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How to Use

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Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
MINOCYCLINE IM		*First*	8101	31-Mar-2017	1-Apr-2017	*USP41–NF36*	*USP41–NF36*	Change

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
HYDROCHLORIDE EXTENDED-RELEASE TABLETS	PURITIES/*Organic Impurities*	*Supplement to USP39–NF34*						*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 solution*: Use the *Sample stock solution* as directed in the *Assay*.
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 *Figure 1*:Change VHC mouthpiece to: Spacer mouthpiece
GRANISETRO	*USP Reference*	*First*	Online	31-Mar-2017	1-Apr-2017	*USP41–NF36*	*USP41–NF36*	Line 3 of USP

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N HYDROCHLORIDE TABLETS	*standards <11>*	*Supplement to USP40–NF35*						Granisetron Related Compound C RS: Change carboxamide. to: carboxamide hydrochloride.
DOBUTAMINE INJECTION	IDENTIFICATION/*A.*	*USP39–NF34*	3561	31-Mar-2017	1-Apr-2017	*USP41–NF36*	*USP41–NF36*	Line 1 of *Sample solution*: Change 10 mg/mL of dobutamine hydrochloride in methanol, clarified by centrifugation to: Use the neat Injection.
SORBITOL SOLUTION	ASSAY/*Procedure/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 *Sample solution* (mg/g) to:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								C_U = concentration of Sorbitol Solution in the *Sample solution* (mg/g)
MILK THISTLE TABLETS	STRENGTH/*Content of Silymarin*	*USP39–NF34*	6767	31-Mar-2017	1-Apr-2017	*USP41–NF36*	*USP41–NF36*	Row 2 of Column 2 of *Table 1*: Change 0 to: 85 AND Row 2 of Column 3 of *Table 1*: Change 0 to: 15
ADAPALENE GEL	ASSAY/*Procedure*	*First Supplement to USP39–NF34*	7983	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 5 of *Sample stock solution*: Change Cool to room temperature and dilute with *Diluent* to volume. to:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Cool to room temperature and dilute with *Mobile phase* to volume.
PROMETHAZINE HYDROCHLORIDE TABLETS	ADDITIONAL REQUIREMENTS/*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. $C_{17}H_{20}N_2S$ 284.42 to: Isopromethazine hydrochloride; *N*,*N* -

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								H-phenothiazin-10-yl)propan-1-amine hydrochloride. $C_{17}H_{20}N_2S \cdot HCl$ 320.88
AMINOPHYLLINE	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP40–NF35*	2735	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2 of USP Theophylline Related Compound D RS: Change *N*-Methyl-5-(methylamino)-1*H*-imidazole-4-carboxamide. $C_6H_{10}N_4O$ 154.17 to: Theophyllidine; *N*-Methyl-5-(methylamino)-1*H*-imidazole-4-carboxamide hydrochloride monohydrate. C

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								$_6H_{10}N_4O \cdot HCl \cdot H_2O$ 208.65
ATROPINE SULFATE	ASSAY/ *Procedure*	*USP39–NF34*	2638	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 4 of *Analysis*: Change $(C_{17}H_{23}NO_{32} \cdot H_2SO_4)$ to: $[(C_{17}H_{23}NO_3)_2 \cdot H_2SO_4]$
LEVETIRACETAM EXTENDED-RELEASE TABLETS	PERFORMANCE TESTS/ *Dissolution <711>/Test 7*	*Revision Bulletin (Official October 01, 2016)*	Online	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2 of *Standard solution*: Change *Buffer A*. to: *Medium*.
EFAVIRENZ	IMPURITIES/*Organic Impurities*	*USP39–NF34*	3656	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Row 4 of Column 1 of *Procedure 1/Impurity Table 1*:Change Efavirenz pent-3-ene-1-yne (*cis*)[c] to: Efavirenz pent-3-ene-1-yne (*trans*)[c] AND Row 5 of

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Column 1 of *Procedure 1*/*Impurity Table 1*:Change Efavirenz pent-3-ene-1-yne (*trans*)[d] to: Efavirenz pent-3-ene-1-yne (*cis*)[d] AND Row 3 of Column 1 of *Procedure 2*/*Impurity Table 2*: Change Efavirenz pent-3-ene-1-yne (*cis*)[a] to: Efavirenz pent-3-ene-1-yne (*trans*)[a] AND Row 4 of Column 1 of *Procedure 2*/*Impurity Table 2*:Change

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								Efavirenz pent-3-ene-1-yne (*trans*)[b] to: Efavirenz pent-3-ene-1-yne (*cis*)[b]
TETRACYCLINE HYDROCHLORIDE	IMPURITIES/*Organic Impurities*	*USP39–NF34*	6080	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 16 of *Analysis*: Delete Calculate the percentage of any unspecified impurity in the portion of Tetracycline Hydrochloride taken: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$ r_U = peak response of any unspecified impurity from the *Sample solution* r_S = peak response of tetracycline from the *Standard solution*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								C_S = concentration of USP Tetracycline Hydrochloride RS in the *Standard solution* (µg/mL) C_U = concentration of Tetracycline Hydrochloride in the *Sample solution* (µg/mL)
POWDERED TURMERIC	COMPOSITION */Content of C urc uminoi ds/ Chromatographi c system*	*USP39–NF34*	6867	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Column*: Change 4.6-mm × 20-cm; to: 4.6-mm × 25-cm;
PROMETHAZINE HYDROCHLORIDE INJECTION	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*Second Supplement to USP39–NF34*	8785	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* -

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								H-phenothiazin-10-yl)propan-1-amine. $C_{17}H_{20}N_2S$ 284.42 to: Isopromethazine hydrochloride; *N*,*N*-
THEOPHYLLINE	ADDITIONAL REQUIREMENTS/*USP Reference Standards RS <11>*	*Second Supplement to USP39–NF34*	8844	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	*H*-phenothiazin-10-yl)propan-1-amine hydrochloride. $C_{17}H_{20}N_2S \cdot HCl$ 320.88 Line 3 of USP Theophylline Related Compound D RS: Change *N*-Methyl-5-(meth

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								ylami no)-1*H* -imidazole-4-car boxamide. $C_6H_{10}N_4O$ 154.17 to: *N* -Methyl-5-(meth ylami no)-1*H* -imidazole-4-car boxamide hydrochloride monohydrate. $C_6H_{10}N_4O \cdot HCl \cdot H_2O$ 208.65
AMINOPHYLLINE TABLETS	PERFORMANCE TESTS/ *Uniformity of Dosage Units <905>*/ *Procedure for content uniformity*	*USP39–NF34*	2483	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Variable definition for C_U in *Analysis*: Change (mg/mL) to: to:(µg/mL)
AMINOPHYLLINE TABLETS	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP40–NF35*	2742	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2 of USP Theophylline Related Compound D RS: Change

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								N-Methyl-5-(methylamino)-1*H*-imidazole-4-carboxamide. $C_6H_{10}N_4O$ 154.17 to: Theophyllidine; *N*-Methyl-5-(methylamino)-1*H*-imidazole-4-carboxamide hydrochloride monohydrate. $C_6H_{10}N_4O \cdot HCl \cdot H_2O$ 208.65
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE CREAM	ASSAY/ *Procedure*	*USP39–NF34*	3262	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	In the first variable definition list in *Analysis*: Change C_S = concentration of USP Clotrimazole RS in the *Clotrimazole*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								stock solution (mg/mL) to: C_S = concentration of USP Clotrimazole RS in the *Standard solution* (mg/mL) AND In the second variable definition list in *Analysis*: Change C_S = concentration of USP Betamethasone Dipropionate RS in the *Betamethasone dipropionate stock solution* (mg/mL) to: C_S = concentration of USP Betamethasone

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Dipropionate RS in the *Standard solution* (mg/mL)
TERAZOSIN TABLETS	ASSAY/ *Procedure*	*USP39–NF34*	6045	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 3 of *System suitability solution*: Change *Standard solution* to: *Standard stock solution*
CURCUMINOIDS TABLETS	STRENGTH/*Content of Curcuminoids/ Chromatographic system*	*USP39–NF34*	6585	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Column*: Change 4.6-mm × 20-cm; to: 4.6-mm × 25-cm;
IMIPRAMINE PAMOATE CAPSULES	PERFORMANCE TESTS/ *Dissolution <711>/Test 1*	*Second Supplement to USP39–NF34*	8681	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 6 of *Analysis*: Change $Result_i = (r_i/r_s) \times C_s \times [M \times (M$

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								$_{r1}/M_{r2})] \times V$ x (1/*L*) × 100 to: Result$_i$ = (r_i/r_s) × C_s × [*M* × (M_{r1}/M_{r2})] × *D* × *V* × (1/*L*) × 100 AND To the variable definition list in *Analysis*: Add *D* = dilution factor of the *Sample solution*
SODIUM CHLORIDE	IM PURITIES/*Limit of Potassium/ Instrumental conditions*	*Second Supplement to USP39–NF34*	8821	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Mode*: Change Atomic absorption spectrophotometry to: Atomic emission spectroscopy
AMINOPHYLLINE INJECTION	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP40–NF35*	2737	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2 of USP Theophylline Related Compound D RS: Change *N* -Methyl-5-(methylami

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								no)-1*H*-imidazole-4-carboxamide. $C_6H_{10}N_4O$ 154.17 to: Theophyllidine; *N*-Methyl-5-(methylamino)-1*H*-imidazole-4-carboxamide hydrochloride monohydrate. $C_6H_{10}N_4O \cdot HCl \cdot H_2O$ 208.65
BRINZOLAMIDE	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP39–NF34*	2788	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2 of USP Brinzolamide Related Compound B RS: Change (*R*-4-Amino)-2,3-dihydro-2-(3-methoxy propyl)-4*H*-thieno[3,2,-e]-thiazine-6-sulfonamide-1,1-dioxi

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								de ethandioate 1:1. to: (*R*)-4-Amino-2-(3-methoxypropyl)-3, 4-di hydro-2*H* -thieno[3,2-e][1, 2]thiazine-6-sulf onamide 1,1-dioxide oxalate.
ANALYTICAL INSTRUMENT QUALIFICATION	ANALYTICAL INSTRUMENT QUALIFICATION PR OCES S/*Qualification Phases*	*First Supplement to USP40–NF35*	8083	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2 of paragraph 1 of *Operational Qualification*: Change OQ demonstrates fitness for the selected use, and should reflect the contents of the DQ document. to: OQ demonstrates fitness for the

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								selected use, and should reflect URS. AND Line 2 of paragraph 3 of *Operational Qualification/Instrument Function Tests*: Change Holistic tests, which involve the entire system, demonstrate that the whole system complies with user specifications in the DQ. to: Holistic tests, which involve the entire system, demonstrate that the whole system

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								complies with URS.
ENTECAVIR	ASSAY/ *Procedure*	*USP39–NF34*	3704	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 5 of *Analysis*: Change Result = $(r_U/r_S) \times (C_S/C_U) \times (M_{r1}/M_{r2}) \times 100$ to: Result = $(r_U/r_S) \times (C_S/C_U) \times 100$ AND Line 14 of *Analysis*: Delete M_{r1}=molecular weight of anhydrous entecavir, 277.28 M_{r2}=molecular weight of entecavir, 295.29
CURCUMINOIDS	COMPOSITION */Content of C urc uminoi ds/ Chromatographi c system*	*USP39–NF34*	6582	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Column*: Change 4.6-mm × 20-cm; to: 4.6-mm × 25-cm;

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POWDERED TURMERIC EXTRACT	COMPOSITION /*Content of C urc uminoi ds/ Chromatographi c system*	*USP39–NF34*	6868	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Column*: Change 4.6-mm × 20-cm; to: 4.6-mm × 25-cm;
PROMETHAZINE HYDROCHLORIDE ORAL SOLUTION	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*Second Supplement to USP39–NF34*	8787	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. $C_{17}H_{20}N_2S$ 282.42 to: Isopromethazine hydrochloride; *N*,*N* -

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								H-phenothiazin-10-yl)propan-1-amine hydrochloride. $C_{17}H_{20}N_2S \cdot HCl$ 320.88
THEOPHYLLINE ORAL SOLUTION	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*Second Supplement to USP39–NF34*	8846	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 3 of USP Theophylline Related Compound D RS: Change *N*-Methyl-5-(methylamino)-1*H*-imidazole-4-carboxamide. $C_6H_{10}N_4O$ 154.17 to: *N*-Methyl-5-(methylamino)-1*H*-imidazole-4-carboxamide hydrochloride

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								monohydrate. $C_6H_{10}N_4O \cdot HCl \cdot H_2O$ 208.65
ATROPINE SULFATE	DEFINITION	*USP39–NF34*	2638	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2: Change ($C_{17}H_{23}NO_{32} \cdot H_2SO_4$), to: [$(C_{17}H_{23}NO_3)_2 \cdot H_2SO_4$],
GALANTAMINE EXTENDED-RELEASE CAPSULES	PERFORMANCE TESTS/ *Dissolution <711>/Test 3*	*USP40–NF35*	4367	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1of *Buffer*: Change To each L of 6.8-g/L potassium phosphate to: To each L of 6.8 g/L of monobasic potassium phosphate
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE CREAM	IMPURITIES/*Organic Impurities: Limit of Clotrimazole Related Compound A*	*USP39–NF34*	3262	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	In the variable definition list in *Analysis*: Change C_S = concentration of USP Clotrimazole Related Compound A

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								RS in the *Clotrimazole related compound A stock solution* (mg/mL) to: C_S = concentration of USP Clotrimazole Related Compound A RS in the *Standard solution* (mg/mL)
TERAZOSIN TABLETS	IMPURITIES/*Organic Impurities*/*Procedure*	*USP39–NF34*	6045	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Sample solution*: Change Transfer 15 mg of the powder from the crushed Tablets to: Transfer a suitable amount of powder, equivalent to 15 mg of terazosin

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								hydrochloride, from the crushed Tablets
TURMERIC	COMPOSITION */Content of C urc uminoi ds/ Chromatographi c system*	*USP39–NF34*	6866	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Column*: Change 4.6-mm × 20-cm; to: 4.6-mm × 25-cm;
PROMETHAZINE HYDROCHLORIDE	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*Second Supplement to USP39–NF34*	8784	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. $C_{17}H_{20}N_2S$ 282.42 to: Isopromethazin

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								e hydrochloride; *N*,*N* - *H* -phenothiazin-10-yl)propan-1-amine hydrochloride. $C_{17}H_{20}N_2S \cdot HCl$ 320.88
TACROLIMUS CAPSULES	PERFORMANCE TESTS/ *Dissolution <711>/Test 5*	*Second Supplement to USP39–NF34*	8834	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Medium*: Change 0.5 g/L to: 0.05 g/L
ADENINE	CHEMICAL INFORMATION	*USP39–NF34*	2346	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2: Change 1*H* -Purin-6-amine; to: 9*H* -Purin-6-amine;
AMINOPHYLLINE ORAL SOLUTION	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP40–NF35*	2739	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2 of USP Theophylline Related Compound D RS: Change *N*

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								-Methyl-5-(methylamino)-1*H*-imidazole-4-carboxamide. $C_6H_{10}N_4O$ 154.17 to: Theophyllidine; *N*-Methyl-5-(methylamino)-1*H*-imidazole-4-carboxamide hydrochloride monohydrate. $C_6H_{10}N_4O \cdot HCl \cdot H_2O$ 208.65
BRINZOLAMIDE OPHTHALMIC SUSPENSION	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*USP39–NF34*	2789	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2 of USP Brinzolamide Related Compound B RS: Change (*R*-4-Amino)-2,3-dihydro-2-(3-methoxypropyl)-4*H*-thieno[3,2,-e]-t

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								hiazine-6-sulfonamide-1,1-dioxide ethandioate 1:1. to: (*R*)-4-Amino-2-(3-methoxypropyl)-3, 4-dihydro-2*H*-thieno[3,2-e][1, 2]thiazine-6-sulfonamide 1,1-dioxide oxalate.
HALCINONIDE	IMPURITIES/*Organic Impurities/ Chromatographic system*	*USP39–NF34*	4175	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Column*: Change 1.8-µm packing L1 to: 1.7-µm packing L1
CURCUMINOIDS CAPSULES	STRENGTH/*Content of Curcuminoids/*	*USP39–NF34*	6583	27-Jan-2017	1-Feb-2017	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Column*: Change 4.6-mm × 20-cm; to: 4.6-mm ×

