

## Supplementary Information

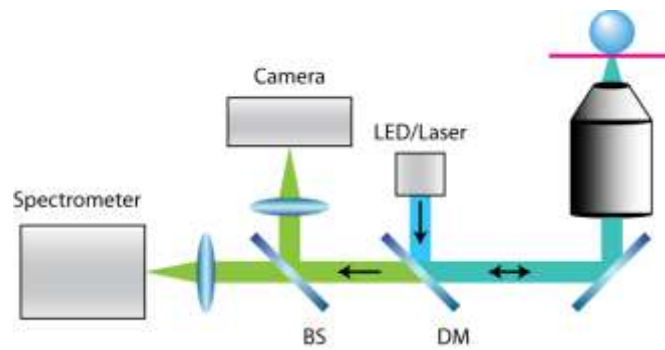


Fig. S1. Optical setup. An inverted microscope is used to image, excite and collect light from the sample. For the bioluminescence experiments the LED/laser source and dichroic mirror (DM) are removed.

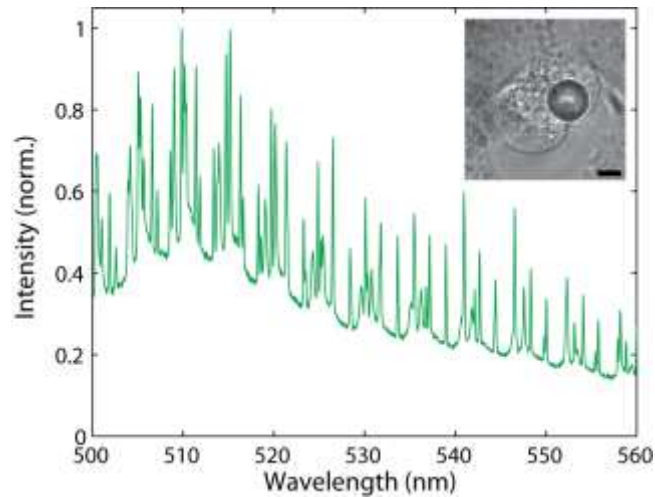


Fig. S2. Spectrum from bead in a cell. WGMs in the spectra of a GFP coated  $\text{BaTiO}_3$  bead in a HeLa cell undergoing cellular division (in Fig. 2). Scale bar, 5  $\mu\text{m}$ .

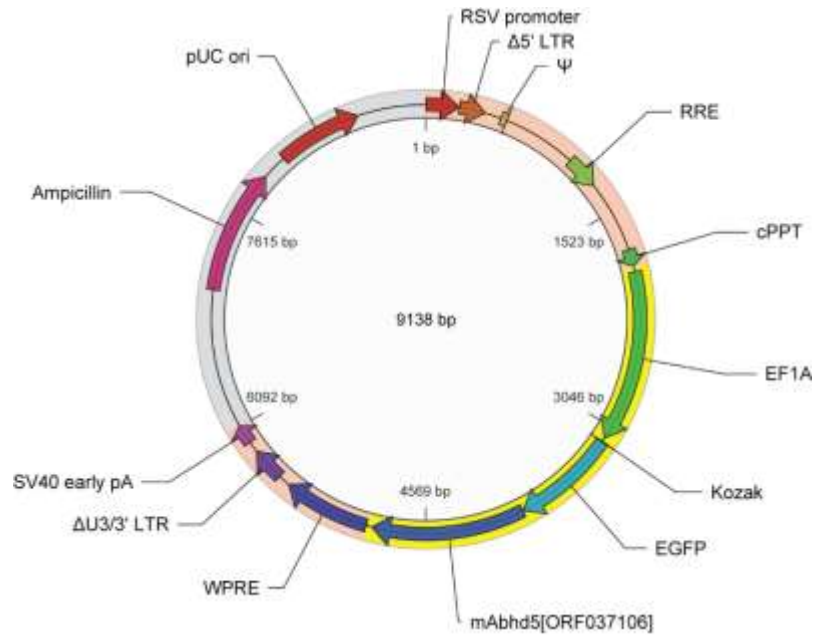


Fig. S3. GFP-ABHD5. A genetic map of the vector used in the transduction of 3T3-L1 adipocytes.

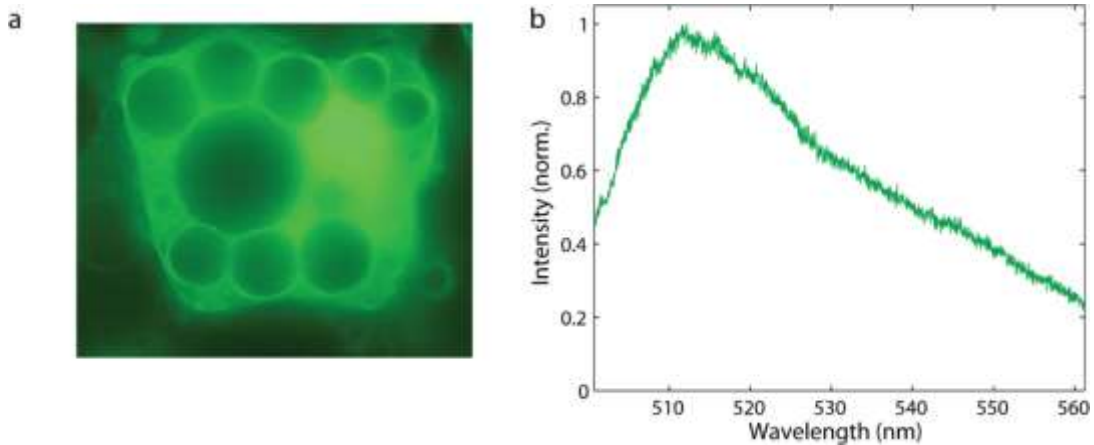


Fig. S4. Fluorescence of a GFP-transduced adipocyte, without ABHD5. (a) A typical fluorescent image showing bright emission from the cytoplasm, whereas fluorescence in adipocytes transfected with GFP-ABHD5 were more predominantly produced at the surface of lipid droplets (Fig. 4b). (b) A typical emission spectrum showing broad spontaneous emission from the GFP and no apparent spectral features associated with the WGMs of lipid droplets.

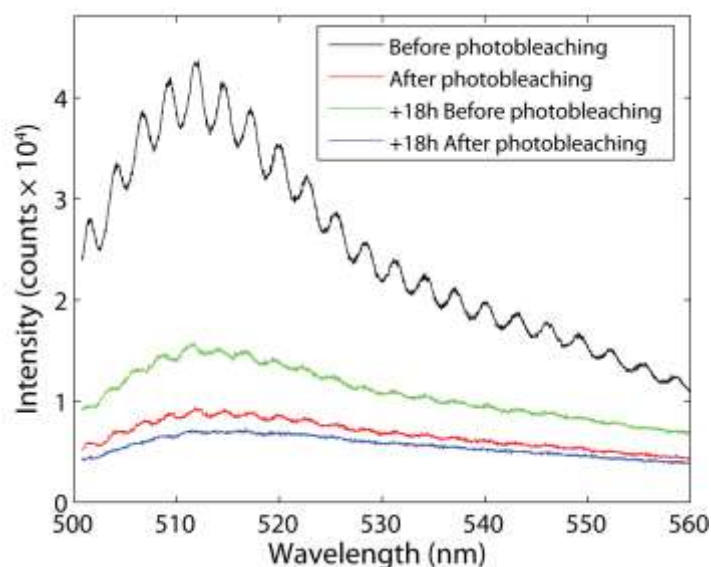


Fig. S5. Time-lapse spectra showing the photobleaching and recovery of GFP fluorescence from a GFP-ABHD5-transfected adipocyte (in Fig. 4c).

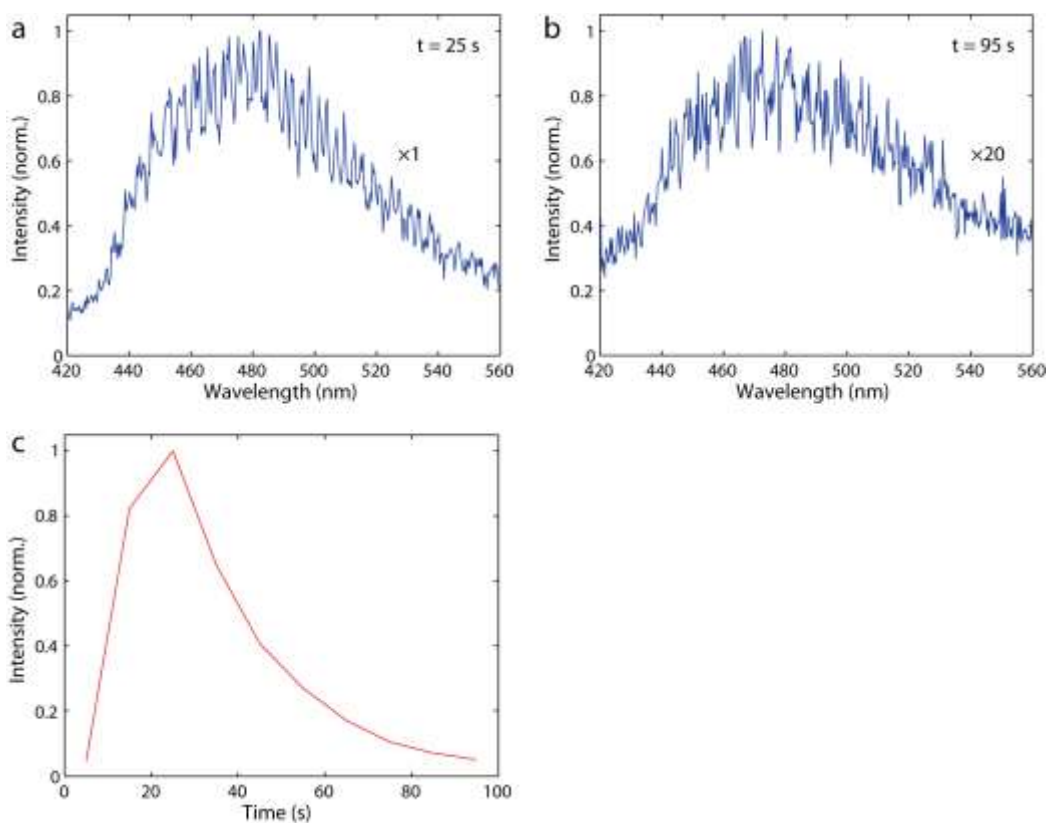


Fig. S6. Cavity-modified bioluminescence. (a,b) Spectra of light emitted by the bead in Fig. 4 at 25 s after the addition of luciferin when the light output is at maximum, (a), and after 95 s, (b). (c) Time trace of the bioluminescence intensity after CTZ was added to the beads at  $t = 0$ . The total intensity initially increases and then decreases exponentially.

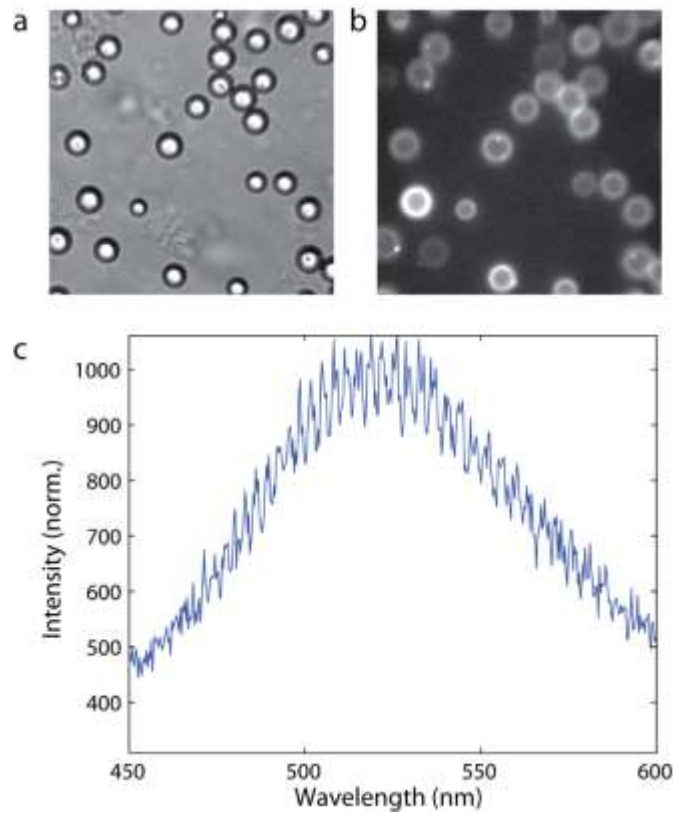


Fig. S7. RLuc8.6-coated glass beads. (a) Bright-field image of soda lime glass beads with diameters ranging from 15 to 19  $\mu\text{m}$  (Cospheric). The beads were functionalized according to a protocol described by Ribeiro *et al.* in “Immobilization of luciferase from a firefly lantern extract on glass strips as an alternative strategy for luminescent detection of ATP,” *J. Biolumin. Chemilumin.* **13**, 371–378, 1998. Briefly, we used 3 mg/ml Renilla luciferase (RLuc8.6) and 3 mg/ml poly-L-lysine hydrobromide (MW 30,000-70,000, Sigma) in Tris buffer (pH 7.4) and incubated beads in this solution at 4°C for 24 h. The beads were then washed five times with Tris buffer and transferred to PBS. (b) Bioluminescent image of the beads. (c) Spectrum of light from a single bead. WGM spectral peaks are visible.

### GFP-ABHD5 vector sequence used in the transduction of 3T3-L1 adipocytes

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