PD 117519

Cat. No.: HY-100032 CAS No.: 96392-15-3 Molecular Formula: $C_{19}H_{21}N_{5}O_{4}$

Molecular Weight: 383.4

Target: Adenosine Receptor Pathway: GPCR/G Protein

Powder -20°C Storage:

3 years 2 years

In solvent -80°C 2 years

> -20°C 1 year

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: ≥ 100 mg/mL (260.82 mM)

* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.6082 mL	13.0412 mL	26.0824 mL
	5 mM	0.5216 mL	2.6082 mL	5.2165 mL
	10 mM	0.2608 mL	1.3041 mL	2.6082 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: ≥ 2.5 mg/mL (6.52 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.52 mM); Clear solution

BIOLOGICAL ACTIVITY

PD 117519 (CI947) is an A_{2A} adenosine agonist which has shown oral antihypertensive activity in pharmacological animal Description $models^{[1][2]}$.

IC₅₀ & Target A_{2A} adenosine^[1]

In Vivo

PD 117519 (2-10 mg/kg; oral administration; 16-24 hours; male beagle dogs) treatment produces significant hemodynamic changes at T_{max} (4 hours) follows by acute coronary vascular injury that is evident at 16 hours postdosing. Treatment with 2 or 10 mg/kg of PD 117519 produces significant increases in mean heart rate and decreases in mean indirectsystolic blood pressure at time of highest drug exposure, 4 hours postdosing^[3].

Animal Model:	24 male beagle dogs (8-12 months old) ^[3]	
Dosage:	2 mg/kg or 10 mg/kg	
Administration:	Oral administration; 16 hours and 24 hours	
Result:	Increased in mean heart rate and decreased in mean indirectsystolic blood pressure time of highest drug exposure. Induced acute coronary arteriopathy. The endotheliu also appears injured.	

REFERENCES

- [1]. Reynolds DL, et al. Liquid chromatographic analysis of the adenosine agonist PD 117519 in dog plasma. J Pharm Biomed Anal. 1991;9(4):345-9.
- [2]. Tobin GA, et al. The role of eNOS phosphorylation in causing drug-induced vascular injury. Toxicol Pathol. 2014 Jun;42(4):709-24.
- [3]. Enerson BE, et al. Acute drug-induced vascular injury in beagle dogs: pathology and correlating genomic expression. Toxicol Pathol. 2006;34(1):27-32.

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

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