Review

A Dynamic Overview of Antimicrobial Peptides and Their Complexes

Viviane Silva de Paula 1 and Ana Paula Valente 2,*

1 Department of Chemistry and Biochemistry, University of California Santa Cruz, Santa Cruz, CA 95064, USA; vsilvade@ucsc.edu
2 Centro de Biologia Estrutural e Bioimagem, Instituto de Bioquímica Médica, Centro Nacional de Resonância Magnética Nuclear Jiri Jonas, Universidade Federal do Rio de Janeiro, Rio de Janeiro 21941-902, Brazil
* Correspondence: valente.anap@gmail.com or anapval@bioqmed.ufrj.br; Tel.: +55-021-3104-2326

Academic Editor: Brian F. Volkman

Received: 20 July 2018; Accepted: 9 August 2018; Published: 15 August 2018

Abstract: In this narrative review, we comprehensively review the available information about the recognition, structure, and dynamics of antimicrobial peptides (AMPs). Their complex behaviors occur across a wide range of time scales and have been challenging to portray. Recent advances in nuclear magnetic resonance and molecular dynamics simulations have revealed the importance of the molecular plasticity of AMPs and their abilities to recognize targets. We also highlight experimental data obtained using nuclear magnetic resonance methodologies, showing that conformational selection is a major mechanism of target interaction in AMP families.

Keywords: antimicrobial peptides; defensins; NMR; structure and dynamics; conformational selection

1. Introduction

The biological functions of proteins are coded in their structures, and dynamic and conformational changes are key events in all processes that involve protein recognition [1–4]. Nuclear magnetic resonance (NMR) and molecular simulations have shown that protein structure is heterogeneous and can adopt various conformational states that are energetically similar, separated by energy barriers on the order of kT (the product of the Boltzmann constant, k, and the temperature, T) [2,3,5,6]. In the presence of a ligand, a specific conformation is trapped, characterizing a conformational selection mechanism [7,8]. In the unbound state, a protein coexists with the bound state in some small fraction, and this fraction can bind to the ligand to reach the fully bound state. Further accommodation can occur, characterizing an induced-fit event [9]. To allow crossing at physiological temperatures, the barriers must be small because the dynamics are driven by the thermal fluctuations of the surrounding liquid [2,5]. Additionally, loss of entropy in the protein, which opposes binding, is often paired with disorder in remote regions, usually not correlated with the binding site in addition to changes in entropy of the solvent. New flexible regions often arise in the relaxed bound state or protecting exposed hydrophobic region and increasing the entropy in the well-known enthalpy-entropy compensation mechanism [2,10]. Regions with multiple states in the conformational space are accessed via local thermal fluctuations, and they play important roles in the entropic-enthalpic balance during recognition and binding. By identifying and analyzing the regions of local flexibility and cooperative motion of the residues inside a protein, we argue that it is possible to determine which parts of the protein will lead the kinetics of biologically relevant processes. These regions are typically characterized by highly conserved residues within a family of proteins with related biological functions and are involved in binding interactions [2,3,10].

This complex behavior occurs across a wide range of time scales and has been challenging to portray. These changes can range from side chain motions to domain reorientation; they therefore
occur at differing time scales and energy barriers. Recent advances in NMR and molecular dynamics have provided experimental evidence and offer direct measurement of protein motion in solution at atomic resolutions [11].

In this review, we focus primarily on applying NMR methods to the elucidation of interactions between antimicrobial peptides (AMPs) and their targets. We summarize the methods that can be used to reveal molecular mechanisms of interaction, beginning with structure determination using NMR and proceeding to a high-resolution view of the structures and dynamics of their complexes.

2. Antimicrobial Peptides

All multicellular organisms produce AMPs as a first line of defense. More than 2500 AMPs have been deposited in the Antimicrobial Peptide Database (http://aps.unmc.edu/AP/main.php), and they represent a diversity of activities against bacteria, yeasts, fungi, viruses, and cancer cells [12,13]. Although their mechanisms of action are not completely understood, a large number of AMPs act with high selectivity and specificity through interactions with membrane lipid components and components of the cellular matrix, targeting cytoplasmic components and interfering with cellular metabolism [12–14]. In solution, these peptides undergo complex conformational changes related to their multiple targets and mechanisms of action [15–17]. This review is mainly about defensins, which are AMPs, but not all AMPs are proteins (but rather short peptides) and not all AMPs adopt a well-defined structure, unless in the presence of a membrane.

Defensins represent a class of cationic AMPs that play pivotal roles in innate and adaptive immunities against microbial and viral infections and play various roles in inflammation, wound repair, expression of cytokines and chemokines, production of histamine, and enhancement of antibody responses [17]. They constitute an ancient and diverse gene family, present in most multicellular organisms ranging from plants, fungi, insects, mollusks, and arachnids to mammals, including humans. Defensins are a family of small, conserved proteins (~5 kDa). They are cysteine-rich AMPs and are classified into three subgroups, α-, β-, and θ-defensins, based on their amino acid sequences and patterns of disulfide bond connectivity [18–22]. All defensin structures contain a β-sheet fold that is stabilized by three disulfide bonds formed by six cysteine residues [23]. β-Defensins contain three intramolecular disulfide bonds in a I–V, II–IV, III–VI arrangement [24,25]. Although genome-based analyses have identified 28 human genes codified for human β-defensins (HBDs) [26], to date, only a few have been isolated and studied at the protein level. Human β-defensins 1–4 are predominantly secreted by epithelial cells and the male reproductive tract [27]. Both HBD5 and HBD6 appear to be specifically expressed in the epididymis [28].

Even though sequence identity between defensins varies from about 20% to 90%, and significant differences are evident in their biological functions, all defensins described thus far adopt essentially the same fold. Figure 1 shows a superposition of five defensins (HBD1, HBD2, HBD3, HBD4, and HBD6) to illustrate this similarity. Each of these structures consists of an N-terminal α helix and a three-stranded antiparallel β sheet. The root mean square deviation (RMSD) value across the Cα backbone atoms for these five β-defensin structures is 1.99 Å, but this varies within the N-terminal region due in part to increased flexibility (see Section 2.1).

The highly conserved cysteine residues pair up to form disulfide bridges that are crucial to maintaining structural integrity, which is a prerequisite for defensin activity.
2.1. Conformational Dynamics of Defensins

Nuclear magnetic resonance is currently the most powerful technique used to characterize the structural and dynamic properties of AMPs in complexes with their potential molecular targets, such as micelles and other membrane mimetics or oligosaccharides and peptides. To apply advanced NMR techniques, relatively large amounts of isotope-labeled proteins containing $^{13}$C, $^{15}$N, or $^2$H are desired [30,31]. Therefore, a number of methods for producing recombinant AMPs in bacteria have been developed [32–34]. Recombinant AMPs can be challenging to express because they can be toxic to their host bacteria and susceptible to proteases. We successfully expressed five β-defensins (HBD1, HBD4, HBD5, HBD6, and HBD11) as soluble fusion proteins using thioredoxin in *Escherichia coli* [32] (for HBD1 and HBD6). The pET32 vector uses thioredoxin as a fusion partner, increasing expression level, stability, and solubility [35].

Recent NMR studies of defensin dynamics [32,36,37] have revealed that these heavily cross-linked small proteins exhibit extensive conformational dynamics. Micro- to millisecond conformational exchanges are extensive, attributable to transient interconversions between the native ground state and energetically less-favorable ‘excited’ state(s). Studies have shown that transitions to such sparsely populated conformers are implicated in many aspects of protein function [38–40].

Interestingly, HBD1 and HBD6 $^{15}$N backbone relaxation analyses (Figure 2) identified regions undergoing microsecond to millisecond motions encompassing the α-helix and the loop between β1 and β2 strands [32]. These data showed that the conformational exchanges undergone by HBD1 and HBD6 could represent a mechanism conserved across the β-defensin family. This study emphasizes the conformational plasticity in β-defensins, which can play an important role in receptor interactions and is an important consideration in engineering defensins. However, only two β-defensin dynamics have been characterized thus far, circumventing extrapolation to other β-defensins.
2.2. Sugar Cane Defensin 5

This complex behavior has also been observed in sugar cane defensin 5 (SD5), which exhibits complex millisecond conformational dynamics involving all secondary structure elements [37]. Temperature-dependent Carr-Purcell-Meiboom-Gill NMR relaxation dispersion measurements probed the sparsely populated “excited” state of SD5, revealing it to be enthalpically unfavorable, suggesting a rearrangement of stabilizing contacts formed by surface-exposed side chains and/or a secondary structure. Overall, the emerging depiction of SD5 dynamics suggests this protein can populate either of two well-ordered conformational states, the excited conformer being more compact than the native state, and having a distinct secondary structure and side chain arrangements. The observation of an energetically unfavorable yet more compact excited state reveals a remarkable evolution of the CS α/β fold to expose and reorganize hydrophobic residues (Figure 3), enabling the creation of versatile binding sites and mechanisms for enthalpic-entropic compensation [41,42].

2.3. Dynamics Obtained Using Molecular Dynamics Simulations

Few defensin dynamics are characterized using NMR parameters; however, interesting results have been obtained using molecular dynamics. Sharadadevi and Nagaraj studied HBD1 and HNP-3 in water, and they observed structural fluctuations even in the presence of disulfide bonds [43].
Furthermore, both these entities unfold during MD. One interesting structural feature of HBD1, according to NMR data, is the lack of strong and well-defined interactions between secondary structure elements [44]. Also, deuterium exchange does not indicate the presence of extensive hydrogen bonding between the β-strands [45]. Also, common structural feature among defensins is their lack of hydrophobic cores and lack of extensive hydrogen bonding, implying conformation plasticity, an important characteristic for biological functions involving differing targets. The structure is stabilized primarily by disulfide bonds, allowing it to tolerate extensive residue substitution without disturbing the final tertiary structure. Molecular dynamics simulations performed by Zhang et al. revealed microsecond-long motions in α-defensin 5 and β-defensin 3 [45]. As also shown by their NMR results, despite the presence of disulfide bonds, these defensins present dynamic properties. Toubar et al. observed sliding monomers in dimer formation among HBD1 and mutants, suggesting that in other forms of oligomers as well, dynamics are observed and found to be important for the biological functions of HBD1 [46].

2.4. Analyzing Defensin Complexes Using NMR

The primary advantage of NMR is its ability to detect and analyze dynamic systems. Biomolecular NMR applications demand a combination of experiments applied to isotope-labeled samples. To map a ligand-protein binding site within a protein, 1H-15N correlation experiments are typically used, and this requires that the protein be 15N labeled. These fingerprint spectra of the protein can be easily recorded, and they allow tracking of the ligand binding event at residue resolution (Figure 4).

![Protein NMR: Binding site mapping probed by amide chemical shift perturbation](image)

**Figure 4.** Nuclear magnetic resonance (NMR) as a tool to determine ligand-defensin complex formation. (A) Once the backbone chemical shift assignment is accomplished, a 1H-15N heteronuclear single quantum coherence (HSQC) spectrum of the free protein containing the assigned amide groups is obtained. (B) Titration of an unlabeled ligand (in this example, a glycosaminoglycan [GAG]) bound to 15N-labeled human β-defensin (HBD) results in a chemical shift perturbation map of HBD6-ligand interactions. The NMR titration revealed large changes in the chemical shift of HBD6 in a fast-exchange equilibrium between free and GAG-bound HBD6, as demonstrated by a gradual shift in peak positions in 1H-15N HSQC spectra. (C) The amide groups significantly perturbed upon GAG binding are highlighted on the HBD6 surface (protein data bank: 2LWL) in blue and cyan to map the binding sites for the GAG. Figure adapted from [41]. For further details see Section 2.5.
Once the backbone chemical shift assignment is accomplished, chemical shift analysis gives information about the structure [47,48]. These chemical shift perturbations (CSPs) are then mapped onto the protein structure to define binding sites—regions that were perturbed in the presence of the ligand (Figure 4C). Changes in peak line width and intensity are also useful, reflecting changes in size and/or dynamics.

