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Cholesterol Expels Ibuprofen from the Hydrophobic Membrane Core and Stabilizes Lamellar Phases in Lipid Membranes Containing Ibuprofen

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(Dated: April 17, 2015)

There is increasing evidence that common drugs, such as aspirin and ibuprofen, interact with lipid membranes. Ibuprofen is one of the most common over the counter drugs in the world, and is used for relief of pain and fever. It interacts with the cyclooxygenase pathway leading to inhibition of prostaglandin synthesis. From X-ray diffraction of highly oriented model membranes containing between 0 and 20 mol% ibuprofen, 20 mol% cholesterol, and dimyristoylphosphatidylcholine (DMPC), we present evidence for a non-specific interaction between ibuprofen and cholesterol in lipid bilayers. At a low ibuprofen concentrations of 2 mol%, three different populations of ibuprofen molecules were found: two in the lipid head group region and one in the hydrophobic membrane core. At higher ibuprofen concentrations of 10 and 20 mol%, the lamellar bilayer structure is disrupted and a lamellar to cubic phase transition was observed. In the presence of 20 mol% cholesterol, ibuprofen (at 5 mol%) was found to be expelled from the membrane core and reside solely in the head group region of the bilayers. 20 mol% Cholesterol was found to stabilize lamellar membrane structure and the formation of a cubic phase at 10 and 20 mol% ibuprofen was suppressed. The results demonstrate that ibuprofen interacts with lipid membranes and that the interaction is strongly dependent on the presence of cholesterol.

Keywords: Ibuprofen, Cholesterol, Lipid Bilayers, Molecular Structure, X-Ray Diffraction

1. INTRODUCTION

In addition to specific interactions with biochemical targets, many drugs and pharmaceuticals are known to interact with lipid membranes through non-specific molecular interactions.1,2 For example, physical interactions with lipid membranes can cause changes to the membrane’s fluidity, thickness, or area per lipid.3,4 As many biological processes, such as cell signalling and adhesion, are mediated by the membrane and membrane bound proteins, changes to membrane processes induced by drugs can lead to significant changes in their biological function.5–9

When assessing the impact of a foreign molecule (such as a drug) on membrane properties, the partitioning of the drug within the membrane is often crucial. As an example, the common analgesic aspirin has been shown to interact with the head group region of the lipid membrane leading to an increase in lipid fluidity.10,11 Aspirin was eventually shown to counteract cholesterol’s condensing effect and to redissolve cholesterol plaques in lipid bilayers at high cholesterol concentrations12,13, and also to inhibit formation of cholesterol rafts at physiological concentrations of cholesterol.14 In contrast, the co-surfactant hexanol partitions into the tail group region leading to profound changes in the membrane structure as it induces a lamellar to hexagonal phase transition.15 In particular, small molecules can change partitioning of peptides in membranes. Melatonin was shown to reduce the population of the membrane-embedded state of amyloid-β25–35, a peptide involved in plaque formation in Alzheimer’s disease.16

Ibuprofen is a non-steroidal anti-inflammatory drug
(NSAID) who’s primary effect is related to the inhibition of prostaglandin synthesis, leading to anti-inflammatory and pain killing properties\textsuperscript{17,18}. Ibuprofen is a non-selective inhibitor of the cyclooxygenase enzyme. However, there is evidence for an interaction of ibuprofen with lipid membranes. Several studies have reported that ibuprofen leads to an increase in area per lipid\textsuperscript{19} and membrane defects\textsuperscript{20}, as well as a decreased membrane bending modulus, \(\kappa\)\textsuperscript{21}.

Here, we determine the location of ibuprofen in saturated lipid bilayers at a concentration of 2 mol% and report experimental evidence for an indirect, non-specific interaction between ibuprofen and cholesterol in membranes containing 5 mol\% ibuprofen and 20 mol\% cholesterol. Through X-ray diffraction in multi-lamellar, oriented membranes, we locate the ibuprofen molecule in the head group region and the hydrophobic core of the bilayers and observe that the presence of cholesterol expels ibuprofen from the membrane core. Cholesterol was also found to stabilize membrane structure, as the formation of an inverse cubic phase at high concentrations of 10 and 20 mol\% ibuprofen was suppressed when 20 mol\% cholesterol was present.

2. RESULTS

Highly oriented, multi-lamellar membrane stacks were prepared on silicon wafers and the molecular structure was studied using high resolution X-ray diffraction imaging, as depicted in Figure 1. By using oriented membranes, the in-plane (\(q_1\)) and out-of-plane (\(q_z\)) structure was determined separately, but simultaneously. All membranes were incubated at 30°C in 100% humidity for 24 h before the measurements and scanned at a temperature of \(T=28^\circ\text{C}\) and 50% relative humidity (RH). Similar to protein crystallography, this dehydrated state suppresses thermal fluctuations, increases the number of higher order Bragg peaks and thereby enhances structural features, allowing for a high spatial resolution\textsuperscript{22}.

Figure 2 shows 2-dimensional reciprocal space maps for a subset of samples in this study. Measurements are taken for \(-0.3 \text{ Å}^{-1} < q_1 < 3 \text{ Å}^{-1}\) and \(0 \text{ Å}^{-1} < q_z < 1.1 \text{ Å}^{-1}\). Pure DMPC membranes are shown in Figure 2 (a). Some qualitative conclusions can be drawn from the scattering patterns. The observed scattering shows a number of well defined intensities along both, the out-of-plane (\(q_z\)) and in-plane (\(q_1\)) axis, indicative of lamellar bilayers with strong in-plane ordering.

The arrangement of the different molecular components in the plane of the membranes can be determined from the in-plane scattering along \(q_1\). As introduced by Katsaras and Raghunathan\textsuperscript{23,24}, different molecular components, such as lipid tails, lipid head groups and also ibuprofen and cholesterol molecules, can form molecular sub-lattices in the plane of the membrane leading to non-overlapping sets of Bragg peaks.

The 100\% DMPC sample in Figure 2 (a) shows a number of well defined intensities along both, the in-plane (\(q_1\)) and out-of-plane (\(q_z\)) axis, indicative of a 2-dimensional reciprocal space structure significantly, concentrations of more than 5 mol\% induce changes in the in-plane and out-of-plane pattern (parts (c) and (d)). Lamellar structure is stabilized in the presence of cholesterol (parts (e) and (f)).

![FIG. 2. Reciprocal space maps of selected samples: (a) pure DMPC bilayers; (b) DMPC+2 mol\% ibuprofen; (c) DMPC+5 mol\% ibuprofen; (d) DMPC+20 mol\% ibuprofen; (e) DMPC+20 mol\% cholesterol; (f) DMPC+20 mol\% cholesterol+20 mol\% ibuprofen. While a small concentration of ibuprofen of 2 mol\% in part (b) does not alter membrane structure significantly, concentrations of more than 5 mol\% induce changes in the in-plane and out-of-plane pattern (parts (c) and (d)). Lamellar structure is stabilized in the presence of cholesterol (parts (e) and (f)).](image-url)
study were measured at \( T = 28^\circ \text{C} \), however, de-hydrated to 50% RH to enhance structural features leading to a more closely packed gel structure.

The sample with 2 mol% ibuprofen in Figure 2 (b) shows a qualitatively similar pattern indicating that small amounts of ibuprofen do not lead to a significant change in membrane structure or topology. However, membranes prepared with 5 mol% and 20 mol% ibuprofen in Figures 2 (c) and (d) show a single in-plane feature at \( q_z = 1.5 \ \text{Å}^{-1} \). This peak is indicative of hexagonal packing of disordered lipid tails\(^{26}\). Additional reflections are observed along \( q_z \), indicative of a change in membrane topology from the lamellar phase. Samples prepared with 20 mol% cholesterol also show disordered in-plane profiles (Figures 2 (e) and (f)), however, a lamellar \( q_z \) pattern.

2.1. Electronic Properties of Ibuprofen

Ibuprofen is an overall hydrophobic drug consisting of a large, hydrophobic body consisting of an aromatic ring and a carbon tail, and a small, hydrophilic head, where the oxygen groups are located. Ibuprofen was found to have low partitioning into water and to locate in the lipid phase\(^{27}\), preferentially in the interfacial region of the bilayer\(^{28}\).

As electromagnetic waves, X-rays mainly interact with the electronic structure of molecules. Electron distributions describing the ibuprofen molecule were calculated using the solved crystal structure of ibuprofen\(^{29}\). The corresponding structure file is deposited in the Crystallography Open Database with reference number 2006278.

To take into account thermal motion of atoms and electrons, the position of each atom was modeled by a Gaussian distribution with a width (FWHM) of 1 Å (or, in the case of samples with cholesterol, 2 Å) and the corresponding electron distributions were then projected onto the \( z \)-axis. The molecule can be rotated to have any orientation with respect to the \( z \)-axis.

When the long axis of the molecule is not tilted with respect to the \( z \)-axis, three Gaussian distributions well describe the averaged profile, as shown in Figure 3 (a).

The first peak is assigned to the tail region of the ibuprofen, the second to the ring structure, and the third peak to the oxygenated head region. When the ibuprofen molecule is tilted between 30° and 60° two Gaussians are required, as depicted in Figure 3 (b)-(c). When the molecule is tilted 90°, only a single Gaussian is required, as depicted in Figure 3 (d). The electronic profiles in Figure 3 describe the molecule when the thermal motion of each atom is modeled by a Gaussian with FWHM of 1 Å.

The Gaussian distributions used to describe the ibuprofen profiles were then shifted and scaled to fit observed changes in membrane electron density with the inclusion of ibuprofen. The orientation and position of all membrane-embedded states can be determined in this fashion with high accuracy. This technique was used previously by Dies et al., who used the atomic structures of amyloid-\( \beta \) peptides and melatonin to determine the location of the peptides and enzyme in lipid membranes of different membrane compositions\(^{16,30}\).
FIG. 4. (a) Out-of-plane X-ray diffraction ($q || = 0 \text{ Å}^{-1}$) of oriented DMPC membranes containing ibuprofen at concentrations of 0 mol% (black), 2 mol% (blue), 5 mol% (green), and 20 mol% (red). (b) Peak indexing for a membrane with 5 mol% ibuprofen. Gaussian peaks were fit to describe the observed reflectivity curve. Peaks drawn in blue correspond to peaks, which scatter solely from cubic phases. Peaks in red agree with scattering from either a cubic phase or an epitaxially related lamellar phase. The inset (c) shows the position of the peaks along $q_z$ vs. assigned peak indices ($h^2 + k^2 + l^2)^{1/2}$ for a cubic phase. The quality of the peak assignments is shown by the perfectly linear behaviour.

2.2. The Interaction of Ibuprofen with DMPC Membranes

For a quantitative analysis, the 2-dimensional data in Figure 2 were cut along the $q_z$ direction. Out-of-plane diffraction for DMPC membranes prepared with ibuprofen concentrations from 0 mol% to 20 mol% is presented in Figure 4 (a). Up to twelve evenly spaced diffraction peaks were observed for pure DMPC bilayers, indicative of a well ordered lamellar structure. The measured lamellar spacing, $d_z$, for the pure DMPC was determined to be 55.1 Å, in agreement with previous reports. A similar pattern is observed for DMPC+2 mol% ibuprofen, which indicates that small amounts of ibuprofen do not change the structure of the bilayers significantly or alter the topology of the membranes. Additional peaks are observed at higher ibuprofen concentrations of 5 mol% and 20 mol% in Figure 4 (a). The structural changes associated with these reflections will be discussed below.

The location of the ibuprofen molecules in the saturated lipid bilayers can be determined by comparing the results for pure DMPC and DMPC+2 mol% ibuprofen.
membrane core, oriented parallel with the bilayer z-axis, with a tilt angle of 0±5°. In addition to changes in the tail regions, two additional increases in electron density were observed in the head group region. A single peak is observed at \( z=17 \) Å, best described by a bound ibuprofen molecule, rotated by 90°±11° with respect to the z-axis, at the interface of the head group and tail group regions. Two peaks at \( z=23 \) Å and \( z=27 \) Å are best described by a molecule which is distributed between the two bilayers and aligned with the z-axis (tilt of 0±5°). The peak at \( z=23 \) Å is described by the electron distribution of both the hydroxyl group and the terminal methyl groups of an ibuprofen molecule, suggesting both portions of the molecule are observed embedded in the head groups (of opposite bilayers). The peak at \( z=27 \) Å suggests the ring-group of ibuprofen observed between bilayers. A cartoon depicting the three membrane bound states for ibuprofen is shown in Figure 9 (a). By integrating the area under the peaks observed in the difference electron density curve, the relative occupation of each bound state can be determined. A relative occupation of 56% is observed for the upright state in the tails, 8% for the rotated state at the head-tail interface, and 36% for the state in the head groups.

Figure 4 (b) shows the out-of-plane diffraction pattern obtained from a membrane prepared with 5 mol% ibuprofen in a pure DMPC membrane. All observed peaks are fit with Gaussian peak profiles. The observed peaks cannot be indexed by a pure lamellar phase, however, may be indexed to a \( \text{Im}3m \) cubic structure with lattice parameter \( a = 134 \) Å. The corresponding cubic phase Miller indices are given on the Figure; however, select peaks are indexed by a lamellar phase with bilayer spacing of \( d_z = 55.1 \) Å. These peaks are indicated by red Gaussian profiles in Figure 4 (b). Peaks solely indexed by the cubic phase are described by blue profiles. Note that the spacing of the [211] plane of cubic phases is often observed to be epitaxially related to the bilayer spacing of the lamellar phase. The position, \( d \)-spacing, and Miller indices for all peaks extracted from Figure 4 (b) are listed in Table 1.

To determine the relation between cubic and lamellar phase and the orientation of both phases, 2-dimensional X-ray maps of the region of interest were obtained for membranes with 0 mol%, 10 mol%, and 20 mol% ibuprofen, and are displayed in Figure 6. The plots show the region \( 0 < q_z < 0.21 \) Å\(^{-1}\) and \( 0 < q_{||} < 0.21 \) Å\(^{-1}\) in more detail, as compared to the overview plots in Figure 2. The pure DMPC bilayers in Figure 6 (a) show the lamellar \([100]_L\) Bragg peak and two diffuse contributions: The lamellar diffuse scattering occurring in horizontal sheets is the result of bilayer undulation dynamics. Bilayers, which are not perfectly oriented parallel to the silicon substrate lead to a faint powder ring, labeled as “defect scattering”. The number of these defect bilayers is typically very small as evidenced by the logarithmic intensity plot.

In addition to the cubic peaks observed in the out-of-

FIG. 6. High resolution reciprocal space maps show the increase in powder scattering with increased ibuprofen. Bilayers were prepared with concentrations of: (a) 0 mol%; (b) 10 mol% and (c) 20 mol% ibuprofen. Only a lamellar peak is observed for pure DMPC. The observed diffuse scattering was attributed to lamellar diffuse scattering due to fluctuations and defect scattering as the result of a small fraction of bilayers not perfectly aligned on the substrate\(^3\). For a membrane with 10 mol% ibuprofen, the defect scattering significantly increased indicative of a large fraction of “misaligned” bilayers. A cubic pattern is observed at 20 mol% ibuprofen. Intensities are shown on a logarithmic scale.
plane curves in Figure 4, there is a drastic increase in the intensity of defect scattering with increasing ibuprofen content (DMPC+10 mol% ibuprofen is shown in Figure 6 (b)), indicating an increase of membranes, which have a random orientation with respect to the perpendicular $z$-axis. While cubic peaks were observed in the specular out-of-plane scans, no diffuse cubic signals are visible in the 2-dimensional data at this ibuprofen concentration, most likely because the volume fraction of the cubic phase is still small. The pattern at 10 mol% ibuprofen is indicative of a coexistence of lamellar and cubic phases.

A distinct cubic peak pattern is observed at 20 mol% ibuprofen in Figure 6 (c). The positions of the broad powder-rings agree with cubic peaks observed in refection measurements: the [211], [411], and [422] peaks are observed corresponding to a cubic phase (see Figure 4 (b)). The faint [110] and [220] peaks observed in out-of-plane curves could not be resolved from the more diffuse in-plane scattering.

### 2.3. The Interaction of Ibuprofen with Membranes Containing Cholesterol

Because ibuprofen molecules were found to partition in pure lipid bilayers, DMPC membranes with cholesterol concentrations of 20 mol% and between 0 mol% to 20 mol% ibuprofen were prepared to study a potential interaction between cholesterol and ibuprofen. Out-of-plane diffraction scans for DMPC+20 mol% cholesterol, DMPC+20 mol% cholesterol+5 mol% ibuprofen and DMPC+20 mol% cholesterol+20 mol% ibuprofen are plotted in Figure 7. The diffraction patterns could all be indexed by lamellar phases. Electron density profiles for 0 mol% and 5 mol% ibuprofen were used to determine the position of the ibuprofen molecule in cholesterol-containing DMPC membranes and are shown in Figure 8 (a). The curve containing ibuprofen in Figure 8 (b) was scaled to represent a DMPC+20 mol% cholesterol bilayer with 5 mol% ibuprofen embedded, as in Section 2.2. The difference between this scaled curve and the curve without ibuprofen was used to locate ibuprofen in membranes with cholesterol. Note that for membranes with cholesterol, when modelling the electronic distribution of ibuprofen for fitting to the difference curve each atom is modelled by a Gaussian distribution with a width of 2 Å, as opposed to 1 Å for membranes without cholesterol. The need for increased Gaussian blurring is most likely a consequence of increased molecular disorder with the addition of cholesterol to gel phase membranes.

Two Gaussian distributions, centred at $z=15$ Å and at 26 Å, were found to well describe the difference in electron density and were modelled as membrane embedded states for ibuprofen. The location of these states is in excellent agreement with the head group states observed in bilayers without ibuprofen in Section 2.2. The peak at $z=15$ Å is best described by an ibuprofen molecule tilted by $90^\circ\pm1$ relative to the bilayer normal, as depicted in the electron distribution calculations in Figure 3 (b). The peak at $z=26$ Å describes an ibuprofen molecule oriented parallel with the $z$-axis, and embedded between bilayers (tilt of $0^\circ\pm1$), similar to the bilayer spanning state observed in membranes without cholesterol. Note that the two membrane electron density profiles in Figure 8 (b) coincide in the tail group region, suggesting that ibuprofen does not occupy this region in the presence of cholesterol. By comparing the integrated intensity of the corresponding Gaussian peaks, the relative occupations of the states

<table>
<thead>
<tr>
<th>$q_z$-position ($\text{Å}^{-1}$)</th>
<th>$d$-spacing (Å)</th>
<th>Miller Index</th>
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<tr>
<td>0.077</td>
<td>88.3</td>
<td>[110]</td>
</tr>
<tr>
<td>0.117</td>
<td>53.7</td>
<td>[211]</td>
</tr>
<tr>
<td>0.131</td>
<td>48.1</td>
<td>[220]</td>
</tr>
<tr>
<td>0.201</td>
<td>31.2</td>
<td>[411]</td>
</tr>
<tr>
<td>0.231</td>
<td>27.2</td>
<td>[422]</td>
</tr>
<tr>
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<td>25.9</td>
<td>[101]</td>
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<td>23.4</td>
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<tr>
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<tr>
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<td>[18 9 9]</td>
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**FIG. 7.** Out-of-plane diffraction for bilayers prepared with 20 mol% cholesterol and ibuprofen concentrations of: (a) 0 mol%, (b) 5 mol%, (c) 20 mol%.
FIG. 8. Electron density profiles for DMPC membranes prepared with 20 mol% cholesterol (black) and 20 mol% cholesterol with 5 mol% ibuprofen (blue). The curves in (a) are on an absolute scale, while the ibuprofen containing curve in (b) was scaled to overlap the profile with that of a 20 mol% cholesterol-containing DMPC membrane (see details in text). The difference between the scaled curve and the black curve is best described by two Gaussian profiles, which are labelled in (b).

