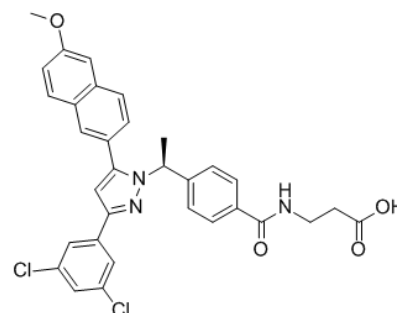


**Product Name** : MK-0893  
**Cat. No.** : PC-45831  
**CAS No.** : 870823-12-4  
**Molecular Formula** : C<sub>32</sub>H<sub>27</sub>Cl<sub>2</sub>N<sub>3</sub>O<sub>4</sub>  
**Molecular Weight** : 588.4805  
**Target** : Glucagon Receptor  
**Solubility** : 10 mM in DMSO



## Biological Activity

A potent, selective, reversible and competitive antagonist of glucagon receptor with high binding affinity (IC<sub>50</sub>=6.6 nM) and functional cAMP activity (IC<sub>50</sub>=15.7 nM); displays selectivity for glucagon receptor relative to other family B GPCRs, IC<sub>50</sub> of 1020 nM for GIPR, 9200 nM for PAC1, and >10000 nM for GLP-1R, VPAC1, and VPAC2; blunts glucagon-induced glucose elevation in hGCGR mice and rhesus monkeys, also lowers ambient glucose levels in both acute and chronic mouse models.

Diabetes

Phase 2 Discontinued

## References

Xiong Y, et al. J Med Chem. 2012 Jul 12;55(13):6137-48.

Guan HP, et al. J Lipid Res. 2015 Nov;56(11):2183-95.

Jazayeri A, et al. Nature. 2016 May 12;533(7602):274-7.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

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