

A simple sensor based on 1,8-naphthalimide with large Stokes shift for detection of hypochlorous acid in living cells

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 2. ^{13}C NMR spectrum of compound 2
 3. ^1H NMR spectrum of Probe 1
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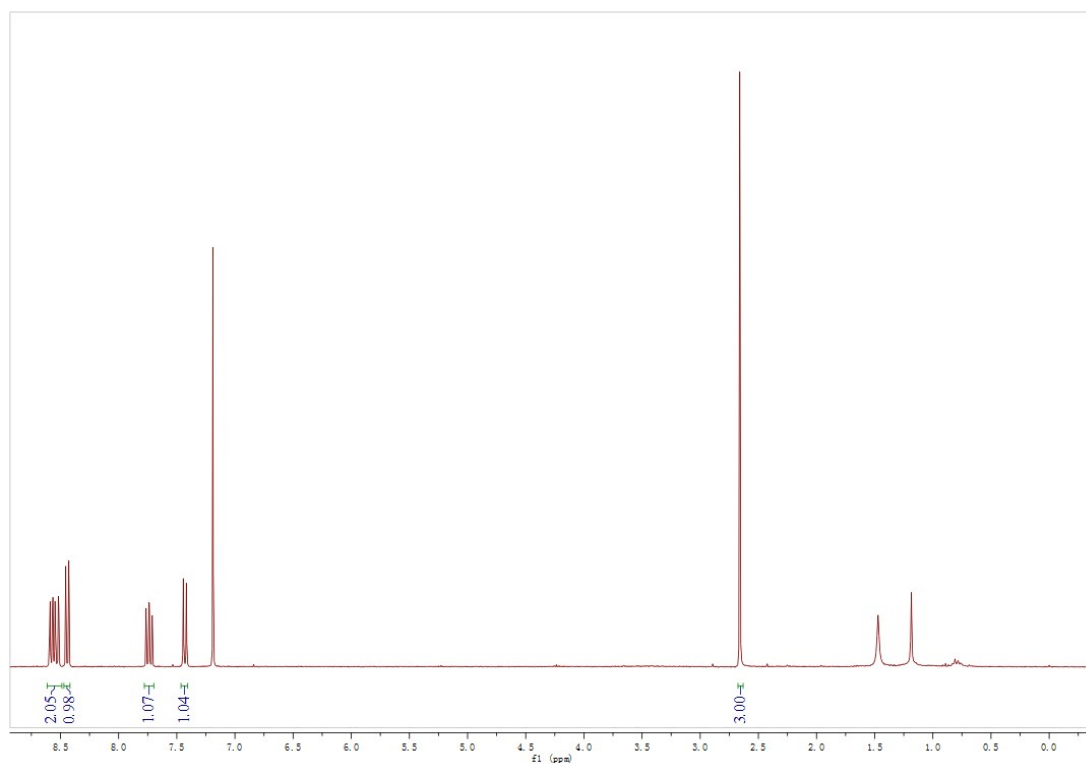


Fig. S1 ^1H NMR spectrum of compound 2 (CDCl_3)

