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# A cucurbit[8]uril-stabilized 3D charge transfer supramolecular polymer with a remarkable confinement effect for enhanced photocatalytic proton reduction and thioether oxidation†

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A highly water-soluble tetrahedral compound T1 that contains one tetraphenylmethane core and four 2-oxynaphthelene-CH<sub>2</sub>-4,4'-bipyridinium (NP-CH<sub>2</sub>-DIPY) side arms has been prepared. Control experiments reveal that NP-CH<sub>2</sub>-BIPY undergoes antiparallel dimerization which is stabilized by two charge transfer interactions with NP as an electron-rich donor and BIPY as an electron-deficient acceptor. This binding motif drives T1 to aggregate into a nanoscale supramolecular polymer SP in water. Adding 2 equivalents of cucurbit[8]uril to the solution of SP leads to the formation of a more stable three dimensional supramolecular polymer CBSP through a 2 + 2 binding motif in which two CB[8] rings encapsulate one charge transfer complex formed by the antiparallel NP-CH<sub>2</sub>-DIPY dimer. As a water soluble porous supramolecular polymer, CBSP exhibits a remarkable confinement effect for visible light-induced proton reduction to produce hydrogen and L-methionine thioether oxidation to the corresponding sulfoxide derivative in water.

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### Introduction

Supramolecular polymers have emerged from the combination of conventional covalently bonded polymers and non-covalent bonding motifs. 1-11 Pioneering works by Lehn and Meijer triggered extensive investigations of hydrogen bonding as a driving force for the generation of supramolecular polymers, 12-15 which were followed by the applications of other non-covalent forces, including coordination, 16,17 donoracceptor interaction, 18,19 solvophobicity 20-22 and ionic pairing electrostatic interaction. 23-26 Most of the early reports of supramolecular polymers have been realized in organic media of low polarity. Hydrophobicity provides the most efficient approach to construct supramolecular polymers in water.<sup>27–29</sup> In this context, many robust connecting patterns have been developed through encapsulation of cyclodextrins and cucurbiturils for hydrophobic monomeric or dimeric species. 21,30-36 The application of polymeric and multitopic building blocks has allowed for the fabrication of three-dimensional (3D)

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supramolecular polymers in water with enhanced stability that is not attainable for linear systems. In recent years, such 3D supramolecular polymers have been vigorously investigated as biocompatible materials for self-healing, drug delivery and bioimaging. Turther development of 3D water-soluble, porous supramolecular polymers should open new opportunities for the exploration of novel porosity-based functions in aqueous media. Currently, examples of the application of supramolecular polymers for homogeneous catalysis in aqueous media are very rare. To expand this potential, the establishment of new binding patterns for the construction of functional supramolecular architectures is highly desirable.

We previously reported the generation of water-soluble, three-dimensional supramolecular polymers from the coassembly of cucurbit[8]uril (CB[8]) and rationally designed multitopic building blocks, 43-45 which is driven by the encapsulation of the CB[8] macrocycle for intermolecular homodimers of peripheral hydrophobic aromatic units of the multitopic building blocks. 30-32 Given the high stability of the 2 + 2 complexation patterns formed between CB[8] and elongated aromatic guests, 46-51 we envisioned that the implementation of this enhanced binding motif might lead to the generation of new, highly stable 3D supramolecular polymers that exhibit unique functions. We herein describe that a tetraphenylmethane-cored compound that bears four donor-acceptor-type side arms can co-assemble with CB[8] in water to form a

highly stable charge transfer complexation-based porous supramolecular polymer, which exhibits a remarkable confinement effect in water for enhanced visible light-induced proton reduction and thioether oxidation to afford hydrogen gas and a sulfone.

