Product Data Sheet

Montelukast sodium

Cat. No.: HY-13315 CAS No.: 151767-02-1 Molecular Formula: $C_{35}H_{35}CINNaO_3S$

Molecular Weight: 608.17

Target: Leukotriene Receptor Pathway: GPCR/G Protein

Storage: 4°C, sealed storage, away from moisture

* In solvent: -80°C, 2 years; -20°C, 1 year (sealed storage, away from moisture)

SOLVENT & SOLUBILITY

 $H_2O : \ge 50 \text{ mg/mL } (82.21 \text{ mM})$ In Vitro

> DMSO: 50 mg/mL (82.21 mM; Need ultrasonic) * "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.6443 mL	8.2214 mL	16.4428 mL
	5 mM	0.3289 mL	1.6443 mL	3.2886 mL
	10 mM	0.1644 mL	0.8221 mL	1.6443 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

1. Add each solvent one by one: PBS

Solubility: 1.25 mg/mL (2.06 mM); Suspended solution; Need ultrasonic

Montelukast (10 $\mu\text{M};$ 18 h) modulates the activation of MMP-9 $^{[3]}.$

BIOLOGICAL ACTIVITY

Description	Montelukast sodium (MK0476) is a potent, selective and orally active antagonist of cysteinyl leukotriene receptor 1 (CysLT ₁). Montelukast sodium can be used for the reseach of asthma and liver injury. Montelukast sodium also has an antioxidant effect in intestinal ischemia-reperfusion injury, and could reduce cardiac damage. Montelukast sodium decreases eosinophil infiltration into the asthmatic airways. Montelukast sodium can also be used for COVID-19 research ^{[1][2][3][4]} .
IC ₅₀ & Target	CysLT ₁
In Vitro	Montelukast (5 μ M; 1 h) inhibits APAP (Acetaminophen) (HY-66005)-induced cell damage ^[1] . Montelukast (0.01-10 μ M; 30 min) diminishes the 5-oxo-ETE-induced cell migration and modulates the activation of the plasmin-plasminogen system ^[3] .

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Migration Assay ^[3]			
Cell Line:	Eosinophils		
Concentration:	0.01-10 μΜ		
Incubation Time:	30 min		
Result:	Diminished the 5-oxo-ETE–induced cell migration.		
Western Blot Analysis ^[3]			
Cell Line:	Eosinophils		
Concentration:	10 μΜ		
Incubation Time:	18 h		
Result:	Reduced the 5-oxo-ETE-boosted MMP-9 secretion.		

In Vivo

 $Montelukast~(3~mg/kg;~oral~gavage)~protects~against~APAP-induced~hepatotoxicity~in~mice^{[1]}.$

Montelukast (1 mg/kg; miniosmotic pump administration) reduces the airway remodeling changes observed in OVA-treated mice and blocks the actions of cysteinyl leukotrienes (LT) C4, D4, and E4 mediated by the CysLT1 receptor^[2].

Montelukast (1 mg/kg; miniosmotic pump administration) reduces the elevated levels of IL-4 and IL-13 found in the BAL fluid of OVA-treated mice^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	C57BL/6J mice (8-week-old; 22-25 g) are induced acute hepatic injury $^{[1]}$		
Dosage:	3 mg/kg		
Administration:	Oral gavage 1 h after saline or APAP administration		
Result:	Decreased serum levels of alanine transaminase (ALT) and aspartate aminotransferase (AST), and alleviated liver damage.		

CUSTOMER VALIDATION

- J Cachexia Sarcopenia Muscle. 2022 Jun 9.
- Artif Cell Nanomed B. 2019 Dec;47(1):4234-4239.
- Eur J Pharmacol. 2022 May 15;923:174892.
- Naunyn Schmiedebergs Arch Pharmacol. 2023 Feb 27.
- Patent. US20230404992A1.

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REFERENCES

- $[1]. \ Langlois\ A, et\ al.\ Montelukast\ regulates\ eosinophil\ protease\ activity\ through\ a\ leukotriene-independent\ mechanism.\ J\ Allergy\ Clin\ Immunol.\ 2006;118(1):113-119.$
- [2]. Khan AR, et al. Montelukast in hospitalized patients diagnosed with COVID-19. J Asthma. 2022 Apr;59(4):780-786.

[3]. Pu S, et, al. Montelukast Prevents Mice Against Acetaminophen-Induced Liver Injury. Front Pharmacol. 2019 Sep 18; 10:1070.						
[4]. William RHJ, et, al. A role for	r cysteinyl leukotrienes in air	way remodeling in a mouse asthi	ma model. Am J Respir Crit Care Med. 20	002 Jan 1; 165(1): 108-16.		
	Caution: Product has no	ot been fully validated for me	dical applications. For research use	only.		
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Page 3 of 3 www.MedChemExpress.com