

Light-powered autonomous and directional molecular motion of a dissipative self-assembling system

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Biomolecular motors convert energy into directed motion and operate away from thermal equilibrium. The development of dynamic chemical systems that exploit dissipative (non-equilibrium) processes is a challenge in supramolecular chemistry and a premise for the realization of artificial nanoscale motors. Here, we report the relative unidirectional transit of a non-symmetric molecular axle through a macrocycle powered solely by light. The molecular machine rectifies Brownian fluctuations by energy and information ratchet mechanisms and can repeat its working cycle under photostationary conditions. The system epitomizes the conceptual and practical elements forming the basis of autonomous light-powered directed motion with a minimalist molecular design.

Molecular machines and motors are extensively used by living organisms to perform crucial functions such as transport of substrates and mechanical actuation^{1,2}. Synthetic molecular machines³⁻⁵ capable of performing tasks as motor proteins do must exhibit directionally controlled motion by dissipating energy from a constant external source in an autonomous fashion³⁻⁶. These aspects are strictly related to the ability of the device to operate away from chemical equilibrium—a condition that is inherent to biochemical systems⁷ but represents a formidable challenge for artificial ones^{5,6,8}. Despite laboratory demonstration of the potential utility of molecular machines for applications in catalysis⁹⁻¹², materials science¹³⁻¹⁶, information and communication technology¹⁷ and nanomedicine¹⁸, the realization of directional and autonomous molecular movements is a very difficult endeavour, usually tackled with sophisticated chemical structures and/or operation procedures³⁻⁵.

We previously described¹⁹ the directionally controlled transit of a non-symmetric molecular axle through a macrocycle, obtained by ratcheting the self-assembly potential energy profile by means of sequential photochemical and chemical stimulation²⁰. Building upon these results we now report the relative unidirectional transit of an oriented axle through a macrocyclic ring using light as the sole stimulus²¹. This molecular machine exhibits autonomous behaviour as it can repeat its operation cycle indefinitely under photostationary conditions. Our experiments demonstrate that the device takes advantage of light energy to operate away from equilibrium, thus providing a significant example of a dissipative self-assembling system⁶. The results also suggest that both energy and information ratchet mechanisms can account for the rectification of thermal fluctuations brought about by photons²². A further element of interest is that the system consists of wholly synthetic unsophisticated compounds and its operation is based on a very efficient, reversible and clean photoreaction, namely, azobenzene $E \rightleftharpoons Z$ photoisomerization²³.

Design, preparation and photoreactivity of the system

The molecular components of the present system are (Fig. 1a) the macrocyclic ring 2,3-dinaphtho[24]crown-8 ether **1**²⁴ and the

molecular axle $E-2^+$, comprising a photoswitchable E -azobenzene unit at one end, a central ammonium recognition site for the macrocycle and a passive methylcyclopentyl pseudo-stopper at the other end¹⁹. As shown schematically in Fig. 1b, the ring should enter exclusively from the E -azobenzene side of the axle for kinetic reasons, affording a pseudorotaxane $[1 \supset E-2]^+$ in which the macrocycle encircles the recognition site on account of hydrogen-bonding interactions between the oxygen atoms and the ammonium centre and, possibly, π -stacking forces involving the naphthalene and azobenzene units²⁵. Subsequently, light irradiation converts the E -azobenzene unit into the bulkier Z form, a process that should also cause a destabilization of the supramolecular complex $[1 \supset Z-2]^+$ and dethreading of the components. If the extrusion from the ring is faster for the pseudo-stopper of the axle than for its Z -azobenzene unit, dethreading should occur preferentially along the same direction originally followed for threading. Reset is obtained by photochemical or thermal conversion of the azobenzene gate back to the E isomer, thereby regenerating the starting form of the axle and causing directionally controlled threading once again (Fig. 1b).

In the present system, autonomous light-powered cycling²⁶⁻²⁹ between the states shown in Fig. 1b under steady irradiation is possible because the same photons can trigger both $E \rightarrow Z$ and $Z \rightarrow E$ photoisomerizations, owing to the overlapping absorption spectra and photoreactivity of E - and Z -azobenzene. Overall, the photo-induced relative unidirectional transit of the axle through the macrocycle would be obtained according to a flashing energy ratchet mechanism (Fig. 1c), in which a single photonic stimulus controls both the relative stabilities of the assembled and disassembled states (that is, switching) and the relative heights of the kinetic barriers for threading/dethreading over either extremities (that is, gating)²².

Directionally controlled threading/dethreading movements

¹H NMR spectra in CD₂Cl₂ (Supplementary Figs 3 and 4) show that both $E-2^+$ and $Z-2^+$ pseudorotaxane-type complexes with macrocycle **1**. The association of **1** and **2**⁺ also causes changes in the UV-vis absorption and luminescence spectra. In particular, the

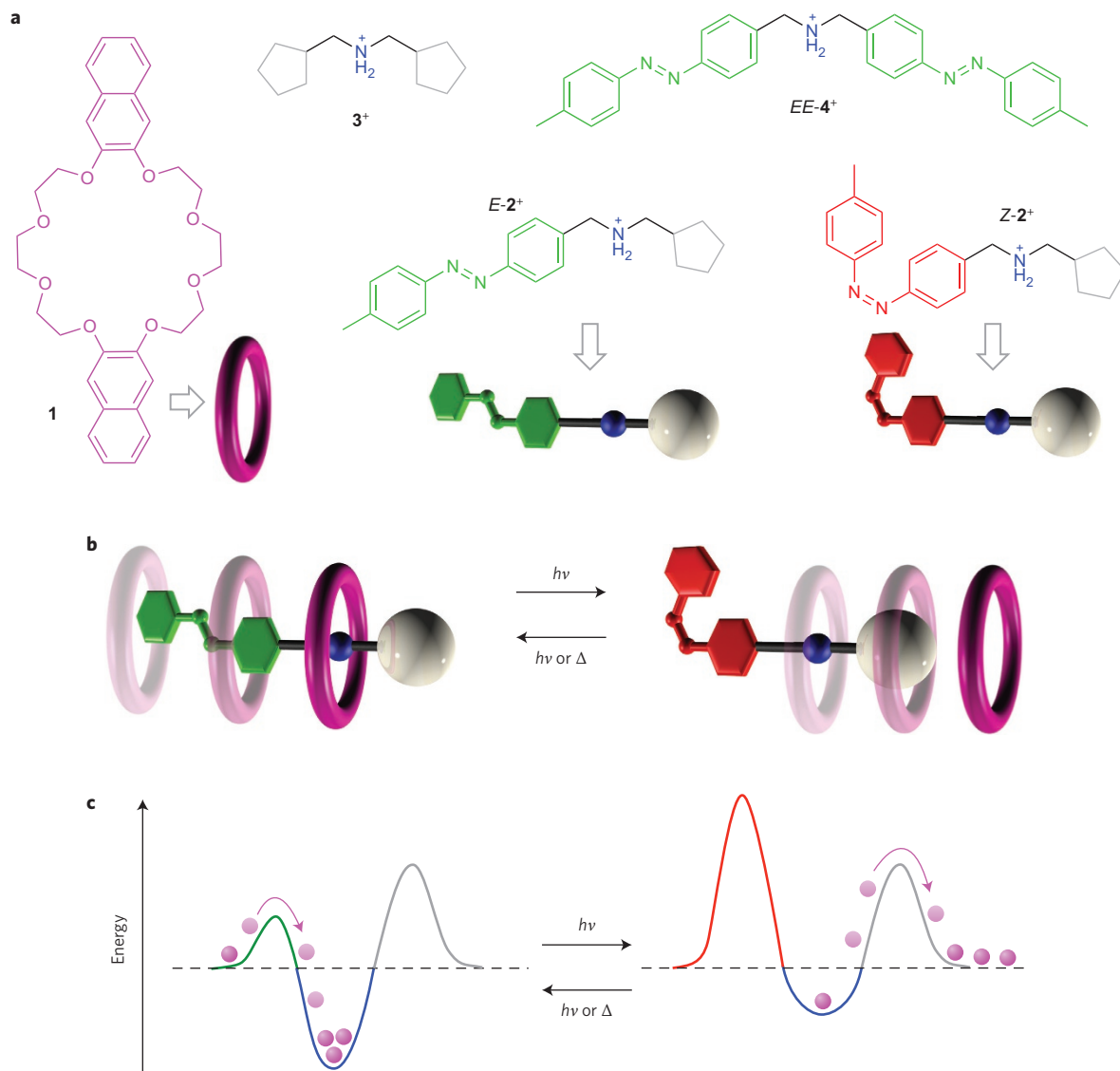


Figure 1 | Design and operation of the system. a, Structure formulae, with cartoon representation, of macrocycle **1**, non-symmetric axes **E-2⁺** and **Z-2⁺**, and symmetric model axes **3⁺** and **EE-4⁺**. **b**, Schematic representation of the relative unidirectional translations of the ring and axle components triggered by light. **c**, Simplified potential energy curves (free energy versus ring-axle distance) for the states shown in **b**, illustrating how periodic modulation of the energy barriers and wells can produce directional and repetitive threading–dethreading movements.

Table 1 | Kinetic and thermodynamic data for self-assembly of the investigated complexes.

Complex	K^* (M^{-1})	$-\Delta G^{\circ\dagger}$ (kcal mol ⁻¹)	$k_{in}^{##}$ ($M^{-1} s^{-1}$)	$\Delta G_{in}^{\ddagger\ddagger}$ (kcal mol ⁻¹)	k_{out} (s^{-1})	$\Delta G_{out}^{\ddagger\ddagger}$ (kcal mol ⁻¹)
[1 ⊃ E-2] ⁺	6.3×10^5	7.8	54	14.8	$2.2 \times 10^{-5\ddagger}$ $8.6 \times 10^{-5 }$	23.4
[1 ⊃ Z-2] ^{+\ddagger}	1.7×10^5	7.0	0.81	17.3	$4.7 \times 10^{-6 }$	24.3
[1 ⊃ 3] ⁺	3.2×10^4	6.0	2.2	16.7	$3.8 \times 10^{-5\ddagger}$ $6.9 \times 10^{-5 }$	23.1
[1 ⊃ EE-4] ⁺	$>10^7$	>9.4	55	14.8	$<5.5 \times 10^{-6 }$	>24
[1 ⊃ ZZ-4] ^{+\ddagger}	#	#	$3.9 \times 10^{-2**}$	19.0	††	††

*Determined by luminescence or absorption titrations. †Free energies of association (ΔG°), calculated from the K values using the expression $\Delta G^{\circ} = -RT \ln K$. ##Determined by UV-vis absorption or luminescence spectroscopy. ‡Free energies of activation for the threading (ΔG_{in}^{\ddagger}) and dethreading ($\Delta G_{out}^{\ddagger}$) processes, calculated using the relationships $\Delta G_{in}^{\ddagger} = -RT \ln(k_{in}/h/kT)$ and $\Delta G_{out}^{\ddagger} = -RT \ln(k_{out}/h/kT)$, respectively, where R , h and k correspond to the gas, Planck and Boltzmann constants, respectively. ||Calculated from the k_{in} and K values using the expression $k_{out} = k_{in}/K$. ¶Obtained upon exhaustive irradiation of the corresponding *E*-isomer at 365 nm; the *E* → *Z* conversion yield at the photostationary state was >96%. †Not measured because equilibrium cannot be reached before substantial thermal *Z* → *E* back isomerization occurs. **Estimated from the initial part of the time-dependent luminescence changes upon mixing **1** and **ZZ-4⁺**, after deconvolution of the contribution of *Z* → *E* isomerization and threading from the *E*-azobenzene terminus. ††Not determined. Conditions: dichloromethane (CH₂Cl₂), 20 °C.

intense fluorescence of **1** ($\lambda_{max} = 345$ nm) is completely quenched when the ring is complexed by either isomer of axle **2⁺**. The thermodynamic and kinetic data for self-assembly of these complexes,

together with those of the complexes formed by the symmetric model axes **3⁺** and **EE-4⁺** (Fig. 1a) with macrocycle **1**, are reported in Table 1 (Supplementary Table 1, equation (1) and Figs 5–24).

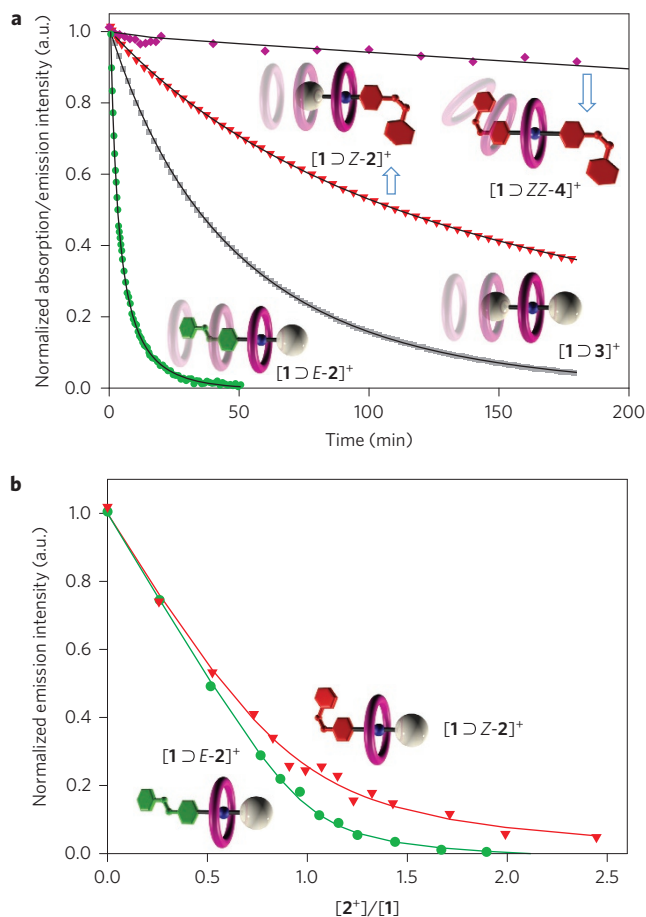


Figure 2 | Kinetic and thermodynamic characterization of the self-assembly process. **a**, Time-dependent luminescence or absorption changes associated with the self-assembly of the complexes studied in this work. Data corresponding to the threading of $[1 \supset EE-4]^+$ are not shown because they fully overlap with those for threading of $[1 \supset E-2]^+$. The solid lines represent the data fitting according to a kinetic model consisting of a second-order threading and an opposed first-order dethreading. Conditions: CH_2Cl_2 , 20 °C, 100 μM for each component. **b**, Titration curves, obtained from luminescence data, observed upon addition of either $E-2^+$ or $Z-2^+$ to a 54 μM solution of **1** in CH_2Cl_2 at 20 °C. Solid lines represent data fitting according to a 1:1 binding model. In all cases, excitation was performed at 272 nm.

Axles $Z-2^+$ and $ZZ-4^+$, as well as their complexes with **1**, can be obtained by exhaustive irradiation of the corresponding E isomers at 365 nm. In all cases $E \rightarrow Z$ conversion at the photostationary state is >96%.

Photochemical experiments performed on the $[1 \supset E-2]^+$ complex show that the $E \rightarrow Z$ isomerization of the azobenzene unit of 2^+ is sensitized upon irradiation in the absorption bands of **1**, indicating that excitation energy is transferred from the naphthalene units of the ring to the E -azobenzene moiety of the axle. Such an energy-transfer process also accounts for the emission quenching, as confirmed by the fact that the luminescence is quenched when **1** is complexed by $EE-4^+$, while it is unaffected in the $[1 \supset 3]^+$ complex. Conversely, the photosensitization of the $Z \rightarrow E$ transformation in the $[1 \supset Z-2]^+$ complex is not efficient because of the poor overlap of the emission spectrum of **1** with the absorption spectrum of the Z isomer of the axle. The significant quenching of the emission of **1** in the $[1 \supset Z-2]^+$ complex can be ascribed to electron transfer from the photoexcited naphthalene moiety of the ring to the lowest unoccupied molecular orbital (LUMO) of the azobenzene axle.

The fact that the threading rate constant for $E-2^+$ is close to that of $EE-4^+$ and 25 times faster than that of 3^+ (Table 1, Fig. 2a) indicates that axle $E-2^+$ enters ring **1** almost exclusively (>96%) with its E -azobenzene terminus. On the other hand, the rate constant for self-assembly of **1** and $Z-2^+$ compares with that for the threading of 3^+ , which, in its turn, is 50 times faster than that for threading of $ZZ-4^+$. All these observations clearly indicate that $Z-2^+$ threads the ring with its methylcyclopentyl terminus, and confirm that the $E \rightleftharpoons Z$ photoisomerization can effectively be used to control the self-assembly kinetics³⁰.

In contrast with the behaviour of an earlier reported system¹⁹, the $E \rightleftharpoons Z$ photoisomerization of 2^+ affects the stability of the resulting pseudorotaxane, as shown by luminescence titration curves (Fig. 2b). The association constant with **1** drops from $6.3 \times 10^5 \text{ M}^{-1}$ for $E-2^+$ to $1.7 \times 10^5 \text{ M}^{-1}$ for $Z-2^+$, most likely as a consequence of a diminished π -stacking of the naphthalene units of **1** with the non-planar Z -azobenzene moiety. Under typical experimental conditions (1:1 mixture of 50 μM **1** and 2^+), exhaustive 365 nm irradiation affords the dethreading of ~15% of the pseudorotaxane species. Hence, the reversible photoisomerization of the axle can also be exploited to switch the system, that is, to change the relative stabilities of the self-assembled and disassembled states (Fig. 1c).

The direction of photoinduced dethreading can be determined by considering that the barriers (or rate constants) for threading with the different termini of a non-symmetric axle give information about the relative heights of the corresponding dethreading barriers. On the basis of the threading rate constants for 3^+ and $ZZ-4^+$ with **1**, it can be calculated that more than 98% of $Z-2^+$ axles will enter the macrocycle with the methylcyclopentyl stopper. It can therefore be concluded that extrusion of the methylcyclopentyl unit is kinetically preferred over Z -azobenzene in the dethreading of $[1 \supset Z-2]^+$. It should also be noted that the latter process ($k = 4.7 \times 10^{-6} \text{ s}^{-1}$) is faster than the thermal $Z \rightarrow E$ isomerization ($k = 1.6 \times 10^{-6} \text{ s}^{-1}$); these numbers indicate that, on average, the complex should have approximately three dethreading chances before its azobenzene gate is thermally converted back to the E configuration. The full reset of the system is determined solely by the $Z \rightarrow E$ isomerization, which restores the original potential energy landscape (Fig. 1c) and promotes the rethreading of $E-2^+$ into **1**, preferentially with the azobenzene terminus (Fig. 1b), thus completing the operation cycle.

Autonomous operation away from equilibrium

It is well known that for a closed path of reactions at thermal equilibrium, detailed balance—a consequence of microscopic reversibility—states that each individual process has to be at equilibrium; that is, no directional flow can occur^{22,31}. In our case, two of the four processes, namely isomerization of the free and complexed axle (Fig. 3, processes 2 and 4), are photochemically driven and hence are not subject to microscopic reversibility^{31–33}. One can define $K_{2,h\nu}$ and $K_{4,h\nu}$ as the ratio of the concentrations of the products and reactants for the respective reactions once the steady state under light irradiation is reached. Our results show that the photostationary states afforded upon irradiation of the free and complexed axle at $\lambda > 400 \text{ nm}$ have the same E/Z composition. This means that $K_{2,h\nu} = K_{4,h\nu}$, but, because $K_1 > K_3$ (the ring associates more strongly with $E-2^+$ than with $Z-2^+$, Table 1), detailed balance, as anticipated above, is not fulfilled (Supplementary Fig. 1 and equation (2)). In other words, at the steady state, the net rates of the individual processes in the closed cycle are equal but non-zero, and the system travels through the cycle one way. In the present case, the cycling path (clockwise in Fig. 3) is dictated by the fact that $K_1 > K_3$. It is worthwhile to remember that, for the reasons described above, the threading–dethreading processes represented in Fig. 3 are directionally controlled; that is, $E-2^+$ and $Z-2^+$ thread/dethread the ring with their E -azobenzene and methylcyclopentyl termini, respectively.

