# Tranylcypromine hemisulfate

| Cat. No.:          | HY-B1496  |  |
|--------------------|---|--|
| CAS No.:           | 13492-01-8  | $\wedge$   |
| Molecular Formula: | C <sub>9</sub> H <sub>12</sub> NO <sub>2</sub> S <sub>0.5</sub>   |  |
| Molecular Weight:  | 182.23  | $\begin{bmatrix} \\ \end{bmatrix}$ , $\mathbb{NH}_2$ |
| Target:            | Monoamine Oxidase; Histone Demethylase  |  |
| Pathway:           | Neuronal Signaling; Epigenetics   |  |
| Storage:           | 4°C, sealed storage, away from moisture<br>* In solvent : -80°C, 1 year; -20°C, 6 months (sealed storage, away from moisture) | 0.5H <sub>2</sub> SO <sub>4</sub>                    |

# SOLVENT & SOLUBILITY

|  | Preparing<br>Stock Solutions  | Solvent Mass<br>Concentration | 1 mg      | 5 mg       | 10 mg      |  |
|--|---|-------------------------------|-----------|------------|------------|--|
|  |   | 1 mM                          | 5.4876 mL | 27.4379 mL | 54.8757 mL |  |
|  |   | 5 mM                          | 1.0975 mL | 5.4876 mL  | 10.9751 mL |  |
|  |   | 10 mM                         | 0.5488 mL | 2.7438 mL  | 5.4876 mL  |  |
|  | Please refer to the solubility information to select the appropriate solvent. |                               |           |            |            |  |

| BIOLOGICAL ACTIVITY |  |  |  |
|---------------------|--|--|--|
|                     |  |  |  |
| Description         | Tranylcypromine (SKF 385) hemisulfate is an irreversible, nonselective monoamine oxidase (MAO) inhibitor used in the treatment of depression. Tranylcypromine hemisulfate is also a lysine-specific demethylase 1 (LSD1) inhibitor, suppresses lesion growth and improves generalized hyperalgesia in mouse with induced endometriosis. Tranylcypromine has antidepressant effects <sup>[1][2]</sup> .   |  |  |
| $IC_{50}$ & Target  | KDM1/LSD1  |  |  |
| In Vitro            | Tranylcypromine (10 nM to 10 μM) exerts neuroprotective effects against toxicity induced by human Aβ(1-42) oligomers<br>independently from the presence of glial cells <sup>[1]</sup> . Tranylcypromine (100 μM) significantly protects RGCs from glutamate<br>neurotoxicity-induced apoptosis as well as apoptosis induced by oxidative stress. Tranylcypromine promotes mitogen-<br>activated protein kinase 12 (p38 MAPKγ) expression under conditions of glutamate (Glu)-induced stress. Besides,<br>tranylcypromine contributes to RGC survival via alterations of p38 MAPKγ activity <sup>[3]</sup> .<br>MCE has not independently confirmed the accuracy of these methods. They are for reference only. |  |  |

**Product** Data Sheet



Tranylcypromine treatment significantly and substantially reduces the lesion size and improves generalized hyperalgesia in a dose-dependent fashion in mice with induced endometriosis. In addition, tranylcypromine treatment results in reduced immunoreactivity to biomarkers of proliferation, angiogenesis, and H3K4 methylation, leading to arrested EMT and lesion growth<sup>[2]</sup>. Tranylcypromine (500 mM) injection exerts neuroprotective effects within intracellular apoptotic signaling pathways and suppresses morphologic changes in the retina of the rat, suppresses caspase 3 activity and recovers p38 MAPK $\gamma$  expression in the retina after NMDA-induced injury, and enhances RGC survival after retinal injury via the attenuation of NMDA neurotoxicity<sup>[3]</sup>. Tranylcypromine (10 µg/g) causes an approximate and significant doubling of labeled cells in the combined brain regions examined, as detected by BrdU immunohistochemistry. Tranylcypromine causes the greatest increase in cell proliferation in the cerebellum<sup>[4]</sup>.

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## PROTOCOL

#### Animal Administration <sup>[3]</sup>

Briefly, the rats are anesthetized with an intraperitoneal injection of a 1:1 mixture of xylazine hydrochloride (4 mg/kg) and ketamine hydrochloride (10 mg/kg). Then, the pupil is dilated with phenylephrine hydrochloride and tropicamide eye drops, and 20 nmol NMDA with or without tranylcypromine is injected into the vitreous cavity. To assess the inhibitory effect of mitogen-activated protein kinase (MAPK), 100 nmol BIRB796 is intravitreally injected at the same time of NMDA injection. The injections are performed under a microscope using a 33-gauge needle connected to a microsyringe; the needle is inserted approximately 1.0 mm behind the corneal limbus. Next, either PBS (vehicle control) or 500 mM tranylcypromine (1000 nmol) mixed with 10 mM NMDA (20 nmol) in a total volume of 2.0 µL is injected into the vitreous cavity. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **CUSTOMER VALIDATION**

- Biomaterials. 2018 Dec 6;193:30-46.
- Biol Reprod. 2020 Dec 1;103(6):1229-1237.
- Biochem Biophys Res Commun. 2019 May 14;512(4):852-858.
- Patent. US20180263995A1.

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#### REFERENCES

[1]. Caraci F, et al. Neuroprotective effects of the monoamine oxidase inhibitor tranylcypromine and its amide derivatives against A $\beta$ (1-42)-induced toxicity. Eur J Pharmacol. 2015 Oct 5;764:256-263.

[2]. Sun Q, et al. Tranylcypromine, a lysine-specific demethylase 1 (LSD1) inhibitor, suppresses lesion growth and improves generalized hyperalgesia in mouse with induced endometriosis. Reprod Biol Endocrinol. 2016 Apr 9;14:17.

[3]. Tsutsumi T, et al. Potential Neuroprotective Effects of an LSD1 Inhibitor in Retinal Ganglion Cells via p38 MAPK Activity. Invest Ophthalmol Vis Sci. 2016 Nov 1;57(14):6461-6473.

[4]. Romanczyk TB, et al. The antidepressant tranylcypromine alters cellular proliferation and migration in the adult goldfish brain. Anat Rec (Hoboken). 2014 Oct;297(10):1919-26.

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

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