Rationalizing Steroid Interactions with Lipid Membranes: Conformations, Partitioning, and Kinetics

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ABSTRACT: Steroids have numerous physiological functions associated with cellular signaling or modulation of the lipid membrane structure and dynamics, and as such, they have found broad pharmacological applications. Steroid–membrane interactions are relevant to multiple steps of steroid biosynthesis and action, as steroids are known to interact with neurotransmitter or membrane steroid receptors, and steroids must cross lipid membranes to exert their physiological functions. Therefore, rationalizing steroid function requires understanding of steroid–membrane interactions. We combined molecular dynamics simulations and isothermal titration calorimetry to characterize the conformations and the energetics of partitioning, in addition to the kinetics of flip–flop transitions and membrane exit, of 26 representative steroid compounds in a model lipid membrane. The steroid classes covered in this study include birth control and anabolic drugs, sex and corticosteroid hormones, neuroactive steroids, as well as steroids modulating the lipid membrane structure. We found that the conformational ensembles adopted by different steroids vary greatly, as quantified by their distributions of tilt angles and insertion depths into the membrane, ranging from well-defined steroid conformations with orientations either parallel or normal to the membrane, to wide conformational distributions. Surprisingly, despite their chemical diversity, the membrane/water partition coefficient is similar among most steroids, except for structural steroids such as cholesterol, leading to similar rates for exiting the membrane. By contrast, the rates of steroid flip–flop vary by at least 9 orders of magnitude, revealing that flip–flop is the rate-limiting step during cellular uptake of polar steroids. This study lays the ground for a quantitative understanding of steroid–membrane interactions, and it will hence be of use for studies of steroid biosynthesis and function as well as for the development and usage of steroids in a pharmacological context.

INTRODUCTION

Steroids are a heterogeneous group of typically hydrophobic organic compounds characterized by a tetracyclic fused-ring core (Figure 1). Steroids have various functions in cells and are involved in numerous metabolic pathways. Certain steroid compounds modulate the structure of biological membranes, typical examples including cholesterol in animals, β-sitosterol in plants, and ergosterol in fungi. Other steroids function as signaling molecules, such as corticosteroid and sex hormones. Steroids have found wide pharmacological applications in, among others, anti-inflammatory drugs, birth control, anesthetics, and cancer treatment, and they are frequently abused to improve performance in work or sports.

The interactions of steroid hormones with biological membranes are relevant to many aspects of their functions. The classical action of steroid hormones entails binding to intracellular steroid receptors, which ultimately results in changes in gene expression. To this end, steroids have to be internalized into cells. According to the free hormone hypothesis, because of their hydrophobicity, steroids are able to freely diffuse across lipid bilayers; however, megalin-dependent endocytosis has been shown to be at least partly responsible for the uptake of sex hormones and vitamin D₃. In addition to their classical genomic action, also nongenomic mechanisms of action are known for certain steroids. This includes neuroactive steroid compounds, which interact with neurotransmitter receptors and modulate neuronal excitability, as well as steroid actions mediated by membrane steroid receptors.
of steroid molecules may be influenced by their interactions with lipid membranes.\textsuperscript{17}

Hence, rationalizing the metabolic functions of steroids requires understanding of steroid–membrane interactions. The best-studied steroids are probably the long-tailed sterols, such as cholesterol. Using both experimental and computational approaches, the effects of cholesterol on membrane structure, the conformations and partitioning of cholesterol in bilayers, cholesterol–lipid interactions, and cholesterol flip–flop transitions have been described in great detail.\textsuperscript{1,17–29} By contrast, the literature on other steroids is less abundant. Partition coefficients have often been reported for octanol or bilayers from cell extracts, while data on partitioning in model lipid bilayers of controlled composition are limited.\textsuperscript{30–38} Computational studies on the properties of steroids in bilayers have usually been done for only a few steroids, some using short simulations, coarse-grained models, or implicit solvent.\textsuperscript{26,39–43}

Here, we provide a comprehensive overview of the conformational, energetic, and kinetic characteristics of steroids in a model lipid membrane of 1-palmitoyl-2-oleoyl-sn-glycerol-3-phosphocholine (POPC). We used all-atom molecular dynamics (MD) simulations to compare the steroid–membrane interactions of 26 steroid compounds, as quantified by their positions in the membrane and distributions of tilt angles, as well as by their membrane/water partition coefficients, and kinetics of flip–flop and membrane exiting (Figure 1). To this end, we derived force field parameters for 26 steroids, and we refined these parameters against membrane/water partition coefficients obtained from isothermal titration calorimetry (ITC) or the literature. The all-atom simulations, complemented by calorimetric data, provide an atomic-level view of the conformations, energetics, and kinetics of the steroids in a lipid membrane.

Figure 1. (A–F) Steroids considered in this study. The groups on C-3 (“head”) and C-17 (“tail”) atoms are color-coded as in the following figures. (G) Nomenclature of the tetracyclic steroid core.
RESULTS AND DISCUSSION

Orientational Diversity of Steroids in the Membrane. Steroids are overall hydrophilic because of their common tetracyclic hydrocarbon core (Figure 1G). However, they carry different functional groups at various positions, which influence the orientation and position of the steroid in the membrane (Figure 1A–E). For instance, the conformation of cholesterol is imposed by the polar hydroxyl group at the C-3 atom and the aliphatic tail attached to C-17 (Figure 1E). This specific configuration of functional groups imposes a vertical orientation (i.e., parallel to the membrane normal), positioned such that the aliphatic tail is solvated by the lipid tails, whereas the hydroxyl group can form hydrogen bonds with the polar lipid head groups or water.26

Analogously to cholesterol, we defined atoms C-3 and C-17 as “head” and “tail” atoms, respectively, and derived the steroid orientation in the membrane (Figure 2). Figure 3 presents the average of the cosine of the tilting angle between (i) the steroid axis, connecting the head and tail atoms, and (ii) the z-axis, normal to the membrane plane. Hence, cos(α) = 1 denotes a vertical orientation in the membrane, with the steroid core oriented as in cholesterol; cos(α) = 0 indicates that the steroid is oriented horizontally, and cos(α) = −1 indicates an inverted vertical orientation, with the A-ring toward the membrane core (see Figure 1G). Values between these three special cases denote a tilted orientation of the steroid with respect to the membrane normal.

Figure 3 demonstrates that a steroid core alone does not impose any consensus orientation in the membrane shared by all steroids. Instead, different steroids adopt different orientations (Figure 4), depending on the functional chemical groups (Figure 3, colored bars). In addition, large standard deviations of cos(α) found for many steroids suggest that they do not assume a single well-defined orientation but instead a wide distribution of orientations (Figure 3, error bars). Indeed, the complete cos(α) distributions presented in Figure S1 reveal wide orientational distributions of, for instance, estrogen and corticosteroid hormones (see also β-estradiol and hydrocortisone in Figure 4A,1). In some cases, the cos(α) distribution exhibits multiple peaks, indicating that these steroids can adopt multiple distinct orientations, as found for testosterone, 4-androstenedione (Figure S1, Figure 4C,D), dihydrotestosterone, and levonorgestrel.

A well-defined vertical orientation is observed for steroids with a clear distinction between the hydrophilicity of the head and tail functional groups, respectively (Figure 3, left bars). These include primarily the sterols with a hydroxyl head group and a long aliphatic tail (i.e., cholesterol, β-sitosterol, and dehydroergosterol, see also Figure 4J). Pregnenolone and pregnenolone sulfate are likewise oriented vertically because of a combination of a hydroxyl or negatively charged head group, respectively, with a relatively hydrophobic tail group with limited capabilities of forming hydrogen bonds (Figure 4F,G). On the other hand, steroids with identical or similar head and tail groups tend to lie horizontally in the membrane, such as β-estradiol with a hydroxyl group on each end, or 4-androstenedione with a keto group on each end (Figure 3, middle bars; Figure 4A,D).

The most frequent substitutions at the head C-3 atom are hydroxyl and keto groups (Figure 3A). The hydroxyl group can form hydrogen bonds with the ester moiety of the POPC lipid heads as well as with water molecules found in this region. Therefore, a hydroxyl group in this position tends to favor a vertical or tilted orientation, such that ring A points toward the membrane surface. The only exceptions are the steroids where the tail atoms also carry hydroxyl groups, leading to a horizontal or even slightly downward-tilted orientation, as seen for estriol with two hydroxyl groups on ring D as opposed to only one hydroxyl group in ring A (Figure 4B). On the other hand, a keto group on the C-3 atom leads to a slight “inverted” tilt, such that ring A points to the membrane core (Figure 3A, light blue bars). This is rationalized by the fact that keto groups are purely hydrogen-bond acceptors but not donors, leading to reduced possibilities of forming hydrogen bonds. Therefore, when coupled with a hydroxyl group at the steroid tail, the head tends to sink deeper into the membrane than the tail, leading to an inverted orientation. Of course, when a similar or more hydrophobic group is present at the tail, such as in 4-androstenedione or progesterone, an average horizontal orientation or noninverted tilted orientation is found (Figure 4D,E). Corticosteroids, which in addition to the hydroxyl-containing group on C-17 and the keto group on C-3 have an additional polar group on atom C-11, similarly adopt a horizontal orientation (Figure 4I).

Insertion Depth of Steroids in the Membrane. In addition to the orientation, another major degree of freedom is given by the vertical position of the steroid relative to the membrane center of mass, that is, the insertion depth in the membrane (Figure 2). Figure 5 shows the mean vertical position (Δz) of the head and tail atoms, where Δz = 0 denotes the membrane center of mass (COM; Figure 5, dots and triangles). As expected, because of their overall amphiphilic nature, many steroids tend to localize below or near the ester groups of POPC (Figure 5, red horizontal line), at the interface between the polar and apolar regions of the membrane. However, the vertical position of the steroid is clearly modulated by the chemical modifications: steroids that have no hydrogen-bond donors sink deeper into the membrane than steroids that carry hydroxyl groups. For instance, pregnenolone acetate with zero hydroxyl groups is localized closer to the membrane COM than estriol with three hydroxyl groups (Figure 4H,B).

Likewise, the vertical positions of the individual head and tail atoms are strongly modulated by the chemical modifications, in line with the orientational diversity of steroids (previous section). For instance, the negatively
Figure 3. Tilting of the steroids with respect to the membrane normal, quantified by the mean of the cosine of the tilting angle between the steroid axis and the membrane normal (mean and SD over 500 ns and 14 steroid molecules). The bar plots are colored according to the functional group on (A) the C-3 ("head") atom and on (B) the C-17 ("tail") atom (see legend and Figure 1). Values of cos(α) of 1, 0, and −1 indicate a vertical orientation (as in cholesterol), a horizontal orientation, and an inverted vertical orientation, respectively. For cholesterol, results from two different force fields are shown (GAFF and Slipids, see the Methods section in the SI).
cyclohexane roughly correlates with the number of hydroxyl groups. Namely, steroids without hydroxyl groups exhibit negative $\Delta G_{\text{part}}^{\circ}$, indicating a preference for the apolar solvent (Figure 7B, bright yellow), while steroids with only one hydroxyl yield mostly $\Delta G_{\text{part}}^{\circ} \approx 0$ (Figure 7B, bright orange). Steroids with two and three hydroxyl groups mostly prefer water over cyclohexane, with corticosteroids, which have additional polar groups (such as carbonyl/keto groups),...
showing the most positive \( \Delta G_{\text{part}}^{\text{exp}} \) values (Figure 7B, orange and red). The number of carbonyl groups on the steroid further modulates \( \Delta G_{\text{part}}^{\text{exp}} \), as indicated by the number of asterisks in Figure 7B. Compared with hydroxyl groups, however, carbonyl groups have a smaller effect on \( \Delta G_{\text{part}}^{\text{exp}} \) owing to their lower polarity.

