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How to Use

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AZITHROMYCI	IMPURITIES	*USPNF Online*	Online	28-Mar-2025	1-Apr-2025	*NA*	*NA*	In *Organic*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
N TABLETS								*Impurities/ Analysis/Samples*: Change *Standard solution*, *Sample solution*, and *Blank* to: *Peak identification solution*, *Standard solution*, *Sample solution*, and *Blank*
IOPAMIDOL INJECTION	USP REFERENCE STANDARDS <11>	*USPNF Online*	Online	28-Mar-2025	1-Apr-2025	*NA*	*NA*	In USP Iopamidol Related Compound B RS: Change *N*,*N* ?-bis[2-hydroxy-1-(hydroxymethyl)ethyl]-2,4,6-tri

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								iodoisophthalamide. $C_{16}H_{20}I_3N_3O_7$ 747.07 to: N^1,N^3-Bis(1,3-dihydroxypropan-2-yl)-5-(2-hydroxyacetamido)-2,4,6-triiodoisophthalamide. $C_{16}H_{20}I_3N_3O_8$ 763.06
COSYNTROPIN	IDENTIFICATION	*USPNF Online*	Online	28-Mar-2025	1-Apr-2025	*NA*	*NA*	In *B. Amino Acid Analysis*/*System suitability*/*Resolution*: Change (*hp* ? *hv*)/*hp* × 100 ± 90%, to: (*hp* ? *hv*)/*hp* × 100 ? 90%,
PSEUDOEPHEDRINE HYDROCHLORIDE TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	28-Mar-2025	1-Apr-2025	*NA*	*NA*	In *Dissolution, Procedure for a Pooled Sample* <*711*>/*Chromatographi*

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								c system: Change **Flow rate:** 1.5 mL/min to: **Flow rate:** 1.5 mL/min **Injection volume:** 10 µL In *Acidity/ Sample solution*: Change 500 mg/mL of Dimethyl Sulfoxide in water to: 50.0 g of Dimethyl Sulfoxide in 100 mL of water
DIMETHYL SULFOXIDE	SPECIFIC TESTS	*USPNF Online*	Online	28-Mar-2025	1-Apr-2025	*NA*	*NA*	
TIOCONAZOLE	CHEMICAL INFORMATION	*USPNF Online*	Online	28-Feb-2025	1-Mar-2025	*NA*	*NA*	Change 1-[2,4-Dichloro-[?-(2-chloro-3-thenyl)-oxy]phenethyl]imidazole to: 1-[2,4-Dichloro-[?-(2-chloro-3-th

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								enyl)oxy]phenethyl]imidazole
TIOCONAZOLE	IMPURITIES	*USPNF Online*	Online	28-Feb-2025	1-Mar-2025	*NA*	*NA*	In *Organic Impurities/ Procedure/Analysis*: Change 1-[2,4-dichloro-?-[(3-thenyl)-oxy]phenethyl]imidazole hydrochloride (tioconazole related compound A), 1-[2,4-dichloro-?-[(2,5-dichloro-3-thenyl)-oxy]phenethyl]imidazole hydrochloride (tioconazole related compound B), and 1-[2,4-dichloro-?-[(5-bromo-2-chloro-3-thenyl)-oxy]phenethyl]imidazole hydrochloride (tioconazole

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								related compound C) to: tioconazole related compound A, tioconazole related compound B, and tioconazole related compound C
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
TIOCONAZOLE	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	28-Feb-2025	1-Mar-2025	*NA*	*NA*	In *USP Reference Standards* <11>/USP Tioconazole Related Compound A RS: Change 1-[2,4-Dichloro-?-[(3-thenyl)-oxy]phenethyl]imidazole

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								hydrochloride. to: 1-[2-(2,4-Dichlorophenyl)-2-(thiophen-3-ylmethoxy)ethyl]-1*H*-imidazole hydrochloride. AND In *USP Reference Standards* <11>/USP Tioconazole Related Compound B RS: Change 1-[2,4-Dichloro-?-[(2,5-dichloro-3-thenyl)oxy]phenethyl]imidazole hydrochloride. to: 1-{2-(2,4-Dichlorophenyl)-2-[(2,5-dichlorothiophen-3-yl)methoxy]ethyl}-1*H*-imidazole

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								hydrochloride. AND In *USP Reference Stan dards* <11>/USP Tioconazole Related Compound C RS: Change 1-[2,4-Dichloro-?-[(5-bromo-2-chloro-3-thenyl)-oxy]-phenethyl]imidazole hydrochloride. to: 1-{2-[(5-Bromo-2-chlorothiophen-3-yl)methoxy]-2-(2,4-dichlorophe nyl)ethyl}-1*H*-imidazole hydrochloride.
RESIDUAL HOST CELL PROTEIN MEASUREMENT IN	1. INTRODUCTION AND SCOPE	*USPNF Online*	Online	31-Jan-2025	1-May-2025	*NA*	*NA*	In paragraph 3: Change Residual HCP ELISA

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BIOPHARMACEUTICALS BY LIQUID CHROMATOGRAPHY–MASS SPECTROMETRY								to: HCP ELISA
RESIDUAL HOST CELL PROTEIN MEASUREMENT IN BIOPHARMACEUTICALS BY LIQUID CHROMATOGRAPHY–MASS SPECTROMETRY	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
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)
HYDROXYPROPYL BETADEX	IMPURITIES	*USPNF Online*	Online	31-Jan-2025	1-Feb-2025	*NA*	*NA*	In *Limit of Betadex, Propylene Glycol, and Other Related*

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								Sub stances/ Chromatographi c system: Change **Column temperature:** 40° to: **Temperatures Detector:** 40° **Columns:** 40°
RESIDUAL HOST CELL PROTEIN MEA SUREMENT IN BIOPHARMAC EUTICALS BY LIQUID CHRO MATOGRAPHY –MASS SPECT ROMETRY	5. QUANTITATI ON OF HCPS	*USPNF Online*	Online	31-Jan-2025	1-May-2025	*NA*	*NA*	In three instances in *5.1 Methods for HCP Quantitation*: Change product to: product protein
DIBUCAINE	CHROMATOG RAPHIC PURITY	*USPNF Online*	Online	27-Dec-2024	1-Jan-2025	*NA*	*NA*	Change Proceed as directed for *Chromatographi c purity* under *Dibucaine Hydrochloride*, to:

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OXALIPLATIN	IMPURITIES	*USPNF Online*	Online	27-Dec-2024	1-Jan-2025	*NA*	*NA*	Proceed as directed for *Organic Impurities* under *Dibucaine Hydrochloride*, In *Organic Impurities, Procedure 1/Analysis*: Change Result = $(r_U/r_S) \times (C_S/(C_U) \times (M_{r1}/M_{r2}) \times 100$ to: Result = $(r_U/r_S) \times (C_S/C_U) \times (M_{r1}/M_{r2}) \times 100$
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

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								hydrochloric acid
INDOMETHACIN SODIUM	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	22-Nov-2024	1-Dec-2024	*NA*	*NA*	In *USP Reference Standards ?11?*/USP Indomethacin Related Compound A RS: Change ($C_{12}H_{13}NO_3$) to: $C_{12}H_{13}NO_3$ AND In *USP Reference Standards ?11?*/USP Indomethacin Related Compound B RS: Change ($C_7H_5ClO_2$) to: $C_7H_5ClO_2$
TAMSULOSIN HYDROCHLORIDE CAPSULES	ASSAY	*USPNF Online*	Online	22-Nov-2024	1-Dec-2024	*NA*	*NA*	In *Procedure 1/Analysis*: Change C_S = concentration of USP

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								Tamsulosin Hydrochloride RS in the *Standard stock solution* (mg/mL) to: C_S = concentration of USP Tamsulosin Hydrochloride RS in the *Standard solution* (mg/mL)
TAMSULOSIN HYDROCHLORIDE CAPSULES	PERFORMANCE TESTS	*USPNF Online*	Online	22-Nov-2024	1-Dec-2024	*NA*	*NA*	In *Dissolution ?711?/Test 1/Analysis*: Change $Result_3 = \{[(C_3 \times V_2) + (C_2 + V_S)] \times (1/L) \times 100\} + Result_1$ to: $Result_3 = \{[(C_3 \times V_2) + (C_2 \times V_S)] \times (1/L) \times 100\} + Result_1$
FLUCONAZOLE	IMPURITIES	*USPNF Online*	Online	25-Oct-2024	1-Nov-2024	*NA*	*NA*	In *Organic Impuri*

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								ties/*Analysis*: Change Calculate the percentage of any individual unspecified impurity in the portion of Fluconazole taken: to: Calculate the percentage of any other specified and unspecified impurity in the portion of Fluconazole taken: AND Change r_U = peak response of any individual impurity from the *Sample solution* to: r_U = peak response of any

