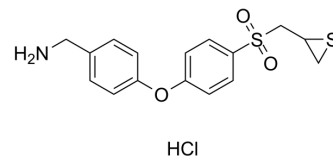


## ND-336

Cat. No.:	HY-124373		
CAS No.:	1807453-83-3		
Molecular Formula:	C <sub>16</sub> H <sub>18</sub> ClNO <sub>3</sub> S <sub>2</sub>		
Molecular Weight:	371.9		
Target:	MMP		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



## BIOLOGICAL ACTIVITY

<b>Description</b>	ND-336 is a selective inhibitor of matrix metalloproteinase (MMP)-2, MMP-9, and MMP-14, with K <sub>i</sub> s of 85, 150, and 120 nM, respectively. ND-336 accelerates diabetic wound healing in mice by lowering inflammation and by enhancing angiogenesis and re-epithelialization of the wound <sup>[1][2]</sup> .										
<b>IC<sub>50</sub> &amp; Target</b>	MMP-2 85 nM (K <sub>i</sub> )	MMP-9 150 nM (K <sub>i</sub> )	MMP-14 120 nM (K <sub>i</sub> )								
<b>In Vivo</b>	<p>ND-336 accelerates diabetic wound healing by decreasing inflammation and by enhancing angiogenesis and re-epithelialization of the wound, thus reversing the pathological condition<sup>[1]</sup>.</p> <p>ND-336 (0.05-0.01 mg; topical application; daily for 14 day) accelerates diabetic wound healing<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>Female diabetic db/db mice<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>0.05, 0.025, and 0.01 mg</td> </tr> <tr> <td>Administration:</td> <td>Topical application; daily for 14 day</td> </tr> <tr> <td>Result:</td> <td>Healed 1.2- to 1.6-fold faster than those treated with ND-322 than those treated with vehicle.</td> </tr> </table>			Animal Model:	Female diabetic db/db mice <sup>[1]</sup>	Dosage:	0.05, 0.025, and 0.01 mg	Administration:	Topical application; daily for 14 day	Result:	Healed 1.2- to 1.6-fold faster than those treated with ND-322 than those treated with vehicle.
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## REFERENCES

- [1]. Gao M, et al. Acceleration of diabetic wound healing using a novel protease-anti-protease combination therapy. Proc Natl Acad Sci U S A. 2015;112(49):15226-15231.
- [2]. Nguyen TT, et al. Validation of Matrix Metalloproteinase-9 (MMP-9) as a Novel Target for Treatment of Diabetic Foot Ulcers in Humans and Discovery of a Potent and Selective Small-Molecule MMP-9 Inhibitor That Accelerates Healing. J Med Chem. 2018;61(19):8825-8837.

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

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