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Anion-templated synthesis of a switchable fluorescent [2]catenane with sulfate sensing capability†

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Anion templation strategies have facilitated the synthesis of various catenane and rotaxane hosts capable of strong and selective binding of anions in competitive solvents. However, this approach has primarily relied on positively charged precursors, limiting the structural diversity and the range of potential applications of the anion-templated mechanically interlocked molecules. Here we demonstrate the synthesis of a rare electroneutral [2]catenane using a powerful, doubly charged sulfate template and a complementary diamidocarbazole-based hydrogen bonding precursor. Owing to the unique three-dimensional hydrogen bonding cavity and the embedded carbazole fluorophores, the resulting catenane receptor functions as a sensitive fluorescent turn-ON sensor for the highly hydrophilic sulfate, even in the presence of a large excess of water. Importantly, the [2]catenane exhibits enhanced binding affinity and selectivity for sulfate over its parent macrocycle and other acyclic diamidocarbazole-based receptors. We demonstrate also, for the first time, that the co-conformation of the catenane may be controlled by reversible acid/base induced protonation and deprotonation of the anionic template, SO_4^{2-} . This approach pioneers a new strategy to induce molecular motion of interlocked components using switchable anionic templates.

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Introduction

Mechanically interlocked molecules (MIMs), such as catenanes, rotaxanes, or knots, have emerged as captivating hosts for molecular recognition,^{1–5} with ever-expanding applications in sensing,^{6–12} catalysis,^{9,13–18} and the construction of molecular machines.^{18–28} They are particularly appealing for their three-dimensional cavities, which can serve as distinctive, solvent-shielded binding sites,^{29,30} and for the unique dynamic properties of their interlocked components (*e.g.* shuttling or pirouetting motions). Anion templation strategies,^{31,32} pioneered by Beer and co-workers, have facilitated the synthesis of numerous hydrogen bonding^{33–37} (HB) and halogen bonding^{38–42} (XB) interlocked hosts capable of strong and selective binding of anions in competitive solvents.^{43–46} Importantly, such receptors often outperform their non-interlocked analogues in terms of both affinity and selectivity, positioning them as highly promising candidates for molecular sensing applications.^{47–51}

To date, however, anion templation strategies have predominantly relied on positively charged precursors,

leveraging their capacity for effective preorganization through strong interactions with anionic templates. This significantly limits both the structural diversity and potential applications of anion templated MIMs. For example, anion transport across lipid bilayers requires highly lipophilic, and hence typically uncharged, hosts,^{52,53} and therefore only very recently has the first [2]catenane anion transporter been described.⁵⁴ Moreover, the ever-present counterions compete for binding sites of charged receptors, diminishing their selectivity. Thus, further development and future applications of the anion templation methodology depend on the availability of charge neutral and synthetically versatile building blocks, capable of forming orthogonal complexes with strongly interacting anionic templates.

Tetrahedral sulfate with its dinegative charge, strong HB-accepting capabilities, and presumed orthogonal coordination preference, is a particularly attractive anionic template (Fig. 1).^{55–57} However, there are only two examples of its use in the synthesis of MIMs thus far, both described by Beer and co-workers. The first is a doubly charged [2]catenane, which binds sulfate strongly but less selectively than its acyclic precursor, presumably due to its higher charge.⁵⁸ The other is an electro-neutral triply interlocked capsule, which sequesters sulfate during synthesis, but cannot release it, even in the presence of water.⁵⁹ In both cases the interlocked cavity proved to be

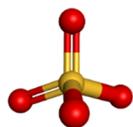
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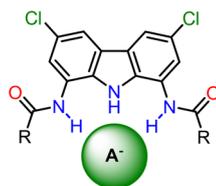


Sulfate Template



- ✓ Strong HB Acceptor
- ✓ Dinegative Charge
- ✓ Orthogonal Coordination Preference

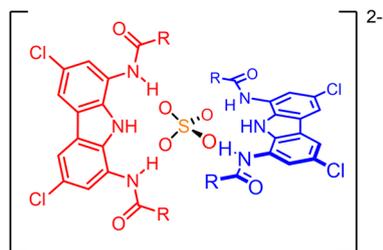
Diamidocarbazole Building Block



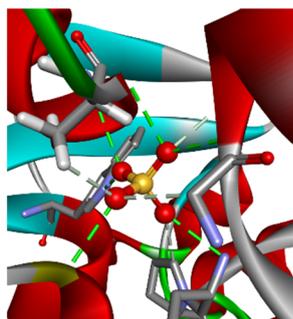
- ✓ Strong HB Donor
- ✓ No Permanent Charge
- ✓ Fluorescent
- ✓ Forms Orthogonal Complexes with Sulfate

Inspiration:

Orthogonal Complexes with Sulfate



Sulfate Binding Protein



Design:

Sulfate Binding Electroneutral Fluorescent MIM

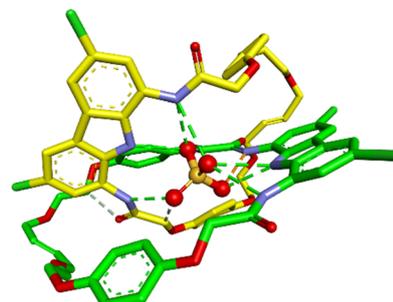


Fig. 1 Sulfate template and high affinity diamidocarbazole building blocks enable the synthesis of the electroneutral fluorescent [2]catenane, capable of binding sulfate inside an interlocked hydrogen bonding cavity reminiscent of the binding site of Sulfate Binding Protein (SBP).

a promising binding site for sulfate, yet with limited functionality and no sensing capabilities.

In a previous report, we showed that simple and readily available diamidocarbazoles exhibit exceptionally high affinity for sulfate.⁶⁰ Despite the lack of permanent charge and the presence of only three hydrogen bond donors, they form robust 2 : 1 (ligand : anion) complexes with this anion, even in highly competitive DMSO–water mixtures (Fig. 1). Most importantly, these complexes adopt orthogonal geometry, which is a prerequisite for the successful assembly of interlocked structures. Together with the outstanding fluorescent sensing properties of diamidocarbazoles, this encouraged us to strategically use these building blocks in the anion templated synthesis of electroneutral fluorescent MIMs capable of binding and sensing sulfate.

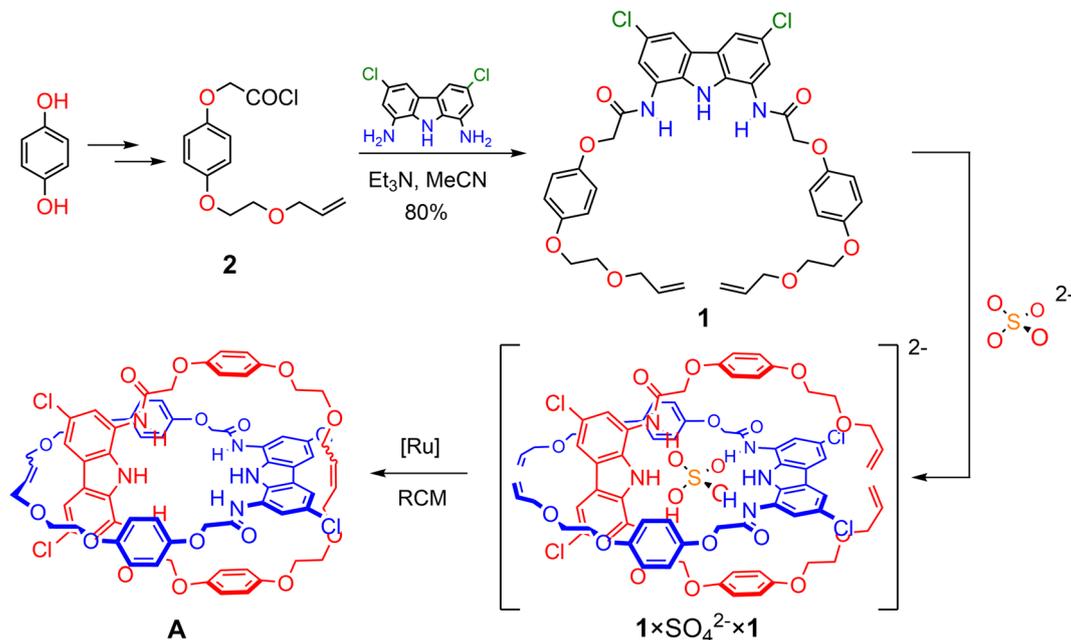
Sulfate recognition in aqueous media draws significant scientific and technological interest due to its important role in biology, medicine, and industry.^{61–83} Sulfate is the fourth most abundant anion in human plasma (serum levels 0.3–0.5 mM) and has important functions in human physiology.^{84,85} For example, it is used in the sulfonation of various endo- and exogenous compounds, which, *inter alia*, regulate functionality of secreted proteins, hormones, and signaling molecules. Sulfonation is involved in the control of various biological processes, including blood clotting and the formation of connective tissues.^{86–88} Unusually low levels of sulfate have been found in the plasma of patients with rheumatoid arthritis and irritable bowel disease.^{89–92} In industry, sulfate is known, for example, to interfere with the remediation of nuclear waste.⁶² Hence, sulfate-selective sensors could be used to elucidate the