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								other specified and unspecified impurity from the *Sample solution*
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 342.16 to: Anhydrous 342.13
CALCITONIN SALMON	SPECIFIC TESTS	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Bioidentity/Medium B*: Change 2 mM *RPMI 1640* to: *RPMI 1640*
METHOTREXATE	ASSAY	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Procedure/Buffer*: Correct the

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								reagent cross reference 1 N sodium hydroxide to: 1 N sodium hydroxide
RALBUMIN HUMAN	IMPURITIES	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Limit of High Molecular Weight Proteins*: Change **Acceptance criteria** **Individual impurities:** NMT 1.0% to: **Acceptance criteria:** NMT 1.0%
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

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FLUCONAZOLE TABLETS	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *USP Reference Standards <11>*/USP Fluconazole Related Compound B RS: Change $C_{13}H_{13}FN_9O$ to: $C_{13}H_{13}FN_6O$
PHENOXYBENZAMINE HYDROCHLORIDE	ASSAY	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Procedure*/*Solution A*: Change sodium phosphate monobasic to: sodium phosphate, monobasic, anhydrous
CETIRIZINE HYDROCHLORIDE	IMPURITIES	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Organic Impurities, Procedure 2*/*Solution A*: Change tetrabutyl ammonium to:

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								tetrabutylammonium
MOXIFLOXACIN TABLETS	IMPURITIES	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Organic Impurities*/*Buffer*: Change tetrabutyl ammonium to: tetrabutylammonium AND In *Organic Impurities*/*Solution C*: Change tetrabutyl ammonium to: tetrabutylammonium
CALCITONIN SALMON	IMPURITIES	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Procedure: Related Peptides and Other Related Substances*/*Test 2*: Change **Resolution solution:** to:

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								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*.
KETOCONAZOLE	ASSAY	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Procedure*/*Buffer*: Change tetrabutyl ammonium to: tetrabutylammonium
RALBUMIN HUMAN	ASSAY	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Albumin Content*/*Native stock running buffer*: Change

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ADENOSINE	ASSAY	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	glycerol to: glycine In *Procedure*/*Buffer*: Change tetrabutyl ammonium to: tetrabutylammonium
CETIRIZINE HYDROCHLORIDE TABLETS	IMPURITIES	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Organic Impurities*/*Buffer*: Change tetrabutyl ammonium to: tetrabutylammonium
PACLITAXEL	IMPURITIES	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Organic Impurities*/*Table 4*: Change [a](2a*R*,4*S*,4a*S*,6*R*,9*S*,11*S*,12*S*,12a*R*,12b*S*)-12b-(Acetyloxy)-12-(benzyloxy)-1,2a,3,4,4a,6

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								,9,10,11,12,12a,12b-dodecahydro-4,6,9,11-tetrahydroxy-4a,8,13,13-tetramethyl-
								H-cyclodeca[3,4]benz[1,2-*b*]oxet-5-one. to: [a](2a*R*,4*S*,4a*S*,6*R*,9*S*,11*S*,12*S*,12a*R*,12b*S*)-12b-(Acetyloxy)-12-(benzoyloxy)-1,2a,3,4,4a,6,9,10,11,12,12a,12b-dodecahydro-4,6,9,11-tetrahydroxy-4a,8,13,13-tetramethyl-7,11-methano-5*H*-cyclodeca[3,4]benz[1,2-*b*]oxet-5-one.
POTASSIUM	ASSAY	*USPNF Online*	Online	30-Aug-2024	1-Sep-2024	*NA*	*NA*	In *Procedure for*

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PHOSPHATES COMPOUNDED INJECTION								*Potassium*/*Chromatographic system*/*Column*: Change 4.6-mm × 25-cm; packing L97 to: 4-mm × 25-cm; packing L97 AND In *Procedure for Phosphate*/*Chromatographic system*/*Column*: Change 4.6-mm × 25-cm; packing L103 to: 4-mm × 25-cm; packing L103
NICARDIPINE HYDROCHLORIDE	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	26-Jul-2024	1-Aug-2024	*NA*	*NA*	In *USP Reference Standards* <*11*>/USP Nicardipine

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								Related Compound B RS: Change 3-{2-[Benzyl(methyl)amino]ethyl }5-methyl 2,6-dimethyl-4-(3-nitrophenyl)pyridine-3,5-dicarboxylate oxalate. to: 3-{2-[Benzyl(methyl)amino]ethyl } 5-methyl 2,6-dimethyl-4-(3-nitrophenyl)pyridine-3,5-dicarboxylate oxalate. AND In *USP Reference Standards* <*11*>/USP Nicardipine Related Compound D RS: Change Bis{2-[benzyl(methyl)amino]ethyl}2,6-dimethyl-4-(3-nitrophenyl

