Sensors & Diagnostics

PAPER

Cite this: Sens. Diagn., 2024, ³, 1062

An imidazo[1,2-a]pyridine-functionalized xanthene fluorescent probe for naked-eye detection of Hg^{2+} and its application in cell imaging and test strips†

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An imidazo[1,2-a]pyridine-functionalized xanthene dye (Rh-Ip) was designed and developed as a fluorescent probe (Rh-Ip-Hy) for Hg^{2+} by introducing spirolactam to the molecule. The probe Rh-Ip-Hy with a ring-closed spirolactam structure reacts with Hg^{2+} to form a fluorescent ring-opened spirolactone structure. Due to its asymmetric structure and the presence of a Lewis base site, the probe exhibits a larger Stokes shift and higher pH tolerance than the traditional xanthene dyes and probes. Furthermore, the probe is highly selective to Hg^{2+} within a wide pH range of 5.0–11.0. The probe Rh-Ip-Hy also exhibits low cytotoxicity and can be used for the detection of Hq^{2+} in living HeLa cells through fluorescence imaging. Finally, a paper-based test strip was prepared and successfully applied for the detection of Hg²⁺ in tap water and lake water samples. PAPER
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Received 23rd March 2024, Accepted 27th April 2024

DOI: 10.1039/d4sd00090k

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Introduction

Mercury is a highly toxic pollutant that poses a significant threat to the environment. Mercury, even when present in low concentrations in the environment, can accumulate through the food chain, $1,2$ leading to a variety of diseases, including neurological damage³ and serious cognitive and motion disorders.4 Therefore, it is important to develop simple tools for the sensitive and selective detection of Hg^{2+} in various water samples.

Over the past decades, several methods such as highperformance liquid chromatography, atomic emission spectroscopy, surface-enhanced Raman spectroscopy, and inductively coupled plasma mass spectroscopy have been developed for detecting Hg^{2+} in environmental samples with high sensitivity, selectivity, and precision.^{5,6} However, most of these methods are time-consuming or need tedious sample preparation steps, for which reason they are not suitable for on-site analysis and in vivo imaging.⁷ Fluorescent probe-based detection methods have been widely used for detecting various analytes both *in vitro* and *in vivo* because of

their advantages such as high selectivity and sensitivity, noninvasiveness and high spatiotemporal resolution. $8-14$

Due to their advantages including high molar absorption coefficient, high fluorescence quantum yield, and good photostability, xanthene-based fluorophores including rhodamines and fluoresceins have been widely used for designing fluorescent probes by following a target-induced ring-opening strategy enabled by the unique property of spirolactam. $15-17$ Initially, these probes are in the spirolactam form, which is nonfluorescent and colourless. However, in the presence of analytes, strong fluorescence and significant colour changes are observed because of the analyte-triggered ring-opening of probes. Although various xanthene-based probes have been developed for detecting Hg^{2+} with high selectivity and sensitivity, there are still some limitations to these probes. For example, classic rhodamine and fluorescein dyes have a small Stokes shift of 20-40 nm,^{18,19} which is susceptible to being affected by self-absorption and background fluorescence. Furthermore, some reported spirolactam-based probes are vulnerable to H^+ (see Table S1; ESI†), causing undesired background signals, which may impede their applications to practical complex samples and in vivo imaging (Scheme 1).^{20,21} These concerns encouraged us to develop a novel fluorescent probe for detecting Hg^{2+} in various aquatic environments.

In the present manuscript, an imidazo $[1,2-a]$ pyridine group is grafted to a xanthene plane to form a probe with an asymmetrical structure and a Lewis base site. The aim is to increase the Stokes shifts and improve the pH tolerance. As expected, the Hg²⁺ fluorescent probe $(Rh-Ip-Hy)$ exhibits a

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[†] Electronic supplementary information (ESI) available: Copies of ¹H/¹³C NMR and HRMS spectra for Rh-Ip and Rh-Ip-Hy, the limit of detection, HRMS spectra of sensing mechanism are included. See DOI: [https://doi.org/10.1039/](https://doi.org/10.1039/d4sd00090k) [d4sd00090k](https://doi.org/10.1039/d4sd00090k)

Scheme 1 (A) Structures of traditional xanthenes. (B) Structures and color changes of Rh-Ip-Hy (5 μ M) before and after exposure to Hg²⁺ (50 μM).

large Stokes shift (75 nm) in the presence of Hg^{2+} and high pH tolerance within the pH range of 1–14. Furthermore, the proposed probe Rh-Ip-Hy is highly selective and sensitive to Hg^{2+} within a wide pH range (5.0–11.0), and can be utilized for the quantitative detection of Hg^{2+} in living HeLa cells.

