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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.
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Monograph Title	Section	Source	Page Number	Errata Post	Errata Official	Target Errata	Target Online	Description
		Publication		Date	Date	Print Publication	Fix Publication	
SODIUM	ASSAY/	USP37–NF32	4727	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	Line 1 of <i>Blank</i> :

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SALICYLATE	<i>Procedure</i>								Change acetic acid to: glacial acetic acid
ESCITALOPRAM TABLETS	PERFORMANCE TESTS/ <i>Dissolution <711>/Test 2</i>	USP37–NF32	2852	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 3 of <i>Buffer</i> : Change phosphoric to: phosphoric acid
THIOTHIXENE	<i>Limit of (E)-thiothixene/Standard preparations</i>	USP37–NF32	4942	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 1 of C: Change Transfer about 200 mg of thiothixene to: Transfer about 200 mg of Thiothixene
HEXACHLOROPHENE LIQUID N/B. SOAP	IDENTIFICATION	USP37–NF32	3231	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 1 of <i>Analysis</i> : Change <i>Sample solution</i> to: <i>Sample</i>
POWDERED ECHINACEA ANGUSTIFOLIA	ADDITIONAL REQUIREMENTS/ <i>Reference Standards <11></i>	USP37–NF32	5346	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 5: Change USP Powdered <i>Echinacea purpurea</i> Extract RS

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MAPROTILINE PERFORMANC HYDROCHLOR E IDE TABLETS TESTS/ <i>Dissolution</i> <711>	USP37–NF32	3655	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	to: USP Powdered <i>Echinacea angustifolia</i> Extract RS Line 3 of <i>Analysis:</i> Change Determine the labeled amount of maprotiline hydrochloride to: Determine the percentage of the labeled amount of maprotiline hydrochloride
POWDERED ECHINACEA PURPUREA EXTRACT	COMPOSITION USP37–NF32	5371	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	Line 14 of <i>Analysis in</i> <i>Content of Total</i> <i>Phenols:</i> Change $C_S =$ concentration of the relevant analyte in the corresponding <i>Standard</i> <i>solution</i>

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							<p>C_U = concentration of <i>Echinacea purpurea</i> in the <i>Sample solution</i> (mg/mL)</p> <p>to:</p> <p>C_S = concentration of the relevant analyte in the corresponding <i>Standard solution</i> (mg/mL)</p> <p>C_U = concentration of Powdered <i>Echinacea purpurea</i> Extract in the <i>Sample solution</i> (mg/mL)</p> <p>AND</p> <p>Line 20 of <i>Analysis in Content of Dodecatetraenoic Acid Isobutylamides</i>: Change</p>

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DESCRIPTION AND SOLUBILITY	<i>Lauric Acid</i>	USP37–NF32 1506	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	<p>C_U = concentration of <i>Echinacea purpurea</i> in the <i>Sample solution</i> (mg/mL)</p> <p>to:</p> <p>C_U = concentration of Powdered <i>Echinacea purpurea</i> Extract in the <i>Sample solution</i> (mg/mL)</p> <p>Line 4: Change emulsifying and/or solubilizing agent; tablet and/or capsule lubricant.</p> <p>to:</p> <p>emulsifying agent; lubricant.</p>
OMEPRAZOLE ASSAY/ ORAL SUSPENSION	<i>Procedure</i>	USP37–NF32 4067	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	<p>Line 2 of <i>Solution A</i>: Change with dilute phosphoric acid</p> <p>to:</p>

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POWDERED HOLY BASIL LEAF EXTRACT	COMPOSITION <i>/Content of Triterpenes</i>	USP37–NF32	5458	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	with dilute sodium hydroxide Line 4 of <i>Standard solution B</i> : Change 0.45-?L to: 0.45-?m
ALCOHOL IN DEXTROSE INJECTION	ASSAY/ <i>Dextrose</i>	USP37–NF32	1637	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 15 of <i>Analysis</i> : Change <i>A</i> = length of the polarimeter tube (mm) to: <i>A</i> = 100 mm divided by the length of the polarimeter tube (mm)
PYRANTEL PAMOATE	ASSAY/ <i>Procedure</i>	USP37–NF32	4491	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 1 of <i>Mobile phase</i> : Change acetic acid to: glacial acetic acid AND Line 1 of <i>Column</i>

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CARBAMAZEPINE EXTENDED-RELEASE TABLETS	USP37–NF32	2123	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	<i>efficiency:</i> Change NLT 8000 theoretical plates to: NLT 8000 theoretical plates for the pyrantel peak Line 2 of <i>Sample stock solution B:</i> Change <i>Standard stock solution</i> to: <i>Sample stock solution A</i>
RIBAVIRIN CAPSULES	PERFORMANCE TESTS/ <i>Dissolution <711>/ Procedure 1/ Chromatographic system</i>	USP37–NF32 4562	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	Line 1 of <i>Column:</i> Change 7-µm packing L17 to: 9-µm packing L17
DIPHENHYDRAMINE CITRATE AND	IMPURITIES/ <i>Limit of Ibuprofen</i>	USP37–NF32 2651	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	Line 7 of <i>Analysis:</i> Change

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IBUPROFEN TABLETS	<i>Related Compound C</i>								R_U = peak area ratio of ibuprofen to valerophenone from the <i>Sample solution</i> R_S = peak area ratio of ibuprofen to valerophenone from the <i>Standard solution</i> to: R_U = peak area ratio of ibuprofen related compound C to valerophenone from the <i>Sample solution</i> R_S = peak area ratio of ibuprofen related compound C to valerophenone from the <i>Standard solution</i>

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SULFACETAMI ASSAY/ DE SODIUM <i>Procedure</i> OPHTHALMIC SOLUTION	USP37–NF32	4766	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	Line 2 of <i>Sample stock solution:</i> Change sulfacetamide to: sulfacetamide sodium AND Line 1 of <i>Sample solution:</i> Change sulfacetamide to: sulfacetamide sodium
GRISEOFULVI N CAPSULES	PERFORMANC E TESTS/ <i>Uniformity of Dosage Units</i> <905>/ <i>Procedure for content uniformity</i>	USP37–NF32 3196	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	Line 6 of <i>Analysis:</i> Change Result = $(A_U/A_S) \times (C_S/C_U) \times P \times 100$ to: Result = $(A_U/A_S) \times (C_S/C_U) \times P \times F \times 100$ AND After <i>P</i> in definition list:

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VALSARTAN	ADDITIONAL REQUIREMENT S/USP Reference Standards <11>	USP37–NF32	5115	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	<p>Add</p> <p><i>F</i> = conversion factor, 0.001 mg/?g</p> <p>Line 2 of USP Valsartan Related Compound A RS: Change (R)-N-Valeryl-N-([2?-(1H-tetrazole-5-yl)bi-phen-4-yl]methyl)valine.</p> <p>to:</p> <p>N-Valeryl-N-([2?-(1H-tetrazole-5-yl)bi-phenyl-4-yl]methyl)-D-valine.</p> <p>AND</p> <p>Line 2 of USP Valsartan Related Compound B RS: Change (S)-N</p>

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							-Butyryl -N -([2?-(1H -tetrazole-5-yl)bi phen-4-yl]methy l)-valine. to: N -Butyryl -N -{[2?-(1H -tetrazole-5-yl)bi phenyl-4-yl]met hyl}-L-valine. AND Line 2 of USP Valsartan Related Compound C RS: Change (S)-N -Valeryl -N-([2'-(1H -tetrazole-5-yl)bi phen-4-yl]methy l)-valine benzyl ester. to: N -Valeryl -N

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LAMIVUDINE AND ZIDOVUDINE TABLETS	IM PURITIES/ <i>Organic Impurities</i>	USP37–NF32	3484	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	-{[2?-(1H-tetrazole-5-yl)bi-phenyl-4-yl]methyl}-L-valine benzyl ester. Line 12 of <i>Analysis</i> : Change unidentified impurity to: unspecified impurity AND Line 19 of <i>Analysis</i> : Change unidentified impurities to: unspecified impurities
POWDERED ECHINACEA PALLIDA	ADDITIONAL REQUIREMENTS/ <i>USP Reference Standards <11></i>	USP37–NF32	5356	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 5: Change USP Powdered <i>Echinacea purpurea</i> Extract RS to: USP Powdered <i>Echinacea pallida</i> Extract

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NYSTATIN	DEFINITION	USP37-NF32	4035	30-May-2014		1-Jun-2014	USP38-NF33	USP38-NF33	RS Line 6: Change extemporaneous solution to: extemporaneous preparation
HOLY BASIL LEAF	COMPOSITION <i>/Content of Triterpenes</i>	USP37-NF32	5454	30-May-2014		1-Jun-2014	USP38-NF33	USP38-NF33	Line 4 of <i>Standard solution B</i> : Change 0.45-?L to: 0.45-?m
DESCRIPTION AND SOLUBILITY	<i>Sodium Acetate</i>	USP37-NF32	1525	30-May-2014		1-Jun-2014	USP38-NF33	USP38-NF33	Line 7: Change transfer liquid. to: transfer ligand.
PROGESTERONE VAGINAL SUPPOSITORIES	ASSAY/ <i>Suppositories in Fatty Acid Base</i>	USP37-NF32	4430	30-May-2014		1-Jun-2014	USP38-NF33	USP38-NF33	Line 1 of <i>System suitability solution</i> : Change Transfer 2.0 mL of <i>System suitability stock solution A</i> and <i>System suitability stock solution B</i> to:

