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The supramolecular assemblies based on heteroatom-containing triangulenes

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The heteroatom-containing derivatives of triangulene feature a rich supramolecular chemistry which results from their (mostly) planar π -conjugated structure – predestined for the self-assembly via $\pi\pi$ -stacking interactions into 1D or 2D architectures. Herein, this behavior will be discussed – also taking the potential applications of these assemblies into account.

Introduction

Parent triangulene (**1**), also referred to as Clar's hydrocarbon, represents the structurally simplest non-Kekulé-type polycyclic aromatic hydrocarbon (PAH) featuring a biradical triplet ground state.¹ Though already hypothesized by Clar in 1953,² the on-surface synthesis and structural characterization of the highly reactive compound **1** was accomplished by Pavlíček *et al.*

very recently (Fig. 1);³ a number of substituted, thus stabilized derivatives was reported already some years earlier.⁴ Open-shell non-Kekulé PAHs, such as **1**, are essentially of academic interest, though applications in the field of organic magnetics⁵ and optoelectronics¹ have been suggested. The magnetic and photo-physical properties are expected to result from the self-assembly of the planar π -conjugated molecules into 1D columnar architectures, driven by $\pi\pi$ -stacking interactions. In such assemblies, coupling of electrons should give rise, *e.g.*, to ferromagnetism or low-energy absorption.⁶

The molecular structure of **1** can be varied basically by replacing carbon atoms – in the center and/or the in outer positions – by heteroatoms, thus configuring heteroatom-containing triangulene derivatives. A broad range of such structures has already

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Fig. 1 Schematic representation of the chemical structure of triangulene **1**. A representative AFM image of **1** deposited onto a Cu(111) surface is also shown (scale bar: 5 Å).³ Figure reproduced with kind permission; © 2017 Springer Nature.



Fig. 2 Schematic representation of the carbon-centered triangulonium and heteroatom-centered triangulene derivatives deduced from the parent compound **1**.

been realized and one can distinguish between two subclasses: the cationic carbon-centered and the neutral heteroatom-centered derivatives (Fig. 2). In general, these types of compounds are much better understood than parent **1** and their fundamental chemistry as well as their photophysical and electrochemical properties will, before focusing on their supramolecular chemistry, briefly be discussed in the following.

Preamble on heteroatom-containing triangulenes

Triangulonium cations are formed when the central C-atom of **1** is practically replaced by an sp^2 -hybridized C^+ center. Thereby, heteroatom-bridged and planarized dyes of the triaryl-methane family are formed (2–5, Fig. 3).⁷ The first example for this, planar sesquixanthylum cation (2, $TOTA^+$) was already reported in 1964,⁸ but this compound was somehow disregarded until the late 1990s when a number of functionalized derivatives⁹ or, even more important, the nitrogen-bridged counterparts to 2

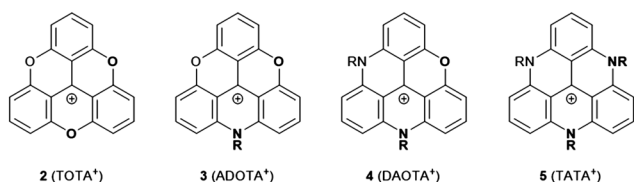


Fig. 3 Schematic representation of O- and/or N-bridged triangulonium cations **2–5** (R denotes an alkyl or aryl residue, the counterions are omitted for clarity).

were reported (3 to 5).^{10,11} Special credit has to be given to Laursen and his co-workers for establishing reliable protocols to prepare these triangulonium cations. All these derivatives are stable and efficient dyes which exhibit UV/vis absorption bands in the region from ca. 450 to 650 nm.⁷ ADOTA⁺ (**3**) and DAOTA⁺ (**4**) are strong emitters with fluorescence quantum yields (ϕ) above 40% in acetonitrile solutions (emission lifetimes (τ) longer than 20 ns). These particular dyes have been considered in the context of bio-imaging applications, *in vitro* as well as *in vivo*.¹² With respect to the electrochemical properties, a distinct trend can be observed for **2–5**: briefly, with increasing number of nitrogen atoms, the oxidation becomes easier. This behavior can be attributed to the stronger donor-character of the nitrogen atoms; consequently, the more oxygen-rich derivatives are much easier to be reduced (reduction potentials of -0.385 and -1.40 V for **2** and **5**, respectively). The ease of oxidation, in the case of TATA⁺ (**5**), was used by Gueret *et al.* for photosensitizing the light-driven hydrogen evolution reaction (HER) in the presence of a Co-based catalyst (remarkably, the TATA⁺ cation turned out to be a better photosensitizer than the benchmark $Ru(bpy)_3^{2+}$ under the experiments' conditions).¹³

For the neutral heteroatom-centered triangulenes a broad structural diversity is known. Concerning their synthesis two alternative strategies can basically be applied – the range of linear syntheses and the more universal divergent synthesis which was introduced recently by Hatakeyama and co-workers.¹⁴

The nitrogen-centered derivatives, either with isopropenyl (**6**) or carbonyl bridges (**7**), are already known since Hellwinkel's pioneering work in the 1970s.^{15–17} Whereas, their counterparts with higher group-15 elements in the center, *i.e.*, phosphorous or arsenic, were published much later.^{18–22} As detailed below, the latter ones possess concave, bowl-shaped molecular structures and, therefore, their self-assembly behavior will significantly differ from that of the planar N-centered molecules. These, in turn, resemble planarized derivatives of the propeller-shaped triaryl-amines which offer a beneficial property: extension of the π -conjugation through the central N-atom and, as a result from, improved materials properties when compared to analogous non-bridged triaryl-amines.²³ The photophysical and electrochemical properties of the core structures mainly depend on the nature of the bridging unit – ranging from a strongly electron-withdrawing carbonyl- (**7**) to a highly electron-donating oxygen-moiety (**8**, Fig. 4).^{24–26} It has been shown that these properties can be tuned over a wide range by appropriate modification of the parent structures.²⁷ The Venkataraman group focused on the

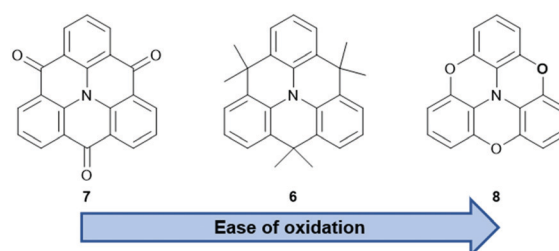


Fig. 4 Schematic representation of the N-centered triangulenes **6–8**.

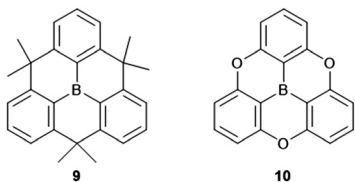


Fig. 5 Schematic representation of the B-centered triangulenes **9** and **10**.

photophysical properties and the self-assembly behavior of **7**²⁸ (*vide infra*) and it was realized that, after functionalization in the phenyl- and/or carbonyl-positions, materials with even semiconductor-like properties can be obtained.²³ The same also holds true for isopropenyl-bridged **6**: the incorporation into more complex architectures afforded materials with high ϕ values and large two-photon absorption (TPA) cross-sections.²⁹ A more detailed discussion of functionalized N-centered triangulenes can be found in Kivala's personal account on this topic.²³

Compared to their N-centered counterparts, the boron-centered derivatives are relatively young members of the heteroatom-containing triangulene family with their syntheses being reported within the last decade, in particular by Yamaguchi and co-workers.^{14,30–34} Basically, triarylboranes represent the electron-acceptor counterparts to the electron-donating triaryl-amines and, thus, they play an important role in various optoelectronic applications: organic-light emitting diodes (OLEDs), TPA materials or sensing of anions.^{35–37} The planarization of the triarylborane framework, as in **9** and **10** (Fig. 5) strongly effects the solid-state properties of the material by enabling intermolecular interactions to occur much more efficiently. It was shown, for example, that π -extended derivatives of **9** functioned as electron-transport materials in OLEDs.³¹ The packing of such discotic species into columnar architectures *via* $\pi\pi$ -stacking interactions afforded a liquid crystalline (LC) assembly with ambipolar charge-carrier characteristics.³²

To sum up this section, the various types of heteroatom-containing triangulenes represent important structures to be employed in optoelectronic applications. This can be attributed to their rich photophysical and electrochemical properties which, in turn, are strongly associated with their intermolecular interactions, *i.e.*, their supramolecular chemistry. The self-assembly behavior of triangulene derivatives, also in the context of potential applications, is in the spotlight of the subsequent sections.

Supramolecular assemblies based on C-centered triangulenium cations

The π -electronic triangulenium cations exhibit a self-assembly behavior that is strongly influenced by the accompanying anions. Thus, the supramolecular interaction of these compounds cannot be rationalized by the concepts of $\pi\pi$ -stacking interactions alone³⁸ – ion-pairing interactions between the π -electronic ions significantly contribute to the supramolecular ordering/packing of the components. Recently, Haketa and Maeda pointed out that three different assembly modes have to be considered in this



Fig. 6 Schematic representation of the three different packing modes known for π -electronic ionic species.³⁹ Figure reproduced with kind permission; © 2017 The Royal Chemical Society.

context (Fig. 6).³⁹ First of all, the two most extreme scenarios have to be named: the charge-by-charge and the charge-segregated ordering. In the former case, columnar heterostacks with alternating cationic and anionic molecules are formed (here, the $\pi\pi$ -stacking is facilitated by electronic attraction of opposite charges); whereas the latter one involves the stacking of molecules of identical charges (in such assemblies, which are rare, the electronic repulsion must be surpassed by even stronger attractive forces). The third (and most common) mode, which is the intermediate one, involves contributions from the other two modes.

In an initial study, Krebs *et al.* investigated the solid-state structures of the TOTA⁺ cations **2** and **11** as a function of their counterions.⁴⁰ Notable differences with respect to solid-state ordering of the molecules were observed: in detail, for the monovalent anions (*i.e.*, I[−], BF₄[−], AsF₆[−], PF₆[−], NO₃[−], CF₃SO₃[−]), a zigzag-type packing of cations and anions was revealed by X-ray diffraction (XRD) analysis (Fig. 7). On the other hand, the bulky divalent anions (*i.e.*, Mo₆Cl₁₄^{2−}, S₂O₆^{2−}) were found to be “sandwiched” between two TOTA⁺ sites. However, in both cases, the anions were detected in close proximity to the cationic centers of the TOTA⁺ molecules. The situation was somehow different in the case of (12)PF₆:⁹ this salt featured a pseudohexagonal-type of stacking with strong contributions from the charge-segregated mode in which the PF₆[−] anions were located close to the amino-substituents.

Laursen and co-workers investigated the formation of densely columnar packed Langmuir films at the water–air interface.⁴¹ For this purpose the amphiphilic TOTA⁺ derivatives **13**, comprising a



Fig. 7 Schematic representation of the TOTA⁺ cations **2**, **11** and **12**. The solid-state structure of (2)BF₄ is shown exemplarily (view along the *b*-axis of the *Pbca* unit cell).⁴⁰ Figure reproduced with kind permission; © 1999 International Union of Crystallography.

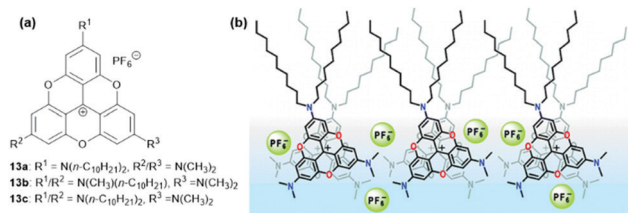


Fig. 8 (a) Schematic representation of the amphiphilic TOTA⁺ derivatives **13**. (b) Representation of the molecular packing of **13a** within the assembled LB films.⁴¹ Figure reproduced with kind permission; © 2009 American Chemical Society.

hydrophilic part, a rigid hydrophobic π -system and flexible hydrophobic alkyl chains were utilized. According to grazing incidence X-ray diffraction (GIXD) analysis, **13a** and **13b** yielded 2D-crystalline monolayers. In these structures, columnar stacks of the rigid-planar TOTA⁺ moieties were found, whereby these aggregates were well segregated from the accompanying PF₆⁻ counterions (a packing mode that was reminiscent of the solid-state structure of (12)PF₆,⁹ Fig. 8). The analytical data indicated an average column width which was almost identical to the diameter of the used molecules, *i.e.*, about 17 Å; further, a remarkably short repeat distance of 3.45 Å along the columns was found, thus suggesting an almost cofacial alignment of the TOTA⁺ cations with only a marginal tilting of the molecules relative to the columnar axis. Later, the same group also prepared multilayer films from **13a** by applying the Langmuir–Blodgett (LB) technique.⁴² These films featured the same type of supramolecular stacking as in the previous case. In-depth UV/vis absorption and emission studies gave evidence for an H-type aggregation of the molecules. Moreover, polarized fluorescence measurements revealed even a macroscopic alignment of the molecules resulting from the strong $\pi\pi$ -stacking interactions; thereby, the counterions were believed to be located between the single layers.

A different type of assembly was found in the case of the TATA⁺ cations **14**. Due to the presence of the aliphatic or etheric chains, smectic-type bilayer assemblies were found in the solid state.⁴³ Thereby, the layer-to-layer distance could be adjusted *via* the lengths of the pending chains which were oriented strictly perpendicular to the planar TATA⁺ moieties; the layers, in turn, were interwoven due to hydrophobic interactions of the chains. As a result, two TATA⁺ units were forced to stack in a face-to-face-type fashion, thereby overcoming their intrinsic electrostatic repulsion, eventually establishing a hexagonally ordered bilayer structure (Fig. 9). The authors processed the compounds into thin-solid films *via* spin-coating from solution. A high anisotropic polycrystallinity was observed in all these films, even on the cm length scale. The photophysical properties of the materials were investigated and suggested a potential use in opto-electronic applications due to their efficient exciton-transport behavior.⁴⁴ Noteworthy, the corresponding triaryl-substituted TATA⁺ derivatives did not show this self-assembly behavior due to the perpendicular orientation of the substituents relative to the planar molecular core.⁴⁵

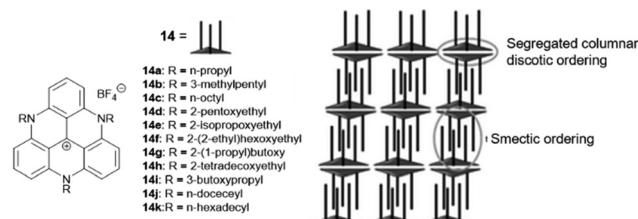


Fig. 9 Schematic representation of the TATA⁺ derivatives **14** that assembled into a smectic-type structure. A representation of the supramolecular packing as evidenced from XRD analyses is also shown.⁴³ Figure reproduced with kind permission; © 2012 The Royal Chemical Society.

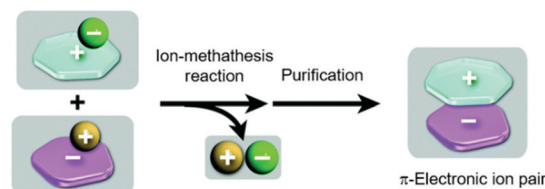


Fig. 10 Schematic representation of the synthesis of π -electronic ion pairs *via* a salt-metathesis reaction of appropriate couples of precursor salts.⁴⁶ Figure reproduced with kind permission; © 2013 The Royal Chemical Society.

In all previous examples “conventional” small-molecule anions were accompanying the triangulenium cations. In addition to these, also the assembly of genuine π -electronic ion pairs, *i.e.*, both the cation and the anion representing π -conjugated molecules, has been studied. Such species can, in principle, be obtained in a salt-metathesis reaction according to Fig. 10.^{39,46} It has to be noted that stable π -electronic anions are rare (compared to their cationic counterparts), since they prone to react with electrophiles. As one of the few exceptions, the cyclopentadienyl anions (Cp⁻) equipped with five strongly electron-withdrawing substituents (as the weak conjugated bases of superstrong organic acids⁴⁷) have to be named in this context.

For example, Maeda and co-workers investigated the self-assembly properties of (14a)(PCCp) in the solid state (PCCp: 1,2,3,4,5-pentacyanocyclopentadienyl).⁴⁸ X-ray crystal structure analysis revealed a regular packing of two pairs of 14a⁺ and PCCp⁻ ions in an alternating fashion (Fig. 11); whereas, the (CV)PCCp ion salt (CV: crystal-violet cation) exhibited a strictly alternating stacking of the CV⁺ and PCCp⁻ ions. The authors concluded that the highly ordered charge-by-charge assembly is disfavored when the π -electronic cation and anion strongly differ in their unit areas (as for the combination of 14a⁺ and PCCp⁻). Furthermore, the systems (14j)PCCp and (14k)PCCp, featuring long alkyl chains attached to the TATA⁺ cation, were investigated. These side chains were introduced to facilitate the dimension-controlled self-assembly of the ion pair due to additional van der Waals interactions.⁴⁸ Micrometer-scaled prisms were obtained in both cases from CH₂Cl₂/methanol solutions. These materials were characterized by (polarized) optical microscopy ((P)OM), atomic force microscopy (AFM) imaging and synchrotron XRD analysis. The latter one gave

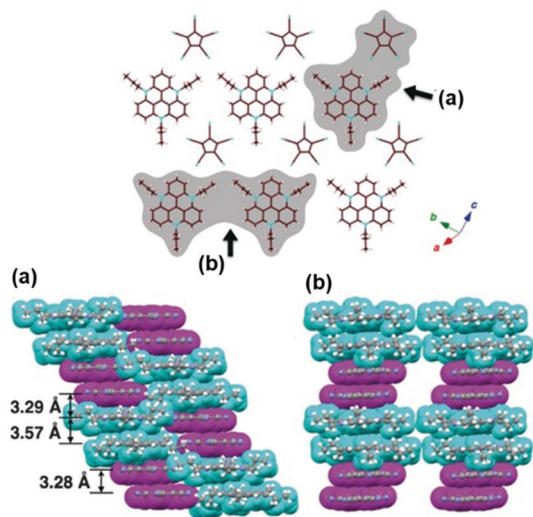


Fig. 11 X-ray single-crystal structure (**14a**)(**15a-Cl**). Top: Representative packing mode as top view of one layer. Bottom: Two different space-filling model side views (cyan and magenta units represent the cations and anions, respectively).⁴⁸ Figure reproduced with kind permission; © 2016 Wiley-VCH.

evidence for the presence of 2D anisotropically ordered lamellar phases within the macroscopic structures. The mesophases were identified by differential scanning calorimetry (DSC) measurements and POM imaging revealed textures with crystalline behavior. The samples melted into isotropic liquids at 169 °C and 176 °C, respectively.

The binding of anions to appropriate π -electronic receptor molecules represents a versatile alternative to obtain planar π -electronic anions.³⁹ Against this background, Maeda *et al.* introduced the BF_2 -complexes of 1,3-bis(pyrrol-2-yl)-1,3-diketones as efficient anion receptors for the formation of such anions and employed them in the context of supramolecular ion-pairing assembly (see *e.g.* ref. 39 and the literature cited therein). Among others, also the TATA^+ cations **14a**⁺, **14c**⁺, **14j**⁺ and **14k**⁺ were combined with the anions (**15-Cl**)[−] and (**15-Br**)[−] (Fig. 12) to yield supramolecular gels in alkane solvents and thermotropic LC materials.⁴⁹ The X-ray

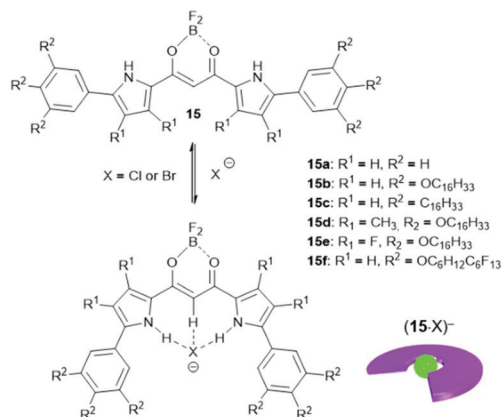


Fig. 12 Schematic representation of the formation of π -electronic anions by binding of a halide to the anion receptor **15**.

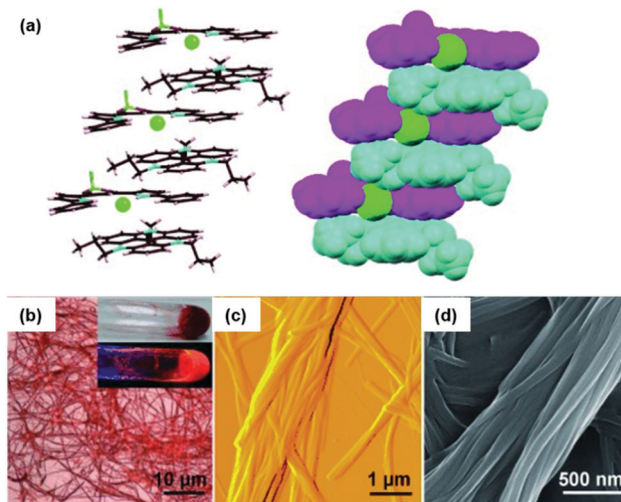


Fig. 13 (a) Solid-state packing of (**14a**)(**15a-Cl**) in the stick-type (left) and space-filling representation (right; **14a**, **15a** and Cl^- are depicted in cyan, magenta and green, respectively). (b–d) Microscopy images of the octane xerogel of (**14a**)(**15b-Cl**). (b) OM image (the inset shows pictures of the bulk sample upon visible (top) and UV light (bottom)), (c and d) representative AFM and SEM images, respectively.⁴⁹ Figure reproduced with kind permission; © 2010 Wiley-VCH.

single crystal analysis of (**14a**)(**15b-X**) ($\text{X} = \text{Cl}$ and Br) confirmed the expected packing of the planar π -electronic ions in a highly ordered charge-by-charge-type fashion (Fig. 13a). In all cases, supramolecular organogels with LC properties, comprising sub- μm -sized fibers, were observed (representative microscopy images are shown in Fig. 13b–d).

The LC behavior of ion-pairing assemblies based on TATA^+ cations was investigated further in particular focusing on the influence of the pending chains of the accompanying π -electronic anion. For (**14a**)(**15b-Cl**), the mesophase featured a focal-conic texture for which a triplet of transition temperatures of 94, 88 and 42 °C upon cooling from the isotropic liquid state (Iso) was identified by POM imaging (Fig. 14a).⁴⁹ The structure of the LC phase was studied by synchrotron XRD measurements and the observed peaks suggested a Col_h phase (Fig. 14b); the proposed stacking mode, *i.e.*, the formation of tetramers which assembled in a charge-by-charge-type fashion, was also present in the crystalline material (Fig. 13a).

The morphology of the assembly could be modified simply *via* the nature of substituents on the anion-receptor molecule. For example, the π -electronic salt (**14a**)(**15c-Cl**) yielded a Col_t mesophase at 50 °C upon cooling from the Iso instead of the Col_h one as in the previous case (Fig. 15a).⁵⁰ The XRD pattern featured a broad peak at which was assigned to a contribution of a charge-segregated packing mode to the overall assembly (Fig. 15b). This behavior was even more pronounced in the systems (**14a**)(**15d-Cl**) and (**14a**)(**15e-Cl**), where the stacking of π -electronic ions of identical charges was even dominant.⁵¹ The authors rationalized the strong contributions by charge-segregated stacking to the pronounced distortion of aryl rings due to the presence of the pyrrolic β -substituents (*i.e.*, CH_3 and F , respectively). As a consequence of this particular stacking behavior, ambipolar



Fig. 14 (a) POM image of **(14a)(15b-Cl)** at 87 °C. (b) XRD pattern of **(14a)(15b-Cl)** in the mesophase at 70 °C (for both measurements the Iso was cooled down with a rate of 2 °C min⁻¹). The proposed supramolecular stacking mode is also shown.⁴⁹ Figure reproduced with kind permission; © 2010 Wiley-VCH.

charge-transport (CT) properties with high mobilities were observed (*e.g.*, **(14a)(15e-Cl)** exhibited a mobility of negative charges in the range of $5 \times 10^{-3} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ at $4 \times 10^3 \text{ V}$). It was concluded that the structure of the counteranion strongly influences the conductivity of the ion-paired assemblies and, in particular, the aggregation in a charge-segregated mode is of importance to achieve high conductivity values (for examples see ref. 52).

For the π -electronic salt **(14a)(15f-Cl)** in which partially fluorinated side chains were present in the anion-receptor molecule, the mesophase was formed below a much higher

temperature as for the analogous non-fluorinated system **(14a)(15b-Cl)** (*i.e.*, 153 °C vs. 80 °C); the increased stability of the mesophase was attributed to the much stronger interaction of the semi-fluorinated aliphatic chains.⁵³ In these assemblies, a transient conductivity of $2.7 \times 10^{-5} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ was observed.

In another study, also the π -conjugated system of the anion-receptor species was extended. For example, the benzo[*c*]-fused analogues of **15** were studied in this context.⁵⁴ The π -electronic salts **(14a)(16-Cl)** were prepared and studied with respect to their self-assembly behavior. In the case of **(14a)(16b-Cl)**, a tetragonal Col_t mesophase was identified by POM imaging and XRD measurements (Fig. 16). In more detail, each disk of the column was formed by four sets of the ion pair; thereby, the flat cation and anion molecules formed the disks with the out-facing pending alkyl chains forming a hydrophobic shell along the columns.

The anion-receptor molecules were also equipped with bowl-shaped corannulene moieties, thereby expanding the π -electronic surface of the resulting anions.⁵⁵ In particular, the ion salt **(14a)(17-Cl)**, in which an unsymmetrical π -electronic anion was used, exhibited a rich self-assembly behavior. In 1,4-dioxane, for example, spherical aggregates with diameters in the range of 30 to 50 μm were identified by SEM imaging (Fig. 17a and b). However, a distinct mesophase could not be identified upon cooling of the Iso phase; instead, XRD measurements at 80 °C revealed a lamellar-type morphology with a much lower degree of ordering (compared to the aforementioned examples, Fig. 17c). Nonetheless, the LC materials showed a significantly increased charge-carrier mobility of up to $0.2 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$; this behavior was attributed to the presence of the fused corannulene units which offered a better overlap of the π -conjugated systems. In a similar fashion, also the phenanthro[*b*]-fused derivatives **18**



Fig. 15 (a) POM image of **(14a)(15c-Cl)** at 62 °C. (b) XRD pattern of **(14a)(15c-Cl)** in the mesophase at 62 °C (for both measurements the Iso was cooled with a rate of 2 °C min⁻¹). The proposed supramolecular stacking mode is also shown.⁵⁰ Figure reproduced with kind permission; © 2016 Wiley-VCH.

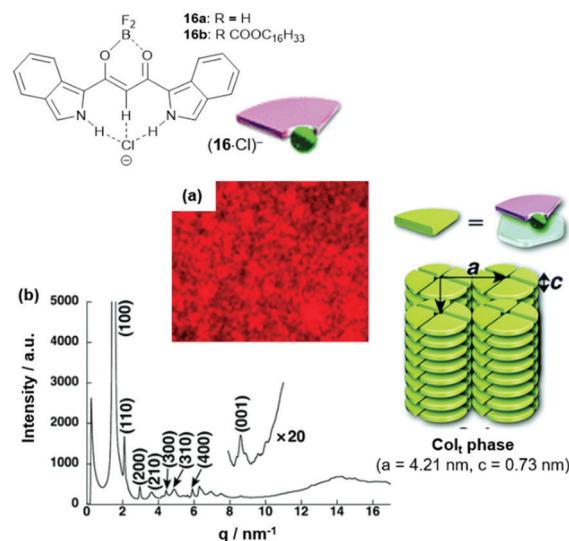


Fig. 16 Top: Schematic representation of the π -electronic anions **(16-Cl)⁻**. Bottom: (a and b) POM image and synchrotron XRD pattern of **(14a)(16b-Cl)**, respectively (both measurements were conducted at 80 °C); the proposed supramolecular stacking within the Col_t structure is also shown.⁵⁴ Figure reproduced with kind permission; © 2012 The Royal Chemical Society.

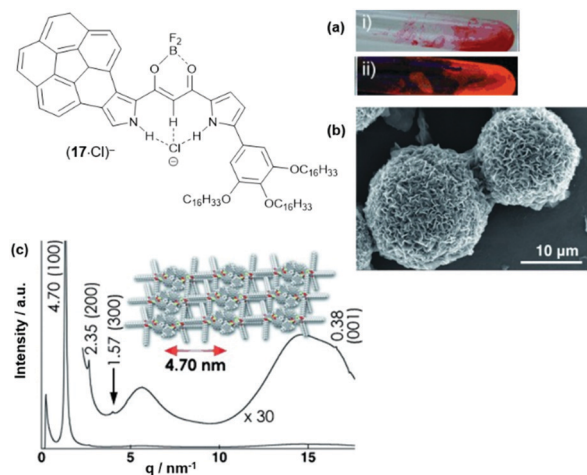


Fig. 17 Schematic representation of the π -electronic anion $(17\text{-Cl})^-$. (a) Photographs of $(14a)(17\text{-Cl})$ in 1,4-dioxane under visible (i) and UV light (ii). (b) SEM image of the precipitated $(14a)(17\text{-Cl})$. (c) XRD pattern of $(14a)(17\text{-Cl})$ at 80 °C on cooling from the iso phase; the proposed supramolecular structure of the assembly is also shown.⁵⁵ Figure reproduced with kind permission; © 2013 Wiley-VCH.

were employed for the complexation of chloride ions and, subsequently, the formation of π -electronic salts with TATA⁺ cations.⁵⁶ The anion-receptor molecules featured an increased affinity towards halide anions (attributed to additional C–H...Cl interactions with the phenanthroline unit) and, as result, the $(17\text{-Cl})^-$ were of highly planar structure. Accordingly, their π -electronic salts $(14a)(18\text{-Cl})$ yielded hexagonal columnar structures with liquid-crystalline mesophases which were analyzed by POM imaging and synchrotron XRD measurements (Fig. 29). For some of the systems $(14a)(18\text{-Cl})$, high photoconductivities were found in their anisotropically aligned mesophases (Fig. 18).

Flood and co-workers pointed out that the aforementioned anion-receptor molecules were able to bind only halide ions; whereas, bulky anions, *e.g.*, BF_4^- or PF_6^- , did not yield planar

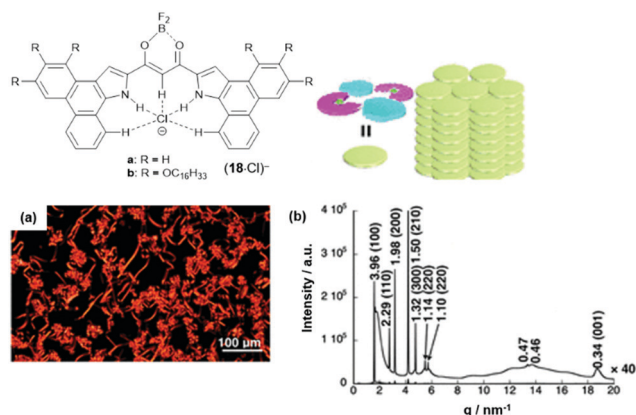


Fig. 18 Schematic representation of the phenanthro[b]-fused π -electronic anion $(18\text{-Cl})^-$. (a) POM image of $(14a)(18b\text{-Cl})$ at 180 °C upon cooling. (b) Synchrotron XRD pattern of $(14a)(18b\text{-Cl})$ at 150 °C upon cooling; the proposed supramolecular packing mode of the assembly is also shown.⁵⁶ Figure reproduced with kind permission; © 2019 American Chemical Society.

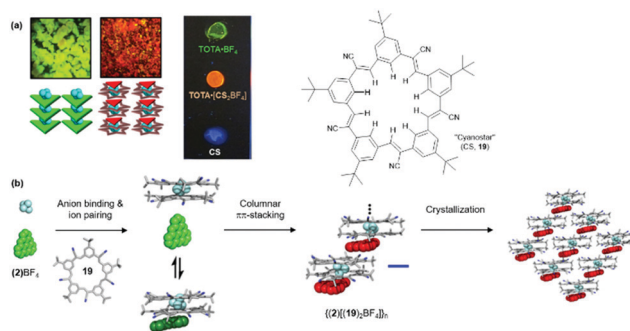


Fig. 19 (a) Photographs of the single crystals (left) and drop-cast films (right) showing the characteristic emission of the individual components or of the supramolecular assembly. (b) Schematic representation of the proposed self-assembly mechanism of $(2)(\text{BF}_4)$ and **19**.⁵⁸ Figure reproduced with kind permission; © 2017 American Chemical Society.

π -electronic anions due to a mismatch with the receptors' binding sites (thus, planar stacking of the chromophores was prevented). To overcome this limitation, these authors used the so-called "cyanostar" receptor (CS, **19**) which forms highly stable 2:1 sandwich-type complexes with large anions, *e.g.*, BF_4^- ($\log \beta_2 \approx 12$).⁵⁷ When combined with TOTA⁺ (**2**), the hierarchical self-assembly of the components gave a red-emitting material – both in solution as well as in the solid state, *i.e.*, single crystals and drop-cast films; note, that $(2)\text{BF}_4$ and **19** alone are green and blue emitting, respectively (Fig. 19a).⁵⁸ A four-step assembly mechanism was identified by NMR as well as UV/vis absorption and emission spectroscopy: binding of the anion to **19**, formation of an ion-paired salt, 1D columnar $\pi\pi$ -stacking and, eventually, crystallization (Fig. 19b). The photophysical studies revealed further that the supramolecular species $\{(2)[(19)_2\text{BF}_4]\}_n$ possessed its own excited-state manifold, significantly different from that of the TOTA⁺ cation. This CT state originated from the $\pi\pi$ -stacking interactions between the π -electronic ions. Thus, hierarchical ion-pairing assembly cannot only be applied to fabricate materials with LC behavior but also to form supramolecular architectures with rich opto-electronic properties. In both cases, the photophysical characteristic arise from the mixing of the ionic components and, therefore, represent a versatile tool in designing optical materials.

Supramolecular assemblies based on nitrogen-centered triangulenes

Due to their neutral structure, the N-heterotriangulenes exhibit a self-assembly behavior that basically results from $\pi\pi$ -stacking interactions. The solid-structure of **7** was reported by Field and Venkataraman in 2002.²⁸ The planar molecules densely packed into 1D columns which reported a herringbone-type ordering between the planes (plane-to-plane distance of 3.46 Å, Fig. 20). This packing mode already suggested the potential applicability of such planar building blocks to assemble nanostructures with opto-electronic and LC properties.

