

Transient self-assembly of metal-organic complexes

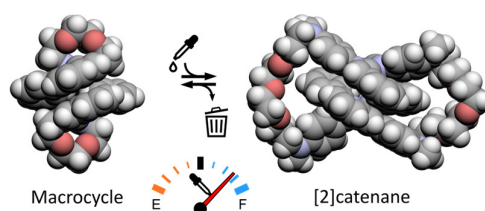
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Abstract



Implementing transient processes in networks of dynamic molecules holds great promise for developing new functional behaviours. Here we report that trichloroacetic acid can be used to temporarily rearrange networks of dynamic imine-based metal complexes towards new equilibrium states, forcing them to express complexes otherwise unfavourable in their initial equilibrium states. Basic design principles were determined for the creation of such networks. Where a complex distribution of products was obtained in the initial equilibrium state of the system, the transient rearrangement temporarily yielded a simplified output, forcing a more structured distribution of products. Where a single complex was obtained in the initial equilibrium state of the system, the transient rearrangement temporarily modified the properties of this complex. By doing so, the mechanical properties of an helical macrocyclic complex could be temporarily altered by rearranging it into a [2]catenane.

Introduction

To rival the capabilities of biological systems, temporal control over chemical reactivity must be achieved in synthetic systems. Most synthetic self-assembly processes are designed to generate well-ordered structures with high thermodynamic or kinetic stabilities—these structures being at the global minimum of the energy landscape or trapped in local minima.¹ By modifying the energy landscape using external stimuli (such as pH, light, or the addition of a chemical species) to create a new minimum, these structures could be forced to rearrange into new ones, yielding stimuli-responsive self-assembly processes.² While this approach produces highly functional systems,³ it requires the repeated intervention of an operator to administer opposing stimuli at appropriate times to switch the system back and forth between its different functional states.

To overcome this limitation and inspired by biological systems,^{1b,4} chemists have coupled self-assembly and energy-dissipating processes so that self-assembly processes could transiently express different structures via an influx of energy in the form of light, heat or chemicals.^{1b,5} These so-called “transient self-assemblies” require a constant input of energy to persist in time. If the energy supply is stopped, these structures dismantle, their components being recaptured by the initial state of the system (*i.e.*, its state prior to the energy influx). Interest in chemically-induced transient self-assembly is rapidly growing encouraged by examples of their unique capabilities, including transient gel formation,⁶ operation of molecular machines,⁷ temporal control over host–guest systems,⁸ transient formation of supramolecular assemblies⁹ and non-equilibrium self-replication.¹⁰ In most current reports, however, chemically-induced transient processes are mostly used to modulate the behaviour of single component self-assemblies. Few examples exist in which the consumption of a chemical reactant regulates the organization of networks of dynamic molecules.^{10c,11} Using classical stimuli-responsive self-assembly, networks of dynamic molecules have enabled the emergence of complex functions inaccessible with simpler systems.^{3a,3d,3g,12,13} Combining the capabilities of such networks with energy-dissipating processes would enable the development of new adaptive behaviours, furthering our understanding of complex systems (including biological ones).

To date, libraries of dynamic metal–organic complexes consisting of amine and 2-formylpyridine components reversibly condensed into imine ligands around labile metal-ions have yielded some of the most advanced artificial networks of dynamic molecules.^{3a,3d,3g,12g,12k,13c–i}

Here we demonstrate that a ‘transient acid’^{5h,7b–g,7i,8c,11f,12j} can force libraries of imine-based metal complexes to temporarily rearrange towards a new equilibrium state, generating complexes otherwise unfavourable in absence of the acid. The study provides basic design principles for creating networks of dynamic complexes with temporarily modifiable functions, showing that the properties of metal-organic complexes can be altered by their transient rearrangement.

