Proteins

WS6

Cat. No.: HY-12461 CAS No.: 1421227-53-3 Molecular Formula: $C_{29}H_{31}F_3N_6O_3$ Molecular Weight: 568.59 Target: Others

Pathway: Others

Powder Storage: -20°C 3 years

2 years

In solvent -80°C 2 years

> -20°C 1 year

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Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: $\geq 100 \text{ mg/mL} (175.87 \text{ mM})$

* "≥" means soluble, but saturation unknown.

| Preparing Stock Solutions | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg |
|------------------------------|-------------------------------|-----------|-----------|------------|
| | 1 mM | 1.7587 mL | 8.7937 mL | 17.5874 mL |
| | 5 mM | 0.3517 mL | 1.7587 mL | 3.5175 mL |
| | 10 mM | 0.1759 mL | 0.8794 mL | 1.7587 mL |

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (4.40 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (4.40 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (4.40 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

WS6 is a novel small molecule that promotes β cell proliferation in rodent and human primary islets with EC50 of 0.28 uM(R7T1 cell viability).EC50 value: 0.28 uM [1]Target: β cell proliferation agonistin vitro: WS6 induced up to 4% of rat β cells to proliferate, with an EC50 of 0.4 μ M. In the same format, WS6 also induced 3% of human β cells to proliferate, with a similar potency to the rat β cells. WS6 induced R7T1 proliferation in dose response, with EC50 value of 0.28 μ M, Proliferation of R7T1 cells, which are cultured in suspension and grow as clusters, was apparent by visible inspection. in vivo: RIP-DTA mice were fed Dox in the drinking water until the onset of overt diabetes (blood glucose reading >300 mg/dL, typically 4-10 days), at

which point Dox treatment was discontinued and treatment with WS6 was initiated (5 mg/kg every other day via intraperitoneal injection). Pharmacokinetic studies with WS6 at 50 mg/kg revealed a CMAX of \boxtimes 5 μ M and T1/2 of \boxtimes 2 h. Treatment with WS6 caused a progressive reduction of blood glucose over time, starting around 2 weeks.

REFERENCES

[1]. Shen W, et al. Small-molecule inducer of β cell proliferation identified by high-throughput screening. J Am Chem Soc. 2013 Feb 6;135(5):1669-72.

[2]. Zhang H, Xiang L, Yang L, et al. WS6 Induces Adult Hippocampal Neurogenesis in Correlation to its Antidepressant Effect on the Alleviation of Depressive-like Behaviors of Rats. Neuroscience. 2021;473:119-129.

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

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