

Effect of Membrane Characteristics on Phase Separation and Domain Formation in Cholesterol-Lipid Mixtures

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ABSTRACT We examine, using an analytical mean-field model, the distribution of cholesterol in a lipid bilayer. The model accounts for the perturbation of lipid packing induced by the embedded cholesterol, in a manner similar to that of transmembrane proteins. We find that the membrane-induced interactions between embedded cholesterol molecules vary as a function of the cholesterol content. Thus, the effective lipid-cholesterol interaction is concentration-dependent. Moreover, it transitions from repulsive to attractive to repulsive as the cholesterol content increases. As the concentration of cholesterol in the bilayer exceeds a critical value, phase separation occurs. The coexistence between cholesterol-rich and cholesterol-poor domains is universal for any bilayer parameters, although the composition of the cholesterol-rich phase varies as a function of the lipid properties. Although we do not assume specific cholesterol-lipid interactions or the formation of a lipid-cholesterol cluster, we find that the composition of the cholesterol-rich domains is constant, independent of the cholesterol content in the bilayer.

INTRODUCTION

Cholesterol is the most abundant molecule in mammalian plasma membranes (Gennis, 1989). The incorporation of cholesterol has been shown to modulate the packing of the phospholipid molecules in the membrane, thereby increasing bilayer rigidity and mechanical durability, and reducing passive permeability (Simons and Ikonen, 2000). Moreover, cholesterol partitioning into cholesterol- and sphingolipids-rich domains has been found to trigger the formation of “rafts”, which are implicated in such diverse membrane processes as signal transduction, protein stabilization, protein and lipid sorting, and membrane fusion (Brown and London, 1998).

Although cholesterol is essential for the proper functioning of cell membranes, excess cholesterol levels could prove detrimental: Excess cholesterol may precipitate to form cholesterol monohydrate crystals, which play a significant role in the course of diseases such as gallstones or plaque deposition in atherosclerosis (Small, 1980, 1988).

Understanding the effect of cholesterol on membrane properties and functions—as well as the triggers for the nucleation of cholesterol crystals—requires understanding of the phase behavior of cholesterol in lipid bilayers. Various studies examined cholesterol-lipid mixing in both biological and synthetic membranes (Bach and Wachtel, 2003; Finegold, 1993; McConnell and Radhakrishnan, 2003; Ohvo-Rekila et al., 2002; Silviu, 2003). As summarized in Fig. 1, in the limit of low cholesterol concentrations, lipids and cholesterol are uniformly mixed. As the concentration of cholesterol increases, the system undergoes a process

reminiscent of phase separation to form cholesterol-rich domains coexisting with the dilute (gaslike) regions (Loura et al., 2001a,b; Veatch et al., 2004; Worthman et al., 1997). The composition of the cholesterol-rich domains, x_o , varies as a function of the lipid type (acyl chain length, headgroup charge, and the system parameters—temperature, composition, and pressure) (Crane and Tamm, 2004; Ohvo-Rekila et al., 2002; Veatch and Keller, 2002; Veatch et al., 2004). Radhakrishnan and McConnell (2003, 1999) find that the interactions between the lipids and the cholesterol in the cholesterol-rich domains are so specific as to suggest the formation of a “molecular complex”.

As the concentration of cholesterol in the bilayer increases, so does the fraction of the area occupied by the cholesterol domains (Crane and Tamm, 2004). Once the overall system composition reaches x_o , the domains occupy the entire membrane area. However, although the composition of the cholesterol-rich domains clearly corresponds to a preferred state, it does not define the maximal cholesterol solubility in the bilayer: The overall concentration of cholesterol may be increased up to a much higher value—defined as the solubility limit—at which cholesterol crystals appear (Huang et al., 1999). The main parameters controlling the solubility limit were found to be the sample preparation method, lipid type, length, and degree of unsaturation of acyl chains, presence of charge on lipid headgroup, and interheadgroup hydrogen bonds (Bach and Wachtel, 2003; Huang et al., 1999). For example, Huang et al. (1999) recently showed that at room temperature, the maximum solubility limit of cholesterol is 66 mol % in phosphatidylcholine bilayers and 51 mol % in phosphatidylethanolamine bilayers. However, the accuracy of these values may be questioned, since the solubility limit is commonly measured through the detection of cholesterol crystallites in solution (Bach and Wachtel, 2003), although recent studies suggest that crystallites initially nucleate and

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1993; Dan and Safran, 1998; Goulian, 1996; Goulian et al., 1993a).

