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Dynamic lanthanide exchange between quadruplestranded cages: the effect of ionic radius differences on kinetics and thermodynamics†

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Seven different $[Ln_2L_4]^{2-}$ (Ln = La, Nd, Eu, Tb, Er, Tm and Lu) lanthanide-based quadruple-stranded helicates are here reported and transmetalation among the pre-assembled cages is studied. Combining two homonuclear helicates, $[Ln_2^AL_4]^{2-}$ and $[Ln_2^BL_4]^{2-}$, leads to the formation of a mixture of homo- and heteronuclear systems due to ion exchange. This dynamic behaviour is studied by electrospray ionization mass spectrometry (ESI-MS) both qualitatively and quantitatively, allowing one to gain information on the thermodynamics and on the kinetics of the process. The rate of the Ln ion exchange is strongly correlated with the difference in the lanthanide effective ionic radius (Δ EIR). Upon increasing the Δ EIR, the kinetic constants grow exponentially: from the minimum to the maximum value of the Δ EIR, the kinetic constants of the forward and backward reactions increase by three orders of magnitude. In contrast, the equilibrium constant is the same for all the $[Ln_2^AL_4]^{2-}/[Ln_2^BL_4]^{2-}$ couples, showing that the transmetalation is mainly entropy driven towards a statistical mixture and not affected by the Δ EIR.

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Introduction

Among metallo-supramolecular systems, lanthanide (Ln)-based architectures are attracting increasing attention due to their remarkable optical and magnetic properties. However, with Ln³+ ions, the design and control of the final supramolecule can be challenging. This is a direct consequence of the inner nature of the 4f orbitals: the ligand-field effects are small, and the coordination chemistry of these ions is similar across the entire series with the coordination number and geometry very variable and basically dependent on the steric pro-

perties of the ligand. There is, however, a steady variation of the effective ionic radius (EIR) across the series, which is the so-called "lanthanide contraction". Although the radius difference (Δ EIR) is quite small (ca. 0.20 Å between La³⁺ and Lu³⁺ and ca. 0.02 Å between two consecutive lanthanides), it can have important chemical consequences on the nature and features of supramolecular complexes. For instance, the Δ EIR coupled to the use of multicompartmental ligands has been used to control the self-assembly of heterobimetallic helicates, 1,2 while the Δ EIR effect has been shown to lead to lanthanide-selective self-assembly through multivalency and cooperativity.3,4 Furthermore, Δ EIR effects were proved to strongly affect the formation of the more stable thermodynamic product and the final metallo-supramolecular structure⁵⁻¹¹ as well as the rate of conversion between different aggregates. 11 More subtle effects have also been reported, such as variation of the twist angles in pseudo-helical Ln tripodal complexes12 and fine-tuning of the helicity and the helical winding in oligo(salamo)-based metal LZn₃Ln helicates. 13

In the past few decades, advances in the coordination chemistry of multinuclear compounds have been exploited to drive the self-assembly of many new discrete supramolecular motifs. Due to the nature of the metal-ligand interactions, many of these systems have a dynamic character with reversible association and dissociation able to generate complex mixtures. To unveil such dynamic behaviours, it is necessary to

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fully understand, control and design their functional properties. Mass spectrometry (MS) is a powerful tool to study such supramolecules 14-16 for determining both the ensembles' structure in terms of composition, shape, and size, and their dynamic behaviour and formation pathways. 17-25 However, only a relatively small number of studies report MS quantitative analyses for metallo-supramolecular complexes. 17-19,23-27 Among the dynamic processes displayed by supramolecular complexes, post-synthetic ion exchange is particularly important since it provides new pathways to synthesize heterometallic systems or species that cannot be achieved by direct metalligand coordination. 20,21,24-36 Despite the importance of this synthesis strategy, little rigorous works have been reported on the kinetics³³ and the thermodynamics^{29,30} of the ion exchange process for supramolecular complexes.

Oligo-β-diketones are versatile ligands for the preparation of metallo-supramolecular architectures^{37,38} ranging from metallocycles, cages and MOFs, 39-43 to triple- and quadruplestranded helicates 44-53 and even to interlocked structures. 54-57 Recently, we reported two $[Ln_2L_4]^{2-}$ quadruple-stranded helicates (Ln = La, Eu and Gd) that show dynamic and adaptive helicity reorganization due to a guest-to-host chirality transfer. 44 Herein, by using a closely related bis-β-diketone ligand, we report a larger series of seven [Ln₂L₄]²⁻ quadruple-stranded helicates (Ln = La, Nd, Eu, Tb, Er, Tm and Lu), Fig. 1, that once assembled undergo transmetalation. The thermodynamics and kinetics of the dynamic ion exchange have been studied and the role of the Ln^{3+} ΔEIR has been evaluated. The process has been qualitatively and quantitatively characterized by time-dependent electrospray ionization mass spectrometry (ESI-MS). Mixing a series of two homonuclear $[Ln^{A}_{2}L_{4}]^{2-}$ and [Ln^B₂L₄]²⁻ with increasing Ln³⁺ ΔEIR always leads to the formation of a statistical mixture of homo- and heteronuclear helicates due to the Ln exchange. All the studied systems have

an equilibrium constant close to K = 4. The Ln³⁺ Δ EIR, hence, does not affect the thermodynamics of the process that is mainly governed by statistical factors and entropy driven. On the other hand, we demonstrate that the rate of the dynamic ion exchange is Ln³⁺ radii dependent, Fig. 1b. The kinetic constants of the forward and backward reactions revealed an exponential trend correlated with the Ln3+ ΔEIR of the two homonuclear pre-assembled cages, Fig. 1c. This fundamental study provides new tools and guidelines to investigate dynamic processes in metallo-supramolecular ensembles, and for the precise preparation and control of lanthanide-based mixed coordination-driven systems.

Results and discussion

[Ln₂L₄]²⁻ cage synthesis and characterization

The bis-β-diketone ligand was synthesized using a two-step protocol: (i) Ulmann coupling followed by (ii) Claisen condensation, and characterized as described in the ESI† (NMR Fig. S1-S4† and single crystal XRD Fig. S6†). The lantern-like cages with the general formula [Ln2L4]2- were prepared by adding an ethanol solution of the deprotonated ligand to an ethanol solution of a Ln3+ salt. Seven different Ln3+ ions were used to self-assemble $[Ln_2L_4]^{2-}$ cages where Ln = La, Nd, Eu, Tb, Er, Tm and Lu. Two bases, tetraethylammoniun hydroxide and dicyclohexylamine, were employed. Cages with countercation NEt4+ were used for ESI-MS and XRD analyses, while cages with the DCHA⁺ (dicyclohexylammonium) countercation were used for NMR studies due to their higher solubility. A metal: ligand molar ratio of 1:2.5 leads to the formation of the pure compound in a good yield (Table S1†). All the cages show very clean ESI-MS spectra with only one signal ascribed to the [Ln₂L₄]²⁻ double-negative charged species. The experi-

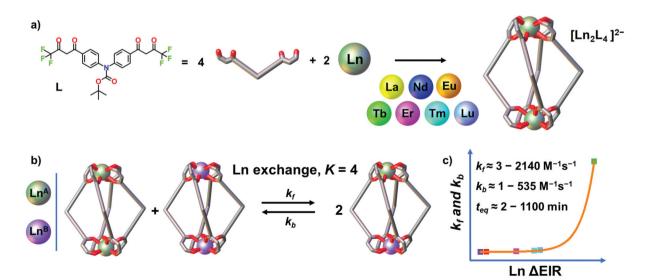
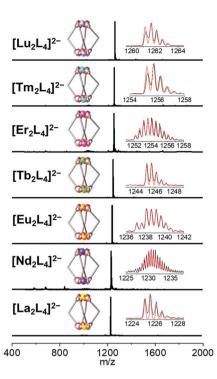


Fig. 1 (a) Self-assembly of seven $[Ln_2L_4]^{2-}$ cages (Ln = La, Nd, Eu, Tb, Er, Tm and Lu). (b) Dynamic Ln^{3+} ion exchange equilibrium between two preassembled cages and (c) exponential trend of the kinetic constants depending on the Ln Δ EIR.



Eu, Tb, Er, Tm and Lu). Insets: experimental (black) and simulated (red) isotopic patterns.

mental isotopic pattern of each species is in good agreement with the calculated one, Fig. 2. The systems were also characterized by ¹H-NMR and Fig. 3a shows the ligand (H₂L, L²⁻) and [La₂L₄]²⁻ cage spectra. Upon deprotonation, all the ligand signals undergo upfield shifting. The strongest effect is observed for the proton H1, in the α position of the β-diketonate moiety, shifting from ca. 7.1 to ca. 5.9 ppm, while the tert-butyl protons (H4) are barely perturbated. After La³⁺ coordination, the cage shows only a single set of signals with a total of four resonances, in agreement with the ligand C_2 symmetry and the mean C_4 symmetry of the quadruple-stranded $[La_2L_4]^{2-}$ architecture. H1, the H atom closer to the metal center, is downfield shifted to ca. 6.2 ppm. Diffusion-ordered NMR spectroscopy (DOSY) was performed to provide dimensional information. Two distinct bands were observed and ascribed to the cage and to the DCHA+ cation. The calculated hydrodynamic diameter for $[La_2L_4]^{2-}$ is 23.5 Å. The NMR spectra for the Eu and Lu systems (Fig. S5†) are also reported in the ESI†, while the Nd, Tb, Er and Tm analogue spectra showed very poor resolution due to these metals' high magnetic moments.

For the Eu cage, it was possible to obtain single crystals suitable for XRD from the slow evaporation of mother liquors (ethanol). Structural determination ultimately confirmed the cage assembly and the formation of helicate systems (Fig. 3b). Both the right-handed $(P, \Delta\Delta)$ and left-handed $(M, \Lambda\Lambda)$ helicates are present in a 1:1 ratio as an enantiomeric pair in the

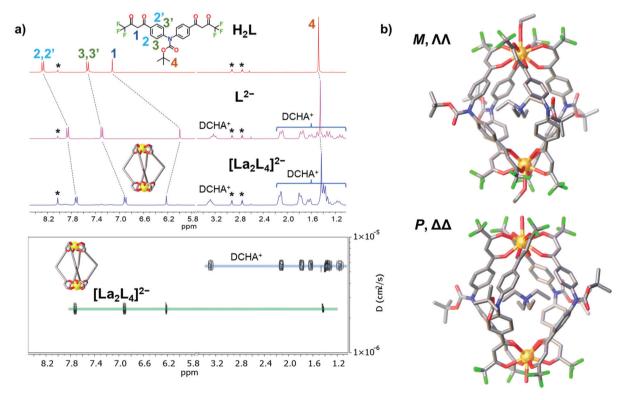


Fig. 3 (a) 1 H-NMR spectra (25 $^{\circ}$ C, 300 MHz, DMF-d₇) of ligand H₂L, deprotonated ligand L²⁻ and the cage $[La_{2}L_{4}]^{2-}$. DOSY of $[La_{2}L_{4}]^{2-}$. * = DMF. (b) Crystal structure of the Eu helicate, M and P forms in the asymmetric unit. H atoms and external NEt_4^+ ions omitted for clarity. Color code: C, grey; O, red; N, blue; F, green; Eu, orange.

asymmetric unit (Fig. S7 and S8†), likewise to what was previously observed for a similar quadruple-stranded helicate.⁴⁴ Each helicate hosts a NEt₄⁺ counterion in its cavity. The Eu-O distances are in the range of 2.486-2.639 Å and the Eu...Eu distances are 12.054 Å and 12.199 Å for the two crystallographic independent helicates, respectively. The H···H distances of the tert-butyl groups of opposite ligands are close to 23 Å. This value is in good agreement with the hydrodynamic diameter calculated from the DOSY experiment (23.5 Å) for the lanthanum cage. Conversely to previous findings,44 the Eu ions do not display an octa-coordination mode since they are nona-coordinated. The Eu ions are tetrakis-chelated by four β-diketonato groups and the ninth site is occupied by a water or ethanol molecule. In particular, with the space group being centrosymmetric, the unit cell contains four helicates: two are ethanol coordinated (one M and one P helicate), the other two are water coordinated (one M and one P helicate), Fig. S7b.† The nona-coordination can be a consequence of the crystallization environment in the presence of good coordinating solvents such as ethanol and water. ESI-MS analysis did not show any adduct with water or ethanol. DFT studies were performed both on octa- and nona-coordinated La cages. The comparison of the X-ray nona-coordinated Eu helicate and the DFT-optimized octa-coordinated La helicate shows that the ligand arrangement in the two systems is very similar as highlighted by the overlay of the two structures (Fig. S10†). Moreover, the solvent molecule binding energy has also been determined: -7.4 and -8.5 kcal mol⁻¹ for water and ethanol, respectively (see the ESI for details, Table S3†). This suggests that these molecules can be easily lost, as for instance during electrospray ionization.

Qualitative and quantitative ESI-MS of the Ln³⁺ ion dynamic

Time-dependent ESI-MS analysis was used to study both qualitatively and quantitatively possible dynamic processes between two pre-assembled cages, $[Ln_2^AL_4]^{2-}$ and $[Ln_2^BL_4]^{2-}$. Preliminary experiments were performed by mixing two equimolar solutions of the $[Eu_2L_4]^{2-}$ and $[Tb_2L_4]^{2-}$ cages in a 1:1 ratio in a sealed vial at 50 °C, and monitoring their ESI-MS spectra at increasing time, Fig. 4b. At time zero, only the patterns ascribed to the two homometallic cages are present. After 60 minutes, the isotopic pattern of the heterometallic cage $[EuTbL_4]^{2-}$ appears, as confirmed by comparison with the calculated one (Fig. S19†), showing that the two homometallic cages undergo ion exchange, forming a mixture of heteronuclear and homonuclear systems. The intensity of the Ln-mixed cage increases over time to the detriment of the homometallic ones. In order to apply ESI-MS to monitor the concentration of the different species at different times, we hypothesized that the $[Ln_2L_4]^{2-}$ species have comparable ionization efficiencies. To confirm this hypothesis, the ESI-MS spectrum of a solution containing equimolar amounts of the two homometallic cages $[Eu_2L_4]^{2-}$ and $[Tb_2L_4]^{2-}$ was repetitively measured just after mixing. If the hypothesis is correct, the two cages must exhibit MS spectra that indicate an equal relative amount. The relative

amount was deduced according to eqn (1). Then, eqn (2) can be easily used to derive the molarity of the species (see the ESI† for details).

