BI-3802

MedChemExpress

Cat. No.:	HY-108705		
CAS No.:	2166387-65-9		
Molecular Formula:	C24H39CIN6C) ₃	
Molecular Weight:	484.98		
Target:	Bcl-2 Family; Molecular Glues		
Pathway:	Apoptosis;	PROTAC	
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month

SOLVENT & SOLUBILITY

In Vitro

		1
Mass		
Solvent	1 mg	5 mg

Preparing Stock Solutions	Solvent Concentration	1 mg	5 mg	10 mg
	1 mM	2.0619 mL	10.3097 mL	20.6194 mL
	5 mM	0.4124 mL	2.0619 mL	4.1239 mL
	10 mM	0.2062 mL	1.0310 mL	2.0619 mL

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

DMSO: 6.67 mg/mL (13.75 mM; ultrasonic and warming and heat to 60°C)

Description	BI-3802 is a highly potent BCL6 degrader and inhibits the Bric-à-brac (BTB) domain of BCL6 with an IC ₅₀ of ≤3 nM. BI-3802 induces the polymerization of BCL6 and promotes BCL6 degration depended on E3 ligase SIAH1. BI-3802 has antitumor activity ^{[1][2]} .	
IC ₅₀ & Target	IC50: ≤ 3 nM (BCL6 BTB) ^[1]	
In Vitro	BI-3802 shows an IC ₅₀ of 43 nM for the cellular BCL6 ^[1] . BI-3802 increases the interaction between BCL6 and SIAH1 (EC ₅₀ = 64 nM) ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

REFERENCES

[1]. Kerres N, et al. Chemically Induced Degradation of the Oncogenic Transcription Factor BCL6. Cell Rep. 2017 Sep 19;20(12):2860-2875.



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

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