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RIVASTIGMINE IMPURITIES TARTRATE		USPNF Online	Online	26-Apr-2024	1-May-2024	NA	NA	In <i>Organic Impurities/Procedure 1/Impurity Table 1/footnote d</i> : Change 3-Nitrophenyl ethyl(methyl)carbamate. to: 4-Nitrophenyl ethyl(methyl)carbamate.
MELOXICAM ORAL SUSPENSION	PERFORMANCE TESTS/ <i>Dissolution</i> <711>	USPNF Online	Online	29-Jul-2022	1-Aug-2022	NA	NA	In <i>Analysis</i> : Change W_U = weight of the Oral Suspension taken (mg) to: W_U = weight of the Oral Suspension taken (g) Change (3 <i>R</i> ,3?S-meso)-zeaxanthin to: (3 <i>R</i> ,3?S
MESO-ZEAXANTHIN	IDENTIFICATION/C.	USPNF Online	Online	27-Jan-2023	1-Jun-2023	NA	NA	

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SITAGLIPTIN PHOSPHATE	IM PURITIES/ <i>Organic Impurities</i>	<i>USPNF Online</i>	Online	30-Jun-2023	1-Jul-2023	NA	NA	)-zeaxanthin In <i>Analysis</i> : Change C_S = concentration of USP Sitagliptin Phosphate in the <i>Standard solution</i> (mg/mL) to: C_S = concentration of USP Sitagliptin Phosphate RS in the <i>Standard solution</i> (mg/mL)
NORFLURANE	IMPURITIES	<i>USPNF Online</i>	Online	17-Nov-2023	1-Dec-2023	NA	NA	In <i>Organic Impurities/</i> Table 2: Change Line No. to: Peak Elution Order AND In <i>Halides/Figure 1</i> : Add label Flow Meter

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PROCHLORPE RAZINE MALEATE TABLETS	PERFORMANCE TESTS/ <i>Dissolution</i> <711>	USPNF Online	Online	26-Aug-2022	1-Sep-2022	NA	NA		In <i>Analysis</i>: Change Results = $(A_U/A_S) \times C_S \times D \times (1/L) \times V \times 100$ $A_U =$ absorbance of the <i>Sample solution</i> $A_S =$ absorbance of the <i>Standard solution</i> $C_S =$ concentration of USP Prochlorpe razine Maleate RS in the <i>Standard solution</i> <i>D</i> = dilution factor for <i>Sample solution</i>, if needed <i>L</i> = label claim (mg/Tablet) <i>V</i> = volume of <i>Medium</i>, 500

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							mL to: $\text{Result} = (A_U/A_S) \times C_S \times V \times D \times (M_{r1}/M_{r2}) \times (1/L) \times 100$ $A_U =$ absorbance of the <i>Sample solution</i> $A_S =$ absorbance of the <i>Standard solution</i> $C_S =$ concentration of USP Prochlorperazine Maleate RS in the <i>Standard solution</i> $V =$ volume of <i>Medium</i> , 500 mL $D =$ dilution factor for the <i>Sample solution</i> , if needed M

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Methyl Acetate	CHEMICAL INFORMATION	USPNF Online	Online	27-Jan-2023	1-Feb-2023	NA	NA	r_1 = molecular weight of prochlorperazine, 373.94 M_{r2} = molecular weight of prochlorperazine maleate, 606.09 <i>L</i> = label claim (mg/Tablet) Change CAS RN®: 74-20-9. to: CAS RN®: 79-20-9. In <i>Leachable Arsenic/ Analysis:</i> Change A 3.0-mL portion of the <i>Sample solution</i> meets the requirements in <i>Arsenic ?211?</i> , <i>Procedures, Procedure.</i> to: A 3.0-mL
PURIFIED SILICEOUS EARTH	IMPURITIES	USPNF Online	Online	28-Jul-2023	1-Aug-2023	NA	NA	

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GLUCAGON	PROCESS-RELATED IMPURITIES AND OTHER COMPONENTS	USPNF Online	Online	29-Dec-2023	1-Jan-2024	NA	NA	portion of the <i>Sample solution</i> meets the requirements in <i>Arsenic ?211?</i>, <i>Procedures, Procedure 1.</i> In <i>Acetic Acid in Peptides/Analysis</i>: Change C_{SPA} = concentration of potassium acetate in each of the <i>Standard solutions</i> (mg/mL) to: C_{SPA} = concentration of potassium acetate in each of the <i>Standard solutions</i> (µg/mL) AND In <i>Ammonium/</i>

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DOXYCYCLINE CAPSULES	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USPNF Online	Online	24-Jun-2022	1-Jul-2022	NA	NA		<i>Analysis:</i> Change C_{SAC} = concentration of ammonium chloride in each of the <i>Standard solutions</i> (mg/mL) to: C_{SAC} = concentration of ammonium chloride in each of the <i>Standard solutions</i> (µg/mL) In USP Doxycycline Related Compound A RS: Change 444.43 to: 444.44 AND Change (4S,4aR,5S, 5aR,6S, 12aS

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ANETHOLE	DEFINITION	USPNF Online	Online	28-Oct-2022	1-Nov-2022	NA	NA		)-4-(Dimethylamino)-1,4,4a,5,5a,6,11,12a-octahydro-3,5,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-2-naphthacenenecarboxamide monohydrochloride. $C_{22}H_{24}N_2O_8 \cdot HCl$ 480.13 to: (4S,4aR,5S,5aR,6S,12aS)-4-(Dimethylamino)-1,4,4a,5,5a,6,11,12a-octahydro-3,5,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-2-naphthacenenecarboxamide hydrochloride. $C_{22}H_{24}N_2O_8 \cdot HCl$ 480.90 Change (E

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BISOPROLOL FUMARATE	CHEMICAL INFORMATION	USPNF Online	Online	28-Apr-2023	1-May-2023	NA	NA		)-1-methyl-4-(1-propenyl) benzene to: (E)-1-methoxy-4-(1-propenyl) benzene Change (±)-1-[[?-(2-Isop <i>p</i>-tolyl]oxy]-3-(iso propyl amino)-2-propanol fumarate (2:1) (salt) to: (±)-1-[[?-(2-Isop <i>p</i>-tolyl]oxy]-3-(iso propylamino)-2-propanol fumarate (2:1) (salt)

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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 ($\mu\text{g/mL}$) to: C_U = nominal concentration of digoxin in the Sample solution ($\mu\text{g/mL}$)
MUPIROCIN NASAL OINTMENT	Related compounds/Table 1	USPNF Online	Online	29-Apr-2022	1-May-2022	NA	NA	In footnote 2: Change 9-[(E)-4-[(2R,3aS,6S,7S,8aRS)-2-[(1RS,2S,3S)-1,3-Dihydroxy-2-methylbutyl]-7-hydroxyhexahydro-2H-furo[3,2-c]pyran-6-yl]-3-methylbut-2-enoyloxy}nonanoic

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								acid. to: 9- <i>{(E)</i> <i>)-4-[(2R,3aS,6S,7S)</i> <i>)-2-[(1RS,2S,3S)</i> <i>)-1,3-Dihydroxy-2-methylbutyl}-7-hydroxyhexahydro-2H-furo[3,2-c]</i> <i>]pyran-6-yl]-3-methylbut-2-enoyloxy}nonanoic acid.</i>
N-Benzoyl-L-arginine Ethyl Ester Hydrochloride	REAGENT SPECIFICATIONS	USPNF Online	Online	30-Sep-2022	1-Dec-2022	NA	NA	Change Crystallized Trypsin (USP Monograph). to: Trypsin (USP Monograph).
GLYCYL-L-TYROSINE	IMPURITIES/Related Compounds	USPNF Online	Online	24-Feb-2023	1-Jun-2023	NA	NA	In <i>Buffer solution</i> : Change Dissolve 6.84 g of potassium

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UREA C 13	IMPURITIES	USPNF Online	Online	25-Aug-2023	1-Sep-2023	NA	NA	phosphate in 1000 mL of water. to: Dissolve 6.84 g of monobasic potassium phosphate in 1000 mL of water. In <i>Isotopic Purity/Chromatographic system</i> : Change Flow rate : Flow rate to: Flow rate : 1 mL/min
METHYLNALT REXONE BROMIDE	CHEMICAL INFORMATION	USPNF Online	Online	23-Feb-2024	1-Mar-2024	NA	NA	Correct the chemical structure
AMMONIUM G LYCYRRHIZATE	CHEMICAL INFORMATION	USPNF Online	Online	24-Jun-2022	1-Jul-2022	NA	NA	Change 840.08 to: 839.97
PSEUDOEPHEDRINE HYDROCHLORIDE	PERFORMANCE TESTS	USPNF Online	Online	28-Apr-2023	1-May-2023	NA	NA	Change Uniformity of

