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### ARTICLE TYPE

#### Lanthanide-cation templated synthesis of rotaxanes

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Received (in XXX, XXX) Xth XXXXXXXX 20XX, Accepted Xth XXXXXXXX 20XX DOI: 10.1039/b000000x

<sup>5</sup> The first lanthanide cation-templated synthesis of an interlocked structure is demonstrated through an interpenetrated assembly between a pyridine *N*-oxide threading component coordinating to a lanthanide cation complexed within a macrocycle. Stoppering of the pseudo-<sup>10</sup> rotaxane assembly allows for preparation of the [2]rotaxane.

The development of new innovative strategic templating routes for the construction of mechanically interlocked molecules with potential molecular switching and machine-like nanotechnological applications is of intense current research 15 interest.<sup>1</sup> In addition to their dynamic behaviour, rotaxanes and catenanes can also be designed to contain topologically unique three-dimensional cavities for molecular recognition and sensing applications.<sup>2</sup> Although transition metal cation, and to a lesser extent alkali and alkaline earth metal cation, templates have been 20 imaginatively exploited in the synthesis of rotaxanes and catenanes,<sup>3</sup> to the best of our knowledge, the ultilization of lanthanides as potential templating reagents for interlocked molecular framework assembly has not been demonstrated.<sup>4</sup> This is somewhat surprising given the usefulness of luminescent <sup>25</sup> lanthanide complexes in imaging and assay.<sup>5</sup> We describe herein the first lanthanide-cation templated synthesis of a rotaxane.

Adapting our recently reported sodium and barium cation templated synthesis of a rotaxane using the pyridine N-oxide motif,<sup>6</sup> the synthetic strategy undertaken for the preparation of the <sup>30</sup> target lanthanide-[2]rotaxane is shown in Scheme 1. A kinetically stable lanthanide complexed DOTA cyclen derivative integrated into a macrocyclic structural framework, forms initially a pseudo-rotaxane assembly with an appropriately functionalized pyridine *N*-oxide threading component, where the pyridine *N*-oxide ligand

<sup>35</sup> serves to satisfy the lanthanide cation's coordination sphere. Subsequent stoppering of the interpenetrated assembly produces the novel lanthanide containing interlocked structure.

The synthetic procedure used for preparation of the lanthanide macrocyclic component **4a** and **4b** of the target [2]rotaxane <sup>40</sup> system and the bis-azide functionalised pyridine *N*-oxide thread

component is outlined in Scheme 2.

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† Electronic Supplementary Information (ESI) available. Experimental details for synthetic and spectroscopic proceedures, and additional characterisation. See DOI: 10.1039/b000000x/



Scheme 1. Schematic representation for the preparation of the target lanthanide-[2]rotaxane.

Two of the nitrogen atoms of cyclen were selectively alkylated with *tert*-butyl bromoacetate to give bis-ester cyclen **1**, using established procedures.<sup>7</sup> The bridging unit **2** was prepared by <sup>55</sup> condensing two equivalents of chloroacetylchloride with the corresponding bis-amine derivative<sup>8</sup> (see SI). Reaction of the 1,7-DO2A diester (**1**) and the bis-amide compound **2** in acetonitrile under basic conditions afforded macrocycle **3**, which was isolated in 70 % yield after purification by recrystallization from <sup>60</sup> diethyl ether. Cleavage of the tert-butyl ester groups in **3** followed by complexation of the resulting product with either lutetium or europium lanthanide trifluoromethanesulfonate salts in methanol, produced the lanthanide complexes **4a** and **4b** in a quantitative yield. The condensation reaction between 3,5-bis-<sup>65</sup> chlorocarbonyl pyridine and two equivalents of 3bromopropylamine hydrobromide in dry CH<sub>2</sub>Cl<sub>2</sub> in the presence ChemComm

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Scheme 2. Synthesis of a) lanthanide macrocycles complexes and b) bisazide pyridine N-oxide thread component.



*Figure 1*. High resolution electrospray ionization mass spectrum of [2]rotaxane **8b** (top) with theoretical isotope model for [M-TfO+Na]<sup>2+</sup> (bottom).

of triethylamine gave the bis-bromo amide pyridine derivative **5** in 50 % yield. Azide nucleophilic substitution followed by oxidation using oxone afforded the bis-azide pyridine *N*-oxide compound **6** in 80 % yield.