Experimental

Materials and instrumentation

All chemicals and reagents were purchased from Innochem Company (Beijing, China) and used without further treatment unless noted. Test solutions of $\text{Zn}(\text{NO}_3)_2, \text{Cu}(\text{NO}_3)_2,$ AgNO₃, Al(NO₃)₃.9H₂O, NaCl, FeCl₂, FeCl₃, KCl, LiCl, $CaCl₂·2H₂O$, $BaCl₂·2H₂O$, $MgSO₄$, $Ni(ClO₄)₂$, $Mn(ClO₄)₂$, $Cd(CIO₄)₂$, $Co(CIO₄)₂$, and $Pb(CH₃COO)₂$ were prepared by dissolving or diluting these compounds in double-distilled water, followed by further dilution if necessary.

Instrumentation and spectroscopic studies

Absorption spectra were recorded on an Agilent Cary 60 spectrophotometer. Fluorescence spectroscopy was conducted on a Hitachi F-4600 spectrometer. 1 H NMR (400 MHz) and 13 C NMR (100 MHz) spectra were recorded on a Bruker AVANCE III 400 MHz spectrometer with chemical shifts reported in ppm (TMS as the internal standard). High-resolution mass spectrometry (HRMS) was performed with an Agilent 6510 Q-TOF LC/MS instrument (Agilent Technologies, Palo Alto, USA) that was equipped with an electrospray ionization (ESI) source. The pH measurements were carried out using a Leici PHS-3C meter. Cells were imaged using a TCS SP8 II confocal laser scanning microscope (Nikon C2plus, Germany).

Sample preparation

The stock solution of **Rh-Ip-Hy** (10 mM) was prepared in N,Ndimethylformamide (DMF). The test solution was set at a concentration of 5 μM by diluting the stock solution with $H₂O/EtOH$ (v/v; 4/1). Stock solutions of various metal ions (10.0 mM) were prepared in distilled water and were diluted with $H₂O/EtOH$ (v/v; 4/1) to the required concentration for specific experiments. The emission spectra were recorded 30 min after the addition of each analyte.

Synthesis of the dye Rh-Ip

A mixture of compound 1 (ref. 22) $(1.2 \text{ g}, 5 \text{ mmol})$ and 2- $[4-$ (diethylamino)-2-hydroxybenzoyl]benzoic acid (1.4 g, 4.47 mmol) was dissolved in CF_3COOH/CH_3SO_3H (20 mL, $v/v = 1/1$). The resulting solution was heated at 80 \degree C for 24 h. After cooling, the mixture was poured into $H₂O$ (80 mL), neutralized using NaHCO₃, and extracted with CH₂Cl₂ (40 mL \times 3). Then, the organic layer was dried over $Na₂SO₄$. The solvent was removed under reduced pressure. The final product was purified using column chromatography (PE/EtOAc = $2/1$) to afford the target compound as a brown solid (762 mg, yield: 35%). HRMS: m/z [M + H⁺] = 488.1973, [M + 2H⁺] = 244.6021; calcd for $[C_{31}H_{25}N_3O_3 + H^+]$: 488.1974, $[C_{31}H_{25}N_3O_3 + 2H^+]$: 244.6053. ¹H NMR (400 MHz, CDCl₃, ppm): δ 1.16 (t, 6H, J = 7.0 Hz, $-CH_3$), 3.34 (q, 4H, $J = 7.0$ Hz, $-CH_2$), 6.34 (dd, 1H, $J = 8.8$ Hz, 2.8 Hz, Ar–H), 6.45 (d, 1H, $J = 2.8$ Hz, Ar–H), 6.56 (d, 1H, $J =$ 8.8 Hz, Ar-H), 6.71 (t, 1H, $J = 7.0$ Hz, Ar-H), 7.10 (t, 1H, $J = 7.0$ Hz, Ar-H), 7.19 (d, 1H, $J = 7.2$ Hz, Ar-H), 7.24 (d, 1H, $J = 6.8$ Hz, Ar–H), 7.31 (d, 1H, $J = 8.4$ Hz, Ar–H), 7.53 (d, 1H, $J = 9.2$ Hz, Ar– H), 7.59 (m, 3H, Ar-H), 8.01 (m, 3H, Ar-H). ¹³C NMR (100 MHz, CDCl3, ppm): δ 12.64, 44.60, 84.06, 97.81, 105.31, 108.02, 108.62, 112.59, 117.39, 117.66, 119.73, 124.21, 124.86, 125.11, 125.55, 125.65, 127.01, 128.60, 128.85, 129.37, 129.62, 135.06, 144.70, 145.58, 149.76, 151.699, 152.77, 153.59, 169.93. **Sensors Articles.**

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Synthesis of Rh-Ip-Hy. To an ethanolic solution (10 mL) of Rh-Ip (250 mg, 0.51 mmol), hydrazine hydrate (1 mL) was added. The mixture was heated at 100 °C for 4 h. After cooling, the solvent was removed under reduced pressure. The residue was purified by column chromatography $\rm (CH_2 Cl_2/C_2H_5OH = 100/1$) to afford the target compound as a pale brown solid (128 mg, yield: 50%). HRMS: m/z [M + H⁺] = 502.2238, $[M + 2H^+]$ = 251.6155; calcd for $[C_{31}H_{27}N_5O_2 + H^+]$: 502.2238, $\left[C_{31}H_{27}N_{5}O_{2} + 2H^{+}\right]:$ 251.6155. ¹H NMR (400 MHz, CDCl₃, ppm, contain H₂O): δ 1.18 (t, 6H, J = 7.0 Hz, -CH₃), 3.35 (q, 4H, $J = 7.0$ Hz, $-CH_2$), 3.79 (s, 2H, $-NH_2$), 6.33 (dd, 1H, $J = 8.8$ Hz, 2.4 Hz, Ar-H), 6.47 (d, 1H, J = 2.4 Hz, Ar-H), 6.49 (d, 1H, $J = 8.8$ Hz, Ar–H), 6.73 (t, 1H, $J = 7.8$ Hz, Ar–H), 7.12 (t, 2H, $J = 7.8$ Hz, Ar–H), 7.21 (d, 1H, $J = 2.0$ Hz, Ar–H), 7.31 (dd, 1H, J = 8.4 Hz, 2.0 Hz, Ar–H), 7.45–7.55 (m, 3H, Ar– H), 7.64 (d, 1H, J = 8.4 Hz, Ar–H), 7.95–7.99 (m, 2H, Ar–H), 8.02 (d, 1H, $J = 6.8$ Hz, Ar-H). ¹³C NMR (100 MHz, CDCl₃, ppm): δ 12.67, 44.48, 65.99, 98.28, 98.60, 104.27, 107.94, 108.63, 112.53, 117.38, 117.82, 119.31, 123.37, 123.89, 124.69, 124.78, 125.57, 125.60, 127.79, 128.54, 129.50, 129.54, 132.92, 144.73, 145.61, 149.16, 151.56, 152.50, 153.35, 166.90.

Results and discussion

Synthesis and characterization of Rh-Ip-Hy

The synthesis procedure for the probe Rh-Ip-Hy is shown in Scheme 2. First, compound 1 was prepared according to the

Scheme 2 Synthesis routes of Rh-Ip-Hy. Reagents and conditions: a) $Na₂CO₃$, EtOH, reflux for 20 h; b) 2-(4-(N,N-diethylamino)-2hydroxybenzoyl)benzoic acid, TFA/MeSO₃H (v/v, 1/1), 80 °C for 24 h; c) hydrazine hydrate, EtOH, 100 °C for 4 h.

literature²² by the reaction of 2-bromo-1-(4-hydroxyphenyl) ethanone with 2-aminopyridine and was allowed to react with 2-(4-(N,N-diethylamino)-2-hydroxyben-zoyl)benzoic acid according to a conveniently acid-promoted Friedel–Crafts acylation protocol in CF_3COOH/CH_3SO_3H (v/v, 1/1) to produce Rh-Ip. Subsequently, the target probe Rh-Ip-Hy was prepared with a yield of 50% through the reaction of Rh-Ip with hydrazine hydrate in anhydrous ethanol. The structures of Rh-Ip and Rh-Ip-Hy were characterized using highresolution mass spectrometry (HRMS) and 1 H- and 13 C-NMR spectroscopy (see the ESI†).

