Giant capsids from lattice self-assembly of cyclodextrin complexes

Proteins can readily assemble into rigid, crystalline and functional structures such as viral capsids and bacterial compartments. Despite ongoing advances, it is still a fundamental challenge to design and synthesize protein-mimetic molecules to form crystalline structures. Here we report the lattice self-assembly of cyclodextrin complexes into a variety of capsid-like structures such as lamellae, helical tubes and hollow rhombic dodecahedra. The dodecahedral morphology has not hitherto been observed in self-assembly systems. The tubes can spontaneously encapsulate colloidal particles and liposomes. The dodecahedra and tubes are respectively comparable to and much larger than the largest known virus. In particular, the resemblance to protein assemblies is not limited to morphology but extends to structural rigidity and crystallinity—a well-defined, 2D rhombic lattice of molecular arrangement is strikingly universal for all the observed structures. We propose a simple design rule for the current lattice self-assembly, potentially opening doors for new protein-mimetic materials.
All living organisms are self-assembled entities where two major kinds of self-assembly are involved: the assembly of lipids into soft, fluidic membranes mainly driven by the hydrophobic interaction and the assembly of proteins into rigid, crystalline structures driven by a combination of hydrophobic, H-bonding and electrostatic interactions. The lipid assembly is extensively reproduced and well extended by synthetic amphiphilic small molecules, polymers and even nanoparticles to form lamellar, tubular, vesicular and micellar structures. The protein assembly that produces crystalline structures such as lamellae, tubules and polyhedra is, however, largely unparalleled by synthetic or non-peptide molecules with a few notable exceptions. The imbalance in the research of lipid and protein mimicry thus calls for more attention to the latter.

Looking beyond biomimetic self-assembly, one can notice that carbon allotropes share the same morphological pattern: graphite, graphene, nanotubes and C₆₀ in analogy to lamellar, tubular and polyhedral assemblies. The properties and functions of carbon allotropes, lipid assemblies and protein assemblies are, however, drastically different as carbon atoms are connected by chemical bonds, lipids by hydrophobic interactions and proteins by a combination of intermolecular interactions. In this context, what has been gradually recognized is the importance of intermolecular interactions, structural flexibility/rigidity and fluidity/crystallinity over that of morphology. For example, peptoids, amphiphilic hexabenzocoronene and catanionic surfactants were found to form, respectively, nanosheets, nanotubes and regular hollow icosahedra, which successfully mimicked protein assemblies’ morphologies but cannot rival their rigidity nor well-defined crystallinity.

Seeking for a synthetic system that parallels the morphology, rigidity and crystallinity of protein assemblies, we chose to study a supramolecular complex, sodium dodecyl sulphate (SDS)@2β-CD (one SDS molecule inside two β-cyclodextrin molecules, Fig. 1c). The complex was recently reported by us to form lamellar and tubular structures, but the study was restricted to morphology on the mesoscopic scale. In this paper, we reveal the existence of another structure, hollow rhombic dodecahedra and, more importantly, scrutinize the system on the microscopic/molecular scale to identify a high structural rigidity and well-defined, in-plane crystallinity.

**Results**

**General phase behaviour.** Depending on concentration, the SDS@2β-CD aqueous solutions can be divided into lamellar (50 to 25 wt%), tubular (25 to 6 wt%) and polyhedral (6 to 4 wt%) phases (Fig. 1c). A general phase diagram and discussion on the SDS/β-cyclodextrin (β-CD) stoichiometry is given in Supplementary Fig. 1. The lamellar and tubular phases were studied in our previous work, so we focus here on the structural rigidity and newly identified nanosheets and polyhedra. Nanosheets can be found in the tubular and polyhedral phases as the minority and possibly the intermediate structure (Fig. 2a–c). The nanosheets are a few hundreds of nm in lateral size, very thin as reflected by the low electron contrast, and of a typical parallelogram shape with sharp, straight edges and an obtuse angle of 104°. When excess salts were added to the SDS@2β-CD solutions in all the 3 phases, flake crystals precipitated with a shape the same as the nanosheets but much larger and thicker (Fig. 2d). The microscopical image of the lamellar phase is full of parallel lines with a uniform interval (Fig. 2e), typical for cross sections of lamellar structures.

In the tubular phase, tubular structures pervades (Fig. 2f) with a monodisperse diameter ~1 μm, a mean length ~40 μm and open ends. A closer inspection reveals that the tube walls are

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**Figure 1 | Stylized view of the lipid-like and protein-like self-assembly.** (a) Lipid molecules form lamellar, tubular and vesicular structures, the flexibility and fluidity of which are emphasized in the illustration. (b) Proteins form lamellar, helical tubular and regular icosahedral structures with rigidity and crystallinity (hexagonal lattice in this case). (c) SDS@2β-CD assemblies, in a protein-mimetic way, into lamellar, helical tubular and rhombic dodecahedral structures with inherent rigidity and in-plane, rhombic crystalline nature. In the molecular view, SDS is an anionic surfactant with a hydrocarbon tail (yellow) and a -(SO₄)⁻ headgroup (blue and red), while β-CD is a ring of hepta-saccharides (green C and red O atoms).
made of multiple, equally spaced and ~4 nm thick (for detailed discussion about layer thickness, please see Supplementary Fig. 2) layers (Fig. 2g–i). In a situation like Fig. 2i, the walls are fractured rather than bended. The straightness of the tubes is strikingly persistent across at least 4 orders of length scale: no mesoscopic bending on the order of 100 μm (Fig. 2f) nor microscopical fluctuation on the order of 10 nm (Fig. 2g,h). The persistence length of the tubes is estimated to be at least on the order of 1 m (Supplementary Fig. 3), order-of-magnitude higher than that of lipid tubules ~10 μm (ref. 23) and microtubules ~1 μm (ref. 24), signifying a high rigidity.

Hollow rhombic dodecahedra. In the polyhedral phase, transmission electron microscopy (TEM) pictures show polygons, mostly hexagons and some octagons, ~1 μm large with sharp edges (Fig. 3a,d). We found that the cross sections of a rhombic dodecahedron can account for the observed polygons. In Fig. 3c, a rhombic dodecahedron is rotated along x and y axis and its cross sections along the xy plane are represented by the dark grey polygons. The indexed polygons are compared with the TEM pictures in Fig. 3d, reaching a good agreement. The freeze-fractured (FF) TEM pictures (Fig. 3d, middle row) reveal that the dodecahedra are hollow and, sometimes, of double shells. The cryogenic transmission electronic microscope (cryo-TEM) pictures (Fig. 3d, bottom row) show a normal dodecahedra and a rare one with a nonconcentric dodecahedron inside.

The rhombic dodecahedral geometry is further consolidated by the atomic force microscopy (AFM) measurements (Fig. 3b), in which bumpy, rhombic objects can be observed. We reason that the observed morphology is resulted by the collapse of hollow dodecahedra on a flat substrate in dry condition (Fig. 3e, see the figure caption). In the case of a single-shell dodecahedron (shell thickness 4 nm), the bumps, sink and pits are expected to be higher than, close to, and lower than 8 nm, respectively. These expectations are confirmed by two examples in Fig. 3f, where the yellow-white, green and green-blue correspond to heights of 16 nm, 8 and 4 nm, respectively. We therefore determined that the structures are of a hollow rhombic dodecahedral geometry, which has not been observed before in self-assembly systems and is of potential uses in templated particle synthesis and colloidal self-assembly.

Universal in-plane molecular arrangement—a 2D rhombic lattice. X-ray scattering technique was employed to resolve the molecular arrangement of SDS@2β-CD inside the observed structures. Small-angle X-ray scattering (SAXS) measurements of the lamellar, tubular and polyhedral phases (microstructures randomly orientated) were conducted in European Synchrotron Radiation Facility, Grenoble, France. The SAXS curves can be divided into 3 regions (Fig. 4a). In the cyan region (0–4 nm⁻¹), 4 curves share a similar decaying trend corresponding to the form factor. The pink region features equally spaced, lamellar peaks that gradually vanishes on the dilution of SDS@2β-CD, indicating the increase of layer-to-layer distance (from 15 to 42 nm) and the lessening of layers in a stack. This is in line with the TEM observations where the lamellar, tubular and polyhedral structures have a dozen of, several and one or two coherent layers, respectively (Figs 2 and 3). In the green region (4–8 nm⁻¹), 3 distinctive structural peaks can be identified for all the curves, implying a single in-plane crystalline arrangement shared by all the microstructures. The scattering curves up to 4 nm⁻¹ were fitted by a bilayer form factor and lamellar structure factor (modified Caillé theory25, Fig. 4b). The corresponding electron density profile suggests a bilayer...
arrangement of SDS@2β-CD with its long axis perpendicular to the layer (Fig. 4c). In Caille theory, the Caille parameter \(Z\) describes the thermo fluctuations of the layers, inversely related to the layer rigidity. The fitted \(Z\) (on the order of 0.01) underlines that the rigidity of SDS@2β-CD microstructures is order-of-magnitude higher than that of lipid membranes (\(Z\) usually on the order of 1)\textsuperscript{25}.

To precisely determine the in-plane lattice structure, we aligned the tubular structures and then studied the sample with wide-angle X-ray scattering. By carefully loading tubular samples into capillaries, we achieved flow-induced alignment of tubes (Supplementary Fig. 4). The scenario of a multilamellar tube diffracting X-ray is schematically illustrated in Fig. 4d, where the lamellar period (10s of nm) gives cyan dots at small \(q\) along the equator and the in-plane lattice (about the molecular size, ~1 nm) yellow dots at large \(q\). In line with this scenario, a few lamellar peaks and a broad form factor peak are located on the equator at small \(q\) (Fig. 4e), while a well-defined diffraction pattern is observed at large \(q\) (Fig. 4f). The pattern’s symmetry along the equator and medial axis signifies a helical nature for the tubes. The pattern is perfectly matched by two mirrored lattices (green and yellow grids).

The in-plane unit cell is thus resolved as a rhombus with \(a = b = 1.52\) nm (comparable to β-CD diameter, 1.5 nm) and \(\gamma = 104^\circ\), in excellent agreement with the reported single-crystal data for other aliphatic chain/β-CD complexes\textsuperscript{26}. With the \(b\) axis-to-equator angle = 3\(^{\circ}\), the lattice scroll up right-handed or left-handed into helical tubes (Fig. 4g). Such a lattice is stabilized by an extensive network of direct (CD to CD) H-bonds and indirect (water mediated) H-bonds, where the O atoms on two neighbour CD rims are close enough to form multiple direct H-bonds (highlighted by magenta in Fig. 4h, the top and side views). In addition, we argue that the \(\gamma = 104^\circ\) is a consequence of maximization of the direct H-bonds with the given 7-fold molecular symmetry of β-CD (Supplementary Fig. 5).

It is noteworthy that the rhombic lattice is universally presented in all the observed microstructures as evidenced by the identical in-plane structural peaks in all the 3 phases (Fig. 4a, 4b, 4c).

