

Hemokinin 1 (mouse)

Catalog No: tcsc7597

Available Sizes

Size: 1mg

Size: 5mg

Size: 10mg

Specifications

CAS No:

208041-90-1

Formula:

C₆₁H₁₀₀N₂₂O₁₅S

Pathway: Neuronal Signaling;GPCR/G Protein

Target:

Neurokinin Receptor; Neurokinin Receptor

Purity / Grade:

Solubility:

10 mM in H2O

Observed Molecular Weight:

1413.65

Product Description

Hemokinin 1 (mouse) is a selective agonist of **neurokinin-1 receptor**, with **K**_i of 0.175 nM and 560 nM for human NK1 receptor and human NK2 receptor, respectively.

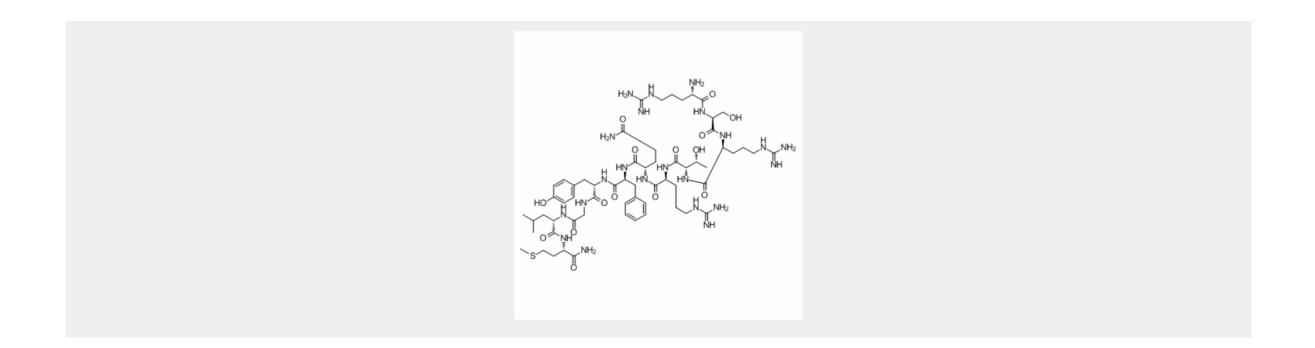
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IC50 & Target: Ki: 0.175 nM (Human NK1 receptor), 560 nM (Human NK2 receptor)^[1]

In Vitro: Hemokinin 1 (mouse) (1 nM-3 μ M) produces concentration-dependent contraction of RUB averaging 66±3% (n=6) of the maximal contraction produced by KCl (80 mM). Hemokinin 1 (mouse) (10 nM-10 μ M) induces a quickly-developing contractile responses of GPI, as does the tachykinin NK3 receptor selective agonist senktide or neurokinin B (NKB). Hemokinin 1 (mouse) induces full agonist responses but with a 500 fold lower potency as compared to NKB^[1].

In Vivo: Hemokinin 1 (mouse) (0.01-100 nmol/kg i.v., n=10) induces a dose-related hypotension that is maximal at the dose of 10 nmol/kg. For systolic blood pressure (SBP), the ED_{50} value is 0.2 nmol/kg (0.1-0.4 nmol/kg) for Hemokinin 1 (mouse). For diastolic blood pressure (DBP), the ED_{50} value is 0.1 nmol/kg (0.07-0.2 nmol/kg) for Hemokinin 1 (mouse). Hemokinin 1 (mouse) (0.1-100 nmol/kg, i.v.) induces a dose-related salivary secretion in atropine-pretreated rats^[1].



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