ARN-3236

Cat. No.: HY-120856 CAS No.: 1613710-01-2 Molecular Formula: $C_{19}H_{16}N_2O_2S$

Molecular Weight: 336.41

Target: Salt-inducible Kinase (SIK) Pathway: Immunology/Inflammation

Powder -20°C Storage: 3 years

4°C 2 years

In solvent -80°C 6 months

> -20°C 1 month

Product Data Sheet

Solvent & Solubility

In Vitro DMSO: 130 mg/mL (386.43 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.9726 mL	14.8628 mL	29.7256 mL
	5 mM	0.5945 mL	2.9726 mL	5.9451 mL
	10 mM	0.2973 mL	1.4863 mL	2.9726 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: ≥ 2.17 mg/mL (6.45 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.17 mg/mL (6.45 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.17 mg/mL (6.45 mM); Clear solution

BIOLOGICAL ACTIVITY

Description ARN-3236 is an oral active and selective inhibitor of salt-inducible kinase 2 (SIK2), with IC50s of <1 nM, 21.63 nM and 6.63 nM for SIK2, SIK1 and SIK3, respectively. Has anti-cancer activity^{[1][2]}.

IC₅₀ & Target SIK2 SIK1 SIK3

<1 nM (IC₅₀) 21.63 nM (IC₅₀) 6.63 nM (IC₅₀)

REFERENCES
[1]. Lombardi MS, et al. SIK inhibition in human myeloid cells modulates TLR and IL-1R signaling and induces an anti-inflammatory phenotype. J Leukoc Biol. 2016 May;99(5):711-21.
[2]. Zhou J, et al. A Novel Compound ARN-3236 Inhibits Salt-Inducible Kinase 2 and Sensitizes Ovarian Cancer Cell Lines and Xenografts. Clin Cancer Res. 2017 Apr 15;23(8):1945-1954.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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