Supporting Information

Synthesis of Fluorosurfactants for Emulsion-Based Biological Applications

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Supplementary Figures



Figure S1. (a) Photograph of the synthesized PFPE-Tris surfactants with increasing the feed molar ratio of Tris to PFPE from 0.3, 0.5, 0.8 to 1.0 and (b) their full-scale FT-IR spectra. (c) Comparison of FT-IR spectra of the synthesized PFPE-Tris surfactants at Tris/PFPE feed molar ratios of 1.0 (PFPE₁-Tris_{1.0}) and 1.2 (PFPE₁-Tris_{1.2}) showed no major difference, indicating a complete conversion of PFPE-Tris occurred at the feed molar ratio of 1.0.



Figure S2. Amount of DNA recovered from w/o emulsion after disrupting the droplets by a droplet breaking agent.



Figure S3. Luciferase activities of the transfected cells. Cells were transfected *in vitro* using jetPEITM/DNA complexes synthesized in w/o emulsion droplets stabilized with different test surfactants. The jetPEITM/DNA polyplexes formulated without emulsion treatment was used as a positive control.



Figure S4. Survival rate of hMSCs cultured in the emulsion droplets. hMSCs maintained their viability (>85%) in the w/o/w droplets for at least 4 days.