2001, **34**, 477–487; examples: (b) C. Romuald, E. Busseron and F. Coutrot, *J. Org. Chem.*, 2010, **75**, 6516–6531; (c) G. Du, E. Moulin, N. Jouault, E. Buhler and N. Giuseppone, *Angew. Chem., Int. Ed.*, 2012, **51**, 12504–12508; (d) X. Yan, B. Zheng and F. Huang, *Polym. Chem.*, 2013, **4**, 2395–2399.
- 13 B. Zheng, M. Zhang, S. Dong, J. Liu and F. Huang, *Org. Lett.*, 2012, **14**, 306–309.
- 14 (a) D.-J. Feng, X.-Q. Li, X.-Z. Wang, X.-K. Jiang and Z.-T. Li, *Tetrahedron*, 2004, **60**, 6137–6144; (b) L. Wu, Y.-M. He and Q.-H. Fan, *Adv. Synth. Catal.*, 2011, **353**, 2915–2919.
- 15 S. Di Stefano, *J. Phys. Org. Chem.*, 2010, **23**, 797–805.
- 16 E. Leize, A. Jaffrezic and A. Van Dorselaer, *J. Mass Spectrom.*, 1996, **31**, 537–544.
- 17 For reviews on multivalency, see: (a) M. Mammen, S. K. Choi and G. M. Whitesides, *Angew. Chem., Int. Ed.*, 1998, **37**, 2754–2794; (b) A. Mulder, J. Huskens and D. N. Reinhoudt, *Org. Biomol. Chem.*, 2004, **2**, 3409–3424; (c) J. D. Badjic, A. Nelson, S. J. Cantrill, W. B. Turnbull and J. F. Stoddart, *Acc. Chem. Res.*, 2005, **38**, 723–732; (d) C. Fasting, C. A. Schalley, M. Weber, O. Seitz, S. Hecht, B. Koksche, J. Dornedde, C. Graf, E.-W. Knapp and R. Haag, *Angew. Chem., Int. Ed.*, 2012, **51**, 10472–10498. For reviews on the thermodynamic analysis of chelate cooperativity in supramolecular complexes, see: (e) G. Ercolani, C. Piguet, M. Borkovec and J. Hamacek, *J. Phys. Chem. B*, 2007, **111**, 12195–12203; (f) C. A. Hunter and H. L. Anderson, *Angew. Chem., Int. Ed.*, 2009, **48**, 7488–7499; (g) G. Ercolani and L. Schiaffino, *Angew. Chem., Int. Ed.*, 2011, **50**, 1762–1768.

