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NEOTAME	ADDITIONAL R	<i>First</i>	8485	28-Jul-2017	1-Aug-2017	<i>USP42–NF37</i>	<i>First</i>	Line 2 of USP

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	EQUIREMENT S/USP Reference Standards <11>	<i>Supplement to USP40–NF35</i>						<i>Supplement to USP41–NF36</i>	Neotame Related Compound A RS: Change <i>N</i> -[3,3-Dimethylbutyl)-L-?-aspartyl]-L-phenylalanine. to: <i>N</i> -[<i>N</i> -(3,3-Dimethylbutyl)-L-?-aspartyl]-L-phenylalanine.
POTASSIUM CITRATE EXTENDED-RELEASE TABLETS	OTHER COMPONENTS/Content of Potassium	<i>Revision Bulletin (Official April 01, 2017)</i>	Online	28-Jul-2017		1-Aug-2017	<i>USP42–NF37</i>	<i>First Supplement to USP41–NF36</i>	Line 1 of <i>Sample solution</i> : Change Transfer 3.0 mL of the clear filtrate, reserved from the Assay, to a 100-mL volumetric flask. to: <i>Sample stock solution</i> : Dilute the clear filtrate, reserved from the Assay, with

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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>	water to obtain a solution containing about 160 µg/mL of potassium citrate monohydrate. <i>Sample solution:</i> Transfer 3.0 mL of the <i>Sample stock solution</i> to a 100-mL volumetric flask. 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>.

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									The version appearing in the <i>First 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.
FLOW CYTOMETRIC ENUMERATION OF CD34+ CELLS	SAMPLE PREPARATION/ <i>Table 1. Dot Plot Descriptions and Gating Instructions</i>	USP40–NF35	216	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Column 4 for Step 6: Add • For counting bead enumeration, proceed to either Step 7 or Step 8 according to the recommendations of the bead manufacturer.

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OLIGOSACCHARIDE ANALYSIS	SEPARATION AND IDENTIFICATION OF OLIGOSACCHARIDES	USP40–NF35	273	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Line 1 of <i>Normal Phase Chromatography/HILIC/Ammonium formate buffer</i> : Change Add 1.4 M ammonia solution to 1.4 M formic acid solution. to: Add 1.4 M ammonia solution to 1.4 M formic acid solution until a pH of 4.4 is obtained.
TOPICAL AEROSOLS	DELIVERED-DOSE UNIFORMITY	USP40–NF35	499	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Line 2: Change proceed as directed in the test for <i>Delivered Dose Uniformity</i> in <i>Metered-Dose Inhalers and Dry Powder Inhalers</i> , as described in

Monograph Title Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							<601>, to: proceed as directed in <i>Inhalation and Nasal Drug Products: Aerosols, Sprays, and Powders—Performance Quality Tests</i> <601>, A. <i>Delivered-Dose Uniformity/A.2 Inhalation Aerosols and Inhalation Sprays</i> and A.4 <i>Inhalation Powders</i>, AND Line 4: Delete Unless otherwise stated in the individual monograph, apply the acceptance criteria for <i>Metered-Dose</i>

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NUCLEIC ACID-BASED TECHNIQUES—GENERAL	USP40–NF35	1471	26-May-2017	1-Jun-2017	USP41–NF36	USP41–NF36	<i>Inhalers and Dry Powder Inhalers as described in <601>.</i> Line 1 of bullet 1: Change "Review Criteria for Nucleic Acid Amplification-Based In Vitro Diagnostic Devices for Direct Detection of Infectious Microorganisms" (1993) to: "Nucleic Acid Based In Vitro Diagnostic Devices for Detection of Microbial Pathogens" (2005)
ACETYLCYSTEINE SOLUTION Procedure	USP40–NF35	2586	26-May-2017	1-Jun-2017	USP41–NF36	USP41–NF36	Line 3 of <i>Sample solution</i>: Change <i>Standard stock</i>

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									<i>solution to:</i>
									<i>Sample stock solution</i>
CONSTRUCT HUMAN FIBROBLASTS IN BILAYER SYNTHETIC SCAFFOLD	SPECIFIC TESTS/ <i>Metabolic Activity Assessment</i>	USP40–NF35	3561	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Line 2 of <i>L-Glutamine solution</i> : Change 29.2 g to: 2.92 g
GLUCONOLAC TONE	ASSAY/ <i>Procedure</i>	USP40–NF35	4412	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Line 5 of <i>Analysis</i> : Change Each mL of <i>Back-titrant</i> to: Each milliliter of <i>Titrant</i>
IOHEXOL	IM PURITIES/ <i>Limit of Free Iodide/Analysis</i>	USP40–NF35	4649	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Line 3 of variable definition list: Change 0.1269 mg/mEq to: 126.9 mg/mEq
MYCOPHENOLATE MOFETIL FOR ORAL SUSPENSION	ASSAY/ <i>Procedure</i>	USP40–NF35	5251	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Line 3 of <i>Sample solution</i> : Change 45-µm pore

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PHENYTOIN ORAL SUSPENSION	IM PUR ITIES/ <i>Organic Impurities</i>	USP40–NF35	5690	26-May-2017	1-Jun-2017	USP41–NF36	USP41–NF36	size. to: 0.45-µm pore size.
								Line 1 of <i>Sample solution</i> : Change 1 mg/mL of Oral Suspension in <i>Diluent</i> to: Nominally 1 mg/mL of phenytoin prepared as follows. Weigh and transfer a suitable volume of Oral Suspension to an appropriate volumetric flask. Add methanol to about 20% of the final flask volume and dissolve. Dilute with <i>Diluent</i> to volume. Dissolve with

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PIOGLITAZONE AND METFORMIN HYDROCHLORIDE TABLETS	PERFORMANCE TESTS/ <i>Dissolution</i> <711>/Test 2	USP40–NF35	5720	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	the aid of sonication, if necessary. Line 3 of <i>Standard solution</i> : Change USP Metformin Hydrochloride RS in <i>Diluent A</i> to: USP Metformin Hydrochloride RS from the <i>Metformin standard stock solution</i> in <i>Diluent A</i>
POLYETHYLENE GLYCOL 3350	SPECIFIC TESTS/ <i>Apparent Weight-Average Molecular Weight and Polydispersity</i>	USP40–NF35	5745	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Line 1 of <i>Standard solution</i> : Change <i>Standard solution</i> : 1.0 mg/mL each of five polyethylene glycol standards with molecular weights of 1000,2000,

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							3000, 4000, and 6000 g/mol (Da) in <i>Mobilephase</i>. Pass a portion of the solution to: <i>Standard solutions:</i> Prepare 1.0 mg/mL each of five polyethylene glycol standards with molecular weights of 1000,2000, 3000, 4000, and 6000 g/mol (Da) in <i>Mobilephase</i> separately in five individual flasks. Pass a portion of each solution AND Line 1 of <i>Analysis/Samples:</i> Change <i>Standard solution</i> and

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									<i>Sample solution to:</i> <i>Standard solutions and Sample solution AND</i> Line 3 of <i>Analysis:</i> Change Separately inject equal volumes of the <i>Standard solution</i> to: Separately inject equal volumes of the <i>Standard solutions</i> Line 5 of <i>Carbohydrate Content/ Acceptance criteria:</i> Change <i>Moisture Content</i> to: <i>Loss on Drying</i> AND Line 3 of <i>Suture</i>
SCAFFOLD BOVINE DERMIS	SPECIFIC TESTS	USP40–NF35	6113	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	

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									Retention Force/Analysis: Change 40 to: 4-0
EVENING PRIMROSE CAPSULES	ST OILREN GTH/Analysis	USP40–NF35	6952	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	In the first variable definition list: Change m_S = weight of USP Methyl Ester RS in the Standard solution (mg) to: m_S = weight of the relevant USP Methyl Ester RS in the Standard solution (mg)
SUCROSE	SPECIFIC TESTS/Color Value/Analysis	USP40–NF35	7938	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Line 6 of the variable definition list: Change The absolute difference between two results is NMT 3.

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									to: <i>Suitability requirements</i> <i>Repeatability:</i> The absolute difference between two results is NMT 3.
METHIONINE	IMPURITIES/Related Compounds	<i>First Supplement to USP40–NF35</i>	8337	26-May-2017		1-Jun-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>	Row 5 of Column 3 of <i>Table 2</i> : Change 0.1 to: 1.0
METHYLDOPA	ASSAY/Procedure/Chromatographic system	<i>First Supplement to USP40–NF35</i>	8339	26-May-2017		1-Jun-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>	Line 1 of <i>Injection volume</i> : Change 1 mL to: 20 µL
TIMOLOL MALEATE	IDENTIFICATION/B.	<i>First Supplement to USP40–NF35</i>	8416	26-May-2017		1-Jun-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>	Line 1: Change major peak of the <i>Sample solution</i> corresponds to that of the <i>Standard solution</i> ,

