

Recognition of Hg²⁺ and Cr³⁺ in Physiological Conditions by a Rhodamine Derivative and Its Application as a Reagent for Cell-Imaging Studies

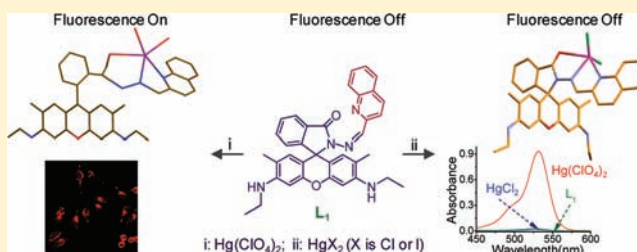
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Supporting Information

ABSTRACT: A new rhodamine-based receptor, derivatized with an additional fluorophore (quinoline), was synthesized for selective recognition of Hg²⁺ and Cr³⁺ in an acetonitrile/HEPES buffer medium of pH 7.3. This reagent could be used as a dual probe and allowed detection of these two ions by monitoring changes in absorption and the fluorescence spectral pattern. In both instances, the extent of the changes was significant enough to allow visual detection. More importantly, the receptor molecule could be used as an imaging reagent for detection of Hg²⁺ and Cr³⁺ uptake in live human cancer cells (MCF7) using laser confocal microscopic studies. Unlike Hg(ClO₄)₂ or Hg(NO₃)₂ salts, HgCl₂ or HgI₂ failed to induce any visually detectable change in color or fluorescence upon interaction with L₁ under identical experimental conditions. Presumably, the higher covalent nature of Hg^{II} in HgCl₂ or HgI₂ accounts for its lower acidity and its inability to open up the spirolactam ring of the reagent L₁. The issue has been addressed on the basis of the single-crystal X-ray structures of L₁·HgX₂ (X⁻ = Cl⁻ or I⁻) and results from other spectral studies.



INTRODUCTION

The design of optical sensors for selective recognition and sensing of desired metal ions is an important and contemporary research area. In this regard, metal ions that are known to have detrimental effects on living organisms or the environment are generally more common as target metal ions for such studies.¹ Apart from this, other critical aspects like signal transduction and the possibility of using these reagents in physiological conditions generally influence the receptor design.² The Hg²⁺ ion is considered to be one of the most toxic metal ions and thus one of the most common among different metal ions that are being studied for recognition studies. The extreme toxicity of mercury and its derivatives results from their affinities toward thiol groups in proteins and enzymes.³ This leads to the malfunctioning of the living cells and eventually leads to serious health hazards. More importantly, the Hg²⁺ ion present in soil or in effluent water is assimilated by the lower aquatic organisms, which are known to convert it to methylmercury, one of the most potent neurotoxins for human race.⁴ Reports also reveal that this methylmercury moves upward in the food chain through the marine predator animals and bioaccumulates in human beings with manifestation of different neurological disorders.⁵ Bioaccumulation of Hg²⁺ from atmospheric deposition is also known to happen in certain mosses and tree leaves, and this adversely affects photosynthesis and transpiration in plants; the former is believed to have an impact

on the global carbon cycle.⁶ Unlike Hg²⁺, Cr³⁺ is less harmful to human life, though chromium present in other higher oxidation states (4+ and 6+) has a grave consequence on human health. Cr³⁺ is an effective nutrient and gives immunity power to the human body to prevent various diseases like diabetes, cardiovascular disease, etc.⁷ Further, its deficiency is known to influence the metabolism of glucose and lipids adversely and causes maturity-onset diabetes, cardiovascular diseases, and nervous system disorders.⁸ However, exposure to higher concentrations of Cr³⁺ is known to inflict a negative effect on normal enzymatic activities. A recent study reveals that soluble Cr³⁺ at pH 6–8 can be found transiently in significant concentrations and has an adverse influence on microorganisms, like *Shewanella* sp. MR-4.⁹ The Cr³⁺ ion, present in the cytoplasm, is known to bind nonspecifically to DNA and other cellular components, and these processes become important when the concentration of Cr³⁺ exceeds a certain threshold value. These are known to inhibit DNA transcription and possibly replication.¹⁰ All of these have made it almost obligatory to develop efficient sensor molecules, which can be used for the efficient and selective detection of these two ions present in trace quantities. These reagents may become even more versatile if these are found to work in an aqueous

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environment and at physiological pH because this opens up the possibility for their *in vivo* application. Both of these metal ions are known to be extensively solvated in an aqueous medium, and the unfavorably high enthalpy of solvation poses a challenge to chemists in developing a suitable receptor for sensing of these ions in an aqueous environment; to date, methods for monitoring the actual concentration of Hg^{2+} or Cr^{3+} *in vivo* are rather limited.^{7c,8a,11}

Current approaches for environmental and clinical samples rely on costly, time-consuming methods like atomic absorption/emission spectroscopy or inductively coupled plasma mass spectrometry,¹² which are not very convenient and handy for “in-field” detection. These limitations have actually set off an enormous interest among chemists for the development of efficient, cost-effective, and reversible chemosensors for Hg^{2+} and Cr^{3+} . Although a number of selective sensors for Hg^{2+} have been devised using various modes of signal transduction processes through detectable changes in either redox¹³ or spectral (electronic and fluorescence)^{14,15} properties, such examples are uncommon for Cr^{3+} in the literature. In this regard, a receptor molecule that could be used as a dual probe for the detection and sensing of the desired metal ions by probing the metal-ion binding induced changes in the electronic as well as fluorescence spectral properties because the output signal has a distinct edge over other receptors because these receptors could be used either as colorimetric or fluorogenic sensors. Colorimetric sensors allow easy in-field detection, while fluorescence-based sensors provide an edge in imaging studies. Hg^{2+} and Cr^{3+} are known to quench effectively the excited state of the fluorophore through an efficient spin-orbit coupling and paramagnetic effect, respectively.¹⁶ Thus, examples of fluorescence-based reversible sensor molecules for these two ions for sensing and imaging applications are not common, except the reagents that were designed based on the spiro lactam to open ring amide equilibrium process of rhodamine derivatives.¹⁷ Appropriately functionalized rhodamine derivatives belong to a class of receptors that allow detection of Hg^{2+} present in an aqueous solution through visually detectable changes in color, as well as through a *switch-on* fluorescence response. This dual detection mode has made various rhodamine derivatives an attractive choice for the design of receptors for Hg^{2+} . However, examples for the use of rhodamine derivatives for the recognition of Cr^{3+} in the aqueous environment are limited.¹⁸ In this article, we report a new rhodamine derivative that binds specifically to Hg^{2+} and Cr^{3+} in the presence of a large excess of other common alkali, alkaline earth, and all common transition-metal ions. All previous references on Hg^{2+} recognition using rhodamine derivatives describe the use of $\text{Hg}(\text{NO}_3)_2$ or $\text{Hg}(\text{ClO}_4)_2$ salts, which induce a spiro lactam ring-opening reaction upon coordination to the Hg^{2+} ion with associated *switch-on* responses at ~ 530 nm for the electronic spectra and at ~ 560 nm for the fluorescence spectra.¹⁷ These allow a visual detection, as well as usage as an imaging agent for the *in vitro* and *in vivo* detection of Hg^{2+} in physiological conditions.¹⁹ However, literature reports on the use of HgCl_2 for recognition studies are very limited. Two independent reports reveal that HgCl_2 initiates an irreversible and stoichiometric reaction between adjacent thiourea and cyclic amide functionalities to yield an oxadiazole derivative with an associated spiro lactam ring-opening reaction that leads to the generation of a strongly luminescent xanthene form.²⁰ The only other two references that described the use of HgCl_2 for sensing studies reported the

use of the same thiospiro derivative of rhodamine B. On the basis of mass spectral evidence, it was proposed that, in both studies, the Hg^{2+} ion was initially coordinated to a thiocarboxylate derivative of the xanthene form of rhodamine B, which subsequently underwent a hydrolysis reaction to generate the corresponding carboxylate derivative.²¹ In one of these four examples, a mechanistic pathway involving HgCl_2 was proposed in an intermediate. However, X-ray structural evidence for such a proposition remained elusive. Literature reports reveal that HgCl_2 and HgI_2 , two commonly available halide salts, are more covalent in nature. HgCl_2 possesses only 23% ionic character, while HgI_2 has even less ionic character.²² Interaction of such a nonionic Hg^{II} center with a spiro lactam derivative is not well understood, and this is discussed in this manuscript with single-crystal structural evidence using a new reagent, L_1 . This new reagent, L_1 , could be used as a reversible and selective fluorogenic, as well as colorimetric sensor for Hg^{2+} and Cr^{3+} in a mixed aqueous medium [CH_3CN /aqueous HEPES buffer (1 mM; 3:2, v/v)] at a physiological pH (pH 7.3). We have also shown that this receptor could be used for detecting uptake of the Hg^{2+} ion by living cells from an aqueous solution containing Hg^{2+} at pH 7.3 through cell imaging studies. In the present study, we have incorporated a quinoline unit with the aim of having another fluorescent fragment, along with the Rhodamine 6G derivative, to enable us to probe the metal-ion recognition processes at two different monitoring wavelengths of the fluorescence spectrum to avoid the possibility of any interference at any particular wavelength. However, no quinoline-based fluorescence was observed for either free L_1 or L_1 bound to Hg^{2+} or Cr^{3+} . The reasons for such an observation are also discussed.

■ EXPERIMENTAL SECTION

Rhodamine 6G hydrochloride, quinoline-2-aldehyde, LiClO_4 , NaClO_4 , KClO_4 , CsClO_4 , $\text{Mg}(\text{ClO}_4)_2$, $\text{Ca}(\text{ClO}_4)_2$, $\text{Ba}(\text{ClO}_4)_2$, $\text{Sr}(\text{ClO}_4)_2$, $\text{Cu}(\text{ClO}_4)_2$, $\text{Zn}(\text{ClO}_4)_2$, $\text{Co}(\text{ClO}_4)_2$, $\text{Ni}(\text{ClO}_4)_2$, $\text{Cr}(\text{ClO}_4)_3$, $\text{Fe}(\text{ClO}_4)_2$, $\text{Fe}(\text{ClO}_4)_3$, $\text{Cd}(\text{ClO}_4)_2$, $\text{Hg}(\text{ClO}_4)_2$, and $\text{Pb}(\text{ClO}_4)_2$ were obtained from Sigma-Aldrich and used as received. All of the other reagents used were procured from S. D. Fine Chemicals, India. Hydrazine hydrates, glacial acetic acid, acetonitrile, methanol (AR; Merck), and ethanol (Spectrosol; Spectrochem, India) were used as solvents. HPLC-grade water (Merck, India) was used for experiments and spectral studies. The Rhodamine 6G hydrazine derivative (L) was synthesized following the reported procedure.²³ Electrospray ionization mass spectrometry (ESI-MS) measurements were carried out on a Waters QToF-Micro instrument. Microanalysis (C, H, and N) was performed using a Perkin-Elmer 4100 elemental analyzer. Fourier transform infrared (FTIR) spectra were recorded as KBr pellets using a Perkin-Elmer Spectra GX 2000 spectrometer. ^1H and ^{13}C NMR spectra were recorded on a Bruker 500 MHz FT NMR (model Avance-DPX 500). Electronic spectra were recorded with a Shimadzu UV-3101 PC/Varian Cary 500 Scan UV-vis-near-IR spectrophotometer. Emission spectra were recorded using an Edinburgh Instruments model Xe-900. Time-resolved fluorescence measurements were carried out using a time-correlated single-photon-counting (TCSPC) technique using an Edinburgh Instruments model OB920. A 360 nm light-emitting diode (LED) was used as the excitation source for the present studies.

Crystals of suitable size were selected from the mother liquor, immersed in paratone oil, then mounted on the tip of a glass fiber, and cemented using epoxy resin. Intensity data for both crystals were collected using $\text{Mo K}\alpha$ ($\lambda = 0.71073$ Å) radiation on a Bruker SMART APEX diffractometer equipped with a CCD area detector at 110 K. The data integration and reduction were processed with *SAINT* software.^{24a} An empirical absorption correction was applied to the collected reflections with *SADABS*.^{24b} The structures were solved by

direct methods using *SHELXTL*^{24c} and were refined on F^2 by the full-matrix least-squares technique using the *SHELXL-97* program.^{24d} All non-hydrogen atoms were refined anisotropically until convergence was reached. Hydrogen atoms attached to the ligand moieties and solvents of crystallization were stereochemically fixed in both complexes. The diagrams of the crystal structures were generated using *Mercury* 2.3. A summary of the crystallographic data and details of data collection for all three compounds are given in Table 1.

Table 1. Crystallographic Data for $L_1 \cdot [HgCl_2]$ and $L_1 \cdot [HgI_2]$

identification code	$L_1 \cdot [HgCl_2]$	$L_1 \cdot [HgI_2]$
CCDC number	CCDC 832870	CCDC 832871
empirical formula	$C_{75}H_{72}Cl_{10}Hg_2N_{10}O_5$	$C_{36}H_{33}HgI_2N_5O_2$
fw	1949.11	1022.06
temperature (K)	110(2)	110(2)
wavelength (Å)	0.710 73	0.710 73
cryst syst	monoclinic	monoclinic
space group	$C2/c$	$C2/c$
<i>a</i> (Å)	32.831	38.4292(17)
<i>b</i> (Å)	17.1862(11)	9.4617(4)
<i>c</i> (Å)	29.3246(18)	19.7407(9)
β (deg)	110.4270(10)	105.3580(10)
volume (Å ³)	15505.8(17)	6921.5(5)
<i>Z</i>	8	8
density (calcd) (Mg m ⁻³)	1.670	1.962
abs coeff (mm ⁻¹)	4.357	6.271
<i>F</i> (000)	7696	3888
crystal size (mm ³)	0.64 × 0.43 × 0.38	0.35 × 0.14 × 0.11
reflections	45 481	20 208
independent	17 947 [R(int) = 0.0297]	8038 [R(int) = 0.0234]
refinement method	full-matrix least squares on F^2	full-matrix least squares on F^2
data/restraints/no. of parameters	17 947/0/945	8038/0/427
GOF on F^2	1.018	1.145
final <i>R</i> indices [$I > 2\sigma(I)$]	<i>R</i> 1 = 0.0380, <i>wR</i> 2 = 0.0996	<i>R</i> 1 = 0.0289, <i>wR</i> 2 = 0.0813
<i>R</i> indices (all data)	<i>R</i> 1 = 0.0469, <i>wR</i> 2 = 0.1035	<i>R</i> 1 = 0.0340, <i>wR</i> 2 = 0.0991
largest diff peak and hole (e Å ⁻³)	1.804 and -1.314	1.323 and -0.656

MCF7 cells were grown in Dulbecco's Modified Eagle Medium supplemented with 10% fetal bovine serum at 37 °C were trypsinized, and 1×10^4 cells were added in each well in a 24-well culture plate. After 48 h of growth, the cells were treated with different concentrations (0–4.0 μ M) of $Cr(ClO_4)_3$ and $Hg(ClO_4)_2$ dissolved in 1X phosphate-buffered saline for 1 h. MCF7 cells were then stained with a 4 μ M solution of L_1 dissolved in acetonitrile/aqueous HEPES (3:2, v/v) for 60 min at 37 °C. Cells were viewed under a laser scanning confocal microscope (FV100, Olympus) at an excitation wavelength of 543 nm.

Synthesis of L_1 . Rhodamine 6G hydrazine (L) (300 mg, 0.646 mmol) and quinoline-2-aldehyde (101 mg, 0.646 mmol) were dissolved in 20 mL of methanol. To this was added approximately 2 drops of acetic acid, and the resulting solution was refluxed for 10 h. An off-white precipitate was found. The reaction mixture was allowed to attain room temperature, and then the precipitate was collected through filtration. The residue was washed thoroughly with methanol to isolate L_1 in pure form with 76% yield (the yield was calculated based on the starting reagents). ¹H NMR [500 MHz, $CDCl_3$, $SiMe_4$, J (Hz), δ (ppm)]: 8.530 (1H, s, $-CH_{10}=N-$), 8.117 (1H, d, $J = 9$, ArH₇), 8.056–8.013 (1H, m, ArH₁₄), 7.975 (1H, d, $J = 9$, ArH₁₃), 7.719 (1H, d, $J = 8$, ArH₁₆), 7.618 (1H, t, $J = 8$, ArH₁₅), 7.513–7.443 (4H, m, ArH₂, H₃, H₄, H₅), 7.08 (1H, d, $J = 7$, ArH₈), 6.453 (2H, s,

ArH₂₀, H₂₉), 6.382 (2H, s, ArH₂₃, H₂₆), 3.477 (2H, s, $-NH-$), 3.21 (4H, q, H₃₂, H₃₄), 1.860 (6H, s, H₃₁, H₃₆), 1.306 (6H, t, H₃₃, H₃₅). ¹³C NMR [125 MHz, $CDCl_3$, $SiMe_4$, δ (ppm)]: 165.804, 155.049, 152.894, 151.458, 147.949, 147.815, 146.433, 135.965, 134.072, 129.525, 129.463, 128.492, 128.406, 128.075, 127.717, 127.544, 126.960, 123.902, 118.242, 106.206, 97.304, 66.209, 38.555, 16.885, 14.948. ESI-MS (positive mode, m/z). Calcd for $C_{36}H_{33}N_5O_2$: 567.58. Found: 568.34 (40%; $M + H^+$), 590.33 (100%; $M + Na^+$). Elem Anal. Calcd: C, 76.17; H, 5.86; N, 12.34; O, 5.64. Exptl: C, 75.8; H, 5.9; N, 12.3.

General Methodology Adopted for Spectroscopic Studies.

A solution of the perchlorate salts of the respective ions (Li^+ , K^+ , Na^+ , Ca^{2+} , Mg^{2+} , Fe^{2+} , Fe^{3+} , Co^{2+} , Ni^{2+} , Cu^{2+} , Zn^{2+} , Cd^{2+} , Pb^{2+} , Ba^{2+} , Sr^{2+} , Cr^{3+} , and Hg^{2+}) was used for the studies. For checking the relative, but qualitative, binding affinity of individual metal ions toward the reagent L_1 , the effective metal-ion concentration in acetonitrile/aqueous HEPES buffer (1 mM; 3:2, v/v) was maintained at 1.0×10^{-4} M, while that for the receptor L_1 was maintained at 7.0×10^{-6} M. The solution pH was found to be 7.3.

Evaluation of the Association Constant for the Formation of $L_1 \cdot Hg^{2+}$ and $L_1 \cdot Cr^{3+}$. Receptor L_1 was dissolved in acetonitrile/aqueous HEPES buffer (1 mM; 3:2, v/v; pH 7.3) and stored in dark conditions. This solution composition was maintained for all spectrophotometric titration studies. A stock solution of $Hg(ClO_4)_2$ having a concentration of 1.18×10^{-3} M, was prepared in the same solvent composition. A solution of L_1 having an effective concentration of 7.05×10^{-6} M was used for spectroscopic titrations, while final concentrations for the Hg^{2+} ion were varied between 0 and 7.13×10^{-4} M. All solutions were stored at 25 °C. The solution pH was adjusted to 7.3 using an aqueous HEPES buffer solution having an effective concentration of 1 mM.

An effective concentration of 7.76×10^{-6} M for L_1 in acetonitrile/aqueous HEPES buffer (1 mM; 3:2, v/v; pH 7.3) was used for titration studies with a Cr^{3+} solution. A 1.46×10^{-3} M stock solution of $Cr(ClO_4)_3$ was prepared in the same solvent composition and was used for spectrophotometric titration studies, while effective concentrations of Cr^{3+} were varied between 0 and 1.02×10^{-3} M. The solution pH was maintained at 7.3 using an aqueous HEPES buffer solution with an effective concentration of 1 mM.

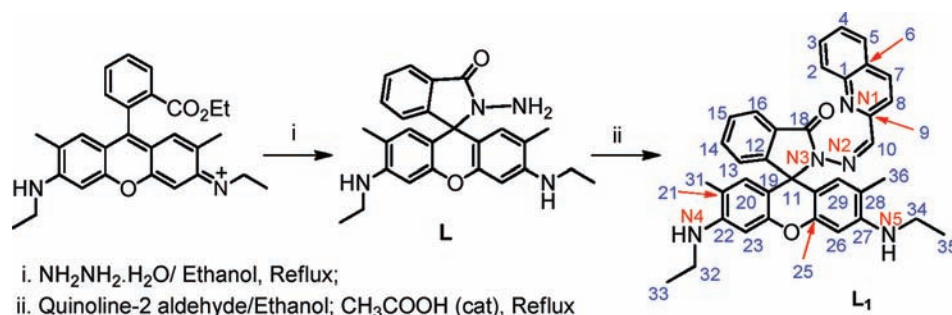
General Methodology Adopted for Fluorescence Studies.

A solution of the perchlorate salts of the respective ions (Li^+ , Na^+ , K^+ , Ca^{2+} , Mg^{2+} , Ba^{2+} , Sr^{2+} , Fe^{2+} , Fe^{3+} , Co^{2+} , Ni^{2+} , Cu^{2+} , Zn^{2+} , Cd^{2+} , Pb^{2+} , Ba^{2+} , Sr^{2+} , Cr^{3+} , and Hg^{2+}) in a spectroscopic-grade acetonitrile/aqueous HEPES buffer (1 mM; 3:2, v/v; pH 7.3) solvent mixture was used for the studies, while maintaining an effective concentration of the respective metal ion at 2.0×10^{-4} M. A solution of the receptor L_1 with an effective concentration of 10.0×10^{-6} M was prepared in the same mixed-solvent medium and was used for checking the relative binding affinity with different metal ions.

Evaluation of the Association Constant for the Formation of $L_1 \cdot Hg^{2+}$ and $L_1 \cdot Cr^{3+}$. Receptor L_1 with an effective concentration of 7.0×10^{-6} M in an acetonitrile/aqueous HEPES buffer (1 mM; 3:2, v/v; pH 7.3) was used for the emission titration studies with a Hg^{2+} solution. A stock solution of $Hg(ClO_4)_2$ having a concentration of 8.41×10^{-3} M in an acetonitrile/aqueous HEPES buffer (3:2, v/v; pH 7.3) solution was used.

The formation constant for $L_1 \cdot Cr^{3+}$ was also evaluated from fluorescence titration data. For this, the effective concentration of L_1 was maintained at 7.76×10^{-6} M in acetonitrile/aqueous HEPES buffer (3:2, v/v; pH 7.3), while a stock solution of Cr^{3+} having a concentration of 2.54×10^{-3} M in an acetonitrile/aqueous HEPES buffer (3:2, v/v) medium of pH 7.3 was used for this study. The effective Cr^{3+} concentration was varied between 0 and 7.93×10^{-4} M for this titration. In both cases, the solution pH was adjusted to 7.3 using an aqueous HEPES buffer solution having an effective concentration of 1 mM.

Calculations for the Binding Constants Using Spectrophotometric Titration Data. The association constant and stoichiometry for the formation of the respective complexes were evaluated using the Benesi–Hildebrand (B–H) plot (eq 1).²⁵

Scheme 1. Methodology Adopted for the Synthesis of L₁

$$\frac{1}{(A - A_0)} = \frac{1}{K(A_{\text{max}} - A_0)}[M^{n+}] + \frac{1}{(A_{\text{max}} - A_0)} \quad (1)$$

A_0 is the absorbance of L₁ at absorbance maxima ($\lambda = 527$ nm for Hg^{2+} and 531 nm for Cr^{3+}), A is the observed absorbance at that particular wavelength in the presence of a certain concentration of the metal ion (M^{n+}), A_{max} is the maximum absorbance value that was obtained at $\lambda = 527$ nm (Hg^{2+}) or 531 nm (Cr^{3+}) during titration with varying $[M^{n+}]$, K is the association constant (M^{-1}) and was determined from the slope of the linear plot, and $[M^{n+}]$ is the concentration of the Hg^{2+} or Cr^{3+} ion added during titration studies. The goodness of the linear fit of the B–H plot of $1/(A - A_0)$ vs $1/[\text{Hg}^{2+}]$ or $1/[\text{Cr}^{3+}]$ for 1:1 complex formation confirms the binding stoichiometry between L₁ and Hg^{2+} or Cr^{3+} .

$$\frac{1}{(I - I_0)} = \frac{1}{K(I_{\text{max}} - I_0)}[M^{n+}] + \frac{1}{(I_{\text{max}} - I_0)} \quad (2)$$

Binding stoichiometries for the respective complex formations were also confirmed from Job's plot. In the case of evaluation of the binding constant from the results of fluorescence titration, a modified B–H equation (eq 2) was used, where I_0 , I_{max} and I represent the emission intensity of free L₁, the maximum emission intensity observed in the presence of added metal ion at 555 nm for Hg^{2+} ($\lambda_{\text{ext}} = 500$ nm) or 557 nm for Cr^{3+} ($\lambda_{\text{ext}} = 500$ nm), and the emission intensity at a certain concentration of the metal ion added, respectively.

RESULTS AND DISCUSSION

An intermediate spirolactam form of the rhodamine derivative (L) was synthesized following a reported procedure by reacting (*E*)-ethyl-2-[6-(ethylamino)-3-(ethylimino)-3*H*-xanthen-9-yl]-benzoate with hydrazine hydrate.²³ The purity of this intermediate was checked by standard analytical and spectroscopic techniques (Supporting Information). Receptor L₁ was synthesized following a Schiff base type reaction between the intermediate spirolactam derivative (L) and quinoline-2-aldehyde in methanol as the solvent (Scheme 1). A small amount (2 or 3 drops) of CH_3COOH was added in the reaction medium as a catalyst. A white precipitate appeared and was isolated through filtration. This white residue was thoroughly washed with methanol and air-dried. The results of the various analytical and spectroscopic data (elemental analysis, ^1H and ^{13}C NMR, FTIR, and ESI-MS) confirmed the desired purity of the reagent L₁, and this was used further for recognition studies. This newly synthesized compound L₁ was found to have a limited solubility in water, and thus an acetonitrile/aqueous HEPES buffer (3:2, v/v) solution was used for our studies. The solution of L₁ in an acetonitrile/aqueous HEPES buffer (3:2, v/v) medium was colorless, which revealed that L₁ was present almost exclusively in the spirolactam form under the experimental conditions. The ^{13}C NMR spectrum was recorded for L₁. A characteristic peak for

the tertiary C₁₁ atom appeared near 66 ppm, which further corroborated this proposition.^{26,27}

Single crystals of L₁ could not be obtained for X-ray structural analysis; however, the crystal X-ray structures for L₁·[HgCl₂] (1a) and L₁·[HgI₂] (1b) were obtained and are shown in Figure 1. X-ray-quality single crystals for 1a and 1b

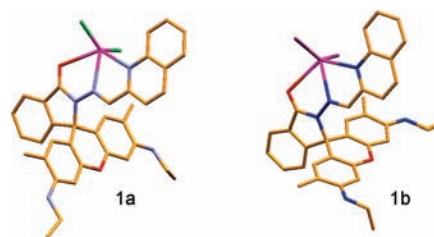


Figure 1. Single-crystal X-ray structures for 1a and 1b. Hydrogen atoms, lattice chloroform, and methanol molecules are omitted for clarity.

were grown from a $\text{CHCl}_3/\text{CH}_3\text{OH}$ solution (1:1, v/v) of L₁ and 1.2 mol equiv of HgCl_2 or HgI_2 at room temperature by a slow evaporation method.

Mercury diagrams of 1a and 1b are depicted in Figure 1, and the structural parameters for both complexes are given in Table 1. Both complexes crystallized in a monoclinic system with space group $C2/c$. In the case of 1a, two molecules of the metal complex are present in the asymmetric unit along with two CHCl_3 and one CH_3OH as solvents of crystallization. As depicted in Figure 1, for both compounds, the coordination environment around the central Hg^{II} showed a highly distorted trigonal-bipyramidal geometry in which the triangular base is constituted by the coordinated halide ions and the central azine nitrogen N2 of the rhodamine receptor. The axial positions are occupied by N1 of the quinoline ring and exocyclic ketonic oxygen O1 of the fused bicyclic ring. Interestingly, the second molecule of L₁·[HgCl₂] in the asymmetric unit of 1a also showed almost identical coordination geometry. Thus, the rhodamine-based receptor acts as a tridentate chelating ligand in the formation of neutral complexes with weak coordination involving the ketonic oxygen O1 with mercury [$\text{Hg}1 \cdots \text{O}1 = 2.709(4)/2.650(3)$ Å for 1a and $\text{Hg} \cdots \text{O}1 = 2.809(4)$ Å for 1b].

The average $\text{Hg} \cdots \text{Cl}$ distance, considering both molecules present in the asymmetric unit, is $2.3726(5)$ Å for 1a, and the $\text{Hg} \cdots \text{I}$ distance for 1b is $2.6435(4)$ Å. The $\text{Hg} \cdots \text{N}$ distance is in the range $2.441(4)–2.450(3)$ Å for 1a and $2.507(3)–2.566(4)$ Å for 1b. The crystal lattices are constructed from different types of hydrogen-bonding interactions involving the metal complex and the lattice CHCl_3 and CH_3OH (via $\text{N–H} \cdots \text{Cl}$, $\text{O–H} \cdots \text{Cl}$, $\text{N–H} \cdots \text{O}$, and $\text{C–H} \cdots \text{O}$ hydrogen bonding) for 1b,

thus stabilizing the molecule. Thus, *N*-ethyl hydrogen atoms of the rhodamine unit form intermolecular hydrogen bonding with the coordinated Cl1 and Cl3; the OH group of the lattice methanol is involved in O–H···Cl interaction with coordinated Cl3 and the alcoholic oxygen atom acts as an acceptor in N–H···O interaction with the hydrogen atom from the *N*-ethyl group. Hydrogen from both CHCl₃ molecules participates in C–H···O interaction with the exocyclic ketonic oxygen atom of the different metal complexes present in the asymmetric unit holding the volatile solvent molecule in the lattice. No classical hydrogen-bonding interaction is observed for **1b**.

