

## RESEARCH ARTICLE

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Cite this: *Org. Chem. Front.*, 2024, **11**, 5754The *N*-biphenyldihydroisoquinolinium scaffold as a novel motif for selective fluorimetric detection of quadruplex DNA†‡Denisa Soost,<sup>a</sup> Gerhard Bringmann<sup>b</sup> and Heiko Ihmels<sup>\*a</sup>

The spectroscopic investigation of the DNA-binding properties of (*S*)-6,8-dimethoxy-2-(4'-methoxy-[1,1'-biphenyl]-4-yl)-1,3-dimethyl-3,4-dihydroisoquinolin-2-ium revealed the cationic donor-acceptor-substituted biaryl unit as a promising motif for selective fluorimetric detection of particular quadruplex DNA (G4-DNA) forms. The title compound exhibits the characteristic solvent-dependent dual emission of biaryl fluorophores; however, the emission quantum yields are very low ( $\Phi_{\text{fl}} \leq 0.01$ ) because of radiationless deactivation of the excited state by conformational changes. In contrast, the emission intensity of the biaryl derivative increases by a factor of 2–10 on association with G4-DNA ( $K_{\text{B}} = 8 \times 10^4 \text{ M}^{-1}$ – $4 \times 10^5 \text{ M}^{-1}$ ), because the structural relaxation of the excited ligand is suppressed in the binding site. The signal pattern of the dual emission, specifically the relative contribution of each band, varies with the different G4-quadruplex forms **22AG**, **c-kit**, **c-myc**, **c-kas**, and **h-ras1**, most likely caused by the different dynamic flexibility of the ligand in the distinct binding sites. These effects enable the fluorimetric identification of sterically constrained binding sites, such as in **c-kit** and **h-ras1**, even with the naked eye.

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## Introduction

The selective staining and detection of nucleic acids is an important bioanalytical technique in chemistry and biology.<sup>1</sup> For this purpose, organic fluorescent probes figure as valuable and versatile tools.<sup>2</sup> Specifically, dyes with a very low emission intensity that increases upon association with the target analyte, often referred to as fluorescent “light-up probes”, have proven to be very useful and provide valuable chemosensors for the fluorimetric detection of DNA and similar targets *in vitro* and *in vivo*.<sup>3</sup> In this context, especially the non-canonical quadruplex DNA (G4-DNA) is an important target.<sup>4</sup> Quadruplex DNA is formed upon folding and assembly of guanine-rich DNA strands to a stack of usually three neighboring guanine quartets.<sup>5</sup> The structures of quadruplex DNAs vary depending mainly on the particular sequence of the parent single strand(s).<sup>6</sup> Most importantly, G4-DNA structures have essential biological functions, for example in the promoter

regions of oncogenes, which influence the growth of cancer cells,<sup>7</sup> or in the single-stranded overhang of telomeric DNA.<sup>8</sup> Therefore, G4-DNA is an attractive target for the development of fluorescent probes that enable the detection and monitoring of this DNA form.<sup>9</sup> Indeed, various fluorescent probes have already been developed that may be used to detect G4-DNA structures and to differentiate them from other DNA forms<sup>10</sup> *in vitro*<sup>11</sup> and in cells.<sup>12</sup> Hence, it has been demonstrated that suitable cationic dyes, for example distyrylpyridinium derivatives such as **1**, bind selectively to particular target G4-DNA structures and that this complex formation is accompanied by a characteristic, G4-DNA-specific fluorescence light-up effect (Scheme 1).<sup>13</sup> As another notable example, the cyanine dye **2** exhibits a strong fluorescence light-up effect upon selective association with the G4-DNA form **c-myc** (Scheme 1),<sup>14</sup> which has been used to identify and distinguish this quadruplex form from other nucleic acid structures.<sup>15</sup> But, despite the highly favorable detection properties of these and other<sup>2,13</sup> fluorescent probes, there is still demand for novel approaches and for structural features that enable the selective detection of G4-DNA. Thus, there are, so far, only few examples of fluorescent probes available that are able to distinguish different topologies of G4-DNA with sufficient selectivity.<sup>16</sup> At the same time, it appears that selectivity is more easily accomplished for some G4-DNA forms, *e.g.* for **c-myc**,<sup>14,15</sup> whereas fluorescent probes for others, such as **c-kit**,<sup>17</sup> are still rare. Against this background, we report the coincidental

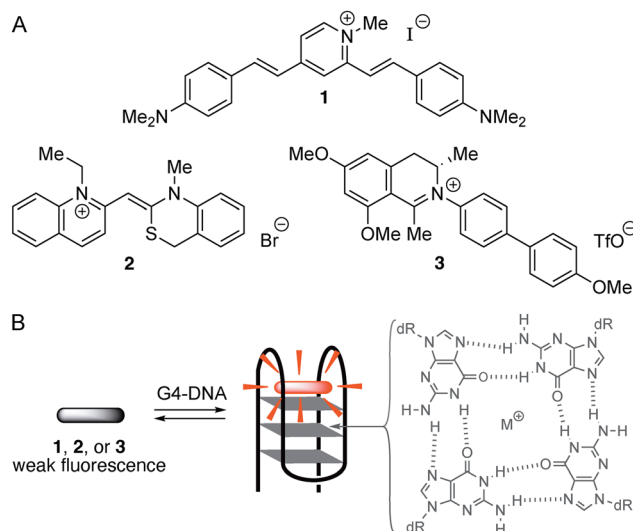
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† Dedicated to Prof. Dr. Frank Würthner on the occasion of his 60th birthday.

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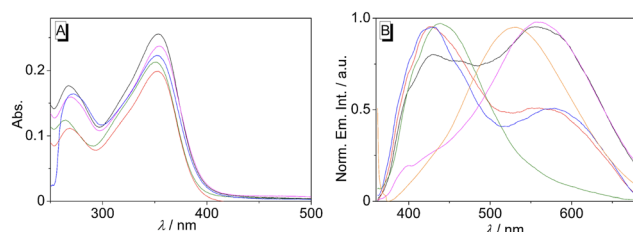


**Scheme 1** (A) Structures of distyrylpyridinium and cyanine dyes **1** and **2** and of the biphenyl-substituted 3,4-dihydroisoquinolin-2-ium **3**. (B) Schematic representation of the fluorescence light-up effect of ligands upon binding to G4-DNA; grey: structure of a guanine quartet, dR = deoxyribose residue of the DNA backbone.

observation of the selective fluorimetric response of an *N*-biphenyldihydroisoquinolinium fluorophore upon binding to G4-DNA, which may contribute to the search for novel functional features of G4-DNA-targeting fluorescent probes. During our studies of the DNA-binding properties of dihydroisoquinolinium-based natural products,<sup>18</sup> we have noticed that the very weakly fluorescent cationic biphenyl-substituted derivative **3** showed a distinct fluorescence in the presence of nucleic acids, especially pronounced with G4-DNA. Therefore, we investigated the photophysical and the DNA-binding properties of this particular biphenyl derivative. And we will demonstrate herein that, indeed, this compound is a promising starting point for the development of G4-DNA-selective fluorescent probes.

## Results and discussion

For a first orientation, the absorption and emission properties of biphenyl **3** were investigated in different solvents (Fig. 1 and



**Fig. 1** Absorption ( $c = 10 \mu\text{M}$ ) (A) and normalized emission spectra (B) ( $\lambda_{\text{ex}} = 350 \text{ nm}$ ) of **3** in H<sub>2</sub>O (green), DMSO (blue), MeOH (black), EtOH (magenta), MeCN (red), and glycerol (orange).

**Table 1** Absorption and emission properties of compound **3** in different solvents

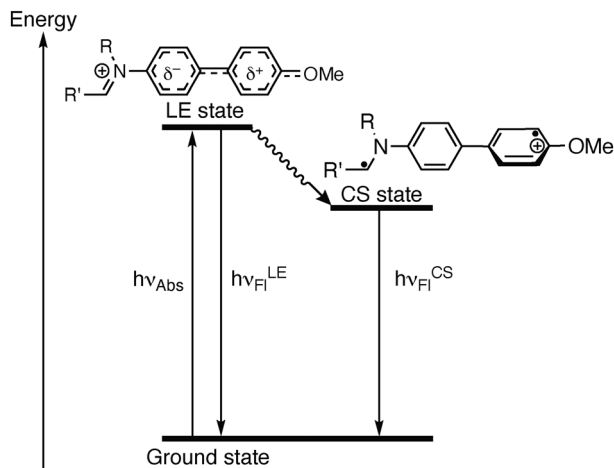
| Solvent          | $\lambda_{\text{abs}}^a/\text{nm}$ | $\epsilon^b/\text{L mol}^{-1} \text{ cm}^{-1}$ | $\lambda_{\text{fl}}^c/\text{nm}$ | $\Phi_{\text{fl}}^d$ |
|------------------|------------------------------------|------------------------------------------------|-----------------------------------|----------------------|
| H <sub>2</sub> O | 350 (264)                          | 21 300                                         | 439                               | <0.01                |
| DMSO             | 352 (272)                          | 22 280                                         | 428 (580)                         | 0.01                 |
| MeOH             | 353 (267)                          | 25 532                                         | 555 (429)                         | <0.01                |
| EtOH             | 354 (269)                          | 23 713                                         | 560                               | 0.01                 |
| MeCN             | 352 (268)                          | 19 849                                         | 426 (552)                         | <0.01                |

<sup>a</sup> Long-wavelength absorption maximum;  $c = 10 \mu\text{M}$ . <sup>b</sup> Molar extinction coefficient. <sup>c</sup> Emission maximum;  $\lambda_{\text{ex}} = 350 \text{ nm}$ . <sup>d</sup> Fluorescence quantum yield relative to anthracene in cyclohexane ( $\Phi_{\text{fl}} = 0.36$ ). The smaller maximum is given in parantheses.

Table 1). The absorption spectra of compound **3** are almost independent of the solvent, and the long-wavelength maxima only changed in a small range, from 350 nm in water to 354 nm in EtOH (Fig. 1A). In contrast, the fluorescence properties of compound **3** were found to depend on the solvent (Fig. 1B). While in water, a blue-shifted emission maximum was detected at 439 nm, a red-shifted maximum at 560 nm was found in EtOH. At the same time, dual emission of biphenyl **3** was observed in MeOH, MeCN, and DMSO, namely with two distinct emission bands at 352–353 nm and 556–580 nm (Fig. 1B and Table 1). In all employed solvents the emission quantum yields of **3** are very low ( $\Phi_{\text{fl}} \leq 0.01$ , Table 1).

Dual emission has been frequently observed in fluorescent biphenyl derivatives that are integrated in a donor-acceptor system. And it is commonly explained by the dynamic photo-physical processes in the excited state and the resulting different emitting excited species.<sup>19</sup> Specifically, the excitation induces an intramolecular charge transfer (CT), which is often referred to as charge shift (CS) in the case of ionic chromophores, and to conformational changes in the excited state leading to twisted (TICT) or planar structures (PICT).<sup>19b,20</sup> Accordingly, the emission properties of compound **3** are also influenced by the donor-acceptor interplay between the electron-donating methoxy group and the electron-withdrawing iminium functionality, in combination with the conformational changes of the biphenyl unit. Thus, compound **3** showed the expected emission band of a biphenyl chromophore at around 435 nm, which was assigned to the locally excited (LE) state, most likely formed after conformational change to an almost planar structure and a resonance-type excited state (Scheme 2).<sup>20f</sup> After further rotation around the biaryl axis, the TICT/CS state is formed with an almost perpendicular biphenyl structure, which results in the red-shifted emission band at *ca.* 550 nm (Scheme 2).<sup>20f,21</sup> Apparently, these two states are stabilized or destabilized to a different extent by several solvent properties, because there is no obvious relationship between the formation of the emission bands with the solvent parameters.<sup>20c,22</sup> As the fluorescence quantum yields are very low in all tested solvents, an additional non-radiative deactivation of the excited state obviously takes place with high efficiency, which may be caused by torsional relaxation around the biphenyl bond or around the C<sub>ar</sub>-N or the C<sub>ar</sub>-O bond of

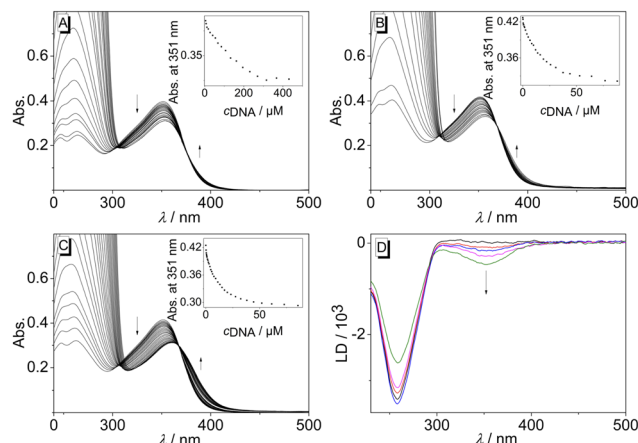




**Scheme 2** State diagram of the photoexcitation and deactivation pathways of the *N*-biphenyldihydroisoquinolinium **3**.

the donor and acceptor substituents.<sup>23</sup> To verify this assumption, the fluorescence spectra were recorded in media with different viscosity, namely in glycerol/water with increasing glycerol content (Fig. S4, *cf.* ESI†), because the viscosity of the solvent has a considerable effect on the rate of conformational changes, thus, on the emission quantum yield of excited molecules.<sup>24</sup> Indeed, it was observed that in glycerol/water mixtures with higher viscosity, namely with glycerol fraction >40%, the emission intensity of **3** increased significantly with increasing glycerol content (Fig. 1B and Fig. S4 *cf.* ESI†). Usually, this observation indicates a non-radiative deactivation of the excited state by conformational changes, such as torsional relaxation,<sup>25</sup> because in highly viscous glycerol the conformational changes are slowed down and emission becomes competitive.<sup>26</sup> Hence, torsional relaxation from either the LE state or the TICT/CS state of **3** to a non-emitting conformer most likely contributes to a significant extent to the non-radiative deactivation of the excited state.

The interactions of biphenyl **3** with calf thymus (ct) DNA, as a typical duplex DNA structure, and with representative quadruplex DNA (G4-DNA) forms **22AG**, **c-kit**, **c-kras**, **c-myc**, and **h-ras1** were investigated. These G4-DNA forms relate to specific, biologically relevant DNA sequences and have a unique sequence and structure, as defined by the arrangement of the G-quartets and loop regions.<sup>27</sup> The ligand-DNA interactions were monitored by photometric and fluorimetric titrations in aqueous buffer solutions. In almost all cases, a con-



**Fig. 2** Photometric titration of **3** ( $c = 20 \mu\text{M}$ ) with ct DNA (A) in BPE buffer ( $C_{\text{Na}^+} = 16 \text{ mM}$ ,  $\text{pH} = 7.0$ ), and with **22AG** (B) and **c-kit** (C) in K-phosphate buffer ( $C_{\text{K}^+} = 95 \text{ mM}$ ,  $\text{pH} = 7.0$ ). The arrows indicate the development of the absorption bands during the titration. Inset: plot of the absorption at  $\lambda = 351 \text{ nm}$  versus  $c_{\text{DNA}}$ . D: LD spectra of **3** in the presence of ct DNA ( $c = 20 \mu\text{M}$ ) in BPE buffer ( $C_{\text{Na}^+} = 16 \text{ mM}$ ,  $\text{pH} = 7.0$ ) at  $\text{LDR} = 0.0$  (black),  $0.2$  (red),  $0.5$  (blue),  $1.0$  (magenta),  $1.5$  (green).

tinuous decrease of the absorption maxima of compound **3** at  $351 \text{ nm}$  and a slight red shift ( $\Delta\lambda_{\text{ctDNA}} = 3 \text{ nm}$ ,  $\Delta\lambda_{22\text{AG}} = 6 \text{ nm}$ ,  $\Delta\lambda_{\text{c-kit}} = 9 \text{ nm}$ ) were observed upon addition of DNA (Fig. 2A, B, C and Fig. S1, *cf.* ESI†). This effect was most pronounced upon addition of G4-DNA **c-kit**. Moreover, during titrations with G4-DNA, an isosbestic point was formed at  $365 \text{ nm}$ , whereas with ct DNA an initially formed isosbestic point faded during titration. The binding isotherms derived from the photometric DNA titrations of compound **3** were used to determine the binding constants  $K_b$  (Table 2 and Fig. S2, *cf.* ESI†),<sup>28</sup> which revealed moderate binding affinities towards ct DNA ( $K_{\text{ct}} = 1.3 \times 10^4 \text{ M}^{-1}$ ) and to G4-DNA ( $K_{22\text{AG}} = 7.6 \times 10^4 \text{ M}^{-1}$ ;  $K_{\text{c-kit}} = 1.6 \times 10^5 \text{ M}^{-1}$ ;  $K_{\text{c-kras}} = 1.8 \times 10^5 \text{ M}^{-1}$ ,  $K_{\text{c-myc}} = 4.3 \times 10^5 \text{ M}^{-1}$ ,  $K_{\text{h-ras1}} = 3.1 \times 10^5 \text{ M}^{-1}$ ). However, the fitting curves of the binding isotherms of ct DNA, **c-kras**, and **h-ras1** showed deviations, presumably because of heterogeneous binding at relatively high or low ligand loading on the DNA. To further assess the binding mode of compound **3** with duplex DNA, the DNA-ligand complexes were analyzed by linear dichroism (LD) spectroscopy in the hydrodynamic field of a rotating cuvette.<sup>29</sup> In the presence of ct DNA, a negative LD signal between  $300 \text{ nm}$  and  $400 \text{ nm}$  was observed, that is in a range where only the ligand **3** absorbs. And the intensity of these LD bands

**Table 2** Binding constants,  $K_b$ , of compound **3** with ct DNA and G4-DNA, and shifts of melting temperature,  $\Delta T_m$ , of G4-DNA in the presence of **3**

|                               | ct DNA       | 22AG <sup>a</sup> /F21T <sup>b</sup> | c-kit <sup>a</sup> /FkitT <sup>b</sup> | c-kras <sup>a</sup> /FkrasT <sup>b</sup> | c-myc <sup>a</sup> /FmycT <sup>b</sup> | h-ras1       |
|-------------------------------|--------------|--------------------------------------|----------------------------------------|------------------------------------------|----------------------------------------|--------------|
| $K_b^a/10^4 \text{ M}^{-1}$   | 1.3          | 7.6                                  | 16                                     | 18                                       | 43                                     | 31           |
| $\Delta T_m^b/^\circ\text{C}$ | <sup>c</sup> | 6                                    | 3                                      | 6                                        | 2                                      | <sup>c</sup> |

<sup>a</sup> Binding constant determined from photometric titrations with **22AG**, **c-kit**, **c-kras**, **c-myc**, and **h-ras1**. <sup>b</sup> Determined from fluorimetric analysis of dye-labeled oligonucleotides; estimated error  $\pm 0.5^\circ\text{C}$ . **F21T** = fluo-G<sub>3</sub>(TTAG<sub>3</sub>)<sub>3</sub>-tamra, **FkitT** = fluo-AG<sub>3</sub>AG<sub>3</sub>CGCTG<sub>3</sub>AG<sub>2</sub>AG<sub>3</sub>-tamra, **FmycT** = fluo-TGAG<sub>3</sub>TG<sub>3</sub>TAG<sub>3</sub>TG<sub>3</sub>TA-tamra, **FkrasT** = fluo-AG<sub>3</sub>CG<sub>2</sub>TG<sub>2</sub>A<sub>2</sub>GAG<sub>3</sub>A-tamra, fluo = fluorescein, tamra = tetramethylrhodamine. <sup>c</sup> Not determined.



increased with increasing ligand-to-DNA ratio (LDR) (Fig. 2D). At the same time, the intensity of LD bands of the DNA bases decreased during the titration.

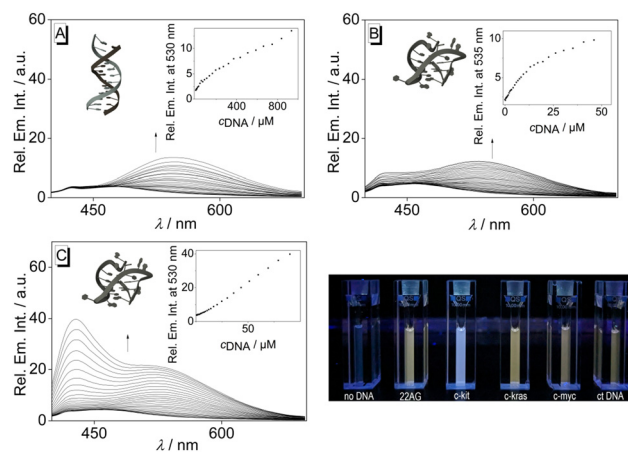
Overall, the photometric analyses indicated the association of the biphenyl **3** with nucleic acids. In particular, the changes in the absorption spectrum, namely hypochromism and the red shift of the absorption bands, are characteristic features of ligands that bind to duplex DNA<sup>30</sup> or G4-DNA.<sup>31</sup> Moreover, there are no large differences between the binding constants with the employed nucleic acids, which showed that there is no pronounced binding selectivity of the ligand towards a particular DNA form. In the case of duplex DNA, the additional LD-spectroscopic studies indicated an intercalation of the ligand, as confirmed by the negative LD bands of the ligand, which are characteristic of DNA intercalators.<sup>32</sup> Accordingly, the observed binding constant of **3** with ct DNA is in a range commonly observed for cationic DNA binders.<sup>33</sup> In contrast, the affinity of this ligand towards G4-DNA is relatively low as compared with the ones of known cationic ligands, which usually have binding constants of  $K_b > 10^5 \text{ M}^{-1}$ .<sup>34</sup> In most cases, ligands that intercalate into duplex DNA tend to bind with G4-DNA by terminal  $\pi$  stacking to the quadruplex unit.<sup>35</sup> Therefore it is assumed that the ligand **3** binds with essentially the same binding mode to **c-kit**, **c-kras**, **c-myc**, and **h-ras1**, and that the low binding constants reflect a somewhat hindered fit of the ligand structure to the binding site. The latter may be caused by the three-dimensional steric demand of the ligand as it does not have the required planar structure.

In addition, the stabilization of the dye-labeled G4-DNA forms **F21T**, **FkitT**, **FkrasT** and **FmycT** towards thermally induced unfolding after binding of ligand **3** was investigated by thermal DNA-denaturation experiments (Table 2 and Fig. S4, cf. ESI†).<sup>36</sup> Analysis of the melting temperature after ligand binding revealed a moderate stabilization of the quadruplex forms **F21T** ( $\Delta T_m = 6^\circ \text{C}$ ) and **FkrasT** ( $\Delta T_m = 6^\circ \text{C}$ ), whereas **FkitT** ( $\Delta T_m = 3^\circ \text{C}$ ) and **FmycT** ( $\Delta T_m = 2^\circ \text{C}$ ) were only slightly stabilized under the same conditions. These results indicated that upon binding the ligand **3** stabilizes the G4-DNA forms to a different extent depending on their topology. Notably, the shifts of the melting temperatures,  $\Delta T_m$ , of the different G4-DNA forms in the presence of ligand **3** do not correlate well with the binding constants. This apparent contradiction has been observed frequently during studies of ligand-quadruplex interactions and may be explained by different dependencies of equilibrium constants and Gibbs free energies on the temperature,<sup>6a</sup> specifically as the binding constant was determined at temperatures below the respective DNA melting temperature. Furthermore, it was shown exemplarily that the stabilization of the G4-DNA **FkitT** by ligand **3** did not significantly change in the presence of the double-stranded DNA **ds26** [ $d(\text{CA}_2\text{TCG}_2\text{ATCGA}_2\text{T}_2\text{CGATC}_2\text{GAT}_2\text{G})$ ; ( $\Delta \Delta T_m = -0.8^\circ \text{C}$ )], which indicated that the ligand binds selectively to the G4-DNA in competition with duplex DNA.<sup>37,38</sup>

