Dynamics of receptor-mediated nanoparticle internalization into endothelial cells.
David Gonzalez-Rodriguez, Abdul I Barakat

To cite this version:
David Gonzalez-Rodriguez, Abdul I Barakat. Dynamics of receptor-mediated nanoparticle internalization into endothelial cells.. PLoS ONE, Public Library of Science, 2014, 10 (4), pp.e0122097. 10.1371/journal.pone.0122097. hal-01178880

HAL Id: hal-01178880
https://hal-polytechnique.archives-ouvertes.fr/hal-01178880
Submitted on 21 Jul 2015

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L’archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d’enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.
Dynamics of Receptor-Mediated Nanoparticle Internalization into Endothelial Cells

David Gonzalez-Rodriguez*, Abdul I. Barakat*

Laboratoire d’Hydrodynamique (LadHyX), École Polytechnique, CNRS UMR 7646, Palaiseau, France

* davidgr@alum.mit.edu (DGR), barakat@ladhyx.polytechnique.fr (AIB)

Abstract

Nanoparticles offer a promising medical tool for targeted drug delivery, for example to treat inflamed endothelial cells during the development of atherosclerosis. To inform the design of such therapeutic strategies, we develop a computational model of nanoparticle internalization into endothelial cells, where internalization is driven by receptor-ligand binding and limited by the deformation of the cell membrane and cytoplasm. We specifically consider the case of nanoparticles targeted against ICAM-1 receptors, of relevance for treating atherosclerosis. The model computes the kinetics of the internalization process, the dynamics of binding, and the distribution of stresses exerted between the nanoparticle and the cell membrane. The model predicts the existence of an optimal nanoparticle size for fastest internalization, consistent with experimental observations, as well as the role of bond characteristics, local cell mechanical properties, and external forces in the nanoparticle internalization process.

Introduction

The pathological complications of atherosclerosis, namely heart attacks and strokes, remain the leading cause of worldwide mortality [1]. Because atherosclerosis is fundamentally a disease that involves inflammation of the endothelium, the monolayer of cells that lines the inside of blood vessels, a particularly promising idea is the use of nanoparticles as cargo vehicles for targeted delivery of anti-inflammatory agents to arterial endothelial cells. Recent studies have established that nanoparticle internalization into endothelial cells depends on a number of factors including nanoparticle size and surface functionalization [2–4]. Elucidating the basis for these observations is of primary interest.

A critical component in the development of an effective nanoparticle-based endovascular drug delivery system is the interaction between particles and the endothelial cell surface. To specifically target inflamed endothelial cells at an atherosclerotic lesion, nanoparticles can be coated with antibodies against endothelial cell adhesion molecules, such as selectins, VCAM-1, PECAM-1, or ICAM-1 [5]. Out of these different receptors, intercellular adhesion molecule-1 (ICAM-1) is a particularly relevant target, since its level of expression in vascular endothelial
cells is enhanced significantly by pathological stimuli such as oxidants, cytokines, and abnormal fluid mechanical shear stresses [3]. Specifically, a 20 to 100 fold increase in ICAM-1 expression in activated over quiescent cells has been reported [6]. ICAM-1-mediated nanoparticle internalization into endothelial cells has been the subject of a number of recent experimental studies [3, 6–10]. Nanoparticles coated with anti-ICAM-1 antibodies activate a specific endocytosis pathway termed CAM-mediated endocytosis [7]. Unlike endocytosis mediated by other membrane receptors, CAM-mediated endocytosis requires multivalent binding: a single anti-ICAM-1 antibody is not internalized by an endothelial cell, whereas a particle carrying several antibodies can be internalized. CAM-mediated endocytosis is actin-dependent, but it involves different protein machineries than clathrin-mediated endocytosis, caveoli, macropinocytosis, or phagocytosis [3, 7].

In this article we develop a mathematical model to describe receptor-mediated nanoparticle internalization, specifically considering the case of ICAM-1-mediated endocytosis. Several theoretical models of nanoparticle internalization have previously been proposed. A first group of theoretical models describes receptor-mediated internalization of spherical and non-spherical particles limited by diffusion of receptors within the cell membrane [11–13]. These models assume the particle ligand density to be much larger than the cell membrane receptor density, thus making receptor diffusion towards the particle wrapping zone a limiting physical mechanism. This assumption does not appear applicable to ICAM-1-mediated nanoparticle endocytosis into inflamed endothelial cells, where receptor and ligand densities are both of the order of 1000 molecules/\(\mu\)m\(^2\) [14], and diffusion of receptors thus becomes negligible. Another group of theoretical models uses energetic approaches to describe membrane wrapping of a nanoparticle (see the recent review by Bahrami et al. [15]). These models have investigated the role of nanoparticle shape and orientation [16, 17], nanoparticle deformability [18], and interactions between multiple nanoparticles [19, 20], but they often leave out the cell’s cytoplasmic rigidity and the bond formation dynamics, as well as the kinetics of the wrapping process. Recent advances in modeling the wrapping of a nanoparticle by a membrane have incorporated stochastic thermal fluctuations to study the kinetics of wrapping [21] or conformational changes of membrane proteins, which are described by particle dynamics simulations [22].

Here we develop a new theoretical model to study the kinetics of nanoparticle internalization under the following premises: (i) we consider the case where receptor and ligand density are comparable, so we neglect receptor diffusion; (ii) we include both membrane bending and viscoelastic deformation of the cytoskeleton; (iii) we account for the dynamics of bond formation under force. Unlike most of the previous theoretical models, which are based on energy formulations, we develop our model in terms of force balances. Our formulation is inspired by that proposed by Dembo et al. to study the kinetics of detachment of a membrane from a surface [23]. Our model enables us to understand how receptor-mediated internalization is affected by particle size, bond characteristics, cell mechanical properties, and external forces exerted on the nanoparticle.

**Model**

Our conceptualization of receptor-mediated nanoparticle internalization is represented in Fig 1. We consider a three-dimensional axisymmetric geometry. Internalization is driven by the formation of bonds between ligands on the particle surface (the total density of free plus bound ligands is denoted by \(\xi_l\)) and receptors on the cell membrane (the total density of free plus bound receptors is denoted by \(\xi_r\)). The density of bonds is denoted by \(\xi_b\), which is a function of the curvilinear coordinate \(s\) and of the elapsed time \(t\). The particle radius is denoted by \(a\). The model aims to predict the evolution of the membrane shape, determined by the variables \(R(s, t)\)
Mechanically, the cell membrane deformation is described as the bending of an inextensible axisymmetric plate subjected to large deformations. Force equilibria in the directions tangential and normal to the membrane yield

\[
\frac{\partial T}{\partial s} - V \frac{\partial \theta}{\partial s} + \sigma_t = 0 \quad (1)
\]

\[
T \frac{\partial \theta}{\partial s} + \frac{\partial V}{\partial r} + T \sin \theta \frac{V \cos \theta}{r} + \sigma_n = 0, \quad (2)
\]

where \(T\) and \(V\) are respectively the shear and normal forces per unit of azimuthal length of the membrane, \(\theta\) is the angle between the membrane and the horizontal, measured in the radial plane (see Fig 1), \(r\) is the radial distance to the axis of symmetry, \(s\) is the curvilinear coordinate, and \(\sigma_t\) and \(\sigma_n\) are respectively the tangential and normal stresses acting on the membrane. In deducing these equations, we have assumed that, due to reorganization of the lipid molecules composing the membrane, the membrane tension remains isotropic, thus its value is the same in the radial and azimuthal directions.

