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The inverse hexagonal – inverse ribbon – lamellar gel phase transition sequence in dehydrated DOPC:DOPE phospholipid mixtures.

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The inverse hexagonal to inverse ribbon phase transition in a mixed phosphatidylcholine-phosphatidylethanolamine system at low hydration is studied using Small and Wide Angle X-ray scattering. It is found that the structural parameters of the inverse hexagonal phase are independent of temperature. By contrast the length of each ribbon of the inverse ribbon phase increases continuously with decreasing temperature over a range of 50 °C. At low temperatures the inverse ribbon phase is observed to have a transition to a gel lamellar phase, with no intermediate fluid lamellar phase. This phase transition is confirmed by Differential Scanning Calorimetry. The biological relevance of these findings to dehydration and freezing damage are discussed.

The transitions of naturally occurring phospholipids from the biologically stable bilayer phase to various non-lamellar phases, has been the subject of considerable interest for decades eg [1–7]. Such phases are known to occur in membrane systems with low water contents, and thus are particularly important in understanding dehydration and freezing damage in biological organisms eg [8–13]. In addition to their importance at low hydrations, these phases may be involved in cell fusion [7, 14–16].

The inverse hexagonal phase (H_{II}), which consists of thin channels of water surrounded by lipid head groups, arranged on a 2D hexagonal lattice, has been widely studied [1–7]. An equally important phase which has received less attention is the inverse ribbon phase (P_{α}), which consists of ribbon-like strips of bilayer packed on a two-dimensional rectangular lattice. A schematic of both phases is shown in Fig. 1. The importance of such phases in biology is that they disrupt the cell membrane, a primary function of which is semipermeability - the ability to allow water to pass through to maintain osmotic balance with its surroundings, but provide a barrier to the passage of almost everything else. When the concentration of solutes around a cell increases, a cell responds by expelling water to re-establish osmotic balance. Loss of semipermeability - when a cell loses the ability to respond to osmotic changes - is a primary cause of damage in cells which have been dehydrated or frozen [8, 17], and has been correlated with transitions of membrane lipids to non-bilayer phases, which are no longer capable of keeping the inside and outside of the cell separated.

Despite their potential importance, inverse ribbon phases have previously been studied in only a small number of very low hydration systems of soaps eg [2] and some phospholipids [3, 18–23]. The data that does exist on this phase is limited in phospholipid systems, partly due to poor resolution of higher order Bragg reflections. At present there is no systematic study of the temperature dependence of the structural parameters, or the kinetics of the transition from the H_{II} phase.

Figure 1: (Color online) Schematic of the H_{II} and P_{α} phases. The structural parameters of the unit cells for both phases are shown. In the ribbon phase the headgroups do not extend to the end of the ribbons, but rather that the space between ribbons is filled with hydrocarbon, represented by the hydrocarbon distance d_h .

In this paper we use synchrotron small and wide angle x-ray scattering (SAXS/WAXS) to study the phase behaviour of a low hydration mixture of Dioleoylphosphatidylcholine (DOPC) and Dioleoylphosphatidylethanolamine (DOPE), representative of the most widely occurring phospholipids in nature. We observe an inverse fluid ribbon phase (P_{α}) which is stable over a wider range of temperatures, and at higher hydrations, than has previously been observed. The system undergoes a transition from P_{α} to H_{II} over the range

40-45 °C. Upon cooling, the P_α phase transforms into the lamellar gel (L_β) phase just below 0 °C, without an intermediate fluid lamellar (L_α) phase. We characterize the structural changes occurring during the H_{II} to P_α transition, and determine the temperature dependence of the structural parameters of the P_α phase over a wide temperature range. The high flux of 3rd generation synchrotrons allows for short acquisition times with high signal to noise, enabling studies of the kinetics of the transition, with some hysteresis observed.

