Screening Libraries

Product Data Sheet

Acetyl Gastric Inhibitory Peptide (human) (TFA)

Cat. No.: HY-P3580A

Molecular Formula: $\mathsf{C_{228}H_{340}N_{60}O_{67}S.C_2HF_3O_2}$

Molecular Weight: 5139.62

 $Ac-Tyr-Ala-Glu-Gly-Thr-Phe-Ile-Ser-Asp-Tyr-Ser-Ile-Ala-Met-Asp-Lys-Ile-His-Gln-Gln-As \\ {}_{Ac-YAEGTFISDYSIAMDKIHQQDFVWW,LIAQKGKKNDWKHNITQ (TFA salt)} \\$ Sequence:

p-Phe-Val-Asn-Trp-Leu-Leu-Ala-Gln-Lys-Gly-Lys-Lys-Asn-Asp-Trp-Lys-His-Asn-Ile-Thr-

Ac-YAEGTFISDYSIAMDKIHOODFVNWLLAQKGKKNDWKHNITQ Sequence Shortening:

Target: Insulin Receptor

Pathway: Protein Tyrosine Kinase/RTK

Sealed storage, away from moisture Storage:

> Powder -80°C 2 years

> > -20°C 1 year

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)

SOLVENT & SOLUBILITY

In Vitro

H₂O: 50 mg/mL (9.73 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	0.1946 mL	0.9728 mL	1.9457 mL
	5 mM	0.0389 mL	0.1946 mL	0.3891 mL
	10 mM			

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description

Acetyl Gastric Inhibitory Peptide (human) TFA is a fatty acid derivatized analog of glucose-dependent insulinotropic polypeptide with improved antihyperglycaemic and insulinotropic properties. Acetyl Gastric Inhibitory Peptide (human) TFA can be used for research of diabetes, insulin resistance and obesity^{[1][2][3]}.

In Vitro

Acetyl Gastric Inhibitory Peptide (human) TFA induces cyclic adenosine 3'5' monophosphate (cAMP) production with an EC 50 value of 1.9 nM in Chinese hamster lung fibroblast cells transfected with the human GIP receptor^[1].

Acetyl Gastric Inhibitory Peptide (human) TFA (10⁻¹³-10⁻⁸ nM) shows potent effect at stimulating insulin release compared to the native GIP in BRIN-BD11 cells^[1].

Acetyl Gastric Inhibitory Peptide (human) TFA improves glucose intolerance, type 2 diabetes, beta-cell glucose insensitivity, insulin resistance and reduced insulin secretion^[2].

Acetyl Gastric Inhibitory Peptide (human) TFA has metabolic stability and hypoglycemic and insulin modulating activities of two fatty acid derivatized N-terminally acetylated GIP analogs were evaluated in in vitro and in vivo^[3].

	MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
In Vivo	Acetyl Gastric Inhibitory Peptide (human) TFA (25 nmol/kg; i.p.; single dose) shows resistance to plasma dipeptidylpeptidase IV degradation, resulting in enhanced biological activity and improved antidiabetic potential in vivo ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

REFERENCES

- [1]. O'Harte FP, et al. Improved stability, insulin-releasing activity and antidiabetic potential of two novel N-terminal analogues of gastric inhibitory polypeptide: N-acetyl-GIP and pGlu-GIP. Diabetologia. 2002 Sep;45(9):1281-91.
- [2]. Gault Victor A, et al. GIP peptide analogues for treatment of diabetes, insulin resistance and obesity: World Intellectual Property Organization, WO2005082928[P]. 2005-12-01.
- [3]. O'Harte, et al. Analogs of gastric inhibitory polypeptide as a treatment for age related decreased pancreatic beta cell function: World Intellectual Property Organization, WO2007028632[P].2007-03-15.

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

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