### Results and discussion

When an electron-rich aromatic unit is connected to an electron-deficient aromatic unit with a short linker that does not allow for intramolecular interaction, the resulting ditopic molecule may stack to form a homodimer stabilized by two intermolecular donor-acceptor interactions or charge transfer complexations. 46-51 The encapsulation of one CB[8] macrocycle for the single donor-acceptor complex can further stabilize the whole dimer through multivalency. 52,53 Zhang et al. utilized this strategy to construct a main-chain supramolecular 1',1"-(butane-1,4-divl)bis(1-(anthracen-2from ylmethyl)-4,4'-bipyridine-1,1'-diium) bromide.54 To develop a new binding motif to fabricate highly stable 3D supramolecular polymers in water, we designed and synthesized a tetrahedral compound T1 which bears four hydrophilic diglycol chains (Scheme 1). It was envisioned that the 2-oxynaphthelene and 4,4'-dipyridinium units in the four peripheral aromatic arms would undergo intermolecular dimerization which could be further stabilized by two CB[8] macrocycles to afford a 3D donor-acceptor-type supramolecular polymer.

The synthetic route to compound T1 is shown in Scheme 1. Compounds 1 and  $2^{55}$  were condensed in acetonitrile under reflux to afford compound 3 in 83% yield. Compound 3 was then reduced with LiAlH<sub>4</sub> in tetrahydrofuran to give compound 4 in 97% yield. The treatment of 4 with phosphorus tribromide produced 5 in 58% yield. Finally, compound 5 was condensed with  $6^{56}$  in  $N_iN_i$ -dimethylformamide to afford T1 in 66% yield, which had good solubility in water (>2.5 mM). Compounds M1, M2 and BVM were prepared as control molecules. The details for their synthesis are provided in the ESI.†

The <sup>1</sup>H NMR spectrum of compound **T1** in D<sub>2</sub>O exhibited one set of signals. When CB[8] was added to the solution, the resolution of the signals continuously decreased and the spectrum of their 1:4 mixture only gave rise to broad unidentifiable signals (Fig. S1†). This observation indicated that strong binding occurred between the two compounds, but the binding stoichiometry could not be established. Thus, <sup>1</sup>H NMR titration experiments were carried out for control M1 and CB[8] in D<sub>2</sub>O. Upon the addition of CB[8], a new set of signals appeared, whereas those of free M1 became weakened (Fig. 1a). With the increase of the amount of CB[8], this set of signals became enhanced, but did not undergo any shifting. When 1.0 equivalent of CB[8] was added, the signals of free M1 vanished completely. Adding another 1.0 equivalent of CB[8] did not cause the generation of new signals. These results confirmed that the two components formed a very stable 2 + 2 complex, which did not undergo quick exchange with the free components on the <sup>1</sup>H NMR timescale. The

Scheme 1 The structures of compounds T1, M1, M2, BVM and  $Na_4[Ru (BDC)_3]$  and the synthetic route to compound T1.

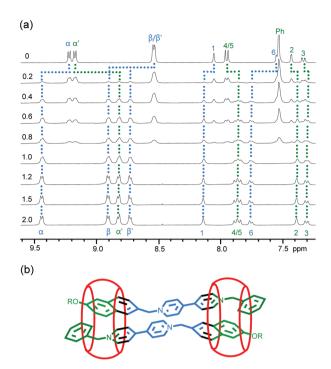


Fig. 1 (a) <sup>1</sup>H NMR spectra of the mixture of M1 (5.0 mM) and CB[8] (0, 0.2, 0.4, 0.6, 0.8, 1.0, 1.2, 1.5 and 2.0 equiv.) in  $D_2O$  at 25 °C. (b) 2 + 2 binding pattern between M1 and CB[8] revealed by the above <sup>1</sup>H NMR spectra in D<sub>2</sub>O, highlighting a slipped antiparallel stacking of the aromatic units of two molecules of M1 in the cavity of the CB[8] macrocycles.

signals of M1 within the downfield area in the CB[8]-free solution and the solution of its 1:1 mixture were assigned using the 2D NMR technique (Fig. S2 and S3†). A comparison of the two spectra revealed that the signals of  $\alpha'$ - and 2-5-H of M1 underwent significant upshifting, which can be attributed to the shielding of the CB[8] macrocycle through encapsulation. In contrast, the signals of  $\alpha$ -,  $\beta$ -,  $\beta$ -, 1- and 6-H shifted downfield pronouncedly, which could be caused by the de-shielding of the CB[8] macrocycle and possibly also by forming intermolecular hydrogen bonding with the carbonyl oxygen atoms of CB[8]. These results supported that the two compounds formed a 2 + 2 complex with the binding pattern shown in Fig. 1b.

UV-vis spectra were then recorded for both M1 and T1 with the addition of an incremental amount of CB[8] (Fig. 2). For both compounds, adding CB[8] caused a significant decrease of the absorption centred at 233 nm, which gave an inflection point when 1.0 or 4.0 equiv. of CB[8] were added. This is because T1 bears four side arms that can bind to CB[8]. These results indicated that both compounds formed stable complexes with CB[8] at the respective low concentration. By assuming a low limit of 95% of the two components that formed the 2 + 2 complexes, the association constant  $(K_a)$  for the four-component complex  $(M2)_2 \subset (CB[8])_2$  could be calculated to be  $1.9 \times 10^{16} \text{ M}^{-3}$ , whereas the apparent  $K_a$  for the similar complexes formed by the side arms of T1 and CB[8]

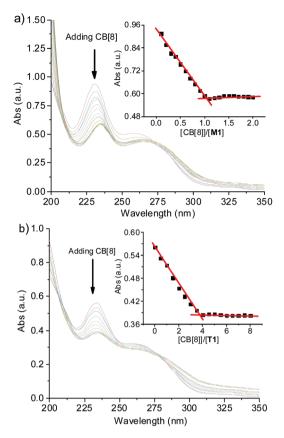


Fig. 2 UV-vis absorption spectra of (a) M1 (20  $\mu$ M) and (b) T1 (2.5  $\mu$ M) in water in the presence of an incremental amount of CB[8] at 25 °C. Inset: the absorbance at 233 nm vs. [CB[8]]/[M1] or [T1].

was determined to be  $1.5 \times 10^{17} \text{ M}^{-3}$ . The latter was substantially larger than the former, well reflecting the multivalency of the tetratopic molecule in binding to CB[8].

2,6-Dialkoxynaphthalene has been demonstrated as an electron-rich aromatic unit to produce intermolecular donoracceptor complexes. 46,57-59 We conjectured that the 2-alkoxynaphthalene unit at a high concentration might also function as an electron-rich donor to form similar complexes, which can be further enhanced through CB[8] encapsulation in water. We thus further recorded the UV-vis absorption spectra of M1 and T1 at high concentrations. The UV-vis absorption spectrum displayed a broad absorption band around 390 and 408 nm (Fig. 3a), respectively. Adding 1.0 or 4.0 equivalents of CB[8] to their solution caused significant enhancement of this absorption band and the maximum absorption wavelength also shifted to 405 and 420 nm, respectively. With or without the addition of CB[8], this absorption band exhibited an identical molar absorption coefficient ( $\varepsilon$ ) that was independent of the concentration (Fig. S4–S7 $\dagger$ ). For **T1**, the  $\varepsilon$  values were 393 and 206, and for M1, the values were 134 and 62. Consistent with the above observations, the aqueous solutions of compounds T1 and M1 were brown and light brown (Fig. 3b), which were dependent on the concentration, while adding CB[8] caused darkening of the colour. Because both controls

