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# Specific colorimetric detection of Fe<sup>3+</sup> ions in aqueous solution by squaraine-based chemosensor†

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A new squaraine based chemosensor **TSQ** was developed for colorimetric detection of Fe<sup>3+</sup> ions. A thymine moiety in **TSQ** was constructed to act as an ion acceptor. The sensor displayed an instant colorimetric response specific to Fe<sup>3+</sup> over the other metal ions in 20% AcOH–H<sub>2</sub>O solution. The limit of detection was much lower than that of the environmental protection agency guideline (5.37 μM) in drinking water. A 1 : 1 binding between **TSQ** and Fe<sup>3+</sup> ion was evidenced by Job's plot measurement, ESI-MS and Fourier transform infrared (IR) measurements. Moreover, the proposed sensing mechanism of the receptor towards Fe<sup>3+</sup> was strongly supported by DFT calculation. Finally, the sensor has proven to be suitable in real sample applications.

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## 1 Introduction

For the past years, the development of highly sensitive and selective chemosensors has been attracting great interests for detecting heavy metals due to their importance in environmental, biology and chemistry domains.<sup>1–10</sup> Iron, the third most abundant element on the earth, is not only the ubiquitous element in the environment, but also an essential element in living cells. It plays crucial roles in functional biological processes, such as cellular metabolism, enzymatic reactions, oxygen transport and gene expression.<sup>11–16</sup> Both deficiency and excess can cause a series of diseases.<sup>17</sup> Current qualitative and quantitative determination methods for Fe<sup>3+</sup> ions include atomic absorption spectroscopy, inductively coupled plasma-atomic emission spectroscopy, mass spectrometry, electrochemical and fluorescence spectroscopic analysis.<sup>18–22</sup> These methods suffer either from the sophisticated nature and high cost of instruments, or from tedious sample preparation procedures.<sup>23,24</sup> It is imperative to develop simple, highly sensitive and selective chemosensors for Fe<sup>3+</sup> detection. To date, a few chemosensors have been developed based on functionalized nanoparticles or conjugated organic chromophores.<sup>25–32</sup> However, few of them can be used as

practical probes limited by their low sensitivity and selectivity, as well as incompatibility with aqueous environments.<sup>33–37</sup> For example, Hg<sup>2+</sup>, Zn<sup>2+</sup> as well as Cu<sup>2+</sup>, which exhibit many properties similar to those of Fe<sup>3+</sup>, could interfere the Fe<sup>3+</sup> detection.<sup>38,39</sup> Also it is noted that not many chemosensors with long absorption wavelength have been reported so far. Hence, to develop efficient colorimetric sensing materials still remains a big challenge.

Squaraines are versatile organic dyes which have shown intense absorption in near infrared region and have been widely applied in the optoelectronics fields such as light emitting diodes, field-effect transistors and chemosensors.<sup>40–46</sup> Taken advantages of their unique properties, squaraine-based ion sensors can be constructed by a combination of squaraine chromophores with ligands. The intramolecular charge transfer will be occurred once metal ions have interacted with designed ion receptor. This could result in a rich modulation of color changes.

Thymine is one of the four nucleobases to perform base pairing with adenine in constructing double helix DNA. It has proven to be one of the most effective ligands binding to the metal ions especially for mercury ion.<sup>47</sup> Therefore, we introduced a thymine group into the squaraine chromophore, leading to a new thymine-squaraine based chemosensor. To our surprise, the resulting chemosensor can specifically detect Fe<sup>3+</sup> over the other metal ions by color change in aqueous solution. The complexation to Fe<sup>3+</sup> and sensing mechanism were further investigated by Job's plot measurement, UV-vis titration, ESI-MS spectrometry analysis, IR analysis and theoretical calculation. The improvement in the color response, rapid accurate recognition of Fe<sup>3+</sup> and low detection limit make this approach very promising for Fe<sup>3+</sup> detection.

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## 2 Experimental section

### 2.1 Reagents and apparatus

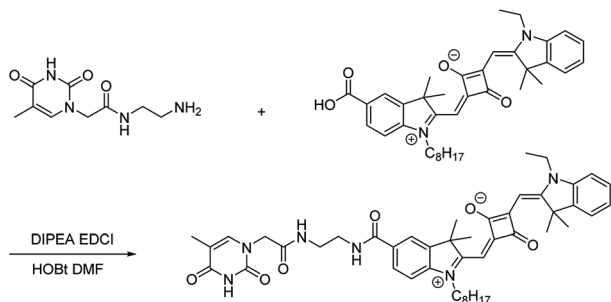
All the reagents related to the starting materials for synthesizing **TSQ** in Scheme 1 and the inorganic salts together with buffer used in the properties study of **TSQ** were obtained from commercial available sources in analytical grade without purification.  $^1\text{H}$  NMR (400 MHz) and  $^{13}\text{C}$  NMR (400 MHz) spectra were recorded on a Bruker AV-400 spectrometer using a deuterated solvent as the lock and TMS as an internal reference. Mass spectrometry analysis was performed on a Q exactive mass spectrometer (Thermo Fisher Scientific, USA). Absorption spectra were measured on Molecular Device Spectrometer 5 (Molecular Devices Corporation, USA).

### 2.2 Synthesis of **TSQ**

The synthetic pathway for **TSQ** was illustrated in Scheme 1. A mixture of A1 (1.351 g, 1.8 mmol) and A2 (0.375 g, 1.8 mmol) were dissolved in *N,N*-dimethylformamide (20 mL) together with amide coupling reagents of *N,N*-diisopropylethylamine (0.225 g, 3.6 mmol), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.342 g, 1.98 mmol) and 1-hydroxybenzotriazole (0.243 g, 1.8 mmol). The mixture was stirred for 4 h at room temperature. It was monitored by thin layer chromatography and purified by column chromatography using DCM : MeOH = 30 : 1 as eluents. The final compound **TSQ** was obtained as dark solid in the yield of 45.2%. The structure was confirmed by NMR and mass spectrometric analysis (Fig. S1 and S2†).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$ : 10.41 (s, 1H), 7.99 (s, 2H), 7.89–7.84 (m, 2H), 7.37–7.30 (m, 2H), 7.19–7.16 (m, 2H), 7.04–7.02 (m, 2H), 6.94–6.92 (d,  $J = 7.9$  Hz), 5.97–5.92 (m, 2H), 4.37 (s, 1H), 4.09–4.07 (d,  $J = 6.6$  Hz, 2H), 3.89 (s, 1H), 3.58 (s, 2H), 3.50 (m, 4H), 1.76–1.70 (m, 16H), 1.35–1.22 (m, 15H), 0.87–0.82 (m, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$ : 181.28, 177.93, 171.20, 168.85, 168.02, 167.93, 164.66, 151.62, 145.46, 142.42, 141.66, 141.09, 129.93, 129.89, 128.54, 128.05, 127.98, 124.43, 122.44, 121.44, 111.07, 109.66, 108.75, 87.37, 86.77, 50.70, 49.72, 48.68, 43.77, 40.29, 38.72, 31.73, 29.70, 29.37, 29.33, 29.17, 27.07, 27.00, 26.80, 22.59, 14.13, 14.09, 12.30, 12.13 ppm. ESI-MS calculated for **TSQ** 788.4261, found 789.5288.

### 2.3 UV-vis spectroscopy

A stock solution of **TSQ** was prepared 10 mM in DMSO. Further dilutions were made to prepare 100  $\mu\text{M}$  of **TSQ** by adding different types of solutions. 11 different metal salts were



Scheme 1 Synthetic pathway for chemosensor **TSQ**.

prepared 10 mM in distilled water and diluted further accordingly. After mixing **TSQ** and metal ion solutions, the absorption measurements were made in 96 well plates on Molecular Device Spectrometer 5 (Molecular Devices Corporation, USA) at the wavelength range of 350 nm to 750 nm.