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CIPROFLOXACPERFORMANC IN EXTENDED- E RELEASE TESTS/ TABLETS	<i>First Supplement to USP37–NF32 Dissolution <711>/Test 2</i>	6619	30-May-2014	1-Jun-2014	<i>USP38–NF33</i>	<i>USP38–NF33</i>	Transfer 2.0 mL of each <i>System suitability stock solution A</i> and <i>System suitability stock solution B</i> AND Line 2 of <i>Standard solution A</i> : Change Mix 5.0 of the solution to: Mix 5.0 mL of the solution Line 1 of <i>Standard solution</i> : Change 0.56 mg/mL of USP Ciprofloxacin Hydrochloride RS in <i>Medium</i> to: 0.62 mg/mL of USP Ciprofloxacin

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AMINOSALICYL ASSAY/ LATE SODIUM <i>Procedure</i>	USP37–NF32	1745	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	<p>Hydrochloride RS in <i>Medium</i> AND Line 8 of <i>Analysis</i>: Change $C_S =$ concentration of ciprofloxacin in the <i>Standard solution</i> (mg/mL) to: $C_S =$ concentration of ciprofloxacin hydrochloride in the <i>Standard solution</i> (mg/mL)</p> <p>Line 15 of <i>Analysis</i>: Change $C_U =$ concentration of aminosalicylate in the <i>Sample solution</i> (mg/mL) to: C</p>

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QUINIDINE GLUCONATE	DEFINITION	USP37–NF32	4512	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	<p>$U =$ concentration of Aminosalicylate Sodium in the <i>Sample solution</i> (mg/mL)</p> <p>Line 6: Change quinidine sulfate to: quinidine gluconate</p>
CODEINE PHOSPHATE ORAL SOLUTION	ASSAY/ <i>Procedure</i>	USP37–NF32	2451	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	<p>Line 6 of <i>Analysis</i>: Change Result = $(R_U/R_S) \times (C_S/C_U) \times 100$ to: Result = $(R_U/R_S) \times (C_S/C_U) \times (M_{r1}/M_{r2}) \times 100$ AND After C_U in definition list: Add M_{r1} = molecular weight of codeine phosphate hemihydrate,</p>

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									406.37 M_r = molecular weight of anhydrous codeine phosphate, 397.37
SUFENTANIL CITRATE INJECTION	ASSAY/ <i>Procedure</i>	USP37–NF32	4759	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 11 of <i>Analysis</i> : Change sufentanil to: sufentanil citrate
FOSPHENYTOIN SODIUM INJECTION	ADDITIONAL REQUIREMENT <i>S/USP Reference Standards <11></i>	USP37–NF32	3096	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Before USP Fosphenytoin Sodium RS: Add USP Endotoxin RS
TRIACETIN	ASSAY/ <i>Procedure/Titrimetric system</i>	USP37–NF32	5031	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 2: Change (See <i>Titrimetry <541></i> , <i>Residual Titrations</i> .) to: (See <i>Titrimetry <541></i> .) AND Line 1 of <i>Mode</i> : Change

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HYDROCHLORIM OTHIAZIDE TABLETS	IM PUR ITIES/ <i>Organic Impurities</i>	USP37–NF32	3247	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Direct titration to: Residual titration Line 11 of <i>Analysis:</i> Change C_U = concentration of the <i>Sample solution</i> (?g/mL) to: C_U = nominal concentration of hydrochlorothia zide in the <i>Sample solution</i> (?g/mL)
POWDERED ECHINACEA ANGUSTIFOLIA EXTRACT	ADDITIONAL R EQUIREMENT S/ <i>USP Reference Standards <11></i>	USP37–NF32	5350	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 5: Change USP Powdered <i>Echinacea purpurea</i> Extract RS to: USP Powdered <i>Echinacea angustifolia</i> Extract RS
METHADONE HYDROCHLOR IDE ORAL	OTHER COMPONENTS	USP37–NF32	3744	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 3 of <i>Alcohol Determination:</i>

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SOLUTION									
POWDERED D COMPOSITION ECAFFEINATE /Content of D GREEN TEA Pol EXTRACT	<i>yphe ols/ Chromatographi c system</i>	USP37–NF32	5438	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Change gas–liquid chromatographic procedure to: gas chromatographic procedure Line 1 of <i>Column</i> : Change 4.6-mm 625-cm; to: 4.6-mm x 25-cm;
DESCRIPTION AND SOLUBILITY	<i>Potassium Alginate</i>	USP37–NF32	1520	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 6: Change suspending and/or viscosity agent. to: suspending and/or viscosity-increasing agent.
OXCARBAZEPI NE TABLETS		USP37–NF32	4119	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Row 5 of Column 1 of <i>Table 1</i> : Change Methoxydibenzazepine ^b to:

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ORANGE OIL	ASSAY/ <i>Total Aldehyde Content</i>	USP37–NF32	6091	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Methoxycarbamazepine ^b Line 1 of <i>Sample:</i> Change 5 mL of Oil to: 5 mL of Orange Oil, accurately weighed
ALUMINA, MAGNESIA, CALCIUM CARBONATE, AND SIMETHICONE CHEWABLE TABLETS	ASSAY/ <i>Magnesium Hydroxide</i>	USP37–NF32	1674	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 5 of <i>Magnesium stock solution:</i> Change Transfer 2.0 mL of this solution to a 100-mL volumetric flask to obtain a solution containing 20 ?g/mL of magnesium (Mg) to: Transfer 1.0 mL of this solution to a 100-mL volumetric flask to obtain a

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PYRANTEL PAMOATE	OTHER COMP ONE NTS/ <i>Content of Pamoic Acid</i>	USP37–NF32	4491	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	solution containing 10 ?g/mL of magnesium (Mg) Line 1 of <i>Mobile phase</i> : Change acetic acid to: glacial acetic acid
CHLOROXYLE NOL	ASSAY/ <i>Procedure</i>	USP37–NF32	2308	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 12 of <i>Analysis</i> : Change $C_S =$ concentration of chloroxylenol in the <i>Standard solution</i> (mg/mL) to: $C_S =$ concentration of USP Chloroxylenol RS in the <i>Standard solution</i> (mg/mL)
RITONAVIR	IMPURITIES	USP37–NF32	4601	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Row 13 of Column 1 of

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DOXEPIN HYD ASSAY/ ROCHLORIDE <i>Procedure</i> ORAL SOLUTION	USP37–NF32	2713	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	<p><i>Table 2:</i> Change 3-Epimer^m to: 3<i>R</i>-Epimer^m AND Row 16 of Column 1 of <i>Table 2:</i> Change 5-Epimer^p to: 5<i>R</i>-Epimer^p AND Row 17 of Column 1 of <i>Table 2:</i> Change Valine urea analog^q to: Diacyl valine urea^q</p> <p>Line 1 of <i>Standard solution:</i> Change Dilute 4.0 of <i>Standard stock solution</i> to:</p>

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TAMSULOSIN HYDROCHLORIDE CAPSULES	PERFORMANCE TESTS/ <i>Dissolution</i> <711>/Test 4	USP37–NF32	4830	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Dilute 4.0 mL of <i>Standard stock solution</i> Line 9 of the second calculation: Change V = volume of <i>Buffer stage medium</i> , 1000 mL to: V = volume of <i>Buffer stage medium</i> , 500 mL
GRISEOFULVIN TABLETS	ASSAY/ <i>Procedure</i>	USP37–NF32	3198	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 13 of <i>Analysis</i> : Change P = potency of griseofulvin in USP Griseofulvin RS (?g/mL) to: P = potency of griseofulvin in USP Griseofulvin RS (?g/mg)
ECHINACEA	ADDITIONAL R	USP37–NF32	5343	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 5: Change

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NGUSTIFOLIA	EQUIREMENT <i>S/USP Reference Standards <11></i>								USP Powdered <i>Echinacea purpurea</i> Extract RS to: USP Powdered <i>Echinacea angustifolia</i> Extract RS
LINDANE CREAM	ASSAY/ <i>Proce dure/ Chromatographi c system/ Temperatures</i>	USP37–NF32	3561	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	After the <i>Injection port</i> subsection: Add a new subsection <i>Detector. 250°</i>
POWDERED ECHINACEA PALLIDA EXTRACT	ADDITIONAL R EQUIREMENT <i>S/USP Reference Standards <11></i>	USP37–NF32	5359	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 5: Change USP Powdered <i>Echinacea purpurea</i> Extract RS to: USP Powd ered <i>Echinacea pallida</i> Extract RS
DESCRIPTION AND SOLUBILITY	<i>Carmellose</i>	USP37–NF32	1486	30-May-2014		1-Jun-2014	USP38–NF33	USP38–NF33	Line 4: Change Suspending and/or viscosity increasing

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OLMESARTAN CHEMICAL MEDOXOMIL INFORMATION	USP37–NF32	4057	30-May-2014	1-Jun-2014	USP38–NF33	USP38–NF33	agent; to: Suspending and/or viscosity-increasing agent; Line 3 of the chemical name: Change methyl ester to: methyl ester; (5-Methyl-2-oxo-1,3-dioxol-4-yl)methyl 1-[[2?-(1H)-tetrazol-5-yl]biphenyl-4-yl]methyl}-4-(2-hydroxypropan-2-yl)-2-propyl-1H-imidazole-5-carboxylate

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