Porcine dynorphin A(1-13)

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®

Cat. No.:	HY-P0088				
CAS No.:	72957-38-1				
Molecular Formula:	$C_{75}H_{126}N_{24}O_{15}$	194 ₀ 3942 194 ₀ 3945			
Molecular Weight:	1603.95				
Sequence:	Tyr-Gly-Gly-Phe-Leu-Arg-Arg-Ile-Arg-Pro-Lys-Leu-Lys				
Sequence Shortening:	YGGFLRRIRPKLK				
Target:	Opioid Receptor				
Pathway:	GPCR/G Protein; Neuronal Signaling				
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:≥60 mg/mL (37.41 mM) * "≥" means soluble, but saturation unknown.				
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
		1 mM	0.6235 mL	3.1173 mL	6.2346 mL
		5 mM	0.1247 mL	0.6235 mL	1.2469 mL
		10 mM	0.0623 mL	0.3117 mL	0.6235 mL
	Please refer to the solubility information to select the appropriate solvent.				
In Vivo	1. Add each solvent Solubility: 100 mg	one by one: PBS /mL (62.35 mM); Clear solution; Need	dultrasonic		

Description	Porcine dynorphin A (1-13) is a potent, endogenous κ opioid receptor agonist and is antinociceptive at physiological concentrations.			
IC ₅₀ & Target	к Opioid Receptor/KOR			
In Vivo	Dynorphin A (1-13) exposure (33 μM) causes a significant loss in neuronal viability at 4 h with a visible destruction in neuronal morphology seen at 16 h. Exposure to dynorphin A (1-13) causes acute increases in [Ca ²⁺] _i in individual neurons similar to increases seen with acute NMDA treatment. Continuous exposure to dynorphin A (1-13) (100 μM) causes a			

significant loss of neurons over time $^{[1]}$.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Hauser KF, et al. Dynorphin A (1-13) neurotoxicity in vitro: opioid and non-opioid mechanisms in mouse spinal cord neurons. Exp Neurol. 1999 Dec;160(2):361-75.

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

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