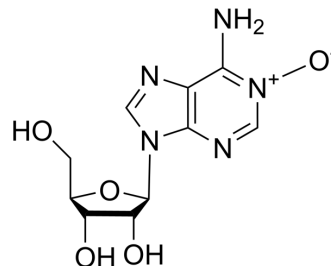


## Adenosine N1-oxide

<b>Cat. No.:</b>	HY-102082
<b>CAS No.:</b>	146-92-9
<b>Molecular Formula:</b>	C <sub>10</sub> H <sub>13</sub> N <sub>5</sub> O <sub>5</sub>
<b>Molecular Weight:</b>	283.24
<b>Target:</b>	Others
<b>Pathway:</b>	Others
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

Description	Adenosine N1-oxide is an oral active anti-inflammatory agent, and can be isolated from royal jelly. Adenosine N1-oxide promotes osteogenic and adipocyte differentiation <sup>[1][2]</sup> .																
In Vitro	<p>Adenosine N1-oxide (1-40 μM, 24 h) inhibits TNF-α and IL-6 secretion by RAW264.7 cells stimulated with 2 μg/mL LPS (HY-D1056)<sup>[2]</sup>.</p> <p>Adenosine N1-oxide (20/100 μM, 30 mins) increases intracellular cAMP production in both peritoneal macrophages and RAW264.7 cells in a dose-dependent manner<sup>[2]</sup>.</p> <p>Adenosine N1-oxide (5-10 μM, 30 mins) increases the protein level of phosphorylation of Akt and GSK-3β in 2 μg/mL LPS (HY-D1056)-stimulated RAW264.7 cells<sup>[2]</sup>.</p> <p>Adenosine N1-oxide (2-10 μM, 6-7 days) promotes osteogenic and adipocyte differentiation in MC3T3-E1 cells<sup>[2]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis<sup>[2]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>RAW264.7 cells</td> </tr> <tr> <td>Concentration:</td> <td>2, 5 and 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>30 mins</td> </tr> <tr> <td>Result:</td> <td>Increased the protein level of phosphorylation of Akt and GSK-3β in 2 μg/mL LPS (HY-D1056)-stimulated RAW264.7 cells.</td> </tr> </table> <p>Cell Differentiation Assay<sup>[2]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>MC3T3-E1 cells</td> </tr> <tr> <td>Concentration:</td> <td>2, 5 and 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>6-7 days</td> </tr> <tr> <td>Result:</td> <td>Promoted osteogenic and adipocyte differentiation in MC3T3-E1 cells</td> </tr> </table>	Cell Line:	RAW264.7 cells	Concentration:	2, 5 and 10 μM	Incubation Time:	30 mins	Result:	Increased the protein level of phosphorylation of Akt and GSK-3β in 2 μg/mL LPS (HY-D1056)-stimulated RAW264.7 cells.	Cell Line:	MC3T3-E1 cells	Concentration:	2, 5 and 10 μM	Incubation Time:	6-7 days	Result:	Promoted osteogenic and adipocyte differentiation in MC3T3-E1 cells
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In Vivo	<p>Adenosine N1-oxide (oral administration, 135 mg/kg for three times) reduces the lethality caused by LPS (HY-D1056)-induced endotoxin shock in BALB/c mice<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>																

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Animal Model:	LPS-induced endotoxin shock in BALB/c mice <sup>[1]</sup>
Dosage:	135 mg/kg for three times
Administration:	Oral administration
Result:	Reduced the lethality caused by LPS (HY-D1056)-induced endotoxin shock in BALB/c mice.

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## REFERENCES

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[1]. Kohno K, et al. Anti-inflammatory effects of adenosine N1-oxide. J Inflamm (Lond). 2015;12(1):2. Published 2015 Jan 20.

[2]. Ohashi E, et al. Adenosine N1-Oxide Exerts Anti-inflammatory Effects through the PI3K/Akt/GSK-3 $\beta$  Signaling Pathway and Promotes Osteogenic and Adipocyte Differentiation. Biol Pharm Bull. 2019;42(6):968-976.

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

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