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The fluorescent biomarkers for lipid droplets with quinolone-coumarin unit

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Fig. S1 Photofading behaviors of dyes 1a-b in acetonitrile.



Fig. S2 Normalized absorption spectra (a) and emission spectra (b, excited at 430nm, slit widths: 1.5 nm/1.5 nm) of dye **1b** (10µM) in different solvents.



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Fig. S4 The reversible absorption (left) and emission (right) responses of dyes **1a–b** with changes between neutral and acidic conditions. (a) absorption spectra of **1a**; (b) emission spectra of **1a**; (c) absorption of **1b**; (d) emission spectra of **1b**.



Fig. S5 Fluorescence confocal images of living I929 (a-d) and HeLa (e-h) cells with dye **1b** (2 μ M) and their ROIs analysis. (a, e) Bright–field images; (b, f) confocal images (red channel) of cells with Nile red (1 μ M); (c, g) confocal images (green channel) of cells with **1b** (2 μ M); (d, h) merged images of green and red channels; (i, j) fluorescence intensities of the regions of interest (ROIs) across the cells. Green channel emission was collected in 470–550 nm upon excitation at 458 nm, and red channel emission was collected in 575–700 nm upon excitation at 561 nm. Scale bar: 25 μ m.



Fig. S6 Fluorescence confocal images of fixed I929 (a-d) and HeLa (e-h) cells with dye **1b** (2 μ M) and their ROIs analysis. (a, e) Bright–field images; (b, f) confocal images (red channel) of cells with Nile red (1 μ M); (c, g) confocal images (green channel) of cells with **1b** (2 μ M); (d, h) merged images of green and red channels; (i, j) fluorescence intensities of the regions of interest (ROIs) across the cells. Green channel emission was collected in 470–550 nm upon excitation at 458 nm, and red channel emission was collected in 575–700 nm upon excitation at 561 nm. Scale bar: 25 μ m.



Fig. S7 Fluorescence confocal images of living HeLa cells with dye **1a** (2 μ M), Nile red (1 μ M), MitoTracker [®] Red CMXRos (0.5 μ M) or Lyso-Tracker Red (0.5 μ M). (a, f, k) Bright–field images; (b, g, l) confocal images (green channels) of cells with dye **1a** (2 μ M); (c) confocal images (red channel) of cells with Nile red (1 μ M); (d) merged images of (b) and (c); (e) fluorescence intensity correlation plot of (d); (h) confocal images (red channel) of cells with MitoTracker [®] Red CMXRos (0.5 μ M); (i) merged images of (g) and (h); (j) fluorescence intensity correlation plot of (i); (m) confocal images (red channel) of cells with Lyso-Tracker Red (0.5 μ M); (n) merged images of (l) and (m); (n) fluorescence intensity correlation plot of (n); Green channel emission was collected in 470–550 nm upon excitation at 458 nm, and red channel emission was collected in 575–700 nm upon excitation at 561 nm. Scale bar: (a–j) 25 μ m, (k–o) 50 μ m.



Fig. S8 Fluorescence confocal images of living HeLa cells with dye **1b** (2 μ M), Nile red (1 μ M), MitoTracker [®] Red CMXRos (0.5 μ M) or Lyso-Tracker Red (0.5 μ M). (a, f, k) Bright–field images; (b, g, l) confocal images (green channels) of cells with dye **1b** (2 μ M); (c) confocal images (red channel) of cells with Nile red (1 μ M); (d) merged images of (b) and (c); (e) fluorescence intensity correlation plot of (d); (h) confocal images (red channel) of cells with MitoTracker [®] Red CMXRos (0.5 μ M); (i) merged images of (g) and (h); (j) fluorescence intensity correlation plot of (i); (m) confocal images (red channel) of cells with Lyso-Tracker Red (0.5 μ M); (n) merged images of (l) and (m); (n) fluorescence intensity correlation plot of (n); Green channel emission was collected in 470–550 nm upon excitation at 458 nm, and red channel emission was collected in 575–700 nm upon excitation at 561 nm. Scale bar: 25 μ m.



Fig. S9¹H NMR of dye **1a**.





Fig. S11 ¹³C NMR of dye 1a.



Fig. S12 ¹³C NMR of dye 1b.



Fig. S13 HRMS(ESI^+) of dye **1a**.



