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Simultaneous Organic and Inorganic Host-Guest Chemistry within Pillararene-Protein Cage Frameworks

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Abstract: Supramolecular self-assembly of biomolecules provides a powerful bottom-up strategy to build functional nanostructures and materials. Among the different biomacromolecules, protein cages offer various advantages including uniform size, versatility, multi-modularity, and high stability. Additionally, protein cage crystals present confined micro-environments with well-defined dimensions. On the other hand, molecular hosts, such as cyclophanes, possess a defined cavity size and selective recognition of guest molecules. However, the successful combination of macrocycles and protein cages to achieve functional co-crystals has remained limited. In this study, we demonstrate electrostatic binding between cationic pillar[5]arenes and (apo)ferritin cages that results in porous and crystalline frameworks. The electrostatically assembled crystals present a face-centered cubic (FCC) lattice and have been characterized by means of small-angle X-ray scattering and cryo-TEM. These hierarchical structures result in a multiadsorbent framework capable of hosting both organic and inorganic pollutants, such as dyes and toxic metals, with potential application in water-remediation technologies.

The design and development of hierarchical structures based on the bottom-up assembly of biological building blocks has received growing interest in nanotechnology and material science. With the correct choice of building blocks, the resulting highly ordered (crystalline) structures can be engineered to present a porous structure with well-defined pore sizes similar to metal–organic frameworks (MOFs),[11] covalent organic framework (COFs),[12] perovskites,[13] and zeolites.[14] Such porous structures hold the potential to be applied in technologies including drug delivery, water remediation, and catalysis.[5–10] Among the plethora of biomolecules available, self-assembled nanocompartments such as protein cages are of high interest due to their uniform size and shape.[11–16] Together with their ability to encapsulate cargo, protein cages provide biocompatibility and biodegradability with chemical and thermal stability.[17–19] Protein cages have been extensively exploited as reaction vessels, templates for materials synthesis, drug carriers, bioimaging, and vaccine development.[19–23] Among many protein cages, ferritin (Ft) has emerged as an exceptional and promising building block. The iron oxide-containing cage can be found in eukarya, archaea, and bacteria and plays a vital role in iron regulation. The iron-free cage (apoferitin, aFt) presents an inner cavity, which can act as a host for a variety of organic and inorganic materials.[24] Due to their uniform shape and size, both cages, that is, Ft and aFt have been ideal candidates to be organized into multifunctional higher-order superlattices. They have already been used to create higher-order crystalline structures by self-assembly.[25] Together with organic dyes,[26] dendrons and dendrimers,[27–29] polypeptides and proteins,[30,31] metal ions or nanoparticles,[32–35] and macrocyclic molecules,[36] with varying potential applications in catalysis,[31,37] storage, and separation. Even though proteins are known for their relatively low stability outside aqueous media, the structural robustness of these materials in dry state has been previously demonstrated.[38] Their stability can also be increased by means of small molecule crosslinking,[38,39] or embedding the structure within polymeric[40] or inorganic matrices.[41]

Macrocycles such as cyclophanes have a cavity of defined size that can bind guest molecules through noncovalent interactions with high selectivity and affinity. Pillar[n]arenes are composed of “n” hydroquinone units that adopt a barrel-shaped configuration with variable cavity dimensions depending upon the number of hydroquinone units.[42,43] A critical parameter for the host-guest chemistry of pillararenes is the derivatization of the hydroquinone units. Versatile functionalization of pillar[n]arenes can provide various modified derivatives with anticipated chemical or physical properties for a wide range of applications in supramolecular chemistry.[44–48] There have been several advances in the controlled assembly of proteins with macrocyclic compounds such as calixarenes acting as molecular glue to yield crystalline frameworks.[49–51]

Recently, we have studied the electrostatic self-assembly of Pyrococcus furiosus aFt and a library of cyclophanes.[50] An optimal cyclophane was identified, which formed crystalline...
porous framework through tuning the electrostatic interactions. However, insights into the practical implications of the structures with two differentiated chemical environments have not been described.

