

TUG-891

Chemical Properties

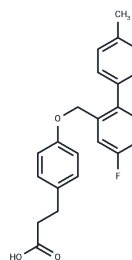
CAS No. : 1374516-07-0

Formula: C₂₃H₂₁FO₃

Molecular Weight: 364.41

Appearance: no data available

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year



Biological Description

| | |
|---------------|--|
| Description | TUG-891 is a G protein-coupled receptor (GPCR) expressed in intestine, adipocytes, and pro-inflammatory macrophages that is activated by long chain free fatty acids. |
| Targets(IC50) | GPCR |
| In vitro | TUG-891 exhibits analogous signaling characteristics to the long-chain fatty acid (LCFA) α -linolenic acid at the human Free Fatty Acid receptor 4 (FFA4), stimulating Ca ²⁺ mobilization, recruiting β -arrestin-1 and β -arrestin-2, and promoting extracellular signal-regulated kinase phosphorylation. Moreover, TUG-891 activation of FFA4 leads to the receptor's swift phosphorylation and internalization[1]. |
| Kinase Assay | Inhibition of recombinant human Mps1 by BAY 1161909 or BAY 1217389 is assessed in TRFRET-based in vitro kinase assays via phosphorylation of a biotinylated peptide (Biotin-Ahx-PWDPDDADITEILG-NH ₂). Under standard assay conditions kinase and test compound are preincubated for 15 min before enzyme reaction is started by addition of substrate and ATP upon 10 μ M[1]. |

Solubility Information

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|------------|---|
| Solubility | DMSO: 40 mg/mL (109.77 mM), Sonication is recommended. H ₂ O: < 1 mg/mL (insoluble or slightly soluble), (< 1 mg/ml refers to the product slightly soluble or insoluble) |
|------------|---|

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|------------|------------|
| 1 mM | 2.7442 mL | 13.7208 mL | 27.4416 mL |
| 5 mM | 0.5488 mL | 2.7442 mL | 5.4883 mL |
| 10 mM | 0.2744 mL | 1.3721 mL | 2.7442 mL |
| 50 mM | 0.0549 mL | 0.2744 mL | 0.5488 mL |

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

Hudson BD, et al. The pharmacology of TUG-891, a potent and selective agonist of the free fatty acid receptor 4 (FFA4/GPR120), demonstrates both potential opportunity and possible challenges to therapeutic agonism. Mol Pharmacol. 2013 Nov;84(5):710-25.

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