CrystEngComm

Accepted Manuscript

This is an *Accepted Manuscript*, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. We will replace this *Accepted Manuscript* with the edited and formatted *Advance Article* as soon as it is available.

You can find more information about *Accepted Manuscripts* in the Information for Authors.

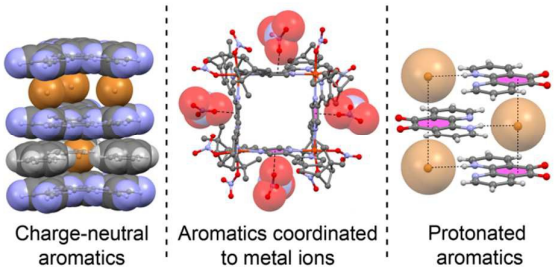
Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal's standard Terms & Conditions and the Ethical guidelines still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this *Accepted Manuscript* or any consequences arising from the use of any information it contains.

ROYAL SOCIETY OF CHEMISTRY

www.rsc.org/crystengcomm

TOC graphic

This highlight article summarizes some of the fundamental aspects of the anion–π interaction leading to several design strategies for generating new receptors. In the main body we highlight some relevant examples that illustrate the viability of these strategies and the importance of anion–π interactions in crystal engineering



CrystEngComm Accepted Manuscript

Journal Name

ARTICLE

Towards design strategies for anion–π interactions in crystal engineering

Received 00th January 20xx,
Accepted 00th January 20xx

DOI: 10.1039/x0xx00000x

www.rsc.org/

Antonio Bauzá,[a] Tiddo J. Mooibroek*,[b] and Antonio Frontera*,[a]

For well over half a century part of the scientific community has been committed to understand and predict how molecules recognize each other. This subject of unceasing interest, 'supramolecular chemistry', relies on the understanding of noncovalent interactions. Among the noncovalent forces, the anion–π interaction has attracted increasing attention ever since its inception about two decades ago. This highlight article first summarizes some of the fundamental aspects of this interaction leading to several design strategies. In the main body we highlight some relevant examples that illustrate the viability of these strategies and the importance of anion–π interactions in crystal engineering.

1. Introduction

Supramolecular chemistry is a highly multidisciplinary field of research. Its development has had a profound impact on the growing efficacy and success of constructing molecular assemblies of different sizes, shapes and functionalities.[1,2] Supramolecular chemists depend on the understanding of the non-covalent forces that govern molecular recognition phenomena. The design strategy of constructing interaction-complementary receptors for a target host molecule has resulted in host-guest systems with high affinities ($K_{ass} \geq 10^4$ M^{-1}),[3,4,5] even in very competitive media such as water (i.e., small, polar and protic).[6] In the solid state the interaction-complementary approach is even more pronounced, as competing solvent molecules are mostly absent. As a result, the manner in which crystalline materials are organized can be seen as a reflection of strong and weak intermolecular forces balancing themselves towards a stable state. Strong and highly directional interactions such as hydrogen bonding and σ-hole bonding,[7,8,9,10,11,12,13,14,15] together with less directional forces like ion pairing are indeed commonly used to engineer crystals.

Interactions involving π-systems (e.g. aromatic rings, nucleobases) are very relevant for molecular interactions. For instance C–H/π, π–π and cation–π interactions are important in protein structure and enzyme catalysis.[16,17,18,19] During the past two decades the anion–π interaction has been established as the attractive interaction between an anion and an electron deficient π-system.[20,21,22,23,24,25] The birth of this increasingly-established interaction can be pinpointed to several publications.[26,27,28,29,30] In 1996 Woollins *et al.* described a close contact between Cl^- and the aromatic ring $[S_4N_3]^+$ in the crystal packing of thiotrithiazylium chloride and named this as a "π-facial interaction".[26] Other early publications are from Schneider *et al.*[27,28] and Yamabe *et al.*[29,30] Recently, this interaction has been described as a subclass of π-hole bonding interactions.[31,15b] Various theoretical[32,33,34,35] and experimental[36,37,38,39,40] investigations showcase that anion–π interactions can be of structural and functional significance. Indeed, the relevance of the anion-π interaction has been noticed in biochemical processes and fields like environmental chemistry, medicine and supramolecular chemistry.[20,41,42,43,44,45]

This highlight article is not just a bibliographic review of the literature related to anion–π interactions since several surveys have been written for that purpose.[20,24,46,47,48,49] Instead, we showcase herein that a fundamental understanding of the interaction leads to design strategies that can be exploited in crystal engineering.

2. Fundamental aspects of anion-π interactions

2.1 Physical nature

The physical nature of the anion–π interaction has been studied and rationalized using high level *ab initio* calculations and several partition energy schemes.[50,51] The general assumption is that ion-induced polarization and electrostatic interactions are the main contributors to the anion-π interaction.[52,53,54]

The molecular polarizability of a compound is denoted as $\alpha_{||}$. The π-cloud on an aromatic compound is readily polarized by an anion, leading to an 'induced dipole' (see Fig. 1A). The anion can then interact with this induced dipole. This polarization contribution to the total interaction energy of an anion-π contact can be substantial.[50–53] Actually, the

[a] Departament de Quimica, Universitat de les Illes Balears, Crta. de Valldemossa km 7.5, 07122 Palma de Mallorca, Baleares, Spain. E-mail: toni.frontera@uib.es
[b] School of Chemistry of the University of Bristol, Cantock's Close, BS8 1TS, Bristol (United Kingdom). E-mail: chtjm@bristol.ac.uk

† Footnotes relating to the title and/or authors should appear here.
Electronic Supplementary Information (ESI) available: [details of any supplementary information available should be included here]. See DOI: 10.1039/x0xx00000x

CrystEngComm Accepted Manuscript

Please do not adjust margins

electrostatic repulsion anticipated in the benzene···Cl^- pair is largely overruled by the favourable induced dipole interaction, resulting in a nearly negligible overall repulsive interaction energy of +2.4 kcal/mol.[55] Induced polarization interactions can also rationalize the dual binding mode exhibited by arenes with a marginal permanent quadrupole moment perpendicular to the ring plane (Q_{zz} in Buckinghams, B).[56,57] That is, molecules with very small Q_{zz} values such as 1,3,5-trifluorobenzene (Q_{zz} = 0.57 B) and s-triazine (Q_{zz} = 0.90 B) are able to interact with both anions and cations because the electrostatic term is negligible and the interaction is thus dominated by the polarization term (always favourable).

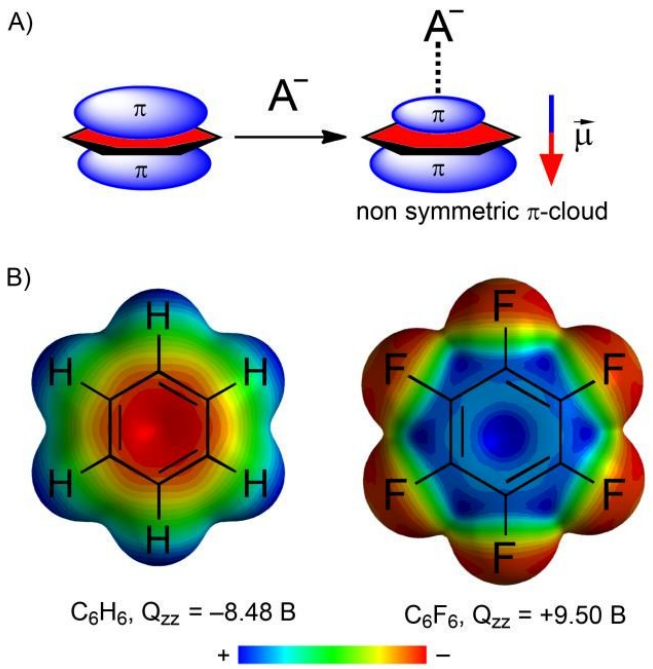


Fig. 1 Illustrations to aid understanding of the physical nature of anion-π interactions: (A) Schematic representation of the ion-induced dipole. (B) Molecular electrostatic potential (MEP) of benzene and hexafluorobenzene; MEP surfaces computed at the B3LYP/6-31G* level of theory.

The electrostatic contribution to an anion–π interaction correlates well with the permanent quadrupole moment perpendicular to the ring plane (Q_{zz} in Buckinghams, B). This physical property is a measure of the electron deficiency of an aromatic system. As one might expect, a more electron poor aromatic tends to display stronger binding to anions. The magnitude and sign of Q_{zz} depend on the substitution pattern on a given aromatic. For example (see Fig. 1B), the negative Q_{zz} found for benzene (Q_{zz} = –8.48 B) predicts a repulsive interaction with anions. If, however, H is replaced by F (i.e. hexafluorobenzene) the Q_{zz} becomes positive (9.50 B) and the interaction becomes attractive.

The anion–π interaction has also been examined using molecular orbital analysis, and compared to the related cation–π interaction.[58,59] Interestingly, a totally different picture emerged: the atomic orbitals of a cation do not participate in the molecular orbitals of the cation–π complex, while conversely, the atomic orbitals of an anion contribute significantly to the molecular orbitals of the anion–π complex.[58,59]

2.2. Energies and directionality

The usefulness of an intermolecular interaction for molecular engineers relies in part on its strength and directionality. For the solid state (where competing solvent molecules are mainly absent), energies computed using high level theoretical models provide a reliable estimate of the actual enthalpic component of an anion-π interaction. Various such inquiries have revealed that a single anion-π interaction varies in strength from very weak (–3.6 kcal/mol for 1,4-difluorobenzene···Cl^- complex) to strong (–24.0 kcal/mol for 1,3,5-trinitrobenzene···Cl^-) compared to hydrogen bonding (typically –2-25 kcal/mol).[60]

In light of such estimated binding enthalpies it can be expected that anion-π interactions are directional. The geometry that is intuitively preferred can be seen as a 'T-shaped' geometry, where the anion is located above/below the centre of an aromatic ring at a ring plane–anion angle of 90°. However, the rather wide electropositive region above/below an aromatic implies that lateral deviation from this geometry has a small energy penalty. The size of the electropositive potential may thus impede on the directionality of anion-π interactions.

A convenient source of information regarding the actual directionality of intermolecular interactions is contained within the ever-growing Cambridge Structure Database (CSD). It is indeed commonly assumed that a proper statistical analysis of the CSD can unveil the geometric preferences of a given supramolecular interaction. This assumption stems from the intuition that even weak intermolecular forces must somehow manifest themselves in the way molecules pack in crystals.

Some CSD surveys have shown close proximity of electron rich/anionic entities above/below electron deficient (hetero) aromatics.[20,61,62,63] The directional character of anion-π (but also lone-pair-π)[64] interactions has only recently been demonstrated by a thorough CSD evaluation of interactions between anionic/electron rich atoms and a pentafluorophenyl *vs*. a phenyl ring.[64] The method used in these inquiries takes account of the non-spherical volume of the arene and corrects for a random scattering of the data. The reasoning behind the method is akin to the 'cone-correction method'[65] that is known to be essential for a proper evaluation of the geometric preferences of hydrogen bonding.[60] These studies clearly showed that the predicted 'T-shaped' binding mode is preferred for pentafluorophenyl rings, while a clustering of data near the phenyl's H-atoms was observed.[64]

2.3. Towards design strategies

From the above it is clear that to engineer anion–π interaction, the π–binding units should have a large molecular polarizability ($\alpha_{||}$) and a large and positive quadrupole moment (Q_{zz}).

The polarizability can be somewhat tuned by the size of the aromatic; larger conjugated systems (e.g. naphtalene) are more polarizable than smaller π-systems (e.g. benzene).

CrystEngComm Accepted Manuscript

This journal is © The Royal Society of Chemistry 20xx

Please do not adjust margins

Enlarging the permanent quadrupole moment of an arene can be achieved in several ways. Attachment of strong electron withdrawing groups like $-NO_2$ and -CN leads to large positive Q_{zz} values (e.g. Fig. 2A). For benzene derivatives this strategy is somewhat limited because it is difficult to install more than three such groups. Using polycyclic aromatic compounds enables one to attach more electron withdrawing groups, with the added advantage of increased molecular polarizability (see e.g. Fig. 2A). An alternative way of rendering an arene ring electron deficient is to have it partake in a cation-π interaction. The induced positive potential on the ring can then interact favourably with anions to give strong 'cation–π–anion complexes' in which the arene effectively bridges the ion pair.[66,67,68,69,70,71,72] These ternary complexes exhibit large binding energies and shorter equilibrium distances than those in the corresponding binary ion–π complexes. Alternatively, the ring itself can contain a positive charge, such as in tropylium or protonated azines.

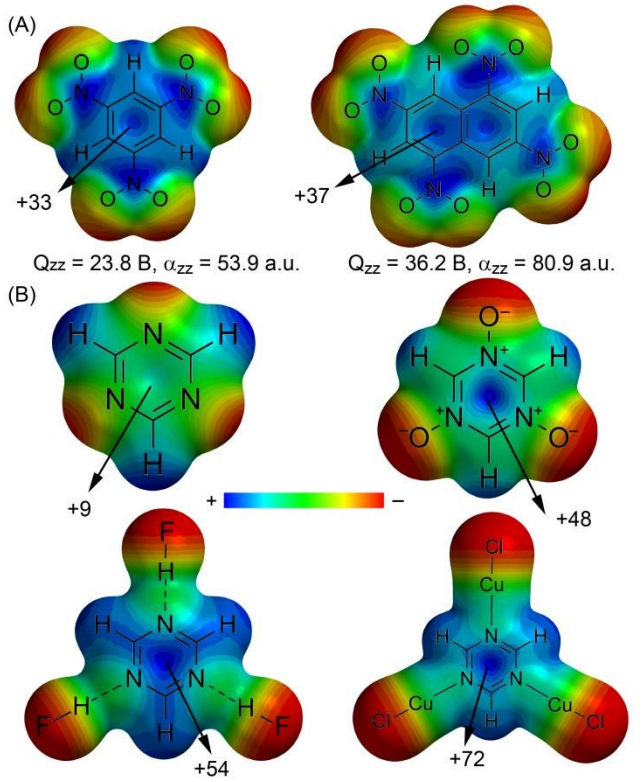


Fig. 2 Illustrations of design strategies to obtain strong anion-pi interactions: (A) MEP of 1,3,5-trinitrobenzene and 1,3,5,7-tetranitronaphtalene, values of MEP, quadrupole moments (Q_{zz}) and molecular polarizabilities (α_{zz}). (B) MEP of s-triazine, [s-triazine···3HF], [s-triazine···3Cu^{I}Cl] and 1,3,5-triazine 1,3,5-trioxide. MEP surfaces computed at the B3LYP/6-31G* level of theory. MEP values over the ring centre in kcal/mol.

N-substituted heteroaromatics can be particularly useful to generate anion-π interactions. While some (e.g. s-triazine, Fig. 2B) already have a positive electrostatic potential over the ring centre, the MEP value can be greatly enhanced when a N-atom is bound to some positively polarized entity or when it is oxidized to an N-oxide. As is illustrated in Fig. 2B for s-triazine, hydrogen bonding, coordination to a metal ion and oxidation all lead to a larger positive MEP value over the ring centre.[48,73,74,75,76,77] Particularly useful about N-heterocyclic arenes is that their coordination chemistry can be exploited to construct larger molecular assemblies. Another design factor is the additivity of the anion–π interaction. For example, for a series of complexes between Cl^-/Br^- and some *s*-triazines it was computed that the ternary "sandwich" complexes were twice as stable as the corresponding binary complexes.[78] Thus, designing systems that can encapsulate an anion using multiple anion-π contacts seems advantageous.

3. Highlighted structures

3.1 General remarks and overview

We have chosen to highlight several structures where anion-π interactions were intended and/or closely studied by the authors of the original manuscripts. For ease of reference we use the CSD reference codes in graphical renderings of these examples. In addition to reported examples, we examined recent entries (post November 2010) in the CSD for particularly evident anion-π interactions. That is, structures where the van der Waals shell of an atom belonging to an anion overlaps with all six atoms of a six-membered aromatic ring comprised of C and/or N.[‡] Nearly all of these close contacts have been overlooked by the authors of the original manuscripts, highlighting just how prevalent yet non-canonical anion-π interactions are. An asterisk (*) has been added to their CSD reference codes to demarcate them from the structures where anion–π interactions were clearly intended and/or studied. Occasionally we also discuss some theoretical studies that are related to the structure highlighted and/or some interesting experimental findings reported by the authors.

Shown in Fig. 3 is an overview of the aromatics used in the crystal structures we mention in this work. The selected structures are grouped under three main headings, based on the type of aromatic used, namely: formally charge neutral arenes rendered electron deficient by covalently attached groups (A), arenes that are coordinated to a metal ion (B), and arenes that bear a formal positive charge. For completion we also indicate in what coordination environment the metal ions are.

3.2 Charge-neutral aromatics

Pyrazine-N,N'-dioxide (PDO) is formally charge-neutral, yet has a substantial electron deficient centre due to the high degree of polarization of the N=O bonds. This has been exploited by using PDO as a linking component in octacyanotugstenate-based networks.[79] The hybrid networks are co-organized by two types of interactions, as indicated in Fig. 4: anion-π interactions between the pyrazine ring centre and a single N_{CN} atom of $[W(CN)_8]^{3-}$, and an interaction where four N_{CN} atoms point towards the $N_{pyrazine}$ atoms.

CrystEngComm Accepted Manuscript

Please do not adjust margins

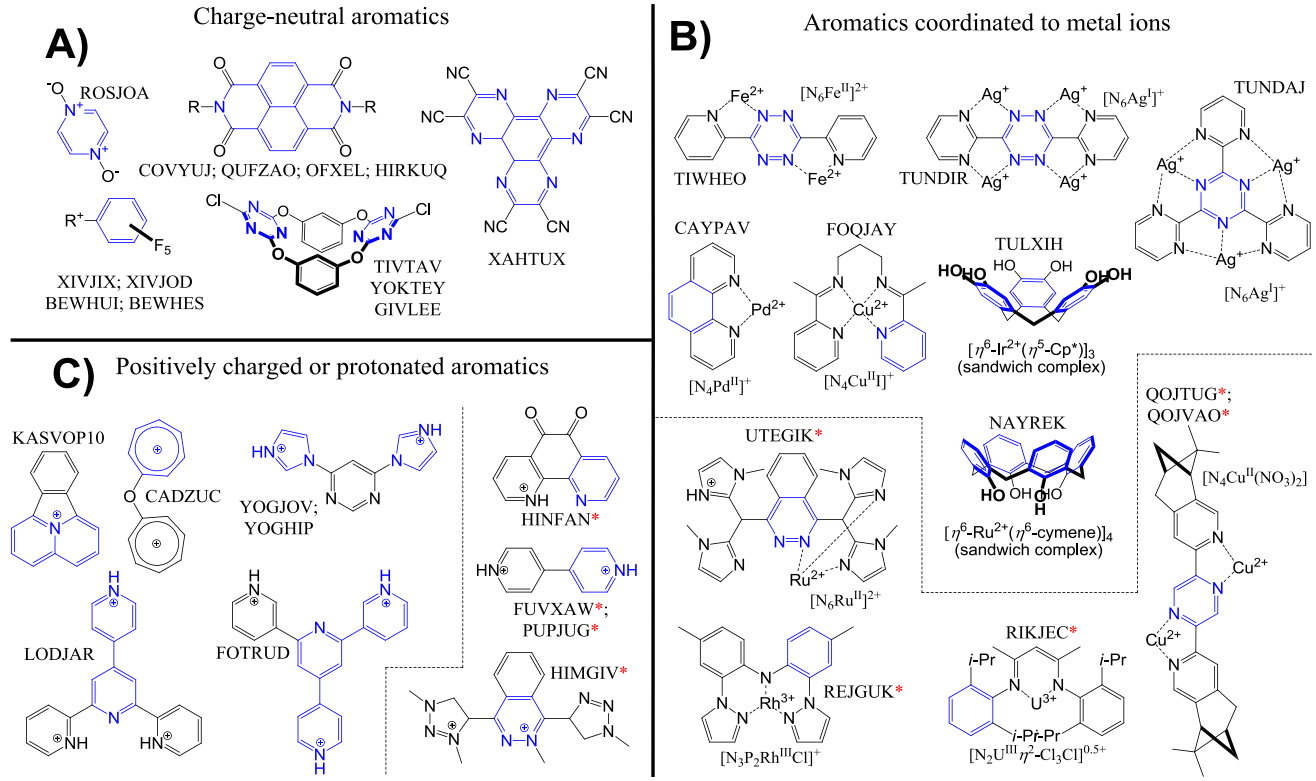


Fig. 3 Overview of examples (including CSD reference codes) highlighted in this paper. Three classes have been distinguished where the interacting aromatic is formally charge neutral (A), coordinated to a metal ion (B) or where the aromatic bears a formal positive charge (C). The part of the aromatic involved in an anion-π interaction is highlighted in blue. The red asterisk (*) accompanying some CSD reference codes denote that the structure was found by the CSD search[‡] and was not intended/studied by the authors of the original work.

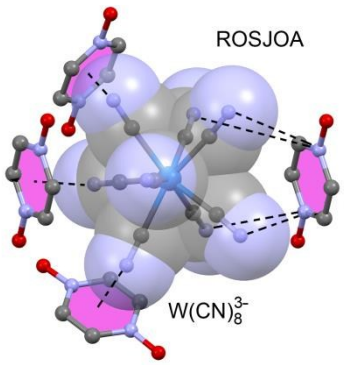


Fig. 4 Fragment of the X-ray crystal structure containing Pyrazine-N,N′-dioxide moieties. The relevant anion–π interactions are indicated by dashed lines. The CSD reference code is indicated. H-atoms omitted for clarity.

The electron deficient pentafluorophenyl (PFP) group is also a good building block for generating anion-π interactions.[64] Giese et al.[80,81,82] have exploited this experimentally by focusing on purely organic salts where the positive charge is located some distance away from the PFP group. From their work it is evident that the position of an anion above the PFP π-system can be tuned. Indeed, in the solid state structures of [N-(pentafluorobenzyl)pyridinium]$^+$[X]$^-$ salts, the way in which the anion is encapsulated by C–H···X hydrogen bonds and anion–π contacts depends on the anion.[81]

In this particular system, both π-acidic rings (C_6F_5 and pyridinium moieties) are involved in the anion–π contacts (see Fig. 5). For X = I_3^- and PF_6^-, there are cooperative anion–π interactions between the two electron-deficient arenes and the anion located between them (Fig 5A), while for X = Br^- or BF_4^- (Fig. 5B), this cooperativity is absent.

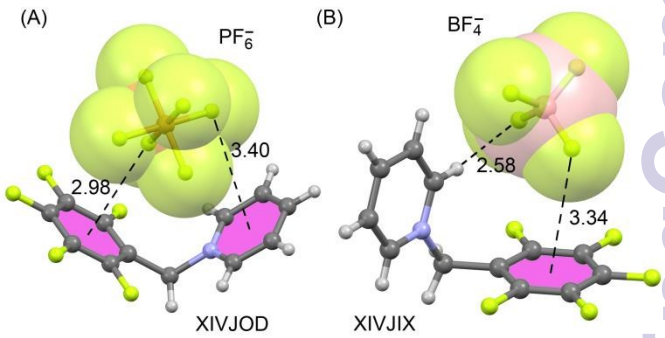


Fig. 5 X-ray crystal structures of pentafluorophenyl derivatives. The CSD reference codes are indicated. Distances in Å.

CrystEngComm Accepted Manuscript

This journal is © The Royal Society of Chemistry 20xx

Please do not adjust margins

In a related investigation[82] they used simple triphenyl(pentafluorobenzyl)-phosphonium and bis(pentafluorobenzyl)-phosphonium salts. The X-ray structures provided crucial data on the influence of anion size on the molecular structure of cations containing two adjacent electron-deficient rings (see Fig. 6).

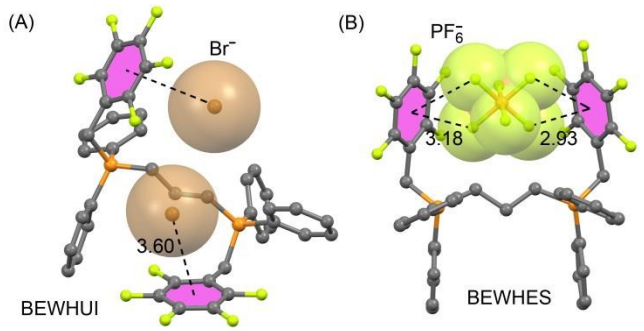


Fig. 6 X-ray crystal structures of pentafluorophenyl derivatives. The CSD reference codes are indicated. Distances in Å. H-atoms omitted for clarity.

In particular, whereas bromide anions interact by means of anion-π interactions in a 1:1 mode with the pentafluorobenzene unit Z-configured (Fig. 6A), the bulkier anions iodide, tetrafluoroborate, and hexafluorophosphate (Fig. 6B) result in a 1:2 tweezer-like anti-configuration in which the anion interacts simultaneously with two pentafluorobenzene units. Apparently, when the spatial separation of the two electron-deficient rings matches the size of the anion, the formation of two concurrent anion–π interactions induce a conformational change from the *anti*-form observed for the smaller anion to the tweezer-like *syn*-form for the greater one.

The naphthalenediimide (NDI) core readily seems to engage in anion-π interactions, likely because NDI is both electron deficient and easily polarized (large, hence large $\alpha_{||}$). Several groups used NDI to construct fascinating assemblies in the solid state and several works deserve special attention. Liu *et al.*[83] have generated a panchromatic hybrid crystal of iodoplumbate nanowires and J-aggregated NDIs. The organic–inorganic hybrid is constituted by anionic iodoplumbate nanowires that strongly bind to the NDIs by means of charge-assisted anion–π interactions (see Fig. 7).

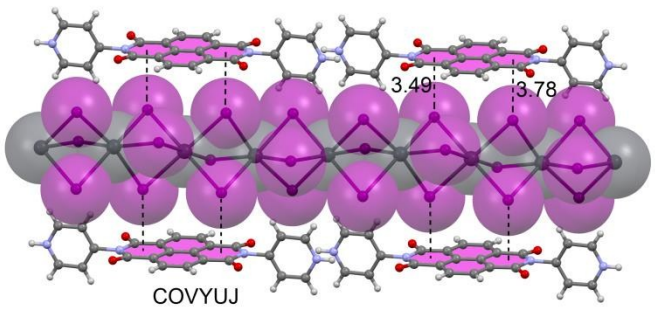


Fig. 7 X-ray structure of the hybrid crystal of iodoplumbate nanowire and J-aggregated NDIs. The CSD reference code is indicated. Distances in Å.

The material has photo-induced long-lived charge-separated states (also when irradiated with sunlight), making this study interesting for research in the area of solar cells and photocatalysis. The solid-state UV-Vis diffuse reflectance spectral analysis revealed that the crystal is a panchromatic hybrid with a broad absorption band from 200 nm to 800 nm, making this material much 'darker' than other iodoplumbate nanowires.[84,85] It is worthy to note that the long-lived charge separation in this hybrid material is basically due to the intense and panchromatic absorption and the close contacts between organic and inorganic units through anion–π interactions that aid the electron transfer (see Fig. 7).

Kumar *et al.*[86] attached two phosphonium groups to an NDI unit (hence 'NDI(PPh_3)$_2$') and were able to characterize the radical cation [(NDI(PPh_3)$_2^{\bullet+}$)BPh_4^-] and its extraordinary π-acidic precursor [(NDI(PPh_3)$_2^{2+}$)·2BF_4^-]. In this latter structure (see Fig. 8) the NDI unit is tightly sandwiched in between two BF_4^- anions through anion-π contacts.

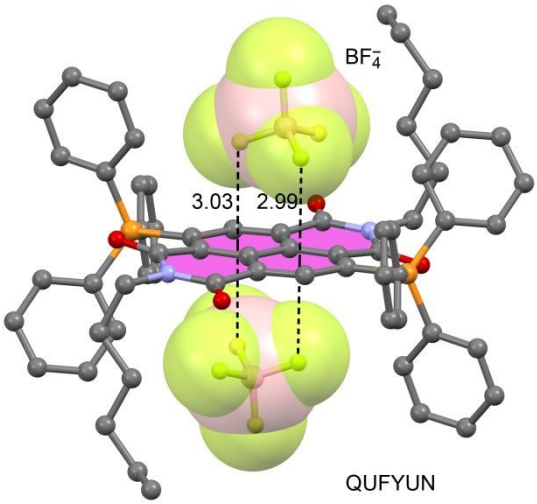


Fig. 8 Fragments of the X-ray crystal structures containing NDI π-acidic moieties. The relevant anion–π interactions are indicated by dashed lines. The CSD reference code is indicated. Distances in Å. H-atoms omitted for clarity.

A remarkable finding of their study is the extraordinary stability of both complexes, which were air-stable and resisted conventional silica column chromatography. What is more, the straightforward electron transfer (even in nonpolar solvent) strongly tunes the optical properties of the radical ion. This implies the usefulness of such materials in research towards switchable panchromatic materials, phosphorous-based stable radical ions and, in general, spin-based research.

NDIs have also been applied to encapsulate/stabilize polyoxoxmetalates (POMs) by anion-π interactions, as is shown in Fig. 9 where $[O_{40}PW_{14}]^{6-}$ is encapsulated by three N,N'-di(4-pyridyl)-1,4,5,8-naphthalenediimide units.[87] This enquiry demonstrated for the first time that anion–π interactions are appropriate for the stabilization and immobilization of functional POM anions.

CrystEngComm Accepted Manuscript

Please do not adjust margins

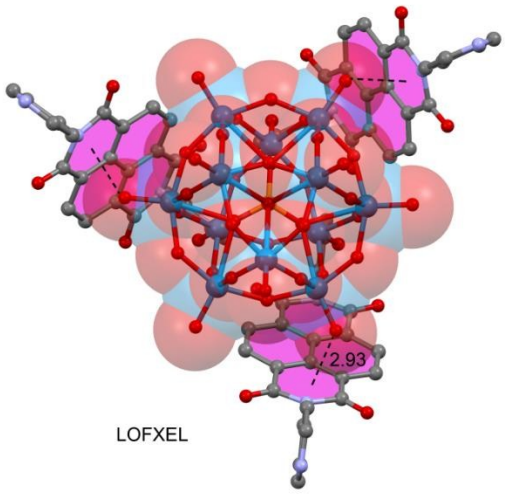


Fig. 9 Fragment of the X-ray crystal structures containing POM and NDI moieties. The relevant anion–π interactions are indicated by dashed lines. The CSD reference code is indicated. Distance in Å.

More recently, the combination of NDIs, POMs, Zn(II) and several co-ligands has been used[88] to synthesize a very rare radical-doped POM-based host–guest crystalline material. Interestingly, it has fast-responsive reversible photochromic properties and photocontrolled tunable luminescence. In addition, this material is able to photocatalytically oxidize benzylic alcohols to aldehydes using air as catalyst re-oxidant. Of particular importance is that the trapped POM anions interact with functional NDIs via directional anion–π contacts. The anion–π interactions stabilize and immobilize the functional POM anions and also promote the charge transfer and exchange among components leading to interesting properties of the crystalline material (i.e., reversible photochromism, photocontrolled tunable luminescence, and photocatalytic activity).

Stoddart et al.[89] reported on a macrocycle linking three NDIs together using 1,2-diaminocyclohexane, leading to molecular triangular prisms able to encapsulate anions (see Fig. 10). They have used both theory and experiment to demonstrate orbital interactions and electron transfer between the anion and the π-acidic surface of the NDIs in their synthesized redox-active prisms. The presence of three NDI moieties symmetrically distributed leads to a large number of individually accessible redox states in the triangular prisms, opening the door to potential applications in molecular electronics. The electron deficient interior of the molecular prism is perfect for studying anion–π interactions. This ability is evidenced by the trapping of linear I_3^- anions inside the receptor cavity, causing a significant change in the packing of the prisms in the extended solid-state architecture. The encapsulation of I_3^- anions provokes π–π stacking of the chiral prisms into supramolecular helices, providing an extraordinary example of anion-induced self-assembly. In addition, the chirality provided by the six stereogenic centers of the cyclohexane corners in the occupied prisms dictate the either right- or left-handedness of their packing in the solid state.

