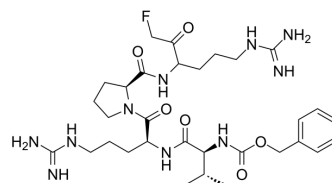


## Val-Arg-Pro-DL-Arg-Fluoromethylketone

Cat. No.:	HY-P4544
CAS No.:	1926163-57-6
Molecular Formula:	C <sub>31</sub> H <sub>49</sub> FN <sub>10</sub> O <sub>6</sub>
Molecular Weight:	676.78
Sequence:	Val-Arg-Pro-{DL-Arg}-{Fluoromethylketone}
Sequence Shortening:	VRP-{DL-Arg}-{Fluoromethylketone}
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>	Val-Arg-Pro-DL-Arg-Fluoromethylketone is a potent MALT1 inhibitor. Val-Arg-Pro-DL-Arg-Fluoromethylketone inhibits cell proliferation and migration. Val-Arg-Pro-DL-Arg-Fluoromethylketone shows anticancer activity <sup>[1]</sup> .								
<b>IC<sub>50</sub> &amp; Target</b>	MALT1 <sup>[1]</sup>								
<b>In Vitro</b>	<p>Val-Arg-Pro-DL-Arg-Fluoromethylketone (75 mol/L; 6, 12, 24, 36, and 48 h) inhibits the proliferation and migration of OCI-LY10 cells<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>OCI-LY10 cells</td> </tr> <tr> <td>Concentration:</td> <td>75 mol/L</td> </tr> <tr> <td>Incubation Time:</td> <td>6, 12, 24, 36, and 48 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited the proliferation and migration of OCI-LY10 cells.</td> </tr> </table>	Cell Line:	OCI-LY10 cells	Concentration:	75 mol/L	Incubation Time:	6, 12, 24, 36, and 48 h	Result:	Inhibited the proliferation and migration of OCI-LY10 cells.
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Concentration:	75 mol/L								
Incubation Time:	6, 12, 24, 36, and 48 h								
Result:	Inhibited the proliferation and migration of OCI-LY10 cells.								
<b>In Vivo</b>	<p>Val-Arg-Pro-DL-Arg-Fluoromethylketone (37.5 M*0.2 mL; every other day for 12 days) inhibits the growth of the tumor in mouse<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>4-6 weeks, 16-20 g, Female BALB/c nude mice (OCI-LY10 cells)<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>37.5 M*0.2 mL</td> </tr> <tr> <td>Administration:</td> <td>Every other day for 12 days</td> </tr> <tr> <td>Result:</td> <td>Inhibited the growth of the xenografts and decreased the expression of P65.</td> </tr> </table>	Animal Model:	4-6 weeks, 16-20 g, Female BALB/c nude mice (OCI-LY10 cells) <sup>[1]</sup>	Dosage:	37.5 M*0.2 mL	Administration:	Every other day for 12 days	Result:	Inhibited the growth of the xenografts and decreased the expression of P65.
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Result:	Inhibited the growth of the xenografts and decreased the expression of P65.								

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## REFERENCES

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

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

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