

Product Data Sheet

Urocortin III (mouse) (free acid)

Cat. No.:	HY-P3597	
Molecular Formula:	$C_{186}H_{311}N_{51}S_{2}$	
Molecular Weight:	4171.26	
Sequence Shortening:	FTLSLDVPTNIMNILFNIDKAKNLRAKAAANAQLMAQI	
Target:	CFTR	
Pathway:	Membrane Transporter/Ion Channel	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

	BIOLOGICAL ACTIVITY				
	Description	Urocortin III (mouse) (free acid) is a selective CRF2 receptor agonist (with high affinity for the CRF2 receptor). Urocortin III (mouse) (free acid) significantly inhibits gastric emptying without modifying colonic transit ^{[1][2]} .			
	IC ₅₀ & Target	CRF2 receptor ^{[1][2]} .			
	In Vivo	Urocortin III (mouse) (free acid) (120 μg/kg; i.p.; single) inhibits gastric emptying of a solid meal without modifying colonic transit in mice ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			
		Animal Model:	Adult male C57BL/6 mice (6 to 8-week-old) ^[1] .		
		Dosage:	12, 60, 120 μg/kg		
		Administration:	Intraperitoneal injection; single		
		Result:	Inhibited gastric emptying of the solid meal only at the highest dose, and did not alter distal colonic transit.		

REFERENCES

[1]. Martínez V, et al. Differential actions of peripheral corticotropin-releasing factor (CRF), urocortin II, and urocortin III on gastric emptying and colonic transit in mice: role of CRF receptor subtypes 1 and 2. J Pharmacol Exp Ther. 2002 May;301(2):611-7.

[2]. Lewis K, et al. Identification of urocortin III, an additional member of the corticotropin-releasing factor (CRF) family with high affinity for the CRF2 receptor. Proc Natl Acad Sci U S A. 2001 Jun 19;98(13):7570-5.

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

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