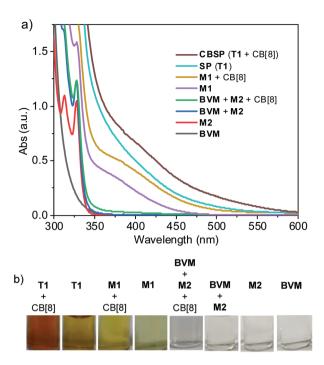


Fig. 3 (a) UV/vis absorption spectrum of the solution of CBSP (T1 + 2CB[8]), SP (T1), M1 + CB[8], M1, BVM + M2 + CB[8], BVM + M2, M2 and BVM in water. For all the solutions, the concentration of the NP and/or BIPY units was kept at 1.0 mM. (b) The photographs of the different solutions highlighting their colours. For all the solutions, the concentration of the BIPY unit was controlled at 5.0 mM.

M2 and BVM did not exhibit significant absorption within the range of long wavelengths (>350 nm) (Fig. 3a and b), these results supported that both M1 and T1 formed charge transfer complexes, which were further strengthened by CB[8]. These observations showed that encapsulation complexation of CB[8] for the donor-acceptor dimer of M1 was remarkably enhanced in the 2 + 2 binding as compared with the simple three-component complexation. The results also showed that the charge transfer complexes formed by T1 was more stable than those of M1 in the absence or presence of CB[8].

Since the UV-vis absorption spectra could not be recorded for the solution of even higher concentrations, we then conducted <sup>1</sup>H NMR experiments for the solution of M1 and T1 at concentrations higher than those used for the above UV-vis spectroscopy. When the concentration of T1 was increased from 0.5 mM to 2.5 mM, which corresponded to the concentration of 2.0 and 10 mM for the appended aromatic arms, the signals of the protons of the bipyridinium and naphthalene units in the <sup>1</sup>H NMR spectrum all underwent significant upshifting (0.1-0.6 ppm) (Fig. S8†). Accompanied by this continuous upshifting of the signals, their resolution also decreased gradually, indicating increasing intermolecular stacking. In contrast, for M1, increasing the concentration from 2 mM to 10 mM did not cause large shifting of the signals of the two aromatic units (Fig. S9†). These results further confirmed that the bipyridinium and naphthalene units of T1 formed more stable charge transfer complexes.

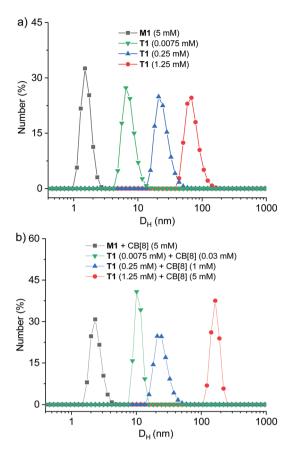


Fig. 4 Dynamic light scattering profile of (a) the solution of M1 and T1 in water and (b) the solution of the mixture of M1 (1:1) and T1 (1:4) with CB[8] in H2O at 25 °C.

Dynamic light scattering (DLS) experiments further revealed the formation of large aggregates by T1 in the absence and presence of CB[8] (Fig. 4). At a high concentration of 5.0 mM, the solution of M1 gave rise to a very small hydrodynamic diameter  $(D_{\rm H})$  of 1.5 nm. With the addition of 1.0 equivalent of CB[8],  $D_H$  was increased slightly to 2.3 nm, which was still very small. In contrast, at a concentration of 1.25 mM, which corresponded to the concentration of 5 mM for the four side arms, the  $D_{\rm H}$  value was determined to be 68 nm. With the addition of CB[8] (5 mM), the  $D_{\rm H}$  value was increased to 165 nm. Both results reflected that T1 itself underwent significant aggregation to afford nanoscale supramolecular entities, which were converted into even large supramolecular entities through binding with CB[8]. As expected, diluting the solution caused the decrease of the  $D_{\rm H}$  value for both T1 and its mixture with CB[8] (Fig. 4 and S10†). However, even at a low concentration of 0.0075 mM, two solutions still gave rise to  $D_{\rm H}$  of 6.5 and 10 nm. Taken together, the above <sup>1</sup>H NMR, UV-vis and DLS experiments consistently supported that T1 aggregated in water to afford a charge transfer-type supramolecular polymer SP and its 1:4 mixture with CB[8] gave rise to a more stable and even larger 3D supramolecular polymer CBSP (Fig. 5). Given the tendency of tetrahedral building blocks to form 3D porous architectures in solutions, we propose that the 3D

