Lipid Landscapes: Vibrational Spectroscopy for Decoding Membrane Complexity

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Abstract
Cell membranes are incredibly complex environments containing hundreds of components. Despite substantial advances in the past decade, fundamental questions related to lipid-lipid interactions and heterogeneity persist. This review explores the complexity of lipid membranes, showcasing recent advances in vibrational spectroscopy to characterize the structure, dynamics, and interactions at the membrane interface. We include an overview of modern techniques such as surface-enhanced infrared spectroscopy (SEIRAS) as a steady-state technique with single-bilayer sensitivity, two-dimensional sum-frequency generation (2D SFG) spectroscopy, and two-dimensional infrared (2D IR) spectroscopy to measure time-evolving structures and dynamics with femtosecond time resolution. Furthermore, we discuss the potential of multiscale MD simulations, focusing on recently developed simulation algorithms which have emerged as a powerful approach to interpret complex spectra. We highlight persistent challenges in accurately sampling heterogeneous ensembles in multicomponent membranes. Overall, this review provides an up-to-
date comprehensive overview of the powerful combination of vibrational spectroscopy and simulations to illuminate lipid-lipid, lipid-protein, and lipid-water interactions in the intricate conformational landscape of cell membranes.

1. Introduction

A century of membrane models

Order and structure at every length scale is one of the hallmarks of biology. Lipid membranes provide molecular-scale barriers that compartmentalize a cell’s internal components as well as separate the delicate machinery of the cell from its surroundings. Their structure and physical, chemical, and mechanical properties, support the multiple roles of membranes, including as a physical barrier to support concentration gradients in energy production, as well as signaling, sensing, and transport (1–6). While the biochemical properties of membranes were relatively well established in early studies, structure and composition were difficult to characterize, mainly due to a lack of experimental techniques that could provide molecular-level structure, a task that remains challenging to this day. The original “bilayer” idea was first proposed in 1925, based on area-per-lipid measurements of compositions extracted from mammalian cells (7, 8). Later, in the mid 1930’s, Davson and Danielli proposed the first molecular model that incorporated proteins as part of the plasma membrane (9). The model was informed by early measurements of differential permeabilities, electron microscopy, thermodynamic property such as phase transition temperature, and, at the time new, X-ray investigation of protein crystals which showed dense cores solvated by water (10). Since proteins were known to be solvated by water at the time, a sandwich-like structure was proposed (Figure 1, left). In this model, a porous “protein film” would be located above and below the bilayer. While this model is too crude by current standards, it incorporates
some key elements that we associate with cell membranes. In addition, despite the crude models, the importance of hydrogen bonding and lipid-lipid interactions was recognized early on, and infrared spectroscopy, among other techniques, played an important role in supporting a dynamic fluid-like bilayer view that we still associate with cell membranes today (11, 12).

![Figure 1. Development of lipid models from 1930’s to 2020’s. The Davson and Danielli protein sandwich model (9), the Singer and Nicolson fluid-mosaic model (13), and the current view of cell membranes are illustrated from left to right. Individual components are indicated in the legend.](image)

The more influential “fluid-mosaic model” was developed in the 1970’s by Singer and Nicolson (Figure 1, middle) (13). Within this picture, membrane proteins are arranged in oriented amphipathic structures partially embedded within the phospholipid matrix, which is organized as a continuous “fluid” bilayer. This was based on X-ray crystallography, freeze-fracture electron microscopy, bulk thermodynamic measurements, and other biochemical studies. While this model has obvious limitations, the basic concept of transmembrane proteins embedded within the bulk bilayer is sufficient to explain many observations, and thus, the fluid-mosaic model remains
currently relevant as a qualitative description of the local membrane environment (14). Initial studies using infrared spectroscopy played, perhaps the most important role, in establishing that the lipids are characterized by liquid-like environments, despite having solid-like properties at the macroscopic scale (15, 16). These measurements informed the “fluid” character of the supporting lipids in the fluid-mosaic model, suggesting a dynamic bilayer environment.

Nanometer-scale order, compartmentalization, and heterogeneity make lipid membranes significantly richer in complexity compared to previous models (Figure 1, right). Modern models incorporate compositions containing a huge diversity of components, as observed in modern lipidomic analyses (17–19). One key aspect of modern lipid membrane models is the presence of compartmentalized domains with distinct lipid and protein compositions, referred to as “lipid rafts” that have been observed in artificial and biological membranes. There are clear implications to biological functions that result from colocalizing proteins, as well as modulating the local environment around certain proteins (20–22). Rafts have remained a topic of discussion for over a decade, and this stems mainly from the difficulty in experimentally characterizing their nanometer-scale structure and composition with current techniques. In addition, the concept of rafts rests on the idea of stable domains enriched in certain lipids. Experiments and simulations currently support a more nuanced picture than the lipid raft model, suggesting the formation of dynamic multicomponent local environments that results in transient interactions with specific lipid species, termed the “functional paralipidome” (22). While transient changes in local membrane composition can influence the protein conformational landscape, the molecular mechanisms connecting environments with protein modulation are not understood. While techniques such as cryo-EM have revolutionized the field of membrane structural biology, measuring lipid heterogeneity remains extremely difficult, due to a lack of techniques that provide species-specific
local environments. Beyond structural measures, there is virtually no time-resolved measurement that can serve to understand the coupling between protein and lipid dynamics across timescales (23, 24).

**What is missing from the current picture?**

While current membrane models include detailed descriptions, one fundamental aspect that remains underexplored is physical chemistry of membranes with biological compositions, which typically contain hundreds to thousands of lipid species when considering the diversity in headgroups and acyl chain structures. *Why so many lipids?* This question remains largely unresolved since biochemical approaches to addressing this question have provided mainly indirect information, and the complex interplay between local lipid composition and membrane protein topology as well as colocalization can be particularly complex (25, 26). The fundamental interactions are well understood: headgroup dipole-dipole interactions, hydrogen bonding between headgroups and with interfacial water, tail packing, and steric effects. Recent techniques and experiments, described below, have begun to shed light on the interplay between acyl chain composition and general trends in membrane protein topologies, but fundamental questions remain unaddressed. Specific questions include: how the balance of forces between lipid-lipid, lipid-protein, and lipid-water interactions, together with packing, dictates the local membrane environment in membranes containing thousands of lipids. Figure 2 shows a schematic representation of different lipid structures to emphasize on the diversity in headgroup structures and highlight hydrogen-bond donors and acceptors, and the steric size of the headgroup, as well as positive and negative charges associated with different functional groups. In brief, this depiction stresses the complex hydrogen bonding interactions present in multicomponent membranes. The main challenges to achieving a molecular-level understanding stem largely from the lack of well-
controlled model systems that incorporate lipid diversity and a range of local interactions, which can be combined with the dearth of experimental techniques to probe species-specific interactions, molecular conformations, and local environments in multicomponent bilayers.

Figure 2. Cartoon structures of common lipids in eukaryotic cells. The structures highlight the complexity in the headgroup structures in terms of charges as well as the number of hydrogen-bond donor and acceptor groups. The hydrophobic acyl chains are depicted in yellow, and the headgroup moieties are represented by colors and shapes based on the charge and their ability to donate and accept hydrogen bonds.
What is the outlook?

Developing quantitative models of intermolecular interactions within membrane components is crucial for establishing a general set of mechanistic principles that govern the coupling between environments and protein functions. Key questions that need to be addressed include the balance of forces stabilizing local lipidic environments around transmembrane proteins, the role of lipid-protein interactions in determining protein structure and stability, and the direct implications for protein function regulation. Formulating hypotheses remains challenging due to insufficient molecular information and the complexity of lipid-protein interactions. To advance beyond oversimplified membrane models requires new experimental approaches. Time-resolved and high-sensitivity vibrational spectroscopic methods discussed in this review show potential for providing molecular-level insights into these issues, revealing critical details about the structural and dynamic aspects of the interactions. In particular, the combination of time-resolved vibrational spectroscopy and molecular dynamics simulations is perfectly suited for investigating structure and dynamics at the interface over a range of timescales from picoseconds to milliseconds. These new approaches can help arrive at a comprehensive, molecular-level understanding of the interface, uncovering the impacts of interplay between lipid environments and proteins on associated functionality and biological processes. In the remainder of this review, we summarize the recent developments in ultrafast spectroscopies of model membranes, discuss the strengths and weaknesses of each approach, and provide some specific examples from recent research.

2. Vibrational Spectroscopy as a Probe of Membrane Structure and Heterogeneity

Vibrational spectroscopy is a powerful technique to measure conformational ensembles and local environments in complex membranes as vibrational frequencies are highly sensitive to molecular
structure, molecular interactions, and local environments. The main challenge in using vibrational spectroscopy lies in the interpretation of spectra within heterogeneous biological mixtures, as overlapping peaks produce ultra-congested lineshapes that obscure many features. As a result, at present model bilayers such as small unilamellar vesicles, which topologically resemble the most fundamental structures of cell membranes, are commonly used as model systems for membrane spectroscopy (27–29). Lipid molecules contain several intrinsic vibrational probes that report on the interfacial interactions and other local environments (Figure 3). The O-H stretch of water is a useful vibrational probe given its ability to probe the extended hydrogen-bond networks in water (30–32). Furthermore, both the phospholipid headgroups and alkyls can serve as sensitive vibrational probes reporting on specific regions of the bilayer. For example, the phosphate P-O stretching modes are useful for investigating the headgroup hydration, orientations, and lipid-lipid interactions (33–35). Interpreting vibrational lineshapes for phosphate groups can be challenging since the region is congested by the partial overlap among symmetric PO$_2^-$, asymmetric PO$_2^-$, and R-O-P-O-R' stretching modes. Additionally, the ester carbonyls are excellent reporters with well-known characteristics for interfacial hydration, interfacial dynamics, and lipid-protein interactions (27, 28, 35–37). The amide groups of proteins are also strong IR probes for protein structure and local environments (36, 38, 39). Given the rich molecular information derived from vibrational spectroscopy, many IR-based techniques including time-resolved methods have been used to investigate membranes and membrane proteins.
Figure 3. Illustration of common vibrational probes in lipid membranes. Top: Cartoon of a lipid bilayer section with a bound helical peptide (green) showing the different vibrational probes. Carbon, oxygen, nitrogen, and phosphate atoms are colored as gray, red, blue, and orange, respectively. Interfacial water molecules are shown in translucent blue. The approximate frequency of each probe is shown below the corresponding chemical structure. Bottom: FTIR spectrum of a dried 1,2-dimyristoyl-sn-glycero-3-phosphocholine (DMPC) lipid film measured under ambient conditions. The shaded areas indicate the frequency regions of the phosphate stretching mode (green), -OH bending mode (blue), amide I stretching mode (blue), ester carbonyl stretching mode (purple), -CH₂- stretching mode (yellow), and -OH stretching mode (brown). The horizontal axis is frequency in cm⁻¹, and the vertical axis represents the IR absorbance in arbitrary units. The absorbances across different regions are scaled for clarity.
2.1. Membrane Structure Characterized by IR Spectroscopy

Infrared techniques, such as Fourier-transform infrared spectroscopy (FTIR) can be used to measure the lineshapes associated with the different vibrational modes in the samples. Figure 3 (lower panel) shows an annotated example of an FTIR spectrum with the vibrational modes from phospholipid headgroup, ester linkage, acyl chain, and water molecule across a broad spectral range from 1000 to 4000 cm\(^{-1}\). The center frequencies and lineshapes of vibrational modes in IR spectra provide molecular information about the phase of the membrane as described by the ordering of the acyl chains, interfacial hydrogen bond networks of water reported by the -OH vibrational mode, and lipid hydration by phosphate headgroup and ester linkage (40–42).

The ester carbonyl stretching mode spans a region between 1725 and 1740 cm\(^{-1}\) depending on the lipid conformation and interfacial hydration level. In solution, the infrared lineshape of lipid ester carbonyl is broad and asymmetric due to the presence of two hydrogen-bonding species: carbonyls with no hydrogen bond and carbonyls with a single hydrogen bond. Abundant lipid species such as phosphatidylcholines (PC) do not contain hydrogen bond donors, and thus, any hydrogen bonds are with interfacial water molecules. Deconvolving the measured lineshape using peak models can quantify the hydrogen-bond populations at the water-lipid interface, thereby providing insights into water penetration into the interfacial region, carbonyl orientations, and interactions with the aqueous environment. As an example, an FTIR spectrum of 1-palmitoyl-2-oleoyl-sn-glycero-3-phospho-L-serine (POPS) and 1,2-dimyristoyl-sn-glycero-3-phosphocholine (DMPC) at a 1 to 9 molar ratio is shown in Figure 4. The ester carbonyl stretching region exhibits an asymmetric IR lineshape described by two overlapping peaks corresponding to two distinct hydrogen-bonded carbonyl species: zero hydrogen bond (0HB) and one hydrogen bond (1HB). The 1HB peak position is red-shifted by
approximately 15 cm\(^{-1}\) from the 0HB peak position as a result of the induced electric field by the partial positive charge on the hydrogen atom (43). Fitting the FTIR spectrum with two Gaussian profiles can help quantify the 0HB and 1HB populations, which involves computing the peak areas from Gaussian fittings and determining the relative extinction coefficients of the two hydrogen-bonding species (44). Quantifying hydrogen bond populations is critical for characterizing water penetration, lipid-water interfaces, and lipid packing efficiency. This quantity can also be compared with molecular dynamics simulations, allowing for a direct and one-to-one connection between experiments and simulations.

![FTIR spectrum of POPS:DMPC](image)

**Figure 4.** Example FTIR spectrum for phosphatidylserine (POPS): phosphatidylcholine (DMPC) vesicles at a 1:9 ratio in MOPS buffer in D\(_2\)O, with an illustration of two hydrogen-bonding species in a lipid. Left: Spectrum was fitted with two Gaussian profiles (1HB in yellow and 0HB in green). Dotted and solid lines show the experimental spectrum and the Gaussian fit spectrum, respectively. Right: Illustration of a PC lipid with one carbonyl hydrogen-bonded to a water molecule (1HB) shown in slate blue spheres, and the other without hydrogen bond. Hydrogen atoms in the lipid are omitted for clarity. Carbon, oxygen, nitrogen, and phosphate atoms are colored as gray, red, blue, and orange, respectively.

Measuring spectra with well-defined compositions, including those lipids shown in Figure 2, enables researchers to examine each component's influence on the interfacial properties of model
systems. Furthermore, temperature-dependent FTIR spectra are useful for investigating lipid phase transitions, and determining enthalpies and entropies associated with hydrogen bond formation (45). In addition to the ester carbonyl stretch, other vibrational modes also provide information about interfacial environments, morphologies, and local interactions. For example, a redshift in \(-\text{CH}_2\)-stretching peaks indicates more ordered acyl chains, as, for example, in the gel phase (46). Thus, this spectral region can be used to track phase transitions in model bilayers. It is important to note that probing intrinsic vibrational modes yields an ensemble-averaged view across oscillators. For instance, the hydrogen-bond number obtained from FTIR spectrum of POPS:DMPC represents the average number for all ester groups in both lipid species, and hydrogen bonds formed specifically with POPS cannot be differentiated from those with DMPC. Consequently, if an altered composition in lipid vesicles has a low concentration or weak signal, discerning the changes in lineshapes solely based on linear IR spectra becomes challenging. Isotope-label is a good approach to isolate the contributions from different lipid compositions and investigate the effects of individual components. For instance, the ester carbonyl groups of PC can be isotope-edited by \(^{13}\text{C}\), which gives rise to a red-shifted carbonyl stretch 40 cm\(^{-1}\) away from the carbonyl stretch of unlabeled PS (47). As a result, the effects of PC or PS can be studied separately, though the method is limited to investigating species with relatively high concentrations.

