



Cite this: *Phys. Chem. Chem. Phys.*, 2015, 17, 6066

Large protonation-gated photochromism of an OPE-embedded difurylperfluorocyclopentene†

Jannic Wolf,‡ Thomas Huhn* and Ulrich E. Steiner

A recently reported protolytic gating effect on the ring closing reaction of an oligo(phenylene ethynylene) (OPE) embedded difurylperfluorocyclopentene (**S**) with a dimethylaminophenyl chain link in each of the side arms, was quantitatively analyzed in detail. The reaction system (S_o , S_oH^+ , $S_oH_2^{2+}$, S_c , S_cH^+ , $S_cH_2^{2+}$) comprising three protolytic forms in both open and closed configuration, is characterized by four protolytic equilibrium constants and six photochemical quantum yields of ring closing and ring opening. The absorption spectra, conductivity, and effective photochemical quantum yields were measured in acetonitrile as functions of solvent acidity varied by addition of trifluoroacetic acid and triethylamine and characterized by an effective pH_{non-aq} . Based on the derivation of a rigorous method for assessing the individual quantum yields of ring closure and ring opening of the six species, it was shown that it is specifically the second protonation step that is responsible for a more than 10-fold increase in the quantum yield of ring closure.

Received 12th November 2014,
Accepted 23rd January 2015

DOI: 10.1039/c4cp05258g

www.rsc.org/pccp

Introduction

Molecules that can be reversibly switched between two well-defined states through optical activation are of special interest for molecular electronics applications. Diarylethenes (DAEs) are particularly suited for this task since they are thermally stable and highly fatigue resistant. During the switching process, DAEs with 5-membered heterocycles connected by the ethene-1,2-diyl-bridge in the 3- and 3'-position, respectively, are converted by a 6π -electrocyclic reaction between two states differing in the π -electron delocalization across the molecule. As borne out by the valence bond structures, in the open state, electronic communication from one substituent attached in 2-position of the aryl moiety with a substituent attached in the 2'-position of the other aryl moiety is not possible. In the closed state, the substituents attached in 2- and 2'-position of the aryl rings electronically communicate efficiently over the π -system that now spans the whole molecule (Scheme 1). In recent years,

efforts have been made to integrate these molecules into mechanically controlled break junction setups, for measuring the conductance difference between the two forms.^{2,3} It was shown that the conjugated ring-closed isomer of diarylperfluorocyclopentenes can exhibit up to 38 times higher conductance compared to its ring-open isomer.⁴ In a recent study aimed at a better understanding of the switching mechanism and the charge transport along these molecules, we presented a family of switchable molecular wires having electron withdrawing or electron donating side chains on long conjugated oligophenylene-ethynylene side arms (OPE).¹ We demonstrated that the electron density in these OPEs influences the efficiency of the photochromic process, as quantified by the quantum yields of ring opening and ring closing. One member of the family, equipped with strongly electron donating *N,N*-dimethylamino groups on the side arm, showed extensive reversible modulation of the quantum yield of ring closing upon protonation. On another note, molecular switches also can be used for a direct control of their environment by changes in their physical and chemical properties upon photoisomerization. After Lehn's seminal work on the photomodulation of a pK_a -value,⁵ Raymo and coworkers have systematically exploited the photomodulation of pH by spiropyranes to establish logical gates on a molecular basis.⁶⁻⁸ Furthermore, the control of catalytic activity⁹ or the switching of material properties in thin film devices^{10,11} have represented a main focus of investigations in the area of DAEs.

Gated photochromism (GP) in its pure definition denotes a situation where chemical or environmental changes of either of two interconvertible photochromic forms of a compound causes it to become non-photochromic.¹² Contrary to the IUPAC

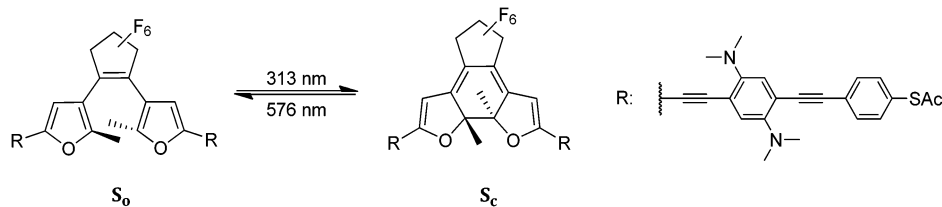
Fachbereich Chemie, Universität Konstanz, D-78457 Konstanz, Germany.

E-mail: thomas.huhn@uni-konstanz.de

† Electronic supplementary information (ESI) available: It refers to the topics: (1) system of equations for calculating the equilibrium concentrations of the species B, BH^+ , BH_2^{2+} , HA and A^- ; (2) absorbance/absorbance diagrams for the spectral change on titration of S_o by TFAH; (3) simulated spectra of the three protolytic forms; (4) defining a pH_{non-aq} in a non-aqueous aprotic solvent; (5) calculation of the fractions of the protolytic forms during forward and back titration; (6) derivations of the pH dependent quantum yields of individual protolytic forms. See DOI: 10.1039/c4cp05258g

‡ Present address: Solar & Photovoltaics Engineering Research Center, King Abdullah University of Science and Technology, Thuwal 23955-6900, Saudi Arabia.





Scheme 1 Reversible photochromism of the bisfurylperfluorocyclopentyl-derived molecular switch with a dibasic phenylenediamine group inserted in each of the conjugated OPE side arms. S_o (open form) and S_c (closed form).

definition, in practice GP often is not a strict on/off phenomenon because of the co-existence of various protolytic species in both photochromic forms. Kobatake *et al.* demonstrated that, upon protonation, an acid-sensitive DAE having a phenyldiethylamino side group could be switched from a thermally irreversible to a thermally reversible photochromic system.¹³ A similar effect for a DAE based on 2-ethynyl-pyridyl substituents was reported by Yumoto *et al.*¹⁴ Upon protonation of the pyridyl moiety, the DAE became unswitchable by visible light, and thus was locked in the ring closed isomer. This effect was not observed if the quaternized pyridyl moiety was connected by a non-conjugated alkyl linker, a fact that clearly demonstrates the long range influence of the remote protonation on the switching core if transmitted by a conjugated linker. The trend that modulation of the electron donating abilities of *para-N,N*-diethylamino-phenyl groups in the 5-position of the DAE significantly affects the switching behavior was first observed by Irie and Uchida.^{15,16} Strong electron donors such as the *para-N,N*-dimethylamino-phenyl group reduced the quantum yield of the ring closing reaction. However, after addition of trifluoroacetic acid (TFAH), resulting in the formation of a protonated species, which the authors assigned to the doubly protonated molecular switch, the protonated DAE smoothly underwent cyclization and cycloreversion. A separation of quantum yields of the mono- and di-protonated form was not attempted. The idea of a “multi-addressable” switching system was pursued by Liu and Chen¹⁷ who studied a dithiophenylethene (DTE) symmetrically conjugated to a 2-(8-hydroxyquinoline) substituent on each of both thiophene rings, thereby introducing at the same time two basic functions with the nitrogen heteroatoms as potential protonation sites and also two chelating functions in a cooperation of the nitrogens in the ring and neighboring hydroxyl groups. Both functions, when addressed, cause a moderate change of the ring closing quantum yield of the DTE unit. A more detailed analysis attempting to separate *e.g.* the effects of single and double protonation as well as a *pK*-determination was not pursued.

In a recent paper by Cipolloni *et al.*¹⁸ a benzobis(imidazol) bearing two aniline substituents and thus functioning as a four-basic chromophore was inserted between two DTE units. Combining the 5 protolytic stages with 3 ring-closing stages (open-open, open-closed, closed-closed) this system would bear the potential of realizing a 15-stage switch. Although the *pK*-values were not exactly determined, the sequence of protonations and the predominant form at various pH values were assigned. The average quantum yields of ring-closing and

opening determined as a function of pH exhibited a pronounced minimum at the point of double protonation. A precise separation of the photochemical quantum yields of all 5 protolytic forms was not feasible. Since the central benzobis(imidazol) chromophore is also fluorescent, the pH-dependent behavior was also observable in the emission quantum yields.¹⁹

More recently, Massaad *et al.* reported the first quantitative GP study of a 1,2-bis(5-carboxy-2-methylthien-3-yl)cyclopentene DAE.²⁰ In this case, the DAE was a dicarboxylic acid. Its quantum yield of ring closing was found to be slightly sensitive to the presence of tetrabutyl ammonium hydroxide, whereas its cycloreversion was not affected. On the other hand, a photo-modulation of the *pK_a* of more than one *pK_a*-unit was observed, the closed isomer being more acidic. The theoretical method of evaluation applied in this work, as well as in another interesting paper by Kutsunugi *et al.*²¹ on protolytic gating involving a bis(dibenzothienyl)ethene with the central double bond bridged by a dimethylaminophenyl thiamide function capable of double protonation at the amino group and the thiamide nitrogen, will be further referred to below in relation to our own work.

