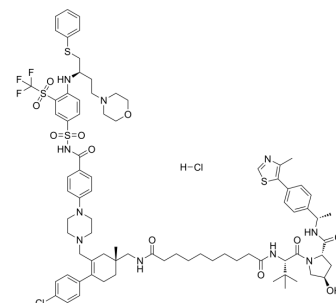


## PZ703b hydrochloride

<b>Cat. No.:</b>	HY-115718B
<b>Molecular Formula:</b>	C <sub>80</sub> H <sub>103</sub> Cl <sub>2</sub> F <sub>3</sub> N <sub>10</sub> O <sub>11</sub> S <sub>4</sub>
<b>Molecular Weight:</b>	1636.9
<b>Target:</b>	PROTACs; Apoptosis; Bcl-2 Family
<b>Pathway:</b>	PROTAC; Apoptosis
<b>Storage:</b>	-20°C, sealed storage, away from moisture and light * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 180 mg/mL (109.96 mM; Need ultrasonic)																	
	<table border="1"> <thead> <tr> <th rowspan="2">Solvent Concentration</th> <th rowspan="2">Mass</th> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> </thead> <tbody> <tr> <td>1 mM</td> <td>0.6109 mL</td> <td>3.0546 mL</td> <td>6.1091 mL</td> </tr> <tr> <td>5 mM</td> <td>0.1222 mL</td> <td>0.6109 mL</td> <td>1.2218 mL</td> </tr> <tr> <td>10 mM</td> <td>0.0611 mL</td> <td>0.3055 mL</td> <td>0.6109 mL</td> </tr> </tbody> </table>	Solvent Concentration	Mass	1 mg	5 mg	10 mg	1 mM	0.6109 mL	3.0546 mL	6.1091 mL	5 mM	0.1222 mL	0.6109 mL	1.2218 mL	10 mM	0.0611 mL	0.3055 mL	0.6109 mL
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	Please refer to the solubility information to select the appropriate solvent.																	
<b>In Vivo</b>	<ol style="list-style-type: none"> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 40% PEG300 &gt;&gt; 5% Tween-80 &gt;&gt; 45% saline Solubility: ≥ 4.5 mg/mL (2.75 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 4.5 mg/mL (2.75 mM); Clear solution</li> </ol>																	

### BIOLOGICAL ACTIVITY

<b>Description</b>	PZ703b hydrochloride is a Bcl-xL PROTAC degrader that induces apoptosis and inhibits cancer cell proliferation. PZ703b hydrochloride can be used for the research of bladder cancer research <sup>[1][2]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	Bcl-xL
<b>In Vitro</b>	<p>PZ703b hydrochloride (0-1 μM, 24 h) synergistically inhibits bladder cancer cell proliferation with Mivebresib with a dose-dependent manner and induces apoptosis in bladder cancer cells via the mitochondrial pathway<sup>[1]</sup>.</p> <p>PZ703b hydrochloride (0-1 μM, 48 h) inhibits cell viability of MOLT-4 and RS4;11 cells with IC<sub>50</sub> values of 15.9 and 11.3 nM, respectively<sup>[2]</sup>.</p> <p>PZ703b hydrochloride (10 nM, 48 h) induces rapid and durable BCL-XL degradation and apoptosis in MOLT-4 cells through the caspase-3 mediated pathway<sup>[2]</sup>.</p>

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

#### Apoptosis Analysis<sup>[1]</sup>

Cell Line:	MOLT-4 cell line
Concentration:	10 $\mu$ M
Incubation Time:	48 hours
Result:	Induced cell apoptosis of MOLT-4 cells.

## 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 Degradar 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.**

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

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