3. DISCUSSION AND CONCLUSIONS

Structural parameters, such as the lamellar spacing, \( d_z \), the cubic spacing and the area per lipid, \( A_L \), were determined for all samples and are listed in Table 2. Ibuprofen was found to not change \( d_z \) and \( A_L \) in gel DMPC bilayers for the concentrations used and within the resolution of this experiment. Addition of 20 mol% cholesterol led to an increase in lipid area and a decrease of lamellar spacing, as reported previously for gel phase bilayers\(^3\). The presence of 20 mol% cholesterol was found to suppress the formation of a cubic phases when up to 20 mol% ibuprofen was incorporated as well.

3.1. Partitioning of Ibuprofen in Saturated Lipid Membranes With and Without Cholesterol

The partitioning of ibuprofen in gel phase DMPC membranes was determined using a combination of X-ray diffraction and electronic structure calculations using crystallographic ibuprofen data. The result is summarized in Figure 9. While the peak amplitudes in the calculated profiles in part (a) appear to be systematically slightly smaller than the measured differences, peak position and peak widths show an excellent agreement. Based on the electronic properties in Section 2.1, the orientation of the ibuprofen molecules can be determined: while 3 peaks in the electron density difference indicate a parallel orientation, a single peak is consistent with a perpendicular, 90° orientation.

Three different membrane bound populations were observed when 2 mol% ibuprofen were added to the DMPC bilayers, as sketched in Figure 9 (a): ① a state in the hydrophobic membrane core, where the ibuprofen molecules align parallel to the lipid acyl chains; 56% of ibuprofen molecules were found in this state; ② 8% of ibuprofen molecules were observed at the interface between head groups-tail groups, and ③ 36% of ibuprofen molecules were found attached to the membrane head group region, situated between the lipid head groups of two bilayers. At ibuprofen concentrations greater than 5 mol% (10 and 20 mol%), disruption of the lamellar membrane phase and the formation of a cubic lyotropic phase was observed.

Based on a fit of the molecular electronic distribution of the ibuprofen molecule to the experimental data, as depicted in Figure 3, the ibuprofen molecules in the hydrophobic membrane core align parallel to the lipid tails, with their hydrophilic head groups located in the head group region of the bilayers (population ①). The 180° position, where the oxygen groups would locate in the bilayer centre, was found to be less favourable with a \( \chi^2 \) value of 6.33 \cdot 10^4, as compared to \( \chi^2 = 4.28 \cdot 10^4 \) for the 0° case. The locations of the ibuprofen molecules are consistent with previous studies, where ibuprofen was reported to associate with PC lipids\(^3,27\). Based on electrostatic considerations, the hydrophilic head of the ibuprofen is likely to locate in the head group region of the bilayers\(^19,28\). Population ③ corresponds to a state, where the ibuprofen molecule appears to be partially embedded in the head groups of two lipid bilayers, with the hydroxyl group in one bilayer and the terminal methyl groups in another. This membrane-spanning state of ibuprofen is likely a consequence of the stacked bilayers used for the diffraction experiments.

Only two populations of ibuprofen molecules were observed in the presence of cholesterol, as depicted in Figure 9 (b). 14% of the ibuprofen molecules were found to occupy a state at the head group-tail group interface (population ②), while 86% of the molecules were found in the space between two bilayers, attached to the head group region (population ③). While the two states are in agreement with states ② and ③ observed with no cholesterol, no membrane embedded ibuprofen state was observed in the cholesterol-containing lipid bilayers. The presence of cholesterol in the membrane seems to suppress partitioning of ibuprofen into the tail region. As ibuprofen and cholesterol molecules compete for the same space, cholesterol seems to have a higher affinity for the lipid acyl chains.

X-ray diffraction has been used previously to determine the position of a similar NSAID, aspirin, in DMPC membranes with and without cholesterol\(^10,12\). Aspirin was found to reside exclusively in the lipid head group region. However, there is a large hydrophobic component to the ibuprofen molecule, which would increase its affinity for the hydrophobic membrane core. Previous simulations of membrane systems incorporating ibuprofen locate the molecule in the tail group regions\(^21,32\). In addition, Langmuir isotherm experiments have also sug-
gested that ibuprofen may partition into the head group regions of lipid monolayers\textsuperscript{33}.

Our results agree qualitatively with other reports. Simulations by Khajeh \textit{et al.} report that the relative position of ibuprofen shifts towards the head groups in DMPC membranes containing 25 mol\% cholesterol\textsuperscript{32}. Additional studies have suggested that drug-membrane interactions are significantly influenced by the presence of cholesterol\textsuperscript{34,35}. Our experiments present experimental evidence that cholesterol influences the position of ibuprofen in the membrane and, as will be described below, also suppresses the cubic phase induced by ibuprofen.

