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Pyridine-pyrazole based Al(III) 'turn on' sensor for MCF7 cancer cell imaging and detection of picric acid†

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We report herein the development of a new pyridine-pyrazole based bis-bidentate asymmetric chemosensor that shows excellent turn-on chelation-enhanced Al³⁺-responsive fluorescence. The presence of two 'hard' phenolic hydroxyl groups plays a pivotal role in switching-on the sensing through coordination to the 'hard' Al³⁺ ion, while the mechanism can be interpreted by the chelation-enhanced fluorescence (CHEF) process. The X-ray single structure show a planar conjugated structure of the ligand, which was further stabilized by extensive H-bonding and π - π stacking. The photophysical studies related to the sensing behavior of the titular ligand toward aluminum was investigated in detail using various spectroscopic techniques like UV-Vis, photoluminescence, fluorescence and time-correlated single-photon count (TCSPC) and time-resolved NMR. The spectroscopic methods also confirm the selective detection of Al³⁺ ion in the presence of other metal ions. The theoretical calculations using Density Functional Theory (DFT) and the Time Dependent Density Functional Theory (TD-DFT) provide further insight on the mechanistic aspects of the turn-on sensing behavior including the electronic spectra of both the ligand and the complex. Interestingly, the as-synthesized H₂DPC-Al complex can also be utilized as a fluorescence-based sensor for various nitroaromatics including picric acid, for which an INHIBIT logic gate can also be constructed. The as synthesized complex was subsequently used as a fluorescent probe for imaging of human breast adenocarcinoma (MCF7) cells using live cell confocal microscopic techniques.

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Introduction

While the jury is still out on the potential health benefits of aluminum (Al) on the human body, it certainly bears the accusation of doing serious harm when somebody is exposed to larger-than-normal amounts.¹⁻⁵ Al, being the most abundant metal in Earth's crust (8.3% by weight), occurs ubiquitously in the environment. Many things that we see and use in our daily life, ranging from basic household utensils, electrical

equipment, cosmetics, to the aviation industry, are made of Al or its alloys.⁶⁻⁸ Al is also one of the most common and often indispensable components of medicine and extensively used in the pharmaceutical industry.⁹⁻¹¹ Wide spread use of Al in various food additives and packaging industry further enhance the risk of oral exposure.¹² The risk primarily comes from the fact that Al is a well-known neurotoxin and excess accumulation can lead to various neurodegenerative and neurological disorders.¹³⁻¹⁵ Two such well-documented detrimental effects of Al accumulation would be Parkinson's disease and Alzheimer's disease, for which to date the best solution relies on the philosophical remedy of "prevention is better than cure".¹⁶⁻¹⁹ Naturally as preventive measures, precise sensing and quantitative estimation of Al remain the first and foremost step in the fight against Al-related diseases. It is noteworthy here that the permissible Al consumption limit set by the World Health Organization (WHO) is 7 mg per week per kg body weight, while that of the European Food Safety Authority (EFSA) is 1 mg per week per kg body weight.^{20,21}

The above points clearly underline the importance of designing selective and sophisticated yet easy techniques for sensing of Al in its most common trivalent state (Al³⁺).²²

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However, sensing of Al^{3+} , because of its unique coordination and chemical characteristics, is somewhat different and challenging from other toxic metal sensing.²³ Some of the major limitations include strong hydration enthalpy, poor coordination ability and lack of spectroscopic characteristics.^{24,25} Notwithstanding above limitations, a vast number of methods for Al-sensing have been developed over the years including atomic absorption spectrometry, inductively coupled plasma atomic emission spectrometry, mass spectrometry, electrochemical detection as well as ^{27}Al NMR technologies.^{26,27} However, most of these methods suffer from serious limitations like time consuming lengthy experimentations, requirement of costly and sophisticated instruments.^{28,29} In contrast to these instrumental methods, optical sensing-based methods has recently gained immense interest for providing low-cost, time-saving, efficient, selective results.^{30,31}

Among the optical sensing-based techniques, fluorescence based chemosensor has gained popularity in recent time due to cost effectiveness, ease of synthesis, robust, accurate, low detection limit.^{32,33} The essence of this method is to use a fluorescent probe as chemosensor which through metal-coordination “turn-on” the Al^{3+} responsive fluorescence.^{34,35} The method usually involves three popular mechanisms in inorganic chemistry: chelation-enhanced fluorescence, photo-induced electron transfer (PET) and metal-ligand charge transfer.^{36–38} Naturally, the successful execution of this method depends on the design of fluorescent probe and its metal-ligand coordination driven luminescent behaviour.^{39–41}

The chelation-enhanced fluorescence of the resultant Al-compounds also exhibits various applications beyond chemosensing.^{42–45} One such popular application would be fluorescence-based bio-imaging and bio-labeling. The ‘turn-on’ fluorescent probes have widely been used to detect the Al-induced neurotoxicity in rodents and zebrafish.^{42,46,47}

The change in luminescent behavior of Al-complexes upon addition of a chemical probe has turned out to be important tool for sensing and detection itself.⁴⁸ The most common examples include the ‘turn-off’ or fluorescence quenching based detection of electron deficient nitroaromatics.⁴⁹ Of particular importance is the detection of picric acid (PA) because of its ambivalent significances. On one hand, picric acid is an essential multipurpose ingredient used in various fields like rocket fuel, leather industry or antiseptic agent.^{50–54} On the other hand, PA is highly explosive in nature and a notorious environmental pollutant causing serious damage to the aquatic system.^{55–57} Interestingly, Al-complexes, *inter alia*, have turned out to be simpler and economical yet efficient analytical technique for faster detection of PA.

We extend our work along these lines using a new pyridine-pyrazole based bis-bidentate asymmetric ligand, (*E*)-*N'*-(2,3-dihydroxybenzylidene)-3-(pyridin-2-yl)-1*H*-pyrazole-5-carbohydrazide (H_2DPC). The motivation comes from our earlier work where similar semi-rigid ligands efficiently coordinate to the transition metals as well as ‘hard’ trivalent lanthanides.^{58–63} Considering comparable coordination behaviors of lanthanides and Al^{3+} , we envisioned that H_2DPC containing $-\text{CONH}$, $\text{C}=\text{N}$, and $-\text{OH}$ groups would lead to similar

coordination complex formation with Al^{3+} , while the presence of two phenolic groups should turn-on the desired chelation-enhanced fluorescence. Indeed, H_2DPC act as a chemosensor and helps to detect through Al^{3+} -responsive fluorescence. The selective detection of Al^{3+} ion in presence of other metal ions is also confirmed. The as-synthesized $\text{H}_2\text{DPC}-\text{Al}$ complex was further successfully utilized for bio-imaging human breast cancer cells (MCF7). Furthermore, applicability of the as-synthesized $\text{H}_2\text{DPC}-\text{Al}$ complex for fluorescence-based sensing of various nitroaromatics including PA have also been investigated in detail.

Result and discussions

Design and synthesis of chemosensor

Clearly, the “turn-on” Al^{3+} responsive fluorescence through metal-coordination is at the heart of the matter, for the current chemosensor design. While the major obstacle remains the metal-ligand coordination, owing to the uncanny coordination behavior of the Al^{3+} ion. Optimization of coordination behavior of the ligand is therefore the most crucial aspect in Al^{3+} chemosensor design. We have recently demonstrated the effectiveness of asymmetric semi-rigid pyridine-pyrazole based Schiff base ligands in lanthanide complex formation.^{58,60} The coordination behavior of lanthanides closely resembles to that of Al^{3+} and led us believe that pyridine-pyrazole based Schiff base ligands will be equally useful for Al^{3+} .

The titular chemosensor of this study, H_2DPC , was synthesized in reasonably good yield following a typical condensation reaction between 2,3-dihydroxybenzaldehyde and 3-(pyridin-2-yl)-1*H*-pyrazole-5-carbohydrazide. We chose *ortho*-dihydroxybenzene moiety for two reasons: (a) presence of two hard O-chelating sites should favour the coordination to trivalent Al and (b) the phenolic OH-group are well known to induce chromophoric effect, which would definitely be beneficial in inducing turn-on fluorescence. For the current ligand, the PET effect expected to occur due to the lone pair of electrons of conjugated azomethine-N and phenolic-O which concomitantly quenches the fluorescence. But upon addition of Al^{3+} , the metal-coordination is expected to stop the PET effect while enhancing the fluorescence.

The H_2DPC molecule was thoroughly characterized by elemental analysis, various spectroscopic techniques like FT-IR, ESI-MS, ^1H , ^{13}C NMR as well as diffraction techniques. The FT-IR spectrum of the ligand confirms the presence of two phenolic ($-\text{OH}$) with a characteristics broad peak at 3284 cm^{-1} . On the other hand, sharp peaks at 1680 cm^{-1} and at 3325.05 cm^{-1} were observed for amide ($\text{O}=\text{C}$) stretch and amide ($\text{N}-\text{H}$) stretch, respectively. The ^1H and ^{13}C NMR data is consisted with the expected values. A characteristic peak at $\delta 14.18\text{ ppm}$ confirms the presence of pyrazole hydrogen, whereas peaks for $-\text{OH}$ proton appear at 11.38 ppm and 10.79 ppm . The peak for amide hydrogen appears at 12.18 ppm . Further corroboration comes from the ESI-MS spectra which shows a single m/z peak at 324.10 assigned to $[\text{H}_2\text{DPC} + \text{H}^+]$ and $[\text{H}_2\text{DPC} + \text{Na}^+]$ at 346.07 (Fig. S1†). For conformation of the suspected complex, we have also collected sequential mass spectra and after addition of



AlCl_3 solution as a source of Al^{3+} ion there was a new peak generate at 385.05 presumably for the Al–DPC complex assigned as $(\text{H}_2\text{DPC} + \text{Al}^{3+} + \text{Cl}^- + \text{H}^+)$ and another at 408.14 assigned as $(\text{H}_2\text{DPC} + \text{Al}^{3+} + \text{Cl}^- + \text{H}^+ + \text{Na}^+)$ (Fig. S2 and S3[†]). Furthermore, comparison of isotopic distribution pattern of the complex system with the simulated mass spectra shows that they are superimposable with each other and henceforth confirms the formation of the complex (Fig. S4[†]).

Suitable single crystals of diffraction quality were also obtained from a MeOH–DMF solvent mixture. The X-ray single crystal structure of H_2DPC (monoclinic, $P2_1/c$) shows an expected planar orientation of the molecule (Fig. 1 & S5[†]). The exo-hydroxyl groups of two adjacent H_2DPC s form strong $\text{O} \cdots \text{H} \cdots \text{O}$ hydrogen bonds with the keto groups to sustain a sideways self-assembly. The sideways arrangement is further supported by pair wise $\text{N} \cdots \text{H} \cdots \text{N}$ hydrogen bonds between two juxtaposed pyrazole–pyridine moieties. Strong π – π interactions involving all the aromatic rings, on the other hand, help the supramolecular self-assembly in the perpendicular directions.

Although the crystal structure provides sufficient information about the connectivity and solid state self-assembly, it

cannot predict the stable solvated form of the ligand. For which, we have carried out theoretical calculations. The experimental crystal structure shows some interesting synergy with the theoretically calculated structure and predicted stable conformer. In the solvent medium, H_2DPC may exist in two tautomeric forms, *viz.*, keto and enol form. Our theoretical exploration suggests that the keto form of the ligand is 0.09 eV more stable than the enol form (Fig. S6[†]). The ground state optimized geometry of keto- H_2DPC reveals $\text{O1} \cdots \text{C1}$ and $\text{C1} \cdots \text{N1}$ bond distances of 1.23 and 1.37 Å respectively, as exhibited in Fig. S6,† with NPA charges -0.689 , 0.689 and -0.464 of O1 , C1 and N1 .

Photophysical studies

Considering the fact that the theme of this work is primarily based on the chelation enhanced ‘turn-on’ Al^{3+} fluorescence, a detailed photophysical behavior of the ligand and ligand–Al complex were investigated using absorbance (UV-visible), photoluminescence (PL), fluorescence and time-resolved spectroscopic methods. The ligand, H_2DPC shows a characteristic absorption band at 302 nm in DMSO– H_2O mixture (1 : 1 v/v). In

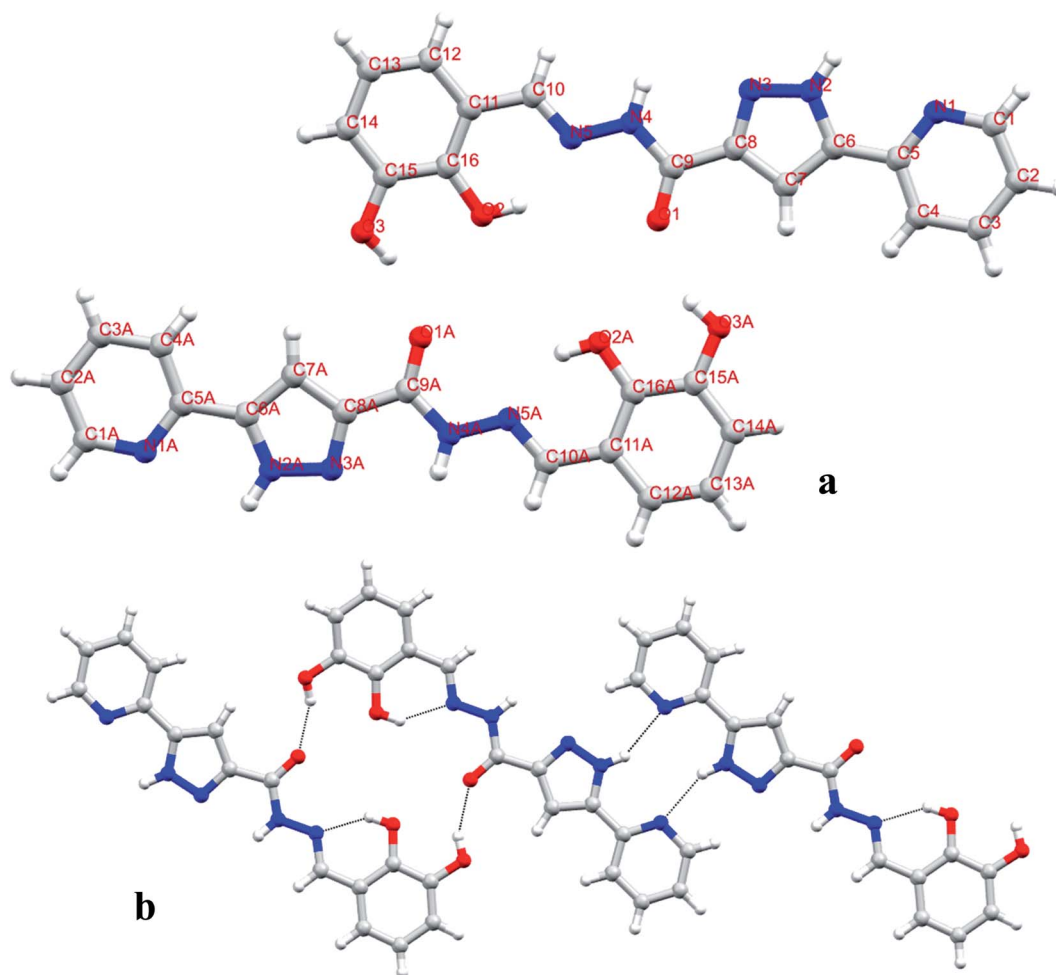


Fig. 1 (a) X-Ray single crystal structures of H_2DPC (light gray–H, dark gray–C, blue–N, red–O) (in (b) intra and inter molecular hydrogen bonds are shown by black dotted lines).



order to check the influence of metal ions on the spectral behavior of the ligand, UV-Vis absorption spectra were collected using the DMSO solution of H₂DPC maintained at pH 7.4 by using 10 mM aqueous solution of HEPES buffer. Accordingly, absorption spectrum of 10⁻⁴ M ligand solution was recorded upon addition of 10⁻⁴ M aqueous solution of different metal ions, such as Na⁺, K⁺, Ca²⁺, Cr³⁺, Al³⁺, Mn²⁺, Ni²⁺, Co²⁺, Zn²⁺, Pb²⁺ and Cd²⁺. We have also taken the images for aqueous solutions of mixture of individual metal salt with ligand, under white light and under UV light (365 nm) to observe the differences in bare eyes (Fig. S7†). As illustrated in Fig. 2, the spectral pattern of free ligand does not show any significant change on adding majority of metal ions except Al³⁺. Addition of Al³⁺ solution to the ligand brings two significant changes in the absorption spectra (a) a red shift of the absorption band from 302 nm to 330 nm and (b) appearance of a new absorption band at 393 nm, with somewhat lower intensity. The newly developed band towards the visible region is further manifested in the change in the color of the solution, from colorless to yellow, and clearly indicates ligand–Al complex formation.

The marked changes in the absorption spectra enhance the prospect of H₂DPC as a turn-on Al³⁺ chemosensor. In order to have better insight of the interaction of H₂DPC with Al³⁺ ion, we have carried out a spectrophotometric titration at room temperature in DMSO–water (1 : 1, v/v, pH = 7.4) by using aqueous 10 mM HEPES buffer solution. For the UV-Vis titration, 0.5 μL aliquot of 1 × 10⁻³ M Al³⁺ solution was added sequentially into the 1.5 ml, 1 × 10⁻⁵ M H₂DPC solution. Thus, we vary the exact concentration of Al³⁺ in the solution from 3.33 × 10⁻⁷ M to 1.2 × 10⁻⁵ M. Upon gradual addition of metal salts (Al³⁺), the strong π–π* absorption band of the free ligand at 302 nm shows a red shift to 330 nm with gradual decrease in absorbance value. Concomitantly, a new weak broad band at 393 nm appeared in the spectrum, which might be attributed to ligand to metal charge transfer (LMCT), showed a gradual increase in absorbance value without any significant shifting of the peak (Fig. 3).

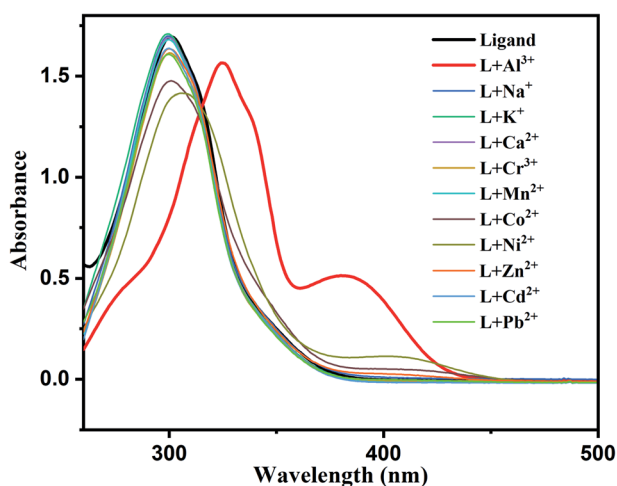


Fig. 2 UV-Vis spectra of H₂DPC on addition of various metal ions and specific change was observed in case of Al³⁺ ion.

