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Supramolecular aggregates as sensory ensembles

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As a new emerging area in chemical sensing, sensing using supramolecular aggregates exhibits unique advantages over that using conventional small-molecule chemical sensors, in terms of high sensitivity and selectivity, and the simplicity of the sensory building blocks. This Feature Article outlines the recent research progress made in sensing based on induced supramolecular aggregation–disaggregation. The reviewed sensory building blocks, in general, in the form of a small molecular sensor, yet with a much simpler structure, which form aggregates, are those of perylene derivatives, pyrene derivatives, tetraphenylethylene derivatives, metallophilic species and metal–organic frameworks.

Introduction

A molecular chemosensor in general consists of a molecular recognition site and a signal reporting group, which are either directly connected or linked by a spacer.¹ In order to achieve high sensing performance, for instance, high sensitivity and selectivity, the sensory molecule needs to be sophisticatedly designed. Inspired by the chelating effect observed in coordination chemistry, such as that observed with metal–EDTA complexes

in which four amino acid binding sites are covalently linked within the same EDTA framework, and the specific bio-recognition which relies on multiple weak and less selective interactions operating in a cooperative manner, sensing using aggregates that hold a more simply designed chemosensor(s) by noncovalent interactions was developed.² Here the chemosensor-like building block itself may exhibit less effective performance than the otherwise molecular chemosensor would have done, yet in its aggregated state a comparable or better performance can be achieved. Supramolecular chemistry, landmarked by the 1987 Nobel Prize of chemistry awarded to Lehn, Cram and Pedersen,³ dealing with a complex of molecules that are held together by noncovalent bonds, has led to the buildup of diverse supramolecular architectures in a bottom-up approach. In a sense of employing noncovalent interactions, instead of

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Fig. 1 Comparison of (a) conventional sensory systems and (b) supramolecular sensory systems based on analyte induced supramolecular aggregation or disaggregation strategies.

covalent bonding, multidentate chelating ligands or enzymes are built in supramolecular chemistry by using one building block or a set of building blocks to form aggregates of them with the hope of achieving the chelating effect and/or the specificity in bio-recognition important for sensing.

Indeed, supramolecular aggregation, driven by weak non-covalent interactions, such as van der Waals, π - π stacking, hydrogen/halogen bonding, metal coordination, electrostatic and even metallophilic interactions,⁴ has been utilized to construct sensing platforms with enhanced selectivity and sensitivity. Supramolecular aggregates as sensing ensembles are advantageous over the conventional small-molecule chemical sensors in the following aspects (Fig. 1). (i) The building blocks are assembled by means of relatively weak and dynamic non-covalent interactions, providing easy and facile approaches for manipulating supramolecular structures. (ii) The supramolecular sensing system can be easily made water compatible by introducing water-soluble substituents into the building blocks, which provides enormous possibilities for such recognition systems to be applied for biological analytes in the aqueous phase.⁵ (iii) The higher local concentration of the binding sites and the lower local concentration of water molecules around the binding sites in the aggregates substantially enhance the originally weak and water-susceptible interactions, which makes those otherwise insufficient binding events in water of the corresponding molecular sensors highly sensitive. Since in many cases the building block molecules are aromatic compounds with large π -systems, strong π - π stacking in the aggregates would make them conjugate-polymer like, affording an additional possibility for signal amplification.⁶ The relative orientations among the binding sites in the aggregates can be tuned good for analyte binding,⁷ resulting in high selectivity. (iv) In particular, the aggregation or disaggregation process

may involve cooperative interactions that also result in signal amplification, leading to higher sensitivity and selectivity as well. Therefore, using supramolecular aggregates from building blocks with easy design and facile syntheses could achieve even better sensing performance than that of molecular sensory systems that rely on sensors with delicate structures.

A variety of supramolecular sensing systems have been developed. Yet, regarding analyte induced aggregation or disaggregation strategies, considerable progress has been made only with limited categories of building blocks, which operate for sensing *via* π - π stacking modulated photophysics of π -conjugated dyes, aggregation induced emission, metallophilic interactions, and more recently those emerging from highly ordered metal-organic frameworks. We highlighted in 2012 supramolecular aggregation/disaggregation-based molecular sensing in China.⁸ With the rapid development of supramolecular sensing, it becomes necessary to update its current state of the art. In this Feature Article we accordingly review recent developments using supramolecular sensory ensembles consisting of the following five representative building blocks, *i.e.* perylene derivatives, pyrene derivatives, tetraphenylethylene derivatives, metallophilic species, and metal-organic frameworks (MOFs).

PBI derivatives

Derivatives of perylene dyes are extensively employed as molecular and supramolecular probes because of their strong absorptivity in the mid-visible spectral region, and thermal and photochemical stability.⁹ In particular, perylene-3,4:9,10-bis(dicarboximides) (PBIs) have attracted much attention as optimal fluorescent dyes and been intensively utilized in fluorescent switches, molecular transistors and sensors.^{10,11} They have been considered to be excellent fluorophores in sensor design because of their excellent electron accepting ability and strong fluorescence in the monomeric and small oligomeric states.^{12,13} Furthermore, the PBI structure and the planar π -electron deficient aromatic nature are known to strongly promote the formation of aggregates through stacking interactions between the π -conjugated cores in a face-to-face fashion, forming H-aggregates, which result in fluorescence quenching and a hypsochromic shift of the absorption.^{14,15} Thus, great efforts have been made to develop PBI based supramolecular sensing systems, by equipping a binding site into the PBI structures. Easy and versatile derivatization makes the PBI framework more attractive.

Jiang's group in 2010 reported the Hg^{2+} selective PBI derivative **1** which bears an imide group similar to that in thymine (T) known to form the so-called "T- Hg^{2+} -T" binding motif. Thus, Hg^{2+} was shown to be able to specifically induce aggregation of **1**, leading to a dramatic fluorescence quenching of **1** and allowing for a highly sensitive and selective sensing of Hg^{2+} . More importantly, it afforded a nonfluorescent chemosensing ensemble "(1- Hg)_n" for thiol-containing species (Fig. 2).¹⁶ Addition of thiol-containing amino acids to the **1**- Hg^{2+} ensemble





Fig. 2 Hg^{2+} induced aggregation of **1** and dissociation of aggregates in the presence of cysteine. Reprinted with permission from *Chem. Commun.*, 2010, **46**, 4938–4940. Copyright 2010 RSC.

solution induced dissociation of the aggregates and recovery of the fluorescence. For cysteine, a detection limit of 9.6 nM was achieved. It therefore represents a sensory system that works for both Hg^{2+} and thiol-containing species. Since both the aggregation and disaggregation processes are cooperative, high sensitivity and selectivity were observed, despite the otherwise weak interaction of Hg^{2+} with a simple imide group. This is the first report that shows that an imide group other than that from thymine could lead to a similar “T- Hg^{2+} -T” binding motif for sensing applications.