Moment equilibrium yields

\[
V = \frac{\cos \theta}{r} (M_s - M_\theta) + \frac{\partial M_r}{\partial s}, \quad (3)
\]

where \(M_s\) and \(M_\theta\) are the moments in the radial and azimuthal directions. Assuming linear elastic membrane deformation, assuming that a planar cross-section of the membrane remains planar after bending, and neglecting the mean circumferential deformation of the cell membrane (i.e., neglecting stresses due to membrane stretching), we deduce the following relations
between the moments and the membrane shape:

\[ M_s = -B \left( \frac{\partial \theta}{\partial s} + \frac{\nu}{r} \sin \theta \right) \]  

(4)

\[ M_\theta = -B \left( \frac{\sin \theta}{r} + \nu \frac{\partial \theta}{\partial s} \right), \]  

(5)

where \( B = E_m h^3/(12(1-\nu^2)) \) is the membrane bending modulus, \( h \) is the membrane thickness, \( E_m \) is the Young’s modulus of the membrane, and \( \nu \) its Poisson’s ratio. Using Eqs 3–5, the force equilibria given by Eqs 1 and 2 can be rewritten in terms of two unknowns, \( \theta \) and \( T \). For reasons of numerical stability, we project the force equilibrium equations onto the radial and vertical axes. The projected equations are:

\[
\begin{align*}
\cos \theta \frac{\partial T}{\partial s} - T \sin \theta \frac{\partial \theta}{\partial s} - \frac{T \sin^2 \theta}{r} + B \left[ \sin \theta \frac{\partial^3 \theta}{\partial s^3} \right] \\
+ \frac{2 \sin \theta \cos \theta \partial^2 \theta}{r} + \cos \theta \frac{\partial \theta \partial^2 \theta}{\partial s \partial s^2} \\
+ \cos (2\theta) \left( \frac{\partial \theta}{\partial s} \right)^2 - \frac{\sin \theta (\sin^2 \theta - 2 \cos^2 \theta) \partial \theta}{r^2} \\
+ \frac{\sin^2 \theta \cos \theta}{r^3} \right] + \sigma_r = 0 
\end{align*}
\]  

(6)

and

\[
\begin{align*}
\sin \theta \frac{\partial T}{\partial s} + T \cos \theta \frac{\partial \theta}{\partial s} + \frac{T \sin \theta \cos \theta}{r} + B \left[ -\cos \theta \frac{\partial^3 \theta}{\partial s^3} \right] \\
- \frac{2 \cos^2 \theta \partial^2 \theta}{r} + \sin \theta \frac{\partial \theta \partial^2 \theta}{\partial s \partial s^2} \\
+ \sin (2\theta) \left( \frac{\partial \theta}{\partial s} \right)^2 + \frac{\cos \theta (\cos^2 \theta - 2 \sin^2 \theta) \partial \theta}{r^2} \\
- \frac{\sin \theta \cos^3 \theta}{r^3} \right] + \sigma_v = 0, 
\end{align*}
\]  

(7)

where \( \sigma_r \) and \( \sigma_v \) are respectively the radial and vertical stresses acting on the membrane.

**Stresses exerted on the membrane**

Stresses exerted on the membrane arise from interactions with the nanoparticle and with the cell’s cytoplasm, as represented in Fig 2. We consider three types of stresses acting on the cell membrane. The first type are the stresses due to viscoelastic deformation of the cytoplasm. The cytoplasm is treated as a Kelvin-Voigt material, i.e., a system comprising an elastic spring, of constant \( K = E/a \), in parallel with a damper, of constant \( M = \mu/a \). Here, \( E \) and \( \mu \) are respectively the Young’s modulus and the viscosity of the cytoplasm, and \( a \) is the particle radius. The cytoplasm is assumed to exert a purely vertical stress on the membrane. The model does not account for the active forces generated by cytoskeletal dynamics, such as those due to actin polymerization, which develop over a time scale of several minutes [3]. Thus, strictly speaking,
the model describes only the early stages of internalization, up to the first few minutes. The sec-
ond type of stresses are due to the deformation of the bonds between the membrane and the
particle. The bonds are treated as linear elastic springs of constant $\kappa$ and undeformed
length $\lambda$. In addition, if the membrane and the particle get very close, a short-range repulsion force (not
represented in the figure) prevents particle-membrane interpenetration. The membrane’s bottom face
sustains a stress arising from the viscoelastic deformation of the cell’s cytoplasm, which is modeled as a
Kelvin-Voigt material of elastic constant $K = E/a$ and viscous constant $M = \mu/a$.

Fig 2. Schematic of the stresses exerted on the membrane. The membrane’s top face sustains a stress
$\sigma_b$ arising from bond deformation. The bonds behave as linear elastic springs of constant $\kappa$ and undeformed
length $\lambda$. Bond orientation is assumed to point towards the nanoparticle center. We denote by $\alpha$ the
angle between the bond direction and the vertical (see Fig 2). The total stress exerted locally by
the bonds is proportional to the local bond density, $\xi_b$. The bonds can exert larger traction
forces than compression forces, since the latter are limited by the bond length remaining posi-
tive, $l \geq 0$. Therefore, to avoid membrane-particle interpenetration, we need a third type of
stresses that can provide sufficiently large contact forces between the particle and the mem-
bane. We thus introduce a short-range repulsion force using the formulation proposed by
[24]. Adding these three components, the total radial and vertical stresses on the membrane are:

$$
\sigma_r = -\xi_b \kappa (l - \lambda) \sin \alpha 
+ \frac{\gamma}{l} \left( \frac{1}{l + \frac{1}{\tau}} \right) \exp \left\{ -\frac{l}{\tau} \right\} \sin \alpha 
$$

(8)

$$
\sigma_v = \left( -KZ - M \frac{\partial Z}{\partial t} \right) \cos \theta 
+ \xi_b \kappa (l - \lambda) \cos \alpha 
- \frac{\gamma}{l} \left( \frac{1}{l + \frac{1}{\tau}} \right) \exp \left\{ -\frac{l}{\tau} \right\} \cos \alpha,
$$

(9)
where $Z$ is the membrane elevation with respect to the undeformed position, $t$ is the time, $l$ is the distance between the membrane and the particle, measured in the direction pointing towards the particle center, and $\gamma < 1$ nN and $\tau / C^2 \approx 5 - 30$ nm are constants in the short-range repulsion force formulation [24].

**Membrane-particle bonds**

The evolution of the bond density at a given membrane position, $s$, and time, $t$, is computed using Bell’s model:

$$\frac{\partial \xi_b}{\partial t} = K_f (\xi_1 - \xi_b)(\xi_r - \xi_b) - K_d \xi_b,$$

where $\xi_b(s, t)$ is the instantaneous, local bond density, $\xi_1$ and $\xi_r$ are respectively the total ligand and receptor densities, and $K_f$ and $K_d$ are respectively the bond formation and dissociation coefficients. Using the formalism proposed by [25], we assume $K_f$ to be constant and $K_d$ to depend on the force applied to the bonds through:

$$K_d = K_d^{(0)} \exp \left\{ \frac{|l| - \lambda x_b}{k_B T} \right\},$$

where $K_d^{(0)}$ is the baseline value at zero force, $k_B \approx 1.38 \cdot 10^{-23}$ J/K is the Boltzmann constant, $T = 310$ K is the temperature, $x$ is the bond spring constant, and $x_0$ is a constant length that characterizes the height of the energy barrier needed to break a bond [25, 26]. Eq 11 assumes the dissociation constant to depend on forces applied to the bonds irrespective of their sign (tension or compression). The rationale underlying this assumption is that there is an optimal distance $\lambda$ for ligand-receptor interaction, whereas when receptors and ligands are compressed into closer distances to each other, they may be forced into conformations that are suboptimal for the interaction between their respective binding sites. Thus, we expect the dissociation constant to increase with an applied compressive force, as is the case under a tensile force.

**Particle position**

The vertical particle position with respect to the membrane, $Z_0$, is determined by balancing the forces acting on the particle: the bond forces, the membrane-particle repulsive forces, and an external vertical force that may be exerted directly on the particle, $F_v$. Vertical force equilibrium yields:

$$F_v - \int_0^\infty \left[ \xi_b K(l - \lambda) \cos \alpha \right. $$

$$- \frac{\gamma}{l} \left( \frac{1}{l} + \frac{1}{l + \tau} \right) \exp \left\{ \frac{-l}{l + \tau} \right\} \sin \alpha \right] 2\pi rdr = 0.$$

The equation is implicit in $Z_0$, since $l$ and $\alpha$ depend on $Z_0$ through straightforward geometric relations.

