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Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
DIVALPROEX SODIUM EXTENDED-RELEASE TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	29-Sep-2023	1-Oct-2023	*NA*	*NA*	In *Dissolution* ?711?/*Test 8*/*Tolerances*: 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*.
PROMETHAZI	IM	*USPNF Online*	Online	26-Aug-2022	1-Sep-2022	*NA*	*NA*	Change

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
NE HYDROCHLORIDE	PURITIES/*Organic Impurities*							**System suitability solution**: 5 µg/mL each of USP Promethazine Hydrochloride RS and USP Promethazine Related Compound B RS from the *Standard stock solution* and *System suitability stock solution*, respectively **Standard solution**: 5 µg/mL of USP Promethazine Hydrochloride RS from the *Standard stock solution* **Sensitivity solution**: 0.25 µg/mL of USP

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								Promethazine Hydrochloride RS from the *Standard solution* to: **System suitability solution**: 5 µg/mL each of USP Promethazine Hydrochloride RS and USP Promethazine Related Compound B RS from the *Standard stock solution* and *System suitability stock solution*, respectively, in *Diluent* **Standard solution**: 5 µg/mL of USP Promethazine

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								Hydrochloride RS from the *Standard stock solution* in *Diluent* **Sensitivity solution**: 0.25 µg/mL of USP Promethazine Hydrochloride RS from the *Standard solution* in *Diluent*
METARAMINOL BITARTRATE	SPECIFIC TESTS/*Content of Tartaric Acid*	*USPNF Online*	Online	27-Jan-2023	1-Feb-2023	*NA*	*NA*	In *Analysis*: Change Result = {[(V_S?V_B) × *N* × *F*]/2 × *W*} × 100 to: Result = {[(V_S?V_B) × *N* × *F*]/(2 × *W*)} × 100
REAGENTS AND REFERENCE TABLES	*Reagent S pec ificatio ns/ Tosylchloramid*	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	Change 127-65-1 to: 7080-50-4

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SECOBARBITAL SODIUM	*e Sodium* OTHER REQUIREMENTS	*USPNF Online*	Online	29-Dec-2023	1-Jan-2024	*NA*	*NA*	Change Where the label states that Secobarbital Sodium is sterile, it 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*

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								endotoxins under *Secobarbital Sodium for Injection*. to: Where the label 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

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								Sodium must be subjected to further processing during the preparation of injectable dosage forms, 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>.
DOXYCYCLINE FOR INJECTION	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	In USP Doxycycline Related Compound A RS: Change 444.43 to: 444.44 AND Change

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								(4*S*,4a*R*,5*S*,5a*R*,6*S*,12a*S*)-4-(Dimethylamino)-1,4,4a,5,5a,6,11,12a-octahydro-3,5,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-2-naphthacenecarboxamide, monohydrochloride. $C_{22}H_{24}N_2O_8$ · HCl 480.13 to: (4*S*,4a*R*,5*S*,5a*R*,6*S*,12a*S*)-4-(Dimethylamino)-1,4,4a,5,5a,6,11,12a-octahydro-3,5,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-2-naphthacenecarboxamide hydrochloride. C

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
NALOXONE HYDROCHLORIDE	IMPURITIES	*USPNF Online*	Online	28-Oct-2022	1-Dec-2022	*NA*	*NA*	$C_{22}H_{24}N_2O_8 \cdot HCl$ 480.90 In *Limit of Naloxone Related Compound D*: Delete **Sensitivity solution**: 1.25 µg/mL of USP Naloxone Related Compound D RS in 0.1 N hydrochloric acid AND In *System suitability*/*Samples*: Delete *Sensitivity solution*, AND In *System suitability*/*Suitability requirements*: Delete **Signal-to-noise**

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								ratio: NLT 10, *Sensitivity solution*
ISOFLURANE	IM PUR ITIES/*Organic Impurities*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	In *Analysis*: Change Result = $(r_U/r_S) \times C_S \times (1/F)$ to: Result = $(r_U/r_S) \times C_F \times (1/F)$ AND Change C_S = final concentration of USP Isoflurane Related Compound B RS in the *Standard solution* (%) to: C_F = final concentration of USP Isoflurane Related Compound B RS in the *Standard solution* (%)
AMMONIUM G	ASSAY/*Content*	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	In *Analysis*:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
LYCYRRHIZATE	*of Ammonium 18?- and 18?-Glycyrrhizate*							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 glycyrrhizate, 839.97 g/mol $M_{W(Acid)}$ = molecular weight of glycyrrhizic acid, 822.94 g/mol
MAFENIDE ACETATE FOR TOPICAL SOLUTION	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	16-Dec-2022	1-Jan-2023	*NA*	*NA*	In *System suitability*: Change **Samples**:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								System suitability solution, *Standard solution A*, and *Standard solution B* to: **Samples**: *System suitability solution* and *Standard 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

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								response is 5%–15% of full-scale deflection. to: **Samples**: *Mobile phase*, *Standard solution A*, *Standard solution B*, and *Sample solution* Chromatograph *Standard solution B* and adjust the integration parameters so that the response is 5%–15% of full-scale deflection.
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

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								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
COLOR AND ACHROMICITY	METHOD II: IN STRUMENTAL (QUANTITATIV E) 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

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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)
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, Indicators, and Solutions—Buffer Solutions*); 900 mL
DEXTROMETHORPHAN HYDROBROMIDE	*Assay*	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Chromatographic system* and

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
	ORAL SOLUTION							*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*;
METOLAZONE	IMPURITIES	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In *Organic Impurities/ Chromatographic system/Column*: Change

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								4.6-mm × 25-cm; 5-µm packing 1 to: 4.6-mm × 25-cm; 5-µm packing L1
POWDERED FORSKOHLII EXTRACT	COMPOSITION /*Content of Forskolin*	*USPNF Online*	Online	29-Jul-2022	1-Aug-2022	*NA*	*NA*	In *Chromatographic system*: Change Column: 4.6-mm × 25-cm; 5-µm, 100 Å to: Column: 4.6-mm × 25-cm; 5-µm, 100 Å; packing L1
MESO-ZEAXANTHIN	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USPNF Online*	Online	27-Jan-2023	1-Jun-2023	*NA*	*NA*	In USP *meso* -Zeaxanthin RS: Change (3*R*,3?*S*-*meso*)-Zeaxanthin. to: (3*R*,3?*S*

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								)-Zeaxanthin.
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 calcium, 1155.36
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**

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								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 Pantoprazole Sodium RS from the *Standard stock solution* in *Acid stage medium*, where *L* is the label claim in mg/Tablet AND In *Dissolution*

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BOTANICAL	PREPARATION	*USPNF Online*	Online	27-May-2022	1-Jun-2022	*NA*	*NA*	<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 Sodium RS from the *Standard stock solution* in *Buffer stage medium*, where *L* is the label claim in mg/Tablet In *General*

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EXTRACTS	S							*Pharmacopeial Requirements/Pesticide Residues*: Change where *L* is the limit in the original article as listed in *Table 4* (see *Pesticide Residue Analysis* under *Articles of Botanical Origin <561>*) to: where *L* is the limit in the original article as listed in *Table 5* (see *Pesticide Residue Analysis* under *Articles of Botanical Origin <561>*)

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DOXORUBICIN HYDROCHLORIDE FOR INJECTION	IMPURITIES/*Organic Impurities*	*USPNF 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)
LEVORPHANOL TARTRATE TABLETS	PERFORMANCE TESTS/ *Uniformity of Dosage Units* ?905?	*USPNF Online*	Online	31-Mar-2023	1-Aug-2023	*NA*	*NA*	In *Sample solution*: Change Nominally about 80 ug/mL of levorphanol tartrate in *Diluent* prepared as follows. to: Nominally about 80 µg/mL of levorphanol

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								tartrate in *Diluent* prepared as follows.
OCTOCRYLENE	IDENTIFICATION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *A./ Acceptance criteria*: Change NMT 3.0%, calculated on the as-is basis to: Absorptivities, calculated on the as-is basis, do not differ by more than 3.0%.
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*

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								system/*Column*: Change 4-mm × 30-cm; packing L1 to: 3.9-mm × 30-cm; packing L1 AND In *System suitability*: Change [Note—The relative retention times for bupivacaine hydrochloride and dibutyl 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

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								1.2, respectively.] AND In System suitability/Suitability requirements: Change **Resolution**: NLT 2.0 between bupivacaine hydrochloride and dibutyl phthalate **Relative standard deviation**: NMT 1.0% for the ratio of bupivacaine to the internal standard from three replicate injections to: **Resolution**: NLT 2.0 between bupivacaine

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								and dibutyl phthalate **Relative standard deviation**: NMT 1.0% for the peak response ratio of bupivacaine to the internal standard from three replicate injections
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,

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								and desethoxy tamsulosin are 0.73, 1.00, 1.80, and 2.80, respectively.] **Suitability 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,

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								and desethoxy tamsulosin are 0.73, 1.00, 1.80, and 2.80, respectively.] **Suitability requirements** **Tailing factor:** NMT 2.0, *Standard solution* **Relative standard deviation:** NMT 5.0%, *Standard solution* **Signal-to-noise ratio:** NLT 10, *Sensitivity solution*
PROMETHAZINE HYDROCHLORIDE TABLETS	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	26-Aug-2022	1-Sep-2022	*NA*	*NA*	Change **System suitability solution**: 5 µg/mL each of USP Promethazine Hydrochloride RS and USP

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								Promethazine Related Compound B RS from the *Standard stock solution* and *System suitability stock solution*, respectively **Standard solution**: 5 µg/mL of USP Promethazine Hydrochloride RS from the *Standard stock solution* **Sensitivity solution**: 0.25 µg/mL of USP Promethazine Hydrochloride RS from the *Standard solution* to: **System suitability**

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								solution: 5 µg/mL each of USP Promethazine Hydrochloride RS and USP Promethazine Related Compound B RS from the *Standard stock solution* and *System suitability stock solution*, respectively, in *Diluent* **Standard solution**: 5 µg/mL of USP Promethazine Hydrochloride RS from the *Standard stock solution* in *Diluent* **Sensitivity solution**: 0.25 µg/mL of USP

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								Promethazine Hydrochloride RS from the *Standard solution* in *Diluent*
METARAMINOL BITARTRATE	ADDITIONAL REQUIREMENTS/*USP Reference Standards ?11?*	*USPNF Online*	Online	27-Jan-2023	1-Feb-2023	*NA*	*NA*	In USP Metaraminol Enantiomer RS: Change 3-[(1*S*,2*R*)-2-Amino-1-hydroxypropyl]phenol. $C_9H_{13}NO_2$ 167.21 to: 3-[(1*S*,2*R*)-2-Amino-1-hydroxypropyl]phenol D-tartrate. $C_9H_{13}NO_2$? $C_4H_6O_6$ 317.29
TRIAZOLAM TABLETS	PERFORMANCE TESTS/*Dissolution* ?711?	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	In *Standard solution*: Change Tablet/mg to: mg/Tablet
AMOXICILLIN	IDENTIFICATIO	*USPNF Online*	Online	26-Jan-2024	1-Feb-2024	*NA*	*NA*	Change

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ORAL SUSPENSION	N							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 Suspension with a mixture of acetone and 0.1 N hydrochloric

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								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 gel mixture (see *Chromatograph y* ?621?). Place the plate in a

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								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 plate by spraying lightly with a solution

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								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.
DOXYCYCLINE TABLETS	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	In USP Doxycycline Related Compound A RS: Change 444.43 to: 444.44 AND Change (4*S*,4a*R*,5*S*,5a*R*,6*S*,12a*S*)-4-(Dimethylamino)-1,4,4a,5,5a

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								,6,11,12a-octahydro-3,5,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-2-naphthacenecarboxamide, monohydrochloride. $C_{22}H_{24}N_2O_8 \cdot HCl$ 480.13 to: (4*S*,4a*R*,5*S*,5a*R*,6*S*,12a*S*)-4-(Dimethylamino)-1,4,4a,5,5a,6,11,12a-octahydro-3,5,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-2-naphthacenecarboxamide hydrochloride. $C_{22}H_{24}N_2O_8 \cdot HCl$ 480.90
FELODIPINE	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	18-Nov-2022	1-Dec-2022	*NA*	*NA*	Change **Buffer**, **Mobile phase**, **System suitability**

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								solution, **Standard solution**, **Sample solution**, and **Chromatographic system**: Proceed as directed in the *Assay*. to: **Buffer**, **Mobile phase**, **System suitability solution**, **Standard solution**, **Sample solution**, and **Chromatographic system**: Proceed as directed in the *Assay*, and use 40 µL injection volume for the *Sensitivity solution*.
AMMONIUM G	SPECIFIC	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	In *Optical*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
LYCYRRHIZAT E	TESTS							*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 basis
PALONOSETR ON HYDROCH LORIDE	IMPURITIES	*USPNF Online*	Online	16-Dec-2022	1-Jan-2023	*NA*	*NA*	In *Limit of Unspecified Impurities*: Change **Mobile phase** and **Chromatographic system**: Proceed as

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								directed in the *Assay*. to: **Mobile phase**, **Standard stock solution**, and **Chromatographic system**: Proceed as directed in the *Assay*.
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,

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								and 317.9 nm from the *Sample solution* correspond to those from the *Standard solution*, as obtained in the *Content of Calcium*.
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
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.*

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

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								$^{19}H_{17}N_5O_7S_3$) in the portion of Ceftiofur Hydrochloride taken: Result = (r_U/r_S) × (C_S/C_U) × P
ERYTHROMYCIN ETHYLSUCCINATE FOR ORAL SUSPENSION	*Identification*	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	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

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								separately 10 µL each of the test solution and the Standard solution to a 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

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								solvent front has moved about 9 cm. Remove the plate from the chamber, mark 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*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								*F*values of the principal spots obtained from the test solution correspond to those obtained from the Standard solution.
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
PROPOFOL	ASSAY	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	**Procedure for 197S** In *Procedure 2*/*Mobile phase*:

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								Change Hexane, acetonitrile, and alcohol (990: 7.5: 1) to: Hexane, acetonitrile, and alcohol, absolute (990: 7.5: 1)
FORSKOHLII	COMPOSITION /*Content of Forskolin*	*USPNF Online*	Online	29-Jul-2022	1-Aug-2022	*NA*	*NA*	In *Chromatographic system*: Change Column: 4.6-mm × 25-cm; 5-µm, 100 Å to: Column: 4.6-mm × 25-cm; 5-µm, 100 Å; packing L1
AZTEC MARIGOLD ZEAXANTHIN EXTRACT	IDENTIFICATION/*C*.	*USPNF Online*	Online	27-Jan-2023	1-Feb-2023	*NA*	*NA*	Change (3*R*,3?*S* *meso*)-zeaxanthin to:

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								(3*R*,3?*S*)-zeaxanthin
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
SODIUM SALICYLATE	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *USP Reference Standards* ?11?: Add USP Phenol RS
NANDROLONE DECANOATE	ASSAY/*Procedure*	*USPNF Online*	Online	27-May-2022	1-Jun-2022	*NA*	*NA*	In *Analysis*: Change Calculate the percentage of Nandrolone Decanoate ($C_{28}H_{44}O_3$) in the portion of Nandrolone Decanoate taken: to: Calculate the percentage of nandrolone

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication Sort ascending	Target Online Fix Publication	Description
								decanoate ($C_{28}H_{44}O_3$) in the portion of Nandrolone Decanoate taken:
DOXORUBICIN HYDROCHLORIDE INJECTION	ASSAY/ *Procedure*	*USPNF 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)
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL TABLETS	PERFORMANCE TESTS/ *Dissolution* ?711?	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	In *Test 2*/*Analysis*: Change C_S = concentration of USP Norethindrone Acetate RS

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								(mg/mL) or USP Ethinyl Estradiol RS (ug/mL) in the *Standard solution* to: C_S = concentration of USP Norethindrone Acetate RS (mg/mL) or USP Ethinyl Estradiol RS (µg/mL) in the *Standard solution*
ZIPRASIDONE ASSAY CAPSULES	*USPNF Online*	Online		25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Procedure/Analysis*: Change 449.40 to: 449.39

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