Compared with the water/cyclohexane partition free energies, water/membrane \( \Delta G_{\text{part}}^{\text{exp}} \) values are much less dependent on the chemical modifications (Figure 7A). The only exceptions are long-tailed steroids that exhibit more negative \( \Delta G_{\text{part}}^{\text{exp}} \) than all other steroids (Figure 7A, dark gray), indicating a strong preference for the membrane. For all other steroids, however, despite their chemical diversity, most \( \Delta G_{\text{part}}^{\text{exp}} \) values vary within \( \sim 10^2 \) kJ mol\(^{-1} \), corresponding to variations in the partition coefficient by a factor of only \( \sim 50 \). This finding is rationalized by the steroid location at the interface between the polar and apolar regions of the membrane. Here, upon changing chemical modifications on the steroid ring, rearrangements of the steroid’s orientation and depth are sufficient to maintain hydrogen bonds of polar groups, while keeping large parts of the apolar surface in contact with the apolar lipid tails. Consequently, chemical modifications have a much smaller effect on \( \Delta G_{\text{part}} \) as compared with \( \Delta G_{\text{part}}^{\text{exp}} \). This demonstrates that \( \Delta G_{\text{part}} \) is not explained by simple determinants such as the number of carbon atoms or number of hydroxyl groups. Instead, the finer modulations of \( \Delta G_{\text{part}} \) may depend on a combination of determinants, including configurational flexibility and specific steroid–lipid interactions, in addition to the overall hydrophobicity of the molecule.

**Kinetics of Steroid Flip–Flop and Membrane Exit.** To obtain the kinetics of steroid transitions at a POPC membrane, we computed transversal diffusion coefficients of the steroids (Figure S5) (see the Methods section in the SI). Rates for steroid flip–flop and for exiting the membrane were estimated following Kramers’ theory (Figure 8C, arrows).\(^{44} \) Figure 8A presents flip–flop rates, \( k_{\text{ff}} \) for all steroids considered in this study except for the anionic steroids, revealing that \( k_{\text{ff}} \) for steroids may span at least 9 orders of magnitude. Evidently, \( k_{\text{ff}} \) anticorrelates with the number of polar groups in the steroid, in particular with the number of hydroxyl groups (Figure 8A, color code). The number of carbonyl groups has only a smaller effect on \( k_{\text{ff}} \) (Figure 8A, asterisks). For steroids with zero or one hydroxyl group, including the long-tailed structural steroids, we found large \( k_{\text{ff}} \) values in the range \( 10^5 - 10^6 \) s\(^{-1} \), corresponding to rapid flip–flop events on the time scale of microseconds up to hundreds of microseconds. These values are in reasonable agreement with previous reports for cholesterol flip–flop.\(^{29,30} \) For steroids with two or three hydroxyl groups, by contrast, \( k_{\text{ff}} \) spans the range \( 10^2 - 10^3 \) s\(^{-1} \), corresponding to flip–flop events on the time scale of milliseconds up to many minutes. As discussed in the Methods section in the SI, for the most polar steroids we cannot exclude the possibility that the PMFs underestimate the true flip–flop barrier. Hence, the flip–flop events of the most polar steroids, such as aldosterone or hydrocortisone, could also occur on the time scale of hours or even longer.

The wide range of \( k_{\text{ff}} \) is readily explained by the wide range of water/cyclohexane partition free energies \( \Delta G_{\text{part}}^{\text{exp}} \) presented above (Figure 7B). Starting from a membrane-bound state (Figure 8C, \( z \approx \pm 1.2 \) nm), steroid flip–flop requires the transition across the hydrophobic membrane core, which involves the removal of most of the steroid–water contacts, similar to a transition from water to cyclohexane. Hence, the free-energy cost for steroid flip–flop correlates with the cost for translocating a steroid from water to cyclohexane (Figure S4). More quantitatively, \( k_{\text{ff}} \) is dictated by the height of the free-energy barrier in the transmembrane PMFs shown in Figure 8C. Here, the barrier height is the difference between (i) the free energy at the hydrophobic core (Figure 8C, \( z \approx 0 \) nm) approximately given via \( \Delta G_{\text{part}}^{\text{exp}} \) and (ii) the free-energy minimum at the membrane-bound state (Figure 8C, \( z \approx \pm 1.2 \) nm), approximately given via \( \Delta G_{\text{part}}^{\text{sim}} \). Hence, \( k_{\text{ff}} \) correlates with \( \exp(-\beta(\Delta G_{\text{part}}^{\text{exp}} - \Delta G_{\text{part}}^{\text{sim}})) \), where \( \beta = 1/RT \) is the inverse temperature. Further, since \( \Delta G_{\text{part}}^{\text{exp}} \) is similar among most steroids, \( k_{\text{ff}} \) is primarily dictated by \( \Delta G_{\text{part}}^{\text{sim}} \) (compare Figure 7B with Figure 8A). Notable exceptions are the long-tailed structural steroids; for these steroids, large negative \( \Delta G_{\text{part}}^{\text{sim}} \) and \( \Delta G_{\text{part}}^{\text{exp}} \) compensate each other, leading to similar flip–flop rates as steroids with one or without any hydroxyl group (Figures 7A,B and 8A, light gray bars).

In contrast with the wide range of \( k_{\text{ff}} \) values, the rates \( k_{\text{exit}} \) for exiting the membrane are highly similar among most of the steroids (Figure 8B). Most \( k_{\text{exit}} \) values are in the order of \( 10^4 \) s\(^{-1} \) corresponding to rapid exit events on the time scale of only hundreds of microseconds. Hence, steroids bind to membranes in a highly transient manner. Exceptions are again the long-tailed steroids that exhibit \( k_{\text{exit}} \) in the range \( 10^{-3} - 10^{-2} \) s\(^{-1} \), indicating that the long-tailed steroids bind tightly to the membranes for minutes up to many hours, in excellent agreement with experimental findings for cholesterol.\(^{19} \) Since exiting the membrane requires overcoming the free energy of membrane/water partitioning, \( k_{\text{exit}} \) strongly correlates with \( \exp(-\beta \Delta G_{\text{part}}^{\text{exp}}) \) (compare Figure 7A with Figure 8B). Hence, the similarity of \( k_{\text{exit}} \) among the nonstructural steroids is a consequence of the similarity of \( \Delta G_{\text{part}}^{\text{exp}} \).

**Kinetics of Membrane Permeation.** Membrane permeation requires (i) steroid entering the membrane, (ii) followed by at least one flip–flop event, and (iii) membrane exiting in
the opposite direction as compared with membrane entry. We estimated rates for membrane entry to be in the order of $10^{-1} \text{s}^{-1}$ for eukaryotic cellular environments, $10^{-2} \text{s}^{-1}$ for planar membranes with pronounced unstirred layers, and $10^{5} \text{s}^{-1}$ for large unilamellar vesicles (LUVs) with a typical radius of $0.1 \mu\text{m}$ (see the Methods section in the SI). Comparing $k_{\text{entry}}$ with $k_{\text{ff}}$ and $k_{\text{exit}}$ shown in Figure 8 A,B reveals that different transitions may become limiting for steroid permeation (Table 1). Namely, membrane permeation for structural long-tailed steroids is limited by slow membrane exit ($k_{\text{exit}}$). Permeation for steroids with 2 or 3 OH groups is limited by flip–flop. Permeation of steroids with 0 or 1 OH groups may be limited by entry (i.e., by unstirred layers) in cells or planar membranes, and by exit in LUVs.

In addition, the ratio $k_{\text{ff}}/k_{\text{exit}}$ determines the average number of flip–flop events before the steroid exits the membrane, and consequently, $k_{\text{ff}}/k_{\text{exit}}$ further determines the probabilities for the two possible directions of membrane exit. A rate ansatz (see the Methods section it the SI) shows that the probability for a full permeation event per membrane entry event is given by

$$P_{\text{perm}} = \left( \frac{k_{\text{exit}}}{k_{\text{ff}}} + 2 \right)^{-1}$$

while the probability that the steroid returns to its original water compartment is $P_{\text{ret}} = 1 - P_{\text{perm}}$. In the case of $k_{\text{ff}} \gg k_{\text{exit}}$, the steroid typically carries out multiple flip–flop events before its exit with equal probability in each direction ($P_{\text{perm}} \approx P_{\text{ret}} \approx 1/2$). By contrast, in the case of $k_{\text{ff}} \ll k_{\text{exit}}$, the steroid will mostly return to its original water compartment before the first flip–flop event occurs ($P_{\text{perm}} \approx 0, P_{\text{ret}} \approx 1$), which may strongly reduce cellular uptake rates (see discussion below).

Among all steroids, $k_{\text{ff}}/k_{\text{exit}}$ varies in the range $10^{-9} - 10^{10}$, and among the nonstructural steroids in the range $10^{-9} - 10^{3}$, demonstrating that the probability for permeation after membrane binding, $P_{\text{perm}}$, greatly varies. Specifically, for steroids without hydroxyl groups we obtain $P_{\text{perm}} \approx 1/2$. For steroids with one hydroxyl group, $P_{\text{perm}}$ drops to values between 0.1 and 0.45. For steroids with two or three hydroxyl groups, $P_{\text{perm}}$ may take values as low as $10^{-10}$, demonstrating that many membrane binding events are needed before the most polar steroids permeate the membrane.

Figure 7. Free energies of partitioning (A) from water to a POPC bilayer, $\Delta G_{\text{part}}^\circ$, and (B) from water to cyclohexane, $\Delta G_{\text{part}}^\circ,\text{cyc}$. Where available, experimentally determined values are shown in part A (all steroids shown in Figure 6); for the rest of the steroids, PMF-derived values are shown. Values in part B were obtained by TI. Error bars represent 95% confidence intervals for the ITC-derived values, and standard errors for the PMF- and TI-derived values. Bars without error bars represent experimental data from the literature for which no errors were available. The coloring indicates the number of hydroxyl groups, long-tailed, or anionic steroids (see legend). The number of asterisks on top of the bars in part B indicates the number of carbonyl groups.
To resolve the apparent discrepancy between (i) experimentally observed rapid responses of cells to steroid exposure and (ii) computationally derived slow partition coexpression, reaching the transmembrane receptors. Therefore, the partitioning of the steroids into the membrane is an important determinant of the interaction and partly determines the kinetics of steroid binding to transmembrane receptors.56 Typically conducted at constant steroid concentration in the bulk solvent. Consequently, because of the hydrophobicity of the steroids, the steroids are greatly enriched in the outer membrane leaflet as quantified by the membrane/water partition coefficients. More quantitatively, using the \( \Delta G^\text{out}_{\text{net}} \) definition shown in the Methods section of the SI, the partition free energies, \( \Delta G^\text{out}_{\text{net}} \), of \(-20\) to \(-35\) kJ mol\(^{-1}\) (Figure 7) suggest that steroids are enriched in the membrane by a factor between 75 and 30 000 as compared with the bulk. This enrichment largely compensates for low flip–flop rates, thus leading to high permeabilities and hence to cell entry of a significant number of steroid molecules within seconds. As such, rapid entry of steroids into the cell is, for polar steroids, not a consequence of “unhindered” diffusion over the membrane, but instead a consequence of steroid enrichment in the outer membrane leaflet.

