C No Droduct /	Specific Test Derformed	Test Mathed Specification	Donas of Tooting	
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Accreditation Standard	ISO/IEC 17025: 2005			
Laboratory	Akums Drugs and Pharmaceuticals Ltd. Plant-1, Quality Control Laboratory, 19-20-21, Sector-6A, I.I.E., SIDCUL, Ranipur, Haridwar			

S.No.	Product /	Specific Test Performed	Test Method Specification	Range of Testing /
	Material of Test		against which tests are	Limits of Detection
			performed	

I. DRUGS & PHARMACEUTICALS

1. General Tests of Pharmaceuticals Dosage Forms

a.	Tablets	i. Description	GTP-QC-003	Comparative (Qualitative)
		ii. Average Weight	GTP-QC-003	-
		iii. Uniformity of Weight	IP 2010 As per respective STPs	-
		iv. Disintegration Test	IP 2010 As per respective STPs	10 seconds - 99 minutes 59 seconds
		v. Uniformity of Dispersion (Only for Dispersible tablets)	IP 2010 As per respective STPs	Qualitative test
b.	Capsules	i. Description of capsule & Contents	GTP-QC-003	Comparative (Qualitative)
		ii. Average Weight	GTP-QC-003	-
		iii. Uniformity of Weight	IP 2010 As per respective STPs	-
		iv. Disintegration Test	IP 2010 As per respective STPs	10 seconds - 99 minutes 59 seconds

Laboratory	Akums Drugs and Pharmaceuticals Ltd. Plant-1, Quality Control Laboratory, 19-20-21, Sector-6A, I.I.E., SIDCUL, Ranipur, Haridwar				
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Discipline	Chemical Testing	Issue Date	04.09.2013		
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are	Range of Testing / Limits of Detection
			performed	

B. Pharmaceuticals Finished Product Having Specific Test

		01		
1.	Albendazole Tablets IP	General Tests for Tablets	S.No. 1a-i,ii, iii, iv	As general test S.No. 1a-i,ii, iii, iv
		Identification	A) By TLC B) By UV IP 2010 STP/FP/ 40000128	Comparative (Qualitative)
		Assay	By UV IP 2010 STP/FP/ 40000128	70% to 130%
2.	Albendazole Tablets USP	General Tests for Tablets	S.No. 1a-i,ii, iii, iv	As general test S.No. 1a-i,ii, iii, iv
		Identification	A) By UV B) By HPLC USP35 NF30 STP/FP/190000347	Comparative (Qualitative)
		Dissolution	By UV USP35 NF30 STP/FP/190000347	D-25% to 100%
		Assay	By HPLC USP35 NF30 STP/FP/190000347	70% to 130%
3.	Ascorbic acid Tablets USP	General Tests for Tablets	S.No. 1a-i,ii, iii	As general test S.No. 1a-i,ii, iii
		Dissolution	A) By chemicallyB) By chemicallyUSP35 NF30STP/FP/40000194	D-25% to 100%
3.		General Tests for Tablets	By HPLC USP35 NF30 STP/FP/190000347 S.No. 1a-i,ii, iii A) By chemically B) By chemically USP35 NF30	As general test S.No. 1a-i,ii, iii

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Disc	ipline	Chemical Testing		Issue Date	e 04.09.2013		
Cert	ificate Number	T-1997		Valid Until	03.09.2015		
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests are performed		ange of Testing / mits of Detection		
	Ascorbic acid Tablets USP	Assay	By HPLC USP35 NF30 STP/FP/40000194	70	% to 130%		
4.	Aceclofenac Tablets IP	General Tests for Tablets	S.No. 1a-i,ii, iii		general test No. 1a-i,ii, iii		
		Identification	By HPLC IP 2010 STP/FP/ 40000034		omparative ualitative)		
		Dissolution	By UV IP 2010 STP/FP/ 40000034	D-	25% to 100%		
		Related Substances	By HPLC IP 2010 STP/FP/ 40000034	0.0	05% to 4.0%		
		Assay	By HPLC IP 2010, P-771 STP-AKUMS/STP/068	70	% to 130%		
5.	Alprazolam Tablets IP	General Tests for Tablets	S.No. 1a-i,ii, iii, iv		general test No. 1a-i,ii, iii, iv		
		Identification	By HPLC IP 2010 STP/FP/ 40000136		omparative ualitative)		
		Uniformity of content	By HPLC IP 2010 STP/FP/ 40000136	70	% to 130%		

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Accr	editation Standard	ISO/IEC 17025: 2005			
Disc	ipline	Chemical Testing	Is	sue Date	04.09.2013
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		
	Alprazolam Tablets IP	Assay	By HPLC IP 2010 STP/FP/ 40000136	70%	to 130%
6.	Amlodipine Besilate Tablets IP	General Tests for Tablets	S.No. 1a-i,ii, iii	-	eneral test). 1a-i,ii, iii
		Identification	By HPLC IP 2010 STP/FP/ 40004941		parative alitative)
		Related substances	By HPLC IP 2010 STP/FP/ 40004941	0.05	% to 1.0%
		Dissolution	By UV IP 2010 STP/FP/ 40004941	D-25	5% to 100%
		Uniformity of content	By HPLC IP 2010 STP/FP/ 40004941	70%	to 130%
		Assay	By HPLC IP 2010 STP/FP/ 40004941	70%	to 130%
7.	Aceclofenac Sustained Release Tablets	General Tests for Tablets	S.No. 1a-i,ii, iii, iv		eneral test 5. 1a-i,ii, iii, iv
		Identification	By HPLC STP/FP/ 40000035		parative llitative)
		Dissolution at different stages	By UV STP/FP/ 40000035	10%	to 150%

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Accr	reditation Standard	ISO/IEC 17025: 2005				
Disc	ipline	Chemical Testing		Issue Date	04.09.2013	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests an performed		nge of Testing / nits of Detection	
	Aceclofenac Sustained Release Tablets	Assay	By HPLC STP/FP/ 40000035	70%	% to 130%	
8.	Amoxycillin Capsules BP	General Tests for Capsules	S.No. 1b-i, ii, iii		general test o. 1b-i, ii, iii	
		Identification	A) By IR B) By TLC BP 2013 STP/FP/40000161		nparative alitative)	
		Related substances	By HPLC BP 2013 STP/FP/40000161	0.0	5% to 10.0%	
		Assay	By HPLC BP 2013 STP/FP/40000161	709	6 to 130%	
9.	Ampicillin Capsules IP	General Tests for Capsules	S.No. 1b-i, ii, iii		general test o. 1b-i, ii, iii	
		Identification	A) By IR B) By HPLC IP 2010 STP/FP/ 40001051		nparative alitative)	
		Dissolution	By UV IP 2010, IP 2010 STP/FP/ 40001051	D-2	5% to 100%	
		Assay	By HPLC IP 2010, IP 2010 STP/FP/ 40001051	709	6 to 130%	

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Ассі	reditation Standard	ISO/IEC 17025: 2005				
Disc	ipline	Chemical Testing		Issue Date	04.09.2013	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificat against which tests are performed		nge of Testing / hits of Detection	
10.	Atenolol Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv	
		Identification	A) By IR B) By UV IP 2010 STP/FP/ 40000218		nparative alitative)	
		Related substances	By HPLC IP 2010 STP/FP/ 40000218	0.05	5% to 1.0%	
		Assay	By UV IP 2010 STP/FP/ 40000218	70%	6 to 130%	
11.	Acetyl salicylic acid & magnesium hydroxide Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv	
	nydroxide 1 ablets	Identification	By HPLC STP/FP/40000076		nparative alitative)	
		Assay	By HPLC STP/FP/40000076	70%	6 to 130%	
12.	Azithromycin Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test o. 1a-i, ii, iii	
		Identification	By HPLC IP 2010 STP/FP/ 40001908		nparative alitative)	
		Related substances	By HPLC IP 2010 STP/FP/ 40001908	0.05	5% to 10.0%	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / ts of Detection
	Azithromycin Tablets IP	Dissolution	By HPLC IP 2010 STP/FP/ 40001908	D-25	% to 100%
		Water	By KF IP 2010 STP/FP/ 40001908	>0.29	%
		Assay	By HPLC IP 2010 STP/FP/ 40001908	70%	to 130%
13.	Amoxicillin Dispersible Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv, v	-	eneral test 1a-i, ii, iii, iv, v
		Identification	By HPLC IP 2010 STP/FP/ 40007707		parative litative)
		Assay	By HPLC IP 2010 STP/FP/ 40007707	70%	to 130%
14.	Amoxycillin & Dicloxacillin Capsules	General Tests for Capsules	S.No. 1b-i, ii, iii, iv		eneral test 1b-i, ii, iii, iv
		Identification	By HPLC STP/FP/ 40000300		parative litative)
		Assay	By HPLC STP/FP/ 40000300	70%	to 130%

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Disc	ipline	Chemical Testing	l	ssue Date	04.09.2013	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificati against which tests are performed		ge of Testing / its of Detection	
15.	Ampicillin & Dicloxacillin Capsules	General Tests for Capsules	S.No. 1b-i, ii, iii, iv		eneral test 5. 1b-i, ii, iii, iv	
		Identification	By HPLC STP/FP/ 40000398		parative llitative)	
		Assay	By HPLC STP/FP/ 40000398	70%	to 130%	
16.	Amoxycillin & Potassium Clavulanate	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 5. 1a-i, ii, iii	
	Tablets IP	Identification	By HPLC IP 2010 STP/FP/ 40000022		parative llitative)	
		Dissolution	By HPLC IP 2010 STP/FP/ 40000022	D-25	5% to 100%	
		Uniformity of Content	By HPLC IP 2010 STP/FP/ 40000022	70%	to 130%	
		Water	By KF IP 2010 STP/FP/ 40000022	>0.2	%	
		Assay	By HPLC IP 2010 STP/FP/ 40000022	70%	to 130%	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specific against which tests a performed		Range of Testing / Limits of Detection		
17.	Aceclofenac & Serratiopeptidase Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		As general test S.No. 1a-i, ii, iii, iv		
	1 ablets	Identification	As per Assay STP/FP/ 40009525		Comparative (Qualitative)		
		Assay	By UV STP/FP/ 40009525		70% to 130%		
18.	Aceclofenac & Paracetamol, Serratiopeptidase	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		As general test S.No. 1a-i, ii, iii, iv		
	Tablets	Identification	As per Assay STP/FP/ 40000038		Comparative (Qualitative)		
		Assay	By UV STP/FP/ 40000038	,	70% to 130%		

19.	Atorvastatin & Fenofibrate Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	As general test S.No. 1a-i, ii, iii, iv
		Identification	By HPLC STP/FP/ 40007503	Comparative (Qualitative)
		Uniformity of Content for Atorvastatin	By HPLC STP/FP/ 40007503	70% to 130%
		Assay	By HPLC STP/FP/ 40007503	70% to 130%
20.	Amlodipine besilate tablets	General Tests for Tablets	S.No. 1a-i, ii, iii	As general test S.No. 1a-i, ii, iii
		Identification	By HPLC (Assay) STP/FP/40000664	Comparative (Qualitative)