DOPC and DOPE were obtained from Avanti Polar lipids and used without further purification. The lipids were measured by weight to give the molar ratio DOPC:DOPE 2:1. Samples were hydrated with excess water and vortex mixed while cycling over a range of temperatures to ensure sample homogeneity. Once mixed, the samples were equilibrated over saturated salts at a relative humidity (RH) of 32% for two weeks at room temperature. Water content of the sample (4.3 wt%, or 1.9 water molecules per lipid) was determined gravimetrically following experiments by drying samples in DSC pans with pin holes over P_2O_5 in a vacuum oven at 80 °C.

Synchrotron SAXS/WAXS experiments were carried out on the ChemMatCARS 15ID-D beamline at the Advanced Photon Source (APS), Argonne National Laboratory. Diffraction patterns were recorded on a Bruker 6000 CCD detector over the q -range 0.046 to 1.7 Å⁻¹, with the lower q -range (SAXS) probing the primary repeats of the inter water core distance for the H_{II} phase and the bilayer repeat spacing for the lamellar and ribbon phases. The higher q -range (WAXS) probes the intra-lipid spacing which indicates lipid fluidity. Samples were placed in 1.5 mm quartz X-ray capillaries (Wolfgang Muller Glas Technik, Berlin) and sealed using Araldite epoxy resin. Temperature control was achieved using a circulating water chiller/heater. Equilibrium measurements (in the range 20 °C - 80 °C) were made after allowing the sample temperature to equilibrate for at least 5 min. Kinetic studies were carried out by equilibrating at 20 °C then temperature jumping to 80 °C (rate ~20 °C/min.), equilibrating for 60 seconds, then temperature jumping back down to 20 °C. Exposures were made at 10 second intervals, and all exposures were 1.2 seconds in duration.

Further equilibrium SAXS measurements below 20 °C were carried out on a Nanostar (Bruker, Germany) covering the q -range 0.035 to 0.85 Å⁻¹. Exposures were of 1 hour duration after equilibrating the sample for 30 min. at the measurement temperature (acquisition times are three orders of magnitude longer than the synchrotron experiments). Samples were mounted in Aluminium sample holders with windows of clear adhesive tape (3M). This instrument is not capable of measuring in the WAXS region, so Differential Scanning Calorimetry (DSC) measurements (Pyris 7 DSC with intracooler (Perkin-Elmer)) were made in order to aid in phase identification. Sam-

Figure 2: (Color online) SAXS scans during cooling from 80 °C (front) to 20 °C (back). Scans are displaced vertically for clarity. The scans were taken at 10 s intervals and are of 1.2 s duration. Cooling rate was approximately 20 °C/min. The inset shows the broad wide angle peak, indicative of fluid chains, which does not change significantly over this temperature range.

ples were scanned at 10 °C/min. and 20 °C/min., during both heating and cooling.

Fig. 2 shows synchrotron SAXS patterns taken during cooling from 80 °C to 20 °C at ~20 °C/min. Phases were assigned based on the characteristic spacing of higher order reflections. At 75 °C, 5 small angle reflections are evident, with the reflections indexing according to the inverse hexagonal phase H_{II} : $q_{hk} = \sqrt{(h^2 + hk + k^2)}$. The broad WAXS reflection centred at 1.36 Å⁻¹ (inset) gives the average intra-lipid spacing (ie the spacing between headgroups). The broadness and absence of distinct sharp peaks in this q region indicates the hydrocarbon chains of the lipid molecules have the high degree of disorder characteristic of a fluid phase.

As the sample is cooled, it undergoes a transition over the range 40-45 °C. At the lowest temperature of 20 °C, 12 major reflections are visible. These reflections index according to a rectangular 2D monoclinic lattice with a systematic absence of (h, k) peaks, $h + k = 2n + 1$ indicating centred symmetry: $q_{hk} = 2\pi\sqrt{\left(\frac{h}{a}\right)^2 + \left(\frac{k}{b}\right)^2}$ where the parameters a and b denote the short and long unit cell axes respectively, as shown in Fig. 1. This phase is identified as the inverse fluid ribbon phase, P_α , which consists of ribbon like strips of alternating layers of lipid bilayer and anti-bilayer with disordered fluid hydrocarbon chains (α conformation). The P_α phase has previously been observed in dehydrated DOPE systems at water contents below 0.6% [18, 20–22]. An inverse ribbon phase with ordered hydrocarbon chains (P_δ) has also been observed

in dehydrated DOPE [19, 23] as well as in PC systems [3]. To the knowledge of the authors, all observations of the ribbon phase in phospholipid systems have been in anhydrous or dehydrated systems with water contents at or below 0.6%, and most of the previous work has been limited by low resolution of the higher order reflections. The detailed nature of the $H_{II} - P_{\alpha}$ transition has not previously been systematically studied.

Fig. 3 shows the structural parameters of both phases as functions of temperature. Both kinetic data (Fig. 2) and equilibrium data are shown. Clearly, the temperature has a greater effect on the structure on the P_{α} phase than the H_{II} phase, with the H_{II} unit cell dimensions remaining essentially constant over the measured temperature range. On cooling to the $H_{II} - P_{\alpha}$ transition, there is an immediate transition into the P_{α} phase with the transition complete within two exposures during scanning (< 20 seconds). The P_{α} short axis, a , is initially equal to twice the H_{II} (1,1) planar distance, and remains essentially constant. The long axis, b , is equal to twice the H_{II} (1,0) planar distance at the transition (Fig. 1). Both the agreement in the structural parameters and the speed of the transition demonstrate that the $H_{II} - P_{\alpha}$ transition involves a distortion of the hexagonal packing of the headgroups into an elongated ribbon, which does not incur a large energy cost. Indeed, DSC measurements show no measurable peak at this temperature. These findings are consistent with the idea [21] that the transition is entropy rather than enthalpy driven.

However, upon cooling there is an extension of the length of the ribbon, with the cell length b following an approximately linear trend as a function of temperature. Both kinetic heating and cooling show the same trends, with some hysteresis, and the equilibration measurements bisect the hysteresis loop. Below 30 °C the ratio $\frac{b}{a} > 2$, allowing the (4,0) and (0,2) reflections of the P_{α} phase to be differentiated in the SAXS data.

The intra-lipid spacing (inset, Fig. 2) decreases from 4.6 Å to 4.5 Å in a roughly linearly fashion as the temperature is reduced, while the FWHM of the intra-lipid WAXS peak remains constant. There is no qualitative change in this peak, indicating there are no gel or crystalline phases in the system. This is in contrast to other reports which have reported an inverse ribbon phase in dehydrated lipid systems with the hydrocarbon chains in the δ conformation – stiff chains with rotational disorder in a two-dimensional square lattice arrangement.

At lower temperatures (< 20 °C - equilibrium measurements), there is a transition to a lamellar phase at -4 °C, identified by a change in the spacing of the bragg reflections to the ratios 1 : 2 : 3 : 4 (data not shown). Intra-lipid (WAXS) data is not available on this instrument, however, DSC heating scans showed a phase transition beginning at -2.4 °C, with an enthalpy of 27.5 kJ/mol, consistent with a transition from fluid to ordered chains, such as the L_{α} to L_{β} (fluid to gel) transition. In ad-

Figure 3: (Color online) Structural dimensions as a function of temperature. Filled symbols are the H_{II} phase, open symbols are the P_{α} phase. The temperature ranges over which the three phases are identified are indicated by horizontal lines. Kinetic runs are indicated by right and left pointing triangles for heating and cooling respectively. Squares are equilibrium measurements (both heating and cooling). Pluses indicate the lamellar gel phase. Measurements at temperatures ≤ 20 °C were carried out on a lab based instrument, the others are synchrotron measurements.

dition, the measured bilayer thickness is 10% larger in the lamellar phase than in the P_{α} phase, again indicative of a fluid-gel transition. The slight offset between the transition temperatures is due to the fact that the SAXS data is measured at equilibrium, whereas the DSC measurements are made during scanning at 10 °C/min.

Interestingly, there is no discontinuity in the cell area as a function of temperature over the phase transition – with the ribbon unit cell area equal to twice the H_{II} unit cell area at the transition. As this is the case, one would assume that at the $H_{II} - P_{\alpha}$ transition, the number of lipids around the H_{II} water core and the number of lipids in half a ribbon should be the same.

The number of lipids around the H_{II} core can be calculated from the water core radius and the repeat spacing. Using literature values for the partial specific volumes of DOPC [24, 25] and DOPE [4], and knowing the mass fractions, the volume fraction of water $\phi_W = \frac{V_W}{V_L + V_W}$ can be calculated. The partial specific volumes are assumed not to change appreciably unless there is a chain freezing/melting transition, such as the $L_{\alpha} - L_{\beta}$ transition [26]. The H_{II} water core radius is then related to the water volume fraction by $R_W = 2d_{11}\sqrt{\phi_W \frac{\sqrt{3}}{2\pi}}$ [4]. For the hexagonal phase at 75 °C this gives $R_W = 4.9$ Å. From this, and the d spacing, one can then calculate the distance from the edge of the water core to the edge of the hexagonal unit cell along the (1,1) and (1,0) axes, i.e. the minimum and maximum lengths of the lipids in

this system. In this case 18.2 Å and 21.7 Å respectively, with an average value of ~19.9 Å.

However, the WAXS peak derives from scattering at some average point along the polar headgroup of the lipid. The polar head group thickness has been estimated as ~8 Å for DOPE at similar hydrations in the H_{II} phase [27], and ~10 Å for phosphatidylcholines in the lamellar phase [28]. Therefore if we assume that the intra-lipid WAXS peak corresponds to halfway along the polar headgroup, this adds ~5 Å to the 4.8 Å water core radius, giving a circumference of ~61 Å. The intra-lipid d spacing at the transition is 4.54 Å, meaning ~14 lipids could fit around the circumference of the water core.

The number of lipids per ribbon unit cell can be determined by assuming that the horizontal hydrocarbon distance between the ribbons (d_h in Fig. 1) can be approximated as the average lipid length in the H_{II} phase (19.9 Å) minus the headgroup length (9 Å). Using the the unit cell length ($b = 84.6$ Å at the transition) and the intra-lipid spacing (4.54 Å), leads to an estimate of ~16 lipids per ribbon, which is consistent, within the assumptions, with the number of lipids around the H_{II} core. Future electron density studies might shed further light onto the nature of this transition.

In this paper we have demonstrated for the first time a P_α phase stable over a wide temperature range, with a direct transition between the P_α phase and gel lamellar phase (with no intermediate lamellar fluid phase). In other words, the P_α phase exists as an intermediate phase between the gel lamellar and H_{II} phases. Furthermore we have shown that the P_α phase can exist at a higher water content in mixed DOPC/DOPE mixtures than in single component systems of either constituent. Of course biological membranes are much more complicated, with many different lipid species, as well as proteins and other components. However, as previous work has shown, differential hydration of different membranes components phase can lead to phase separations [11, 12, 29], concentrating lipids with tendencies to form inverse phases. Similar effects are expected to occur as membranes are brought into close proximity during fusion.

The results presented here are important for our understanding of freezing and dehydration damage in biological membranes, as well as being potentially relevant to membrane fusion. The fact that this relatively simple mixture exhibits a stable P_α phase over such a wide temperature range is an indication that even simple two component model systems behave in ways very different from the single component lipid systems which have been most widely studied to date. This highlights the importance of studying more complex model membrane systems, in order to better understand biological membranes.

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