Describing the Mechanism of Antimicrobial Peptide Action with the Interfacial Activity Model

William C. Wimley*
Department of Biochemistry SL43, Tulane University Health Sciences Center, New Orleans, Louisiana 70112-2699

The field of antimicrobial peptide research originated in the early 1980s when Hans Boman reported (1) that the humoral immune system of silk moths (Hyalophora cecropia) contained peptides (cecropins) with potent and broad-spectrum (active against multiple distinct classes of microbes, e.g., Gram-positive bacteria, Gram-negative bacteria, and fungi) antimicrobial activity. Shortly thereafter, other AMPs were identified. For example, Shunji Natori described sarcotoxins from fly larvae (2), Robert Lehrer described defensins from mammalian macrophages (3), and Michael Zasloff described the magainins found in the skin of the frog Xenopus laevis (4). Although the primary and secondary structures of these first antimicrobial peptides are very different, the early reports described shared characteristics of biological activity and biophysical behavior that are still appropriate today to identify and categorize antimicrobial peptides; AMPs are cationic, amphipathic peptides that have broad spectrum microbicidal activity that is associated with membrane permeabilization. Unlike nonspecific membrane lytic peptide toxins, such as melittin, from the venom of the honey bee (Apis mellifera), or alamethicin from the fungus Trichoderma viride, AMPs have little cytolytic or cytotoxic activity against host cells.

Since those influential early publications, the field has undergone incredible growth, with nearly 1000 known examples of antimicrobial peptides (5–7). Naturally occurring examples are found in all classes of organisms: vertebrate animals including humans, invertebrate animals, plants, and microbes. Descriptions of newly discovered AMPs are common occurrences in the literature. In addition to the naturally occurring AMPs, peptides with the same biological activity have been de-

ABSTRACT Antimicrobial peptides (AMPs) have been studied for three decades, and yet a molecular understanding of their mechanism of action is still lacking. Here we summarize current knowledge for both synthetic vesicle experiments and microbe experiments, with a focus on comparisons between the two. Microbial experiments are done at peptide to lipid ratios that are at least 4 orders of magnitude higher than vesicle-based experiments. To close the gap between the two concentration regimes, we propose an “interfacial activity model”, which is based on an experimentally testable molecular image of AMP–membrane interactions. The interfacial activity model may be useful in driving engineering and design of novel AMPs.

*Corresponding author, wwimley@tulane.edu.

Received for review May 30, 2010 and accepted August 10, 2010.

10.1021/cb1001558
© XXXX American Chemical Society
signed de novo (8), engineered from natural sequences (9, 10), or selected from combinatorial libraries (11, 12).

Much of the vast literature on antimicrobial peptides is devoted to biophysical characterization and structure–function studies in model systems such as lipid vesicles or detergents. Yet despite the large volume of data available, compelling structure–function relationships are very rare in antimicrobial peptide research. In fact, recent literature suggests that antimicrobial activity is not dependent on specific amino acid sequences or on specific peptide structures (e.g., the defensins) having disulfide cross-linked tertiary structure. Some AMPs are cyclic (18). Some small AMPs (< ca. 15 residues) are often poorly or irregularly structured (19). In terms of structure, AMPs are about as diverse as possible (Figure 1). However, one of the few conserved characteristics of antimicrobial peptides is their cationic and hydrophobic composition. Most AMPs are cationic overall with some examples having net charges as high as +9. From databases of AMP sequences (6, 7), we observe that the basic amino acids, Arg and Lys, are roughly 50% more abundant in AMPs than in genomes overall, while the acidic amino acids, Glu and Asp, are ~75% less abundant than expected. The hydrophobic amino acids are also more abundant in AMPs, while polar residues (especially Gln) are less abundant than expected. Smaller AMPs often have higher abundances of aromatic and basic amino acids. Consistent with these observations, AMPs have been successfully engineered using reduced amino acid alphabets (17) that contain only basic and hydrophobic amino acids. Some simple cationic/hydrophobic peptides with good antimicrobial activity include RWn (n = 1–5) (20) and WALRLYLV (11). Even some cationic/hydrophobic acylated tetra-peptides (21) or modified dipeptides (20) have potent antimicrobial activity.

