

**A simple camphor based AIE fluorescent probe for highly specific and sensitive
detection of hydrazine and its application in living cells**

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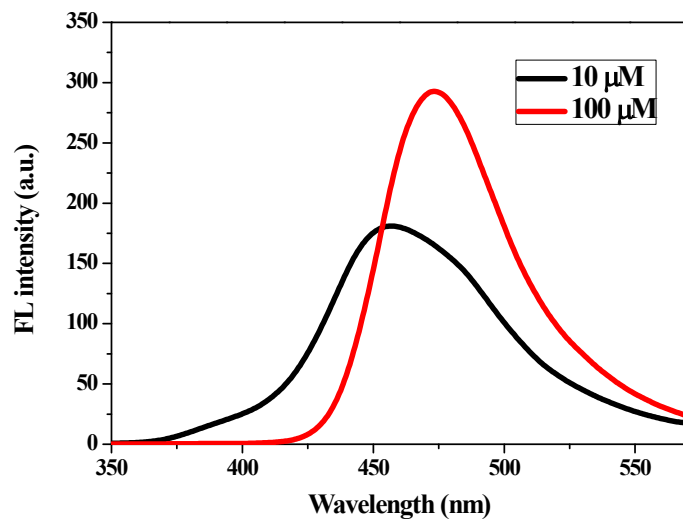


Fig. S1. Fluorescence spectra of probe **CDA** in different concentrations with $f_w = 90\%$.

$\lambda_{\text{ex}} = 300 \text{ nm}$.

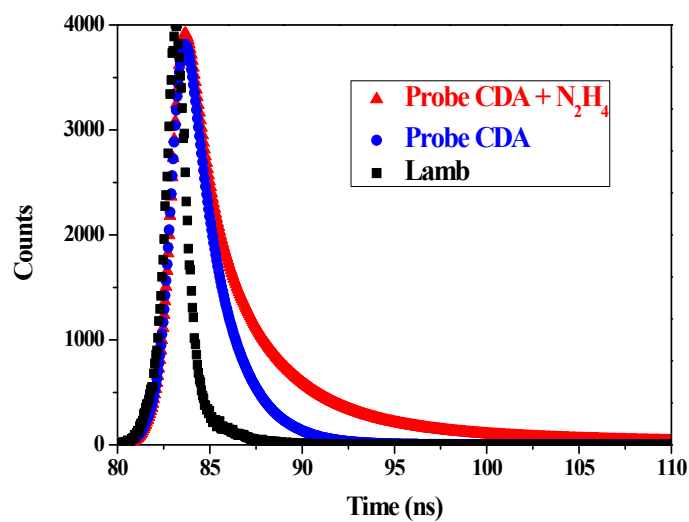


Fig. S2. Fluorescence decay curves of probe **CDA** in the absence and presence of N_2H_4 .

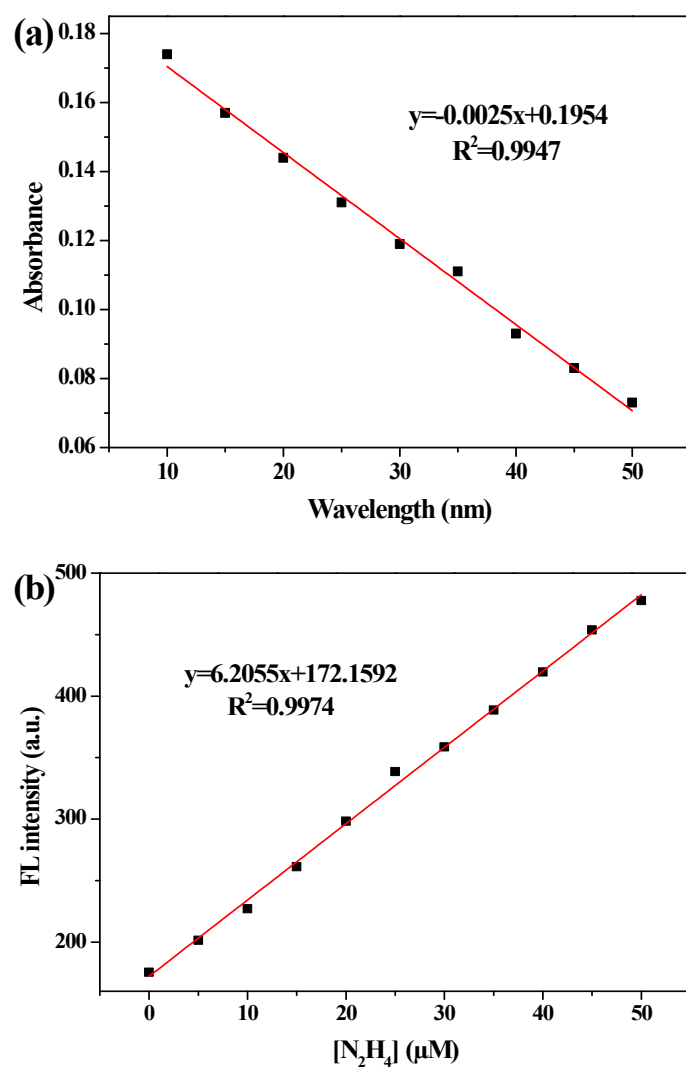


Fig. S3. The linear relationships of the absorption intensity (a) and emission intensity (b) of probe **CDA** against the concentration of N_2H_4 . $\lambda_{ex} = 300$ nm.

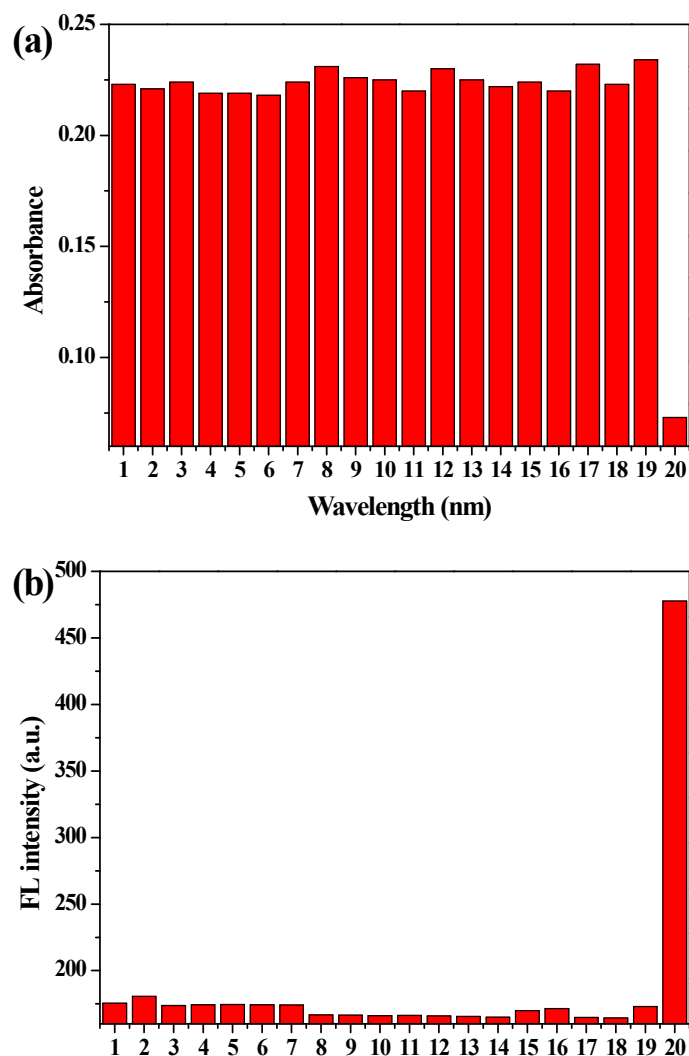


Fig. S4. The absorption intensity (a) and emission intensity (b) of probe **CDA** in the presence of N₂H₄ (40 μ M) and other analytes (100 μ M) in PBS buffer solution (pH = 7.4, 10 mM, containing 10% EtOH). (1) Blank, (2) Alanine, (3) Asparagine, (4) Lysine, (5) Glutathione, (6) Cysteine, (7) Tyrosine, (8) Valine, (9) Urea, (10) Acetylhydrazide, (11) Hydroxylamine, (12) Thiosemicarbazide, (13) o-Phenylenediamine, (14) Al³⁺, (15) Zn²⁺, (16) Fe³⁺, (17) Hg²⁺, (18) NO₃²⁻, (19) SO₃²⁻, (20) N₂H₄. $\lambda_{\text{ex}} = 300$ nm.

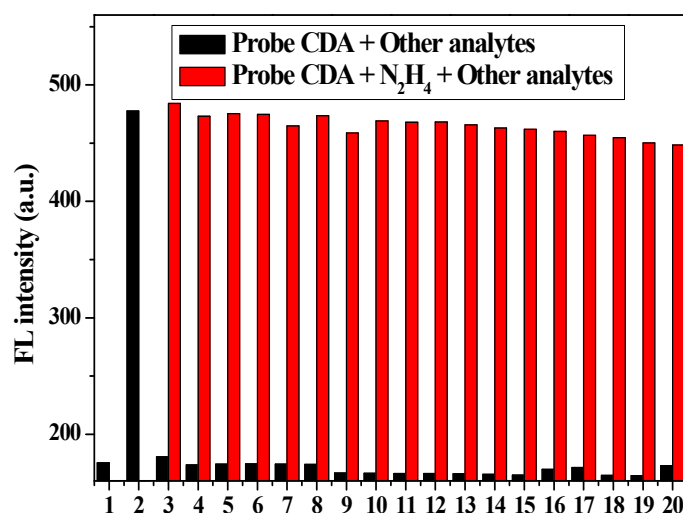


Fig. S5. The emission intensity of probe **CDA** toward N₂H₄ (40 μ M) in the absence and presence of other competitive analytes (100 μ M) in PBS buffer solution (pH = 7.4, 10 mM, containing 10% EtOH). (1) Blank, (2) N₂H₄, (3) Alanine, (4) Asparagine, (5) Lysine, (6) Glutathione, (7) Cysteine, (8) Tyrosine, (9) Valine, (10) Urea, (11) Acetylhydrazide, (12) Hydroxylamine, (13) Thiosemicarbazide, (14) o-Phenylenediamine, (15) Al³⁺, (16) Zn²⁺, (17) Fe³⁺, (18) Hg²⁺, (19) NO₃²⁻, (20) SO₃²⁻. $\lambda_{\text{ex}} = 300$ nm.

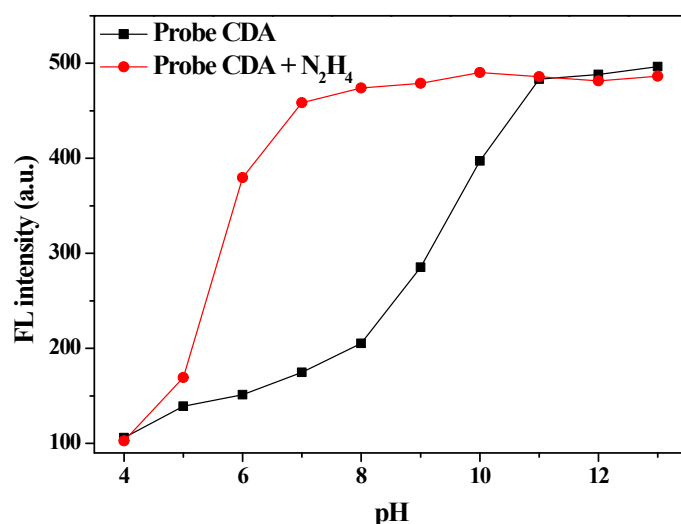


Fig. S6. Effect of pH on the fluorescence intensity of probe **CDA** in the absence and presence of N₂H₄ (40 μ M). $\lambda_{\text{ex}} = 300$ nm.

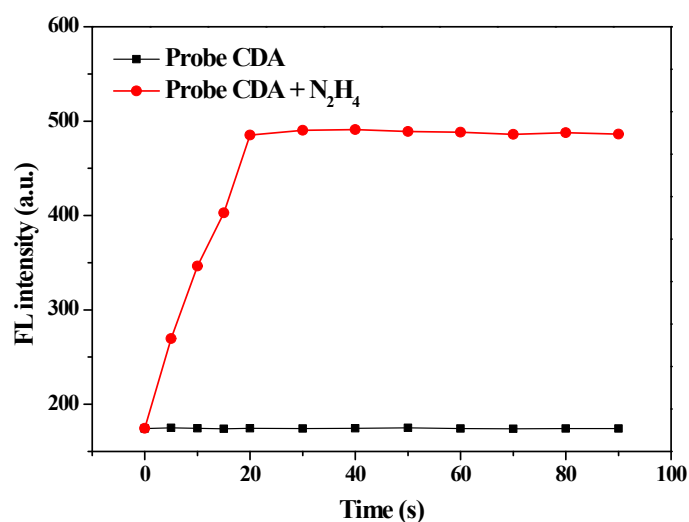
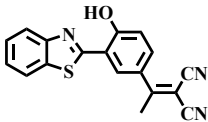
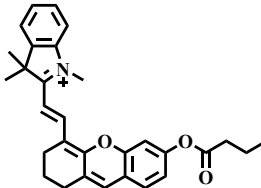
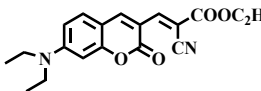
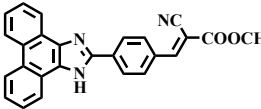
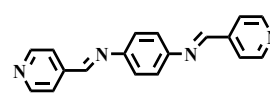
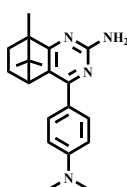


Fig. S7. Time-dependent fluorescence intensity changes of probe **CDA** upon addition of N₂H₄ (40 μ M) in PBS buffer solution (pH = 7.4, 10 mM, containing 10% EtOH). λ_{ex} = 300 nm.

