Fluorescent squaramide ligands for cellular imaging and their encapsulation in cubosomes

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Figure S1 Normalized emissions (A and B) and absorptions (C and D) for L1 and L2, respectively of DMSO solution (1 mM) diluted in water (25 μ M). λ_{Ex} 355 nm.



Figure S2. 293T (up) and Caco-2 cells (down) incubated for 10 min with L1 ligand 200 μ M.



Figure S3 Normalized emissions of L2 (red) and L2 in the presence of 1 eq of TBAH₂PO₄ (black) of a DMSO solution (1 mM) diluted in water (25 μ M). λ_{Ex} 480 nm.



Figure S4. 293T cells incubated for 30 min with cubosomes containing L1 ligand 0.035 µM. A) Bright filed B) Green filed C) merge image of bright and green filed.



Figure S5. Image of 293T cells without cubosomes (left), treated with 10 μ l (middle) and 20 μ l (right) of cubosomes and stained with 0.1 % Trypthan Blue after 24 h. iLED, 40x.





Figure S7. ¹³C-NMR spectrum of **L1** in DMSO- d_6 .



Figure S8. ¹H-NMR spectrum of **L2** in DMSO- d_6 .



Figure S9. ¹³C-NMR spectrum of L2 in DMSO- d_6 .



Figure S10. ESI-MS spectrum of L1.



Figure S11. ESI-MS spectrum of L2.