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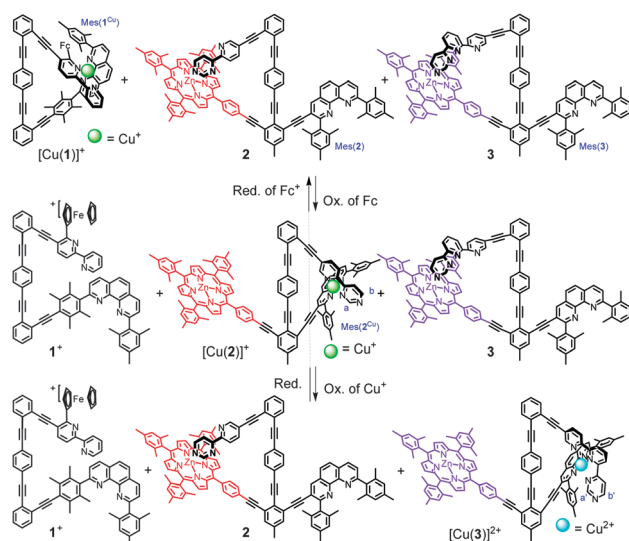
A trio of nanoswitches in redox-potential controlled communication†

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A potential-controlled two-step bidirectional communication protocol between the nanoswitches [Cu(1)]⁺, 2 and 3 is set up, in which ligand followed by metal-ion oxidation drives two subsequent metal ion translocations (self-sorting) changing the switching state at each switch. The communication is reset to its starting point by a two-electron reduction.

Life processes are sustained through intricate communication and signaling networks involving recognition, processing and amplification of information, sometimes through cascading pathways.¹ While multistep signaling, such as in the metallothionein protein,² is often encountered in biology, artificial switches³ or machines⁴ are rarely networked by chemical communication,⁵ and if, they are engaged in single-step protocols.⁶ Intermolecular communication involving major rearrangement at two switches has been reported by the group of Aprahamian^{6c} and more recently by us.^{6d} In the former case, being reminiscent of a biological proton relay, binding of Zn²⁺ ions induced a toggling at the first switch triggering proton release, which stimulated a change in the switching state at the second switch. Our recent work^{6d} demonstrated bidirectional reversible communication between the nanoswitches 2 and 3 using redox equivalents as input signals (Scheme 1). The strategy was based on two orthogonal⁷ coordination scenarios developed by us over the years. Switch 2 equipped with its intramolecular bipyridine and phenanthroline (HETPHEN⁸) coordination site prefers Cu⁺ ions, whereas 3 with its terpyridine and phenanthroline (HETTAP⁸) binding motif favours Cu²⁺ or Zn²⁺.⁹ Due to these preferences, the nanoswitch duo [Cu(2)]⁺ and 3 experienced a copper translocation generating 2 and [Cu(3)]²⁺ upon oxidation, while the starting point was reset by reduction.^{6d}

Here, we describe a two-step bidirectional communication protocol between the three nanoswitches 1–3 using two subsequent



Scheme 1 Reversible cascade switching among three nanoswitches.

one-electron redox inputs. Despite integrating an already reported redox-controlled communication scenario, *i.e.* between switches 2 and 3,^{6d} the realisation of two-step communication between the trio 1–3 adds a new level of complexity requiring to devise a perfectly matching nanoswitch 1 and a potential control for the two-step translocation. The key criteria for our design were: (1) the two oxidation steps have to occur at separate potentials each followed by a clean self-sorting¹⁰ propelling the communication cascade. (2) The new electron-rich switch 1 should have a higher affinity toward Cu⁺ than the known HETPHEN- and HETTAP-based switches 2 and 3 and (3) the lowest oxidation potential. (4) After oxidation, nanoswitch 1 should be the weakest binding ligand for Cu⁺ and (5) also should rapidly release Cu⁺.

Taking all the requirements into account, we considered 2-ferrocenyl-[1,10]-phenanthroline and 6-ferrocenyl-2,2'-bipyridine as possible binding motifs in 1, expecting that the electron-donating ferrocenyl group¹¹ next to the chelate site would not only provide a sizeable gain in binding strength but also ensure

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release of the metal ion upon oxidation of the ferrocene. In an initial screening, 2,9-dimesityl-[1,10]-phenanthroline provided a stronger binding heteroleptic copper(i) complex in combination with 2-ferrocenylphenanthroline, but after oxidation copper remained strongly bound. As a result, we turned to 6-ferrocenyl-2,2'-bipyridine despite its lower binding quality for Cu^+ .

Based on these considerations, we devised a new nanoswitch **1** following our successful triangular design¹² (Scheme 1), expecting the kinetics of metal-ion release to be fast. Its main competitor for Cu^+ , *i.e.* nanoswitch **2**, should be at disadvantage in the competition, because copper(i) complexation at **2** goes along with loss of the pyrimidine \rightarrow zinc(ii) porphyrin interaction.

While the preparation of nanoswitches **2**^{12b} and **3**^{6d} has been reported previously, nanoswitch **1** was synthesised in 60% yield from its direct precursor *via* a Sonogashira coupling (for details, see the ESI†). Upon reacting switch **1** with Cu^+ , the ferrocene protons at the substituted cyclopentadienyl ring lost their symmetry and shifted upfield to 3.32, 3.67, 4.54 and 4.92 ppm in the ¹H NMR spectrum due to the proximity of the shielding phenanthroline. Additionally, the mesityl protons shifted upfield to 6.35 and 6.43 ppm (Fig. S5, ESI†). The ESI-MS spectrum displays a single peak at m/z = 1155.5 Da with its isotopic distribution exactly matching the theoretical one of complex $[\text{Cu}(\mathbf{1})]^+$ (Fig. S7, ESI†).

The cyclic voltammograms show fully reversible redox $\text{Fc}^{0/+}$ waves for **1**^{0/+} at $E_{1/2}$ = 450 mV_{SCE} (Fig. S3, ESI†) and for $[\text{Cu}(\mathbf{1})]^{+/2+}$ at $E_{1/2}$ = 610 mV_{SCE}, while $\text{Cu}^{+/2+}$ oxidation in $[\text{Cu}(\mathbf{1})]^{2+/3+}$ emerged at $E_{1/2}$ = 940 mV_{SCE} (Fig. S6, ESI†). On the basis of the data, we found that Cu^+ is less strongly bound in $[\text{Cu}(\mathbf{1})]^{2+}$ than in $[\text{Cu}(\mathbf{1})]^+$ by $\Delta\log K$ = 2.7.

As a model for paramagnetic $[\text{Cu}(\mathbf{1})]^{2+}$, we characterised diamagnetic $[\text{Zn}(\mathbf{1})]^{2+}$, prepared from stoichiometric amounts of **1** and $\text{Zn}(\text{OTf})_2$. The splitting patterns of the diagnostic protons are similar as for $[\text{Cu}(\mathbf{1})]^+$ but they appear at different positions in the ¹H NMR spectrum. For instance, protons from the substituted cyclopentadienyl ring of ferrocene appear at 3.35, 3.84, 4.37 and 5.48 ppm. Moreover, mesityl protons are shifted to 6.03 and 6.43 ppm (Fig. S8, ESI†). Formation of the complex was also proved from ESI-MS data with a peak at m/z = 578.6 Da and an isotopic distribution matching the one expected for $[\text{Zn}(\mathbf{1})]^{2+}$ (Fig. S9, ESI†).

To address the first step of the new communication protocol, switches **1** and **2** were treated with Cu^+ (1:1:1). After equilibration at 40 °C for a few minutes, 90% of copper(i) was found on $[\text{Cu}(\mathbf{1})]^+$ (Fig. 1a). The selectivity was ascertained from ¹H NMR and UV-Vis data. Reciprocally, addition of switch **1** to a solution of $[\text{Cu}(\mathbf{2})]^+$ (2.70 mM) led to the same results as documented by the appearance of new signals at 2.87 and 3.31 ppm being typical for the free nanoswitch **2** with its pyrimidine ring being immersed into the shielding current of the zinc porphyrin ring (Fig. S11, ESI†). At the same time, emergence of signals at 3.32, 3.67, 4.54 and 4.92 ppm along with upfield shifted mesityl protons (6.35 and 6.43 ppm) suggests formation of $[\text{Cu}(\mathbf{1})]^+$. For the communication step $[\text{Cu}(\mathbf{2})]^+ + \mathbf{1}$, the NMR spectrum shows that full equilibrium was reached 58 min after mixing at room temperature (Fig. S12, ESI†). At a lower concentration (c = 0.1 mM) metal translocation from $[\text{Cu}(\mathbf{2})]^+$ to **1**, followed by UV-Vis spectroscopy at room temperature, took over a day to reach the equilibrium (Fig. S13, ESI†).

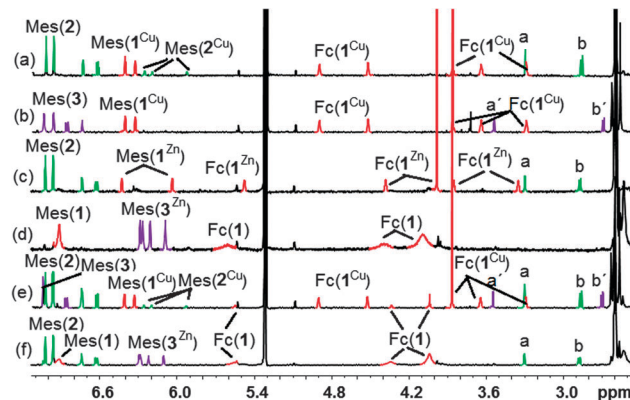


Fig. 1 Partial ¹H NMR spectra (400 MHz, CD_2Cl_2 , 298 K) of switches **1–3** in the presence of metal ions after equilibration. The traces represent: (a) Formation of a 90:10 ratio of $[\text{Cu}(\mathbf{1})]^+$ and $[\text{Cu}(\mathbf{2})]^+$ from a 1:1:1 mixture of **1**, **2** and Cu^+ . (b) Sole formation of $[\text{Cu}(\mathbf{1})]^+$ from a 1:1:1 mixture of **1**, **3** and Cu^+ . (c) Exclusive formation of $[\text{Zn}(\mathbf{1})]^{2+}$ from a 1:1:1 mixture of **1**, **2** and Zn^{2+} . (d) Complete formation of $[\text{Zn}(\mathbf{3})]^{2+}$ from a 1:1:1 mixture of **1**, **3** and Zn^{2+} . (e) Mixing of **1**, **2**, **3** and Cu^+ (1:1:1:1) generates 90% of $[\text{Cu}(\mathbf{1})]^+$ and 10% of $[\text{Cu}(\mathbf{2})]^+$ leaving **3** free in solution. (f) Exclusive formation of $[\text{Zn}(\mathbf{3})]^{2+}$ when **1**, **2**, **3** and Zn^{2+} (1:1:1:1) were mixed.

The exclusive selectivity for $[\text{Cu}(\mathbf{1})]^+$ over switch **3** is warranted due to the fact that **2** has a much higher binding constant for Cu^+ than switch **3** (Fig. S14, ESI†). For a kinetic analysis, we used UV-Vis spectroscopy (Fig. S15, ESI†) at c = 0.1 mM, because the metal-ion transfer is readily detectable due to the reorganisation of the azaterpyridine arm from the phenanthroline to the zinc porphyrin station. Addition of **1** to a solution of $[\text{Cu}(\mathbf{3})]^+$ at room temperature entailed a shift of the Q band absorption from 550 to 561 nm within 3 min in dichloromethane (DCM).

We chose Zn^{2+} as a diamagnetic input to check the face-to-face preferences of the switches for Cu^{2+} . Although in the competition between nanoswitches **1** and **2**, it was complex $[\text{Zn}(\mathbf{1})]^{2+}$ that formed exclusively with switch **2** remaining untouched (Fig. 1c and Fig. S16, ESI†), the pentacoordinated complex $[\text{Zn}(\mathbf{3})]^{2+}$ was afforded as a single product (Fig. 1d and Fig. S17, ESI†) from a mixture of **1**, **3** and Zn^{2+} (1:1:1). The latter result is quite surprising because formation of $[\text{Zn}(\mathbf{3})]^{2+}$ goes along with the loss of a azaterpyridine \rightarrow zinc(ii)porphyrin interaction. The outcome thus reflects the low stability of tetracoordinated zinc(ii) ions. Indeed, the typical ¹H NMR signals of the free switch **1** were preserved at 4.06, 4.32 and 5.57 ppm (from the ferrocene unit), whereas the diagnostic mesityl protons of switch **3** shifted upfield and splitted into four sets appearing at 6.07, 6.18, 6.24, and 6.27 ppm. The same set of signals is seen in the separately prepared $[\text{Zn}(\mathbf{3})]^{2+}$ complex.^{6d} Thus, Cu^{2+} , due to its similar coordination demands as Zn^{2+} , should bind selectively to nanoswitch **3** over **1** and even more over **1**⁺.

