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METFORMIN H PERFORMANC		*USP39–NF34*	4766	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 3 of *Test 3*:

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YDROCHLORIDE EXTENDED-RELEASE TABLETS	E TESTS/ *Dissolution* <*711*>							Change *Medium, Apparatus 1, Apparatus 2*, and *Analysis*: to: *Medium, Apparatus 1*, and *Apparatus 2*:
PALIPERIDONE	CHEMICAL INFORMATION	*USP39–NF34*	5253	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 5: Change (9*RS*)-3-[2-[4-(6-Fluoro-1,2-benzisoxazol-3-yl)piperidin-1-yl]]ethyl]-9-hydroxy-2-methyl-6,7,8,9-tetrahydro-4*H*-pyrido[1,2-*a*]pyrimidin-4-one to: (9*RS*)-3-[2-[4-(6-Fluoro-1,2-benzisoxazol-3-yl)piperidin-1-yl]]ethyl]-9-hydroxy-2-methyl-6,7,8,9-tetrahydro-4*H*-pyri

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								do[1,2-*a*]pyrimidin-4-one
RANITIDINE IN SODIUM CHLORIDE INJECTION	*USP Reference standards <11>*	*USP39–NF34*	5673	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 2 of USP Ranitidine Related Compound C RS: Change *N* -[2-[[[5-[(Dimeth ylamino)methyl] -2-furanyl]methy l] su lfinyl]ethyl]-*N* -methyl-2-nitro- 1,1-ethenediami ne. to: *N* -{2-[({5-[(Dimeth ylamino)methyl] -2-furanyl}meth
								N ?-methyl-2-nitro -1,1-ethenedia mine.
CHONDROITIN	SPECIFIC	*USP39–NF34*	6570	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 2 of

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SULFATE SODIUM, SHARK	TESTS/*Clarity and Color of Solution*							*Instrumental conditions*: Change (See *Spectrophotometry and Light-Scattering <851>*.) to: (See *Ultraviolet-Visible Spectroscopy <857>*.)
OLEYL ALCOHOL	ASSAY/ *Procedure*	*USP39–NF34*	7424	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 1 of *Standard solution*: Change Prepare 1.0 mg/mL of USP Oleyl Alcohol RS in *Internal standard solution*, and heat the solution in a sealed container in a 50° water bath until oleyl alcohol is dissolved. Allow

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								the solution to cool to room temperature, and mix well. to: 1.0 mg/mL of USP Oleyl Alcohol RS in *Internal standard solution* AND Line 1 of *Sample solution*: Change Prepare 1.0 mg/mL of Oleyl Alcohol in *Internal standard solution*, and heat the solution in a sealed container in a 50° water bath until oleyl alcohol is dissolved. Allow the solution to

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								cool to room temperature, and mix well. to: 1.0 mg/mL of Oleyl Alcohol in *Internal standard solution*
DIAZEPAM INJECTION	*Assay*	*USP39–NF34*	3445	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 7 of *Procedure*: 50C / $V(R_U / R_S)$ to: $50(C / V)(R_U / R_S)$
LORAZEPAM ORAL CONCENTRATE	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP39–NF34*	4621	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 3 of Lorazepam Related Compound B RS: Change $C_{13}H_9ClNO$ to: $C_{13}H_9Cl_2NO$
NICOTINE POLACRILEX	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP39–NF34*	5061	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Add USP Polacrilex Resin RS
RANITIDINE ORAL SOLUTION	*USP Reference standards <11>*	*USP39–NF34*	5671	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 2 of USP Ranitidine Related

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								Compound C RS: Change *N* -[2-[[[5-[(Dimethylamino)methyl]-2-furanyl]methyl] su lfinyl]ethyl]-*N* -methyl-2-nitro-1,1-ethenediamine. to: *N* -{2-[({5-[(Dimethylamino)methyl]-2-furanyl}meth
								N ?-methyl-2-nitro-1,1-ethenediamine.
TROSPIUM CHLORIDE	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP39–NF34*	6287	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 3 of USP Trospium Chloride Related Compound C RS:Change

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								(1*R*,3*r*,5*S*)-3-Hydroxyspiro[8-azoniabicyclo[3.2.1]octane-8,1?-pyrrolidinium] chloride. to: (1*R*,3*r*,5*S*)-3-Hydroxyspiro[8-azoniabicyclo[3.2.1]octane-8,1?-pyrrolidinium] chloride, or (1*R*,3*r*,5*S*)-3-Hydroxyspiro[bicyclo[3.2.1]octane-8,1?-pyrrolidin]-1?-ium chloride.
CETYL ALCOHOL	IM PURITIES/*Limit of Related Fatty Alcohols*	*USP39–NF34*	7239	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 1 of *Sample solution*: Change 1 mg/mL of Cetyl Alcohol in ethanol to: Prepare 1.0 mg/mL of Cetyl Alcohol in ethanol, and

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								heat the solution in a sealed container in a 50° water bath until cetyl alcohol is dissolved. Allow the solution to cool to room temperature, and mix well.
ALPRAZOLAM EXTENDED-RELEASE TABLETS	PERFORMANCE TESTS/ *Dissolution <711>*	*USP39–NF34*	2389	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Variable definition list of second equation in *Test 2/Analysis*: Change V_S = volume of the *Sample solution* withdrawn at each time point and replaced with *Medium* (mL) to: V_S = volume of the *Sample solution* withdrawn at

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								each time point (mL) AND Variable definition list of second equation in *Test 3*/*Analysis*: Change V_S = volume of the *Sample solution* withdrawn at each time point and replaced with *Medium* (mL) to: V_S = volume of the *Sample solution* withdrawn at each time point (mL)
PAROXETINE EXTENDED-RELEASE TABLETS	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*First Supplement to USP39–NF34*	8121	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	USP Paroxetine Related Compound B RS: Add $C_{19}H_{21}NO_3 \cdot HCl$ 347.84

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LORAZEPAM	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP39–NF34*	4618	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 3 of USP Lorazepam Related Compound B RS: Change $C_{13}H_9ClNO$ to: $C_{13}H_9Cl_2NO$
MYCOPHENOLATE SODIUM	ASSAY/*Procedure*	*USP39–NF34*	4965	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 3: Change *Solvent A* to: *Solution A* AND Line 5: Change *Solvent B* to: *Solution B*
PALIPERIDONE	IMPURITIES/*Organic Impurities*/*System suitability*/*Suitability requirements*	*USP39–NF34*	5253	27-May-2016	1-Jun-2016	*USP40–NF35*	*USP40–NF35*	Line 3 of *Resolution*: Change hydroxybenzyl to: hydroxybenzoyl
PLASTIC MATERIALS OF CONSTRUCTION	TEST METHODS/*Extractions*/*Table 3*	*USP39–NF34*	493	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Column 4 of *S3* row: Change Extractable metals: Al, Sb, As, Ba, Cd, Co, Ge, Hg, Mn, Ni,

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								Pb, Ti, V, and Zn to: Extractable metals: Al, As, Ba, Cd, Co, Hg, Mn, Ni, Pb, Ti, V, and Zn
DIPHENHYDR AMINE HYDRO CHLORIDE INJECTION	ASSAY/ *Procedure*	*USP39–NF34*	3529	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 1 of *System suitability solution*:Change USP Diphenhydramine Hydrochloride Related Compound A RS to: USP Diphenhydramine Related Compound A RS AND Line 4 of *System suitability*: Change for diphenhydramine hydrochloride

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								related compound A and diphenhydramine hydrochloride are to: for diphenhydramine related compound A and diphenhydramine are
PLASTIC MATERIALS OF CONSTRUCTION	SPECIFICATIONS/ *Polyethylene/Extractable Metals*	*USP39–NF34*	493	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 1 of *Zirconium*: Change 1 µg/g. to: 0.1 µg/g.
DISSOLUTION	INTERPRETATION/*Immediate-Release Dosage Forms/ Immediate-Release Dosage Forms Pooled Sample*	*USP39–NF34*	540	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Row 3 of Column 1 of *Acceptance Table for a Pooled Sample*: Change S_1 to: S_2 AND Row 4 of Column 1 of *Acceptance*

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								Table for a Pooled Sample: Change S_1 to: S_3
DIGOXIN	IM PURITIES/*Related Glycosides*/*System suitability*	*Interim Revision Announcement (Official November 01, 2015)*	Online	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 2: Change *Sample: System suitability solution* to: *Samples: System suitability solution* and *Standard solution* AND Line 2 of *Suitability requirements*: Change *Resolution:* NLT 1.5 between the digoxin and lanatoside C peaks *Relative standard deviation:* NMT

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								2.0%, determined from the digoxin peak in replicated injections to: *Resolution:* NLT 1.5 between the digoxin and lanatoside C peaks, *System suitability solution* *Relative standard deviation:* NMT 2.0%, determined from the digoxin peak in replicated injections, *Standard solution*
SELENOMETHIONINE	CHEMICAL INFORMATION	*USP38–NF33*	6226	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 3: Change [1464-42-2] to: [3211-76-5]
PLASTIC MATERIALS	TEST ME	*USP39–NF34*	493	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 1 of footnote b:

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OF CONSTRUCTION	THODS/ *Extractions/Table 3*							Change For nonplasticized polyethylene only. to: For polyethylene only.
FLUTICASONE PROPIONATE AND SALMETEROL INHALATION POWDER	ASSAY/ *Procedure*	*USP39–NF34*	4020	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 3 of *System suitability*: Change for salmeterol and fluticasone propionate are to: for fluticasone propionate and salmeterol are
CONTAINERS--PERFORMANCE TESTING	MOISTURE VAPOR TRANSMISSION/*Packaging System Classification for Multiple-Unit Containers and Unit-Dose Containers for Liquid Oral*	*USP38–NF33*	465	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 1 of the Equation: Change $[(W_{1i} ? W_T) ? (W_{14i} ? W_T) ? (W_{C1} ? W_{C14})] \times 365 \times \{[100/(W_{1i} ? W_T)] \times 14\}$ to: $[(W_{1i} ? W_T) ?$

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	Dosage Forms/ Procedure							$(W_{14i}$? $W_T)$? $(W_{C1}$? $W_{C14})]$ × 365 × 100/$(W_{1i}$? $W_T)$ × 14
PLASTIC MATERIALS OF CONSTRUCTION	SPECIFICATIONS/ *Polyethylene Terephthalate and Polyethylene Terephthalate G/Extractable Metals*	*USP39–NF34*	493	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 1 of *Titanium*: Change 0.1 µg/g. to: 1 µg/g.
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE TABLETS	IMPURITIES/*Organic Impurities*	*USP39–NF34*	2895	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Row 3 of Column 1 of *Table 6*:Change Candesartan related compound A[b,c] to: Candesartan cilexetil related compound A[b,c]
MEMANTINE HYDROCHLORIDE TABLETS	IMPURITIES/*Organic Impurities*	*Revision Bulletin (Official October 01, 2015)*	Online	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 3 of *Analysis*: Change of USP Memantine Related Compound E RS or

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								to: of memantine related compound E or AND In the variable definition list: Change r_U = peak response of USP Memantine Related Compound E RS or any individual degradation product from the *Sample solution* to: r_U = peak response of memantine related compound E or any individual degradation product from the *Sample solution*

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RESIDUAL HOST CELL PROTEIN MEASUREMENT IN BIOPHARMACEUTICALS	4. HCP IMMUNOASSAY METHOD VALIDATION/*4.3 Sample Linearity/Table 4*	*Second Supplement to USP38–NF33*	7647	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	*Product* column: Change 10.00 (neat), 5.00, 2.50, 1.25, 0.63, 0.31, 0.16 to: 10.00 (neat), 5.00, 2.50, 1.25, 0.625, 0.3125, 0.15625 AND *Sample 1/HCP ratio* column: Change 4.9, 5.7, 4.8, 5.9, 5.0, 5.1, <6 to: 4.90, 5.70, 4.80, 5.92, 4.96, 5.12, <6 AND *Sample 2/HCP ratio* column: Change 2.0, 3.3, 4.0, 5.9, 5.3, 6.1, <6 to: 2.00, 3.30, 4.00, 5.92, 5.28, 6.08, <6 AND

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								Sample 3/HCP ratio column: Change 0.3, 0.5, 0.6, 0.9, 1.4, <6, <6 to: 0.32, 0.50, 0.60, 0.88, 1.44, <6, <6 AND *Sample 3/% max ratio value* column: Change 83% to: 61%
PLASTIC MATERIALS OF CONSTRUCTION	TEST METHODS/ *Physicochemical Tests/ Absorbance*	*USP39–NF34*	493	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 3 of *Plasticized polyvinyl chloride*: Delete Additionally, for nonplasticized polyvinyl chloride materials only, determine the spectrum between 250 and 330 nm in the alcohol