2.2. Transient Interactions in Lipid Membranes

Static spectra offer ensemble-averaged structural views of equilibrium conformations. Obtaining a more complete view of membranes requires measuring local environments and interactions, specifically molecular motions on sub-picosecond timescales. These measurements require ultrafast time-resolved spectroscopies such as two-dimensional infrared (2D IR) spectroscopy and two-dimensional sum-frequency generation (2D SFG) spectroscopy, which have become valuable
tools for detecting and characterizing dynamic processes in biological systems, due to the ability to combine molecular information with high time resolution.

**Structure and Dynamics by 2D IR Spectroscopy**

2D IR spectroscopy uses three ultrafast pulses, two pump pulses and one probe pulse, each with a temporal width of approximately 100 fs. The absorbance in 2D IR spectroscopy is correlated with the peak intensity of three laser pulses, thus improving its capability in detecting weak signals and discerning small changes in the IR lineshape. In 2D IR spectra (Figure 5, left), the horizontal and vertical axes represent the pump and probe frequencies, respectively. The positive peaks (red contours) correspond to the transition between ground and first-excited states, known as the ground state bleach (GSB). The negative peaks (blue contours) denote the vibrational transition between first- and second-excited state, namely the excited state absorption (ESA). The frequency of ESA is downshifted from GSB due to the anharmonicity of oscillation. The spectral widths of contoured peaks along the diagonal and antidiagonal of 2D IR spectra report on the inhomogeneous and homogeneous broadening mechanisms. Overall, the 2D lineshapes report on the distribution of local environments and their fluctuations (48, 49).

Figure 5. Example 2D IR spectra and dynamics analysis for POPS:DMPC vesicles at a 1:9 molar ratio in MOPS buffer in D$_2$O. Left: 2D IR spectra at selected waiting times: 0.15 and 4.0 ps. The solid black line represents the diagonal where pump and probe frequencies are equal. Dotted white
Dynamics of local environments surrounding the IR probes can be detected on a sub-picosecond timescale by varying the delay between pump and probe pulses ($t_2$). In 2D IR spectra, peaks are elongated along the diagonal at early waiting time and become rounder at late waiting time as pump and probe frequencies gradually decorrelate due to the frequency fluctuations when ensembles initially excited at a given frequency become randomized. This process is referred to as spectral diffusion. In essence, waiting-time-dependent 2D IR spectra measure the frequency-frequency correlation function (FFCF) of each vibrational probe within the system (48, 50, 51). The frequency decorrelation times, encoded within the 2D lineshapes, can be extracted using numerical analysis methods such as center line slope (52), nodal line slope (53, 54), ellipticity parameter (55), or fitting to a model response function (56, 57).

Compared to FTIR spectra, 2D IR spectra spread the information onto two frequency axes, which not only capture additional structural information through the 2D lineshapes and cross-peak intensities, but also access local environments around vibrational probes by the evolution of the lineshapes as a function of pump-probe delay (36, 58, 59). In addition, while FTIR spectra require measuring a separate background of the buffer solution for subtraction, 2D IR spectra are often measured without a separate background. This is because, in water, the low oscillator strengths of the solvent vibrations and broad lineshapes naturally suppress the background, thus removing the necessity of background subtraction. Furthermore, given the narrow lineshapes in 2D IR spectra, small differences that can be difficult to observe in FTIR spectra are often easier to analyze in the corresponding 2D IR spectra, as such, it is often useful to extract a corresponding “1D” spectrum.
from a 2D IR spectrum (60, 61). Recently, a robust method for extracting IR lineshape is the pump slice amplitude (PSA), which calculates the difference between the maxima and minima across a series of pump-axis slices through the 2D IR spectrum (62). The PSA method produces lineshapes comparable to FTIR spectra of the same system but with the advantages of eliminating artifacts due to anharmonicity and suppressing the background.

