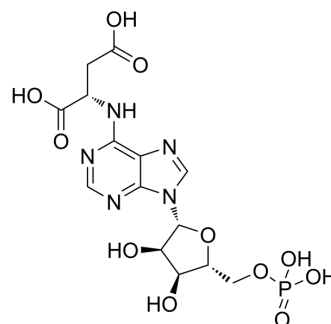


Adenylosuccinic acid

Cat. No.:	HY-127137
CAS No.:	19046-78-7
Molecular Formula:	C ₁₄ H ₁₈ N ₅ O ₁₁ P
Molecular Weight:	463.29
Target:	Endogenous Metabolite
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Adenylosuccinic acid (Adenylosuccinate; Aspartyl adenylate) is a purine ribonucleoside monophosphate and plays a role in nucleotide cycle metabolite. Adenylosuccinic acid can be converted into fumaric acid through adenylosuccinate lyase. Adenylosuccinic acid has the potential for the study of duchenne muscular dystrophy(DMD) ^[1] .								
IC₅₀ & Target	Human Endogenous Metabolite								
In Vivo	<p>Adenylosuccinic acid (oral administration; 3-3000 µg/mL; 6 weeks) significantly improves the features of murine DMD, it decreases the number of centronucleated fibres, lipid accumulation, connective tissue infiltration and Ca²⁺ content of mdx tibialis anterior^[1].</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>C57Bl/10ScSn (normal wild-type strain; Con) and C57Bl/10mdx (mdx) mice^[1]</td> </tr> <tr> <td>Dosage:</td> <td>3 µg/mL, 30 µg/mL, 300 µg/mL, 3000 µg/mL</td> </tr> <tr> <td>Administration:</td> <td>3 µg/mL for 3 days and 30 µg/mL for the next 4 days, and 300 µg/mL for one week, and then 3000 µg/mL for 6 weeks</td> </tr> <tr> <td>Result:</td> <td>Ameliorated the symptoms of murine duchenne muscular dystrophy.</td> </tr> </table>	Animal Model:	C57Bl/10ScSn (normal wild-type strain; Con) and C57Bl/10mdx (mdx) mice ^[1]	Dosage:	3 µg/mL, 30 µg/mL, 300 µg/mL, 3000 µg/mL	Administration:	3 µg/mL for 3 days and 30 µg/mL for the next 4 days, and 300 µg/mL for one week, and then 3000 µg/mL for 6 weeks	Result:	Ameliorated the symptoms of murine duchenne muscular dystrophy.
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REFERENCES

[1]. Timpani CA, et al. Adenylosuccinic acid therapy ameliorates murine Duchenne Muscular Dystrophy. *Sci Rep.* 2020 Jan 24;10(1):1125.

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

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