Alfuzosin Hydrochloride Extended Release Tablets

Type of Posting	Revision Bulletin
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Official Date	01–Jun–2016
Expert Committee	Chemical Medicines Monographs 5
Reason for Revision	Compliance

In accordance with the Rules and Procedures of the 2015-2020 Council of Experts, the Chemical Medicines Monographs 5 Expert Committee has revised the Alfuzosin Hydrochloride Extended-Release Tablets monograph. The purpose for the revision is to add a dissolution test for a generic product approved by the FDA. The liquid chromatographic procedure in *Dissolution Test 7* is based on analyses performed with an Inertsil C8-3 brand of L7 column. The typical retention time for alfuzosin is about 2.6 min.

The Alfuzosin Hydrochloride Extended Release Tablets Revision Bulletin supersedes the current official monograph. The Revision Bulletin will be incorporated into *USP 40–NF 35*.

Should you have any questions, please contact Mary P. Koleck, Ph.D., Scientific Liaison (301-230-7420 or mpk@usp.org).