One might expect that, since cholesterol perturbs the lipid packing, the membrane perturbation energy will be minimal when no cholesterol is present. However, we find that the free energy of the bilayer is minimal at a fixed cholesterol composition, x_o , which varies as a function of the lipid type. As a result, increasing the cholesterol molar concentration in the bilayer above a relatively low value leads to the formation of domains whose composition is set by x_o (see Fig. 1): Further addition of cholesterol simply increases the fraction of bilayer area occupied by the domains, but does not affect the domain composition. When the overall cholesterol concentration in the bilayer exceeds x_o (i.e., the entire membrane is taken up by the cholesterol domains), we find a transition into a one-phase cholesterol-condensed region where cholesterol is uniformly distributed in the bilayer.

MEMBRANE MODEL

Consider a membrane section of a cholesterol/phospholipid bilayer (Fig. 2). The membrane is taken to be composed of only one type of lipid, and the cholesterol to be distributed uniformly between the two monolayers. As a result, the system is symmetrical around the bilayer midplane.

The embedded cholesterol distorts the arrangement of the lipids in two ways: the first is due to a thickness mismatch, since the thickness of a cholesterol molecule (~ 1.7 nm) is smaller than that of a typical monolayer (~ 2 nm) (Yeagle, 1988). The second is due to the cholesterol structure: Molecular dynamics simulations show that the hydrophobic core of cholesterol is buried in the hydrocarbon region of the bilayer and that, on average, the molecule is tilted with respect to the bilayer normal, with a tilt angle of $\sim 14^\circ$ (Kessel et al., 2001). The rearrangement, or perturbation of membrane lipids incurred due to the thickness and angle mismatch with the cholesterol inclusion, increases the membrane energy when compared to a uniform (cholesterol-free) membrane, in the same way that protein inclusions do (Bezrukov, 2000; Dan et al., 1994, 1993; Dan and Safran, 1995, 1998; May 2000b, 2002; Nielsen et al., 1998).

For simplicity, we apply a one-dimensional model, namely, examining the perturbation as a function of distance in one dimension only (Fig. 2): Although obviously an oversimplification, previous analysis has shown that the one-dimensional model yields qualitatively and quantitatively similar results to that of the two-dimensional one in liposome-protein systems (Aranda-Espinoza et al., 1996).

The lipid bilayer is a self-assembled structure: properties such as the area per lipid Σ_0 and monolayer thickness L_m are determined by the lipid chemistry (Israelachvili, 1992), setting a free energy per lipid in the membrane of f_0 . The incorporation of an inclusion may lead to a perturbation in

the lipid packing, which incurs an energetic penalty. In the case of thickness mismatched inclusions, one may define the perturbation profile through a reduced thickness parameter: $\Delta(z) = (L(z) - L_m)/L_m$, where $L(z)$ is the thickness of the perturbed monolayer, L_m is the equilibrium monolayer thickness, and z is the distance from the inclusion boundary. At the inclusion boundary, Δ is defined by the difference between the inclusion thickness and the lipid bilayer thickness. In the case of a “packing” perturbation such as that incurred by cholesterol, the inclusion perturbs the packing of the neighboring lipids, namely, the area per molecule. However, the area per lipid and membrane thickness are coupled through an equation of state (Dan et al., 1994, 1993; Nielsen et al., 1998), so that $\Sigma = \Sigma(\Delta)$. Thus, any type of membrane perturbation may be described through the local thickness profile (see Fig. 2).

The membrane thickness profile varies as a function of the distance from the perturbation focal point: At the inclusion boundary, it is set by the inclusion properties; far away it decays to the unperturbed membrane value, i.e., $\Delta = 0$ or $\Sigma = \Sigma_0$. However, in membranes where the density of inclusions (e.g., proteins, cholesterol) is high and the average distance between inclusions small, the thickness perturbation may remain nonzero throughout the system. The change in monolayer energy (per inclusion) due to insertion of two inclusions, a distance $2d$ apart, is (Dan et al., 1993)

$$U_d = \int_0^d \frac{L_m}{\nu} \left(B\Delta^2 + (\kappa - \kappa'\Sigma_0)L_m \frac{d^2\Delta}{dz^2} + KL_m^2 \left[\frac{d^2\Delta}{dz^2} \right]^2 \right) dz. \quad (1)$$

The first contribution in Eq. 1 is due to packing constraints; B , the monolayer compressibility, describes the energy penalty for perturbation of the local density from equilibrium (namely, $\Delta = 0$ or $\Sigma = \Sigma_0$). B is defined as $\Sigma_0^2(\partial^2 f_0/\partial \Sigma^2)/2$ evaluated at Σ_0 (Dan et al., 1993). The second term accounts for the energy cost when the interface curvature does not match the preferred spontaneous curvature: The monolayer spontaneous curvature, κ , determines the sign and magnitude of the free interface curvature of the monolayer at an oil-water interface. The change in spontaneous curvature is given by $\kappa' = \partial\kappa/\partial\Sigma$, evaluated at the equilibrium bilayer surface density Σ_0 . In this article, we focus on bilayer-forming lipids, and in particular on molecules for which the spontaneous curvature, κ , and its derivative, κ' , are zero. The third contribution is due to the bending energy of the monolayer, where K is the bending modulus. $\nu \approx L_m\Sigma_0$ is the volume of a lipid molecule. All energies are given in units of kT , where k is the Boltzmann coefficient and T the temperature.

The separation, d , between the inclusions depends on their concentration in the membrane. For a single inclusion, the number of lipid molecules per inclusion is $\sim d/\sqrt{\Sigma_0}$. So, the

total number of molecules per inclusion is $1+d/\sqrt{\Sigma_0}$ and the inclusion mole fraction is

$$x = 1/(1 + d/\sqrt{\Sigma_0}) \quad (2)$$

To calculate the inclusion-induced bilayer perturbation profile and perturbation energy, the free energy (Eq. 1) must be minimized consistently with respect to the optimal perturbation profile. Boundary conditions for the system include a thickness mismatching condition at the inclusion/bilayer boundary (Δ_0 defines the reduced thickness difference) and a symmetry-enforcing condition, namely, $d\Delta/dz = 0$ at the midpoint between two inclusions ($z = d$). The third boundary condition is set by the slope of the inclusion at the inclusion-monolayer boundary, $d\Delta/dz = \tan\theta/a$ at $z = 0$, where a is a molecular length scale, and the fourth boundary condition is set by minimization requirements and reads

$$\left[\frac{d}{dz} \left(\frac{2L_m K}{v} \frac{d^2 \Delta}{dz^2} \right) \right]_{z=0} = 0 \quad (\text{Fox, 1950}).$$

Equation 1 may be minimized to yield the perturbation profile and energy in a general way for any value of K , B , and the inclusion mole fraction $x(d)$; however, although the values of the membrane thickness and surface density are well known for a variety of systems, measurements of K and B are not readily available. Thus, we use a molecular model to relate lipid properties to the bilayer parameters (Milner and Witten, 1988; Szleifer et al., 1988; Viovy et al., 1987). Although the model was developed for block copolymers (namely, both the head and tail are taken to be flexible chains), this approach has been shown (Szleifer et al., 1990, 1988) to yield qualitatively, and even reasonably quantitatively, correct results for several amphiphilic systems. For amphiphiles where the spontaneous curvature is zero (namely, $\kappa = 0$), the free energy coefficients are given by (Milner and Witten, 1988; Szleifer et al., 1988; Viovy et al., 1987)

$$f_0(\Sigma) = \gamma\Sigma + a_1 \frac{v}{\Sigma^2} \quad (3a)$$

$$K(\Sigma) = \frac{a_3 v^3}{4\Sigma^4}, \quad (3b)$$

where a_1 and a_3 have the dimensions of a length scale of molecular size (Milner and Witten, 1988). Using the above equations and the relationship $v = \Sigma_0 L_m$, we obtain

$$B \equiv \frac{\Sigma_0^2 f_0''}{2} = \frac{3a_1 L_m}{\Sigma_0} \quad (4a)$$

$$KL_m^2 = \frac{a_3 L_m^5}{4\Sigma_0}. \quad (4b)$$

RESULTS

Minimization of the bilayer free energy with respect to Δ yields the “optimal” perturbation profile—that is, the profile that minimizes the energetic cost associated with inclusion-induced membrane perturbation. For a single inclusion (namely, cholesterol at infinite dilution), we find that the energetic penalty associated with bilayer perturbation is given by

$$U_{\theta, \Delta_0} = \frac{2^{1/2} L_m B}{v A^{1/4}} \left[\Delta_0^2 + \frac{\tan^2 \theta}{a^2 A^{1/2}} + \frac{2^{1/2} \Delta_0 \tan \theta}{a A^{1/4}} \right], \quad (5)$$

where Δ_0 defines the reduced thickness mismatch at the cholesterol boundary and θ the contact angle between the membrane and the inclusion (see Fig. 1). A is the characteristic perturbation decay length—namely, the distance at which the membrane regains, more or less, its unperturbed characteristics. The perturbation length varies as a function of the membrane compressibility and bending modulus through the relationship $A = B/KL_m^2$. Thus, for similar bending moduli, the distance at which the bilayer regains its equilibrium thickness increases with increasing compression modulus, or resistance to area changes.

Equation 5 defines the energetic penalty for embedding a single cholesterol inclusion in a membrane. It constitutes three contributions that correspond to the three terms: The first is due to the thickness mismatch between the membrane and the inclusion. The second is due to the packing mismatch (i.e., “contact angle”) between the membrane and the inclusion, and the last is a cross term that accounts for the interrelationship between the two types of penalties. In the case of cholesterol inclusions, we may assign some values to the different terms in Eq. 5: Δ_0 is typically of order -0.15 (Yeagle, 1988) and θ is of order 14° (Tu et al., 1998). The membrane correlation length, A , depends on the lipid type, but is usually of order 1 nm^{-4} (Nagle and Tristram-Nagle, 2000). Thus, we find that, for cholesterol, the dominant term in Eq. 5 by far is the $\tan^2 \theta$ term. In further discussions, we will therefore neglect the contributions arising from the thickness mismatch, focusing instead on the contact angle.

The perturbation profile associated with inclusion of a cholesterol molecule into a membrane, neglecting the thickness mismatch, is then given by

$$\Delta(z) = \frac{2^{1/2} \tan \theta}{a A^{1/4}} \exp(-A^{1/4} z/2^{1/2}) \sin(A^{1/4} z/2^{1/2}). \quad (6)$$

In Fig. 3, we plot the thickness profile of a monolayer containing a single cholesterol inclusion. Despite the fact that we neglect the thickness mismatch between the cholesterol and the lipid monolayer, we see that the thickness of the monolayer is indeed perturbed by the cholesterol inclusion. This is due to the coupling between the membrane packing (surface area per molecule) and the thickness. Indeed, as in the case of thickness-mismatched inclusion/

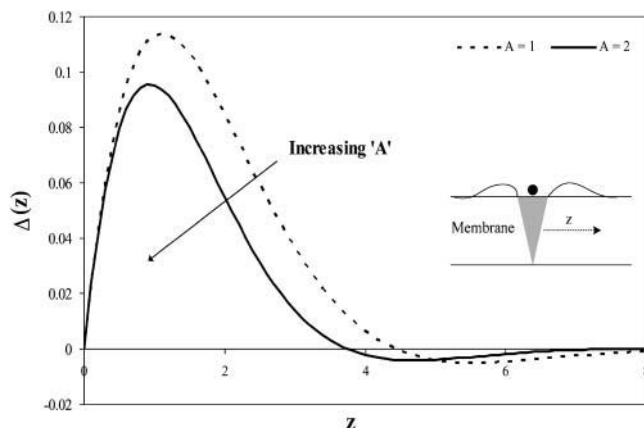


FIGURE 3 Perturbation profile of a monolayer containing a single cholesterol molecule (Eq. 6), as a function of distance from the inclusion boundary, z . The thickness profile of the monolayer is defined as $\Delta(z) = L(z) - L_m/L_m$. Despite the thickness matching, there is a monolayer thickness perturbation near the inclusion, the amplitude of which depends on A , the ratio of the monolayer compression modulus to the bending stiffness.

membrane systems (Aranda-Espinoza et al., 1996; Dan et al., 1993; Nielsen et al., 1998), the membrane thickness decays to its unperturbed value within a distance of order $(3-4 L_m)$ from the cholesterol boundary, which typically corresponds to 6–8 nm. Thus, in systems where the cholesterol spacing in the bilayer is $> \sim 8$ nm (which corresponds to ~ 0.08 mol fraction cholesterol), interactions between the cholesterol molecules may be neglected. However, above this relatively low cholesterol content, interactions between the cholesterol inclusions must be accounted for.

