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Characterization of Solid Supported Dipalmitoylphosphatidylcholine Membranes Containing Cholesterol

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ABSTRACT: The incorporation of cholesterol into dipalmitoylphosphatidylcholine (DPPC) membranes, even in small amounts, has been shown to significantly alter the properties of the membrane. In this work, force-distance interaction profiles of DPPC membranes containing 8 mole% cholesterol obtained using the surface force apparatus are analyzed in the context of high resolution structural characterization by atomic force microscopy and neutron reflectometry. The adhesion between the mixed membranes was greater than that for pure DPPC and was variable – depending on the number of defects in the outer membrane leaflets. These defects were only detectable by atomic force microscopy and had an average size of 230 ± 30 nm and 1-5% surface density in the outer leaflet. The adhesion between the membranes monotonically increased as the thickness of the membrane decreased – in direct correlation with the number of defects present (exposed hydrophobic groups) in the membrane contact region. Due to the low diffusion rate of gel phase membranes, the interaction force profiles were stable and no membrane restructuring was observed.

Key Words: DPPC, cholesterol, supported lipid bilayer, SFA, AFM, zeta potential, fluorescence microscopy, neutron scattering

INTRODUCTION

Cholesterol is an important building block in biological membranes and is found in every cell in the human body. Cholesterol's ubiquitousness underlines its important role in altering the physical^{1,2} and thermodynamic properties of membranes by changing the packing or the ordering of lipids, reducing membrane permeability^{3,4,5}, and in the formation of lipid rafts^{6,7,8,9,10}. In particular, cholesterol is known to have a condensing effect on fluid phase lipids and to conversely fluidize solid phase lipids which helps maintain the integrity of cell membrane when stressed.^{11,12} The concentration of cholesterol varies between different cell types and organelles with concentrations ranging from 5%¹³ to as high as 50%¹⁴ of the membrane. A plethora of studies have looked at the function of cholesterol in cell membranes and the effect of the addition of cholesterol in altering the intermolecular interactions in biomimetic

membrane systems.^{3, 15, 16} For example, phase diagrams of binary and ternary lipid mixtures containing cholesterol are well-documented^{17, 18}, and studies have also looked at the alteration of membrane mechanical properties due to preferential segregation of cholesterol^{2, 8}. In terms of high resolution structural characterization, Mills *et al.*¹⁹ quantified changes in the lamellar repeat distance in multilamellar DPPC vesicles with cholesterol using wide angle x-ray scattering. They found that, in general, addition of cholesterol resulted in the tighter packing of DPPC lipids which induced an increase in membrane thickness and lamellar spacing. Further, 8 mole% cholesterol had the largest effect on altering the packing of DPPC as the mixture remains in the gel phase. Higher cholesterol concentrations result in coexisting gel and liquid ordered phases.

The alteration of lateral intermolecular interactions by cholesterol is well-established from an array of earlier work, however much less is known regarding how the presence of cholesterol modifies the interactions between membranes. In this work, DPPC membranes with 8 mole% cholesterol were studied to quantify and understand how the addition of cholesterol to a DPPC membrane impacts supported membrane structure and membrane-membrane interactions in the gel phase. A comparison is made between measured membrane interaction force-distance profiles by surface force apparatus (SFA), neutron scattering structural characterization, vesicle zeta-potential (ZP), and membrane structure/topology as determined by atomic force microscopy (AFM).

MATERIALS AND METHODS

Chemicals. 1,2-dihexadecanoyl-*sn*-glycero-3-phosphoethanolamine (DPPE, melting point, $T_M = 63^\circ\text{C}$), 1,2-dipalmitoyl-*sn*-glycero-3-phosphocholine (DPPC, $T_M = 41^\circ\text{C}$), and cholesterol (ovine wool, >98%, $T_M = 148^\circ\text{C}$) were purchased from Avanti Polar Lipids, Inc. (Alabaster, AL) and used as received. 2-(4,4-Difluoro-5,7-Diphenyl-4-Bora-3a,4a-Diaza-s-Indacene-3-Pentanoyl)-1-Hexadecanoyl-*sn*-Glycero-3-Phosphocholine (β -BODIPY® 530/550 C₅-HPC) was purchased from Life Technologies (Carlsbad, CA). Electrolyte solutions used NaNO₃ 99.995% (Sigma, St. Louis, MO). Water was purified with a MilliQ gradient water purification system to a resistivity of 18 M Ω ·cm.

