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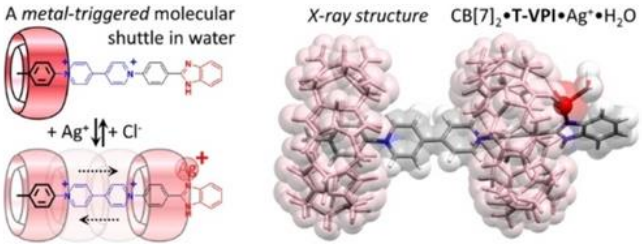
[a] A. Kriat, Dr. D. Gigmes, Dr. A. Kermagoret, Dr. D. Bardelang
Aix Marseille Univ, CNRS, ICR, AMUtech, Marseille, France
E-mail: anthony.kermagoret@univ-amu.fr
E-mail: david.bardelang@univ-amu.fr
[b] Dr. S. Pascal, Dr. O. Siri,
Aix Marseille Univ, CNRS, CINAM, AMUtech, Marseille, France
[c] Dr. B. Kauffmann,
Univ. Bordeaux, CNRS, INSERM, Institut Européen de Chimie et Biologie (UAR3033/US001), 2 rue Escarpit, 33600 Pessac, France
[d] Dr. Y. Ferrand,
CNRS, Bordeaux Institut National Polytechnique, CBMN (UMR 5248), Université de Bordeaux, Institut Européen de Chimie et Biologie, 33600 Pessac, France



Abstract: The structure of the Viologen-Phenylene-Imidazole (**VPI**) guest, previously shown to be bound by cucurbit[7]uril (CB[7]) with binding modes depending on pH and silver ions, has been extended by adding hydrophobic groups on the two extremities of **VPI** before investigations of CB[7] binding by NMR, ITC, X-ray diffraction, UV-vis and fluorescence spectroscopies. With an imidazole station extended by a naphthalene group (**VPI-N**), binding modes of CB[7] are similar to those previously observed. However, with the viologen extended by a tolyl group (**T-VPI**), CB[7] preferentially sits on station **T**, shuttling between the **T** and **P** stations at acid pH or after Ag^+ addition. The CB[7]•**T-VPI** complex thus behaves as a metal-actuated thermodynamic *stop-and-go* molecular shuttle featured by fast and autonomous ring translocation between two stations and a continuum for fractional station occupancy solely and easily controlled by Ag^+ concentration.

Introduction

Since the discovery of molecular shuttles by Stoddart and coworkers in the early 1990s,[1] there have been many reports[2] describing for example restricted ring shuttling,[3] modulation of shuttling speed[4] or tuning of station occupancy.[5] In this field, one could distinguish those for which ring shuttling is reversibly actuated by an external stimulus, in a switch-like manner,[6] from those, often "rapid" for which ring shuttling occurs autonomously between two degenerate or near degenerate stations. However, few have been described, to the best of our knowledge, enabling to control the entry in the shuttle mode (*stop-and-go* molecular shuttles),[7] and the majority of molecular shuttles have been studied in organic solvents. The discovery of cucurbituril macrocycles[8] and especially CB[7] in 2000 (Figure 1)[9] featured by relatively high affinities for guest molecules[10] enabled to prepare several molecular switches[11] and shuttles[12] in water. Yet and again, very few reports are available about cucurbituril based molecular shuttles in water. Kaifer and coworkers, the first, shown in 2006[13] that a "fixed" CB[7] could reversibly enter a fast

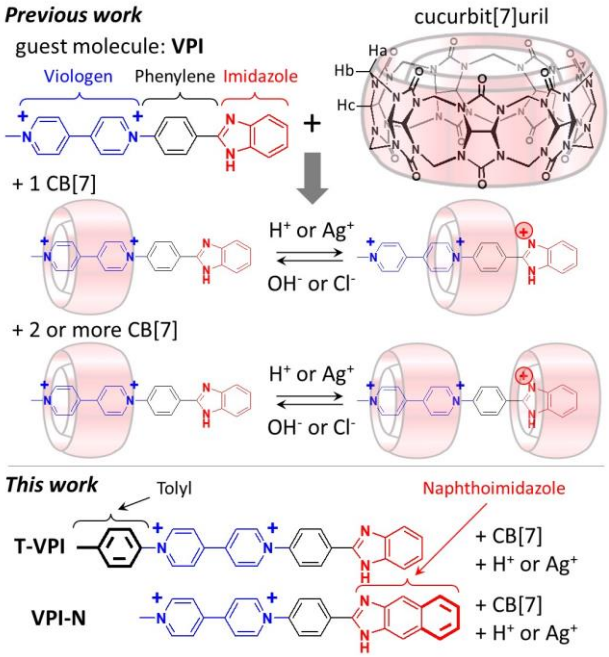


Figure 1. Molecular structures of **VPI** and CB[7] and main binding modes upon addition of H^+ or Ag^+. In this work, we extended the hydrophobic part of the guest on the viologen side by a tolyl group (**T**) or on the imidazole side by a naphthalene (**N**) and studied how this impacted CB[7] binding.

shuttling mode by acidification of a viologen dicarboxylate guest molecule, followed by similar results with a more rigid pseudo-rotaxane.[14] In 2016, Ma and coworkers reported a pH gated CB[7] shuttle with tunable room-temperature phosphorescence.[15] Recently, McClenaghan, Parola, Basilio and coworkers reported a nice example of a CB[7] based molecular shuttle gated by light, pH and temperature.[16] Beside the light-gated system mentioned above raising or lowering a kinetic barrier to exit or enter the shuttling mode, the other examples are based on one (or two) particular station(s) that is (are) rendered much less attractive for the host by a relevant stimulus thereby blocking the ring on another station. Strangely, there is no molecular shuttle in water sensitive to a stimulus improving ring binding on a disfavored station *enough* to reach the affinity of the first station (entry in autonomous shuttling), but *not enough* to trigger full ring translocation (quantitative single shuttling as in a switch).

