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HYOSCYAMIN	*USP Reference*	*USP37–NF32*	3293	30-Jan-2015	1-Feb-2015	*USP39–NF34*	*Second*	Line 2 of USP

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E SULFATE	*standards <11>*						*Supplement to USP38–NF33*	Hyoscyamine Related Compound A RS: Change (1*R*,3*R*,5*S*)-8-azabicyclo[3.2.1]oct-3-yl(2*S*)-3-hydroxy-2-phenylpropanoate. to: (1*R*,3*r*,5*S*)-8-Azabicyclo[3.2.1]oct-3-yl(2*S*)-3-hydroxy-2-phenylpropanoate sulfate (2:1).
POTASSIUM SODIUM TARTRATE	*Identification/C:*	*USP37–NF32*	4369	30-Jan-2015	1-Feb-2015	*USP39–NF34*	*Second Supplement to USP38–NF33*	Change A solution (1 in 10) responds to the tests for *Tartrate <191>*. to: Responds to the tests for *Tartrate <191>*.

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TRIBASIC CALCIUM PHOSPHATE	IMPURITIES/*Dibasic Salt and Calcium Oxide*	*USP37–NF32*	5883	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Delete the subsection *Blank*: 25.0 mL of *Titrant*
TRAZODONE HYDROCHLORIDE	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*First Supplement to USP37–NF32*	6708	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 2 of USP Trazodone Related Compound D RS: Change (2-{3-[4-(4-Bromophenyl)piperazin-1-yl]propyl}-[1*?*]pyridin-3(2*H*)-one hydrochloride. to: 2-{3-[4-(3-Bromophenyl)piperazin-1-yl]propyl}-[1*?*]pyridin-3(2*H*)-one hydrochloride.

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PACKAGING AND STORAGE REQUIREMENTS	GENERAL DEFINITIONS	*USP37–NF32*	315	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Single-Dose* (see also *Injections <1>, Containers for Injections*): Change A single-unit package for an article intended for parenteral administration. Examples of single-dose containers include prefilled syringes, cartridges, fusion-sealed containers, and closure-sealed containers when so labeled. to: A single-unit package for an article intended for parenteral administration. A single-dose container is

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								labeled as such. Examples of single-dose containers include prefilled syringes, cartridges, fusion-sealed containers, and closure-sealed containers when so labeled.
SODIUM PICOSULFATE	CHEMICAL INFORMATION	*Second Supplement to USP37–NF32*	7253	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 5: Change Disodium 4,4?-(pyridin-2-ylmethanediyl)dibenzenesulfonate to: Disodium 4,4?-(pyridin-2-ylmethanediyl)dibenzenesulfate
FLUDARABINE PHOSPHATE	IM PURITIES/*Limit of Sodium*	*USP37–NF32*	3003	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Standard solution*: Change 1 µg/mL of sodium chloride in water to:

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NIFEDIPINE EXTENDED-RELEASE TABLETS	PERFORMANCE TESTS/ *Dissolution <711>/Test 2*	*Second Supplement to USP37–NF32*	Online	21-Nov-2014	1-Dec-2014	*USP38–NF33*	*USP38–NF33*	1 µg/mL of sodium in water Line 1 of *Solution A*: Change sodium phosphate to: dibasic sodium phosphate
MINOCYCLINE FOR INJECTION	IMPURITIES	*USP37–NF32*	3843	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 6 of *Limit of Epiminocycline*: Change [Note—The relative retention times for epiminocycline and minocycline are 0.86 and 1.0, respectively.] to: [Note—The relative retention times for epiminocycline and minocycline are 0.7 and 1.0,

