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HYDROXYZINEASSAY/	USP43–NF38	2271	28-May-2021	1-Jun-2021	NA	NA	In <i>System suita</i>

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
PAMOATE CAPSULES	<i>Procedure</i>								<i>bility/Suitability requirements/Resolution: Change 4-chlorobenophenone, to: 4-chlorobenzop</i>
CARBOMER H OMOPOLYMER	IMPURITIES	<i>USPNF 2021 ISSUE 1</i>	Online	28-May-2021		1-Jun-2021	NA	NA	<i>In Limit of Ethyl Acetate and Cyclohexane/Analysis: Change Samples: Standard stock solution, Standard solution A, Standard solution B, Standard solution C, and Sample solution to: Samples: Standard solution A, Standard solution B, Standard solution C, and</i>

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									<i>Sample solution AND In Limit of Benzene/Analysis: Change Samples: Standard stock solution, Standard solution A, Standard solution B, Standard solution C, and Sample solution to: Samples: Standard solution A, Standard solution B, Standard solution C, and Sample solution</i>
CLOZAPINE	CHEMICAL INFORMATION	USPNF 2021 ISSUE 1	Online	28-May-2021		1-Jun-2021	NA	NA	Change 326.82 to: 326.83
OIL- AND WATER-SOLUBLE VITAMINS	STRENGTH	USPNF 2021 ISSUE 1	Online	30-Apr-2021		1-May-2021	NA	NA	In <i>Vitamins B3 (as Niacinamide)</i> ,

Monograph Title Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
WITH MINERALS CHEWABLE GELS							<p><i>B6, and Folic Acid, Method 1/System suitability.</i> Change [Note—The relative retention times for niacinamide, pyridoxine, and folic acid are about 1.0, 1.6, 2.0, and 3.0 respectively.] to: [Note—The relative retention times for niacinamide, pyridoxine, and folic acid are about 1.0, 2.0, and 3.0, respectively.] AND In <i>Vitamins B1 (as Thiamine Ion) and B2 (as Riboflavin), Method 1; Vitamin C (as</i></p>

Monograph Title Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							<p><i>Ascorbic Acid), Vitamins B3 (as Niacinamide), B6 (as Pyridoxine), and Folic Acid, Method 2/Analysis: Change Calculate the percentage of the labeled amount of vitamin B1 as thiamine ion (C₁₂H₁₇N₄OS⁺), vitamin B2 as riboflavin (C₁₇H₂₀N₄O₆), vitamin B3 as niacinamide (C₆H₆N₂O), vitamin B6 as pyridoxine (C₈H₁₁NO₃), and folic acid (C₁₉H₁₉N₇O₆), in the portion of Chewable Gels taken: to:</i></p>

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							<p>Calculate the percentage of the labeled amount of vitamin C as ascorbic acid (C₆H₈O₆), vitamin B1 as thiamine ion (C₁₂H₁₇N₄OS⁺), vitamin B2 as riboflavin (C₁₇H₂₀N₄O₆), vitamin B3 as niacinamide (C₆H₆N₂O), vitamin B6 as pyridoxine (C₈H₁₁NO₃), and folic acid (C₁₉H₁₉N₇O₆), in the portion of Chewable Gels taken: AND In <i>Vitamins B1 (as Thiamine Ion) and B2 (as Riboflavin), Method 1; Vitamin C (as</i></p>

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							<p><i>Ascorbic Acid), Vitamins B3 (as Niacinamide), B6 (as Pyridoxine), and Folic Acid, Method 2/Acceptance criteria: Change 90.0%–150.0% of the labeled amounts of vitamin B3 as niacinamide, vitamin B6 as pyridoxine, riboflavin, and thiamine as thiamine ion (C₁₂H₁₇N₄OS⁺); and NLT 90.0% and NMT 245.0% of the labeled amount of folic acid to: 90.0%–250.0% of the labeled amount of vitamin C as ascorbic acid;</i></p>