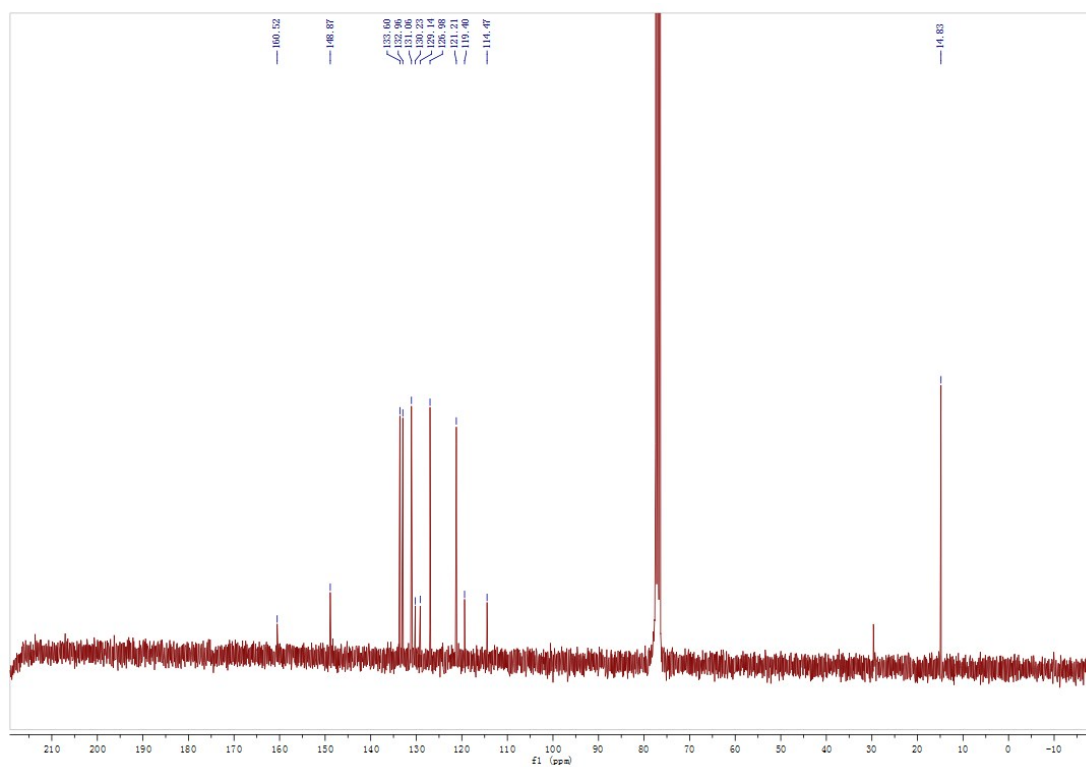


Fig. S2 ^{13}C NMR spectrum of compound 2 (CDCl_3)

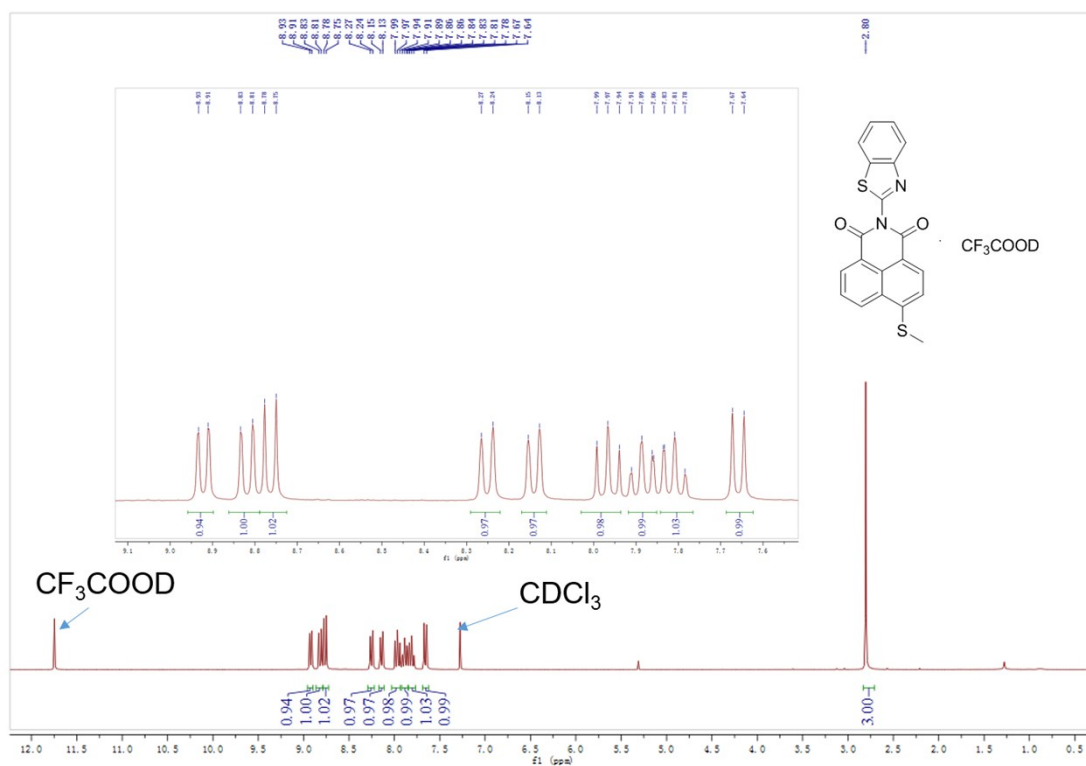


Fig. S3 ¹H NMR spectrum of Probe 1 (CDCl₃ + CF₃COOD)

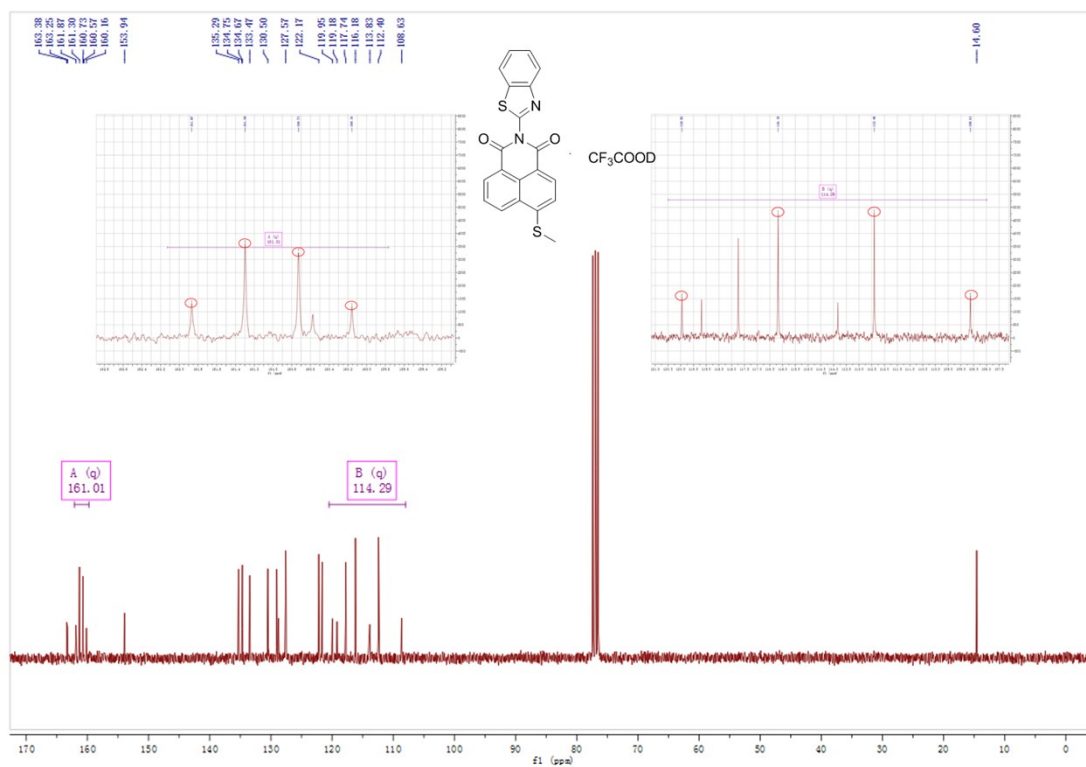


Fig. S4 ¹³C NMR spectrum of Probe 1 (CDCl₃ + CF₃COOD)

