Buspirone Hydrochloride Tablets

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In accordance with the Rules and Procedures of the Council of Experts, the Small Molecules 4 Expert Committee has revised the Buspirone Hydrochloride Tablets monograph. The purpose for the revision is to modify the existing Dissolution Test to include the use of a suitable sinker to accommodate FDA-approved drug products with different dissolution conditions, and to add an alternate wavelength for the UV-based analytical procedure.

The Buspirone Hydrochloride Tablets Revision Bulletin supersedes the currently official monograph.

Should you have any questions, please contact Devendra Pratap Singh, Associate Scientific Liaison (404-448-8975 or dxp@usp.org).