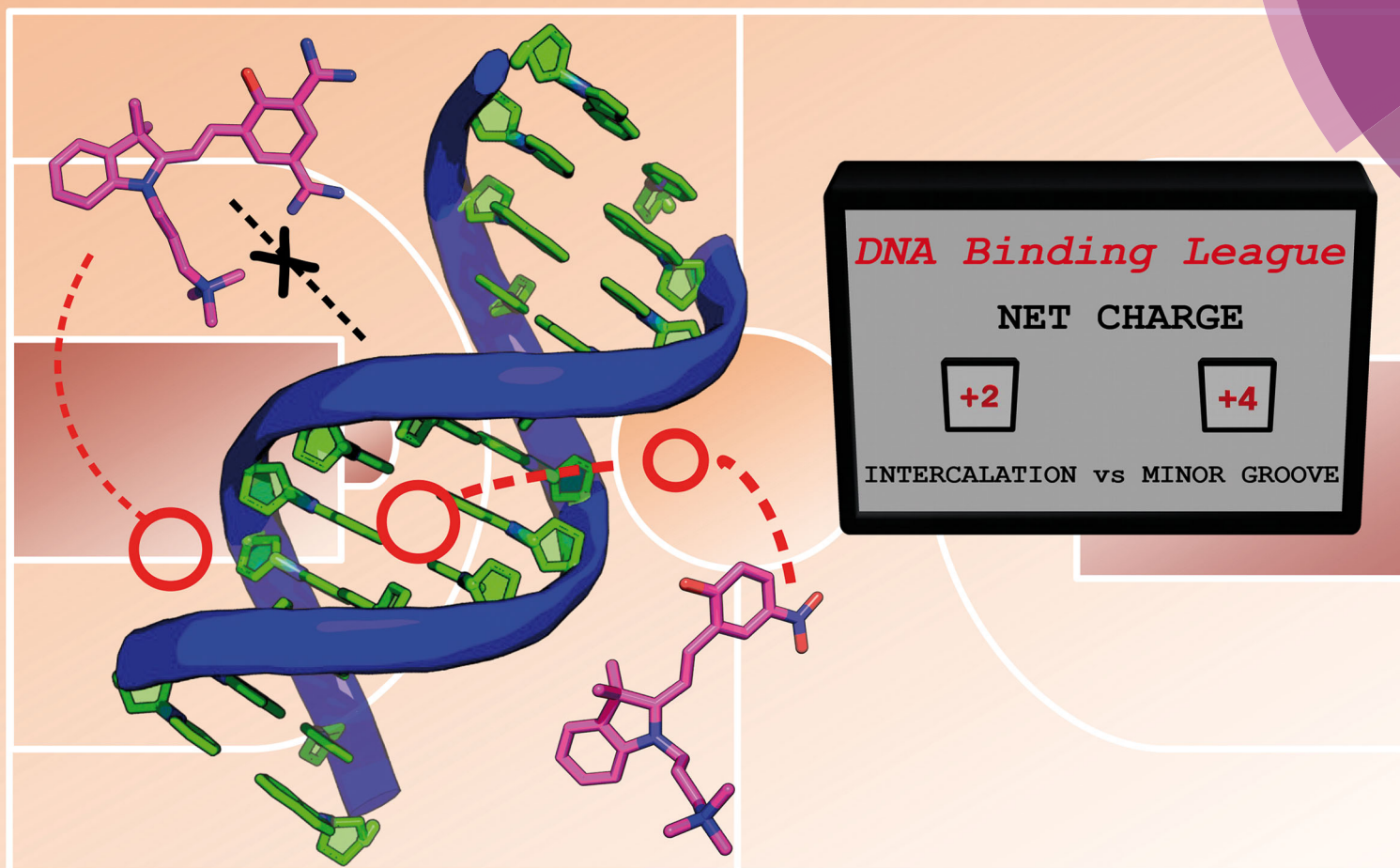


PCCP

Physical Chemistry Chemical Physics
rsc.li/pccp



ISSN 1463-9076



ROYAL SOCIETY
OF CHEMISTRY

Celebrating
IYPT 2019

COMMUNICATION

Pedro A. Sánchez-Murcia, Leticia González *et al.*
DNA-binding mechanism of spiropyran photoswitches:
the role of electrostatics





Cite this: *Phys. Chem. Chem. Phys.*,
2019, 21, 8614

Received 7th December 2018,
Accepted 15th February 2019

DOI: 10.1039/c8cp07508e

rsc.li/pccp

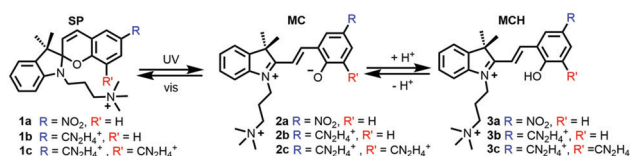
DNA-binding mechanism of spiropyran photoswitches: the role of electrostatics†

Davide Avagliano, Pedro A. Sánchez-Murcia * and Leticia González *

The binding mechanism of the protonated open form of three spiropyran derivatives into a 12-mer (poly-dAT)₂ has been unveiled by means of computational methods. It is found that the electrostatic term in the probe:DNA binding energy, modulates the binding mode, providing new guidelines for the design of spiropyran photoswitches with specific binding modes to DNA.

Molecular photoswitches – molecules that change their chemical configuration upon light absorption – are useful tools to monitor the conformational state of biomolecules, in both temporal and spatial scales.¹ The advantages of these probes are that the implicated species in the photochemical equilibrium highly differ in the binding affinity to the macromolecule upon light absorption, that these species are interconvertible in a reversible manner, and most importantly, that the two species can be selectively accessed by illumination at different wavelengths and/or by thermal activation. Spirogyrans (SP) (Scheme 1) belong to a well-known class of photoswitches, which after light irradiation undergo heterocyclic cleavage to yield the *trans*-merocyanine form (MC).^{2–6}

Spirogyrans have an indoline and a chromene moiety oriented perpendicular to each other and bound together *via* a C_{sp}³ spiro-junction. Upon absorption of UV light, the sigma C–O bond on the spiranic carbon cleaves and builds the MC form. Thereafter, MC can reversibly turn to SP with heat or after irradiation with visible light. Whereas the SP species has moderate lipophilicity, the conjugated coplanar system MC is a zwitterionic species with increased polarity favouring intercalation to polyanions such as DNA *via* π – π stacking interactions. In contrast, the SP form shows no or very little affinity to DNA. The half-time of the MC form increases upon substitution on the phenyl ring with electron withdrawing groups, such as NO₂.^{9–12} For instance, almost half of the population in solution of **1a** (Scheme 1), which contains the scaffold 3',3'-dimethyl-6-nitrospiro[2H-1-benzopyran-2,2'-indoline] (6-nitro-BIPS) with a



Scheme 1 Spirocyclic (SP) compounds **1a**,⁷ **1b**,⁸ the proof of concept **1c** and their corresponding unprotonated (MC, **2**) and protonated (MCH, **3**) merocyanine open forms. Compounds **3** are studied in this work.

trimethylammoniumpropyl group on the indole nitrogen, is converted to **2a** after irradiation at 254 nm.⁷ Despite its efficiency, the binding mode of this family of photoswitches into DNA is still under debate. Experiments with flow-oriented linear dichroism (LD) spectroscopy strongly support that **2a** intercalates into DNA given the dramatic blue-shift of the absorption spectrum of the drug in solution.⁷ A few years later, the same authors found¹³ that the maximum of the absorption spectrum of the protonated form **3a** in buffered aqueous solution is blue-shifted compared to **2a**, in analogy to what was previously observed after binding of **2a** to DNA.⁷ Therefore, it is likely that **2a** gets protonated to form **3a** before binding to DNA.

The latter hypothesis was also supported by the measurement of the DNA binding constants of the amidinium-substituted derivatives **1b**–**3b**.⁸ The protonated **3b** form intercalates stronger into DNA by a factor of *ca.* 50 compared to the unprotonated **2b** one. Again, and in analogy to **3a**, the protonated form **3b** was proposed to be the relevant DNA binder with a similar angle in the LD experiment (*ca.* 70°). Based on this experimental evidence, the authors ascribed the differences in the binding affinities of **3a** and **3b** to the total net positive charge on the photoprobe.⁸ However, the mechanism of the intercalation of **3a** and **3b** remains uncertain since before intercalation these species could bind to different grooves of the DNA.¹⁴ This is, for example, the case of the drug daunomycin (aka. doxorubicin) that can visit up to four states before intercalation, some of them pre-intercalative states,¹⁵ which were identified by molecular dynamics (MD) simulations as local minima bound in the minor groove of DNA.^{16,17}