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
	Chromatographic system							25-cm;
GUAR GUM	ASSAY/*Content of Galactomannan and Ratio of Constituting Mannose and Galactose*	*First Supplement to USP39–NF34*	7964	18-Nov-2016	1-Dec-2016	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 4 of *Analysis*: Change *Standard solution B* to: *Sample solution B*
DEXTROSE	IDENTIFICATION/*C. Water Determination <921>*	*Second Supplement to USP39–NF34*	8612	18-Nov-2016	1-Dec-2016	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1: Change *Water Determination <921>* to: *Water Determination <921>, Method I*
PLASTIC MATERIALS OF CONSTRUCTION	TEST METHODS	*USP40–NF35*	542	18-Nov-2016	1-Dec-2016	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Physicochemical Tests*/*Acidity or Alkalinity*/*BRP indicator solution*: Change 1.0 mg/mL of bromophenol blue, to:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								1.0 mg/mL of bromothymol blue, AND Line 3 of *Plastic Additives/Polyethylene, Cyclic Olefins, and Polypropylene/Phenolic Antioxidants/Test B*: Change tris(2,4-di-*tert*-butylphenyl) phosphate; to: tris(2,4-di-*tert*-butylphenyl) phosphite; AND Line 1 of *Plastic Additives/Polyethylene,*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Cyclic Olefins, and P oly propyl ene/Phenolic Anti oxidants/Test C/Mobile phase: Change (55:45:5, v/v/v) to: (50:45:5, v/v/v) AND Line 1 of *Plastic Additiv es/ Polyethylene, Cyclic Olefins, and P oly propyl ene/ Nonphenolic Anti oxidant s/ Chromatographi c system/*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Application: Change 20 µL of *Sample solution S10,* reference solution (*o*) and the reference solutions corresponding to to: 20 µL of *Sample solution S10* and the reference solutions corresponding to AND Line 2 of *Plastic Additives/Plasticized Polyvinyl Chloride/USP Reference Standards <11>/USP Plastic Additive 05 RS*: Change Tris(2,4-di-*tert*

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								-butylphenyl) phosphate. to: Tris(2,4-di-*tert* -butylphenyl) phosphite.
BETAXOLOL OPHTHALMIC SOLUTION	IM PUR ITIES/*Organic Impurities*	*USP39–NF34*	2749	18-Nov-2016	1-Dec-2016	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 14 of *Analysis*: Change M_{r1} = molecular weight of betaxolol hydrochloride, 343.89 M_{r2} = molecular weight of betaxolol, 307.43 to: M_{r1} = molecular weight of betaxolol, 307.43 M_{r2} = molecular weight of betaxolol hydrochloride, 343.89
DIGOXIN ORAL SOLUTION	IDENTIFICATIO N/*B.*	*USP39–NF34*	3493	18-Nov-2016	1-Dec-2016	*USP41–NF36*	*Second Supplement to*	Line 1 of *Procedure*:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
							USP40–NF35	Change Proceed as directed for *Procedure* in the test for *Related glycosides* under *Digoxin*, except to omit the use of the *Gitoxin standard solution*. to: Apply 10 µL of the *Test solution* and 10 µL of the *Standard solution* on a line parallel to and about 2.5 cm from the bottom edge of a reversed-phase thin-layer chromatographic plate coated with a 0.25-mm layer of chromatographic silica

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								gel mixture to which is permanently bonded octadecylsilane (C18). Allow the spots to dry, and place the plates in a developing chamber containing a mixture of methanol and water (7:3). Develop the chromatogram until the solvent front has moved about 15 cm above the line of application. Remove the plate, and allow the solvent to evaporate. Spray the plate with *Chloramine T–trichloroacetic acid reagent*, freshly mixed,

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								and heat in an oven at 110° for 10 minutes.
CHONDROITIN SULFATE SODIUM	IMPURITIES	*USP39–NF34*	6566	18-Nov-2016	1-Dec-2016	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 1 of *Residue on Ignition <281>*: Change 20.0%–30.0% to: 20.0%–30.0% on the dried basis
CIPROFLOXACIN	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*Second Supplement to USP39–NF34*	8597	18-Nov-2016	1-Dec-2016	*USP41–NF36*	*Second Supplement to USP40–NF35*	Line 2 of USP Ciprofloxacin Ethylenediamine Analog RS: Change 7-(2-Aminoethylamino)-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-quinolinec

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								arboxylic acid hydrochloride. $C_{15}H_{16}FN_3O_3$ · HCl 341.77

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