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A Focus on Triazolium as a Multipurpose Molecular Station for pH-Sensitive Interlocked Crown-Ether-Based Molecular Machines

Frédéric Coutrot*^[a]

The control of motion of one element with respect to others in an interlocked architecture allows for different co-conformational states of a molecule. This can result in variations of physical or chemical properties. The increase of knowledge in the field of molecular interactions led to the design, the synthesis, and the study of various systems of molecular machinery in a wide range of interlocked architectures. In this field, the discovery of new molecular stations for macrocycles is an attrac-

tive way to conceive original molecular machines. In the very recent past, the triazolium moiety proved to interact with crown ethers in interlocked molecules, so that it could be used as an ideal molecular station. It also served as a molecular barrier in order to lock interlaced structures or to compartmentalize interlocked molecular machines. This review describes the recently reported examples of pH-sensitive triazolium-containing molecular machines and their peculiar features.

1 Introduction

Since the conformational state of a molecule is closely related to its intrinsic properties, interlocked molecular architectures, or more precisely, interlocked molecular machines,^[1] for which a component can be mechanically moved with control relative to another, can be considered of high interest. Among interlocked molecules, rotaxanes occupy an important place. They comprise a macrocycle which surrounds a molecular axle that is usually ended by bulky stoppers in order to avoid any disassembly of the structure.

If the chemical system is designed with the aim of acting as a molecular machine, different sites of interactions (i.e. molecular stations) must be present in the threaded axle so that each of them can bind the macrocycle with different affinities. Decreasing the interactions of the best molecular station for the macrocycle or increasing those of the poorest molecular station can result in the controlled shuttling of the macrocycle along the molecular thread. More exactly, this change excites the molecular system out of equilibrium, before it relaxes to the energetically favored co-conformation via the so-called Brownian motion. In these singular chemical species, not only the presence of the macrocycle around the threaded axle but also its controlled variable localization can give rise to the

modulation of the chemical and physical properties of the molecule. Hence, novel molecular stations for various macrocycles have been the subject of various research endeavors in the last two decades.

In order to reversibly tune the affinity of a molecular site of interaction for the macrocycle, different kinds of stimuli have been used to trigger the shuttling of the macrocycle along the thread: pH,^[2] temperature^[3] or solvent variation,^[4] and photo-,^[5] electro-,^[6] or chemical^[7] reactions. Among the plethora of systems that have been described until now, we would like to focus in this review on the pH-dependent molecular machines that are exclusively based on the interactions of crown ethers with ammonium and triazolium^[8] molecular stations. For a long time, the ammonium motif has been known to interact quite strongly with crown ethers such as 24-crown-8,^[9] so that it can serve as a template for driving the rotaxane formation. On the other hand, the triazolium moiety was only reported in 2008 as a weak molecular station for dibenzo-24-crown-8 (DB24C8).^[13,39] Therefore, systems containing ammonium and triazolium molecular stations for crown ethers can be quite efficiently used for designing new molecular machines. More precisely, because of their quite different affinities for crown ethers, the ammonium moiety can act as the best site of interactions, whereas the triazolium becomes the predominant site of localization of the macrocycle in basic medium after deprotonation of the ammonium, because no more interaction remains between the revealed amine and the crown ether.

The easy access to triazolium compounds makes the synthetic route to such interlocked molecular machines straightforward and versatile. A particular asset of the system is the fact that triazolium can be obtained from azide and alkyne fragments, which allows any kind of chemical variations on the triazole moiety depending on the precursors used. When

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making the triazolium molecular station through nucleophilic substitution, it is interesting to note the possibility to graft on the triazole nitrogen atom any moiety which can play different roles. Thus, so far, different triazolium-containing molecular architectures, from rotaxane to the more sophisticated double-lasso rotamacrocycle via molecular muscles, have been synthesized and studied.

Some of these molecular machines have only been prepared to overcome a synthetic challenge or for a purely “chemical design” aim, whereas others have already found utility as switchable catalysts for organic reactions, switchable fluorescent probes, or extendable materials. In all the reported structures, different molecular motions arising from the interlocked architecture have been accurately studied. Some of them directly concern the translation of the macrocycle in a [2]rotaxane, which can be accompanied in some cases by a rotary motion, whereas others relate the tightening/loosening of lasso compounds, or even a molecular “jump rope” motion in a double-lasso structure. Triazolium can also be used as a multi-purpose molecular station that can act as a kinetic barrier in order to trap the macrocycle in a compartment of the threaded axle, making the system unbalanced and not at equilibrium.

In this review, we propose to give the current state of art on triazolium-based crown ether molecular machines. We will first describe the general synthetic accesses to triazolium-based interlocked molecules. We will then successively highlight examples that have been reported with a pure chemical design aim, and those designed for a defined application. A following section will be devoted to triazolium-based molecular muscles and their use as building blocks in polymers. The relative affinity of the DB24C8 for triazolium and pyridinium amide molecular stations will then be exemplified. This will lead us to discuss about the compartmentalization of three-station-containing molecular machines by using triazolium as both a molecular station and a barrier for the macrocycle. The final sections will relate on more sophisticated mono- and double-lasso architectures with their relative motions.

Frédéric Coutrot received his PhD in 1999 from the University Henri Poincaré (Nancy, France). He then joined the group of Dr. Michel Marraud (Ecole Nationale Supérieure des Industries Chimiques, Nancy) as a temporary lecturer. In 2000, he moved to David Leigh's laboratory with a Marie-Curie Fellowship, first at Warwick University (England), then at King's Building in Edinburgh (Scotland). Since 2002, he has been an Assistant Professor at the University of Montpellier (France) and is currently the team leader of the Supramolecular Machines and Architectures Team at the Institut des Biomolécules Max Mousseron (IBMM). His research focuses on the synthesis and study of new interlocked molecular machines.



2 Synthetic Accesses to Triazolium-Based Interlocked Compounds and ^1H NMR Characterization

2.1 General synthetic access to triazolium-containing rotaxane molecular machines

Triazolium-based rotaxane molecular machines are usually obtained through a threading-capping strategy using a two-step sequence beginning with a copper(I)-catalyzed Huisgen^[10] alkyne–azide 1,3-dipolar cycloaddition,^[11] also called “CuAAC click chemistry”, then by the subsequent alkylation of the triazole (Figure 1).

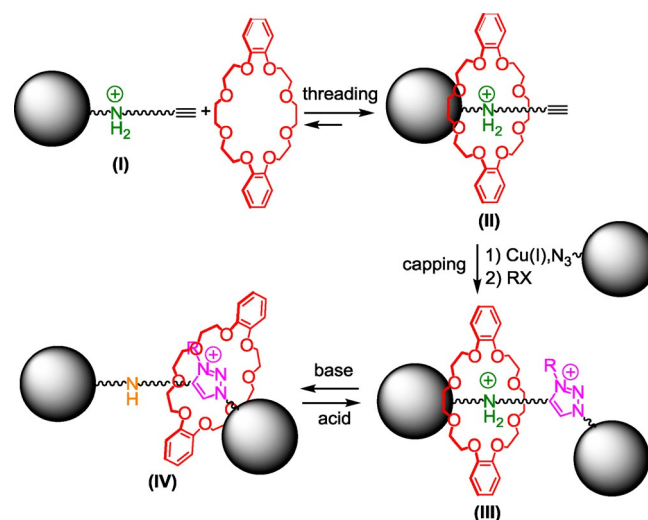


Figure 1. General chemical route to triazolium-containing molecular machines.

The classical straightforward chemical route to triazolium-containing molecular machines begins from a molecular axle I that bears a secondary ammonium moiety as the template to thermodynamically drive the formation of the semi-rotaxane II and an alkyne function at one extremity of the thread for further chemical connection. The dibenzo-24-crown-8 interacts quite well with the ammonium template, so that the conversion to the semi-rotaxane II can be generally quantitative. The CuAAC click reaction between the alkyne-containing semi-rotaxane II and the azido “stopper” groups then allows the capping of the threaded axle and affords the locked [2]rotaxane architecture III. The inverse was also found in the literature, that is to say an alkyne stopper group with an azido-containing semi-rotaxane. Subsequent regioselective N-alkylation of the triazole provides the triazolium as the second molecular station. This latter has a much poorer affinity for the crown ether than the ammonium. As a consequence, in the two-station-based [2]rotaxane III, the macrocycle mainly resides around the ammonium site. However, deprotonation of the ammonium molecular station triggers the translational motion of the crown ether toward the triazolium station (in IV). The process can be reversed by adding an acid. Interestingly, al-

though triazolium interacts well with crown ether in a locked rotaxane molecular architecture, this moiety cannot be used as a template to prepare rotaxane from disassembled elements, because the intermolecular interactions between crown ether and triazolium are much too weak. Indeed, when attempting to thread a crown ether with a triazolium-containing axle, one may assume that the enthalpic energy (this latter being directly related to the weak noncovalent interactions between the elements) cannot overcome the opposite change in entropic energy. On contrary, in a mechanically interlocked [2]rotaxane, the change in entropies between the deprotonated and the protonated rotaxanes seems much lower. In that case, the enthalpic factor which is driven by interactions between triazolium and the crown ether become predominant, which corroborates the fact that triazolium is a good molecular station for the macrocycle in such a locked structure. Taking into account such considerations, we recently imagined a diverted strategy to prepare a wide range of triazolium-based rotaxanes that are devoid of any other template.

2.2 A diverted strategy to yield triazolium rotaxane devoid of any other template

Since the triazolium moiety cannot be utilized as an efficient template for the formation of crown-ether-based rotaxane, we recently reported a general diverted strategy to yield triazolium-based rotaxanes (Figure 2).^[12]

The strategy relies on the use of a so-called macrocycle “transporter” **A** which contains an ammonium template in order to efficiently bind a crown ether. Compound **A** has also been designed to be further extended at one of its extremities with a triazolium-containing axle through a *N*-hydroxysuccinimide (NHS) ester bond. Once connected, the extended ammonium- and triazolium-based rotaxane **C** can undergo co-conformational changes through pH variation. Indeed, deprotonation

of the ammonium station triggers the shuttling of the macrocycle from its initial ammonium location to the triazolium **D**. The NHS ester moiety, which has been chosen as a linker between the two initial axle fragments, allows for the contraction of the axle when reacting with an amine. This last reaction (step 4) provides triazolium-based rotaxane **E**, while the macrocycle “transporter” is recycled at the same time (step 5). One of the advantages of this synthesis remains in the fact that the loaded macrocycle transporter **B** can be purified and stored over time. Moreover, the possibility to recycle the transporter is one more asset that makes this strategy very attractive and quite direct from **B**.

Because it is very recent, this second strategy has not yet been widely used. On contrary, the strategy reported in 2008 has now been used by several groups to prepare triazolium-based molecular machines which are described hereafter.

2.3. Usual ¹H NMR characterization of ammonium and *N*-methyltriazolium in interactions with DB24C8 in a pH-sensitive [2]rotaxane molecular machine

The weak interactions between the DB24C8 for the ammonium and the triazolium moieties induce typical displacements in ¹H NMR chemical shifts with respect to the free elements (Figure 3).

In all the interlocked structures, the NMR signals of the hydrogen atoms of the crown ether are split because they are facing the two nonsymmetrical ends of the threaded axle. Moreover, the hydrogen atoms H^F, and to a lesser extent H^D, are usually shielded because they are located in the shielding cavity of the benzylammonium moiety. There are multiple influences of the crown ether on the thread. At the protonated state, the DB24C8 mainly interacts with the ammonium station through hydrogen bonds and ion-dipole interactions. In this co-conformation, hydrogen atoms H¹⁻³, which belong to the ammonium site, are all dramatically shifted downfield, due to their hydrogen-bonding interactions with the oxygen atoms of the DB24C8. At the same time, methylene hydrogens H⁴⁻⁹, which are next to the ammonium, experience more or less the shielding effect of the aromatic rings of the DB24C8. No other chemical changes are observed for the other part of the threaded molecular axle, corroborating the localization of the macrocycle. At the deprotonated state, the DB24C8 interacts with the triazolium site. This new localization can be evidenced by the usual main ¹H NMR variations. With respect to the free molecular axle, H^{2'} and in a lesser extent H^{4'} are both shifted downfield in

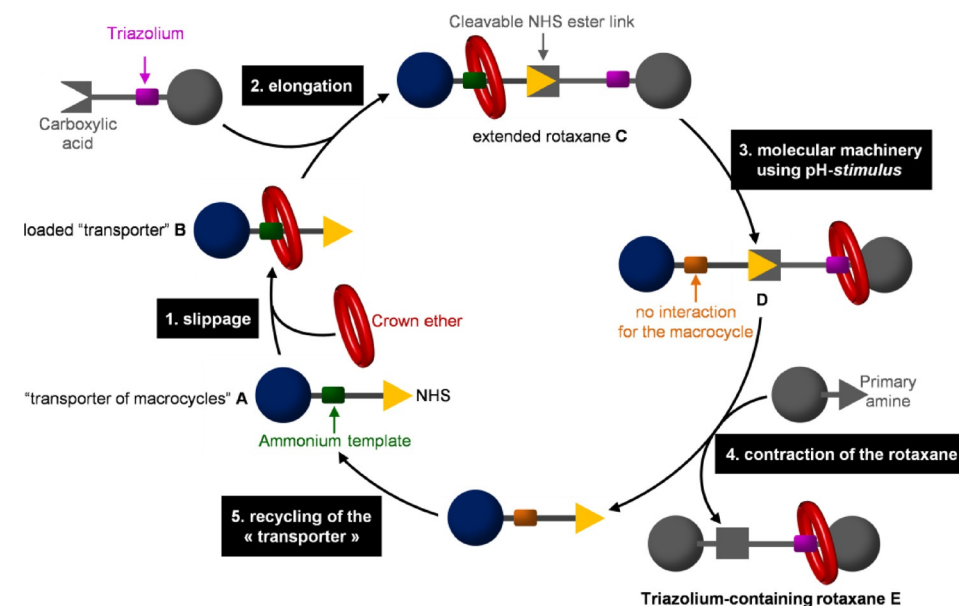


Figure 2. A diverted route to triazolium-containing rotaxanes.

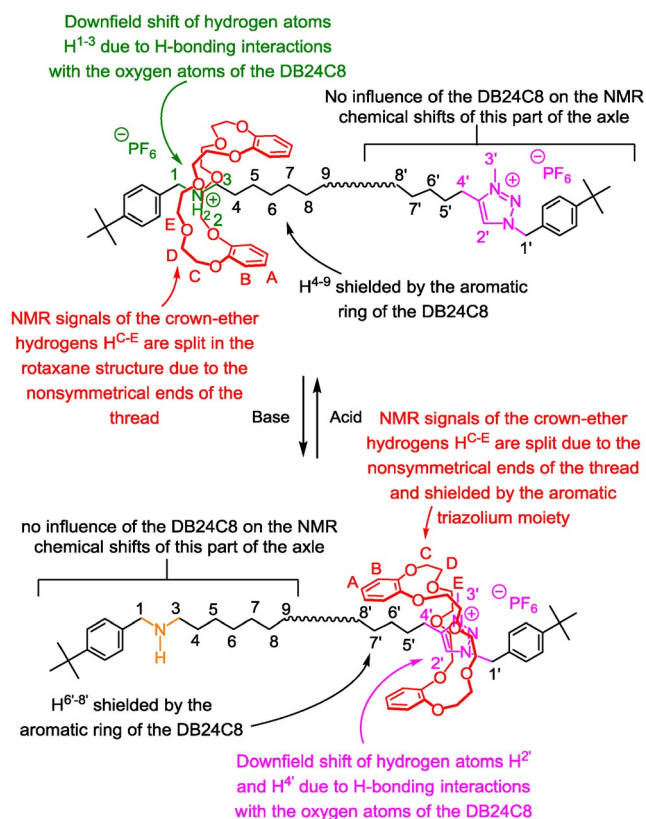


Figure 3. Usual main influence of the DB24C8 on the ^1H NMR chemical shifts of the ammonium and *N*-methyltriazolium stations.

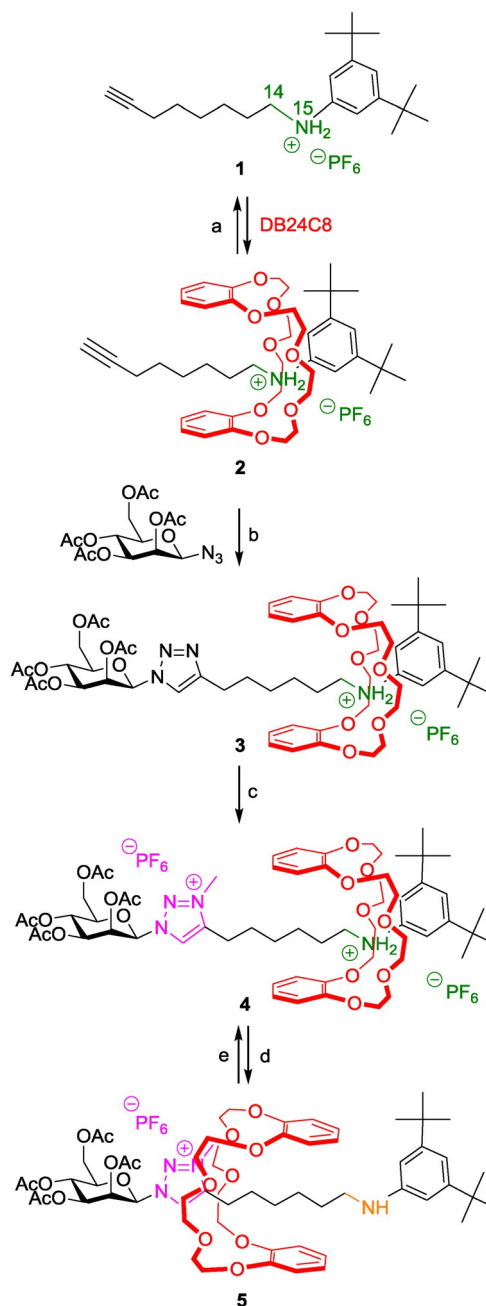
the rotaxane, due to hydrogen-bonding interactions with the oxygen atoms of the DB24C8. At the same time, hydrogens $\text{H}^{6-8'}$ are shifted upfield due to their localization in the shielding cavity of the aromatic rings of the DB24C8.

In other triazolium-containing rotaxanes, some variations of the exposed rules can be observed for hydrogens H^1 and $\text{H}^{4'}$ especially if the triazolium is located further to the bulky end. In that less constrained case, one may instead observe a shielding effect for these hydrogens.

3 Molecular Design in the [2]Rotaxane Series

3.1 A pH-sensitive mannosyl [2]rotaxane molecular machine

We reported the first example of a pH-sensitive [2]rotaxane molecular machine possessing a triazolium molecular station for the DB24C8 in 2008 (Scheme 1).^[13] Its synthesis appeared quite straightforward and efficient, and the molecular machinery easy to drive and characterize. The targeted DB24C8-based [2]rotaxane **4** consists of an encircled axle which contains an anilinium and a *N*-methyltriazolium molecular stations. Two bulky extremities have been chosen for the thread in order to prevent the macrocycle from any unthreading. One extremity is a mannose derivative, whereas the other one is a di-*tert*-butyl anilinium moiety. The anilinium was chosen as the pH-dependent station and as the template to drive the rotaxane formation. The triazolium serves as the second station



Scheme 1. Preparation of a mannosyl pH-sensitive [2]rotaxane molecular machine. *Reagents and conditions:* a) DB24C8 (2 equiv), CH_2Cl_2 , RT; b) 2,6-lutidine, $\text{Cu}(\text{MeCN})_4\text{PF}_6$, CH_2Cl_2 , RT, 24 h, 72%; c) 1) CH_3I , 2) NH_4PF_6 , quantitative; d) DIEA; e) 1) $\text{HCl}/\text{Et}_2\text{O}$, 2) NH_4PF_6 .

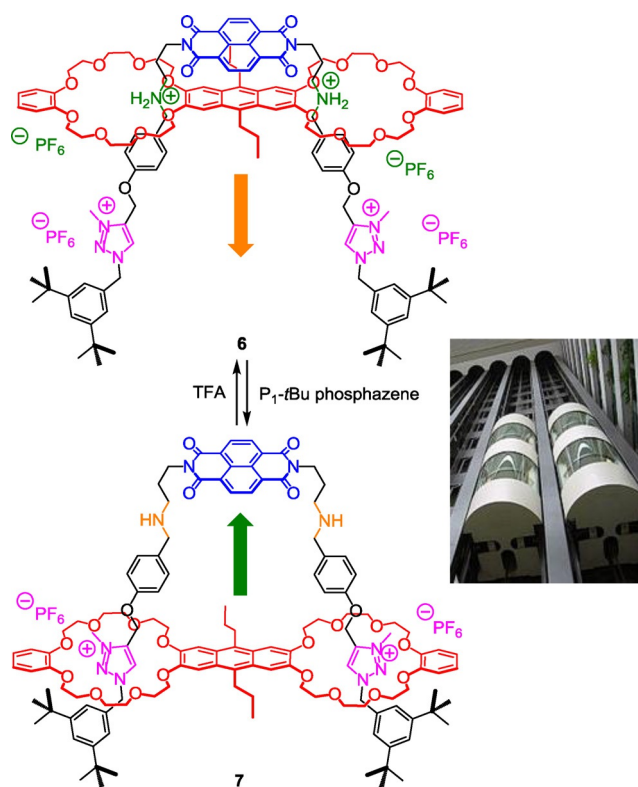
of much poorer affinity for the DB24C8. The preparation of the [2]rotaxane molecular architecture was achieved according to the threading and end-capping strategy described in Section 2.1. The anilinium alkyne **1** was able to thread quite instantaneously through the DB24C8 because of the hydrogen-bonding and ion-dipole interactions between the oxygen atoms of the crown ether and both the ammonium charge and the hydrogen atoms H^{14} and H^{15} . The obtained semi-rotaxane **2** was then capped at its alkyne extremity using CuAAC click chemistry with the mannosyl azide derivative. The subse-

quent methylation of the triazole revealed the triazolium station and afforded the pH-sensitive [2]rotaxane molecular machine **4**. After deprotonation of the anilinium station, the DB24C8 has no reason to sit anymore over the aniline because it has no more interaction with it. Therefore, the DB24C8 moves toward the *N*-methyltriazolium and interacts with it. Protonation of **5** inverts the translational process of the DB24C8 toward the anilinium.

This system appeared very appealing, first because of its quite easy synthetic access, second because of the efficiency of the molecular machinery, and third because of the easy characterization of the molecular machinery by ^1H NMR. Numerous other molecules based on this system have been reported since then. As a closely related example, Chen et al. proposed in 2010 the synthesis of a [3]rotaxane molecular machine, in which two triptycene derivatives are linked together thanks to two DB24C8 spacers, the latter being each threaded by two distinct axes.^[14] Other aesthetic examples are highlighted below.

3.2 A pH-sensitive molecular elevator

Using a similar approach (i.e. CuAAC click reaction followed by methylation of the triazole) although with different precursors, Liu et al. reported in 2013 a double-leg donor-acceptor molecular elevator where the distance between two aromatic platforms could be controlled thanks to the translational motion of the macrocycles along a bis-threaded axle (Scheme 2).^[15]



Scheme 2. Actuation of a triazolium-containing double-leg molecular elevator.

At the protonated state **6**, the bis-crown ether resides around the ammonium molecular stations where it interacts through hydrogen-bonding, ion-dipole, and additional charge-transfer interactions. Indeed, in this peculiar co-conformation, in addition to the already discussed weak interactions, and as already reported with a simpler example devoid of triazolium units,^[16] the electron-deficient naphthalenediimide interacts with the electron-rich anthracene moiety through a charge transfer. Noteworthy, the fluorescence of the interlocked molecule **6** is quenched by the proximity of these two moieties. Furthermore, in addition to hydrogen-bonding and ion-dipole interactions, this charge transfer improves the prior semi-rotaxane formation, so that the latter can be efficiently prepared using only a stoichiometric amount of each element to be assembled. The deprotonation of rotaxane **6** necessitated a strong base derived from phosphazene, because of the higher pK_a of the ammonium group with respect to anilinium. The action of the base induces the shuttling of the bis-crown ether toward the triazolium stations, moving the anthracene and the naphthalenediimide away from each other, therefore restoring at the same time the fluorescence intensity of the anthracene. In summary, the molecular machine acts as a molecular “elevator”^[17] in which the distance between the anthracene and the naphthalenediimide moieties platform can be adjusted precisely. Authors suggest using the switchable cavity of this molecule in order to bind a guest molecule between the two rigs under the assistance of the two platforms.

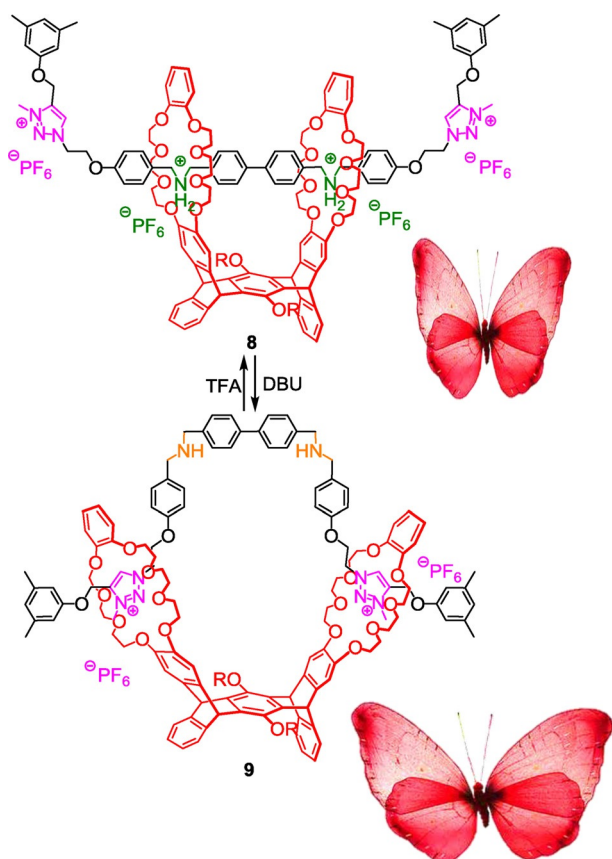
3.3 Mimicking the flipping motion of butterfly wings

Another example of “chemical design” was reported by Chen et al. in 2014.^[18] Inspired by smart bionic machines, they proposed the synthesis, through CuAAC click reaction followed by methylation of the triazoles, of a 2rotaxane containing a pentiptycene-derived bis(crown ether), two ammonium stations, and two *N*-methyltriazolium stations. The pH-sensitive molecular machinery that is inherent to this doubly-threaded molecule mimicks the flapping motion of the wings of a butterfly (Scheme 3).

Deprotonation of the ammonium-containing rotaxane **8** forces the DB24C8 rings to interact with the triazolium stations. Since the pure bending of the molecular axle seems impossible to reach without any torsion of the pentiptycene derivative, this being probably due to steric hindrance and constraints, the authors proposed that the twist of the host is necessary for the DB24C8 to reside around the triazolium units. The process can be reversed by simply adding trifluoroacetic acid (TFA). This sequential pH-dependent moving apart or approaching from each other of the two linked DB24C8 moieties can closely approximate the wing-flapping motion of a butterfly, where the DB24C8 are the wings.

3.4 A pH-sensitive molecular pulley

The knowledge acquired by Chen et al. on triptycene-derived crown ether hosts^[19] gave them the opportunity to mimic a molecular pulley (Scheme 4).^[20] In their ingenious example,



Scheme 3. Mimicking the wing-flapping motion of a butterfly using a triazolium-containing pentiptycene bis(crown ether)-based 2rotaxane.

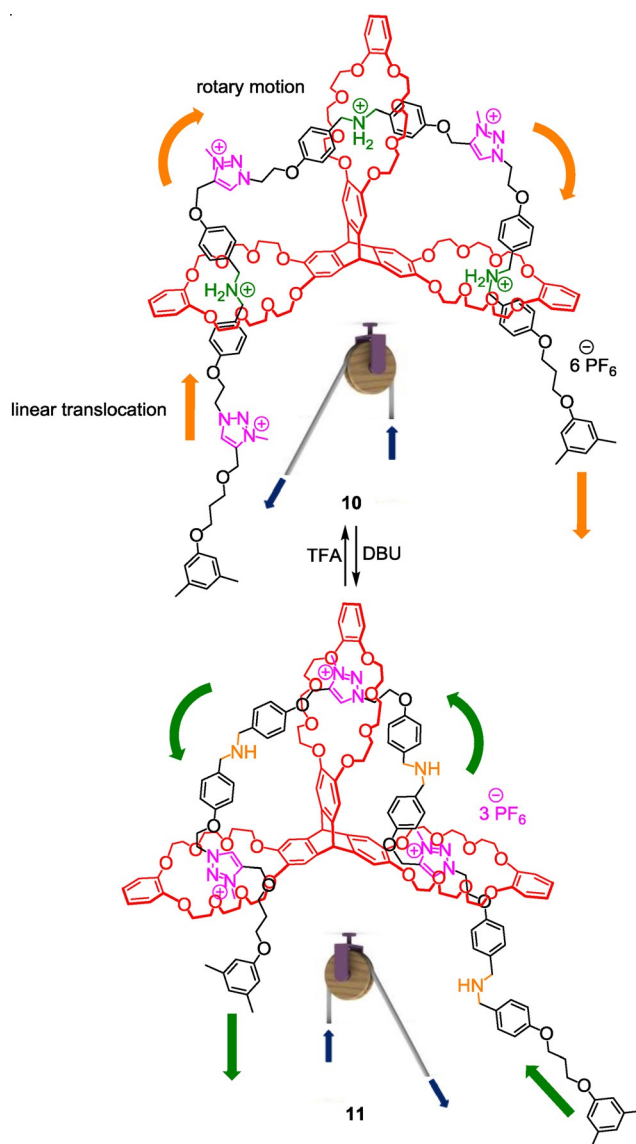
the triptycene moiety serves as a D_{3h} symmetrical scaffold that bears three crown ethers. The three DB24C8 are all threaded by one single molecular axle that contains alternatively three ammonium and three *N*-methyltriazolium stations. In **10**, the three DB24C8 derivatives encircle the ammonium stations. Deprotonation of the ammoniums triggers the forward shuttle of the cable-like thread until each DB24C8 interacts with the triazolium units. The reverse motion is possible through reprotonation of the amines using TFA. Due to the tridimensional geometry of the triptycene, the axle undergoes both translation and rotation motions, like in bi-stable [2]rotaxane and [2]catenane. The authors suggest the future incorporation of loads on either the molecular cable or on the wheel component, in analogy to the operation of a pulley.

Examples of triazolium-containing rotaxane molecular machines that have found utility as switchable organic catalysts or as switchable fluorophores are highlighted in the following section.

4 Utilization of Triazolium-Containing Molecular Machines

4.1 As switchable pH-sensitive [2]rotaxane catalysts

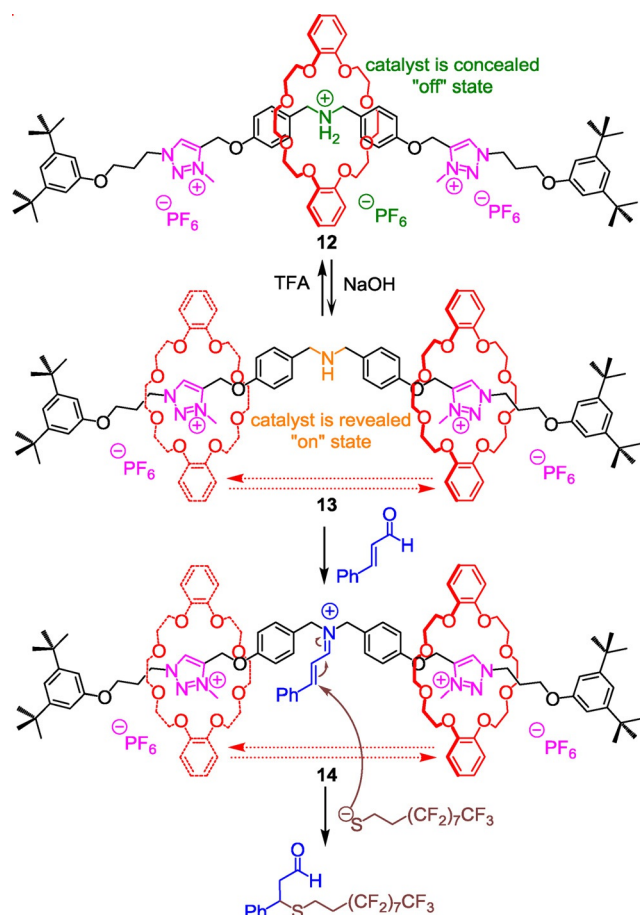
Inspired by how nature can control enzymatic syntheses through trigger-induced effects, Leigh et al. successfully used



Scheme 4. Mimicking a molecular pulley in a triply interlocked triazolium-containing rotaxane molecular machine.

the crown ether- and triazolium-ammonium-based molecular machinery in order to prepare appealing switchable organic rotaxane catalysts.^[21] The aim was to turn “on” or “off” the activity of a [2]rotaxane catalyst through the concealing or the revealing of the catalytic motif, depending on the pH-sensitive localization of a macrocycle along a thread. The first switchable catalyst **13** they proposed consists of a [2]rotaxane molecular machine that contains one ammonium and two *N*-methyltriazolium stations for the DB24C8 (Scheme 5).^[22]

The ammonium moiety was chosen because of its ability, in a non-interlocked molecule, to catalyze Michael-type reactions either at the protonated or the deprotonated state. After having prepared the [2]rotaxane molecular machine **12** according to the two-step sequential chemical route highlighted in section 2.1, the authors investigated the possibility to reveal or conceal the secondary amino function via the shuttling of the DB24C8 along the threaded molecular axle. At the protonated

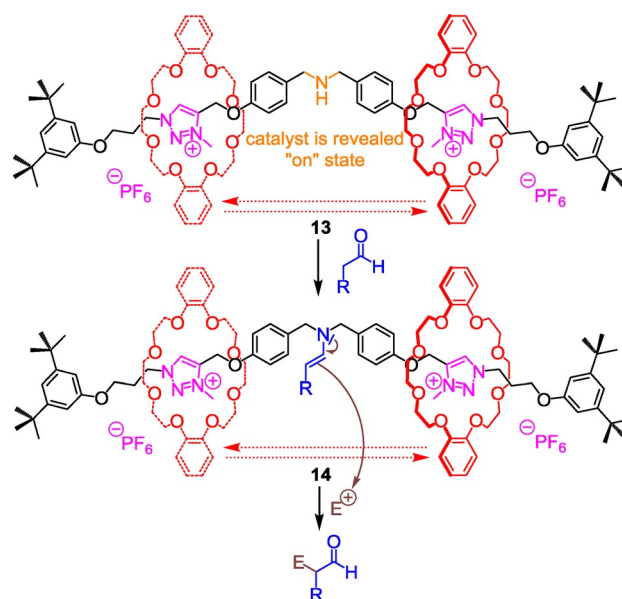


Scheme 5. Switching “on” or “off” a triazolium-containing [2]rotaxane catalyst for Michael-type addition.

state, the ammonium is encircled by the crown ether, which results in the concealing of the ammonium catalyst, hence its deactivation. At this stage, mixing 5% of the [2]rotaxane **12** with *trans*-cinnamaldehyde and an aliphatic thiol did not yield to any Michael-type adduct, even after five days of reaction. However, washing the previous mixture with a 1 M aqueous solution of sodium hydroxide induces the motion of the DB24C8 toward the triazolium stations (rotaxane **13**). In this new co-conformation, the amino moiety is revealed, bringing about the “on” state of the catalyst, hence the iminium catalysis.^[23] In this basic condition, the Michael addition could be carried out in a 66% yield after only one hour!

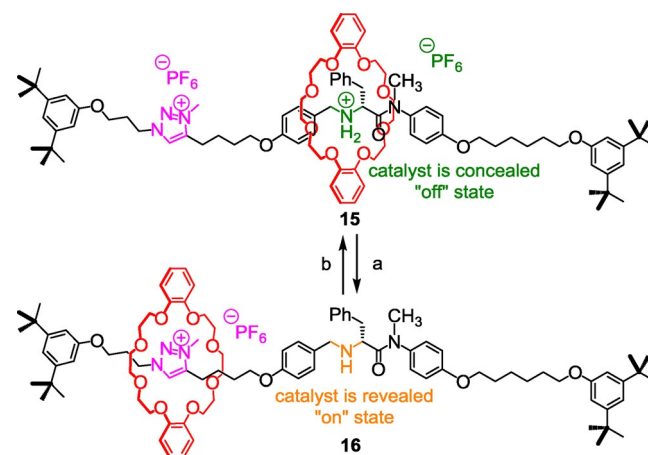
By continuing their efforts, the same authors successfully extended the catalyzed Michael-type reaction of the same thiol compound to different α,β -unsaturated carbonyl compounds.^[24] They also reported the Michael addition using 1,3-dicarbonyl compounds as the nucleophilic agent. Enamine catalysis^[25] was also investigated using the same switchable rotaxane catalyst for nucleophilic substitution and nucleophilic addition reactions in the α position of an aldehyde (Scheme 6).

In this case, both the α -chlorination of saturated aldehydes using *N*-chlorosuccinimide and the Michael addition to vinyl bis-sulfone were achieved using **13** as the catalyst (20 mol%) with respective ranges of conversion of 40–61% and 23–70%



Scheme 6. Utilization of a switchable triazolium-containing rotaxane catalyst for the introduction of an electrophile at the α position of an aldehyde.

depending on the substitution of the aldehyde. No such reactions were observed in the absence of **13** or in the presence of the protonated rotaxane **12**. One drawback still remains in the impossibility to extend these two catalyzed reactions to cyclic or acyclic ketones. The extension to tandem iminium–enamine catalyzed reaction was achieved in a 50% overall conversion in that peculiar case, the *trans*-cinnamaldehyde was first subjected to iminium catalysis allowing the introduction of a thiol in the β position of the aldehyde. This led to an enamine which subsequently reacted with vinyl bis-sulfone in the α position of the aldehyde. Eventually, the efficacy of the switchable rotaxane catalyst **13** was successfully tested in a Diels–Alder reaction between a dialdehyde and a dienophile.



Scheme 7. A chiral switchable triazolium-containing rotaxane catalyst for stereoselective conjugated addition. Reagents and conditions: a) NaOH; b) 1) HCl, 2) KPF₆.

The same authors recently extended their work by synthesizing the chiral pH-sensitive rotaxane molecular switch **15/16**, which proved to catalyze conjugated additions, only at the deprotonated state, through the iminium ion activation and notably with high stereoselectivity (Scheme 7).^[26]

The principle is the same as already discussed, that is to say based on the revealing of the amine moiety through the shuttling of the DB24C8 in a basic medium. Although the literature mentioned that conjugated addition could be generally carried out efficiently with high stereoselectivity using cyclic amine as catalyst, here, it is interesting to note that it could be realized with an acyclic secondary amine derived from a (*R*)-phenylalanine residue. Enantiomeric excess as high as 88% was obtained. This very attractive contribution now opens new avenues toward the selective sequential reactions of a mixture of a wide range of compounds using successive activation of switchable catalysts.

4.2 As switchable pH-sensitive fluorescent rotaxanes

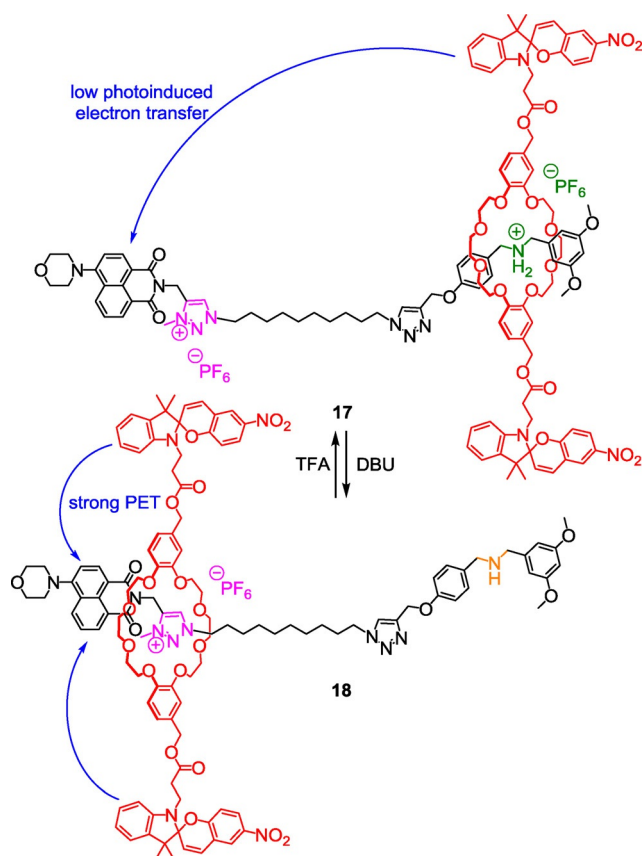
4.2.1 In triazolium-containing [2]rotaxanes

The possibility for a macrocycle to be localized at a different place along a thread gives the opportunity to tune the fluorescence of an interlocked molecule depending on a stimulus. Qu et al. synthesized a switchable spiropyran-containing [2]rotaxane in which the threaded axle is end-capped by a fluorescent 4-morpholin-naphthalimide.^[27] The targeted pH-sensitive molecular machine **17** contains a DB24C8 macrocycle derivative and two ammonium and triazolium molecular stations. The DB24C8 macrocycle is substituted by two nitroaromatic spiropyran moieties that can tune the fluorescence of the 4-morpholin-naphthalimide stopper depending on their distance (Scheme 8).

