Proteins

Screening Libraries

Product Data Sheet

Aristeromycin

Cat. No.: HY-112639 CAS No.: 19186-33-5 Molecular Formula: C₁₁H₁₅N₅O₃ Molecular Weight: 265.27

Target: Bacterial Pathway: Anti-infection

Storage: Powder -20°C 3 years

> 2 years -80°C 6 months In solvent

> > -20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 50 mg/mL (188.49 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.7697 mL	18.8487 mL	37.6974 mL
	5 mM	0.7539 mL	3.7697 mL	7.5395 mL
	10 mM	0.3770 mL	1.8849 mL	3.7697 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: 2.5 mg/mL (9.42 mM); Suspended solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (9.42 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (9.42 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	Aristeromycin, an adenosine analog, is an antibiotic and a potent S-adenosylhomocysteine hydrolase (AHCY) inhibitor ^{[1][2]} .		
IC ₅₀ & Target	S-adenosylhomocysteine hydrolase $^{[1]}$		
In Vitro	The IC $_{50}$ value of Aristeromycin against AHCY is 38.5 nM at 50 μ M S-adenosylhomocysteine (SAH) (approximately equal to the Km: 48 μ M), but 271 nM at 1000 μ M SAH (20× Km). With 60 min of preincubation, the mean IC $_{50}$ value of Aristeromycin at 50 μ M SAH is 12.7 nM ^[1] .		

Aristeromycin has IC_{50} values of 3.2 μ M for LNCaP-FGC cell growth and 0.88 μ M for LNCaP-hr cell growth [1]. At least in part, Aristeromycin can regulate oncogenic EZH2 expression by inducing miR-26a [1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Uchiyama N, et al. Aristeromycin and DZNeP cause growth inhibition of prostate cancer via induction of mir-26a. Eur J Pharmacol. 2017 Oct 5;812:138-146.

[2]. Ishikura T, et al. Inhibition of S-adenosylhomocysteine hydrolase by purine nucleoside analogues. Nucleic Acids Symp Ser. 1983;(12):119-22.

Caution: Product has not been fully validated for medical applications. For research use only.

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