In this article, a protein crystal combining two types of molecular hosts is presented. On one hand, aFt and Ft present the ability to naturally host ionic metals. We studied abundantly available horse spleen Ft and aFt (Figure 1a), which are composed of a different ratio of the heavy chain (21 kDa) and light chain (19 kDa) subunits depending upon the type of tissue, and present a pI of ~5.4. The cavity can accommodate up to ~4500 Fe atoms in the form of a hydrous ferric oxide mineral core with a variable amount of phosphate.\[52\] The size of the outer shell of ferritin is 12 nm, while the inner cavity is approximately 8 nm in size (Figure 1a). On the other hand, pillar[5]arene (P10⁺), a cyclophane with ten positive charges, was used as a molecular glue. P10⁺ presents high affinity and selectivity to hydrophobic moieties due to the hydrophobic cavity with an inner diameter of approximately 1.5 nm (Figure 1b). Such porous materials have added advantage of improved mass diffusion properties and can be used to develop efficient catalysts and scavengers for water remediation purposes. We have characterized these cyclophane–protein cage frameworks (CPF) through different techniques, which detail the electrostatically driven self-assembly into highly symmetrical order structures. The CPF adopts a crystal structure where protein cages form a face-centered cubic (FCC) lattice, and the pillar[5]arene hosts are electrostatically bridging the protein lattice and provide additional functionality for the pores between the proteins. Furthermore, CPFs demonstrate selective host-guest chemistry by binding common water pollutants such as methyl orange and heavy metal ions.

The synthesis of P10⁺ has been described elsewhere.\[56\] The formation of large complexes between the protein cages (Ft and aFt) and the molecular host P10⁺ was first studied using dynamic light scattering (DLS) by monitoring the increase in particle count rate as well as the hydrodynamic diameter (Dh). A constant concentration of aFt or Ft (0.1 mgmL⁻¹) in Tris buffer (20 mm, pH 7.5) was titrated with an increasing concentration of P10⁺ (Figure 1c, left). First, Ft shows a sharp intensity increase, which corresponds to the presence of aggregates. This is achieved at [P10⁺] of 10 mgL⁻¹, whereas for aFt, it is achieved at 17.5 mgL⁻¹. The small variation in the point of aggregation can be explained by the significantly smaller molecular weight of aFt due to the absence of the iron oxide core, which results in a higher number of cages per volume. Therefore, a higher P10⁺ concentration is required to induce complex formation. To prove the electrostatic interaction as the key driving force for recognition,\[53,54\] the complexes are disassembled by increasing [NaCl] for both protein cages (Figure 1c, right). This process occurs at similar [NaCl] (i.e., above 50 mm) for both cages. Secondly, Ft shows a change in

![Figure 1. Self-assembly of protein cage–P10⁺ complexes.](image-url)
shows well-resolved diffraction peaks at low NaCl concentrations (0–40 mM). However, increasing the electrolyte concentration results in a decrease in the attractive interactions and prevents assembly formation, which is also in good agreement with the DLS data. With Ft–P10+ complexes (Figure 2b), the Bragg reflections were observed even at higher ionic strength (up to 90 mM). In both cases, the relative peak positions fit with the first allowed reflections of a FCC lattice (Figures 2c and S2). They are assigned to the (111), (200), (220), and (311) reflections, which correspond to $q_{\text{Miller}}/q^* = \sqrt{3}/2$, $\sqrt{4}/2$, $\sqrt{8}/2$, and $\sqrt{11}/2$, respectively. Plotting the square root of the sum of the squared Miller indexes, $(\sqrt{h^2 + k^2 + l^2})$ versus the assigned magnitude of the scattering vector ($q$), results in a linear trend that yields a slope of $26.69 \, \text{Å}^{-1}$, and for the fitted FCC system, a lattice parameter of $a_{\text{FCC}} = 18.6$ Å (Figure 2c, inset). The calculated nearest-neighbor aFt center-to-center distance is 13.2 nm, which is in good agreement with the reported aFt diameter ($d_{\text{at}} \sim 12$ nm) and the observed DLS dimensions (14.7 nm; DLS data in Figure S2). Also, the comparison to a simulated scattering curve from a finite FCC structure confirms the Bravais lattice space group $\text{Fm\tilde{3}m}$ (no. 225). Similar lattice constant and dimensions were determined for Ft–P10+ complexes ($a_{\text{Ft+P10}} = 18.2$ nm and calculated nearest-neighbor Ft center-to-center = 13.1 nm) and is presented in Figure S2.

Figure 2. Structural characterization of (a)Ft–P10+ complexes. SAXS diffractograms measured for a) aFt–P10+ and b) Ft–P10+ complexes at various NaCl concentrations. c) SAXS data for the aFt–P10+ complex at 20 mM of NaCl, compared to the fitted FCC model (offset in the y-direction for clarity). Inset: Square root of the sum of the square of the Miller indexes in the assigned reflections for the FCC structure vs. the measured q-vector positions. Unit cell parameter of $a = 18.6$ Å (space group $\text{Fm\tilde{3}m}$, no. 225). The concentrations of the protein cages (aFt and Ft) and P10+ are constant in all SAXS experiments, that is, 4 and 2.67 mg mL$^{-1}$, respectively; they were mixed in 20 mM Tris buffer (pH 7.5). d) Optical microscopy image of Ft–P10+ crystals showing octahedral habit and sizes over 5 μm. Cryo-TEM image of vitrified aqueous solutions containing e) Ft–P10+ with 20 mM NaCl, f) Ft–P10+ with 80 mM NaCl, and g) aFt–P10+ with 20 mM NaCl. Insets: Corresponding fast Fourier transform (FFT).
By using the hanging drop method (Figure S3), Ft–P10 complexes with octahedral habit and dimensions of approximately 5 μm were observed (Figure 2d). For all other protein cage complexes at different NaCl concentrations, no faceted macrocrystals were observed under optical microscopy (Figure S4). To visualize and further analyze the formed lattices in solution, cryogenic transmission electron microscopy (cryo-TEM) was used. A vitrified sample from the freshly prepared solution of both CPFs, that is, Ft–P10 and aFt–P10+, was imaged (see the Supporting Information for sample preparation). After optimizing the crystallization conditions for microscopy, Ft–P10+ complexes were imaged at 20 mM and 80 mM of NaCl based on better crystallinity observed through SAXS results. A similar selection was also chosen for aFt–P10+ with 20 mM NaCl. Figure 2e–g shows typical small crystallites with a highly ordered arrangement of individual Ft or aFt particles. Low magnification images are shown in Figure S5. Note that samples were prepared at final protein concentration of 4 mg mL⁻¹, and subsequently diluted to 1.33 mg mL⁻¹ after crystal formation, indicating that the structures remain largely unchanged upon dilution.

The pore size of Ft–P10+ crystals ($d_{oct} = 6.2$ nm; Figure S6) allows hosting of relatively large functional moieties within the interstitial space of the crystalline structure. In the CPFs, the voids between the protein cages are only partially occupied by the P10+ bridging the proteins, thus being available for binding suitable small-molecular guests (Figure 3a). The Ft cage, on the other hand, is well known for its ability to oxidize, nucleate, and host various metal ions,[55] which has been utilized for water treatment solutions even at industrial scale.[56,57] To test our hypothesis, the selective binding of organic guest molecules was initially studied by following the changes in the absorption spectra of two dye molecules: methyl orange (MO) and fluorescein. MO is also a typical dyeing agent in the fabric industry as well as a potentially carcinogenic pollutant, and therefore a relevant model molecule for water remediation applications.[58] Further, the dimensions of MO are suitable for binding inside cationic pillararenes, displaying a solvatochromic effect, whereas fluorescein can fit the cavity only partially.

When MO is titrated with free P10+, a clear decrease in the UV-Vis absorption spectrum of MO is observed at 470 nm, together with an increase at 400 nm; this indicates that the MO

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**Figure 3.** Host-guest binding of organic and inorganic materials. a) Schematic presentation of the Ft–P10+ composite unit cell showing full particles (top left) and highlighting the different areas for the simultaneous binding of small organic molecules by the cyclophane (yellow, bottom left) and inorganic materials inside the ferritin cage (green, bottom right). UV-Vis titration of MO (2.5 μg mL⁻¹) with b) P10+ ($[\text{P}10+] = 0–213 \mu$g mL⁻¹) and c) aFt–P10+ ($[\text{P}10+] = 0–213 \mu$g mL⁻¹) shows clear binding. Inset: 400/470 nm absorbance ratio for P10+, aFt–P10+, and aFt–PAMAM-G2. The latter, showing no binding, is plotted as a reference. d) Control titration of fluorescein (2.5 μg mL⁻¹), which is too large to bind efficiently inside the P10+ cavity ($[\text{P}10+] = 0–213 \mu$g mL⁻¹), shows very minor changes in the spectrum. e) Adsorption isotherms of CdII and AsV on Ft–P10+ complexes obtained by inductive coupled plasma–optical emission spectrometry (ICP-OES). Solid lines present Langmuir adsorption isotherm fits. Inset: schematic presentation of the CdII and AsV (generalized as M) adsorption on the ferritin mineral core. f) Elemental analysis of aFt co-loaded with Fe and CdII or AsV. Inset: a schematic presentation of the co-loading inside aFt.
is binding inside the host (Figure 3b). Such changes are in good agreement with previously reported observations of MO-pillar[5]arene complexes.\textsuperscript{50,51} Binding inside the cavity is further supported by i) the broadening of the MO and P10+ NMR signals caused by the interaction, ii) the shielding of the aromatic signals of MO up to +0.88 ppm (Figure S8a), and iii) the deshielding of the aromatic signals of the host upon intake.\textsuperscript{52} A similar change in absorption is observed when the titration is carried out by using the aFt–P10+ crystals. Extracting the crystal-induced background scattering (Figure S9b) yields similar changes in the absorption spectra (Figure 3c). Plotting the absorbance ratios against the host concentration indicates that the crystals are somewhat less efficient binders compared to freely soluble host in terms of the P10+ concentration. This can be expected since the binding of aFt to P10+ can negatively influence the host’s ability to accept guests. However, binding inside the crystals still clearly takes place. If the aFt crystals are formed with a generation two polyamidoamine dendrimer (PAMAM–G2), which is not able to complex MO, there are no changes in UV-Vis spectra during the titration (Figure S9i). The UV-Vis titration data show only minor changes for the fluorescein absorbance upon binding to the P10+ (Figure 3d). Together with the very limited shift of the NMR signals of fluorescein (Figure S8b), the results verify our hypothesis that small molecules can be selectively bound by free hosts, provided that the host is able to interact efficiently with the guest molecule.

To exploit the ferritin cavities in the complexes, we studied their ability to bind toxic metal ions such as arsenate. This is relevant as groundwater contaminated by cadmium and arsenic is a major problem, especially in South (East) Asia, and typical adsorption purification methods rely on the binding or co-precipitation with iron oxide or hydroxides.\textsuperscript{61} Native ferritin hosts possess an amorphous iron hydroxide (FeOOH) core, which is known to adsorb heavy metals, such as arsenate and cadmium. This approach of binding arsenic was also used very recently for leukemia cells featuring high CD71 expression, which resulted from the loading of these inorganic materials, we turned to porous CPFs with iron and Cd II and As V pollutants, enabled by the porous protein–molecular host network. We emphasize that these results not only demonstrate that the system is multifunctional but also pave the way for the successful design of protein assemblies with confined and synergetic chemical environments. Such supramolecular approaches and binding capabilities can be extended to other molecular guests like cyclodextrins, with a rich host-guest chemistry.

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**Conflict of Interest**

The authors declare no conflict of interest.

**Data Availability Statement**

The data that support the findings of this study are available from the corresponding author upon reasonable request.

**Keywords:** host-guest chemistry · protein cages · protein crystals · pillararene · water remediation
