Bisoprolol Fumarate Tablets

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In accordance with the Rules and Procedures of the Council of Experts, the Small Molecules 2 Expert Committee has revised the Bisoprolol Fumarate Tablets monograph. The purpose of this revision is to add *Dissolution Test 3* to accommodate FDA-approved drug products with different dissolution conditions and/or tolerances than the existing dissolution test(s). Existing references to reagents have been updated for consistency with the reagent entry.

- *Dissolution Test 3* was validated using the Waters Symmetry C8 brand of column with L7 packing. The typical retention time for bisoprolol is about 2.7 min.

The Bisoprolol Fumarate Tablets Revision Bulletin supersedes the currently official monograph.

Should you have any questions, please contact Jasmine Lawrence, Scientist III (+1-301-230-6363 or jasmine.lawrence@usp.org).