Calculating the membrane (optimal) perturbation profile for membranes containing cholesterol molecules that are separated by a distance $2d$ yields

$$\Delta(z) = \frac{2^{1/2} \tan \theta}{a A^{1/4}} \left[\frac{\sin\left(\frac{A^{1/4}(2d-z)}{2^{1/2}}\right) \sinh\left(\frac{A^{1/4}z}{2^{1/2}}\right) + \sin\left(\frac{A^{1/4}z}{2^{1/2}}\right) \sinh\left(\frac{A^{1/4}(2d-z)}{2^{1/2}}\right)}{\sin(2^{1/2} A^{1/4} d) + \sinh(2^{1/2} A^{1/4} d)} \right], \quad (7)$$

where $x = 1/(1+d/\sqrt{\Sigma_0})$ is the cholesterol mole fraction. Substituting this profile into the expression for the energy (relative to the dilute limit) yields

$$U_{\text{eff}} = \frac{2^{1/2} \pi^2 \tan^2 \theta L_m^4}{64 a \Sigma_0^2} \left[\frac{\cos(2\sqrt{2\Sigma_0}(1-x)/L_m x) + \cosh(2\sqrt{2\Sigma_0}(1-x)/L_m x)}{\sin(2\sqrt{2\Sigma_0}(1-x)/L_m x) + \sinh(2\sqrt{2\Sigma_0}(1-x)/L_m x)} - 1 \right]. \quad (8)$$

U_{eff} defines the membrane energetic penalty, per inclusion, due to the incorporation of x mole fraction cholesterol. It is composed of two contributions: one is the energetic penalty

associated with the insertion of the individual cholesterol into the bilayer (i.e., Eq. 5); the other component is due to the membrane-induced interactions between cholesterol molecules.

In Fig. 4, we plot the membrane perturbation energy as a function of the cholesterol mole fraction (which is inversely proportional to the average distance between cholesterol molecules). The value of $U_{\text{eff}} = 0.708$ kT/nm defines the energy required to insert a single cholesterol molecule (Eq. 5). We see that for low cholesterol mole fractions, U_{eff} remains constant, thereby indicating that the energetic penalty associated with cholesterol incorporation remains constant, or that the cholesterol perturbation domains do not overlap (i.e., no membrane-induced interactions between cholesterol molecules). As x increases, we see a very slight increase in the energy corresponding to a region where the perturbed areas of the membrane start to overlap. Quite surprisingly, however, we see a wide region (in the case shown in Fig. 4, it corresponds to $0.22 \leq x \leq 0.5$) where U_{eff} decreases below the value of the dilute, single inclusion, reaching a minimum at $x = x_0$ (≈ 0.36 in this case). The location of this minimum depends on the bilayer characteristics, and is given approximately by $x_0 \approx (1 + 1/\sqrt{2\Sigma_0}/L_m)^{-1}$.

The minimum in U_{eff} as a function of x indicates an effectively attractive interaction between the embedded cholesterol molecules—namely, that the penalty associated with membrane perturbation is reduced due to interactions between the perturbation profiles of neighboring cholesterol inclusions.

Why would the bilayer, which is obviously perturbed by the cholesterol inclusions, favor a moderately close packing rather than complete phase separation (between pure cholesterol and pure lipid domains)? To understand this,

we must reexamine the membrane thickness profile. As shown in Fig. 3, in the dilute cholesterol case, the membrane thickness increases and then decreases to the unperturbed

value. However, if the cholesterol density is higher (see inset in Fig. 4), the membrane thickness does not decay back to the unperturbed value, thereby reducing the curvature penalty.

