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How to Use

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Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
OIL- AND WAT	STRENGTH	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Vitamin A,*

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ER-SOLUBLE VITAMINS WITH MINERALS CAPSULES								*Method 3*/*Analysis*: Change C_S = concentration of retinyl acetate ($C_{23}H_{32}O_2$) from USP Vitamin A RS in the *Standard solution* (?g/mL) to: C_S = concentration of retinyl acetate ($C_{22}H_{32}O_2$) from USP Vitamin A RS in the *Standard solution* (?g/mL)
UREA	SPECIFIC TESTS	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Alcohol-Insoluble Matter*/*Sample solution*: Change 100 mg/mL of Urea dissolved in warm alcohol to: Dissolve 5.0 g of Urea in 50

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

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

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								telmisartan and telmisartan related compound A **Tailing factor:** NMT 2.0 for telmisartan **Relative standard deviation:** NMT 2.0% for telmisartan
PARTICLE SIZE ANALYSIS BY DYNAMIC LIGHT SCATTERING	GLOSSARY	*USPNF Online*	Online	27-Oct-2023	1-May-2024	*NA*	*NA*	In *Average particle diameter*: Change expressed in nanometers. to: expressed in meters. AND In *Viscosity*: Change in millipascal-seconds (mPa?s). to: in pascal-seconds (Pa?s).

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

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

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

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								Standard solution **Relative standard deviation:** NMT 5.0%, *Standard solution* **Signal-to-noise ratio:** NLT 10, *Sensitivity solution*
DIGOXIN TABLETS	ASSAY	*USPNF Online*	Online	29-Sep-2023	1-Oct-2023	*NA*	*NA*	In *Procedure/Analysis*: Change C_U = nominal concentration of in the *Sample solution* (µg/mL) to: C_U = nominal concentration of digoxin in the *Sample solution* (µg/mL)
DIVALPROEX SODIUM EXTENDED-RELEASE TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	29-Sep-2023	1-Oct-2023	*NA*	*NA*	In *Dissolution* ?711?/*Test 8*/*Tolerances*: Change The percentage

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								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*.
UREA C 13	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Isotopic Purity/ Chromatographic system*: Change **Flow rate**: Flow rate

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								to: **Flow rate**: 1 mL/min
BLACK CUMIN SEED THYMOQUINONE OIL	SPECIFIC TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Fats and Fixed Oils* ?401?, *Procedures*, *Fatty Acid Composition*/*Table 2*: Change Linoleic to: Linoleic acid
CRANBERRY FRUIT DRY JUICE	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *USP Reference Standards* ?11?: Change USP Procyanidin A_2 RS to: USP Procyanidin A2 RS
ZIPRASIDONE CAPSULES	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Organic Impurities*/*Solution B*: Change potassium hydroxide

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								to: potassium hydroxide solution AND In *Organic Impurities/Analysis*: Change 449.40 to: 449.39
VALGANCICLOVIR HYDROCHLORIDE	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Organic Impurities/Table 3*/footnote c: Change 2-[(2-Amino-6-oxo-1,6-dihydro-9*H*-purin-9-yl)methoxy]-2-hydroxypropyl methyl-L-valinate hydrochloride. to: 3-[(2-Amino-6-oxo-1,6-dihydropurin-9-yl)methoxy]-2-hydroxypropyl L-valinate hydrochloride.

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ARIPIPRAZOLE TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Dissolution <711>/Test 1/ Procedure/ Chromatographic procedure/Analysis*: Change Result = $(R_U/R_S) \times C_S \times V \times (1/L) \times 100$ to: Result = $(R_U/R_S) \times C_S \times V \times D \times (1/L) \times 100$ AND Add *D* = dilution factor of the *Sample solution*, 2
CRANBERRY FRUIT DRY JUICE	IDENTIFICATION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *A./Standard solution B*: Change USP Procyanidin A_2 RS to:

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								USP Procyanidin A2 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_2 and epicatechin. *Standard solution C* exhibits two brown bands in the upper-half section corresponding in *R*

