

Understanding the nanoscale structure of hexagonal phase lyotropic liquid crystal membranes

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Introduction

Nanostructured membrane materials have become increasingly popular for aqueous separations applications such as desalination and biorefinement because they offer the ability to control membrane architecture at the atomic scale permitting the design of solute-specific separation membranes. [?] Most membrane-based aqueous separations of small molecules can be achieved using reverse osmosis (RO) or nanofiltration (NF). [?]

While RO and NF have seen many advances in the past few decades, they are far from perfect separation technologies. Current commercial RO membranes are thin film composite membranes with a porous support layer and an active layer made of a dense polyamide polymer matrix. The membranes are unstructured with tortuous and polydisperse diffusion pathways. Separation occurs as solutes dissolve into and diffuse through voids in the polymer matrix at different rates.[?] With optimization, one can exploit these differences to create a functional selective barrier. High feed pressures are required in order to achieve economical solvent flux. The largest application of reverse osmosis is desalination. A typical seawater desalination process requires feed pressures in the range of 800 to 1000 psi in order to overcome osmotic pressure. [?] Only 0.5 % of the world's water is fresh. Desalination is an important source of potable water, necessary in order to keep up with demand generated by a rapidly growing global population. Although RO is currently the cheapest form of desalination, the cost must be further decreased in order to meet demand. By designing better membranes for desalination, we can achieve higher water fluxes and lower feed pressure requirements which will decrease the cost of fresh water production. [?].

NF was introduced as an intermediate between RO and ultrafiltration, having the ability to separate

organic matter and salts on the order of one nanometer in size. Explicit pore pathways running through the thickness of the membrane permit high solvent fluxes at lower pressures than RO. NF membranes typically have a surface charge resulting from ionizable groups which affords separation of ions smaller than the pore radius by Donnan exclusion. [?] NF is often used as a precursor to reverse osmosis since it can efficiently remove a significant portion of dissolved solutes. Following NF pretreatment, RO can further purify the permeate at lower pressure. Unfortunately, NF membranes, like RO, possess a pore size distribution which limits their ability to perform precise separations [?]. An NF membrane with high selectivity could offer the versatility to separate ions as well as small charged and uncharged molecules to the same degree as RO and at a significantly lower cost.

Nanostructured membranes can bypass many of the performance issues which plague traditional NF and RO membranes. One can accomplish targeted separations with high selectivity by tuning shape, size and functionality of the molecular building blocks which form these materials. As a result, solute rejecting pores can have their sizes tuned uniformly, resulting in strict size cut-offs. Entirely different mechanisms may govern transport in a given nanostructured material which can inspire novel separation techniques.

Development of nanostructured materials has been limited by the ability to synthesize and scale various fundamentally sound technologies. Graphene sheets are atomically thick which results in excellent water permeability but defects during manufacturing severely impact selectivity. [?] Molecular dynamics simulations of carbon nanotubes show promise [?] but synthetic techniques are unable to achieve scalable alignment and pore monodispersity. [?, ?] Zeolites have sub-nm pores with a narrow pore size distribution and MD simulations exhibit complete rejection of solvated ions, [?] however, experimental rejection was low and attributed to interstitial defects formed during membrane synthesis. [?] There is a need for a scalable nanostructured membrane.

Self assembling lyotropic liquid crystals (LLCs) are a suitable candidate for aqueous separation applications. LLCs share the characteristic ability of nanostructured membrane materials to create highly ordered structures with the added benefits of low cost and synthetic techniques feasible for large scale production. [?] LLC systems created by the monomer named Na-GA3C11 have been extensively studied experimentally. [?, ?, ?, ?, ?] Neat liquid crystal monomer forms the thermotropic, Col_h phase. The presence of ca. 10 wt % water results in the H_{II} phase. In both cases, monomers assemble into mesophases made of hexagonally packed, uniform size, cylinders with hydrophilic head groups oriented inward towards the cylinder center. The hydrophilic region can act as a pore for aqueous separations. One can envision tailoring the pore region for specific separations by changing the monomer chemistry.

Research into LLC membranes has been revived in recent years. During early stages of exploration, mesophases formed by Na-GA3C11 could not be macroscopically aligned, resulting in low flux membranes,

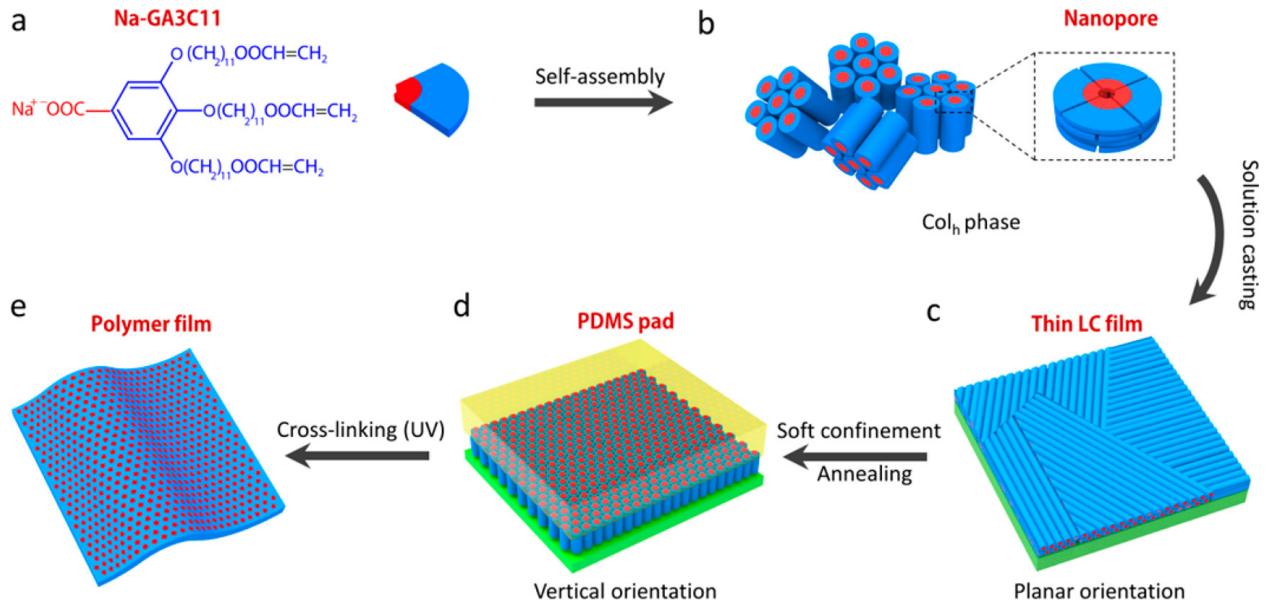


Figure 1: The wedge shaped liquid crystal monomer (a) self assembles into mesophases with hexagonally packed pores (b). The pores are made of stacked monomer disks. A sub-micron-thick film is created by casting a dilute solution of Na-GA3C11/THF solution onto a silicon substrate and allowing the solvent to evaporate. The thin film contains nanoporous columns which lie parallel to the film plane. (d) When a soft PDMS pad is imposed to the thin film, with subsequent thermal annealing, the columns align perpendicular to the film plane. (e) Photo-cross-linking of the aligned film creates a mechanically stable thin film with vertically aligned nanopores

and no clear route towards scalable and economical filtration. In 2014, Feng et al. showed that the mesophases could be aligned using a magnetic field with subsequent crosslinking to lock the structure in place. [?] In 2016, Feng et al. showed that the same result could be obtained using a technique termed soft confinement. [?]

Our current understanding of LLC systems is not rich enough to be able to precisely design membranes for specific separations. Over the past 20 years, H_{II} phase LLC membrane studies have been limited primarily to Na-GA3C11 with some characterization done after minor structural modifications. Resel et al. varied the length of the monomer tails and the counterion used and observed its affect on pore spacing [?]. Their study offers no insight into dynamics within the pore. Rejection studies show that membranes formed by NA-GA3C11 can not perform separations of solutes less than 1.2 nm in diameter because the pores are too large [?]. We do not yet understand how to controllably reduce the effective pore size or how to tune the chemical environment in the nanopores for effective water desalination or small organic molecule separations. The only source of predictive modeling for LLC systems have been macroscopic models which likely do not adequately describe transport at these length scales. [?] It will be challenging to efficiently narrow down the large design space in a laboratory setting without a robust model.

