## **Product** Data Sheet

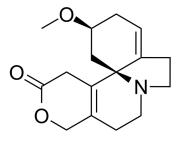
# Dihydro-β-erythroidine hydrobromide

Cat. No.: HY-107670 CAS No.: 29734-68-7 Molecular Formula: C<sub>16</sub>H<sub>22</sub>BrNO<sub>3</sub> Molecular Weight: 356.25 nAChR Target:

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

4°C, sealed storage, away from moisture Storage:

\* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



**HBr** 

## **SOLVENT & SOLUBILITY**

In Vitro

H<sub>2</sub>O: 35 mg/mL (98.25 mM; Need ultrasonic and warming) DMSO: 8 mg/mL (22.46 mM; Need ultrasonic and warming)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.8070 mL	14.0351 mL	28.0702 mL
	5 mM	0.5614 mL	2.8070 mL	5.6140 mL
	10 mM	0.2807 mL	1.4035 mL	2.8070 mL

Please refer to the solubility information to select the appropriate solvent.

### **BIOLOGICAL ACTIVITY**

Description

Dihydro-β-erythroidine (DHβE) hydrobromide is a potent, orally active, and competitive antagonist of neuronal nAChRs. Dihydro- $\beta$ -erythroidine hydrobromide shows selectivity for  $\alpha4\beta4$  and  $\alpha4\beta2$  nAChRs, with IC<sub>50</sub>s of 0.19 and 0.37  $\mu$ M, respectively. Antidepressant-like activities<sup>[1][2][3]</sup>.

In Vivo

Dihydro-Beta-Erythroidine hydrobromide (DH $\beta$ E hydrobromide; 10 nmol/0.5  $\mu$ L) is infused into the respective areas prior to a systemic nicotine injection (0.2 mg/kg; SC). DHßE infused into the ventral tegmental area (VTA), nucleus accumbens (NAcc), or infralimbic (IL) cortex, but not prelimbic (PrL) cortex, attenuated nicotine-enhanced responding for a conditioned reinforcer (CRf)<sup>[2]</sup>.

The co-administration of Dihydro-Beta-Erythroidine hydrobromide (5.0 mg/kg; s.c.) with nicotine (0.2 and 0.4 mg/kg; s.c.) prevents the development of conditioned taste aversions (CTAs)<sup>[4]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male hooded Lister rats <sup>[4]</sup>
Dosage:	0.5, 1.6 and 5.0 mg/kg

Administration:	S.c.
Result:	A complete blockade of the nicotine effect produced by the training dose of nicotine (0 mg/kg).

#### **REFERENCES**

- [1]. Harvey SC, et al. Multiple determinants of dihydro-beta-erythroidine sensitivity on rat neuronal nicotinic receptor alpha subunits. J Neurochem. 1996 Nov;67(5):1953-9.
- [2]. Tabbara RI, et al. Nicotine enhances responding for conditioned reinforcement via  $\alpha 4\beta 2$  nicotinic acetylcholine receptors in the ventral tegmental area, but not the nucleus accumbens or the prefrontal cortex. Neuropharmacology. 2019 Apr;148:68-76.
- [3]. Clementson S, et al. Enantioselective Total Synthesis of (+)-Dihydro-β-erythroidine. J Am Chem Soc. 2019 Jun 5;141(22):8783-8786.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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