Supporting Information

A facile, low-cost bimetallic iron-nickel MOF nanozyme-propelled ratiometric fluorescent sensor for highly sensitive and selective uric acid detection and its smartphone application

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Fig. S1. (A) XRD pattern of Fe-MOF-NH₂. Fe-MOF-NH₂ exhibited four major peaks at 8°, 10°, 16° and 19° which correspond to (002), (101), (103) and (201) peaks of Fe-MOF-NH₂ phases, respectively. (B) SEM image of Fe-MOF-NH₂ at the scale of 500 nm.



Fig. S2. (A) XRD pattern of Ni-MOF-NH₂. Ni-MOF-NH₂ showed four major peaks at 10°, 16°, 18° and 22° which correspond to (100), (110), (101) and (210) peaks of Ni-MOF-NH₂ phases, respectively. (B) SEM image of Ni-MOF-NH₂ at 200 nm.



Fig. S3. SEM image of Fe₃Ni-MOF-NH₂ at 500 nm scale.



Fig. S4. (A) SEM elemental mapping image of oxygen element. (B) Map Sum Spectrum of Fe₃Ni-MOF-NH₂.



Fig. S5. (A) High-resolution XPS spectra of C 1s, N 1s (B) and O 1s (C).



Fig. S6. Comparison of peroxidase (POD)-like catalytic activity of $Fe_3Ni-MOF-NH_2$, $Fe-MOF-NH_2$ and $Ni-MOF-NH_2$, and the corresponding UV-visible spectra (7 μ L $Fe_xNi_y-MOF-NH_2$, 0.2 mM TMB and 10 mM H_2O_2 were used).



Fig. S7. EPR spectra of the DMPO/•OH spin adduct of Fe₃Ni-MOF-NH₂, Fe-MOF-NH₂ and Ni-MOF-NH₂.



Fig. S8. (A) Lineweaver-Burk double reciprocal plots of Fe₃Ni-MOF-NH₂ for TMB and (B) H₂O₂.



Fig. S9. (A) Optimization of catalytic time. (B) Optimization of uricase's concentration.



Fig. S10. (A) 3D normalized fluorescent column bars of Fe₃Ni-MOF-NH₂ with different concentrations of UA in a 2-month storage period at 4°C (bright-blue columns: day 0, pale-blue: after 2 months). (B) 3D normalized fluorescent column bars of DAP with different concentrations of UA in a 2-month storage period at 4°C (bright-orange columns: day 0, pale-orange: after 2 months).



Fig. S11. (A) Fluorescence spectra of Fe₃Ni-MOF-NH₂ with various concentrations of UA on day 0. (B) Fluorescence spectra of Fe₃Ni-MOF-NH₂ after 2 months with same concentrations of UA as on day 0. (C) Fluorescence spectra of DAP with various concentrations of UA on day 0. (D) Fluorescence spectra of DAP after 2 months with various concentrations of UA.



Fig. S12. 3D normalized ratiometric fluorescent column bars of the platform in the absence and presence of PO (orange column: 0.5 mg/mL uricase, blue column: 0.5 mg/mL uricase + 200 μ M UA, green columns: 0.5 mg/mL uricase + 200 μ M UA + PO).



Fig. S13. (A) Equivalent logic symbol of YES^NOT logic pair. (B) Corresponding truth table of the YES^NOT logic pair. (C) 3D normalized fluorescent column bars of Fe_3Ni -MOF-NH₂ (blue columns) and that of DAP (orange columns) in the absence and presence of UA. (D) Fluorescence spectra of Fe_3Ni -MOF-NH₂ and DAP under different input combinations of YES^NOT logic pair.

Catalyst	Substrate	$K_{m}(mM)$	$V_{max} (10^{-7} \text{ M} \cdot \text{s}^{-1})$	Ref.
Fe ₃ Ni-MOF-NH ₂	TMB	0.1458	1.3727	This work
	H_2O_2	0.1532	1.6838	This work
HRP	TMB	0.434	1.00	(43)
	H_2O_2	3.70	0.871	(43)

Table S1. Comparison of kinetic parameters (K_m and V_{max}) of Fe₃Ni-MOF-NH₂ and HRP.