Neurotensin

Cat. No.:	HY-P0234			
CAS No.:	39379-15-2	— Но		
Molecular Formula:	C ₇₈ H ₁₂₁ N ₂₁ O ₂₀			
Molecular Weight:	1672.92			
Sequence:	{Pyr}-Leu-Tyr-Glu-Asn-Lys-Pro-Arg-Arg-Pro-Tyr-Ile-Leu			
Sequence Shortening:	{Pyr}-LYENKPRRPYIL			
Target:	Neurotensin 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

	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
		1 mM	0.5978 mL	2.9888 mL	5.9776 mL
		5 mM	0.1196 mL	0.5978 mL	1.1955 mL
Pleas		10 mM	0.0598 mL	0.2989 mL	0.5978 mL
	Please refer to the solubility information to select the appropriate solvent.				
In Vivo	1. Add each solvent Solubility: 20 mg/	one by one: PBS mL (11.96 mM); Clear solution; Need	ultrasonic		

BIOLOGICAL ACTIVITY				
Description	Neurotensin, a gut tridecapeptide, acts as a potent cellular mitogen for various colorectal and pancreatic cancers which possess high-affinity neurotensin receptors (NTR).			
IC ₅₀ & Target	Neurotensin receptors (NTR) ^[1]			
In Vitro	Neurotensin induces the expression of MIP-2, MCP-1, IL-1β and TNFα in murine microglial cells and stimulates IL-8 secretion in a non-transformed colon epithelial cell line stably transfected with the NTR. The high-affinity NTR, a member of the G- protein coupled receptor (GPCR) family, is present in a majority of human pancreatic and colorectal cancers, suggesting that Neurotensin (NT) may act in an endocrine fashion to affect tumor growth. Acting through the NTR1, Neurotensin is known to stimulate various signal transduction pathways, including intracellular calcium ([Ca ²⁺] _i), the mitogen-activated protein			

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kinases (MAPKs), ERK and JNK, and various PKC isoforms. Treatment of HCT116 cells with Neurotensin (100 nM) significantly increases HCT116 cell migration (~3-fold) compared with vehicle treatment; pretreatment with Curcumin (10 μ M) blocksthe stimulatory effect of NT on HCT116 cell migration. ctivation of MEK/ERK by NT and downstream induction of AP-1 transcription factors contributes to the proliferative effects of Neurotensin^[1].

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

CUSTOMER VALIDATION

• J Am Soc Mass Spectrom. 2020 Jul 5.

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REFERENCES

[1]. Wang X, et al. Curcumin inhibits neurotensin-mediated interleukin-8 production and migration of HCT116 human colon cancer cells. Clin Cancer Res. 2006 Sep 15;12(18):5346-55.

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