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CHROMATOGRAPHY	ADJUSTMENT OF CHROMATOGRAPHIC CONDITIONS	*USPNF Online*	Online	26-Aug-2022	1-Dec-2022	*NA*	*NA*	In *Liquid Chromatography: Isocratic Elution/Injection volume*: Change L_2 = internal diameter of the column used (mm) dc_1 = particle size indicated in the monograph (µm) dc_2 = particle size of the column used (µm) to: L_2 = new column length (mm) dc_1 = column internal diameter indicated in the monograph (mm) dc_2 = new column internal

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								diameter (mm)
METHYLENE BLUE	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USPNF Online*	Online	26-Aug-2022	1-Sep-2022	*NA*	*USPNF 2023 Issue 2*	In USP Azure B RS: Change 3-(Dimethylamino)-7-(methylamino)-phenothiazine-5-ium chloride. to: 3-(Dimethylamino)-7-(methylamino)phenothiazin-5-ium chloride.
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*

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Test Method, Spectral Transmission Requirements for Light-Resistant Components and Systems.
DOXORUBICIN HYDROCHLORIDE INJECTION	IMPURITIES/*Organic Impurities*	*USPNF 2022 Online*	Online	28-Oct-2022	1-Nov-2022	*NA*	*NA*	In *Analysis*: Change *P* = potency of doxorubicin in USP Doxorubicin Hydrochloride RS (µg/mg) to: *P* = potency of doxorubicin hydrochloride in USP Doxorubicin Hydrochloride RS (µg/mg)
ESCITALOPRAM TABLETS	PERFORMANCE TESTS/ *Dissolution ?711?*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	In *Test 1*/*Medium*: Change 0.1 N hydrochloric acid VS; 900

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								mL to: 0.1 N hydrochloric acid; 900 mL AND In *Test 2*/*Medium*: Change 0.1 N hydrochloric acid VS; 900 mL to: 0.1 N hydrochloric acid; 900 mL
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-Dideoxy-3-*C*-methyl-3-*O*-methyl-?-L-*ribo*-hexopyranosyl)oxy]-2-ethyl-3,4,

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								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*-hexopyranosyl)oxy]-2-ethyl-3,4,10-trihydroxy-3,5,6,8,10,12,14-heptamethyl-11-

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OIL-SOLUBLE VITAMINS CAPSULES	STRENGTH	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	[[3-[*N*-(4-methylphenylsulfonyl)-*N*-methylamino]-3,4,6-trideoxy-ß-D-*xylo*-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one. 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

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								Standard solution (?g/mL)
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?-fluorobenzophenone hydrochloride. to: 5-Chloro-2-(2-diethylaminoethylamino)-2?-fluorobenzophenone hydrochloride.
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE TABLETS	ASSAY/ *Procedure*	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	In *Buffer*: Change Adjust with phosphoric acid to a pH of 6.4 ± 0.01. to: Adjust with phosphoric acid to a pH of 6.4 ± 0.1.
SODIUM	IM	*USPNF Online*	Online	30-Sep-2022	1-Oct-2022	*NA*	*NA*	In *Buffer*:

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
PICOSULFATE	PURITIES/*Organic Impurities*							Change cetylrimethylammonium bromide, to: cetyltrimethylammonium bromide,
MOMETASONE FUROATE TOPICAL SOLUTION	IDENTIFICATION	*USPNF Online*	Online	18-Nov-2022	1-Dec-2022	*NA*	*NA*	In test *A*: Change The retention time of the major peak of the *Sample solution* corresponds to that of the *Standard solution*, both relative to the internal standard, as obtained in the *Assay*. to: The retention time of the major peak of the *Sample solution*

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								corresponds to that of the *Standard solution*, as obtained in the *Assay*.
MIRTAZAPINE TABLETS	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	24-Feb-2023	1-Mar-2023	*NA*	*NA*	In *System suitability/Suitability requirements/Resolution*: Change 10-ketomirtazpine to: 10-ketomirtazapine AND In *Table 1*: Change 1-Ketomirtazpine[c] to: 1-Ketomirtazapine[c]
VIGABATRIN TABLETS	IMPURITIES/*Organic*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	In *Analysis*: Change

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
	Impurities							C_S= concentration of USP Vigabatrin Related Compound A in the *Standard solution* to: C_S = concentration of USP Vigabatrin Related Compound A RS in the *Standard solution*
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

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

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								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 mg of amoxicillin per mL by adding 0.1 N hydrochloric acid to Amoxicillin for Injectable Suspension. Allow the solution to stand for 5

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

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

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								minutes: the R_F value of the principal spot obtained from the test solution corresponds to that obtained from the Standard solution.
IODIXANOL INJECTION	IMPURITIES	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In *Organic Impurities, Procedure 2*/footnote 4: 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-hydrox