3.2. The Suppression of Cubic Phases by Cholesterol

Inverse (type II) phases, such as inverse cubic or inverse-hexagonal phases, are frequently observed in amphiphile-water systems, including systems with lipids, surfactants, and block co-polymers\textsuperscript{36–38}. A lamellar to cubic phase transition may be induced in membranes by temperature or pressure jumps in systems containing lipids with negative curvature\textsuperscript{39,40}. Alternatively, inverse phases can be induced by the addition of a largely hydrophobic co-surfactant\textsuperscript{15,41,42}. The fingerprint of a lamellar to cubic phase transition is the appearance of Bragg peaks in diffraction experiments which require 3-fold symmetry to properly index\textsuperscript{43}.

Oriented membranes with ibuprofen concentrations less than 5 mol\% formed lamellar phases, while samples with concentrations greater than 5 mol\% could not be indexed by a single 1-dimensional lamellar phase and re-
quired a 3-dimensional cubic phase to index all peaks. The observed Bragg peaks for all samples in the cubic phase are consistent with either Im\(\text{3m}\) or \text{Pn3m}\) space groups, which are frequently observed in membrane systems\(^{36,43,44}\). The \([111]\) peak, which we did not observe, is systematically absent for \(\text{Im3m}\) but not \text{Pn3m}, suggesting \text{Im3m} is the best candidate. Another frequently observed cubic phase, with space group \text{Ia3d}, did not describe the peaks as the [110] reflection (absent for \text{Ia3d}) was observed\(^{44,45}\).

In membranes prepared on a solid substrate, where the lamellar phase is oriented, Bragg scattering from the membrane stack is observed solely along the out-of-plane axis, \(q_z\). However, the formation of 3-dimensional cubic phases leads to the appearance of off-specular scattering. Typically, cubic phases form as grains with random orientation, resulting in powder scattering, although oriented cubic phases have been prepared\(^{46}\). Two-dimensional measurements of reciprocal space were collected to observe off-specular scattering in the presence of ibuprofen and are depicted in Figure 6. The maps highlight the monotonic increase in powder scattering with increasing ibuprofen concentration. This suggests that increasing ibuprofen leads to cubic phases with grains at random orientation.

While specular peaks can unambiguously be indexed by cubic phases for ibuprofen concentrations greater than 5 mol\%, we note that a subset of those peaks may be indexed by a lamellar phase with bilayer spacing in close agreement with samples displaying a pure lamellar phase. The [211] plane of cubic phases is often observed to be epitaxially related to a bilayer spacing in systems with a lamellar to cubic transition\(^{19,44,47}\). Figure 4 (b) demonstrates how the observed peaks are indexed by either a cubic phase, or a cubic phase and a lamellar phase. The experiments, therefore, do not rule out the possibility of a lamellar phase coexisting with the cubic phase. From the 2-dimensional diffraction data in Figure 6 it seems that the formation of cubic phases is accompanied by a distortion of the lipid bilayers phase and the occurrence of bilayers not parallel to the \(z\)-axis.

There is evidence that the impact of certain drugs on the lipid membrane is dependent on membrane composition. For example, negatively charged lipids have been shown to accelerate the binding of the antimicrobial peptide Lacticin Q\(^{48}\). In addition, the anti-cancer drug Taxol has a different impact on saturated model membranes and unsaturated membranes\(^{49}\). In a recent paper by Khajeh et al., molecular dynamics (MD) simulations were performed on membranes with cholesterol and ibuprofen\(^{42}\) and report that the permeation of ibuprofen across the membrane is decreased by an increased stiffness of the membrane caused by cholesterol.

An increase in chain rigidity and decrease in permeability with the inclusion of cholesterol could explain the reduced penetration depth of Ibuprofen into the membrane\(^{32,50}\). In addition, a change in the position of ibuprofen could explain why cholesterol suppresses cubic phase formation. Cholesterol itself has not been shown to suppress cubic phases in membranes with inherently negative curvature\(^{31}\). However, by causing ibuprofen to partition in the head groups as opposed to the tail groups, cholesterol may prevent the negative curvature or decrease in bending modulus induced by ibuprofen. Our results demonstrate how a membrane constituent, such as cholesterol, can influence the membrane impact of a drug, such as ibuprofen, by changing the partitioning of the drug. Cholesterol can, therefore, act as a protective agent, by inhibiting cubic phases even when ibuprofen is present in high concentration.

### 4. MATERIALS AND METHODS

#### 4.1. Preparation of the Multi-Lamellar Membranes

Highly oriented, multi-lamellar membranes were prepared on polished 2 \(\times\) 2 cm\(^2\) silicon wafers. The wafers were first pre-treated by sonication in dichloromethane (DCM) at 310 K for 25 minutes to remove all organic contamination and create a hydrophobic substrate. After removal from the DCM post-sonication, each wafer was thoroughly rinsed three times by alternating with \(\sim 50\) mL of ultra pure water and methanol. 1,2-dimyristoyl-sn-glycero-3-phosphocholine (DMPC) and cholesterol were obtained from Avanti Polar Lipids and individually dissolved in 1:1 mixtures of chloroform and tri-fluoro-ethanol (TFE). Ibuprofen was also dissolved in a mixture of 1:1 chloroform and TFE. The DMPC, cholesterol and ibuprofen solutions were then mixed in the appropriate ratios to achieve the desired membrane compositions for the experiment. All samples prepared for this study are listed in Table 2. Molecular representations of the components are shown in Figure 1 (a).