Examples of 2D IR spectra and lineshape analysis for POPS:DMPC at a 1:9 molar ratio are depicted in Figure 5. The left panel displays the 2D IR spectra in the ester carbonyl region at early (0.15 ps) and late (4.0 ps) waiting times. The signal is centered at approximately 1740 cm⁻¹, consistent with the FTIR spectrum (Figure 4). The broad band along the diagonal originates from the two hydrogen-bonding species (0HB and 1HB), as described previously (Figure 4), with 1HB red-shifted by approximately 15 cm⁻¹. The evolution of the lineshapes as a function of waiting time \( t_2 \) is used to extract the dynamics of the local environments around lipid ester carbonyl groups. In Figure 5, we provide an example of 2D lineshape analysis using the center line slope (CLS) method (63). In brief, CLS is the slope of the line connecting the maxima of peaks, which are a series of cuts through the 2D spectrum parallel to the pump frequency axis. CLS decays indicate the loss of frequency correlation. In the right panel of Figure 5, CLS for a lipid vesicle sample is calculated and plotted versus waiting time up to 5 ps. Since CLS decay measures the FFCF decay \( C(t_2) \), it can be represented by the following equation using the standard Bloch model (48):

\[
C(t_2) = \langle \delta \omega(t_2) \delta \omega(0) \rangle = \frac{\delta(t_2)}{T_2} + \Delta^2 e^{-\frac{t_2}{\tau_1}} + \Delta_0^2,
\]

where \( \delta \omega(t), \Delta^2, \tau_1 \) and \( \Delta_0^2 \) is the frequency fluctuation at time \( t \), the amplitude of frequency fluctuations, decay constant, and the static inhomogeneity components, respectively. \( T_2 \) is the
constant giving rise to the homogeneous width and $t_2$ is the waiting time between pump and probe pulses. The FFCF decay constant for the ester carbonyl stretch is determined to be approximately 1.07 ps for the 1:9 POPS:DMPC vesicles.

In membranes, the interfacial hydrogen bond network is disrupted due to interactions with the lipid headgroups and imbalance of donors and acceptors. In bulk water, the hydrogen bond dynamics are as fast as hundreds of femtoseconds (32, 64), but the dynamics become slower at the interface as a few picoseconds. In general, disordered water networks exhibit slower reorganization dynamics because the hydrogen bond switching process is a concerted process involving multiple water molecules (36, 64). However, since the spectrum measures frequency fluctuations, not only hydrogen bond switching, but also local fluctuations in hydrogen bond distances and angles significantly contribute to the measured loss in frequency correlation (29, 65). Measured dynamics from 2D IR can indicate the changes of water networks at the lipid-water interface and hence the effects of variables such as membrane composition, crowding, or interfacial localization of ions.

Here we show an example of how ester carbonyl 2D IR provides direct access to environment and dynamics at the lipid-water interface. It is important to emphasize that while 2D IR spectroscopy is not a surface-specific technique, the vibrational reporters are located precisely at the ~1 nm interface between the hydrophobic acyl chain region and bulk-like water beyond the headgroup region, and thus the measurements provide interfacial information. There are multiple avenues for applying 2D IR spectroscopy to understand how headgroup composition and lipid-lipid interactions determine the interfacial environments, contributing to a more complete understanding of membranes, ranging from the effects of headgroup charges (57) or ion localization (66), to the effects of transmembrane crowding on interfacial water networks (67).
**Interfacial Environment by SFG spectroscopy**

The membrane interaction with its environment is crucial for membrane function and stability. 2D IR can detect the local environment of the lipid interface by probing the frequency fluctuations of lipid ester carbonyl groups, where the lineshape and dynamics of 2D IR spectra are correlated with the water network at the interface. However, direct detection of the interfacial environment around the cell membrane can be difficult to perform with 2D IR, as the bulk water dominates the final signal, it is not possible to measure the interface-specific signals. Thus, a technique with interface selectivity is essential for isolating interfacial waters and measuring specific interactions involving waters at the membrane interface.

![Diagram of 2D SFG configuration for measuring interfacial water molecules with a lipid monolayer at the air-water interface. A “pump” beam (red) is introduced to resolve the excitation frequency. The other two input beams, green for visible light and pink for IR, act as the “probe”. The output signal (gray blue) is generated at the sum frequency of visible and IR wavelengths. A local oscillator (yellow) is introduced for heterodyned detection to resolve phase information.](image_url)

Figure 6. Diagram of 2D SFG configuration for measuring interfacial water molecules with a lipid monolayer at the air-water interface. A “pump” beam (red) is introduced to resolve the excitation frequency. The other two input beams, green for visible light and pink for IR, act as the “probe”. The output signal (gray blue) is generated at the sum frequency of visible and IR wavelengths. A local oscillator (yellow) is introduced for heterodyned detection to resolve phase information.

One-dimensional SFG, developed in the last 30 years, provides access to ensemble-averaged local environments at the interface, including dipole orientation, and specific hydrogen-bond networks,
though heterodyne-detection needs to be incorporated to obtain an unambiguous measure of the vibrational response (68). The SFG method uses a narrowband visible pulse combined with a broadband IR pulse, which generates an output signal at the sum frequency of two input pulses if vibrational probes in the system resonant with the IR pulse. The output SFG pulse is mixed with a local oscillator to deconvolve the phase information. Two-dimensional vibrational sum-frequency generation (2D SFG) spectroscopy, which incorporates additional pulses to resolve dynamics, has advanced significantly over the past five years, becoming a powerful tool to investigate interfacial dynamics of membranes (69). An IR pump pulse is implemented in SFG to provide a frequency-resolved excitation axis, similar to the pump axis in 2D IR spectroscopy. 2D SFG can be interpreted as a frequency-resolved IR “pump” and SFG “probe” technique. 2D SFG has the advantage of high selectivity towards interfacial regions with anisotropic molecular orientations, inherent to second-order nonlinear optical processes (70). To avoid the interference of multiple interfaces, a monolayer at the air-water interface is usually prepared (Figure 6) (36, 71). Using 2D SFG, the dynamics of the hydrogen-bonded and free \(-\text{OH}\) stretches of water are probed through characteristic peaks in the region of 3200 to 3700 cm\(^{-1}\), providing direct access to the hydrogen bond networks of water within the first few solvation shells around the lipid headgroups.