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CHLORIDE TABLETS									Dosage Units ?905? Procedure for content uniformity to: Uniformity of Dosage Units ?905? : Meets the requirements Procedure for content uniformity, chewable tablets only AND In <i>Uniformity of Dosage Units</i> : Delete Acceptance criteria : Meet the requirements for chewable tablets In <i>Dissolution ?711?/Test 1/Analysis</i> :
TELMISARTAN TABLETS	PERFORMANCE TESTS	USPNF Online	Online	27-Oct-2023	1-Nov-2023	NA	NA		

Monograph Title Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication _Sort descending_	Description
							Change Determine the percentage of telmisartan ($C_{33}H_{30}N_4O_2$) dissolved: $\text{Result} = (A_U \times C_S \times V \times 100) / (A_S \times D \times L)$ to: Calculate the percentage of the labeled amount of telmisartan ($C_{33}H_{30}N_4O_2$) dissolved: $\text{Result} = (A_U / A_S) \times C_S \times V \times D \times (1/L) \times 100$ AND Change $C_S =$ concentration of the <i>Standard solution</i> (mg/mL) to: C

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

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APREPITANT CAPSULES	PERFORMANCE TESTS/ <i>Dissolution</i> <711>	USPNF Online	Online	27-May-2022	1-Jun-2022	NA	NA		<i>solution</i> AND In <i>Dissolution</i> ?711?/Test 3/Analysis: Change C_S = concentration of the <i>Standard</i> <i>solution</i> (mg/mL) to: C_S = concentration of USP Telmisartan RS in the <i>Standard</i> <i>solution</i> (mg/mL) Change Test 1 Dilute 1 mL of phosphoric acid with water to 1 L. to: Test 1 AND Change

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DOXORUBICIN ASSAY/ HYDROCHLORIDE FOR INJECTION	USPNF Online	Online	28-Oct-2022	1-Nov-2022	NA	NA	Dilute phosphoric acid: to: Dilute phosphoric acid: Dilute 1 mL of phosphoric acid with water to 1 L. In <i>Analysis</i>: Change <i>P</i> = potency of doxorubicin in USP Doxorubicin Hydrochloride RS (µg/mg) to: <i>P</i> = potency of doxorubicin hydrochloride in USP Doxorubicin Hydrochloride RS (µg/mg) Change Calcium lactate
CALCIUM LACTATE	CHEMICAL INFORMATION	USPNF Online	Online	31-Mar-2023	1-Apr-2023	NA	NA

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FOSAMPRENA PERFORMANCE TESTS VIR CALCIUM TABLETS	USPNF Online	Online	25-Aug-2023	1-Sep-2023	NA	NA	(1:2) hydrate to: Calcium lactate hydrate AND Change Calcium lactate (1:2) pentahydrate to: Calcium lactate pentahydrate In <i>Dissolution</i> ? 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: 0.02 M sodium acetate buffer, pH 3.5, prepared as

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RIVASTIGMINE TARTRATE	ADDITIONAL REQUIREMENTS	USP NF Online	Online	26-Apr-2024	1-May-2024	NA	NA		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. In <i>USP Reference Standards</i> 1171: Change USP Rivastigmine Tartrate R-Isomer RS to: USP Rivastigmine Tartrate R-Isomer RS (<i>R</i>)-3-[1-(Dimethyl amino)ethyl]phenyl ethylmethylcarbamate, hydrogen

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SUTURES--NE PROCEDURE EDLE ATTACHMENT		USPNF Online	Online	29-Jul-2022	1-Aug-2022	NA	NA	tartrate. C ₁₄ H ₂₂ N ₂ O ₂ · C ₄ H ₆ O ₆ 400.42 In Removable Needle Attachment: Change For USP sizes 5-0 through 2-0, to: For USP sizes 5-0 through 2, In System suitability: 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
MESO- ZEAXANTHIN	COMPOSITION /Stereoisomeric Composition	USPNF Online	Online	27-Jan-2023	1-Jun-2023	NA	NA	

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							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)-zeaxanthin, and (3R,3?R,6?R)-lutein are 0.94, 1.00, 1.06, and 1.11, respectively.] AND In System suitability/Suitability requirements/Resolution: Change

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ATORVASTATIN ASSAY/ N CALCIUM TABLETS	USPNF Online	Online	30-Jun-2023	1-Jul-2023	NA	NA	(3R,3?S-meso)-zeaxanthin to: (3R,3?S)-zeaxanthin AND In <i>Analysis</i> : Change (3R,3?S-meso)-zeaxanthin to: (3R,3?S)-zeaxanthin AND In <i>Acceptance criteria</i> : Change (3R,3?S-meso)-Zeaxanthin to: (3R,3?S)-Zeaxanthin In <i>Analysis</i> : Change M_{r1} = molecular weight of atorvastatin, 558.64

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									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
OLMESARTAN MEDOXOMIL TABLETS	PERFORMANCE TESTS	USPNF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA		In <i>Dissolution</i> ?711?/Test 5/Apparatus 2: Change 50 rpm. Use peak vessels. to: 50 rpm. Use apex vessels.
PROMETHAZINE HYDROCHLORIDE	IMPURITIES/Organic Impurities	USPNF Online	Online	26-Aug-2022	1-Sep-2022	NA	NA		Change System suitability solution : 5 µg/mL each of USP

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							Promethazine Hydrochloride RS and USP Promethazine Related Compound B RS from the <i>Standard stock solution</i> and <i>System suitability stock solution</i> , respectively Standard solution: 5 µg/mL of USP Promethazine Hydrochloride RS from the <i>Standard stock solution</i> Sensitivity solution: 0.25 µg/mL of USP Promethazine Hydrochloride RS from the <i>Standard solution</i>

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							to: System suitability solution: 5 µg/mL each of USP Promethazine Hydrochloride RS and USP Promethazine Related Compound B RS from the <i>Standard stock solution</i> and <i>System suitability stock solution</i> , respectively, in <i>Diluent</i> Standard solution: 5 µg/mL of USP Promethazine Hydrochloride RS from the <i>Standard stock solution</i> in <i>Diluent</i>

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METARAMINO L BITARTRATE	SPECIFIC TESTS/ <i>Content of Tartaric Acid</i>	USPNF Online	Online	27-Jan-2023	1-Feb-2023	NA	NA	Sensitivity solution: 0.25 µg/mL of USP Promethazine Hydrochloride RS from the <i>Standard solution</i> in <i>Diluent</i> In <i>Analysis</i> : Change Result = $\{(V_S \cdot V_B) \times N \times F\} / 2 \times W \times 100$ to: Result = $\{(V_S \cdot V_B) \times N \times F\} / (2 \times W) \times 100$ Change 127-65-1 to: 7080-50-4
REAGENTS AND REFERENCE TABLES	<i>Reagent Specificity/ Tosylchloramide Sodium</i>	USPNF Online	Online	28-Jul-2023	1-Aug-2023	NA	NA	Change 127-65-1 to: 7080-50-4
SECOBARBITAL SODIUM	OTHER REQUIREMENTS	USPNF Online	Online	29-Dec-2023	1-Jan-2024	NA	NA	Change Where the label states that Secobarbital Sodium

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							is sterile, it meets the requirements for <i>Sterility Tests</i> ?71? and for <i>Bacterial endotoxins</i> under <i>Secobarbital Sodium for Injection</i>. 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 <i>Bacterial endotoxins</i> under <i>Secobarbital Sodium for Injection</i>.

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							to: Where the label states that Secobarbital Sodium is sterile, it meets the requirements for <i>Sterility Tests</i> ?71? and the level of bacterial endotoxins is not more than 0.9 USP Endotoxin Units per mg of secobarbital sodium tested per <i>Bacterial Endotoxins Test</i> <85>. Where the label states that Secobarbital Sodium must be subjected to further processing during the

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DOXYCYCLINE FOR INJECTION	ADDITIONAL R EQUIREMENT S/USP Reference Standards <11>	USPNF Online Online	24-Jun-2022	1-Jul-2022	NA	NA	preparation of injectable dosage forms, the level of bacterial endotoxins is not more than 0.9 USP Endotoxin Units per mg of secobarbital sodium tested per <i>Bacterial Endotoxins Test</i> <85>. In USP Doxycycline Related Compound A RS: Change 444.43 to: 444.44 AND Change (4S,4aR,5S,5aR,6S,12aS)-4-(Dimethylamino)-1,4,4a,5,5a

