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Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
BOTANICAL EXTRACTS	PREPARATIONS	*USPNF Online*	Online	27-May-2022	1-Jun-2022	*NA*	*NA*	In *General 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*

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								<*561*>)
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
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

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MESO-ZEAXANTHIN	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USPNF Online*	Online	27-Jan-2023	1-Jun-2023	*NA*	*NA*	RS (µg/mg) In USP *meso* -Zeaxanthin RS: Change (3*R*,3?*S*-*meso*)-Zeaxanthin. to: (3*R*,3?*S*)-Zeaxanthin.
CALCIUM LACTATE	CHEMICAL INFORMATION	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	Change Calcium lactate (1:2) hydrate to: Calcium lactate hydrate AND Change Calcium lactate (1:2) pentahydrate to: Calcium lactate pentahydrate
ATORVASTATIN CALCIUM TABLETS	ASSAY/ *Procedure*	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	In *Analysis*: Change M_{r1} = molecular weight of atorvastatin,

Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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
FOSAMPRENAVIR CALCIUM TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Dissolution* ?711?/*Medium*: Change 26.7 g/L of sodium acetate trihydrate in water. Add 133 mL of glacial acetic acid to this solution, and then dilute with water to 10 L; 900 mL. to:

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								0.02 M sodium acetate buffer, pH 3.5, prepared as follows. Dissolve 2.67 g of sodium acetate in 100 mL of water. Add 13.3 mL of glacial acetic acid and then dilute with water to 1000 mL; 900 mL.
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: 50 rpm. Use apex vessels.
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

Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								tartrate in *Diluent* prepared as follows. to: Nominally about 80 µg/mL of levorphanol tartrate in *Diluent* prepared as follows.
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,

Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								558.65 M_{r2} = molecular weight of atorvastatin calcium, 1155.36
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%.
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

Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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* <711>/*Test 2*/*Buffer stage*: Change **Buffer stage standard solution:**

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

Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Nandrolone Decanoate taken: to: Calculate the percentage of nandrolone decanoate ($C_{28}H_{44}O_3$) in the portion of Nandrolone Decanoate taken:
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
DOXORUBICIN HYDROCHLORIDE INJECTION	ASSAY/ *Procedure*	*USPNF Online*	Online	28-Oct-2022	1-Nov-2022	*NA*	*NA*	In *Analysis*: Change *P* = potency of

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								doxorubicin in USP Doxorubicin Hydrochloride RS (µg/mg) to: *P* = potency of doxorubicin hydrochloride in USP Doxorubicin Hydrochloride RS (µg/mg)
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: (3*R*,3?*S*)-zeaxanthin
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 (mg/mL) or USP Ethinyl Estradiol

Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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*
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
ZIPRASIDONE CAPSULES	ASSAY	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Procedure/Analysis*: Change 449.40 to: 449.39
SODIUM	ADDITIONAL R	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *USP*

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SALICYLATE	EQUIREMENTS							*Reference Standards* ?11?: Add USP Phenol RS
AZITHROMYCIN FOR ORAL SUSPENSION	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USPNF Online*	Online	27-May-2022	1-Jun-2022	*NA*	*NA*	In USP Azithromycin Related Compound F RS: Change 762.97 to: 762.98 AND In USP Desosaminylazithromycin RS: Change 590.79 to: 590.80
POWDERED 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 ×

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AZTEC MARIGOLD ZEAXANTHIN EXTRACT	COMPOSITION /*Procedure 4: Stereoisomeric Composition*	*USPNF Online*	Online	27-Jan-2023	1-Feb-2023	*NA*	*NA*	25-cm; 5-µm, 100 Å; packing L1 In *System suitability/Suitability requirements/Resolution*: Change (3*R*,3?*S meso*)-zeaxanthin to: (3*R*,3?*S*)-zeaxanthin AND In *Analysis*: Change (3*R*,3?*S meso*)-zeaxanthin to: (3*R*,3?*S*)-zeaxanthin AND In *Acceptance criteria*: Change (3*R*,3?*S*

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								meso)-Zeaxanthin to: (3*R*,3?*S*)-Zeaxanthin
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE TABLETS	IM PURITIES/*Limit of 2-Methylbenzhydrol*	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	In *Standard solution*: Change 7.5 µg/mL of USP 2-Methylbenzhydrol RS in methanol to: 7.5 µg/mL of USP Methylbenzhydrol RS in methanol AND In *Analysis*: Change C_S = concentration of USP 2-Methylbenzhydrol RS in the *Standard solution* (mg/mL) to: C_S = concentration of

Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								USP Methylbenzhydrol RS in the *Standard solution* (mg/mL)
RISEDRONATE SODIUM DELAYED-RELEASE TABLETS	PERFORMANCE TESTS/ *Dissolution* ?711?	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	In *Buffer stage*/*Analysis*: Change Calculate the percentage of the labeled amount of risedronate sodium ($C_7H_{10}NNaO_7P_7$) dissolved: to: Calculate the percentage of the labeled amount of risedronate sodium ($C_7H_{10}NNaO_7P_2$) dissolved:
ZIPRASIDONE CAPSULES	PERFORMANCE TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Dissolution* ?711?/*Test 1*/*Tier*

