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Anion-templated 2D frameworks from hexahydroxytriphenylene†

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Hexahydroxytriphenylene (HHTP) forms 2D frameworks through O–H...anion hydrogen bonds with a range of anions. In all cases, 1:1 co-crystals of HHTP and tetraalkylammonium-anion salts are obtained, which have a layered 2D structure. When HSO_4^- was used as anion, an unprecedented methylation reaction was observed giving crystals containing the methylsulfate anion (MeOSO_3^-).

The vast majority of research into the supramolecular chemistry of anions has focused on their solution phase behaviour spurred by the key role negatively-charged species play in a range of biological and environmental processes.¹ However recently, a handful of reports have demonstrated that anions can also be used to template the formation of self-assembled structures.²

While the interaction of anions with O–H hydrogen bond donors has received comparatively little attention, the groups of Smith,³ and Kass and Wang,⁴ have demonstrated that receptors containing these groups can be potent anion hosts. Very recently, White and MacLachlan have demonstrated that O–H...anion hydrogen bonds can also be used to form extended structures including 1D polymers and nanotubes.⁵

Hexahydroxytriphenylene (**HHTP**, Scheme 1) is an important precursor for a range of supramolecular framework structures. When reacted with transition metals in the presence of base, **HHTP** gives two or three-dimensional metal organic frameworks (MOFs) containing the sextuply-deprotonated form of the ligand [*i.e.* triphenylene-hexakis(olate)].⁶ Some of these MOFs show very high electrical or proton conductivity. Additionally, **HHTP** has been used to form highly stable covalent organic frameworks (COFs) with boronate derivatives, which show impressive gas storage properties.⁷

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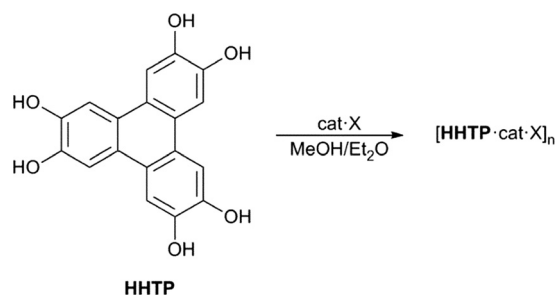
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Herein, we show that O-H...anion hydrogen bonding from the six hydroxyl groups of **HHTP** can be used to prepare 2D layered structures. The complexes form sheets with one sheet comprised of anions and **HHTP** ligands, and the second made up of cations and solvent molecules.

We initially tried to prepare **HHTP** directly from catechol using ammonium persulfate as oxidant,⁸ but recovered only unreacted catechol. Instead we found that high purity **HHTP** could be readily prepared by FeCl₃-mediated oxidation of 1,2-dimethoxybenzene (veratrole),⁹ followed by demethylation using BBr₃ (see ESI†).

We attempted to determine if **HHTP** binds anions in solution using ^1H NMR spectroscopy, but were unable to obtain quantitative data. We have found **HHTP** only shows solubility in acetone or polar solvents (alcohols, DMSO), however in polar solvents, solution binding is too weak to be observed, while addition of anions to **HHTP** in acetone causes precipitation. Previous studies have shown that the interactions between catechol³ or catechol-containing^{5b} molecules and anions are modest in acetonitrile, so we suggest that more competitive solvents such as DMSO or alcohols disrupt the interactions between **HHTP** and anions allowing them to remain soluble. Less competitive acetone presumably does not do this, and therefore some kind of oligomeric or polymeric **HHTP**-anion complex forms and precipitates from solution.



Scheme 1 Structure of HHTP and its assembly into anion-templated 2D polymeric structures (cat = tetraalkylammonium cation, X = anion).

We then investigated the crystallisation of **HHTP** with anions: dissolving 1:1 stoichiometric mixtures of **HHTP** and TBA salts or TEA-Cl in methanol gave clear solutions, and vapour diffusion of diethyl ether into these solutions gave single crystals when $X = \text{Cl}^-$, I^- , OAc^- or H_2PO_4^- and twinned crystals when $X = \text{HSO}_4^-$. When $X = \text{Br}^-$, crystals were obtained but diffraction studies show that these appear to be modulated (see ESI†).