$$\%[Ln^{A}Ln^{B}L_{4}] = \frac{A[Ln^{A}Ln^{B}L_{4}]}{\sum A[Ln^{A}Ln^{B}L_{4}]} \cdot 100$$
 (1)

$$|Ln^{A}Ln^{B}L_{4}| = \frac{\% \left[Ln^{A}Ln^{B}L_{4}\right] \cdot (|Ln_{2}^{A}L_{4}|_{0} + |Ln_{2}^{B}L_{4}|_{0})}{100}$$
 (2)

where %[LnALnBL4] is the relative amount of the cage, A [Ln^ALn^BL₄] is the integrated area of the species isotopic pattern, and |LnALnBL4| is the molarity of the homonuclear cage if $Ln^A = Ln^B$, and the heteronuclear cage if $Ln^A \neq Ln^B$. $|Ln^{A}_{\ 2}L_{4}|_{0}$ and $|Ln^{B}_{\ 2}L_{4}|_{0}$ are the initial molar concentrations of the homometallic species, respectively. Application of eqn (1) always gave values of about 50% for the relative amounts of the $[Eu_2L_4]^{2-}$ and $[Tb_2L_4]^{2-}$ cages (Table S4†), as expected for an equimolar mixture of the two cages, confirming that they have similar ionization efficiencies.

The ESI-MS spectra of Fig. 4b and Fig. S18† were used to derive the relative amount and concentration of the different cages during the ion exchange. At equilibrium, a statistical mixture of homometallic (25%) and heterometallic (50%) cages is obtained, Fig. 5b and Table S6.† Then, the same approach was applied to study the ion exchange with other $[Ln_{2}^{A}L_{4}]^{2-}/[Ln_{2}^{B}L_{4}]^{2-}$ couples, namely $[Tm_{2}L_{4}]^{2-}/[Lu_{2}L_{4}]^{2-}$, $[Eu_2L_4]^{2-}/[Tm_2L_4]^{2-}, [La_2L_4]^{2-}/[Eu_2L_4]^{2-}, [Nd_2L_4]^{2-}/[Er_2L_4]^{2-}$ and $[La_2L_4]^{2-}/[Lu_2L_4]^{2-}$. Homonuclear cages bearing different Ln³⁺ ions were chosen in order to have Ln³⁺ ions spread across the lanthanide series, providing a range of effective ionic radii (EIR) and ionic radius differences (Δ EIR), as detailed in Table 1. The couples were also chosen in order to give similar values of ΔEIR but the Ln^{3+} ions were combined across the lanthanide series, such as $[Tm_2L_4]^{2-}/[Lu_2L_4]^{2-}$ and $[Eu_2L_4]^{2-}/[Lu_2L_4]^{2-}$ $[Tb_2L_4]^{2-}$ (Δ EIR 0.02 and 0.03 Å, respectively) or $[La_2L_4]^{2-}$ $[Eu_2L_4]^{2-}$ and $[Nd_2L_4]^{2-}/[Er_2L_4]^{2-}$ ($\Delta EIR~0.11$ and 0.12 Å, respectively). The use of ESI-MS allows one to investigate also $[Ln_2^AL_4]^{2-}/[Ln_2^BL_4]^{2-}$ couples that cannot be analyzed using NMR due to poor resolution. The only important constraint in the choice of the $[Ln^{A}_{2}L_{4}]^{2-}/[Ln^{B}_{2}L_{4}]^{2-}$ couple is that there must be no significant overlaps between the isotopic patterns of the homonuclear and heteronuclear cages. The same protocol used for the couple $[Eu_2L_4]^{2-}/[Tb_2L_4]^{2-}$ was applied. First, the two homonuclear cages were tested to verify whether they possess the same ionization efficiencies (Table S6, and Fig. S11†) in order to draw quantitative information from ESI-MS. Second, time-dependent analyses were performed to follow the dynamic ion exchange (Fig. 5, Tables S5-S10†). Fig. 5 clearly shows that all the $[Ln_2^AL_4]^{2-}/[Ln_2^BL_4]^{2-}$ couples give a statistical mixture of homonuclear and heteronuclear cages, but the times to reach it are different.

Thermodynamics and kinetics analysis of the Ln3+ ion dynamic exchange

Starting from two homometallic cages with a 1:1 ratio, two homometallic and one heterometallic cages with a ratio 1:1:2

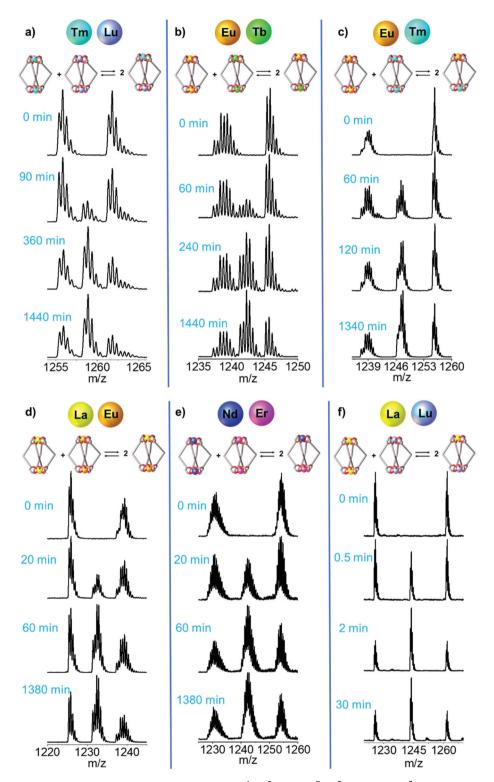


Fig. 4 Time-dependent ESI-MS spectra of an equimolar solution of $[Ln^A_2L_4]^2$ and $[Ln^B_2L_4]^2$. Dynamic Ln^{3+} ion exchange between the couples with increasing ionic radius difference (Δ EIR): (a) $[Tm_2L_4]^2$ / $[Lu_2L_4]^2$, (b) $[Eu_2L_4]^2$ / $[Tb_2L_4]^2$, (c) $[Eu_2L_4]^2$ / $[Tm_2L_4]^2$ / $[Tm_2L_4]^2$ / $[Eu_2L_4]^2$ / $[Eu_2L_4]^2$ / $[Eu_2L_4]^2$ - $[Eu_2L_$

are obtained, eqn (3). Based on a statistical analysis,⁵⁹ we can anticipate that the equilibrium constant will be K = 4. Indeed, at equilibrium, the experimental concentrations are all very

close to the following values: $|Ln_2^AL_4| = |Ln_2^BL_4| = 5 \times 10^{-6} \text{ M}$ and $|Ln_4^AL_1^BL_4| = 1 \times 10^{-5} \text{ M}$. All the $Ln_4^AL_1^BL_2^A$ ion exchange reactions, hence, have the same equilibrium constant (Table 1).