Changes in the rotational correlation time (determined using NMR relaxation times) or the translational self-diffusion coefficient (measured using pulsed-field gradient NMR diffusion methods) have also been used to detect binding and dimerization and to assess the $K_d$ value of the dimer. Analysis of NMR data offers structural insights into ligand-protein interactions at residue resolution. The information obtained is used to define areas of mutagenesis, guiding the design of functional studies, etc.

2.5. Defensin Interactions with Glycosaminoglycans and the Role of Dimerization

Glycosaminoglycans (GAGs) are sulfated polysaccharides, important components of the extracellular matrix. Several studies have established that cellular trafficking is orchestrated through chemokine-GAG interactions, which dictate the duration and steepness of chemokine gradients [49–52]. Although the roles of chemokine oligomerization and GAG binding have been known for some time, details of how defensins are recognized and immobilized by GAGs remain elusive. Of the five β-defensins for which 3-dimensional structures are currently known, only HBD2 and HBD6 have been characterized, via NMR, in their abilities to bind to GAGs [41,53].

For a range of GAGs, including heparin/heparan sulfate, dermatan sulfate, and chondroitin sulfate, Seo et al. elucidated the sites where binding on HBD2 takes place. Using NMR CSP analysis, five basic residues (Arg22, Arg23, Lys25, Lys39, and Lys40) of HBD2 were found to be involved in the formation of complexes [53]. In addition, HBD2 dimerization has been characterized using mass spectrometry and hydrodynamic calculations, establishing that it is induced by GAGs. Interestingly, the three basic residues in the $^{22}$RRY$^{25}$ stretch form a BBXB (B representing a basic amino acid, and X representing any amino acid) heparin-binding motif, which together with K39 and K40, creates a positive patch on the HBD2 surface (Figure 5A). This identified binding site is closely similar to the heparin binding site of the chemokine RANTES [54], which also contains the BBXB motif ($^{44}$RKNR$^{47}$) together with two additional spatially proximate basic residues, R20 and K25 [53].

In our studies of HBD6, we used a combination of structural and in silico analyses to characterize its interactions with GAGs [41]. We found that the HBD6-GAG complex reveals residues located at the α-helix and the loop between the β and β strands comprising the oligosaccharide binding interface. Interestingly, residues Lys-29, Ser-30, and Lys-32 showed increased R2 values in the presence of GAGs compared to free HBD6 in solution. Another interesting feature of this defensin is that the picosecond to nanosecond dynamics observed in the first residues located in the α-helix in the free state were reduced in the GAG-bound form, indicating that the α-helix becomes more structured upon GAG binding. These data suggest that Ser-30, Lys-5, Lys-29, and Lys-32, are parts of the BBXB motif, a known feature observed in heparin-binding proteins (Figures 4 and 5A). Additionally, our data suggest a dynamical compensation in the structure upon complex formation. Finally, high-pressure NMR and relaxation parameters showed that GAG binding modulates multiple structural and dynamic properties of HBD6, inducing formation of a ternary complex with 2:1 HBD6:GAG stoichiometry (see Figure 5).

The low sequence identity (17%) between HBD2 and HBD6 imposes difficulties in identifying regions important for dimerization and GAG interaction; nevertheless, the same basic residues patch-mapped in HBD2 are conserved in HBD3 and HBD6 (Figure 5B). The basic patch is formed by various residue types and could explain differences in interactions.
2.6. Modeling Defensin Complexes Using High Ambiguity Driven Docking

Nuclear magnetic resonance allows in vitro structural studies of protein complexes [55]; however, elucidation of entire structures is challenging because they are large in size and dynamics at interfaces between subunits leads to exchange-induced line broadening in the spectra. Determining and quantifying binding and cooperativity in protein complexes is challenging and requires a combination of biophysical methods and integrated structural biology methodologies. Computational docking approaches using experimental data have been used to generate models of a complex based on known 3-dimensional structures of the components in their free states. The high ambiguity driven docking (HADDOCK) method makes use of CSPs to model the structures of protein complexes [43,56–60]. Specifically, differences in the backbone and side chain chemical shifts upon complex formation are used to derive ambiguous distance restraints [61,62], thus driving the subunits to dock, given the changes are localized to the binding surface.

The defensin system complexed with various ligands has been explored using a combination of structural biology methodologies and docking guided by NMR CSP data [32,41]. Using HADDOCK, a rendition was created, and it indicates that two HBD6 monomers sandwich the GAG. When combined with high-pressure NMR and $^{15}$N relaxation data, HADDOCK is a powerful tool, capturing intermediate stages that could represent snapshots of the first stages of recognition and ligand binding [41] (Figure 6).

The importance of both defensin-GAG interactions and defensin oligomerization remains poorly understood. To define the defensin-GAG functional role in innate and adaptive immune responses, additional structural and cellular studies are necessary.
**Figure 6.** Dimerization and cooperativity during glycosaminoglycan (GAG) recognition by human β-defensin (HBD)6. (A) NMR relaxation experiments can reveal conformational exchanges and oligomerization (determined using NMR relaxation times) of defensins alone or complexed with GAGs. The $R_2/R_1$ ratios are translated during changes in the rotational correlation time, revealing the size of the complex. Note the grey areas for a direct comparison of the $R_2$ values between the two HBD6 states (free and GAG-bound). (B) ITC measurements dissect the binding mechanism of HBD6 and confirm the 2:1 stoichiometry. (C) Rendition of the “sandwich-like” dimerization mechanism. The lowest-energy structure of the HBD6-GAG complex within the highest-scoring NMR-restrained high ambiguity driven docking alternatives is chosen. Residues with substantial chemical shift perturbations upon GAG titration are highlighted in cyan (α-helix) and magenta (β2-β3). Hydrogen bonds and salt bridges formed by the sulfate moieties are represented by yellow dashed lines. Hydrophobic interactions between the aromatic rings of F1 residues from each monomer of HBD6 are represented by dark blue dots (right panel). The electrostatic potential of the ternary complex is indicated. Surface polarity is indicated as positive (blue), negative (red), or nonpolar (white; left panel). (D) High-pressure NMR confirms the sandwich-like dimerization mechanism and the induced dissociation of the HBD6-GAG ternary complex, revealing an intermediate stage. A series of $^1$H, $^{15}$N heteronuclear single quantum coherence spectra were recorded, varying the pressure. This data can reveal a snapshot of the first stage of GAG recognition whereby HBD6 exhibits features of a cooperative binding mechanism. This panel is adapted from [41] with permission from the original publisher.
Recently, Jarva et al. [63] showed that the dimerization interface of HBD2 is one of the binding sites of the lipid phosphatidylinositol 4,5-biphosphate (PIP2). The X-ray structure of HBD2 in the presence of PIP2 revealed two binding sites and shed important information in the mechanism of action of defensins.

2.7. Challenges to Understanding AMPs and Membrane Interaction

Antimicrobial peptides are usually short and cationic and exhibit extensive conformational diversity. Structure determination is often problematic because of their intricate dynamics, resulting in calculated structures with low convergence and geometrical violations [12,64–66]. On the other hand, characterization of AMP structures and dynamics is important so that their mechanisms of action can be understood, leading to the strategic design of new pharmacologically active compounds [15,16,67].

Membrane interaction is the first event in the AMP mechanism of action, and it is accompanied by a decrease in dynamics and acquisition of an amphipathic conformation. The AMP mechanism of action has been modeled by the barrel-stave model, the carpet model, and the toroidal pore model, among others. Solid-state NMR makes important contributions to the understanding of AMP organization and disruption in membranes [68–72]. One of the most-studied AMPs, magainin, lies in the bilayer interface and loosens the packing of the hydrophobic region, reducing the bilayer thickness [67]. Various studies confirm that the order parameter of the bilayer core decreases using deuterium NMR, possibly suggesting the most probable mechanism of disruption, but the complete picture is not yet known. Mani et al. used $^{19}$F to show lipid disorder near the toroidal pore formed by protegrin-1 [69].

Paramagnetic Relaxation Enhancement (PRE) measurements can also provide information about AMPs inserted in micelles and membranes [73]. Taken together, these parameters can aid in the interpretation of the relationships between PRE values and peptide orientations in micelles.

To contribute to this work, we examined tritripticin (TRF3), a cathelicidin AMP that exists as an ensemble of four conformations in solution [74]. The indole region therefore produces 12 peaks in $^1$H NMR spectra rather than the expected three. In the presence of micelles, one major conformation is stable [74–77].

Using NMR to evaluate the structure of TRF3 in micelles with various chemical features reveals interactions occurred through conformational selection, stabilizing similar structures despite differences in detergent headgroups or dipole orientations [78]. These data also reveal the imperative constraint of the WWW-F motif in the TRF3 structure, acquired at interfaces, and show that pre-organization in the free state is crucial for membrane interaction.

A large number AMPs act with high selectivity and specificity through interactions with membrane lipid components. These peptides undergo complex conformational changes in solution, but upon binding to an interface, one major conformation is stabilized.

A number of theoretical and computer simulation approaches have been established to describe AMP and lipid interactions. The MD is an “in silico” approach that can characterize the interaction between AMPs and membranes, in which the dynamics of the molecule into the lipid bilayer and its affects on surrounding lipids are explored. MD simulations have succeeded in precisely describing the process of peptide binding, folding and partitioning into lipid bilayers, at atomic resolution [79,80].

Recently, Wang et al. (2015) used experimentally guided unbiased MD in order to characterize the molecular mechanisms of pore formation by maculatin, a well-known AMP from the Australian tree frog *Litoria genimaculata*, revealing that rather than a single pore or detergent-like holes, maculatin forms an ensemble of structurally different channels that continually form and disperse at equilibrium [81,82]. Membrane permeabilization is dominated by oligomeric forms of maculatin, which conduct water, ions and small dyes.

Lee et al. (2016) explored the adsorption mechanism of human β-defensin 3 (HBD3) using Gram-negative (GN), Gram-positive (GP), and mammalian membrane models, through MD simulations [83]. They reach to the conclusion that Arg17, Arg36 and Arg38 form both polar
and nonpolar interactions and are potentially the key residues for HBD3 antibacterial activity. Also, they concluded that high concentration of POPG lipids in the GP membrane leads to strong electrostatic interaction with HBD3 and stabilize its adsorption rapidly as compared with HBD3-GN membrane complex.

3. Conclusions

Antimicrobial peptides are important for innate and adaptive immune responses, and they can provide a variety of avenues in drug discovery. The universe of biological functions is linked to the binding promiscuity of AMPs, which is related to their ability to oligomerize and to interact with membranes and elements of the cellular matrix. All possibilities are correlated to their dynamical properties because regions that have multiple states are associated with recognition and binding. By identifying and analyzing the regions of local flexibility and cooperative motion of the residues inside a protein, those parts of the protein that lead the kinetics of the biologically relevant processes can be elucidated.

