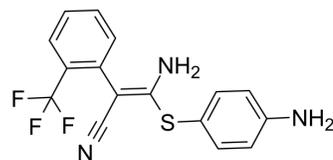


SL327

Cat. No.:	HY-15437		
CAS No.:	305350-87-2		
Molecular Formula:	C ₁₆ H ₁₂ F ₃ N ₃ S		
Molecular Weight:	335.35		
Target:	MEK		
Pathway:	MAPK/ERK Pathway		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro

DMSO : 68 mg/mL (202.77 mM; Need ultrasonic)
 Ethanol : 0.1 mg/mL (0.30 mM; Need ultrasonic and warming)

	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.9820 mL	14.9098 mL	29.8196 mL
	5 mM	0.5964 mL	2.9820 mL	5.9639 mL
	10 mM	0.2982 mL	1.4910 mL	2.9820 mL

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

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
 Solubility: ≥ 2.5 mg/mL (7.45 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
 Solubility: ≥ 2.5 mg/mL (7.45 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
 Solubility: ≥ 2.5 mg/mL (7.45 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

SL327 inhibits MEK1 and MEK2, with IC₅₀ values of 180 nM and 220 nM, respectively.

IC₅₀ & Target

MEK1 180 nM (IC ₅₀)	MEK2 220 nM (IC ₅₀)
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In Vitro

The specificity of SL327 for MEK is investigated. Kinase activity is assessed by measuring the incorporation of

[³²P]phosphate during phosphorylation of substrate peptides specific for each kinase. Although SL327 inhibits MEK with an IC₅₀ of 0.27 μM, 10 μM SL327 has no significant effect on PKA, CaMKII, or PKC^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

SL327, which crosses the blood-brain barrier, is administered intraperitoneally at several concentrations to animals prior to cue and contextual fear conditioning. Administration of SL327 completely blocks contextual fear conditioning and significantly attenuates cue learning when measure 24 hr after training. Animals treated with SL327 exhibit significant attenuation of water maze learning; they take significantly longer to find a hidden platform compared with vehicle-treated controls and also fail to use a selective search strategy during subsequent probe trials in which the platform is removed. Mice are injected with various concentrations of SL327 (10, 30, 50 mg/kg i.p.), and 1 hr later their hippocampi are removed and assayed for activated MAPK. SL327 attenuates phosphorylated MAPK levels in a dose-dependent manner. Administration of 10, 30, or 50 mg/kg SL327 significantly attenuates p42 phospho-MAPK levels (F=20.90, P<0.0001; 10 mg/kg SL327 vs. vehicle, P<0.05, and 30 and 50 mg/kg SL327 vs. vehicle, P<0.001). Injection with 30 or 50 mg/kg SL327 also significantly reduces p44 phospho-MAPK levels (F=5.627, P<0.005; 30 mg/kg vs. vehicle, P<0.05, and 50 mg/kg SL327 vs. vehicle, P<0.01)^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

PROTOCOL

Kinase Assay ^[2]

Protein kinase assays are performed. All kinase assays are started by adding enzyme to a mixture that included [^γ-³²P]ATP and substrate. This mixture is then incubated at 30°C or 37°C for 10 min. The reaction is stopped by spotting aliquots of the reaction mixture onto Whatman P-81 phosphocellulose filter paper. The papers are then washed in 150 mM H₃PO₄, dried, and subjected to scintillation counting. The catalytic subunit of PKA is assayed by measuring [³²P]phosphate incorporation into the substrate Kemptide (100 μM). The activity of CaMKII is determined by measuring phosphorylation of the synthetic peptide Autocamtide (100 μM) in the presence of 100 μM Calcium and 10 μg/mL Calmodulin. A synthetic peptide analog of a fragment of neurogranin, NG₍₂₈₋₄₃₎ (10 μM) is used as a specific substrate for the catalytic subunit of PKC. In all cases, substrate phosphorylation was linear with respect to time and enzyme concentration^[2].

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Animal Administration ^[2]

Mice^[2]

Adult male 129S3/SvImJ mice are used. In the 1x-pairing paradigm of cue and contextual fear conditioning, animals are placed in the fear conditioning apparatus for 3 min, then a 30-sec acoustic conditioned stimulus (CS; white noise, 80 dB) is delivered. During the last second of the CS, a 1-sec shock unconditioned stimulus (US; 0.5 mA) is applied to the grid floor. To assess contextual learning, the animals are returned to the training context 24 hr post-training, and freezing behavior is scored for 5 min. To assess cue learning, the animals are placed in a different context (novel odor, lighting, cage floor, and visual cues) following contextual testing. Baseline behavior is measured for 3 min in the novel context (Pre-CS), then the tone is presented for 3 min. Freezing behavior is assessed with a time sampling procedure whereby the animal was observed for ~1 sec every 5 sec. The experimenter is blind to drug treatment. Animals are injected with either vehicle (2 mL/kg, 100% DMSO) or SL327 (10, 30, and 50 mg/kg; at 2 mL/kg, dissolved in 100% DMSO) intraperitoneally 1 hr before training^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Brain Behav Immun. 19 August 2022.
- Eur J Pharmacol. 2023 Nov 7:961:176161.
- Front Mol Neurosci. 2018 Aug 21;11:287.
- Neuropharmacology. 2023 Sep 5, 109693.

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- Brain Res Bull. 2016 Jun;124:40-7.

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REFERENCES

- [1]. Cheng Y, et al. Current Development Status of MEK Inhibitors. Molecules. 2017 Sep 26;22(10). pii: E1551.
- [2]. Selcher JC, et al. A necessity for MAP kinase activation in mammalian spatial learning. Learn Mem. 1999 Sep-Oct;6(5):478-90.
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Caution: Product has not been fully validated for medical applications. For research use only.

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