# **Product** Data Sheet

# Bemnifosbuvir hemisulfate

Cat. No.: HY-137958
CAS No.: 2241337-84-6

Molecular Formula:  $C_{24}H_{33}FN_7O_7P._1/_2H_2O_4S$ 

Molecular Weight: 630.58

Target: HCV; SARS-CoV
Pathway: Anti-infection

**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: 180 mg/mL (285.45 mM; Need ultrasonic)

 $H_2O : \ge 100 \text{ mg/mL} (158.58 \text{ mM})$ 

\* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.5858 mL	7.9292 mL	15.8584 mL
	5 mM	0.3172 mL	1.5858 mL	3.1717 mL
	10 mM	0.1586 mL	0.7929 mL	1.5858 mL

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 5 mg/mL (7.93 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 5 mg/mL (7.93 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility:  $\geq 5~\rm mg/mL$  (7.93 mM); Clear solution

### **BIOLOGICAL ACTIVITY**

Description

Bemnifosbuvir hemisulfate (AT-527), a hemisulfate salt of AT-511, a guanosine nucleotide proagent, is a potent and orally active HCV viral replication inhibitor. Bemnifosbuvir hemisulfate is highly effective in the control of SARS-CoV-2 (COVID-19) infection in vitro (EC<sub>90</sub>=0.47  $\mu$ M). Bemnifosbuvir hemisulfate has pangenotypic antiviral activity<sup>[1][2][3]</sup>.

IC<sub>50</sub> & Target

EC50: 5-28 nM (HCV)<sup>[1]</sup>

EC90: 0.47 μM (SARS-CoV-2)<sup>[2]</sup>

In Vitro	Bemnifosbuvir hemisulfate has pan-genotypic antiviral activities that inhibits HCV genotype 1a (HCV GT1a), HCV GT1b, HCV GT2a, HCV GT3a, HCV GT4a, and HCV GT5a replication with $EC_{50}$ values of 12.8 nM, 12.5 nM, 9.2 nM, 10.3 nM, 14.7 nM, and 28.5 nM, respectively <sup>[1]</sup> . In normal human airway epithelial cells, the concentration of Bemnifosbuvir hemisulfate required to inhibit replication of SARS-CoV-2 by $EC_{90}$ is 0.47 $\mu$ M, very similar to its $EC_{90}$ against HCoV-229E, HCoV-OC43 and SARS-CoV in Huh-7 cells <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	When given orally to rats (500 mg/kg) and monkeys (30 mg/kg, 100 mg/kg or 300 mg/kg), Bemnifosbuvir hemisulfate preferentially delivers high levels of AT-9010 in the liver in vivo <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **CUSTOMER VALIDATION**

• bioRxiv. December 23, 2021.

See more customer validations on  $\underline{www.MedChemExpress.com}$ 

#### **REFERENCES**

- [1]. Steven S Good, et al. Preclinical evaluation of AT-527, a novel guanosine nucleotide prodrug with potent, pan-genotypic activity against hepatitis C virus. PLoS One. 2020 Jan 8;15(1):e0227104.
- [2]. Steven S Good, et al. AT-527, a double prodrug of a guanosine nucleotide analog, is a potent inhibitor of SARS-CoV-2 in vitro and a promising oral antiviral for treatment of COVID-19. Antimicrob Agents Chemother. 2021 Feb 8;AAC.02479-20.
- [3]. Elina Berliba, et al. Safety, pharmacokinetics and antiviral activity of AT-527, a novel purine nucleotide prodrug, in HCV-infected subjects with and without cirrhosis. Antimicrob Agents Chemother. 2019 Sep 30;63(12):e01201-19.

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

Page 2 of 2 www.MedChemExpress.com