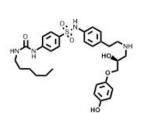


CRISPR Editing Enhancer - L755,507

Chemical Name: (S)-4-(3-hexylureido)-N-(4-(2-((2-hydroxy-3-(4-hydroxyphenoxy)propyl)amino)ethyl)phenyl)benzenesulfonamide



Molecular Weight:	584.73
Formula:	$C_{30}H_{40}N_4O_6S$
Purity:	≥98%
CAS#:	159182-43-1
Solubility:	DMSO up to 100 mM
Storage	Powder: 4 °C 1 year
	DMSO: 4 °C 3 months
	-20 °C 1 year

Biological Activity:

L-755,507 was previously characterized as a potent and selective β 3 adrenergic receptor partial agonist with EC₅₀~0.43 nM. It has > 1000 fold selectivity over β 1- and β 2-adrenoceptors (EC₅₀~580 nM and >10000 nM for β 1- and β 2-adrenoceptors respectively). It stimulates lipolysis in rhesus adipocytes in vitro (EC₅₀ = 3.9 nM). In a recent study, L-755,507 was identified to enhance CRISPR-mediated homology-directed repair (HDR) efficiency in human induced pluripotent stem cells (iPSCs) and other cell types.

How to Use:

In vitro: L-755,507 was used at 1-5 µM final concentration in various in vitro assays.

In vivo: L-755,507 stimulates metabolic rate by 30% after acute bolus intravenous administration of 0.1 mg/kg to rhesus monkeys.

Reference:

- 1. Fisher MH, et al. A selective human beta3 adrenergic receptor agonist increases metabolic rate in rhesus monkeys. (1998) J Clin Invest.101(11):2387-93.
- 2. Parmee ER, et al. Discovery of L-755,507: a subnanomolar human beta 3 adrenergic receptor agonist. (1998) Bioorg Med Chem Lett. 8(9):1107-12.
- 3. Yu C, et al. Small Molecules Enhance CRISPR Genome Editing in Pluripotent Stem Cells. (2015) Cell Stem Cell 16(2):142-7.

Products are for research use only. Not for human use.