**Model parameters and nondimensionalization**

The model parameters and their typical values are summarized in Table 1. The bond characteristics correspond to values found in the literature for ICAM-1 receptor-antibody pairs. The tabulated binding constants, $K_f$ and $K_d^{(0)}$, are typical values for membrane-bound ICAM-1 receptors and surface-adsorbed anti-ICAM-1 antibody ligands. These values are estimated from the
experimental results by [27], which, it should be noted, were performed using a soluble form of ICAM-1 and surface-adsorbed anti-ICAM-1 antibodies. To deduce the tabulated value range for $K_f$ and $K_0 d$, we apply the relationship between soluble and surface binding kinetics proposed by [25]. To apply Bell’s formulas, we assume a diffusion coefficient for soluble ICAM-1 of $D_s = 50 \mu m^2$/s [25], a diffusion coefficient for membrane-bound ICAM-1 in the range $D_m = 0.01 - 0.4 \mu m^2$/s [28, 29], and a maximum distance for receptor-ligand interaction of 1 nm [25]. The value of the repulsion constant $\gamma$ is chosen so that repulsion becomes negligible (compared to bond elasticity) at a distance equal to the bond’s equilibrium length, $\lambda$. This leads us to set $\gamma = 1 fN$. We note that our model results are insensitive to changes in $\gamma$ of up to one order of magnitude.

The equations are nondimensionalized using the particle radius, $a \approx 100$ nm, as the unit of length and the ligand density $\xi_l \approx 1000$ molecules/$\mu m^2$, as the unit of bond density. Since internalization is driven by bond formation, the unit of stress is taken as $\xi_l k_B T / a \approx 10^{10}$ Pa, which is the bond stress corresponding to a characteristic bond elongation of $a k_B T / (a \gamma x_b)$. There are two time scales, given by the rate of bond formation and by the rate of viscoelastic cytoplasmic deformation. Since the cytoplasm’s viscoelastic time scale, $\mu / E \approx 1$ s, is usually longer than the bond formation time scale, $1 / (K_f \xi_l) \approx 0.1$ s, the former is chosen as the unit of time for nondimensionalization. By nondimensionalizing the equations with the aforementioned units of length, bond density, stress, and time, we deduce the nondimensional parameters characterizing the problem, which are listed in Table 2. The tabulated values show that the three forces resisting deformation, due to cell viscoelasticity ($C_E$), membrane bending ($C_B$), and membrane tension ($T_0$), are of comparable magnitude, and therefore they should all be accounted for in the model. The nondimensional parameter $t_b$ is the ratio of the bond formation time to the viscoelastic time, which is typically smaller than 1, since bond formation is fast compared to viscoelastic deformation.

**Numerical solution**

We need to solve the system formed by Eqs 6 and 7, which determine the instantaneous membrane shape; Eq 12, which determines the instantaneous particle position; and Eq 10, which

| parameter          | symbol | best estimate | range            | reference |
|--------------------|--------|---------------|------------------|-----------|
| particle radius    | $a$    | 0.1 $\mu m$   | 0.05 – 2         | [3]       |
| ligand density     | $\xi_l$| $10^3 \mu m^{-2}$ | 300 – 7000      | [14]      |
| receptor density   | $\xi_r$| $10^3 \mu m^{-2}$ | 100 – 5000      | [8, 14]   |
| bond elasticity    | $\kappa$| $10^{-2} N/m$ | $10^{-5} – 1$   | [14, 24, 38, 39] |
| bond length        | $\lambda$| 20 nm        | 15 – 35         | [14, 38, 40] |
| repulsion constant | $\gamma$| 1 fN         | $0 – 10^6$      | [24]      |
| repulsion length   | $\tau$ | 5 nm          | 5 – 20           | [24]      |
| binding rate       | $K_f$  | $10^{-2} \mu m^2$/s | 0.002 – 0.06   | [27]      |
| bond equilibrium constant | $K_{eq} / K_f$ | 0.05 $\mu m^{-2}$ | 0.02 – 0.10   | [27]      |
| Bell’s length constant | $x_b$  | 0.5 nm        | 0.1 – 1.0        | [25, 41] |
| membrane tension   | $T_0$  | 30 pN/$\mu m$ | 10 – 50         | [42–44]  |
| cytoplasm elasticity| $E$    | 1 kPa         | 1 – 7            | [45, 46] |
| cytoplasm viscosity| $\mu$  | 1 kPa · s     | 0.5 – 6          | [47–49]  |
| membrane bending   | $B$    | $10^{-19}$ J  | $10^{-20} – 10^{-19}$ | [11, 23] |
determines the evolution of the bond density. The equations are solved iteratively in time. Initially \((t = 0)\), the membrane is assumed flat \((R(s, 0) = 0, Z(s, 0) = 0)\), the particle is located above the membrane at \(Z_0(t = 0) = a + \lambda_s\), and the particle is assumed to be weakly attached to the membrane through an initial bond density \(\xi_0(s, 0) = 0.01\xi_1(1 - 5s/a)\) for \(s < 0.2a\), and \(\xi_0(s, 0) = 0\) otherwise. Provided that the initial bond distribution is weak, and due to swift bond formation at early times, the precise structure of the initial bond distribution has virtually no effect on the model results presented here. At each time step, \(t + \Delta t\), we start by updating the bond density. With the membrane shape and particle position from the previous time, \(t\), we analytically solve Eq 10 to compute the updated bond density, \(\xi(s, t + \Delta t)\). The time increment \(\Delta t\) is variable, and it is chosen by the solver so that the change in bond density is smaller than given relative (2%) and absolute (0.02) thresholds. The values of these thresholds have been chosen so that further reductions of the thresholds do not alter the final results by more than 5%. The time steps obtained in this manner are very small at early times (of the order of \(10^{-6}\) time units), when bond formation progresses rapidly, and significantly larger at late times (of the order of 0.05 time units), when the internalization process approaches the final equilibrium state. Once the updated bond density at \(t + \Delta t\) has been obtained, the new membrane shape and particle position are calculated by solving Eqs 6, 7, and 12. To obtain the membrane shape, axial symmetry conditions are imposed at the membrane center \((s = 0)\): \(R = 0, \theta = 0,\) and \(\partial^2 \theta/\partial s^2 = 0\). At the outer edge of the membrane, \(s = s_{\text{max}}\), the following boundary conditions are imposed: \(Z = 0, M_s = 0, T \cos \theta - V \sin \theta = T_o\). We note that the radial position of the outer edge, \(R(s_{\text{max}})\), is left free; it will decrease during the wrapping process to satisfy the membrane inextensibility assumption. The outer edge is taken as \(s_{\text{max}} = 20a\). We have verified that further increasing the value of \(s_{\text{max}}\) does not affect the simulation results. Moreover, we also verify that the membrane has recovered its planar horizontal shape at \(s = s_{\text{max}}\), i.e., that \(dZ/ds \approx 0\). Eqs 6 and 7 are linearized using a Newton-Raphson iteration scheme, then discretized in space and time using a finite difference scheme. The time step is taken equal to \(\Delta t\) defined above. The spatial grid is not uniform. Rather, the grid size is given by \(\Delta s = 0.04 a/g(s)\), with \(g(s) = 1 + 10 \exp \left(-s^2/(2a^2)\right)\). This relation yields a grid size that is about 10 times finer under the particle center \((\Delta s(s \rightarrow 0) \approx 4 \cdot 10^{-3} a)\) than far from the particle \((\Delta s(s_{\text{max}}) \approx 4 \cdot 10^{-2} a)\). We have verified that further grid refinement does not change the final results by more than 5%. The results are advanced in time following this scheme until the membrane shape converges to a final state. Once the updated bond density at \(\Delta t\) has been obtained, the new membrane shape and particle position are calculated by solving Eqs 6, 7, and 12. To obtain the membrane shape, axial symmetry conditions are imposed at the membrane center \((s = 0)\): \(R = 0, \theta = 0,\) and \(\partial^2 \theta/\partial s^2 = 0\). At the outer edge of the membrane, \(s = s_{\text{max}}\), the following boundary conditions are imposed: \(Z = 0, M_s = 0, T \cos \theta - V \sin \theta = T_o\). We note that the radial position of the outer edge, \(R(s_{\text{max}})\), is left free; it will decrease during the wrapping process to satisfy the membrane inextensibility assumption. The outer edge is taken as \(s_{\text{max}} = 20a\). We have verified that further increasing the value of \(s_{\text{max}}\) does not affect the simulation results. Moreover, we also verify that the membrane has recovered its planar horizontal shape at \(s = s_{\text{max}}\), i.e., that \(dZ/ds \approx 0\). Eqs 6 and 7 are linearized using a Newton-Raphson iteration scheme, then discretized in space and time using a finite difference scheme. The time step is taken equal to \(\Delta t\) defined above. The spatial grid is not uniform. Rather, the grid size is given by \(\Delta s = 0.04 a/g(s)\), with \(g(s) = 1 + 10 \exp \left(-s^2/(2a^2)\right)\). This relation yields a grid size that is about 10 times finer under the particle center \((\Delta s(s \rightarrow 0) \approx 4 \cdot 10^{-3} a)\) than far from the particle \((\Delta s(s_{\text{max}}) \approx 4 \cdot 10^{-2} a)\). We have verified that further grid refinement does not change the final results by more than 5%. The results are advanced in time following this scheme until the membrane shape converges to a final state.

### Table 2. Nondimensional coefficients and typical values.

| parameter | definition | typical value |
|-----------|------------|--------------|
| \(\xi_1\) | \(\xi_1/\xi_0\) | 1 |
| \(K_s\) | \(K_s^{(s)}/(K_s^{(s)}/\xi_1)\) | \(5 \cdot 10^{-5}\) |
| \(\lambda\) | \(\lambda/a\) | 0.2 |
| \(\xi\) | \(\xi/a\) | 0.05 |
| \(C_{\xi}\) | \(E_{\xi}/(\kappa /k_{B} T)\) | 10 |
| \(C_{\xi}\) | \(B_{\xi}/(\kappa /a^3 /k_{B} T)\) | 1 |
| \(\xi_{\text{rep}}\) | \(\alpha_{\kappa} /(k_{B} T)\) | 100 |
| \(T_{\text{rep}}\) | \(T_{\text{rep}}/\kappa /a^3 /k_{B} T\) | 3 |
| \(C_{\kappa}\) | \(\kappa a_{\kappa} /k_{B} T\) | 100 |
| \(t_0\) | \(E/(\mu K_s \xi_0)\) | 0.1 |

DOI:10.1371/journal.pone.0122097.t002
equilibrium wrapping state, which is typically reached in less than 40 time units. The solution algorithm is implemented in Matlab R2012a.

Results

Fig 3 illustrates the typical internalization kinetics predicted by the model. Fig 3a shows how the internalization process, quantified through the ratio of the particle’s penetration depth \( d \) to the particle diameter \( 2a \) (see figure inset), evolves in time. The process first advances very rapidly, and then it slows down to asymptotically reach a maximum particle penetration, at which an equilibrium is reached and internalization stops. At this final state, the energy gained in bond formation by further wrapping of the particle does not compensate the energetic cost to further bend the membrane and deform the cytoplasm. The final \( d/(2a) \) value depends thus on the local cell deformability, as we will discuss below. Fig 3b–3e show snapshots of the internalization process. We can distinguish three phases. The early phase, up to snapshot 3c, is a short phase characterized by rapid bond formation and development of firm adhesion between the particle and the cell membrane. In this early phase, the bond density is not yet sufficiently high to fully bend the membrane around the particle, and thus we observe bonds that are both under strong compression (dark blue) and under strong tension (red; see snapshot 3b). The second phase, between snapshots 3c and 3d, is characterized by a relatively constant density of bonds along the particle-membrane contact region. These bonds, represented in cyan and green, are all close to their neutral state (i.e., their length \( l \) is close to the equilibrium value \( \lambda \)). Over this contact region, the membrane is wrapped around the particle and follows its spherical shape. Internalization progresses driven by bond formation within a small leading region, where the bonds are under strong tension (bonds represented in red). The final phase, represented by the snapshot 3e, corresponds to internalization approaching the final equilibrium state, where further wrapping is no longer favorable.

Fig 4 illustrates the effect of the nanoparticle size on the internalization process. Fig 4a shows how the time required for internalization depends on the nanoparticle radius. We define the time for internalization as the time required for reaching \( d/(2a) = 1 \). For typical parameter values (the best estimates in Table 1), our model predicts an optimal radius at which internalization is fastest of \( a \approx 50 \) nm, which is consistent with experimental observations [4]. The time required for internalization remains comparable for a range of radii between 30 and 90 nm. Outside this range, the time required to reach \( d/(2a) = 1 \) becomes asymptotically longer, and at \( a = 20 \) nm or \( a = 100 \) nm the stage \( d/(2a) = 1 \) is no longer reached, indicating that internalization efficiency decreases significantly for very small and very large particles. Fig 4b shows the maximum value of \( d/(2a) \) that is reached as a function of the particle size. When the particle radius decreases below \( a = 20 \) nm, \( d/(2a) \) rapidly approaches zero, indicating that no particle wrapping occurs. This suggests that receptor-mediated internalization becomes impossible for small radii. In contrast, the maximum amount of wrapping attained for particles larger than \( a = 100 \) nm decreases slowly with increasing particle size, indicating a larger degree of membrane wrapping around the particle and thus suggesting that receptor-mediated internalization may still be possible up to a micrometric particle size (i.e., \( a \approx 500 \) nm), as observed in experiments [3].

Fig 5 shows the dependence of the particle internalization time (defined as the time to reach \( d/(2a) = 1 \)) on the time required for bond formation, \( t_b \). In the figure both axes are expressed in seconds, but they can equivalently be regarded as normalized by the viscoelastic time scale, \( \mu/E \), which we have adopted as our time unit, and whose typical value is indeed of the order of 1 s. The figure shows that the time required for particle internalization increases proportionally to both the bond formation time and to the viscoelastic time. Interestingly, whereas both the
Fig 3. Kinetics of nanoparticle internalization. a. Evolution of the particle internalization, measured by the depth-to-diameter ratio, $d/(2a)$ (see inset), as a function of the elapsed non-dimensional time (one non-dimensional time unit corresponds typically to one second). The computation is performed with $a = 50$ nm, $K_0 = 0.01 \mu m^{-2}$ and otherwise with the best estimate of the parameter values listed in Table 1. The horizontal dashed line corresponds to the value $d/(2a) = 1$, which is the threshold where we define a particle as internalized. Four snapshots of the internalization process, indicated on plot a by circle markers denoted by ’b’ to ’e’ and corresponding to times $t = 5 \cdot 10^{-3}$, $t = 10^{-2}$, $t = 2.0$, and $t = 10.0$ s, respectively, are presented in the plots b to e. Each snapshot shows the instantaneous membrane shape (full line), the initial membrane shape (dashed line), the particle (gray shaded circle), and the bonds formed between the particle and the membrane (short color lines). The density of color lines is proportional to the bond density. Dark blue, cyan, green, and red lines correspond respectively to bonds that are strongly compressed ($l < 0.9\lambda$), slightly compressed ($0.9\lambda \leq l < \lambda$), slightly stretched ($\lambda < l \leq 1.1\lambda$), or strongly stretched ($l > 1.1\lambda$).
viscoelastic time and the bond formation time are at most of the order of a second, the particle internalization time is much longer, of the order of a few tens of seconds and up to minutes, as observed in experiments [3]. The reason for the emergence of this longer internalization time scale is that, whereas bond formation induces deformations at the scale of the bond length, internalization requires deformations at the length scale of the particle. Thus, the ratio of time scales of particle internalization to bond formation is proportional to the ratio of length scales of the particle to the bond.

Fig 6 shows curves of maximum particle internalization depth (maximum $d/(2a)$) versus particle radius ($a$) for different values of several bond parameters. Fig 6a shows the effect of changes in the bond spring constant, $\kappa$. Interestingly, there appears to be an optimal value of $\kappa$, of about $5 \cdot 10^{-3}$ N/m, for which particle internalization is most efficient. The existence of this optimum arises from two opposing effects of bond rigidity, $\kappa$, on particle wrapping. On the one hand, reducing bond rigidity reduces the force provided by each bond to deform the cell
and wrap the nanoparticle. On the other hand, increasing bond rigidity makes the bond dissociation constant increase very rapidly with bond length (due to Eq 11), thus impairing bond formation. Fig 6b and 6c respectively show the effect of changes in the bond reaction constant, $K_0 / K_f$, and in the ligand density, $\xi_l$, which is in all cases assumed equal to the receptor density, $\xi_r$. As expected, particle internalization becomes more efficient with decreasing $K_0 / K_f$ and with increasing $\xi_l$, since in both cases the available energy gain from bond formation increases.

Fig 7 shows curves of maximum particle internalization depth (maximum $d/(2a)$) versus particle radius ($a$) for different values of the cell membrane properties (its bending rigidity $B$ and its tension $T_0$) and of the cytoplasm properties (its Young’s modulus $E$). As expected, internalization is favored by a smaller resistance to cell deformation, thus by smaller $B$, $T_0$, or $E$. However, changes in these different parameters affect internalization differently. Within the range of physiological values, the effect of changes in $T_0$ on internalization are much smaller than those of changes in $B$ or in $E$. Changes in $B$ strongly affect the internalization of smaller particles, since membrane bending stresses, which scale as $B/a^3$, are the dominant resistance to small particle internalization. Large particle internalization, on the other hand, is virtually independent of the value of $B$ and strongly controlled by the value of $E$. Fig 7 thus shows that cytoplasmic rigidity becomes the dominant resisting mechanism at large $a$, as one would expect from comparing the dependence on $a$ of the nondimensional coefficients $C_B$ and $^\wedge T_0$ in Table 2.

Fig 8a shows the dependence of the time required for the internalization of a particle of radius $a = 50$ nm on the magnitude of a normal, nondimensional force, $\tilde{F}_v = F_a \xi_l/(a^2k_BT)$, exerted onto the particle (see figure inset). Consistent with experimental observations using magnetic nanoparticles subjected to a magnetic field [30], we predict the internalization time to decrease with the applied force. The figure shows that the nondimensional force required to significantly decrease the time for particle internalization is of the order of one nondimensional unit, which for typical parameter values corresponds in dimensional terms to a stress applied onto the particle of about 2 kPa (or equivalently a total force of about 20 pN exerted onto the
Fig 6. Effect of bond properties on the maximum internalization depth as a function of the particle radius, a. Different marker shapes and colors correspond to different values of the bond spring constant $\kappa$ (in plot a), of the bond reaction constant $K_0^d/K_0$ (in plot b), or of the ligand density $\xi_l$ (in plot c). In all plots, the black curve corresponds to the best estimate of the varied parameter ($\kappa = 10^{-3}$ N/m, $K_0^d/K_0 = 5 \cdot 10^{-2}$ $\mu$m$^{-2}$, and $\xi_l = 10^2$ $\mu$m$^{-2}$, respectively). In each plot, all parameter values not indicated are taken equal to their best estimates (Table 1), except for $\xi_l$ = $\xi_f$ in plot c. The gray horizontal line indicates the internalization threshold, $d/(2a) = 1$.

doi:10.1371/journal.pone.0122097.g006
Fig 7. Effect of cell mechanical properties on the maximum internalization depth as a function of the particle radius, a. Different marker shapes and colors correspond to different values of the membrane bending modulus \( B \) (in plot a), of the initial membrane tension \( T_0 \) (in plot b), or of the cell’s Young’s modulus \( E \) (in plot c). In each plot, the black curve corresponds to the best estimate of the varied parameter (\( B = 10^{-19} \) J, \( T_0 = 30 \) pN/\( \mu \)m, and \( E = 10^3 \) Pa, respectively). On each plot, all parameter values not indicated are taken equal to their best estimates (Table 1). The gray horizontal line indicates the internalization threshold, \( d/(2a) = 1 \).

doi:10.1371/journal.pone.0122097.g007
Fig 8. Effect of a vertical force on internalization. a. Time required for particle internalization (defined as the time required to reach \( \frac{d}{2a} = 1 \)) as a function of a normalized vertical compressive force, \(-\hat{F}_v = -F_v r_0/(\kappa a^2 k_BT)\), for \( a = 50 \text{ nm} \) and all other parameter values equal to their best estimates (Table 1). The inset shows a snapshot of the membrane deformation and particle position for the case \( \hat{F}_v = -1 \). The snapshot corresponds to the time when \( \frac{d}{a} = 1 \). b. Time required for particle internalization (defined as the time required to reach \( \frac{d}{2a} = 1 \)) as a function of the particle radius \( a \) in the absence of force (black circles) and in the presence of a normalized force \( \hat{F}_v = -0.5 \) (blue squares).

doi:10.1371/journal.pone.0122097.g008
50 nm particle). Fig 8b illustrates the effect on the internalization time of applying a nondimensional force \( F_v = -0.5 \) onto particles of different radii \( a \). Dimensionally, \( F_v = -0.5 \) corresponds to a stress of about 1 kPa applied onto the particle top surface. The figure shows that a compressive stress more significantly improves the internalization of larger particles. As discussed above, the internalization of small particles is mainly limited by the membrane bending rigidity, whereas the internalization of large particles is mainly limited by the rigidity of the cell’s cytoplasm. Because the main effect of a compressive force is to deform the cell’s cytoplasm, external compressive stresses particularly facilitate the internalization of large particles. The figure thus suggests that external forces acting on nanoparticles—induced for example by electric charges, magnetic fields, or the flow pressure—can favor the internalization of larger particles.

We have also investigated the effect of a shear flow on particle internalization. To this end, we have developed a two-dimensional version of the model, where the particle shape becomes an infinitely long cylinder. The two-dimensional model allows us to consider the fore-aft asymmetric case where a horizontal force and torque act on the nanoparticle. We found that the flow-induced drag only affects the early stage of particle adhesion to the membrane, when the particle is most exposed to the flow. A sufficiently strong flow drag will rapidly detach the particle and no internalization will take place. However, if the initial adhesion is sufficiently strong to withstand the flow drag, the flow does not affect the particle internalization dynamics, which are then virtually identical to the no-flow case. The effect of the flow on the results is thus bimodal and abrupt, and it depends critically on the initial bond distribution. This result is consistent with numerous studies that have described the important role of flow in particle adhesion [14, 31–33]. However, recent studies have shown that flow also has a direct effect on nanoparticle internalization [9, 34]. Our two-dimensional computations of internalization under shear flow suggest that the dominant process by which flow modulates nanoparticle internalization is not a purely mechanical one, as included in our model, but rather an active biological phenomenon involving cellular mechanotransduction, by which the flow affects cytoskeletal organization and dynamics [9] and biochemical pathways involved in particle endocytosis [34].

Discussion

In this section we propose simple scaling arguments to interpret our numerical results and thus to gain insight into the physics underlying the internalization process. Our goal in this section is to understand the time scales governing the kinetics of internalization, as well as the dependence of the nanoparticle internalization depth on the membrane rigidity and cytoplasmic Young’s modulus.

