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A Molecular View of the Cholesterol Condensing Effect in DOPC Lipid Bilayers

Mohammad Alwarawrah, Jian Dai, and Juyang Huang*

Department of Physics, Texas Tech University, Lubbock, Texas 79409

Received: February 15, 2010; Revised Manuscript Received: May 5, 2010

The condensing effect of cholesterol in dioleoylphosphatidylcholine (DOPC) lipid bilayers was systematically investigated via atomistic molecular dynamics (MD) simulation. Fourteen independent 200 ns simulations, spanning the entire range of cholesterol mole fraction (x_c) in DOPC bilayers (i.e., from $x_c = 0$ to 0.66), were performed at 323 K. The molecular areas occupied by DOPC and cholesterol at different distances from the bilayer center were analyzed using a slicing method based on the VDW radii of atoms. Curiously, while the average area per lipid and the cholesterol tilt angle, with respect to the bilayer normal, both show monotonic decreases as x_c increases, the average bilayer height shows a significant decrease for $x_c > 0.35$, following an initial increase. The calculated partial-specific areas of lipids clearly show the condensing effect of cholesterol. The VDW areal analysis showed that the condensing effect is limited only to the cholesterol sterol ring region, where the acyl chains of DOPC are severely compressed by cholesterol. As x_c increases, the headgroups of DOPC gradually expand along the bilayer–aqueous interface to occupy more lateral area. Thus, it confirmed a key prediction of the umbrella model. At high cholesterol mole fractions, the calculated area per DOPC and area per cholesterol using some existing methods showed an inconsistent result: both increase while the overall area per lipid decreases. The inconsistency stems from the problematic assumption that cholesterol and DOPC have cylindrical shape and the same height. Our results showed that the total area of a PC/cholesterol bilayer is primarily determined by the molecular packing in the cholesterol sterol ring region. An alternative analysis of area per molecule within this region is proposed, which takes into account the cholesterol tilt angle and the practical incompressibility of cholesterol sterol rings. The new calculation shows that the majority of the area lost due to the cholesterol condensing effect is taken from PC molecules.

Introduction

The lipid bilayer is the fundamental structural scaffold of the cell plasma membrane. Phospholipids and cholesterol are the most abundant lipid constituents within these membranes, and the molecular interactions between the two molecules can greatly affect the biochemical and biophysical properties of a lipid bilayer. In the past decade, molecular dynamics (MD) simulation has been widely utilized in the study of cholesterol-containing lipid bilayers at the atomistic scale.^{1–9} One of the most important effects of cholesterol on lipid membranes is the cholesterol condensing effect: The surface area of a cholesterol-containing lipid bilayer is less than the sum of areas of the individual bilayer components.^{10–12} The cholesterol condensing effect can be quantitatively described in terms of the average area per molecule, which reflects the molecular organization within a lipid bilayer. A seemingly straightforward question that interests many researchers is “what is the average area occupied by a phospholipid or by a cholesterol in a lipid bilayer?” For a pure PC bilayer simulation, the answer can be easily obtained by dividing the total area of the bilayer in the simulation box by the total number of PC molecules; however, there is no simple answer for a PC bilayer containing cholesterol. A number of methods have been developed for the purpose of calculating the area per PC and area per cholesterol from an MD simulation.^{13–16} These methods have been applied to various phospholipid bilayers and to different cholesterol mole fractions as well as at different system temperatures. One of these approaches was proposed by Hofsäuss et al.,¹⁵ in which the

volume of cholesterol was considered to be constant. Additionally, phospholipids and cholesterol were assumed to have the same height in a lipid bilayer. Edholm and Nagle improved the volumetric component of the analysis but kept the same height assumption.¹³ Another method applied Voronoi analysis in two dimensions to find average areas per PC and per cholesterol as a function of x_c ;¹⁶ however, it has been suggested that this method may intrinsically overestimate the area occupied by smaller molecules and underestimate the area occupied by larger molecules.¹⁷ Finally, Falck et al. divided a lipid bilayer into horizontal slices and analyzed the molecular area within each slice. They showed that there is indeed a z -dependence of the molecular areas when examined at an atomic level.¹⁴

Edholm and Nagle recognized the existence of arbitrary features in the above methods and introduced the formalism of partial-specific area.¹³ Partial-specific area is defined as the increase of total bilayer area by adding one more lipid molecule at a particular bilayer composition. This method has been applied to DPPC/cholesterol, DOPC/cholesterol, and POPC/cholesterol bilayers.^{5,13} For example, in a DPPC bilayer containing 10 mol % of cholesterol, the calculated partial-specific area of cholesterol was about -0.19 nm^2 . The negative value results from the well-known cholesterol condensing effect. Therefore, a systematic analysis of bilayer area can shed light on the molecular origin of the cholesterol condensing effect.

In this study, the cholesterol condensing effect in DOPC bilayers was systematically investigated using atomistic MD simulation. Fourteen independent 200 ns simulations, spanning the entire possible range of cholesterol mole fraction in DOPC bilayers, were performed at 323 K. The molecular areas occupied by DOPC and cholesterol at different distances from

* To whom correspondence should be addressed. E-mail: juyang.huang@ttu.edu. Phone: (806) 742-4780. Fax: (806) 742-1182.

the bilayer center were analyzed using a slicing method based on the VDW radii of atoms. The analysis revealed that as x_c increases, the areas occupied by PC and cholesterol both decrease in the cholesterol sterol ring region; however, PC headgroups actually occupy more lateral area at the bilayer–aqueous interface. Thus, it confirmed an important prediction of the umbrella model.¹⁸ The reduction of area per cholesterol in the cholesterol sterol ring region is primarily due to the decrease of cholesterol tilt angle with respect to the bilayer normal, while the reduction of area per PC in the same region is due to the compression of acyl chains by the cholesterol sterol rings. The area per molecule was calculated using the method of Hofsäss et al. as well as the method of Edholm and Nagle over the entire range of cholesterol mole fraction. Interestingly, in the high cholesterol region, which has been rarely studied by MD simulation, the calculated area per cholesterol as well as the area per PC both increases, while the average area per lipid decreases. This inconsistency stems from the built-in assumption by both methods that PC and cholesterol were of cylindrical shape and of the same height. Our analysis shows that total area of a PC/cholesterol bilayer is primarily determined by the packing of cholesterol and PC in the cholesterol sterol ring region. We propose an alternative method for calculating area per PC and area per cholesterol in this region that gives a more informative description of the cholesterol condensing effect at the molecular level.

