

Supporting Information

Protection/Deprotection of Surface Activity and Its Applications in the Controlled Release of Liposomal Contents

Xueshu Li and Yan Zhao *

Department of Chemistry, Iowa State University, Ames, Iowa 50011-3111, USA

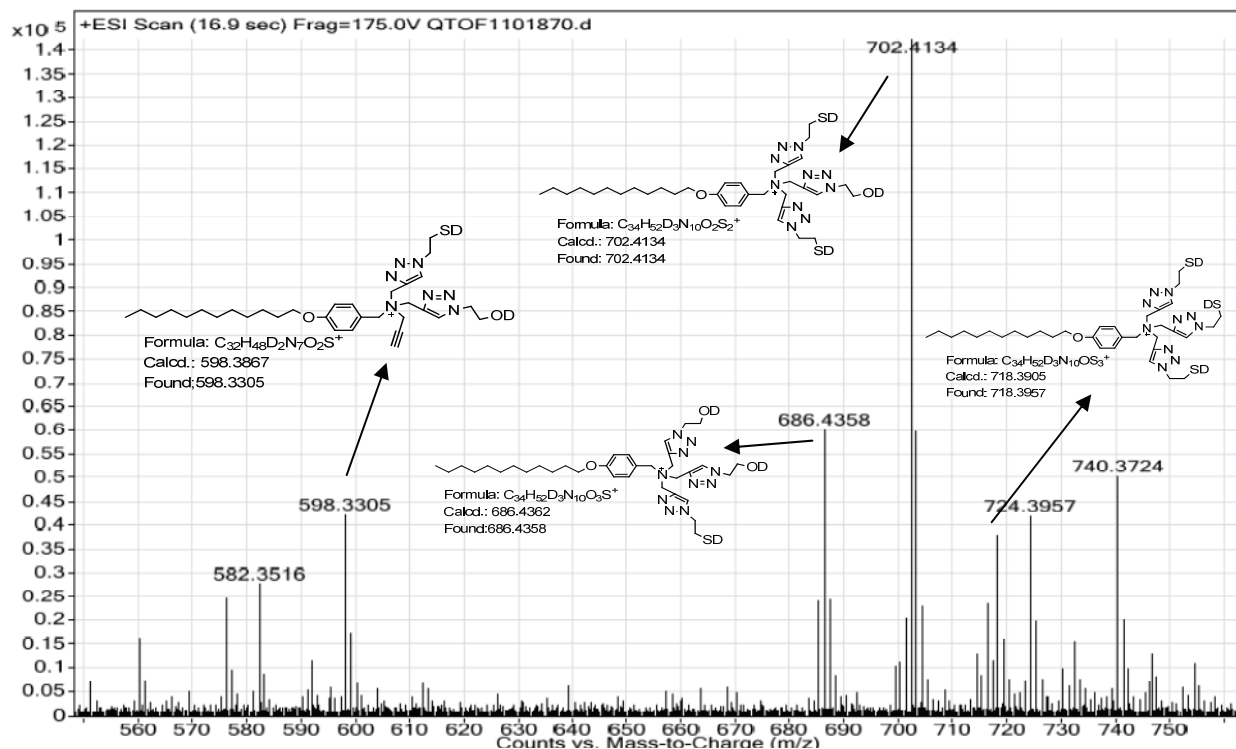


Figure 1S. ESI-MS spectra of DTT-digested nonfluorinated SCM-CH₂CH₂OH. Deuterated peaks appeared because D₂O/H₂O (3/1) was used in the preparation of SCM (to facilitate monitoring of the reaction by ¹H NMR spectroscopy).

