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# Development of tissue paper-based chemosensor and demonstration for the selective detection of Cu<sup>2+</sup> and Hg<sup>2+</sup> ions†

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Heavy metals emanate from natural and man-made sources, such as agricultural chemicals including fertilisers and pesticides, medical waste, and chemicals released from industries. Detection and monitoring toxic metal ions is one of the challenges confronting scientists in biological, environmental, and chemical systems. This study describes the design and synthesis of a new imidazole-based fluorescent and colourimetric chemosensor (DPICDT) for highly selective sensing of Hg<sup>2+</sup> and Cu<sup>2+</sup> ions in aqueous acetonitrile medium. The probe was synthesised by coupling benzil and substituted aldehyde using ethanolic ammonium acetate. The structure of DPICDT was confirmed via IR spectra, NMR, and HR-MS spectra. The DPICDT probe displayed a rapid naked-eye response towards Cu<sup>2+</sup> ions from colourless to red-purple and significant fluorescence quenching response towards Hg<sup>2+</sup> over other competitive metal ions in both solution and solid support. The binding modes of DPICDT with Cu<sup>2+</sup> and Hg<sup>2+</sup> ions were found to be at a 1:1 ratio as determined using Job plot, ESI HR-MS, and the sensing mechanism was evolved by <sup>1</sup>H NMR titrations, HR-MS spectra, and DFT calculations. The lower detection limit was 15.1 nM for Cu<sup>2+</sup>, eventually far less than the World Health Organization guideline for drinking water (Cu<sup>2+</sup> – 31.5 μM) and 1.17 μM for Hg<sup>2+</sup> (permissible concentration 2 ppb). Promisingly, the tissue paper-based DPICDT test strips and silica-supported DPICDT were developed and demonstrated for on-site application without resorting to expensive instruments.

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## Introduction

In recent years, heavy metals have been an environmental threat and are of grave concern worldwide and a growing threat to humanity. Among all heavy metals, copper (Cu) and mercury (Hg) are potentially toxic. Copper is a critically important transition metal for many fundamental physiological processes and in regulating the action of some of the coenzymes in humans, such as superoxide dismutase, tyrosinase, and cytochrome C oxidase.<sup>1–3</sup> Copper can be stored in the living tissues of plants, animals, and humans. However, overloading of copper in

humans would induce a sequence of neurodegenerative diseases, such as Wilson, Alzheimer, Parkinson's, and metabolic disorders including obesity and diabetes.<sup>4,5</sup> In addition, gratuitous metal ions lead to environmental pollution, which soberly threatens the health of microorganisms, plants, animals, and even humans.<sup>6</sup> Mercury, a dormant contaminant in the environment, has been a global concern because of its toxicity. Notably, Hg<sup>2+</sup> ions can quickly accumulate in living beings. They can bind with thiols and amino groups in proteins and enzymes, which may further induce diseases such as kidney failure, tumour formation, motor disorders, and neurological damage such as prenatal brain damage.<sup>7,8</sup> Considering an overview of these ions in day-to-day life, developing techniques for sensing and monitoring these ions is in great demand.

Various analytical methods have been used for the determination of metal ions, including nuclear techniques,<sup>9,10</sup> gravimetry,<sup>11</sup> electrochemical techniques,<sup>12,13</sup> ion chromatography,<sup>14</sup> γ-ray spectrometry,<sup>15</sup> inductively coupled plasma/mass spectrometry,<sup>16</sup> capillary zone electrophoresis,<sup>17</sup> and using cation-exchange resin<sup>18</sup> and extraction.<sup>19</sup> However, these methods have many drawbacks, including high cost, intricate sample preparation, time-consuming, and require skilled human resources. To overcome these limitations, chemical sensors were used to determine

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metal ions.<sup>20,21</sup> The alternative and more sensitive methods that can swiftly and selectively measure the metal ions in low concentrations in different media are difficult. Mainly, amphoteric imidazole is a well-known organic compound, and its substituted analogues have been ardently studied due to its simple synthetic tactic, superior yield, effortless purification, desirable photophysical properties, and ease of integration of the artificial binding sites towards various analytes.

Although some of the sensors have been reported to detect  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$ , most of them either could not access discrimination within the same measurement method or differentiate both using masking reagents.<sup>22,23</sup> Unfortunately, to the best of our knowledge, a simple sensor that can detect both  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  within the same sample with various responses is yet to be reported. With this in mind herewith, we developed a simple, facile, and rapid DPICDT chemosensor with specific selectivity and sensitivity towards  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  over other interfering metal ions, leading to predominant naked eye colour change from colourless to pink for  $\text{Cu}^{2+}$  and fluorescence off response for  $\text{Hg}^{2+}$  under UV light ( $\lambda = 365 \text{ nm}$ ). Interestingly, the tissue paper-based DPICDT test strips and solid-supported DPICDT@ $\text{SiO}_2$  were developed and demonstrated the scope of DPICDT for practical sensing applications for  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  ions.

## Experimental section

### Materials and instrumentation

The chloride and nitrate salts of  $\text{Hg}^{2+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Ca}^{2+}$ ,  $\text{Ce}^{3+}$ ,  $\text{Co}^{2+}$ ,  $\text{Ba}^{2+}$ ,  $\text{Bi}^{2+}$ ,  $\text{Li}^+$ ,  $\text{Mg}^{2+}$ ,  $\text{Mn}^{2+}$ ,  $\text{Mo}^{2+}$ ,  $\text{Na}^+$ ,  $\text{Ni}^{2+}$ ,  $\text{Pb}^{2+}$ ,  $\text{Sr}^{2+}$ ,  $\text{Al}^{3+}$ ,  $\text{Zr}^{2+}$ ,  $\text{Zn}^{2+}$ , all reagents, and solvents (analytical and spectroscopic grade) were purchased from Sigma Aldrich, TCI chemicals, Avra and were used without further purification. Deionised water was used throughout the experiments. The stock solutions were prepared by dissolving the analytes in deionised water. The reaction monitoring and purity determination of the substrates were accompanied by thin layer chromatography using silica gel 60 F254 plates (Merck Chemical Company, USA). The melting point was measured on an electro-thermal melting point apparatus. For column chromatography, 60–120 mesh silica gel was used to purify the product. FT-IR spectra were measured using a JASCO-4100 spectrophotometer.  $^1\text{H}$  NMR and  $^{13}\text{C}$  NMR spectra were obtained on Bruker 400 MHz spectrometer, and the chemical shifts are expressed in  $\delta$  ppm using tetramethylsilane (TMS) as an internal standard. High-resolution mass spectrometry (HRMS) was obtained on a Joel GC Mate II GC-Mass spectrometer. Absorption spectra were recorded using a Jasco V-750 UV-Vis spectrophotometer, and emission spectra were obtained on a Jasco FP-8300 instrument. Atomic absorption spectrometer (AAS – Varian Spectra 240) was used to determine the metal concentration. Fluorescent quantum yields were measured against anthracene ( $\phi_{\text{fl}} = 0.28$  in ethanol) as the fluorescence standard.

### Preparation of stock solutions

For spectroscopic measurements, a stock solution of  $20 \mu\text{M}$  DPICDT was prepared in ( $\text{CH}_3\text{CN}:\text{H}_2\text{O}$ ) (8 : 2, v/v) and various

metal ion solutions,  $\text{Hg}^{2+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Ca}^{2+}$ ,  $\text{Ce}^{3+}$ ,  $\text{Co}^{2+}$ ,  $\text{Ba}^{2+}$ ,  $\text{Bi}^{2+}$ ,  $\text{Li}^+$ ,  $\text{Mg}^{2+}$ ,  $\text{Mn}^{2+}$ ,  $\text{Mo}^{2+}$ ,  $\text{Na}^+$ ,  $\text{Ni}^{2+}$ ,  $\text{Pb}^{2+}$ ,  $\text{Sr}^{2+}$ ,  $\text{Al}^{3+}$ ,  $\text{Zr}^{2+}$ ,  $\text{Zn}^{2+}$  (1 mM), were prepared using deionised water.

### Fabrication DPICDT loaded paper strips

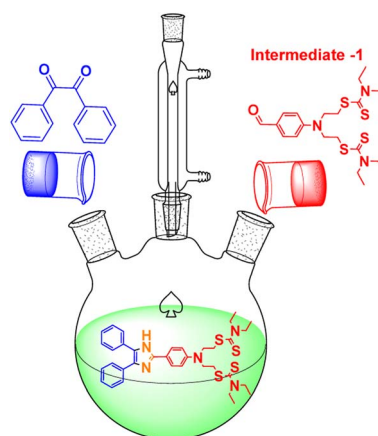
DPICDT (10.0 mg) was dissolved in 10 mL of ethanol. Strips of tissue paper ( $2 \times 2 \text{ cm}$ ) were immersed in the above solution for 1 min, then placed on a clean glass plate, and allowed to dry naturally at ambient temperature. In this way, we obtained the DPICDT-loaded tissue paper strips.

## Results and discussion

### Chemistry

**Synthesis of ((4-formylphenyl)azanediyl)bis(ethane-2,1-diyl)bis(diethylcarbamodithioate) (intermediate-1).** A mixture of 4-(bis(2-chloroethyl)amino)benzaldehyde (2.5 g, 10 mmol) and sodium diethyldithiocarbamate (4.3 g, 25 mmol) was dissolved in acetonitrile (25 mL) at room temperature. The mixture was heated to reflux for about 24 h. The reaction mixture was cooled to room temperature, and the solution was poured into 100 mL of ice water and filtered to give a light brown solid. The crude product was recrystallized in ethanol to yield intermediate-1 as crystalline light brown solid<sup>24</sup> (4 g, 88%)  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ ,  $\delta$ ): 9.74 (s 1H, CHO) 7.76–7.73 (d, 2H, Ar H), 7.07–7.05 (d, 2H, Ar H), 4.07–4.02 (t, 4H,  $\text{CH}_2$ ), 3.80–3.55 (m, 8H,  $\text{CH}_2$ ), 3.53–3.51 (t, 4H,  $\text{CH}_2$ ), 1.31–1.27 (t, 12H,  $\text{CH}_3$ ) (Fig. S1†);  $^{13}\text{C}$  NMR (400 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 194.47, 190.23, 152.34, 132.34, 125.79, 111.60, 49.86, 49.78, 46.94, 32.93, 12.50, 11.60 (Fig. S2†). HRMS (ESI): calcd for  $\text{C}_{21}\text{H}_{34}\text{N}_3\text{OS}_4$  472.1579 found 472.1593 (Fig. S3†).

**Synthesis of ((4-(4,5-diphenyl-1H-imidazole-2-yl)phenyl)azanediyl)bis(ethane-2,1-diyl)bis(diethylcarbamodithioate) [DPICDT].** The DPICDT chemosensor was achieved by a previously reported literature procedure with minor modifications using a simple single-step condensation reaction of aldehyde and ketone with base, as illustrated in Scheme 1.<sup>25</sup> To a stirred solution of benzil (1 g, 2.1 mmol) in ethanol (20 mL), intermediate-1 [0.45 g, 2.1 mmol] was added, followed by the addition of



**Scheme 1** Synthesis of ((4-(4,5-diphenyl-1H-imidazole-2-yl)phenyl)azanediyl)bis(ethane-2,1-diyl)bis(diethylcarbamodithioate) [DPICDT].



ammonium acetate (1.6 g, 21 mmol). The reaction mixture was stirred and heated to reflux for 4 h. The course of the reaction was monitored by thin layer chromatography. After completion, the reaction mixture was cooled and poured into crushed ice. The precipitate was filtered and dried. The obtained solid was further purified by column chromatography using hexane and ethyl acetate (15%) as eluent. The desired DPICDT was obtained as a dirty white solid with an 85% yield.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ ,  $\delta$ ): 12.3 (s 1H, NH) 7.99–7.77 (d, 2H, Ar H), 7.54–7.52 (d, 2H, Ar H), 7.33–7.25 (m, 2H, Ar H), 7.01–6.99 (d, 2H, Ar H), 4.05–4.02 (t, 4H,  $\text{CH}_2$ ), 3.78–3.70 (m, 8H,  $\text{CH}_2$ ), 3.54–3.50 (t, 4H,  $\text{CH}_2$ ), 1.31–1.27 (t, 12H,  $\text{CH}_3$ ) (Fig. S4†);  $^{13}\text{C}$  NMR (400 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 194.92, 147.87, 128.51, 127.84, 127.18, 126.83, 117.91, 112.23, 49.97, 49.69, 46.91, 33.31, 12.53, 11.63 (Fig. S5†). HRMS (ESI): calcd for  $\text{C}_{35}\text{H}_{43}\text{N}_5\text{S}_4$  662.004 found 662.2448 (Fig. S6†).

### Colourimetric and spectral response of DPICDT to $\text{Cu}^{2+}$

The spectroscopic properties of DPICDT in ( $\text{CH}_3\text{CN}:\text{H}_2\text{O}$ ) (8 : 2, v/v) solution were investigated with absorption and fluorescence studies. Absorption spectra of the DPICDT were measured in the presence of different metal ions like  $\text{Hg}^{2+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Ca}^{2+}$ ,  $\text{Ce}^{3+}$ ,  $\text{Co}^{2+}$ ,  $\text{Ba}^{2+}$ ,  $\text{Bi}^{2+}$ ,  $\text{Li}^+$ ,  $\text{Mg}^{2+}$ ,  $\text{Mn}^{2+}$ ,  $\text{Mo}^{2+}$ ,  $\text{Na}^+$ ,  $\text{Ni}^{2+}$ ,  $\text{Pb}^{2+}$ ,  $\text{Sr}^{2+}$ ,  $\text{Al}^{3+}$ ,  $\text{Zr}^{2+}$ , and  $\text{Zn}^{2+}$ . As shown in (Fig. 1), the UV-Vis absorption of DPICDT (20  $\mu\text{M}$ ) exhibits two bands, high energy  $\lambda_{\text{max}}$  325 nm attributed to  $\pi$ - $\pi^*$  transitions and a shoulder peak at 283 nm.

The binding interactions of DPICDT towards diverse metal ions show a selective response and significant absorption changes towards  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  ions. However, none of the other metal ions caused a substantial effect in the absorption

band of DPICDT. When DPICDT was treated with  $\text{Cu}^{2+}$  ion, the peaks at 283 and 325 nm disappeared and induced a significant bathochromic shift with a new strong absorption band at 525 nm due to ligand-metal charge transfer. Meanwhile, the colour of the DPICDT probe solution turned from colourless to red-purple after the addition of  $\text{Cu}^{2+}$  ions.

The response was reasonably fast, and we could observe a distinct colour change while adding. Thus, DPICDT can be used to detect  $\text{Cu}^{2+}$  ions by the naked eye.<sup>26</sup> In addition, the colour change could be explained by the LMCT mechanism. The band with the molar extinction coefficient in the thousands ( $26 \times 10^3 \text{ M}^{-1} \text{ cm}^{-1}$  at 520 nm) is too large to be a Cu-based d-d transition and thus must be a ligand-based transition.<sup>27</sup> An isosbestic point was identified at 355 nm, which indicates a DPICDT- $\text{Cu}^{2+}$  complex formation.

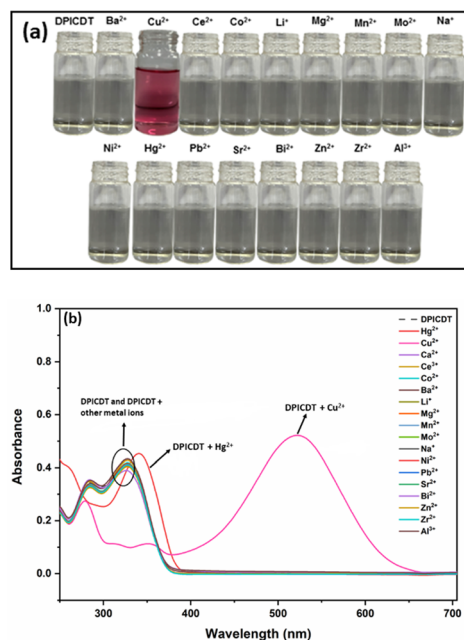
To investigate the DPICDT- $\text{Cu}^{2+}$  interaction, the absorption titration of DPICDT (20  $\mu\text{M}$ ) by incremental addition of  $\text{Cu}^{2+}$  ion (1 mM) was recorded in ( $\text{CH}_3\text{CN}:\text{H}_2\text{O}$ ) (8 : 2, v/v) solution (Fig. 2). UV-Vis titration showed a gradual extinction of the 325 nm signal in the presence of increasing amounts of  $\text{Cu}^{2+}$  and a slight increase of the 495 nm signal. Effectively, the absorbance at 325 nm decreases and that of 495 nm increases gradually upon adding one equivalent of  $\text{Cu}^{2+}$ . Therefore, an M1L complex seems to be formed during the titration.

Fluorescence emission spectra of DPICDT are recorded upon excitation at 325 nm to understand the nature of interactions of metal ions in the excited state (Fig. 3). DPICDT shows an emission band at 421 nm, and the emission peak decreased completely upon the addition of  $\text{Cu}^{2+}$  ion. The paramagnetic nature of  $\text{Cu}^{2+}$  ions can be expected to quench the excited state of DPICDT.<sup>28</sup> Adding 2.5 equivalence of  $\text{Cu}^{2+}$  ions led to a 30-fold decrease in the 421 nm fluorescence intensity of DPICDT.

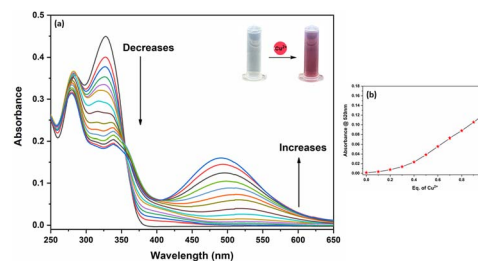
For a better understanding of the quenching mechanism, the fluorescence spectral titrations of DPICDT (20  $\mu\text{M}$ ) were performed with incremental addition of  $\text{Cu}^{2+}$  in ( $\text{CH}_3\text{CN}:\text{H}_2\text{O}$ ) (8 : 2, v/v). As shown in (Fig. 4), the fluorescence intensity of DPICDT centered at 421 nm gradually reduced and shifted to 403 nm with increasing concentration of  $\text{Cu}^{2+}$ , which eventually became constant at two equivalents of  $\text{Cu}^{2+}$ .

### Spectral response of DPICDT to $\text{Hg}^{2+}$

The sensing ability of DPICDT was checked against  $\text{Hg}^{2+}$  ions using UV-Vis and fluorescence spectroscopy techniques. As



**Fig. 1** (a) DPICDT (20  $\mu\text{M}$ ) in the presence of various analytes (1 mM) in ( $\text{CH}_3\text{CN}:\text{H}_2\text{O}$ ) (8 : 2, v/v) under daylight. (b) Specific selectivity of DPICDT (20  $\mu\text{M}$ ) in the presence and absence of diverse cations (1 mM) in ( $\text{CH}_3\text{CN}:\text{H}_2\text{O}$ ) (8 : 2, v/v).



**Fig. 2** (a) UV-Vis titration of (20  $\mu\text{M}$ ) DPICDT in ( $\text{CH}_3\text{CN}:\text{H}_2\text{O}$ ) (8 : 2, v/v) with increasing amounts of  $\text{Cu}^{2+}$  (1 mM) (b) absorbance plotted vs. equiv. of  $\text{Cu}^{2+}$  at 500 nm.



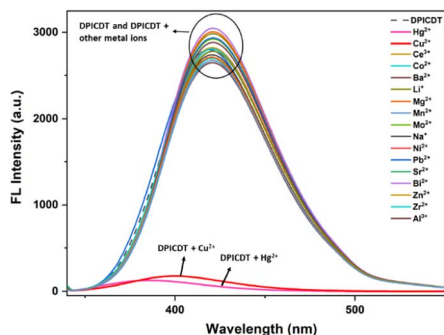


Fig. 3 Fluorescence spectra of DPICDT (20  $\mu$ M) in the presence and absence of diverse cations (1 mM) in ( $\text{CH}_3\text{CN} : \text{H}_2\text{O}$ ) (8 : 2, v/v).

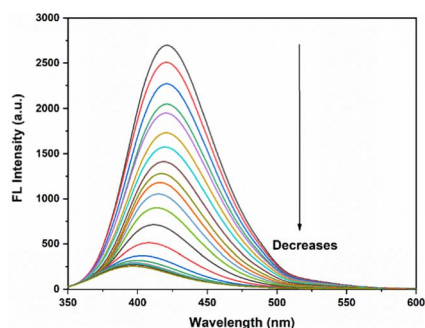


Fig. 4 Fluorescence titration of (20  $\mu$ M) DPICDT in ( $\text{CH}_3\text{CN} : \text{H}_2\text{O}$ ) (8 : 2, v/v) with increasing amounts of  $\text{Cu}^{2+}$  (1 mM)  $\lambda_{\text{ext}} = 325$  nm.

shown in (Fig. 1) the peak at 283 nm disappeared, and the peak at 325 nm was red-shifted to 340 nm with a slight increase in absorbance while adding  $\text{Hg}^{2+}$ . Furthermore, the UV-Vis titration of DPICDT with  $\text{Hg}^{2+}$  is shown in (Fig. 5a). With the addition of increasing amounts of  $\text{Hg}^{2+}$  to a solution of DPICDT in ( $\text{CH}_3\text{CN} : \text{H}_2\text{O}$ ) (8 : 2, v/v), the isosbestic point at 333 nm and 270 nm confirmed the formation of the complex between DPICDT and  $\text{Hg}^{2+}$ .

The selectivity of the DPICDT sensor to  $\text{Hg}^{2+}$  was further investigated by fluorometric detection in ( $\text{CH}_3\text{CN} : \text{H}_2\text{O}$ ) (8 : 2, v/v), which shows a significant decrease in the emission intensity with a slight shift observed upon adding  $\text{Hg}^{2+}$ . After the successive addition of 0–2.5 equivalents of  $\text{Hg}^{2+}$ , the emission at 420 nm was entirely quenched, and a weak emission peak at 400 nm was found (Fig. 5b).

In contrast, the fluorescence behaviour of DPICDT in THF decreased with increasing concentration of DMSO. For instance, upon excitation, fluorescence spectrum of dilute DPICDT in THF showed a fluorescence emission band at 410 nm (Fig. S7†). Interestingly, increasing the concentration of DMSO in DPICDT exhibits a minor spectral shift from 410 to 415 nm with a decrease in emission intensity. This decrease in fluorescence intensity was ascribed to an increase in the dipole moment of the TICT state, which on interaction with a polar solvent (DMSO) promotes the intramolecular motions.<sup>29</sup>

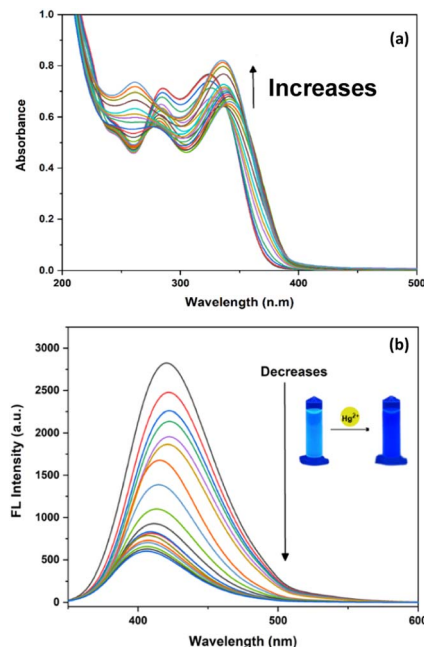


Fig. 5 (a) UV-Vis titration of (20  $\mu$ M) DPICDT in ( $\text{CH}_3\text{CN} : \text{H}_2\text{O}$ ) (8 : 2, v/v) with increasing amounts of  $\text{Hg}^{2+}$ ; (b) fluorescence titration of (20  $\mu$ M) DPICDT in ( $\text{CH}_3\text{CN} : \text{H}_2\text{O}$ ) (8 : 2, v/v) with increasing amounts of  $\text{Hg}^{2+}$  (1 mM)  $\lambda_{\text{ext}} = 325$  nm.