In order to improve the solubility and, by the same time, modulate the photophysical properties as well as the aggregation

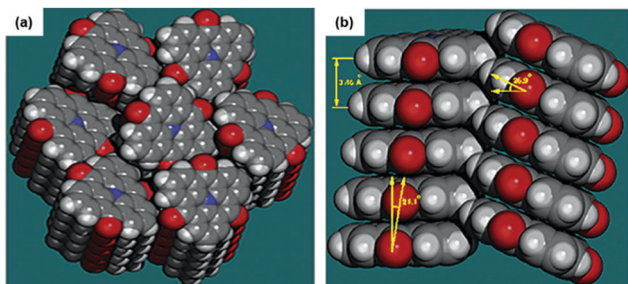


Fig. 20 Illustration of the solid-state structure of **7** (a: top view; b: side view).²⁸ Figure reproduced with kind permission; © 2017 American Chemical Society.

behavior, the parent N-heterotriangulene **7** was modified by introducing substituents on the phenyl moieties.²⁷ In one of the earliest examples in this respect, Zhang *et al.* attached carbazole-based dendrons (1st (G1) and 2nd generation (G2)) to the core structure of **7**.⁵⁹ These donor-acceptor-type featured an intramolecular CT transitions and aggregation-induced emission (AIE) was observed. The latter one is indicative for the planarization of the molecules due to intermolecular non-covalent interactions at high concentrations or in thin-solid films. For the derivatives with G1-substituents, a head-to-tail interaction (the so-called J-type aggregation) and not a columnar packing due to $\pi\pi$ -stacking interactions was concluded from UV/vis absorption and emission spectroscopy results, the derivatives with the large G2-substituents did not show this behavior, since the bulky dendrons prevented any non-covalent intermolecular interactions. Thus, placing sterically demanding moieties onto the core of **7** is beneficial for increasing the solubility and enables to adjust the photophysical and electrochemical properties, but is rather counterproductive with respect to the ability of the molecules to assemble into highly ordered arrays *via* non-covalent interactions.

Chen and co-workers introduced slim ethynylphenyl moieties by the Sonogashira-type cross-coupling reaction and investigated the self-assembly behavior of the resultant triangulene **20a** in a binary methanol/dichloromethane solvent mixture.⁶⁰ A two-level supramolecular polymerization of **20a**, driven by concerted $\pi\pi$ -stacking and hydrophobic interactions, yielded nanowires which eventually afforded well-organized microrods (Fig. 21). XRD and X-ray scattering measurements confirmed the long-range

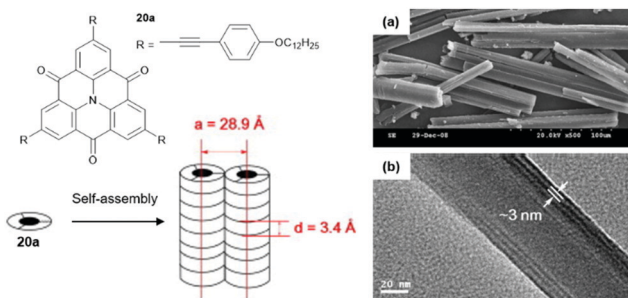


Fig. 21 Schematic representation of the two-step self-assembly of **20** into microrods. A representative SEM (a) and a TEM image (b) are also shown.⁶⁰ Figure reproduced with kind permission; © 2009 Elsevier B.V.

ordering within the assemblies. The diameter of the nanowires, as determined by TEM imaging, was in good agreement with the one for the 1D columnar stack as derived from the XRD analysis (*ca.* 3 nm).

Against the same background, Müllen and co-workers synthesized a small library of carbonyl-bridged N-centered triangulenes **20** in which long alkyl chains were attached to the core *via* different types of spacer units (Fig. 22).²⁴ The aliphatic chains enabled a good solubility of all derivatives in broad range of organic solvents; however, the chemical structure and bulkiness of the substituents as well as the processing conditions had a profound influence on the supramolecular self-assembly of **20**. In detail, the phenyl-bridged derivative **20c** yielded columnar disc-like assemblies in solution and in the solid state; the derivatives with ethynylphenyl spacers (**20d/e**), however, revealed such a behavior only in the solid state. Within the series, **20b** showed the lowest degree of aggregation in solution; whereas, well-aligned fibers could be fabricated by dip-coating from solution (Fig. 23a).^{24,61} As concluded from the electron-diffraction pattern, the highly crystalline fibers were formed by 1D columnar arrays of $\pi\pi$ -stacked molecules. In case of **20g**, the solid-state structure could be visualized by XRD analysis of single crystals. The cofacial $\pi\pi$ -stacking of the molecules gave rise to an overall columnar arrangement (Fig. 23b).²⁴ Thermotropic LC behavior was observed for the ethynylphenyl-equipped derivatives **20e/f**

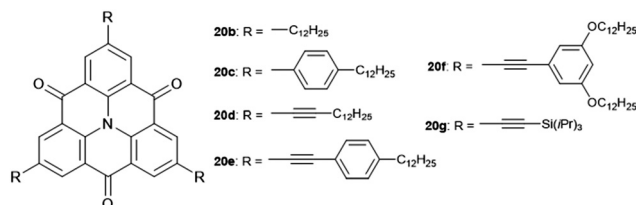


Fig. 22 Schematic representation of the two-step self-assembly of **20** into microrods. A representative SEM (a) and a TEM image (b) are also shown.⁶⁰ Figure reproduced with kind permission; © 2009 Elsevier B.V.

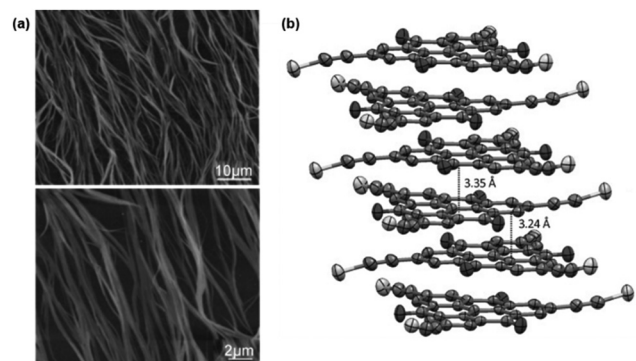


Fig. 23 (a) Representative tapping-mode AFM images of the fibers fabricated from **21b**.⁶¹ (b) Representation of the solid-state structure of **21g** (side view).²⁴ Schematic representation of the two-step self-assembly of **20** into microrods. A representative SEM (a) and a TEM image (b) are also shown.⁶⁰ Figure reproduced with kind permission; © 2011 and 2009 Wiley-VCH, respectively.

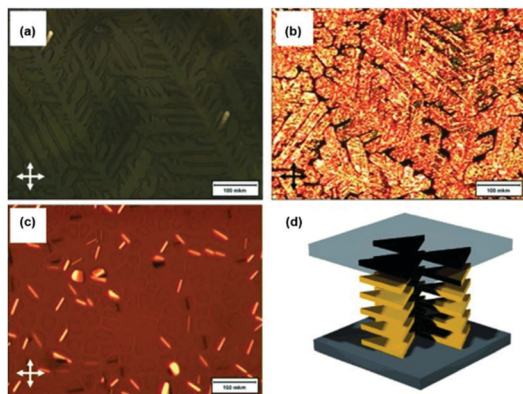


Fig. 24 POM images of **20e** (a: at 140 °C; b: at 30 °C) and **20f** (c: at 30 °C). (d) Illustration of the homeotropic packing of the discotic molecules.²⁴ Figure reproduced with kind permission; © 2009 Wiley-VCH.

which ordered themselves in homeotropic fashion (Fig. 24, no LC properties were found for the crystalline materials **20b–d**).

The multichromophoric compounds **21** were synthesized by Haedler *et al.* and the photophysical investigations revealed intramolecular CT processes (the direction of this depended on the chromophores in the periphery).⁶² The derivative **21b** formed a fluorescent organo-gel even in the sub-mM concentration regime; thereby, the CT characteristics, as observed in solution, were retained. According to AFM and TEM imaging, this gel comprised 1D nanofibers with molecular-scale diameters (Fig. 25a).⁶³ The fiber formation was attributed to a highly ordered cofacial $\pi\pi$ -stacking of the molecules, supported by hydrogen-bonding interactions involving the amide moieties of the pending groups. Remarkably, the fibers exhibited the ability to efficiently transport energy over μm distances, even at room temperature (*i.e.*, basically limited only by the fiber lengths). Photoluminescence imaging was applied to quantify this long-range energy transport (Fig. 25b and c).



Fig. 25 Schematic representation of the multichromophoric derivatives **21**. (a) Representative TEM image of the organo-gel formed by **21b**. (b) Representative photoluminescence image of a selected fiber. (c) Intensity profile along the orange dashed arrow in (b); the distance corresponds to the length along the arrow.⁶³ Figure reproduced with kind permission; © 2015 Springer-Nature.

The supramolecular polymerization of the chiral carbonyl-bridged N-heterotriangulenes **22** was studied by the Schmidt and the Meijer group.⁶⁴ For both enantiomers, two distinctly different self-assembly pathways were identified (Fig. 26). The authors proposed an isodesmic supramolecular process, which could be trapped by cooling to 7 °C (state A). Moreover, a cooperative nucleation–elongation gave rise to a thermodynamically stable state B at the same temperature (induced by first undercooling to –5 °C). An in-depth discussion of the mechanisms of these two prominent types of supramolecular polymerizations can be found elsewhere.⁶⁵ In both these states, a helical superstructure – arising from the same type of aggregation as for **21b** – was formed. Further, the supramolecular polymerization proceeded in a controlled fashion when two samples of the same enantiomer, one in state A and the other in state B, were combined; thereby, the kinetically trapped state evolved into the thermodynamically favored one. Remarkably, this process was found to be highly enantioselective – no such transition was identified if the states formed of different enantiomers were mixed (*e.g.*, combining (*R*)-**22** in the state A with (*S*)-**22** in the state B). It was concluded that a fundamental understanding of the self-assembly mechanism is as important as the structural design of the molecular components in order to create highly ordered functional supramolecular materials.

In 2014, Yamaguchi and co-workers reported that π -extension of parent **7** can also be achieved *via* the bridging positions.⁶⁶ This finding enabled the synthesis of π -extended derivatives featuring a rich photophysical and electrochemical behaviour.²⁷ In view of the supramolecular chemistry of these materials, the steric crowding around the exocyclic alkenyl moieties often prevents a planarization and, thus, the self-assembly into highly ordered architectures

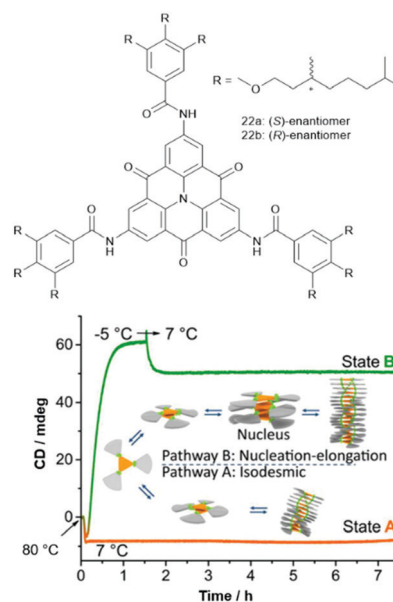


Fig. 26 Top: Schematic representation of the chiral N-heterotriangulenes **21**. Bottom: Illustration of the supramolecular polymerization of **21** which could be trapped in two different states by cooling.⁶⁴ Figure reproduced with kind permission; © 2015 Springer-Nature.



