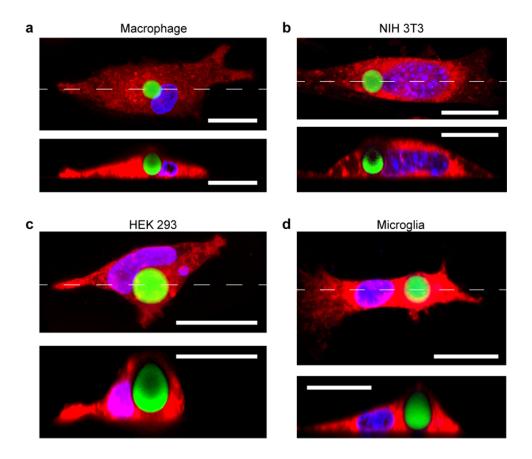
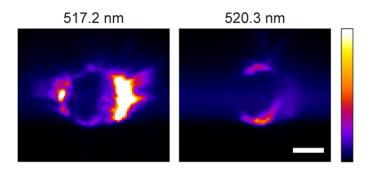
Lasing within live cells containing intracellular optical microresonators for barcode-type cell tagging and tracking

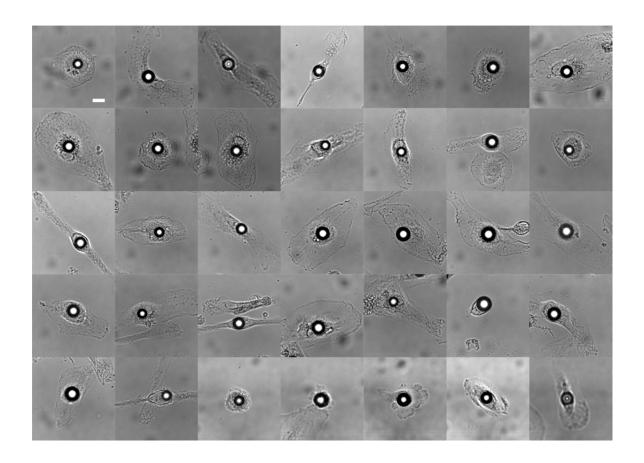
Supplementary Information



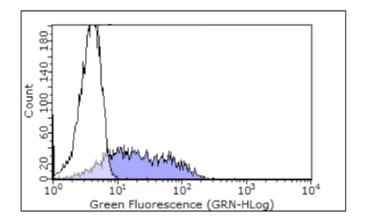
Supplementary Fig. 1: Confocal laser scanning microscopy (CLSM) data of different cells with cytoplasm (red fluorescence), cell nucleus (blue fluorescence) and internalized micro-sphere resonator (green fluorescence). Maximum intensity projection (top) and cross section along the dashed line (bottom). a, primary human macrophage cells, b, NIH 3T3 fibroblasts, c, Human Embryonic Kidney 293 and d, primary mouse microglia. For all four cell types, spontaneous internalization of micro-sphere resonators was observed. Scale bars in all panels, 20 µm.



Supplementary Fig. 2: Hyperspectral images (false colour) of the spatial emission pattern of a 5.5 µm radius cell internalized micro-sphere laser at 517.2 and 520.3 nm wavelength. Scale bar, 5 µm. Emission at 517.2 nm is attributed to a TE mode, emission at 520.3 nm is attributed to a TM mode. The definition of TM and TE is with respect to the surface of the micro-sphere.



Supplementary Fig. 3: Bright field images of the cell lasers discussed in Fig. 3c of the main text. Scale bar, $20~\mu m$.



Supplementary Fig. 4: CD14 expression of purified macrophages. Plastic adherent cells from isolated peripheral blood mononuclear cells were recovered and stained with CD14-FITC antibody (eBioscience, Hatfield, UK), shown shaded, compared to auto-fluorescence from the same cell population without CD14-FITC antibody staining (unshaded). Analysis was performed on a Guava 8HT flow cytometer (Millipore, UK).

Supplementary Video 1: Representative time-lapse video of a macrophage cell, starting 1 h after addition of micro-spheres to the culture dish. Video 50x accelerated. During its random migration, the cell reached a micro-sphere and within less than five minutes internalized the entire sphere. Afterwards, the cell remained motile, dragging the internalized sphere along.

Supplementary Video 2: Confocal Laser Scanning Microscopy image stack of macrophage with cytoplasm (red fluorescence), cell nucleus (blue fluorescence) and internalized micro-sphere resonator (green fluorescence).

Supplementary Video 3: Time lapse video of the macrophage cells investigated in Fig. 3a and b of the main text over the whole 19h time period of the experiment.