Single crystal X-ray diffraction† (SCXRD) studies revealed that all of the crystals had the formula $[\text{HHTP}\cdot X\text{-alkylammonium salt}]_n$ and contained very similar supramolecular structures (when we started with TBA- H_2SO_4 , we obtained crystals containing the methylated anion MeOSO_3^- , see later). In all crystals, **HHTP** and the anion form a 2D sheet architecture, which is assembled through O-H $\cdots$ anion hydrogen bonding between **HHTP**'s O-H groups and the anion, as well as intermolecular O-H $\cdots$ O hydrogen bonds between adjacent **HHTP** molecules. This 2D sheet alternates with layers comprised of the alkylammonium cations (these various levels of supramolecular assembly are shown in Fig. 1 for the structure of $[\text{HHTP}\cdot\text{TBA}\cdot\text{Cl}]_n$; full diagrams of all of the complexes are provided in the ESI†).

These O-H $\cdots$ anion hydrogen bonds are typically short: with H $\cdots$ X distances as little as 1.74 Å, 64%vdW (where %vdW refers to the distance as a percentage of the sum of the van der Waals radii¹⁰ of hydrogen and the interacting atom on the anion§). Generally the H $\cdots$ X distances expressed as %vdW increase slightly with decreasing anion basicity, so that the mean H $\cdots$ X bond length is 67%vdW for OAc^- , 68%vdW for H_2PO_4^- , 71%vdW for MeOSO_3^- , 77%vdW for Cl^- and 82%vdW for I^- . The intermolecular O-H $\cdots$ O hydrogen bonds are comparable in length in all structures, and are relatively short, H $\cdots$ O distances: 1.91–2.13 Å, 70–79%vdW_{H,X}.

More specifically, in the three complexes containing halide anions, $[\text{HHTP}\cdot\text{TBA}\cdot\text{Cl}]_n$, $[\text{HHTP}\cdot\text{TBA}\cdot\text{I}]_n$ and $[\text{HHTP}\cdot\text{TEA}\cdot\text{Cl}]_n$, each anion receives three O-H hydrogen bonds from three different **HHTP** molecules (Fig. 2a). In $[\text{HHTP}\cdot\text{TBA}\cdot\text{H}_2\text{PO}_4]_n$ (Fig. 2b) and $[\text{HHTP}\cdot\text{TBA}\cdot\text{MeOSO}_3]_n$ (Fig. 2c), a similar structure is observed with the larger oxoanions again sitting between three **HHTP** molecules. In the crystals containing MeOSO_3^- , the anion receives one hydrogen bond from each of the three **HHTP** moieties, while in the H_2PO_4^- structure, two **HHTP** molecules and a methanol solvate act as O-H donors to the anion, which in turn donates two hydrogen bonds to **HHTP**. The structure of $[\text{HHTP}\cdot\text{TBA}\cdot\text{OAc}]_n$ (Fig. 2d) is slightly different with the acetate anion and a methanol solvent molecule located in a pocket between four **HHTP** molecules. In this pocket, the acetate anion receives hydrogen bonds from two of these molecules, as well as the methanol solvent, while the other two **HHTP** groups in this pocket donate H-bonds to the methanol molecule.

Despite these differing **HHTP** $\cdots$ anion bonding arrangements, 2D sheets are formed in each case. In the structure of $[\text{HHTP}\cdot\text{TBA}\cdot\text{Cl}]_n$ (Fig. 1), these sheets are close to planar (the angle between the mean plane of the **HHTP** moieties is 9°), while in the other TBA-containing structures they show varying degrees of corrugation (angles between mean planes range from 27–40°, see Fig. S13–S18 of the ESI†). In the structure containing the TEA cation, the corrugation is even more pronounced (angle between mean planes = 50°). Presumably this buckling, or lack of it, occurs to maintain favourable hydrogen bonding interactions between **HHTP** and the anion, and between adjacent **HHTP** molecules, while also accommodating the differently-sized anions and cations as well as the solvent or absence of it.