Fig. 27 Schematic representation of the N-heterotriangulenes **23** and **24** featuring a π -extension via their bridging positions.

is obsolete. For example, the structure of the derivatives **23** and **24** was found to be slightly bowl-shaped and not planar (as for their aforementioned carbonyl-bridged analogues **20b** and **20g**, Fig. 27).⁶⁷

Very recently, Dorca *et al.* demonstrated the pathway complexity of the 1D supramolecular polymerization of a chiral, dicyanovinyl-substituted derivative of **22**.⁶⁸ The authors claimed that their study represented the first example where the conformational change of a monomer enabled the pathway complexity of a supramolecular polymerization, *i.e.*, the formation of two different aggregates from one type of molecular unit. The bowl-shaped monomer **22** existed also in a planar, yet metastable conformation. Both of these conformers were found to assemble into different types of aggregates – a kinetic as well as a thermodynamic one (Fig. 28). The stability of these aggregates was high enough to enable a co-existence in solution at room temperature (the interconversion between the different aggregates was energetically unfavored due to the steric hindrance arising from the substituents in the bridging positions). However, heating of the mixture resulted in the disassembly of the aggregates and **22** adopted only the thermodynamically favored C_3 -symmetric conformation. Upon cooling down to room temperature, the conformer afforded exclusively helically shaped columnar assemblies with distinct circular dichroism (CD) signatures which could be correlated to the chiral monomers.

In contrast to the rigid-planar carbonyl-bridged N-heterotriangulene **7**, its dimethylmethylene-bridged counterpart **6** is slightly more flexible and a shallow bowl-shaped structure was verified by X-ray single crystal analysis.²⁹ Moreover, the sp^3 -hybridized bridge atoms introduce an out-of-plane steric effect which basically excludes any cofacial $\pi\pi$ -stacking interactions (at least in the solid state). On the other hand, **6** is much more soluble and represents, due to the electron-rich nature,

a useful precursor for introducing substituents in the *para*-positions of the phenyl rings. Consequently, a broad range of derivatives has been synthesized in the last decade and were, due to their planarized structure, found to exhibit a much higher degree of π -conjugation when compared to the members of the related triarylamine family.²⁷ The π -extended derivatives of **6** have, for instance, been used as hole-transport material in organic light-emitting devices (OLEDs), as photosensitizers in solar-cell devices or in non-linear optics due to their two-photon absorption behavior.

The dimethylmethylene-bridged N-heterotriangulenes do not enable a supramolecular self-assembly into 1D columnar materials. However, the straightforward threefold symmetrical functionalization of parent **6** yields derivatives suited for an assembly into 2D structures, *i.e.*, sheet-like motives. This approach can be realized on the basis of covalent as well as supramolecular chemistry. As an example for the formation of covalent 2D structures derived from small-molecule N-heterotriangulene precursor molecules, the work of Bieri *et al.* has to be mentioned: the formation of such a sheet-like structure was achieved by assembling the symmetrical tribromo-derivative of **6** onto a Ag(111) surface, followed by thermal treatment at 300 °C.⁶⁹ The same strategy was followed by Steiner *et al.* to prepare a covalent 2D network of carbonyl-bridged N-heterotriangulenes on an Au(111) surface.⁷⁰ In order to utilize supramolecular chemistry for the very same purpose, the substituents attached to **6** need to function as supramolecular binding sites to force the molecular components into highly-ordered arrays on the basis of strong and directional, yet reversible non-covalent interactions (Fig. 29). In such a supramolecular process, the non-covalent bonds are formed and cleaved until eventually an equilibrium is reached in which the assembly features a high degree of order and – ideally – only a small number of structural defects can be found.⁷¹

In this respect, the cyano group represents, despite its chemically simple structure, already a highly versatile functionality due to the highly asymmetric charge distribution that can be utilized to form ordered structures *via* intermolecular dipole-dipole⁷² or hydrogen-bonding interactions⁷³ as well as, even more efficiently, by metal-to-ligand coordination.⁷⁴ Against this background, Kivala *et al.* prepared the threefold cyano-functionalized N-centered triangulene **25** and studied its adsorption onto coinage-metal surfaces, *i.e.*, Cu(111), Ag(111) and Au(111).^{75,76} Scanning tunneling microscopy (STM), low-energy electron diffraction, X-ray photoelectron spectroscopy (XPS) and density functional



Fig. 28 Illustration of the pathway complexity of a dicyanovinyl-equipped N-centered triangulene – two different types of supramolecular aggregates are accessible from the two conformers.⁶⁸ Figure reproduced with kind permission; © 2019 American Chemical Society.

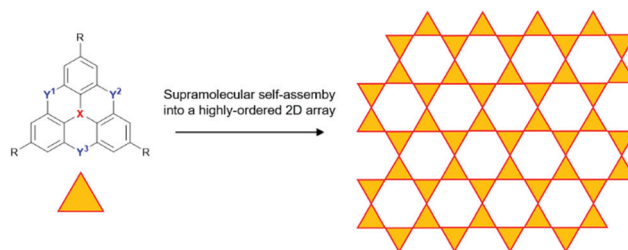


Fig. 29 Idealized schematic representation of the self-assembly of heteroatom-containing triangulenes into a supramolecular sheet-like array.

theory (DFT) calculations allowed the authors to identify all three above-mentioned bonding motifs. A hexagonally densely packed pattern of **25**, stabilized by hydrogen-bonding interactions, was observed in the case of Ag(111). In contrast to this, two distinctly different partially porous phases were identified in the submonolayer coverage of Au(111) – the densely-packed α -phase originated from an interplay between dipolar-coupling and hydrogen-bonding interactions; whereas, the formation of the (filled) β -phase was explained by additional metal-to-ligand coordination involving metal centers of the Au(111) substrate (Fig. 30). However, on a Cu(111) surface only small areas of hexagonally close-packed sheets stabilized by metal-to-ligand coordination and areas of rather disordered molecules could be found. Thus, the role of the substrate also needs to be considered carefully when aiming for the assembly of supramolecular sheets on solid supports. The particular relevance of such porous sheets with respect to materials science, in particular when aiming for nanoelectronic devices, is highlighted by the ferromagnetic half-metallicity reported for a covalent material composed of a related N-centered triangulene monomer.⁷⁷

The Kivala group also transformed the CN-groups into carboxylic acids (**26a**) and 2,4-diamino-1,3,5-triazin-6-yl (DAT) units (**26b**), thereby introducing moieties designated for the self-assembly into 2D structures *via* self-complementary hydrogen-bonding interactions.²³ Porous 2D and 3D structures formed by hydrogen-bonding interactions have attracted considerable attention and have recently been reviewed elsewhere.⁷⁸ The solid-state structure of **26b** was investigated by X-ray crystallographic analysis and confirmed the formation of regular planar sheets in which the molecules were linked by DAT-dimers (supported by hydrogen-bonding interactions involving co-crystallized solvent molecules). Besides the formation of supramolecular sheets on Au(111) of either of the monomers, also the co-adsorbance of **26a** and **26b**, representing a donor- and an acceptor-type molecule, was reported (Fig. 31).⁷⁹ Presumably due to the maximization of the overall number of hydrogen bonds on the cost of the optimization of the hydrogen-bonding strength a mixed network was established. DFT calculations revealed that the band gap between the highest occupied and the lowest unoccupied molecular orbitals (*i.e.*, the HOMO and the LUMO)



Fig. 30 Schematic representation of the CN-functionalized N-centered triangulene **25**. The adsorption onto an Au(111) substrate yielded two different phases as visualized by STM imaging (scale bar: 12.5 nm).^{75,76} Figure reproduced with kind permission; © 2014 Wiley-VCH.



Fig. 31 Top: Schematic representation of the functionalized N-centered heterotriangulenes **26**; the proposed hydrogen-bonding interaction between the two entities is also shown. Bottom: STM images of the supramolecular sheets formed by **26a**, **26b** as well as a 1:1 mixture of **26a** and **26b** (from left to right).⁷⁹ Figure reproduced with kind permission; © 2018 The Royal Chemical Society.

was defined by both molecules, a fundamental characteristic of donor-acceptor assemblies.

Supramolecular assemblies based on boron-centered triangulenes

In 2012, Yamaguchi and co-workers reported on the synthesis of the alkyl-bridged boron-centered triangulene **9**; the proposed planar structure of this bridged triphenylborane molecules could be verified by single-crystal XRD analysis (Fig. 32).³³ However, intermolecular $\pi\pi$ -stacking interactions were prevented due to steric hindrance of the dimethylmethylene bridges (similar to the N-centered analogue **6**); instead, a face-to-edge-type orientation of the molecules was found. The B-centered derivative **9** was highly stable against air, moisture, amines and heat. This behavior was rationalized by the structural constraint in **9** (which is absent in the parent triphenylborane), not requiring any further stabilizing bulky substituents. Moreover, the electron-deficient boron atoms retained their Lewis acidity and could reversibly coordinate fluoride ions; thereby, the molecular structure of **9** experienced a plane-to-bowl interconversion.



Fig. 32 Top (left) and side view (right) of the solid-state structure of **9**.³³ Figure reproduced with kind permission; © 2012 American chemical Society.

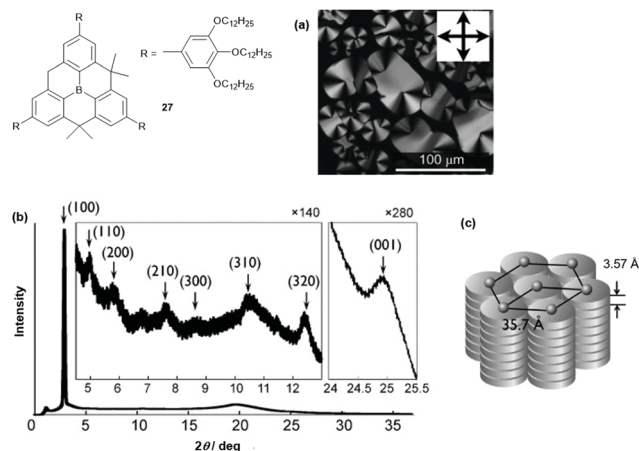


Fig. 33 Schematic representation of **27**. (a) POM image of **27** (at 70 °C upon cooling from the isotropic liquid; the inset shows the directions of the polarizer and analyzer). (b) XRD pattern of **27** (at room temperature after cooling from the isotropic liquid). (c) Schematic representation of the proposed hexagonal columnar phase of **27**. Figure reproduced with kind permission; © 2012 American chemical Society.

Decreasing the steric demand of the bridges allows for a much stronger intermolecular interaction and columnar assemblies can be obtained. This was shown by Yamaguchi *et al.* who studied the unsymmetrical derivative **27**.³² According to DSC and XRD measurements, a $\pi\pi$ -stacked hexagonal columnar LC phase was assembled from the discotic compound at room temperature (disc-to-disc distance of *ca.* 3.6 Å, Fig. 33). The photophysical properties were investigated and revealed a blue-shifted and reduced fluorescence of **27** in the LC phase relative to that found in THF solution (LC phase: $\lambda_{\text{em}} = 427$ nm ($\phi_{\text{F}} = 0.61$); THF solution: $\lambda_{\text{em}} = 457$ nm ($\phi_{\text{F}} = 0.99$)). This finding was attributed to the dense packing of **27** in the LC phase, where the interconversion between the planar and the bowl-shaped conformation was impeded. The ordered columnar stacking and the short intermolecular distances in the LC assembly resulted in an ambipolar charge-carrier behavior: electron and hole mobilities of 10^{-3} and 3×10^{-5} cm² V⁻¹ s⁻¹, respectively, were determined by time-of-flight measurements.