The mixed cationic and hydrophobic composition of AMPs makes them well suited for interacting with and perturbing microbial cytoplasmic membranes that typically present anionic surfaces, rich in lipids such as...
phosphatidylglycerol or cardiolipin, to the outside environment. One of the most commonly cited explanations for the selectivity of AMPs for microbes over host cells is the difference in membrane interactions due to differences in exposed anionic lipid content. Because of the contribution of electrostatic interactions, binding of AMPs to microbial membranes is significant, while binding of AMPs to the neutral phosphatidylcholine/cholesterol/sphingomyelin-rich surfaces of animal plasma membranes is weaker. There is ample direct evidence that most AMPs permeabilize microbial cytoplasmic membranes and that the membranes are often permeabilized with increasing severity with time (22). AMPs can dissipate the electrochemical gradient across microbial plasma membranes, within a few seconds of addition (11, 22). Thus, AMPs must be able to rapidly pass through the thick proteoglycan layer of Gram-positive bacteria and the outer membrane lipopolysaccharide (LPS) layer of Gram-negative bacteria. Permeation of larger markers, including dye markers, metabolites, and cytosolic proteins, through the cytoplasmic membrane occurs on the time scale of minutes to tens of minutes (11, 13). After an hour or more in contact with AMPs, gross disruption of microbial membrane structure and morphology is often noted, including membrane blebbing, vesiculation, fragmentation, release of DNA, cell aggregation, and destruction of cell morphology.

The Membrane Is Not the Only Site of AMP Action. While most AMPs interact with and influence the integrity of microbial membranes, it is not known if membrane permeabilization is always the lethal event or if the membrane is the only site of action. There are AMPs that may have alternate modes of action. For example the bovine neutrophil defensin indolicidin, a 13-mer rich in basic and aromatic residues, has typical low micromolar antibacterial activity and the degree of membrane permeabilization is often associated with AMP activity, it is apparently not always required for activity.

It is not known whether small amounts of membrane permeabilization alone can account for lethality of AMPs. Those peptides that do not extensively permeabilize microbe membranes may be having effects on microbe viability that depend on interactions with intracellular components, perhaps in addition to effects on the membrane. Given the similarity of physical chemistry between AMPs and some cell-penetrating peptides (26–28), it should not be surprising that AMPs can also translocate across microbial membranes and can sometimes do so without extensive permeabilization. Although translocation of peptides into microbes has not often been studied carefully, there are reports that peptide translocation is coupled to leakage. For example, the human defensin cryptdin-4 and the frog AMP magainin 2 both translocate across bilayers with a half-time of about 10 min (24, 29). It seems likely that many AMPs will translocate across microbial membranes. Axelsen reported changes in gene transcription in multiple genes in microorganisms treated with subpermeabilizing concentrations of cecropin A (30) and concluded that systemic metabolic effects were taking place prior to membrane permeabilization. This work suggested that translocation is occurring at concentrations that do not induce permeabilization. Other intracellular targets of AMPs have been proposed, including DNA and chaperonins.

Almeida and Pokorny have suggested that it may be appropriate to recognize a very broad overlap between membrane permeabilization and membrane translocation activity (31, 32). They have described a quantitative, predictive model based on hydrophobicity scales (31). It is likely that most AMPs will also be cell-penetrating peptides when interacting with living microbes and will have access to the cytoplasm. Multiple simultaneous mechanisms of AMP action (e.g., membrane permeabilization as well as intracellular effects) may help explain their broad-spectrum activity and the rarity of inducible resistance.

Model Membrane Studies. There are countless studies showing that AMPs interact with and perturb lipid bilayer membranes in vitro. Within these many experiments, lipid compositions, buffer conditions, and peptide and lipid concentrations vary widely. Even the phase state of the lipids and the degree of bilayer hydration vary between experiments. Combined with the inherently variable nature of AMP activity, these factors confuse the interpretation and comparison of results.
While lipid compositions vary widely, mixtures of anionic and zwitterionic lipids are often used to mimic microbial membranes. For example, some laboratories use mixtures of phosphatidylglycerol and phosphatidylethanolamine to specifically mimic the E. coli inner membrane (33), while others use pure phosphatidylglycerol to specifically mimic Gram-positive bacteria (34). Other commonly used systems include phosphatidylcholine mixed with phosphatidylglycerol, phosphatidylserine, or cardiolipin to broadly mimic the anionic surface of a microbe (12, 17). Experiments have also been performed with lipids extracted from E. coli or other organisms (35). To mimic exposed mammalian or host membranes, zwitterionic phosphatidylcholine (PC) or PC-cholesterol membranes are often used (8). Because of the variety of protocols and experimental conditions used, a single most appropriate lipid composition has never emerged from the literature.

In model membranes and living cells, lipid composition will affect peptide binding as well as the inherent susceptibility of the bilayer to permeabilization. Other than electrostatic effects, there are few, if any, examples of specific binding of an AMP to a particular lipid species. Cationic/hydrophobic peptides with good interfacial activity are expected to perturb any fluid phase lipid bilayer membrane to which they bind well enough, although bilayers with some lipid compositions may be inherently more or less stable than others. In practical terms AMPs that are less hydrophobic and more cationic will require anionic lipids for binding. For example, human defensins, which are highly cationic, must be studied in bilayers with anionic lipids, or else they are inactive due to poor binding (36). The more hydrophobic AMPs, such as the magainins, are active in zwitterionic lipid bilayers as well as anionic bilayers. It is likely that any model system that comprises a fully hydrated, fluid phase bilayer to which one can achieve reasonable bound peptide concentration is appropriate as a model system. Recently we showed that large unilamellar vesicles (11, 12, 25) composed of 90% phosphatidylycholine and 10% phosphatidylglycerol in 50 mM sodium phosphate buffer could be used to specifically select non-hemolytic, broad-spectrum antimicrobial peptides from combinatorial libraries, suggesting that this may be a good consensus membrane composition to study AMPs.

Upon binding of AMPs to anionic lipid vesicles through electrostatic and hydrophobic interactions, many types of perturbations can occur, including membrane permeabilization, but AMPs can also drive vesicle aggregation, vesicle fusion, formation of nonbilayer phases, lipid phase separation, transbilayer movement (flip-flop) of lipids, and complete solubilization of membranes. Some of these effects may not relate to antimicrobial activity and should be considered experimental artifacts. Although not often reported in the literature, vesicle aggregation and fusion are very common occurrences when cationic peptides are added to anionic vesicles (13), as evidenced by rapid increases in turbidity upon peptide addition. It is likely that some (or many) reports of peptide-induced “leakage” from anionic vesicles observed at very high peptide to lipid (P:L) ratios (such as P:L = 1:10) include leakage that is due to leaky fusion/aggregation events in addition to (or instead of) direct membrane permeabilization. Few attempts have been made to distinguish leakage that is purely a consequence of fusion from legitimate leakage that occurs without vesicle fusion. In a recent paper (13) we examined AMP-induced fusion and leakage independently and found that leakage and vesicle fusion both occurred at P:L > 1:50 while only leakage occurred at lower P:L (up to 1:500). We concluded that leakage measured at P:L = 1:50 may have included a contribution from fusion as well as membrane permeabilization.

As shown by the example data in Figure 2, most antimicrobial peptides are active against vesicles when bound peptide to lipid ratios are from about 1:500 to 1:50, or about 200—2000 peptides per vesicle. In contrast, ideal pore-forming peptides can permeabilize lipid vesicle with as few as 10 peptides bound per vesicle, or 1 peptide per 10,000 lipids (37). We speculate that AMPs are not more potent because the very potent peptides lack selectivity. On the other end of the spectrum, any membrane-interacting molecule can disrupt membranes at very high concentrations. For this reason, it is likely that some published results for putative AMPs permeabilizing vesicles are essentially artifacts that arise from extremely high peptide:lipid ratios. The literature

**KEYWORDS**

All-or-none release: Release of entrapped markers from vesicles during which a portion of the vesicles release all of their contents and the remainder of the vesicles release none.

Antimicrobial peptide (AMP): A peptide of less than 50 residues that has microbicidal activity at micromolar concentrations.

Barrel-stave pore: A membrane-spanning pore formed by peptides that interact laterally to form a pore lined by peptides.

Broad-spectrum activity: Microbicidal activity against multiple classes of microbes.

Carpet model: A commonly cited descriptive model of antimicrobial peptide action that is used to describe the fact that antimicrobial peptides only permeabilize membranes when the membranes are carpeted with peptide.
contains many experiments which show significant leakage of vesicle entrapped contents at P:L of 1:10, 1:1, or even 10:1. Any in vitro membrane permeabilization measurements that show activity only at more than 1 peptide bound per 50 lipids (P:L = 1:50) should be viewed with caution.

The Prevailing Transmembrane Pore Models. The simplest models of membrane permeation by peptides involve the formation of transbilayer pores or channels through the membrane as shown by the models in Figure 3. In a barrel-stave pore, peptides interact laterally with one another to form a specific structure that is reminiscent of a membrane protein ion channel. In the toroidal pore model, specific peptide–peptide interactions are not present. Instead, peptides affect the local curvature of the bilayer in a cooperative manner such that a toroid of high curvature forms. These two structures are fundamentally different. Barrel stave pores work with the bilayer hydrocarbon core, using it as a template for peptide self-assembly, whereas toroidal pores work against the hydrocarbon core, disrupting the normal segregation of polar and nonpolar parts of the membrane by providing alternate surfaces for lipid hydrocarbon and headgroups to interact favorably with.

A classical example of a lytic peptide toxin that almost certainly forms transmembrane pores is alamethicin, a 20-residue fungal peptide that folds into an amphipathic α-helix. Alamethicin can exist, depending on hydration and concentration, either parallel or perpendicular to the lipid bilayer normal (38). The perpendicular structure (39, 40) is consistent with transmembrane barrel stave or toroidal pore. Huang has given evidence that the pore is a barrel-stave pore (39); however, we have shown that alamethicin catalyzes very rapid transbilayer equili-
brane pores is scarce or lacking entirely, as we discuss microbial peptides, evidence for water-filled, transmembrane pores for alamethicin. The lytic peptide toxin melittin, from honey bee venom, is another classical example of a pore-forming peptide that forms a membrane-spanning pore, but in the realm of membrane-permeabilizing peptides, these classical “examples” are actually exceptions. In any case, alamethicin and melittin are not really antimicrobial peptides, which are defined by their host defense function, but instead are nonspecific, membrane-permeabilizing peptide toxins. For the vast majority of so-called “pore-forming” antimicrobial peptides, evidence for water-filled, transmembrane pores is scarce or lacking entirely, as we discuss below.

The Prevailing Non-pore Models. In addition to the pore models described above, AMP activity has also been described using some common nonpore models that have been proposed to explain or categorize the mechanism of action of AMPs. The so-called “carpet model” is the most commonly cited phenomenological model and was proposed in 1996 by Shai (42) to explain the mechanism of action of mammalian cecropin P1 on model membranes. Cecropin P1 is always oriented parallel to the membrane surface and is active only at high P:L ratios. Shai and co-workers concluded that the peptide is active only when it forms a “carpet” on the bilayer surface. The “detergent model!” is also often cited to explain the catastrophic collapse of membrane integrity and probe-size-independent leakage observed with some AMPs at high peptide concentration (35).

While these catchphrases and others have been useful toward providing a common lexicon to a diverse field, they lack a specific molecular or physical-chemical basis for membrane-permeabilizing activity. Although these models may have broadly inspired successful design efforts (16), they have rarely been used to make specific molecular predictions or to drive engineering and design. In contrast some recent discussions have laid a physical chemical foundation that should be much more useful. For example, Bechinger and Lohner (43, 44) recently presented a model based on molecular shape to describe antimicrobial peptide activity. They suggested that the physical chemistry of AMPs and membranes could be described with phase diagrams, as one would describe mixtures of lipids. Recently, Epand and colleagues (45, 46) have proposed that lateral lipid phase separation, or lipid clustering, induced by cationic antimicrobial peptides explains their biophysical properties as well as their selectivity and biological activity. They have provided experimental evidence to support this idea suggesting that leakage could occur at phase boundary defects (45). Almeida and colleagues have described AMP activity in terms of binding, insertion, and perturbation (32, 47–49) using kinetic and thermodynamic models. Such quantitative descriptions of AMP activity, once they are parametrized to correlate peptide sequence or composition to activity, could allow for predictions of activity to be made and thus could be useful for engineering. We suggest here that the physical-chemical concept of “interfacial activity”, perhaps in conjunction with these other concepts, will be useful to explain, predict, and engineer the activity of AMPs.

