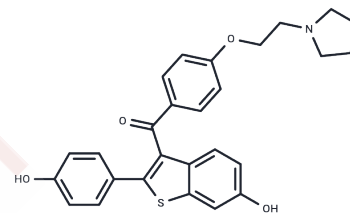


LY117018

Chemical Properties

CAS No. : 63676-25-5
 Formula: C₂₇H₂₅NO₄S
 Molecular Weight: 459.56
 Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year
 Actual storage temperature shall be subject to the COA.



Biological Description

Description	LY117018 shows antiproliferative effects on breast cancer cell lines. LY117018 is a Raloxifene analog and is a selective estrogen receptor modulator.
Targets(IC50)	Estrogen Receptor/ERR,PI3K
In vitro	LY117018 (1 μM) treatment, caused a predominantly hypophosphorylated pRb. At a higher concentration of LY117018 (1 μM), the level of p53 appeared to decline. LY117018 (0.01-1000 nM; 24 hours) at lower concentrations (0.01-10 nM) caused an E2-like increase in p53 levels when compared to its effects on cells grown in the stripped medium. LY117018 did not block E2-induced pRb phosphorylation, at lower concentrations[1]. LY117018 (1 μM; 96 hours) suppresses MCF-7 cell proliferation with an IC50 of 1 μM[2]. LY117018 inhibits oxidative stress-induced endothelial cell apoptosis through activation of ERK1/2 signaling pathway[3].

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.176 mL	10.880 mL	21.7599 mL
5 mM	0.4352 mL	2.176 mL	4.352 mL
10 mM	0.2176 mL	1.088 mL	2.176 mL
50 mM	0.0435 mL	0.2176 mL	0.4352 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.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Dinda S, et al. Effects of LY117018 (a SERM analog of raloxifene) on tumor suppressor proteins and proliferation of breast cancer cells. *Horm Mol Biol Clin Investig.* 2010 Aug 1;2(1):211-7.

Baumann KH, et al. Effects of celecoxib and ly117018 combination on human breast cancer cells in vitro. *Breast Cancer (Auckl).* 2009 Apr 7;3:23-34.

Yu J, et al. Raloxifene analogue LY117018 suppresses oxidative stress-induced endothelial cell apoptosis through activation of ERK1/2 signaling pathway. *Eur J Pharmacol.* 2008 Jul 28;589(1-3):32-6.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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