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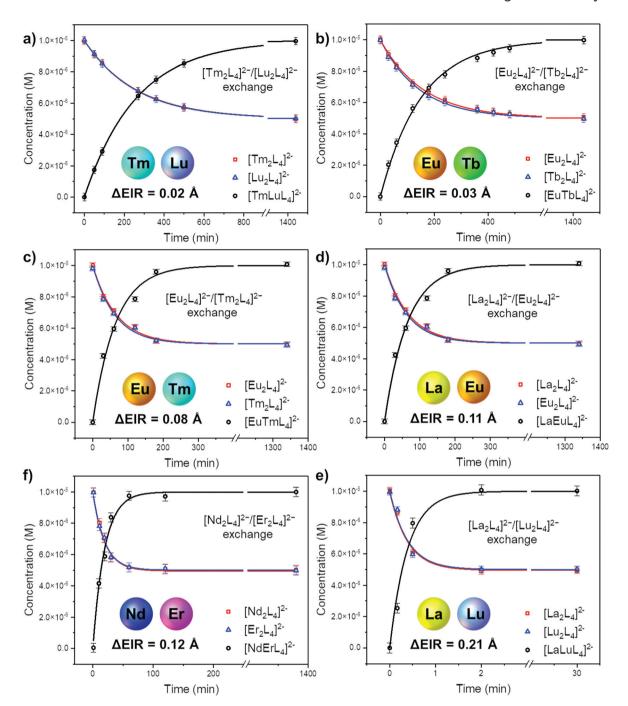


Fig. 5 Concentration over time during the Ln^{3+} ion exchange reaction as obtained by time-dependent ESI-MS of equimolar $[Ln^{A}_{2}L_{4}]^{2-}/[Ln^{B}_{2}L_{4}]^{2-}$ mixtures. Kinetic profiles according to eqn (4) for the homometallic cages (red and blue lines) and eqn (5) for the heterometallic cages (black lines). Fitting details are reported in Fig. S15-S17, S20-S22, S25-S27, S30-S32, S35-37, and S40-S42.

This suggests that the ion exchange process is essentially entropy driven and hence governed by statistical factors. If the enthalpic contribution is negligible, we can infer that the different homometallic and heterometallic cages have similar stabilities across the lanthanide series.

$$\left[Ln_2^AL_4\right]^{2-} + \left[Ln_2^BL_4\right]^{2-\frac{k_f}{\overleftarrow{k_b}}} 2\left[Ln^ALn^BL_4\right]^{2-} \tag{3}$$

In contrast, the kinetics of the Ln³⁺ ion exchange reaction are very different. Fig. 5 highlights that by increasing the ionic radius difference (Δ EIR) between the involved Ln³⁺ ions, the time to reach the statistical mixture decreases. Some previous works¹⁷⁻¹⁹ treated the kinetics of dynamic ligand exchange between pre-assembled metallo-supramolecular architectures as first- or second-order direct reactions without considering the reversible nature of the equilibrium. This simplification

Table 1 Ln³⁺ ion EIR and Δ EIR for the couples $[Ln^A_2L_4]^2$ - $[Ln^B_2L_4]^2$ -and K, k_f , k_b and t_{eq} for the Ln^{3+} ion exchange reactions

	La ³⁺	Nd^{3+}	$Eu^{3^{+}}$	Tb^{3+}	Er^{3+}	$\mathrm{Tm}^{3^{+}}$	$Lu^{3^{+}}$
$\operatorname{Ln}^{3+}\operatorname{EIR}^{a}(\mathring{A})$	1.18	1.12	1.07	1.04	1.00	0.99	0.97
$[Ln^{A}_{2}L_{4}]^{2-}/[Ln^{B}_{2}L_{4}]^{2-}$	ΔEIR (Å)		$k_{\rm f} ({\rm M}^{-1} {\rm s}^{-1})$		$k_{\rm b} ({\rm M}^{-1} {\rm s}^{-1})$		t _{eq} (min)
$[Tm_2L_4]^{2-}/[Lu_2L_4]^{2-}$	0.02	3	.97 ± 0.35	3.20 ± 0.01	$0.8 \pm 0.$	01	1199.3 ± 4.7
$[Eu_2L_4]^{2-}/[Tb_2L_4]^{2-}$	0.03	4	$.05 \pm 0.26$	5.49 ± 0.07	1.37 ± 0	0.02	700.2 ± 9.3
$[Eu_2L_4]^{2-}/[Tm_2L_4]^{2-}$	0.08	3	$.98 \pm 0.29$	12.78 ± 0.46	3.20 ± 0.11		300.9 ± 10.8
$\left[\operatorname{La_{2}L_{4}}\right]^{2-}/\left[\operatorname{Eu_{2}L_{4}}\right]^{2-}$	0.11	3	$.98 \pm 0.29$	30.09 ± 1.02	7.52 ± 0.25		127.7 ± 4.3
$[Nd_2L_4]^{2-}/[Er_2L_4]^{2-}$	0.12	4	$.00 \pm 0.40$	42.77 ± 1.46	10.69 ± 0.36		89.8 ± 3.1
$[La_2L_4]^{2-}/[Lu_2L_4]^{2-}$	0.21	4	$.02 \pm 0.35$	2140.31 ± 166.50	535.08	± 41.63	1.8 ± 0.1

^a For octa-coordinated Ln³⁺ ions.⁵⁸

can be adopted only under specific conditions (see below), otherwise it can lead to misleading results (see the ESI, Fig. S12†). As a matter of fact, the dynamic exchange reaction (ion or ligand) is a reversible reaction and its kinetics must be treated as a pair of forward and backward reactions (kinetic constants $k_{\rm f}$ and $k_{\rm b}$, respectively). These two reactions occur simultaneously and are related to the equilibrium constant by $K = k_f/k_b$. Only if $k_f \gg k_b$ (high value for K), the reverse reaction can be neglected, and the kinetics analysis simplifies to a rate law for a direct reaction. However, in the cases here discussed K = 4 and k_b will be one quarter of k_f : the backward reaction cannot be neglected. The kinetics of the dynamic Ln3+ ion exchange has been treated as two opposing second-order reactions (see the ESI† for the detailed treatment). This leads to eqn (4) and (5) that give the concentrations of the homometallic and heterometallic species over time, respectively.

$$\left| L n_2^A L_4 \right| = \left| L n_2^B L_4 \right| = \frac{a_0}{2} (1 + e^{-k_f t 2 a_0}) \tag{4}$$

$$|Ln^{A}Ln^{B}L_{4}| = a_{0}(1 - e^{-k_{f}t2a_{0}})$$
 (5)

where a_0 is the initial concentration of the homometallic cages, k_f is the kinetic constant of the forward reaction, and tis the time. Moreover, the performed kinetic analysis allows one to estimate t_{eq} , the time to reach the equilibrium, *i.e.* the statistical mixture, eqn (6) (see the ESI† for details).

$$t_{\rm eq} = \frac{1}{2a_0 k_{\rm f}} \ln \frac{1}{\alpha} \tag{6}$$

where α is an arbitrary small number (see the ESI† for details). For calculations of $t_{\rm eq}$, we assumed α = 0.01, which corresponds to a homometallic cage concentration that is 1% higher than the equilibrium concentration.

Fitting the experimental data with eqn (4) and (5), Fig. 5, allowed us to determine k_f and then k_b for the different $[Ln_{2}^{A}L_{4}]^{2-}/[Ln_{2}^{B}L_{4}]^{2-}$ couples involved in the ion dynamic exchange reaction. Table 1 summarizes the obtained results as the average of the values for the two homometallic cages and the heterometallic one (Table S9 \dagger). The more the Δ EIR between the two Ln^{3+} ions increases, the more k_f (and k_b) increases and consequently $t_{\rm eq}$ decreases. For instance, $k_{\rm f}$ is 3.20 \pm 0.01 ${\rm M}^{-1}$ ${\rm s}^{-1}$ ($t_{\rm eq}$ 1199.3 ± 4.7 min) for the couple ${\rm [Tm_2L_4]^{2-/[Lu_2L_4]^{2-}}}$

with the minimum $\Delta EIR = 0.02$ Å. For the exchange reaction $[Eu_2L_4]^{2-}/[Tb_2L_4]^{2-}$, $\Delta EIR = 0.03 \text{ Å and } k_f \text{ is } 5.49 \pm 0.07 \text{ M}^{-1} \text{ s}^{-1}$ $(t_{\rm eq} 700.2 \pm 9.2 \text{ min})$, for $[Eu_2L_4]^{2-}/[Tm_2L_4]^{2-}$, $\Delta EIR = 0.08 \text{ Å}$ and $k_{\rm f}$ is 12.78 \pm 0.46 M⁻¹ s⁻¹ ($t_{\rm eq}$ 300.9 \pm 10.8 min), for $[{\rm La_2L_4}]^{2-}/$ $[Eu_2L_4]^{2-}$, $\Delta EIR = 0.11$ Å and k_f is 30.09 ± 1.02 M^{-1} s⁻¹ (t_{eq} 127.7 ± 4.3 min), and for $[Nd_2L_4]^{2-}/[Er_2L_4]^{2-}$, Δ EIR = 0.12 Å and k_f is 42.77 ± 1.46 M⁻¹ s⁻¹ (t_{eq} 89.8 ± 3.1 min). The couple $[La_2L_4]^{2-}/[Lu_2L_4]^{2-}$ with the highest $\Delta EIR = 0.21$ Å shows a dramatic increase of k_f (2140.31 \pm 166.50 M^{-1} s⁻¹) with a $t_{\rm eq}$ value of only 1.8 \pm 0.1 minutes.

Two considerations deserve to be mentioned. First, mononuclear Ln β-diketonate complexes are known to be kinetically labile and to easily undergo ligand exchange. 60-63 For instance, Eu β-diketonate solutions were found to give a mixture of the tetrakis $[EuL_4]^-$, tris $[EuL_3]$, bis $[EuL_2]^+$, and mono $[EuL]^{2+}$ β-diketonate systems depending on the solvent and concentration.⁶⁴ Second, among Ln metallosupramolecular systems, it has been previously demonstrated that Ln tetrahedra architectures display fast conversion to helicates when using large ions, the middle Ln ones give moderate conversion while the smaller ones show slow conversion rates. 11 Hence, based on these considerations, it can be hypothesized that the Ln transmetalation may involve the partial dissociation of the two homometallic $[Ln_2L_4]^{2-}$ helicates to allow the ion exchange and that the overall kinetics may be governed by the homometallic helicate with the slower rate of dissociation even if we did not observe any undissociated species by either NMR or ESI-MS analyses. However, the experimental data do not evidence a clear relationship between the exchange rate and the effective ionic radius of one of the two Ln3+ ions of the [LnA2L4]2-/[LnB2L4]2- couple in terms of more or less labile homometallic systems and hence faster or slower dissociation. Indeed, one may argue that if the exchange kinetics was governed by the dissociation rate of the helicate with the smaller ion, the expected k_f trend should be: $[La_2L_4]^{2-}/[Lu_2L_4]^{2-} \approx [Tm_2L_4]^{2-}/[Lu_2L_4]^{2-} > [Eu_2L_4]^{2-}/[Tm_2L_4]^{2-}$ $> [Nd_2L_4]^{2-}/[Er_2L_4]^{2-} > [Eu_2L_4]^{2-}/[Tb_2L_4]^{2-} > [La_2L_4]^{2-}/[Eu_2L_4]^{2-}.$ On the other hand, an overall kinetic governed by the dissociation rate of the helicate with the bigger ion should give the following order: $[La_2L_4]^{2-}/[Lu_2L_4]^{2-} \approx [La_2L_4]^{2-}/[Eu_2L_4]^{2-} >$ $[\mathrm{Nd}_2\mathrm{L}_4]^{2-}/[\mathrm{Er}_2\mathrm{L}_4]^{2-} > [\mathrm{Eu}_2\mathrm{L}_4]^{2-}/[\mathrm{Tb}_2\mathrm{L}_4]^{2-} \approx [\mathrm{Eu}_2\mathrm{L}_4]^{2-}/[\mathrm{Tm}_2\mathrm{L}_4]^{2-} >$ $[Tm_2L_4]^{2-}/[Lu_2L_4]^{2-}$. Neither a more complex lability pattern like that displayed by hydrated Ln³⁺ ions for water dissociation rates

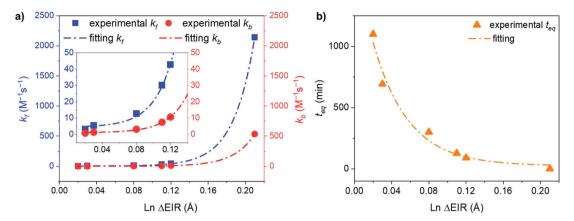


Fig. 6 Exponential trends of (a) k_f and k_b , and (b) t_{eq} for the Ln³⁺ ion exchange kinetics correlated with the Ln Δ EIR. Inset: magnification of the 0.01-0.14 Å ∆EIR region. Fitting details are reported in Fig. S43.†

showing a maximum for the central ions and lower values at the beginning and end of the Ln series⁶⁵ allows one to rationalize the exchange rates here observed. Indeed, experimental data show the following kinetic constant order: $[La_2L_4]^{2-}/[Lu_2L_4]^{2-}\gg$ $\lceil Nd_2L_4 \rceil^{2-} / \lceil Er_2L_4 \rceil^{2-} > \lceil La_2L_4 \rceil^{2-} / \lceil Eu_2L_4 \rceil^{2-} > \lceil Eu_2L_4 \rceil^{2-} / \lceil Tm_2L_4 \rceil^{2-} > \lceil Eu_2L_4 \rceil$ $[Eu_2L_4]^{2-}/[Tb_2L_4]^{2-} > [Tm_2L_4]^{2-}/[Lu_2L_4]^{2-}$, suggesting a more complex mechanism that possibly involves both the homometallic helicates as suggested by the second-order kinetics.

