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CARBINOXAMINE MALEATE	IMPURITIES/*Organic Impurities*	*Second Supplement to USP41–NF36*	8786	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In *Standard stock solution*: Change (equivalent to 0.05 mg/mL of USP Carbinoxamine Maleate RS free base) and 0.05 mg/mL each of USP Carbinoxamine Related Compound A RS, USP Carbinoxamine Related Compound B RS, and USP Carbinoxamine Related Compound C RS free base to: (equivalent to 0.05 mg/mL of carbinoxamine) and 0.05 mg/mL each of USP Carbinoxamine Related Compound A RS, USP Carbinoxamine Related Compound B RS, and USP Carbinoxamine Related Compound C RS (as the free base) AND In the *Standard solution*: Change (equivalent to 0.001 mg/mL of USP Carbinoxamine Maleate RS free base) and 0.001 mg/mL each of USP Carbinoxamine Related Compound A RS, USP Carbinoxamine Related Compound B RS, and USP Carbinoxamine Related Compound C RS free base to: (equivalent to 0.001 mg/mL of carbinoxamine) and 0.001 mg/mL each of USP Carbinoxamine Related Compound A RS, USP Carbinoxamine Related Compound B RS, and USP Carbinoxamine Related Compound C RS (as the free base) AND In the *Analysis*: Change C_S = concentration of USP Carbinoxamine Maleate RS free base to: C_S = concentration of USP Carbinoxamine Maleate RS (as the free base)
METAXALONE	IDENTIFICATION/*B.*	*USP41–NF36*	2611	29-Mar-2019	1-Apr-2019	*NA*	*NA*	Change The retention time of the major peak of the *Sample solution* corresponds to that of the *Sample solution*, as obtained in the *Assay*. to: The retention time of the major peak of the *Sample solution* corresponds to that of the *Standard solution*, as obtained in the *Assay*.

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SODIUM BICARBONATE	IMPURITIES/*Carbonate*	*USP42–NF37*	4019	28-Jun-2019	1-Jul-2019	*NA*	*NA*	In *Analysis*: Change and promptly add 10 g of sodium bicarbonate to: and promptly add 10 g of Sodium Bicarbonate
2,5-DIHYDROXYBENZOIC ACID	REAGENTS AND REFERENCE TABLES/ *Reagent Specifications*	*USP42–NF37*	6097	22-Nov-2019	1-Dec-2019	*NA*	*NA*	Change [303-07-1]. to: [490-79-9].
TRIAZOLAM TABLETS	*Uniformity of dosage units <905>*	*USP41–NF36*	4202	28-Dec-2018	1-Jan-2019	*NA*	*NA*	In *Mobile phase* and *Chromatographic system*: Change Proceed as directed in the *Assay* under *Triazolam*. to: Proceed as directed in the *Assay*. AND In *Procedure*: Change Proceed as directed for *Procedure* in the *Assay* under *Triazolam*. to: Proceed as directed in the *Assay*.
CLOMIPHENE CITRATE	ADDITIONAL REQUIREMENTS/ *USP Reference Standards <11>*	*USP42–NF37*	1068	31-May-2019	1-Jun-2019	*NA*	*NA*	In USP Clomiphene Related Compound A RS: Change (*E*,*Z*)-2-[4-(1,2-Diphenylethyl)phenoxy]-*N*,*N*-diethylethanamine hydrochloride. to: (*E*,*Z*)-2-[4-(1,2-Diphenylvinyl)phenoxy]-*N*,*N*-diethylethanamine hydrochloride. Also known as (*E*,*Z*)-2-[4-(1,2-Diphenylethenyl)phenoxy]-*N*,*N*-diethylethanamine hydrochloride.
FEXOFENADINE HYDROCHLORIDE TABLETS	ADDITIONAL REQUIREMENTS/ *USP Reference Standards <11>*	*Revision Bulletin (Official November 01, 2018)*	Online	30-Aug-2019	1-Sep-2019	*NA*	*NA*	In USP Fexofenadine Related Compound A RS: Change Benzeneacetic acid, 4-[1-oxy-4-[4-(hydroxydiphenylmethyl)-1-piperidinyl]butyl]-?,?-dimethyl. to: 2-(4-{4-[4-(Hydroxydiphenylmethyl)piperidin-1-yl]butanoyl}phenyl)-2-methylpropanoic acid; Also known as Benzeneacetic acid, 4-[1-oxy-4-[4-(hydroxydiphenylmethyl)-1-piperidinyl]butyl]-?,?-dimethyl.
LEVALBUTEROL HYDROCHLORIDE	ADDITIONAL REQUIREMENTS/ *USP Reference Standards <11>*	*USP42–NF37*	2518	26-Apr-2019	1-May-2019	*NA*	*NA*	In USP Levalbuterol Related Compound D RS: Change 5-{2-[(1,1-Dimethylethyl)amino]-1-hydroxyethyl}-2-hydroxy-benzaldehyde. $C_{13}H_{19}NO_3$ 237.29 ?[NOTE—This may be available as the sulfate salt (2:1).]? (USP 1-May-2019) to: 5-[2-(*tert*-Butylamino)-1-hydroxyethyl]-2-hydroxybenzaldehyde sulfate (2:1) (salt); Also known as 5-[2-{(1,1-Dimethylethyl)amino}-1-hydroxyethyl]-2-hydroxy-benzaldehyde sulfate (2:1). $(C_{13}H_{19}NO_3)_2 \cdot H_2SO_4$ 572.67
ALUMINA, MAGNESIA, AND SIMETHICONE ORAL SUSPENSION	SPECIFIC TESTS/ *Microbial Enumeration Tests <61> and Tests for Specified Microorganisms <62>*	*USP42–NF37*	170	26-Jul-2019	1-Aug-2019	*NA*	*NA*	Change cfu/g to: cfu/mL

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<121> INSULIN ASSAYS	ASSAY/*Rabbit Blood Sugar Method—Quantitative*	*Revision Bulletin (Official May 01, 2019)*	Online	27-Dec-2019	1-Jan-2020	*NA*	*NA*	In *Standard stock solution*: Change of USP Insulin RS of the appropriate species to: of the USP Insulin Reference Standard of the appropriate species AND In *Sample stock solution*: Change of USP Insulin RS of the appropriate species. to: of the USP Insulin Reference Standard of the appropriate species.
SALIX SPECIES BARK POWDER	ADDITIONAL REQUIREMENTS	*USP42–NF37*	5189	25-Jan-2019	1-Feb-2019	*NA*	*NA*	In *Labeling*: Change The label states the Latin binomial(s) of one or several *Salix* species included in the article. to: The label states the Latin binomial(s) of one or several *Salix* species included in the article. Dosage forms prepared with this article should bear the following statement: Not for use in children, women who are pregnant or nursing, or by persons with known sensitivity to aspirin. AND In *USP Reference Standards <11>*: Delete (This monograph is postponed indefinitely.)
EPRINOMECTIN	IMPURITIES/*Organic Impurities*/*Acceptance criteria*	*USP42–NF37*	1627	31-May-2019	1-Jun-2019	*NA*	*NA*	In *Total unknown impurities*: Change NMT 1.0 to: NMT 1.0%
BENDAMUSTINE HYDROCHLORIDE FOR INJECTION	ADDITIONAL REQUIREMENTS/ *USP Reference Standards <11>*	*USP42–NF37*	487	27-Sep-2019	1-Oct-2019	*NA*	*NA*	In USP Bendamustine Related Compound B RS: Change 4-(1-Methyl-5-morpholino-1*H*-benzimidazol-2-yl)butanoic acid. $C_{16}H_{21}N_3O_3$ 303.36 to: 4-(1-Methyl-5-morpholino-1*H*-benzimidazol-2-yl)butanoic acid hydrochloride. $C_{16}H_{21}N_3O_3 \cdot x$HCl
BACILLUS COAGULANS CAPSULES	ASSAY/*Enumeration*	*USP42–NF37*	4749	31-May-2019	1-Jun-2019	*NA*	*NA*	In *Peptone diluent*: Change Dispense into sterile containers as needed for preparing samples. Trace mineral solution. Prepare a solution containing to: Dispense into sterile containers as needed for preparing samples. *Trace mineral solution*: Prepare a solution containing
HYPROMELLOSE ACETATE SUCCINATE	ASSAY	*USP42–NF37*	5772	25-Oct-2019	1-Nov-2019	*NA*	*NA*	In *Content of Methoxy and 2-Hydroxypropoxy Groups*/*Analysis*: Change Result = $(r_{UM}/r_{SM}) \times (C_S/C_U) \times (M_{r1}/M_{r2})$ to: Result = $(r_{UM}/r_{SM}) \times (C_S/C_U) \times (M_{r1}/M_{r2}) \times 100$ AND Change Result = $(r_{UI}/r_{SI}) \times (C_S/C_U) \times (M_{r1}/M_{r2})$ to: Result = $(r_{UI}/r_{SI}) \times (C_S/C_U) \times (M_{r1}/M_{r2}) \times 100$

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REAGENTS AND REFERENCE TABLES	REAGENT SPECIFICATIONS	*USP42–NF37*	6079	26-Apr-2019	1-May-2019	*NA*	*NA*	In *Beef Extract/Microbial Content*: Change MT to: NMT
LOVASTATIN TABLETS	PERFORMANCE TESTS/*Dissolution <711>*	*First Supplement to USP42–NF37*	8727	26-Jul-2019	1-Aug-2019	*NA*	*NA*	In *Mobile phase*: Change Processed as directed in the *Assay* to: Proceed as directed in the *Assay*.