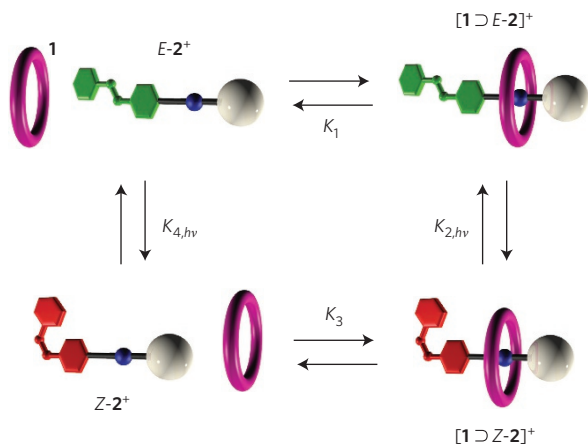


Figure 3 | Square cycle of chemical and photochemical reactions representing operation of the system. Horizontal and vertical arrows denote the self-assembly equilibria and photochemical isomerization reactions, respectively. The indicated parameters refer to the reactions read from left to right and from top to bottom.

Gaining direct experimental evidence of photostationary cycling, however, is not straightforward, because the collective chemical and physical properties of the ensemble exhibit no change at the steady state. On the basis of the above discussion, the supply of light energy is expected to cause cycling—that is, to establish a net flux around the cycle in one way even if the concentration of each species remains static—thus bringing the system out of equilibrium. Therefore, a proof for autonomous light-driven cycling would be the observation that the concentration reached at the photostationary state for any species involved is not consistent with its expected equilibrium value^{28,34,35}.

As the fluorescence of **1** is completely quenched when the ring is complexed by either $E-2^+$ or $Z-2^+$, the fluorescence signal measured under appropriate spectral conditions³⁶ can be correlated to the amount of uncomplexed macrocycle present in the solution. We therefore monitored the concentration of the free ring in a solution containing **1** and **2**⁺ kept under steady irradiation at 365 nm. In an optimized experiment, a solution containing 50 μM **1** and 150 μM $E-2^+$ was allowed to equilibrate in the dark at 20 °C; the self-assembly of the components was revealed by a strong decrease in the emission intensity. Irradiation of the solution with relatively high-intensity (1.7×10^{-6} Einstein min^{-1}) 365 nm light afforded nearly complete $E \rightarrow Z$ conversion in such a short time (5 min) that the self-assembly equilibria could not keep up with the photoisomerization changes. In fact, the concentration of the free ring was unchanged, within error, after irradiation, confirming that the dethreading of a fraction of the $[1 \supset Z-2]^+$ species (expected because its stability constant is lower than that of $[1 \supset E-2]^+$) does not take place on this timescale.

Once the system had been completely photoisomerized (that is, when the absorption spectrum became invariant, Supplementary Figs 20 and 21), the irradiation was resumed with a lower light intensity (1.0×10^{-7} Einstein min^{-1}) to prevent any photodecomposition effect. A decrease in the concentration of the free macrocycle was observed, which reached a plateau at $\sim 6\%$ after ~ 2 h of continuous irradiation (Fig. 4a). This observation indicates a degree of complexation even larger than that expected for $E-2^+$ (that is, the guest form exhibiting the highest affinity for the ring) and it is clearly incompatible with any equilibrium mixture of **1** and (E - or Z -) 2^+ . A control experiment performed on a solution containing the macrocycle and the deprotonated guest $E-2$ (lacking the ammonium recognition site and thus unable to associate with **1**) under otherwise identical conditions showed no change in the emission

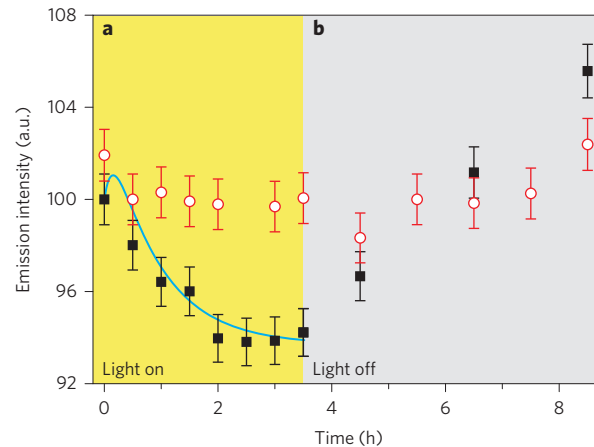


Figure 4 | Observation of photostationary cycling of the system away from equilibrium. **a**, Fluorescence intensity values of uncomplexed macrocycle **1** upon prolonged 365 nm irradiation of a mixture of ring and axle at the $E \rightleftharpoons Z$ photostationary state in CH_2Cl_2 at 20 °C (black squares). The solution was deoxygenated to exclude photodecomposition effects and was prepared by mixing 50 μM **1** and 150 μM $E-2^+$ (98% complexation of **1**) followed by exhaustive irradiation at 365 nm (99% $E \rightarrow Z$ conversion after 5 min). Open circles show the fluorescence intensity values in an identical experiment in which the deprotonated axle **2**, lacking the ammonium centre and thus unable to associate with **1**, is used in place of 2^+ . The solid line is a computer simulation of the process, based on the scheme shown in Fig. 3 and performed using the experimentally determined values for threading, dethreading, photoisomerization and thermal isomerization rates. **b**, Luminescence changes observed in the experiments described in **a**, after irradiation at 365 nm has been switched off. Significant absorption changes, due to $Z \rightarrow E$ thermal isomerization, are observed. Inner filter effects prevent the analysis of data points after ~ 8 h. In all cases, excitation was performed at 365 nm. Error bars represent the standard deviation of the luminescence intensity measurement under the experimental conditions. See main text for more details.

intensity upon prolonged irradiation (Fig. 4a), confirming that the observed effect arises from the interplay of photoisomerization and self-assembly processes (Fig. 3).

Such an effect can be understood by considering that dethreading of $[1 \supset Z-2]^+$ through the methylcyclopentyl group is the slowest of all the processes of the cycle (Table 1). Accordingly, a larger concentration of this species at the photostationary state with respect to dark equilibrium conditions is expected. In practice, while the cycle operates, reaction 2 (Fig. 3) populates the $[1 \supset Z-2]^+$ state, characterized by very slow dissociation kinetics, scavenging $[1 \supset E-2]^+$ and thus fostering the complexation of $E-2^+$ generated in reaction 4 with the concomitant decrease of free ring concentration. This mechanism is efficient because, under steady irradiation, light continuously and repeatedly triggers both $E \rightarrow Z$ and $Z \rightarrow E$ isomerization.

When the light is switched off, $[1 \supset Z-2]^+$ begins to dethread because its concentration is higher than the equilibrium value and the amount of the free ring initially increases (Fig. 4b). As the slow concomitant $Z \rightarrow E$ thermal isomerization regenerates the E -isomer, the ring concentration is expected to level off as time goes on. Unfortunately, the absorption changes associated with the isomerization prevent an accurate determination of the fluorescence intensity because of the strong reabsorption of the emitted light³⁶. Nevertheless, after 20 days in the dark at room temperature—that is, when all the isomerization and self-assembly processes have reached equilibrium—the observed emission intensity is identical to that measured for the equilibrated solution of **1** and $E-2^+$ before irradiation. This result highlights the full reversibility and remarkable fatigue resistance of this ensemble.

Energetic and mechanistic considerations

To verify our interpretation of the observed results we performed numeric simulations of the cycle shown in Fig. 3 using the experimentally determined values for threading, dethreading, photoisomerization and thermal isomerization rates (Table 1, Supplementary Fig. 2 and Supplementary equations (3) to (10)). The excellent match between the simulation and the measured data points (Fig. 4a) confirms the validity of our mechanistic hypotheses.

A few interesting facts can be extracted from the simulation to assess the performance of the cycle (Supplementary equations (11) to (13)). The cycling rate is $1.7 \times 10^{-10} \text{ M s}^{-1}$ and the efficiency is 2.3×10^{-3} cycles per photon; that is, about 430 photons need to be absorbed on average to complete a cycle. The product of the rate constants of the clockwise processes divided by the product of the rate constants of the anticlockwise processes (Fig. 3) indicates that, on average, the system loops in the 'wrong' way (that is, anticlockwise), once every 160 cycles. The maximum amount of work that could be performed in a cycle, which can be estimated from the free energy change of the system^{22,37}, is $5.1k_{\text{B}}T$ ($3.0 \text{ kcal mol}^{-1}$ at $20 \text{ }^{\circ}\text{C}$), that is, about a quarter of the energy provided by adenosine triphosphate (ATP) hydrolysis. Considering that, under the conditions employed, one cycle uses ~ 430 photons of 365 nm light (78 kcal mol^{-1}), the upper limit for the energy conversion efficiency is 9×10^{-5} .

A closer inspection of the photochemical data (Supplementary Table 2) enables a discussion about the mechanisms used by this system to rectify Brownian fluctuations²². First, we found that the $E \rightarrow Z$ conversion at the photostationary state upon irradiation at 365 nm is slightly larger for the complex than for the free axle ($K_{2,h\nu} > K_{4,h\nu}$). As the photoisomerization quantum yields are not affected by the presence of the macrocycle, the more efficient $E \rightarrow Z$ transformation in the complex must arise from its slightly higher molar absorption coefficient at 365 nm in comparison with the free axle. The situation is even more interesting when the system is irradiated with light also absorbed by the ring component (for example, 287 nm). Because excitation energy is efficiently transferred from **1** to $E\text{-}2^+$ but not to $Z\text{-}2^+$ in their respective complexes (and, as noted above, the inherent photoisomerization efficiency of the axle does not change when it is surrounded by the ring), irradiation of the ring-axle mixture at this wavelength generates a photostationary state with a larger Z/E composition than for the axle alone and, again, $K_{2,h\nu} > K_{4,h\nu}$.

The device operation is based on pure energy ratcheting^{22,38} if irradiation is performed with $\lambda > 400 \text{ nm}$ ($K_{2,h\nu} = K_{4,h\nu}$ as discussed above), whereas a contribution from information ratcheting^{22,34,35} takes place with irradiation at 365 or 287 nm , although via different physical phenomena. The ring does not absorb light at 365 nm , but its presence enhances the absorption of the E -axle, leading to the preferential light-triggered closure of the azobenzene gate in the pseudorotaxane rather than the free axle. On the other hand, the light absorbed by the ring at 287 nm causes the transfer of excitation energy to the E -axle, a process that can efficiently occur only if the components are bound, and triggers the $E \rightarrow Z$ isomerization of the latter³⁴. In both cases, information about the location of the ring with respect to the axle can control the gating³¹.

Conclusions

We have described a dissipative self-assembling ensemble that uses light energy to perform directed molecular movements in a repetitive fashion out of equilibrium⁶. Although the system as such cannot be used to perform work in a bulk solution³⁷, it provides a viable route to the construction of rotary motors based on catenanes³⁹ or linear motors⁴⁰ based on polymeric motifs^{9,11}, which are capable of moving—and possibly transporting cargo—over long distances. It can also be regarded as a precursor of artificial molecular pumps³⁸

capable of generating concentration gradients across membranes^{41–44}. With this aim in mind we are applying the above strategy to three-dimensional macrocycles, which can be included in bilayer membranes^{45,46} and in which face-selective threading is possible^{47,48}.

Besides its practical applications, this system is of great interest because it epitomizes the conceptual and practical elements at the basis of repetitive directionally controlled molecular motion powered by light^{22,31,37}. Its minimalist design, facile synthesis, convenient stimulation and excellent reversibility are essential features that will foster further research developments as well as a fundamental premise for real-world applications.

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Author contributions

M.B. synthesized the compounds. G.R., S.S. and M.B. performed the physico-chemical experiments. G.R. carried out numerical simulations. A.C. conceived the project and wrote the paper. M.V. discussed the results and commented on the manuscript, together with all authors.

Competing financial interests

The authors declare no competing financial interests.