Phenindione

Cat. No.:HY-B0325CAS No.:83-12-5Molecular Formula: $C_{15}H_{10}O_2$ Molecular Weight:222.24Target:OthersPathway:Others

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 : ≥ 100 mg/mL (449.96 mM)

H₂O: < 0.1 mg/mL (insoluble)

* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	4.4996 mL	22.4982 mL	44.9964 mL
	5 mM	0.8999 mL	4.4996 mL	8.9993 mL
	10 mM	0.4500 mL	2.2498 mL	4.4996 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 (11.25 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (11.25 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Phenindione is an anticoagulant which functions as a Vitamin K antagonist. Target: OthersPhenindione (Rectadione) is an anticoagulant which functions as a Vitamin K antagonist. A lymphocyte transformation test showed proliferation of T-cells from the hypersensitive patient, but not from four controls on exposure to phenindione in vitro. Drug-specific T-cell clones were generated and characterized in terms of their phenotype, functionality, and mechanism of antigen presentation. Forty-three human leukocyte antigen class II restricted CD4+ $\alpha\beta$ T-cell clones were identified. T-cell activation resulted in the secretion of interferon- γ and interleukin-5 [1].

REFERENCES				
1]. Naisbitt, D.J., et al., Characterization of the T-cell response in a patient with phenindione hypersensitivity. Journal of Pharmacology and Experimental Therapeutics 313(3): p. 1058-1065.				
	Caution: Product has not been fully validated for medical applications. For research use only.			
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