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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
TRAMADOL HY	PERFORMANC	*USP42–NF37*	4409	26-Apr-2019	1-May-2019	*NA*	*NA*	In *Cell*: 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
DROCHLORIDE EXTENDED-RELEASE TABLETS	E TESTS/*Dissolution <711>/Test 1/Instrumental conditions*							5 cm to: 5 mm
NEPHELOMETRY AND TURBIDIMETRY	4. INSTRUMENTATION	*USP42–NF37*	7059	26-Apr-2019	1-May-2019	*NA*	*NA*	In paragraph 2: Change silicone diodes to: silicon diodes
CLONIDINE TRANSDERMAL SYSTEM	PERFORMANCE TESTS/*Drug Release <724>/Test 1*	*USP42–NF37*	1084	26-Apr-2019	1-May-2019	*NA*	*NA*	*Apparatus 7*: Change (see *Figure 4a*). to: (see *Figure 5a*).
LEVALBUTEROL INHALATION SOLUTION	IM PURITIES/*Limit of S-Albuterol*	*USP42–NF37*	Online	26-Apr-2019	1-May-2019	*NA*	*NA*	In *Mobile phase*: Change ?*Acetonitrile*, *methanol*, and acetic acid (50:50). To each liter of the solution add 3 mL of acetic acid and 1 mL of triethylamine.? (USP 1-May-2019) to: Acetonitrile and

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								methanol (50:50). To each liter of the solution add 3 mL of acetic acid and 1 mL of triethylamine.
SODIUM BICARBONATE COMPOUNDED INJECTION	ASSAY/ *Procedure for Sodium Bicarbonate*	*USP42–NF37*	4023	26-Apr-2019	1-May-2019	*NA*	*NA*	In *Analysis*: Change Result = [(V_S ? V_B) x N_A x F] x 100 to: Result = [(V_S ? V_B) x N_A x F x 100]/W AND Change F = equivalency factor, 84.01 mg/mL to: F = equivalency factor, 84.01 mg/mEq W = sample weight (mg)
REAGENTS AND REFERENCE TABLES	REAGENT SPECIFICATIONS	*USP42–NF37*	6079	26-Apr-2019	1-May-2019	*NA*	*NA*	In *Beef Extract/ Microbial Content*:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
PHARMACEUTICAL CALCULATIONS IN PHARMACY PRACTICE	19. MEAN KINETIC TEMPERATURE/*19.2 MKT Equation*	*USP42–NF37*	7831	26-Apr-2019	1-May-2019	*NA*	*NA*	Change MT to: NMT In the variable definition list: Change T_n = value for the total number of storage temperatures recorded during the observation period temperature recorded during the nth time period, e.g., *n*th week to: T_n = value for the temperature recorded during the *n*th time period, e.g., *n*th week
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE TABLETS	ASSAY/*Procedure/Chromatographic system*	*First Supplement to USP42–NF37*	Online	26-Apr-2019	1-May-2019	*NA*	*NA*	In *Column*: Change 4.6-mm × 15-cm; 5-µm packing L7. [Not

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								e—Conditioning of the *Column* with *Solution A* and *Solution B* $^{?}(90:10)_{?}$ (ERR 1-Mar-2019) for about 30 min is recommended prior to use.] to: 4.6-mm × 15-cm; 5-µm packing L7
IMIPRAMINE PAMOATE CAPSULES	IMPURITIES/*Organic Impurities*	*USP42–NF37*	Online	26-Apr-2019	1-May-2019	*NA*	*NA*	In *Solution A*: Change *Chromatographic acetonitrile* to: Acetonitrile AND In *Solution B*: Change *chromatographic acetonitrile* to: acetonitrile
MORPHINE SULFATE EXTENDED-RELEASE	IMPURITIES/*Organic Impurities*	*Revision Bulletin (Official November 01, 2018)*	Online	26-Apr-2019	1-May-2019	*NA*	*NA*	In Row 4 of Column 1 of *Table 5*: 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
CAPSULES								Morphine related compound B[b] to: Morphine related compound B (anhydrous)[b]
HYPROMELLOSE PHTHALATE	IMPURITIES/*Chloride and Sulfate <221>, Chloride*	*Harmonization (Official May 01, 2019)* Online		26-Apr-2019	1-May-2019	*NA*	*NA*	In *Analysis*: Change ?Add 1 mL of silver nitrate TS to the *Standard solution* and then add a 50-mL portion of the *Sample solution*. Mix and allow to stand for 5 min protected from direct sunlight. Compare the turbidity of the solutions.?(NF 1-May-2019) to: Add 1 mL of silver nitrate TS to the *Standard solution*. Add 1

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								mL of silver nitrate TS to a 50-mL portion of the *Sample solution*. After mixing, allow each solution to stand for 5 min protected from direct sunlight. Compare the turbidity of the solutions.
ISOPHANE INSULIN HUMAN SUSPENSION	ASSAY/ *Procedure*	*Interim Revision Announcement (Official January 01, 2019)*	Online	29-Mar-2019	1-Apr-2019	*NA*	*NA*	In *Standard solution*: Change USP Insulin Beef RS to: USP Insulin Human RS
SCOPOLAMINE HYDROBROMIDE	IDENTIFICATION/*B.*	*First Supplement to USP41–NF36*	8420	29-Mar-2019	1-Apr-2019	*NA*	*NA*	In *Sample solution*: Change 50 mg/mL of alcohol to: 50 mg/mL in water
PREDNISOLONE SODIUM	*Related compounds*	*USP41–NF36*	3416	29-Mar-2019	1-Apr-2019	*NA*	*NA*	In *Table 1*: Add Prednisolone

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PHOSPHATE								sodium phosphate 1.00 — —
MERCAPTOPURINE	IMPURITIES/*Organic Impurities*	*USP41–NF36*	2587	29-Mar-2019	1-Apr-2019	*NA*	*NA*	Change *Sample solution*: 0.12 mg/mL of Mercaptopurine in *Solution A*. [NOTE—Inject the *Sample solution* within 1 h of preparation.] to: *Sample stock solution*: 0.5 mg/mL of mercaptopurine in a mixture of methanol and *Solution A* (1:9) prepared as follows. Transfer a suitable quantity of Mercaptopurine to an appropriate volumetric flask,

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								add methanol equivalent to 10% of the final volume, and shake to dissolve. Dilute with *Solution A* to volume. *Sample solution*: 0.12 mg/mL of mercaptopurine in *Solution A* from the *Sample stock solution*. [NOTE—Inject the *Sample solution* within 1 h of preparation.]
BUMETANIDE TABLETS	ASSAY/ *Procedure*	*Second Supplement to USP41–NF36*	Online	29-Mar-2019	1-Apr-2019	*NA*	*NA*	In *Sample solution*: Change Nominally 0.05 mg/mL of bumetanide prepared as follows. to: Nominally 125

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
RUTIN	CHEMICAL INFORMATION	*USP41–NF36*	4841	29-Mar-2019	1-Apr-2019	*NA*	*NA*	µg/mL of bumetanide prepared as follows. Change 3-Rhamnoglucoside of 5,7,3',4'-tetrahydroxyflavonol; 2-(3,4-Dihydroxyphenyl)-5,7-dihydroxy-4*H*-chromen-4-one-3-yl 6-*O*-?-L-rhamnopyranosyl-?-D-glucoside [250249-75-3]. to: 3-Rhamnoglucoside of 5,7,3',4'-tetrahydroxyflavonol trihydrate; 2-(3,4-Dihydroxyphenyl)-5,7-dihydroxy-4*H*-chromen-4-one-3-yl 6-*O*-?-L-rhamno

