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How to Use

- **Searching:** Type keyword in search field at top of page. Search by all or part of a monograph title. For searches using multiple criteria, you will find items that match each of the specified criteria unless quotation marks are used.
 - For example, a search on Aminosalicyclic Acid Tablets will result in anything that contains “Aminosalicyclic” OR “Acid” OR “Tablets”
 - A search for “Aminosalicyclic Acid Tablets” will result in anything that specifically contains “Aminosalicyclic Acid Tablets”
- **Sorting:** Click on any column header title to sort alphabetically or chronologically in ascending or descending order. Note: the page load column is sorted alphabetically so that a number is ordered by first digit vs. by the actual number; thus, numbers will not always be in order.
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ENZACAMENE ASSAY/ <i>Procedure</i>	<i>USPNF Online</i>	Online	28-Oct-2022	1-Dec-2022	NA	NA	In <i>Analysis</i> : Change C_U = concentration of enzacamene in the <i>Sample solution</i> (mg/mL) to: C_U = concentration of Enzacamene in the <i>Sample solution</i> (mg/mL)
POWDERED ROSEMARY COMPOSITION <i>/Content of Phenolic Diterpenes</i>	<i>USPNF Online</i>	Online	28-Apr-2023	1-May-2023	NA	NA	In <i>Sample solution</i> : Change Before injection, pass through a membrane filter of 0.45- μ L or finer pore size, to: Before injection, pass through a membrane filter of 0.45- μ m or finer pore size, Change
AZITHROMYCI CHEMICAL	<i>USPNF Online</i>	Online	25-Aug-2023	1-Sep-2023	NA	NA	Change

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N	INFORMATION								748.98 to: 749.00 AND Change 767.00 to: 767.01 AND Change 785.02 to: 785.03 Change Proceed as directed in <i>Identification</i> test A under <i>Metaproterenol Sulfate Inhalation Solution</i> , beginning with “Allow the spots to dry”: to: Allow the spots to dry, and develop the chromatogram
METAPROTERENOL SULFATE ORAL SOLUTION	<i>Identification</i> /A.	USPNF Online	Online	24-Feb-2023	1-Mar-2023	NA	NA		

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TRIMETHOBE	CHEMICAL	USPNF Online	Online	28-Jul-2023	1-Aug-2023	NA	NA	in a solvent system consisting of the upper layer of a freshly prepared mixture of butyl alcohol, water, and formic acid (50:25:7) until the solvent front has moved about three-fourths of the length of the plate. Remove the plate from the developing chamber, mark the solvent front, and allow the solvent to evaporate. Locate the spots on the plate by examination under short-wavelength UV light: Change

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NZAMIDE HYD INFORMATION ROCHLORIDE							_Sort descending_
							<i>N</i>-[<i>p</i>-[2-(Dimethylamino)ethoxy]benzyl]-3,4,5-trimethoxy benzamide monohydrochloride to: <i>N</i>-[4-[2-(Dimethylamino)ethoxy]benzyl]-3,4,5-trimethoxybenzamide monohydrochloride
COCOYL CAP SPECIFIC RYLOCAPRAT TESTS E	USPNF Online Online		23-Feb-2024	1-May-2024	NA	NA	In <i>Fats and Fixed Oils</i> ?401?, <i>Procedures, Hydroxyl Value/Analysis:</i> Change Calculate the acid value as directed in the chapter. to: Calculate the hydroxyl value as directed in

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MILK THISTLE CAPSULES	PERFORMANCE TESTS/ <i>Disintegration and Dissolution</i> <2040>, <i>Dissolution</i>	USPNF Online	Online	24-Jun-2022	1-Aug-2022	NA	NA	the chapter. In <i>Medium</i> : Change <i>Buffer</i> containing 2% lauryl sulfate; 900 mL to: <i>Buffer</i> containing 2% sodium lauryl sulfate; 900 mL
PENICILLIN G BENZATHINE INJECTABLE SUSPENSION	IDENTIFICATION	USPNF Online	Online	18-Nov-2022	1-Dec-2022	NA	NA	Change It responds to the <i>Identification test</i> under <i>Penicillin G Benzathine Oral Suspension</i> . to: Mix a portion of it with methanol to obtain a solution containing about 3000 Penicillin G Units per mL. Apply 20 µL of

