



Cite this: *RSC Adv.*, 2019, 9, 11865

Received 19th March 2019  
 Accepted 11th April 2019

DOI: 10.1039/c9ra02119a

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# Aggregation-induced emission enhancement (AIEE)-active tetraphenylethene (TPE)-based chemosensor for Hg<sup>2+</sup> with solvatochromism and cell imaging characteristics†

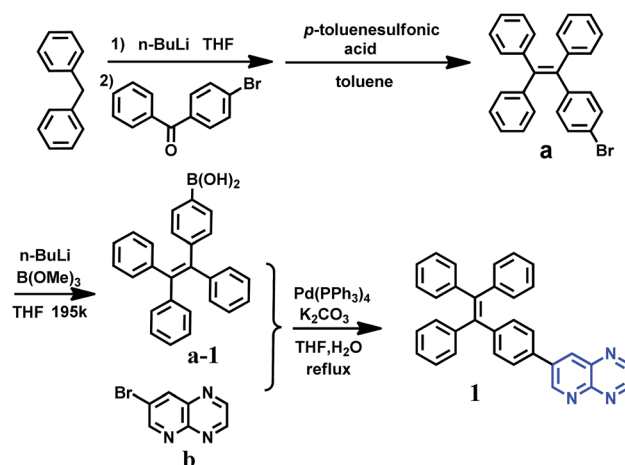
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An aggregation-induced emission enhancement (AIEE)-active fluorescent sensor based on a tetraphenylethene (TPE) unit has been successfully designed and synthesized. Interestingly, the luminogen could detect Hg<sup>2+</sup> with high selectivity in an acetonitrile solution without interference from other competitive metal ions, and the detection limit was 7.46 × 10<sup>-6</sup> mol L<sup>-1</sup>. Furthermore, the luminogen also showed interesting solvatochromic behavior and superior cell imaging performance.

The design and synthesis of molecular sensors for the detection of metal ions, especially transition-metal ions, has attracted much attention.<sup>1–3</sup> Among all transition-metal ions, mercury (Hg<sup>2+</sup>) is identified as one of the most dangerous and ubiquitous heavy metals. Indeed, it is not biodegradable, and can cause extreme injury to the environment as well as human health.<sup>4–8</sup> Additionally, it can be accumulated through the food chain in the human body, consequently giving rise to several deleterious effects such as central nervous system defects, kidney damage, endocrine system disease and so on.<sup>9–12</sup> Although many governments around the world have adopted strict regulations to limit Hg<sup>2+</sup> emission, the global Hg<sup>2+</sup> pollution caused by human activities is still serious.<sup>13,14</sup> Therefore, it is highly desirable to develop a new method for the detection of Hg<sup>2+</sup> with high selectivity and sensitivity.<sup>15–17</sup>

Most traditional fluorescent sensors suffer from a detrimental phenomenon called aggregation-caused quenching (ACQ), which usually results in a poor solid-state emission efficiency. Fortunately, in 2001, Tang *et al.* reported a fluorescent molecule named 1-methyl-1,2,3,4,5-pentaphenylsilole. Interestingly, the fluorescence emission of this luminogen was induced by aggregation, a phenomenon referred to as aggregation-induced emission (AIE).<sup>18</sup> Subsequently, in 2002, Park *et al.* reported an interesting phenomenon named aggregation-induced emission enhancement (AIEE).<sup>19</sup> In fact, both AIE and AIEE can achieve highly efficient emission in the solid state or aggregated state.<sup>20–27</sup> In the past years, AIE (or AIEE) phenomenon has attracted considerable research interest owing to the potential applications

in a lot of fields, including bioimaging, fluorescence sensors, organic lighting emitting diode (OLED) devices and organic lasers.<sup>28–33</sup> Meanwhile, many stimuli-responsive materials have been reported, including photochromism, mechanochromism, and solvatochromism.<sup>34–46</sup> At present, the solvatochromism materials have been generally used in chemical and biological systems.<sup>47</sup> To date, many fluorescent chemosensors for Hg<sup>2+</sup> have been reported. In contrast, the corresponding chemosensors with AIE or AIEE effect are rare, not to mention solvatochromic AIE or AIEE-active fluorescent sensors for Hg<sup>2+</sup> with good cell imaging behavior. Indeed, preparing such multifunctional sensors is challenging and significant. In this paper, we reported an AIEE-active tetraphenylethene (TPE)-based fluorescent sensor (Scheme 1) for the detection of Hg<sup>2+</sup>. Furthermore, the luminogen also showed remarkable solvatochromism and good cell imaging characteristics.



Scheme 1 The synthetic route of luminogen 1.

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† Electronic supplementary information (ESI) available: Experimental section, NMR spectra, mass spectrum, and characterization data mentioned in the paper. See DOI: 10.1039/c9ra02119a



To investigate the AIEE phenomenon of luminogen **1**, we initially studied the UV-visible absorption spectra and photoluminescence (PL) spectra in acetonitrile–water mixtures with different water fractions ( $f_w$ ). The results indicated that the absorption spectra exhibited level-off tails in the long wavelength region as the water content increased (Fig. S1†). It is well-known that such tails are usually associated with the Mie scattering effect, which is the key signal of nano-aggregate formation.<sup>48,49</sup> As presented in Fig. 1, luminogen **1** showed weak fluorescence and the luminescence quantum yield ( $\Phi$ ) was as low as 1.35%. Interestingly, when the  $f_w$  in the acetonitrile solution was increased to 80%, an obvious emission band was formed, and a yellow-green fluorescence was observed. As the water content was increased to 90%, the emission intensity was further increased, and a bright yellow-green luminescence ( $\Phi = 27.81\%$ ) with a  $\lambda_{\text{max}}$  at 545 nm could be observed.

Clearly, water is a poor solvent of luminogen **1**. As a result, the generation of the yellow-green emission can be attributed to the aggregate formation.<sup>50–52</sup> Moreover, as shown in Fig. 2, the nano-aggregates obtained were verified by dynamic light scatterings (DLS). Therefore, **1** is a typical AIEE-active fluorescent molecule, and its AIEE behavior is caused by the restricted intramolecular rotation. As shown in Fig. S2,† solid-state compound **1** showed a strong green emission ( $\Phi = 14.50\%$ ) with a  $\lambda_{\text{max}}$  at 497 nm, and the corresponding fluorescence lifetime is 2.66 ns (Fig. S3†).

On the other hand, the luminogen **1** also displayed interesting solvatochromism effect. As presented in Fig. 3, the absorption spectra and fluorescence spectra of luminogen **1** in different solvents were investigated, respectively. Obviously, the absorption spectra was barely affected by the polarity of solvents. However, the photoluminescence peaks were gradually red-shifted from 515 nm to 625 nm, and thus **1** exhibited remarkable solvatochromism behavior. Obviously, the

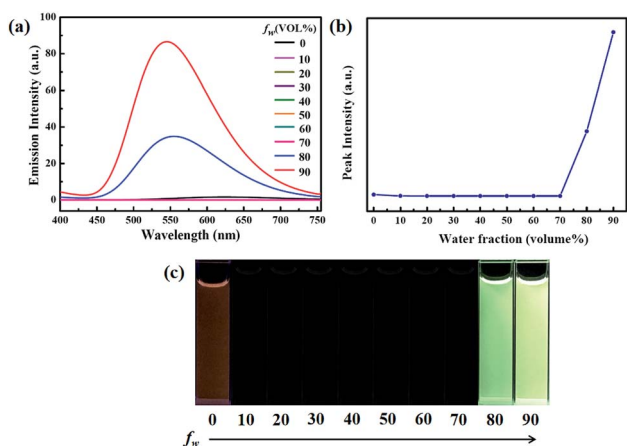


Fig. 1 (a) Fluorescence spectra of the dilute solutions of **1** ( $2.0 \times 10^{-5} \text{ mol L}^{-1}$ ) in acetonitrile–water mixtures with different water content (0–90%). Excitation wavelength = 380 nm. (b) Changes the emission intensity of **1** at 545 nm in acetonitrile–water mixtures with different volume fractions of water (0–90%). (c) PL images of **1** ( $2.0 \times 10^{-5} \text{ mol L}^{-1}$ ) in acetonitrile–water mixtures with different  $f_w$  values under 365 nm UV illumination.



Fig. 2 Size distribution curve of luminogen **1** ( $2.0 \times 10^{-5} \text{ mol L}^{-1}$ ) in acetonitrile–water mixtures with  $f_w = 90\%$ .

molecular structure of luminogen **1** is distorted due to the presence of TPE unit, and the conjugation degree of molecule **1** is different in various solvents, and the intramolecular charge transfer (ICT) effect is possibly responsible for the solvatochromism behavior of **1**.

Subsequently, the changes in the fluorescence of luminogen **1** induced by  $\text{Hg}^{2+}$  were investigated in acetonitrile ( $2.0 \times 10^{-5} \text{ mol L}^{-1}$ ) at room temperature. In the fluorometric titration experiments, as shown in Fig. 4, the emission intensity significantly decreased when the  $\text{Hg}^{2+}$  concentration increased from 0 to 7.0 equivalents in acetonitrile. Meanwhile, the fluorescent color changed from orange-red to colorless, and followed by a plateau upon further titration (Fig. 5). Remarkably, the emission intensity of luminogen **1** was almost quenched completely. Furthermore, based on the titration experiments, the detection limit of luminogen **1** for  $\text{Hg}^{2+}$  on the basis of  $\text{LOD} = 3 \times \sigma/B$  (where  $\sigma$  is the standard deviation of blank sample and  $B$  is the slope between the fluorescence intensity versus  $\text{Hg}^{2+}$  concentration) was  $7.46 \times 10^{-6} \text{ M}$  (Fig. S4†). Moreover,



Fig. 3 (a) UV-Vis absorption spectra and (b) normalized fluorescence spectra (Excitation wavelength = 380 nm) of luminogen **1** in different solvents ( $2.0 \times 10^{-5} \text{ mol L}^{-1}$ ). (c) Photographs of **1** under 365 nm UV illumination in different solvents: tol (toluene); THF (tetrahydrofuran); EA (ethyl acetate); BT (acetone); MeCN (acetonitrile); DMSO (dimethyl sulfoxide).





Fig. 4 Fluorescence titration spectra of luminogen 1 ( $2.0 \times 10^{-5} \text{ mol L}^{-1}$ ) induced by  $\text{Hg}^{2+}$  (0–7.0 equiv.) in an acetonitrile solution. Excitation wavelength = 380 nm.



Fig. 5 The emission intensity changes of luminogen 1 at 625 nm with different equivalents of  $\text{Hg}^{2+}$ .

a good linear relationship could be obtained ( $R = -0.9933$ ) and the quenching constant of luminogen 1 with  $\text{Hg}^{2+}$  was  $1.9 \times 10^4 \text{ M}^{-1}$  (Fig. S5<sup>†</sup>). On the other hand, the binding ratio of luminogen 1 for  $\text{Hg}^{2+}$  was established through Job's plot and the results showed 1 : 1 stoichiometric complexation (Fig. S6<sup>†</sup>). Next, the binding interactions between 1 and  $\text{Hg}^{2+}$  were further investigated by NMR in acetonitrile- $d_3$ . As presented in Fig. 6, the signal of Ha from 9.42 ppm shifted to 9.62 ppm and the Hb changed from 8.43 ppm to 9.24 ppm. Besides, the Hc or Hd was slightly shifted for 0.04 ppm or 0.06 ppm, respectively. These consequences revealed that the N on the pyridine and the N on the pyrazine (near the N on the pyridine) are the most probable binding with  $\text{Hg}^{2+}$ . Mass spectra were utilized to further demonstrate the binding mode of luminogen 1 toward  $\text{Hg}^{2+}$ . The peak located at  $m/z = 820.0$  was coincided well with the ensemble  $[\text{1} + \text{Hg}^{2+} + 2\text{NO}_3^- + \text{Cl}^-]^-$ , confirming the binding ratio of luminogen 1 for  $\text{Hg}^{2+}$  with 1 : 1 stoichiometry (Fig. S7<sup>†</sup>).

Next, in order to study the selectivity behavior of luminogen 1 as a fluorescent sensor for  $\text{Hg}^{2+}$ , other metal ions, such as  $\text{Zn}^{2+}$ ,  $\text{Cd}^{2+}$ ,  $\text{Ba}^{2+}$ ,  $\text{Sr}^{2+}$ ,  $\text{Mn}^{2+}$ ,  $\text{Mg}^{2+}$ ,  $\text{Ca}^{2+}$ ,  $\text{Pb}^{2+}$ ,  $\text{Ni}^{2+}$ ,  $\text{Co}^{2+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Al}^{3+}$ ,  $\text{Fe}^{3+}$ ,  $\text{Cr}^{3+}$ ,  $\text{Ag}^+$  and  $\text{K}^+$  were also measured in acetonitrile

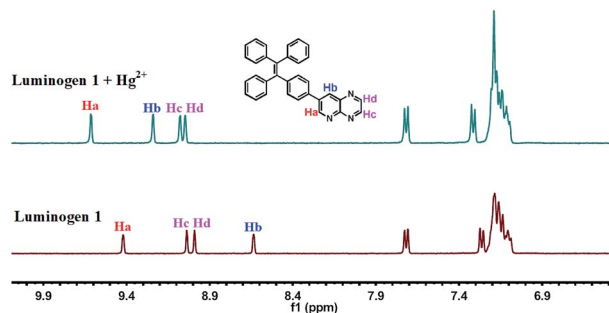


Fig. 6  $^1\text{H}$  NMR (acetonitrile- $d_3$ , 400 MHz) spectra changes of luminogen 1 in the presence of  $\text{Hg}^{2+}$ .

under the same experimental conditions. The corresponding UV-Vis absorption spectra were shown in Fig. S8<sup>†</sup>. Furthermore, as showed in Fig. 7, when these cations were added separately into the solution containing luminogen 1, no obvious fluorescence changes were observed. Indeed, as noticed in Fig. S9<sup>†</sup>, no obvious interference was observed when  $\text{Hg}^{2+}$  (7.0 equiv.) was added with other ions (7.0 equiv.). These results indicated that luminogen 1 could be served as a highly selective fluorescent sensor for detection of  $\text{Hg}^{2+}$ .

Fluorescent probe is a powerful tool for optical imaging, which allows direct visualization of biological analytes.<sup>53</sup> Luminogen 1 was AIEE-active due to the restriction of intramolecular rotation in the aggregate state, which is beneficial for cell imaging. Indeed, the viability of HeLa cells incubated with luminogen 1 was evaluated by the standard MTT method (Fig. 8), and the result indicated that compound 1 exhibited low cytotoxicity. Next, cell imaging behavior of luminogen 1 was investigated using a confocal laser scanning microscopy (CLSM). HeLa cells were incubated with luminogen 1 (20  $\mu\text{M}$ ) for 30 min at 37  $^\circ\text{C}$  and the fluorescence images were obtained by CLSM.

As presented in Fig. 9, an intense yellow-green fluorescence, which was consistent with the fluorescence of luminogen 1 in

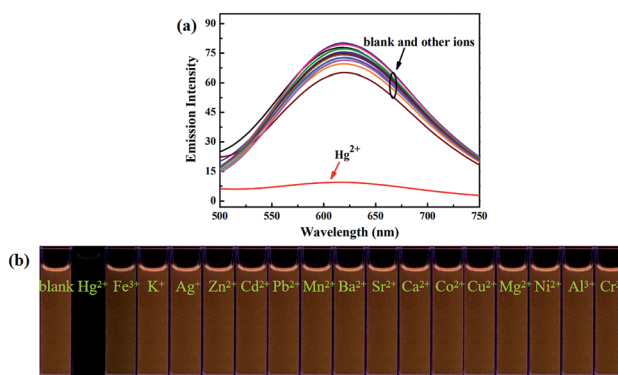


Fig. 7 (a) Fluorescence spectra of luminogen 1 ( $2.0 \times 10^{-5} \text{ mol L}^{-1}$ ) towards various cations including  $\text{Zn}^{2+}$ ,  $\text{Cd}^{2+}$ ,  $\text{Hg}^{2+}$ ,  $\text{Ba}^{2+}$ ,  $\text{Sr}^{2+}$ ,  $\text{Mn}^{2+}$ ,  $\text{Mg}^{2+}$ ,  $\text{Ca}^{2+}$ ,  $\text{Pb}^{2+}$ ,  $\text{Ni}^{2+}$ ,  $\text{Co}^{2+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Al}^{3+}$ ,  $\text{Fe}^{3+}$ ,  $\text{Cr}^{3+}$ ,  $\text{Ag}^+$  and  $\text{K}^+$ . Excitation wavelength = 380 nm. (b) Fluorescence photographs of luminogen 1 after addition of various metal ions (7.0 equiv.) under 365 nm light.





Fig. 8 The MTT assay of compound 1 for measuring cell viability.

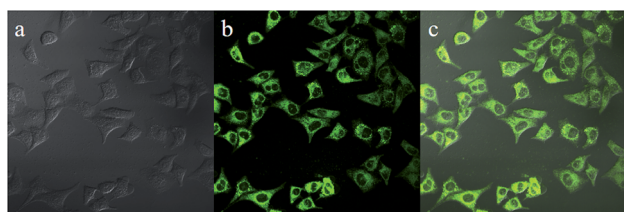


Fig. 9 Fluorescence images of HeLa cells incubated with luminogen 1 (20 μM) for 30 min at 37 °C: (a) bright field image; (b) fluorescence image; (c) merge image (a and b).

acetonitrile–water mixture with high water content (80% or 90%), was displayed inside the cells. This result indicated that luminogen 1 tended to aggregate inside the cells, and thus the bright yellow-green luminescence was clearly observed. Furthermore, the merged picture c demonstrated that picture a and picture b overlapped very well. These results indicated that luminogen 1 showed superior cell imaging performance.

In summary, a TPE-based fluorescent molecule was designed and synthesized. The luminogen exhibited obvious AIEE phenomenon. Moreover, luminogen 1 could be used as a highly selective fluorescence turn-off chemosensor for  $\text{Hg}^{2+}$ , and the detection limit was  $7.46 \times 10^{-6} \text{ mol L}^{-1}$ . Furthermore, 1 also displayed interesting solvatochromic behavior and superior cell imaging performance. Further studies on the design of new AIE or AIEE-active fluorescent chemosensors are in progress.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The authors are grateful for the financial support from the National Natural Science Foundation of China (21702079 and 41867053), the “5511” Science and Technology Innovation Talent Project of Jiangxi Province (20165BCB18015), the key project of Natural Science Foundation of Jiangxi Province (20171ACB20025), the youth project of Natural Science

Foundation of Jiangxi Province (20171BAB213004), the Young Talents Project of Jiangxi Science and Technology Normal University (2017QNBjRC005), and the Project of the Science Fund of Jiangxi Education Office (GJJ180593).

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