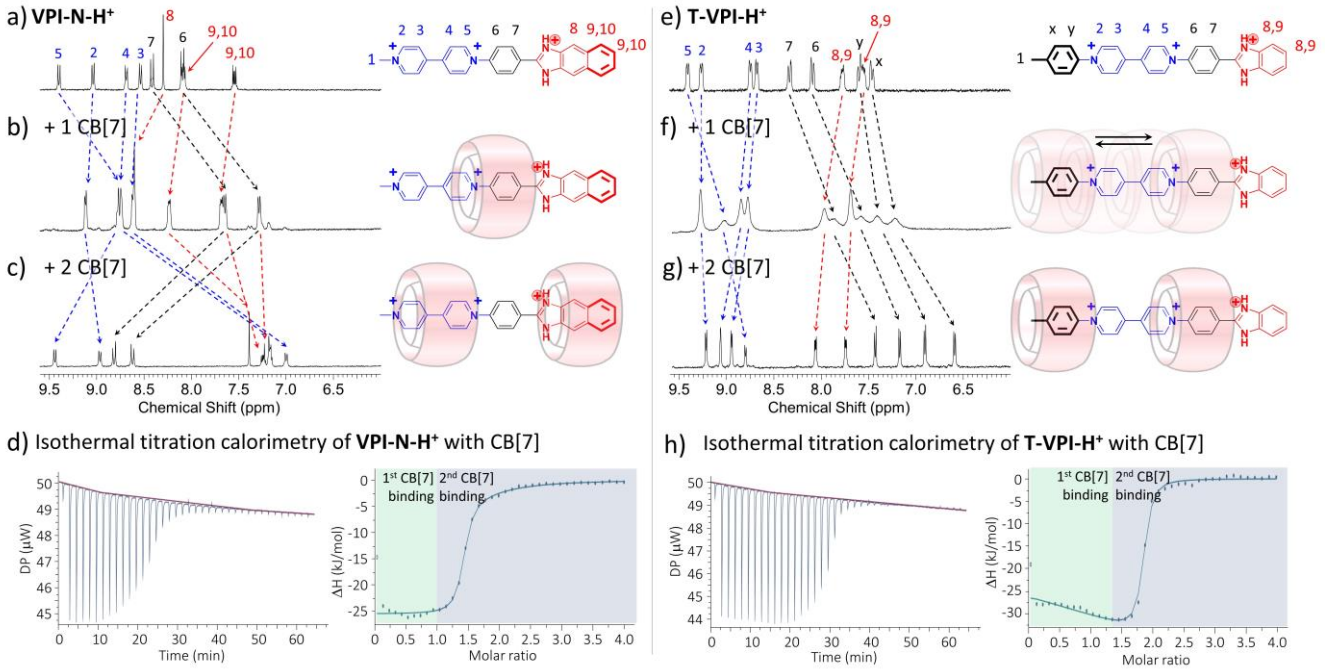


Figure 3. Aromatic part of ^{1}H NMR spectra of **VPI-N-H$^+$** (a) alone, (b) with 1 equiv. of CB[7] and (c) with 5 equiv. of CB[7] at 300 K. Microcalorimetric titration (d) of **VPI-N-H$^+$** with CB[7] in line with two binding events (see text). Aromatic part of ^{1}H NMR spectra of **T-VPI-H$^+$** (e) alone, (f) with 1 equiv. of CB[7] and (g) with 2.5 equiv. of CB[7] at 300 K. Microcalorimetric titration (h) of **T-VPI-H$^+$** with CB[7] in line with two binding events (see text).

Recently, we reported the pH-responsive viologen derivative **VPI** because of the three different stations CB[7] can dock on (Figure 1).[17] While CB[8] formed a host•guest 2:2 complex with this guest,[18] and CB[10], a host•guest 2:3 complex,[19] results were very different with CB[7]. At one equivalent of host, with H^+ or Ag^+ cations, CB[7] quantitatively moved from station **V** to station **P** affording a pH- or metal-triggered ring translocation switch in water.[17, 20] Fostered by these findings, we wondered how structural extensions by hydrophobic fragments would impact CB[7] binding considering that the action of chemical stimuli (H^+ or Ag^+) was expected to modulate the outcome. We thus prepared two new compounds, extended on the imidazole side by a naphthyl group (**VPI-N**) and on the viologen side by a tolyl group (**T-VPI**) before investigating their interactions with CB[7] and the impact of pH and $AgNO_3$ on the binding modes. Finally, we found that H^+ or Ag^+ cations could enable the ring of a new host•guest complex to enter a fast autonomous shuttling mode by improving binding on a previously disfavored station resulting in a new kind of thermodynamic molecular shuttle in water.

Results and Discussion

Synthesis. The synthesis of **VPI-N** is largely inspired from that used for **VPI**,[20] and the preparation of **T-VPI** relies on well-known Zincke chemistry. Details of these syntheses and characterizations are described in supporting information.

CB[7] binding on VPI-N and T-VPI. The ^{1}H NMR spectrum of **VPI-N** in D_2O (Figure 2a) is featured by chemical shifts of all aromatic resonances that are upfield shifted (7.1-8.8 ppm range) compared to analogous signals of **VPI** (7.3-9.4 pm range)[17] suggesting possible aggregation but the signals remaining sharp, we assigned this spectrum to the possible formation of a **VPI-N** dimer in solution at 1 mM.[21] In the presence of 3 equiv. of TFA (trifluoroacetic acid, *vide infra* Figure 3a), resonances in the aromatic region shifted downfield (7.5-9.4 ppm range) matching with the range of chemical shifts observed for **VPI** or **VPI**-H$^+$,[17] each signal closely positioned with respect to those of analogous protons of **VPI** and thereby suggesting that **VPI-N-H$^+$** is present as a monomer in solution. As for **VPI**, when one equiv. of CB[7] was added in a solution of **VPI-N**, signals corresponding to protons H_3 and H_4 are shifted upfield by 1.0 ppm (Figure 2b) in line with viologen complexation.[20] However and contrary to the situation with **VPI**,[17, 20] a second binding is possible, on station **N** (Figure 2c) but requires an excess of CB[7]. In parallel of NMR studies, we investigated CB[7] binding by isothermal titration calorimetry (ITC) and observed an unusual V-shape thermogram for the titration of **VPI-N** with CB[7] (Figure 2d). Indeed, the ΔH values between 0 and 1 equiv. of host decreased before increasing after 1 equiv. of host. This behavior could be due to a dissociation of **VPI-N** dimers during an initial CB[7] binding, before a second complexation occurs with a larger enthalpic contribution after addition of more than 1 equiv. of host. Considering a 2:1 binding model, a good fit with experimental points was obtained affording stepwise binding constants $K_1 = 2.91 \times 10^5$ M^{-1} and $K_2 = 2.86 \times 10^3$ M^{-1}. The K_1 value for binding of CB[7] on the viologen station is close to the one determined for **VPI** by ITC for the same binding ($K = 7.25 \times 10^5$ M^{-1}).[20] The relatively small value of K_2 for binding of the second CB[7] on station **N** is consistent with the need of several equivalents of host to saturate this station as monitored by NMR (Figure 2c). On the other hand, CB[7] behaved differently in the presence of **T-VPI**. Contrary to **VPI** or **VPI-N**, at 1 equiv. of host, CB[7] prefers sitting on station **T** of **T-VPI**, next to the viologen, as sometimes observed on viologen derivatives.[12e, 22] Indeed, 0.8 ppm upfield shifts of signals corresponding to protons Hx and Hy (Figure 2f) are assigned to the presence of CB[7] on station **T**. As for **VPI-N** though, a slight excess of host triggered a second binding of CB[7] but this time on station **P** as suggested by the upfield shifted signals of protons H_6 (1.1 ppm) and H_7 (0.8 ppm, Figure 2g). Titrations by ITC of

Table 1. Binding constants and thermodynamic parameters obtained by Isothermal Titration Calorimetry in water.[a]

Species	K_1/M^{-1}	ΔH_1/kJ.mol^{-1}	$-T\Delta S_1$/kJ.mol^{-1}	K_2/M^{-1}	ΔH_2/kJ.mol^{-1}	$-T\Delta S_2$/kJ.mol^{-1}
CB[7]•**VPI-N**	2.91 (±0.61) × 10^5	-6.2 (± 0.6)	-24.9 (± 1.1)	2.86 (±0.21) × 10^3	-45.6 (± 5.1)	+25.9 (± 5.3)
CB[7]•**T-VPI**	5.85 (±0.14) × 10^7	-25.1 (± 0.1)	-19.2 (± 0.2)	1.59 (±0.09) × 10^5	-21.6 (± 0.1)	-8.1 (± 0.2)
CB[7]•**VPI-N** (acid pH)	4.86 (±0.10) × 10^6	-25.6 (± 0.2)	-12.6 (± 0.3)	2.10 (±0.15) × 10^4	-4.3 (± 0.4)	-20.4 (± 0.6)
CB[7]•**T-VPI** (acid pH)	4.54 (±0.21) × 10^6	-11.3 (± 4.2)	-26.8 (± 4.3)	3.16 (±0.06) × 10^6	-54.3 (± 5.1)	+17.2 (± 5.2)

[a] Binding model: two sets of sites.

solutions of CB[7] in solutions of **T-VPI** (Figure 2h) afforded thermograms consistent with a stepwise binding of 2 CB[7] and binding constants K_1 = 5.85 × 10^7 M^{-1} and K_2 = 1.59 × 10^5 M^{-1}. These values are close to those determined by competition NMR K_1 = 3.2 × 10^8 M^{-1} (with respect to phenyl-trimethylammonium, Figure S31) and K_2 = 2.9 × 10^5 M^{-1} (compared to 1-ethyl-3-methylimidazolium, Figure S32). These results are consistent with NMR data which suggested a quantitative binding of 1 equiv. of host on station **T**, and an about quantitative binding (requiring only a slight excess of host) on station **P**.

