

Cyclic Activity of an Osmotically Stressed Liposome in a Finite Hypotonic Environment

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Cite This: *Langmuir* 2020, 36, 3659–3666

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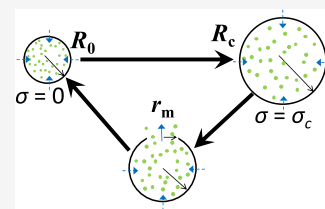


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ABSTRACT: A lipid vesicle, or simply called a liposome, represents a synthetic compartment for the examination of transmembrane transport and signaling phenomena. Yet, a liposome is always subjected to size and shape fluctuations due to local and global imbalance of internal and external osmotic pressures. Here, we show that an osmotically stressed liposome placed within a hypotonic spherical bath undergoes cyclic dynamics described by a periodic sequence of swelling and relaxation phases. These two phases are interfaced by the appearance of a transient transmembrane pore through which chemical delivery occurs. An analytical model was formulated for the recurrent differential equations that convey the time-dependent swelling phase of a pulsatory liposome during individual cycles. We demonstrate that the time-dependent swelling phases of the last several cycles of a pulsatory liposome are strongly dependent on the size of the external bath. Furthermore, decreasing the size of the hypotonic medium reduces the number of cycles of a pulsatory liposome. Comparisons and contrasts of an infinite hypotonic bath with finite external baths of varying radii are discussed.



INTRODUCTION

Cellular response to local and global fluctuations in the osmotic pressure of the external environment is a fundamental property of living systems. There are numerous mechanisms by which cells operate for regulating the strain of the cellular membrane, including mechanosensitive and water channels. In a much more simplified context, a unilamellar liposome undergoes shape and size fluctuations upon rapid changes in the environmental osmotic pressure. It has already been documented that a liposome undergoes a cyclic activity when it is filled with an aqueous solution of an increased osmotic pressure and it is placed within a finite hypotonic environment.^{1–3} In each cycle, the osmotic influx of water swells the liposome up to a critical point at which the membrane attains the maximum strain that is characterized by a critical (or lytic) tension, σ_c (Figure 1A). In this way, the radius of a spherical liposome expands from an initial value, R_0 , to a critical value, R_c . At this point, the nucleation of a transient transmembrane pore occurs,^{4–8} sometimes leading to an irreversible rupture of the liposomal membrane.^{9–12} This phase, also called the swelling phase,¹³ is followed by solute extrusion through the transient pore and a fast relaxation (or compression) phase until the liposome reaches its initial size. Then, a new cycle begins and the liposome activity becomes cyclic. This is the reason why we call this liposome a pulsatory liposome. Therefore, a pulsatory liposome operates like a two-stroke engine. The transmembrane gradient of osmotic solute assures the potential energy that generates the functional force of the engine.

The succession of swelling and relaxation phases is mediated by the appearance of a transmembrane pore formed in a stretched liposomal membrane (Figure 1B). A significant

number of theoretical,^{4,7,10,11,14–24} computational,^{25–27} and experimental^{5,11,28–32} studies have reported the presence of transient pores in liposomes and planar lipid membranes. Moreover, there is an increasing interest in the exploration of these metastable transmembrane pores because of their pivotal role in the cyclic activity of an osmotically stressed liposome.^{1–3,7,33–37}

In this paper, we put an emphasis on the swelling phase of the cyclic activity of a liposome when it is inserted into a finite hypotonic environment. If the external bath is filled with water, then the external concentration of the solute increases in a cyclic fashion.^{1,2,37–39} The cyclic activity of the liposome depends on the transmembrane gradient of the solute. The relaxation phase is short-lived, and it does not affect the swelling phase.² Our expectation is that the size of the hypotonic environment has a significant impact on the number of cycles, the swelling time of a pulsatory liposome, as well as the total amount of solute released into the external environment. On the other hand, we expect that other biophysical parameters of a pulsatory liposome, such as the relaxation time, are not significantly affected. Therefore, we wanted to explore the swelling phase of a pulsatory liposome of varying cycle order.