**Partitioning between the Extracellular Bulk Solution and the Plasma Membrane Can Be an Important Determinant of Steroid Potency.** Several steroids are known to influence the function of transmembrane proteins. In most cases where evidence has been obtained, e.g., in the metabotropic CB1 receptors,49 the bacterial channel GLIC50 and the ionotropic GABA\(_A\)51,52 nicotinic ACh,53 and NMDA receptors,54,55 it has been shown that steroids interact with these targets on transmembrane helices. For many other steroid receptor transmembrane proteins, a binding site in the transmembrane region also seems likely, although the location of the binding site has not yet been established with certainty. To reach their target, steroids supplied with the bloodstream must first partition into the membrane and then, by lateral diffusion, reach the transmembrane receptors. Therefore, the partitioning of the steroids into the membrane is an important determinant of the interaction and partly determines the kinetics of steroid binding to transmembrane receptors.

**Significance of Membrane Permeation for Steroid Function.** At first sight, the wide range of flip–flop rates might seem at odds with the textbook assertion that steroid hormones pass biological membranes “freely” or “unhindered”. Experimentally, it has indeed been observed that membrane crossing of the classical steroid hormones is a fairly rapid process. For instance, the movement of intracellular mineralocorticoid receptors after steroid binding into the cell nucleus can be detected within 3 min after extracellular aldosterone application.46 Furthermore, nonclassical effects such as an aldosterone-induced rise of intracellular Ca\(^{2+}\) concentration have been observed even within seconds.47,48 In either case, complete membrane traversal of aldosterone is necessary, as the presumed target molecules of aldosterone, whether they be mineralocorticoid receptors or nonclassical targets, are located intracellularly. The rapid action of aldosterone is particularly remarkable in this respect, as our simulations indicated that the flip–flop rate of aldosterone is the lowest among all steroids tested (apart from the anionic, sulfonated steroids that would require either protonation or an aqueous defect to flip–flop).

To resolve the apparent discrepancy between (i) experimentally observed rapid responses of cells to steroid exposure and (ii) computationally derived slow flip–flop rates for polar steroids, it is important to notice that such experiments are typically conducted at constant steroid concentration in the bulk solvent. Consequently, because of the hydrophobicity of the steroids, the steroids are greatly enriched in the outer membrane leaflet as quantified by the membrane/water partition coefficients. More quantitatively, using the \( \Delta G^\text{out}_{\text{net}} \) definition shown in the Methods section of the SI, the partition free energies, \( \Delta G^\text{out}_{\text{net}} \), of \(-20\) to \(-35\) kJ mol\(^{-1}\) (Figure 7) suggest that steroids are enriched in the membrane by a factor between 75 and 30 000 as compared with the bulk. This enrichment largely compensates for low flip–flop rates, thus leading to high permeabilities and hence to cell entry of a significant number of steroid molecules within seconds. As such, rapid entry of steroids into the cell is, for polar steroids, not a consequence of “unhindered” diffusion over the membrane, but instead a consequence of steroid enrichment in the outer membrane leaflet.

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**Table 1. Rate-Limiting Steps (Membrane Entry, Flip–Flop, or Membrane Exit) for Membrane Permeation Depending on the Steroid Structure and the Type of Membrane**

| OH groups | cell entry | planar membrane exit | LUV (radius 0.2 \( \mu \)m) exit |
|-----------|------------|----------------------|----------------------------------|
| long-tailed | exit | exit | exit |
| 0–1 OH groups | entry | entry | exit |
| 2–3 OH groups | flip–flop | flip–flop | flip–flop |

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**Figure 8.** Rates (A) for steroid flip–flop and (B) for exiting the membrane. The coloring indicates the number of hydroxyl groups or long-tailed steroids (see legend). The number of asterisks on top of the bars in part A indicates the number of carbonyl groups. (C) PMFs for four selected steroids along the membrane normal \( z \), where \( z = 0 \) is the membrane center (see legend for color code). Arrows illustrate the transitions for flip–flop and membrane exiting.
For instance, the effects of the structurally very similar steroids pregnenolone sulfate and dehydroepiandrosterone sulfate (DHEAS) have been studied in GABA<sub>A</sub> receptors and TRPM3 channels. While in TRPM3 channels, the EC<sub>50</sub> for pregnenolone sulfate is 13–25 times lower than for DHEAS,<sup>57</sup> in GABA<sub>A</sub> receptors, DHEAS has been reported to be approximately equally efficient in inhibiting Cl<sup>−</sup> currents through these receptors. Because our results indicate that the concentration of pregnenolone sulfate in the plasma membrane is about 40 times larger than the DHEAS concentration (at the same bulk concentration in the extracellular solution), these findings indicate that the binding site of steroids on TRPM3 channels<sup>59</sup> is only poorly discriminating between pregnenolone sulfate and DHEAS. On the other hand, our data suggest that membrane-bound DHEAS has stronger effects on GABA<sub>A</sub> compared to pregnenolone sulfate. This example demonstrates that a quantitative understanding of steroid–membrane interactions, as derived in this work, is needed for a detailed interpretation of the experimentally observed receptor response.

**Functional Consequences of the Position and Orientation of Steroids in the Membrane.** Before binding to a proteinaceous binding site on transmembrane segments, steroids must adopt an orientation and an insertion depth that matches the binding site. In addition, if the binding site is located at the intracellular membrane leaflet, at least one flip–flop event is required for binding. Previously, these requirements have complicated a molecular interpretation of experiments. For instance, if a certain steroid shows no (or weak) activity on a receptor, it remains unclear whether (i) the affinity for the steroid is low, (ii) steroid binding does not trigger a relevant conformational transition of the protein, or (iii) whether the steroid does not reach the binding site because of unfavorable orientations adopted in the membrane. Our simulations showed that most (but not all) steroids adopt wide conformational distributions (Figures S1 and S2), in terms of both steroid orientation and insertion depth. Hence, unfavorable conformations may, for most steroids, be excluded as an underlying reason for weak steroid activity.

