



YM-201636

Catalog No: tcsc0592

Available Sizes
Size: 2mg
Size: 5mg
Size: 10mg
Size: 50mg
Size: 100mg
Specifications
CAS No: 371942-69-7
Formula: C ₂₅ H ₂₁ N ₇ O ₃
Pathway: PI3K/Akt/mTOR;Autophagy;PI3K/Akt/mTOR
Target: PI3K;Autophagy;PIKfyve
Purity / Grade: >98%
Solubility: DMSO : ≥ 47 mg/mL (100.54 mM)
Observed Molecular Weight: 467.48





Product Description

YM-201636 is a potent and selective **PIKfyve** inhibitor with an IC_{50} of 33 nM. YM-201636 also inhibits p110 α with IC_{50} of 3.3 μ M.

IC50 & Target: IC50: 33 nM (PIKfyve), 3.3 μ M (p110 α)^[1]

In Vitro: Acute treatment of cells with YM-201636 shows that the PIKfyve pathway is involved in the sorting of endosomal transport, with inhibition leading to the accumulation of a late endosomal compartment and blockade of retroviral exit. The yeast orthologue of PIKfyve, Fab1, is found to be insensitive to YM-201636 (IC₅₀>5 μM). YM-201636 does not inhibit a type IIγ PtdInsP kinase even at 10 μM and inhibits a mouse type Iα PtdInsP kinase with an IC₅₀>2 μM^[1]. YM-201636 almost completely inhibits basal and insulinactivated 2-deoxyglucose uptake at doses as low as 160 nM, with IC₅₀=54 nM for the net insulin response. YM-201636 also completely inhibits insulin-dependent activation of class IA PI 3-kinase^[2]. At low doses (10-25 nM), YM-201636 inhibits preferentially PtdIns5P rather than PtdIns(3,5)P2 production, whereas at higher doses, the two products are similarly inhibited. YM-201636 at 160 nM inhibits PtdIns5P synthesis twice more effectively compared with PtdIns(3,5)P2 synthesis^[3]. MDCK cells treated with YM-201636 accumulate the tight junction protein claudin-1 intracellularly. YM-201636 treatment blocks the continuous recycling of claudin-1/claudin-2 and delays epithelial barrier formation^[4].

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!