Simulation Methods

Fourteen independent 200 ns simulations were performed at 323 K, with a time step of 2.0 fs. In the simulations of DOPC bilayers containing 0, 1.95, 5.08, 10.16, 14.84, 20.31, 25, 30.08, 35.16, 39.84, 44.92, and 50 mol % of cholesterol, the total number of lipids (i.e., DOPC + cholesterol) was kept at 512. For the systems containing 57% and 65.63% of cholesterol, the total numbers of lipids were 200 and 128, respectively. All systems contained 28.625 water molecules for each lipid. The phospholipid force field was from Berger et al.,¹⁹ and the force field for cholesterol was based on the GROMOS force field from Holtje et al.²⁰ The simple point charge (SPC) model was used for water molecules. The force field parameter files for phospholipids were obtained from Dr. Peter Tieleman's Web site, and the file for cholesterol was obtained from the GROMACS Web site. The MD simulations were performed using GROMACS 4.0.5,²¹ and the analysis was done using the GROMACS tools as well as our own programs. The LINCS algorithm was used to keep the lengths of all bonds constant. The cutoff distance for Lennard-Jones interactions and electrostatic interactions were both set at 1.0 nm. Long-range electrostatic interactions were handled with the particle mesh Ewald (PME) method.²² All systems were run for 200 ns in the NPT ensemble using a Berendsen thermostat and barostat with a coupling time constant of 0.1 and 1.0 ps, respectively. The pressure normal and parallel to the bilayer were coupled separately at 1 bar.

The pure DOPC bilayer was constructed by first obtaining the PDB coordinates file of a single DOPC molecule from the Dundee PRODRG2 Server.²³ A single DOPC was placed inside a solvent box, and a simulation was performed to relax the lipid. Afterward, bilayers of 128 and 200 DOPCs were constructed from this relaxed DOPC. Initial equilibration simulations of the two systems consisting of fully solvated 128 and 200 DOPC lipids were performed for 200 ns. Following this, the GROMACS²¹ utility "genconf" was employed to replicate the system of 128 DOPC and thus generate a new starting

configuration consisting of 512 DOPC lipids. To insert cholesterol molecules into the bilayers, a number of DOPCs in each leaflet were randomly selected and replaced with cholesterol. During the replacement, the COM of cholesterol coincided with the COM of the replaced DOPC, and the cholesterol was rotated by a random angle in respect to its principal axis. The VMD program was then used to remove any bad links or overlaps that could happen after cholesterol insertion. In constructing the systems with 57% and 65.63% cholesterol from pure DOPC bilayers, DOPCs were replaced by cholesterol molecules gradually: after replacing about 20 DOPC molecules in each leaflet, the systems were equilibrated for 50 ns before the next replacement. All initial structures went through energy minimization using both the steepest descent and the conjugate gradient algorithms; each ran for 10 000 steps to eliminate any bad contacts between atoms.

Area per Molecule Calculation

Method 1. The first method was proposed by Hofsäss et al.¹⁵ In this method, the volume of cholesterol was considered to be a constant ($v_{\text{CHOL}} = 0.593 \text{ nm}^3$), independent of cholesterol concentration. The volume per PC was calculated by subtracting the volumes of water and cholesterol from the total volume of the simulation box:

$$v_{\text{DOPC}}(x_c) = \frac{V(x_c) - N_w v_w - x_c N_{\text{lipid}} v_{\text{CHOL}}}{(1 - x_c) N_{\text{lipid}}} \quad (1)$$

where N_{lipid} is the total number of lipids in the simulation box, N_w is the number of water molecules, and $V(x_c)$ is the volume of the simulation box. v_w is the volume of a water molecule; this value ($0.031 546 \text{ nm}^3$) was obtained from a separated simulation of pure water at 323 K under the same conditions. The average thickness of the lipid bilayer $h(x_c)$ was calculated using the equation¹⁵

$$h(x_c) = \frac{V(x_c) - N_w v_w}{A(x_c)} \quad (2)$$

The area occupied by a PC and by a cholesterol molecule in the bilayer can be written in terms of the volume and thickness as

$$a_{\text{DOPC}}(x_c) = \frac{2v_{\text{DOPC}}(x_c)}{h(x_c)} \quad \text{and} \quad a_{\text{CHOL}}(x_c) = \frac{2v_{\text{CHOL}}}{h(x_c)} \quad (3)$$

An important assumption, which is built into eq 3, is that PC and cholesterol molecules are assumed to have the same height.

Method 2. Edholm and Nagle improved the volumetric part of the analysis of method 1:¹³ $v_{\text{DOPC}}(x_c)$ and $v_{\text{CHOL}}(x_c)$ are obtained by fitting volume per lipid vs x_c using the linear equation

$$v(x_c) = (1 - x_c)v_{\text{DOPC}}(x_c) + x_c v_{\text{CHOL}}(x_c) \quad (4)$$

Assuming a constant volumetric ratio of cholesterol to PC $f(x_c) = v_{\text{CHOL}}(x_c)/v_{\text{DOPC}}(x_c)$, Edholm and Nagle calculated area per PC and area per cholesterol using

$$a_{\text{DOPC}}(x_c) = \frac{a(x_c)}{1 - x_c(1 - f(x_c))} \quad \text{and} \quad a_{\text{CHOL}}(x_c) = f(x_c)a_{\text{DOPC}}(x_c) \quad (5)$$

where $a(x_c)$ is the area per lipid. Note that eq 5 also contains the built-in assumption that PC and cholesterol have the same height.

Partial-Specific Areas. Edholm and Nagle proposed the formulism of partial-specific molecular area for PC and cholesterol.¹³ The partial-specific areas a_i for a bilayer consisting of $i = 1, \dots, m$ types of molecules is defined as

$$a_i(X) = \left(\frac{\partial A(N_1, \dots, N_m)}{\partial N_i} \right)_{N_j \neq i} \quad (6)$$

where A is the total area, N_i is the number of i type lipid per leaflet, X indicates a set of m lipid mole fractions (x_1, \dots, x_m), and the partial derivative is taken with N_j constant for all $j \neq i$. A convenient graphical way to obtain the partial-specific areas is first plotting $a(x_c)/(1 - x_c)$ vs $x_c/(1 - x_c)$ using

$$\frac{a(x_c)}{1 - x_c} = \frac{A(x_c)}{N_{\text{DOPC}}} = a_{\text{DOPC}}(x_c) + \frac{x_c}{1 - x_c} a_{\text{CHOL}}(x_c) \quad (7)$$

then $a_{\text{CHOL}}(x_c)$ can be estimated directly from the slope of the tangent and $a_{\text{DOPC}}(x_c)$ can be estimated from the intercept of the tangent at $x_c = 0$.

Results and Discussion

The chemical structures of cholesterol and DOPC are shown in Figure 1. The orientation of a cholesterol molecule is characterized by a vector connecting C₂₁ and C₅ atoms of the cholesterol.

Area per Lipid. Area per lipid, a_{pl} , is an essential parameter for describing the state of molecular packing within a lipid bilayer. In an MD simulation, it is calculated by dividing the total area of the simulation box in x - y plane by the total number of lipid molecules in one leaflet of the bilayer. Table 1 lists the average area per lipid for 14 simulations that covered the entire possible range of cholesterol mole fraction in DOPC bilayers. The maximum solubility of cholesterol in DOPC bilayers determined by light scattering, fluorescence anisotropy, and cholesterol oxidase activity assay was 0.66.^{24,25} Some lower values have also been reported, but these likely resulted from problematic sample preparation.^{26,27} In most simulations, the area per lipid reached a stable value quickly (data not shown); however, in certain simulations with high cholesterol content, a stable value was not reached until about 150 ns because replacing PCs with a large number of cholesterol significantly perturbed the system. Slow diffusion of molecules at high cholesterol content also slowed the re-establishment of equilibrium.¹⁵ Therefore, the average values of area per lipid (Table 1 and Figure 3) were calculated using only the last 50 ns of the 200 ns simulations.

Volume per Lipid and Bilayer Thickness. Figure 2 shows volume per lipid $v_{\text{pl}}(x_c)$ and the bilayer thickness $h(x_c)$ as functions of cholesterol mole fraction. Volume per lipid was calculated by subtracting the volume of water from the volume of the simulation box and then dividing it by the total number of lipids. The relation between $v_{\text{pl}}(x_c)$ and x_c is almost linear,

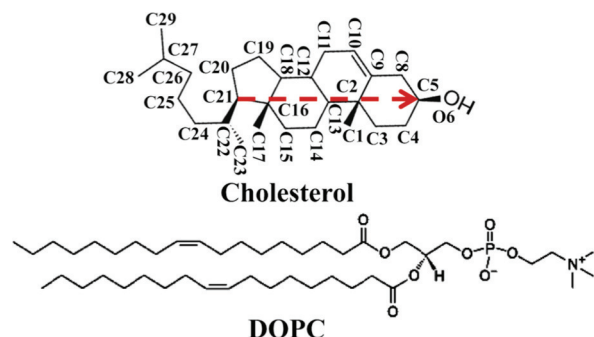


Figure 1. Chemical structures of cholesterol and DOPC. The vector connecting C₂₁ and C₅ atoms of cholesterol (red arrow) is used to represent the orientation of a cholesterol molecule, and the angle between this vector and the bilayer normal is defined as the cholesterol tilt angle.

which is consistent with the measured volumes of DOPC/cholesterol bilayers.²⁸ The bilayer thickness was calculated using eq 2. An important feature of Figure 2 is that the bilayer thickness initially increases as x_c increases, but reaches a maximum around $x_c = 0.35$, and then shows a gradual decrease at higher x_c values. From x_c of 0.4 to 0.66, the total decrease of bilayer thickness is about 0.4 nm (or 0.2 nm per leaflet). As to be discussed later, the decrease of bilayer thickness at high cholesterol mole fraction is due to the movement of DOPC headgroups toward the bilayer–aqueous interface in order to cover more cholesterol.

Partial-Specific Areas. Figure 3 shows the area per lipid and partial-specific area of the DOPC and cholesterol. Since the cross-sectional area of cholesterol is less than that of a DOPC, as expected, area per lipid shows a *monotonic* decrease as a function of cholesterol mole fraction. Intriguingly, partial-specific area of cholesterol, $a_{\text{CHOL}}(x_c)$, is negative for $x_c < 0.2$. By definition, $a_{\text{CHOL}}(x_c)$ is the change of total bilayer area at the bilayer composition x_c by adding one more cholesterol molecule to the bilayer, while keeping all other parameters constant. Therefore, at low cholesterol concentrations, adding cholesterol to DOPC bilayers actually causes a *net reduction* of total bilayer area. The negative partial-specific area of cholesterol was first reported in DPPC(di16:0PC)/cholesterol systems by Edholm and Nagle, who interpreted it as a manifestation of the condensation effect of cholesterol on the surrounding lipids.¹³ We believe this to be the correct interpretation. Experimentally, a negative partial-specific area of cholesterol has been observed in DPPC/cholesterol monolayers at the air–water interface.²⁹ Although there are some important differences between a lipid monolayer and a lipid bilayer, interestingly, the changing manner of the partial-specific areas of both the cholesterol and phospholipid in lipid monolayers is quite similar to that calculated for DOPC lipid bilayers.

Cholesterol Tilt Angle. Figure 4 shows the distributions of cholesterol tilt angle at some selected cholesterol mole fractions. The tilt angle is defined as the angle between the vector connecting C₂₁ and C₅ atoms of cholesterol (see Figure 1) and the bilayer normal. As shown in Figure 4A, the distributions of cholesterol tilt angle are quite broad at low cholesterol concentrations. As x_c increases, the distributions become sharper and the peaks also shift toward a smaller angle. Figure 4B shows the average cholesterol tilt angle as a function of x_c . The average cholesterol tilt angle monotonically decreases from 29.7° at x_c of 0.0195 to 10.7° at x_c of 0.6563. Thus, cholesterol becomes more orientationally ordered and more parallel to the bilayer normal at higher cholesterol mole fractions.

