



taking advantage of the AIEE feature. In addition, the applications in intracellular Zn<sup>2+</sup> imaging and two-photon fluorescence cell imaging were investigated.

## 2. Experimental

### 2.1 Materials

The synthetic routes to **SPF-1** and **SPF-2** are shown in Scheme 1. Synthetic intermediates **1** and **2** were prepared according to our previously reported methods.<sup>21</sup> All chemicals were of analytical grade and were used as received.

### 2.2 Measurements

The melting points of the compounds were determined using WRX-4 microscope melting point apparatus. All nuclear magnetic resonance (NMR) spectra were obtained on a Bruker DRX 400 MHz spectrometer. Elemental analyses were performed using Atlantic Microlab. UV-Vis absorption spectra were determined using a Perkin Elmer Lambda 35 spectrophotometer. Fluorescence spectra were recorded on a Perkin Elmer LS55 instrument.

The TPA cross-section of **SPF-2** was measured at 800 nm using the open-aperture Z-scan technique using a high-power femtosecond laser (PHAROS High-Power Femtosecond Lasers sp). The pulse duration was about 190 fs (estimated as the full width half maximum (FWHM) of a Gaussian temporal profile). The light beam used was Top-hat and the focal length ( $f$ ) was 250 mm. The TPA cross-section ( $\sigma$ ) value can be calculated by using the equation:  $\sigma = hv\beta N_0$ , in which  $N_0 = N_A C$  is the number density of the centers ( $N_A$  is the Avogadro constant,  $C$  represents the solute molar concentration), and  $\beta$  is the TPA coefficient.<sup>22</sup>

### 2.3 Living cell cultures and imaging

A549 cells were incubated in DMEM medium with 10% fetal bovine serum at 37 °C under a 5% CO<sub>2</sub> atmosphere. One day before the fluorescence imaging, the A549 cells were passaged and plated in 30 mm glass culture dishes. During the logarithmic growth phase, they were cultured to 50–70% culture fit in the presence of complete medium (2 mL). Afterwards, the cells on glass slides were washed with DMEM and fed with culture medium (contain 1% DMSO) containing the probe (50 μM) for 30 min at 37 °C under a 5% CO<sub>2</sub> atmosphere. Subsequently, A549 cells were washed with PBS solution three times. One of the groups cells stained with **SPF-1** was treated with 100 μM Zn<sup>2+</sup> for another 30 min and washed three times with PBS solution.

Two-photon fluorescence cell images were taken on an Ultime IV confocal microscope equipped with a femtosecond Ti:sapphire laser.

### 2.4 Preparation of the compounds

**Synthesis of 2,2'-(1Z,1'Z)-((2'-(tert-butyl)-9,9'-spirobi[fluorene]-2,7'-diyl)bis(azanylylidene))bis(methanylylidene)diphenol (SPF-1).** To a 50 mL flask, 0.402 g (1 mmol) **2**, 10 mL absolute ethyl alcohol, 5 mL ethyl acetate, and 0.22 mL (2.1 mmol) 2-hydroxybenzaldehyde

were added. The mixture was stirred at room temperature for 1 h. The resulting precipitate was filtered and washed with cold ethanol and finally dried under vacuum. Yield 0.51 g (83.6%). MP: 243–245 °C. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) 13.21 (s, 2H), 7.94 (d,  $J = 8.1$  Hz, 1H), 7.89 (t,  $J = 8.1$  Hz, 2H), 7.82 (d,  $J = 8.1$  Hz, 1H), 7.50 (dd,  $J = 8.1, 1.7$  Hz, 1H), 7.44 (t,  $J = 7.8$  Hz, 1H), 7.39 (dd,  $J = 8.1, 1.8$  Hz, 1H), 7.32 (q,  $J = 7.3, 5.6$  Hz, 5H), 7.17 (t,  $J = 7.8$  Hz, 1H), 6.97 (dd,  $J = 8.0, 2.5$  Hz, 2H), 6.88 (td,  $J = 7.4, 2.7$  Hz, 2H), 6.81 (d,  $J = 7.8$  Hz, 2H), 6.81 (d,  $J = 7.8$  Hz, 2H), 6.74 (s, 1H), 6.66 (s, 1H), 1.21 (s, 9H). <sup>13</sup>C NMR (101 MHz, CDCl<sub>3</sub>)  $\delta$  161.77, 161.62, 161.17, 151.79, 150.43, 150.40, 149.13, 148.54, 148.02, 147.65, 141.09, 140.81, 140.76, 138.67, 133.03, 132.97, 132.21, 132.16, 128.09, 128.03, 125.51, 124.25, 121.95, 121.82, 120.91, 120.87, 120.61, 120.15, 119.64, 119.20, 118.99, 118.96, 117.21, 116.49, 116.32, 77.07, 76.76, 66.15, 35.01, 31.47. Elemental analysis (%) calcd for C<sub>43</sub>H<sub>34</sub>N<sub>2</sub>O<sub>2</sub>: C 84.56, H 5.61, N 4.59, O 5.24. Found: C 84.81, H 5.62, N 4.57.

**Synthesis of 2'-(tert-butyl)-7,7'-dinitro-9,9'-spirobi[fluorene]-2-carbaldehyde (3).** To a 250 mL flask, 3.2 g (6.96 mmol) compound **1** and 30 mL anhydrous dichloromethane were added under a nitrogen atmosphere. After the mixture was stirred at 0 °C for 10 min, a 30 mL CH<sub>2</sub>Cl<sub>2</sub> solution of TiCl<sub>4</sub> (16 mL) and 20 mL CH<sub>2</sub>Cl<sub>2</sub> solution of dichloro(methoxy)methane (4 mL, 4.2 mmol) were added dropwise over 30 min, respectively. The reaction mixture was continually stirred at room temperature for 24 h. Then, the mixture was poured into iced water and extracted using CH<sub>2</sub>Cl<sub>2</sub> (20 mL × 3). The combined organic solution was washed with 50 mL saturated sodium bicarbonate solution and 50 mL water, respectively. The organic solvent was removed using a rotary evaporator. The residue was purified by column chromatography on silica (petroleum ether/ethyl acetate, 5:1, v/v) to give a yellow solid (1.56 g). Yield: 46.0%. MP: 175–178 °C. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>,  $\delta$ ): 9.91 (s, 1H), 8.40 (dd,  $J = 8.4, 2.0$  Hz, 1H), 8.34 (dd,  $J = 8.4, 2.0$  Hz, 1H), 8.14 (dd,  $J = 15.0, 8.2$  Hz, 2H), 8.04 (dd,  $J = 8.0, 1.1$  Hz, 1H), 7.98 (d,  $J = 8.5$  Hz, 1H), 7.92 (d,  $J = 8.2$  Hz, 1H), 7.59 (m,  $J = 9.0, 8.2$  Hz, 2H), 7.47 (d,  $J = 2.0$  Hz, 1H), 7.31 (s, 1H), 6.73 (s, 1H), 1.18 (s, 9H).

**Synthesis of (E)-4-(2-(2'-(tert-butyl)-2,7'-dinitro-9,9'-spirobi[fluorene]-7-yl)vinyl)-N,N-diphenylaniline (4).** To a 50 mL flask, 0.123 g (0.25 mmol) **3**, 0.324 g (0.5 mmol) 4-(iodotriphenylphosphoranyl)methyl)-N,N-diphenylaniline, 0.03 g (0.75 mmol) NaH and 5 mL dried THF were added under a nitrogen atmosphere. Then, the mixture was stirred at 0 °C for 24 h. The solvent was evaporated, and the residue was subjected to column chromatography on silica gel eluting with petroleum ether and ethyl acetate (4:1, V/V) to give 98 mg of a red solid (yield 54%). MP: 155–158 °C. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  8.36 (ddd,  $J = 7.9, 5.6, 2.1$  Hz, 2H), 8.03–7.92 (m, 4H), 7.65 (dd,  $J = 8.1, 1.5$  Hz, 1H), 7.60 (dd,  $J = 8.2, 1.7$  Hz, 1H), 7.56 (d,  $J = 2.0$  Hz, 2H), 7.30 (d,  $J = 3.3$  Hz, 1H), 7.27 (s, 2H), 7.25 (s, 1H), 7.13–6.97 (m, 10H), 6.95 (s, 1H), 6.90–6.79 (m, 3H), 1.22 (s, 9H).

**Synthesis of (E)-7-(tert-butyl)-7'-(4-(diphenylamino)styryl)-9,9'-spirobi[fluorene]-2,2'-diamine (5).** To a mixture of 0.366 g 4 (0.5 mmol) and 0.558 g iron powder (1 mmol) in 25 mL ethanol, 2 mL of concentrated hydrochloric acid was added dropwise over 30 min under a nitrogen atmosphere. After refluxing for 3 h,

the solution was neutralized dropwise with  $\text{NH}_4\text{OH}$  solution, and the resulting precipitate was filtered off with suction. The solvent was evaporated, and the residue was purified by column chromatography (petroleum ether/ethyl acetate, 10:7, v/v) to afford 0.252 g of a yellow solid **5** (75%). MP: 181–183 °C.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ ) 7.68 (d,  $J = 7.9$  Hz, 1H), 7.63 (d,  $J = 8.0$  Hz, 1H), 7.60–7.49 (m, 3H), 7.40 (d,  $J = 8.4$  Hz, 2H), 7.40 (d,  $J = 8.4$  Hz, 2H), 7.35–7.23 (m, 6H), 7.06–6.95 (m, 8H), 7.06–6.95 (m, 8H), 6.93 (s, 1H), 6.85 (d,  $J = 8.5$  Hz, 2H), 6.69 (s, 1H), 6.57–6.51 (m, 3H), 5.85 (dd,  $J = 6.3, 2.0$  Hz, 2H), 5.14 (d,  $J = 35.9$  Hz, 4H), 1.10 (s, 9H).

**Synthesis of 2,2'-((1Z,1'Z)-((2-(*tert*-butyl)-2'-((*E*)-4-(diphenylamino)styryl)-9,9'-spirobifluorene]-7,7'-diyl)bis(azanylylidene))-bis(methanylylidene)diphenol (SPF-2).** To a 50 mL flask, 0.134 g (0.2 mmol) **5**, 15 mL absolute ethyl alcohol, 5 mL ethyl acetate, and 43.8  $\mu\text{L}$  (0.42 mmol) 2-hydroxybenzaldehyde were added. The mixture was refluxed at 85 °C. The reaction progress was monitored on TLC. When the reaction was over (about 1 hour), the solvent was evaporated. A yellow solid was precipitated, which was filtered and washed with cold ethanol and finally dried under vacuum. Yield 0.138 g (79%). MP: 173–175 °C.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  13.2 (s, 2H), 8.52 (d,  $J = 5.9$  Hz, 2H), 7.91 (dd,  $J = 8.1, 3.1$  Hz, 2H), 7.86 (t,  $J = 8.0$  Hz, 2H), 7.58 (dd,  $J = 8.1, 1.6$  Hz, 1H), 7.52 (dd,  $J = 8.1, 1.8$  Hz, 1H), 7.41–7.29 (m, 7H), 7.26–7.22 (m, 4H), 7.12–7.06 (m, 4H), 7.06–6.94 (m, 7H), 6.94–6.82 (m, 6H), 6.71 (dd,  $J = 7.3, 1.9$  Hz, 2H), 1.22 (s, 9H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  161.70, 161.63, 161.14, 151.87, 150.72, 150.44, 149.65, 148.54, 147.89, 147.75, 147.51, 147.34, 140.71, 140.56, 140.34, 138.69, 137.76, 133.02, 132.97, 132.18, 131.31, 129.28, 128.16, 127.28, 126.77, 126.52, 125.60, 124.56, 123.37, 123.07, 122.00, 121.97, 121.91, 120.94, 120.80, 120.62, 120.38, 119.65, 119.17, 118.97, 118.92, 117.19, 117.17, 116.46, 116.34, 77.38, 77.26, 77.06, 76.75, 66.08, 31.47. Elemental analysis (%) calcd for  $\text{C}_{60}\text{H}_{49}\text{N}_3\text{O}_2$ : C 85.98, H 5.61, N 4.77, O 3.64. Found: C 85.72, H 5.62, N 4.78.

### 3. Results and discussion

#### 3.1 One-photon absorption

The one-photon absorption spectra of **SPF-1** and **SPF-2** in various organic solvents were investigated. The maximum absorption wavelength of **SPF-2** is longer than that of **SPF-1** (Table 1), which can be attributed to the incorporation of a diphenylaminoaryl-vinyl group into the spirobifluorene core of **SPF-2**. It can be seen from Table 1 that the absorption spectra of **SPF-1** and **SPF-2** exhibit almost no change with different polar solvents. However, the water content has a significant effect on the absorption response of the two compounds. It was noted that the absorption spectra of **SPF-1** in the DMF/water and **SPF-2** in 1,4-dioxane/water mixtures were all red-shifted and broadened.

Table 1 One-photon absorption properties of **SPF-1** and **SPF-2**

Compound	<b>SPF-1</b>				<b>SPF-2</b>			
	Toluene	THF	DCM	DMF	Toluene	THF	DCM	DMF
$\lambda_{\text{max}}$ (nm)	362	360	363	361	397	396	398	400

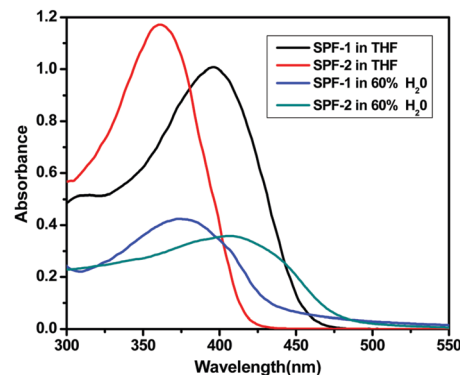


Fig. 1 One-photon absorption spectra of **SPF-1** and **SPF-2** in THF and organic solvent/water mixtures (60% water in volume).

In addition, the absorbance of **SPF-1** and **SPF-2** decreases with the increase of the water content (Fig. 1), respectively. The phenomena may be caused by the formation of their nanoaggregates.<sup>23,24</sup>

#### 3.2 AIEE properties

**SPF-1** and **SPF-2** are soluble in common organic solvents (such as toluene, THF, dichloromethane, 1,4-dioxane and DMF), but insoluble in pure water. We used anhydrous DMF as a good solvent and water as a poor solvent to investigate the AIEE attributes of **SPF-1** by the precipitation method.<sup>23,25</sup> **SPF-1** exhibited a relatively weak fluorescence emission (excited at 362 nm) in DMF solution (10  $\mu\text{M}$ ). As shown in Fig. 2, the fluorescence emission of **SPF-1** began to increase when the water was added slowly. When the water content reached 50%, a significant fluorescent enhancement (at 538 nm) could be observed. At a water content of more than 60%, the fluorescence emission intensity reached the maximum (Fig. 2, right). Increasing water fractions can reduce the solubility of **SPF-1** and the formation of **SPF-1** aggregates in the DMF/water mixture was confirmed using dynamic light scattering (DLS) analysis (Fig. S1, ESI†). In the aggregated state, the intramolecular rotations are restricted, which blocks the nonradiative relaxation channel and activates the radiative decay.<sup>7</sup> Apparently, the emission enhancement was caused by aggregation-induced effects. However, a further increase in the water content (>80%) caused the AIEE properties to undergo a slight decrease in the PL intensity. This phenomenon could be caused by the variation in the packing mode of the molecules in the aggregates.<sup>23,26</sup>

Similar AIEE properties were obtained for **SPF-2**. When excited at 400 nm, **SPF-2** emits weak fluorescence in pure 1,4-dioxane solution (10  $\mu\text{M}$ ). Addition of water into the 1,4-dioxane solution, however, aggregates the **SPF-2** molecules and enhances the fluorescence emission intensity (at 540 nm) (Fig. 3, left). **SPF-2** gave a maximum luminescence at  $f_w = 60\%$  (Fig. 3, right). Compared to **SPF-1**, a more significant fluorescence enhancement was observed for **SPF-2** when the water content was increased from 40% to 50% in volume. This result can presumably be attributed to the comparatively lower solubility of **SPF-2** in the aqueous mixture. Thus, it can be confirmed that **SPF-1** and **SPF-2** were endowed with AIEE attributes.<sup>27</sup>

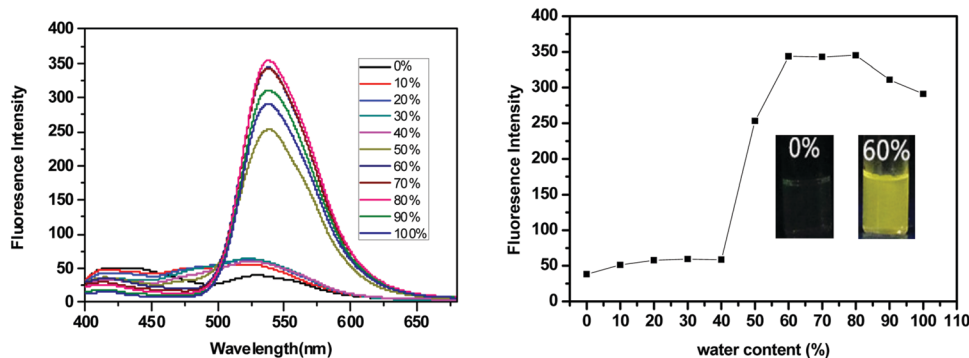


Fig. 2 Emission spectra of **SPF-1** in DMF and DMF/water mixtures with different water fractions (left). Plot of the emission intensity ( $I$ ) versus the composition of the aqueous mixtures of the **SPF-1** DMF solution (right). Solution concentration:  $10 \mu\text{M}$ ;  $\lambda_{\text{ex}}$ :  $362 \text{ nm}$ . Inset (right): photographs of **SPF-1** in pure DMF and DMF/water mixtures (60%) under  $365 \text{ nm}$  UV irradiation.