CB[7] binding on VPI-N and T-VPI at acid pH. As for **VPI**,[17] when TFA was added on a stoichiometric solution of **VPI-N** and CB[7], the CB[7] host moved on station **P** (Figure 3b) as reflected by marked upfield shifts for signals of H_5 ($\Delta\delta$ = 0.25 ppm), H_6 ($\Delta\delta$ = 0.8 ppm), and H_7 ($\Delta\delta$ = 0.7 ppm). And as for **VPI** again, when 2 equiv. of host were used in the presence of TFA, there are 2 CB[7] sitting on the viologen and imidazole stations as shown by the diagnostic upfield shifts of signals of protons H_3 ($\Delta\delta$ = 1.5 ppm), H_4 ($\Delta\delta$ = 1.5 ppm), H_8 ($\Delta\delta$ = 0.8 ppm), H_9 ($\Delta\delta$ = 0.8 ppm), and H_{10} ($\Delta\delta$ = 0.25 ppm, Figure 3c). We next recorded ITC thermograms for the titration of CB[7] with **VPI-N** at acid pH which confirmed these results (Figure 3d). Simulations according to a 2:1 binding mode afforded a good match with experimental points and binding constants K_1 = 4.85 × 10^6 M^{-1} and K_2 = 2.10 × 10^4 M^{-1}. This K_1 value, higher than the one recorded in water at neutral pH (2.91 × 10^5 M^{-1}) is consistent with ring translocation on station **P** (Figure 3b) and also with the one recorded previously for binding of CB[7] on station **P** of **VPI** with Ag^+ (6.45 × 10^6 M^{-1}),[20] or H^+ (10.4 × 10^6 M^{-1}).[17] The K_2 value is also higher at acid pH (2.10 × 10^4 M^{-1}) than the one at neutral pH (2.86 × 10^3 M^{-1}) reflecting better binding on station **N** when the imidazole is protonated. A large range of metal ions have previously been tested with **VPI**.[20] As we found that only Ag^+ was impacting NMR spectra, we focused on this cation with the new guest molecules. With this stimulus, the results are very similar to those obtained with TFA (see supporting information). Therefore, the extension of the benzimidazole into a naphthimidazole did not change much the way CB[7] binds **VPI-N**, H^+ and Ag^+ cations providing results consistent with those previously obtained with **VPI**.

We then tested the influence of pH and added TFA to solutions of **T-VPI** containing 1 or 2 equiv. of CB[7]. With one host molecule, the ^{1}H NMR spectrum is featured by broad peaks (Figure 3f), the lack of resonance in the region corresponding to the signals of the phenylene possibly indicating that their position has been shifted upfield (other resonances at down field correspond to signals of viologen protons). The absence of sharp peaks also suggested that CB[7] could shuttle between at least 2 stations. Station **T** being largely preferred at neutral pH and station **P** being one of possible stations at acid pH (as for **VPI**), we hypothesized a shuttling of CB[7] between the **T** and **P** stations caused by protonation of the imidazole group of **T-VPI**. This is in line with previous reports on structurally similar guest compounds for CB[7].[17] With two CB[7] in an acidic solution of **T-VPI**, the ^{1}H NMR spectrum (Figure 3g) showed sharp peaks in line with tolyl and phenylene complexation, thereby supporting the previous hypothesis.

Indeed, upfield shifts of 0.8 and 1.1 ppm for signals of H_6 and H_7, respectively (station **P**), and of 0.7 and 0.9 ppm for signals of Hx and Hy, respectively (station **T**), are in line with 2 CB[7] hosts firmly sitting on stations **T** and **P**. This shows that once protonated, **T-VPI-H$^+$** has a clear preference to host CB[7] molecules on the stations **T** and **P** with a similar affinity. As previously done, we performed ITC titrations, this time of CB[7] in solutions of **T-VPI** at acid pH and recorded thermograms in line with double host binding. Simulations according to a 2:1 binding mode afforded a good fit with experimental points and similar binding constants K_1 = 4.54 × 10^6 M^{-1} and K_2 = 3.16 × 10^6 M^{-1}, supporting the proposed shuttling of CB[7] between two stations of **T-VPI-H$^+$**. Results obtained by ITC are summarized in table 1.

As for **VPI-N**, the addition of $AgNO_3$ instead of TFA in solutions of **T-VPI** with CB[7] showed pretty similar ^{1}H NMR spectra leading to similar conclusions (see supporting information). Therefore, adding another (tolyl) binding site on the viologen resulted in different CB[7] binding modes, the first CB[7] residing on station **T**, the second on station **P** showcasing a shifting in CB[7] docking from stations **V**+**I** (for **VPI**) to stations **T**+**P** (for **T-VPI)**. The matching of affinities between stations **T** and **P** (after guest protonation) afforded a new molecular shuttle with fast autonomous ring translocation in water due to the near degeneracy of binding sites for CB[7] on **T-VPI-H$^+$** and Brownian motion,[23] temperature only marginally impacting host shuttling (288-348 K, Figure S43).

A short read at Table 1 shows large differences in enthalpic and entropic behaviour. For the CB[7]•**VPI-N** system, the first binding on station **V** is largely entropy-driven with an unusually small enthalpic value of -6.2 kJ.mol^{-1} ($\Delta H_{CB[7]\bullet viologen}$ = -16.7 kJ.mol^{-1},[24] $\Delta H_{CB[7]\bullet \mathbf{VPI}}$= -14.7 kJ.mol^{-1})[20] which may originate from the (necessary) dissociation of the **VPI-N** dimer (energy penalty) before CB[7] can bind **VPI-N**. The second binding on uncharged station **N** is enthalpy-driven with a ΔH_2 value of -45.6 kJ.mol^{-1} and is also featured by a large unfavourable entropy change. This trend is similar to the one observed for the binding of neutral guests, such as 3-diazabicycloc[2.2.2]oct-2-ene or cyclopentanone,[25] by CB[7] and explained by the release of high-energy water molecules from the cavity of the host.[26] This non-classical hydrophobic effect ($-T\Delta S_2 > 0$) suggests that an entropic contribution, resulting from the release of water molecules from the naphthalene moiety to bulk water during complexation of station **N**, is significant.[27, 25]

For **VPI-N-H$^+$**, this is the opposite. The first binding on station **P** is enthalpy-driven (ΔH_1 = -25.6 kJ.mol^{-1}), but the second

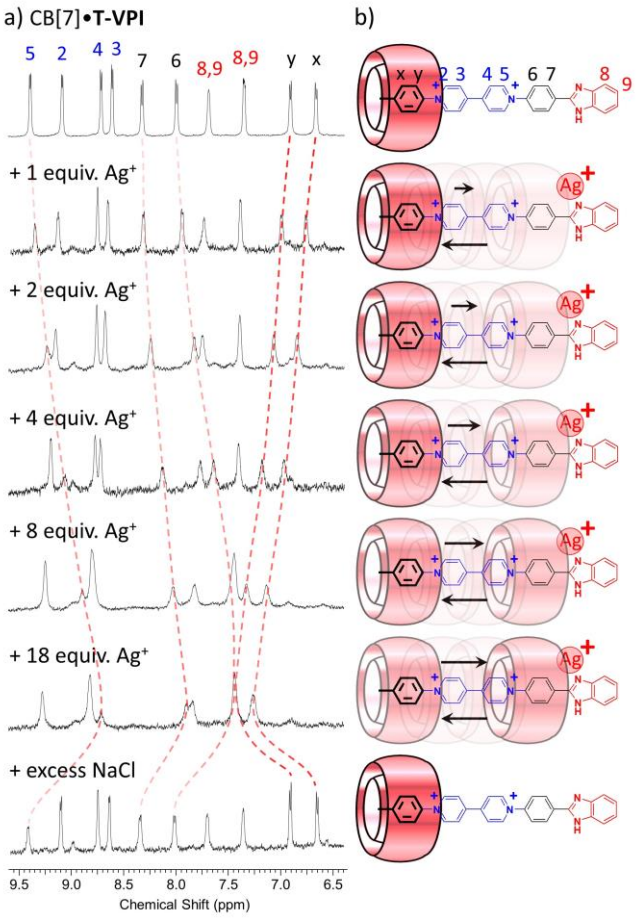


Figure 4. (a) Aromatic region of the 500 MHz ^{1}H NMR spectrum in D_2O (300 K) of the CB[7]•**T-VPI** complex with gradual addition of Ag^+ cations and (b) proposed gradual shuttling from no shuttle (top), CB[7] residing on station **T** up to rapid shuttling of the ring over the two **T** and **P** stations (bottom).

binding is entropy-driven ($-T\Delta S_2$ = -20.4 kJ.mol^{-1}) with low ΔH_2 (-4.3 kJ.mol^{-1}), perhaps due to the energetic penalty of relocating the first CB[7] on station **V** to allow the second CB[7] to dock on station **N** (or the opposite).

Without dimerization or host relocation for **T-VPI**, the two CB[7] bindings at neutral pH are both enthalpy-driven (ΔH_1 = -25.1 kJ.mol^{-1} and ΔH_2 = -21.6 kJ.mol^{-1}) with a high total Gibbs enthalpy change ($\Delta G°$ = -74 kJ.mol^{-1}). The similarity of thermodynamic signatures for CB[7] binding on station **P** for (i) the CB[7]•**T-VPI** system and (ii) the CB[7]•**VPI-N-H**$^+$ system is consistent (~70% $\Delta H°$ driven and ~30% $\Delta S°$ driven) in spite of the different protonated state of the guest. However, at acid pH, the two bindings (K_1 and K_2) are equal in magnitude but the first (binding and shuttling) is entropy-driven ($-T\Delta S_1$ = -26.8 kJ.mol^{-1}) with a small value of ΔH_1 = -11.3 kJ.mol^{-1}, while the second is largely enthalpy-driven (ΔH_2 = -54.3 kJ.mol^{-1}) and entropy compensated ($-T\Delta S_2$ = +17.2 kJ.mol^{-1}). This may reflect some energy penalty at precisely locating a host which prefers shuttling (small ΔH_1 value), before the two CB[7] are fixed on stations **T** and **P**. Or, the CB[7] shuttling (non-specific interaction due to the energetic equivalence of binding sites **T** and **P**) could perturb guest solvation, enough to make the axle resembling an hydrophobic one since the thermodynamic signature ($\Delta H << 0$, $\Delta S > 0$) looks like the one for pure hydrophobic compounds.[25] Unexpectedly, the thermodynamic behaviour is similar to the one observed for CB[7] binding on **VPI-N** even if the complexes involved are different.

A metal-actuated molecular shuttle. During our investigation of the CB[7]•**T-VPI** complex, and more specifically the molecular shuttle with excess H^+, we wondered if we could progressively make the CB[7] ring shuttling from the **T** to the **P** station with Ag^+. Recording ^{1}H NMR spectra of the 1:1 CB[7]•**T-VPI** complex with gradual addition of Ag^+ showed it is indeed the case (Figure 4) with progressive deshielding of signals corresponding to Hx and Hy protons suggesting that the **T** station is less and less occupied while at the same time, signals of protons 5, 6, and 7 progressively shifted upfield (CB[7] on station **P**). Variable-temperature (VT) NMR (Figure S56) did not allow to observe coalescence that would have enable to determine activation energies and rates of exchange. However, the titration with increasing concentrations of Ag^+ unambiguously showed the autonomous and gradual, cation-triggered ring shuttling between the **T** and **P** stations.

X-ray diffraction. Single crystals of a host:guest 2:1 complex with silver could be grown from an aqueous solution of **T-VPI** with two CB[7] and excess $AgNO_3$ (^{1}H NMR spectrum, Figure 5a). The single-crystal structure (Figure 5b, CCDC number 2217466) confirmed the position of the first CB[7] on station **T** and that of the second CB[7] on station **P**. Beside the host:guest arrangement, one silver atom was found close to the imidazole function of **T-VPI** (Ag-N distance of 2.24 Å). The silver atom also interacts with two oxygen atoms of one carbonyl rim of the CB[7] ring docked on station **P** (Ag-O distances of 2.33 and 2.70 Å). The coordination sphere of silver is completed by one ordered water molecule (Ag-O distance of 2.42 Å). The structure of this $CB[7]_2$•**T-VPI**•Ag^+ complex confirms the previously postulated type of interactions for Ag^+ ions with imidazole containing guest molecules and CB[7].[20] Indeed, for **VPI**, CB[7] was postulated to be bound on station **P** with one silver ion

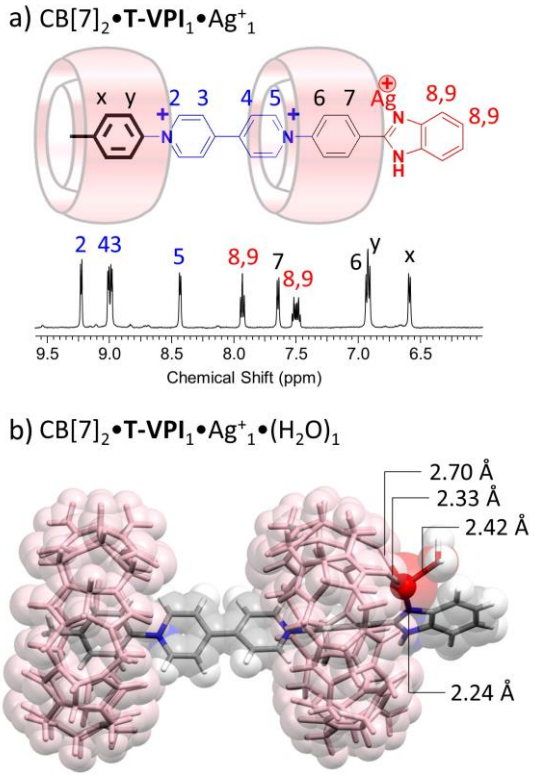


Figure 5. (a) Aromatic region of the 600 MHz ^{1}H NMR spectrum in D_2O (300 K) of the $CB[7]_2$•**T-VPI**• Ag^+ complex and (b) X-ray structure of the same complex.

coordinated by both, the guest by the imidazole nitrogen atom, and the host by two oxygen atoms of the closest carbonyl rim.[20] Consistent with NMR results, the two CB[7] rings are found sitting on the **T** and **P** stations.
This type of double host binding implying *separated stations* (**V**+**I** or **T**+**P**) observed with **VPI**, **VPI-N** and **T-VPI** is most likely due to the impossibility for two CB[7] to sit simultaneously on adjacent stations.

UV-visible absorption and fluorescence spectroscopies. Contrary to **VPI** which is weakly or not fluorescent, **VPI-T** is emissive in solution which incited us to investigate the formation of these complexes by absorption and fluorescence spectroscopies (Figure 6 and supporting information). First, the comparison of the UV-vis spectra in water solution showed no important shift of absorption band upon addition of CB[7] for **T-VPI** and the corresponding protonated or coordinated species. Only a slight hypsochromic shift of the lower energy band was noticed for **T-VPI** in presence of 2 CB[7], presumably due to the presence of a macrocycle on the **P** station, interacting with the benzimidazole ring. Second, regarding the emission properties, **T-VPI** displays a weak fluorescence in the violet-blue range, peaking at 380 nm. The binding of a first CB[7] unit does not shift the band, nevertheless, a second equivalent triggers a bathochromic shift of the emission band toward 398 nm. This observation is also consistent with the presence of a CB[7] at the proximity of the imidazole.
In the case of **T-VPI-H$^+$**, a small redshift can be noticed upon addition of 1 equiv. of CB[7], which presumably reveals that the single host is shuttling between the **T** and **P** stations, in contrast with **T-VPI** that lacks such redshift. In the case of **T-VPI•**Ag$^+$, the progressive bathochromic shift of the emission in presence of CB[7] also suggests that a comparable assumption can be formulated, but care has to be taken since the emissions are sensibly quenched by the coordination to silver, as can be noticed by the lower signal-to-noise ratio recorded.