Figure 3 | Morphology of the SDS@2β-CD polyhedra. (a) A TEM picture of the negatively stained polyhedra, scale bar, 1\(\mu\)m. (b) An AFM image of the polyhedra on mica substrate in a dry condition, scale bar, 2\(\mu\)m. (c) A diagram showing the top view and cross-sectional view (along the xy plane, the dark grey polygons) of a rhombic dodecahedron with different orientations. The orientation is indexed to be compared with TEM images. (d) Selected TEM pictures corresponding to the rhombic dodecahedral geometry with varied orientations. From top to bottom rows are images by negatively stained TEM, FF-TEM (highlighting a double-shell configuration), and cryo-TEM (featuring a rare, nonconcentric configuration). (e) A simple scenario of the collapse of hollow dodecahedra. First, one of the 12 equal faces rather than any vertex will land on a flat substrate, then the 2 faces parallel to the substrate are labelled cyan, the 2 faces perpendicular to the substrate yellow and the other 8 faces green. Second, we consider the dodecahedron to experience a vertical pressure due to gravity and dehydration, then the cyan faces may remain intact, the yellow faces may be forced to fracture, and the green faces squeezed. Finally, the dodecahedron collapses into a somewhat rhombic structure with bumps along the equator, a sink in the center, and pits near two poles. (f) Selected AFM images are accounted by the scenario. Please note that the dodecahedra are about 1\(\mu\)m large so the scale bars for d and f are not shown.
Precipitates depending on the concentration of SDS into hollow rhombic dodecahedra, and to stack up into flake building blocks or intermediate structures to laterally expand into well-defined in-plane crystalline nature. The nanosheets serve as lattice self-assembly, the resultant structures of which feature a lamellar nor polyhedral structures. The observed lanreotide was reported to form monodisperse nanotubes (although not similar in shape, the formers are much larger in size). The SDS@2b-CO3 resembles the protein assembly such as viral capsids. Although lattice self-assembly of SDS@2b-CO3 lipids or amphiphilic molecules that are fluidic and soft, the and salts (Fig. 1c). In contrast to the traditional self-assembly of b-CD structures.

Figure 4 | X-ray scattering of the SDS@2b-CD structures. (a) SAXS curves of the lamellar, tubular and polyhedral phases with different SDS@2b-CD concentrations. The curves (vertically shifted for clarity) are divided into pink, cyan (overlapping with pink) and green regions, corresponding to lamellar structure factor, form factor and in-plane crystallinity, respectively. (b) The SAXS curve (30 wt%, black dots) up to 4 nm–1 is fitted by a bilayer form factor (cyan line) and lamellar structure factor (blue line) with a good result (red line). The cyan and blue lines are vertically shifted for clarity. (c) The electron density profile across the bilayer. The thickness 3.1 nm and electron density of the core part (the green region) matches a vertical stack of 4 b-CD molecules (0.75 nm high), the thickness and electron density of the outer part suggests two SDS molecules with their tails threading the CD cavities and headgroups pointing outwards. Threaded b-CD molecules usually stack in a head-to-head fashion to maximize the inter-CD H-bonding along the threading direction. (d) A scheme of wide-angle X-ray scattering of aligned multilamellar tubes with in-plane crystallinity. (e) Scattering pattern along the equator at small angle, corresponding to the blue box in f. (f) Scattering pattern at large angle is super-positioned by the mirrored yellow and green grids. (g) The in-plane lattice with a rhombic unit cell, a = b = 1.52 nm, γ = 104 ° and the b axis-to-equator angel = 3°. (h) Top and side views of the SDS@2b-CD lattice, highlighting the possible, inter-CD H-bonds (magenta).

Possible pores at the vertices of the rhombic dodecahedra. We noticed in the EM pictures that the dodecahedra vertices are usually rounded, indicative of pores at the vertices. Following Dubois and Zemb et al.'s arguments4,5, we reason two possible ways to understand the formation of pores. One is to remove the highly undesired curvature singularity at the vertices by forming pores. Another is to minimize total curvature energy by creating negative Gaussian curvature at the pores and leaving the spontaneous curvature and Gaussian curvature to be close to zero at the facets. The zero curvature is clearly favoured as evidenced from the predominate lamellar and tubular (nearly planar due to the large diameter) structures at higher concentrations. The negative local curvature could be partially relieved by higher surface charge at the pores. In the previous catanionic icosahedral system4,5, the excess anionic surfactant molecules enrich themselves at the pores to form half micelles to cover edges of the pores. In the current case, if the hydrophilic outer rims of CD are exposed to water at the edges of pores or planar structures, its energy penalty is much less than that of exposing hydrophobic chains to water in the catanionic surfactant system. It is, therefore, not clear at this stage about the exact molecular arrangement at the pores. For extended discussion on the rhombic dodecahedral geometry, please see the Supplementary Discussion28,29.

