

Guanfacine 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 Guanfacine Tablets 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(s). *Labeling* information has been incorporated to support the inclusion of *Dissolution Test 2*.

• Dissolution Test 2 was validated using the Microbondapack C18 brand of column with L1 packing. The typical retention time for guanfacine is about 3 min.

Additional editorial changes were made to update the monograph to current *USP* style.

The Guanfacine 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).