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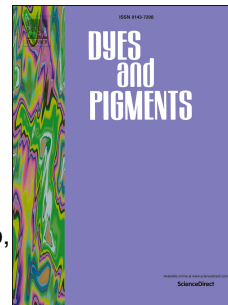
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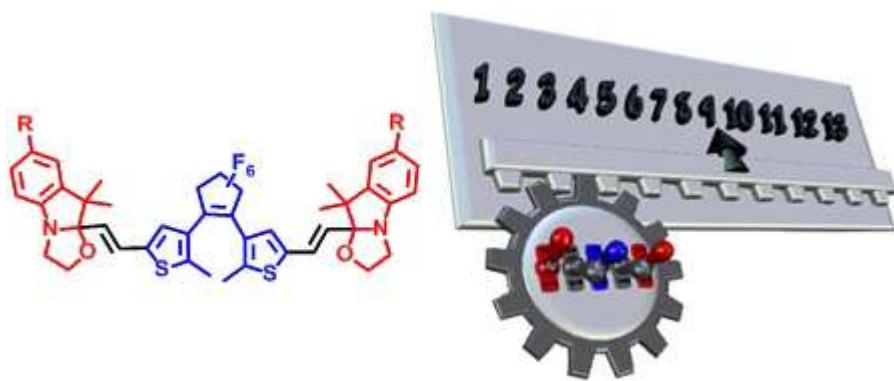
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ACCEPTED MANUSCRIPT

13 metastable states arising from a simple multifunctional unimolecular system

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ABSTRACT: A diarylethene core decorated with two benzooxazolidine side-arms through ethylenic spacers represents a smart example of multi-addressable system whose reversible responses could be selectively activated on demand. UV-Visible and NMR spectroscopies

and electrochemical studies allow to overview its performances when stimulated by light, acid/base and electrons, then underlining its photochromic, acidochromic and electrochromic properties. The multichromophoric combination could be considered as multifunctional and multistate systems, as interconversions can be performed by different stimuli whereas each stimulus provides a specific metastable state.

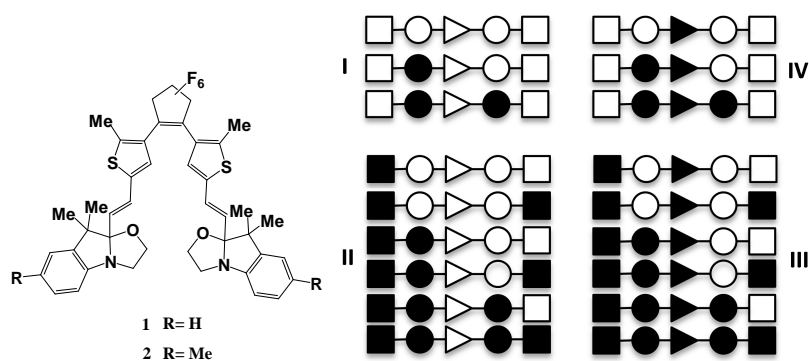
1. Introduction.

Photochromic diarylethenes possessing reversible sets of physicochemical properties have been largely exploited for applications such as switches, rectifiers and memories [1-3]. In fact, their excellent photochromic performances encompassing fatigue resistance, short time response, high quantum yields and no thermal isomerization, ensure to switch upon irradiation between colorless and closed colored forms which exhibit different properties. Furthermore, diarylethenes have been reported to exhibit electrochromic behavior performing cyclisation/ ring opening by application of an electrochemical potential [4-6]. Indolino[2,1-b]oxazolidine derivatives [7-9] belong to a more recent family of photochromic dyes displaying multimodal photo- and acidochromic properties. In addition, the substitution of an indolino[2,1-b]oxazolidine unit with a bithiophene redox probe generated a system capable to be commuted reversibly and indifferently either under UV irradiation, acidity changes or electrochemical stimulation, and thus, confers to this molecular system a pretty and uncommon trimodal switching behavior. [10]

In this context, the combination of diarylethene and indolino-oxazolidine units through covalent linkers offers a multi-addressable system expected to provide up to 2^n distinct states (with n the number of addressable units). The main challenge of such system is not only to keep the selective and controlled addressing of individual units but also to find the right

equilibrium between multifunctional (the same interconversion can be performed by different stimuli) and multistate (each stimulus provides a specific metastable state) properties.

The present paper deals with the design and the investigation of the behavior of a novel molecular system **1** associating the two types of photochromic compounds through ethylenic junction (Scheme 1) and its bis-methyl substituted derivative **2**. As consequence, the presence of five addressable units should lead to 2^5 different states according to the closed or open forms of indolino-oxazolidine, associated with trans or cis ethylenic junction to the open or cyclized diarylethene and excluding the absolute stereochemistry of the molecular system (due to the presence of an asymmetric carbon atom in the closed forms of indolino-oxazolidine). The 32 possible states are reduced to 18 due to symmetry consideration. Each unit exhibiting different and complementary multimodal switching abilities, their association should lead to the development of a promising molecular multistate system where the individual units could be selectively addressed by selecting the appropriate stimulation (light, chemicals, and electron). This keypoint is particularly relevant for molecular memory or logic gate operators [11-14]. As a consequence of their mutual influence, the commutation of the different addressable units is expected to induce a drastic change of the optical properties of the whole molecular system. In this context, the UV-Visible absorption spectroscopy is certainly the simplest characterization technique to monitor the different commutation states under various stimuli. In parallel, NMR spectroscopy offers an elegant way to structurally characterize the various isomers.



Scheme 1. Target compounds **1** and **2** and the eighteen possible combinations (\square Box, \circ C=C, \triangleright DAE)

2. Results and discussion.

2.1 Investigations of processes between all-trans isomers of targets in their initial state **I_{tt}**

Addition of trifluoroacetic acid (TFA) in CH_3CN solution of target **1** (**2**) in its initial state **I_{tt}** induced the opening of both oxazolidine rings into the corresponding protonated form **II_{tt}** (See scheme 2 for structures of states) concomitantly with color change of solution from colorless to yellow (Figure 1, **1**; data for **2** are in supplementary materials). The successive addition of some acid aliquots revealed an isosbestic point evidencing that the oxazolidine ring opening did not occur in a stepwise manner. Indeed, under their closed forms, the indoline heterocycles are generally almost orthogonal to their olefinic substituent [15-16]. As a consequence, no interaction through bonds between them could be expected leading to an independent behavior and then a concomitant commutation of them when the indolino-oxazolidine units, hereafter named BOX for convenience, are directly stimulated. Absorption band is bathochromically shifted to 442 nm (443 nm) with a concomitant decrease of the band at 280 nm, translating the extension of the π conjugated system and the establishment of a charge transfer band between thienyl donor and indoleninium acceptor parts. The indoleninium moieties ($\text{N-CH}_2\text{-CH}_2\text{-OH}$) are clearly identified by two well-resolved triplets at 4.0

and 4.6 ppm in ^1H NMR spectrum, characterizing **II_{tt}**, having its ethylenic junctions in *trans* isomery, as indicated by the two doublets at 7.2 (I=2H) and 8.4 (I=2H) ppm with a vicinal scalar coupling of 15.9 Hz (Figure 2). Closure of BOX units upon neutralization with triethylamine (TEA) vapor was confirmed by the disappearance of absorption band at 442 nm and the initial absorption spectrum was recovered entirely. This was nicely evidenced by NMR studies where a concomitant disappearance of triplet signals at 4.0 and 4.6 ppm and the reappearance of CH_2 signals for closed BOX between 3.3 and 3.8 ppm were observed. Consequently, the stimulation of the system by pH change allows to switch between states **I_{tt}** and **II_{tt}** in a selective, quantitative and reversible manner.

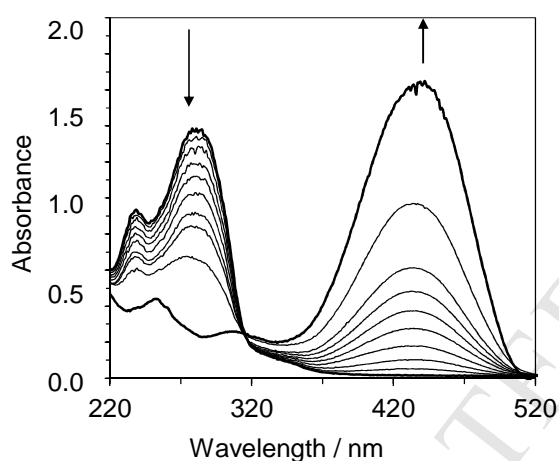


Figure 1. Absorption changes **I_{tt}**→**II_{tt}** (**1**, 2.5×10^{-5} M, CH_3CN) upon acid addition.

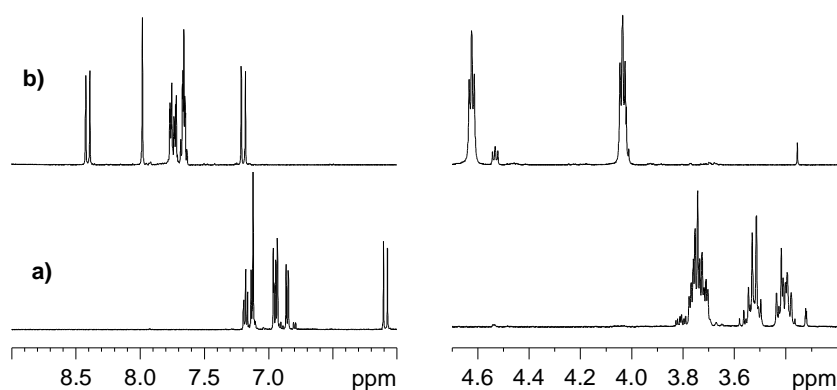


Figure 2. ^1H NMR spectrum a) of state **ltt** b) of state **II** (**1**, 10^{-3} M, CD_3CN).

Both DAE and BOX exhibit photochromic behavior, nevertheless irradiation of target **1** (**2**) in its initial state **ltt** in acetonitrile with 254 nm or 312 nm light conducts to selective cyclization of the diarylethene unit (abbreviated as DAE) as supported by the appearance of a band at 585 nm (583 nm), and the concomitant decrease of the band at 280 nm (285 nm) (Figure 3). This process is accompanied by a color change of solution from colorless to blue. From ^1H NMR spectrum, 90 % of conversion is quantified from the CH_3 and the thienyl ^1H signals shifted from 1.95 to 2.1 ppm, and from 7.1 to 6.5 ppm, respectively, evidencing the cyclization of DAE in **IV** (Figure 4). State **IV** is thermally stable for weeks at ambient temperature, while bleached completely back to colorless solution with visible light (577 nm) irradiation.

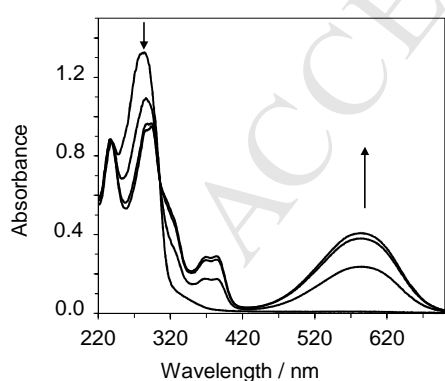


Figure 3. Absorption changes **ltt**→**IV** (**1**, 2.5×10^{-5} M, CH_3CN) upon 312nm light irradiation.