Encapsulation by the SDS@2b-CD tubes. The main function of viral capsids and bacterial compartments is to encapsulate DNA, RNA, proteins and even magnets2,5,16. In this context, we recently
reported that the SDS@β-CD tubes can spontaneously enclose colloidal particles with different shapes and chemistry to form 1D straight chains, double/triple helices and hybrid structures. An interesting case is the haematite chain inside a tube similar to the magnetosome chain inside magnetotactic bacteria. In this work, we take a step forward by including soft liposomes into the tubes. Since SDS is a surfactant capable of solubilizing lipids, liposomes with a high cholesterol content were chosen to resist SDS. The added liposomes ~0.8 μm remain intact for at least several hours and are indeed trapped in the tubes (Fig. 5a). This hierarchical, liposome-in-tube construction reminds us of bacterial walls where the lipid membrane is covered by a surface layer of crystalline proteins (S-layer). It is envisioned that viable bacteria (typically ~1 μm large) could be included into the tubes, enabling the research of bacterial confinement and assembly.

**Discussion**

The generalization of the lattice self-assembly is established in a matrix of CD complexes with the guest molecules ranging from ionic, zwitterionic, to nonionic amphiphilic molecules and the host molecules being α-CD and β-CD. Replacing β-CD with 2-hydroxypropyl-β-CD or methyl-β-CD makes the complexes fully dissolvable in water and unable to assemble, underlining the necessity of inter-CD H-bonds (see also the discussion of Fig. 4h and Supplementary Fig. 5 for the importance of the H-bonding network). Recalling that excess of salt results in precipitation, we argue a simple design rule for the lattice self-assembly: the presence of strong, directional in-plane attraction (H-bonds in this case) and out-of-plane repulsion (for example, electrostatic or steric). The insensitivity of the assembly behaviour to the head group of the guest molecules suggests that one can functionalize the head group to mimic bacterial compartments for bio-mineralization and enzyme catalysis. Different host molecules are expected to produce fairly different lattices: hexagonal for α-CD, rhombic for β-CD and square for γ-CD depending on the molecular symmetry (6-, 7- and 8-folds, respectively, Fig. 5b). Looking ahead to possible applications, fixation of the CD networks by polymerization and subsequent removal of the guest molecules could form selective membranes as well as porous tubes and polyhedra with hydrophobic channels inside CD molecules and hydrophilic channels between CD molecules (Fig. 5c, the yellow and cyan regions). The pore geometry might be precisely tuned by employing different CD molecules. Such membranes are, in principle, similar to the reported colloidal Kagome lattice but with much smaller pore sizes.

**Methods**

**Materials.** SDS (99%) was purchased from Acros Organics and used as received. β-CD was purchased from Sinopharm Chemical Reagent with a water content of 14%. The silica particles with polydispersity <5% and haematite particles with polydispersity <10% were synthesized in the van’t Hoff Laboratory for Physical and Colloid Chemistry in Utrecht University. 1-Palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine and cholesterol (98%) were obtained from Avanti Polar Lipids Inc. β-BODIPY 530/550 C8, HPC was purchased from Molecular Probes. HEPES (sodium salt) was obtained from Acros Organics.

**Preparation of SDS@2β-CD suspensions.** Desired amounts of β-CD, SDS and water were weighed into tubes to give a constant β-CD/SDS molar ratio of 2:1 and different total concentrations of β-CD and SDS (in wt%). The mixtures were heated to ~60 °C to obtain transparent and isotropic solutions, where SDS@β-CD is formed. The solutions were then cooled to 25 °C and incubated at least 48 h to allow SDS@β-CD to form structures. According to the morphology of the structures, the whole studied concentration range is divided into three phases, lamellar (25–50 wt%, semi-transparent and highly viscous), tubular (6–25 wt%, turbid and viscous) and polyhedral (4–6 wt%, slightly turbid and watery), respectively. The structures therein are stable for months.

**Co-assembly of colloidal particles or liposomes with SDS@β-CD tubes.** Aqueous suspensions of colloids were centrifuged at 2,000–3,000 r.p.m. for 15 min, followed by removal of the supernatant water. A 10 wt% SDS@β-CD tubular suspension was added to the centrifuge tube to give a final tube/colloid mixture containing 1–20 wt% colloids. The sample was heated to ~60 °C to melt the tubes and was sonicated to disperse the particles. Then the centrifuge tube was cooled to room temperature and incubated with gentle vibration to avoid sedimentation of the particles. On cooling, the tubes formed and the particles were spontaneously included into the tubes.

**Transmission electron microscopy.** TEM images were recorded on a JEM-100 CX II transmission electron microscope (JEOL, Japan, 80 kV). The samples were prepared by dropping solutions onto copper grids coated with Formvar film. Excess water was removed by filter paper, and the samples were dried in ambient environment at room temperature for TEM observation. In case of the polyhedral samples, they were negatively stained by uranyl acetate before water removal.

**Freeze-fracture transmission electron microscopy.** A small drop of a sample was placed between two copper disks. The mounted sample was plunged into liquid propane that was cooled by liquid nitrogen. Microstructures are preserved by this procedure most of cases. A freeze-fracture apparatus (BalzersBAC400, Germany) was employed to fracture and replicate the specimen at ~140 °C. A thin layer of platinum-carbon was cast onto the specimen to obtain the replicas, which were then ready for TEM observation.

**Cryogenic transmission electronic microscope.** A few microliters of samples were mounted onto a copper TEM grid and the excess solution was removed with filter paper. The specimen was immersed into a cryo-box (Carl Zeiss NTS GmbH, filled with liquid ethane) to be rapidly cooled to ~170 to ~180 °C. The specimen was then transferred to a Zeiss EM922 FEFTEM (Zeiss NTS GmbH, Oberkochen, Germany) by a cryo-transfer holder (Cf3500, Gatan, Munich, Germany). TEM observation was made at ~160 °C with an acceleration voltage of 200 kV. Reduced electron doses (500–2,000 e nm^-2_) were used to obtain zero-loss filtered images.
A bottom-mounted CCD camera system (UltraScan 1,000, Gatan) was employed to record all images, which were then processed by Digital Micrograph 3.9.

**Confocal laser scanning microscopy.** The SDS@β-CD structures were stained by Nile red in the following way. A drop of 15 μl Nile red in acetonitrile solution (1 mg ml⁻¹) was added to a test tube, followed by volatilization of the acetonitrile. A desired amount of SDS@β-CD aqueous solution was then added to the tube. The samples were then incubated at least 24 h for the dye molecules to diffuse into the SDS@β-CD structures. The stained SDS@β-CD, colloid-in-tube, or liposome-in-tube suspensions were loaded into glass capillaries (Vitrocom, 0.1 x 2 x 50 mm), which were then sealed by a ultraviolet-curing epoxy glue. An inverted confocal laser scanning microscopy (Leica, True Confocal Scanner SP, Germany) was used to conduct experiments in fluorescence and differential interference contrast modes.

**Atomic force microscopy.** AFM measurements were conducted by Nanoscope IIIa (Digital Instruments Inc., USA) in tapping mode under ambient conditions on mica substrates.

**Small-angle X-ray scattering.** Measurements were performed at the DUBBLE beamline of the European Synchrotron Radiation Facility (ESRF, Grenoble, France). The microradian resolution setup was used. A PILATUS-1M detector (detected by SASfit software package) was fitted by SASfit software package.

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**Acknowledgements**

We appreciate the insightful suggestions from Thomas Zemb. L.J. acknowledges support by the startup funding from Jinan University. L.J. thanks helpful discussion with Albert P. Philipe and Samia Oushaji. S.G. acknowledges support by the Institute for Basic Science, project code IBS-R020-D1. The personnel of the DUBBLE beamline is thanked for the support during SAXS measurements. The Nederlandse Organisatie voor Wetenschapelijk Onderzoek (NWO) is acknowledged for the provided synchrotron beam-time.

