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SODIUM NITR <i>Identification</i>	USP39–NF34	5880	29-Jul-2016	1-Aug-2016	USP41–NF36	First	Line 1 of

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OPRUSSIDE							<i>Supplement to USP40–NF35</i>	Identification C: Change A solution (1 in 4) responds to the flame test for <i>Sodium</i> <191>. to: A solution (1 in 4) imparts an intense yellow color to a nonluminous flame.
BISOCTRIZOLE	IMPURITIES/Organic Impurities	<i>First Supplement to USP39–NF34</i>	8008	29-Jul-2016	1-Aug-2016	<i>USP41–NF36</i>	<i>First Supplement to USP40–NF35</i>	Footnote b of Table 2: Change Phenol, 2,2-methyl-6-(2H-benzotriazol-2-yl)-4-(1,1,3,3-tetramethylbutyl)]. to: Phenol, 2,2-methyl-6-(2H-benzotriazol-2-yl)-4-(1,1,3,3-tetramethylbutyl)].

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ASSESSMENT REFERENCES OF DRUG PRODUCT LEACHABLES ASSOCIATED WITH PHARMACEUTICAL PACKAGING/DELIVERY SYSTEMS	USP39–NF34	1850	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	-benzotriazol-2-yl)-4-(1,1,3,3-tetramethylbutyl)]. Delete reference 8.
GRANISETRO N HYDROCHLORIDE	USP Reference standards <11>	USP39–NF34 4151	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	Line 2 of USP Granisetron Related Compound B RS: Change (N-[(1R,3r,5)-9-Methyl-9-azabicyclo[3.3.1]non-3-yl]-1H-indazole-3-carboxamide). to: (N-[(1R,3r,5S)-9-Methyl-9-azabicyclo[3.3.1]non-3-yl]-1H

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KETOROLAC T ADDITIONAL R ROMETHAMIN EQUIREMENT E INJECTION S/USP Reference Standards <11>	USP39–NF34	4468	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	-indazole-3-carb oxamide). AND Line 2 of USP Granisetron Related Compound E RS: Change ((1 <i>R</i> ,3 <i>r</i> <i>S</i> ,5 <i>S</i>)-9-Methyl-9-az abicyclo[3.3.1]n onan-3-amine, acetate salt). to: ((1 <i>R</i> ,3 <i>r</i> ,5 <i>S</i>)-9-Methyl-9-az abicyclo[3.3.1]n onan-3-amine, acetate salt). Line 4 of USP Ketorolac Related Compound A RS: Change 358.15 to: 358.39 AND Line 3 of USP Ketorolac Related

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							Compound B RS: Change 227.09 to: 227.26 AND Line 3 of USP Ketorolac Related Compound C RS: Change 225.09 to: 225.24 AND Line 3 of USP Ketorolac Related Compound D: Change 211.1 to: 211.3 Line 1 of Samples: Change Standard stock solution, Standard solution, Sample stock
SIMETHICONE ASSAY/ Proce dure/Analysis	USP39–NF34	5843	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	

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BANABA LEAF IDENTIFICATION DRY EXTRACT N	USP39–NF34	6494	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	<i>solution, and Sample solution to:</i> <i>Standard solution and Sample solution</i> Delete Identification A. AND Line 1 of Identification B: Change B. to: A. AND Line 1 of Identification C: Change C. to: B.	
OMEGA-3-ACID ETHYL ESTERS CAPSULES	ASSAY/Content of EPAee, DHAee, and Total Omega-3-Acid Ethyl Esters/Analysis	Second Supplement to USP39–NF34	8755	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	Line 16 of the third equation: Change L = label claim of total omega-3-acids ethyl esters (g/Capsule) to:

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FEXOFENADINE HYDROCHLORIDE	CHEMICAL INFORMATION	USP39–NF34 3895	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	L = label claim of total omega-3-acids ethyl esters (mg/Capsule) Line 7: Change [138452-21-8]. to: [153439-40-8].
HALCINONIDE CREAM	IMPURITIES/ <i>Organic Impurities/Chromatographic system</i>	USP39–NF34 4176	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	Line 1 of <i>Column</i> : Change 1.8-μm packing L1 to: 1.7-μm packing L1
NAPROXEN TABLETS	IMPURITIES/ <i>Organic Impurities</i>	USP39–NF34 4993	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	Line 6 of <i>Analysis</i> : Change Result = $(r_U/r_S) \times (C_U/C_S) \times 100$ to: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$
TRIHEXYPHENIDYL HYDROCHLORIDE EXTENDED-RELEASE CAPSULES	Assay	USP39–NF34 6265	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	Line 1 of <i>Mobile phase and Chromatographic system</i> : Change Prepare as

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							directed in the Assay under <i>Trihexyphenidyl Hydrochloride</i> . to: <i>Mobile phase</i> —Prepare a mixture of acetonitrile, water, and triethylamine (920:80:0.2), adjust with phosphoric acid to a pH of 4.0, mix, filter, and degas. Make adjustments if necessary (see <i>System Suitability</i> under <i>Chromatography</i> <621>). <i>Chromatographic system</i> (see <i>Chromatography</i> <621>)—The liquid chromatograph is equipped with a 210-nm

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							detector and a 4.6-mm × 8-cm column that contains 3-μm packing L1. The flow rate is about 2 mL per minute. Chromatograph the <i>Standard preparation</i>, and record the peak responses as directed for <i>Procedure</i>: the column efficiency determined from the analyte peak is not less than 1300 theoretical plates, the tailing factor for the analyte peak is not more than 3.0, and the relative standard deviation for replicate

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							injections is not more than 1.0%. AND Line 1 of <i>Procedure</i>: Change Proceed as directed for <i>Procedure</i> in the Assay under <i>Trihexyphenidyl Hydrochloride</i>. to: Separately inject equal volumes (about 10 µL) of the <i>Standard preparation</i> and the Assay <i>preparation</i> into the chromatograph, record the chromatograms, and measure the responses for the major peaks. AND

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									Line 6 of <i>Procedure</i>: Change and the other terms are as defined therein. to: C is the concentration, in mg per mL, of USP Trihexyphenidyl Hydrochloride RS in the <i>Standard preparation</i>, r_U and r_S are the trihexyphenidyl peak responses obtained from the Assay <i>preparation</i> and the <i>Standard preparation</i>, respectively. Line 2 of USP Bisotrizole Resolution Mixture RS: Change A mixture of
BISOCTRIZOLE	ADDITIONAL REQUIREMENT	<i>First Supplement to SIUSP Reference Standards <11></i>	8008	29-Jul-2016		1-Aug-2016	USP41–NF36	<i>First Supplement to USP40–NF35</i>	

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									approximately 1.5% of bisoctrizole isomer [phenol, 2,2-methylenebis[6-(2H-benzotriazol-2-yl)-4-(1,1,3,3-tetramethylbutyl)]] in a matrix of bisoctrizole. to: A mixture of approximately 1.5% of bisoctrizole isomer [phenol, 2,2'-methylene bis[6-(2H-benzotriazol-2-yl)-4-(1,1,3,3-tetramethylbutyl)]] in a matrix of bisoctrizole.
ORALLY INHALED AND NASAL DRUG PRODUCTS	REFERENCES	USP39–NF34	1862	29-Jul-2016		1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	Delete reference 3.
GRANISETRO N HYDROCHLORIDE	USP Reference standards <11>	USP39–NF34	4153	29-Jul-2016		1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	Line 2 of USP Granisetron Related

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INJECTION									Compound B RS: Change (<i>N</i> -[(1 <i>R</i> ,3 <i>r</i> ,5)-9-Methyl-9-azabicyclo[3.3.1]non-3-yl]-1 <i>H</i> -indazole-3-carboxamide). to: <i>N</i> -[(1 <i>R</i> ,3 <i>r</i> ,5 <i>S</i>)-9-Methyl-9-azabicyclo[3.3.1]non-3-yl]-1 <i>H</i> -indazole-3-carboxamide.
LOVASTATIN	<i>USP Reference standards <11></i>	<i>USP39–NF34</i>	4631	29-Jul-2016		1-Aug-2016	<i>USP41–NF36</i>	<i>First Supplement to USP40–NF35</i>	Line 2 of USP Lovastatin Related Compound A RS: Change [Dihydro-lovastatin][butanoic acid, 2-methyl-, 1,2,3,4,4a,7,8,8a-octahydro-3,7-dimethyl-8-[2(tetrahydro-4-hydroxy-6-oxo

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SIMVASTATIN IM TABLETS	PURITIES/Organic Impuri	USP39–NF34 5848	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	o-2H -pyran-2-yl)-ethyl]-1-naphthalenyl ester, [1S-[1?(R*), 3?,7?, ,8?(2S*,4S*),-8???]]-. to: (1S,3S,4aR,7S,8S,8aS)-8-{2-[(2R,4R)-4-hydroxy-6-oxotetrahydro-2H-pyran-2-yl]ethyl}-3,7-dimethyl-1,2,3,4,4a,7,8,8a-octahydronaphthalen-1-yl (S)-2-methylbutanoate. Line 5: Change Result = $(r_U/r_S) \times (C_S/C_U) \times$

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	<i>ties/Analysis</i>								$\frac{(M_{r1} - M_{r2})}{M_{r1}} \times 100$ to: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$ AND Delete the variable definitions for M_{r1} and M_{r2} . Line 1: Change 0.5% to: NMT 0.5% Line 6 of Analysis: Change Result = $(r_U/r_S) \times (C_U/C_S) \times 100$ to: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$
FUMARIC ACIDS	SPECIFIC TESTS/ <i>Water Determination, Method I <921></i>	USP39–NF34	7309	29-Jul-2016		1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	Line 1: Change 0.5% to: NMT 0.5%
NAPROXEN SODIUM TABLETS	IMPURITIES/ <i>Organic Impurities</i>	Second Supplement to USP39–NF34	Online	29-Jul-2016		1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	Line 6 of Analysis: Change Result = $(r_U/r_S) \times (C_U/C_S) \times 100$ to: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$
BACITRACIN ZINC	IMPURITIES	USP39–NF34	2674	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Delete the <i>Residue on Ignition <281></i> test.
LORAZEPAM INJECTION	ADDITIONAL REQUIREMENT S/ <i>USP Reference Standards <11></i>	USP39–NF34	4620	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 3 of Lorazepam Related Compound B RS: Change C

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NALTREXONE Related HYDROCHLORIDE		USP39–NF34	4985	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	$^{13}\text{H}_9\text{ClNO}$ to: $\text{C}_{13}\text{H}_9\text{Cl}_2\text{NO}$ Line 5: Change $10F(C/W)(r_U / r_S)$ to: $1000F(C/W)(r_U / r_S)$
RANITIDINE INJECTION	USP Reference standards <11>	USP39–NF34	5670	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 2 of USP Ranitidine Related Compound C RS: Change N -[2-[[[5-[(Dimethylamino)methyl]-2-furanyl]methyl]sulfinyl]- N -methyl-2-nitro-1,1-ethenediamine. to: N -{2-[[{5-[(Dimethylamino)methyl]-2-furanyl}meth

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TETRACYCLIN ASSAY/ E HYDROCHL ORIDE CAPSULES	USP39–NF34	6082	27-May-2016	1-Jun-2016	USP40–NF35	USP40–NF35	<i>N</i> ?-methyl-2-nitro -1,1-ethenedia mine. Line 6 of <i>Sample solution:</i> Change dilute with <i>Diluent</i> to volume. to: dilute with <i>Solution A</i> to volume.
VINPOCETINE IM PUR ITIES/ <i>Organic Impurities</i>	USP39–NF34	6880	27-May-2016	1-Jun-2016	USP40–NF35	USP40–NF35	Footnote a of <i>Table 1:</i> Change Ethyl (12<i>RS</i>,13a SR,13bSR)-13a -ethyl-12-hydrox y-2,3,5,6,12,13, 13a,13b- octa hydro-1<i>H</i> -in dolo[3,2,1-<i>de</i>

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							]py rido[3 ,2,1- <i>ij</i>][1,5]naphthyridi ne-12-carboxyla te (ethyl vincamate). to: Ethyl (12 <i>S</i> ,13 <i>aS</i> ,13 <i>bS</i>)-13 <i>a</i> -ethy l-12-hydroxy-2,3 ,5,6,12,13,13 <i>a</i> ,1
							<i>H</i> -in dolo[3,2,1- <i>de</i>]py rido[3 ,2,1- <i>ij</i>][1,5]naphthyridi ne-12-carboxyla te (ethyl vincamate). AND Footnote b of <i>Table 1</i> : Change

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							Methyl (13aS,13bS)-13a-ethyl-9-methoxy-2,3,5,6,13a,13b-hexahydro-1 <i>H</i> -in dolo[3,2,1- <i>de</i>]pyrido[3,2,1- <i>ij</i>][1,5]naphthyridine-12-carboxylate (apovincamine). to: Methyl (13aS,13bS)-13a-ethyl-2,3,5,6,13a,13b-hexahydro-1 <i>H</i> -in dolo[3,2,1- <i>de</i>]pyrido[3,2,1- <i>ij</i>][1,5]naphthyridine-12-carboxylate

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							ne-12-carboxylate (apovincamine). AND Footnote d of <i>Table 1</i> : Change Ethyl (12 <i>RS</i> ,13a <i>RS</i> ,13b <i>RS</i>)-13a-ethyl-2,3,5,6,12,13,13a,13b-octa hydro-1 <i>H</i> -in dolo[3,2,1- <i>de</i>]pyrido[3,2,1- <i>ij</i>][1,5]naphthyridine-12-carboxylate (dihydrovinpocetine). to: Ethyl (12 <i>R</i> ,13a <i>S</i> ,13b <i>S</i>)-13a-ethyl-2,3,5,6,12,13,13a,13b-octa

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REAGENTS, INDICATORS AND SOLUTIONS	REAGENTS/6. General Tests for Reagents/6.2 Amino Nitrogen Test in Reagents	USP39–NF34	2080	27-May-2016	1-Jun-2016	USP40–NF35	USP40–NF35		hydro-1<i>H</i>-in dolo[3,2,1-<i>de</i>]pyrido[3,2,1-<i>ij</i>][1,5]naphthyridine-12-carboxylate (dihydrovinpocetine). In the numerator of the equation: Change 2.8 to: 14 AND Add $\times f$ AND Line 10: Change where %LOD is the percentage of loss on drying. to: where <i>f</i> is the correction factor

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HAZARDOUS DRUGS—HANDLING IN HEALTHCARE SETTINGS	5. FACILITIES ENGINEERING CONTROLS/5.3 Compounding	<i>First Supplement to USP39–NF34</i>	7721	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	obtained in the standardization of 0.2 N sodium hydroxide and %LOD is the percentage of loss on drying. First bullet in second paragraph: Change • Be externally vented through high-efficiency particulate air (HEPA) filtration to: • Be externally vented
IMIQUIMOD CREAM	IMPURITIES/Organic Impurities	USP39–NF34	4289	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Row 2 of Column 3 of <i>Table 2</i>: Change 1.5 to: 1.15 AND Row 3 of Column 3 of <i>Table 2</i>: Change

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METFORMIN H PERFORMANC YDROCHLORI E DE EXTENDED-TESTS/ RELEASE <i>Dissolution</i> TABLETS <i><711></i>	USP39–NF34	4766	27-May-2016	1-Jun-2016	USP40–NF35	USP40–NF35	1.15 to: 1.5 Line 3 of <i>Test 3</i> : Change <i>Medium</i> , <i>Apparatus 1</i> , <i>Apparatus 2</i> , and <i>Analysis</i> : to: <i>Medium</i> , <i>Apparatus 1</i> , and <i>Apparatus 2</i> :
PALIPERIDON CHEMICAL E INFORMATION	USP39–NF34	5253	27-May-2016	1-Jun-2016	USP40–NF35	USP40–NF35	Line 5: Change (9 <i>RS</i>)-3-[2-[4-(6-Fluo ro-1,2-benzisox azol-3-yl)piperid in-1-yl]]ethyl]-9- hydroxy-2-meth yl-6,7,8,9-tetra hydro-4 <i>H</i> -pyrido [1,2- <i>a</i>]pyrimidin-4-one to: (9 <i>RS</i>)-3-[2-[4-(6-Fluo ro-1,2-benzisox azol-3-yl)piperid in-1-yl]]ethyl]-9-

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RANITIDINE IN USP Reference Standards <11> SODIUM CHLORIDE INJECTION	USP39–NF34	5673	27-May-2016	1-Jun-2016	USP40–NF35	USP40–NF35	hydroxy-2-methyl-6,7,8,9-tetrahydro-4H-pyrido[1,2-a]pyrimidin-4-one Line 2 of USP Ranitidine Related Compound C RS: Change <i>N</i> -[2-[[[5-[(Dimethylamino)methyl]-2-furanyl]methyl]sulfinyl]ethyl]- <i>N</i> -methyl-2-nitro-1,1-ethenediamine. to: <i>N</i> -{2-[[[5-[(Dimethylamino)methyl]-2-furanyl]meth
							<i>N</i>

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CHONDROITIN SPECIFIC SULFATE SODIUM, SHARK	TESTS/ <i>Clarity and Color of Solution</i>	USP39–NF34	6570	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	?-methyl-2-nitro-1,1-ethenediamine. Line 2 of <i>Instrumental conditions</i> : Change (See <i>Spectrophotometry and Light-Scattering <851></i> .) to: (See <i>Ultraviolet-Visible Spectroscopy <857></i> .)
OLEYL ALCOHOL	ASSAY/ <i>Procedure</i>	USP39–NF34	7424	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 1 of <i>Standard solution</i> : Change Prepare 1.0 mg/mL of USP Oleyl Alcohol RS in <i>Internal standard solution</i> , and heat the solution in a sealed container in a

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

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									until oleyl alcohol is dissolved. Allow the solution to cool to room temperature, and mix well. to: 1.0 mg/mL of Oleyl Alcohol in <i>Internal standard solution</i>
DIAZEPAM INJECTION	Assay	USP39–NF34	3445	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 7 of <i>Procedure</i> : 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 ₉ ClNO to: C ₁₃ H ₉ Cl ₂ NO Add USP Polacrilex Resin RS
NICOTINE POLACRILEX	ADDITIONAL REQUIREMENTS / USP Reference	USP39–NF34	5061	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	

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RANITIDINE ORAL SOLUTION	<i>Standards <11></i> <i>USP Reference standards <11></i>	USP39–NF34	5671	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 2 of USP Ranitidine Related Compound C RS: Change <i>N</i> -[2-[[[5-[(Dimethylamino)methyl]-2-furanyl]methyl]sulfinyl]- <i>N</i> -methyl-2-nitro-1,1-ethenediamine. to: <i>N</i> -{2-[[[5-[(Dimethylamino)methyl]-2-furanyl]methyl]sulfinyl]- <i>N</i> -methyl-2-nitro-1,1-ethenediamine.
TROSPIMUM CHLORIDE	ADDITIONAL REQUIREMENT	USP39–NF34	6287	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 3 of USP Trospium

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		<i>SIUSP Reference Standards <11></i>							Chloride Related Compound C RS:Change (1 <i>R</i> ,3 <i>r</i> ,5 <i>S</i>)-3-Hydroxyspiro[8-azoniabicyclo[3.2.1]octane-8,1?-pyrrolidinium] chloride. to: (1 <i>R</i> ,3 <i>r</i> ,5 <i>S</i>)-3-Hydroxyspiro[8-azoniabicyclo[3.2.1]octane-8,1?-pyrrolidinium] chloride, or (1 <i>R</i> ,3 <i>r</i> ,5 <i>S</i>)-3-Hydroxyspiro[bicyclo[3.2.1]octane-8,1?-pyrrolidin]-1?-ium chloride.
CETYL ALCOHOL	IM PURITIES/ <i>Limit of Related Fatty Alcohols</i>	USP39–NF34	7239	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 1 of <i>Sample solution</i> : Change 1 mg/mL of Cetyl Alcohol in ethanol 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
ALPRAZOLAM EXTENDED- RELEASE TABLETS	PERFORMANC E TESTS/ <i>Dissolution</i> <711>	USP39–NF34	2389	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Prepare 1.0 mg/mL of Cetyl Alcohol in ethanol, and 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. Variable definition list of second equation in <i>Test 2/Analysis</i>: Change V_S = volume of the <i>Sample solution</i> withdrawn at each time point and replaced with <i>Medium</i> (mL) 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
									V_S = volume of the <i>Sample solution</i> withdrawn at each time point (mL) AND Variable definition list of second equation in <i>Test 3/Analysis</i> : Change V_S = volume of the <i>Sample solution</i> withdrawn at each time point and replaced with <i>Medium</i> (mL) to: V_S = volume of the <i>Sample solution</i> withdrawn at each time point (mL)
PAROXETINE EXTENDED-RELEASE	ADDITIONAL REQUIREMENT	R <i>First Supplement to USP39–NF34</i> S/ <i>USP</i>	8121	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	USP Paroxetine Related

Monograph Title Section		Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
TABLETS	<i>Reference Standards <11></i>							Compound B RS: Add $C_{19}H_{21}NO_3 \cdot HCl$ 347.84
LORAZEPAM	ADDITIONAL REQUIREMENT <i>S/USP Reference Standards <11></i>	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/ <i>Procedure</i>	USP39–NF34	4965	27-May-2016	1-Jun-2016	USP40–NF35	USP40–NF35	Line 3: Change <i>Solvent A</i> to: <i>Solution A</i> AND Line 5: Change <i>Solvent B</i> to: <i>Solution B</i>
PALIPERIDONE	IMPURITIES/ <i>Organic Impurities/System suitability/Suitability requirements</i>	USP39–NF34	5253	27-May-2016	1-Jun-2016	USP40–NF35	USP40–NF35	Line 3 of <i>Resolution</i> : Change hydroxybenzyl to: hydroxybenzoyl
SAMARIUM 153	Sm <i>Other requirements</i>	USP39–NF34	5791	27-May-2016	1-Jun-2016	USP40–NF35	USP40–NF35	Line 1: Change <i>Injections and</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
LEXIDRONAM INJECTION									<i>Implanted Drug Products <1>; not subject to Container Content.</i> to: Meets the requirements of <i>Injections and Implanted Drug Products <1>; not subject to Container content.</i>
CHONDROITIN SULFATE SODIUM, SHARK	COMPOSITION <i>/Disaccharide Composition</i>	USP39–NF34	6570	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 2 of <i>Chondroitinase ABC solution</i> : Change 10.0 mL of <i>Buffer solution</i> to: 1.0 mL of <i>Buffer solution</i> AND Line 4 of <i>Analysis</i> : Change and 1.0 mL to: and 0.1 mL
SEMISOLID	IN VITRO PER	USP39–NF34	1869	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 7 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
DRUG PRODU	FORMANCE								<i>Application of Drug Release:</i>
CTS—PERFOR	TESTS								Change
MANCE TESTS									The individual amounts of drug released from <i>R</i> is plotted versus time,
									to:
									The individual amounts of drug released from <i>R</i> are plotted versus the square root of time,
SODIUM CETO IM		USP39–NF34	7518	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 1 of
STEARYL	PURITIES/ <i>Limit</i>								<i>Endpoint</i>
SULFATE	<i>of Sodium Chloride and Sodium Sulfate/Sodium sulfatel/ Titrimetric system</i>								<i>detection:</i>
									Change
									Potentiometric
									to:
									Visual
DICLOFENAC IM		USP39–NF34	3460	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Line 1 of
SODIUM EXTE PUR									<i>Standard</i>
NDED-	ITIES/ <i>Organic</i>								<i>solution:</i>
RELEASE	<i>Impurities</i>								Change
TABLETS									0.001 mg/mL of USP Diclofenac

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LORAZEPAM TABLETS	ADDITIONAL REQUIREMENT S/USP Reference Standards RS <11>	USP39–NF34	4622	27-May-2016		1-Jun-2016	USP40–NF35	USP40–NF35	Sodium RS in <i>Diluent</i> to: 0.001 mg/mL each of USP Diclofenac Sodium RS and USP Diclofenac Related Compound A RS in <i>Diluent</i> Line 3 of USP Lorazepam Related Compound B RS: Change C ₁₃ H ₉ ClNO to: C ₁₃ H ₉ Cl ₂ NO

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