Cecropin P1, porcine

Cat. No.:	HY-P2317			
CAS No.:	125667-96-1			
Molecular Formula:	C ₁₄₇ H ₂₅₃ N ₄₅ O ₄₃			
Molecular Weight:	3338.86 SWLSKTAKKLENSAKKRISEGIAIAIQGGPR			
Sequence:	Ser-Trp-Leu-Ser-Lys-Thr-Ala-Lys-Lys-Leu-Glu-Asn-Ser-Ala-Lys-Lys-Arg-Ile-Ser-Glu-Gly- Ile-Ala-Ile-Ala-Ile-Gln-Gly-Gly-Pro-Arg			
Sequence Shortening:	SWLSKTAKKLENSAKKRISEGIAIAIQGGPR			
Target:	Endogenous Metabolite; Bacterial			
Pathway:	Metabolic Enzyme/Protease; Anti-infection			
Storage:	Sealed storage, away from moisture and light, under nitrogen Powder -80°C 2 years -20°C 1 year * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light, under nitrogen)			

SOLVENT & SOLUBILITY

	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
		1 mM	0.2995 mL	1.4975 mL	2.9950 mL
		5 mM	0.0599 mL	0.2995 mL	0.5990 mL
		10 mM	0.0300 mL	0.1498 mL	0.2995 mL

BIOLOGICAL ACTIVITy Description Cecropin P1, porcine is an antibacterial peptide that can be isolated from the upper part of the small intestine of the pig. Cecropin P1, porcine shows antibacterial activity against Gram-negative bacteria. Cecropin P1, porcine shows antiviral activity and inhibits PRRSV infection^{[11][2]}. In Vitro Cecropin P1, porcine (0-480 µg/mL, 36-96 h) markedly inhibits CH-1a infection and replication in Marc-145 cells^[2]. Cecropin P1, porcine (0-480 µg/mL, 36 h) not only displays extracellular virucidal activity against PRRSV (porcine reproductive and respiratory syndrome virus), but also exerts a potent inhibitory effect when added either before, simultaneously with, or after viral inoculation^[2]. Cecropin P1, porcine (480 µg/mL, 0-72 h) blocks CH-1a-induced apoptosis during the late phase of infection^[2]. Cecropin P1, porcine (0-480 µg/mL, 0-4 h) inhibits viral particle release^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Product Data Sheet



Cell	Cell Viability Assay ^[2]					
Cell	Line:	Marc-145 cells				
Cor	centration:	160, 320, and 480 μg/mL				
Incu	ubation Time:	36, 48, 72, 96 h				
Res	ult:	Significantly inhibited viral infection in a dose-dependent manner at 36 h postinfection. Inhibited CH-1a infection in Marc-145 cells with a 50% effective concentration (EC ₅₀) of 112 μg/mL. The 50% cytotoxic concentration (CC ₅₀) of Cecropin P1 for Marc-145 cells was estimated to be 719 μg/mL.				
Wes	Western Blot Analysis ^[2]					
Cell	Line:	Marc-145 cells				
Cor	centration:	160, 320, and 480 μg/mL				
Incu	ubation Time:	36 h				
Res	ult:	Significantly inhibited viral infection in a dose-dependent manner at 36 h postinfection. Significantly reduced the expression of the viral N protein when administered with either the pre-, co-, or posttreatment method.				

In Vivo

Cecropin P1 (1 mg/kg, IP, once) prevents bacterial growth, endotoxemia, and mortality in rats with septic shock^[3]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Andersson M, et al. Ascaris nematodes from pig and human make three antibacterial peptides: isolation of cecropin P1 and two ASABF peptides. Cell Mol Life Sci. 2003 Mar;60(3):599-606.

[2]. Guo C, et al. Cecropin P1 inhibits porcine reproductive and respiratory syndrome virus by blocking attachment. BMC Microbiol. 2014 Nov 18;14:273.

[3]. Giacometti A, et al. Effect of mono-dose intraperitoneal cecropins in experimental septic shock. Crit Care Med. 2001 Sep;29(9):1666-9.

[4]. Jiang R, et al. Expression of antimicrobial peptide Cecropin P1 in Saccharomyces cerevisiae and its antibacterial, antiviral activity in vitro. Electronic Journal of Biotechnology, 2020.

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

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