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Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
IPRATROPIUM IMPURITIES		*USPNF Online*	Online	29-Mar-2024	1-Apr-2024	*NA*	*NA*	In *Organic*

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BROMIDE INHALATION SOLUTION								*Impurities*: Change **Column:** 4.6-mm × 25-µm; 5-µm packing L1 to: **Column:** 4.6-mm × 25-cm; 5-µm packing L1
SUCROSE	IMPURITIES	*USPNF Online*	Online	29-Mar-2024	1-Apr-2024	*NA*	*NA*	In *Sulfite/ Analysis*/footnote 1: Change Test kit for sulfite determination may be ordered from the following suppliers: Megazyme Ltd. (Product code: K-ETSULPH); R-Biopharm (Enzytec) (Article No.: E6275); BioSen Tec/Nzytech

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								(Catalogue No.: AK00071). to: Test kit for sulfite determination may be ordered from the following suppliers: Megazyme Ltd. (Product code K-ETSULPH); R-Biopharm (Enzytec) (Article No. E6275); Nzytech (Catalogue No. AK00071) and BioSenTec (Product reference 040-E).
METHYLNALTREXONE BROMIDE	CHEMICAL INFORMATION	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	Correct the chemical structure
IODIXANOL INJECTION	IMPURITIES	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In *Organic Impurities, Procedure 2*/footnote 4:

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								Change 5-[[3-[[3-[[[3-[[3-[[3,5-Bis-[[[2,3-dihydroxypropyl]amino]carbonyl]-2,4,6-triiodophenyl](acetylimino)]-2-hydroxypropyl](acetylimino)]-5-[[[2,3-dihydroxypropyl]amino]carbonyl]-2,4,6-triiodophenyl]carbonyl]amino]-2-hy droxypropyl]oxy]-2-hydroxypropyl](acetylimino)]-*N*,*N*'-bis(2,3-dihy droxypropyl)-2,4,6-triiodo-1,3-benzendicarboxamide. to: 5-[[3-[[3-[[[3-[[3-[[3,5-Bis-[[[2,3-dihydroxypropyl]amino]carbonyl]-2,4,6-triiodophenyl](acetylamino)]-2-hydroxypr

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								opyl](acetylamino)]-5-[[[2,3-dihydroxypropyl]amino]carbonyl]-2,4,6-triiodophenyl]carbonyl]amino]-2-hydroxypropyl]oxy]-2-hydroxypropyl](acetylamino)]-*N*,*N*'-bis(2,3-dihydroxypropyl)-2,4,6-triiodo-1,3-benzenedicarboxamide.
THALIDOMIDE CAPSULES	ASSAY	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In *Procedure*: Change $1000C(R_U/R_S)$ to: $500C(R_U/R_S)$
COCOYL CAPRYLOCAPRATE	SPECIFIC TESTS	*USPNF Online*	Online	23-Feb-2024	1-May-2024	*NA*	*NA*	In *Fats and Fixed Oils* ?401?, *Procedures*, *Hydroxyl Value*/*Analysis*: Change Calculate the acid value as directed in the chapter.

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								to: Calculate the hydroxyl value as directed in the chapter.
METOLAZONE	IMPURITIES	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In *Organic Impurities/ Chromatographic system/Column*: Change 4.6-mm × 25-cm; 5-µm packing 1 to: 4.6-mm × 25-cm; 5-µm packing L1
IODIXANOL INJECTION	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In *USP Reference Standards* ?11?/USP Iodixanol Related Compound ERS: Change 5-[[3-[[3-[[(2,3-Dihydroxypropyl)amino]carbonyl]-5- [[amino]carb

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								onyl]-2,4,6-triiodophenyl](acetylimino)]-2-hydroxypropyl]-(acetylimino)]-N,N?-bis(2,3-dihydroxypropyl)-2,4,6-triiodo-1,3-benzenedicarboxamide. to: 5-{*N*-[3-(*N*-{3-Carbamoyl-5-[(2,3-dihydroxypropyl)carbamoyl]-2,4,6-triiodophenyl}acetamido)-2-hydroxypropyl]acetamido}-N^1,N^3-bis(2,3-dihydroxypropyl)-2,4,6-triiodoisophthalamide.
FLURAZEPAM HYDROCHLORIDE	USP REFERENCE STANDARDS ?11?	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In USP Flurazepam Related Compound C RS: Change 5-Chloro-2-(2-diethylaminoethyl(amino)-2?-fluor

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								obenzophenone hydrochloride. to: 5-Chloro-2-(2-diethylaminoethylamino)-2?-fluorobenzophenone hydrochloride.
PROPOFOL	ASSAY	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In *Procedure 2*/*Mobile phase*: Change Hexane, acetonitrile, and alcohol (990: 7.5: 1) to: Hexane, acetonitrile, and alcohol, absolute (990: 7.5: 1)
DRY HEAT DEPYROGENATION	INTRODUCTION	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	Change parental manufacturing to: parenteral manufacturing
DICYCLOMINE HYDROCHLORIDE	IDENTIFICATION	*USPNF Online*	Online	26-Jan-2024	1-Feb-2024	*NA*	*NA*	In *B. Identification Tests—General* ?191?,

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AMOXICILLIN ORAL SUSPENSION	IDENTIFICATION	*USPNF Online*	Online	26-Jan-2024	1-Feb-2024	*NA*	*NA*	*Chemical Identification Tests*, *Chloride*: Change Meets the requirements when tested as specified in test *B*. to: Meets the requirements of the test for amine hydrochlorides 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

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								responds to the *Identification* test under *Amoxicillin Capsules*. 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 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

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 methanol, chloroform, water, and pyridine (90:80:30:10). When the solvent front has moved

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								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 corresponds to that obtained from the Standard solution.
LINEZOLID TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	26-Jan-2024	1-Feb-2024	*NA*	*NA*	In *Dissolution* ?711?/*Test 1*/*Analysis*:

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								Change Result = (r_U/r_S) × C_S × V × (1/L) × 100 r_U = peak response of linezolid from the *Sample solution* r_S = peak response of linezolid from the *Standard solution* C_S = concentration of USP Linezolid RS in the *Standard solution* (mg/mL) V = volume of *Medium*, 900 mL L = label claim (mg/Tablet) to: Result = (r_U/r_S) × C_S × V × (1/L) × D × 100 r

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CARVEDILOL TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	26-Jan-2024	1-Feb-2024	*NA*	*NA*	A_U = peak response of linezolid from the *Sample solution* r_S = peak response of linezolid from the *Standard solution* C_S = concentration of USP Linezolid RS in the *Standard solution* (mg/mL) *V* = volume of *Medium*, 900 mL *L* = label claim (mg/Tablet) *D* = dilution factor of the *Sample solution*, as needed In *Dissolution* <711>/*Test 3*/ *Chromatographi*

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								c system/*Column*: Change 4.6-mm × 15-mm; 5-?m packing L7 to: 4.6-mm × 15-cm; 5-?m packing L7
THEOPHYLLINE CAPSULES	IDENTIFICATION	*USPNF Online*	Online	26-Jan-2024	1-Feb-2024	*NA*	*NA*	Change **A:** The contents of the Capsules respond to *Identification* tests *A* and *B* under *Theophylline Tablets*. **B:** The retention time of the major peak in the chromatogram of the *Assay preparation* corresponds to that in the chromatogram of the *Standard*

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								preparation, as obtained in the *Assay*. to: **A:** Triturate a quantity of the contents of Capsules, equivalent to about 500 mg of theophylline, with 10-mL and 5-mL portions of solvent hexane, and discard the solvent hexane. Triturate the residue with two 10-mL portions of a mixture of equal volumes of 6 N ammonium hydroxide and water, and filter each time. Evaporate the combined filtrates to about 5 mL, neutralize, if

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								necessary, with 6 N acetic acid, using litmus, and then cool to about 15°, with stirring. Collect the precipitate on a filter, wash it with cold water, and dry at 105° for 2 hours: the theophylline so obtained melts between 270° and 274° (see *Melting Range or Temperature* ?741?, *Procedures*, *Procedure for Class I*). Retain the remaining portion of the theophylline for use in *Identification* test *B*. **B:** The IR absorption spectrum of a

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								potassium bromide dispersion of the residue obtained in *Identification* test *A* exhibits maxima only at the same wavenumbers as that of a potassium bromide dispersion of USP Theophylline RS. **C:** The retention time of the major peak in the chromatogram of the *Assay preparation* corresponds to that in the chromatogram of the *Standard preparation*, as obtained in the *Assay*.

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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*)
GLUCAGON	PROCESS-RELATED IMPURITIES AND OTHER COMPONENTS	*USPNF Online*	Online	29-Dec-2023	1-Jan-2024	*NA*	*NA*	In *Acetic Acid in Peptides/Analysis*: Change C_{SPA} = concentration of potassium acetate in each of the *Standard solutions* (mg/mL) to: C_{SPA} = concentration of potassium acetate in each of the *Standard solutions* (µg/mL) AND In

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Am monium/ Analysis: Change C_{SAC} = concentration of ammonium chloride in each of the *Standard solutions* (mg/mL) to: C_{SAC} = concentration of ammonium chloride in each of the *Standard solutions* (µg/mL)
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
SECOBARBITAL SODIUM	OTHER REQUIREMENTS	*USPNF Online*	Online	29-Dec-2023	1-Jan-2024	*NA*	*NA*	Change Where the label states that Seco barbital Sodium is sterile, it

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								meets the requirements for *Sterility Tests* ?71? and for *Bacterial endotoxins* under *Secobarbital Sodium for Injection*. Where the label states that Secobarbital Sodium must be subjected to further processing during the preparation of injectable dosage forms, it meets the requirements for *Bacterial endotoxins* under *Secobarbital Sodium for Injection*. to: Where the label

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								states that Secobarbital Sodium is sterile, it meets the requirements for *Sterility Tests* ?71? and the level of bacterial endotoxins is not more than 0.9 USP Endotoxin Units per mg of secobarbital sodium tested per *Bacterial Endotoxins Test* <85>. Where the label states that Secobarbital Sodium must be subjected to further processing during the preparation of injectable dosage forms,

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								the level of bacterial endotoxins is not more than 0.9 USP Endotoxin Units per mg of secobarbital sodium tested per *Bacterial Endotoxins Test* <85>.
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 solution of USP Amoxicillin RS in 0.1 N hydrochloric acid containing 4 mg per mL. Use within 10 minutes after

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 methanol, chloroform, water, and pyridine (90:80:30:10). When the solvent front has moved

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								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 corresponds to that obtained from the Standard solution.
NORFLURANE	IMPURITIES	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Organic Impurities*/*Table 2*: Change

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CEFDINIR	IMPURITIES	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	Line No. to: Peak Elution Order AND In *Halides*/*Figure 1*: Add label Flow Meter 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.
AMIKACIN SULFATE	ASSAY	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Procedure*/*Analysis*: Change C_U = concentration of amikacin in the

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								Sample solution (mg/mL) to: C_U = concentration of Amikacin Sulfate in the *Sample solution* (mg/mL)
DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE CAPSULES	PERFORMANCE TESTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Dissolution* ?711?/*Test for dutasteride*/*Tier 2*/*Medium*: Change 10 g/L of cetyltrimethylammonium bromide and 750,000 USP units of activity/mg of pepsin, purified in 0.1 N hydrochloric acid; 900 mL to: Dissolve 10 g of cetyltrimethylammonium bromide and 1.6 g of pepsin, purified in 1000

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								mL of 0.1 N hydrochloric acid; 900 mL
SODIUM SALICYLATE	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *USP Reference Standards* ?11?: Add USP Phenol RS
CEFDINIR FOR ORAL SUSPENSION	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *USP Reference Standards* ?11?/USP Cefdinir Related Compound A RS: Change 413.43 to: 413.42 AND In *USP Reference Standards* ?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

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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
AZITHROMYCIN	IMPURITIES	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Organic Impurities*/*Table 2*: Change 3'-*N*-Demethyl-3'-*N*-[(4-methylphenyl)sulfonyl]azithromycin[m] to: 3?-*N*-{[4-(Acetylamino)phenyl]sulfonyl}-3?-demethylazithromycin[m]

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								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*-(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*

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								,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.
OLMESARTAN MEDOXOMIL TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Dissolution* ?711?/*Test 5*/*Apparatus 2*: Change 50 rpm. Use peak vessels. to:

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								50 rpm. Use apex vessels.
CEFDINIR	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *USP Reference Standards* ?11?/USP Cefdinir Related Compound A RS: Change 413.43 to: 413.42
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 volume*—5 µL. *Developing solvent system*—a

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								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 minutes.
DUTASTERIDE AND TAMSULOSIN	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *USP Reference Standards*

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HYDROCHLORIDE CAPSULES								?11?/USP Dihydrodutasteride RS: Change *N*-[2,5-Bis(trifluoromethyl)phenyl]-3-oxo-4-aza-5?-androst-1-ene-17?-carboxamide. to: *N*-[2,5-Bis(trifluoromethyl)phenyl]-3-oxo-4-aza-5?-androstane-17?-carboxamide.
Strychnine Sulfate	REAGENTS AND REFERENCE TABLES	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Reagent Specifications*: Change CAS RN®: 60-41-3. to: CAS RN®: 60491-10-3.
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 DICLOFENAC SODIUM AND MISOPROSTOL DELAYED-RELEASE TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Dissolution* ?711?: Move *Test 2* after *Test 1*
PANTOPRAZOLE SODIUM DELAYED-RELEASE TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Dissolution* ?711?/*Test 2*/*Acid stage*: Change **Acid stage standard solution:** (*L*/10) mg/mL of USP Pantoprazole Sodium RS from the *Standard stock solution* in *Acid stage medium*, where *L* is the label claim in mg/Tablet to: **Acid stage standard solution:** (*L*/10000) mg/mL of USP

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								Pantoprazole Sodium RS from the *Standard stock solution* in *Acid stage medium*, where *L* is the label claim in mg/Tablet AND In *Dissolution* <711>/*Test 2*/*Buffer stage*: Change **Buffer stage standard solution:** (*L*/1000) of USP Pantoprazole Sodium RS where *L* is the label claim in mg/Tablet to: **Buffer stage standard solution:** (*L*/1000) mg/mL of USP Pantoprazole

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								Sodium RS from the *Standard stock solution* in *Buffer stage medium*, where *L* is the label claim in mg/Tablet
CEFDINIR CAPSULES	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *USP Reference Standards* ?11?/USP Cefdinir Related Compound A RS: Change 413.43 to: 413.42 AND In *USP Reference Standards* ?11?/USP Cefdinir Related Compound B RS: Change $C_{14}H_{13}N_4O_4S_2$ 365.41 to: C

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								$^{14}H_{14}N_4O_4S_2$ 366.41
TELMISARTAN TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Dissolution* ?711?/*Test 1*/*Analysis*: Change Determine the percentage of telmisartan ($C_{33}H_{30}N_4O_2$) dissolved: Result = (A_U × C_S × V × 100)/(A_S × D × L) to: Calculate the percentage of the labeled amount of telmisartan ($C_{33}H_{30}N_4O_2$) dissolved: Result = (A_U/A_S) × C_S × V × D × (1/L) × 100 AND Change C_S = concentration of the *Standard*

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								solution (mg/mL) to: C_S = concentration of USP Telmisartan RS in the *Standard solution* (mg/mL) AND In *Dissolution* ?711?/*Test 2*/*Analysis*: Change r_U = peak response from the *Sample solution* r_S = peak response from the *Standard solution* to: r_U = peak response of telmisartan from the *Sample solution* r_S = peak response of

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								telmisartan from the *Standard solution* AND In *Dissolution* ?711?/*Test 3*/*Analysis*: Change C_S = concentration of the *Standard solution* (mg/mL) to: C_S = concentration of USP Telmisartan RS in the *Standard solution* (mg/mL)
METHYLBENZETHONIUM CHLORIDE	ASSAY	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Procedure*/*Analysis*: Change Calculate the percentage of methylbenzethonium chloride ($C_{28}H_{44}ClNO_2 \cdot H$

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AMOXICILLIN FOR INJECTABLE SUSPENSION	Identification	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	$_2$O) in the portion of Methylbenzethonium Chloride taken: to: Calculate the percentage of methylbenzethonium chloride ($C_{28}H_{44}ClNO_2$) in the portion of Methylbenzethonium Chloride taken: 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

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								minutes before use: the solution responds to the *Identification* test under *Amoxicillin Capsules*. 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. Allow the solution to stand for 5 minutes before use. Prepare a Standard solution of USP Amoxicillin RS in 0.1 N

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								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 methanol, chloroform, water, and

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								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 corresponds to that obtained from the