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								respectively.]
SULFAMETHOXAZOLE AND TRIMETHOPRIM INJECTION	ASSAY/*Procedure*	*USP37–NF32*	4784	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 3 of *Standard solution*: Change in *Mobile phase* from *Sample stock solution* to: in *Mobile phase* from *Standard stock solution*
SAW PALMETTO EXTRACT	COMPOSITION /*Content of Long-Chain Alcohols and Sterols*	*USP37–NF32*	5545	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 3 of *Acceptance criteria*:Add The lipophilic Extract contains 0.15%–0.35% of long-chain alcohols, and the hydroalcoholic Extract contains 0.01%–0.15% of long-chain alcohols on the anhydrous basis.
ERYTHROMYCIN DELAYED-	PERFORMANCE	*First Supplement to*	6633	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to*	Line 1 of *Standard*

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RELEASE TABLETS	TESTS/ *Dissolution <711>/Test 1/Buffer stage*	*USP37–NF32*					*USP38–NF33*	*solution*: Change 0.28 mg/mL of USP Erythromycin RS in *Medium* to: Dissolve USP Erythromycin RS in *Medium* to obtain a concentration similar to that of the *Sample solution*. AND Line 1 of *Sample solution*: Change Pass portions of the solution under test through a suitable filter. to: If necessary, dilute a filtered portion of the solution under test with

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								Medium to obtain a solution containing about 0.28 mg/mL of erythromycin.
POLYSORBATE 80	SPECIFIC TESTS/*Fats and Fixed Oils, Acid Value <401>*	*Second Supplement to USP37–NF32*	7089	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Analysis*: Change with 0.1 N potassium hydroxide or 0.1 N sodium hydroxide to: with 0.1 N potassium hydroxide VS or 0.1 N sodium hydroxide VS
CEFUROXIME AXETIL FOR ORAL SUSPENSION	ASSAY/ *Procedure*	*USP37–NF32*	2243	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Change *Buffer*: 23 mg/mL of monobasic ammonium phosphate in water to: *Solution A*: 23 g/L of

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								monobasic ammonium phosphate in water AND Line 4 of *System suitability solution*: Change Dilute with *Buffer* to volume. to: Dilute with *Solution A* to volume. AND Line 3 of *Standard solution*: Change and dilute with *Buffer* to volume. to: and dilute with *Solution A* to volume. AND Line 3 of

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								Sample solution: Change and dilute with *Buffer* to volume. to: and dilute with *Solution A* to volume.
GANODERMA LUCIDUM FRUITING BODY POWDER	COMPOSITION /*Content of Triterpenoic Acids*	*Revision Bulletin (Official August 01, 2014)*	Online	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Standard solution A*: Change USP Ganoderic Acid RS to: USP Ganoderic Acid A RS AND Line 3 of variable definition list in *Analysis*: Change r_S = peak area of ganoderic acid in *Standard solution A* to: *r*

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								$_S$= peak area of ganoderic acid A in *Standard solution A* AND Line 11 of variable definition list in *Analysis*: Change *F* = relative response factor, with respect to ganoderic acid A (see *Table 3*) to: *F* = relative response factor, with respect to ganoderic acid A (see *Table 2*)
MANNITOL INJECTION	*Assay*	*USP37–NF32*	3653	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Change: *Mobile phase*, *Resolution solution*, and *Chromatographic system* —Proceed as directed in the *Assay* under

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								Mannitol. to: *Mobile phase*—Use degassed water. *Resolution solution*—Dissolve sorbitol and USP Mannitol RS in water to obtain a solution having concentrations of about 4.8 mg per mL of each. *Chromatographic system* (see *Chromatography <621>*)—The liquid chromatograph is equipped with a refractive index detector that is maintained at a constant temperature and a 4-mm ×

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								25-cm column that contains packing L19. The column temperature is maintained at a temperature between 30° and 85° controlled within ±2° of the selected temperature, and the flow rate is about 0.5 mL per minute. Chromatograph the *Standard preparation*, and record the peak responses as directed for *Procedure*: the relative standard deviation for replicate injections is not more than 2.0%. In a similar manner,

