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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”
  - A search for “Aminosalicyclic Acid Tablets” will result in anything that specifically contains “Aminosalicyclic Acid Tablets”
- **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.
  - For example, page 2178 will come before page 74 on a page sort.
- **Downloading:** You can download the Errata table in Comma-separated Value (.csv). The download will include the Errata that you have filtered on.
- **Importing:** You will need to import the file into Excel or Open Office with UTF-8 encoding, as opposed to simply opening it. To import, open Excel or Open Office and select import from the File drop-down. Depending on the version you are using, you should be presented with import formatting options to include UTF-8 as one of the first steps. Importing via UTF-8 should eliminate odd character conversions.

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GLYCERYL TRIASSAY/ <i>Content</i>	<i>First</i>	Online	28-Sep-2018	1-Oct-2018	<i>USP43–NF38</i>	<i>Second</i>	This erratum

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CAPRYLATE	<i>of Triglycerides</i>	<i>Supplement to USP41–NF36</i>						<i>Supplement to USP41–NF36</i>	applies to the new USP-NF ONLINE platform only. Line 1 of <i>System suitability solution</i> : Change 20 mUSP Glyceryl Monocaprylate RSUSP Glyceryl Monocaprylate RSg/mL each of 1-monooctanoyl-rac-glycerol and 1-monodecanoyl-rac-glycerol in tetrahydrofuran to: 20 mg/mL each of 1-monooctanoyl-rac-glycerol and 1-monodecanoyl-rac-glycerol in tetrahydrofuran In the <i>Analysis</i> : Change
METHADONE ASSAY/ HYDROCHLOR	<i>Procedure</i>	<i>USP41–NF36</i>	2628	28-Sep-2018		1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to</i>	

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IDE INJECTION								USP41–NF36	Result = $(R_U/R_S) \times W \times 100$ to: Result = $(R_U/R_S) \times W$
NIFEDIPINE EXPERFORMANC TENDED- RELEASE TABLETS	E TESTS/ Dissolution <711>/Test 5	First Supplement to USP41–NF36	8369	28-Sep-2018		1-Oct-2018	USP43–NF38	Second Supplement to USP41–NF36	Line 1 of Standard stock solution: Change 50 mg of USP Nifedipine RS in Diluent A and water (50:50) to: 0.50 mg/mL of USP Nifedipine RS prepared as follows. Transfer a suitable amount of USP Nifedipine RS to an appropriate volumetric flask. Dissolve in 50% of the flask volume of Diluent A. Dilute with water to volume.

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PRAVASTATIN SODIUM	ADDITIONAL REQUIREMENTS	<i>First Supplement to USP41–NF36</i>	Online	28-Sep-2018		1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	<p>AND Line 1 of <i>Instrumental conditions</i>/ <i>Analytical wavelength</i>: Change 238 nm to: 338 nm</p> <p>This erratum applies to the new <i>USP-NF ONLINE</i> platform only. Line 2 of <i>USP Reference Standards</i> &lt;11&gt;: Change <i>USP Pravastatin Related Compound A RS</i> to: <i>USP Pravastatin 1,1,3,3-Tetramethyl butylamine RS</i></p>
TRIAMTERENE	ADDITIONAL REQUIREMENTS	<i>First Supplement to</i>	Online	28-Sep-2018		1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to</i>	This erratum applies to the

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	S	USP41–NF36						USP41–NF36	new USP-NF ONLINE platform only. Line 4 of <i>USP Reference Standards</i> <11>: Change USP Doxazosin Related Compound A RS to: USP Triamterene Related Compound A RS
ORDINARY IMPURITIES	KEY FOR VISUALIZATION TECHNIQUES	USP41–NF36	Online	28-Sep-2018		1-Oct-2018	USP43–NF38	<i>Second Supplement to USP41–NF36</i>	This erratum applies to the new USP-NF ONLINE platform only. Change 3. <i>Solution A–Mix</i> 850 mg of bismuth subnitrate with 40 mL of water and 10 mL of glacial acetic acid.

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							<p>4. <i>Solution B</i>—Dissolve 8 g of potassium iodide in 20 mL of water. Mix A and B together to obtain a Stock Solution which can be stored for several months in a dark bottle. Mix 10 mL of the Stock Solution with 20 mL of glacial acetic acid, and dilute with water to make 100 mL, to prepare the spray reagent.</p> <p>to:</p> <p>3. <i>Solution A</i>—Mix 850 mg of bismuth subnitrate with 40 mL of water and 10 mL of glacial acetic acid.</p>

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LOXAPINE SUCCINATE	IM PUR ITIES/ <i>Organic</i>	<i>USP41–NF36</i>	2486	28-Sep-2018		1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	<p><i>Solution</i>  <i>B</i>–Dissolve 8 g of potassium iodide in 20 mL of water. Mix A and B together to obtain a Stock Solution which can be stored for several months in a dark bottle. Mix 10 mL of the Stock Solution with 20 mL of glacial acetic acid, and dilute with water to make 100 mL, to prepare the spray reagent.  AND  Update list numbers 5–23 to: 4–22  In the variable definition list in <i>Analysis</i>: Add</p>

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									F = relative response factor (see <i>Impurity Table 1</i> )
	<i>Impurities/Procedure</i>								
INSULIN ASSAYS	ADDITIONAL REQUIREMENTS/USP Reference Standards <11>	USP41–NF36	6054	28-Sep-2018		1-Oct-2018	USP43–NF38	Second Supplement to USP41–NF36	This erratum applies to the new USP-NF ONLINE platform only. Delete USP Insulin RS
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE OPHTHALMIC SOLUTION	IMPURITIES/Organic Impurities: Dorzolamide Hydrochloride	First Supplement to USP41–NF36	Online	28-Sep-2018		1-Oct-2018	USP43–NF38	Second Supplement to USP41–NF36	This erratum applies to the new USP-NF ONLINE platform only. In the first variable definition list in <i>Analysis</i> : Change $C_S =$ concentration of USP Doxazosin Related Compound D RS in the <i>Standard solution</i> (mg/mL) to:



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							<p><math>C_S</math> = concentration of USP Dorzolamide Related Compound D RS in the <i>Standard solution</i> (mg/mL) AND In the second variable definition list in <i>Analysis</i>: Change <math>C_S</math> = concentration of USP Doxazosin Related Compound B RS in the <i>Standard solution</i> (mg/mL) to: <math>C_S</math> = concentration of USP Dorzolamide Related</p>

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TRANLYCYPR OMINE TABLETS	PERFORMANC E TESTS/ Dissolution <711>	<i>First Supplement to USP41–NF36</i>	Online	28-Sep-2018		1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	Compound B RS in the <i>Standard solution</i> (mg/mL) This erratum applies to the new <i>USP-NF ONLINE</i> platform only. Change <i>System suitability</i> ? ?1S ( <i>USP41</i> ) <i>Sample: Standard solution Suitability requirements Tailing factor.</i> NMT 2.0 to: <i>System suitability Sample: Standard solution Suitability requirements</i> ? ?1S ( <i>USP41</i> ) <i>Tailing factor.</i>

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ISOLEUCINE	IM PUR ITIES/ <i>Related Compounds</i>	<i>First Supplement to USP41–NF36</i>	Online	28-Sep-2018	1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	NMT 2.0 This erratum applies to the new <i>USP-NF ONLINE</i> platform only. Line 1 of <i>Standard solution</i> : Change USP L-Leucine RS to: USP L-Isoleucine RS In the variable definition in <i>Analysis</i> : Change $C_S =$ concentration of USP Cyclophosphamide RS in the <i>Standard solution</i> (mg/mL). [Note—Concentration is calculated on the anhydrous basis.] C
CYCLOPHOSPHAMIDE	ASSAY/ <i>Procedure</i>	<i>USP41–NF36</i>	1123	28-Sep-2018	1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	

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NAPROXEN TABLETS	IM PURITIES/ <i>Organic Impurities</i>	USP41–NF36	2865	28-Sep-2018		1-Oct-2018	USP43–NF38	<i>Second Supplement to USP41–NF36</i>	<p>U = concentration of Cyclophosphamide in the <i>Sample solution</i> (mg/mL).  [Note—Nominal concentration is calculated on the anhydrous basis.]  to:  C<sub>S</sub> = concentration of USP Cyclophosphamide RS in the <i>Standard solution</i> (mg/mL)  C<sub>U</sub> = concentration of Cyclophosphamide in the <i>Sample solution</i> (mg/mL)</p> <p>Line 3 of <i>System suitability solution</i>:  Change 0.5 mg/mL of</p>

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VITAMIN D ASSAY	ASSAY/ Procedure 8	<i>First Supplement to USP41–NF36</i>	Online	28-Sep-2018		1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	<p>USP Naproxen RS and 0.5 µg/mL of USP Naproxen Related Compound A RS in <i>Diluent</i>, from <i>Standard stock solution 1</i> and <i>Standard stock solution 2</i>, respectively to:</p> <p>0.5 µg/mL of USP Naproxen Related Compound A RS from <i>Standard stock solution 2</i> and 0.5 mg/mL of USP Naproxen RS in <i>Diluent</i></p> <p>Line 1 of <i>Aqueous potassium hydroxide solution</i>: Change 800 mg to:</p>

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SUMATRIPTAN ASSAY/ NASAL SPRAY <i>Procedure</i>	<i>First Supplement to USP41–NF36</i>	Online	28-Sep-2018	1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	800 g This erratum applies to the new <i>USP-NF ONLINE</i> platform only. In the variable definition in <i>Analysis</i> : Change $C_S =$ concentration of USP Sumatriptan Succinate Related Compound A RS to: $C_S =$ concentration of USP Sumatriptan Succinate RS
TRANEXAMIC ACID INJECTION SPECIFIC TESTS	<i>First Supplement to USP41–NF36</i>	Online	28-Sep-2018	1-Oct-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	This erratum applies to the new <i>USP-NF ONLINE</i> platform only. Change <i>Dissolution</i>

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ACETAMINOPHEN ORAL SUSPENSION	IM PURITIES/Organic Impurities	<i>Revision Bulletin (Official August 01, 2018)</i>	Online	28-Sep-2018		1-Oct-2018	USP43–NF38	<i>Second Supplement to USP41–NF36</i>	<p>&lt;711&gt;: Meets the requirements to: <i>Sterility Tests</i></p> <p>&lt;71&gt;: Meets the requirements</p> <p>This erratum applies to the new USP-NF ONLINE platform only. Change <i>Solution B</i>: 0.2% trifluoroacetic acid in water to: <i>Solution B</i>: 0.2% trifluoroacetic acid in acetonitrile</p> <p>Change <i>Test preparation</i>: Into a 50-mL color-comparison tube place 10.0 mL of <i>Solution S</i>.</p>
ELASTOMERIC CLOSURES FOR INJECTIONS	PHYSICOCHEMICAL TESTS/Heavy Metals	<i>Revision Bulletin (Official January 01, 2018)</i>	Online	31-Aug-2018		1-Sep-2018	USP43–NF38	<i>Second Supplement to USP41–NF36</i>	<p>Change <i>Test preparation</i>: Into a 50-mL color-comparison tube place 10.0 mL of <i>Solution S</i>.</p>

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AMIKACIN SULFATE INJECTION	ASSAY/ <i>Procedure</i>	USP41–NF36 203	31-Aug-2018	1-Sep-2018	USP43–NF38	<i>Second Supplement to USP41–NF36</i>	<p>to: <i>Test preparation:</i> Into a 50-mL color-comparison tube pipet 10.0 mL of <i>Solution S</i> and dilute with water to 25 mL. Using a pH meter or short-range pH indicator paper as external indicator, adjust with 1 N acetic acid or 6 N ammonium hydroxide to a pH of between 3.0 and 4.0, dilute with water to 40 mL, and mix.</p> <p>Variable definition list in <i>Analysis:</i> Change <math>C_U</math> = nominal concentration of amikacin in the</p>



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FISH OIL CONTAINING OMEGA-3 ACIDS CAPSULES	DEFINITION	USP41–NF36	4620	31-Aug-2018		1-Sep-2018	USP43–NF38	Second Supplement to USP41–NF36	<p><i>Sample solution</i> to:  <math>C_U</math> = nominal concentration of amikacin in the <i>Sample solution</i> (mg/mL)</p> <p>Line 6: Change Scombroidea, to: Scombridae,</p>
INSULIN LISPRO INJECTION	PRODUCT-RELATED SUBSTANCES AND IM PURITIES/ <i>Related Substances</i>	USP41–NF36	2180	31-Aug-2018		1-Sep-2018	USP43–NF38	Second Supplement to USP41–NF36	<p>Line 3 of <i>Analysis</i>: Change Calculate the percentage of insulin lispro, A-21 desamido insulin lispro, and other impurities in the portion of Insulin Lispro taken: to: Calculate the percentage of insulin lispro, A-21 desamido insulin lispro,</p>

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HAZARDOUS DRUGS - HANDLING IN HEALTHCARE SETTINGS	REFERENCES	<i>Revision Bulletin (Official December 01, 2017)</i>	Online	31-Aug-2018		1-Sep-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	and other impurities in the portion of Injection taken: Reference 37: Delete <a href="http://www.osha.gov/Publications/OSHA3151.html">http://www.osha.gov/Publications/OSHA3151.html</a> .
AMINO BENZOATE POTASSIUM	IMPURITIES/Limit of Aniline and p-Toluidine	<i>USP41–NF36</i>	209	31-Aug-2018		1-Sep-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	Line 3 of System suitability: Change [Note—The relative retention times of aniline and p-toluidine are about 4.1 and 5.1 min, respectively.] to: [Note—The relative retention times of aniline and p-toluidine are 0.8 and 1.0, respectively.]
FISH OIL	DEFINITION	<i>USP41–NF36</i>	4622	31-Aug-2018		1-Sep-2018	<i>USP43–NF38</i>	<i>Second</i>	Line 7: Change

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CONTAINING OMEGA-3 ACIDS DELAYED-RELEASE CAPSULES								<i>Supplement to USP41–NF36</i>	Scombroidea, to: Scombridae,
MILRINONE LACTATE INJECTION	IM PURITIES/ <i>Organic Impurities</i>	<i>USP41–NF36</i>	2748	31-Aug-2018		1-Sep-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	Equation in Analysis: Change Result = $(r_U/r_S) \times (C_S/C_U) \times (Mr_1/Mr_2) \times 100$ to: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$ AND In the variable definition list: Delete $Mr_1$ = molecular weight of milrinone free base, 211.22 $Mr_2$ = molecular weight of milrinone lactate, 151.16
HYDROCHLOROTHIAZIDE CAPSULES	PERFORMANCE TESTS/ <i>Dissolution/ Test 2</i>	<i>USP41–NF36</i>	2049	31-Aug-2018		1-Sep-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	Line 1 of <i>Empty capsules solution</i> : Change Place 10

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ASSESSMENT REFERENCES OF EXTRACTABLES ASSOCIATED WITH PHARMACEUTICAL PACKAGING/DELIVERY SYSTEMS		USP41–NF36	7910	27-Jul-2018		1-Aug-2018	USP43–NF38	Second Supplement to USP41–NF36	Capsules into to: Place 10 empty capsules into Line 2 of reference 7: Delete <a href="http://webstore.ansi.org/RecordDetail.aspx?sku=ASTM%20F1980-07(2011)&amp;source=msn&amp;adgroup=astm">http://webstore.ansi.org/RecordDetail.aspx?sku=ASTM%20F1980-07(2011)&amp;source=msn&amp;adgroup=astm</a> . Accessed 19 March 2013.
FISH OIL CONTAINING OMEGA-3 ACIDS	DEFINITION	USP41–NF36	4617	27-Jul-2018		1-Aug-2018	USP43–NF38	Second Supplement to USP41–NF36	Line 5: Change Scombroidea, to: Scombridae,
SODIUM FLUORIDE AND ACIDULATED PHOSPHATE TOPICAL SOLUTION	ASSAY	First Supplement to USP41–NF36	8423	27-Jul-2018		1-Aug-2018	USP43–NF38	Second Supplement to USP41–NF36	Line 1 of Sample solution: Change 1.1 µg/mL of sodium fluoride to: 0.5 µg/mL of fluoride ion AND In the variable