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								1/*Phosphate buffer, pH 7.5*: Change sodium hydroxide to: sodium hydroxide solution AND In *Dissolution* ?711?/*Test 1*/*Tier 1*/*Analysis*: Change 449.40 to: 449.39 AND In *Dissolution* ?711?/*Test 1*/*Tier 2*/*Solution A* and *Solution B*: Change sodium hydroxide to: sodium hydroxide

Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								solution AND In *Dissolution* ?711?/*Test 2*/*Tier 2*/*Analysis*: Change 449.40 to: 449.39 AND In *Dissolution* ?711?/*Test 3*/*Tier 2*/*Analysis*: Change 449.40 to: 449.39
DOXYCYCLINE HYCLATE	CHEMICAL INFORMATION	*USPNF Online*	Online	27-May-2022	1-Jun-2022	*NA*	*NA*	Change 1025.87 to: 1025.88
LATANOPROST	ASSAY/ *Procedure*	*USPNF Online*	Online	29-Jul-2022	1-Dec-2022	*NA*	*NA*	Change **Standard solution**: Transfer 2.0 mg/mL of USP Latanoprost RS into a suitable

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								volumetric flask, dissolve in dehydrated alcohol equivalent to 20% of the final volume, and dilute with chro matographic hexane to volume. **Sample solution**: Transfer 2.0 mg/mL of Latanoprost into a suitable volumetric flask, dissolve in dehydrated alcohol equivalent to 20% of the final volume, and dilute with chro matographic hexane to volume. to: **Standard**

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								solution: 2.0 mg/mL of USP Latanoprost RS prepared as follows. Transfer USP Latanoprost RS into a suitable volumetric flask, dissolve in dehydrated alcohol equivalent to 20% of the final volume, and dilute with chro matographic hexane to volume. **Sample solution**: 2.0 mg/mL of Latanoprost prepared as follows. Transfer Latanoprost into a suitable volumetric flask,

Monograph Title	Section	Source Publication Sort ascending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								dissolve in dehydrated alcohol equivalent to 20% of the final volume, and dilute with chro matographic hexane to volume.
DOFETILIDE	CHEMICAL INFORMATION	*USPNF Online*	Online	28-Oct-2022	1-Nov-2022	*NA*	*NA*	Change ?-[(*p* -Methanesulfon amidophenethyl)methylamino]m ethane sulfono-*p* -phenetidide to: *N* -{4-[2-(Methyl{2- [4-(methylsulfon amido)phenoxy] ethyl}amino)eth yl]phenyl}metha nesulfonamide
CHLOROPROC AINE HYDROC HLORIDE INJECTION	ASSAY/ *Procedure*	*USPNF Online*	Online	27-Jan-2023	1-Feb-2023	*NA*	*NA*	In *Sample solution*: Change Transfer about

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								100 mg of chloroprocaine hydrochloride to a 100-mL volumetric flask, dissolve in 40 mL of methanol, and dilute with water to volume. to: Transfer a volume of Injection, equivalent to about 100 mg of chloroprocaine hydrochloride to a 100-mL volumetric flask, add 40 mL of methanol, and dilute with water to volume.
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE TABLETS	ADDITIONAL REQUIREMENTS/*USP Reference Standards* ?11?	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	Change USP 2-Methylbenzhydrol RS to: USP Methylbenzhydrol RS

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TIOCONAZOLE	ADDITIONAL REQUIREMENTS/*USP Reference Standards* ?11?	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	In USP Tioconazole Related Compound A RS: Change 389.73 to: 389.72 AND In USP Tioconazole Related Compound B RS: Change 458.62 to: 458.60 AND In USP Tioconazole Related Compound C RS: Change $C_{16}H_{13}BrCl_2N_2OS \cdot HCl$ 468.63 to: $C_{16}H_{12}BrCl_3N_2OS \cdot HCl$ 503.06

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ZIPRASIDONE CAPSULES	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Organic Impurities/Solution B*: Change potassium hydroxide to: potassium hydroxide solution AND In *Organic Impurities/Analysis*: Change 449.40 to: 449.39
DOXYCYCLINE HYCLATE	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	27-May-2022	1-Jun-2022	*NA*	*NA*	In *Table 2*, footnote b: Change (4*S*,4a*R*,5*S*,5a*R*,6*R*,12a*S*)-2-Acetyl-4-(dimethylamino)-4a,5a,6,12a-tetrahydro-3,5,10,12,12a-pentahydroxy-6-methyl-tet

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								racene-1,11-dioxo-2-naphthacenecarboxamide. to: (4*S*,4a*R*,5*S*,5a*R*,6*R*,12a*S*)-2-Acetyl-4-(dimethylamino)-3,5,10,12,12a-pentahydroxy-6-methyl-4a,5a,6,12a-tetrahydrotetracene-1,11(4*H*,5*H*)-dione.
METHSUXIMIDE	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	26-Aug-2022	1-Sep-2022	*NA*	*NA*	In *System suitability*/*Column efficiency*: Change NTL 5800 theoretical plates to: NLT 5800 theoretical plates
PRIMIDONE	ASSAY/	*USPNF Online*	Online	28-Oct-2022	1-Dec-2022	*NA*	*NA*	In *Sample*

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TABLETS	*Procedure*							*solution*: Change 0.05 mg/mL of primidone from the *Standard stock solution* in *Diluent* to: 0.05 mg/mL of primidone from the *Sample stock solution* in *Diluent*
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.
POTASSIUM GLUCONATE ORAL SOLUTION	SPECIFIC TESTS	*USPNF Online*	Online	27-Jan-2023	1-Feb-2023	*NA*	*NA*	Change **Alcohol Determination** ?611?, *Method II—Distillation Method*: to: **Alcohol Determination**

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ROSEMARY	COMPOSITION /*Content of Phenolic Diterpenes*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	?611?, *Method I—Distillation Method*: In *Sample solution*: Change Before injection, pass through a membrane filter of 0.45-µL or finer pore size, to: Before injection, pass through a membrane filter of 0.45-µm or finer pore size,
METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER	ADDITIONAL REQUIREMENTS/*USP Reference Standards* ?11?	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	Change USP Methacrylic Acid and Ethyl Acrylate Copolymer (1:1) RS (USP Methacrylic Acid Copolymer, Type C RS) to:

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ZIPRASIDONE HYDROCHLORIDE	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	USP Methacrylic Acid and Ethyl Acrylate Copolymer (1:1) RS (USP Methacrylic Acid Copolymer Type C RS) In *Organic Impurities*/*Solution B*: Change Acetonitrile, methanol, and *Buffer* (55:5:40). Adjust with potassium hydroxide TS to a pH of 6.0. to: Acetonitrile, methanol, and *Buffer* (55:5:40). Adjust with potassium hydroxide solution to a pH of 6.0. AND

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								In both equations in *Organic Impurities/Analysis*: Change 449.40 to: 449.39
MAGNESIUM OXIDE	IMPURITIES/*Limit of Calcium*	*USPNF Online*	Online	25-Feb-2022	1-Dec-2022	*NA*	*NA*	In *Analysis*: Change C_U = concentration of Magnesium Hydroxide in the *Sample solution* (mg/mL) to: C_U = concentration of Magnesium Oxide in the *Sample solution* (mg/mL)
DOXYCYCLINE HYCLATE	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USPNF Online*	Online	27-May-2022	1-Jun-2022	*NA*	*NA*	In USP Doxycycline Related Compound A RS: Change 444.43

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								to: 444.44 AND Change 480.13 to: 480.90
ENZACAMENE	ASSAY/ *Procedure*	*USPNF Online*	Online	28-Oct-2022	1-Dec-2022	*NA*	*NA*	In *Analysis*: Change C_U = concentration of enzacamene in the *Sample solution* (mg/mL) to: C_U = concentration of Enzacamene in the *Sample solution* (mg/mL)
CEFDINIR	IMPURITIES	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *Organic Impurities*/*Table 2*/footnote a: Change 1*N*-[(*Z*)-2-(2-Aminothiazol-4-yl)-2-(hydroxyimino)acetyl]

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								glycine. to: *N*-[(*Z*)-2-(2-Aminothia zol-4-yl)-2-(hydr oxyimino)acetyl] glycine.
LEAD	PROCEDURES /*Procedure 1: Chemical Method*	*USPNF Online*	Online	27-Jan-2023	1-Jun-2023	*NA*	*NA*	In *Analysis*: Change Add to the acid solution 5.0 mL of *Standard dithizone solution* and 4 mL of *Ammonia cyanide solution*, to: Add to the acid solution 5.0 mL of *Standard dithizone solution* and 4 mL of *Ammonium cyanide solution*,
POWDERED ROSEMARY	COMPOSITION /*Content of Phenolic*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	In *Sample solution*: Change

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	Diterpenes							Before injection, pass through a membrane filter of 0.45-µL or finer pore size, to: Before injection, pass through a membrane filter of 0.45-µm or finer pore size,
METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER	IDENTIFICATION/*A*.	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	Change USP Methacrylic Acid and Ethyl Acrylate Copolymer (1:1) RS to: USP Methacrylic Acid and Ethyl Acrylate Copolymer (1:1) RS (USP Methacrylic Acid Copolymer Type C RS)
AZITHROMYCIN	CHEMICAL INFORMATION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change 748.98

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CHROMATOGRAPHY	SYSTEM SUITABILITY	*USPNF Online*	Online	25-Mar-2022	1-Dec-2022	*NA*	*NA*	to: 749.00 AND Change 767.00 to: 767.01 AND Change 785.02 to: 785.03 Change **System Repeatability—Assay of an Active Substance or an Excipient** to: **System Repeatability**

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