Packing and Mobility of Hydrocarbon Chains in Phospholipid Lyotropic Liquid Crystalline Lamellar Phases and Liposomes, Characterised Using Positron Annihilation Lifetime Spectroscopy (PALS)

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Supplementary Information



Figure S1:A) Generalised phase behaviour for phosphocholine phospholipids with lipid chain length C12-18. n represents the mol ratio of water to lipid molecules.¹ B) and C) The binary lipid-water phase diagram of bulk DMPC² and DPPC³, respectively. The red line represents the composition at which PALS measurements were made with increasing temperature.



Figure S2: Cryo-TEM images of the liposome samples for A) DLPC, B) DOPC, C) DPPC and D) DSPC.



Figure S3: oPs lifetime in the organic region (τ_4) for DPPC and its corresponding unsaturated lipid (C16:1) as a function of temperature.



Figure S4: oPs lifetime in the organic region (τ_4) for DSPC and DOPC as a function of temperature.



Figure S5: oPs intensity in the organic region (I_4) for DPPC and its corresponding unsaturated lipid (C16:1) as a function of temperature.



Figure S6: oPs intensity in the organic region (I_4) for DPPC and DOPC as a function of temperature.



Figure S7: oPs intensity in the aqueous region (I_3) for DPPC and its corresponding unsaturated lipid (C16:1) as a function of temperature.



Figure S8: oPs intensity in the aqueous region (I_3) for DSPC and DOPC as a function of temperature.



Figure S9: I₄ versus standard deviation of τ_4 for the lamellar mesophase and liposomes of DPPC. The plot shows that reducing oPs intensity values (I₄) results in increasing uncertainties for the corresponding oPs lifetime (τ_4).

References

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