An L-DOPA functionalized amphiphilic PBI **2** was reported in 2014 by Govindaraju *et al.*,¹⁷ as a reversible fluorescence switching probe for the detection and sensing of cationic surfactants and $\text{Fe}^{3+}/\text{Cu}^{2+}$ in aqueous solutions, following, respectively, the host-guest interactions driven aggregation and disaggregation of **2**. Photophysical studies of **2** showed that the compound existed as a monomer in dimethyl sulfoxide (State-I), while in pure aqueous solutions in the form of self-assembled aggregates, displaying very weak fluorescence along with red-shifted broad absorption (State-II). Interestingly, the cationic surfactant cetyltrimethylammonium bromide (CTAB), but not neutral and anionic surfactants, could disassemble **2** under micellar conditions, restoring the bright yellow fluorescence and vibronically well-defined absorption bands of **2** (State-III). **2** was able to sense Fe^{3+} and Cu^{2+} owing to the metal chelating capacity of its DOPA moiety, so that metal binding drove the assembly of **2** (State-IV). Fluorescence of **2** was quenched by Fe^{3+} and Cu^{2+} by 3.5 and 2.5-fold, respectively. Such metallosupramolecular aggregates were found to revert back to the fluorescence “ON” state by the introduction of a metal ion chelator, diethylene triamine pentaacetic acid (DTPA, State-III), further signifying the role of metal ions toward the assembly of **2** (Fig. 3).

Würthner *et al.*¹⁸ reported an interesting metal templated assembly of crown ether functionalized PBIs **3a** and **3b**. The templated self-assembly of these PBIs was shown to be dependent on the size of the crown ether receptor and the nature of metal ions. Ba^{2+} selectively templates the assembly of 15-crown-5 functionalized PBI **3a** into a dimeric PBI- Ba^{2+} complex through a positive homotropic cooperative binding, so that the fluorescence of **3a** is efficiently and selectively

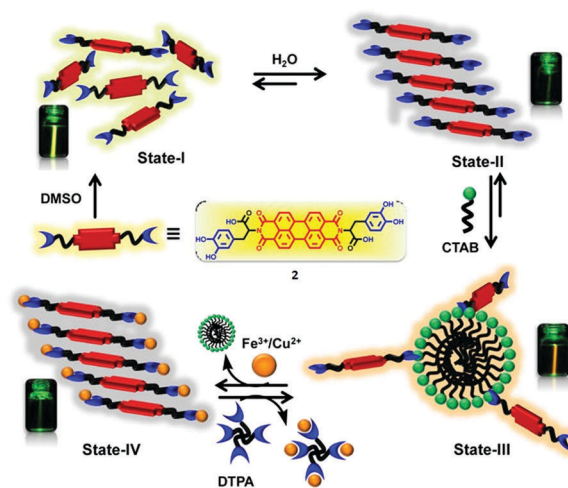


Fig. 3 Schematic representation of solvent driven molecular organization of **2** and transformation of aggregated State-II of **2** to molecularly dissolved State-III in the presence of CTAB micelles, and the transformation of State-III into State-IV via metal ion induced reorganization of **2** in the presence of CTAB micelles. Reprinted with permission from *ACS Appl. Mater. Interfaces.*, 2014, **6**, 21369–21379. Copyright 2014 ACS.

quenched by Ba^{2+} , whereas it is not quenched or quenched very little by the other alkali and alkaline earth metal ions such as Na^+ , K^+ , Rb^+ , Mg^{2+} and Ca^{2+} . These results show the high selectivity of Ba^{2+} templating of the assembly of PBI, giving a binding constant of $(1.51 \pm 0.04) \times 10^5 \text{ M}^{-1}$ for Ba^{2+} , despite the fact that 15-crown-5 is Na^+ selective. In contrast, both absorption and emission of the 18-crown-6 functionalized PBI **3b** were not influenced by those metal ions of up to 7 equivalents. This observation suggests the vital importance of the size of the crown ether receptor in the metal ion templated assembly of the PBI derivatives, actually implying the subtle balance of multiple interactions involved in the templating and advantages of the supramolecular sensing.



The essential role of carbohydrate-protein interactions in maintaining diverse biological processes has attracted much



attention to develop carbohydrate-based sensors for these interactions. Among them, fluorescent glycoclusters hold great potential because of their intrinsic fluorescence, high sensitivity to minor external stimuli and good biocompatibility. A water-soluble mannose functionalized PBI **4**, developed by Wang *et al.*,¹⁹ exhibited a specific response towards Concanavalin A (Con A) by *ca.* 3 times fluorescence enhancement, resulting from disaggregation of **4** in the presence of Con A. Whereas peanut agglutinin (PNA) and bovine serum albumin (BSA) led to no significant changes in the fluorescence of **4**. The binding constant of **4** with Con A, $(9.87 \pm 0.3) \times 10^6 \text{ M}^{-1}$, is three orders of magnitude higher than that ($8.2 \times 10^3 \text{ M}^{-1}$) with the monosaccharide ligand in Con A. Fluorescence imaging of **4** for macrophage cells also demonstrated that the sensor could be applied as a protein-specific probe for diagnosis and molecular imaging under live cell conditions.



Amphiphilic PBI dyes **5** and **6** functionalized with *para*- and *ortho*-phenylboronic acid moieties, for interacting with 1,2-/1,3-*cis*-diol to form a cyclic boronate ester, were designed as chirality sensors for α -hydroxycarboxylates.²⁰ The achiral compounds **5** and **6** existed as dimers or small oligomers in the absence of an α -hydroxy carboxylate, yet formed large aggregates upon binding it, resulting in a decrease in the absorption of the perylene chromophore around 500 nm. Strong induced Cotton effects were observed in the absorption window of the achiral perylene chromophore and were utilized for enantiomeric excess (ee) determination and discrimination of seven α -hydroxy carboxylates that bear no chromophore, with an average absolute error of 2% in ee determination and 100% correctness in analyte identification. Note that the boronic acid group that interacts with the chiral analyte is located far away from the chromophore, yet an effective chirality induction was shown by the coupled exciton CD signals from the latter. This example highlights the potential of using induced helical aggregation of achiral dyes for CD measurement of the ee of those chiral analytes bearing no chromophore. In principle it should also work for a chiral species that bears a chromophore but is far away from the chiral centre. This expands the scope of applications in chirality sensing of classic CD spectroscopy that normally requires the presence of a chromophore next to the chiral centre for direct measurements. It is the aggregation of

the achiral building blocks upon interacting with the chiral species that leads to the transformation of the molecular chirality into supramolecular structures.



