Interaction of Nucleic Acids with the Glycocalyx

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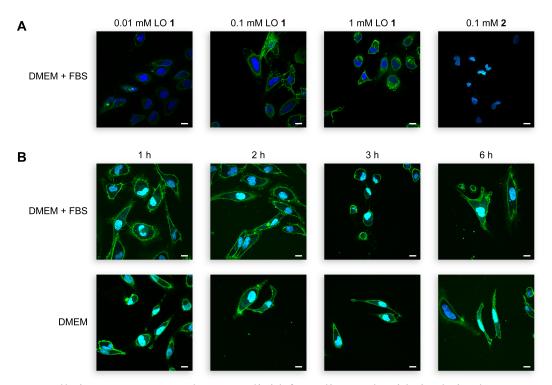


Figure S1. Cells incorporate a LO but not a lipid-free oligonucleotide in their plasma membrane. Images were obtained by confocal microscopy. LO **1** and oligonucleode **2** are green (fluorescein); cell nuclei are blue (Hoechst 33342). (A) LO **1** was suspended in DMEM containing 10% v/v FBS and incubated with HeLa cells at the indicated concentration for 12 h. Oligonucleotide **2** was likewise incubated with HeLa cells for 4 h. (B) LO **1** (0.1 mM) was incubated with or without 10% v/v FBS for the indicated times. Scale bars: 10 μm.

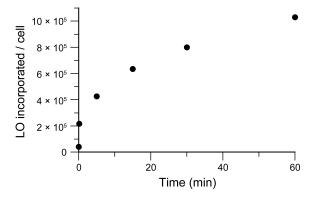


Figure S2. Cellular incorporation of a LO depends on time. LO **3** (10 mM) was incubated with HeLa cells for the indicated times, and incorporation was measured by flow cytometry.

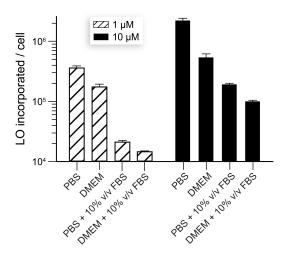


Figure S3. Cellular incorporation of a LO is affected by the medium. LO **3** (1 or 10 μ M) was incubated with HeLa cells for 15 min in the indicated media, and incorporation was measured by flow cytometry. n = 3; error bars: \pm SD.

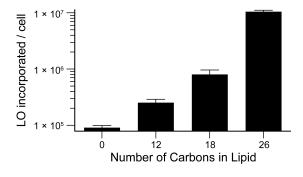


Figure S4. Cellular incorporation of LOs is affected by hydrophobicity. LO **9** (no lipid), LO **10** (C_{12}), LO **3** (C_{18}), and LO **11** (C_{26}) at 10 μ M were incubated with HeLa cells for 15 min, and incorporation was measured by flow cytometry. Error bars: \pm SD.

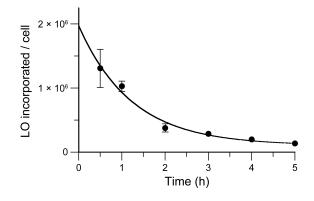


Figure S5. Cells release a long LO after incorporation. Fluorescent LO **11** (10 μ M), which has a C26 lipid tail, was incorporated into the plasma membrane of HeLa cells, and retention was measured by flow cytometry. Error bars: \pm SD.

