

GIP (1-30) amide, porcine TFA

Cat. No.:	HY-P2541A
Molecular Formula:	C ₁₆₄ H ₂₄₆ F ₃ N ₄₁ O ₄₉ S
Molecular Weight:	3665.02
Sequence:	Tyr-Ala-Glu-Gly-Thr-Phe-Ile-Ser-Asp-Tyr-Ser-Ile-Ala-Met-Asp-Lys-Ile-Arg-Gln-Gln-Asp-Phe-Val-Asn-Trp-Leu-Leu-Ala-Gln-Lys-NH ₂ <small>YAEGTFISDYSIAMDKIRQQDFVNWLLAQK-NH₂ (TFA salt)</small>
Sequence Shortening:	YAEGTFISDYSIAMDKIRQQDFVNWLLAQK-NH ₂
Target:	Insulin Receptor
Pathway:	Protein Tyrosine Kinase/RTK
Storage:	Sealed storage, away from moisture and light Powder -80°C 2 years -20°C 1 year * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)

SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (27.28 mM; Need ultrasonic)																			
	H ₂ O : < 0.1 mg/mL (ultrasonic) (insoluble)																			
	<table border="1"> <thead> <tr> <th rowspan="2">Concentration</th> <th colspan="3">Mass</th> </tr> <tr> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> </thead> <tbody> <tr> <td>1 mM</td> <td>0.2728 mL</td> <td>1.3642 mL</td> <td>2.7285 mL</td> </tr> <tr> <td>5 mM</td> <td>0.0546 mL</td> <td>0.2728 mL</td> <td>0.5457 mL</td> </tr> <tr> <td>10 mM</td> <td>0.0273 mL</td> <td>0.1364 mL</td> <td>0.2728 mL</td> </tr> </tbody> </table>	Concentration	Mass			1 mg	5 mg	10 mg	1 mM	0.2728 mL	1.3642 mL	2.7285 mL	5 mM	0.0546 mL	0.2728 mL	0.5457 mL	10 mM	0.0273 mL	0.1364 mL	0.2728 mL
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Please refer to the solubility information to select the appropriate solvent.																				
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (0.68 mM); Suspended solution; Need ultrasonic Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (0.68 mM); Clear solution 																			

BIOLOGICAL ACTIVITY

Description	GIP (1-30) amide, porcine TFA is a full glucose-dependent insulinotropic polypeptide (GIP) receptor agonist with high affinity equal to native GIP(1-42) ^[1] . GIP (1-30) amide, porcine is a weak inhibitor of gastric acid secretion and potent stimulator of insulin.
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REFERENCES

[1]. L S Hansen, et al. N-terminally and C-terminally truncated forms of glucose-dependent insulinotropic polypeptide are high-affinity competitive antagonists of the human GIP receptor. Br J Pharmacol. 2016 Mar;173(5):826-38.

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

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