CSV October 4, 2022 through October 4, 2024 See all Errata for USP–NF How to use

×

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 Aminosalicylic Acid Tablets will result in anything that contains “Aminosalicyclic” OR “Acid” OR “Tablets”
    - A search for “Aminosalicylic Acid Tablets” will result in anything that specifically contains “Aminosalicylic 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.
    - For example, page 2178 will come before page 74 on a page sort.
- **Downloading:** You can download the Errata table in Comma-separated Value (.csv). The download will include the Errata that you have filtered on.
- **Importing:**You will need to import the file into Excel or Open Office with UTF-8 encoding, as opposed to simply opening it. To import, open Excel or Open Office and select import from the File drop-down. Depending on the version you are using, you should be presented with import formatting options to include UTF-8 as one of the first steps. Importing via UTF-8 should eliminate odd character conversions.

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
RALBUMIN	IMPURITIES	*USPNF Online*	Online	27-Sep-2024	1-Oct-2024	*NA*	*NA*	In *Limit of High*

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
HUMAN								*Molecular Weight Proteins*: Change **Acceptance criteria Individual impurities:** NMT 1.0% to: **Acceptance criteria:** NMT 1.0%
REAGENTS AND REFERENCE TABLES	*Solutions/Test Solutions and Indicator Solutions/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 procedures. Apply the *Test for sensitivity* to confirm suitability for freshly or previously prepared

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 prepared solutions. AND In *Test for sensitivity*: Delete Use the *Test for sensitivity* to

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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
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
RISEDRONATE SODIUM DELA YED-RELEASE TABLETS	PERFORMANC E 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

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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:
RISEDRONATE SODIUM DELAYED-RELEASE TABLETS	ASSAY	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	Delete [Note—Use a non-metallic liquid chromatography system for analysis.]
RIVASTIGMINE TARTRATE	IMPURITIES	*USPNF Online*	Online	26-Apr-2024	1-May-2024	*NA*	*NA*	In *Organic Impurities/Procedure 1/Impurity Table 1*/footnote d: Change 3-Nitrophenyl ethyl(methyl)carb

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								amate. to: 4-Nitrophenyl ethyl(methyl)carbamate.
RIVASTIGMINE TARTRATE	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	26-Apr-2024	1-May-2024	*NA*	*NA*	In *USP Reference Standards* ?11?: Change USP Rivastigmine Tartrate R-Isomer RS to: USP Rivastigmine Tartrate R-Isomer RS (*R*)-3-[1-(Dimethylamino)ethyl]phenyl ethylmethylcarbamate, hydrogen tartrate. $C_{14}H_{22}N_2O_2 \cdot C_4H_6O_6$ 400.42
ROCURONIUM BROMIDE	CHEMICAL INFORMATION	*USPNF Online*	Online	18-Nov-2022	1-Dec-2022	*NA*	*USPNF 2023 Issue 3*	Change 609.68 to: 609.69

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
ROCURONIUM BROMIDE	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	18-Nov-2022	1-Dec-2022	*NA*	*USPNF 2023 Issue 3*	In footnote b of *Table 1*: Change 2?-(Morpholin-4-yl)-16?-(pyrrolidin-1-yl)-5?-androstan-3?,17?-diol. to: 2?-(Morpholin-4-yl)-16?-(pyrrolidin-1-yl)-5?-androstane-3?,17?-diol.
ROSEMARY	COMPOSITION /*Content of Phenolic Diterpenes*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	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,
ROSEMARY LEAF DRY AQUEOUS	COMPOSITION /*Content of Rosmarinic Acid*	*USPNF Online*	Online	28-Apr-2023	1-May-2023	*NA*	*NA*	In *Standard solution B*: Change

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
EXTRACT								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, AND 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, Change Where the label states that Seco barbital Sodium is sterile, it
SECOBARBITAL SODIUM	OTHER REQUIREMENTS	*USPNF Online*	Online	29-Dec-2023	1-Jan-2024	*NA*	*NA*	

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 endotoxins* under *Secobarbital Sodium for Injection*. to: Where the label

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 Sodium must be subjected to further processing during the preparation of injectable dosage forms,

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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>.
SITAGLIPTIN PHOSPHATE	CHEMICAL INFORMATION	*USPNF Online*	Online	28-Oct-2022	1-Nov-2022	*NA*	*NA*	Change 523.32 to: 523.33 AND Change (3*R*)-3-Amino-1-[3-(trifluoromethyl)-5,6-dihydro[1,2,4]triazolo[4,3-*a*]pyrazin-7(8*H*)-yl-4-(2,4,5-trifluorophenyl)butan-1-one phosphate

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								monohydrate to: (3*R*)-3-Amino-1-[3-(trifluoromethyl)-5,6-dihydro[1,2,4]triazolo[4,3-*a*]pyrazin-7(8*H*)-yl]-4-(2,4,5-trifluorophenyl)butan-1-one phosphate monohydrate
SITAGLIPTIN PHOSPHATE	IMPURITIES/*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 in the *Standard*

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								solution (mg/mL)
SODIUM ALGINATE	ASSAY	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Procedure/Analysis*: Change Result = $(V_2 \times N \times W_E)/(W \times D)$ to: Result = $(V_2 \times N \times W_E \times 10)/(W \times D)$
SODIUM SALICYLATE	ADDITIONAL REQUIREMENTS	*USPNF Online*	Online	17-Nov-2023	1-Dec-2023	*NA*	*NA*	In *USP Reference Standards* ?11?: Add USP Phenol RS
SOTALOL HYDROCHLORIDE	IMPURITIES/*Organic Impurities*	*USPNF Online*	Online	24-Feb-2023	1-Mar-2023	*NA*	*NA*	In *Analysis*: Change Calculate the percentage of any unspecified impurities in the portion of Sotalol Hydrochloride taken: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$ r_U= sum of all the peak

