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- **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.
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AZITHROMYCIN FOR INJECTION	ADDITIONAL REQUIREMENTS	USPNF Online	Online	28-Jul-2023	1-Aug-2023	NA	USPNF 2024 Issue 2	In USP Reference Standards ?11?/USP Azithromycin A RS: Change 734.96 to: 734.97 AND In USP Azithromycin N-oxide RS: Change 764.98 to: 765.00 AND In USP N-Demethylazithromycin RS: Change 734.96 to: 734.97 AND In USP Desominylazithromycin RS: Change 590.79

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AZITHROMYCI IMPURITIES N FOR INJECTION	USPNF Online Online		28-Jul-2023	1-Aug-2023	NA	USPNF 2024 Issue 2	to: 590.80 In footnote m in Table 2: Change (2R,3S,4R,5R ,8R,10R,11R ,12S,13S,14R)-13-[(2,6-Dideo xy-3-C -meth yl-3-O-methyl-?- L-ribo -hexopyranosyl) oxy]-2-ethyl-3,4, 10-trihydroxy-3, 5,6,8,10,12,14- heptamethyl-11- [[3-[N -(4-acetamidop h en ylsulf onyl)-N -methylamino]-3 ,4,6-trideoxy-?- D-xylo -hexopyranosyl] oxy]-1-oxa-6-az acyclopentadec

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						—	an-15-one. to: (2 <i>R</i> ,3 <i>S</i> ,4 <i>R</i> ,5 <i>R</i> ,8 <i>R</i> ,10 <i>R</i> ,11 <i>R</i> ,12 <i>S</i> ,13 <i>S</i> ,14 <i>R</i>)-13-[(2,6-Dideoxy-3- <i>C</i> -meth yl-3- <i>O</i> -methyl-?- <i>L</i> -ribo -hexopyranosyl)oxy]-2-ethyl-3,4,10-trihydroxy-3,5,6,8,10,12,14-heptamethyl-11-[[3- <i>N</i> -(4-methylphenylsulfon yl)- <i>N</i> -methylamino]-3,4,6-trideoxy-β- <i>D</i> -xylo -hexopyranosyl]oxy]-1-oxa-6-azacyclopentadecan-15-one.
ROCURONIUM CHEMICAL BROMIDE	USPNF Online Online INFORMATION		18-Nov-2022	1-Dec-2022	NA	USPNF 2023 Issue 3	Change 609.68 to: 609.69

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ROCURONIUM IM BROMIDE	IMPURITIES/ <i>Organic Impurities</i>	USPNF Online	Online	18-Nov-2022	1-Dec-2022	NA	USPNF 2023 Issue 3	In footnote b of Table 1: Change 2?-(Morpholin-4-yl)-16?-(pyrrolidin-1-yl)-5?-and rostan-3?,17?-diol. to: 2?-(Morpholin-4-yl)-16?-(pyrrolidin-1-yl)-5?-and rostane-3?,17?-diol.
METHYLENE BLUE	IMPURITIES/ <i>Organic Impurities</i>	USPNF Online	Online	26-Aug-2022	1-Sep-2022	NA	USPNF 2023 Issue 2	In footnote a in Table 2: Change 3-(Dimethylamino)-7-(methyldamino)-phenothiazine-5-ium chloride. to: 3-(Dimethylamino)-7-(methyldamino)phenothiazin-5-ium chloride.
METHYLENE BLUE	ADDITIONAL REQUIREMENT	USPNF Online	Online	26-Aug-2022	1-Sep-2022	NA	USPNF 2023 Issue 2	In USP Azure B RS: Change

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							—	3-(Dimethylamino)-7-(methy- l- amino)-phenothiazine-5-ium chloride. to: 3-(Dimethylamino)-7-(methy- l- amino)phenothiazine-5-ium chloride.
ECONAZOLE NITRATE	ADDITIONAL REQUIREMENT <i>S/USP Reference Standards <11></i>	USP-NF Online	Online	26-Aug-2022	1-Sep-2022	NA	USP-NF 2023 Issue 2	In USP Econazole Related Compound C RS: Change 1-(4-Chlorobenzyl)-3-{2-[(4-chlorobenzyl)oxy]-2-(2,4-dichlorophenyl)-1H-imidazol-3-ium nitrate (salt). C ₂₅ H ₂₁ Cl ₁₄ N ₃ O ₄ 569.26 to: 1-(4-Chlorobenzyl)-3-{2-[(4-chlorobenzyl)oxy]-2-(

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							—	2,4-dichlorophenyl)-1H-imidazol-3-ium chloride. C ₂₅ H ₂₁ Cl ₅ N ₂ O 542.71
NAPROXEN SODIUM TABLETS	IM PURITIES/Organic Impurities	<i>First Supplement to USP41–NF36</i>	8363	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	In the System suitability solution: Change 0.5 mg/mL of USP Naproxen Sodium 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: 0.5 µg/mL of USP Naproxen Related Compound A RS from

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							—	Standard stock solution 2 and 0.5 mg/mL of USP Naproxen Sodium RS in Diluent
CUPRIC SULFATE INJECTION	Assay	USP41–NF36	1112	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	Line 8 of Assay preparation: Change to bring the total sodium content of this flask to 13.5 mg.
								to: to bring the total sodium chloride content of this flask to 13.5 mg.
OCTOCRYLEN E	SPECIFIC TESTS/Acidity	First Supplement to USP41–NF36	8379	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	Line 1 of Acceptance criteria: Change NMT 0.18 mL of Titrant is required to: NMT 0.18 mL of Titrant/g is required
FLUTICASONE IM		USP41–NF36	1836	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	In the Tailing

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PROPIONATE INHALATION POWDER	PURITIES/Organic Impurities/System suitability						—	factor: Change NLT 1.3 to: NMT 1.3
OXYBUTYNIN CHLORIDE EXTENDED-RELEASE TABLETS	PERFORMANCE TESTS/Dissolution <711>/Test 8	Revision Bulletin (Official April 01, 2018)	Online	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	In the variable definition list in Analysis: Change C_s = concentration of oxybutynin chloride in the Standard solution (mg/mL) to: C_s = concentration of USP Oxybutynin Chloride RS in the Standard solution (mg/mL)
GLUTARAL CO NCENTRATE	ASSAY/Procedure	USP41–NF36	1960	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	In the variable definition list in Analysis: Change W = weight of

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							—	Concentrate taken (g) to: W = nominal weight of glutaral taken (g)
LIDOCAINE	ASSAY/ Procedure	<i>First Supplement to USP41–NF36</i>	Online	30-Nov-2018	1-Dec-2018	<i>USP43–NF38</i>	<i>USP42–NF37</i>	This erratum applies to the new <i>USP-NF</i> ONLINE platform only. Line 2 of <i>Standard solution</i> : Change 1 N sodium hydroxide, to: 1 N hydrochloric acid, AND Line 2 of <i>Sample solution</i> : Change 1 N sodium hydroxide, to: 1 N hydrochloric

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AZEOTROPIC ISOPROPYL ALCOHOL	ADDITIONAL REQUIREMENT <i>S/USP Reference Standards <11></i>	USP41–NF36	2257	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	acid, Line 1 of USP 2-Propanol System Suitability RS: Change It contains 0.1% of each of the following: ethyl ether, acetone, isopropyl alcohol, diisopropyl ether, 1-propanol, and 2-butanol. to: It is a mixture of the following: ethyl ether (0.1%), acetone (0.1%), diisopropyl ether (0.1%), 1-propanol (0.1%), 2-butanol (0.1%), and isopropyl alcohol (99.5%).

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OXYBUTYNIN ASSAY/ CHLORIDE EX Procedure TENDED- RELEASE TABLETS	<i>Revision Bulletin (Official April 01, 2018)</i>	Online	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	This erratum applies to the Revision Bulletin posted on www.uspnf.com only. In System suitability: Add [Note—The relative retention times for oxybutynin and oxybutynin related compound A are about 1.0 and 1.6, respectively.]
METOPROLOL CHEMICAL TARTRATE INFORMATION	USP41–NF36	2712	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	Line 4: Change (±)-1-(Isopropyl amin o)-3-[p -(2-methoxyethy l)phenoxy]-2-pr opanol l-(+)-tartrate (2:1) (salt); 1-(Isopropylami no)-3-[p

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
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							—	-(2-methoxyethyl)phenoxy]-2-propanol (2:1) dextro-tartrate salt to: (±)-1-(Isopropylamino)-3-[p-(2-methoxyethyl)phenoxy]-2-propanol (+)-tartrate (2:1) (salt); 1-(Isopropylamino)-3-[4-(2-methoxyethyl)phenoxy]propan-2-ol L-tartrate
OXYBUTYNIN CHLORIDE EX-TENDED-RELEASE TABLETS	IMPURITIES/Organic Impurities	Revision Bulletin (Official April 01, 2018)	Online	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	This erratum applies to the Revision Bulletin posted on www.uspnf.com only. In the variable definition list in <i>Analysis</i> : Change

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						—	C_U = nominal concentration of the <i>Sample solution</i> (mg/mL) to: C_U = nominal concentration of the <i>Sample solution</i> (mg/mL) [Note—Disregard any peak less than 0.1%.] Line 2 of USP Salmeterol Related Compound H RS: Change 1-Hydroxy-4-[2-hydroxy-5-(1-hydroxy-2-[[6-(4-phenylbutoxy)hexyl]amino}ethyl)benzyl]-2-naphthoic acid. $C_{36}H_{43}NO_6$ 585.73 to: 1-Hydroxy-4-[2-
FLUTICASONE ADDITIONAL R PROPIONATE EQUIREMENT AND <i>S/USP</i> SALMETEROL <i>Reference</i> INHALATION <i>Standards <11></i> POWDER	USP40–NF35	4309	30-Mar-2018	1-Apr-2018	USP42–NF37	USP42–NF37	

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							—	hydroxy-5-(1-hydroxy-2-[[6-(4-phenylbutoxy)hexyl]amino]ethyl)benzyl]-2-naphthoic acid, monohydrate. C ₃₆ H ₄₃ NO ₆ · H ₂ O 603.76
PANTOPRAZOLE SODIUM DEE LAYED-RELEASE TABLETS	PERFORMANCE TESTS/ <i>Dissolution Test</i> <i>1</i>	USP41–NF36	3157	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	In the <i>Analysis</i> : Change Result = (r _U /r _S) × C _S × (M _{r1} /M _{r2}) × V × (100/L) to: Result = (r _U /r _S) × C _S × (M _{r1} /M _{r2}) × V × (100/L) × D AND In the variable definition list in <i>Analysis</i> : Add D = dilution factor for the <i>Sample solution</i> , 2
PERINDOPRIL ERBUMINE	IMPURITIES/ <i>Limit of Perindopril</i>	USP40–NF35	5644	30-Mar-2018	1-Apr-2018	USP42–NF37	USP42–NF37	Line 3: Add [Note—The relative

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							—	retention times for perindopril and perindopril related compound I are 1.0 and 1.6, respectively.]
AMLODIPINE AND ATORVAS TATIN TABLETS	DEFINITION	<i>First Supplement to USP41–NF36</i>	8270	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	Line 4: Change (C ₃₃ H ₃₄ FN ₂ O ₅) to: (C ₃₃ H ₃₅ FN ₂ O ₅)
SALMETEROL INHALATION POWDER	ADDITIONAL REQUIREMENT S/USP	USP40–NF35	6096	30-Mar-2018	1-Apr-2018	USP42–NF37	USP42–NF37	Line 2 of USP Salmeterol Related Compound H RS: Change 1-Hydroxy-4-[2-hydroxy-5-(1-hydroxy-2-[[6-(4-p henylbutoxy)he xyl]amino}ethyl) benzyl]-2-napht hoic acid. C ₃₆ H ₄₃ NO ₆ 585.73 to: 1-Hydroxy-4-[2-hydroxy-5-(1-hydroxy-2-[[6-(4-p henylbutoxy)he

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							Sort ascending	
							—	xyl]amino}ethyl) benzyl]-2-napht hoic acid, monohydrate. $C_{36}H_{43}NO_6 \cdot H_2O$ 603.76
ALLOPURINOL	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USP41–NF36	121	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	Line 3 of USP Allopurinol Related Compound A RS: Change $(C_5H_6N_4O)_2 \cdot H_2SO_4$ 350.32 to: $(C_4H_6N_4O)_2 \cdot H_2SO_4$ 350.31
AMLODIPINE AND ATORVASTATIN TABLETS	ASSAY/Procedure	First Supplement to USP41–NF36	8270	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	In the second Calculate statement in Analysis: Change $(C_{33}H_{34}FN_2O_5)$ to: $(C_{33}H_{35}FN_2O_5)$
PLASTIC MATERIALS OF CONSTRUCTION	TEST METHODS/Tin in Non-Tin-Stabilized Materials	USP41–NF36	6403	30-Mar-2018	1-Apr-2018	USP42–NF37	USP42–NF37	Line 3 of Sample solution: Change If the solution is not colorless,

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						—	add the sodium sulfate to: If the solution is not colorless, add the sodium sulfite
CIPROFLOXAC IMPURITIES IN AND DEXAMETHASONE OTIC SUSPENSION	USP41–NF36	951	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	In the variable definition list of <i>Ciprofloxacin Related Compound Analysis</i> : Change C_U = nominal concentration of ciprofloxacin in the Otic Suspension (mg/mL) to: C_U = nominal concentration of ciprofloxacin in the <i>Sample solution</i> (mg/mL) AND In the variable

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							—	definition list of <i>Dexamethasone Related Compound Analysis: Change C_U = nominal concentration of dexamethasone in the Otic Suspension (mg/mL) to: C_U = nominal concentration of dexamethasone in the Sample solution (mg/mL)</i>
AMLODIPINE AND ATORVASE TATIN TABLETS	PERFORMANCE TESTS/ <i>Dissolution</i> <711>	<i>First Supplement to USP41–NF36</i>	8270	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	In the second Calculate statement in <i>Analysis: Change (C₃₃H₃₄FN₂O₅) to: (C₃₃H₃₅FN₂O₅) AND In Tolerances:</i>

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							—	Change (C ₃₃ H ₃₄ FN ₂ O ₅) to: (C ₃₃ H ₃₅ FN ₂ O ₅) Line 1 of <i>Analysis:</i> Change To the <i>Sample solution</i> to: To 50 mL of the <i>Sample solution</i>
CUPRIC CHLORIDE	ASSAY/ <i>Procedure</i>	USP41–NF36	1109	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	
ACAMPROSAT IM E CALCIUM	<i>First</i> PURITIES/ <i>Limit Supplement to of Acamprosate USP41–NF36 Related Compound A</i>		8263	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	In the variable definition list in <i>Analysis:</i> Change C _S = concentration of USP Acamprosate Calcium Related Compound A RS in the <i>Standard solution</i> (µg/mL) to: C _S = concentration of USP

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							—	Acamprosate Related Compound A RS in the <i>Standard solution</i> (µg/mL)
CUPRIC CHLORIDE INJECTION	ASSAY/ <i>Procedure</i>	USP41–NF36	1111	30-Nov-2018	1-Dec-2018	USP43–NF38	USP42–NF37	Line 3 of <i>Sample solution</i> : Change to bring the total sodium content of this flask to 13.5 mg. to: to bring the total sodium chloride content of this flask to 13.5 mg.
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

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							—	<i>M formic acid solution.</i> to: Add 1.4 M ammonia solution to 1.4 M formic acid solution until a pH of 4.4 is obtained.
SCAFFOLD BOVINE DERMIS	SPECIFIC TESTS	USP40–NF35	6113	26-May-2017	1-Jun-2017	USP41–NF36	USP41–NF36	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 Retention Force/Analysis</i>: Change 40 to: 4-0
MYCOPHENOLATE MOFETIL FOR ORAL	ADDITIONAL REQUIREMENT S/USP	USP39–NF34	4961	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	Line 5 of <i>Mycophenolate Mofetil Related</i>

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SUSPENSION	Reference Standards <11>						—	Compound A RS: Change $C_{23}H_{31}NO_7$ to: $C_{22}H_{29}NO_7$
ROPINIROLE XTENDED- RELEASE TABLETS	E IM PUR ITIES/Organic Impurities/Analysis	Second Supplement to USP39–NF34	8814	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	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 Sample solution (µg/mL) to: C_U = nominal concentration of ropinirole in the Sample solution (µg/mL)
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 Delivered Dose

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						—	<i>Uniformity in Metered-Dose Inhalers and Dry Powder Inhalers, as described in <601>, to:</i> <i>proceed as directed in Inhalation and Nasal Drug Products: Aerosols, Sprays, and Powders—Performance Quality Tests <601>, A. Delivered-Dose Uniformity/A.2 Inhalation Aerosols and Inhalation Sprays and A.4 Inhalation Powders, AND</i> <i>Line 4: Delete Unless otherwise</i>

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							—	stated in the individual monograph, apply the acceptance criteria for <i>Metered-Dose Inhalers and Dry Powder Inhalers</i> as described in <601>.
EVENING PRIMROSE OIL REN CAPSULES	ST GTH/ <i>Analysis</i>	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 <i>Standard solution</i> (mg) to: m_S = weight of the relevant USP Methyl Ester RS in the <i>Standard solution</i> (mg)
SORBITOL SOLUTION	ASSAY/ <i>Proce</i>	USP39–NF34	5897	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	In the variable definition list:

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							—	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)
RIBOFLAVIN ASSAY	ASSAY	USP40–NF35	386	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	Line 2 of the variable definition list in <i>Chemical Methods, Procedure 1/Analysis</i> : Change sodium hydrofluorite to: sodium hydrosulfite
NUCLEIC ACID-BASED TECHNIQUES—G		USP40–NF35	1471	26-May-2017	1-Jun-2017	USP41–NF36	USP41–NF36	Line 1 of bullet 1: Change "Review Criteria

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ENERAL	Appendix 1: Regulations and Standards						—	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)
VINORELBINE TARTRATE	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USP39–NF34	6370	31-Mar-2017	1-Apr-2017	USP41–NF36	USP41–NF36	Line 2 of USP Vinorelbine Related Compound A RS: Change 4-O -Deacetylvinorelbine. to: 4-O -Deacetylvinorelbine tartrate.

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TIMOLOL MALEATE	MULTIPLE SECTIONS	<i>Second Supplement to USP40–NF35</i>	Online	28-Jul-2017	1-Aug-2017	USP41–NF36	USP41–NF36	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 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.uspn

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							—	f.com/notices/se cond-suppleme nt-usp-40-nf-35- online-timolol- maleate.
SUCROSE	SPECIFIC TESTS/ <i>Color Value/Analysis</i>	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. to: <i>Suitability requirements</i> <i>Repeatability:</i> The absolute difference between two results is NMT 3.
SPACERS AND VALVED HOLDING CHAMBERS USED WITH INHALATION AEROSOLS—CH	1. INTRODUCTION/1.5 <i>Definitions of Key Terms</i> <i>Relating to This Chapter</i>	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

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CHARACTERIZATION TESTS							—	mouthpiece
ACETYLCYSTEINE SOLUTION	ASSAY/ <i>Procedure</i>	USP40–NF35	2586	26-May-2017	1-Jun-2017	USP41–NF36	USP41–NF36	Line 3 of <i>Sample solution</i> : Change <i>Standard stock solution</i> to: <i>Sample stock solution</i>
VINORELBINE INJECTION	ADDITIONAL REQUIREMENTS/ <i>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 -Deacetylvinorelbine. to: 4-O -Deacetylvinorelbine tartrate.
METHIONINE	IMPURITIES/Related Compounds	<i>First Supplement to</i> USP40–NF35	8337	26-May-2017	1-Jun-2017	USP41–NF36	USP41–NF36	Row 5 of Column 3 of <i>Table 2</i> : Change 0.1 to: 1.0

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
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PENICILLIN G PROCAINE	SPECIFIC TESTS/ <i>Content of Penicillin G and Procaine/Analysis</i>	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 <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

Monograph Title	Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							Sort ascending	
							—	Sample solution (mg) to: C_U = concentration of Penicillin G Procaine in the Sample solution (mg/mL)
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
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 <i>Table 1</i> : Change 0

Monograph Title Section	Source Publication	Page Number	Errata Post Date	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							_ Sort ascending
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