

Produktinformation



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SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic in



Product Information



Edrophonium (chloride)

Item No. 15928

CAS Registry No.: 116-38-1

N-ethyl-3-hydroxy-N,N-dimethyl-Formal Name:

benzenaminium, monochloride

Synonyms: Enlon®, Tensilon® MF: C₁₀H₁₆NO • Cl

FW: 201.7 **Purity:** ≥95%

Stability: ≥2 years at -20°C Supplied as: A crystalline solid UV/Vis.: λ_{max} : 223, 277 nm

Laboratory Procedures

For long term storage, we suggest that edrophonium (chloride) be stored as supplied at -20°C. It should be stable for at least two years.

Edrophonium (chloride) is supplied as a crystalline solid. A stock solution may be made by dissolving the edrophonium (chloride) in the solvent of choice. Edrophonium (chloride) is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide, which should be purged with an inert gas. The solubility of edrophonium (chloride) in these solvents is approximately 10, 2, and 0.5 mg/ml, respectively.

Further dilutions of the stock solution into aqueous buffers or isotonic saline should be made prior to performing biological experiments. Ensure that the residual amount of organic solvent is insignificant, since organic solvents may have physiological effects at low concentrations. Organic solvent-free aqueous solutions of edrophonium (chloride) can be prepared by directly dissolving the crystalline solid in aqueous buffers. The solubility of edrophonium (chloride) in PBS, pH 7.2, is approximately 10 mg/ml. We do not recommend storing the aqueous solution for more than one day.

Edrophonium (chloride) is an acetylcholinesterase (AChE) inhibitor that is known to prevent the breakdown of the neurotransmitter acetylcholine by binding specifically to its catalytic site. 1 It has been shown to inhibit AChE activity in human red blood cells, purified calf forebrain, and purified octopus brain with K; values of 0.2, 0.2, and 0.4 μM, respectively.² Edrophonium is often used as part of a battery of pharmacological tests to confirm a diagnosis of the autoimmune neuromuscular junction disorder, myasthenia gravis.3

References

- 1. Roufogalis, B.D. and Wickson, V.M. Acetylcholinesterase: specificity of the peripheral anionic site for cholinergic ligands. Mol. Pharm. 11, 352-60 (2011).
- Boyle, N.A., Talesa, V., Giovannini, E., et al. Synthesis and study of thiocarbonate derivatives of choline as potential inhibitors of acetylcholinesterase. J. Med. Chem. 40(19), 3009-3013 (1997).
- Juel, V.C. and Massey, J.M. Myasthenia gravis. Orphanet J. Rare Dis. 2, 44 (2007).

Related Products

For a list of related products please visit: www.caymanchem.com/catalog/15928

WARNING: This product is for laboratory research only: not for administration to humans. Not for human or veterinary DIAGNOSTIC OR THERAPEUTIC USE

This material should be considered hazardous until information to the contrary becomes available. Do not ingest, swallow, or inhale. Do not get in eyes, on skin, or on clothing. Wash thorou r handling. This information contains some, but not all, of the information required for the safe and proper use of this material. Before use, the user must review the complete Safety Data Sl which has been sent via email to your institution.

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Said refund or replacement is conditioned on Buyer giving written notice to Cayman within thirty (30) days after arrival of the material at its destination. Failure of Buyer to give said notice within rty (30) days shall constitute a waiver by Buyer of all claims hereunder with respect to said material.

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Cayman Chemical

Mailing address

1180 E. Ellsworth Road Ann Arbor, MI 48108 USA

Phone

(800) 364-9897 (734) 971-3335

(734) 971-3640

custserv@caymanchem.com

www.caymanchem.com