Institute of Theoretical Chemistry, Faculty of Chemistry, University of Vienna,
Währinger Strasse 17, 1090, Vienna, Austria. E-mail: leticia.gonzalez@univie.ac.at

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8cp07508e



In this work, we unravel the intercalation modes of spiro-pyran derivatives from the solvent into DNA using MD and umbrella sampling simulations. To this aim, a 12-mer (poly-dAT)₂ has been selected as these compounds are known to bind to AT.⁷ Upon light-driven ring opening and concomitant protonation of the phenoxy moiety (MCH in Scheme 1), **3a** intercalates into (poly-dAT)₂. However, before inserting into DNA, **3a** has to travel from the bulk solvent into the nucleic acid. Potentially, the probe can visit a few binding modes before intercalation, *i.e.* entering DNA by the major (path M) or by the minor (path m) groove, as sketched in Fig. 1a. Experimentally, it is known⁷ from flow-oriented LD that **3a** in the presence of DNA shows one negative peak around 420 nm. A single peak indicates that **3a** has only one binding mode. Furthermore, it was determined that the photoprobe forms an angle of $80 \pm 5^\circ$ with the main Z-axis of DNA, meaning that the transition dipole moment of **3a** is parallel to the plane of the sandwiched nucleobases, and thus, the molecule is intercalated in a sandwich-like way between the DNA base pairs. Unfortunately, and to the best of our knowledge, neither kinetics nor structural data of the intercalation process of any of the derivatives of the SP family have been reported to date. Therefore, we explore here the two possible binding pathways to the major (M) and minor (m) groove of **3a** by means of umbrella sampling MD simulations.

As the reaction coordinate (RC) for binding to both grooves, we define the distance between the center-of-mass of **3a** and the center-of-mass of the four nucleobases in between **3a** inserts.

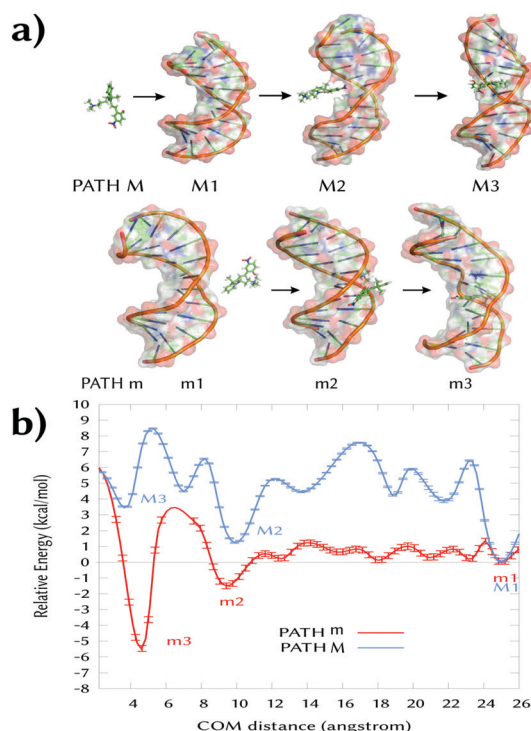


Fig. 1 (a) Intercalation process of **3a** into a 12-mer (poly-dAT)₂ through path M (major groove) and path m (minor groove). (b) Free energy profiles (kcal mol⁻¹) of path M (major) and m (minor) computed with umbrella sampling MD simulations. The bootstrap error values are shown over the plots.

From the initial intercalated geometries (named M3 and m3 in Fig. 1a), we pull out the probe from the DNA (RC ~ 2 Å) into the bulk solvent up to RC = 26 Å (M1 and m1). The whole RC was divided into 50 windows separated by 0.5 Å and the system was allowed to oscillate around the RC for 5 ns per window – the time at which the energy curves were converged (Sections S1 and S2, ESI†). The initial guesses for M3 and m3 were obtained by manual docking after structural superimposition of our system with the crystal structure of daunomycin intercalated into DNA (PDB id. 1VTH).¹⁸ Then, the two systems were minimized and equilibrated in explicit solvent. Due to the presence of the two methyl groups on the indoline ring of **3a** (Scheme 1), there was only one possibility in each case for the relative orientation of these groups with respect to the methyl group of the two thymine units, in between the probe intercalates in the states M3 and m3.