Shuttling the macrocycle toward the triazolium recognition site was achieved through the addition of 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) on rotaxane **17**. Concomitantly, the fluorescent emission intensity decreased by 45% in **18**, with respect to **17**. This can be ascribed to the higher proximity between the fluorophore and the two photochrome units in the deprotonated rotaxane **18**. In this state, the quenching of the 4-morpholin-naphthalimide fluorophore, via the photo-induced transfer of one electron of the highest occupied molecular orbital (HOMO) of the spiropyran moieties to the vacancy of the HOMO of the excited 4-morpholin-naphthalimide, becomes more effective.

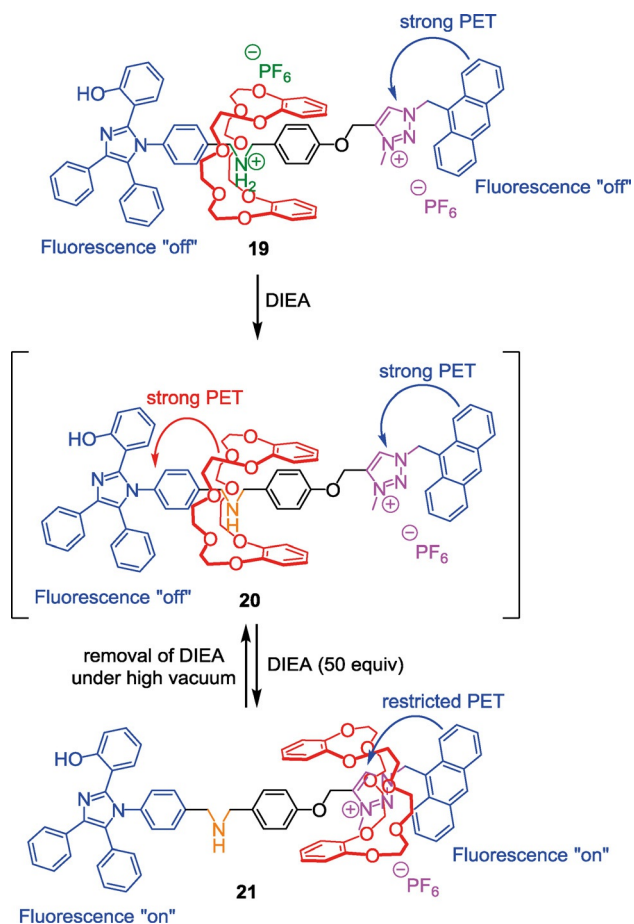
A somewhat similar example of controlled fluorescence in a [2]rotaxane molecular machine was published by Li et al. in 2011.^[28] They synthesized a tight rotaxane based on the ammonium/triazolium/DB24C8 system which operates in a basic environment. Here, the threaded axle holds at each extremity two chromophores as bulky stoppers (Scheme 9).

Surprisingly, the molecular machine **19** operated slightly differently with respect to the other reported examples. On one hand, deprotonation using the weak base *N,N*-diisopropylethylamine (DIEA) did not lead to the immediate displacement of the crown ether around the triazolium station in **20**, unless



Scheme 8. A triazolium-containing [2]rotaxane with fluorescent output.

a large amount of 50 equivalents of DIEA was used. On the other hand, removal of DIEA under high vacuum leads to the reversible motion of the DB24C8 around the NH site. The authors explained this result by the fact that anthracene is a bulky moiety which sterically disturbs the solvation of the *N*-methyltriazolium cation by acetonitrile. This results in a tighter ion-pair between the triazolium cation and the hexafluorophosphate anion that prevents the DB24C8 to interact optimally with the triazolium station. Adding the stronger Lewis base DIEA causes a better solvation of the triazolium cation, inducing the dissociation of the ion-pair, hence promoting the interaction between the DB24C8 and the triazolium recognition site (compound **21**). Another reasonable explanation is that a small amount of DIEA is just not sufficient to deprotonate the ammonium quantitatively. Removal of the DIEA under vacuum increases the concentration of diisopropylethylammonium, thus leading to the reprotonation of the rotaxane. In any cases, what is interesting here is that the fluorescence of the two extremities can be switched on or off using a pH stimulus. In the protonated rotaxane **19**, the fluorescence of the anthracenyl moiety is quenched because of the strong photoinduced electron transfer of the anthracene to the triazolium moiety. At the same time, the close distance between the DB24C8 and the hydroxyl-substituted tetraphenylimidazole (HPI) results in the weak fluorescence emission of the HPI moiety. At the deprotonated state, the DB24C8 moves toward the triazolium station. The concealing of the triazolium by the crown ether



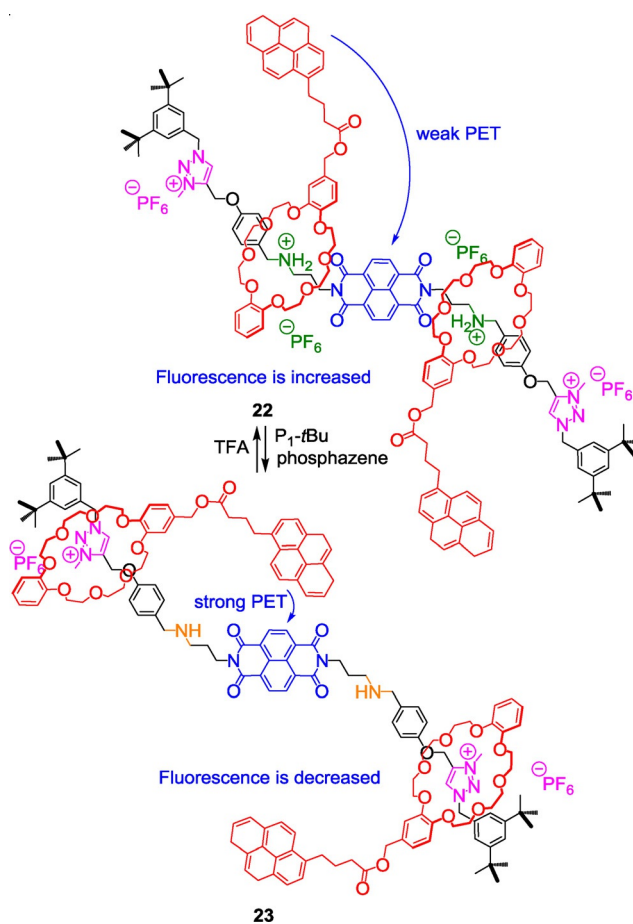
Scheme 9. A triazolium-containing [2]rotaxane with two fluorescent outputs.

now prevents from any photoinduced electron transfer (PET) between the anthracenyl group and the triazolium. Therefore, fluorescence of the anthracene is restored. The same increase in fluorescence emission is observed for the HPI moiety because of the higher distance from the DB24C8 which now sits over the triazolium station.

4.2.2 In triazolium-containing [3]rotaxanes

Another example of tunable fluorescence in a more sophisticated [3]rotaxane molecular machine was reported by Liu et al.^[29] The internal motion of the DB24C8 along the thread is once again based on its different affinities for the ammonium and triazolium stations as anteriorly explained (Scheme 10).

An electron-deficient naphthalenediimide (NDI) is used as a linker between two branched [2]rotaxanes that both contain an ammonium and a triazolium station. The ammoniums are located close to the NDI core, whereas triazolium units lie at each extremity of the [3]rotaxane. Two DB24C8 initially surround the ammoniums of the thread at the protonated state **22**. They both bear an electron-rich pyrene moiety that can be involved in a charge-transfer complex with the NDI moiety. The obtained results indicate that the fluorescence intensity of **22** decreases upon addition of the phosphazene strong base, whereas addition of TFA on **23** increases the fluo-

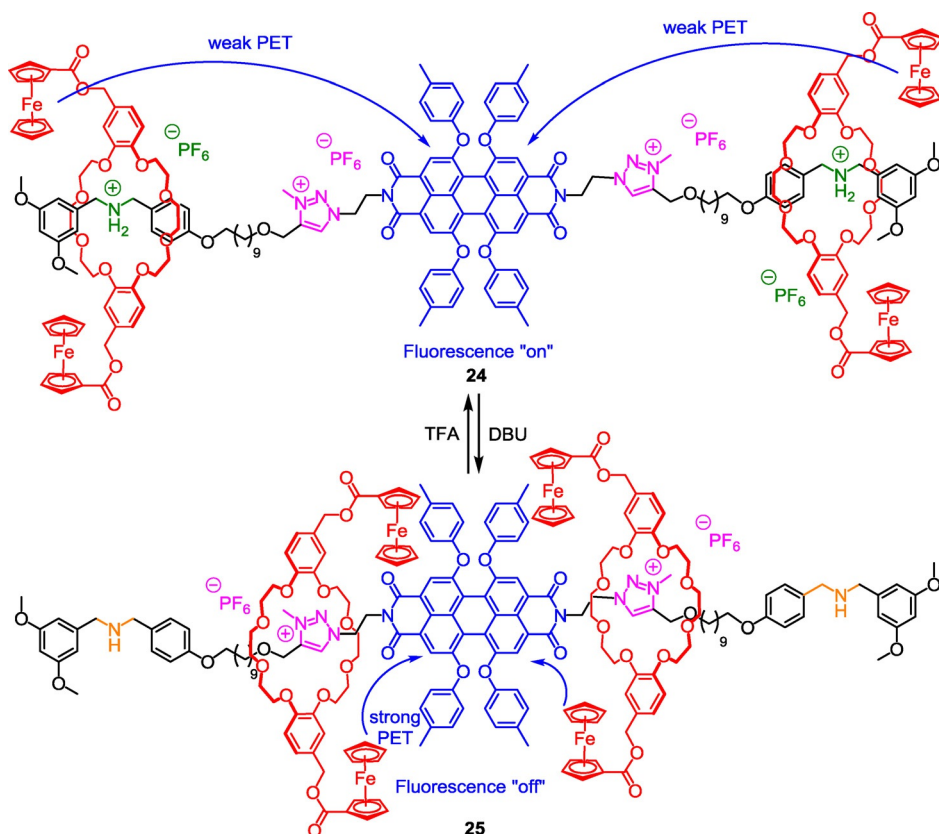


Scheme 10. A triazolium-containing [3]rotaxane with fluorescent output.

rescence. This can be explained by an internal charge-transfer (ICT) process which depends on the spatial distance between NDI and pyrene moieties. Further experiments showed that the measured luminescence is only due in this case to the free pyrene. The authors used molecular energy minimization to evaluate the distance between the electron-rich and the electron-deficient moieties. In the protonated system **22**, the distance between the pyrenes and the NDI appears larger than in **23**, although the DB24C8 are located much closer to the NDI. This corroborates the π - π stacking interactions observed in **23** by ^1H NMR and by UV/Vis absorption spectroscopies between the NDI and one pyrene unit.

Another fluorescent [3]rotaxane was published by Qu et al. in 2015. The symmetrical bi-stable bis-branched [3]rotaxane molecular machine contains a perylene bisimide (PBI) chromophore at the middle of the threaded axle and ferrocenyl units on the DB24C8.^[30] The molecular machinery is once again based on DB24C8 derivative and ammonium/triazolium molecular stations, so that two distinct co-conformational states can be easily reached upon variation of pH (Scheme 11).

However, contrary to the previous example, the ammonium station is located at the two extremities of the thread, while triazoliums are next to the PBI core. The remarkable pH-dependent tuning of the visual fluorescence of the PBI was achieved through its distance-dependent quenching by the ferrocenyl



Scheme 11. A triazolium-containing [3]rotaxane with fluorescent output.

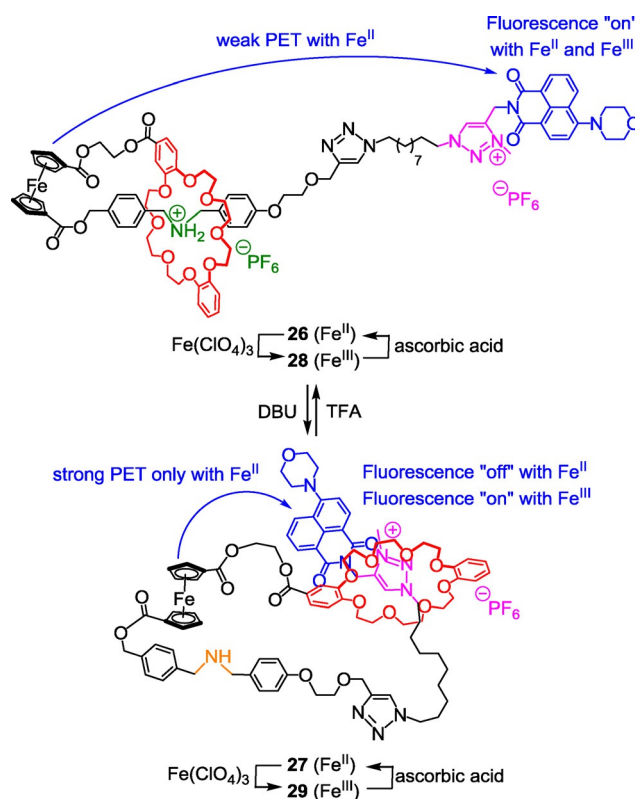
units. The change of this photophysical property was possible thanks to both the molecular machinery and the photoinduced electron transfer from the electron-rich ferrocene donors to the electron-deficient PBI fluorophore acceptor. At the protonated state (compound **24**), the two macrocycles reside around the ammonium stations: in this co-conformation, the ferrocenyl moieties are located too far away from the PBI to allow any efficient PET. Hence, the fluorescence of the PBI is "on". However, deprotonation of the ammoniums using 3 equiv of DBU causes the shuttling of the two macrocycles along the thread until the triazolium sites, which are close to the fluorescent PBI moiety. It results from these motions that ferrocenyl groups are now near the PBI core of the [3]rotaxane molecular machine, hence inducing a strong PET which is responsible for the decrease (80%) of the emission intensity at 620 nm with respect to the original spectrum. Interestingly, this fluorescence quenching could be visualized with the naked eye. The authors also demonstrated that protonation of **25** using 6 equiv of TFA restored the fluorescence intensity. This cycle (protonation/deprotonation) could be repeated several times without noticing any degradation.

4.2.3 In triazolium-containing [1]rotaxanes

In 2013, Qu et al. published a fluorescence switch in the [1]rotaxane series.^[31] The principle is quite similar to the [3]rotaxane described above, although the motion of the molecular ma-

chine relies on the tightening or the loosening of a lasso (Scheme 12).

The ferrocenyl unit, which is part of the lasso's loop, can be localized more or less close to the fluorescent stopped molecular tail depending on the molecular machinery, using the difference of affinity of the DB24C8 for the ammonium and the triazolium station. In the protonated state, the DB24C8 has more affinity for the ammonium moiety, therefore forcing the lasso to adopt a tightened co-conformation. At this point, the ferrocenyl unit remains too far away from the morpholin-naphthalimide to quench its fluorescence, the latter being monitored by the naked eye. Deprotonation of the ammonium results in the shuttling of the DB24C8 around the triazolium, hence loosening the lasso. In this specific co-conformation, the ferrocenyl unit is much closer to the tail, now authoriz-



Scheme 12. A triazolium-containing [1]rotaxane with fluorescent output.

ing a strong PET from the electron-rich ferrocene to the electron-deficient naphthalimide. This PET affects the fluorescence emission intensity at 521 nm in **27** which decreases by 80% with respect to the protonated tightened lasso **26**. Using theoretical calculations, the authors managed to get data relative to the distance separating the iron atom and the nitrogen atom of the naphthalimide. The results obtained are in total agreement with the fluorescence observation. Indeed, in **26**, the calculated distance between the two atoms is 35.66 Å, whereas in **27** it decreases to only 12.87 Å, therefore allowing a more efficient PET between the ferrocene and the naphthalimide, thus the quenching of the fluorescence.

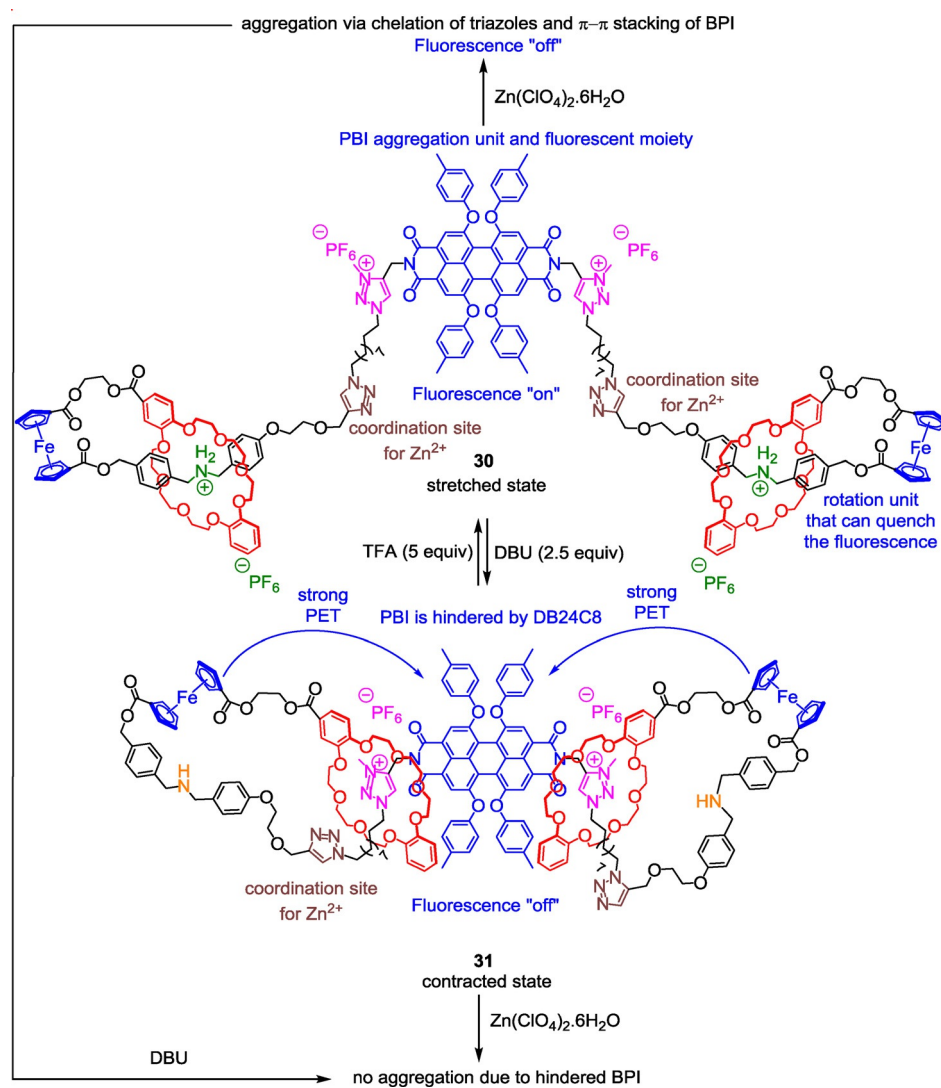
The authors then explored the possible dual pH- and redox-dependent output signal after oxidizing/reducing the ferrocene unit, using respectively $\text{Fe}(\text{ClO}_4)_3$ or ascorbic acid. In rotaxane **29**, after that the ferrocene unit of rotaxane **27** is oxidized, the fluorescence is dramatically restored to its original level. Furthermore, the oxidation of the protonated fluorescent rotaxane **26** leads to compound **28** without any significant change of emission intensity.

These two last results can be ascribed to the fact that oxidation of the ferrocene units obviously affects its electron-donating ability, thus not allowing efficient PET in this case, whatever the distance between the considered moieties. Reducing the ferrocene unit with ascorbic acid gave rise to the controllable pH-switchable fluorescence output signal (rotaxanes **26/27**). In summary, this molecular machine proved to operate upon a pH variation through the motion of the macrocycle along the threaded axle. The fluorescent output signal was found to be both pH- and redox-dependent. Quenching the fluorescence could be achieved using a pH stimulus (deprotonation)

through actuation of the machinery, whereas restoring the original level of fluorescence was possible either by oxidation or by protonation. The authors present their system as an INHIBIT logic gate combining a "silent" and an "active" signal output which could be of real interest for the conception of logic circuits with memories or sequential functions. The same group extended this work to the synthesis and the study of two pH-sensitive star-shaped [1](*n*)-ro-

taxane molecular machines with tri and tetrabranch cores that are linked to triazolium stations.^[32] Using the features of the two previously reported examples, **24** and **26**, Qu et al. recently published an appealing bis-branched [1]rotaxane that can combine dual-mode molecular motions and tunable nanostructural morphologies (Scheme 13).^[33]

The molecular machine **30** is based on the difference in affinity of ammonium and triazolium stations for the DB24C8 macrocycles. Moreover, it combines two ferrocenyl units that are used to connect the thread and the macrocycle for the bis[1]rotaxane architecture, on one hand, and for its electron-rich donor ability on the other hand. An electron-deficient PBI fluorophore lies in the core of the thread and possesses aggregation potential that can be tuned. 1,2,3-triazoles are located between ammonium and triazolium molecular stations and serve as coordinating sites for Zn^{2+} in order to tune the aggregation degree of the PBI moiety. As for the previously described examples, the DB24C8 can move along the thread upon a pH stimulus, here triggering the contraction and the



Scheme 13. A triazolium-containing bis-branched [1]rotaxane with reversible fluorescence, stretching/contraction motion, and reversible aggregation.

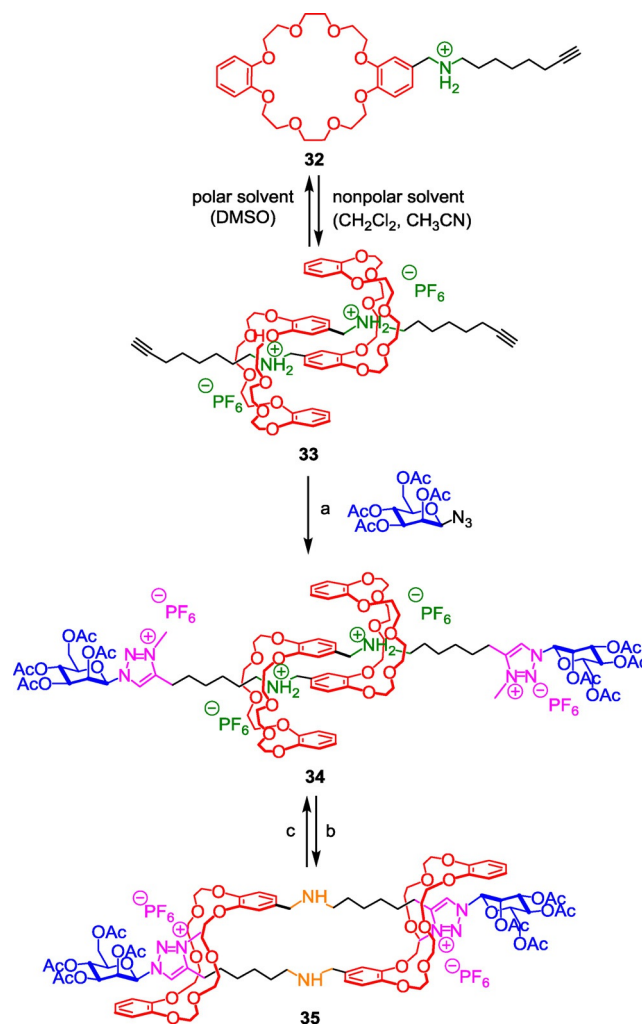
stretching of the bis[1]rotaxane **30/31** because of its single molecular [1]rotaxane architecture. Interestingly, it was shown that the translational movement of the DB24C8 was accompanied by a simultaneous rotational motion of the ferrocene units, which makes this machine acting as a dual-mode movement type. The fluorescence of the PBI could be tuned either upon change in solvent polarity or variation in pH. Indeed, increasing the proportion of the apolar methylcyclohexane in a solution of **30** in acetone results in the dramatic decrease of the fluorescence, due to π - π stacking aggregation of the PBI. On the contrary, in **31**, very little π - π stacking aggregation of the PBI was noticed. On the other hand, the fluorescence can be also controlled by the molecular machinery, via the distance-dependent photoinduced electron transfer of the ferrocene moieties to the PBI. Upon deprotonation, the fluorescence of PBI is quenched because the ferrocene units are located closer to the PBI. At the protonated state, it is also interesting to notice that addition of Zn^{2+} could induce the aggregation of the molecule through the cross-linking chelation of the cationic transition metal with the neutral triazole moieties. This chelation phenomenon helps the PBI moieties to self-interact through π - π stacking. Addition of the strong base DBU decreased the aggregation because of the shuttling of the DB24C8 around the triazolium, thus hampering π - π stacking due to steric hindrance. Using theoretical calculations, the authors estimated a percentage of molecular length change of 28.3% between the two conformational states. This value is very close to the value observed in human muscles. Although the contraction and the stretching of the molecule can be compared with a muscle, the first molecular muscle reported by Sauvage et al., which was based on metal-ligand coordination, holds a slightly different doubly interlocked molecular architecture.^[34] Examples of such a topology, based on the system of molecular machinery that is highlighted in this review, are given in the following section.

5 Triazolium-Containing Molecular Muscles and Their Use as Building blocks in Polymers

5.1 Synthesis and study of a pH-sensitive molecular muscle

"Molecular muscle"^[35] is a generic term used for a molecule designed to adopt a contracted or a stretched co-conformation in response to a stimulus, in a similar manner to human muscles. Inspired by the pioneering work of Jean-Pierre Sauvage^[36] on doubly interlocked molecular architectures and by those of Fraser Stoddart on DB24C8-based [c2]daisy chain,^[37] we reported in 2008 the first pH-sensitive molecular muscle. A few other examples based on other systems have been reported since then.^[38] We actually extended the ammonium-triazolium system of recognition sites for DB24C8 that we previously published^[13] in the [2]rotaxane series to the preparation of a dimannosyl pH-sensitive molecular muscle (Scheme 14).^[39]

The so-called hermaphrodite compound **32** was synthesized and dissolved in several different solvents which differ in polarity. Whereas **32** remains unassembled in the more polar solvent DMSO, it self-assembles as a pseudo-rotaxane dimer in



Scheme 14. An ammonium/triazolium-containing pH-sensitive molecular muscle. *Reagents and conditions:* a) 1) mannosyl azide (pictured), $\text{Cu}(\text{CH}_3\text{CN})_4\text{PF}_6$, lutidine, 2) CH_3I then NH_4PF_6 ; b) NaOH (aq)/ CH_2Cl_2 ; c) 1) HCl (Et_2O , 2) NH_4PF_6 .