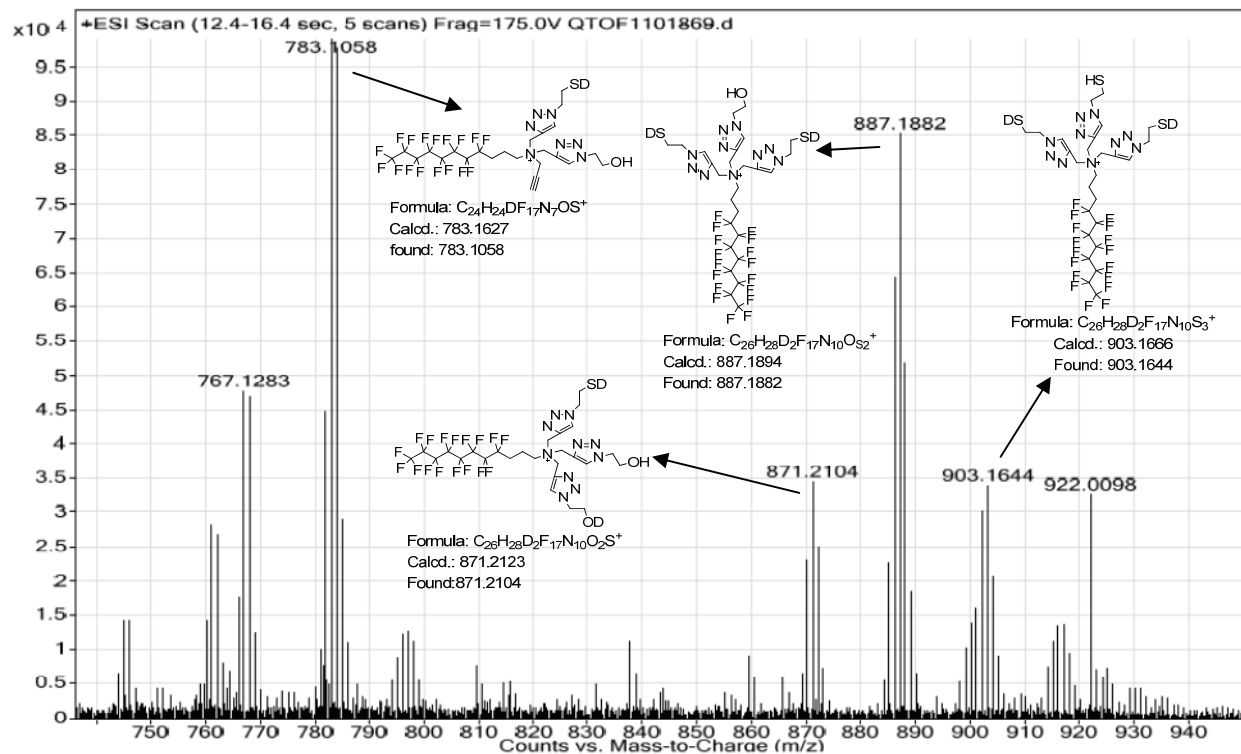


Figure 2S. ESI-MS spectra of DTT-digested fluorinated SCM-CH₂CH₂OH. Deuterated peaks appeared because D₂O/H₂O (3/1) was used in the preparation of SCM (to facilitate monitoring of the reaction by ¹H NMR spectroscopy).

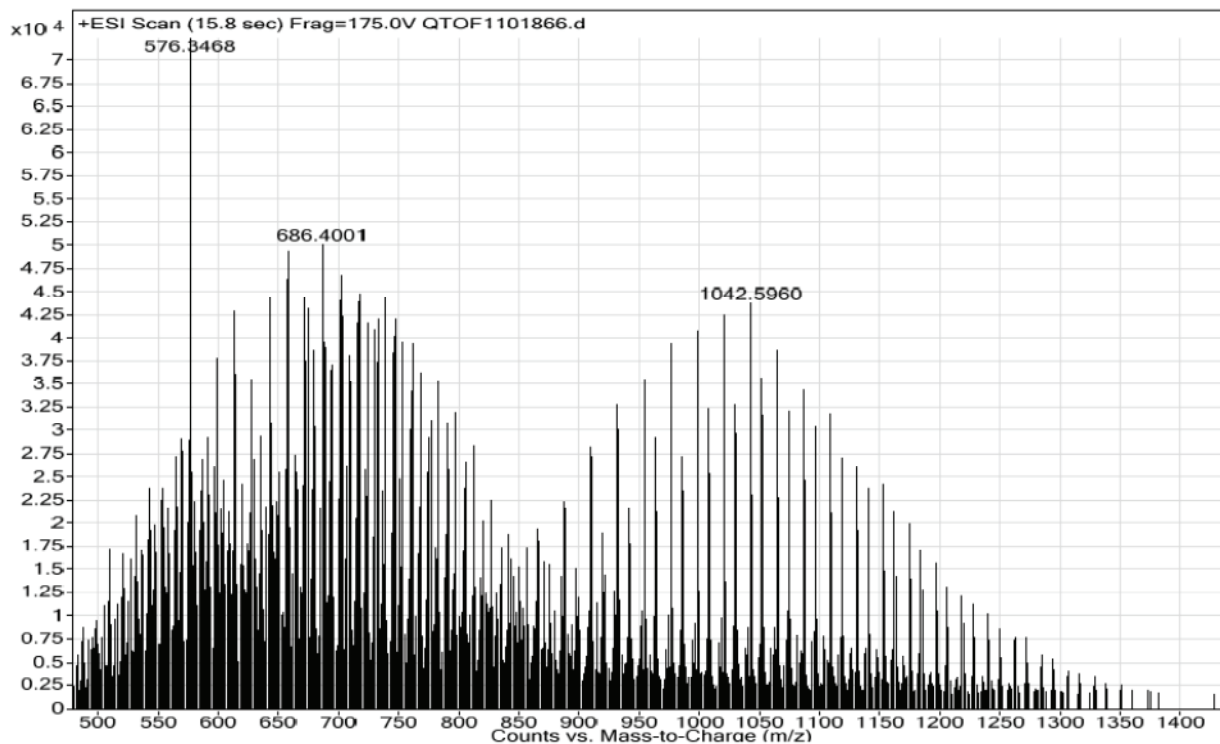


Figure 3S. ESI-MS spectra of DTT-digested nonfluorinated SCM-PEG.