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								sample associated with *Solution S6*.
METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER	IMPURITIES/*Limit of Methacrylic Acid and Ethyl Acrylate*	*First Supplement to USP39–NF34*	Online	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	Line 5 of *Standard solution*: Change Mix 10.0 mL of this solution to: Mix 5.0 mL of this solution AND Line 7 of *Standard solution*: Change about 0.67 µg/mL to: about 0.5 µg/mL Line 4 of *Sample solution*: Change 10.0 mL of this solution to: 5.0 mL of this solution AND

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ARGININE HYDROCHLORIDE	SPECIFIC TESTS/*Chloride Content*	*USP38–NF33*	2279	25-Mar-2016	1-Apr-2016	*USP40–NF35*	*USP40–NF35*	In the variable definition list in *Analysis*: Change V_F = final volume of the *Sample solution*, 15 mL D = dilution factor for preparation of the *Sample solution*, 5 to: V_F = final volume of the *Sample solution*, 10 mL D = dilution factor for preparation of the *Sample solution*, 10 Delete the subsection *Blank*: 140 mL of water and 1 mL of dichlorofluorescein TS AND The equation in

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								the *Analysis*: Change Result = [(*V* ? *B*) × *N* × *F* × 100]/*W* to: Result = (*V* × *N* × *F* × 100)/*W* AND Line 10 of *Analysis*: Delete *B* = *Blank* titrant volume (mL)
DEXTROAMPHETAMINE SULFATE	ASSAY/ *Procedure*	*USP38–NF33*	3060	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 1 of *System suitability solution*: Change 0.02 µg/mL each of USP Dextroamphetamine Related Compound A RS and USP Dextroamphetamine Related Compound B RS in *Standard solution* to: Transfer about

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								40 mL of the *Standard solution* to a 50-mL volumetric flask. Using a microliter syringe, add 1 µL each of USP Dextroamphetamine Related Compound A RS and USP Dextroamphetamine Related Compound B RS. Dilute with *Standard solution* to volume.
TAMSULOSIN HYDROCHLORIDE CAPSULES	ASSAY/ *Procedure 8*	*USP38–NF33*	5442	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 1 of *Standard solution*: Change Prepare a solution containing 1.0 mg/mL of USP Tamsulosin Hydrochloride RS in methanol.

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								to: Prepare a solution containing 0.1 mg/mL of USP Tamsulosin Hydrochloride RS in methanol. AND Line 6 of *Sample solution*: Change Add 30 mL of *Mobile phase*, and shake by mechanical means for 30 min. to: Add 30 mL of *Mobile phase*, shake by mechanical means for 30 min, and dilute with *Mobile phase* to volume.
ALTEPLASE	ASSAY/ *Biological*	*USP39–NF34*	2398	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to*	Line 1 of *Human*

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	Potency						*USP39–NF34*	*thrombin solution*: Change 33 Units in terms of the U.S. Standard Thrombin/mL in *Buffer* to: 33 U.S. Units in terms of the U.S. Standard Thrombin/mL in *Buffer* AND Line 5 of *Analysis*: Change *Standard solution* and *Sample solution*, to: *Standard solution* or *Sample solution*,
NIFEDIPINE EXTENDED-RELEASE TABLETS	PERFORMANCE TESTS/ *Dissolution*	*Revision Bulletin (Official December 01, 2015)*	Online	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 4 of *Medium*: Change 6 g/L

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	<711>/Test 9							to: 10 g/L
HYDROCHLORIC ACID INJECTION	ASSAY/ *Procedure*	*USP38–NF33*	3770	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 8 of *Analysis*: Change *F* = equivalency factor, 18.23 mg/mEq to: *F* = equivalency factor, 36.46 mg/mEq
DONEPEZIL HYDROCHLORIDE	IMPURITIES/*Organic Impurities/Procedure 2*	*First Supplement to USP38–NF33*	7384	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Footnote h of *Table 3*: Change 1-Benzyl-4-[(5,6-dimethoxyindan-2-yl)methyl]piperidine. to: 1-Benzyl-4-[(5,6-dimethoxyinden-2-yl)methyl]piperidine.
DEXAMETHASONE SODIUM PHOSPHATE INJECTION	ASSAY/ *Procedure*	*Interim Revision Announcement (Official May 01, 2015)*	Online	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 9 of *Analysis*: Change C_S = concentration of USP Dexameth

