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How to Use

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CELL	5. CELL BANK	*USPNF Online*	Online	31-Mar-2023	1-May-2023	*NA*	*NA*	In *Table 4*:

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
BANKING PRACTICES FOR RECOMBINANT BIOLOGICS	CHARACTERIZATION							Change In vitro assay[d,e] + + +[f] to: In vitro assay[d,e] + ?[f] +
DRY HEAT DEPYROGENATION	INTRODUCTION	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	Change parental manufacturing to: parenteral manufacturing
LOW MOLECULAR WEIGHT HEPARIN MOLECULAR WEIGHT DETERMINATIONS	PROCEDURE	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	In footnote 1 in *Molecular Weight Measurements of Low Molecular Weight Heparins by Gel Permeation Chrom atogr aphy/ Chromatographic system/ Colum ns/Analytical*: Change guard column

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
LEAD	PROCEDURES /*Procedure 1: Chemical Method*	*USPNF Online*	Online	27-Jan-2023	1-Jun-2023	*NA*	*NA*	TSK SWXL 6-mm × 4-mm; to: guard column TSK SWXL 6-mm × 4-cm; 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 *Ammonium cyanide solution*,
LEAD	REQUIREMENTS FOR PROCEDURE VALIDATION	*USPNF Online*	Online	27-Jan-2023	1-Jun-2023	*NA*	*NA*	Change • **Precision Repeatability** to: • **Precision**

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
TOPICAL AND TRANSDERMAL DRUG PRODUCTS—PRODUCT QUALITY TESTS	SPECIFIC TESTS FOR TDS	*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.
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

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 94.811 100.000 107.304
PACKAGING AND STORAGE REQUIREMENTS	GENERAL *Packaging Definitions*	*Second Supplement to USP43–NF38*	Online	26-Mar-2021	1-Dec-2025	*NA*	*NA*	In *Light-resistant container*: Change ?661.2?, *Functionality,*

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Spectral Transmission Requirements for Light-Resistant Components and Systems. to: ?661.2?, *Functionality Test Method, Spectral Transmission Requirements for Light-Resistant Components and Systems*.
ANTIBIOTICS—MICROBIAL ASSAYS	APPENDICES	*USPNF Online*	Online	29-Dec-2023	1-Jan-2024	*NA*	*NA*	In two instances in *Appendix 1* equations: Change 14.020 to: 14.022
MEASUREMENT OF STRUCTURAL STRENGTH OF SEMISOLIDS BY PENETRO	APPARATUS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Figure 2*: Change 66±0.25 Ø to: 65±0.25 Ø

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METRY								
ACARBOSE	IMPURITIES	*USPNF Online*	Online	29-Dec-2023	1-Jan-2024	*NA*	*NA*	In *Chromatographic Purity/Analysis*: Change Result = $(r_U/r_A) \times (1/F) \times 100$ to: Result = $(r_U/r_A) \times (1/F)$
ACARBOSE TABLETS	IDENTIFICATION/*B.*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	Change The spectrum obtained from the *Sample solution* shows IR maxima in the regions of 3500–3200, 2950–2890, 1653–1633, and 1070–1000 cm^{-1}. to: The spectrum obtained from the sample preparation shows IR maxima in the regions of

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								3500–3200, 2950–2890, 1653–1633, and 1070–1000 cm^{-1}.
ACESULFAME POTASSIUM	SPECIFIC TESTS	*USPNF Online*	Online	27-Dec-2024	1-Jan-2025	*NA*	*NA*	In *Acidity or Alkalinity*/*Analysis*: Change If the solution is blue, titrate with 0.01 N hydrochloride acid to: If the solution is blue, titrate with 0.01 N hydrochloric acid
ACYCLOVIR	ASSAY/ *Procedure*	*USPNF Online*	Online	29-Apr-2022	1-May-2023	*NA*	*NA*	In the *Sample solution*: Change 0.1 N sodium hydroxide to: 0.01 N sodium hydroxide
ADENOSINE	ASSAY	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Procedure*/*Buffer*:

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Change tetrabutyl ammonium to: tetrabutylammonium
AMANTADINE HYDROCHLORIDE	IDENTIFICATION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *A.*: Change *Spectroscopic Identification Tests* ?197?, *Infrared Spectroscopy*: 197A, 197K, and 197S to: *Spectroscopic Identification Tests* ?197?, *Infrared Spectroscopy*: 197A, 197K, or 197S
AMIKACIN SULFATE	ASSAY	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	**Procedure for 197S** In *Procedure/Analysis*: Change C_U = concentration of amikacin in the

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Sample solution (mg/mL) to: C_U = concentration of Amikacin Sulfate in the *Sample solution* (mg/mL)
AMLODIPINE AND BENAZEPRIL HYDROCHLORIDE CAPSULES	ASSAY	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Procedure*/*Buffer 1*: Change tetrabutyl ammonium to: tetrabutylammonium
AMOXICILLIN BOLUSES	IDENTIFICATION	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	Change *Application volume*, *Developing solvent system*, *Procedure*—Proceed as directed for the *Identification* test under *Amoxicillin Tablets*.

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								to: *Application volume*—5 µL. *Developing solvent system*—a mixture of methanol, chloroform, water, and pyridine (90:80:30:10). *Procedure*—Proceed as directed in *Thin-Layer Chromatographic Identification Test* <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

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								per mL, and dry at 110° for 15 minutes.
AMOXICILLIN FOR INJECTABLE SUSPENSION	Identification	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	Change 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. Allow the solution to stand for 5 minutes before use: the solution responds to the *Identification* test under *Amoxicillin Capsules*. to: Prepare a test solution

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								containing the equivalent of 4 mg of amoxicillin per mL by adding 0.1 N hydrochloric acid to Amoxicillin for Injectable Suspension. 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 c hromatographic

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								plate coated with a 0.25-mm layer of chromatographic silica gel mixture (see *Chromatography* <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-fourths of the length of the plate, remove the plate from the chamber,

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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.
AMOXICILLIN INTRAMAMMARY INFUSION	IDENTIFICATION	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	Change The solution obtained responds to the *Identification* test under *Amoxicillin Capsules*. to:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
Sort descending								
								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 *Chromatography* <621>). Place the plate in a suitable chromatographic chamber, and develop the chromatogram in a solvent system

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 R_F value of the principal spot

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								obtained from the test solution corresponds to that obtained from the Standard solution.
AMOXICILLIN ORAL SUSPENSION	IDENTIFICATION	*USPNF Online*	Online	26-Jan-2024	1-Feb-2024	*NA*	*NA*	Change Shake a portion of Oral Suspension 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 *Identification* test under *Amoxicillin Capsules*. to: Prepare a test solution by

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								shaking a portion of Oral Suspension 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. 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 c hromatographic plate coated with a 0.25-mm

Monograph Title	Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								layer of chromatographic silica gel mixture (see *Chromatography* ?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-fourths of the length of the plate, remove the plate from the chamber, and dry with warm air for 10

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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.
ARIPIPRAZOLE TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Dissolution <711>/Test 1/ Procedure/ Chromatographic procedure/Analysis*: Change Result =

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ATORVASTATIN CALCIUM	ADDITIONAL REQUIREMENTS/*USP Reference Standards* ?11?	*USPNF Online* Online		26-May-2023	1-Jun-2023	*NA*	*NA*	$(R_U/R_S) \times C_S \times V \times (1/L) \times 100$ to: Result = $(R_U/R_S) \times C_S \times V \times D \times (1/L) \times 100$ AND Add *D* = dilution factor of the *Sample solution*, 2 Change USP Atorvastatin Related Compound H RS (lactone impurity) to: USP Atorvastatin Related Compound H RS Also known as Lactone impurity; AND Change

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
ATORVASTATIN CALCIUM TABLETS	PERFORMANCE TESTS/ *Dissolution* ?711?	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	USP Atorvastatin Related Compound I RS (acetonide impurity) to: USP Atorvastatin Related Compound I RS Also known as Acetonide impurity; In *Test 1*, *Test 3*, *Test 4*, *Test 5*, and *Test 6*/*Analysis*: Change M_{r1} = molecular weight of atorvastatin, 558.64 M_{r2} = molecular weight of atorvastatin calcium, 1155.34 to: M_{r1} = molecular weight of

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 atorvastatin, 558.64 M_{r2} = molecular weight of atorvastatin calcium, 1155.34 to: M_{r1} = molecular weight of atorvastatin, 558.65

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								M_{r2} = molecular weight of atorvastatin calcium, 1155.36
ATORVASTATIN CALCIUM TABLETS	ASSAY/ *Procedure*	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	In *Analysis*: 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
ATOVAQUONE	IMPURITIES	*USPNF Online*	Online	26-May-2023	1-Jun-2023	*NA*	*NA*	Change **Related Compounds System suitability**

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								solution and **Sample solution**: Prepare as directed in the *Assay*. **Analysis Samples**: *System suitability solution* and *Sample solution* Using the chromatograms of the *Sample solution* and the *System suitability solution*, calculate the percentage of atovaquone related compounds in the portion of Atovaquone taken: to: **Organic Impurities**

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AZITHROMYCIN	IMPURITIES	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	**Mobile phase**, **Diluent**, **System suitability solution**, **Standard solution**, **Sample solution**, **Chromatographic system** and **System suitability**: Proceed as directed in the *Assay*. **Analysis Sample**: *Sample solution* Calculate the percentage of atovaquone related compounds in the portion of Atovaquone taken: In *Organic Impurities*/*Table 2*: Change

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								3'-*N*-Demethyl-3'-*N*-[(4-methylphenyl)sulfonyl]azithromycin[m] to: 3?-*N*-{[4-(Acetylamino)phenyl]sulfonyl}-3?-demethyl azithromycin[m] AND In *Table 2*/footnote m: Change (2*R*,3*S*,4*R*,5*R*,8*R*,10*R*,11*R*,12*S*,13*S*,14*R*)-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*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								-(4-methylphenylsulfonyl)-*N*-methylamino]-3,4,6-trideoxy-ß-D-*xylo*-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one. to: (2*R*,3*S*,4*R*,5*R*,8*R*,10*R*,11*R*,12*S*,13*S*,14*R*)-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-acetamidophenylsulfonyl)-*N*-methylamino]-3

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								,4,6-trideoxy-?-D-*xylo*-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one.
AZITHROMYCIN	CHEMICAL INFORMATION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change 748.98 to: 749.00 AND Change 767.00 to: 767.01 AND Change 785.02 to: 785.03
AZITHROMYCIN	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Organic Impurities*/*Table 2*: Change: 3?-*N*-{[4-(Acetylamino)phenyl]sulfonyl}-3?-demethylazithromycin[m] to: 3’-*N*-D

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								emethyl-3'-*N*-[(4-methylphenyl)sulfonyl]azithromycin[m] AND In *Organic Impurities*/*Table 2*/footnote m: Change (2*R*,3*S*,4*R*,5*R*,8*R*,10*R*,11*R*,12*S*,13*S*,14*R*)-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-acetamidophenylsulfonyl)-*N*-methylamino]-3,4,6-trideoxy-?-

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								D-*xylo*-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one. to: (2*R*,3*S*,4*R*,5*R*,8*R*,10*R*,11*R*,12*S*,13*S*,14*R*)-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-methylphenylsulfonyl)-*N*-methylamino]-3,4,6-trideoxy-ß-D-*xylo*-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one.
AZITHROMYCI	ADDITIONAL R	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *USP*

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N	EQUIREMENTS							*Reference Standards* ?11?/USP Azaerythromycin A RS: Change 734.96 to: 734.97 AND In USP Azithromycin Related Compound F RS: Change 762.97 to: 762.98 AND In USP Desosaminylazithromycin RS: Change 590.79 to: 590.80
AZITHROMYCIN FOR INJECTION	IMPURITIES	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*USPNF 2024 Issue 2*	In footnote m in *Table 2*: Change (2*R*,3*S*,4*R*,5*R*,8*R*,10*R*,11*R*,12*S*,13*S*,14*R*)-13-[(2,6-Dideo

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								xy-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-acetamidophenylsulfonyl)-*N*-methylamino]-3,4,6-trideoxy-?-D-*xylo*-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one. to: (2*R*,3*S*,4*R*,5*R*,8*R*,10*R*,11*R*,12*S*,13*S*,14*R*)-13-[(2,6-Dideoxy-3-*C*-methyl-3-*O*-methyl-?-L-*ribo*

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								-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-ß- D-*xylo* -hexopyranosyl] oxy]-1-oxa-6-az acyclopentadec an-15-one.
AZITHROMYCIN FOR INJECTION	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*USPNF 2024 Issue 2*	In *USP Reference Standards* ?11?/USP Azaerythromycin A RS: Change 734.96 to: 734.97 AND In USP Azithromycin *N*-oxide RS: Change 764.98

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								to: 765.00 AND In USP *N* -Demethylazithromycin RS: Change 734.96 to: 734.97 AND In USP Desosaminylazithromycin RS: Change 590.79 to: 590.80
AZTEC MARIGOLD ZEAXANTHIN EXTRACT	IDENTIFICATION/*C*.	*USPNF Online*	Online	27-Jan-2023	1-Feb-2023	*NA*	*NA*	Change (3*R*,3?*S* *meso*)-zeaxanthin to: (3*R*,3?*S*)-zeaxanthin
AZTEC MARIGOLD ZEAXANTHIN EXTRACT	COMPOSITION /*Procedure 4: Stereoisomeric Composition*	*USPNF Online*	Online	27-Jan-2023	1-Feb-2023	*NA*	*NA*	In *System suitability*/*Suitability requirements*/*Resolution*:

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
BISOPROLOL FUMARATE	CHEMICAL INFORMATION	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	Change (3*R*,3?*S* *meso*)-zeaxanthin to: (3*R*,3?*S*)-zeaxanthin AND In *Analysis*: Change (3*R*,3?*S* *meso*)-zeaxanthin to: (3*R*,3?*S*)-zeaxanthin AND In *Acceptance criteria*: Change (3*R*,3?*S* *meso*)-Zeaxanthin to: (3*R*,3?*S*)-Zeaxanthin Change (±)-1-[[?-(2-Isop *p*

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								-tolyl]oxy]-3-(iso propyl amino)-2 -propanol fumarate (2:1) (salt) to: (±)-1-[[?-(2-Isop *p* -tolyl]oxy]-3-(iso propylamino)-2- propanol fumarate (2:1) (salt)
BLACK CUMIN SEED THYMO QUINONE OIL	DEFINITION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change carvacol to: carvacrol
BLACK CUMIN SEED THYMO QUINONE OIL	SPECIFIC TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Fats and Fixed Oils* ?401?, *Procedures*, *Fatty Acid Co mposition*/*Table 2*: Change Linoleic to: Linoleic acid

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BROMPHENIRAMINE MALEATE	SPECIFIC TESTS	*USPNF Online*	Online	26-Apr-2024	1-May-2024	*NA*	*NA*	Change **Optical Rotation** ?781? to: **Optical Rotation**, *Angular Rotation* ?781A?
BUPIVACAINE HYDROCHLORIDE INJECTION	ASSAY/ *Procedure*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	In *Chromatographic system*/*Column*: Change 4-mm × 30-cm; packing L1 to: 3.9-mm × 30-cm; packing L1 AND In *System suitability*: Change [Note—The relative retention times for bupivacaine hydrochloride and dibutyl

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								phthalate are about 1.0 and 1.2, respectively.] to: [Note—The relative retention times for bupivacaine and dibutyl phthalate are about 1.0 and 1.2, respectively.] AND In System suitability/Suitability requirements: Change **Resolution**: NLT 2.0 between bupivacaine hydrochloride and dibutyl phthalate **Relative standard deviation**: NMT 1.0% for the

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								ratio of bupivacaine to the internal standard from three replicate injections to: **Resolution**: NLT 2.0 between bupivacaine and dibutyl phthalate **Relative standard deviation**: NMT 1.0% for the peak response ratio of bupivacaine to the internal standard from three replicate injections
CALCITONIN SALMON	IMPURITIES	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Procedure: Related Peptides and Other Related Sub stances*/*Test 2*:

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CALCITONIN SALMON	SPECIFIC TESTS	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	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 C*. In *Bioid entity*/*Medium B*: Change 2 mM *RPMI 1640* to: *RPMI 1640*

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CALCIUM ASCORBATE	IDENTIFICATION/*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 *Sample solution* correspond to those from the *Standard solution*, as obtained in the *Assay*. to: Characteristic emission lines for calcium at 184.0, 315.9, and 317.9 nm from the *Sample solution* correspond to those from the *Standard solution*, as obtained in the *Content of Calcium*.
CALCIUM ASCORBATE	ASSAY/ *Procedure*	*USPNF Online*	Online	30-Jun-2023	1-Aug-2023	*NA*	*NA*	In *Analysis*: Change

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								M_{r1} = molecular weight of calcium ascorbate dihydrate, 426.43 to: M_{r1} = molecular weight of calcium ascorbate dihydrate, 426.34
CALCIUM LACTATE	CHEMICAL INFORMATION	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	Change Calcium lactate (1:2) hydrate to: Calcium lactate hydrate AND Change Calcium lactate (1:2) pentahydrate to: Calcium lactate pentahydrate
CARVEDILOL TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	26-Jan-2024	1-Feb-2024	*NA*	*NA*	In *Dissolution* <711>/*Test 3*/ *Chromatographi*

Monograph Title Sort descending	Section	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*	*c* *system*/*Column*: Change 4.6-mm × 15-mm; 5-?m packing L7 to: 4.6-mm × 15-cm; 5-?m packing L7 In *Organic Impurities*/*Table 2*/footnote a: Change 1*N*-[(*Z*)-2-(2-Aminothiazol-4-yl)-2-(hydroxyimino)acetyl] glycine. to: *N*-[(*Z*)-2-(2-Aminothiazol-4-yl)-2-(hydroxyimino)acetyl] glycine.

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