### Metal binding and sensing mechanism

The binding ratio of DPICDT to  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  was further confirmed by the Job's plot. The Job's plot results are illustrated in (Fig. S8†), and 1 : 1 stoichiometry was calculated for both DPICDT- $\text{Cu}^{2+}$  and DPICDT- $\text{Hg}^{2+}$ . The stoichiometry was further confirmed by HR-MS analysis. The positive-ion ESI mass indicated peaks at  $m/z$  721.2675 and  $m/z = 1059.4265$ , which were assigned to  $[\text{DPICDT} + \text{Cu}^{2+} - 3\text{H}^+]^+$  (calcd, 724.1697) (Fig. S9†) and  $[\text{DPICDT} + \text{Hg}(\text{ClO}_4)_2 + \text{H}^+]^+$  (calcd, 1061.1089) (Fig. S10†), respectively.

From the above outcomes, it was feasible to determine the dissociation constants ( $K_a$ ) for the interaction of DPICDT with  $\text{Cu}^{2+}$  and DPICDT with  $\text{Hg}^{2+}$  using the Benesi-Hildebrand plot. The dissociation constant value of  $\text{Hg}^{2+}$  with DPICDT was found to be  $3.85 \times 10^6 \text{ M}^{-1}$  and  $2.71 \times 10^6 \text{ M}^{-1}$  for  $\text{Cu}^{2+}$  (Fig. S11†). From this equation,  $\text{LOD} = 3\sigma/K$  (where  $\sigma$  is the standard deviation of the blank measurement, and  $K$  is the slope of the linear plot, which is emission intensity *versus* concentration), and the limit of detection was computed to be 15.1 nM for  $\text{Cu}^{2+}$  and 1.17  $\mu\text{M}$  for  $\text{Hg}^{2+}$  (Fig. S12†). The fluorescence quantum yield was determined using anthracene as a reference with a known  $\Phi_R$  of 0.27 in ethanol. The quantum yield values were found to be 0.155 for DPICDT, 0.0051 for DPICDT- $\text{Cu}^{2+}$ , and for DPICDT- $\text{Hg}^{2+}$  is 0.0087. The obtained 30-fold and 17-fold decrease in quantum yield supports the fluorescence quenching of DPICDT on binding with  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$ , respectively.

To further investigate the binding mechanism, the expected complex formation was further confirmed using FT-IR and  $^1\text{H}$ -NMR spectra. The FT-IR spectrum of DPICDT with and





without  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  were measured and illustrated in (Fig. S13†). The DPICDT peaks at  $694\text{ cm}^{-1}$ ,  $1263\text{ cm}^{-1}$ ,  $1348\text{ cm}^{-1}$ , and  $2975\text{ cm}^{-1}$  were attributed to the stretching vibration of C–S, C=S,  $(\text{CH}_3)_2\text{N}-\text{C}=\text{S}$  and NH, respectively. However, in the FT-IR spectrum of DPICDT- $\text{Cu}^{2+}$ , the NH peak broadened and was observed at  $3150\text{ cm}^{-1}$ ; this further confirms that the imine group was involved in coordination with  $\text{Cu}^{2+}$ . In addition, in the FT-IR spectrum of DPICDT- $\text{Hg}^{2+}$ , the characteristic peaks of C–S, C=S, and  $(\text{CH}_3)_2\text{N}-\text{C}=\text{S}$  were broadened and merged. These obvious changes clearly indicate that the C=S in the carbamodithioate group of DPICDT was involved in coordination with  $\text{Hg}^{2+}$  ions.

To clarify the coordination mode of DPICDT,  $^1\text{H}$  NMR titration was executed in  $\text{DMSO}-d_6$ . The DPICDT showed signals at 12.3 ppm due to NH proton ( $\text{H}_e$ ); the peaks at 7.9–7.0 ppm and 4.0–3.3 ppm are attributed to aromatic and methylene protons ( $\text{H}_b$ ,  $\text{H}_c$ , and  $\text{H}_d$ ), respectively. The peak at 1.2 ppm corresponds to the methyl protons ( $\text{H}_a$ ). The spectral data before and after adding  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  are depicted in (Fig. 6a and b). Upon complexation with  $\text{Cu}^{2+}$ , the imine proton signal at 12.3 ppm decreased gradually and slightly shifted downfield to 12.52 ppm. Similarly, with the addition of  $\text{Hg}^{2+}$  to DPICDT, the aliphatic proton ( $\text{H}_a$ ,  $\text{H}_b$ ,  $\text{H}_c$ , and  $\text{H}_d$ ) signals broadened and shifted downfield. These results showed that the coordination occurs with thione containing dithiocarbamate group. According to Pearson's hard – soft acid – base theory,  $\text{Hg}^{2+}$  ions have a high affinity towards

sulfur atoms, which was distinctly determined by the fluorescence quenching effect. Thus, based on the above facts, Scheme 2 depicts the plausible coordination modes of DPICDT with  $\text{Hg}^{2+}$  and  $\text{Cu}^{2+}$  ions.

### Theoretical calculation methods

The electronic properties of DPICDT and its complexes with  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  were investigated through theoretical analysis using DFT calculations. The optimized structure of DPICDT is shown in (Fig. 7).

To determine the reactive sites of the ligand, molecular electrostatic potential surface (ESP) calculations were performed for the metal atom binding, as illustrated in (Fig. 8). The ESP plot revealed regions of higher negative potential around the sulfur ( $-\text{C}=\text{S}$ ) and nitrogen atoms of the ligand, suggesting that these sites are the most likely binding sites for the  $\text{Hg}^{2+}$  and  $\text{Cu}^{2+}$  ions, respectively. Additionally, the Frontier molecular orbitals (FMOs) of the ligand and its  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  complexes (DPICDT- $\text{Cu}^{2+}$ , DPICDT- $\text{Hg}^{2+}$ ) were analysed and are presented in (Fig. 8). The FMO analysis of DPICDT and its  $\text{Cu}^{2+}$  complex indicated that the charge density of the free ligand primarily resides on the imidazole part. However, the FMO plots of  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  complexes showed L–M charge transfer for  $\text{Cu}^{2+}$  and twisted intramolecular charge transfer for  $\text{Hg}^{2+}$ . This charge transfer, accompanied by a significant reduction in the HOMO–LUMO band gap, is likely responsible for the shift in the UV absorption band.