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specific against which tests a performed		nge of Testing / hits of Detection			
	Amlodipine besilate tablets	Dissolution	By UV STP/FP/40000664	D-2	5% to 100%			
		Related substances	By HPLC STP/FP/40000664	0.05	5% to 10.0%			
		Assay	By HPLC STP/FP/40000664	70%	o to 130%			
21.	Amlodipine & Atenolol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv			
		Identification	By HPLC STP/FP/40000155		nparative alitative)			
		Assay	By HPLC STP/FP/40000155	70%	o to 130%			
22.	Amlodipine & Atorvastatin Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv			
		Identification	By HPLC STP/FP/40009392		nparative alitative)			
		Uniformity of Content	By HPLC STP/FP/40009392	70%	o to 130%			
		Assay	By HPLC STP/FP/40009392	70%	o to 130%			
23.	Amlodipine & Losartan potassium Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv			
	1 abicio 11	Identification	By HPLC IP-2010 STP/FP/400056		nparative alitative)			

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Accr	reditation Standard	ISO/IEC 17025: 2005				
Disc	ipline	Chemical Testing	Issue	e Date	04.09.2013	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / ts of Detectior	
	Amlodipine & Losartan potassium Tablets IP	Uniformity of Content	By HPLC IP-2010 STP/FP/40005667	70%	to 130%	
	Tablets Ir	Dissolution	By HPLC IP-2010 STP/FP/40005667	D-25	% to 100%	
		Assay	By HPLC IP-2010 STP/FP/40005667	70%	to 130%	
24.	Amlodipine & Telmisartan Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	As general test S.No. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/ 40008064		parative litative)	
		Uniformity of Content	By HPLC STP/FP/ 40008064	70%	to 130%	
		Assay	By HPLC STP/FP/ 40008064	70%	to 130%	
25.	Atorvastatin calcium tablets	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test . 1a-i, ii, iii, iv	
		Identification	By HPLC (Assay) STP/FP/40001168		parative litative)	
		Dissolution	By UV STP/FP/40001168	D-25	% to 100%	
		Related substances	By HPLC STP/FP/40001168	0.05%	% to 10.0%	
		Uniformity of Content	By HPLC STP/FP/40001168	70%	to 130%	

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Ассі	editation Standard	ISO/IEC 17025: 2005					
Disc	ipline	Chemical Testing		Issue Date	04.09.2013		
Cert	ificate Number	T-1997		Valid Until	03.09.2015		
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests ar performed		ge of Testing / its of Detection		
		Assay	By HPLC STP/FP/40001168	70%	to 130%		
26.	Amlodipine & Olmesartan	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
	Medoxomil Tablets	Identification	By HPLC STP/FP/ 40010522		nparative alitative)		
		Assay	By HPLC STP/FP/ 40010522	70%	to 130%		
27.	Artemether & Lumefantrine Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/ 40000735		nparative alitative)		
		Assay	By HPLC STP/FP/ 40000735	70%	to 130%		
28.	Aceclofenac & Paracetamol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	By UV STP/FP/40000051	0051 Comparative (Qualitative)			
		Assay	By UV STP/FP/40000051	1 70%	to 130%		
29.	Aceclofenac & Tramadol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	As per Assay STP/FP/ 40009910		nparative alitative)		

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Disc	cipline	Chemical Testing	Is	sue Date	04.09.2013		
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection		
	Aceclofenac & Tramadol Tablets	Uniformity of Content	By HPLC STP/FP/ 40009910	70%	to 130%		
		Assay	By HPLC STP/FP/ 40009910	70%	to 130%		
30.	Aceclofenac, Paracetamol & Chlorzoxazone	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
	Tablets	Identification	As per assay STP/FP/40003724		Comparative (Qualitative)		
		Assay	By UV STP/FP/40003724	70%	to 130%		
31.	Aceclofenac, Paracetamol &	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 5. 1a-i, ii, iii		
	Rabeprazole Tablets	Identification	By HPLC STP/FP/40009172		parative llitative)		
		Dissolution	By HPLC STP/FP/40009172	D-25	D-25% to 100%		
		Uniformity of Content for Rabeprazole & Aceclofenac	By HPLC STP/FP/40009172	70%	to 130%		
		Assay	By HPLC STP/FP/40009172	70%	to 130%		
32.	Albendazole & Ivermectin Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	A) For Albendazole- By UV B) For Ivermectin By HPLC STP/FP/ 40001089		parative litative)		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection
	Aceclofenac & Tramadol Tablets	Uniformity of Content for Ivermectin	By HPLC STP/FP/ 40001089	70%	to 130%
		Assay	A) For Albendazole- By UV B) For Ivermectin By HPLC STP/FP/ 40001089	70%	to 130%
33.	Amlodipine Besilate & Metoprolol Succinate (ER)	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test o. 1a-i, ii, iii
	Tablets	Identification	By HPLC STP/FP/ 40012171		parative llitative)
		Dissolution	By HPLC STP/FP/ 40012171	Aml 10%	5% to 100% (For odipine) to 100% Metoprolol)
		Uniformity of Content	By HPLC STP/FP/ 40012171	70%	to 130%
		Assay	By HPLC STP/FP/ 40012171	70%	to 130%
34.	Bisacodyl Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv
		Identification	By HPLC STP/FP/40000329		parative llitative)
		Uniformity of Content	By HPLC STP/FP/40000329	70%	to 130%

