

## **Data Sheet**

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**Product Name** : SCH23390 hydrochloride

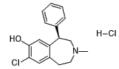
 Cat. No.
 :
 PC-22147

 CAS No.
 :
 125941-87-9

 Molecular Formula
 :
 C<sub>17</sub>H<sub>19</sub>Cl<sub>2</sub>NO

 Molecular Weight
 :
 324.25

Target : Dopamine Receptor Solubility : 10 mM in DMSO



## **Biological Activity**

SCH23390 hydrochloride is a potent and selective dopamine D1-like receptor antagonist with Kis of 0.2 nM and 0.3 nM for the D1 and D5 receptor, respectively.

SCH23390 shows high affinity to human 5-HT2C receptor with a Ki of 9.3 nM, also binds with high affinity to the 5-HT2 and 5-HT1C receptors.

SCH23390 inhibits G protein-coupled inwardly rectifying potassium (GIRK) channel with IC50 of 268 nM.

SCH23390 (1 uM) reverses the inhibitory effects of Isosibiricin on NLRP3 expression and the cleavages of caspase-1 and IL- $1\beta$  in the LPS-induced BV-2 cells.

SCH23390 abolishes generalized seizures evoked by the chemoconvulsants.

## References

Bourne JA, et al. CNS Drug Rev. 2001 Winter;7(4):399-414.

Millan MJ, et al. Psychopharmacology (Berl). 2001 Jun;156(1):58-62.

Kuzhikandathil EV, et al. Mol Pharmacol. 2002 Jul;62(1):119-26.

Caution: Product has not been fully validated for medical applications. Lab Use Only!

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