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# Bicolour fluorescent molecular sensors for cations: design and experimental validation

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Molecular entities whose fluorescence spectra are different when they bind metal cations are termed bicolour fluorescent molecular sensors. The basic design criteria of this kind of compound are presented and the different fluorescent responses are discussed in terms of their chemical behaviour and electronic features. These latter elements include intramolecular charge transfer (ICT), formation of intramolecular and intermolecular excimer/exciple complexes and Förster resonance energy transfer (FRET). Changes in the electronic properties of the fluorophore based on the decoupling between its constitutive units upon metal binding are also discussed. The possibility of generating fluorescent bicolour indicators that can capture metal cations in the gas phase and at solid–gas interfaces is also discussed.

## 1. Introduction

Detection of metal cations by means of molecular sensors has become a very active research field, with important applications in analytical chemistry,<sup>1</sup> materials<sup>2</sup> and environmental sciences,<sup>3</sup> and biological chemistry,<sup>4</sup> among other scientific areas. Very recently, this technique has emerged as a promising tool in particle physics since luminescence

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sensing of cations is very well-suited for the detection of special events such as neutrinoless double beta decay nuclear processes.<sup>5</sup>

Different techniques developed so far for the detection of metallic ions include atomic absorption spectroscopy, voltammetry, electrochemical analysis, inductively coupled plasma spectrometry, laser induced breakdown spectroscopy, neutron activation analysis and UV-Vis spectroscopy, among others.<sup>6,7</sup> However, fluorescence<sup>8</sup> is perhaps the most frequently used because of the relative simplicity of the associated devices, its very high sensitivity and its excellent spatial and temporal

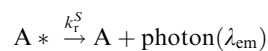
resolution.<sup>9–15</sup> Since in many samples different metal cations can coexist, the corresponding sensors must exhibit high selectivity. In this respect, fluorescence sensing arrays have been developed as a useful alternative based on ensembles of different less selective sensors that exhibit characteristic responses for different analytes.<sup>16</sup>

Many different types of fluorescent sensors able to detect metal cations have been described, including nanoparticles,<sup>17</sup> carbon-based quantum dots,<sup>18,19</sup> and dendrimers,<sup>20</sup> as well as metal–organic frameworks (MOFs),<sup>21</sup> and biomolecules such as DNA<sup>22</sup> and proteins.<sup>23</sup> These clustered and macromolecular sensors are not included in the present Perspective Article, which is focused on discrete fluorescent molecular sensors.

The fluorescence emission<sup>24</sup> spectrum of a given fluorophore is produced after excitation at a certain wavelength (normally in the UV region), followed by emission of photons at larger wavelengths. The Stokes shift expresses the difference between the maximum absorption wavelength  $\lambda_{\text{abs}}^{\text{max}}$  and the maximum emitted wavelength  $\lambda_{\text{em}}^{\text{max}}$ . If this difference is expressed in wavenumbers, the corresponding  $\Delta\bar{\nu}_{\text{Stokes}}$  is given by

$$\Delta\bar{\nu}_{\text{Stokes}} = \frac{1}{\lambda_{\text{abs}}^{\text{max}}} - \frac{1}{\lambda_{\text{em}}^{\text{max}}}$$

Since  $\lambda_{\text{em}}^{\text{max}} > \lambda_{\text{abs}}^{\text{max}}$ ,  $\Delta\bar{\nu}_{\text{Stokes}} > 0$ . The excitation process generates a transition from the ground state  $S_0$  of molecule A to the first singlet excited state  $S_1(A^*)$ . The decay from  $S_1(A^*)$  to  $S_0(A)$  is determined by the radiative kinetic constant  $k_r^S$



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and follows a usual first-order kinetics in the form

$$[{}^1\text{A}^*]_t = [{}^1\text{A}^*]_0 \exp\left(-\frac{t}{\tau_S}\right)$$

In the latter equation,  $[{}^1\text{A}^*]_0$  is the concentration of excited molecules at  $t = 0$ , which corresponds to the instant of the pulse light excitation, and  $\tau_S$  is the lifetime of the excited state  $S_1$ :

$$\tau_S = \frac{1}{k_r^S + k_{nr}^S}$$

where  $k_{nr}^S$  is the total kinetic constant associated with non-radiative processes that do not result in fluorescence emission.

The time-dependent intensity of the fluorescence emission  $i_F(t)$  after excitation at  $t = 0$  is proportional to the concentration of molecule A in the singlet excited state and to the kinetic constant associated with the fluorescence emission:

$$i_F(t) = k_r^S [{}^1\text{A}^*]_t = k_r^S [{}^1\text{A}^*]_0 \exp\left(-\frac{t}{\tau_S}\right)$$

It is important to note that the measured fluorescence intensity  $I_F$  is obtained from the total decay at different wavelengths and is expressed in arbitrary units (a.u., not to be confused with atomic units) that depend on the experimental conditions.

The fluorescence quantum yield for a given  $\text{A}^* \rightarrow \text{A}$  transition is the fraction of excited molecules  $S_1(\text{A}^*)$  that decay to the ground state  $S_0(\text{A})$  with emission of photons (*vide supra*):

$$\phi_F = k_r^S \tau_S = \frac{k_r^S}{k_r^S + k_{nr}^S}$$

Thus, the quantum yield gives the fraction of emitted photons during the total decay with respect to the absorbed photons. This magnitude gives crucial information to measure the efficiency of fluorescence emission. Another relevant parameter is the brightness<sup>4,23</sup> of the fluorescence emission  $B_F$ , which is the product of the quantum yield and the molar extinction coefficient  $\epsilon_{\text{abs}}$  at the excitation light wavelength.

$$B_F = \phi_F \epsilon_{\text{abs}}$$

All these combined magnitudes characterize the photophysical behaviour of a given fluorophore.

The key point in fluorescence sensing of different metallic cations is the change of these magnitudes upon coordination or binding of the sensor with the analyte. In a fluorescent monocolour sensor the emitted photons are of identical wavelengths  $\lambda_{\text{em}}$  in the free state and upon coordination or binding with the metal cation. This can occur *via* photoinduced electron transfer<sup>1,25</sup> (PET) phenomena *via* an electronic donation from one heteroatom of the sensor that quenches the fluorescence relaxation of the fluorophore in the free  $S_1$  state. Upon coordination, this donation is directed to the metal and the fluorescence decay takes place efficiently thus resulting in an off-on transition

(turn-on enhancement) on going from the free to the metal-coordinated state. Another less frequently observed PET consists of the reverse on-off process (turn-off quenching). In most cases, the PET perturbations of the relaxation processes result in preferential emission of photons with  $\lambda_{\text{em}}$  values similar for free and bound states, the only difference being the very low and very high intensities in the off and on states, respectively. Therefore, the larger the difference between the intensities of the off and on states, the higher the sensitivity of the sensor. These monocolour probes require very high intensity-based separation factors between their 'off' and 'on' states; such separation factors are challenging to attain, given various analyte-independent factors which affect the probe intensity (*e.g.* fluctuations in the excitation source, the effect of the microenvironment, variations in the probe concentration, *etc.*)<sup>26,27</sup> In this Perspective Article we rather focus on bicolor probes which rely on wavelength separation and are not affected by such factors.

Molecular bicolor sensors emit photons with different wavelengths in the free ( $\lambda_{\text{em}}$ ) and bound ( $\lambda'_{\text{em}}$ ) states. They can also be described as dual emission<sup>28</sup> ratiometric probes with signal changes associated with two independent or correlated emitters.<sup>27</sup> Depending on the magnitude of  $\lambda'_{\text{em}}$  with respect to  $\lambda_{\text{em}}$ , blue (hypsochromic) or red (bathochromic) shifts will be observed. Thus, both free and bound bicolor sensors can be active fluorescent emitters within an on-on' scheme. In this case, the sensitivity will be determined by the respective intensities and the corresponding photophysical parameters. Two relevant parameters are the wavelength difference  $\Delta\lambda_F = \lambda'_{\text{em}} - \lambda_{\text{em}}$  and the peak discrimination factor  $f_\lambda$ , defined as

$$f_\lambda = \frac{I'_\lambda - I_\lambda}{I_\lambda}$$

where  $I'_\lambda$  and  $I_\lambda$  are the intensities of the signals characterized by a wavelength  $\lambda$  at the bound and free states, respectively. This factor is an indicator of the expected signal cleaning of the metal-bound sensor with respect to its free unbound state, especially when the concentration of the metallic cation is very low (*vide infra*). Thus, the greater the separation dictated by  $f_\lambda$ , the lesser the overlap between both signals. Aside the intrinsic values of quantum yield, brightness and intensity, these last two factors will determine the sensitivity of the molecular sensor with respect to the background. This is particularly important in cation-tagging experiments with extremely low concentration of the metallic cation (eventually, only one!), in which the concentration of the free molecular bicolor sensor is therefore orders of magnitude more higher than that of the bound sensor.

Another important aspect associated with the design of molecular fluorescent bicolor sensors is the necessity of developing sensors able to capture cations in the gas phase. This in turn implies the exploration of supramolecular chemistry on solid-gas interfaces, a research field much less



developed than its in-solution analogues. The use of single cation identification has been proposed as a way to dramatically reduce the radioactive background<sup>29</sup> in xenon detectors looking for neutrinoless double beta decay. The  $^{136}\text{Xe}$  isotope decays into barium, emitting two electrons (and two anti-neutrinos in the standard process). Identification of a coincidence signal of the two electrons with the appropriate energy with the appearance in the active volume of a  $\text{Ba}^{2+}$  ion will be an unequivocal signature of the neutrinoless process and, consequently, a demonstration that neutrinos are their own anti-particles (Majorana<sup>30</sup> particles). Demonstrating this hypothesis would have deep consequences in particle physics and in cosmology.<sup>31,32</sup> A promising “wet” barium tagging experiment has been reported as a proof of concept.<sup>33</sup> In order to reproduce the required “dry” conditions imposed by these experiments, several interesting barium-tagging sensors have been reported so far,<sup>34,35</sup> all of them based on PET off-on fluorescence emission. Our approach to barium-tagging is instead based on the development of suitable on-on’ bicolor fluorescent sensors, which benefit from a wavelength-shifting mechanism.

The many contributions to the development of ratiometric molecular fluorescent sensors for cations have been comprehensively described in leading reviews and book chapters in the area.<sup>9,26,27,36–42</sup> In this Perspective Article, we shall present and discuss the main features and design criteria of molecular bicolor fluorescent sensors for metallic cations, intended for the detection of extremely low cation concentrations (eventually only one) on a surface covered by unbounded sensor molecules, in dry media. In the following sections, we will describe the main modes of sensor-cation interactions and the electronic phenomena responsible for the change in the fluorescence spectra upon metal coordination or after interaction with the cationic centre. We will also report on recent results obtained in our laboratories on barium tagging in dry media involving solid-gas phases.