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								ypropyl](acetyli mino)]-*N*,*N* '-bis(2,3-dihy dr oxypropyl)-2,4,6 -triiodo-1,3-ben zendicarboxami de. to: 5-[[3-[[3-[[[3-[[3 -[[3,5-Bis-[[[2,3-dihydroxypropyl]amino]carbonyl]-2,4,6-triiodoph enyl](acetylamin o)]-2-hydroxypr opyl](acetylamin o)]-5-[[[2,3-dihy droxypropyl]ami no]carbonyl]-2,4 ,6-triiodophenyl] carbonyl]amino] -2-hydroxypropy l]oxy]-2-hydroxy propyl](acetyla mino)]-*N*,*N* '-bis(2,3-dihydro xypropyl)-2,4,6-t riiodo-1,3-benze ndicarboxamide .

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HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE TABLETS	IMPURITIES/*Limit of Homatropine Hydrobromide and Related Substances*	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	In *Buffer*: Change Adjust with phosphoric acid to a pH of 6.4 ± 0.01. to: Adjust with phosphoric acid to a pH of 6.4 ± 0.1.
TERAZOSIN CAPSULES	ASSAY/*Procedure*	*USPNF Online*	Online	30-Sep-2022	1-Oct-2022	*NA*	*NA*	In *Hydrochloric acid solution*: Change 0.1 N methanolic hydrochloric acid to: 0.01 N methanolic hydrochloric acid
ROCURONIUM BROMIDE	CHEMICAL INFORMATION	*USPNF Online*	Online	18-Nov-2022	1-Dec-2022	*NA*	*USPNF 2023 Issue 3*	Change 609.68 to: 609.69
LIDOCAINE HYDROCHLORIDE AND	*Assay for lidocaine hydrochloride*	*USPNF Online*	Online	24-Feb-2023	1-Mar-2023	*NA*	*NA*	Change Proceed with Injection as

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DEXTROSE INJECTION								directed in the *Assay for lidocaine hydrochloride* under *Lidocaine and Epinephrine Injection*. to: Proceed with Injection as directed in the *Assay for lidocaine hydrochloride* under *Lidocaine Hydrochloride and Epinephrine Injection.*
FLUVOXAMINE MALEATE TABLETS	ASSAY/ *Procedure*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	In *Solution A*: Change 8 g/L of 1-pentanesulfonic acid sodium salt and 1 g/L of monobasic potassium phosphate in water.

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								to: 8 g/L of 1-pentanesulfonic acid sodium salt and 1.1 g/L of monobasic potassium phosphate in water.
REAGENTS AND REFERENCE TABLES	*Solutions*/*Test Solutions and Indicator Solutions*/*Starch TS*	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change Prepare this solution by one of the following procedures: to: Prepare this solution by one of the following procedures. Apply the *Test for sensitivity* to confirm suitability for freshly or previously prepared solutions or commercially bought solutions.

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								AND In *Storage*: Delete Use the *Test for sensitivity* to confirm suitability for use. AND In *Procedure with No Preservative*: Delete Apply the *Test for sensitivity* to confirm suitability for freshly or previously prepared solutions. AND In *Test for sensitivity*: Delete Use the *Test for sensitivity* to confirm suitability for use.

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TELMISARTAN ASSAY TABLETS		*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	AND In *Procedure with Salicylic Acid as Preservative*: Change Mix 1 g of soluble starch with 50 mL of cold water to: Mix 1 g of soluble starch with 5 mL of cold water In *Procedure/System suitability/Suitability requirements*: Change **Resolution:** NLT 3 between telmisartan and telmisartan related compound A **Tailing factor:**

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								NMT 2.0 for the telmisartan peak **Capacity factor:** NLT 1.5 **Relative standard deviation:** NMT 2.0% 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
IODIXANOL INJECTION	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In *USP Reference Standards* ?11?/USP Iodixanol

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								Related Compound ERS: Change 5-[[3-[[3-[[(2,3-Dihydroxypropyl)amino]carbonyl]-5- [[amino]carbonyl]-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.