CARBINOXAMINE MALEATE TABLETS	IMPURITIES/*Organic Impurities*	*Second Supplement to USP41–NF36*	8788	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In the *Standard stock solution*: Change USP Carbinoxamine Maleate RS free base) to: carbinoxamine) AND In the *Standard solution*: Change USP Carbinoxamine Maleate RS free base) to: carbinoxamine) AND In the *Analysis*: Change C_S = concentration of USP Carbinoxamine Maleate RS free base to: C_S = concentration of USP Carbinoxamine Maleate RS (as the free base)
PREDNISOLONE SODIUM PHOSPHATE	*Related compounds*	*USP41–NF36*	3416	29-Mar-2019	1-Apr-2019	*NA*	*NA*	In *Table 1*: Add Prednisolone sodium phosphate 1.00 — —
VORICONAZOLE	IMPURITIES/ *Voriconazole Related Compounds C and D*	*USP42–NF37*	4601	28-Jun-2019	1-Jul-2019	*NA*	*NA*	In *System suitability solution*: Change 0.25 µg/mL of USP Voriconazole RS to: 0.25 µg/mL of USP Voriconazole RS in *Mobile phase*
0.1 N POTASSIUM HYDROXIDE VS	REAGENTS AND REFERENCE TABLES/ *Solutions*	*USP42–NF37*	6185	22-Nov-2019	1-Dec-2019	*NA*	*NA*	Change *Standardization*: Add 2 drops of phenolphthalein TS to 20 mL of 0.1 N potassium hydroxide VS. Titrate with 0.1 N hydrochloric acid VS until a permanent pale-pink color is produced. to: *Standardization*: Add 2 drops of phenolphthalein TS to 20 mL of 0.1 N hydrochloric acid VS. Titrate with the potassium hydroxide solution until a permanent pale-pink color is produced.

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VITAMIN A	ADDITIONAL REQUIREMENTS	*USP41–NF36*	4327	28-Dec-2018	1-Jan-2019	*NA*	*NA*	Change Delete the following ?•*USP Reference Standards <11>* USP Retinyl Acetate RS USP Retinyl Palmitate RS ?(CN 1-May-2018) to: •*USP Reference Standards <11>* USP Retinyl Acetate RS USP Retinyl Palmitate RS
DACTINOMYCIN	IDENTIFICATION/ *A. Ultraviolet Absorption <197U>/Acceptance criteria*	*USP42–NF37*	1203	31-May-2019	1-Jun-2019	*NA*	*NA*	In *Absorptivity*: Change The absorptivity of the *Sample solution* at 445 nm is NLT 95.0% and NMT 103.0% that of the *Standard solution*. to: The absorptivity, calculated on the dried basis, of the *Sample solution* at 445 nm is NLT 95.0% and NMT 103.0% that of the *Standard solution*.
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE EXTENDED-RELEASE TABLETS	ADDITIONAL REQUIREMENTS/ *USP Reference Standards <11>*	*Revision Bulletin (Official August 01, 2018)*	Online	30-Aug-2019	1-Sep-2019	*NA*	*NA*	In USP Fexofenadine Related Compound A RS: Change Benzeneacetic acid, 4-[1-oxy-4-[4-(hydroxydiphenylmethyl)-1-piperidinyl]butyl]-?,?-dimethyl. to: 2-(4-{4-[4-(Hydroxydiphenylmethyl)piperidin-1-yl]butanoyl}phenyl)-2-methylpropanoic acid; Also known as Benzeneacetic acid, 4-[1-oxy-4-[4-(hydroxydiphenylmethyl)-1-piperidinyl]butyl]-?,?-dimethyl.
LEVALBUTEROL INHALATION SOLUTION	IMPURITIES/*Limit of S-Albuterol*	*USP42–NF37*	Online	26-Apr-2019	1-May-2019	*NA*	*NA*	In *Mobile phase*: Change ?*Acetonitrile, methanol,* and acetic acid (50:50). To each liter of the solution add 3 mL of acetic acid and 1 mL of triethylamine.? (USP 1-May-2019) to: Acetonitrile and methanol (50:50). To each liter of the solution add 3 mL of acetic acid and 1 mL of triethylamine.
