

Product Data Sheet

Product Name: Tankyrase Inhibitors (TNKS) 49

Cat. No.: GC11548

Chemical Properties

Cas. No.

Chemical Name N-((1r,4r)-4-(4-cyanophenoxy)cyclohexyl)-3-((4-oxo-3,4-dihydroquinazolin-2-yl)thio)prop anamide

SMILES O=C1NC(SCCC(N[C@H]2CC[C@H](OC3=CC=C(C#N)C=C3)CC2)=O)=NC4=CC=CC=C41

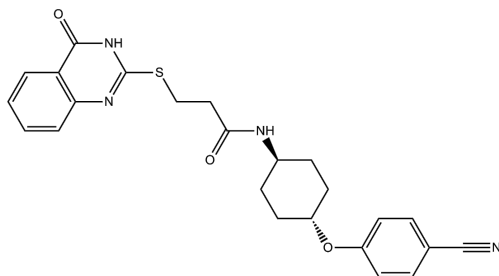
Formula $C_{24}H_{24}N_4O_3S$ M.Wt 448.54

Solubility $\geq 22.45\text{mg/mL}$ in DMSO Storage Store at -20°C

General tips For obtaining a higher solubility , please warm the tube at 37°C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure



Protocol

Cell experiment [1]:

Cell lines SW480-TBC cell lines

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: (909) 407-4943 Fax: (626) 353-8530 E-mail: tech@glpbio.com

Address: 10292 Central Ave. #205, Montclair, CA, USA

Product Data Sheet

Preparation method The solubility of this compound in DMSO is >10 mM. General tips for obtaining a higher concentration: Please warm the tube at 37 °C for 10 minutes and/or shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Reaction Conditions 24 h; IC₅₀=1.9 nM

Applications Tankyrase Inhibitors (TNKS) 49, phenoxy compounds, has a good enzymatic potency and cellular potency with IC₅₀ value of 1.9 nM in the SW480-TBC cellular assay. The compound demonstrated excellent potencies in TNKS2 autoparsylation assay and the two additional functional cellular assays.

Animal experiment [1]:

Animal models Athymic nude mice.

Dosage form 10 and 50 mg/kg; q.d.; oral taken

Applications Tankyrase Inhibitors (TNKS) 49 was evaluated for Wnt-pathway specific pharmacological activity in mouse tumor pharmacodynamic (PD) models. Upon once daily oral administration (at 10 and 50 mg/kg) to mice (n=4) bearing human DLD-1 tumors for 3 days, both compounds exhibited statistically significant, dose-dependent axin2 accumulation (2.1- to 3-fold) and inhibition of STF (70–79%) at day 3 (24 h after the last dose) .

Other notes Please test the solubility of all compounds indoor, and the actual solubility may slightly differ with the theoretical value. This is caused by an experimental system error and it is normal.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: (909) 407-4943 Fax: (626) 353-8530 E-mail: tech@glpbio.com

Address: 10292 Central Ave. #205, Montclair, CA, USA

Product Data Sheet

References:

[1] Hua Z, Bregman H, Buchanan J L, et al. Development of Novel Dual Binders as Potent, Selective, and Orally Bioavailable Tankyrase Inhibitors[J]. Journal of medicinal chemistry, 2013, 56(24): 10003-10015.

Background

Tankyrase Inhibitors (TNKS) 49 is a potent, selective and orally bioavailable inhibitor of tankyrase with IC₅₀ value of 0.1nM [1].

Tankyrase 1 and tankyrase 2 are members of PARP family. They can use NAD⁺ as substrates to transfer ADP-ribose polymers onto target proteins. The tankyrase are found to bind to PARSylate axin proteins which are the negative regulator of Wnt pathway. It makes tankyrase to be targets in treatment for adenomatous polyposis coli. Tankyrase inhibitors 49 is an optimization of the previous hit compound inhibitor 8 with improved potency and selectivity. It has excellent effects in both tankyrase assay and cellular assay (total β -catenin degradation assay in SW480 cells) with IC₅₀ values of 0.1nM and 1.9nM, respectively. In addition, it is found to be a dual binder with both the nicotinamide pocket and the induced pocket of the enzymes [1].

In the in vivo studies in rodents, tankyrase inhibitors 49 is found to potently inhibit TNKS2 autoparsylation with IC₅₀ value of 7.6nM. It also causes stabilization and accumulation of axin protein in SW480 cells with EC₅₀ value of 4nM. In DLD-1 cells with truncated APC, the inhibitor inhibits the STF reporter transcription with IC₅₀ value of 0.3nM suggesting its downstream inhibitory activity on Wnt-associated transcription [1].

References:

[1] Hua Z, Bregman H, Buchanan J L, et al. Development of Novel Dual Binders as Potent, Selective, and Orally Bioavailable Tankyrase Inhibitors. Journal of medicinal chemistry, 2013, 56(24): 10003-10015.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: (909) 407-4943 Fax: (626) 353-8530 E-mail: tech@glpbio.com

Address: 10292 Central Ave. #205, Montclair, CA, USA