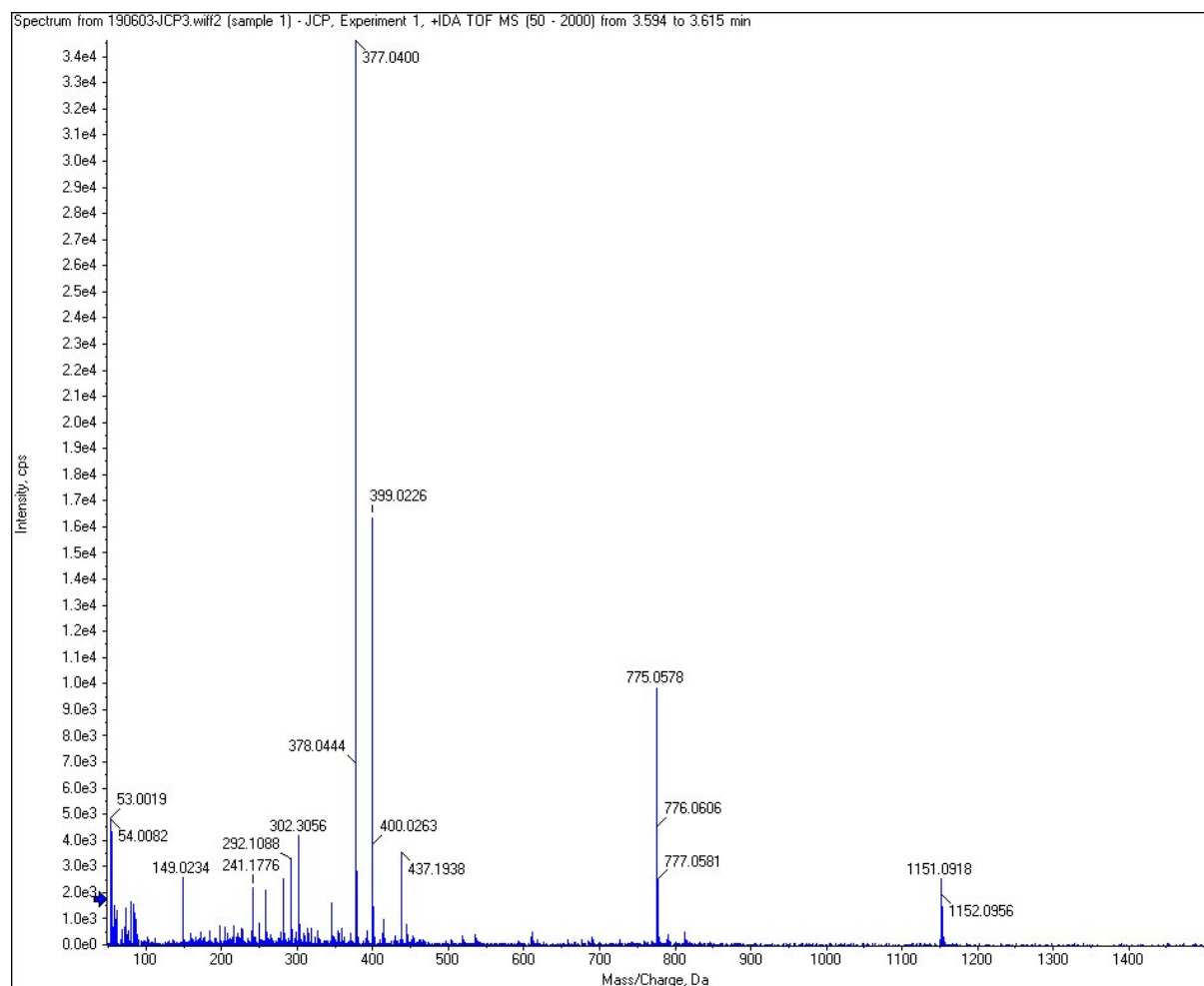


Fig. S5 HRMS spectra characterizations of Probe 1

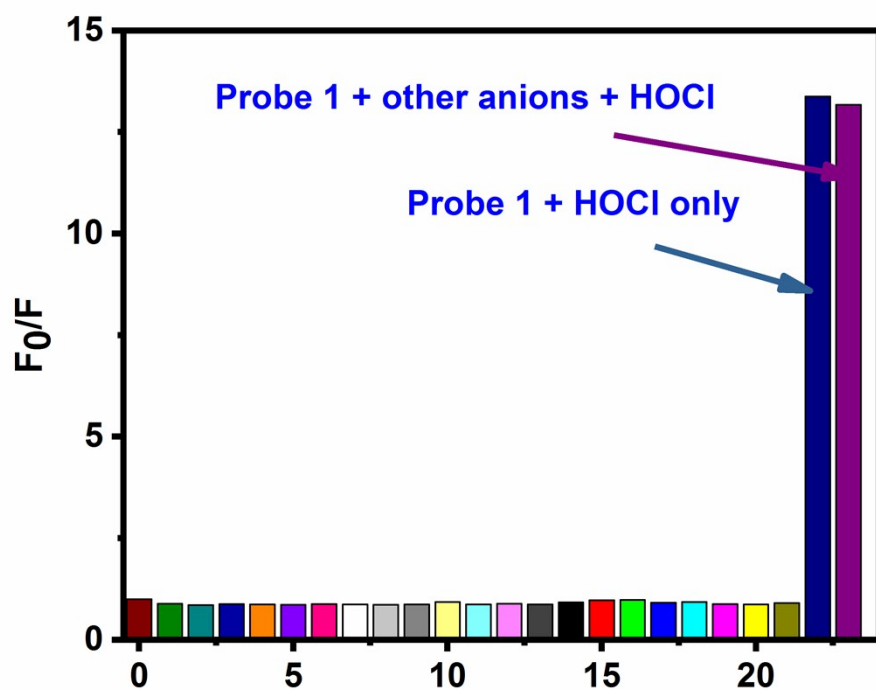


Fig. S6 Fluorescence spectra of Probe 1 to various relevant species (all 10eq; NO_3^- , NO_2^- , Ag^+ , Al^{3+} , Ca^{2+} , Cr^{3+} , Co^{2+} , Fe^{2+} , Fe^{3+} , Mn^{2+} , Ni^{2+} , Pb^{2+} , Zn^{2+} , Cu^{2+} , Hg^{2+} , Cd^{2+} , H_2O_2 , HOCl, NO, ONOO $^-$, $\bullet\text{OH}$, T-Buoo $^-$).

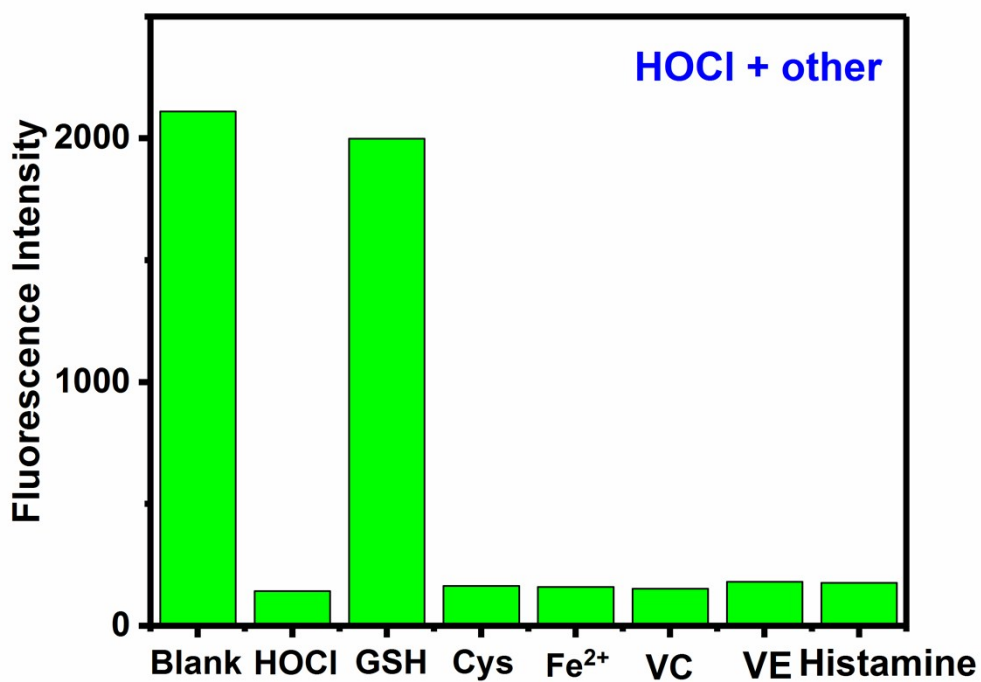


Fig. S7 Comparison of the fluorescence intensity with various reducing materials (10 eq) to **Probe 1** (10^{-5}M) + HOCl.

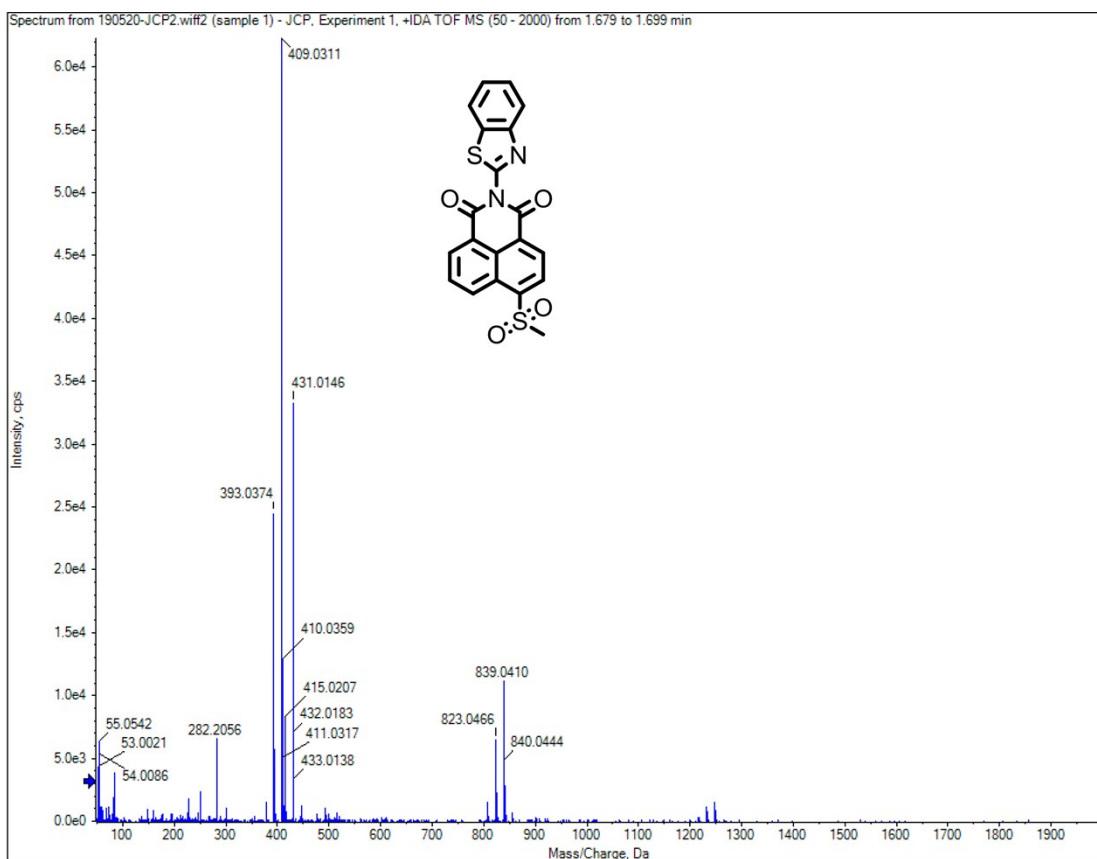
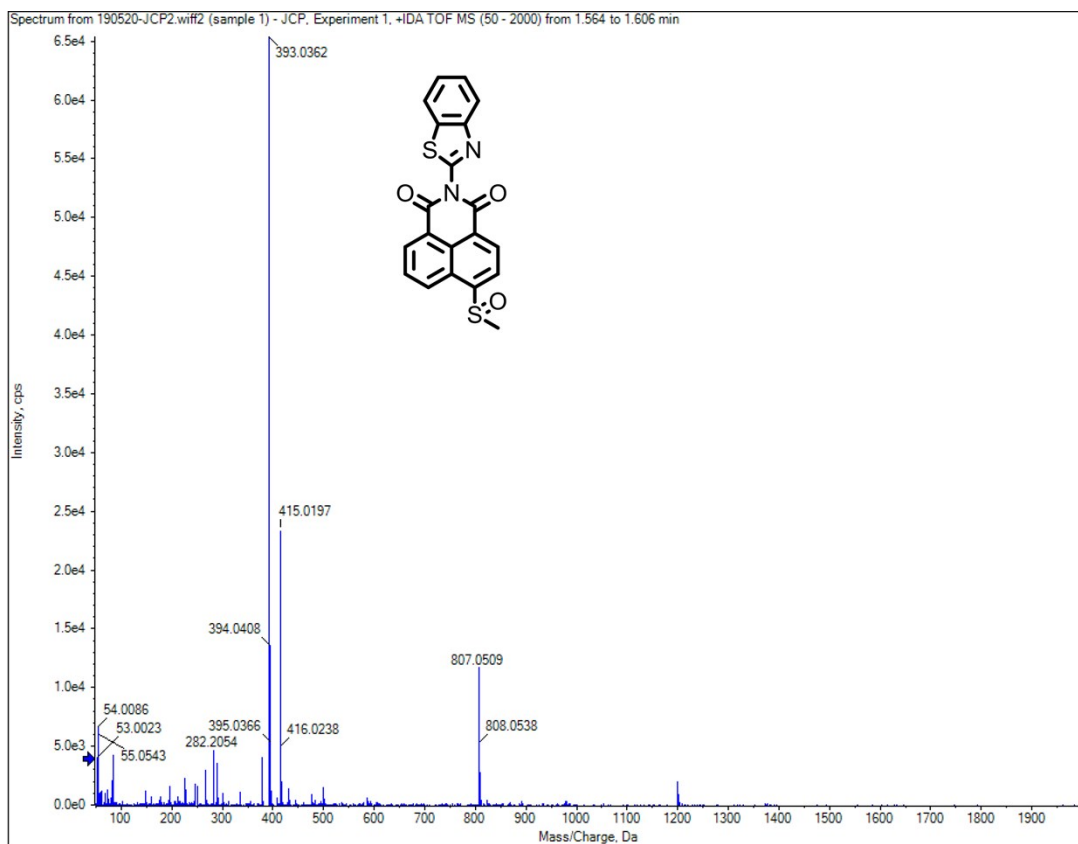


Fig. S8 Mass spectrum of Probe 1 after oxidation by HOCl

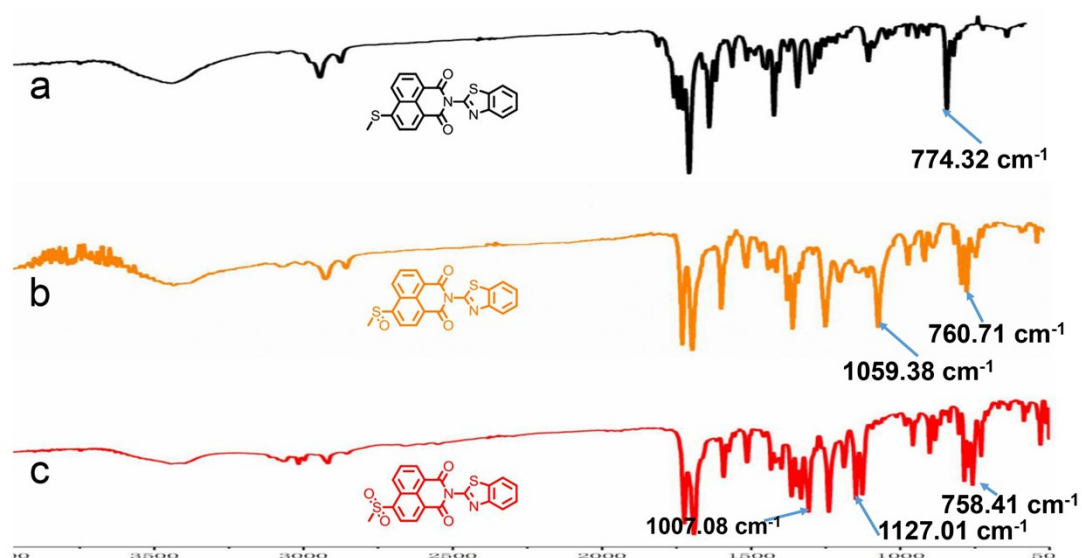


Fig. S9 FT-IR Spectra of Probe 1 and product after reaction with HOCl

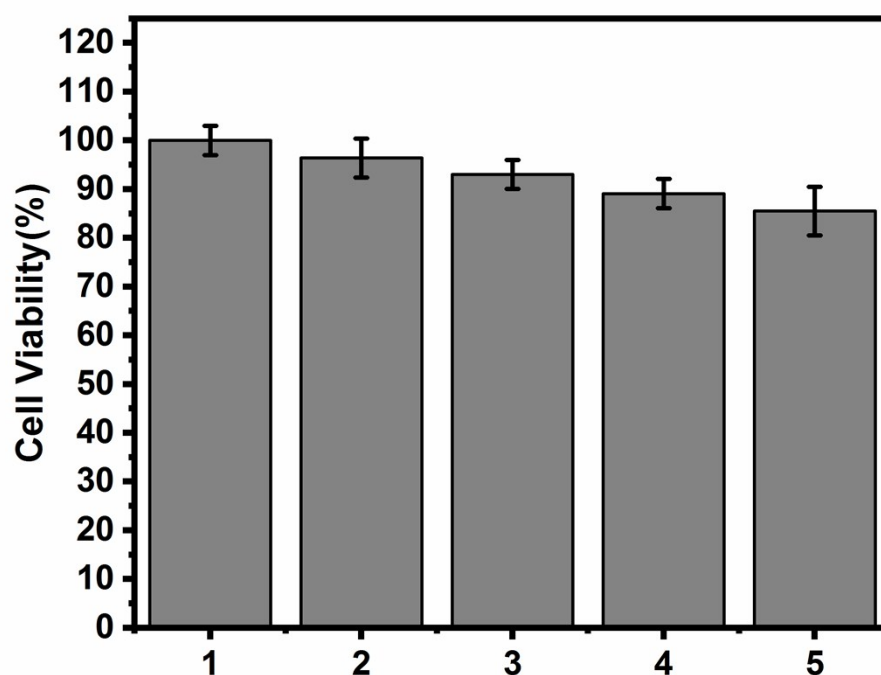
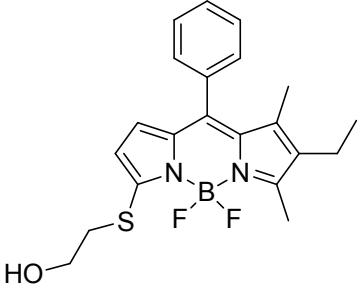
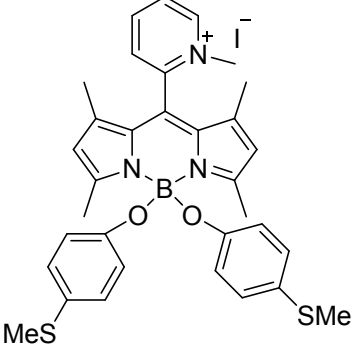
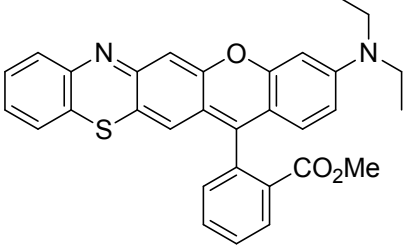
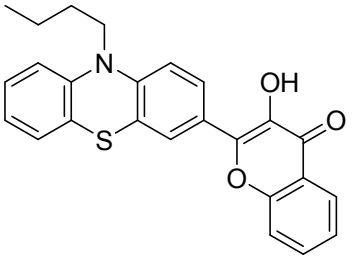
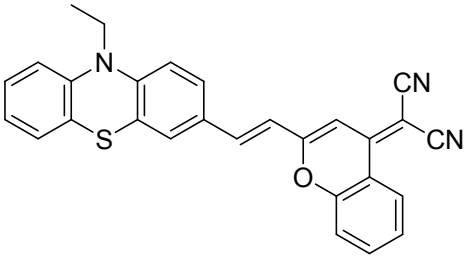
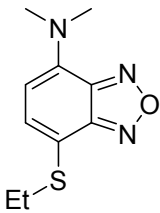
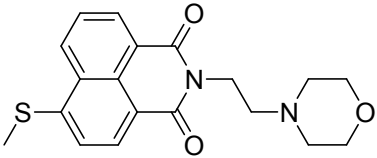
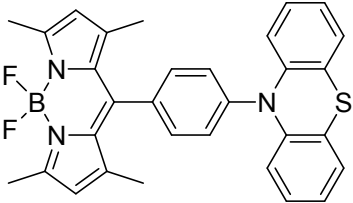
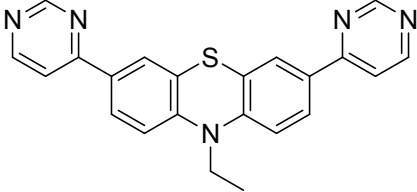
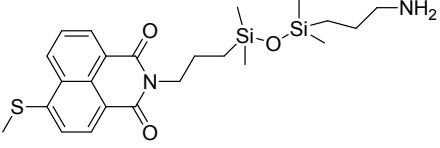
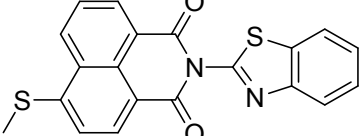


Fig. S10 MTT assay of HeLa cells treated in the presence of Probe 1 (1. 0μM 2. 5μM 3. 10μM 4. 20μM 5. 30μM) and incubated for 24 hours.

Table S1 Comparison of highly specific fluorescent probes for HOCl

Probe	Detection limit	Reaction time	Stokes shift	Reference
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	0.4 μ M	fast	12nm	1
	53nM	10min	40nm	2
	2.3nM	10min	97 nm	3
	6.6nM	10min	140nm	4
	0.72 μ M	null	150nm	5
	0.34 μ M	60min	170nm	6

	0.674 μM	2.5min	105nm	7
	0.72 μM	15min	14nm	8
	0.453 μM	20min	68nm	9
	1.6 μM	30s	95nm	10
	0.237 μM	2min	120nm	This work

Reference

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