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Acc	reditation Standard	ISO/IEC 17025: 2005	ISO/IEC 17025: 2005						
Disc	cipline	Chemical Testing		Issue Date	04.09.2013				
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificat against which tests are performed		nge of Testing / hits of Detection				
		Assay	By HPLC STP/FP/40000329	70%	5 to 130%				
35.	Betamethasone Sodium Phosphate Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv				
		Identification	A) By TLC B) By Chemically IP 2010 STP/FP/ 40000308		nparative alitative)				
		Uniformity of Content	By HPLC IP 2010 STP/FP/ 40000308	70%	6 to 130%				
		Assay	By HPLC IP 2010 STP/FP/ 40000308	70%	5 to 130%				
36.	Bisoprolol Fumarate Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test o. 1a-i, ii, iii				
		Identification	A) By TLC USP35 NF30 STP/FP/ 40011334	Comparative (Qualitative)					
		Dissolution	By HPLC USP35 NF30 STP/FP/ 40011334	D-2	5% to 100%				
		Uniformity of Dosage unit	By HPLC USP35 NF30 STP/FP/ 40011334	70%	6 to 130%				

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Accr	reditation Standard	ISO/IEC 17025: 2005					
Disc	ipline	Chemical Testing	lse	sue Date	04.09.2013		
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / ts of Detection		
	Bisoprolol Fumarate Tablets USP	Assay	By HPLC USP35 NF30 STP/FP/ 40011334	70%	to 130%		
37.	Calcium Carbonate & Vitamin D ₃ Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	-	eneral test . 1a-i, ii, iii, iv		
		Identification	As per Assay STP/FP/ 40000369		parative litative)		
		Assay	By Titration (For Calcium) By HPLC (For Vitamin D ₃) STP/FP/ 40000369	70%	to 130%		
38.	Ciprofloxacin Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test . 1a-i, ii, iii, iv		
		Identification	A) By HPLC B) By TLC IP 2010 STP/FP/ 40000476		parative litative)		
		Dissolution	By UV IP 2010 STP/FP/ 40000476	D-25	% to 100%		
		Assay	By HPLC IP 2010 STP/FP/ 40000476	70%	to 130%		
39.	Clopidogrel Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii	-	eneral test . 1a-i, ii, iii		
		Identification	By HPLC IP 2010 STP/FP/ 40000562		parative litative)		

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	Clopidogrel Tablets IP	Dissolution	By UV IP 2010 STP/FP/ 40000562	D-25	5% to 100%	
		Related substances	By HPLC IP 2010 STP/FP/ 40000562	0.05	% to 5.0%	
		Assay	By HPLC IP 2010 STP/FP/ 40000562	70%	to 130%	
40.	Cefixime Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 5. 1a-i, ii, iii	
		Identification	By HPLC IP 2010 STP/FP/ 40012002		parative litative)	
		Dissolution	By UV IP 2010 STP/FP/ 40012002	D-25	5% to 100%	
		Water	By KF IP 2010 STP/FP/ 40012002	>0.2	%	
		Assay	By HPLC IP 2010 STP/FP/ 40012002	70%	to 130%	
41.	Cefixime Dispersible Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, v		eneral test 9. 1a-i, ii, iii	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection	
	Cefixime Dispersible Tablets	Identification	By HPLC STP/FP/ 40000445		parative litative)	
		Dissolution	By UV STP/FP/ 40000445	D-25	5% to 100%	
		Assay	By HPLC STP/FP/ 40000445	70%	to 130%	
42.	Cefixime & Ofloxacin Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 9. 1a-i, ii, iii	
		Identification	By HPLC STP/FP/ 40000444		parative litative)	
		Dissolution	By HPLC STP/FP/ 40000444	D-25	5% to 100%	
		Assay	By HPLC STP/FP/ 40000444	70%	to 130%	
43.	Cefixime & Ornidazole Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 9. 1a-i, ii, iii, iv	
		Identification			parative litative)	
		Assay	By HPLC STP/FP/ 40001269	70%	to 130%	
44.	Cefixime & Dicloxacillin Sodium Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 9. 1a-i, ii, iii	
		Identification	By HPLC STP/FP/ 40003762	Comparative (Qualitative)		

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S.No.	Product / Material of Test				ge of Testing / its of Detection			
	Cefixime & Dicloxacillin Sodium Tablets	Dissolution	By HPLC STP/FP/ 40003762	D-25% to 100%				
	Tablets	Assay	By HPLC STP/FP/ 40003762	70% to 130%				
45.	Cefixime & Potassium Clavulanate Tablets	General Tests for Tablets S.No. 1a-i, ii, iii			eneral test 9. 1a-i, ii, iii			
		Identification	By HPLC STP/FP/ 40000441		parative litative)			
		Dissolution	By HPLC STP/FP/ 40000441	D-25	% to 100%			
		Water	By KF STP/FP/ 40000441	>0.2	%			
		Assay	By HPLC STP/FP/ 40000441	70%	to 130%			
46.	Cefadroxil Dispersible Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv, v		eneral test 9. 1a-i, ii, iii, iv, v			
		Identification	By HPLC STP/FP/ 40009664		parative litative)			
		Water	By KF STP/FP/ 40009664	>0.2	%			
		Assay	By HPLC STP/FP/ 40009664	70%	to 130%			

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests are performed		nge of Testing / nits of Detection
47.	Cefadroxil Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test No. 1a-i, ii, iii
		Identification	By TLC IP 2010 STP/FP/ 40000588		mparative ualitative)
		Dissolution	By HPLC IP 2010 STP/FP/ 40000588	D-	25% to 100%
		Water	By KF IP 2010 STP/FP/ 40000588	>0	2%
		Assay	By HPLC IP 2010 STP/FP/ 40000588	70	% to 130%
48.	Ciprofloxacin & Tinidazole Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test No. 1a-i, ii, iii, iv
		Identification	By UV (Assay) STP/FP/ 40009357		mparative ualitative)
		Assay	By UV STP/FP/ 40009357	70	% to 130%
49.	Ciprofloxacin & Ornidazole Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii		general test No. 1a-i, ii, iii
		Identification	By HPLC STP/FP/ 40010241		mparative ualitative)

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	Cefadroxil Dispersible Tablets	Dissolution	By HPLC STP/FP/ 40010241	D-2.	5% to 100%
		Related substances	By HPLC STP/FP/ 40010241	0.05	% to 10%
		Assay	By HPLC STP/FP/ 40010241	70%	to 130%
50.	Citicoline Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test 5. 1a-i, ii, iii, iv
		Identification	By HPLC STP/FP/ 40000519		nparative alitative)
		Assay	By HPLC STP/FP/ 40000519	70%	to 130%
51.	Clarithromycin Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test 5. 1a-i, ii, iii
		Identification	By HPLC USP 35 NF30 STP/FP/40008188		nparative alitative)
		Dissolution	By HPLC USP 35 NF30 STP/FP/40008188	D-2.	5% to 100%
		Loss on Drying	USP 35 NF30 STP/FP/40008188	<6.0	9%
		Assay	By HPLC USP 35 NF30 STP/FP/40008188	70%	to 130%

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificatior against which tests are performed		ge of Testing / ts of Detection		
52.	Cefuroxime Axetil Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 1a-i, ii, iii		
		Identification	A) By IR B) By HPLC IP 2010 STP/FP/ 40001952		parative litative)		
		Dissolution	By UV IP 2010 STP/FP/ 40001952	D-25	% to 100%		
		Related substances	By HPLC IP 2010 STP/FP/ 40001952	0.059	% to 4.0%		
		Assay	By HPLC IP 2010 STP/FP/ 40001952	70%	to 130%		
53.	Cefpodoxime Proxetil Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 1a-i, ii, iii		
		Identification	By HPLC USP 35 NF30 STP/FP/190000565		parative litative)		
		Dissolution	By HPLC USP 35 NF30 STP/FP/190000565	D-25	% to 100%		
		Loss on Drying	USP 34 NF30 STP/FP/190000565	<5.0	%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection		
	Cefpodoxime Proxetil Tablets USP	Assay	By HPLC USP 35 NF30 STP/FP/190000565	70%	to 130%		
54.	Cephalexin Capsules BP	General Tests for Capsules	S.No. 1b-i, ii, iii		eneral test 5. 1b-i, ii, iii		
		Identification	A) By IRB) By TLCC) By chemicallyBP2013STP/FP/40000429		parative llitative)		
		Dissolution	By UV BP2013 STP/FP/40000429	D-25	5% to 100%		
		Related substances	By TLC BP2013 STP/FP/40000429		parative llitative)		
		Assay	By HPLC BP2013 STP/FP/40000429	70%	to 130%		
55.	Cetirizine Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 5. 1a-i, ii, iii		
		Identification	By HPLC IP 2010 STP/FP/ 40007190		parative llitative)		
		Dissolution	By UV IP 2010 STP/FP/ 40007190	D-25	5% to 100%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificatio against which tests are performed		ige of Testing / its of Detectior	
	Cetirizine Tablets IP	Related substances	By HPLC IP 2010 STP/FP/ 40007190	0.05	% to 2.0%	
		Uniformity of content	By HPLC IP 2010 STP/FP/ 40007190	70%	to 130%	
		Assay	By HPLC IP 2010 STP/FP/ 40007190	70%	to 130%	
56.	Chloramphenicol Capsules IP	General Tests for Capsules	S.No. 1b-i, ii, iii		general test 5. 1b-i, ii, iii	
		Identification	A) By IR B) By Chemically IP 2010 STP/FP/ 40009764		nparative alitative)	
		Specific Optical Rotation	IP 2010 STP/FP/ 40009764	0° -	360°	
		Dissolution	By UV IP 2010 STP/FP/ 40009764	D-2:	5% to 100%	
		Assay	By UV IP 2010 STP/FP/ 40009764	70%	to 130%	
57.	Chlorpheniramine Maleate, Paracetamol & Phenylephrine HCl Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii		general test 5. 1a-i, ii, iii	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
	Chlorpheniramine Maleate, Paracetamol & Phenylephrine HCl	Identification	By HPLC STP/FP/ 40008182	Comparative (Qualitative)		
	Tablets	Uniformity of content (Chlorpheniramine & Phenylephrine HCl)	By HPLC STP/FP/ 40008182	70% to 130%		
		Assay	By HPLC STP/FP/ 40008182	70% to 130%		
58.	Calcium Citrate Maleate & Vitamin	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	As general test S.No. 1a-i, ii, iii, iv		
	D ₃ Tablets	Identification	By HPLC STP/FP/ 40001072	Comparative (Qualitative)		
		Assay	By HPLC STP/FP/ 40001072	70% to 130% (For Calcium Citrate Maleate), Not less than 70% (For Vitamin D_3)		
59.	Cetirizine & Ambroxol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	As general test S.No. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/ 40002204	Comparative (Qualitative)		
		Uniformity of content (Cetirizine)	By HPLC STP/FP/ 40002204	70% to 130%		
		Assay	By HPLC STP/FP/ 40002204	70% to 130%		