Absorption spectral titrations for L₁ (7.05 × 10^{−6} M) in an acetonitrile/aqueous HEPES buffer (3:2, v/v) medium with varying pH revealed that the spirocyclic form for L₁ was retained for a pH range of 5.0–11.0. Below pH 5.0, a new distinct band appeared at 527 nm, and its intensity was found to increase with a further decrease in the solution pH. This signified the spiro lactam ring opening because the acyclic forms of rhodamine 6G derivatives are known to absorb strongly at around 530 nm. As expected, the solution of L₁ was found to be nonfluorescent in an acetonitrile/aqueous HEPES buffer (3:2, v/v) solution at neutral pH because of the nonluminescent nature of the spiro lactam form of the rhodamine derivative. However, this solution showed a strong fluorescence band with a maximum at 555 nm upon excitation at 500 nm for pH < 3.0. This also corroborated the appreciable spiro lactam ring-opening reaction of L₁ at pH < 3.0 because the xanthene forms of rhodamine 6G derivatives are known to be strongly fluorescent with an emission maximum at around 560 nm (Figure 2).²⁸ Thus, the pH range of 5–11 is suitable for using

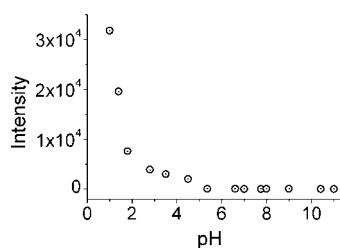


Figure 2. Change in the emission intensity at 555 nm with variation in the pH of the acetonitrile/aqueous HEPES buffer (3:2, v/v) solution for L₁.

this reagent as a probe molecule for the detection of any metal ion.

Earlier reports reveal that certain transition-metal ions bind selectively with appropriate derivatives of rhodamine, where metal-ion binding induces opening of the spiro lactam ring and generation of the xanthene form with associated changes in the electronic and fluorescence spectral patterns. Thus, we checked the binding affinity of the reagent L₁ toward a series of groups 1A (Li⁺, Na⁺, K⁺, and Cs⁺) and 2A (Mg²⁺, Ba²⁺, Ca²⁺, and Sr²⁺) and all common transition-metal ions (Fe²⁺, Fe³⁺, Co²⁺, Ni²⁺, Cu²⁺, Zn²⁺, Cd²⁺, Pb²⁺, Cr³⁺, and Hg²⁺) by monitoring the change in the electronic and fluorescence spectral patterns in a CH₃CN/aqueous HEPES buffer (3:2, v/v; pH 7.3) medium. Absorption spectra (Figure 3a) revealed a distinct change and the appearance of a new spectral band with a maximum at 527 nm only for Hg²⁺ and a maximum at 531 nm for Cr³⁺; the extent of the change in the absorption intensity was more prominent for Hg²⁺ (Figure 3). For all of the other ions used for this study, no change in the absorption spectra of L₁ was

observed. For Hg²⁺ and Cr³⁺, spectral changes were also associated with a visually detectable change from colorless to pink-red, while no such change could be visually detected for any other cations studied. These tend to suggest that the reagent L₁ could bind selectively to Hg²⁺ and Cr³⁺ in a CH₃CN/HEPES buffer (3:2, v/v) medium of pH 7.3. Interestingly, the use of a comparable concentration of HgX₂ (X[−] = Cl[−] or I[−]) failed to induce any detectable change in the absorption spectra for a predried and distilled acetonitrile solution of the reagent L₁. Spectra for this receptor in the presence of comparable concentrations of Hg(ClO₄)₂ and HgCl₂ are shown in the inset in Figure 3a. For HgCl₂, no spectral change was observed when studies were done in a CH₃CN/HEPES buffer medium (3:2, v/v) of pH 7.3.

The selective binding of L₁ to Hg²⁺ and Cr³⁺ among all other metal ions was also studied using the emission spectral response of the solution of L₁ (10.0 × 10^{−6} M) in the absence and presence of an excess (14.3 mol equiv) of each of these metal ions in CH₃CN/aqueous HEPES buffer (3:2, v/v) having a pH of 7.3 (Figure 3b). Among the various metal ions studied, a new strong emission band with a maximum at 555 nm appeared only for Hg²⁺ and Cr³⁺. Quantum yield evaluation for L₁ was not possible because it shows negligible absorbance at 530 nm in CH₃CN/aqueous HEPES buffer (3:2, v/v) of pH 7.3; however, the quantum yields (Φ) for L₁ in the presence of Hg(ClO₄)₂ and Cr(ClO₄)₃ were found to be 0.85 and 0.75, respectively, when the acyclic xanthene form of rhodamine 6G was used as a reference (λ_{ext} = 500 nm). Earlier reports reveal that the spiro lactam ring in rhodamine 6G derivatives opens up and converts to the xanthene form upon coordination to a metal ion and the corresponding xanthene derivatives absorb and emit strongly at around 530 and 560 nm, respectively.²⁹ Thus, the new emission band at 556 nm could be ascribed to the binding of the reagent L₁ only to Hg²⁺ or Cr³⁺ in a CH₃CN/aqueous HEPES buffer (3:2, v/v) medium of pH 7.3. In general, Hg²⁺ is known to quench the emission of a fluorophore because of an efficient spin–orbit coupling.^{19c,30} Cr³⁺, being a paramagnetic species, is also expected to cause an emission quenching of a fluorophore.³¹ Thus, considering these, the “switch-on” fluorescence response of the reagent L₁ upon selective binding to these two ions is of significance. Interestingly, the “switch-on” fluorescence response was observed for Hg²⁺ only when Hg^{II} salts in the form of Hg(ClO₄)₂ or Hg(NO₃)₂ were used.

The relative affinities of Hg²⁺ and Cr³⁺ (both as perchlorate salts) toward L₁ were evaluated from the data of the systematic spectrophotometric titration experiments in a CH₃CN/aqueous HEPES buffer (pH 7.3; 3:2, v/v) medium (Figure 4). For these, [L₁] was maintained at 7.05 × 10^{−6} M for studies with Hg(ClO₄)₂ and [L₁] was maintained at 7.76 × 10^{−6} M for studies with Cr(ClO₄)₃, while [Hg(ClO₄)₂] or [Cr(ClO₄)₃] was varied between 0 and 7.14 × 10^{−4} M or between 0 and 1.02 × 10^{−3} M, respectively (Figure 4). A new band appeared with a gradual increase in the absorbance intensity at 527 nm for Hg(ClO₄)₂ and at 531 nm for Cr(ClO₄)₃ (Figure 4). In both instances, a change in the solution color from colorless to pink-red was observed. A B–H plot of 1/(A – A₀) against 1/[Hg²⁺] or 1/(A – A₀) against 1/[Cr³⁺] was linear. The goodness of linear fit confirms the 1:1 binding stoichiometry between the reagent L₁ and Hg²⁺ or Cr³⁺, and the respective association constants for L₁·Hg(ClO₄)₂ and L₁·Cr(ClO₄)₃ were found to be 3.11 × 10³ and 2.0 × 10³ M^{−1}, respectively. Spectral response and association constant values remain unchanged

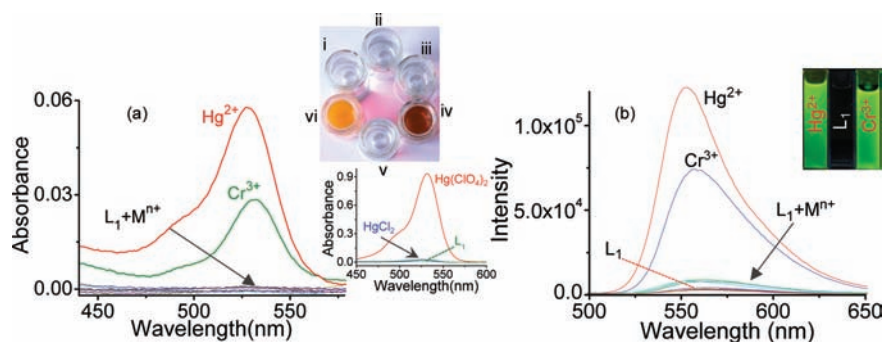


Figure 3. Electronic spectra of (a) L_1 (7.06×10^{-6} M) with different metal ions (1.0×10^{-4} M) and emission spectra of (b) L_1 (10.0×10^{-6} M) with different metal ions (2.0×10^{-4} M) in CH_3CN /aqueous HEPES buffer (3:2, v/v) of pH 7.3; $M^{n+} = \text{Li}^+, \text{Na}^+, \text{K}^+, \text{Cs}^+, \text{Mg}^{2+}, \text{Ca}^{2+}, \text{Ba}^{2+}, \text{Sr}^{2+}, \text{Cu}^{2+}, \text{Ni}^{2+}, \text{Zn}^{2+}, \text{Cd}^{2+}, \text{Co}^{2+}, \text{Fe}^{2+}, \text{Fe}^{3+}, \text{and Pb}^{2+}$. Inset: Change in the color of L_1 (7.06×10^{-6} M) in CH_3CN /aqueous HEPES buffer (3:2, v/v) of pH 7.3 in the absence and presence of different metal ions: (i) L_1 , (ii) in the presence of alkali metal ions ($\text{Li}^+, \text{Na}^+, \text{K}^+$, and Cs^+); (iii) in the presence of alkaline-earth metal ions ($\text{Mg}^{2+}, \text{Ca}^{2+}, \text{Ba}^{2+}$, and Sr^{2+}); (iv) in the presence of Hg^{2+} (1.0×10^{-4} M); (v) in the presence of any one of the transition-metal ions, except Cr^{3+} ; (vi) in the presence of Cr^{3+} (1.0×10^{-4} M). Inset: Electronic spectra of L_1 (1.12×10^{-5} M) in the presence of $\text{Hg}(\text{ClO}_4)_2$ (6.25×10^{-4} M) and HgCl_2 (6.25×10^{-4} M) in a CH_3CN medium. Inset: Change in the observed emission of L_1 (7.06×10^{-6} M) in CH_3CN /aqueous HEPES buffer (3:2, v/v) of pH 7.3 in the absence and presence of Hg^{2+} and Cr^{3+} ions added as the perchlorate salt.

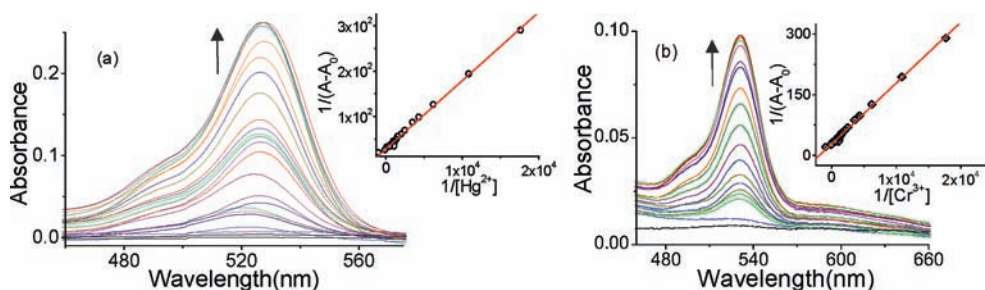


Figure 4. Electronic spectra of (a) L_1 (7.05×10^{-6} M) in the absence and presence of varying $[\text{Hg}(\text{ClO}_4)_2]$ ($0-7.14 \times 10^{-4}$ M) and (b) L_1 (7.76×10^{-6} M) in the absence and presence of varying Cr^{3+} ($0-1.02 \times 10^{-3}$ M) in an acetonitrile/aqueous HEPES buffer (1 mM) (3:2, v/v) solution of pH 7.3. Insets: linear fits of the B-H plots in support of the 1:1 binding stoichiometry for $L_1 \cdot \text{Hg}^{2+}$ and $L_1 \cdot \text{Cr}^{3+}$ formation. The ClO_4^- salt of both metal ions was used for these studies.

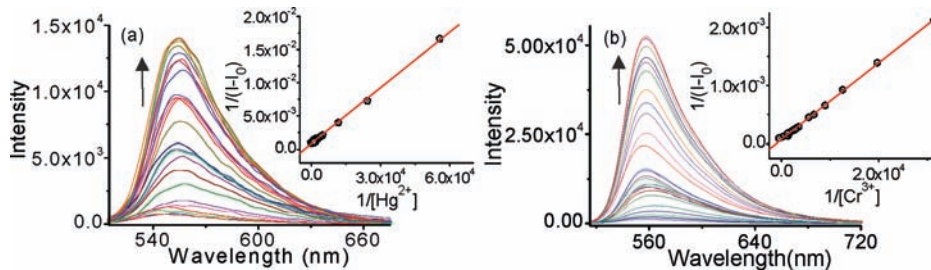


Figure 5. Plot of the change in the fluorescence spectral pattern for receptor (a) L_1 (7.0×10^{-6} M) in the presence of varying $[\text{Hg}(\text{ClO}_4)_2]$ ($0-6.39 \times 10^{-4}$ M) or (b) L_1 (7.76×10^{-6} M) in the presence of varying $[\text{Cr}(\text{ClO}_4)_3]$ ($0-7.93 \times 10^{-4}$ M) in a CH_3CN /HEPES buffer medium (1 mM; 3:2, v/v) of pH 7.3; $\lambda_{\text{ext}} = 500$ nm was used for the studies. Insets: Goodness of the linear fits of the B-H plot confirming the 1:1 binding stoichiometry.

when the spectrophotometric titration experiment was repeated using $\text{Hg}(\text{NO}_3)_2$, instead of $\text{Hg}(\text{ClO}_4)_2$ in CH_3CN /aqueous HEPES buffer (3:2, v/v) of pH 7.3. This revealed that counteranions did not have any influence on the binding process with receptor L_1 and the affinity constants for Hg^{II} salts, which were primarily ionic in nature. The results of the ESI-MS studies further supported a 1:1 binding stoichiometry for the reaction between L_1 and $\text{Hg}(\text{ClO}_4)_2$ or $\text{Cr}(\text{ClO}_4)_3$ (Supporting Information).