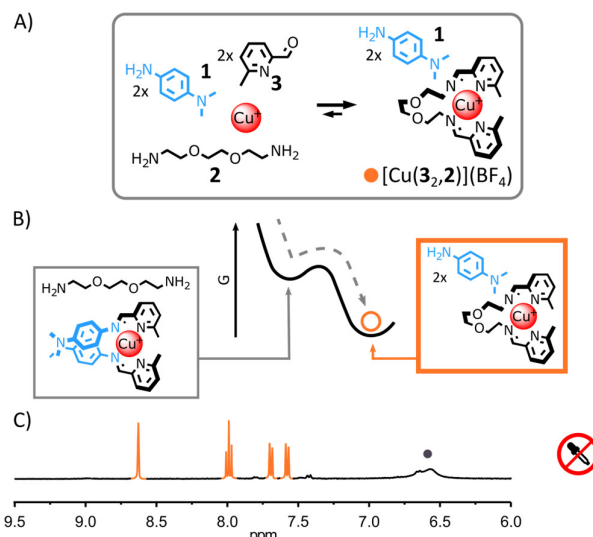


Figure 1. (A) Self-assembly of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ under thermodynamic control. Reaction conditions: $\mathbf{1}:\mathbf{2}:\mathbf{3}:\text{Cu}(\text{BF}_4)$ (2:2:2:1), CD_3CN , 60 °C, 12 h. (B) Schematic representation of the Gibbs free energy landscape of the reaction; $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ only forms as a metastable intermediate during the self-assembly process. (C) Partial ^1H NMR spectrum (400 MHz, CD_3CN , 297 K) of the crude reaction mixture after 12 h at 60 °C; the diagnostic signals of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ are coloured in orange and two of the diagnostic signals of $\mathbf{1}$ are highlighted by a grey circle.

Results and Discussion

Rational and main design

Due to the differences of basicity between arylamine and alkylamine, we envisaged that trichloroacetic acid—an acid capable of temporarily acidifying a medium^{5h,7c,7d,7i,11f,12j}—could be used to induce the transient rearrangement of imine-based metal complexes.

Here we will use the notation **(a,b)** to refer to the imine-based ligand obtained by the condensation of aldehyde **a** with amine **b**. The more basic ditopic amine **2** (Fig. 1A and 1B) should yield the most thermodynamically stable complex with Cu(I) and 2-formylpyridine **3** under neutral conditions, based on previous work.^{11f,13f,14} Complex $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ should therefore be the unique complex observed when Cu(I) is added to an equimolar mixture of **1**, **2** and **3**. Upon addition of trichloroacetic acid, the initial acidification of the medium should displace **2** from the complex by protonation, allowing the incorporation of the less basic arylamine **1** (Fig. 2A and 2B). A new transient equilibrium state should quickly be achieved, expressing $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ as its most thermodynamically stable complex. The slow decarboxylation of trichloroacetate^{5h,7c,7d,7i,11f,12j} should then incrementally return the system of the metal complexes to its initial equilibrium state (*i.e.*, prior to the addition of acid) via a quasi-static process continually maintaining the metal complexes at equilibrium (the rearrangement of the metal complexes being faster than the decarboxylation of trichloroacetate). The decarboxylation of trichloroacetate generates a strong base—trichloromethyl anions—that should restore neutral alkylamine **2**, regenerating $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ with chloroform and carbon dioxide as the only side products.

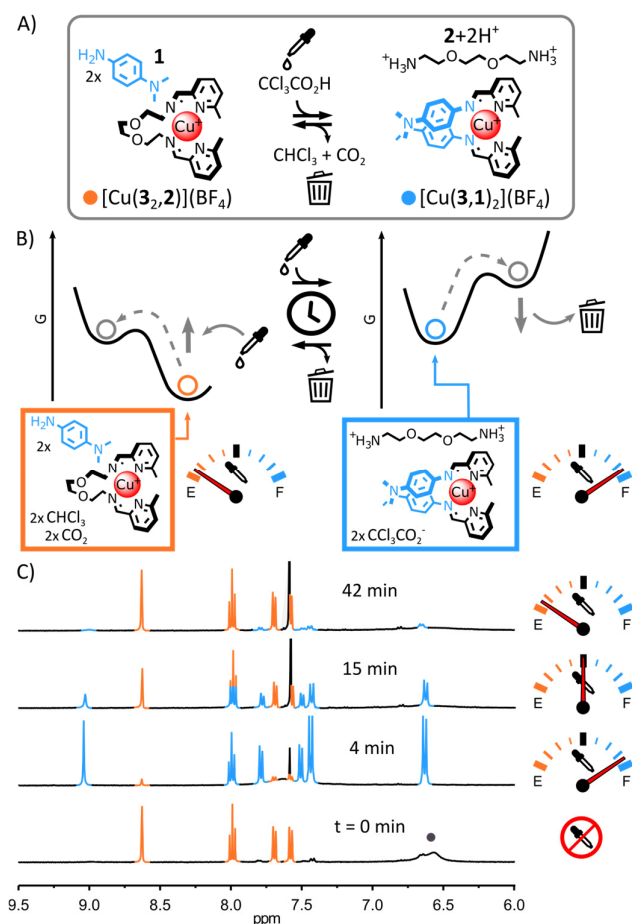


Figure 2. (A) Transient rearrangement of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ into $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$. Conditions: CCl_3COOH (2.5 eq.), CD_3CN , r. t. (B) Schematic representation of the Gibbs free energy landscape of the rearrangement; $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ forms under thermodynamic control as part of a quasi-static process and $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ forms under thermodynamic control in the default equilibrium state of the system. (C) Partial ^1H NMR spectra (400 MHz, CD_3CN , 297 K) showing the evolution of the crude reaction mixture upon addition of CCl_3COOH ; the diagnostic signals of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ and $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ are coloured in orange and blue, respectively, two of the diagnostic signals of $\mathbf{1}$ are highlighted by a grey circle.

This rationale was tested. Combining alkylamine $\mathbf{2}$, 2-formylpyridine $\mathbf{3}$ and Cu(I) in a 2:2:1 ratio in CD_3CN led to the quantitative formation of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ after 12 h at 60°C (see ESI for details and full characterization data of new compounds.) When arylamine $\mathbf{1}$ (2 eq.) was added to the solution at room temperature, $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ remained the only complex observable by ^1H NMR spectroscopy (Fig. 1C), hinting at its greater stability compared to $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$. Addition of trichloroacetic acid (2.5 eq.) to this stable mixture of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ and $\mathbf{1}$ immediately caused the rearrangement of the system, yielding $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ and $[\mathbf{2}+2\text{H}^+]$ as unique products observable by ^1H NMR spectroscopy (Fig. 2C). After a few minutes, the signals of $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ and $[\mathbf{2}+2\text{H}^+]$ started to diminish giving place to those of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ and $\mathbf{1}$ —the decarboxylation of trichloroacetate slowly returning the system to its initial state (ESI, Fig. S18). After ca. 40 min, the ^1H NMR spectra of the reaction mixture closely matched that prior to the addition of acid, differing only by the emergence of a strong signal at 7.58 ppm attributed to the side product CHCl_3 (Fig. 2C).

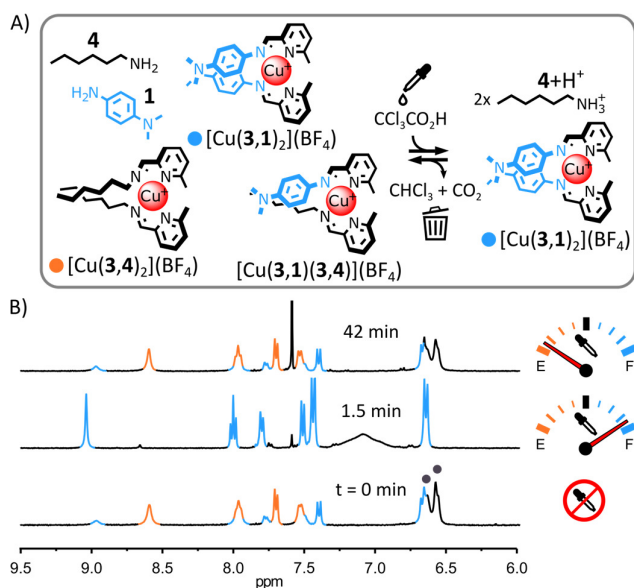


Figure 3. (A) Transient ‘simplification’ of a mixture of complexes into $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$. Conditions: CCl_3COOH (2.5 eq.), CD_3CN , r. t. (B) Partial ^1H NMR spectra (400 MHz, CD_3CN , 297 K) showing the evolution of the crude reaction mixture upon addition of CCl_3COOH ; the diagnostic signals of $[\text{Cu}(\mathbf{3},\mathbf{4})_2]^+$ and $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ are coloured in orange and blue, respectively, two of the diagnostic signals of $\mathbf{1}$ are highlighted by a grey circle, the diagnostic signals of $[\text{Cu}(\mathbf{3},\mathbf{1})(\mathbf{3},\mathbf{4})]^+$ overlap with those of $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ and $[\text{Cu}(\mathbf{3},\mathbf{4})_2]^+$.

The robustness of the system was tested by subjecting the same sample to five consecutive rearrangement cycles (ESI, Fig. S18 – 23). The evolution of the system was monitored by ^1H NMR spectroscopy, invariably showing behaviours identical to those observed during the first rearrangement cycle (ESI, Fig. S18 – 23) and showing only minimal deterioration of the system even after five consecutive cycles (ESI, Fig. S24). Having established that trichloroacetic acid could induce the transient rearrangement of a library of metal complexes—temporarily forming a complex unfavourable in the initial equilibrium state of the system—we sought to probe the tolerance of the system to modifications.

Structural tolerance of the design

The structural tolerance of the system shown in Figure 2 was studied by systematic modification of the nature of (i) the alkylamine, (ii) the tetrahedral metal ion and (iii) the arylamine.

(i) The substitution of ditopic alkylamine $\mathbf{1}$ for monotopic alkylamine $\mathbf{4}$ resulted in a distribution of homoleptic and heteroleptic complexes in the initial equilibrium state of the system (Fig. 3); however, the acid-induced rearrangement of the system remained unchanged. Upon addition of arylamine $\mathbf{1}$ (2 eq.) to $[\text{Cu}(\mathbf{3},\mathbf{4})_2]^+$ in CD_3CN , ^1H NMR signals corresponding to $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ (and to the heteroleptic complex $[\text{Cu}(\mathbf{3},\mathbf{1})(\mathbf{3},\mathbf{4})]^+$) appeared besides those of $[\text{Cu}(\mathbf{3},\mathbf{4})_2]^+$ (Fig. 3B). The lack of the self-sorting in the initial equilibrium state of the system could be attributed to the reduced entropic driving force for the formation of $[\text{Cu}(\mathbf{3},\mathbf{4})_2]^+$ compared to the formation of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ (loss of the chelate effect).^{14b,15} Adding trichloroacetic acid (2.5 eq.) to the system immediately

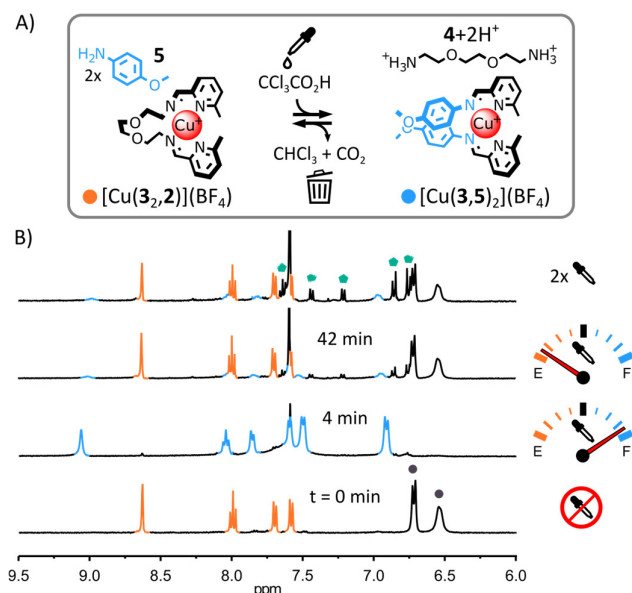


Figure 4. (A) Transient rearrangement of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ into $[\text{Cu}(\mathbf{3},\mathbf{5})_2]^+$. Conditions: $\text{CCl}_3\text{CO}_2\text{H}$ (2.5 eq.), CD_3CN , r. t. (B) Partial ^1H NMR spectra (400 MHz, CD_3CN , 297 K) showing the evolution of the crude reaction mixture upon addition of $\text{CCl}_3\text{CO}_2\text{H}$ and after two rearrangement cycles; the diagnostic signals of $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ and $[\text{Cu}(\mathbf{3},\mathbf{5})_2]^+$ are coloured in orange and blue, respectively, two of the diagnostic signals of **5** are highlighted by a grey circles, the diagnostic signals of trichloromethylated imine (**3,5**) are highlighted by green pentagons.

simplified the reaction mixture, with only $[\text{Cu}(\mathbf{3},\mathbf{1})_2]^+$ and $[\mathbf{2}+2\text{H}^+]$ being observable by ^1H NMR spectroscopy. This ‘simplified state’ persisted for a few minutes before slowly returning to the initial distribution of products (after ca. 40 min, ESI Fig. S26). This transient ‘simplified state’ could be recalled three times with little signs of fatigue (ESI, Fig. S26 – 30). This system exemplifies how a network of dynamic complexes can be forced to temporarily express fewer products—a simplified output—by populating a different more structured equilibrium state via an energy-dissipating process.

(ii) The system behaves similarly when Cu(I) is replaced by Ag(I). The increased plasticity of the coordination geometry of Ag(I)¹⁶ yielded a distribution of complexes in the initial equilibrium state of the system, that immediately simplified upon addition of trichloroacetic acid. Following the addition of acid (2.5 eq.), $[\text{Ag}(\mathbf{3},\mathbf{1})_2]^+$ and $[\mathbf{2}+2\text{H}^+]$ were the sole products observed in solution by ^1H NMR spectroscopy (ESI, Fig. S31–32). The system gradually returned to its initial distribution of products over ca. 40 min (ESI, Fig. S32). This process could be repeated for five consecutive cycles with minimal deterioration (ESI, Fig. S32–38), indicating the compatibility of Ag(I) with our transient rearrangement process.

(iii) The use of the less electron-rich *p*-anisidine **5** in place of 4-(dimethylamino)aniline **1** was uninfluential on the initial self-assembly of the system or its transient rearrangement (Fig. 4). From ^1H NMR spectra, $[\text{Cu}(\mathbf{3},\mathbf{2})]^+$ and **5** were the only species present initially and $[\text{Cu}(\mathbf{3},\mathbf{5})_2]^+$ and $[\mathbf{2}+2\text{H}^+]$ were the only species present upon addition of trichloroacetic acid (Fig. 4B). However, in subsequent rearrangement cycles (Fig. 4B, top spectrum), **5** led to the rapid deterioration of the system via the trichloromethylation of imine (**3,5**) (ESI, Fig. S40–S42).¹⁷ This result indicates that in our original system (Fig. 2) the strong

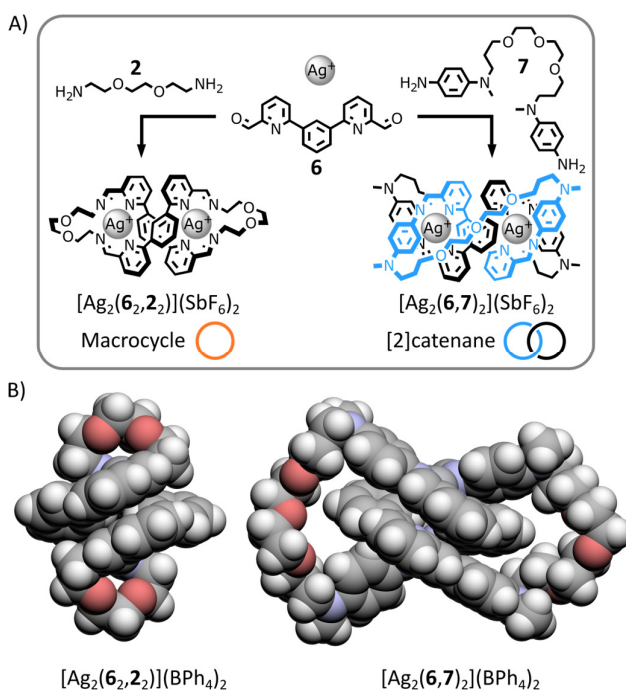


Figure 5. (A) Self-assembly of macrocycle $[\text{Ag}_2(\mathbf{6},\mathbf{2})_2]^{2+}$ and [2]catenane $[\text{Ag}_2(\mathbf{6},\mathbf{7})_2]^{2+}$ under thermodynamic control. Reaction conditions: **6**:**2**: $\text{Ag}(\text{SbF}_6)$ (1:1:1) or **6**:**7**: $\text{Ag}(\text{SbF}_6)$ (1:1:1), CD_3CN : CDCl_3 3:1, 60 °C, 12 h. (B) Single crystal X-ray structures of macrocycle $[\text{Ag}_2(\mathbf{6},\mathbf{2})_2](\text{BPh}_4)_2$ and of [2]catenane $[\text{Ag}_2(\mathbf{6},\mathbf{7})_2](\text{BPh}_4)_2$; solvent molecules and counterions have been omitted for clarity.

electron-donating *N,N*-dimethyl group of **1** must reduce the reactivity of imine (**3,1**) towards nucleophilic attack by the trichloromethyl anions generated by the decarboxylation of trichloroacetate.

Having established basic design principles, we envisaged that the transient rearrangement of metal-organic complexes described in Figure 2 could be used to temporarily modify the properties of supramolecular architectures, enabling a macrocyclic complex to become a [2]catenane—two mechanically interlocked macrocycles.

Transient rearrangement of a macrocyclic complex into a [2]catenane

Dialdehyde **6** (Fig. 5A) is known to self-assemble with *p*-anisidine **5** and tetrahedral metal ions to produce dinuclear double-helical complexes (e.g., $[\text{Cu}_2(\mathbf{6},\mathbf{5})_2]^{2+}$).^{13b} Based on previous work by Nitschke,^{14b} we anticipated that both a macrocyclic complex and a [2]catenane could be generated from this helical motif if appropriate diamines were used in place of **5** (Fig. 5A).

In the presence of Ag(I) and aided by the gauche effect,¹⁸ dialkylamine **2** should have the right length and flexibility, to preferentially bridge two distinctive dialdehydes **6** while being incorporated in the helical complex, generating the dinuclear helical macrocycle $[\text{Ag}_2(\mathbf{6},\mathbf{2})_2]^{2+}$ (Fig. 5A). In contrast, the longer length and increased rigidity of diarylamine **7** should favour its condensation with the same dialdehyde **6** in the helical complex, yielding the dinuclear helical [2]catenane $[\text{Ag}_2(\mathbf{6},\mathbf{7})_2]^{2+}$ (Fig. 5A).

To confirm their architectures, both complexes were prepared by heating an equimolar mixture of dialdehyde **6**,

diamine **2** or **7** and $\text{Ag}(\text{SbF}_6)$ in $\text{CD}_3\text{CN}:\text{CDCl}_3$ 3:1 at 60 °C for 12 h. In both cases, a single product was obtained with NMR spectra and ESI-MS analyses consistent with the structures shown in Figure 5 (ESI Section 2.4.2 and 2.4.3). X-ray-quality crystals of both architectures were grown by liquid–liquid diffusion (ESI, Section 4.1 and 4.2). The solid-state structures of both complexes unambiguously confirmed their topologies (Fig. 5B and ESI, Section 4.1 and 4.2).

The organic ligand of macrocycle $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ forms a 42-atom-loop twisted in a double-helical shape by two $\text{Ag}(\text{I})$ ions each coordinated to two iminopyridine domains of the ligand in a flattened tetrahedral geometry. The architecture of [2]catenane $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ consists of two interlocked 36-membered rings entwined by an helical core created by two $\text{Ag}(\text{I})$ ions coordinating each to one of the iminopyridine domain of each macrocycles in an irregular four-coordinate geometry.

