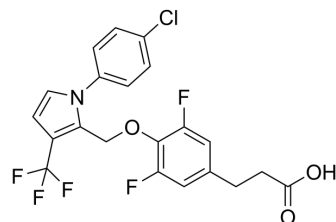


GPR120 Agonist 4

Cat. No.:	HY-160628
CAS No.:	1628641-89-3
Molecular Formula:	C ₂₁ H ₁₅ ClF ₅ NO ₃
Molecular Weight:	459.79
Target:	Free Fatty Acid Receptor
Pathway:	GPCR/G Protein
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	GPR120 Agonist 4 (example 1) is a GPR120 agonist with the EC ₅₀ values of 1 μM and 0.35 μM for β-arrestin A and Calcium A. GPR120 Agonist 4 can be used for the research of type II diabetes mellitus ^[1] .
IC₅₀ & Target	GPR120 1 μM (EC50)
In Vitro	GPR120 Agonist 4 (example 1) (0.012-25 μM, 90 min) can activate GPR120 through recruitment of β-Arrestin in CHO-K1 GPR120 β-Arrestin cells ^[1] . GPR120 Agonist 4 (0.012-25 μM, 90 min) promotes the release of Ca ²⁺ in HEK293 with human GPR120 clone, which verifies the activation effect of GPR120 Agonist 4 on GPR120 ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	GPR120 Agonist 4 (example 1) (0.2/0.3/1/3/10 mg/kg, i.p., 15/30/45/60/90 min) dose-dependently promotes a decrease in glucose levels of the high fat diet mice model ^[1] . GPR120 Agonist 4 (1/3/10 mg/kg, i.p., 15/30/45/60/90 min) inhibits the glucose level of starvation mouse model in a dose-dependent manner ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. SUI, Zhihua, et al. Bicyclic pyrrole derivatives useful as agonists of gpr120. WO2015134038. 2014-03-07

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA