

Use of Rhodamine-allyl Schiff base by chemodosimetric process for total Palladium estimation and application for live cell imaging

Anup Kumar Bhanja, Snehasis Mishra, Ketaki Kar, Kaushik Naskar, Suwendu Maity, Krishna Das Saha, Chittaranjan Sinha*

Fig.S1: Powder X-ray diffraction pattern of RD-2.

Fig.S2. FT-IR spectrum of RD-2 in KBr disc

Fig.S3. ¹HNMR spectrum of RD-2 in CDCl₃

Fig.S4. ¹³CNMR spectrum of RD-2 in CDCl₃

Fig.S5. Mass spectrum of RD-2

Fig.S6. Fluorescence intensity changes profiles of 100 μM RD-2 in CH₃CN–water (HEPES buffer, pH =7.4; v/v, 1/4) in presence of selected metal ions at excitation wavelength 505 nm.

Fig.S7. The linear dynamic response of RD-2 for Pd²⁺ and the determination of the detection limit (LOD) for Pd²⁺.

Fig.S8. Plausible mechanism of Pd²⁺ induced spirolactam ring opening and fluorescence emission change strategy of probe(our previous report).

Fig.S9. Mass spectrum of RD-2 with Pd²⁺.

Fig.S10. ¹HNMR spectrum (300MHz) of RD-2 in CD₃CN with Pd²⁺.

Fig.S11. FT-IR spectrum of RD-2 in KBr disc with Pd²⁺.

Fig.S12. ¹³C NMR spectrum of RD-2 in CD₃CN-CDCl₃ with Pd²⁺.

Fig.S13. Effect of pH on the fluorescence activity of RD and RD with Pd²⁺ in (CH₃CN/H₂O, ¼, v/v, HEPES buffer).

Table S1. Comparative information of different probes for detection of Pd²⁺ and their LOD and Reference.

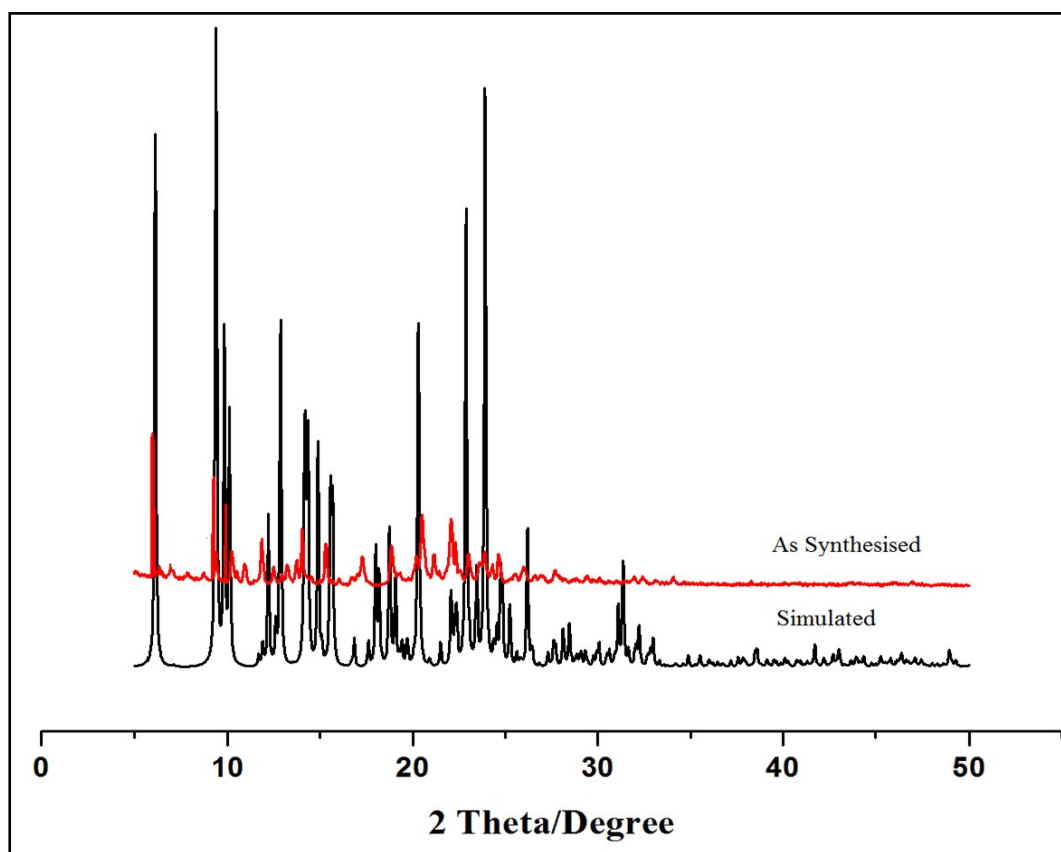


Fig S1: Powder X-ray diffraction pattern of RD-2.

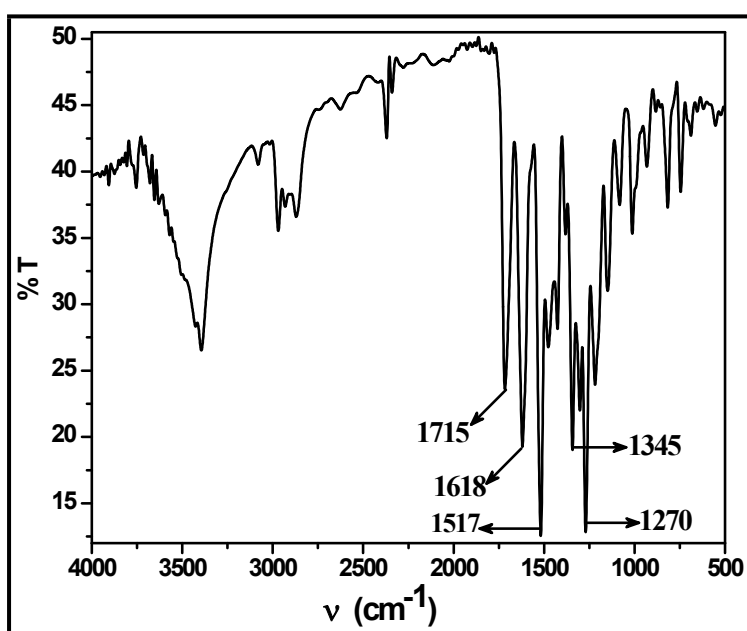


Fig. S2. FT-IR spectrum of RD-2 in KBr disc.

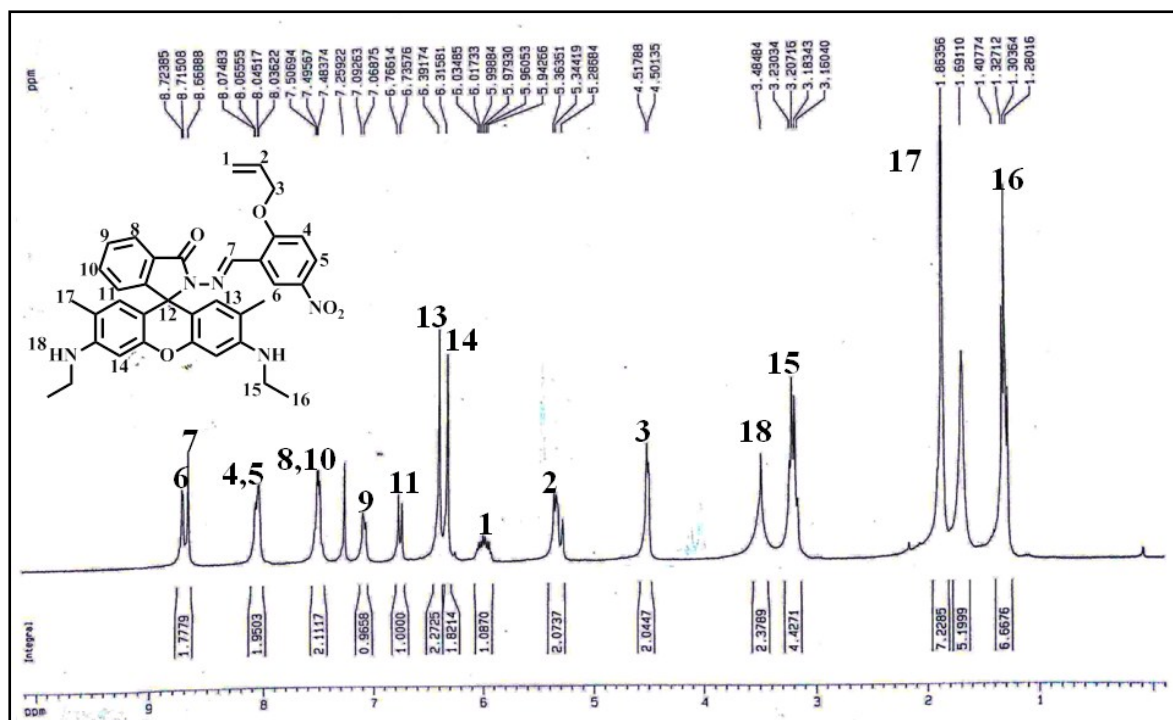


Fig. S3. ^1H NMR spectrum(300MHz) of RD-2 in CDCl_3 .

