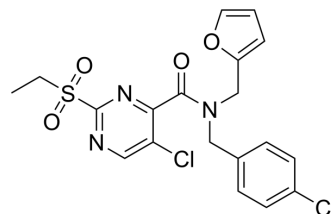


ZAP-180013

Cat. No.:	HY-136179	
CAS No.:	873080-25-2	
Molecular Formula:	C ₁₉ H ₁₇ Cl ₂ N ₃ O ₄ S	
Molecular Weight:	454.33	
Target:	Tyrosinase	
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 : 250 mg/mL (550.26 mM; Need ultrasonic)																								
	Preparing Stock Solutions	<table border="1"> <thead> <tr> <th>Solvent Concentration</th> <th>Mass</th> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> </thead> <tbody> <tr> <td>1 mM</td> <td></td> <td>2.2010 mL</td> <td>11.0052 mL</td> <td>22.0104 mL</td> </tr> <tr> <td>5 mM</td> <td></td> <td>0.4402 mL</td> <td>2.2010 mL</td> <td>4.4021 mL</td> </tr> <tr> <td>10 mM</td> <td></td> <td>0.2201 mL</td> <td>1.1005 mL</td> <td>2.2010 mL</td> </tr> </tbody> </table>	Solvent Concentration	Mass	1 mg	5 mg	10 mg	1 mM		2.2010 mL	11.0052 mL	22.0104 mL	5 mM		0.4402 mL	2.2010 mL	4.4021 mL	10 mM		0.2201 mL	1.1005 mL	2.2010 mL			
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Please refer to the solubility information to select the appropriate solvent.																									
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (4.58 mM); Clear solution																								
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.08 mg/mL (4.58 mM); Suspended solution; Need ultrasonic																								
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.58 mM); Clear solution																								

BIOLOGICAL ACTIVITY

Description	ZAP-180013 is a zeta-chain-associated protein kinase 70 (ZAP-70) inhibitor with an IC ₅₀ of 1.8 μM. ZAP-180013 inhibits the interaction of ZAP-70 SH2 domain with immunoreceptor tyrosine-based activation motif (ITAMs) ^[1] .
IC ₅₀ & Target	IC ₅₀ : 1.8 μM (ZAP-70) ^[1]
In Vitro	ZAP-70 is a critical molecule in the transduction of T cell antigen receptor signaling and the activation of T cells. Upon activation of the T cell antigen receptor, ZAP-70 is recruited to the intracellular ζ-chains of the T cell receptor, where ZAP-70 is activated and colocalized with its substrates. Inhibitors of ZAP-70 could potentially function as treatments for

autoimmune diseases or organ transplantation. ZAP-180013 disrupts the interaction between ZAP-70 and the T cell antigen receptor. The IC_{50} s in both FP and TR-FRET assays for ZAP-180013 are 9.6 μ M and 16.841 μ M, respectively^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Patrick R Visperas, et al. Identification of Inhibitors of the Association of ZAP-70 With the T Cell Receptor by High-Throughput Screen. SLAS Discov. 2017 Mar;22(3):324-331.

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

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