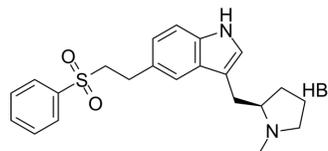


## Eletriptan hydrobromide

<b>Cat. No.:</b>	HY-A0010
<b>CAS No.:</b>	177834-92-3
<b>Molecular Formula:</b>	C <sub>22</sub> H <sub>27</sub> BrN <sub>2</sub> O <sub>2</sub> S
<b>Molecular Weight:</b>	463.43
<b>Target:</b>	5-HT Receptor
<b>Pathway:</b>	GPCR/G Protein; Neuronal Signaling
<b>Storage:</b>	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 100 mg/mL (215.78 mM; Need ultrasonic)					
	<b>Preparing Stock Solutions</b>	<b>Solvent</b>	<b>Mass</b>	<b>1 mg</b>	<b>5 mg</b>	<b>10 mg</b>
		<b>Concentration</b>				
		<b>1 mM</b>		2.1578 mL	10.7891 mL	21.5782 mL
		<b>5 mM</b>		0.4316 mL	2.1578 mL	4.3156 mL
	<b>10 mM</b>		0.2158 mL	1.0789 mL	2.1578 mL	
Please refer to the solubility information to select the appropriate solvent.						
<b>In Vivo</b>	<ol style="list-style-type: none"> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 40% PEG300 &gt;&gt; 5% Tween-80 &gt;&gt; 45% saline Solubility: ≥ 2.75 mg/mL (5.93 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.75 mg/mL (5.93 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2.75 mg/mL (5.93 mM); Clear solution</li> </ol>					

### BIOLOGICAL ACTIVITY

<b>Description</b>	Eletriptan hydrobromide (Eletriptan HBr) is a selective 5-HT <sub>1B</sub> and 5-HT <sub>1D</sub> receptor agonist with K <sub>i</sub> of 0.92 nM and 3.14 nM, respectively.	
<b>IC<sub>50</sub> &amp; Target</b>	5-HT <sub>1B</sub> Receptor 0.92 nM (K <sub>i</sub> )	5-HT <sub>1D</sub> Receptor 3.14 nM (K <sub>i</sub> )

### REFERENCES

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- [1]. Napier C, et al. Characterisation of the 5-HT receptor binding profile of eletriptan and kinetics of [3H]eletriptan binding at human 5-HT<sub>1B</sub> and 5-HT<sub>1D</sub> receptors. *Eur J Pharmacol*, 1999, 368(2-3), 259-268.
- [2]. MaassenVanDenBrink A, et al. Craniovascular selectivity of eletriptan and sumatriptan in human isolated blood vessels. *Neurology*, 2000, 55(10), 1524-1530.
- [3]. Gupta P, et al. The in vivo pharmacological profile of eletriptan (UK-116,044): a potent and novel 5-HT<sub>1B/1D</sub> receptor agonist. *Eur J Pharmacol*, 2000, 398(1), 73-81.
- [4]. Hoskin KL, et al. The 5-hydroxytryptamine<sub>1B/1D/1F</sub> receptor agonists eletriptan and naratriptan inhibit trigeminovascular input to the nucleus tractus solitarius in the cat. *Brain Res*, 2004, 998(1), 91-99.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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