PRODUCT INFORMATION



Tenocyclidine (hydrochloride)

Item No. 17014

CAS Registry No.: 1867-65-8

1-[1-(2-thienyl)cyclohexyl]-piperidine, Formal Name:

monohydrochloride

Synonym:

MF: C₁₅H₂₃NS ● HCl

285.9 FW: **Purity:** ≥98% UV/Vis.: λ_{max} : 235 nm Supplied as: A crystalline solid

Storage: -20°C Stability: ≥5 years

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



Tenocyclidine (TCP), known more formally as N-[1-(2-thienyl)cyclohexyl] piperidine, is an analog of phencyclidine (PCP) in which the phenyl substituent is replaced with a thiophene group. TCP is a potent antagonist of the NMDA receptor-gated ion channel ($K_d = 9 \text{ nM}$). TCP blocks NMDA-activated excitatory postsynaptic potential in brain more potently than PCP. PCP and many analogs, including TCP, block the uptake and enhance the release of dopamine both in vitro and in vivo.3 TCP is regulated as a Schedule I compound in the United States. This product is intended for forensic and research applications.

References

- 1. Yeh, G.C., Bonhaus, D.W., Nadler, J.V., et al. N-methyl-D-aspartate receptor plasticity in kindling: Quantitative and qualitative alterations in the N-methyl-p-aspartate receptor-channel complex. Proc. Natl. Acad. Sci. USA 86(20), 8157-8160 (1989).
- 2. Bartschat, D.K. and Blaustein, M.P. Phencyclidine in low doses selectively blocks a presynaptic voltage-regulated potassium channel in rat brain. Proc. Natl. Acad. Sci. USA 83(1), 189-192 (1986).
- Chaudieu, I., Vignon, J., Chicheportiche, M., et al. Role of the aromatic group in the inhibition of phencyclidine binding and dopamine uptake by PCP analogs. Pharmacol. Biochem. Behav. 32(3), 699-705 (1989).

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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