Fexofenadine Hydrochloride Tablets

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Expert Committee	Small Molecules 5

In accordance with the Rules and Procedures of the Council of Experts, the Small Molecules 5 Expert Committee has revised the Fexofenadine Hydrochloride Tablets monograph. The purpose of this revision is to widen the acceptance criteria for *Fexofenadine related compound A* from NMT 0.4% to NMT 1.0% and *Total impurities* from NMT 0.5% to NMT 1.3% in the test for *Organic Impurities*, to accommodate FDA-approved drug products with different limits.

Existing references to reagents and reagent names have been updated for consistency with official reagent entry names.

The Fexofenadine Hydrochloride Tablets 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).