Sensing performance of the probe for mercury ions

To determine the appropriate solvent environment for the interaction of Rh-Ip-Hy and Hg^{2+} , the fluorescence response of Rh-Ip-Hy towards Hg^{2+} was determined in mixtures of water and different organic solvents $(v/v, H₂O/organic)$ solvent = $4/1$). As shown in Fig. S1, \dagger after the addition of Hg^{2+} to the probe solutions, no obvious emission changes were observed in DMF/H_2O , $DMSO/H_2O$, and CH_3CN/H_2O systems. In contrast, there was a significant fluorescence enhancement in the $EtOH/H₂O$ system, indicating interactions between Rh-Ip-Hy and Hg^{2+} . Therefore, the EtOH/H₂O (v/v, 1/4) solvent system was chosen to investigate the spectral response of the probe to Hg^{2+} .

Firstly, the spectra of $Rh-Ip-Hy$ (5 μ M) after adding increasing amounts of mercury ions in PBS buffer (10 mM, EtOH/H₂O = $1/4$, pH 7.4) were investigated. As shown in Fig. 1a, the probe Rh-Ip-Hy exhibited almost no absorption within the range of 400–700 nm, which is attributed to the closed ring structure of the spirolactam. After the addition of Hg^{2+} (0-80 μ M), the absorbance of the probe at 500 nm gradually increased with the increasing concentrations of Hg^{2+} , indicating that Hg^{2+} induced the ring-opening form of Rh-Ip-Hy (Scheme 1B). For fluorescence titration, as shown in Fig. 1b, the gradual addition of Hg^{2+} to the probe solution resulted in an increased emission band at 575 nm, which reached the maximum and remained stable when the concentration of Hg^{2+} ions exceeded 50 μM. The data

Fig. 1 a) UV-vis and b) fluorescence spectra of Rh-Ip-Hy (5 μ M) after adding increasing amounts of Hg^{2+} in the PBS buffer (10 mM, EtOH/ $H₂O = 1/4$, pH of 7.4). Insets: the corresponding intensity changes of the probe as a function of Hq^{2+} concentration.

suggested that the interactions between the probe and the Hg^{2+} ions led to a colored and fluorescent compound (Scheme 1B). Furthermore, Rh-Ip-Hy exhibited a larger Stokes shift (75 nm) than the classic rhodamine and fluorescein dyes with a typical Stokes shift of less than 30 nm, which could reduce self-absorption and prevent selfquenching from excitation light. A linear relationship was observed between the fluorescence intensities of Rh-Ip-Hy and Hg²⁺ within the concentration range of 0–50 μ M. The detection limit of Hg^{2+} ions was found to be 57.57 nM $(R² = 0.9981, \sigma = 0.6094, k = 3.176 \times 10⁷, n = 10)$ based on 3 times the standard deviation rule (Fig. S2†).

Then, the spectral properties of Rh-Ip-Hy in the absence and presence of various metal ions were investigated. As shown in Fig. 2a, Rh-Ip-Hy $(5 \mu M)$ showed almost no emission beyond 500 nm, indicating the closed spirolactam ring structure of the probe. After the addition of various metal ions (50 μ M) including Ag⁺, Al³⁺, Ca²⁺, Cd²⁺, Co²⁺, Cu²⁺, Fe³⁺, Fe²⁺, K⁺, Li⁺, Mg²⁺, Mn²⁺, Na⁺, Ni⁺, Pb²⁺, and Zn^{2+} to the solution of the probe, no significant emission changes were observed. In contrast, the addition of Hg^{2+} triggered a distinct emission intensity increase at 575 nm, with a more than 20-fold enhancement in the intensity. Since some spirolactam-based xanthene probes have been reported to react with ClO[−] or other reactive oxygen species (ROS) , $23-25$ the effect of ROS on the fluorescence of Rh-Ip-Hy was also investigated. As shown in Fig. 2a, the addition of ROS (including H_2O_2 , NaClO, and \cdot OH) did not produce any obvious changes in the fluorescence of the probe. The

Fig. 2 a) Fluorescence spectra of Rh-Ip-Hy (5 μ M) in the presence of 50 μM of various interfering species. Inset: histogram representing the fluorescence enhancement of the probe at 575 nm in the presence of various interfering species. 1: probe, 2: Cu^{2+} , 3: Ag⁺, 4: Fe²⁺, 5: Zn²⁺, 6: Pb²⁺, 7: Ni²⁺, 8: Mg²⁺, 9: Al³⁺, 10: Hg²⁺, 11: Co²⁺, 12: Cd²⁺, 13: Ca²⁺ 14: Ba²⁺, 15: K⁺, 16: Fe³⁺, 17: Li⁺, 18: ClO[−], 19: H₂O₂, and 20: \cdot OH; b) changes ($(F_i - F_0)/F_0$) in the fluorescence intensity (575 nm) of the probe (5 μ M) upon the addition of each metal ion (50 μ M) followed by Hg²⁺ (50 μM). $λ_{ex}$: 500 nm. Slit: 10 nm, 10 nm.

results indicated the high selectivity of Rh-Ip-Hy to Hg^{2+} among various metal ions. Furthermore, the antiinterference experiment revealed that other metal ions did not have any significant effect on the sensing of mercury ions by the probe (Fig. 2b). These results suggested that Rh-**Ip-Hy** could be employed as a tool for tracing Hg^{2+} in biological samples.

For practical and bioimaging applications, the timedependent fluorescence responses of $\mathbf{Rh}\text{-}\mathbf{Ip}\text{-}\mathbf{Hy}$ to Hg^{2+} were recorded. As shown in Fig. 3a, the addition of Hg^{2+} to the solution of the probe induced a significant fluorescence enhancement at 575 nm within the first 10 min. The fluorescence gradually stabilized within 30 min, indicating the reaction equilibrium between the probe and the Hg^{2+} ions. Then, experiments to assess the pH dependency of the fluorescence were conducted in the presence and absence of Hg^{2+} . As shown in Fig. 3b, the probe exhibited weak fluorescence within the pH range of 1–14, which was attributed to the closed spirolactam ring structure of the probe. Compared with many reported spirolactam ring structure probes, $26,27$ the probe Rh-Ip-Hy exhibited high pH tolerance (Table S1†). After the addition of Hg^{2+} , the probe exhibited an obvious and stable fluorescence enhancement at 575 nm in the pH range of 5–11. These results showed that Rh-Ip-Hy could detect mercury ions within a wide pH range (5.0–11.0), which also covers the physiological pH conditions.

Fig. 3 a) Time-dependent fluorescence change (at 575 nm) of Rh-Ip-Hy (5 μ M) with the addition of Hg²⁺ (10 equiv.) in the PBS buffer solution (10 mM, EtOH/H₂O = 1/4, pH of 7.4). b) Profile of pH dependence of the fluorescence intensity of the probe (5 μ M) at 575 nm in the absence and presence of Hg^{2+} (10 equiv.) in the PBS buffer solution (10 mM, EtOH/H₂O = $1/4$).

Investigation of the sensing mechanism

To elucidate the sensing mechanism of Rh-Ip-Hy for Hg^{2+} , HRMS of the probe before and after its reaction with Hg^{2+} was conducted. As shown in Fig. S3a,† the probe alone exhibited a peak at (m/z) 502.22, which corresponds to the Rh-Ip-Hy $[M + H]^{+}$ ion (calcd.: 502.22). After reacting with three equivalents of Hg^{2+} for 20 min, the intensity of the peak at (m/z) 502.22 was considerably lower, and a new peak was observed at (m/z) 488.19, which corresponds to the hydrolyzed product of Rh-Ip (Fig. S3b†). Based on the HRMS results, we therefore suggest that the fluorescence of the probe can be switched on and off because of the selective Hg^{2+} -promoted opening of the spirolactam ring and the subsequent hydrolysis process (Scheme 3). Similar reaction mechanisms have also been reported in previous studies.²⁸⁻³²

Scheme 3 Suggested recognition mechanism of Rh-Ip-Hy for Hg^{2+} .