The binding constant was also calculated from the absorption titration experiment using the Benesi–Hildebrand (B–H) plot:

$$1/(A - A_0) = 1/\{K_a(A_{\max} - A_0)[Al^{3+}]\} + 1/(A_{\max} - A_0)$$

where, A₀ and A are the absorbance of the ligand in the absence and presence of Al³⁺ ions respectively. A_{max} is the saturated absorbance of the ligand in the presence of Al³⁺ ions. K_a is the binding constant, which for the above UV-Vis titration turn out to be 3.50 × 10⁵ M⁻¹ (Fig. S8a†).

On the other hand, appearance of an isosbestic point at 321 nm clearly indicates that the reaction is clean and straightforward. Interestingly, the metal–ligand complex formation can also be monitored visually as gradual addition of the metal solution keep intensifying a yellow tinge to the initial colorless solution of the free ligand which ultimately turns it to a bright yellow solution.

Photoluminescence studies

The Photoluminescence spectra of 1 × 10⁻³ M H₂DPC in 3 ml DMSO–water 1 : 1 mixture solution (v/v) was collected after exciting the sample at λ_{ex} = 417 nm. The free ligand however shows negligible fluorescence intensity. In order to evaluate the possibility of chelation enhance fluorescence, the emission spectra of H₂DPC were recorded in presence of other metal salts. Accordingly, spectra were collected for 1.5 ml of aqueous 1 × 10⁻³ M chloride salt solutions of individually Na⁺, K⁺, Ca²⁺, Cr³⁺, Al³⁺, Mn²⁺, Ni²⁺, Co²⁺, Zn²⁺, Pb²⁺, Hg²⁺ and Cd²⁺, mixed with 1.5 ml DMSO solution of 1 × 10⁻³ M ligand solution. All the spectra were collected in buffer (pH=7.4) by using aqueous 10 mM HEPES buffer solution in DMSO water mixture (1 : 1 v/v) at room temperature using excitation wavelength 417 nm. As illustrated in Fig. 4, except Zn²⁺ and Al³⁺ ion all the other cations did not show any significant enhancement of intensity than that of the ligand. Only a slight enhancement in emission intensity

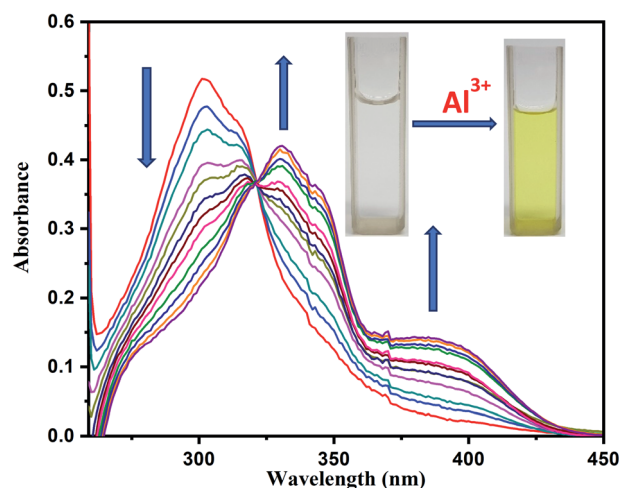


Fig. 3 On gradual addition of Al³⁺ the change in absorbance value of H₂DPC spectra. Inset: image of free ligand solution and mixture of ligand and Al³⁺ solution.



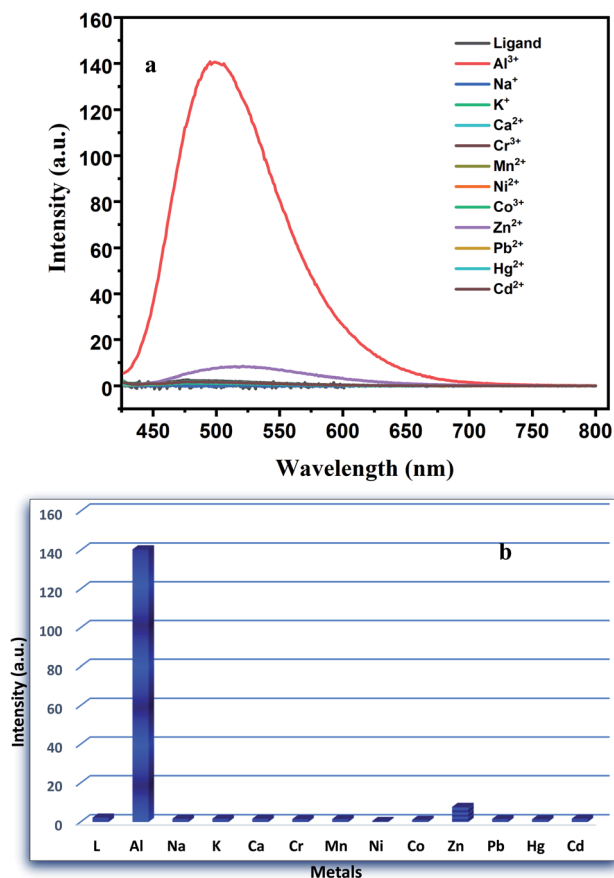


Fig. 4 (a) PL spectra of ligand in presence of various metal ions and (b) the corresponding bar plot.

is observed upon the addition of Zn^{2+} salt. To eliminate the participation of Zn^{2+} ion in the detection process EDTA has been used as a chelating ligand. The EDTA- Zn^{2+} complex has higher binding constant than the corresponding Al^{3+} complex. Naturally, EDTA can easily form complex with Zn^{2+} and separate it from Al^{3+} in the solution phase. To support that we have taken the PL spectra of Zn^{2+} - H_2DPC complex and after that we added $1 \text{ ml } 10^{-3} \text{ M}$ concentration of aqueous EDTA solution. As a result, the quenching of intensity of the solution was observed because of forming Zn-EDTA complex. Subsequently when some aqueous solution of Al^{3+} was added the intensity of photoluminescence in the mixture increases abruptly (Fig. S9[†]).

As illustrated in the Fig. 4a, we can observe that for $\text{H}_2\text{DPC} + \text{Al}^{3+}$ solution, a remarkable *ca.* 18-fold increase of the turn-on fluorescence intensity was observed at 502 nm with a red shift of 18 nm, a subsequent fluorometric titration was performed in $\text{DMSO-H}_2\text{O}$ (1 : 1, v/v, pH 7.4) by using aqueous 10 mM HEPES buffer solution to know more about the reaction of H_2DPC and Al^{3+} (Fig. 5). As per our expectation, sequential addition of Al^{3+} to the ligand solution leads to a gradual enhance of the fluorescence intensity till it reaches to the saturation point. For the PL titration $0.5 \mu\text{L}$ aliquots Al^{3+} solution with the concentration of $1 \times 10^{-3} \text{ M}$ were added sequentially into the 1.5 ml solution of the H_2DPC having the concentration of $1 \times 10^{-5} \text{ M}$. Thus, we vary the exact concentration of Al^{3+} in the solution

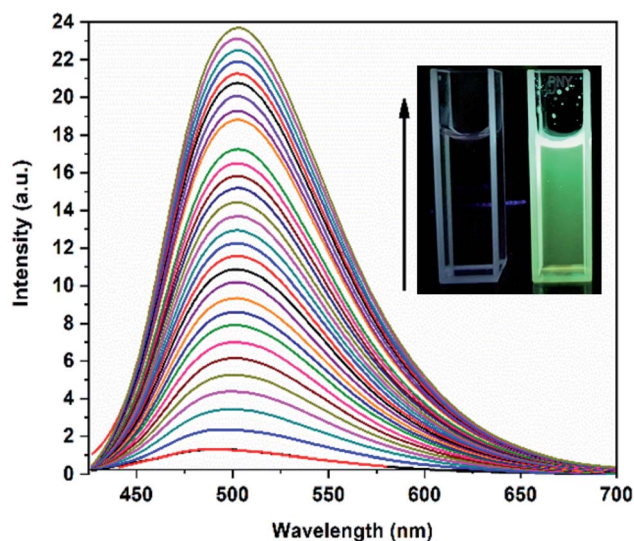
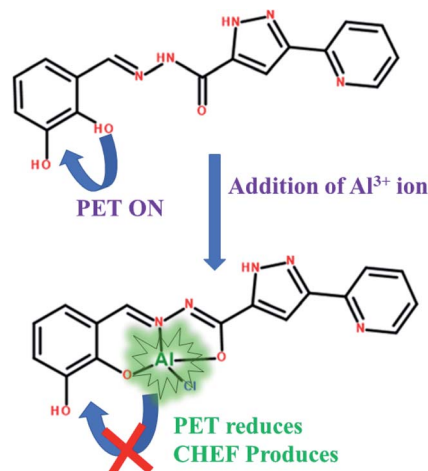


Fig. 5 The change of PL spectra of the ligand after gradual addition of Al^{3+} metal ion. The inset pictures show the comparative photographs of free ligand and green-colored luminescent ligand-metal complex after illuminating with UV light of 365 nm.

from $3.33 \times 10^{-7} \text{ M}$ to $1.2 \times 10^{-5} \text{ M}$. The binding constant of the ligand metal complex was calculated by using the Benesi-Hildebrand (B-H) plot:

$$1/(I - I_0) = 1/\{K(I_{\text{max}} - I_0)[\text{Al}^{3+}]\} + 1/(I_{\text{max}} - I_0)$$

where, I_0 is the emission intensity of ligand observed at 500 nm wavelength, I is the observed emission intensity at 500 nm in the presence of Al^{3+} , I_{max} is the maximum value of emission intensity that was obtained at 500 nm when saturation point reaches, K is the binding constant $[(I_{\text{max}} - I_0)/(I - I_0)]$ vs. $1/[\text{Al}^{3+}]$ has been plotted following the above equation and a straight line was obtained. The binding constant was determined from the slope of the linear plot and the value was $1.37 \times 10^5 \text{ M}^{-1}$



Scheme 1 The possible mode of interaction between H_2DPC and Al^{3+} ion by PET and CHEF mechanism.



