

Helospectin I

Cat. No.:	HY-P3053
CAS No.:	93438-37-0
Molecular Formula:	C ₁₈₃ H ₂₉₃ N ₄₇ O ₅₉
Molecular Weight:	4095.63
Sequence Shortening:	HSDATFTAEYSKLLAKLALQKYLESilGSSTSPRPSS
Target:	Others
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

BIOLOGICAL ACTIVITY

Description	Helospectin I is a neuropeptide of the vasoactive intestinal peptide (VIP) family. Helospectin I has vasodilatory and antihypertensive activities, and decreases blood pressure. Helospectin I is originally isolated from the salivary gland venom of the lizard <i>Heloderma suspectum</i> ^{[1][2]} .															
IC₅₀ & Target	Vasoactive intestinal peptide (VIP) ^[1]															
In Vitro	<p>Helospectin I (suffusion at 0.1 nM, in bicarbonate buffer for 30 min) evokes significant, sustained and similar vasodilation in the intact hamster cheek pouch^[1].</p> <p>Helospectin I relaxes the Phenylephrine-contracted rat femoral arteries with pEC₅₀ of 6.82^[2].</p> <p>Helospectin I (0.1 nM-1 μM) inhibits the binding of 125I-labeled VIP and 125I-secretin to dispersed chief cells^[4].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>															
In Vivo	<p>Helospectin I (0.03-2 nmol/kg, 100 μL of infusion at the left jugular vein) reduces blood pressure in rats^[2].</p> <p>Helospectin I (0.1-0.8 nmol /kg, i.v.) increases plasma levels of glucagon in mice^[3].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 30%;">Animal Model:</td> <td>SD rats^[2]</td> </tr> <tr> <td>Dosage:</td> <td>0.03, 0.3, 1, 2 nM/kg</td> </tr> <tr> <td>Administration:</td> <td>100 μL of infusion at the left jugular vein, followed by washing the catheter with 100 μL saline.</td> </tr> <tr> <td>Result:</td> <td>Reduced the blood pressure, but was less effective than vasoactive intestinal peptide (VIP) in the low dose range.</td> </tr> </table> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 30%;">Animal Model:</td> <td>Mice^[3]</td> </tr> <tr> <td>Dosage:</td> <td>0.1, 0.2, 0.4, 0.8 nM /kg</td> </tr> <tr> <td>Administration:</td> <td>Intravenous injection (i.v.)</td> </tr> </table>		Animal Model:	SD rats ^[2]	Dosage:	0.03, 0.3, 1, 2 nM/kg	Administration:	100 μL of infusion at the left jugular vein, followed by washing the catheter with 100 μL saline.	Result:	Reduced the blood pressure, but was less effective than vasoactive intestinal peptide (VIP) in the low dose range.	Animal Model:	Mice ^[3]	Dosage:	0.1, 0.2, 0.4, 0.8 nM /kg	Administration:	Intravenous injection (i.v.)
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Result:	Markedly stimulated glucagon secretion, and had no direct action on insulin secretion.
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REFERENCES

- [1]. Tsueshita T, et al. Helospectin I and II evoke vasodilation in the intact peripheral microcirculation. *Peptides*. 2004 Jan;25(1):65-9.
- [2]. Grundemar L, et al. Vascular effects of helodermin, helospectin I and helospectin II: a comparison with vasoactive intestinal peptide (VIP). *Br J Pharmacol*. 1990 Mar;99(3):526-8.
- [3]. Ahrén B. Effects of helospectin I on insulin and glucagon secretion in the mouse. *Br J Pharmacol*. 1991 Apr;102(4):916-8.
- [4]. Rai A, et al. Actions of Helodermatidae venom peptides and mammalian glucagon-like peptides on gastric chief cells. *Am J Physiol*. 1993 Jul;265(1 Pt 1):G118-25.
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Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA