

# Nitrate anion templated assembly of a [2]rotaxane for selective nitrate recognition in aqueous solvent mixtures†

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**The first nitrate anion templated assembly of an interlocked molecular architecture is demonstrated through the preparation of a [2]rotaxane. Removal of the discrete nitrate anion template from the [2]rotaxane reveals an interlocked host system capable of strong and selective recognition of nitrate, in aqueous–organic solvent mixtures, over a range of more basic mono-charged oxoanions.**

The design of anion receptors capable of highly selective recognition is a key challenge for supramolecular chemistry.<sup>1</sup> Surprisingly, little attention has been paid to the selective recognition of nitrate, despite the significant importance of this anion in environmental and medical contexts. The over use of nitrate in fertilizers has led to accumulation of the anion in water courses, leading to eutrophication (excessive plant growth) and the subsequent disruption of the aquatic ecosystem.<sup>2</sup> Furthermore, increased nitrate levels has been implicated in the formation of carcinogenic nitrosamines and causes methemoglobinemia (blue-baby syndrome) in infants.<sup>3</sup> With this in mind, the development of novel approaches toward selective nitrate recognition is of importance. However, the design of nitrate receptors is challenging due to a combination of high hydration energy and low basicity of the anion, which results in a low affinity for hydrogen bonds.<sup>4</sup> The trigonal planar geometry of the anion has been exploited in a small number of tripodal, macrocyclic and cage-like host systems, which position the hydrogen bond donors in a complementary trigonal arrangement, and can recognize nitrate with modest selectivity in polar organic solvents.<sup>5</sup> However, recognition of nitrate in aqueous media by synthetic organic host molecules remains, to the best of our knowledge, elusive and high levels of nitrate selectivity have yet to be achieved.

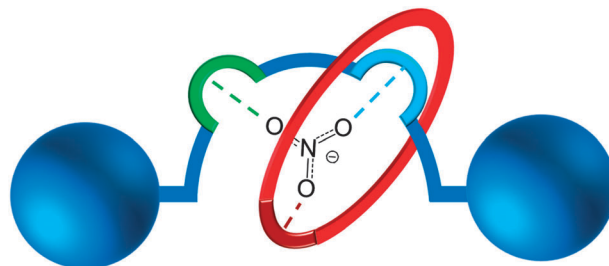
The exquisite anion guest selectivity displayed by nature in phosphate and sulfate binding proteins is accomplished through a three-dimensional convergent array of hydrogen bond donors and acceptors, arranged in an optimized geometry for recognition of the complementary oxoanion.<sup>6</sup> In our group we have developed the use of discrete

anion templates for the formation of interlocked molecular architectures, utilising chloride,<sup>7</sup> bromide<sup>8</sup> and sulfate<sup>9</sup> as the templating anions. The resulting interlocked host molecules can encapsulate anions between the interlocked components, with a high degree of selectivity for the templating anion, through convergent hydrogen bond donors reminiscent of those found in anion binding proteins in nature.

Herein we report the first example of the use of nitrate as a template for the formation of interpenetrated and interlocked molecular architectures.<sup>10</sup> Discrete nitrate anion templated pseudorotaxane formation of a suitably designed threading component within a macrocycle is demonstrated initially. Stoppering of the pseudorotaxane assembly led to the construction of a [2]rotaxane host system, which displays excellent selectivity for nitrate in aqueous–polar organic solvent mixtures over a range of other, more basic, mono-charged oxoanions.

Our strategy was to design a complementary threading component that contains two hydrogen bonding recognition sites for forming hydrogen bonds to two of the oxygen atoms of the nitrate anion. The remaining oxygen atom would thus be free to interact with a suitable hydrogen bonding motif that is integrated into a macrocycle component and facilitate the trigonal nitrate anion templated assembly of a rotaxane (Fig. 1).

The possibility of using nitrate as a pseudorotaxane templating anion was investigated initially using the asymmetrical bidentate isophthalamide-3,5-bis-amide pyridinium containing thread **2-PF<sub>6</sub>**,



**Fig. 1** Schematic representation of a nitrate templated [2]rotaxane, with two hydrogen bonding recognition sites in the axle (blue and green) and one in the macrocycle (red), forming a complementary binding site for the trigonal nitrate anion between the interlocked components.

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**Scheme 1** Assembly of nitrate templated pseudorotaxane **1·2·NO<sub>3</sub>**.



**Fig. 2**  $^1\text{H}$  NMR spectra of (a) macrocycle **1**, (b) macrocycle **1** + 1 equiv. TBANO<sub>3</sub>, (c) pseudorotaxane **1·2·NO<sub>3</sub>**, (d) thread **2·PF<sub>6</sub>** in  $d_6$ -acetone (500 MHz). For atom labels see Scheme 1.

(Scheme 1), terminated with non-interacting hexyl chains (see ESI† for synthesis). Macrocycle **1**, incorporating an isophthalamide motif to coordinate to the nitrate anion, and hydroquinone groups to provide secondary stabilization through aromatic donor–acceptor interactions with the electron deficient pyridinium moiety of the thread, was prepared according to literature procedures.<sup>11</sup>

Initial  $^1\text{H}$  NMR pseudorotaxane assembly studies were undertaken in  $d_6$ -acetone (Fig. 2). Addition of TBANO<sub>3</sub> to a solution of macrocycle **1** led to downfield shifts of the isophthalamide amide protons and internal proton b, indicating coordination of the oxoanion within the amide binding cleft. Upon addition of one equivalent of thread **2·PF<sub>6</sub>**, downfield shifts of the thread protons 1, 4 and amides were observed, which is indicative of nitrate binding, and demonstrates that both the isophthalamide and pyridinium isophthalamide moieties are involved in hydrogen bonding to the nitrate anion. Importantly, the observed upfield perturbation and increased splitting of the macrocycle hydroquinone protons c and d

is characteristic of aromatic donor–acceptor interactions between the electron rich hydroquinone groups in the macrocycle and the positively charged electron deficient pyridinium group in the thread, confirming the formation of the pseudorotaxane **1·2·NO<sub>3</sub>** (Scheme 1).

The successful formation of the nitrate templated pseudorotaxane **1·2·NO<sub>3</sub>** suggested that the synthesis of a [2]rotaxane would be possible using nitrate templation, *via* a stoppering strategy. To this end bis-azide functionalized axle precursor **3·NO<sub>3</sub>** was prepared (see ESI†) for a copper(i) catalysed azide–alkyne (CuAAC) click stoppering reaction with a suitable alkyne functionalized stopper. Synthesis of rotaxane **5·NO<sub>3</sub>** was achieved by mixing 1 equiv. of **3·NO<sub>3</sub>** with 1.1 equiv. of macrocycle **1** in 4 : 1  $\text{CH}_2\text{Cl}_2$ –acetone to form the initial pseudorotaxane assembly. Addition of catalytic  $\text{Cu}(\text{CH}_3\text{CN})_4\text{PF}_6$  and 2.2 equiv. of stopper alkyne **4** gave a crude product whose  $^1\text{H}$  NMR spectrum revealed that the rotaxane was formed in approximately 35% yield (Scheme 2). Purification by size exclusion chromatography and silica gel chromatography gave rotaxane **5·NO<sub>3</sub>** in an isolated yield of 24%, which was fully characterized by  $^1\text{H}$ ,  $^{13}\text{C}$  and ROESY NMR, and high resolution electrospray mass spectrometry (see ESI†). Importantly, an analogous reaction conducted in the absence of nitrate with **3·PF<sub>6</sub>** gave no evidence of rotaxane formation, which highlights the crucial templating role of the nitrate anion.<sup>12</sup>

The  $^1\text{H}$  spectra of rotaxane **5·NO<sub>3</sub>**, macrocycle **1** and axle precursor **3·NO<sub>3</sub>** are compared in Fig. 3. It is noteworthy that the



**Scheme 2** Synthesis of rotaxane **5·NO<sub>3</sub>** *via* nitrate templation.





**Fig. 3**  $^1\text{H}$  NMR spectra of (a) macrocycle **1**, (b) rotaxane **5-NO<sub>3</sub>** (c) axle precursor **3-NO<sub>3</sub>** in 1 : 1  $\text{CDCl}_3$ - $\text{CD}_3\text{OD}$  (500 MHz). For atom labels see Scheme 2.

macrocycle hydroquinone protons c and d are split and shifted upfield, which are diagnostic of the aromatic donor-acceptor interactions between the hydroquinones in the macrocycle and the axle pyridinium motif, and confirms the interlocked nature of the rotaxane. Further evidence is obtained in the  $^1\text{H}$  NMR ROESY spectrum, in which multiple through space interactions between the macrocycle and axle are observed (see ESI†). Anion exchange to the non-coordinating hexafluorophosphate salt, in preparation for anion recognition studies, was achieved by washing a solution of the rotaxane **5-NO<sub>3</sub>** in  $\text{CH}_2\text{Cl}_2$  with aqueous  $\text{NH}_4\text{PF}_6$ . The anion recognition properties of rotaxane **5-PF<sub>6</sub>** were investigated using  $^1\text{H}$  NMR titration experiments in a competitive aqueous-polar organic solvent mixture of 45 : 45 : 10  $\text{CDCl}_3$ - $\text{CD}_3\text{OD}$ - $\text{D}_2\text{O}$ . Upon binding of nitrate the internal pyridinium axle proton 1 is strongly perturbed upfield ( $\Delta\delta = 0.19$  ppm after 10 equiv.) which is diagnostic of the nitrate anion binding within the rotaxane cavity. Addition of other oxoanions led to smaller perturbations of axle proton 1 (Table 1). WinEQNMR<sup>213</sup> analysis of the titration data, monitoring proton 1, enabled the determination of 1 : 1 stoichiometric anion association constants shown in Table 1.

The trigonal nitrate anion was found to bind strongly within the rotaxane host's complementary tridentate hydrogen bond donor binding cavity in this competitive aqueous-polar organic solvent mixture. Impressive selectivity for nitrate was observed over a range of other more basic oxoanions: the pseudo-trigonal hydrogen carbonate anion bound considerably more weakly, and acetate and dihydrogenphosphate resulted in very weak binding, which in the case of acetate was too weak to be quantified.

The selectivity for nitrate over acetate is particularly impressive, given that acetate is  $10^5$  times more basic,<sup>14</sup> and reflects in part the geometric complementarity of the rotaxane binding cavity for

nitrate. To the best of our knowledge this is the first example of a synthetic anion receptor capable of this level of nitrate selectivity in aqueous solvent mixtures over other mono-charged oxoanions.

The spherical chloride anion, which is of comparable size to nitrate but lacks the trigonal geometric preference, was found to bind with a similar affinity to nitrate. Addition of the larger  $\text{Br}^-$  anion led to very small perturbations of the internal binding cavity proton 1, and the binding could not be quantified. Analogous titrations with axle precursor **3-PF<sub>6</sub>** and  $\text{NO}_3^-$ ,  $\text{AcO}^-$  and  $\text{Cl}^-$ , revealed no binding of the two oxoanions in the same solvent mixture, but significantly stronger binding of  $\text{Cl}^-$  ( $150 \text{ M}^{-1}$ ). This serves to further highlight the importance of the interlocked cavity in remarkably enhancing the strength of nitrate binding with respect to the non-interlocked pyridinium axle, and the crucial role the unique three-dimensional binding domain plays in achieving the shape selectivity for the trigonal nitrate anion, by encapsulating the oxoanion guest.

In summary, we have demonstrated the first example of nitrate anion templation for the formation of interlocked molecular architectures, through the preparation of a [2]rotaxane. Removal of the nitrate template affords an interlocked host system which displays unprecedented binding affinity and selectivity for nitrate over other oxoanions of significantly higher basicity, in a competitive aqueous-organic solvent mixture. The exploitation of nitrate as a templating reagent in the synthesis and development of interlocked anion receptors and sensors is continuing in our laboratories.

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**Table 1** Anion association constants of rotaxane **5-PF<sub>6</sub>** in 45 : 45 : 10  $\text{CDCl}_3$ - $\text{CD}_3\text{OD}$ - $\text{D}_2\text{O}$ , free energy values and complexation induced chemical shift changes of proton 1

Anion <sup>a</sup>	$\text{NO}_3^-$	$\text{HCO}_3^-$	$\text{H}_2\text{PO}_4^-$	$\text{AcO}^-$	$\text{Cl}^-$	$\text{Br}^-$
$K_a \text{ (M}^{-1}\text{)}$	430	100	50	— <sup>c</sup>	490	— <sup>c</sup>
$\Delta G \text{ (kJ mol}^{-1}\text{)}$	−15.0	−11.4	−9.7	— <sup>c</sup>	−15.3	— <sup>c</sup>
$\Delta\delta(1)^d$	0.19	0.10	0.07	0.02	0.21	0.02

$T = 298 \text{ K}$ . <sup>a</sup> Anions added as TBA salts, except for  $\text{HCO}_3^-$  which was added as the TEA salt. <sup>b</sup> Calculated using chemical shift data of proton 1. Errors estimated to be <10%. <sup>c</sup> Binding too weak to be quantified. <sup>d</sup> Chemical shift change of proton 1 after addition of 10 equiv. of anion.