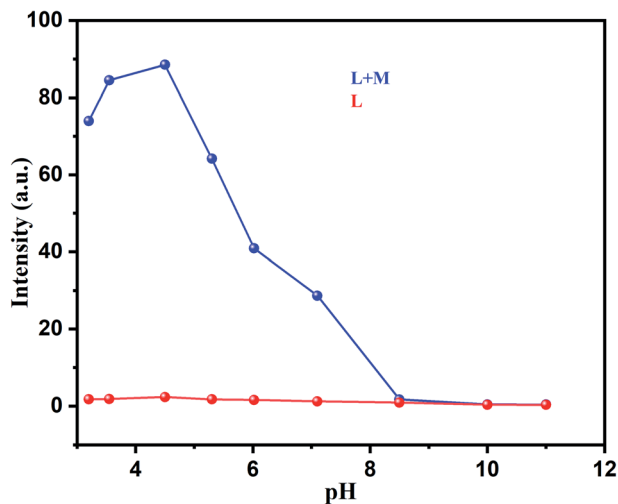


Fig. 6 The change of intensity of free H₂DPC and H₂DPC–Al solution in various range of pH.

(Fig. S8b†). The detection limit was also calculated using the equation:

$$\text{limit of detection (LOD)} = 3\sigma/k$$

where, σ is the standard deviation of blank measurement, and k is the previously determined slope. The standard deviation for the current system is turned out to be 0.141902 and henceforth the limit of detection (LOD) of Al³⁺ has been calculated as 2.15565×10^{-7} M or 0.216 μ M by using the above 3σ method (Fig. S10†). We also compared our values of binding constant and limit of detection (LOD) with some previous works done by other groups and by comparing the results we can conclude that our designed molecule is very much handful to detect the Al³⁺ ion from the aqueous solution (Table S1†).

Furthermore, the Job's plot analysis was carried out by plotting fluorescence intensity against different mole fractions of Al³⁺ while keeping volume of solution constant (Fig. S11a†). A

maximum in this plot was obtained at 0.5 mole fraction, which indicates a 1 : 1 complex formation of H₂DPC and Al³⁺. The same experiment was also performed by UV-Vis data (Fig. S11b†).

The enhancement in the fluorescence intensity can be attributed to the most common two-step processes of turn-on sensing: (a) suspension of photoinduced electron transfer (PET) and (b) activation of chelation enhanced fluorescence (CHEF) through the co-ordination of azomethine-N and phenolic-O with the metal ion. For free ligand, the electrons of both azomethine-N and phenolic-O lone pairs take part in conjugation process and activates the photoinduced electron transfer (PET) and concomitantly quenches the fluorescence. However, as soon as the Al–ligand chelation take place the lone pairs participate in metal coordination which stops PET but activate CHEF with consequential enhancement in fluorescence intensity (Scheme 1).

Effect of pH variation on fluorescence intensity of free ligand and Al-complex has also been studied (Fig. 6) in detail. As illustrated in Fig. 6 there was no significant fluorescence emission for free ligand observed between the pH range of 4 to 12. The emission spectra of the ligand were, however, completely different in presence of Al³⁺ ion. In comparison to the ligand, the Al–ligand chelation leads to the increase in emission intensity between the pH ranges of 3 to 8 (Fig. 6). Though the sharp decrease in PL intensity at the higher pH range more specifically at alkaline medium can be explained in terms of Al(OH)₃ precipitation. Interestingly, the occurrence of CHEF at the physiological pH is immensely important for any biological studies, and as such enhances the potential of the as-synthesized Al-complex as a probe for bioimaging, as discussed in later section.

Another important issue of chemosensing is the selectivity towards the concerned metal ion. In order to evaluate the selectivity of H₂DPC towards Al, the intensity of photoluminescence of H₂DPC + Al in presence of other competitive metal ions (Co²⁺, Ca²⁺, Cd²⁺, Cr³⁺, Hg²⁺, K⁺, Mn²⁺, Na⁺, Pb²⁺, Zn²⁺) were investigated. As evident from Fig. 7, addition of

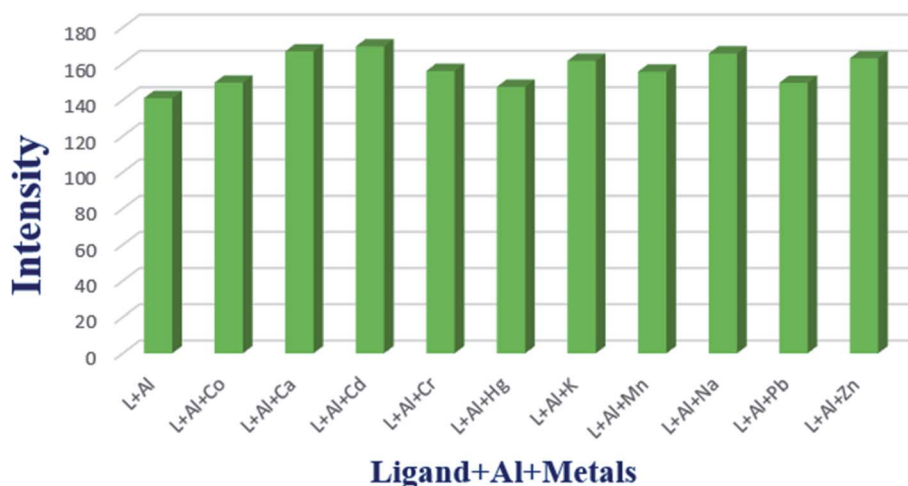


Fig. 7 The bar plot of intensity in presence of various metal ions mixture in ligand–Al solution.



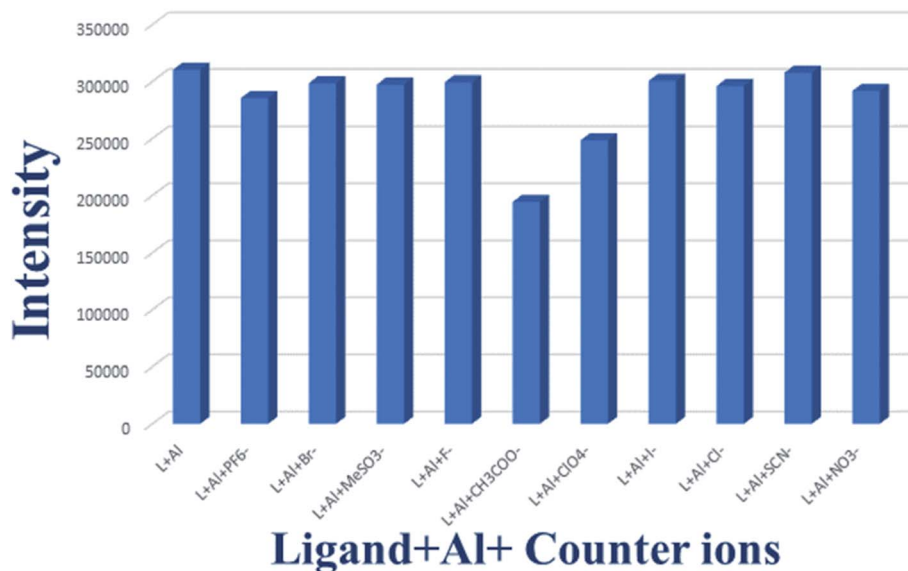


Fig. 8 The bar plot of intensity in presence of various counter ions mixture in ligand–Al solution.

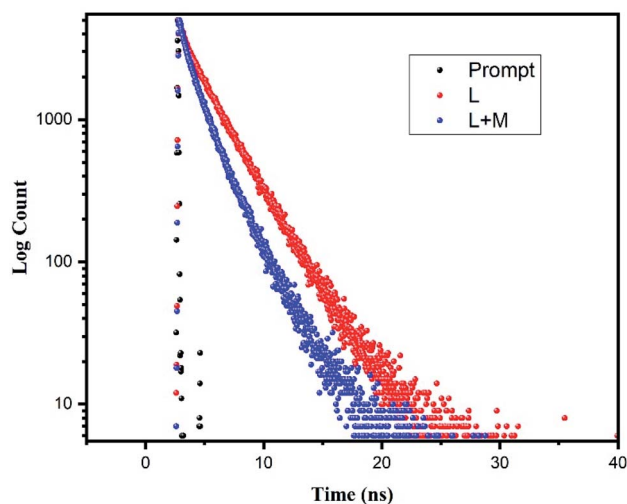


Fig. 9 Decay profile of H₂DPC and Al–H₂DPC complex.

competitive metal ions did not yield any significant change to the original intensity of the H₂DPC + Al solution. The result clearly indicates that H₂DPC can detect Al³⁺ selectively even in presence of other metal ions.

