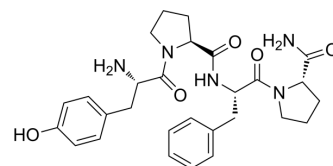


Morphiceptin

Cat. No.:	HY-P1701
CAS No.:	74135-04-9
Molecular Formula:	C ₂₈ H ₃₅ N ₅ O ₅
Molecular Weight:	521.61
Sequence Shortening:	YFPF-NH2
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

H₂O : ≥ 250 mg/mL (479.29 mM)
 DMSO : 125 mg/mL (239.64 mM; Need ultrasonic)
 * "≥" means soluble, but saturation unknown.

	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	1.9171 mL	9.5857 mL	19.1714 mL
	5 mM	0.3834 mL	1.9171 mL	3.8343 mL
	10 mM	0.1917 mL	0.9586 mL	1.9171 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- Add each solvent one by one: PBS
Solubility: 100 mg/mL (191.71 mM); Clear solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: ≥ 2.08 mg/mL (3.99 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
Solubility: ≥ 2.08 mg/mL (3.99 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 2.08 mg/mL (3.99 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Morphiceptin is a potent and specific agonist for morphine (μ) receptors. Morphiceptin, as a synthetic peptide, is the amide of a fragment of the milk protein β-casein. Morphiceptin has morphinelike activities and is highly specific for morphine (μ)

	receptors but not for enkephalin (κ) receptors ^[1] .
IC ₅₀ & Target	μ Opioid Receptor/MOR
In Vivo	Morphiceptin potently inhibits the binding of [³ H]dihydromorphine, ¹²⁵ I-labeled FK33824, and [³ H]naloxone to rat brain membrane preparations with a 50 percent inhibition concentration (IC ₅₀) of about 20 nM ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Chang KJ, et al. Morphiceptin (NH₄-tyr-pro-phe-pro-COHN₂): a potent and specific agonist for morphine (mu) receptors. Science. 1981;212(4490):75-77.

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

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