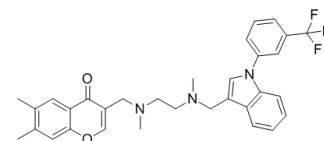


SPD304

Cat. No.:	HY-111255	
CAS No.:	869998-49-2	
Molecular Formula:	C ₃₂ H ₃₂ F ₃ N ₃ O ₂	
Molecular Weight:	547.61	
Target:	TNF Receptor	
Pathway:	Apoptosis	
Storage:	Powder	-20°C 3 years
	In solvent	-80°C 6 months
		-20°C 1 month



SOLVENT & SOLUBILITY

In Vitro

DMSO : 25 mg/mL (45.65 mM; Need ultrasonic)
 H₂O : 20 mg/mL (36.52 mM; Need ultrasonic)

Solvent	Mass	Concentration		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	1.8261 mL	9.1306 mL	18.2612 mL
	5 mM	0.3652 mL	1.8261 mL	3.6522 mL
	10 mM	0.1826 mL	0.9131 mL	1.8261 mL

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

BIOLOGICAL ACTIVITY

Description	SPD304 is a selective TNF- α inhibitor, which promotes dissociation of TNF trimers and therefore blocks the interaction of TNF and its receptor. SPD304 has an IC ₅₀ of 22 μ M for inhibiting in vitro TNF receptor 1 (TNFR1) binding to TNF- α ^{[1][2]} .
IC₅₀ & Target	IC ₅₀ : 22 μ M (TNF α) ^[1] .
In Vitro	SPD304 (2 μ M) significantly rescues the survivability of aHSCs, reduces the production of lipid hydroxides, and increased intracellular GSH. The co-treatment of GA (75 μ M) and SPD304 (2 μ M), down-regulate TRADD almost 2-fold (w/o inhibitor vs. w/ inhibitor) and p-RIP3 1.4-fold compared to GA alone, and promotes caspase 8 activation ^[4] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	SPD304 cannot be used in vivo due to its high toxicity ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Aging Cell. 2020 Oct;19(10):e13217.
- Cell Death Dis. 2020 Dec 11;11(12):1050.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Molly M. He, et al. Small-Molecule Inhibition of TNF- α . Science 11 Nov 2005.

[2]. Alexiou P, et al. Rationally designed less toxic SPD-304 analogs and preliminary evaluation of their TNF inhibitory effects. Arch Pharm (Weinheim). 2014 Nov;347(11):798-805.

[3]. Mouhsine H, et al. Identification of an in vivo orally active dual-binding protein-protein interaction inhibitor targeting TNF α through combined in silico/in vitro/in vivo screening. Sci Rep. 2017 Jun 13;7(1):3424.

[4]. Gallic acid induces necroptosis via TNF- α signaling pathway in activated hepatic stellate cells. Chang YJ, et al. PLoS One. 2015 Mar 27;10(3):e0120713.

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