BioVision

05/16

RELATED PRODUCTS:

FAAH Inhibitor, PF-622 (1928)

JNJ-1661010 (2413) EZSolution™ MFAP (2811)

PRODUCT:

JZL195

ALTERNATE NAME: 4-[(3-Phenoxyphenyl)methyl]-1-piperazinecarboxylic acid 4nitrophenyl ester

CATALOG #:

AMOUNT:

B1064-5, 25

STRUCTURE:



MOLECULAR FORMULA:	$C_{24}H_{23}N_3O_5$
MOLECULAR WEIGHT:	433.46
CAS NUMBER:	1210004-12-8
APPEARANCE:	White to off-white solid
SOLUBILITY:	
SOLUBILITT.	DMSO (>20 mg/ml)
PURITY:	≥98% by HPLC

STORAGE: Store at -20 °C. Protect from moisture

DESCRIPTION: JZL195 is a potent dual inhibitor of Monoacylglycerol lipase (MAGL) (IC₅₀ = 2 nM) and fatty acid amide hydrolase (FAAH) $(IC_{50} = 4 \text{ nM})$, enzymes that degrade the endocannabinoids 2arachidonoylglycerol (2-AG) and anandamide (AEA), the endogenous ligands for the cannabinoid G-protein coupled receptors CB1 and CB2. It poorly inhibits neuropathy target esterase and ABHD6 and does not inhibit other brain serine hydrolases. JZL 195 displays time-dependent inhibition of FAAH and MAGL in vivo, consistent with a covalent mechanism of activation.

REFERENCE: Long, J.Z., et al. (2009). Proc. Natl. Acad. Sci. USA 106, 20270-20275.

HANDLING: Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of exposure.

USAGE:

FOR RESEARCH USE ONLY! Not to be used in humans