

Dermaseptin

Cat. No.:	HY-P0263
CAS No.:	136212-91-4
Molecular Formula:	$C_{152}H_{257}N_{43}O_{44}S_2$
Molecular Weight:	3455.1
Sequence:	Ala-Leu-Trp-Lys-Thr-Met-Leu-Lys-Lys-Leu-Gly-Thr-Met-Ala-Leu-His-Ala-Gly-Lys-Ala-Ala-Leu-Gly-Ala-Ala-Ala-Asp-Thr-Ile-Ser-Gln-Gly-Thr-Gln ALWKTMLKKLGTMLHAGKAALGAAADTISQGTQ
Sequence Shortening:	ALWKTMLKKLGTMLHAGKAALGAAADTISQGTQ
Target:	Bacterial; Fungal; Antibiotic
Pathway:	Anti-infection
Storage:	Sealed storage, away from moisture Powder -80°C 2 years -20°C 1 year * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)

SOLVENT & SOLUBILITY

In Vitro	H ₂ O : 25 mg/mL (7.24 mM; Need ultrasonic)				
	Preparing Stock Solutions	<div>Solvent Concentration</div> <div>Mass</div>	1 mg	5 mg	10 mg
		1 mM	0.2894 mL	1.4471 mL	2.8943 mL
		5 mM	0.0579 mL	0.2894 mL	0.5789 mL
		10 mM	---	---	---
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: PBS Solubility: 50 mg/mL (14.47 mM); Clear solution; Need ultrasonic				

BIOLOGICAL ACTIVITY

Description	Dermaseptin, a peptide isolated from frog skin, exhibits potent antimicrobial activity against bacteria, fungi, and protozoa at micromolar concentration ^[1] .
In Vitro	Dermaseptin is a water-soluble, thermostable, and nonhemolytic peptide endowed with highly potent antimicrobial activity against pathogenic fungi at micromolar concentration. Circular dichroism spectra of dermaseptin in hydrophobic media indicated 80% alpha-helical conformation, and predictions of secondary structure suggested that dermaseptin can be configured as an amphiphatic alpha-helix spanning over residues 1-27, a structure that perturbs membrane functions regulating water flux ^[1] . Dermaseptin exerts a lytic action upon bacteria, protozoa, yeasts, and filamentous fungi at micromolar concentrations. Molecular elements responsible for the exceptional antimicrobial potency of dermaseptin are

to be traced to the NH₂-terminal alpha-helical amphipathic segment spanning residues 1-18 of the molecule^[1]. Dermaseptin (5-100 µg/ml; 48 hours) inhibits by 100% the proliferation of most microorganisms tested, including Gram-positive or Gram-negative bacteria, parasites, yeasts, and filamentous fungi, at micromolar concentrations^[2]. Dermaseptin (5-100 µg/ml; 48 hours) does not inhibit the proliferation of human KJ3 cells after a 48 h incubation, and dermaseptin treatment for 1 h does not permeate guinea pig lymphocytes up to the highest concentration assayed (200 µg/ml). Hemolysis of rabbit erythrocytes occurs after 1 h of treatment at doses above 200 µg/ml, with 50% hemolysis at 350 µg/ml^[2]. Dermaseptin has antimicrobial activities and is against *Aeromonas cauiiae*, *Pseudomonas aeruginosa*, *Escherichia coli*, *Enterococcus faecalis*, *L. mezicana* (NF^α strain) and *Microsporum canis* (IP1194) with MIC values of 50 µg/ml; 100 µg/ml; 25 µg/ml; 15 µg/ml; and 50 µg/ml, respectively^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

- [1]. Mor A, et al. Isolation, amino acid sequence, and synthesis of dermaseptin, a novel antimicrobial peptide of amphibian skin. *Biochemistry*. 1991 Sep 10;30(36):8824-30.
- [2]. Mor A, et al. The NH₂-terminal alpha-helical domain 1-18 of dermaseptin is responsible for antimicrobial activity. *J Biol Chem*. 1994 Jan 21;269(3):1934-9.

Caution: Product has not been fully validated for medical applications. For research use only.

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