### Reversibility studies

To increase the applicability of the chemosensor DPICDT, it was significant to test its reversibility for the detection of cations. The reversible performance of DPICDT towards  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  was studied using ethylenediamine tetraacetic acid (EDTA) and KI in UV-Vis and FL experiments, respectively, as shown in (Fig. 9). The DPICDT solution is colourless and when

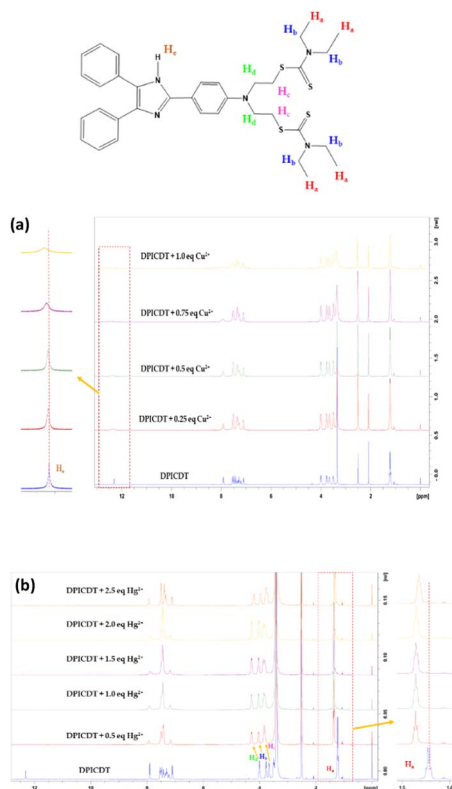
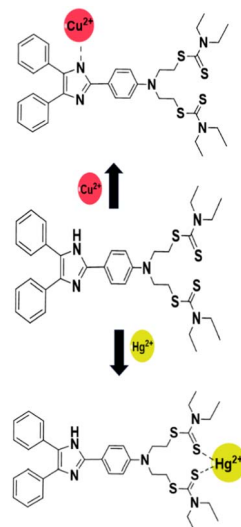


Fig. 6  $^1\text{H}$  NMR spectra of DPICDT (400 Hz,  $\text{DMSO}-d_6$ ) in the absence and presence of  $\text{Cu}^{2+}$  (0 to 1 equiv.) (a) and  $\text{Hg}^{2+}$  (0 to 2.5 equiv.) (b).



Scheme 2 Plausible binding of DPICDT with  $\text{Hg}^{2+}$  and  $\text{Cu}^{2+}$  ions.

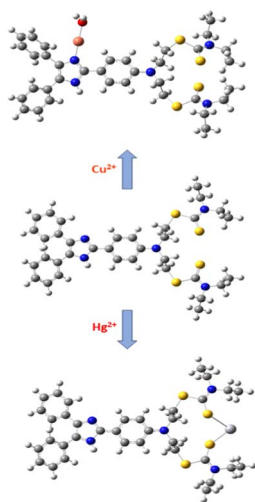


Fig. 7 DFT optimised DPICDT, DPICDT- $\text{Cu}^{2+}$ , and DPICDT- $\text{Hg}^{2+}$  complexes.

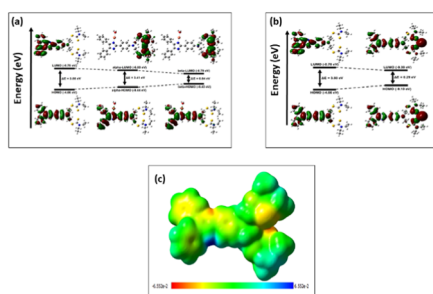


Fig. 8 (a) Frontier molecular orbitals of the DPICDT probe and its  $\text{Cu}^{2+}$  complex (DPICDT- $\text{Cu}^{2+}$ ). (b) Frontier molecular orbitals of DPICDT and its  $\text{Hg}^{2+}$  complex (DPICDT- $\text{Hg}^{2+}$ ). (c) Molecular electrostatic potential (MEPS) formed by mapping the total density over electrostatic potential in the gas phase of DPICDT.

it binds  $\text{Cu}^{2+}$  ions, the colour is changed to red-purple. However, adding the chelating agent ethylenediamine tetraacetic acid into the  $\text{Cu}^{2+}$ -DPICDT solution resulted in a colour change from colourless to red-purple, indicating the recovery of DPICDT. Upon successive addition of ethylenediamine tetraacetic acid solution to  $\text{Cu}^{2+}$ -DPICDT, the maximum absorption peak at 520 nm decreased, and it almost resorted to the original DPICDT signal, indicating the de-complexation.

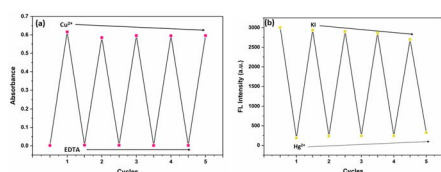


Fig. 9 (a) Reversible changes in colour and of DPICDT at 520 nm in  $\text{CH}_3\text{CN} : \text{H}_2\text{O}$  (8 : 2, v/v) upon alternate addition of  $\text{Cu}^{2+}$  and EDTA; (b) Reversible changes in fluorescence intensity of DPICDT upon alternate addition of  $\text{Hg}^{2+}$  and KI.

$\text{Hg}^{2+}$  induces a remarkable fluorescence change by showing OFF behaviour through complex formation ( $\text{Hg}^{2+}$ -DPICDT). In addition, the fluorescence ON response corresponding to DPICDT was recovered upon adding the KI solution. The KI addition increased the intensity and again decreased with negligible intensity loss. This could be attributed to the formation of stable mercury iodide and the consequent release of free DPICDT. The repeated demonstration of the ON/OFF behaviour of the system by fluorescence as well as the absorbance change, clearly suggests that DPICDT is reversible and a reusable sensor.

### Interference study

DPICDT was treated with various interfering cations to test its practical ability toward  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  using UV-Vis spectrometer and fluorescence spectrometer. As shown in Fig. 10, various metal ions had negligible effects.<sup>30</sup> These findings suggested that DPICDT could be used as a better colorimetric sensor for  $\text{Cu}^{2+}$  and fluorometric sensor for  $\text{Hg}^{2+}$  ions.

### Solid matrix – extraction performance of DPICDT towards $\text{Cu}^{2+}$ & $\text{Hg}^{2+}$

The extraction efficiency is an essential factor in the design and synthesis of novel chemosensors to improve their practical applicability in industries. A solid-supported material (DPICDT@ $\text{SiO}_2$ ) was fabricated to study the extraction performance of DPICDT towards  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$ , which is displayed in (Fig. 11). The solid-supported matrix was prepared by adsorbing DPICDT onto commercially available silica gel ( $\text{SiO}_2$ , 60–120 mesh) in a 1 : 4 w/w ratio. The resulting light brown powders (DPICDT@ $\text{SiO}_2$ ) were collected and utilised as extractants in column chromatography. The extraction ability of DPICDT was estimated by measuring the amount of  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  individually using AAS, resulting in >99% of  $\text{Cu}^{2+}$  and 100% of  $\text{Hg}^{2+}$  ions being extracted by DPICDT@ $\text{SiO}_2$ . Upon the addition of 100 ppm of  $\text{Cu}^{2+}$  solution through solid-supported DPICDT@ $\text{SiO}_2$ , the colour of solid support changed from light brown (DPICDT@ $\text{SiO}_2$ ) to chestnut ( $\text{Cu}^{2+}$ -DPICDT@ $\text{SiO}_2$ ), and the resulting copper-free fractions were collected. Furthermore, the ethylenediamine tetraacetic acid (EDTA) solution was passed through the column; as expected, the blue colour confirms the  $\text{Cu}^{2+}$ -EDTA complex formation, and the  $\text{Cu}^{2+}$  free DPICDT@ $\text{SiO}_2$  bed was fully recovered. Similarly, by adding 100 ppm of  $\text{Hg}^{2+}$  solution, light brown DPICDT@ $\text{SiO}_2$  changed to bright yellow, instantly confirming the ( $\text{Hg}^{2+}$ -DPICDT@ $\text{SiO}_2$ )

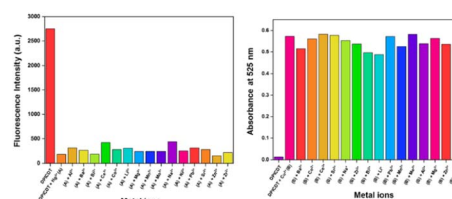


Fig. 10 Bar diagram showing interference from other metal ions during detection of  $\text{Hg}^{2+}$  (left) and  $\text{Cu}^{2+}$  (right) using DPICDT.



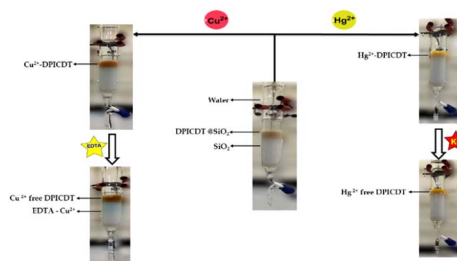


Fig. 11 Demonstration of extracting metal ions through column chromatography method.

complex formation. The addition of KI solution into the column resulted in  $\text{Hg}^{2+}$  free DPICDT@ $\text{SiO}_2$ . The AAS result suggests that DPICDT is potentially useful as a stationary phase for separating  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  using column chromatography (Fig. S14†).

### Cost-effective tissue paper-based sensor application

We prepared simple tissue paper-based test strips to demonstrate the DPICDT sensor application towards the on-site application of spiked samples. The  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  spiked solutions were prepared individually using deionised water. The DPICDT was dissolved in ethanol, and then ordinary tissue papers were immersed into the solution and incubated for 1 min. The collected tissue papers were dried in air at room temperature and utilised to sense various metal ions. A noticeable change was only observed for  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  solutions. As depicted in (Fig. 12), the colour of the DPICDT-coated tissue paper strip changed from colourless to red-purple with  $\text{Cu}^{2+}$  solution under the naked eye. Similarly, the colour intensity decreased when  $\text{Hg}^{2+}$  was added to the DPICDT tissue paper strip under the 365 nm UV lamp. Furthermore, to study the sensitivity of DPICDT (Fig. 13), DPICDT fabricated tissue paper was immersed in different concentrations of  $\text{Cu}^{2+}$  &  $\text{Hg}^{2+}$  ( $1 \times 10^{-3}$  M,  $1 \times 10^{-6}$  M,  $1 \times 10^{-9}$  M). The color change occurred quickly, and the rate of changing the color of the tissue paper strip increased proportionally with increasing  $\text{Cu}^{2+}$  concentration, and the color intensity was decreased proportionally with increasing  $\text{Hg}^{2+}$  concentration. Therefore, tissue paper-based DPICDT test strips can instantly detect  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  ions in water samples without the requirements of any analytical instruments.

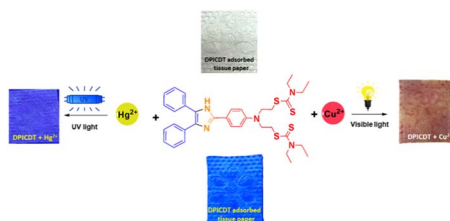


Fig. 12 Photographs of tissue paper strips containing DPICDT for detecting  $\text{Cu}^{2+}$  in natural light (right) and  $\text{Hg}^{2+}$  under 365 nm UV light (left).

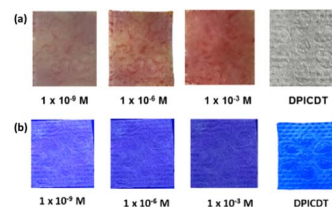


Fig. 13 Colorimetric response and fluorometric response of DPICDT on tissue paper with various concentrations of (a)  $\text{Cu}^{2+}$  and (b)  $\text{Hg}^{2+}$  ions.

