U-73343

Cat. No.:	HY-108630			
CAS No.:	142878-12-4			
Molecular Formula:	$C_{29}H_{42}N_{2}O_{3}$			
Molecular Weight:	466.66			
Target:	Phospholipase			
Pathway:	Metabolic Enzyme/Protease			
Storage:	Powder	-20°C	3 years	
	In solvent	-80°C	6 months	
		-20°C	1 month	

SOLVENT & SOLUBILITY

In Vitro

DMSO : 3.33 mg/mL (7.14 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.1429 mL	10.7144 mL	21.4289 mL
	5 mM	0.4286 mL	2.1429 mL	4.2858 mL
	10 mM			

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

Diologicality			
Description	U-73343, works as a protonophore, is an inactive analog of U-73122 and can be used as a negative control. U-73343 dose- dependently inhibits acid secretion irrespective of the stimulant. U-73122 is a phospholipase C (PLC) and 5-LO (5- lipoxygenase) inhibitor with an IC ₅₀ of 1-2.1 μM for PLC ^{[1][2]} .		
In Vitro	U73343, which potently inhibits acid secretion, shows no inhibitory effect on either K ⁺ -pNPPase or H ⁺ ,K ⁺ -ATPase activity. U73343 dose-dependently inhibits, histamine (Hist)-, carbachol (CCh)-, and dbcAMPstimulated aminopyrine accumulation in gastric glands ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

CUSTOMER VALIDATION

- Sci Signal. 2021 Dec 14;14(713):eabj4243.
- J Biol Chem. 2022 May;298(5):101847.

Product Data Sheet

REFERENCES

[1]. Muto Y, et al. The putative phospholipase C inhibitor U73122 and its negative control, U73343, elicit unexpected effects on the rabbit parietal cell. J Pharmacol Exp Ther. 1997 Sep;282(3):1379-88.

[2]. A Tatrai, et al. U-73122, a phospholipase C antagonist, inhibits effects of endothelin-1 and parathyroid hormone on signal transduction in UMR-106 osteoblastic cells. Biochim Biophys Acta. 1994 Dec 30;1224(3):575-82.

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

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