Ziftomenib

Cat. No.:	HY-132001				
CAS No.:	2134675-36-6				
Molecular Formula:	$C_{_{33}}H_{_{42}}F_{_{3}}N_{_{9}}O_{_{2}}S_{_{2}}$				
Molecular Weight:	717.87				
Target:	Epigenetic Reader Domain				
Pathway:	Epigenetics				
Storage:	Powder	-20°C	3 years		
		4°C	2 years		
	In solvent	-80°C	6 months		
		-20°C	1 month		

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SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (139.30 mM; Need ultrasonic)						
Preparing Stock Solutions		Solvent Mass Concentration	1 mg	5 mg	10 mg		
	1 mM	1.3930 mL	6.9650 mL	13.9301 mL			
		5 mM	0.2786 mL	1.3930 mL	2.7860 mL		
		10 mM	0.1393 mL	0.6965 mL	1.3930 mL		
	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 (2.90 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (2.90 mM); Clear solution						

BIOLOGICAL ACTIVITY					
Description	Ziftomenib (KO-539) is an orally active menin-MLL interaction inhibitor with antitumor activities (WO2017161028A1, compound 151) ^[1] .				
In Vitro	The mixed-lineage leukemia (MLL) protein is a histone methyltransferase critical for the epigenetic regulation of gene transcription. Many acute leukemias, including acute myeloblastic leukemia (AML), acute lymphoblastic leukemia (ALL) and mixed-lineage leukemia (MLL), are characterized by the presence of chimeric MLL fusion proteins that result from chromosomal translocations of the MLL gene located at chromosome 11, band q23 (11q23). MLL fusion proteins lack the original histone methyltransferase activity of the C-terminus of MLL and gain the ability to regulate transcription of numerous oncogenes, including HOX and MEIS1, resulting in increased cell proliferation and decreased cell differentiation, ultimately leading to leukemogenesis ^[1] .				

Product Data Sheet

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MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

• Leukemia. 2023 Mar 28.

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REFERENCES

[1]. Tao Wu, et al. Substituted inhibitors of menin-mll and methods of use. WO2017161028A1.

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

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