Supporting Information

Synthesis of lipo-glycopolymers for cell surface engineering

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Figure S1. ¹H NMR spectrum of MAL-DPPE. MAL-DPPE: ¹H NMR (600 MHz, CDCl₃) δ 6.70 (s, 2H), 5.19 (s, 1H), 3.90 (s, 4H), 3.69-3.74 (m, J=6 Hz, 2H), 3.46-3.48 (m, 2H), 3.16-3.20 (q, J=6 Hz, 2H), 2.27-2.31 (q, J=6 Hz, 4H), 2.19-2.21 (m, 2H), 1.41-1.47 (m, 10H), 1.24-1.30 (m, 48H), 0.86-0.88 (t, J=6 Hz, 6H). MS for C₄₇H₈₅N₂O₁₁P [M + H] ⁺ m/z: 886.1700 (calculated); 885.9731 (found). MS for C₄₇H₈₅N₂O₁₁P [M + Na] ⁺ m/z: 908.1600 (calculated); 907.5648 (found).



Figure S2. ¹³C NMR spectrum of MAL-DPPE. MAL-DPPE: ¹³C NMR (600 MHz, CDCl₃) δ 174.24, 173.97, 171.11, 134.12, 70.32, 63.93, 62.74, 43.52, 40.03, 37.28, 34.09-34.22, 31.92, 29.16-29.75, 28.00, 25.87, 24.92, 22.68, 14.10.



Figure S3. ³¹P NMR spectrum of MAL-DPPE. ³¹P NMR (600 MHz, CDCl₃) $\delta \Box$ 4.21.



Figure S4. IR spectra of DPPE and MAL-DPPE. FT-IR spectrum showed that the peak at 1700 cm⁻¹ attribute to C=O from amide bonds and C=C bonds, and the peaks at 3106 cm⁻¹, 3207 cm⁻¹ and 1435 cm⁻¹ correspond to CH stretches from carbon-carbon double bonds, N–H and C–N bonds, respectively.



Figure S5. UV spectra of pSMF and pSMF-SH over a wavelength range of 260-400 nm. The disappearance of the peak of thioester groups at 296 nm demonstrated a successful reduction of copolymers.

Sample	Feed ratio	Ratio of ¹ H-NMR	M _n	PDI
	(SS : MAG)	(SS : MAG)	(GPC)	
pSM	1:1	1.4 : 1	4600	1.17



Figure S6. GPC traces of pSMF-SH and DPPE-pSMF.



Figure S7. (a) ¹H-NMR spectrum of DPPE-pSM. (b) Infrared spectra of pSM and DPPE-pSM.

Table S2. The density of thiol groups of pSM-SH and DPPE-pSM.



Figure S8. Dynamic light scattering plot of DPPE-pSM ($C = 10 \mu M$) at 37°C in water.



Figure S9. Confocal micrographs of HeLa cells incubated for 1 h with 3.4 μ M of DPPE-pSMF. The cells were stained with DAPI, a nuclear counterstain, before imaging (scale bar: 20 μ m).



Figure S10. Flow cytometry analysis. HeLa cells were incubated with different concentrations of DPPE-pSMF and then analyzed by flow cytometry. Cell surface incorporation of lipid-anchored glycopolymers onto cell membranes depends on polymer concentrations.