### 2.4 Job's plot measurements

The stock solution of sensor **TSQ** (10 mM) in DMSO and  $\text{FeCl}_3$  (10 mM) in distilled water were prepared, respectively. When the mole fraction of  $\text{Fe}^{3+}$  was equal to 0.1, the sensor **TSQ** solution (1.8  $\mu\text{L}$ ) and  $\text{FeCl}_3$  solution (0.2  $\mu\text{L}$ ) were added to each 20% AcOH solution to make a total volume of 200  $\mu\text{L}$ . After stirring for a few seconds, UV-vis spectra were recorded at room temperature. When the mole fraction gradually increased to 1.0, **TSQ** volume decreased by 0.2  $\mu\text{L}$  and  $\text{Fe}^{3+}$  volume increased by 0.2  $\mu\text{L}$ , respectively. The absorption spectra were recorded at absorption maximum wavelength. The plots were drawn by plotting  $A_0/(A_0 - A)$  vs.  $1/[\text{Fe}^{3+}]$ , where  $A_0$  equaled to absorption intensity of **TSQ** without  $\text{Fe}^{3+}$ ,  $A$  was corresponding to the absorption intensity of **TSQ** with different concentrations of  $\text{Fe}^{3+}$ .

### 2.5 Competition tests

2  $\mu\text{L}$  of  $\text{Na}^+$ ,  $\text{K}^+$ ,  $\text{Li}^+$ ,  $\text{Ag}^+$ ,  $\text{Zn}^{2+}$ ,  $\text{Hg}^{2+}$ ,  $\text{Cd}^{2+}$ ,  $\text{Fe}^{2+}$ ,  $\text{Co}^{2+}$ ,  $\text{Ca}^{2+}$  ion solutions (10 mM) were extracted individually and mixed with 2  $\mu\text{L}$   $\text{Fe}^{3+}$  (2 mM), 2  $\mu\text{L}$  **TSQ** (10 mM) and filled up with 20% AcOH– $\text{H}_2\text{O}$  solution to total volume of 200  $\mu\text{L}$ . After stirring the solutions for a few seconds, UV-vis spectra were recorded at room temperature.

## 3 Results and discussion

### 3.1 Spectral properties of **TSQ**

The absorption properties of **TSQ** were investigated in different aqueous solutions, *i.e.* distilled pure water, phosphate buffer (PBS buffer, pH = 7.4), borate buffer (pH = 6.8) and acetic acid. The results were recorded as shown in Fig. 1a. Two broad absorption peaks of **TSQ** were observed at 605 nm and 660 nm once the water was present (distilled pure water, phosphate buffer and borate buffer). The sensor has shown a cyan color in the solution. Only one strong and symmetric peak was observed at 635 nm which has been shown as blue color in the presence of pure acetic acid. It is well known that squaraines have great tendency to aggregate in a different pattern which corresponding to the interesting spectroscopic response. A blue shifted absorption and broaden absorption peaks will be observed as in H-aggregation (paralleled oriented arrangement) and red shifted absorption and sharp absorption peaks will be shown as in J-aggregation (head to tail fashion). In this case, it is assumed that the different extent and H-aggregation occur in the presence of water by **TSQ**. To further validate the water induced aggregation effect, the different percentage of water was added into acetic acid to form acetic acid solution. As shown in Fig. 1b, as the percentage of water in acetic acid increased, the absorption performances were almost preserved even in the presence of 60%  $\text{H}_2\text{O}$  in acetic acid solution. With further increased water percentage to 80%, the intensity of new weak band at 650 nm



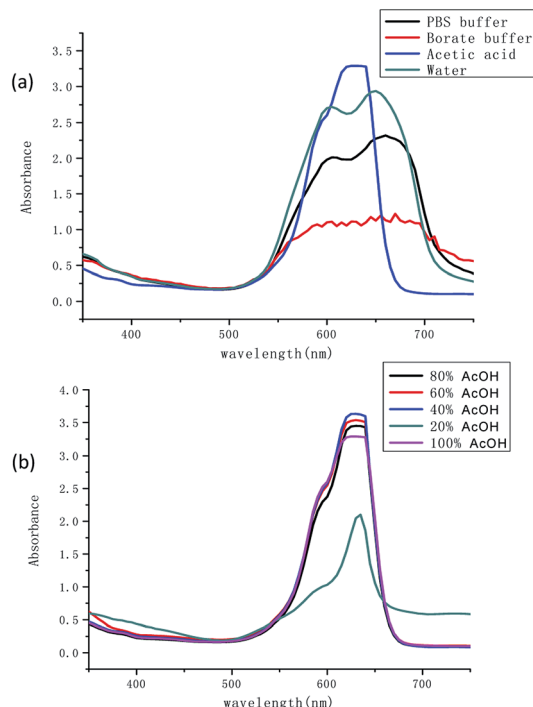


Fig. 1 (a) UV-vis spectra of TSQ (100 μM) in different aqueous solutions and (b) in the mixture of AcOH–H<sub>2</sub>O.

concomitantly increased (blue shift) which was consistent with results in Fig. 1a. For both PBS buffer (pH = 7.4) and borate buffer (pH = 6.8), the pH of the solvent system has no significant influence on the absorption spectrum.

### 3.2 Colorimetric sensing for Fe<sup>3+</sup>

Preliminary results have shown that TSQ could possess good absorption properties in acetic acid, which were selected for identifying the metal ions in the following experiments.

Colorimetric responses to various metal ions including Na<sup>+</sup>, K<sup>+</sup>, Li<sup>+</sup>, Ag<sup>+</sup>, Zn<sup>2+</sup>, Hg<sup>2+</sup>, Cd<sup>2+</sup>, Fe<sup>2+</sup>, Co<sup>2+</sup>, Ca<sup>2+</sup>, Fe<sup>3+</sup> were investigated. It has shown in Fig. 2a that in 20% AcOH–H<sub>2</sub>O solution of 100 μM TSQ, the color change was dramatic and specific to Fe<sup>3+</sup>, while addition of other metal ions did not show any significant change. The Fe<sup>3+</sup> induced a spontaneous color change from blue to green, which could be easily detected by the naked eyes. In the corresponding UV-vis spectrum, the intensity of strong absorption at 635 nm was decreased after the addition of Fe<sup>3+</sup>.

The solvent effects on detection of Fe<sup>3+</sup> for TSQ in AcOH–H<sub>2</sub>O solution were further evaluated (Fig. 3). By varying the portion of AcOH in the distilled water, the decreased absorption change of TSQ only can be observed in 20% AcOH–H<sub>2</sub>O solution. There were no similar observations in other solutions, which indicating that TSQ was a highly sensitive and selective colorimetric sensor for Fe<sup>3+</sup> ions.

### 3.3 Time-resolved colorimetric study

For the purpose of exploring the relationship between absorption intensity and response time, a dynamic study of TSQ in the detection of Fe<sup>3+</sup> was monitored. The green color of TSQ was

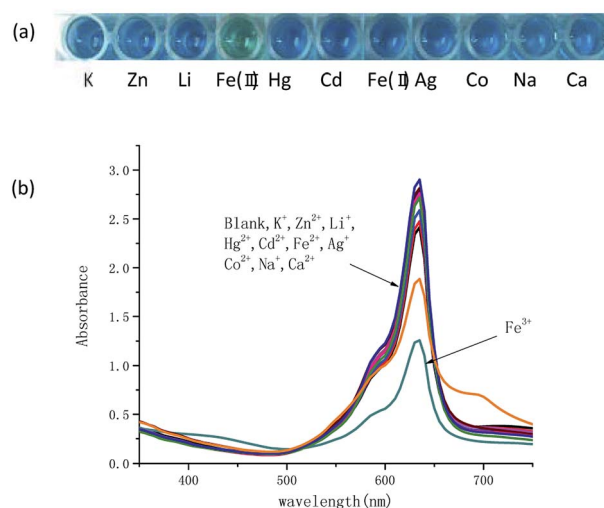


Fig. 2 (a) Visualization and (b) absorption spectra for TSQ solution (100 μM) with/without addition of different metal ions (100 μM) in 20% AcOH–H<sub>2</sub>O.

kept even after 4 hours which clearly indicated that the detection for Fe<sup>3+</sup> was instant and the complexation reaction was stable (Fig. 4).

### 3.4 Binding constant $K_a$ and limit of detection (LOD) for Fe<sup>3+</sup>

To get insight into the binding mode between TSQ and Fe<sup>3+</sup>, the UV-vis absorption spectra of 100 μM TSQ in 20% AcOH–H<sub>2</sub>O solution were recorded during the titration of various concentrations of Fe<sup>3+</sup>. The binding constant ( $K_a$ ) was estimated using a Benesi–Hildebrand plot, which was calculated by absorption changes of consequent titration ( $A_0/A_0 - A$ ) against  $1/[Fe^{3+}]$ . The magnitude of  $K_a$  was calculated from the intercept and slope of the straight line, and the estimated value is about  $9.6 \times 10^5 M^{-1}$  (Fig. 5). More importantly, the absorption change of TSQ corresponded to the concentration of Fe<sup>3+</sup> in a linear manner in the range of 1–100 μM. The high sensitivity of this sensor in 20% AcOH–H<sub>2</sub>O solution allowed detection limit reaching to be 1 μM. The detection limit of Fe<sup>3+</sup> was much lower than the EPA guidelines for drinking water of 5.37 μM.<sup>48</sup> The sensing ability of

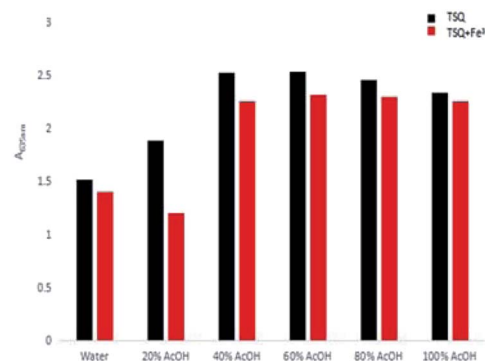


Fig. 3 Bar chart showing the absorption changes of TSQ (100 μM) in the different AcOH–H<sub>2</sub>O solution with adding 100 μM Fe<sup>3+</sup> (red bar) or without Fe<sup>3+</sup> (black).



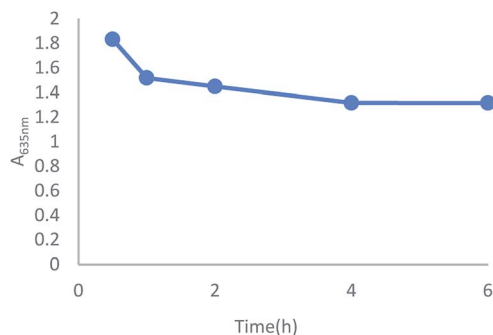


Fig. 4 Dynamic study on the absorption change by mixing TSQ (100  $\mu\text{M}$ ) and  $\text{Fe}^{3+}$  (100  $\mu\text{M}$ ) in 20% AcOH– $\text{H}_2\text{O}$  solution.

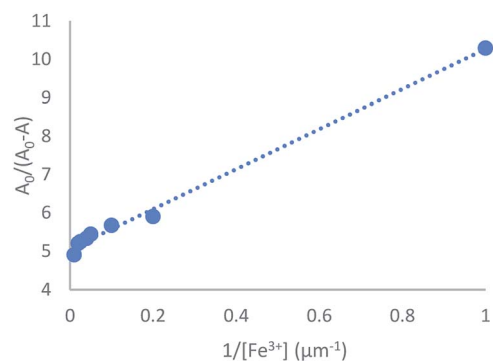


Fig. 5 Benesi–Hildebrand plot analysis of the absorption changes for the complexation between TSQ and  $\text{Fe}^{3+}$ ,  $R^2 = 0.9947$ .

TSQ towards  $\text{Fe}^{3+}$  was also compared to those for coumarin and rhodamine derivatives, which considering as other two types of promising near infrared colorimetric sensing probes. The LOD of present probe TSQ is superior to those of coumarin derivative<sup>49,50</sup> and rhodamine based chemosensors<sup>31</sup> (Table S1†).

### 3.5 Complexation mechanism of TSQ– $\text{Fe}^{3+}$

Job's plot measurement was carried out to explore the binding ratio of TSQ and  $\text{Fe}^{3+}$  (Fig. 6). The plots were done by recording the absorption intensity  $A$  against  $[\text{Fe}^{3+}]/([\text{Fe}^{3+}] + [\text{TSQ}])$ . The results revealed that the TSQ probe bind with  $\text{Fe}^{3+}$  in 1 : 1 stoichiometry, which was further confirmed by ESI-MS data. A solution containing TSQ and 1 equiv. of  $\text{FeCl}_3$  has shown a strong peak at  $m/z$  905.5341, assigned to  $[\text{TSQ} + \text{Fe}^{3+} + \text{CH}_3\text{-COOH} + \text{H}^+]$  ion (Fig. S3†). IR spectra can provide valuable information on the complexation. To gain additional insight into the  $\text{Fe}^{3+}$  binding properties, the FT-IR spectra of TSQ and TSQ– $\text{Fe}^{3+}$  complex were measured (Fig. 7). The shift of the carbonyl absorption band indicated the change of structure. The characteristic amide carbonyl ( $\text{C}=\text{O}$ ) stretching vibrations at  $1685 \text{ cm}^{-1}$  shifted to  $1608 \text{ cm}^{-1}$  in the presence of  $\text{Fe}^{3+}$ , while the characteristic carbonyl ( $\text{C}=\text{O}$ ) stretching frequencies in thymine moiety appeared at  $1584 \text{ cm}^{-1}$  instead of  $1596 \text{ cm}^{-1}$  in the complex of TSQ– $\text{Fe}^{3+}$ . In addition, a new broad peak was generated at  $3400 \text{ cm}^{-1}$  which corresponding to the N–H vibration. All these results strongly indicate the amide carbonyl O atom ( $\text{C}=\text{O}$ ) and the carbonyl O atom in thymine moiety were

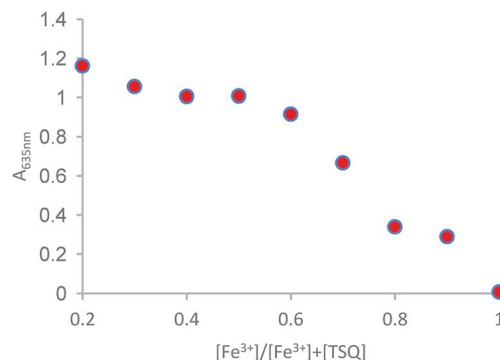


Fig. 6 Job's plot for the complexation of TSQ with  $\text{Fe}^{3+}$  in AcOH– $\text{H}_2\text{O}$  (2 : 8, v/v) solution.

involved in the recognition of  $\text{Fe}^{3+}$ . The density functional theory (DFT) study was further in proceed to get insight into details of TSQ– $\text{Fe}^{3+}$  complexation. All calculations were carried out with the Gaussian 09 package.<sup>51</sup> The density functional theory (DFT) hybrid model with the B3LYP was used for the gas-phase geometry optimizations, Lan12dz basis set with effective core potential (ECP) for Fe, and the 6-31G(d) basis set was used for all remaining atoms. Based on the calculations, the favourability binding mode between TSQ (a) and its  $\text{Fe}^{3+}$  complex (b) was depicted in Fig. 8. It is favourable for the thymine moiety, which was vertical to the planar of asymmetrical squaraine dyes. And the  $\text{Fe}^{3+}$  ion was tended to bind with one carbonyl group in thymine and another carbonyl group on the oxygen atom of amide bond. All conformations were calculated at the same level to confirm their stability (no imaginary frequencies).

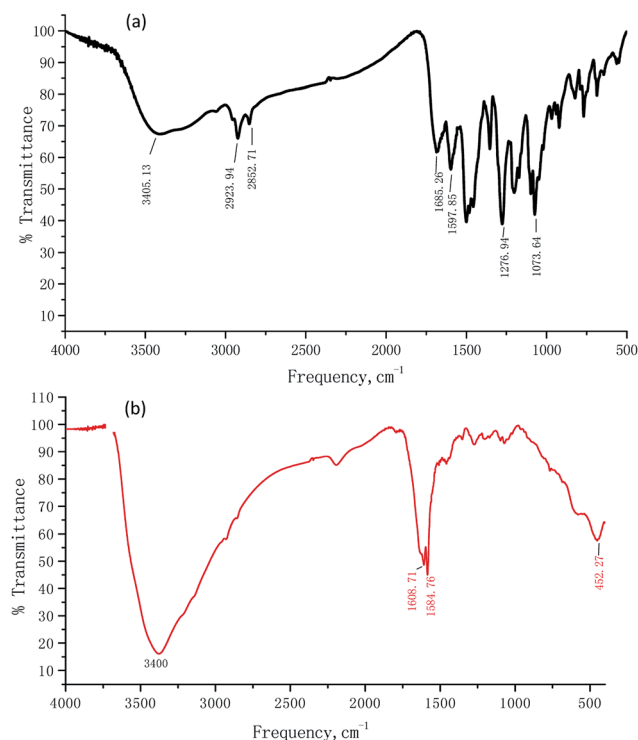


Fig. 7 The IR spectra of TSQ (Black line) and TSQ– $\text{Fe}^{3+}$  (Red line).





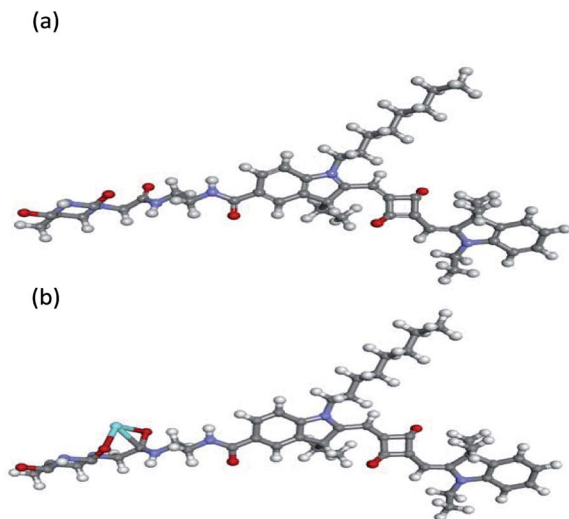


Fig. 8 The geometry-optimized structures of TSQ (a) and its Fe<sup>3+</sup> complex (b) at the SMD (H<sub>2</sub>O)-TD-PBE0/TZVP + LANL2DZ level.

### 3.6 Ions interference effects on the TSQ towards Fe<sup>3+</sup>

Counterion effects on the sensitive colorimetric sensor TSQ for Fe<sup>3+</sup> have also been investigated. The results have shown that the anions with weak coordination abilities towards Fe<sup>3+</sup> such as Cl<sup>-</sup>, SO<sub>4</sub><sup>2-</sup>, SO<sub>3</sub><sup>2-</sup>, NO<sub>3</sub><sup>-</sup>, CO<sub>3</sub><sup>2-</sup> had no predominant effect on the absorption change (Fig. 9). In addition, the colorimetric responses towards other environmentally relevant metal ions were recorded as well. Upon addition of 5 equiv. Na<sup>+</sup>, K<sup>+</sup>, Li<sup>+</sup>, Ag<sup>+</sup>, Zn<sup>2+</sup>, Hg<sup>2+</sup>, Cd<sup>2+</sup>, Fe<sup>2+</sup>, Co<sup>2+</sup>, Ca<sup>2+</sup> in TSQ-Fe<sup>3+</sup> complex solution, no apparent absorption changes were observed, indicating that extra amount of other cations have no significant interference to the response of TSQ towards Fe<sup>3+</sup> (Fig. 10).

### 3.7 Reversible study of TSQ-Fe<sup>3+</sup> complex towards EDTA

In order to test if the proposed TSQ-Fe<sup>3+</sup> complex could be reversed, EDTA as a strong chelator was added to the solution containing of 1 equiv. TSQ and 1 equiv. Fe<sup>3+</sup>. As seen in Fig. 11, the addition of 1 equiv. EDTA restored the absorption signal of TSQ to its original level. Additional 1 equiv. Fe<sup>3+</sup> was added in again to make another round of detection by colorimetric

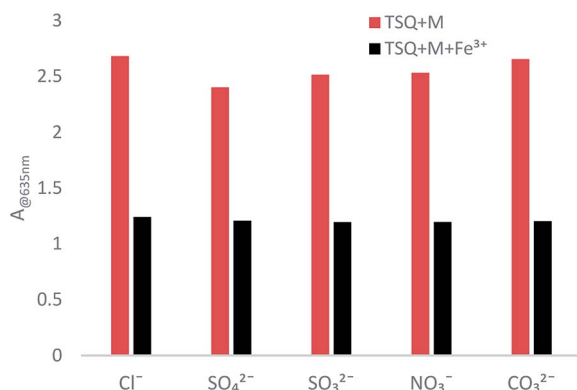


Fig. 9 Absorption changes of TSQ-Fe<sup>3+</sup> in the presence of various test anions in AcOH-H<sub>2</sub>O (2 : 8, v/v).