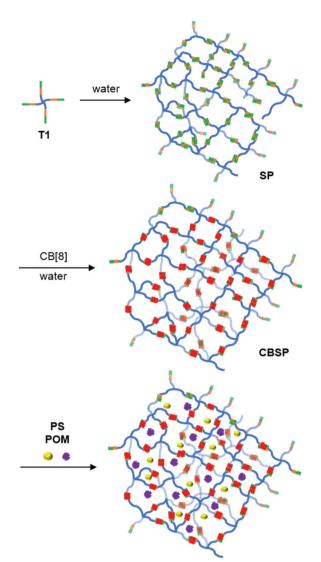


Fig. 5 Representation of the self-assembly of the 3D supramolecular polymer CBSP from T1 and CB[8] in water and its inclusion of the photosensitizer and the POM catalyst.

supramolecular polymers formed by T1 and CB[8] should possess intrinsic porosity. By assuming an ideal 3D diamondoid structure, CBSP would form porous channels with an aperture diameter of 4.0 nm (Fig. S11†).

Porous organic polymers are ideal materials for the attainment of the confinement effect for catalysing various heterogeneous chemical transformations. 60-65 We have also demonstrated that homogeneous 3D porous supramolecular organic frameworks can exhibit an important confinement effect for homogeneous photocatalysis in aqueous media. 66-69 The high stability of CBSP allows it to maintain a nanoscale size at micromolar concentrations of the building blocks. We thus studied the capability of CBSP for simultaneous enrichment of the photosensitizer Na<sub>4</sub>[Ru(BDC)<sub>3</sub>] (PS, BDC = 2,2'-bipyridyl-4,4'-carboxylate) (Scheme 1) and the Keggin-type polyoxometallate catalyst Na<sub>3</sub>PW<sub>12</sub>O<sub>40</sub> (POM), which forms an important integrated photocatalytic system for visible light induced

reduction of protons into H<sub>2</sub>.67,70 This enrichment was expected to take place due to the multianionic character of both species, which should lead to the inclusion of the two species in the interior of CBSP driven by multivalent ionic pairing electrostatic attraction. Dialysis experiments using a membrane filter (molecular weight cut-off: 2.0 kDa) for the solution of CBSP ([T1] = 0.1 mM) and  $Na_4[Ru(BDC)_3]$  (20  $\mu$ M) showed that CBSP could completely suppress the diffusion of the photosensitizer,67 which confirmed the efficient adsorption of the photosensitizer by the porous supramolecular polymer (Fig. S12†). This adsorption or inclusion would increase their efficient concentration in the interior of the polymer and thus facilitate electron transfer once the photosensitizer molecules are excited by visible light. Hydrogen evolution reaction (HER) experiments were then conducted by irradiating an aqueous solution (2 mL) that contained PS (20 µM), POM (2 µM) and CBSP at different concentrations. The results are presented in Table 1. In the absence of CBSP, the turnover number (TON) for the HER was determined to be 43 (entry 1). Adding 25, 50 and 100 µM CBSP could improve the TON to 208, 538 and 1683 (entries 2-4), respectively, which represents a 4.8, 12.5 and 39.1-fold increase of the TON. Further addition of CBSP did not cause the increase of the TON (entries 5 and 6), which may be rationalized by considering that both PS and POM had been completely included in the interior of CBSP and further addition of CBSP would rather decrease the apparent concentration of PS and POM in its interior. With the addition of SP at 100 µM, the TON was determined to be 78 (entry 7), which indicated that SP (T1) only slightly promoted the catalytic efficiency of the PS-POM system. The important confinement effect displayed by CBSP should reasonably reflect its increased stability and probably the pore rigidity due to the encapsulation of CB[8] for T1. When M1 or its 1:1 mixture with CB[8] was added at an identical concentration of the donor and acceptor, the TON was found to be 36 and 61 (entries 8 and 9). Given their high concentration, these results showed that they did not impose significant influence on the catalysis of the **PS-POM** system.

Table 1 Visible light-induced proton reduction into hydrogen gas in the presence of CBSP or the control compound or the mixture<sup>a</sup>

|       | CBSP or    | Conc.b        | Time | PS   | POM  |                    |
|-------|------------|---------------|------|------|------|--------------------|
| Entry | control    | (μ <b>M</b> ) | (h)  | (μM) | (μM) | $\mathrm{TON}^{c}$ |
| 1     | CBSP       | 0             | 24   | 20   | 2    | 43                 |
| 2     | CBSP       | 25            | 24   | 20   | 2    | 208                |
| 3     | CBSP       | 50            | 24   | 20   | 2    | 538                |
| 4     | CBSP       | 100           | 24   | 20   | 2    | 1683               |
| 5     | CBSP       | 150           | 24   | 20   | 2    | 1323               |
| 6     | CBSP       | 200           | 24   | 20   | 2    | 987                |
| 7     | SP         | 100           | 24   | 20   | 2    | 78                 |
| 8     | M1         | 400           | 24   | 20   | 2    | 36                 |
| 9     | M1 + CB[8] | 400           | 24   | 20   | 2    | 61                 |

<sup>&</sup>lt;sup>a</sup> The hydrogen evolution reaction was conducted under irradiation of a xenon lamp (300 W, light power density: 200 mW cm<sup>-2</sup>) for 24 h in the acid solution (pH = 1.8) containing methanol (20%, v/v) with a total volume of 2.0 mL. <sup>b</sup> For CBSP, referred to as [T1]. <sup>c</sup> TON determined by analysing the headspace gas with GC.

It has been reported that the bipyridinium radical cation (BIPY\*+) can produce reactive oxygen species (ROS) in the presence of aerobic O2.71 We thus further studied the potential of CBSP for photocatalytic aerobic oxidation of organic sulfides to sulfoxide derivatives, for which L-methionine (Met-S) was used as a model substrate. The photocatalytic oxidation of Met-S to the sulfoxide derivative Met-S=O has been recently investigated by Stoddart et al. with BIPY-derived [2]catenane as a catalyst.72 Under the standard reaction conditions, no oxidation of Met-S took place. In the presence of CBSP (10 mol%), the reaction proceeded smoothly (Table 2, entries 1-7). After 18 h, 90% of Met-S was consumed to afford Met-S=O in 88% yield. When the reaction time was  $\leq$ 18 h, no sulfone derivative was detected from the reaction. When the reaction time was increased to 24 h, Met-S could be consumed completely, but the yield of Met-S=O was reduced to 74%, due to further oxidation of Met-S=O to the corresponding sulfone derivative which was confirmed by observing the formation of the sulfone derivative. In the presence of SP with an identical concentration (10 mol%) of T1, irradiating the solution for 18 h led to 68% consumption of Met-S and Met-S=O was produced in 65% yield (Table 2, entry 8). Clearly, the formation of the BIPY\*+ radical cations by CBSP through visible lightinduced electron transfer from Met-S was more efficient than that by SP. That is, CBSP also exhibited an important confinement effect for this organic transformation as compared with SP. With the decrease in the amount of both SP and CBSP, the corresponding yield of Met-S=O was reduced (Table 2, entries 9-12). However, CBSP consistently displayed higher catalytic

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Table 2 Oxidation of L-methionine in the presence of CBSP or different controls

Me S OH Catalyst, 
$$O_2$$
 Me S OH NH<sub>2</sub> OH

Met-S Met-S=O

Met-S=O

| Entry | CBSP or control | Amount (mol%) | Time<br>(h) | <b>Met-S</b><br>Conv. <sup>b</sup> (%) | <b>Met-S≔O</b> <sup>b</sup><br>Yield (%) |
|-------|-----------------|---------------|-------------|--|--|
| 1     | CBSP            | 10            | 6           | 29                                     | 26                                       |
| 2     | CBSP            | 10            | 9           | 53                                     | 46                                       |
| 3     | CBSP            | 10            | 12          | 67                                     | 59                                       |
| 4     | CBSP            | 10            | 15          | 75                                     | 71                                       |
| 5     | CBSP            | 10            | 18          | 90                                     | 88                                       |
| 6     | CBSP            | 10            | 21          | 93                                     | 79                                       |
| 7     | CBSP            | 10            | 24          | 100                                    | 74                                       |
| 8     | SP (T1)         | 10            | 18          | 68                                     | 65                                       |
| 9     | CBSP            | 5             | 24          | 43                                     | 34                                       |
| 10    | SP (T1)         | 5             | 24          | 15                                     | 12                                       |
| 11    | CBSP            | 2.5           | 24          | 31                                     | 22                                       |
| 12    | SP (T1)         | 2.5           | 24          | 10                                     | 8  |
| 13    | M1              | 40            | 18          | 8                                      | 7  |
| 14    | M1 + CB[8]      | 40            | 18          | 12                                     | 9  |

<sup>a</sup> Standard reaction conditions: Met-S (5 mmol), CBSP (0.5 mmol, 10%) in H<sub>2</sub>O (2 mL), irradiation with a blue LED strip (total 18 W, light power density: 7 mW cm<sup>-2</sup>) under an aerobic oxygen atmosphere at 25 °C. b The conversion of Met-S and the yield of Met-S=O were determined by HPLC.

activity. With an identical BIPY2+ concentration (40 mol%), M1 or its 1:1 mixture with CB[8] only caused very low conversion of Met-S and the formation of Met-S=O (Table 2, entries 13 and 14). All the results are consistent with those revealed above for the photocatalytic reduction of protons to hydrogen, again supporting the important confinement effect of CBSP. In water, CBSP should not be able to adsorb hydrophilic Met-S in its interior. Thus, this confinement effect might be rationalized by considering that CBSP produces a high local concentration of BIPY<sup>2+</sup> units in the supramolecular polymer, which facilitated its single electron reduction to BIPY\*+ by Met-S molecules in the interior of the supramolecular polymer and subsequent electron transfer from BIPY\*+ to O2 to produce O2. In this way, the oxidation of Met-S could be accelerated remarkably in the interior of CBSP.

### Conclusions

In summary, we have demonstrated that the cucurbit[8]uril encapsulation-based 2 + 2 donor-acceptor binding pattern can be utilized to construct a highly stable, water-soluble 3D porous supramolecular polymer from a tetrahedral building block. The new supramolecular polymer possesses a nanoscale size at micromolar scale concentrations. The new porous supramolecular polymer has been revealed to exhibit a remarkable confinement effect not only for visible light-induced proton reduction to hydrogen gas by simultaneously including a ruthenium complex photosensitizer and a polyoxometalate catalyst, but also for aerobic oxygen oxidation of methionine thioether to the corresponding sulfoxide derivative. This work opens up a new possibility for the application of the cucurbit [8] encapsulation-based 2 + 2 binding pattern to fabricate functional supramolecular architectures.

### Conflicts of interest

There are no conflicts to declare.

### Author contributions

ZTL and DWZ conceptualized the ideas and supervised the investigation. ZL, QL, JDS, ZKW and HW performed the experiments and collected the data. ZL and ZTL analysed the data. ZL wrote the original draft. ZTL and DWZ finished the writing. All authors read and contributed to the revisions.

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