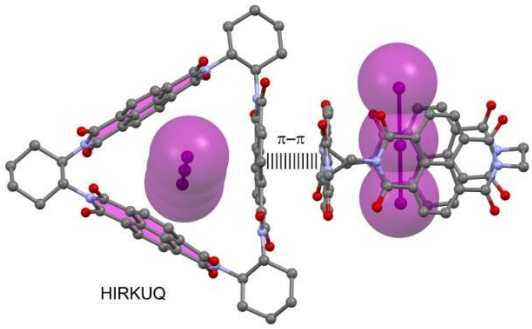


Fig. 10 The X-ray crystal structure of the NDI based triangular prism. The CSD reference code is indicated. H-atoms are omitted for clarity.

Another family of neutral anion receptors that exploited the additivity of anion-π interactions is based on tetraoxacalix[2]arene[2]triazine macrocyclic host reported by Wang *et al.*[90] They combined several techniques (electrospray ionization mass spectrometry, fluorescence titration and X-ray crystallography) to demonstrate that tetraoxacalix[2]arene[2]triazine forms 1:1 complexes with several polyatomic and monoatomic anions. The solid state structures with polyatomic anions like NO_3^-, BF_4^-, PF_6^- and SCN^- reveal that two opposing s-triazine rings act as a pair of tweezers to interact with the anions through cooperative anion–π interactions (see Fig. 11A). In monoatomic (smaller) anions like Cl^- and Br^-, a water molecule is also involved in the anion binding and the tweezers interact with the anion-water pair through anion–π and lp–π interactions (see Fig. 11B).[91] Interestingly, Wang et al. also altered the tetraoxacalix[2]arene[2]triazine host by introducing hydroxyl groups in the meta-position of the arene rings. In the NMe_4Br co-crystal of this novel macrocycle (see Fig. 11C),[92] the introduced –OH goups act like water in the unsubstituted version (Fig. 11B), hydrogen bonding to Br^- and interacting with an s-triazine ring by a lone-pair-π interaction.

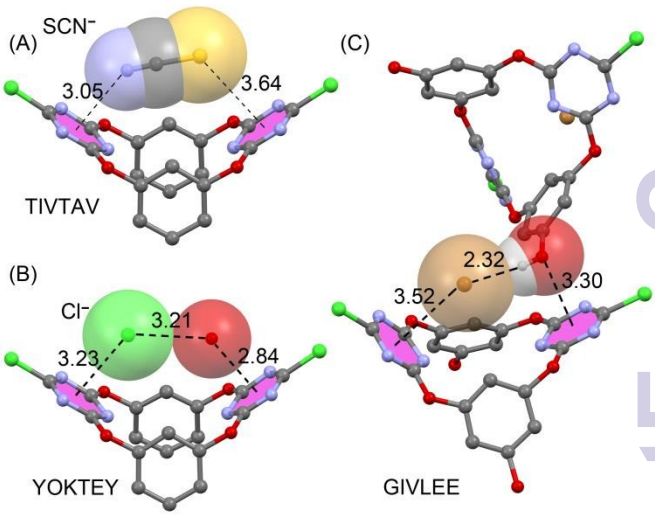


Fig. 11 X-ray crystal structures of tetraoxacalix[2]arene[2]triazine hosts. The CSD reference codes are indicated. Distances in Å. H-atoms omitted for clarity.

CrystEngComm Accepted Manuscript

This journal is © The Royal Society of Chemistry 20xx

Please do not adjust margins

1,4,5,8,9,12-hexaazatriphenylene (HAT) is a large (and thus easily polarizable) electron deficient aromatic known to generate anion-π interaction. For example, K. Dunbar *et al*. constructed a HAT derivative bearing six (electron withdrawing) cyanogroups.[93] This neutral yet strongly π-acidic molecule co-crystallized with N(*n*-Pr)$_4$Br from benzene, leading to a packing where HAT interacts with four anions simultaneously (three anions by one side and one by the opposite site, see Fig. 12). Furthermore, Ballester's group[94] has studied anion–π complexes of HAT(CN)$_6$ both theoretical and experimentally and demonstrated that the formation of anion–π interactions, charge-transfer or electron transfer adducts strongly depends on the type of anion.

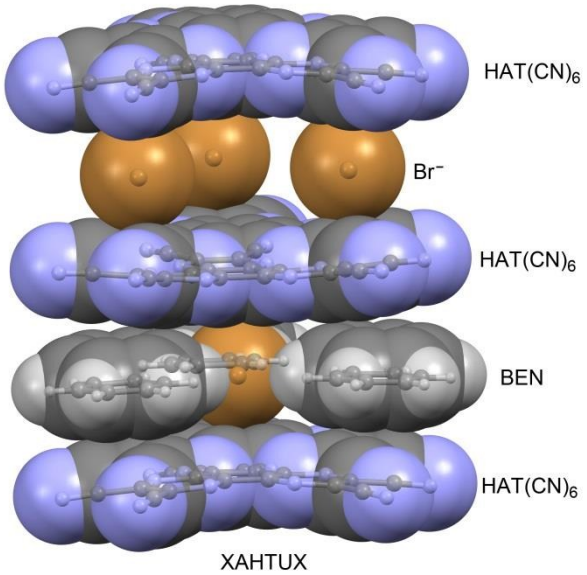


Fig. 12 X-ray structure of $[HAT(CN)_6]_2[Br^-]_3$. The CCDC reference code is indicated.

Finally, it is worth pointing out some theoretical studies with the bicycle pyridazino[4,5-d]pyridazine and larger polycyclic analogues thereof.[76,77] These inquiries demonstrated that the anion–π interaction strengthens as the number of fused rings increases. Interestingly, when such extended heteroaromatics were hydrogen bonded to water, even stronger complexes were predicted to form with anions.

3.3 Aromatics coordinated to metal ions

Dunbar's group has greatly contributed to highlight the crucial role of the anion–π interaction in the formation of self-assembled metallacycles.[95,96,97] Taking advantage of the structural versatility of metal ions and the directionality of metal–ligand interactions[98,99,100,101,102,103] they have recently described[104] the spontaneous assembly of elegant supramolecular architectures with unusual properties and applications or intriguing host–guest behaviour. They have provided unambiguous evidence that anion–π interactions are the main driving force in the templation process leading to the formation of Fe(II) metallacycles with π-acidic cavities, using 3,6-bis(2-pyridyl)-1,2,4,5-tetrazine as chelating ligand in the solid state. The anions play a decisive role in the formation and size of the supramolecular polygons, establishing multiple anion–π interactions with the electron deficient walls, as is exemplified in Fig. 13 for TIWHEO.

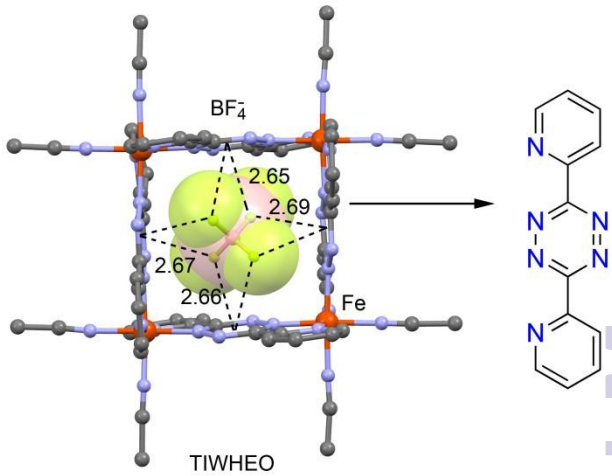


Fig. 13 Fragment of the X-ray crystal structures containing 3,6-bis(2-pyridyl)-1,2,4,5-tetrazine ligands. The relevant anion–π interactions are indicated by dashed lines. The CSD reference code is also indicated. H-atoms omitted for clarity.

Safin *et al*.[105] recently reported anion-induced self-assemblies using Ag^I salts and the two multidentate ligands 3,6-bis(2'-pyrimidyl)-1,2,4,5-tetrazine and 2,4,6-tris(2-pyrimidyl)-1,3,5-triazine. Interestingly, the selective use of PF_6^-, ClO_4^- and OTf^- anions leads to the formation of different solid state architectures, providing clear evidence of the pivotal role played by different anions (size and shape) and their interactions with the ligand. Two structures where anion-π directed assembly is most evident are depicted in Fig. 14A and B.

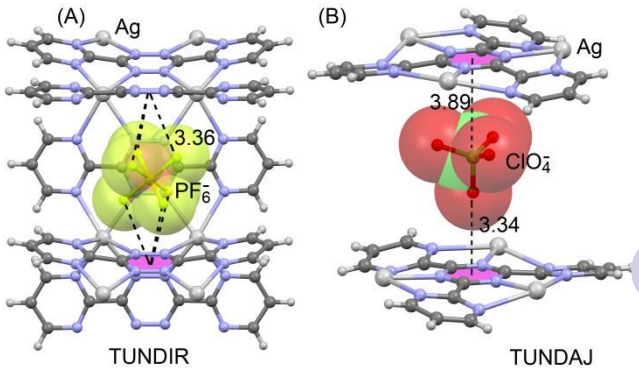


Fig. 14 Fragments of the X-ray crystal structures containing Pyrazine-N,N'-dioxide moieties. The relevant anion–π interactions are indicated by dashed lines. The CSD reference codes are indicated. Distances in Å.

In TUNDIR, the PF_6^- anion interacts with three central tetrazine rings of the 3,6-bis(2'-pyrimidyl)-1,2,4,5-tetrazine ligand, and in TUNDAJ the ClO_4^- is sandwiched between two central rings of the 2,4,6-tris(2-pyrimidyl)-1,3,5-triazine ligand.

Another clear example of this strategy has been reported by Domasevitch's group.[74] In particular, several X-ray crystal structures of *s*-tetrazine μ^4–coordinated to Ag^I have been

CrystEngComm Accepted Manuscript

Please do not adjust margins

reported, exhibiting very close contacts between anions and the *s*-tetrazine ring indicating strong anion–π interactions. Computational models of their structures corroborated that strong anion-π interactions are present between nitrate anions and the tetrazine ring. The polycyclic heteroaromatic arene pyridazino[4,5-d]pyridazine mentioned earlier has been studied as anion-π donor when coordinated to metal ions by Gural'skiy *et al.*,[74] clearly revealing the formation several anion–π interactions. Qin *et al.*[106] have reported the aqueous synthesis of an octanuclear macrocycle that includes fluorescent carbazole-based dipyrazole ligands, which coordinate to dipalladium corners. The X-ray diffraction analysis (see Fig. 15) reveals that this hybrid metallomacrocycle traps BF_4^- anions in the dipalladium-phenantroline clips through short anion–π contacts between BF_4^- anions and coordinated phenanthroline ligands.

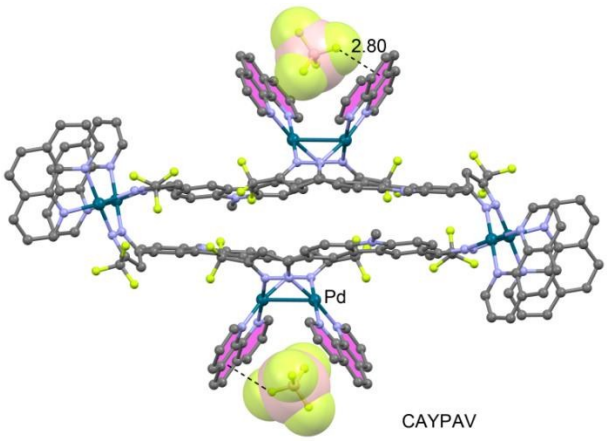


Fig. 15 Fragment of the X-ray crystal structure showing the anions trapped in the dipalladium clips through anion–π interactions. The relevant anion–π interactions are indicated by dashed lines. The CSD reference code is indicated. Distance in Å.

A recent experimental and theoretical study[107] has evidenced the formation of anion–π/π–π/π–anion assemblies in the solid state structures of several copper(II) complexes with a tetradentate Schiff base (see Fig. 16 for a selected example). Similar anion–π/π–π/π–anion assemblies in the solid state have been also found in N^6-decyladenine hydrochloride salts and bisadenine derivatives.[108]

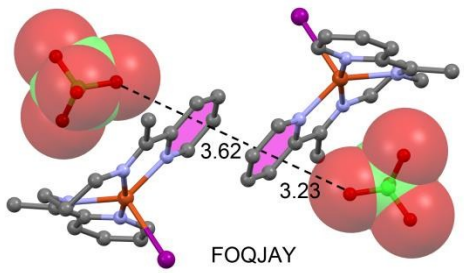


Fig. 16 Fragment of the X-ray crystal structure exhibiting anion–π/π–π/π–anion assembly. The CSD reference code is indicated. Distances in Å.

The interplay between π–π and anion–π interactions has also been investigated by others by combining theoretical studies and searches in the CSD.[109,110,111] These studies too revealed that the mutual influence between ion–π and π–π interactions can lead to strong cooperativity effects.

Atwood and co-workers reported on calixarenes and cyclotriveratrylenes macrocycles that have their arene rings μ^6–coordinated to transition metal ions (Ru, Ir, Rh).[69–72] This renders these molecules (which normally act as a host for cations) into an electron deficient bowl to accommodate anions, as is illustrated in Fig. 17.

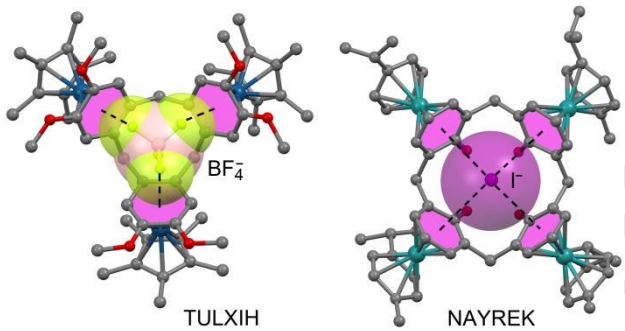


Fig. 17 Fragments of the X-ray crystal structures containing quinolizinium and tropylium moieties. The relevant anion–π interactions are indicated by dashed lines. The CSD reference codes are indicated.

In addition to the examples highlighted above, we next summarize some structures we found by our CSD inquiry[‡] where very strong anion-π contacts have been overlooked by the authors of the original manuscripts. Graphical illustrations of these examples are collected in Fig. 18A-D, and in all cases the van der Waals shell of an atom belonging to an anion overlaps with all six atoms of the interacting arene, representing η^6 anion-π interactions. In one example, UTEGII (see Fig. 18A), a PF_6^- anion interacts with phtalazine with an exceptionally short F···ring centroid distance of 2.64 Å; the sum of the van der Waals radii of F (1.47) and C (1.70) equals 3.17 Å. The shortest F···C distance of 2.908 Å is 0.262 Å shorter than the benchmark of overlapping van der Waals shells.

Another example is REJGEK as depicted in Fig. 18B, where the electrochemically active Rh(III) coordination complex exhibit a η^6 anion–π interaction between the phenyl aromatic ring and one chlorine atom of the anion. The Cl^-···ring centroid distance of 3.09 Å is again exceptionally short when compared to the sum of the van der Waals radii of 3.45 Å for Cl (1.75) and C (1.70). Kiplinger and co-workers recently investigated the coordination behaviour of bulky β-diketiminate ligands toward UCl_4.[112] The reaction of two equivalents of the ligand with UCl_4 led to an interesting cationic diuranium complex (see Fig 18C). The X-ray structure demonstrates that the β-diketiminate ligand is symmetrically bound to the U and an interesting combination of anion–π and halogen bonding interactions is crucial to rationalize the solid state architecture of this compound. That is, the Cl^- counterion is located almost perfectly over the ring centre of the aromatic ring of one complex and simultanously establishes several Cl···Cl interactions with the coordinated U chlorido ligands belonging to another complex generating a remarkably supramolecular assembly (see Fig 18C). Finally, Li *et al.*[113] reported on the self

CrystEngComm Accepted Manuscript

This journal is © The Royal Society of Chemistry 20xx

Please do not adjust margins

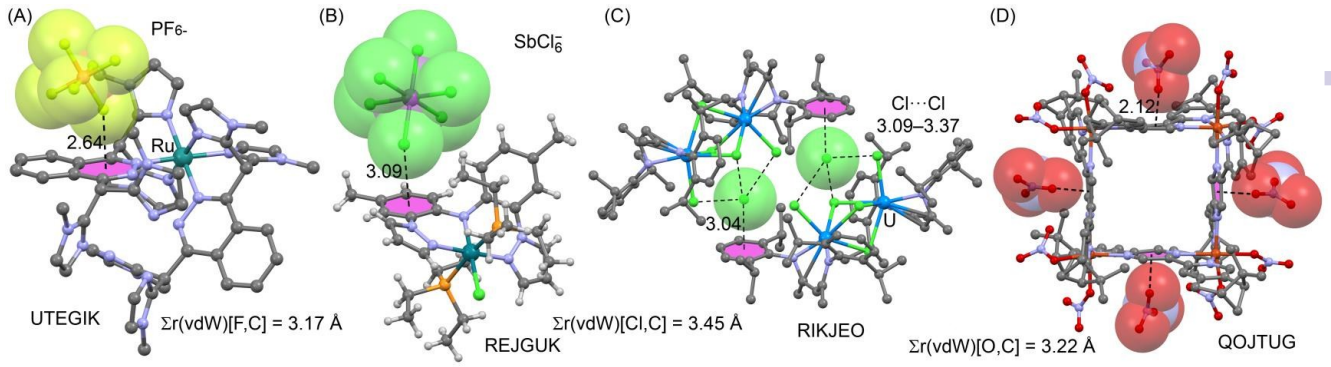


Fig. 18 (A-D) Illustrations of examples where a η^6 anion–π interaction involving a metal-coordinated heteroaromatic was found in the CSD,‡ which has escaped the attention of the authors of the original manuscripts. The CSD reference codes are given. Distances in Å. The sum of van der Waals radii [Σr(vdW)] values are also given. In A, B and D structures the H-atoms have been omitted for clarity.

assembly of enantiopure 2,5-bis(4,5-pinene-2-pyridyl)pyrazine ligands with copper(II) nitrate, leading to second-order nonlinear optically (NLO)-active square Cu(II) enantiomeric pairs (one enantiomer is shown in Fig 18D). The X-ray structure presents four crystallographically equivalent Cu(II) ions and four chiral ligands forming a molecular square. Each Cu(II) ion in these squares forms an octahedron by coordinating to four nitrogen atoms from the ligands and two oxygen atoms of two monodentate NO_3^- anions. Remarkably, each pyrazine ring forms extraordinary short anion–π interactions with the monodentate NO_3^- anion of the neighbouring complex (only the naked anions are shown in Fig 18D for clarity), thereby gluing together the individual squares. At this point, it is worth mentioning that in some of the examples shown above illustrating this section the anion gets as close to the cation as sterically possible. The π–system is basically in the way and it is difficult to separate the contribution of the cation and the π–system. To this respect, several manuscripts have theoretically studied this matter, generally using different theoretical models and the contribution of the anion–π interaction is usually large due to the enhanced acidity of the arene due to the coordination.[114] Moreover, since the anion–π interaction is directional and, conversely, the electrostatic attraction is non-directional, the final position of the (poly)anion is determined by the anion–π interaction.

3.4 Positively charged or protonated aromatics

The geometric and energetic features of anion–π^+ complexes of several aromatic cations (tropylium, quinolizinylium) have been investigated theoretically and by analysis of the CSD.[115,116,117,118] Two selected examples retrieved from the literature (KASVOP10[119] and CADZUC[120]) are shown in Fig. 19A and B.

The importance of the anion–π^+ interaction in protonated purine and pyrimidine bases has been recently reviewed.[121] As expected, interaction energies of anion–π^+ complexes are dominated by strong electrostatic effects exhibiting very large binding energies (> 80 kcal/mol).

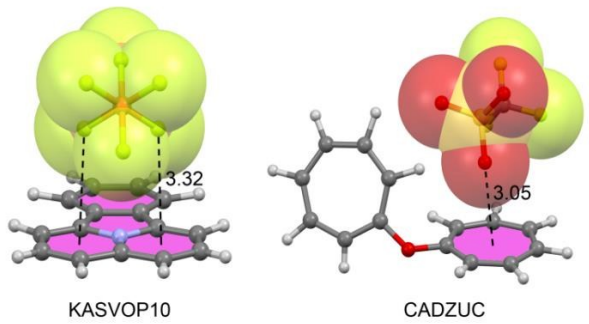


Fig. 19 Fragments of the X-ray crystal structures containing quinolizinium and tropylium moieties. The relevant anion–π interactions are indicated by dashed lines (distances in Å). The CSD reference codes are indicated.

Orvay *el al.*[122] also exploited anion–π^+ interactions in their design, synthesized and X-ray characterization of five proton transfer compounds using 4,6-di(1H-imidazol-1-yl)pyrimidine and different counterions. In the crystal structures of these five salts, anion–π interactions involving the aromatic rings play a fundamental role for the generation of three-dimensional supramolecular frameworks in the solid state. Two selected examples are illustrated in Fig. 20A,B.

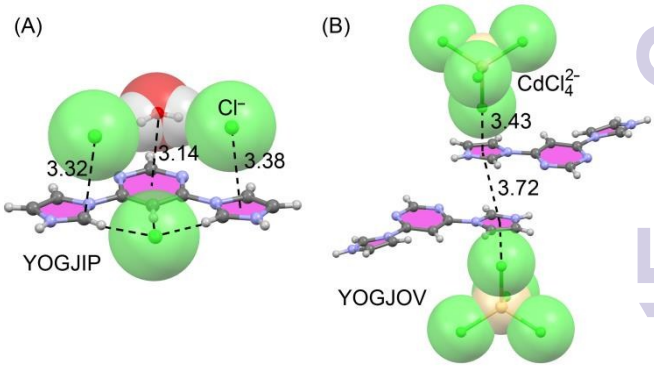


Fig. 20 Fragments of the X-ray crystal structures containing 4,6-di(1H-imidazol-1-yl)pyrimidine moieties. The relevant anion–π interactions are indicated by dashed lines. The CSD reference codes are indicated. Distances in Å.

CrystEngComm Accepted Manuscript

Please do not adjust margins

The first one (YOGJIP) is characterized by the formation of a remarkable supramolecular architecture through two anion–π^+ (chloride ions) and one lp–π (water molecule) interactions involving all three aromatic rings of the diprotonated 4,6-di(1H-imidazol-1-yl)pyrimidine moiety. That is, an oxygen atom lone pair of the water molecule is oriented toward the π-cloud of the pyrimidine ring forming a strong lp–π interaction. In addition, two chloride ions interact with the imidazolium rings forming two anion–π^+ interactions. In the second one (YOGJOV) two $[CdCl_4]^{2-}$ anions and two imidazolium moieties participate in the generation of anion–π/π–π/π–anion assemblies (see Fig. 20B).

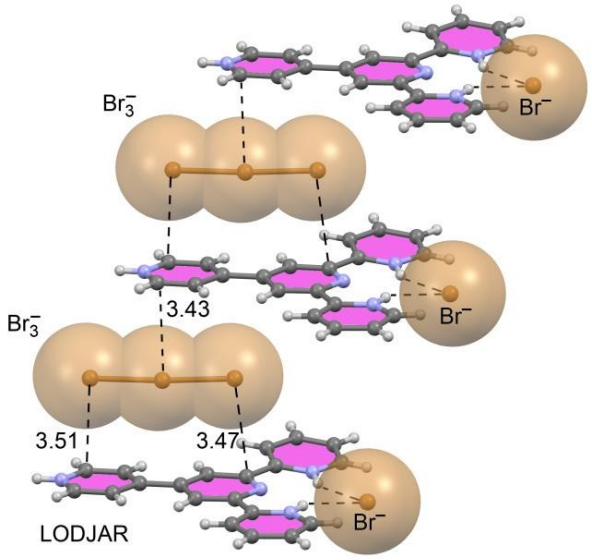


Fig. 21 Fragment of the X-ray crystal structure containing 4′-(4-pyridyl)-2,2′:6′,2″-terpyridine moieties. The relevant anion–π interactions are indicated by dashed lines. The CSD reference code is also indicated. Distances in Å.

Several interesting crystal structures of 4′-(4-pyridyl)-2,2′:6′,2″-terpyridine (PTP) salts have been recently reported by Manna *et al.*[123] They found that the structural architecture depends on the type of anion (mononuclear or trinuclear) and on the pH from which the solid crystallized. As is illustrated in Fig 21 for one of the structures (LODJAR), the $[\text{pyridine-H}]^+$ rings concurrently participate in hydrogen-bonding (N–H···Br^-) and anion–π^+ interactions.

In this experimental work, the role of the anion–π^+ interaction in solid state has also been investigated theoretically, providing additional evidence to its role as a decisive supramolecular force responsible for the packing geometry of these salts. The same research group[124] has further investigated the role of the anion–π^+ interactions in the generation of supramolecular assemblies using a similar ligand, i.e. the triply protonated 4′-(4-pyridyl)-3,2′:6′,3″-terpyridine. Combining X-ray crystallography with theoretical studies, they showed that π^+–π and various anion–π^+ interactions are the major driving forces in the stabilization of the assemblies observed in these solid state structure (see Fig. 22).

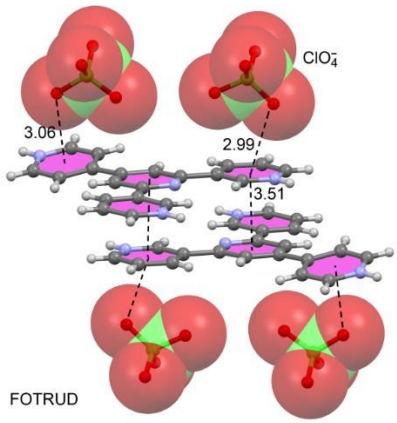


Fig. 22 Fragment of the X-ray crystal structure containing 4′-(4-pyridyl)-3,2′:6′,3″-terpyridine moieties. The relevant anion–π interactions are indicated by dashed lines. The CSD reference code is also indicated. Distances in Å.

Shown in Fig. 23A-C are illustrations of solid state structures where anion–π^+ interactions (found by our CSD analysis)[‡] are particularly relevant (η^6 anion-π^+ interactions) but not described by the original authors. For instance Chen *et al.*[125] have reported the X-ray structure of the 5,6-Dioxo-1,10-phenanthrolinium bromide (see Fig. 23A). The bromide anions

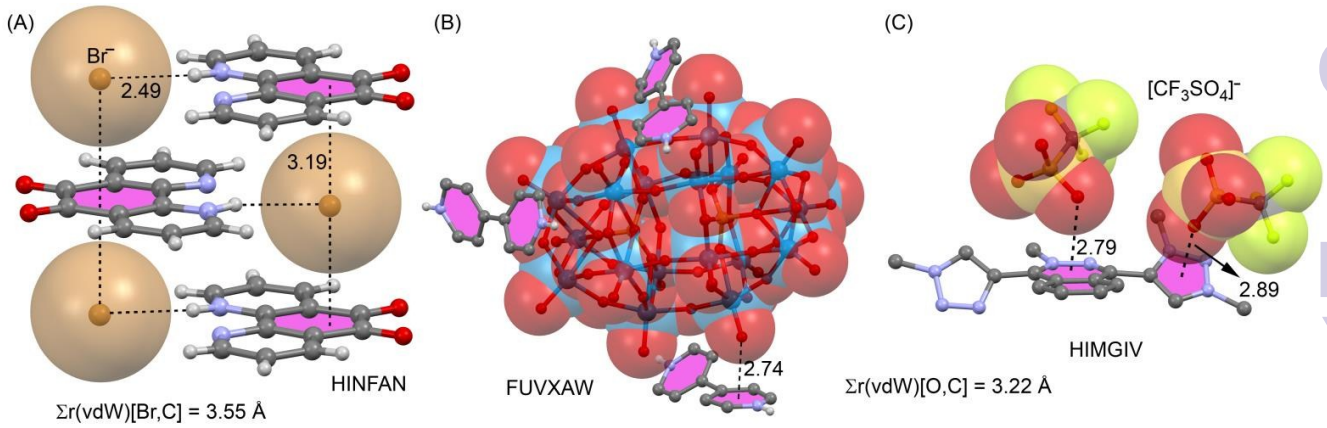


Fig. 23 Illustrations of examples where a η^6 anion–π interaction involving a formally cationic aromatic was found in the CSD,[‡] which has escaped the attention of the authors of the original manuscripts. Distances in Å. The sum of van der Waals radii [Σr(vdW)] values are also given

CrystEngComm Accepted Manuscript

This journal is © The Royal Society of Chemistry 20xx

Please do not adjust margins

induce the formation of sandwich complexes in the solid state, forming two symmetrically equivalent anion–π^+ interactions with the dioxo ring. The shortest Br^-···ring centroid distance of 3.22 Å is 0.33 Å shorter than the sum of the van der Waals radii of B (1.85) and C (1.70) equals 3.55 Å. The shortest Br^-···C distance of 3.36 Å is 0.19 Å shorter than the benchmark of overlapping van der Waals shells. 4,4'-bipyridine has been combined with polyoxometalates POMs to generate several organic-inorganic supramolecular hybrid crystal, prepared from acidic solutions, that are organized through anion-π^+ interactions.[126] In Fig. 23B we show a selected example where a Wells–Dawson polyoxoanion is involved. This type of POM has 54 surface oxygen atoms including 18 terminal oxygen atoms (only bonded to W atom) and 36 μ^2-O atoms (bonded to two W atom). The terminal O atoms establish extensive anion–π^+ interactions in the crystal structure (see Fig 23B). These interactions along with hydrogen bonds connect the POMs and the diprotonated bipyridine cations generating a remarkable 3D supramolecular network. Finally, Albretch's and co-workers[127] have synthesised and characterized of a novel phthalazine-triazole ligand (1,4-bis(1-methyl-1H-1,2,3-triazol-4-yl)phthalazine) with the purpose to generate new ruthenium(II) complexes. They also studied its reactivity in several methylation reactions. Particularly, when the ligand was reacted with MeOTf (OTf = trifluoromethyl sulfonate) a double methylation was observed and the resulting product was characterized by X-ray analysis (see Fig. 23C). Interestingly, the double methylated ligand exhibit anion–π^+ interactions with both the six and five membered rings that are positively charged in the solid state

Concluding remarks

There clearly remains an element of unpredictability in crystal engineering, as even the coordination geometry of a transition metal complex is not always as predicted. The degree of serendipity logically increases with weaker interactions, which include the anion-π interaction. Nevertheless, anion-π interactions are undoubtedly a force to be reckoned with for a proper post factum rationalization of supramolecular assemblies in crystals. Moreover, while serendipity remains, our recently acquired understanding of anion-π interactions – as highlighted in this work– has led to a degree of predictability that was absent only about two decades ago. It is our hope that the design strategies outlined in this highlight will inspire (supra)molecular designers to advance this predictability even more. We furthermore anticipate that with an increased control over anion-π interactions follow opportunities for the design of novel materials, catalysts, sensors, and biologically active molecules.

Acknowledgements

The financial support from MINECO of Spain (projects CTQ2014-57393-C2-1-P and CONSOLIDER INGENIO 2010 CSD2010-00065, FEDER funds) is gratefully acknowledged.

Notes and references

‡ The CSD version 5.36 (November 2014, including three updates) was consulted using ConQuest version 1.17, while excluding entries before November 2010 (CSD version 5.32). A subset was first created where an aromatic six-membered ring consisting of C and/or N was present. This subset was then subjected to a search for structures where the *inter*molecular distance between an electron rich atom (N, P, As, O, S, Se, TE, F, Cl, Br, I, At) and all six of the rings atoms have overlapping van der Waals shells. Manual inspection of the resulting 68 structures revealed several clear examples of anion-π interactions that were not mentioned by the authors of these structures, as highlighted in this contribution. Several of the structures were too disordered for reliable interpretation, some contained clear evidence of lone-pair–π interactions, and some structures contained anio-π features that the authors of the structure intended and/or mentioned. Relaxing the search criteria to 6 × overlapping van der Waals shells + 0.1 Å gave 170 structures and relaxing further to 0.25 Å gave 1,057 structures that might well contain numerous additional examples of anion-π interactions.

1 H. J. Schneider, *Angew. Chem. Int. Ed*., 2009, **48**, 3924.
2 H. J. Schneider and A. Yatsimirski, Principles and methods in supramolecular chemistry, Wiley, Chichester, 2000.
3 J. M. Lehn, Supramolecular chemistry concepts and perspectives, Wiley–VCH, Weinheim, 1995.
4 F. Vögtle Supramolecular chemistry: an introduction, Wiley, New York, 1993.
5 a) P. D. Beer, P. A. Gale and D. K. Smith, Supramolecular chemistry, Oxford University Press, Oxford, 1999; b) J. W. Steed and J. L. Atwood, Supramolecular chemistry, Wiley, Chichester, 2000.
6 H. Destecroix, C. M. Renney, T. J. Mooibroek, T. S. Carter, P. F. N. Stewart, M. P. Crump and A. P. Davis, *Angew. Chem. Int. Ed*., 2015, **54**, 2057.
7 P. Murray-Rust and W. D. S. Motherwell, *J. Am. Chem. Soc*., 1979, **101**, 4374.
8 P. Murray-Rust, W. C. Stallings, C. T. Monti, R. K. Preston and J. P. Glusker, *J. Am. Chem. Soc.*, 1983, **105**, 3206.
9 N. Ramasubbu, R. Parthasarathy and P. Murray-Rust, *J. Am. Chem. Soc*., 1986, **108**, 4308.
10 P. Metrangolo, H. Neukirch, T. Pilati and G. Resnati, *Acc. Chem. Res*., 2005, **38**, 386.
11 P. Politzer and J. S. Murray, *ChemPhysChem*, 2013, **14**, 278.
12 P. Politzer, J. S. Murray and T. Clark, *Phys. Chem. Chem. Phys*., 2013, **15**, 11178.
13 P. Politzer, J. S. Murray and T. Clark, *Phys. Chem. Chem. Phys*., 2010, **12**, 7748.
14 A. Bundhun, P. Ramasami, J. S. Murray and P. Politzer, *J. Mol. Model*., 2013, **19**, 2739.
15 a) A. Bauzá, T. J. Mooibroek and A. Frontera, *Angew. Chem. Int. Ed*., 2013, **52**, 12317; b) A. Bauzá, T. J. Mooibroek and A. Frontera, *ChemPhysChem*., 2015, **16**, 2496; c) A. Bauzá and A. Frontera, *ChemPhysChem*., 2015, **16**, 3108.
16 J. C. Ma and D. A. Dougherty, *Chem. Rev*., 1997, **97**, 1303.
17 N. Zacharias and D. A. Dougherty, *Trends Pharmacol. Sci*., 2002, **23**, 281.
18 N. S. Scrutton and A. R. C. Raine, *Biochem J*., 1996, **319**, 1.
19 J. P. Gallivan and D. A. Dougherty, *Proc. Nat. Acad. Sci*., 1999, **96**, 9459.
20 A. Frontera, P. Gamez, M. Mascal, T. J. Mooibroek and J. Reedijk, *Angew. Chem. Int. Ed*., 2011, **50**, 9564.
21 B. L. Schottel, H. T. Chifotides and K. R. Dunbar, *Chem. Soc. Rev*., 2008, **37**, 68.
22 C. Caltagirone and P. A. Gale, *Chem. Soc. Rev*., 2009, **38**, 520.

CrystEngComm Accepted Manuscript

23 P. Gamez, T. J. Mooibroek, S. J. Teat and J. Reedijk, *Acc. Chem. Res*., 2007, **40**, 435.
24 B. P. Hay and V. S. Bryantsev, *Chem. Commun*., 2008, **44**, 2417.
25 A. Bauzá, D. Quiñonero, P. M. Deyà and A. Frontera *New J. Chem*., 2013, **37**, 2636.
26 J. R. Galán–Mascarós, A. M. Z. Slawin, J. D. Woollins and D. J. Williams, *Polyhedron* 1996, **15**, 4603.
27 H. J. Schneider, F. Werner and T. Blatter, *J. Phys. Org. Chem*., 1993, **6**, 590.
28 H. J. Schneider, *Angew. Chem. Int. Ed. Engl*., 1991, **30**, 1417.
29 K. Hiraoka, S. Mizuse and S. Yamabe, *J. Phys. Chem*., 1987, **91**, 5294.
30 K. Hiraoka, S. Mizuse and S. Yamabe, *J. Chem. Phys.*, 1987, **86**, 4102.
31 J. S. Murray, P. Lane, T. Clark, K. E. Riley and P. Politzer, *J. Mol. Model*., 2012, **18**, 541.
32 D. Quiñonero, C. Garau, C. Rotger, A. Frontera, P. Ballester, A. Costa and P. M. Deyà, *Angew. Chem. Int. Ed*., 2002, **41**, 3389.
33 M. Mascal, A. Armstrong and M. D. Bartberger, *J. Am. Chem. Soc*., 2002, **124**, 6274.
34 I. Alkorta, I. Rozas and J. Elguero, *J. Am. Chem. Soc*., 2002, **124**, 8593.
35 D. Y. Kim, N. J. Singh and K. S. Kim, *J. Chem. Theory Comput*., 2008, **4**, 1401.
36 B. L. Schottel, J. Bacsa and K. R. Dunbar, *Chem. Commun*., 2005, **41**, 46.
37 B. Han, J. J. Lu and J. K. Kochi, *Cryst. Growth Des*., 2008, **8**, 1327.
38 M. Mascal, I. Yakovlev, E. B. Nikitin and J. C. Fettinger, *Angew. Chem. Int. Ed*., 2007, **46**, 8782.
39 R. J. Götz, A. Robertazzi, I. Mutikainen, U. Turpeinen, P. Gamez and J. Reedijk, *Chem. Commun*., 2008, **44**, 3384.
40 M. Albrecht, M. Müller, O. Mergel, K. Rissanen and A. Valkonen, *Chem. Eur. J*., 2010, **16**, 5062.
41 C. Estarellas, A. Frontera, D. Quiñonero and P. M. Deyà, *Angew. Chem. Int. Ed*., 2011, **50**, 415.
42 S. Chakravarty, Z. Z. Sheng, B. Iverson and B. Moore, *FEBS Lett*., 2012, **586**, 4180.
43 D. D. Jenkins, J. B. Harris, E. F. Howell, R. J. Hinde and J. Baudry, *J Comput. Chem*., 2013, **34**, 518.
44 a) R. E. Dawson, A. Hennig, D. P. Weimann, D. Emery, V. Ravikumar, J. Montenegro, T. Takeuchi, S. Gabutti, M. Mayor, J. Mareda, C. A. Schalley and S. Matile, *Nat. Chem*., 2010, **2**, 533; b) N. Sakai, J. Mareda, E. Vauthey and S. Matile, *Chem. Commun*., 2010, **46**, 4225.
45 K. E. Ranaghan, J. E. Hung, G. J. Bartlett, T. J. Mooibroek, J. N. Harvey, D. N. Woolfson, W. A. van der Donk and A. J. Mulholland, *Chem. Sci*., 2014, **5**, 2191.
46 P. Gamez, T. J. Mooibroek, S. J. Teat and J. Reedijk, *Acc. Chem. Res*., 2007, **40**, 435.
47 P. Ballester, Recognition of Anions, Springer Berlin, Heidelberg, 2008
48 A. Frontera, *Coord. Chem. Rev*., 2013, 257, 1716.
49 M. Giese, M. Albrecht and Kari Rissanen, *Chem. Rev*., 2015, **115**, 8867.
50 D. Kim, P. Tarakeshwar and K. S. Kim, *J. Phys. Chem. A*, 2004, **108**, 1250.
51 D. Y. Kim, N. J. Singh, J. W. Lee and K. S. Kim, *J. Chem. Theory Comput.*, 2008, **4**, 1162.
52 C. Garau, A. Frontera, D. Quiñonero, P. Ballester, A. Costa and P. M. Deyà, *Chem. Phys. Chem*. 2003, **4**, 1344.
53 D. Quiñonero, C. Garau, A. Frontera, P. Ballester, A. Costa and P. M. Deyà, *Chem. Phys. Lett*., 2002, **359**, 486.
54 A. Bauzá, P. M. Deyà, A. Frontera and D. Quiñonero, *Phys. Chem. Chem. Phys*., 2014 **16**, 1322.
55 C. Garau, D. Quiñonero, A. Frontera, P, Ballester, A. Costa, P. M. Deyà, *New J. Chem*., 2003, **27**, 211.
56 C. Garau, D. Quiñonero, A. Frontera, P. Ballester, A. Costa and P. M. Deyà, *Org. Lett*., 2003, **5**, 2227.
57 C. Garau, A. Frontera, D. Quiñonero, P. Ballester, A. Costa and P. M. Deyà, *J. Phys. Chem. A*, 2004, **108**, 9423.
58 C. Garau, A. Frontera, D. Quiñonero, P. Ballester, A. Costa and P. M. Deyà, *J. Phys. Chem. A*, 2004, **108**, 9423.
59 C. Garau, A. Frontera, D. Quiñonero, P. Ballester, A. Costa and P. M. Deyà, *Chem. Phys. Lett*., 2004, **399**, 220.
60 T. Steiner, *Angew. Chem. Int. Ed*., 2002, **41**, 48.
61 T. J. Mooibroek, P. Gamez, *Inorg. Chim. Acta* 2007, **360**, 381.
62 T. J. Mooibroek, C. A. Black, P. Gamez and J. Reedijk, *Cryst. Growth Des*., 2008, 8, 4, 1082.
63 a) C. Murcia-García, A. Bauzá, A. Frontera and R. Streubel, *CrystEngComm*, 2015, **17**, 6736; b) C. Murcia-García, A. Bauzá, G. Schnakenburg, A. Frontera and R. Streubel, *CrystEngComm*, 2015, **17**, 1769.
64 a) T. J. Mooibroek and P. Gamez, *CrystEngComm* 2012, **14**, 1027; b) T. J. Mooibroek and P. Gamez, *CrystEngComm*, 2013, **15**, 1802.
65 J. Kroon, and J. A. Kanters, Nature, 1975, **248**, 667.
66 I. Alkorta and J. Elguero, *J. Phys. Chem. A* 2003, **107**, 9428.
67 I. Alkorta, D. Quiñonero, C. Garau, A. Frontera, J. Elguero and P. M. Deyà, *J. Phys. Chem. A* 2007, **111**, 3137.
68 D. Quiñonero, A. Frontera, P. M. Deyà, I. Alkorta and J. Elguero, *Chem. Phys. Lett*., 2008, **460**, 406.
69 J. W. Steed, R. K. Juneja and J. L. Atwood, *Angew. Chem. Int. Ed*., 1995, **33**, 2456.
70 M. Staffilani, K. S. B. Hancock, J. W. Steed, K. T. Holman, J. L. Atwood, R. K. Juneja and R. S. Burkhalter, *J. Am. Chem. Soc*., 1997, **119**, 6324.
71 K. T. Holman, M. M. Halihan, S. S. Jurisson, J. L. Atwood, R. S. Burkhalter, A. R. Mitchell and J. W. Steed, *J. Am. Chem. Soc*., 1996, **118**, 9567.
72 M. Staffilani, G. Bonvicini, J. W. Steed, K. T. Holman, J. L. Atwood and M. R. J. Elsegood, *Organometallics* 1998, **17**, 1732.
73 D. Quiñonero, A. Frontera and P. M. Deyà, *ChemPhysChem*., 2008, **9**, 397.
74 I. A. Gural'skiy, D. Escudero, A. Frontera, P. V. Solntsev, E. B. Rusanov, A. N. Chernega, H. Krautscheid and K. V. Domasevitch, *Dalton Trans*., 2009, **38**, 2856.
75 A. Bauzá, D. Quiñonero, P. M. Deyà and A. Frontera, *Theor. Chem. Acc.*, 2012, **131**, 1219.
76 D. Escudero, A. Frontera, D. Quiñonero and P. M. Deyà, *J. Comput. Chem*., 2009, **30**, 75.
77 X. Lucas, C. Estarellas, D. Escudero, A. Frontera, D. Quiñonero and P.M. Deyà, *ChemPhysChem*, 2009, **10**, 2256.
78 C. Garau, D. Quiñonero, A. Frontera, P. Ballester, A. Costa and P. M. Deyà, *J. Phys. Chem. A*, 2005, **109**, 9341.
79 R. Podgajny, D. Pinkowicz, B. Czarnecki, M. Kozieł, S. Chorąży, B. Wis, W. Nitek, M. Rams and B. Sieklucka, *Cryst. Growth Des*., 2014, **14**, 4030.
80 M. Giese, M. Albrecht, A. Valkonen and K. Rissanen, *Eur. J. Org. Chem*., 2013, 3247.
81 M. Giese, M. Albrecht, T. Repenko, J. Sackmann, A. Valkonen and K. Rissanen, *Eur. J. Org. Chem*., 2014, 2435

CrystEngComm Accepted Manuscript

This journal is © The Royal Society of Chemistry 20xx

82 M. Giese, M. Albrecht, A. Valkonen and K. Rissanen, *Chem. Sci*., 2015, **6**, 354.
83 J.-J. Liu, Y.-F. Guan, C. Jiao, M.-J. Lin, C.-C. Huang and W.-X. Dai, *Dalton Trans*., 2015, **44**, 5957-5960.
84 J. Fujisawa, N. Tajima, K. Tamaki, M. Shimomura and T. Ishihara, *J. Phys. Chem. C* 2007, **111**, 1146.
85 T. Fukumoto, M. Hirasawa and T. Ishihara, *J. Lumin*., 2000, **497**, 87.
86 S. Kumar, M. R. Ajayakumar, G. Hundal and P. Mukhopadhyay, *J. Am. Chem. Soc*., 2014, **136**, 12004.
87 J.-Z. Liao, X.-J. Dui, H.-L. Zhang, X.-Y. Wu, C.-Z. Lu *CrystEngComm*, 2014, **16**, 10530.
88 J.-Z. Liao, H.-L. Zhang, S.-S. Wang, J.-P. Yong, X.-Y. Wu, R. Yu, and C.-Z. Lu, *Inorg. Chem*., 2015, **54**, 4345–4350
89 S. T. Schneebeli, M. Frasconi, Z. Liu, Y. Wu, D. M. Gardner, N. L. Strutt, C. Cheng, R. Carmieli, M. R. Wasielewski and J. R. Stoddart, *Angew. Chem. Int. Ed*., 2013, **52**, 13100.
90 D.-X. Wang and M.-X. Wang, *J. Am. Chem. Soc*., 2013, **135**, 892.
91 D.-X. Wang, Q.-Y. Zheng, Q.-Q. Wang and M.-X. Wang, *Angew. Chem. Int. Ed*., 2008, **47**, 7485.
92 W. Liu, Q.-Q. Wang, Y. Wang, Z.-T. Huang, D.-X. Wang, *RSC Adv*., 2014, **4**, 9339.
93 H. T. Chifotides, B. L. Schottel and K. R. Dunbar, *Angew. Chem. Int. Ed*., 2010, **49**, 7202.
94 G. Aragay, A. Frontera, V. Lloveras, J. Vidal-Gancedo and P. Ballester, *J. Am. Chem. Soc*., 2013, **135**, 2620.
95 C. S.Campos-Fernandez, B. L. Schottel, H. T. Chifotides, J. K. Bera, J. Bacsa, J. M. Koomen, D. H. Russell and K. R. Dunbar, *J. Am. Chem. Soc.*, 2005, **127**, 12909.
96 B. L. Schottel, H. T. Chifotides, M. Shatruk,A. Chouai, J. Bacsa, L. M. Perez and K. R. Dunbar, *J. Am. Chem. Soc.*, 2006, **128**, 5895.
97 P. S. Szalay, J. R. Galan-Mascaros, B. L. Schottel, J. Basca, L. M. Perez, I. S. Ichimura, A. Chouai and K. R. Dunbar, *J. Cluster Sci*., 2004, **15**, 503.
98 R. Chakrabarty, P. S. Mukherjee and P. J. Stang, *Chem. Rev*., 2011, **111**, 6810.
99 M. Fujita, M. Tominaga, A. Hori and B. Therrien, *Acc. Chem. Res*., 2005, **38**, 371.
100 M. Fujita, K. Umemoto, M. Yoshizawa, N. Fujita, T. Kusukawa and K. Biradha, *Chem. Commun*., 2001, 509.
101 M. D. Pluth, R. G. Bergman and K. N. Raymond, *Acc. Chem. Res*., 2009, **42**, 1650.
102 R. A. Bilbeisi, J. K. Clegg, N. Elgrishi, X. de Hatten, M. Devillard, B Breiner, P. Mal and J. R. Nitschke, *J. Am. Chem. Soc*., 2012, **134**, 5110.
103 J. Rebek, Jr. *Acc. Chem. Res*., 2009, **42**, 1660.
104 H. T. Chifotides, I. D. Giles and K. R. Dunbar, *J. Am. Chem. Soc*., 2013, **135**, 3039.
105 D. A. Safin, A. Pialat, A. A. Leitch, N. A. Tumanov, I. Korobkov, Y. Filinchuk, J. L. Brusso and M. Murugesu, *Chem. Commun*., 2015, **51**, 9547-9550
106 L. Qin, L.-Y. Yao and S.-Y. Yu, Inorg. Chem., 2012, **51**, 2443–2453
107 A. Bhattacharyya, P. K. Bhaumik, A. Bauzá, P. P. Jana, A. Frontera, M. G. B. Drew and S. Chattopadhyay, *RSC Adv*., 2014, **4**, 58643.
108 A. Garcia-Raso, F. M. Albertí, J. J. Fiol, A. Tasada, M. Barceló-Oliver, E. Molins, D. Escudero, A. Frontera, D. Quiñonero, and P. M. Deyà, *Inorg. Chem*., 2007, *46*, 10724.
109 A. Frontera, D. Quiñonero, C. Garau, A. Costa, P. Ballester and P. M. Deyà, *J. Phys. Chem. A*, 2006, **110**, 9307.
110 D. Quiñonero, A. Frontera, C. Garau, P. Ballester, A. Costa and P. M. Deyà, *ChemPhysChem*, 2006, **7**, 2487.
111 A. Frontera, D. Quiñonero, A. Costa, P. Ballester and P. M. Deyà, *New J. Chem*., 2007, **31**, 556.
112 M. J. Monreal, R. J. Wright, D. E. Morris, B. L. Scott, J. T. Golden, P. P. Power and J. L. Kiplinger, *Organometallics*, 2013, **32**, 1423.
113 X.-L. Li, J.-L. Kang, X.-L. Zhang, H.-P. Xiao, A.-L. Wang, L. Zhou, S.-M. Fanga and C.-M. Liu, *Dalton Trans*., 2014, **43**, 17226.
114 a) S. Brooker, N. G. White, A. Bauza, P. M. Deyà and A. Frontera, Inorg. Chem., 2012, 51, 10334; b) C. Biswas, M. G. B. Drew, D. Escudero, A. Frontera and A. Ghosh, *Eur. J. Inorg. Chem*., 2009, **2009**, 2238-2246; c) A. Das, S. M. Choudhury, C. Estarellas, B. Dey, A. Frontera, J. Hemming, M. Helliwell, P. Gamez and S. Mukhopadhyay, *CrystEngComm*, 2011, **13**, 4519; d) P. Kar, R. Biswas, M. G. R. Drew, A. Frontera and A. Ghosh, *Inorg. Chem*. 2012, **51**, 1837.
115 C. Estarellas, A. Frontera, D. Quiñonero and P.M. Deyà, *J. Chem. Theory Comput.*, 2008, **4**, 1981.
116 D. Quiñonero, A. Frontera, D. Escudero, P. Ballester, A. Costa and P. M. Deyà, *ChemPhysChem*., 2007, **8**, 1182.
117 A. Das, S. Ray Choudhury, B. Dey, S. K. Yalamanchili, M. Helliwell, P. Gamez, S. Mukhopadhyay, C. Estarellas and A. Frontera, *J. Phys. Chem. B* 2010, **114**, 4998.
118 A. Das, S. Ray Choudhury, C. Estarellas, B. Dey, A. Frontera, J. Hemming, M. Helliwell, P. Gamez and S. Mukhopadhyay, *CrystEngComm*, 2011, **13**, 4519.
119 M. Fourmigue, K. Boubekeur, P. Batail, K. Bechgaard, *Angew. Chem., Int. Ed*., 1989, **28**, 588
120 R. F. Childs, R. Faggiani, C. J. L. Lock, C. V. Rogerson, *J. Org. Chem*., 1983, **48**, 3043
121 J. J. Fiol, M. Barceló-Oliver, A. Tasada, A. Frontera, À. Terrón and Á. García-Raso, *Coord. Chem. Rev*., 2013, **257**, 2705.
122 F. Orvay, A. Bauzá, M. Barceló-Oliver, A. García-Raso, J. J. Fiol, A. Costa, E. Molins, I. Mata and A. Frontera, *CrystEngComm*, 2014, **16**, 9043.
123 P. Manna, S. K. Seth, A. Bauzá, M. Mitra, S. R. Choudhury, A. Frontera and S. Mukhopadhyay, *Cryst. Growth Des*., 2014, **14**, 747.
124 P. Manna, S. K. Seth, M. Mitra, S. R. Choudhury, A. Bauzá, A. Frontera and S. Mukhopadhyay, *Cryst. Growth Des*., 2014, **14**, 5812.
125 M.-H. Chen, S.-F. Xue and Z. Tao, *Hecheng Huaxue*, 2009, **17**, 722
126 (a) J.-W. Zhao, L.-J. Chen, J.-P. Wang, *Huaxue Yanjiu*, 2008, **19**, 14; (b) H. Yang, X. Lin, B. Xu, Y. You, M. Cao, S. Gao and R. Cao, *J. Mol. Struct*., 2010, **966**, 33
127 J. Aguiló, A. Naeimi, R. Bofill, H. Mueller-Bunz, A.Llobet, L. Escriche, X. Sala and M. Albrecht, *New J. Chem*., 2014, **38**, 1980

CrystEngComm Accepted Manuscript