TABLE 1: Area per Lipid ($a_{\text{pl}}(x_c)$), Bilayer Height ($h(x_c)$), Average Tilt Angle of Cholesterol (θ_{avg}), and Area per DOPC (a_{DOPC}) and per Cholesterol (a_{CHOL}) Calculated Using Methods 1 and 2 and in the “Areal-Determining Plane” (ADP)

x_c (mol %)	$a_{\text{pl}}(x_c)$ (\AA^2)	$h(x_c)$ (\AA)	θ_{avg} (deg)	method 1		method 2		in ADP	
				a_{DOPC} (\AA^2)	a_{CHOL} (\AA^2)	a_{DOPC} (\AA^2)	a_{CHOL} (\AA^2)	a_{DOPC} (\AA^2)	a_{CHOL} (\AA^2)
0	67.1 \pm 0.5	39.4 \pm 0.3		67.1 \pm 0.6		67.1 \pm 0.6		67.1 \pm 0.5	
1.95	65.7 \pm 0.5	39.8 \pm 0.3	29.7 \pm 4.7	66.4 \pm 0.5	29.8 \pm 0.2	66.5 \pm 0.5	27.3 \pm 0.2	66.2 \pm 2.1	43.7 \pm 2.1
5.08	63.3 \pm 0.4	40.5 \pm 0.3	28.7 \pm 2.7	65.1 \pm 0.5	29.3 \pm 0.2	65.3 \pm 0.5	26.8 \pm 0.2	64.4 \pm 1.2	43.3 \pm 1.1
10.16	59.6 \pm 0.6	41.6 \pm 0.4	27.3 \pm 2.0	63.1 \pm 0.7	28.5 \pm 0.3	63.3 \pm 0.7	26.0 \pm 0.3	61.5 \pm 1.0	42.7 \pm 0.7
14.84	55.8 \pm 0.4	43.1 \pm 0.3	25.6 \pm 1.6	60.7 \pm 0.5	27.5 \pm 0.2	61.1 \pm 0.5	25.1 \pm 0.2	58.1 \pm 0.7	42.1 \pm 0.6
20.31	52.2 \pm 0.8	44.3 \pm 0.7	23.5 \pm 1.2	58.7 \pm 1.0	26.8 \pm 0.4	59.3 \pm 1.0	24.4 \pm 0.4	54.9 \pm 0.9	41.4 \pm 0.4
25	48.8 \pm 0.3	45.8 \pm 0.3	21.1 \pm 1.1	56.4 \pm 0.4	25.9 \pm 0.2	57.2 \pm 0.4	23.5 \pm 0.2	51.5 \pm 0.4	40.7 \pm 0.3
30.08	46.6 \pm 0.2	46.2 \pm 0.2	18.9 \pm 0.8	55.6 \pm 0.3	25.7 \pm 0.1	56.6 \pm 0.3	23.3 \pm 0.1	49.4 \pm 0.3	40.2 \pm 0.2
35.16	44.3 \pm 0.2	46.7 \pm 0.3	16.6 \pm 0.8	54.6 \pm 0.3	25.4 \pm 0.1	55.9 \pm 0.3	23.0 \pm 0.2	46.9 \pm 0.3	39.6 \pm 0.2
39.84	43.2 \pm 0.2	46.4 \pm 0.2	14.8 \pm 0.8	54.8 \pm 0.3	25.6 \pm 0.1	56.4 \pm 0.3	23.2 \pm 0.1	45.7 \pm 0.2	39.3 \pm 0.1
44.92	41.6 \pm 0.1	46.4 \pm 0.2	13.7 \pm 1.0	54.6 \pm 0.3	25.6 \pm 0.1	56.5 \pm 0.3	23.2 \pm 0.1	43.6 \pm 0.2	39.1 \pm 0.2
50	40.6 \pm 0.1	45.7 \pm 0.2	12.4 \pm 0.5	55.2 \pm 0.3	26.0 \pm 0.1	57.5 \pm 0.3	23.7 \pm 0.1	42.3 \pm 0.2	38.9 \pm 0.1
57	39.5 \pm 0.2	44.6 \pm 0.3	11.5 \pm 0.7	56.5 \pm 0.5	26.6 \pm 0.2	59.4 \pm 0.4	24.4 \pm 0.2	40.4 \pm 0.2	38.8 \pm 0.1
65.63	38.7 \pm 0.2	42.5 \pm 0.4	10.7 \pm 0.7	59.4 \pm 0.7	27.9 \pm 0.3	63.1 \pm 0.5	26.0 \pm 0.3	38.8 \pm 0.2	38.7 \pm 0.1

Acyl Chain Order Parameter. The order parameter of DOPC acyl chains was calculated using the last 50 ns of the simulation and was averaged from the two leaflets. Figure 5 shows the order parameter of the two chains at each carbon atom position. In general, as observed in some other MD simulation studies,^{5,14} the higher the cholesterol mole fraction, the higher the chain order parameter. Also, the *cis* double bond at C9 position produces a very sharp drop of chain order parameter. A new result in Figure 5 is that the chain order parameter at $x_c = 0.6563$ become slightly lower than that at $x_c = 0.50$ at some positions. To verify this result, we ran the simulation at $x_c = 0.6563$ for another 100 ns (i.e., 300 ns total). The result remained the same. x_c of 0.6563 is practically at the

maximum solubility of cholesterol in DOPC bilayer (0.66). Above this mole fraction, excess cholesterol precipitates from the bilayer to form cholesterol monohydrate crystals.²⁶ According to the umbrella model, the ability of DOPC to cover nonpolar bodies of cholesterol has been stretched to the limit at the solubility limit, and each DOPC needs to cover two cholesterol molecules on average.¹⁸ The bilayer is forced to adapt a few lateral distributions of lipids that can still satisfy the coverage requirement. Thus, the acyl chains of DOPC become slightly more disordered at the maximum solubility of cholesterol.

Slicing Bilayers. In order to understand the cholesterol condensing effect at atomic scale, we used the slicing method of Falck et al.¹⁴ with a modification. Instead of making a grid and counting the numbers of grids occupied by PC, cholesterol, water, and void, we counted the VDW areas that belong to these molecules. We first “sliced” the bilayer in z direction at 0.1 nm intervals. z is the distance from the bilayer center in the direction of bilayer normal. Within each slice, we counted the numbers

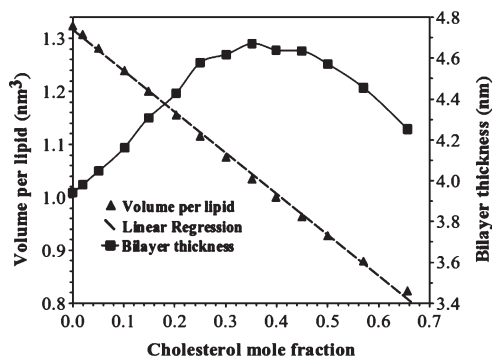


Figure 2. Volume per lipid $v_{\text{pl}}(x_c)$ and bilayer thickness $h(x_c)$ vs cholesterol mole fraction in DOPC bilayers. The linear fit to the volume per lipid data is $v_{\text{pl}}(x_c) = 1.316 - 0.775x_c$.

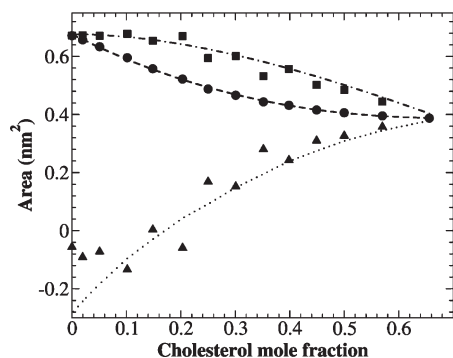


Figure 3. Area per lipid and partial-specific areas of DOPC and cholesterol vs cholesterol mole fraction. Circles: average area per lipid; square: DOPC partial-specific area; triangles: cholesterol partial-specific area.

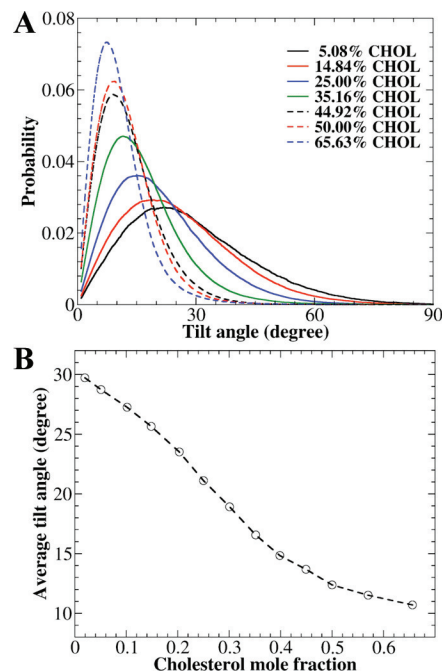


Figure 4. (A) Tilt angle distributions between the bilayer normal and the vector connecting C21 and C5 atoms of cholesterol in DOPC bilayers at some selected cholesterol concentrations. (B) Average cholesterol tilt angle versus cholesterol mole fraction.

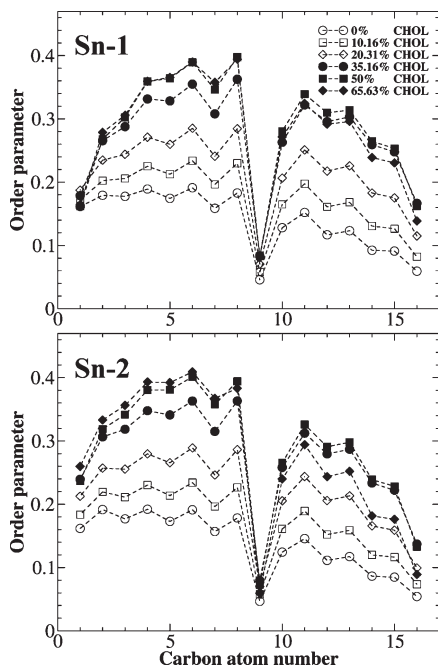


Figure 5. Acyl chain order parameter vs the carbon atom number in DOPC bilayers at some selected cholesterol concentrations.

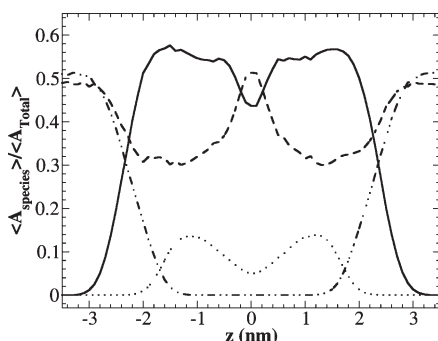


Figure 6. VDW areal fractions as functions of z for a DOPC bilayer containing 20.31 mol % of cholesterol. z is the distance from the bilayer center in the direction of bilayer normal. The areal fractions are normalized by total bilayer area. DOPC VDW areal fraction $\langle A_{\text{DOPC}}(z) \rangle / \langle A_{\text{total}} \rangle$ (solid line); cholesterol VDW areal fraction $\langle A_{\text{CHOL}}(z) \rangle / \langle A_{\text{total}} \rangle$ (dotted line); water areal fraction $\langle A_{\text{water}}(z) \rangle / \langle A_{\text{total}} \rangle$ (dot-dot-dash line); and free area fraction $\langle A_{\text{free}}(z) \rangle / \langle A_{\text{total}} \rangle$ (dashed line).

of atoms that belong to PC, cholesterol, or water. The area associated with each atom was calculated using the van der Waals radius.³⁰ The total VDW area occupied by DOPC, cholesterol, or water in each slice was obtained by summing together all corresponding VDW areas. The remaining areas that were not occupied by DOPC, cholesterol, or water were counted as free areas. Overall, our results were similar to those obtained by Falck et al. for DPPC/cholesterol bilayers using their grid counting method. Figure 6 shows the calculated area profiles as functions of z for a DOPC bilayer containing 20.31 mol % of cholesterol. Figure 7 shows a few selected areal slice snapshots for a DOPC bilayer containing 20.31 mol % of cholesterol at the end of a 200 ns simulation. At the bilayer center (Figure 7A), the free area is quite large and almost no water is present. Some cholesterol tails from one leaflet extend into the opposite leaflet. Figure 7B shows the region where DOPC acyl chains and cholesterol sterol rings are tightly packed. The amount of free area is lowest in this region. Figure 7C is a cross-sectional slice of the bilayer at a distance $z \sim 1.8$ nm from the bilayer center. Many cholesterol hydroxyl headgroups

