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TERAZOSIN	ASSAY/	*USPNF Online*	Online	30-Sep-2022	1-Oct-2022	*NA*	*NA*	In *Hydrochloric*

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CAPSULES	*Procedure*							*acid solution*: Change 0.1 N methanolic hydrochloric acid to: 0.01 N methanolic hydrochloric acid
TETRACYCLINE HYDROCHLORIDE	SPECIFIC TESTS/ *Bacterial Endotoxins Test <85>*	*USPNF Online*	Online	28-Oct-2022	1-Nov-2022	*NA*	*NA*	Change Where the label states 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

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								form monograph(s) in which tetracycline hydrochloride is used can be met. to: Where the label states 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

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								Hydrochloride is used can be met.
TETRACYCLINE HYDROCHLORIDE	ADDITIONAL REQUIREMENTS/*Labeling*	*USPNF Online*	Online	28-Oct-2022	1-Nov-2022	*NA*	*NA*	Change Where tetracycline hydrochloride must be sterile or subjected to further processing during the preparation of injectable dosage forms to ensure acceptable levels of bacterial endotoxins, it is so labeled. to: Where Tetracycline Hydrochloride must be sterile or subjected to further processing during the preparation of

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								injectable dosage forms to ensure acceptable levels of bacterial endotoxins, it is so labeled.
THALIDOMIDE CAPSULES	ASSAY	*USPNF Online*	Online	23-Feb-2024	1-Mar-2024	*NA*	*NA*	In *Procedure*: Change $1000C(R_U/R_S)$ to: $500C(R_U/R_S)$
THEOPHYLLINE CAPSULES	IDENTIFICATION	*USPNF Online*	Online	26-Jan-2024	1-Feb-2024	*NA*	*NA*	Change **A:** The contents of the Capsules respond to *Identification* tests *A* and *B* under *Theophylline Tablets*. **B:** The retention time of the major peak in the chromatogram of the *Assay preparation* corresponds to

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								that in the chromatogram of the *Standard preparation*, as obtained in the *Assay*. to: **A:** Triturate a quantity of the contents of Capsules, equivalent to about 500 mg of theophylline, with 10-mL and 5-mL portions of solvent hexane, and discard the solvent hexane. Triturate the residue with two 10-mL portions of a mixture of equal volumes of 6 N ammonium hydroxide and water, and filter each time. Evaporate the combined

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								filtrates to about 5 mL, neutralize, if necessary, with 6 N acetic acid, using litmus, and then cool to about 15°, with stirring. Collect the precipitate on a filter, wash it with cold water, and dry at 105° for 2 hours: the theophylline so obtained melts between 270° and 274° (see *Melting Range or Temperature* ?741?, *Procedures*, *Procedure for Class I*). Retain the remaining portion of the theophylline for use in *Identification* test *B*.

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								B: The IR absorption spectrum of a potassium bromide dispersion of the residue obtained in *Identification* test *A* exhibits maxima only at the same wavenumbers as that of a potassium bromide dispersion of USP Theophylline RS. **C:** The retention time of the major peak in the chromatogram of the *Assay preparation* corresponds to that in the chromatogram

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								of the *Standard preparation*, as obtained in the *Assay*.
THYROID	ASSAY/ *Procedure*	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	In *Chromatographic system*: Change **Column**: 4.6-cm × 25-cm; packing L1 to: **Column**: 4.6-mm × 25-cm; packing L1
THYROID TABLETS	ASSAY/ *Procedure*	*USPNF Online*	Online	31-Mar-2023	1-Apr-2023	*NA*	*NA*	In *Chromatographic system*: Change **Column:** 4.6-cm × 25-cm; packing L1 to: **Column:** 4.6-mm × 25-cm; packing L1
TIOCONAZOLE	ADDITIONAL REQUIREMENT	*USPNF Online*	Online	30-Jun-2023	1-Jul-2023	*NA*	*NA*	In USP Tioconazole

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	S/*USP Reference Standards* ?11?							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
TRIAZOLAM TABLETS	PERFORMANCE TESTS/	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	In *Standard solution*: Change

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	Dissolution ?711?							Tablet/mg to: mg/Tablet
TRIETHYL CITRATE	ASSAY/ *Procedure*	*USPNF Online*	Online	24-Feb-2023	1-May-2023	*NA*	*NA*	In *System suitability*/*Suitability requirements*/*Tailing factor*: Change NMT 1.5 for the triethyl citrate to trimethyl citrate peaks to: NMT 1.5 for the triethyl citrate and trimethyl citrate peaks
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]b

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								enzyl]-3,4,5-trimethoxybenzamide monohydrochloride
TRYPTOPHAN	ADDITIONAL REQUIREMENTS/*USP Reference Standards ?11?*	*USPNF Online*	Online	24-Jun-2022	1-Jul-2022	*NA*	*NA*	In USP Tryptophan Related Compound A RS: Change 3,3?-[Ethylidenebis(1*H*-indole-1,3-diyl)]bis[2*S*)-2-aminopropanoic]acid. $C_{24}H_{26}N_4O_4$ 432.49 to: (2*S*,2'*S*)-3,3'-[Ethane-1,1-diyl bis(1*H*-indole-1,3-diyl)]bis(2-aminopropanoic acid). $C_{24}H_{26}N_4O_4$ 434.50
UREA	SPECIFIC TESTS	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Alcohol-Insoluble Matter*/*Sample solution*:

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								Change 100 mg/mL of Urea dissolved in warm alcohol to: Dissolve 5.0 g of Urea in 50 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

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								residue remains, pass the *Sample solution* through a tared filter, wash the residue and the filter with 20 mL of warm alcohol, and dry at 105° for 1 h.
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 to: **Flow rate**: 1 mL/min
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-hydroxyp

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								ropyl methyl-L-valinate hydrochloride. to: 3-[(2-Amino-6-oxo-1,6-dihydropurin-9-yl)methoxy]-2-hydroxypropyl L-valinate hydrochloride.
VIGABATRIN TABLETS	IM PUR ITIES/*Organic Impurities*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	In *Analysis*: Change C_S = concentration of USP Vigabatrin Related Compound A in the *Standard solution* to: C_S = concentration of USP Vigabatrin Related Compound A RS in the *Standard solution*
ZINC UNDECYLENATE	ASSAY/ *Procedure*	*USPNF Online*	Online	18-Nov-2022	1-Dec-2022	*NA*	*NA*	In *Solution A*: Change 0.15 N

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								hydrochloric acid in water prepared as follows. Transfer 150 mL of hydrochloric acid to a 500-mL volumetric flask, dilute with water to volume, and mix well. to: 0.15 N hydrochloric acid in water prepared as follows. Transfer 150 mL of 0.5 N hydrochloric acid to a 500-mL volumetric flask, dilute with water to volume, and mix well.
ZIPRASIDONE CAPSULES	ASSAY	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Procedure/Analysis*:

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ZIPRASIDONE CAPSULES	PERFORMANCE TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change 449.40 to: 449.39 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 AND In *Dissolution* ?711?/*Test 1*/*Tier 2*/*Solution A* and *Solution B*:

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ZIPRASIDONE CAPSULES	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	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 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
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 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

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								of 6.0. AND In both equations in *Organic Impurities/Analysis*: Change 449.40 to: 449.39

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