

CVN417

## Chemical Properties

CAS No. :

Formula:

Molecular Weight:

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Actual storage temperature shall be subject to the COA.

## Biological Description

Description	CVN417 is an orally active antagonist of nAChR containing the $\alpha 6$ subunit, modulating phasic dopaminergic neurotransmission in an impulse-dependent fashion. It inhibits Ca (2+) efflux through nAChR subunits, displaying IC50 values of 0.086 $\mu$ M ( $\alpha 6$ ), 2.56 $\mu$ M ( $\alpha 3$ ), and 0.657 $\mu$ M ( $\alpha 4$ ). CVN417 has shown efficacy in reducing resting tremor in rodent models, suggesting a potential to ameliorate movement disorders in conditions like Parkinson's disease [1].
Targets(IC50)	Others,AChR
In vitro	CVN417 (10 $\mu$ M; 0-2 h) exhibits lower metabolic turnover in human liver microsomes or hepatocytes [1]. In vitro ADME data for CVN417 [1] show intrinsic clearance rates (Cl int) in human liver microsomes of 2.8 $\mu$ L/min/mg, in rat 31.2, mouse 33.3, and dog 27.7; and in hepatocytes (mL/min/10 <sup>6</sup> cells) in human 3.7, rat 20.8, mouse 25.1, and dog 32.3.

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Tel:781-999-4286

E\_mail:info@targetmol.com

Address:34 Washington Street,Wellesley Hills,MA 02481