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AMMONIUM GLYCYRRHIZATE	CHEMICAL INFORMATION	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	Change 840.08 to: 839.97
N-Benzoyl-L-arginine Ethyl Ester Hydrochloride	REAGENT SPECIFICATIONS	*USPNF Online*	Online	30-Sep-2022	1-Dec-2022	*NA*	*NA*	Change Crystallized Trypsin (USP Monograph). to: Trypsin (USP Monograph).
ROCURONIUM BROMIDE	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	18-Nov-2022	1-Dec-2022	*NA*	*USPNF 2023 Issue 3*	In footnote b of *Table 1*: Change 2?-(Morpholin-4-yl)-16?-(pyrrolidin-1-yl)-5?-androstan-3?,17?-diol. to: 2?-(Morpholin-4-yl)-16?-(pyrrolidin-1-yl)-5?-androstane-3?,17?-diol.
GLYCYL-L-TYROSINE	IMPURITIES/*Related Compounds*	*USPNF Online*	Online	24-Feb-2023	1-Jun-2023	*NA*	*NA*	In *Buffer solution*: Change Dissolve 6.84 g of potassium

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								phosphate in 1000 mL of water. to: Dissolve 6.84 g of monobasic potassium phosphate in 1000 mL of water.
PSEUDOEPHEDRINE HYDROCHLORIDE TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	Change **Uniformity of Dosage Units ?905? Procedure for content uniformity** to: **Uniformity of Dosage Units ?905?: Meets the requirements Procedure for content uniformity, chewable tablets only** AND

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UREA C 13	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Uniformity of Dosage Units*: Delete **Acceptance criteria**: Meet the requirements for chewable tablets
TELMISARTAN TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Isotopic Purity/ Chromatographic system*: Change **Flow rate**: Flow rate to: **Flow rate**: 1 mL/min In *Dissolution* ?711?/*Test 1*/*Analysis*: Change Determine the percentage of telmisartan ($C_{33}H_{30}N_4O_2$) dissolved: Result = (A_U ×

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								$C_S \times V \times 100)/(A_S \times D \times L)$ to: Calculate the percentage of the labeled amount of telmisartan ($C_{33}H_{30}N_4O_2$) dissolved: Result = $(A_U/A_S) \times C_S \times V \times D \times (1/L) \times 100$ AND Change C_S = concentration of the *Standard solution* (mg/mL) to: C_S = concentration of USP Telmisartan RS in the *Standard solution* (mg/mL) AND

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								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 telmisartan from the *Standard solution* AND In *Dissolution* ?711?/*Test 3*/*Analysis*: Change C_S = concentration of

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								the *Standard solution* (mg/mL) to: C_S = concentration of USP Telmisartan RS in the *Standard solution* (mg/mL)
AMMONIUM GLYCYRRHIZATE	ASSAY/*Content of Ammonium 18?- and 18?-Glycyrrhizate*	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	In *Analysis*: Change $M_{W(Salt)}$ = molecular weight of ammonium glycyrrhizate, 840.08 g/mol $M_{W(Acid)}$ = molecular weight of glycyrrhizic acid, 821.59 g/mol to: $M_{W(Salt)}$ = molecular weight of ammonium

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								glycyrrhizate, 839.97 g/mol $M_{W(Acid)}$ = molecular weight of glycyrrhizic acid, 822.94 g/mol
LANSOPRAZOLE DELAYED-RELEASE CAPSULES	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	30-Sep-2022	1-Oct-2022	*NA*	*NA*	Delete **Blank**: Methanol and *Diluent* (1:9)
MAFENIDE ACETATE FOR TOPICAL SOLUTION	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	16-Dec-2022	1-Jan-2023	*NA*	*NA*	In *System suitability*: Change **Samples**: *System suitability solution*, *Standard solution A*, and *Standard solution B* to: **Samples**: *System suitability solution* and *Standard*

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								solution A AND In *Analysis*: Change **Samples**: *Mobile phase*, *Standard solution A*, and *Sample solution* Chromatograph the *Standard solution* and adjust the integration parameters so that the response is 5%–15% of full-scale deflection. to: **Samples**: *Mobile phase*, *Standard solution A*, *Standard solution B*, and *Sample solution* Chromatograph *Standard*

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								solution B and adjust the integration parameters so that the response is 5%–15% of full-scale deflection.
METHYLNALTREXONE BROMIDE	CHEMICAL INFORMATION	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	Correct the chemical structure
ERYTHROMYCIN DELAYED-RELEASE TABLETS	PERFORMANCE TESTS/ *Dissolution* ?711?	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	In *Test 1*: Change **Buffer stage Medium**: 0.05 M pH 6.8 phosphate buffer (see *Reagents, Indicators, and Solutions—Buffer Solutions*) to: **Buffer stage Medium**: 0.05 M pH 6.8 phosphate buffer (see *Reagents,*

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								Indicators, and Solutions—Buffer Solutions); 900 mL
OSELTAMIVIR PHOSPHATE	IMPURITIES/*Organic Impurities*/*Procedure 3: Limit Of Tributyl Phosphine Oxide*	*USPNF Online*	Online	26-May-2023	1-Jun-2023	*NA*	*NA*	In *System suitability*/*Suitability requirements*/*Relative standard deviation*: Change NMT 10.0% for the tributyl phosphine oxide and osteltamivir phosphate peaks to: NMT 10.0% for the tributyl phosphine oxide and oseltamivir phosphate peaks
DEXTROMETHORPHAN HYD	*Assay*	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Chromatographi*

