

Product Data Sheet

(R)-Mephenytoin

Cat. No.:HY-126043CAS No.:71140-51-7Molecular Formula: $C_{12}H_{14}N_2O_2$ Molecular Weight:218.25

Target: Cytochrome P450

Pathway: Metabolic Enzyme/Protease

Storage: Powder -20°C 3 years
In solvent -80°C 6 months

-20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 220 mg/mL (1008.02 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	4.5819 mL	22.9095 mL	45.8190 mL
	5 mM	0.9164 mL	4.5819 mL	9.1638 mL
	10 mM	0.4582 mL	2.2910 mL	4.5819 mL

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
 - Solubility: ≥ 5.5 mg/mL (25.20 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 5.5 mg/mL (25.20 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	$(R)-Mephenytoin \ ((-)-Mephenytoin), the \ R-enantiomer \ of \ Mephenytoin. \ Mephenytoin \ is \ an \ Anticonvulsant \ agent \ [1][2].$
In Vitro	(R)-Mephenytoin can be N-demethylated by the cytochrome P450 (CYP) isoform CYP2C9 to form the metabolite 5-phenyl-5-ethylhydantoin (nirvanol) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Meier UT, et, al. Assay of mephenytoin metabolism in human liver microsomes by high-performance liquid chromatography. Anal Biochem. 1985 Dec;151(2):286-91.

2]. Relling MV, et, al. Tolbutamide	e and mephenytoin hydroxylation by human cytochrome P	450s in the CYP2C subfamily. J Pharmacol Exp Ther. 1990 Jan;252(1):442-7.
(Caution: Product has not been fully validated for me	edical applications. For research use only.
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