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60.	Calcium Citrate Maleate, Calcitriol, Magnesium & Zinc Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		As general test S.No. 1a-i, ii, iii, iv
		Identification	As per Assay STP/FP/ 40011955		Comparative (Qualitative)
		Assay	By HPLC (Calcitriol), By Titration (Calcium Citrate Maleate Magnesium), By UV (Zinc) STP/FP/ 40011955),	70% to 130%
61.	Cinnarizine & Domperidone Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		As general test S.No. 1a-i, ii, iii, iv
		Identification	By HPLC (Assay) STP/FP/ 40000713		Comparative (Qualitative)
		Assay	By HPLC STP/FP/ 40000713		70% to 130%
62.	Clonazepam & Propranolol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		As general test S.No. 1a-i, ii, iii, iv
		Identification	As per Assay STP/FP/ 40004698		Comparative (Qualitative)
		Assay	By HPLC STP/FP/ 40004698		70% to 130%
63.	Clonazepam Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		As general test S.No. 1a-i, ii, iii

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	Clonazepam Tablets IP	Identification	A) By IR B) By HPLC IP 2010 STP/FP/ 40008584		parative llitative)		
		Dissolution	By HPLC IP 2010 STP/FP/ 40008584	D-25	5% to 100%		
		Related substances	By HPLC IP 2010 STP/FP/ 40008584	0.05	% to 4.0%		
		Uniformity of content	By HPLC IP 2010 STP/FP/ 40008584	70%	to 130%		
		Assay	By HPLC IP 2010 STP/FP/ 40008584	70%	to 130%		
64.	Cyproheptadine Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	 A) By IR B) By TLC (Related substances) C) By Chemically IP 2010 STP/FP/ 40001223 		parative litative)		
		Related substances	By TLC IP 2010 STP/FP/ 40001223		parative litative)		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection	
	Cyproheptadine Tablets IP	Uniformity of content	By UV IP 2010 STP/FP/ 40001223	70%	to 130%	
		Assay	By UV IP 2010 STP/FP/ 40001223	70%	to 130%	
65.	Diacerein Capsules IP	General Tests for Capsules	S.No. 1b-i, ii, iii, iv		eneral test b. 1b-i, ii, iii, iv	
		Identification	By HPLC IP 2010 STP/FP/ 40010635		parative llitative)	
		Assay	By HPLC IP 2010 STP/FP/ 40010635	70%	to 130%	
66.	Diclofenac Potassium, Serratiopeptidase	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv	
	Tablets	Identification	By UV(Assay) STP/FP/ 40001988		parative llitative)	
		Assay	By UV STP/FP/ 40001988	70%	to 130%	
67.	Diclofenac Sodium & Serratiopeptidase	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv	
	Tablets	Identification	By UV (Assay) STP/FP/ 40004115		parative litative)	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests an performed		ige of Testing / its of Detection
		Assay	By UV STP/FP/ 40004115	70%	to 130%
68.	Slow Diclofenac Tablets BP (Prolonged-release Diclofenac Tablets BF	General Tests for Tablets			general test 5. 1a-i, ii, iii
	Diclofenac Tablets BP)	Identification	By IR BP2013 STP/FP/40000650		nparative alitative)
		Dissolution	By UV BP2013 STP/FP/40000650	5%	to 100%
		Related substances	By HPLC BP2013 STP/FP/40000650	0.05	% to 1.0%
		Assay	By HPLC BP2013 STP/FP/40000650	70%	to 130%
69.	Diclofenac Sodium & Paracetamol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test 5. 1a-i, ii, iii, iv
		Identification	By UV STP/FP/ 40008484		nparative alitative)
		Uniformity of content for Diclofenac	By UV STP/FP/ 40008484	70%	to 130%
		Assay	By UV STP/FP/ 40008484	70%	to 130%

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70.	Diclofenac Sodium, Paracetamol & Chlorzoxazone	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
	Tablets	Identification	By UV (Assay) STP/FP/ 40008399		nparative alitative)		
		Uniformity of content for Diclofenac	By UV STP/FP/ 40008399	70%	to 130%		
		Assay	By UV (Assay) STP/FP/ 40008399	70%	to 130%		
71.	Dicyclomine & Mefenamic Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	As per Assay STP/FP/ 40003852		nparative alitative)		
		Dissolution for Mefenamic Acid	By HPLC STP/FP/ 40003852	D-25	5% to 100%		
		Assay	By Titration (For Dicyclomin By HPLC (For Mefenamic) STP/FP/ 40003852	ne) & 70%	to 130%		
72.	Dicyclomine & Paracetamol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	As per Assay STP/FP/ 40004134		nparative alitative)		
		Uniformity of content for Dicyclomine	By Titration STP/FP/ 40004134	70%	to 130%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection	
	Dicyclomine & Paracetamol Tablets	Assay	By Titration (For Dicyclomine) & By UV (For Paracetamol) STP/FP/ 40004134	70%	to 130%	
73.	Divalproex Sodium Delayed Release Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii	As general test S.No. 1a-i, ii, iii		
		Identification	By HPLC USP 35 NF30 STP/FP/ 40006015		parative litative)	
		Dissolution	By HPLC USP 35 NF30 STP/FP/ 40006015	D-25	5% to 100%	
		Assay	By HPLC USP 35 NF30 STP/FP/ 40006015	70%	to 130%	
74.	Aluminium, Magnesium & Simethicone Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	iii, iv As general test S.No. 1a-i, ii, iii, ir		
	Sinetificone Tablets	Identification	A) By IR (Simethicone)B) By Chemically(Aluminium & Magnesium)STP/FP/ 40000661		parative litative)	
		Assay	A) By IR (Simethicone)B) By Titration(Aluminium & Magnesium)STP/FP/ 40000661	70%	to 130%	

