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Certificate of Analysis

 $\begin{array}{lll} \text{Product Name} & \text{Lovastatin, EP grade} \\ \text{Glentham Code} & \text{GP2911} \\ \text{CAS Number} & 75330\text{-}75\text{-}5 \\ \text{MDL Number} & \text{MFCD00072164} \\ \text{Batch Number} & 030\text{FYG} \\ \\ \text{Molecular Weight} & 404.56 \\ \text{Molecular Formula} & \text{C}_{24}\text{H}_{36}\text{O}_{5} \\ \text{Storage Temp.} & +4^{\circ}\text{C} \\ \end{array}$

Property	Specification	Batch 030FYG
Physical Description	White or almost white crystalline powder	Conforms
Identification (IR)	To conform to standard	Conforms
Specific Optical Rotation	+325 - +340 ° (c=0.5, CH3CN)	330.4 °
Impurity E	≤ 0.5%	0.21%
Impurity A	≤ 0.3%	0.05%
Impurity B	≤ 0.3%	0.12%
Impurity C	≤ 0.3%	Conforms
Impurity D	≤ 0.3%	0.13%
Impurity F	≤ 0.15%	0.11%
Any Unspecified Impurity	≤ 0.10%	0.06%
Total Impurities	≤ 1.0%	0.47%
Loss on Drying	≤ 0.5%	0.19%
Sulphated Ash	≤ 0.2%	0.08%
Assay	97.0 - 102.0 % (dried substance)	99.46%
Pharmacopoeia Specification(s)	EP	Conforms to EP

Specification Version v1.0

Manufacture Date 2018-10-01

Re-Test Date 2023-11-05

Glentham Life Sciences confirm that the above referenced product conformed to the information displayed in this document on the quality release date. Please check www.glentham.com or contact us using the details above for the current version of this document.

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