As a matter of fact, plotting k_f and k_b versus the ΔEIR , Fig. 6a, reveals an exponential dependence of the kinetic constant's values related to the Ln3+ ionic radius difference of the two homonuclear cages participating in the ion exchange. An exponential relationship was also found by plotting t_{eq} versus Δ EIR, Fig. 6b. The collected data allow one to span across the entire Ln series, studying the exchange reaction with [Ln^A₂L₄]²⁻/[Ln^B₂L₄]²⁻ couples having different or very close values of ΔEIR (for instance: $[La_2L_4]^{2-}/[Eu_2L_4]^{2-}$ $[Nd_2L_4]^{2-}/[Er_2L_4]^{2-}$ or $[Tm_2L_4]^{2-}/[Lu_2L_4]^{2-}$ and $[Eu_2L_4]^{2-}/[Lu_2L_4]^{2-}$ $[Tb_2L_4]^{2-}$), choosing Ln³⁺ ions from the beginning, the middle, and the end of the Ln series. The analyzed combinations show that the values of ΔEIR between Ln^A and Ln^B can be used as an empirical parameter to predict the exchange reaction rate. This means that based on the Δ EIR it is possible to know the exchange rate and that, on working with small and intermediate values of Δ EIR (slow/moderate Ln exchange rates), it is experimentally possible to precisely control the homometallic/ heterometallic mixture composition by simply tuning the mixing time and then stopping the reaction by evaporating the solvent or embedding the system in some polymers to prepare a solid solution. This fine control is particularly appealing for the preparation of mixed Ln systems to exploit their luminescence properties (for instance Eu/Tb systems).

Conclusions

In summary, a series of quadruple-stranded dinuclear helicates $[Ln_2L_4]^{2-}$ were prepared using a bis- β -diketone ligand

and seven different Ln3+ ions (Ln = La, Nd, Eu, Tb, Er, Tm and Lu) and fully characterized by NMR, ESI-MS, DFT and single crystal XRD. The dynamic behavior of these systems was studied qualitatively and quantitatively by ESI-MS, providing information on how fast the ion exchange process between two pre-assembled homonuclear cages is completed, leading to a statistical mixture of homonuclear and heteronuclear cages. The thermodynamics of the transmetalation reaction is not affected by the Ln3+ ionic radius difference (ΔEIR) of the two homonuclear pre-assembled cages participating in the ion exchange. This suggests that the process is mainly entropy driven and that both the homometallic and heterometallic cages have similar stabilities across the Ln series. In contrast, the rate of Ln3+ ion exchange is strongly correlated with the Δ EIR, which can be used as an empirical parameter to know the exchange rate. The more the ΔEIR increases, the more the exchange rate increases. Both the forward and the backward kinetic constants ($k_{\rm f}$ and $k_{\rm b}$) and the equilibration time (t_{eq}) show an exponential trend related to the Ln^{3+} Δ EIR. We envisage that this fundamental study will advance the supramolecular chemistry of Ln coordination-driven cages in terms of the synthesis, properties, and applications, in particular, of Ln-mixed systems where the control of the composition is paramount for the final functional properties such as magnetism and luminescence. In fact, rigorous knowledge of the ion exchange kinetics will allow the preparation of Ln-mixed systems with precise control over their composition in terms of homonuclear and heteronuclear systems by simply tailoring the mixing time of different pre-assembled cages.

Author contributions

M.R.^a conceptualization and supervision. M.R.^a, M.R.^b, A.C., L.B., R.S., S.C. and J.T. investigation. M.R. a and S.C formal analysis. L.A. and G.H.C funding acquisition. M.R. a writing - original draft. All authors writing - review & editing.

Conflicts of interest

There are no conflicts to declare.

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References

- N. André, R. Scopelliti, G. Hopfgartner, C. Piguet and J.-C.
 G. Bünzli, Discriminating between lanthanide ions: self-assembly of heterodimetallic triple-stranded helicates, *Chem. Commun.*, 2002, 214–215.
- 2 N. André, T. B. Jensen, R. Scopelliti, D. Imbert, M. Elhabiri, G. Hopfgartner, C. Piguet and J. C. G. Bünzli, Supramolecular recognition of heteropairs of lanthanide ions: A step toward self-assembled bifunctional probes, *Inorg. Chem.*, 2004, 43, 515–529.
- 3 A. M. Johnson, M. C. Young, X. Zhang, R. R. Julian and R. J. Hooley, Cooperative thermodynamic control of selectivity in the self-assembly of rare earth metal-ligand helices, *J. Am. Chem. Soc.*, 2013, 135, 17723–17726.
- 4 X.-Z. Li, L.-P. Zhou, L.-L. Yan, Y.-M. Dong, Z.-L. Bai, X.-Q. Sun, J. Diwu, S. Wang, J.-C. Bünzli and Q.-F. Sun, A supramolecular lanthanide separation approach based on multivalent cooperative enhancement of metal ion selectivity, *Nat. Commun.*, 2018, **9**, 547.
- 5 Y. Wu, S. Morton, X. Kong, G. S. Nichol and Z. Zheng, Hydrolytic synthesis and structural characterization of lanthanide-acetylacetonato/hydroxo cluster complexes A systematic study, *Dalton Trans.*, 2011, **40**, 1041–1046.
- 6 S. Zebret, C. Besnard, G. Bernardinelli and J. Hamacek, Thermodynamic discrimination in the formation of tetranuclear lanthanide helicates, *Eur. J. Inorg. Chem.*, 2012, 2012, 2409–2417.
- 7 K. P. Carter, K. E. Thomas, S. J. A. Pope, R. J. Holmberg, R. J. Butcher, M. Murugesu and C. L. Cahill, Supramolecular assembly of molecular rare-earth-3,5dichlorobenzoic acid-2,2':6',2"-terpyridine materials: Structural systematics, luminescence properties, and magnetic Behavior, *Inorg. Chem.*, 2016, 55, 6902–6915.
- 8 A. Vuillamy, S. Zebret, C. Besnard, V. Placide, S. Petoud and J. Hamacek, Functionalized triptycene-derived tripodal ligands: Privileged formation of tetranuclear cage assemblies with larger Ln(III), *Inorg. Chem.*, 2017, **56**, 2742–2749.

- 9 H.-Y. Wong, W. T. K. Chan and G.-L. Law, Assembly of lanthanide(III) cubanes and dimers with single-molecule magnetism and photoluminescence, *Inorg. Chem.*, 2018, 57, 6893–6902.
- 10 C. Chen and A. Zhang, Radii-dependent self-assembly polynuclear lanthanide complexes as catalysts for CO2 transformation into cyclic carbonates, *New J. Chem.*, 2021, 45, 20155–20163.
- 11 K. H. Yim, C. T. Yeung, M. R. Probert, W. T. K. Chan, L. E. Mackenzie, R. Pal, W. T. Wong and G. L. Law, Helicate-to-tetrahedron transformation of chiral lanthanide supramolecular complexes induced by ionic radii effect and linker length, *Commun. Chem.*, 2021, 4(116), 1–10.
- 12 S. Mizukami, H. Houjou, M. Kanesato and K. Hiratani, Adjustment of twist angles in pseudo-helical lanthanide complexes by the size of metal ions, *Chem. Eur. J.*, 2003, **9**, 1521–1528.
- 13 S. Sairenji, S. Akine and T. Nabeshima, Lanthanide contraction for helicity fine-tuning and helix-winding control of single-helical metal complexes, *Dalton Trans.*, 2016, 45, 14902–14906.
- 14 H. Wang, C. Guo and X. Li, Multidimensional mass spectrometry assisted metallo-supramolecular chemistry, CCS Chem., 2021, 3161–3184.
- 15 Z. Qi, T. Heinrich, S. Moorthy and C. A. Schalley, Gas-phase chemistry of molecular containers, *Chem. Soc. Rev.*, 2014, 44, 515–531.
- 16 E. Kalenius, M. Groessl and K. Rissanen, Ion mobility—mass spectrometry of supramolecular complexes and assemblies, *Nat. Rev. Chem.*, 2019, 3, 4–14.
- 17 Y.-R. Zheng and P. J. Stang, Direct and quantitative characterization of dynamic ligand exchange between coordination-driven self-assembled supramolecular polygons, *J. Am. Chem. Soc.*, 2009, **131**, 3487–3489.
- 18 S. Sato, Y. Ishido and M. Fujita, Remarkable stabilization of M 12 L 24 spherical frameworks through the cooperation of 48 Pd(II)–pyridine interactions, *J. Am. Chem. Soc.*, 2009, 131, 6064–6065.
- 19 L. Wang, B. Song, S. Khalife, Y. Li, L.-J. Ming, S. Bai, Y. Xu, H. Yu, M. Wang, H. Wang and X. Li, Introducing seven transition metal ions into terpyridine-based supramolecules: self-assembly and dynamic ligand exchange study, *J. Am. Chem. Soc.*, 2020, 142, 1811–1821.
- 20 M. Rancan, J. Tessarolo, P. L. Zanonato, R. Seraglia, S. Quici and L. Armelao, Self-assembly of a constitutional dynamic library of Cu(II) coordination polygons and reversible sorting by crystallization, *Dalton Trans.*, 2013, 42, 7534–7538.
- 21 B. Akhuli, L. Cera, B. Jana, S. Saha, C. A. Schalley and P. Ghosh, Formation and transmetalation mechanisms of homo- and heterometallic (Fe/Zn) trinuclear triple-stranded side-by-side helicates, *Inorg. Chem.*, 2015, **54**, 4231–4242.
- 22 B. Jana, L. Cera, B. Akhuli, S. Naskar, C. A. Schalley and P. Ghosh, Competitive transmetalation of first-row transition-metal ions between trinuclear triple-stranded side-by-side helicates, *Inorg. Chem.*, 2017, **56**, 12505–12513.

Research Article

- 23 E. O. Bobylev, D. A. Poole, B. de Bruin and J. N. H. Reek, Selective formation of Pt12L24 nanospheres by ligand design, Chem. Sci., 2021, 12, 7696-7705.
- 24 E. O. Bobylev, B. de Bruin and J. N. H. Reek, Catalytic formation of coordination-based self-assemblies by halide impurities, Inorg. Chem., 2021, 60, 12498-12505.
- 25 E. O. Bobylev, D. A. Poole, B. de Bruin and J. N. H. Reek, How to prepare kinetically stable self-assembled Pt12L24 nanocages while circumventing kinetic traps, Chem. - Eur. J., 2021, 27, 12667-12674.
- 26 F. J. Rizzuto, M. Kieffer and J. R. Nitschke, Quantified structural speciation in self-sorted CoII6L4 cage systems, Chem. Sci., 2018, 9, 1925-1930.
- 27 C. Bravin, A. Guidetti, G. Licini and C. Zonta, Supramolecular cages as differential sensors for dicarboxylate anions: Guest length sensing using principal component analysis of ESI-MS and 1H-NMR raw data, Chem. Sci., 2019, 10, 3523-3528.
- 28 X. Z. Li, L. P. Zhou, S. J. Hu, L. X. Cai, X. Q. Guo, Z. Wang and Q. F. Sun, Metal ion adaptive self-assembly of photoactive lanthanide-based supramolecular hosts, Chem. Commun., 2020, 56, 4416-4419.
- 29 S. Akine, T. Taniguchi, T. Saiki and T. Nabeshima, Ca²⁺and Ba²⁺-selective receptors based on site-selective transmetalation of multinuclear polyoxime-zinc(II) complexes, J. Am. Chem. Soc., 2005, 127, 540-541.
- 30 S. Akine, T. Taniguchi and T. Nabeshima, Helical metallohost-guest complexes via site-selective transmetalation of homotrinuclear complexes, J. Am. Chem. Soc., 2006, **128**, 15765-15774.
- 31 Y.-B. Dong, P. Wang, J.-P. Ma, X.-X. Zhao, H.-Y. Wang, B. Tang and R.-O. Huang, Coordination-driven nanosized lanthanide "molecular lantern" with tunable luminescent properties, J. Am. Chem. Soc., 2007, 129, 4872-4873.
- 32 M. E. Carnes, M. S. Collins and D. W. Johnson, Transmetalation of self-assembled, supramolecular complexes, Chem. Soc. Rev., 2014, 43, 1825-1834.
- 33 D. Zare, Y. Suffren, H. Nozary, A. Hauser, C. Piguet, D. Zare, D. Nozary, P. rof DrC. Piguet, D. Suffren and D. Hauser, Controlling lanthanide exchange in triple-stranded helicates: A way to optimize molecular light-upconversion, Angew. Chem., Int. Ed., 2017, 56, 14612-14617.
- 34 J. R. Shakirova, E. V. Grachova, V. V. Gurzhiy, S. K. Thangaraj, J. Jänis, A. S. Melnikov, A. J. Karttunen, S. P. Tunik and I. O. Koshevoy, Heterometallic clustercapped tetrahedral assemblies with postsynthetic modification of the metal cores, Angew. Chem., Int. Ed., 2018, 57, 14154-14158.
- 35 H. Ube, K. Endo, H. Sato and M. Shionoya, Synthesis of hetero-multinuclear metal complexes by site-selective redox switching and transmetalation on a homo-multinuclear complex, J. Am. Chem. Soc., 2019, 141, 10384-10389.
- 36 X. Hu, J. Chai, C. Zhang, J. Lang, S. P. Kelley, S. Feng, B. Liu, D. A. Atwood and J. L. Atwood, Biomimetic selfassembly of Co II-seamed hexameric metal-organic nanocapsules, J. Am. Chem. Soc., 2019, 141, 9151-9154.

- 37 J. K. Clegg, F. Li and L. F. Lindoy, Oligo-β-diketones as versatile ligands for use in metallo-supramolecular chemistry: Recent progress and perspectives, Coord. Chem. Rev., 2022, 455, 214355.
- 38 R. W. Saalfrank, H. Maid and A. Scheurer, Supramolecular coordination chemistry: The synergistic effect of serendipity and rational design, Angew. Chem., Int. Ed., 2008, 47, 8794-
- 39 J. K. Clegg, K. Gloe, M. J. Hayter, O. Kataeva, L. F. Lindoy, B. Moubaraki, J. C. McMurtrie, K. S. Murray and D. Schilter, New discrete and polymeric supramolecular architectures derived from dinuclear (bis-β-diketonato) copper(II) metallocycles, Dalton Trans., 2006, 3977-3984.
- 40 J. K. Clegg, S. S. Iremonger, M. J. Hayter, P. D. Southon, R. B. Macquart, M. B. Duriska, P. Jensen, P. Turner, K. A. Jolliffe, C. J. Kepert, G. V. Meehan and L. F. Lindoy, Hierarchical self-assembly of a chiral metal-organic framework displaying pronounced porosity, Angew. Chem., Int. Ed., 2010, 49, 1075-1078.
- 41 M. Rancan, A. Dolmella, R. Seraglia, S. Orlandi, S. Quici and L. Armelao, A templating guest sorts out a molecular triangle from a dimer-trimer constitutional dynamic library, Chem. Commun., 2012, 48, 3115-3117.
- 42 J. K. Clegg, F. Li, K. A. Jolliffe, L. F. Lindoy, G. V. Meehan, S. Parsons, P. A. Tasker and F. J. White, Hierarchical assembly of discrete copper(II) metallo-structures from preassembled dinuclear (bis-β-diketonato)metallocycles and flexible difunctional co-ligands, Dalton Trans., 2013, 42, 14315-14323.
- 43 M. Rancan, J. Tessarolo, M. Casarin, P. L. Zanonato, S. Quici and L. Armelao, Double level selection in a constitutional dynamic library of coordination driven supramolecular polygons, Inorg. Chem., 2014, 53, 7276-7287.
- 44 M. Rancan, J. Tessarolo, A. Carlotto, S. Carlotto, M. Rando, L. Barchi, E. Bolognesi, R. Seraglia, G. Bottaro, M. Casarin, G. H. Clever and L. Armelao, Adaptive helicity and chiral recognition in bright europium quadruple-stranded helicates induced by host-guest interaction, Cell Rep. Phys. Sci., 2022, 3, 100692.
- 45 G. Han, Y. Zhou, Y. Yao, Z. Cheng, T. Gao, H. Li and P. Yan, Preorganized helical chirality controlled homochiral selfassembly and circularly polarized luminescence of a quadruple-stranded Eu₂L₄ helicate, Dalton Trans., 2020, 49, 3312-3320.
- 46 Y. B. Tan, Y. Okayasu, S. Katao, Y. Nishikawa, F. Asanoma, M. Yamada, J. Yuasa and T. Kawai, Visible circularly polarized luminescence of octanuclear circular Eu(III) helicate, J. Am. Chem. Soc., 2020, 142, 17653-17661.
- 47 N. Suko, H. Itamoto, Y. Okayasu, N. Okura and J. Yuasa, Helix-mediated over 1 nm-range chirality recognition by ligand-to-ligand interactions of dinuclear helicates, Chem. Sci., 2021, 12, 8746-8754.
- 48 A. P. Bassett, S. W. Magennis, P. B. Glover, D. J. Lewis, N. Spencer, S. Parsons, R. M. Williams, L. De Cola and Z. Pikramenou, Highly luminescent, triple- and quadruple-

- stranded, dinuclear Eu, Nd, and Sm(III) lanthanide complexes based on bis-diketonate ligands, J. Am. Chem. Soc., 2004, 126, 9413-9424.
- 49 A. J. Brock, I. M. Etchells, E. G. Moore and J. K. Clegg, stranded phenyl-spaced Dinuclear triple 1,3-bisβ-diketonato lanthanide(III) complexes: synthesis, structures and spectroscopy, Dalton Trans., 2021, 50, 4874-4879.
- 50 R. Chen, Q. Q. Yan, S. J. Hu, X. Q. Guo, L. X. Cai, D. N. Yan, L. P. Zhou and Q. F. Sun, Dinuclear helicate or mononuclear pincer lanthanide complexes from one ligand: stereo-controlled assembly and catalysis, Org. Chem. Front., 2021, 8, 2576-2582.
- 51 X. Gao, H. Li, P. Chen, W. Sun and P. Yan, A series of triplestranded lanthanide(III) helicates: Syntheses, structures and single molecular magnets, Polyhedron, 2017, 126, 1-7.
- 52 P. Chen, H. Li, W. Sun, J. Tang, L. Zhang and P. Yan, Crystallization of triple- and quadruple-stranded dinuclear bis-β-diketonate-Dy(III) helicates: Single molecule magnetic behavior, CrystEngComm, 2015, 17, 7227-7232.
- 53 H. Li, P. Chen, W. Sun, L. Zhang and P. Yan, Solvent triggered structural diversity of triple-stranded helicates: Single molecular magnets, Dalton Trans., 2016, 45, 3175-3181.
- 54 H. Ju, J. K. Clegg, K.-M. Park, L. F. Lindoy and S. S. Lee, Formation of a dicopper platform based polyrotaxane whose "string" and "bead" are constructed from the same components, J. Am. Chem. Soc., 2015, 137, 9535-9538.
- 55 G. Truccolo, Z. Tessari, J. Tessarolo, S. Quici, L. Armelao and M. Rancan, A Cu(II) metallocycle for the reversible selfassembly of coordination-driven polyrotaxane-like architectures, Dalton Trans., 2018, 47, 12079-12084.

- 56 M. Rancan, G. Truccolo, A. Carlotto, S. Quici and L. Armelao, A Zn(II) metallocycle as platform to assemble a $1D+1D \rightarrow 1D$ polyrotaxane via $\pi \cdots \pi$ stacking of an ancillary ligand, Inorganics, 2019, 7, 137.
- 57 F. Li, J. K. Clegg, L. F. Lindoy, R. B. MacQuart and G. V. Meehan, Metallosupramolecular self-assembly of a universal 3-ravel, Nat. Commun., 2011, 2, 1-5.
- 58 CRC Handbook of Chemistry and Physics, ed. W. M. Havnes, D. R. Lide and T. J. Bruno, CRC Press, 97th edn, 2016.
- 59 G. Ercolani, C. Piguet, M. Borkovec and J. Hamacek, Symmetry numbers and statistical factors in self-assembly and multivalency, J. Phys. Chem. B, 2007, 111, 12195-12203.
- M. L. Bhaumik, Exchange of rare-earth ions in chelate solu-60 tions, J. Inorg. Nucl. Chem., 1965, 27, 243-244.
- 61 N. Serpone and R. Ishayek, Ligand-exchange equilibrium studies of some octacoordinate yttrium(III)-β-diketonate complexes by hydrogen-1 nuclear magnetic resonance spectroscopy, Inorg. Chem., 1971, 10, 2650-2656.
- 62 F. A. Cotton, P. Legzdins and S. J. Lippard, Nuclear magnetic resonance studies of eight-coordinate β-diketonate complexes related to laser chelates, J. Chem. Phys., 1966, 45, 3461-3462.
- 63 K. Binnemans, Rare-earth beta-diketonates, Handb. Phys. Chem. Rare Earths, 2005, 35, 107-272.
- 64 H. Samelson, C. Brecher and A. Lempicki, Spectroscopy and coordination chemistry of europium chelate solutions: Concentration and solvent effects, J. Mol. Spectrosc., 1966, 19, 349-371.
- 65 D. P. Fay, D. Litchinsky and N. Purdie, Ultrasonic absorption in aqueous salts of the lanthanides, J. Phys. Chem., 1969, 73, 544-552.