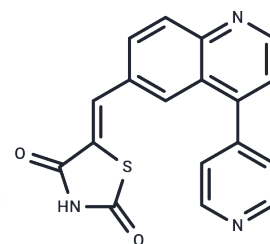


GSK1059615

Chemical Properties

CAS No. : 958852-01-2
 Formula: C₁₈H₁₁N₃O₂S
 Molecular Weight: 333.36
 Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year
 Actual storage temperature shall be subject to the COA.



Biological Description

Description	GSK1059615 has been used in trials studying the treatment of Lymphoma, Solid Tumours, Endometrial Cancer, Solid Tumor Cancer, and Metastatic Breast Cancer.
Targets(IC50)	Apoptosis,mTOR,PI3K
In vitro	GSK1059615 (25 mg/kg) effectively inhibited tumor growth in a xenograft mouse model of BT474 or HCC1954 breast cancer cells.
In vivo	GSK1059615 inhibited the PI3K pathway and induced cell arrest in G1 phase, and apoptosis was observed.GSK1059615 inhibited PI3K and decreased intracellular AKT phosphorylation at Ser473 in T47D and BT474 cancer cells with an IC50 of 40 nM. GSK1059615 also inhibited mTOR with an IC50 of 12 nM.
Kinase Assay	HTRF In vitro Profiling Assays for PI3K Inhibition: The measurement of the GSK1059615-dependent inhibition of the PI3Ks is accessed using a HTRF based PI3K profiling assay kit. 400 pM enzyme is used in PI3K α and δ assays, 200 pM in PI3K β assays, and 1 nM in PI3K γ assay. In addition, the PI3K α , β , and δ assays are run with 150 mM NaCl and 100 μ M ATP, while the PI3K γ assay is run with no NaCl and 15 μ M ATP. All reactions are run at 10 μ M PIP2. GSK1059615 is serially diluted (3-fold in DMSO), and 50 nL is transferred to a 384-well low-volume assay plate. PI3K Reaction Buffer is prepared by diluting the stock 1:4 with de-ionized water. Freshly prepared DTT is added at a final concentration of 5 mM on the day of use. Enzyme addition and GSK1059615 pre-incubation are initiated by the addition of 2.5 μ L of PI3K in reaction buffer. Plates are incubated at room temperature for 15 min. Reactions are initiated by addition of 2.5 μ L of 2 \times substrate solution (PIP2 and ATP in 1 \times reaction buffer). Plates are incubated at room temperature for one hour. Reactions are quenched by the addition of 2.5 μ L of stop solution. The quenched reactions are then processed to detect product formation by adding 2.5 μ L of Detection Solution. Following a 1-hour incubation in the dark, the HTRF signal is measured on the Envision plate reader set for 330 nM excitation and dual emission detection at 620 nM (Eu) and 665 nM (APC). The IC50 value is then obtained.
Cell Research	Cells are plate at a density of 1 \times 10 ⁴ cells per well in clear flat-bottomed 96-well plates and incubated overnight. Then, GSK1059615 is added and the plates are incubated for 30 min. At the end of incubation, media is aspirated from the plates, and the plate is wash once with cold PBS. 80 μ L MSD Lysis buffer is added into each well and the plates are incubated on a shaker at 4 $^{\circ}$ C for at least 30 min. For Akt duplex assay, plates are washed with 200 μ L/well wash buffer for 4 times and tapped on paper towel to blot.

A DRUG SCREENING EXPERT

Cell Research	Then, 60 μ L lysates is added to each well and the plates are incubated on shaker at room temperature for 1 hour. After another 4 times washing, antibody is added (25 μ L per well) and the plates are incubated on shaker for 1 hour and washed again. Finally, Read Buffer is added (150 μ L per well) and the plates are read immediately. IC50 values are then obtained.(Only for Reference)
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Solubility Information

Solubility	DMSO: 5 mg/mL (15 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.9998 mL	14.9988 mL	29.9976 mL
5 mM	0.600 mL	2.9998 mL	5.9995 mL
10 mM	0.300 mL	1.4999 mL	2.9998 mL
50 mM	0.060 mL	0.300 mL	0.600 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.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Carnero A. Expert Opin Investig Drugs, 2009, 18(9), 1265-1277.

Auger KR, et al. EORTC-NCI-AACR, 2008.

Bei S, et al. Inhibition of gastric cancer cell growth by a PI3K-mTOR dual inhibitor GSK1059615. Biochem Biophys Res Commun. 2019 Mar 26;511(1):13-20.

Knight SD, et al. ACS Med Chem Lett, 2010, 1(1), 39-43.

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