Author Contributions: All authors (V.S.d.P. and A.P.V.) contributed equally to this work. All authors critically reviewed the manuscript. All the authors read and approved the final manuscript.

Funding: This work was supported by the Brazilian agencies Fundação de Amparo a Pesquisa do Estado do Rio de Janeiro (FAPERJ), and Conselho Nacional de Pesquisa e Desenvolvimento (CNPq) and Instituto Nacional de Ciência e Tecnologia de Biologia Estrutural e Bioimagem (INCT-INBEB).

Acknowledgments: We thank Centro Nacional de Ressonância Magnética Nuclear (CNRMN) for support.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. James, L.C.; Tawfik, D.S. Conformational diversity and protein evolution—A 60-year-old hypothesis revisited. Trends Biochem. Sci. 2003, 28, 361–368. [CrossRef]
2. Copperman, J.; Guenza, M.G. Mode localization in the cooperative dynamics of protein recognition. J. Chem. Phys. 2016, 145, 015101.
3. Okazaki, K.I.; Takada, S. Dynamic energy landscape view of coupled binding and protein conformational change: Induced-fit versus population-shift mechanisms. Proc. Natl. Acad. Sci. USA 2008, 105, 11182–11187. [CrossRef] [PubMed]
4. Clore, G. Interplay between conformational selection and induced fit in multidomain protein–ligand binding probes by paramagnetic relaxation enhancement. Biophys. Chem. 2014, 186, 3–12. [CrossRef] [PubMed]
5. Mobley, D.L.; Dill, K.A. Binding of Small-Molecule Ligands to Proteins: “What You See” Is Not Always “What You Get”. Structure 2009, 17, 489–498. [CrossRef] [PubMed]
6. Boehr, D.; Nussinov, R.; Wright, P. The role of dynamic conformational ensembles in biomolecular recognition. Nat. Chem. Biol. 2009, 5, 789–796. [CrossRef] [PubMed]
7. Monod, J.; Wyman, J.; Changeux, J.P. On the nature of allosteric transitions—A plausible model. J. Mol. Biol. 1965, 12, 88–118. [CrossRef]
8. Nussinov; Ma, B.; Tsai, C.J. Multiple conformational selection and induced fit events take place in allosteric propagation. Biophys. Chem. 2014, 186, 22–30. [CrossRef] [PubMed]
9. Koshland, D. Application of a theory of enzyme specificity to protein synthesis. Proc. Natl. Acad. Sci. USA 1958, 44, 98–104. [CrossRef] [PubMed]
10. Popovych, N.; Sun, S.; Ebright, R.H.; Kalodimos, C.G. Dynamically driven protein allostery. Nat. Struct. Mol. Biol. 2006, 13, 831–838. [CrossRef] [PubMed]
11. Sekhar, A.; Velyvis, A.; Zoltsman, G.; Rosenzweig, R.; Bouvignies, G.; Kay, L.E. Conserved conformational selection mechanism of Hsp70 chaperone-substrate interactions. eLife 2018, 7, e32764. [CrossRef] [PubMed]
12. Epand, R.; Vogel, H. Diversity of antimicrobial peptides and their mechanisms of action. Biochim. Biophys. Acta 1999, 1462, 11–28. [CrossRef]
13. Zhang, L.J.; Gallo, R.L. Antimicrobial peptides. Curr. Biol. 2016, 26, R14-9. [CrossRef] [PubMed]
14. Travkova, O.G.; Moehwald, H.; Brezesinski, G. The interaction of antimicrobial peptides with membranes. *Adv. Colloid Interface Sci.* 2017, 247, 521–532. [CrossRef] [PubMed]

15. Gomes-Neto, F.; Valente, A.P.; Almeida, F.C.L. Modeling the interaction of dodecylphosphocholine micelles with anticoagulant peptide PW2 guided by NMR data. *Molecules* 2013, 18, 10056–10080. [CrossRef] [PubMed]

16. Valente, A.P.; Miyamoto, C.; Almeida, F.C.L. Implications of protein conformational diversity for binding and development of new biological active compounds. *Curr. Med. Chem.* 2006, 13, 3697–3703. [CrossRef] [PubMed]

17. Mattar, E.H.; Almehdar, H.A.; Yacoub, H.A.; Uversky, V.N.; Redwan, E.M. Antimicrobial potentials and structural disorder of human and animal defensins. *Cytokine Growth Factor Rev.* 2016, 28, 95–111. [CrossRef] [PubMed]

18. Ganz, T. Defensins: Antimicrobial peptides of innate immunity. *Nat. Rev. Immunol.* 2003, 3, 710–720. [CrossRef] [PubMed]

19. Lehrer, R.I. Primate defensins. *Nat. Rev. Microbiol.* 2004, 2, 727–738. [CrossRef] [PubMed]

20. Selsted, M.E.; Ouellette, A.J. Mammalian defensins in the antimicrobial immune response. *Nat. Immunol.* 2005, 6, 551–557. [CrossRef] [PubMed]

21. Hazlett, L.; Wu, M. Defensins in innate immunity. *Cell Tissue Res.* 2011, 343, 175–188. [CrossRef] [PubMed]

22. Verma, C.; Seebah, S.; Low, S.M.; Zhou, L.; Liu, S.P.; Li, J.; Beuerman, R.W. Defensins: Antimicrobial peptides for therapeutic development. *Biotechnol. J.* 2007, 11, 1353–1359. [CrossRef] [PubMed]

23. Yang, D.; Biragyn, A.; Kwak, L.W.; Oppenheim, J.J. Mammalian defensins in immunity: More than just microbicidal. *Trends Immunol.* 2002, 23, 291–296. [CrossRef] [PubMed]

