A highly selective colorimetric fluorescent probe for detection of Hg$^{2+}$ and its application on test strips

Geng Yang, Xia Meng, Shimin Fang, Hongdong Duan, Lizhen Wang and Zhenzheng Wang

An efficient fluorescent probe Pyr-Rhy based on pyrazole was developed, which can detect Hg$^{2+}$ in water. Its fluorescence properties were studied by UV-vis and fluorescence spectroscopy, and the study results indicated that this probe can selectively detect Hg$^{2+}$ via complexation reaction, and then cause a remarkable color change from colorless to pink and a strong fluorescence enhancement can be observed. Furthermore, this probe showed high sensitivity with the detection limit down to $2.07 \times 10^{-8}$ M, and its stoichiometric ratio toward Hg$^{2+}$ ions was 1:1. The sensing mechanism was investigated by Job’s plot $^1$H NMR titrations, and FT-IR spectra analysis, which demonstrated a chelation-enhanced fluorescence (CHEF) mechanism. More importantly, obvious color changes of sensor Pyr-Rhy can be observed when it was impregnated on filter paper testing strips and immersed in Hg$^{2+}$ solution (water as solution), indicating its potential application for trace Hg$^{2+}$ detection in environmental samples.

1. Introduction

Mercury (Hg), one of the most toxic pollutants,$^1$ is widely distributed in the environment and exists in the state of mercury metal, inorganic mercury and organic mercury compounds.$^2$ The organic mercury is the most harmful of the mercury-containing compounds, because it is hardly biodegradable and can be biomagnified in the food chain.$^4$ Even at very low levels, organic mercury can seriously destroy the activity of proteins and enzymes, thereby causing various diseases, including brain damage, movement disorders, cognitive dysfunction and kidney failure.$^5$ There are many previous studies for the detection of mercury ions with different methods, and we have compared the present method with the others, and a table was added (Table 1). While other methods can detect Hg$^{2+}$, they can’t be reused; and the synthesis methods of other methods are cumbersome and not easy to operate. Therefore, it is necessary to develop a sensitive, fast and low-cost method for detection of Hg$^{2+}$.

In recent years, there have been many ways to detect cations and anions such as ratiometric fluorescent paper sensor,$^1$ silica-based SERS chip.$^{11,14}$ Schiff base fluorescent probes have been drawn great attention and widely used in detection of heavy metal ions.$^15$ The C=N structure in Schiff base has strong chelating ability, and more importantly, these chelating processes always produce strong fluorescence release.$^16,17$ Pyrazole derivatives are interesting heterocyclic compounds bearing five membered heterocycles and useful raw materials to synthesize various biological molecules. Due to their strong coordination ability with metal ions, pyrazole based derivatives have been widely used for the development of fluorescent probes to detect heavy metal ions, such as Zn$^{2+}$, Cd$^{2+}$, and Hg$^{2+}$.$^{18-20}$ Furthermore, pyrazole derivatives are always employed as important capping motifs in probing molecules, which can be used as ligands for selective recognition metal ions in coordination chemistry. Therefore, pyrazoles are attracting building blocks for designing and synthesizing a series of heterocyclic compounds, such as optical brighteners, and UV stabilizers, and fluorescent probes.$^{21}$

In this paper, a novel fluorescent probe Pyr-Rhy was described, which was synthesized by modification of rhodamine with a pyrazole derivative (Fig. 1). This probe exhibited high fluorescent “turn-on” response toward Hg$^{2+}$ in aqueous DMSO solution, and this process was accompanied with a sensitive fluorescence color change that could not be interfered by other coexisting metal ions. Test results showed that the recognition process of sensor Pyr-Rhy toward Hg$^{2+}$ was completed instantly, and the detection limit was found to be as low as $2.07 \times 10^{-8}$ M. Therefore, sensor Pyr-Rhy could be used for quantitative estimation of Hg$^{2+}$ in aqueous media.
2. Materials and methods

2.1. General information

In this experiment unless otherwise noted, all reagents and materials were derived from commercial, which without further refinement. Pyrazole, p-fluorobenzaldehyde, rhodamine 6G, hydrazine hydrate and glacial acetic acid were purchased from Aladdin. Distilled water was produced in our lab. Different metal ion solutions were prepared with the corresponding inorganic salts. Nuclear magnetic spectrometer is Bruker AV-400 spectrometer; infrared spectrometer is Bruker ALPHA FT-IR spectrometer; fluorospectrophotometer is Hitachi F-4600 and scanning speed was 2400 nm/min.