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								chromatograph the *Resolution solution*: the resolution, *R*, between the sorbitol and mannitol peaks is not less than 2.0. AND Line 1 of *Procedure*: Change Proceed as directed for *Procedure* in the *Assay* under Mannitol. to: Separately inject equal volumes (about 20 µL) of the *Assay preparation* and the *Standard preparation* into the chromatograph, record the chromatograms,

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								and measure the responses for the major peaks.
SACCHARIN SODIUM	IDENTIFICATION/*B. Procedure*	*USP37–NF32*	4638	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Analysis*: Change To the *Sample solution* to: To 10 mL of the *Sample solution*
ZANAMIVIR	ASSAY/*Procedure/Chromatographic system*	*USP37–NF32*	5197	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Column*: Change 5-?m packing L##[1] to: 5-?m packing L82 AND Delete footnote 1
COPOVIDONE	IMPURITIES/*Limit of Monomers (1-Vinyl-2-Pyrrolidone, Vinyl Acetate, and 2-Pyrrolidone)*	*USP37–NF32*	5938	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 29 of *Analysis*: Change Calculate the content of 2-pyrrolidinone to:

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GLYCERYL BEHENATE	DEFINITION	*Second Supplement to USP37–NF32*	7077	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Calculate the content of 2-pyrrolidone Line 7: Change behenic (docosanic) acid to: behenic (docosanoic) acid
METHODS FOR THE DETERMINATION OF PARTICULATE MATTER IN INJECTIONS AND OPHTHALMIC SOLUTIONS	LIGHT OBSCURATION PARTICLE COUNT TEST/ *Instrument Standardization Tests/Particle Counting Accuracy—System Suitability*	*USP37–NF32*	1301	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 11 of *Method 2—Multichannel Instruments*: Change NMT ±10% of stated size. to: NMT ±10% of stated concentration.
GANODERMA LUCIDUM FRUITING BODY	COMPOSITION /*Content of Triterpenoic Acids*	*Revision Bulletin (Official August 01, 2014)*	Online	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Standard solution A*: Change USP Ganoderic Acid RS to: USP Ganoderic Acid A RS

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								AND Line 3 of variable definition list in *Analysis*: Change r_S = peak area of ganoderic acid in *Standard solution A* to: r_S = peak area of ganoderic acid A in *Standard solution A* AND Line 10 of variable definition list in *Analysis*: Change *F* = relative response factor, with respect to ganoderic acid A (see *Table 3*) to: *F* = relative response factor, with respect to

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								ganoderic acid A (see *Table 2*)
FLUTAMIDE	ASSAY/ *Procedure*	*USP37–NF32*	3057	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 2 of *Sample solution*: Change Dissolve the sample in acetonitrile to: Dissolve a previously dried sample in acetonitrile
PETROLATUM	IM PUR ITIES/*Organic Impuri ties*/*Procedure: Organic Acids*	*USP37–NF32*	4253	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Sample solution*: Change 20.0 g of Petrolatum in 100 mL of neutralized alcohol and water (1:2). to: 20.0 g of Petrolatum in 100 mL of a 1 in 2 mixture of neutralized alcohol and

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								water.
SULFAMETHOXAZOLE AND TRIMETHOPRIM ORAL SUSPENSION	ASSAY/ *Procedure*	*USP37–NF32*	4785	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 3 of *Standard solution*: Change in *Mobile phase* from *Sample stock solution* to: in *Mobile phase* from *Standard stock solution*
TRIBASIC CALCIUM PHOSPHATE	ASSAY/ *Procedure*	*USP37–NF32*	5883	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Delete the subsection *Blank*: Proceed as directed in the *Analysis*, omitting the test specimen. AND Equation in *Analysis*: Change Result = {[V_S ? V_B) × M × F]/W} × 100 to: Result = [(V_S × M × F)/W] × 100 AND

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								Line 2 of the variable definition list in *Analysis*: Delete V_B = *Titrant* volume consumed by the *Blank* (mL)
TRAZODONE HYDROCHLORIDE	IM PURITIES/*Limit of Trazodone Related Compound F and Cyclophosphamide Related Compound A/ Chromatographic system/MS conditions*	*First Supplement to USP37–NF32*	6708	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 4 of *Acquisition mode*:Change 272 to: 273
MEBENDAZOLE	ASSAY/ *Procedure*	*Second Supplement to USP37–NF32*	7199	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 3 of *Analysis*: Change Calculate the percentage of each impurity in the portion of Oral Suspension taken: to:

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DOLASETRON MESYLATE	CHEMICAL INFORMATION	*USP37–NF32*	2693	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Calculate the percentage of mebendazole ($C_{16}H_{13}N_3O_3$) in the portion of Mebendazole taken: Line 7: Change [115956-13-3]. to: [878143-33-0] Anhydrous [115956-13-3].
FENTANYL	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*Second Supplement to USP37–NF32*	Online	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Delete USP Fentanyl Related Compound C RS AND Delete USP Fentanyl Related Compound F RS
METFORMIN HYDROCHLORIDE EXTENDED-RELEASE TABLETS	PERFORMANCE TESTS/*Dissolution <711>*	*USP37–NF32*	3732	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Equation in *Test 1*: Change Result = $[(A_U/A_S) \times C_S \times (V ? V_S) + (C_{60} \times V_S) + (C_{180} \times V$

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								$_S$) ×100]/*L* to: Result = [(A_U/A_S) × C_S × (*V* ? V_S) + (C_{60} ×V_S) + (C_{180} ×V_S)] ×(100/*L*) AND Equation 3 in *Test 2*: Change Result = [C_2 × (*V* ? SV_1) + C_1 ×SV_1 ×100]/*L* to: Result = [C_2 × (*V* ? SV_1) + C_1 ×SV_1] ×(100/*L*) AND Equation 4 in *Test 2*: Change Result = {C_n ×[*V* ? (*n* ? 1)V_S] + (C_1 + C_2 + ... + C_{n-1}) × V_S ×100}/*L* to: Result = {C_n ×[*V* ? (*n* ? 1)V_S] + (C_1 + C_2 + ... + C_{n-1}) × V_S} ×(100/*L*)

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SACCHARIN SODIUM	IM PUR ITIES/*Organic Impuri ties/Procedure 1: Limit of Tolue nesulfonamides*	*USP37–NF32*	4638	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 4 of *Acceptance criteria*: Change of the *Internal standard solution* to: of the caffeine (internal standard)
ZANAMIVIR	IM PUR ITIES/*Organic Impuri ties/ Chromatographi c system*	*USP37–NF32*	5197	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Column*: Change 5-?m packing L##[1] to: 5-?m packing L82
CHLORPHENI RAMINE MALEATE	SPECIFIC TESTS/*Optical Rotation, Specific Rotation <781>*	*First Supplement to USP37–NF32*	6617	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Sample*: Change 100 mg/mL in water to: 100 mg/mL in water at 20°
HYDROXYPRO PYL CELLULOSE	ASSAY	*Second Supplement to USP37–NF32*	7080	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Internal standard solution*: Change

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								Methycyclohexane to: Methylcyclohexane
CEFADROXIL FOR ORAL SUSPENSION	IDENTIFICATION/*Thin-Layer C / Chromatographic system*	*USP37–NF32*	2182	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 2 of *Developing solvent system*: Change (60:40:15) to: (60: 40: 1.5)
GANODERMA LUCIDUM FRUITING BODY	SPECIFIC TESTS/ *Botanical Characteristics*	*Revision Bulletin (Official August 01, 2014)*	Online	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 6 of *Macroscopic*: Change concentrically culcate to: concentrically sulcate
LAMIVUDINE AND ZIDOVUDINE TABLETS	IMPURITIES/*Organic Impurities*	*USP37–NF32*	3484	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Row 17 of Column 1 of *Table 2*: Change (the limit includes individual unidentified impurities) to:

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								(the limit includes individual unspecified impurities)
WHITE PETROLATUM	IMPURITIES/*Organic Impurities*/*Procedure: Organic Acids*	*USP37–NF32*	4254	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 1 of *Sample solution*: Change 20.0 g in 100 mL of neutralized alcohol and water (1:2). to: 20.0 g of White Petrolatum in 100 mL of a 1 in 2 mixture of neutralized alcohol and water.
SULFAMETHOXAZOLE AND TRIMETHOPRIM TABLETS	ASSAY/*Procedure*	*USP37–NF32*	4787	21-Nov-2014	1-Dec-2014	*USP39–NF34*	*Second Supplement to USP38–NF33*	Line 3 of *Standard solution*: Change in *Mobile phase* from *Sample stock solution* to: in *Mobile*

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								phase from *Standard stock solution*
STERILE WATER FOR IRRIGATION	CHEMICAL INFORMATION	*USP37–NF32*	5175	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	Add the chemical formula and molecular weight: H_2O 18.02
POWDERED CELLULOSE	IDENTIFICATION/*B. Procedure*	*USP37–NF32*	5923	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	Fourth equation in *Analysis*: Change Result = 95 × $[?]_c/W_S$ × [(100 ? %LOD)/100] to: Result = [95 × $[?]_c$]/{W_S × [(100 ? %LOD)/100]}
BETAMETHASONE SODIUM PHOSPHATE	IM PURITIES/*Limit of Free Betamethasone*	*USP37–NF32*	1965	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	Line 1 of *Sample stock solution*: Change 1.0 mg/mL of Betamethasone Sodium Phosphate in water to: 1.0 mg/mL of Betamethasone

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								Sodium Phosphate in water, prepared as follows. Dissolve 25.0 mg of Betamethasone Sodium Phosphate in water to make 25.0 mL.
DROSPIRENONE AND ETHINYL ESTRADIOL TABLETS	IMPURITIES/*Organic Impurities/ Chromatographic system*	*USP37–NF32*	2739	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	Line 2 of *Detector 2*: Change Monitor the signal at 344 nm between 37 and 42 min. to: Monitor the signal at 344 nm for ethinyl estradiol related compound B (typically between 37 and 42 min).
PSEUDOEPHEDRINE HYDROCHLORIDE	ASSAY/ *Procedure*	*USP37–NF32*	4481	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	Line 2 of *System suitability solution*:

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								Change 0.02 mg/mL of USP Ephedrine Sulfate RS to: 0.002 mg/mL of USP Ephedrine Sulfate RS
STERILE WATER FOR INJECTION	CHEMICAL INFORMATION	*USP37–NF32*	5175	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	Add the chemical formula and molecular weight: H_2O 18.02
STERILE PURIFIED WATER	ADDITIONAL REQUIREMENTS	*USP37–NF32*	5176	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	Add a section: *USP Reference Standards <11>* USP 1,4-Benzoquinone RS USP Sucrose RS
DEXCHLORPHENIRAMINE MALEATE	IMPURITIES/*Organic Impurities*	*First Supplement to USP37–NF32*	6626	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	Line 1 of *Standard solution*: 2.2 ?g/mL of USP Dexchlorpheniramine Maleate RS in *Diluent*, to: 2.8 ?g/mL of

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
ESCITALOPRAM ORAL SOLUTION	IMPURITIES/*Organic Impurities*	*USP37–NF32*	2580	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	USP Dexchlorpheniramine Maleate RS in *Diluent*, Row 6 of Column 1 of *Table 3*: Change Desfluorocitalopram[f] to: Desfluorocitalopram[f,c]
KETOPROFEN EXTENDED-RELEASE CAPSULES	PERFORMANCE TESTS/ *Uniformity of Dosage Units <905>/System suitability/Suitability requirements*	*USP37–NF32*	3469	26-Sep-2014	1-Oct-2014	*USP39–NF34*	*First Supplement to USP38–NF33*	Line 1 of *Tailing factor*: Change NLT 1.5 to: NMT 1.5

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