In this paper, we present a detailed quantitative study of the protonation-gated photochromic behavior of the long, conjugated molecular switch S_o in solution (Scheme 1). This compound is characterized by its extended π -system with the photoreactive chromophore embedded between two conjugated side arms comprising phenylenediamine groups in each side arm for efficient pH-gating. The overall length of *ca.* 3 nm in the closed state and the thiol endgroups render S_o a switchable molecular wire (SMW). Upon treatment of S_o with TFAH, the tetra-basic molecular switch considerably changes its photophysical behavior. A more than tenfold increase in the ring-closing quantum yield, accompanied by a high conversion rate at the photo stationary state (PSS) was observed in the presence of 70 equiv. of TFAH. The modulation is fully reversible upon addition of an equimolar amount of triethylamine (TEA). By modeling the coupled protolytic equilibria in the S_o - S_c -TFAH-TEA system on the basis of the concept of an effective pH_{non-aq} , we succeeded in assessing the relative fractions of mono- and di-protonated species as functions of the TFAH and TEA concentrations and in assigning individual quantum yields of photoswitching to the two protolytic species. Thereby, we clearly demonstrate that double protonation of the SMW is required to achieve the drastic quantum yield increases that are observed.



Experimental

All sample solutions were air-saturated and no special measures were taken to exclude moisture during the experiments. Trifluoroacetic acid (TFAH) from Sigma-Aldrich and spectroscopic grade MeCN from Merck were used as received. The synthesis of the investigated molecular switch is reported in ref. 1.

1. Continuous irradiation in solution

A high-pressure mercury lamp (Osram HBO 200 W/2) was used. The light was passed through a heat filter (optical length 5 cm, filled with a saturated aqueous solution of CuSO₄), a collimating lens and an electronic shutter. Irradiation experiments were performed in a setup with a 313 nm or 576 nm interference filter (Schott), a thermostated optical cell holder with a cell of 1 cm path length (Hellma QS) equipped with a magnetic stirring bar and filled with 3 mL of sample solution. At 313 nm, the photon irradiance I_0 was determined by azobenzene actinometry,^{22,23} at 576 nm by mesodiphenylhelianthrene actinometry.²⁴ UV-vis absorption spectra were recorded before and after irradiation on a Cary 50 UV-vis spectrophotometer. Quantum yields of the photoreactions were determined according to the method of Gauglitz.^{22,23,25}

2. Acid–base titration experiments with S_o

These experiments were carried out in Quartz cuvettes and monitored by UV-vis spectroscopy. In each experiment, a sample of the SMW in its open form in MeCN (3 mL, $c = 1.25 \times 10^{-5}$ M) was used. The titrations were carried out with a micropipette, and the cuvette was shaken after each addition. Stock solutions in MeCN of either TFAH (“St-TFAH”; $c = 3.75 \times 10^{-1}$ M) or TEA (“St-TEA”; $c = 3.75 \times 10^{-1}$ M) were used for titrations. The concentration was chosen such that 1 μ L of each stock solution equals 10 equivalents of either acid or base with respect to the content of S_o in the UV-cuvette. For the forward titration, St-TFAH was used. The back titration was carried out with St-TEA on a sample pre-treated with 7 μ L of St-TFAH. The basicity of the ring-closed isomer was estimated by adding 7 μ L of St-TFAH to the sample before it was irradiated at 313 nm to yield a PSS, where 97% of the switch is present in the closed form.²⁶ This solution was then back titrated with St-TEA in portions of 10 equivalents. For determining the pH_{non-aq} dependence of the quantum yields, pH_{non-aq} values were set by adding suitable amounts of St-TFAH to freshly prepared samples.

3. Conductometric test of protonation constants

Modeling the observed spectral changes during the titration with TFAH, indicated the formation of free ions. Conductometric titrations were performed in order to yield independent evidence of free-ion formation during the titration. A regime of higher concentrations of the SWM S_o (1.00×10^{-4} M in MeCN) compared to the UV-vis spectroscopic analyses was chosen in order to yield higher conductivity values matching the sensitivity range of the conduction bridge. The measurements were performed under ambient conditions at room temperature. The conductance of a stirred solution of SMW S_o in MeCN (20 mL)

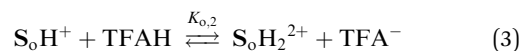
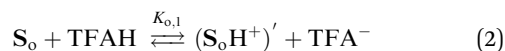
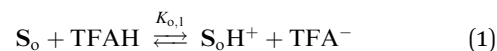
was monitored during the addition of aliquots of 10 equiv. of TFAH (53.3 μ L St-TFAH).

Results and discussion

1. Equilibrium constants of protonation

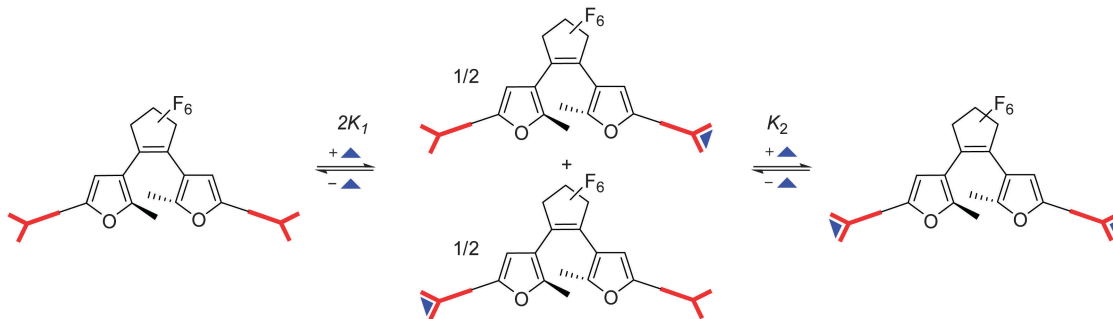
Details of the light induced switching between the non-conjugated open ring isomer S_o and the conjugated ring closed isomer S_c (cf. Scheme 1) have been reported in ref. 1. In summary, the absorption maximum of the ring open isomer in MeCN was found at 331 nm. Upon irradiation with 313 nm light, an absorption band in the visible developed with a maximum at 589 nm. However, the degree of conversion to the closed ring isomer in the PSS was rather low (46%), due to a rather small quantum yield ϕ_{o-c} (0.4%) for the ring closing process. Upon addition of 70 equiv. of TFAH to the solution, this quantum yield increased by more than a factor of 10 up to a value of 5% and conversion to the closed SMW in the PSS was almost complete (97%) under such conditions. The protonation dependent gating effect was found to be fully reversible upon addition of equimolar amounts of TEA. Such back titrated samples showed the same photophysical behavior as untreated samples, indicating that the furan rings of the molecular switch are stable under non-nucleophilic but acidic conditions. Furthermore, it was observed that the other molecular switches investigated in our previous study that did not have NMe₂ groups showed neither an enhancement of the ring closing quantum yield nor decomposition in the presence of TFAH.¹ Thus, it can be concluded that the behavior of the molecular switch S is uniquely controlled by protonation/deprotonation of the molecule at the NMe₂ groups in the side arms. NMR spectra of the protonated S_o and S_c are provided in the SI of ref. 1. The ring opening quantum yield ϕ_{c-o} was not affected by the presence of acid or base.¹

Two distinct sites sensitive to protonation are located at each phenylenediamine in the side arms of S_o. In view of the Coulomb repulsion energy, it can be assumed that a second protonation will be unlikely to occur at the second NMe₂ group at the same site. Instead, the phenylenediamine of the other side arm will be protonated. Consequently, the following protonation scheme was adopted, where TFAH stands for the undissociated trifluoroacetic acid and TFA⁻ for its anion:



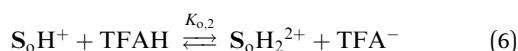
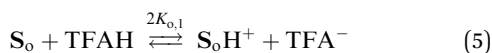
The symbols S_oH⁺ and (S_oH⁺)' in eqn (1) and (2) denote the two forms with single protonation at the left or right OPE arm, respectively. Since both sides of the molecule are spectroscopically and chemically identical, these two equilibria can





Scheme 2 Cartoon illustrating the protonation processes. The red “Y” represents the dibasic phenylenediamine and the blue triangle a proton.

be treated as one but with double statistical weight. Thus, the equations for the protonation equilibria can be reduced to



The cartoon shown in Scheme 2 illustrates the situation.

