

Product Name: MK-8776 (SCH-900776)

Revision Date: 01/10/2021

Product Data Sheet

MK-8776 (SCH-900776)

Cat. No.: A8477

CAS No.: 891494-63-6
Formula: C15H18BrN7

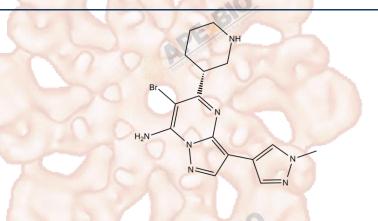
M.Wt: 376.25

Synonyms:

Target: Cell Cycle/Checkpoint

Pathway: Chk

Storage: Store at -20°C



Solvent & Solubility

insoluble in H2O; insoluble in EtOH; ≥18.8 mg/mL in DMSO

In Vitro

In Vitro

Preparing Stock Solutions	Solvent Concentration	1mg	5mg	10mg
	1 mM	2.6578 mL	13.2890 mL	26.5781 mL
	5 mM	0.5316 mL	2.6578 mL	5.3156 mL
	10 mM	0.2658 mL	1.3289 mL	2.6578 mL

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

Biological Activity

Shortsummary	Chk1 inhibitor,potent and selective	
IC ₅₀ & Target	3 nM (Chk1), 0.16 μM (CDK2), 1.5 μM (Chk2)	
	# A 7/300	

Cell Viability Assay

10"	
Cell Line:	U2OS cells
Preparation method:	The solubility of this compound in DMSO is >18.8mg/mL. 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.
Reacting conditions:	2h

	Applications:	In U2OS cells, SCH 900776 induced a dose-dependent loss of DNA replication capability 24 hours after hydroxyurea exposure. SCH 900776 enhanced apoptosis for at least 48 hours following release from hydroxyurea blockade.			
	Animal experiment				
In Vivo	Animal models:	BALB/c mice bearing A2780 xenografts			
	Dosage form:	4-32 mg/kg			
	Applications:	SCH-900776 (8 mg/kg) led to enhanced tumor pharmacodynamic and regression responses. SCH 900776 (16 and 32 mg/kg) induced incremental improvements in tumor response.			
	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.			

Product Citations

1. Bourgeois A, Bonnet S, et al. "Inhibition of CHK 1 (Checkpoint Kinase 1) Elicits Therapeutic Effects in Pulmonary Arterial Hypertension." Arterioscler Thromb Vasc Biol. 2019 Aug;39(8):1667-1681.PMID:31092016

See more customer validations on www.apexbt.com.

References

[1] Guzi T J, Paruch K, Dwyer M P, et al. Targeting the replication checkpoint using SCH 900776, a potent and functionally selective CHK1 inhibitor identified via high content screening[J]. Molecular cancer therapeutics, 2011, 10(4): 591-602.

Caution

FOR RESEARCH PURPOSES ONLY.

NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

Specific storage and handling information for each product is indicated on the product datasheet. Most APExBIO products are stable under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Shortterm storage of many products are stable in the short-term at temperatures that differ from that required for long-term storage. We ensure that the product is shipped under conditions that will maintain the quality of the reagents. Upon receipt of the product, follow the storage recommendations on the product data sheet.



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