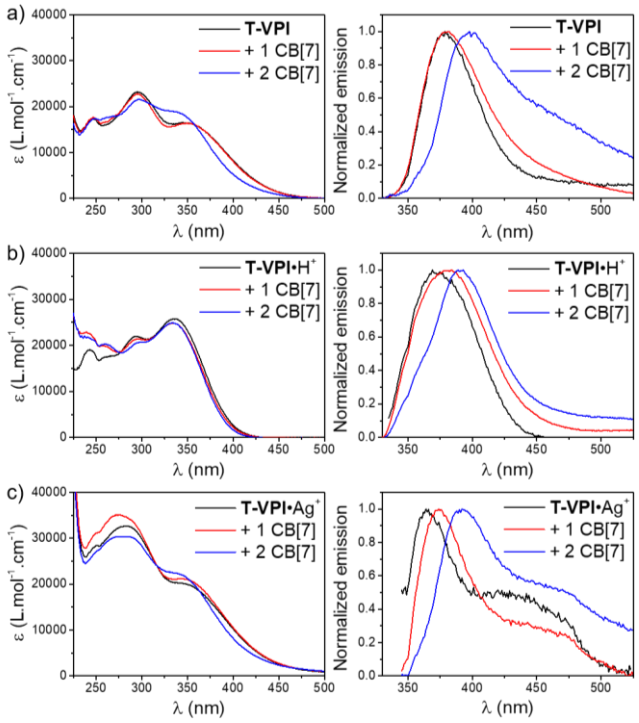


Figure 6. Electronic absorption (left) and fluorescence (right) spectra of **T-VPI** (10^{-5} M in H_2O) alone and with 1 or 2 CB[7] (a) at neutral pH, (b) at acid pH and (c) in the presence of 10 equiv. of Ag$^+$ ions (λ_{exc} = 290 nm).

Discussion. While binding of CB[7] on hydrophobic groups next to viologens (i. e. tolyl, phenylene) has been observed (in agreement with many previous reports),[13, 12e, 28, 14, 22] this trend remains difficult to explain. A possible reason may be that charge assisted N$^+$C-H•••O=C hydrogen bonds between the guest and the carbonyl crowns of the host may be stronger (more directional) when CB[7] is on the tolyl than when it is on the viologen for which such interactions would become more orthogonal so less stabilizing. At one equiv. of CB[7], H$^+$ or Ag$^+$ stimuli actuate CB[7] translocation from station **V** to station **P** (for **VPI** and **VPI-N**) or activate ring shuttling between stations **T** and **P** (for **T-VPI**). However, with 2 equiv. of host in solution, one CB[7] is always bound (either on station **V** or station **T**) before capture of the second CB[7], only at excess host, or when H$^+$ or Ag$^+$ cations are added, improving affinity of the second host for relevant stations. Even if representations of the CB[7] position along the guests are simplified for the sake of clarity throughout the paper, the atomic representation provided by the crystal structure affords more precise insights about CB[7] positioning. The unique interplay between Ag$^+$ (or H$^+$), CB[7] and **VPI** derivatives is probably, at least in part, responsible for the results observed, this kind of cooperative interaction being able to (i) actuate ring translocation, (ii) trigger shuttling between two stations or (iii) capture another macrocycle over a linear track. Figure 7 summarizes the supramolecular complexes discovered and the binding modes of CB[7] whether H$^+$ or Ag$^+$ ions are present or not. Hence, compared to previous results obtained with **VPI**, we found that hydrophobic extension by a naphthyl group on the imidazole did not change much the outcome regarding CB[7] binding with H$^+$ or Ag$^+$. The main difference being the possibility to complex the **N** station at high excess host, without H$^+$ or Ag$^+$. However, the introduction of a tolyl group on the other side (on the viologen) changed the site of the first binding to station **T**, created a shuttling motion in the presence of H$^+$ or

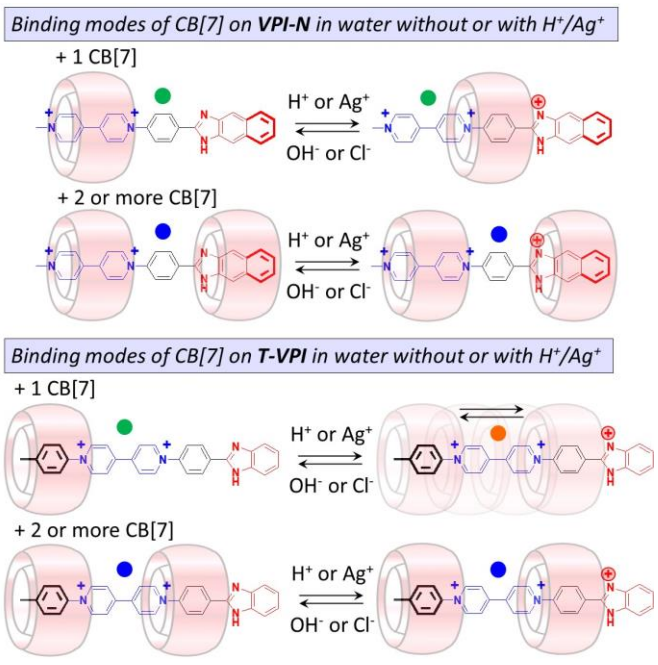


Figure 7. New supramolecular complexes obtained with **VPI-N** and with **T-VPI** in the presence of CB[7], without (left column) and with (right column) H$^+$ or Ag$^+$ stimuli (●: 1:1 complexes, ●: 2:1 complexes, ●: molecular shuttle).

Ag^+, and shifted the previously seen double binding from the **V**+**I** stations to the **T**+**P** stations. The serendipitous discovery of a metal-actuated molecular shuttle shows that other stimuli (here a metallic cation) can be used to enter the autonomous shuttle mode. This is possible because the asymmetric guest molecule **T-VPI** possesses one tolyl favoured station for CB[7] and another station becoming about as attractive for CB[7] as the tolyl after addition of H^+ (quantitative binding of H^+) or Ag^+ (equilibrated Ag^+:imidazole:CB[7] binding). Interestingly, there is a continuum accessible in the "degree of shuttling" due to the lability of the tridentate Ag^+•••N(guest)•••O_2(CB[7]) interaction. Indeed, one can continuously manipulate the degree of shuttling by the simple control of the quantity of Ag^+ added from the "stop state" to the about equally populated "autonomous shuttling state" with all intermediate situations.

Conclusions

In conclusion, we have explored how hydrophobic extensions on either end of a viologen-phenylene-imidazole (**VPI**) guest molecule impact CB[7] binding. Replacement of the initial benzimidazole unit by a naphthimidazole part (**VPI-N**) resulted in similar binding behaviours for CB[7], the only noticeable difference is the possibility for a second binding on the newly formed **N** station at excess host concentration. However, introduction of a tolyl group on the viologen side (**T-VPI**) showed marked changes, CB[7] binding first on the tolyl group (station **T**) followed by binding of a second CB[7] on the phenylene (station **P**). At acid pH, ring translocation or ring capture is observed for **VPI-N** (as for **VPI**) but for **T-VPI**, shuttling or second binding promotion is observed. For the three guest molecules, the addition of Ag^+ ions led to the same complexation modes with CB[7] as with H^+, but required excess metal ion where H^+ could be used stoichiometrically. Beside NMR enabling to propose binding modes of CB[7], ITC allowed to quantify the affinity of the hosts for precise stations of the guests confirming double host binding and strongly supporting the proposed shuttling of one CB[7] on protonated **T-VPI**. Finally, the single-crystal structure of the $CB[7]_2$•**T-VPI**•Ag^+ complex allowed to confirm the binding mode of silver, linking simultaneously the host and the guest to stabilize a CB[7] on the best station.

Experimental Section

General. Chemicals and solvents were purchased from commercial sources (Aldrich, Acros, ABCR or TCI) and used without further purification. CB[7] was prepared according to a previous paper[29] with extra caution regarding possible remaining acid traces. NMR measurements were recorded on Bruker AVL 300, 400, 500 or 600 spectrometers (see supporting information). When necessary for D_2O solutions, acetone was used as internal reference (2.22 ppm). CHN elemental analyses were performed using a Flash EA analyser, 1112 series Thermo Finnigan driven by the Eager 300 software (see supporting information). Isothermal Titration Calorimetry (ITC) experiments were performed on a Malvern MicroCal PEAQ-ITC titrating aqueous solutions of neutral CB[7] at 2 mM into aqueous solutions of guests (100 or 150 μM). For experiments at acid pH, solutions of CB[7] (2 mM) in aqueous HCl (0.5 mM) were titrated into solutions of guests (0.1 mM) in aqueous HCl (0.5 mM) carefully checking the pH was done decreasing below 2. Results were auto-analyzed using the Malvern MicroCal PEAQ-ITC Analysis Software. UV-visible absorption spectra were recorded in spectrophotometric grade water (ca. 10^{-5} M) on a VARIAN CARY 50 SCAN spectrophotometer at room temperature with a 300 nm/min scan rate. Emission spectra were measured using a Horiba-Jobin Yvon Fluorolog-3 spectrofluorimeter equipped with a three-slit double-grating excitation and a spectrograph emission mono-chromator with dispersions of 2.1 $nm.mm^{-1}$ (1200 grooves per mm). A 450 W xenon continuous wave lamp provided excitation. The luminescence of diluted solutions was detected at right angle using 10 mm quartz cuvettes.

Synthesis. The syntheses of **VPI-N** and **T-VPI** are detailed in supporting information. All products were characterized by ^1H and ^{13}C NMR (see supporting information, Figures S1 to S19). A counter anion metathesis was performed on **VPI-N** using a NO_3^- ion-exchange resin (see supporting information).

^1H NMR of **VPI-N** (500 MHz, D_2O, 3.3 10^{-4} M) δ 8.95 (d, *J* = 5.3 Hz, 2H, ***H5***), 8.76 (d, *J* = 5.9 Hz, 2H, ***H2***), 8.19 (d, *J* = 5.4 Hz, 2H ***H4***), 8.13 (d, *J* = 4.6 Hz, 2H, ***H3***), 8.08 (d, *J* = 8.0 Hz, 2H, ***H7***), 7.90 (br s, 2H, ***H8***), 7.73 (br s, 2H, ***H9*** or ***H10***), 7.67 (d, *J* = 8.2 Hz, 1H, ***H6***), 7.21 (dd, *J* = 6.5, Hz *J* = 3.0 Hz, 2H, ***H9*** or ***H10***), 4.31 (s, 3H, ***H1***). Figure S7.
^1H NMR of **T-VPI** (300 MHz, D_2O) δ 9.38 (d, *J* = 7.1 Hz, 2H, ***H5***), 9.32 (d, *J* = 7.1 Hz, 2H, ***H2***), 8.74 (d, *J* = 7.0 Hz, 2H, ***H4***), 8.69 (d, *J* = 7.0 Hz, 2H, ***H3***), 8.31 (d, *J* = 8.8 Hz, 2H, ***H7***), 7.96 (d, *J* = 8.8 Hz, 2H, ***H6***), 7.68 (dd, *J* = 5.8, 2.8 Hz, 4H, overlapped signals of ***Hy*** and ***H8***), 7.57 (d, *J* = 8.2 Hz, 2H, ***Hx***), 7.32 (dd, *J* = 6.1, 3.2 Hz, 2H, ***H9***), 2.47 (s, 3H, ***H1***). Figure S15.

NMR of VPI-N with CB[7]: 1:1 complex. A 1 mM solution of CB[7]•**VPI-N** was prepared by addition of 125 μL of a 4 mM solution of **VPI-N** in D_2O, 100 μL of 5 mM solution of CB[7] in D_2O and 275 μL of D_2O. ^1H NMR of CB[7]•**VPI-N** (300 MHz, D_2O, 1 mM) δ 9.38 (d, *J* = 6.4 Hz, 2H, ***H5***), 9.02 (d, *J* = 6.5 Hz, 2H, ***H2***), 8.48 (d, *J* = 8.7 Hz, 2H, ***H7*** or ***H6***), 8.38 (d, *J* = 8.7 Hz, 2H, ***H7*** or ***H6***), 8.19 (s, 2H, ***H8***), 8.06 (br s, 2H, ***H9*** or ***H10***), 7.51 (br s, 2H, ***H9*** or ***H10***), 7.21 (d, *J* = 6.4 Hz, 2H, ***H4***), 7.08 (d, *J* = 6.4 Hz, 2H, ***H3***), 5.73 (d, *J* = 15.4 Hz, 14H, CB[7]), 5.54 (s, 14H, CB[7]), 4.25 (d, *J* = 15.5 Hz, 14H, CB[7]), 2.22 (acetone, ref). Figure S20.
2:1 complex. A 0.2 mM solution of CB[7]•**VPI-N** was prepared by addition of 25 μL of a 4 mM solution of **VPI-N** in D_2O, 20 μL of a 5 mM solution of CB[7] in D_2O and 455 μL of D_2O; then were added successively 4, 16, 60 and 100 μL of a 5 mM solution of CB[7] in D_2O to prepare solutions presenting **VPI-N**/CB[7] ratios of 1/1.2, 1/2, 1/5 and 1/10, respectively (Figure S22). NMR of **VPI-N** + 5 equiv CB[7]: ^1H NMR (500 MHz, D_2O) δ 9.49 (d, *J* = 5.7 Hz, 2H, ***H5***), 8.99 (d, *J* = 5.7 Hz, 2H, ***H2***), 8.79 (br s, 2H, ***H7*** or ***H6***), 8.57 (br s, 2H, ***H7*** or ***H6***), 7.22 (d, *J* = 6.4 Hz, 2H, ***H4***), 7.14 (br m, 6H, 2H, overlapped signals of ***H8***, ***H9*** and ***H10***), 7.06 (d, *J* = 5.7 Hz, 2H, ***H3***), 5.90 – 5.60 (m, 75H, CB[7]), 5.60 – 5.36 (m, 74H, CB[7]), 4.34 – 4.03 (m, 74H, CB[7]). Figure S23.
NMR of T-VPI with CB[7]: 1:1 complex. A 0.5 mM solution of CB[7]•**T-VPI** was prepared by addition of 125 μL of a 2 mM solution of **T-VPI** in D_2O (2.5 10^{-7} mol), 50 μL of 5 mM solution of CB[7] in D_2O (2.5 10^{-7} mol) and 325 μL of D_2O. ^1H NMR of CB[7]•**T-VPI** (500 MHz, D_2O, 0.5 mM) δ 9.49 (d, *J* = 6.4 Hz, 2H, ***H5***), 9.15 (d, *J* = 6.4 Hz, 2H, ***H2***), 8.79 (d, *J* = 4.7 Hz, 2H, ***H4***), 8.68 (d, *J* = 6.2 Hz, 2H, ***H3***), 8.43 (d, *J* = 8.4 Hz, 2H, ***H7***), 8.10 (d, *J* = 8.4 Hz, 2H, ***H6***), 7.78 (br s, 2H, ***H8*** or ***H9***), 7.44 (dd, *J* = 5.9, 3.1 Hz, 2H, ***H8*** or ***H9***)), 6.99 (d, *J* = 8.0 Hz, 2H, ***Hy***), 6.75 (d, *J* = 8.0 Hz, 2H, ***Hx***), 5.80 (two d, *J* = 15.4 Hz, 14H, CB[7]), 5.50 (s, 14H, CB[7]), 4.21 (app dd, *J* = 15.4, 9.2 Hz, 14H, CB[7]), 2.22 (acetone, ref), 2.09 (br s, 3H, ***H1***). Figure S25.

2:1 complex. A 0.2 mM solution of $CB[7]_2$•**T-VPI** was prepared by addition of 50 µL of a 2 mM solution of **T-VPI** in D_2O (10^{-7} mol), 50 µL of a 5 mM solution of CB[7] (2.5 10^{-7} mol) in D_2O and 400 µL of D_2O. ^1H NMR of $CB[7]_2$•**T-VPI** (500 MHz, D_2O, 0.2 M, 298 K) δ 9.34 (d, *J* = 6.0 Hz, 2H, ***H2***), 9.06 (br m, 4H, ***H3*** and ***H4***), 8.08 (br s, 2H, ***H5***), 7.89 (br m, ***H8*** and ***H9***), 7.47 (br s, 2H, ***H7***), 7.01 (d, *J* = 8.2 Hz, 2H, ***Hy***), 6.83 (br s, 2H, ***H6***), 6.65 (d, *J* = 8.0 Hz, 2H, ***Hx***), 5.78 (m, 32H, CB[7]), 5.53 (m, 33H, CB[7]), 4.22 (app t, *J* = 15.4 Hz, 36H, CB(7]), 2.22 (acetone, ref), 2.02 (br s, 3H, ***H1***). Figure S28.

NMR of VPI-N-H$^+$ with CB[7]: **VPI-N-H$^+$**. A solution of TFA-D (12.5 mM, 120 µl, 1.5 10^{-6} mol) was added to a 1 mM solution of **VPI-N** in D_2O (500 µL, 0.5 10^{-6} mol), leading to a pale-yellow solution. ^1H NMR of **VPI-N-H$^+$** (300 MHz, D_2O, 1 mM) δ 9.40 (d, *J* = 7.1 Hz, 2H, ***H5***), 9.04 (d, *J* = 6.9 Hz, 2H, ***H2***), 8.69 (d, *J* = 7.0 Hz, 2H, ***H4***), 8.54 (d, *J* = 6.9 Hz, 2H, ***H3***), 8.42 (d, *J* = 8.9 Hz, 2H, ***H7***), 8.30 (s, 2H, ***H8***), 8.14 – 8.05 (m, 4H, overlapped signals of ***H6*** and ***H9***), 7.55 (dd, *J* = 6.5, 3.3 Hz, 2H, ***H10***), 4.48 (s, 3H, ***H1***). Figure S18.

1:1 complex. A 1 mM solution of CB[7]•**VPI-N-H$^+$** was prepared by addition of 125 µL of a 4 mM solution of **VPI-N** (5 10^{-7} mol) in D_2O, 100 µL of a 5 mM solution of CB[7] (5 10^{-7} mol) in D_2O, 120 µL of a 12.5 mM solution of TFA in D_2O (1.5 10^{-6} mol) and 155 µL of D_2O. ^1H NMR of CB[7]•**VPI-N-H$^+$** (400 MHz, D_2O, 1mM) δ 9.15 (d, *J* = 6.1 Hz, 2H, ***H5***), 8.80 (m, 4H, overlapped signals of ***H2*** and ***H3***), 8.65 (m, 4H, overlapped signals of ***H4*** and ***H8***), 8.26 (m, 2H, ***H9*** or ***H10***), 7.70 (m, 4H, overlapped signals of ***H7*** and ***H9*** or ***H10***), 7.32 (d, *J* = 8.2 Hz, 2H, ***H6***), 5.71 (dd, *J* = 23.2, 15.5 Hz, 14H, CB[7]), 5.55 (s, 13H, CB[7]), 4.58 (s, ***H1***), 4.26 (dd, *J* = 15.4, 8.4 Hz, 13H, CB[7]), 2.22 (acetone, ref). Figure S33.

2:1 complex. A 1 mM solution of $CB[7]_2$•**VPI-N-H$^+$** was prepared by addition of 125 µL of a 4 mM solution of **VPI-N** (5 10^{-7} mol) in D_2O, 200 µL of a 5 mM solution of CB[7] (10^{-6} mol) in D_2O, 120 µL of a 12.5 mM solution of TFA in D_2O (1.5 10^{-6} mol) and 60 µL of D_2O. ^1H NMR of $CB[7]_2$•**VPI-N-H$^+$** (300 MHz, D_2O, 1 mM) δ 9.51 (d, *J* = 7.0 Hz, 2H, ***H5***), 9.04 (d, *J* = 6.7 Hz, 2H, ***H2***), 8.89 (d, *J* = 8.8 Hz, 2H, ***H7***), 8.69 (d, *J* = 8.8 Hz, 2H, ***H6***), 7.46 (s, 1H, ***H8***), 7.32 (dd, *J* = 6.4, 3.3 Hz, 2H, ***H9*** or ***H10***), 7.29 – 7.20 (m, 4H, overlapped signals of ***H4*** and ***H9*** or ***H10***), 7.07 (d, *J* = 6.5 Hz, 2H, ***H3***), 5.86 – 5.65 (m, 28H, CB[7]), 5.67 (s, 3H, ***H1***), 5.56 (s, 14H, CB[7]), 5.45 (s, 14H, CB[7]), 4.27 (d, *J* = 15.4 Hz, 14H, CB[7]), 4.17 (dd, *J* = 15.4, 2.3 Hz, 14H, CB[7]), 2.22 (acetone, ref). Figure S36.

NMR of T-VPI-H$^+$ with CB[7]: **T-VPI-H$^+$**. a 5 µL of 3.5% DCl solution (5 10^{-6} mol) were added to a 1 mM solution of **T-VPI** in D_2O (500 µL, 0.5 10^{-6} mol), leading to a colorless solution. ^1H NMR of **T-VPI-H$^+$:** (300 MHz, D_2O) δ 9.44 (d, *J* = 5.3 Hz, 2H, ***H5***), 9.30 (d, *J* = 5.1 Hz, 2H, ***H2***), 8.78 (d, *J* = 5.4 Hz, 2H, ***H4***), 8.72 (d, *J* = 5.5 Hz, 2H, ***H3***), 8.37 (d, *J* = 7.1 Hz, 2H, , ***H7***), 8.13 (d, *J* = 7.3 Hz, 2H, , ***H6***), 7.80 (m, 2H, , ***H8***), 7.66 – 7.54 (m, 4H,overlapped signals of , ***H6*** and ***Hy***), 7.50 (d, *J* = 8.0 Hz, 2H, , ***Hx***), 2.40 (s, 3H, , ***H1***). Figure S19.

1:1 complex. A 1 mM solution of CB[7]•**T-VPI-H$^+$** was prepared by addition of 250 µL of a 2 mM solution of **T-VPI** (5 10^{-7} mol) in D_2O, 100 µL of a 5 mM solution of CB[7] (5 10^{-7} mol) in D_2O, 10 µL of a 200 mM solution of TFA in D_2O (2 10^{-6} mol) and 150 µL of D_2O. The solution was diluted to 0.1 mM. ^1H NMR of CB[7]•**T-VPI**-H$^+$ (500 MHz, D_2O, 0.1 mM) δ 9.33 (br s, ***H2***), 9.08 (br s, ***H5***), 8.89 (br s, ***H4***), 8.81 (br s, ***H3***), 8.02 (br s, ***H8*** or ***H9***), 7.91 (s, ***H7***), 7.74 (br s, ***H8*** or ***H9***), 7.62 (s, ***H6***), 7.46 (br s, ***Hy***), 7.29 (br s, ***Hx***), 5.81 – 5.67 (m, CB[7]), 5.52 (two s, CB[7]), 4.24 (d, *J* = 15.4 Hz, CB[7]), 2.36 (s, ***H1***), 2.22 (acetone, ref). Figure S39.

2:1 complex. A 0.5 mM solution of $CB[7]_2$•**T-VPI-H$^+$** was prepared by addition of 125 µL of a 2 mM solution of **T-VPI** (2.5 10^{-7} mol) in D_2O, 50 µL of a 5 mM solution of CB[7] (2.5 10^{-7} mol) in D_2O, 12 µL of a 100 mM solution of TFA in D_2O (1.2 10^{-6} mol) and 320 µL of D_2O. ^1H NMR of $CB[7]_2$•**T-VPI-H$^+$** (400 MHz, D_2O, 298 K, 0.5 mM) δ 9.27 (d, *J* = 7.0 Hz, 2H, ***H2***), 9.12 (d, *J* = 6.7 Hz, 2H, ***H4***), 9.00 (d, *J* = 6.9 Hz, 2H, ***H3***), 8.86 (d, *J* = 7.0 Hz, 2H, ***H5***), 8.12 (dd, *J* = 6.2, 3.2 Hz, 2H, ***H8*** or ***H9***), 7.80 (dd, *J* = 6.2, 3.2 Hz, 2H, ***H8*** or ***H9***), 7.49 (d, *J* = 8.6 Hz, 2H, ***H7***), 7.23 (d, *J* = 8.6 Hz, 2H, ***H6***), 6.97 (d, *J* = 8.4 Hz, 2H, ***Hy***), 6.65 (d, *J* = 8.3 Hz, 2H, ***Hx***), 5.75 (app ddd, *J* = 22.9, 15.4, 9.8 Hz, 28H, CB[7]), 5.65 – 5.43 (m, 28H, 28H, CB[7]), 4.24 (app ddd, *J* = 23.7, 15.5, 10.6 Hz, 28H, 28H, CB[7]), 2.22 (acetone, ref), 2.02 (br s, 3H, ***H1***). Figure S44.

NMR of VPI-N•Ag$^+$ with CB[7]: 1:1:1 complex. a 0.5 mM solution of CB[7]•**VPI-N**•Ag^+ was prepared by addition of 62.5 µL of a 4 mM solution of **VPI-N** (2.5 10^{-7} mol) in D_2O, 50 µL of a 5 mM solution of CB[7] (2.5 10^{-7} mol) in D_2O, 25 µL of a 200 mM solution of $AgNO_3$ in D_2O (5 10^{-6} mol) and 380 µL of D_2O. ^1H NMR of CB[7]•**VPI-N**•Ag^+ (500 MHz, D_2O, 300 K, 0.5 mM) δ 9.15 (d, *J* = 6.0 Hz, 2H, ***H2***), 8.76 (m, overlapped signals of ***H4*** and ***H3***), 8.51 (br s, 2H, ***H8***), 8.45 (d, *J* = 5.8 Hz, 2H, ***H5***), 8.21 (br s, 2H, ***H9*** or ***H10***), 7.86 (d, *J* = 8.0 Hz, 2H, ***H7***), 7.60 (br s, 2H, ***H9*** or ***H10***), 7.06 (d, *J* = 7.7 Hz, 2H, ***H6***), 5.69 (app dd, *J* = 23.2, 15.5 Hz, 14H, CB[7]), 5.53 (s, 14H, CB[7]), 4.58 (br s, ***H1***), 4.24 (d, *J* = 15.5 Hz, 14H, CB[7]), 2.22 (acetone, ref). Figure S48.

2:1:1 complex. To a 0.5 mM solution of $CB[7]_2$•**VPI-N**•Ag^+ (2.5 10^{-7} mol) was added 50 µL of a 5mM solution of CB[7] in D_2O (2.5 10^{-7} mol). ^1H NMR of $CB[7]_2$•**VPI-N**•Ag^+ (500 MHz, D_2O, 300 K, 0.5 mM) δ 9.49 (d, *J* = 5.9 Hz, 2H, ***H5***), 9.03 (d, *J* = 5.5 Hz, 2H, ***H2***), 8.98 (d, *J* = 8.3 Hz, 2H, ***H7***), 8.57 (d, *J* = 8.1 Hz, 2H, ***H6***), 7.50 (d, *J* = 7.6 Hz, 1H, ***H9***), 7.42 – 7.29 (m, 3H, overlapped signals of ***H8***, ***H10*** and ***H9'***), 7.24 (two d, *J* = 16.8, 6.8 Hz, 3H, overlapped signals of ***H10'*** and ***H4***), 7.16 (br s, 1H, ***H8'***), 7.07 (d, *J* = 5.5 Hz, 2H, ***H3***), 5.73 (dd, *J* = 15.0, 7.2 Hz, 28H, CB[7]), 5.56 (s, 14H, CB[7]), 5.46 (s, 14H, CB[7]), 4.27 (d, *J* = 15.6 Hz, 14H, CB[7]), 4.17 (dd, *J* = 15.4, 3.0 Hz, 14H, CB[7]), 2.22 (acetone, ref). Figure S52.

NMR of T-VPI•Ag$^+$ with CB[7]: 1:1:1 complex. A 0.5 mM solution of CB[7]•**T-VPI**•Ag^+ was prepared by addition of 125 µL of a 2 mM solution of **T-VPI** (2.5 10^{-7} mol) in D_2O, 50 µL of a 5 mM solution of CB[7] (2.5 10^{-7} mol) in D_2O, 25 µL of a 200 mM solution of $AgNO_3$ in D_2O (5 10^{-6} mol) and 300 µL of D_2O. ^1H NMR of CB[7]•**T-VPI**•Ag^+ (500 MHz, D_2O, 298 K, 0.5 mM) δ 9.35 (br s, 2H, ***H2***), 8.89 (br s, 4H, overlapped signals of ***H3*** and ***H4***), 8.80 (br s, 2H, ***H5***), 7.95 (br m, 4H, overlapped signals of ***H7*** and ***H8*** or ***H9***), 7.52 (br s, 6H, overlapped signals of ***H6***, ***Hy*** and ***H8*** or ***H9***), 7.34 (br s, 2H, ***Hx***), 5.71 (d, *J* = 15.2 Hz, 14H, CB[7]), 5.52 (br s, 14H, CB[7]), 4.23 (d, *J* = 15.4 Hz, 14H, CB[7]), 2.37 (br s, 3H, ***H1***), 2.22 (acetone, ref). Figure S54.

2:1:1 complex. A 0.5 mM solution of $CB[7]_2$•**T-VPI**•Ag^+ was prepared by addition of 125 µL of a 2 mM solution of **T-VPI** (2.5 10^{-7} mol) in D_2O, 100 µL of a 5 mM solution of CB[7] (5 10^{-7} mol) in D_2O, 25 µL of a 200 mM solution of $AgNO_3$ in D_2O (5 10^{-6} mol) and 250 µL of D_2O. ^1H NMR (500 MHz, D_2O) δ 9.28 (d, *J* = 6.8 Hz, 2H, ***H2***), 9.07 (d, *J* = 6.9 Hz, 2H, ***H4***), 9.04 (d, *J* = 6.7 Hz, 2H, ***H3***), 8.49 (d, *J* = 6.9 Hz, 2H, ***H5***), 7.99 (app t, *J* = 8.1 Hz, 2H, ***H8-H8'*** or ***H9-H9'***), 7.70 (d, *J* = 8.5 Hz, 7H, ***H7***), 7.61 – 7.52 (m, 2H, ***H8-H8'*** or ***H9-H9'***), 6.98 (two d, *J* = 8.5 Hz, 4H, overlapped signals of ***H6*** and ***Hy***), 6.65 (d, *J* = 8.1 Hz, 2H, ***Hx***), 5.83 (d, *J* = 15.5 Hz, 7H, CB[7]), 5.79 (d, *J* = 15.5 Hz, 7H, CB[7]), 5.72 (d, *J* = 15.5 Hz, 14H, CB[7]), 5.52 (d, *J* = 17.8 Hz, 28H, CB[7]), 4.29 – 4.16 (m, 28H, CB[7]), 2.22 (acetone, ref), 2.01 (br s, 3H, ***H1***). Figure S57.

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