diverse roles of sulfate in living organisms, allow improved diagnostics, and facilitate monitoring of environmental samples.^{93–108}

Despite the growing understanding of the biological, environmental, and technological importance of sulfate, examples of its fluorescent sensing in aqueous media are very limited.^{99–107} In fact, sulfate remains one of the most challenging anionic targets for molecular recognition in water, primarily due to its extremely high hydration energy ($\Delta G = -1090 \text{ kJ mol}^{-1}$). Therefore, inspired by the sulfate binding protein, which strongly and selectively binds sulfate in water by encapsulating the anion using a network of hydrogen bonds,^{109–111} we envisaged that strategic incorporation of strong hydrogen bond donors inside a three-dimensional cavity of an electroneutral fluorescent catenane could mimic these characteristics and allow strong binding and selective sensing of sulfate in aqueous media (Fig. 1).

Herein we show that by using powerful, doubly charged anionic template, SO_4^{2-} , and complementary HB precursors, it is possible to synthesize charge-neutral [2]catenane **A** (Scheme 1), which is a rare example of an electroneutral MIM achieved by the anion templation strategy.^{38,54,59,112,113} Importantly, catenane **A** binds sulfate more strongly and more selectively than its parent macrocycle, its acyclic precursor, and a model receptor containing two diamidocarbazole units linked covalently. Catenane **A** serves as a sensitive turn-ON fluorescent sensor for the extremely hydrophilic sulfate, even in highly competitive aqueous-organic medium. Additionally, we demonstrate, for the first time, that the relative disposition of the macrocyclic components of the catenane (its co-conformation) can be





Scheme 1 Synthesis of [2]catenane A via RCM of the orthogonal assembly of $1 \times \text{SO}_4^{2-} \times 1$.

controlled by reversible acid/base induced protonation–deprotonation of SO_4^{2-} . This approach pioneers a new strategy to induce reversible molecular motions using switchable anionic templates.

Results and discussion

Synthesis of the [2]catenane precursor and its sulfate-directed self-assembly

Based on our previous studies,⁶⁰ we envisaged that sulfate directed orthogonal assembly of two appropriately designed diamidocarbazole ligands **1** would enable the synthesis of homo[2]catenane **A** in a one-pot, double ring-closing metathesis (RCM) reaction (Scheme 1). Alkene-terminated precursor **1** was obtained in a convergent synthesis, ending with the reaction of acyl chloride **2** with 1,8-diamino-3,6-dichlorocarbazole (Scheme 1). The alkene-terminated arm **2** was synthesized in 4 steps from 1,4-hydroquinone (Scheme S1†), and 1,8-diamino-3,6-dichlorocarbazole was obtained in 3 steps from carbazole, according to the previously reported procedure.^{114,115}

The sulfate-directed self-assembly of precursor **1** was investigated by ¹H NMR titrations. The addition of tetrabutylammonium (TBA) sulfate to the solution of **1** in DMSO-*d*₆ induced large downfield perturbations of the carbazole (α , $\Delta\delta \approx 2.5$ ppm) and amide (β , $\Delta\delta \approx 2.0$ ppm) NH proton signals, indicating that the anion forms strong hydrogen bonds with these donors (Fig. 2). Interestingly, the signals of the hydroquinone protons B and C shifted significantly upfield (B, $\Delta\delta \approx 0.2$ ppm; C, $\Delta\delta \approx 0.4$ ppm) until 0.5 equivalent of TBA₂SO₄ was added, whereas further addition of sulfate shifted them in the opposite direction. At around 1 : 1 ligand-to-sulfate ratio these signals returned almost to their starting positions and at a host : guest ratio > 1.2 no

further changes could be observed (Fig. 2c). A similar perturbation pattern was observed for signals D–F, corresponding to the –OCH₂– protons, but the magnitude of these changes decreased with increasing distance from the hydroquinone ring. This characteristic behavior has been previously observed during titrations of simple diamidocarbazoles with TBA sulfate, and attributed to the consecutive formation of 2 : 1 and 1 : 1 ligand : anion complexes.⁶⁰ In the orthogonal 2 : 1 complex, protons positioned close to the carbazole core of the other ligand experience substantial shielding. However, as the concentration of sulfate increases, the orthogonal 2 : 1 complex gives way to the 1 : 1 complex, in which these close contacts are very unlikely.

The lack of strong auxiliary interactions (e.g. π – π stacking of electron-deficient and electron-rich aromatic rings) suggests that the orthogonal assembly depends mostly on the strong hydrogen bonding between the two ligands and the tetrahedral sulfate. Indeed, despite the highly competitive solvent – DMSO-*d*₆, the association constant of electrically neutral receptor **1** with SO_4^{2-} turned out to be too high to be reliably measured by ¹H NMR titrations ($K_{1:1} > 10^5 \text{ M}^{-1}$, $K_{2:1} > 10^3 \text{ M}^{-1}$). As DMSO is not compatible with RCM, the analogous binding studies were also conducted in CDCl₃. The results confirmed that strong orthogonal assembly also takes place in this medium (Fig. S42 and S43;† $K_{1:1} > 10^5 \text{ M}^{-1}$, $K_{2:1} > 10^3 \text{ M}^{-1}$), boding well for the sulfate-templated synthesis of [2]catenane *via* ring-closing metathesis.

Sulfate-templated synthesis of [2]catenane A

Catenane precursor **1** is poorly soluble in typical solvents used for RCM, such as DCM or toluene. However, it dissolves well in the presence of 0.5 equivalents of TBA sulfate, demonstrating again the strong affinity of diamidocarbazoles to the anionic template. 2nd Generation Grubbs catalyst, which is commonly used in the anion templated synthesis of interlocked structures, turned out to



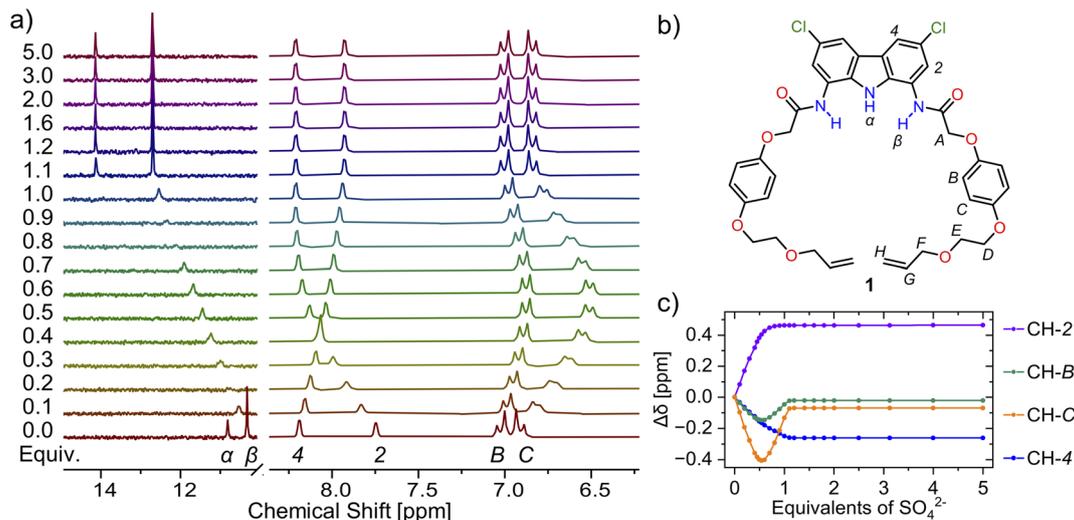


Fig. 2 (a) Truncated ^1H NMR spectra of precursor **1** (b) with increasing amount of TBA_2SO_4 in $\text{DMSO}-d_6 + 0.5\% \text{H}_2\text{O}$. (c) Corresponding binding isotherms. The solid lines are visual aid only.

