Levetiracetam Tablets

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Expert Committee	Small Molecules 4

In accordance with the Rules and Procedures of the Council of Experts, the Small Molecules 4 Expert Committee has revised the Levetiracetam Tablets monograph. The purpose of this revision is to add *Dissolution Test 5* to accommodate FDA-approved drug products with different dissolution conditions and/or tolerances than the existing dissolution test(s). Additionally, minor editorial changes have been made to update the monograph to current USP style.

- *Dissolution Test 5* was validated using the Symmetry C18 brand of column with L1 packing. The typical retention time for levetiracetam is about 3 min.

The Levetiracetam Tablets Revision Bulletin supersedes the currently official monograph.

Should you have any questions, please contact Robyn Fales, Senior Scientist I (240-221-2047 or rnp@usp.org).