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								)-1,4-dihydropyridine-3,5-dicarboxylate dihydrochloride. to: Bis{2-[benzyl(methyl)amino]ethyl} 2,6-dimethyl-4-(3-nitrophenyl)-1,4-dihydropyridine-3,5-dicarboxylate dihydrochloride.
EXENATIDE INJECTION	PRODUCT-RELATED SUBSTANCES AND IMPURITIES	*USPNF Online*	Online	26-Jul-2024	1-Aug-2024	*NA*	*NA*	In *Procedure/Chromatographic system/Column*: Change 4.6-mm × 10-cm; 3-µm packing L52 to: 4.6-mm × 10-cm; 3-µm packing L52; two columns in series
CRANBERRY FRUIT JUICE	IDENTIFICATION	*USPNF Online*	Online	26-Jul-2024	1-Aug-2024	*NA*	*NA*	In *A. HPTLC for Articles of*

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DRY EXTRACT								*Botanical Origin <203>*/*Sample solution*: Change Mix 50 mg of Cranberry Fruit Juice Dry Extract with 7 mL of water. Sonicate for 10 min at room temperature, centrifuge, and save the supernatant. Precondition an SPE cartridge (L2; 500 mg/6 mL) with 3 mL of methanol, and drain it with suction. Load 3 mL of water onto the cartridge and apply suction until the solvent reaches 1–2 mm above the top of the cartridge's

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								stationary phase. Then load the cartridge with 1.5 mL of the supernatant, apply suction, and discard the eluate. Add 1 mL of a mixture of water and methanol (80:20). Load 1 mL of methanol and collect the eluate as the *Sample solution*. to: Mix 50 mg of Cranberry Fruit Juice Dry Extract with 7 mL of water. Sonicate for 10 min at room temperature, centrifuge, and save the supernatant. Precondition an

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								SPE cartridge (L2; 500 mg/6 mL) with 3 mL of methanol, and drain it with suction. Load 3 mL of water onto the cartridge and apply suction until the solvent reaches 1–2 mm above the top of the cartridge's stationary phase. Then load the cartridge with 1.5 mL of the supernatant, apply suction, and discard the eluate. Add 1 mL of a mixture of water and methanol (80:20), apply suction, and discard the eluate. Load 1

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								mL of methanol and collect the eluate as the *Sample solution*.
NAPROXEN SODIUM	SPECIFIC TESTS	*USPNF Online*	Online	28-Jun-2024	1-Aug-2024	*NA*	*NA*	In *Loss on Drying <731>/Analysis*: Change Dry at 105° for 3 h. to: Dry in vacuum at 105° for 3 h.
EDETATE DISODIUM COMPOUNDED OPHTHALMIC SOLUTION	DEFINITION	*USPNF Online*	Online	28-Jun-2024	1-Jul-2024	*NA*	*NA*	In two instances: Change to a pH between 6.5 and 7.5. to: to a pH between 6.1 and 7.1.
DESCRIPTION AND SOLUBILITY	Hydrogenated Castor Oil	*USPNF Online*	Online	28-Jun-2024	1-Aug-2026	*NA*	*NA*	Change White, crystalline wax. Freely soluble in water, in saline TS, and in dextrose

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								solutions; very slightly soluble in alcohol; practically insoluble in chloroform and in ether. Insoluble in water and in most common organic solvents. *NF category*: Stiffening agent; lubricant; film-forming agent; r elease-modifying agent. to: White, crystalline wax. Insoluble in water and in most common organic solvents. *NF category*: Stiffening agent; lubricant; film-forming agent; r

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								elease-modifying agent.
NEOMYCIN AND POLYMYXIN B SULFATES AND LIDOCAINE CREAM	ASSAY	*USPNF Online*	Online	28-Jun-2024	1-Jul-2024	*NA*	*NA*	Change **Assay for ne omyc in—**Proceed with Cream as directed in the *Assay for neomycin* under *Neomycin and Polymyxin B Sulfates Cream*. **Assay for polymyxin B—**Proceed with Cream as directed in the *Assay for polymyxin B* under *Neomycin and Polymyxin B Sulfates Cream*. to: **Assay for ne omyc**

