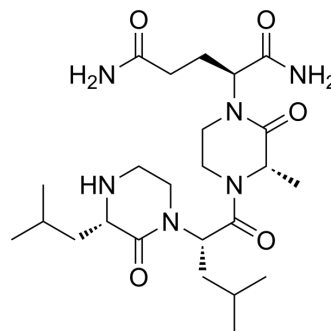


## OHM1

Cat. No.:	HY-124722
CAS No.:	1450995-09-1
Molecular Formula:	C <sub>24</sub> H <sub>42</sub> N <sub>6</sub> O <sub>5</sub>
Molecular Weight:	494.63
Target:	HIF/HIF Prolyl-Hydroxylase
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	OHM1 is an analog of HIF1α CTAD that inhibits its binding with p300/CBP. OHM1 targets CH1 domain with an affinity of 0.53 μM <sup>[1]</sup> .								
<b>In Vitro</b>	<p>OHM1 (1-20 μM; 24 h) results in a dose-dependent reduction in the HIF promoter activity, and reduces the level of HIF1α transcriptional activity under hypoxia to that observed under normoxia at 20 μM in MDA-MB-231 cells<sup>[1]</sup>.</p> <p>OHM1 down-regulates hypoxia-inducible gene expression<sup>[1]</sup>.</p> <p>OHM1 (10 μM) down-regulates multiple genes implicated in angiogenesis, apoptosis, cell proliferation, and invasion, along with several cancer-specific markers in the A549 non-small-cell lung cancer cell line<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>RT-PCR<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>A549</td> </tr> <tr> <td>Concentration:</td> <td>10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Down-regulated the mRNA expression levels of the critical angiogenesis regulator vascular endothelial growth factor (VEGFA) by 80%. Decreased lysyl oxidase (LOX) and glucose transporter 1 (GLUT1) expression.</td> </tr> </table>	Cell Line:	A549	Concentration:	10 μM	Incubation Time:	24 h	Result:	Down-regulated the mRNA expression levels of the critical angiogenesis regulator vascular endothelial growth factor (VEGFA) by 80%. Decreased lysyl oxidase (LOX) and glucose transporter 1 (GLUT1) expression.
Cell Line:	A549								
Concentration:	10 μM								
Incubation Time:	24 h								
Result:	Down-regulated the mRNA expression levels of the critical angiogenesis regulator vascular endothelial growth factor (VEGFA) by 80%. Decreased lysyl oxidase (LOX) and glucose transporter 1 (GLUT1) expression.								
<b>In Vivo</b>	<p>OHM1 (15 mg/kg; i.p.; every other day for 15 injections) reduces MDA-MB-231 tumor volume in mice<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>BALB/c mice, MDA-MB-231 xenograft model<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>15 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intraperitoneal injection, every other day for 15 injections</td> </tr> <tr> <td>Result:</td> <td>Reduced the median tumor volume by roughly 50% compared with the untreated group. Did not cause measurable changes in animal body weight or other signs of toxicity in tumor-bearing animals, nor increased the metastasis rate.</td> </tr> </table>	Animal Model:	BALB/c mice, MDA-MB-231 xenograft model <sup>[1]</sup>	Dosage:	15 mg/kg	Administration:	Intraperitoneal injection, every other day for 15 injections	Result:	Reduced the median tumor volume by roughly 50% compared with the untreated group. Did not cause measurable changes in animal body weight or other signs of toxicity in tumor-bearing animals, nor increased the metastasis rate.
Animal Model:	BALB/c mice, MDA-MB-231 xenograft model <sup>[1]</sup>								
Dosage:	15 mg/kg								
Administration:	Intraperitoneal injection, every other day for 15 injections								
Result:	Reduced the median tumor volume by roughly 50% compared with the untreated group. Did not cause measurable changes in animal body weight or other signs of toxicity in tumor-bearing animals, nor increased the metastasis rate.								

---

## REFERENCES

---

[1]. Lao BB, et al. In vivo modulation of hypoxia-inducible signaling by topographical helix mimetics. Proc Natl Acad Sci U S A. 2014 May 27;111(21):7531-6.

---

**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