Supporting Information for

A Lysosome-targeted and Ratiometric Fluorescent Probe for Imaging Exogenous and Endogenous Hypochlorous Acid in Living Cells

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Figure S1 The absorption spectral changes of **Lyso-HA** (10 μ M) upon addition of increasing concentrations of HOC1 (0-10 equiv) in PBS buffer, pH 5.0, containing 30 % DMF as a cosolvent. Inset: Photographs showing the color changes of the probe Lyso-HA (1 mM) before and after addition of 10 equiv. HOCl to the solution.



Figure S2 The emission intensity changes (at 585 nm) of **SP** at different pH PBS buffer, containing 30 % DMF as a cosolvent ($\lambda_{ex} = 420$ nm).



Figure S3 The fluorescence responses of the probe Lyso-HA (5.0 μ M) to various relevant species (100 μ M) in pH 5.0, PBS buffer (30 % DMF)



Figure S4 Cytotoxicity assays of **Lyso-HA** at different concentrations (0 μ M; 5 μ M; 10 μ M; 20 μ M; 30 μ M; 50 μ M) for HeLa cells.



Figure S5 Quantified relative fluorescence intensity ratio of I_{red}/I_{blue} images analyzed using Nikon NIS Element software and presented as mean \pm s.d, n = 3.



Figure S6 Brightfield and fluorescence images of HeLa cells stained with the probe **Lyso-HA** and LysoTracker Greeen. a) brightfield image; b) from green channel (lysosomes staining); c) from the red channel; d) overlay of brighfield, green and red channels; e) overlay of green and red channels; f) Intensity profile of linear region of interest across in the HeLa cell costained with LysoTracker Green and Red channel of **Lyso-HA** imaging of HOCl; g) Intensity scatter plot of green and red channels.



Figure S7 ¹H-NMR (DMSO-*d*₆) spectrum of compound 3



Figure S8 ¹³C-NMR (DMSO-*d*₆) spectrum of compound 3



Figure S10 ¹³C-NMR (DMSO-*d*₆) spectrum of compound **4**.



Figure S12 ¹³C-NMR (DMSO-*d*₆) spectrum of compound Lyso-HA.