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A highly selective pyridoxal-based chemosensor for the detection of Zn(II) and application in live-cell imaging; X-ray crystallography of pyridoxal-TRIS Schiff-base Zn(II) and Cu(II) complexes†

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In a simple, one-step reaction, we have synthesized a pyridoxal-based chemosensor by reacting tris(hydroxymethyl)aminomethane (TRIS) together with pyridoxal hydrochloride to yield a Schiff-base ligand that is highly selective for the detection of Zn(II) ion. Both the ligand and the Zn(II) complex have been characterized by ¹H & ¹³C NMR, ESI-MS, CHN analyses, and X-ray crystallography. The optical properties of the synthesized ligand were investigated in an aqueous buffer solution and found to be highly selective and sensitive toward Zn(II) ion through a fluorescence turn-on response. The competition studies reveal the response for zinc ion is unaffected by all alkali and alkaline earth metals; and suppressed by Cu(II) ion. The ligand itself shows a weak fluorescence intensity (quantum yield, $\Phi = 0.04$), and the addition of zinc ion enhanced the fluorescence intensity 12-fold (quantum yield, $\Phi = 0.48$). The detection limit for zinc ion was 2.77×10^{-8} M, which is significantly lower than the WHO's guideline (76.5 μ M). Addition of EDTA to a solution containing the ligand–Zn(II) complex quenched the fluorescence, indicating the reversibility of Zn(II) binding. Stoichiometric studies indicated the formation of a 2 : 1 L₂Zn complex with a binding constant of 1.2×10^9 M⁻² ($\pm 25\%$). The crystal structure of the zinc complex shows the same hydrated L₂Zn complex, with Zn(II) ion binding with an octahedral coordination geometry. We also synthesized the copper(II) complex of the ligand, and the crystal structure showed the formation of a 1 : 1 adduct, revealing 1-dimensional polymeric networks with octahedral coordinated Cu(II). The ligand was employed as a sensor to detect zinc ion in HEK293 cell lines derived from human embryonic kidney cells grown in tissue culture which showed strong luminescence in the presence of Zn(II). We believe that the outstanding turn-on response, sensitivity, selectivity, lower detection limit, and reversibility toward zinc ion will find further application in chemical and biological science.

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Introduction

Zinc is the second most abundant metal in the human body, and plays a critical role in many biological processes such as gene expression, protein and DNA synthesis, enzymatic

reactions, growth, and development processes, as well as in many pathological processes such as epilepsy, Alzheimer's disease, infantile diarrhea, and ischemic stroke.^{1–6} In addition, a high concentration of chelatable zinc ion is found in the brain, spermatozoa, and pancreas.^{7–9} Chelatable Zn(II) has numerous benefits in biological systems such as regulating neuron transmission, suppressing apoptosis, immune function, *etc.*; however, the mechanism of chelatable Zn(II) is less understood as compared to other cations such as Na(I), K(I), and Ca(II). On the other hand, Zn(II) is spectroscopically silent due to its closed-shell [Ar]3d¹⁰4s⁰ electronic configuration; hence detecting zinc with conventional techniques such as nuclear magnetic resonance (NMR), electron paramagnetic resonance (EPR), and electronic absorption spectroscopy is ineffective.⁵ Atomic absorption spectroscopy can detect zinc selectively and sensitively; however, this technique is destructive to the sample

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† Electronic supplementary information (ESI) available: ¹H NMR, ¹³C NMR, COSY NMR, ¹³C-DEPT135 NMR, IR and ESI-MS spectrum of ligand, EDTA reversibility plot, pH effect plot, detection limit plot, UV-Vis and emission plot on addition of anions, ESI-MS of metal complex, quantum yield and detection limit calculation. CCDC 2095325 (1), 2095330 ([1⁻]₂Zn²⁺·5H₂O) and 2095332 ([1]Cu²⁺·2ClO₄⁻). For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d1ra05763d



and has no spatial resolution.¹⁰ Analytical methods such as UV-Vis spectroscopy, potentiometry, and electrochemical analysis have also been used for the detection of Zn(II), Cu(II), and Co(II).^{11–13} Fluorescent probes are the most effective tools for sensing applications, however, exhibiting high sensitivity and selectivity, and have generally short response times when imaging biological samples. Several fluorescent probes have been reported for the detection of zinc ion in a wide variety of cells such as A549 cells, SW620 cells, Vero cells, HeLa cells, MDA-MB 468 cells, HEK 293 cells, zebrafish, and several other cells with applications in bio-imaging.^{14–22} Additional studies have been carried out in similar alcohol/water mixtures as employed in the bio-imaging studies here.^{20,23–25}

Dipicolylamine (DPA), coumarin, quinoline, bipyridine, triazole, iminodiacetic acid, acyclic and cyclic polyamines, and Schiff-bases are commonly used receptors to detect Zn(II) ion.^{5,26–30} These receptors often contain nitrogen atoms which serve as good binding sites for Zn(II), and when linked with various fluorophores, they can induce effective signaling transduction to recognize the binding event through a variety of photophysical mechanisms. In this regard, we have focused on an easily prepared Schiff-base ligand (one-step synthesis), using readily available reagents, that yields a fluorescent probe based on pyridoxaldehyde. Schiff-bases are generally poor/non-fluorescent due to conventional modes of non-radiative decay such as C=N bond isomerization and/or excited-state intramolecular proton transfer (ESIPT) involving phenolic protons.³¹ Pyridoxaldehyde is one form of vitamin B6 which acts as a co-enzyme in various enzyme reactions and is involved in aerobic metabolism.³² Pyridoxaldehyde also acts as a coenzyme in many enzymatic reactions and is involved in many aspects of macro-nutrient metabolism, neurotransmitter synthesis, histamine synthesis, hemoglobin synthesis and function, and gene expression. This suggests that pyridoxaldehyde is biocompatible and can be readily used in bio-applications. Several Schiff-bases based on pyridoxaldehyde have been reported in the literature that forms complexes with a wide variety of metal cations, including transition metals,^{33–43} heavy metals,^{44,45} lanthanides,^{46–49} and actinides.^{49–51} However, only a few reports show pyridoxal-based Schiff-bases act as fluorometric or colorimetric chemosensors for the detection of transition metals, biomolecules, and anions.^{31,52–64}

In this report, we explore a pyridoxal-based, Zn(II) selective fluorescent probe combined with TRIS that exhibits, ease of synthesis, cost efficiency, chemical and photostability, fluorescence selectivity and sensitivity, and fast response time for spectral imaging of live-cells (Scheme 1). The coordination

environment for Zn(II) ion complexation is clearly delimited by stoichiometric studies and X-ray crystallography, and the TRIS functionality of the Schiff-base provides alternate hydroxyl binding sites for competing metal cations that are more oxophilic than Zn(II), yielding exceptional selectivity and sensitivity.

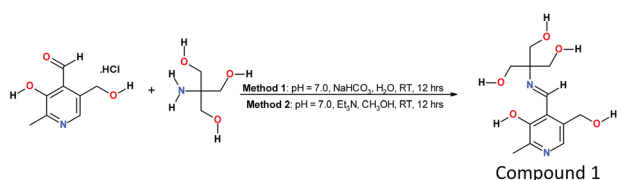
Materials and physical methods

Pyridoxal hydrochloride, 2-amino-2-(hydroxymethyl)-1,3-propanediol (TRIS), and triethylamine were purchased from Sigma-Aldrich and used without further purification. All other materials were of reagent grade. All cations used in this study are perchlorate salts, except tetrakis(acetonitrile)copper(I) tetrafluoroborate. Nuclear magnetic resonance (NMR) spectra were collected in DMSO-*d*₆ at room temperature on a Bruker Ascend™ 400 MHz, and the chemical shifts were reported in part per million (ppm). Fourier transform-infrared spectroscopy (FT-IR) spectra were collected on a Bruker Alpha FT-IR spectrometer, and the wavenumber was reported in cm⁻¹. Element analyses were carried out using an Exeter Analytic CE-440 and helium gas was used as the carrier gas. UV-Vis spectra were collected using a Varian Cary 50 Bio UV-Vis spectrometer and luminescence spectra were collected using a Photon Technology International (PTI) steady-state spectrofluorometer and Fluoromax-4. Binding constant determinations were conducted using Bindfit – Host/Guest modelling software freely available at supramolecular.org.⁸⁰ Electrospray ionization-mass spectroscopy (ESI-MS) data were collected using a Varian 500-MS Ion Trap Mass spectrometer in dilute methanol. Crystallographic data were recorded using a Bruker Venture D8 Apex III diffractometer at 100 K using MoK_α radiation and CuK_α. The crystallographic data were integrated by Apex III software, and the structures were solved using SIR97 (ref. 65) and refined with SHELXL-97 (ref. 66) followed by further refinement and completion of the structure on WinGX⁶⁷ and Mercury 2020.3 Windows was used to visualize the structures. The alcohol group at C12 of **1** was modeled as disordered over two positions with 50 : 50 occupancies. Graphical figures were plotted using OriginPro 9.0 software. The reaction schemes and chemical structure were drawn on BIOVIA Draw for academic use by Dassault Systèmes.

Synthesis of pyridoxal-TRIS Schiff-base ligand (1)

Method 1. 0.5 g (2.45 mmol) of pyridoxal hydrochloride was dissolved in 25 mL of water, and the pH of the solution changed from 3 to 7 by the addition of sodium bicarbonate as monitored by pH indicator strips. 0.3 g (2.45 mmol) of TRIS is added and the reaction mixture was left overnight to stir at room temperature. After the completion of the reaction, the solvent was evaporated and the solid residue dissolved in methanol, and any undissolved solid residue is removed using gravity filtration. Excess diethyl ether is added to the methanol solution to obtain the product by precipitation, which is filtered and washed.

Method 2. 0.2 g (0.98 mmol) of pyridoxal hydrochloride is dissolved in 10 mL of methanol, and the pH of the solution



Scheme 1 Synthesis of pyridoxal-TRIS Schiff's Base.



changed from 3 to 7 by adding approximately 20 drops of triethylamine as monitored by pH indicator strips. 0.12 g (0.98 mmol) of TRIS is added, and the reaction mixture was left overnight at room temperature. After the reaction was complete, excess diethyl ether was added and left in the refrigerator overnight to obtain the product as yellow crystals.

Compound 1. Percent yield: 84% (method 1) and 86% (method 2), melting points: 233–235 °C, molar mass: 270.28 g mol⁻¹, elemental analyses for C₁₂H₁₈N₂O₅: theoretical C = 53.33%; H = 6.71%; N = 10.36% and experimental C = 53.17%; H = 6.69%; N = 10.42%, ¹H NMR (400 MHz, DMSO-*d*₆, 25 °C): δ 2.34 (3H, s, methyl), 3.62–3.64 (6H, dd, *J* = 5.20 Hz, methylene), 4.60–4.62 (2H, dd, *J* = 5.04 Hz, methylene), 4.88–4.91 (3H, t, *J* = 5.34 Hz, alkoxy), 5.30–5.33 (1H, t, *J* = 5.24 Hz, alkoxy), 7.72 (1H, s, aromatic), 8.89 (1H, s, carbimine), 15.44 (1H, s, zwitterion proton), ¹³C NMR (400 MHz, DMSO-*d*₆, 25 °C): δ 162.48, 158.75, 150.76, 134.99, 133.07, 117.89, 68.02, 61.54, 58.96, 193.4, ESI-MS (CH₃OH): 271.2 *m/z* [M + H]⁺.

Synthesis of the Zn complex (2)

A methanolic solution of ligand (0.1 g, 0.37 mmol) was added dropwise to a solution of hydrated zinc perchlorate (Zn(ClO₄)₂·6H₂O, 0.14 g, ~0.37 mmol), and the mixture was stirred at room temperature. The resulting reaction was filtered, and the supernatant liquid was kept at room temperature for slow evaporation. A suitable pale-yellow single-crystal was obtained for X-ray structure determination. Percent yield: 35%.

Synthesis of Cu complex (3)

A 5 mL methanolic solution of ligand (0.1 g, 0.37 mmol) was added dropwise to the 5 mL acetonitrile solution of hydrated copper perchlorate (Cu(ClO₄)₂·6H₂O, 0.14 g, ~0.37 mmol), and the mixture was refluxed for 4 h. Excess of diethyl ether was added to the reaction mixture and left in the refrigerator overnight. A green crystal was obtained, which is dissolved into acetonitrile and diffused into diethyl ether to get an X-ray quality crystal. Percent yield: 82%. Molar mass: 532.72 g mol⁻¹, elemental analyses for C₁₂H₁₈N₂O₁₃Cl₂Cu₁: theoretical C = 27.06%; H = 3.41%; N = 5.26% and experimental C = 27.50%; H = 3.28%; N = 5.36%. ESI-MS (*m/z*) = 553.2 [M + Na]⁺ (cal. 553.93).

UV-Vis and fluorescence studies

Both UV-Vis and fluorescence spectra were collected using the same solutions without altering the conditions. A stock solution of cation and anion salts were prepared in 10 mL (1.2 × 10⁻² M) of acetonitrile or H₂O, and a stock solution of compound **1** in 100 mL (5 × 10⁻⁵ M) in various solvent systems such as 100% methanol, 10% and 50% water (0.1 M HEPES buffer, pH = 7.3) in methanol were prepared. UV-Vis and fluorescence studies were executed by titration of 25 μL aliquots of stock cations or anions salt solution to 3.0 mL of **1** at room temperature. For the stoichiometric and binding constant study of Zn(II) with **1**, the stock solution of Zn(II) is diluted to 6.0 × 10⁻⁴ M, and titration was carried by successive addition of 25 μL aliquots of Zn(II) salt solution to 3.0 mL of **1**.

Quantum yield determination

We used anthracene in ethanol as the reference, and the following equation was used for the calculation of the quantum yield.^{68,69}

$$\phi_{\text{sample}} = \phi_{\text{reference}} \times \frac{A_{\text{sample}}}{A_{\text{reference}}} \times \frac{\text{Abs}_{\text{reference}}}{\text{Abs}_{\text{sample}}} \times \frac{\eta_{\text{sample}}^2}{\eta_{\text{reference}}^2} \quad (1)$$

where, ϕ , Abs, *A*, and η represent quantum yield, absorbance, area under the fluorescence curve, and refractive index of the medium, respectively.

Detection limit determination

The detection limit of **1** for Zn(II) was determined by using fluorescence titration data at 472 nm based on a reported method.⁷⁰ The detection limit was calculated based on eqn (2), where σ is the standard deviation of blank measurement, and *K* is the slope between the fluorescence intensity versus Zn(II) concentration. The fluorescence spectrum of compound **1** was recorded 6 times to achieve the standard deviation of blank.

$$\text{Detection limit} = \frac{3\sigma}{K} \quad (2)$$

Bioimaging study

Human embryonic kidney (HEK) 293 cells were maintained in Dulbecco's Modified Eagle's Medium purchased from Sigma Aldrich Inc. and supplemented with 10% fetal bovine serum (FBS) purchased from Atlas biologicals, DMEM – high glucose medium, and antibiotic/antimycotic solution (100×) purchased from Hyclone laboratories.

Imaging system

Brightfield and fluorescence images were captured at 20× magnification by Leica Microsystems's Leica DMI8 confocal microscope using LAS X software.

Cell culture

The stock solution of compound **1** was prepared in the mixture of 80% methanol and 20% water (0.1 M HEPES buffer, pH = 7.3) and used for incubation in cells. Cells were adherent-cultured in 8-well culture plates for 72 h. Subsequently, the first group was incubated with probe compound **1** (20.0 mg L⁻¹) solution at 37 °C for 40 min, then washed 3 times with phosphate-buffered saline solution (PBS, pH = 7.20). The initial image was acquired using a confocal laser scanning microscope (Leica DMI8) without disturbing the cell positions. The second group was preincubated with probe compound **1** (20.0 mg L⁻¹) solution at 37 °C for 40 min, then with DMEM containing zinc acetate (40.0 mg L⁻¹) for another 40 min, and then washed with PBS for 3 times. These cells were underwent imaging measurement by a confocal microscope (Leica DMI8). The exciting light was 410 nm. The emission range of the blue channel was 420–600 nm.



Results and discussion

Synthesis and characterization

The synthesis of **1** was carried out in two different solvent systems, aqueous and organic solvent (CH₃OH), as shown in Scheme 1, and leads to the same product. In aqueous solvent, the pyridoxal hydrochloride was first neutralized with NaHCO₃ to pH = 7 using pH indicator strips; whereas, in an organic solvent, it is neutralized with triethylamine. In either case, the colorless solution turned yellow in color, and the yield of the product was above 84%. The final product, compound **1**, was recrystallized by slow evaporation in a mixture of methanol and ether and characterized using ¹H NMR, ¹³C NMR, COSY NMR, CHN analyses, ESI-MS, and X-ray crystallography.

¹H NMR of the free ligand (**1**, Fig. S1†) was carried out using DMSO-*d*₆ as a solvent. The ¹H NMR spectra of **1** showed a singlet at 8.89 ppm due to the presence of the CH=N- linkage, which confirms the formation of the Schiff-base. In the free ligand, the phenolic proton has been transferred to the imine nitrogen to form the zwitterion, which has also been confirmed by single-crystal X-ray crystallography (below). A singlet zwitterion proton on the imine nitrogen appears at 15.44 ppm (donated from the nearby phenol group) and the carbimine proton appears at 8.89 ppm and are coupled to each other as indicated

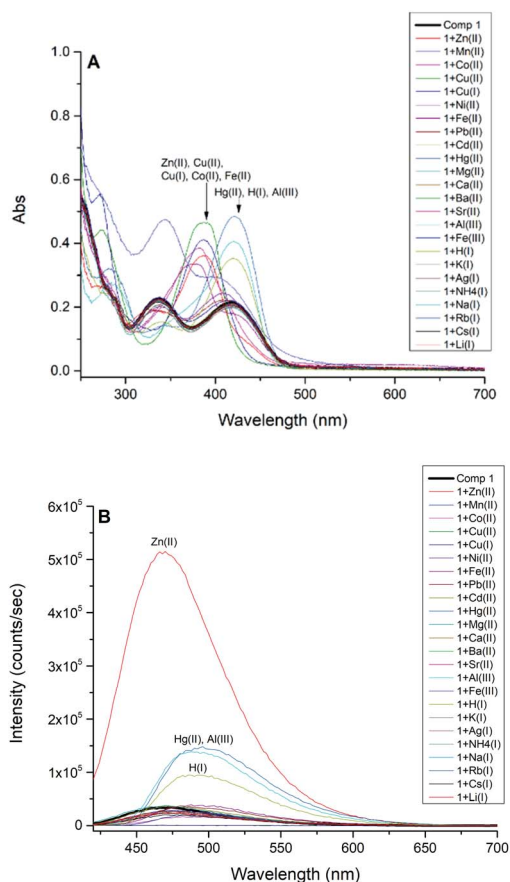


Fig. 1 (A) UV-Vis spectra of compound **1** (5.0×10^{-5} M). (B) Emission spectra of compound **1** (5.0×10^{-5} M, $\lambda_{\text{ext}} = 413$ nm) in CH₃OH before (black) and after the addition of 2.0 equivalents of cations.

by COSY NMR in Fig. S3.† The aromatic proton from the pyridine ring appears at 7.72 ppm. The triplet peaks at 5.32 ppm and 4.89 ppm displayed the presence of the aliphatic -OH groups and the doublet peaks at 4.60 ppm and 3.63 ppm displayed the aliphatic methylene -CH₂ groups from the pyridoxal and TRIS moieties, respectively, with identical coupling for the -CH₂-OH pairs. The singlet peak at 2.34 ppm displays the presence of the methyl group from the pyridoxal ring.

The ¹³C NMR spectra (Fig. S4†) of **1** displayed a sharp characteristic peak due to the azomethine carbon, which resonates at 162.48 ppm. The signal observed at 19.34 ppm belongs to the methyl group. The signals observed at 117.89, 133.07, 134.99, 150.76, and 158.75 ppm are assigned due to pyridoxal ring carbon moieties. The signal observed at 68.02 ppm is due to the quaternary carbon and the signals observed at 58.96 and 61.54 ppm are due to the -CH₂ groups. The ¹³C-DEPT NMR reveals positive signals at 135.0 ppm (aromatic -CH) and 19.3 ppm (aliphatic -CH₃) and negative signals for 58.9 and 61.6 ppm (aliphatic -CH₂ groups for pyridoxal and TRIS).

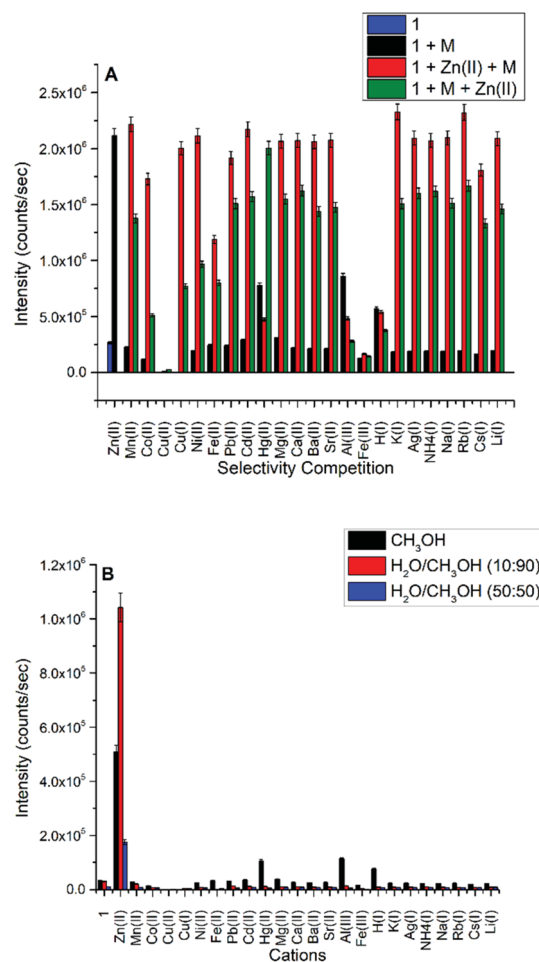


Fig. 2 (A) Selectivity – competition study of **1** with addition of 2.0 equivalents of cations at 470 nm. (B) Selectivity based on the fluorescence intensity profile of **1** with addition of 2.0 equivalents of cations in the different solvent systems at 470 nm. $\lambda_{\text{ext}} = 413$ nm. H₂O contains 0.1 M HEPES buffer, pH = 7.3.



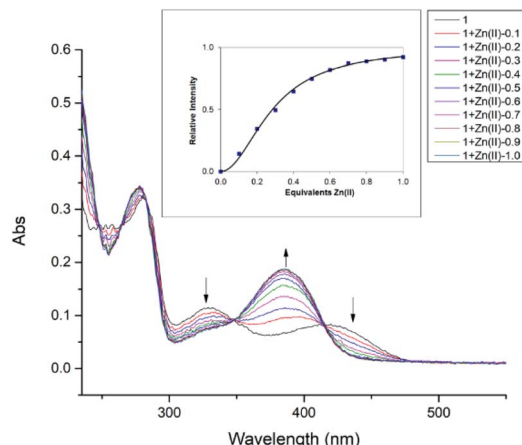


Fig. 3 Changes in the UV-Vis spectra of compound **1** in CH₃OH upon addition of Zn(II) in CH₃CN. Inset: relative absorbance intensity at 385 nm with added equivalents of Zn(II) and best fit for 2 : 1 **1** : Zn(II) association. The association constant for a 2 : 1 best fit equals $1.2 \times 10^9 \text{ M}^{-2}$ ($\pm 25\%$).

Optical properties

The absorbance spectra of **1** in methanol displayed two distinctive absorbance bands centered at 277 nm and 329 nm, which are assigned to $\pi-\pi^*$ transitions, and a third absorbance peak at 420 nm attributed to $n-\pi^*$ transition due to the azomethine group. However, the addition of 2.0 equivalents of Zn(II), Co(II), Cu(II), Cu(I), Fe(II), Hg(II), Al(III), Fe(III), and H(I) induced dramatic bathochromic shifts to longer wavelengths, indicating the formation of ligand-metal adducts with **1**

(Fig. 1A). The fluorescence activity of **1** was also examined by adding 2.0 equivalents of metal salts and collecting the resulting emission spectra (Fig. 1B), indicating high selectivity for Zn(II) ion which exhibits an intense fluorescent band at 470 nm. The addition of Cu(II) quenched the fluorescence intensity of **1** completely. The addition of H(I), Hg(II), and Al(III) have moderate fluorescence enhancements but are not biocompeting cations under physiological conditions.

A competition study for Zn(II) (Fig. 2A) with the same cations shows that Zn(II) selectivity has increased ~ 6 fold over the ligand, and the luminescence, for the most part, is maintained for Zn(II) even in the presence of most of the cations, except for Cu(II), Al(III), Fe(III), Hg(II), and H(I). In the case of Cu(I) and Cu(II), both quenched the weakly fluorescent **1**; however, only Cu(II) can quench the fluorescence of **1** in the presence of Zn(II).

Subsequently, we studied the selectivity of compound **1** towards Zn(II) in different solvent systems (Fig. 2B), including 100% methanol, 90% methanol/10% water (0.1 M HEPES buffer, pH = 7.3), and 50% methanol/50% water (0.1 M HEPES buffer, pH = 7.3). The Schiff base is hydrolyzed by water, hence we tested the fluorescence intensity to detect Zn(II) in different fractions of an aqueous (0.1 M HEPES buffer, pH = 7.3) mixture of CH₃OH and DMSO presented in Fig. S16 and S17[†] respectively which suggest the loss of emission intensity at higher content of water.

The selectivity of Zn(II) in all studied solvent systems is maintained with the highest sensitivity/intensity exhibited using HEPES buffer, potentially due to pH differences between these buffered and unbuffered solvents. Enhanced fluorescence due to H(I), Hg(II), and Al(III) cations were not observed in the

Table 1 Crystallographic data table showing important parameters such as unit cell, crystal system, space group, etc.

	Compound 1	[1] ₂ Zn ²⁺ ·5H ₂ O	[1]Cu ²⁺ ·2ClO ₄ ⁻
Empirical formula	C ₁₂ H ₁₈ N ₂ O ₅	C ₂₄ H ₄₄ N ₄ O ₁₅ Zn	C ₁₂ H ₁₈ N ₂ O ₁₃ Cl ₂ Cu ₁
Formula weight	270.29	694.02	532.72
Temperature (K)	100	100	100
Crystal system	Triclinic	Orthorhombic	Triclinic
Space group	<i>P</i> $\bar{1}$	<i>Pna</i> 2 ₁	<i>P</i> $\bar{1}$
Unit cell (Å, °)	<i>a</i> = 7.5493(3) <i>b</i> = 9.2659(3) <i>c</i> = 9.8603(3) α = 91.084(2) β = 104.586(1) γ = 109.587(1)	<i>a</i> = 20.3659(17) <i>b</i> = 8.7855(9) <i>c</i> = 16.9217(19) α = 90 β = 90 γ = 90	<i>a</i> = 5.8767(23) <i>b</i> = 9.1251(39) <i>c</i> = 18.2139(84) α = 86.630(17) β = 80.725(15) γ = 74.305(14)
Volume (Å ³)	624.83(4)	3027.70(5)	927.90(4)
Z	2	4	2
Density diffraction	1.44	1.522	1.91
Absorption coefficient	0.946	0.889	1.540
<i>F</i> (000)	288.0	1464.0	542.0
θ range	4.7–74.3	2.0–25.4	2.3–26.4
Index ranges	$\pm 9 \pm 11 \pm 12$	$\pm 24 \pm 10 \pm 20$	$\pm 7 \pm 11 \pm 22$
Reflection collected	19 191	23 510	24 629
Independent reflections	2534	5476	3771
Observed reflections	2269	3725	2240
Goodness-of-fit (GOOF)	1.112	1.045	1.031
Final <i>R</i> indices [<i>I</i> > 2 σ (<i>I</i>)]	0.049	0.058	0.079
<i>R</i> indices (all data)	0.053	0.101	0.139
CCDC deposit number	2 095 325	2 095 330	2 095 332



other solvent systems, only in methanol. Compound **1** is highly selective and sensitive towards Zn(II) ion, and acts as a fluorometric and colorimetric sensor with large fluorescence enhancement in the presence of Zn(II), and the color of **1** (yellow in CH₃OH) turns green in the presence of Cu(II), red in the presence of Co(II), and burgundy in the presence of Mn(II).

pH-dependence and reversibility

The enhanced fluorescence response of **1** with Zn(II) in 10% H₂O (0.1 M, HEPES buffer, pH = 7.3) revealed that compound **1** may be pH-dependent. Fig. S10† shows that fluorescence intensity maximizes at pH = 7 and quenches under acidic and basic conditions. We have also studied the UV-Vis and fluorescence responses of **1** with added anions, and relative nondescript spectral changes occur with different strength anionic bases, as shown in Fig. S11 and S12† as well. In addition, reversible binding of guests is a defining principle of chemosensors. Hence, the reversibility of Zn(II) binding with **1** in methanol was investigated using the strong chelating ligand EDTA. The addition of an aqueous solution of EDTA quenched the enhanced fluorescence response of compound **1** with Zn(II) which reappeared with the following addition of Zn(II), as shown in Fig. S8,† supporting the binding of Zn(II) with **1** as reversible. The ESI-MS spectrum of recovered **1** after the addition of an aqueous solution of EDTA shows the molecular ion peaks (*m/z*) of 271.1 [M + H]⁺ as shown in Fig. S9.†

Binding constant determination

To further investigate the interaction between compound **1** with Zn(II), the UV-Vis titration of **1** with an increasing amount of added Zn(II) was carried out in methanol, until the absorbance and fluorescence intensities saturate as shown in Fig. 3 and S15† respectively. The absorption peaks at 329 nm and 420 nm gradually decrease, giving a new strong absorption peak at 385 nm, possibly related to phenolate (O⁻) and Zn(II) coordination. Similar spectral behavior was observed between 240–390 cm⁻¹ for the perturbation of ligand moieties (–(OH)C=C–C=N–) due to the coordination of Zn(II) during the reaction period.³¹ These spectral changes are also accompanied by clean isosbestic points at 265 nm, 282 nm, 348 nm, and 413 nm, which indicates the formation of a single zinc complex. The change in absorption at 385 nm was used to calculate the binding constant as shown in Fig. 3 (inset). The data points (squares) follow a sigmoidal curve for 2 : 1 complexation of compound **1** with Zn(II), with saturation occurring at approximately 0.5 equivalents of added Zn(II). The 2 : 1 association constant represents the equilibrium 2L + Zn(II) ↔ L₂Zn(II) and equals 1.2 × 10⁹ M⁻² (±25%) resulting in the best fit.⁸⁰ The inflection point in Job's plot (Fig. S19†) *via* fluorescence titration at the mole fraction of ~0.33 further confirms the formation of the L₂Zn(II) complex.

Quantum yield and detection limit

The quantum yield of compound **1** and its Zn(II) complex at λ_{ex} = 377 nm in ethanol (referenced to anthracene as standard, φ = 0.27 in ethanol), produced a quantum yield that was ~12 times

larger in the presence of Zn(II) ion (φ = 0.48) compared to the compound **1** (φ = 0.04). These values are comparable to the quantum yields of the pyridoxal derivatives previously reported.³¹ The titration between **1** and Zn(II) results in a linear equation $y = 260.29x + 5649.5$ ($R^2 = 0.992$) in the range of 5–25 × 10⁻⁶ M concentration of Zn(II) at 472 nm as shown in Fig. S13† and the detection limit was found to be 2.8 × 10⁻⁷ M (28 μM) which is observable below the World Health Organizations guideline (76.5 μM) for Zn(II) in water.⁷¹

Crystallography

Single crystals of **1**, the Zn(II) complex, and a related Cu(II) complex were selected for X-ray crystallographic analysis, and data were collected on a Bruker D8 Venture at 100 K. Crystal system, unit cell, space group, and other parameters are presented in Table 1.

Compound **1** adopts a zwitterion structure (see Fig. 4) with the proton transferred from the pyridoxal phenol to the imine nitrogen (N₂). The zwitterion structure is generally observed in pyridoxal-containing derivatives; however, in this case, the

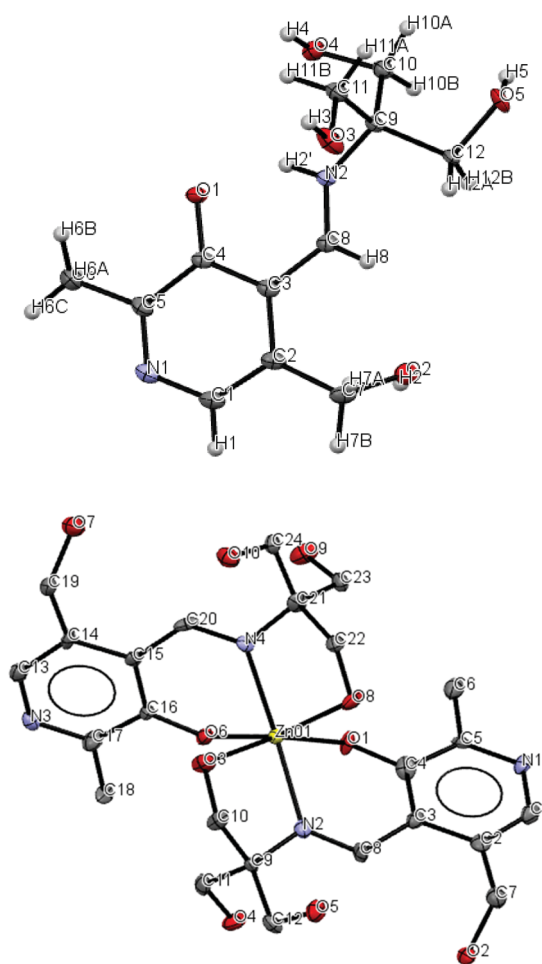


Fig. 4 Thermal ellipsoid diagram (50%) of compound **1** and its Zn(II) complex. (Grey: carbon, red: oxygen, purple: nitrogen, yellow: zinc.) Hydrogen atoms, disordered atoms, and water molecules are not shown for clarity.



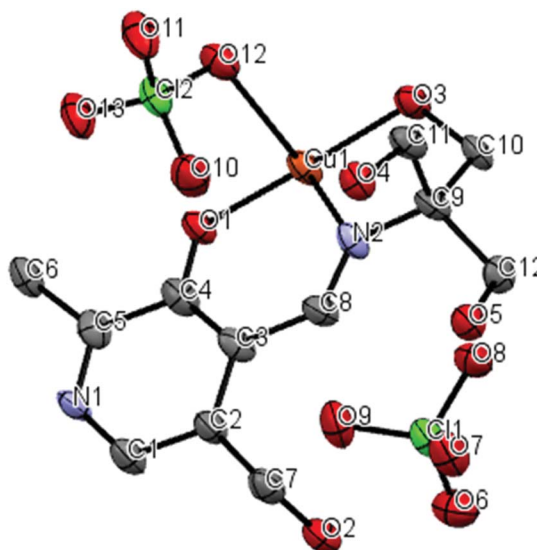
Table 2 Important bond lengths and angles of different connectivity in the crystal structure

	Compound 1	$[1^-]_2\text{Zn}^{2+}\cdot 5\text{H}_2\text{O}$	$[1]\text{Cu}^{2+}\cdot 2\text{ClO}_4^-$
C4–O1	1.302	1.332	1.280
C8–N2	1.287	1.282	1.271
N2–C9	1.471	1.472	1.494
C3–C2	1.419	1.432	1.436
C3–C4	1.416	1.412	1.416
C1–C2	1.370	1.352	1.386
C1–N1	1.357	1.361	1.359
N1–C5	1.327	1.332	1.324
C5–C4	1.443	1.412	1.433
C8–N2–C9	127.29	120.55	127.1
C3–C4–O1	123.07	123.67	126.16
C5–C4–O1	120.19	118.20	116.63
C1–N1–C5	119.78	118.10	125.46

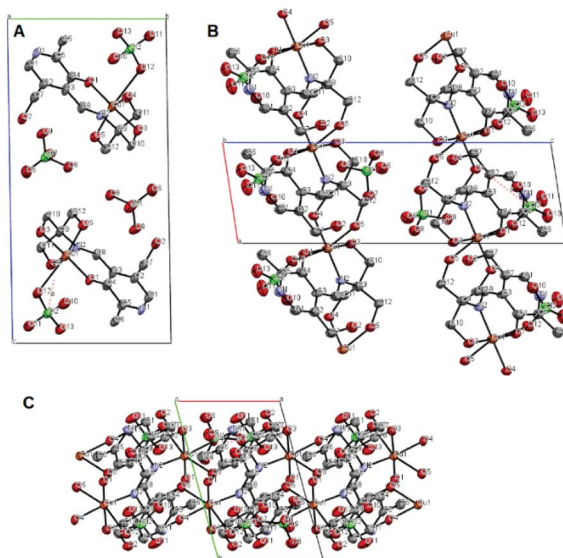
imine nitrogen is protonated instead of the nitrogen atom of the pyridoxal ring.⁷² The bond angles and distances presented in Table 2 are in good agreement with those compounds where the zwitterion is adopted.⁷³ The C8–N2–C9 angle (127.29°) is significantly greater than the normal imine bond angle and is characteristic of the protonated nitrogen. This is also in agreement with the C4–O1 distance (1.302 Å), which is intermediate between C=O distance (1.23 Å) and C–O distance (1.43 Å), and O1–N2 distance (2.587 Å), a suitable distance for intramolecular hydrogen bonding. The C and O atoms of one alkoxy group at C12 are disordered with equal occupancies.

We report the crystal structures of the Zn(II) and Cu(II) complexes of compound 1 to gain comprehensive knowledge of the structural variations of these two metal complexes. As shown in the crystal structure in Fig. 4, the reaction between 1 and hydrated zinc perchlorate in either 1 : 1 or 2 : 1 ratios generated crystals of a complex containing two molecules of 1 and one zinc(II) ion, confirming the previous 2 : 1 stoichiometry. Five water molecules were also found in the structure, hydrogen-bonded to the hydroxyl and nitrogen-containing moieties. The center zinc ion is coordinated through the imine nitrogen and the phenolate and alkoxy oxygen atoms from each of the two ligands, achieving a distorted six-coordinate octahedral geometry. The two deprotonated phenoxy oxygen atoms and two protonated alkoxy oxygen atoms occupy the equatorial position, whereas the two imino nitrogen atoms adopt an axial, trans arrangement. The angle between the mean planes through the pyridine rings of the two ligands in Zn-complex is 78.36° . Other fluorescent probes have been reported which bind with Zn(II),⁷⁴ Cu(II),^{75,76} and Co(II),⁷⁷ and a similar structure ((6-methoxy-2-[tris-(hydroxymethyl)-methyl-imino-methyl]-phenolate), incorporating benzene instead of pyridine) to 1 has been reported by Zhang *et al.*, making a complex with Mn(II).⁷⁸

A crystal structure of the corresponding copper(II) complex reveals a different 1-dimensional polymeric structure of the building-block shown in Fig. 5. In this case, 1 forms a 1 : 1 complex with Cu(II) ion; one molecule of 1 chelate the Cu(II) ion with the imine nitrogen, one phenolic oxygen, and one alkoxy oxygen, combined with two alkoxy oxygen atoms from

**Fig. 5** Thermal ellipsoid diagram (50%); asymmetric unit of copper complex, hydrogen atoms were not shown for clarity.

a neighboring complex, plus one perchlorate ion to form a coordination polymer with distorted six-coordinate octahedral geometry, *i.e.* each ligand is linked to two copper centers through different parts of the molecule, propagating along the *a*-axis (Fig. 6). The pyridine rings of two different ligands are parallel and the distance between them is 3.33 Å. We speculate that Cu(II) ion is more oxophilic and binds to the extra alkoxy groups of the TRIS rather than to another imine functional group.

**Fig. 6** Ortep diagram of the polymeric form of 3 units of $[1]\text{Cu}^{2+}\cdot 2\text{ClO}_4^-$, hydrogen atoms were not shown for clarity, (A) along the *a*-axis, (B) along the *b*-axis, and (C) along the *c*-axis. (Grey: carbon, red: oxygen, purple: nitrogen, orange: copper, and green: chlorine.)

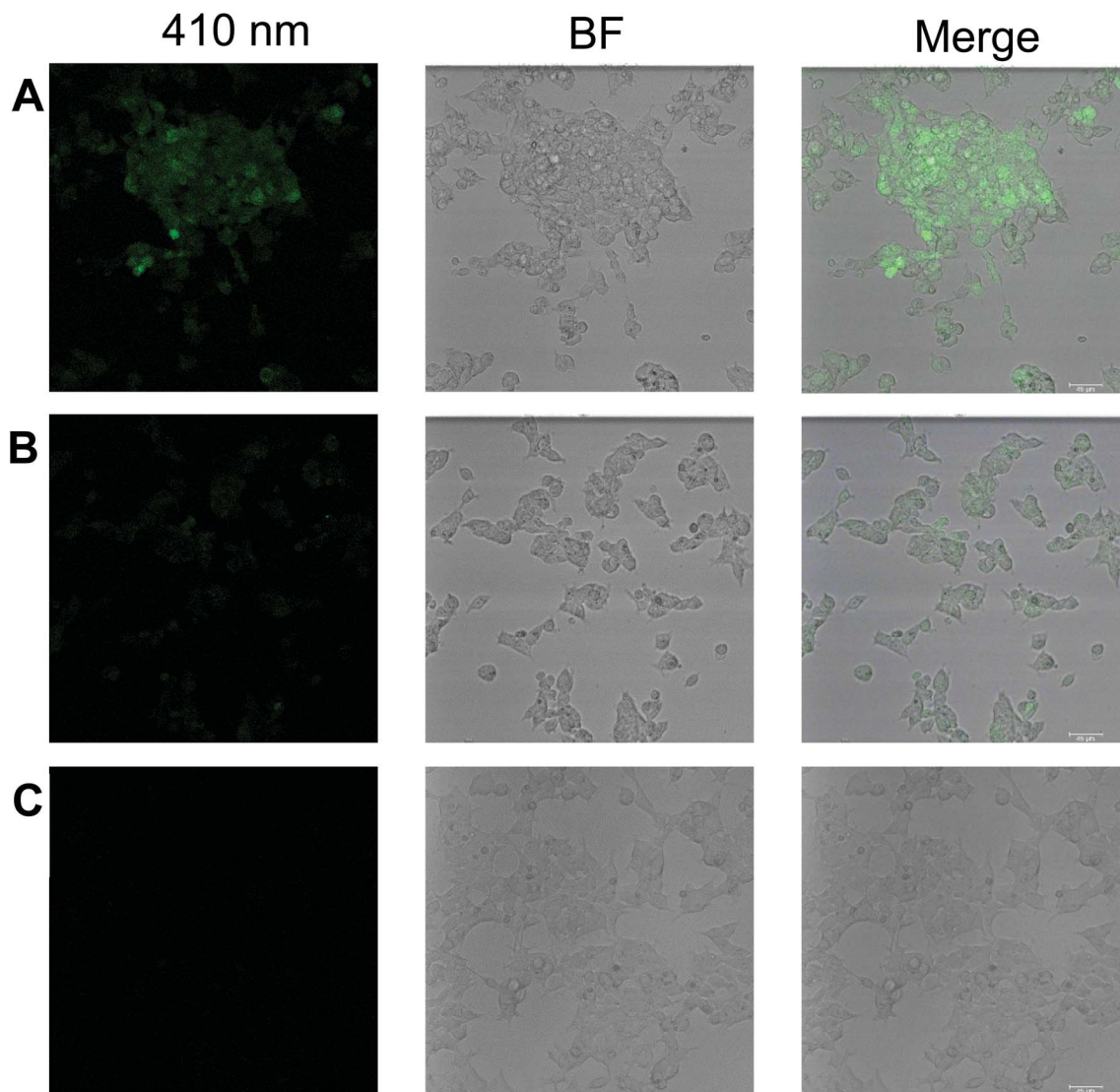


Fig. 7 The fluorescence (410 nm) and brightfield (BF) images of human embryonic kidney cells referred as HEK293 or HEK cells; (A) preincubated with 20 ppm compound **1** at 37 °C for 40 min and then incubated with 40 ppm of zinc acetate at 37 °C for another 40 min, (B) incubated with 20 ppm of compound **1** at 37 °C for 40 min, and (C) before incubation; scale: 45 µm.