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									to: timolol peak of the <i>Sample solution</i> corresponds to that of the <i>System suitability solution</i> , Line 6 of <i>Chromatographic system</i> : Delete <i>Autosampler temperature</i> : 4°
TIMOLOL MALEATE	IM PURITIES/ <i>Enantiomeric Purity</i>	<i>First Supplement to USP40–NF35</i>	8416	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	
DOCETAXEL	IM PURITIES/ <i>Organic Impurities, Procedure 1</i>	<i>Revision Bulletin (Official August 01, 2016)</i>	Online	26-May-2017		1-Jun-2017	USP41–NF36	USP41–NF36	Footnote c of <i>Table 2</i> : Change (2aR,4R,4aS,9S,11S,12S,12aR,12bS)-1,2a,3,4,4a,6,9,10,11,12,12a,12b-Dodecahydro-4,9,11,12,12b-pentahydroxy-4a,8,13,13-tetramethyl-7,11-methano-5H-cyclodeca[3,4]benz[1

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							,2- <i>b</i>]oxet-5,6-dione 12b-acetate, 12-benzoate, 9-ester with (2 <i>R</i> ,3 <i>S</i>)- <i>N</i> - -formyl-3-phenyl isoserine. to: (2 <i>aR</i> ,4 <i>S</i> ,4 <i>aS</i> ,6 <i>R</i> ,9 <i>S</i> ,11 <i>S</i> ,12 <i>S</i> ,12 <i>aR</i> ,12 <i>bS</i>)?12b?Acetoxy? 9?(((2 <i>R</i> ,3 <i>S</i>)?3?formamido? 2?hydroxy?3?ph enylpropanoyl)o xy)?4,6,11?trihy droxy?4 <i>a</i> ,8,13,1 3?tetramethyl?5 ?oxo?2 <i>a</i> ,3,4,4 <i>a</i> , 5,6,9,10,11,12,1 2 <i>a</i> ,12 <i>b</i> ?dodeca hydro?1 <i>H</i> ?7,11?methano cyclodeca[3,4] <i>b</i> enzo[1,2?b]oxet-12?yl benzoate.

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							AND Footnote d of <i>Table 2</i>: Change (2aR,4R,4aS,6R,9S,11S,12S,12aR,12bS)-6-[[[(2,2-Dichloroethoxy)carbonyl]oxy]-1,2a,3,4,4a,6,9,10,11,12,12a,12b-dodecahydro-4,9,11,12,12b-pentahydroxy-4a,8,13,13-tetramethyl-7,11-methano-5H-cyclodeca[3,4]benz[1,2-b)oxet-5-one 12b-acetate, 12-benzoate, 9-ester with (2R,3S)-N-tert-butoxycarbonyl-L-phenylisoserine. to: (2aR,4S,4aS

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DOBUTAMINE INJECTION	IDENTIFICATION	USP39–NF34	3561	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	,6R,9S,11S ,12S,12aR ,12bS)?6?{[(2,2?Dichl oroethoxy)carbo nyl]oxy}?1,2a,3, 4,4a,6,9,10,11,1 2,12a,12b?dode cahydro?4,9,11, 12,12b?pentahy droxy?4a,8,13,1 3?tetramethyl?7 , 1 1? meth ano?5H ?cyclodeca[3,4] benz[1,2?b]oxet?5?one 12b?acetate, 12?benzoate, 9?ester with (2R,3S)?N?tert- butoxycarbonyl? 3?phenylisoseri ne. Line 1 of Sample solution:

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									Change 10 mg/mL of dobutamine hydrochloride in methanol, clarified by centrifugation to: Use the neat Injection.
EZETIMIBE	IM PURITIES/ <i>Organic Impurities, Procedure 1/System suitability/Suitability requirements</i>	USP39–NF34	3840	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Line 2 of <i>Resolution</i> : Change <i>Standard solution</i> to: <i>System suitability solution</i>
MYCOPHENOLATE MOFETIL	ADDITIONAL REQUIREMENT <i>S/USP Reference Standards <11></i>	USP39–NF34	4957	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Line 5 of USP Mycophenolate Mofetil Related Compound A RS: Change $C_{23}H_{31}NO_7$ to: $C_{22}H_{29}NO_7$
MYCOPHENOLATE MOFETIL FOR INJECTION	ADDITIONAL REQUIREMENT <i>S/USP Reference</i>	USP39–NF34	4960	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Line 5 of Mycophenolate Mofetil Related Compound A

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<i>Standards <11></i>									RS: Change $C_{23}H_{31}NO_7$ to: $C_{22}H_{29}NO_7$
MYCOPHENOL ADDITIONAL R ATE MOFETIL EQUIREMENT FOR ORAL S/USP SUSPENSION Reference <i>Standards <11></i>		USP39–NF34	4961	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Line 5 of Mycophenolate Mofetil Related Compound A RS: Change $C_{23}H_{31}NO_7$ to: $C_{22}H_{29}NO_7$
SORBITOL SOLUTION	ASSAY/ Proce dure/Analysis	USP39–NF34	5897	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	In the variable definition list: Change C_U = nominal concentration of Sorbitol Solution in the <i>Sample solution</i> (mg/g) to: C_U = concentration of Sorbitol Solution in the <i>Sample solution</i> (mg/g)
VINORELBINE TARTRATE	ADDITIONAL R EQUIREMENT S/USP	USP39–NF34	6370	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Line 2 of USP Vinorelbine Related

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	<i>Reference Standards <11></i>								Compound A RS: Change 4-O -Deacetylvinorel bine. to: 4-O -Deacetylvinorel bine tartrate.
VINOIRELBINE INJECTION	ADDITIONAL R EQUIREMENT <i>S/USP Reference Standards <11></i>	USP39–NF34	6371	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Line 2 of USP Vinorelbine Related Compound A RS: Change 4-O -Deacetylvinorel bine. to: 4-O -Deacetylvinorel bine tartrate.
POWDERED MILK THISTLE EXTRACT	COMPOSITION <i>/Content of Silymarin</i>	USP39–NF34	6764	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Row 2 of Column 2 of <i>Table 1:</i> Change 0 to: 85 AND Row 2 of Column 3 of

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									<i>Table 1:</i> Change 0 to: 15 Row 2 of Column 2 of <i>Table 1:</i> Change 0 to: 85 AND Row 2 of Column 3 of <i>Table 1:</i> Change 0 to: 15
MILK THISTLE CAPSULES	ST REN GTH/ <i>Content of Silymarin</i>	USP39–NF34	6765	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Row 2 of Column 2 of <i>Table 1:</i> Change 0 to: 85 AND Row 2 of Column 3 of <i>Table 1:</i> Change 0 to: 15
MILK THISTLE TABLETS	ST REN GTH/ <i>Content of Silymarin</i>	USP39–NF34	6767	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Row 2 of Column 2 of <i>Table 1:</i> Change 0 to: 85 AND Row 2 of Column 3 of <i>Table 1:</i>

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MAGNESIUM A ASSAY/ LUMINOMETA Magnesium SILICATE Oxide	USP39–NF34	7375	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	Change 0 to: 15 Line 5 of Analysis: Change eriochrome black TS trituration to: eriochrome black T trituration
MINOCYCLINE IM HYDROCHLOR PUR IDE EXTENDE ITIES/Organic D-RELEASE Impurities TABLETS	First Supplement to USP39–NF34	8101	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	Change Buffer, Mobile phase, Diluent, and Sample solution: Prepare as directed in the Assay. to: Buffer, Mobile phase, and Diluent: Prepare as directed in the Assay. AND Add Sample

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REAGENTS, INDICATORS AND SOLUTIONS	<i>Solutions/Volumetric Solutions</i>	<i>Second Supplement to USP39–NF34</i>	8458	31-Mar-2017	1-Apr-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>		<i>solution:</i> Use the <i>Sample stock solution</i> as directed in the Assay. Line 3 of <i>0.07 N Ferrous Ammonium Sulfate VS:</i> Change In a 100-mL volumetric flask, to: In a 1000-mL volumetric flask,
MAGNESIUM ALUMINUM SILICATE	SPECIFIC TESTS/ <i>Viscosity/Acceptance criteria</i>	<i>Second Supplement to USP39–NF34</i>	8558	31-Mar-2017	1-Apr-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>		Line 1 of <i>Type IA:</i> Change 225–600 to: 225–600mPa · s AND Line 1 of <i>Type IB:</i> Change 150–450 to: 150–450 mPa · s AND Line 1 of <i>Type IC:</i> Change 800–2200

