

Product Name: Go 6983 Revision Date: 01/10/2021 Product Data Sheet

Go 6983

Cat. No.:	A8343	H _0 _0-
CAS No.:	133053-19-7	
Formula:	C26H26N4O3	
M.Wt:	442.51	
Synonyms:	Goe 6983;Go6983;Go-6983	
Target:	TGF- β / Smad Signaling	
Pathway:	PKC	N
Storage:	Store at -20°C	
	810	810
Solvent 8	& Solubility	BEE

≥22.15 mg/mL in DMSO: insoluble in EtOH: insoluble in H2O

In Vitro	Preparing Stock Solutions	Solvent Concentration	1mg	5mg	10mg
		1 mM	2.2598 mL	11.2992 mL	22.5984 mL
	PEBIO	5 mM	0.4520 mL	2.2598 mL	4.5197 mL
		10 mM	0.2260 mL	1.1299 mL	2.2598 mL

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

Biological Activity

Shortsummary pan-PKC inhibitor

IC₅₀ & Target

In Vitro

6 nM (PKCγ), 7 nM (PKCα), 7 nM (PKCβ), 10 nM (PKCδ), 60 nM (PKCζ), 20 μM (PKCμ)

Cell Viability Assay	To and
Cell Line:	ARCcPE cells
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.
Reacting conditions:	200 and 1000 nM; 48 hrs

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	Applications:	In ARCaPE cells, Go 6983 at the dose of 200 nM significantly inhibited the		
		up-regulation of PKCs(PKCa, PKC β and PKC γ) stimulated by the PMA		
		treatment. In addition, Go 6983 showed complete inhibition at the dose of 1000		
		nM.		
	Animal experiment			
In Vivo	Animal models:	Mice bearing B16BL6 tumors		
	Dosage form:	22 μg/mouse, i.v.		
	Applications:	In a mouse pulmonary B16BL6 tumor model, Go6983 at the dose of 22 μ g/mouse significantly inhibited tumor metastasis by 51.2 %.		
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility material		
		slightly differ with the theoretical value. This is caused by an experimental		
		system error and it is normal.		

Product Citations



1. Lee E, Wang J, et al. "DNMT1 Regulates Epithelial-Mesenchymal Transition and Cancer Stem Cells, Which Promotes Prostate Cancer Metastasis."Neoplasia. 2016 Sep;18(9):553-66.PMID:27659015

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References

[1]. Gschwendt, M., et al., Inhibition of protein kinase C mu by various inhibitors. Differentiation from protein kinase c isoenzymes. FEBS Lett, 1996. 392(2): p. 77-80.

[2]. He, H., et al., Phorbol ester phorbol-12-myristate-13-acetate induces epithelial to mesenchymal transition in human prostate cancer ARCaPE cells. Prostate, 2010. 70(10): p. 1119-26.

[3]. Kim HR, Lee KH, Park SJ, Kim SY, Yang YK, Tae J, Kim J. Anti-cancer activity and mechanistic features of a NK cell activating molecule. Cancer Immunol Immunother. 2009 Oct;58(10):1691-700.

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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