The planar heteroatom-bridged B-centered derivatives **28** and **29** were both reported very recently (Fig. 34).^{14,34} In contrast to the carbon-bridged analogue **9**, the oxygen-bridged triangulene **28** exhibited intermolecular interactions in the solid state: due to the absence of steric hindrance, a $\pi\pi$ -stacking motive with a parallel offset and a plane-to-plane distance of *ca.* 3.38 Å was formed (Fig. 35).³⁴ However, the phenyl substituents of **29**, oriented perpendicular to the planar triangulene unit, prevented such a packing for steric reasons.



Fig. 34 Schematic representation of the heteroatom-bridged boron-centered triangulenes **28** and **29**.



Fig. 35 (a) Solid-state structure of **28** (left: top view; right: side view). (b) Cross section of the solid-state packing.³⁴ Figure reproduced with kind permission; © 2016 The Royal Chemical Society.

Supramolecular assemblies based on bowl-shaped heterotriangulenes

In contrast to the planar N-centered triangulenes, their phosphorous-centered counterparts were persistent bowl-shaped or convex/concave structures. This difference results from the presence of the P-atom in a threefold respect: firstly, the significantly larger covalent radius of phosphorous (when compared to B, C and N) urges the atom out of plane even when incorporated into a π -conjugated system.⁸⁰ Secondly, the molecular structure is configurationally rather stable due to the large energy barrier for the pyramidal inversion.⁸¹ Finally, Bent's rule prevents phosphorous from being hybridized easily (P is by far more electropositive as C or N).⁸² The dimensions of the bowl-shaped P-centered triangulenes are defined by two parameters – the bowl depth and the cone angle (d and θ , respectively, Fig. 36). These parameters, in turn, are of relevance when considering the binding of appropriate guest molecules within the bowl (*e.g.*, fullerenes, *vide infra*). Moreover, the structural and electronic features of such molecules also depend on the oxidation state of the central phosphorous atom: the lone electron pair of a trivalent P-center gives rise to a Lewis basic character; whereas, electronegative substituents in pentavalent species are known to induce Lewis acidity.



Fig. 36 Schematic representation of triphenylphosphine and its axially occupied derivatives. The bowl depth (d) and the cone angle (θ) represent the key parameters when structural constraining the parent structure.²⁷ Figure reproduced with kind permission; © 2019 American Chemical Society.



Fig. 37 Schematic representation of the fundamental packing modes of bowl-shaped molecules.²² Figure reproduced with kind permission; © 2016 American Chemical Society.

Yamamura *et al.* pointed out that the molecular structure has a profound impact on the packing behavior and two different types of packing modes can basically be obtained (Fig. 37): ordered 1D concave/convex stacks (A and B) and less ordered assemblies (*e.g.*, C).²²

A variety of O- and/or S-bridged P-centered triangulenes **30** was prepared and their self-assembly behavior in solution as well as in the solid state was investigated (Fig. 38). The C_{3v} -symmetric O-bridged derivative **30a** with pyroelectric properties was reported by Krebs *et al.* already in 1997.²¹ The low sublimation temperature of **30a** suggested only weak intermolecular interactions; however, a highly ordered packing mode due to $\pi\pi$ -stacking interactions was observed in the crystalline state (Fig. 39). The authors proposed a sufficiently high dipole moment (μ) as the main driving force for this self-assembly behavior. This assumption was verified later by applying multipolar modelling studies: the dipole moment of **30a** was enhanced by *ca.* 42% when going from the solubilized to the crystalline state ($\mu = 4.7$ D).⁸³

The S-containing derivatives **30b–d** were reported by Yamamura *et al.*²² With increasing number of S-bridges, the cone angles increased and, consequently, the bowl depths decreased



Fig. 38 Schematic representation of the O- and/or S-bridged P-centered triangulenes **30**.



Fig. 39 Representation of the molecular packing of **30a** in the crystalline state. Figure reproduced with kind permission; © 1997 American Chemical Society.

(**30a**: $d = 2.43$ Å; **30d**: 2.03 Å). Not only the molecular parameters, but also the packing behavior was affected by the nature of the bridges. The C_{3v} -symmetric derivative **30d** again assembled into highly ordered 1D columnar stacks of type B (according to Fig. 37). On the contrary, the C_s -symmetric representatives **30b** and **30c** stacked in a less ordered zig-zag-type mode.

The same authors also investigated the self-assembly behavior of various “bowl chiral” derivatives of **30a** (Fig. 40). Basically, the self-assembly of such chiral bowl-shaped molecules yields materials with supramolecular chirality which, in turn, might be used to create materials applicable in polarization-related photonics or chiral organic conducting systems (the chiroptical switching represents the base in this context).⁸⁴

In the crystalline state, *rac*-**31a** featured an ordered hexagonal array of anisotropic 1D columns of either enantiopure (*M*)-**31a** or (*P*)-**31a** molecules $\pi\pi$ -stacked in a convex/concave-type fashion.²⁰ Thereby, the alignment of the columns corresponded to the type-B packing (Fig. 37) in which each enantiopure column was directly opposed to the neighboring one, which was formed by molecules of the second enantiomer (Fig. 41a). The XRD analysis of single-crystalline (*M*)-**31a**, however, revealed a different, *i.e.*, an orthorhombic packing mode: each convex/concave columnar

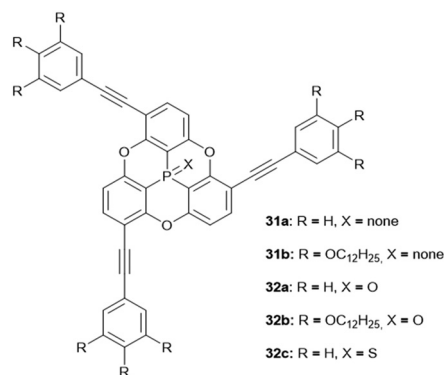


Fig. 40 Schematic representation of the chiral bowl-shaped P-centered triangulenes **31**.



Fig. 41 Representation of the packing of *rac*-**31a** (a) and (*M*)-**31a** (b) in the crystalline state.²⁰ Figure reproduced with kind permission; © 2015 The Royal Chemical Society.

$\pi\pi$ -stack was surrounded by five columns – four of these aligned in the opposite and one aligned in the same direction (Fig. 41b).

The derivative **31b** possessing alkoxy side chains assembled into columnar LC phases.²⁰ According to DSC and XRD measurements, the LC phases derived from the enantiopure compounds were more ordered than the one formed by racemic **31b**. Remarkably, the circular dichroism (CD) signal of the thin-solid film of enantiopure **31b** had the opposite signature as in solution. This difference could directly be attributed to the self-assembly of the molecules in the condensed state. The reversibility of the chiroptical switching of enantiopure **31b** was demonstrated by the multiple transition between the anisotropic liquid and the LC phases.

The N-bridged heterotriangulene **32** was reported only recently and did, due to steric hindrance of the pending phenyl moieties, not yield any ordered assemblies.¹⁴ The arsenic-centered triangulenes possess similar molecular structures as their phosphorous-containing counterparts;¹⁸ however, their self-assembly behavior in solution or in the solid state has not been discussed in the scattered reports on these compounds. The first example for a silicon-centered triangulene derivative was published by Hatakeyama and co-workers; the values of bowl parameters were in the same range as those reported for the similar P-centered derivatives.¹⁴ Thus, an application of these young members of the heterotriangulene family in the field of supramolecular host-guest complexation appears feasible. On the contrary, the steric hindrance arising from the pending substituents in concert with the presence of an axial substituent on the heteroatom is expected to prevent the formation of ordered supramolecular convex/concave $\pi\pi$ -stacks. The same also holds true for the various examples for axially-substituted phosphorous-containing triangulenes.²⁷

Beyond 1D and 2D assemblies: miscellaneous supramolecular architectures based on heteroatom-containing triangulenes

Bowl-shaped molecules have attracted considerable attention in the field of supramolecular chemistry due to their ability to bind spherical (convex) guest molecules to their concave face *via* van der Waals, $\pi\pi$ -electronic and charge-transfer interactions.⁸⁵ In particular, the recognition of pristine fullerene C_{60} represents an intensive field of research in this respect.⁸⁶ For example, the bowl-shaped hydrocarbons corannulene⁸⁷ and sumanene⁸⁸ have been reported as potent host molecules, since their curved structures resemble fullerene subunits and, thus, exactly match C_{60} 's convex surface.

Within this context, Yamamura *et al.* utilized the chiral P-centered triangulene P-oxide (*P*)-**32a** as bowl-shaped host for the complexation of C_{60} .¹⁹ In the crystalline state four host molecules were found to almost perfectly cover the guest's surface (Fig. 42). Remarkably, the host-guest interaction was strong enough to withstand ionization: a species assigned to

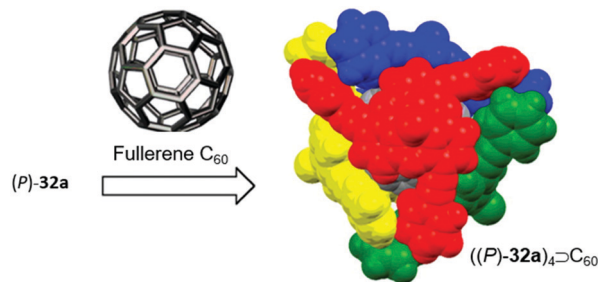


Fig. 42 Schematic representation of the formation of a host-guest complex by coordination of four chiral (*P*)-**32a** host molecules to a fullerene C_{60} guest.¹⁹ Figure reproduced with kind permission; © 2014 American Chemical Society.

the (*P*)-**32a** $\supset C_{60}$ 1.1 complex was observed in the matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) mass spectrum of the crystalline sample. Since the *P*-sulfide **32c** featured almost the same bowl dimensions as **32a**, (*P*)-**32c** yielded a similar host-guest complex of 4 : 1 stoichiometry with C_{60} in the solid state.²⁰ Due to the lone electron pair, **31a** possessed a deeper bowl and, consequently, a sandwich-type 2 : 1 complex was formed with C_{60} (Fig. 43). This type of architecture is reminiscent of Haino's biscalix[5]arene- C_{60} adducts; the CT interaction between biscalix[5]arene-based molecular tweezers and fullerene derivatives was utilized by these authors to assemble discrete supramolecular architectures as well as linear, side-chain or cross-linked supramolecular polymers.⁸⁹ It may, thus, be concluded that also the ((*P*)-**31a**)₂ $\supset C_{60}$ motive can be adapted for the very same purpose. However, reports targeting this application are not yet at hand.

From the highly directional types of non-covalent interactions, halogen bonding (XB) has emerged as a powerful tool in various fields of chemical research – ranging from medical or catalytic applications to crystal engineering or materials science.^{90,91} Though being still under scientific debate, XB interactions are



Fig. 43 Top: Illustration of the coordination of a P-centered triangulene with a deep (**31a**) and a shallow bowl (**32a/c**) to C_{60} . Bottom: Space-filling model of the ((*P*)-**31a**)₂ $\supset C_{60}$ complex.²⁰ Figure reproduced with kind permission; © 2015 The Royal Chemical Society.

generally described by Clark's σ -hole concept: basically, the attractive interaction results from an electrophilic region on a molecule's halogen atom (the so-called σ -hole) and an electron-rich sites in another (or the same) molecule.^{92,93} 1,2,4,5-Tetrafluoro-3,6-diiodobenzene represents a prominent ditopic XB donor which was employed by Schaub *et al.* for the co-crystallization with **33** in a 1:2 ratio.⁹⁴ X-Ray crystallographic analysis of the single crystals revealed a packing mode in which XB-donor molecules were surrounded by in total six molecules of **33** (Fig. 44). Two of these were coordinated in a linear fashion to the aryl iodide by strong $\text{P}=\text{O} \cdots \text{I}$ XB-type interactions (the strength of this interaction was reflected by the very short bonding length of *ca.* 2.683 Å). The other four molecules of **33** contributed by $\pi\pi$ -stacking, $\text{H} \cdots \text{F}$ as and $\text{Ar}-\text{H} \cdots \pi$ interactions to the overall packing mode. This (up to now) exclusive example shows how non-covalent interactions, beyond the $\pi\pi$ -stacking of planar or bowl-shaped molecules, can be utilized to force triangulene derivatives into ordered arrays.

Finally, the planar triangulene scaffold has been used as template for the formation of metal complexes. Lacour and co-workers designed a tripodal ligand for the coordination of lanthanide(III) ions in which three tridentate binding sites were attached to the planar TATA⁺ unit (**34**, Fig. 45).⁹⁵ The resultant Eu(III) complex of **34** was characterized by ¹H NMR spectroscopy, mass spectrometry and single-crystal XRD analysis.



Fig. 44 Top: Schematic representation of the P-centered triangulene **33**. Bottom: X-ray single-crystal structure of the 2:1 complex of **33** and 1,2,4,5-tetrafluoro-3,6-diiodobenzene (thermal ellipsoids at 50% probability, hydrogen atoms omitted for clarity). (a) Illustration of the packing into chain-like assemblies. (b) Illustration of the linear halogen-bonding interaction contributing to the overall packing mode.⁹⁴ Figure reproduced with kind permission; © 2017 Wiley-VCH.

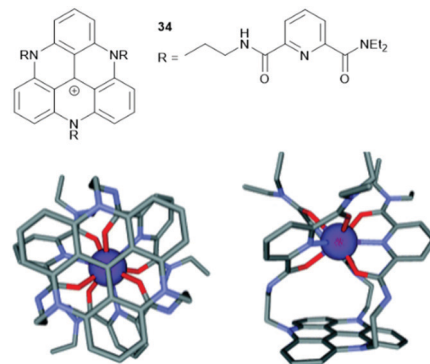


Fig. 45 Top: Schematic representation of the tripodal ligand **34**. Bottom: Illustration of the structural model [Eu(**34**)]⁴⁺ (Scigress software, hydrogen atoms are omitted for clarity); a top view (left) and a side view (right) are shown.⁹⁵ Figure reproduced with kind permission; © 2012 The Royal Chemical Society.

Molecular-modelling studies were performed and supported the experimental data (Fig. 45). Further, [Eu(**34**)]⁴⁺ exhibited the emission behavior as **34** itself; thus, energy transfer from the excited TATA⁺ core to the Eu(III) center could be excluded.

The last examples show how hydrogen-/halogen-bonding interactions and metal-to-ligand coordination can be combined with heteroatom-containing triangulene moieties. It can be speculated that the proper combination of these non-covalent forces with the $\pi\pi$ -stacking interactions offered by π -conjugated molecular structure might be of high virtue – not only in crystal engineering but also for the controlled supramolecular assembly of triangulene-based hierarchical 2D and even 3D supramolecular architectures.⁹⁶ Such (metal-) organic frameworks might be employed in various opto-electronic applications, such as the harvesting of light or photocatalysis.⁹⁷

Conclusions and outlook

The basic structures of the various types of heteroatom-containing triangulenes have already been known since some decades; however, applying their supramolecular chemistry as a tool to fabricate materials with tunable liquid-crystalline or opto-electronic properties has emerged as field of research rather recently. In particular, the planar boron- or nitrogen-centered representatives have been utilized for the formation dimension-controlled columnar assemblies.²⁷ The same also holds true for the triangulenium cations which have been combined with appropriate anions to afford π -electronic salts.³⁹ For these assemblies opto-electronic applications have already been envisioned. Moreover, the stacking of concave/convex or bowl-shaped derivatives offers one further opportunity to obtain 1D columnar assemblies. The C_3 -symmetric derivatives equipped with appropriate functionalities have been utilized for the self-assembly into 2D nanosheets *via* directional non-covalent interactions (*e.g.*, dipole-dipole and hydrogen-bonding interactions, metal-to-ligand coordination). In addition to this, various non-covalent interactions play a pivotal role in the solid-state packing of functionalized triangulene derivatives,

an aspect which is of relevance with respect to crystal engineering (*i.e.*, tuning the structural or photophysical properties in the solid state).

A broad range of coordination nanosheets (CONASHs) has been prepared from symmetrical planar building blocks and (transition) metal ions.⁹⁷ Such materials are known to exhibit electron or proton conductivities, electrochromism as well as photo- or (photo)-electrocatalytic activities. Surprisingly, triangulene (or triangulenium) dyes have not yet been discussed in this context despite their rich photophysical properties predestined for such purposes.¹³ The same holds true for metal-organic frameworks, the 3D analogues to CONASHs, which are currently of particular interest regarding light-harvesting⁹⁸ and water-splitting applications.⁹⁹ We expect that such functional materials can also be derived from heteroatom-containing triangulene (or triangulenium) derivatives. However, the assembly of such hierarchically structured supramolecular 2D and 3D assemblies is non-trivial and requires a cautious control over the interplay between the $\pi\pi$ -stacking interactions and the other involved non-covalent forces (*i.e.*, hydrogen/halogen bonding interactions, metal-to-ligand coordination). We look forward to see such complex supramolecular assemblies to be realized and, even more, to be brought into action.

Conflicts of interest

The authors declare no conflicts of interest.

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References

- 1 Z. Zeng, X. Shi, C. Chi, J. T. López Navarrete, J. Casado and J. Wu, *Chem. Soc. Rev.*, 2015, **44**, 6578–6596.
- 2 E. Clar and D. G. Stewart, *J. Am. Chem. Soc.*, 1953, **75**, 2667–2672.
- 3 N. Pavlíček, A. Mistry, Z. Majzik, N. Moll, G. Meyer, D. J. Fox and L. Gross, *Nat. Nanotechnol.*, 2017, **12**, 308.
- 4 Y. Morita, S. Suzuki, K. Sato and T. Takui, *Nat. Chem.*, 2011, **3**, 197.
- 5 I. Ratera and J. Veciana, *Chem. Soc. Rev.*, 2012, **41**, 303–349.
- 6 V. Sharma, N. Som, S. D. Dabhi and P. K. Jha, *Chemistry-Select*, 2018, **3**, 2390–2397.
- 7 J. Bosson, J. Gouin and J. Lacour, *Chem. Soc. Rev.*, 2014, **43**, 2824–2840.
- 8 J. C. Martin and R. G. Smith, *J. Am. Chem. Soc.*, 1964, **86**, 2252–2256.
- 9 B. W. Laursen, F. C. Krebs, M. F. Nielsen, K. Bechgaard, J. B. Christensen and N. Harrit, *J. Am. Chem. Soc.*, 1998, **120**, 12255–12263.
- 10 B. W. Laursen and F. C. Krebs, *Angew. Chem., Int. Ed.*, 2000, **39**, 3432–3434.
- 11 B. W. Laursen and F. C. Krebs, *Chem. – Eur. J.*, 2001, **7**, 1773–1783.
- 12 B. P. Maliwal, R. Fudala, S. Raut, R. Kokate, T. J. Sørensen, B. W. Laursen, Z. Gryczynski and I. Gryczynski, *PLoS One*, 2013, **8**, e63043.
- 13 R. Gueret, L. Poulard, M. Oshinowo, J. Chauvin, M. Dahmane, G. Dupeyre, P. P. Lainé, J. Fortage and M.-N. Collomb, *ACS Catal.*, 2018, **8**, 3792–3802.
- 14 S. Nakatsuka, H. Gotoh, K. Kinoshita, N. Yasuda and T. Hatakeyama, *Angew. Chem., Int. Ed.*, 2017, **56**, 5087–5090.
- 15 S. Bamberger, D. Hellwinkel and F. A. Neugebauer, *Chem. Ber.*, 1975, **108**, 2416–2421.
- 16 D. Hellwinkel and M. Melan, *Chem. Ber.*, 1971, **104**, 1001–1016.
- 17 D. Hellwinkel and M. Melan, *Chem. Ber.*, 1974, **107**, 616–626.
- 18 D. Hellwinkel, A. Wiel, G. Sattler and B. Nuber, *Angew. Chem.*, 1990, **102**, 677–680.
- 19 M. Yamamura, T. Saito and T. Nabeshima, *J. Am. Chem. Soc.*, 2014, **136**, 14299–14306.
- 20 M. Yamamura, K. Sukegawa and T. Nabeshima, *Chem. Commun.*, 2015, **51**, 12080–12083.
- 21 F. C. Krebs, P. S. Larsen, J. Larsen, C. S. Jacobsen, C. Boutton and N. Thorup, *J. Am. Chem. Soc.*, 1997, **119**, 1208–1216.
- 22 M. Yamamura, T. Hasegawa and T. Nabeshima, *Org. Lett.*, 2016, **18**, 816–819.
- 23 N. Hammer, T. A. Schaub, U. Meinhardt and M. Kivala, *Chem. Rec.*, 2015, **15**, 1119–1131.
- 24 M. Kivala, W. Pisula, S. Wang, A. Mavrinskiy, J.-P. Gisselbrecht, X. Feng and K. Müllen, *Chem. – Eur. J.*, 2013, **19**, 8117–8128.
- 25 F. Schlütter, F. Rossel, M. Kivala, V. Enkelmann, J.-P. Gisselbrecht, P. Ruffieux, R. Fasel and K. Müllen, *J. Am. Chem. Soc.*, 2013, **135**, 4550–4557.
- 26 M. Kuratsu, M. Kozaki and K. Okada, *Angew. Chem., Int. Ed.*, 2005, **44**, 4056–4058.
- 27 M. Hirai, N. Tanaka, M. Sakai and S. Yamaguchi, *Chem. Rev.*, 2019, **119**, 8291–8331.
- 28 J. E. Field and D. Venkataraman, *Chem. Mater.*, 2002, **14**, 962–964.
- 29 Z. Fang, T.-L. Teo, L. Cai, Y.-H. Lai, A. Samoc and M. Samoc, *Org. Lett.*, 2009, **11**, 1–4.
- 30 T. Kushida and S. Yamaguchi, *Organometallics*, 2013, **32**, 6654–6657.
- 31 A. Shuto, T. Kushida, T. Fukushima, H. Kaji and S. Yamaguchi, *Org. Lett.*, 2013, **15**, 6234–6237.
- 32 T. Kushida, A. Shuto, M. Yoshio, T. Kato and S. Yamaguchi, *Angew. Chem., Int. Ed.*, 2015, **54**, 6922–6925.
- 33 Z. Zhou, A. Wakamiya, T. Kushida and S. Yamaguchi, *J. Am. Chem. Soc.*, 2012, **134**, 4529–4532.
- 34 Y. Kitamoto, T. Suzuki, Y. Miyata, H. Kita, K. Funaki and S. Oi, *Chem. Commun.*, 2016, **52**, 7098–7101.
- 35 G. Turkoglu, M. E. Cinar and T. Ozturk, *Molecules*, 2017, **22**, 1522.
- 36 C. D. Entwistle and T. B. Marder, *Chem. Mater.*, 2004, **16**, 4574–4585.
- 37 F. Jäkle, *Chem. Rev.*, 2010, **110**, 3985–4022.

