A fluorometric and colorimetric dual-mode sensor based on nitrogen and iron co-doped graphene quantum dots for detection of ferric ion in biological fluids and cellular imaging

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**Fig. S1** (A) UV-vis absorption spectra of N, Fe-GQDs, inset is the photographs of N, Fe-GQDs under the 365 nm UV light (left) and sunlight (right). (B) Fluorescence emission spectra of N, Fe-GQDs with excitation by different wavelengths. (C) The high-resolution TEM (HRTEM) image of N, Fe-GQDs. (D) The size distribution of N, Fe-GQDs.



**Fig. S2** (A) XRD patterns of N, Fe-GQDs. (B) The high-resolution XPS spectrum of C 1s in N, Fe-GQDs. (C) The high-resolution XPS of N 1s in N, Fe-GQDs. (D) The high-resolution XPS of Fe 2p in N, Fe-GQDs.



Fig. S3 Fluorescence decay spectra of N, Fe-GQDs in the absence (black) and presence (red) of  $Fe^{3+}$  ions.



Fig. S4 The photograph of corresponding color changes of N, Fe-GQDs in the presence of different  $Fe^{3+}$  concentrations.



Fig. S5 Absorption spectra response of N, Fe-GQDs at 335 nm in the presence of different biomolecules and Fe<sup>3+</sup> (300  $\mu$ M). A<sub>0</sub> and A are the absorbance at 335 nm in the absence and presence of biomolecules and Fe<sup>3+</sup>, respectively.



**Fig. S6** (A) UV-vis absorption spectra of N, Fe-GQDs in the presence of different  $Fe^{3+}$  concentrations in lake water,  $Fe^{3+}$  concentration is 170, 200, 250, 300, 350 and 400  $\mu$ M (from bottom to top), respectively. Inset is the relationship between the absorbance at 335 nm and the  $Fe^{3+}$  ions concentrations in lake water. (B) The photograph of N, Fe-GQDs in the presence of the corresponding  $Fe^{3+}$  concentrations in lake water under sunlight.

Products	Detect method	Linear range (µM)	Detect limit (µM)	Ref	
CDs	Fluorometry	12.5-100	9.97	1	
N-CQDs	Fluorometry	0-20	-	2	
CDs	Fluorometry	8-100	0.8	3	
CQDs	Fluorometry	0-50	1.3	4	
CDs	Fluorometry	16-166	6.05	5	
P <sub>2</sub> O <sub>7</sub> <sup>4-</sup> -AuNPs	Colorimetry	10-60	5.6	6	
AuNPs	Colorimetry	1–37	0.85	7	
AgNPs	Colorimetry	0.08-80	0.08	8	
N, Fe-GQDs	Fluorometry	10-110	3.21		
	Colorimetry	0-450	1.34	This work	

Table S1 Comparison of different fluorescence probes for  $Fe^{3+}$  detection.

CDs: Carbon dots

CQDs: Carbon quantum dots

N-CQDs: N-dope carbon quantum dots

AgNPs: Ag nanoparticles

AuNPs: Au nanoparticles

Sample	Spike (µM)	Found (µM)	Recovery (%)	RSD (%, n=3)
serum	20.00	20.84 (19.91ª)	104.20	2.17
urine	50.00	48.81 (51.01 <sup>a</sup> )	97.62	1.92
	100.00	93.60 (97.79 <sup>a</sup> )	93.60	3.31
	20.00	18.74 (20.89 <sup>a</sup> )	93.70	2.58
	50.00	51.17 (48.63 <sup>a</sup> )	102.34	4.90
	100.00	108.20 (102.31ª)	108.20	2.04

 Table S2 Fe<sup>3+</sup> determination results in human serum and urine samples.

<sup>a</sup> The concentration in bracket was found by ICP-OES.



**Fig. S7** The fluorescence microscopy images of HeLa cells treated with N, Fe-GQDs, (A) the bright-field images, (B) the fluorescent images.

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