BUSPIRONE HYDROCHLORIDE	IDENTIFICATION/*B.*	*USP42–NF37*	618	26-Jul-2019	1-Aug-2019	*NA*	*NA*	Change relative retention time to: retention time
POLYETHYLENE GLYCOL	CHEMICAL INFORMATION	*USP42–NF37*	5882	31-Jan-2020	1-Feb-2020	*NA*	*NA*	See https://www.uspnf.com/sites/default/files/usp_pdf/EN/january-2020-errata-with-image.pdf for correction
ARGATROBAN	CHEMICAL INFORMATION	*USP41–NF36*	346	22-Feb-2019	1-Mar-2019	*NA*	*NA*	See https://www.uspnf.com/sites/default/files/usp_pdf/EN/february_2019_errata_image1.pdf for correction
GUANABENZ ACETATE	IMPURITIES/ *Limit of 2,6-Dichlorobenzaldehyde/ Chromatographic system*	*USP42–NF37*	2129	31-May-2019	1-Jun-2019	*NA*	*NA*	In *Column*: Change 1.8-mm × 3-mm; to: 1.8-m × 3-mm;

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DESFLURANE	CHEMICAL INFORMATION	*USP42–NF37*	1230	27-Sep-2019	1-Oct-2019	*NA*	*NA*	Change (±)-2-Difluoromethyl 1,2,2,2-tetrafluoroethyl ether to: (±)-2-Difluoromethyl 1,2,2,2-tetrafluoroethyl ether; 2-(Difluoromethoxy)-1,1,1,2-tetrafluoroethane.
FLUDROCORTISONE ACETATE TABLETS	IMPURITIES/*Organic Impurities*/*Table 1*	*Second Supplement to USP41–NF36*	8843	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In footnote a: Change 9-Fluoro-11?,17,21-trihydroxypregn-4-ene-3,20-dione 21-acetate. to: 9-Fluoro-11?,17,21-trihydroxypregn-4-ene-3,20-dione.
LIQUID GLUCOSE	ASSAY/*Reducing Sugars*	*USP42–NF37*	5741	31-May-2019	1-Jun-2019	*NA*	*NA*	In the variable definition list in *Analysis*: Change C_U = concentration of dextrose equivalent to: C_U = concentration of Liquid Glucose
<55> BIOLOGICAL INDICATORS—RESISTANCE PERFORMANCE TESTS	D-VALUE DETERMINATION	*USP42–NF37*	6385	25-Oct-2019	1-Nov-2019	*NA*	*NA*	In the third paragraph in *Procedure*: Change stated spore filter to: stated spore titer
REAGENTS AND REFERENCE TABLES	REAGENT SPECIFICATIONS	*USP42–NF37*	6104	26-Apr-2019	1-May-2019	*NA*	*NA*	In *Ferric Nitrate*: Change [10421-48-4]. to: [7782-61-8].
OXALIPLATIN INJECTION	IMPURITIES/*Limit of Oxalic Acid*/*Chromatographic system*	*First Supplement to USP42–NF37*	8781	26-Jul-2019	1-Aug-2019	*NA*	*NA*	In *Column*: Change L31 to: L81
RUTIN	CHEMICAL INFORMATION	*USP41–NF36*	4841	29-Mar-2019	1-Apr-2019	*NA*	*NA*	Change 3-Rhamnoglucoside of 5,7,3',4'-tetrahydroxyflavonol; 2-(3,4-Dihydroxyphenyl)-5,7-dihydroxy-4*H*-chromen-4-one-3-yl 6-*O*-?-L-rhamnopyranosyl-?-D-glucoside [250249-75-3]. to: 3-Rhamnoglucoside of 5,7,3',4'-tetrahydroxyflavonol trihydrate; 2-(3,4-Dihydroxyphenyl)-5,7-dihydroxy-4*H*-chromen-4-one-3-yl 6-*O*-?-L-rhamnopyranosyl-?-D-glucoside trihydrate [250249-75-3].
