



## **Cutamesine dihydrochloride**

**Catalog No: tcsc3288** 

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## **Available Sizes**

Size: 10mg

Size: 50mg



## **Specifications**

CAS No:

165377-44-6

Formula:

 $C_{23}H_{34}CI_2N_2O_2$ 

**Pathway:** 

GPCR/G Protein

**Target:** 

Sigma Receptor

**Purity / Grade:** 

>98%

**Solubility:** 

DMSO: 30 mg/mL (67.96 mM; Need ultrasonic and warming)

**Alternative Names:** 

SA4503 (dihydrochloride); AGY94806 dihydrochloride

**Observed Molecular Weight:** 

441.43

## **Product Description**

Cutamesine dihydrochloride (SA4503 dihydrochloride) is a potent **Sigma 1** receptor agonist with an  $IC_{50}$  of 17.4 nM in guinea pig brain membranes.



IC50 & Target: IC50: 17.4 nM (σ1receptor, guinea pig brain membranes)[1]

In Vitro: The sigma receptor might be involved in several diseases in the central nervous system. Cutamesine, a potent  $\sigma$ 1 receptor agonist, has 103-fold higher affinity for  $\sigma$ 1 (IC<sub>50</sub>=17.4 nM) than  $\sigma$ 2 (IC<sub>50</sub>=1,784 nM) sites in guinea pig brain membranes. Cutamesine is 14-fold selective for  $\sigma$ 1 (K<sub>i</sub>=4.6 nM) over  $\sigma$ 2 (K<sub>i</sub>=63.1 nM) sites in guinea pig brain homogenates<sup>[1]</sup>. Cutamesine protects motor neuron NSC34 cells against superoxide dismutase 1 and serum free neurotoxicity. It upregulates the phosphorylation levels of Akt and extracellular signal-regulated kinase (ERK) 1/2<sup>[2]</sup>. Cutamesine reduces the activation of the MAPK/ERK pathway and down-regulated the ionotropic glutamate receptor, GluR1<sup>[3]</sup>.

In Vivo: Cutamesine extends the survival time in the SOD1G93A mice<sup>[2]</sup>.

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!