Supplementary Information

A fast-response and highly specific Si-Rhodamine probe for endogenous peroxynitrite detection in living cells[†]

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1 Supplementary Spectra and chart



Fig. S1. Proposed reaction mechanism and HRMS of compound SiMH.



Fig. S2. Fluorescence intensity changes of SiNH (5 μ M) for ONOO⁻ over various species (100 equiv) in HEPES buffer solution (50 mM, pH=7.4 H₂O:CH₃CN = 7:3) at 595 nm (1) Probe only (5 μ M), (2) Cys, (3) NO, (4) HClO, (5) H₂O₂, (6) NaNO₂, (7) GSH, (8) Glu, (9) Pro, (10) ONOO⁻. $\lambda_{ex} = 450$ nm.



Fig. S3. Average size of nanoprobe measured by dynamic light scattering.



Fig. S4. Fluorescence spectra of nanoprobe (5 μ M) ($\lambda_{ex} = 450$ nm) in the presence of different amount of ONOO⁻ (0 – 10 μ M) in aqueous solution, and the corresponding linear relationship between the fluorescent intensity and ONOO⁻ concentrations. Conditions: HEPES buffers (50 mM, pH = 7.4).



Fig. S5. Fluorescence intensity of micelles in the absence and presence of ONOO⁻ at various pH values ($\lambda_{em} = 595$ nm).



Fig. S6. Percentage of viable Hela cells after treatment with indicated concentrations of micelles after 8 hours.

2 ¹H NMR, ¹³C NMR and HRMS charts.



Figure S7. ¹H NMR of compound SiX (CDCl₃, 400 MHz).



Figure S8. ¹³C NMR of compound SiX (CDCl₃, 100 MHz).



Figure S9. HRMS of compound SiX.



Figure S10. ¹H NMR of compound SiCl (CDCl₃, 400 MHz).



Figure S11. HRMS of compound SiCl.



Figure S12. ¹H NMR of compound SiNH (CDCl₃, 400 MHz).



Figure S13. ¹³C NMR of compound SiNH (CDCl₃, 100 MHz).



Figure S14. HRMS of compound SiNH.



Figure S15. ¹H NMR of compound SiMH (CDCl₃, 400 MHz).



Figure S16. HRMS of compound SiMH.