Application in cell imaging

To further explore the potential biological properties of Rh-Ip-Hy, its cytotoxicity to HeLa cells was evaluated using the CCK-8 assay. As shown in Fig. 4, after co-incubation with the probe (40 μM) for 24 hours, the survival rate of the cells was greater than 80%, which indicated that the probe had low cytotoxicity to cells and could be used for bio-imaging. Cell imaging for Hg^{2+} was conducted using **Rh-Ip-Hy** as a tracer. As shown in Fig. 5, the HeLa cells incubated with the probe $(5 \mu M)$ exhibited almost no fluorescence in the green channel (500–550 nm). However, significant and increased fluorescence was observed when the above-mentioned cells were treated with 2 equivalents of Hg^{2+} . These results demonstrated that **Rh-Ip-Hy** can permeate the cell and be used as a tracer for the detection of mercury ions in living cells. **Example the statistical on the common access Article is licensed to the properties of Kh sp. 3024. Downloaded on 9/10/2024 9:40:57 AM. This article is licensed under the properties of Kh sp. 3024. Downloaded under the pr**

Detection of Hg^{2+} with paper strips in real water samples

To facilitate the convenient on-site use of the fluorescence method, test strips were prepared by immersing filter paper strips into an ethanolic solution of $Rh-Ip-Hy$ (100 μ M) for 1 h. The strips were then dried with a blow-dryer. Hg^{2+} solutions of different concentrations (0, 100, 150, 200, and 250 μ M) were dropwise pipetted onto the dried paper strips.

Fig. 4 Cell viability of HeLa cells after incubation with different concentrations of Rh-Ip-Hy for 12 hours.

Fig. 5 Confocal fluorescence images of HeLa cells. a–c) Brightfield, fluorescence and merged images of HeLa cells after incubation with Rh-Ip-Hy (5 μM) for 30 min. d–f) Brightfield, fluorescence and merged images of HeLa cells pretreated with $Rh-Ip-Hy$ (5 μ M) for 30 min, and then incubated with Hg²⁺ (10 μ M) for 20 min.

Fig. 6 Visualization of Hg^{2+} using the test strips prepared with Rh-Ip-Hy in distilled water, tap water and lake water, respectively.

After natural drying, the color changes in the paper strips were observed under natural light. As shown in Fig. 6, the addition of Hg^{2+} induced noticeable color changes in the paper strips from sage green to red under natural light. Furthermore, the colors of the strips deepened with the increase in the concentration of Hg^{2+} . When the test strips were used for sensing Hg^{2+} -containing tap water or lake water, similar color changes were observed (Fig. 6). These results indicated that test strips with Rh-Ip-Hy as a probe can be easily prepared for the simple and convenient detection of Hg^{2+} in real water samples.

Conclusions

In summary, an imidazo[1,2-a]pyridine-functionalized xanthene fluorescent probe (Rh-Ip-Hy) was designed and prepared for detecting Hg^{2+} in various environments. Due to its asymmetric structure, the probe shows a larger Stokes shift (75 nm) than the classic rhodamine and fluorescein dyes. Because of the presence of a Lewis base site, the probe maintains a nonfluorescent ring-closed spirolactam structure within the pH range of 1–14, and shows selective fluorescence off-on signals to Hg^{2+} within a wide pHresponse range (5–11). Rh-Ip-Hy can permeate the cell membrane and can be used for the imaging of Hg^{2+} in HeLa cells. Finally, Rh-Ip-Hy-loaded strips can be easily prepared and exhibit good performance for the detection of Hg^{2+} in various water samples.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We gratefully acknowledge the National Natural Science Foundation of China (Grant No. 22274068, 22076073), the Major Basic Research Project of Natural Science Foundation of Shandong Province (No. ZR2023ZD27) and the Natural Science Foundation of Shandong Province (No. ZR2022QB123).

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