Sample Preparation. Mica substrate supported lipid bilayers were used in SFA, fluorescence microscopy (FM), and AFM studies. Neutron scattering studies used ultra smooth single crystal quartz. The membranes were constructed using Langmuir-Blodgett (LB) deposition (Nima Coventry, U.K.). The inner monolayer for SFA, AFM and FM experiments was DPPE LB deposited at 45 mN/m and dipping speed of 1 mm/min. Mixtures for FM imaging contained 0.5% of Bodipy-HPC. Previous studies show that LB deposited DPPE forms an almost defect free, robust and strongly physisorbed monolayer on mica with

transfer ratios of 0.997 ± 0.004 ²⁰ with a thickness of 2.56 ± 0.05 nm under these conditions.²¹ The tight packing and stability of the gel phase DPPE inner monolayer minimizes molecular exchange between the two leaflets and provides an ideal base for the 92:8 monolayer, which is the primary interest of this study. In all cases, the outer monolayer was 92:8 DPPC-cholesterol deposited at 30mN/m. For the SFA, FM, and AFM work, the outer leaflet was LB deposited with a dipping speed of 4 mm/min. The transfer ratio of the outer monolayer was 0.843 ± 0.057 . For neutron scattering measurements a symmetric membrane of 92:8 DPPC-cholesterol was used for ease of modeling and analysis. The inner leaflet was deposited by LB at 30 mN/m and dipping speed of 1 mm/min on quartz substrates. Because of the weaker adhesion of the membrane to quartz vs. mica, the outer leaflet was deposited by Langmuir-Schaefer (LS) method where the monolayer coated quartz was stamped horizontally through the lipid monolayer on the air-water interface to form the symmetric bilayer. The isotherm for 92:8 DPPC-cholesterol is shown in Figure 1.

Figure 1. Isotherm of 92:8 DPPC-cholesterol at 25°C. (Inset) FM image of the 92:8 DPPC-cholesterol membrane at 40x magnification with a photobleached spot. The diffusivity of the DPPC:cholesterol mixtures was very low (almost no recovery observed after 3 hours).

Vesicle solutions for zeta potential measurements were prepared using the extrusion method. 92:8 DPPC-cholesterol mixtures were prepared in chloroform, dried under nitrogen and then placed under vacuum for at least 4 hours. The dried lipids were hydrated with 0.5mM NaNO₃ solution prepared

in MilliQ-water to a concentration of 0.4 mg/mL, heated to a temperature of 50°C, and then extruded through a 100 nm pore size polycarbonate membrane for 10 passes while maintaining the temperature around 50°C. After the extrusion process, the vesicle solution was placed in a warm water bath (~50°C) and then cooled down slowly to 25°C by placing the bath with the vesicle solution in a temperature controlled room at 25°C.

Surface Force Measurements (SFA). The SFA technique has been used extensively to measure the interaction forces between surfaces and details of the technique can be found in the following references^{22, 23, 24}. Based on multiple-beam interferometry (MBI), the SFA provides a definitive reference for the surface separation (± 0.2 nm in this work). Briefly, one of the membrane coated mica surfaces was mounted on a fixed stage and the other on a double cantilever spring of known stiffness ($\sim 2.6 \times 10^5$ mN/m) which can be displaced vertically. The back of the mica substrates was coated with a 55 nm thick, evaporated silver layer. The silver layer on each disk partially transmits light directed normally through the surfaces which constructively interferes producing fringes of equal chromatic order (FECO). The distances between the surfaces can be measured by observation of the position and displacement of FECO peak wavelengths within a spectrometer. A custom automated SFA Mark-II was used for data collection. The system enables constant and/or variable surface displacements via a computer-controlled motor system. A sensitive CCD camera (Princeton SPEC-10:2K Roper Scientific, Trenton, NJ) was interfaced with the spectrometer and computer acquisition system to allow automated FECO wavelength determination, which was used to determine the separation distance.

After lipid bilayer deposition, the surfaces were transferred and mounted into the SFA under water. The water in the SFA was saturated with lipids to minimize desorption from the surface during the course of the measurements. After the surfaces were mounted, the SFA was placed in a temperature-controlled room at 25.0°C for at least two hours to allow equilibration. The membrane thickness was determined using the FECO wavelength shift from membrane contact relative to bare mica substrates after completing each experiment. As the membranes were asymmetric with inner leaflets of DPPE and outer leaflets of DPPC containing 8 mole% cholesterol, we treat the two outer leaflets, which we are primarily interested in, as an equivalent membrane of the mixture composition. Force profiles shown in the results section are from three independent experiments which represent the range of typical force profiles obtained from six completely independent SFA experiments.

Atomic Force Microscopy (AFM). AFM images were acquired using a MFP3D-SA system (Asylum Research, Santa Barbara, CA). A silicon nitride cantilever (model AC240, Bruker, Santa Barbara, CA) with force constant of 1 N/s was used for imaging. All the images were acquired in contact mode in MilliQ

gradient water with force of 37nN. AFM images were analyzed using Gwyddion Version 2.31 (<http://gwyddion.net/>).

Fluorescence Microscopy (FM). FM images were acquired using a Nikon Eclipse E600 microscope connected with CoolSNAP-Pro CCD camera at 10x and 40x magnification.

Zeta Potential Measurements (ZP). ZP was used to quantify the magnitude of the electrical charge based on measurements of vesicle mobility in an electric field in 0.5 mM NaNO₃ electrolyte solutions (Brookhaven Zeta Plus, Holtsville, NY). Although not identical with the surface charge or surface potential determination from force profile measurements by SFA, ZP provides the sign of the electrical charge and its relative magnitude referenced between the hydrodynamic shear plane at the vesicle surface and the bulk solution. ZP results were obtained from six independent samples with ten measurements per sample.

Neutron Reflectivity (NR). Reflectivity, R , is defined as the ratio of the number of neutrons elastically and specularly scattered from a surface to that of the incident beam. When measured as a

function of wave-vector transfer, Q_z , where θ is the angle of incidence and λ

$$Q_z = |k_{out} - k_{in}| = \frac{4\pi \sin\theta}{\lambda}$$

is the wavelength of the beam, the reflectivity curve contains information regarding the sample-normal profile of the in-plane averaged scattering length density (SLD) and is therefore most suited for studies of interfacial, layered films. From the measured reflectivity profile, the thickness, SLD, and roughness of the membrane normal to the substrate can be determined by minimizing the difference between the measured reflectivity and that obtained from a modeled SLD profile of the membrane²⁵.

NR experiments were performed on the SPEAR beamline at the Manuel Lujan Neutron Scattering Center (Los Alamos National Laboratory). Neutron reflectivities down to $R \approx 1 \times 10^{-6}$, and momentum transfers out to $Q_z = \sim 0.2 \text{ \AA}^{-1}$ were obtained. The uncertainty of the Q_z resolution, σ_{Q_z}/Q_z , including instrumental resolution was approximately 3% for the entire range of scattering vectors. NR data was modeled using Motofit (Rev. 409), a module in IgorPro 6.31. All measurements were done in D₂O at room temperature which varied between 24-28°C, well below the melting point of the membrane. Three independent experiments were done and about the same membrane thickness was obtained from each of the experiments. The NR profile shown in the results section is the data set from the most complete (highest quality) membrane.