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									to: 800–2200 mPa · s AND Line 1 of <i>Type IIA</i> : Change 100–300 to: 100–300 mPa · s	
ROPINIROLE	EXTENDED-RELEASE TABLETS	IMPURITIES/ <i>Organic Impurities/Analysis</i>	<i>Second Supplement to USP39–NF34</i>	8814		31-Mar-2017	1-Apr-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>	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) to: C_U = nominal concentration of ropinirole in the <i>Sample solution</i> (µg/mL)
RIBOFLAVIN	ASSAY		<i>USP40–NF35</i>	386		31-Mar-2017	1-Apr-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>	Line 2 of the

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ASSAY									variable definition list in <i>Chemical Methods, Procedure 1/Analysis</i> : Change sodium hydrofluorite to: sodium hydrosulfite
SPACERS AND VALVED HOLDING CHAMBERS USED WITH INHALATION AEROSOLS—CHARACTERIZATION TESTS	1. INTRODUCTION/1.5 Definitions of Key Terms Relating to This Chapter	USP40–NF35	1988	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Bottom right corner of <i>Figure 1</i> : Change VHC mouthpiece to: Spacer mouthpiece
PENICILLIN G PROCAINE	SPECIFIC TESTS/Content of Penicillin G and Procaine/Analysis	USP40–NF35	5607	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Line 7 of the variable definition list for the first equation: Change $C_U =$ concentration of Penicillin G Procaine in the

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							<i>Sample solution</i> (mg) to: C_U = concentration of Penicillin G Procaine in the <i>Sample solution</i> (mg/mL) AND Line 7 of the variable definition list for the second equation: Change C_U = concentration of Penicillin G Procaine in the <i>Sample solution</i> (mg) to: C_U = concentration of Penicillin G Procaine in the <i>Sample solution</i> (mg/mL)
RINGER'S	ASSAY	USP40–NF35 6019	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	Line 1 of

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INJECTION							<i>Calcium/ Instrumental condit ions/Mode:</i> Change spectrophotometer to: spectrophotometry AND Line 1 of <i>Potas sium/Solution A:</i> Change Suitable wetting agent to: Suitable nonionic wetting agent
DESCLORATADIM NE ORALLY DI PUR SINTEGRATIN ITIES/ G TABLETS <i>Organic Impurities</i>	<i>First Supplement to USP40–NF35</i>	Online	31-Mar-2017	1-Apr-2017	<i>USP41–NF36</i>	<i>USP41–NF36</i>	Line 5 of <i>Sample solution:</i> Change Add 70% of the flask volume of <i>Mobile phase</i> to: Add 70% of the flask volume of

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
GRANISETRO N HYDROCHLORIDE INJECTION	USP Reference standards <11>	First Supplement to USP40–NF35	Online	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	<i>Diluent</i> Line 3 of USP Granisetron Related Compound C RS: Change carboxamide. to: carboxamide hydrochloride.
GRANISETRO N HYDROCHLORIDE TABLETS	USP Reference standards <11>	First Supplement to USP40–NF35	Online	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Line 3 of USP Granisetron Related Compound C RS: Change carboxamide. to: carboxamide hydrochloride.
ALMOTRIPTAN TABLETS	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	Revision Bulletin (Official February 01, 2017)	Online	31-Mar-2017		1-Apr-2017	USP41–NF36	USP41–NF36	Line 4 of USP Almotriptan Related Compound B RS: Change $\frac{1}{2}\text{C}_4\text{H}_4\text{O}$ to: $\frac{1}{2}\text{C}_4\text{H}_4\text{O}_4$
ADENINE	CHEMICAL INFORMATION	USP39–NF34	2346	27-Jan-2017		1-Feb-2017	USP41–NF36	Second Supplement to USP40–NF35	Line 2: Change 1H -Purin-6-amine; to:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
AMINOPHYLLINE TABLETS	PERFORMANCE TESTS/ <i>Uniformity of Dosage Units</i> <905>/ <i>Procedure for content uniformity</i>	USP39–NF34	2483	27-Jan-2017		1-Feb-2017	USP41–NF36	<i>Second Supplement to USP40–NF35</i>	9H -Purin-6-amine; Variable definition for C_U in <i>Analysis</i> : Change (mg/mL) to: to:(µg/mL)
ATROPINE SULFATE	DEFINITION	USP39–NF34	2638	27-Jan-2017		1-Feb-2017	USP41–NF36	<i>Second Supplement to USP40–NF35</i>	Line 2: Change ($C_{17}H_{23}NO_{32} \cdot H_2SO_4$), to: [($C_{17}H_{23}NO_3$) ₂ · H_2SO_4],

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