To illustrate this, it is instructive to pick two extreme examples: Pregnenolone acetate is completely inactive on TRPM3 channels, while pregnenolone sulfate is a strong agonist.<sup>57,59</sup> Pregnenolone sulfate is predominantly oriented perpendicular to the plasma membrane, whereas pregnenolone acetate has a strong preference for the orientation parallel to the phospholipid bilayer, but also samples other orientations. Also, these molecules can be found at various depths inside the membrane. These observations indicate that pregnenolone acetate is incapable of activating TRPM3 channels not only because of its unfavorable orientation and position within the membrane. Rather, because this substance does not have any appreciable effect on TRPM3,<sup>59</sup> either this steroid cannot bind to TRPM3 proteins, or its binding does not induce channel opening. For agonist activity on TRPM3 channels, either bulky or negatively charged (or both) head groups on the C3 position of pregnenolone appear to be indispensable.<sup>59</sup> More generally, though, the observed rapid and wide-ranging movements observed for many of the steroids indicate that these molecules are capable of approaching and docking to membrane-embedded binding sites regardless of their average orientation. Unfavorably oriented binding sites, however, would exhibit reduced rates of binding.

**CONCLUSIONS**

The broad spectrum of steroid compounds encompassed by this study provides a global view of the range of steroid–membrane interactions, highlighting similarities and differences among the family of steroids. Although steroids share a common structural core, they reveal greatly different conformational ensembles in a lipid membrane, imposed by the chemical modifications on the tetracyclic steroid core. Namely, certain steroids adopt well-defined conformational orientations, by orienting strictly either parallel or normal to the membrane, whereas other steroids reveal high orientational flexibility, hence adopting wide conformational ensembles.

For steroids that are neither long-tailed nor anionic, free-energy calculations revealed that the cyclohexane/water partition coefficients vary by 16 orders of magnitude. By contrast, membrane/water partition coefficients are surprisingly similar, varying by only 2–3 orders of magnitude. Further, we derived the kinetics of steroids in membranes, that is, the rates of steroid flip–flop and membrane exiting. We found that rates of membrane exiting are remarkably similar among many steroids, whereas flip–flop rates vary by many orders of magnitude. These trends for steroid flip–flop and exiting rates are rationalized by the trends of cyclohexane/water and membrane/water partition coefficients.

Exceptions are given by the long-tailed steroids such as cholesterol or dehydroergosterol, as well by the anionic steroids such as pregnenolone sulfate; namely, long-tailed steroids exhibit greatly increased membrane affinity and greatly decreased membrane exiting rates, but they display similar flip–flop rates compared to most other steroids. Anionic steroids exhibit greatly reduced flip–flop rates because flip–flop would either involve translocation of the anionic group across the hydrophobic core or require protonation of the steroid; however, anionic steroids show similar membrane/water partitioning compared to most other steroids.

This study provides quantitative understanding of steroid–bilayer interactions, relevant to steroid permeation across the bilayer, as well as for steroid binding to transmembrane receptors and to other membrane proteins. The topologies for all steroids with refined partial atomic charges are available for download at https://biophys.uni-saarland.de/steroids.html.

**ASSOCIATED CONTENT**

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acscentsci.8b00332.

Additional experimental and computational methods, data, and figures including cos(α) distributions, Δz distributions, experimental vs calculated free energies, cyclohexane/water partition free energies vs the transfer free energies for steroids between the bulk and the membrane center, and transversal diffusion coefficients (PDF)

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Notes
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REFERENCES
(1) Yeagle, P. L. Cholesterol and the Cell Membrane. Biochim. Biophys. Acta, Rev. Biomembr. 1985, 822, 267–287.
(2) Bernsdorf, C.; Winter, R. Differential Properties of the Sterols Cholesterol, Ergosterol, β-Sitosterol, Trans-7-Dehydrocholesterol, Stigmasterol and Lanosterol on DPPC Bilayer Order. J. Phys. Chem. B 2003, 107, 10658–10664.
(3) Orme, M. L.; Back, D. J.; Breckenridge, A. M. Clinical Pharmacokinetics of Oral Contraceptive Steroids. Clin. Pharmacokinet. 1983, 8, 95–136.
(4) Haupt, H. A.; Rovere, G. D. Anabolic Steroids: A Review of the Literature. Am. J. Sports Med. 1984, 12, 469–484.
(5) Ericson-Neilsen, W.; Kaye, A. D. Steroids: Pharmacology, Complications, and Practice Delivery Issues. Ochsner J. 2014, 14, 203–207.
(6) Blickenstaff, R. T. Antitumor Steroids; Academic Press, 2012.
(7) Sjögqvist, F.; Garle, M.; Rane, A. Use of Doping Agents, Particularly Anabolic Steroids, in Sports and Society. Lancet 2008, 371, 1872–1882.
(8) Evans, R. M. The Steroid and Thyroid Hormone Receptor Superfamily. Science 1988, 240, 889–895.
(9) Beato, M.; Chávez, S.; Truss, M. Transcriptional Regulation by Steroid Hormones. Steroids 1996, 61, 240–251.
(10) Nykjaer, A.; Dragun, D.; Walther, D.; Vorum, H.; Jacobsen, C.; Herz, J.; Melsen, C.; Christensen, I. E.; Willnow, T. E. An Endocytic Pathway Essential for Renal Uptake and Activation of the Steroid 25-(OH) Vitamin D3. Cell 1999, 96, 507–515.
(11) Hammes, A.; Andreasen, T. K.; Spoelgen, R.; Raja, J.; Hubner, N.; Schulz, H.; Metzger, J.; Schweigert, F. J.; Lupa, P. B.; Nykjaer, A.; Willnow, T. E. Role of Endocytosis in Cellular Uptake of Sex Steroids. Cell 2005, 122, 751–762.
(12) Rupprecht, R.; Holstboer, F. Neuroactive Steroids: Mechanisms of Action and Neuropsychopharmacological Perspectives. Trends Neurosci. 1999, 22, 410–416.
(13) Falkenstein, E.; Tillmann, H. C.; Christ, M.; Feuring, M.; Wehling, M. Multiple Actions of Steroid Hormones—a Focus on Rapid, Nongenomic Effects. Pharmacol Rev. 2000, 52, 513–556.
(14) Losel, R. M.; Falkenstein, E.; Feuring, M.; Schultz, A.; Tillmann, H.-C.; Rossoll-Haserth, K.; Wehling, M. Nongenomic Steroid Action: Controversies, Questions, and Answers. Physiol. Rev. 2003, 83, 965–1016.
(15) Norman, A. W.; Mizwicki, M. T.; Norman, D. P. G. Steroid-Hormone Rapid Actions, Membrane Receptors and a Conformational Ensemble Model. Nat. Rev. Drug Discovery 2004, 3, 27–41.
(16) Levin, E. R.; Hammes, S. R. Nuclear Receptors Outside the Nucleus: Extranuclear Signalling by Steroid Receptors. Nat. Rev. Mol. Cell Biol. 2016, 17, 783–797.
(17) Alfsen, A. Biophysical Aspects of the Mechanism of Action of Steroid Hormones. Prog. Biophys. Mol. Biol. 1983, 42, 79–93.
(18) Bloch, K. E. Sterol Structure and Membrane Function. CRC Crit Rev. Biochem 1983, 14, 47–92.
(19) Ipsen, J. H.; Mourtis, O. G.; Bloom, M. Relationships Between Lipid Membrane Area, Hydrophobic Thickness, and Acyl-Chain Orientational Order. The Effects of Cholesterol. Biophys. J. 1990, 57, 405–412.
(20) McMullen, T. P.; McElhaney, R. N. Physical Studies of Cholesterol-Phospholipid Interactions. Curr. Opin. Colloid Interface Sci. 1996, 1, 83–90.
(21) Chiu, S. W.; Jakobsson, E.; Mashl, R. J.; Scott, H. L. Cholesterol-Induced Modifications in Lipid Bilayers: A Simulation Study. Biophys. J. 2002, 83, 1842–1853.
(22) Veatch, S. L.; Keller, S. L. Organization in Lipid Membranes Containing Cholesterol. Phys. Rev. Lett. 2002, 89, 268101.
(23) Ohvo-Reikälä, H.; Ramstedt, B.; Leppimäki, P.; Slote, J. P. Cholesterol Interactions with Phospholipids in Membranes. Prog. Lipid Res. 2002, 41, 66–97.
(24) Mourtis, O. G.; Zuckermann, M. J. What’s So Special About Cholesterol? Lipids 2004, 39, 1101–1113.
(25) de Meyer, F.; Smit, B. Effect of Cholesterol on the Structure of a Phospholipid Bilayer. Proc. Natl. Acad. Sci. U. S. A. 2009, 106, 3654–3658.
(26) Rög, T.; Pasenkiewicz-Gierula, M.; Vattulainen, I.; Karttunen, M. Order Effects of Cholesterol and Its Analogues. Biochim. Biophys. Acta, Biomembr. 2009, 1788, 97–121.
(27) Wennberg, C. L.; van der Spoel, D.; Hub, J. S. Large Influence of Cholesterol on Solute Partitioning into Lipid Membranes. J. Am. Chem. Soc. 2012, 134, 5351–5361.
(28) Bennett, W. F. D.; MacCallum, J. L.; Hinner, M. J.; Marrink, S. J.; Tieleman, D. P. Molecular View of Cholesterol Flip-Flop and Chemical Potential in Different Membrane Environments. J. Am. Chem. Soc. 2009, 131, 12714–12720.
(29) Jo, S.; Rui, H.; Lim, J. B.; Klauuda, J. B.; Im, W. Cholesterol Flip-Flop: Insights from Free Energy Simulation Studies. J. Phys. Chem. B 2010, 114, 13342–13348.
(30) Tipping, E.; Ketterer, B.; Christodoulides, L. Interactions of Small Molecules with Phospholipid Bilayers. Binding to Egg Phosphatidylcholine of Some Uncharged Molecules (2-Acetylamino-fluorene, 4-Dimethylaminoazobenzene, Oestrone and Testosterone) That Bind to Ligandin and Aminoazo-Dye-Binding Protein A. Biochem. J. 1979, 180, 319–326.
(31) Sato, B.; Huseby, R. A.; Matsumoto, K.; Samuels, L. T. Molecular Nature of Interaction of Steroids with Biomembranes Related to Androgen Biosynthesis. J. Steroid Biochem. 1979, 11, 1353–1359.
