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GPR10 agonist 1

Cat. No.:	HY-P10378
Molecular Formula:	C ₂₀₀ H ₃₂₄ N ₅₈ O ₅₇ S ₂
Molecular Weight:	4517.2
Sequence:	Ser-Arg-Ala-His-Gln-Cys-Ser-{Nle}-Glu-Thr-Arg-Thr-Cys-Asp-Ile-Asn-Pro-Ala-Trp-Tyr-Thr-Gly-{hArg}-Gly-Ile-Arg-Pro-Val-Gly-Arg-Phe-NH ₂ (Disulfide bridge: Cys6-Cys13; multiple ethylene glycol-fatty acid; hArg=Homoarginine)
Sequence Shortening:	SRAHQCS-{Nle}-ETRTCDINPAWYTG-{hArg}-GIRPVGRF-NH ₂ (Disulfide bridge: Cys6-Cys13; multiple ethylene glycol-fatty acid; hArg=Homoarginine)
Target:	GnRH Receptor
Pathway:	GPCR/G Protein
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

BIOLOGICAL ACTIVITY

Description	GPR10 agonist 1 (compound 18-S4) is a potent GPR10 agonist with EC ₅₀ values of 80, 7.8 nM in the presence (10%) or absence (0%) of FBS, respectively. GPR10 agonist 1 has the potential for the research of chronic obesity ^[1] .				
In Vivo	GPR10 agonist 1 (compound 18-S4) (0.5, 5 mg/kg; s.c.; daily for 12 day) decreases diet-induced obesity (DIO) mouse body weight ^[1] . Pharmacokinetic Parameters (1 mg/kg; s.c.; 7-9 weeks, male C57 mice) ^[1] .				
	t _{1/2} (h)	C _{max} (ng/mL)	AUC _{last} (ng·h/mL)	AUC _∞ (ng·h/mL)	AUC _{0-24h} (ng·h/mL)
	7.81	1670	17700	17900	15500
	MCE has not independently confirmed the accuracy of these methods. They are for reference only.				
	Animal Model:	24 weeks male mice (a diet-induced obesity (DIO) mouse model) ^[1]			
	Dosage:	0.5, 5 mg/kg			
	Administration:	S.c.; daily for 12 daily			
	Result:	Significantly decreased the body weight at 0.5 mg/kg and 5 mg/kg daily injection gave similar efficacy to the 0.5 mg/kg dose.			

REFERENCES

[1]. Pflimlin E, et al. Design of a Long-Acting and Selective MEG-Fatty Acid Stapled Prolactin-Releasing Peptide Analog. ACS Med Chem Lett. 2019 Jul 5;10(8):1166-1172.

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

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