Molecular sensors for analytes with multiple functional groups are normally designed to have several binding sites that are well arranged to match the structures of the analytes, to achieve high sensitivity and selectivity. In terms of the supramolecular sensory ensemble, a set of monofunctional building blocks may form co-aggregates that thereby bear multiple binding sites as well and thus function similarly to the molecular sensors that contain a set of well-defined binding sites, yet greatly simplify the structure and syntheses of the building blocks. It should be emphasized that achieving a synergistic effect to overcome the entropic cost of forming such multi-component co-aggregates is challenging. A successful example for sensing L-DOPA (**7**) was recently reported, employing the induced aggregation of a mixture of boronic acid and aldehyde functionalized PBIs **5** and **8**.²¹ Here the boronic acid group in **5** was expected to interact with the *cis*-diol moiety in L-DOPA, while the aldehyde group in **8** was expected to form an imine bond with the amine group in L-DOPA, both being reversible and dynamic. The three-component **5**-L-DOPA-**8** complex formed at low concentrations in aqueous solutions underwent aggregation, exhibiting a strong synergistic effect between the two orthogonal dynamic covalent interactions. The CD-based sensing of **7** at a micromolar concentration level can be carried out by simply mixing with two simple building blocks **5** and **8**. The high affinity towards **7** results from the stabilization by aromatic π -stacking between **5** and **8**. The divalent binding of **7** to the **5**-**8** ensembles results in the formation of long fibrous PBI aggregates with a significantly larger size and stronger induced CD signals compared to the aggregates formed, respectively, by the **7**-**5** and **7**-**8** complexes in which only monovalent binding of **7** is possible. This report confirms that using a mixture of building blocks bearing different binding groups could indeed operate, at least to the same extent, as the molecular sensor that bears all those binding groups, since these much more easily designed and facily synthesized building blocks may function cooperatively. This is important since many analyte targets may have more than one functional group so that the molecular sensors could be structurally complicated to ensure high binding efficiency.





Wang *et al.*²² proposed a label free ultrasensitive fluorescence turn-on sensing scheme for DNA, using the cationic polyelectrolytes-induced aggregation of the negatively charged PBI dye **9** that is shown by the severe quenching of the emission of **9**. The tested four cationic polymers **10–13** could induce almost complete quenching of the fluorescence of **9**, but with varying efficiency as indicated by the concentrations of the polymers required to fully quench the emission, 300, 1150, 175, 50 nM, respectively. When a single stranded DNA (a polyanion) was added to the solution of the cationic polymer–**9** aggregates, strong electrostatic interactions between the single stranded DNA and the cationic polymers released **9** from the aggregates, recovering the emission of **9**. Aggregates **12–9** exhibited the best fluorescence recovery up to 89.8%, by only 1.5–2 nM DNA in *ca.* 5 minutes. This smart scheme allows a highly sensitive assay, with a limit of the detection of 2 pM for DNA. Based on the fact that alkaline phosphatase (ALP) could remove the phosphate group of the 5'-phosphorylated single-stranded DNA, thus preventing the DNA sample from being degraded by λ -exonuclease, an assay for the highly sensitive detection of ALP activity was established using the **12–9** supramolecular sensory ensemble. Upon mixing with the DNA samples, the fluorescence of the aggregates **12–9** was turned on, reaching a limit of detection down to 0.02 mU mL⁻¹. This represents the rare example of induced disaggregation of sensory building blocks, releasing them in an amplified manner that ensures high sensitivity.



Pyrene derivatives

As a simple aromatic fluorophore, pyrene has a large extinction coefficient, high quantum yield and good stability in aqueous

solutions. Pyrene is known to form an excimer, due to its long lifetime of 10² ns orders of magnitude and its strong tendency for $\pi \cdots \pi$ stacking, thereby emitting structureless excimer fluorescence red-shifted by *ca.* 100 nm from the monomer emission. Pyrene has thus been used for a long time to design building blocks for supramolecular aggregates that form in the presence of an analyte.

Yam's group²³ in 2009 developed a pyrene-based boronic acid fluorescence sensing ensemble for glucose, in which (i) the water-soluble boronic acid functionalized polymer **14** transformed into a polyanion upon binding with glucose and (ii) the positively charged pyrene derivative **15** electrostatically attached along the polymer backbone, in the aggregated form as indicated by the strong excimer emission at 492 nm. A linear response was found for glucose between 0 and 17 mM. Among a number of tested sugars, glucose produced the strongest excimer emission, which is likely due to the formation of 1:2 glucose–boron acid complexes that bring two pendant boronic acid moieties into close proximity, thus generating two adjacent negative charges which favour the aggregation of positively charged **15**.



An aggregation-based glucose sensor **16** developed by Jiang's group clearly manifests the multivalent effect, with higher apparent affinity to glucose than to fructose, in such a simple phenylboronic acid derivative containing a pyrene fluorophore (Fig. 4).²⁴ In an alkaline (pH 10) aqueous solution **16** at 0.1 mM concentration exists in small aggregates that emit only pyrene monomer fluorescence at 390 nm. Glucose binding leads to more ordered aggregates of **16**, since one glucose molecule



Fig. 4 Cartoon illustration of the aggregates formed, respectively, from the 1:1 fructose complex and the 1:2 glucose complex with **16**. Reprinted with permission from *J. Am. Chem. Soc.*, 2013, **135**, 1700–1703. Copyright 2013 ACS.





Fig. 5 Schematic representation of fluoride sensing by the **16**-PCA boronate ester **17**. Reprinted with permission from *Chem. Commun.*, 2014, **50**, 13987–13989. Copyright 2014 RSC.

binds two boronic acid groups, thus the resultant complex is more hydrophobic, enabling the observation of pyrene excimer emission. With fructose, 1:1 binding stoichiometry with the boronic acid group results in a neutral zwitterionic boronate ester of higher hydrophobicity, however, electrostatic repulsions between the zwitterionic boronates lead to disordered aggregates of **16**, unfavourable for excimer formation. Despite the simple phenylboronic acid structure, **16** in its aggregate state exhibits a highly selective and sensitive response toward glucose against fructose, opposite to what is known for phenylboronic acid that shows preferential binding to fructose.