A tilting incubator was heated to 313 K and the lipid solutions placed inside to equilibrate. 200 \(\mu\)L of lipid solution was deposited on each wafer and the solvent was then allowed to slowly evaporate for \(\sim 10\) minutes while being gently rocked, such that the lipid solution spread evenly on the wafers. After drying, the samples were

<table>
<thead>
<tr>
<th>DMPC (mol %)</th>
<th>Ibuprofen (mol %)</th>
<th>Cholesterol (mol %)</th>
<th>(d_z) (Å)</th>
<th>cubic spacing ((\AA^2))</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0</td>
<td>0</td>
<td>55.1</td>
<td>40.84</td>
</tr>
<tr>
<td>2</td>
<td>2</td>
<td>0</td>
<td>55.3</td>
<td>40.84</td>
</tr>
<tr>
<td>3</td>
<td>5</td>
<td>0</td>
<td>55.3</td>
<td>40.5</td>
</tr>
<tr>
<td>4</td>
<td>10</td>
<td>0</td>
<td>56</td>
<td>137</td>
</tr>
<tr>
<td>5</td>
<td>20</td>
<td>0</td>
<td>–</td>
<td>135.7</td>
</tr>
<tr>
<td>6</td>
<td>0</td>
<td>20</td>
<td>51.3</td>
<td>42.5</td>
</tr>
<tr>
<td>7</td>
<td>5</td>
<td>20</td>
<td>51.7</td>
<td>42.5</td>
</tr>
<tr>
<td>8</td>
<td>20</td>
<td>20</td>
<td>50.9</td>
<td>42.5</td>
</tr>
</tbody>
</table>

TABLE 2. Lamellar spacings and area per lipid are provided for all samples examined. For samples with cubic symmetry, the bilayer repeat distance was calculated using peaks which fit a lamellar spacing.
placed in vacuum at 313 K for 12 hours to remove all traces of solvent. Samples were then placed in a sealed container containing an open vial of pure water and allowed to equilibrate to 293 K. The temperature was then slowly increased to 303 K over a period of 24 hours. This procedure results in highly oriented, multi-lamellar membrane stacks an uniform coverage of the silicon substrates. About 3000 highly oriented stacked membranes with a total thickness of ~10 μm are produced using this protocol. The high sample quality and high degree of order is a prerequisite to determine in-plane and out-of-plane structure of the membranes separately, but simultaneously.

4.2. X-Ray Scattering Experiment

Out-of-plane and in-plane X-ray scattering data was obtained using the Biological Large Angle Diffraction Experiment (BLADE) in the Laboratory for Membrane and Protein Dynamics at McMaster University. BLADE uses a 9kW (45 kV, 200 mA) CuK-α Rigaku Smartlab rotating anode at a wavelength of 1.5418 Å. Both source and detector are mounted on moveable arms such that the membranes stay horizontal during measurements. Foucsing, multi layer optics provide a high intensity parallel beam with monochromatic X-ray intensities up to 10^{10} counts/(s × mm²). This beam geometry provides optimal illumination of the membrane samples to maximize the scattered signal. By using highly-oriented stacks, the in-plane (q_||) and out-of-plane (q_z) structure of the membranes could be determined independently. A sketch of the scattering geometry is depicted in Figure 1 (b). Full 2-dimensional reciprocal space maps are shown in Figure 2.

The X-ray scattering experiments determine three pieces of information relevant to molecular structure of the membranes. Firstly, out-of-plane diffraction scans allow for the identification of the phase of the membranes (lamellar or cubic) and also permit the reconstruction of electron density profiles (for lamellar samples). Electron density profiles were used to determine the position of the molecular constituents. Secondly, in-plane scattering measurements at high q_|| allow for the organization of the lipid molecules in the plane of the membrane to be determined. The area per lipid may be determined from the in-plane structure, as detailed in Barrett et al.\textsuperscript{10}. Thirdly, scans performed at low q_|| and low q_z can be used to measure the degree of orientation within the samples.

The 2-dimensional X-ray data in Figure 2 show well-defined peaks along the q_||-axis, which allow the determination of the lateral membrane structure. Several correlation peaks were observed in the in-plane data for ibuprofen concentrations of less than 2 mol%, and were well fit by Lorentzian peak profiles. The intensity has a distinct rod-like shape, typical for a 2-dimensional system. Membranes containing more than 2 mol% ibuprofen showed one broad Lorentzian peak, centered at ∼1.5 Å⁻¹ due to the organization of the lipid tails in the hydrophobic membrane core. The area per lipid molecule can be determined from the in-plane diffraction data, when assuming that the lipid tails form a densely packed structure with hexagonal symmetry (planar group p6\textsubscript{3}). As reported from, e.g., neutron diffraction,\textsuperscript{26} in the absence of fluctuations (in gel state lipid bilayers), the area per lipid can be determined from the position of the in-plane Bragg peak at q_T to \( A_L = 16\pi^2/\sqrt{3q_T^2} \)^{10,13,52}. The distance between two acyl tails is determined to be \( q_T = 4\pi/\sqrt{3q_T} \), with the area per lipid simplified to \( A_L = 3a_T^2 \), as listed in Table 2. The area per lipid for the pure DMPC and DMPC+2 mol% ibuprofen samples, which show a highly organized lateral membrane structure with additional in-plane Bragg peaks in Figure 2, were determined from the lattice parameters of the corresponding orthogonal tail lattice.

