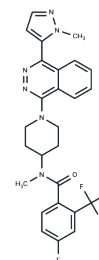


Taladegib

Chemical Properties

| | |
|-------------------|---|
| CAS No. : | 1258861-20-9 |
| Formula: | C ₂₆ H ₂₄ F ₄ N ₆ O |
| Molecular Weight: | 512.5 |
| Appearance: | no data available |
| Storage: | Powder: -20°C for 3 years In solvent: -80°C for 1 year |



Biological Description

| | |
|---------------|--|
| Description | Taladegib (LY2940680) is an orally bioavailable small molecule antagonist of the Hedgehog (Hh)-ligand cell surface receptor smoothened (Smo) with potential antineoplastic activity. |
| Targets(IC50) | Hedgehog/Smoothened,Smo |
| In vitro | LY2940680 inhibits cancer growth in cell lines containing a mutation in the gene encoding Smoothened that researchers had previously observed in patient with cancer who developed resistance to vismodegib. [1] |
| In vivo | Taladegib has excellent pharmacokinetic properties in rodent and non-rodent species. Taladegib administrated orally treats Ptch+/- p53-/- transgenic mice which spontaneously develop medulloblastoma, produces remarkable efficacy and significantly improves their survival. Taladegib reveals rapid kinetics of anti-tumor activity through magnetic resonance imaging of these mice, and Taladegib induces Caspase-3 activity and reduces proliferation by immunohistochemistry analysis of medulloblastoma tumors. Taladegib inhibits Hh regulated gene expression in the subcutaneous xenograft tumor stroma and produces significant anti-tumor activity. [2] |

Solubility Information

| | |
|------------|--|
| Solubility | DMSO: 5.64 mg/mL (11 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble) |
|------------|--|

A DRUG SCREENING EXPERT

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|-----------|------------|
| 1 mM | 1.9512 mL | 9.7561 mL | 19.5122 mL |
| 5 mM | 0.3902 mL | 1.9512 mL | 3.9024 mL |
| 10 mM | 0.1951 mL | 0.9756 mL | 1.9512 mL |
| 50 mM | 0.039 mL | 0.1951 mL | 0.3902 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.

Reference

Redmond EM et al. Expert Opin Investig Drugs, 2011, 20(12),1649-1664.
Mark H. Bender, Cancer Research, 2011, Volume 71, Issue 8, Supplement1

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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Tel:781-999-4286 E_mail:info@targetmol.com Address:36 Washington Street,Wellesley Hills,MA 02481