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REAGENTS	CHROMATOG RAPHIC CO LUMN S/Packings	USP41–NF36	5776	27-Jul-2018		1-Aug-2018	USP43–NF38	Second Supplement to USP41–NF36	definition list in <i>Analysis:</i> Change $C_U$ = nominal concentration of sodium fluoride in the <i>Sample solution</i> ( $\mu\text{g/mL}$ ) to: $C_U$ = nominal concentration of fluoride ion in the <i>Sample solution</i> ( $\mu\text{g/mL}$ ) Line 2 of L73: Delete [Note—Available as Jordi-Gel DBV from <a href="http://www.jordiflp.com">www.jordiflp.com</a> .] Delete USP Endotoxin RS
DEXAMETHAS ONE SODIUM PHOSPHATE INJECTION	ADDITIONAL R EQUIREMENT S/USP Reference Standards <11>	USP41–NF36	1204	27-Jul-2018		1-Aug-2018	USP43–NF38	Second Supplement to USP41–NF36	Delete USP Endotoxin RS
RACEPINEPHR INE INHALATION SOLUTION	ADDITIONAL R EQUIREMENT S	USP41–NF36	3564	27-Jul-2018		1-Aug-2018	USP43–NF38	Second Supplement to USP41–NF36	Add USP Reference Standards <11> USP Epinephrine

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MICONAZOLE NITRATE	IM PUR ITIES/ <i>Organic Impurities</i>	<i>First Supplement to USP41–NF36</i>	Online	27-Jul-2018		1-Aug-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	Bitartrate RS (Note: This errata applies to the new <i>USP-NF ONLINE</i> platform only.) Line 1 of <i>Standard solution</i> : Change 1.2 µg/mL each of USP Miconazole Nitrate RS, USP Econazole Nitrate RS, USP Doxazosin Related Compound C RSUSP Doxazosin Related Compound C RS, USP Miconazole Related Compound F RS, and USP Miconazole Related Compound I RS

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							<p>in <i>Diluent</i> to: 1.2 µg/mL each of USP Miconazole Nitrate RS, USP Econazole Nitrate RS, USP Miconazole Related Compound C RS, USP Miconazole Related Compound F RS, and USP Miconazole Related Compound I RS in <i>Diluent</i> AND In the first variable definition list in <i>Analysis:</i> Change <math>C_S =</math> concentration of USP Doxazosin Related Compound C</p>

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							RS, USP Doxazosin Related Compound F RS, USP Miconazole Related Compound I RS, or USP Miconazole Nitrate RS in the <i>Standard            solution</i> (µg/mL) to: $C_S =$ concentration of USP Miconazole Related Compound C RS, USP Miconazole Related Compound F RS, USP Miconazole Related Compound I RS, or USP Econazole Nitrate RS in



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GLUCAGON BIOIDENTITY TESTS <i>B. In Vitro Cell-Based Bioidentity Test</i>	<i>First Supplement to USP41–NF36</i>	8627	27-Jul-2018	1-Aug-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	the <i>Standard solution</i> (µg/mL) Line 1 of <i>Medium B</i> : Change Kreb's salt solution containing 0.3% (v/v) human serum albumen to: Krebs' salt solution containing 0.3% (v/v) human serum albumin AND Line 1 of <i>Medium C</i> : change Kreb's salt solution containing 0.3% (v/v) human serum albumen to: Krebs' salt solution containing 0.3% (v/v) human serum albumin

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FOLIC ACID INJECTION		<i>USP Reference standards &lt;11&gt;</i>	USP41–NF36 1866	27-Jul-2018		1-Aug-2018	USP43–NF38	<i>Second Supplement to USP41–NF36</i>	Delete USP Endotoxin RS
ZIPRASIDONE IM HYDROCHLORIDE	PURITIES/Organic Impurities	<i>Second Supplement to USP41–NF36</i>	8992	27-Jul-2018		1-Aug-2018	USP43–NF38	<i>Second Supplement to USP41–NF36</i>	In the second variable definition in <i>Analysis</i> : Change $r_U$ = peak response of chl oroindolinone or ziprasidone related compound F from the <i>Sample solution</i> to: $r_U$ = peak response of chl oroindolinone, ziprasidone related compound F, or any unspecified impurity from the <i>Sample solution</i>
PLASTIC MATERIALS OF CONSTRUCTION	TEST METHODS		USP41–NF36 6403	27-Jul-2018		1-Aug-2018	USP43–NF38	<i>Second Supplement to USP41–NF36</i>	Line 1 of <i>Plastic Additives/Reference solution</i>

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							<p><i>ns/Reference solution I:</i> Change 0.24 mg/mL of USP Plastic Additive 4 RS prepared in the <i>Solvent mixture</i> to: 0.24 mg/mL of USP Plastic Additive 4 RS prepared in methylene chloride</p> <p>AND</p> <p>Line 1 of <i>Reference solution J:</i> Change 0.24 mg/mL of USP Plastic Additive 5 RS prepared in the <i>Solvent mixture</i> to: 0.24 mg/mL of USP Plastic Additive 5 RS prepared in methylene</p>

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									chloride AND Line 1 of <i>Poly(ethylene-vinyl acetate)/Polyvinyl chloride, plasticized/Reference solutions U, V, W: Change 0.1-mg/mL solutions to: 10.0-mg/mL solutions</i>
DIVALPROEX SODIUM DELA E YED-RELEASE TESTS/ CAPSULES	PERFORMANC	<i>USP41–NF36</i>	1354	27-Jul-2018		1-Aug-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	Line 1 of <i>Tolerances: Change NMT 20% (Q) of the labeled amount of to: NMT 20% of the labeled amount of</i>
	<i>Dissolution &lt;711&gt;/Test 2</i>								
RACEPINEPHRINE HYDROCHLORIDE	<i>USP Reference standards &lt;11&gt;</i>	<i>USP41–NF36</i>	3565	27-Jul-2018		1-Aug-2018	<i>USP43–NF38</i>	<i>Second Supplement to USP41–NF36</i>	Add USP Epinephrine Bitartrate RS (Note: This errata applies to
MICONAZOLE NITRATE	ADDITIONAL R EQUIREMENT	<i>First Supplement to</i>	Online	27-Jul-2018		1-Aug-2018	<i>USP43–NF38</i>	<i>Second Supplement to</i>	

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	S	USP41–NF36						USP41–NF36	the new <i>USP-NF ONLINE</i> platform only.) Line 4 of <i>USP Reference Standards</i> <11>: Change USP Doxazosin Related Compound C RS to: USP Miconazole Related Compound C RS AND Line 7 of <i>USP Reference Standards</i> <11>: Change USP Econazole Nitrate RS to: USP Miconazole Related Compound F RS
REAGENTS	REAGENTS,	USP41–NF36	5737	27-Jul-2018		1-Aug-2018	USP43–NF38	Second	Line 4 of

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	INDICATORS, AND S OL UTION S/Reagent Specifications							Supplement to USP41–NF36	N-(2-Tetrahydro furoyl)piperazin e: Change EMS- DOTTIKON, <a href="http://www.ems-dottikon.ch">www.ems- dottikon.ch</a> . to: Oakwood Chemical, <a href="http://www.oakwoodc&lt;br/&gt;hemical.com">www.oakwoodc hemical.com</a> .
BUPROPION H IDENTIFICATIO YDROCHLORI N/C. DE	Identification Tests—General <191>, Chloride	USP41–NF36	573	27-Jul-2018		1-Aug-2018	USP43–NF38	Second Supplement to USP41–NF36	Line 1 of Acceptance criteria: Change Meets the requirements of test to: Meets the requirements of test A
RACEPINEPHR INE	USP Reference standards <11>	USP41–NF36	3564	27-Jul-2018		1-Aug-2018	USP43–NF38	Second Supplement to USP41–NF36	Add USP Epinephrine Bitartrate RS
MICONAZOLE NITRATE	ASSAY/ Procedure	First Supplement to USP41–NF36	Online	27-Jul-2018		1-Aug-2018	USP43–NF38	Second Supplement to USP41–NF36	(Note: This errata applies to the new USP- NF ONLINE platform only.)

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							Line 1 of <i>System suitability solution</i> : Change 0.1 mg/mL of USP Miconazole Nitrate RS and 6 µg/mL of USP Econazole Nitrate RS in <i>Diluent</i> . to: 0.1 mg/mL of USP Miconazole Nitrate RS and 6 µg/mL of USP Miconazole Related Compound F RS in <i>Diluent</i> .
POLYDEXTRO ASSAY/ SE Procedure	<i>First Supplement to USP41–NF36</i>	8491	29-Jun-2018	1-Jul-2018	<i>USP42–NF37</i>	<i>Second Supplement to USP41–NF36</i>	Line 1 of <i>Sample solution</i> : Change Polydextrose, calculated on the anhydrous and ash-free

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HYDROGENATED POLYDEXTROSE ROSE	USP41–NF36	5495	29-Jun-2018	1-Jul-2018	USP42–NF37	Second Supplement to USP41–NF36	<p>basis, in <i>Mobile phase</i> to: Polydextrose in <i>Mobile phase</i> AND Line 1 of <i>Acceptance criteria</i>: Change NLT 90.0% to: NLT 90.0% on the anhydrous and ash-free basis Change <i>Standard solution</i>: 4.0 mg/mL of USP Polydextrose RS, calculated on the anhydrous and ash-free basis, in <i>Mobile phase Sample solution</i>: 4.0 mg/mL of Hydrogenated Polydextrose, calculated on</p>



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NUCLEAR MAGNETIC RESONANCE SPECTROSCOPY IDENTIFICATION TESTING OF BACTERIAL POLYSACCHARIDES USED IN VACCINE MANUFACTURE	2. PROCEDURE	<i>First Supplement to USP41–NF36</i>	8633	29-Jun-2018		1-Jul-2018	<i>USP42–NF37</i>	<i>Second Supplement to USP41–NF36</i>	the anhydrous and ash-free basis, in <i>Mobile phase</i> to: <i>Standard solution:</i> 4.0 mg/mL of USP Polydextrose RS in <i>Mobile phase Sample solution:</i> 4.0 mg/mL of Hydrogenated Polydextrose in <i>Mobile phase</i> Line 3 of 2.1 <i>Equipment R equipment s/Processing Parameters:</i> Change adsorption to: absorption
CALCIUM CITRATE MALATE	IM PURITIES/ <i>Limit of Fluoride</i>	<i>First Supplement to USP41–NF36</i>	8299	29-Jun-2018		1-Jul-2018	<i>USP42–NF37</i>	<i>Second Supplement to USP41–NF36</i>	Line 1 of <i>Standard solution:</i>

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CHORIONIC GONADOTROPIN FOR INJECTION	USP41–NF36	1982	29-Jun-2018	1-Jul-2018	USP42–NF37	Second Supplement to USP41–NF36	Change Transfer 1.0 mL of <i>Standard stock solution</i> to: Transfer 2.0 mL of <i>Standard stock solution</i> Line 2 of <i>USP Reference Standards</i> <11>: Delete USP Endotoxin RS
ESZOPICLONE ADDITIONAL REQUIREMENT S/USP Reference Standards	Second Supplement to USP41–NF36	Online	25-May-2018	1-Jun-2018	USP42–NF37	Second Supplement to USP41–NF36	Line 2 of USP Eszopiclone Related Compound A RS: [Note—This material may be available in the free base or salt form.] 6-(5-Chloropyridin-2-yl)-7-oxo-6,7-dihydro-5H-pyridolo[3,4-b]pyrazin-5-yl 4

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							<p>-methylpiperazine-1-carboxylate 4-oxide.  <math>C_{17}H_{17}ClN_6O_4</math>  404.81  6-(5-Chloropyridin-2-yl)-7-oxo-6,7-dihydro-5H-pyrido[3,4-b]pyrazin-5-yl 4-methylpiperazine-1-carboxylate 4-oxide, 3-chlorobenzoic salt (1:1).  <math>C_{17}H_{17}ClN_6O_4 \cdot C_7H_5ClO</math> 561.38  to:  [Note—This material may be available in the free base or salt form.]  6-(5-Chloropyridin-2-yl)-7-oxo-6,7-dihydro-5H-pyrido[3,4-</p>

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							-b]pyrazin-5-yl 4 -methylpiperazine-1-carboxylate 4-oxide. $C_{17}H_{17}ClN_6O_4$ 404.81 6-(5-Chloropyridin-2-yl)-7-oxo-6,7-dihydro-5H-pyridolo[3,4-b]pyrazin-5-yl 4-methylpiperazine-1-carboxylate 4-oxide, 3-chlorobenzoate salt (1:1). $C_{17}H_{17}ClN_6O_4 \cdot C_7H_5ClO_2$ 561.38

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