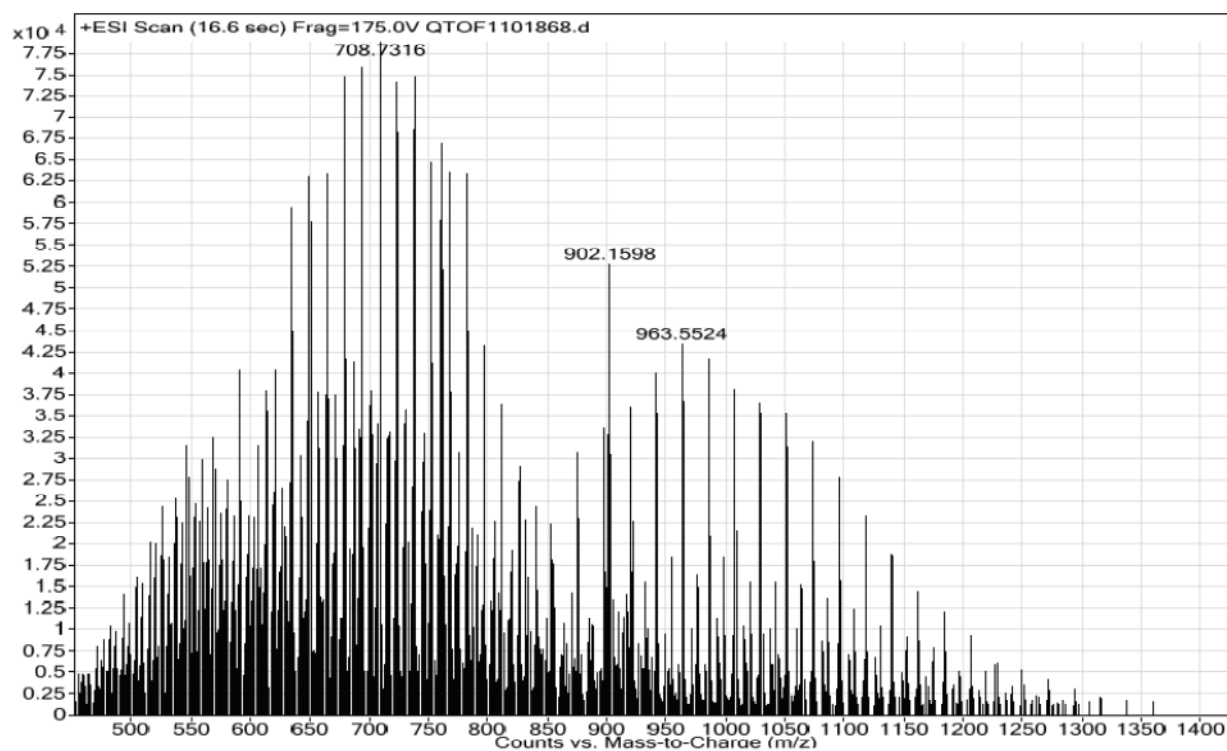


Figure 4S. ESI-MS spectra of DTT-digested fluorinated SCM-PEG.

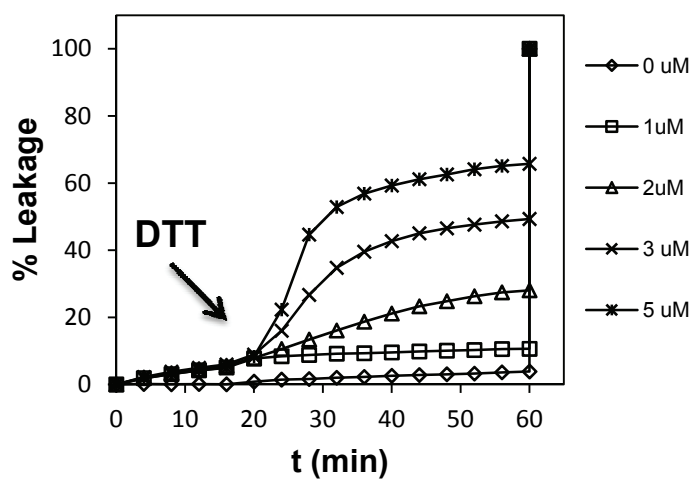


Figure 5S. Percent leakage of CF from POPC/POEPC LUVs with 30 mol % cholesterol triggered by SCM-PEG and DTT. [DTT] = 1 mM. The concentrations of (cross-linked) surfactant in SCM are shown on the side.