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								asone Sodium Phosphate RS in the *Standard solution* (µg/mL) to: C_S = concentration of USP Dexameth asone Sodium Phosphate RS in the *Standard solution* (mg/mL)
GALANTAMINE HYDROBROMIDE	IMPURITIES/*Organic Impurities*	*USP38–NF33*	3646	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 8 of *Analysis*: Change Result = $(r_U/r_S) \times (C_S/C_U) \times (1/F) \times (100/100 ? L)$ to: Result = $(r_U/r_S) \times (C_S/C_U) \times (1/F) \times (100/[100 ? L])$
GLUCOSAMINE SULFATE POTASSIUM CHLORIDE	CHEMICAL INFORMATION	*USP38–NF33*	6074	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 5: Change [38899-05-7]. to: [1296149-08-0] .
ALTEPLASE	ASSAY/	*USP39–NF34*	2401	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second*	Line 1 of

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FOR INJECTION	*Biological Potency*						*Supplement to USP39–NF34*	*Human thrombin solution*: Change 33 Units in terms of the U.S. Standard Thrombin/mL in *Buffer* to: 33 U.S. Units in terms of the U.S. Standard Thrombin/mL in *Buffer* AND Line 5 of *Analysis*: Change *Standard solution* and *Sample solution*, to: *Standard solution* or *Sample solution*,
RISPERIDONE TABLETS	*Assay*	*USP38–NF33*	5195	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 4 of *Procedure*: Change

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IMIQUIMOD CREAM	SPECIFIC TESTS/*pH <791>*	*First Supplement to USP38–NF33*	7409	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Calculate the quantity, in mg, of risperidone to: Calculate the percentage of the labeled amount of risperidone Line 1 of *Sample*: Change Nominally 50 mg/mL of imiquimod from Cream in water. to: Nominally 2.5 mg/mL of imiquimod from Cream in water.
DEXAMETHASONE SODIUM PHOSPHATE OPHTHALMIC SOLUTION	ASSAY/ *Procedure*	*Interim Revision Announcement (Official May 01, 2015)*	Online	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 9 of *Analysis*: Change C_S = concentration of USP Dexamethasone Sodium Phosphate RS in the *Standard solution* (µg/mL)

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								to: C_S = concentration of USP Dexamethasone Sodium Phosphate RS in the *Standard solution* (mg/mL)
GALANTAMINE TABLETS	PERFORMANCE TESTS/ *Dissolution*	*USP38–NF33*	3649	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 6 of *Analysis* in *Test 1*: Change Result = $(A_U/A_S) \times (C_S/L) \times (M_{r1}/M_{r2}) \times 100$ to: Result = $(A_U/A_S) \times (C_S/L) \times (M_{r1}/M_{r2}) \times V \times 100$ AND Add to the variable definition list in *Test 1* V = volume of *Medium*, 500 mL AND Equation in *Test 3*: Change

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Result = (r_U/r_S) × (C_S/L) × (M_{r1}/M_{r2}) × 100 to: Result = (r_U/r_S) × (C_S/L) × (M_{r1}/M_{r2}) × V × 100 AND Add to the variable definition list in *Test 3* V = volume of *Medium*, 500 mL
GLUCOSAMINE SULFATE SODIUM CHLORIDE	CHEMICAL INFORMATION	*USP38–NF33*	6075	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 5: Change [38899-05-7]. to: [1296149-13-7].
ROPINIROLE TABLETS	IM PURITIES/*Organic Impurities*	*USP39–NF34*	5756	29-Jan-2016	1-Feb-2016	*USP40–NF35*	*Second Supplement to USP39–NF34*	Footnote b of *Table 2*: Change 4-[2-(Dipropylnitroryl)ethyl]-1,3-dihydrox-2*H*-indol-2-one. to: *N*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								-[2-(2-Oxoindolin-4 -yl)ethyl]-*N* -propylpropan-1 -amine oxide.
DESCRIPTION AND SOLUBILITY	SUNFLOWER OIL	*Second Supplement to USP38–NF33*	7761	20-Nov-2015	1-Dec-2015	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 5: Change *NF category*: Coating agent; emollient; solvent; tablet and/or capsule diluent; vehicle (oleaginous). to: *NF category*: Coating agent; emollient; solvent; diluent; vehicle (oleaginous).
REPOSITORY CORTICOTROPIN INJECTION	ASSAY/ *Procedure*	*Second Supplement to USP38–NF33*	8063	20-Nov-2015	1-Dec-2015	*USP40–NF35*	*Second Supplement to USP39–NF34*	Line 7 of *Replication*: Change (see <111>, *Confidence Intervals for Individual Assays*). to: (see <111>,

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								The Confidence Interval and Limits of Potency).

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