Fig. 1b shows the obtained energy profiles for path M (blue line) and m (red line). In both cases, the chemical probe visits several pre-intercalative states before intercalation. However, our calculation clearly predicts that binding to the major groove (pathway M) is unfavorable. When **3a** approaches to the DNA through the major groove it visits only metastable states since all the local minima between 25 and 3 Å are higher in energy compared to the minimum M1. Moreover, the intercalated state M3 is less stable than both the solvated state M1 (RC = 25 Å, $\Delta E > 3$ kcal mol⁻¹) and the local minimum M2 (at ~ 10 Å). Inspection of the M3 geometry reveals that the ligand's rings are coplanar and inserted between four nucleobases parallel to them. In such a conformation, the binding site is highly distorted and the hydrogen bonds between the surrounding base pair are broken (A7:T6' and A5:T8', Fig. S1, ESI†). This explains the unfavorable energy for the state M3.

In contrast, the intercalated state m3 is the most thermodynamically stable binding mode. Along the minor groove binding pathway, we identified several metastable pre-intercalative states of **3a** (between RC = 8 Å and RC = 23 Å) that end up in the local minimum m2 at RC = 9 Å. Since the energy of m2 is quite similar to m1 ($\Delta E = -1.5$ kcal mol⁻¹), the binding mode m2 is reversible and **3a** could return to the bulk solvent. However, supplying only *ca.* 5 kcal mol⁻¹, **3a** can overcome the energy barrier between m2 and m3 to reach the latter intercalative state. This extra energy is used to open the space between the nucleobases.

Fig. 2 illustrates the geometry of **3a** in the m2 state. The indoline and the phenol rings are almost co-planar with a small rotation (*ca.* 15° with respect to each other, Fig. S2, ESI†) and the molecule is parallel to the DNA backbone. The nitro group is oriented to the same side of the ammoniumpropyl tail and projected outwards to the solvent. In such a disposition, the phenolic hydroxyl group establishes a hydrogen-bond with the carbonyl oxygen of a thymine T6' (dashed line, Fig. 2a). This interaction remains when the system is further simulated during 100 ns without restraints. This observation, together with the orientation of the phenol of **3a** in m2 and m3, could explain the higher experimental binding constant values for the MCH form (**3a**) compared to the MC one (**2a**).⁷ Indeed, as can be seen in Fig. 2b and 3a, the phenoxy group in **2a** would face a negatively-charged region in both m3 and m2 states, respectively.



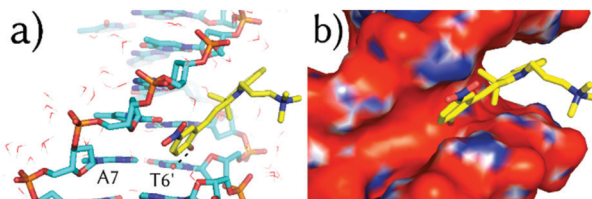


Fig. 2 (a) Representative binding geometry of **3a** (sticks, carbon atoms colored in yellow) in m2 after 100 ns of unrestrained MD simulation. Only the closest water molecules (lines) are shown. (b) The APBS-generated electrostatic potential on the van der Waals surface of the DNA in m2 in the presence of **3a**.

In addition, the computed adaptive Poisson–Boltzmann solver (APBS) electrostatic potential on the surface of the DNA for m2 (Fig. 2b) shows that **3a** binds to a region with high negative potential (red contour) and that the nitro group is oriented to a positive region (blue contour). Thus, **3a** establishes a large number of contacts in the short range (*ca.* 243 in an inner sphere of 4 Å) along the MD simulation, including the interactions between the trimethylammoniumpropyl group of **3a** and the phosphate groups in the DNA.

Once **3a** visits m2, the system requires *ca.* 5 kcal mol^{−1} to make room between the four stacking nucleobases A7:T6' and A5:T8' it binds to. Once in m3, only the phenol ring of **3a** is parallel to the planes defined by the former four nucleobases and almost perpendicular to the main Z-axis of the DNA (Fig. 3a). In m3, **3a** establishes face-to-face π – π stacking interactions between the phenol ring and the base pairs T6:A7' and A5:T8', as observed for other intercalative agents.¹⁹ The distances between the phenol ring and each of the former base pairs are, in fact, below 4 Å. The hydrogen-bonds of the Watson–Crick base pairs T6:A7' and A5:T8' and the surrounding A7:T6' and T8:A5' are kept in the presence of **3a** but the distance between the nucleobases on the same strand T6 and A5 and A7' and T8' is increased from the non-intercalative crystallographic distance of 5 Å (like between A7 and T6) to 7.1 Å and 6.3 Å, respectively.