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NALOXONE HYIMPURITIES DROCHLORIDE	USPNF Online	Online	28-Oct-2022	1-Dec-2022	NA	NA	,6,11,12a-octahydro-3,5,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-2-naphtacenecarboxamide, monohydrochloride. C ₂₂ H ₂₄ N ₂ O ₈ · HCl 480.13 to: (4S,4aR,5S,5aR,6S,12aS)-4-(Dimethylamino)-1,4,4a,5,5a,6,11,12a-octahydro-3,5,10,12,12a-pentahydroxy-6-methyl-1,11-dioxo-2-naphtacenecarboxamide hydrochloride. C ₂₂ H ₂₄ N ₂ O ₈ · HCl 480.90 In Limit of Naloxone Related Compound D:

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ISOFLURANE	IM PUR	USPNF Online	Online	28-Apr-2023	1-May-2023	NA	NA		Delete Sensitivity solution: 1.25 µg/mL of USP Naloxone Related Compound D RS in 0.1 N hydrochloric acid AND In System suitability/Samples: Delete Sensitivity solution, AND In System suitability/Suitability requirements: Delete Signal-to-noise ratio: NLT 10, Sensitivity solution In Analysis: Change

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		ITIES/Organic Impurities						Result = $(r_U/r_S) \times C_S \times (1/F)$ to: Result = $(r_U/r_S) \times C_F \times (1/F)$ AND Change C_S = final concentration of USP Isoflurane Related Compound B RS in the <i>Standard solution</i> (%) to: C_F = final concentration of USP Isoflurane Related Compound B RS in the <i>Standard solution</i> (%) In <i>Dissolution</i> ?711?/Test 8/Tolerances: Change The percentage of the labeled
DIVALPROEX SODIUM EXTENDED-RELEASE TABLETS	PERFORMANCE TESTS	USPNF Online	Online	29-Sep-2023	1-Oct-2023	NA	NA	

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								amount of valproic acid (C ₈ H ₁₆ O ₂) dissolved at the times specified conform to <i>Dissolution</i> <711>, <i>Acceptance Table 1</i> . to: The percentage of the labeled amount of valproic acid (C ₈ H ₁₆ O ₂) dissolved at the times specified conform to <i>Dissolution</i> <711>, <i>Acceptance Table 2</i> .
MUPIROCIN CALCIUM	CHEMICAL INFORMATION	USPNF Online	Online	29-Apr-2022	1-May-2022	NA	NA	Change 1075.34 to: 1075.35
LANSOPRAZOLE DELAYED-RELEASE	IM PUR ITIES/ <i>Organic</i>	USPNF Online	Online	30-Sep-2022	1-Oct-2022	NA	NA	Delete Blank: Methanol and

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CAPSULES ERYTHROMYCIN DELAYED- RELEASE TABLETS	<i>Impurities</i> PERFORMANCE TESTS/ <i>Dissolution</i> ?711?	USPNF Online Online	31-Mar-2023	1-Apr-2023	NA	NA	<i>Diluent</i> (1:9) In Test 1: Change Buffer stage Medium: 0.05 M pH 6.8 phosphate buffer (see <i>Reagents, Indicators, and Solutions—Buffer Solutions</i>) to: Buffer stage Medium: 0.05 M pH 6.8 phosphate buffer (see <i>Reagents, Indicators, and Solutions—Buffer Solutions</i>); 900 mL In <i>Chromatographic system and Procedure:</i> Change C is the
DEXTROMETHORPHAN HYDROBROMIDE ORAL SOLUTION	Assay	USPNF Online Online	25-Aug-2023	1-Sep-2023	NA	NA	

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METOLAZONE IMPURITIES	USPNF Online	Online	23-Feb-2024	1-Mar-2024	NA	NA	concentration, in mg per mL, of USP Dextromet horphan Hydrobromide RS, on the anhydrous basis, in the <i>Standard preparation</i> ; to: C is the concentration, in mg per mL, of USP Dextromet horphan Hydrobromide RS in the <i>Standard preparation</i> ; In <i>Organic Impurities/Chromatographic system/Column</i> : Change 4.6-mm × 25-cm; 5-μm packing 1

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							Sort descending	
AMMONIUM G LYCYRRHIZATE	ASSAY/Content of Ammonium 18?- and 18?- Glycyrrhizate	USPNF Online	Online	24-Jun-2022	1-Jul-2022	NA	NA	to: 4.6-mm × 25-cm; 5-μm packing L1 In Analysis: Change $M_{W(Salt)}$ = molecular weight of ammonium glycyrrhizate, 840.08 g/mol $M_{W(Acid)}$ = molecular weight of glycyrrhizic acid, 821.59 g/mol to: $M_{W(Salt)}$ = molecular weight of ammonium glycyrrhizate, 839.97 g/mol $M_{W(Acid)}$ = molecular weight of glycyrrhizic acid, 822.94

