Hydroxychloroquine Sulfate Tablets

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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 Hydroxychloroquine Sulfate 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). Existing references to reagents have been updated for consistency with the reagent entry. In addition, an option for deaeration of media in Dissolution Test 1 was added to accommodate FDA-approved drug products utilizing this dissolution test. The format of this test was updated to current USP technical style, including the addition of the necessary equations.

The Hydroxychloroquine Sulfate Tablets Revision Bulletin supersedes the currently official monograph.

Should you have any questions, please contact Brice Wagner, Scientist III (301-998-6832 or brice.wagner@usp.org).