These two structures exemplify the importance of controlling crossing point connectivity while synthesizing interlocked architectures.¹⁹ Both complexes have the same helical core, providing the same initial three crossing points. The difference of chemical topology between macrocycle $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ and [2]catenane $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ arises from the retention of a different number of these initial crossing points when connecting the end-groups of the helical core with diamine **2** or **7**. By connecting the closest end-groups of the helical core, diamine **2** retains none of the initial crossing points, yielding a topologically trivial macrocycle. By connecting the end-groups of the same strand of the helical core, diamine **7** retains one of the initial crossing points, yielding a [2]catenane.

Having established the nature of both complexes, we attempted the transient rearrangement of $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ and **7** into $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ and $[\mathbf{2}+2\text{H}^+]$.

As shown in Figure 6, when diarylamine **7** (1 eq.) was added to $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ in $\text{CD}_3\text{CN}:\text{CDCl}_3$ 3:1, ^1H NMR spectroscopy provided no evidence of reaction between these two species, indicating that [2]catenane $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ is unfavourable in the initial equilibrium state of the system. Addition of trichloroacetic acid (10 eq.) triggered the transient rearrangement of the system: macrocycle $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ was gradually converted into [2]catenane $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ and dialkylamine **2** was trapped in its protonated form (Fig. 6B). The concentration of $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ peaked at ca. 240 min before starting to decrease—slowly at first, but accelerating with depletion of the acid (ESI, Fig. S46 and S48). After ca. 870 min, macrocycle $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ was again the only complex visible by ^1H NMR spectroscopy.

Compared to the systems of mononuclear complexes in Figures 2–4, the rate of rearrangement of this system was considerably lower ($[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ peaked after ca. 240 min whereas $[\text{Ag}(\mathbf{3}, \mathbf{1})_2]^+$ peaked after ca. 2 min), reflecting the increased complexity of this rearrangement process. Metal-organic complexes involving dialdehyde and diamine components are known to self-assemble via the initial formation of metastable ill-defined oligomers and polymers slowly rearranging into (kinetically or thermodynamically) more stable structures.^{18b,20} A similar scenario is likely when the present system is rearranging towards [2]catenane $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$

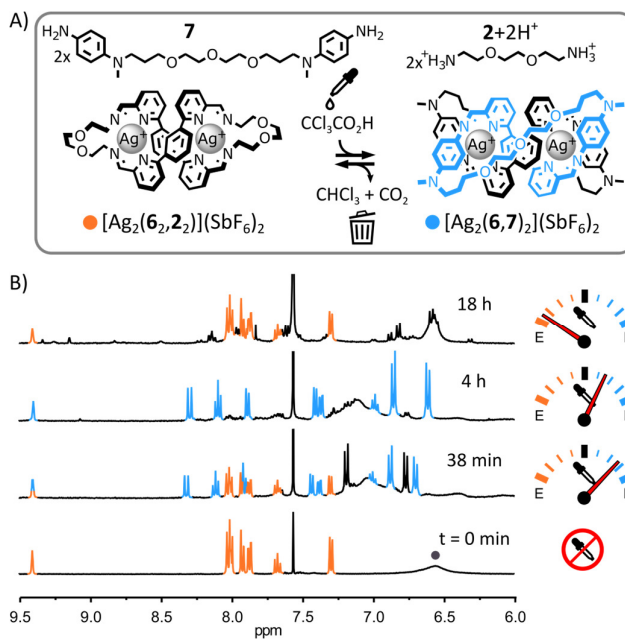


Figure 6. (A) Transient rearrangement of macrocycle $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ into [2]catenane $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$. Conditions: CCl_3COOH (10 eq.), $\text{CD}_3\text{CN}:\text{CDCl}_3$ 3:1, r. t. (B) Partial ^1H NMR spectra (400 MHz, $\text{CD}_3\text{CN}:\text{CDCl}_3$ 3:1, 297 K) showing the evolution of the crude reaction mixture upon addition of CCl_3COOH ; the diagnostic signals of $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ and $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ are coloured in orange and blue, respectively, two of the diagnostic signals of **7** are highlighted by a grey circle.

following the addition of trichloroacetic acid. Indications of the formation of ill-defined oligomeric and polymeric species were found by monitoring the rearrangement of the system by ^1H NMR spectroscopy. Despite $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ and $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ being the only species visible by ^1H NMR spectroscopy throughout the rearrangement process (Fig. 6B and ESI, Fig. 46), their combined populations remained under ca. 80% of the initial population of $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ (i.e., prior to the addition of trichloroacetic acid), the missing material being presumably trapped in ill-defined oligomers and polymers undetectable by ^1H NMR spectroscopy. After the concentration of $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ peaked, the baseline of the ^1H NMR spectra of the reaction became broad and uneven in the aromatic region (Fig. 6B and ESI, Fig. 46), indicating an increased formation of oligomeric and polymeric species. This observation is consistent with the progressive deprotonation of $[\mathbf{2}+2\text{H}^+]$ during the decomposition process of trichloroacetate into CHCl_3 and CO_2 , making both diamines **2** and **7** available to form oligomeric and polymeric species as the system returns to its initial equilibrium state.

Compared to the systems of mononuclear complexes in Figures 2–4, a large excess of trichloroacetic acid was required to maximize the formation of [2]catenane $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ (10 eq. of trichloroacetic acid were needed in this case vs. 2.5 eq. in the case of the simpler systems). The amount of $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ generated during the transient process and its lifetime were both found to be roughly proportional to the amount of acid added (ESI, Fig. S44–S46 and S48). When the amount of acid added was increased from 7.5 to 10 eq., the peak population of $[\text{Ag}_2(\mathbf{6}, \mathbf{7})_2]^{2+}$ increased from ca. 35 to 70% of the initial population of $[\text{Ag}_2(\mathbf{6}_2, \mathbf{2}_2)]^{2+}$ and the amount of time required for

the system to return to its initial equilibrium state increased from ca. 450 to 810 min. This high requirement in acid suggests that the rearrangement of the system is slow compared to the rate of decarboxylation of trichloroacetate. An excess of acid is therefore needed to reach a steady state in the system, maximizing the transient formation of catenane $[\text{Ag}_2(\mathbf{6},\mathbf{7})_2]^{2+}$ under thermodynamic control.

Higher amounts of acid (up to 12 eq., ESI Fig. S47 and S48) did not increase the peak population of $[\text{Ag}_2(\mathbf{6},\mathbf{7})_2]^{2+}$ further, hinting that some of the oligomeric and polymeric intermediates formed during the rearrangement of the system may be too kinetically stable to rearrange in the time scale of the transient process (in the conditions used).

While all the initial macrocycle $[\text{Ag}_2(\mathbf{6}_2,\mathbf{2}_2)]^{2+}$ could be recovered at the end of the rearrangement using 7.5 eq. of trichloroacetic acid (ESI Fig. S44 and S48), the amount of macrocycle recovered at the end of the rearrangement cycle diminished when more acid was used (90 and 70% of the initial macrocycle were recovered when 8.5 or 12 eq. of trichloroacetic acid were used, respectively, ESI, Fig. S45 – S48). Performing a second rearrangement cycle led to significant deterioration of the system (ESI, Fig. S50).

Conclusions

Trichloroacetic acid has been used to control in time the composition of imine-based metal complexes via energy-dissipation. By exploiting the base-promoted decarboxylation of trichloroacetic acid, it was possible to cycle between the preferential incorporation of an arylamine or an alkylamine component in a metal complex, enabling the transient expression of complexes otherwise unfavourable in absence of trichloroacetic acid.

The electronic properties of the arylamine component appeared to be crucial to the robustness of the transient rearrangement process, whereas the nature of the alkylamine component and tetrahedral metal ions had little influence. Transient processes can temporarily simplify the output of a self-assembly process by populating a different, more structured, equilibrium state. The properties of metal complexes can be controlled in time via the transient modification of their compositions. We showed that the mechanical properties of a macrocyclic complex could be temporarily altered by rearranging it into a topologically more complex [2]catenane.

By providing a generalizable method to transiently access different equilibrium states in systems of metal-organic complexes (or architectures), the information gleaned from our investigation will facilitate the development of new adaptive behaviours in these systems; behaviours exploiting both the capabilities of the default equilibrium (or kinetically trapped) state of the system and the capabilities of transient equilibrium states accessible via energy-dissipation.

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