NADH-IN-1

Cat. No.:	HY-146036		
CAS No.:	1432445-15	-2	
Molecular Formula:	C ₁₉ H ₂₁ F ₃ N ₂ O	S	
Molecular Weight:	382.44		
Target:	Endogenou	s Metabo	lite
Pathway:	Metabolic E	nzyme/P	rotease
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month

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SOLVENT & SOLUBILITY

Sto		Solvent Mass Concentration	1 mg	5 mg	10 mg	
	Preparing Stock Solutions	1 mM	2.6148 mL	13.0739 mL	26.1479 mL	
		5 mM	0.5230 mL	2.6148 mL	5.2296 mL	
		10 mM	0.2615 mL	1.3074 mL	2.6148 mL	
	Please refer to the so	Please refer to the solubility information to select the appropriate solvent.				
n Vivo	1. Add each solvent	one by one: 10% DMSO >> 90% cor	n oil			

BIOLOGICAL ACTIV	ИТҮ
Description	NADH-IN-1 has NADH:ubiquinone oxidoreductase inhibitory activity with an IC ₅₀ value of 27 μM. NADH-IN-1 can effectively stimulate glucose uptake in vitro. NADH-IN-1 is readily metabolised by the liver. NADH-IN-1 can be used for researching diabetes ^[1] .
IC₅₀ & Target	IC ₅₀ : 27 μM (NADH:ubiquinone oxidoreductase) ^[1]
In Vivo	NADH-IN-1 (1 μM) exhibits a short half-life and fast intrinsic clearance indicating that it is readily metabolised by the liver in vivo; shows no adverse effects on primary rat hepatocytes, and does not inhibit the hERG channel ^[1] . NADH-IN-1 (10 mg/kg; IV or PO; single dosage) produces no observable toxic effects at 10 mg/kg by IV or PO; exhibits a short half-life and high plasma clearance; exhibits high mouse and human serum protein binding, as well as moderate bioavailability ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Product Data Sheet

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Animal Model:	Male C57BL/6 mice ^[1]
Dosage:	10 mg/kg
Administration:	IV or PO; single dosage (Pharmacokinetics Analysis)
Result:	Produced no observable toxic effects at 10 mg/kg by IV or PO; exhibited a short half-life of 0.45 h and high plasma clearance; exhibited high mouse and human serum protein binding, as well as moderate bioavailability (21.4%).

REFERENCES

[1]. Devine R, et al. Design, synthesis, and biological evaluation of aryl piperazines with potential as antidiabetic agents via the stimulation of glucose uptake and inhibition of NADH:ubiquinone oxidoreductase. Eur J Med Chem. 2020;202:112416.

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

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