To assess the influence of the association of ligand **3** with DNA on its emission properties, fluorimetric titrations with duplex and quadruplex DNA were conducted. In aqueous

buffer solution, compound **3** showed a weak, hardly detectable emission band with a maximum at 460 nm. Upon addition of ct DNA, **22AG**, **c-kit**, **c-kras**, **c-myc**, and **h-ras1**, the emission intensity of the bound ligand **3** increased; however, the development of the emission bands varied distinctly with the different DNA forms (Fig. 3A, B, C and Fig. S1, cf. ESI†). Specifically, the contribution of the two emission maxima resulting from the dual fluorescence of the ligand **3** (see above) depends on the respective DNA form. Upon addition of ct DNA, only one clear emission band developed with a maximum at 530 nm and with an increase of emission intensity by a factor of 2 (at ligand : DNA = 2), whereas the band at ca. 460 nm remained very weak. A similar development of the red-shifted fluorescence band at 535–538 nm with an intensity increased by a factor of 4–5 was observed with G4-DNA **22AG**, **c-kras**, **c-myc**, respectively. But in these cases, the intensity of the initial blue-shifted band also gradually increased, albeit to a smaller extent than the red-shifted band. In sharp contrast, the addition of **c-kit** and **h-ras1** to compound **3** led to a much stronger increase of the emission intensity of the blue-shifted band at 431 nm (**c-kit**) and 450 nm (**h-ras1**) by a factor of 5 and 10 (at ligand : DNA = 2), along with the rise of the red-shifted band with smaller intensity. In each case, the fluorescence light-up effect of the ligand **3** upon addition of DNA can be followed by the naked eye (Fig. 3). Moreover, the solutions with different DNA forms could be distinguished by different emission colors, whereas the distinct clear blue emission in the presence of G4-DNA **c-kit**, and **h-ras1** stood out as a special indicative element.

The increased emission intensity of compound **3** after addition of DNA is most likely caused by the limited conformational freedom of the ligand within the DNA binding site,<sup>3b</sup>



**Fig. 3** Fluorimetric titration of **3** ( $c = 20 \mu\text{M}$ ) with ct DNA (A), **22AG** (B), and **c-kit** (C); (A: in BPE buffer,  $c_{\text{Na}^+} = 16 \text{ mM}$ ,  $\text{pH} = 7.0$ , B and C: in K-phosphate buffer,  $c_{\text{K}^+} = 95 \text{ mM}$ ,  $\text{pH} = 7.0$ ). The arrows indicate the development of the absorption or emission bands during the titration. Inset: plot of the emission at  $\lambda = 330 \text{ nm}$  or  $\lambda = 431 \text{ nm}$  versus  $c_{\text{DNA}}$ : of ct DNA (A), **22AG** (B), and **c-kit** (C). Structures in insets taken from the Protein Data Bank (PDB, <https://www.rcsb.org/>; PDB ID: A, 2k0v; B, 1kf1; C, 6gh0).



which in turn suppresses the torsional relaxation of the biphenyl unit as non-radiative deactivation pathway of the excited molecule. Such a fluorescent light-up effect based on this mechanism has been frequently observed for cationic DNA binders that contain a biaryl substructure.<sup>21</sup> Moreover, this effect of the binding site on the fluorescence properties of the ligand may also explain the varying development of its emission bands upon association with the different DNA forms (Fig. 3). Specifically, the size and shape of the binding sites of the employed DNA forms vary distinctly,<sup>39</sup> so that the bound ligand experiences different degrees of restricted conformational freedom in each case. Such an effect of the binding sites on the emission intensity and energy has already been demonstrated with the structurally resembling compound 9-(4-dimethylamino)benzo[*b*]quinolizinium, which exhibits a fluorescence light-up effect with different emission maxima in duplex and quadruplex DNA.<sup>40</sup> Likewise, this influence of the binding sites on the emission color has been explained by the different conformational flexibility of the ligand within the binding sites.<sup>40</sup>

In the presence of ct DNA, the emission bands resemble the ones in glycerol and indicate the restricted free volume within the DNA binding site, which, in turn, should hinder conformational changes. Accordingly, the relaxation to the non-emissive conformation is suppressed under these conditions. But obviously, minor structural changes to a relaxed TICT/CS state are still possible, so that the related red-shifted emission band is also formed. In contrast, upon binding to G4-DNA a clear dual emission of the bound ligand was observed, which indicates that both the LE state and the TICT/CS state are populated to a significant extent within the respective binding sites. However, as a striking difference the contribution of each emission band varies with the different G4-DNA forms. And in the case of **c-kit** and **h-ras1**, the blue-shifted emission band dominates the overall spectrum, and this combination of bands results in an overall blue emission. For an explanation of this special effect, it is proposed that this effect is caused by the exceptional binding pocket provided by the quadruplex structure.<sup>39c</sup> Specifically, in the case of **c-kit** the particular loop structures allow the formation of a cleft at the 3'-end of the quadruplex in which organic ligands may be accommodated.<sup>41</sup> Indeed, simple docking studies<sup>42</sup> with the ligand **3** and **c-kit** revealed that the biphenyl **3** may bind in this binding pocket (Fig. 4) and point to a suppression of conformational freedom of the biphenyl unit in this constrained environment. As a result, the emission occurs mainly from the LE state leading to a distinct blue emission.<sup>43</sup> Likewise, it is proposed that the conformational flexibility of ligand **3** is similarly restricted in the binding site of **h-ras1**. In this particular case, the available free volume is probably even more confined because the fluorescence light-up factor is larger. The "regular" terminal  $\pi$  stacking of the ligand at the other G4-DNA forms **c-kras**, **c-myc**, **22AG**, by contrast, leaves sufficient conformational freedom to give mainly the red-shifted emission of the relaxed state. Notably, these latter G4-DNA forms have different quadruplex structures regarding the

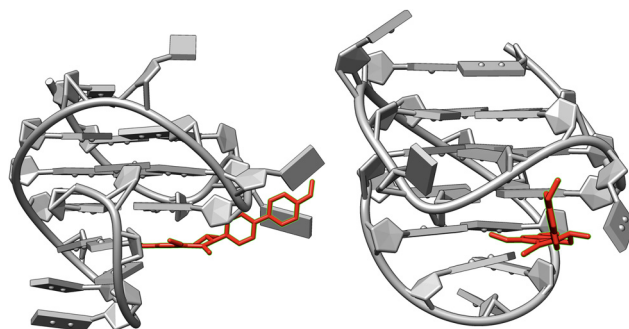


Fig. 4 Structure of the complex between ligand **3** and G4-DNA **c-kit** (PDB ID: 2O3M) as estimated from molecular docking (AutoDock Vina, UCSF Chimera 1.15).

DNA-strand orientation, namely parallel (**c-kras**, **c-myc**) and hybrid (**22AG**); and likewise, the quadruplex forms, with which a more pronounced blue shift was observed, have different strand alignment (**c-kit**: parallel, **h-ras1**: antiparallel). Therefore, it may be concluded that this particular structural parameter does not influence significantly the emission properties of the bound ligand **3**. At the same time, it cannot be excluded that the different polarity and/or availability of hydrogen bonds within the binding sites of the different G4-DNA forms also contributes to the varying emission properties of the bound ligand, because similar effects were observed in different solvents (Fig. 1). In this context, it should be emphasized that the different fluorescence response of the ligand **3** upon association with different DNA forms is not the result of a selective DNA recognition, because the binding studies revealed similar affinities to all tested DNA forms (see above). Instead, the varying emission properties are most likely caused by the different binding modes in combination with the high sensitivity of the dual emission of the biphenyl fluorophore on the surrounding medium.

## Conclusions

In summary, we have identified the cationic, donor-acceptor-substituted biphenyl structure **3** as a promising complementary motif of quadruplex DNA binders. It has, to the best of our knowledge, not been introduced to this research field so far. However, biaryl-type fluorophores are already known that may be used for fluorimetric DNA detection, with thioflavin T (ThT) being the most prominent example.<sup>44</sup> But these fluorescent probes do not exhibit dual emission, so that their response to DNA binding is a fluorescence light-up effect *without* a color change.

In the present case, neither the affinity nor the selectivity of the ligand towards a particular DNA form was especially pronounced. But still, the high sensitivity of the emission properties of the biphenyl unit, specifically the pronounced dual emission, to the surrounding medium may be used to detect and characterize different binding sites of G4-DNA by distinct



emission properties of the bound ligand. Specifically, sterically constrained binding sites, such as the one in **c-kit**, may be identified by the characteristic stronger contribution of the blue-shifted emission band. Therefore, it is proposed that similar cationic biphenyl derivatives with optimized structures for stronger binding affinities will figure as useful and efficient tools for the fluorimetric analysis of quadruplex DNA.

## Data availability

The data supporting this article have been included as part of the ESI.†

## Conflicts of interest

There are no conflicts to declare.

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