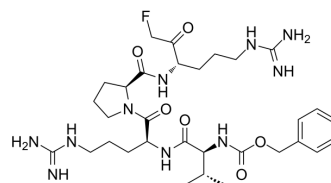


## Z-VRPR-FMK

Cat. No.:	HY-120231
CAS No.:	1381885-28-4
Molecular Formula:	C <sub>31</sub> H <sub>49</sub> FN <sub>10</sub> O <sub>6</sub>
Molecular Weight:	676.78
Target:	MALT1
Pathway:	Metabolic Enzyme/Protease; NF-κB
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Z-VRPR-FMK is an irreversible MALT1 protein inhibitor. Z-VRPR-FMK inhibits the growth and invasion of diffuse large B-cell lymphoma by inhibiting MALT1-induced NF-κB activation and MMP expression <sup>[1]</sup> .
<b>In Vitro</b>	Z-VRPR-FMK (50 μM, 24 h) can effectively reduce the level of c-REL in the nucleus of HBL-1 cells and inhibit the signal transduction of NF-κB <sup>[2]</sup> . Z-VRPR-FMK (50 μM, 48 h) can inhibit the proliferation of ABC-DLBCL cell lines HBL-1, TMD8, OCI-Ly3 and OCI-Ly10 <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Jianglong Feng, et al. Z-VRPR-FMK can inhibit the growth and invasiveness of diffuse large B-cell lymphoma by depressing NF-κB activation and MMP expression induced by MALT1. *Int J Clin Exp Pathol*. 2019 Jun 1;12(6):1947-1955.

[2]. Lorena Fontan, et al. MALT1 small molecule inhibitors specifically suppress ABC-DLBCL in vitro and in vivo. *Cancer Cell*. 2012 Dec 11;22(6):812-24.

**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