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ROBROMIDE ORAL SOLUTION								*c system* and *Procedure*: Change *C* is the concentration, in mg per mL, of USP Dextromethorphan Hydrobromide RS, on the anhydrous basis, in the *Standard preparation*; to: *C* is the concentration, in mg per mL, of USP Dextromethorphan Hydrobromide RS in the *Standard preparation*;
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
AMMONIUM GLYCYRRHIZATE	SPECIFIC TESTS	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	In *Optical Rotation, Specific Rotation ?781?*: Change ?781? to: ?781S? AND Change **Acceptance criteria**: +49.0 to +55.0 on the anhydrous basis to: **Acceptance criteria**: +49.0° to +55.0° on the anhydrous

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								basis
CEFTIOFUR HYDROCHLORIDE	ASSAY/ *Procedure*	*USPNF Online*	Online	30-Sep-2022	1-Oct-2022	*NA*	*NA*	In *Analysis*: Change Calculate the percentage of ceftiofur ($C_{19}H_{17}N_5O_7S_3$) in the portion of Ceftiofur Hydrochloride taken: Result = (r_U/r_S) × (C_S/C_U) × P × 100 to: Calculate the quantity, in µg/mg, of ceftiofur ($C_{19}H_{17}N_5O_7S_3$) in the portion of Ceftiofur Hydrochloride taken: Result = (r_U/r_S) × (C_S/C_U) × P
PALONOSETRON HYDROCHLORIDE	IMPURITIES	*USPNF Online*	Online	16-Dec-2022	1-Jan-2023	*NA*	*NA*	In *Limit of Unspecified Impurities*: Change

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
METOLAZONE IMPURITIES		*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	**Mobile phase** and **Chromatographic system**: Proceed as directed in the *Assay*. to: **Mobile phase**, **Standard stock solution**, and **Chromatographic system**: Proceed as directed in the *Assay*. In *Organic Impurities/ Chromatographic system/Column*: Change 4.6-mm × 25-cm; 5-µm packing 1 to: 4.6-mm × 25-cm; 5-µm

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ERYTHROMYCIN ETHYLSUCCINATE FOR ORAL SUSPENSION	*Identification*	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	packing L1 Change Proceed as directed in the *Identification* test under *Erythromycin Ethylsuccinate Oral Suspension*, beginning with “Prepare a Standard solution.” to: Prepare a Standard solution of USP Erythromycin Ethylsuccinate RS in methanol containing about 3 mg per mL. Apply separately 10 µL each of the test solution and the Standard solution to a

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								suitable thin-layer chromatographic plate (see *Chromatography* ?621?) coated with a 0.25-mm layer of chromatographic silica gel mixture, and allow to dry. Place the plate in an unlined chromatographic chamber, and develop the chromatograms in a solvent system consisting of a mixture of methanol and chloroform (85:15) until the solvent front has moved about 9 cm. Remove the plate from the chamber, mark

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								the solvent front, and allow the solvent to evaporate. Spray the plate with a mixture of dehydrated alcohol, *p* -methoxybenzaldehyde, and sulfuric acid (90:5:5). Heat the plate at 100° for 10 minutes, and examine the chromatograms, in which the erythromycin and succinic acid moieties appear as black-to-purple spots: the R_F values of the principal spots obtained from the test solution correspond to those obtained

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								from the Standard solution.
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

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Content of Calcium.
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 **Procedure for 197S**
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
MAGNESIUM OXIDE	IMPURITIES/*Limit of Calcium*	*USPNF Online*	Online	25-Feb-2022	1-Dec-2022	*NA*	*NA*	In *Analysis*: Change *C*

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NONOXYNOL 9	CHEMICAL INFORMATION	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	$_U$= concentration of Magnesium Hydroxide in the *Sample solution* (mg/mL) to: C_U = concentration of Magnesium Oxide in the *Sample solution* (mg/mL) Update the chemical structure AND Change ?-(*p* -Nonylphenyl)-?-hydroxynona(oxyethylene) CAS RN®: 26027-38-3. to: ?-(4-Nonylphenyl)-?-hydroxynona(oxyethylene).
CALCIUM L-5-	IDENTIFICATIO	*USPNF Online*	Online	28-Oct-2022	1-Nov-2022	*NA*	*NA*	In *A.*

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METHYLTETRAHYDROFOLATE								*Spectroscopic Identification Tests <197>, Infrared Spectroscopy: 197K*: Change USP Calcium D L-5-Methyltetrahydrofolate RS to: USP Calcium D, L-5-Methyltetrahydrofolate RS

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