RESULTS AND DISCUSSIONS

Force - distance profiles of 92:8 DPPC:Cholesterol membranes. Figure 2 shows the measured force-distance profile between opposing 92:8 DPPC-cholesterol membranes in 0.5mM NaNO₃. The three different profiles capture the variation in measurements from six independent experiments. The average thickness obtained across all the measurements for an equivalent 92:8 DPPC-cholesterol bilayer was 5.9±0.3nm.

Figure 2. Force-distance profiles between 92:8 DPPC-cholesterol membranes in 0.5mM NaNO₃ solution and fit of the electrostatic contribution (solid line) with the origin of charge at the membrane interface. $D = 0$ is defined as contact between the mica substrates (open symbols: approach; filled symbols: separation). . (Inset) The same force profiles after subtracting the electrostatic contribution to quantify the magnitude of the attractive adhesion between the membranes.

A weak, long-range electrostatic repulsion was apparent in the force profiles with a decay length consistent with the electrolyte concentration. The electrostatic repulsion was fit using the Poisson-Boltzmann equation yielding a surface charge density of 1.0 ± 0.5 mC/m² or surface potential of 19 ± 3 mV

with the origin of charge at the membrane surface.²⁶ Although the ionization constant of cholesterol and PC lipids^{27,28} would suggest an overall neutral charge for the membranes, the presence of charged lipid impurities has been previously reported.^{20,29} The measured surface charge corresponds to 1 negative charge per 400 lipids, or 0.25mole% of impurities present in the mixture. In order to corroborate and further quantify the electrostatic repulsion in the 92:8 DPPC-cholesterol system, the zeta potential of vesicle suspensions was measured. The zeta potential of 92:8 DPPC-cholesterol vesicles was -9.8 ± 3.8 mV,³⁰ which is in good agreement with the electrostatic charge measured in the SFA and our previous zeta potential measurements of similar lipid compositions²⁰. The zeta potential and low incidence of membrane spanning defects observed from AFM scans, as described later, are consistent with the source of the electrostatic repulsion being due to the presence of lipid impurities with the origin of charge at the membrane surface.

To better quantify the adhesion between the membranes, the electrostatic repulsion was subtracted from the force profile as shown in the inset of Figure 2. The adhesion between the 92:8 DPPC-cholesterol membranes varied from -1.1 ± 0.1 to -4.7 ± 0.1 mN/m and was reproducible for a given contact region between the membranes. The strength of adhesion at the low end is comparable to pure gel phase DPPC membranes.^{31,32} The higher adhesion values of some membrane contact positions are attributed to hydrophobic attraction due to membrane defects. Specifically, AFM scans of the 92:8 DPPC-cholesterol membrane (Figure 3) revealed nanoscopic monolayer defects in the outer 92:8 monolayer that extend down to the inner DPPE monolayer. The size range of the defects was remarkably uniform, diameter = 230 ± 30 nm. AFM scans of cholesterol devoid membranes (a pure DPPC leaflet on an inner DPPE leaflet) showed considerably fewer defects strongly suggesting that the incorporation of cholesterol correlates with more defects (see supplemental information).³³ These defects expose hydrophobic moieties and a commensurate increase in the measured adhesion between the two membranes due to hydrophobic attraction. The variability in the adhesion is attributed to inhomogeneities in the membrane defect density, which ranged from 1 to 5% surface coverage³⁴. In one case, a membrane spanning hole that reached the mica substrate was detected by AFM (Figure 3B). The height scan thickness of the membrane was ~ 6 nm, which is in good agreement with the membrane thickness obtained from SFA measurements.

Figure 3. (A) 10 $\mu\text{m} \times 10 \mu\text{m}$ and (Inset) 5 $\mu\text{m} \times 5 \mu\text{m}$ AFM topography images of the 92:8 DPPC-cholesterol system. (B) and (C) Corresponding cursor profiles as indicated in (A) and inset of image (A) respectively to obtain height profiles. (D) and (E) Histogram of defect depth and width across the sample respectively.

In all cases, membrane restructuring or thinning of the membrane was negligible during the course of an experiment; even under increased loads at contact. Reproducible force profiles and adhesive minima for a given contact position strongly suggests that the system does not rearrange during the time scale of the experiment. This observed trend is consistent with the small size of the defects and the gel phase behavior of the membrane (see Figure 1). Hemifusion was not observed in any of the experiments due to the low diffusion and relatively low applied force during the course of the experiments. Conversely, fluid phase membranes containing defects have been shown to restructure in contact.²⁰

Neutron Scattering Studies. The structure of supported 92:8 DPPC-cholesterol membranes on quartz was also characterized using NR and Figure 4 shows the best data set from three independent experiments. The thickness of the membrane was consistent between the three measurements, but the SLD of the hydrocarbon region was somewhat higher for the other two cases. The hydrocarbon region SLD is indicative of membrane defects as the presence of D₂O in the defects increases the SLD. Due to the resolution of the NR data, which was primarily sensitive to the hydrocarbon region (acyl chains) of the bilayer, a simple 1-layer model of was used to fit the data. The thickness of the box was 4.15 ± 0.10 nm with an SLD of $-0.3 \pm 0.2 \times 10^{-6} \text{ \AA}^{-2}$. This value is in agreement with the SLD of pure PC hydrocarbon tails of $-0.41 \times 10^{-6} \text{ \AA}^{-2}$ by Fragneto *et al.*³⁵ given that the neutron SLD of cholesterol is $0.21 \times 10^{-6} \text{ \AA}^{-2}$ and a defect density less than 5%.^{36, 37} The other samples had somewhat higher hydrocarbon SLDs indicative of higher defect densities. The uniform thickness of the hydrocarbon region for all the samples is also consistent with nanoscopic rather than macroscopic defects.

Figure 4. (Top image) Fit of RQ^4 vs Q from the neutron reflectivity data. (Bottom image) Schematic of the supported membrane and corresponding neutron scattering length density profile of the membrane obtained from fitting the data.

CONCLUSIONS

As one of the major components in biological membranes, the functions and properties of cholesterol have been widely studied and documented. In this work, a simplified system of a substrate

supported membrane was used to measure the interaction forces between two cholesterol containing membranes and the structure of the membranes with ultra-high resolution. The interaction force-distance profiles between DPPC-cholesterol membranes measured using SFA and AFM imaging of the membrane revealed the presence of nanoscopic defects leading to an enhanced hydrophobic attraction between the membranes in contact. The non-homogeneity in the lateral defect distribution across the membrane, as demonstrated by AFM and neutron scattering studies, leads to variation in the thickness of the membrane and membrane-membrane adhesion. A weak electrostatic repulsion due to the presences of charged lipid impurities was detected and further quantified by vesicle zeta potential measurements. Such studies demonstrate that cholesterol alters the intersurface interaction and membrane structure in complex and important ways.

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REFERENCES

1. Drolle, E.; Kucerka, N.; Hoopes, M. I.; Choi, Y.; Katsaras, J.; Karttunen, M.; Leonenko, Z. Effect of melatonin and cholesterol on the structure of DOPC and DPPC membranes. *Bba-Biomembranes* **2013**, 1828 (9), 2247-2254.
2. Rawicz, W.; Smith, B. A.; McIntosh, T. J.; Simon, S. A.; Evans, E. Elasticity, strength, and water permeability of bilayers that contain raft microdomain-forming lipids. *Biophys J* **2008**, 94 (12), 4725-4736.
3. Jurak, M. Thermodynamic Aspects of Cholesterol Effect on Properties of Phospholipid Monolayers: Langmuir and Langmuir-Blodgett Monolayer Study. *J Phys Chem B* **2013**, 117 (13), 3496-3502.
4. Almeida, P. F. F. Thermodynamics of lipid interactions in complex bilayers. *Bba-Biomembranes* **2009**, 1788 (1), 72-85.