be inactive in the RCM of $1 \times \text{SO}_4^{2-} \times 1$. We suspect that the high basicity of sulfate in anhydrous and aprotic medium is responsible for the rapid deactivation of the catalyst. Screening of other commercially available catalysts showed that only the relatively fast-initiating Hoveyda-type complexes, particularly nitro-Grela (SIPr), allowed for the complete conversion of the substrate. To further limit the exposure of the catalyst to sulfate, the catalyst was slowly added to the reaction mixture using a syringe pump. Also, tetrafluoro-1,4-benzoquinone was added to the reaction mixture to prevent isomerization of the terminal allyloxy groups to unreactive vinyl ethers. The latter unwanted reaction often accompanies olefin metathesis catalysed by Ru complexes.¹¹² Ultimately, two major products were isolated and separated from SO_4^{2-} template by flash chromatography: [2]catenane **A** and macrocycle **3** (Fig. 3).

The product ratio was found to strongly depend on substrate concentration. While the high concentration of both the precursor and the template favors the formation of the orthogonal 2:1 complex, it also lowers the overall macrocyclization yield due to the formation of insoluble oligomers.

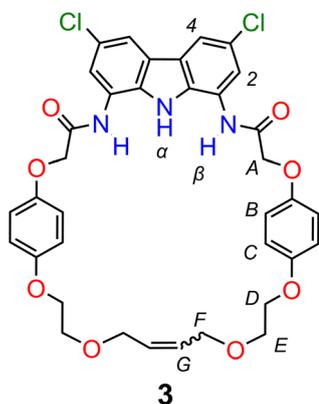


Fig. 3 Macrocycle **3** – a side product of the synthesis of [2]catenane **A**.

On the other hand, very high dilution also decreased the yield of catenane **A**, likely due to the dissociation of the 2:1 complex. Under optimized conditions (1 mM of **1** in DCM, RT), catenane **A** was synthesized in 26% yield and macrocycle **3** in 14% yield.

Sulfate templation was found to be indispensable for obtaining catenane **A** in considerable yield (Table 1). Attempts to synthesize **A** in the absence of any anionic template failed due to the poor solubility of precursor **1** in DCM. However, the solubility of **1** in chloroform was sufficient to carry out RCM without the addition of any anion. The main product under these conditions was macrocycle **3** (40%) with only trace amounts of catenane **A** (2%). Chloride, which has been the most popular anionic template for the synthesis of catenanes and rotaxanes,⁴³ did not significantly improve the solubility of **1** in DCM. In chloroform, however, RCM in the presence of 0.5

Table 1 Effects of concentration and template on the synthesis of catenane **A** from precursor **1**^a

Entry	Concentration of 1	Template	Solvent	Yield of A
1	1 mM	0.5 equiv. SO_4^{2-}	DCM	26%
2	0.5 mM	0.5 equiv. SO_4^{2-}	DCM	23%
3	20 mM	0.5 equiv. SO_4^{2-}	DCM	0%
4	1 mM	1.0 equiv. SO_4^{2-}	DCM	4%
5	1 mM	—	CHCl_3	2%
6	1 mM	0.5 equiv. Cl^-	CHCl_3	5%

^a Isolated yields of a single run.



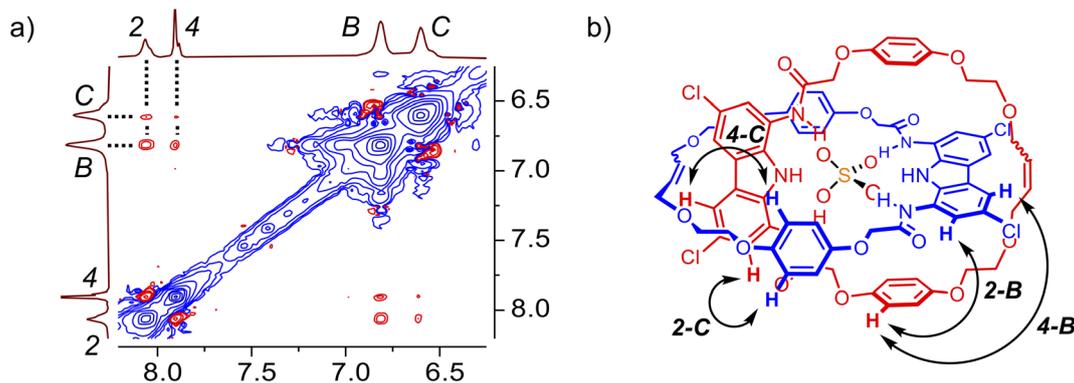


Fig. 4 (a) Fragment of the 2D ROESY spectra of $A \times SO_4^{2-}$. (b) Contacts between protons of the carbazole ring and the hydroquinone ring, which are possible due to the interlocked nature of the receptor.

equivalent of chloride afforded similar product distribution as in the absence of any anion: 43% of macrocycle **3** and 5% of catenane **A**.

Catenane **A** and macrocycle **3** were characterized by high-resolution mass spectrometry (HR-MS) and a series of NMR experiments. The HR-MS of **A** revealed an intense signal at $m/z = 1435.54$, corresponding to a single-charged ion with an isotopic profile matching $[A \times Na]^+$. The 1H NMR spectra of catenane **A** and macrocycle **3** in DMSO were very similar (Fig. S11†). The only differences were minor upfield perturbations ($\Delta\delta \approx 0.05$ ppm) of aromatic CH signals of carbazole and hydroquinone moieties in catenane **A**, when compared to its parent macrocycle **3**. This small shift may be attributed to the proximity of aromatic rings in some conformations of the mechanically interlocked structure. Nevertheless, the interlocked nature of **A** was further confirmed by 2D 1H - 1H ROESY NMR and titration experiments, as described below.

The ROESY NMR spectra of **A** and **3** in DMSO- d_6 did not reveal any specific through-space contacts, which would be unequivocally attributed to the interlocked structure (Fig. S10†). This is likely due to rapid (co-)conformational changes of **A** on the NMR time scale. However, in the presence of 1 equivalent of TBA_2SO_4 , distinct cross-peaks appeared in the ROESY spectrum of **A**, due to close contacts between hydroquinone protons B and C, as well as ether protons D-F, with carbazole protons CH-2 and CH-4 (Fig. 4 and S12†). The interlocked nature of **A** allows the hydroquinone rings of one macrocycle to be positioned above/below the carbazole ring of the other, what becomes clearly visible when sulfate binding rigidifies the whole structure.

Anion binding studies of [2]catenane **A** and macrocycle **3**

1H NMR titrations. The anion binding properties of [2]catenane **A** and macrocycle **3** were studied by 1H NMR titrations with TBA salts. Addition of 1 equivalent of TBA_2SO_4 to the solution of **A** in DMSO- d_6 + 0.5% H_2O caused a significant downfield shift and broadening of the carbazole (α , $\Delta\delta \approx 2$ ppm) and amide (β , $\Delta\delta \approx 1$ ppm) NH signals due to the formation of strong hydrogen bonds with the anion (Fig. 5). Hydroquinone proton signals B and C moved steadily upfield as

a result of shielding caused by the proximity of the carbazole moiety of the second macrocyclic component (Fig. 5b). Addition of the second equivalent of sulfate led to the formation of a new set of signals (slow exchange on the NMR timescale). NH signals were further shifted downfield by *ca.* 1 ppm, while hydroquinone signals B and C returned to the vicinity of their starting positions. No further changes were observed after addition of more than 2.5 equivalents of SO_4^{2-} . These results showed that the catenane forms a robust 1 : 1 complex with sulfate, in which the anion is bound by two carbazole moieties. Slow exchange on the NMR timescale suggests that sulfate is particularly tightly bound in the interlocked cavity, in contrast to the complexes formed by macrocycle **3** (*vide infra*). The 1 : 2 (receptor : anion) complex, which dominates in the presence of excess of sulfate, appears to have a distinct co-conformation, in which the two carbazole subunits individually bind different sulfate ions and presumably move away from each other due to electrostatic repulsion between the anions (Fig. 5a). Another proof of this notable co-conformational change was provided by the ROESY spectrum of catenane **A** in the presence of 2.5 equivalents of sulfate (Fig. S13†). Specifically, no cross-peaks were found between hydroquinone and carbazole CH signals, in stark contrast to the results obtained in the presence of 1 equivalent of sulfate (*vide supra*).

The co-conformational change can also be tracked by diffusion-ordered NMR spectroscopy (DOSY). The addition of 1 equiv. of TBA_2SO_4 to the solution of free catenane **A** in DMSO- d_6 results in an increase of the diffusion coefficient by 8%, from $1.03 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$ to $1.11 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$. This increase in mobility despite increasing mass may be attributed to locking of the flexible free catenane in a more compact, contracted co-conformation, characteristic of the 1 : 1 complex with sulfate. Further addition of sulfate (to 2.5 equiv.) leads to a significant decrease of the diffusion coefficient by 20% (to $0.89 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$), much higher than might be expected based on the increasing molecular mass. This is in line, however, with the adoption of the expanded 1 : 2 complex with a much larger hydrodynamic radius (Fig. S14–S17†).¹¹⁶

Increase of water content significantly simplifies the sulfate binding equilibria of **A**. In DMSO- d_6 /H₂O 9 : 1, only the 1 : 1



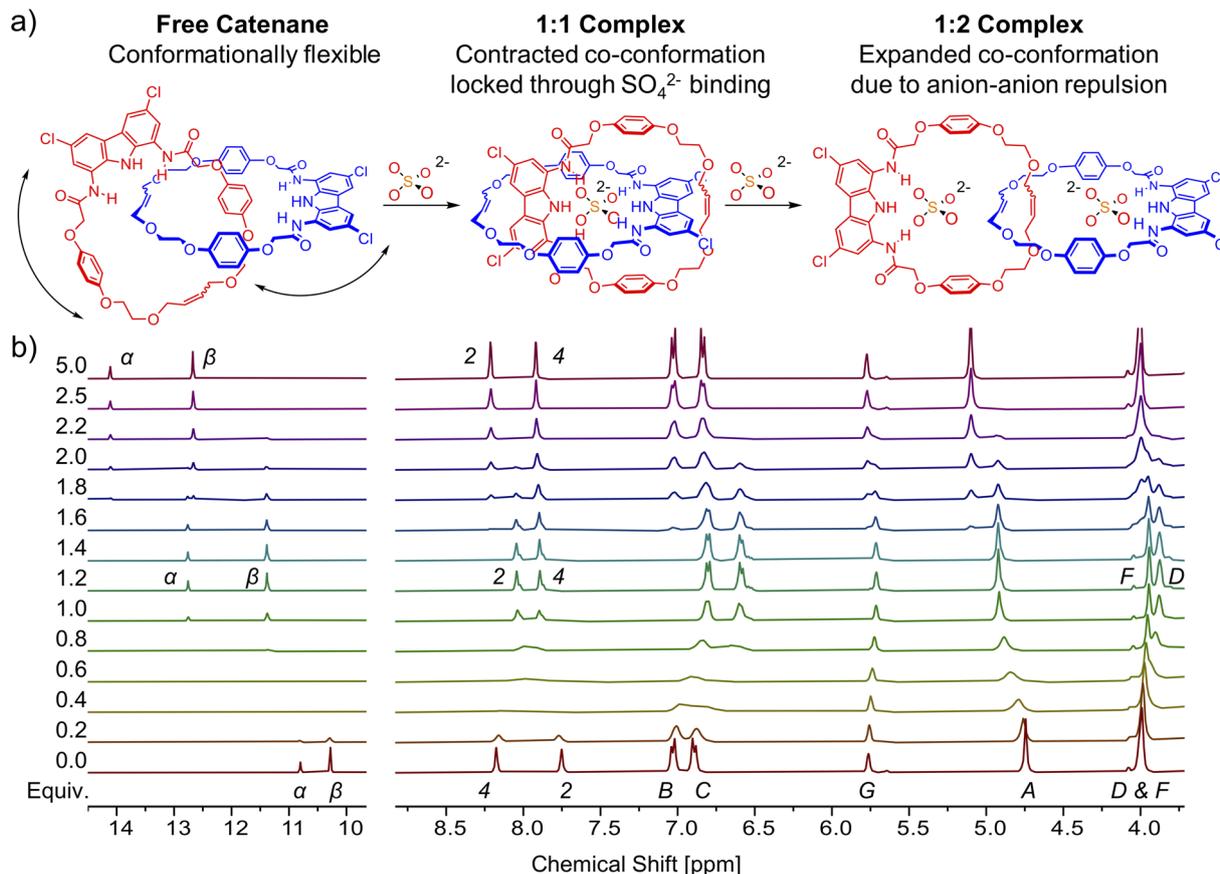


Fig. 5 (a) Graphical representation of the complexes formed during the titration of [2]catenane A with TBA₂SO₄. (b) Truncated ¹H NMR spectra of [2]catenane A with increasing amount of TBA₂SO₄ in DMSO-*d*₆ + 0.5% H₂O.

complex is formed (Section S5.1.2[†]), even in the presence of excess of sulfate. This is presumably due to stronger stacking of the catenane's aromatic surfaces in the 1 : 1 A × SO₄²⁻ complex as a consequence of the hydrophobic effect.^{117,118} Moreover, the interlocked cavity efficiently shields the bound guest from the surrounding solvent molecules, further increasing the stability of the complex.¹¹⁹ This result clearly illustrates the potential of interlocked microenvironments in the design of molecular receptors aimed at binding in aqueous media.

Similar to catenane A, titration of TBA₂SO₄ to the solution of macrocycle 3 in DMSO-*d*₆ + 0.5% H₂O broadens both the NH signals and shifts them significantly downfield (α, Δδ ≈ 3 ppm; β, Δδ ≈ 2.5 ppm). In case of 3, however, no further changes were observed after the addition of 1.2 equivalents of sulfate (Fig. 6). Hydroquinone signals B and C shifted only marginally during the course of the titration, but close inspection of the binding isotherm for signal B revealed a shallow minimum at around 0.5 equivalents of sulfate (Fig. 6c), likely due to its weak shielding by an aromatic ring. Apparently, alongside the 1 : 1 complex, the macrocycle 3 forms also a 2 : 1 (receptor : anion) complex. Molecular modeling of the 2 : 1 complexes using Conformer-Rotamer Ensemble Sampling Tool with the GFN2-xTB approach revealed multiple low energy arrangements,^{120,121} including orthogonal and sandwich type structures, suggesting

that several binding modes could contribute to the overall association (Section S6.3[†]).

UV-Vis titrations. While NMR titrations provided invaluable structural insights into the binding properties of receptors A and 3, the quantitative analysis was primarily performed using UV-Vis spectroscopy, which allowed determination of higher binding constants. UV-Vis titrations were conducted in a highly competitive aqueous-organic solvent medium: DMSO/H₂O 9 : 1. The addition of oxoanions such as sulfate (SO₄²⁻), dihydrogen phosphate (H₂PO₄⁻) and benzoate (PhCOO⁻), as TBA salts, elicited a sharp increase of absorbance in the 340–390 nm region of the UV-Vis spectra of catenane A (Fig. 7, S50–S53[†]), macrocycle 3 (Fig. S56–S61[†]) and acyclic precursor 1 (Fig. S62–S65[†]). In contrast, no significant changes were observed upon the addition of chloride (Cl⁻), indicating that it is not appreciably bound by these receptors in this competitive medium.

Numerical analysis performed using the HypSpec software suite revealed that catenane A binds sulfate stronger than other oxoanions with a high binding constant of $K = 776\,000\text{ M}^{-1}$ for the 1 : 1 complex. This value is almost 10 times higher than that of acyclic precursor 1 and more than 20 times higher than that of macrocycle 3 (Table 2). Surprisingly, macrocycle 3 binds sulfate more weakly than its acyclic precursor 1, presumably due to its increased rigidity and energetic cost associated with



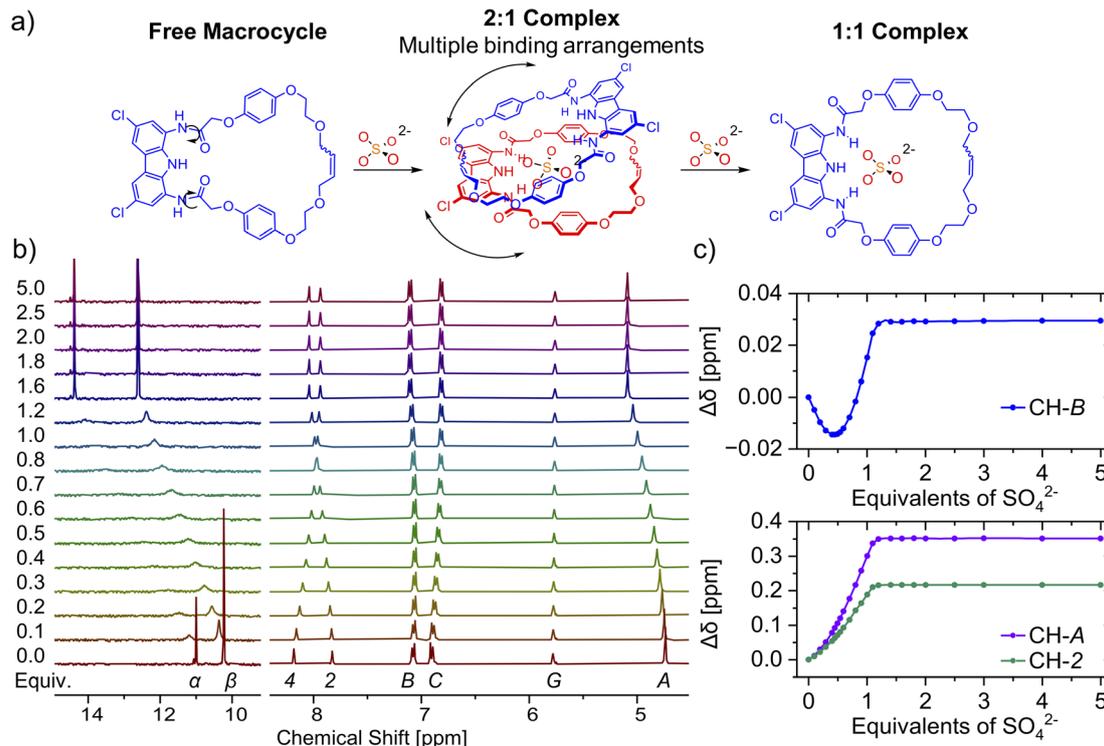


Fig. 6 (a) Graphical representation of the complexes formed during the titration of macrocycle 3 with TBA₂SO₄. (b) Truncated ¹H NMR spectra of macrocycle 3 with increasing amount of TBA₂SO₄ in DMSO-*d*₆ + 0.5% H₂O. (c) Corresponding binding isotherms. The solid lines are visual aid only.

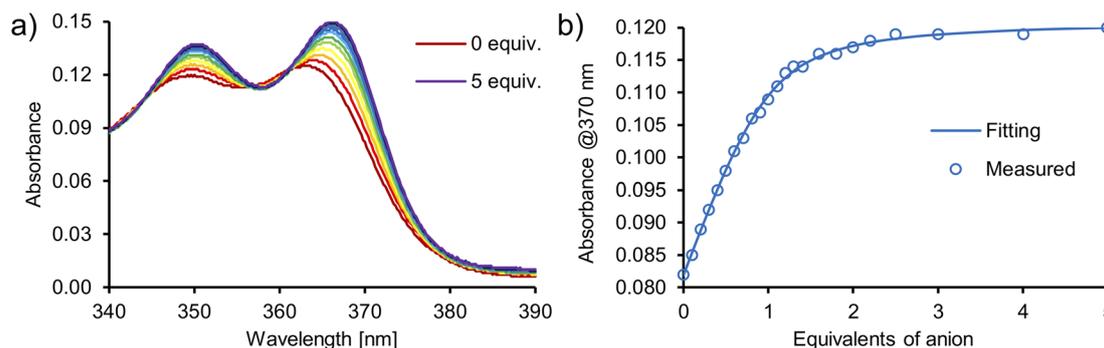


Fig. 7 (a) UV-Vis spectra recorded during the titration of [2]catenane A with TBA₂SO₄ in DMSO/H₂O 9 : 1. (b) 1 : 1 host-guest binding curve fitted to the results obtained during the titration of [2]catenane A with TBA₂SO₄ in DMSO/H₂O 9 : 1.

conformational changes and solvent rearrangement upon binding. This supposition was supported by the results of DFT calculations (Section S6.4†). Notably, catenane A binds sulfate 2.5 times stronger than the previously reported receptor 4 (Fig. 8), which binds sulfate with two diamidocarbazole moieties connected covalently, rather than by a mechanical bond.¹⁰³ The observed selectivity trend: SO₄²⁻ > H₂PO₄⁻ > PhCOO⁻ > Cl⁻ follows the series of HB acceptor parameters for anions.¹²² However, the 1 : 1 association constant of catenane A with sulfate is almost 100 times higher than that of dihydrogen phosphate and over 3000 times higher than that of benzoate, revealing excellent selectivity, which is also significantly enhanced in comparison to acyclic precursor 1, macrocycle 3, and bis(diamidocarbazole)

receptor 4. The high affinity and selectivity of catenane A for sulfate could be attributed to exceptionally good size and shape complementarity between the interlocked binding cavity of A and tetrahedral SO₄²⁻.

Fluorescent sensing of anions by [2]catenane A

The carbazole moiety directly incorporated into the interlocked cavity serves as a convenient fluorescent binding probe. Addition of TBA₂SO₄ to a 10 μM solution of catenane A in DMSO/H₂O 9 : 1 elicited a vivid increase of fluorescence emission intensity in the 360–440 nm range (up to 60%, Fig. 9). Strong sulfate binding resulted in saturation of the response after only 1.5 equivalents of SO₄²⁻ (10 μM) were added. In comparison,



Table 2 Association constants K_a [M^{-1}] of [2]catenane **A**, macrocycle **3**, acyclic precursor **1** and bis(diamidocarbazole) receptor **4** with model anions

Receptor	SO_4^{2-} ^a	$H_2PO_4^-$ ^a	$PhCOO^-$ ^a	Selectivity for SO_4^{2-} vs. $H_2PO_4^-$ ^b	Selectivity for SO_4^{2-} vs. $PhCOO^-$ ^b
Catenane A	$K_{1:1} = 776\,000$	$K_{1:1} = 8130$ $K_{1:2} = 3240$	$K_{1:1} = 230$	95	3400
Macrocycle 3	$K_{1:1} = 32\,400$ $K_{2:1} = 2290$	$K_{1:1} = 760$	$K_{1:1} = 300$ $K_{2:1} = 4900$	43	110
Acyclic precursor 1	$K_{1:1} = 83\,200$ $K_{2:1} = 1170$	$K_{1:1} = 2750$	$K_{1:1} = 250$	30	330
Bis(diamidocarbazole) receptor 4	$K_{1:1} = 295\,000$	$K_{1:1} = 10\,700$ $K_{1:2} = 1230$	$K_{1:1} = 810$	28	360

^a Measured by UV-Vis titrations in DMSO/ H_2O 9 : 1 at 298 K. Error < 15%. ^b Calculated as a ratio of $K_{1:1}(SO_4^{2-})$ and $K_{1:1}(H_2PO_4^-)$ or $K_{1:1}(PhCOO^-)$ for a given receptor. Error < 20%.