Sensing of the fluoride ion in aqueous solutions using a boronic acid based sensor has been challenging because of the high hydration enthalpy of the fluoride ion and the severe interference from the hydroxide anion. Jiang's group reported²⁵ a supramolecular amphiphilic pyrene-based boronic acid sensor for direct fluoride sensing in aqueous solutions (Fig. 5). Binding of the fluoride ion to the boronate ester of **16** and protocatechuic acid (PCA) triggered the formation of supramolecular aggregates of ester **17**, leading to presumably lower local concentration of water and higher local density of binding sites, which enhance the fluoride affinity and therefore the selectivity against OH^- . To enhance the Lewis acidity of boronic acid toward the fluoride ion and reduce the interference of OH^- , PCA was chosen to interact with **16** to form ester **17** which aggregates under highly acidic conditions ($\text{pH} \leq 2$), ruling out the interference of OH^- . The excimer emission of **16** undergoes a dramatic enhancement upon addition of fluoride ions, together with an increase in solution turbidity, indicative of supramolecular aggregation, allowing sensitive detection of fluoride at the ppm level in aqueous solutions. The fact that the addition of CTAB micelles leads to a dramatic quenching of the excimer emission supports the conclusion that aggregation strengthens the boron-fluoride interaction in aqueous solutions. In the presence of 10 mM PCA, an apparent binding constant for fluoride at 10^3 M^{-1} orders of magnitude was obtained, a high value unprecedented for boronic acid sensors in water.

Tetraphenylethylene derivatives

The concept of aggregation induced emission (AIE)²⁶ or aggregation induced enhanced emission (AIEE)²⁷ has been proposed

and well received as a new promising strategy for creating potential fluorescence sensors. Compared with the typical aggregation-caused quenching (ACQ) effect which is unfavourable for chemical sensing in terms of sensitivity, AIE holds great potential in expanding the applications of traditional organic luminophores into new “turn-on” luminescence chemo-/bio-sensors. Tang's group²⁸ was the first to report the AIE-active molecules, represented by the star molecule TPE, tetraphenylethylene. TPE is non-emissive in the dissolved state, whereas it turns highly emissive in the aggregated state. Tang *et al.*^{26,28,29} attributed the AIE to the block of the radiationless pathways by the restriction of intramolecular rotations (RIR) in the aggregated state.

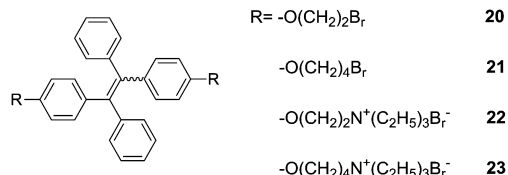
Yang *et al.*³⁰ prepared a TPE-based sensor **18** for DNA, employing the coordination of Zn^{2+} with the tridentate ligands substituted in the TPE core that results in a positively charged TPE derivative. Upon the addition of a 10 nt ssDNA (X10), a significant fluorescence enhancement in $\text{H}_2\text{O}/\text{DMSO}$ (99:1, v/v) was observed. The complementary sequence DNA of X10, Y10, and the hybridized dsDNA of X10/Y10 was also tested and fluorescence enhancement to a similar extent was shown for Y10 but slightly less for X10/Y10, despite the same emission maximum at 465 nm. The control TPE molecule **19**, interacts with DNA in an electrostatic manner; its fluorescence enhancement is in general weaker than that observed for **18**- Zn^{2+} , suggesting that the coordination of Zn^{2+} with ligands in **18** contributes cooperatively with the electrostatic interactions of the cationic complex that forms to the total interaction of **18** with DNA.



Tang *et al.*³¹ reported a set of AIE-active TPE derivatives **20–23**. Cationic salts **22** and **23** are soluble in water and in DMSO and DMF, as well; **22** and **23** were tested as sensors for protein and DNA, by means of hydrophobic and electrostatic interactions. In dilute solutions of **20** and **21** in acetonitrile (AN) they are practically nonfluorescent, whereas they become highly fluorescent in 1:99 (v/v) AN/water solution. The intensity of **20** at 476 nm is increased by 240-folds and its absorption shifted to red from 310 nm in AN to 330 nm in the suspension. Similar observations were made with **21**, therefore, both **20** and **21** are AIE-active. Because of the amphiphilic nature of **22** and **23**, the addition of organic solvents such as methanol, AN, THF and 1,4-dioxane into their aqueous solutions did not cause salting-out or aggregation of them. However, their fluorescence quantum yields increased with increasing concentration, suggesting that **22** and **23** are AIE-active too. Calf thymus (ct) DNA or bovine



serum albumin (BSA) were shown to enhance the fluorescence of **22** and **23** in aqueous phosphate buffer of pH 7.0 at 25 °C and red-shift its emission to 462 nm, which differs from its absorption maximum by 134 nm. **23** is more sensitive for the detection of DNA and BSA, for example, its fluorescence is enhanced by 16.3 and 23.8 times by 300 $\mu\text{g mL}^{-1}$ DNA and 500 $\mu\text{g mL}^{-1}$ BSA, respectively. These two systems of **18-Zn²⁺** and **22/23** can be similarly understood to that of **14/15** for the sensing of glucose.¹⁸

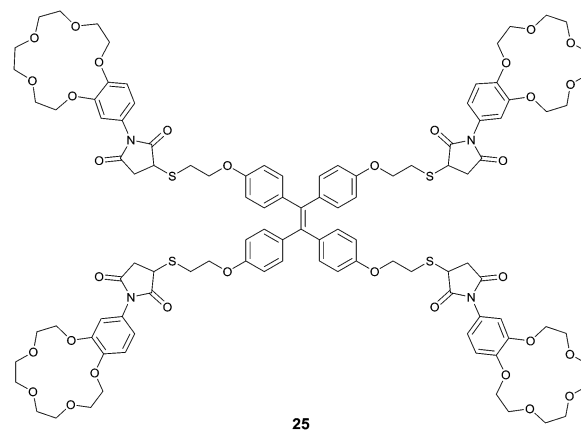


Hierarchical self-assembly of positively charged metallacycle **24** and negatively charged linear polysaccharides was observed by Yang *et al.*,³² in which they showed that multiple electrostatic interactions were driving forces. NMR, ESI-MS and morphology studies show that the elaborately designed tris-TPE metallacycle form entangled bead-like aggregates upon the threading of the linear heparin through the metallacycles, nicely supported by the enhanced emission of the TPE fluorophores that allow the detection of heparin at the clinical dosage level. The excellent selectivity for heparin over its analogues such as sodium polyacrylate and sodium poly(*p*-styrenesulfonate) is impressive, considering the multiple electrostatic interaction nature. The size matching appears to be a key factor which is amplified upon aggregation, explaining the observed selectivity for heparin.



Integrating host-guest molecular recognition and the AIE effect, Liu *et al.*³³ developed a highly sensitive and selective fluorescence sensor **25** for K⁺. As a crown-ether functionalized TPE derivative, the four pendant benzo-15-crown-5 moieties and the TPE core serve, respectively, as a K⁺ binding group and an AIE-active motif. **25** could dissolve well in THF, thus exhibiting faint fluorescence, whereas in aqueous THF solution, fluorescence of

25 was enhanced gradually with increasing water volume percentage until 70%, when a dramatic enhancement starts to happen because of the aggregation of **25**. Meanwhile, the emission of **25** at 460 nm in THF also experienced a dramatic enhancement by 9.5 fold when K⁺ was introduced at 0.0–19.2 μM concentrations. A detection limit of 1.0 μM was obtained for K⁺ in THF at a concentration of **25** of 4.8 μM , demonstrating the high sensitivity of the sensing. The binding of K⁺ with the peripheral crown ether moieties in **25**, forming K⁺/15-C-5 sandwich complexes, promoted the cross-linked networking of the TPE molecules and thus restricted the internal rotation, accounting for the observed fluorescence enhancement similar to that observed for **25** itself in water-rich THF solutions. Indeed, DLS data indicated the existence of the aggregate nanoparticles of size 200 nm when **25** was mixed with 1 equivalent of K⁺ ions in THF, and this was confirmed by the AFM observations. It also is worth pointing out that **25** is highly selective toward K⁺ in THF, since enhancement of the fluorescence of **25** was only observed by K⁺ among the tested metal cations of Li⁺, Na⁺, K⁺, NH₄⁺, Ca²⁺, Mg²⁺ and Pb²⁺, despite the fact that 15-C-5 binds Na⁺ better.



Attaching metal binding groups to the TPE core, Zhu's group³⁴ developed two fluorescence chemosensors **26** and **27** for Ag⁺ and Hg²⁺, respectively. Here adenine (A) and thymine (T) are the respective binding groups for Ag⁺ and Hg²⁺, following the well-established "A-Ag⁺-A" and "T-Hg²⁺-T" binding motifs. Enhancement of the fluorescence of **26** and **27** occurs in the presence of Ag⁺ and Hg²⁺ due to the induced aggregation that restricts the intramolecular rotation in the TPE core (Fig. 5). **26** itself showed rather faint emission in H₂O/THF (5 : 1, v/v), however, its emission at 470 nm increased linearly upon the addition of AgClO₄ that can be visualized by the naked eye. This was attributed to the coordination of Ag⁺ with **26** to form the oligomer- or polymer-like complexes (Fig. 6), rather than the intramolecular coordination complex of **26** and Ag⁺. Despite the weak and less selective binding of free adenine with Ag⁺, the response of fluorescence of **26** in aqueous THF solutions was highly sensitive and highly selective as well since only Ag⁺, and slightly Hg²⁺, could induce emission enhancement, among the tested metal ions Ba²⁺, Ca²⁺, Cd²⁺, Co²⁺, Cu²⁺, Fe³⁺, Fe²⁺, Hg²⁺, Mg²⁺, Mn²⁺, Ni²⁺, Pb²⁺, Zn²⁺, Ag⁺, Cs⁺, and K⁺. Similar observations were made with fluorescence sensing by **27** for Hg²⁺.





Fig. 6 Design rationale for Ag^+ and Hg^{2+} chemosensors **26** and **27**. Reprinted with permission from *Org. Lett.*, 2008, **10**, 4581–4584. Copyright 2008 ACS.



Ceroni and co-workers observed switchable phosphorescence upon complexation of Mg^{2+} to **28**.³⁵ Compound **28**, another kind of AIE molecule, is nonluminescent in fluid solution, whereas it exhibits red-shifted absorption and phosphorescence in the presence of Mg^{2+} . The assembled polymers display similar phosphorescence spectral profiles and lifetimes to those of their powder, suggesting that aggregation restricts the intramolecular rotations.³⁶ Subsequent addition of fluoride disassembles the aggregates because of the formation of MgF_2 and the vanishing of the phosphorescence. This process is reversible by adding Mg^{2+} and F^- alternately, making it a tunable luminescence sensor for both Mg^{2+} and F^- .



Tanaka *et al.*³⁷ developed an interesting TPE-based sensor for Cu^{2+} , by using an azide-modified TPE **29** that interacts with diethylene glycol dipropionate **30** *via* click chemistry. This reaction between **29** and **30** in 1:1 (v/v) $\text{H}_2\text{O}/\text{THF}$ forms the intermolecular cross-linked products (like aggregates) in the

presence of trace amounts of Cu^+ which is *in situ* generated by reduction of Cu^{2+} by sodium ascorbate. Thus the fabricated assay establishes a sensitive and selective sensing scheme for Cu^{2+} with a limit of detection of $1\ \mu\text{M}$, sensitive enough for the level of Cu^{2+} in drinking water according to US EPA standard. Sample solutions containing various other metal ions (Li^+ , Na^+ , K^+ , Mg^{2+} , Ca^{2+} , Mn^{2+} , Fe^{2+} , Fe^{3+} , Co^{2+} , Ag^+ and Zn^{2+}) and a mineral water sample containing Na^+ , K^+ , Mg^{2+} , and Ca^{2+} ions under the same treatment procedures produced no significant enhancement in the fluorescence, it only happened when Cu^{2+} ions were added. This opens a new avenue for employing the AIE phenomenon to achieve high selectivity, a character of chemodosimeters.³⁸



Employing the click reaction, He and co-workers developed a new AIE fluorescence chemosensor **31** for Fe^{3+} .³⁹ The TPE derivative **31** started to aggregate in $\text{H}_2\text{O}/\text{THF}$ binary solvents containing 60% water, in which strong emission was observed with two bands that peaked at 408 nm and 432 nm. **31** in $\text{H}_2\text{O}/\text{THF}$ (2:1, v/v) exhibited significant fluorescence quenching only in the presence of Fe^{3+} , whereas the tested other cations (Li^+ , Na^+ , K^+ , Mg^{2+} , Ca^{2+} , Ba^{2+} , Mn^{2+} , Ni^{2+} , Fe^{2+} , Pb^{2+} , Al^{3+} , Cr^{3+} , Hg^+ , Ce^{2+} and Cd^{2+}) did not. This was attributed to the coordination of Fe^{3+} to **31** that disfavors the aggregation of **31**. Job's plot and ^1H NMR titrations led to tentative assignment of the structure of the 2:1 $\text{Fe}^{3+}/\text{31}$ complex which is multi-charged and thus more hydrophilic and tends to aggregate less in aqueous solutions. Again the system is sensitive with a limit of detection of $0.7\ \mu\text{M}$.





Fig. 7 Rationale for the fluorescence turn-on detection of H_2O_2 using **32** which upon oxidative hydrolysis leads to the neutral TPE product **33** that undergoes aggregation. Reprinted with permission from *Tetrahedron Lett.*, 2014, **55**, 1471–1474. Copyright 2014 Elsevier.

Hydrogen peroxide is the least reactive and mild oxidant among reactive oxygen species (ROS), which are toxic to cells if the level of ROS exceeds the tolerable physiological range. Developing sensitive sensors to monitor the level of H_2O_2 both in living cells and in the environment has attracted intensive attention. Zhang and co-workers⁴⁰ designed a H_2O_2 -selective fluorescence sensor **32** which contains a TPE core and a H_2O_2 -reactive group, arylboronate ester, which are linked *via* a pyridinium bridge (Fig. 7). The pyridinium bridge makes **32** water-soluble, thus enabling it to exist in the monomer form and being nonfluorescent. Oxidative hydrolysis of **32** by H_2O_2 generates the much less water-soluble product **33**, which undergoes aggregation and therefore becomes highly fluorescent, allowing turn-on fluorescence sensing of H_2O_2 with high selectivity as a chemodosimeter would normally exhibit. Indeed, among the tested reactive oxygen species such as H_2O_2 , singlet oxygen, the hydroxyl radical, the superoxide anion and nitric oxide, enhancement of emission was only observed in the case of H_2O_2 . The assay is highly sensitive as well, with a limit of detection for H_2O_2 down to 180 nM. More importantly, since D-glucose can be oxidized in the presence of glucose oxidase to produce H_2O_2 , the scheme shown in Fig. 6 can be applied for D-glucose as well.

Metallophilic interactions

Two closed-shell metal cations (such as Au^+ , Ag^+ and Cu^+) tend to have attractive interactions rather than repulsion when their distance is smaller than the sum of the van der Waals radii, as is the so-called metallophilic interaction (metal \cdots metal interaction). Metal-coordinated complexes with metallophilic interactions exhibit unique photochemical and photophysical properties, making them useful in many aspects, such as functional nanomaterials,^{18,22,41} luminescent materials for ion recognition and sensing,^{21,42,43} and chiral nanoclusters.^{26,27,44} Our group has been interested in establishing a series of coordination polymers facilitated by argentophilic interactions, with the aim of achieving versatile sensors, choosing zigzag structured $(\text{Ag}^+\text{-SR})_n$ backbones.



Fig. 8 Formation of $\text{Ag}(\text{I})\text{-NCys}$ coordination polymers allowing for the ratiometric fluorescence sensing of Ag^+ . Reprinted with permission from *Chem. Commun.*, 2011, **47**, 5900–5902. Copyright 2011 RSC.

A ratiometric luminescence sensor with outstanding sensitivity and selectivity for Ag^+ was established by forming coordination polymers with the thiol ligand NCys, which is an *N*-(1-naphthyl-acetyl)cysteine derivative designed to bear an aromatic fluorophore (Fig. 8).⁴⁵ The *in situ* formed coordination polymers were assumed to be facilitated by the $\text{Ag}^+\cdots\text{Ag}^+$ interaction which is probed by a new absorption at 350 nm, assigned to ligand-to-metal charge transfer transition modified by the argentophilic interaction (LMMCT, $\text{S} \rightarrow \text{Ag}^+$). $\pi\cdots\pi$ stacking of the naphthyl moieties in the side chains may contribute additionally to the stabilization of the polymers. The naphthyl fluorophore was also designed to harvest light and emit fluorescence. Dual emission from the naphthyl and $\text{Ag}^+\cdots\text{Ag}^+$ fluorophores allows establishing a sensitive ratiometric assay for Ag^+ at the sub- μM level, in particular with high selectivity so that much more thiophilic Hg^{2+} exhibits little interference.

CD signals corresponding to $\text{Ag}^+\cdots\text{Ag}^+$ absorption were also observed and the signs of the CD signals are related to the chirality of the cysteine residue.^{42,45} A possible reason is that the $\text{Ag}^+\cdots\text{Ag}^+$ interaction and the sidechain/sidechain interactions in the coordination polymers form a network including the $\text{Ag}^+\cdots\text{Ag}^+$ chromophore and the chiral cysteine residues. We therefore extended the study to employ coordination polymers of Ag^+ with an achiral thiol ligand that bears a binding group for chiral species, for chiral sensing. A first attempt was made with a thiophenol that bears a boronic acid group for interacting with saccharides (Fig. 9).⁴² CD spectra of varying profiles were observed with the tested saccharides such as glucose and fructose, confirming that the chirality of the saccharide is transferred into the $\text{Ag}^+\cdots\text{Ag}^+$ chromophore in the supramolecular sensory ensembles upon interacting with the boronic acid group to form the cyclic boronate ester. Glucose that interacts with the supramolecular sensor in a multivalent manner results in stronger CD signals whereas the monovalent fructose induced practically no or very weak CD signals. The CD spectral profile could be acquired to correlate to the saccharide identity. The observation made here is of relevance for understanding signal transduction in biological systems. Applying coordination polymers featuring metal \cdots metal interactions remains at its early stage and it deserves to be explored in a broad scope by designing new sensory ligands and employing other metal \cdots metal interactions.





Fig. 9 Interactions of glucose with Ag^+ -MPBA coordination polymers. Reprinted with permission from *Chem. Commun.*, 2015, **51**, 8017–8019. Copyright 2015 RSC.

Metal–organic frameworks

Metal–organic frameworks (MOFs) with well-defined pore-structures have evolved into functional materials for many potential applications.⁴⁶ Yet because of their poor stability in biological media, few of them have been applied to biosensing. Chen's group⁴⁷ is among the first to construct a water-stable Cu^{2+} -based zwitterionic carboxylate three-dimensional MOF (**34**, Fig. 10), for the detection of DNA and RNA. This elaborately fabricated MOF affords multiple interactions with probe DNAs attached with the fluorophore FAM (P-DNAs), forming two P-DNA@**34** systems. Fluorescence of the P-DNA is efficiently quenched by Cu^{2+} , presumably *via* a photo-induced electron-transfer (PET) mechanism. Upon introducing the human immunodeficiency virus (HIV) double-stranded DNA (ds-DNA) sequence (T_0) or Sudan virus (SUDV) RNA sequence (T_0'), relevant P-DNA gets rid of the MOF and hybridizes with the complementary sequence, resulting in fluorescence recovery up to 91% and ensuring high sensitivity with detection limits of 196 pM for T_0 and 73 pM for T_0' . In addition, the detection is hardly interfered by analogous DNA or RNA sequences with one base pair mismatching, demonstrating its high selectivity. The reasons for the excellent sensing performance remain undisclosed, while the size of the pores in MOF was proposed to be a possibility.

Wang *et al.*⁴⁴ carried out a noteworthy conceptual MOF-based “turn-on” fluorescence sensing using encapsulated AIE fluorophores for five-membered-ring energetic heterocyclic compounds (5MR-HCs, Fig. 11). 5MR-HCs are a kind of explosive, whose detection is urgently demanded to be explored. Adjusting metal nodes from Mg^{2+} to Ni^{2+} and Co^{2+} , the weak luminescence of MOFs declines to be nonfluorescent, arising from the ligand-to-metal



Fig. 10 Proposed mechanism for the detection of target DNR or RNA based on a fluorescence biosensor formed from MOF **34** and fluorophore-labelled probe DNA. Reprinted with permission from *Anal. Chem.*, 2015, **87**, 12206–12214. Copyright 2015 ACS.



Fig. 11 Crystal structures of Mg^{2+} , Ni^{2+} , and Co^{2+} constructed TABD-MOFs and a schematic representation of the mechanism for detection of the energetic heterocyclic compounds by the three TABD-MOFs. LMCT: ligand-to-metal charge transfer. Reprinted with permission from *J. Am. Chem. Soc.*, 2014, **136**, 15485–15488. Copyright 2014 ACS.

charge transfer (LMCT) effect. The $\text{C}=\text{N}$ or $\text{N}=\text{N}$ group in the 5MR-HCs could destroy the coordination bonds in MOFs through competitive coordination to the metal nodes, releasing the encapsulated AIE fluorophore TABD-COOH which aggregates spontaneously and emits enhanced fluorescence. The method is sensitive in that 5MR-HCs down to 40 nM can be detected *via* the fluorescence signal while that at 6.5 ng cm^{-2} can be identified by the naked eye in several seconds. Releasing the building components from the polymeric MOFs affords signal amplification by an exponential expansion of the amount of small components and by the subsequent AIE of these components.

Actually luminescent MOFs have become excellent candidates for the detection of explosives, thanks to their easy syntheses and structural diversity. Zheng *et al.*⁴⁸ built two luminescent MOFs, whose photoluminescence is quenched only by picric acid (PA), a highly electron-deficient nitrobenzene derivative. Electron-rich π -conjugated fluorescent ligands are introduced as electron donors, which transfer their electrons to the electron-deficient analytes leading to fluorescence quenching. The resonance energy transfer mechanism is also operative, making it an effective sensor for PA. A similar mechanism of electron transfer is shown to operate in Lu's system with high thermostability up to 500°C .⁴⁹ The MOF material with high porosity allows its framework to accommodate guests of varying sizes. The highly stable MOF fluorescence sensor is able to detect a trace amount of nitroaromatic explosives, selectively with no interference from water. Fluorophore modified MOFs represent highly promising sensors as their porous structures accommodate guests of suitable size with the possibility of close contact, leading to highly effective electron and/or energy transfer that ensure high sensitivity, together with high selectivity.

Conclusions and perspectives

Despite the unique advantages of supramolecular sensors and the tremendous success they have achieved, they are still at a very early stage for practical applications. This Feature Article is thus hoped to provide an overview of the current state of the art; particular attention is given to how supramolecular



aggregates have been applied for sensing to achieve a better performance, while the designing of chemosensor-like building blocks is much easier. Given the fact that supramolecular aggregates have been extensively studied for many decades and many of the outcomes of those efforts can be extended to the new applications outlined here, we thus wish to invite those working in the field of supramolecular aggregates to join together, bringing in their expertise to facilitate the exploration of aggregates for sensing. There are indeed several concerns that remain to be clarified, for example, the size distributions and concentrations, and even the morphology of the aggregates in complicated systems, to ensure whether they may or may not influence the sensing signaling. High enough local concentration of the building blocks needs to be loaded into bio-systems such as cells to allow aggregation and consequently to perform *in vivo* sensing and imaging. Future exploration of supramolecular aggregate based sensors is perhaps directed towards detecting diverse biologically important species, to meet the requirement for practical applications such as clinical detection and medical examinations. For example, the development of simple and reliable sensors for point-of-care personalized medicine holds great potential.