In vitro cell studies of compound **1**

We believe that the outstanding turn-on response, sensitivity, selectivity, low detection limit, and reversibility of compound **1** toward zinc ion can find a practical application in biological cell imaging. In this regard, we studied the *in vitro* fluorescence response of **1** to detect zinc ion in human embryonic kidney cells referred as HEK293 or HEK cells. In this study, HEK293 cells are incubated with 20 ppm **1** alone and **1** plus 40 ppm Zn(II), and compared to cells alone using confocal microscopy with corresponding brightfield and fluorescence images with an objective of 20×. Here we find that even without preincubating with Zn(II), compound **1** exhibits weak fluorescence with living cells as shown in Fig. 7B, and a stronger fluorescence signal is observed when 40 ppm zinc ion is incubated along with 20 ppm **1** as shown in Fig. 7A as monitored using the 410 nm blue excitation channel. Cells without **1** and Zn(II) do not show

fluorescence, Fig. 7C. The mean fluorescence intensity (MFI) was calculated according to the method described previously⁷⁹ by using Fiji software as shown in Fig. S18† which suggests the MFI of compound **1** with Zn(II) is increased ~5 fold over compound **1** in HEK293 cells. Previously, pyridoxal-based chemosensors were used for bio-imaging A549, human lung cancer cells, by fluorescence microscopy, but only exhibited fluorescence when incubated together with added zinc ion.³¹ This suggests that **1** has the potential to be used in incubated cells spiked with Zn(II).

Conclusion

The synthesis of compound **1** and the resulting Zn(II) and Cu(II) metal complexes were carried out and fully characterized by standard analytical techniques. Compound **1** was found to be highly selective and sensitive towards Zn(II) in aqueous solution



via fluorescence turn-on response. Zn(II) ion titrations and the crystal structure of the Zn(II) complex showed the formation of a 2 : 1 complex with Zn(II) with a distorted octahedral geometry and a high binding affinity. The crystal structure of the copper complex, the only major competing cation, reveals a different 1 : 1 binding motif and the formation of a coordination polymer instead. The bio-imaging of compound **1** was conducted using HEK 293 cells via blue-channel fluorescence at 410 nm, and even before incubating with added Zn(II) ion, weak fluorescence of these cells was observed. We believe that the outstanding turn-on response and sensitivity toward zinc ion will find future applications in chemical and biological science.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Notes and references

- D. S. Auld, Zinc coordination sphere in biochemical zinc sites, *Zinc Biochemistry, Physiology, and Homeostasis*, 2001, 85–127.
- E. H. Cox and G. L. McLendon, Zinc-dependent protein folding, *Curr. Opin. Chem. Biol.*, 2000, **4**(2), 162–165.
- X. Xie and T. G. Smart, A physiological role for endogenous zinc in rat hippocampal synaptic neurotransmission, *Nature*, 1991, **349**(6309), 521–524.
- L. A. Finney and T. V. O'Halloran, Transition metal speciation in the cell: insights from the chemistry of metal ion receptors, *Science*, 2003, **300**(5621), 931–936.
- E. L. Que, D. W. Domaille and C. J. Chang, Metals in neurobiology: probing their chemistry and biology with molecular imaging, *Chem. Rev.*, 2008, **108**(5), 1517–1549.
- Z. Xu, J. Yoon and D. R. Spring, Fluorescent chemosensors for Zn²⁺, *Chem. Soc. Rev.*, 2010, **39**(6), 1996–2006.
- C. J. Frederickson, Neurobiology of zinc and zinc-containing neurons, *Int. Rev. Neurobiol.*, 1989, **31**, 145–238.
- P. D. Zalewski, S. H. Millard, I. J. Forbes, O. Kapaniris, A. Slavotinek, W. H. Betts, A. D. Ward, S. F. Lincoln and I. Mahadevan, Video image analysis of labile zinc in viable pancreatic islet cells using a specific fluorescent probe for zinc, *J. Histochem. Cytochem.*, 1994, **42**(7), 877–884.
- P. Zalewski, X. Jian, L. Soon, W. Breed, R. Seemark, S. Lincoln, A. Ward and F. Sun, Changes in distribution of labile zinc in mouse spermatozoa during maturation in the epididymis assessed by the fluorophore zinquin, *Reprod., Fertil. Dev.*, 1996, **8**(7), 1097–1105.
- S. Assaf and S.-H. Chung, Release of endogenous Zn²⁺ from brain tissue during activity, *Nature*, 1984, **308**(5961), 734–736.
- B. Bansod, T. Kumar, R. Thakur, S. Rana and I. Singh, A review on various electrochemical techniques for heavy metal ions detection with different sensing platforms, *Biosens. Bioelectron.*, 2017, **94**, 443–455.
- F. Zhou, C. Li, H. Zhu and Y. Li, Simultaneous determination of trace metal ions in industrial wastewater based on UV-vis spectrometry, *Optik*, 2019, **176**, 512–517.
- L. A. Malik, A. Bashir, A. Qureshi and A. H. Pandith, Detection and removal of heavy metal ions: a review, *Environ. Chem. Lett.*, 2019, **17**(4), 1495–1521.
- S. Mondal, S. M. Mandal, D. Ojha, D. Chattopadhyay and C. Sinha, Water soluble sulfaguanidine based Schiff base as a “turn-on” fluorescent probe for intracellular recognition of Zn²⁺ in living cells and exploration for biological activities, *Polyhedron*, 2019, **172**, 28–38.
- X. Wang, G. Ding, Y. Duan, M. Wang, G. Zhu, X. Li, Y. Zhang and X. Qin, Novel ‘naked-eye’ bis-Schiff base fluorescent chemosensors for sensitive detection of Zn²⁺ and bio-imaging in living cells, *Tetrahedron*, 2020, **76**(16), 131108.
- Z.-Y. Li, H.-K. Su, K. Zhou, B.-Z. Yang, T. Xiao, X.-Q. Sun, J. Jiang and L. Wang, Oxo-spirocyclic structure bridged ditopic Schiff base: a turn-on fluorescent probe for selective recognition of Zn(II) and its application in biosensing, *Dyes Pigm.*, 2018, **149**, 921–926.
- M. Fang, S. Xia, J. Bi, T. P. Wigstrom, L. Valenzano, J. Wang, M. Tanasova, R. L. Luck and H. Liu, Detecting Zn(II) ions in live cells with near-infrared fluorescent probes, *Molecules*, 2019, **24**(8), 1592.
- S. Heng, P. Reineck, A. K. Vidanapathirana, B. J. Pullen, D. W. Drumm, L. J. Ritter, N. Schwarz, C. S. Bonder, P. J. Psaltis and J. G. Thompson, Rationally designed probe for reversible sensing of zinc and application in cells, *ACS Omega*, 2017, **2**(9), 6201–6210.
- J. M. Goldberg, F. Wang, C. D. Sessler, N. W. Vogler, D. Y. Zhang, W. H. Loucks, T. Tzounopoulos and S. J. Lippard, Photoactivatable sensors for detecting mobile zinc, *J. Am. Chem. Soc.*, 2018, **140**(6), 2020–2023.
- J. Mandal, P. Ghorai, K. Pal, P. Karmakar and A. Saha, 2-Hydroxy-5-methylisophthalaldehyde based fluorescent-colorimetric chemosensor for dual detection of Zn²⁺ and Cu²⁺ with high sensitivity and application in live cell imaging, *J. Lumin.*, 2019, **205**, 14–22.
- Z. Lu, W. Fan, Y. Lu, C. Fan, H. Zhao, K. Guo, W. Chu and Y. Lu, A highly sensitive fluorescent probe for bioimaging zinc ion in living cells and zebrafish models, *New J. Chem.*, 2018, **42**(14), 12198–12204.
- N. Kumar, V. Bhalla and M. Kumar, Beyond zinc coordination: bioimaging applications of Zn(II)-complexes, *Coord. Chem. Rev.*, 2021, **427**, 213550.
- J. Fu, Y. Chang, B. Li, X. Wang, X. Xie and K. Xu, A dual fluorescence probe for Zn²⁺ and Al³⁺ through differentially



- response and bioimaging in living cells, *Spectrochim. Acta, Part A*, 2020, **225**, 117493.
- 24 S.-Y. Jiao, L.-M. Kong, G.-Q. Liu, X. Jia, J. Tian, Y.-G. Liu, L.-X. Zhang, W.-X. Zhang, Y.-H. Li and Z. Huang, A simple and an easy-to-synthesize turn-on fluorescent probe for rapid detection of Zn²⁺ and its application in bioimaging, *Tetrahedron Lett.*, 2020, **61**(46), 152507.
- 25 L. Yan, X. Wen and Z. Fan, A large-Stokes-shift fluorescent probe for Zn²⁺ based on AIE, and application in live cell imaging, *Anal. Bioanal. Chem.*, 2020, **412**(6), 1453–1463.
- 26 A. P. De Silva, H. N. Gunaratne, T. Gunnlaugsson, A. J. Huxley, C. P. McCoy, J. T. Rademacher and T. E. Rice, Signaling recognition events with fluorescent sensors and switches, *Chem. Rev.*, 1997, **97**(5), 1515–1566.
- 27 P. Jiang and Z. Guo, Fluorescent detection of zinc in biological systems: recent development on the design of chemosensors and biosensors, *Coord. Chem. Rev.*, 2004, **248**(1–2), 205–229.
- 28 P. Carol, S. Sreejith and A. Ajayaghosh, Ratiometric and Near-Infrared Molecular Probes for the Detection and Imaging of Zinc Ions, *Chem.-Asian J.*, 2007, **2**(3), 338–348.
- 29 E. Kimura and S. Aoki, Chemistry of zinc(II) fluorophore sensors, *Zinc Biochemistry, Physiology, and Homeostasis*, 2001, 5–18.
- 30 N. C. Lim, J. V. Schuster, M. C. Porto, M. A. Tanudra, L. Yao, H. C. Freake and C. Brückner, Coumarin-based chemosensors for zinc(II): toward the determination of the design algorithm for CHEF-type and ratiometric probes, *Inorg. Chem.*, 2005, **44**(6), 2018–2030.
- 31 S. Mandal, Y. Sikdar, D. K. Maiti, G. P. Maiti, S. K. Mandal, J. K. Biswas and S. Goswami, A new pyridoxal based fluorescence chemo-sensor for detection of Zn(II) and its application in bio imaging, *RSC Adv.*, 2015, **5**(89), 72659–72669.
- 32 K. M. Kim, T. Qin, Y.-Y. Jiang, L.-L. Chen, M. Xiong, D. Caetano-Anollés, H.-Y. Zhang and G. Caetano-Anollés, Protein domain structure uncovers the origin of aerobic metabolism and the rise of planetary oxygen, *Structure*, 2012, **20**(1), 67–76.
- 33 S. Mondal, M. Chakraborty, A. Mondal, B. Pakhira, A. J. Blake, E. Sinn and S. K. Chattopadhyay, Cu(II) complexes of a tridentate N,N,O-donor Schiff base of pyridoxal: synthesis, X-ray structures, DNA-binding properties and catecholase activity, *New J. Chem.*, 2018, **42**(12), 9588–9597.
- 34 S. Mandal, S. Chatterjee, R. Modak, Y. Sikdar, B. Naskar and S. Goswami, Syntheses, crystal structures, spectral studies, and DFT calculations of two new square planar Ni(II) complexes derived from pyridoxal-based Schiff base ligands, *J. Coord. Chem.*, 2014, **67**(4), 699–713.
- 35 S. A. Elsayed, A. M. Noufal and A. M. El-Hendawy, Synthesis, structural characterization and antioxidant activity of some vanadium(IV), Mo(VI)(IV) and Ru(II) complexes of pyridoxal Schiff base derivatives, *J. Mol. Struct.*, 2017, **1144**, 120–128.
- 36 V. Murašková, N. Szabó, M. Pižl, I. Hoskovcová, M. Dušek, Š. Huber and D. Sedmidubský, Self assembly of dialkoxo bridged dinuclear Fe(III) complex of pyridoxal Schiff base with CC bond formation-structure, spectral and magnetic properties, *Inorg. Chim. Acta*, 2017, **461**, 111–119.
- 37 U. Basu, I. Pant, A. Hussain, P. Kondaiah and A. R. Chakravarty, Iron(III) complexes of a pyridoxal Schiff base for enhanced cellular uptake with selectivity and remarkable photocytotoxicity, *Inorg. Chem.*, 2015, **54**(8), 3748–3758.
- 38 M. Greb, J. Hartung, F. Köhler, K. Špehar, R. Kluge and R. Csuk, The (Schiff base) vanadium(V) complex catalyzed oxidation of bromide – a new method for the in situ generation of bromine and its application in the synthesis of functionalized cyclic ethers, *Eur. J. Org. Chem.*, 2004, (18), 3799–3812.
- 39 S. Bagheri, F. Pazoki, N. Esfandiary, M. M. Fadaei and A. Heydari, Synthesis and characterization of Pd (II)-vitamin B6 complex supported on magnetic nanoparticle as an efficient and recyclable catalyst system for C–N cross coupling of amides in deep eutectic solvents, *Appl. Organomet. Chem.*, 2020, **34**(8), e5723.
- 40 M. Guricová, M. Pižl, Z. Smékal, L. Nádherný, J. Čejka, V. Eigner and I. Hoskovcová, Template synthesis and structure of Co(II), Ni(II), and Cu(II) complexes with pyridoxilydenetaurinate Schiff base ligand, *Inorg. Chim. Acta*, 2018, **477**, 248–256.
- 41 L. A. Fontana, J. D. Siqueira, J. Ceolin, B. A. Iglesias, P. C. Piquini, A. Neves and D. F. Back, Peroxidase activity of new mixed-valence cobalt complexes with ligands derived from pyridoxal, *Appl. Organomet. Chem.*, 2019, **33**(7), e4903.
- 42 S. Naskar, S. Naskar, R. J. Butcher and S. K. Chattopadhyay, Synthesis, X-ray crystal structures and spectroscopic properties of two Ni(II) complexes of pyridoxal Schiff's bases with diamines: importance of steric factor in stabilization of water helices in the lattices of metal complex, *Inorg. Chim. Acta*, 2010, **363**(2), 404–411.
- 43 S. Naskar, S. Naskar, H. M. Figgie, W. S. Sheldrick and S. K. Chattopadhyay, Synthesis, crystal structures and spectroscopic properties of two Zn(II) Schiff's base complexes of pyridoxal, *Polyhedron*, 2010, **29**(1), 493–499.
- 44 J. M. Galván-Hidalgo, T. Ramírez-Apan, A. Nieto-Camacho, S. Hernández-Ortega and E. Gómez, Schiff base Sn(IV) complexes as cytotoxic agents: synthesis, structure, isosteric and bioisosteric replacement, *J. Organomet. Chem.*, 2017, **848**, 332–343.
- 45 S. Asijaa, N. Malhotra and R. Malhotra, Bioactive penta-coordinated diorganotin(IV) complexes of pyridoxalimine Schiff bases, *Phosphorus, Sulfur Silicon Relat. Elem.*, 2012, **187**(12), 1510–1520.
- 46 M. Pižl, O. Jankovský, P. Ulbrich, N. Szabó, I. Hoskovcová, D. Sedmidubský and V. Bartůněk, Facile preparation of nanosized yttrium oxide by the thermal decomposition of amorphous Schiff base yttrium complex precursor, *J. Organomet. Chem.*, 2017, **830**, 146–149.
- 47 M. K. Nair, P. S. Ajitha and P. K. Rejimon, Synthesis, Structural Characterization and Antibacterial studies of new Schiff base 5-(Hydroxymethyl)-4-((1Z)-[2-N(3-