Because of the rather slow rate preventing a kinetic study using cyclic voltammetry, we initiated redox-triggered communication by adding $\text{TBPA}^+\text{SbCl}_6^-$ (tris(4-bromophenyl)aminium hexachloroantimonate) for oxidation and **dmfc** (decamethylferrocene) for reduction. To evaluate the subsequent step, a solution of $[\text{Cu}(\mathbf{1})]^+$ and **2** (at 1:1 ratio, 0.1 mM) in DCM was treated with one equivalent of $\text{TBPA}^+\text{SbCl}_6^-$. As a result, the Q band of **2** shifted from 561 to 550 nm and became constant after 4 min at room



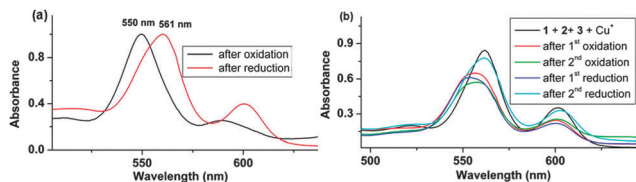


Fig. 2 (a) A solution of $[\text{Cu}(\text{CH}_3\text{CN})_4]\text{B}(\text{C}_6\text{F}_5)_4$, **1** and **2** = 1 : 1 : 1 at 10^{-4} M in dry DCM was treated with 1 equiv. of $\text{TBPA}^+\text{SbCl}_6^-$ (black trace), then with 1 equiv. of **dmfc** (red trace, also see the text). (b) UV-Vis changes at the Q band showing the communication between **1–3** and Cu^+ (1 : 1 : 1 : 1; at 2×10^{-5} M) as a result of two oxidation and two reduction steps (see text).

temperature (Fig. 2a, black trace). The change is in full agreement with a Cu^+ -induced swing of the nanoarm in **2** from the zinc(II) porphyrin to the phenanthroline station and thus formation of $[\text{Cu}(\text{2})]^+$ and **1**⁺. This interpretation was further corroborated by ESI-MS peaks at $m/z = 1093.4$ and 1799.2 Da, indicating the presence of switches **1**⁺ and $[\text{Cu}(\text{2})]^+$, respectively (Fig. S20, ESI†). For a reversal of the process, the above solution was reacted with **dmfc** (one equiv.). The absorption peak shifted back to 561 nm and became constant after 18 h (Fig. 2a, red trace). The ESI-MS spectrum of the solution confirmed the reverse translocation of Cu^+ as it displays mainly the peak at $m/z = 1155.5$ Da for $[\text{Cu}(\text{1})]^+$ and a small one at 1799.2 Da for $[\text{Cu}(\text{2})]^+$, reflecting the starting situation (Fig. S21, ESI†). To monitor the copper ion transfer by NMR, we chose a soluble **BFD**¹³ derivative for reduction of $[\text{Cu}(\text{2})]^+$ and **1**⁺ as it produces a diamagnetic oxidation product. The ¹H NMR spectrum shows the presence of complexes $[\text{Cu}(\text{1})]^+$ and $[\text{Cu}(\text{2})]^+$ in a ratio of 90 : 10 (Fig. S18, ESI†).

