



# SZABO SCANDIC

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## Produktinformation



Forschungsprodukte & Biochemikalien



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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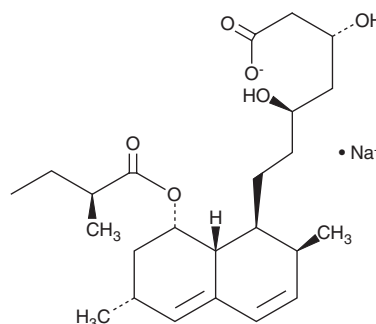
# PRODUCT INFORMATION



## Lovastatin Hydroxy Acid (sodium salt)

Item No. 10010339

**CAS Registry No.:** 75225-50-2  
**Formal Name:** 2-methyl-1S,2,3R,7S,8S,8aR-hexahydro-3,7-dimethyl-8-[[2-[(2R,4R)-tetrahydro-4-hydroxy-6-oxo-2H-pyran-2-yl]ethyl]-1-naphthalenyl ester, butanoic acid, monosodium salt  
**MF:**  $C_{24}H_{37}O_6 \cdot Na$   
**FW:** 444.5  
**Purity:**  $\geq 95\%$   
**UV/Vis.:**  $\lambda_{max}$ : 231, 238, 246 nm  
**Supplied as:** A crystalline solid  
**Storage:**  $-20^{\circ}C$   
**Stability:**  $\geq 2$  years



Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

### Laboratory Procedures

Lovastatin hydroxy acid (sodium salt) is supplied as a crystalline solid. A stock solution may be made by dissolving the lovastatin hydroxy acid (sodium salt) in an organic solvent purged with an inert gas. Lovastatin hydroxy acid (sodium salt) is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide. The solubility of lovastatin hydroxy acid (sodium salt) in these solvents is approximately 10 mg/ml.

If aqueous stock solutions are required for biological experiments, they can best be prepared by diluting the organic solvent into aqueous buffers or isotonic saline. Ensure that the residual amount of organic solvent is insignificant, since organic solvents may have physiological effects at low concentrations. We do not recommend storing the aqueous solution for more than one day.

### Description

Lovastatin hydroxy acid is a potent and competitive inhibitor of HMG-CoA reductase ( $K_i = 0.6$  nM).<sup>1</sup> It is the more potent, open ring, hydroxy acid form of its prodrug, lovastatin (Item No. 10010338).<sup>2</sup>

### References

1. Alberts, A.W., Chen, J., Kuron, G., *et al.* Mevinolin: A highly potent competitive inhibitor of hydroxymethylglutaryl-coenzyme A reductase and a cholesterol-lowering agent. *Proc. Natl. Acad. Sci. USA* **77**(7), 3957-3961 (1980).
2. Duggan, D.E., Chen, I.W., Bayne, W.F., *et al.* The physiological disposition of lovastatin. *Drug Metab. Dispos.* **17**(2), 166-173 (1989).

#### WARNING

THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

#### SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the [complete](#) Safety Data Sheet, which has been sent via email to your institution.

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