Bicolor fluorescent probes can be considered a special type of the so-called single-fluorophore ratiometric probes with two analyte-sensitive emission signals, as they change the emission intensity and wavelength upon interaction with the analyte. They can also be described as dual emission materials composed of two independent emitters (free and bound) in equilibrium.<sup>28,43</sup> In general, these molecules incorporate different components that include a synthetic ionophore (the cation recognition-binding domain), the fluorophore and, optionally, a spacer and a linker to anchor the sensor with suitable surfaces or biomolecules. The covalent connections among these components can be different, thus giving rise to different sensor architectures, the most usual ones being those shown in Fig. 1A.

The binding groups (ionophores) are responsible for the efficient and preferably highly selective capture of the metal cation and can consist of chelating groups<sup>4</sup> such as polycarboxylic acids **B1**, podands **B2**, coronands such as crown ethers<sup>44–46</sup> **B3** and cryptands **B4**. Other ionophores<sup>47,48</sup> such as calixarenes, cyclodextrins and pillararenes, not shown in Fig. 1B, have been described.

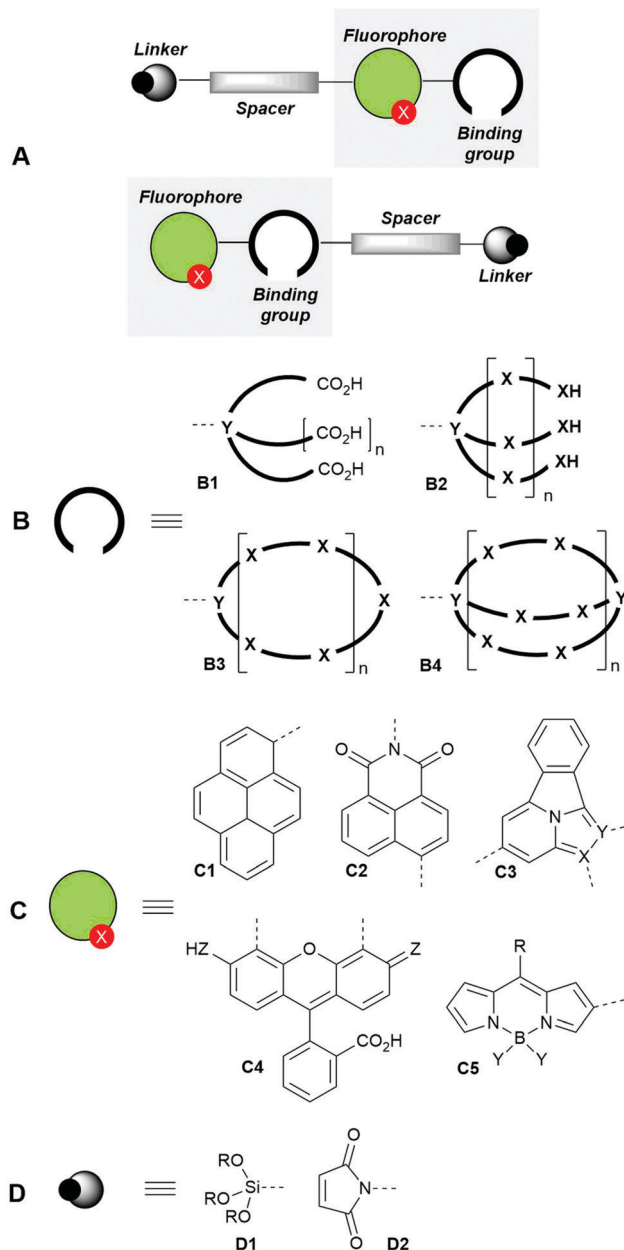


Fig. 1 General design patterns of fluorescent molecular sensors for cations (A), showing their main components: binding groups (ionophores, B), fluorophores (C) and linkers (D).

The structures of fluorophore components (Fig. 1C) are very varied<sup>4</sup> and comprise aromatic hydrocarbons such as pyrenes **C1**, and polyheterocyclic aromatic moieties such as naphthylimides **C2**, tetracyclic structures based on imidazo[1,2(5*a*)]pyridines<sup>49,50</sup> **C3**, fluorescein derivatives **C4** and BIODIPY-based structures **C5**, among others.

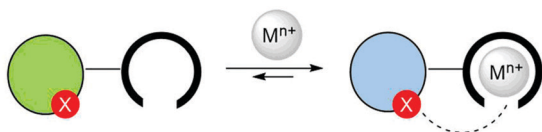
When required, the spacers can consist of aliphatic linear chains or polyethylene glycol (PEG) derivatives, which can intercalate with aromatic hydrocarbons or heterocycles. Finally, the linkers can be trialkoxysilyl derivatives **D1** (very useful for anchoring the fluorescent molecular sensors to solid surfaces) or maleimides **D2**, which can bind – with low selectivity – cysteine-containing



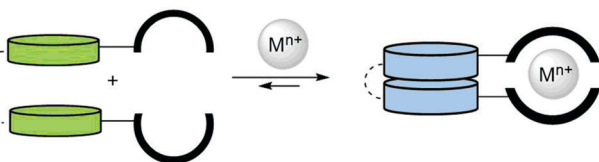
## A. Chelation-based bicolour sensors

### A.1. Fluorophore + chelating group

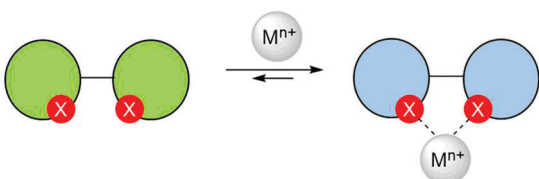
– Monomeric



– Dimeric



### A.2. Chelating fluorophore



## B. Reactivity-based bicolour sensors

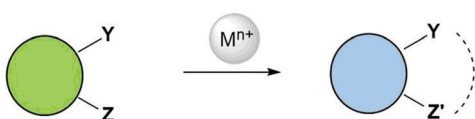


Fig. 2 Main types of bicolour sensors classified according to their structure and interaction with the metal. Different chelation modes (A, monomeric and dimeric chelating groups, as well as chelating fluorophores) are shown. Bicolour sensors in which the change in the fluorophore is generated by a chemical reaction with the metallic ion are also shown (mode B). Optional spacer and linker components are omitted.

proteins or biothiols, thus giving rise to the corresponding conjugated derivatives.<sup>51</sup>

Molecular fluorescent bicolour sensors can change the intensity and emission wavelength of their fluorescence signals in different ways after interacting with metal cations (Fig. 2). When an ionophore is present, metal binding can modify the photophysical properties of the sensor *via* formation of monomeric and dimeric complexes (Fig. 2, mode A.1). In some cases, when the fluorophore itself is able to bind directly the metallic cation, the photophysical properties of the chelate can be different to those of the free fluorophore (Fig. 2, mode A.2), thus yielding a bicolour sensor. In many cases, a heteroatom X possessing a suitable lone-pair present in the fluorophore can also bind the metallic centre thus forming a complex in which *both* the synthetic ionophore and the heteroatom are bound to the cationic analyte (Fig. 2, note the dashed line in monomeric complex A.1). Finally, reactive functional groups decorating the fluorophore can react selectively with the metal to be transformed into a different fluorophore, thus giving rise to an ionophore-free fluorescent bicolour sensor (Fig. 2B). These considerations apply to not only single emitters from one

excited electronic state, but also to dual emitters with a sole fluorophore possessing two emitting states and dual emitters with two independent or correlated emission centres.<sup>28</sup>

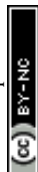
## 2. Mechanisms of bicolour sensing

### 2.1. Intramolecular charge transfer (ICT)

If changes in the electronic structure of the fluorescent bicolour sensor are considered, one possibility consists of modifying the polarization patterns between substituents of the fluorophore in the free and bound states. This process is named Photo-induced Charge Transfer (PCT), which almost in all cases takes place intramolecularly. Therefore, the latter process will be referred to as Intramolecular Charge Transfer (ICT). In these systems, an electron-releasing group (ERG), which acts as a donating group, transfers a certain electron density to an electron-withdrawing group (EWG), which is the acceptor of this charge transfer. This process involves complex electronic interactions. Within a simplified scheme involving delocalized frontier molecular orbitals, two situations can be distinguished, depending on the proximity of ERG and EWG moieties to the metallic cation. Of course, specific electronic interactions can be important, especially when there is a direct union between the fluorophore and the metal ion. When the ERG is close to the  $M^{n+}$  centre bound to the ionophore, the two-electron  $ERG \rightarrow M^{n+}$  interaction will compete with the  $ERG \rightarrow EWG$  donation thus increasing the  $\nu'_{em}$  frequency associated with the fluorescence  $S_1 \rightarrow S_0$  transition (Fig. 3A). Therefore, a blue shift will be observed, in many cases with a certain decrease in brightness. Alternatively, when the EWG is closer to the cation, the  $ERG \rightarrow EWG$  donation will increase in magnitude and the  $EWG-M^{n+}$  interaction will result in the lowering of the lowest unoccupied molecular orbital (LUMO) energy of the fluorophore, thus resulting in a lower  $\nu'_{em}$  frequency and the corresponding red shift, usually associated with a brighter signal (Fig. 3B). In both cases, the sensitivity of the ICT sensor correlates with the respective differences in brightness, as well as with the peak discrimination factors (*vide supra*).

### 2.2. Excimers

Another way to obtain radiometric bicolour sensors consists of designing fluorophores that, upon binding the metallic cation through the adequate ionophore, form dimers (homo- or heterodimers) that are more stable in the  $S_1$  excited state. Thus, an excimer (excited dimer) originates from the interaction of an excited fluorophore with a nearby congener in the ground state. If the two interacting units are non-identical, the whole chemical entity is named an exciplex (excited complex). The two units can be connected by a spacer and incorporate their respective incomplete ionophore components that, upon coordination with the metallic cation, generate the optimal binding group (Fig. 4A). An analysis of the frontier molecular orbitals (FMOs) of the monomeric units of the sensor at the  $S_0$  and  $S_1$  (Fig. 4B) states shows that the energy levels of the FMOs of the excimer (or the exciplex) are modified upon dimerization



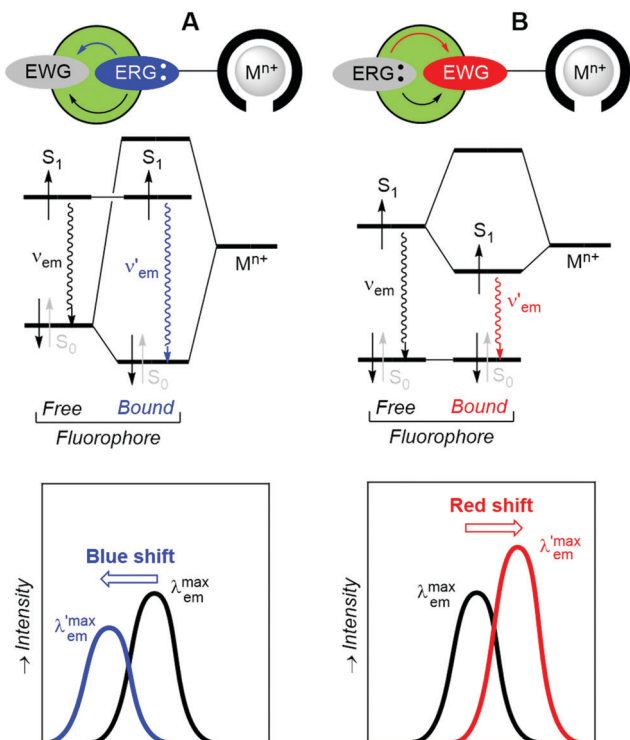


Fig. 3 Schematic representation of intramolecular charge transfer processes (ICT) that modify the fluorescence emission of bicolour sensors in the presence of cations  $M^{n+}$ . (A) Blue shift effect of an electron-releasing group (ERG) close to the ionophore. (B) Red shift effect of an electron-withdrawing group (EWG) close to the ionophore.

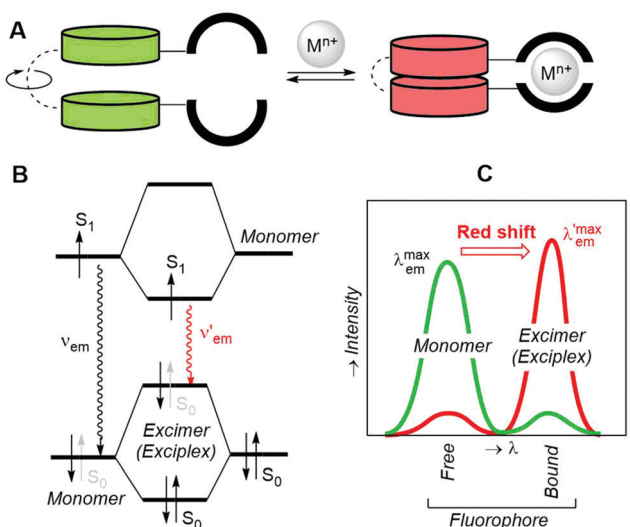


Fig. 4 (A) Cartoon showing the formation of excimers/excplexes in the presence of a metallic cation. (B) Qualitative diagram showing the distortion of the FMO energy levels in the  $S_1$  state of the excimer with respect to those of the monomer. (C) Red shift of the emission spectra of monomers and excimers/excplexes associated with the diagram shown in (B).

and usually result in a shorter HOMO–LUMO gap (HOMO stands for the highest occupied molecular orbital).<sup>52</sup> In turn,

this shrinkage in energy levels generates a red shift that permits to discriminate the free and bound states of the sensor. The binding ability of the partial ionophores in the free monomeric state and the stability of the excimer/excplex can result in a background signal that, depending upon the respective intensities, permits to establish an accurate measurement of the radiometric response (Fig. 4C). In other cases, the excimer/excplex is relevant in the absence of the metallic cation, whose coordination results in the total or partial loss of fluorescent response associated with excimer inhibition promoted by cation coordination.

### 2.3. FRET

A related phenomenon in which the emission involves an interaction between two fluorophores (Fig. 5A) is the Förster resonance energy transfer (FRET).<sup>53–55</sup> This process consists of a non-radiative excitation transfer<sup>56</sup> from the excited donating (D\*) group to the acceptor group (A) in its ground state (Fig. 5B). The latter group relaxes *via* an  $A^*(S_1) \rightarrow A(S_0)$  process thus emitting with a different wavelength (Fig. 5C).<sup>57</sup> Any FRET requires appropriate relative orbital energies associated with the FMOs of D and A. In the case of ratiometric bicolour sensors, the FRET is usually observed when the sensor is bound to the cationic centre. It is noteworthy that the distance  $R$  required for FRET is significantly larger than the average values of the combined van der Waals radii of the D and A groups. The Förster distance  $R_0$  is defined as the distance at which the FRET efficiency  $\phi_{\text{FRET}} = [1 + (R/R_0)^6]^{-1}$  is 0.5. For instance, if D = pyrene and A = coumarin,  $R_0 = 3.9$  nm.<sup>58</sup> The FRET efficiency also depends on the relative orientation of D and A, as well as

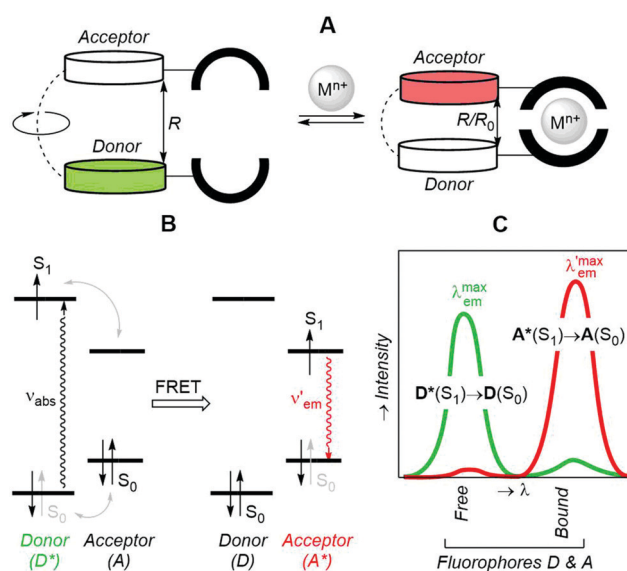


Fig. 5 (A) Schematic representation of a fluorescent sensor incorporating donor and acceptor units.  $R_0$  is the Förster radius (see the text). (B) Förster-resonance electron transfer (FRET) associated with the frontier molecular orbitals of the donor and the acceptor. (C) A representation of the fluorescence emission spectra of the FRET sensor in the free and bound states.



on the overlap between the emission spectrum of D and the absorption spectrum of A.

#### 2.4. Two-component fluorophores

In the previous sections, we have presented bicolour sensors based on fluorophores that act as indivisible photoactive ensembles. Another possibility consists of splitting a given fluorophore into two separate rotatable units connected by a single covalent bond (Fig. 6). In turn, this two-unit fluorophore **1** is covalently attached to an ionophore. When a metallic cation is captured by the latter binding moiety, the nature of the interaction is reinforced by additional coordination with a heteroatom of fluorophore **1**, thus yielding an ICT-complex (Fig. 2, A.1 pattern) that in addition incorporates a  $\pi$ -cation interaction<sup>59</sup> between the metallic centre  $M^{n+}$  and the  $Ar_2$  group gathered in Fig. 25. This  $Ar_2-M^{n+}$  complex promotes a  $Ar_2 \rightarrow M^{n+}$  charge transfer, as well as a rotation of  $Ar_2$  with respect to the polyheterocyclic  $HetAr_1$  unit. The outcome of these metal-induced changes is the decoupling between the  $HetAr_1$  and  $Ar_2$  groups. Therefore, the new  $HetAr_1$  fluorophore **2** thus formed (Fig. 25) is less conjugated than its predecessor fluorophore **1** in the unbound state and will emit photons with shorter wavelength. If an optical filter is installed in the detection system to hide the emission associated with fluorophore **1**, this blue shift should permit an essentially background-free signal upon cation complexation if the peak discrimination factor  $f_\lambda$  is large enough.

### 3. Design and validation of bicolour probes

All of the above discussed bicolour sensors have been classified in terms of the chemical interactions involved

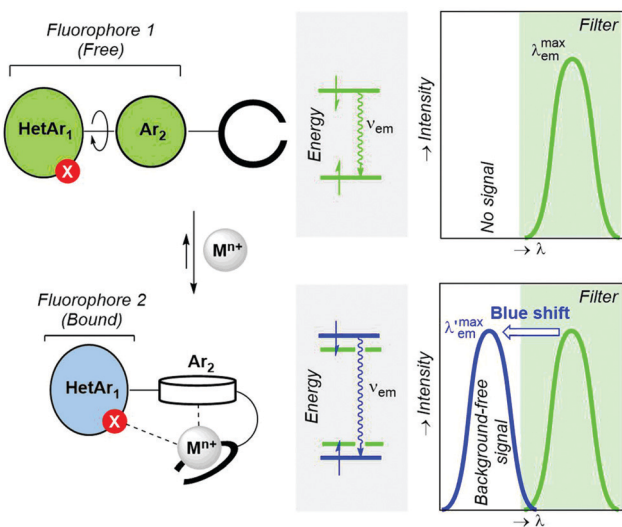


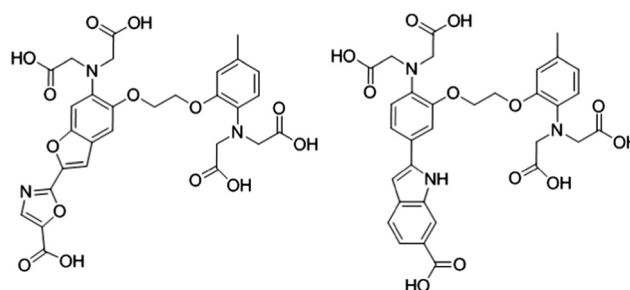
Fig. 6 Design criteria of two-component fluorescent bicolour indicators (FBI) based on the decoupling of one component of the starting fluorophore upon metal coordination. The metal-induced hypsochromic shift and the recording of a background-free signal by means of an optical filter are also highlighted.  $HetAr$  and  $Ar$  denote polyheterocyclic aromatic and aromatic groups, respectively.

(fluorophore–ionophore, direct chelation by the fluorophore and a chemical reaction with the cation) or the electronic effects that promote the change in the wavelength of the emitted photons (ICT, excimer/excplex, FRET and related phenomena). In the following sections, an overview of selected examples of these strategies for the construction of bicolour fluorescent probes for cations is presented and discussed.

#### 3.1. ICT-Based sensors

Many review articles and books describe the photo-induced ICT phenomenon and the potential applications of ICT-based compounds.<sup>24,60</sup> In polar solvents, rearrangement of the solvation shell around the ICT excited state produces its stabilization and the concomitant destabilization of the landing ground state, raising its energy level.<sup>41</sup> Therefore, these compounds may show dual emission: a peak in the most energetic region of the emission spectra that originates at the locally excited state (LE), and a red-shifted signal from the solvent-stabilized ICT excited state. Additionally, if the donor and the acceptor are connected through a rotating  $\pi$ -electron bridge, a relaxation of the ICT excited state can occur through an intramolecular rotation, decoupling the orbitals of the donor and acceptor, forming a highly polar excited state. This phenomenon is referred to as “twisted intramolecular charge transfer” (TICT), and it is characterized by extra-large Stokes shifts in polar solvents.

Both ICT- and TICT-based fluorophores are suitable scaffolds for the construction of bicolour probes for cations by integrating a cation receptor either on the ERG or on the EWG sites of the fluorophore (Fig. 3). The simplicity of the design makes ICT-based sensors the most widely used ratiometric probes for cations. Among the earliest examples are **Indo-1** and **Fura-2** (Fig. 7), developed in 1985 by Tsien and collaborators.<sup>61</sup> Nowadays, **Indo-1** and **Fura-2** are the leading compounds of a series of commercially available fluorescent ratiometric probes for  $Ca^{2+}$  and other cations.<sup>62</sup> They combine a stilbene skeletal fluorophore (reinforced by extra heterocyclic bonds) with the well-established BAPTA fragment as an ion-recognition site. These molecular probes are characterized by a strong fluorescent ratiometric response to  $Ca^{2+}$ . Because of the inhibition of the ICT upon coordination of the  $Ca^{2+}$  to the



	Fura-2	Fura-2 + $Ca^{2+}$	Indo-1	Indo-1 + $Ca^{2+}$
$\lambda_{abs}$	362 nm	335 nm	349 nm	331 nm
$\lambda_{em}$	512 nm	505 nm	485 nm	410 nm

Fig. 7 **Fura-2** and **Indo-1** ICT-based sensors for  $Ca^{2+}$ .



aniline nitrogen of the BAPTA receptor, a blue-shifted fluorescence emission is observed (Fig. 3A). One of the main limitations of these probes was the need for UV-light excitation. It was circumvented using two-photon absorption (TPA) microscopy, which permitted the study of cation distribution in live tissues.<sup>63</sup>

More recent ICT-based sensors are based on fluorophores that can be excited in the visible region. In 2019 Woodruff and O'Halloran synthesized a family of BODIPY-based fluorophores containing an appended *N,N,N'*-tris(2-pyridylmethyl)-1,2-ethanediamine (trispicen) chelator as a Zn<sup>2+</sup> receptor site.<sup>64</sup> Among them, **ZincBY-4** (Fig. 8), containing a pending acetamide on the 5-position of the BODIPY core outperformed other derivatives due to intramolecular hydrogen bonding between the amide hydrogen and the fluorine atoms of the BF<sub>2</sub> centre, thus increasing the Zn<sup>2+</sup> affinity. In polar solvents, the ICT-based fluorescence of the free probe is strongly quenched, whereas a blue-shifted emission is recovered upon Zn<sup>2+</sup> chelation due to disruption of the ICT process.

In 2017, Han, Zang and colleagues published ratiometric probes for Cd<sup>2+</sup>, also based on cation-induced ICT depletion, characterized by a long wavelength fluorescence emission (**Me<sub>4</sub>BOPHY-BPA** in Fig. 8).<sup>65</sup> The sensor consisted of a tetramethylated BOPHY fluorophore conjugated to bis(pyridine-2-yl) amine (BPA) podand acting as a Cd<sup>2+</sup> selective receptor. Cd<sup>2+</sup>-Chelation caused a clear hypsochromic shift of the fluorescence emission revealing the characteristic yellow emission of the **Me<sub>4</sub>BOPHY** dye. In 2019, Zeng *et al.* pushed this concept further developing a Hg<sup>2+</sup>-probe (**L** in Fig. 8) with NIR emission in both the holo and the apo form.<sup>66</sup> This sensor consists of a chromenyl-cyanine dye containing a conjugated NS<sub>2</sub>O<sub>2</sub> Hg<sup>2+</sup>-chelating unit. The strong ICT process between the N-atom of the cation receptor and the fluorescent dye, together with the NIR emission of the naked fluorophore, permitted the authors to construct this ratiometric NIR sensor operative under visible-light excitation.

The clear advantages of TPA microscopy compared to conventional one-photon absorption (OPA) microscopy for the analysis of cation distribution in live cells, tissues, and organisms (*i.e.* improved penetration, reduced toxicity, reduced autofluorescence, and minimal light scattering) drove the development of molecular probes optimized for TPA.<sup>67–69</sup> When considering ICT-based sensors, this has been pursued by binding the cation recognition site to the acceptor domain of an ICT-fluorophore. Therefore, upon metal binding, rather than ICT depletion (as observed in the examples mentioned above) the polarization of the ICT excited state will increase. This probe configuration should not only produce a bathochromic shift in the fluorescence emission of the cation-bound probe (Fig. 3B), but also enhance the TPA cross-section, and the brightness of the fluorescence emission.<sup>69</sup>

Based on this premise, in the last few years a series of ratiometric TPA probes appeared in the literature. Several Zn<sup>2+</sup>-selective sensors have been developed based on a BPA podand bound to different fluorophores. Representative examples are shown in Fig. 9.<sup>70–73</sup> **MPVQ**-series, initially developed by Meng's

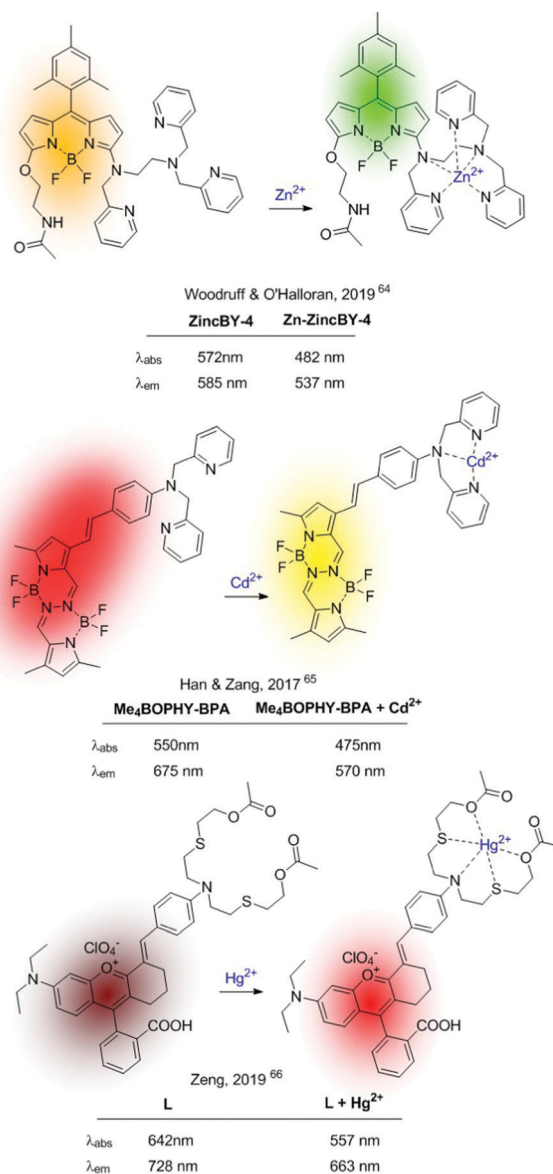


Fig. 8 Recent examples of metal-induced ICT depletion probes.

group,<sup>73</sup> consisted of BPA-appended quinoline as the fluorescent skeleton with a conjugated donor on the 6-position, whereas **chromis-1** contains 2-arylthiazole as a push-pull fluorophore. In all cases, a red-shifted fluorescence was observed in the presence of Zn<sup>2+</sup>, which was accompanied by an increased TPA cross-section.

More complex ratiometric sensors for cations, optimized for TPA, have been constructed using centrosymmetric donor- $\pi$ -acceptor- $\pi$ -donor quadrupolar configurations, in which the cation recognition site is integrated into the central acceptor unit, thus constituting a chelating fluorophore (Fig. 2, pattern A.2). The **GBC** probe (Fig. 10), developed in 2014,<sup>74</sup> was a ground-breaking example of sensors based on this structural motif and inspired further developments.<sup>75,76</sup> The cation recognition is achieved through a central bipyridine unit with two appended carbazole donors linked through vinylic  $\pi$ -electron



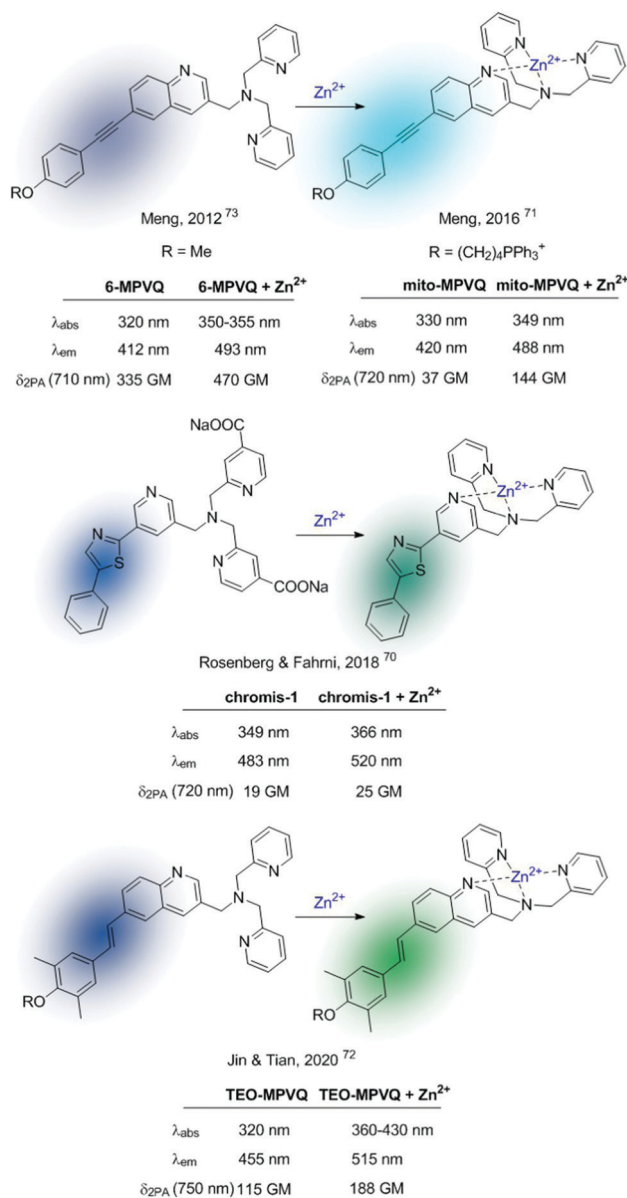


Fig. 9 TPA ICT-based ratiometric probes using BPA as a selective Zn<sup>2+</sup> receptor.

