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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] +
RESIDUAL HOST CELL PROTEIN MEASUREMENT IN BIOPHARMACEUTICALS BY LIQUID CHROMATOGRAPHY–MASS SPECTROMETRY	1. INTRODUCTION AND SCOPE	*USPNF Online*	Online	31-Jan-2025	1-May-2025	*NA*	*NA*	In paragraph 3: Change Residual HCP ELISA to: HCP ELISA
RESIDUAL HOST CELL PROTEIN MEASUREMENT IN BIOPHARMACEUTICALS BY LIQUID CHROMATOGRAPHY–MASS SPECTROMETRY	4. SAMPLE PREPARATION, CHROMATOGRAPHIC SEPARATION, AND MASS SPECTROMETRY ANALYSIS	*USPNF Online*	Online	31-Jan-2025	1-May-2025	*NA*	*NA*	In paragraph 4 in *4.1 Sample Preparation*: Change product (or polysorbate) to: product protein (or polysorbate)
RESIDUAL HOST CELL PROTEIN MEASUREMENT IN BIOPHARMAC	5. QUANTITATION OF HCPS	*USPNF Online*	Online	31-Jan-2025	1-May-2025	*NA*	*NA*	In three instances in *5.1 Methods for HCP Quantitation*:

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
EUTICALS BY LIQUID CHRO MATOGRAPHY –MASS SPECT ROMETRY								Change product to: product protein
RESIDUAL HOST CELL PROTEIN MEA SUREMENT IN BIOPHARMAC EUTICALS BY LIQUID CHRO MATOGRAPHY –MASS SPECT ROMETRY	7. BEST PRACTICES FOR REPORTING DATA FROM ELISA VERSUS LC-MS/MS	*USPNF Online*	Online	31-Jan-2025	1-May-2025	*NA*	*NA*	In paragraph 1 in *7.3 Comparison of ELISA and LC-MS/MS HCP Results*: Change the comparing to: comparing
DRY HEAT DE PYROGENATI ON	INTRODUCTIO N	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	Change parental manufacturing to: parenteral manufacturing
LEAD	PROCEDURES /*Procedure 1: Chemical Method*	*USPNF Online*	Online	27-Jan-2023	1-Jun-2023	*NA*	*NA*	In *Analysis*: Change Add to the acid solution 5.0 mL of *Standard dithizone solution* and 4 mL of *Ammonia cyanide solution*,

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								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**
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

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								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 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	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Table 5*: Change Sum 98.809 100.000 107.307 White point

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	AND COLOR MATCHES							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, Spectral Transmission Requirements for Light-Resistant Components and Systems.* to: ?661.2?, *Functionality Test Method, Spectral Transmission Requirements for Light-Resistant*

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								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 PENETROMETRY	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
ACARBOSE	IMPURITIES	*USPNF Online*	Online	29-Dec-2023	1-Jan-2024	*NA*	*NA*	In *Chromatographic Purity/Analysis*: Change Result = (r_U/r_A) × (1/*F*) × 100 to: Result = (r_U/r_A) × (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

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								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 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

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								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*: Change tetrabutyl ammonium to: tetrabutylammonium
ALUMINUM SULFATE	CHEMICAL INFORMATION	*USPNF Online*	Online	25-Oct-2024	1-Nov-2024	*NA*	*NA*	Change $Al_2(SO_4)_3 \cdot xH_2O$ (anhydrous) 342.15 to: $Al_2(SO_4)_3 \cdot xH_2O$ AND Change Anhydrous

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AMANTADINE HYDROCHLORIDE	IDENTIFICATION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	342.16 to: Anhydrous 342.13 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 *Sample solution* (mg/mL)

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								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*. to: *Application*

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Sort descending								*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 per mL, and dry at 110° for 15

Monograph Title Sort descending	Section	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*	minutes. 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 containing the equivalent of 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
								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 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.
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: Prepare a Standard

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 consisting of a mixture of

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 obtained from the test solution

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								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 shaking a portion of Oral

Monograph Title	Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 layer of chromat ographic silica

Monograph Title	Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 minutes. Locate the spots on 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
								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.
APIGENIN	CHEMICAL INFORMATION	*USPNF Online*	Online	28-Feb-2025	1-Aug-2025	*NA*	*NA*	Change $C_{15}H_{12}O_5$ 272.26 to: $C_{15}H_{10}O_5$ 270.24
ARIPIPRAZOLE TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Dissolution <711>/Test 1/ Procedure/ Chromatographic*

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

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ATORVASTATIN CALCIUM TABLETS	ASSAY/ *Procedure*	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	Lactone impurity; AND Change USP Atorvastatin Related Compound I RS (acetonide impurity) to: USP Atorvastatin Related Compound I RS Also known as Acetonide impurity; In *Analysis*: Change M_{r1} = molecular weight of atorvastatin, 558.64 M_{r2} = molecular weight of atorvastatin calcium, 1155.34 to: *M*

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								M_{r1}= molecular weight of atorvastatin, 558.65 M_{r2} = molecular weight of atorvastatin calcium, 1155.36
ATORVASTATIN CALCIUM TABLETS	PERFORMANCE TESTS/ *Dissolution* ?711?	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	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 atorvastatin, 558.65 M_{r2} = molecular weight of atorvastatin

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								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 M_{r2} = molecular weight of atorvastatin calcium, 1155.36

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ATOVAQUONE IMPURITIES		*USPNF Online*	Online	26-May-2023	1-Jun-2023	*NA*	*NA*	Change **Related Compounds System suitability 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

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								Atovaquone taken: to: **Organic Impurities** **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

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AZITHROMYCIN	IMPURITIES	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	Atovaquone taken: In *Organic Impurities*/*Table 2*: Change 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)

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								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. 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-acetamidop

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								h en ylsulf onyl)-*N* -methylamino]-3 ,4,6-trideoxy-?- D-*xylo* -hexopyranosyl] oxy]-1-oxa-6-az acyclopentadec an-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-(Acetylamin o)phenyl]sulfon

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								yl}-3?-demethyl azithromycin[m] to: 3'-*N*-Demethyl-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-acetamidoph

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								en ylsulf onyl)-*N* -methylamino]-3 ,4,6-trideoxy-?- D-*xylo* -hexopyranosyl] oxy]-1-oxa-6-az acyclopentadec an-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-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-ß- D-*xylo*

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								-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one.
AZITHROMYCIN	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *USP 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	IMPURITIES	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*USPNF 2024 Issue 2*	In footnote m in *Table 2*:

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INJECTION								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-?-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*

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								)-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.
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

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AZITHROMYCIN TABLETS	IMPURITIES	*USPNF Online*	Online	28-Mar-2025	1-Apr-2025	*NA*	*NA*	In USP Azithromycin *N*-oxide RS: Change 764.98 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 In *Organic Impurities/ Analysis/Samples*: Change *Standard solution*, *Sample*

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								solution, and *Blank* to: *Peak identification solution*, *Standard solution*, *Sample solution*, and *Blank*
BISOPROLOL FUMARATE	CHEMICAL INFORMATION	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	Change (±)-1-[[?-(2-Isop *p* -tolyl]oxy]-3-(iso propyl amino)-2 -propanol fumarate (2:1) (salt) to: (±)-1-[[?-(2-Isop *p* -tolyl]oxy]-3-(iso propylamino)-2- propanol

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								fumarate (2:1) (salt)
BLACK CUMIN SEED THYMOQUINONE OIL	DEFINITION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change carvacol to: carvacrol
BLACK CUMIN SEED THYMOQUINONE OIL	SPECIFIC TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Fats and Fixed Oils* ?401?, *Procedures*, *Fatty Acid Composition*/*Table 2*: Change Linoleic to: Linoleic acid
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*:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 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.]

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 ratio of bupivacaine to the internal standard from three replicate injections to: **Resolution**: NLT 2.0 between bupivacaine and dibutyl phthalate **Relative**

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								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*: Change **Resolution solution:** to: **System suitability solution:** AND In *Procedure: Related Peptides and Other Related Sub stances*/*Test*

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								2/*Sample solution*: Change 100 mL of *Buffer C*. to: 100 ?L of *Buffer C*.
CALCITONIN SALMON	SPECIFIC TESTS	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Bioid entity*/*Medium B*: Change 2 mM *RPMI 1640* to: *RPMI 1640*
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:

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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*.

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