NV-5138

Cat. No.:	HY-114384		
CAS No.:	2095886-80	-7	
Molecular Formula:	C ₇ H ₁₃ F ₂ NO ₂		
Molecular Weight:	181.18		
Target:	mTOR		
Pathway:	PI3K/Akt/m	TOR	
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month

®

MedChemExpress

SOLVENT & SOLUBILITY

	Mass Solvent Concentration	1 mg	5 mg	10 mg
Preparing Stock Solut	1 mM	5.5194 mL	27.5969 mL	55.1937 m
	5 mM	1.1039 mL	5.5194 mL	11.0387 ml
	10 mM	0.5519 mL	2.7597 mL	5.5194 mL

DIOLOGICALACTIV		
Description	NV-5138, a leucine analog, is for antidepressant studies ^[1]	s the first selective and orally active brain mTORC1 activator, binding to Sestrin2. NV-5138 is used][2].
IC ₅₀ & Target	mTORC1 ^[1] .	
In Vivo	NV-5138 is found to be esser following intravenous and o NV-5138 (160 mg/kg, po, sing phospho-p70S6K1, and pho NV-5138 (80 mg/kg, po, daily MCE has not independently	ntially 100% orally bioavailable with an elimination half-life in plasma of ~ 3 h determined oral dosing in rats ^[1] . gle dose) rapidly increases levels of phospho-mTOR as well as the downstream targets, sphor-4EB-P1, in synaptosomal preparations of PFC ^[2] . y for a total of 7 days) also produces antidepressant effects ^[2] . confirmed the accuracy of these methods. They are for reference only.
	Animal Model:	Male Sprague-Dawley rats weighing 250-260 g ^[2] .
	Dosage:	40, 80, 160 mg/kg.

Product Data Sheet

∠F

 $\bar{N}H_2$

F<

OH

Administration:	PO, single dose (160 mg/kg) or daily for a total of 7 days (40, 80 mg/kg).
Result:	Produced antidepressant effects.
Animal Model:	Male Sprague–Dawley (SD) rats weighed 250-400 g ^[1] .
Dosage:	1 mg/kg, 5 mg/kg (Pharmacokinetic Design).
Administration:	I.V at 1 mg/kg and PO at 5 mg/kg.

CUSTOMER VALIDATION

- McGill University. 2023 Jul 5.
- bioRxiv. 2023 Apr 24.

See more customer validations on <u>www.MedChemExpress.com</u>

REFERENCES

[1]. Sengupta S, et al. Discovery of NV-5138, the first selective Brain mTORC1 activator. Sci Rep. 2019 Mar 11;9(1):4107.

[2]. Kato T, et al. Sestrin modulator NV-5138 produces rapid antidepressant effects via direct mTORC1 activation. J Clin Invest. 2019 Apr 16;129(6):2542-2554.

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

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA