PRODUCT INFORMATION



Cortistatin-14 (trifluoroacetate salt)

Item No. 25074

Formal Name: L-prolyl-L-cysteinyl-L-lysyl-L-asparaginyl-L-

phenylalanyl-L-phenylalanyl-L-tryptophyl-

L-lysyl-L-threonyl-L-phenylalanyl-Lseryl-L-seryl-L-cysteinyl-L-lysine, cyclic (2→13)-disulfide, trifluoroacetate salt

 $C_{81}H_{113}N_{19}O_{19}S_2 \bullet XCF_3COOH$

FW: 1,721.0 **Purity:** ≥95%

MF:

Supplied as: A lyophilized powder

Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

Cortistatin-14 (trifluoroacetate salt) is supplied as a lyophilized powder. A stock solution may be made by dissolving the cortistatin-14 (trifluoroacetate salt) in water. The solubility of cortistatin-14 (trifluoroacetate salt) in water is approximately 1 mg/ml. We do not recommend storing the aqueous solution for more than one day.

Description

Cortistatin-14 is an endogenous neuropeptide agonist of somatostatin receptors (IC₅₀s = 0.5-18.2 nM for human SST₁-SST₅ receptors expressed in CCL39 cells).^{1,2} It is expressed in GABAergic interneurons in the cortex and hippocampus.^{2,3} Cortistatin-14 competitively inhibits hexarelin binding to human growth hormone secretagogue receptors (GHS-R) in pituitary membranes (K_i = 0.54 μM).⁴ It also activates the human Mas-related gene receptor MrgX2 expressed in HEK293 cells leading to quantifiable GTP- γ S binding to the G protein subunit $G\alpha_0$ (EC₅₀ = 1.3 μ M).⁵ Cortistatin-14 (0.01-1 μ M), similarly to somatostatin-14 (Item No. 20809), completely inhibits thyrotropin-releasing hormone (TRH; Item No. 22917), and partially inhibits vasoactive intestinal peptide (VIP; Item No. 24996), stimulation of cAMP production in GH_4 cells.³ In mice, cortistatin-14 dose-dependently (0.05-5 μ g, i.c.v.) impairs memory retention when administered after foot shock avoidance training.⁶ Cortistatin-14 (40-320 µg/kg, s.c.) dose-dependently reduces growth hormone secretion in rats by greater than 80% after 10 minutes when administered at doses of 40-320 μg/kg.⁷

References

- 1. Criado, J.R., Li, H., Jiang, X., et al. J. Neurosci. Res. 56(6), 611-619 (1999).
- 2. Spier, A.D. and de Lecea, L. Brain Res. Brain Res. Rev. 33(2-3), 228-241 (2000).
- 3. de Lecea, L., Criado, J.R., Prospero-Garcia, O., et al. Nature 381(6579), 242-245 (1996).
- 4. Deghenghi, R., Papotti, M., Ghigo, E., et al. J. Endocrinol. Invest. 24(1), RC1-RC3 (2001).
- 5. Burstein, E.S., Ott, T.R., Feddock, M., et al. Br. J. Pharmacol. 147(1), 73-82 (2006) 6. Flood, J.F., Uezu, K., and Morley, J.E. Brain Res. 775(1-2), 250-252 (1997).
- 7. Deghenghi, R., Avallone, R., Torsello, A., et al. J. Endocrinol. Invest. 24(11), RC31-RC33 (2001).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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H-Pro-Cys-Lys-Asn-Phe-Phe-Trp-Lys-Thr-Phe-

• XCF₃COOH

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