Reversible communication between $[\text{Cu}(\text{1})]^+$ and **3** upon oxidation is more complicated because on the product side the redox potentials, *i.e.* $E_{1/2}(\text{1}^{0/+}) = 450 \text{ mV}_{\text{SCE}}$ and $E_{1/2}([\text{Cu}(\text{3})]^{+/2+}) = 250 \text{ mV}_{\text{SCE}}$, invert.^{6d} As a result, a follow-up redox disproportionation of **1**⁺ and $[\text{Cu}(\text{3})]^+$ furnished **1** and $[\text{Cu}(\text{3})]^{2+}$. In the experiment, a 1 : 1 solution of $[\text{Cu}(\text{1})]^+$ and **3** (0.1 mM) in DCM was reacted with one equiv. of $\text{TBPA}^+\text{SbCl}_6^-$ and the metal exchange process was monitored at 561 nm (Q band of the free nano-switch **3**). The absorption gradually shifted to 550 nm with full completion after 3 min (Fig. S23, ESI†). Metal translocation was ascertained from ESI-MS peaks at 1093.4 and 1876.2 Da attributed to **1**⁺ and $[\text{Cu}(\text{3})]^+$, respectively, but equally a signal at 938.6 Da suggestive of $[\text{Cu}(\text{3})]^{2+}$ was noticed (Fig. S24, ESI†). After reduction with **dmfc** (1 equiv.) the Q band was restored at 561 nm within 3 min (Fig. S23, ESI†). ESI-MS supported the reversal by showing a signal at 1155.5 Da revealing the formation of $[\text{Cu}(\text{1})]^+$ (Fig. S25, ESI†). Similarly, ¹H NMR data, now using the **BFD**¹³ derivative for reduction, indicate complete reverse translocation of the Cu^+ from switch $[\text{Cu}(\text{3})]^+$ to **1** (Fig. S22, ESI†). Thus, the battery of results is clearly in full accord with the required selectivity changes upon oxidation/reduction.

Finally, we focused on stepwise switching with all three nano-switches being present in solution. In both directions of metal translocation, at first the least stable complex was prepared and then the other switches with increasing binding ability were added sequentially. Thus, to test Cu^+ transfer, complex $[\text{Cu}(\text{3})]^+$ was prepared first, then switches **2** and **1** were added sequentially,

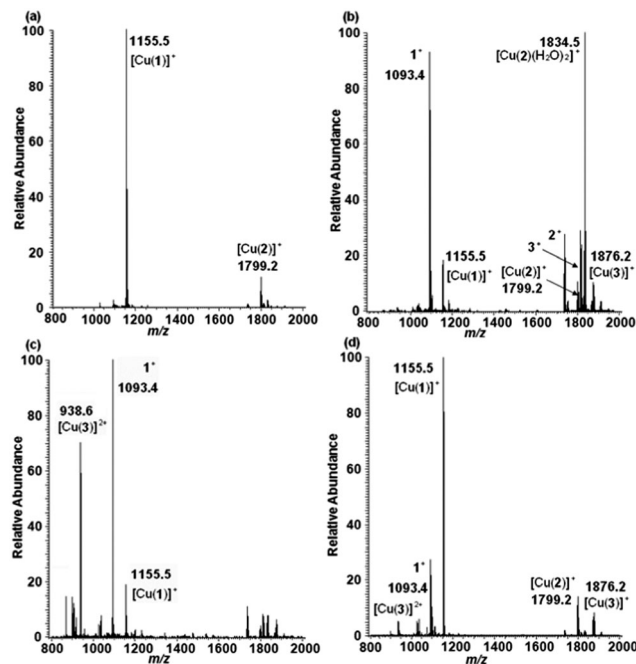


Fig. 3 ESI-MS signature (a) of a solution containing **1**, **2**, **3** and Cu^+ (1 : 1 : 1) in DCM. The solution after oxidation with (b) one or (c) two equiv. of $\text{TBPA}^+\text{SbCl}_6^-$. (d) After reduction of (c) with two equiv. of **dmfc**. Full spectra, see Fig. S28 and S31–S34 (ESI†).

each addition recorded by ¹H NMR. After the first addition,^{6d} a 90 : 10 ratio between $[\text{Cu}(\text{2})]^+$ and $[\text{Cu}(\text{3})]^+$ was established (Fig. S26, ESI†). Finally, after adding **1**, a new equilibrium emerged with $[\text{Cu}(\text{1})]^+$ being formed predominantly in 90% yield, whereas $[\text{Cu}(\text{2})]^+$ was present in only 10% yield and switch **3** was completely devoid of Cu^+ ions (Fig. 1e and Fig. S27, ESI†). Both ¹H NMR and ESI-MS spectra, the latter (Fig. 3a) displaying signals at 1155.5 Da for $[\text{Cu}(\text{1})]^+$ and at 1799.2 Da for $[\text{Cu}(\text{2})]^+$, thus fully match with the data obtained from individual metal complexation studies involving two switches only. A similar protocol was employed for the Zn^{2+} translocation. As expected, in the presence of all switches, the metal ion distributions were similar to those found in the individual dual-switch studies. As a matter of fact, nano-switches **1** and **2** remained completely uncomplexed in the presence of Zn^{2+} with $[\text{Zn}(\text{3})]^{2+}$ being the only coordination complex (Fig. 1f and Fig. S29, ESI†), also proven from the ESI-MS data showing only one signal at 939.1 Da (Fig. S30, ESI†).

Lastly, to investigate multistep communication by using only electron transfer as the input, chemical oxidation and reduction steps were implemented (*vide supra*). For characterisation, we applied the ESI-MS and UV-Vis techniques as each step produced paramagnetic component(s). The equilibrium mixture of switches **1**, **2** and **3** in the presence of Cu^+ (ratio 1 : 1 : 1 : 1, Fig. 3a) was treated with one equiv. of $\text{TBPA}^+\text{SbCl}_6^-$ and the resulting solution was injected into the ESI-MS instrument. The spectrum (Fig. 3b) shows the presence of **1**⁺, $[\text{Cu}(\text{2})]^+$, and $[\text{Cu}(\text{2})(\text{H}_2\text{O})_2]^+$, which were assigned from their signals at 1093.4, 1799.2, and 1834.5 Da. Only traces of $[\text{Cu}(\text{1})]^+$ and $[\text{Cu}(\text{3})]^+$ were visible at 1155.5 and 1876.2 Da, respectively. The relative abundance pattern clearly implies a translocation of Cu^+ from switch **1**⁺



to 2, which goes along with a reorganisation of the arm in 2 from the porphyrin to the phenanthroline station as visible in the UV-Vis spectrum (Fig. 2b, black \rightarrow red trace). Subsequently, the mixture was reacted with one more equivalent of **TBPA** $^{+}$ SbCl_6^{-} to oxidise $[\text{Cu}(2)]^{+}$, which triggered Cu^{2+} translocation as seen from ESI-MS signals at $m/z = 1093.4$ and 938.6 Da for **1** $^{+}$ and $[\text{Cu}(3)]^{2+}$, respectively (Fig. 3c). Despite the reorganisation at both nanoswitches 2 and 3, the amount of $N \rightarrow \text{zinc(II)}$ porphyrin coordination remains in essence constant (Fig. 2b, red \rightarrow green trace). The backward process was initiated by adding one equiv. of **dmfc** as a reducing agent, which, however, should reduce **1** $^{+}$ and not $[\text{Cu}(3)]^{2+}$ by oxidation potential considerations. Indeed, no reorganisation of the arm and thus no severe changes in the UV-Vis spectrum were observed (Fig. 2b, green \rightarrow blue trace). Finally, after treatment with one additional equivalent of **dmfc** to reduce $[\text{Cu}(3)]^{2+}$, Cu^{+} translocation from $[\text{Cu}(3)]^{+}$ to **1** generated $[\text{Cu}(1)]^{+}$ as demonstrated by the ESI-MS spectrum (Fig. 3d). Diagnostically, the process was accompanied by a large nanomechanical reorganisation of the moving arm as evidenced from UV-Vis data at the Q band (Fig. 2b, blue \rightarrow cyano trace). Thus, the trio of nanoswitches has been reset to its initial position after two oxidation and two reduction steps.

Herein, we establish an unprecedented two-step communication cascade between three nanoswitches by programming both metal ion and ligand binding properties depending on their redox states (self-sorting). Ligand oxidation in $[\text{Cu}(1)]^{+}$ at $E_{1/2} = 610 \text{ mV}_{\text{SCE}}$ triggered the release of Cu^{+} acting as an input signal for the next switch. Upon second one-electron oxidation at $[\text{Cu}(2)]^{+}$ ($E_{1/2} = 750 \text{ mV}_{\text{SCE}}$), 6d translocation of Cu^{2+} changed the switching state of **3** $\rightarrow [\text{Cu}(3)]^{2+}$. A two-electron reduction reversed the system to its initial state. Thus, this example not only introduces multistep communication in systems chemistry 14 but also advances the interconnection of molecular gates and machinery.

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Notes and references

- 1 D. L. Nelson and M. M. Cox, *Lehninger Principles of Biochemistry*, W. H. Freeman and Company, New York, 5th edn, 2008.
- 2 (a) S. G. Bell and B. L. Vallee, *ChemBioChem*, 2009, **10**, 55; (b) H. J. Forman, M. Maiorino and F. Ursini, *Biochemistry*, 2010, **49**, 835.
- 3 (a) M.-M. Russew and S. Hecht, *Adv. Mater.*, 2010, **22**, 3348; (b) *Molecular Switches*, ed. B. L. Feringa and W. R. Browne, Wiley-VCH, Weinheim, 2nd edn, 2011; (c) A. C. Fahrenbach, S. C. Warren, J. T. Incorvati, A.-J. Avestro, J. C. Barnes, J. F. Stoddart and B. A. Grzybowski, *Adv. Mater.*, 2013, **25**, 331.
- 4 Some recent reviews: (a) M. von Delius and D. A. Leigh, *Chem. Soc. Rev.*, 2011, **40**, 3656; (b) A. Coskun, M. Banaszak, R. D. Astumian, J. F. Stoddart and B. A. Grzybowski, *Chem. Soc. Rev.*, 2012, **41**, 19; (c) S. F. M. van Dongen, S. Cantekin, J. A. A. W. Elemans, A. E. Rowan and R. J. M. Nolte, *Chem. Soc. Rev.*, 2014, **43**, 99; (d) P. Ceroni, A. Credi and M. Venturi, *Chem. Soc. Rev.*, 2014, **43**, 4068.
- 5 (a) F. M. Raymo and S. Giordani, *Org. Lett.*, 2001, **3**, 3475; (b) S. Silvi, E. C. Constable, C. E. Housecroft, J. E. Beves, E. L. Dunphy, M. Tomasulo, F. M. Raymo and A. Credi, *Chem. – Eur. J.*, 2009, **15**, 178.
- 6 (a) S. Silvi, A. Arduini, A. Pochini, A. Secchi, M. Tomasulo, F. M. Raymo, M. Baroncini and A. Credi, *J. Am. Chem. Soc.*, 2007, **129**, 13378; (b) H. Kai, S. Nara, K. Kinbara and T. Aida, *J. Am. Chem. Soc.*, 2008, **130**, 6725; (c) D. Ray, J. T. Foy, R. P. Hughes and I. Aprahamian, *Nat. Chem.*, 2012, **4**, 757; (d) S. Pramanik, S. De and M. Schmittel, *Angew. Chem., Int. Ed.*, 2014, **53**, 4709.
- 7 M. L. Saha, S. De, S. Pramanik and M. Schmittel, *Chem. Soc. Rev.*, 2013, **42**, 6860.
- 8 M. L. Saha, S. Neogi and M. Schmittel, *Dalton Trans.*, 2014, **43**, 3815.
- 9 J.-P. Sauvage, *Acc. Chem. Res.*, 1998, **31**, 611.
- 10 (a) M. M. Safont-Sempere, G. Fernández and F. Würthner, *Chem. Rev.*, 2011, **111**, 5784; (b) M. L. Saha and M. Schmittel, *Org. Biomol. Chem.*, 2012, **10**, 4651.
- 11 S. Kume and H. Nishihara, *Chem. Commun.*, 2011, **47**, 415.
- 12 (a) M. Schmittel, S. De and S. Pramanik, *Angew. Chem., Int. Ed.*, 2012, **51**, 3832; (b) M. Schmittel, S. Pramanik and S. De, *Chem. Commun.*, 2012, **48**, 11730; (c) S. De, S. Pramanik and M. Schmittel, *Dalton Trans.*, 2014, **43**, 10977.
- 13 3-(11-Bromoundecyl)-1,1'-biferrocenylene is used in this study as a reductant. $E_{1/2}(\text{BFD}) = 0.09 \text{ V}_{\text{SCE}}$ and $E_{1/2}(\text{BFD}^{+}) = 0.76 \text{ V}_{\text{SCE}}$ (see (a) R. Breuer and M. Schmittel, *Organometallics*, 2012, **31**, 6642; (b) R. Breuer and M. Schmittel, *Organometallics*, 2013, **32**, 5980).
- 14 J. R. Nitschke, *Nature*, 2009, **462**, 736.