The effect of protonation is clearly represented by a change of the absorption spectra during titration with TFAH (Fig. 1). The spectral change is characterized by several isosbestic points and almost linear relationships between the absorbance values at different wavelengths (*cf.* ESI† Fig. S1), which generally is taken as an indication that there are only two different forms involved in the observed conversion. In the following, we describe the evaluation of the equilibrium constants for the conversion between three protolytic forms S_o , S_oH^+ and $S_oH_2^{2+}$ and the assignment of individual spectra to these species. Actually, the apparent bimodal character of the spectral change can be accounted for by the fact that the spectrum of S_oH^+ as such very closely corresponds to a linear combination of the spectra of S_o and $S_oH_2^{2+}$ (*cf.* Fig. S2, ESI†).

Applying the law of mass action to eqn (5) and (6) and taking into account the stoichiometric relations in the titrated reaction

mixture, a system of eqn (S1)–(S5) (ESI†) was obtained. It was solved with the help of a *Mathematica* computer program²⁷ for various trial values of K_1 and K_2 , to yield the equilibrium concentrations of TFAH, TFA^- , S_o , S_oH^+ and $S_oH_2^{2+}$. Two representative examples of the calculated equilibrium compositions as a function of the amount of acid are shown in Fig. 2.

Values for K_1 and K_2 were assumed to be equal since it is plausible that the first protonation in one half of the molecule does not significantly influence the second protonation step in the other half. In any case, for similar values of K_1 and K_2 it would be difficult to assess their difference from the limited information at hand.²⁸

Each measured spectrum $A(\lambda, [\text{TFAH}]_0)$ during the stepwise titration of S_o with TFAH (Fig. 1) represents the superposition of the weighted individual spectra of the three protolytic forms S_o , S_oH^+ and $S_oH_2^{2+}$ at this specific point of the titration:

$$A(\lambda, [\text{TFAH}]_0) = A_{S_o}(\lambda) \frac{[S_o]([\text{TFAH}]_0)}{[S_o]_{\text{tot}}} + A_{S_oH^+}(\lambda) \frac{[S_oH^+]([\text{TFAH}]_0)}{[S_o]_{\text{tot}}} + A_{S_oH_2^{2+}}(\lambda) \frac{[S_oH_2^{2+}]([\text{TFAH}]_0)}{[S_o]_{\text{tot}}} \quad (7)$$

The concentrations of the various protolytic forms $[S_oH_n^{n+}]([\text{TFAH}]_0)$ are functions of the total concentration $[\text{TFAH}]_0$ of the acid. The wavelength dependent coefficients $A_{S_o}(\lambda)$, $A_{S_oH^+}(\lambda)$ and $A_{S_oH_2^{2+}}(\lambda)$ represent the individual spectra of the $[S_oH_n^{n+}]$ if present in the total concentration $[S_o]_{\text{tot}}$.

Based on the calculated concentrations of the three protolytic forms in dependence of $[\text{TFAH}]_0$ for certain values of $K_{o,1}$ and $K_{o,2}$, the measured spectra were used to fit the three absorbances A_i for each wavelength. The $[\text{TFAH}]_0$ dependence of the measured spectra can be fitted well with the described procedure such that the differences between measured and simulated spectra were always found below 1%. However, a rather wide range of preset trial values for $K_{o,1} = K_{o,2}$ gave equally good results for the fit. Therefore, the quality of the fit does not represent a sufficient criterion to determine the most realistic values for $K_{o,1} = K_{o,2}$. In Fig. S2 (ESI†), four representative examples of simulated spectra of the individual protolytic forms for different K -values are shown. For higher values of

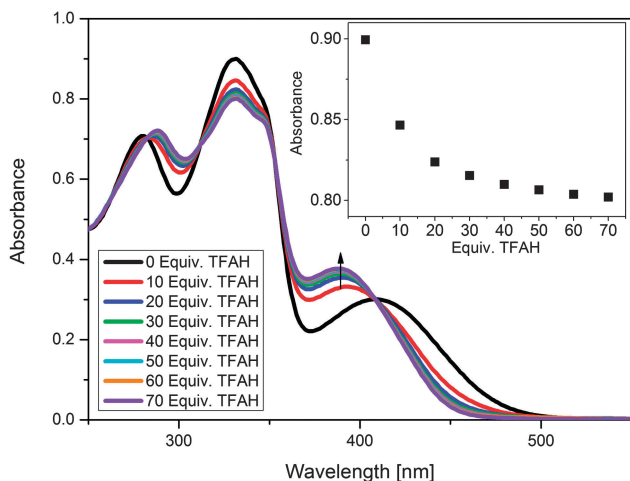


Fig. 1 UV-vis spectra recorded during the titration of S_o (3 mL, 1.25×10^{-5} M, MeCN) with TFAH (3.75×10^{-1} M, MeCN). 10 equiv. aliquots TFAH (1 μ l) were used for each step. The inset shows the volume corrected titration curve extracted at 330 nm.



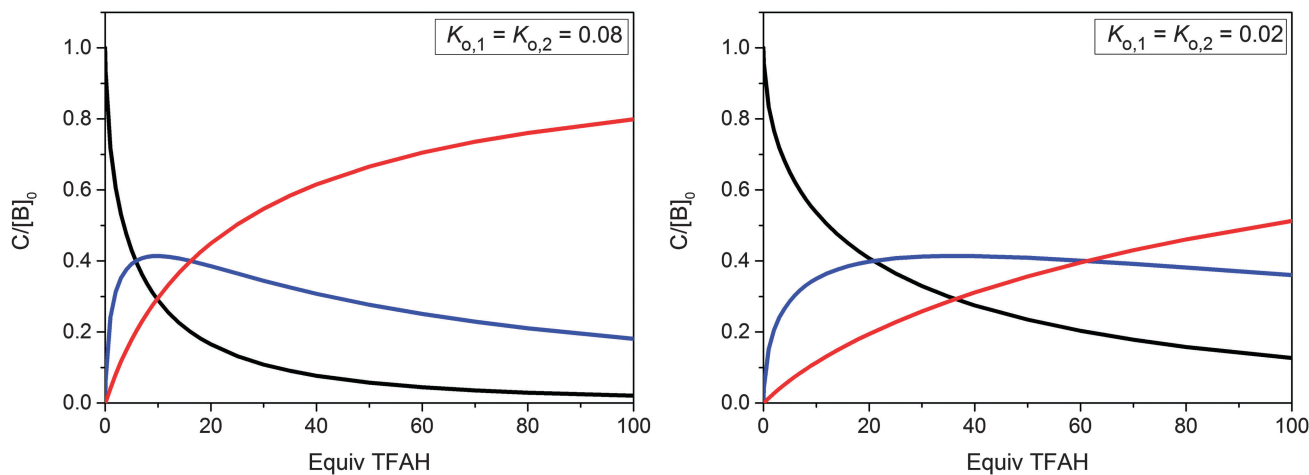


Fig. 2 Calculated equilibrium concentrations of S_o (black), S_oH^+ (blue) and $S_oH_2^{2+}$ (red) as functions of added acid ($[TFAH]_0$) for two representative values of $K_{o,1} = K_{o,2}$. Concentrations are given in units of $[S_o]_{tot} = 1.25 \times 10^{-5}$ M. The horizontal scale represents the number of added TFAH equivalents.