Emission spectra and the intensity at 556 nm remained unchanged for the pH ranges of 5–8.9 for Hg^{2+} and 5–8.5 for Cr^{3+} (Supporting Information). Thus, the binding constants for the formation of $L_1 \cdot \text{Hg}(\text{ClO}_4)_2$ and $L_1 \cdot \text{Cr}(\text{ClO}_4)_3$ in a

CH_3CN /HEPES buffer (3:2, v/v) medium were evaluated at pH 7.3 using systematic emission titration studies (Figure 5). As mentioned earlier, free L_1 did not show any fluorescence upon excitation at 500 nm, and the new emission band at around 556 nm appeared due to formation of the corresponding xanthene form upon coordination to the Hg^{2+} or Cr^{3+} ion. The respective binding constants were also evaluated. The formation constants for $L_1 \cdot \text{Hg}(\text{ClO}_4)_2$ ($K_{\text{Hg}^{2+\text{Flu}}}$) and $L_1 \cdot \text{Cr}(\text{ClO}_4)_3$ ($K_{\text{Cr}^{3+\text{Flu}}}$) were found to be 3.84×10^3 and $2.1 \times 10^3 \text{ M}^{-1}$, respectively, while the lowest detection limit was 10.72 ppm for Hg^{2+} and 5.6 ppm for Cr^{3+} . These values were close to those calculated from the absorption titration data. For both cations, namely, Hg^{2+} and Cr^{3+} , a 1:1

binding stoichiometry was confirmed from the goodness of the linear fit of the B–H plot (insets in Figure 5), as well as from Job's plot (Supporting Information), using data obtained from either the emission or absorption spectral titration for the respective ions. A higher solvation energy for Cr^{3+} ($H_{\text{hyd}} = 4639 \text{ kJ mol}^{-1}$) in water compared to that of Hg^{2+} ($H_{\text{hyd}} = 1824 \text{ kJ mol}^{-1}$) could have contributed to a weaker binding of Cr^{3+} to the reagent L_1 than that of Hg^{2+} .

As mentioned earlier, the electronic and emission spectral responses were insignificant when either HgCl_2 or HgI_2 , instead of $\text{Hg}(\text{ClO}_4)_2$ or $\text{Hg}(\text{NO}_3)_2$ were used for studies under identical experimental conditions. For HgCl_2 , much weaker absorption (inset in Figure 3a) and emission bands with maxima at around 526 and 552 nm, respectively, were observed. This tends to suggest that the interaction between Hg^{II} in HgX_2 ($\text{X}^- = \text{Cl}^-$ or I^-) and L_1 was too weak to be able to induce the ring-opening reaction to the spirolactam form and convert the reagent L_1 to the corresponding xanthene form. This apparent contradiction was resolved by ^{13}C NMR spectral studies of L_1 in the absence and presence of varying $[\text{Hg}(\text{ClO}_4)_2]$ and HgCl_2 . For the spirolactam ring form, the signal for the tertiary carbon atom (C_{11} in Scheme 1) for free L_1 appeared at 66 ppm. The intensity of this signal at 66 ppm was found to decrease with an increase in $[\text{Hg}(\text{ClO}_4)_2]$ with the simultaneous appearance and growth of a new signal at 116 ppm. This signified the opening of the lactam ring with simultaneous xanthene ring formation of L_1 upon coordination to Hg^{II} of $\text{Hg}(\text{ClO}_4)_2$. No such changes were observed when ^{13}C NMR spectra were recorded in the presence of increasing $[\text{HgCl}_2]$, and the signal for the tertiary carbon atom was retained even in the presence of 1.2 mol equiv of HgCl_2 (Supporting Information). Thus, the results of the ^{13}C NMR studies also corroborate our results of the absorption and emission spectra, which reveal that HgCl_2 fails to induce the lactam ring-opening reaction in L_1 and generation of the xanthene form. Further and more conclusive proof for HgCl_2 in failing to induce the ring-opening reaction for L_1 even after coordination to the Hg^{II} center of HgX_2 was apparent from the single-crystal X-ray structures for $\text{L}_1 \cdot \text{HgX}_2$ ($\text{X}^- = \text{Cl}^-$ or I^- ; Figure 1). HgCl_2 possesses only 23% ionic character, and that for HgI_2 is even less.²² Presumably, this lower ionic character for HgX_2 ($\text{X}^- = \text{Cl}^-$ or I^-) accounts for the lower acidity of the Hg^{II} center and its ability to induce the ring-opening reaction. The binding constant and stoichiometry for L_1 formation were also evaluated based on the small changes that were observed for systematic absorption and emission titration, keeping $[\text{L}_1]$ ($1.128 \times 10^{-5} \text{ M}$) constant and varying $[\text{HgCl}_2]$ between 0 and $1.6 \times 10^{-3} \text{ M}$. Binding constant values that were evaluated from absorption and emission titration data are 90 and 125 M^{-1} , respectively (Supporting Information). In both cases, 1:1 binding stoichiometry was established by the goodness of the linear fit of the B–H plot.

The reversibility of the binding process between L_1 and Hg^{2+} or Cr^{3+} was established when the original spectrum for L_1 was restored upon the addition of either I^- as an aqueous solution of KI to the solution of $\text{L}_1 \cdot \text{Hg}(\text{ClO}_4)_2$ or EDTA^{2-} to the solution of $\text{L}_1 \cdot \text{Cr}(\text{ClO}_4)_3$ in a CH_3CN /aqueous HEPES buffer (3:2, v/v) medium of pH 7.3. I^- and EDTA^{2-} have a strong affinity for Hg^{2+} and Cr^{3+} , respectively, and the respective binding constants are much higher than that for L_1 . This caused demetalation of the reagent L_1 and regeneration of the spirolactam ring with bleaching of the absorption band at $\sim 530 \text{ nm}$ and the emission band at around 555 nm. Thus,

reagent L_1 could be used as a reversible sensor for the recognition of Hg^{2+} and Cr^{3+} in a mixed-solvent medium under physiological conditions (Supporting Information). The reversibility in the binding of L_1 to Hg^{2+} in the presence of an excess of KI could be further extended for use of the reagent L_1 as a selective chemosensor for Cr^{3+} . In the presence of an excess of KI, HgI_2 was formed preferentially, and this caused restoration of the original spectra for L_1 with formation of the cyclic lactam form of the reagent (*vide infra*) L_1 . Again, upon the addition of Cr^{3+} to this solution mixture having KI, the absorbance band at 530 nm and the emission band at 557 nm reappeared (Supporting Information) and the pink color of the solution was also restored. Preferential binding of the iodide ion to the Hg^{2+} ion led to the formation of HgI_2 and the cyclic lactam form of the reagent L_1 . However, the absorption and emission spectral bands, as well as the solution color for the acyclic xanthene form of the reagent, were restored upon coordination of the Cr^{3+} ion to L_1 . Further experiments revealed that the absorbance spectra for $\text{L}_1 \cdot \text{Cr}^{3+}$ and its solution color remained unchanged upon the addition of an excess of the iodide ion in the form of KI. Thus, in the presence of an excess of KI, reagent L_1 could also be used for the selective recognition of Cr^{3+} from all other metal ions.

Weaker binding of L_1 to the Hg^{2+} center in 1a compared to that in $\text{L}_1 \cdot \text{Hg}(\text{ClO}_4)_2$ was also evident from the FTIR spectra recorded for L_1 in the presence of 5 mol equiv of HgCl_2 and $\text{Hg}(\text{ClO}_4)_2$. The stretching frequency for the $\text{C}=\text{O}$ bond for L_1 appeared at 1695 cm^{-1} , while it appeared at 1684 cm^{-1} in the presence of HgCl_2 and at 1647 cm^{-1} in the presence of $\text{Hg}(\text{ClO}_4)_2$ (Supporting Information). A larger shift signifies a higher polarization of the $\text{C}=\text{O}$ bond upon more efficient binding to the Hg^{2+} center in $\text{Hg}(\text{ClO}_4)_2$.

The influence of the binding of Hg^{2+} or Cr^{3+} to the receptor L_1 on the lifetime of its excited state was further checked from the results of the TCSPC studies using a 360 nm nano-LED as the excitation source. Emission decay traces ($\lambda_{\text{Mon}} = 550 \text{ nm}$) for receptor L_1 ($7.76 \times 10^{-6} \text{ M}$) could be best fitted with a biexponential function with time constants $\tau_1 = 0.19 \pm 0.01$ (37.08%) and $\tau_2 = 2.80 \pm 0.07$ (62.09%) with $\chi^2 = 1.241 \text{ ns}$, whereas the same receptor showed a biexponential decay with the time constants $\tau_1 = 0.02 \pm 0.05$ (12.41%) and $\tau_2 = 2.8 \pm 0.05$ (87.59%) having $\chi^2 = 1.02 \text{ ns}$ in the presence of the Hg^{2+} ion, added as $\text{Hg}(\text{ClO}_4)_2$. The minor and short-lived component was assigned to the decay time for the excited state related to the spirolactam moiety, while the long-lived major component was attributed to the xanthene form of the reagent L_1 .^{19b} The relative increase in the percentage of the open-ring xanthene form upon the addition of Hg^{2+} confirms the interaction of Hg^{2+} with the receptor. A similar experiment performed with comparable $[\text{Cr}(\text{ClO}_4)_3]$ showed a biexponential decay trace with time constants $\tau_1 = 0.97 \pm 0.03$ (44%) and $\tau_2 = 4.13 \pm 0.12$ (56%).

A comparison of the absorption spectra for free reagent L and L_1 revealed additional absorption bands at 332 and 347 nm for L_1 . Analogous quinoline derivatives are known to have an absorption band within the range 310–350 nm. Thus, an exact assignment of the absorption band for the quinoline fragment in L_1 is difficult. However, one of these two bands at 332 and 347 nm originates from the quinoline fragment and neither free L_1 nor $\text{L}_1 \cdot \text{Hg}(\text{ClO}_4)_2 / \text{L}_1 \cdot \text{Cr}(\text{ClO}_4)_3$ showed any characteristic emission band for the quinoline fragment even upon excitation at ~ 332 or $\sim 347 \text{ nm}$, where the quinoline fragment is expected to absorb (Supporting Information).^{16a,32} Let us address this

apparently anomalous observation. In the case of L_1 , the flexibility and rapid C=N isomerization of the imine functionality coupled to the quinoline fragment could be responsible for the nonradiative deactivation of the quinoline-based excited state and quenching of the quinoline-based emission for L_1 .^{19c,33} Further, literature reports reveal that the ring nitrogen atom of the quinoline moiety is strongly hydrogen-bonded with water molecules in aqueous or mixed aqueous medium and forms a stoichiometric complex between the excited state of quinoline and a solvent molecule (exciplex),^{19c,34} which could also have contributed to the observed quenching of the quinoline-based emission. For $L_1 \cdot \text{Hg}(\text{ClO}_4)_2$ and $L_1 \cdot \text{Cr}(\text{ClO}_4)_3$, it may be presumed that the efficient spin-orbit coupling for Hg^{2+} and the strong paramagnetic influence of the Cr^{3+} ion could be responsible for quenching of the quinoline-based emission.

Cell Imaging Study. Hg^{2+} is known to exert an estrogenic effect on breast cancer cells by binding to estrogen receptors and causing increased cell growth. Among heavy metal ions, chromium is found to be in higher amounts in human breast cancer tumor samples than in benign breast tissue.³⁵ These reports relate these metal ions to malignancy and led us to check the possibility of using the reagent L_1 as an imaging agent for the detection of these toxic metal ions in human breast cancer cells.

The L_1 -stained breast cancer cell MCF7 treated with either Cr^{3+} or Hg^{2+} was visualized under the laser scanning confocal microscope FV1000 (Olympus), as mentioned in the Materials and Methods section. MCF7 cells treated with different concentrations of Cr^{3+} (Figure 6A) or Hg^{2+} (Figure 6B)

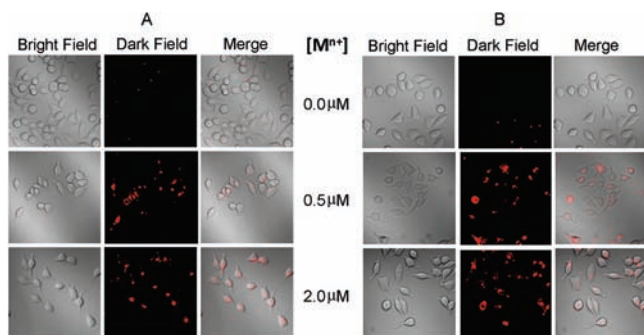


Figure 6. Confocal images of (A) Cr^{3+} - and (B) Hg^{2+} -treated MCF7 cells. The cells were supplemented with different concentrations of $\text{Cr}(\text{ClO}_4)_3$ or $\text{Hg}(\text{ClO}_4)_2$ (0.0 μM ; A, 0.5 μM ; B, 2.0 μM) in the growth media for 1.0 h at 37 °C followed by staining with 4.0 μM L_1 for 1.0 h at 37 °C ($\lambda_{\text{ext}} = 543 \text{ nm}$).

showed brighter fluorescence in comparison to untreated cells with no or very little fluorescence. The fluorescence was mostly localized in the cytoplasm of the cells. The results also indicate that L_1 could detect Cr^{3+} or Hg^{2+} at the 0.5 μM level. The reversibility of the binding of the reagent L_1 to Hg^{2+} or Cr^{3+} ions present in MCF7 cells could also be established by recording confocal images of the stained cells in the presence of an excess (5 mol equiv with respect to L_1) of KI [for $\text{Hg}(\text{ClO}_4)_2$] or Na_2EDTA [for $\text{Cr}(\text{ClO}_4)_3$]; a substantial decrease in fluorescence of the MCF7 cells, previously treated with the respective metal ions and the reagent L_1 , upon exposure to iodide or EDTA^{2-} ions was observed (Supporting Information).

Further, the cytotoxicity of L_1 on MCF7 cells was determined by a conventional MTT assay (Supporting Information),³⁶ which revealed that, upon exposure to a 5 μM concentration of L_1 (a concentration that was comparable to that used for confocal imaging studies; Figure 6) for 6 h, ~87% of the MCF7 cells remained viable. This nullified the possibility of any significant cytotoxic influence of the reagent L_1 on MCF7 cells. Therefore, it may be concluded that L_1 could be used as a viable chemosensor for Cr^{3+} or Hg^{2+} in biological samples.

In brief, we have reported a new rhodamine-based reagent (L_1) that showed remarkable preference toward Hg^{2+} and Cr^{3+} . Upon binding to either of these two ions, a visually detectable change in the color and emission were observed because of conversion of the lactam form of the rhodamine derivative to the acyclic xanthene form. This offered the possibility of using this reagent for use either as a colorimetric or as a fluorescence-based reagent for the detection of larger fluctuations of these ions, especially in the case of Hg^{2+} , which binds at a micromolar level under physiological conditions. Further studies revealed that conversion from the cyclic lactam form to the acyclic xanthene form was reversible, and thus this reagent could be used as a reversible sensor for the detection of Hg^{2+} or Cr^{3+} . More importantly, reagent L_1 could be used as an imaging agent for detection of these ions, which are known for their deleterious influence in cell malignancy, in breast cancer cell MCF7. More importantly, this is the first report on the single-crystal structure for HgX_2 ($X = \text{Cl}$ or I) bound to a lactam form of the rhodamine derivative and explicitly explains the inability of these lactam reagents to recognize mercury halides through detectable changes in optical or fluorescence spectra.

■ ASSOCIATED CONTENT

📄 Supporting Information

Characterization data of the receptor L_1 , spectral crystallographic details, biological studies, and X-ray crystallographic data in CIF format. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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