

Rapamycin (CAS: 53123-88-9)

Catalog #: EBC51016

Biological Activity

Chemical Name

Synonyms AY 22989, Sirolimus, NSC-2260804

3S,6R,7E,9R,10R,12R,14S,15E,17E,19E,21S,23S,26R,27R,34aS)-

9,10,12,13,14,21,22,23,24,25,26,27,32,33,34,34a-Hexadecahydro-9,27-dihydroxy-3-[(1R)-2-[(1S,3R,4R)-4-hydroxy-3-methoxycyclohexyl]-1-methylethyl]-10,21-dimethoxy-

6,8,12,14,20,26-hexamethyl-23,27-epoxy-3*H*-pyrido[2,1-

c][1,4]oxaazacyclohentriacontine-1,5,11,28,29(4H,6H,31H)-pentone

Application Rapamycin is a potent and specific mTOR inhibitor with an IC50 of 0.1 nM in HEK293 cells

CAS No. 53123-88-9Purity $\geq 98.0\%$ Molecular Weight 914.17Molecular Formula $C_{51}H_{79}NO_{13}$ Shipping Gel Pack

Storage Store at -20° C

Target & IC_{50} mTOR: IC50 = 0.1 nM

Molecular Structure

Solubility

DMSO: 100 mg/mL (109.39 mM) Ethanol: 100 mg/mL (109.39 mM)

PS: < 1 mg/ml refers to the product insoluble

Description

Rapamycin is a member of the macrolide immunosuppressant family and a FRAP inhibitor. Rapamycin exhibits binding and inhibitory actions to the FK506 binding protein (FKBP5) proline rotamase via simultaneous binding by FKBP12 and FRAP. FRAP (RAFT1) proteins exhibit homology to PI 4- and PI 3-kinases, which have PI 4-kinase and autophosphorylating activities. The rapamycin/FKBP complex does not inhibit the FRAP PI 4-kinase activity, but does inhibit FRAP autophosphorylation. Rapamycin is unique in its ability to inhibit lymphokine induced cell proliferation at the G1 and S phase as well as an irreversible cellular arrest at the G1 phase in Scerevisiae cells. Rapamycin also exhibits selective signal blocking leading to the activation of p70/85 S6 kinase, which is potentially due to the inhibition of FRAP autophosphorylation or protein kinase activity. Rapamycin also exhibits Angiogenesis inhibition, possibly through the inhibition of the Akt pathway. Rapamycin is an inhibitor of mTOR. Rapamycin is also known as Rapamune, 23,27-Epoxy-3H-pyrido[2,1-c][1,4]oxaazacyclohentriacontine, AY 22989, and mTOR Inhibitor I

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