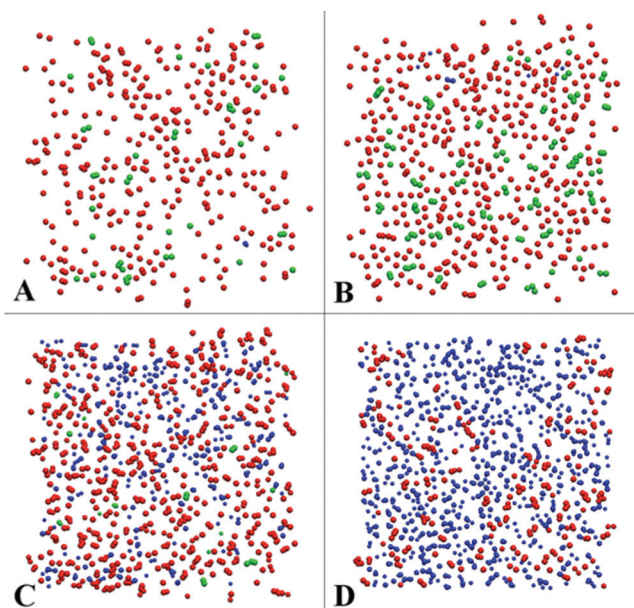


Figure 7. Snapshots of cross-sectional slices of a DOPC bilayer with 20.31 mol % of cholesterol at 200 ns. DOPC elements are colored in red, cholesterol in green, and water in blue. The remaining area is the free area. (A) $z \approx 0$, i.e., at bilayer center; (B) $z \approx 1.2$ nm, where many cholesterol sterol rings resided; (C) $z \approx 1.8$ nm, where many cholesterol headgroups resided; (D) $z \approx 2.3$ nm, where many DOPC headgroups resided.

are present in this slice, but they occupy a small fraction of total area. The amount of free area in this slice is smaller than the free area in the bilayer center. Figure 7D shows a cross-sectional slice at $z \sim 2.3$ nm. In this slice, there are substantial amounts of water and some DOPC headgroups, but no cholesterol.

Figure 8 shows the VDW area occupied by DOPC, VDW area of cholesterol, and free space as functions of location z at various cholesterol concentrations. Together, Figures 6–8 provide some important details about the packing of cholesterol in DOPC bilayers and the cholesterol condensing effect. The structure of a cholesterol-containing PC bilayer can be roughly divided into three regions: the bilayer center region, cholesterol sterol rings region, and PC headgroup region. (i) In the bilayer center region, as shown in Figure 8B, the density of cholesterol is low, since cholesterol has only a short acyl chain. The free space is largest at the bilayer center, as shown in Figure 8C. Thus, the molecular packing is quite loose at the bilayer center, and molecular interactions in this region *do not* determine the total area of the bilayer. (ii) In the cholesterol sterol ring region, the density of cholesterol is at its peak due to its dense sterol rings. As cholesterol mole fraction increases, the density of cholesterol (normalized to per cholesterol) shows little change (Figure 8B), indicating the sterol rings are rigid and not compressible. The largest change for cholesterol is its tilt angle with respect to the bilayer normal (Figure 4). As the cholesterol mole fraction increases, the average tilt angle decreases and the tilt angle distribution becomes narrower, which indicate that cholesterol is more ordered and becomes more parallel to the bilayer normal. In contrast, as shown in Figure 8A, the density of DOPC (normalized to per PC) shows a significant decrease as cholesterol mole fraction increases. The rigid sterol rings of cholesterol compress the acyl chains of DOPC, causing the tight packing and the increase of order parameter of acyl chains (Figure 5), which is a well-known effect of cholesterol.³¹ As expected, the free space is the lowest in this region (Figure 8C). The cholesterol condensing effect is most significant in this

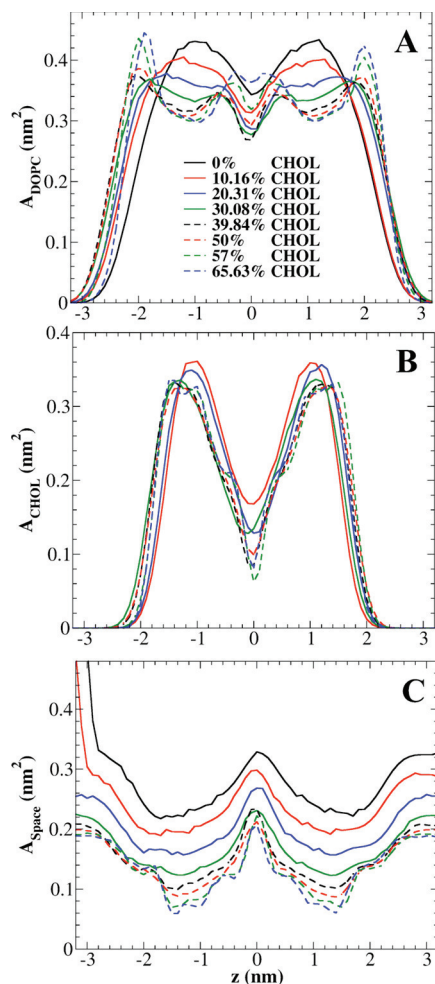


Figure 8. VDW area per DOPC (A), per cholesterol (B), and free space per lipid (C) versus location z at some selected cholesterol concentrations.

region, where cholesterol's sterol rings cut into the space originally occupied by the acyl chains of PC and reduce the overall bilayer area. (iii) In the PC headgroup region, the density of cholesterol is very low. However, the behavior of the PC density profile is more complex: As a result of increasing cholesterol mole fraction, the density of PC increases in this region as the acyl chains of DOPC become more ordered and the thickness of the bilayer increases. When x_c is above 0.2, the density profile of PC shows a small peak in this region. Furthermore, for $x_c > 0.39$, the peak becomes large and sharp and also shifts slightly toward the bilayer center. This shift is directly related to the decrease of bilayer height at high cholesterol shown in Figure 2. It is unknown whether such shift also exists in a DPPC/cholesterol system, since the highest x_c in the previous study was 0.50.¹⁴ Therefore, cholesterol does not produce a condensing effect in the PC headgroup region: In contrast, PC headgroups gradually expand to cover more lateral area along the bilayer–aqueous interface as x_c increases.

The condensing of PC acyl chains and the lateral expansion of PC headgroup can be elegantly explained by the umbrella model.^{18,32} The model suggests that the small headgroup/body ratio of cholesterol determines its mixing behavior in a lipid membrane. The cholesterol OH is a small polar head that cannot cover its larger nonpolar steroid ring body. In a bilayer, nonpolar cholesterol relies on polar phospholipid headgroup coverage to avoid the unfavorable free energy of cholesterol contact with water. Driven by the coverage requirement, cholesterol mol-

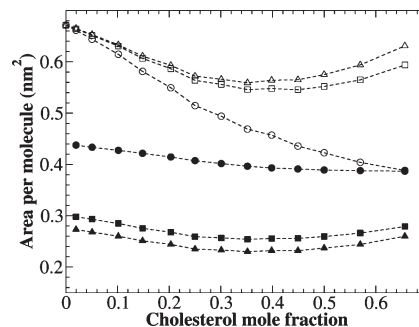


Figure 9. Area per DOPC and area per cholesterol calculated using different methods. Area per DOPC: by method 1 (open squares) and by method 2 (open triangles). Area per cholesterol: by method 1 (filled squares) and by method 2 (filled triangles). Area per cholesterol in the areal-determining planes (i.e., cholesterol sterol ring region): filled circles. Area per DOPC in the areal-determining planes: open circles.

ecules need to squeeze into the acyl chain region and partially hide under the headgroups of neighboring PCs, which will restrict the motions of the acyl chains and increase the chain order parameter. While this requirement causes the condensing of PC acyl chains, the headgroups of PC need to reorient and expand at bilayer–aqueous interface in order to cover more cholesterol. Curiously, the expansion of PC headgroups shown in Figure 8A was explicitly predicted in the umbrella model.¹⁸ Recently, MD simulation studies showed that the umbrella effect of cholesterol is present in both saturated and unsaturated PC bilayers.³³ Furthermore, a coarse grain simulation demonstrated that no cholesterol condensing effect could be observed if cholesterol structure is modified slightly by the addition of an extra hydrophilic headgroup or a reduction in size of the hydrophobic body.² The study confirmed that the cholesterol condensing effect originates directly from the mismatch between cholesterol's small polar headgroup and its large nonpolar body, not from the VDW attraction between cholesterol and phospholipids.

Area per PC and Area per Cholesterol. Figure 9 and Table 1 show the area per DOPC and area per cholesterol calculated using different methods of analysis. The value of $a_{\text{CHOL}}(x_c)$ calculated by method 1 varies between 0.298 and 0.254 nm², and $a_{\text{DOPC}}(x_c)$ varies between 0.671 and 0.546 nm². We noticed that volume per lipid vs x_c in our system is quite linear (Figure 2), similar to DPPC/cholesterol system. Therefore, following the procedure used by Edholm and Nagle,¹³ we first fit the volume per lipid using a linear function that assumes v_{DOPC} and v_{CHOL} are independent of x_c . The linear regression shown in Figure 2 gives $v_{\text{DOPC}} = 1.316$ nm³ and $v_{\text{CHOL}} = 0.541$ nm³. The area per DOPC and area per cholesterol calculated by method 2 were plotted in Figure 9 as well. The values of $a_{\text{CHOL}}(x_c)$ lie in the range 0.273–0.230 nm², and $a_{\text{DOPC}}(x_c)$ is within 0.671–0.559 nm², similar to the result obtained using method 1. The most important feature in Figure 9 is that both area per DOPC and area per cholesterol initially decrease with x_c but start to increase when x_c is greater than 0.40. The increase has not been observed in previous studies as the highest cholesterol mole fraction attempted was 0.50.⁵ The decreases of area per PC and area per cholesterol have been interpreted as the result of the cholesterol condensing effect. However, *this interpretation becomes problematic at high cholesterol mole fractions, since both calculated area per cholesterol and area per PC increase for $x_c > 0.40$, while the area per lipid continues to decrease (Figure 3).* The inconsistency was caused by the assumption built into methods 1 and 2: Both PC and cholesterol have cylindrical shapes and the same height. As shown in Figures

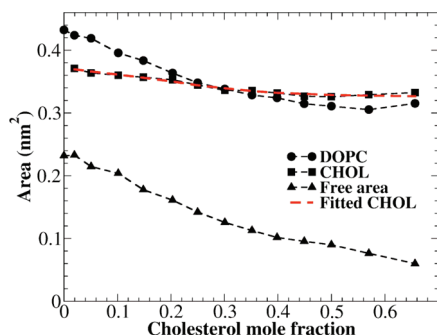


Figure 10. VDW areas of DOPC and cholesterol and free area per lipid as functions of cholesterol mole fraction in the areal-determining planes. The red dashed line is the fitting function $A_0/\cos(\theta(x_c))$, where $\theta(x_c)$ is the average cholesterol tilt angle plotted in Figure 4B, and A_0 is the fitting constant ($0.3179 \pm 0.0026 \text{ nm}^2$).

6–8, this is a problematic assumption: Cholesterol is shorter than DOPC, and the cross-sectional area of molecule varies widely as a function of z . Thus, both methods intrinsically overestimate area per DOPC and underestimate area per cholesterol. Fundamentally, the lipid bilayer is a three-dimensional structure: Treating it as a two-dimensional structure will yield unpredictable results.

Lipid Areas in the Areal-Determining Plane. Since the heights of the two molecules are different, Figures 6–8 show that there is a strong z -dependence between the cross-sectional areas of PC and cholesterol. Therefore, the average area per lipid and area per cholesterol are poorly defined parameters. Partial-specific areas of lipids correctly assess the change of total bilayer area when a lipid is added to or removed from the bilayer. On the basis of the discussion above, it can be concluded that the molecular packing in the cholesterol sterol ring region determines the total area of a cholesterol-containing lipid bilayer. Here, we define the horizontal plane, in which the VDW cross-sectional area of cholesterol reaches the maximum (Figure 8B), as the “areal-determining plane”. *The packing density of cholesterol and PC together is highest within this plane, and the PC–cholesterol interactions in this plane essentially determine the total area of the bilayer.* Figure 10 shows the VDW areas of DOPC and cholesterol as functions of cholesterol mole fraction within the areal-determining plane. The data were averaged over the last 50 ns of simulation. At each cholesterol mole fraction, the peak position of the VDW area of cholesterol (Figure 8B) was used to locate the areal-determining plane. The VDW area of DOPC in the areal-determining planes decreases from 0.431 to 0.308 nm^2 due to the compression by cholesterol sterol rings. For the same reason, the free area per lipid also decreases continuously. An important feature of Figure 10 is that the VDW area of cholesterol in this region decreases from 0.366 to 0.322 nm^2 . Since cholesterol sterol ring structure is rigid and does not change shape in a PC bilayer, this decrease must be caused by the decrease of cholesterol tilt angle with respect to the bilayer normal as cholesterol mole fraction increases. In order to visualize the effect of cholesterol tilt angle, the VDW area of cholesterol in the areal-determining plane was fitted by the function $A_0/\cos(\theta(x_c))$ in Figure 10 (the red dashed line). $\theta(x_c)$ is the average cholesterol tilt angle in Table 1 and Figure 4B, and A_0 is a fitting constant, which is the VDW area of cholesterol sterol rings at zero tile angle. The best fit was obtained with $A_0 = 0.3179 \pm 0.0026 \text{ nm}^2$. This good fit between cholesterol VDW area and $A_0/\cos(\theta(x_c))$ indicates that the decrease of VDW area of cholesterol with the increasing of x_c indeed results from the decreasing of cholesterol tilt angle.

On the basis of the above analysis, we reached the conclusion that the area lost due to the cholesterol condensing effect is primarily the VDW area and the free area originally belonged to PC. Therefore, we propose the following approach to quantitatively analyze the cholesterol condensing effect *within the areal-determining planes*: (i) The volume and shape of cholesterol fused rings are essentially independent of cholesterol mole fraction in a lipid bilayer. However, the average cholesterol tile angle has a large influence on the bilayer area. The area per cholesterol in the areal-determining plane is simply calculated using

$$a_{\text{CHOL}}(x_c) = \frac{a_0}{\cos(\theta(x_c))} \quad (8)$$

where a_0 is the cross-sectional area of cholesterol within the areal-determining plane with zero tilt angle plus some free area associated with the cholesterol, and $\theta(x_c)$ is the average tilt angle of the cholesterol as a function of x_c . (ii) The rest of the area is assigned to PC: The area per DOPC in the areal-determining plane is calculated by

$$a_{\text{DOPC}}(x_c) = \frac{A(x_c) - N_{\text{CHOL}}a_{\text{CHOL}}(x_c)}{N_{\text{DOPC}}} \quad (9)$$

where $A(x_c)$ is the total lateral area of the simulation box and N_{CHOL} and N_{DOPC} are the numbers of cholesterol and PC in each leaflet, respectively.

The VDW areas of PC and cholesterol shown in Figure 10 are bare areas. As pointed out by Edholm and Nagle,¹³ the free area or free volume is naturally a part of lipid bilayers. For pure PC lipid bilayers, all free area or free volume should be assigned to PC. The difficulty is how to divide the free area between PC and cholesterol when a bilayer contains cholesterol. The physical meaning of the parameter a_0 is the area occupied by the cholesterol sterol rings in a *hypothetical* pure cholesterol bilayer with zero tilt angle, which includes some free area associated with a cholesterol. Since the maximum solubility of cholesterol in many PC bilayers is 0.66,^{24,25} the value of a_0 cannot be obtained from either experiment or simulation. However, the possible range of a_0 can be narrowed down through following considerations: (i) The cross-sectional area of cholesterol in cholesterol monohydrate crystals was determined to be 0.36 nm^2 by Craven using X-ray diffraction.³⁴ (ii) At the air–water interface, in pure cholesterol monolayers at a lateral pressure similar to that of a lipid bilayer (30 mN/m),³⁵ the measured area per cholesterol is 0.37–0.39 nm^2 .^{36,37} This area includes the free area surrounding a cholesterol in the monolayers. (iii) The bare area of cholesterol sterol rings (without any free area) at zero tilt angle (i.e., A_0) was extrapolated to be $0.3179 \pm 0.0026 \text{ nm}^2$ in this study. This value should be considered as the lower limit of a_0 . Although the relevance of the molecular area of cholesterol in a crystal or a monolayer to that in a bilayer is a concern, we estimate that the reasonable value for a_0 should be between 0.35 and 0.40 nm^2 . In this study, we choose $a_0 = 0.38 \text{ nm}^2$ for the calculation.

The calculated area per DOPC and area per cholesterol in the areal-determining plane are also plotted in Figure 9. Because our definition is obviously different from that of methods 1 and 2, the obtained values are very different as well. The area per cholesterol in the areal-determining plane decreases from $\sim 0.438 \text{ nm}^2$ at low cholesterol to $\sim 0.387 \text{ nm}^2$ near its maximum

solubility limit ($x_c = 0.66$). The decrease is caused by the decrease of cholesterol tilt angle with respect to the bilayer normal. On the other hand, area per DOPC in the areal-determining plane decreases from the pure DOPC bilayer value of ~ 0.671 to ~ 0.388 nm² near $x_c = 0.66$. This large decrease of area per DOPC, about 40% of its original area, is caused by the compression of DOPC acyl chains by cholesterol's rigid sterol rings and reflects the true magnitude of the cholesterol condensing effect.

Conclusion

Based on the analysis of cross-sectional areas of molecules using the slicing method, the condensing effect of cholesterol is found to be limited to the cholesterol sterol ring region where the acyl chains of DOPC are severely compressed by cholesterol. As x_c increases, the headgroups of DOPC gradually expand along the bilayer–aqueous interface to cover more cholesterol. The results are in good agreement with the umbrella model. Our analysis showed that the total area of a PC/cholesterol bilayer is primarily determined by the molecular packing in the cholesterol sterol ring region, more precisely, in the areal-determining plane. The calculated partial-specific areas of lipids clearly show the condensing effect of cholesterol. A cholesterol-containing PC bilayer is indeed a 3-dimensional structure; therefore, average area per PC and area per cholesterol are not well-defined quantities. An alternative approach for calculating area per molecule within the areal-determining plane is proposed. In this region, the cholesterol tilt angle largely determines the area occupied by a cholesterol molecule, and the compression of PC acyl chains by cholesterol sterol rings is directly responsible for the well-known cholesterol condensing effect. Although the new method still has some arbitrary features, the calculated areas are more meaningful in describing the molecular packing and cholesterol condensing effect in a lipid bilayer: Our calculation showed that the majority of the area lost due to the cholesterol condensing effect is taken from PC molecules.

Acknowledgment. This work was supported by the National Institute of Health grant 1 R01 GM077198-01A1 subaward 49238-8402 and National Science Foundation grant MCB-0344463. The authors thank Mr. Charlie Huang for assistance in editing. The authors also acknowledge the High Performance Computing Center (HPCC) at Texas Tech University for providing HPC resources for this work.

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JP101415G