Supporting Information

A Sole Multi-Analyte receptor responds with three distinct fluorescence signals: Traffic signal like sensing of Al³⁺, Zn²⁺ and F⁻

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Fig. S1: Changes in the absorption spectra of L upon the addition of different metal ions. INSET: Visual colour change upon the addition of Al^{3+} and Zn^{2+} to L.



Fig. S2: Changes in the emission spectra of L upon the addition of different metal ions. INSET: Visual colour change upon the addition of Al^{3+} and Zn^{2+} to L under UV lamp (λ_{ex} = 365 nm).



Fig. S3: Changes in the absorption spectra of L upon the addition of different anions.



Fig. S4: Changes in the emission spectra of L upon the addition of different anions.



Fig S5: Normalized fluorescence responses of L (10 μ M) to various cations in mixed solvent. The red bars represent the emission intensities of L in the presence of cations of interest (5 eqv.). The black bars represent the change of the emission that occurs upon the subsequent addition of Al³⁺ to the above solution.



Fig S6: Normalized fluorescence responses of L (10 μ M) to various cations in mixed solvent. The red bars represent the emission intensities of L in the presence of cations of interest (5 eqv.). The black bars represent the change of the emission that occurs upon the subsequent addition of Zn²⁺ to the above solution.



Fig S7: Job's plot between L and Al^{3+} ions. X_{Host} is the mole fraction of L and ΔI is the change (I-I₀) in the intensity of the emission spectra in presence of guest i.e; Al^{3+} .



Fig S8: Job's plot between L and Zn^{2+} ions. X_{Host} is the mole fraction of L and ΔI is the change (I-I₀) in the intensity of the emission spectra in presence of guest i.e; Zn^{2+} .



Fig S9: Bensei-Hildebrand plot obtained for Al³⁺ from the emission experiment (emission intensity calculated from 500 nm) studies.



Fig S10: Bensei-Hildebrand plot obtained for Zn^{2+} from the emission experiment (emission intensity calculated from 550 nm) studies.



Fig S11: Effect of pH on the fluorescence intensity of L.



Fig S12: MTT assay to determine the cytotoxic effects of compounds L, L–Al and L–Zn complex on HeLa cells.



Fig S13: ¹H-NMR spectra of L in CDCl₃.



Fig S14: Expanded ¹H-NMR spectra of L in CDCl₃.



Fig S15: ¹³C-NMR spectra of L in CDCl₃.



Fig S16: Mass spectrum of **L**, Calculated $[L +H]^+= 503.1832$, Found 503.1867 (Mass spectrum obtained in positive mode).



Fig. S17: ¹H-NMR titration spectras of L with Al³⁺ in DMSO-d₆.



Fig. S18: ¹H-NMR titration spectras of L with Zn^{2+} in DMSO-d₆.



Fig. S19: ¹H-NMR titration spectras of L with F⁻ in CDCl₃.



Fig. S20: Crystal structure of L and various interactions presents in it.