## Conclusion

In summary, we have successfully designed and synthesised a novel imidazole fragment (DPICDT), which could serve as a dual chemosensor for  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  in the presence of various other competitive ions. DPICDT displayed a colourimetric response towards  $\text{Cu}^{2+}$  and a fluorescence turn-off response towards  $\text{Hg}^{2+}$  with two different sensing approaches. The analytical detection limit of  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  were 15.1 nM and 1.17  $\mu\text{M}$ , respectively, obtained from the linear plot, which is relatively lower upon comparing recent literature reviews (Tables S1 & S2†). Furthermore, DPICDT has been utilised successfully for  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  extraction through a solid-supported matrix. UV-Vis, ESI-MS, and NMR studies thoroughly examined the complex formation, stoichiometry, and binding mode, which show the 1:1 complex formation of DPICDT with both  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$ . Moreover, the easy-to-prepare tissue paper test strip provided a convenient and reliable detection of  $\text{Cu}^{2+}$  and  $\text{Hg}^{2+}$  for practical applications. Here, a single molecular system acts as a colourimetric sensor for  $\text{Cu}^{2+}$  and an ON-OFF fluorescence sensor for  $\text{Hg}^{2+}$ , integrating the chemosensor principle.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- M. H. Mahnashi, A. M. Mahmoud, S. A. Alkahtani, R. Ali and M. M. El-Wakil, *Spectrochim. Acta, Part B*, 2020, **228**, 117846.
- Y. Cao, Y. Liu, F. Li, S. Guo, Y. Shui, H. Xue and L. Wang, *Microchem. J.*, 2019, **150**, 104176.
- F. N. Moghadam, M. Amirnasr, S. Meghdadi, K. Eskandari, A. Buchholz and W. Plass, *Spectrochim. Acta, Part A*, 2019, **207**, 6–15.



- 4 A. Mohammadi and Z. Ghasemi, *Spectrochim. Acta, Part A*, 2020, **228**, 117730.
- 5 Y. S. Kim, G. J. Park, S. A. Lee and C. Kim, *RSC Adv.*, 2015, **5**, 31179–31188.
- 6 L. L. Yang, A. L. Tang, P. Y. Wang and S. Yang, *Org. Lett.*, 2020, **22**, 8234–8239.
- 7 P. Marimuthu, B. Umamahesh, S. Baskar, K. Sathiyarayanan, B. Venkatachalapathy, C. Ravichandran and N. S. Karthikeyan, *New J. Chem.*, 2018, **42**, 8530–8536.
- 8 V. Raju, R. S. Kumar, S. A. Kumar, G. Madhu, S. Bothra and S. K. Sahoo, *J. Chem. Sci.*, 2020, **132**, 1–11.
- 9 M. B. Rodríguez, F. V. Tomé, J. C. Lozan and V. G. Escobar, *Appl. Radiat. Isot.*, 2000, **52**, 705–710.
- 10 E. J. Bouwer, J. W. McKlveen and W. J. McDowell, *Nucl. Technol.*, 1979, **42**, 102–111.
- 11 C. Kósa, M. Danko and P. Hrdlovič, *J. Fluoresc.*, 2012, **22**, 1371–1381.
- 12 J. N. Li, F. Y. Yi, Z. M. Jiang and J. J. Fei, *Mikrochim. Acta*, 2003, **143**, 287–292.
- 13 V. K. Gupta, A. K. Jain and P. Kumar, *Sens. Actuators, B*, 2006, **120**, 259–265.
- 14 E. H. Borai and A. S. Mady, *Appl. Radiat. Isot.*, 2002, **57**, 463–469.
- 15 A. Pantelica, I. I. Georgescu, M. D. Murariu-Magureanu, I. Margaritescu and E. Cincu, *Radiat. Prot. Dosim.*, 2001, **97**, 187–191.
- 16 E. A. Casartelli and N. Miekeley, *Anal. Bioanal. Chem.*, 2003, **377**, 58–64.
- 17 B. F. Liu, L. B. Liu and J. K. Cheng, *Talanta*, 1998, **47**, 291–299.
- 18 A. K. Jain, V. K. Gupta, U. Khurana and L. P. Singh, *Electroanalysis*, 1997, **9**, 857–860.
- 19 N. El-Hefny and J. Daoud, *J. Radioanal. Nucl. Chem.*, 2004, **261**, 357–363.
- 20 B. K. Rani and S. A. John, *New J. Chem.*, 2017, **41**, 12131–12138.
- 21 K. Tayade, A. Kaur, S. Tetgure, G. K. Chaitanya, N. Singh and A. Kuwar, *Anal. Chim. Acta*, 2014, **852**, 196–202.
- 22 V. Merz, J. Merz, M. Kirchner, J. Lenhart, T. B. Marder and A. Krueger, *Chem. - Eur. J.*, 2021, **27**, 8118–8126.
- 23 H. Cho, J. B. Chae and C. Kim, *ChemistrySelect*, 2019, **4**, 2795–2801.
- 24 X. M. Meng, L. Liu, H. Y. Hu, M. Z. Zhu, M. X. Wang, J. Shi and Q. X. Guo, *Tetrahedron Lett.*, 2006, **47**, 7961–7964.
- 25 J. Jayram and V. Jeena, *RSC Adv.*, 2018, **8**, 37557–37563.
- 26 M. P. Bhat, M. Kigga, H. Govindappa, P. Patil, H. Y. Jung, J. Yu and M. Kurkuri, *New J. Chem.*, 2019, **43**, 12734–12743.
- 27 Y. S. Kim, G. J. Park, S. A. Lee and C. Kim, *RSC Adv.*, 2015, **5**, 31179–31188.
- 28 Y. Xiang, A. Tong, P. Jin and Y. Ju, *Org. Lett.*, 2006, **8**, 2863–2866.
- 29 S. Sharma, A. Gupta, C. P. Pradeep and A. Dhir, *ChemistrySelect*, 2017, **2**, 10517–10523.
- 30 R. Arumugaperumal, V. Srinivasadesikan, M. C. Lin, M. Shellaiah, T. Shukla and H. C. Lin, *RSC Adv.*, 2016, **6**, 106631–106640.

