



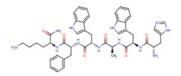
Growth hormone releasing peptide

Chemical Properties

CAS No.: 87616-84-0 Formula: C46H56N12O6

Molecular Weight: 873
Appearance: N/A

Storage: 0-4°C for short term (days to weeks), or -20°C for long term (months).



Biological Description

| Description | Growth hormone releasing peptide ia a natual product ,and has anti-inflammatory effects, due to its inhibitory effect on PKC-induced activation of p38, JNK and NF-κB, possibly by targeting to MKP-1 and PP2A. | | | | |
|----------------------------|---|--|--|--|--|
| Targets(IC ₅₀) | PP2A: None | | | | |
| In vitro | Growth hormone releasing peptide-2 (GHRP-2) attenuated phorbol 12, 13-didecanoate (PDD)-induced expression of protein and mRNA, the promoter activity of COX-2 and IL-8 genes, and the secretion of prostaglandin E2 (PGE) and IL-8. GHRP-2 promoted the degradation of PDD-induced COX-2 and IL-8 proteins with the involvement of proteasomal and lysosomal pathways. PDD-mediated COX-2 production acts via the p38, c-Jun N-terminal kinase (JNK), extracellular signal-regulated kinase (ERK) and nuclear factor kappa-light-chain-enhancer of activated B cells (NF-κB) pathways; PDD-mediated IL-8 production acts via the p38, JNK and ERK pathways. GHRP-2 reduced the PDD-induced phosphorylation of p38 and JNK and activator protein 1 (AP-1) reporter activation and PDD-induced NF-κB nuclear translocation and reporter activation. The inhibitors of mitogen-activated protein kinase phosphatase-1 (MKP-1) and protein phosphatase 2 (PP2A) reduced the inhibitory effect of GHRP-2 on PDD-induced COX-2 and IL-8 expression[1]. | | | | |
| Cell Research | KGN cells were pretreated with Growth hormone releasing peptide-2(GHRP-2)(1 M) or with GHRP-2 in combination with an inhibitor, either a MKP-1 inhibitor (sanguinarine; 0.01, 0.1, and 1 M) or a PP2A inhibitor (okadaic acid; 10, and 30 M) for 2 h before the addition of PDD for an additional 12 h. Sanguinarine at 0.1 or 1 M was able to attenuate the suppression effect of GHRP-2 on PDD-induced COX-2 expression and at 1 M, it was also able to attenuate the inhibitory effect of GHRP-2 on PDD-induced IL-8 expression[1]. | | | | |
| Animal Research | | | | | |

Solubility Information

| Solubility | < 1 mg/ml refers to the product slightly soluble or insoluble |
|------------|---|
|------------|---|

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Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|----------|----------|-----------|
| 1 mM | 1.145 mL | 5.727 mL | 11.455 mL |
| 5 mM | 0.229 mL | 1.145 mL | 2.291 mL |
| 10 mM | 0.115 mL | 0.573 mL | 1.145 mL |
| 50 mM | 0.023 mL | 0.115 mL | 0.229 mL |

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. The storage conditions and period of the stock solution: - 80 °C for 6 months; - 20 °C for 1 month. Please use it as soon as possible.

Reference

- 1. Yi-Ning C, David S, Yen-Chun P, et al. Growth Hormone Releasing Peptide-2 Attenuation of Protein Kinase C-Induced Inflammation in Human Ovarian Granulosa Cells[J]. International Journal of Molecular Sciences, 2016, 17(8):1359-.
- 2. Martínez, Rebeca, Hernández, Liz, Gil, Lázaro, et al. Growth hormone releasing peptide-6 enhanced antibody titers against subunit antigens in mice (BALB/c), tilapia (Oreochromis niloticus) and African catfish (Clarias gariepinus)[J]. Vaccine, 2017, 35(42):5722-5728.

Inhibitors · Natural Compounds · Compound Libraries

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