PRODUCT INFORMATION



Tolperisone-d₁₀ (hydrochloride)

Item No. 22592

CAS Registry No.: 1185160-65-9

Formal Name: 3-2,2,3,3,4,4,5,5,6,6-decadeuteriopiperidin-1-

yl)-2-methyl-1-(4-methylphenyl)propan-1-one,

monohydrochloride)

MF: C₁₆H₁₃D₁₀NO ● HCI

291.9 FW:

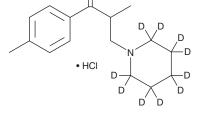
Chemical Purity: ≥98% (Tolperisone)

Deuterium

Incorporation: \geq 99% deuterated forms (d₁-d₁₀); \leq 1% d₀

Supplied as: A solid 2-8°C Storage: ≥4 years Stability:

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



Laboratory Procedures

Tolperisone-d₁₀ is intended for use as an internal standard for the quantification of tolperisone 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).

Description

Tolperisone is a piperidine that acts as an inhibitor of voltage-gated sodium channels with an IC₅₀ value of 198 µM, for inhibition of peak sodium currents evoked by step depolarizations in dorsal root ganglion (DRG) neurons, in vitro. Tolperisone acts as an antispastic agent that attenuates monosynaptic, disynaptic, and polysynaptic reflux response to dorsal root stimulation in vivo. Formulations containing tolperisone are used as analgesics and muscle relaxants.

Reference

1. Kocsis, P., Farkas, S., Fodor, L., et al. Tolperisone-type drugs inhibit spinal reflexes via blockade of voltagegated sodium and calcium channels. J. Pharmacol. Exp. Ther. 315(3), 1237-1246 (2005).

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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