PZ703b TFA

Cat. No.:	HY-115718A
Molecular Formula:	C ₈₂ H ₁₀₃ ClF ₆ N ₁₀ O ₁₃ S ₄
Molecular Weight:	1714.46
Target:	PROTACs; Bcl-2 Family
Pathway:	PROTAC; Apoptosis
Storage:	-20°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)

BIOLOGICAL ACTIVITY			
Description	PZ703b TFA is a Bcl-xl PROTAC degradation agent that induces apoptosis and inhibits cancer cell proliferation for bladder cancer research ^{[1][2]} .		
IC ₅₀ & Target	Bcl-xL		
In Vitro	PZ703b (0-1 μM, 24 h) TFA can synergistically inhibit bladder cancer cell proliferation with Mivebresib in a dose-dependent manner and induce apoptosis in bladder cancer cells via the mitochondrial pathway ^[1] . PZ703b (0-1 μM, 48 h) TFA inhibits MOLT-4 and RS4;11 cells with the IC ₅₀ values of 15.9 and 11.3 nM respectively ^[2] . PZ703b (10 nM, 48 h)TFA induces rapid and durable BCL-XL degradation and apoptosis in MOLT-4 cells through the caspase- 3 mediated pathway ^[2] .		

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

Apoptosis Analysis ^[1]	
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Cell Line:	Bladder cancer cell lines 5637, SW780, and HT-1977
Concentration:	1 μΜ
Incubation Time:	24 hours
Result:	Induction of Bcl-xl degradation increased the association between Mcl-1 and Bim (a pro- apoptotic Bcl-2 protein). It was further shown that forced expression of Bcl-xl or Mcl-1 significantly reduced PZ703b-induced apoptosis. Resulted in a slight activation of Bax and Bak.

CUSTOMER VALIDATION

• Biochem Biophys Res Commun. 16 July 2022.

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REFERENCES

[1]. Yi Xu, et al. Mivebresib synergized with PZ703b, a novel Bcl-xl PROTAC degrader, induces apoptosis in bladder cancer cells via the mitochondrial pathway. Biochem Biophys Res Commun. 2022 Oct 1;623:120-126.

[2]. Pratik Pal, et al. Discovery of a Novel BCL-XL PROTAC Degrader with Enhanced BCL-2 Inhibition. J Med Chem. 2021 Oct 14;64(19):14230-14246.

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

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