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Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
RESIDUAL	1. INTRODUCT	USPNF Online Online	31-Jan-2025	1-May-2025	NA	NA	In paragraph 3:

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
HOST CELL ION AND PROTEIN MEA SCOPE SUREMENT IN BIOPHARMAC EUTICALS BY LIQUID CHRO MATOGRAPHY –MASS SPECT ROMETRY							Change Residual HCP ELISA to: HCP ELISA
RESIDUAL 4. SAMPLE PR HOST CELL EPARATION, C PROTEIN MEA HROMATOGR SUREMENT IN APHC BIOPHARMAC SEPARATION, EUTICALS BY AND MASS SP LIQUID CHRO ECTROMETRY MATOGRAPHY ANALYSIS –MASS SPECT ROMETRY	USPNF Online Online		31-Jan-2025	1-May-2025	NA	NA	In paragraph 4 in <i>4.1 Sample Preparation</i> : Change product (or polysorbate) to: product protein (or polysorbate)
RESIDUAL 5. QUANTITATI HOST CELL ON OF HCPS PROTEIN MEA SUREMENT IN BIOPHARMAC EUTICALS BY LIQUID CHRO MATOGRAPHY –MASS SPECT ROMETRY	USPNF Online Online		31-Jan-2025	1-May-2025	NA	NA	In three instances in <i>5.1 Methods for HCP Quantitation</i> : Change product to: product protein
RESIDUAL 7. BEST HOST CELL PRACTICES	USPNF Online Online		31-Jan-2025	1-May-2025	NA	NA	In paragraph 1 in 7.3

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
PROTEIN MEA FOR SUREMENT IN REPORTING BIOPHARMAC DATA FROM EUTICALS BY ELISA VERSUS LIQUID CHRO LC-MS/MS MATOGRAPHY –MASS SPECT ROMETRY DRY HEAT DE INTRODUCTIO PYROGENATI N ON	USPNF Online Online		23-Feb-2024	1-Mar-2024	NA	NA	<i>Comparison of ELISA and LC- MS/MS HCP Results: Change the comparing to: comparing Change parental manufacturing to: parenteral manufacturing In Analysis: Change Add to the acid solution 5.0 mL of Standard dithizone solution and 4 mL of Ammonia cyanide solution, to: Add to the acid solution 5.0 mL of Standard dithizone solution and 4 mL of</i>
LEAD	PROCEDURES <i>/Procedure 1: Chemical Method</i>	USPNF Online Online	27-Jan-2023	1-Jun-2023	NA	NA	

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LEAD	REQUIREMENTS FOR PROCEDURE VALIDATION	USPNF Online Online	27-Jan-2023	1-Jun-2023	NA	NA	Ammonium cyanide solution, Change • Precision Repeatability to: • Precision
TOPICAL AND SPECIFIC TRANSDERMA TESTS FOR L DRUG PROD TDS UCTS—PRODU CT QUALITY TESTS		USPNF Online Online	28-Jul-2023	1-Dec-2023	NA	NA	In Release Liner Peel Test: Change The product fails the test if the mean peel force is outside the acceptable range determined during product development. to: The product fails the test if the overall mean peel force is outside the acceptable range determined during product development.

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PARTICLE SIZE ANALYSIS BY DYNAMIC LIGHT SCATTERING	GLOSSARY	USPNF Online	Online	27-Oct-2023	1-May-2024	NA	NA	In Average particle diameter: Change expressed in nanometers. to: expressed in meters. AND In Viscosity: Change in millipascal-seconds (mPa?s). to: in pascal-seconds (Pa?s).
COLOR AND ACHROMICITY	METHOD II: INSTRUMENTAL (QUANTITATIVE) ASSESSMENT OF COLOR AND COLOR MATCHES	USPNF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	In Table 5: Change Sum 98.809 100.000 107.307 White point 98.811 100.000 107.304 to: Sum 94.809 100.000 107.307 White point

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PACKAGING AND STORAGE REQUIREMENTS	GENERAL <i>Packaging Definitions</i>	<i>Second Supplement to USP43–NF38</i>	Online	26-Mar-2021	1-Dec-2025	NA	NA	94.811 100.000 107.304 In <i>Light-resistant container</i> : Change ?661.2?, <i>Functionality, Spectral Transmission Requirements for Light-Resistant Components and Systems</i> . to: ?661.2?, <i>Functionality Test Method, Spectral Transmission Requirements for Light-Resistant Components and Systems</i> .
ANTIBIOTICS—MICROBIAL ASSAYS	APPENDICES	<i>USPNF Online</i>	Online	29-Dec-2023	1-Jan-2024	NA	NA	In two instances in <i>Appendix 1</i> equations: Change 14.020

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MEASUREME NT OF STRUCTURAL STRENGTH OF SEMISOLIDS BY PENETRO METRY	APPARATUS	USPNF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	to: 14.022 In <i>Figure 2</i> : Change 66±0.25 Ø to: 65±0.25 Ø
ACARBOSE	IMPURITIES	USPNF Online	Online	29-Dec-2023	1-Jan-2024	NA	NA	In <i>Chromatographi c Purity/Analysis</i> : Change Result = (r_U/r_A) × (1/F) × 100 to: Result = (r_U/r_A) × (1/F)
ACESULFAME POTASSIUM	SPECIFIC TESTS	USPNF Online	Online	27-Dec-2024	1-Jan-2025	NA	NA	In <i>Acidity or Alkalini ty/Analysis</i> : Change If the solution is blue, titrate with 0.01 N hydrochloride acid to: If the solution is blue, titrate with

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ADENOSINE	ASSAY	USPNF Online	Online	27-Sep-2024	1-Oct-2024	NA	NA	0.01 N hydrochloric acid In <i>Proce</i> <i>dure/Buffer:</i> Change tetrabutyl ammonium to: tetrabutylammo nium
ALUMINUM SULFATE	CHEMICAL INFORMATION	USPNF Online	Online	25-Oct-2024	1-Nov-2024	NA	NA	Change $\text{Al}_2(\text{SO}_4)_3 \cdot x\text{H}_2\text{O}$ (anhydrous) 342.15 to: $\text{Al}_2(\text{SO}_4)_3 \cdot x\text{H}_2\text{O}$ AND Change Anhydrous 342.16 to: Anhydrous 342.13
AMANTADINE HYDROCHLORIDE	IDENTIFICATION	USPNF Online	Online	25-Aug-2023	1-Sep-2023	NA	NA	In A.: Change <i>Spectroscopic Identification Tests</i> ?197?, <i>Infrared</i>

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AMIKACIN SULFATE	ASSAY	USPNF Online Online	17-Nov-2023	1-Dec-2023	NA	NA	<i>Spectroscopy:</i> 197A, 197K, and 197S to: <i>Spectroscopic</i> <i>Identification</i> <i>Tests</i> ?197?, <i>Infrared</i> <i>Spectroscopy:</i> 197A, 197K, or 197S Procedure for 197S In <i>Proce</i> <i>dure/Analysis:</i> Change C_U = concentration of amikacin in the <i>Sample solution</i> (mg/mL) to: C_U = concentration of Amikacin Sulfate in the <i>Sample solution</i> (mg/mL) In <i>Proce</i>
AMLODIPINE AND	ASSAY	USPNF Online Online	27-Sep-2024	1-Oct-2024	NA	NA	

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BENAZEPRIL H YDROCHLORI DE CAPSULES								<i>dure/Buffer 1:</i> Change tetrabutyl ammonium to: tetrabutylammo nium
AMOXICILLIN BOLUSES	IDENTIFICATIO N	USPNF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	Change <i>Application</i> volume, <i>Developing</i> solvent system, <i>Proce</i> <i>dure</i> —Proceed as directed for the <i>Identification</i> test under <i>Amoxicillin</i> <i>Tablets</i> . to: <i>Application</i> volume—5 µL. <i>Developing</i> solvent system—a mixture of methanol, chloroform, water, and pyridine

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
AMOXICILLIN FOR INJECTABLE SUSPENSION	Identification	USPNF Online Online	27-Oct-2023	1-Nov-2023	NA	NA	(90:80:30:10). <i>Proce</i> <i>dure</i> —Proceed as directed in <i>Thin-Layer Chro</i> <i>matographic</i> <i>Identification</i> <i>Test</i> <201>. Dry the plate with the aid of a current of warm air for 10 minutes. Locate the spots on the plate by spraying lightly with a solution of ninhydrin in alcohol containing 3 mg per mL, and dry at 110° for 15 minutes. Change Prepare a test solution containing the equivalent of 4 mg of amoxicillin per mL by adding

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							0.1 N hydrochloric acid to Amoxicillin for Injectable Suspension. Allow the solution to stand for 5 minutes before use: the solution responds to the <i>Identification</i> test under <i>Amoxicillin Capsules</i>. to: Prepare a test solution containing the equivalent of 4 mg of amoxicillin per mL by adding 0.1 N hydrochloric acid to Amoxicillin for Injectable Suspension.

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							Allow the solution to stand for 5 minutes before use. Prepare a Standard solution of USP Amoxicillin RS in 0.1 N hydrochloric acid containing 4 mg per mL. Use within 10 minutes after preparation. Apply separately 5 µL of each solution on a thin-layer chromatographic plate coated with a 0.25-mm layer of chromatographic silica gel mixture (see <i>Chromatography</i> <621>). Place the plate in a suitable chromatographic chamber, and

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							develop the chromatogram in a solvent system consisting of a mixture of methanol, chloroform, water, and pyridine (90:80:30:10). When the solvent front has moved about three-fourths of the length of the plate, remove the plate from the chamber, and dry with warm air for 10 minutes. Locate the spots on the plate by spraying lightly with a solution of ninhydrin in alcohol containing 3 mg per mL, and dry

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
AMOXICILLIN IDENTIFICATION NTRAMAMMA N RY INFUSION	USP NF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	at 110° for 15 minutes: the R_F value of the principal spot obtained from the test solution corresponds to that obtained from the Standard solution. Change The solution obtained responds to the <i>Identification</i> test under <i>Amoxicillin Capsules</i>. to: Prepare a Standard solution of USP Amoxicillin RS in 0.1 N hydrochloric acid containing 4 mg per mL. Use within 10 minutes after preparation.

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							Apply separately 5 µL of each solution on a thin-layer chromatographic plate coated with a 0.25-mm layer of chromatographic silica gel mixture (see <i>Chromatography</i> <621>). Place the plate in a suitable chromatographic chamber, and develop the chromatogram in a solvent system consisting of a mixture of methanol, chloroform, water, and pyridine (90:80:30:10). When the solvent front has moved about three-

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AMOXICILLIN ORAL SUSPENSION	IDENTIFICATIO N	USPNF Online	26-Jan-2024	1-Feb-2024	NA	NA	fourths of the length of the plate, remove the plate from the chamber, and dry with warm air for 10 minutes. Locate the spots on the plate by spraying lightly with a solution of ninhydrin in alcohol containing 3 mg per mL, and dry at 110° for 15 minutes: the R_F value of the principal spot obtained from the test solution corresponds to that obtained from the Standard solution. Change Shake a portion of Oral Suspension

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							with a mixture of acetone and 0.1 N hydrochloric acid (4:1) to obtain a solution containing about 1 mg of amoxicillin per mL. The solution responds to the <i>Identification</i> test under <i>Amoxicillin Capsules</i>. to: Prepare a test solution by shaking a portion of Oral Suspension with a mixture of acetone and 0.1 N hydrochloric acid (4:1) to obtain a solution containing

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							about 1 mg of amoxicillin per mL. Prepare a Standard solution of USP Amoxicillin RS in 0.1 N hydrochloric acid containing 1 mg per mL. Use within 10 minutes after preparation. Apply separately 5 µL of each solution on a thin-layer chromatographic plate coated with a 0.25-mm layer of chromatographic silica gel mixture (see <i>Chromatography</i> ?621?). Place the plate in a suitable chromatographic chamber, and develop the chromatogram

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							in a solvent system consisting of a mixture of methanol, chloroform, water, and pyridine (90:80:30:10). When the solvent front has moved about three-fourths of the length of the plate, remove the plate from the chamber, and dry with warm air for 10 minutes. Locate the spots on the plate by spraying lightly with a solution of ninhydrin in alcohol containing 3 mg per mL, and dry at 110° for 15 minutes: the <i>R</i>

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							F value of the principal spot obtained from the test solution corresponds to that obtained from the Standard solution. In <i>Dissolution</i> <711>/Test 1/ Proce dure/ Chromatographi c proce dure/Analysis: Change Result = $(R_U/R_S) \times C_S \times V \times (1/L) \times 100$ to: Result = $(R_U/R_S) \times C_S \times V \times D \times (1/L) \times 100$ AND Add D = dilution
ARIPIRAZOL E TABLETS	PERFORMANC E TESTS	USPNF Online	Online	25-Aug-2023	1-Sep-2023	NA	NA

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ATORVASTATIN CALCIUM	ADDITIONAL REQUIREMENTS/USP Reference Standards ?11?	USPNF Online Online	26-May-2023	1-Jun-2023	NA	NA	factor of the <i>Sample solution</i> , 2 Change USP Atorvastatin Related Compound H RS (lactone impurity) to: USP Atorvastatin Related Compound H RS Also known as Lactone impurity; AND Change USP Atorvastatin Related Compound I RS (acetone impurity) to: USP Atorvastatin Related

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ATORVASTATIN CALCIUM TABLETS ASSAY/ <i>Procedure</i>	USPNF Online	Online	30-Jun-2023	1-Jul-2023	NA	NA	Compound I RS Also known as Acetonide impurity; In <i>Analysis</i> : Change M_{r1} = molecular weight of atorvastatin, 558.64 M_{r2} = molecular weight of atorvastatin calcium, 1155.34 to: M_{r1} = molecular weight of atorvastatin, 558.65 M_{r2} = molecular weight of atorvastatin calcium, 1155.36
ATORVASTATIN CALCIUM TABLETS PERFORMANCE TESTS/ <i>Dissolution</i> ?711?	USPNF Online	Online	30-Jun-2023	1-Jul-2023	NA	NA	In <i>Test 1</i> , <i>Test</i> <i>3</i> , <i>Test 4</i> , <i>Test</i> <i>5</i> , and <i>Test</i> <i>6/Analysis</i> : Change <i>M</i>

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							r_1 = molecular weight of atorvastatin, 558.64 M_{r2} = molecular weight of atorvastatin calcium, 1155.34 to: M_{r1} = molecular weight of atorvastatin, 558.65 M_{r2} = molecular weight of atorvastatin calcium, 1155.36
ATORVASTATIN CALCIUM TABLETS	ADDITIONAL REQUIREMENTS/USP Reference Standards ?11?	USPNF Online	Online	30-Jun-2023	1-Jul-2023	NA	NA In USP Atorvastatin Related Compound B RS: Change 1155.34 to: 1155.36
ATORVASTATIN CALCIUM TABLETS	IMPURITIES/Organic Impurities	USPNF Online	Online	30-Jun-2023	1-Jul-2023	NA	NA In Analysis: Change M_{r1} = molecular weight of

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ATOVAQUONE IMPURITIES	USPNF Online	Online	26-May-2023	1-Jun-2023	NA	NA	atorvastatin, 558.64 M_{r2} = molecular weight of atorvastatin calcium, 1155.34 to: M_{r1} = molecular weight of atorvastatin, 558.65 M_{r2} = molecular weight of atorvastatin calcium, 1155.36 Change Related Compounds System suitability solution and Sample solution: Prepare as directed in the Assay. Analysis Samples:

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							<i>System suitability solution and Sample solution</i> Using the chromatograms of the <i>Sample solution</i> and the <i>System suitability solution</i>, calculate the percentage of atovaquone related compounds in the portion of Atovaquone taken: to: Organic Impurities Mobile phase, Diluent, System suitability solution, Standard solution, Sample solution