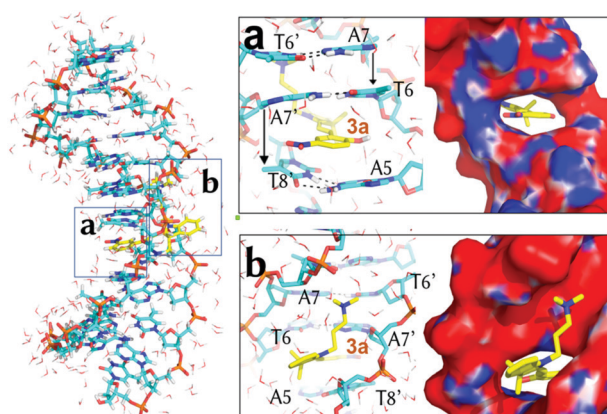


Fig. 3 Representative binding geometry of **3a** in m3 after 100 ns of unrestrained MD simulation. Boxes (a) and (b) zoom-in on the detailed orientation of the phenol and the indoline rings, respectively. Only the closest water molecules (lines) are shown. The APBS-generated electrostatic potential is also shown on the van der Waals surface of the DNA.

Finally, the nitro group of **3a** is again looking at a positive potential region (Fig. 3a) and the ammoniumpropyl tail interacts with a region with negative potential in between the phosphate groups (Fig. 3b).

The intercalation of **3a** increases the angle between the indoline and the chromene moieties (Fig. S2, ESI†) and enlarges the amplitude of this hampered rotation between the two rings. This finding agrees with the larger experimental wavelength shift in the absorption spectrum upon intercalation⁷ than when going from water to chloroform in the absence of DNA, what may indicate a conformational change to a nonplanar structure of **3a** in m3.

The substitution of the nitro group in **3a** by an amidinium group (compound **3b**)⁸ is known to increase the binding constant of the photoprobe to DNA. This effect could be due to the specific interactions of the amidinium group, the increment in the net positive charge in **3b**, or a combination of both factors. In order to discern between these explanations, we simulated the binding states m2 and m3 of **3b** with the (poly-dAT)₂. We directly employed the already calculated binding states m2 and m3 of **3a**, as there is no experimental evidence of different binding geometries or orientations between the nitro and the amidinium substituted MCH (**3a** and **3b**), except for a slight difference in the measured angle in the flow-oriented linear dichroism (LD) spectra.⁸

Our computations show that **3b** in both m2 and m3 states shares the main binding spots found in the nitro derivative (Fig. S3, ESI†) with the amidinium group projected to the solvent. In the m2 binding mode, the amidinium group turned out to have no specific interaction with the DNA backbone or with the nucleobases. Therefore, the first possibility highlighted above cannot explain why **3b** binds stronger to DNA than **3a**. Thus, we computed the binding energies of **3a** and **3b** to DNA using the molecular mechanics Poisson–Boltzmann surface area (MM-PBSA) method.²⁰

We considered both m2 and m3 binding modes for each of the compounds. Table 1 collects the total solvated binding energies of **3a** and **3b** to DNA and their individual terms. In both cases, the intercalative state m3 is the most stable state (Fig. S4, ESI†). **3b** shows the largest binding energy value due to a smaller positive (repulsive) electrostatic energy component (ΔE^{elec}) than in **3a**. This term has two contributions: the electrostatic component of the Poisson–Boltzmann solvation energy (ΔE^{PB}) and the electrostatic term in the gas phase (ΔE^{EEL}) (Table S1, ESI†). In the pre-intercalative state m2, ΔE^{elec} strongly decreases between **3a** and **3b** (109%) so that the electrostatic component is so severely reduced in **3b** that it turns negative (attractive by -1.02 kcal mol^{−1}). Therefore, one is left to conclude that the increase of the positive charge on the probe strengthens the binding of **3** to DNA.

