

Product Data Sheet

His-{Aib}-Gln-Gly-Thr-Phe-Thr-Ser-Asp-

Ala-Arg-Glu-Phe-Val-Gln-Trp-Leu-Leu-{Lys(γGlu-γGlu-Ser-Glu-Ser-γGlu-γGlu-

C₁₈ diacid)}-Thr-NH₂ (TFA)

NN1177 TFA

Cat. No.: HY-P10032A

Molecular Formula: $C_{206}H_{323}N_{51}O_{66}.xC_2HF_3O_2$

Sequence: His-{Aib}-Gln-Gly-Thr-Phe-Thr-Ser-Asp-Leu-Ser-Lys-Tyr-Leu-Glu-Ser-Lys-Arg-Ala-Arg-

NH2

Sequence Shortening: H-{Aib}-QGTFTSDLSKYLESKRAREFVQWLL-{Lys(\gammaGlu-\gammaGlu-Ser-\gammaGlu-\g

diacid)}-T-NH2

Target: GCGR; Cytochrome P450; GLP Receptor

Pathway: GPCR/G Protein; Metabolic Enzyme/Protease

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 H₂O: 100 mg/mL (Need ultrasonic)

BIOLOGICAL ACTIVITY

Description	NN1177 (NNC9204-1177) TFA is a long-acting GLP-1/glucagon receptor co-agonist. NN1177 TFA can induce a dose-dependent body weight loss in diet-induced obese (DIO) mice $^{[1][2]}$.
IC ₅₀ & Target	CYP3A4
In Vitro	NN1177 (100 nM, 3 days) TFA reduces CYP3A4 mRNA expression (57.2-71.7%) and activity (18.5-51.5%) in freshly isolated human hepatocytes ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	NN1177 (3 or 5 nmol/kg, s.c.) TFA induces body weight loss, loss of fat mass, and improvement in glucose tolerance in dietinduced obese (DIO) mice ^[1] . NN1177 (0.75-4 nmol/kg, s.c., once daily, 8 weeks) TFA reduces liver fat and inflammatory and fibrosis relevant biomarkers in C57Bl/6 mice fed a fructose and high fat rich diet (NASH model) ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Simonsen L, et al. Preclinical evaluation of a protracted GLP-1/glucagon receptor co-agonist: Translational difficulties and pitfalls. PLoS One. 2022 Mar 4;17(3):e0264974.

[2]. Monfeuga T, et al. Evaluation of long acting GLP1R/GCGR agonist in a DIO and biopsy-confirmed mouse model of NASH suggest a beneficial role of GLP-1/glucagon

agonism in NASH patients. Mol Metab. 2023 Dec 7;79:101850. [3]. Säll C, et al. In vitro CYP450 enzyme down-regulation by GLP-1/glucagon co-agonist does not translate to observed drug-drug interactions in the clinic. Drug Metab Dispos. 2022 Jun 9:DMD-AR-2022-000865. Caution: Product has not been fully validated for medical applications. For research use only. Fax: 609-228-5909 Tel: 609-228-6898 E-mail: tech@MedChemExpress.com Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA

Page 2 of 2 www.MedChemExpress.com