Proteins



Product Data Sheet

LY223982

Cat. No.: HY-112737 CAS No.: 117423-74-2 Molecular Formula: $C_{30}H_{30}O_{7}$ Molecular Weight: 502.56

Target: Leukotriene Receptor Pathway: GPCR/G Protein

Storage: Powder -20°C 3 years

> In solvent -80°C 6 months

> > -20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (198.98 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.9898 mL	9.9491 mL	19.8981 mL
	5 mM	0.3980 mL	1.9898 mL	3.9796 mL
	10 mM	0.1990 mL	0.9949 mL	1.9898 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 (4.97 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (4.97 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	LY223982 is a potent and specific inhibitor of leukotriene B4 receptor, with an IC ₅₀ of 13.2 nM against [³ H]LTB4 binding to LTB4 receptor.
IC ₅₀ & Target	LTB ₄ 13.2 nM (IC ₅₀)
In Vitro	LY223982 is a potent and specific inhibitor of leukotriene B4 (LTB4) receptor, with an IC $_{50}$ of 13.2 nM against [3H]LTB4 binding to LTB4 receptor. LY223982 is also a potent antagonist of the aggregation of human neutrophils by LTB4 (IC $_{50}$, 100 nM) $^{[1]}$. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

LY223982 inhibits transient leukopenia induced in rabbits with LTB4 (ED $_{50}$, 3 mg/kg) but not with FMLP, and shows no agonist activity in any of the test systems^[1].

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

REFERENCES

[1]. Jackson WT, et al. Specific inhibition of leukotriene B4-induced neutrophil activation by LY223982. J Pharmacol Exp Ther. 1992 Dec;263(3):1009-14.

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

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