As shown in Fig 3, the internalization dynamics can be described as consisting of three distinct stages: development of a firm adhesion, wrapping, and stabilization at an equilibrium wrapped state. What sets the duration of these different stages? Let us first consider the time needed for completing the firm adhesion stage. The firm adhesion stage is characterized by the development of a sufficiently large bond density to deform the membrane and initiate the wrapping process. Since the initial bond density \( \xi_0 \) is small compared to the total ligand and receptor densities, \( \xi_l \) and \( \xi_r \), we can characterize the short-time dynamics of bond formation by the following simplified version of Eq 10:

\[
\frac{\partial \xi_b}{\partial t} \approx K \xi_l \xi_r.
\]

We deduce from Eq 13 that the time needed to complete the firm adhesion stage and start
deforming the membrane scales as
\[ t_{\text{firm}} \sim \frac{\xi_{\text{b, firm}}}{K_r^{1/2} \varepsilon}, \]  \hspace{1cm} (14)

where \( \xi_{\text{b, firm}} \) is the bond density that needs to be attained in order to start deforming the membrane. Deformation of the membrane becomes possible when the energy stored by the bond springs reaches the energy needed to bend the membrane, i.e.:
\[ \frac{1}{2} \kappa (l_{\text{max}} - \lambda)^2 \xi_{\text{b, firm}} \sim \frac{1}{2} B \frac{1}{a^2}, \]  \hspace{1cm} (15)

from which we deduce that \( \xi_{\text{b, firm}} \sim B/(\kappa (l_{\text{max}} - \lambda)^2 a^2) \). Here, \( l_{\text{max}} \) is the maximum distance between the particle and the membrane at which bonds can form. Net bond formation occurs if the bond dissociation rate remains smaller than the bond formation rate, i.e., if
\[ \frac{K_r}{K} = \frac{K_r}{(K_f \xi)} \leq 1. \]  

The value of \( l_{\text{max}} \) is thus given by \( K_r = 1 \). Considering the dependence of \( K_d \) on \( l \) given by Eq 11, we deduce
\[ (l_{\text{max}} - \lambda) \sim \frac{k_B T}{k_B \xi} \ln \left( \frac{K_f^{1/2}}{K_D^{1/2}} \right). \]  \hspace{1cm} (16)

For a bond spring constant \( \kappa = 0.01 \text{ N/m}, \) Eq 16 yields a maximum bond deformation of the order of \( (l_{\text{max}} - \lambda) \approx 10 \text{ nm} \), which is consistent with our numerical results. Combining Eqs 14 and 15, with \( (l_{\text{max}} - \lambda) \) given by Eq 16, we conclude that the duration of the firm adhesion stage scales as
\[ t_{\text{firm}} \sim \frac{B}{K_r^{1/2} \varepsilon a^2 \kappa (l_{\text{max}} - \lambda)^2}. \]  \hspace{1cm} (17)

For typical parameter values, Eqs 15 and 17 yield a typical bond density to start particle wrapping of \( \xi_{\text{b, firm}} \approx 0.05 \xi_0 \), which is attained at a time \( t_{\text{firm}} \approx 10^{-3} \text{ s} \), consistent with the results of the numerical model (Fig 3).

To interpret the typical duration of the wrapping phase, we propose a simplified view of the wrapping process, schematized in Fig 9. We consider a simplified geometry where the membrane is wrapped around the particle over the contact zone, adopting a spherical shape of radius \( a + \lambda \), whereas outside the contact zone the membrane remains flat. Bonds follow a radial orientation. We note that this simplified view is limited to representing wrapping up to the

\[ \text{Fig 9. Schematics illustrating a simplified view of the geometry of early particle wrapping.} \]

doi:10.1371/journal.pone.0122097.g009
particle’s equator, since beyond $\alpha = \pi/2$ formation of new bonds becomes geometrically impossible (see Fig 9). At a given instant, the internalization depth $d$ is related to the wrapping angle $\alpha$ by $d = a(1 - \cos \alpha)$. New bonds are being formed over an annulus of width $p$, where bond length remains smaller than $l_{\text{max}}$ defined above. Geometric considerations yield $p = (l_{\text{max}} - \lambda) / \tan \alpha$. Since bond formation requires a time of the order of $t_b \sim K_i \xi_b$, we can write the following equation of the simplified internalization dynamics:

$$\frac{dz}{dt} \approx \frac{p}{t_b} \frac{(a + \lambda)}{(a + \lambda) l_{\text{max}} - \lambda} \tan \alpha. \quad (18)$$

This differential equation yields $d/(2a) \sim (\cos \alpha)/2 \sim \exp[(-l - \lambda)(a + \lambda)/t_b t)],/2$, which is an exponentially asymptotic evolution equation towards an equilibrium value, here of $d/2a = 0.5$, and conceptually similar to the asymptotic evolution described by our computational model and shown in Fig 3. This simple reasoning suggests that the time for wrapping scales as

$$t_{\text{wrapping}} \sim \frac{(a + \lambda)}{(l - \lambda)} t_b, \quad (19)$$

which, for $a = 50$ nm, yields $t_{\text{wrapping}} \sim 7t_b$. The full computational model predicts indeed a proportionality between the two time scales, as shown in Fig 5 above, but with a coefficient of proportionality of about 25 rather than 7, corresponding to the more accurate wrapping geometry considered by the computational model.

Next we discuss the dependence of the maximum internalization depth on the membrane and cytoplasm rigidity ($B$ and $E$), presented in Fig 7 above. As shown in Fig 7a, membrane bending rigidity sets a minimum size for nanoparticle internalization, and the minimum particle size that can be internalized decreases with decreasing $B$. Here we propose a scaling argument to interpret the observed dependence. For small particles ($a < 30$ nm), the main resisting force to internalization is due to membrane bending, whereas the driving force is provided by the bonds forming at the edge of the wrapping region. Momentum balance per unit of azimuthal length yields the scaling $B \theta/\theta ds \sim F_b(\xi_p) w$, where $F_b$ is the force per bond, $p$ is the width of the annulus of stretched bonds at the edge of the wrapping zone (see Fig 9), and $w$ is the horizontal extent of the deformed region of the membrane (see Fig 10a). The rate of change of membrane inclination scales as $d\theta/ds \sim a/w \sim (d/a)/w$. For small particles, the two dominant terms defining the deformed shape of the membrane (see Fig 10b) are the membrane bending rigidity and the cytoplasm elasticity. Balance between these two yields a typical extent of membrane deformation of $w \sim (B/K)^{1/4} = (Ba/E)^{1/4}$. Combining these scalings leads to

$$\frac{d}{2a} \sim \sqrt{\frac{a}{BE}}. \quad (20)$$

Fig 11a represents the data of Fig 7a rescaled according to the dependence on $B$ suggested by Eq 20. In Fig 7a the curves for different $B$ collapsed at large $a$, since for large particles membrane bending rigidity becomes small compared to membrane tension and cytoplasmic elasticity, and thus its effect on internalization becomes negligible. In Fig 11a, the rescaled curves collapse at small $a$, suggesting that the scaling argument given by Eq 20 correctly captures the effect of $B$ on the internalization of small particles.

As shown in Fig 7, the internalization of large particles ($a > 100$ nm) is largely controlled by the cytoplasmic Young’s modulus, $E$. We can also interpret this dependence using a scaling argument. The elastic force per unit of azimuthal length required to vertically indent the cytoplasm scales as $(Kd) \cdot w$, where the first factor represents the elastic force per unit area and the
second factor is the horizontal extent of the deformed region. For large particles \((a > 100 \text{ nm})\), \(w \sim d\) (see Fig 10c). Balance between the elastic force and the bond force yields the scaling \(Kd^2 \sim F_b \xi p\). Since \(K = E/a\), we deduce a simple scaling law for the dependence of \(d\) on \(a\) and \(E\), valid for large particle radius:

\[
\frac{d}{2a} \sim \frac{1}{\sqrt{Ea}}.
\] (21)

Fig 11b represents in log-log scale the data of Fig 7c rescaled according to Eq 21. This rescaling makes the different curves nearly collapse for large \(a\), suggesting that Eq 21 correctly captures the effect of \(E\) on the internalization of large particles. Only the scaled curve corresponding to \(E = 500 \text{ Pa}\) is far from collapsing with the others; this is attributed to the fact that, for this smaller value of \(E\), the cell’s elastic rigidity is no longer the dominant resisting mechanism to nanoparticle internalization. Fig 11b also shows that, for large particles, the maximum internalization depth \(d/(2a)\) seems to decrease with \(1/\sqrt{a}\), consistent with Eq 21.