AMP May Not Form Transmembrane Pores. Despite innumerable studies of AMP activity in cells, vesicles, and other synthetic bilayer systems, compelling evidence for specific, transmembrane pores or channels in vesicle bilayers, even transient pores, is rare. While some structural studies have indirectly supported the idea of discrete pores (50–52), vesicle-based leakage experiments are mostly inconsistent with this idea. AMPs bind to vesicles rapidly (<30 s) (17) and remain bound at equilibrium. Because large unilamellar vesicles (LUV) contain roughly 100,000 lipids each, a typical experiment (P:L of 1:100) there are about 1000 peptides bound to each vesicle, capable of forming a hundred or more small pores. To examine what would happen to a vesicle’s contents if a pore were present, we have performed numerical simulations of uncharged small molecule release from single vesicles based on random walk diffusion through single pores. These simulations predict the release kinetics of entrapped small probes from a large unilamellar vesicle (53) through a single water-filled channel of 10 Å diameter (Figure 4). The calculated release rate of 50,000 ions per second per pore is consistent with the known ability of protein ion channels to pass as many as 107 ions per second (54). Because the interior of a “large” unilamellar vesicle is only about 10−19 L, there are only a few hundred to a few thousand probe molecules inside each vesicle. The simulation results, in Figure 4, show that a single aqueous channel through a bilayer will release all the contents of a vesicle in a few tenths of a second, at most. A vesicle with 1000 bound peptides (P:L = 1:100)
could conceivably contain more than 100 pores. In reality, vesicles with 1000 bound antimicrobial peptides typically release only a portion of their contents over tens of minutes, with maximum initial rates rates from 0.1 to 100 ions per second per vesicle. Commonly, initial rates are 1–10 ions per second per vesicle (i.e., per 1000 peptides). The observed release rates are thus at least 3 orders of magnitude slower than predicted for even a single aqueous channel. Furthermore, leakage from vesicles caused by AMPs is often incomplete, suggesting that true pores do not form, even transiently. This analysis suggests that most AMPs do not ever form true pores in large unilamellar vesicles and leakage occurs by a general disruption of membrane integrity.

**Giant Unilamellar Vesicle Studies.** Studies of AMP activity in giant unilamellar vesicles (GUV) are revealing because they allow individual vesicles and individual events to be monitored. Not surprisingly, different AMPs studied in GUVs in different laboratories behave quite variably. In some cases, individual vesicles accumulate AMPs on their surface until a catastrophic burst event occurs (55). Bursting occurs only above a threshold peptide concentration and is a stochastic event in a collection of vesicles. This behavior is consistent with the all-or-none release behavior observed in LUVs (described above). In other GUV experiments, vesicles remain intact after AMP addition but steadily lose entrapped contents over the course of a few hundred seconds (56, 57). Toroidal or barrel stave pores have been invoked to explain this type of observation. Given the 100-fold larger size of a GUV relative to an LUV, leakage through a *single pore* is expected to occur in the observed time range (Figure 4), but a single GUV actually has enough bound peptide to have *millions* of peptide pores. In a typical GUV experiment, a small concentration of GUVs (very low lipid concentration, see Figure 5) are exposed to 1–10 μM peptide. The bound peptide to lipid ratio will be high, minimally P:L = 1:50 for a peptide that binds moderately well (see Figure 5). A 10 μM GUV will contain about 10^6 lipids (compared to 10^5 for an LUV) meaning that a GUV will have roughly 2 × 10^7 peptides bound to it in a typical experiment. That is
enough peptide bound to a single giant vesicle to form about two million pores, assuming decameric pores. The fact that release of entrapped contents is very slow even when there are twenty million peptides bound to each vesicle indicates that transmembrane pore formation, if it occurs at all, is a very rare event. It is much more likely that there is a low flux of markers through the membrane caused by the global disruption of the lipid packing caused by the peptides. Tamba and Yamazaki called this a “two-state transition” model of external binding followed by critical destabilization of the bilayer. This is the same idea used to explain “carpet model” leakage from LUVs.

The Enigma of Partial Transient Release. Studies of vesicle permeabilization by AMPs (e.g., Figure 4, panel C) frequently show an enigmatic and still unexplained behavior: upon addition of peptide to a homogeneous population of vesicles (or vesicles to peptide), a rapid burst of leakage occurs which then stops with a halftime of just a few minutes. Leakage often ceases before complete loss of entrapped contents has taken place. A second addition of peptide will cause a second burst of leakage which also stops (W.C.W. unpublished observations). Increasing the peptide concentration will increase the fractional loss of contents. Perhaps even more enigmatic is the observation that partial transient release is sometimes graded (13), in which all of the vesicles release a fraction of their contents, and is sometimes all-or-none (17), in which a fraction of the vesicles release all of their contents and the remainder release none of their contents. Furthermore, AMP-induced leakage of probes is sometimes probe-size-dependent and sometimes probe-size-independent. For many AMPs the rate of leakage after the rapid initial burst is very slow, sometimes immeasurably slow. For example, we have shown that some antimicrobial peptides (13, 17) release almost no measurable amount of entrapped markers for as long as 24 h after the initial burst, despite the fact that hundreds of peptides remain bound to each vesicle and that the peptides have the same secondary structure as during the initial leakage phase. See Figure 4, panel C for an example of actual release data. Published leakage curves for many AMPs are very similar to these curves. In sharp contrast, true pore-forming peptides, such as alamethicin, cause rapid and continual leakage of vesicle contents that does not stop until complete release has occurred. As shown diagrammatically in Figure 2, leakage caused by true pore-forming peptides occurs at P:L ratios as low as 1:10000 (10 peptide/vesicle), whereas most AMPs have good activity only in the range of P:L = 1:100 (1000 peptides per vesicle) or higher.

Transient Failure. Partial transient release shows that vesicle permeabilization by most AMPs must be driven by nonequilibrium events. It has been suggested that it is driven by the disequilibrium state that immediately follows the addition of peptide to the vesicles in which only the outer monolayer of the vesicle has peptide bound to it. The resulting imbalance of charge, area, and surface tension may drive the transient, sometimes catastrophic, failure of the bilayer structure and concomitant leakage of entrapped contents. Depending on the magnitude and lifetime of the failure, it is possible to imagine either graded or all-or-none release, and it is possible to imagine size-dependent or size-independent release.

In any case, release driven by AMPs over a wide range of peptide concentration usually stops or slows significantly before all vesicle contents have been released. This probably means that once transbilayer equilibrium of peptides has been reached, the driving force for large-scale bilayer destabilization is decreased. Recently, we reported a family of AMPs that causes partial, all-or-none release of vesicle contents (17). Assuming that all peptides ultimately reach transbilayer equilibrium, this observation suggests that in addition to the stochastic, transient event that relieves transbilayer asymmetry of peptides and concomitantly causes complete release of entrapped contents, there can also be a "silent" non-permeabilizing pathway to equilibrium. At least for this family of peptides, a fraction of the vesicles reach equilibrium by a catastrophic event, losing all their contents, while the remaining vesicles reach equilibrium by a process that involves no leakage whatsoever. The silent pathway to equilibrium is presumably peptide translocation across the bilayer. At higher peptide concentration, the fraction of vesicles releasing all their contents increases, consistent with an increase in probability of catastrophic bilayer permeabilization.

A Comparison of Biophysical and Biological Experiments. Although the biological activity of AMPs and their vesicle permeabilizing activity are correlated, it is not known how closely the mechanisms of action overlap in the two systems. For example, there are no data to suggest that transient release phenomena occur in microbial permeabilization. The overlap in mech-
anism is a critical question in engineering and design studies, because these are almost always based on vesicle permeabilization assays. To begin comparing mechanisms of action we have contrasted a typical in vitro experiment using synthetic lipid bilayers with a typical biological experiment measuring on microbe sterilization. Experimentally verified stoichiometry information from the two classes of experiments is shown in Table 1. In both experiments it is common to use peptide concentrations in the low micromolar range. Vesicle experiments are typically carried out at 0.1–1 mM lipid, which gives system P:L ratios of roughly 1:100. Biological assays are carried out at 10^4–10^6 bacterial cells mL\(^{-1}\) (58). Assuming that an \(E.\ coli\) cell is tubular and is 1 \(\mu\)m wide and 2 \(\mu\)m long and taking into account the area of a lipid molecule (~0.7 nm\(^2\)) (59) and the fact that there are two membranes, we calculate roughly 2.5 \(\times\) 10^7 lipids in the bacterial cell. Thus, at 10^7 cells mL\(^{-1}\) the total lipid concentration is about 5 nM. This is 4–5 orders of magnitude lower lipid concentration than in a typical vesicle-based experiment. The total (system) peptide to lipid ratio in a microbe sterilization assay there are roughly 10–100 peptides bound for each bacterial lipid molecule (Table 1) amounting to as many as 10^9 peptides bound to each cell. Importantly, such high concentrations of bound peptide are required for activity; a 10-fold decrease in peptide concentration (e.g., from 5 to 0.5 \(\mu\)M peptide) eliminates antimicrobial activity for most AMPs despite the fact that there are still huge numbers of peptides bound to each cell. Of course, it is not physically possible to have 100 peptides bound per lipid in a membrane; peptide must be binding in large amounts to other cellular components. AMPs are known to bind lipopolysaccharide, cell wall components, and DNA (61, 62). Whatever the actual membrane binding, it is reasonable to assume the microbial membranes are saturated with peptide and that there is a large reservoir of nonmembrane peptide bound to each cell. With as many as 10^9 peptides bound to each cell, AMPs will actually outnumber all proteins, ATP, and most electrolytes and metabolites. (For example, there are probably several million ATP molecules per bacterial cell). The practical consequence of these numbers is that microbe killing can result from leakage

<table>
<thead>
<tr>
<th>Property</th>
<th>Large unilamellar vesicle</th>
<th>(E.\ coli) sterilization assay</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dimension</td>
<td>0.1 (\mu)m diameter</td>
<td>1 (\mu)m (\times) 2 (\mu)m diameter</td>
</tr>
<tr>
<td>Aqueous volume</td>
<td>(10^{-19}) L</td>
<td>(10^{-15}) L</td>
</tr>
<tr>
<td>Number of lipids</td>
<td>100,000 per vesicle</td>
<td>25,000,000 per cell</td>
</tr>
<tr>
<td>Typical concentrations</td>
<td>500 (\mu)M lipid</td>
<td>(10^6) cells mL(^{-1})</td>
</tr>
<tr>
<td></td>
<td>(1 \times 10^{12}) vesicles mL(^{-1})</td>
<td>0.004 (\mu)M lipid</td>
</tr>
<tr>
<td></td>
<td>5 (\mu)M peptide</td>
<td>5 (\mu)M peptide</td>
</tr>
<tr>
<td>Relative peptide</td>
<td>(1 \times 10^7) peptides per vesicle</td>
<td>(1 \times 10^9) peptides per cell</td>
</tr>
<tr>
<td>Total peptide:lipid</td>
<td>1 peptide:100 lipids</td>
<td>1000 peptide:1 lipid</td>
</tr>
<tr>
<td>Bound peptide:lipid</td>
<td>1 peptide:200 lipids</td>
<td>100 peptides:1 lipid</td>
</tr>
</tbody>
</table>

*Bound peptide to lipid ratios are conservative estimates based on direct, experimentally measured binding (see text).
that is very inefficient. For example, one ATP released for every 100 bound peptides is enough to deplete ATP completely. Given the conditions described in Table 1, it should not be surprising that bacteria do not frequently develop inducible resistance to AMPs, and when they do it involves changes in gross membrane architecture (63). Most importantly, these numbers show that the formation of explicit transmembrane pores is not necessary to explain the biological activity of AMPs.

For comparison, a similar calculation can be done for a typical erythrocyte hemolysis assay (e.g., 50 μM peptide added to 5 x 10⁸ red blood cells (8)). This calculation gives a system P:L ratio of about 1:1. Because binding of AMPs to erythrocytes is weak, the bound P:L ratio may be 1:20 or less. Thus, in terms of overall peptide to lipid ratio, a typical hemolysis assay is performed under conditions that are comparable to lipid vesicle leakage assays. While the overall peptide to lipid ratio in a hemolysis assay is quite dissimilar to the conditions of a microbe sterilization assay, it is likely that the membranes in both assays are close to saturation and thus may be comparable to one another.

The Interfacial Activity Model. The hydrocarbon core of an unperturbed lipid bilayer membrane is one of the most hydrophobic microenvironments found in nature, with physical-chemical properties that are very similar to a liquid alkane phase (64). Accordingly, the hydrocarbon core (25–30 Å thick) normally imparts a strict barrier to the permeation of polar or charged solutes through the bilayer. Yet, it is positioned between the two bilayer interfacial zones, called “zones of tumultuous chemical heterogeneity” by Wiener and White (64). Each interfacial zone is 10–15 Å thick and is composed of the lipid polar groups, water, and solution counterions, as well as small amounts of the hydrocarbon groups.

The property that we define as “interfacial activity” is the ability of a peptide to perturb the permeability barrier imposed by the hydrocarbon core by partitioning into the interfacial region of the bilayer and driving local rearrangements in vertical lipid packing (i.e., normal to the bilayer plane) that alter the segregation between the hydrocarbon core and the interfacial groups. This activity is driven by peptides that bind to membranes and are amphipathic, but with imperfect segregation of polar and nonpolar groups. Unlike a very hydrophobic peptide or one with ideal amphipathicity, when an interfacially active peptide is bound to a bilayer, the bilayer must be deformed and the hydrocarbon disrupted (i.e., intermingled with polar lipid headgroup moieties) to simultaneously accommodate the nonpolar and polar/charged groups of the peptide (11). Examination of the structures and sequences of AMPs (Figure 1) reveals that most of them have a hydrophobic segment or patch, but one that is not nearly large enough to span a bilayer and is interrupted by or bounded by at least one polar residue, frequently arginine or lysine. Such “imperfect amphipathicity” is necessary for interfacial activity. Much of the available data suggests that the exact structure or spatial arrangement of the hydrophobic and polar groups is not as important as the physical chemical balance between the two types of interactions. When such a peptide binds to a bilayer, the hydropho-
bic portion, often dominated by aromatic residues, drives a deep partitioning into the interfacial zones, but the nearby polar residues promote the incursion of lipid polar groups deeper in the membrane, along with the polar residues of the peptide. We suggest that the signature of an interfacially active peptide in a lipid bilayer experiment will be bilayer translocation of peptide and lipid, even at low peptide concentration. At higher peptide concentrations, the nonequilibrium distribution causes a transient, cooperative transbilayer movement of peptide, lipids, and polar solutes, which defines the leakage activity of the peptide.

In Figure 6, we show the results of molecular dynamics simulations of a peptide “pore” conducted by Sen-gupta and Marrink (65). These simulations capture the essence of the interfacial activity model in several aspects. The presence of peptides, which are mostly partitioned into the interfacial zone, dramatically perturbs the normally strict segregation of polar and nonpolar moieties across the bilayer. Compare, for example, the distribution of water and lipid polar groups in the vicinity of the peptides (bilayer middle) and away from the peptides (bilayer left/right). The leakage of polar solutes across the bilayer occurs because they are carried across the bilayer along with peptide and lipid molecules when the perturbed bilayer structure is formed. The most significant difference between the simulation results and the interfacial activity model is that some interfacially active antimicrobial peptides can act without self-assembly, as shown experimentally by Almeida (49).

When one considers interfacial activity in the context of a peptide-saturated bacterial membrane, the details will be different from a lipid vesicle experiment, but the interactions are driven by the same basic physical chemistry. This is the reason there is a strong correlation between the two experiments but not an exact overlap in mechanism. We have proposed here that the interfacial activity of imperfectly amphipathic peptides, as measured in lipid vesicle experiments, is a reasonable surrogate measurement for the likelihood that a peptide will have broad-spectrum antimicrobial activity.

**Experimentally Testable Predictions.** The interfacial activity model can be expressed in terms of experimentally testable predictions:

- Interfacially active peptides partition into the bilayer interface and locally perturb polar-nonpolar segregation of lipid moieties.
- Interfacial activity requires imperfect amphipathicity and the proper balance of hydrophobic and polar amino acids.
- Interfacial activity will depend more on amino acid composition than peptide structure or sequence.
- Interfacial activity does not require peptide self-assembly.
- The permeation pathway is mostly made of lipid and peptide, not an aqueous channel.
- Only a few molecules are transported across the membrane per peptide.
- At similar bound peptide to lipid ratios, an interfacially active peptide will behave similarly in all fluid phase membranes.
- At low bound peptide to lipid ratios AMPs will translocate across membranes.
- Lipid translocation, peptide translocation, and membrane leakage will always be coupled.
- Leakage from vesicles will depend on the rate of peptide addition.
- Nonpeptide molecules (peptide mimetics or other polymers) that are imperfectly amphipathic can have AMP-like activity.

**Conclusions.** In terms of detailed molecular mechanisms, multiple overlapping mechanisms of AMP activity probably exist and it may not be possible to define a single unifying description. What we have done here is to define the physical-chemical commonalities that seem to be important in antimicrobial peptide activity. While there is overlap between the interfacial activity model and others in the literature, we believe it is beneficial to begin to think about the problem of antimicrobial peptide mechanism, engineering, and design using an experimentally testable, mechanistic, and semimolecular model of AMP action.

**Acknowledgment:** For many interesting antimicrobial peptide discussions over the course of many years, I thank Kalina Hristova, Paulo Almeida, Antje Pokorny, Mikhail Merzliakov, Yechiel Shai, Jack Blazyk, Bill Walkenhorst, Josh Rausch, Chris Bishop, Ramesh Rathinakumar, Aram Krauson, Jessica Marks, Thomas Freeman, Drew Hoffmann, Jing He, Andrew Wimley, and the members of the New Orleans Protein Folding Intergroup. This work is supported by the National Institutes of Health GM060000 and the Louisiana Board of Regents Support Fund RC/EEP-05 (2007-2010).

**Supporting Information Available:** This material is available free of charge via the Internet at http://pubs.acs.org.
REFERENCES