24. Lehrer, R.I.; Ganz, T. Defensins of vertebrate animals. *Curr. Opin. Immunol.* 2002, 1, 96–102. [CrossRef]

25. Schibli, D.J.; Hunter, H.N.; Aseyev, V.; Starner, T.D.; Wienczek, J.M.; McCray, P.B., Jr.; Tack, B.F.; Vogel, H.J. The solution structures of the human beta-defensins lead to a better understanding of the potent bactericidal activity of HBD3 against *Staphylococcus aureus*. *J. Biol. Chem.* 2002, 277, 8279–8289. [CrossRef] [PubMed]

26. Schuttle, B.C.; Mitros, J.P.; Bartlett, J.A.; Walters, J.D.; Jia, H.P.; Welsh, M.J.; Casavant, T.L.; McCray, P.B., Jr. Discovery of five conserved beta-defensin gene clusters using a computational search strategy. *Proc. Natl. Acad. Sci. USA* 2002, 99, 2129–2133. [CrossRef] [PubMed]

27. Yang, D.; Biragyn, A.; Hoover, D.M.; Lubkowski, J.; Oppenheim, J.J. Multiple roles of antimicrobial defensins, cathelicidins, and eosinophil-derived neurotoxin in host defense. *Annu. Rev. Immunol.* 2004, 22, 181–215. [CrossRef] [PubMed]

28. Yamaguchi, Y.; Nagase, T.; Makita, R.; Fukuhara, S.; Tomita, T.; Tominaga, T.; Kurihara, H.; Ouchi, Y. Identification of multiple novel epididymis-specific beta-defensin isoforms in humans and mice. *J. Immunol.* 2002, 169, 2516–2523. [CrossRef] [PubMed]

29. Ye, Y.; Godzik, A. Flexible structure alignment by chaining aligned fragment pairs allowing twists. *Bioinformatics* 2003, 19, ii246–255. [CrossRef] [PubMed]

30. Hiroaki, H. Recent applications of isotopic labeling for protein NMR in drug discovery. *Expert Opin. Drug Discov.* 2013, 8, 523–536. [CrossRef] [PubMed]

31. Sugiki, T.; Fujiwara, T.; Kojima, C. Latest approaches for efficient protein production in drug discovery. *Expert Opin. Drug Discov.* 2014, 9, 1189–1204. [CrossRef] [PubMed]

32. De Paula, V.S.; Gomes, N.S.; Lima, L.G.; Miyamoto, C.A.; Monteiro, R.Q.; Almeida, F.C.; Valente, A.P. Structural basis for the interaction of human beta-defensin 6 and its putative chemokine receptor CCR2 and breast cancer microvesicles. *J. Mol. Biol.* 2013, 425, 4479–4495. [CrossRef] [PubMed]

33. Ishida, H.; Nguyen, L.T.; Gopal, R.; Aizawa, T.; Vogel, H.J. Overexpression of Antimicrobial, Anticancer, and Transmembrane Peptides in Escherichia coli through a Calmodulin-Peptide Fusion System. *J. Am. Chem. Soc.* 2016, 138, 11318–11326. [CrossRef] [PubMed]

34. Haney, E.F.; Mansour, S.C.; Hancock, R.E. Antimicrobial Peptides: An Introduction. *Methods Mol. Biol.* 2017, 1548, 3–22. [PubMed]

35. LaValle, E.R.; DiBlasio, E.A.; Kovacic, S.; Grant, K.L.; Schendel, P.F.; McCoy, J.M. A thioredoxin gene fusion expression system that circumvents inclusion body formation in the *E. coli* cytoplasm. *Biotechnology* 1993, 11, 187–193. [CrossRef] [PubMed]

36. De Medeiros, L.N.; Angeli, R.; Sarzedas, C.G.; Barreto-Bergter, E.; Valente, A.P.; Kurtenbach, E.; Almeida, F.C.L. Backbone dynamics of the antifungal Psd1 pea defensin and its correlation with membrane interaction by NMR spectroscopy. *Biochim. Biophys. Acta* 2010, 1798, 105–113. [CrossRef] [PubMed]
37. De Paula, V.S.; Razzera, G.; Barreto-Berгер, E.; Almeida, F.C.L.; Valente, A.P. Portrayal of complex dynamic properties of sugarcane defensin 5 by NMR: Multiple motions associated with membrane interaction. *Structure* 2011, 19, 26–36. [CrossRef] [PubMed]

38. Mittermaier, A.K.; Kay, L.E. Observing biological dynamics at atomic resolution using NMR. *Trends Biochem. Sci.* 2009, 34, 601–611. [CrossRef] [PubMed]

39. Henzler-Wildman, K.; Kern, D. Dynamic personalities of proteins. *Nature* 2007, 450, 964–972. [CrossRef] [PubMed]

40. Korzhnev, D.M.; Religa, T.L.; Banachewicz, W.; Fersht, A.R.; Kay, L.E. A Transient and Low-Populated Protein-Folding Intermediate at Atomic Resolution. *Science* 2010, 329, 1312–1316. [CrossRef] [PubMed]

41. De Paula, V.S.; Pomin, V.H.; Valente, A.P. Unique properties of human β-defensin 6 (hBD6) and glycosaminoglycan complex: Sandwich-like dimerization and competition with the chemokine receptor 2 (CCR2) binding site. *J. Biol. Chem.* 2014, 289, 22969–22979. [CrossRef] [PubMed]

42. Machado, L.E.S.F.; De Paula, V.S.; Pustovalova, Y.; Bezsonova, I.; Valente, A.P.; Korzhnev, D.M.; Almeida, F.C.L. Conformational Dynamics of a Cysteine-Stabilized Plant Defensin Reveals Evolutionary Mechanism to Expose Hydrophobic Residues. *Biochemistry* 2018, in press.

