PRODUCT INFORMATION



Raclopride

Item No. 17422

CAS Registry No.: 84225-95-6

3,5-dichloro-N-[[(2S)-1-ethyl-2-pyrrolidinyl] Formal Name:

methyl]-2-hydroxy-6-methoxy-benzamide

Synonym: (S)-(-)-Raclopride

MF: C₁₅H₂₀Cl₂N₂O₃

347.2 FW: **Purity:** ≥98% 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

Raclopride is supplied as a solid. A stock solution may be made by dissolving the raclopride in the solvent of choice, which should be purged with an inert gas. Raclopride is soluble in organic solvents such as ethanol and DMSO. The solubility of raclopride in these solvents is approximately 20 and 100 mM, respectively.

Description

Raclopride is a selective dopamine D_2/D_3 receptor antagonist with K_i values of 1.8, 3.5, 2,400, and 18,000 nM for D_2 , D_3 , D_4 , and D_1 receptors, respectively. It passes the blood brain barrier and can be used in in vivo binding and autoradiography studies of the dopamine system under normal and pathological conditions such as Huntington's disease.^{2,3}

References

- 1. Seeman, P. and Van Tol, H.H.M. Dopamine receptor pharmacology. Trends Pharmacol. Sci. 15(7), 264-270 (1994).
- 2. Lidow, M.S., Goldman-Rakic, P.S., Rakic, P., et al. Dopamine D2 receptors in the cerebral cortex: Distribution and pharmacological characterization with [3H]raclopride. Proc. Natl. Acad. Sci. USA 86(16), 6412-6416 (1989).
- 3. Ginovart, N. Imaging the dopamine system with in vivo [11C] raclopride displacement studies: Understanding the true mechanism. Mol. Imaging Biol. 7(1), 45-52 (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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CAYMAN CHEMICAL

1180 EAST ELLSWORTH RD ANN ARBOR, MI 48108 · USA PHONE: [800] 364-9897

[734] 971-3335

FAX: [734] 971-3640 CUSTSERV@CAYMANCHEM.COM WWW.**CAYMANCHEM**.COM