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CARBOMER IN CHEMICAL TERPOLYMER INFORMATION	<i>USPNF 2021</i>	Online	30-Apr-2021	1-May-2021	NA	NA	90.0%–150.0% of the labeled amounts of vitamin B3 as niacinamide, vitamin B6 as pyridoxine, riboflavin, and thiamine as thiamine ion (C ₁₂ H ₁₇ N ₄ OS ⁺); and NLT 90.0% and NMT 245.0% of the labeled amount of folic acid Please see the updated chemical structure at online.uspnf.com
DOXEPIN HYDROCHLORIDE CAPSULES PERFORMANCE TESTS/ Dissolution <711>	<i>Revision Bulletin (Official May 01, 2021)</i>	Online	30-Apr-2021	1-May-2021	NA	NA	<i>In Test 1/Analysis:</i> Change <i>L</i> = label claim of doxepin hydrochloride (mg/Capsule) to: <i>L</i> = label claim (mg/Capsule)

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AZITHROMYCI PERFORMANC N FOR ORAL E SUSPENSION TESTS/ <i>Dissolution</i> <711>	USPNF 2021 ISSUE 1	Online	30-Apr-2021	1-May-2021	NA	NA	In <i>Medium</i> : Change Sodium phosphate buffer, pH of 6.0 (14.2 g/L of disodium hydrogen orthophosphate anhydrous in <i>water</i> , adjusted with dilute hydrochloric acid to a pH of 6.0) to: Sodium phosphate buffer, pH 6.0 (14.2 g/L of sodium phosphate, dibasic, anhydrous in <i>water</i> , adjusted with dilute hydrochloric acid to pH 6.0) AND In <i>Solution A</i> : Change

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MALTITOL SOLUTION	IM PURITIES/ <i>Limit of Nickel</i>	USPNF 2021	Online	30-Apr-2021		1-May-2021	NA	NA	orthophosphoric acid to: phosphoric acid Change <i>Solution A:</i> [Note—Prepare this solution fresh weekly.] to: <i>Standard nickel solution A:</i> [Note—Prepare this solution fresh weekly.]
CARBOMER IN IM TERPOLYMER	PURITIES/ <i>Limit of Acrylic Acid</i>	USPNF 2021	Online	30-Apr-2021		1-May-2021	NA	NA	In <i>Analysis:</i> Change $C_U =$ concentration of Carbomer Interpolymer in the <i>Sample solution</i> (g) to: $C_U =$ concentration of Carbomer Interpolymer in the <i>Sample solution</i> (mg/g)
AZITHROMYCI ASSAY/		USPNF 2021	Online	30-Apr-2021		1-May-2021	NA	NA	In <i>Solution A:</i>

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N FOR ORAL Procedure SUSPENSION	ISSUE 1						Change orthophosphoric acid to: phosphoric acid
CARBOPROST CHEMICAL TROMETHAMI INFORMATION NE	USP43–NF38 771		30-Apr-2021	1-May-2021	NA	NA	Change 489.64 to: 489.65 AND Change (Z)-7-[(1R,2R,3R,5S)-3,5-Dihydroxy-2-[(E)-(3S)-3-hydroxy-3-methyl-1-octenyl]cyclopentyl]-5-heptenoic acid compound with 2-amino-2-(hydroxymethyl)-1,3-propanediol (1:1); to: (Z)-7-[(1R,2R,3R,5S)-3,5-Dihydroxy-2-[(E)-(3S)-3-hydroxy-3-methyloct-1-enyl]

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SERTRALINE HYDROCHLORIDE	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USPNF 2021 ISSUE 1	Online	26-Mar-2021		1-May-2021	NA	NA	cyclopentyl]-5-h eptenoic acid compound with 2-amino-2-(hydr oxymethyl)-1,3- propanediol (1:1); In USP Sertraline Hydrochloride Racemic Mixture RS: Change $C_{17}H_{17}Cl_2 \cdot HCl$ to: $C_{17}H_{17}Cl_2N \cdot HCl$
RILUZOLE TABLETS	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USP43–NF38	3883	26-Mar-2021		1-Apr-2021	NA	NA	In USP Riluzole Related Compound A RS: Change 4-Trifluorometh oxyaniline. $C_7H_6F_3NO$ 177.12 to: 4-(Trifluorometh oxy)aniline. $C_7H_6F_3NO$ 177.13
MIRTAZAPINE	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USP43–NF38	Online	26-Mar-2021		1-Apr-2021	NA	NA	In USP

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TABLETS	EQUIREMENT								<p>Mirtazapine Resolution Mixture RS: Change Impurity D: 1,2,3,4,10,14b-Hexahydropyrazino[2,1-a]pyrido[2,3-c][2]benzazepine .</p> <p>to:</p> <p>Impurity D: [Note—This impurity may be available either as the free base form or as the hydrochloride salt form.] 1,2,3,4,10,14b-Hexahydropyrazino[2,1-a]pyrido[2,3-c][2]benzazepine or 1,2,3,4,10,14b-Hexahydropyrazino[2,1-a]</p>

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PACKAGING AND STORAGE REQUIREMENTS	GENERAL <i>Packaging Definitions</i>	<i>Second Supplement to USP43–NF38</i>	Online	26-Mar-2021		1-Dec-2025	NA	NA]pyri do[2,3-c][2]benzazepine hydrochloride. In <i>Light-resistant container</i> . Change ?661.2?, <i>Functionality, Spectral Transmission Requirements for Light-Resistant Components and Systems</i> . to: ?661.2?, <i>Functionality Test Method, Spectral Transmission Requirements for Light-Resistant Components and Systems</i> .
GENERAL NOTICES AND REQUIREMENTS	2. OFFICIAL STATUS AND LEGAL	<i>USPNF 2021 ISSUE 1</i>	Online	26-Mar-2021		1-May-2021	NA	NA	In 2.10. <i>Official Text</i> . Change http://www.uspn

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TS	RECOGNITION								f.com . to: https://online.uspnf.com .
CARBOMER COPOLYMER	CHEMICAL INFORMATION	<i>USPNF 2021 ISSUE 1</i>	Online	26-Mar-2021		1-May-2021	NA	NA	See online.uspnf.com for correction
GLYBURIDE AND METFORMIN HYDROCHLORIDE TABLETS	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	<i>USP43-NF38</i>	2128	26-Mar-2021		1-Apr-2021	NA	NA	In USP Glyburide Related Compound A RS: Change 368.84 to: 368.83
AMPICILLIN	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	<i>USP43-NF38</i>	316	26-Mar-2021		1-Apr-2021	NA	NA	In USP Amoxicillin Related Compound A RS: Change (2S,5R,6R)-6-Amino-3,3-dimethyl-7-oxo-4-thia-1-azabicyclo[3.2.0]heptane-2-carboxylic acid. C ₁₆ H ₁₄ N ₂ O ₂ 266.29 to: (2S,5R,6R

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TRIHXYPHENIM IDYL HYDROC PUR HLORIDE ITIES/ <i>Organic</i> TABLETS <i>Impurities</i>	USP43–NF38	4519	26-Mar-2021	1-Apr-2021	NA	NA)-6-Amino-3,3-di methyl-7-oxo-4- thia-1-azabicycl o[3.2.0]heptane -2-carboxylic acid. C ₈ H ₁₂ N ₂ O ₃ S 216.26 In <i>Sample</i> <i>solution</i> : Change Nominally 1 mg/mL of trihexyphenidyl hydrochloride in <i>Diluent</i> prepared as follows. to: Nominally 1 mg/mL of trihexyphenidyl hydrochloride prepared as follows.
RILUZOLE ADDITIONAL R EQUIREMENT S/USP <i>Reference</i> <i>Standards <11></i>	USP43–NF38	3882	26-Mar-2021	1-Apr-2021	NA	NA	In USP Riluzole Related Compound A RS: Change 4-Trifluorometh oxyphenylamine

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MIRTAZAPINE	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USP43–NF38	Online	26-Mar-2021		1-Apr-2021	NA	NA	<p>; 4-trifluoromethoxyaniline. $C_7H_6F_3NO$ 177.12 to: 4-(Trifluoromethoxy)aniline. $C_7H_6F_3NO$ 177.13 In USP Mirtazapine Resolution Mixture RS: Change Impurity D: 1,2,3,4,10,14b-Hexahydro-1H-pyrido[2,1-a]pyridine do[2,3-c][2]benzazepine . to: Impurity D: [Note—This impurity may be available either as the free base form or as the hydrochloride salt form.] 1,2,3,</p>

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PACKAGING AND STORAGE REQUIREMENTS	GENERAL <i>Packaging Definitions</i>	<i>Second Supplement to USP43–NF38</i>	Online	26-Mar-2021	1-Apr-2021	NA	NA	4,10,14b-Hexahydropyrazino[2,1-a]pyrido[2,3-c][2]benzazepine or 1,2,3,4,10,14b-Hexahydropyrazino[2,1-a]pyrido[2,3-c][2]benzazepine hydrochloride. In <i>Light-resistant container</i> . Change ?671?, <i>Spectral Transmission</i> to: ?671?, <i>Spectral Transmission for Light-Resistant Packaging Components or Systems</i>
NADOLOL AND IMPURITIES BENDROFLUMETHIAZIDE		<i>USPNF 2021 ISSUE 1</i>	Online	26-Mar-2021	1-May-2021	NA	NA	In <i>Organic Impurities, Procedure 2: Be</i>

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TABLETS									<i>ndroflumethiazi de Related Impurities/Acceptance criteria:</i> Change The reporting threshold is NMT 0.05%. to: The reporting threshold is 0.05%.
ZILEUTON	<i>USP Reference standards <11></i>	<i>USP43–NF38</i>	4679	26-Mar-2021		1-Apr-2021	NA	NA	In USP Zileuton Related Compound A RS: Change <i>N</i> -(1-B enzo-[<i>b</i>]thien-2-ylethyl) urea. $C_{11}H_{12}N_2OS$ 220.30 to: <i>N</i> -(1-B enzo-[<i>b</i>]thien-2-ylethyl) urea; Also known as 1-[1

Monograph Title Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							<p>-(Benz o[<i>b</i>]thiophen-2-yl)et hyl]urea. C₁₁H₁₂N₂OS 220.29 AND In USP Zileuton Related Compound B RS: Change 2-(Be nzo[<i>b</i>]thien-2-oyl)ben zo[<i>b</i>]thiopene. C₁₇H₁₀OS₂ 294.40 to: 2-(Be nzo[<i>b</i>]thien-2-oyl)ben zo[<i>b</i>]thiophene; Also known as Bis(b enzo[<i>b</i>]thiophen-2-yl)m ethanone. C₁₇H₁₀OS₂ 294.39 AND In USP Zileuton</p>

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									Related Compound C RS: Change 1-Ben zo-[<i>b</i>]thien-2-ylethan one. C ₁₀ H ₈ OS 176.24 to: 1-Ben zo-[<i>b</i>]thien-2-ylethan one; Also known as 1-(Be nzo[<i>b</i>]thiophen-2-yl)et han-1-one. C ₁₀ H ₈ OS 176.23 In USP Amlodipine Related Compound A RS: Change 522.93 to: 522.94 Change
AMLODIPINE AND OLMESARTAN MEDOXOMIL TABLETS	ADDITIONAL R EQUIREMENT S/USP <i>Reference</i> <i>Standards <11></i>	USP43–NF38	Online	26-Feb-2021		1-Mar-2021	NA	NA	
STRONTIUM CHLORIDE Sr	CHEMICAL INFORMATION	USP43–NF38	4126	26-Feb-2021		1-Mar-2021	NA	NA	

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89 INJECTION									⁸⁹ SrCl ₂ to: ⁸⁹ SrCl ₂
FENTANYL CITRATE	CHEMICAL INFORMATION	USPNF 2021 ISSUE 1	Online	26-Feb-2021		1-May-2021	NA	NA	Change 528.59 to: 528.60
PACLITAXEL INJECTION	TITLE	USPNF 2021 ISSUE 1	Online	26-Feb-2021		1-May-2021	NA	NA	Change Paclit 1axel Injection to: Paclitaxel Injection
TOTAL ORGANIC CARBON	PROCEDURES /2. Sterile Water	USPNF 2021 ISSUE 1	Online	26-Feb-2021		1-May-2021	NA	NA	In Column 2 in Table 2: Change 1,4-Benzoquinone (mL/L) to: 1,4-Benzoquinone (mg/L) AND In Column 2 and 4 in Table 3: Change Sucrose Concentration (mg/mL) to: Sucrose Concentration

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COPOVIDONE IM PURITIES/ <i>Limit of Monomers (1 2021) -Vinyl-2-Pyrrolidone and Vinyl Acetate)</i>	<i>Harmonization Online (Official May 01,</i>		26-Feb-2021	1-May-2021	NA	NA	(mg/L) In <i>Standard solution:</i> Change 0.25 ?g/mL of 1-vinyl-2-pyrrolidone and 5 ?g/mL of vinyl acetate, respectively, diluted from the <i>Standard stock solution</i> in <i>Mobile phase</i> to: 0.25 ?g/mL of 1-vinyl-2-pyrrolidone and 0.25 ?g/mL of vinyl acetate, respectively, diluted from the <i>Standard stock solution</i> in <i>Mobile phase</i>
AMITRIPTYLIN ASSAY/ E HYDROCHLORIDE	<i>USP43–NF38 Procedure</i>	261	26-Feb-2021	1-Mar-2021	NA	NA	In <i>System suitability solution:</i> Change 0.5 µg/mL of USP Amitriptyline

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							<p>Related Compound A RS, 1 µg/mL of USP</p> <p>Amitriptyline Hydrochloride RS, and 1.5 µg/mL each of USP</p> <p>Amitriptyline Related Compound B RS, USP Cyclo benzaprine Hydrochloride RS, and USP Nortriptyline Hydrochloride RS from suitable volumes of <i>Standard solution, System suitability stock solution A, and System suitability stock solution B</i> in <i>Mobile phase</i> to:</p>

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							0.5 µg/mL of USP Amitriptyline Related Compound A RS, 1 µg/mL of USP Amitriptyline Hydrochloride RS, and 1.5 µg/mL each of USP Amitriptyline Related Compound B RS, USP Cyclo benzaprine Hydrochloride RS, and USP Nortriptyline Hydrochloride RS from suitable volumes of <i>System suitability stock solution A</i> and <i>System suitability stock solution B</i> in <i>Mobile phase</i>

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VENLAFAXINE PERFORMANC HYDROCHLORE IDE EXTENDE TESTS/ D-RELEASE CAPSULES		USP43–NF38	4598	26-Feb-2021		1-Mar-2021	NA	NA	In Test 2/Analysis: Change Result _i = (r _U /r _S) × C _S × (M _{r1} /M _{r2}) to: Result _i = (A _U /A _S) × C _S × (M _{r1} /M _{r2}) AND Change r _U = peak response from the <i>Sample solution</i> r _S = peak response from the <i>Standard solution</i> to: A _U = absorbance from the <i>Sample solution</i> A _S = absorbance from the <i>Standard solution</i>
FENTANYL CITRATE	IM PUR	USPNF 2021 ISSUE 1	Online	26-Feb-2021		1-May-2021	NA	NA	In both variable definition lists in

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									<i>Analysis:</i> Change M_{r1} = molecular weight of fentanyl citrate, 528.59 M_{r2} = molecular weight of fentanyl, 336.47 to: M_{r1} = molecular weight of fentanyl citrate, 528.60 M_{r2} = molecular weight of fentanyl, 336.48
AMLODIPINE AND OLMESARTAN MEDOXOMIL TABLETS	IM PUR ITIES/ <i>Organic Impurities</i>	USP43–NF38 Online		26-Feb-2021		1-Mar-2021	NA	NA	In the first variable definition list in <i>Analysis:</i> Change 522.93 to: 522.94
CLONIDINE	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USP43–NF38	1100	26-Feb-2021		1-Mar-2021	NA	NA	In USP Clonidine Related Compound A RS: Change 1-Acetyl-2-(2,6-

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CLOZAPINE	IM PURITIES/Organic Impurities	USPNF 2021 ISSUE 1	Online	26-Feb-2021	1-May-2021	NA	NA	dichlorophenylamino)-2-(4,5-dihydroimidazol). to: 1-Acetyl-2-(2,6-dichlorophenylamino)imidazolidine. In <i>System suitability/Suitability requirements/Resolution:</i> Change NLT 2.5 between demethyl clozapine and clozapine to: NLT 2.5 between demethyl clozapine and clozapine, <i>System suitability solution</i> Change <i>Quantitative Assessment of Cleaning</i>
CLEANING GLASS APPARATUS	CLEANING VALIDATION BEST PRACTICES	USPNF 2021 ISSUE 1	Online	26-Feb-2021	1-May-2021	NA	NA	

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ANALYTICAL METHODS BASED ON THE TESTING OF PARTICLE COUNTING VIA LIGHT SCATTERING	4. FACTORS THAT AFFECT THE TESTING OF PHENOMENA—PARTICLE COUNTING VIA LIGHT SCATTERING	<i>Second Supplement to USP43–NF38</i>	Online	26-Feb-2021		1-May-2021	NA	NA	Procedure to: Quantitative Assessment of Cleaning Process In paragraph 1: Change an airborne liquid counter. to: an airborne counter.
FEXOFENADINE HYDROCHLORIDE	ADDITIONAL REQUIREMENTS/USP Reference Standards <11>	<i>USP43–NF38</i>	1869	29-Jan-2021		1-Feb-2021	NA	NA	In USP Fexofenadine Related Compound B RS: Change 3-[1-Hydroxy-4-[4-(hydroxydiphenylmethyl)-1-piperidiny]butyl]-?, ?-dimethyl benzeneacetic acid hydrochloride. $C_{32}H_{39}NO_4 \cdot HCl$ 538.12 to:

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HYDROMORPHINE HYDROCHLORIDE ORAL SOLUTION	USP43–NF38	2252	29-Jan-2021	1-Feb-2021	NA	NA	<p>3-[1-Hydroxy-4-[4-(hydroxydiphenylmethyl)-1-piperidiny]butyl]-?, ?-dimethyl benzeneacetic acid hydrochloride monohydrate; Also known as 2-(3-{1-Hydroxy-4-[4-(hydroxydiphenylmethyl)piperidin-1-yl]butyl}phenyl)-2-methylpropanoic acid hydrochloride monohydrate. $C_{32}H_{39}NO_4 \cdot HCl \cdot H_2O$ 556.14</p> <p>In footnote g of <i>Table 3</i>: Change 2,2?-Bihydromorphone. to: (5?)-3-Hydroxy-2-[(5?)-3-hydroxy-17-methyl-6-o</p>

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RISKS AND MITIGATION STRATEGIES FOR THE STORAGE AND TRANSPORTATION OF FINISHED DRUG PRODUCTS	4. RISK MITIGATION CATEGORIES AS QMS ELEMENTS	<i>Second Supplement to USP43–NF38</i>	Online	29-Jan-2021		1-Feb-2021	NA	NA	xo-4,5-epoxymorphinan-2-yl]-17-methyl-4,5-epoxymorphinan-6-one dihydrochloride. In <i>4.1 Documentation and Procedures/4.1.3 Labels</i> : Change The use of symbols that are recognized by international organizations is strongly recommended. to: The use of symbols that are recognized by international organizations is strongly recommended. See <i>General Notices, 10.20. Labeling</i> . In <i>Cation-</i>
DOPAMINE HY	<i>Limit of 5-hydro</i>	<i>USP43–NF38</i>	1495	18-Dec-2020		1-Jan-2021	NA	NA	

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
DROCHLORID E AND DEXTROSE INJECTION	<i>xymethylfurfural and related substances</i>								<i>exchange column: Change Proceed as directed under Column Partition Chromatography (see Chromatography ?621?), to: Proceed as directed under Chromatography ?621?, General Procedures, Column Chromatography, In Table 3/footnote b: Change 1-(2-(2-Methoxyphenoxy)ethylamino)-3-(6,7,8,9-tetrahydro-5H-carbazol-4-yl)propan-2-ol. to:</i>
CARVEDILOL	IM PURITIES/ <i>Organic Impurities, Procedure 2</i>	USP43–NF38	Online	18-Dec-2020		1-Jan-2021	NA	NA	<i>Change 1-(2-(2-Methoxyphenoxy)ethylamino)-3-(6,7,8,9-tetrahydro-5H-carbazol-4-yl)propan-2-ol. to:</i>

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NOREPINEPHRINE BITARTRATE	IDENTIFICATION N/C. Procedure	USP43–NF38	Online	18-Dec-2020		1-Jan-2021	NA	NA	1-(2-(2-Methoxyphenoxy)ethylamino)-3-(2,3,4,9-tetrahydro-1H-carbazol-5-yl)propan-2-ol. In <i>Sample solution</i> : Change 0.01 µg/mL to: 0.1 mg/mL
ELASTOMERIC COMPONENT FUNCTIONAL SUITABILITY IN PARENTERAL PRODUCT PACKAGING/DELIVERY SYSTEMS	5. NEEDLE AND SPIKE ACCESS FUNCTIONAL SUITABILITY TESTS	<i>Second Supplement to USP43–NF38</i>	Online	18-Dec-2020		1-Jan-2021	NA	NA	In paragraph 4 of 5.1 <i>Frame nomenclature/Cartridge system s/Procedure A</i> : Change graticule to: graticule
PINDOLOL TABLETS	ASSAY/ Procedure	<i>Second Supplement to USP43–NF38</i>	Online	18-Dec-2020		1-Jan-2021	NA	NA	In <i>Chromatographic system/Run time</i> : Change

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FENTANYL	CHEMICAL INFORMATION	USP43–NF38	Online	18-Dec-2020		1-Jan-2021	NA	NA	NLT 2 times the retention time of the nortriptyline peak to: NLT 2 times the retention time of the nortriptyline peak Change 336.47 to: 336.48
DOBUTAMINE IN DEXTROSE INJECTION	ASSAY/ Procedure 1: Dextrose	USP43–NF38	1470	18-Dec-2020		1-Jan-2021	NA	NA	In <i>Analysis</i> : Change Result = [(100 × a) × (I/?)] × (1/C _U) × (M _{r1} /M _{r2}) × 100 to: Result = [(100 × a)/(I × ?)] × (1/C _U) × (M _{r1} /M _{r2}) × 100
CARVEDILOL	ADDITIONAL REQUIREMENT S/USP Reference Standards	USP43–NF38	Online	18-Dec-2020		1-Jan-2021	NA	NA	In USP Carvedilol Related Compound A RS: Change 629.74 to:

Monograph Title Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							629.75 AND In USP Carvedilol Related Compound B RS: Change 645.74 to: 645.76 AND In USP Carvedilol Related Compound C RS: Change 496.60 to: 496.61 AND In USP Carvedilol Related Compound E RS: Change 2-(2-Methoxyphenoxy)ethyl amine. $C_9H_{13}NO_2$ 167.21 to:

Monograph Title Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							<p>[Note—This material may be available in the free base or salt form.]</p> <p>2-(2-Methoxyphenoxy)ethyl amine. $C_9H_{13}NO_2$ 167.21</p> <p>2-(2-Methoxyphenoxy)ethyl amine hydrochloride monohydrate. $C_9H_{13}NO_2 \cdot HCl$? H_2O 221.68</p> <p>AND</p> <p>In USP Carvedilol System Suitability Mixture RS: Change Mixture of approximately 0.1% carvedilol related compound F (1-(2-(2-Methoxyphenoxy)ethylam</p>

Monograph Title Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							<p>ino)-3-(2,3,4,9-tetrahydro-1<i>H</i>-carbazol-5-yl)propan-2-ol) in a matrix of carvedilol drug substance.</p> <p>to:</p> <p>Contains a mixture of carvedilol related compound F in a matrix of carvedilol drug substance: Carvedilol. Carvedilol related compound F.</p> <p>[Note—This material may be available in the free base or salt form.]</p> <p>1-(2-(2-Methoxyphenoxy)ethylamino)-3-(2,3,4,9-tetrahydro-</p>

Monograph Title Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							o-1H -carbazol-5-ylox y)propan-2-ol. C ₂₄ H ₃₀ N ₂ O ₄ 410.51 1-(2-(2-Methoxy phenoxy)ethyla mino)-3-(2,3,4,9 -tet rahydr o-1H -carbazol-5-ylox y)propan-2-ol acetate. C ₂₄ H ₃₀ N ₂ O ₄ ? C ₂ H ₄ O ₂ 470.57

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