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							this test solution and 20 µL of a Standard solution of USP Penicillin G Benzathine RS in methanol containing 2.5 mg per mL to a suitable thin-layer chromatographic plate (see <i>Chromatography</i> ?621?) coated with a 0.25-mm layer of chromatographic silica gel, and allow the spots to dry. Using an unlined developing chamber, develop the chromatogram in a solvent system consisting of a mixture of

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							methanol, acetonitrile, and ammonium hydroxide (70:30:3) until the solvent front has moved about three-fourths of the length of the plate. Remove the plate from the developing chamber, and allow to air-dry. Spray the plate uniformly with a spray reagent prepared as follows. Dissolve 20 g of tartaric acid and 1.7 g of bismuth subnitrate in 80 mL of water. Add 2.5 mL of this solution, 2.5 mL of potassium iodide solution

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ESCITALOPRA ASSAY/ M TABLETS <i>Procedure</i>	USPNF Online Online		28-Apr-2023	1-May-2023	NA	NA	(4 in 10), and 10 g of tartaric acid to 50 mL of water, and mix. Examine the chromatograms: the principal spot obtained from the test solution corresponds in R_F value to that obtained from the Standard solution. In <i>Buffer</i> : Change Adjust with 1 N sodium hydroxide VS to a pH of 5.2. to: Adjust with 1 N sodium hydroxide to a pH of 5.2.
OIL- AND WAT STRENGTH ER-SOLUBLE VITAMINS WITH	USPNF Online Online		27-Oct-2023	1-Nov-2023	NA	NA	In <i>Vitamin A, Method 3/Analysis</i> : Change

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MINERALS CAPSULES								C_S = concentration of retinyl acetate ($C_{23}H_{32}O_2$) from USP Vitamin A RS in the <i>Standard solution</i> (?g/mL) to: C_S = concentration of retinyl acetate ($C_{22}H_{32}O_2$) from USP Vitamin A RS in the <i>Standard solution</i> (?g/mL)
METHOTREXATE	ADDITIONAL REQUIREMENTS/USP Reference Standards ?11?	USPNF Online	Online	24-Jun-2022	1-Jul-2022	NA	NA	In USP Methotrexate System Suitability Mixture RS: Change $C_{22}H_{26}N_8O_5 \cdot HCl$ 518.95 to: $C_{22}H_{26}N_8O_5 \cdot x$ HCl
MESO-ZEAXANTHIN	COMPOSITION /Stereoisomeric	USPNF Online	Online	27-Jan-2023	1-Feb-2023	NA	NA	In System suitability:

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PREPARATION							Change [Note—The approximate relative retention times for (3S,3?S)-zeaxanthin, (3R,3?S-meso)-zeaxanthin, (3R,3?R)-zeaxanthin, and (3R,3?R,6?R)-lutein are 0.94, 1.00, 1.06, and 1.11, respectively.] to: [Note—The approximate relative retention times for (3S,3?S)-zeaxanthin, (3R,3?S)-zeaxanthin, (3R,3?R

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							)-zeaxanthin, and (3R,3?R,6?R)-lutein are 0.94, 1.00, 1.06, and 1.11, respectively.] AND In System <i>suitabil</i> <i>ity/Suitability</i> <i>re</i> <i>quire</i> <i>ments/</i> <i>Resolution:</i> Change (3R,3?S- <i>meso</i>)-zeaxanthin to: (3R,3?S)-zeaxanthin AND In Analysis: Change (3R,3?S- <i>meso</i>)-zeaxanthin to: (3R,3?S

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METHYLPHENIDATE HYDROCHLORIDE EXTENDED-RELEASE TABLETS	ADDITIONAL REQUIREMENT S/USP Reference Standards ?11?	USPNF Online	Online	26-May-2023	1-Jun-2023	NA	NA	)-zeaxanthin AND In Acceptance criteria: Change (3R,3?S-meso)-Zeaxanthin to: (3R,3?S)-Zeaxanthin In USP Methylphenidate Related Compound A RS: Change ?-Phenyl-2-piperidineacetic acid hydrochloride. to: (RS)-2-Phenyl-2-[(RS)-piperidin-2-yl]acetic acid hydrochloride. In Dissolution ?711?: Move Test 2 after
DICLOFENAC SODIUM AND MISOPROSTOL	PERFORMANCE TESTS	USPNF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	