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Notes and references

- B. Daly, J. Ling and A. P. de Silva, *Chem. Soc. Rev.*, 2015, **44**, 4203–4211.
- J. Sun, M. Cai and J. J. Lavigne, *Supramol. Chem.*, John Wiley & Sons, Ltd, 2012.
- L. You, D. Zha and E. V. Anslyn, *Chem. Rev.*, 2015, **115**, 7840–7892.
- M. F. Geer and L. S. Shimizu, *Supramol. Chem.*, John Wiley & Sons, Ltd, 2012.
- D. Gorl, X. Zhang and F. Wurthner, *Angew. Chem., Int. Ed.*, 2012, **51**, 6328–6348.
- S. Rochat and T. M. Swager, *ACS Appl. Mater. Interfaces*, 2013, **5**, 4488–4502.
- Z. Chen, Q. Wang, X. Wu, Z. Li and Y.-B. Jiang, *Chem. Soc. Rev.*, 2015, **44**, 4249–4263.
- J.-S. Shen, D.-H. Li, Y.-B. Ruan, S.-Y. Xu, T. Yu, H.-W. Zhang and Y.-B. Jiang, *Luminescence*, 2012, **27**, 317–327.
- C. Li, J. Schöneboom, Z. Liu, N. G. Pschirer, P. Erk, A. Herrmann and K. Müllen, *Chem. – Eur. J.*, 2009, **15**, 878–884.
- T. Ye, R. Singh, H. J. Butt, G. Floudas and P. E. Keivanidis, *ACS Appl. Mater. Interfaces*, 2013, **5**, 11844–11857.
- X. Feng, Y. An, Z. Yao, C. Li and G. Shi, *ACS Appl. Mater. Interfaces*, 2012, **4**, 614–618.
- B. A. Jones, M. J. Ahrens, M. H. Yoon, A. Facchetti, T. J. Marks and M. R. Wasielewski, *Angew. Chem., Int. Ed.*, 2004, **43**, 6363–6366.
- C. Zhao, Y. Zhang, R. Li, X. Li and J. Jiang, *J. Org. Chem.*, 2007, **72**, 2402–2410.
- T. Tang, K. Peneva, K. Mullen and S. E. Webber, *J. Phys. Chem. A*, 2007, **111**, 10609–10614.
- Y. Zheng, H. Long, G. C. Schatz and F. D. Lewis, *Chem. Commun.*, 2005, 4795–4797.
- Y.-B. Ruan, A.-F. Li, J.-S. Zhao, J.-S. Shen and Y.-B. Jiang, *Chem. Commun.*, 2010, **46**, 4938–4940.
- A. K. Dwivedi, M. Pandeewar and T. Govindaraju, *ACS Appl. Mater. Interfaces*, 2014, **6**, 21369–21379.
- A. Weissenstein and F. Würthner, *Chem. Commun.*, 2015, **51**, 3415–3418.
- K.-R. Wang, H.-W. An, R.-X. Rong, Z.-R. Cao and X.-L. Li, *Biosens. Bioelectron.*, 2014, **58**, 27–32.
- X. Wu, X.-X. Chen, B.-N. Song, Y.-J. Huang, Z. Li, Z. Chen, T. D. James and Y.-B. Jiang, *Chem. – Eur. J.*, 2014, **20**, 11793–11799.
- X.-X. Chen, X. Wu, P. Zhang, M. Zhang, B.-N. Song, Y.-J. Huang and Y.-B. Jiang, *Chem. Commun.*, 2015, **51**, 13630–13633.
- Y. Wang, J. Chen, H. Jiao, Y. Chen, W. Li, Q. Zhang and C. Yu, *Chem. – Eur. J.*, 2013, **19**, 12846–12852.
- C. Yu and V. W.-W. Yam, *Chem. Commun.*, 2009, 1347–1349.
- Y.-J. Huang, W.-J. Ouyang, X. Wu, Z. Li, J. S. Fossey, T. D. James and Y.-B. Jiang, *J. Am. Chem. Soc.*, 2013, **135**, 1700–1703.
- X. Wu, X.-X. Chen, B.-N. Song, Y.-J. Huang, W.-J. Ouyang, Z. Li, T. D. James and Y.-B. Jiang, *Chem. Commun.*, 2014, **50**, 13987–13989.
- J. Luo, Z. Xie, J. W. Lam, L. Cheng, H. Chen, C. Qiu, H. S. Kwok, X. Zhan, Y. Liu, D. Zhu and B. Z. Tang, *Chem. Commun.*, 2001, 1740–1741.
- B. K. An, S. K. Kwon, S. D. Jung and S. Y. Park, *J. Am. Chem. Soc.*, 2002, **124**, 14410–14415.
- Y. Hong, M. Haussler, J. W. Lam, Z. Li, K. K. Sin, Y. Dong, H. Tong, J. Liu, A. Qin, R. Renneberg and B. Z. Tang, *Chem. – Eur. J.*, 2008, **14**, 6428–6437.
- Y. Hong, J. W. Lam and B. Z. Tang, *Chem. Soc. Rev.*, 2011, **40**, 5361–5388.
- Z. Zhu, L. Xu, H. Li, X. Zhou, J. Qin and C. Yang, *Chem. Commun.*, 2014, **50**, 7060–7062.
- H. Tong, Y. Hong, Y. Dong, M. Haussler, J. W. Lam, Z. Li, Z. Guo, Z. Guo and B. Z. Tang, *Chem. Commun.*, 2006, 3705–3707.
- L.-J. Chen, X.-Y. Ren, N.-W. Wu, B. Sun, J.-Q. Ma, L. Zhang, H. Tan, M. Liu, X. Li and H.-B. Yang, *J. Am. Chem. Soc.*, 2015, **137**, 11725–11735.
- X. Wang, J. Hu, T. Liu, G. Zhang and S. Liu, *J. Mater. Chem.*, 2012, **22**, 8622–8628.
- L. Liu, G. Zhang, J. Xiang, D. Zhang and D. Zhu, *Org. Lett.*, 2008, **10**, 4581–4584.
- A. Fermi, G. Bergamini, M. Roy, M. Gingras and P. Ceroni, *J. Am. Chem. Soc.*, 2014, **136**, 6395–6400.
- I. G. Occhiuto, M. Samperi, M. Trapani, G. De Luca, A. Romeo, R. F. Pasternack and L. M. Scolaro, *J. Inorg. Biochem.*, 2015, **153**, 361–366.
- T. Sanjji, M. Nakamura and M. Tanaka, *Tetrahedron Lett.*, 2011, **52**, 3283–3286.
- A. T. Aron, K. M. Ramos-Torres, J. A. Cotruvo, Jr. and C. J. Chang, *Acc. Chem. Res.*, 2015, **48**, 2434–2442.
- J. H. Ye, J. Liu, Z. H. Wang, Y. Bai, W. C. Zhang and W. J. He, *Tetrahedron Lett.*, 2014, **55**, 3688–3692.
- F. Hu, Y. Y. Huang, G. X. Zhang, R. Zhao and D. Q. Zhang, *Tetrahedron Lett.*, 2014, **55**, 1471–1474.
- F. Cheng, E. M. Bonder and F. Jäkle, *J. Am. Chem. Soc.*, 2013, **135**, 17286–17289.
- Q. Zhang, Y. Hong, N. Chen, D. D. Tao, Z. Li and Y.-B. Jiang, *Chem. Commun.*, 2015, **51**, 8017–8019.
- Z. Yao, Y. Yang, X. Chen, X. Hu, L. Zhang, L. Liu, Y. Zhao and H.-C. Wu, *Anal. Chem.*, 2013, **85**, 5650–5653.
- Y. Guo, X. Feng, T. Han, S. Wang, Z. Lin, Y. Dong and B. Wang, *J. Am. Chem. Soc.*, 2014, **136**, 15485–15488.
- D. H. Li, J. S. Shen, N. Chen, Y. B. Ruan and Y. B. Jiang, *Chem. Commun.*, 2011, **47**, 5900–5902.
- Y. Cui, B. Li, H. He, W. Zhou, B. Chen and G. Qian, *Acc. Chem. Res.*, 2016, **49**, 483–493.
- S.-P. Yang, S.-R. Chen, S.-W. Liu, X.-Y. Tang, L. Qin, G.-H. Qiu, J.-X. Chen and W.-H. Chen, *Anal. Chem.*, 2015, **87**, 12206–12214.
- Z.-Q. Shi, Z.-J. Guo and H.-G. Zheng, *Chem. Commun.*, 2015, **51**, 8300–8303.
- Y.-N. Gong, L. Jiang and T.-B. Lu, *Chem. Commun.*, 2013, **49**, 11113–11115.