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								pyranosyl-?-D-glucoside trihydrate [250249-75-3].
METAXALONE	IDENTIFICATION/*B.*	*USP41–NF36*	2611	29-Mar-2019	1-Apr-2019	*NA*	*NA*	Change The retention time of the major peak of the *Sample solution* corresponds to that of the *Sample solution*, as obtained in the *Assay*. to: The retention time of the major peak of the *Sample solution* corresponds to that of the *Standard solution*, as obtained in the *Assay*.
MESNA TABLETS	ADDITIONAL REQUIREMENTS/*USP*	*Second Supplement to USP41–NF36*	8906	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In USP Mesna Related Compound A

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
	Reference Standards <11>							RS: Change 2-(Acetylthio)ethane-1-sulfonic acid. $C_4H_8O_4S_2$ 184.22 to: 2-(Acetylthio)ethane-1-sulfonic acid, potassium salt, crystal adduct with potassium chloride. $C_4H_7KO_4S_2$? KCl 296.86 AND In USP Mesna Related Compound B RS: Change 2,2?-Disulfanediylbis(ethane-1-sulfonic acid). $C_4H_{10}O_6S_4$ 282.36 to: 2,2?-Disulfanediylbis(ethane-1-sulfonic acid), dipotassium

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								salt, crystal adduct with sodium chloride. $C_4H_8K_2O_6S_4$? NaCl 416.98
DEXMEDETOMIDINE HYDROCHLORIDE	CHEMICAL INFORMATION	*USP41–NF36*	Online	22-Feb-2019	1-Mar-2019	*NA*	*NA*	This erratum applies to the *USP-NF* ONLINE platform only. See https://www.usp nf.com/sites/def ault/files/usp_pd f/EN/february_2 019_errata_ima ge1.pdf for correction
REAGENTS	*Solutions/Volumetric Solutions/0.01 N Sodium Hydroxide VS*	*USP41–NF36*	5770	22-Feb-2019	1-Mar-2019	*NA*	*NA*	See https://www.usp nf.com/sites/def ault/files/usp_pd f/EN/february_2 019_errata_ima ge1.pdf for correction
FLUDROCORTISONE ACETATE TABLETS	IMPURITIES/*Organic Impurities/Table*	*Second Supplement to USP41–NF36*	8843	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In footnote a: Change 9-Fluoro-11?,17 ,21-trihydroxypr

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							egn-4-ene-3,20-dione 21-acetate. to: 9-Fluoro-11?,17,21-trihydroxypregn-4-ene-3,20-dione.
ANALYTICAL DATA—INTERPRETATION AND TREATMENT	MEASUREMENT PRINCIPLES AND VARIATION	*USP42–NF37*	7129	22-Feb-2019	1-Mar-2019	*NA*	*NA*	See https://www.usp nf.com/sites/default/files/usp_pdf/EN/february_2019_errata_image1.pdf for correction
OXANDROLONE	*Related compounds*	*USP41–NF36*	3072	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In footnote 4 of the second table: Change Methyl-(1,17?-dihydroxy-17?-methyl-1,3-seco-2-nor-5?-androstane-3-oate. to: Methyl 1,17?-dihydroxy-17?-methyl-1,3-seco-2-nor-5?-androstan-3-oate.
CARBINOXAMINE		*Second*	8786	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In *Standard*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
NE MALEATE	PUR ITIES/*Organic Impurities*	*Supplement to USP41–NF36*						*stock solution*: Change (equivalent to 0.05 mg/mL of USP Carbinoxamine Maleate RS free base) and 0.05 mg/mL each of USP Carbinoxamine Related Compound A RS, USP Carbinoxamine Related Compound B RS, and USP Carbinoxamine Related Compound C RS free base to: (equivalent to 0.05 mg/mL of carbinoxamine) and 0.05 mg/mL each of USP Carbinoxamine Related Compound A

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								RS, USP Carbinoxamine Related Compound B RS, and USP Carbinoxamine Related Compound C RS (as the free base) AND In the *Standard solution*: Change (equivalent to 0.001 mg/mL of USP Carbinoxamine Maleate RS free base) and 0.001 mg/mL each of USP Carbinoxamine Related Compound A RS, USP Carbinoxamine Related Compound B RS, and USP Carbinoxamine

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Related Compound C RS free base to: (equivalent to 0.001 mg/mL of carbinoxamine) and 0.001 mg/mL each of USP Carbinoxamine Related Compound A RS, USP Carbinoxamine Related Compound B RS, and USP Carbinoxamine Related Compound C RS (as the free base) AND In the *Analysis*: Change C_S = concentration of USP Carbinoxamine Maleate RS free

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								base to: C_S = concentration of USP Carbinoxamine Maleate RS (as the free base)
PRAZOSIN HYDROCHLORIDE COMPOUNDED ORAL SUSPENSION	ASSAY/ *Procedure*	*Second Supplement to USP41–NF36*	8945	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In the *Mobile phase*: Change tetramethylammonium hydrochloride to: tetramethylammonium hydroxide
DIVALPROEX SODIUM EXTENDED-RELEASE TABLETS	ASSAY/ *Procedure*	*USP41–NF36*	1358	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In *Buffer*: Change 0.5 g/L of citric acid and 0.4 g/L of dibasic sodium phosphate in water to: 0.5 g/L of anhydrous citric acid and 0.4 g/L of anhydrous dibasic sodium

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								phosphate in water
AMITRIPTYLINE HYDROCHLORIDE TABLETS	IMPURITIES/*Organic Impurities*	*Second Supplement to USP41–NF36*	8759	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In *Analysis*: Change r_U = peak response of amitriptyline related compound A, amitriptyline related compound B, or nortriptyline hydrochloride from the *Sample solution* r_S = peak response of amitriptyline related compound A, amitriptyline related compound B, or nortriptyline hydrochloride from the *Standard solution* to: *r*

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								$_U$= peak response of amitriptyline related compound A, amitriptyline related compound B, or nortriptyline from the *Sample solution* r_S = peak response of amitriptyline related compound A, amitriptyline related compound B, or nortriptyline from the *Standard solution*
MESNA	ADDITIONAL REQUIREMENTS/*USP Reference Standards <11>*	*Second Supplement to USP41–NF36*	8904	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In USP Mesna Related Compound A RS: Change 2-(Acetylthio)ethane-1-sulfonic acid. $C_4H_8O_4S_2$

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								184.22 to: 2-(Acetylthio)ethane-1-sulfonic acid, potassium salt, crystal adduct with potassium chloride. $C_4H_7KO_4S_2$? KCl 296.86 AND In USP Mesna Related Compound B RS: Change 2,2?-Disulfanediylbis(ethane-1-sulfonic acid). $C_4H_{10}O_6S_4$ 282.36 to: 2,2?-Disulfanediylbis(ethane-1-sulfonic acid), dipotassium salt, crystal adduct with sodium chloride. 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
ARGATROBAN	CHEMICAL INFORMATION	*USP41–NF36*	346	22-Feb-2019	1-Mar-2019	*NA*	*NA*	$^4H_8K_2O_6S_4$? NaCl 416.98 See https://www.usp nf.com/sites/def ault/files/usp_pd f/EN/february_2 019_errata_ima ge1.pdf for correction
CALCIUM SILICATE	IM PURITIES/*Limit of Lead*	*USP41–NF36*	5240	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In *Lead standard solution*: Change 1000 mg of lead/mL^4 to: 1000 mg of lead/L^4
CARBINOXAMINE MALEATE TABLETS	IM PUR ITIES/*Organic Impurities*	*Second Supplement to USP41–NF36*	8788	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In the *Standard stock solution*: Change USP Carbinoxamine Maleate RS free base) to: carbinoxamine) AND In the *Standard solution*:

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								Change USP Carbinoxamine Maleate RS free base) to: carbinoxamine) AND In the *Analysis*: Change C_S = concentration of USP Carbinoxamine Maleate RS free base to: C_S = concentration of USP Carbinoxamine Maleate RS (as the free base)
REAGENTS	*Reagent S pec ificatio ns/ 7,8-Dihydrofolic Acid*	*Second Supplement to USP41–NF36*	9052	22-Feb-2019	1-Mar-2019	*NA*	*NA*	Change *(L-Glutamic Acid, N-[4-[[(2-Amino-3,4,7,8-tetrahydro-4-pteridinyl)methyl]amino]benzoyl]-),* 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
								(L-Glutamic Acid, N-[4-[[(2-Amino-3,4,7,8-tetrahydro-4-oxo-6-pteridinyl)methyl]amino]benzoyl]-),
METHYLDOPA	SPECIFIC TESTS/*Optical Rotation <781S>*	*USP41–NF36*	2666	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In the *Sample solution*: Change aluminum chloride to: aluminum chloride hexahydrate
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE TABLETS	ASSAY/*Procedure/Chromatographic system*	*Second Supplement to USP41–NF36*	8781	22-Feb-2019	1-Mar-2019	*NA*	*NA*	In *Column*: Change [Note—Conditioning of the *Column* with *Solution A* and *Solution B* (80:20) to: [Note—Conditioning of the *Column* with *Solution A* and *Solution B* (90:10)

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
L104	CHROMATOGRAPHIC COLUMNS/*Packings*	*First Supplement to USP41–NF36*	8503	26-Jan-2019	1-Feb-2019	*USP42–NF37*	*Second Supplement to USP41–NF36*	Add L104—Triazol groups chemically bonded to porous silica particles, 1.5–10 ?m in diameter.
LEFLUNOMIDE	IMPURITIES/*Organic Impurities*/*Procedure 2*	*USP41–NF36*	2353	25-Jan-2019	1-Feb-2019	*NA*	*NA*	Change *Standard solution*: 0.5 µg/mL of USP Leflunomide RS, from the *Standard solution* in *Mobile phase* to: *Standard stock solution*: Proceed as directed in the *Standard solution* in the *Assay*. *Standard solution*: 0.5 µg/mL of USP Leflunomide RS from the

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								Standard stock solution in *Mobile phase*
SALIX SPECIES BARK DRY EXTRACT	INTRODUCTION	*USP42–NF37*	5187	25-Jan-2019	1-Feb-2019	*NA*	*NA*	Delete (This monograph is postponed indefinitely.)
SALIX SPECIES BARK	INTRODUCTION	*USP42–NF37*	5185	25-Jan-2019	1-Feb-2019	*NA*	*NA*	Delete (This monograph is postponed indefinitely.)
SALIX SPECIES BARK POWDER	ADDITIONAL REQUIREMENTS	*USP42–NF37*	5189	25-Jan-2019	1-Feb-2019	*NA*	*NA*	In *Labeling*: Change The label states the Latin binomial(s) of one or several *Salix* species included in the article. to: The label states the Latin binomial(s) of one or several *Salix* species included in the article. Dosage

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								forms prepared with this article should bear the following statement: Not for use in children, women who are pregnant or nursing, or by persons with known sensitivity to aspirin. AND In *USP Reference Standards* <*11*>: Delete (This monograph is postponed indefinitely.)
SALMETEROL INHALATION POWDER	ASSAY/ *Procedure/Analysis*	*First Supplement to USP41–NF36*	Online	25-Jan-2019	1-Feb-2019	*NA*	*NA*	In the definition list: Change M_{r1} = molecular weight of salmeterol free base, 415.75 to: *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
SALIX SPECIES BARK DRY EXTRACT	ADDITIONAL REQUIREMENTS	*USP42–NF37*	5187	25-Jan-2019	1-Feb-2019	*NA*	*NA*	r_1= molecular weight of salmeterol free base, 415.57 In *Labeling*: Change It meets the labeling requirements of *Botanical Extracts* <*565*>. to: It meets the labeling requirements of *Botanical Extracts* <*565*>. Dosage forms prepared with this article should bear the following statement: Not for use in children, women who are pregnant or nursing, or by persons with known sensitivity 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
								aspirin. AND In *USP Reference Standards* <*11*> : Delete (This monograph is postponed indefinitely.)
BENZETHONIUM CHLORIDE	IMPURITIES/*Organic Impurities/Acceptance criteria*	*First Supplement to USP41–NF36*	8297	25-Jan-2019	1-Feb-2019	*NA*	*NA*	In *Total impurities*: Change 1.0% to: NMT 1.0%
SALIX SPECIES BARK	ADDITIONAL REQUIREMENTS	*USP42–NF37*	5185	25-Jan-2019	1-Feb-2019	*NA*	*NA*	In *Labeling*: Change The label states the Latin binomial(s) of one or several *Salix* species included in the article. to: The label states the Latin binomial(s) of one or several *Salix* species

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								included in the article. Dosage forms prepared with this article should bear the following statement: Not for use in children, women who are pregnant or nursing, or by persons with known sensitivity to aspirin. AND In *USP Reference Standards* <*11*>: Delete (This monograph is postponed indefinitely.)
SALMETEROL INHALATION POWDER	PERFORMANCE TESTS	*First Supplement to USP41–NF36*	Online	25-Jan-2019	1-Feb-2019	*NA*	*NA*	In the definition list in *Particle Size Distribution by Cascade Impaction/Analysis*:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
								Change M_{r1} = molecular weight of salmeterol free base, 415.75 to: M_{r1} = molecular weight of salmeterol free base, 415.57 AND In the definition list in *Delivered-Dose Uniformity <601>/Analysis*: Change M_{r1} = molecular weight of salmeterol free base, 415.75 to: M_{r1} = molecular weight of salmeterol free base, 415.57
SALIX SPECIES BARK POWDER	INTRODUCTION	*USP42–NF37*	5189	25-Jan-2019	1-Feb-2019	*NA*	*NA*	Delete (This monograph is postponed indefinitely.)
PANTOPRAZO	IM	*First*	8392	28-Dec-2018	1-Jan-2019	*NA*	*NA*	In *Column*:

Monograph Title	Section	Source Publication	Page Number	Errata Post Date Sort ascending	Errata Official Date	Target Errata Print Publication	Target Online Fix Publication	Description
LE SODIUM	PUR ITIES/*Organic Impurities*/*Test 2*/ *Chromatographic system*	*Supplement to USP41–NF36*						Change 4.6-mm x 12.5-cm; 5-?m packing L1 to: 4-mm x 12.5-cm; 5-?m packing L1
METACRESOL	IDENTIFICATION/*B.*	*USP41–NF36*	2605	28-Dec-2018	1-Jan-2019	*NA*	*NA*	Change The retention time of the major peak of the *Sample solution* corresponds to that of the *Standard solution*, as obtained in the *Assay*. to: The retention time of the metacresol peak of the *Sample solution* corresponds to that of the *Standard solution*, as obtained in 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
VITAMIN A	ADDITIONAL REQUIREMENTS	*USP41–NF36*	4327	28-Dec-2018	1-Jan-2019	*NA*	*NA*	*Assay*. Change Delete the following ?•*USP Reference Standards <11>* USP Retinyl Acetate RS USP Retinyl Palmitate RS ?(CN 1-May-2018) to: •*USP Reference Standards <11>* USP Retinyl Acetate RS USP Retinyl Palmitate RS
ESOMEPRAZOLE MAGNESIUM DELAYED-RELEASE CAPSULES	PERFORMANCE TESTS/ *Dissolution <11>/Test 3/Buffer stage*	*Revision Bulletin (Official March 01, 2018)*	Online	28-Dec-2018	1-Jan-2019	*NA*	*NA*	In the *Standard solution*: Change 0.25 M sodium hydroxide, to: 0.25 N sodium hydroxide, AND In the *Sample 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
								Change 0.25sodium hydroxide, to: 0.25 N sodium hydroxide,

Pagination

- First page « First
- Previous page ‹ Previous
- …
- Page 13
- Page 14
- Page 15
- Page 16
- Page 17
- Page 18
- Page 19
- Page 20
- Page 21
- …
- Next page Next ›
- Last page Last »