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							Chromatographic system and System suitability: Proceed as directed in the Assay. Analysis Sample: <i>Sample solution</i> Calculate the percentage of atovaquone related compounds in the portion of Atovaquone taken: In <i>Organic Impurities/Table 2</i>: Change 3'-N-Demet hyl-3'-N-[(4-methylphenyl)sulfonyl]azithromycin^m to:
AZITHROMYCI IMPURITIES N	USPNF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	

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							3?-N -[4-(Acetylamin o)phenyl]sulfon yl}-3?-demethyl azithromycin ^m AND In <i>Table</i> 2/footnote m: Change (2R,3S,4R,5R ,8R,10R,11R ,12S,13S,14R)-13-[(2,6-Dideo xy-3-C -meth yl-3-O-methyl-?- L-ribo -hexopyranosyl) oxy]-2-ethyl-3,4, 10-trihydroxy-3, 5,6,8,10,12,14- heptamethyl-11- [[3-[N -(4-methylpheny lsulfon yl)-N -methylamino]-3 ,4,6-trideoxy-β- D-xyl o -hexopyranosyl] oxy]-1-oxa-6-az

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							acyclopentadecan-15-one. to: (2 <i>R</i> ,3 <i>S</i> ,4 <i>R</i> ,5 <i>R</i> ,8 <i>R</i> ,10 <i>R</i> ,11 <i>R</i> ,12 <i>S</i> ,13 <i>S</i> ,14 <i>R</i>)-13-[(2,6-Dideoxy-3-C-methyl-3- <i>O</i> -methyl-?- <i>L</i> -ribo-hexopyranosyl)oxy]-2-ethyl-3,4,10-trihydroxy-3,5,6,8,10,12,14-heptamethyl-11-[[3-[<i>N</i> -(4-acetamidophenylsulfonyl)- <i>N</i> -methylamino]-3,4,6-trideoxy-?- <i>D</i> -xylo-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one. Change 748.98
AZITHROMYCIN	CHEMICAL INFORMATION	USPNF Online	25-Aug-2023	1-Sep-2023	NA	NA	

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AZITHROMYCI IMPURITIES N	USPNF Online Online		25-Aug-2023	1-Sep-2023	NA	NA	to: 749.00 AND Change 767.00 to: 767.01 AND Change 785.02 to: 785.03 In <i>Organic Impurities/</i>Table 2: Change: 3'-N -[4-(Acetylamino)phenyl]sulfonyl}-3'-demethyl azithromycin^m to: 3'-N -D emet hyl-3'-N -[(4-methylphenyl)sulfonyl]azithromycin^m AND In <i>Organic Impurities/</i>Table

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							2/footnote m: Change (2 <i>R</i> ,3 <i>S</i> ,4 <i>R</i> ,5 <i>R</i> ,8 <i>R</i> ,10 <i>R</i> ,11 <i>R</i> ,12 <i>S</i> ,13 <i>S</i> ,14 <i>R</i>)-13-[(2,6-Dideo xy-3-C -meth yl-3- <i>O</i> -methyl-?- <i>L</i> -ribo -hexopyranosyl) oxy]-2-ethyl-3,4, 10-trihydroxy-3, 5,6,8,10,12,14- heptamethyl-11- [[3- <i>N</i> -(4-acetamidop h en ylsulf onyl)- <i>N</i> -methylamino]-3 ,4,6-trideoxy-?- <i>D</i> -xylo -hexopyranosyl] oxy]-1-oxa-6-az acyclopentadec an-15-one. to: (2 <i>R</i> ,3 <i>S</i> ,4 <i>R</i> ,5 <i>R</i> ,8 <i>R</i> ,10 <i>R</i> ,11 <i>R</i>

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AZITHROMYCIN	ADDITIONAL REQUIREMENTS	USPNF Online	Online	25-Aug-2023	1-Sep-2023	NA	NA	,12S,13S,14R)-13-[(2,6-Dideoxy-3-C-methyl-3-O-methyl-?-L-ribo-hexopyranosyl)oxy]-2-ethyl-3,4,10-trihydroxy-3,5,6,8,10,12,14-heptamethyl-11-[[3-[N-(4-methylphenylsulfon-yl)-N-methylamino]-3,4,6-trideoxy-β-D-xyl-o-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one. In USP Reference Standards ?11?/USP Azithromycin A RS: Change 734.96 to: 734.97

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AZITHROMYCI IMPURITIES N FOR INJECTION	USPNF Online Online		28-Jul-2023	1-Aug-2023	NA	USPNF 2024 Issue 2	AND In USP Azithromycin Related Compound F RS: Change 762.97 to: 762.98 AND In USP Desosa minylazithromyc in RS: Change 590.79 to: 590.80 In footnote m in Table 2: Change (2R,3S,4R,5R ,8R,10R,11R ,12S,13S,14R)-13-[(2,6-Dideo xy-3-C -meth yl-3-O-methyl-?- L-ribo -hexopyranosyl) oxy]-2-ethyl-3,4, 10-trihydroxy-3, 5,6,8,10,12,14-

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							heptamethyl-11- [[3-[<i>N</i> -(4-acetamidop h en ylsulf onyl)- <i>N</i> -methylamino]-3 ,4,6-trideoxy-?- D-xylo -hexopyranosyl] oxy]-1-oxa-6-az acyclopentadec an-15-one. to: (2 <i>R</i> ,3 <i>S</i> ,4 <i>R</i> ,5 <i>R</i> ,8 <i>R</i> ,10 <i>R</i> ,11 <i>R</i> ,12 <i>S</i> ,13 <i>S</i> ,14 <i>R</i>)-13-[(2,6-Dideo xy-3- <i>C</i> -meth yl-3- <i>O</i> -methyl-?- L-ribo -hexopyranosyl) oxy]-2-ethyl-3,4, 10-trihydroxy-3, 5,6,8,10,12,14- heptamethyl-11- [[3-[<i>N</i> -(4-methylpheny lsulfon

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AZITHROMYCIN FOR INJECTION	ADDITIONAL REQUIREMENTS	USPNF Online	28-Jul-2023	1-Aug-2023	NA	USPNF 2024 Issue 2	yl)-N -methylamino]-3 ,4,6-trideoxy-β- D-xyl -hexopyranosyl] oxy]-1-oxa-6-az acyclopentadec an-15-one. In USP <i>Reference</i> <i>Standards</i> ?11?/USP Azae rythromycin A RS: Change 734.96 to: 734.97 AND In USP Azithromycin N-oxide RS: Change 764.98 to: 765.00 AND In USP N -Demethylazithr omycin RS: Change 734.96

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							to: 734.97 AND In USP Desosa minylazithromyc in RS: Change 590.79 to: 590.80 Change carvacol to: carvacrol In <i>Fats and</i> <i>Fixed Oils</i> ?401?, <i>Procedures,</i> <i>Fatty Acid</i> <i>Co</i> <i>mposition/Table</i> 2: Change Linoleic to: Linoleic acid Change Optical Rotation ?781? to: Optical Rotation,
BLACK CUMIN SEED THYMO QUINONE OIL	DEFINITION	USPNF Online	25-Aug-2023	1-Sep-2023	NA	NA	
BLACK CUMIN SEED THYMO QUINONE OIL	SPECIFIC TESTS	USPNF Online	25-Aug-2023	1-Sep-2023	NA	NA	
BROMPHENIR AMINE MALEATE	SPECIFIC TESTS	USPNF Online	26-Apr-2024	1-May-2024	NA	NA	

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
CALCITONIN SALMON	IMPURITIES	USPNF Online Online	27-Sep-2024	1-Oct-2024	NA	NA	<i>Angular Rotation ?781A?</i> <i>In Procedure: Related Peptides and Other Related Sub stances/Test 2: Change Resolution solution: to: System suitability solution: AND In Procedure: Related Peptides and Other Related Sub stances/Test 2/Sample solution: Change 100 mL of Buffer C. to: 100 ?L of Buffer</i>

Monograph Title __Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
CALCITONIN SALMON	SPECIFIC TESTS	USPNF Online	Online	27-Sep-2024	1-Oct-2024	NA	NA	C. In <i>Bioid</i> <i>entity/Medium</i> B: Change 2 mM RPMI 1640 to: RPMI 1640
CALCIUM ASCORBATE	IDENTIFICATIO N/A.	USPNF Online	Online	26-May-2023	1-Aug-2023	NA	NA	Change Characteristic emission lines for calcium at 184.0, 315.9, and 317.9 nm from the <i>Sample solution</i> correspond to those from the <i>Standard</i> <i>solution</i> , as obtained in the Assay. to: Characteristic emission lines for calcium at 184.0, 315.9, and 317.9 nm from the <i>Sample solution</i>

Monograph Title __Sort descending__	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
CALCIUM ASCORBATE	ASSAY/ Procedure	USPNF Online	Online	30-Jun-2023	1-Aug-2023	NA	NA	correspond to those from the <i>Standard solution</i> , as obtained in the <i>Content of Calcium. In Analysis:</i> Change M_{r1} = molecular weight of calcium ascorbate dihydrate, 426.43 to: M_{r1} = molecular weight of calcium ascorbate dihydrate, 426.34
CARVEDILOL TABLETS	PERFORMANCE TESTS	USPNF Online	Online	26-Jan-2024	1-Feb-2024	NA	NA	<i>In Dissolution <711>/Test 3/ Chromatographic system/Column:</i> Change 4.6-mm × 15-mm; 5-?m

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
CEFDINIR	IMPURITIES	USPNF Online Online	17-Nov-2023	1-Dec-2023	NA	NA	packing L7 to: 4.6-mm × 15-cm; 5-?m packing L7 In <i>Organic Impurities/Table 2/footnote a</i> : Change 1N-[(Z)2-(2-Aminothia zol-4-yl)-2-(hydr oxyimino)acetyl] glycine. to: N-[(Z)2-(2-Aminothia zol-4-yl)-2-(hydr oxyimino)acetyl] glycine.
CEFDINIR	ADDITIONAL R EQUIREMENT S	USPNF Online Online	17-Nov-2023	1-Dec-2023	NA	NA	In <i>USP Reference Standards</i> ?11?/USP Cefdinir Related Compound A RS: Change 413.43 to: 413.42
CEFDINIR	ADDITIONAL R	USPNF Online Online	17-Nov-2023	1-Dec-2023	NA	NA	In <i>USP</i>

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
CAPSULES	EQUIREMENTS							Reference Standards ?11?/USP Cefdinir Related Compound A RS: Change 413.43 to: 413.42 AND In <i>USP Reference Standards</i> ?11?/USP Cefdinir Related Compound B RS: Change $C_{14}H_{13}N_4O_4S_2$ 365.41 to: $C_{14}H_{14}N_4O_4S_2$ 366.41
CEFDINIR FOR ADDITIONAL ORAL SUSPENSION	ADDITIONAL REQUIREMENTS	USPNF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	In <i>USP Reference Standards</i> ?11?/USP Cefdinir Related Compound A RS: Change 413.43 to:

Monograph Title Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
CEFUROXIME ASSAY AXETIL FOR ORAL SUSPENSION	<i>USPNF Online</i> Online		31-May-2024	1-Jun-2024	NA	NA	413.42 AND In <i>USP</i> <i>Reference</i> <i>Standards</i> ?11?/USP Cefdinir Related Compound B RS: Change $C_{14}H_{13}N_4O_4S_2$ 365.41 to: $C_{14}H_{14}N_4O_4S_2$ 366.41 In <i>Proce</i> <i>dure/System</i> <i>suitability:</i> Change [Note—The relative retention times for acetanilide, cefuroxime axetil diastereoisomer B, cefuroxime axetil diastereoisomer A, and cefuroxime

Monograph Title Section __Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							axetil delta-3 isomers are 0.4, 0.8, 0.9, and 1.0, respectively.] to: [Note—The relative retention times for cefuroxime axetil diastereoisomer B, cefuroxime axetil diastereoisomer A, and cefuroxime axetil delta-3 isomers are 0.8, 0.9, and 1.0, respectively.]

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