# Lipidomic and in-gel analysis of maleic acid co-polymer nanodiscs reveals differences in composition of solubilized membranes

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#### **Supplementary Method**

IR-spectra of the xMA polymers

## Styrene maleic acid (2.3:1)

IR (ATR) v: 2927, 1566, 1492, 1411, 759, 698 cm<sup>-1</sup>.



**Diisobutylene maleic acid (1:1)** 

IR (ATR) v: 2947, 1709, 1366, 1186, 921 cm<sup>-1</sup>.



#### Styrene maleic anhydride (2.3:1)



IR (ATR) v: 2923, 1855, 1774, 1493, 1453, 914, 698 cm<sup>-1</sup>.

## Styrene trimethyl-ethylamime maleimide (SMA-QA)







Superimposition of SMA-QA (in orange) and SMAnh IR (in grey) spectra.



**Figure S1**. Saturation of PL species present in *E. coli* classified depending on their headgroup, being: PG, PE, PI, PS or PA. The number of double bonds indicated are those found in the total fatty acid chains of a single PL species<sup>‡</sup>.



**Figure S2**. Fatty acid chain length distribution in *E. coli* samples represented by the number of carbons present in the fatty acid chains of the total PI, PG, PS or PA species<sup>‡</sup>.



Figure S3. Distribution of species present in the SL fraction of solubilized Jurkat membranes<sup>‡</sup>.



**Figure S4**. Distribution of species present in the GL fraction of solubilized Jurkat membranes<sup>‡</sup>. DG stands for diacylglycerol and TG for triacylglycerol<sup>‡</sup>.



**Figure S5.** Analysis of the PC and PE fractions of Jurkat membranes. The found different species are: intact PE or PC, being PE or PC with two acyl chains; lysoPE or lysoPC if one of the acyl chain is lost; PE(P) or PC(P) if one of the esther bonds is replaced by an alkenyl ether linkage and PE(O) or PC(O) when the esther linkage is replaced by an alkyl ether bond <sup>‡</sup>.



Figure S6. Saturation degree of Jurkat membranes. Represented from top to bottom, saturation profile of: all lipid species; PL; SL and GL<sup>‡</sup>.



**Figure S7**. Saturation of PL species present in Jurkat membranes classified depending on their headgroup, being: PC, PG, PE or PI. The number of double bonds indicated are those found in the total fatty acid chains of a single PL species<sup>‡</sup>.



**Figure S8**. PL fatty acid chain length distribution in Jurkat samples represented by the number of carbons present in the fatty acid chains of the total PG, PE or PI species<sup>‡</sup>.





**Figure S9**. Fatty acid chain length distribution of the SL fraction present in Jurkat samples represented by the number of carbons present in the fatty acid chains<sup>‡</sup>.

## Notes

 $\ddagger$  Data points correspond to three technical replicates. Error bars represent  $\pm$  S.D. Significant differences (upon one-way ANOVA) are denoted as \*(p < 0.05), \*\*(p < 0.01), \*\*\*(p < 0.001), \*\*\*\*(p < 0.001).