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How to Use

- **Searching:** Type keyword in search field at top of page. Search by all or part of a monograph title. For searches using multiple criteria, you will find items that match each of the specified criteria unless quotation marks are used.
 - For example, a search on Aminosalicyclic Acid Tablets will result in anything that contains “Aminosalicyclic” OR “Acid” OR “Tablets”
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- **Sorting:** Click on any column header title to sort alphabetically or chronologically in ascending or descending order. Note: the page load column is sorted alphabetically so that a number is ordered by first digit vs. by the actual number; thus, numbers will not always be in order.
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HYDROXYPRO ASSAY PYL CELLULOSE	<i>Second Supplement to USP37–NF32</i>	7080	21-Nov-2014	1-Dec-2014	<i>USP39–NF34</i>	<i>Second Supplement to USP38–NF33</i>	Line 1 of <i>Internal standard solution:</i> Change Methycyclohexane to: Methylcyclohexane
POLYSORBAT SPECIFIC E 80 TESTS/ <i>Fats and Fixed Oils, Acid Value <401></i>	<i>Second Supplement to USP37–NF32</i>	7089	21-Nov-2014	1-Dec-2014	<i>USP39–NF34</i>	<i>Second Supplement to USP38–NF33</i>	Line 1 of <i>Analysis:</i> 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
DACARBAZINE IM FOR PURITIES/ <i>Limit of 2-Azahypoxanthine</i> INJECTION	<i>Second Supplement to USP37–NF32</i>	Online	26-Sep-2014	1-Oct-2014	<i>USP39–NF34</i>	<i>First Supplement to USP38–NF33</i>	Line 2 of <i>Analysis:</i> Change 2-azahypoxanthine monohydrate

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MEBENDAZOLE	ASSAY/ Procedure	<i>Second Supplement to USP37–NF32</i>	7199	21-Nov-2014	1-Dec-2014	<i>USP39–NF34</i>	<i>Second Supplement to USP38–NF33</i>	to: 2-azahypoxanthine Line 3 of Analysis: Change Calculate the percentage of each impurity in the portion of Oral Suspension taken: to: Calculate the percentage of mebendazole (C ₁₆ H ₁₃ N ₃ O ₃) in the portion of Mebendazole taken: to: Calculate the percentage of mebendazole (C ₁₆ H ₁₃ N ₃ O ₃) in the portion of Mebendazole taken:
SODIUM PICOSULFATE	CHEMICAL INFORMATION	<i>Second Supplement to USP37–NF32</i>	7253	21-Nov-2014	1-Dec-2014	<i>USP39–NF34</i>	<i>Second Supplement to USP38–NF33</i>	Line 5: Change Disodium 4,4?-(pyridin-2-ylmethanediyl)dibenzene sulfonate to: Disodium 4,4?-(pyridin-2-ylmethanediyl)dibenz

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RISPERIDONE ORAL SOLUTION	ADDITIONAL REQUIREMENT <i>SI/USP Reference Standards <11></i>	<i>Second Supplement to USP37–NF32</i>	Online	25-Jul-2014	1-Aug-2014	<i>USP39–NF34</i>	<i>First Supplement to USP38–NF33</i>	enesulfate Line 2 of USP Risperidone Related Compounds Mixture RS: Change Contains a mixture of the following four compounds: 98.9% of <i>Risperidone</i> . 0.5% of <i>Risperidone cis-N-oxide: cis</i> -3-[2-[4-(6-Fluoro-1,2-benzisoxazol-3-yl)-1-piperidinyl]ethyl]-6,7,8,9-tetrahydro-2-methyl-4H-pyrido[1,2-a]pyrimidin-4-N-oxide. 0.3% of <i>Bicyclorisperido</i>

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	—						ne: 3-(4-Fluoro-2-hydroxyphenyl)-1-[2-(6,7,8,9-tetrahydro-2-methyl-4-oxo-4H-pyridin-1(2H)-ylidene)-1H-imidazo[4,5-b]pyridin-3-yl)ethyl]-1-aza-2-azoniabicyclo[2.2.2]oct-2-ene iodide. 0.3% of Z-<i>Oxime</i>: (Z)-3-[2-[4-(2,4-Difluorophenyl)(hydroxyimino)methyl]-1-piperidinylethyl]-6,7,8,9-tetrahydro-2-methyl-4H-pyridin-4-one . to: Contains a

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	—						mixture of the following four compounds: Risperidone. Risperidone <i>cis</i>-<i>N</i>-oxide: <i>cis</i> -3-[2-[4-(6-Fluoro-1,2-benzisoxazol-3-yl)-1-piperidinyl]ethyl]-6,7,8,9-tetrahydro-2-methyl-4<i>H</i>-pyrido[1,2-<i>a</i>]pyrimidin-4-one, <i>N</i>-oxide. Bicyclorisperidone: 3-(4-Fluoro-2-hydroxyphenyl)-1-[2-(6,7,8,9-tetrahydro-2-methyl-4-oxo-4<i>H</i>-pyrido[1,2-<i>a</i>]pyrimidin-3-yl)ethyl]-2-aza-1-az

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		—						oniabicyclo[2.2.2]oct-2-ene iodide. Z-Oxime: (Z)-3-[2-[4-(2,4-Difluorophenyl)(hydroxyimino)methyl]-1-piperidinylethyl]-6,7,8,9-tetrahydro-2-methyl-4H-pyrido[1,2-a]pyrimidin-4-one
TIZANIDINE TABLETS	ASSAY/ Procedure/System suitability	Second Supplement to USP37–NF32	Online	25-Jul-2014	1-Aug-2014	USP39–NF34	First Supplement to USP38–NF33	Change Sample: Standard solution to: Samples: System suitability solution and Standard solution AND Change Resolution: NLT

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	—						4.0 between tizanidine and tizanidine related compound C; NLT 4.0 between tizanidine and tizanidine related compound B <i>Tailing factor:</i> NMT 2.0 <i>Relative standard deviation:</i> NMT 2.0% to: <i>Resolution:</i> NLT 4.0 between tizanidine and tizanidine related compound C; NLT 4.0 between tizanidine and tizanidine related compound B,

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		—						System suitability solution Tailing factor: NMT 2.0, Standard solution Relative standard deviation: NMT 2.0%, Standard solution
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
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	Line 1 of Solution A: Change sodium phosphate to: dibasic sodium

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CHLORPHENIRAMINE MALEATE	SPECIFIC TESTS/Optical Rotation, Specific Rotation <781>	Second Supplement to USP38–NF33	Online	25-Sep-2015	1-Oct-2015	USP40–NF35	First Supplement to USP39–NF34	phosphate Line 1: Delete Specific Rotation
RESIDUAL HOST CELL PROTEIN MEASUREMENT IN V 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

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		—						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 <i>Sample 3/HCP ratio</i> 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 <i>Sample 3/% max ratio value</i> column: Change 83% to: 61% Line 5: Change <i>NF category</i> : Coating agent; emollient; solvent; tablet and/or capsule
DESCRIPTION AND SOLUBILITY	SUNFLOWER OIL	<i>Second Supplement to USP38–NF33</i>	7761	20-Nov-2015	1-Dec-2015	<i>USP40–NF35</i>	<i>Second Supplement to USP39–NF34</i>	

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		—						diluent; vehicle (oleaginous). to: <i>NF category:</i> Coating agent; emollient; solvent; diluent; vehicle (oleaginous). Line 3 of <i>Sample stock solution:</i> Change Transfer the thimble to a continuous-extraction apparatus fitted with a 250-mL round-bottom flask containing 150 mL of hexane, and heat the flask on a heating mantle for 4 h. After the extraction, detach the round-bottom
MILK THISTLE	COMPOSITION <i>/Content of Silymarin</i>	<i>Second Supplement to USP38–NF33</i>	7878	20-Nov-2015	1-Dec-2015	<i>USP40–NF35</i>	<i>Second Supplement to USP39–NF34</i>	

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	—						flask from the extraction apparatus, and discard the hexane. Dry the extraction thimble to remove residual hexane, to: Transfer the thimble to a continuous-extraction apparatus fitted with a 250-mL round-bottom flask containing 150 mL of solvent hexane, and heat the flask on a heating mantle for 4 h. After the extraction, detach the round-bottom flask from the extraction apparatus, and

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		—						discard the solvent hexane. Dry the extraction thimble to remove residual solvent hexane, Line 3 of <i>Sample stock solution</i> : Change Transfer the thimble to a continuous-extraction apparatus fitted with a 250-mL round-bottom flask containing 150 mL of hexane, and heat the flask on a heating mantle for 4 h. After the extraction, detach the round-bottom flask from the extraction
POWDERED MILK THISTLE	COMPOSITION <i>/Content of Silymarin</i>	<i>Second Supplement to USP38–NF33</i>	7880	20-Nov-2015	1-Dec-2015	<i>USP40–NF35</i>	<i>Second Supplement to USP39–NF34</i>	

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	—						apparatus, and discard the hexane. Dry the extraction thimble to remove residual hexane, to: Transfer the thimble to a continuous-extraction apparatus fitted with a 250-mL round-bottom flask containing 150 mL of solvent hexane, and heat the flask on a heating mantle for 4 h. After the extraction, detach the round-bottom flask from the extraction apparatus, and discard the solvent hexane.

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CORTICOTRO ASSAY/ PIN INJECTION	<i>Second Supplement to USP38–NF33</i>	8059	20-Nov-2015	1-Dec-2015	<i>USP40–NF35</i>	<i>Second Supplement to USP39–NF34</i>	Dry the extraction thimble to remove residual solvent hexane, Line 7 of <i>Replication: Change (see <111>, Confidence Intervals for Individual Assays).</i> to: (see <111>, <i>The Confidence Interval and Limits of Potency</i>).
CORTICOTRO ASSAY/ PIN FOR INJECTION	<i>Second Supplement to USP38–NF33</i>	8061	20-Nov-2015	1-Dec-2015	<i>USP40–NF35</i>	<i>Second Supplement to USP39–NF34</i>	Line 7 of <i>Replication: Change (see <111>, Confidence Intervals for Individual Assays).</i> to: (see <111>, <i>The Confidence</i>

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REPOSITORY ASSAY/ CORTICOTRO Procedure PIN INJECTION	<i>Second Supplement to USP38–NF33</i>	8063	20-Nov-2015	1-Dec-2015	USP40–NF35	<i>Second Supplement to USP39–NF34</i>	<i>Interval and Limits of Potency).</i> Line 7 of <i>Replication:</i> Change (see <111>, <i>Confidence Intervals for Individual Assays).</i> to: (see <111>, <i>The Confidence Interval and Limits of Potency).</i>
LOPINAVIR AND RITONAVIR ORAL SOLUTION	IM PURITIES/ <i>Organic Impurities</i> <i>Second Supplement to USP38–NF33</i>	8139	20-Nov-2015	1-Dec-2015	USP40–NF35	<i>Second Supplement to USP39–NF34</i>	Row 17 of Column 3 of <i>Table 2:</i> Change 0.2 to: 0.2 ^p AND Add footnote ^p Disregard any peak less than 0.01%.
ELEMENTAL I DRUG	<i>Second</i>	7594	29-May-2015	1-Jun-2015	USP39–NF34	USP39–NF34	Line 1 of Row 1

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MPURITIES--LI MITS	SUBSTANCE AND E XCI PIENTS/ <i>Table 2</i>	<i>Supplement to USP38–NF33</i>						of Column 2: Change (g/g) to: (µg/g) AND Line 1 of Row 1 of Column 3: Change (g/g) to: (µg/g) AND Line 1 of Row 1 of Column 4: Change (g/g) to: (µg/g)
MOEXIPRIL HY IM DROCHLORID PUR E AND HYDRO ITIES/ <i>Organic</i> CHLOROTHIAZ <i>Impurities</i> IDE TABLETS		<i>Second Supplement to USP38–NF33</i>	8158	20-Nov-2015	1-Dec-2015	<i>USP40–NF35</i>	<i>Second Supplement to USP39–NF34</i>	Row 6 of Column 2 of <i>Table 5</i> : Change 0.62 to: 0.82
MANNITOL INJECTION	<i>Specific rotation <781></i>	<i>Second Supplement to USP38–NF33</i>	Online	20-Nov-2015	1-Dec-2015	<i>USP40–NF35</i>	<i>Second Supplement to USP39–NF34</i>	Line 1: Change Transfer an accurately

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	—						measured volume of Injection, equivalent to about 1 g of mannitol as determined by the Assay, to a 100-mL volumetric flask:it meets the requirements of the test for <i>Specific rotation</i> under <i>Mannitol</i>. to: +137° to +145°.Transfer an accurately measured volume of Injection, equivalent to about 1 g of mannitol as determined by the Assay, to a 100-mL volumetric flask.

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		—						Add 40 mL of a 1-in-10 ammonium molybdate solution, previously filtered if necessary. Add 20 mL of 1 N sulfuric acid, and dilute with water to volume.
SODIUM CHLORIDE	IM PURITIES/ <i>Limit of Potassium/Instrumental conditions</i>	<i>Second Supplement to USP39–NF34</i>	8821	27-Jan-2017	1-Feb-2017	<i>USP41–NF36</i>	<i>Second Supplement to USP40–NF35</i>	Line 1 of <i>Mode</i> : Change Atomic absorption spectrophotometry to: Atomic emission spectroscopy
OMEGA-3-ACID ETHYL ESTERS CAPSULES	SPECIFIC TESTS/ <i>Concentration of Omega-3-Acid Ethyl Esters</i>	<i>Second Supplement to USP39–NF34</i>	8755	18-Nov-2016	1-Dec-2016	<i>USP41–NF36</i>	<i>Second Supplement to USP40–NF35</i>	Line 21: Change C_U = nominal concentration of the total omega-3-acid ethyl esters in the <i>Sample</i>

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								<i>solution (g/mL) to:</i> C_U = Capsule fill content of the <i>Sample solution</i> (g/mL)
TACROLIMUS CAPSULES	PERFORMANCE TESTS/ <i>Dissolution <711>/Test 5</i>	<i>Second Supplement to USP39–NF34</i>	8834	27-Jan-2017	1-Feb-2017	<i>USP41–NF36</i>	<i>Second Supplement to USP40–NF35</i>	Line 1 of <i>Medium:</i> Change 0.5 g/L to: 0.05 g/L
POVIDONE	IMPURITIES/ <i>Formic Acid</i>	<i>Second Supplement to USP39–NF34</i>	8778	18-Nov-2016	1-Dec-2016	<i>USP41–NF36</i>	<i>Second Supplement to USP40–NF35</i>	Line 3 of <i>Sample solution:</i> Change column of about 80 mm to: column of about 8 mm
THEOPHYLLINE	ADDITIONAL REQUIREMENTS/ <i>USP Reference Standards RS <11></i>	<i>Second Supplement to USP39–NF34</i>	8844	27-Jan-2017	1-Feb-2017	<i>USP41–NF36</i>	<i>Second Supplement to USP40–NF35</i>	Line 3 of USP Theophylline Related Compound D RS: Change <i>N</i> -Methyl-5-(methanamine)-1 <i>H</i>