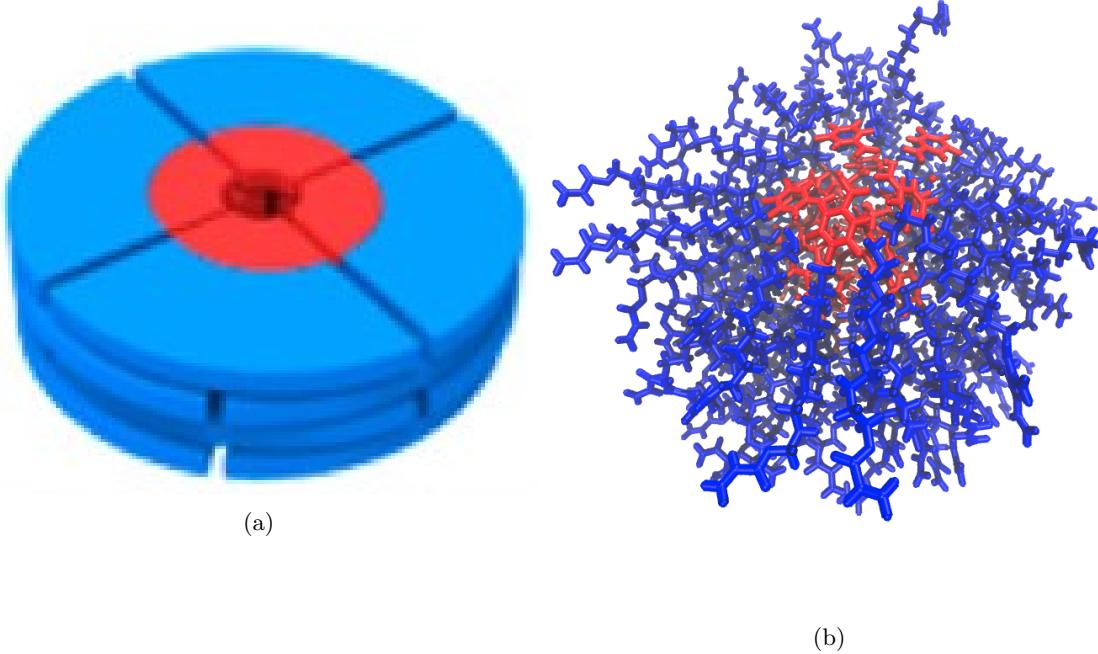


Figure 2: (a) Our previous understanding of the pore structure allows us to speculate about separation behavior. (b) A detailed molecular model will allow us to directly observe solute transport. Here, four stacked layers of 5 monomers are pictured atomistically. The hydrophilic region is in red and the hydrophobic region is colored blue.

A molecular level understanding of LLC membrane structure, enabled by molecular dynamics simulations, will provide guidelines to reduce the large chemical space available to design monomers for creation of separation-specific membranes. A good molecular model should incorporate a detailed picture of the nanoscopic pore structure which will be crucial to understanding the role of monomer structure in solute transport and membrane design. Models resulting from molecular dynamics simulations will provide the required level of detail. We can directly observe solute transport and suggest governing mechanisms. We can observe how the choice of head group may influence pore size for size exclusion driven separations. We can interchange counterions which may influence both the pore size and the strength of the Donnan potential which affects the degree to which the membrane can exclude charged species.

In order to appropriately model transport, we must gain a thorough understanding of the nanoscopic structure of LLC membranes. Our approach to constructing a general model will follow the development of a model of the assembly formed by Na-GA3C11 since it has sufficient experimental characterization. We have also narrowed our scope to the development of a model of the Col_h phase membrane. Compared to the H_{II} phase, the Col_h phase is a simpler starting point, due to the absence of water, and has detailed experimental

Wide-angle X-ray scattering (WAXS) patterns useful for reconstructing structural data.

Despite having structural data, there is still information which experiment cannot definitively answer. There are several key questions that we will investigate which will be laid out and numbered in subsequent paragraphs.

Monomers in the Col_h system are theorized to be partitioned into stacked layers which form columnar pores. We want to know

1. If layers do exist, how many monomers constitute a single layer?

A simple molecular simulation study of a similar molecule suggested that there are 4 monomers in each layer. Their estimation is based on a simulated system containing only 16 total monomers which likely does not sufficiently model the chemical environment present in the real system. [?]. A separate calculation based on the volume of the liquid crystal monomers proposes that there are seven monomers in each layer [?]. A molecular model orders of magnitude larger than any other reported atomistic liquid crystal membrane simulations has the best chance of directly answering this question. We will directly change the layer composition and note its effect on membrane structure.

2. Does our model support the existence of layers and if so, how well defined are the layers?

Experimentally, their existence is supported by evidence of strong π - π stacking interactions in the direction perpendicular to the membrane plane. π - π stacking will only occur between the aromatic monomer head groups which leaves no description of what is happening in the monomer tail region. The tails may entangle isotropically while stacking order is maintained among headgroups.

3. How do monomers in each layer position themselves with respect to surrounding layers?

The π - π stacking interactions may be a driving force of self assembly in this system. [?] Gas phase ab initio studies of benzene dimers have shown a clear energetic advantage for parallel displaced and T-shaped π - π stacking conformations versus a sandwiched conformation [?]. Substituted benzene rings exhibit an even stronger π - π stacking attraction which favors the parallel displaced configuration in all cases except where the substitutions are extremely electron withdrawing. [?, ?]. We can use simulated X-ray diffraction patterns to compare the two stacking configurations.

4. Can the system exist in other metastable states or phases that are not accessed during experiments?

There remains the possibility that there is more than one metastable state associated with a given LLC system. Simulating a membrane atomistically will require many atoms which limits the timescales accessible with MD. It is reasonable to expect that we will generate configurations which are kinetically

trapped in a metastable free energy basin. We must be able to identify which state is produced experimentally and why others are not.

5. What constitutes a pore and how well-defined are the pore regions?

The limited picture that experiment provides tells us that there are hexagonally packed, hydrophilic regions where transport is likely to occur. One may instinctively assume that these regions are tube-like pathways. We will explore the composition of the pores and the partition between the hydrophilic and hydrophobic regions.

6. Is it necessary to include any water in order to appropriately model the Col_h phase?

While the Col_h phase is described as dry, it has been suggested by experimentalists in unpublished communications that it is likely that small amounts of ambient water are leached into neat monomer. Experimentally, achieving a hexagonal phase with a completely dry system has proven difficult. If neat monomer is allowed to sit in ambient conditions, its color turns from transparent to slightly opaque and a hexagonal phase forms. Although we will not explore whether water is necessary for self assembly, we hypothesize that the hydrogen bonding network formed by the water may play a role in structuring the pores and holding together the hexagonal phase. We can use simulated X-ray diffraction patterns to see if there is any meaningful structural difference between a "dry" and "wet" system.

In this study, we build a significantly more realistic atomistic model of LLC membranes than, to our knowledge, has ever previously been done, and explore what new structural information can be gained and what structure hypotheses are supported by this model. We validate the model using as much experimental information as possible. We are most interested in reproducing the conclusions about structure which have been drawn from X-ray diffraction (XRD) experiments and in matching ionic conductivity measurements [?].

We used experimental wide angle X-ray scattering (WAXS) data (produced as described in [?]) and small-angle X-ray scattering (SAXS) data from [?] to inform our choices of some initial structural parameters (Figure 3). We rely primarily on the 2D WAXS data since it encodes all structural details down to the sub-nm scale. There are five major features of interest present in the 2D experimental pattern shown in Figure 3b. The first is located at $q_z = 1.7 \text{ \AA}^{-1}$, corresponding to a real space separation of 3.7 \AA . The reflection is attributed to $\pi-\pi$ stacking between aromatic rings in the direction perpendicular to the membrane plane, or z-axis [?]. For simplicity, this reflection will be referred to as R- π . A weak intensity line is located at exactly half the q_z value of R- π ($q_z = 0.85 \text{ \AA}^{-1}$), corresponding to a real space periodic spacing of 7.4 \AA . This reflection has been interpreted as 2_1 helical ordering of aromatic rings along the z axis meaning if the

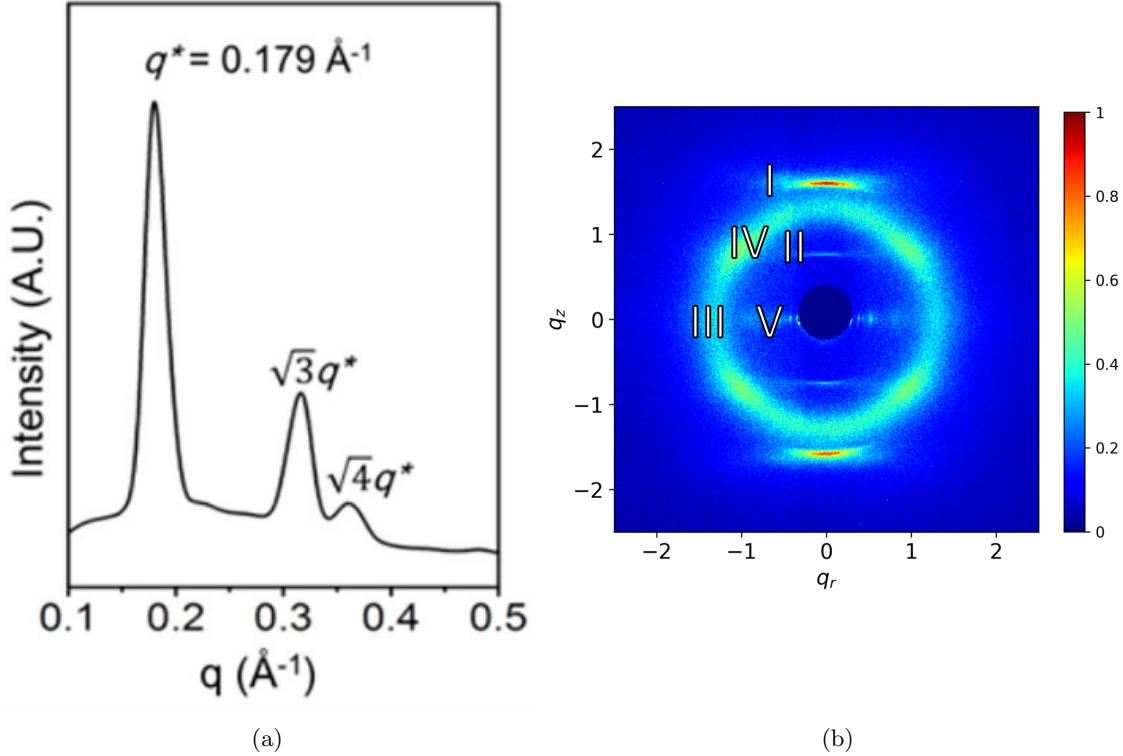


Figure 3: (a) 1D small angle X-ray scattering indicates hexagonal packing of pores as well as the spacing between pores (b) 2D wide angle X-ray scattering gives details about repeating features less than 1 nanometer apart