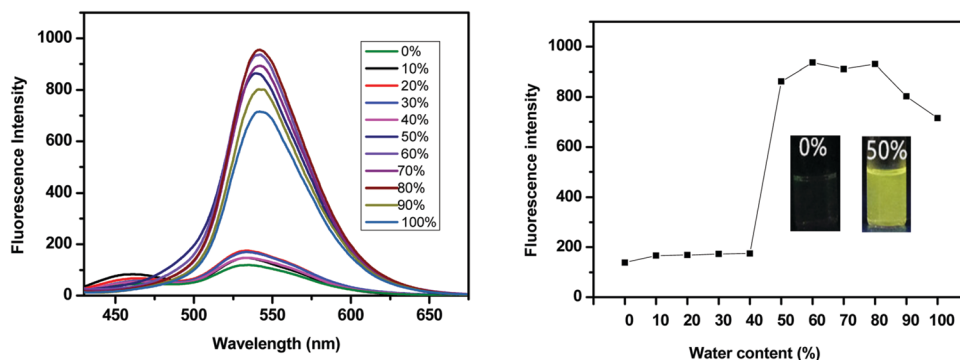
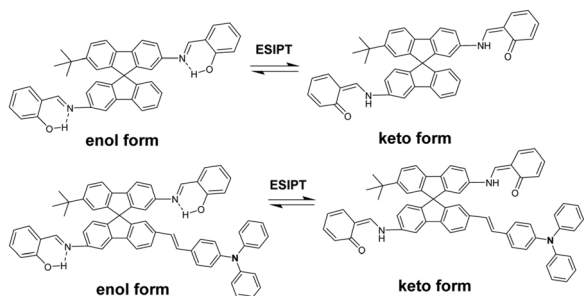


Fig. 3 Emission spectra of **SPF-2** in 1,4-dioxane and 1,4-dioxane/water mixtures with different water fractions (left). Plot of the emission intensity ( $I$ ) versus the composition of the aqueous mixtures of the **SPF-2** DMF solution (right). Solution concentration:  $10 \mu\text{M}$ ;  $\lambda_{\text{ex}}$ :  $400 \text{ nm}$ . Inset (right): photographs of **SPF-2** in pure 1,4-dioxane and 1,4-dioxane/water mixtures (50%) under  $365 \text{ nm}$  UV irradiation.

### 3.3 Excited-state intramolecular proton transfer characteristics of SPF

The process of excited-state intramolecular proton transfer (ESIPT) consists of proton transfer between an enol ( $\text{E}^*$ ) and a keto form ( $\text{K}^*$ ) upon photoexcitation (Scheme 2).<sup>28–31</sup> In general, an ESIPT-based luminophore shows dual emission bands: the normal emission from the enol-form (high energy, low Stokes shift) and the ESIPT emission from the keto-form (low energy, large Stokes shift).<sup>30,32,33</sup> AIE mechanisms involving ESIPT have been reported in the literature.<sup>30,32</sup> In the aggregate state, the keto emission used to be the preferred emission owing to



Scheme 2 Illustration of the ESIPT process for **SPF-1** and **SPF-2**.

strong intramolecular hydrogen bonding and other favorable parameters.<sup>32,34,35</sup> In the DMF/ $\text{H}_2\text{O}$  mixture (Fig. 2, left), **SPF-1** showed dual weak emissions at a lower fraction of water (0–40%), in which the shorter-wavelength emission occurred at  $425 \text{ nm}$  and the longer-wavelength emission occurred at  $528 \text{ nm}$  (excitation at  $362 \text{ nm}$ ). As the ratio of water increased from 50% to 100%, the emission at  $528 \text{ nm}$  becomes the more prominent emission and is associated with the red-shift to  $538 \text{ nm}$ . The red shift of the emission band could be attributed to the result of aggregation-induced charge-transfer enhancement.<sup>36–38</sup> A similar trend was observed for compound **SPF-2** (Fig. 3, left). It showed weak fluorescence at  $460 \text{ nm}$  (enol emission) and  $530 \text{ nm}$  (keto emission) in a 1,4-dioxane/water mixture with water fractions ranging from 0 to 40 vol%. Upon increasing the amount of water to more than 40%, the fluorescence intensity of the shorter-wavelength emission ( $460 \text{ nm}$ ) disappeared completely, while that of the longer-wavelength emission ( $530 \text{ nm}$ ) increased significantly with a red shift from  $530$  to  $542 \text{ nm}$ .

To gain a better understanding of the ESIPT phenomena of **SPF-1** and **SPF-2**, a theoretical calculation based on density functional theory (DFT) and time-dependent density functional theory (TDDFT) methods was carried out using the Gaussian 16 program. The geometry optimizations of the  $\text{S}_0$  and  $\text{S}_1$  states were performed with the B3LYP hybrid DFT and the basis set

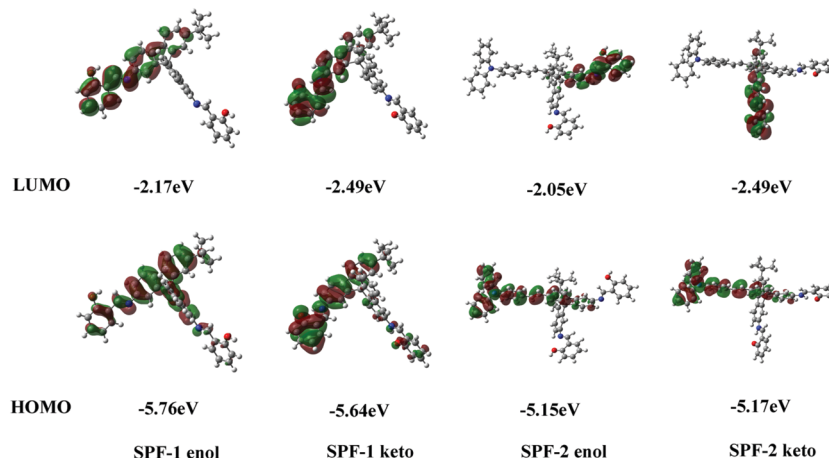


Fig. 4 The calculated frontier molecular orbitals for the HOMO and LUMO and the orbital energies of **SPF-1** and **SPF-2** in DMF.

used was 6-31G\*. DMF was employed as solvent. Fig. 4 displays the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) diagrams for the two compounds, and the calculated HOMO and LUMO energies are correspondingly listed. It is noteworthy that for **SPF-1** and **SPF-2**, the K\* form is more stable than the E\* form by more than 0.32 eV.<sup>39</sup> Therefore, the E\* → K\* transformation could occur through a barrier-less path owing to the lack of energy barrier in the S<sub>1</sub> excited state.<sup>40</sup> After decaying to the ground state, a reverse proton transfer process from the keto to the enol takes place to produce the original E form. For **SPF-2**, the occurrence of the reverse proton transfer process is not as easy because of the existence of the 0.4 kcal mol<sup>-1</sup> (0.02 eV) energy barrier.<sup>40</sup> This result implied that the ESIPT process was feasible in the **SPF-1** and **SPF-2** molecules.

Moreover, the fluorescence properties of **SPF-1** and **SPF-2** in the enol and keto forms in DMF were simulated. The obtained fluorescence peaks of **SPF-1**-enol and **SPF-1**-keto are 404.4 and 522.72 nm, which is in excellent agreement with the experimental data of 425 and 528 nm (see Fig. 2, left). In the calculated fluorescence peaks of **SPF-2**, the emission bands 521.24 and 600.73 nm correspond to the enol and keto forms of **SPF-2**, respectively.<sup>41</sup> However, only a single emission band at 495 nm was observed in the experimental data. The average absolute

deviation between the theoretical (521.24 nm) and experimental (495) wavelengths was found to be 26.24 nm. The longer-wavelength emission band of **SPF-2** (> 495 nm) disappeared in DMF, which may be caused by the presence of intermolecular hydrogen bonding between the **SPF-2** and DMF molecules. The enol isomer of **SPF-2** is better stabilized by DMF molecules than the keto isomer and the ESIPT process is not possible.<sup>40</sup>

The time-resolved fluorescence decay behaviors of **SPF-1** (in DMSO/water solution, 9 : 1, v/v) and **SPF-2** (in 1,4-dioxane/water solution, 13 : 7, v/v) were investigated (Fig. S2 and S3, ESI<sup>†</sup>). Two relaxation pathways are involved in the decay process. The quick decay lifetimes are 3.63 and 0.82 ns, respectively, while the long decay lifetimes are 37.30 and 2.75 ns, respectively.<sup>30</sup> Therefore the ESIPT emission of **SPF-1** and **SPF-2** was further confirmed.

### 3.4 Detection of Zn<sup>2+</sup>

The corresponding solutions of **SPF-1** in DMF/H<sub>2</sub>O (9 : 1, v/v, 10 μM) and **SPF-2** in 1,4-dioxane/water (13 : 7, v/v, 10 μM) were prepared, respectively, in which the water volume percentage was less than 40% and the probe did not exhibit AIEE activity on its own.<sup>9</sup>

As depicted in Fig. 5, the **SPF** solutions show very weak fluorescence in the absence of Zn<sup>2+</sup>.<sup>42</sup> Upon addition of Zn<sup>2+</sup>,

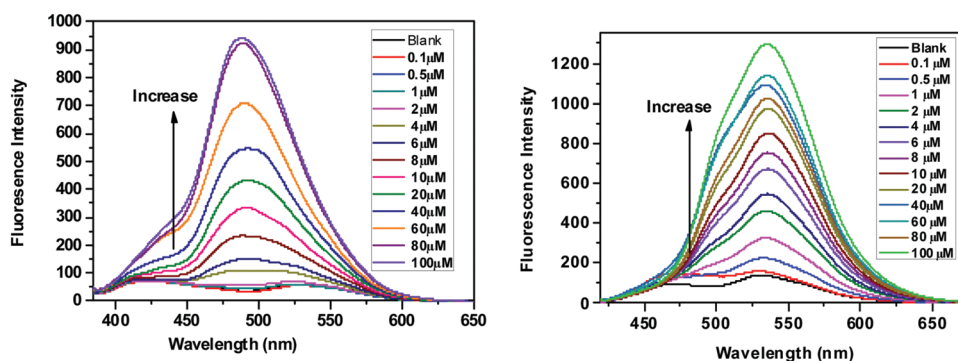


Fig. 5 Fluorescence spectra of **SPF-1** (10 μM) ( $\lambda_{\text{ex}} = 362$  nm) with varying concentrations of Zn<sup>2+</sup> in DMF/H<sub>2</sub>O (9 : 1, v/v) (left). Fluorescence spectra of **SPF-2** (10 μM) ( $\lambda_{\text{ex}} = 400$  nm) with varying concentrations of Zn<sup>2+</sup> in 1,4-dioxane/H<sub>2</sub>O (13 : 7, v/v) (right).