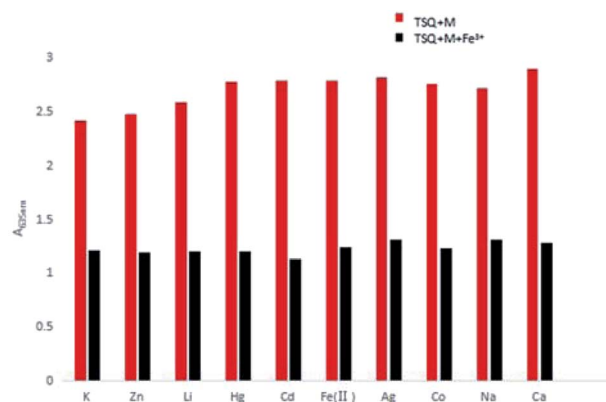


Fig. 10 Absorption changes of TSQ-Fe<sup>3+</sup> in the presence of various test cations in AcOH-H<sub>2</sub>O (2 : 8, v/v).

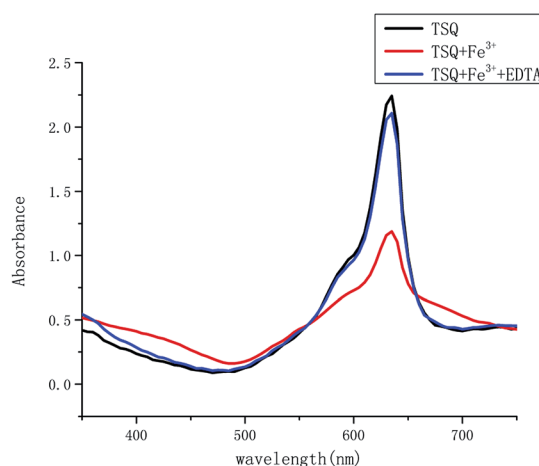


Fig. 11 Absorption spectra of TSQ (100 μM) + Fe<sup>3+</sup> (100 μM) upon the addition of EDTA (1 equiv.) in AcOH-H<sub>2</sub>O (2 : 8, v/v).

change from blue to green (Fig. S4<sup>†</sup>). All these results show that the process of titrating sensor TSQ with Fe<sup>3+</sup> is reversible and that sensor TSQ could be used as an on-off-on switch chemosensor.

### 3.8 Preliminary analytical applications

In order to examine the capability of the proposed chemosensor in environmental samples, TSQ was applied to the determination of the Fe<sup>3+</sup> in waste water from an electroplating factory. As shown in Table 1, the initial Fe<sup>3+</sup> concentrations were measured by the atomic absorption spectrometry (AAS) method and three levels of standard concentrations of Fe<sup>3+</sup> were added into real

Table 1 Determination of Fe<sup>3+</sup> ion in waste water samples with TSQ

Sample	AAS (μM)	Added (mM)	Found (mM)	Recovery (%)
Waste water 1	2.08	1.20	1.22	101.2
Waste water 2	3.78	2.31	2.38	103.3
Waste water 3	6.02	3.62	3.58	99.0



samples. By applying our **TSQ** test, the content of  $\text{Fe}^{3+}$  in waste water was in good agreement with that added amount by a relative error in less than 5%. Therefore, the present **TSQ** was initially proven to be well used in the determination of  $\text{Fe}^{3+}$  in real samples.

## 4 Conclusions

In summary, we have successfully constructed a thymine-squaraine based colorimetric chemosensor for  $\text{Fe}^{3+}$  detection in aqueous solution. It has shown an excellent selectivity for  $\text{Fe}^{3+}$  over the other cations (blue to green). The limited of detection ( $1 \mu\text{M}$ ) was determined lower than the EPA guidelines for drinking water of  $5.37 \mu\text{M}$  and no other ion interferences have shown significant effects on the  $\text{Fe}^{3+}$  detection. Further study of mechanism by Job's plot measurement, ESI-MS indicated that 1 : 1 complex was formed between **TSQ** and  $\text{Fe}^{3+}$ . IR spectrum and DFT calculation further revealed the detailed binding mode of **TSQ**- $\text{Fe}^{3+}$  complex in the way of binding on the amide carbonyl O atom (C=O) and the carbonyl O atom on thymine moiety towards  $\text{Fe}^{3+}$ . Reversible study and preliminary analytical application study have shown strong potent of **TSQ** as a good chemosensor to  $\text{Fe}^{3+}$  in real sample applications. Considering that the developed thymine-squaraine based chemosensor **TSQ** was quite effective and selective, we expect that it will potentially serve as colorimetric probe for more  $\text{Fe}^{3+}$ -related environmental studies.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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