ERRATA

INTERIM REVISION ANNOUNCEMENT

Following is a list of errata and corrections to *USP–NF*. The page number indicates where the item is found and in which official or pending official publication of *USP–NF*. If necessary, this list will be updated with every issue of *PF*. This information will also be available as a cumulative table in future *Supplements* and will appear in its corrected form in a future annual edition of *USP–NF*. An erratum consists of content erroneously published that does not accurately reflect the intended official or effective requirements as approved by the Council of Experts. USP staff is available to respond to questions regarding the accuracy of a particular requirement by calling 1-800-822-USPC.

USP32–NF27 Page	Title	Section	Description
182	〈561〉 Articles of Botanical Origin	Methods of Analysis	Line 5 under Alcohol-Soluble Extractives, Method 2: Change "and then allowing to stand for 18 hours." to:
			and then allowing to stand.
261	(699) Density of Solids	Gas Pycnometry for the Measurement of Density	Change: $V_s = V_c + \frac{V_r}{1 - \left[\frac{P_i - P_r}{P_f - P_r}\right]} $ (1)
			to:
			$V_{S} = V_{C} - \frac{V_{r}}{\left[\frac{P_{i} - P_{r}}{P_{f} - P_{r}}\right] - 1}$
1174	Benzalkonium Chloride Solution	Limit of foreign amines	Change ", meets the requirements of the test for Foreign amines under Benzalkonium Chloride." to: To 5 mL of this solution add 3 mL of 1 N sodium hydroxide: no precipitate is formed. Heat to boiling: the odor of amines is not perceptible.
1621	Bacitracin Ointment	Assay	Line 9: Change "and quantitatively dilute with <i>Test Dilution</i> " to: and quantitatively dilute with <i>Buffer No. 1</i> to obtain a <i>Test Dilution</i>
online	Cefazolin	Assay	Lines 1–2 under <i>Mobile phase</i> , change " <i>Mobile phase</i> —Prepare a suitable mixture of <i>pH 7.0 Buffer</i> and acetonitrile (9:1)." to: <i>Mobile phase</i> —Prepare a suitable mixture of <i>pH 3.6 Buffer</i> and acetonitrile (9:1).
1858	Ceftazidime for Injection	Assay	Line 6 under <i>Procedure</i> : Change the formula "25,000[C/W (100- $m-s$)](r_U/r_s)" to: 250,000[C/W (100- $m-s$)](r_U/r_s)
2807	Loratadine Oral Solution	Related compounds	Line 4 under System suitability solution 1: Change "0.002 mg per mL." to: 0.002 mg per mL in Diluent.
2985	Minocycline Hydrochloride	Identification, Infrared Absorption ⟨197K⟩	Line 1: Change "previously dried at 100° for 2 hours." to: [NOTE—Dry the <i>Standard</i> and <i>Sample</i> at 100° for 2 hours prior to use.]
3338	Potassium Chloride Extended-Release Tablets	Assay	Line 8 under Assay preparation 1: Add to the end of the paragraph "Transfer 5.0 mL of the resulting solution to a 100-mL volumetric flask, add 2.0 mL of sodium chloride solution (1 in 5) and 1.0 mL of hydrochloric acid, dilute with water to volume, and mix."
3691	Tetracaine Hydrochloride for Injection	Residue on ignition	Line 7 under <i>Residue on ignition</i> : Change "Heat, gently at first." to: "Heat gently at a temperature as low as practicable."

USP32–NF27 Page	Title	Section	Description
First Supplem	ent to USP32-NF27		
4039	Cefaclor Capsules	Assay	Line 6 under Chromatographic system: Change "which are about 0.8 and 1.0 for the delta-3 isomer and cefaclor, respectively;" to: which are about 0.8 and 1.0 for cefaclor and the delta-3 isomer, respectively;
Second Suppl	ement to USP32-NF2	27	
4220	Azithromycin for Injection	Limit of azithromycin N-oxide	Line 1 under <i>Resolution solution</i> : Change "Dissolve an accurately weighed portion of USP Azithromycin <i>N</i> -Oxide RS in <i>Standard solution</i> to obtain a solution having a concentration of about 0.0015 mg of azithromycin <i>N</i> -oxide and 0.45 mg of azithromycin per mL." to: 0.0015 mg/mL of azithromycin <i>N</i> -oxide and 0.45 mg/mL of azithromycin in <i>Diluent</i> .
4238	Fexofenadine Hydrochloride and Pseudoephedrine Hydrochloride Extended-Release Tablets	Related compounds	Test 3, line 2 under <i>Standard solution</i> : Change "100-mL volumetric flask," to: 50-mL volumetric flask