Lovastatin Tablets

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Expert Committee	Small Molecules 2

In accordance with the Rules and Procedures of the Council of Experts, the Small Molecules 2 Expert Committee has revised the Lovastatin Tablets monograph. The purpose of this revision is to widen the *Acceptance criteria* of lovastatin acid from NMT 0.85% to NMT 1.5% in the test for *Organic Impurities* to accommodate FDA-approved drug products.

The Lovastatin Tablets Revision Bulletin supersedes the currently official monograph.

Should you have any questions, please contact Donald Min, Staff Scientist (301-230-7457 or ddm@usp.org).