

Supplementary Information for

In Cellulo and In Vivo Comparison of Cholesterol, Beta-Sitosterol and Dioleoylphosphatidylethanolamine for Lipid Nanoparticle Formulation of mRNA

Ayoub Medjmedj ^{1,†}, Albert Ngalle-Loth ^{1,†}, Rudy Clemençon ¹, Josef Hamacek ^{1,2}, Chantal Pichon ^{1,2} and Federico Perche ^{1,*}

¹ Centre de Biophysique Moléculaire UPR4301 CNRS, Rue Charles Sadron, 45071 Orléans, France; ayoub.medjmedj@cnrs-orleans.fr (A.M.); albert.ngalle-loth@cnrs-orleans.fr (A.N.-L.); rudy.clemencon@cnrs.fr (R.C.); josef.hamacek@cnrs.fr (J.H.); pichon@cnrs.fr (C.P.)

² Centre de Biophysique Moléculaire, University of Orléans, 45100 Orléans, France

* Correspondence: federico.perche@cnrs-orleans.fr; Tel.: +33-2-38-25-55-44

† These authors contributed equally to this work.

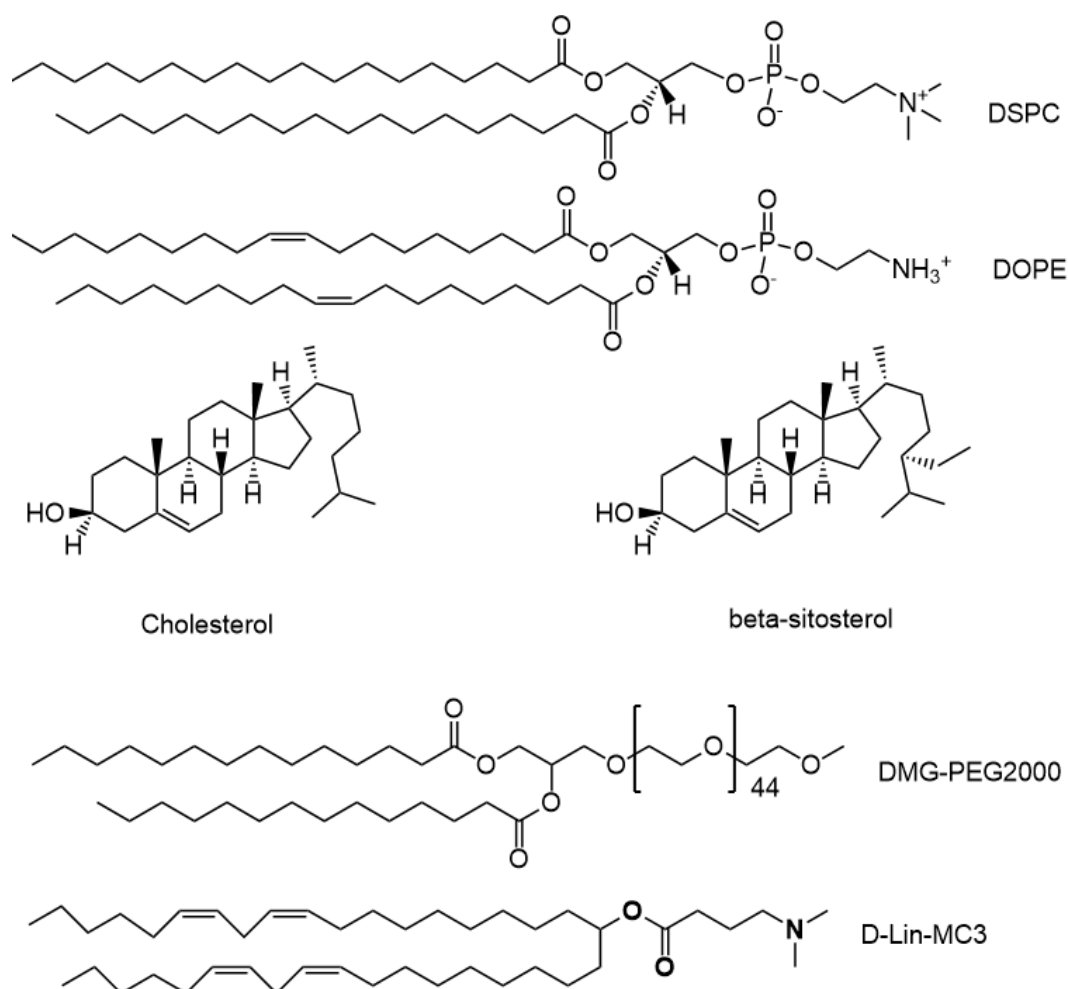
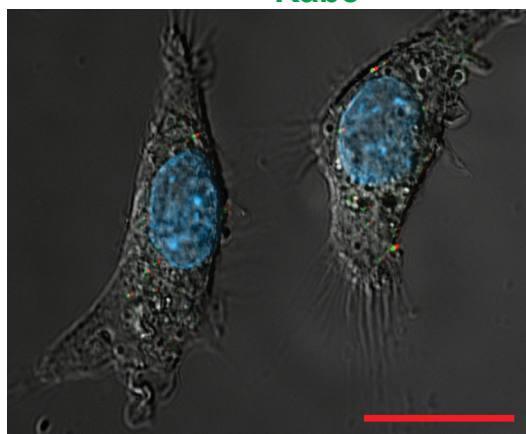


