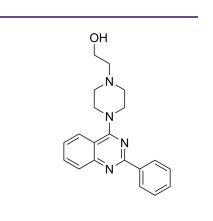
## **BVDV-IN-1**

Cat. No.:	HY-131976
CAS No.:	345651-04-9
Molecular Formula:	$C_{20}H_{22}N_4O$
Molecular Weight:	334.41
Target:	DNA/RNA Synthesis
Pathway:	Cell Cycle/DNA Damage
Storage:	4°C, protect from light * In solvent : -80°C, 6 months: -20°C, 1 month (protect from light)
	<i>,</i>

## SOLVENT & SOLUBILITY

	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	2.9903 mL	14.9517 mL	29.9034 mL		
		5 mM	0.5981 mL	2.9903 mL	5.9807 mL		
		10 mM	0.2990 mL	1.4952 mL	2.9903 mL		
	Please refer to the so	lubility information to select the ap	propriate solvent.				
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (6.22 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (6.22 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (6.22 mM); Clear solution						

BIOLOGICAL ACTIVITY				
Description	BVDV-IN-1 is a non-nucleoside inhibitor (NNI) of bovine viral diarrhea virus (BVDV), with an EC <sub>50</sub> of 1.8 μM. BVDV-IN-1 directly binds to a hydrophobic pocket of the BVDV RdRp. BVDV-IN-1 has antiviral activity against BVDV resistant to NNI thiosemicarbazone (TSC) <sup>[1]</sup> .			
In Vitro	BVDV-IN-1 (compound 1.9) binds to a hydrophobic pocket of the BVDV RNA-dependent RNA polymerase (RdRp) <sup>[1]</sup> . BVDV-IN-1 inhibits the replication of TSC-resistant BVDV variants in vitro <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			



Product Data Sheet

## REFERENCES

[1]. Gabriela A Fernández, et al. Design and Optimization of Quinazoline Derivatives: New Non-nucleoside Inhibitors of Bovine Viral Diarrhea Virus. Front Chem. 2020 Dec 10;8:590235.

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

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