To check the effect of counter anion on the fluorescence of Al–ligand complex similar experiments were performed using different counter anions. The H₂DPC molecule showed no

significant change in fluorescence intensity when different counter anions were added to the Al³⁺ solution (Fig. 8). For these experiments, 10^{−3} M solution of each anion (PF₆[−], Br[−], MeSO₃[−], F[−], CH₃COO[−], ClO₄[−], I[−], Cl[−], SCN[−], NO₃[−]) was added into the 2 ml of 10^{−3} M solution of the ligand–metal system. Interestingly, for CH₃COO[−] and ClO₄[−], there was very little decrease in the original intensity of the complex (Fig. S12†) were observed. Furthermore, the fluorescence intensities in presence various inorganic acids, such as HCl, HNO₃ and H₂SO₄ (Fig. S13†) were also investigated. The result shows that the intensity was increased in presence of all acids while maximum effect was observed for H₂SO₄.

Time resolved fluorescence spectroscopy

The fluorescence lifetime measurements using time-correlated single-photon count (TCSPC) experiments were carried out to determine and compare the average life-time free H₂DPC molecule and Al–H₂DPC complex. Lifetime measurement data were taken on excitation at 405 nm and as depicted in Fig. 9, the fluorescence decay plot exhibits a bi-exponential nature. As expected for a CHEF process, the life time of the free ligand (H₂DPC, 3.74510 ns) decrease in presence of Al³⁺ ion, with a corroborating life time 1.8 ns for the Al–H₂DPC complex (Table 1). Further support comes from the theoretically calculated values show that the Δ*E* value for free ligand is higher than the Δ*E* value of complex. In the theoretical study we have seen that the Δ*E* value for ligand is 4.12 eV whereas for metal

Table 1 The values for average life time of H₂DPC and Al–H₂DPC complex

Environment	<i>a</i> ₁	<i>a</i> ₂	<i>τ</i> ₁ (ns)	<i>τ</i> ₂ (ns)	<i>χ</i> ²	<i>⟨τ⟩</i> (ns)
H ₂ DPC	0.2255	0.7745	0.599995	4.66082	1.507279991	3.74510
H ₂ DPC–Al	0.7272	0.2728	1.54612	3.00989	1.002613126	1.94543



complexes it is 3.72 eV so due to the low energetic HOMO–LUMO gap for complex than ligand. The overall number of transitions between two state getting increase as a result the intensity getting higher after adding Al^{3+} ion in ligand solution but overall, the average life time for metal complex also getting decrease than only ligand. The values of a_1 and a_2 are also determined using the equation,

$$a_1 = \frac{B_1}{B_1 + B_2}$$

and

$$a_2 = \frac{B_2}{B_1 + B_2}$$

The value of B_1 and B_2 are given as 22.55% and 77.45% for ligand and 72.72% and 27.28% for ligand metal complex and B is referred as relative amplitude and the average life time was calculated with the following equation $\langle \tau \rangle = a_1\tau_1 + a_2\tau_2$.

$^1\text{H-NMR}$ titration analysis

For better understanding of the interaction of H_2DPC with Al^{3+} ion, $^1\text{H-NMR}$ spectroscopic titration was carried out in $\text{DMSO-}d_6$ solvent. The titration was done with sequential addition of AlCl_3

(D_2O solution) in H_2DPC ($\text{DMSO-}d_6$) solution. Upon gradual addition of Al^{3+} solution into the ligand solution led to the disappearance of the peaks for phenolic–OH protons at δ 11.383 ppm and δ 10.792 ppm (Fig. 10). It actually implies the breaking of intra molecular hydrogen bonding between the phenolic–OH and imine $-\text{C}=\text{N}$ and also the deprotonation of phenolic–OH. We have observed in theoretical structure that the $-\text{NH}$ beside the carbonyl group get tautomerized and form $\text{C}=\text{N}$ in complex formation. The time-resolved NMR spectra also exhibits the disappearance of peak for amide H in δ 12.181 ppm upon step by step increase of metal ion concentration, which actually confirms the complex formation. Furthermore, a downfield shift was observed for the proton at δ 8.645 ppm. Based on the titration data we can conclude that because of the high ionic potential of Al^{3+} there is hard–hard interaction between Al^{3+} and phenolic (O^-), as well as with imine $\text{N}(-\text{C}=\text{N})$, which actually driven the complex formation process.

Theoretical studies

Even after numerous attempts, we could not find a diffraction quality single crystal of the $\text{H}_2\text{DPC-Al}$ complex. This prompts us to carry out theoretical calculation was done on $\text{H}_2\text{DPC-Al}$. As

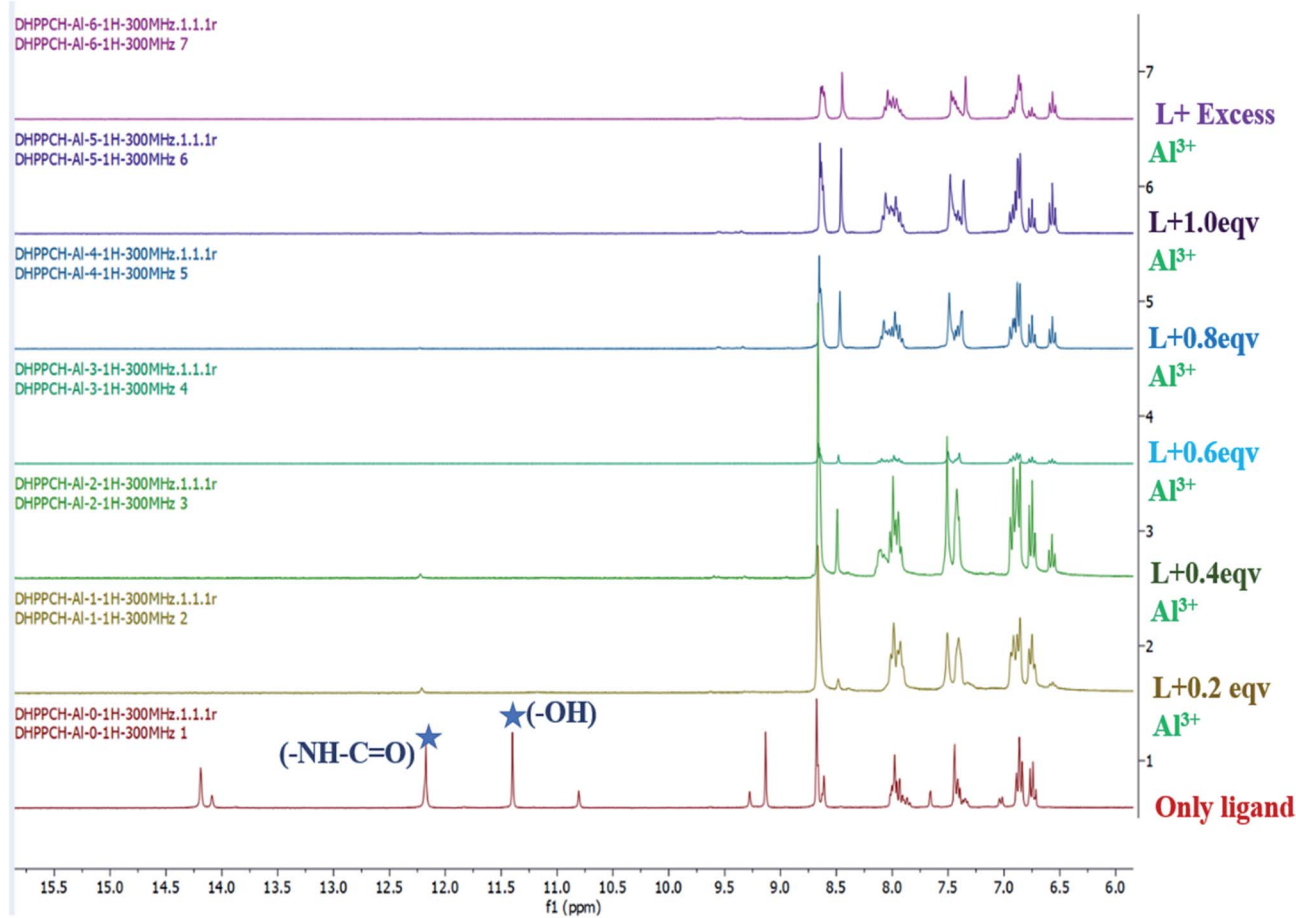


Fig. 10 The NMR titration of H_2DPC (in $\text{DMSO-}d_6$ solution) with gradual addition of Al^{3+} (in D_2O solution).



