

# MK 0893

Catalog No: tcsc0324



## Available Sizes

**Size:** 5mg

**Size:** 10mg

**Size:** 50mg



## Specifications

**CAS No:**

870823-12-4

**Formula:**

$C_{32}H_{27}Cl_2N_3O_4$

**Pathway:**

GPCR/G Protein

**Target:**

Glucagon Receptor

**Purity / Grade:**

>98%

**Solubility:**

H<sub>2</sub>O :

**Observed Molecular Weight:**

588.48

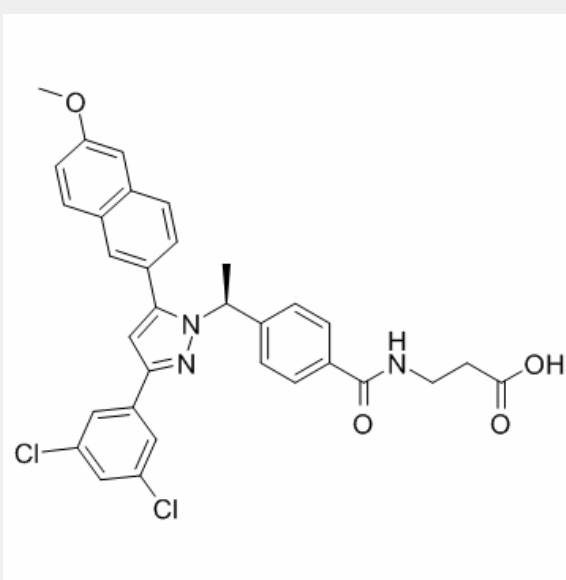
## Product Description

MK 0893 is a potent, selective **glucagon receptor** antagonist with **IC<sub>50</sub>** of 6.6 nM, and > 200 fold selectivity against GIPR, PAC1, GLP-1R, VPAC1 and VPAC2.

IC50 & Target: IC50: 6.6 nM (glucagon receptor)

**In Vitro:** MK 0893 is selective for glucagon receptor relative to other family B GPCRs, showing IC<sub>50</sub> values of 1020 nM for GIPR, 9200 nM for PAC1, and >10000 nM for GLP-1R, VPAC1, and VPAC2. MK 0893 is active against the rhesus monkey GCGR, showing an IC<sub>50</sub> of 56 nM in a cAMP assay with CHO cells expressing the rhesus GCGR<sup>[1]</sup>.

**In Vivo:** MK 0893 blunts glucagon-induced glucose elevation in hGCGR mice and rhesus monkeys. It also lowers ambient glucose levels in both acute and chronic mouse models: in hGCGR ob/ob mice it reduces glucose (AUC 0-6 h) by 32% and 39% at 3 and 10 mpk single doses, respectively. In hGCGR mice on a high fat diet, MK 0893 at 3, and 10 mpk po in feed lowers blood glucose levels by 89% and 94% at day 10, respectively, relative to the difference between the vehicle control and lean hGCGR mice<sup>[1]</sup>.



All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!