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								*F*and color to procyanidin A_2 and epicatechin in *Standard solution B*. *Standard solution C* also exhibits a series of faint or indistinct brown bands of differing intensities below procyanidin A_2 in the lower-half section. to: *Standard solution B* exhibits two brown bands in the upper-half section corresponding to procyanidin A2 and epicatechin. *Standard solution C*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								exhibits two brown bands in the upper-half section corresponding in R_F and color to procyanidin A2 and epicatechin in *Standard solution B*. *Standard solution C* also exhibits a series of faint or indistinct brown bands of differing intensities below procyanidin A2 in the lower-half section. AND In *A./ Acceptance criteria/ Derivatization reagent B/White light*: Change The *Sample*

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								solution exhibits two faint brown bands corresponding in R_F to procyanidin A_2 and epicatechin in *Standard solution B*. The *Sample solution* also exhibits a series of faint or indistinct brown bands of differing intensities in the lower-half section, corresponding to the same bands in *Standard solution C*. No bands corresponding in R_F to epigallocatechin-3-*O*-gallate, procyanidin B_2, or procyanidin

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								B_1 appear below procyanidin A_2. to: The *Sample solution* exhibits two faint brown bands corresponding in R_F to procyanidin A2 and epicatechin in *Standard solution B*. The *Sample solution* also exhibits a series of faint or indistinct brown bands of differing intensities in the lower-half section, corresponding to the same bands in *Standard solution C*. No bands corresponding in *R*

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								Fto epigallocatechin-3-*O*-gallate, procyanidin B2, or procyanidin B1 appear below procyanidin A2. AND In *C./Acceptance criteria/Profile at 520 nm*: Change cyanidin-3-*O*-arabinose, to: cyanidin-3-*O*-arabinoside, AND In *C./Acceptance criteria/Profile at 520 nm*: Change peonidin-3-*O*-arabinose to:

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								peon idin-3-*O* -arabinoside
ZIPRASIDONE CAPSULES	ASSAY	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Proce dure*/*Analysis*: Change 449.40 to: 449.39
AZITHROMYCIN	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Organic Impurities*/*Table 2*: Change: 3?-*N* -{[4-(Acetylamino)phenyl]sulfonyl}-3?-demethylazithromycin[m] to: 3'-*N* -D emet hyl-3'-*N* -[(4-methylphenyl)sulfonyl]azithromycin[m] AND In *Organic Impurities*/*Table 2*/footnote m: Change

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								(2*R*,3*S*,4*R*,5*R*,8*R*,10*R*,11*R*,12*S*,13*S*,14*R*)-13-[(2,6-Dideoxy-3-*C*-methyl-3-*O*-methyl-?-L-*ribo*-hexopyranosyl)oxy]-2-ethyl-3,4,10-trihydroxy-3,5,6,8,10,12,14-heptamethyl-11-[[3-[*N*-(4-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-Dideo

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								xy-3-*C*-methyl-3-*O*-methyl-?-L-*ribo*-hexopyranosyl)oxy]-2-ethyl-3,4,10-trihydroxy-3,5,6,8,10,12,14-heptamethyl-11-[[3-[*N*-(4-methylphenylsulfonyl)-*N*-methylamino]-3,4,6-trideoxy-ß-D-*xylo*-hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one.
DEXTROMETH ORPHAN HYDROBROMIDE ORAL SOLUTION	*Assay*	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Chromatographic system* and *Procedure*: Change *C* is the concentration, in mg per mL, of USP Dextromethorphan Hydrobromide

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SODIUM ALGINATE	ASSAY	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	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*; In *Procedure/Analysis*: Change Result = $(V_2 \times N \times W_E)/(W \times D)$ to: Result = $(V_2 \times N \times W_E \times 10)/(W \times D)$
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

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								water. Add 133 mL of glacial acetic acid to this solution, and then dilute with water to 10 L; 900 mL. to: 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.
ZIPRASIDONE HYDROCHLORIDE IMPURITIES	*USPNF Online*	Online		25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Organic Impurities*/*Solution B*: Change Acetonitrile, methanol, and *Buffer* (55:5:40). Adjust with

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								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 In both equations in *Organic Impuri ties*/*Analysis*: Change 449.40 to: 449.39
REAGENTS AND REFERENCE TABLES	*Solutions*/*Test Solutions and Indicator Solutio ns*/*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

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								procedures. Apply the *Test for sensitivity* to confirm suitability for freshly or previously prepared solutions or commercially bought solutions. 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

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								prepared solutions. AND In *Test for sensitivity*: Delete Use the *Test for sensitivity* to confirm suitability for use. 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
BLACK CUMIN SEED THYMO QUINONE OIL	DEFINITION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change carvacol to: carvacrol
CRANBERRY	COMPOSITION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Content of Pr*

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FRUIT DRY JUICE								
								/*Standard stock solution*: Change USP Procyanidin A_2 RS to: USP Procyanidin A2 RS AND In *Content of Pr*
								/*Analysis*: Change Use the absorbance recorded for *Standard solutions 1–5* to obtain a calibration curve (absorbance vs. concentration, in µg/mL, of

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								procyanidin A_2) and perform a linear regression analysis. Determine the concentration (*C*), in µg/mL, of total proanthocyanidins as procyanidin A_2 in the *Sample solution*.
								to: Use the absorbance recorded for *Standard solutions 1–5* to obtain a calibration curve (absorbance vs. concentration, in µg/mL, of procyanidin A2) and perform a linear regression analysis. Determine the

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								concentration (*C*), in µg/mL, of total proanthocyanidins as procyanidin A2 in the *Sample solution*. AND In *Content of Pr* */Analysis*: Change *C* = concentration of the *Sample solution* as procyanidin A_2 from the regression line (µg/mL) to: *C* = concentration of the *Sample solution* as procyanidin A2 from the regression line (µg/mL)

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ZIPRASIDONE CAPSULES	PERFORMANCE TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	AND In *Content of Pr* */Acceptance criteria*: Change procyanidin A_2 to: procyanidin A2 In *Dissolution* ?711?/*Test 1*/*Tier 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

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								AND In *Dissolution* ?711?/*Test 1*/*Tier 2*/*Solution A* and *Solution B*: Change sodium hydroxide to: sodium hydroxide 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

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AZITHROMYCIN	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *USP Reference Standards* ?11?/USP Azaerythromycin A RS: Change 734.96 to: 734.97 AND In USP Azithromycin Related Compound F RS: Change 762.97 to: 762.98 AND In USP Desosaminylazithromycin RS: Change 590.79 to: 590.80
AMANTADINE HYDROCHLORIDE	IDENTIFICATION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *A.*: Change *Spectroscopic Identification Tests* ?197?, *Infrared Spectroscopy*:

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								197A, 197K, and 197S to: *Spectroscopic Identification Tests* ?197?, *Infrared Spectroscopy*: 197A, 197K, or 197S **Procedure for 197S**
CRANBERRY FRUIT DRY JUICE	DEFINITION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change procyanidin A_2 to: procyanidin A2
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%.
AZITHROMYCIN	CHEMICAL INFORMATION	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change 748.98

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								to: 749.00 AND Change 767.00 to: 767.01 AND Change 785.02 to: 785.03
TOPICAL AND SPECIFIC TRANSDERMA TESTS FOR L DRUG PROD TDS UCTS—PRODU CT QUALITY TESTS		*USPNF Online*	Online	28-Jul-2023	1-Dec-2023	*NA*	*NA*	In *Release Liner Peel Test*: Change The product fails the test if the mean peel force is outside the acceptable range determined during product development. to: The product fails the test if the overall mean peel force is outside the acceptable

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								range determined during product development.
PIPERACILLIN	SPECIFIC TESTS	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	In *Sterility Tests* ?71?: Change Where the label states that Piperacillin is sterile or that it must be subjected to further processing during the preparation of injectable dosage forms, it meets the requirements when tested as directed in *Test for Sterility of the Product to Be Examined*, *Membrane Filtration*. to: Where the label states that Piperacillin is

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								sterile, it meets the requirements when tested as directed in *Test for Sterility of the Product to Be Examined*, *Membrane Filtration*.
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 764.98 to: 765.00 AND In USP *N*-Demethylazithromycin RS:

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								Change 734.96 to: 734.97 AND In USP Desosaminylazithromycin RS: Change 590.79 to: 590.80
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
TRIMETHOBENZAMIDE HYDROCHLORIDE	CHEMICAL INFORMATION	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	Change *N*-[*p*-[2-(Dimethylamino)ethoxy]benzyl]-3,4,5-trimethoxy benzamide monohydrochloride to: *N*-[4-[2-(Dimethylamino)ethoxy]benzyl]-3,4,5-trimethoxybenzami

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								de monohydrochloride
PURIFIED SILICEOUS EARTH	IMPURITIES	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	In *Leachable Arsenic/ Analysis*: Change A 3.0-mL portion of the *Sample solution* meets the requirements in *Arsenic* ?211?, *Procedures*, *Procedure*. to: A 3.0-mL portion of the *Sample solution* meets the requirements in *Arsenic* ?211?, *Procedures*, *Procedure 1*.
CLOMIPHENE CITRATE TABLETS	IM PUR ITIES/*Organic Impurities*	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	In *Table 2*: Change Clomiphene related compound A 0.87 – 2.0 to: Clomiphene

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								related compound A 0.87 1.0 2.0
CHLORTETRACYCLINE HYDROCHLORIDE	CHEMICAL INFORMATION	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	Change 7-Chloro-4-(dimethylamino)-1,4,4a,5,5a,6,11,12a-octahydro-3,6,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-2-naphthacenecarboxamide monohydrochloride to: (4*S*,4a*S*,5a*S*,6*S*,12a*S*)-7-Chloro-4-(dimethylamino)-3,6,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-1,4,4a,5,5a,6,11,12a-octahydrotetracene-2-carboxamide monohydrochloride
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*

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								,8*R*,10*R*,11*R* ,12*S*,13*S*,14*R*)-13-[(2,6-Dideoxy-3-*C* -methyl-3-*O*-methyl-?-L-*ribo* -hexopyranosyl)oxy]-2-ethyl-3,4,10-trihydroxy-3,5,6,8,10,12,14-heptamethyl-11-[[3-[*N* -(4-acetamidophen ylsulfonyl)-*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*

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								-meth yl-3-*O*-methyl-?-L-*ribo* -hexopyranosyl) oxy]-2-ethyl-3,4, 10-trihydroxy-3, 5,6,8,10,12,14-heptamethyl-11-[[3-[*N* -(4-methylpheny lsulfon yl)-*N* -methylamino]-3 ,4,6-trideoxy-ß-D-*xylo* -hexopyranosyl] oxy]-1-oxa-6-az acyclopentadec an-15-one.
REAGENTS AND REFERENCE TABLES	*Reagent S pec ificatio ns/ Tosylchloramid e Sodium*	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	Change 127-65-1 to: 7080-50-4
POLYVINYL ALCOHOL	IMPURITIES	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	In *Limit of Methanol (Methyl Alcohol) and Methyl Acetate/*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Analysis: Change C_U = concentration of methanol (methyl alcohol) or methyl acetate in the *Sample solution* (mg/mL) to: C_U = concentration of Polyvinyl Alcohol in the *Sample solution* (mg/mL)
SITAGLIPTIN PHOSPHATE	IM PURITIES/*Organic Impurities*	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	In *Analysis*: Change C_S = concentration of USP Sitagliptin Phosphate in the *Standard solution* (mg/mL) to: C_S = concentration of USP Sitagliptin Phosphate RS

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								in the *Standard solution* (mg/mL)
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

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								$^{16}H_{12}BrCl_3N_2OS \cdot HCl$ 503.06
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
CALCIUM ASCORBATE	ASSAY/ *Procedure*	*USPNF Online*	Online	30-Jun-2023	1-Aug-2023	*NA*	*NA*	In *Analysis*: Change M_{r1} = molecular weight of calcium ascorbate dihydrate, 426.43 to: M_{r1} = molecular weight of calcium ascorbate dihydrate, 426.34
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	Page Number	Errata Post Date Sort ascending	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
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: USP Methacrylic Acid and Ethyl

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DIGOXIN TABLETS	ASSAY/ *Procedure*	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	Acrylate Copolymer (1:1) RS (USP Methacrylic Acid Copolymer Type C RS) In *Chromatographic system*: Change **Column:** 4.2-mm × 25-cm; 5-?m packing L1 to: **Column:** 4.6-mm × 25-cm; 5-µm packing L1
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 ($C_7H_{10}NNaO_7P_7$)

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								dissolved: to: Calculate the percentage of the labeled amount of risedronate sodium ($C_7H_{10}NNaO_7P_2$) dissolved:

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