

defensins and cancer

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Abstract

Introduction

Antimicrobial peptides (AMP) have been isolated and characterized from a wide range of animal, plant and bacterial species and are known to play important roles in the host defence system and innate immunity [1](#)

(AMP) are polypeptides of fewer than 100 amino acids contain a positive net charge (from +2 to +9), high lysine and arginine content, and a significant number of hydrophobic residues ($\geq 30\%$) [2](#).

These properties permit the peptides to fold into amphipathic conformations upon contact with cell membranes, and the positively charged polar face help the molecules bind to the membrane through electrostatic interaction with the negatively charged head groups of phospholipids [3](#)

The history of AMPs dates back to 1922 with the discovery of lysozyme by Alexander Flemming. The first reported animal originated AMP is defensin, which was isolated from rabbit leukocytes in 1956 [4](#). at the same time, it was also proven that human leukocytes contain AMPs in their lysosomes.

They are found two families of antimicrobial peptides have been implicated in antimicrobial activity of phagocytes, inflammatory and epithelial secretions. Defensins are widely distributed in mammalian epithelial cells and phagocytes, [5](#). Cathelicidins are structurally and evolutionarily distinct antimicrobial peptides that are similar to defensins in abundance and distribution [???](95)01050-o

References

1. **Design of an α -helical antimicrobial peptide with improved cell-selective and potent anti-biofilm activity**
Shi-Kun Zhang, Jin-wen Song, Feng Gong, Su-Bo Li, Hong-Yu Chang, Hui-Min Xie, Hong-Wei Gao, Ying-Xia Tan, Shou-Ping Ji
Scientific Reports (2016-06-08) <https://doi.org/ggt4kh>
DOI: [10.1038/srep27394](https://doi.org/10.1038/srep27394) · PMID: [27271216](https://pubmed.ncbi.nlm.nih.gov/27271216/) · PMCID: [PMC4897634](https://pubmed.ncbi.nlm.nih.gov/PMC4897634/)
2. **Defensins: antimicrobial peptides of innate immunity**
Tomas Ganz
Nature Reviews Immunology (2003-09) <https://doi.org/cx6gzg>
DOI: [10.1038/nri1180](https://doi.org/10.1038/nri1180) · PMID: [12949495](https://pubmed.ncbi.nlm.nih.gov/12949495/)
3. **Animal antimicrobial peptides: an overview.**
D Andreu, L Rivas
Biopolymers (1998) <https://www.ncbi.nlm.nih.gov/pubmed/10333735>
DOI: [10.1002/\(sici\)1097-0282\(1998\)47:6<415::aid-bip2>3.0.co;2-d](https://doi.org/10.1002/(sici)1097-0282(1998)47:6<415::aid-bip2>3.0.co;2-d) · PMID: [10333735](https://pubmed.ncbi.nlm.nih.gov/10333735/)
4. **PHAGOCYTIN: A BACTERICIDAL SUBSTANCE FROM POLYMORPHONUCLEAR LEUCOCYTES**
James G. Hirsch
The Journal of Experimental Medicine (1956-05-01) <https://doi.org/dzvswf>
DOI: [10.1084/jem.103.5.589](https://doi.org/10.1084/jem.103.5.589) · PMID: [13319580](https://pubmed.ncbi.nlm.nih.gov/13319580/) · PMCID: [PMC2136631](https://pubmed.ncbi.nlm.nih.gov/PMC2136631/)
5. **Defensins. Natural peptide antibiotics of human neutrophils.**
T Ganz, ME Selsted, D Szklarek, SS Harwig, K Daher, DF Bainton, RI Lehrer
Journal of Clinical Investigation (1985-10-01) <https://doi.org/bb8wz4>
DOI: [10.1172/jci112120](https://doi.org/10.1172/jci112120) · PMID: [2997278](https://pubmed.ncbi.nlm.nih.gov/2997278/) · PMCID: [PMC424093](https://pubmed.ncbi.nlm.nih.gov/PMC424093/)