$K_{o,1} = K_{o,2}$, the S_oH^+ spectrum is closer to the S_o spectrum, whereas for lower values of $K_1 = K_2$, the S_oH^+ spectrum is closer to the $S_oH_2^{2+}$ spectrum. If it is assumed that the spectrum of the mono protonated form should be in the middle between the spectra of S_o and $S_oH_2^{2+}$, one would expect the values of K_1 and K_2 to range between 0.06 and 0.1. This conclusion is confirmed by conductometric titration experiments and by the analysis of the acidity dependence of the photochemical quantum yield (*vide infra*).

2. Conductometric validation of protonation constant

Neat MeCN showed a specific conductivity of $0.3 \mu S cm^{-1}$. Adding TFAH to neat MeCN did not result in an increase of conductivity, indicating no dissociation of this acid in this solvent. The specific conductivity of a 10^{-4} M solution of S_o in dry MeCN was determined to be $2.1 \mu S cm^{-1}$. Upon the addition of TFAH, the specific conductivity increased asymptotically to reach saturation at $24.3 \mu S cm^{-1}$ (7 added aliquots, Fig. 3). The observed saturation in the measurement of conductivity correlates with the saturation observed in the UV-vis spectra during titration. Under the prevailing experimental conditions, the charge carriers are represented by the ions S_oH^+ , $S_oH_2^{2+}$ and TFA^- . According to eqn (5) and (6), protonation of S_o or S_oH^+ yields the equivalent amount of anions TFA^- assuming that, according to their charges, the molar conductances of S_oH^+ , TFA^- and $S_oH_2^{2+}$ (TFA^-)₂ are in a ratio of 1:2, the specific conductance κ of the solution should be described by eqn (8)

$$\kappa = \kappa_0 + \lambda[S_oH^+] + 2\lambda[S_oH_2^{2+}] \quad (8)$$

where κ_0 is the initial value of the specific conductivity and λ is the molar conductivity of the salt $S_oH^+ TFA^-$. The conductivity of a solution of S_o before addition of TFAH indicates that a small fraction of the sample is already protonated as a remnant of its preparation method. In Fig. 3, the observed variation of κ as a function of equivalents of TFAH is shown together with theoretical curves for κ according to eqn (8) using various

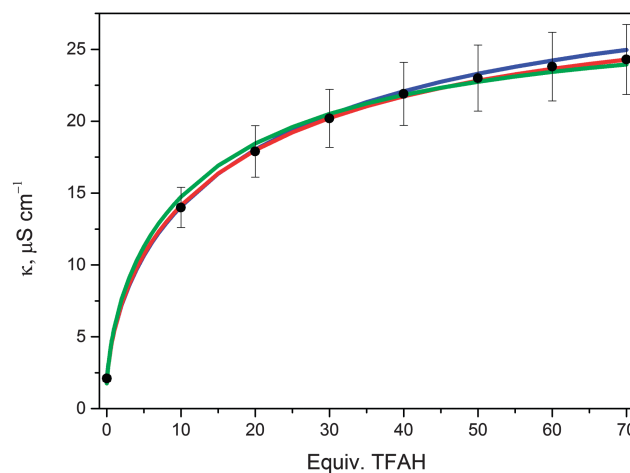


Fig. 3 Conductometric titration curve of a 1.0×10^{-4} M sample solution of S_o in MeCN. Black points: experimental data with error bars; colored curves: simulations according to eqn (8) assuming $K_{o,1} = K_{o,2} = 0.05$ and $\lambda = 155 \Omega^{-1} cm^2 mol^{-1}$ (blue curve), $K_{o,1} = K_{o,2} = 0.06$ and $\lambda = 146 \Omega^{-1} cm^2 mol^{-1}$ (red curve), $K_{o,1} = K_{o,2} = 0.08$ and $\lambda = 138 \Omega^{-1} cm^2 mol^{-1}$ (green curve).

values of $K_{o,1} = K_{o,2}$ together with pertinent best fit values of λ . The diagram shows that $K_{o,1}$ values in the range of 0.5 and 0.08 fit the observed conductivity curve very well. The assumed molar conductivities of about $140 \Omega^{-1} cm^2 mol^{-1}$ are of an order of magnitude perfectly reasonable for ions of the present size in MeCN.²⁹

These experiments confirm that ion pairs between the protonated compound S_oH^+ and the anion TFA^- in MeCN dissociate rather completely. The same is probably true when the dimethylaminophenylthioamide modified DAE described by Kutsunugi *et al.*²¹ reacts with TFAH in MeCN. This issue was under discussion in that work.

3. Equilibrium constants of protonation of the closed form S_c

Since the electronic structure of the closed-ring isomer S_c differs from its ring-open counterpart S_o , it can be expected



that both forms show different basicity. The basicity of the open form S_o is characterized by the equilibrium constants $K_{o,1}$ and $K_{o,2}$ given in the previous sections. The ring closed isomer S_c is not readily prepared by irradiation in the UV, due to the too low quantum yield of ring closure of S_o . But in previous experiments, it has been found that S_o undergoes photochemical ring closure much more efficiently in the protonated state. Furthermore, the PSS for the protonated form lies almost exclusively on the side of the closed form (97%), indicating that the quantum yield of ring closure strongly increases on protonation,¹ without a concomitant increase of the quantum yield of the reverse photoreaction.

To determine the protolytic equilibria of S_c , in principle one can start with the photochemically converted acidified solution and then add a base such as TEA. However, in such mixtures the acid base equilibria of S_c corresponding to those of S_o described by eqn (1)–(4) are coupled to further equilibria between TFAH and TEA, which turned out not to be simple since they also involve non-dissociated ion pairs. In order to avoid the complication of having to unravel these additional equilibria, it was useful to apply the concept of pH in non-aqueous solutions which we here define as (*cf.* ESI[†])

$$\text{pH}_{\text{non-aq}} = -\log \frac{[\text{TFAH}]}{[\text{TFA}^-]} \quad (9)$$

As detailed in the ESI,[†] the degree of protonation of S_o as well as of S_c is uniquely determined by the value of $\text{pH}_{\text{non-aq}}$. But the converse is also true: the value of $\text{pH}_{\text{non-aq}}$ is a unique function of the degree of protonation of the two forms of S . Hence our strategy to determine the $\text{pH}_{\text{non-aq}}$ in the mixed acid–base solutions, was as follows: forward titration of a solution of S_o by TFAH, followed by back titration with TEA. From the forward titration, we can assess the value of the total absorbance at a given total concentration $[\text{TFAH}]_0$ of the acid, for which we know the $\text{pH}_{\text{non-aq}}$ according to our previous analysis of the protonation equilibria of S_o according to eqn (1)–(4). Then, from the total absorbance of the mixed protolytic forms of S_o measured during the back titration, we can assess the $\text{pH}_{\text{non-aq}}$ values as a function of total concentrations of TFAH and TEA. In the next experiment, we follow the same procedure, but after the end of the forward titration, irradiate the solution to the PSS, yielding the closed form almost quantitatively. During the subsequent back titration we determine the absorbances of the mixtures of the protolytic forms of S_c as a function of $\text{pH}_{\text{non-aq}}$ which now is known from the first titration run carried out exclusively with the open form.

The spectra obtained during back titration of S_o pretreated with 70 equiv. TFAH are depicted in Fig. 4.

As evident from the inset of Fig. 4, the back titration of the acidified samples of S_o with TEA in MeCN does not simply yield the reverse curve of the forward titration with TFAH. This is due to the fact that not only the concentration of non-dissociated TFAH is diminished, but also free TFA^- anions are created. Thus, there is a twofold effect in the law-of-mass-action according to eqn (1)–(4): (i) through the decrease of $[\text{TFAH}]$ in the denominator, and (ii) through the increase of $[\text{TFA}^-]$ in the

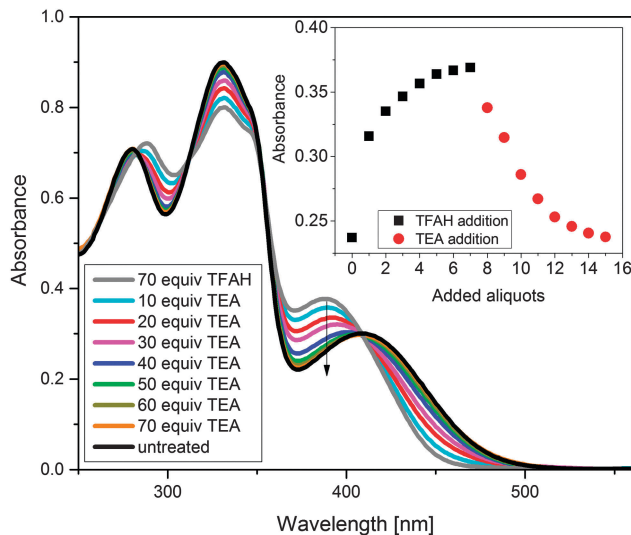


Fig. 4 Spectral series of the back titration of an acidic sample (70 equiv. TFAH) of S_o (3 mL, 1.25×10^{-5} M in MeCN) with 10 equiv. aliquots (1 μL) of TEA (3.75×10^{-1} M, MeCN). The inset shows a series of absorption values extracted at 374 nm for the forward and back titration of S_o with 10 equiv. aliquots of TFAH and TEA, respectively.

nominator. The change of $\text{pH}_{\text{non-aq}}$ during the back titration, obtained in the way described above, is shown in Fig. 5. A reference value of 0.08 was adopted for $K_{o,1} = K_{o,2}$ when calculating the concentrations of TFAH and TFA^- needed for $\text{pH}_{\text{non-aq}}$ determination according to eqn (9), because this value of $K_{o,1} = K_{o,2}$ is in good agreement with all three criteria: spectral middle argument for $S_o\text{H}^+$, conductivity, and quantum yield (*cf.* below).

Taking full advantage of the assignment of a $\text{pH}_{\text{non-aq}}$ value to any of the titration stages, the fractions of the protolytic forms, $S_c\text{H}^+$ and $S_c\text{H}_2^{2+}$ during each stage of the titration can be obtained solely from the pertinent protonation constants and the $\text{pH}_{\text{non-aq}}$ value (*cf.* ESI[†]). As in the case of the open form, the equilibrium constants for the first and second protonation by TFAH are treated as equal. Likewise, the statistical factor of 2 is included in the definition of $K_{1,c}$. The equilibrium constants $K_{1,c} = K_{2,c}$ have been determined in the same manner as shown for the open form. The fractions of the protolytic forms have been calculated according to eqn (S15)–(S17) given in the ESI.[†] Using these fractions, the absorbance at any wavelength is represented as given in eqn (10):

$$A_c(\lambda, \text{pH}_{\text{non-aq}}) = A_{S_c}(\lambda)X(\text{pH}_{\text{non-aq}}) + A_{S_c\text{H}^+}(\lambda)Y(\text{pH}_{\text{non-aq}}) + A_{S_c\text{H}_2^{2+}}(\lambda)Z(\text{pH}_{\text{non-aq}}) \quad (10)$$

where the coefficients A_i are the specific absorbances referring to the total concentration and X , Y , Z the fractions of the individual protolytic forms. Trial values for $K_{1,c} = K_{2,c}$ were assumed and the $A_{i,\text{closed}}$ values were fitted against the titration data shown in Fig. 6. Similar to the fitting procedure of the open form, the quality of the fit did not represent a sufficiently sharp criterion to determine the proper K -values unequivocally since a wide range of equilibrium constants gives equally good



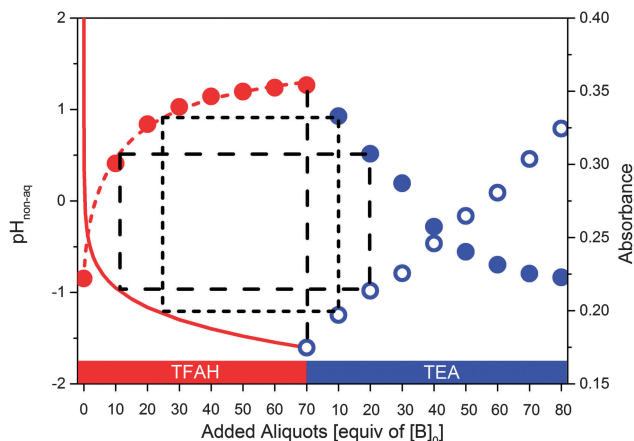


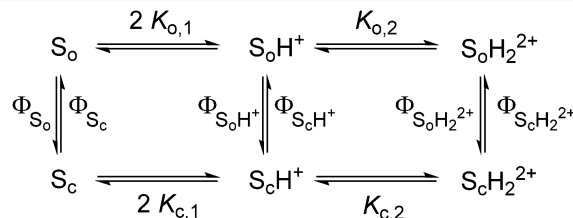
Fig. 5 Variation of $\text{pH}_{\text{non-aq}}$ during forward (red line) and back (blue open circles) titration. During the forward titration, $\text{pH}_{\text{non-aq}}$ was calculated according to the definition in eqn (9) using the values of $[\text{TFAH}]$ and $[\text{TFA}^-]$ resulting from calculations of the equilibria (1)–(4) with $K_{\text{o},1} = K_{\text{o},2} = 0.08$. The filled data points indicate the absorbance at 374 nm during forward (red) and back titration (blue). The rectangles demonstrate the procedure to convert the absorbance values during back titration into $\text{pH}_{\text{non-aq}}$ values. For details cf. text.

fits. Adopting again the spectroscopic argument that at each wavelength the absorbance of S_cH^+ should be intermediate between the absorbance of the two other forms (S_c and $\text{S}_c\text{H}_2^{2+}$), the best fit was found to be $K_{1,c} = K_{2,c} = 0.043$. Thus, within the present accuracy of determining the K -values, the basicity of the ring-closed isomer does not significantly deviate from that of the ring-open isomer.

4. pH dependent quantum yields

Depending on $\text{pH}_{\text{non-aq}}$, variable amounts of the three protolytic forms S_o , S_oH^+ , and $\text{S}_o\text{H}_2^{2+}$ of **S** are present in the sample solution. For each of them, a forward and backward photoisomerization with a distinct quantum yield is possible. Isomer S_c is involved in protolytic equilibria, too, with its protolytic forms being denoted S_c , S_cH^+ , and $\text{S}_c\text{H}_2^{2+}$. Consequently, there are four equilibrium constants and six individual quantum yields to be considered for a comprehensive description of the reactions in the sample solution (Scheme 3). Analyzing the photochemical kinetics of ring opening and closing at different $\text{pH}_{\text{non-aq}}$, it is possible to single out the six individual quantum yields and to distinguish the protolytic gating effect of the first and second stage of protonation. In general, when it comes to the determination of photochemical parameters from stationary photokinetics, particularly in the case of photo-reversible reactions, there are two basic strategic philosophies: (i) one can set up all relevant rate equations, integrate them numerically and fit the unknown parameters to the observed kinetic curves (cf. e.g. ref. 21 and 30), or (ii) one can aim for an analytical solution as worked out by Gauglitz and coworkers.^{22,23} Simplifications can be achieved, by irradiating at isosbestic points and by exploiting directly the slopes of kinetic curves

(cf. e.g. ref. 18), although the slope-based methods are less accurate, since they use fewer kinetic data points than when using a time-integrated simulation. Compared to numerical integration methods, the analytical integration method appears to be more general and more transparent and therefore will be the method of choice in our approach.



Scheme 3 Reaction scheme of coupled protolytic and photochemical conversions with assignment of equilibrium constants $K_{\text{o},1}$, $K_{\text{o},2}$, $K_{\text{c},1}$, $K_{\text{c},2}$ and quantum yields Φ_{S_o} , $\Phi_{\text{S}_o\text{H}^+}$, $\Phi_{\text{S}_o\text{H}_2^{2+}}$ for the open and Φ_{S_c} , $\Phi_{\text{S}_c\text{H}^+}$, and $\Phi_{\text{S}_c\text{H}_2^{2+}}$ for the closed forms.