2.2. Synthesis

2.2.1. Synthesis of compound 1. Pyrazole (1.36 g, 20 mmol), p-fluorobenzaldehyde (2.48 g, 20 mmol) and anhydrous potassium carbonate (6.90 g, 50 mmol) were dissolved in 50 mL of DMF, the mixture was allowed to warm to 110 °C and stirred at this ambient temperature for 24 h. After TLC showed the conversion of the starting material, the mixture was cooled to room temperature, filtered. The filtrate was diluted with 400 mL of distilled water and the mixture was extracted with 100 mL of acetic ether, the organic layer was dried with anhydrous sodium sulfate and then evaporated, the resultant residue was purified by a column chromatography (petroleum/ethyl acetate, 7/1) to give compound 1 was a white powder (3.34 g, yield 91%); mp 82–84 °C; 1H NMR (400 MHz, DMSO-d$_6$) δ 10.00 (s, 1H), 8.68 (d, J = 2.5 Hz, 1H), 8.06 (dd, J = 8.7 Hz, 4H), 7.85 (t, J = 6.4 Hz, 1H), 6.67–6.58 (m, 1H); 13C NMR: (100 MHz, DMSO-d$_6$) δ 191.87, 143.67, 142.21, 133.65, 131.15, 128.44, 118.27, 108.88 ppm; ESI-TOF HRMS (m/z): calcd for C$_{10}$H$_8$N$_2$O$^+$, [M+H$^+$], 172.0729; found, 172.0731.

2.2.2. Synthesis of compound 3. To the mixture of rhodamine 6G 2 (9.85 g, 20 mmol) in 300 mL of ethanol was added anhydrous potassium carbonate (6.90 g, 50 mmol) and stirred for 24 h. After TLC showed the conversion of the starting material, the mixture was cooled to room temperature, filtered. The filtrate was diluted with 400 mL of distilled water and the mixture was extracted with 100 mL of acetic ether, the organic layer was dried with anhydrous sodium sulfate and then evaporated, the resultant residue was purified by a column chromatography (petroleum/ethyl acetate, 7/1) to give compound 3 was a white powder (3.34 g, yield 91%); mp 82–84 °C; 1H NMR (400 MHz, DMSO-d$_6$) δ 10.00 (s, 1H), 8.68 (d, J = 2.5 Hz, 1H), 8.06 (dd, J = 8.7 Hz, 4H), 7.85 (t, J = 6.4 Hz, 1H), 6.67–6.58 (m, 1H); 13C NMR: (100 MHz, DMSO-d$_6$) δ 191.87, 143.67, 142.21, 133.65, 131.15, 128.44, 118.27, 108.88 ppm; ESI-TOF HRMS (m/z): calcd for C$_{10}$H$_8$N$_2$O$^+$, [M+H$^+$], 172.0729; found, 172.0731.

Table 1. Compared the present method with the others

<table>
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<tr>
<th>Structural of compound</th>
<th>Molecular formula</th>
<th>Solvent</th>
<th>Repeatability</th>
<th>Reference</th>
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<tr>
<td>C$<em>{18}$H$</em>{22}$N$_4$S</td>
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<tr>
<td>C$<em>{27}$H$</em>{29}$N$_3$OSR</td>
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<tr>
<td>C$<em>{18}$H$</em>{43}$N$_5$O$_3$S</td>
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<td>10</td>
<td></td>
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<tr>
<td>C$<em>{13}$H$</em>{12}$N$_2$O$_2$S</td>
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<td>11</td>
<td></td>
</tr>
<tr>
<td>C$<em>{19}$H$</em>{34}$N$_6$O$_2$</td>
<td>DMSO/H$_2$O (1/1, v/v)</td>
<td>Yes</td>
<td>This work</td>
<td></td>
</tr>
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</table>

