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## **Product Information**

Product ID R5974

CAS No. 147098-20-2

Chemical Name (S-((R\*,S\*-(E)))-7-(4-(4-fluorophenyl)-6-(1-methylethyl)-2-

(methyl(methylsulfonyl) amino)-5-pyrimidinyl)-3,5-dihydroxy-

6-heptenoic acid, calcium salt (2:1)

Synonym

Formula 2C<sub>22</sub>H<sub>26</sub>FN<sub>2</sub>O<sub>6</sub>S) Ca<sup>2+</sup>

Formula Wt. 1001.14

Melting Point 156-160°C dec°mp

Purity ≥98%

Solubility Slightly soluble in water, marginally soluble in ethanol, soluble

in DMSO(100mg/mL), DMF

## **Pricing and Availability**

Bulk quanitites available upon request

Product ID	Size	List Price
R5974	25 mg	\$169.10
R5974	100 mg	\$563.60
R5974	250 mg	\$1070.70

Store Temp Ambient Ship Temp Ambient

Description Rosuvastatin is an inhibitor of HMG-CoA reductase that decreases serum LDL and VLDL and is used to treat atherosclerosis in clinical settings. In addition to anti-atherosclerotic benefit, rosuvastatin also exhibits anti-inflammatory, cardioprotective, and anti-platelet activities. In animal models, rosuvastatin decreases levels of plasma lipids and angiotensin II (ATII) by inhibiting expression of the angiotensin II type 1 (AT1) receptor and p-ERK1/2 and upregulating scavenger receptor B1. Separately, rosuvastatin downregulates ERK, p38MAPK, matrix metalloproteinases 2 and 9 (MMP2/9), and PAI expression, inhibiting migration and proliferation of vascular smooth muscle cells. This compound also downregulates expression of the nucleotidebinding oligomerization domain-like receptor protein 3 (NLRP3) inflammasome, inhibiting cardiac remodeling and dysfunction in animal models of diabetic cardiomyopathy. Additionally, rosuvastatin inhibits activation of NADPH oxidase and PKC, decreasing platelet recruitment and activation in vitro and ex vivo.

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Caution: This product is intended for laboratory and research use only. It is not for human or drug use.