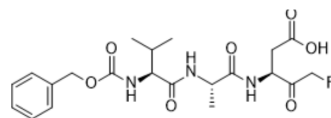


## Z-VAD-FMK

<b>Cat. No.:</b>	HY-16658B
<b>CAS No.:</b>	161401-82-7
<b>Molecular Formula:</b>	C <sub>21</sub> H <sub>28</sub> FN <sub>3</sub> O <sub>7</sub>
<b>Molecular Weight:</b>	453.46
<b>Target:</b>	Caspase; Apoptosis
<b>Pathway:</b>	Apoptosis
<b>Storage:</b>	4°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 : 100 mg/mL (220.53 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>2.2053 mL</td> <td>11.0263 mL</td> <td>22.0527 mL</td> </tr> <tr> <td>5 mM</td> <td>0.4411 mL</td> <td>2.2053 mL</td> <td>4.4105 mL</td> </tr> <tr> <td>10 mM</td> <td>0.2205 mL</td> <td>1.1026 mL</td> <td>2.2053 mL</td> </tr> </tbody> </table>	Solvent Concentration	Mass	1 mg	5 mg	10 mg	1 mM	2.2053 mL	11.0263 mL	22.0527 mL	5 mM	0.4411 mL	2.2053 mL	4.4105 mL	10 mM	0.2205 mL	1.1026 mL	2.2053 mL
Solvent Concentration	Mass			1 mg	5 mg	10 mg												
		1 mM	2.2053 mL	11.0263 mL	22.0527 mL													
5 mM	0.4411 mL	2.2053 mL	4.4105 mL															
10 mM	0.2205 mL	1.1026 mL	2.2053 mL															
	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: ≥ 2.08 mg/mL (4.59 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% (20% SBE-β-CD in saline) Solubility: 2.08 mg/mL (4.59 mM); Suspended solution; Need ultrasonic</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2.08 mg/mL (4.59 mM); Clear solution</li> </ol>																	

### BIOLOGICAL ACTIVITY

<b>Description</b>	Z-VAD-FMK (Z-VAD(OH)-FMK) is a well-know pan caspase inhibitor, which does not inhibit ubiquitin carboxy-terminal hydrolase L1 (UCHL1) activity even at concentrations as high as 440 μM <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	Caspase
<b>In Vitro</b>	Z-VAD-FMK (40 μM) reverses the apoptotic effect exerted by total saponin of Solanum lyratum Thunb (TSSLT) in Hela cells. HeLa cells are pretreated with Z-VAD-FMK (40 μM) for 30 min and exposed to TSSLT (6 μg/mL) for 48 h <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### Cell Viability Assay<sup>[2]</sup>

Cell Line:	HeLa cells
Concentration:	40 $\mu$ M
Incubation Time:	Prtreated for 30 minutes
Result:	Prevented TSSLT-induced cell death. More than 80% cell survival was observed.

## CUSTOMER VALIDATION

- Cell. 2024 Feb 1;187(3):624-641.e23.
- Science. 2021 Mar 5;371(6533):eabb2224.
- Cell Host Microbe. 2023 Nov 8;31(11):1820-1836.e10.
- Nat Microbiol. 2022 Jul;7(7):1041-1053.
- Cell Mol Immunol. 2023 Aug 17.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

## REFERENCES

[1]. Davies CW, et al. The co-crystal structure of ubiquitin carboxy-terminal hydrolase L1 (UCHL1) with a tripeptide fluoromethyl ketone (Z-VAE(OMe)-FMK). Bioorg Med Chem Lett. 2012 Jun 15;22(12):3900-4.

[2]. Liu HR, et al. Antiproliferative activity of the total saponin of Solanum lyratum Thunb in Hela cells by inducing apoptosis. Pharmazie. 2008 Nov;63(11):836-42.

**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