5. Needham, D.; Nunn, R. S. Elastic-Deformation and Failure of Lipid Bilayer-Membranes Containing Cholesterol. *Biophys J* **1990**, *58* (4), 997-1009.
6. Dietrich, C.; Bagatolli, L. A.; Volovyk, Z. N.; Thompson, N. L.; Levi, M.; Jacobson, K.; Gratton, E. Lipid rafts reconstituted in model membranes. *Biophys J* **2001**, *80* (3), 1417-1428.
7. Silvius, J. R. Role of cholesterol in lipid raft formation: lessons from lipid model systems. *Bba-Biomembranes* **2003**, *1610* (2), 174-183.
8. Crane, J. M.; Tamm, L. K. Role of cholesterol in the formation and nature of lipid rafts in planar and spherical model membranes. *Biophys J* **2004**, *86* (5), 2965-2979.
9. Cinek, T.; Horejsi, V. The Nature of Large Noncovalent Complexes Containing Glycosyl-Phosphatidylinositol-Anchored Membrane-Glycoproteins and Protein Tyrosine Kinases. *J Immunol* **1992**, *149* (7), 2262-2270.
10. Ahmed, S. N.; Brown, D. A.; London, E. On the origin of sphingolipid/cholesterol-rich detergent-insoluble cell membranes: Physiological concentrations of cholesterol and sphingolipid induce formation of a detergent-insoluble, liquid-ordered lipid phase in model membranes. *Biochemistry-Us* **1997**, *36* (36), 10944-10953.
11. Smaby, J. M.; Brockman, H. L.; Brown, R. E. Cholesterols Interfacial Interactions with Sphingomyelins and Phosphatidylcholines - Hydrocarbon Chain Structure Determines the Magnitude of Condensation. *Biochemistry-Us* **1994**, *33* (31), 9135-9142.
12. Wydro, P.; Knapczyk, S.; Lapczynska, M. Variations in the Condensing Effect of Cholesterol on Saturated versus Unsaturated Phosphatidylcholines at Low and High Sterol Concentration. *Langmuir* **2011**, *27* (9), 5433-5444.
13. Fliesler, S. J.; Schroepfer, G. J. Sterol Composition of Bovine Retinal Rod Outer Segment Membranes and Whole Retinas. *Biochim Biophys Acta* **1982**, *711* (1), 138-148.
14. Bloch, K. Cholesterol: evolution of structure and function. In *Biochemistry of lipids, lipoproteins, and membranes*, Vance, D. E.; Vance, J. E., Eds.; Elsevier: Amsterdam ; New York, 1991, pp 363-381.
15. Anderson, T. G.; McConnell, H. M. Condensed complexes and the calorimetry of cholesterol-phospholipid bilayers. *Biophys J* **2001**, *81* (5), 2774-2785.
16. Brown, D. A.; London, E. Structure and function of sphingolipid- and cholesterol-rich membrane rafts. *J Biol Chem* **2000**, *275* (23), 17221-17224.
17. Marsh, D. Cholesterol-induced fluid membrane domains: A compendium of lipid-raft ternary phase diagrams. *Bba-Biomembranes* **2009**, *1788* (10), 2114-2123.
18. Marsh, D. Liquid-ordered phases induced by cholesterol: A compendium of binary phase diagrams. *Bba-Biomembranes* **2010**, *1798* (3), 688-699.
19. Mills, T. T.; Huang, J. Y.; Feigenson, G. W.; Nagle, J. F. Effects of cholesterol and unsaturated DOPC lipid on chain packing of saturated gel-phase DPPC bilayers. *Gen Physiol Biophys* **2009**, *28* (2), 126-139.
20. Kurniawan, J.; Yin, N. N.; Liu, G. Y.; Kuhl, T. L. Interaction Forces between Ternary Lipid Bilayers Containing Cholesterol. *Langmuir* **2014**, *30* (17), 4997-5004.
21. Kienle, D. F.; de Souza, J. V.; Watkins, E. B.; Kuhl, T. L. Thickness and refractive index of DPPC and DPPE monolayers by multiple-beam interferometry. *Anal Bioanal Chem* **2014**, *406* (19), 4725-4733.
22. Israelachvili, J. N.; Adams, G. E. Measurement of Forces between 2 Mica Surfaces in Aqueous-Electrolyte Solutions in Range 0-100 Nm. *J Chem Soc Farad T 1* **1978**, *74*, 975-&.
23. Israelac.Jn. Thin-Film Studies Using Multiple-Beam Interferometry. *J Colloid Interf Sci* **1973**, *44* (2), 259-272.
24. Kuhl, T. L.; Leckband, D. E.; Lasic, D. D.; Israelachvili, J. N. Modulation of Interaction Forces between Bilayers Exposing Short-Chained Ethylene-Oxide Headgroups. *Biophys J* **1994**, *66* (5), 1479-1488.
25. Kjaer, K. Some Simple Ideas on X-Ray Reflection and Grazing-Incidence Diffraction from Thin Surfactant Films. *Physica B* **1994**, *198* (1-3), 100-109.

26. Grabbe, A.; Horn, R. G. Double-Layer and Hydration Forces Measured between Silica Sheets Subjected to Various Surface Treatments. *J Colloid Interf Sci* **1993**, *157* (2), 375-383.
27. Tsui, F. C.; Ojcius, D. M.; Hubbell, W. L. The Intrinsic P_{ka} Values for Phosphatidylserine and Phosphatidylethanolamine in Phosphatidylcholine Host Bilayers. *Biophys J* **1986**, *49* (2), 459-468.
28. Moncelli, M. R.; Becucci, L.; Guidelli, R. The Intrinsic P_{k(a)} Values for Phosphatidylcholine, Phosphatidylethanolamine, and Phosphatidylserine in Monolayers Deposited on Mercury-Electrodes. *Biophys J* **1994**, *66* (6), 1969-1980.
29. Pincet, F.; Cribier, S.; Perez, E. Bilayers of neutral lipids bear a small but significant charge. *Eur Phys J B* **1999**, *11* (1), 127-130.
30. A positive shift in the measured ZP value was observed over time in some of the measurements. Typically, such shifts are associated with changes in the vesicle suspension with the application of an electric field, consistent with gel phase vesicle instability and fusion. Therefore, the reported value of Ψ for the 92:8 DPPC-cholesterol vesicles was the average of all measurements before the positive shift was observed.
31. Marra, J.; Israelachvili, J. Direct Measurements of Forces between Phosphatidylcholine and Phosphatidylethanolamine Bilayers in Aqueous-Electrolyte Solutions. *Biochemistry-Us* **1985**, *24* (17), 4608-4618.
32. Orozco-Alcaraz, R.; Kuhl, T. L. Interaction Forces between DPPC Bilayers on Glass. *Langmuir* **2013**, *29* (1), 337-343.
33. Our measurements of six DPPC on DPPE system were consistent with earlier studies by Marra and Israelachvili yielding an adhesion of -0.9 ± 0.2 mN/m.
34. The defect density was calculated from more than 60 10x10 μ m AFM scans. The analysis scan size was chosen to be about the same size as the membrane contact region in the SFA.
35. Fragneto, G.; Graner, F.; Charitat, T.; Dubos, P.; Bellet-Amalric, E. Interaction of the third helix of Antennapedia homeodomain with a deposited phospholipid bilayer: A neutron reflectivity structural study. *Langmuir* **2000**, *16* (10), 4581-4588.
36. Greenwood, A. I.; Tristram-Nagle, S.; Nagle, J. F. Partial molecular volumes of lipids and cholesterol. *Chem Phys Lipids* **2006**, *143* (1-2), 1-10.
37. The SLD of cholesterol was calculated using the partial molar volume of cholesterol embedded in a DPPC membrane assuming that cholesterol was oriented normal to the bilayer plane. The subphase D₂O has large SLD contrast to the acyl chain of lipids which enable us to quantify the defect density. The SLD of the hydrocarbon region indicates a complete bilayer was deposited with less than $1 \pm 3\%$ defects.

TOC Figure