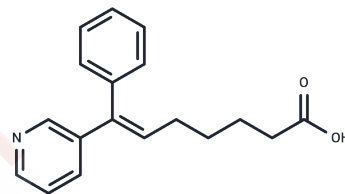


Isbogrel

Chemical Properties

CAS No. :	89667-40-3
Formula:	C ₁₈ H ₁₉ NO ₂
Molecular Weight:	281.35
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year Actual storage temperature shall be subject to the COA.



Biological Description

Description	Isbogrel (CV4151) is a small molecule thromboxane A ₂ synthase (TXA ₂ synthase) inhibitor that can be used to study cardiac arrhythmias, transient ischemic attacks and thrombosis.
Targets(IC ₅₀)	Prostaglandin Receptor
In vivo	Isbogrel significantly inhibited photochemically induced MCA thrombosis by oral (1 and 10 mg/kg) and intravenous (1 mg/kg) administration.[2] Isbogrel (10 mg/kg, p.o.), furthermore, significantly reduced the infarct size and inhibited the increase in lactate content. Isbogrel might be a useful drug for the treatment of cerebral thrombosis and for the prevention of cerebral infarction.[2]

Solubility Information

Solubility	DMSO: 55 mg/mL (195.49 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	3.5543 mL	17.7715 mL	35.5429 mL
5 mM	0.7109 mL	3.5543 mL	7.1086 mL
10 mM	0.3554 mL	1.7771 mL	3.5543 mL
50 mM	0.0711 mL	0.3554 mL	0.7109 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Terashita Z, et al. Inhibition of arachidonic acid induced-aggregation of rabbit platelets with CV-4151 (isbogrel), a selective thromboxane A2 (TXA2) synthase inhibitor: modulation of the antiplatelet action and prostanoid metabolism by rat aortic rings. *J Lipid Mediat Cell Signal*. 1996 Jan;13(1):1-8.

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Asai F, et al. In vitro antiplatelet profiles of the new thromboxane synthetase inhibitor sodium 2-(1-imidazolylmethyl)-4,5-dihydrobenzo[b]thiophene-6-carboxylate. *Arzneimittelforschung*. 1991 May;41(5):506-10.

Terashita Z, et al. Effects of thromboxane A2 synthase inhibitors (CV-4151 and ozagrel), aspirin, and ticlopidine on the thrombosis caused by endothelial cell injury. *Thromb Res*. 1995 Mar 1;77(5):411-21.

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