Table S1. Comparison with other reported N₂H₄ fluorescence probes.

Probe	Response time	LOD	Detection medium	Reference
	55 min	9.3 ppb	DMF/H ₂ O (v/v 8/2)	[62]
	30 min	5.09 ppb	DMF/PBS (v/v 1/19)	[63]
	12 min	5.2 ppb	DMF/PBS (v/v 3/7)	[64]
	20 min	51.2 ppb	DMSO/H ₂ O (v/v 6/4)	[65]

	2 min	3.2 ppb	MeOH/H ₂ O (v/v 4/6)	[65]
	20 s	2.26 ppb	EtOH/PBS (v/v 1/9)	This work

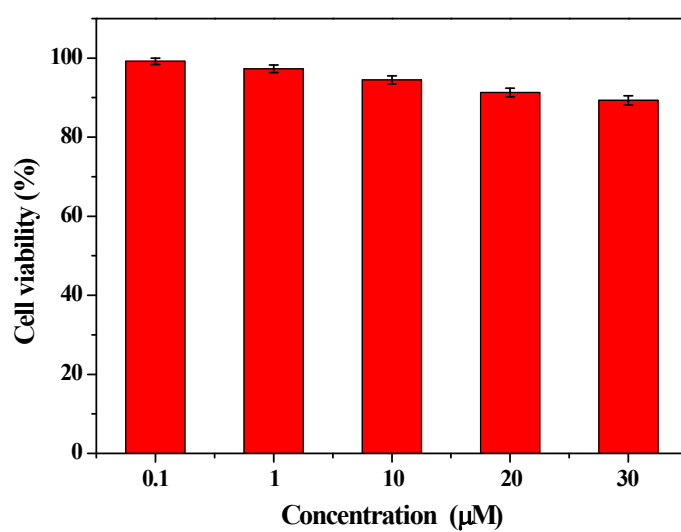


Fig. S8. MTT assay of HeLa cells was incubated with 0.1, 1, 10, 20 and 30 μM probe **CDA** for 48 h.

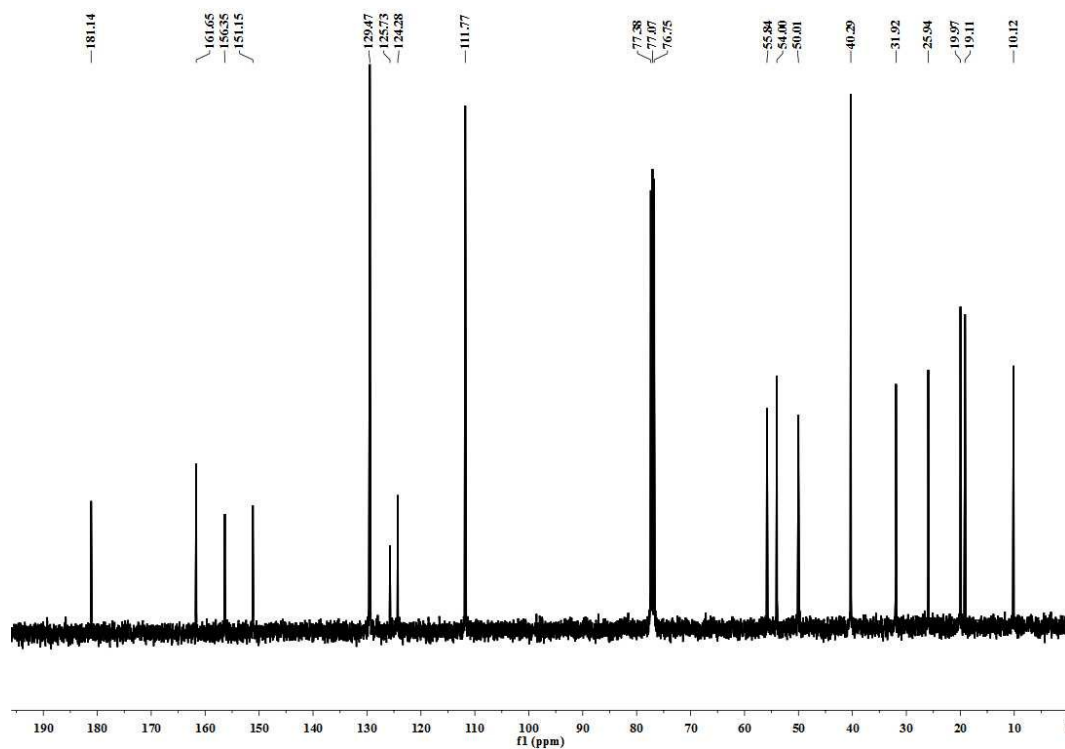


Fig. S9. ^{13}C NMR spectra of probe **CDA** in CDCl_3 .

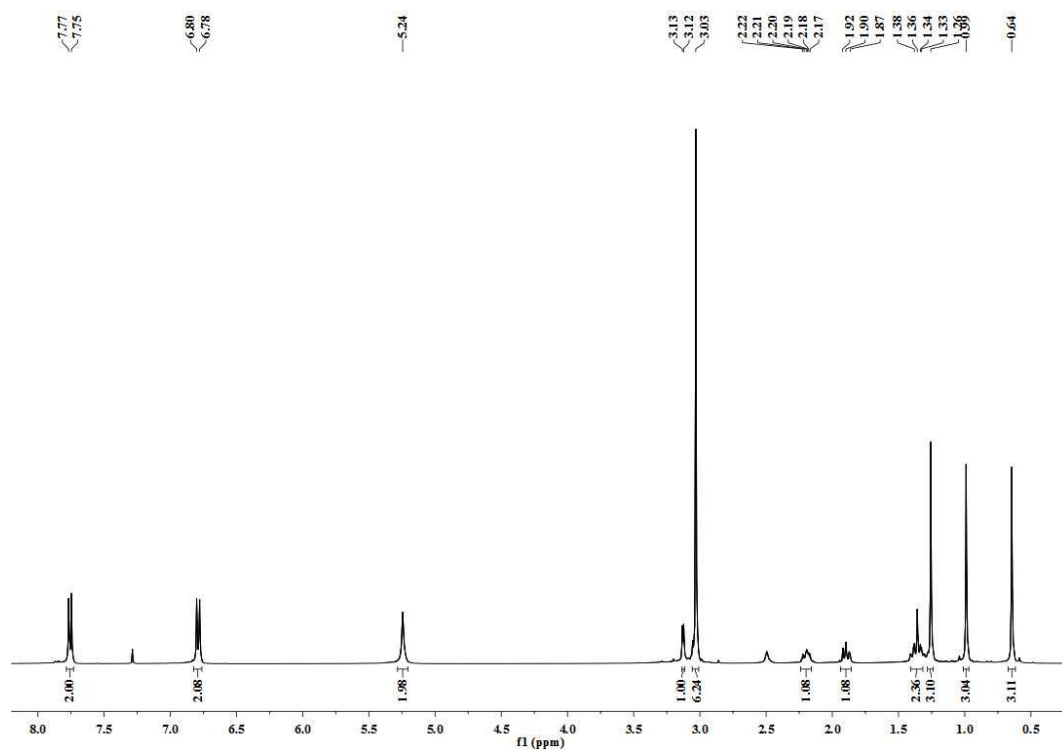


Fig. S10. ^1H NMR spectra of probe **CDA** in CDCl_3 .

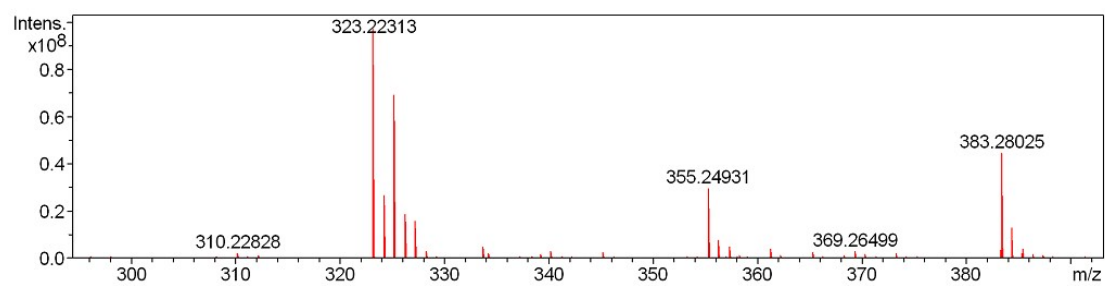


Fig. S11. HRMS of probe **CDA**.