

Anamorelin (hydrochloride)

Catalog No: tcsc1037



Available Sizes

Size: 5mg

Size: 10mg

Size: 50mg

Size: 100mg



Specifications

CAS No:

861998-00-7

Formula:

$C_{31}H_{43}ClN_6O_3$

Pathway:

GPCR/G Protein

Target:

GHSR

Purity / Grade:

>98%

Solubility:

DMSO : ≥ 28 mg/mL (48.01 mM)

Alternative Names:

RC-1291 hydrochloride;ONO-7643 hydrochloride

Observed Molecular Weight:

583.16

Product Description

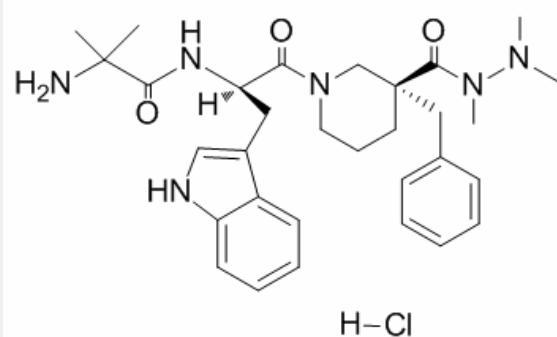
Anamorelin hydrochloride is a novel **ghrelin receptor** agonist with **EC₅₀** value of 0.74 nM in the FLIPR assay.

IC₅₀ & Target: Ki: 0.7 nM (ghrelin receptor)^[1]

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In Vitro: In the FLIPR assay, Anamorelin (ANAM) shows significant agonist activity on the ghrelin receptor, with EC₅₀ value of 0.74 nM. No significant antagonist activity is observed with Anamorelin at concentrations of up to 1,000 nM. In the binding experiments, Anamorelin binds to the ghrelin receptor with a binding affinity constant (K_i) of 0.70 nM. In the competition assay with radiolabeled ibutamoren (³⁵S-MK-677; another ghrelin receptor agonist) Anamorelin (ANAM) is also found to bind with high affinity to the ghrelin receptor (IC₅₀=0.69 nM). In rat pituitary cells incubated with Anamorelin, there is a dose-dependent stimulatory effect on GH release and the potency (EC₅₀) is 1.5 nM. Anamorelin is screened for activity against a set of over 100 receptors, ion channels, transporters, and enzymes. Anamorelin demonstrates binding to the tachykinin neurokinin 2 (NK₂) site (IC₅₀=0.021 μM); however, a subsequent NK₂ functional assay demonstrates no functional activity^[1].

In Vivo: In rats, Anamorelin (ANAM) at an oral dose of 3, 10, or 30 mg/kg once daily significantly increases both food intake and body weight from Day 2 to Day 7 of treatment compared with the vehicle control. The cumulative change in food intake and weight gain increases dose-dependently, and these changes are significant at all dose levels (P0-6h in rats)^[1].



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