Received: December 21, 2019

Revised: March 17, 2020

Published: March 18, 2020



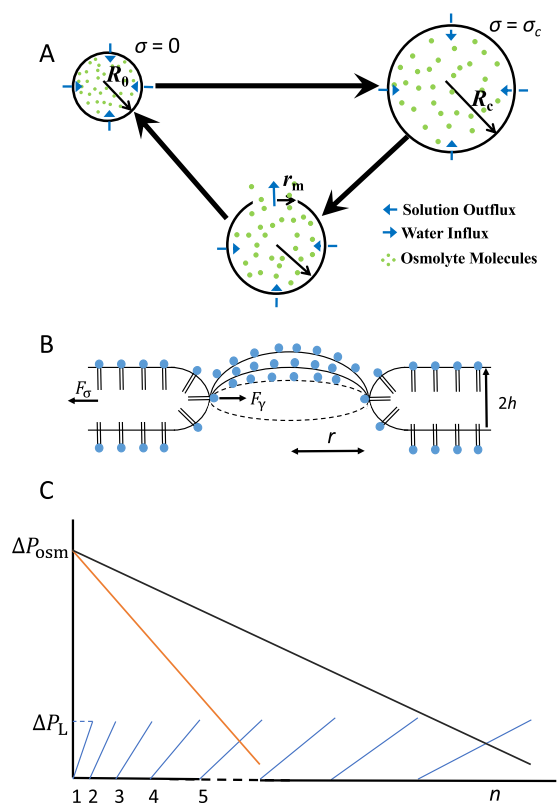


Figure 1. Diagrams that show details of the dynamics of a pulsatory liposome during a single cycle. (A) Each cycle encompasses two phases: swelling phase (top) and relaxation phase (bottom). The initial radius of the liposome, R_0 , is reached when the surface tension is zero. In contrast, the critical radius of the liposome, R_c , is attained at a critical surface tension, σ_c . At this point, the liposomal membrane undergoes a lytic tension, leading to a transient transmembrane pore. The maximum radius of the transient pore is r_m . (B) This model assumes the osmotic stress-induced formation of a single transient pore across the liposomal membrane. The dynamics of the transient pore is affected by the balance between the driving force for the pore opening, F_σ , which corresponds to the mechanical surface tension of the membrane, σ , and the driving force for pore closure, F_γ , which corresponds to edge tension (or line tension), γ . The former force is created by the Laplace pressure, whereas the latter force is created by the strong hydrophobic interactions among the alkyl tails of the liposomal phospholipids. r is the pore radius. h is the thickness of a monolayer of the liposomal bilayer membrane. (C) Schematic that shows the balance of the osmotic and Laplace pressures during the succession of various cycles. ΔP_{osm} and ΔP_L are the osmotic and Laplace pressures, respectively. n is the cycle order. A pulsatory liposome becomes active when $\Delta P_L < \Delta P_{\text{osm}}$. The tilted line in black corresponds to a pulsatory liposome located within an infinite hypotonic bath (e.g., a larger number of cycles). The tilted line in orange corresponds to a pulsatory liposome located within a finite hypotonic bath (e.g., a smaller number of cycles).

Here, we report that the size of the finite hypotonic bath has a substantial impact on the total number of cycles and the amount of solute released during the active life of the liposome. A pulsatory liposome becomes inactive at a point in which the internal osmotic pressure is equal to the Laplace pressure (Figure 1C). The outcomes of this study represent a theoretical platform for further developments with a practical impact. Specifically, the solute might be substituted by a chemical agent of pharmacological significance for a healthcare application. In this case, a pulsatory liposome can release

drugs in a programmable fashion at a strategic location of a targeted tissue.^{40,41}

METHODS

Background. Let us consider a large liposome filled with a solute-containing aqueous solution and assume that it is located within a sufficiently large, water-filled spherical bath. In the first phase, the liposome expands from the initial state up to a critical state when a single transient transmembrane pore appears. Let us set the initial liposomal radius equal to R_0 . In this case, the liposomal membrane is unstretched so that the mechanical surface tension, σ , is zero. In the critical state, the liposomal radius and surface tension are R_c and σ_c , respectively. In the second phase, the cycle is determined by the appearance of the transmembrane pore. The time trajectory of the transient pore has two phases: in the first part, the pore radius increases up to a maximum value, r_m , and in the second part, the pore radius declines up to the pore closure, leading to the recovery of the liposomal membrane (Figure 1A,B). During this process, some solute amount is extruded through the transient pore.

As soon as the membrane is recovered, the liposome returns to its initial state and a new cycle begins. In our model, we assumed that the critical membrane tension, σ_c , at which the relaxation phase starts depends on the intrinsic properties of the liposomal membrane so that this is assumed unaltered by increasing the cycle order.^{1,10,11} Because of the solute release through the transient transmembrane pore, the internal solute concentration is changed at the end of each cycle. The liposomal expansion is mediated by the balance between the osmotic and Laplace pressures. This is the conceptual mechanism by which an osmotically stressed liposome placed within a finite, hypotonic environment can undergo a cyclic activity as a result of the balance between the osmotic and Laplace pressures (Figure 1C). Overall, each cycle of a pulsatory liposome is characterized by a differential equation for the liposomal radius during the swelling (expansion) phase and a system of three differential equations for the liposomal radius, transient pore radius, and solute internal concentration during the relaxation (compression) phase.