positions of the aromatic rings can be traced by a helix, then for each turn in the helix, there should be two aromatic rings. For this reason it will be referred to as R-helix. A third major reflection is marked by a low intensity ring located at $r = 1.4 \text{ \AA}^{-1}$. The real space separation corresponds to 4.5 \AA which is characteristic of the average spacing between packed alkane chains. This reflection will be called R-alkanes. Within R-alkanes, are four spots of higher relative intensity which will be called R-spots. All are located ≈ 40 degrees from the q_z axis in their respective quadrants. In many liquid crystal systems this can be explained by the tilt angle of the alkane chains with respect to the xy plane. The final feature corresponds to the spacing and symmetry of the d_{100} plane which can be related to the distance between pores. The feature, which will be called R-pores, is characterized by dots along $q_z = 0$. The spacing between dots is indicative of the hexagonal symmetry of the packed pores. The same information at higher resolution is obtained using a SAXS setup. By radially integrating the 2D data one gets a 1D curve which is shown in Figure 3a.

Methods

Liquid crystal monomers were parameterized using the Generalized AMBER Forcefield [?] with the Antechamber package [?] provided with AmberTools16 [?]. Atomic charges were assigned using the am1bccsym method of molcharge shipped with QUACPAC from Openeye Scientific Software. All molecular dynamics simulations were run using Gromacs 2016 [?, ?, ?, ?]

An ensemble of characteristic, low-energy vacuum monomer configurations were constructed by applying a simulated annealing process to a parameterized monomer.

- Monomers were cooled from 1000K to 50K over 10 nanoseconds.
- A low energy configuration was randomly pulled from the trajectory and charges were reassigned with molcharge.
- Using the new charges, the monomer system was annealed again and monomer configurations were pulled from the trajectory to be used for full system construction (Figure 4a).

The timescale for self assembly of monomers into the hexagonal phase is unknown and likely outside of a reasonable length for an atomistic simulation, calling for a more efficient way to build the system.

- Previous work has shown a coarse grain model self assemble into the H_{II} phase configuration in ≈ 1000 ns [?].
- We attempted atomistic self-assembly by packing monomers into a box using Packmol [?].
- Simulations of greater than 100 ns show no indicators of progress towards an ordered system.
- To bypass the slow self-assembly process, python scripts are used to assemble monomers into a structure close to one of a number of hypothesized equilibrium configurations (Figure 4).

A typical simulation volume contains four pores in a monoclinic unit cell, the smallest unit cell that maintains hexagonal symmetry when extended periodically.

- Each pore is made of twenty stacked monomer layers with periodic continuity in the z direction, avoiding any edge effects and creating an infinite length pore ideal for studying transport.
- A small number of layers is preferred in order to reduce computational cost and to allow us to look at longer timescales.
- Ultimately, we chose to build a system with 20 monomer layers in each pore in order to obtain sufficient resolution when simulating X-ray diffraction patterns. This point will be explained in more detail later.

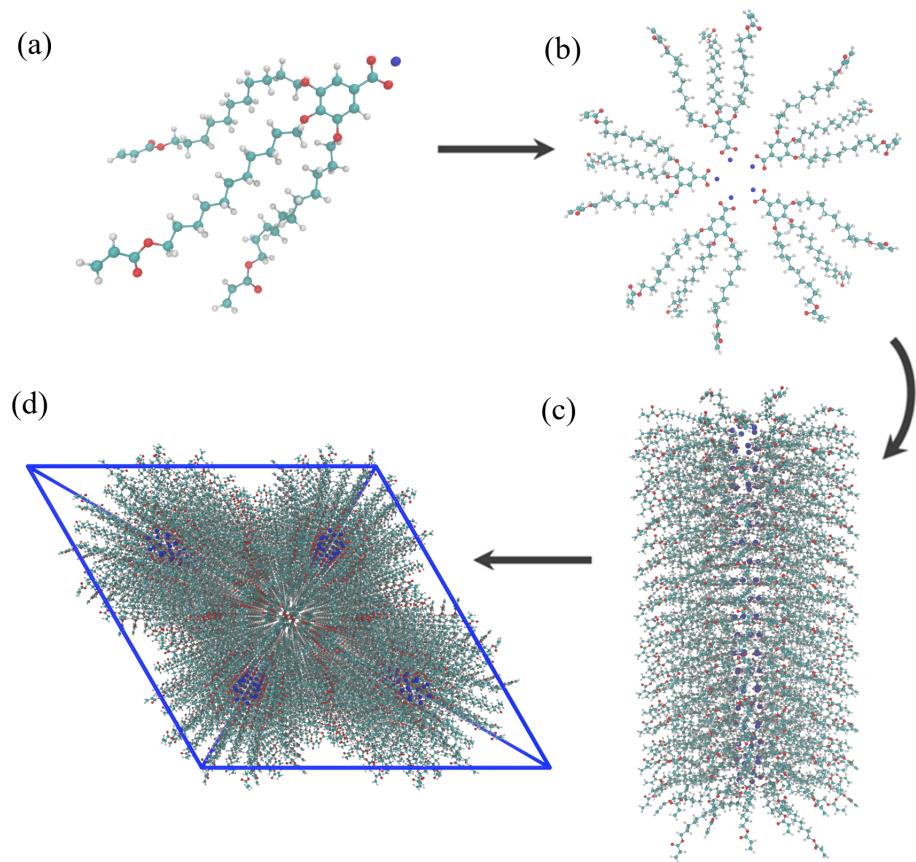


Figure 4: (a) A single monomer was parameterized and annealed to produce a low energy configuration. (b) Monomers are rotated and assembled into layers with hydrophobic centers. (c) Twenty layers are stacked on top of each other to create a pore. (d) Pores are duplicated and placed into a monoclinic unit cell

When constructing an initial configuration, there are a number of variables which require careful consideration while placing monomers. The equilibrium configuration is sensitive to some while insensitive to others.

- The starting pore radius, defined as the distance of a chosen head group carbon from the pore's central axis, does not influence the equilibrium structure when a reasonable value is chosen (See Supplemental)
- The pore radius is chosen to be 0.6 nm in our initial configurations because the pore size is estimated to be \approx 1.2 nm
- The initial distance between pores also has little effect on the equilibrated structure. However, one should not start them too close or there will likely be unintended high energy repulsions during early equilibration
- We chose an initial pore spacing of 4.5 nm, \approx 10 % larger than the experimental value of 4.12 nm.
- The distance between layers and the rotation of the layers with respect to adjacent layers, and the number of monomers per layer do influence the equilibrium structure and require further justification for their choices.
- We rely on experimental data to inform our choices

We chose the layer spacing for the initial configuration based on $R-\pi$ and then allowed system to readjust during equilibration.

- Each monomer was rotated so the plane of the aromatic head groups would be coplanar with the xy plane.
- We explore two different initial layer spacings
- The first is exactly equal to $R-\pi$ with layers placed so aromatic rings are stacked 3.7 Å apart in the z-direction.
- A second system is explored with an initial layer spacing of 5 Å
- A third system with an initial layer spacing of 10 Å was briefly explored
- When layers are spaced out sufficiently, they tend to collapse on each other while simultaneously slipping in between layers of other pores which lead to an artificially thick membrane with pores spaced closely together.
- The interested reader can learn more about it in the Supplemental Information.

The relative interlayer orientation was chosen based on clues from diffraction data as well as the various known stacking modes of benzene and substituted benzene rings: sandwiched, parallel-displaced and T-shaped [?] (Figures 5a to 5c).

- The T-shaped configuration was ruled out based on the inconsistency of its $\approx 5 \text{ \AA}$ equilibrium stacking distance [?].
- The system's preference towards the sandwiched vs. parallel displaced stacking modes will be explored.
- Both have reported stacking distances near the $R-\pi$ value of 3.7 \AA
- Headgroups in our sandwiched initial configuration are stacked directly on top of each other while stacked headgroups in the parallel displaced initial configuration are offset by $180/nmon$ degrees where $nmon$ equals the number of monomers per layer.

As outlined in 1 the number of monomers in each layer is unknown so we tested configurations constructed with a varied number of monomers per layer.

- Systems were built in the offset and parallel displaced configurations with 4, 5, 6, 7 and 8 monomers per layer

We developed equilibration schemes to create dry and wet configurations. Both schemes start with an initial configuration generated according to the previous guidelines.

- Dry equilibration scheme:
 - Restraints with a force constant of $1\text{e}6 \text{ KJ mol}^{-1} \text{ nm}^{-2}$, fix monomer head groups in the sandwiched or parallel-displaced configurations while allowing monomer tails to settle.
 - Doing so also mitigates system dependence on initial monomer configuration.
 - The restrained portion of the equilibration scheme is run in the NVT ensemble.
 - Every 50 ps, we reduce the force constants by the square root of its previous value
 - Once the force constant is below $10 \text{ KJ mol}^{-1} \text{ nm}^{-2}$, the restraints are released linearly until there is no more restraining potential.
 - The resulting unrestrained structure is allowed to equilibrate for 5 ns in the NPT ensemble with pressure controlled by the berendsen barostat.
 - Long, NPT equilibration simulations are run for at least 400 ns using the Parrinello-Rahman barostat with a time constant of 10 ps.

- Wet equilibration scheme:
 - In order to create a wet system, we solvated an initial configuration with water using gmx solvate
 - All water molecules placed outside the pore region are removed
 - Waters inside the pore region are randomly removed until the desired concentration of water in the pores is reached.
 - The remainder of the equilibration is the same as the dry system
- In all cases, the v-rescale thermostat was used with tau-t = 0.1

Using an equilibrated structure, a crosslinking procedure was performed in order to match synthetic procedures.

- The purpose of crosslinking is to maintain macroscopic alignment of the crystalline domains, ensuring aligned, hexagonally packed pores.
- For that reason, we are not concerned with replicating the kinetics of the reaction, but instead emphasize the consistency of the final structure with experimental structural data.
- The algorithm was developed based on the known reaction mechanism.
- Crosslinking of this system is a free radical polymerization (FRP) taking place between terminal vinyl groups present on each of the three monomer tails.
- FRPs require an initiator which bonds to the system, meaning new atoms were introduced into the system.
- For simplicity, the initiator was simulated as hydrogen and made present in the simulation by including them in all possible bonding positions as dummy atoms.
- The crosslinking procedure is carried out iteratively.
- During each iteration, bonding carbon atoms are chosen based on a distance cut-off.
- The topology is updated with new bonds and dummy hydrogen atoms are changed to appropriate hydrogen types.
- Head-to-tail addition was the only propagation mode considered due to its dominance in the real system.
- Direction of attack was not considered because the resultant mixture is racemic.

Using equilibrated structures, we carry out various calculations to characterize the system.

- We determine the point at which a system is equilibrated when the distance between pores stops changing

To calculate the equilibrated pore spacing, we measured the distance between pore centers.

- Pore centers were located by averaging the coordinates of sodium ions in their respective pores.
- Statistics were generated using the bootstrapping technique (See Supplemental Information)
 - We are interested in 5 pore-to-pore distances which should all be equal in a perfect hexagonal array, however only 4 distances are independent
 - Each pore spacing has its own trajectory of spacing vs. time. Using data collected after the system is equilibrated, we calculate how long it takes for the data in each of the 5 trajectories to become uncorrelated using `pymbar.timeseries.integratedAutocorrelationTime()`
 - We break the full trajectories down into sub-trajectories based on the maximum autocorrelation time of those found in the previous step.
 - For each bootstrap trial, we recreate an equilibrium trajectory by randomly sampling pore spacings from the sub-trajectories
 - We get an average value for each pore spacing by finding the mean of the bootstrapped data
 - We calculate the overall average as the mean of all bootstrapped pore spacings
 - The uncertainty for each pore spacing is calculated as $\frac{<x> - \bar{x}}{\sqrt{4}}$ where $<x>$ is the average spacing from the bootstrap trial and \bar{x} is the average value of one of the pore spacings.
 - We report the mean of these uncertainties

To quantify the degree of layering and the equilibrium distance between layers in our system, we calculate a spatial correlation function, $g(z)$, measured along the z-axis (perpendicular to the membrane plane).

- To calculate $g(z)$, we binned the z component distances between the center of mass of each component and all others of the same pore over at least 50 ns of equilibrated trajectory and then normalized by the average number density.
- To extract the average distance between layers we applied a discrete fourier transform to $g(z)$ and extracted the highest intensity frequency

Simulated X-ray diffraction patterns were generated based on atomic coordinates for a direct experimental comparison.

- All atomic coordinates were simulated as gaussian spheres of electron density corresponding to each atom's atomic number.
- A three dimensional fourier transform (FT) of the array of electron density results in a three dimensional structure factor which represents the unit cell in reciprocal space. z axis to generate a 2D cross section close to what one would see experimentally.
- We matched experimental 2D WAXS patterns by adjusting the initial spacing between layers and the orientation of the head groups with respect to adjacent layers.

We explored the pore composition by measuring the average number densities of various monomer components with respect to the pore centers.

- The centers of the pores were located using the same method to calculate the equilibrium pore spacing
- We looked at the average number density of sodium ions, aromatic rings and carbon atoms making up the monomer tails
- The radial distance from the pore center of all atoms in each group are binned then normalized by area

We calculated ionic conductivity using two different methods for robustness.

- The Nernst-Einstein relation relates the DC ionic conductivity to ion diffusivity, D , concentration, C , ion charge, q , the boltzmann constant, k_b , and temperature, T :

$$\sigma = \frac{q^2 C D}{k_b T}$$

- Sodium ion diffusion coefficients were found by calculating the slope of the linear region of the z-direction mean square displacement curve as indicated by the einstein relation [?].]
- We looked at the MSD plot to determine where to begin and end a linear fit
- Ion concentration was measured with respect to the entire unit cell.
- The second method, termed the 'Collective Diffusion' model, measures the movement of the collective variable, Q , which is defined as the amount of charge transfer through the system and can be thought to represent the center of charge of the system.
- The conductance, γ of the system can be calculated as:

