Fluorescence “Turn-off” Sensing of Iron (III) Ions Utilizing Pyrazoline Based Sensor: Experimental and Computational Study

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Received: 13 June 2022 / Accepted: 2 September 2022 / Published online: 22 September 2022
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Abstract
A simple pyrazoline-based ‘‘turn off’’ fluorescent sensor 5-(4-methoxyphenyl)-3-(5-methylfuran-2-yl)-1-phenyl-4,5-dihydro-1H-pyrazole (PFM) was synthesized and well characterized by different techniques such as FT-IR, 1H-NMR, 13C-NMR, and mass spectrometry. The synthesized sensor PFM was utilized for the detection of Fe3+ ions. Fluorescence emission selectively quenched by Fe3+ ions compared to other metal ions (Mn2+, Al3+, Fe2+, Hg2+, Cu2+, Co2+, Ni2+, Cd2+, Pb2+, and Zn2+) via paramagnetic fluorescence quenching and showed good anti-interference ability over the existence of other tested metals. Under optimum conditions, the fluorescence intensity of sensor quenched by Fe3+ in the range of 0 to 3 μM with detection limit of 0.12 μM. Binding of Fe3+ ions to PFM solution were studied by fluorescent titration, revealed formation of 1:1 PFM-Fe metal complex and binding constant of complex was found to be of 1.3 × 105 M−1. Further, the fluorescent sensor has been potentially used for the detection of Fe3+ in environmental samples (river water, tap water, and sewage waste water) with satisfactory recovery values of 99–101%.

Keywords Chalcone · Pyrazoline · Fluorescence quenching · Iron · DFT

Introduction
Iron, as one of the most fundamental trace elements, is known for cell formation in the human body system and growth in plants. Ferric ions (Fe3+) play an imperative role in lots of critical biological processes (e.g., oxygen transfer in haemoglobin, electron-proton transfer, RNA and DNA synthesis, nerve conduction, enzyme synthesis, and regulation of acid–base balance). However, excess deposition of iron in the human body can cause some severe diseases including hemochromatosis, Parkinson’s and Alzheimer’s disease, and diabetes [1]. Moreover, the lack of iron in the human body decreases immunity throughout the developmental periods. Subsequently, a trace level of iron plays a vital role in the health of the living organism. Considering importance of evaluation of Fe3+ ions concentration, WHO and European legislation have set the permissible limit of iron in drinking water and food as 0.3 ppm (~5.4 μM) and 0.2 ppm (~3.8 μM) respectively [2]. Thus, quantitative detection of Fe3+ ion at an ultra-trace level in environmental samples is of prominent concern using efficient analytical methods. Numerous methods (e.g. atomic absorption spectrometry, spectrophotometry, colorimetry, high-performance liquid chromatography, inductively coupled plasma optical emission spectrometry, and electrochemical analysis) for the detection of Fe3+ ions have been used with good aspects [3, 4]. However, all these methods emerged with various limitations such as complicated technologies, sophisticated handling, expensive devices, and a long time for operating systems. Therefore, compared with these methods, the fluorescence sensing strategy has gained considerable attention, as it has protruded optimistic method for monitoring and detection of diverse ions in environmental, and biological samples [5]. This consideration is due to a lot of highlights...
like simplicity, cost-effectiveness, rapid responses, high selectivity, and sensitivity [6].

Pyrazolines derivatives, as the most recent fluorescent sensor have generated much excitement because of their multipurpose applicability in various fields compared with other fluorescent emitters [7, 8]. The synthetic versatility and the extended synthesis potential of pyrazoline with intrinsic biological and pharmacological activity (e.g. antimalarial, anti-inflammatory, antibacterial) have made pyrazolines and its derivatives as one of the most well-known precursors to fungal, anti-inflammatory, antibacterial) have made pyrazolines and its derivatives as one of the most well-known precursors to chemistry [9, 10]. Specifically, the assurance of spectroscopic properties of pyrazoline dyes widely used as pH sensors, metal ion fluorescent sensor, living cell imaging probes, and logic-based devices is of great importance [11, 12]. Now a days pyrazolines fluorescence sensors are also contributing in the direction of various other specifications like reusability, reversibility, good quantum yield and suitable fluorescence life time in addition to high sensitivity and selectivity [13–16].

In this context, and in continuation of our research on pyrazoline based fluorescent sensors [17], we synthesized and characterized a “turn Off” fluorescent sensor, namely, 5-(4-methoxyphenyl)-3-(5-methylfuran-2-yl)-1-phenyl-4,5-dihydro-1H-pyrazole (PFM). This molecule, detects Fe$^{3+}$ over other metal cations (Al$^{3+}$, Fe$^{3+}$, Fe$^{2+}$, Mn$^{2+}$, Cu$^{2+}$, Co$^{2+}$, Hg$^{2+}$, Ni$^{2+}$, Cd$^{2+}$, Pb$^{2+}$, and Zn$^{2+}$). Besides, the DFT calculations were used to confirm the experimental results.

### Experimental

#### Reagents and Measurements

All the metal salts and 4-methoxybenzaldehyde were purchased from Loba Chemie Pvt. Ltd. (Mumbai, India). 2-acetyl-5-methylfuran and phenyl-hydrazine were purchased from Sigma-Aldrich (Mumbai, India). The synthesized compounds were characterized by FT-IR, $^1$H-NMR, $^{13}$C-NMR, and mass spectrometry. FT-IR spectra were scanned on a Perkin Elmer Spectrum Infrared Spectrophotometer Version (10.6.0), Japan. $^1$H-NMR and $^{13}$C-NMR spectra were recorded on a 500 MHz Bruker spectrometer, Switzerland. LC–MS Spectrometer Model Q-ToF Micro Waters was used to record the mass spectrum of the PFM. To record absorption spectra of the compounds, UV-1800 Shimadzu UV–Visible spectrophotometer (Shimadzu, Japan) was used. All fluorescence experiments were performed with a Shimadzu RF-5301PC spectrophotofluorometer, (Shimadzu, Japan). Triply distilled water (TDW) was used for the experimental work.

#### Synthesis and Characterization of 5-(4-methoxyphenyl)-3-(5-methylfuran-2-yl)-1-phenyl-4,5-dihydro-1H-pyrazole (PFM)

The synthesis of pyrazoline derivative PFM is presented in Scheme 1. The starting material chalcone (CFM) was synthesized from 2-acetyl-5-methylfuran and 4-methoxybenzaldehyde as reported in the literature [18]. Then, a mixture of chalcone CFM (2.42 g, 0.01 mol), phenylhydrazine (1.5 g, 0.01 mol), KOH (0.5 g, 0.01 mol), and ethanol (20.0 ml) was refluxed continuously for 6 h and reaction progress was monitored by TLC. After reaction completion, the resulting mixture was neutralized with iced-HCl to yield a dark brown solid mass. Then, the product PFM was recrystallized from MeOH [19].

Dark Brown solid; Yield: 72%; m.p. 179-180°C; FT-IR: $\nu_{\text{max}}$ (cm$^{-1}$): 3100 (aromatic C-H) & 1595 (C=N); $^1$H-NMR (500 MHz, CDCl$_3$): $\delta$ 7.17 (4H, m), 7.03 (2H, m), 6.80 (2H, d), 6.73 (1H, d), 6.37 (1H, d), 6.00 (1H, d), 5.11 (1H, dd, $J_{\text{AX}}$ = 12 Hz, $J_{\text{AX}}$ = 12 Hz, H–X), 2.97 (1H, dd, $J_{\text{AX}}$ = 7 Hz, $J_{\text{AM}}$ = 17 Hz, H–A), 2.33 (3H, s, 1''-CH$_3$); $^{13}$C-NMR (125 MHz, CDCl$_3$): $\delta$ 159.04, 153.86, 146.63, 144.85, 139.36, 134.40, 128.89, 127.12, 119.04, 114.53, 113.58, 111.013, 107.96, 63.4, 55.27, 43.27, 13.91; Calculated ESI–MS: m/z 332.39 for C$_{21}$H$_{20}$N$_2$O$_3$.

#### Analytical Procedure

Stock solutions (1 µM) of metal salts i.e. AlCl$_3$, FeCl$_3$,6H$_2$O, FeCl$_2$H$_2$O, MnCl$_2$, CuCl$_2$.2H$_2$O, Co(NO$_3$)$_2$, HgCl$_2$, NiCl$_2$, Cd(NO$_3$)$_2$, Pb(NO$_3$)$_2$, and ZnCl$_2$ were prepared in triply distilled water. The stock solution of probe PFM (2 x 10$^{-5}$ M) in methanol: water (1:9, v/v) was prepared.
Investigated the fluorescent behavior of the sensor PFM, the fluorescence excitation and emission wavelength were found to be 350 nm and 484 nm respectively. Afterwards, for selectivity, the absorption and fluorescence spectrum of PFM in the presence of each metal salt was recorded. For the experiment, 100 µL of PFM solution, 100 µL of metal salt was taken in cuvette and then diluted up to 3 ml with triply distilled water. For the better considerations of the quenching behavior of sensor PFM, fluorescence titration was performed in the presence of different concentrations of Fe$$^{3+}$$ ions (0–3 µM). The limit of detection (LOD) value was obtained from the 3σ/K (where σ is the standard deviation of the blank solution and K represents the slope of the calibration curve between fluorescent intensity and the Fe$$^{3+}$$ concentrations).

**Binding Measurement’s**

Job’s plot analysis was carried out to identify the binding stoichiometry between sensor PFM and Fe$$^{3+}$$ ions. The solution of sensor PFM and Fe$$^{3+}$$ ions were prepared to carry out Job’s plot experiments. The plot was constructed from the emission profile by maintaining the sum of the concentration of Fe$$^{3+}$$ ions and the PFM constant. The fluorescence spectrum was recorded by varying the mole fraction of PFM and Fe$$^{3+}$$ ions at an excitation wavelength of 350 nm. The emission intensity was plotted against the mole fraction of the Fe$$^{3+}$$ ions. The molar ratio corresponding to the highest point or intensity was plotted against the mole fraction of the ions at an excitation wavelength of 350 nm. The fluorescence spectrum was recorded by varying the mole fraction of PFM and Fe$$^{3+}$$ ions (0–3 µM). The limit of detection (LOD) value was obtained from the 3σ/K (where σ is the standard deviation of the blank solution and K represents the slope of the calibration curve between fluorescent intensity and the Fe$$^{3+}$$ concentrations).

The association constant of sensor PFM with Fe$$^{3+}$$ ions was calculated according to the fluorescence emission intensity data using the modified Benesi–Hildebrand equation:

$$\frac{F_{0} - F_{\text{min}}}{F - F_{\text{min}}} = \frac{1}{K_{a} [M]} + 1$$

where, $$F_{0}$$, $$F$$, and $$F_{\text{min}}$$ are the fluorescence intensities of PFM in the absence of Fe$$^{3+}$$ ions, at an intermediate concentration of Fe$$^{3+}$$ ions, and a concentration of complete interaction of Fe$$^{3+}$$ ions, respectively. $$K_{a}$$ is the association constant and [M] represents the concentration of the metal ion (Fe$$^{3+}$$) [21, 22].

**Computational Study**

To gain insight into the structures and fluorescence properties of PFM, before and after the addition of metals, density functional theory (DFT) computations were performed on B3LYP/6-311G(d,p)/LANL2DZ using Gaussian 09 software. The optimized geometrical parameters, net charges on active centres, and energetic of the ground state for intermediate chalcone CFM, the sensor (PFM), and its binding with iron metal (PFM-Fe) were calculated. The spectral theoretical results of vibration analysis, $^{13}$C-NMR, and $^{1}$H-NMR of the ligand molecule were also examined.

**Detection of Iron in Water Samples**

Different water samples (river water, tap water, and sewage waste water) were used for the practical applicability of synthesized sensor. The river water was collected from the Ghaggar River (Pathiala, Punjab, India). The tap water was collected from the chemistry lab (Khalsa College, Pathiala) and sewage waste water was taken from the Punjabi University, Pathiala (Punjab, India). All the water samples were filtered through a Grade 1 Whatman filter paper (pore size: 11 µm) and nylon-6,6 membrane filter (0.2 µm per 47 mm) before analysis. All the samples were tested with the proposed method before spiking.

**Results and Discussion**

**Synthesis and Structural Characterizations**

The absence of stretching frequency of α, β-unsaturated carbonyl group and the presence of (C=N) stretching frequencies at 1595 cm$^{-1}$ and 1244 cm$^{-1}$ in the IR-spectrum (Figs. S1 and S2) of the sensor PFM confirmed the subsequent cyclization of chalcone to form the pyrazoline derivative PFM [23]. In the 500 MHz instruments, the $^{1}$H-NMR coupling constant analysis of compound CFM indicated that hydrogen atoms of the olefinic carbon–carbon bond were in a trans arrangement ($J = 15$ Hz) (Fig. S3). $^{1}$H-NMR spectra of the compound PFM exhibit the presence of two non-equivalent protons of a methylene group ($H_{A}/H_{M}$) at 2.97 ppm and 3.64 ppm, because of the (H–X) proton at vicinal asymmetric carbon. The methene proton (H–X) appeared as a doublet of doublets at 5.11 ppm, because of vicinal coupling with the two magnetically non-equivalent protons of the methylene group at position 4 of the pyrazoline ring (Fig. S4) [23]. The carbonyl carbon of the chalcone CFM appeared at 177.39 ppm (Fig. S5). A signal due to C=N carbon of the pyrazoline ring was observed in PFM at 159.04 ppm. C4 and C5 carbons of the pyrazoline ring resonated at 63.41 ppm and 43.27 ppm respectively (Fig. S6) [24]. The characteristic peaks of masses were observed at m/z 243.44 (Fig. S7) and m/z 332.39 in the mass spectra of chalcone CFM and ligand PFM (Fig. S8).

**Absorption and Fluorescence Experiments of PFM for the Sensing of Fe$$^{3+}$$ Ion**

UV–Vis absorption and fluorescence spectra of compound PFM were presented together in Fig. 1a to determine the
optical properties of the compound PFM in methanol:water (1:9). The PFM exhibited absorption bands at 286 nm and 366 nm and relatively high intensity fluorescence band at 484 nm. The n–π* transition may be due to conjugation between a lone pair electron of the nitrogen atom in the pyrazoline moiety and the π–bond of the benzene ring.

The effects of several metal ions having biological and ecological importance (Al³⁺, Fe³⁺, Fe²⁺, Mn²⁺, Cu²⁺, Co²⁺, Hg²⁺, Ni²⁺, Cd²⁺, Pb²⁺, and Zn²⁺) on the absorption properties of the compound PFM were investigated. For this purpose, solutions having selected metal ions were added into the compound PFM’s 2×10⁻⁵ M aqueous solution. Then absorption measurements of metal ions were realized and recorded in Fig. 1b. Upon addition of metal ions to the solution containing PFM, the band observed at 286 nm diminished while the absorption of the band at 366 nm is enhanced. These changes can be manifest due to charge transfer properties of the pyrazoline moiety [25]. However, in the case of Fe³⁺ ions, the absorption spectra exposed significant changes. A bathochromic shift for band at 366 nm, and a new band at 625 nm was identified. The changes in band position and intensity of several characteristic absorbing peaks in the UV–Vis absorption spectrum of PFM-Fe can be attributed to n→π* and π→π* transitions of the aromatic π system containing the C=N bonds[26–28].

Further, to gain insight for the selectivity of sensor PFM, fluorescence measurements were done. Different excitation wavelengths (320–400 nm) were optimized to get the maximum fluorescence emission intensity for PFM. The maximum fluorescence emission intensity (484 nm) was observed at excitation wavelength of 350 nm (Fig. S9). The selective recognition of different metal ions having concentration 1 μM (Al³⁺, Fe³⁺, Fe²⁺, Mn²⁺, Cu²⁺, Co²⁺, Hg²⁺, Ni²⁺, Cd²⁺, Pb²⁺, and Zn²⁺) was examined (λex=350 nm; λem=484 nm). For the selectivity experiment, 100 μL of different metal solution and 100 μL of PFM were taken and diluted to 3 mL with TDW. The solution was stirred for 3 min and kept undisturbed at room temperature for 15 min. A large decrease in fluorescence intensity was observed for Fe³⁺ comparative to other metal cations, showing a selective recognition of Fe³⁺ ion by pyrazoline-based ligand PFM (Fig. 2).

**Effect of Response Time**

The effect of the response time of the PFM sensor in the presence of Fe³⁺ was investigated at different times (1–20 min). As shown in Fig. S10, the fluorescence intensity of PFM sensor with Fe³⁺ reached equilibrium within 1 min and then almost no change in the fluorescence intensity within 20 min was observed. It showed that PFM quickly coordinates with Fe³⁺ ions in almost 1 min. The rapid and stable response of the sensor PFM could serve as an efficient and consistent probe for Fe³⁺ ions.

**Fluorescence Life Time and Quantum Yield**

The average lifetime of the PFM sensor in the absence and presence of Fe³⁺ was found to be 1.94×10⁻⁹ s and 1.55×10⁻¹⁰ s respectively. Figure S11 represents the

![Fluorescence spectra of PFM (2×10⁻⁵ M) in methanol:water (1:9) upon addition of various metal ions (1 μM) (Ex. Wavelength 350 nm)](image)
fluorescence lifetime decay curve of PFM in the absence and presence of Fe\(^{3+}\) fitted with three-component exponential decay curve.

The fluorescence quantum yield of the sensor was calculated using quinine sulphate in 0.5 M H\(_2\)SO\(_4\) as reference standard. The fluorescence quantum yield of PFM sensor in the absence of Fe\(^{3+}\) was measured as 0.69. However, upon the addition of Fe\(^{3+}\), the quantum yield decreases and measured as 0.27.

**Reusability and Reversibility**

To endorse the reversible and reusability performance of the sensor PFM towards Fe\(^{3+}\) ions, the effect of addition of EDTA on the fluorescence was examined. When EDTA was added in to the solution, the fluorescence intensity of compound PFM returned to the earlier state (93%) in which there was no Fe\(^{3+}\) ions. This indicates that compound PFM fluorescence sensor system detecting Fe\(^{3+}\) ions is reversible and can be reused for further findings. The fluorescence response of subsequent additions of Fe\(^{3+}\) or EDTA on PFM was carried out. On adding Fe\(^{3+}\), the fluorescence intensity of PFM decreases while addition of EDTA on PFM increased the fluorescence intensity. These cycles were repeated five times without losing much sensitivity of the sensor (Fig. S12).

**Sensitivity Study of PFM for Fe\(^{3+}\) Ion**

To quantify the sensitivity and fluorescence quenching behavior between PFM and Fe\(^{3+}\), the sensing proficiency of PFM towards Fe\(^{3+}\) was further explored in the range of 0–3 \(\mu\)M. On the addition of Fe\(^{3+}\) ions in a sequential manner in the PFM solution, the fluorescence emission intensity of the sensor PFM gradually decreases with the increase in concentration of Fe\(^{3+}\) (Fig. 3a). The emission intensity of sensor PFM was specifically quenched by Fe\(^{3+}\) ions via paramagnetic fluorescence quenching [5, 29]. The quenching phenomena were further analyzed by the Stern–Volmer equation:

\[
\frac{F_0}{F} = 1 + K_{SV}\left[Fe^{3+}\right]
\]

where, \(F_0\) is the initial fluorescence intensity of the PFM solution in the absence of Fe\(^{3+}\), \(F\) is the fluorescence emission intensity in the presence of Fe\(^{3+}\), and \(K_{SV}\) is the Stern–Volmer constant. The Stern–Volmer plot (\(F_0/F\) versus \([Fe^{3+}]\)) depicts that the quenching ratio increase linearly with the increase in Fe\(^{3+}\) concentration (\(R^2 = 0.99\)) (Fig. 3b). The \(K_{SV}\) was calculated from the slope of plot and found to be 4.45 \(\times\) 10\(^{-5}\) M\(^{-1}\).

The limit of detection (LOD) was calculated 0.12 \(\mu\)M for Fe\(^{3+}\) using the equation \(LOD = 3\sigma/K\), which are far lower than most extreme toxin levels for Fe\(^{3+}\) (5.4 \(\mu\)M) in drinking water given by EPA guidelines (Fig. 3c) [2].

**Competitive Selectivity of PFM for Fe\(^{3+}\) Ions**

To investigate the selectivity and efficiency of PFM towards Fe\(^{3+}\) ions, competitive experiment was carried out. For the
competitive study, 1 mL of Fe³⁺ (1 μM) was added to the 100 μL of PFM (2 × 10⁻⁵ M) solution containing other metal ions (Al³⁺, Fe²⁺, Mn²⁺, Cu²⁺, Co²⁺, Hg²⁺, Ni²⁺, Cd²⁺, Pb²⁺, and Zn²⁺ at concentration of 2 × 10⁻⁶ M) were taken and the fluorescence emission spectra was obtained. The interfering metal ions induced no significant changes in the fluorescence intensity of the sensor PFM (Fig. 4). As a result, the PFM can be presented as a highly selective and reliable fluorescent sensor for Fe³⁺ ion recognition [25]. Moreover, relative error (%) for various metal ions was calculated:

\[
\text{Relative error} (\%) = \left( \frac{F - F_0}{F_0} \right) \times 100\%
\]

where, F₀ and F are the fluorescence emission intensities in the absence and presence of interfering ion. Table 1 validated the relative error (%) values showing great tolerance of other metals over the Fe³⁺ ion. The relative error is also found to be less than ±5. These results suggest that the metal binding of PFM shows an evident preference for ferric ions over other competing ions.

**Proposed Binding and Sensing Mechanism**

A Job’s Plot was performed to calculate the binding stoichiometry of PFM and Fe³⁺ ion (Fig. 5). A turning point at 0.5 mol fractions indicates 1:1 metal–ligand binding interactions between sensor PFM and Fe³⁺. The association constant (K_a) was calculated to be 1.3 × 10⁵ M⁻¹ according to the modified Benesi–Hildebrand equation (Fig. 6).

The mechanism of interaction between the Fe³⁺ and PFM was studied through FT-IR (Fig. S13). It can be proposed that the formation of the coordination between Fe³⁺ and sensor PFM resulted from electronegative atom nitrogen of the pyrazoline ring and the oxygen of the furyl ring (Fig. 7). This interaction causes the fluorescence quenching of the sensor PFM. The FTIR spectrum of PFM exhibits a peak at 1595 cm⁻¹ (stretching vibration C=N), and peak at 1245 cm⁻¹ (C-O stretching vibration). However, in the PFM-Fe complex, the characteristic stretching vibration C-O peak of PFM partially disappear while the C = N stretching vibration of pyrazoline ring shifted from to 1595 cm⁻¹ to 1561 cm⁻¹. This can be due to the interaction of Fe³⁺ ions with the nitrogen and oxygen atom of PFM. The binding of Fe³⁺ to PFM resulted in the electron or energy transfer from PFM to Fe³⁺ metal ion causes fluorescence quenching of

\[
\begin{array}{|c|c|}
\hline
\text{Interferent ion} & \text{Relative Error} \% (\Delta F/F_0 \times 100) \\
\hline
\text{Zn}^{2+} & 1.411 \\
\text{Hg}^{2+} & -1.411 \\
\text{Cd}^{2+} & -1.376 \\
\text{Fe}^{2+} & 0.882 \\
\text{Mn}^{2+} & 4.4107 \\
\text{Co}^{2+} & 2.681 \\
\text{Cu}^{2+} & -3.0345 \\
\text{Ni}^{2+} & -3.493 \\
\text{Pb}^{2+} & 4.234 \\
\text{Al}^{3+} & 4.869 \\
\hline
\end{array}
\]

**Fig. 4** a Fluorescence response of PFM upon addition of Fe³⁺ ion in the presence of other competing metal ions (Ex. Wavelength 350 nm). b Dark grey bars represent the fluorescence intensity of PFM in the presence of 1 μM of metal ion, light grey bars represent the fluorescence intensity in the presence of various metal ions after the addition of Fe³⁺.
PFM. The fluorescence emission spectra of PFM overlaps with the absorption spectra of Fe$^{3+}$, suggesting the fluorescence resonance energy transfer (FRET) mechanism for the fluorescence quenching (Fig. S14).

**Computational Study**

**Molecular Geometry Optimization**

The optimized structures of these compounds along with the labeling of atoms are shown in Fig. 8. After optimization, the binding energies ($\Delta E = E($complex$) - E($PFM$)$) were calculated for all metals with PFM to obtain the strongest binding of the metal cation with PFM. The calculated result, $\Delta E = -564$ kcal mol$^{-1}$ showed the minimum energy changes for iron complex as compared to other metals (Table S1). Also, more negative energy value (-1194.45 a.u.) of the PFM-Fe than the free ligand (-1072.42 a.u.) confirms the stability of the PFM-Fe system compared to other metals showing high selectivity towards iron ions [5, 30, 31].

Also, for chalcone CFM, ligand PFM, and ligand complex (PFM-Fe$^{3+}$) with iron metal, C–C bond distances are found to be in the range from 1.528–1.533 Å, 1.529–1.539 Å, and 1.531–1.546 Å while for C-N, these values are 1.469 Å, 1.470 Å, and 1.478 Å respectively [30].

**Mulliken Population Analysis and Molecular Electrostatic Potential**

The Mulliken population analysis is correlated to the vibrational properties and nature of chemical bonds present in the molecule. The Mulliken charge distribution structure and horizontal bar diagram of comparative mulliken atomic charges of the title compounds are shown in Fig. 9a, b, respectively. All the hydrogen atoms in the compounds carry a net positive charge. The atomic charge distribution shows that the hydrogen atoms of the methoxy group have a bigger positive atomic charge (0.2e to 0.32e) than the other hydrogen atoms. As expected, the charge of the nitrogen atom (N31 $=-0.1826$, -0.2933 and N32 $=-0.2574$, -0.2008 in PFM and PFM-Fe respectively) is negative. Additionally, the results illustrate that the charge of the oxygen atoms in the carbonyl group of CFM and furyl ring exhibits a negative charge, which acts as donor atoms. The oxygen atom of -OCH$_3$ group enforces a large negative charge on the carbon (C11 $=-0.1285$, C26 $=-0.3822$, C26 $=-0.4545$ in CFM, PFM, and PFM-Fe respectively) attached to it. Iron atom (Fe46 $=0.7185$) has a high positive charge showing electropositive character.

The electrostatic potential surfaces are correlated with the charge density, shape, dipole moment, and position of chemical reactivity of the molecules. As inspected from the MEPs map of the title compounds (Fig. 10), the negative regions are localized over the electronegative oxygen and nitrogen atoms. The maximum positive regions are localized on the hydrogen atoms and the metal ion [32].

**Molecular Reactivity**

The electron donor–acceptor properties of various types of molecules can be defined by using the energy of HOMO and LUMO. FMO’s also helped to interpret the kinetic stability, charge transfer, and chemical reactivity of a molecule. The frontier molecular orbital distribution of the compounds CFM, PFM, and PFM-Fe were represented in Fig. 11a. The smaller value of the HOMO and LUMO energy gap showed that the studied molecule has high polarizability, chemical reactivity, and biological activity.
Fig. 8  Optimized geometric structures of CFM, PFM, and PFM-Fe at B3LYP/6-311G (d,p)
In the PFM molecule, the HOMO (–6.621 eV) and the LUMO (–2.506 eV) are situated at the benzene and pyrazoline ring, respectively. For the PFM-Fe, the electron density of HOMO (–16.079 eV) is mainly situated at the molecular framework. Whereas, the electron density of LUMO (–14.062 eV) is situated at the coordination.
center. The calculated energy gap ($\Delta$) between HOMO and LUMO for PFM-Fe is found to be 2.02 eV, which is lower than that of unbound PFM (4.12 eV). Based on the DFT calculations, the binding between PFM and Fe is energetically favorable. These DFT results implied that the interaction of Fe to PFM effectively decreases the HOMO–LUMO energy gap of the PFM-Fe and intensively stabilizes the sensing of Fe by forming PFM-Fe compound [33]. Furthermore, the chemical reactivity parameters of the compounds (Table 2) were also calculated with the help of the energy of HOMO and LUMO orbitals. Using FMOs energies, the ionization potential (I) and electron affinity (A) can be measured as: $I = -E_{\text{HOMO}}$ and $A = -E_{\text{LUMO}}$.

Moreover, Density of States (DOS) plots of PFM are examined to study the electronic structure of the molecule via population analysis of orbitals. DOS plot represents the energy level of each orbital. DOS plot (Fig. 11b) for PFM shows that FMOs and the energy gap of HOMO–LUMO are in complete agreement with the result obtained from the DFT study [34]. To investigate the optimization process Fig. S15a, a graph displaying deviation from the target was plotted (Fig. S15b). Also, the graph energy vs. optimization step was plotted as shown in Fig. S15c. In these plots, the path of structure convergence was established. As inspected from these plots, the self-consistent field is converging as the line is directed towards zero [35].

Note: Some other important thermodynamics parameters, NBO, Vibrational analysis, and NMR of the compounds based on theoretical results are discussed in Sect. 1 of the SI.

**Detection of Fe$^{3+}$ in Water Samples**

The synthesized PFM sensor was utilized for the detection of Fe$^{3+}$ in different water samples (river water, tap water, and sewage waste water) by using the spike recovery method. All the samples were tested before spiking and the results revealed that the concentrations of Fe$^{3+}$ were either below the LOD or not present in the tested samples. So, all the samples were spiked with different concentrations of Fe$^{3+}$ and the results obtained were shown in Table 3. It can be seen that the detected Fe$^{3+}$ ions concentration is close to the spiked value with a satisfactory recovery value (99.0–101.0%) and low RSD values (0.78–2.03%). This signifies the great practical potential of the present method for the detection of Fe$^{3+}$ in water samples.
Comparison with Other Reported Fe³⁺ Fluorescent Sensors

To provide quantifiable achievement for the design of a new fluorescent sensor, the sensing performance of PFM was compared with some reported pyrazoline based fluorescent sensors for Fe³⁺ detection (Table 4). As shown in Table 4, the synthesized fluorescent ligand PFM showed a better detection limit for Fe³⁺ (0.12 μM) in comparison to other reported sensors. Pyrazoline based fluorescent sensors such as P [36], Q [37], R [38], and S [39] showed good selectivity for Fe³⁺ ions, but were not used for any real sample analysis. Pyrazole-pyrazoline based sensor (U) [40] showed a better detection limit but the synthetic process was tedious and involved 4 step reaction. On the other hand, the present procedure involves a simple procedure with 2 step reaction (Scheme 1). Moreover, the sensor showed better extraction recovery of 99–101% (tap, river, and sewage waste water) with RSD <2.1% than the other reported methods.

### Table 2 Chemical Reactivity Parameters of the Compounds Based on HOMO–LUMO

| S. No | CFM     | PFM     | PFM-FE   |
|-------|---------|---------|----------|
| E\textsubscript{HOMO} (eV)  | -7.308  | -6.6216 | -16.0792 |
| E\textsubscript{LUMO} (eV)  | -2.133  | -2.5064 | -14.0623 |
| ΔE\textsubscript{HOMO-LUMO} (eV) | 5.1747  | 4.11517 | 2.0169   |
| Ionization potential (IP) (eV) | 7.308   | 6.6216  | 16.0792  |
| Electron affinity (EA) (eV)  | 2.133   | 2.5064  | 14.0623  |
| Chemical potential (eV)      | -4.7207 | -4.564  | -15.0707 |
| Electronegativity(eV)        | 4.7207  | 4.564   | 15.0707  |
| Chemical hardness (eV)⁻¹     | 2.5873  | 2.0575  | 1.008454 |
| Chemical Softness (eV)⁻¹     | 0.1932  | 0.243   | 0.495808 |
| Electrophilic global index (eV) | 4.3065  | 5.0618  | 112.611  |

### Table 3 Detection of Fe³⁺ in different water samples (tap, river, and sewage waste water) by the proposed PFM sensor

| Order | Matrix          | Amount spiked (μM) | Amount found (μM) | Recovery % | RSD % |
|-------|-----------------|--------------------|-------------------|------------|-------|
| 1     | River water     | 0.89               | 0.878             | 98.6       | 0.85  |
|       |                 | 1.79               | 1.77              | 98.8       | 0.91  |
|       |                 | 2.50               | 2.43              | 97.2       | 0.78  |
| 2     | Tap water       | 0.89               | 0.884             | 99.3       | 1.23  |
|       |                 | 1.79               | 1.79              | 100.0      | 1.11  |
|       |                 | 2.50               | 2.48              | 99.2       | 2.03  |
| 3     | Sewage waste water | 0.89             | 0.88              | 98.8       | 0.59  |
|       |                 | 1.79               | 1.79              | 100.0      | 0.96  |
|       |                 | 2.50               | 2.50              | 100.0      | 1.20  |
Conclusions

The synthesized organic fluorescent probe based on pyrazoline 5-(4-methoxyphenyl)-3-(5-methylfuran-2-yl)-1-phenyl-4,5-dihydro-1H-pyrazole (PFM) has been designed for the selective detection of Fe³⁺ ions. The sensor PFM displayed a “turn off” fluorescence response towards Fe³⁺ ion with a detection limit of 0.12 µM. The binding stoichiometry of Fe³⁺ ions with PFM was 1:1 confirmed by Job’s plot, and their binding mechanism of them was demonstrated by paramagnetic enhanced quenching, FRET and density functional theory (DFT) study. The proposed PFM-sensor was satisfactory applied for quantitative and cost-effective detection of Fe³⁺ with good precision in real samples.

Supplementary Information The online version contains supplementary material available at https://doi.org/10.1007/s10895-022-03024-y.

Acknowledgements The authors (PS, SB, IM, MY, and AKM) acknowledge the UGC-SAP and Chemistry Department, Punjabi University, Patiala, for providing lab and instrument facilities.

Authors’ Contributions Promila Sharma: Performed the experimentation. Shikha Bhogal, Irshad Mohiuddin: Helped in performing the experiments. Mohammad Yusuf and Ashok Kumar Malik: Helped in writing and supervised the research work.

Funding The authors did not receive support from any organization for the submitted work and have no financial or non-financial interests to disclose.

Availability of Data and Material Data and material information is provided and will be shared on request.

Declarations

Ethics Approval There are no ethic approvals required for this research work.

Consent to Participate All authors will participate in the revision of the manuscript.

Consent for Publication All authors agree for the publication.

Conflicts of Interest/Competing Interests All the authors declare that there is no conflict of interest.

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Table 4 Comparison of sensor PFM with other sensor for detection of iron(III)

| Order | Fluorescent chemosensor                                                                 | Linear range (µM) | Detection limit (µM) | Recovery % | R.S.D % | Matrix                          | Ref   |
|-------|----------------------------------------------------------------------------------------|-------------------|----------------------|------------|---------|---------------------------------|-------|
| 1     | 2-(5-(4-Chlorophenyl)-3-(pyridin-2-yl)-4,5-dihydropyrazol-1-yl) 4 benzod[1]thiazole (P) | 10–200            | 3.0                  |            |         |                                 | [36]  |
| 2     | 3-(2,5-dimethylthiophen-3-yl)-5-(9-ethyl-9H-carbazol-3-yl)-4,5-dihydro-1H-pyrazol-1-ylbenzo[d]thiazole (Q) | 10–90             |                      |            |         |                                 | [37]  |
| 3     | 2-(1-(benzo[d]thiazol-2-yl)-5-(3,4-dimethoxyphenyl)-4,5-dihydro-1Hpyrazol-3-yl)phenol (R) | 10–50             |                      |            |         |                                 | [38]  |
| 4     | (2-(3-(pyridin-2-yl)-5-(3,4,5-trimethoxyphenyl)-4,5-dihydro-1H-pyrazol-1-yl)benzo[d]thiazole (S) | 0–40              |                      |            |         |                                 | [39]  |
| 5     | 4-(2-(3-Methyl-5-oxo-1-tosyl-1H-pyrazol-4(5H)-ylidene)hydrazinyl)-N-(pyrimidin-2-yl)benzenesulphonamide (T) | 20–100            | 17                   | 94.8–97    | 1.3–2.34 | River, ground, and tap water    | [2]   |
| 6     | coumarin-based pyrazoline                                                                | 0–120             | 0.101                |            |         | Imaging in HeLa cells           | [41]  |
| 7     | Pyrazole-pyrazoline (U)                                                                  | 0–10              | 0.00039              |            |         |                                 | [40]  |
| 8     | 5-(4-methoxyphenyl)-3-(5-methylfuran-2-yl)-1-phenyl-4,5-dihydro-1H-pyrazole (PFM)       | 0–3               | 0.12                 | 97.2–100   | 0.59–2.03 | Tap, river and sewage waste water | This work |
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