(32) Golden, G.; Rubin, R.; Mason, R. Steroid Hormones Partition to Distinct Sites in a Model Membrane Bilayer: Direct Demonstration by Small-Angle X-Ray Diffraction. Biochim. Biophys. Acta, Biomembr. 1998, 1368, 161–166.
(33) Mitragotri, S. In Situ Determination of Partition and Diffusion Coefficients in the Lipid Bilayers of Stratatum Corneum. Pharm. Res. 2000, 17, 1026–1029.
(34) Yamamoto, H.; Liljestrand, H. M. Partitioning of Selected Estrogenic Compounds Between Synthetic Membrane Vesicles and Water: Effects of Lipid Components. Environ. Sci. Technol. 2004, 38, 1139–1147.
(35) Kwon, J.-H.; Liljestrand, H. M.; Katz, L. E.; Yamamoto, H. Partitioning Thermodynamics of Selected Endocrine Disruptors Between Water and Synthetic Membrane Vesicles: Effects of Membrane Compositions. Environ. Sci. Technol. 2007, 41, 4011–4018.
(36) Estronca, L. M. B. B.; Moreno, M. J.; Vaz, W. L. C. Kinetics and Thermodynamics of the Association of Dehydroergosterol with Lipid Bilayer Membranes. Biophys. J. 2007, 93, 4244–4253.
(37) Lokajóvá, J.; Tišlová, H.; Vítila, T.; Riekkola, M.-L.; Wiedmer, S. K. Covalent Binding of Phospholipid Vesicles on Fused Silica Capillaries for Electrochromatography. Soft Matter 2011, 7, 6041–6050.
(38) Modi, S.; Anderson, B. D. Bilayer Composition, Temperature, Speciation Effects and the Role of Bilayer Chain Ordering on Partitioning of Dexamethasone and Its 21-Phosphate. Pharm. Res. 2013, 30, 3154–3169.
(39) Oren, I.; Fleishman, S. J.; Kessel, A.; Ben-Tal, N. Free Diffusion of Steroid Hormones Across Biomembranes: A Simplex Search with Implicit Solvent Model Calculations. Biophys. J. 2004, 87, 768–779.
(40) Vijayan, B.; Biggin, P. C. A Steroid in a Lipid Bilayer: Localization, Orientation, and Energetics. Biophys. J. 2008, 95, L45–L47.
(41) Orsi, M.; Essex, J. W. Permeability of Drugs and Hormones Through a Lipid Bilayer: Insights from Dual-Resolution Molecular Dynamics. Soft Matter 2010, 6, 3797–3808.
(42) Parisio, G.; Sperotto, M. M.; Ferrari, A. Flip-Flop of Steroids in Phospholipid Bilayers: Effects of the Chemical Structure on Transbilayer Diffusion. J. Am. Chem. Soc. 2012, 134, 12198–12208.
(43) Khelashvili, G.; Harries, D. How Sterol Tilt Regulates Properties and Organization of Lipid Membranes and Membrane Insertions. Chem. Phys. Lipids 2013, 169, 113–123.
(44) Hänggi, P.; Talkner, P.; Borkovec, M. Reaction-Rate Theory: Fifty Years After Kramers. Rev. Mod. Phys. 1990, 62, 251.
(45) Fugler, L.; Clejan, S.; Bittman, R. Movement of Cholesterol Between Vesicles Prepared with Different Phospholipids or Sizes. J. Biol. Chem. 1985, 260, 4098–4102.
(46) Grossman, C.; Ruhs, S.; Langenbruch, L.; Mildenberger, S.; Sträts, N.; Schumann, K.; Gekle, M. Nuclear Shuttling Precedes Dimerization in Mineralocorticoid Receptor Signaling. Chem. Biol. 2012, 19, 742–751.
(47) Harvey, B. J.; Higgins, M. Nongenomic Effects of Aldosterone on Ca2+ in M-1 Cortical Collecting Duct Cells. Kidney Int. 2000, 57, 1395–1403.
(48) Haseroth, K.; Gerdes, D.; Berger, S.; Feuring, M.; Günther, A.; Herbst, C.; Christ, M.; Wehling, M. Rapid Nongenomic Effects of Aldosterone in Mineralocorticoid-Receptor-Knockout Mice. Biochem. Biophys. Res. Commun. 1999, 266, 257–261.
(49) Vallée, M.; et al. Pregnenolone Can Protect the Brain from Cannabis Intoxication. Science 2014, 343, 94–98.
(50) Cheng, W. W. L.; Chen, Z.-W.; Bracamontes, J. R.; Budelier, M. M.; Krishnan, K.; Shin, D. J.; Wang, C.; Jiang, X.; Covey, D. F.; Akk, G.; Evers, A. S. Mapping Two Neurosteroid-Modulatory Sites in the Prototypic Pentameric Ligand-Gated Ion Channel GLIC. J. Biol. Chem. 2018, 293, 3013–3027.
(51) Laverty, D.; Thomas, P.; Field, M.; Andersen, O. J.; Gold, M. G.; Biggin, P. C.; Gielen, M.; Smart, T. G. Crystal Structures of a GABA<sub>A</sub> Receptor Chimera Reveal New Endogenous Neurosteroid-Binding Sites. Nat. Struct. Mol. Biol. 2017, 24, 977–985.
(52) Miller, P. S.; Scott, S.; Masulis, S.; De Colibus, L.; Pardon, E.; Steyaert, J.; Aricescu, A. R. Structural Basis for GABA<sub>A</sub> Receptor Potentiation by Neurosteroids. Nat. Struct. Mol. Biol. 2017, 24, 986–992.
(53) Baier, C. J.; Fantini, J.; Barrantes, F. J. Disclosure of Cholesterol Recognition Motifs in Transmembrane Domains of the Human Nicotinic Acetylcholine Receptor. Sci. Rep. 2011, 1, 69.
(54) Kostakis, E.; Jang, M.-K.; Russe, S. J.; Gibbs, T. T.; Farb, D. H. A Steroid Modulatory Domain in NR2a Collaborates with NR1 Exon 5 to Control NMDAR Modulation by Pregnenolone Sulfate and Protons. J. Neurochem. 2011, 119, 486–496.
(55) Borovska, J.; Vytkicky, V.; Stastna, E.; Kapras, V.; Slavikova, B.; Horak, M.; Chodounska, H.; Vytkicky, L. Access of Inhibitory Neurosteroids to the NMDA Receptor. Br. J. Pharmacol. 2012, 166, 1069–1083.
(56) Heerklotz, H.; Keller, S. How Membrane Partitioning Modulates Receptor Activation: Parallel Versus Serial Effects of Hydrophobic Ligands. Biophys. J. 2013, 105, 2607–2610.
(57) Wagner, T. F. J.; Loch, S.; Lambert, S.; Straub, I.; Mannebach, S.; Mathar, I.; Düser, M.; Lis, A.; Flockerzi, V.; Philipp, S. E.; Oberwinkler, J. Transient Receptor Potential M3 Channels Are Ionotropic Steroid Receptors in Pancreatic Beta Cells. Nat. Cell Biol. 2008, 10, 1421–1430.