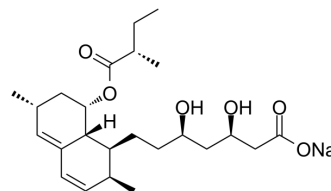


## Lovastatin hydroxy acid sodium

<b>Cat. No.:</b>	HY-123672
<b>CAS No.:</b>	75225-50-2
<b>Molecular Formula:</b>	C <sub>24</sub> H <sub>37</sub> NaO <sub>6</sub>
<b>Molecular Weight:</b>	444.54
<b>Target:</b>	HMG-CoA Reductase (HMGCR)
<b>Pathway:</b>	Metabolic Enzyme/Protease
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 12.5 mg/mL (28.12 mM; ultrasonic and warming and heat to 60°C)

Solvent	Mass	Concentration		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.2495 mL	11.2476 mL	22.4952 mL
	5 mM	0.4499 mL	2.2495 mL	4.4990 mL
	10 mM	0.2250 mL	1.1248 mL	2.2495 mL

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

### BIOLOGICAL ACTIVITY

#### Description

Lovastatin hydroxy acid sodium (Mevinolinic acid sodium) is a highly potent inhibitor of HMG-CoA reductase with a K<sub>i</sub> of 0.6 nM<sup>[1]</sup>.

#### IC<sub>50</sub> & Target

Ki: 0.6 nM (HMG-CoA reductase)<sup>[1]</sup>

#### In Vitro

Mevinolin in the hydroxy-acid form, mevinolinic acid, is a potent competitive inhibitor of 3-hydroxy-3-methylglutaryl-coenzyme A reductase [mevalonate: NADP<sup>+</sup> oxidoreductase (CoA-acylating)]<sup>[1]</sup>.  
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. A W Alberts, et al. Mevinolin: A Highly Potent Competitive Inhibitor of Hydroxymethylglutaryl-Coenzyme A Reductase and a Cholesterol-Lowering Agent. Proc Natl Acad Sci U S A. 1980 Jul;77(7):3957-61.

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

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