

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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# PRODUCT INFORMATION



## Memantine-d<sub>6</sub> (hydrochloride)

Item No. 25463

CAS Registry No.: 1189713-18-5

3,5-di(methyl-d<sub>3</sub>)-tricyclo[3.3.1.1<sup>3,7</sup>] Formal Name:

decan-1-amine, monohydrochloride

MF:  $C_{12}H_{15}D_6N \bullet HCI$ 

FW: 221.8

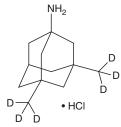
**Chemical Purity:** ≥98% (Memantine)

Deuterium

Incorporation:  $\geq$ 99% deuterated forms (d<sub>1</sub>-d<sub>6</sub>);  $\leq$ 1% d<sub>0</sub>

Supplied as: A solid Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



### **Laboratory Procedures**

Memantine-d<sub>6</sub> (hydrochloride) is intended for use as an internal standard for the quantification of memantine (Item No. 14184) by GC- or LC-MS. The accuracy of the sample weight in this vial is between 5% over and 2% under the amount shown on the vial. If better precision is required, the deuterated standard should be quantitated against a more precisely weighed unlabeled standard by constructing a standard curve of peak intensity ratios (deuterated versus unlabeled).

Memantine-d<sub>6</sub> (hydrochloride) is supplied as a solid. A stock solution may be made by dissolving the memantine-d, (hydrochloride) in the solvent of choice, which should be purged with an inert gas. Memantine-d<sub>6</sub> (hydrochloride) is soluble in DMSO and methanol.

### Description

Memantine is an NMDA receptor antagonist that blocks NMDA-induced currents in rat retinal ganglion cells by 90% when used at a concentration of 12  $\mu$ M. It reverses inhibition of dephosphorylation of the synthetic tau phosphopeptide p17 (tau<sub>194-207</sub>) induced by the endogenous inhibitor of protein phosphatase 2A (PP2A) I<sub>1</sub>PP2A in vitro.<sup>2</sup> In vivo, memantine (2 mg/kg) restores PP2A activity, decreases GSK-3β and amyloid- $\beta$  ( $\hat{A\beta}$ ) levels in the hippocampus, cerebral cortex, and ventricular areas, and attenuates spatial learning and memory deficits in the AAV1-I<sub>1</sub> PP2A rat model of Alzheimer's disease. Memantine (20 mg/kg) reduces responding on the ethanol-associated lever in a cue-induced ethanol-seeking test in rats.<sup>3</sup> It also decreases secretion of matrix metalloproteinase-9 (MMP-9), degradation of collagen IV, the size of cerebral ischemia-induced brain infarcts, and neuronal cell death in a mouse model of focal cerebral ischemia.<sup>4</sup>

#### References

- 1. Chen, H.S.V., Pellegrini, J.W., Aggarwal, S.K., et al. Open-channel block of N-methyl-D-aspartate (NMDA) responses by memantine: Therapeutic advantage against NMDA receptor-mediated neurotoxicity. J. Neurosci. 12(11), 4427-4436 (1992).
- 2. Wang, X., Blanchard, J., Grundke-Iqbal, I., et al. Memantine attenuates Alzheimer's disease-like pathology and cognitive impairment. PLoS One 10(12), e0145441, (2015).
- Vangeliene, V., Olevska, A., and Spanagel, R. Long-lasting effect of NMDA receptor antagonist memantine on ethanol-cue association and relapse. J. Neurochem. 135(6), 1080-1085 (2015).
- Chen, Z.-Z., Yang, D.-D., Zhao, Z., et al. Memantine mediates neuroprotection via regulating neurovascular unit in a mouse model of focal cerebral ischemia. Life Sci. 150, 8-14 (2016).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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