
Product Data Sheet

Product Name: Urocortin II (human) (trifluoroacetate salt)
Cat. No.: GC18231

Chemical Properties

Cas. No. 398001-88-2

Chemical Name N/A

SMILES N/A

Formula $C_{194}H_{339}N_{63}O_{54}S.XCF_3COOH$ M.Wt 4450.2

Solubility Formic Acid: 1 mg/ml Storage Store at -20°C

General tips For obtaining a higher solubility, please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT, or blue ice upon request.

Structure

Background

Urocortin II is a neuropeptide hormone and member of the corticotropin-releasing factor (CRF) family which includes mammalian CRF, urocortin, urocortin III, frog sauvagine, and piscine urotensin I. Human urocortin II shares 34, 43, and 37-40% sequence homology with rat and human CRF, human urocortin, and human urocortin III. Urocortin II increases cell shortening and accelerates relaxation of rabbit ventricular myocytes in a time- and concentration-dependent manner. In vivo, urocortin reduces arterial blood pressure in normotensive and spontaneously hypertensive rats via peripheral CRF2 receptor agonism. It induces dose-dependent tachycardia and hypotension in rats when administered at doses of 3 and 30 pmol/kg. Urocortin (10 and 20 µg/kg) also reduces the visceral pain response to colorectal distension in conscious rats and delays gastric emptying in mice.

References:

[1]. Adão, R., Santos-Ribeiro, D., Rademaker, M.T., et al. Urocortin 2 in cardiovascular

Caution: Product has not been fully validated for medical applications. For research use only.

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health and disease Drug Discov. Today 20(7), 906-914 (2015).

[2]. Gardiner, S.M., March, J.E., Kemp, P.A., et al. Regional hemodynamic actions of selective corticotropin-releasing factor type 2 receptor ligands in conscious rats J. Pharmacol. Exp. Ther. 312(1), 53-60 (2005).

[3]. Million, M., Wang, L., Wang, Y., et al. CRF2 receptor activation prevents colorectal distension induced visceral pain and spinal ERK1/2 phosphorylation in rats Gut 55(2), 172-181 (2006).

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