- hydroxyphenyl]ethanimidoyl)-2-ethyl}pyridine-3-ol-hydrochloride and its Lanthanide(III) perchlorate complexes, *Res. J. Pharm., Biol. Chem. Sci.*, 2013, **4**, 1145–1153.
- 48 M. Chakraborty, S. Mondal, C. Cardin, A. L. Rheingold, C. D. Mukhopadhyay and S. K. Chattopadhyay, Yb(III), Sm(III) and La(III) complexes of a tetradentate pyridoxal Schiff base ligand: their DNA-binding activity and bio-imaging applications, *Polyhedron*, 2020, **175**, 114167.
- 49 D. F. Back, E. Bonfada, G. M. de Oliveira and E. S. Lang, Chelation of ThIV, EuIII and NdIII by dianionic N,N'-bis(pyridoxylideneiminato)ethylene, (Pyr₂en)²⁻. On the search of feasible modelings for heavy metals damage inhibition in living beings, *J. Inorg. Biochem.*, 2007, **101**(4), 709–714.
- 50 D. F. Back, G. M. de Oliveira, E. S. Lang and J. P. Vargas, Assembly of new Schiff base ligands derived from vitamin B6 and stabilization through complexation of N,N'-bis(pyridoxylideneimine)-o-phenylene: synthesis and X-ray structural features of pyridoxal/o-phenylenediamine adducts and of [UO₂(H₂pyr₂phen)Cl]NO₃ and [UO₂(Hpyr₂phen)Cl]₂{pyr₂phen = N,N'-bis(pyridoxylideneiminato)²⁻ phenylene}, *Polyhedron*, 2008, **27**(12), 2551–2556.
- 51 D. F. Back, G. M. de Oliveira and E. S. Lang, Chelation of UO₂²⁺ by vitamin B6 complex derivatives: synthesis and characterization of [UO₂(β-pyracinate)₂(H₂O)] and [UO₂(Pyr₂en)DMSO]Cl₂{Pyr₂en=N,N'-ethylenebis(pyridoxylideneiminato)}. A useful modeling of assimilation of uranium by living beings, *J. Inorg. Biochem.*, 2006, **100**(10), 1698–1704.
- 52 S. Mandal, S. K. Mandal, A. R. Khuda-Bukhsh and S. Goswami, Pyridoxal based fluorescent chemosensor for detection of copper(II) in solution with moderate selectivity and live cell imaging, *J. Fluoresc.*, 2015, **25**(5), 1437–1447.
- 53 C. Das, B. Pakhira, A. L. Rheingold and S. K. Chattopadhyay, Turn on ESIPT based chemosensor for histidine: application in urine analysis and live cell imaging, *Inorg. Chim. Acta*, 2018, **482**, 292–298.
- 54 D. Sharma, A. Moirangthem, R. Kumar, S. A. Kumar, A. Kuwar, J. F. Callan, A. Basu and S. K. Sahoo, Pyridoxal-thiosemicarbazide: its anion sensing ability and application in living cells imaging, *RSC Adv.*, 2015, **5**(63), 50741–50746.
- 55 D. Sharma, S. Bothra, S. M. Roy, R. Kumar, J. P. Nandre, U. D. Patil and J. F. Callan, Pyridoxal derived chemosensor: its application in anion sensing and molecular logic gate building, *Indian J. Chem., Sect. A: Inorg., Bio-inorg., Phys., Theor. Anal. Chem.*, 2020, **55**(1), 44–50.
- 56 D. Sharma, S. K. Sahoo, S. Chaudhary, R. K. Bera and J. F. Callan, Fluorescence 'turn-on' sensor for F⁻ derived from vitamin B6 cofactor, *Analyst*, 2013, **138**(13), 3646–3650.
- 57 D. Sharma, S. A. Kumar and S. K. Sahoo, Vitamin B6 cofactor derived chemosensor for the selective colorimetric detection of acetate anions, *Tetrahedron Lett.*, 2014, **55**(4), 927–930.
- 58 S. K. Sahoo, D. Sharma, A. Moirangthem, A. Kuba, R. Thomas, R. Kumar, A. Kuwar, H.-J. Choi and A. Basu, Pyridoxal derived chemosensor for chromogenic sensing of Cu²⁺ and fluorogenic sensing of Fe³⁺ in semi-aqueous medium, *J. Lumin.*, 2016, **172**, 297–303.
- 59 S. Bothra, R. Kumar and S. K. Sahoo, Pyridoxal conjugated gold nanoparticles for distinct colorimetric detection of chromium(III) and iodide ions in biological and environmental fluids, *New J. Chem.*, 2017, **41**(15), 7339–7346.
- 60 Y. Upadhyay, T. Anand, L. T. Babu, P. Paira, G. Crisponi, R. Kumar and S. K. Sahoo, Three-in-one type fluorescent sensor based on a pyrene pyridoxal cascade for the selective detection of Zn(II), hydrogen phosphate and cysteine, *Dalton Trans.*, 2018, **47**(3), 742–749.
- 61 C. Xu, L. Liao, Y. He, R. Wu, S. Li and Y. Yang, Resonance light scattering detection of uranium based on its reaction with a Schiff base containing tetradentate ligand and phosphate groups to form supramolecular polymer, *J. Radioanal. Nucl. Chem.*, 2015, **304**(3), 1163–1169.
- 62 S. Mondal, P. Adak, C. Das, S. Naskar, B. Pakhira, A. L. Rheingold, E. Sinn, C. S. Eribal and S. K. Chattopadhyay, Synthesis, X-ray crystal structures, spectroscopic and electrochemical studies of Zn(II), Cd(II), Ni(II) and Mn(II) complexes of N1,N4-bis(pyridoxylidene) triethylenetetramine, *Polyhedron*, 2014, **81**, 428–435.
- 63 T. Mukherjee, J. C. Pessoa, A. Kumar and A. R. Sarkar, Synthesis, spectroscopic characterization, insulin-enhancement, and competitive DNA binding activity of a new Zn(II) complex with a vitamin B6 derivative—a new fluorescence probe for Zn(II), *Dalton Trans.*, 2012, **41**(17), 5260–5271.
- 64 M. Strianese, S. Milione, V. Bertolasi and C. Pellecchia, Iron and manganese pyridoxal-based complexes as fluorescent probes for nitrite and nitrate anions in aqueous solution, *Inorg. Chem.*, 2013, **52**(20), 11778–11786.
- 65 A. Altomare, M. C. Burla, M. Camalli, G. L. Casciarano, C. Giacovazzo, A. Guagliardi, A. G. Moliterni, G. Polidori and R. Spagna, SIR97: a new tool for crystal structure determination and refinement, *J. Appl. Crystallogr.*, 1999, **32**(1), 115–119.
- 66 G. M. Sheldrick and T. R. Schneider, [16] SHELXL: high-resolution refinement, in *Methods in enzymology*, Elsevier, 1997, vol. 277, pp. 319–343.
- 67 L. J. Farrugia, WinGX suite for small-molecule single-crystal crystallography, *J. Appl. Crystallogr.*, 1999, **32**(4), 837–838.
- 68 G. Jones, W. R. Jackson, C. Y. Choi and W. R. Bergmark, Solvent effects on emission yield and lifetime for coumarin laser dyes. Requirements for a rotatory decay mechanism, *J. Phys. Chem.*, 1985, **89**(2), 294–300.
- 69 J. R. Lakowicz, *Principles of fluorescence spectroscopy*, Springer Science & Business Media, 2013.
- 70 A. Shrivastava and V. Gupta, Methods for the determination of limit of detection and limit of quantitation of the analytical methods, *Chron. Young Sci.*, 2011, **2**(1), 21.
- 71 A. Kim, H. Lee, D. Yun, U. Jung, K.-T. Kim and C. Kim, Developing a new chemosensor targeting zinc ion based on two types of quinoline platform, *Spectrochim. Acta, Part A*, 2020, **241**, 118652.



- 72 V. M. Leovac, M. D. Joksović, V. Divjaković, L. S. Jovanović, Ž. Šaranović and A. Pevec, Synthesis, spectroscopic and X-ray characterization of a copper(II) complex with the Schiff base derived from pyridoxal and aminoguanidine: NMR spectral studies of the ligand, *J. Inorg. Biochem.*, 2007, **101**(7), 1094–1097.
- 73 M. F. Belicchi, G. F. Gasparri, E. Loporati, C. Pelizzi, P. Tarasconi and G. Tosi, Thiosemicarbazones as coordinating agents. Solution chemistry and X-ray structure of pyridoxal thiosemicarbazone trihydrate and spectroscopic properties of its metal complexes, *J. Chem. Soc., Dalton Trans.*, 1986, **11**, 2455–2461.
- 74 F. Wang, K. Wang, Q. Kong, J. Wang, D. Xi, B. Gu, S. Lu, T. Wei and X. Chen, Recent studies focusing on the development of fluorescence probes for zinc ion, *Coord. Chem. Rev.*, 2020, 213636.
- 75 S. Sharma and K. S. Ghosh, Overview on recently reported fluorometric sensors for the detection of copper ion based on internal charge transfer (ICT), paramagnetic effect and aggregation induced emission (AIE) mechanisms, *J. Mol. Struct.*, 2021, **1237**, 130324.
- 76 D. Udhayakumari, S. Naha and S. Velmathi, Colorimetric and fluorescent chemosensors for Cu²⁺. A comprehensive review from the years 2013–15, *Anal. Methods*, 2017, **9**(4), 552–578.
- 77 J. Liu, J. Liang, C. Wang and J. Ma, Electrospun CoSe@N-doped carbon nanofibers with highly capacitive Li storage, *J. Energy Chem.*, 2019, **33**, 160–166.
- 78 X. Zhang, P. Wei, J. Dou, B. Li and B. Hu, Bis{2-methoxy-6-[tris(hydroxymethyl)methyliminomethyl]phenolato-[kappa]³O,N,O'}manganese(II) dimethanol solvate hemihydrate, *Acta Crystallogr., Sect. E: Struct. Rep. Online*, 2009, **65**(3), m293–m294.
- 79 M. H. Shihan, S. G. Novo, S. J. Le Marchand, Y. Wang and M. K. Duncan, A simple method for quantitating confocal fluorescent images, *Biochemistry and Biophysics Reports*, 2021, **25**, 100916.
- 80 D. B. Hibbert and P. Thordarson, The death of the Job plot, transparency, open science and online tools, uncertainty estimation methods and other developments in supramolecular chemistry data analysis, *Chem. Commun.*, 2016, **52**, 12792–12805.