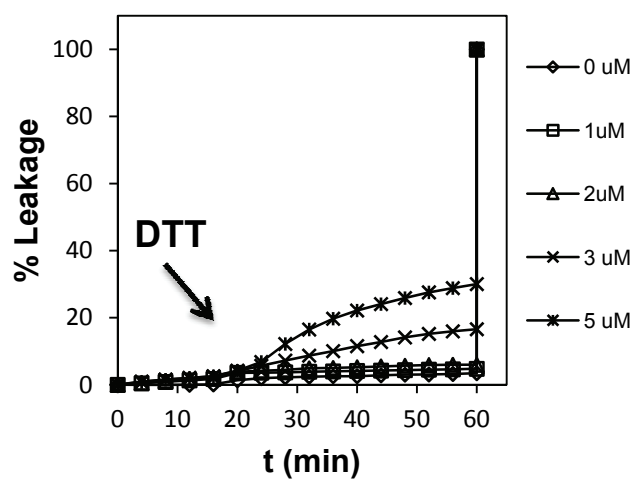


Figure 6S. Percent leakage of CF from POPC/POEPC LUVs with 50 mol % cholesterol triggered by SCM-PEG and DTT. [DTT] = 1 mM. The concentrations of (cross-linked) surfactant in SCM are shown on the side.

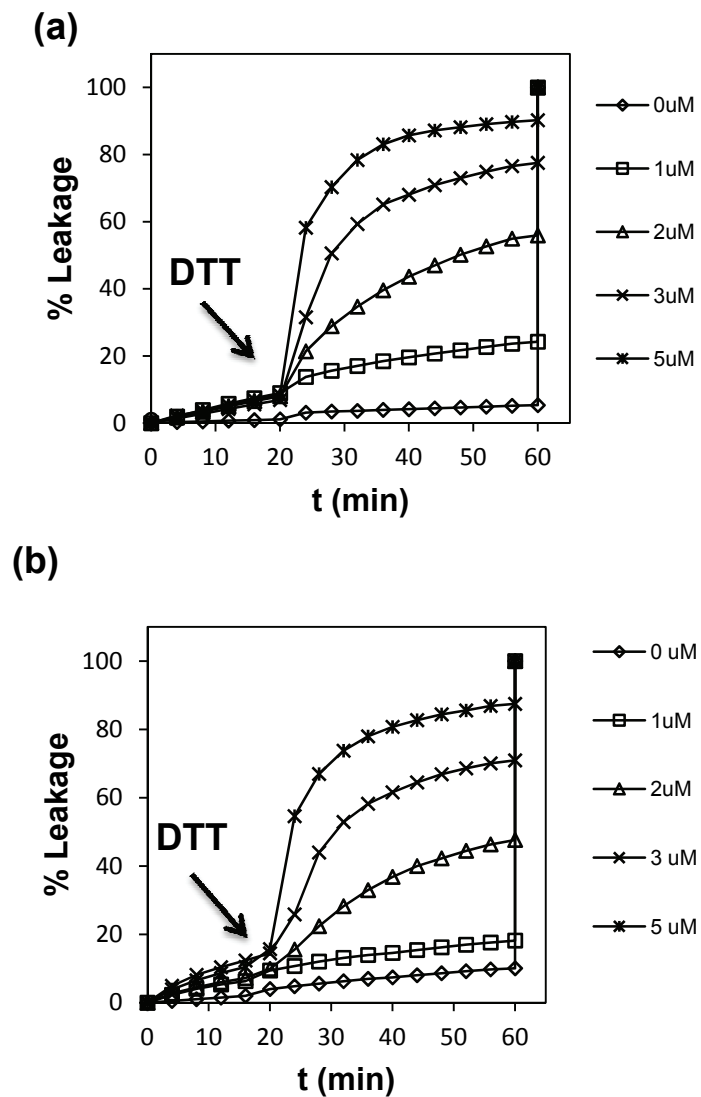


Figure 7S. Percent leakage of CF from POPC/POEPC LUVs triggered by (a) SCM-CH₂CH₂OH and (b) SCM-PEG. [DTT] = 1 mM. The concentrations of (cross-linked) surfactant in SCM are shown on the side.