# Data Sheet (Cat.No.T10250)



# ADRA1D receptor antagonist 1 HCl

#### **Chemical Properties**

CAS No.: 1191908-14-1

Formula: C15H14Cl2N4O

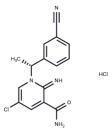
Molecular Weight: 337.2

Appearance: no data available

keep away from direct sunlight, keep away from

Storage: moisture

Powder: -20°C for 3 years | In solvent: -80°C for 1 year



### **Biological Description**

Description	ADRA1D receptor antagonist 1 HCl is an orally active, potent and selective α1D adrenergic receptor antagonist (Ki: 1.6 nM) with potential antiproliferative activity.		
Targets(IC50)	Adrenergic Receptor		
In vitro	ADRA1D receptor antagonist 1 exhibits low hERG inhibition and higher selectivity for $\alpha$ 1D-AR over $\alpha$ 1A- and $\alpha$ 1B-ARs, without affecting $\alpha$ 1A- and $\alpha$ 1B-ARs[1].		
In vivo	In the rat model of bladder outlet obstruction (BOO), ADRA1D receptor antagonist 1 (4.4 $\mu$ g/kg; intravenous injection) dose-dependently decreased the occurrence of non-voiding bladder contractions during the urinary storage phase[1].		
Animal Research	Animal Model: Rat with bladder outlet obstruction (BOO). Dosage: 4.4 µg/kg. Administration: Intravenous injection		

#### **Solubility Information**

Solubility	DMSO: 80 mg/mL (237.25 mM), Sonication is recommended.	
	(< 1 mg/ml refers to the product slightly soluble or insoluble)	

## **Preparing Stock Solutions**

	1mg	5mg	10mg
1 mM	2.9656 mL	14.828 mL	29.656 mL
5 mM	0.5931 mL	2.9656 mL	5.9312 mL
10 mM	0.2966 mL	1.4828 mL	2.9656 mL
50 mM	0.0593 mL	0.2966 mL	0.5931 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Page 1 of 2 www.targetmol.com

#### Reference

Sakauchi N,et al. Discovery of 5-Chloro-1-(5-chloro-2-(methylsulfonyl)benzyl)-2-imino-1,2-dihydropyridine-3carboxamide (TAK-259) as a Novel, Selective, and Orally Active α1D Adrenoceptor Antagonist with Antiurinary

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Page 2 of 2 www.targetmol.com

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