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USP 37 DELIVERABLE VOLUME (698): Meets the requirements for Oral Suspension packaged in multiple-unit containers LIMIT OF 4-AMINOPHENOL A N/øthnnnl fnrmir and wafer (7 S' 2 '42 S h Official Monographs / Acetaminophen 1569 sonicate for 5 min, and dilute with Mobile phase to vol-

20140418141324-DM-DM 20140418141309 M99873 - USP-NF

31 be carefully checked to ascertain that they will function properly See USP general 32 chapters Aerosols, Nasal Sprays, Metered-Dose Inhalers, and Dry Powder Inhalers 33 601 , Deliverable Volume 698 , Density of Solids 699 , Osmolality and Osmolarity 34 785 , pH 791 , Pharmaceutical Compounding—Nonsterile Preparations 795 ,

<698> DELIVERABLE VOLUME - 2012-10-01

<698> DELIVERABLE VOLUME - 2012-10-01 Monograph Title <698> DELIVERABLE VOLUME Errata Identifier

0f3f3944-7368-941c-4e43-45627374a185 Figure 1, right branch, left box: Change Volume of 1 more containers is less than 95% LV to: Volume of 1 or more containers is less than 95% LV Section ACCEPTANCE CRITERIA/For Multiple-Unit Containers Page Number

Oxford Textbook Sports Medicine Medical

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theunitedstate pharmacopeia 2 -volume

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<1151> PHARMACEUTICAL DOSAGE FORMS

USP 35 General Information / □1151□ Pharmaceutical Dosage Forms765 maceutical preparations, which are given elsewhere in thistent uniformity does not rely on the assumption of blend Pharmacopeia uniformity and can be applied in all cases

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General Chapters Overview Anthony J DeStefano, PhD Vice President, General Chapters USP 2010-2015 Council of Experts Expert Committee Orientation

Sample Sizes in Uniformity Measurements - The Role of USP

Sample Sizes in Uniformity Measurements - The Role of USP Anthony J DeStefano, PhD Walter W Hauck, PhD Vice President, General Chapters Sr Scientific Fellow

» Ibuprofen contains not less than 97.0 percent and not ...

ing 20 mg per mL of USP Ibuprofen RS in chloroform to a thin-the requirements layer chromatographic plate (see Chromatography □621□) coatedDeliverable volume □698□— with a 025-mm layer of chromatographic silica gel and previ-FOR ORAL SUSPENSION PACKAGED IN MULTIPLE-UNIT CONTAINERS: ously activated by heating at 105° for 30 minutes

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1USP 38 THE UNITED STATES PHARMACOPEIA 1NF 33 THE NATIONAL FORMULARY Volume 3/a By authority of the United States Pharmacopeia1 Convention Prepared by the Council of Experts and its Expert Committees Official from May 1, 2015 The designation on the cover of this publication, "USP NF 2015," is for ease of identification only

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