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Supporting Information for

The Diversity of The Coordination Bond Generated A POSS-based Fluorescent Probe for The Reversibly Detection of Cu(II), Fe(III) and Amino Acids

Kun Zhang, Xiaoni Wang, Minggang Tian, Zhiming Gou, Yujing Zuo*

School of Chemistry and Chemical Engineering, School of Materials Science and Engineering, University of Jinan, Shandong 250022, P.R. China.

*Corresponding Author. *E-mail address:* <u>zuoyujing0888@163.com</u>

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Sensor	Detection limit	Fluorescence	Cell imaging	Imaging
		photostability	photostability	application
RhQA	Cu ²⁺ (1.8 × 10 ⁻⁸ mol/L) Fe ³⁺ (3.3 × 10 ⁻⁸ mol/L)	Not mentioned	Not mentioned	Zebrafish
Ру	Fe ³⁺ (1.18 × 10 ⁻⁸ mol/L)	Not mentioned	Not mentioned	Cell
L	Zn ²⁺ (1.34 × 10 ⁻⁷ mol/L) Fe ³⁺ (1.39 × 10 ⁻⁷ mol/L)	Not mentioned	Not mentioned	Test strips
1	Fe ³⁺ (4.2 × 10 ⁻⁸ mol/L)	Not mentioned	Not mentioned	Cell Zebrafish
This work	Fe ³⁺ (3.2×10^{-9} mol/L) Cu ²⁺ (1.9×10^{-9} mol/L)	2 h	14 min	Test strips Cell Zebrafish

Table. S1 The comparison of PSI-A with some other sensor probes for ions.



Fig. S1. ¹H NMR spectrums of PSI-A in CDCl₃.



Fig. S2. ¹³C NMR spectrum of PSI-A in CDCl₃.



Fig. S3. The UV fluorescence spectrum of PSI-A.



Fig. S4. PSI-A was used to prepare a strip for detecting Cu^{2+} and Fe^{3+}



Fig. S5. Add Asp, Trp, Ser, Ace, Arg, Cys amino acid solution to the fluorescent probe solution of **PSI-A** quenched by Fe³⁺.



Fig. S6. (a)-(f) Add Asp, Ace, Arg, Trp, Ser, Cys amino acid solution to the fluorescent probe solution of **PSI-A** quenched by Cu^{2+} .



Fig. S7. PSI-A was used to dynamically detect cycles of Cu²⁺, Ace, Arg, Asp, Ser, Trp, and Fe³⁺, Asp.



Fig. S8. The fluorescence spectra of the probe PSI-A at different pH values.



Fig. S9. The fluorescence spectra of the probe PSI-A at ATP and Glu.



Fig. S10. PSI-A fluorescence in culture media.



Fig. S11. Cytotoxicity of PSI-A on HepG2 cells determined by MTT.