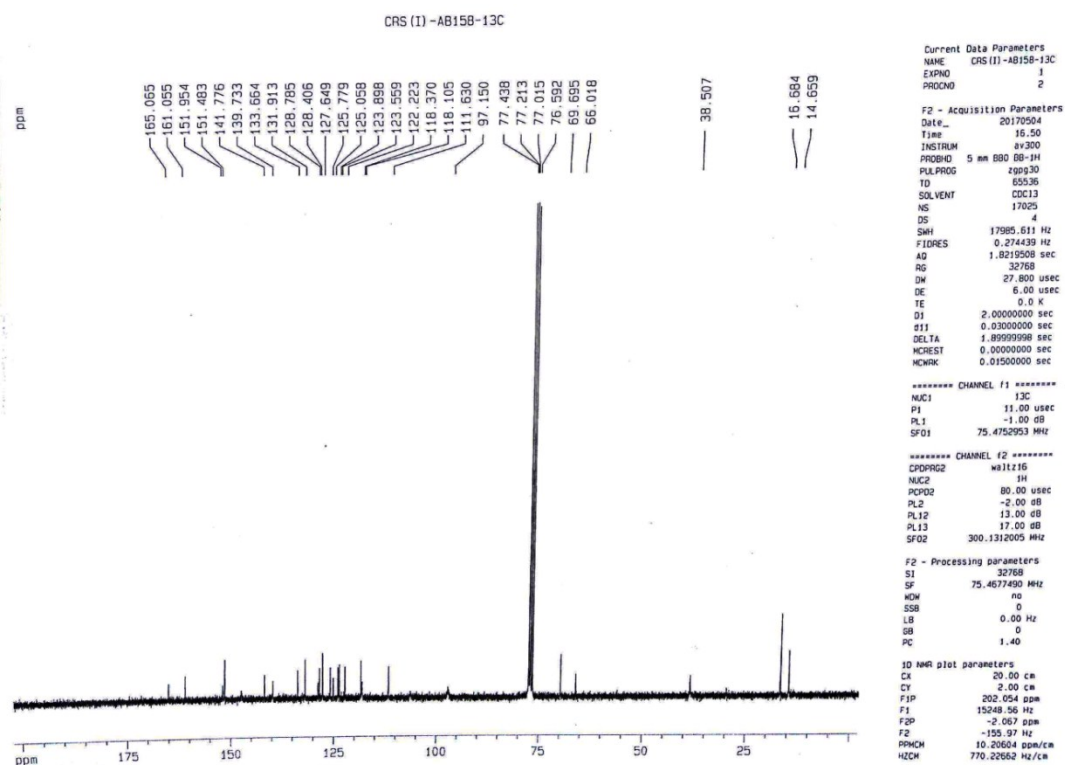


Fig. S4. ^{13}C NMR spectrum of RD-2 in CDCl_3

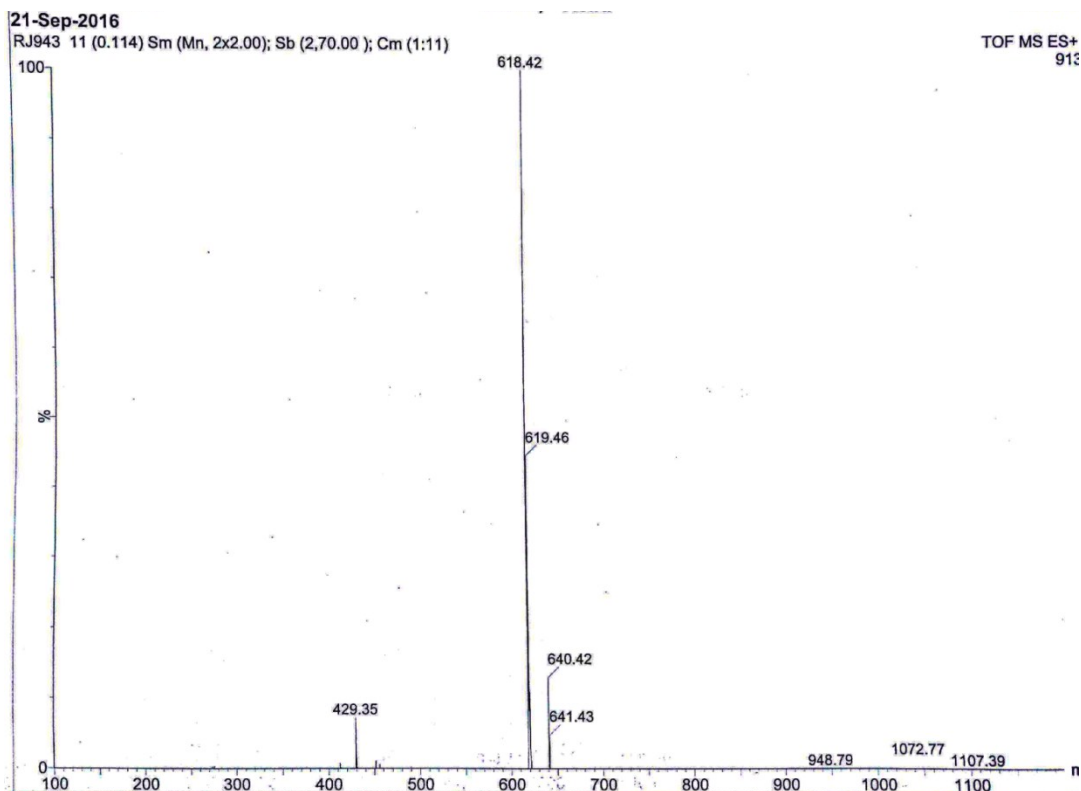


Fig. S5. Mass spectrum of RD-2.

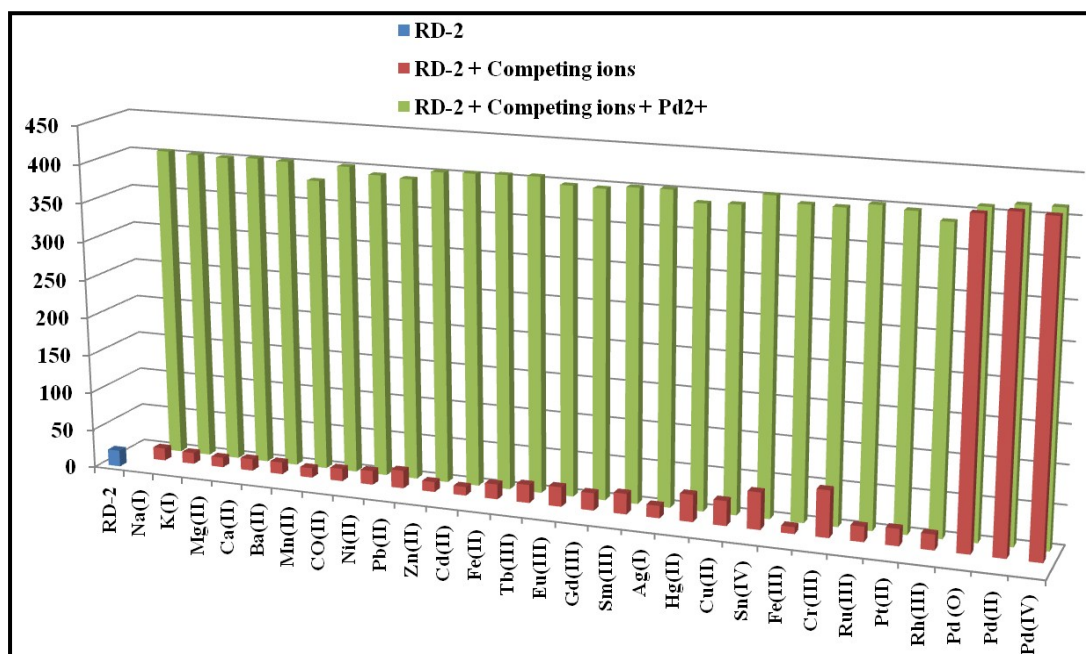


Fig. S6. Fluorescence intensity changes profiles of 100 μM RD-2 in CH_3CN –water (HEPES buffer, pH =7.4; v/v, 1/4) in presence of selected metal ions at excitation wavelength 505 nm.

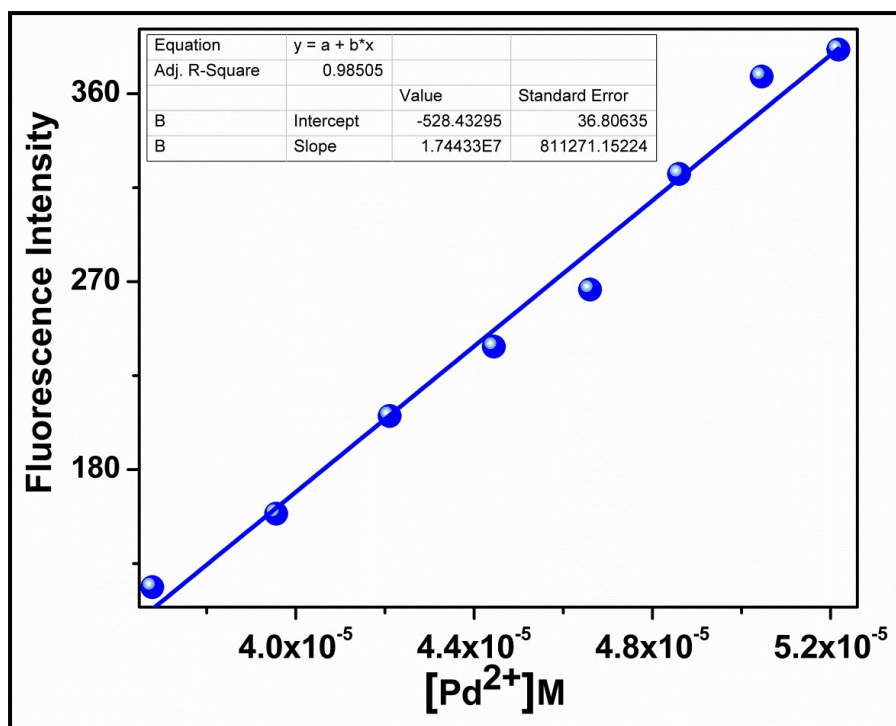


Fig. S7. The linear dynamic response of RD-2 for Pd^{2+} and the determination of the detection limit (LOD) for Pd^{2+} .

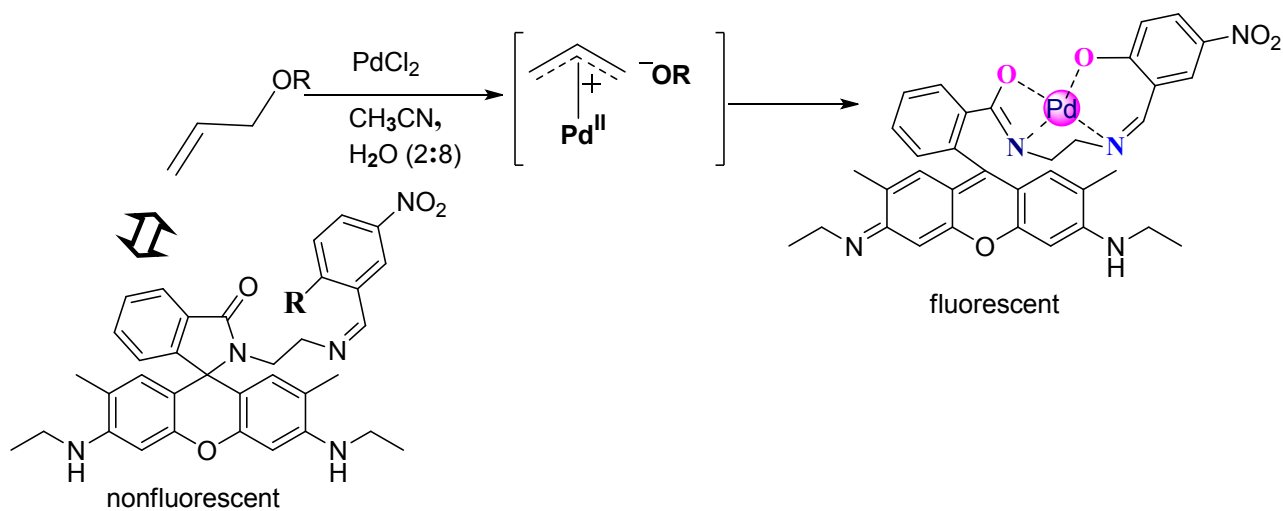


Fig. S8. Plausible mechanism of Pd^{2+} induced spiro lactam ring opening and fluorescence emission change strategy of probe (our previous report).

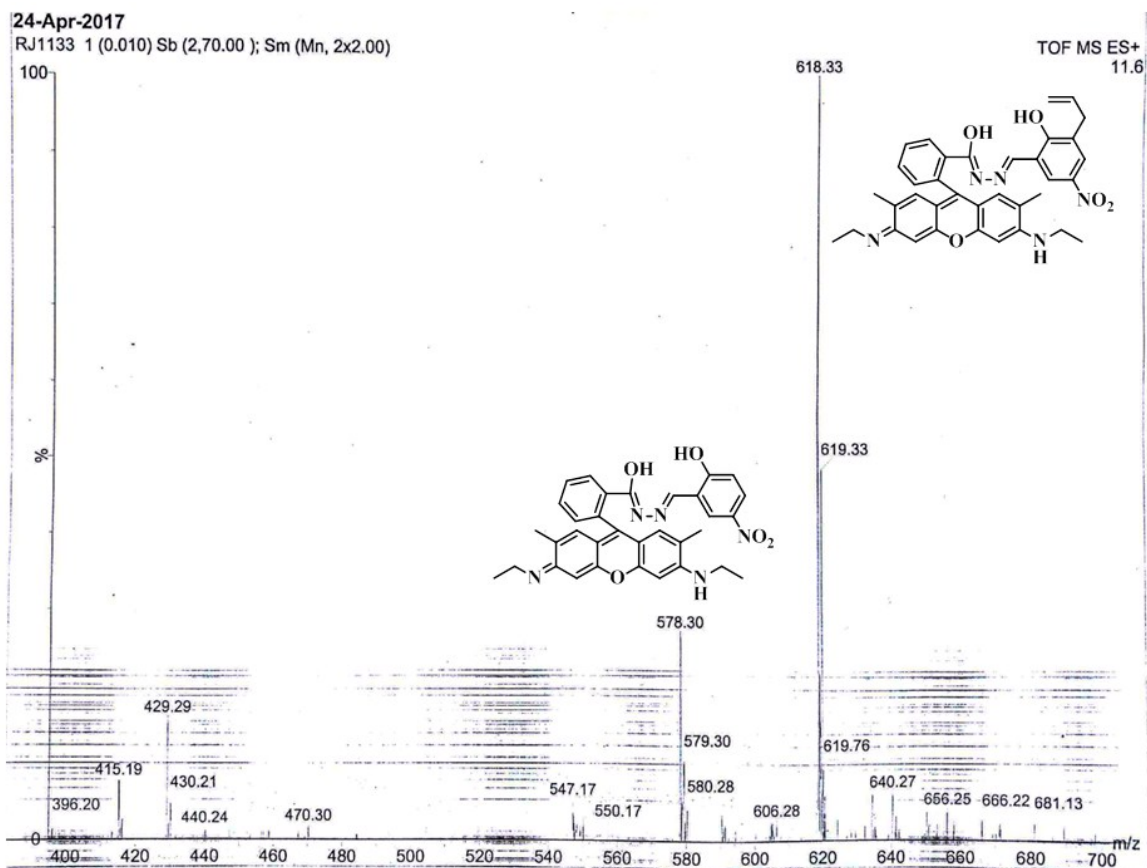


Fig. S9. Mass spectrum of RD-2 with Pd²⁺.

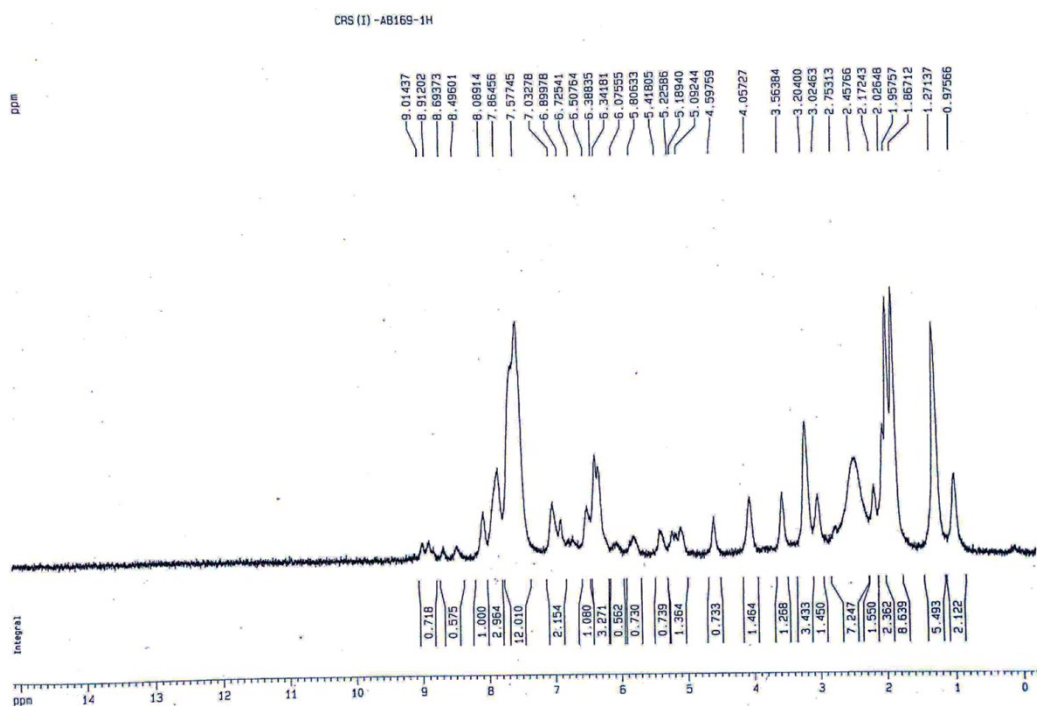


Fig. S10. ^1H NMR spectrum (300MHz) of RD-2 in CD_3CN with Pd^{2+} .

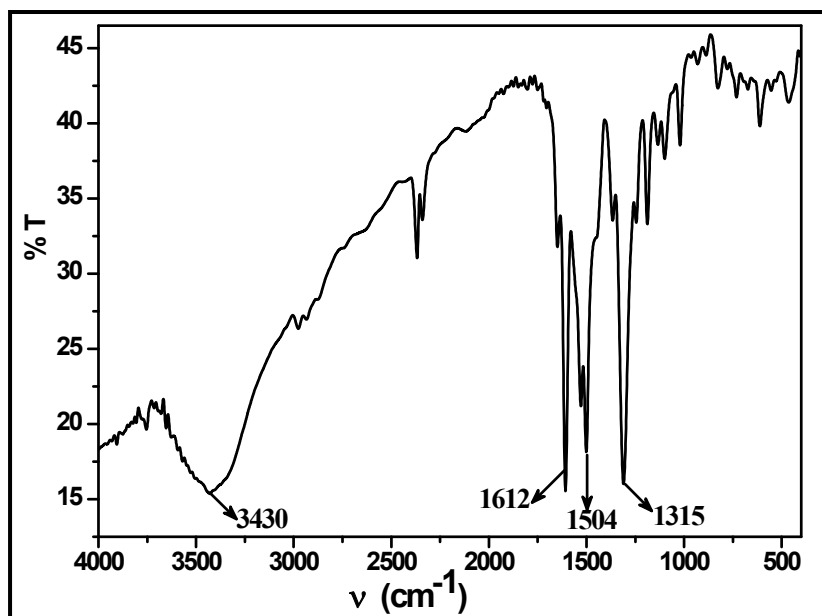


Fig. S11. FT-IR spectrum of RD-2 in KBr disc with Pd^{2+} .

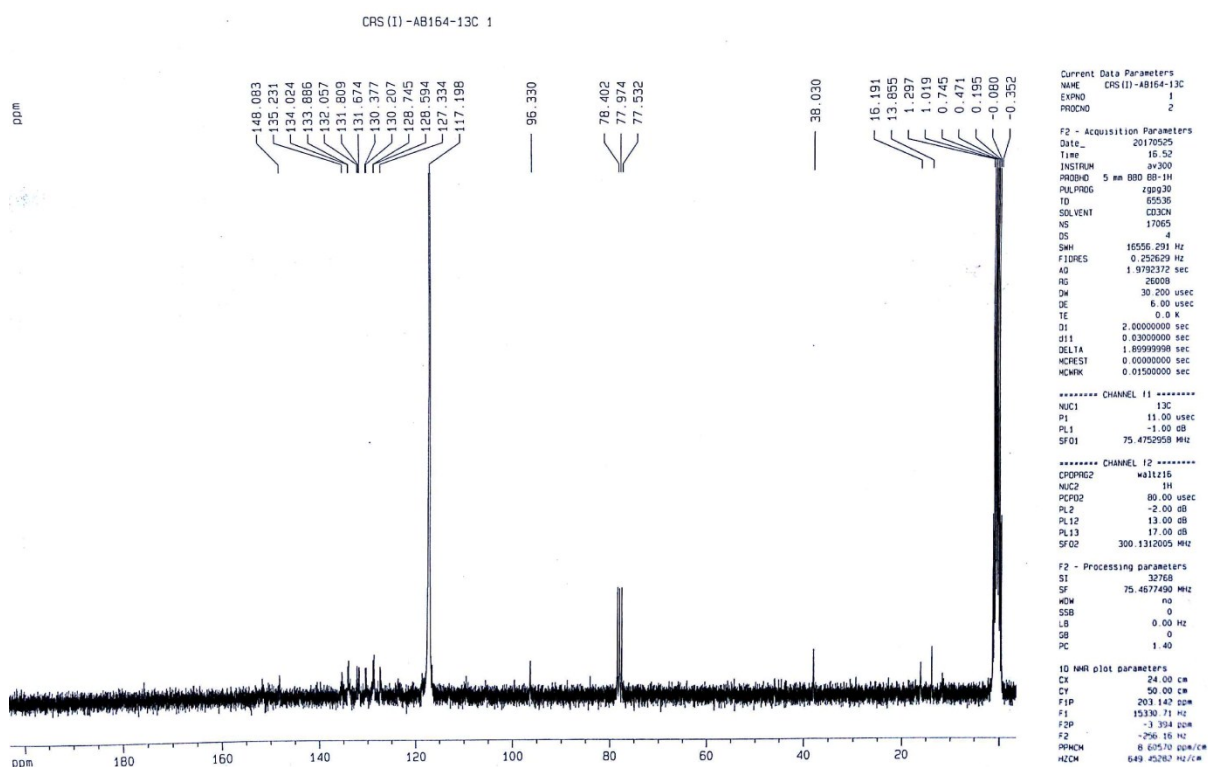


Fig. S12. ^{13}C NMR spectrum of RD-2 in $\text{CD}_3\text{CN}-\text{CDCl}_3$ with Pd^{2+} .

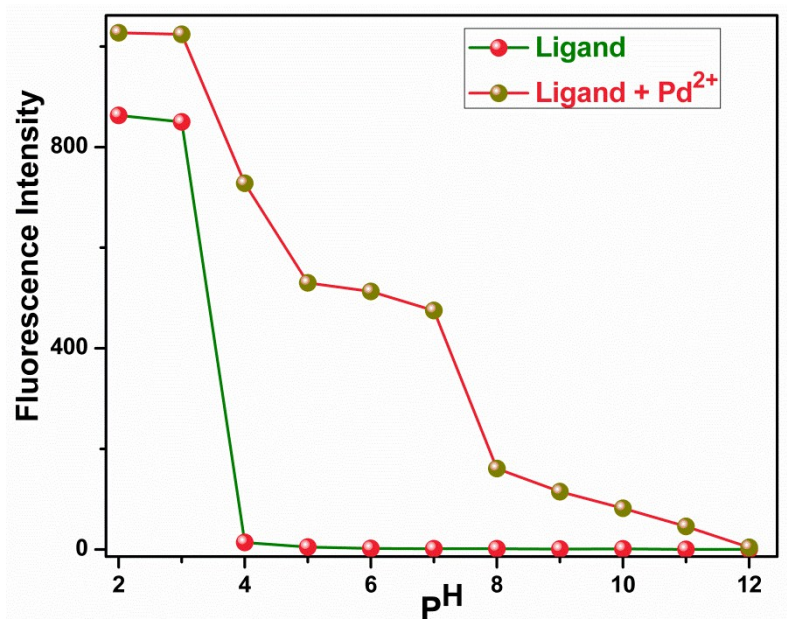
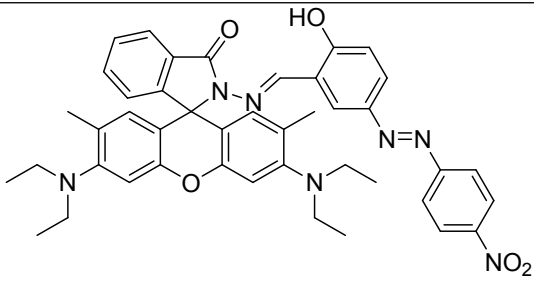
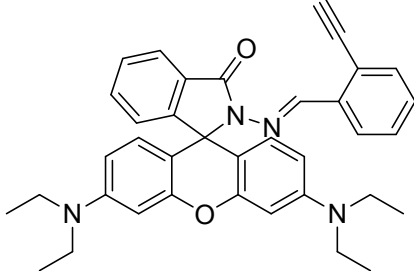
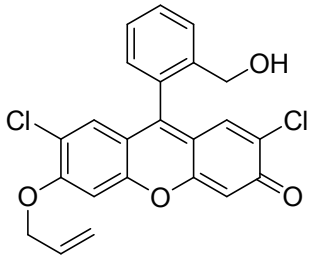
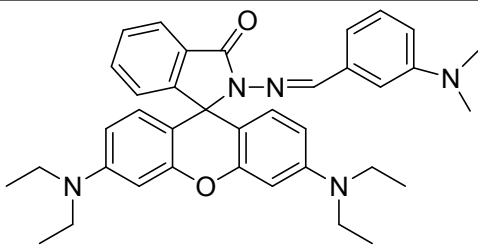
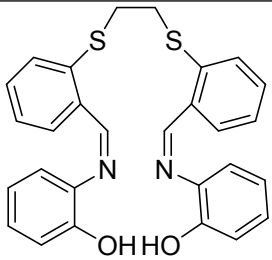
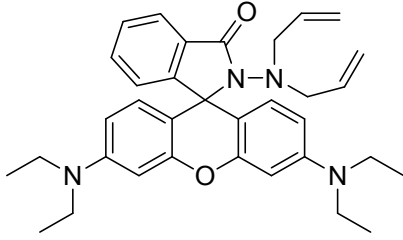
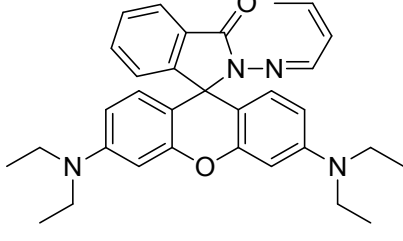
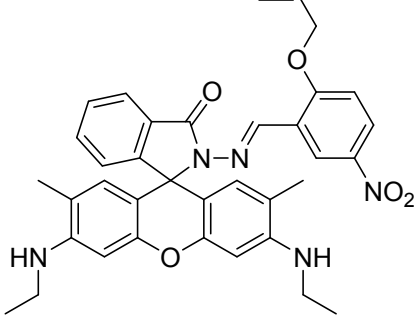


Fig. S13. Effect of pH on the fluorescence activity of RD-2 and RD-2 with Pd²⁺ in (CH₃CN/H₂O, 1/4, v/v, HEPES buffer).

Table S1. Comparative information of different probes for detection of Pd²⁺ and their LOD and Reference

Sl.No.	Fluorophore	LOD, nM	Ref.
1.		100	33
2.		55	34
3.		1650	35

4.		450	36
5.		191	37
6.		3900	38
7.		280	39
8.		120	40

9.		185	41
10.		190	42
11.		95	This Work