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75.	Drotaverine & Diclofenac Potassium Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 9. 1a-i, ii, iii, iv			
		Identification	By HPLC STP/FP/ 40009280		parative litative)			
		Assay	By HPLC STP/FP/ 40009280	70%	to 130%			
76.	Duloxetine Hydrochloride Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	-	eneral test 9. 1a-i, ii, iii, iv			
		Identification	By HPLC STP/FP/ 40002099		parative litative)			
		Assay	By HPLC STP/FP/ 40002099	70%	to 130%			
77.	Esomeprazole 40 mg (As enteric coated pellets) & Domperidone 30 mg (As sustained release pellets) Capsules	General Tests for Capsules	S.No. 1b-i, ii, iii		eneral test b. 1b-i, ii, iii			
		Identification	By HPLC STP/FP/ 40002060		parative litative)			
		Dissolution	By HPLC STP/FP/ 40002060	10%	to 100%			
		Assay	By HPLC STP/FP/ 40002060	70%	to 130%			
78.	Enalapril Maleate Tablets BP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 9. 1a-i, ii, iii, iv			
		Identification	By TLC By HPLC BP2013 STP/FP/40007002		parative litative)			

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	Enalapril Maleate Tablets BP	Related substances	By HPLC BP2013 STP/FP/40007002	0.05	% to 2.0%		
		Assay	By HPLC BP2013 STP/FP/40007002	70%	to 130%		
79.	Escitalopram & Clonazepam Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	-	As general test S.No. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/ 40007514		nparative alitative)		
		Uniformity of content	By HPLC STP/FP/ 40007514	70%	to 130%		
		Assay	By HPLC STP/FP/ 40007514	70%	to 130%		
80.	Escitalopram Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv		
		Identification	By HPLC IP-2010 STP/FP/ 40000675		nparative alitative)		
		Uniformity of content	By HPLC IP-2010 STP/FP/ 40000675	70%	to 130%		
		Assay	By HPLC IP-2010 STP/FP/ 40000675	70%	to 130%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		nge of Testing / its of Detection		
81.	Esomeprazole Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test o. 1a-i, ii, iii		
		Identification	By HPLC IP 2010, P-1296 STP/FP/40000821		nparative alitative)		
		Dissolution	By HPLC IP 2010, P-1296 STP/FP/40000821	D-2	5% to 100%		
		Assay	By HPLC IP 2010, P-1296 STP/FP/40000821	70%	o to 130%		
82.	Ferrous Ascorbate & Folic Acid Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv		
		Identification	As per Assay, Iron (By Chemically) STP/FP/ 40001885		nparative alitative)		
		Assay	By Titration (Ferrous Ascorbate & By UV (Folic Acid) STP/FP/ 40001885) 70%	o to 130%		
83.	Fexofenadine Hydrochloride Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii	As general test S.No. 1a-i, ii, iii			
		Identification	By IR By HPLC USP34 NF30 STP/FP/190000117		nparative alitative)		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ige of Testing / its of Detection	
	Fexofenadine Hydrochloride Tablets USP	Dissolution	By HPLC USP34 NF30 STP/FP/190000117	D-2:	5% to 100%	
		Related substances	By HPLC USP34 NF30 STP/FP/190000117	0.05	% to 2%	
		Assay	By HPLC USP34 NF30 STP/FP/190000117	70%	to 130%	
84.	Fluconazole Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	-	As general test S.No. 1a-i, ii, iii, iv	
		Identification	By HPLC IP-2010 STP/FP/ 40000907		nparative alitative)	
		Assay	By HPLC IP-2010 STP/FP/ 40000907	70%	to 130%	
85.	Fluoxetine Capsules IP	General Tests for Capsules	S.No. 1b-i, ii, iii		general test 5. 1b-i, ii, iii	
		Identification	By HPLC IP 2010 STP/FP/ 40008385		nparative alitative)	
		Dissolution	By HPLC IP 2010 STP/FP/ 40008385	D-25	5% to 100%	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificat against which tests are performed		nge of Testing / its of Detection			
	Fluoxetine Capsules IP	Related substances	By HPLC IP 2010 STP/FP/ 40008385	0.05	5% to 2.0%			
		Assay	By HPLC IP 2010 STP/FP/ 40008385	70%	o to 130%			
86.	Propranolol Hydrochloride SR & Flunarizine Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii	As general test S.No. 1a-i, ii, iii				
		Identification	By HPLC STP/FP/ 40000406		nparative alitative)			
		Dissolution	By HPLC STP/FP/ 40000406	10%	o to 100%			
		Uniformity of content	By HPLC STP/FP/ 40000406	70% to 130%				
		Related substances	By HPLC STP/FP/ 40000406	0.05	% to 5.0%			
		Assay	By HPLC STP/FP/ 40000406	70%	o to 130%			
87.	Flunarizine Dihydrochloride & Sustained release Propranolol hydrochloride Capsules	General Tests for Capsules	S.No. 1b-i, ii, iii		general test o. 1b-i, ii, iii			
		Identification	By HPLC STP/FP/ 40000406	Comparative (Qualitative)				
		Dissolution	By HPLC STP/FP/ 40000406	10%	o to 100%			

