Proteins

Product Data Sheet



AKT-IN-6

Cat. No.: HY-19982 CAS No.: 1430056-54-4 Molecular Formula: $\mathsf{C}_{22}\mathsf{H}_{20}\mathsf{FN}_5\mathsf{O}$ Molecular Weight: 389.43 Target: Akt

Pathway: PI3K/Akt/mTOR

Storage: Powder -20°C 3 years

2 years

In solvent -80°C 6 months

> -20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 125 mg/mL (320.98 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.5679 mL	12.8393 mL	25.6786 mL
	5 mM	0.5136 mL	2.5679 mL	5.1357 mL
	10 mM	0.2568 mL	1.2839 mL	2.5679 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: ≥ 2.08 mg/mL (5.34 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (5.34 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (5.34 mM); Clear solution

BIOLOGICAL ACTIVITY

Description AKT-IN-6 (Example 13) is a potent Akt inhibitor. AKT-IN-6 inhibits Akt1, Akt2 and Akt3 with IC₅₀s < 500nM, respectively. (patent WO2013056015A1).

Akt1 Akt2 Akt3 IC₅₀ & Target

In Vitro Akt is a central node in cell signaling downstream of growth factors, cytokines, and other cellular stimuli. Aberrant loss or gain of Akt activation underlies the pathophysiological properties of a variety of complex diseases, including type-2 diabetes

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

REFERENCES

[1]. Taisheng Huang; et al. Isoindolinone and pyrrolopyridinone derivatives as akt inhibitors. WO2013056015A1.

[2]. Manning BD, et al. AKT/PKB signaling: navigating downstream. Cell. 2007;129(7):1261-1274.

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

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