ZONISAMIDE	IMPURITIES/*Organic Impurities*	*USP42–NF37*	4675	28-Jun-2019	1-Jul-2019	*NA*	*NA*	In *Analysis*: Change C_U = concentration of zonisamide related compound A in the *Sample solution* (mg/mL) to: C_U = concentration of zonisamide in the *Sample solution* (mg/mL)
25% TETRABUTYLAMMONIUM HYDROXIDE TS	REAGENTS AND REFERENCE TABLES/*Solutions*	*Second Supplement to USP42–NF37*	9336	22-Nov-2019	1-Dec-2019	*NA*	*NA*	Change Transfer about 34.82 g to: Transfer about 77.1 g

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BENZETHONIUM CHLORIDE	IMPURITIES/*Organic Impurities/Acceptance criteria*	*First Supplement to USP41–NF36*	8297	25-Jan-2019	1-Feb-2019	*NA*	*NA*	In *Total impurities*: Change 1.0% to: NMT 1.0%
DIDANOSINE	IMPURITIES/*Related Compounds*	*USP42–NF37*	1336	31-May-2019	1-Jun-2019	*NA*	*NA*	In *System suitability solution*: Change 0.5 mg/mL of of didanosine from USP Didanosine System Suitability Mixture RS in *Diluent* to: 0.5 mg/mL of USP Didanosine System Suitability Mixture RS in *Diluent*
MECLOFENAMATE SODIUM	CHEMICAL INFORMATION	*USP42–NF37*	2706	30-Aug-2019	1-Sep-2019	*NA*	*NA*	Change 336.15 to: 336.14 AND Change [6385-02-0]; to: [67254-91-5]; AND Change UNII: 9MMQ0YER4E. to: UNII: 94NJ818U2W.
LEVALBUTEROL INHALATION SOLUTION	IMPURITIES/*Organic Impurities*	*USP42–NF37*	2520	26-Apr-2019	1-May-2019	*NA*	*NA*	In Row 3 of *Table 3*: Change Levalbuterol?—? —?— to: Levalbuterol?1.0?—?—
BUSPIRONE HYDROCHLORIDE TABLETS	IDENTIFICATION/*B.*	*USP42–NF37*	621	26-Jul-2019	1-Aug-2019	*NA*	*NA*	Change relative retention time to: retention time
ALFADEX	CHEMICAL INFORMATION	*USP42–NF37*	5561	31-Jan-2020	1-Feb-2020	*NA*	*NA*	Change 972.84 to: 972.85
DEXMEDETOMIDINE HYDROCHLORIDE	CHEMICAL INFORMATION	*USP41–NF36*	Online	22-Feb-2019	1-Mar-2019	*NA*	*NA*	This erratum applies to the *USP-NF* ONLINE platform only. See https://www.uspnf.com/sites/default/files/usp_pdf/EN/february_2019_errata_image1.pdf for correction
LEVOCARNITINE	IMPURITIES/ *Enantiomeric Purity*	*USP42–NF37*	2541	31-May-2019	1-Jun-2019	*NA*	*NA*	In the second equation in *Analysis*: Change Result = $(R_L$? $P_B)/(P_A$? $P_B)$ to: Result = $(R_L$? $P_B)/(P_A$? $P_B) \times 100$

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DESFLURANE	ADDITIONAL REQUIREMENTS/ *USP Reference Standards <11>*	*USP42–NF37*	1230	27-Sep-2019	1-Oct-2019	*NA*	*NA*	In USP Desflurane Related Compound A RS: Change Bis-(1,2,2,2-tetrafluoroethyl)ether. to: Bis-(1,2,2,2-tetrafluoroethyl)ether; Also known as: 1,1,1,2-Tetrafluoro-2-(1,2,2,2-tetrafluoroethoxy)ethane.
MESNA	ADDITIONAL REQUIREMENTS/ *USP Reference Standards <11>*	*Second Supplement to USP41–NF36*	8904	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In USP Mesna Related Compound A RS: Change 2-(Acetylthio)ethane-1-sulfonic acid. $C_4H_8O_4S_2$ 184.22 to: 2-(Acetylthio)ethane-1-sulfonic acid, potassium salt, crystal adduct with potassium chloride. $C_4H_7KO_4S_2$? KCl 296.86 AND In USP Mesna Related Compound B RS: Change 2,2?-Disulfanediylbis(ethane-1-sulfonic acid). $C_4H_{10}O_6S_4$ 282.36 to: 2,2?-Disulfanediylbis(ethane-1-sulfonic acid), dipotassium salt, crystal adduct with sodium chloride. $C_4H_8K_2O_6S_4$? NaCl 416.98
REAGENTS AND REFERENCE TABLES	SOLUTIONS/*0.01 M Edetate Disodium VS*	*USP42–NF37*	6179	31-May-2019	1-Jun-2019	*NA*	*NA*	In *Standardization*: Change previously dried at 100° to: previously dried at 110°
<1222> TERMINALLY STERILIZED PHARMACEUTICAL PRODUCTS—PARAMETRIC RELEASE	INTRODUCTION	*USP42–NF37*	8021	25-Oct-2019	1-Nov-2019	*NA*	*NA*	In paragraphs 4 and 5: Change a probability of a PNSU to: a PNSU

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