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	—						-imidazole-4-carboxamide. C ₆ H ₁₀ N ₄ O 154.17 to: N -Methyl-5-(meth ylami no)-1 <i>H</i> -imidazole-4-carboxamide hydrochloride monohydrate. C ₆ H ₁₀ N ₄ O · HCl · H ₂ O 208.65
ROPINIROLE E XTENDED- RELEASE TABLETS	PERFORMANC E TESTS/ <i>Dissolution</i> <711>/Test 1	<i>Second Supplement to USP39–NF34</i>	8814	18-Nov-2016	1-Dec-2016	<i>USP41–NF36</i>	<i>Second Supplement to USP40–NF35</i> Line 2 of <i>Buffer</i> 1: Adjust with <i>Solution A</i> to a pH of 4.0. to: Adjust with <i>Solution A</i> to a pH of 4.0. Dilute with water to 1 L.
THEOPHYLLIN E ORAL SOLUTION	ADDITIONAL R EQUIREMENT S/USP <i>Reference Standards</i> <11>	<i>Second Supplement to USP39–NF34</i>	8846	27-Jan-2017	1-Feb-2017	<i>USP41–NF36</i>	<i>Second Supplement to USP40–NF35</i> Line 3 of USP Theophylline Related Compound D RS: Change

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		—						<i>N</i> -Methyl-5-(methylamino)-1 <i>H</i> -imidazole-4-carboxamide. $C_6H_{10}N_4O$ 154.17 to: <i>N</i> -Methyl-5-(methylamino)-1 <i>H</i> -imidazole-4-carboxamide hydrochloride monohydrate. $C_6H_{10}N_4O \cdot HCl \cdot H_2O$ 208.65
DES LORATADINE	ADDITIONAL REQUIREMENTS/USP Reference Standards <11>	<i>Second Supplement to USP39–NF34</i>	8607	30-Sep-2016	1-Oct-2016	<i>USP41–NF36</i>	<i>First Supplement to USP40–NF35</i>	Line 2 of USP Desloratadine Related Compound B RS: Change 8-Chloro-11-(1,2,3,6-tetrahydropyridin-4-yl)-6,11-di hydro-5 <i>H</i> -benzo[5,6]cycl

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		—						ohe pta[1,2-<i>b</i>] pyridine. $C_{19}H_{19}ClN_2$ 310.82 to: 8-Chloro-11-(1, 2,3,6-tetrahydro pyridin-4-yl)-6,1 1-di hydro-5<i>H</i> -benzo[5,6]cycl ohe pta[1,2-<i>b</i>] pyridine hydrochloride. $C_{19}H_{20}Cl_2N_2$ 347.28
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: L = label claim of total omega-3-acids

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NAPROXEN SODIUM TABLETS	IMPURITIES/Organic Impurities	Second Supplement to USP39–NF34	Online	29-Jul-2016	1-Aug-2016	USP41–NF36	First Supplement to USP40–NF35	ethyl esters (mg/Capsule) Line 6 of Analysis: Change Result = $(r_U/r_S) \times (C_U/C_S) \times 100$ to: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$
REAGENTS, INDICATORS AND SOLUTIONS	Solutions/Volumetric Solutions	Second Supplement to USP39–NF34	8458	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	Line 3 of 0.07 N Ferrous Ammonium Sulfate VS: Change In a 100-mL volumetric flask, to: In a 1000-mL volumetric flask,
MAGNESIUM ALUMINUM SILICATE	SPECIFIC TESTS/Viscosity/Acceptance criteria	Second Supplement to USP39–NF34	8558	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	Line 1 of Type IA: Change 225–600 to: 225–600mPa · s AND Line 1 of Type IB: Change 150–450 to:

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ROPINIROLE E IM XTENDED- RELEASE TABLETS	PUR ITIES/ <i>Organic Impuri ties/Analysis</i>	<i>Second Supplement to USP39–NF34</i>	8814	31-Mar-2017	1-Apr-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>	150–450 mPa · s AND Line 1 of <i>Type /C</i> : Change 800–2200 to: 800–2200 mPa · s AND Line 1 of <i>Type /IA</i> : Change 100–300 to: 100–300 mPa · s Line 8 of the variable definition list of the first equation: Change C_U = nominal concentration of ropinirole related compound B free base in the <i>Sample solution</i> (µg/mL)