**Conclusion**

Our model results show the existence of an optimal radius for receptor-mediated nanoparticle internalization of the order of 50 nm. Below this optimal radius, receptor-mediated internalization becomes rapidly impaired by membrane bending rigidity. Above the optimal radius, cytoplasmic rigidity makes internalization less efficient, but the effect of increasing the radius above the optimum is more gradual, as the maximum depth to which the particle can be internalized decreases only with the inverse of the square root of the particle size. We have also shown that bond characteristics play an important role in internalization. Notably, we have identified an optimum value of the bond elastic constant, of the order of 5 nN/\(\mu\text{m}\), for which internalization is most efficient.
Whereas experimental studies have evidenced an effect of flow on nanoparticle internalization [9, 34], our model suggests that the mechanical effect of a shear force applied on a nanoparticle is only important in determining nanoparticle adhesion to the cell membrane. Once firm adhesion is attained, our model predicts that internalization dynamics become independent of the direct mechanical action of the shear. This result suggests that shear affects nanoparticle internalization by modulating the active dynamics of the cellular cytoskeleton, rather than by a direct mechanical effect on the nanoparticle. In contrast, our model predicts that normal compressive stresses applied to nanoparticles can significantly improve their internalization, especially of those particles larger than about 100 nm.

A limitation of the model presented here is that it disregards active forces generated by the cellular cytoskeleton during nanoparticle internalization. Indeed, experiments show that ICAM-1-mediated internalization is dependent on the formation of actin stress fibers, a phenomenon that occurs over a time scale of about 10 minutes [3]. In contrast, the internalization mechanism described here is completed over a much shorter time scale (of a minute or less). Our model thus only describes the initial step of receptor-mediated nanoparticle internalization.
internalization, whereas later stages of internalization are controlled by cytoskeletal dynamics. Thus, our model can be directly validated against experiments of receptor-mediated nanoparticle uptake into artificial systems that lack a dynamic cytoskeleton, such as vesicles [35, 36] or polymersomes [37]. As our model accounts for the main internalization mechanisms with the exception of cytoskeletal dynamics, comparison between our model’s predictions and experiments of nanoparticle cell uptake can be used to isolate the contribution of cytoskeletal dynamics to the internalization process. Based on such additional experimental knowledge, cytoskeletal dynamics can be included in the model by making the cytoplasmic rheology or the bond properties depend on the internalization stage, or by adding an additional, time-dependent force acting on the membrane, to account for the active forces generated by the cytoskeleton.

Acknowledgments
D.G.-R. was funded by the Fondation pour la Recherche Médicale. This work was supported in part by a permanent endowment in cardiovascular cellular engineering from the AXA Research Fund.

Author Contributions
Conceived and designed the experiments: DGR AIB. Performed the experiments: DGR. Analyzed the data: DGR AIB. Wrote the paper: DGR AIB.

References
1. Mendis S, Puska P, Norrving B. Global atlas on cardiovascular disease prevention and control. World Health Organization; 2011.
2. Chithrani BD, Ghazani AA, Chan WCW. Determining the size and shape dependence of gold nanoparticle uptake into mammalian cells. Nano Letters. 2006; 6:662–666. PMID: 16608261
3. Muro S, Garnacho C, Champion JA, Leferovich J, Gajewski C, Schuchman EH, et al. Control of endothelial targeting and intracellular delivery of therapeutic enzymes by modulating the size and shape of ICAM-1-targeted carriers. Molecular Therapy. 2008; 16:1450–1458. doi: 10.1038/mt.2008.127 PMID: 18560419
4. Albanese A, Tang PS, Chan WCW. The effect of nanoparticle size, shape, and surface chemistry on biological systems. Annual Review of Biomedical Engineering. 2012; 14:1–16. doi: 10.1146/annurev-bioeng-071811-150124 PMID: 22524388
5. Muzykantov VR. Targeted drug delivery to endothelial adhesion molecules. ISRN Vascular Medicine. 2013; 2013:916254. doi: 10.1155/2013/916254
6. Calderon AJ, Bhowmick T, Leferovich J, Burman B, Pichette B, Muzykantov V, et al. Optimizing endothelial targeting by modulating the antibody density and particle concentration of anti-ICAM coated carriers. Journal of Controlled Release. 2011; 150:37–44. doi: 10.1016/j.jconrel.2010.10.025 PMID: 21047540
7. Muro S, Wierwoldt R, Thomas A, Koniaris L, Albelda SM, Muzykantov VR, et al. A novel endocytic pathway induced by clustering endothelial ICAM-1 or PECAM-1. Journal of Cell Science. 2003; 116:1599–1609. doi: 10.1242/jcs.00367 PMID: 12640043
8. Muro S, Dzubla T, Qiu W, Leferovich J, Cui X, Berk E, et al. Endothelial targeting of high-affinity multivalent polymer nanocarriers directed to intercellular adhesion molecule 1. The Journal of Pharmacology and Experimental Therapeutics. 2006; 317:1161–1169. doi: 10.1124/jpet.105.098970 PMID: 16505161
9. Bhowmick T, Berk E, Cui X, Muzykantov VR, Muro S. Effect of flow on endothelial endocytosis of nanocarriers targeted to ICAM-1. Journal of Controlled Release. 2012; 157:485–492. doi: 10.1016/j.jconrel.2011.09.067 PMID: 21951807
10. Serrano D, Bhowmick T, Chadhra R, Garnacho C, Muro S. Intercellular adhesion molecule 1 engagement modulates sphingomyelinase and ceramide, supporting uptake of drug carriers by the vascular endothelium. Arteriosclerosis, Thrombosis, and Vascular Biology. 2012; 32:1178–1185. doi: 10.1161/ATVBAHA.111.244186 PMID: 22328778
11. Gao H, Shi W, Freund LB. Mechanics of receptor-mediated endocytosis. Proceedings of the National Academy of Sciences of the USA. 2005; 102:9469–9474. doi: 10.1073/pnas.0503879102 PMID: 15972807

12. Decuzzi P, Ferrari M. The receptor-mediated endocytosis of nonspherical particles. Biophysical Journal. 2008; 94:3790–3797. doi: 10.1529/biophysj.107.120238 PMID: 18234813

13. Li X, Xing D. A simple method to evaluate the optimal size of nanoparticles for endocytosis based on kinetic diffusion of receptors. Applied Physical Letter. 2010; 97:153704. doi: 10.1063/1.3502489

14. Calderon AJ, Muzykantov V, Muro S, Eckmann DM. Flow dynamics, binding and detachment of spherical carriers targeted to ICAM-1 on endothelial cells. Biotherapy. 2009; 46:323–341. doi: 10.3233/BIR-2009-0544 PMID: 19721193

15. Bahrami AH, Raatz M, Aguado-Canalejo J, Michel R, Curtis E, Hall CK, et al. Wrapping of nanoparticles by membranes. Advances in Colloid and Interface Science. 2014; 208:214–224. doi: 10.1016/j.cis.2014.02.012 PMID: 24703299

16. Dasgupta S, Auth T, Gompper G. Wrapping of ellipsoidal nano-particles by fluid membranes. Soft Matter. 2013; 9:5473–5482. doi: 10.1039/c3sm50351h

17. Dasgupta S, Auth T, Gompper G. Shape and orientation matter for the cellular uptake of nonspherical particles. Nano Letters. 2014; 14:687–693. doi: 10.1021/nl403949h PMID: 24383757

18. Yi X, Shi X, Gao H. Cellular uptake of elastic nanoparticles. Physical Review Letters. 2011; 107:098101. doi: 10.1103/PhysRevLett.107.098101 PMID: 21929271

19. Chaudhuri A, Battaglia G, Golestanian R. The effect of interactions on the cellular uptake of nanoparticles. Physical Biology. 2011; 8:046002. doi: 10.1088/1478-3975/8/4/046002 PMID: 21508440

20. Raatz M, Lipowsky R, Weikl TR. Cooperative wrapping of nanoparticles by membrane tubes. Soft Matter. 2014; 10:3570–3577. doi: 10.1039/c3sm52496a PMID: 24656648

21. Mirgian S, Muthukumar M. Kinetics of particle wrapping by a vesicle. The Journal of Chemical Physics. 2013; 139:044908. doi: 10.1063/1.4813921 PMID: 23902020

22. Yue T, Zhang X. Molecular understanding of receptor-mediated membrane responses to ligand-coated nanoparticles. Soft Matter. 2011; 7:9104–9112. doi: 10.1039/c1sm05398a

23. Dembo M, Torney DC, Saxman K, Hammer D. The reaction-limited kinetics of membrane-to-surface adhesion and detachment. Proc R Soc Lond B. 1988; 234:55–83. doi: 10.1098/rspb.1988.0038 PMID: 2901109

24. Bell GI, Dembo M, Bongrand P. Cell adhesion. Competition between nonspecific repulsion and specific bonding. Biophys J. 1984; 45:1051–1064. doi: 10.1016/S0006-3495(84)84252-6 PMID: 6743742

25. Bell GI. Models for the specific adhesion of cells to cells. Science. 1978; 200:618–627. doi: 10.1126/science.347575 PMID: 347575

26. Seifert U. Rupture of multiple parallel molecular bonds under dynamic loading. Physical Review Letters. 2000; 84:2750–2753. doi: 10.1103/PhysRevLett.84.2750 PMID: 11017316

27. Casasnovas JM, Springer TA. Kinetics and thermodynamics of virus binding to receptor. The Journal of Biological Chemistry. 1995; 270:13216–13224. doi: 10.1074/jbc.270.22.13216 PMID: 7768920

28. Zhou X, Perez F, Han K, Jurivich DA. Clonal senescence alters endothelial ICAM-1 function. Mechanisms of Ageing and Development. 2006; 127:779–785. doi: 10.1016/j.mad.2006.07.003 PMID: 16930678

29. Barreiro O, Zamai M, Yañez-Mó M, Tejera E, López-Romero P, Monk PN, et al. Endothelial adhesion receptors are recruited to adherent leukocytes by inclusion in preformed tetraspanin nanoplatforms. Journal of Cell Biology. 2008; 183:527–542. doi: 10.1083/jcb.200805076 PMID: 18955551

30. MacDonald C, Barbee K, Polyak B. Force dependent internalization of magnetic nanoparticles results in highly loaded endothelial cells for use as potential therapy delivery vectors. Pharmaceutical Research. 2012; 29:1270–1281. doi: 10.1007/s11095-011-0663-7 PMID: 22234617

31. Chang KC, Tees DFJ, Hammer DA. The state diagram for cell adhesion under flow: Leukocyte rolling and firm adhesion. Proceedings of the National Academy of Sciences of the USA. 2000; 97:11262–11267. doi: 10.1073/pnas.200240897 PMID: 11005837

32. Pierres A, Touchard D, Benoliel AM, Bongrand P. Dissecting streptavidin-biotin interaction with a laminar flow chamber. Biophysical Journal. 2002; 82:3214–3223. doi: 10.1016/S0006-3495(02)75664-6 PMID: 12023246

33. Verdier C, Couzon C, Duperray A, Singh P. Modeling cell interactions under flow. Journal of Mathematical Biology. 2009; 58:235–259. doi: 10.1007/s00285-008-0164-4 PMID: 18292997

34. Han J, Zern BJ, Shuvvaev VV, Davies PF, Muro S, Muzykantov V. Acute and chronic shear stress differentially regulate endothelial internalization of nanocarriers targeted to Platelet-Endothelial Cell Adhesion Molecule-1. ACS Nano. 2012; 6:8824–8836. doi: 10.1021/nn302687n PMID: 22957767
35. Fery A, Moya S, Puch PH, Brochard-Wyart F, Mohwald H. Interaction of polyelectrolyte coated beads with phospholipid vesicles. Comptes Rendus Physique. 2003; 4:259–264. doi: 10.1016/S1631-0705 (03)00030-6

36. Le Bihan O, Bonnafous P, Marak L, Bickel T, Trépout S, Mornet S, et al. Cryo-electron tomography of nanoparticle transmigration into liposome. Journal of Structural Biology. 2009; 168:419–425. doi: 10.1016/j.jsb.2009.07.006 PMID: 19596070

37. Jaskiewicz K, Larsen A, Schaeffel D, Koynov K, Lieberwirth I, Fytas G, et al. Incorporation of nanoparticles into polymersomes: size and concentration effects. ACS Nano. 2012; 6:7254–7262. doi: 10.1021/nn302367m PMID: 2279259

38. Agrawal NJ, Radhakrishnan R. The role of glycocalyx in nanocarrier-cell adhesion investigated using a thermodynamic model and Monte Carlo simulations. Journal of Physical Chemistry C: Nanomaterials and interfaces. 2007; 111:15848–15856. doi: 10.1021/jp074514x PMID: 19081797

39. Liu J, Weller GER, Zern B, Ayyaswamy PS, Eckmann DM, Muzykantov VR, et al. Computational model for nanocarrier binding to endothelium validated using in vivo, in vitro, and atomic force microscopy experiments. Proceedings of the National Academy of Sciences of the USA. 2010; 107:16530–16535. doi: 10.1073/pnas.1006611107 PMID: 20823296

40. Schiavo VL, Robert P, Limozin L, Bongrand P. Quantitative modeling assesses the contribution of bond strengthening, re-binding and force sharing to the avidity of biomolecule interactions. PLOS One. 2012; 7:e44070. doi: 10.1371/journal.pone.0044070 PMID: 23024747

41. Evans E, Kinoshita K, Simon S, Leung A. Long-lived, high-strength states of ICAM-1 bonds to \( \beta_2 \) integrin, I: Lifetimes of bonds to recombinant \( \alpha_5 \beta_2 \) under force. Biophysical Journal. 2010; 98:1458–1466. doi: 10.1016/j.bpj.2009.09.067 PMID: 20409464

42. Mogilner A, Oster G. Cell motility driven by actin polymerization. Biophysical Journal. 1996; 71:3030–3045. doi: 10.1016/S0006-3495(96)79496-1 PMID: 8968574

43. Raucher D, Sheetz MP. Cell spreading and lamellipodial extension rate is regulated by membrane tension. Journal of Cell Biology. 2000; 148:127–136. doi: 10.1083/jcb.148.1.127 PMID: 10629223

44. Boulant S, Kural C, Zeeh JC, Ubelmann F, Kirchhausen T. Actin dynamics counteract membrane tension during clathrin-mediated endocytosis. Nature Cell Biology. 2011; 13:1124–1131. doi: 10.1038/ncb2307 PMID: 21841790

45. Sato M, Nagayama K, Kataoka N, Sasaki M, Hane K. Local mechanical properties measured by atomic force microscopy for cultured bovine endothelial cells exposed to shear stress. Journal of Biomechanics. 2000; 33:127–135. doi: 10.1016/S0021-9290(99)00178-5 PMID: 10609525

46. Mathur AB, Collinsworth AM, Reichert WM, Kraus WE, Truskey GA. Endothelial, cardiac muscle and skeletal muscle exhibit different viscous and elastic properties as determined by atomic force microscopy. Journal of Biomechanics. 2001; 34:1545–1553. doi: 10.1016/S0021-9290(01)00149-X PMID: 11716856

47. Sato M, Theret DP, Wheeler LT, Ohshima N, Nerem RM. Application of the micropipette technique to the measurement of cultured porcine aortic endothelial cell viscoelastic properties. Journal of Biomechanical Engineering. 1990; 112:263–268. doi: 10.1115/1.2891183 PMID: 2214707

48. Sato M, Oshima N, Nerem RM. Viscoelastic properties of cultured porcine aortic endothelial cells exposed to shear stress. Journal of Biomechanics. 1996; 29:461–467. doi: 10.1016/0021-9290(95)00069-0 PMID: 8964775

49. Koay EJ, Shieh AC, Athanasiou KA. Creep indentation of single cells. Journal of Biomechanical Engineering. 2003; 125:334–341. doi: 10.1115/1.1572517 PMID: 12929237