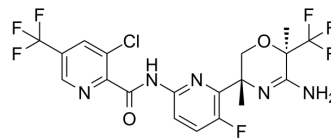


## Umibecestat

Cat. No.:	HY-119689
CAS No.:	1387560-01-1
Molecular Formula:	C <sub>19</sub> H <sub>15</sub> ClF <sub>7</sub> N <sub>5</sub> O <sub>2</sub>
Molecular Weight:	513.8
Target:	Beta-secretase
Pathway:	Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Umibecestat (CNP520) is a beta-site amyloid precursor protein cleaving enzyme-1 (BACE-1) inhibitor with IC <sub>50</sub> s of 11 nM and 10 nM for human BACE-1 and mouse BACE-1, respectively <sup>[1]</sup> . Umibecestat can be used for the research of alzheimer's disease.																
<b>IC<sub>50</sub> &amp; Target</b>	IC <sub>50</sub> : 11 nM (human BACE-1), 10 nM (mouse BACE-1) <sup>[1]</sup>																
<b>In Vitro</b>	Umibecestat (CNP520) is a potent BACE-1 inhibitor that is selective for BACE-1 over other human pepsin-like aspartic proteases, including BACE-2 and cathepsin D <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.																
<b>In Vivo</b>	<p>Umibecestat (CNP520) (1.5-51.3 mg/kg; given by oral gavage; 72 hours) shows a dose-dependent effects on Aβ<sub>40</sub> and a long duration of action in both rat brain and CSF<sup>[1]</sup>.</p> <p>Umibecestat (CNP520) (3.1 mg/kg; oral administration; 7 days) shows a &gt; 75% reduction on Aβ<sub>40</sub> and Aβ<sub>42</sub> in CSF after dosing and returns slowly to baseline over the next 7 days<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>Male rats (3-4 months old)<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>1.5 mg/kg (3 μM/kg)-51.3 mg/kg (100 μM/kg)</td> </tr> <tr> <td>Administration:</td> <td>Given by oral gavage; 72 hours</td> </tr> <tr> <td>Result:</td> <td>Reduced 89.3±4.5% Aβ<sub>40</sub> at the highest dose in brain tissue, and 50% lowering of rat brain Aβ<sub>40</sub> (ED<sub>50</sub>) was 2.4±0.31 mg/kg. Reduced ~50% Aβ<sub>40</sub> at a single oral 30 μM/kg (15.4 mg/kg) dose after 24 hours in both rat brain and CSF</td> </tr> </table> <table border="1"> <tr> <td>Animal Model:</td> <td>3-month-old beagle dogs<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>3.1 mg/kg (6 μM/kg)</td> </tr> <tr> <td>Administration:</td> <td>Oral administration; 7 days</td> </tr> <tr> <td>Result:</td> <td>Both Aβ<sub>40</sub> and Aβ<sub>42</sub> concentrations in CSF showed a &gt; 75% reduction at 12-48 h after</td> </tr> </table>	Animal Model:	Male rats (3-4 months old) <sup>[1]</sup>	Dosage:	1.5 mg/kg (3 μM/kg)-51.3 mg/kg (100 μM/kg)	Administration:	Given by oral gavage; 72 hours	Result:	Reduced 89.3±4.5% Aβ <sub>40</sub> at the highest dose in brain tissue, and 50% lowering of rat brain Aβ <sub>40</sub> (ED <sub>50</sub> ) was 2.4±0.31 mg/kg. Reduced ~50% Aβ <sub>40</sub> at a single oral 30 μM/kg (15.4 mg/kg) dose after 24 hours in both rat brain and CSF	Animal Model:	3-month-old beagle dogs <sup>[1]</sup>	Dosage:	3.1 mg/kg (6 μM/kg)	Administration:	Oral administration; 7 days	Result:	Both Aβ <sub>40</sub> and Aβ <sub>42</sub> concentrations in CSF showed a > 75% reduction at 12-48 h after
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## REFERENCES

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[1]. Neumann U, et al. The BACE-1 inhibitor CNP520 for prevention trials in Alzheimer's disease. EMBO Mol Med. 2018 Nov;10(11). pii: e9316.

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

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