Ampicillin Capsules

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In accordance with the Rules and Procedures of the Council of Experts, the Small Molecules 1 Expert Committee has revised the Ampicillin Capsules monograph. The purpose of this revision is to add Dissolution Test 2 to accommodate FDA-approved drug products with different dissolution conditions and/or tolerances than the existing dissolution test. Labeling information has been incorporated to support the inclusion of Dissolution Test 2.

- Dissolution Test 2 was validated using the Triart C18 brand of column with L1 packing. The typical retention time for ampicillin is about 6 min.

The Ampicillin Capsules Revision Bulletin supersedes the currently official monograph.

Should you have any questions, please contact Yanyin Yang, Senior Scientist II, (301-692-3623 or yanyin.yang@usp.org).