Data Sheet (Cat.No.T17114)



Tolrestat

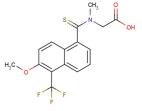
Chemical Properties

CAS No.: 82964-04-3

Formula: C16H14F3NO3S

Molecular Weight: 357.35 Appearance: N/A

Storage: 0-4°C for short term (days to weeks), or -20°C for long term (months).



Biological Description

Description	Tolrestat is a potent and orally active inhibitor of aldose reductase (IC50: 35 nM).	
Targets(IC ₅₀)	Aldose Reductase: 35 nM	
In vivo	Tolrestat (1.8 mg/kg per day) causes a reversal to normal RBC sorbitol levels of diabetic rats. The estimated ID in the sciatic nerve and lenses is 4.8 and about 20 for tolrestat, and 1.7 and 2.2 for (±)sorbinil, respectively in 21-day diabetic rats. Either tolrestat or sorbinil inhibits tissue AR activity but does not significantly affect plasma lipoprotein levels, or affect the bodyweight of the mice or their general health [1][2][3].	

Solubility Information

Solubility	DMSO: 100 mg/mL (279.84 mM) (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.798 mL	13.992 mL	27.984 mL
5 mM	0.56 mL	2.798 mL	5.597 mL
10 mM	0.28 mL	1.399 mL	2.798 mL
50 mM	0.056 mL	0.28 mL	0.56 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. The storage conditions and period of the stock solution: - 80 °C for 6 months; - 20 °C for 1 month. Please use it as soon as possible.

Reference

- 2. Simard-Duquesne N, et al. The effects of a new aldose reductase inhibitor (tolrestat) in galactosemic and diabetic rats. Metabolism. 1985 Oct;34(10):885-92.
- 3. Srivastava S, et al. Aldose reductase protects against early atherosclerotic lesion formation in apolipoprotein E-null mice. Circ Res. 2009 Oct 9;105(8):793-802.

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