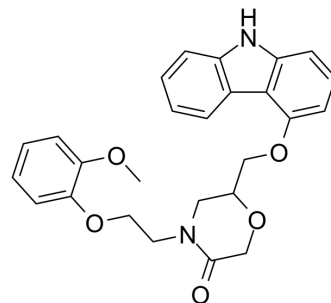


VK-II-36

Cat. No.:	HY-111014		
CAS No.:	955371-66-1		
Molecular Formula:	C ₂₆ H ₂₆ N ₂ O ₅		
Molecular Weight:	446.5		
Target:	Calcium Channel		
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro

DMSO : 100 mg/mL (223.96 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	2.2396 mL	11.1982 mL	22.3964 mL
5 mM	0.4479 mL	2.2396 mL	4.4793 mL
10 mM	0.2240 mL	1.1198 mL	2.2396 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 (5.60 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 2.5 mg/mL (5.60 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

VK-II-36 is a carvedilol analog that suppresses sarcoplasmic reticulum Ca²⁺ release but does not block the β-receptor. VK-II-36 inhibits triggered activities evoked by both early and delayed after depolarizations^[1].

REFERENCES

- [1]. Maruyama M, et al. Carvedilol analogue inhibits triggered activities evoked by both early and delayed afterdepolarizations. Heart Rhythm. 2013 Jan;10(1):101-7.

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

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