- <sup>10</sup> The synthesis of the target lanthanide [2]rotaxanes was achieved by a copper (I) catalysed cycloaddition azide-alkyne (CuAAC) click stoppering reaction as shown in Scheme 3. A solution of the appropriate lanthanide macrocyclic complex (**4a** and **4b**), bis-azide pyridine *N*-oxide axle precursor **6** and two
- <sup>15</sup> equivalents of alkyne-functionalised terphenyl stopper **7** in dichloromethane in the presence of a catalytic amount of  $Cu(CH_3CN)_4PF_6$  was stirred at room temperature for 48 hours. After purification using size exclusion chromatography, the rotaxanes **8a** and **8b** were isolated in 20 % yield, and were





characterized by high-resolution electrospray mass spectroscopy (Figure 1) and  ${}^{1}$ H NMR.<sup>9</sup>

A comparison of the <sup>1</sup>H NMR spectra of the [2]rotaxane 8a, <sup>25</sup> macrocycle 4a and stoppered axle is shown in Figure 2. The upfield shifts of the macrocycle hydroquinone protons observed with 8a are diagnostic of aromatic donor – acceptor stacking interactions between the electron rich hydroquinone groups of the macrocycle and the electron deficient pyridine *N*-oxide axle

<sup>30</sup> motif. Two-dimensional <sup>1</sup>H-<sup>1</sup>H-ROESY spectroscopy was also used to confirm the interlocked nature of rotaxane **8a** with through-space correlations between protons in the respective axle and macrocycle components being identified. (see Figure SI6 in the Supporting Information).<sup>10</sup>An equimolar solution of cyclen Published on 01 August 2013. Downloaded by Universidade Federal do Parana on 05/08/2013 10:16:51



*Figure 2.* Partial <sup>1</sup>H NMR spectra in CDCl<sub>3</sub>/CD<sub>3</sub>OD 1:1 at 293 K of (a) macrocycle **4a**, (b) [2] rotaxane **8a**, and (c) axle. For atom labels see scheme 3.

macrocycle **3** and pyridine *N*-oxide derivative **6** in  $CD_2Cl_2$ 5 revealed no upfield perturbations of the macrocycle's hydroquinone protons which suggests the contribution of aromatic donor – acceptor interactions to the overall mechanical bond formation process is minimal and highlights the crucial templating role of the lanthanide cation<sup>11</sup>.

- <sup>10</sup> The luminescence from the europium centre was also used to probe the structure of macrocycle **4b** and rotaxane **8b** in 1:1 CH<sub>2</sub>Cl<sub>2</sub>:CH<sub>3</sub>OH and in 1:1 CD<sub>2</sub>Cl<sub>2</sub>:CD<sub>3</sub>OD. In the former solvent mixture, the observed luminescence lifetime of **8b** was found to be 0.94 ms, while in deuterated media the luminescence lifetime
- <sup>15</sup> increased to 1.18 ms. Understanding solvation of even simple systems in binary solvent mixtures is not straightforward, but given the residual charge on the lanthanide centre, it is reasonable to assume that any inner sphere solvent at the europium ion will be comprised of methanol rather than DCM molecules. As such it
- $_{20}$  is possible to estimate q, the number of inner sphere solvent molecules bound to the lanthanide, from the equation q =  $2.4(\tau_{CH3OH}{}^{-1} \tau_{CD3OD}{}^{-1} 0.125 0.0375x)$ , where x is the number of exchangeable amide N-H oscillators close to the metal centre.  $^{12}$  In this case, the calculated value of q is 0 if we assume
- <sup>25</sup> that all four amide N-H oscillators contribute (including the two on the axle), which certainly equates to exclusion of methanol from the inner coordination sphere in **8b**, and suggests that the pyridine *N*-oxide oxygen atom acts as an axial donor to the lanthanide. The luminescence lifetime of **4b** in 1:1
  <sup>30</sup> CH<sub>2</sub>Cl<sub>2</sub>:CH<sub>3</sub>OH is shorter and a biexponential fit gives 0.20 and
- <sup>30</sup> CH<sub>2</sub>Cl<sub>2</sub>:CH<sub>3</sub>OH is shorter and a biexponential fit gives 0.20 and 0.63 ms, indicating solvation at the europium centre in contrast to **8b**.
- In summary, the first lanthanide-cation templated synthesis of an <sup>35</sup> interlocked structure has been demonstrated. A pyridine *N*-oxide ligand threading component coordinates to a lutetium or europium lanthanide complexed DOTA cyclen motif which is itself incorporated into a macrocycle. The novel lanthanide [2]rotaxanes are prepared through stoppering of the <sup>40</sup> pseudorotaxane assemblies.
- F.Z. acknowledges the Ministry of Education of Spain for a Postdoctoral contract (Programa Nacional de Movilidad de Recursos Humanos del Plan Nacional I+D+I 2008-2011). O. A. B. thanks the European Research Council for funding under the

<sup>45</sup> European Union's Framework Program (FP7/2007-2013) ERC Advanced Grant Agreement n<sup>o</sup> 267426. M. J. L. thanks the EPSRC for a DTA studentship.

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  - 9 The by-products from the reaction were non-interlocked macrocycle and axle.
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