

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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PRODUCT INFORMATION



Lamotrigine-d2

Item No. 37196

CAS Registry No.: 1132746-94-1

Formal Name: 6-(5,6-dichlorophenyl-2,3,4-d₂)-1,2,4-

triazine-3,5-diamine

Synonym: LTG-d₂

MF: C₉H₄Cl₂D₃N₅

FW: 259.1

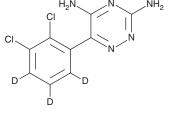
Chemical Purity: ≥98% (Lamotrigine)

Deuterium

Incorporation: \geq 99% deuterated forms (d₁-d₃); \leq 1% d₀

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

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



Laboratory Procedures

Lamotrigine-d₃ is intended for use as an internal standard for the quantification of lamotrigine (Item No. 15428) 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).

Lamotrigine- d_3 is supplied as a solid. A stock solution may be made by dissolving the lamotrigine- d_3 in the solvent of choice, which should be purged with an inert gas. Lamotrigine-d₃ is soluble in the organic solvent ethanol at a concentration of approximately 25 mg/ml.

Description

Lamotrigine is an anticonvulsant.¹ It inhibits voltage-gated sodium channels (Na_v) in HEK293 cells expressing recombinant human Na_v1.2, Na_v1.5, or Na_v1.8 (IC₅₀s = 10, 62, and 96 μ M, respectively), as well as high voltage-activated calcium currents in isolated rat cortical neurons (IC₅₀ = 12.3 μ M), an effect that can be reversed by the N-type calcium channel blocker ω-conotoxin GVIA (Item No. 24114) and P-type calcium channel blocker ω-agatoxin IVA (Item No. 21605).^{1,2} Lamotrigine protects against seizures induced by maximal electroshock (MES) in mice and rats (ED₅₀s = 10.1 and 7.4 μ mol/kg, respectively).³ It also decreases mechanical allodynia in a rat model of neuropathic pain induced by spinal nerve ligation $(ED_{50} = 47 \mu mol/kg)$. Formulations containing lamotrigine have been used in the treatment of epilepsy and bipolar disorder.

References

- 1. Drizin, I., Gregg, R.J., Scanio, M.J., et al. Discovery of potent furan piperazine sodium channel blockers for treatment of neuropathic pain. Bioorg. Med. Chem. 16(12), 6379-6386 (2008).
- Stefani, A., Spadoni, F., Siniscalchi, A., et al. Lamotrigine inhibits Ca²⁺ currents in cortical neurons: Functional implications. Eur. J. Pharmacol. 307(1), 113-116 (1996).
- 3. Miller, A.A., Wheatley, P., Sawyer, D.A., et al. Pharmacological studies on lamotrigine, a novel potential antiepileptic drug: I. Anticonvulsant profile in mice and rats. Epilepsia 27(5), 483-489 (1986).

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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