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication _Sort descending_	Description
MAFENIDE ACETATE FOR PUR TOPICAL SOLUTION	IM ITIES/ <i>Organic Impurities</i>	USPNF Online	Online	16-Dec-2022	1-Jan-2023	NA	NA	g/mol In System suitability: Change Samples: System suitability solution, Standard solution A, and Standard solution B to: Samples: System suitability solution and Standard solution A AND In Analysis: Change Samples: Mobile phase, Standard solution A, and Sample solution Chromatograph the Standard

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	_Sort descending_	Description
OSELTAMIVIR PHOSPHATE	IM PURITIES/Organic Impurity	USPNF Online	Online	26-May-2023	1-Jun-2023	NA	NA		<i>solution</i> and adjust the integration parameters so that the response is 5%–15% of full-scale deflection. to: Samples: <i>Mobile phase, Standard solution A, Standard solution B, and Sample solution</i> Chromatograph <i>Standard solution B</i> and adjust the integration parameters so that the response is 5%–15% of full-scale deflection. In System <i>suitability/Suitability</i>

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							Sort descending	
	ties/Procedure 3: Limit Of Tributyl Phosphine Oxide							quire ments/Relative standard deviation: Change NMT 10.0% for the tributyl phosphine oxide and osteltamivir phosphate peaks to: NMT 10.0% for the tributyl phosphine oxide and oseltamivir phosphate peaks
COLOR AND ACHROMICITY	METHOD II: IN STRUMENTAL (QUANTITATIV E) ASSESSMENT OF COLOR AND COLOR MATCHES	USPNF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	In Table 5: Change Sum 98.809 100.000 107.307 White point 98.811 100.000 107.304 to: Sum 94.809

Monograph Title Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication _Sort descending_	Description
PANTOPRAZO PERFORMANC LE SODIUM DEE TESTS LAYED- RELEASE TABLETS	USPNF Online	Online	17-Nov-2023	1-Dec-2023	NA	NA	100.000 107.307 White point 94.811 100.000 107.304 In Dissolution ?711?/Test 2/Acid stage: Change Acid stage standard solution: (L/10) mg/mL of USP Pantoprazole Sodium RS from the <i>Standard stock solution in Acid stage medium,</i> where L is the label claim in mg/Tablet to: Acid stage standard solution: (L/10000) mg/mL of USP Pantoprazole

Monograph Title Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication _Sort descending_	Description
							Sodium RS from the <i>Standard stock solution</i> in <i>Acid stage medium</i>, where <i>L</i> is the label claim in mg/Tablet AND In <i>Dissolution</i> <711>/Test 2/Buffer stage: Change Buffer stage standard solution: (<i>L</i>/1000) of USP Pantoprazole Sodium RS where <i>L</i> is the label claim in mg/Tablet to: Buffer stage standard solution: (<i>L</i>/1000) mg/mL of USP Pantoprazole

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication _Sort descending_	Description
BOTANICAL EXTRACTS	PREPARATION S	USPNF Online	Online	27-May-2022	1-Jun-2022	NA	NA	Sodium RS from the <i>Standard stock solution</i> in <i>Buffer stage medium</i> , where <i>L</i> is the label claim in mg/Tablet In <i>General Pharmacopeial R equ irement s/Pesticide Residues: Change</i> where <i>L</i> is the limit in the original article as listed in <i>Table 4</i> (see <i>Pesticide Residue Analysis under Articles of Botanical Origin <561></i>) to: where <i>L</i> is the

Monograph Title Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication _Sort descending_	Description
DOXORUBICIN IM HYDROCHLOR PUR IDE FOR ITIES/ <i>Organic</i> INJECTION <i>Impurities</i>	USPNF Online Online		28-Oct-2022	1-Nov-2022	NA	NA	limit in the original article as listed in <i>Table 5</i> (see <i>Pesticide Residue Analysis</i> under <i>Articles of Botanical Origin</i> <561>) In <i>Analysis</i> : Change <i>P</i> = potency of doxorubicin in USP Doxorubicin Hydrochloride RS (µg/mg) to: <i>P</i> = potency of doxorubicin hydrochloride in USP Doxorubicin Hydrochloride RS (µg/mg)

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