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L DELAYED- RELEASE TABLETS									<i>Test 1</i>
CARBAMAZEPI NE TABLETS	ADDITIONAL R EQUIREMENT <i>S/USP</i> <i>Reference</i> <i>Standards <11></i>	<i>USPNF Online</i>	Online	27-May-2022	1-Jun-2022	NA	NA		In USP Carbamazepine Related Compound A RS: Change 238.28 to: 238.29 AND In USP Carbamazepine Related Compound B RS: Change 193.24 to: 193.25 AND In USP 9-Methy lacridine RS: Change 193.24 to: 193.25 Change USP Calcium D L-5-Methyltetra
CALCIUM L-5- METHYLTETR AHYDROFOLA	ADDITIONAL R EQUIREMENT <i>S/USP</i>	<i>USPNF Online</i>	Online	28-Oct-2022	1-Nov-2022	NA	NA		

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TE	<i>Reference Standards <11></i>							hydrofolate RS to: USP Calcium D, L-5-Methyltetrahydrofolate RS In footnote 1 in <i>Molecular Weight Measurements of Low Molecular Weight Heparins by Gel Permeation Chromatography</i> system/ Columns/Analytical: Change guard column TSK SWXL 6-mm × 4-mm; to: guard column TSK SWXL 6-mm × 4-cm;
LOW MOLECULAR WEIGHT HEPARIN MOLECULAR WEIGHT DETERMINATIONS	PROCEDURE	<i>USPNF Online</i> Online		31-Mar-2023	1-Apr-2023	NA	NA	

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CRANBERRY FRUIT DRY JUICE	DEFINITION	USPNF Online	Online	25-Aug-2023	1-Sep-2023	NA	NA	Change procyanidin A ₂ to:
SUCROSE	IMPURITIES	USPNF Online	Online	29-Mar-2024	1-Apr-2024	NA	NA	procyanidin A ₂ In <i>Sulfite/Analysis</i> s/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 (Catalogue No.: AK00071). to: Test kit for sulfite determination may be ordered

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								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).
DACARBAZINE IM FOR INJECTION	PURITIES/ <i>Organic Impurities</i>	USPNF Online	Online	26-Aug-2022	1-Sep-2022	NA	NA	In System suitability/Suitability requirements/Signal-to-noise ratio: Change NTL 10, to: NLT 10, Change
LEAD	REQUIREMENTS FOR	USPNF Online	Online	27-Jan-2023	1-Jun-2023	NA	NA	Change • Precision

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	PROCEDURE VALIDATION							Repeatability to: • Precision In <i>Analysis</i> : Change M_{r1} = molecular weight of calcium ascorbate dihydrate, 426.43 to: M_{r1} = molecular weight of calcium ascorbate dihydrate, 426.34
CALCIUM ASCORBATE	ASSAY/ <i>Procedure</i>	<i>USPNF Online</i>	Online	30-Jun-2023	1-Aug-2023	NA	NA	
CEFDINIR FOR ADDITIONAL ORAL SUSPENSION	ADDITIONAL REQUIREMENT S	<i>USPNF Online</i>	Online	17-Nov-2023	1-Dec-2023	NA	NA	In <i>USP Reference Standards</i> ?11?/USP Cefdinir Related Compound A RS: Change 413.43 to: 413.42 AND In <i>USP</i>

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									<i>Reference Standards</i> ?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 In <i>Analysis</i> : Change Result = $(R_U/R_S) \times C_S \times V$ $\times (D/W_U) \times (1/L)$ $\times 100$ R_U = peak area ratio of ibuprofen to benzophenone from the <i>Sample solution</i> R_S = peak area ratio of ibuprofen to benzophenone from the <i>Standard solution</i>
IBUPROFEN ORAL SUSPENSION	PERFORMANC E TESTS/ <i>Dissolution</i> <711>	USPNF Online	Online	27-May-2022	1-Jun-2022	NA	NA		

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						Sort descending	C_S = concentration of USP Ibuprofen RS in the <i>Standard solution</i> (mg/mL) V = volume of <i>Medium</i>, 900 mL D = density of Oral Suspension (g/mL) W_U = weight of the portion of Oral Suspension added to the <i>Medium</i> (g) L = label claim (mg/mL) to: Result = $(R_U/R_S) \times C_S \times V \times (d/W_U) \times D \times (1/L) \times 100$ R_U = peak area ratio of ibuprofen to

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							benzophenone from the <i>Sample solution</i> R_S = peak area ratio of ibuprofen to benzophenone from the <i>Standard solution</i> C_S = concentration of USP Ibuprofen RS in the <i>Standard solution</i> (mg/mL) V = volume of <i>Medium</i> , 900 mL d = density of Oral Suspension (g/mL) W_U = weight of the portion of Oral Suspension added to the <i>Medium</i> (g)

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DIAZOXIDE	IMPURITIES/ <i>Organic Impurities</i>	USPNF Online	Online	28-Oct-2022	1-Nov-2022	NA	NA	<i>D</i> = dilution factor of the <i>Sample solution</i>, 2 <i>L</i> = label claim (mg/mL) In footnote a in <i>Table 2</i>: Change 7-Chloro-2H-benzothiazine-3(4H)-one 1,1-dioxide. to: 7-Chloro-2H-1,2,4-benzothiazine-3(4H)-one 1,1-dioxide.
ROSEMARY LEAF DRY AQUEOUS EXTRACT	COMPOSITION / <i>Content of Rosmarinic Acid</i>	USPNF Online	Online	28-Apr-2023	1-May-2023	NA	NA	In <i>Standard solution B</i>: Change Before injection, pass through a membrane filter of 0.45-µL or finer pore size,

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AZITHROMYCI IMPURITIES N	USPNF Online	Online	25-Aug-2023	1-Sep-2023	NA	NA	to: Before injection, pass through a membrane filter of 0.45-µm or finer pore size, AND In <i>Sample solution</i>: 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, In <i>Organic Impurities/</i>Table 2: Change: 3? -N -[4-(Acetylamin o)phenyl]sulfonyl}-3?-demethyl azithromycin^m to:

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							3'-N -D emet hyl-3'-N -[(4-methylphenyl)sulfonyl]azithromycin ^m AND In <i>Organic Impurities/</i> Table 2/footnote m: Change (2 <i>R</i> ,3 <i>S</i> ,4 <i>R</i> ,5 <i>R</i> ,8 <i>R</i> ,10 <i>R</i> ,11 <i>R</i> ,12 <i>S</i> ,13 <i>S</i> ,14 <i>R</i>)-13-[(2,6-Dideoxy-3-C-methyl-3- <i>O</i> -methyl-?- <i>L</i> -ribo-hexopyranosyl)oxy]-2-ethyl-3,4,10-trihydroxy-3,5,6,8,10,12,14-heptamethyl-11-[[3-[<i>N</i> -(4-acetamidophenyl)sulf

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							onyl)- <i>N</i> -methylamino]-3 ,4,6-trideoxy-?- D- <i>xyl</i> -hexopyranosyl] oxy]-1-oxa-6-az acyclopentadec an-15-one. to: (2 <i>R</i> ,3 <i>S</i> ,4 <i>R</i> ,5 <i>R</i> ,8 <i>R</i> ,10 <i>R</i> ,11 <i>R</i> ,12 <i>S</i> ,13 <i>S</i> ,14 <i>R</i>)-13-[(2,6-Dideo xy-3- <i>C</i> -meth yl-3- <i>O</i> -methyl-?- L- <i>ribo</i> -hexopyranosyl) oxy]-2-ethyl-3,4, 10-trihydroxy-3, 5,6,8,10,12,14- heptamethyl-11- [[3-[<i>N</i> -(4-methylpheny lsulfon yl)- <i>N</i> -methylamino]-3 ,4,6-trideoxy-β- D- <i>xyl</i> -hexopyranosyl]

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OIL-SOLUBLE VITAMINS CAPSULES	STRENGTH	USPNF Online	Online	27-Oct-2023	1-Nov-2023	NA	NA	oxy]-1-oxa-6-azacyclopentadecan-15-one. In <i>Vitamin A, Method 3/Analysis:</i> Change C_S = concentration of retinyl acetate ($C_{23}H_{32}O_2$) from USP Vitamin A RS in the <i>Standard solution</i> (?g/mL) to: C_S = concentration of retinyl acetate ($C_{22}H_{32}O_2$) from USP Vitamin A RS in the <i>Standard solution</i> (?g/mL)
TRIETHYL CITRATE	ASSAY/ <i>Procedure</i>	USPNF Online	Online	24-Feb-2023	1-May-2023	NA	NA	In <i>System suitability/Suitability requirements/Tailing</i>