It is interesting that the same gross structure is observed when TEA-Cl or TBA-Cl is used; this is in contrast to related systems prepared from tetrahydroxytriptycene and anions, in

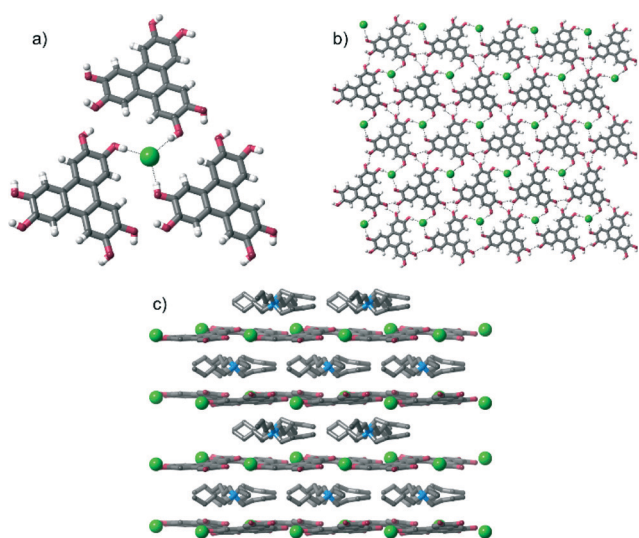


Fig. 1 Views of the solid state structure of $[\text{HHTP}\cdot\text{TBA}\cdot\text{Cl}]_n$: a) environment around Cl^- anion; b) 2D sheet formed from **HHTP** molecules and anions; c) packing viewed along c axis (with hydrogen atoms omitted for clarity). TBA cation disorder is omitted for clarity.

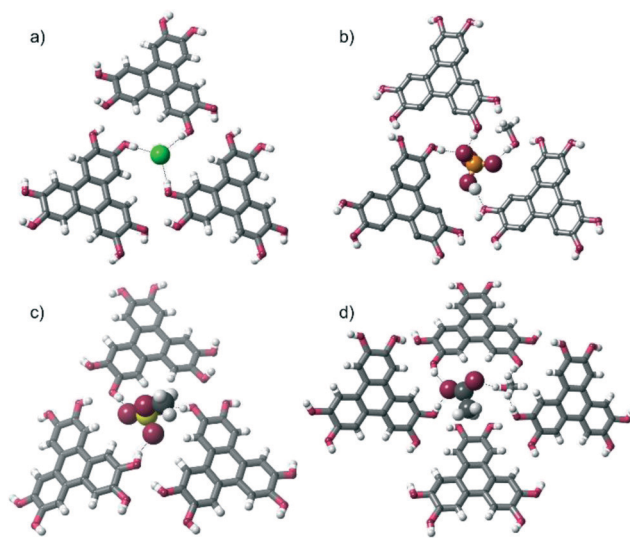


Fig. 2 Comparison of the anion environments in solid state structures of $[\text{HHTP}\cdot\text{TBA}\cdot X]_n$: a) $[\text{HHTP}\cdot\text{TBA}\cdot\text{Cl}]_n$ (the anion environments in $[\text{HHTP}\cdot\text{TBA}\cdot\text{I}]_n$ and $[\text{HHTP}\cdot\text{TEA}\cdot\text{Cl}]_n$ are very similar); b) $[\text{HHTP}\cdot\text{TBA}\cdot\text{H}_2\text{PO}_4]_n$; c) $[\text{HHTP}\cdot\text{TBA}\cdot\text{MeOSO}_3]_n$; d) $[\text{HHTP}\cdot\text{TBA}\cdot\text{OAc}]_n$.



PXRD patterns for the bulk materials containing TBA cations were generally consistent with those simulated from the single crystal structures, albeit with some loss of crystallinity upon drying (Fig. 3a and S19–S24, see ESI† for further details regarding PXRD experiments). In the TEA-containing structure [HHTP-TEA-Cl]_n there is a significant loss of crystallinity upon drying (Fig. 3b), which is perhaps not surprising given the large

In conclusion, we have demonstrated that O-H $\cdots$ anion hydrogen bonds can be used to assemble the triphenylene-containing ligand **HHTP** into framework materials. The materials contain 2D sheets formed from **HHTP** and anions, which adopt different levels of corrugation dependent on the anion and cation size. Despite the fact that O-H $\cdots$ anion interactions are weak in solution (in polar solvents), they reproducibly give crystalline frameworks in good yields. These results demonstrate the importance of weak interactions in crystal engineering and we suggest that if anion-templated materials can be prepared using stronger anion $\cdots$ receptor interactions, these systems may prove useful for a range of materials applications.

§ This parameter allows useful comparison between hydrogen bonds to differently-sized anions. It is relatively imprecise, as the position of hydrogen atoms is poorly-determined by SCXRD and so values are given to only two significant figures. For a more detailed discussion, see the ESI,[†] and N. G. White, C. J. Serpell and P. D. Beer, *Cryst. Growth Des.*, 2014, 14, 3472–3479.

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