The time-dependence of the total concentration of all three open forms $[\text{S}_o]_{\text{tot}}$ is described by eqn (11).³¹

$$\begin{aligned}
 \frac{d[\text{S}_o]_{\text{tot}}}{dt} = I_0 \frac{F}{V} (1 - 10^{-A}) & \left\{ -\Phi_{\text{S}_o} \frac{\varepsilon_{\text{S}_o} \ell [\text{S}_o]}{A} - \Phi_{\text{S}_o\text{H}^+} \frac{\varepsilon_{\text{S}_o\text{H}^+} \ell [\text{S}_o\text{H}^+]}{A} \right. \\
 & - \Phi_{\text{S}_o\text{H}_2^{2+}} \frac{\varepsilon_{\text{S}_o\text{H}_2^{2+}} \ell [\text{S}_o\text{H}_2^{2+}]}{A} + \Phi_{\text{S}_c} \frac{\varepsilon_{\text{S}_c} \ell [\text{S}_c]}{A} \\
 & \left. + \Phi_{\text{S}_c\text{H}^+} \frac{\varepsilon_{\text{S}_c\text{H}^+} \ell [\text{S}_c\text{H}^+]}{A} + \Phi_{\text{S}_c\text{H}_2^{2+}} \frac{\varepsilon_{\text{S}_c\text{H}_2^{2+}} \ell [\text{S}_c\text{H}_2^{2+}]}{A} \right\} \quad (11)
 \end{aligned}$$

Here I_0 denotes the photon flux density, F the illuminated area of the cuvette, V the (stirred) volume of the sample solution, ℓ the optical path length, A the total absorbance at the wavelength of irradiation and the various ε parameters are the molar absorption coefficients of the respective open and closed species. As shown in Section 6 of the ESI† (eqn (S20)–(S28)), eqn (11) can be reduced to the form

$$\frac{d[\text{S}_o]_{\text{tot}}}{dt} = -F_{\text{pk}}^{-1} I_0 \frac{F}{V} \left\{ Q[\text{S}_o]_{\text{tot}} - \bar{Q}[\text{S}]_{\text{tot}} \right\} \quad (12)$$

where the photokinetic factor F_{pk} is given by

$$F_{\text{pk}} = \frac{A}{1 - 10^{-A}} \quad (13)$$

Q is the pseudo quantum yield, comprising forward and backward reaction

$$Q = \vec{Q} + \bar{Q} \quad (14)$$

with³²

$$\vec{Q} = \Phi_{\text{S}_o} A_{\text{S}_o} X_{\text{S}_o} + \Phi_{\text{S}_o\text{H}^+} A_{\text{S}_o\text{H}^+} Y_{\text{S}_o\text{H}^+} + \Phi_{\text{S}_o\text{H}_2^{2+}} A_{\text{S}_o\text{H}_2^{2+}} Z_{\text{S}_o\text{H}_2^{2+}} \quad (15)$$

$$\begin{aligned}
 \bar{Q} = & \Phi_{\text{S}_c} A_{\text{S}_c} X_{\text{S}_c} + \Phi_{\text{S}_c\text{H}^+} A_{\text{S}_c\text{H}^+} Y_{\text{S}_c\text{H}^+} \\
 & + \Phi_{\text{S}_c\text{H}_2^{2+}} A_{\text{S}_c\text{H}_2^{2+}} Z_{\text{S}_c\text{H}_2^{2+}} \quad (16)
 \end{aligned}$$



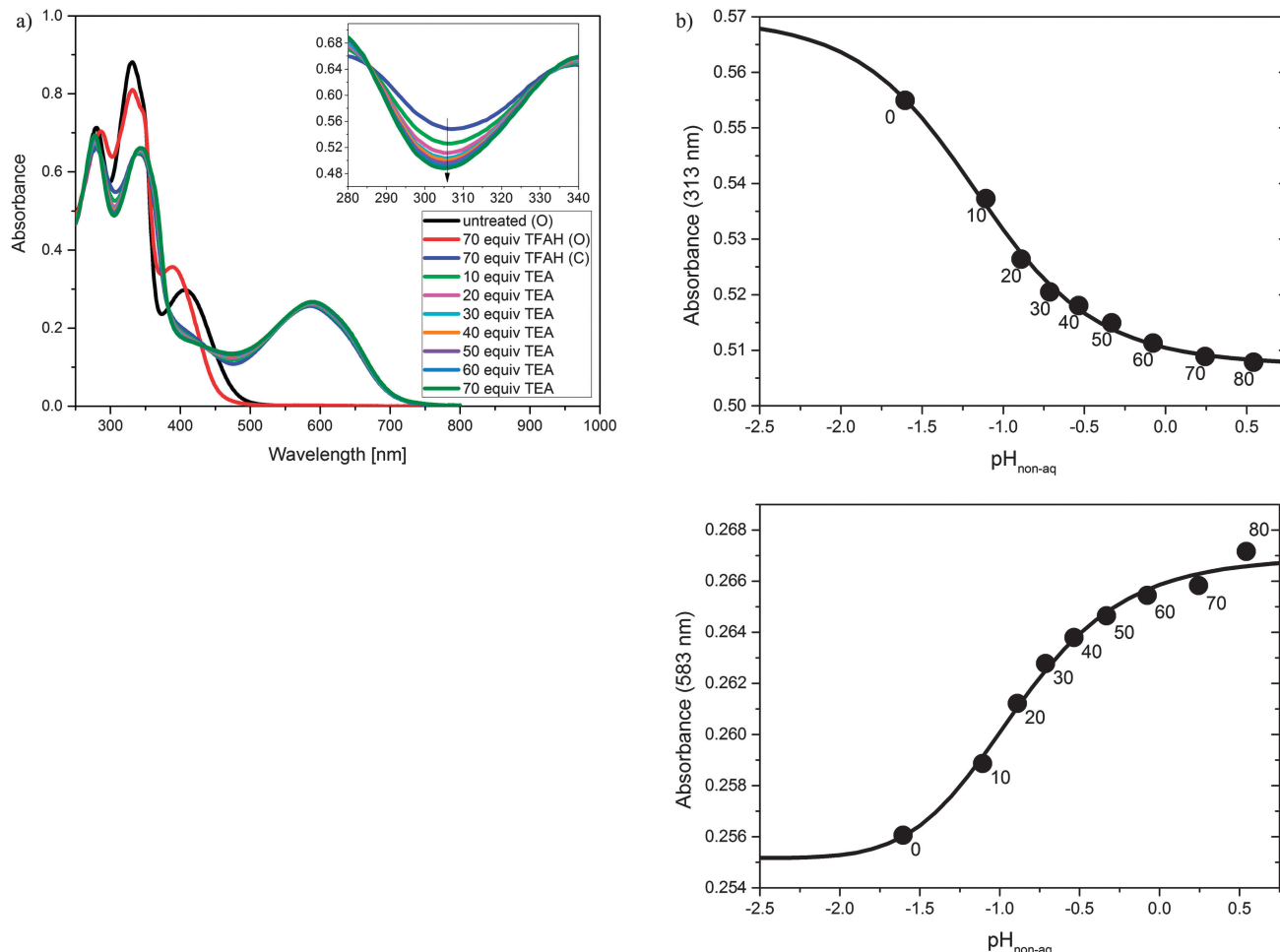


Fig. 6 (a) Absorption spectra during back titration of S_c with TEA after its photochemical generation from S_o in a solution with 70 equivalents of TFAH. The inset shows the spectral region used for the analysis of the protonation equilibria. (b) The upper panel shows the $\text{pH}_{\text{non-aq}}$ dependence of the absorbance of the closed form at 313 nm. For $K_{1,c} = K_{2,c} = 0.043$, the best fit values of the specific absorbances of S_c , S_cH^+ and $S_cH_2^{2+}$ obeying the criterion that $A_{S_cH^+}$ should be intermediate between A_{S_c} and $A_{S_cH_2^{2+}}$ are $A_{S_c} = 0.509$, $A_{S_cH^+} = 0.541$ and $A_{S_cH_2^{2+}} = 0.571$. The lower panel depicts the $\text{pH}_{\text{non-aq}}$ dependence at 583 nm with the fit of $A_{S_c} = 0.266$, $A_{S_cH^+} = 0.255$ and $A_{S_cH_2^{2+}} = 0.255$ calculated assuming the same $K_{1,c} = K_{2,c} = 0.043$ as in the upper panel. In this case we are close to an isosbestic point of all three forms.