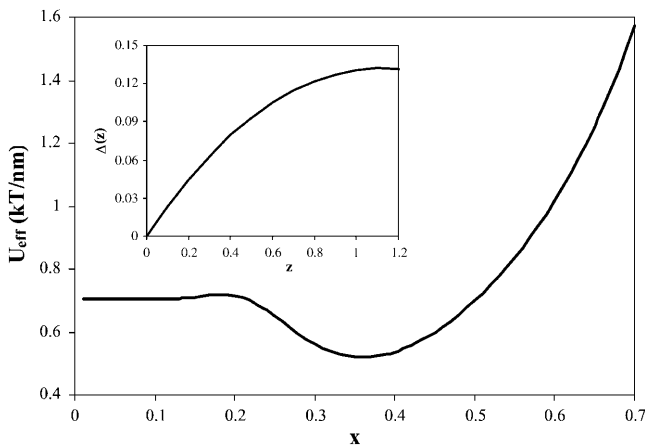


FIGURE 4 Membrane perturbation penalty per inclusion, Eq. 8, as a function of the cholesterol mole fraction, x . For a given membrane, the free energy of the system is at a global minimum at a finite composition, defined by x_o , which depends on the type of lipid. Here $L_m = 1.7$ nm and $\Sigma_o = 0.4$ nm². The inset shows the membrane perturbation profile at $x = 0.36$, namely, where U_{eff} is minimal. The perturbation decay length is taken to be $A = 1$ nm⁻⁴.

Once the concentration of cholesterol increases above a critical value (in this case, when $x \approx 0.5$), the perturbation energy increases sharply, indicating a region where increasing the cholesterol concentration is highly unfavorable, or, alternately, where the membrane-induced cholesterol-cholesterol interactions become strongly repulsive.

The membrane perturbation energy, U_{eff} , indicates that there is an effectively attractive interaction between cholesterol molecules embedded in a lipid bilayer, which favors regions with a specific cholesterol mole fraction: The energy associated with the membrane perturbation may be reduced by forming cholesterol domains whose composition is given by x_o . This suggests that, if the overall cholesterol content in the bilayer is lower than x_o , the membrane will phase-separate into domains whose composition corresponds to x_o and regions where the cholesterol content is low.

To correctly evaluate the mixing phase diagram of cholesterol in the bilayer, we cannot focus only on the membrane-induced interactions. We need to calculate the overall system free energy, which includes the interaction energy, U_{eff} , the two-dimensional mixing entropy, and the direct interactions between cholesterol molecules in the bilayer.

The direct interactions between membrane-embedded cholesterol molecules are composed of two contributions: The first, which is attractive, is due to van der Waals forces (Yeagle, 1988). The second, which is effectively repulsive, is due to an interfacial penalty associated with the increased cholesterol exposure to water in cholesterol-only domains when compared to the lipid “shielding” effect (Huang and Feigenson, 1999). Both of these are negligible when

compared to the membrane perturbation penalty, except in extremely high concentrations when $x \approx 1$ or $d \approx 0$. Since in this analysis we focus on moderate cholesterol concentrations, we neglect these contributions. Thus, the membrane mixing free energy per unit width is given by

$$\Delta F_{\text{mix}} = x(U_{\text{eff}} - T\Delta S_{\text{mix}}), \quad (9)$$

where $-T\Delta S_{\text{mix}} = \ln(x) + ((1-x)/x)\ln(1-x)$ is the dimensionless entropy of mixing in a two-component system, per inclusion (see, for example, Dill and Bromberg, 2002).

The mixing behavior, or phase diagram at a fixed temperature T of the cholesterol-lipid systems, is set by the value of $\partial^2(\Delta F_{\text{mix}})/\partial x^2$ at the given mixture composition, x : If the second derivative is positive, the mixed system is stable. If it is negative, the system separates into coexisting domains. As shown in Fig. 5, we find that (as expected from U_{eff}) in the limit of either dilute cholesterol limit or high concentrations, the system favors mixing (regions α and δ). However, in an intermediate range, we find a region, β , where the system undergoes phase separation into cholesterol-rich and cholesterol-poor domains. The composition of the cholesterol-rich domains is set by x_o , the composition at which the bilayer energy is minimal. (The composition of the coexisting phases may be calculated using the Maxwell construction. Due to the distinct minimum in U_{eff} at x_o , which dominates ΔF_{mix} , we find that the composition of the cholesterol-rich phase is closely associated with x_o .) The range of compositions at which domain coexistence appears varies as a function of the lipid characteristics, namely, area per molecule and bilayer thickness.

As the average mole fraction of cholesterol in the phase-separated region increases, so does the fraction of bilayer area occupied by the cholesterol-rich domains. Once x exceeds that of the cholesterol-rich domains ($\sim x_o$), the membrane enters another uniformly mixed phase—the cholesterol-dense phase. As shown by Fig. 4, in this region

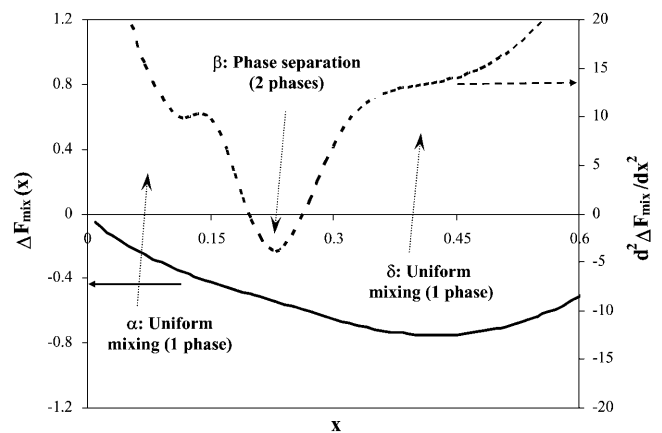


FIGURE 5 Second derivative of the system free energy (Eq. 9) as a function of cholesterol mole fraction x for $\sqrt{\Sigma_o}/L_m = 0.354$.

the membrane perturbation energy per cholesterol increases almost linearly with the cholesterol mole fraction.

One issue of interest is the nucleation of cholesterol crystals. Crystallite nucleation can take place via two routes. In the first one, excess cholesterol remains in solution, thereby nucleating crystallites in the bulk suspension (Huang et al., 1999; Huang and Feigenson, 1999). Alternately, the excess cholesterol may segregate in the bilayer into condensed cholesterol domains, which coalesce with time and precipitate into solution over time (Troup et al., 2003). In either case, cholesterol will accumulate in the bilayer until the chemical potential of the cholesterol in the membrane becomes equal to that of cholesterol in the monohydrate crystals, or (neglecting the mixing entropy in all phases) when $U_{\text{eff}} = \mu_{\text{crystal}}$. Unfortunately, estimating the chemical potential of cholesterol in monohydrate crystals (whether within the bilayer or in bulk suspension) is outside the domain of this study: It accounts for a combination of the molecular, short-range cholesterol-cholesterol interactions as a function of the cholesterol organization in the crystal, as well as the interfacial tension between the crystal and the surrounding solution. However, we may qualitatively estimate the effect of bilayer characteristics on the maximum solubility limit through evaluation of the effect of these parameters on U_{eff} in the limit of high cholesterol content. As shown in Fig. 6, we find that (for a given bilayer thickness) U_{eff} decreases with increasing lipid surface density Σ_0 . Thus, we conclude that the critical solubility limit shifts toward higher values for lipids with a larger headgroup area (Σ_0). This result is in qualitative agreement with the experimental results of Huang et al. (1999), where it was shown that, comparing phosphatidylcholines versus phosphatidylethanolamine bilayers, the maximum solubility limit increases with the headgroup area.

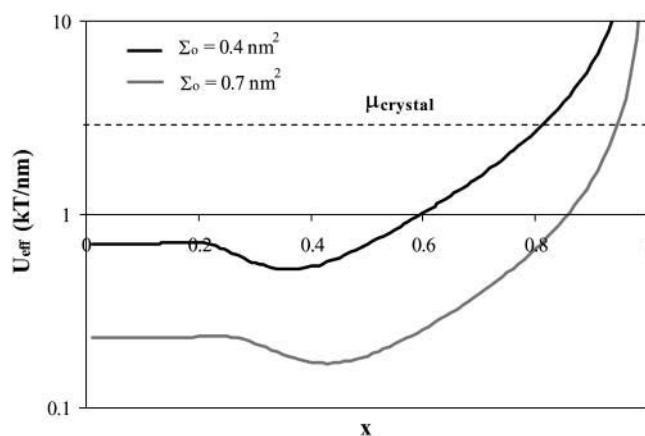


FIGURE 6 U_{eff} as a function of x for two different lipid surface densities (but L_m is the same). The maximal solubility limit is defined as the x value at which U_{eff} exceeds the chemical potential of the cholesterol in the crystal. We see that (regardless of the value of μ_{crystal}) the solubility limit will be higher for the higher density bilayer. Note that the y axis is on a log scale.

DISCUSSION AND CONCLUSIONS

In this article, we develop a mean-field model for cholesterol-lipid bilayers. The model accounts for the cholesterol-induced perturbation of the bilayer, which gives rise to an energetic penalty whose magnitude is sensitive to the cholesterol content in the bilayer. One of the main conclusions from our analysis is that the enthalpy of mixing between cholesterol and lipids in the bilayer cannot be accurately described using a constant interaction potential that is concentration-independent, as done in standard mean-field lattice models (i.e., writing the energy as $Ux(1-x)$, where U is independent of x) (Chiu et al., 2002; Huang, 2002; Huang and Feigenson, 1999; Pasenkiewicz-Gierula et al., 2000). In fact, we find that the nature of the cholesterol interactions, let alone their numerical value, changes as a function of the cholesterol content: At low cholesterol concentrations, the membrane-induced interactions between cholesterol molecules are repulsive, favoring uniform distribution. As the concentration exceeds above a certain value, the interactions turn positive, favoring a specific distance between cholesterol molecules embedded in the bilayer, thereby leading to phase separation between cholesterol-rich and cholesterol-poor domains. As the concentration increases above a second critical value, the interactions become repulsive again, favoring uniform mixing.

The composition of the cholesterol-rich domains, x_0 , is found to vary as a function of the lipid type (Fig. 4), but to be largely independent of the overall cholesterol composition in the system. This is in agreement with the experiments of Radhakrishnan and McConnell (McConnell and Radhakrishnan, 2003; Radhakrishnan et al., 2000; Radhakrishnan and McConnell, 1999), who find that the interactions between the lipids and the cholesterol in the domains formed in monolayers at the air-liquid interface are so specific as to suggest the formation of a “molecular complex”. This is in qualitative agreement with our analysis, which demonstrates that the system energy is greatly minimized upon the formation of regions characterized by a specific cholesterol/lipid ratio. In summary, we developed a model to understand the influence of membrane characteristics on domain formation and phase separation in binary mixtures of lipids and cholesterol. We find that the nature and magnitude of the membrane-mediated interactions between cholesterol molecules change as a function of the cholesterol content. As a result (Fig. 7), we find that the cholesterol-lipid bilayer transitions from a homogeneous, dilute (gaslike) at low cholesterol content to a phase-separated coexistence between cholesterol-poor and cholesterol-rich domains, followed at higher cholesterol concentrations by another transition to a uniform phase. The composition of the cholesterol-rich domains in the phase-separated regime is largely set by the lipid characteristics and is independent of the cholesterol concentration.

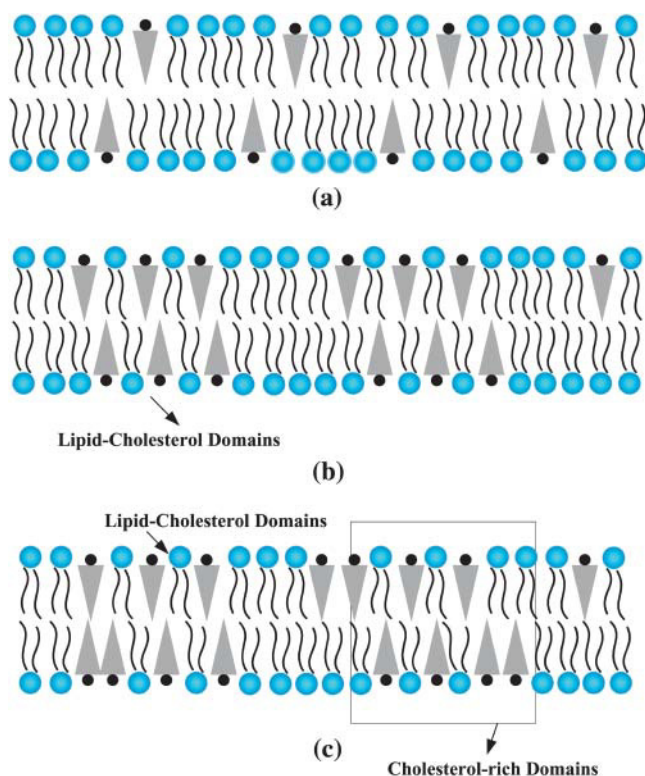


FIGURE 7 Schematic of the coexisting phases in a lipid-cholesterol bilayer: (a) at low concentrations, cholesterol is uniformly distributed in the bilayer; (b) as the concentration is increased, lipid-cholesterol domains start to form; and (c) above a certain cholesterol concentration, cholesterol-rich domains coexist with the lipid-cholesterol domains.

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