Supplementary Msterials

Cd(II) Coordination Polymers Constructed from Tris-pyridyl-trisamide and Polycarboxylic acid: Synthesis, Structures and Sensing Properties

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Fig. S1. IR spectrum of complex 1.



Fig. S2. IR spectrum of complex 2.



Fig. S3. IR spectrum of complex 3.



Fig. S4. The PXRD patterns of complex 1.



Fig. S5. The PXRD patterns of complex 2.



Fig. S6. The PXRD patterns of complex 3.



Fig. S7. TGA curve of complex 1.



Fig. S8. TGA curve of complex 2.



Fig. S9. TGA curve of complex 3.



Fig. S10. The PXRD patterns of complex **1** before and after immersion in different solvents.



Fig. S11. The PXRD patterns of complex 2 before and after immersion in different solvents.



Fig. S12. The photoluminescence spectra of L.



Fig. S13. The photoluminescence spectra of 1,4-H₂BDC.



Fig. S14. The photoluminescence spectra of 4,4'-H₂BDC.



Fig. S15. The photoluminescence spectra of complex 1.



Fig. S16. The photoluminescence spectra of complex 2.



Fig. S17. The solid-state luminescence spectra of complex 1 before and after immersion into different metal cations.



Fig. S18. The solid-state luminescence spectra of complex 2 before and after immersion into different metal cations.



Fig. S19. The emission spectra of complex 1 in different concentrations of Fe^{3+} ion.



Fig. S20. The emission spectra of complex 2 in different concentrations of Fe^{3+} ion.



Fig. S21. PXRD patterns for complex 1 after Fe^{3+} sensing for five cycles.



Fig. S22. PXRD patterns for complex 2 after Fe^{3+} sensing for five cycles.



Fig. S23. The PXRD patterns of complex **1** before and after immersion in different metal cations.



Fig. S24. The PXRD patterns of complex **2** before and after immersion in different metal cations.



Fig. S25. UV-vis absorption spectra of Fe^{3+} and the excitation and emission spectra of (a) 1 and (b) 2.



Fig. S26. The EDX data of complex 1. (a) The selected part for point scan.
(b) The result of point scan. (c) The element mapping scan of Fe³⁺. (d) The element percentage analysis of point scan.





(a)

(b)



(c)

Element	Weight (%)	Percentage (%)
C K	49.47	75.97
O K	19.99	23.05
Fe L	2.67	0.88
Cd L	0.62	0.10

(d)

Fig. S27. The EDX data of complex 2. (a) The selected part for point scan. (b) The result of point scan. (c) The element mapping scan of Fe^{3+} . (d) The element percentage analysis of point scan.







(c)

(b)

Element	Weight (%)	Percentage (%)
C K	38.66	67.49
O K	18.02	23.63
Fe L	22.40	8.41
Cd L	2.60	0.49

(d)

Fig. S28. IR spectra of L, 1,4-H₂BDC and complex 1 before and after immersion in Fe³⁺.



Fig. S29. IR spectra of L, 4,4'-H₂BDC and complex 2 before and after immersion in Fe³⁺.



Fig. S30. Selected XPS patterns before and after addition of Fe^{3+} ion to complex 2.



Fig. S31. Competitive sensing experiments in the presence of the other cations for complexes (a) **1** and (b) **2**.





(b)

Fig. S32. (a) N_2 adsorption-desorption isotherms for complex 1. (b) Poresize distribution curve for complex 1.



Fig. S33. (a) N_2 adsorption-desorption isotherms for complex 2. (b) Poresize distribution curve for complex 2.



Fig. S34. PXRD patterns for complex 1 before and after N_2 adsorption and desorption.



Fig. S35. PXRD patterns for complex 2 before and after N_2 adsorption and desorption.

