Tranexamic Acid 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 Tranexamic Acid 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).

- *Dissolution Test 3* was validated using the Nucleosil 100 C18 brand of column with L1 packing (4.6-mm × 15-cm, 5 µm). The typical retention time for tranexamic acid is about 5.5 min.

The Tranexamic Acid Tablets Revision Bulletin supersedes the currently official monograph.

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