**Author contributions**

L.J. conceived the experiment and wrote the manuscript; S.Y. performed the experiments; L.J., Y.Y. and J.H. discussed microscopy results; L.J., A.V.P. and L.M.J.K.-B. performed the X-ray measurements and discussed the results; M.D. took the cryo-TEM pictures. All authors contributed to analysing the results and writing the paper.

**Additional information**

**Supplementary Information** accompanies this paper at http://www.nature.com/naturecommunications

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**How to cite this article:** Yang, S. et al. Giant capsids from lattice self-assembly of cyclodextrin complexes. *Nat. Commun.* 8, 15856 doi: 10.1038/ncomms15856 (2017).

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Supplementary Figure 1. The general phase diagram of the SDS/β-CD aqueous solution at room temperature. Along the SDS/β-CD ratio axis, plate crystal precipitates tend to form at low ratio because β-CD is of very limited solubility in water, and the solution is clear without any observable aggregates at high ratio because SDS/β-CD 1/1 complex is of excellent solubility in water. Near the 1/2 stoichiometry line, the complexes form lamellar, tubular, and polyhedral structures depending on the concentration. Although the current paper is focused on the assembly behavior along the stoichiometry line, we did measure the SDS/β-CD ratio inside assembly structures using PGSE-NMR (pulsed-gradient Spin-Echo nuclear magnetic resonance) method for a few selected samples that are a bit off the stoichiometry line (crosses in the phase diagram). Briefly, this method can measure the concentrations of free SDS and β-CD molecules that do not participate into the assembly structures because free molecules are of high diffusivity. For details of this method, please see our previous publication. According the results, we conclude that the SDS/β-CD ratio inside the assembly structures is always very close to 1/2 (at least 98% of the complexes are in the 1/2 form) even when the bulk ratio is a bit off 1/2.
Supplementary Figure 2. The thickness of the SDS@2β-CD layer. a, The swelling behavior in lamellar (I) and tubular (II) phases. b and c, AFM measurements of a single-walled, collapsed tube. Scale bar: 2 μm. d, The apparent, projected, and true thicknesses in cryo-TEM observations. d, AFM measurements of a single-shell, collapsed polyhedron. The layer thickness of the lamellar, tubular, and polyhedral structures is determined to be about 4 nm by a battery of independent methods. The lamellar phase is governed by the classic swelling relationship, \( d = t / \varphi \), where the repeat distance \( d \) is determined by the first harmonic peak \( q_1 \) (\( d = 2\pi / q_1 \)), \( t \) is the thickness of the SDS@2β-CD layer, and \( \varphi \) is the volume fraction of SDS@2β-CD. A linear fitting of the points in Supplementary Figure 2a gives a \( t \) 3.9 nm. For the tubes, \( t \) is directly determined from AFM measurements (b-c) to be 4 nm. The layer thickness observed from the cryo-TEM pictures (Figure 2g-2i) ~10 nm is actually a sum of projected and true thicknesses (Figure S1d), thus largely overestimating \( t \). The shell thickness of the polyhedral is identified to be 4 nm by AFM measurements (e). The fitting of the SAXS curves reaches a thickness of 4.1 nm (Figure 4b and 4c).
Supplementary Figure 3. Optical and Confocal Microscopy Images of SDS@2β-CD tubes for the Estimation of Persistence Length. The tube diameter is ~1.1 μm in all the three images. Since the tubes barely bend in the entire view of optical microscopy, it is very difficult to precisely measure the persistence length, $P$. We thus try to roughly estimate $P$. The average bending of the tubes (averagely ~40 μm long) is smaller than 1 °, so it takes 2 m long to bend 90 °. $P$ is estimated to be at least on the order of 1 m, so the Young’s modulus is on the order of 1 GPa.
Supplementary Figure 4. Alignment of SDS@2β-CD tubes. a, The capillary loaded with aligned tubes under polarized optical microscopy, scale bar 1 mm. b and c, longitudinal and radial cross-sectional views of the tubes under CLSM, scale bars 15 and 3 μm, respectively. The alignment is confirmed by the uniform color in the birefringence image (a) and the longitudinal and radial cross-sectional views of the tubes (b and c).
Supplementary Figure 5. The magic angel 104 ° as a consequence of maximization of the direct H-bonds with the given 7-fold molecular symmetry of β-CD. This angel appears in the nanosheets (Figure 2a-2c), precipitated flakes (Figure 2d), and the tube diffraction (Figure 4f). Here we try to use simple model to account for the angel. β-CD is a ring of 7 saccharide moieties with H-bonding sites on the rim, we thus treat it as a regular heptagon with H-bonding sites on the 7 vertices. The heptagons are arranged in a rhombic lattice, where $a$ and $b$ ($a = b$) are constants, $\gamma$ is a variant, and the self-orientation of each heptagon $\theta$ is the other variant ($a$). It is reasonable to assume that the β-CD molecules would arrange themselves to maximize direct H-bonding by minimize the distances between H-bonding sites. The sum of the distances between two nearest H-bonding sites (highlighted by the two circles) is a function of $\gamma$ and $\theta$, which is plotted as a color map (b). Clearly, the minimum (dark blue) corresponds to $\gamma = 104 °$. Therefore, we argue that the $\gamma 104 °$ is a consequence of maximization of the direct H-bonds with the given 7-fold molecular symmetry of β-CD. Following this argument, we expect the lattice of α-CD and γ-CD to be hexagonal and square, respectively. In addition, we note that the exposed edges of nanosheets and flake crystals are predominantly (10) and (01) edges as a result of maximizing the direct, inter-CD H-bonding.
**Supplementary Discussion**

The current dodecahedra were formed by CD complexes (the building units) at a desired concentration via reversible self-assembly: at high temperature ~ 60 °C the building units are fully dissolved in water, forming no dodecahedra nor planar lattice, at room temperature the building units assemble into dodecahedra (up to 1 μm) in coexistence with a minority of planar structures (a few hundreds of nm). There are three possible pathways of dodecahedron formation: the building units add in one by one to form the dodecahedra, small planar structures (several hundreds of nm, like what we observed) assemble into the dodecahedra, and a single large planar structure (has to be several μm large) buckles into a single rhombic dodecahedron. As we did not observe any planar structure that large, we speculate that the buckling pathway is of lower possibility and that the former two pathways or their combination are of higher possibility.

It is pleasing to see the present self-assembly of hollow rhombic dodecahedra as a manifestation of the quasi-equivalence principle—the association of identical building units into highly symmetric structures due to strong intermolecular H-bonding with high efficiency and no templates\(^2\)—although the current polyhedron size and free energy associated to pore formation might exceed the valid range of the principle. Following Lidmar et al.’s arguments,\(^3\) the Foppel–von Karman number is defined as

\[
\gamma = \frac{E^{2D} R^2}{\kappa},
\]

where \(E^{2D}\) is the 2D Young’s modulus, \(R\) the facet size, and \(\kappa\) the bending rigidity. Larger \(\gamma\) favours buckled facets over spherical geometry. In our case, the 3D Young’s modulus is estimated to be on the order of 1 GPa, so \(E^{2D} \approx 4 \text{ Pa*m}\) and \(\kappa \approx 6\text{E-18 J}\). Given \(R \approx 1\text{E-6 m}\), the \(\gamma\) is on the order of 1E6. Such a large value indicates that the quasi-equivalence principle is not relevant to the formation of these shapes. It is also surprising that, unlike the icosahedral geometry favoured by catanionic surfactants and many capsid proteins, the rhombic dodecahedral geometry is dominant possibly as a consequence of the in-plane rhombic lattice.
Supplementary References

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