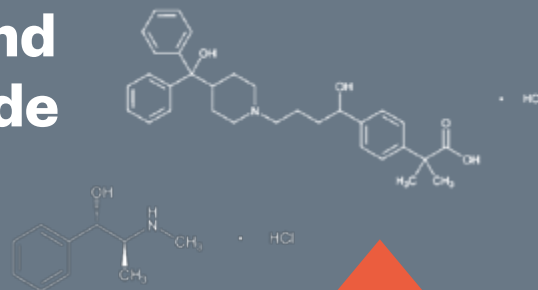


Fexofenadine Hydrochloride and Pseudoephedrine Hydrochloride

Quality Solutions

Category: Antihistamine



USP can support your development and manufacturing activities on Fexofenadine Hydrochloride and Pseudoephedrine Hydrochloride-based medicines with these existing and upcoming standards.

Official and Proposed Documentary Standards and Associated Physical Materials

MONOGRAPH 1

FEXOFENADINE HYDROCHLORIDE AND PSEUDO-EPHEDRINE HYDROCHLORIDE EXTENDED-RELEASE TABLETS | NEW

Official as of 01-Nov-2023

USP Benzoic Acid RS	USP-1055002
USP Fexofenadine Hydrochloride RS	USP-1270377
USP Pseudoephedrine Hydrochloride RS	USP-1581005
USP Fexofenadine Related Compound A RS	USP-1270388

PHARMACEUTICAL ANALYTICAL IMPURITY(IES)*

USP Fexofenadine Olefin PAI	USP-1A00731
USP Fexofenadine Chloro Analog PAI New	USP-1A00980
USP Fexofenadine N-oxide PAI	USP-1A01920

* Documents in PF Online are not official and not suitable to demonstrate compliance. They may never become official.

* Pharmaceutical Analytical Impurities (PAI products) are released using a process developed by USP's subject matter experts. The release process is based on internal policies, standard operating procedures, and requirements as defined by USP's Quality Management System. USP is an ISO 9001:2015 certified facility. PAI products are different from official USP Reference Standards. PAI products are not required for compendial compliance.

01-Dec-2023. For publications related to **Fexofenadine Hydrochloride and Pseudoephedrine Hydrochloride** in PF online after this date, please visit the integrated [USP-NF/PF](#) online.

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