Analytical Model for the Swelling Phase. Let us consider that the initial state of the liposome is defined by the surface area, A_0 , and the volume, V_0 . At the same time, the initial state of the liposome is characterized by the initial internal concentration of the solute, C_{01} . In a hypotonic environment, the liposomal volume increases because of the osmotic pressure difference between the interior and exterior of the lipid vesicle. The direct outcome of the volume growth is the stretching of the bilayer membrane and the appearance of the Laplace pressure. During liposomal swelling, two pressures occur in directions against each other: osmotic pressure, ΔP_{osm} , and Laplace pressure, ΔP_L , corresponding to the transmembrane gradient of the solute concentration and surface tension, respectively (Figure 1B).

The rate of change in the liposomal volume, V , is described by the following equation

$$\frac{dV}{dt} = P_w V_{\mu w} \beta A (\Delta P_{\text{osm}} - \Delta P_L) \quad (1)$$

Here, P_w and $V_{\mu w}$ denote the water permeability across the liposomal membrane and the water molar volume, respectively. A is the total area of the liposome.

Here,

$$\beta = \frac{1}{N_A k_B T} \quad (2)$$

where N_A , k_B , and T are the Avogadro number, the Boltzmann constant, and the absolute temperature, respectively. The osmotic pressure, ΔP_{osm} ,^{42,43} and Laplace pressure, ΔP_L ,^{29,30} are given by the following equations

$$\Delta P_{\text{osm}} = N_A k_B T \Delta C_m \quad (3)$$

$$\Delta P_L = \frac{2\sigma}{R} \quad (4)$$

where

$$\Delta C_m = C_{in} - C_{out} \quad (5)$$

is the transmembrane gradient of the solute concentration. R denotes the liposomal radius. C_{in} and C_{out} are the solute concentrations inside and outside the liposome, respectively. According to Hooke's law of elasticity in a two-dimensional formulation, if a closed spherical membrane is stretched by surface tension, σ , the radius of the swelled state, $R(\sigma)$, is

$$R(\sigma) = R_0 \sqrt{1 + \frac{\sigma}{E}} \quad (6)$$

where R_0 and R are the liposomal radii in the initial (untensed) state and expanded (tensed) state, respectively. In eq 6, E is the elastic modulus for the stretching and compression of the liposomal membrane in two dimensions. Therefore, if a liposome is stretched, then the mechanical tension is given by the following formula

$$\sigma = E \frac{R^2 - R_0^2}{R_0^2} \quad (7)$$

Let us assume a spherical geometry of the liposome. Using eqs 2–7, we rewrite eq 1 in the following form

$$\frac{dR}{dt} = P_w V_{\mu w} \left[\Delta C_m - \frac{2\beta E}{R} \left(\frac{R^2}{R_0^2} - 1 \right) \right] \quad (8)$$

Let us consider two cases. In the first case, the liposome is placed within an infinite water bath with a time-independent solute concentration ($C_{out} = 0$). In the second case, the liposome is placed within a finite (closed) hypotonic environment with a time-dependent solute concentration. In the latter case, C_{out} increases during the pulsatory solute exchange with the hypotonic environment, whereas C_{in} decreases. Using the law of mass conservation during the swelling process of the first cycle (Supporting Information, eqs S1–S7), eq 8 can be analytically solved for the first cycle, as follows

$$\frac{8\alpha\beta EP_w V_{\mu w}}{R_0^2} t = (\alpha + 1) \ln \left| \frac{\alpha - 1}{2x^2 - \alpha - 1} \right| + (\alpha - 1) \ln \left| \frac{\alpha + 1}{2x^2 + \alpha - 1} \right| \quad (9)$$

where

$$\alpha = \sqrt{1 + \frac{2C_{01}R_0}{\beta E}} \quad (10)$$

and

$$x(t) = \frac{R(t)}{R_0} \quad (11)$$

Here, C_{01} is the initial internal solute concentration.

Differential eq 9 has a solution that is a bijective function

$$t = t(x), \text{ with } x \in \left[1, \frac{R_c}{R_0} \right] \text{ and } t \geq 0 \quad (12)$$

where R_c is the critical radius of the liposome. Here, we define the critical swelling ratio, x_c , by the following formula

$$x_c = \frac{R_c}{R_0} \quad (13)$$

One might calculate the inverse function $R = R(t)$. Yet, the differential equation of the swelling process of a pulsatory liposome in the n -th cycle is given by the following expression (Supporting Information, eqs S8–S16)

$$\frac{dx}{dt} = \frac{P_w V_{\mu w} C_{01}}{R_0} \left[\frac{f^{n-1}}{x^3} - \frac{2\beta E}{R_0 C_{01}} \left(x - \frac{1}{x} \right) - \frac{1 - f^{n-1}}{F - x^3} \right] \quad (14)$$

where

$$f = \frac{R_0^3}{R_c^3} \quad (15)$$

The differential equations that describe the swelling phases of a pulsatory liposome, except the equation for the first cycle, can be solved using numerical methods.