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								in—Proceed as directed under *Antibiotics—Microbial Assays* ?81?, using an accurately weighed portion of Cream, equivalent to about 1.75 mg of neomycin, shaken in a separator with about 50 mL of ether, and extracted with four 20-mL portions of *Buffer B.3*. Combine the aqueous extracts, and dilute with *Buffer B.3* to an appropriate volume to obtain a stock solution of convenient concentration. Dilute this stock

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								solution quantitatively and stepwise with *Buffer B.3* to obtain a *Test Dilution* having a concentration assumed to be equal to the median dose level of the Standard. **Assay for polymyxin B—**Proceed as directed under *Antibiotics—Microbial Assays* ?81?, using an accurately weighed portion of Cream shaken with about 50 mL of ether in a separator, and extracted with four 25-mL portions of *Buffer B.6*. Combine the

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								aqueous extracts, and dilute with *Buffer B.6* to an appropriate volume to obtain a stock solution. Dilute this stock solution quantitatively and stepwise with *Buffer B.6* to obtain a *Test Dilution* having a concentration assumed to be equal to the median dose level of the Standard (10 Polymyxin B Units per mL). Add to each test dilution of the Standard a quantity of USP Neomycin Sulfate RS, dissolved in *Buffer B.6*, to

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								obtain the same concentration of neomycin present in the *Test Dilution*.
TYLOSIN INJECTION	SPECIFIC TESTS	*USPNF Online*	Online	31-May-2024	1-Jun-2024	*NA*	*NA*	Change *pH* ?921? to: *pH* ?791?
CEFUROXIME AXETIL FOR ORAL SUSPENSION	ASSAY	*USPNF Online*	Online	31-May-2024	1-Jun-2024	*NA*	*NA*	In *Procedure*/*System suitability*: Change [Note—The relative retention times for acetanilide, cefuroxime axetil diastereoisomer B, cefuroxime axetil diastereoisomer A, and cefuroxime axetil delta-3 isomers are 0.4, 0.8, 0.9, and 1.0, respectively.]

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								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.]
METRONIDAZOLE CAPSULES	IMPURITIES	*USPNF Online*	Online	31-May-2024	1-Jun-2024	*NA*	*NA*	In *Organic Impurities*: Change **Mobile phase**, **Sample solution**, and **Chromatographic system:** Proceed as directed in the *Assay*. to: **Mobile phase** and

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Chromatographic system: Proceed as directed in the *Assay*. AND Change **Standard solution:** 1 µg/mL of metronidazole from USP Metronidazole RS and 2 µg/mL of tinidazole related compound A from USP Tinidazole Related Compound A RS in *Mobile phase* to: **Standard solution:** 1 µg/mL of metronidazole from USP Metronidazole

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								RS and 2 µg/mL of tinidazole related compound A from USP Tinidazole Related Compound A RS in *Mobile phase* **Sample soluti on:** Nominally 1 mg/mL of metronidazole prepared as follows. Mix the contents of Capsules (NLT 20). Transfer an amount equivalent to 100 mg of metronidazole to a 100-mL volumetric flask, add 80 mL of *Mobile phase*, and sonicate with intermittent shaking for 10

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								min. Shake for 30 min, and dilute with *Mobile phase* to volume. Centrifuge a portion of the solution.
PROCAINAMIDE HYDROCHLORIDE	ASSAY	*USPNF Online*	Online	31-May-2024	1-Jun-2024	*NA*	*NA*	In *Resolution solution*: Change *p*-aminobenzoic acid to: *p*-aminobenzoic acid
DOCETAXEL	IMPURITIES	*USPNF Online*	Online	31-May-2024	1-Jun-2024	*NA*	*NA*	In *Organic Impurities*, *Procedure 1*: Change **System suitability solution, Standard solution, Sample solution,** and **Chromatographic system:** Proceed as directed in the

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Assay. to: **Mobile phase, Diluent, System suitability solution, Standard solution, Sample solution,** and **Chromatographic system:** Proceed as directed in the *Assay*.
RIVASTIGMINE TARTRATE	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	26-Apr-2024	1-May-2024	*NA*	*NA*	In *USP Reference Standards* ?11?: Change USP Rivastigmine Tartrate R-Isomer RS to: USP Rivastigmine Tartrate R-Isomer RS (*R*)-3-[1-(Dimethyl

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								amino)ethyl]phenyl ethylmethylcarbamate, hydrogen tartrate. $C_{14}H_{22}N_2O_2 \cdot C_4H_6O_6$ 400.42

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