Structural parameters measured using the diffraction measurements, such as d_\textsubscript{z} spacing and A_L, for all samples are provided in Table 2.

4.3. Out-of-Plane Structure and Electron Densities

The out-of-plane structure of the membranes was determined using out-of-plane X-ray diffraction. The membrane electron density, \( \rho(z) \), is approximated by a 1-dimensional Fourier analysis:

\[
\rho(z) = \rho_W + \frac{F(0)}{d_z z} + \frac{2}{d_z} \sum_{n=1}^{N} F(q_n) \nu_n \cos(nz) \]

\[
= \rho_W + \frac{F(0)}{d_z z} + \frac{2}{d_z} \sum_{n=1}^{N} \sqrt{\nu_n} q_n \nu_n \cos\left(\frac{2\pi n z}{d_z}\right) \tag{1}
\]

\( N \) is the highest order of the Bragg peaks observed in the experiment and \( \rho_W \) is the electron density of bulk water. The integrated peak intensities, \( I_n \), are multiplied by \( q_n \) to generate the form factors, \( F(q_n) \). The bilayer

FIG. 10. Out-of-plane diffraction data for all samples for which Fourier analysis was performed. \( T(q_\nu) \), which is proportional to the membrane form factor, is shown in each inset and was used to determine the phases \( \nu_n \). (a) pure DMPC; (b) 2 mol% ibuprofen; (c) 20 mol% cholesterol; (d) 20 mol% cholesterol and 20 mol% ibuprofen.
form factor which is in general a complex quantity, is real-valued when the structure is centro-symmetric. The phase problem of crystallography, therefore, simplifies to the sign problem \( F(q_z) = \pm |F(q_z)| \) and the phases, \( \nu_n \), can only take the values \( \pm 1 \). The phases, \( \nu_n \), are needed to reconstruct the electron density profile from the scattering data following Eq. (1). When the membrane form factor \( F(q_z) \) is measured at several \( q_z \) values in a continuous fashion, \( T(q_z) \), which is proportional to \( F(q_z) \), can be fit to the data:

\[
T(q_z) = \sum_n \sqrt{n} q_n \sin\left(\frac{1}{2} d_z q_z - \pi n\right). \tag{2}
\]

In order to determine the phases quantitatively, the form factor has to be measured at different \( q_z \)-values using the so-called swelling technique or by measuring the bilayer at different contrast conditions when using neutron diffraction. In this paper, by fitting the experimental peak intensities and comparing them to the analytical expression for \( T(q_z) \) in Eq. (2), the phases, \( \nu_n \), could be assessed. Good agreement was obtained, and the results shown in Figure 10.

The calculated electron densities, \( \rho(z) \), which are initially on an arbitrary scale, were then transformed to an absolute scale. The curves were vertically shifted to fulfill the condition \( \rho(0) = 0.22 \, \text{e}^-/\AA^3 \) (the electron density of a CH\(_3\) group) in the centre of a bilayer. The curves were then scaled until the total number of electrons \( e^- = A_L \int_{0}^{d_z/2} \rho(z) \, dz \) across a membrane leaflet agrees with the total number of electrons expected based on the sample composition, with the addition of 7 water molecules, in agreement with\(^{16,53}\).

The \( d_z \)-spacing between two neighbouring membranes in the stack was determined from the distance between the Bragg reflections \( (d_z = 2\pi/\Delta q_z) \) along the out-of-plane axis, \( q_z \). Up to a peak order of 12 was observed from DMPC membranes, and up to 14 for DMPC membranes with cholesterol. Note that not all diffraction orders are necessarily observed for the different samples as the scattering intensity depends on the form factor of the bilayers.

**ACKNOWLEDGMENTS**

This research was funded by the Natural Sciences and Engineering Research Council (NSERC) of Canada, the National Research Council (NRC), the Canada Foundation for Innovation (CFI), and the Ontario Ministry of Economic Development and Innovation. R.J.A. is the recipient of an Ontario Graduate Scholarship, L.T. is the recipient of a Canada Graduate Scholarship, M.C.R. is the recipient of an Early Researcher Award from the Province of Ontario.

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Our experiments provide evidence for a non-specific interaction between ibuprofen and cholesterol in lipid membranes. Ibuprofen was found to reside in both the head group and tail group regions of the saturated DMPC bilayers. However, when cholesterol was incorporated in the membranes, ibuprofen was found to reside in the head group region, only. At the same time, cholesterol was found to stabilize the lamellar membrane phase by suppressing the transition into a cubic phase.