Comparisons between the First Cycle and Subsequent Cycles. The major question in this study is how the “finite” nature of the spherical environment with a radius, R_b , impacts the parameters of the liposome. Specifically, we solve the differential equation for the swelling phase of each cycle. For the first cycle, $C_{out} = 0$, and the differential equation is provided by eq 9. The final form of eq 8 for the expansion phase of a pulsatory liposome during the n -th cycle ($n \geq 2$, $C_{out} > 0$) is eq 14. The functional operation of a pulsatory liposome is contingent upon a specific imbalance between competing pressures: the osmotic pressure is greater than the Laplace pressure

$$RT\Delta C > \frac{2\sigma}{R} \quad (16)$$

This is the running condition of a pulsatory liposome. During the cyclic activity of the liposome, its osmotic pressure declines continuously until this becomes equal to the Laplace pressure. At this point, the cyclic activity of a pulsatory liposome stops. The number of cycles achieved by a pulsatory liposome is calculated by the following formula

$$N = \text{INT} \left[\frac{\ln F - \ln Z}{3 \ln x_c} \right] \quad (17)$$

where

$$F = \frac{R_b^3}{R_0^3} \quad (18)$$

and

$$Z = 1 + \frac{2E\beta}{C_{01}R_c} (F - x_c^3)(x_c^2 - 1) \quad (19)$$

RESULTS

Our analytical model can be used to acquire quantitative information regarding the cyclic activity of an osmotically stressed liposome. Let us consider that such a giant unilamellar vesicle (GUV) has the radius in the initial state, $R_0 = 19.7 \mu\text{m}$, and the radius of the critical state, $R_c = 20.6 \mu\text{m}$.⁵ In addition, we consider a liposome that contains a solution of a nonpermeant solute. Let us consider that the initial internal solute concentration is $C_{01} = 0.5 \text{ M}$. Using experimentally determined parameters,⁴⁴ the membrane permeability coefficient for water, P_w , is equal to $3 \times 10^{-5} \text{ m/s}$ and the molecular volume of a water molecule is $V_{\mu w} = 18.04 \times 10^{-6} \text{ m}^3/\text{mol}$. A representative value of the two-dimensional stretch modulus of the lipid bilayer is $E = 0.2 \text{ N/m}$.^{29,30} Typical values for the surface tension, σ , of a phospholipid membrane in a stretched state are in the range of 0.1–1 mN/m.^{28,33} Yet, GUVs undergo much greater values of their surface tension due to their large spherical surfaces. To examine the effect of a finite water bath on a pulsatory liposome, we considered three closed spherical boxes with a radius, R_b , where R_b values are $4R_0$, $8R_0$, and $12R_0$. We formulate the following parameters that characterize the cyclic activity of a liposome: the number of cycles, the amount of solute released at the end of the cyclic liposomal activity, the duration of each cycle, and the total lifetime of the cyclic activity.

Using eq 17, we found that the total numbers of cycles for finite hypotonic environments with radii $4R_0$, $8R_0$, and $12R_0$ are 30, 42, and 46, respectively. If a liposome is placed into an infinite hypotonic environment, namely, for $C_{\text{out}} = 0$ and an indefinite duration, then the cyclic activity has 47 cycles. This is a value closely similar to the maximum cycle order of a spherical hypotonic bath of radius, $R_b = 12R_0$. This outcome was obtained using eq 17 and employing the condition $R_b \rightarrow \infty$. Theoretical predictions of this analytical model are in accord with prior experimental observations acquired by Karatekin and colleagues.^{29,30} They have experimentally detected a succession of 30–40 pores in a giant liposome tensed by intense optical illumination, each of them corresponding to a certain cycle order.

Figure 2 shows the time dependence of the swelling ratio of a pulsatory liposome, R/R_0 , for the first part of the cyclic

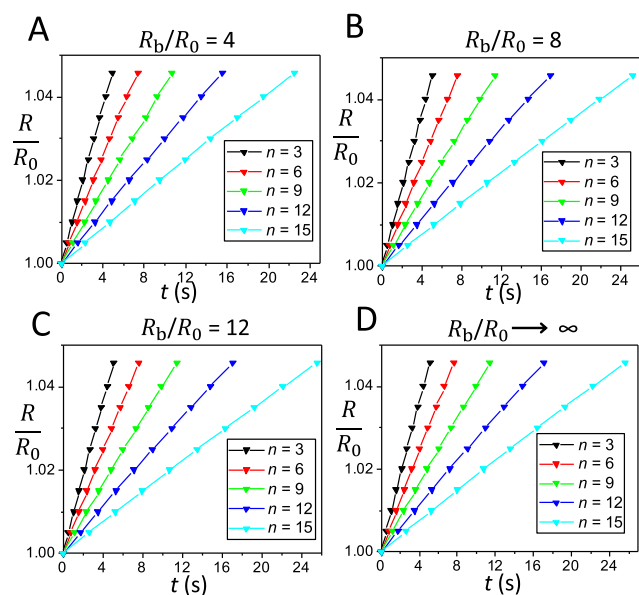


Figure 2. Time dependence of the liposomal radius during the swelling phase for cycles of varying cycle order, n , at the beginning of the cyclic activity. Different panels indicate cases when the liposome was placed within a finite water bath of radii, $R_b = 4R_0$ (A), $R_b = 8R_0$ (B), and $R_b = 12R_0$ (C), as well as when the liposome was placed within an infinite water bath, $R_b \rightarrow \infty$ (D). In (A)–(D), the cycle orders of the liposomal activity are 3, 6, 9, 12, and 15.

activity of the liposome, which includes 15 cycles. These calculations were performed for finite hypotonic spherical baths with radii $R_b = 4R_0$ (Figure 2A), $R_b = 8R_0$ (Figure 2B), and $R_b = 12R_0$ (Figure 2C), as well as for an infinite hypotonic environment ($R_b \rightarrow \infty$; Figure 2D). The swelling phase occurred at a swelling ratio, R_c/R_0 , in the range of 1–1.045. These panels show that the time dependences of the swelling phases for cycles from the beginning of the cyclic activity are fairly linear and slightly dependent on the size of the external hypotonic environment. For example, for $n = 3$, the swelling times of a pulsatory liposome were 4.90, 5.07, and 5.09 s, for the radii of the external bath $4R_0$, $8R_0$, and $12R_0$, respectively. If the liposome was inserted into an infinite hypotonic environment, then the swelling time was 5.10 s, a closely similar value to those acquired for finite hypotonic water baths. On the other hand, for $n = 15$, the swelling times were 22.46, 25.17, and 25.47 s, for the radii of the external bath of $4R_0$, $8R_0$, and $12R_0$, respectively. Yet, the swelling time was 25.60 s when the

liposome was inserted within an infinite hypotonic environment.

Figure 3 illustrates the time dependence of the swelling ratio of a pulsatory liposome for the last part of the cyclic activity,

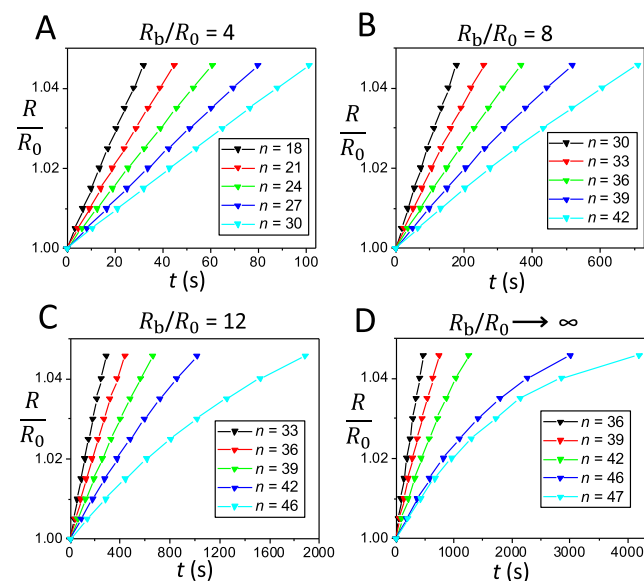


Figure 3. Time dependence of the liposomal radius during the swelling phases that correspond to the last cycles of the pulsatory activity. This time dependence is illustrated for three distinct cases of a finite hypotonic environment and an infinite hypotonic water bath, as follows: (A) $R_b = 4R_0$, cycles 18, 21, 24, 27, and 30; (B) $R_b = 8R_0$, cycles 30, 33, 36, 39, and 42; (C) $R_b = 12R_0$, cycles 33, 36, 39, 42, and 46; and (D) infinite hypotonic medium, cycles 36, 39, 42, 46, and 47.

which encompasses 12–14 cycles. This data was acquired for the cases presented in Figure 2, either for a finite spherical bath or for an infinite environment under hypotonic conditions ($R_b = 4R_0$, Figure 3A; $R_b = 8R_0$, Figure 3B; $R_b = 12R_0$, Figure 3C; $R_b \rightarrow \infty$; Figure 3D). In both cases, the finite hypotonic medium and opened water bath, the expanded liposomal radius is a linear function on the swelling time, except for the last cycles. This observation is highlighted for the case of hypotonic environments of larger sizes, e.g., for $n = 46$ when $R_b = 12R_0$ as well as for $n = 46$ and 47 when $R_b \rightarrow \infty$. Moreover, in contrast to the findings pertaining to Figure 2, the swelling phases of the last cycles were drastically slower in large external baths (e.g., for $R_b = 12R_0$ and $R_b \rightarrow \infty$) with respect to smaller finite environments (e.g., for $R_b = 4R_0$ and $8R_0$). This phenomenon was amplified at the highest cycle orders. For instance, the durations of the swelling phases were 16.95 and 31.42 min for $n = 42$ and 46, respectively, when R_b was $12R_0$ (Figure 3C). The slowest swelling phase, in the duration of 69.97 min, was obtained for $n = 47$ when a pulsatory liposome was placed within an infinite water bath (Figure 3D).

Figure 4A presents the durations of the swelling phase of the 30th cycle of pulsatory liposomes located in hypotonic water baths of varying radii. One can note a significant distinction between the swelling rate acquired for the external bath with the smallest radius and the others. If we define the swelling rate as the critical swelling ratio divided by the total swelling time, meaning $(R_c/R_0)/t_{\text{swell}}$, where t_{swell} is the duration of the swelling phase for a certain cycle order, then its value was $4.52 \times 10^{-4} \text{ s}^{-1}$ for the 30th cycle of a finite bath with $R_b = 4R_0$. In contrast, slower swelling rates of 2.56×10^{-4} , 2.37×10^{-4} , and

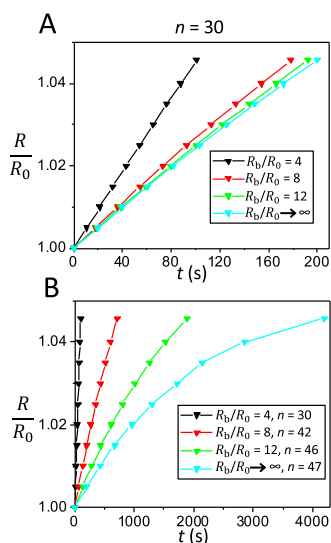


Figure 4. Comparisons of the durations of the swelling phase of the 30th cycle and final cycle for distinct sizes of the hypotonic water bath. (A) Duration of the swelling phase of the 30th cycle for the three sizes of finite hypotonic water baths ($R_b/R_0 = 4, 8,$ and 12) and for an infinite hypotonic environment. (B) Duration of the swelling phase of the last cycle for the three sizes of finite hypotonic water baths ($R_b/R_0 = 4, 8,$ and 12) and for an infinite hypotonic environment.

$2.29 \times 10^{-4} \text{ s}^{-1}$ were determined for external baths with the radii equal to $8R_0, 12R_0,$ and infinity, respectively.

Figure 4B illustrates the time dependence of the swelling ratio, R/R_0 , of a pulsatory liposome during the last cycle. The durations of the swelling phase of the last cycles were equal to 1.68, 11.84, and 31.42 min when a pulsatory liposome was inserted into finite hypotonic media with radii $4R_0, 8R_0,$ and $12R_0$, respectively. Yet, the duration of the swelling phase of the last cycle was equal to 69.97 min when the liposome was placed within an infinite hypotonic environment. These are the durations required for a pulsatory liposome to attain the critical swelling ratio, R_c/R_0 , of 1.045 during the last cycle. This observation reveals that an increase in the size of the external hypotonic bath drastically declines the swelling rate of a pulsatory liposome in the terminal cycle.

We found that the durations of the first half of the swelling phase of the last cycles (e.g., those corresponding to $R/R_0 = 0.5 \times (R_c/R_0 + 1) = 1.0228$) were equal to 0.90, 5.88, and 13.42 min when a pulsatory liposome was inserted within finite hypotonic media of radii $4R_0, 8R_0,$ and $12R_0$, respectively. However, the duration of the first half of the swelling phase of the last cycle was equal to 21.77 min when the liposome was placed within an infinite hypotonic bath. Notably, in this case, the duration of the second half of the swelling phase of the last cycle was equal to 48.19 min, a significantly longer value than that of the first half of the swelling phase. This outcome results from a strong nonlinear time dependence of the liposomal swelling ratio in the last cycle of an infinite external bath (Figure 4B, $F \rightarrow \infty, n = 47$).

Figure 5 shows the duration of the swelling phase as a function of the cycle order for hypotonic baths of smaller sizes (Figure 5A) and for larger environments (Figure 5B). It is clear that cycles of low-order numbers ($n < 15$) are significantly shorter than those of high-order numbers ($n > 30$). The duration of the swelling phase drastically increased to tens of

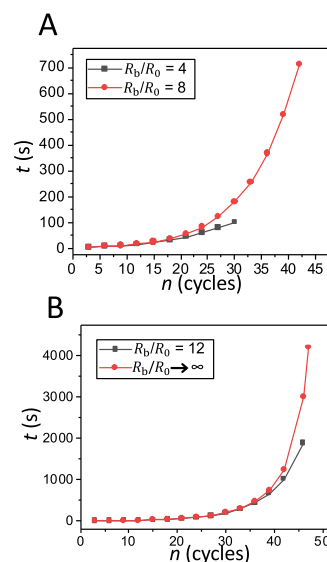


Figure 5. Duration of the swelling phase as a function of the cycle order for a pulsatory liposome. (A) Comparison of hypotonic water baths with $R_b = 4R_0$ and $8R_0$. (B) Comparison of hypotonic water baths with $R_b = 12R_0$ and $R_b \rightarrow \infty$.

minutes for cycles with an order number greater than 40. On the contrary, the total swelling time for the maximum cycle order was only 100.95 s in the case of the smallest hypotonic bath.

If the pulsating liposome functions as a device for the controlled release of a chemical agent, then the most important parameter is the amount of solute released into the external environment during its functional operation. Let us assume that the initial internal concentration of the solute is equal to 0.5 M, so the liposome contains $Q_{01} = C_{01}V_0 = 9.643 \times 10^{12}$ molecules. At the end of the cyclic activity, a pulsatory liposome releases the amounts of solute equal to $Q_{30} = 9.4997 \times 10^{12}$ molecules, $Q_{42} = 9.6083 \times 10^{12}$ molecules, and $Q_{46} = 9.6227 \times 10^{12}$ molecules, if this liposome is placed within closed hypotonic environments with radii $R_b = 4R_0, 8R_0,$ and $12R_0$, respectively. For the case of an infinite hypotonic bath, the amount of delivered solute is equal to $Q_{47} = 9.625 \times 10^{12}$ molecules, which is $\sim 99.81\%$ of the initial solute amount. We highlight that the amount of solute released during a single cycle of the same order is unchanged regardless of whether a pulsatory liposome is placed into a finite or an infinite hypotonic environment.

DISCUSSION

In the Results section, we provided a detailed quantitative description of the swelling phase of a pulsatory liposome when this is placed within a hypotonic bath of varying size. The energy required for the liposomal swelling is provided by the osmotic transmembrane gradient. The swelling phase is an active process of each cycle and is described by a differential equation for the radius of the liposome (eq 8). In contrast, the liposomal relaxation is a passive and fast process,^{38,39} resulting from the appearance of a transient pore of a pulsatory liposome at a point that reaches a critical radius, R_c , and a critical surface tension, σ_c . The appearance of the transient pore is a critical event for the cyclic activity of the liposome because at this point the swelling phase stops and the relaxation phase starts.

After this event, some internal solute is released into the external hypotonic bath.

The lifetime of the transient pore is ~ 1.725 s.⁴⁵ This value was previously inferred using similar conditions to those employed in this study. The transient pore undergoes two phases, the expansion and shrinkage phases. The durations of these phases are 225 and 1500 ms, respectively. The lifetime of the pore is independent of the cycle order of the pulsatile liposome. Kinetics of the two phases of the transient pore depends on the properties of the liposomal membranes. Because of the very short duration of the transient pore, we expect that this parameter does not significantly affect the overall dynamics of the liposomal cyclic activity.

The pore energy, E_p , is given by the following formula⁴⁶

$$E_p = 2\pi r\gamma - \sigma\pi r^2 \quad (20)$$

This shows the balance between the line tension and surface tension. Here, r denotes the pore radius. γ and σ are the line tension and surface tension coefficients, respectively. The only stable energetic minimum of E_p is reached at a radius $r = 0$. The transient pore undergoes membrane rupture at a pore radius greater than a critical radius, $r_c = \gamma/\sigma$. However, this phenomenon is rare because the activation free energy for this process is high ($\Delta E = \pi\gamma^2/\sigma$).⁴⁷ For smaller tensions, a transient pore can rapidly shrink, evolving toward a closed pore state. On the contrary, for tensions greater than a critical value, a transient pore can rapidly expand until it reaches membrane rupture, leading to liposomal lysis. Therefore, it is not possible to detect stable transmembrane pores with long lifetimes. Yet, Zhelev and Needham have demonstrated for the very first time the appearance of long-lived quasi-stable transmembrane pores with a radius of $1 \mu\text{m}$ in GUVs.⁴⁸ A giant liposome was aspirated into a micropipette and short-lived electrical stimuli were applied across the vesicle. In this way, the lifetime of relatively stable pores spanned a duration range from tenths of a second to several seconds. These studies enabled the determination of the pore size and pore line tension. In parallel with these studies, Wilhelm and co-workers have studied the kinetics of pore appearance followed by mechanical rupture using high-voltage, charge-pulse technique.⁴⁹ A large number of theoretical,^{4,5,7,10,11,14-24,28-30,46} computational,^{7,16,21,23,25-27,50} and experimental^{5,11,28-32} studies have documented the existence of these transient pores across bilayer membranes.