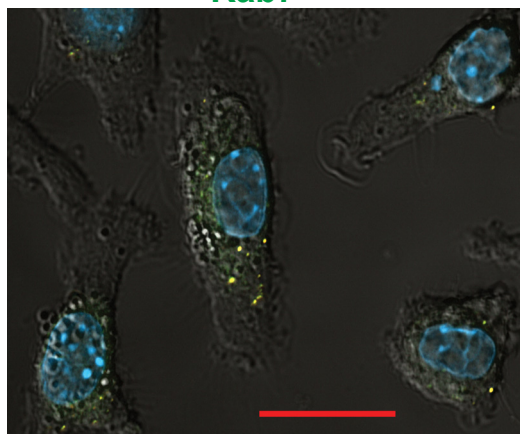
Figure S1. Structures of the lipids used in this study.

LFM / Cy3mRNA

Rab5



Rab7



Rab11

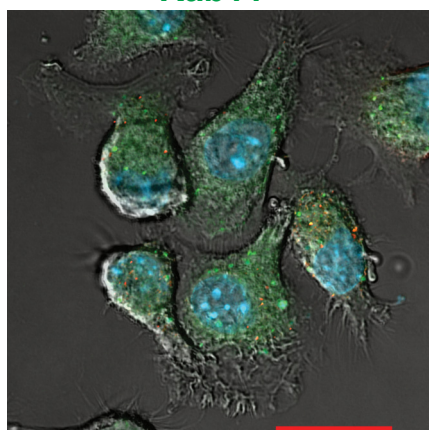


Figure S2. Intracellular trafficking of Cy3 mRNA-LFM complexes in DC 2.4 cells by confocal microscopy with either Rab5-GFP, Rab7-GFP or Rab11-GFP. Nuclei were stained with DAPI. Bar represents 20 μ m.

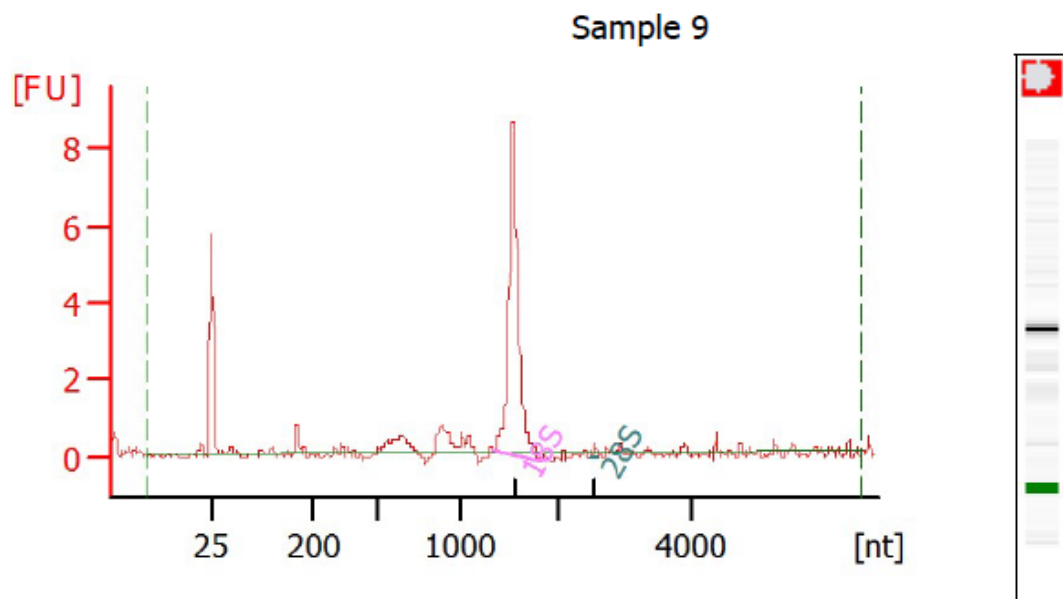


Figure S3. Agilent analysis of the GFP mRNA produced by *in vitro* transcription.