43. Sharadadevi, A.; Nagaraj, A. A Molecular dynamics study of human defensins HBD-1 and HNP-3 in water. *J. Biomol. Struct. Dyn.* 2010, 27, 541–550. [CrossRef] [PubMed]

44. Zhang, L. Different dynamics and pathway of disulfide bonds reduction of two human defensins, a molecular dynamics simulation study. *Proteins* 2017, 85, 665–681. [CrossRef] [PubMed]

45. Toubar, R.A.; Zhmurov, A.; Barsegov, V.; Marx, K.A. Comparative simulation studies of native and single-site mutant human beta-defensin-1 peptides. *J. Biomol. Struct. Dyn.* 2013, 31, 174–194. [CrossRef] [PubMed]

46. Gao, G.; Williams, J.G.; Campbell, S.L. Protein–protein interaction analysis by nuclear magnetic resonance spectroscopy. *Methods Mol. Biol.* 2004, 261, 79–92. [PubMed]

47. Palmer, A.G. NMR characterization of the dynamics of biomacromolecules. *Chem. Rev.* 2004, 104, 3623–3640. [CrossRef] [PubMed]

48. Hoogewerf, A.J.; Kuschert, G.S.; Proudfoot, A.E.; Borlat, F.; Clark-Lewis, I.; Power, C.A.; Wells, T.N. Glycosaminoglycans mediate cell surface oligomerization of chemokines. *Biochemistry* 1997, 36, 13570–13578. [CrossRef] [PubMed]

49. Crown, S.E.; Yu, Y.; Sweeney, M.D.; Leary, J.A.; Handel, T.M. Heterodimerization of CCR2 chemokines and regulation by glycosaminoglycan binding. *J. Biol. Chem.* 2006, 281, 25438–25446. [CrossRef] [PubMed]

50. Proudfoot, A.E.; Handel, T.M.; Johnson, Z.; Lau, E.K.; LiWang, P.; Clark-Lewis, I.; Borlat, F.; Wells, T.N.; Kosco-Vilbois, M.H. Glycosaminoglycan binding and oligomerization are essential for the in vivo activity of certain chemokines. *Proc. Natl. Acad. Sci. USA* 2003, 100, 1885–1890. [CrossRef] [PubMed]

51. Salanga, C.I.; Handel, T.M. Chemokine oligomerization and interactions with receptors and glycosaminoglycans: The role of structural dynamics in function. *Exp. Cell Res.* 2011, 317, 590–601. [CrossRef] [PubMed]

52. Seo, E.S.; Blaum, B.S.; Vargues, T.; DeCecco, M.; Deakin, J.A.; Lyon, M.; Barran, P.E.; Campopiano, D.J.; Uhrin, D. Interaction of human β-defensin 2 (HBD2) with glycosaminoglycans. *Biochemistry* 2010, 49, 10486–10495. [CrossRef] [PubMed]

53. Proudfoot, A.E.L.; Fritchley, S.; Borlat, F.; Shaw, J.P.; Vilbois, F.; Zwahlen, C.; Trkola, A.; Marchant, D.; Clapham, P.R.; Wells, T.N.C. The BBXB motif of RANTES is the principal site for heparin binding and controls receptor selectivity. *J. Biol. Chem.* 2001, 276, 10620–10626. [CrossRef] [PubMed]

54. Gujjarro, J.I.; Morton, C.J.; Flaxco, K.W.; Campbell, I.D.; Dobson, C.M. Folding kinetics of the SH3 domain of PI3 kinase by real-time NMR combined with optical spectroscopy. *J. Mol. Biol.* 1998, 276, 657–667. [CrossRef] [PubMed]

55. Van Dijk, A.D.J.; Boelens, R.; Bonvin, A.M.J.J. Data-driven docking for the study of biomolecular complexes. *FEBS J.* 2005, 272, 293–312. [CrossRef] [PubMed]

56. Dominguez, C.; Boelens, R.; Bonvin, A.M.J.J. HADDOCK: A protein-protein docking approach based on biochemical or biophysical information. *J. Am. Chem. Soc.* 2003, 125, 1731–1737. [CrossRef] [PubMed]