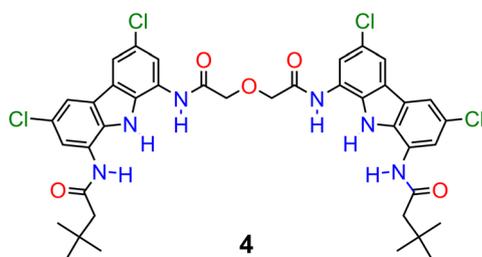


Fig. 8 Bis(diamidocarbazole) receptor **4**.

benzoate did not noticeably affect the fluorescence emission of the catenane in a similar concentration range, and led to only a 15% increase of emission intensity after the addition of 1000 equivalents (Fig. S68[†]). The turn-ON sensing response is likely due to the restriction of vibrational and rotational modes of the receptor upon anion binding. Such rigidification limits non-radiative internal conversion, which competes with the fluorescence emission. Interestingly, sensing selectivity for sulfate vs. dihydrogen phosphate is even more pronounced, owing to a distinct optical response elicited by phosphate binding. More specifically, the addition of $TBAH_2PO_4$ caused a decrease of emission intensity in the 360–400 nm range and the appearance

of a new band between 400 and 500 nm, which was saturated after the addition of 125 equivalents of the anion.¹²³ Accordingly, catenane **A** is a rare example of a turn-ON fluorescent sensor with high sensitivity and selectivity for sulfate (Fig. 9c).

Co-conformational switching of [2]catenane **A**

Although catenane **A** binds sulfate (SO_4^{2-}) strongly, it shows virtually no affinity towards bisulfate (HSO_4^-) in DMSO- d_6 (Section S5.1.6[†]). Since co-conformations of the 1 : 1 and 1 : 2 sulfate complexes differ significantly and can be easily distinguished by ¹H NMR, we decided to investigate switching between these two well-defined states using acid and base.¹²⁴ In DMSO, SO_4^{2-} is much more basic than in water, and therefore can be protonated even by rather weak acids, such as benzoic or acetic acid. However, to simplify NMR spectra, we decided to use strong, NMR-silent trifluoromethanesulfonic (triflic) acid.

In a model experiment, the addition of 2 equivalents of sulfate to the 2 mM solution of catenane **A** in DMSO- d_6 led to the formation of the 1 : 2 (receptor : anion) complex, in which each carbazole moiety binds its own sulfate. Then, the addition of 1 equivalent of triflic acid protonated one of the sulfates to the poorly bound HSO_4^- , enforcing rotation of the interlocked subunits to form the 1 : 1 complex, in which both carbazole

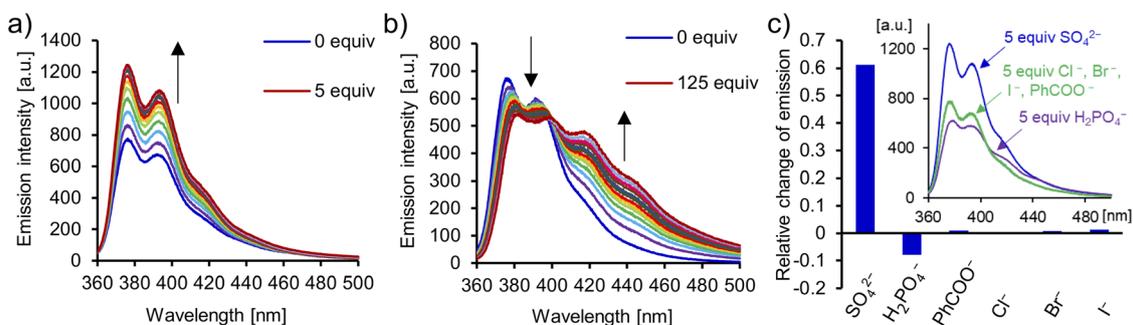


Fig. 9 Emission spectra recorded during titration of [2]catenane **A** with TBA_2SO_4 (a) and $TBAH_2PO_4$ (b) in DMSO/ H_2O 9 : 1 (ex: 350 nm). (c) Relative change of emission intensity: $(I - I_0)/I_0$ of catenane **A** at 370 nm in the presence of 5 equiv. of TBA salts. Inset: comparison of emission spectra of [2]catenane **A** in the presence of 5 equiv. of TBA salts of SO_4^{2-} , $H_2PO_4^-$, $PhCOO^-$, and halides (Cl^- , Br^- , and I^-) in DMSO/ H_2O 9 : 1 (ex: 350 nm).



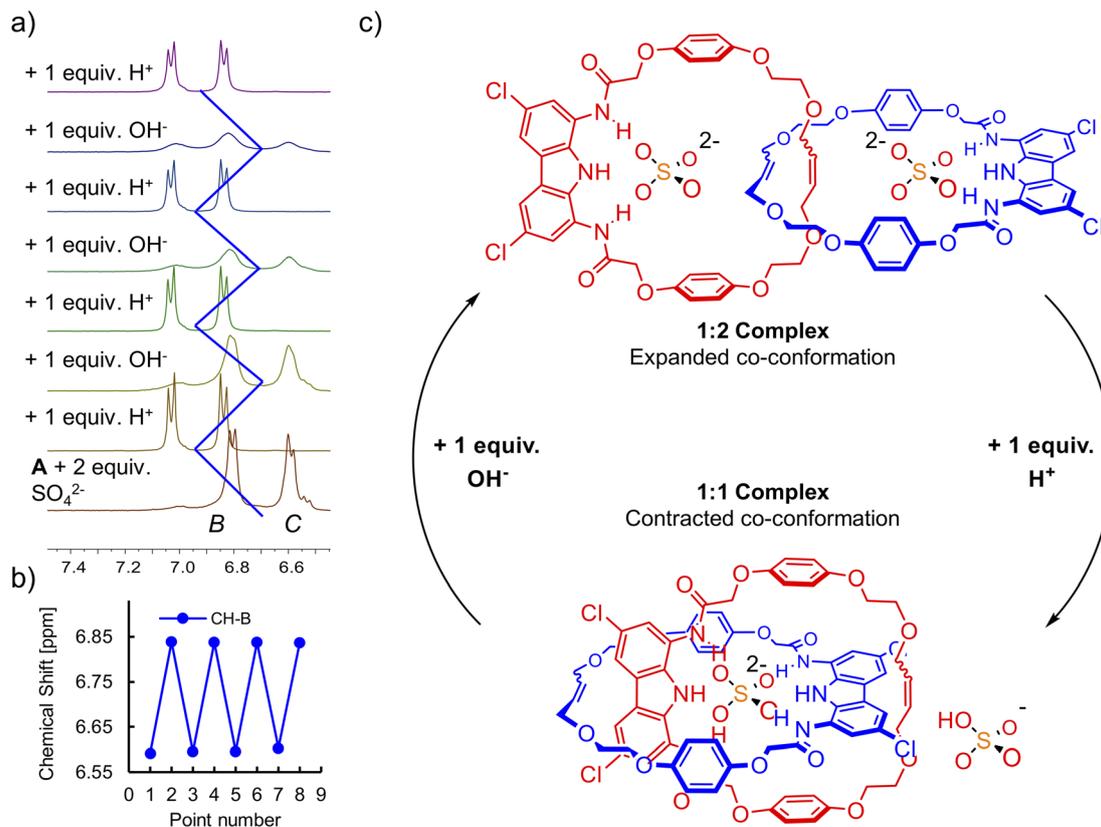


Fig. 10 (a) Acid-base triggered switching between the 1 : 1 and 1 : 2 complexes of [2]catenane A and sulfate tracked by ¹H NMR. (b) Chemical shift of signal CH–B during the ¹H NMR switching experiment. (c) Graphical representation of the switching cycle.