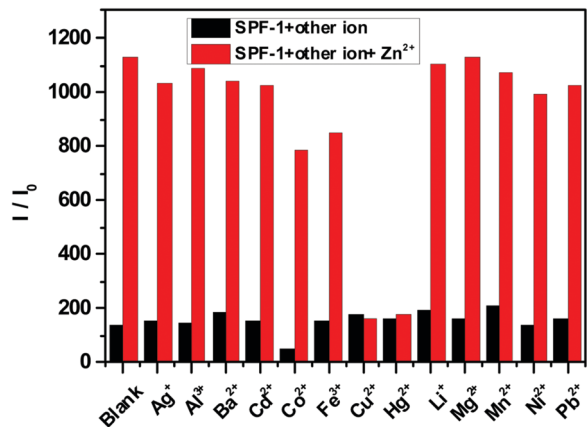


Fig. 8 The relative fluorescence intensity profiles ( $I/I_0$ ) of **SPF-1** (10  $\mu\text{M}$ , DMF/ $\text{H}_2\text{O}$ , 9:1, v/v) in the presence of 1 equiv.  $\text{Zn}^{2+}$  and the interfering ions ( $\text{Ag}^+$ ,  $\text{Al}^{3+}$ ,  $\text{Ba}^{2+}$ ,  $\text{Cd}^{2+}$ ,  $\text{Co}^{2+}$ ,  $\text{Fe}^{3+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Hg}^{2+}$ ,  $\text{Li}^+$ ,  $\text{Mg}^{2+}$ ,  $\text{Mn}^{2+}$ ,  $\text{Ni}^{2+}$ ,  $\text{Pb}^{2+}$ , respectively). Red bars: the addition of  $\text{Zn}^{2+}$  and various other metal ions. Black bars: the addition of various other metal ions.

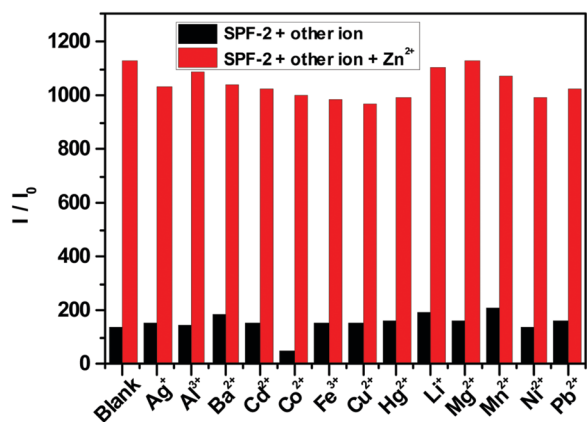


Fig. 9 The relative fluorescence intensity profiles ( $I/I_0$ ) of **SPF-2** (10  $\mu\text{M}$ , 1,4-dioxane/ $\text{H}_2\text{O}$ , 13:7, v/v) in the presence of 1 equiv.  $\text{Zn}^{2+}$  and the interfering ions ( $\text{Ag}^+$ ,  $\text{Al}^{3+}$ ,  $\text{Ba}^{2+}$ ,  $\text{Cd}^{2+}$ ,  $\text{Co}^{2+}$ ,  $\text{Fe}^{3+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Hg}^{2+}$ ,  $\text{Li}^+$ ,  $\text{Mg}^{2+}$ ,  $\text{Mn}^{2+}$ ,  $\text{Ni}^{2+}$ ,  $\text{Pb}^{2+}$ , respectively). Red bars: the addition of  $\text{Zn}^{2+}$  and various other metal ions. Black bars: the addition of various other metal ions.

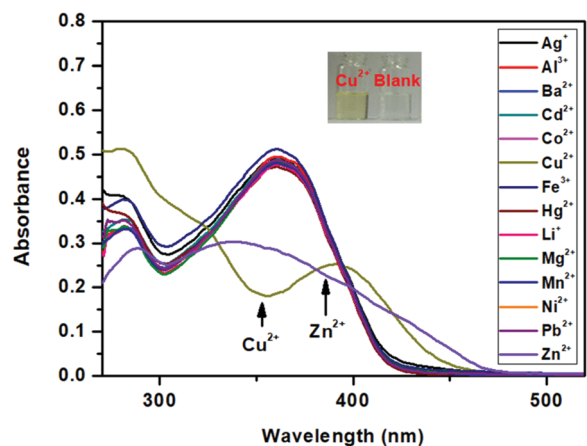


Fig. 10 The absorption spectra of **SPF-1** (10  $\mu\text{M}$ ) upon addition of various metal ions (10  $\mu\text{M}$ ) in DMF/ $\text{H}_2\text{O}$  (9:1, v/v).

fluorescence (CHEF), photoinduced electron transfer (PET), and C=N isomerization have been reported.<sup>9</sup>

The coordination of **SPF** with  $\text{Zn}^{2+}$  was confirmed by HRMS analysis. The reaction mixture of probe **SPF-2** with  $\text{Zn}^{2+}$  exhibited a prominent peak at  $m/z$  950.7093 (Fig. S8, ESI<sup>†</sup>), which is assigned to the species  $[\text{SPF-2} + 2\text{Zn}^{2+} - \text{C}(\text{CH}_3)_3]$ . As shown in Fig. S7 (ESI<sup>†</sup>), the molecular ion peaks corresponding to  $[\text{SPF-1} + 2\text{Zn}^{2+} - \text{C}(\text{CH}_3)_3 - 2\text{H}]$  at  $m/z$  679.4360,  $[\text{SPF-1} + 2\text{Zn}^{2+} - 2\text{CH}_3 + \text{H}]$  at  $m/z$  709.4486, and  $[\text{SPF-1} + 2\text{Zn}^{2+} - \text{CH}_3]$  at  $m/z$  723.4612 could be clearly observed. To expound the reason for the fluorescence light-up, DLS analysis was performed. As depicted in Fig. 11, the intensity-average hydrodynamic diameter,  $\langle D_h \rangle$ , of **SPF** is approximately 2.0 nm, which is in line with the molecular size. This indicates that **SPF-1** can be molecularly dissolved in DMF/water and **SPF-2** can be dissolved in a 1,4-dioxane/water mixture, respectively. However, the mean diameter was approximately 12 nm for **SPF-1** and 193 nm for **SPF-2** upon introduction of 2 equivalents of  $\text{Zn}^{2+}$ . Increasing the concentration of  $\text{Zn}^{2+}$  resulted in a significant enhancement in the average size of the aggregates (Fig. 11) and the fluorescence intensity, which supports the observed  $\text{Zn}^{2+}$ -triggered AIEE activity of the **SPF** probe.<sup>9,51</sup> The formation of aggregates can also be confirmed by the observation of a long absorption tail (see Fig. 6). On the basis of the experimental results described above, we proposed that the phenomenon caused by the addition of  $\text{Zn}^{2+}$  and resulting in the fluorescence enhancement could be attributed to the following reasons: (1) the coordination of **SPF** with  $\text{Zn}^{2+}$  can reduce the solubility of **SPF** and induce the formation of **SPF** aggregates; (2) the ESIPT process can be suppressed through a metal ion binding process;<sup>52–56</sup> (3) the binding of the nitrogen atom of C=N with  $\text{Zn}^{2+}$  restricted C=N isomerization in the excited states and inhibited the PET process of the C=N group within **SPF**.<sup>43</sup>

### 3.6 pH effect

We investigated the effect of pH on the sensing behavior of **SPF** towards  $\text{Zn}^{2+}$ . The variation of the fluorescence intensity with different pH values in the absence and presence of  $\text{Zn}^{2+}$  are shown in Fig. 12.

The fluorescence intensity of pure **SPF-1** (at 488 nm) remains almost unchanged in a wide pH range of 3–13. After addition of 1 equivalent of  $\text{Zn}^{2+}$ , the fluorescence intensity of **SPF-1** is enhanced gradually when the pH is increased from 3.0 to 6.1. At pH 6.1, the fluorescence intensity reached a plateau and did not change significantly upon changing the pH values from 6.1 to 10.5. At pH values higher than 10.5, a slight decline in the emission occurred, which could be attributed to the formation of  $\text{Zn}(\text{OH})_2$ .<sup>42</sup> In strong acidic solutions (pH < 3.0), the introduction of  $\text{Zn}^{2+}$  caused no changes in the fluorescence intensity of **SPF-1**, which could be attributed to the instability of the C=N bond under low pH value conditions.<sup>3</sup>

In acidic environments (pH < 6.0), the fluorescence of pure **SPF-2** (at 535 nm) increased significantly, which could be due to the suppressed PET processes from the protonated triphenylamine unit.<sup>6</sup> When the pH increased from 6.1 to 7.4, a slight decline in the emission can be observed. In basic conditions (pH > 7.4),

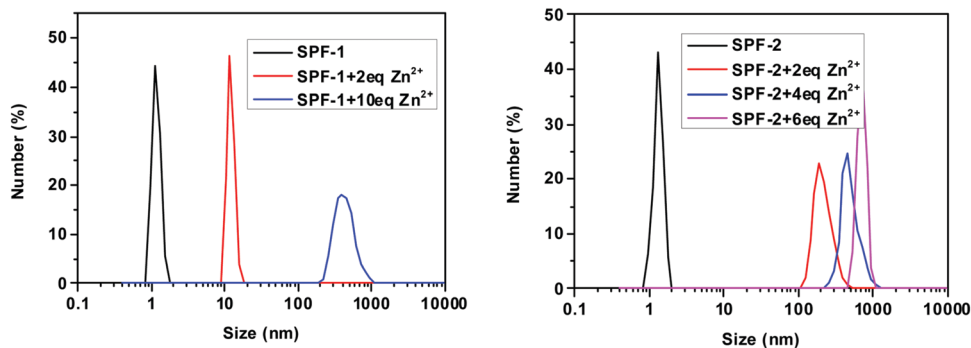


Fig. 11 DLS profile for **SPF-1** (left) (50  $\mu\text{M}$  in aqueous solution) and in the presence of 2.0 equiv. of  $\text{Zn}^{2+}$ , and 10.0 equiv. of  $\text{Zn}^{2+}$ , respectively. DLS profile for **SPF-2** (right) (50  $\mu\text{M}$  in aqueous solution) and in the presence of 2.0 equiv. of  $\text{Zn}^{2+}$ , 4.0 equiv. of  $\text{Zn}^{2+}$ , and 6.0 equiv. of  $\text{Zn}^{2+}$ , respectively.

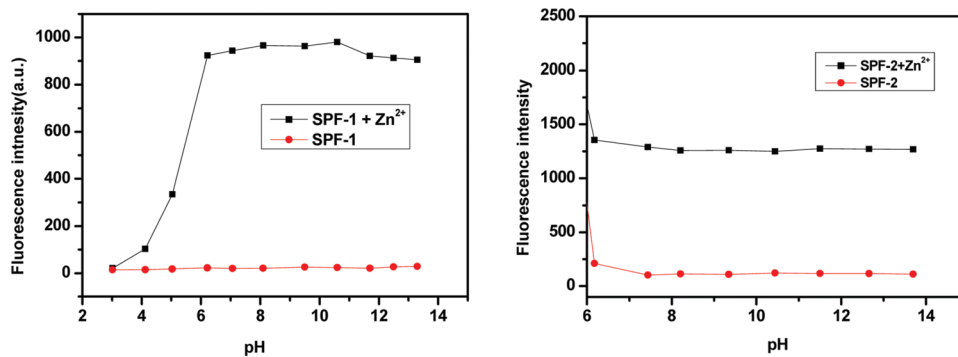


Fig. 12 Variation of fluorescence intensity at 488 nm for **SPF-1** solution with and without  $\text{Zn}^{2+}$  (1 equiv.) as a function of pH (left). Variation of fluorescence intensity at 535 nm for **SPF-2** solution with and without  $\text{Zn}^{2+}$  (1 equiv.) as a function of pH (right).

almost no changes in the fluorescence intensity were found. In a pH greater than 6.1, the emission intensity of **SPF-2** increased dramatically after addition of 1 equivalent of  $\text{Zn}^{2+}$ . The pH study shows that SPF could be used to monitor  $\text{Zn}^{2+}$  at a wide range of pH values and has the potential for application in conditions with a physiological pH (6.0–9.0).<sup>9</sup>

### 3.7 Two-photon absorption properties of SPF-2

In the structure of **SPF-2**, the triphenylamine group functions as an electron donor (D), the 2-(iminomethyl)phenol moiety acts as an electron acceptor (A) and the fluorene-vinyl moiety as a conjugation bridge. Therefore, the biphenyl branch containing a triphenylamine group is a donor- $\pi$ -acceptor (D- $\pi$ -A) type chromophore and is expected to serve as a source for the two-photon absorption. The TPA cross-section value of **SPF-2** was determined using a femtosecond open-aperture Z-scan technique in THF.<sup>23</sup> The open Z-scan data are shown in Fig. 13 and the  $\sigma$  of **SPF-2** was found to be 200 GM at a wavelength of 800 nm.

### 3.8 Cell imaging studies

The application of **SPF-1** for intracellular  $\text{Zn}^{2+}$  imaging (one-photon fluorescence) and **SPF-2** for two-photon fluorescence cell imaging were demonstrated, respectively. Before this, the cytotoxicities of **SPF-1** and **SPF-2** were first evaluated using MTT assays. After incubating the A549 cells with 50  $\mu\text{M}$  **SPF-1** or **SPF-2** for 24 h,

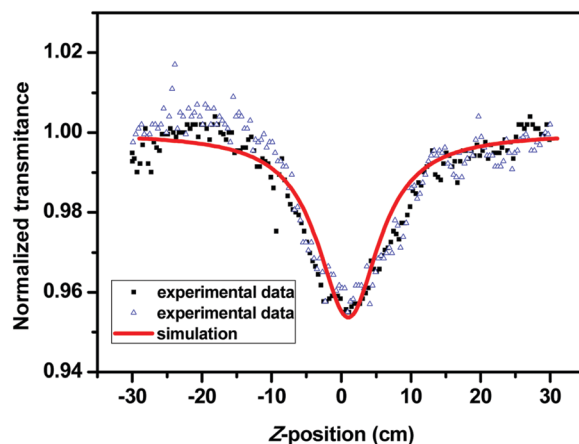


Fig. 13 Open Z-scan data of **SPF-2** in THF.

the cell viability remained at more than 95%, indicating the low cytotoxicity of the probe.<sup>57</sup>

Almost no intracellular fluorescence was observed for the **SPF-1** endocytosed cells. However, an intense green fluorescence was detected after these cells were treated with 100  $\mu\text{M}$   $\text{Zn}^{2+}$  solutions (Fig. 14). The results of this experiment indicated that **SPF-1** is potentially applicable for use in intracellular  $\text{Zn}^{2+}$  imaging.<sup>3</sup>

Two-photon excited fluorescence microscopy is a powerful method that is used for three-dimensional imaging in biological

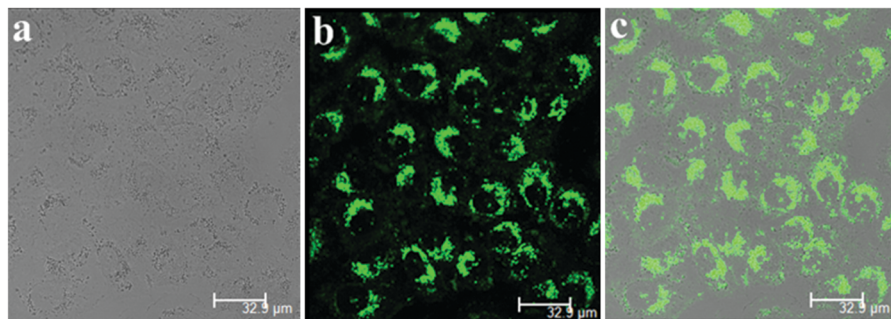


Fig. 14 Confocal fluorescence images of A549 cells after incubation with **SPF-1** (50  $\mu\text{M}$ ) for 30 min, and upon further incubation with  $\text{Zn}^{2+}$  (100  $\mu\text{M}$ ) for another 30 min. (a) Bright-field transmission image; (b) fluorescence image; and (c) overlay images of the A549 cells.

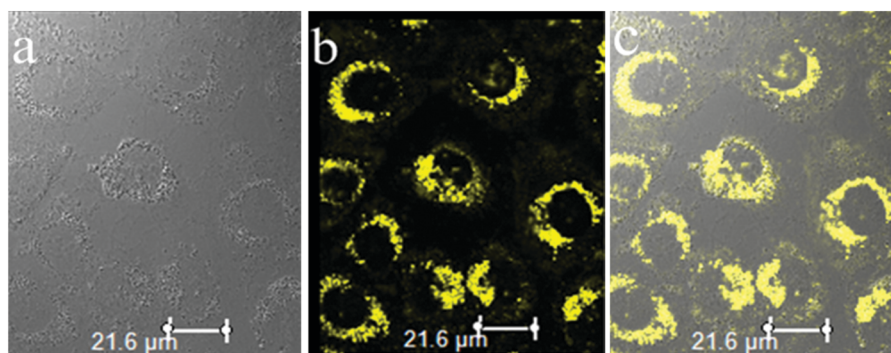


Fig. 15 Two-photon fluorescence images (at 800 nm) for A549 cells incubated with **SPF-2** (50  $\mu\text{M}$ ) for 30 min. (a) Bright field; (b) fluorescence; and (c) overlay images of the A549 cells.

systems owing to several distinct advantages, such as an increased penetration depth, lower tissue auto-fluorescence and self-absorption, and so on. The A549 cells treated with **SPF-2** (50  $\mu\text{M}$ ) show a bright yellow fluorescence using two-photon microscopy (at 800 nm) (Fig. 15). The results proved that **SPF-2** is suitable for two-photon fluorescence imaging of cells. As **SPF-2** is more hydrophobic than **SPF-1**, the **SPF-2** molecules aggregate more easily in the intracellular environment and emit a strong fluorescence signal.

## 4. Conclusions

In summary, two spirobifluorene-based probes for  $\text{Zn}^{2+}$  have been developed by taking advantage of their AIEE features. The sensing mechanism is ascribed to the chelation mediated AIEE behavior of **SPF**. **SPF-1** was successfully used for intracellular  $\text{Zn}^{2+}$  imaging and **SPF-2** was used for two-photon fluorescence cell imaging. The results demonstrated that spirobifluorene is an ideal building block for AIEE-based bioprobes. We hope that this molecular design triggers further interest in using spiro-configured compounds in AIEE-based bioprobes for further applications.

## Conflicts of interest

There are no conflicts to declare.

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