## MCE RedChemExpress

## **Tiapride**

 $\begin{array}{lll} \textbf{Cat. No.:} & \textbf{HY-B1196A} \\ \textbf{CAS No.:} & 51012-32-9 \\ \textbf{Molecular Formula:} & \textbf{C}_{15}\textbf{H}_{24}\textbf{N}_2\textbf{O}_4\textbf{S} \\ \end{array}$ 

Molecular Weight: 328.43

Target: Dopamine Receptor

Pathway: GPCR/G Protein; Neuronal Signaling

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	Tiapride, an atypical neuroleptic agent, is a selective dopamine D2-receptor antagonist with little propensity for causing catalepsy and sedation. The IC <sub>50</sub> values of Tiapride are 1440, 45.8, >100, and 11.7 $\mu$ M for D1; D2; D3; D4, respectively <sup>[1]</sup> .	
IC <sub>50</sub> & Target	D <sub>2</sub> Receptor 110-320 nM (IC <sub>50</sub> )	D <sub>3</sub> Receptor 180 nM (IC <sub>50</sub> )

## **REFERENCES**

[1]. D H Peters, et al. Tiapride. A review of its pharmacology and therapeutic potential in the management of alcohol dependence syndrome. Drugs. 1994 Jun;47(6):1010-32.

[2]. T Arima, et al. Comparison of effects of tiapride and sulpiride on D-1, D-2, D-3 and D-4 subtypes of dopamine receptors in rat striatal and bovine caudate nucleus membranes. Jpn J Pharmacol. 1986 Jul;41(3):419-23.

Caution: Product has not been fully validated for medical applications. For research use only.

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