units bind the same anion (Fig. 10). The change of co-conformation manifested as the upfield shift of hydroquinone proton signals in ¹H NMR. Next, 1 equivalent of TBAOH was added to regenerate sulfate and trigger rotation of the subunits back to the 1 : 2 complex. This switching cycle was repeated eight times with excellent reproducibility. The switching between SO₄²⁻ with strong HB accepting capabilities and HSO₄⁻ with poor HB accepting capabilities¹²² is reminiscent of the much better established switching of cationic templates, such as Cu²⁺/Cu⁺ switching elicited by redox reactions.¹²⁵

Computational studies of [2]catenane A

Further insights into sulfate complexation and anion-induced rotary motions of catenane A were obtained through conformational analysis and molecular dynamics (MD) simulations undertaken with the MacroModel suite, using OPLS 2005/2015 and Charmm General v. 3.0.1 force fields (Sections S6.1 and S6.2†). The initial conformational search performed for catenane A revealed three major conformations (Fig. 11). In the lowest-energy conformation both diamidocarbazole moieties are situated orthogonally (Fig. 11a). One of the C=O groups of each diamidocarbazole moiety is pointing towards the center of the cavity, forming an intramolecular hydrogen bond with the carbazole NH. Additionally, it also forms hydrogen bonds with two NHs of the other diamidocarbazole, which leads to the formation of an array of 6 intramolecular hydrogen bonds. The

conformation is additionally stabilized by π -stacking interactions between the carbazole moiety of one macrocycle and the hydroquinone moieties of the other.

In the second most stable conformation the two diamidocarbazoles are situated in a parallel manner and are held together by π - π stacking interactions and four HBs (Fig. 11b). Three of these HBs are formed between the three NH groups of one diamidocarbazole and the C=O HB acceptor of the other, and the fourth HB binds together the central carbazole NH and the hydroquinone oxygen atom of the other macrocycle. This conformation is additionally stabilized by π - π stacking interactions of three hydroquinone moieties, where two hydroquinones of one macrocycle sandwich the hydroquinone ring of the other. The conformation was found to be 6.3 kcal mol⁻¹ energetically less stable than the first conformation, apparently due to the lower number of HBs.

The third conformation is not stabilized by any HBs, and is therefore 15.5 kcal mol⁻¹ higher in energy than the first one. It is only stabilized by π - π stacking interactions between two hydroquinone moieties of one macrocycle and the carbazole moiety of the other (Fig. 11c).

No other significantly different conformations were found in the 25 kcal mol⁻¹ window. However, MD simulations of the three conformations revealed that they are very dynamic, with multiple low-energy states observed even during 100 ns simulations at 25 °C. While the hydrogen-bonded cores of the two



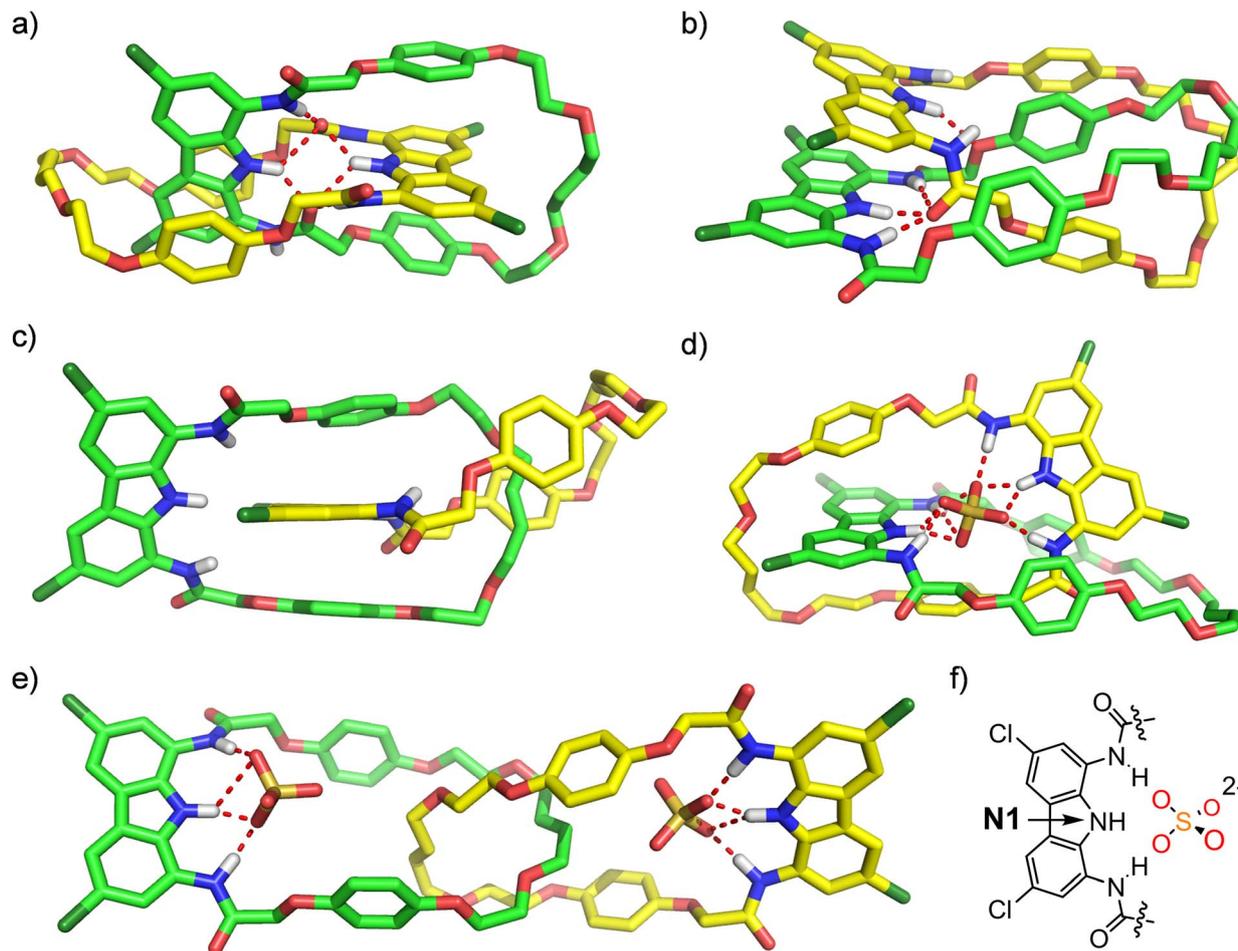


Fig. 11 Low-energy conformations (a–c) of free catenane **A** obtained from conformational search. Snapshots from MD simulations showing stable conformations of (d) $A \times SO_4^{2-}$ and (e) $SO_4^{2-} \times A \times SO_4^{2-}$. (f) Atom labels used in the discussion of MD simulations.

most stable conformations are largely preserved during the MD runs, the π - π stacking interactions are easily broken. This might be responsible for the lack of specific contacts in 1H ROESY spectrum of free catenane **A** (*vide supra*).

A similar MD analysis was performed for the 1:1 and 1:2 sulfate complexes. In the 1:1 complex, sulfate is bound inside the interlocked cavity of the catenane by nine strong hydrogen bonds, formed by all six NH donors from both diamidocarbazole moieties (Fig. 11d). It is worth noting that while sulfate binding disrupts the array of hydrogen bonds observed in the lowest-energy conformation of **A**, it does not induce any major changes in the mutual orientation of the macrocyclic components. Also, despite the breaking of 6 intramolecular hydrogen bonds, sulfate binding is still energetically favorable by approximately $79.9 \text{ kcal mol}^{-1}$. The complex is very stable over the simulation time and unlikely to undergo any major conformational changes. During the entire 100 ns run the two N1(diamidocarbazole)-S(sulfate) distances vary only between 3.2 and 3.5 Å. During this time however, the sulfate rotates in the cavity quickly (Fig. S72b†).

The co-conformation of the 1:2 complex is significantly different. Each sulfate anion is bound by a single diamidocarbazole moiety, forming 4 hydrogen bonds in total, with

either two (edge binding) or three (plane binding) oxygen atoms of the anion (Fig. 11e). Due to the electrostatic repulsion between the anions, both diamidocarbazoles are pushed away from each other to an 'expanded' co-conformation, which is in agreement with the ROESY and DOSY spectra of the complex. During the 100 ns simulation run, diamidocarbazole moieties maintain the relatively long distance of 15 to 22 Å (measured between the nitrogen atoms of the carbazole rings), as the two macrocycles move to a certain extent with respect to each other. This means that the 1:2 complex is significantly more flexible than the 1:1 complex. However, both the N1(diamidocarbazole)-S(sulfate) distances are very stable during the entire 100 ns simulation and remain in the 3.0–3.7 Å range. This means that both sulfates are tightly bound in their respective binding sites, although rotate even faster than in the 1:1 complex (Fig. S68b†).

To further investigate the conformational change between the 1:1 and 1:2 sulfate complexes, we performed MD runs for the 1:2 complex from which one sulfate anion and two random TBA cations were removed, keeping the system electrically neutral. The exemplary MD run resulted in rapid change of the catenane co-conformation, which converged to the low-energy 1:1 complex in less than 1 ns (Fig. 12). This change from the 'expanded' to the



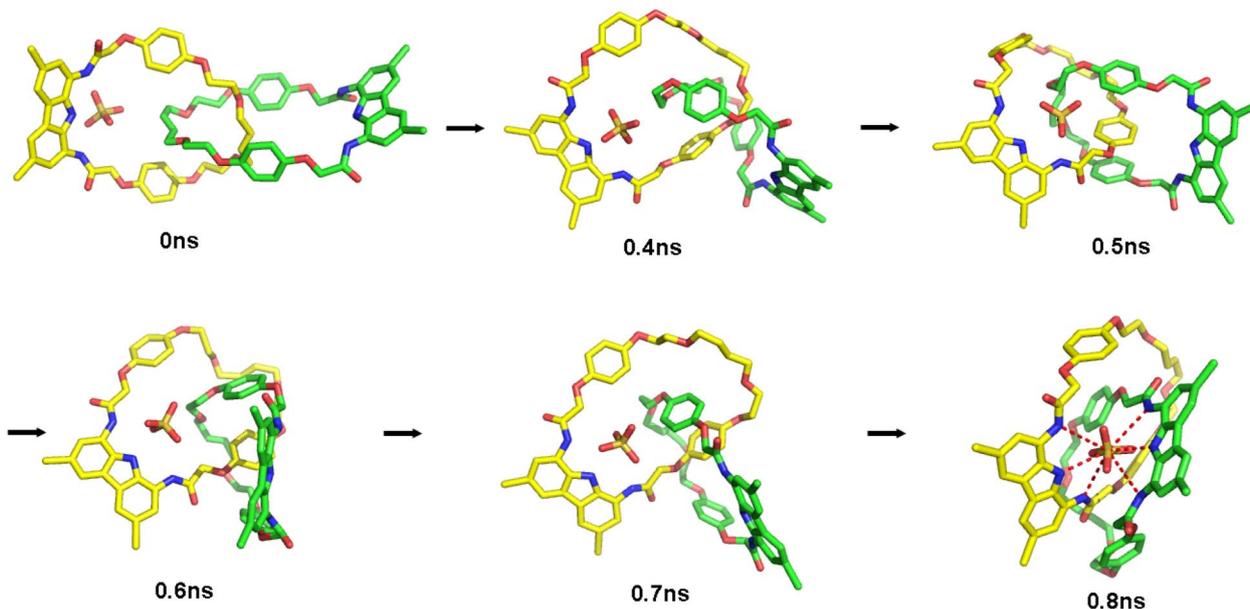


Fig. 12 Snapshots from MD runs showing switch from the 'expanded' to the 'contracted' co-conformation of catenane A.

'contracted' co-conformation, driven by chelation of the sulfate by the second diamidocarbazole moiety, leads to a drastic decrease (from *ca.* 20 Å to *ca.* 9 Å) of the distance between the N1 nitrogen atoms of the two carbazoles, as well as of the distance between one of the carbazole's N1 atoms and the sulfate's sulfur atom (from 17 Å to 3.5 Å).

Upon the transition to the 'contracted' co-conformation, the system becomes energetically and geometrically stable. In some MD runs, however, removal of one sulfate from the 1 : 2 complex resulted in a conformation, in which one of the macrocycles binds sulfate using all three NH groups, while the other uses only a single amide NH (Fig. S77[†]). As a result, the sulfate is bound more weakly than in the 'contracted' co-conformation described above, and the distance between sulfur and nitrogen of one carbazole ring remains relatively large: on average 6.5 Å. Energy data suggest that this conformation is energetically less favorable than the most stable one by approximately 30–40 kcal mol⁻¹, but might be quite stable kinetically, since no major conformational/energy changes were observed during the entire 50 ns MD run. This is because the metastable conformation is stabilized by the intramolecular HB between amide C=O and carbazole NH, and this HB has to be broken to allow transition to the most stable arrangement. It is likely therefore, that following sulfate removal from the 1 : 2 complex the resulting 1 : 1 complex becomes temporarily trapped in this high-energy, metastable state, although ultimately converges to the lowest-energy conformation. This computational result highlights challenges associated with the development of fast molecular switches based on hydrogen bonding interactions.

Conclusions

In this work, we have developed an anion-templated synthesis of electroneutral [2]catenane with remarkable affinity, selectivity,

and a strong turn-ON fluorescence response towards sulfate anion, SO₄²⁻. Anion binding studies revealed that the catenane binds the extremely hydrophilic sulfate strongly even in the presence of large excess of water, and with high selectivity over other oxoanions, notably over dihydrogen phosphate H₂PO₄⁻. Sulfate is bound inside the interlocked cavity by two diamidocarbazole moieties, forming strong, hydrogen bonded 1 : 1 complex. The excellent size and shape complementarity between the interlocked host and sulfate guest resulted in a significant increase of binding strength and selectivity when compared to the acyclic precursor, the parent macrocycle, and the acyclic receptor containing two similar diamidocarbazole moieties connected covalently by a flexible linker.

Our results indicate that by using a powerful anionic template and complementary high-affinity building blocks, it is possible to expand the scope of anion templated synthesis, which is therefore no longer limited to positively charged building blocks. Our findings showcase sulfate as a highly effective anionic template for the synthesis of MIMs and potentially also other complex supramolecular structures, such as helicates, foldamers and capsules, from electroneutral components. Moreover, diamidocarbazole has emerged from this study as a privileged building block for the anion templated synthesis, programmed self-assembly and the construction of highly sensitive fluorescent receptors.

Furthermore, sulfate may be used as a unique switchable template, able to enforce large amplitude molecular motions within an interlocked structure. Therefore, SO₄²⁻ fills the gap of missing switchable anionic templates and further expands the applications of anion templation strategies.

The unique properties of catenane A illustrate the utility of electroneutral ion binding MIMs as receptors, sensors and molecular switches, with promising future applications in extraction, transmembrane transport, and the development of responsive devices and materials.



Data availability

General information, experimental procedures, characterization data for all new compounds, raw data and fitting procedures for ^1H NMR, UV-Vis and fluorescence titration, detailed MD analysis and additional figures are available in the ESI.†

Author contributions

Krzysztof M. Bąk: conceptualization, data curation, formal analysis, investigation (synthesis, binding studies), developing methodology, validation of results, visualization, writing – original draft, writing – reviewing & editing. Bartosz Trzaskowski: formal analysis (MD simulations), investigation (MD simulations), visualization (MD simulations), writing – review & editing. Michał J. Chmielewski: conceptualization, funding acquisition, project administration, supervision, writing – original draft, writing – reviewing & editing.

Conflicts of interest

There are no conflicts to declare.

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