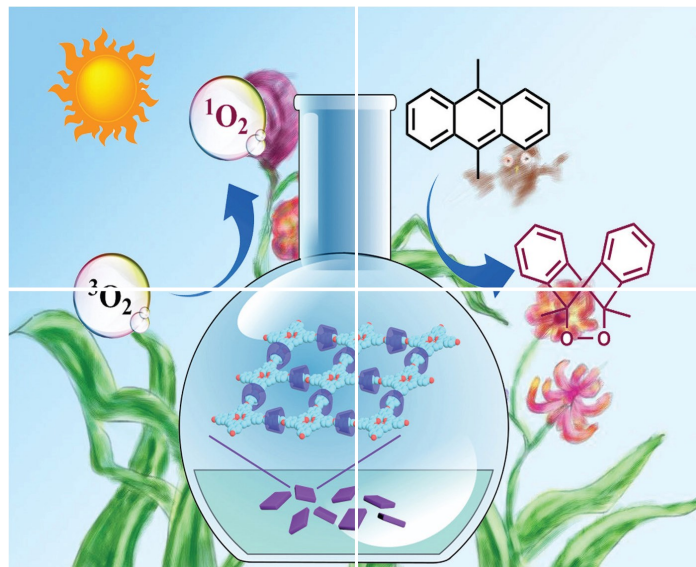


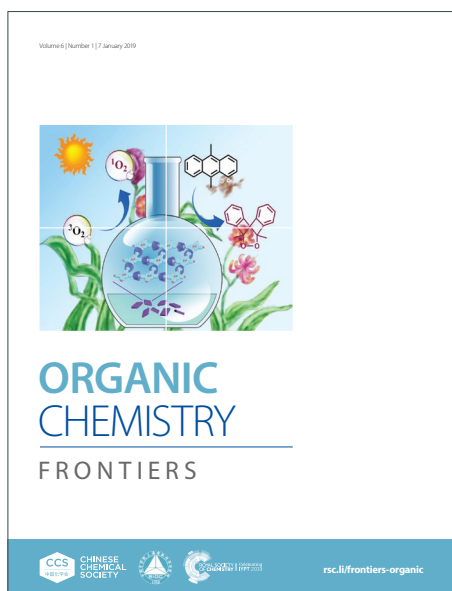
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ARTICLE

Fluorescent visualization of cucurbit[8]uril-triggered dynamic host-guest assemblies

Xiaodong Zhang, Tao Sun and Xin-Long Ni*

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Cucurbit[8]uril (Q[8] or CB[8]), as a synthetic macrocyclic host, play an important role in constructing various water soluble dynamic supramolecular polymers and materials. Here we report host-guest interactions of Q[8] with π -conjugated bipyridinium guests in aqueous solution that resulted in a series of unprecedented dynamic 2:2, 2:3, and 1:2 assemblies (guest/Q[8]). Most importantly, these subtly different host-guest binding modes could be distinguished by naked-eye observation of the resulting quantitative fluorescence emissions. Further characterization suggested that the Q[8] host-stabilized charge-transfer interactions could be dissociated and controlled by the N-substituted alkyl chains on the cationic guests via the formation of 2:3 quaternary complex or U-shaped conformation, which leading to the discovery of new possible binding fashions for the Q[8] host.

Introduction

Cucurbit[8]uril (Q[8] or CB[8]),^{1, 2} a large homologue of the cucurbituril family, is unique because of its ability to form ternary complexes by encapsulating two hetero- or homo-guests in its rigid macrocyclic cavity. In particular, host-stabilized charge-transfer (HSCT),³ as the dominant interaction force, has resulted in the wide use of Q[8] to construct various water soluble dynamic materials in the past decade.⁴⁻¹⁵ Meanwhile, it was also found that Q[8] is able to act as molecular handcuffs to simultaneously include two chromophore dyes in its cavity to form dimer assembly. This not only increases the π - π stacking of the aromatic moieties,¹⁶ but also enhances intra/inter-molecular charge transfer (ICT) of the inclusion guest dyes.¹⁷

Recently, it was discovered that π -conjugated bipyridinium derivatives could serve as both electron-deficient and electron-rich groups that could dimerize inside the Q[8] cavity to form highly stable Q[8]-enhanced ternary complexes¹⁸⁻²⁰ and supramolecular polymers²¹⁻²³ with enhanced light efficiency. More recently, Scherman's group,²⁴⁻²⁶ Schalley's group,²⁷ and Ma's group²⁸ revealed that complexation of diaryl viologens bearing electron-donating groups with Q[8] could be used to form a series of unique dimer emissions in 2:2 binding mode. These studies indicated the potential of Q[8] as a host in the fabrication of smart tunable luminescent materials. However, to our knowledge, there is few attention in Q[8] encapsulation-triggered

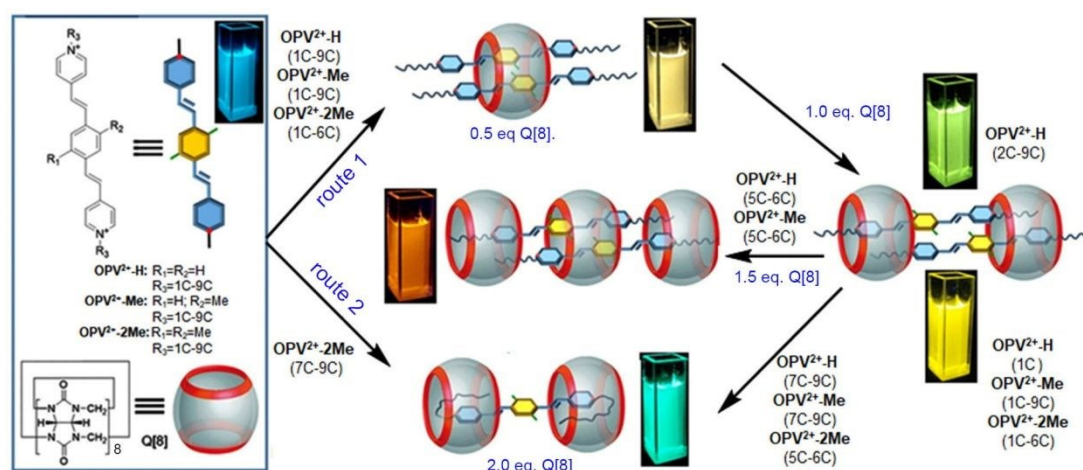
fluorescent signals for monitoring of the accompanying dynamic host-guest interactions.

Fluorescence has been widely used to explore dynamic processes, such as self-assembly,^{29,30} molecular motion³¹ and conformational changes.³² The high sensitivity of fluorescence methods allows for tracking of molecular interactions even at nanomolar concentrations. In particular, it is possible to visualize directly and in real time to monitor processes in situ. For example, Tang et al. exploited aggregation-induced emission fluorescent imaging to successfully demonstrate the transition processes of surfactant micelles and microemulsion droplets.³³ Li and co-workers directly observed the distribution of gelatin in calcium carbonate crystals by super-resolution fluorescence microscopy.³⁴ Yang et al. found that the process and dynamics of coordination-driven self-assembly could be tracked in real-time by fluorescence-resonance energy transfer.³⁵

In this work, a series of different length alkyl chains were introduced into π -conjugated bipyridinium species as guest molecule (Scheme 1). We found that the alkyl chains as the terminal moieties played a crucial role in driving the Q[8] host to form various dynamic host-guest assemblies including 2:2, 2:3, and 1:2 assemblies. In particular, the multiple dynamic Q[8]-based host-guest interactions could be distinguished by naked-eye observation of the resulting quantitative fluorescence emissions. Further studies revealed that the Q[8]-based HSCT interaction could be dissociated or controlled by the alkyl chain lengths and the slightly modification of the π -conjugated aromatic core of the guests.

Key Laboratory of Macrocyclic and Supramolecular Chemistry of Guizhou Province, Guizhou University, Guiyang, 550025, China. E-mail: longni333@163.com.

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Scheme 1. Chemical structures of OPV²⁺ derivatives and Q[8], and representation of Q[8]-based dynamic host-guest assemblies with multiple tunable fluorescent signals. Route 1: Q[8] concentrations dependent host-guest interactions of 2:1, 2:2, 2:3, and 1:2 (guest/Q[8]); route 2: the formation of U-shaped conformation of Q[8] with OPV²⁺-2Me (7C-9C).

Results and discussion

Our previous work revealed that the photophysical properties of oligo(*p*-phenylenevinylene) (OPV²⁺)-based bispyridinium guest molecules could be sensitively switched by the Q[8] host.²¹ Here, to conveniently monitor multimodular host-guest assemblies of Q[8] with the alkyl chain-appended guests, the π -conjugated OPV²⁺ core was selected as the fluorescent reporting moiety. In order to capture and distinguish dynamic host-guest interactions, such as whether the Q[8] host could enclose the phenyl part of the guest, larger barrier groups, such as one or two methyl (Me) groups, were appended to the phenyl moiety to serve as a “brake.” The molecular structure of guests together with plausible fluorescent visible host-guest assemblies are shown in Scheme 1. Guests OPV²⁺-H, OPV²⁺-Me, and OPV²⁺-2Me with different alkyl chains (containing carbon atoms from 1C to 9C), in which I[−] as counter anions for 1C appended guests, and Br[−] as counter anions for other guests. The related guests hereafter abbreviated as OPV²⁺-(H, Me, 2Me) (1C to 9C).

The fluorescence and UV-vis spectra of the guests appeared similar emission peak around 480 nm and absorption peak around 390 nm in aqueous solution, respectively, indicating that the appended alkyl chains did not affect the optical properties of the OPV²⁺ core. In the presence of Q[8] host, OPV²⁺-H and OPV²⁺-Me derived guests appeared similar optical response, while some different fluorescent response behaviour was observed in the case of OPV²⁺-2Me based guests (Fig 1, Figs. S1-S3 and Table S1). For example, an enhanced bathochromic fluorescent emission (yellow colour) shift was observed for all of the guest solutions when the concentration of Q[8] was fixed at 0.5 eq., except for the OPV²⁺-2Me guests (7C-9C), where slight red-shifts were noted

(Figs. 1f). When the concentration of Q[8] was increased to 1.0 eq., considerably enhanced emission around at 550 nm was observed (Fig. 1a, 1d, and Fig. S1), except for guests OPV²⁺-H (2C-9C), where a tunable enhanced blue-shift emission was seen (Fig. S1 and Table S1).

As the concentration of Q[8] was increased to 1.5 eq., no changes in the fluorescence spectra were observed for any of the 1C-4C appended guests (Fig. S1 and Table S1). However, OPV²⁺-H guests (5C-6C) (Fig. 1b) and OPV²⁺-Me (5C-6C) underwent significant red-shift emissions (Table S1), and a blue-shift emission change was observed in the OPV²⁺-2Me (5C-6C) solution (Figs. 1e).

Interestingly, OPV²⁺-H guests (7C-9C) and OPV²⁺-Me (7C-9C) exhibited blue-shift with enhanced emission intensity (Fig. 1c and Table S1), and a slight red-shift of OPV²⁺-2Me (7C-9C) in the case of concentrations of Q[8] was increased from 0.5 eq. to 2.0 eq., respectively (Figs. 1f and Table S1). Essentially, the different fluorescent responses of the same alkyl-attached guests when Q[8] was added is attributed to modification by the Me group on the OPV²⁺ core (Scheme 1).

Compared to the multiple fluorescent signals triggered by different concentrations of Q[8] host, similar changes in the UV-vis absorption spectra were observed (Figs. S2-S3). At first glance, all of the maximum absorbance peaks of the guests at 390 nm decreased remarkably, with bathochromic shift of the band at 400-470 nm in the presence of Q[8] host. Close inspection revealed considerably enhanced red-shifted absorbance peaks for OPV²⁺-H (5C-6C) and OPV²⁺-Me (5C-6C) guests with two isoabsorption points around 413 and 432 nm, and enhanced blue-shifted absorbance peaks (but still red-shifted compared to the free guest) with isoabsorption points around 428 and 441 nm for OPV²⁺-2Me (5C-6C) guests.

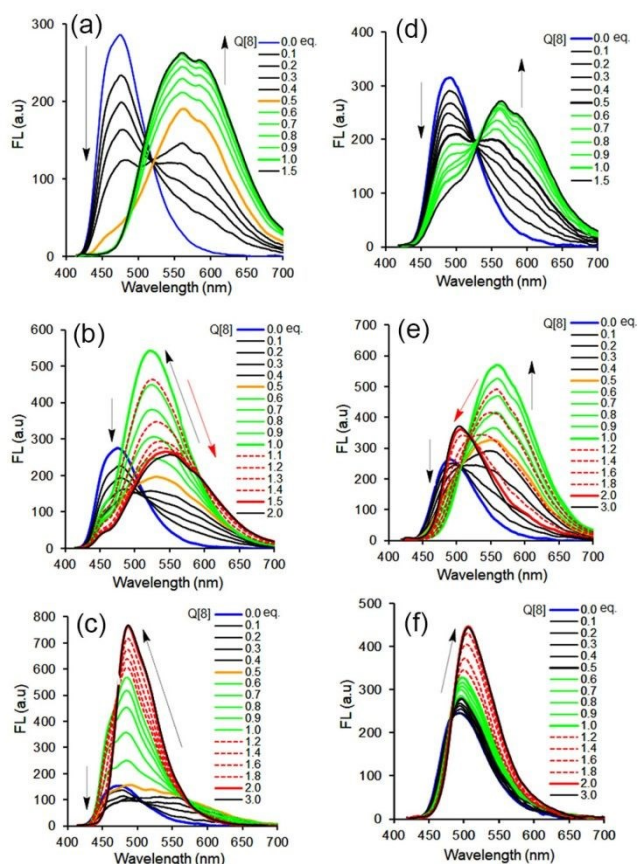


Fig. 1 Fluorescence spectrum changes of **OPV²⁺** core based guests (a-c) **OPV²⁺-H** (1C, 5C, 9C), (d-f) **OPV²⁺-2Me** (1C, 5C, 9C) (each of 10.0 μ M in aqueous solution, pH 7.2) in the presence of increasing concentrations of Q[8] host, respectively. (λ_{exc} = 400 nm)

Generally, guest complexation within the cavity of Q[8] typically results in an upfield shift of the encapsulated protons (shielding effect of the hydrophobic cavity) and a downfield shift of protons in close proximity to the portals of the host (deshielding effect of the carbonyl-rimmed portal). For example, upon titration of 0.5 eq. Q[8] into a guest **OPV²⁺-H** (1C) solution (neutral D₂O), nearly all of the protons on the parent **OPV²⁺** core underwent upfield shifts (Fig. 2a). Particular protons on the ethylene moieties and the phenyl group, including the methyl groups on the **OPV²⁺-Me** and **OPV²⁺-2Me** guests (Figs. S4-S7, apart from **OPV²⁺-2Me** (7C-9C), which will be discussed later on), were upfield shifted significantly. In contrast, no obvious proton chemical shift changes were observed for the N-substituted alkyl chains. These results indicated that the phenyl aromatic and ethylene moieties were encapsulated in the Q[8] cavity to form 2:1 host-guest complexes (guest/Q[8]).²¹ It is believed that the Q[8] host is threaded onto the guests such that it can move back and forth along the aromatic **OPV²⁺** core of the guests. In particular, ion dipole interactions between the positively charged nitrogen atoms of the guests and the carbonyl oxygen atoms of the Q[8] host prevent dissociation. This shuttling motion is relatively fast on the NMR time scale, so all of the signals for the protons

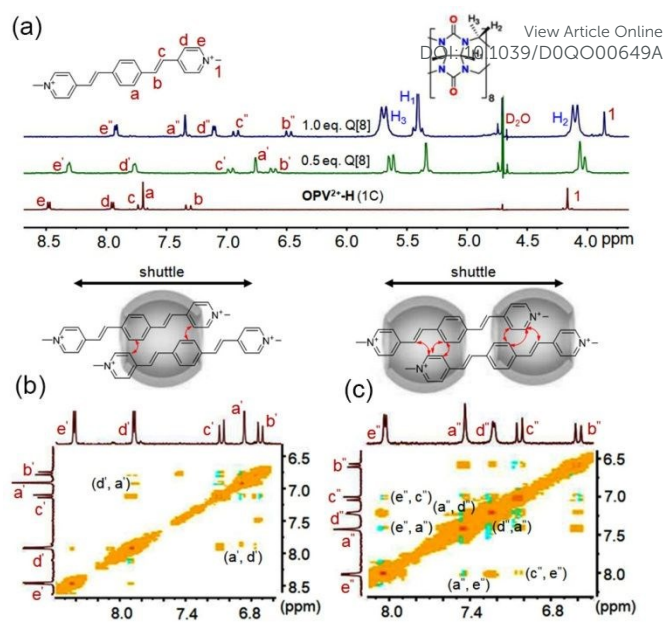


Fig. 2 (a) ¹H NMR spectra of **OPV²⁺-H** (1C) (1.0 mM in D₂O) with different concentrations of Q[8] (0~1.0 mM); ¹H-¹H NOESY of **OPV²⁺-H** (1C)/Q[8] at molar ratios of 2:1 (b) and 2:2 (c)

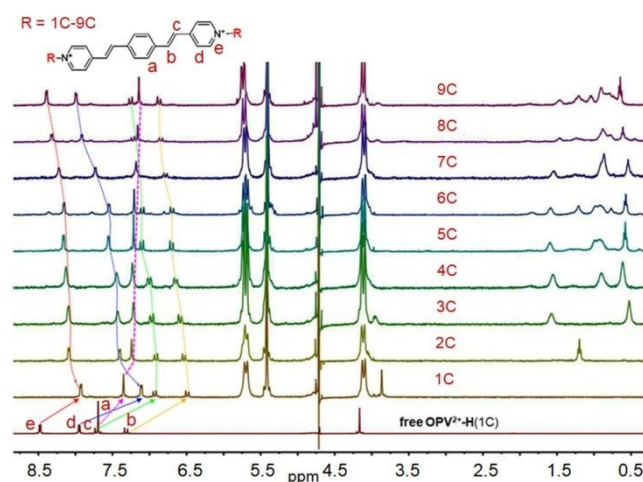


Fig. 3 ¹H NMR spectra of Q[8]/**OPV²⁺-H** (1C-9C) at a molar ratio of 2:2.

on the aromatic core are shifted upfield in the spectrum, and no proton chemical shift changes are observed for the N-substituted alkyl chains. Similar proton signal changes were apparent in the host-guest complexes of **OPV²⁺-Me** (Fig. S5) and **OPV²⁺-2Me** (Fig. S6) systems, but less upfield shift was observed for the protons on the **OPV²⁺** core (Fig. S7), suggesting that the barrier Me and 2Me groups on the phenyl moiety as “brakes” slowed the rate of the Q[8] shuttling motion.

As an example, 2D NMR spectra such as ¹H-¹H NOESY (nuclear Overhauser effect spectroscopy) of Q[8] with guest **OPV²⁺-H** (1C) showed NOE correlation of H_d-H_a, implying the formation of head-to-tail dimer complexes (Fig. 2b). Therefore, complexation inside the confined space of the Q[8] cavity results in distinct π - π stacking (substantial electron-

delocalization due to dimeric stacking along the long molecular axis), which leads to the significant red-shift fluorescence emission (yellow colour) as mentioned in Fig. 1. Compared to the shorter fluorescence life time (around 0.13 ns) and lower quantum yield (around 5%) of those free guests, all of the Q[8]-encapsulated guest assemblies in solution had a long lifetime of 11–15 ns (Table S2) and an enhanced quantum yield (Table S3), apart from OPV²⁺-H (7C–9C), where the fluorescence lifetime was 6.7–8.6 ns. This may be attributed to the long alkyl chain placing the π -conjugated OPV²⁺ fluorophore near to the Q[8] carbonyl portals, which was confirmed by the lower upfield chemical shift of the phenyl protons of OPV²⁺-H (7C–9C) (Fig. S7a).

As the concentration of Q[8] was increased to ~1.0 eq., for OPV²⁺-H guests, the signals for all guest protons, including those of the N-substituted alkyl chains, were located in the upfield area (Figs. 2 and S4). Protons on the pyridinium (H_d and H_e) appeared to be shifted further upfield, whereas the ethylene protons were still in the same position and phenyl protons were shifted downfield (still upfield shifted compared to the free guest). The NOE correlations between H_e–H_{c'}, H_e–H_{a''}, and H_{a''}–H_{d''}, (Fig. 2c) indicated that a dynamic 2:2 complex was formed in this process. Interestingly, closer inspection revealed that the H_d and H_e protons exhibited a gradual downfield shift as the length of the N-substituted alkyl chains increased (Fig. 3). In particular, for OPV²⁺-H (7C–9C) guests, it was observed that the signal for H_d underwent a downfield shift compared to 0.5 eq. of Q[8] (H_d was even downfield shifted, to 7.98 ppm from 7.90 ppm in the free state, suggesting that H_d was in close proximity to the Q[8] portal). This result indicated that the longer alkyl chains might play an important role in switching the dynamic 2:2 host-guest complex from head-to-tail geometries to head-to-head arrangements, which is also supported by the fluorescence spectra changes (weak intensity around at 550 nm) and shorter life time (Table S2). Similar NMR and fluorescence spectra changes were observed in the OPV²⁺-Me and OPV²⁺-2Me systems, suggesting similar host-guest assembly modes. However, the significant proton downfield shift of the Me group and strong fluorescence intensity around 555 nm for OPV²⁺-2Me (1C–6C) implied that the double Me moieties can control and retard the shuttling rate of Q[8] on the guest and favor the head-to-tail complex (Fig. S7c).

Upon increasing the concentration of Q[8] up to 1.5 eq., no further chemical shift changes of the protons were observed in any of the 1C–4C-appended host-guest complexes. However, the proton peaks of the ethylene (H_b and H_c), phenyl (H_a) moieties and the alkyl chains on the 5C–6C-based OPV²⁺-H and OPV²⁺-Me guests, including the monomer Me group on OPV²⁺-Me, exhibited significant upfield shifts (Figs. 4a and S4–S5). The signals of the other protons, such as H_d and H_e, exhibited slightly downfield shifts compared to 1.0 eq. Q[8], but were still upfield relative to their original positions in the free state. There were no further proton peak shifts when the concentration of Q[8] was increased above 1.5 eq. The 2D NOE

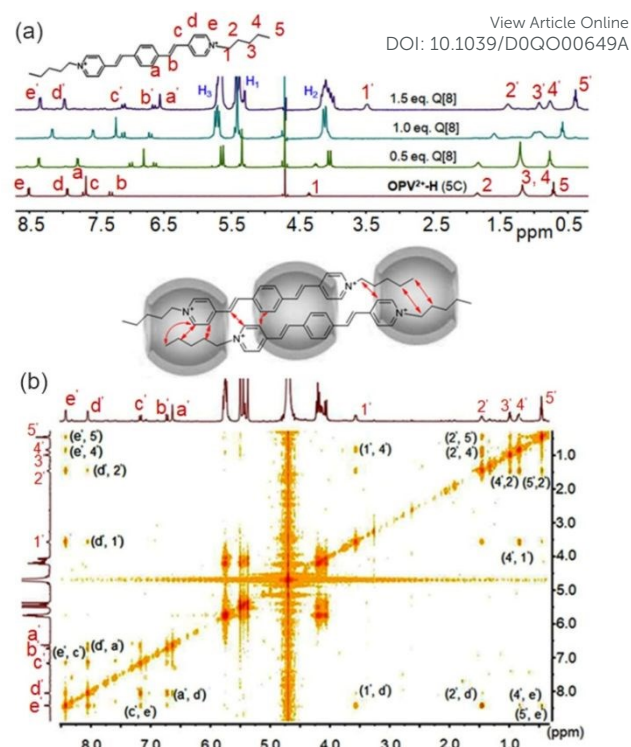


Fig. 4 (a) ¹H NMR spectra of OPV²⁺-H (5C) (1.0 mM in D₂O) with different concentrations of Q[8] (0–1.5 mM); (b) ¹H-¹H NOESY of OPV²⁺-H (5C)/Q[8] at a molar ratio of 2:3.

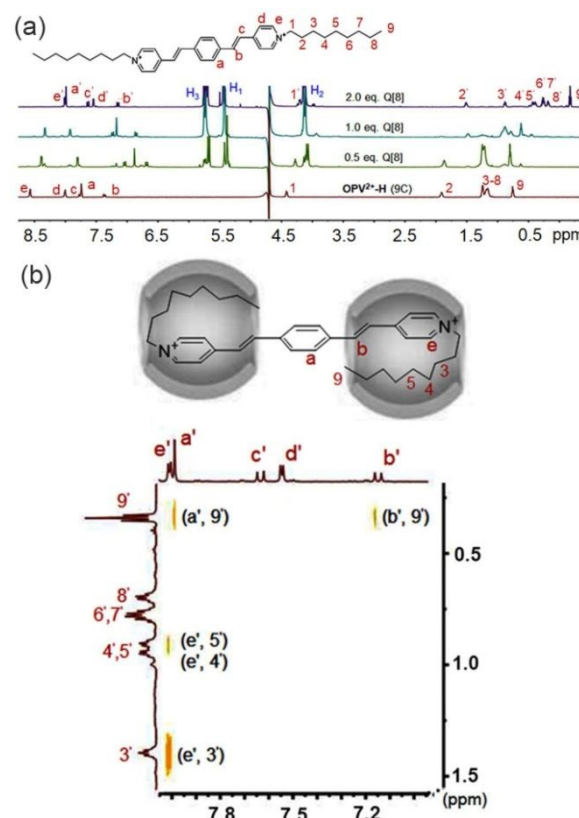


Fig. 5 (a) ¹H NMR spectra of OPV²⁺-H (9C) (0.7 mM in D₂O) with different concentrations of Q[8] (0–1.4 mM); (b) ¹H-¹H NOESY of OPV²⁺-H (9C)/Q[8] at a molar ratio of 1:2.

correlations for **OPV²⁺-H** (5C) ($H_{e^+}-H_c$, $H_{d^+}-H_a$, $H_{e^+}-H_5$, $H_{e^+}-H_4$, $H_{d^+}-H_2$, $H_{d^+}-H_1$, H_1-H_4 , H_2-H_5 , and H_2-H_4), suggested a unique 2:3 assembly of the guests with Q[8]³⁶ (Figs. 4b and S9-10). From a structural viewpoint, this binding mode would strengthen the acceptor-donor-acceptor (A-D-A) electronic interaction of the π -conjugated **OPV²⁺**-derived cationic guest. Especially, the central Q[8] encapsulation of the dimeric guest not only increases the π -surface of the **OPV²⁺** core, but also greatly enhances the electron-donor group promoting the ICT effect of the guest. Therefore, a further red-shifted emission was observed in that host-guest system.

In contrast, in the case of **OPV²⁺-2Me** (5C-6C), it was found that the signals for protons on the ethylene and phenyl groups (including the two Me groups on the phenyl parent) underwent downfield shifts even compared to their original proton peaks in the free guest (Fig. S6), suggesting that the moieties were located at the carbonyl portals of the Q[8] host. The proton signals of the pyridinium and the alkyl chains exhibited remarkable upfield shifts, implying that these groups were deep in the Q[8] cavity. The detailed ¹H NMR titration experiment results revealed that a 1:2 host-guest interaction (Q[8]/guest) was formed in this system as well as with all of the 7C-9C appended guests (Figs. S4-S6).

Interestingly, the NOE correlations (H_a-H_g , $H_{e^+}-H_5$, $H_{e^+}-H_4$, $H_{e^+}-H_3$, and H_b-H_g) from the 1:2 complex of **OPV²⁺-H** (9C) (Fig. 5) provided evidence that the N-substituted alkyl chain was folded into the cavity of Q[8] to form a U-shaped conformation together with partial inclusion the pyridinium moiety. This is slightly different to the observation by Kim,³⁷ where the alkyl chains directly adopted the U-shape themselves in the Q[8] cavity. In addition, for **OPV²⁺-2Me** (7C-9C) guests (Fig. S6), the NMR titration results demonstrated that all the signals of the guest protons were upfield shifted from the initial concentration of Q[8] (below 1.0 eq.). The proton peaks for the Me and parent phenyl moieties were downfield shifted when the concentration of Q[8] was increased from 1.0-2.0 eq. In particular, the continued upfield shifts of the proton signals for the alkyl chains and the pyridinium groups in the NMR spectra suggested that no ternary complexes were formed throughout the Q[8] titration. This result implied that there were no π - π interactions between the **OPV²⁺-2Me** (7C-9C) guests in the Q[8] cavity (Fig. S12), which is consistent with the observation from the fluorescence emission (Fig. 1f). Evidently, the two Me groups on the phenyl of these guests was the key factor in controlling such molecular assembly modes.

Recently, Scherman and co-workers described pioneering work exploring the properties of Q[8]-based complexes by isothermal titration calorimetry (ITC). Their studies revealed that the enthalpies of the 2:2 quaternary complexes of Q[8] with guest were typically below -50 kJ mol⁻¹.²⁴ The large binding enthalpies can be attributed to occupation of the whole Q[8] cavity by guest to release the cavity-bound high energy water.³⁸ In an effort to gain more detailed information on the binding interactions between Q[8] and the **OPV²⁺**-derived guests, ITC studies were also carried out in this work. As shown in Table S4, the obtained enthalpy data from

representative guests were found to range from -95 kJ mol⁻¹ to -50 kJ mol⁻¹, suggesting that the Q[8] cavities were fully occupied by guests, except for **OPV²⁺-Me**(1C)/Q[8] and **OPV²⁺-2Me**(1C)/Q[8] complexes, where the enthalpies were below -40 kJ mol⁻¹. This can be attributed to the Me group and shorter alkyl chain making the guest less included in the Q[8] cavity and leading to less displacement of high energy water molecules from the cavity. Similarly, the much lower binding enthalpies of **OPV²⁺-H** (5C)/Q[8] (-95.12 kJ mol⁻¹) and **OPV²⁺-Me** (5C)/Q[8] (-88.75 kJ mol⁻¹) indicated that a larger number of cavity-bound water molecules were displaced by those novel host-guest binding modes. Therefore, the 2:2, 2:3, and U-shaped 1:2 binding modes of guests with Q[8] were reasonably confirmed by the ITC data and in agreement with the related ¹H NMR titration experiment. Additionally, the binding constants (K_a) for the guests and Q[8] complexes were found to be between 1.0×10^5 M⁻¹ and 9.0×10^6 M⁻¹, indicating strong host-guest binding.

Fluorescence lifetime data systematically determined by time-correlated single photon counting (TCSPC) was fully in line with the above observations. As shown in Table S2, after addition of Q[8] to the relevant guest solution, the lifetime was significantly increased (in the range of 7 to 15 ns), except in the case of the 1:2 complex (guest/Q[8]) and all host-guest interactions of **OPV²⁺-2Me** (7C-9C) where shorter lifetimes were observed. The longest fluorescence lifetimes were observed for the 2:3 complexes of Q[8] with the 5C-6C-appended **OPV²⁺-H** and **OPV²⁺-Me** guests. Essentially, longer lifetimes imply that the π -conjugated **OPV²⁺** core (fluorophore) is deeply encapsulated in the Q[8] cavity, while shorter lifetimes indicate that the fluorophore is less deeply buried or further from the Q[8] cavity.³⁹ Most importantly, the fluorescence lifetime changes were fully in line with the host-guest assembly modes confirmed by the fluorescence and NMR spectra, suggesting that fluorescence decay times may have potential in the characterization of Q[8]-chromophore system binding modes. In particular, the solubility limit of the Q[8] host and the intrinsic hydrophobic effect of the chromophore in water sometimes hinder the performance of the NMR titration experiment. For example, in the present work, the poor solubility of **OPV²⁺-2Me** (7C-9C) in aqueous solution meant that the NMR signal changes were not clear enough to detail the host-guest interactions. The shorter fluorescence lifetimes from these complexes in dilute solution, however, gave strong supplementary evidence to support the host-guest binding modes.

In addition to the above, the Q[8]-derived dynamic host-guest assemblies and the fluorescent signals, such as those of the **OPV²⁺-Me** (5C)/Q[8] complex, can be used as competitive fluorescent indicator displacement chemosensors for the discrimination of spermine and spermidine from other species such as cations, anions and amino acids in pure aqueous solution (Fig. S14). This suggests that Q[8]-based dynamic assemblies and their resulting fluorescent signals have further applications.

Conclusions

In summary, we have successfully prepared a series of Q[8]-derived multiple dynamic host-guest assemblies, including 2:2, 2:3, and 1:2 complexes, that exhibited quantitative fluorescence. These signals were not only sensitively captured in the fluorescence spectra, but could also be visualized directly using the naked eye under UV light. The host-guest binding modes were further characterized by their 1D and 2D NMR spectra. Interestingly, the systematically determined fluorescence lifetimes obtained from the Q[8] complexes indicated that they could be used to help characterize the host-guest binding modes. Most importantly, the Q[8]-based dynamic assemblies induced by N-substituted alkyl chains and subtle modification by Me groups on the OPV²⁺ core had different binding properties according to their HSCT interactions. These dynamic assemblies included the 2:3 quinary complex and the 1:2 interaction-triggered U-shaped conformation have not been reported previously. We expect that the Q[8]-based new binding fashions and the resulting photophysical properties will help to extend application of Q[8] as a macrocyclic host in supramolecular polymers and functional materials.

Conflicts of interest

The authors have no conflicts to declare.

Acknowledgements

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