bridges. GBC also contains oxyethylene side chains to increase its solubility in aqueous media and its cell permeability. The emission red shift and the enhanced TPA observed upon metal binding were initially attributed to ICT, because of the increased electron-withdrawing character of the Zn-coordinated bipyridine unit. The mechanism responsible for the large TPA characteristic of these systems has been further analysed from a theoretical perspective aiming at establishing structure–property relationships.<sup>77–79</sup> These studies revealed that fluorescence emission under two-photon excitation originates from the S<sub>2</sub> excited state rather than from S<sub>1</sub>. Additionally, coordination of the Zn<sup>2+</sup> cation requires the rotation of the central bipyridine unit from the *trans* to the *cis* coplanar form. This conformational change results in an increase of the dipole and quadrupole moments of

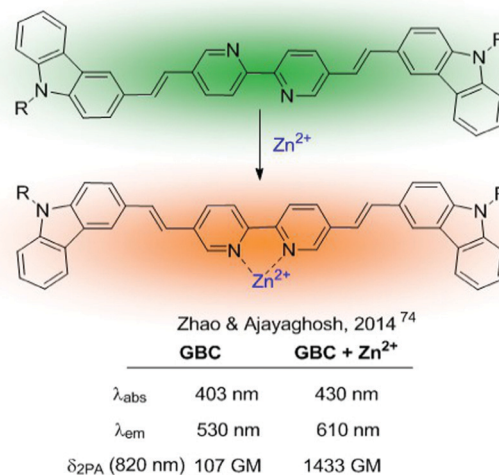


Fig. 10 GBC ratiometric probe for Zn<sup>2+</sup>.

the molecule by a loss of the centre of inversion present in the most stable conformation of the unbound probe.<sup>77</sup>

**3.1.1. Multi-target sensors.** The electronic origin of the shift in the emission observed in ICT-based sensors makes it feasible to develop multi-target sensors, able to discriminate between similar cations due to the different emission colours of their corresponding complexes. The cation-dependent polychromic emission observed in some ICT-based sensors has been attributed to differences in electronic affinities,<sup>80</sup> binding modes,<sup>81</sup> or ion sizes.<sup>81</sup> Representative examples are shown in Fig. 11. It is worth pointing out that sensor **L2** showed an extremely low detection limit for Cd<sup>2+</sup> (0.24 nM), and that both **L1** and **L2** were tested on solid supports, as the active components of indicator paper.<sup>82</sup> In the case of **HQZn**, the cation-dependent emission colour of this TPA sensor, together with the higher affinity of the sensor for Cd<sup>2+</sup> compared to Zn<sup>2+</sup>, permitted the authors to use the Zn-chelated ligand as a Cd<sup>2+</sup> sensor, eliminating the recurrent problem of Zn<sup>2+</sup> interference during Cd<sup>2+</sup> detection in biological systems.<sup>83</sup>

**3.1.2. Reactivity-based ICT bicolour sensors: chemodosimeters.** Highly efficient ratiometric probes for cations based on ICT can also be constructed relying on cation-induced irreversible bond-cleavage/bond-formation reactions on the molecular probe (Fig. 2, pattern B). Such cation-mediated chemical changes in the probe can produce an enhancement or depletion of the charge-transfer process, which eventually affects the colour of the fluorescence emission. The examples range from probes based on the chemical reactivity of the cation with specific functional groups to form a cation-coordinated ICT fluorophore,<sup>84,85</sup> to those based on cation-mediated umpolung reactions on the probe (usually protection/deprotection of functional groups) releasing a cation-free ICT fluorophore.<sup>86–92</sup> Selected examples are shown in Fig. 12. One of the earliest examples was developed by Lin *et al.* in 2009.<sup>88</sup> The probe was based on a phenanthroimidazole dye containing an appended electron-rich 1,3-dithiane. Cu<sup>2+</sup>-Promoted desulfurization of the thioacetal rendered a formyl EWG, with the



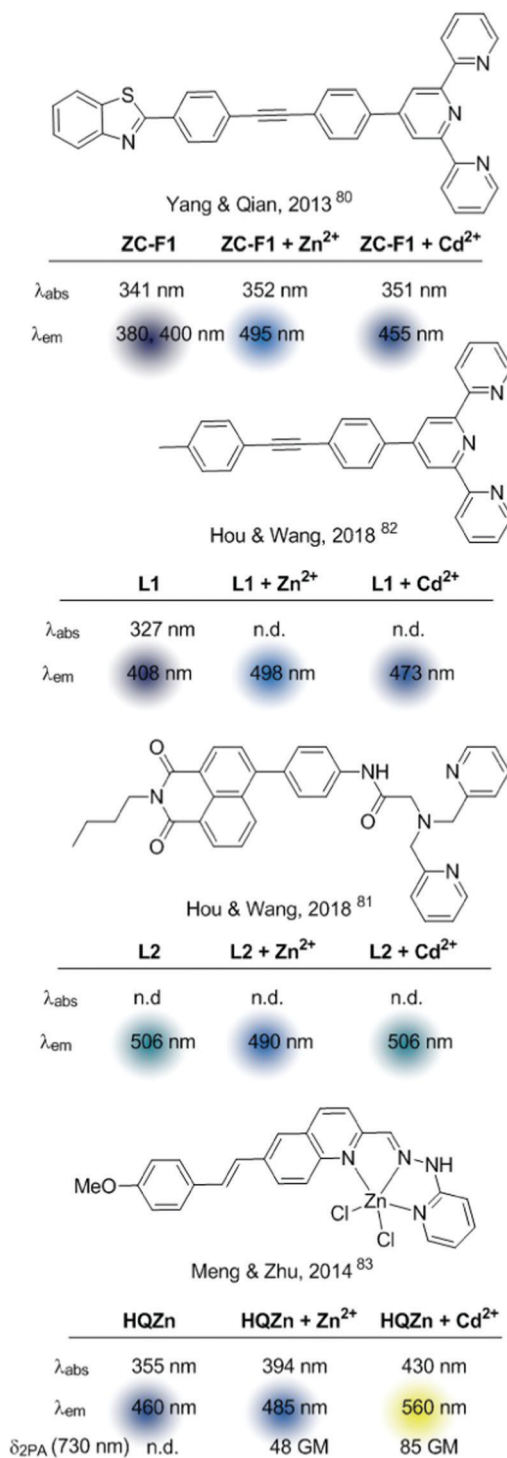


Fig. 11 Multi-target sensors based on ICT.

concomitant bathochromic shift in the fluorescence emission due to ICT between the phenanthroimidazole (ERG) and the formed aldehyde (EWG). More recently, an acyclic thioacetal version of the same probe has been published for Hg<sup>2+</sup>-sensing.<sup>87</sup> This cation-mediated deprotection strategy has been extended to the construction of a large number of ICT-based chemodosimeters using different fluorescent dyes (dansyl, coumarin or naphthalimide,

among others) (see examples in Fig. 12).<sup>86,88–92</sup> Among these probes, one can distinguish two types depending on the effect (ICT enhancement or depletion) that the cation-induced reactivity produces on the probe. The former (metal-induced ICT enhancement) are *a priori* more appropriate for TPA applications. In a very recent example, Kim and Yoon developed **NAP-PS** (Fig. 12), a TPA probe for sensing intracellular Hg<sup>2+</sup> in live cells and tissues.<sup>86</sup> In this example, the presence of Hg<sup>2+</sup> triggers the cleavage of the thiophosphinate P–O bond, forming the ICT fluorophore **NAP-O<sup>-</sup>**. Additionally, grinding solid **NAP-PS** in the presence of Hg(ClO<sub>4</sub>)<sub>2</sub> also produced a red-shift in the fluorescence emission of the resulting powder, demonstrating its response even in the solid state (Fig. 13).

All chemodosimeters described above are based on metal-induced bond-cleavage reactions. Alternatively, they can also be constructed using the so-called “covalent assembly principle”. It refers to those probes in which the analyte triggers a cascade reaction on the probe, thus furnishing the conjugated backbone of a push–pull fluorescent dye.<sup>93</sup> A coumarin-based probe (**ATC-Hg**), shown in Fig. 14, is a recent example of such a strategy.<sup>94</sup> Hg<sup>2+</sup> induced the deprotection of 1,3-dithiolane to a formyl group that condensates intramolecularly with the nearby amino group. This cascade reaction renders a heterocyclic aromatic donor– $\pi$ –acceptor ICT fluorescent dye. This probe showed high selectivity to Hg<sup>2+</sup> and MeHg<sup>+</sup>, with sensitivities of 27 nM and 5.8  $\mu$ M, respectively.

Ratiometric sensors can also be constructed using metal-induced cascade reactions that reduce the conjugation on the probe. The group of Emrulloğlu employed this strategy for the construction of BODIPY-based ratiometric probes for sensing Au<sup>3+</sup> and Hg<sup>2+</sup> (Fig. 15).<sup>95,96</sup> Although strictly speaking these sensors cannot be considered ICT-based probes, it is convenient to include them here under the chemical pattern B shown in Fig. 2. In these examples, metal-mediated activation of the triple bond and subsequent intramolecular cyclisation and protonolysis generates the corresponding unconjugated BODIPY-furan fluorophores, as evidenced by the characteristic green emission after the cation-promoted reaction.

