

Rapamycin

Catalog Number: 5318893

RUO: For Research Use Only. Not for use in diagnostic procedures.

Product Information

Synonyms: Sirolimus, Rapamune, AY 22989

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

 $9,10,12,13,14,21,22,23,24,25,26,27,32,33,34,\ 34a-hexade cahydro-9,27-dihydroxy-3-[(1R)-2[(1S,3R,4R)-4-hydroxy-3-methoxycyclohexyl]-1-methylethyl]-10,21-dimethoxy6,8,12,14,20,26-hexamethyl-23,27-epoxy-3H-pyrido[2,1-c][1,4]$

oxaazacyclohentriacontine1,5,11,28,29 (4H,6H,31H)-pentone

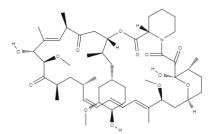
Molecular Formula: $C_{51}H_{79}NO_{13}$

Molecular Weight: 914.2 **CAS Number:** 53123-88-9

Purity: ≥95% **Applications:** FA

Formulation: Crystalline solid

Storage: Product should be kept at -20°C.



Description

Rapamycin was first evaluated as an antifungal agent, but is now studied for its potent immunosuppressive and antiproliferative properties. It forms a complex with FKBP12 and directly binds to mTORC1, inhibiting the functions of the mTOR pathway. Rapamycin is reported to block protein synthesis, arrest cell cycle progression, and inhibit IL-2 signal transduction mechanisms.

Preparation & Storage

Soluble in organic solvents such as ethanol or DMSO. DMSO up to 10mg/ml.

References

- 1.Nourse, J., Firpo, E., Flanagan, W. M., Coats, S., Polyak, K., Lee, M. H., ... Roberts, J. M. (1994). Interleukin-2-mediated elimination of the p27Kipl cyclin-dependent kinase inhibitor prevented by rapamycin.
- 2. Kuo, C. J., Chung, J., Fiorentino, D. F., Flanagan, W. M., Blenis, J., Crabtree, G. R. (1992). Rapamycin selectively inhibits interleukin-2 activation of p70 S6 kinase.; Nature,; 358(6381), 70-73.
- 3. Hidalgo, M., Rowinsky, E. K. (2000). The rapamycin-sensitive signal transduction pathway as a target for cancer therapy.; Oncogene,; 19(56), 6680-6686.