Our analytical model involves a spherical symmetry. Yet, in a given experimental context, ample alterations in external mechanical stress can lead to membrane deformations⁵¹⁻⁵³ and thickness fluctuations.⁷ We considered the case of a spherical liposome because this geometry represents the equilibrium conformation with enhanced shape stability of the membrane. This also means that the membrane tension, σ , has a uniform value across the entire liposomal surface. If transient undulations occur,^{50,54} then the amplitude of these membrane motions would normally decrease during the liposomal swelling phase. Quantitative assessment of local and global membrane undulations, some of which might be thermally activated, is experimentally challenging. The spherical geometry represents the average experimental substate of the undulatory dynamics of the liposomal membrane.^{2,36,37} It is also a reason for which the outcomes of our analytical model quantitatively recapitulate experimental results (see below). However, if an amplified out-of-membrane

pressure occurs, this would lead to stable membrane deformations. In this case, the membrane tension, σ , has no longer a uniform value across the entire liposomal membrane. In addition, the swelling phase of the liposome is no longer described by a single differential equation. The transient transmembrane pore appears on the liposomal membrane, where the critical surface tension, σ_c , is reached. Moreover, the liposome may feature a cyclic activity when stable membrane deformations take place.

Here, we highlighted the quantitative aspects of the swelling phases of the cycles of a pulsatory liposome under finite hypotonic conditions for one major reason. Because the duration of the relaxation phase is very short, the alteration in the internal concentration of the solute is negligible during this process. Indeed, recent experimental studies of the cyclic activity of GUVs provided evidence for very short-lived relaxation phase^{38,39} and unchanged initial radius of the vesicle, R_0 , of a pulsatory liposome for successive cycles.^{2,33} Yet, there is experimental evidence that indicates fluctuations in the critical radius, R_c , of a pulsatory liposome for successive cycles.^{2,33} This finding suggests a more complex dependence of the lytic tension on the cycle order, which was not included in our analytical model (see [Methods](#)). Indeed, Evans and co-workers,⁵⁵ as well as Chabanon and co-workers,² have shown that the lytic tension for the pore formation is a function that depends on the load rate. Therefore, we expect that the amplitude of R_c and the time of pore appearance are affected by the cycle order. If the lytic tension depends on the cycle order, then the corresponding critical value of liposome radius, R_c , is introduced in [eqs 14 and 15](#). Dependence of the lytic tension on the load rate will certainly affect the swelling phase of the liposome. Therefore, this will affect the overall duration of large-order cycles. If the liposomes are exposed to a surfactant-containing environment, then the partitioning of surfactant molecules into the membrane likely changes the lytic tension at which short-lived pores occur.³⁶

The periodic behavior of osmotically stressed liposomes has been first predicted by Koslov and Markin.⁹ Later, other groups have developed theoretical approaches for acquiring quantitative information on this fundamental process.^{1,38} Our theoretical formulation includes low-amplitude size fluctuations of the swelling and relaxation phases between R_0 and R_c , which are characterized by a critical liposomal radius of $\sim 4.5\%$ out of that value of an unstretched lipid vesicle. This assumption is in good accord with the cyclic size fluctuations of GUVs, as previously reported.^{3,5,39} Our model quantitatively recapitulates the number of the swelling–relaxation cycles observed experimentally. For example, we acquired a total number of cycles, n_{max} of 47 for our infinite hypotonic bath. This value is close to the numbers of swell/burst cycles noted with GUVs in hypotonic media.³⁸ Furthermore, our model calculations predict long-lived swelling phases of larger-order number cycles, which is in quantitative agreement with recently reported experimental determinations of osmotically stressed lipid vesicles.^{2,37}

In summary, we formulated an analytical model that provides quantitative information on the cyclic activity of a lamellar liposome in a finite hypotonic environment. The swelling phase of the last several cycles of a pulsatory liposome is strongly dependent on the size of the external bath. A reduced number of cycles of a pulsatory liposome, as well as a lower amount of chemical agent released into the external environment, were found by decreasing the size of the external

hypotonic medium. Furthermore, the duration of the swelling phase is strongly dependent on the cycle order, which is in good accord with recent experimental explorations of the pulsatory GUVs.^{2,33} For example, the swelling phases of the last few cycles last several hundreds of seconds, contrasting the cycles of low-order numbers, which are in the second range. From a practical point of view, the consumed fuel of this two-stroke liposomal engine might be substituted by an active biotherapeutic agent. Finally, there is great potential for the use of lipid vesicles in clinical applications, specifically for controlled drug release of targeted chemicals at precise locations of diseased tissues. In this case, the osmotically stressed liposomes would be located within a finite hypotonic environment.

■ ASSOCIATED CONTENT

SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.langmuir.9b03923>.

Determination of the final state of the first cycle of a pulsatory liposome, where the law of mass conservation applies during the first cycle of the swelling process; and determination of the swelling phase of the n -th cycle of a pulsatory liposome (PDF)

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Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

The authors are grateful to Lauren Mayse and Jiaxin Sun for their comments and stimulating discussions. This work was supported by the U.S. National Institutes of Health grants GM088403 (to L.M.) and GM129429 (to L.M.).

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