### 3.2. Excimer/excimer-based sensors

An excimer was defined by Birks as “a dimer which is associated in an excited electronic state and which is dissociative (*i.e.* would dissociate in the absence of external restraints) in its ground electronic state”<sup>97</sup> (Fig. 4). Therefore, excimer formation is an irradiation-induced phenomenon, strongly dependent on the fluorophore–fluorophore distance. In general, in this kind of bicolour sensor the fluorophore is a polyaromatic system, being pyrene the one most frequently used. Since excimer emission is often red-shifted compared to the fluorescence of the monomer, excimer-based sensors can be used for the construction of ratiometric probes. This type of molecular sensor is designed in such a manner that the interaction of the analyte with the recognition unit of the probe alters the fluorophore–fluorophore distance. Based on this premise, one can distinguish between two types of probes: (A) intermolecular excimers formed by cation-assembled dimers, and (B) intramolecular excimers based



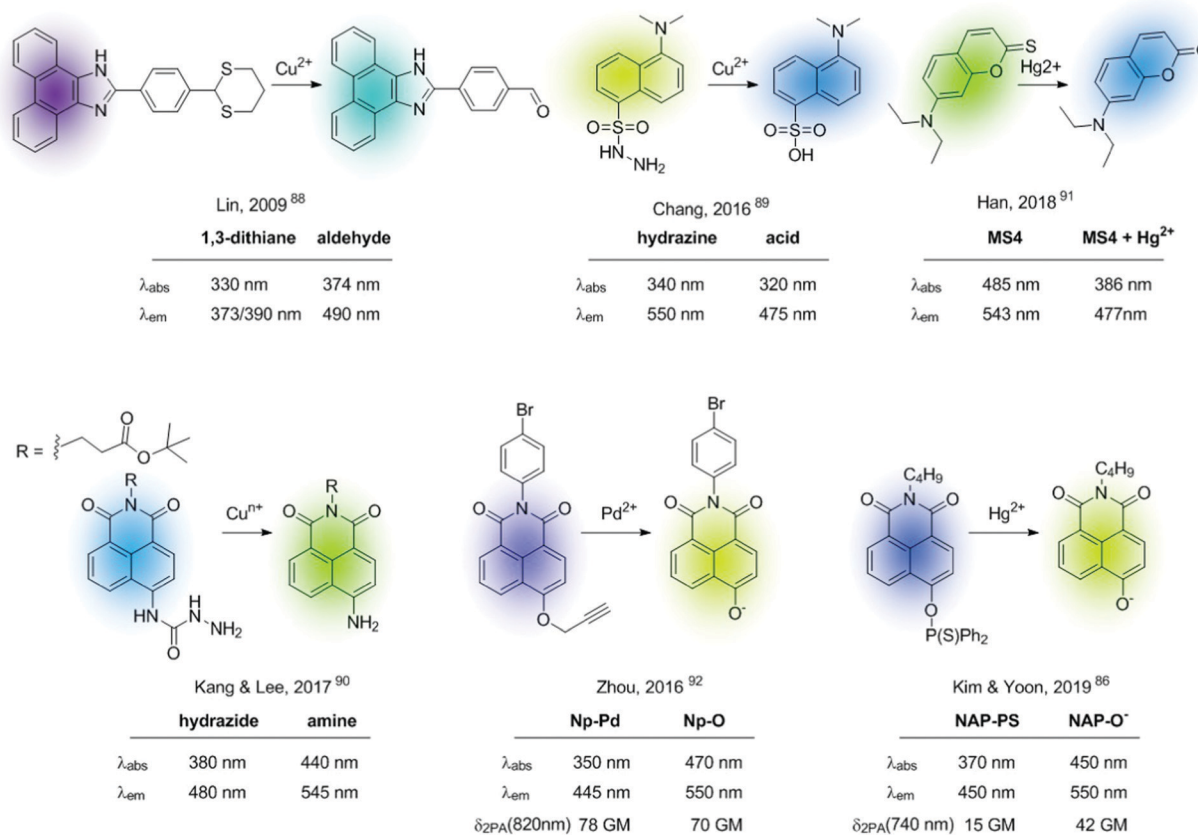


Fig. 12 Representative examples of fluorescent chemodosimeters for cations.

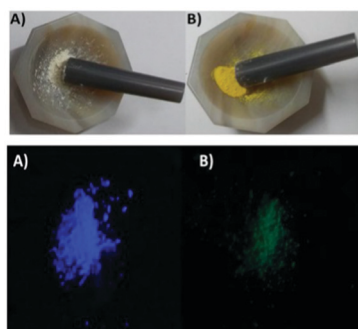


Fig. 13 NAP-PS chemodosimeter before (A) and after grinding with  $\text{Hg}(\text{ClO}_4)_2$  (B). Reproduced from ref. 86 with permission from The Royal Society of Chemistry, copyright 2019.

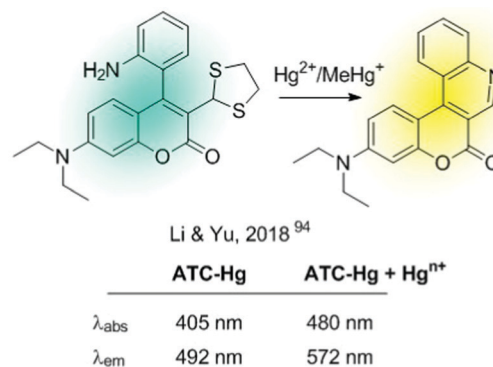


Fig. 14 Chemodosimeter based on covalent assembly.

on probes containing two fluorophores whose conformation is altered by cation binding.<sup>98</sup>

Some recent examples of sensors of type A are shown in Fig. 16.<sup>99–102</sup> Within this configuration, the binding of two sensors to the cation places the fluorophores close enough to form an intramolecular excimer. Among the chosen examples, it is worth highlighting the molecular sensor developed by Misra *et al.* in 2015,<sup>99</sup> which was modified and immobilized onto silica supports showing a ratiometric response to  $\text{Hg}^{2+}$  in the solid-state. Additionally, the interaction can be reversed by

treatment of the silica-supported sensor with EDTA allowing their use as  $\text{Hg}^{2+}$  extractors.

An early example of intramolecular excimer-based sensors of type B is the  $\text{Ba}^{2+}$  and  $\text{Sr}^{2+}$  probe developed by Ushakov and Barigelletti in 1999 (Fig. 17).<sup>103</sup> The probe consists of a bis(styryl) fluorophore containing two crown ether units. It relies on the formation of intramolecular sandwich complexes with the sensing ions (with radius larger than the cavity size of the crown ether), which facilitates excimer emission from the close-lying fluorophores (Fig. 17). Although in the example the fluorescence of the  $\text{Ba}^{2+}$ -complexed probe presents a blue-shifted



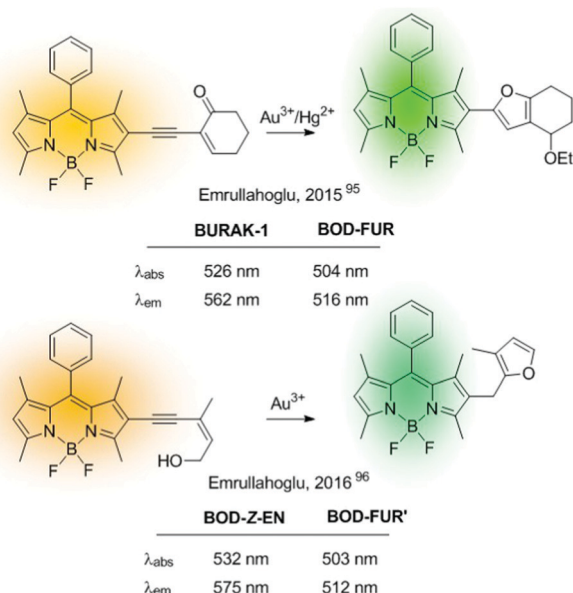


Fig. 15 Example of ratiometric probes based on metal-induced cascade reactions.

band compared to that of the unbound probe, a strong fluorescence attributed to excimer formation was also observed at *ca.* 600 nm.

In 2017 Lee *et al.* developed a ratiometric sensor for  $\text{Zn}^{2+}$  based on an intramolecular excimer inhibition.<sup>104</sup> It consists of a symmetric pyrene-peptidyl probe. In this example, the

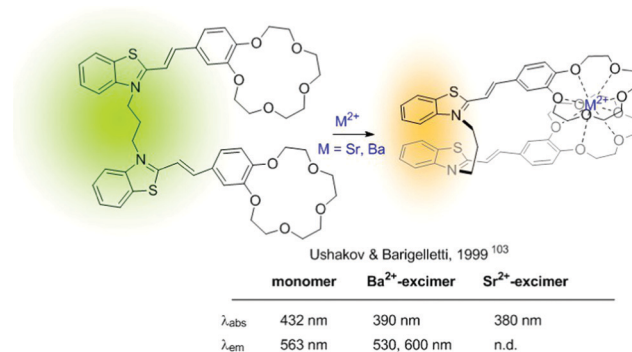


Fig. 17 Intramolecular excimer-based probe for  $\text{Ba}^{2+}$ .

interaction of the probe with the cation modifies its conformation inhibiting pyrene-pyrene excimer formation (Fig. 18).

### 3.3. FRET-based sensors

Since the German scientist T. Förster described the FRET effect in 1948,<sup>105</sup> many fluorescent sensors based on this effect have been developed,<sup>42,106</sup> which have been widely applied in diverse fields, such as biology, physiology, medicine, and pharmacology. This type of sensor presents ideal ratiometric sensing properties as it benefits from a simple structural design. This section describes some FRET-based fluorescent sensors and the combination of these with other mechanisms. As described in Fig. 5, these sensors are based on an energy donor/acceptor system. In many of them, rhodamine appears as a fundamental part of the fluorescent sensor

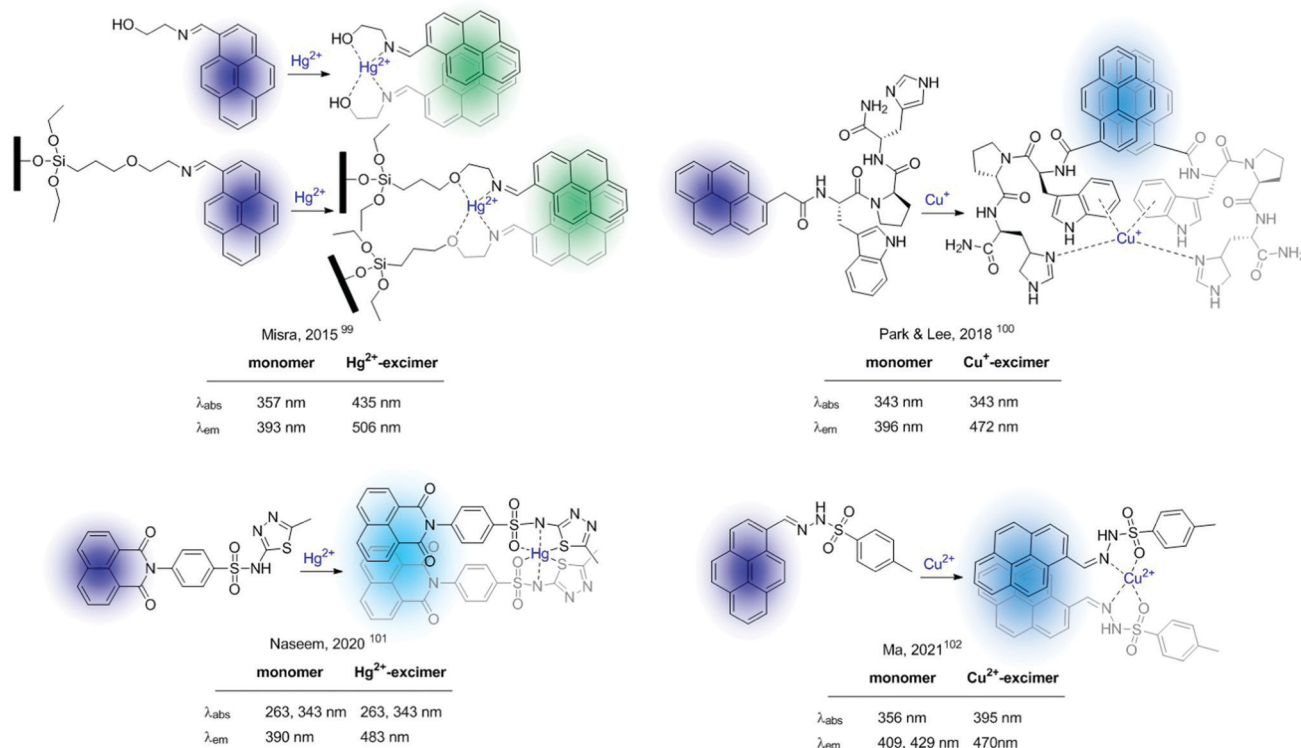


Fig. 16 Excimer-based probes for cations based on 2:1 sensor/cation supramolecular complexes.



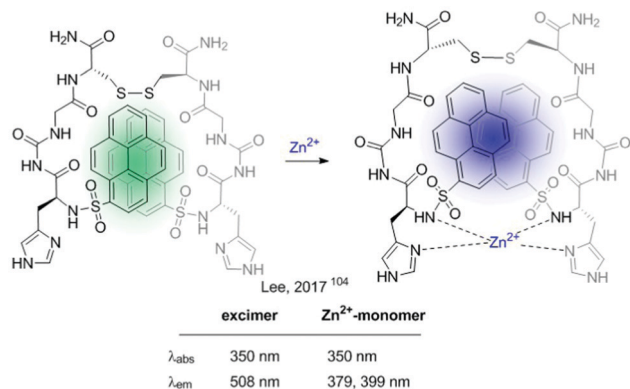


Fig. 18 Intramolecular excimer inhibition-based probe for Zn<sup>2+</sup>.

(acceptor), and it is integrated in the sensor in the form of the non-fluorescent rhodamine spirolactam. The analyte's reaction promotes spiro opening, revealing rhodamine fluorescent dye, which results in a FRET-induced rhodamine-centered emission.

In this context, Meng described in 2018 a two-photon ratiometric fluorescent probe (**RN3**) for Pd<sup>2+</sup> based on coumarin (donor) and rhodamine B (acceptor) connected by a diethylene-triamine linker (Fig. 19).<sup>107</sup> The emission of a 7-substituted coumarin was determined in the range of 370–650 nm, which fully covered the rhodamine B's absorption (450–590 nm). **RN3** showed high sensitivity and selectivity towards Pd<sup>2+</sup> in aqueous solutions, with a clear ratiometric fluorescent response in a

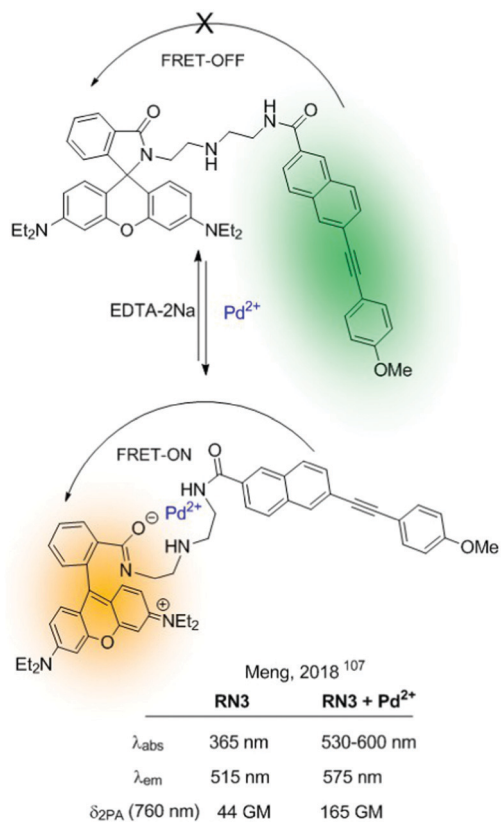


Fig. 19 **RN3** based on the FRET-effect to detect Pd<sup>2+</sup> in aqueous media.

one- and two-photon excitation regime. Additionally, interactions with the cation boosted the TPA cross-section, which was attributed to ICT enhancement on the coumarin upon coordination to Pd<sup>2+</sup>. **RN3** was tested for the detection of Pd<sup>2+</sup> in deep tissues and living cells under TPA, showing low cytotoxicity.

Different rhodamine-based FRET-dyads have been used for the construction of ratiometric sensors (*i.e.* using BODIPY or quinoline-benzothiazole as a donor). They have been widely used for sensing highly polluting cations such as Hg<sup>2+</sup><sup>108</sup> or Cd<sup>2+</sup>. For the latter, in 2015 Goswami described **RQBT**, a rhodamine/quinoline-benzothiazole-based sensor (Fig. 20).<sup>109</sup> The interaction of Cd<sup>2+</sup> with the probe results in the cleavage of the spiro ring and formation of the Cd<sup>2+</sup>-chelated structure, as confirmed by single-crystal X-ray analysis. Probe **RQBT** also showed ratiometric fluorescence in dry media (silica-gel TLC plates), which was used for the construction of portable Cd<sup>2+</sup> sensing kits (Fig. 21).

Alternatively, FRET-probes for cations can be constructed by introducing an ICT sensor as part of a FRET dyad. Interaction of the cation with the ICT sensor will produce a shift in its absorption and/or emission properties, modifying the efficiency

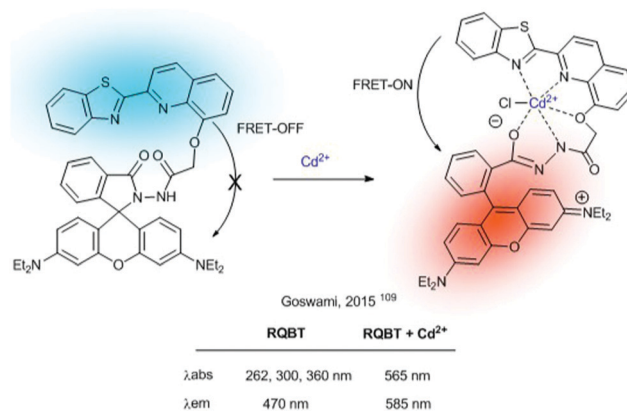


Fig. 20 Proposed FRET-based Cd<sup>2+</sup>-sensing mechanism of **RQBT**.

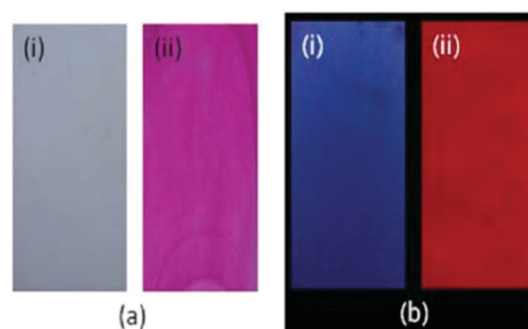


Fig. 21 Photographs of TLC plates after immersion in an **RQBT**-MeOH solution (i) and after immersion in an **RQBT**-Cd<sup>2+</sup> methanol solution (ii) taken in ambient light (a) and under hand-held UV light (b). Excitation wavelength of the UV light is 365 nm. Adapted from ref. 109 with permission from American Chemical Society, copyright 2015.



of the FRET process. A recent example of such a strategy is the CPBT probe for  $Zn^{2+}$  developed by Chen, He, Guo and colleagues.<sup>110</sup> CPBT integrates a TPA donor coumarin derivative with an ICT fluorophore 4-amine-7-sulfamoylbenzo[*c*][1,2,5]-oxadiazole linked to the ionophore (Fig. 22). Within this type of construction, the interaction of the cation with the ICT fluorophore will cause a blue-shift in its emission, diminishing the spectral overlap between the donor and acceptor and weakening the FRET process. CPBT was tested for 2D and 3D ratiometric mapping of  $Zn^{2+}$  in living cells and in the head of zebrafish. Additionally, it showed TPA cross-section >40 GM, which makes it a suitable candidate to be used in two-photon excited fluorescence imaging.

CPBT can be described as an FRET-on  $\rightarrow$  FRET-off ratiometric probe. The reverse mechanism (FRET-off  $\rightarrow$  FRET-on) can also be envisioned, and the QA sensor, developed by Mao *et al.*, serves as a representative example (Fig. 23).<sup>111</sup> This probe contains an anthracene unit as a donor and a quinoline ICT sensor as an acceptor. In the absence of the analyte ( $Zn^{2+}$ ), the probe shows the characteristic emission of anthracene. Interactions of  $Zn^{2+}$  with the quinolone sensor produce a red shift in its absorption band, together with a concomitant enhancement of the effective overlap with the anthracene emission, thus triggering the FRET process.

Closely related to FRET sensors, one can encounter the TBET-based probes (through bond energy transfer).<sup>112</sup> They comprise sensors based on donor-acceptor dyads in which the donor and acceptor are connected through a conjugated bond. These probes can be used for the construction of sensors

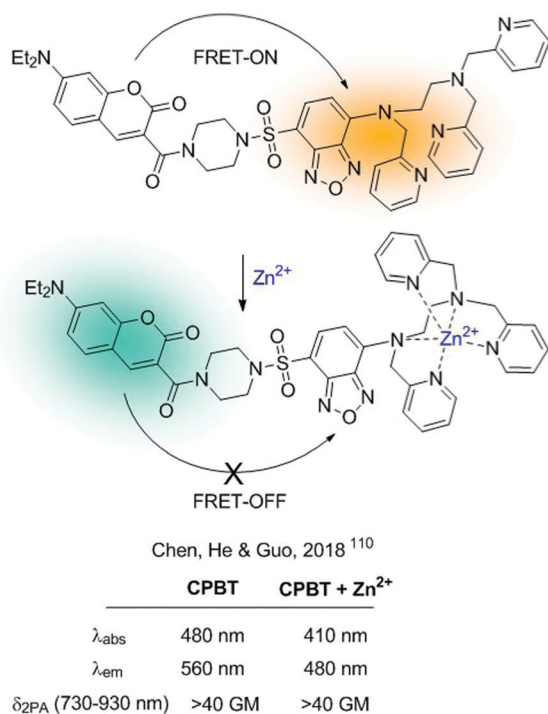


Fig. 22 FRET-ICT probe for  $Zn^{2+}$ , showing an FRET-ON/FRET-OFF mechanism.

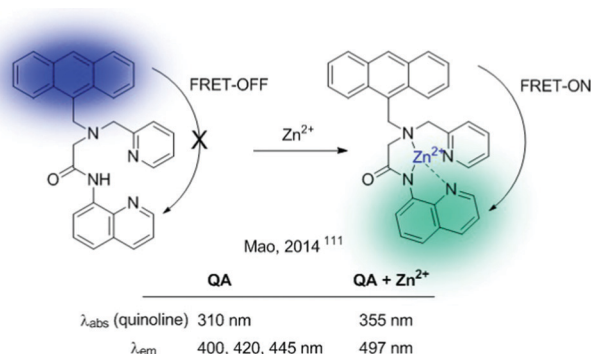


Fig. 23 FRET-ICT probe for  $Zn^{2+}$ , showing an FRET-OFF/FRET-ON mechanism.

as long as the cation interaction alters the extent of the FRET process. The simplest strategy consists of revealing a “dormant” fluorophore (rhodamine), as described before for FRET probes. A representative example is **Np-Rh**, a TPA ratiometric TBET-based probe for  $Cu^{2+}$  shown in Fig. 24.<sup>113</sup>

### 3.4. Two-component fluorescent bicolour indicators

We applied the concept outlined in Section 2. 4 to the issue associated with the background-free detection of individual barium cations in  $\beta\beta 0\nu$  experiments (*vide supra*). Application of indicators to this kind of problem requires a series of developments to enable its use. Although several molecular fluorescent sensors for  $Ba^{2+}$  capture and detection have been described in solution,<sup>114-120</sup> the features of the  $Xe \rightarrow Ba^{2+} + 2e$  experiment at a high pressure require the development of fluorescent sensors able to operate efficiently in the interface between a solid substrate and ultra-dry noble gas.

The Xenon Chamber must incorporate “dry” molecular sensors capable of capturing the  $Ba^{2+}$  ion in a pressurised xenon environment. In addition, a very stable binding that prevents release of the barium cation before optical interrogation is required. As we have commented above, one possibility consists of developing monocolour ICT turn-on sensors.<sup>23,24</sup> Our approach<sup>121</sup> was based on the above-mentioned two-component design. Benzo[*a*]imidazo[5,1,2-*cd*]indolizine was chosen as a suitable fluorophore of type C3 (Fig. 1). To this HetAr<sub>1</sub> group a *para*-phenylene group was added as an Ar<sub>2</sub> component. Finally, a pentaoxa-16-azacyclooctadecane aza-crown ether unit was included as a **B3** type coronand ionophore. This compound was named **FBI**, Fluorescent Bicolour Indicator (Fig. 25A). DFT calculations showed that the HetAr<sub>1</sub>-Ar<sub>2</sub> group gives rise to the fluorophore 1 ensemble, in which the dihedral angle formed by both components was close to coplanarity.

Capture of  $Ba^{2+}$  by adding barium perchlorate in acetonitrile solution or in the gas phase led to a complex in which the  $Ba^{2+} \cdots Ph$  interaction disconnects both components of fluorophore 1 (Fig. 25B). This coordination pattern in which the barium cation, aside the cation- $\pi$  interaction, complexes the five oxygen atoms of the aza-crown ether and one nitrogen atom of fluorophore 2 was elucidated by DFT calculations and NMR experiments.<sup>121</sup> As expected, the **FBI**- $Ba^{2+}$  complex shows a



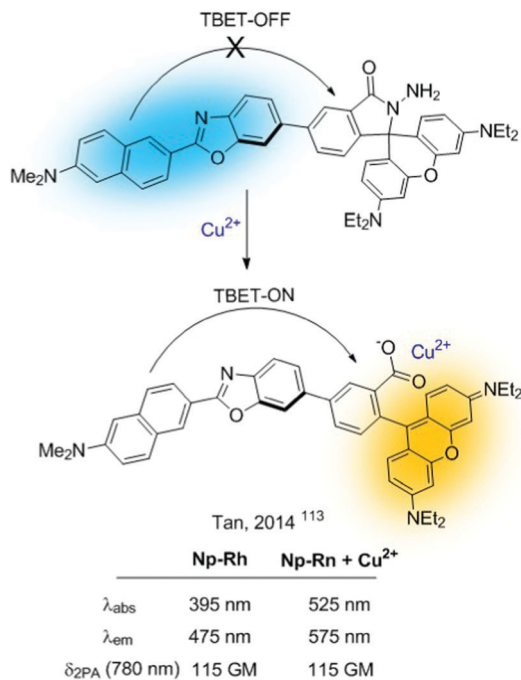


Fig. 24 TBET-based probe for sensing Cu<sup>2+</sup>.

noticeable blue shift (Fig. 25C), with a peak discrimination factor  $f_{\lambda} = 179.74$  and a brightness value of  $B_{\lambda} = 8.06 \text{ M}^{-1} \text{ cm}^{-1}$ . The fluorescence spectra in solution and in the gas phase show intense emission signals for **FBI** and **FBI·Ba<sup>2+</sup>** (Fig. 25D) compatible with the FMO scheme shown in Fig. 25. Application of an optical filter for  $\lambda > 450 \text{ nm}$  to both compounds shows a no signal pattern in the case of **FBI**. This separation of the chelated/unchelated photophysical properties results fundamental for  $\beta\beta 0\nu$  experiments. In order to detect the Ba<sup>2+</sup> cations produced by the decay, they will be driven by electric fields to the cathode where a region of a few centimetres will need to be scanned to look for a positive signal of the complexed molecule. This region will be fully covered, with a density of  $10^5\text{--}10^6 \text{ molecules } \mu\text{m}^{-2}$ , by the bicolour indicator to guarantee a high efficiency to capture the cation. The discrimination factor between the response of the complexed indicator and the response of the residual background molecules on the surface must be large to allow separation of the single chelated molecule from the background emission produced by the surrounding molecules that will also be illuminated during the scan.

## 4. Conclusions and outlook

In this paper, we have highlighted significant advances in the design and experimental validation of fluorescent bicolour sensors. Most of these photo-indicators are based on binding ionophores and fluorophores connected by covalent bonds. If required, suitable spacer units and linkers are incorporated. In addition, some fluorescent bicolour sensors merge the ionophore and the fluorophore components, thus yielding chelating fluorophores whose photophysical properties change

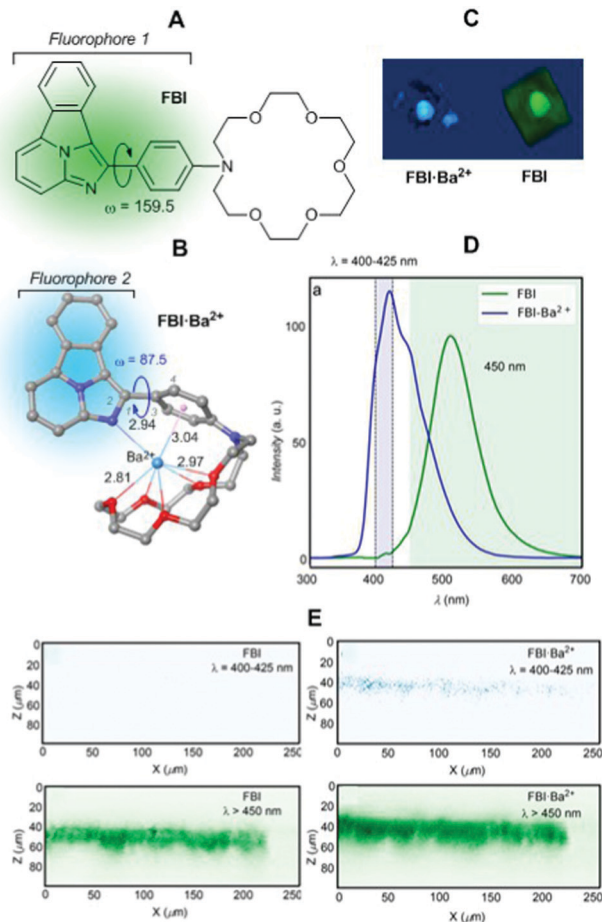


Fig. 25 (A) Structure of the **FBI** molecule showing the approximate coplanarity of the two components of fluorophore **1**. (B) DFT optimized structure of complex **FBI·Ba<sup>2+</sup>**. Bond angles and distances are given in deg and Å, respectively. (C) 3D tomography of **FBI** and **FBI·Ba<sup>2+</sup>** obtained by means of a two-photon absorption (TPA) microscopy device. (D) Solid-gas phase emission spectra of **FBI** and **FBI·Ba<sup>2+</sup>** adsorbed on silica. (E) Z–X profiles showing emission profiles (after TPA excitation) from both complexes with and without a  $\lambda > 450 \text{ nm}$  filter. Adapted from ref. 121 with permission from Springer Nature, copyright 2020.

significantly upon cation binding. In other cases, the ionophore (most frequently a coronand) and the fluorophore work together upon cation coordination and induce a bathochromic or hypsochromic shift in their emission spectra. Finally, several fluorescent sensors generate their bicolour response by means of a chemical reaction promoted by the metallic cation that induces a change in the chemical structure of the fluorophore.

These structural and chemical features result in a bicolour response as an attractive alternative to on–off or off–on changes *via* photoinduced electron transfer (PET) phenomena upon cation coordination. Instead, the bicolour response requires changes in the photophysical properties of the sensor, which is fluorescent in both the free and coordinated states within an on–on' conceptual scheme. These dual patterns include intramolecular charge transfer processes (ICT), excimer/exciple complex and Förster resonance energy transfer (FRET). As an additional design feature, the



fluorophore can incorporate different rotatable units that can be decoupled in the presence of a suitable metal cation thus resulting in different fluorophores in the free and coordinated states.

It is expected that in the years to come these designs will permit the development of more sensitive and selective fluorescent bicolour sensors able to detect specifically metal cations in very low concentrations. This situation will permit further advances in fields ranging from cell biology to particle physics. In addition, the development of fluorescent sensors able to detect metallic cations in the gas phase and at solid-gas interfaces would permit the exploration of supramolecular chemistry in the absence of solvents, thus expanding the field from “wet” to “dry” chemistry.

## Conflicts of interest

There are no conflicts to declare.

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