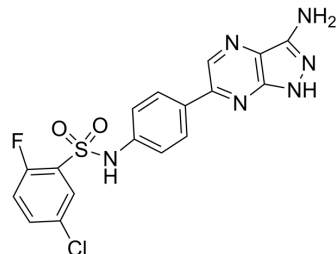


SGK1-IN-1

Cat. No.:	HY-18607		
CAS No.:	1279829-87-6		
Molecular Formula:	C ₁₇ H ₁₂ ClFN ₆ O ₂ S		
Molecular Weight:	418.83		
Target:	SGK		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 50 mg/mL (119.38 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
			10 mg	
Preparing Stock Solutions	1 mM	2.3876 mL	11.9380 mL	23.8760 mL
	5 mM	0.4775 mL	2.3876 mL	4.7752 mL
	10 mM	0.2388 mL	1.1938 mL	2.3876 mL
Please refer to the solubility information to select the appropriate solvent.				
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (5.97 mM); Clear solution			

BIOLOGICAL ACTIVITY

Description	SGK1-IN-1 is a highly active and selective inhibitor of SGK-1, with an IC ₅₀ of 1 nM.
IC₅₀ & Target	SGK1
In Vitro	SGK1-IN-1 (14n) shows an acceptable SGK isoform selectivity with a good activity on hSGK2 and a moderate activity on the hSGK3, with IC ₅₀ s of 1, 41 nM at 10 μM and 50 μM ATP concentration, respectively. SGK1-IN-1 also displays cellular activity in a SGK1-dependent phosphorylation of GSK3β assay in U2OS cells with activities of 0.69 μM ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

-
- Food Chem Toxicol. 2023 Mar 25;113743.

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REFERENCES

[1]. ARYL SULFONOHYDRAZIDES. WO 2013041119 A1.

[2]. Halland N, et al. Discovery of N-[4-(1H-Pyrazolo[3,4-b]pyrazin-6-yl)-phenyl]-sulfonamides as Highly Active and Selective SGK1 Inhibitors. ACS Med Chem Lett. 2014 Oct 23;6(1):73-8.

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

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