
Product Data Sheet

Product Name: Oleoyl Serotonin

Cat. No.: GC11147

Chemical Properties

Cas. No. 1002100-44-8

Chemical Name N-[2-(5-hydroxy-1H-indol-3-yl)ethyl]-9Z-octadecenamide

SMILES O=C(NCCC1=CNC2=CC=C(O)C=C12)CCCCCCC/C=C\CCCCCCCC

Formula $C_{28}H_{44}N_2O_2$ M.Wt 440.1

Solubility $\leq 15\text{mg/ml}$ in DMSO; 15mg/ml in dimethyl formamide Storage Store at -20°C

General tips For obtaining a higher solubility, please warm the tube at 37°C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution: ship with blue ice All other available size: ship with RT, or blue ice upon request.

Structure

Background

Oleoyl Serotonin is a TRPV1 antagonist with IC_{50} value of $2.57\ \mu\text{M}$ for human TRPV1 [1].

The transient receptor potential vanilloid-type 1 (TRPV1) channel is a nonselective cation channel that may be activated by a variety of exogenous and endogenous physical and chemical stimuli. TRPV1 is decreased in the injured nerve fibers but increased in those proximal to the site damage. TRPV1 is a potential new target for the development of analgesic and anti-inflammatory drugs [1].

Oleoyl serotonin is a hybrid molecule patterned after arachidonoyl serotonin. Arachidonoyl serotonin is a dual antagonist of TRPV1 and fatty acid amide hydrolase (FAAH) with IC_{50} values of 0.27 and $8\ \mu\text{M}$, respectively. Arachidonoyl serotonin was highly effective against both acute and chronic peripheral pain [1][2]. In TRPV1 and FAAH assays, Oleoyl serotonin inhibited anandamide hydrolysis by FAAH and capsaicin-induced intracellular Ca^{2+} elevation in HEK293 cells overexpressing the human recombinant TRPV1 receptor with IC_{50} values of $> 50\ \mu\text{M}$ and $2.57\ \mu\text{M}$, respectively [1].

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

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References:

- [1]. Ortar G, Cascio MG, De Petrocellis L, et al. New N-arachidonoylserotonin analogues with potential "dual" mechanism of action against pain. J Med Chem. 2007 Dec 27;50(26):6554-69.
- [2]. Maione S, De Petrocellis L, de Novellis V, et al. Analgesic actions of N-arachidonoylserotonin, a fatty acid amide hydrolase inhibitor with antagonistic activity at vanilloid TRPV1 receptors. Br J Pharmacol. 2007 Mar;150(6):766-81.

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