- 38 C. A. Hunter and J. K. M. Sanders, *J. Am. Chem. Soc.*, 1990, **112**, 5525–5534.
- 39 Y. Haketa and H. Maeda, *Chem. Commun.*, 2017, **53**, 2894–2909.
- 40 F. C. Krebs, B. W. Laursen, I. Johannsen, A. Faldt, K. Bechgaard, C. S. Jacobsen, N. Thorup and K. Boubekeur, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1999, **55**, 410–423.
- 41 J. B. Simonsen, K. Kjær, P. Howes, K. Nørgaard, T. Bjørnholm, N. Harrit and B. W. Laursen, *Langmuir*, 2009, **25**, 3584–3592.
- 42 J. B. Simonsen, F. Westerlund, D. W. Breiby, N. Harrit and B. W. Laursen, *Langmuir*, 2011, **27**, 792–799.
- 43 T. J. Sørensen, C. B. Hildebrandt, J. Elm, J. W. Andreasen, A. Ø. Madsen, F. Westerlund and B. W. Laursen, *J. Mater. Chem.*, 2012, **22**, 4797–4805.
- 44 T. J. Sørensen, C. B. Hildebrandt, M. Glyvradal and B. W. Laursen, *Dyes Pigm.*, 2013, **98**, 297–303.
- 45 P. Hammershøj, T. J. Sørensen, B.-H. Han and B. W. Laursen, *J. Org. Chem.*, 2012, **77**, 5606–5612.
- 46 B. Dong and H. Maeda, *Chem. Commun.*, 2013, **49**, 4085–4099.
- 47 A. Kütt, T. Rodima, J. Saame, E. Raamat, V. Mäemets, I. Kaljurand, I. A. Koppel, R. Y. Garlyauskayte, Y. L. Yagupolskii, L. M. Yagupolskii, E. Bernhardt, H. Willner and I. Leito, *J. Org. Chem.*, 2011, **76**, 391–395.
- 48 Y. Bando, Y. Haketa, T. Sakurai, W. Matsuda, S. Seki, H. Takaya and H. Maeda, *Chem. – Eur. J.*, 2016, **22**, 7843–7850.
- 49 Y. Haketa, S. Sasaki, N. Ohta, H. Masunaga, H. Ogawa, N. Mizuno, F. Araoka, H. Takezoe and H. Maeda, *Angew. Chem., Int. Ed.*, 2010, **49**, 10079–10083.
- 50 Y. Haketa, D. Katayama, S. Fukunaga, Y. Bando, T. Sakurai, W. Matsuda, S. Seki and H. Maeda, *Chem. – Asian J.*, 2016, **11**, 2025–2029.
- 51 Y. Haketa, Y. Honsho, S. Seki and H. Maeda, *Chem. – Eur. J.*, 2012, **18**, 7016–7020.
- 52 R. Yamakado, T. Sakurai, W. Matsuda, S. Seki, N. Yasuda, S. Akine and H. Maeda, *Chem. – Eur. J.*, 2016, **22**, 626–638.
- 53 Y. Terashima, T. Sakurai, Y. Bando, S. Seki and H. Maeda, *Chem. Mater.*, 2013, **25**, 2656–2662.
- 54 Y. Bando, S. Sakamoto, I. Yamada, Y. Haketa and H. Maeda, *Chem. Commun.*, 2012, **48**, 2301–2303.
- 55 Y. Bando, T. Sakurai, S. Seki and H. Maeda, *Chem. – Asian J.*, 2013, **8**, 2088–2095.
- 56 S. Sugiura, W. Matsuda, W. Zhang, S. Seki, N. Yasuda and H. Maeda, *J. Org. Chem.*, 2019, 8886–8898, DOI: 10.1021/acs.joc.9b00754.
- 57 S. Lee, C.-H. Chen and A. H. Flood, *Nat. Chem.*, 2013, **5**, 704.
- 58 B. Qiao, B. E. Hirsch, S. Lee, M. Pink, C.-H. Chen, B. W. Laursen and A. H. Flood, *J. Am. Chem. Soc.*, 2017, **139**, 6226–6233.
- 59 H. Zhang, S. Wang, Y. Li, B. Zhang, C. Du, X. Wan and Y. Chen, *Tetrahedron*, 2009, **65**, 4455–4463.
- 60 H. Zhang, Y. Li, X. Wan and Y. Chen, *Chem. Phys. Lett.*, 2009, **479**, 117–119.
- 61 S. Wang, M. Kivala, I. Lieberwirth, K. Kirchhoff, X. Feng, W. Pisula and K. Müllen, *ChemPhysChem*, 2011, **12**, 1648–1651.
- 62 A. T. Haedler, S. R. Beyer, N. Hammer, R. Hildner, M. Kivala, J. Köhler and H.-W. Schmidt, *Chem. – Eur. J.*, 2014, **20**, 11708–11718.
- 63 A. T. Haedler, K. Kreger, A. Issac, B. Wittmann, M. Kivala, N. Hammer, J. Köhler, H.-W. Schmidt and R. Hildner, *Nature*, 2015, **523**, 196.
- 64 A. T. Haedler, S. C. J. Meskers, R. H. Zha, M. Kivala, H.-W. Schmidt and E. W. Meijer, *J. Am. Chem. Soc.*, 2016, **138**, 10539–10545.
- 65 T. F. A. De Greef, M. M. J. Smulders, M. Wolffs, A. P. H. J. Schenning, R. P. Sijbesma and E. W. Meijer, *Chem. Rev.*, 2009, **109**, 5687–5754.
- 66 C.-M. Chou, S. Saito and S. Yamaguchi, *Org. Lett.*, 2014, **16**, 2868–2871.
- 67 N. Hammer, T. E. Shubina, J.-P. Gisselbrecht, F. Hampel and M. Kivala, *J. Org. Chem.*, 2015, **80**, 2418–2424.
- 68 Y. Dorca, J. Cerdá, J. Aragón, E. Ortí and L. Sánchez, *Chem. Mater.*, 2019, **31**, 7024–7032.
- 69 M. Bieri, S. Blankenburg, M. Kivala, C. A. Pignedoli, P. Ruffieux, K. Müllen and R. Fasel, *Chem. Commun.*, 2011, **47**, 10239–10241.
- 70 C. Steiner, J. Gebhardt, M. Ammon, Z. Yang, A. Heidenreich, N. Hammer, A. Görling, M. Kivala and S. Maier, *Nat. Commun.*, 2017, **8**, 14765.
- 71 H. Liang, Y. He, Y. Ye, X. Xu, F. Cheng, W. Sun, X. Shao, Y. Wang, J. Li and K. Wu, *Coord. Chem. Rev.*, 2009, **253**, 2959–2979.
- 72 P. A. Wood, S. J. Borwick, D. J. Watkin, W. D. S. Motherwell and F. H. Allen, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2008, **64**, 393–396.
- 73 S. Klyatskaya, F. Klappenberger, U. Schlickum, D. Kühne, M. Marschall, J. Reichert, R. Decker, W. Krenner, G. Zoppellaro, H. Brune, J. V. Barth and M. Ruben, *Adv. Funct. Mater.*, 2011, **21**, 1230–1240.
- 74 U. Schlickum, R. Decker, F. Klappenberger, G. Zoppellaro, S. Klyatskaya, M. Ruben, I. Silanes, A. Arnau, K. Kern, H. Brune and J. V. Barth, *Nano Lett.*, 2007, **7**, 3813–3817.
- 75 S. Gottardi, K. Müller, J. C. Moreno-López, H. Yildirim, U. Meinhardt, M. Kivala, A. Kara and M. Stöhr, *Adv. Mater. Interfaces*, 2014, **1**, 1300025.
- 76 K. Müller, J. C. Moreno-López, S. Gottardi, U. Meinhardt, H. Yildirim, A. Kara, M. Kivala and M. Stöhr, *Chem. – Eur. J.*, 2016, **22**, 581–589.
- 77 E. Kan, W. Hu, C. Xiao, R. Lu, K. Deng, J. Yang and H. Su, *J. Am. Chem. Soc.*, 2012, **134**, 5718–5721.
- 78 Y.-F. Han, Y.-X. Yuan and H.-B. Wang, *Molecules*, 2017, **22**, 266.
- 79 C. Steiner, Z. Yang, B. D. Gliemann, U. Meinhardt, M. Gurrath, M. Ammon, B. Meyer, M. Kivala and S. Maier, *Chem. Commun.*, 2018, **54**, 11554–11557.
- 80 B. Cordero, V. Gómez, A. E. Platero-Prats, M. Revés, J. Echeverría, E. Cremades, F. Barragán and S. Alvarez, *Dalton Trans.*, 2008, 2832–2838, DOI: 10.1039/B801115J.
- 81 R. D. Baechler and K. Mislow, *J. Am. Chem. Soc.*, 1970, **92**, 3090–3093.
- 82 I. V. Alabugin, S. Bresch and M. Manoharan, *J. Phys. Chem. A*, 2014, **118**, 3663–3677.
- 83 G. K. H. Madsen, F. C. Krebs, B. Lebech and F. K. Larsen, *Chem. – Eur. J.*, 2000, **6**, 1797–1804.

- 84 B. L. Feringa, R. A. van Delden, N. Koumura and E. M. Geertsema, *Chem. Rev.*, 2000, **100**, 1789–1816.
- 85 M. Saito, H. Shinokubo and H. Sakurai, *Mater. Chem. Front.*, 2018, **2**, 635–661.
- 86 T. Kawase, in *Supramolecular Chemistry of Fullerenes and Carbon Nanotubes*, ed. N. Martín and J.-F. Nierengarten, VCH-Wiley, Weinheim, 2012, pp. 55–78.
- 87 L. T. Scott, M. M. Hashemi and M. S. Bratcher, *J. Am. Chem. Soc.*, 1992, **114**, 1920–1921.
- 88 H. Sakurai, T. Daiko and T. Hirao, *Science*, 2003, **301**, 1878.
- 89 T. Hirao, M. Tosaka, S. Yamago and T. Haino, *Chem. – Eur. J.*, 2014, **20**, 16138–16146.
- 90 P. Metrangolo, F. Meyer, T. Pilati, G. Resnati and G. Terraneo, *Angew. Chem., Int. Ed.*, 2008, **47**, 6114–6127.
- 91 G. Cavallo, P. Metrangolo, R. Milani, T. Pilati, A. Priimagi, G. Resnati and G. Terraneo, *Chem. Rev.*, 2016, **116**, 2478–2601.
- 92 P. Politzer, J. S. Murray and T. Clark, *Phys. Chem. Chem. Phys.*, 2010, **12**, 7748–7757.
- 93 T. Clark, M. Hennemann, J. S. Murray and P. Politzer, *J. Mol. Model.*, 2007, **13**, 291–296.
- 94 T. A. Schaub, R. Sure, F. Hampel, S. Grimme and M. Kivala, *Chem. – Eur. J.*, 2017, **23**, 5687–5691.
- 95 J. Hamacek, C. Besnard, N. Mehanna and J. Lacour, *Dalton Trans.*, 2012, **41**, 6777–6782.
- 96 R. Thakuria, N. K. Nath and B. K. Saha, *Cryst. Growth Des.*, 2019, **19**, 523–528.
- 97 R. Sakamoto, K. Takada, T. Pal, H. Maeda, T. Kambe and H. Nishihara, *Chem. Commun.*, 2017, **53**, 5781–5801.
- 98 M. C. So, G. P. Wiederrecht, J. E. Mondloch, J. T. Hupp and O. K. Farha, *Chem. Commun.*, 2015, **51**, 3501–3510.
- 99 K. Meyer, M. Ranocchiari and J. A. van Bokhoven, *Energy Environ. Sci.*, 2015, **8**, 1923–1937.