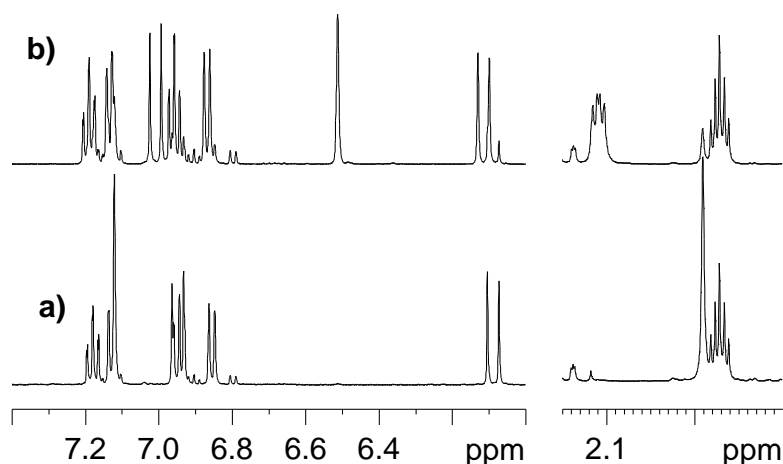


Figure 4. ^1H NMR spectrum a) of state **I** b) of state **IV** (**1**, 10^{-3} M, CD_3CN).

Consequently, the stimulation of the system by UV then Visible light induces the switch between the states **I** and **IV** in a selective, almost quantitative and reversible manner. Whatever the wavelength selected no traces of photoinduced ring-opening of BOX accompanied the DAE cyclization. To stimulate the BOX motif, chlorobenzene was added as a photo-sensitizer [7,17]. Irradiation with 254 nm light of state **I** in acetonitrile/chlorobenzene (90/10) solution reveals the appearance of new bands in 420-440nm range (open BOX, **II**) and in 580-600 nm (cyclized DAE, **IV**) (Figure 5). When irradiation time is prolonged, the band characterizing cyclized DAE is shifted into a new band in 700-770 nm area. The same experiment was repeated and followed by NMR spectroscopy, confirming the transformation of state **I** into states **II** and **IV**, the latter being then converted into state **III'**, with only one of the BOX in open configuration. This means that addressability of **I** by 254 nm light in acetonitrile/chlorobenzene solution is not selective as it conducts to a mixture of states (**II**, **IV** and **III'**).

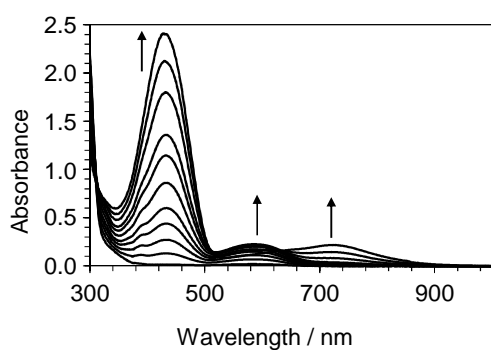


Figure 5. Absorption changes **I_{tt}→II_{tt}+IV_{tt}(+III'_{tt})** (**1**, 2.5×10^{-5} M, CH₃CN/CIPh) upon 254nm light irradiation.

This selective addressability issue can be circumvented by changing the nature of the stimulation to acidity conditions (*vide supra*) or by applying irradiation wavelengths within an appropriate sequence. In a first step, 312 nm light was applied to acetonitrile/chlorobenzene solution of targets leading selectively to the DAE cyclisation (**I_{tt}→IV_{tt}**, 90%, $\lambda = 590$ nm). Clearly, no typical signals for the open oxazolidine ring or for the E/Z isomerization of double bond can be detected. In a second step, the BOX units of **IV_{tt}** were activated following two different ways: i) by acid addition leading to a color change of the solution from blue to green, characterizing the appearance of the state **III_{tt}** (both BOX are open) whose absorption bands are centered at 450 and 780 nm for target **1** (455 and 773 nm for **2**) (Figure 6a).

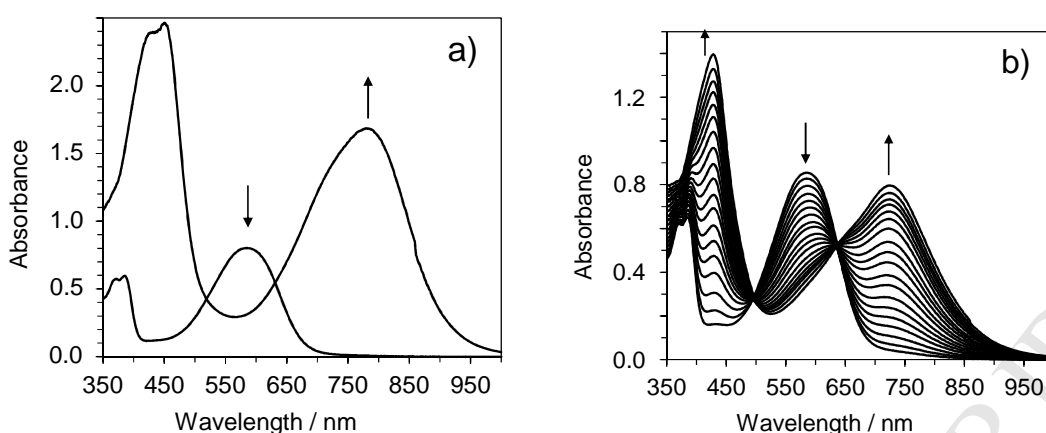
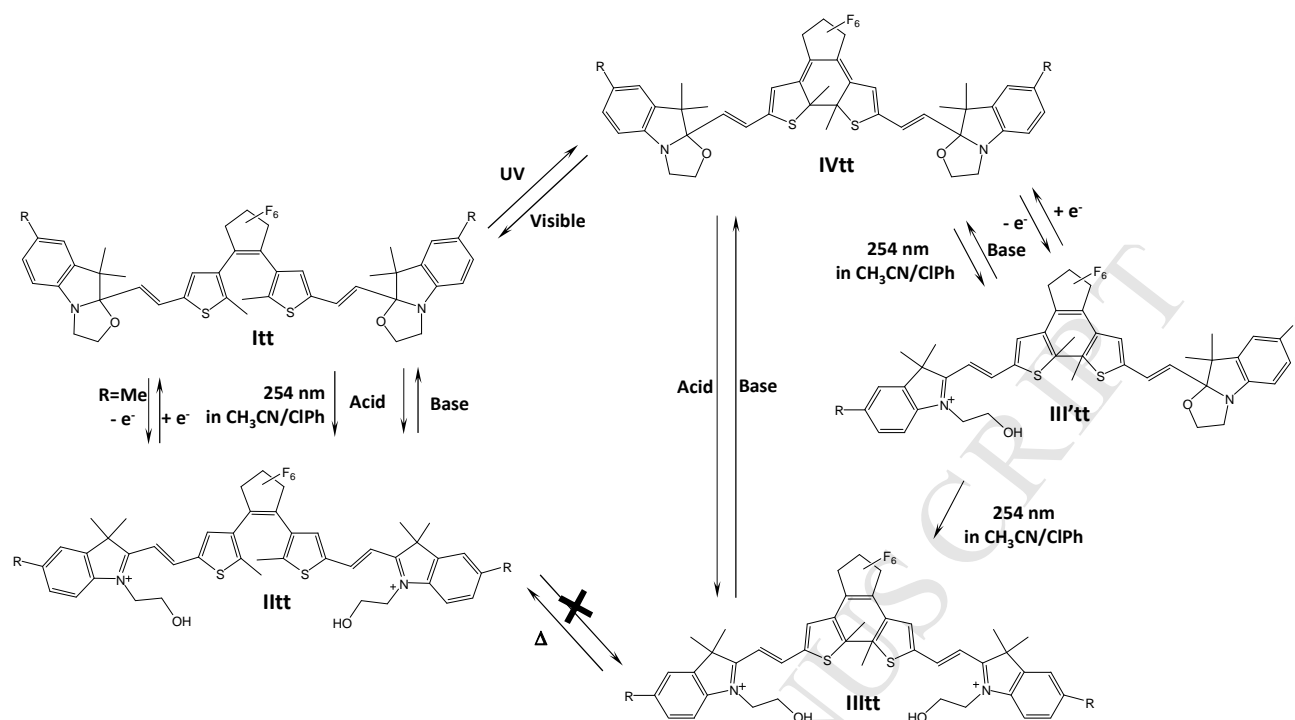


Figure 6. Absorption changes (**1**, 2.5×10^{-5} M, CH₃CN/CIBz) a) **IVtt** → **III'tt** upon acid addition b) **IVtt** → **III'tt** upon 254 nm light irradiation.

The simultaneous opening of both oxazolidine rings is well confirmed by NMR spectroscopy with the CH₂ triplet signals at 3.85 and 4.55 ppm (open BOX) and the CH₃ signal at 2.25 ppm (Cyclized DAE). ii) by irradiation with 254 nm light of state **IVtt** in presence of chlorobenzene, inducing a strong bathochromic shift of the absorption bands (Figure 6b). More important, the opening of the two BOX units is no anymore simultaneous but occurs in stepwise manner when the DAE is closed. It is clearly indicated by the appearance of new absorption bands centered at 428 and 723 nm (432 and 722 nm for **2**) with a concomitant decrease of the absorption at 585 nm (582 nm for **2**), and the clear evidence of isosbestic points supporting the presence of only two species state **IVtt** and state **III'tt** (cyclized DTE with one open and one closed BOX). When the duration of irradiation is prolonged, a bathochromic shift of the band at lower energy followed by a rapid decrease of its absorption intensity, while a continuous increase and hypsochromic shift of the band at higher energy are observed. The shape modification accompanied by the absence of isosbestic point let presume that if the opening of the second BOX occurs well leading to the state **IIItt** (cyclized DTE with two open

BOX), the stability of this last one is limited in our experimental conditions. Indeed, while similar compounds associating the DAE unit with one BOX unit showed no thermal evolution [18-19], here the substitution with two open BOX leads to a slow thermal opening of DAE as observed by UV and NMR spectroscopy ($^{293}k_{\Delta} = 5.3 \times 10^{-5} \text{ s}^{-1}$) towards state **II_{tt}**. Then, the restoration of the molecular system under its initial state **I_{tt}** can be achieved : from **III_{tt}** or **III'_{tt}** upon base treatment leading to **IV_{tt}**, then bleached with visible light, or from **III_{tt}** upon thermal relaxation leading to **II**, then neutralized with base.

As a consequence, the status of the central DAE moiety strongly influences the photochemical behavior of the BOX units. When DAE is open (State **I_{tt}**), irradiation in presence of chlorobenzene with 254 nm light activates the simultaneous opening of both BOX units (State **II_{tt}**). In contrast, when DAE is cyclized (State **IV_{tt}**), stepwise opening of BOX occurred (States **III_{tt}** and **III'_{tt}**). In the same way, the response of the DAE unit is dependent on the BOX status. In fact, all attempts to cyclize the DAE when both BOX are open (state **II_{tt}**) using UV light at different wavelengths were unsuccessful. Contrary to previous investigations on various DAE substituted with one BOX unit where the photocyclization efficiency of diarylethene is strongly reduced by the status of BOX, here, its photochromic behavior is totally suppressed with both open BOX. In resume, the different evidenced interconversions between the all-trans states are displayed in Scheme 2.



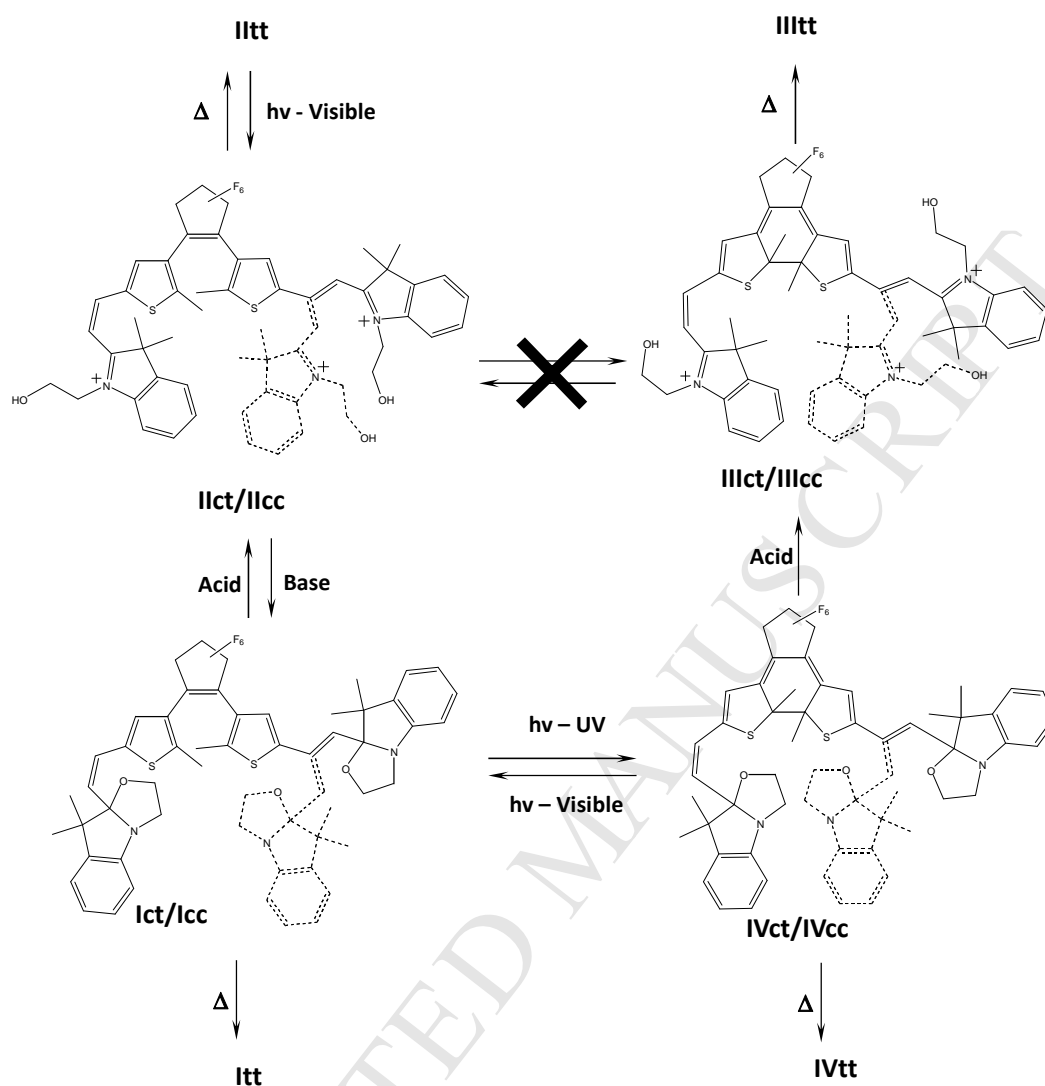
Scheme 2. Reaction pathway between all-trans isomers

2.2 Investigations of processes between cis isomers of targets

Based on previous investigations on similar molecular system [18,19], the photo-isomerization of the ethylenic junction between BOX and DAE is expected, especially when the pending BOX is open. In this context, the *trans-cis* isomerization of both ethylenic junctions has been studied by ^1H NMR spectroscopy. When the state **II'tt** is generated from initial state **I'tt** (previously shown to have all-trans junctions), both ethylenic junctions exhibit a *trans* geometry, as clearly evidenced by two doublets at 7.2 and 8.4 ppm with a scalar coupling constant of 15.8 Hz. Under UV light (254 or 313 nm) no cyclization of DAE from **II'tt** to **III'tt** occurred, but *trans-cis* isomerization of the ethylenic bonds was observed, and was more efficient when UV is replaced by visible light ($\lambda > 400$ nm). The state **II'tt** is then photo-

transformed into two isomers: **IIct** (20% at PSS; isomerization of only one ethylenic bridge) characterized by four doublets (7.1. and 8.4 ppm with $^3J=15.8$ Hz, and 6.5 and 7.5 ppm with $^3J=13$ Hz) and **IIcc** (70% at PSS; isomerization of both ethylenic bridges) characterized by two doublets (6.45 (I=2H) and 7.45 (I=2H) ppm with $^3J=13$ Hz). These two isomers with at least one *cis*-isomery bond relaxed thermally into the state **IIItt** ($k_{IIcc \rightarrow IIct} = 9 \cdot 10^{-4} \text{ s}^{-1}$ and $k_{IIct \rightarrow IIItt} = 5 \cdot 10^{-4} \text{ s}^{-1}$ at rt). As expected, these two *cis*-isomers of state **II** (**IIct** and **IIcc**) can also be neutralized with TEA leading to the closing of BOX into the corresponding states **Ict** and **Icc**. In contrast, they are photochemically silent, as no irradiation enabled to cyclize the diarylethene unit. Concerning the two *cis*-isomers of state **I** (**Ict** and **Icc**), they evolved very slowly through the thermal *cis-trans* isomerization into initial full-*trans* **IItt** ($t_{1/2}$ at rt : **Icc** = 5 days; **Ict** = 10 days). When they are submitted to UV irradiation, the cyclization of DAE (**Ict** \rightarrow **IVct** and **Icc** \rightarrow **IVcc**) is induced. The two isomers **IV** with at least one *cis*-isomery bond relaxed thermally into the state **IVItt** ($k_{IVcc \rightarrow IVct} = 3 \cdot 10^{-4} \text{ s}^{-1}$ and $k_{IVct \rightarrow IVItt} = 1.5 \cdot 10^{-4} \text{ s}^{-1}$ at rt). Surprisingly, upon acid addition on *cis*-isomers of state **IV** (**IVcc** + **IVct**), the state **IIIItt** was directly detected while its corresponding *cis* isomers (**IIIcc** + **IIIct**) would have been expected. They were not observed probably due to a high thermal instability. It should be also noted that the reverse reaction of *trans-cis* isomerization from **IIIItt** was never observed whatever the stimulus used.

From this study, we can conclude that all the states having at least one ethylenic junction in *cis* isomery present a thermal reversibility and no photoreaction occurred between the states **II** and **III**. The set of processes between all *cis*-states is summarized in Scheme 3.



Scheme 3. Reaction pathway between cis isomers

2.2 Electrochromic response of targets

BOX [10,20] and DAE [4-6] units can also be electrochemically addressed. These electroinduced behaviors represent an additional and alternative way to stimulate them selectively. In this context, the electrochemical behaviors of **1** and **2** were investigated by cyclic voltammetry (CV). The CV of both compounds under their initial state **I_{tt}** reveal one irreversible oxidation process at 0.78 and 0.77V for compounds **1** and **2** (Figure 7a-b)

respectively and are assigned to the oxidation of the BOX units. As previously observed [20], this oxidation conducts, when no substituent is present in position 5 of the indoline heterocycle (Target **1**), to a C-C oxidative coupling between BOX unit either in an intra or intermolecular manner and translated on the CV by the emergence of new redox systems at lower potential (0.46V) on the scan back (Figure 7a). This assumption is clearly confirmed by spectroelectrochemistry experiment. In fact, once the oxidation potential reaches 0.78V, one absorption band centered at 494 nm appears, being the spectral signature of the formation of the oxidized BOX dimer [20] (Figure 7c).

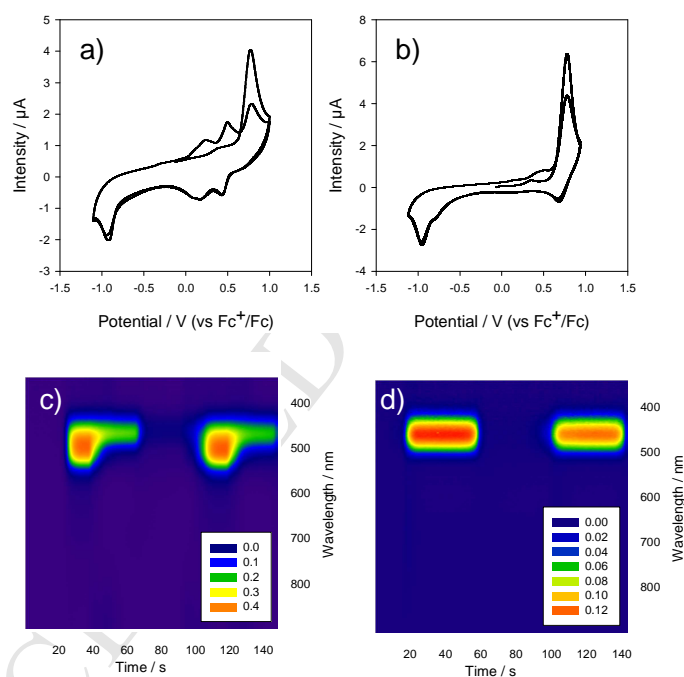


Figure 7. Thin layer cyclic voltammetry of a) **1** b) **2**, and corresponding spectroelectrochemistry of c) **1** d) **2**, in ACN under state **Itt**.

At the opposite, the presence of a methyl group (Target **2**) avoids the C-C oxidative coupling, then permitting the electroinduced ring opening of the oxazolidine as evidenced by the

irreversibility of oxidation process at 0.77V and any appearance of new signal at lower potential (Figure 7 b). Moreover, the electrochemical oxazolidine opening is confirmed by the spectro-electrochemical experiment. Indeed, an absorption band at 460 nm is generated when reaching 0.77V as oxidation potential (Figure 7d). Subsequent reduction of the system at -0.96V restores the initial state. As a consequence, the electrostimulation of the target **2** allows to switch between states **I^{tt}** and **II^{tt}** in a reversible manner as confirmed by the spectro-electrochemical experiments with the successive appearance and disappearance of the absorption band at 460 nm as function of the electrochemical potential.

The electrochemical properties of both targets under their state **IV^{tt}** (generated initially by UV irradiation of state **I^{tt}** with 312 nm light to photocyclize the DAE unit) were also studied. A strong cathodic shift of the first oxidation potential is observed to reach ca. 0.61 and 0.63V for **1** and **2** respectively (Figure 8), whatever the substitution pattern is, and thus, is assigned to the oxidation of the DAE core in agreement with previous studies on electrochemical behavior of various DAE derivatives [5-6].

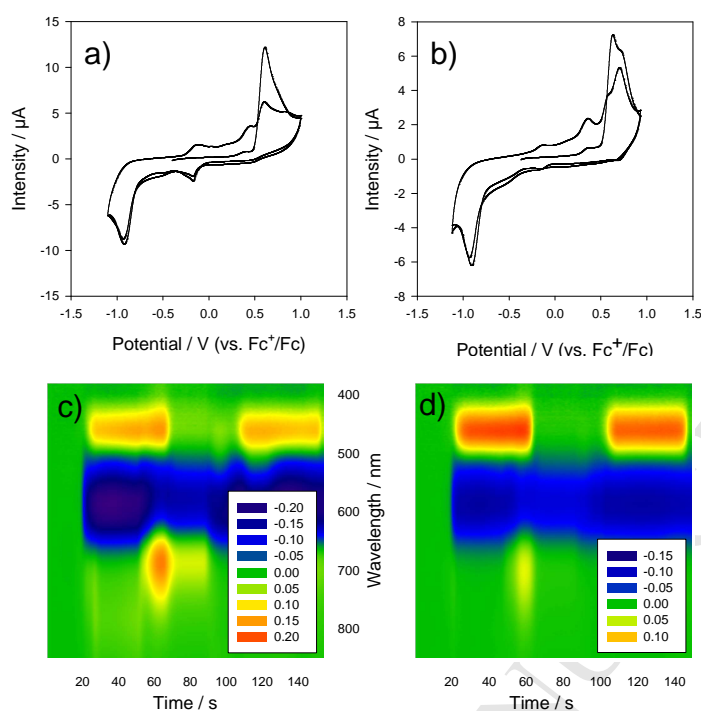


Figure 8. Thin layer cyclic voltammetry of a) **1** and b) **2**, and spectroelectrochemistry experiment of c) **1** and d) **2**, under the state **IV^{tt}** with TBAP as electrolyte at 20mV/s on Pt.

When the positive charges are not efficiently stabilized, the generation of closed dicationic DAE leads generally to the opening of the DAE which is translated on CV by an irreversible oxidation process [5-6] and the appearance of a signal at higher potential corresponding to the oxidation of the open form of DAE. In the present study, the oxidation of the state **IV^{tt}** conducts to the appearance of a new redox system at lower potential (-0.14 V for **1** and -0.11V for **2**) that could result from the opening of one BOX unit. In fact, some of us have recently demonstrated that the electrochemical stimulation of a redox probe associated to a BOX such as a bithiophene induced the oxazolidine ring opening [10]. In order to verify this assumption, the electrochemical properties of **1** under the state **III^{'tt}** (prepared by irradiation of **I^{tt}** with 312 nm light (**I**→**IV**), then with 254 nm light (**IV^{tt}**→**III^{'tt}**) in

acetonitrile/chlorobenzene) were studied. The CV (Figure 9a/b) exhibit a reversible oxidation process at -0.14V, then supporting the electrochemical opening of one oxazolidine ring via the oxidation of the central DAE unit observed previously. Moreover, this assumption is nicely confirmed by the spectroelectrochemistry experiments (Figure 8 c/d) with the successive disappearance of the band characteristic of the state **IV^{tt}** (585 and 593nm for compounds **1** and **2** respectively) and the appearance of the band corresponding to the state **III^{tt}** (428/723 nm and 432/722 nm for compounds **1** and **2** respectively). In addition, the irreversible reduction at -0.920 et -0.911V for **1** and **2**, leads to the disappearance of these characteristic bands, translating as expected the closure of the oxazolidine ring towards the state **IV^{tt}** under electrochemical stimulation, as previously observed with the process of closing from **II^{tt}** to **I^{tt}**.

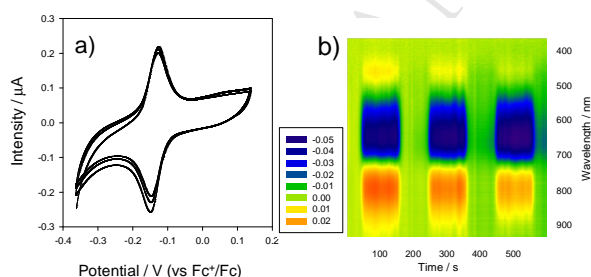


Figure 9. a) Thin layer cyclic voltammetry and b) corresponding spectroelectrochemistry experiment of **1** previously irradiated at 313nm then at 254nm (state **III^{tt}**).

However, the corresponding oxidized species of the state **III^{tt}** present a strong instability in our experimental conditions especially when higher potential is applied. At the time scale of the electrochemistry experiments, this instability (larger with **2** than with **1**) results to the opening of the central DAE moiety as confirmed by the increase of the absorption intensity at 460 nm. The hypothesis that the oxidation of the state **III^{tt}** induces the opening of the second BOX unit providing the state **III^{tt}** then after relaxes into state **II^{tt}** as observed by UV and NMR

spectroscopy is not consistent with electrochemical data. First, the spectro-electrochemical experiment performed on the state **III'tt** obtained photochemically reveals some absorption variations which are not matching with the state **IIItt**. Second, opening of the second BOX unit requests the formation of a radical cation on the central DAE which is not consistent with general effect of the substituent nature on redox potential. In fact, the opening of one BOX unit leading to the formation of an indoleninium moiety acting as strong electron withdrawing group, leads to an impressive cathodic shift (almost 500mV). However, such an unexplained cathodic shift was already observed on DAE. In 2005, Browne et al. studied the electrochemical behavior on various substituted DAE and reported the existence of a competitive oxidation process to the classical oxidation of cyclized form (at -0.1 V for the 5-(p-methoxyphenyl)-5'-phenyl-perfluoroDAE as example) leading to an unknown species referenced as X form. [5,6]. As consequence, we can presume that an identical process occurs here. The authors suggested that the X intermediate results from a reversible intramolecular rearrangement of the dicationic species of the closed form into a more stable species which would explain its appearance only on the scan back of the closed form oxidation and its irreversibility. At the opposite, we can observe the reversible oxidation of this intermediate in solution of compound **1** under the state **III'tt** obtained by successive irradiation at 313 then 254 nm. This fact lets us envisage its formation directly from the neutral closed DAE and strongly correlated to the donor or acceptor nature of the substituent borne by the DAE unit. As consequence, the complete clarification and understanding of the formation of this X form request deeper investigation.

3. Conclusions.

The designed targets incorporate five addressable units which could generate up to 32 different states, constituting the most sophisticated system to date. However, their number is here restricted to eighteen due to the symmetrical substitution of the diarylethene central unit. In the present work, we have demonstrated that thirteen of them can be effectively produced and detected. The way to access to each of them was also extracted by playing with selected stimuli on the multimodal properties of the system, such as light with specific wavelengths associated with the nature of solvent (Photochromism), pH variation (acidochromism), temperature (thermochromism) and electrons (electrochromism). Thus, the present target represents a nice example of system having multifunctional and multistate properties. In addition, a gated photochromism of the DAE unit was underlined. Indeed, the photoinduced cyclization with UV light of DAE from its open form towards the closed one (**I**→**IV**) is selective and almost quantitative in acetonitrile solution when the BOX units are closed and the reverse reaction is achieved with visible light. Once BOX are open, the photochromism properties of the DAE are completely inhibited and no cyclization from **II** to **III** occurred.

At the opposite, the BOX addressability by pH change is unaffected by the status of the central DAE and its opening and closure (**I**→**II** and **IV**→**III**) can be achieved selectively and quantitatively upon acidification and neutralization of solution respectively. Addressing of BOX opening with light is impeded in acetonitrile solution, but is revealed in the presence of chlorobenzene as photosensitizer and upon 254 nm light irradiation. Interestingly, the simultaneous opening of both BOX is achieved from irradiation of the state **I**, (**I**→**II**) whereas the process is successive from irradiation of the state **IV** (**IV**→**III'**→**III**). Contrary to previous investigated derivatives joining one DAE and one BOX, the state **III** is not thermally stable and slowly evolves towards the state **II**, by decyclization of its DAE part. Such a thermal unstability

was also encountered with all the states having at least one of the ethylenic junctions in cis isomery.

Beside their photo- and acidochromic properties, BOX moieties are also electroactive and their electrochemical behavior is strongly affected by the status of the central DAE. Oxidation of the targets in their states **I** conducts to a C-C oxidative coupling for **1**, while to the simultaneous opening of the oxazolidine ring (**I**→**II**) for methyl-substituted **2**. Such behavior between **1** and **2** compound is no longer observed when the central DAE is closed. Indeed, the oxidation occurring mainly on the cyclized DAE in the state **IV**, leads surprisingly to the selective opening of only one BOX unit (**IV**→**III'**) in both targets. The impossibility to induce the opening of the second BOX was assigned to the oxidation of **III'** into an **X** form already observed on few DAE based compounds, and being under deeper investigation to identify it.

4. Experimental section

4.1 Synthesis of targets

The synthesis of the targets is pretty straightforward and start by the elaboration of the 4,4'-bis(2-carboxaldehyde-5-methylthiophene)perfluorocyclopentene from corresponding dichloro derivative according to the procedure described by Feringa et al. [21]. Its further functionalization lays down on the high reactivity of the indoleninium or indolinooxazolidine derivatives with different aromatic aldehydes or nitroso compounds [22]. Hayami et al. reported that the reaction can be performed under several different experimental conditions (protic and aprotic solvents, solvent-free conditions, addition of a base) [23]. Nevertheless, the majority of the reported examples are carried out in boiling alcoholic solvents without any additives and gave satisfactory yields. However in our case, the reflux in ethanol provides the

diacetal and not the targeted molecules. In order to circumvent this issue, a simple solution consisted to replace the ethanol by a more hindered solvent such as t-butanol. The targets **1** and **2** have been fully characterized by NMR and the ethylenic junctions are both in *trans* isomery, as confirmed by the two doublets at 6.1 (2H) and 6.95 (2H) ppm with a vicinal scalar coupling of 15.8 Hz.

4.2 NMR and UV spectroscopies.

NMR spectra were recorded on Avance Bruker 500 or 300 spectrometer (^1H , 500 MHz, ^{13}C , 125 MHz, or ^1H , 300 MHz, ^{13}C , 75 MHz) equipped with TXI or QNP probe, using standard sequences. Data sets were processed using Bruker Topspin 3.2 software. Samples are dissolved in acetonitrile- d_3 or in acetonitrile- d_3 /chlorobenzene- d_5 in NMR tubes. Photoirradiation was carried out directly into the NMR tube in a home-built apparatus with a 1000W high-pressure Hg – Xe lamp equipped with filters. Monochromic UV or visible light was obtained by passing the light through a first filter, then through an interferential one. Irradiation with 254 nm light was achieved in rotating quartz NMR tubes (5 mm) at 295 K with a Bioblock Scientific VL-6LC lamp (12 W).

UV-visible spectra were recorded with a Perkin-Elmer lambda 950 spectrophotometer. Photoirradiation were carried out directly into the quartz absorption cuvettes from Hellma using TLC observation lamp allowing to switch between two different UV tubes (8W) purchased from Vilber-Lourmat emitting respectively at 254 and 312 nm.

4.3 Electrochemical experiments.

Cyclic Voltammetry. Acetonitrile (HPLC grade) and tetra-n-butylammonium hexafluorophosphate (TBAP, electrochemical grade, Fluka was recrystallised from ethanol). Cyclic voltammetry (CV) was performed in a three-electrode cell equipped with a platinum millielectrode, a platinum wire counter-electrode and a silver wire used as a quasi-reference electrode. The voltammograms were recorded on a potentiostat/galvanostat (BioLogic – SP150) driven by the EC-Lab software with positive feedback compensation. All the potentials reported were calibrated versus ferrocene/ferrocenium oxidation potential (+0.405V vs SCE or +0.425V vs Ag/AgCl).

Time-resolved spectroelectrochemistry was performed using the already described experimental setup [24]. The home self-made spectro-electrochemical cell body as well as all the parts are made of Teflon in order to be compatible with a broad variety of solvents. Viton O-rings are used where needed in order to ensure gas-tightness. The working electrode is a *c.a* 2 mm diameter disk of polished platinum inserted in a Teflon rod. It is mounted in the central well on a micrometer screw that permits fine adjustment of the distance between the electrode and the optical window, while maintaining the surface of the electrode parallel to the window. It was completed by a platinum wire as counter electrode and a silver wire used as a quasireference electrode. Its potential was checked vs. the ferrocene/ferrocenium couple before and after each experiment. Electrochemical measurements were carried out with a Biologic SP-150 potentiostat driven by the EC-Lab software including ohmic drop compensation. Spectrophotometric measurements were carried out in direct reflexing mode on the working electrode with a homemade bench composed of different Princeton Instruments modules (light sources, fibers, monochromators, spectroscopy camera and software) using a “Y-shaped” optical fiber bundle to connect the light source, the spectro-electrochemical cell and the spectrophotometer

5. Supplementary data

Synthesis, ^1H NMR and absorption spectra

6. Acknowledgements

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7. References

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Highlights

- A diarylethene joined to two benzooxazolidine by ethylenic spacers was synthesized
- Photochromic, acidochromic and electrochromic responses were evidenced
- The system is multistate (among the 18 possible states, 13 were underlined)
- The system is multifunctional (same interconversion performed by different stimuli)