57. De Vries, S.J.; Van Dijk, M.; Bonvin, A.M.J.J. The HADDOCK web server for data-driven biomolecular docking. *Nat. Protoc.* 2010, 5, 883–897. [CrossRef] [PubMed]
59. Van Zundert, G.C.; Bonvin, A.M.J. Modeling Protein–Protein Complexes Using the HADDOCK Webserver, Modeling Protein Complexes with HADDOCK. Methods Mol. Biol. 2014, 1137, 163–179. [PubMed]
60. Van Zundert, G.C.P.; Rodrigues, J.P.G.L.M.; Trellet, M.; Schmitz, C.; Kastritis, P.L.; Karaca, E.; Melquiond, A.S.J.; Van Dijk, M.; De Vries, S.J.; Bonvin, A.M.J. The HADDOCK2.2 web server. User-friendly integrative modeling of biomolecular complexes. J. Mol. Biol. 2016, 428, 720–725. [CrossRef] [PubMed]
61. Dominguez, C.; Bonvin, A.M.J.J.; Winkler, G.S.; Van Schaik, F.M.A.; Timmers, H.T.M.; Boelens, R. Structural model of the UbcH5B/CNOT4 complex revealed by combining NMR, mutagenesis, and docking approaches. Structure 2004, 12, 633–644. [CrossRef] [PubMed]
62. Van Drogen-Petit, A.; Zwahlen, C.; Peter, M.; Bonvin, A.M. Insight into molecular interactions between two PBI domains. J. Mol. Biol. 2004, 336, 1195–1210. [CrossRef] [PubMed]
63. Järvå, M.; Phan, T.K.; Lay, F.T.; Caria, S.; Kvensakul, M.; Hulett, M.D. Human β-defensin 2 kills Candida albicans through phosphatidylinositol 4,5-bisphosphate-mediated membrane permeabilization. Sci. Adv. 2018, 4, eaat0979. [CrossRef] [PubMed]
64. Yeaman, M.; Yount, N. Mechanisms of Antimicrobial Peptide Action and Resistance. Pharmacol. Rev. 2003, 55, 27–55. [CrossRef] [PubMed]
65. Reddy, K.; Yedery, R.; Aranha, C. Antimicrobial peptides: Premises and promises. Int. J. Antimicrob. Agents 2004, 24, 536–547. [CrossRef] [PubMed]
66. Melo, M.; Ferre, R.; Castanho, M. Antimicrobial peptides: Linking partition, activity and high membrane-bound concentrations. Nat. Rev. Microbiol. 2009, 7, 245–250. [CrossRef] [PubMed]
67. Marquette, A.; Bechinger, B. Biophysical Investigations Elucidating the Mechanics of Action of Antimicrobial Peptides and Their Synergism. Biomolecules 2018, 8, 18. [CrossRef] [PubMed]
68. Bechinger, B.; Salnikov, E.S. The membrane interactions of antimicrobial peptides revealed by solid-state NMR spectroscopy. Chem. Phys. Lipids 2012, 165, 282–301. [CrossRef] [PubMed]
69. Mani, R.; Cady, S.D.; Tang, M.; Waring, A.J.; Lehrer, R.I.; Hong, M. Membrane-dependent oligomeric structure and pore formation of a β-hairpin antimicrobial peptide in lipid bilayers from solid-state NMR. Proc. Natl. Acad. Sci. USA 2006, 103, 16242–16247. [CrossRef] [PubMed]
70. Tang, M.; Hong, M. Structure and mechanism of beta-hairpin antimicrobial peptides in lipid bilayers from solid-state NMR spectroscopy. Mol. Biosyst. 2009, 5, 317–322. [CrossRef] [PubMed]
71. Tang, M.; Waring, A.; Hong, M. Phosphate-mediated arginine insertion into lipid membranes and pore formation by a cationic membrane peptide from solid-state NMR. J. Am. Chem. Soc. 2007, 129, 11438–11446. [CrossRef] [PubMed]
72. Porcelli, F.; Ramamoorthy, A.; Barany, G.; Veglia, G. On the role of NMR spectroscopy for characterization of antimicrobial peptides. Methods Mol. Biol. 2013, 1063, 159–180. [PubMed]
73. Respondek, M.; Madl, T.; Göbl, C.; Golser, R.; Zangger, K. Mapping the Orientation of Helices in Micelle-Bound Peptides by Paramagnetic Relaxation Waves. J. Am. Chem. Soc. 2007, 129, 5228–5234. [CrossRef] [PubMed]
74. Schibli, D.; Hwang, P.; Vogel, H. Structure of the antimicrobial peptide trittrippticin bound to micelles: A distinct membrane-bound peptide fold. Biochemistry 1999, 38, 16749–16755. [CrossRef] [PubMed]
75. Schibli, D.J.; Nguyen, L.T.; Vogel, H.J. Structure-function analysis of trittrippticin analogs: Potential relationships between antimicrobial activities, model membrane interactions, and their micelle-bound NMR structures. Biophys. J. 2006, 91, 4413–4426. [CrossRef] [PubMed]
76. Arias, M.; Jensen, K.V.; Vogel, H.J. Hydroxy-tryptophan containing derivatives of trittripticin: Modification of antimicrobial activity and membrane interactions. Biochim. Biophys. Acta 2015, 1848, 277–288. [CrossRef] [PubMed]
77. Bozelli, J.; Sasahara, E.T.; Pinto, M.R.; Nakaie, C.R.; Schreier, S. Effect of head group and curvature on binding of the antimicrobial peptide trittripticin to lipid membranes. Chem. Phys. Lipids 2012, 165, 365–373. [CrossRef] [PubMed]
78. Santos, T.L.; Moraes, A.; Nakaie, C.R.; Almeida, F.C.L.; Valente, A.P. Structural and dynamics insights of the interaction between Trittripcticin and micelles: An NMR study. Biophys. J. 2016, 111, 2676–2688. [CrossRef] [PubMed]
79. Wang, Y.; Zhao, T.; Wei, D.; Strandberg, E.; Ulrich, A.S.; Ulmschneider, J.P. How reliable are molecular dynamics simulations of membrane active antimicrobial peptides? Biochim. Biophys. Acta 2014, 1838, 2280–2288. [CrossRef] [PubMed]
80. Deleu, M.; Crowet, J.M.; Nasir, M.N.; Lins, L. Complementary biophysical tools to investigate lipid specificity in the interaction between bioactive molecules and the plasma membrane: A review. *Biochim. Biophys. Acta* 2014, 1838, 3171–3190. [CrossRef] [PubMed]

81. Wang, Y.; Chen, C.H.; Hu, D.; Ulmschneider, M.B.; Ulmschneider, J.P. Spontaneous formation of structurally diverse membrane channel architectures from a single antimicrobial peptide. *Nat. Commun.* 2016, 7, 13535. [CrossRef] [PubMed]

82. Ulmschneider, J.P.; Ulmschneider, M.B. Molecular Dynamics Simulations Are Redefining Our View of Peptides Interacting with Biological Membranes. *Acc. Chem. Res.* 2018, 51, 1106–1116. [CrossRef] [PubMed]

83. Lee, J.; Jung, S.W.; Cho, A.E. Molecular Insights into the Adsorption Mechanism of Human β-Defensin-3 on Bacterial Membranes. *Langmuir* 2016, 32, 1782–1790. [CrossRef] [PubMed]