

GLP-1(7-37) acetate

Cat. No.:	HY-P0055A												
CAS No.:	1450806-98-0												
Molecular Formula:	C ₁₅₃ H ₂₃₂ N ₄₀ O ₄₉												
Molecular Weight:	3415.72												
Sequence:	His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-Gly												
Sequence Shortening:	HAEGTFTSDVSSYLEGQAAKEFIAWLKGRG-R-G												
Target:	Glucagon Receptor												
Pathway:	GPCR/G Protein												
Storage:	Protect from light												
	<table border="0"> <tr> <td>Powder</td> <td>-80°C</td> <td>2 years</td> </tr> <tr> <td></td> <td>-20°C</td> <td>1 year</td> </tr> <tr> <td>In solvent</td> <td>-80°C</td> <td>6 months</td> </tr> <tr> <td></td> <td>-20°C</td> <td>1 month</td> </tr> </table>	Powder	-80°C	2 years		-20°C	1 year	In solvent	-80°C	6 months		-20°C	1 month
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HAEGTFTSDVSSYLEGQAAKEFIAWLKGRG



BIOLOGICAL ACTIVITY

Description	GLP-1(7-37) acetate is an intestinal insulinotropic hormone that augments glucose induced insulin secretion ^[1] .																
In Vivo	<p>GLP-1(7-37) (0.5, 5 or 50 pmol/min/kg) infused during the second hour of a 2-hour 11-mM hyperglycemic clamp produces a dose-related enhancement of the glucose-stimulated increase in plasma insulin concentration and an increased rate of glucose infusion in rats^[2].</p> <p>Infusion of GLP-1(7-37) (5 pmol/min/kg) from 1 hour through 7 hours produces a sustained increase in plasma insulin concentration relative to levels in rats infused with vehicle in rats with maintained glucose concentration at 11 mM^[2].</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Male Sprague-Dawley rats weighing 300 to 350 g with glucose IV at a variable rate for 7 hours to maintain plasma glucose concentration at 11 mM^[2].</td> </tr> <tr> <td>Dosage:</td> <td>5 pmol/min/kg.</td> </tr> <tr> <td>Administration:</td> <td>IV from 1 hour through 7 hours^[2].</td> </tr> <tr> <td>Result:</td> <td>Produced a sustained increase in plasma insulin concentration relative to levels in rats infused with vehicle.</td> </tr> </table> <table border="1"> <tr> <td>Animal Model:</td> <td>Male Sprague-Dawley rats weighing 300 to 350 g with maintained plasma glucose concentration at 11 mM^[2].</td> </tr> <tr> <td>Dosage:</td> <td>0.5, 5 or 50 pmol/min/kg.</td> </tr> <tr> <td>Administration:</td> <td>IV during the second hour of a 2-hour 11-mmol/L hyperglycemic clamp.</td> </tr> <tr> <td>Result:</td> <td>Produced a dose-related enhancement of the glucose-stimulated increase in plasma</td> </tr> </table>	Animal Model:	Male Sprague-Dawley rats weighing 300 to 350 g with glucose IV at a variable rate for 7 hours to maintain plasma glucose concentration at 11 mM ^[2] .	Dosage:	5 pmol/min/kg.	Administration:	IV from 1 hour through 7 hours ^[2] .	Result:	Produced a sustained increase in plasma insulin concentration relative to levels in rats infused with vehicle.	Animal Model:	Male Sprague-Dawley rats weighing 300 to 350 g with maintained plasma glucose concentration at 11 mM ^[2] .	Dosage:	0.5, 5 or 50 pmol/min/kg.	Administration:	IV during the second hour of a 2-hour 11-mmol/L hyperglycemic clamp.	Result:	Produced a dose-related enhancement of the glucose-stimulated increase in plasma
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insulin concentration and an increased rate of glucose infusion.

REFERENCES

- [1]. Sarrauste de Menthiere, C. et al. Structural requirements of the N-terminal region of GLP-1-[7-37]-NH₂ for receptor interaction and cAMP production. *European journal of medicinal chemistry* 39, 473-480, doi:10.1016/j.ejmech.2004.02.002 (2004).
- [2]. Hargrove DM, et al. Glucose-dependent action of glucagon-like peptide-1 (7-37) in vivo during short- or long-term administration. *Metabolism*. 1995 Sep;44(9):1231-7.
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