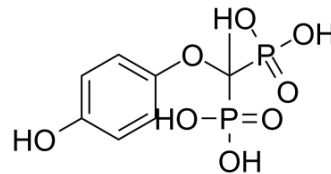


L-690330

Cat. No.:	HY-101075		
CAS No.:	142523-38-4		
Molecular Formula:	C ₈ H ₁₂ O ₈ P ₂		
Molecular Weight:	298.12		
Target:	Phosphatase		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro
 H₂O : 33.33 mg/mL (111.80 mM; Need ultrasonic)
 DMSO : 25 mg/mL (83.86 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
	1 mM	3.3544 mL	16.7718 mL	33.5435 mL
	5 mM	0.6709 mL	3.3544 mL	6.7087 mL
	10 mM	0.3354 mL	1.6772 mL	3.3544 mL

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

In Vivo

- Add each solvent one by one: **10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline**
 Solubility: ≥ 2.5 mg/mL (8.39 mM); Clear solution
- Add each solvent one by one: **10% DMSO >> 90% (20% SBE-β-CD in saline)**
 Solubility: ≥ 2.5 mg/mL (8.39 mM); Clear solution
- Add each solvent one by one: **10% DMSO >> 90% corn oil**
 Solubility: ≥ 2.5 mg/mL (8.39 mM); Clear solution

BIOLOGICAL ACTIVITY

Description
 L-690330 is a competitive inhibitor of **inositol monophosphatase (IMPase)** with K_is of 0.27 and 0.19 μM for recombinant human and bovine IMPase, 0.30 and 0.42 μM for human and bovine frontal cortex IMPase, respectively. L-690330 exhibits 10-fold more sensitive than mouse and rat IMPase^[1].

IC₅₀ & Target
 Ki: 0.27 μM (Recombinant human IMPase), 0.19 μM (Recombinant bovine IMPase), 0.30 μM (Human frontal cortex)

IMPase), 0.42 μ M (Bovine frontal cortex IMPase)^[1]

REFERENCES

[1]. Atack JR, et al. In vitro and in vivo inhibition of inositol monophosphatase by the bisphosphonate L-690,330. J Neurochem. 1993 Feb;60(2):652-8.

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

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