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Cation binding resorcinarene bis-crowns: The effect of lower rim alkyl chain length to crystal packing and solid lipid nanoparticles

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A group of seven resorcinarene bis-crown ethers (CNBC5) with two polyether bridges at the upper rim and either propyl, butyl, pentyl, heptyl, nonyl, decyl or undecyl groups at the lower rim were synthesized and their binding properties with Cs^+ were investigated by NMR titration. The bis-crowns form 1:2

10 complexes with Cs^+ with binding constants of log K 4–5. Crystal structures of bis-crowns and their Cs^+ and K^+ complexes were studied and different packing motifs were found depending on the alkyl chain length. Short ethyl, propyl and butyl alkyl chains gave a layer or pillar packing where the polar and non-polar regions cannot be distinguished, whereas, longer pentyl and decyl chains formed bilayers. Amphiphilic properties and self-assembly in water was studied by preparing solid lipid nanoparticles

15 (SLN) from the bis-crowns. All investigated compounds formed stable SLNs showing amphiphilic character, which in the case of the short chain bis-crowns probably rises from their locked boat conformation separating the polar face of the molecule from the non-polar face.

Introduction

Combining host-guest chemistry and surfactant properties into a
20 single molecule by structural design of supramolecules, nanoscale materials, i.e. films, particles or gels with host-guest functionality can be obtained. Calixarenes and resorcinarenes are macrocyclic supramolecular hosts well suited for this task since they have a concave binding cavity capable of binding various guest
25 molecules or ions.[1-3] Resorcinarenes can be easily converted into amphiphilic molecules using long aliphatic aldehydes in their synthesis resulting in hydrophobic chains below the binding site. The upper rim of the resorcinarene bowl has innate hydrophilic character because of the OH-functionalities derived from the
30 resorcinol, and in addition, the upper rim is readily available for further functionalization to improve the binding affinity and selectivity. The self-assembly of amphiphilic calixarenes and resorcinarenes in water and at interfaces into mono/bilayers, thin films, vesicles and micelles, and properties of these assemblies
35 have been studied avidly to extend the use of calixarenes.[2, 4] Some recent examples of their potential applications include gene delivery,[5, 6] catalytic activity,[7] liquid crystals[8] and VOC sensing.[9] In addition to host-guest properties, environmentally responsive functionalities, which change the organized structures from
40 micelles into larger vesicle according to pH, have been

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45 † Electronic Supplementary Information (ESI) available: Synthetic procedures, SEM images for SLN and details of crystal structue refinement. See DOI: 10.1039/b000000x/

prepared.[10-12]

50 Solid lipid nanoparticles (SLN), or particles prepared from solid lipids, are the latest addition to the family of drug carrier structures since the introduction of liposomes and polymeric nanoparticles, which are prepared from liquid lipids.[13] The preparation process of SLNs leads to particles with diameters
55 from tens to few hundred or thousand nanometers, which can be loaded with drugs or other sensitive compounds and used for their protection and transport. Since calixarenes and resorcinarenes are usually solid materials at room temperature, they can be used for preparation of solid lipid nanoparticles and studies of their
60 potential in encapsulation of biologically important guests, DNA for cell transfection and surface modification for drug targeting have been published.[14-17]

Calixarenes and resorcinarenes with crown ether bridges connecting the hydroxyl groups are very selective cation
65 receptors called calixcrowns.[18] Depending on the number of oxygen donors and thus the length and geometry of the crown bridge, calixcrowns have very good affinity towards alkali and alkaline earth metal cations and ammonium ions.[19] Resorcinarene bis-crowns and their K^+, Cs^+, Rb^+ and Ag^+ complexes have
70 shown very interesting structural properties such as formation of layers, capsules and nanorods.[20-22] Therefore, we have been interested in studying the effect of alkyl chain length in the crystal packing of the resorcinarene bis-crowns and their metal complexes, where it can (a) influence the twisting of the
75 resorcinarene framework and (b) induce a layer or bilayer packing when the hydrophobic effect of the alkyl chains becomes strong enough. The alkyl chain length also affects the amphiphilic properties and self-assembly in water, which was studied by

preparing SLNs out of series of resorcinarene bis-crowns with short, medium and long alkyl chains.

Results

Synthesis and complexation studies

A group of seven resorcinarene bis-crown ethers or CNBC5, where N denotes to the number of carbons at the lower rim alkyl group and 5 to the number of oxygen donors in each polyether bridge, were prepared by O-alkylation of the free hydroxyl groups of various tetramethoxy resorcinarenes[24] (Fig. 1). Propyl, butyl, pentyl, heptyl, nonyl, decyl and undecyl groups at the resorcinarene lower rim were chosen for structural comparison with the previously synthesized C2BC5[21,22] and to create amphiphilic bis-crowns. Synthesis was carried out in dry dimethyl formamide (DMF) using Cs_2CO_3 and ditosylated tetra(ethylene glycol) yielding tetramethoxy resorcinarene bis-crown ethers after purification as 15–30 % yields.

Complexation of the bis-crowns with an alkali metal cation, cesium hexafluorophosphate, was carried out using NMR titration in order to investigate if the lower rim alkyl chain length has an effect on the binding affinity. C3BC5, C5BC5, C9BC5 and C11BC5 bind Cs^+ with the affinity of log K_{11} 1.0–2.2 and total binding constant of log $K_{11}K_{12}$ 4.0–5.0 (Table 1). C2BC5 has binding constant of log K_{11} 1.75 for 1:1 complex,[21] which falls at the same magnitude of order as now determined log K_{11} values. The second binding constant log K_{12} is larger than the first binding constant for all investigated complexes. In case of C2BC5, K_{12} was not determined because Job plot showed that intrinsic water concentration over 1 molar equivalent relative to the host leads to 1:1 complexation. For C3BC5–C11BC5 such a strong trend was not observed, water content being 1–2.5 molar equivalents except for the 13 mol. eq. for C11BC5. In all cases a 1:2 binding model gave better fits than the 1:1 model.

Crystal structures

Resorcinarene bis-crowns

Single crystals of resorcinarene bis-crowns were grown by slow evaporation from alcohol solutions. C4BC5 (structure **C4**) and C5BC5 (**C5**) crystallized in a triclinic P-1 without any solvent in their binding cavity or in the crystal lattice. Analysis of the conformational properties of individual molecules (Table 2)

revealed that the bis-crowns are in a boat or slightly twisted boat conformation and upright aryl rings (A and C) are tilted towards the cavity with -8.3– -10.7 dihedral angles. Crown ether bridges are folded on top of the binding cavities closing the space inside.

Table 1. Binding constants for Cs^+ complexes in acetone-D6.[a]

	C3BC5	C5BC5	C9BC5	C11BC5
log K_{11}[b]	1.04 ± 0.18	1.59 ± 0.41	1.49 ± 0.60	2.21 ± 0.09
log $K_{11}K_{12}$[c]	4.43 ± 0.03	4.62 ± 0.10	4.06 ± 0.19	5.03 ± 0.04
log K_{12}[d]	3.39	3.03	2.57	2.82

[a] NMR titration at 30 °C, R-values <7 %. [b] Binding constant for reaction H+G ↔ HG. [c] Total binding constant for H+2G ↔ HG2. [d] K_{12}= $K_{11}K_{12}/K_{11}$.

The crystal packing of **C5** can be described as "squeezed bilayer" where the upper rim interface of two opposing rows appears as if compressed into one layer (Fig. 2). However, the polar and non-polar layers can still be distinguished. Rows are aligned parallel to A/C aryl plane direction (later A/C direction, Fig. 1). The clockwise (*cw*) counterclockwise (*ccw*) enantiomers of the bis-crowns are related by inversion symmetry. In **C5**, *cw* and *ccw* enantiomers alternate in each bilayer in such a way that each *cw* is facing up and is surrounded by a *ccw* facing down on both sides. **C4** has two molecules in the asymmetric unit, *I* and *II*, which could be assigned either to a boat or a slightly twisted boat conformation. Molecules pack in a pillar assembly with alternating *I* and *II* molecules (Fig. 3). Each pillar consists of either *cw* enantiomers or *ccw* enantiomers, which in turn form layers of *cw* and *ccw* enantiomers, but separation into polar and non-polar regions does not occur.

Long chain bis-crowns C9BC5, C10BC5 and C11BC5 crystallized readily from ethanol in a monoclinic lattice (*Z*=8). Interestingly, most of these structures showed straight alignment of the alkyl groups in the bilayer assembly without disorder but had unresolvable disorder at the crown ether bridges and therefore these structures can only be considered as preliminary structures.[*]

Alkali metal complexes

Alkali metal complexes of the bis-crowns were studied by crystallizing CNBC5s with excess of potassium or cesium hexafluorophosphate in alcohols yielding 1:2 (host-guest) complexes. When the binding pockets of the host are filled with

Table 2 Conformational properties of the resorcinarene bis-crowns and their alkali metal complexes.

	C4[†]		C5	C2K2	C3Cs2	C5K2	C10Cs2
	I	*II*					
Crystal packing arrangement	*pillar*	*pillar*	*squeezed bilayer*	*layer*	*layer / shifted capsule*	*bilayer / shifted capsule*	*bilayer / shifted capsule*
Conformation	*boat*	*boat*	*twisted boat*	*boat*	*twisted boat*	*boat*	*boat*
Tilt/°	3.7	5.7	8.9	1.4	7.0	3.3	1.1
Twist/°	3.9	5.7	8.7	2.1	6.4	3.5	0.5
Distance/Å[a]	4.79 / 8.01	4.87 / 8.00	4.80 / 7.98	5.28 / 7.98	5.46 / 7.95	5.26 / 7.98	5.49 / 7.94
Dihedral angle between opposite rings/°[b]	-10.7 / 175.2	-8.3 / 173.3	-10.3 / 167.3	14.4 / 151.4	22.2 / 148.8	11.8 / 147.0	21.1 / 148.25
Dihedral angle against methine plane/°[c]	80.0 / 89.1, 175.6/176.5	88.9 / 82.9, 176.2/176.3	86.4 / 83.2, 173.2/174.0	96.5 / 97.9, 165.9/165.5	100.3 / 101.5, 168.3/160.5	96.4 / 95.3, 166.5/160.5	100.6 /100.5, 163.2 /165.1
Cavity diameter/Å[d]	5.17 / 5.11	5.26 / 5.21	4.59 / 4.79	4.80 / 5.15	5.42 / 5.28	4.90 / 4.69	5.08 / 5.23

[†] **C4** has two molecules in the asymmetric unit. [a] Between opposite aromatic ring centroids. [b] Between opposite aromatic ring planes A and C, B and D. [c] Aromatic rings plane A,C,B,D against methine plane C7-C14-C21-C28. [d] Average cavity diameter measured as O–O distances.

cations the dihedral angles of A/C aryl rings are 11.8–22.2° and cavity diameters between 4.69–5.23 Å. The binding pocket of the bis-crowns is flexible and able to adjust its size slightly according to the size of the guest cation giving smaller cavity diameters for K^+ than Cs^+ complexes. Cations bind to the host with cation–π (η^6) interaction with a centroid-cation distance of 3.0–3.1 Å for K^+ and 3.2–3.3 for Cs^+, and with M^+–O interactions with methoxy group oxygens and 3–4 coordination bonds to the bridge (O–M^+ 2.70–3.54 Å). One of the PF_6^- anions is coordinated between the two cations inside the binding pocket, which helps to reduce the charge repulsion between the cations. The other anion is located outside the cavity and creates short contacts between the complexes. Depending on the structure, alkyl chain length and the cation, the complexes pack in shifted capsule, layer or bilayer assemblies.

C2BC5 has been previously crystallized as KPF_6 complex in a capsule assembly.[20] Now, another packing for C2BC5•2KPF_6 complex (**C2K2**) was obtained, forming layered packing without the capsule formation. When viewed on top, each *cw* enantiomer alternates with *ccw* enantiomers within a layer, and therefore polar and non-polar sides cannot be distinguished. A side view (B/D direction) of the packing reveals that *cw* and *ccw* enantiomers are separated on their own stacks in a parallel alignment (Fig. 4).

C3BC5•2$CsPF_6$ (**C3Cs2**) crystallized with ethanol as a solvate in the crystal lattice. The conformation of the host is a twisted boat in contrast to the boat conformation of all the other complexes, which can be understood by analyzing the anion/solvent coordination of the cation. One of the PF_6^- anions is coordinated between the two Cs^+ inside the cavity. In addition, a water molecule, not found in the other structures, is coordinated to Cs2 with 3.13 Å Cs–O distance and has a short contact of 2.99 Å to the F10A of the (disordered) second PF_6^-, which in turn is located close to the crown ether bridge of the opposite enantiomer with F8A–C59 distance of 3.12 Å (Fig 5). A pair of complexes, *cw* and *ccw* enantiomers, form a shifted capsule connected by the solvent-anion contacts. The top view of the complexes shows similar alternating pattern of *cw* and *ccw* enantiomers as in **C2K2**, and side view from the B/D direction shows a layered packing, where shifted capsules form diagonal lines through the crystal. The role of the ethanol solvate is to fill the voids at the B/D edges of the complexes, where they form H-bonded circles of four ethanol molecules without connecting to the host-guest complex.

C5BC5•2KPF_6 complex (**C5K2**) forms layers of single *cw* or *ccw* enantiomer which consist of rows aligned in the A/C direction with 2.43 Å shift between the molecules (15 % of complex width) and in the B/D direction with a 3.28 Å (35 %) shift. In contrast to the **C5,** all alkyl chains are oriented straight below the methine plane forming a clear bilayer packing. The upper rim interface of **C5K2** forms shifted capsules with a 2.80 Å dislocation of the B/D planes accounting for 30 % of the width of a molecule (Fig. 6A). Similar packing was obtained for the C10BC5•2$CsPF_6$ complex (**C10Cs2**) despite the difference in cation size and thus larger cavity diameter, but the longer alkyl groups expand the thickness of the bilayer up to 27.39 Å (Fig. 6B).

Solid lipid nanoparticles

The ability of CNBC5's to form solid lipid nanoparticles was tested to assess their self-assembling properties in water. Previously, calixarenes and resorcinarenes bearing long alkyl chains and hydrophilic functionalities at the upper rim have been used to prepare stable SLN's by solvent diffusion (solvent replacement) method.[25, 26] The same method was applied for the CNBC5s, where approximately 5–7 mg of CNBC5 was dissolved in a small amount of THF and water was added to the solution by vigorous stirring, after which a cloudy suspension was formed. The size of the SLN's was analyzed using dynamic light scattering, which gave hydrodynamic diameters of 220–320 nm for the particles with polydispersity indexes of 0.04–0.34. The particle shape and size was confirmed by SEM images, which revealed spherical particles at a size distribution corresponding to the DLS measurements (Fig. 7). For calixarenes and resorcinarenes, it has been discovered that the size of the SLNs is affected by several parameters: THF/water ratio, stirring speed, pH of the solution, and length of the alkyl chains of the calixarene.[25, 26] However, changes on the particle size are mostly affected by the final concentration of the calixarene in the suspension. Therefore in this study, other parameters except the length of the alkyl chains and final concentration of the resorcinarene suspensions were kept constant. In the first series (Fig. 8) the molar concentration of the bis-crowns was constant and the diameters of the particles increase when the amount of carbon atoms at the alkyl chains increase. In the second series SLNs were prepared keeping the mg/L concentration constant to make sure that the increased particle diameter was not originating from the increased amount (in milligrams) of bis-crown in the suspensions. The second series (Fig. 8) has very similar particle sizes than the first one, which indicates that the change in the amount of bis-crowns between the two series has negligible effect. The variation in the particle size, although quite modest, is most likely caused by the different alkyl chain length of the bis-crowns, which affects their amphiphilic properties.

Discussion

Resorcinarene bis-crowns were shown to bind Cs^+ as 1:1 and 1:2 complexes in solution, 1:2 being the dominant species. There is some variation especially between the log K_{11} values, giving 1.04 for C3BC5 and 3.31 for C11BC5. However, the higher log K_{12} value for C3BC5 partly compensates this difference when the total binding constants are examined. In contrast to the previously examined C2BC5, all compounds gave 1:2 complexes when more than 1 molar equivalent of water relative to the host was present in the solution. Therefore, what at first glance appears to be the effect of the alkyl chain length, may well be the indirect result of desolvation of the cation. The sensitivity of the measurement towards water may explain some of the observed differences in the binding constants between the experiments.

Since the affinity of C2BC5 towards K^+ is very low, log K of 0.23 for the 1:1 complex,[21] the binding constant was not determined for the other bis-crown potassium complexes. However, the structural properties of solid state K^+ complexes were compared to the Cs^+ complexes with the purpose of exploring alternative crystal packing forms due to different cation size. Based on the results it seems that the size of the cation does not have a direct influence on the packing, since similar

structures were obtained for the different cations (**C2K2** and C2BC5•CsPF₆ complex;[21] **C5K2** and **C10Cs2**). Instead, solvent coordination together with the cation size has more important role in the packing, which is seen by comparing the **C2K2** and **C3Cs2** and a capsule structure of C2BC5•2KPF₆[20] with a water molecule coordinated inside the cavity. **C3Cs2** also contains water, which is involved in the shifted capsule coordination. The larger diameter of Cs⁺ probably prevents similar coordination of water and the tilted angle of PF₆⁻ between the cations inside the binding pocket as in C2BC5•2KPF₆ capsules, and now coordination happens outside the binding pocket.

Twisting of the resorcinarene skeleton was observed in **C4**, **C5** and **C3Cs2**, and is therefore not limited to short alkyl chain bis-crowns. Rather, all interaction in the lattice determine the conformation of the resorcinarene to provide optimal close packing.

The effect of the lower rim alkyl chain length is connected to the amphiphilic nature of the bis-crowns. When alkyl chains are 2–4 carbons long, they have not been found to form bilayer packing with separated polar and non-polar parts. Instead, layers with alternating upper and lower rims in neighboring molecules or complexes are seen, and in addition, **C4** formed a pillar type assembly with tilted methine carbon planes. For C5BC5 a squeezed bilayer in **C5** and a bilayer in **C5K2** were found, which shows that five carbons is a limiting alkyl chain length for the bilayer type packing. For the long chain bis-crowns increased molecule size made crystallization more difficult and the structures also tend to show more disorder in either at the lower rim alkyl chains or at the crown ether bridges or both. Since C9BC5–C11BC5 had a very bad unresolvable disorder at the upper rim polyether bridges but quite well organized alkyl groups at the lower rim, this can be interpreted as an indication of stronger hydrophobic interactions at the lower rim, which drives the alkyl chains in the ordered packing whereas, the crown ether bridges have more conformational freedom. As a result, complexation of cations in the binding pocket was attempted in order to rigidify the crown ether bridges and, thus, enable the crystal structure analysis of these molecules. This strategy has so far provided the structure of Cs⁺ complex of C10BC5 which has very similar packing compared to **C5K2**.

The second goal of this study was to investigate the self-assembly of CNBC5s in water by preparing SLNs. Bis-crowns are neutral amphiphilic molecules without H-bond donors at their polar part. It is noteworthy, that also bis-crowns with C_2, C_3 and C_4 alkyl chains, which do not arrange in a bilayered packing, indicating their amphiphilic character in the solid state, form these particles. In addition, SLNs with shorter than C_4 alkyl or acyl chains have not been previously reported for calixarenes or resorcinarenes to the best of our knowledge. The ability of these molecules to form SLNs probably arises from the macrocyclic effect and the locked boat conformation, which separates the hydrophilic side of the compound from the hydrophobic side.

The diameter of the SLNs increased for the longer alkyl chains, which is in fact opposite result to those Coleman *et al.*[27] have obtained for the *para*-acyl-calix[9]arene SLNs. *para*-Acyl-calix[4]arenes on the other hand have not shown clear trend in the SLN size for different alkyl chain lengths.[25] Calix[9]arenes have larger, nine membered macrocyclic rings with enhanced conformational flexibility compared to resorcinarenes and calix[4]arenes, which can lead to different arrangement of the amphiphiles during their self-assembly.

The prepared SLNs were stable enough to survive the vacuum treatment needed for the SEM sample preparation (gold coating) and in most cases the SLN suspensions were stable over several months. Some exceptions to this rule were observed, a couple of times the samples formed a visible precipitation within a week or two accompanied with particle size increase. In the SEM images some of the samples showed signs of early aggregation tendency, and in a couple of samples complete assimilation of the smaller < 1 μm particles into the substrate was seen. Also, the nature of the substrate has an effect during the sample preparation since on the hydrophobic carbon tape most of the particles deformed in contrast to the hydrophilic SiOₓ surface, where hydrophilic interactions with the surface could help to maintain the integrity of the particles. According to the preliminary results C11BC5 seemed to give the most stable SLN's. However, further studies are needed to establish a more systematic survey on the properties of the resorcinarene bis-crown SLNs.

As a conclusion, resorcinarene bis-crowns bind Cs⁺ in solution and form solid state complexes with Cs⁺ and K⁺. The lower rim alkyl chains affect the crystal packing and the amphiphilic properties since C2BC5–C4BC5 form layered packing, whereas, a bilayered packing typical for amphiphilic molecules is observed with C_5 and longer alkyl chains. The locked boat conformation of CNBC5 makes also the short chain derivatives behave as amphiphilic molecules, which form stable solid lipid nanoparticles with slight dependence between the SLN size and the alkyl chain length.

Experimental

X-ray crystallography

Single crystal X-ray data were recorded on a Nonius Kappa CCD diffractometer with Apex II detector using graphite monochromatized CuK$_\alpha$ (λ = 1.54178 Å) radiation at a temperature of 173 K. The data were processed and absorption correction was made to all structures with Denzo-SMN v.0.97.638[28] unless otherwise mentioned. The structures were solved by direct methods (SHELXS-97) and refined (SHELXL-97) against F² by full-matrix least-squares techniques using SHELX-97 software package (Table 3).[29] The hydrogen atoms were calculated to their idealized positions with isotropic temperature factors (1.2 or 1.5 times the C temperature factor) and refined as riding atoms. Crystal structure analysis was done using Mercury CSD 2.4 software.[30] Crystallographic data (excluding structure factors) for the structures in this paper have been deposited with the Cambridge Structural Data Center as supplementary publication nos. CCDC 854053–854058. Copies of the data can be obtained free of charge from The Cambridge Crystallographic Data Centre *via* www.ccdc.cam.ac.uk/data_request/cif.

NMR titration

4 mM CNBC5 was titrated with CsPF₆ solution in acetone-D6 and the ¹H NMR spectra were recorded after each addition at 30 °C. The shift the in aromatic resorcinarene signal at 6.005 ppm for the free host was followed and the binding constants were

calculated using WinEQNMR2 software.[31]

SLNs

Solid lipid nanoparticles were prepared by a solvent replacement method.[32] 5 mg (or 5.27 µmol, 5.2–7.8 mg) of CNBC5 was dissolved in 1.5 ml of THF and 50 ml of purified water (Millipore, resistivity >18 MΩ) was added at a constant flow during 10 s into the organic solution under vigorous stirring at 800 rpm with a magnetic stirrer. A cloudy suspension formed immediately. The suspension was stirred for an additional minute and THF was removed under reduced pressure by a rotary evaporator (44 °C, 60–70 mbar). The volume of the suspension was adjusted to 50 ml giving the final nanoparticle concentration of 100 mg/L (or 0.1 mM). The hydrodynamic diameter of the nanoparticles was measured using dynamic light scattering (Beckman Coulter N5 Submicron Particle Size Analyzer) at 90° angle in water using plastic cuvettes (3 min equilibration, 3 min measurement). Three samples for each SLN were measured. The morphology and size of the SLN's were analyzed using scanning electron microscopy (Zeiss EVO 50). Sample preparation: a drop of SLN suspension was pipetted on a piece of silicon wafer attached by carbon tape to the sample holder and dried at ambient conditions. Samples were coated with gold (JEOL Fine coat Ion Sputter JFC-1100) prior to imaging.

Acknowledgement

and *ccw* enantiomers are shown; A/C plane runs through the upright aryl rings (A and C) and B/D plane through parallel aryl rings (B and D) respective to the methine plane C7-C14-C21-C28.
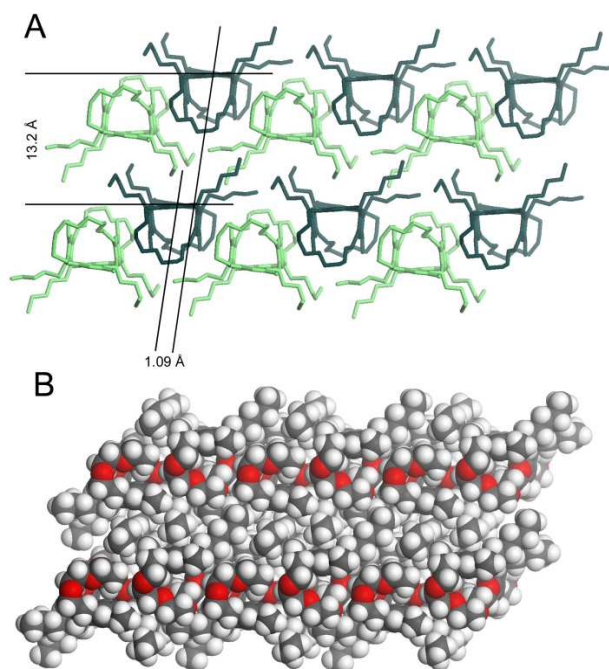
Fig. 2 Crystal packing of **C5**, flattened bilayer, in a stick model (A) with *cw* enantiomers with light and *ccw* in dark green, respectively, and a CPK representation (B), where the polar and non-polar regions are clearly visible. Disorder not shown for clarity.
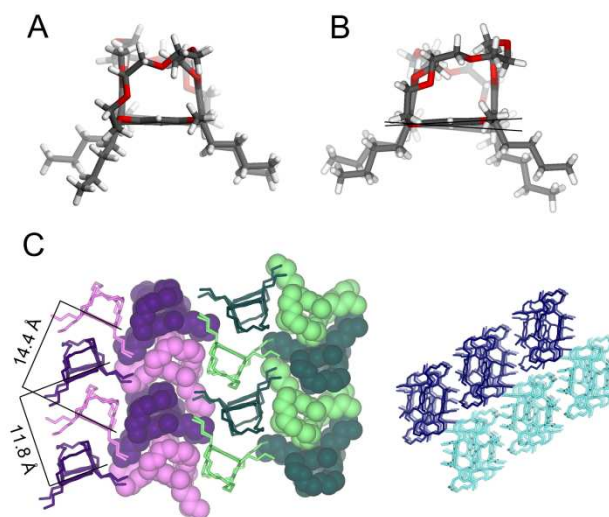
Fig. 3 Crystal structure of **C4** (disorder not shown). **C4** has two molecules, *I* (A) and *II* (B) in the asymmetric unit, the twist angle in *II* indicated. C) Pillar packing of **C4**: *cw* (up, purple) and *ccw* (down, green) enantiomers are separated in pillars with alternating *I* (dark shade) and *II* (light shade). Each pillar consists of a single enantiomer with methine plane angle (43.6°) and distances indicated; a top view shows pillars of *cw* and *ccw* enantiomers in dark and light color, respectively.

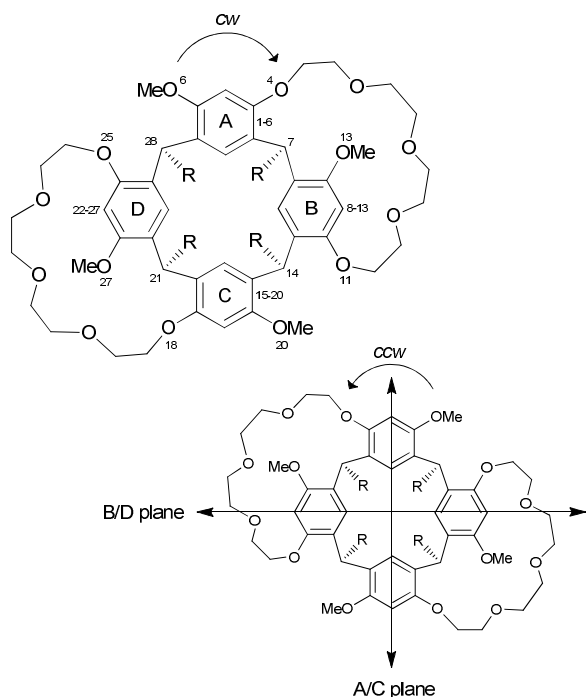**Fig. 1** Structure of the resorcinarene bis-crowns CNBC5, R=C_NH_{2N+1} where N= 2,3,4,5,7,9,10,11 with selected crystallographic numbering. *cw*
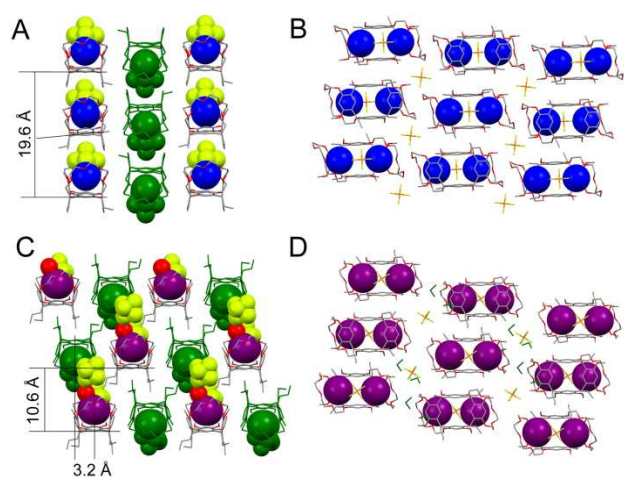
Fig. 4 Side and top views of the crystal packing of **C2K2** layers (A, B) and **C3Cs2** shifted capsules/ layers (C, D); side view: *ccw* enantiomers facing down (green); top view of a layer: outlying anions and ethanol molecules (in D, green) shown as a stick model.
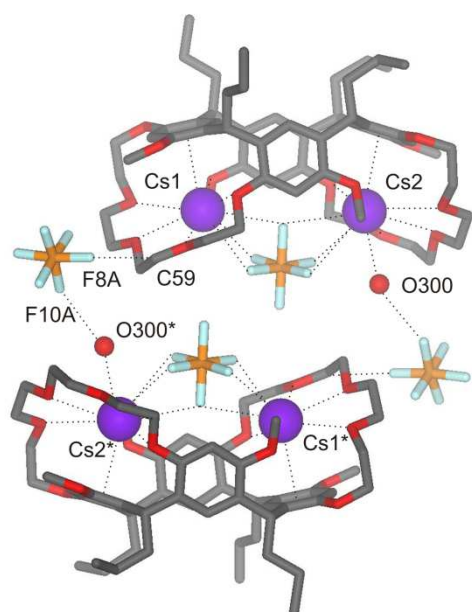

5

Fig. 5 C3Cs2 shifted capsule consisting of *cw* and *ccw* enantiomers, short contacts to the solvent and anions shown with dashed lines, atoms labelled with an asterisk are generated by a symmetry operation –x+1, –y+2, –z+1. Disorder not shown for clarity.
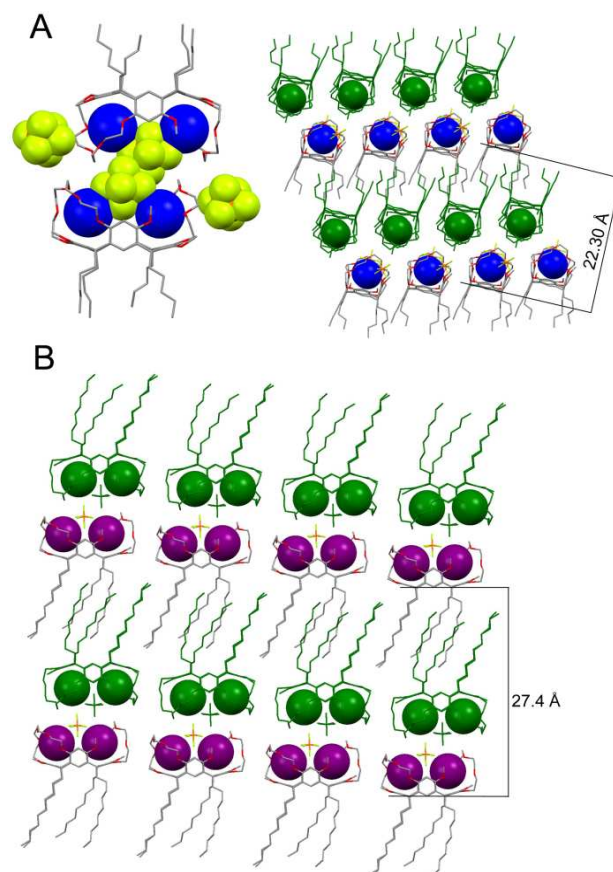
10

15

Fig. 6 C5K2: a front view of an offset capsule and a bilayer packing (A); 20 **C10Cs2**: a front view of a bilayer (B). *ccw* enantiomers in green color; outlying anions and disorder not shown.
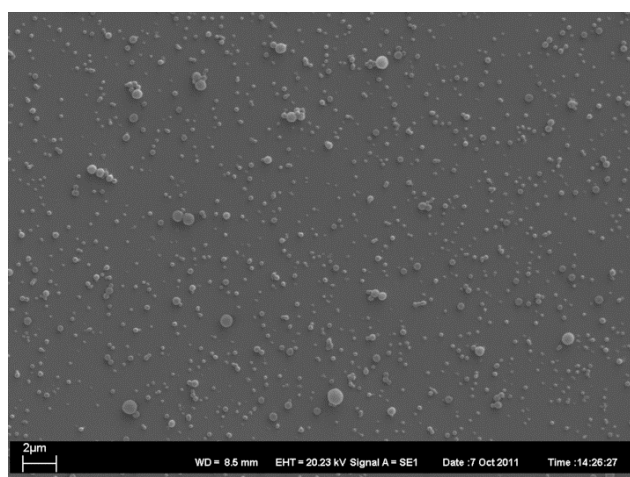

Fig. 7 SEM image of C11BC5 SLN on SiO$_x$; spherical particles with mean diameter of 300 nm are shown.
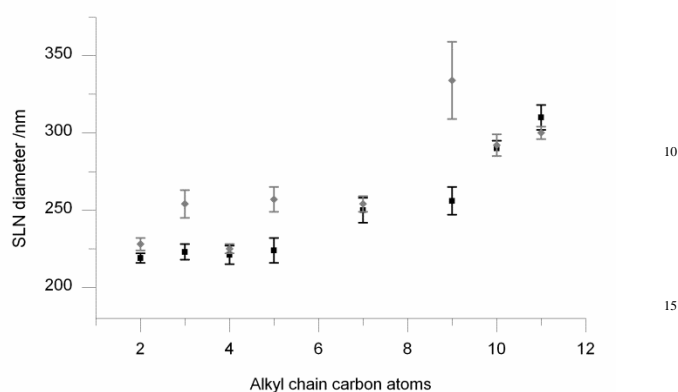
25

Fig. 8 SLN diameters for all CNBC5 (mean of the size distribution from DLS showing standard deviation). SLNs with a constant 0.1 mM concentration (black) and with a constant mass 100 mg/L (grey) show increasing diameter for longer alkyl chains.

Table 3 Crystal structure parameters.[a]

	C4	**C5**[b]	**C2K2**	**C3Cs2**	**C5K2**	**C10Cs2**
Composition	$C_{64}H_{92}O_{14}$	$C_{68}H_{100}O_{14}$	$[K_2(PF_6)$ $(C_{56}H_{76}O_{14})]PF_6$	$[Cs_2(PF_6)(C_{60}H_{84}O_{14})$ $(H_2O)]PF_6 \cdot 2C_2H_6O$	$[K_2(PF_6)$ $(C_{68}H_{100}O_{14})]PF_6$	$[Cs_2(PF_6)$ $(C_{88}H_{140}O_{14})]PF_6$
FW	1085.38	1141.48	1341.31	1695.18	1509.62	1977.76
Crystal system	Triclinic	Triclinic	Monoclinic	Triclinic	Triclinic	Triclinic
Space group	P-1	P-1	C2/c	P-1	P-1	P-1
a/Å	15.5428(4)	14.0441(5)	36.7201(7)	12.5169(3)	10.1107(5)	10.4675(4)
b/Å	16.2070(5)	16.3664(6)	19.6013(5)	17.8145(5)	17.014(1)	16.8329(7)
c/Å	28.1062(9)	16.5780(6)	19.4874(6)	18.0005(5)	22.918(1)	27.588(1)
α/°	78.135(2)	99.948(2)	90	70.468(2)	109.330(3)	96.549(2)
β/°	88.922(2)	104.812(2)	116.801(1)	79.987(2)	95.789(3)	97.404(2)
γ/°	62.066(2)	112.067(2)	90	82.768(1)	89.952(3)	93.961(2)
Volume/Å^3	6096.8(3)	3256.7(2)	12519.6(6)	3715.3(2)	3698.9(3)	4771.6(3)
Z	4	2	8	2	2	2
D_{calcd}/Mg·m^{-3}	1.182	1.164	1.423	1.515	1.355	1.377
F(000)	2352	1240	5600	1732	1592	2056
μ/mm^{-1}	0.661	0.641	2.676	8.867	2.323	6.958
Crystal size/mm	0.15×0.10×0.07	0.26×0.20×0.10	0.15×0.10×0.05	0.30×0.20×0.10	0.10×0.10×0.05	0.20×0.08×0.06
Meas. reflns	29260	15745	17010	45717	16668	19576
Indep. reflns	20072	10590	10802	12755	11625	13472
R_{int}	0.0909	0.0593	0.0589	0.0587	0.1246	0.0927
R_1 [$I > 2\sigma(I)$]	0.0792	0.0852	0.0788	0.0449	0.0969	0.0826
wR_2 [$I > 2\sigma(I)$]	0.1761	0.2238	0.2001	0.1125	0.2355	0.1656
GooF on F^2	1.063	1.025	1.029	1.044	1.074	1.055
Largest diff. peak and hole/ e Å^{-3}	0.470, -0.378	0.580, -0.335	0.713, -0.646	0.882, -1.338	0.828, -0.457	1.027, -0.634

[a] Unit cell dimensions for C9BC5, C10BC5 and C11BC5 at the endnotes. [b] Isomorphous structures obtained from *tert*-butanol/methanol, *iso*-butanol/methanol, and ethanol solutions.

Notes and references

* **C9**: C2/c, *a*=52.218(1), *b*=14.6337(3), *c*=21.2980(5), *β*=101.245(1), V=15962.2(6); **C10**: C2/c, *a*=54.931(2), *b*=14.5212(5) *c*=21.3574(8), *β*=102.354(2), V= 16641.4(1); **C11**: C2/c, *a*=57.434(2), *b*=14.5337(4), *c*=21.1864(7), *β*=100.590(1), V=17383.6(1).

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