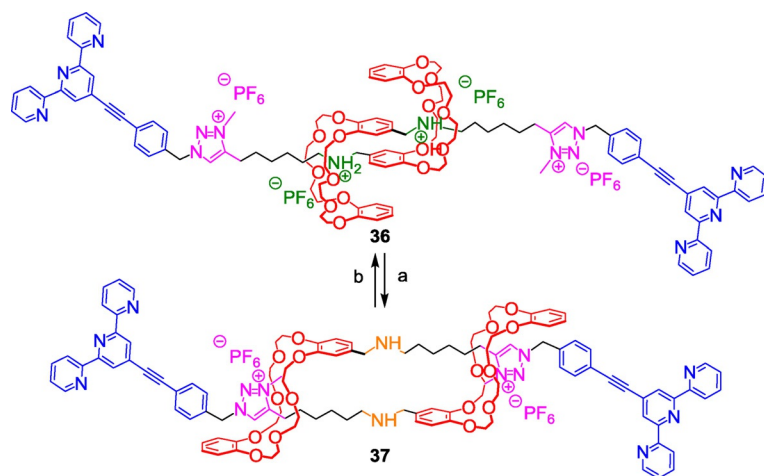
the less polar hydrogen-bond-promoting solvents like dichloromethane or acetonitrile. We pointed out the *meso* stereochemistry of the self-assembled [c2]daisy chain supramolecular arrangement **33**, which is probably due to both steric hindrance and dipole-moment repulsion between the ionic chains linked to the threaded crown ether. The subsequent two-step sequence, 1) CuAAC click reaction and 2) methylation of the triazole, locked the interwoven supramolecular structure by capping the threads with mannosyl stopping groups and also revealed the triazolium stations for the DB24C8. The isolated rotaxane dimer **34** was then studied in order to evidence the actuation of the molecular muscle. As usually observed with simpler interlocked architecture, at the protonated state **34**, the molecule remains in an extended co-conformation because the two DB24C8 prefer to interact with the best ammonium molecular stations. However, deprotonation of the ammonium moieties using sodium hydroxide quantitatively yields to the contracted molecular muscle **35** through the gliding motion of the two DB24C8 along the treads until they reach the triazoli-

um sites. The very efficient molecular machinery from the contracted to the stretched co-conformations matches with a maximum contraction ratio of 48%. The distance between the two anomeric carbons of the stoppering mannose was evaluated up to 23 and 34 Å, respectively. This initial work paved the way to applications in the domain of molecular-muscle-containing supramolecular polymers.

5.2 Use of molecular muscles as building blocks for the synthesis of supramolecular polymers

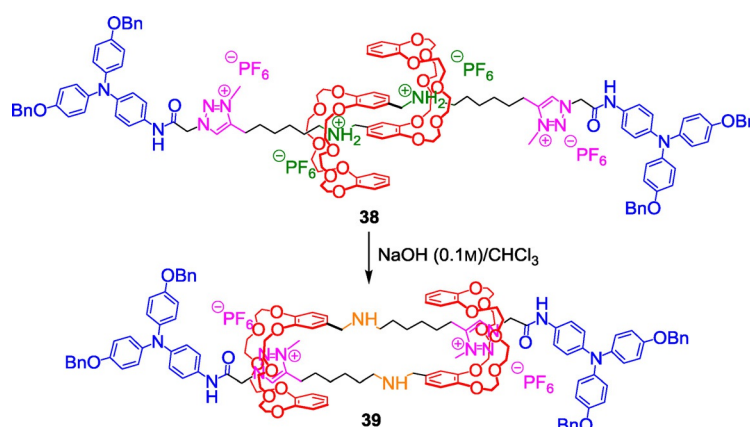
The same specific system of molecular muscle, but with different stoppers, was employed later by Buhler, Giuseppone, et al. with the aim to construct a supramolecular polymer.^[40] To this end, they used terpyridine moieties both to lock the interwoven structure and to further have the possibility to assemble the molecular muscle building blocks through metal coordination (Scheme 15).

The terpyridine moiety was already known to coordinate strongly Fe^{II}, Zn^{II},^[41] or other metals such as Ru^{II} or Os^{II}.^[42] Moreover, Che et al., as well as Schubert et al., previously used terpyridines to prepare supramolecular polymers.^[43] In the present case, the molecular muscle building block **36** was prepared according to Coutrot's strategy and acted upon pH variation like the previously published compound **34**. Adding either Fe^{II} or Zn^{II} to **36** triggered the polymerization thanks to the formation of stable octahedral 2:1 complexes. A longer supramolecular polymer was obtained with Fe^{II} because of its better coordination for terpyridine than Zn^{II}. Of particular interest is the length of the polymer that was obtained this way. No less than 3000 units have been assembled through coordination, which is a real achievement with respect to already published molecular-muscle-containing polymers that usually do not contain more than 22 units.^[44] The change of size of the metallosupramolecular polymer upon variation in pH was evaluated by small-angle neutron scattering. The contour length of the protonated polymer is long (15.9 μm) with respect to the



Scheme 15. An ammonium/triazolium-containing pH-sensitive muscle-like monomer for the synthesis of a metallo-supramolecular polymer. Reagents and conditions: a) NaOH (aq)/CH₂Cl₂; b) 1) HCl/Et₂O, 2) NH₄PF₆.

deprotonated polymer (9.4 μm), while the decrease of the linear density is observed simultaneously. An amplification of almost four orders of magnitude in the mechanical output is observed in the polymer in comparison with the monomer. In other words, the length variation of the polymer between the two co-conformations is about 6.5 μm, while that of the monomer is only around 1 nm. With this work, a real step toward the macroscopic world has been done, even though these polymers should be bundled together in order to better mimic myofibrils of sarcomers. Very recent improvements in this way have been achieved by the same group. They reported the possibility to induce the fabrication of micrometer-long supramolecular fibers from the molecular muscle building block **38** (Scheme 16).^[45]



Scheme 16. Triazolium-based building block for microfiber-induced formation.

The system works as an INHIBIT logic gate where both light and acidic medium trigger the formation of the supramolecular fiber. The Coutrot [c2]daisy chain **33** was functionalized this time with a triarylamine moiety that holds an amide functional group. The usual two-step chemical strategy was applied to lock the structure and to incorporate a triazolium station in each encircled thread. Importantly, light exposure in presence of chlorinated solvent was found to induce the self-assembly of the sole building block **38** through hydrogen bond, π - π stacking, and van der Waals interactions. The formation of bundled fibrils of several microns length was evidenced by NMR, dynamic light scattering, UV/Vis-near infrared (NIR) absorption, and transmission electron microscopy (TEM). In this case, the self-assembly is attributed to the formation and the stabilization of triarylamine radical cations, as well as to the necessary hydrogen bonding between the amide moieties. This latter point

is crucial, since no aggregation of the triarylamine is possible in the absence of amide moiety. Interestingly, the same result is observed if the amide is hindered by the DB24C8 through molecular machinery at basic pH. For this reason, no such supramolecular self-assembly of the deprotonated molecular muscle building block **39** could be observed.

In order to enhance the differentiation, hence the physical property, of the two co-conformational states of supramolecular polymers, the increase of both the rigidity and the length of the spacer which links the two molecular stations might be envisaged in the next future.

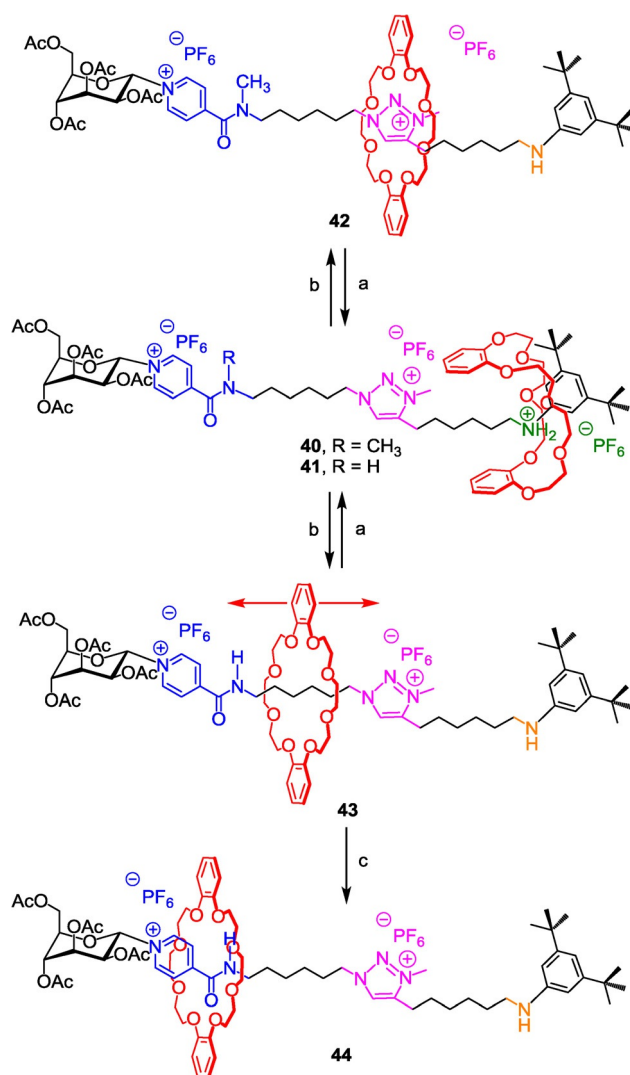
Other methodological studies focused on the relative affinity of the triazolium station for the DB24C8 in three-station-based rotaxanes.

6 Relative Affinity of the DB24C8 for the Triazolium Station With Respect to Pyridinium Amide Station in Three-Station-Based Rotaxane Molecular Machines

6.1 In three-station-based [2]rotaxane molecular machines

The synthesis of three-station-based rotaxanes containing an anilinium, a pyridinium amide, and *N*-methyltriazolium stations provided new interesting behavior of the system and gave insights into the relative affinity of the DB24C8 for the two different weak interaction sites.^[46] Each efficiently synthesized rotaxane **40** or **41** holds three different molecular sites of interaction for the DB24C8 (Scheme 17).

The only difference between these two components relies on the substitution of the amide group. At the protonated state, the behavior of the two rotaxanes **40** and **41** is identical because the substitution of the amide does not affect the localization of the DB24C8, which resides around the best anilinium station. However, the pH-sensitive molecular machines react very differently after the addition of diisopropylethylamine. Deprotonation of the anilinium induces distinct co-conformations in **42** and **43**, because of the quite different affinities of the mono- and the disubstituted pyridinium amide moieties for the macrocycle. In **42**, NMR studies proved that the DB24C8 is localized around the sole triazolium molecular station. It follows that the affinity of the DB24C8 is better for the *N*-methyltriazolium than for the disubstituted pyridinium amide. Deprotonation of rotaxane **41** resulted in a very different compartment of the molecular machine. In this case, the DB24C8 interacts with both the triazolium and the pyridinium amide molecular stations, and the two translational isomers are in fast exchange with respect to the NMR time scale. Extensive NMR studies were achieved to better characterize the translational Brownian motion along the thread between the two stations. At room temperature, in acetonitrile, the DB24C8 spends 62% of its time around the monosubstituted pyridinium amide station for only 38% around the triazolium. This clearly indicates that the monosubstituted pyridinium amide has a slightly better affinity for the DB24C8 than the triazolium. Decreasing the temperature triggers the gradual decrease of the proportion of the translational isomer in which the DB24C8 is localized around

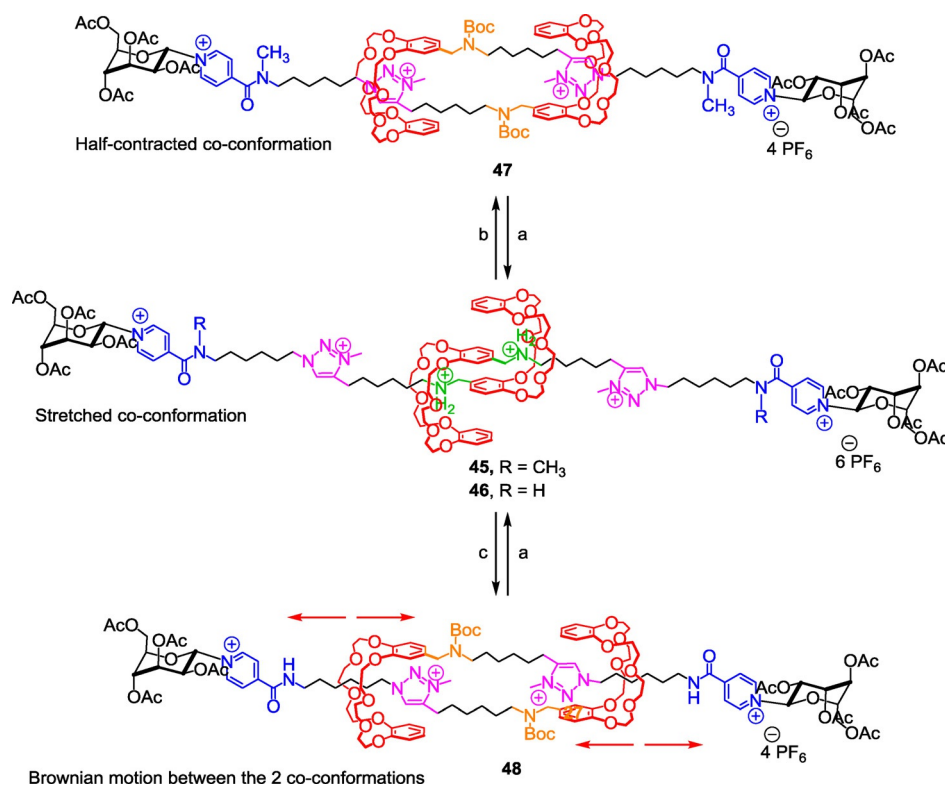


Scheme 17. *N*-methyltriazolium as a molecular station for the DB24C8 in a three-station-based [2]rotaxane. Reagents and conditions: a) 1) HCl/Et₂O, 2) NH₄PF₆, CH₂Cl₂/H₂O, 30 min; b) DIEA; c) CD₂Cl₂, 223 K.

the triazolium site. At 223 K, the DB24C8 does not spend any time around the triazolium (**44**). In summary, in the disubstituted pyridinium amide series, a bi-stable molecular machine is observed in which the DB24C8 can only shuttle between the anilinium and the triazolium. On contrary, in the monosubstituted pyridinium amide series, depending on pH, the machine adopts either a stable protonated state or undergoes a controlled Brownian movement of the DB24C8 between the two weakest stations. In this latter case, the molecular machinery appears both pH- and temperature-dependent, with the two stimuli (i.e. acidification and temperature decrease) leading to two opposite localizations of the macrocycle along the thread.

6.2 In a molecular muscle architecture

The previous work was also extended to the more sophisticated molecular muscle architecture and furnished complementary results on the behavior of the triazolium moiety (Scheme 18).^[47]



Scheme 18. *N*-methyltriazolium as a molecular station for the DB24C8 in a three-station based molecular muscle. Reagents and conditions: a) 1) HCl/Et₂O, 2) NH₄PF₆, H₂O/CH₂Cl₂, RT, 30 min; b) DIEA, Boc₂O, CH₃CN, RT, 5 d; c) DIEA, Boc₂O, CH₃CN.

In the disubstituted pyridinium amide series (compounds **45** and **47**), similarity was observed with respect to **40/42**—in other words, the ammonium is the best molecular station, and among the weakest stations, the triazolium is a better station than the disubstituted pyridinium amide. Therefore, at the protonated state **45**, the DB24C8 sits over the ammonium stations forcing the molecular muscle to adopt a stretched co-conformation. Unsurprisingly, in the carbamoylated^[48] molecular muscle **47**, the DB24C8 reside around the triazolium stations, triggering a half-contracted co-conformation. In contrast, carbamoylation of the monosubstituted pyridinium amide-containing rotaxane dimer **46** results in a Brownian motion between the two triazolium and pyridinium amide sites as already observed in the simpler [2]rotaxane **43**. The two translational isomers are in fast equilibrium between the contracted and the half-contracted co-conformations. However, in comparison with **43**, it is noteworthy that important changes are noticed. Indeed, the ratio of the two translational isomers are inverted in **48**, as if the triazolium seems to be now a better site of recognition for the DB24C8 than the monosubstituted pyridinium amide. In fact, this variation is directly linked to the doubly threaded architecture and is not correlated to a change of affinity of the two stations for the DB24C8. In the peculiar molecular muscle architecture, intramolecular repulsions are possible between the two triazolium cationic charges. The interactions between the DB24C8 and the monosubstituted pyridinium amide (which have been proved to be stronger than

with the triazolium) should logically force the structure to adopt a contracted state. However, in this state, the two triazolium moieties would face each other, which is thermodynamically unfavorable. Then, the molecular muscle **48** rather prefers to adopt a favored half-contracted co-conformation where no such repulsion remains. Unsurprisingly, decreasing the temperature led to the displacement of the equilibrium between the two translational isomers **48** as observed in **43**. Interestingly, the doubly interlocked architecture also makes possible the displacement of the translational equilibrium by tuning the repulsion through changing the polarity of the solvent. The repulsion between triazoliums proved to be the highest in a more polar solvent such as DMSO, hence favoring the half-contracted co-conformation at 97.5%, whereas the lowest repulsion was noticed in the less polar solvent CDCl₃, favoring the contracted co-conformation in a 88.5% ratio. In summary, this study gave new complementary information concerning the relative affinity of a triazolium moiety for the DB24C8. Moreover, we have demonstrated that it is possible to trigger a wide range of co-conformational states upon pH-stimulus. Among them, a co-conformational state in which a Brownian motion between two stations can be generated, controlled by a temperature stimulus or a change in solvent polarity, while it can be switched off by acidification.

The repulsion between triazoliums in the doubly interlocked structure is a very interesting feature that has dramatic consequences on some motions observed in double-lasso architectures (see Section 8). The conversion of the triazole to the triazolium, in order to get a molecular station for crown ether, also gave us the opportunity to introduce a group of interest at the side-chain of the thread.

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7 Dual Role of the Triazolium as Both a Molecular Station and a Kinetic Barrier for a Crown Ether

Triazolium has not only been utilized as a molecular station in a rotaxane molecular machine. It has also served as a kinetic molecular barrier that is able to compartmentalize^[1a,49] molecular machines by trapping the macrocycle around one of the two sides of a threaded axle. For that, we investigated the selective *N*-benzylation of triazole in order to incorporate such

a molecular barrier and station at the middle of a threaded axle that contains two other molecular stations.^[50] The aim was either to trap the macrocycle around stations of weaker affinity than that of anilinium, or to know if and how the DB24C8 could interact only over one of the two sides of the bulky triazolium moiety (Section 7.1). A further reported application of this methodological study consisted in locking a mono-lasso molecular machine possessing a bigger benzometaphenylene-25-crown-8 (BMP25C8) macrocycle (Section 7.2).^[51]

7.1 In a [2]rotaxane molecular architecture

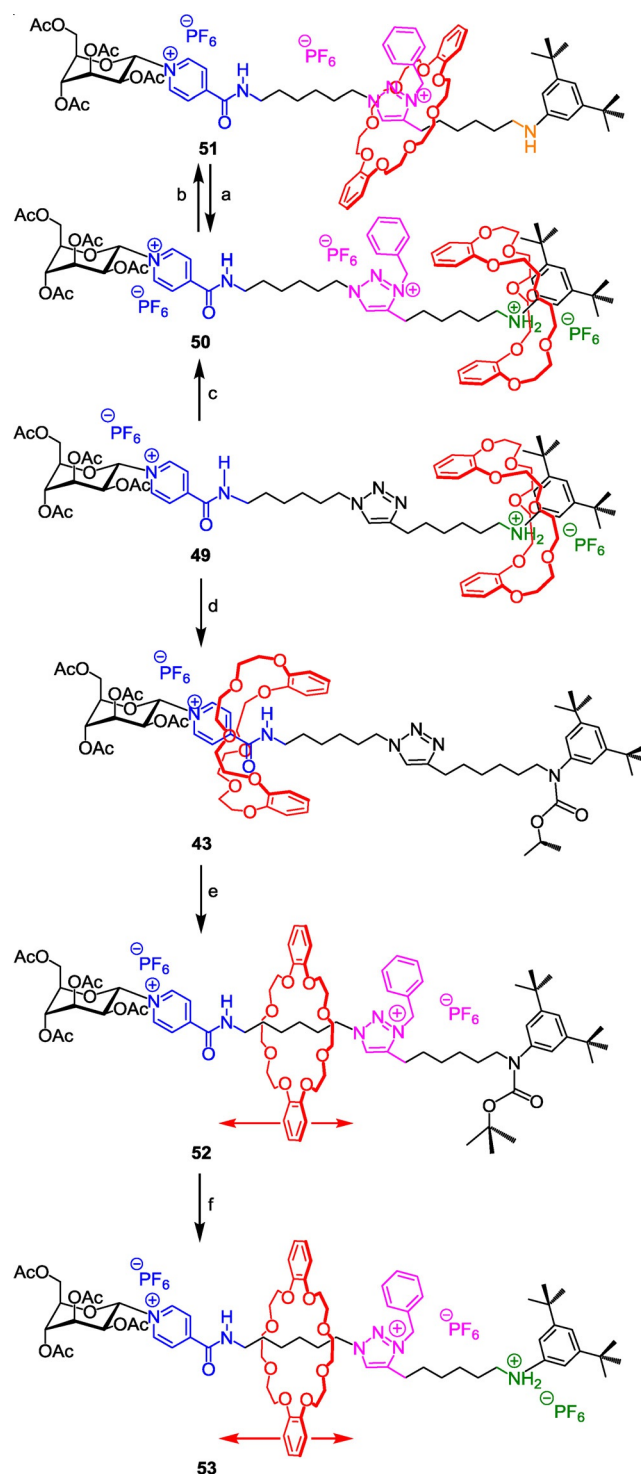
First, the two-station-containing [2]rotaxane molecular machine **49** was efficiently prepared following the two-step sequence synthetic strategy we described in 2008 (Scheme 19).

Given the difference of the relative affinities of the DB24C8 for the two present sites of interactions,^[52] the DB24C8 resides around the best anilinium molecular station. The *N*-benzylation of the triazole afforded the three-station [2]rotaxane molecular machine **50**, in which the *N*-benzyltriazolium proved to act as a molecular station. Indeed, deprotonation of **50** triggers the shuttling of the macrocycle as a result of the disappearance of the anilinium moiety. Whereas a kind of oscillating Brownian motion between the two stations of similar affinity for the DB24C8 (i.e. monosubstituted pyridinium amide and triazolium) was observed in the analogous *N*-methyltriazolium compound **43**, the behavior of the macrocycle appears quite different in the present case. In **51**, the DB24C8 only interacts with the *N*-benzyltriazolium station because the side bulky *N*-benzyl group acts as a molecular barrier that prevents the macrocycle from gliding across it. A so-called bi-stable pH-sensitive compartmentalized molecular machine is thus obtained, in which the DB24C8 shuttles from the ammonium to the triazolium site upon a pH-stimulus, without being able to reach the pyridinium amide site.

A diverted strategy from **43** was also applied in order to obtain **53**, that is to say the translational isomer of **50**, in which the DB24C8 is trapped on the left side of the threaded axle with respect to the benzyltriazolium barrier. A first carbonylation of the aniline moiety triggered the large-amplitude displacement of the DB24C8 around the pyridinium amide molecular station. The subsequent *N*-benzylation of the triazole afforded, after decarbonylation, the new compartmentalized molecular machine **53**. This time, and contrary to **51**, the DB24C8 undergoes a continuous motion between the pyridinium amide and the triazolium station, as already observed in the *N*-methyltriazolium [2]rotaxane **43**. Nevertheless, the molecular machine **53** is here independent of any variation in pH, even if the anilinium is still the best site of interactions for the DB24C8. Indeed, a persistent Brownian co-conformational state is observed in **53** whatever the presence of the best anilinium station, due to the presence of the bulky molecular barrier.

Therefore, to summarize, the triazolium moiety can be used as both a molecular station and a barrier in the same molecular machine, bringing new insights for the conception of new molecular machines. A particular interest of this newly de-

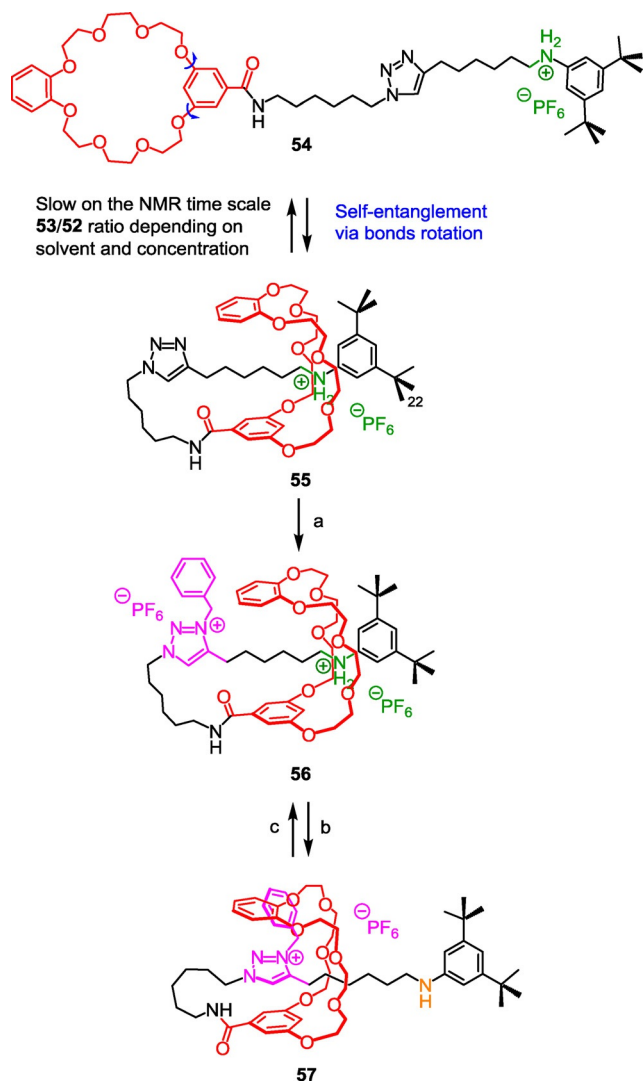
scribed feature of the triazolium moiety is the possibility to trap lasso molecules.



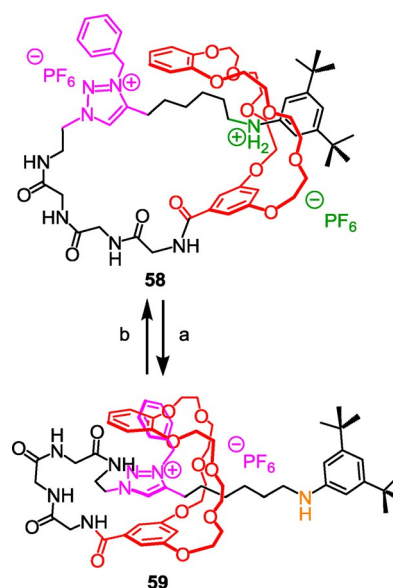
Scheme 19. *N*-benzyltriazolium as both a molecular station and a barrier for the DB24C8 in a [2]rotaxane molecular machine. *Reagents and conditions:* a) 1) HCl, 2) NH_4PF_6 ; b) DIEA; c) 1) BnBr, CH_2Cl_2 , 4 d, RT, 2) NH_4PF_6 , $\text{CH}_2\text{Cl}_2/\text{H}_2\text{O}$, 30 min, 79%; d) Boc_2O , DIEA, 15 h, RT, 93%; e) 1) BnBr, CH_2Cl_2 , 5 d, RT, 2) NH_4PF_6 , $\text{CH}_2\text{Cl}_2/\text{H}_2\text{O}$, 30 min, 65%; f) 1) HCl, 1 h, RT, 2) NH_4PF_6 , $\text{CH}_2\text{Cl}_2/\text{H}_2\text{O}$, 30 min, 60%.

7.2 In lasso structures

Taking advantage of the possibility to use *N*-benzyltriazolium as both a barrier and a station at the same time in an interlocked molecular machine, we reported, in 2013, the preparation of lasso molecular architectures^[53,54] using a self-entanglement strategy^[55] of a “hermaphrodite” compound (Schemes 20 and 21).



In the synthesized compound **54**, the thread is linked to a BMP25C8 crown ether macrocycle and contains an anilinium station as a necessary template to envisage the efficient formation of the rotaxane. Contrary to the strategies previously described, the tail of the thread cannot go through the macrocycle because it already holds a stoppering *tert*-butylphenyl end. Here, the only way to self-interlock is to use a large crown ether such as the BMP25C8 which allows the internal ro-



Scheme 21. *N*-benzyltriazolium as both a molecular station and a barrier in a peptide-containing lasso molecular architecture. Reagents and conditions: a) DIEA; b) 1) HCl/Et₂O, 2) NH₄PF₆, CH₂Cl₂/H₂O.

tation of the metaphenylene aromatic ring. In this case, the rotation of the *meta*-substituted aromatic ring of the BMP25C8 around two σ -bonds allows the intramolecular threading of the anilinium-containing tail through self-entanglement. Obviously, the interactions between the template and the crown ether are weaker than with the shorter DB24C8 which fits much better. Nevertheless, they are sufficient to get, at high dilution, the pseudo[1]rotaxane **55** in a satisfactory yield of 45%. The subsequent incorporation of the benzyl group on the triazole trapped the crown ether, hence locking the [1]rotaxane architecture **56** in a loosened conformation, due to the localization of the BMP25C8 around the best anilinium station. Treatment of the loosened lasso with a base causes the dramatic tightening of the lasso due to the shuttling of the BMP25C8 around the triazolium molecular station.

Using the same strategy, we then extended our studies to a peptide-containing lasso in which a GlyGlyGly tripeptide sequence is part of the loop of the lasso (Scheme 21).

Rotating-frame Overhauser enhancement (ROE) NMR experiments and drift-tube ion-mobility mass spectrometry proved the possibility to constrain more or less the peptide backbone of the lasso depending on molecular machinery. This first example of peptide-containing lasso now paves the way to many further possible applications in the selective targeting using peptide-containing lassos.

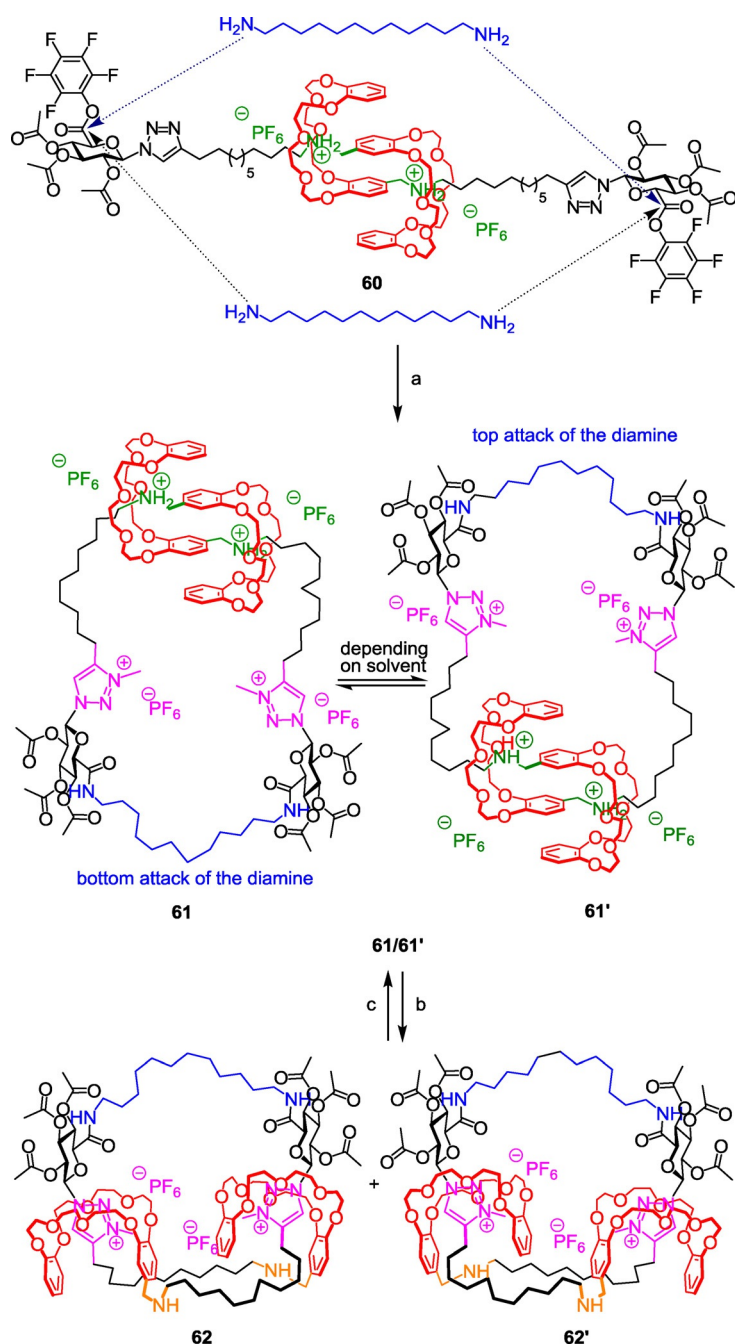
8 Triazolium-Containing Molecular Double Lassos: A Unique Spherical Molecular Muscle

In Sections 5 and 6, we described several examples of linear molecular muscles and their use as building blocks for polymers. In this section, we highlight a unique double-lasso^[56] molecular architecture that can act as a spherical muscle.^[57] Two distinct motions have been observed: a loosening–tightening

of the double-lasso and a controllable molecular “jump rope” movement of a molecular chain around a [c2]daisy^[58] arrangement.

8.1 Tightening–loosening the double lasso

The molecular spherical muscle relies on a double-lasso molecular architecture that can be obtained by the chemical connection of the two ends of a linear molecular muscle (Scheme 22).



Scheme 22. Triazolium-based double-lasso molecular architecture and loosening–tightening motion. *Reagents and conditions:* a) 1) H₂N(CH₂)₁₂NH₂ (1 equiv), CH₂Cl₂ (0.5 mM), 25 °C, 3 d, 2) CH₃I, 4 d, 3) NH₄PF₆, H₂O/CH₂Cl₂, 30 min; b) NaOH/H₂O/CH₂Cl₂; c) 1) HCl/Et₂O, 2) NH₄PF₆/H₂O/CH₂Cl₂, quanti **61/61'**.

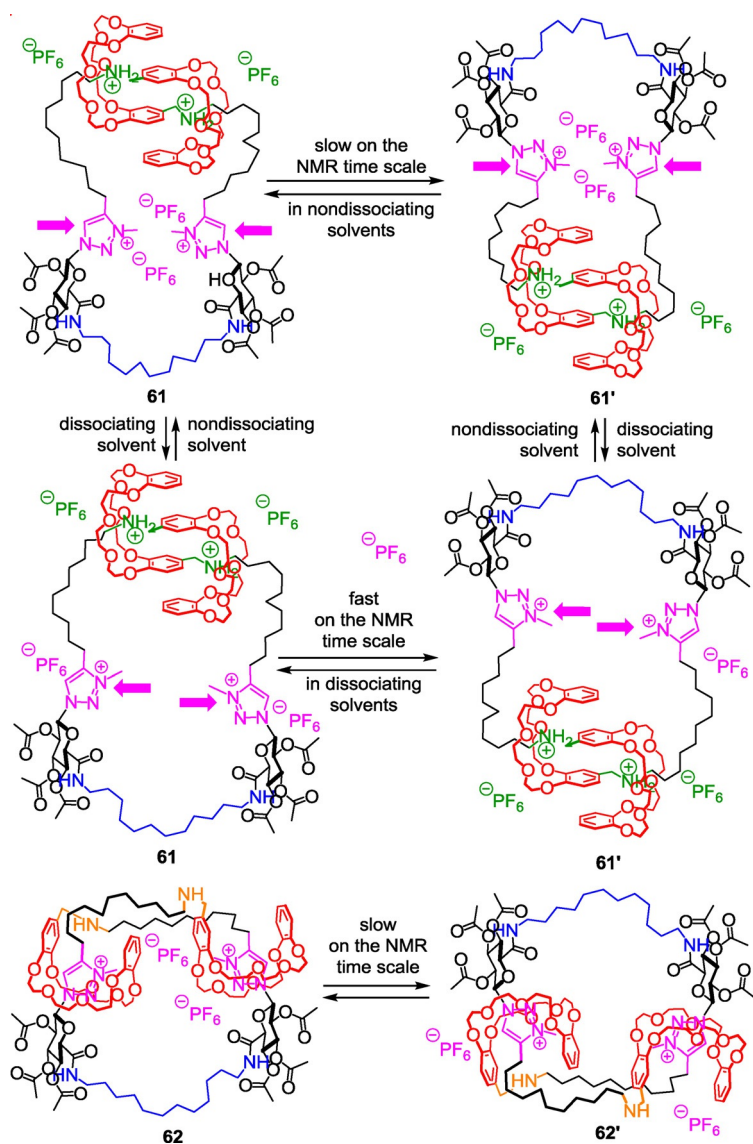
The synthesized [c2]daisy-chain-incorporating compound **60** was designed in order to give enough flexibility to the two extremities in order to permit the cyclization. Thus, the length of the alkyl chain between the ammonium station and the triazole moiety was increased with respect to the previously published molecular muscle **34**. Furthermore, two pentafluorophenyl active ester functions on the glucuronic acid derivative units serve as bulky stopper to lock the interwoven structure. They are also present as highly sensitive moieties toward amines, so that the dodecandiamine can close the double-lasso structure at high dilution, through the linking of the two extremities of the [c2]daisy chains **60**. Subsequent alkylation of the triazoles reveals, in **61**, the second molecular stations for the DB24C8 macrocycles. At this stage, it is interesting to notice that the attack of the diamine by the top or by the bottom of the molecule provides two atropoisomers, whose exchange can be controlled. This exchange is explained in detail in the following section. What interested us first was the deprotonation of the loosened double-lasso **61**. As expected, it triggers the tightening of the double lasso through the shuttling of the macrocycle around the triazolium sites. In **62**, two distinct double-lasso diastereomers were obtained. The size of the double-lasso molecules at the protonated and deprotonated states was evaluated thanks to diffusion-ordered spectroscopy (DOSY) experiments and molecular mechanics calculations. The results corroborate the impressive tightening–loosening internal motion of the lasso.

Designing new double-lasso molecular machines which are able to host a guest molecule with preferential interactions in one of the two co-conformational states is still an issue to tackle. One may imagine different kinds of applications of this new cyclic architecture in the future, like making molecular carriers that respond to a stimulus in order to release a guest molecule, or making switchable nanoreactors for asymmetric synthesis by taking advantage of the chiral helix-type double lasso. One may also think about using this sort of compounds for closing or opening a channel or even capturing molecules to decontaminate a medium.

8.2 Molecular jump-rope motion around the [c2]daisy moiety

Another interesting motion that can be controlled in the double lasso **61** concerns the rotation of the triazolium-containing chain around the [c2]daisy arrangement. We compared this movement to a molecular “jump rope” or to a molecular “turnstile”^[59] depending on if we consider the rotation of the cyclic chain with respect to the [c2]daisy arrangement or the inverse (Scheme 23).

As mentioned before, the possibility of the triazolium-containing chain to clear the [c2]daisy arrange-



Scheme 23. Molecular jump-rope motion in a triazolium-based double-lasso spherical molecular muscle.

ment allows the exchange between isomers **61** and **61'**. As long as the rotation is possible and fast, **61** and **61'** remain conformers, and only one set of ^1H NMR resonances is observed. However, tightening the cavity of the double lasso results in the deceleration or the stopping of this internal rotation. In that case, one set of distinguishable ^1H NMR signals is observed for each isomer.

Two different ways to tighten the double lasso were found in this unique structure. The first is a consequence of a pH stimulus, while the second results from the variation of the polarity of the solvent. Obviously, on one hand, adding a base induces the tightening of the spherical molecular muscle and therefore the impossibility for the cyclic chain to turn around the [c2]daisy chain. On the other hand, tuning the repulsion between the triazoliums that are facing each other in this singular structure enables us to disturb or favor the jump-rope movement in the loosened co-conformation **61**. This was achieved through the variation of the polarity of the solvent, as al-

ready mentioned in the three-station-containing molecular muscle **48**. The shape of the cavity of the double lasso appears larger in polar solvents like acetonitrile because the repulsion between triazoliums is higher. This causes a fast rotation of the molecular jump rope on the NMR time scale because no much steric hindrance disturbs the rotation around the [c2]daisy chain. It is exactly the contrary in dichloromethane where the two triazoliums sit much closer to each other, thus tightening the cavity of the double lasso in a new way (i.e. different from the tightening motion of the lasso ascribed to the gliding of the DB24C8) and to another shape than a basic reagent enforces. This singular tightening of the double lasso induces a slower rotation of the "jump rope" around the [c2]daisy arrangement due to steric hindrance. Interestingly, increasing progressively the proportion of one of the two solvents allows the progressive acceleration or deceleration of the motion.

9 Summary and Outlook

Since we first used the triazolium moiety in 2008 as a molecular station for the DB24C8 in a [2]rotaxane molecular machine, several examples of original molecular machines have emerged. From pure molecular design to applications in stereoselective catalysis or fluorescence output, from motion at the molecular scale to enhanced supramolecular polymer extension or contraction, and from simpler [2]rotaxane architectures to molecular muscles or lasso compounds, triazolium proved to be a molecular station and/or a kinetic molecular barrier of choice for the construction of a wide range of more or less sophisticated interlocked molecular machines. Its very easy access through a two-step sequence strategy, combined with its easy ^1H NMR characterization in both its naked state or in interaction with crown ethers makes the triazolium a multipurpose and very friendly moiety to handle. In addition, the molecular machinery appears very simple to monitor. Although triazolium has a very poor intermolecular affinity for crown ethers, it can interact pretty well in a locked molecular architecture in the absence of a better site of affinity, which makes triazoliums an ideal second molecular station. Moreover, the possibility to reach triazolium-based interlocked components devoid of any other efficient template using a diverted strategy based on a "transporter" of macrocycles has been recently published, opening the way to a wide variety of new triazolium-containing interlocked molecules. Finally, the possibility to form triazolium through the linking of various moieties to the nitrogen atom of a triazole group opens a wide range of further possibilities. Investigating new properties of the triazolium moiety, like its behavior as a redox molecular station in an interlocked machine, should certainly be of high interest for the construction of multiresponsive systems.

Keywords: lasso structures · macrocycles · molecular machines · molecular muscles · rotaxane · triazolium

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