(R)-Fasiglifam

MedChemExpress

| Cat. No.: | HY-10480A | | |
|--------------------|--|-------------|----------|
| CAS No.: | 1234474-57 | ' -7 | |
| Molecular Formula: | C ₂₉ H ₃₂ O ₇ S | | |
| Molecular Weight: | 524.63 | | |
| Target: | Others | | |
| Pathway: | Others | | |
| Storage: | Powder | -20°C | 3 years |
| | | 4°C | 2 years |
| | In solvent | -80°C | 6 months |
| | | -20°C | 1 month |

SOLVENT & SOLUBILITY

| | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg |
|------------------------------|-------------------------------|-----------|-----------|------------|
| Preparing Stock Solutions | 1 mM | 1.9061 mL | 9.5305 mL | 19.0611 mL |
| | 5 mM | 0.3812 mL | 1.9061 mL | 3.8122 mL |
| | 10 mM | 0.1906 mL | 0.9531 mL | 1.9061 mL |

BIOLOGICAL ACTIVITY Description (R)-Fasiglifam is the isomer of Fasiglifam (HY-10480), and can be used as an experimental control. Fasiglifam (TAK-875) is a potent, selective and orally bioavailable GPR40 agonist with EC₅₀ of 72 nM.

REFERENCES

[1]. Nagatake T, et al. 17,18-EpETE-GPR40 axis ameliorates contact hypersensitivity by inhibiting neutrophil mobility in mice and cynomolgus macaques. J Allergy Clin Immunol. 2017 Dec 26. pii: S0091-6749(17)32949-4.

[2]. Yoshiyuki Tsujihata, et al. TAK-875, an Orally Available GPR40/FFA1 Agonist Enhances Glucose-Dependent Insulin Secretion and Improves Both Postprandial and Fasting Hyperglycemia in Type 2 Diabetic Rats. JPET July 13, 2011

[3]. Tsujihata Y,et al. TAK-875, an orally available G protein-coupled receptor 40/free fatty acid receptor 1 agonist, enhances glucose-dependent insulin secretion and improves both postprandial and fasting hyperglycemia in type 2 diabetic rats. J Pharmacol Exp

Product Data Sheet

[4]. Urano Y, et al. Comparative hepatic transcriptome analyses revealed possible pathogenic mechanisms of fasiglifam (TAK-875)-induced acute liver injury in mice. Chem Biol Interact. 2018 Sep 20;296:185-197.

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

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