

Product Name: GS-9620 Revision Date: 01/10/2021 Product Data Sheet

NH<sub>2</sub>

# **GS-9620**

Cat. No.:	A3444
CAS No.:	1228585-88-3
Formula:	C22H30N6O2
M.Wt:	410.51
Synonyms:	GS 9620;GS9620
Target:	Microbiology & Virology
Pathway:	HBV
Storage:	Store at -20°C
	810

## Solvent & Solubility

	insoluble in H2O; $\geq$	soluble in H2O; $\geq$ 20.55 mg/mL in DMSO; $\geq$ 9.9 mg/mL in EtOH with gentle warming and ultrasonic			
In Vitro	Preparing Stock Solutions	Mass Solvent Concentration	1mg	5mg	10mg
	Slock Solutions	1 mM	2.4360 mL	12.1800 mL	24.3599 mL
	<b>elo</b>	5 mM	0.4872 mL	2.4360 mL	4.8720 mL
	PELE	10 mM	0.2436 mL	1.2180 mL	2.4360 mL

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

### **Biological Activity**

Shortsummary

TLR-7 agonist

#### IC<sub>50</sub> & Target

	Cell Viability Assay	Part and	
	Cell Line:	Peripheral blood mononuclear cells (PBMCs) or plasmacytoid dendritic cells	
		(pDCs)	
In Vitro	Preparation method:	Soluble in DMSO. General tips for obtaining a higher concentration: Please	
		warm the tube at 37°C for 10 minutes and/or shake it in the ultrasonic bath for a	
		while. Stock solution can be stored below -20°C for several months.	
	Reacting conditions:	14 ~ 66 nM	
	Reacting conditions:		

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	Applications:	In human and cynomolgus monkey PBMCs and/or pDCs, GS-9620 induced interferon (IFN)-alpha and other cytokines, with a minimum effective concentration ranging from 14 to 66 nM in humans and with 5-fold less potency in monkeys.
	Animal experiment	-
	Animal models:	Cynomolgus monkeys
	Dosage form:	single doses of 0.1 ~ 2.0 mg, daily doses of 0.1 ~ 1.0 mg for 7 days or every
	A Carlo Carlos	other day doses of 0.05 ~ 1.5 mg for 28 days; p.o.
	Applications:	In cynomolgus monkeys, GS-9620 was well tolerated even at the highest oral
		doses (1.5 mg every other day for 28 days). GS-9620 increased IFN-alpha,
In Vivo		immunomodulatory cytokines, chemokines and peripheral blood cell IFN
		stimulated genes (ISGs) in a dose-dependent manner. In addition, there was
		no evidence of tachyphylaxis following every other day dosing, and oral
		administration resulted in limited systemic bioavailability but high oral
	810	absorption.
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may
	Contraction of the second	slightly differ with the theoretical value. This is caused by an experimental
		system error and it is normal.

### **Product Citations**

See more customer validations on www.apexbt.com.



[1]. Turnas P, Zheng X, Lu B, et al. 1129 Preclinical characterization of GS-9620, a potent and selective oral TLR7 agonist[J]. Journal of Hepatology, 2011, 54: S446-S447.

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#### NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

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