Fig. 1. Synthesis of pyrazole-based fluorescent probe Pyr-Rhy.
excess of hydrazine hydrate (0.95 g, 20 mmol), the mixture was heated to 80 °C and refluxed for 12 h to give a dark red clear transparent solution. Then, the mixture was subjected to decompression and evaporation till a few solid was formed. After which, the mixture was poured into 200 mL of deionized water, rested for 3 hours, and filtered. The solid was washed alternately with distilled water and ethanol several times, and dried in vacuo to afford compound 3 as a pink solid (7.88 g, yield 70%). 1H NMR (400 MHz, DMSO-d$_6$) δ 7.79–7.71 (m, 1H), 7.50–7.41 (m, 2H), 6.98–6.87 (m, 1H), 6.26 (s, 2H), 6.10 (s, 2H), 4.99 (t, J = 5.3 Hz, 2H), 4.21 (s, 2H), 3.32 (s, 1H), 1.88 (d, J = 9.9 Hz, 6H), 1.21 (t, J = 7.0 Hz, 6H); 13C NMR (100 MHz, DMSO-d$_6$) δ 165.19, 152.05, 151.33, 147.34, 132.27, 129.47, 127.98, 126.98, 123.40, 122.11, 117.77, 105.00, 95.86, 64.95, 37.44, 17.01, 14.19 ppm; ESI-TOF HRMS (m/z): calcd for C$_{28}$H$_{34}$N$_6$O$_2$, [M + H]$^+$, 534.66; found, 534.68.

2.2.3 Synthesis of sensor Pyr-Rhy. Compound 3 (1.72 g, 10 mmol) and compound 1 (4.56 g, 10 mmol) were dissolved in 40 mL of ethanol, the mixture was heated to 80 °C and refluxed for 2 h after a catalytic amount of acetic acid (3.46 mL, 10 mmol) was added. Then the mixture was cooled to room temperature to yield a rose pink precipitate. The mixture was filtered, the solid was washed with ethanol for several times and dried in vacuo to give sensor Pyr-Rhy (4.98 g, yield 79.3%); 1H NMR (400 MHz, DMSO-d$_6$) δ 8.60 (s, 1H), 8.36 (d, J = 2.4 Hz, 1H), 7.78 (d, J = 7.0 Hz, 1H), 7.70 (d, J = 8.6 Hz, 2H), 7.61 (d, J = 1.3 Hz, 1H), 7.52–7.73 (m, 4H), 6.92 (d, J = 7.3 Hz, 1H), 6.44–6.37 (m, 1H), 6.21 (s, 2H), 6.05 (s, 2H), 4.94 (t, J = 5.3 Hz, 2H), 4.22 (t, J = 5.1 Hz, 1H), 1.07 (t, J = 7.1 Hz, 6H), 0.92 (t, J = 7.0 Hz, 4H); 13C NMR: (100 MHz, DMSO-d$_6$) δ 164.36–164.05, 152.00–151.71, 151.48, 148.30, 146.45–146.22, 142.08–141.78, 141.17–140.81, 134.52–134.16, 132.50, 129.50–129.00, 128.29, 127.35, 124.49–124.07, 123.62–122.94, 118.82, 108.75, 105.45, 96.22, 66.05, 37.95, 19.18–18.89, 17.48, 14.64 ppm; ESI-TOF HRMS (m/z): calcd for C$_{28}$H$_{34}$N$_6$O$_2$, [M + H]$^+$, 534.66; found, 534.68.

3. Results and discussion

3.1. UV-vis absorption spectral studies of sensor Pyr-Rhy

The UV-vis absorption spectra of sensor Pyr-Rhy toward different metal ions (Ag$^+$, Al$^{3+}$, Cd$^{2+}$, Co$^{3+}$, Cr$^{3+}$, Cu$^{2+}$, Hg$^+$, Mg$^{2+}$, Ni$^{2+}$, and Zn$^{2+}$) were studied through using UV-vis spectroscopy. As shown in Fig. 2, free sensor solution in DMSO/H$_2$O (1/1, v/v) showed one distinct absorption band at 320 nm. However, with the addition of Hg$^+$ (10 μM, water as solution) to the above solution, a new the observed absorption band at 320 nm decreased obviously. Whereas, with the addition of other metal cations (Ag$^+$, Al$^{3+}$, Cd$^{2+}$, Co$^{3+}$, Cr$^{3+}$, Cu$^{2+}$, Mg$^{2+}$, Ni$^{2+}$ and Zn$^{2+}$, water as solution) showed no significant decrease, which indicated the complexion of sensor Pyr-Rhy with Hg$^{2+}$. These test results displayed that the UV-vis response of sensor Pyr-Rhy has highly specific selection for Hg$^{2+}$.

3.2. Fluorescence study of sensor Pyr-Rhy

The fluorescence emission behavior of sensor Pyr-Rhy toward different metal ions was investigated in DMSO/H$_2$O (1/1, v/v) solution, and the test results can be seen from Fig. 3. Clearly, free sensor Pyr-Rhy exhibited one fluorescence emission band at about 575 nm with an excitation at 525 nm. But after the addition of Hg$^{2+}$ (1.0 equiv.), a significant fluorescence enhancement can be observed with the emission maximum at 575 nm. In contrast, addition of other metal ions exhibited no significant fluorescence enhancement. This fluorescence response sensor Pyr-Rhy to Hg$^{2+}$ might be ascribed to the blocked photoinduced electron transfer (PET) process and the chelation-enhanced fluorescence (CHEF) behavior. Before adding metal ions, the isomerization of C=N double bond of sensor Pyr-Rhy exhibited weak fluorescence emission. While, addition of metal ions led to a new π-conjugated structure, which was accompanied with a selective chelation-enhanced fluorescence (CHEF) effect.

The specificity of sensor Pyr-Rhy toward Hg$^{2+}$ was investigated by colorimetric experiments. As we can see from Fig. 4a, addition of Hg$^{2+}$ to the solution of sensor Pyr-Rhy in DMSO/H$_2$O solution resulted in a rapid color change from dark green to deep red under visible light illumination. Further, the solution was subjected to UV–vis spectroscopic analysis, as shown in Fig. 4b, the absorption peaks at 320 nm and 575 nm were observed, which indicated the presence of the complexation of sensor Pyr-Rhy with Hg$^{2+}$. The fluorescence emission spectrum (Fig. 4c) was also recorded, the emission intensity showed a significant increase at 575 nm with a slight shoulder at 525 nm, and a new emission band appeared. These results suggested that the fluorescence enhancement was due to the chelation of both the metal ion and the nitrogen double bond of the sensor Pyr-Rhy.
(10 μmol L\(^{-1}\), 1/99, v/v) resulted in a color change from colorless to pink in a few seconds, which suggested that sensor Pyr-Rhy could be used for naked eye recognition of Hg\(^{2+}\) with concentration as low as 10 μmol L\(^{-1}\). In addition, the fluorescence color change from blond to bright yellow was clearly observed under 365 nm UV light after addition of Hg\(^{2+}\). Addition of other metal cations showed no distinct influence on the fluorescence color (Fig. 4b). These outstanding findings demonstrated that

Fig. 4  (a) Colorimetric performance of sensor Pyr-Rhy (10 μM) upon addition of different metal ions (1.0 equiv.) in DMSO/H\(_2\)O (1/1, v/v) solution; (b) color change induced upon addition of Hg\(^{2+}\) under 365 nm UV lamp.

Fig. 5  (a) The fluorescence spectra of sensor Pyr-Rhy (10 μM) with the increasing concentration of Hg\(^{2+}\) ion (0–2.0 equiv.) in ethanol solution; (b) the linear fit between sensor Pyr-Rhy and Hg\(^{2+}\) ion.

Fig. 6  Effect of pH on the fluorescence intensity of sensor Pyr-Rhy toward Hg\(^{2+}\) in DMSO/H\(_2\)O (1/1, v/v) solution.

Fig. 7  The black bars represent the fluorescent intensity of sensor Pyr-Rhy and various metal ions (1.0 equiv.) in DMSO/H\(_2\)O (1/1, v/v) solution; the red bars represent the fluorescent intensity of sensor Pyr-Rhy and Hg\(^{2+}\) (1.0 equiv.) in DMSO/H\(_2\)O (1/1, v/v) solution.
sensor Pyr-Rhy could be used to detect Hg\(^{2+}\) qualitatively and quantitatively by changing the color and doubling the spectral signals.

To explore the sensitivity of sensor Pyr-Rhy toward Hg\(^{2+}\), the fluorescence titration experiments were carried out and the results were shown in Fig. 5a. The fluorescence intensity of sensor Pyr-Rhy was gradually enhanced with the increasing of Hg\(^{2+}\) concentration. It was clearly found that the saturation state was reached in presence of 1.0 equiv. of Hg\(^{2+}\), which proved the 1 : 1 binding stoichiometry of sensor Pyr-Rhy with Hg\(^{2+}\). This fluorescent behavior was ascribed to the coordination of sensor Pyr-Rhy with Hg\(^{2+}\), which decreased the PET process and C=N isomerization. Moreover, there was an excellent linear relationship between the fluorescence intensity and the concentration of Hg\(^{2+}\) \((R^2 = 0.99163)\) (Fig. 5b). Based on this, the detection limits of sensor Pyr-Rhy with Hg\(^{2+}\) was calculated to be as low as \(2.07 \times 10^{-8}\) M (DL = 3Sb/S).

3.3. pH effects

The effect of pH on fluorescence response of sensor Pyr-Rhy toward Hg\(^{2+}\) was determined over a wide range of pH (1.0–14.0). In this test, hydrochloric acid and sodium hydroxide were used to adjust the pH values. As shown in Fig. 6, the fluorescence intensity of sensor Pyr-Rhy increased with the increasing of pH value (pH \(\leq 4.0\)). And with the increasing of pH (pH = 5–12), the fluorescence intensity was observed to be held steady. These results proved that this probe has potential application value to test Hg\(^{2+}\) within a biological scale of pH values.

3.4. Competition experiment and reversibility study

High selectivity toward Hg\(^{2+}\) over the other metal ions is an important parameter, which is helpful to study the performance of sensor Pyr-Rhy. Therefore, the competition experiments were carried out in the presence of Hg\(^{2+}\) (1.0 equiv.) mixed with other metal ions (1.0 equiv.) in aqueous DMSO media (DMSO/H\(_2\)O, 1/1, v/v), and the result was shown in Fig. 7. The selectivity of sensor Pyr-Rhy toward Hg\(^{2+}\) ions over other competitive cations was remarkably high, illustrating that sensor Pyr-Rhy was capable of fluorescent recognition of Hg\(^{2+}\) with high selectivity and sensitivity in aqueous DMSO solution (DMSO/H\(_2\)O, 1/1, v/v).

Reversibility study was carried out to explored the reuse property of sensor Pyr-Rhy toward Hg\(^{2+}\). Reversibility is a key prerequisite in designing and developing novel chemosensors for practical applications. The reversibility of the recognition process of sensor Pyr-Rhy was performed by adding a strong chelating agent EDA (Fig. 8).\(^{31-32}\) The addition of excessive EDA to a mixture of sensor Pyr-Rhy and Hg\(^{2+}\) in ethanol solution...
resulted in diminution of the fluorescence intensity, due to demetalation of \( \text{Hg}^{2+} \) from the corresponding complex. Meanwhile, the solution color was changed from fluorescent yellow to colorless, which indicated the regeneration of chemosensor \text{Pyr-Rhy}. However, when \( \text{Hg}^{2+} \) continued to be added to the mixed solution, a strong fluorescence enhancement was observed and the solution color was changed from colorless to fluorescent yellow, which demonstrated the excellent reversible property of sensor \text{Pyr-Rhy}.

3.5. Contact mode detection between sensor Pyr-Rhy and Hg\(^{2+}\)

The recognition of sensor \text{Pyr-Rhy} toward \( \text{Hg}^{2+} \) was tested on pretreated filter paper to explore its versatility application as a simple and efficient solid-state probe. The test strips were prepared by immersing the filter paper into a saturated solution of sensor \text{Pyr-Rhy} in DMSO for a few seconds. Then, these strips dried in air and subsequently treated with \( \text{Hg}^{2+} \) solution (1.0 \( \times 10^{-2} \) mol L\(^{-1}\)) in water for solid state experiments. The filter

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Fig. 10 (a) The FT-IR spectrum of sensor \text{Pyr-Rhy}; (b) binding FT-IR spectrum of \text{Pyr-Rhy–Hg}^{2+}.

Fig. 11 \(^1\text{H NMR spectra of sensor Pyr-Rhy and complex Pyr-Rhy–Hg}^{2+}\) in DMSO–\(d_6\).
3.6. Proposed binding mode of sensor Pyr-Rhy with Hg$^{2+}$ ion

The binding mode of sensor Pyr-Rhy toward Hg$^{2+}$ was investigated through IR spectra (Fig. 10). Free sensor Pyr-Rhy showed stretching vibration bands assignable to -OH and CH=N at 3422 and 1674 cm$^{-1}$, respectively (Fig. 10a). While complexation of sensor Pyr-Rhy with Hg$^{2+}$ resulted in the obvious shift of -OH stretching vibration band from 3422 cm$^{-1}$ to 3432 cm$^{-1}$ (Fig. 10b). Furthermore, the characteristic stretching vibration band of CH=N and phenolic O-H changed obviously, which indicated that the aldimine nitrogen atom of sensor Pyr-Rhy was involved in the chelating with Hg$^{2+}$. This result was further confirmed by the $^1$H NMR spectra recorded in DMSO-$d_6$ as shown in Fig. 11. The H proton signals of phenolic OH and aldimine CH=N in free sensor Pyr-Rhy were located at δ 8.74 and 8.50 ppm, respectively. However, upon addition of Hg$^{2+}$ (1.0 equiv.), these two signals shifted to lower δ values (8.59 ppm for OH, and 8.36 ppm for CH=N). These data strongly supported the previous conclusion that complexation reaction of sensor Pyr-Rhy with Hg$^{2+}$ occurred at the CH=N position, and thereby leading to the obvious migration of intramolecular electrons and H proton signals of CH=N and OH.

In addition, the Job’s plot of sensor Pyr-Rhy with Hg$^{2+}$ in DMSO/H$_2$O (1/1, v/v) solution was also measured. As depicted in Fig. 12, the maximum emission intensity was measured for a molar fraction of 0.5, which gave a solid evidence for the formation of a 1 : 1 complex of sensor Pyr-Rhy with Hg$^{2+}$. Furthermore, the $R$ value also proved the 1 : 1 stoichiometric ratio for the formation of sensor Pyr-Rhy–Hg$^{2+}$ complex. The 1 : 1 binding stoichiometry was further confirmed by Benesi-Hildebrand method, the plot of 1/(F – $F_0$) against 1/[Hg$^{2+}$] showed an excellent linearity ($R^2 = 0.99163$) (Fig. 5b), which strongly supported the 1 : 1 binding stoichiometry of the sensor Pyr-Rhy and Hg$^{2+}$. The proposed binding mechanism of sensor Pyr-Rhy with metal ions is shown in Fig. 13. Free sensor Pyr-Rhy contains two equilibrium conformations owing to the isomerization of C=N bond, which can lead to weak fluorescence emission. After adding specific metal ions, the isomerization of C=N double bond is severely restrained due to the block of photoinduced electron transfer (PET). This process generates a new π-conjugated structure, which can result in the increase of structure rigidity and the enhancement of fluorescence intensity.

4. Conclusions

In summary, a novel efficient fluorescent probe Pyr-Rhy for naked-eye detection of Hg$^{2+}$ had been successfully synthesized, using the commercial available pyrazole and rhodamine 6G as the starting materials. This probe exhibits high anti-interference performance, rapid response, high sensitivity and excellent selectivity toward Hg$^{2+}$ over other coexisted metal ions within wide pH range. The filter paper test strips demonstrate that probe Pyr-Rhy has high value in the practical applications.
Importantly, the probe can be used to trace amounts of hazardous Hg\(^{2+}\) ions in water with good recoveries and less relative standard deviations, indicating that the developed probe has good accuracy and precision for the analysis of trace Hg\(^{2+}\) in practical samples.

Conflicts of interest

The authors declare no competing interests.

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References