Inhibitors



Product Data Sheet

NSC689857

Cat. No.: HY-123578 CAS No.: 241127-79-7

Molecular Formula: C₂₅H₂₉NO₄ Molecular Weight: 407.5

Target: EGFR; E1/E2/E3 Enzyme

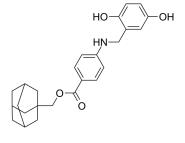
Pathway: JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Metabolic Enzyme/Protease

Storage: Powder -20°C 3 years

> 4°C 2 years

In solvent -80°C 6 months

> -20°C 1 month



SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (245.40 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.4540 mL	12.2699 mL	24.5399 mL
	5 mM	0.4908 mL	2.4540 mL	4.9080 mL
	10 mM	0.2454 mL	1.2270 mL	2.4540 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (6.13 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.13 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.13 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	NSC689857 is a potent EGFR and SCF ^{SKP2} inhibitor with an IC ₅₀ value of 36 μ M for Skp2-Cks1. NSC689857 can inhibit p27 ubiquitylation (IC ₅₀ =30 μ M). NSC689857 has varied activity across cancer types, with more activity against leukemia cell lines than others ^{[1][2]} .
IC ₅₀ & Target	IC $_{50}$: 36 μM (Skp2-Cks1), 30 μM (p27 ubiquitylation) $^{[1]}$

REFERENCES				
1]. Vuong W, et al. High-Throughput Screen for Inhibitors of Androgen Receptor-RUNX2 Transcriptional Regulation in Prostate Cancer. J Pharmacol Exp Ther. 2016 lov;359(2):256-261.				
2]. Lough L, et al. Chemical probes of Skp2-mediated p27 ubiquitylation and degradation. Medchemcomm. 2018 Jun 11;9(7):1093-1104.				

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA

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