$$\gamma = \frac{D_Q}{k_b T}$$

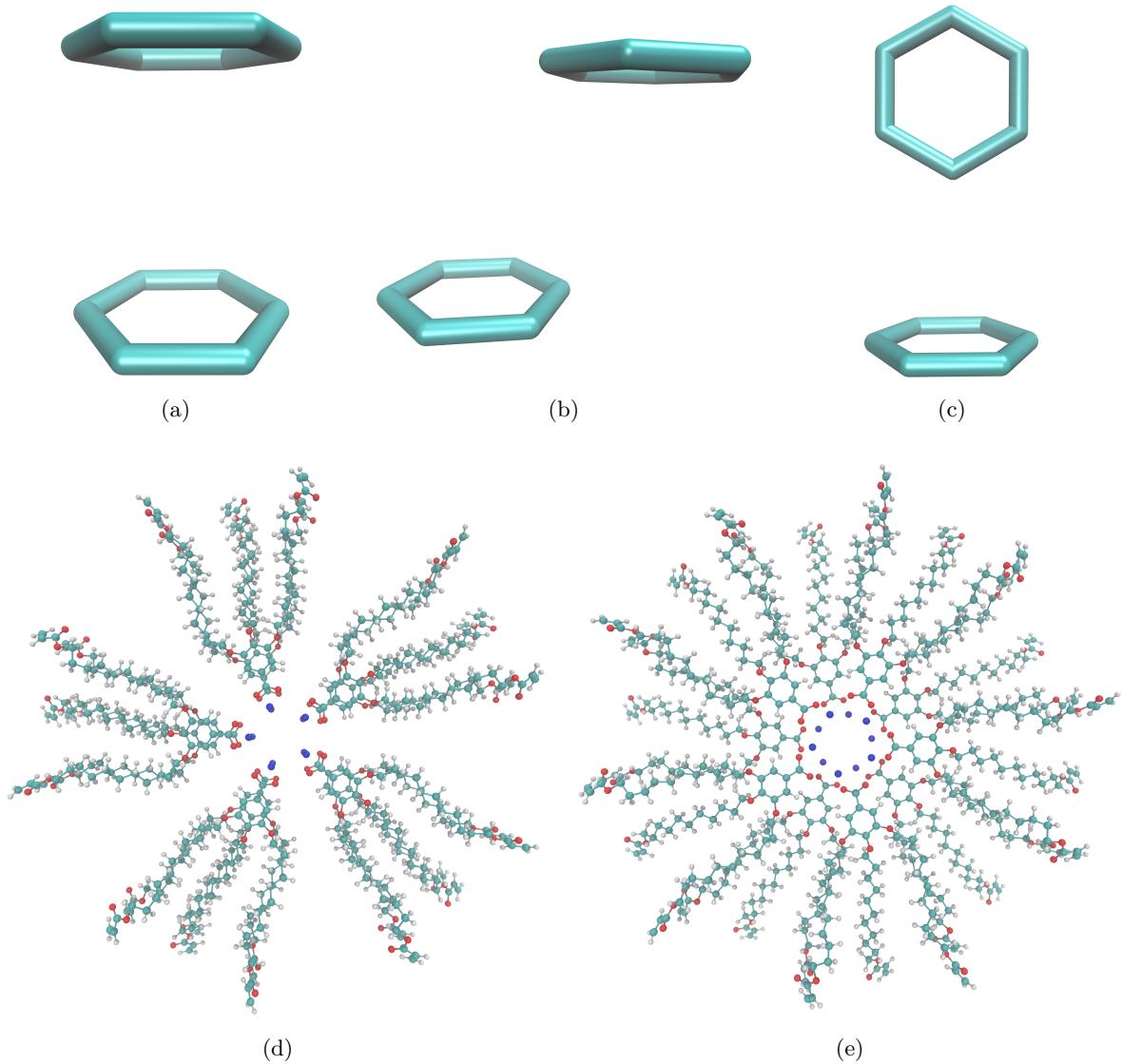


Figure 5: (a) Sandwiched benzene dimers stack 3.8 Å apart. (b) Parallel-Displaced benzene dimers stack 3.4 Å vertically and 1.6 Å horizontally apart. (c) T-shaped benzene dimers stack 5.0 Å apart. (d) Two monomer layers stacked in the sandwiched configuration (e) Two monomer layers stacked in the parallel-displaced configuration

Conversion to ionic conductivity is achieved by multiplying by channel length and dividing by the membrane cross sectional area.

- D_Q is the diffusion coefficient of the collective variable Q. It can be calculated using the einstein relation.
- A full derivation of the model can be accessed elsewhere [?].

Results and Discussion

Determination of Nanoscopic Structural Details

We will now address the questions raised in the introduction in the order that they were asked.

The simulations best support a model built with 5 monomers per layer based on the measured pore-to-pore distances. To discern the composition of the monomer layers, addressing (1), we ran simulations of systems created with 4 - 8 monomers per layer.

- We created systems in both the parallel displaced and sandwiched configurations
- All systems were prepared according to the dry equilibration procedure
- All systems are stable after 400 ns of simulation.
- Figure 7 shows the pore spacing for all systems tested.
- Systems built with 5 monomers in each layer equilibrate to a pore spacing that is most consistent with the experimental value of 4.12 nm derived from SAXS measurements (Figure 3a).
- The remainder of this discussion will focus on the analysis of systems built with 5 monomers per layer.

We learned that layers are well-defined and persistent, answering (2) We verified our conclusion by plotting the pair correlation function, $g(z)$ calculated between atoms along the length of the pores (Fig 6).

- We measured $g(z)$ with respect to aromatic rings in the head groups and, separately, with respect to carbon atoms on the alkane chains.
- Using the head groups as reference, we see that sandwiched configuration layers stack 4.29 Å apart while parallel displaced configuration layers stack 4.37 Å apart.
- Using the alkane chain carbons as reference we see a layer spacing of 4.39 Å and 4.38 Å in the sandwiched and parallel displaced configurations respectively.

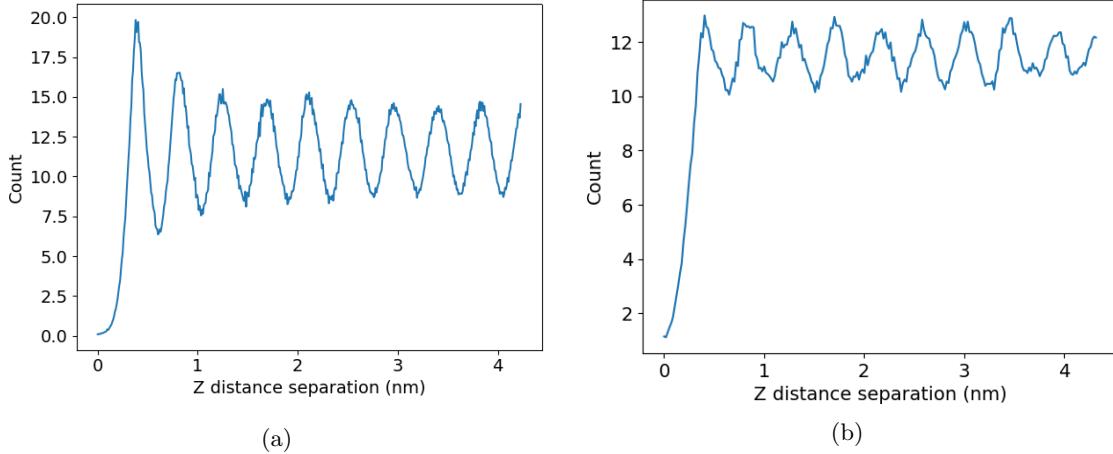


Figure 6: Pair distribution functions of aromatic carbons for the (a) 5 monomer per layer, sandwiched and (b) 5 monomer per layer, parallel displaced configurations. Clear periodic maxima in the z probability density indicate distinct layers. The magnitude of the spikes with respect to the average suggest that the 5 monomer per layer, sandwiched configuration possesses a higher degree of layer partitioning.

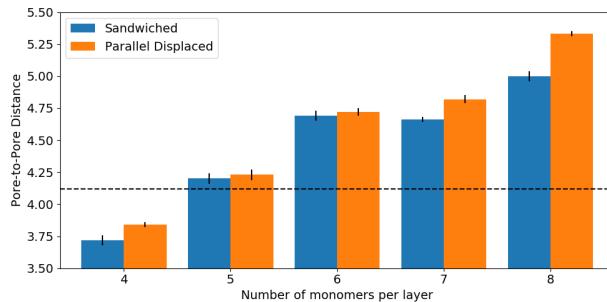


Figure 7: The pore spacing (given in nm) of the model increases as number of monomers in each layer increases. The pore spacing of a system starting in the sandwiched configuration is systematically lower than that started in an offset configuration. Systems built with 5 monomers per layer in a parallel displaced configuration result in a pore spacing closest to the experimental 4.12 nm

- The power spectrums used to calculate the reported layer spacings are given in the Supplemental Information.

We answer question (3) by simulating X-ray diffraction patterns produced from equilibrated MD trajectories.

- Systems of 5 monomers per layer in the parallel displaced and sandwiched configurations were tested.
- Simulated patterns were generated using portions of simulation trajectory after equilibration.
- The patterns for both structures are shown and compared to experiment in Figure ??.

The parallel displaced configuration results in a simulated XRD pattern with the closest match to experiment.

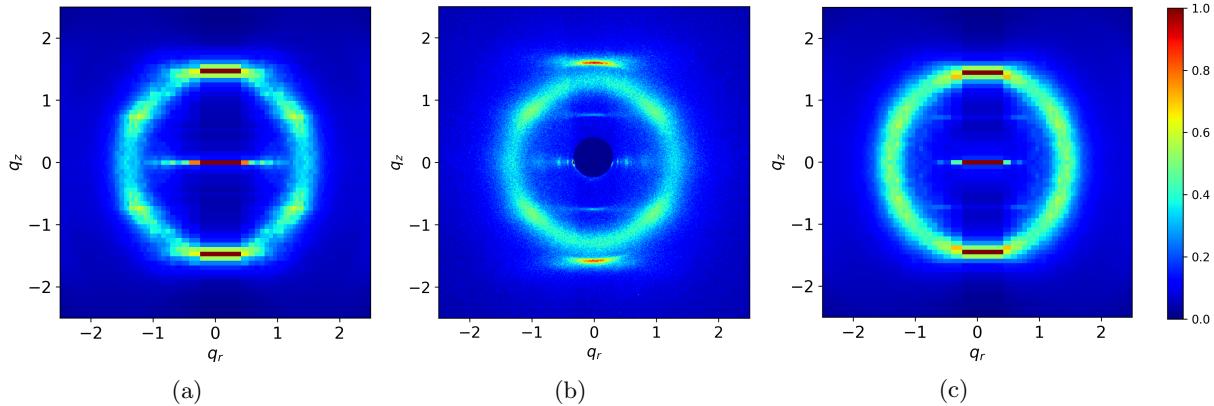


Figure 8: The simulated X-ray diffraction patterns for simulations run in the sandwiched (a) and parallel-displaced (c) configurations are compared to experiment (b). The parallel displaced configuration is the only one that exhibits all major reflections of interest to some degree

- It produces the only pattern that exhibits all major reflections
- R-alkanes, R-pores and R- π appear as they do in the sandwiched configuration.
- R-spots appears, however with a lower intensity relative to R-alkanes when compared to the sandwiched configuration.
- R-helix appears apparently due to the parallel displaced aromatic rings

Simulated XRD of the sandwiched configuration contains all experimental features except for R-helix.

- R-alkanes and R-pores appear in the expected locations.
- R- π is also present, intersecting R-alkanes at a lower q value than in experiment. The rings prefer to stack $\approx 4.2\text{\AA}$ apart as opposed to 3.7\AA .
- Most notably, R-spots appears in the expected location, which suggests that there is something intrinsic to closer packing that gives rise to such features.

R-spots, which appears in both simulated XRD patterns, is a result of the way alkane tails pack together.

- Previously, the spots in the diffraction pattern had been explained as the product of tilted alkane chains.
- We measured the tilt angle of the alkane chains and showed that our system equilibrates to an average tilt angle close to zero degrees 9
- To understand the origin of the spots, we determined which atoms gave rise to the feature

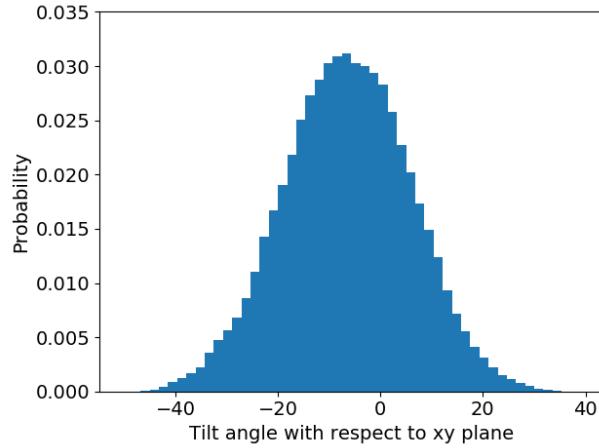


Figure 9: The tilt angle distribution of alkane chain tails with respect to the membrane plane indicates an average tilt angle near 7° which is far from the 37° tilt angle previously used to explain R-spots

- Since R-spots are present as higher intensity spots within R-alkanes, it is likely that the spots arise as a consequence of the tails.
- By removing atoms from the trajectory and simulating a diffraction pattern, we were able to isolate the cause of the spots to the tails (Figure 10).
- Since the tails stay nearly flat, we plotted the centroids of the tails and measured the angle between each centroid and its nearest neighbors with respect to the plane of the membrane (Figure ??).
- We see distinct peaks in the distribution of these angles (Figure 11)

The peaks in the tilt angle distribution are consistent with the location of R-spots.

- The peaks of interest in Figures 11a and ?? are located at $\pm 33^\circ$ which is the same location where the highest intensity of spots are located on the simulated patterns
- We confirmed this conclusion by radially integrating the 2D WAXS pattern for q values between 1.57 and 1.4 (real space between 4 and 4.5 Å)
- We observe that distinct peaks appear c.a. 30° , in close agreement with the previously measured angle distribution (11b 11d)
- We performed the same integration on the raw experimental data and found the angle at which R-spots reaches its highest intensity to be $\pm 37^\circ$ which is a reconcilable difference with our simulated results.

The z -direction correlation functions show that layers in our model prefer to stack further apart than the 3.7 Å suggested by experiment. We attempted equilibration with layers stacked 5.0 nm apart and to our

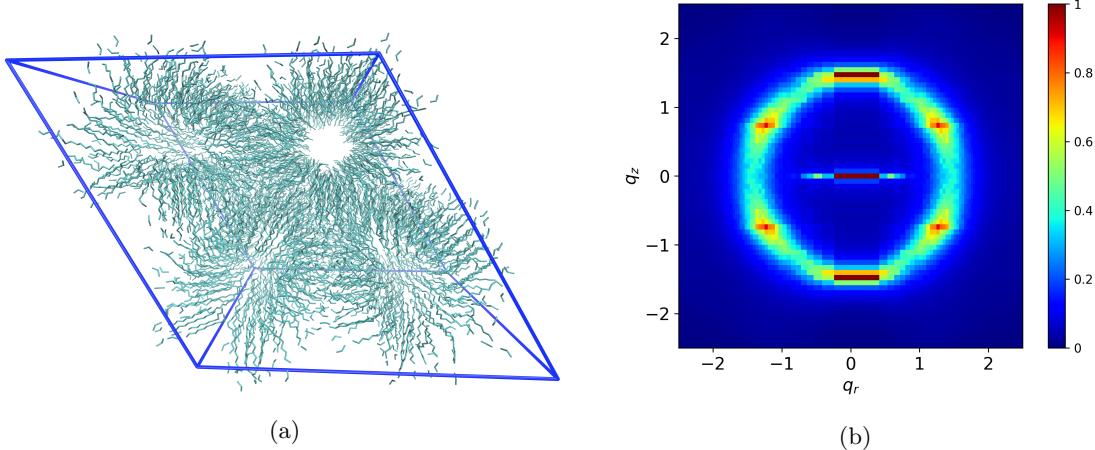


Figure 10: (a) All atoms except carbon atoms making up the tails are removed from the trajectory. (b) The simulated diffraction pattern of the tail-only trajectory still shows R-spots

surprise, we observed long-term stability of a qualitatively different configuration suggesting that we have found more than one metastable free energy basin, corroborating 4.

- Equilibrated systems built according to the 3.7 \AA layer spacing implied by $R-\pi$ are characterized by a defined, cylindrical and open pore structure.
- We will refer to this large set of configurations, with an open pore, as the ordered pore basin open pore (Figure 12a).
- Simulations of systems built with layers stacked 5 \AA apart results in a pore structure characterized by high radial disorder, while still maintaining partitioning between hydrophobic and hydrophilic regions.
- This will be called the disordered pore basin (Figure 12b).
- This LLC membrane may exist in at least two metastable states.
- The distinct difference in pore structure exhibited by each phase will likely lead to different transport mechanisms.
- Because the pore structure will vary between each basin, understanding which phase exists experimentally is necessary in order to ensure we are studying the system which actually dominates.

In order to address 5 and to further understand the structural difference between the ordered pore and disordered pore basins, we plotted the number of densities of heavy atoms in the head group and tail regions of the monomers in addition to the sodium ions. (See Supplemental Information for definition of head group and tail regions)

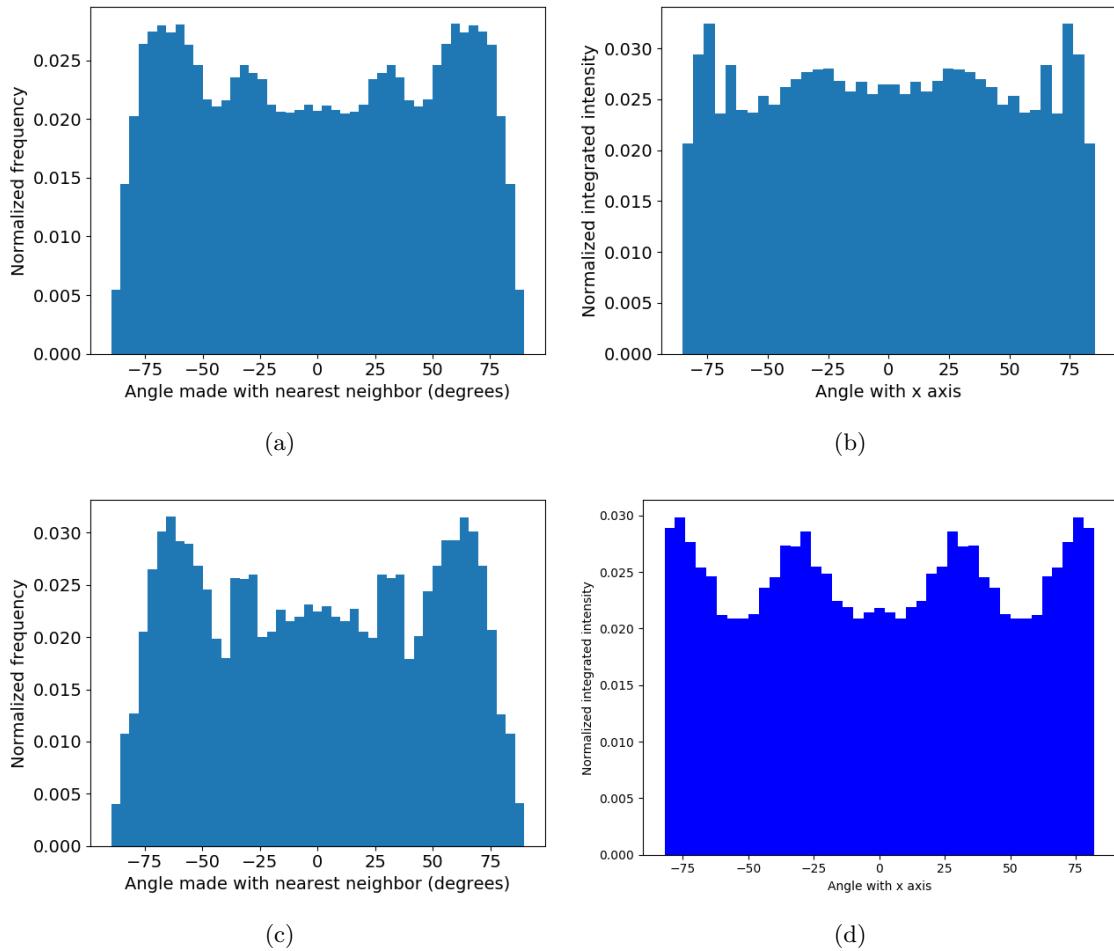


Figure 11: The distribution of angles w.r.t. the xy plane between alkane chain tail centroids and nearest neighbor centroids for equilibrated parallel displaced (a) and sandwiched (c) configurations. The same peaks are visible when the 2D simulated diffraction data is radially integrated in the R-alkanes region, (b) and (d) respectively.

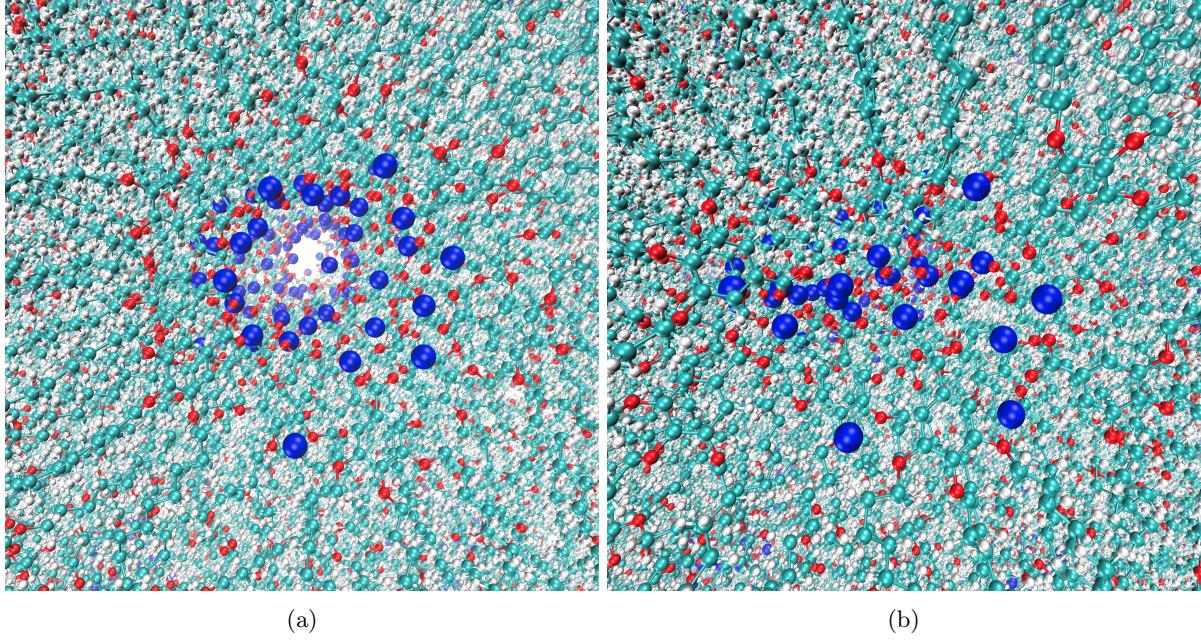


Figure 12: From a qualitative standpoint, the ordered pore basin (a) is characterized by a hollow cylindrical pore while pores in the disorder pore basin (b) are characterized by a higher degree of radial disorder

- For the hydrophilic region, we used the carbon atoms making up the aromatic ring
- For the hydrophobic components we used all carbon atoms of the monomer tails

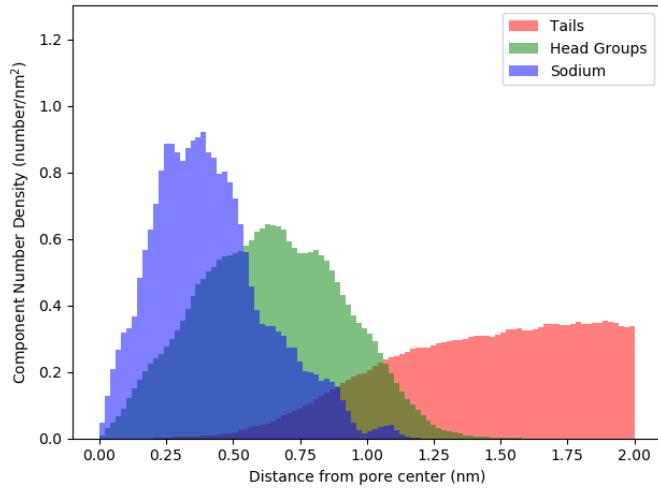
We answer (6) and further support (3) by preparing parallel displaced and sandwiched configurations according to the wet equilibration procedure.

- We do not have an experimental measurement of water in the pores
- We tested a range of water concentrations from 0.5 to 5 percent
- Our lower bound represents a system with on average 2 water molecules for each monomer layer.
- Figure ?? shows results for each configuration.

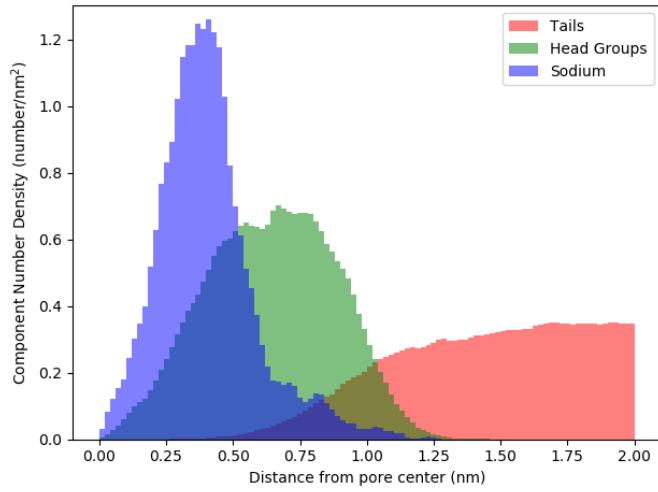
In all cases, water disrupts structuring of the model.

- The plots in Figure ?? are normalized to have the same colorbar as the dry systems.
- The intensity of the reflections decrease when water is added to the system.
- In systems built with 5 weight percent water, R- π and R-spots become nearly indistinguishable from R-alkanes.

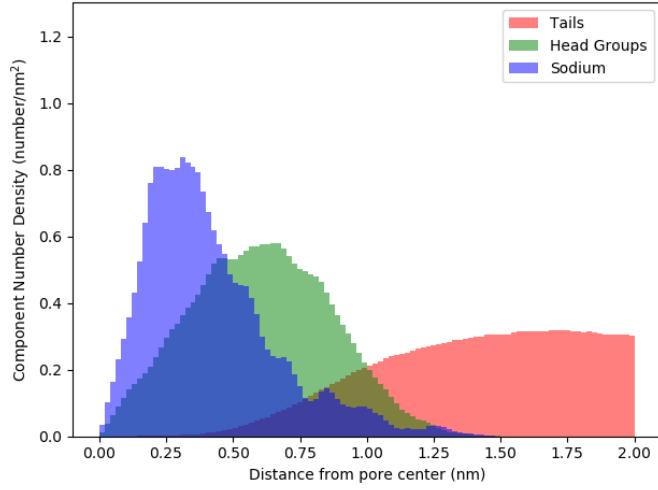
Water is not needed to obtain and simulate the correct equilibrium membrane structure.



(a)



(b)



(c)

Figure 13: (a) Parallel Displaced (b) Sandwiched (c) “Disordered”

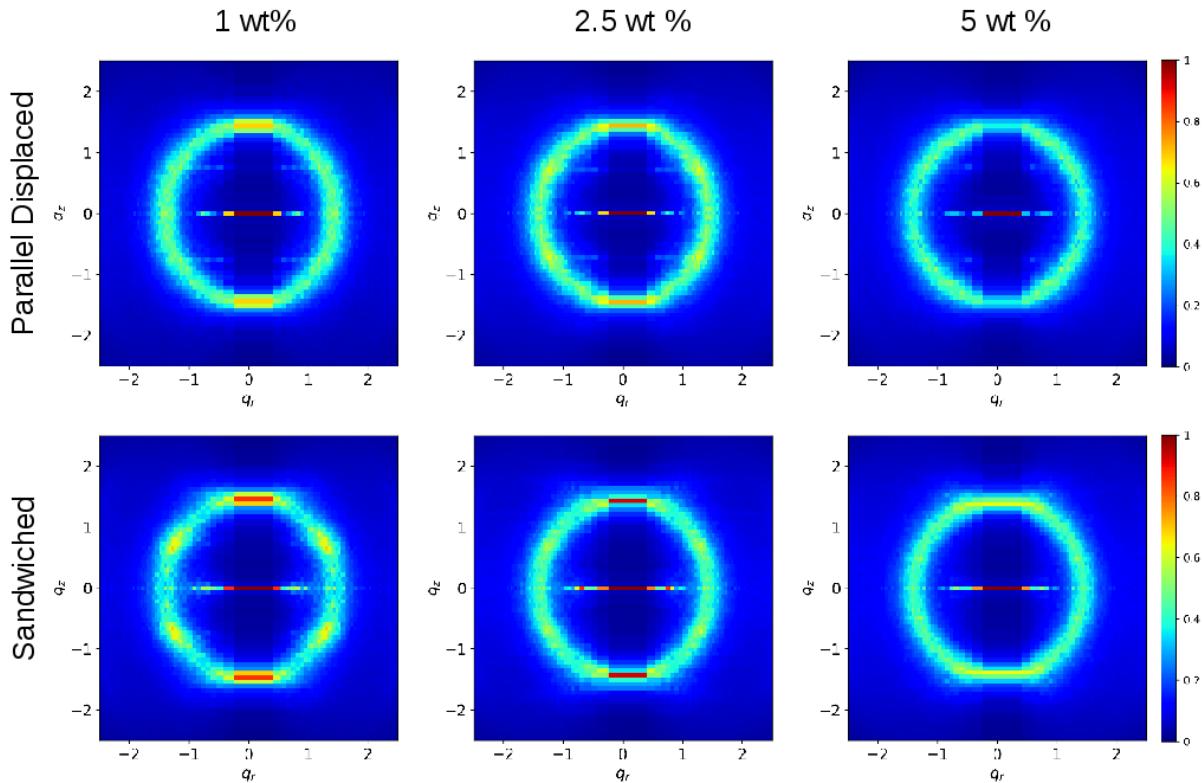


Figure 1: Simulated diffraction patterns generated from trajectories of systems built in the parallel displaced and sandwiched configurations as a function of water content shows weaker reflections than the dry system. All plots are scaled to use the same colorbar as their dry system counterpart

- We note that water absorbed during membrane synthesis may be present in far lower concentrations than those tested here.
- We do not eliminate the possibility that water is necessary in order to drive self-assembly. Studying the mechanisms of self assembly is beyond the scope of this work.
- However, once the system has formed the Col_h phase, adding water only drives disorder in the structure
- In the equilibrium configuration, if water exists, it is confined to the pore region where there is no driving force for aggregation of water molecules.
- In the case of trace water, water molecules will be too sparse to form a hydrogen bonding network.

Ionic conductivity calculation

We use the equilibrated parallel displaced system to calculate ionic conductivity since its structure is the closest match to experiment. The model gives reasonable estimates of ionic conductivity when compared to experiment.

- Calculated values of ionic conductivity obtained using the Nernst Einstein relation and Collective Diffusion model are compared in Figure 15.
- The two methods agree with each other within error, although the uncertainty obtained using the Collective Diffusion model is much higher.
- Much longer simulations are needed to lower the uncertainty, however it is not feasible to do so with a large system.
- For this reason we will likely only use the Nernst Einstein relation in future calculations of this type.

The calculated values of ionic conductivity are higher than experiment by an order of magnitude.

- One can justify the reason for this result by considering the real system studied experimentally
- The ionic conductivity measurement to which we are comparing was done on a 80 μm thick film, nearly 10,000 times thicker than our simulated system.
- The thick film likely has defects leading to non-contiguous pores and imperfect alignment.
- The 2D SAXS pattern shows the small degree of tortuosity present in the system (14)
- It has been shown that there is a large dependence of Ionic conductivity on the alignment of the pores

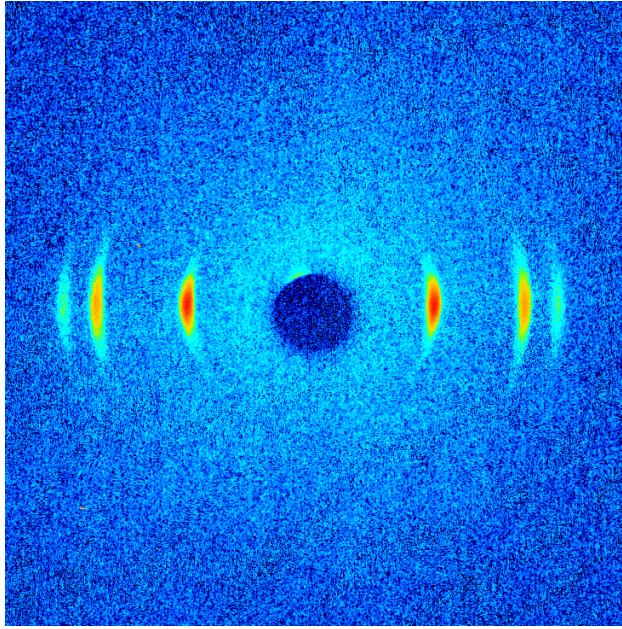


Figure 14: The experimental two dimensional SAXS pattern exhibits reflections resulting from highly aligned pore regions. A perfectly aligned system would exhibit sharp dots at the same locations of the reflections shown. An isotropically aligned system the reflections would appear as rings centered about the origin

- The ionic conductivity of an unaligned film is 85 times lower than that of a nearly aligned film referenced here
- It is not unreasonable that a thin, perfectly aligned film would have a value of ionic conductivity in agreement with our model

Implementation of the crosslinking algorithm

We applied the crosslinking algorithm to the equilibrated parallel-displaced configuration with 5 monomers per layer.

- We reach 95 % conversion of terminal vinyl groups
- The distance between pores shrinks by 0.4\AA after the system is crosslinked
- We simulated the crosslinked system in the NPT ensemble for 100 ns
- Major features are still present in the X-ray diffraction, however at lower intensities 16a
- The ionic conductivity is lower in the crosslinked system 16b

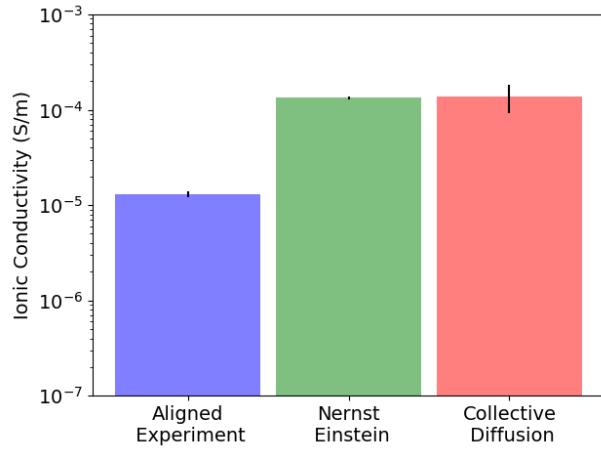


Figure 15: Calculated ionic conductivity using the Nernst-Einstein relation and Collective Diffusion model agree with error. Both methods give calculated values of ionic conductivity an order of magnitude higher than the experimental value

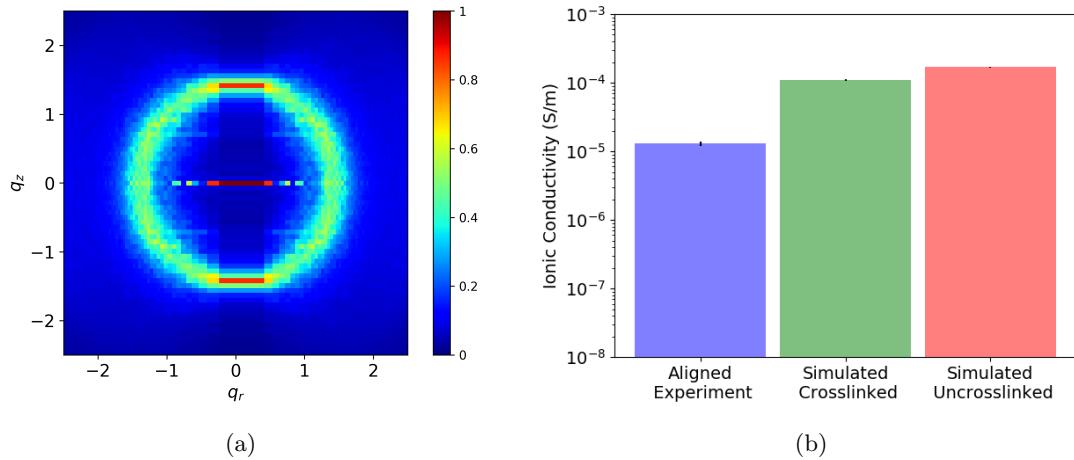


Figure 16: (a) Reflections produced by the crosslinked configuration fade relative to the uncrosslinked system. The colorbar shown is the same used for the uncrosslinked system. (b) The ionic conductivity is smaller relative to the uncrosslinked system, but still much larger than the experimental value

Conclusion

We have used a detailed molecular model of the Col_h phase formed by NA-GA3C11 in order to study the nanoscopic structure.

- While there have been efforts to model formation of various liquid crystalline phases with molecular dynamics, to our knowledge there have been no studies which attempt to examine their structure with the same level of detail presented here.
- Evidence strongly supports that layers are composed of 5 monomers
- We have confirmed that monomers stay partitioned in layers.
- We have explored the affect of two different π - π stacking modes on the equilibrated membrane strucure.
- Simulated diffraction patterns generated from MD trajectories suggest that the parallel-displaced configuration produces a structure with the closest match to experiment.
- Finally, water is not needed to maintain membrane structure our data shows that it actually disorders the pore
- Now that we have a good idea of what the membrane structure should look like, we will evaluate transport of various solutes within the system.
- We will apply the knowledge gained from this study in order to suggest improvements to the existing system as well as to evaluate new unsynthesized LLC systems.