involving the definitions

$$X_{S_o} = \frac{[S_o]}{[S_o]_{\text{tot}}} \quad Y_{S_oH^+} = \frac{[S_oH^+]}{[S_o]_{\text{tot}}} \quad Z_{S_oH_2^{2+}} = \frac{[S_oH_2^{2+}]}{[S_o]_{\text{tot}}} \quad (17)$$

$$X_{S_c} = \frac{[S_c]}{[S_c]_{\text{tot}}} \quad Y_{S_cH^+} = \frac{[S_cH^+]}{[S_c]_{\text{tot}}} \quad Z_{S_cH_2^{2+}} = \frac{[S_cH_2^{2+}]}{[S_c]_{\text{tot}}} \quad (18)$$

$$A_{S_o} = \varepsilon_{S_o} \ell [S_o]_{\text{tot}} \quad A_{S_oH^+} = \varepsilon_{S_oH^+} \ell [S_o]_{\text{tot}} \quad A_{S_oH_2^{2+}} = \varepsilon_{S_oH_2^{2+}} \ell [S_o]_{\text{tot}} \quad (19)$$

$$A_{S_c} = \varepsilon_{S_c} \ell [S_c]_{\text{tot}} \quad A_{S_cH^+} = \varepsilon_{S_cH^+} \ell [S_c]_{\text{tot}} \quad A_{S_cH_2^{2+}} = \varepsilon_{S_cH_2^{2+}} \ell [S_c]_{\text{tot}} \quad (20)$$

$[S_o]_{\text{tot}}$ and $[S_c]_{\text{tot}}$ denote the sum of concentrations of all protolytic forms of S_o and S_c , respectively.

It should be noted that unless the absorption coefficients of the reacting species involved are all equal, the use of ‘‘average

quantum yields’’ (*i.e.* omitting the absorption coefficients in eqn (15) and (16)) in photokinetic equations, is not exact.³²

From eqn (12) it can be deduced that in the photostationary (PSS) state ($d[S_o]/dt = 0$) the following equation holds:

$$\bar{Q}[S]_{\text{tot}} = Q[S_o]_{\text{tot,pss}} \quad (21)$$

and similarly, combining eqn (21) with eqn (14):

$$\bar{Q}[S]_{\text{tot}} = Q[S_c]_{\text{tot,pss}} \quad (22)$$

Using eqn (21), eqn (12) can be rewritten as

$$\frac{d[S_o]_{\text{tot}}}{dt} = -F_{\text{pk}}^{-1} I_0 \frac{F}{V} Q \{ [S_o]_{\text{tot}} - [S_o]_{\text{tot,pss}} \} \quad (23)$$

Since the total concentration of the open form S_o (as well as of the closed form) is linearly related to the absorbance A at any wavelength, eqn (23) is also valid in the form

$$\frac{dA}{dt} = -F_{\text{pk}}^{-1} I_0 \frac{F}{V} Q \{ A - A_{\text{pss}} \} \quad (24)$$



Of course, the observation wavelength should be best chosen far from an isosbestic point, in order to render changes of A relatively high.

This form of the photokinetic rate law corresponds exactly to that one which the approximate analytical method by Gauglitz,^{22,23} employing piecewise linearization of the photokinetic factor, is based on. Thus it is straight forward to measure the pseudo quantum yield Q for any pH of the solution. To decompose Q into the partial pseudo quantum yields \bar{Q} and \bar{Q} , one has to determine the composition of the photostationary state, which can be done by any convenient analytical method (*cf.* below). According to eqn (15) and (16), each of the partial pseudo quantum yields is a linear combination of the individual quantum yields of the pertinent protolytic open or closed forms.³² Since the coefficients $A_{S_o}X_{S_o}$, $A_{S_oH^+}Y_{S_oH^+}$, $A_{S_oH_2^{2+}}Z_{S_oH_2^{2+}}$ and $A_{S_c}X_{S_c}$, $A_{S_cH^+}Y_{S_cH^+}$, $A_{S_cH_2^{2+}}Z_{S_cH_2^{2+}}$ are known for any $\text{pH}_{\text{non-aq}}$, the individual quantum yields Φ_{S_o} , $\Phi_{S_oH^+}$, $\Phi_{S_oH_2^{2+}}$ and Φ_{S_c} , $\Phi_{S_cH^+}$, $\Phi_{S_cH_2^{2+}}$ can be obtained by fitting the observed values for \bar{Q} and \bar{Q} , as a function of $\text{pH}_{\text{non-aq}}$ according to eqn (15) and (16).

The data entering the fit as non-variable parameters is summarized in Table S1 (ESI[†]). Specifically, the values of $\text{pH}_{\text{non-aq}}$ were calculated according to eqn (9) with various trial values of the equilibrium constants for the open-ring isomer, assuming again $K_{o,1} = K_{o,2}$ (*cf.* ESI[†]). The value of Q was derived from the photokinetics observed under irradiation at 313 nm in 1.25×10^{-5} M solutions of S_o in the presence of various amounts of TFAH as described in the experimental part. The PSS composition of each sample was derived from its $A_{583,\text{pss}}$ value with the help of a reference sample analyzed by ¹H-NMR, where $A_{583,\text{pss}} = 0.273$ correlated to 97% conversion to the closed form in the PSS. The variation of the absorbance of the three individual protolytic forms at 583 nm ($A_1 = 0.266$, $A_2 = 0.256$, $A_3 = 0.255$, obtained from the fit shown in Fig. 6b lower panel) was very small, hence a $\text{pH}_{\text{non-aq}}$ dependence of the reference value was considered negligible.

As shown in detail in the ESI,[†] a consistent fit of the partial quantum yield \bar{Q} of the ring closing reaction as a function of $\text{pH}_{\text{non-aq}}$ fixes the acceptable value of $K_{o,1} = K_{o,2}$ to a rather narrow range around 0.08, which is in compliance with the conductometric results and places the spectrum of S_oH^+ right in the middle between those of S_o and $S_oH_2^{2+}$. From the fit shown in Fig. 7, the individual quantum yields $\Phi_{S_o} = 0.0037$, $\Phi_{S_oH^+} < 0.01$ and $\Phi_{S_oH_2^{2+}} = 0.11$ were obtained. A more precise value of the quantum yield after single protonation cannot be given, because the best fit is obtained if we neglect its contribution altogether, or even taken it slightly negative (*cf.* Fig. S3, ESI[†]). In Fig. 7, we also show the contribution of form S_oH^+ , that one would obtain for $\Phi_{S_oH^+} = 0.01$. The deviation from the data points is clearly outside the uncertainty of the experimental data, thus fixing an upper bound to $\Phi_{S_oH^+}$. Probably its real value is close to that of the unprotonated S_o . In any case, however, from the fit, there is unequivocal evidence that the overall rise of \bar{Q} in acidified solutions is dominated by the diprotonated form which exhibits an increase of the ring closing quantum yield by at least a factor of more than 10 relative to the single protonated and

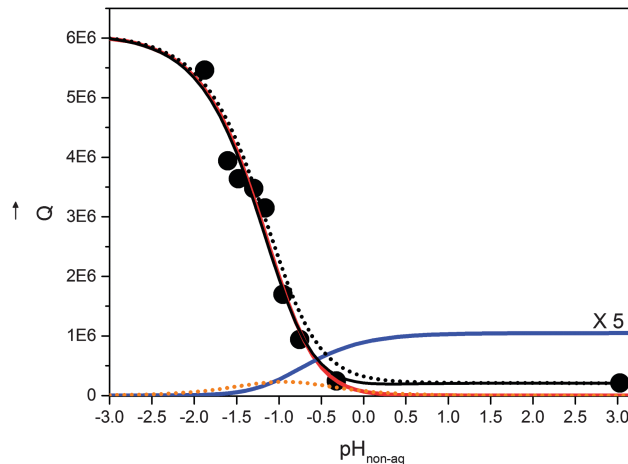


Fig. 7 Partial pseudo quantum yield \bar{Q} as a function of $\text{pH}_{\text{non-aq}}$. Experimentally observed: black data points, simulation: black solid line. The individual contributions of the various protolytic forms of S_o to the simulated value of \bar{Q} are: $\Phi_{S_o}A_{S_o}X_{S_o}$ (blue, for better visibility multiplied by a factor of 5), $\Phi_{S_oH_2^{2+}}A_{S_oH_2^{2+}}Z_{S_oH_2^{2+}}$ (red), the latter dominating the overall value of \bar{Q} . In calculating the solid black curve for the total value of \bar{Q} , the contribution of the intermediate form (S_oH^+) was completely neglected ($\Phi_{S_oH^+} = 0$). On the other hand, the dotted orange line represents the contribution of S_oH^+ assuming $\Phi_{S_oH^+} = 0.01$. The black dotted curve, representing the overall \bar{Q} in that case, is clearly outside the experimental error limits. For details, *cf.* ESI,[†] Fig. S3.

almost 30 relative to the unprotonated form. So far, to the best of our knowledge, comparably strong protolytic gating effects have not been reported in the literature.²⁰

In a similar procedure, the $\text{pH}_{\text{non-aq}}$ dependence of \bar{Q} at 313 nm irradiation wavelength was fitted using eqn (S27) (ESI[†]). The result of the fitting procedure is shown in Fig. S4 (ESI[†]). The individual quantum yields for the back irradiation from the closed to the open form under irradiation with 313 nm light were determined to be: $\Phi_{S_c} = 0.007$, $\Phi_{S_cH^+} = 0.001$ and $\Phi_{S_cH_2^{2+}} = 0.004$ with error margins of $\pm 30\%$ due to the large relative scatter of the data (*cf.* ESI,[†] Fig. S4). In this case, the protonation does not show a pronounced increase of the quantum yield. Since it had been found that the quantum yield of ring opening induced by irradiation in the visible band at 576 nm did not show a protonation-gated effect either, the finer details of the pH-dependence of partial quantum yields at that wavelength were not further investigated in this work. The fact that photochemical ring closing of the open form of DAEs is much more sensitive to changes of substitution patterns than ring opening of the closed form has been pointed out by Uchida *et al.*¹⁶ It is paralleled by effects on the absorption spectra which are much more pronounced in the open form than in the closed form, a fact also apparent for the compound investigated in this work.

Conclusion

In this paper, the method for the photostationary kinetic analysis of a coupled system of equilibrium forms of two photoisomers was rigorously developed in the framework of



the Gauglitz method of photostationary kinetics. For protolytic equilibria, it was demonstrated how the individual quantum yields for each protolytic form can be isolated from the pH-dependence of the overall pseudo quantum yield Q . But in general, the method would be also applicable to other equilibria, potentially causing photochemical gating effects, such as *e.g.* complex formation or thermal isomerization equilibria. In case of the studied switchable molecular wire, it could be proven that it needs protonation on both side arms to achieve the large increase of the quantum yield of ring closure, whereby it is worth of note that the sites of protonation are located at some distance from the switching center.

Acknowledgements

Financial support of this work by the Deutsche Forschungsgemeinschaft through the SFB 767 is gratefully acknowledged. The authors thank M $\text{C}at$ Company for a generous gift of palladium based cross coupling catalysts.

References

- J. Wolf, I. Eberspächer, U. Groth and T. Huhn, *J. Org. Chem.*, 2013, **78**, 8366–8375.
- D. Dulić, S. J. van der Molen, T. Kudernac, H. T. Jonkman, J. J. D. de Jong, T. N. Bowden, J. van Esch, B. L. Feringa and B. J. van Wees, *Phys. Rev. Lett.*, 2003, **91**, 207402.
- S. J. van der Molen and P. Liljeroth, *J. Phys.: Condens. Matter*, 2010, **22**, 133001.
- Y. Kim, T. J. Hellmuth, D. Sysoiev, F. Pauly, T. Pietsch, J. Wolf, A. Erbe, T. Huhn, U. Groth, U. E. Steiner and E. Scheer, *Nano Lett.*, 2012, **12**, 3736–3742.
- S. H. Kawai, S. L. Gilat and J.-M. Lehn, *Eur. J. Org. Chem.*, 1999, 2359–2366.
- F. M. Raymo and S. Giordani, *J. Am. Chem. Soc.*, 2002, **124**, 2004–2007.
- S. Sortino, S. Petralia, S. D. Bella, M. Tomasulo and F. M. Raymo, *New J. Chem.*, 2006, **30**, 515–517.
- S. Silvi, E. C. Constable, C. E. Housecroft, J. E. Beves, E. L. Dunphy, M. Tomasulo, F. M. Raymo and A. Credi, *Chem. – Eur. J.*, 2009, **15**, 178–185.
- B. M. Neilson and C. W. Bielawski, *ACS Catal.*, 2013, **3**, 1874–1885.
- M. M. Russew and S. Hecht, *Adv. Mater.*, 2010, **22**, 3348–3360.
- C. Sciascia, R. Castagna, M. Dekermenjian, R. Martel, A. R. Srimath Kandada, F. Di Fonzo, A. Bianco, C. Bertarelli, M. Meneghetti and G. Lanzani, *J. Phys. Chem. C*, 2012, **116**, 19483–19489.
- S. E. Braslavsky, *Pure Appl. Chem.*, 2007, **79**, 293–465.
- S. Kobatake and Y. Terakawa, *Chem. Commun.*, 2007, 1698–1700.
- K. Yumoto, M. Irie and K. Matsuda, *Org. Lett.*, 2008, **10**, 2051–2054.
- M. Irie, K. Sakemura, M. Okinaka and K. Uchida, *J. Org. Chem.*, 1995, **60**, 8305–8309.
- K. Uchida, T. Matsuoka, S. Kobatake, T. Yamaguchi and M. Irie, *Tetrahedron*, 2001, **57**, 4559–4565.
- H. H. Liu and Y. Chen, *J. Mater. Chem.*, 2009, **19**, 706–709.
- M. Cipolloni, A. Heynderickx, F. Maurel, A. Perrier, D. Jacquemin, O. Siri, F. Ortica and G. Favaro, *J. Phys. Chem. C*, 2011, **115**, 23096–23106.
- F. Ortica, M. Cipolloni, A. Heynderickx, O. Siri and G. Favaro, *Photochem. Photobiol. Sci.*, 2012, **11**, 785–793.
- J. Massaad, J.-C. Micheau, C. Coudret, R. Sanchez, G. Guirado and S. Delbaere, *Chem. – Eur. J.*, 2012, **18**, 6568–6575.
- Y. Kutsunugi, C. Coudret, J. C. Micheau and T. Kawai, *Dyes Pigm.*, 2012, **92**, 838–846.
- G. Gauglitz, *J. Photochem.*, 1976, **5**, 41–47.
- G. Gauglitz and S. Hubig, *J. Photochem.*, 1981, **15**, 255–257.
- H. D. Brauer, R. Schmidt, G. Gauglitz and S. Hubig, *Photochem. Photobiol.*, 1983, **37**, 595–598.
- D. Sysoiev, A. Fedoseev, Y. Kim, T. E. Exner, J. Boneberg, T. Huhn, P. Leiderer, E. Scheer, U. Groth and U. E. Steiner, *Chem. – Eur. J.*, 2011, **17**, 6663–6672.
- The 95% conversion rate reported in ref. 4 was carefully rechecked for the present experiment and found to be slightly higher (97%) than previously reported.
- Wolfram Research Inc., *Mathematica. Version 9.0 ed.*, Wolfram Research, Inc., Champaign, Illinois, 2012.
- In case of the dicarboxylic acid of the photoactive dithienylethene unit, the authors of ref. 20 attempted to evaluate the two deprotonation constants independently. The result of the mono-anion being more acidic than the neutral diacid is not really convincing, the more so, since the spectrum of the monoanion resulting from the analysis exhibits a sharp spectral maximum, where both, neutral form and dianion exhibit only weak shoulders, and although the spectral series on base titration appears to change in a monotonic way at all wavelengths. Also, apart from the charge, the intramolecular hydrogen bridge in the monoacid anion would be expected to exhibit a further weakening effect on the acidity of this species.
- J. F. Coetzee and G. P. Cunningham, *J. Am. Chem. Soc.*, 1965, **87**, 2529–2534.
- S. Delbaere, G. Vermeersch and J. C. Micheau, *J. Photochem. Photobiol., C*, 2011, **12**, 74–105.
- A detailed introduction to the mathematical description of a reversible photoreaction according to the method described by Gauglitz^{22,23} can be found in the Supporting Information given for ref. 25.
- The corresponding equation for decomposing the effective quantum yield as a function of pH, as given by Massaad *et al.*²⁰ (*cf.* Supporting Information of that paper, page 5, Section 2) applies only for irradiations at isosbestic points, *i.e.* in situations where the absorption coefficients of the species involved are equal. In the presence of more than two species such wavelengths are rarely found.