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FLURAZEPAM HYDROCHLORIDE	USP REFERENCE STANDARDS ?11?	USPNF Online	Online	23-Feb-2024	1-Mar-2024	NA	NA	<i>factor</i>: Change NMT 1.5 for the triethyl citrate to trimethyl citrate peaks to: NMT 1.5 for the triethyl citrate and trimethyl citrate peaks In USP Flurazepam Related Compound C RS: Change 5-Chloro-2-(2-diethylaminoethyl (amino)-2?-fluorobenzophenone hydrochloride. to: 5-Chloro-2-(2-diethylaminoethyl amino)-2?-fluorobenzophenone hydrochloride.
MILK THISTLE TABLETS	PERFORMANCE TESTS/ <i>Disintegration</i>	USPNF Online	Online	24-Jun-2022	1-Aug-2022	NA	NA	In <i>Medium</i>: Change <i>Buffer</i> containing 2%

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								lauryl sulfate; 900 mL to: <i>Buffer</i> containing 2% sodium lauryl sulfate; 900 mL
PENICILLIN G BENZATHINE AND PENICILLIN G PROCAINE INJECTABLE SUSPENSION	IDENTIFICATION	USPNF Online	Online	18-Nov-2022	1-Dec-2022	NA	NA	In <i>Test A</i> : Change It responds to the <i>Identification</i> test under <i>Penicillin G Benzathine Oral Suspension</i> : the spot obtained from the test solution, corresponding in R_F value to that obtained from the Standard solution, is completely resolved from a second spot, produced by penicillin G

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							procaine. to: Mix a portion of it with methanol to obtain a solution containing about 3000 Penicillin G Units per mL. Apply 20 µL of this test solution and 20 µL of a Standard solution of USP Penicillin G Benzathine RS in methanol containing 2.5 mg per mL to a suitable thin-layer chromatographic plate (see <i>Chromatography</i> ?621?) coated with a 0.25-mm layer of chromatographic silica gel, and allow

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							the spots to dry. Using an unlined developing chamber, develop the chromatogram in a solvent system consisting of a mixture of methanol, acetonitrile, and ammonium hydroxide (70:30:3) until the solvent front has moved about three-fourths of the length of the plate. Remove the plate from the developing chamber, and allow to air-dry. Spray the plate uniformly with a spray reagent prepared as

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							follows. Dissolve 20 g of tartaric acid and 1.7 g of bismuth subnitrate in 80 mL of water. Add 2.5 mL of this solution, 2.5 mL of potassium iodide solution (4 in 10), and 10 g of tartaric acid to 50 mL of water, and mix. Examine the chromatograms: the principal spot obtained from the test solution corresponds in R_F value to that obtained from the Standard solution. The spot obtained from the test solution, corresponding

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ESCITALOPRA PERFORMANC M TABLETS	E TESTS/ <i>Dissolution</i> ?711?	USPNF Online	Online	28-Apr-2023	1-May-2023	NA	NA		in R_F value to that obtained from the Standard solution, is completely resolved from a second spot, produced by penicillin G procaine. In <i>Test 1/Medium:</i> Change 0.1 N hydrochloric acid VS; 900 mL to: 0.1 N hydrochloric acid; 900 mL AND In <i>Test 2/Medium:</i> Change 0.1 N hydrochloric acid VS; 900 mL

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TRYPTOPHAN	ADDITIONAL REQUIREMENT <i>S/USP Reference Standards ?11?</i>	<i>USPNF Online</i>	Online	24-Jun-2022	1-Jul-2022	NA	NA		to: 0.1 N hydrochloric acid; 900 mL In USP Tryptophan Related Compound A RS: Change 3,3'-[Ethylidene bis(1 <i>H</i> -indole-1,3-diyl)] bis[2 <i>S</i>)-2-aminopropionic]acid. $C_{24}H_{26}N_4O_4$ 432.49 to: (2 <i>S</i> ,2' <i>S</i>)-3,3'-[Ethane-1,1-diyl bis(1 <i>H</i> -indole-1,3-diyl)] bis(2-aminopropionic acid). $C_{24}H_{26}N_4O_4$ 434.50
MESO-ZEAXANTHIN PREPARATIONS	ADDITIONAL REQUIREMENT <i>S/USP</i>	<i>USPNF Online</i>	Online	27-Jan-2023	1-Feb-2023	NA	NA		In USP <i>meso</i> -Zeaxanthin RS:

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									Change (3R,3?S-meso)-Zeaxanthin. to: (3R,3?S)-Zeaxanthin.
ATORVASTATIN CALCIUM	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USPNF Online	Online	26-May-2023	1-Jun-2023	NA	NA		Change USP Atorvastatin Related Compound H RS (lactone impurity) to: USP Atorvastatin Related Compound H RS Also known as Lactone impurity; AND Change USP Atorvastatin Related Compound I RS (acetonide

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DUTASTERIDE PERFORMANCE AND TAMSULOSIN HYDROCHLORIDE CAPSULES	USP NF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	impurity) to: USP Atorvastatin Related Compound I RS Also known as Acetonide impurity; In <i>Dissolution</i> ?711?/Test for dutasteride/Tier 2/Medium: Change 10 g/L of cetyltri methylammonium 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,

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FLUOCINOLON E ACETONIDE	<i>Organic Impurities</i> <i>USPNF Online</i>	Online	27-May-2022	1-Jun-2022	NA	NA	purified in 1000 mL of 0.1 N hydrochloric acid; 900 mL In Acceptance criteria/Total impurities: Change NMT 2.5%. Disregard any peak below 0.05% of the peak area of fluocinolone acetone from the <i>Standard solution</i> . to: NMT 2.5%. Disregard any peak below 0.05% of the peak area of fluocinolone acetone from the <i>Sample solution</i> .
TETRACYCLIN E HYDROCHL ORIDE	SPECIFIC TESTS/ <i>Bacterial</i> <i>USPNF Online</i>	Online	28-Oct-2022	1-Nov-2022	NA	NA	Change Where the label states

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<i>Endotoxins Test</i> <85>							tetracycline hydrochloride must be subjected to further processing during the preparation of injectable dosage forms, the level of bacterial endotoxins are such that the requirement under the relevant dosage form monograph(s) in which tetracycline hydrochloride is used can be met. to: Where the label states Tetracycline Hydrochloride must be

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DEXTRATES	ASSAY/ <i>Dextrose Equivalent</i>	USPNF Online	Online	31-Mar-2023	1-May-2023	NA	NA	subjected to further processing during the preparation of injectable dosage forms, the level of bacterial endotoxins are such that the requirement under the relevant dosage form monograph(s) in which Tetracycline Hydrochloride is used can be met. <i>In Analysis:</i> Change r_A = peak response of each saccharide degree of polymerization (DP

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							$1-DP_3$) in the <i>Sample solution</i> (If any peaks of DP_4 and above are observed in the sample, take the summation of all peak responses DP_{4+} and use this value as r_U.) to: r_A = peak response of each saccharide degree of polymerization (DP_1-DP_3) in the <i>Sample solution</i> (If any peaks of DP_4 and above are observed in the sample, take the summation of all peak responses DP_{4+} and use this

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CRANBERRY FRUIT DRY JUICE	IDENTIFICATION	USPNF Online	Online	25-Aug-2023	1-Sep-2023	NA	NA	value as r_A .) In A./Standard solution B: Change USP Procyanidin A ₂ RS to: USP Procyanidin A ₂ RS AND In A./System suitability/Suitability requirements/ Derivatization reagent B/White light: Change Standard solution B exhibits two brown bands in the upper-half section corresponding to procyanidin A

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						Sort descending	2and epicatechin. <i>Standard solution C</i> exhibits two brown bands in the upper-half section corresponding in R_F and color to procyanidin A_2 and epicatechin in <i>Standard solution B</i>. <i>Standard solution C</i> also exhibits a series of faint or indistinct brown bands of differing intensities below procyanidin A_2 in the lower-half section. to: <i>Standard solution B</i>

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							exhibits two brown bands in the upper-half section corresponding to procyanidin A2 and epicatechin. <i>Standard solution C</i> exhibits two brown bands in the upper-half section corresponding in R_F and color to procyanidin A2 and epicatechin in <i>Standard solution B</i>. <i>Standard solution C</i> also exhibits a series of faint or indistinct brown bands of differing intensities below

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							procyanidin A2 in the lower-half section. AND In A./ <i>Acceptance criterial</i> <i>Derivatization reagent B/White light</i>: Change The <i>Sample solution</i> exhibits two faint brown bands corresponding in R_F to procyanidin A₂ and epicatechin in <i>Standard solution B</i>. The <i>Sample solution</i> also exhibits a series of faint or indistinct brown bands of differing intensities in the lower-half section, corresponding

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						Sort descending	to the same bands in <i>Standard solution C</i>. No bands corresponding in R_F to epigallocatechin-3-O-gallate, procyanidin B₂, or procyanidin B₁ appear below procyanidin A₂. to: The <i>Sample solution</i> exhibits two faint brown bands corresponding in R_F to procyanidin A₂ and epicatechin in <i>Standard solution B</i>. The <i>Sample solution</i> also exhibits a series of faint or indistinct brown

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							bands of differing intensities in the lower-half section, corresponding to the same bands in <i>Standard solution C</i> . No bands corresponding in R_F to epigallocatechin-3-O-gallate, procyanidin B2, or procyanidin B1 appear below procyanidin A2. AND In <i>C./ Acceptance criteria/Profile at 520 nm</i> : Change cyanidin-3-O-arabinose,

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DRY HEAT DE INTRODUCTION PYROGENATION	USPNF Online	Online	23-Feb-2024	1-Mar-2024	NA	NA	to: cyanidin-3-O-arabinoside, AND In C./ Acceptance criteriaProfile at 520 nm: Change peonidin-3-O-arabinose to: peonidin-3-O-arabinoside Change parental manufacturing to: parenteral manufacturing Change NTL 90.0% and NMT 110.0% to: NLT 90.0% and NMT 110.0% In Analysis:
IRINOTECAN HYDROCHLORIDE INJECTION	USPNF Online	Online	26-Aug-2022	1-Sep-2022	NA	NA	
ISOBUTYL ASSAY/	USPNF Online	Online	27-Jan-2023	1-Feb-2023	NA	NA	

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ALCOHOL	<i>Procedure</i>								Change r_S = sum of all the peaks except those each of which with an area less than 0.1 times the area of the major peak from the <i>Reference solution</i> to: r_T = sum of all the peaks except those each of which with an area less than 0.1 times the area of the major peak from the <i>Reference solution</i>
ATORVASTATIN CALCIUM TABLETS	IMPURITIES/ <i>Organic Impurities</i>	USPNF Online	Online	30-Jun-2023	1-Jul-2023	NA	NA		In <i>Analysis</i> : Change M_{r1} = molecular weight of atorvastatin, 558.64

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ANTIBIOTICS—APPENDICES MICROBIAL ASSAYS		USPNF Online	Online	29-Dec-2023	1-Jan-2024	NA	NA		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 In two instances in <i>Appendix 1</i> equations: Change 14.020 to: 14.022 Change 2-(2-Carboxyethyl)-1,2,3,4-tetrahydro-6,7-dimethoxy-2-methyl-1-veratrylisoquinolinium benzenesulfonate,
ATRACURIUM BESYLATE	CHEMICAL INFORMATION	USPNF Online	Online	27-May-2022	1-Jun-2022	NA	NA		

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DOXAZOSIN MESYLATE	IMPURITIES/ <i>Organic Impurities</i>	USPNF Online	Online	28-Oct-2022	1-Nov-2022	NA	NA	pentamethylene ester to: 2-(2-Carboxyethyl)-1,2,3,4-tetrahydro-6,7-dimethoxy-2-methyl-1-veratrylisoquinolinium benzenesulfonate, pentamethylene ester In <i>Analysis</i>: Change C_S = concentration of the corresponding USP Doxazosin Related Compound RS or USP Doxazosin Mesylate RS (for calculating unspecified impurities) in the <i>Standard solution</i> (mg/mL)

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RISEDRONATE ASSAY SODIUM DELA YED-RELEASE TABLETS	USPNF Online	Online	28-Apr-2023	1-May-2023	NA	NA	to: C _S = concentration of the corresponding USP Reference Standard or USP Doxazosin Mesylate RS (for calculating unspecified impurities) in the <i>Standard solution</i> (mg/mL) Delete [Note—Use a non-metallic liquid chromatography system for analysis.]
AZITHROMYCIN ADDITIONAL R EQUIREMENT S	USPNF Online	Online	25-Aug-2023	1-Sep-2023	NA	NA	In <i>USP Reference Standards</i> ?11?/USP Azae rythromycin A RS: Change 734.96 to:

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								734.97 AND In USP Azithromycin Related Compound F RS: Change 762.97 to: 762.98 AND In USP Desosa minylazithromyc in RS: Change 590.79 to: 590.80

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