2D SFG has been recently used to measure interfacial water dynamics for the negatively charged lipid, 1,2-dipalmitoyl-sn-glycero-3-phosphoglycerol (DPPG), versus the positively charged surfactant, 1,2-dipalmitoyl-3-trimethylammonium-propane (DPTAP) (72). The lipid monolayers are prepared on isotopically diluted water, where the \(-\text{OH}\) stretch of HOD gives rise to the majority of SFG signal in the \(-\text{OH}\) stretching region. Interestingly, steady-state SFG spectra of DPPG and DPTAP show opposite signs (Figure 7a-b), indicating the opposite net orientation of interfacial water due to the negatively and positively charged lipid headgroups. However, the peak position
and lineshape in the SFG spectra are similar for DPPG and DPTAP, demonstrating the similar hydrogen bond environment, despite the opposite charges.

Figure 7. Heterodyne-detected SFG and 2D SFG spectra of water at the air-water interface of positively and negatively charged lipids. (a, c) Steady-state and 2D SFG spectra at the DPPG interface; (b, d) Steady-state and 2D SFG spectra at the DPTAP interface. The horizontal and vertical axes represent the $\omega_2$ (SFG probe) and $\omega_{\text{pump}}$ (IR pump) frequencies, respectively. (e) Center line slopes for DPPG (blue) and DPTAP (red) interfaces as a function of IR to SFG delay. Schematic models for DPPG (left) and DPTAP (right) interfaces are also shown at the bottom. Figure is reprinted with permission from reference (68). Copyright 2023 American Chemical Society.
In addition, 2D SFG spectroscopy probes the picosecond spectral diffusion of the -OH stretch (Figure 7c-d). The interpretation of the lineshapes is similar to that of 2D IR spectra, where the evolution of the lineshapes provides a measure of the frequency-fluctuation correlation function of the interfacial water molecules. The time-dependent 2D SFG spectra are analyzed by the center line slope method, where different dynamics are observed for DPPG and DPTAP. In brief, the negatively charged lipid layer disrupts the water network and suppresses the fast HB dynamics due to strong hydrogen bonds involving the hydrophilic DPPG phosphatidylglycerol headgroup (Figure 7). In contrast, the positively charged DPTAP layer does not have significant effects on the interfacial water ordering and dynamics, as evidenced by the fast dynamics that is comparable to bulk water (32, 64). In summary, 2D SFG directly probes interfacial environments, providing a specific view of the relation between membrane composition and water dynamics. The technique can also be used to study membrane proteins (73, 74), pH effects (75), and cation effects (76) in the context of lipid membranes.

2.3. Surface-Enhanced Infrared Absorption Spectroscopy (SEIRAS)

Surface-enhanced infrared absorption spectroscopy (SEIRAS) has emerged as a useful technique for studying lipid membranes, as it significantly enhances detection sensitivity for monitoring conformation and structure at single bilayer interfaces without interference from bulk solvent backgrounds. In SEIRAS, a thin, nanoscale-roughened, metal film is deposited on an attenuated total reflectance (ATR) crystal, enabling the IR beam to excite local surface plasmons in the metal. The plasmonic near-field enhancement partially penetrates ~100 nm from the surface into the sample (Figure 8). The IR signal is greatly enhanced for molecules adsorbed on the metal surface, with surface plasmon polariton modulation resulting in a 10-100× increase in IR absorption (77).
Specifically-engineered nanostructured materials can provide even higher enhancement factors (78–81).

![Diagram of SEIRAS setup](image)

Figure 8. Illustration of an experimental setup of SEIRAS with an adsorbed lipid bilayer (red) on a metal surface (yellow). The blue trapezoid represents the ATR prism. The dashed lines represent the enhancement in electromagnetic field from the generated surface plasmon resonances. The box on the right shows a zoomed-in view of the lipid-metal interface. Figure is adapted with permission from (77).

Initial developments provided a 20× enhancement in the IR absorption of organic monolayers on thin metal films. Osawa and coworkers later noted general properties and the theory for SEIRAS: the IR enhancement is general for nonvolatile molecules; the enhancement increases for molecules strongly chemisorbed on the metal film; and the effect is observed only for the first few adsorbed monolayers (82–84). More recent models provide a surface-average enhancement factor and IR penetration depth from experiment as the ratio of molar absorptivities of SEIRAS to the bulk (85).

A general experimental procedure for SEIRAS involves the formation of a metal layer, typically 100-200 nm of Au or Ag, on a polished ATR substrate through electrochemical deposition (86, 87), metal vapor deposition in vacuum (88–92), or electroless (chemical) deposition (93, 94). The latter technique yields more stable metal films and larger islands on the surface. The enhancement factor depends on the thickness of the film, as thicker films exhibit increased IR reflectivity.
Because of the rapid decay in the enhancement penetration depth, the solvent absorption is minimized. This allows for direct study of specific regions in the membrane with the resolved intrinsic vibrational probes in the lipid membrane. SEIRAS is useful for studying interfacial properties, such as water structure near a supported lipid bilayer (95–97), with very low penetration depth into the bulk. Lipid membranes have large potentials because of concentration or pH gradients, and SEIRAS enables studying the effects of fields and potentials in biological systems using vibrational probes to measure the local electrostatic environments. Additionally, SEIRAS provides the ability to rapidly change sample conditions such as pH or ion concentrations and the ability to measure kinetics on the timescale of seconds to hours (98).

Figure 9. SEIRAS spectra of the titration of a pH-low insertion peptide (pHLIP) inserting/folding into a POPC bilayer. (a) Amide I absorbance increases as pH is decreased (purple to red). (b) The change in absorbance at 1663 cm\(^{-1}\) is plotted as a function of pH. The insets are cartoons showing pHLIP in an inserted/folded state in the membrane at low pH compared to at high pH.

In typical SEIRAS measurements, a lipid vesicle solution is prepared and deposited on the metal surface. A spacer of adsorbed molecules can be used to form “floating” lipid bilayers (96, 97),
mitigating any effects from the metal surface on the membrane properties. The vesicles rupture and self-assemble as an ordered monolayer on the surface, eventually forming a bilayer (99, 100). The bulk solvent can be easily replaced with different solvents, and solutions containing transmembrane proteins can also be added to study protein insertion (101) (Figure 9) or potential-dependent structural changes (102). SEIRAS can also be used to monitor time-resolved \textit{in-situ} processes, such as the expression and folding/inserting of transmembrane proteins into lipid membranes (98). The folding/insertion process and conformational changes can be tracked by the lineshape of amide bands over time.

Relevant advancements in SEIRAS techniques include time-resolved SEIRAS (103, 104), surface-enhanced ultrafast IR (SEIR) (105) and 2D IR (SE2DIR) spectrosopies (102, 106–109). SE2DIR enables the measurement of species with low molar absorptivity in both transmission and reflection geometries with deposited metals < 10 nm of thickness (105, 106). A new plasmonic design has been reported to maximize surface coverage and enhancement while reducing background signals and lineshape distortions (107). The capabilities and recent developments of SEIRAS techniques demonstrate that this technique is suited to address key questions surrounding lipid membranes. The low penetration depth into the sample allows one to study the membrane structure, morphology, interface, and local membrane environments with enhanced single-bilayer detection sensitivity while minimizing signal contributions from bulk solvent. The ability to resolve signals of protein amide bands allows one to elucidate the effect of lipid-protein interactions on protein conformation and the mechanism and kinetics of the folding/inserting of transmembrane proteins.

For example, one can envision differential measurements using isotope-labeled lipids in multicomponent bilayers to examine the functional paralipidome with model peptides. Other variables such as pH, peptide composition, or ion concentrations can be easily altered by replacing
the solvent after lipid bilayers are formed on the SEIRAS substrate. Time-resolved and ultrafast SEIRAS, similar to 2D IR, can measure kinetics on the picosecond timescale, and SE2DIR further improves the signal-to-noise ratio in measurements. Furthermore, there are other possibilities to investigate intact plasma membranes, for example, by combining SEIRAS with cell-unroofing techniques (110).

3. Interpretation of Experiments through Molecular Dynamics Simulations

3.1 Molecular Dynamics Simulations

Interpreting spectra is one of the main challenges due to complex lineshape and spectral congestion in multicomponent systems. For example, when analyzing a mixture, the resulting spectra provide an average of all vibrational modes across different species, making it difficult to separate and identify individual contributions (57, 66, 111). Isotope labeling can be used to isolate individual species provided that the labeled species are present at high concentrations. In addition, interface-specific spectra can be difficult to interpret as coupling and delocalization make vibrational assignments difficult, even for well-defined interfaces such as the air-water interface (68, 112).

Evidently, lipid membranes add complexity as the diversity of lipids, proteins, cholesterol, and carbohydrates makes for a highly heterogeneous interfacial environment. For these reasons, molecular insights are most often derived by computing spectra based on atomistic modeling, and molecular dynamics simulation is an ideal technique to provide molecular ensembles for spectra computation (113–117).

Computed spectra provide the most direct route to interpret spectral features arising from interactions with lipids, water, proteins, ions, and other small molecules. Computing spectra from structural ensembles requires the use of models, such as electrostatic maps, which translate the
local electrostatic environment around each oscillator to a frequency shift (118). The advantage of these maps is that they provide a computationally tractable means of generating spectra for systems that contains hundreds of oscillators due to their computational efficiency (119). These structure-based vibrational models provide a one-to-one comparison between experiment and simulation, and a route to interpret complex lineshapes in terms of local environments. Only short trajectories are required, allowing for very large or complex systems to be simulated. Overall, MD simulations for relatively uniform membranes have become increasingly reliable, with simulated results now achieving near-quantitative agreement with experiment in certain systems (111, 120, 121).

3.2 Challenges of All-atom Simulations

Sampling is one of the ongoing challenges for simulations. Obtaining equilibrated ensembles is difficult because of the large number of degrees of freedom, relative lack of experimental reference to dictate initial ensembles, and the slow lipid reorganization timescales (122–124). New algorithms such as Molecular Dynamics with Alchemical Steps (MDAS) have been developed to mitigate the conformational sampling problem(122). The MDAS algorithm combines conventional MD simulations with alchemical trajectories, which swap the positions of randomly selected pairs, analogous to Monte Carlo simulations, but using a short trajectory where the parameters of two lipids are exchanged. In addition, within the functional paralipidome model described above, different proteins can recruit distinct lipid environments, including specific subtypes with various headgroups. The local membrane properties can be modulated by external factors to accommodate protein structure and function in membranes (125). Enhanced sampling methods are useful for equilibrating such complex systems.
Another challenge is the parameterization of lipid force fields (124, 126, 127). Several all-atom lipid force fields, including CHARMM36, AMBER (LIPID21), Slipids, and polarizable models, have been published and benchmarked in combination with various water models (128–131). However, due to the complex lipid interactions and the number of lipid species, achieving quantitative agreement between simulated and experimental observables beyond simple bilayers remains a challenge (124, 127). Moreover, bilayer properties are sensitive to weak and long-range interactions, which are difficult to accurately parameterize (132–134). Modern computer software and recent multiscale models combined with machine learning present new opportunities for further development (126, 135, 136). There are many opportunities to use vibrational techniques for characterizing interactions in heterogeneous samples, and provide useful molecular-level information for benchmarking simulations (43, 121, 137, 138). Using spectroscopic techniques described on a library of lipid compositions, and modern MD simulations could be a powerful approach to measure the balance of lipid-lipid, lipid-protein, and lipid-water forces that lead to the heterogeneity in multicomponent membranes.

4. Ongoing Challenges, Prospects, and Opportunities

Vibrational spectroscopy has advanced enormously over the past decade. The spectroscopic approaches discussed above have offered exciting opportunities to investigate membrane structure and dynamics in increasingly complex systems. One key challenge is achieving spatial resolution, and the ability to map the nanometer-scale heterogeneity for lipid organization. Nano-IR microscopy, which uses an AFM tip to localize electric fields, and thereby achieve spatial resolution, could be coupled with time-domain techniques to directly probe membrane structure and dynamics from single bonds to micron-scale length (139–142).
The second challenge is membrane asymmetry. Specialized lipids are often enriched in microdomains and unevenly distributed across leaflets. For example, phosphatidylserine (PS) is mainly distributed in the inner leaflet of the plasma membrane in healthy eukaryotic cells due to the activity of transporter proteins. Therefore, characterizing PS distribution is crucial for understanding the functions of lipids in cellular homeostasis and membrane trafficking. Sum-frequency generation spectroscopy could provide a direct measure of net-orientation, and thus measure enrichment across the leaflets. However, challenges would certainly arise in future experimental design of in-vivo SFG measurements of plasma membranes. Another challenge, as discussed above, is the large diversity of lipid species present in cell membranes. While simple lipid compositions have been extensively investigated, building more complex or realistic lipid membranes remains challenging. Vibrational spectroscopy provides ensemble-averaged results across all oscillators, irrespective of lipid species, and isotope-labeling, e.g., $^{13}$C (41, 47, 143, 144), or functional-group-labeling, e.g., nitrile or thiocyanate groups (145, 146), can help identify the contribution of each species individually. MD simulations can also help interpret complex spectra. However, building heterogeneous membranes introduces complicated interactions, and requiring more sensitive vibrational techniques to probe the subtle yet important effects introduced by minor lipid species. Meanwhile, building synthetic systems with artificial lipids (147, 148), lipid nanodiscs (149), and artificial channels (150) can serve as useful platforms to study lipid-protein interactions in minimalistic or well-defined environments.

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**Terms and Definitions:**

*Fluid-mosaic model:* an influential membrane model consisting of proteins embedded within the lipid bilayer.

*Lipid raft:* dynamic regions in biological membranes enriched in certain components, influencing membrane organization and protein localization.

*Functional paralipidome:* a general model or framework for describing the mutual regulation between membrane proteins and their surrounding lipid microenvironments.

*Hydrogen bond:* a directed electrostatic interaction between polar atoms, involving a protic hydrogen.

*Vibrational probe:* a functional group with a well-defined vibrational frequency response that can be used to measure local environments and dynamics.

*Two-dimensional infrared (2D IR) spectroscopy:* a frequency-resolved “pump-probe” technique to measure molecular conformations and dynamics on picosecond timescales.

*Sum-frequency generation (SFG) spectroscopy:* an interface-specific spectroscopy used to measure the environments, water orientations, and hydrogen bond dynamics at single interfaces.

*Pump-slice amplitude (PSA) analysis:* an analysis method to extract FTIR-like lineshapes from 2D IR spectra.
Frequency-fluctuation correlation function (FFCF): a measure of the evolution of the local environment as reported by the frequency fluctuations around a vibrational probe.

Surface-enhanced infrared absorption spectroscopy (SEIRAS): an approach to measure surface-sensitive vibrational spectra with a penetration depth of ~100 nm using gold nanoparticles.

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