

Lateral mobility of proteins in liquid membranes revisited

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The biological function of transmembrane proteins is closely related to their insertion, which has most often been studied through their lateral mobility. For >30 years, it has been thought that hardly any information on the size of the diffusing object can be extracted from such experiments. Indeed, the hydrodynamic model developed by Saffman and Delbrück predicts a weak, logarithmic dependence of the diffusion coefficient D with the radius R of the protein. Despite widespread use, its validity has never been thoroughly investigated. To check this model, we measured the diffusion coefficients of various peptides and transmembrane proteins, incorporated into giant unilamellar vesicles of 1-stearoyl-2-oleoyl-*sn*-glycero-3-phosphocholine (SOPC) or in model bilayers of tunable thickness. We show in this work that, for several integral proteins spanning a large range of sizes, the diffusion coefficient is strongly linked to the protein dimensions. A heuristic model results in a Stokes-like expression for D , ($D \propto 1/R$), which fits literature data as well as ours. Diffusion measurement is then a fast and fruitful method; it allows determining the oligomerization degree of proteins or studying lipid–protein and protein–protein interactions within bilayers.

bilayers | transmembrane proteins | diffusion | peptides | sponge phase

In the hydrodynamic model of Saffman and Delbrück (1), transmembrane peptides and proteins are described as diffusing in a perfectly continuous medium, ignoring the finite size of the lipids. This model predicts that the diffusion coefficient D of a simple cylinder embedded in a thin sheet of fluid matching its height (Fig. 1) is given by

$$D_{\text{Saffman}} = \frac{k_B T}{4\pi\mu_m h} \left(\ln\left(\frac{\mu_m h}{\mu_w R}\right) - 0.5772 \right). \quad [1]$$

In this expression, the adjustable parameters are by order of importance: the thickness h and viscosity μ_m of the liquid membrane, the radius R of the diffusing cylinder, and the viscosity of the surrounding aqueous phase μ_w . This result follows from solving the flow field in the membrane and in the surrounding fluid, assuming no-slip boundary conditions at the surface of the cylinder, which is considered as large compared with the bilayer components (i.e., $R > h$).

Numerous biological studies, both in model systems (2–4) and living cells (5, 6), refer to this continuum approach (7). Because D depends only weakly on R , the characterization of protein or rafts radii is delicate (8); for example, increasing the radius from 10 to 100 Å changes the mobility by a mere 30% [for $h = 30$ Å and $\mu_m = 10$ poise (P ; 1 P = 0.1 Pa·s)].

To check the applicability of the Saffman–Delbrück formula (Eq. 1), we have used fringe pattern photobleaching under the microscope (9) to measure precisely the self-diffusion of transmembrane peptides and proteins of well characterized dimensions.

Results and Discussion

The weight of the bilayer thickness, h , has never been investigated. Rather than using lipids of various lengths, we opted for a unique system where the bilayer thickness can be continuously tuned, leaving the bilayer viscosity constant.

We use a phase of model bilayers made of nonionic surfactants (penta-monododecylether; $C_{12}E_5$) diluted by water. These bilayers can be swollen at will by a hydrophobic solvent, which increases h from 16 Å [unswollen bilayer (10)] up to 40 Å, as checked by small-angle x-ray scattering experiments. To simplify the analysis, we wish to keep the viscosity of the membrane unchanged upon swelling. To do so, we choose dodecane, which matches the hydrophobic tails of the surfactant. We tested the properties of each monolayer by labeled surfactants (C_{12} -FITC) and those of the whole bilayer by longer probes: labeled lipids [1-stearoyl-2-oleoyl-*sn*-glycero-3-phosphocholine (SOPC)–4-chloro-7-nitrobenz-2-oxa-1,3-diazole (NBD)]. As detailed in *Materials and Methods*, the fact that the diffusion coefficients of these two different molecules remained constant upon swelling indicates that the viscosity of this “model membrane” was mainly unaltered, with the mid-plane dodecane displaying the same apparent viscosity as the $C_{12}E_5$ monolayers.

We have synthesized model peptides of increasing hydrophobic length d_π : L_{12} , L_{18} , and L_{24} (Fig. 1), and radius of 5.5 Å. Poly-leucine sequences, capped at both ends by polar heads, create stable and well defined α -helices as models of transmembrane “cylinders.”

Upon swelling, the diffusion coefficient varies in a similar manner for all of the peptides studied (Fig. 1). The value of D has a maximum D_{max} when the hydrophobic membrane thickness, h , matches the peptide hydrophobic length d_π , corresponding to Saffman's conditions. The variation of D_{max} with h is best fitted with a simple $1/h$ law (solid line), whereas the dotted line gives the predicted D_{Saffman} of Eq. 1.

To investigate the dependence of D on the radius R , a model transmembrane protein, bacteriorhodopsin (BR), was incorporated into giant unilamellar vesicles (GUVs) of SOPC (11). The peptide L_{18} , perfectly adapted to the membrane ($h = d_\pi = 28$ Å), calibrates the bilayer's properties.

BR, made of seven transmembrane α -helices [$R_{\text{BR}} = 18$ Å (12)], diffuses 3.8-times slower than the single helix L_{18} -peptide. ($D_{\text{BR}} = 0.08$ and $D_{L_{18}} = 0.31 \mu\text{m}^2/\text{s}$). Such a difference, according to the Saffman–Delbrück formula (Eq. 1), indicates that the proteins are aggregated in giant clusters of radius

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Abbreviations: BR, bacteriorhodopsin; $C_{12}E_5$, penta-monododecylether; GUV, giant unilamellar vesicle; P, poise; SOPC, 1-stearoyl-2-oleoyl-*sn*-glycero-3-phosphocholine.

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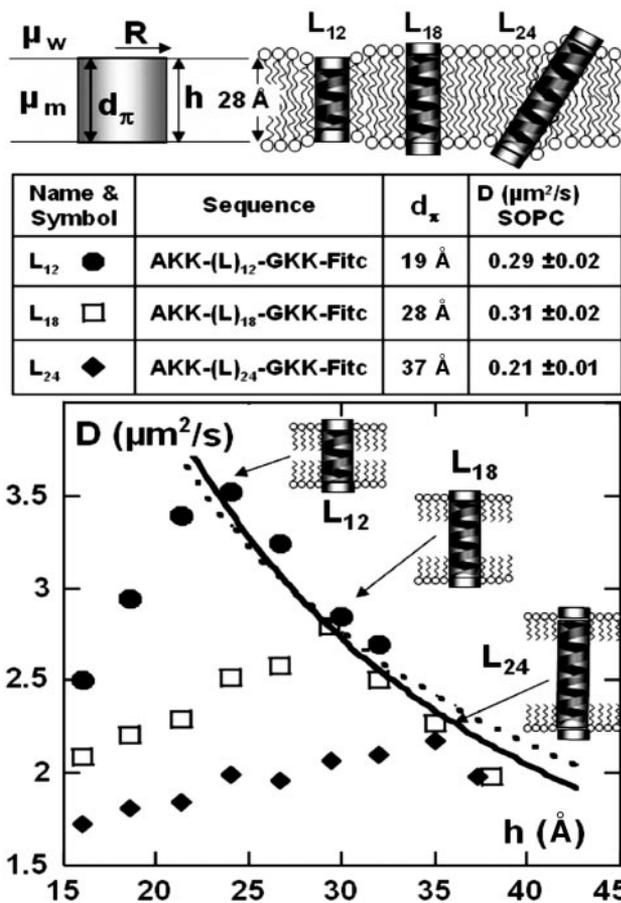


Fig. 1. Peptides used and their D variations versus bilayer thickness h . (Top) The parameters used in the Saffman–Delbrück model (Eq. 1), in the case of peptides diffusing in a giant unilamellar vesicle (GUV) made of SOPC. (Middle) A summary of the properties of the peptides used, their sequences, and their hydrophobic length d_x . The right-hand column gives the measured diffusion coefficient of these peptides in single GUVs. The diffusion coefficient was determined by using evanescent fluorescence recovery after pattern photobleaching technique. The data are typically averaged >200 vesicles. (Bottom) The variation of the diffusion due to the swelling of the $C_{12}E_5$ bilayer for the three analog peptides L_{12} (●), L_{18} (□), and L_{24} (◆). For each peptide, five sets of experiments allowed us to obtain average values with a reproducibility of >5% (the symbol size). The dotted line is the fit with the Saffman–Delbrück model (Eq. 1), using only one adjustable parameter, μ_m . The radius was taken as 5.5 Å and the viscosity of water as 0.01 P, leading to $\mu_m = 2.94$ P. The solid line represents a simple $1/h$ dependence. Note that the relative D variations are the same in $C_{12}E_5$ bilayers and in SOPC membranes: for $h = 28$ Å, L_{24} diffuses 30% slower than L_{18} ; L_{12} and L_{18} have similar mobilities.

$R_{\text{Saffman}} = 0.48 \mu\text{m}$. However, confocal microscopy shows no such large objects, but a homogeneous fluorescence distribution in agreement with previous electron-microscopy results (13, 14) that favor a monomeric state (in the dark state of the light-activated protein). Thus, the $D_{\text{BR}}/D_{L_{18}} \approx R_{L_{18}}/R_{\text{BR}}$.

This behavior, in contrast to the Saffman–Delbrück prediction, is supported by literature. In Fig. 2, we gather published

Considering the peptide L_{18} ($d_p = h = 28$ Å, $R = 5.5$ Å), the viscosity μ_m of the SOPC bilayer at 20°C is calculated at $\mu_m = 33$ P. The apparent radius of BR was then obtained from:

$$\frac{R_{\text{BR}}}{R_{L_{18}}} = \exp \left[\left(1 - \frac{D_{\text{BR}}}{D_{L_{18}}} \right) \times \left(\ln \left(\frac{\mu_m h}{\mu_w R_{L_{18}}} \right) - 0.5772 \right) \right] = 880;$$

$$R_{\text{BR}} = 880 \times 5.5 = 4,840 \text{ Å}.$$

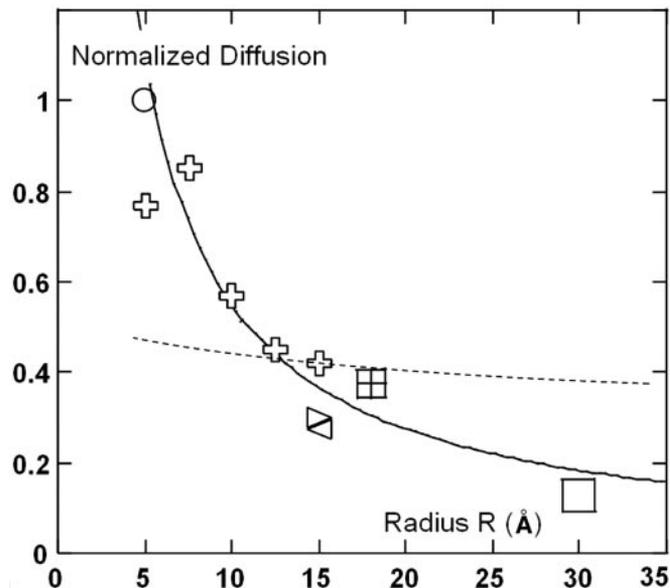


Fig. 2. Normalized diffusion coefficient (D/D_{lipid}) vs. peptide radius R in lipid bilayers. Crosses correspond respectively from the left to monomers, dimers, trimers, tetramers, and hexamers of transmembrane peptides (15). The square symbols at $R = 15, 18,$ and 30 Å correspond, respectively, to acetylcholine receptor (AChR), BR, and SR-ATPase (16). The solid line is a $1/R$ fit, and the dashed line represents the prediction of Saffman's model, using $h = 28$ Å, $\mu_m = 1.75$ P, and $\mu_w = 1$ cP as in ref 15.

diffusion coefficients of transmembrane peptides (15) and various proteins (16). The D variations, normalized by the lipid diffusion, show a continuous decrease of mobility, fitted by a simple $1/R$ law.

Furthermore, two recent studies allow the comparison of diffusion coefficients of two large transmembrane objects embedded in the same membrane. In the first (17), the mobilities of MscL [$R = 25$ Å (18)] and LacS [$R = 32$ Å (19)] differ by a factor of 1.3 corresponding to the ratio of their radii. In the second (14), a $1/R$ law fits correctly the data obtained by FCS and freeze-fracture electron microscopy on photo-activated, oligomerized BR.

We then switched back to the $C_{12}E_5$ bilayers, with incorporated proteins, for further investigation (Fig. 3). A simple test was first performed by using biotinylated L_{12} peptides embedded in bilayers adapted to match their length ($h = d_x = 21$ Å).

When water-soluble streptavidin is grafted onto one transmembrane peptide, the observed decrease of $D_{L_{12}}$ is only 3%. However, when the peptide concentration is twice as large as the streptavidin concentration, stable dimers are formed. The diffusion coefficient of such dimers is half that found for monomers. The object created is anisotropic, but the equivalent size calculated using Eq. 1 seems grossly overestimated: the dimer would be equivalent to a cylinder of radius $R_{\text{Saffman}} = 138$ Å (considering the viscosity of the model bilayers $\mu_m = 2.94$ P, its thickness $h = 21$ Å, and $\mu_w = 10^{-2}$ P). On the contrary, as indicated Fig. 3, the variation observed is compatible with a $1/R$ dependence.**

The above experiments confirm that the diffusion coefficient is rather insensitive to the size R_w of the polar heads, because of the high viscosity contrast ($\approx \mu_m h / \mu_w R_w$) between the bilayer

**Peptides are separated by 20 Å because of the structure of streptavidin. The dimer thus forms an anisotropic object of minor axis 5.5 Å and major axis 15.5 Å. From the area covered: $11 \times (5.5 + 20 + 5.5) = 341 \text{ Å}^2$, and using $R = \sqrt{\text{Area}/\pi}$, we estimated at 10.5 ± 1 Å the radius of the assembly created.

It should be emphasized that, on molecular length-scales, the membrane is not a featureless continuum, because the lipid chains possess finite sizes and internal degrees of freedom. Molecular dynamics “snapshots” of bilayers show a very heterogeneous medium (26) with defects in the membrane packing. Supporting this view, a recent mesoscopic simulation (27) showed that “the diffusive process occurs through collective and correlated motions of lipids,” with a correlation length scale of ≈ 10 Å. On larger length scales, thermal fluctuations and undulations also may contribute to the dissipation of velocity gradients.

One can expect, however, that in the limit of macroscopic inclusions, such that $R \gg \lambda$, the analysis of Saffman–Delbruck still applies. At this stage, more theoretical work seems to be the next logical step.

Conclusion

Our data raise physical questions concerning our understanding of the bilayer “fluid.” The heuristic law we propose opens perspectives for simple and widespread photobleaching or correlation spectroscopy techniques, which will give information about the peptide conformation or protein oligomerization in model systems of giant vesicles. In living cells, the fact that, even in the absence of physical barriers, clusters of proteins should diffuse much slower than dispersed receptors may be decisive when analyzing single particle tracking experiments.

Materials and Methods

Peptides. The synthetic peptides mimic the hydrophobic transmembrane α -helical segments of integral membrane proteins (28). They consist of a simple hydrophobic α -helix, capped at one or both ends by polar heads and labeled with a fluorescent group (FITC) or with biotin to enable an association reaction. A sequence of poly-leucines of variable length (12, 18, or 24 aa) creates a robust and well characterized transmembrane domain; the anchoring to the surface of the bilayer is ensured by positively charged lysine residues. Fig. 1 illustrates and summarizes their properties.

Synthesis. Commercial solvents of analytical grade were redistilled before use. Peptides were synthesized and purified as trifluoroacetic acid (TFA) salts, using previously published solid-phase synthesis and reversed-phase high-performance liquid chromatographic procedures (29, 30).

Proteins.

- (i) OprM is a large protein of the outer membrane of *Pseudomonas aeruginosa*, whose structure is close to that of TolC; its hydrophobic domain is a β -barrel of 12 strands (21).
- (ii) Eight strands constitute the transmembrane domain of OmpA (20); its purification and handling are described in ref. 31.
- (iii) BR (12) is a protein with multitransmembrane segments, purified from purple membranes of *Halobacterium salinarium*. The BR was kept in its dark-adapted state during all experiments.

These proteins were labeled as described in ref. 11 or using Fluoreporter kits purchased from Molecular Probes and following the manufacturer’s procedures.

Preparation of the GUVs. The vesicles were made of SOPC as the main component. The phospholipids used in this study were purchased from Avanti Polar Lipids and used without further purification.

Peptides were dissolved with lipids in chloroform, and the mixture (1 peptide for 10^3 lipids) was spread and evaporated on rough Teflon substrates. After lipid hydration, giant vesicles

were formed in 270 milliosmolar (mOsm) glucose solution (32, 33). The vesicle suspension was added to an aqueous sucrose solution or PBS buffer of a slightly higher osmolarity (290 mOsm) than that of the vesicles. The deflated vesicles can be micromanipulated and present a flat area in contact with the surfaces suitable for evanescent photobleaching. For the BR incorporation, an electroformation procedure was used. See ref. 11 for details.

Lyotropic Phases. Choosing $C_{12}E_5$ as a suitable component, we found that the common surfactant used with transmembrane proteins, β -octylglucopyranoside (β -OG), acts as a perfect co-surfactant. Simply added to a mixture of water and $C_{12}E_5$, β -OG helps the surfactant to create spontaneously a regular network of multiconnected bilayers and controls the properties of these “sponge” or L_3 phases over the wide range of conditions (buffer, temperature, etc.) desired. The phases used in these experiments are mixtures of water: $C_{12}E_5$: β -OG in a 100:30:1 volume ratio.

The structure of the sponge phases has been extensively studied (34, 35). Small-angle x-ray scattering experiments were performed as a careful check of the phase properties, as described in refs. 36 and 37.

The thickness of such bilayers is perfectly controlled and can be increased up to 3-fold through the precise addition of a hydrophobic solvent (10). To simplify the analysis, we chose the solvent to keep the viscosity of the membrane constant upon swelling.

We tested the properties of the monolayers by labeled surfactants (C_{12} -FITC), and the bilayer by a labeled lipid (SOPC-4-chloro-7-nitrobenz-2-oxa-1,3-diazole). The lipid inserts into one hydrophobic monolayer (8 Å) and tests the viscosity of an extra 6 Å. For dry bilayers [$h = 16$ Å (38)], the lipid tail contacts the opposite leaflet; upon swelling, midplane solvent progressively replaces that contribution to the friction.

Using dodecane, which matches the hydrophobic tails of the surfactant, the swelling has no apparent effect on the viscosities. D_{SOPC} remains constant ($5 \pm 0.2 \mu\text{m}^2/\text{s}$), and $D_{C_{12}}$ stays at $8 \pm 0.2 \mu\text{m}^2/\text{s}$. Using shorter chains, the viscosity of the monolayer is decreased, whereas branched hydrocarbons cause an increase of the midplane friction.

To avoid insertion defects, the peptides were incorporated into dry bilayers of $C_{12}E_5$; dodecane then was added in a step-by-step process. To eliminate the possibility of destructive interactions between dodecane and the complex transmembrane domains of the proteins, OprA, OprM, and BR were embedded in dry bilayers.

In this case, the protein hydrophobic height is larger than the hydrophobic thickness of the bilayer, creating a local deformation. Different studies (26, 39) estimate that this perturbation extends on the length scale of one to two surfactant molecules. Such a surfactant annulus may either diffuse along with the protein as a compact block or be constantly regenerated as surfactant molecules adjacent to or near the protein exchange with those further away. Thus, the pertinent dimension of the diffusing “object” could be either equal to the protein radius or 10 Å larger.

The hypothesis of aggregation of diffusing species, leading to much larger radii, is unlikely at our protein/surfactant ratio of $1:10^6$, and the recovery of fluorescence would not be purely monoexponential as observed.

Streptavidin-peptide assemblies were prepared as follows.

- (i) in the absence of peptides, we checked that the high mobility ($15 \mu\text{m}^2/\text{s}$) of streptavidin corresponds to this large soluble protein (50 Å in diameter), confined between the bilayers but without interactions with these bilayers. (Streptavidin labeled with FITC was purchased from Interchim, Montluçon, France.)

- (ii) By adding peptides, we followed the grafting of streptavidin; when one equivalent of peptide is incorporated, we observed a monoexponential recovery of fluorescence, with a characteristic time matching the one obtained with labeled L₁₂ peptides under the same conditions. In that case, the streptavidin molecules are mainly bound to a unique peptide.
- (iii) By adding an excess of peptides (from 2 up to 10 equivalents of peptide), we observed a different diffusion coefficient, showing that streptavidin now associates laterally two peptides. The water spacing between the bilayers is kept at 300 Å for this set of experiments, avoiding interactions between molecules embedded in two different bilayers.

Preparation of the Samples and Concentrations Used. To focus on effects arising solely from individual behavior, we performed concentration-dependent measurements. As the detection is made over large areas, the peptides and proteins were added at minimal concentrations, removing the effects of protein–protein interaction. Typically, for one peptide, 10⁴ to 10⁶ surfactants were used in the lyotropic phase, and 10³ lipids for the GUVs experiments. A typical sample is only 5 μl in volume, introduced by suction into 200-μm-thick flat microchannels. The temperature was controlled at 20°C, while the sponge phases are stable over a 10–35°C temperature range. Any phase transition is easily detectable, given its well known defect features observable under the microscope between crossed polarizers; systematic controls were performed between each experiment.

Diffusion Measured by Fluorescence Recovery After Pattern Photo-bleaching. The fluorescence recovery after pattern photobleaching technique is described in detail in ref. 40. Briefly, when

illuminated by a high-intensity laser flash (200 mW), the fluorescent molecules dispersed within the sample are irreversibly bleached. The recovery of fluorescence intensity with time, $I(t)$, in the bleached areas is governed by the self-diffusion of the unbleached probes and monitored by a low-intensity laser beam. To improve the signal-to-noise ratio, we created a macroscopic gradient of fluorescence following a fringe pattern, using two laser beams intersecting exactly in the focal plane of a light microscope. A piezoelectric crystal made the monitoring beam sweep the bleached fringes at a frequency, ω , and the signal was detected at ω and 2ω using a lock-in amplifier. A nonzero signal at ω would betray convection, while at 2ω the enhanced recovery signal was obtained. In all experiments, monoexponential recoveries of fluorescence were measured: the signal at 2ω is proportional to $I_0[1 - \exp(-t/\tau)]$. A typical measure of diffusion consists of five characteristic times τ corresponding to five different fringe sizes i , with each τ being determined from an average of 10 experiments. We checked that all of the diffusive behavior followed Brownian motion, and the diffusion coefficients were deduced by the classical relation $D = i^2/(4\pi^2\tau)$, where the interfringe values i are in the range 1–300 μm and the typical time values are from 0.1 to 10³ s.

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