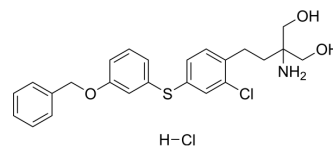


## Mocravimod hydrochloride

Cat. No.:	HY-13660
CAS No.:	509088-69-1
Molecular Formula:	C <sub>24</sub> H <sub>27</sub> Cl <sub>2</sub> NO <sub>3</sub> S
Molecular Weight:	480.45
Target:	LPL Receptor
Pathway:	GPCR/G Protein
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



### SOLVENT & SOLUBILITY

In Vitro	DMSO : 200 mg/mL (416.28 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.0814 mL	10.4069 mL	20.8138 mL
		5 mM	0.4163 mL	2.0814 mL	4.1628 mL
		10 mM	0.2081 mL	1.0407 mL	2.0814 mL
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	<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: ≥ 2.5 mg/mL (5.20 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (5.20 mM); Suspended solution; Need ultrasonic</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2.5 mg/mL (5.20 mM); Clear solution</li> </ol>				

### BIOLOGICAL ACTIVITY

Description	Mocravimod hydrochloride (KRP-203), an immunosuppressant, is a potent and orally active S1PR1 (sphingosine 1-phosphate receptor type 1) agonist <sup>[1][2]</sup> .
IC <sub>50</sub> & Target	S1PR1 <sup>[1]</sup>
In Vivo	<p>Mocravimod hydrochloride (KRP-203) ameliorates atherosclerosis in LDL-R / Mice<sup>[1]</sup>.</p> <p>Mocravimod hydrochloride (KRP-203) (orally; 0.1 and 1 mg/kg/day; for 100 days) prolongs graft survival and attenuates chronic rejection in mHC-disparate rat heart allografts<sup>[2]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>

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Animal Model:	Inbred male DA (MHC haplotype: RT1 <sup>a</sup> ) rats <sup>[2]</sup>
Dosage:	0.1 and 1 mg/kg
Administration:	Orally; daily; for 100 days
Result:	Prolonged graft survival and attenuated chronic rejection in mHC-disparate rat heart allografts.

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## REFERENCES

- [1]. Potì F, et al. KRP-203, sphingosine 1-phosphate receptor type 1 agonist, ameliorates atherosclerosis in LDL-R<sup>-/-</sup> mice. *Arterioscler Thromb Vasc Biol.* 2013 Jul;33(7):1505-12.
- [2]. Shimizu H, et al. KRP-203, a novel synthetic immunosuppressant, prolongs graft survival and attenuates chronic rejection in rat skin and heart allografts. *Circulation.* 2005 Jan 18;111(2):222-9. Epub 2005 Jan 10.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA