

Supporting Information

Characterization of Lipid-Based Hexosomes as Versatile Vaccine Carriers

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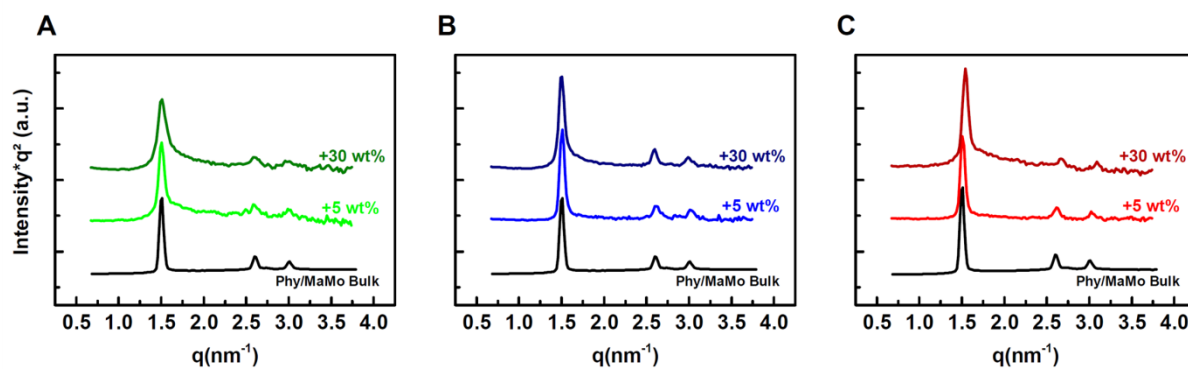


Figure S1. Structure characterization using SAXS of Phy/MaMo (14 wt%) formulations containing poloxamer 407 (A), Pluronic F108 (B), and Myrj 59 (C) at 5 and 30 wt% of total lipid. The black curve represents the non-dispersed bulk phase without stabilizer. Measurements were performed at 25°C. The curves were vertically shifted for clarity.

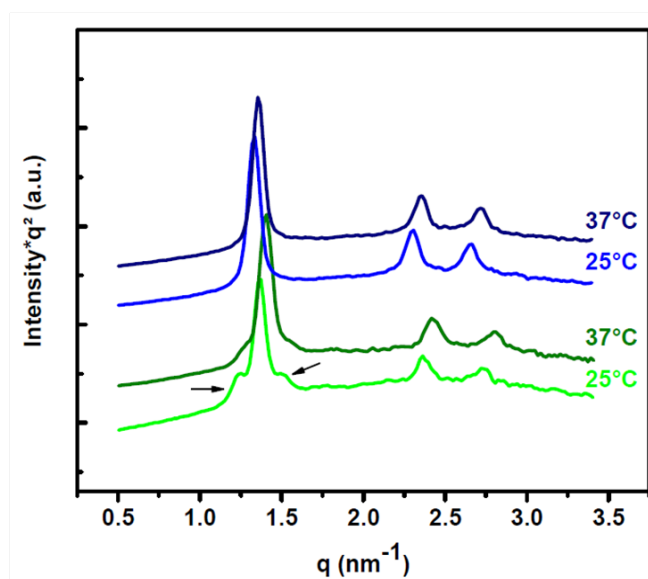


Figure S2. Effect of the headgroup dimension on the internal structure of Phy/MaMo (14 wt%) hexosomes stabilized with 10 wt% poloxamer 407. SAXS data is displayed for hexosomes containing the same molar concentration (ca. 14 mol%) of DDA (green) and Dotap (blue) at 25°C and 37°C. Arrows indicate perturbation of the pure HII peak profile. The curves were vertically shifted for clarity.

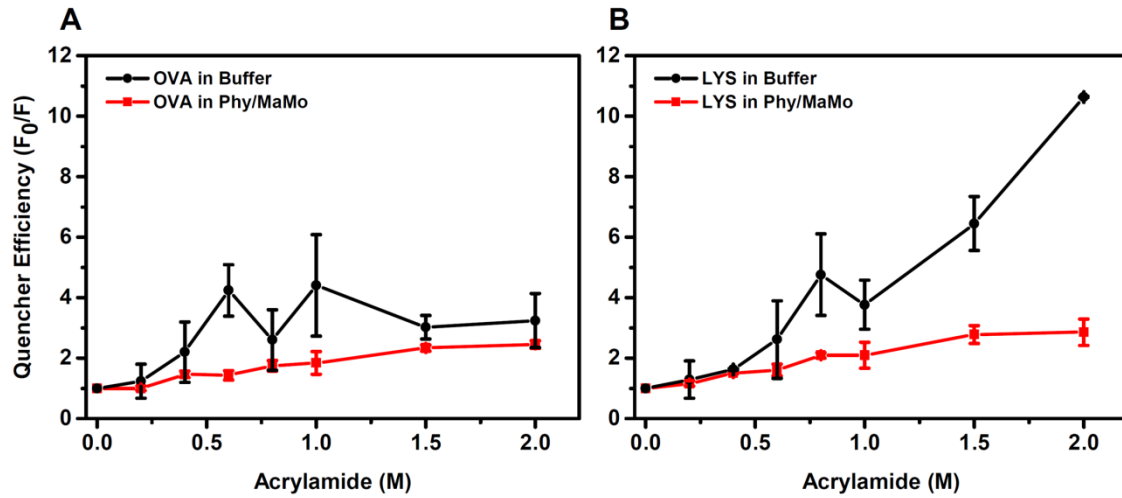


Figure S3. Tryptophan accessibility assay with acrylamide in Phy/MaMo (14 wt%) hexosomes stabilized with 25 wt% poloxamer 407, and loaded with 0.1 wt% (of total lipid) ovalbumin (A) or lysozyme (B). Unquenched protein (F_0); Quenched protein (F). Data is shown as mean \pm SD of two independent experiments performed in triplicate.