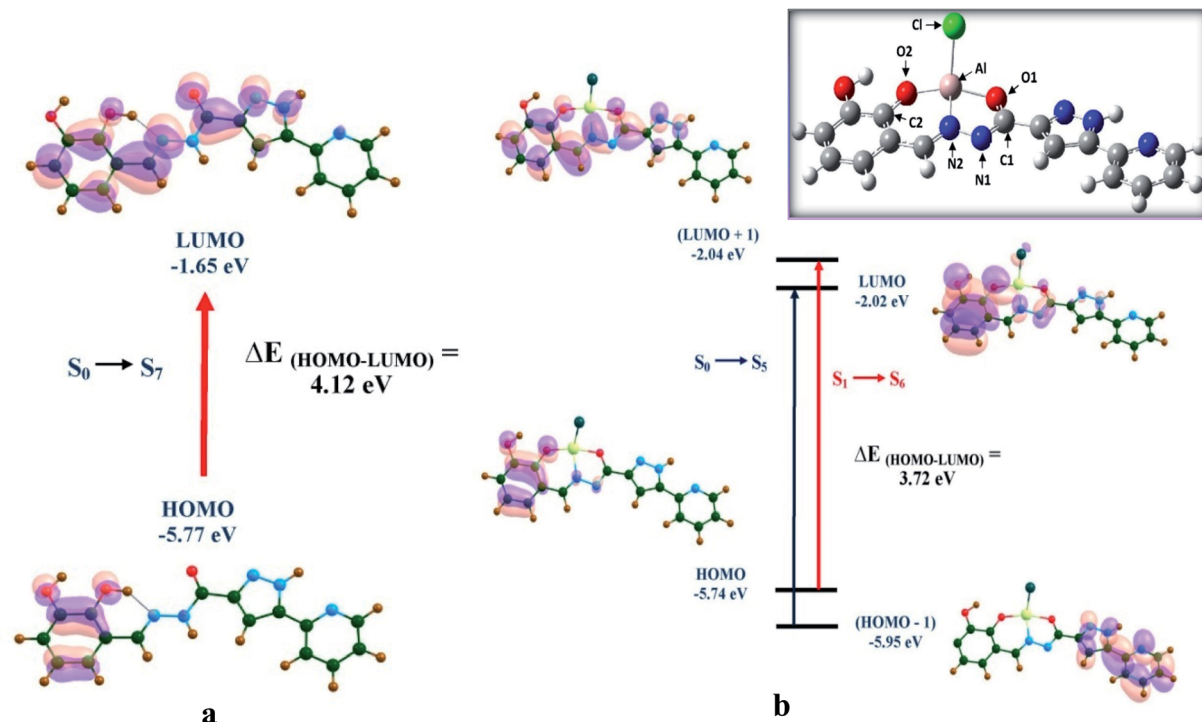


Fig. 11 (a) Frontier molecular orbitals involved in electronic transition of keto-H₂DPC. (b) Frontier molecular orbitals involved in electronic transition of H₂DPC–Al (inset: optimized geometry of the complex H₂DPC–Al).

evident from Fig. 11 (inset) the phenolic oxygen (O2), carbonyl oxygen (O1) and the nitrogen (N2) which is next to amide nitrogen act as the chelating sites in the complex H₂DPC–Al, with O–Al–O bond angle of 156.7°. Upon complex formation with Al³⁺, the distance between C1 and O1 increases to 1.31 Å, while that between N1 and C1 decreases slightly to 1.32 Å, as displayed in (Table S2[†]). The NPA charges of O1, C1, N1 and Al are found to be –0.880, 0.605, –0.436 and 0.241 respectively, while –0.926, 0.334 and –0.514 of O2, C2, N2.

Theoretical outcome shows that the HOMO of keto-H₂DPC, in Fig. 11a, is congested over the 2-hydroxyphenol part of the ligand and the LUMO mainly over the 2-hydroxyphenol part, with a slight localization over the pyrazole part, as well. As depicted in Fig. 11b, the HOMO of H₂DPC–Al complex is congested over the 2-hydroxyphenol part of the ligand, similar to that of keto-H₂DPC, while the (HOMO–1) is distributed solely over the pyrazole part. On the other hand, both the LUMO and

(LUMO+1), are distributed nearly over the entire ligand, and not on the Al centre. Thus, it is evident, that the absorption bands are due to intra-ligand n–π* and π–π* electronic transitions and not because of any metal to ligand or ligand to metal charge transfer.

The HOMO–LUMO gap for the ligand is calculated to be 4.12 eV, which gets lowered to 3.72 eV in the complex H₂DPC–Al. The decrease in HOMO–LUMO gap by 0.40 eV accounts for the red shift in the UV-visible spectra. The electronic transition, S₀ → S₅, at 310.8 nm, corresponding to electron transfer from HOMO to LUMO accounts for the experimentally observed peak at 301.52 nm in the UV-visible spectra. The absorption bands at 330.6 and 390.11 nm in the spectra result from the electronic transitions, S₀ → S₅ (at 346.75 nm) and S₁ → S₆ (at 370.36 nm), corresponding to (HOMO–1) to LUMO and HOMO to (LUMO+1) electron transfers. The results obtained using TZVP basis set are in accordance with the 6-31++G** values. The TZVP

Table 2 The values of TZVP along with all the oscillator strength value

Molecule	Basis set	Energy (eV)	Electronic transition state	Excitation (nm)	Oscillator strength (f)	Experimentally observed transition (nm)	Corresponding MOs
Keto-H ₂ DPC	6-31++ G**	3.99	S ₀ –S ₇	310.80	1.0790	301.52	HOMO → LUMO
	TZVP	3.91	S ₀ –S ₇	317.48	1.0866		
DPC–Al	6-31++ G**	3.58	S ₀ –S ₅	346.75	1.0657	330.60	(HOMO–1) → LUMO
	TZVP	3.54	S ₀ –S ₅	349.80	1.0589		
	6-31++ G**	3.35	S ₁ –S ₆	370.36	1.1015	390.11	HOMO → (LUMO+1)
	TZVP	3.32	S ₁ –S ₆	373.99	1.1158		



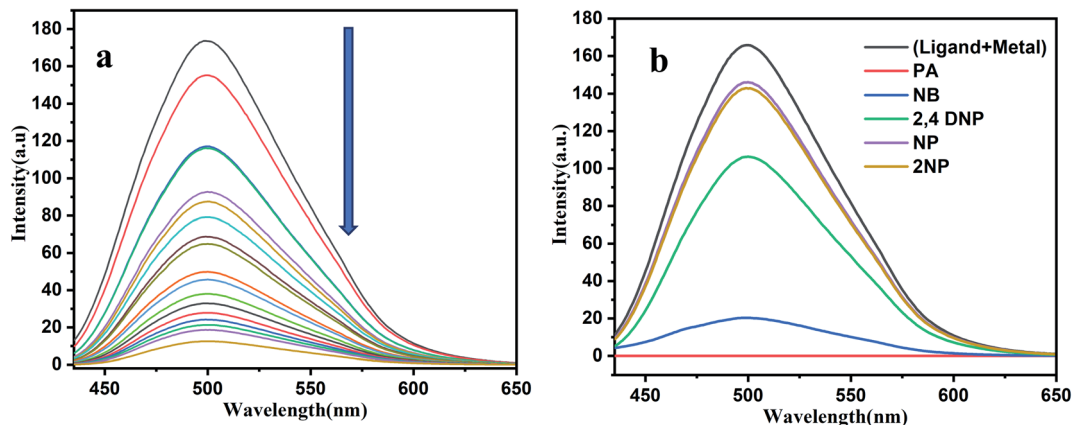


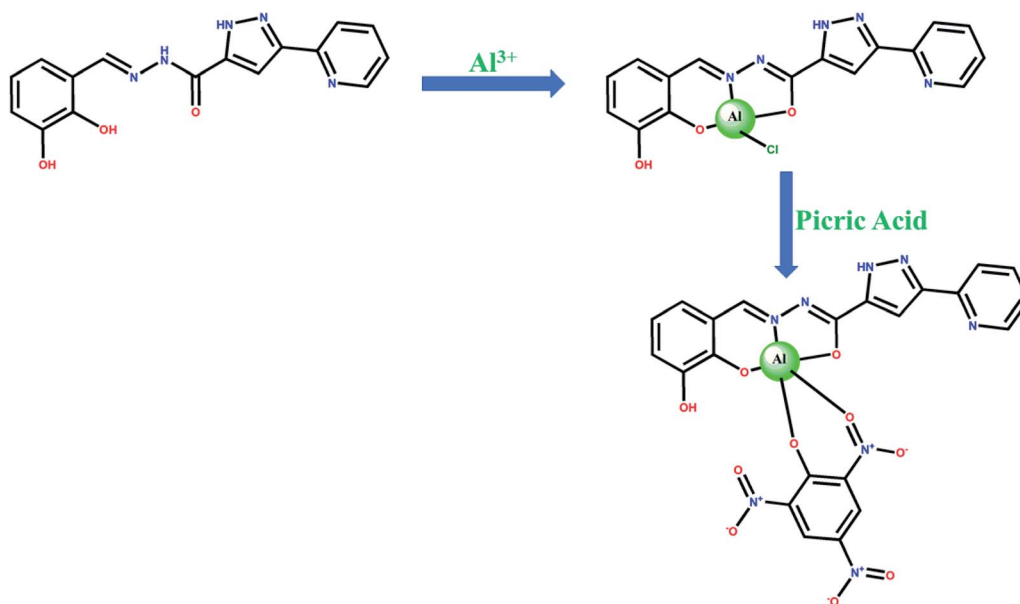
Fig. 12 (a) Change of intensity of H₂DPC–Al complex solution after gradual addition of picric acid into the solution. (b) PL spectra of H₂DPC–Al complex in presence of various nitro aromatic molecules.

values, along with all the oscillator strength values have been exhibited in Table 2.

Application of H₂DPC–Al in picric acid sensing

Interestingly, fluorescence properties of H₂DPC–Al ensemble are found to be dependent on the presence of nitroaromatic (NA) compounds and therefore can be employed for their detection. The fluorescence of H₂DPC–Al compound exhibits the quenching of emission upon the addition of (100 μ L) various nitro aromatic compounds including nitrobenzene, 2,4-dinitrophenol, 2-nitrophenol, 2,4,6-trinitrophenol and 4-nitrophenol (Fig. 12b). Such a quenching effect should be attributed to the charge transfer electron transitions from the H₂DPC–Al ensembles to NA due to the electron-deficient property of NA and the π – π interactions between complex and NA molecules. Among the different NAs, 2,4,6-trinitrophenol or picric acid,

which can also be envisaged as the most important variant of NAs for various ambivalent reasons, shows the most prominent emission quenching. We observed that under UV light the change in intensity for picric acid solution is most pronounced (Fig. S14[†]). This supports the candidature of H₂DPC–Al as a promising luminescent probe for detecting picric acid among all the NAs. For detailed study, the fluorescence intensity of the H₂DPC–Al solution was monitored upon sequential addition of small aliquot of picric acid solutions. As depicted in Fig. 12a, quenching of the fluorescence intensity was observed. While the detection limit of the H₂DPC–Al ensemble toward PA was calculated using the formula of detection limit = $3\sigma/k$. Here, σ (standard deviation) = 1.384999 and k (slope of the plot) = 33895.77. The value of detection limit is 1.2257×10^{-4} (M) (Fig. S15[†]). We have investigated the mechanism for this quenching process. For this, we have collected the mass spectra



Scheme 2 The probable quenching mechanism of Al–H₂DPC complex by picric acid.



of the ($\text{H}_2\text{DPC} + \text{Al} + \text{PA}$) complex and found that peak at 669.28 is probably due to the $\text{H}_2\text{DPC} + \text{Al}^{3+} + \text{PA} + 4\text{Na}^+$ assembly (Fig. S16[†]). The probable quenching mechanism was shown in Scheme 2. From the mechanism we can easily conclude that due to the strong -I and -R effect the electron cloud is getting shifted towards PA and as a result the luminescent intensity of the overall complex is getting decreased.

Formation of logic gate

Applying the reversible behavior of our ligand H_2DPC towards Al^{3+} ion in presence of picric acid in the solution we actually constructed an INHIBIT logic gate with two inputs, Al^{3+} and picric acid (Fig. 13a). Four combinations are possible. In the first case we only use the ligand but both the inputs are not added to the solution so there is no such fluorescence was observed. We can conclude the result as zero. In the second combination we added only one input that is Al^{3+} ion and we get an observable fluorescence. We can conclude the result as one. For the third combination we added only the second input that is picric acid. We did not observe any fluorescence, so we can conclude the result as zero. In the fourth combination we added both the inputs. Though Al^{3+} helps the increase in fluorescence but due to the quenching ability of PA we did not observed any luminescence so we can conclude the result as zero. Thus, in this way we actually constructed a circuit for the INHIBIT gate which ideally follow the truth table of INHIBIT gate (Fig. 13c). We have also taken the picture of all four combinations under UV lamp (Fig. 13b).

Biological studies

The luminescent properties, particularly the characteristics green luminescence of $\text{H}_2\text{DPC}-\text{Al}$ was highly encouraging and prompts us to explore the potential of the complex as a fluorescent probe for cellular imaging. Accordingly, the synthesized $\text{H}_2\text{DPC}-\text{Al}$ complex has been used as a fluorescent probe for the live cell labeling of human breast adenocarcinoma

(MCF7) cells. To be an effective probe for cellular sensing internalization into cell as well as low cytotoxicity are the two most important prerequisites. Accordingly, chances of cytotoxic effects of $\text{H}_2\text{DPC}-\text{Al}$ were evaluated using the conventional cell cytotoxicity (MTT) survival assay experiments. In survival assays, MCF7 cells treated with increasing concentrations of $\text{H}_2\text{DPC}-\text{Al}$ for 72 hours manifested least cytotoxicity as >90% of the cells remained viable even at 50 μM concentration of the compound. In other words, the $\text{H}_2\text{DPC}-\text{Al}$ complex does not exert any adverse effect on the cell and therefore can be termed as nontoxic (Fig. S17[†]).

Having confirmed its non-toxic nature, human breast adenocarcinoma (MCF7) cells seeded in cover glass bottom dish were exposed to 20 μM $\text{H}_2\text{DPC}-\text{Al}$ for 6 h and subjected to live cell confocal microscopy thereafter. As illustrated in Fig. 14, the $\text{H}_2\text{DPC}-\text{Al}$ complex was achieving the critical cell-permeability as such easily pass through the cell membrane during the incubation period. Following the conventional cellular sensing protocol, the nuclei of the cells were stained with Hoechst 33342 (Blue) nuclear marker. Treatment with $\text{H}_2\text{DPC}-\text{Al}$ revealed significant cellular uptake as it permeated easily through the MCF7 cells without causing any considerable damage. Moreover, the nuclei remained intact and showed no significant deformation. The $\text{H}_2\text{DPC}-\text{Al}$ (green) localized both in the nucleus and cytoplasm as observed in confocal microscopy. In order to check whether ligand itself can be used as a fluorescence probe, the live cell labelling of human breast adenocarcinoma (MCF7) cells experiment was repeated using the ligand solution of same concentration. However, the ligand did not show appreciable fluorescence intensity. The result has another implication that the ligand is unable to detect the presence of inherent aluminum in MCF7 cells. This could be attributed to the fact that the concentration of aluminum in the cytoplasm of cell and nucleus is in the range of femtomolar⁶⁴ whereas our ligand has the limit of detection for Al^{3+} ion is in the range of 10^{-7} M.

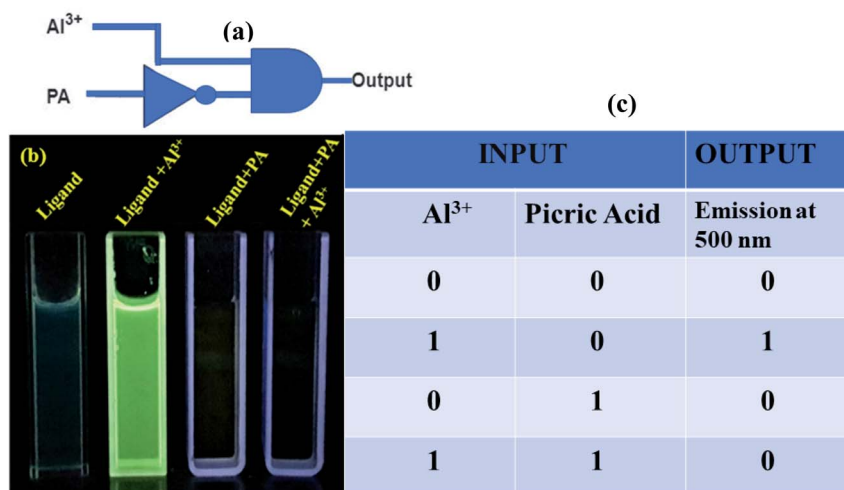


Fig. 13 Operation of inhibit logic gate. (a) Pictorial representation of the logic gate (b) visual color outputs. (c) Truth table corresponding to the INHIBIT logic gate.



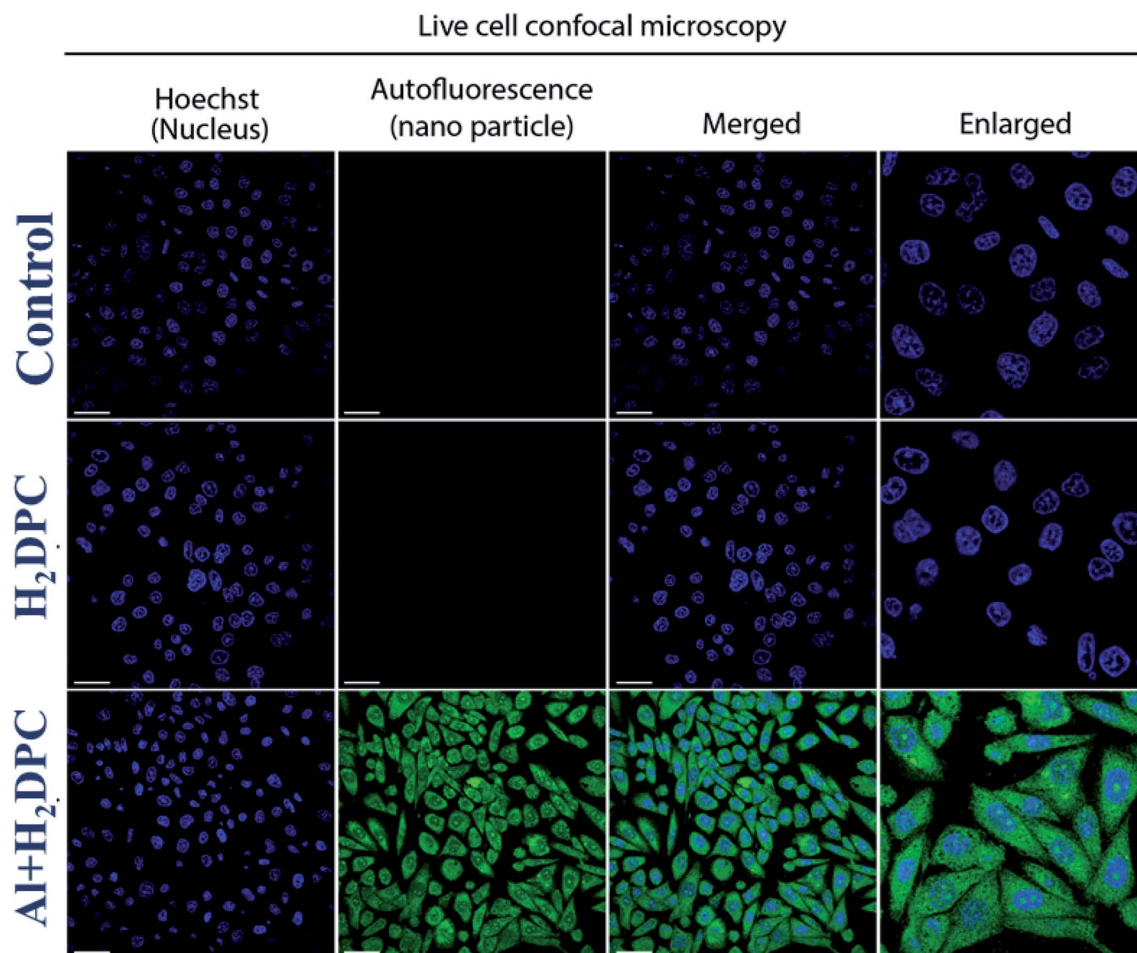


Fig. 14 Live cell confocal microscopy showing delivery of compounds H₂DPC and H₂DPC–Al (green) in the nucleus and cytoplasm. Representative confocal images showing localization of compounds respectively (20 μ M) in human breast adenocarcinoma (MCF7) cells as monitored by intrinsic fluorescence of the drugs (H₂DPC–Al green channel, middle panel) and with Hoechst 33342 (blue channel, left panel) used as a nuclear marker. H₂DPC shows no luminescence inside cells. Right panel shows the merged images.

Conclusions

The impetus for the work presented herein was to develop a pyridine-pyrazole based 'turn on' chemosensor for Al(III). Accordingly, we have successfully synthesized a new pyridine-pyrazole based bis-bidentate asymmetric ligand fitted with two 'hard' –OH groups. The presence of two phenolic groups indeed triggers a turn-on chelation-enhanced Al³⁺-responsive fluorescence. The sensing mechanism based on the chelation-enhanced fluorescence (CHEF) process was studied in detail using various spectroscopic techniques including UV-Vis, PL, fluorescence as well as NMR titration. The spectroscopic methods also confirm the selective detection of Al³⁺ ion in presence of other metal ions. Further corroborative support in favor of sensing mechanism comes from the DFT calculations. On the other hand, theoretical results obtained from the Time Dependent Density Functional Theory (TD-DFT) calculations show an excellent agreement with the electronic spectra corresponding to the ligand (H₂DPC) and the complex (H₂DPC–Al).

The as-synthesized H₂DPC–Al complex shows an excellent fluorescence-based sensing of various nitroaromatics including

picric acid, for which an INHIBIT logic gate can also be constructed. With no apparent cytotoxic adverse effects, the as-synthesized H₂DPC–Al complex was further successfully utilized for bio-imaging human breast cancer cells (MCF7) using live cell confocal microscopic techniques.

Experimental section

Materials and general methods

All the reagents and solvents used in this work were purchased from commercial sources & used without further purification. Solid state FT-IR spectra of 400–4000 cm⁻¹ range were collected from Nicolet MAGNA-IR 750 spectrometer with sample prepared as KBr pellets. Elemental analyses were performed with Perkin-Elmer 2400 series CHN analyser. ¹H and ¹³C NMR spectrum were carried out by 400 MHz and 500 MHz Bruker NMR spectrometers respectively dissolving sample in DMSO-d₆ solvent. Mass spectra was collected by Q-ToF Micro YA263 high resolution (Waters Corporation) mass spectrometer by positive ion mode electrospray ionization and the spectra were collected in methanol. Absorption spectra were collected with a Jasco V-



630 spectrophotometer. Photoluminescence spectra were collected at room temperature using Agilent Cary Eclipse & Hitachi, F-2500 fluorescence spectrophotometer and Horiba scientific TCSPC.

Synthesis of (*E*)-*N'*-(2,3-dihydroxybenzylidene)-3-(pyridin-2-yl)-1*H*-pyrazole-5-carbohydrazide (H_2DPC)

The ligand, (*E*)-*N'*-(2,3-dihydroxybenzylidene)-3-(pyridin-2-yl)-1*H*-pyrazole-5-carbohydrazide (H_2DPC) was synthesized by condensation reaction between 2,3-dihydroxybenzaldehyde (1 mmol, 138 mg) & 3-(pyridin-2-yl)-1*H*-pyrazole-5-carbohydrazide (1 mmol, 203 mg) in ethanol medium. 3-(Pyridin-2-yl)-1*H*-pyrazole-5-carbohydrazide was synthesized using previously reported procedure.⁵⁸ The mixture was stirred for 30 minutes and a drop of glacial acetic acid was added to the suspension. Then the mixture was refluxed for 6 hours under 80 °C. The white precipitate was found. The mixture was cooled to room temperature. Then it was filtered to collect the white precipitate. It was washed with ethanol and recrystallized from methanol (Scheme S1†). The compound was characterized by elemental analysis, FT-IR, ESI-MS & NMR spectrometer. (Yield: 245 mg, 71%). Calcd for $C_{16}H_{13}N_5O_3$ (%): C, 59.44; H, 4.05; N, 21.66. Found: C, 59.22; H, 3.65; N, 22.53. IR (400–4000 cm^{-1}): 3325.05(s), 3284(s), 3161(s), 2997(s), 1679(s), 1548(s), 1473(s), 1406(s), 1267(s), 1205(s), 1159(s) (Fig. S18†). ¹H NMR (500 MHz, [(CD₃)₂SO], δ) 14.183 (s, 1Hpyz-H), 12.181 (s, 1H, NH), 11.383 (s, 1H, phenolic-OH), 10.792 (s, 1H, phenolic-OH), 9.133 (s, 1H, imine-H), 7.867–8.512 (4H, py-H), 6.895–7.212 (4H, Ar-H), 7.615 (1H, pz-CH) (Fig. S19†). ESI-MS m/z (M + H): 324.1064 (Fig. S1†).

Synthesis of Al³⁺ complex of H_2DPC ($H_2DPC-Al$)

To the solution of H_2DPC (0.01 mmol, 3.23 mg) in 0.5 ml DMSO, 0.5 ml of aqueous solution of AlCl₃ (0.01 mmol, 1.33 mg) was added. As a result, the colorless ligand solution turned to yellow. After evaporation of the solvent a yellow mass was extracted. The solid mass was washed for several time with water and dried in vacuum. The compound was characterized by elemental analysis, FT-IR, ESI-MS & NMR spectrometer. (Yield: 69%). Calcd for $C_{16}H_{11}AlClN_5O_3$ (%): C, 50.08; H, 2.89; N, 18.25. Found: C, 49.22; H, 2.50; N, 17.97. IR (400–4000 cm^{-1}): 3375.20(b), 3168(b), 1662(s), 1625(s), 1606(s), 1585(s), 1465(s), 1286(s), 1205(s), (Fig. S20†) ¹H NMR (500 MHz, [(CD₃)₂SO], δ) 12.193(s, 1H, NH), 7.867–8.512 (4H, py-H), 6.895–7.212 (4H, Ar-H), 7.615 (1H, pz-CH) (Fig. S21†) ESI-MS m/z (M + H): 385.0522 (Fig. S2 and S3†).

Live-cell confocal microscopy

Live-cell imaging was carried out using confocal laser-scanning microscope (Leica TCS SP8) with a UV-laser and 63X/1.4 NA oil objective equipped with a heated environmental chamber set to 37 °C with optimal CO₂ facility. Briefly, MCF7 cells seeded on cover glass bottom dish (Genetix, Biotech Asia Pvt. Ltd.) were incubated with compounds H_2DPC and $H_2DPC-Al$ individually for 6 h. Nuclei were subsequently stained with Hoechst 33342 (Blue) (Sigma). This was followed by live cell confocal

microscopy. Images were collected and processed using the Leica software and sized in Adobe Photoshop 7.0.⁶⁵

Cell survival assay

MCF7 cells (6×10^3) were seeded in 96-well plate (BD Biosciences, USA) and treated with compound $H_2DPC-Al$ at the indicated concentrations. After 72 h treatment, cell survival was assessed by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay. Briefly, cells were washed with 1 × PBS and treated with MTT reagent (Sigma) for 3 h at 37 °C and the resulting formazan was dissolved in 100 μ L of DMSO. Plates were analysed on Molecular Devices Spectra Max M2 Microplate Reader at 570 nm. The percent inhibition of viability for each concentration of the compounds was calculated with respect to the control. Data represent mean values \pm s.e.m. for three independent experiments.⁶⁶

Computational details

In order to get a better understanding of the sensing mechanism, we have carried out Density Functional Theory (DFT) calculations. All electronic structures and frequency calculations have been performed using Gaussian 09 (ref. 67) suite of quantum chemistry program. The DFT functional considered for the systems under investigation is B3LYP,^{68,69} in conjunction with 6-31++G** basis set. The self-consistent reaction field (SCRf) method has been implemented using COSMO^{70,71} to take into account the effect of bulk solvent medium, considering dimethyl sulfoxide (DMSO) as the solvent. To interpret electronic spectra corresponding to the ligand H_2DPC and the complex $H_2DPC-Al$, we have carried out Time Dependent Density Functional Theory (TD-DFT) calculations using B3LYP method and basis set 6-31++G**. In order to further validate the obtained results, we have considered TZVP basis set also in conjunction with the same method.

Single crystal X-ray crystallography

X-Ray diffraction intensities for the ligand H_2DPC was collected at 100 K on a Bruker D8 VENTURE Microfocus diffractometer equipped with PHOTON II Detector, with Mo-K α radiation ($\lambda = 0.71073 \text{ \AA}$), controlled by the APEX3 (v2017.3-0) software package and processed using SAINT. Raw data were integrated and corrected for Lorentz and polarization effects using the Bruker APEX III program suite. The structures were solved by direct methods in SHELXS and refined by full-matrix least squares on F^2 in SHELXL. Crystallographic data are summarized in reference no. 72 and the CIF has been deposited with the Cambridge Crystallographic Data Centre (CCDC).⁷²

Conflicts of interest

There are no conflicts to declare.

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