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		—						to: C_U = nominal concentration of ropinirole in the <i>Sample solution</i> (µg/mL) Line 6 of <i>Analysis</i> : Change $\text{Result}_i = (r_i/r_s) \times C_s \times [M \times (M_{r1}/M_{r2})] \times V \times (1/L) \times 100$ to: $\text{Result}_i = (r_i/r_s) \times C_s \times [M \times (M_{r1}/M_{r2})] \times D \times V \times (1/L) \times 100$ AND To the variable definition list in <i>Analysis</i> : Add <i>D</i> = dilution factor of the <i>Sample solution</i>
IMIPRAMINE PAMOATE CAPSULES	PERFORMANCE TESTS/ <i>Dissolution</i> <711>/Test 1	<i>Second Supplement to USP39–NF34</i>	8681	27-Jan-2017	1-Feb-2017	USP41–NF36	<i>Second Supplement to USP40–NF35</i>	Line 6 of <i>Analysis</i> : Change $\text{Result}_i = (r_i/r_s) \times C_s \times [M \times (M_{r1}/M_{r2})] \times V \times (1/L) \times 100$ to: $\text{Result}_i = (r_i/r_s) \times C_s \times [M \times (M_{r1}/M_{r2})] \times D \times V \times (1/L) \times 100$ AND To the variable definition list in <i>Analysis</i> : Add <i>D</i> = dilution factor of the <i>Sample solution</i>
PROMETHAZINE HYDROCHLORIDE	ADDITIONAL REQUIREMENTS/ USP <i>Reference Standards</i> <11>	<i>Second Supplement to USP39–NF34</i>	8784	27-Jan-2017	1-Feb-2017	USP41–NF36	<i>Second Supplement to USP40–NF35</i>	Line 2 of USP Promethazine Related Compound B RS: Change

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		—						Isopromethazine; <i>N,N</i> -
								<i>H</i> -phenothiazin-10-yl)propan-1-amine. $C_{17}H_{20}N_2S$ 282.42 to: Isopromethazine hydrochloride; <i>N,N</i> -
								<i>H</i> -phenothiazin-10-yl)propan-1-amine hydrochloride. $C_{17}H_{20}N_2S \cdot HCl$ 320.88
CIPROFLOXACIN	ADDITIONAL REQUIREMENT	<i>Second Supplement to</i>	8597	18-Nov-2016	1-Dec-2016	<i>USP41–NF36</i>	<i>Second Supplement to</i>	Line 2 of USP Ciprofloxacin Et

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
		<i>S/USP Reference Standards <11></i>					<i>USP40–NF35</i>	hylene diamine Analog RS: Change 7-(2-Aminoethyl amino)-1-cyclopropyl-6-fluoro-4-oxo-1,4-dihydroquinoline-3-carboxylic acid. $C_{15}H_{16}FN_3O_3$ 305.30 to: 1-Cyclopropyl-6-fluoro-1,4-dihydro-4-oxo-7-[(2-aminoethyl)amino]-3-quinolinecarboxylic acid hydrochloride. $C_{15}H_{16}FN_3O_3 \cdot HCl$ 341.77
PROMETHAZINE HYDROCHLORIDE INJECTION	ADDITIONAL REQUIREMENT	<i>Second Supplement to S/USP Reference Standards <11></i>	8785	27-Jan-2017	1-Feb-2017	<i>USP41–NF36</i>	<i>Second Supplement to USP40–NF35</i>	Line 2 of USP Promethazine Related Compound B RS: Change Isopromethazine; <i>N,N</i> -

