

Product Name: TG003 Revision Date: 01/10/2021 Product Data Sheet

TG003

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Cat. No.:	B1431	
CAS No.:	300801-52-9	s o
Formula:	C13H15NO2S	
M.Wt:	249.33	
Synonyms:		
Target:		
Pathway:		
Storage:	Store at -20°C	
	BIO	BIO
Solvent & Solubility		
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	insoluble in H2O; \geq	insoluble in H2O; \geq 12.45 mg/mL in DMSO; \geq 14.67 mg/mL in EtOH with ultrasonic				
In Vitro	Preparing Stock Solutions	Mass Solvent Concentration	1mg	5mg	10mg	
		1 mM	4.0107 mL	20.0537 mL	40.1075 mL	
		5 mM	0.8021 mL	4.0107 mL	8.0215 mL	
	PERMI	10 mM	0.4011 mL	2.0054 mL	4.0107 mL	

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

Biological Activity

Shortsummary

Cdc2-like kinase (Clk) inhibitor

IC₅₀ & Target

In Vitro

Cell Viability Assay	and the second
Cell Line:	HeLa cells
Preparation method:	Soluble in DMSO >12.5mg/mL. General tips for obtaining a higher
	the ultrasonic bath for a while. Stock solution can be stored below -20°C for
	several months.
Reacting conditions:	$2\mu I$ of 10 mM TG003 dissolved in Me2SO, final concentration at $10\mu M,$ 3 days

1 | www.apexbt.com

	Applications:	TG003 had a potent inhibitory effect on the activity of Clk1(Cdc2 like kinase1).			
		TG003 inhibited SF2(Splicing factor2) -dependent splicing of β -globin			
		pre-mRNA in vitro by suppression of Clk-mediated phosphorylation. This drug			
		also suppressed serine/arginine-rich protein phosphorylation, dissociation of			
		nuclear speckles, and Clk1-dependent alternative splicing in cells.			
	Animal experiment	310			
In Vivo	Animal models:	Seven-week-old, male Jcl:TCR mice			
	Dosage form:	30mg/kg TG003 suspended in 5% DMSO, 5% Solutol, 9% Tween-80, and 81% saline, subcutaneously injection			
	Applications:	TG003, an inhibitor of CLK1 in mice, could act as a splice-modifying compound for exon-skipping therapy. TG003 promoted skipping of dystrophin exon 31 with the c.4303G > T mutation.			
	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.			
Produc	et Citations	APE			

See more customer validations on www.apexbt.com.

References



PE-BIC

[1]. Muraki M, Ohkawara B, et al. Manipulation of alternative splicing by a newly developed inhibitor of Clks. J Biol Chem, 2004, 279(23): 24246-24254.

[2]. Sako Y1, Ninomiya K1, et al, Development of an orally available inhibitor of CLK1 for skipping a mutated dystrophin exon in Duchenne muscular dystrophy. Sci Rep. 2017 May 30;7:46126. doi: 10.1038/srep46126.

Caution

FOR RESEARCH PURPOSES ONLY.

NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

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