Monograph Title	Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								responses of all unspecified impurities from the *Sample solution* to: Calculate the percentage of any unspecified impurity in the portion of Sotalol Hydrochloride taken: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$ r_U = peak response of each unspecified impurity from the *Sample solution* AND In *Table 1*: Change Any unspecified impurities to: Any unspecified impurity

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
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.
SUCROSE	IMPURITIES	*USPNF Online*	Online	29-Mar-2024	1-Apr-2024	*NA*	*NA*	In *Sulfite/ Analysis*/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

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								sulfite determination may be ordered 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).
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

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

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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, *Standard solution* **Relative standard deviation:** NMT 5.0%, *Standard solution* **Signal-to-noise ratio:** NLT 10, *Sensitivity solution*
TELMISARTAN ASSAY TABLETS	*USPNF Online*	Online		27-Oct-2023	1-Nov-2023	*NA*	*NA*	In *Procedure*/*System suitability*/*Suitability requirements*:

Monograph Title	Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 telmisartan and telmisartan related compound A **Tailing factor:** NMT 2.0 for telmisartan **Relative standard deviation:** NMT

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
TELMISARTAN TABLETS	PERFORMANCE TESTS	*USPNF Online*	Online	27-Oct-2023	1-Nov-2023	*NA*	*NA*	2.0% for telmisartan In *Dissolution* ?711?/*Test 1*/*Analysis*: Change Determine the percentage of telmisartan ($C_{33}H_{30}N_4O_2$) dissolved: Result = (A_U × C_S × V × 100)/(A_S × D × L) to: Calculate the percentage of the labeled amount of telmisartan ($C_{33}H_{30}N_4O_2$) dissolved: Result = (A_U/A_S) × C_S × V × D × (1/L) × 100 AND Change C_S = concentration of the *Standard*

Monograph Title	Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								solution (mg/mL) to: C_S = concentration of USP Telmisartan RS in the *Standard solution* (mg/mL) AND In *Dissolution* ?711?/*Test 2*/*Analysis*: Change r_U = peak response from the *Sample solution* r_S = peak response from the *Standard solution* to: r_U = peak response of telmisartan from the *Sample solution* r_S = peak response of

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								telmisartan from the *Standard solution* AND In *Dissolution* ?711?/*Test 3*/*Analysis*: Change C_S = concentration of the *Standard solution* (mg/mL) to: C_S = concentration of USP Telmisartan RS in the *Standard solution* (mg/mL)
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

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 subjected to further processing during the preparation of injectable dosage forms, the level of

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								bacterial endotoxins are such that the requirement under the relevant dosage form monograph(s) in which Tetracycline 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.

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								to: 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.
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*

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Tablets. **B:** The retention time of the major peak in the chromatogram of the *Assay preparation* corresponds to 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

Monograph Title	Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								10-mL portions of a mixture of equal volumes of 6 N ammonium hydroxide and water, and filter each time. Evaporate the combined 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*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								?741?, *Procedures*, *Procedure for Class I*). Retain the remaining portion of the theophylline for use in *Identification* test *B*. **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:**

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								The retention time of the major peak in the chromatogram of the *Assay preparation* corresponds to that in the chromatogram 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

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Column: 4.6-cm × 25-cm; packing L1 to: **Column:** 4.6-mm × 25-cm; packing L1
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

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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/ *Dissolution* ?711?	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	In *Standard solution*: Change 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
TRIMETHOBE	CHEMICAL	*USPNF Online*	Online	28-Jul-2023	1-Aug-2023	*NA*	*NA*	Change

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
NZAMIDE HYDROCHLORIDE	INFORMATION							*N*-[*p*-[2-(Dimethylamino)ethoxy]benzyl]-3,4,5-trimethoxy benzamide monohydrochloride to: *N*-[4-[2-(Dimethylamino)ethoxy]benzyl]-3,4,5-trimethoxybenzamide monohydrochloride
TYLOSIN INJECTION	SPECIFIC TESTS	*USPNF Online*	Online	31-May-2024	1-Jun-2024	*NA*	*NA*	Change *pH* ?921? to: *pH* ?791?
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 mL of warm

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 filter with 20 mL

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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-hydroxypropyl methyl-L-valinate hydrochloride. to: 3-[(2-Amino-6-oxo-1,6-dihydropurin-9-yl)methoxy]-2-hydroxypro

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								pyl L-valinate hydrochloride.
VIGABATRIN TABLETS	IMPURITIES/*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 hydrochloric acid in water prepared as follows. Transfer 150 mL of hydrochloric acid to a

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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	PERFORMANCE TESTS	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Dissolution* ?711?/*Test 1*/*Tier 1*/*Phosphate buffer, pH 7.5*: Change sodium hydroxide to: sodium hydroxide

Monograph Title	Section Sort descending	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								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 solution AND In *Dissolution* ?711?/*Test 2*/*Tier 2*/*Analysis*: Change 449.40 to:

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								449.39 AND In *Dissolution* ?711?/*Test 3*/*Tier 2*/*Analysis*: Change 449.40 to: 449.39
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
ZIPRASIDONE CAPSULES	ASSAY	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	In *Procedure*/*Analysis*:

Monograph Title Sort descending	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
ZIPRASIDONE HYDROCHLORIDE	IMPURITIES	*USPNF Online*	Online	25-Aug-2023	1-Sep-2023	*NA*	*NA*	Change 449.40 to: 449.39 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 In both equations in *Organic Impurities/Analysis*: Change

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
Sort descending								449.40 to: 449.39

Pagination

- First page « First
- Previous page ‹ Previous
- Page 1
- Page 2
- Page 3
- Page 4
- Page 5