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
		—						<i>H</i> -phenothiazin-1 0-yl)propan-1-a mine. C₁₇H₂₀N₂S 284.42 to: Isopromethazin e hydrochloride; <i>N,N</i> - <i>H</i> -phenothiazin-1 0-yl)propan-1-a mine hydrochloride. C₁₇H₂₀N₂S · HCl 320.88 Line 2 of USP Promethazine Related Compound B RS: Change Isopromethazin
PROMETHAZINE HYDROCHLORIDE ORAL SOLUTION	ADDITIONAL REQUIREMENT	<i>Second Supplement to USP Reference Standards <11></i>	8787	27-Jan-2017	1-Feb-2017	USP41–NF36	<i>Second Supplement to USP40–NF35</i>	

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
		—						e; N,N -
								H -phenothiazin-1 0-yl)propan-1-a mine. $C_{17}H_{20}N_2S$ 282.42 to: Isopromethazin e hydrochloride; N,N -
								H -phenothiazin-1 0-yl)propan-1-a mine hydrochloride. $C_{17}H_{20}N_2S \cdot HCl$ 320.88
DEXTROSE	IDENTIFICATIO N/C. Water Determination	Second Supplement to USP39–NF34	8612	18-Nov-2016	1-Dec-2016	USP41–NF36	Second Supplement to USP40–NF35	Line 1: Change Water Determination

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
		<921>						<921> to: Water Determination <921>, Method I
PROMETHAZINE HYDROCHLORIDE TABLETS	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	Second Supplement to USP39–NF34	8788	27-Jan-2017	1-Feb-2017	USP41–NF36	Second Supplement to USP40–NF35	Line 2 of USP Promethazine Related Compound B RS: Change Isopromethazine; N,N
								-
								H -phenothiazin-10-yl)propan-1-amine. C ₁₇ H ₂₀ N ₂ S 284.42 to: Isopromethazine hydrochloride; N,N
								-

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
		Sort descending						
								<i>H</i> -phenothiazin-10-yl)propan-1-amine hydrochloride. $C_{17}H_{20}N_2S \cdot HCl$ 320.88
OMEGA-3-ACID ETHYL ESTERS CAPSULES	ASSAY/Content of EPAee, DHAee, and Total Omega-3-Acid Ethyl Esters	Second Supplement to USP39–NF34	8755	18-Nov-2016	1-Dec-2016	USP41–NF36	Second Supplement to USP40–NF35	Line 21 of Analysis: Change C_U = nominal concentration of the total omega-3-acid ethyl esters in the <i>Sample solution</i> (g/mL) to: C_U = Capsule fill content of the <i>Sample solution</i> (g/mL)
CIPROFLOXACIN HYDROCHLORIDE	ADDITIONAL REQUIREMENT SI/USP Reference Standards <11>	Second Supplement to USP39–NF35	8600	18-Nov-2016	1-Dec-2016	USP41–NF36	Second Supplement to USP40–NF35	Line 2 of USP Ciprofloxacin Ethylenediamine Analog RS: Change 7-(2-Aminoethyl

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
		—						amino)-1-cyclopropyl-6-fluoro-4-oxo-1,4-dihydroquinoline-3-carboxylic acid. C ₁₅ H ₁₆ FN ₃ O ₃ 305.30 to: 1-Cyclopropyl-6-fluoro-1,4-dihydro-4-oxo-7-[(2-aminoethyl)amino]-3-quinolinecarboxylic acid hydrochloride. C ₁₅ H ₁₆ FN ₃ O ₃ · HCl 341.77
SODIUM LAURYL SULFATE	IDENTIFICATION	Second Supplement to USP40–NF35	8946	17-Nov-2017	1-Dec-2017	USP42–NF37	Second Supplement to USP41–NF36	Line 1 of A.: Change Infrared Absorption <197K> or <197A> to: ?A. Infrared Absorption <197K> or <197A>?
IDENTIFICATION TESTS—CHEMICAL IDENTIFICATION	IDENTIFICATION	Second Supplement to	Online	17-Nov-2017	1-Dec-2017	USP42–NF37	Second Supplement to	Line 1 of A.: Change

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
NERAL	TESTS/ <i>Thiosulfate</i>	— <i>USP40–NF35</i>					<i>USP41–NF36</i>	yellow; with the addition of sulfur dioxide, filter paper moistened with mercurous nitrate TS blackens. to: yellow, and evolve sulfur dioxide, which blackens filter paper moistened with mercurous nitrate TS.
CANDESARTAN CILEXETIL TABLETS	ADDITIONAL REQUIREMENTS/ <i>USP Reference Standards <11></i>	<i>Second Supplement to USP40–NF35</i>	8730	29-Sep-2017	1-Oct-2017	<i>USP42–NF37</i>	<i>First Supplement to USP41–NF36</i>	Line 2 of USP Candesartan Cilexetil Related Compound D RS: Change 1-[[[(Cyclohexyloxy)carbonyloxy]ethyl

Monograph Title Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
	—						-tetrazol-5-yl)biphenyl-4-yl]methyl}-2-oxo-2,3-dihydro-1 <i>H</i> -benzimidazole-4-carboxylate. to: 1-{[(Cyclohexyloxy)carbonyl]oxy}ethyl
FLUVOXAMINE CHEMICAL MALEATE	<i>Second Supplement to USP40–NF35</i> INFORMATION	8797	29-Sep-2017	1-Oct-2017	<i>USP42–NF37</i>	<i>First Supplement to USP41–NF36</i>	<i>H</i> -tetrazol-5-yl)-[1,1'-biphenyl]-4-yl]methyl)-2-oxo-2,3-dihydro-1 <i>H</i> -benzimidazole-4-carboxylate. Line 4:Change 5-Methoxy-4'-(trifluoromethyl)valerophenone (<i>E</i>)- <i>O</i> -(2-aminoethyl)oxime, maleate (1:1)

Monograph Title	Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
FLUVOXAMINE IM MALEATE	PUR ITIES/Organic Impurities/Table 1	Second Supplement to USP40–NF35	8797	29-Sep-2017	1-Oct-2017	USP42–NF37	First Supplement to USP41–NF36	to: (E)-5-Methoxy-4'- (trifluoromethyl) valerophenone O -(2-aminoethyl) oxime, maleate (1:1) Footnote b:Change 5-Methoxy-1-[4- (trifluoromethyl) phenyl]-1-penta none (E)-O -[2-[(2-succinyl) amino]ethyl]oxi me. to: (E)-5-Methoxy-1-[4-(trifluorometh yl)phenyl]-1-pen tanone O -[2-[(2-succinyl) amino]ethyl]oxi me. AND Footnotes c– g:

Monograph Title	Section	Source Publication Sort descending —	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
EFAVIRENZ	SPECIFIC TESTS/ <i>Enantiomeric Purity</i>	<i>Second Supplement to USP40–NF35</i>	Online	29-Sep-2017	1-Oct-2017	<i>USP42–NF37</i>	<i>First Supplement to USP41–NF36</i>	Delete the space before oxime Line 1 of <i>Mobile phase</i> : Change Hexane and ethanol (97:3) to: Hexane and absolute alcohol (97:3)
TIMOLOL MALEATE	MULTIPLE SECTIONS	<i>Second Supplement to USP40–NF35</i>	Online	28-Jul-2017	1-Aug-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>	The version of the Timolol Maleate monograph which appeared in the <i>Second Supplement to USP 40–NF 35</i> did not include the revisions approved in the version appearing in the <i>First Supplement to USP 40–NF 35</i> . The version appearing in the <i>First</i>

Monograph Title Section	Source Publication Sort descending	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
	—						<i>Supplement</i> should be used. The file as it should have appeared in the <i>Second Supplement</i> is attached to the compendial notice found at http://www.uspnf.com/notices/second-supplement-usp-40-nf-35-online-timolol-maleate.

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