Electronic Supplementary Information

A redox reversible endoplasmic reticulum targeted fluorescent probe

for revealing the redox status of living cells

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1. The preparation of ROS

2,2'-azobis(2-2-ylpropane) dihydrochloride was dissolve in deionized water to prepare ROO•, sodium nitrite was dissolved in deionized water to prepare NO_2^- , NOC-13 was dissolved in deionized water to obtain NO and potassium peroxide was dissolved in DMSO to obtain $O_2^{\bullet-}$. Hydroxyl radicals (•OH) were produced by ferrous ammonium sulfate and hydrogen peroxide (Fenton reaction). Peroxynitrite (ONOO⁻) was synthesized from sodium nitrite and H₂O₂. ¹O₂ was generated by the reaction of H₂O₂ and Na₂MoO₄. ClO⁻ and H₂O₂ were prepared by diluting the purchased stock solution.

2. MTT assays

Cytotoxicity was attained by standard MTT assays. MCF-7 cells were incubated in 96-well culture plates in the presence of 200 μ L DMEM supplemented with 10% FBS and 1% penicillin and streptomycin at 37°C for 24 h in 5% CO₂. After discarding the culture medium, the cells were incubated in the solution of N-Se at different concentrations (0, 0.01, 0.05, 0.1, 0.5, 1, 5, 10, 20, and 40 μ M). Following incubation for 24 h, the cells were rinsed with PBS for three times, and then 180 μ L culture solution and 20 μ L MTT (5 mg/mL) were added to each well for 4 h. Subsequently, the MTT solution was supplanted by 0.2 mL DMSO in each well. The absorbance of each well was acquired at 570 nm. 8 replicate wells per concentration were set and the viability of cells without N-Se was set to 100%.

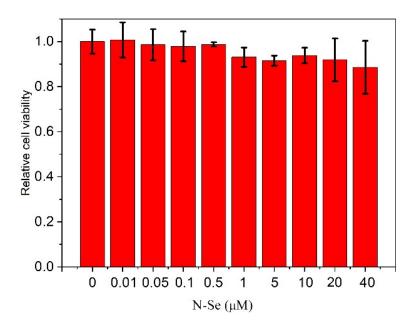


Fig. S1 Cytotoxicity assays of N-Se at different concentrations for MCF-7 cells.



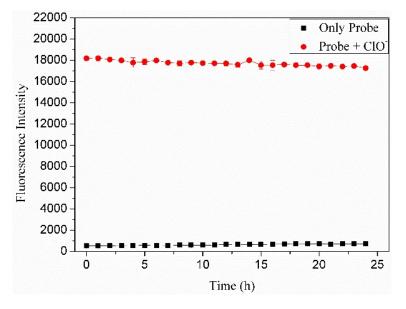


Fig. S2 Photostability of 10 μ M N-Se and N-SeO. Slit widths: 3 and 5 nm for excitation and emission spectra. $\lambda_{ex}/\lambda_{em} = 417/540$ nm.

4. NMR and HRMS data

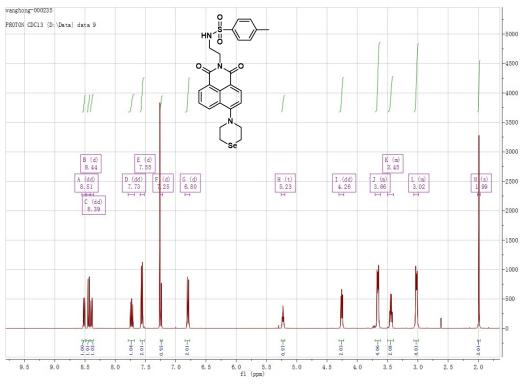
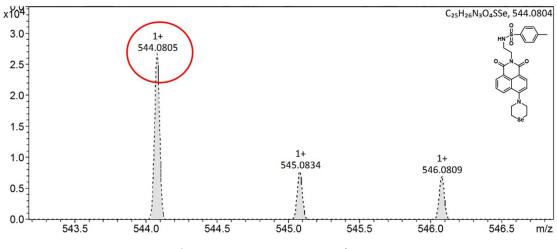
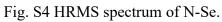


Fig. S3 1H NMR spectrum of N-Se.





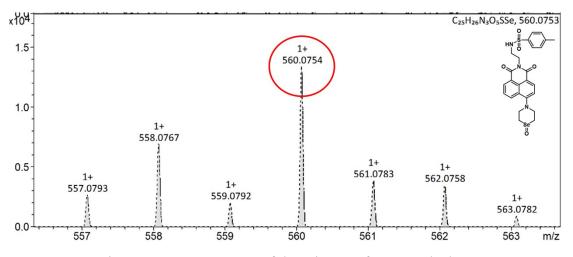


Fig. S5 HRMS spectrum of the mixture of N-Se and ClO⁻.