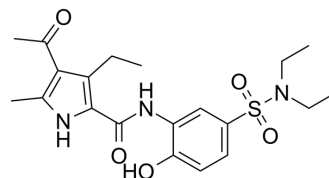


XD14

Cat. No.:	HY-110215		
CAS No.:	1370888-71-3		
Molecular Formula:	C ₂₀ H ₂₇ N ₃ O ₅ S		
Molecular Weight:	421.51		
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



SOLVENT & SOLUBILITY

In Vitro	DMSO : 125 mg/mL (296.55 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
			10 mg	
	Preparing Stock Solutions	1 mM	2.3724 mL	11.8621 mL
	5 mM	0.4745 mL	2.3724 mL	4.7448 mL
	10 mM	0.2372 mL	1.1862 mL	2.3724 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 (4.93 mM); Clear solution			
	2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.93 mM); Clear solution			

BIOLOGICAL ACTIVITY

Description	XD14 is a potent BET inhibitor with antitumor effect. It binds to BRD2, BRD3, and BRD4 with K _d s of 170, 380, and 160 nM, respectively ^[1] .		
IC ₅₀ & Target	BRD2	BRD3	BRD4
	170 nM (Kd)	380 nM (Kd)	160 nM (Kd)

REFERENCES

[1]. Schäker-Hübner L, et al. 4-Acyl Pyrrole Capped HDAC Inhibitors: A New Scaffold for Hybrid Inhibitors of BET Proteins and Histone Deacetylases as Antileukemia Drug Leads. J Med Chem. 2021 Oct 14;64(19):14620-14646.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA