BAY 73-1449

Cat. No.:	HY-118941		
CAS No.:	693790-96-4	ļ	
Molecular Formula:	$C_{26}H_{23}N_{3}O_{3}$		
Molecular Weight:	425.48		
Target:	Prostaglanc	lin Recept	tor
Pathway:	GPCR/G Pro	tein	
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month

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SOLVENT & SOLUBILITY

		Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solution	Preparing Stock Solutions	1 mM	2.3503 mL	11.7514 mL	23.5029 mL
		5 mM	0.4701 mL	2.3503 mL	4.7006 mL
		10 mM	0.2350 mL	1.1751 mL	2.3503 mL
	Please refer to the so	lubility information to select the app	propriate solvent.		
n Vivo		one by one: 10% DMSO >> 90% (20 g/mL (4.89 mM); Suspended solutior			
		one by one: 10% DMSO >> 90% cor ng/mL (4.89 mM); Clear solution	n oil		

BIOLOGICAL ACTIV	
Description	BAY 73-1449 is a selective antagonist of prostacyclin receptor (IP), with high potency (IC ₅₀ of less than 0.1 nM) in cAMP assays in Human HEL cells and rat DRG. BAY 73-1449 can be used in the research of lowering blood pressure ^[1] .
IC₅₀ & Target	IP Receptor <0.1 nM (IC ₅₀)
In Vivo	BAY 73-1449 (0.1-1 mg/kg; i.v.) does not significantly reduce mesenteric inflow, but significantly reduces splenic shunt vessel outflow in rats ^[1] . BAY 73-1449 (1-5 mg/kg, s.c. once daily for 7 d) has no effects on the degree of porto-systemic shunting in rats ^[1] . BAY 73-1449 (1 mg/kg, s.c. once daily for 7 d), has no effects on portal pressures in rats ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Product Data Sheet

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Animal Model:	Male Wistar rats (250-350 g) are ligated portal vein $^{[1]}$
Dosage:	0.1, 1 mg/kg
Administration:	A single i.v.
Result:	Significantly reduced shunt flow without affecting mesenteric flow.

REFERENCES

[1]. Bexis S, et, al. Vascular actions of the prostacyclin receptor antagonist BAY 73-1449 in the portal hypertensive rat. Eur J Pharmacol. 2008 Aug 20;590(1-3):322-6.

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

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