With this conclusion in hand, and as a proof of concept, we designed the photoprobe 6,4-diamidinium-MCH (**3c**, Scheme 1), where an extra amidinium group was introduced in the *ortho* position to the hydroxyl group of the phenol ring. This compound – with an extra net charge – should validate our hypothesis that additional electrostatic contributions could



Table 1 Computed MM-PBSA mean binding energies (kcal mol^{−1}) at 300 K of **3a–3c** to (poly-dAT)₂. The errors represent the standard deviations of the mean values. The total binding energy includes the entropy contribution

Compound	Mode	ΔE^{elec}	Non-electrostatic terms		$T\Delta S$	ΔG^{bind}
		$(\Delta E^{\text{EEL}} + \Delta E^{\text{PB}})$	ΔE^{vdW}	ΔE^{nonpol}		
3a	m3	15.81 ± 0.14	−55.17 ± 0.25	−4.32 ± 0.01	−21.73 ± 0.09	−21.96 ± 0.24
3b		10.10 ± 0.40	−55.78 ± 0.23	−4.62 ± 0.40	−20.96 ± 0.17	−29.35 ± 0.33
3c		11.09 ± 0.59	−50.71 ± 0.42	−4.57 ± 0.03	−26.42 ± 0.37	−17.77 ± 0.74
3a	m2	10.82 ± 0.22	−37.72 ± 0.41	−3.23 ± 0.01	−20.97 ± 0.02	−9.29 ± 0.47
3b		−1.02 ± 0.60	−34.16 ± 0.26	−3.22 ± 0.01	−23.88 ± 0.13	−14.52 ± 0.36
3c		−12.07 ± 0.39	−29.91 ± 0.49	−3.19 ± 0.02	−24.71 ± 0.26	−20.46 ± 0.61

then further increase the binding energy to DNA, and thereby, its binding constant. The *ortho* position was selected since it is chemically modifiable²¹ and the orientation of the group does not interfere with the proposed m2 and m3 binding modes. The values for the total binding energy are also shown in Table 1. Surprisingly, we observed that the extra amidinium group of **3c** increases the binding energy only for the m2 mode, and the m3 state is less stable than the corresponding state in **3a** or **3b**. Note that the role of the entropy, although significant, is not the decisive term for the relative stability between states m2 and m3.

Fig. 4 depicts the binding energies for both the m2 and m3 binding modes of all three derivatives **3a–3c**. Whereas in **3a** and **3b** the m2 state is always below in energy compared to m3, in **3c** the pre-intercalative state m2 competes with m3 as the most stable binding mode. This means that, from the thermodynamical point of view, **3c** would not intercalate, but rather bind in the minor groove, as other agents do.²² Our results are supported by the fact that **3c** has similar size and charge compared to other minor groove binders (Fig. S5, ESI†) and also, like most of them, shows pronounced AT selectivity.

These results indicate that care has to be taken in the modulation of the electrostatic component in photoactivatable DNA-targeting spiropyran. Although experiments suggest that MCH species bind to DNA better than MC due to its increased electrostatics (e.g. **2b** versus **3b**),⁸ an excess of net positive charge on MCH can stabilize the pre-intercalative state m2 such that

this binding site becomes competitive (e.g. **3c** versus **3b**); this is e.g. the case in polyamides or diamidines, which are species known to selectively bind AT pairs in the minor groove but not to intercalate.²²

In summary, this work indicates that the modulation of the electrostatic potential in this family of spiropyran can be used as a tool to control their binding mode to DNA (minor groove binder or intercalation). And most importantly, the current assumption that increasing the net positive charge on the spiropyranic photoswitcher always will favor the intercalation process is, at least, not generalizable and may require further experimentation.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

D. A. thanks the H2020-MSCA-ITN-2017 ETN programme LightDyNAMics, under grant agreement no. 765266. P. A. S.-M. thanks the Austrian FWF (Project M 2260). The authors would like to thank Dr J. J. Nogueira for preliminary discussions on spiropyran. The Vienna Scientific Cluster (VSC) is acknowledged for kind allocation of computational resources.

Notes and references

- W. Szymański, J. M. Beierle, H. A. V. Kistemaker, W. A. Velema and B. L. Feringa, *Chem. Rev.*, 2013, **113**, 6114–6178.
- A. Mustafa, *Chem. Rev.*, 1948, **43**, 509–523.
- (a) R. Klajn, *Chem. Soc. Rev.*, 2014, **43**, 148–184; (b) C. L. Fleming, S. Li, M. Grøtli and J. Andréasson, *J. Am. Chem. Soc.*, 2018, **140**, 14069–14072.
- V. I. Minkin, *Chem. Rev.*, 2004, **104**, 2751–2776.
- S. Prager, I. Burghardt and A. Dreuw, *J. Phys. Chem. A*, 2014, **118**, 1339–1349.
- F. Liu and K. Morokuma, *J. Am. Chem. Soc.*, 2013, **135**, 10693–10702.
- J. Andersson, S. Li, P. Lincoln and J. Andréasson, *J. Am. Chem. Soc.*, 2008, **130**, 11836–11837.
- M. Hammarson, J. R. Nilsson, S. Li, P. Lincoln and J. Andréasson, *Chem. – Eur. J.*, 2014, **20**, 15855–15862.

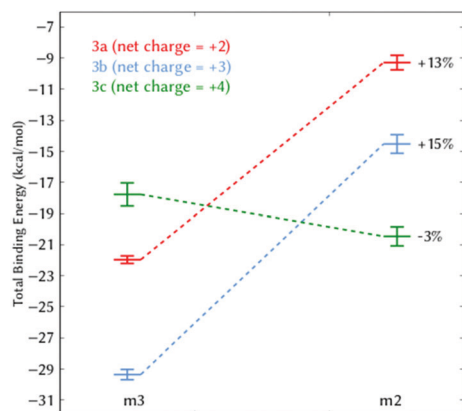


Fig. 4 Total binding energy (kcal mol^{−1}) of **3a–3c** between the binding modes m2 and m3. The percentage indicates the relative energetic change of m2 compared to m3.



- 9 C. Lenoble and R. S. Becker, *J. Phys. Chem.*, 1986, **90**, 62–65.
- 10 M. Gehrtz, C. Braeuchle and J. Voitaender, *J. Am. Chem. Soc.*, 1982, **104**, 2094–2101.
- 11 Y. Sheng, J. Leszczynski, A. A. García, R. Rosario, D. Gust and J. Springer, *J. Phys. Chem. B*, 2004, **108**, 16233–16243.
- 12 S. Aldridge and C. Jones, *Chem. Soc. Rev.*, 2016, **45**, 763–764.
- 13 M. Hammarson, J. R. Nilsson, S. Li, T. Beke-Somfai and J. Andréasson, *J. Phys. Chem. B*, 2013, **117**, 13561–13571.
- 14 A. A. Almaqwashi, T. Paramanathan, I. Rouzina and M. C. Williams, *Nucleic Acids Res.*, 2016, **44**, 3971–3988.
- 15 V. Rizzo, N. Sacchi and M. Menozzi, *Biochemistry*, 1989, **28**, 274–282.
- 16 A. Mukherjee, R. Lavery, B. Bagchi and J. T. Hynes, *J. Am. Chem. Soc.*, 2008, **130**, 9747–9755.
- 17 M. Wilhelm, A. Mukherjee, B. Bouvier, K. Zakrzewska, J. T. Hynes and R. Lavery, *J. Am. Chem. Soc.*, 2012, **134**, 8588–8596.
- 18 C. M. Nunn, L. Van Meervelt, S. Zhang, M. H. Moore and O. Kennard, *J. Mol. Biol.*, 1991, **222**, 167–177.
- 19 G. Barone, C. F. Guerra, N. Gambino, A. Silvestri, A. Lauria, A. M. Almerico and F. M. Bickelhaupt, *J. Biomol. Struct. Dyn.*, 2008, **26**, 115–129.
- 20 N. Homeyer and H. Gohlke, *Mol. Inf.*, 2012, **31**, 114–122.
- 21 (a) C. Kaiser, T. Halbritter, A. Heckel and J. Wachtveitl, *ChemistrySelect*, 2017, **2**, 4111–4123; (b) H. Görner, *Phys. Chem. Chem. Phys.*, 2001, **3**, 416–423.
- 22 (a) R. R. Tidwell and D. W. Boykin, in *Dicationic DNA Minor Groove Binders as Antimicrobial Agents*, in *Small Molecule DNA and RNA Binders*, ed. M. Demeunynck, C. Bailly, W. D. Wilson, John Wiley & Sons, Ltd, Chichester, 2002, pp. 414–460; (b) B. Nguyen, D. Hamelberg, C. Bailly, P. Colson, J. Stanek, R. Brun, S. Neidle and W. David Wilson, *Biophys. J.*, 2004, **86**, 1028–1041; (c) D. D. V. H. Xuemei Cai and P. J. Gray Jr., *Cancer Treat. Rev.*, 2009, **35**, 437–450.