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								Standard solution.
OIL- AND WATER-SOLUBLE VITAMINS WITH MINERALS CAPSULES	STRENGTH	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Vitamin A, Method 3*/*Analysis*: Change C_S = concentration of retinyl acetate ($C_{23}H_{32}O_2$) from USP Vitamin A RS in the *Standard solution* (?g/mL) to: C_S = concentration of retinyl acetate ($C_{22}H_{32}O_2$) from USP Vitamin A RS in the *Standard solution* (?g/mL)
UREA	SPECIFIC TESTS	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Alcohol-Insoluble Matter*/*Sample solution*: Change 100 mg/mL of Urea dissolved in warm alcohol

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								to: Dissolve 5.0 g of Urea in 50 mL of warm alcohol. AND In *Alcohol-Insoluble Matter/Analysis*: Change If any insoluble residue remains, pass the *Sample solution* through a tared filter, wash the residue and the filter with 20 mL of warm alcohol per 50 mL of *Sample solution*, and dry at 105° for 1 h. to: If any insoluble residue remains, pass the *Sample solution* through

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TELMISARTAN ASSAY TABLETS		*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	a tared filter, wash the residue and the filter with 20 mL of warm alcohol, and dry at 105° for 1 h. In *Procedure*/*System suitability*/*Suitability requirements*: Change **Resolution:** NLT 3 between telmisartan and telmisartan related compound A **Tailing factor:** NMT 2.0 for the telmisartan peak **Capacity factor:** NLT 1.5 **Relative standard deviation:** NMT 2.0%

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								to: **Resolution:** NLT 3 between telmisartan and telmisartan related compound A **Tailing factor:** NMT 2.0 for telmisartan **Relative standard deviation:** NMT 2.0% for telmisartan
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).

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								to: in pascal-seconds (Pa?s).
OIL-SOLUBLE VITAMINS CAPSULES	STRENGTH	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Vitamin A, Method 3*/*Analysis*: Change C_S = concentration of retinyl acetate ($C_{23}H_{32}O_2$) from USP Vitamin A RS in the *Standard solution* (?g/mL) to: C_S = concentration of retinyl acetate ($C_{22}H_{32}O_2$) from USP Vitamin A RS in the *Standard solution* (?g/mL)
PANTOPRAZOLE SODIUM DELAYED-RELEASE TABLETS	ASSAY	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Procedure*/*System suitability*/*Tailing factor*: Change NMT 2.0,

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								System suitability solution to: NMT 2.0 for pantoprazole, *System suitability solution*
TAMSULOSIN HYDROCHLORIDE CAPSULES	IMPURITIES	*USPNF Online*	Online	29-Sep-2023	1-Oct-2023	*NA*	*NA*	In *Organic Impurities*/*System suitability*: Change **Sample:** *Standard solution* [Note—The relative retention times for methoxy tamsulosin, tamsulosin, ethoxyphenoxy ethyl bromide, and desethoxy tamsulosin are 0.73, 1.00, 1.80, and 2.80, respectively.] **Suitability**

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								requirements **Tailing factor:** NMT 2.0 **Relative standard deviation:** NMT 5.0% **Signal-to-noise ratio:** NLT 10 to: **Samples:** *Standard solution* and *Sensitivity solution* [Note—The relative retention times for methoxy tamsulosin, tamsulosin, ethoxyphenoxy ethyl bromide, and desethoxy tamsulosin are 0.73, 1.00, 1.80, and 2.80, respectively.] **Suitability requirements**

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Tailing factor: NMT 2.0, *Standard solution* **Relative standard deviation:** NMT 5.0%, *Standard solution* **Signal-to-noise ratio:** NLT 10, *Sensitivity solution*
DIGOXIN TABLETS	ASSAY	*USPNF Online*	Online	29-Sep-2023	1-Oct-2023	*NA*	*NA*	In *Procedure/Analysis*: Change C_U = nominal concentration of in the *Sample solution* (µg/mL) to: C_U = nominal concentration of digoxin in the *Sample solution* (µg/mL)
DIVALPROEX SODIUM EXTENDED-	PERFORMANCE TESTS	*USPNF Online*	Online	29-Sep-2023	1-Oct-2023	*NA*	*NA*	In *Dissolution* ?711?/*Test 8*/*Tolerances*:

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RELEASE TABLETS								Change The percentage of the labeled amount of valproic acid ($C_8H_{16}O_2$) dissolved at the times specified conform to *Dissolution* <711>, *Acceptance Table 1*. to: The percentage of the labeled amount of valproic acid ($C_8H_{16}O_2$) dissolved at the times specified conform to *Dissolution* <711>, *Acceptance Table 2*.

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