### Acknowledgments:

Polish National Science Centre MAESTRO grant decision number DEC2012/04/A/ST5/00609, DSM grant No. 501-D112-86-DSM-115 100 are acknowledged.

#### References:

Coppens, P. (1997). X-Ray Charge Densities and Chemical Bonding; International Union of Crystallography.

Dittrich, B.; Weber, M.; Kalinowski, R.; Grabowsky, S.; Hübschle, C. B.; Luger, P. (2009). Acta Crystallogr. 65 (6), 749–756. Jarzembska, K. N.; Dominiak, P. M. (2012). Acta Crystallogr. A. 68 (Pt 1), 139–147.

Keywords: 9-aminoacridine hemihydrate, charge density, TAAM refinement

## **MS21-P02**

# Preparation of non-covalent organic frameworks using dodecasubstituted porphyrins

Keith Flanagan<sup>1</sup>, Mathias O. Senge<sup>1</sup>

 School of Chemistry, Trinity College Dublin the University of Dublin, Dublin, Ireland

### email: kflanaga@tcd.ie

Nowadays the use of porphyrins to create highly porous, sponge-like frameworks is a well-studied field, established by the seminal work of Byrn et al. coining the phrase "porphyrin sponge" in 1991.[1] This initial study used a porphyrin host, trapping a variety of guests within the crystal lattice. Since then, many contributions have focused on the manipulation of planar porphyrins containing a variety of functional groups (namely carboxylic acids or halogens) to obtain directly linked porphyrin frameworks. [2] However, as of yet, there are no specific examples of the use of highly distorted porphyrins [3] being used to obtain such highly porous materials. In this work, we have synthesized a series of 5,10,15,20-tetraaryl-2,3,7,8,12,13,17,18-octaethylporphyrins with alternating halogen, alkyl, and aryl substitution patterns to probe the effects of non-planarity on the formation of non-covalent organic frameworks.

The target porphyrins were synthesized according to a modified Lindsey condensation reaction followed by metal(II) insertion. To investigate substituent interactions, modified aldehydes were used. For halogen effects, a family of derivatives with F, Cl, Br or I in either the ortho- or para-position of the meso-aryl residues were selected. Hydrogen bonding interactions were examined by incorporating a nitrile group. Steric effects were examined through the incorporation of a benzyloxy group at either the *meta*- or *para*-positions. Finally, investigations were conducted into the use of arm-extended porphyrins using a triazole linker group to further extend their functionality. Solid-state structures were determined via small molecule X-ray crystallography to elucidate the major distortion modes and to characterize any non-covalent interactions. These were then compared to their planar derivatives to determine effects the substitution patterns may have on the formation of non-covalent organic frameworks.

### References:

- [1] Byrn, M. P. et al. (1991). J. Am. Chem. Soc., 113, 6549–6557.
- [2] Goldberg, I. (2008). Cryst. Eng. Comm., 10, 637-645.
- [3] Senge, M. O. et al. (2015). Chem. Commun., 51, 17031-17063.

Keywords: Porphyrin, Halogens, Macrocycles