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / ts of Detection			
	Flunarizine Dihydrochloride & Sustained release	Uniformity of content	By HPLC STP/FP/ 40000406	70%	to 130%			
	Propranolol hydrochloride Capsules	Degradation product	By HPLC STP/FP/ 40000406	0.059	% to 5.0%			
	Capsuits	Assay	By HPLC STP/FP/ 40000406	70%	to 130%			
88.	Folic Acid Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test . 1a-i, ii, iii, iv			
		Identification	By UV USP 34 NF30 STP/FP/40005531		parative litative)			
		Dissolution	By HPLC USP 34 NF30 STP/FP/40005531	D-25	% to 100%			
		Related substances	By HPLC USP 34 NF30 STP/FP/40005531	0.059	% to 2.0%			
		Assay	By HPLC USP 34 NF30 STP/FP/40005531	70%	to 130%			
89.	Gatifloxacin Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 1a-i, ii, iii			
		Identification	By HPLC IP-2010 STP/FP/ 40000311		parative litative)			

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificati against which tests are performed		ge of Testing / its of Detection	
	Gatifloxacin Tablets IP	Dissolution	By HPLC IP-2010 STP/FP/ 40000311	D-25	5% to 100%	
		Related substances	By HPLC IP-2010 STP/FP/ 40000311	0.05	% to 6.0%	
		Assay	By HPLC IP-2010 STP/FP/ 40000311	70%	to 130%	
90.	Glimepiride & Metformin	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 9. 1a-i, ii, iii	
	Hydrochloride SR Tablets	Identification	As per Assay STP-AKUMS/STP/303		parative litative)	
		Dissolution	By HPLC (Glimepiride), By UV (Metformin) STP/FP/ 40001065	10%	to 100%	
		Uniformity of content for Glimepiride	By HPLC STP/FP/ 40001065	70%	to 130%	
		Assay	By HPLC (Glimepiride), By UV (Metformin) STP/FP/ 40001065	70%	to 130%	
91.	Glibenclamide & Cefixime Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 9. 1a-i, ii, iii, iv	
		Identification	By HPLC As per Assay STP/FP/40010392		parative litative)	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection
	Glibenclamide & Cefixime Tablets	Assay	By HPLC As per Assay STP/FP/40010392	70%	to 130%
92.	Ibuprofen Tablets BP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	-	general test 5. 1a-i, ii, iii, iv
		Identification	A) By IR B) By Melting point BP2013 STP/FP/40001021		nparative alitative)
		Related substances	By HPLC BP2013 STP/FP/40001021	0.05	% to 2.0%
		Assay	By HPLC BP2013 STP/FP/40001021	70%	to 130%
93.	Ibuprofen & Paracetamol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv
		Identification	As per Assay STP/FP/ 40001020		nparative alitative)
		Assay	By Titration (For Ibuprofen), B UV (Paracetamol) STP/FP/ 40001020	Sy 70%	to 130%
94.	Lactitol Monohydrate & Ispaghula husk Granules	General Tests for Tablets	S.No. 1a-i, ii		eneral test 5. 1a-i, ii

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / ts of Detection	
	Lactitol Monohydrate & Ispaghula husk Granules	Identification	By HPLC (As per Assay for Lactitol), By Swelling power (For Ispaghula husk) STP/FP/ 40001056		parative litative)	
		Assay	By HPLC STP/FP/ 40001056	70%	to 130%	
95.	Loratadine Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 1a-i, ii, iii	
		Identification	A) By TLC B) By HPLC USP 35 NF30 STP/FP/ 40004128		parative litative)	
		Dissolution	By UV USP 35 NF30 STP/FP/ 40004128	D-25	% to 100%	
		Related compounds	By HPLC USP 35 NF30 STP/FP/ 40004128	0.059	% to 1.0%	
		Assay	By HPLC USP 35 NF30 STP/FP/ 40004128	70%	to 130%	
96.	Loperamide Hydrochloride Tableta ID	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 1a-i, ii, iii	
	Tablets IP	Identification	A) By Chemically B) By HPLC IP 2010 STP/FP/ 40009659		parative litative)	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificat against which tests are performed		ge of Testing / its of Detection			
	Loperamide Hydrochloride Tablets IP	Dissolution	By HPLC IP 2010 STP/FP/ 40009659	D-25	% to 100%			
		Uniformity of content	By HPLC IP 2010 STP/FP/ 40009659	70%	to 130%			
		Assay	By HPLC IP 2010 STP/FP/ 40009659	70%	to 130%			
97.	Lisinopril Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii	-	eneral test 9. 1a-i, ii, iii			
		Identification	By HPLC USP 34 NF 30 STP/FP/40001175		parative litative)			
		Dissolution	By HPLC USP 35 NF 30 STP/FP/40001175	D-25	% to 100%			
		Related substances	By HPLC USP 35 NF 30 STP/FP/40001175	0.05	% to 2.0%			
		Uniformity of content	By HPLC USP 35 NF 30 STP/FP/40001175	70%	to 130%			
		Assay	By HPLC USP 35 NF 30 STP/FP/40001175	70%	to 130%			

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests are performed		nge of Testing / lits of Detection				
98.	Lisinopril & Hydrochlorthiazide Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test o. 1a-i, ii, iii				
	Tablets USP	Identification	By HPLC USP 35 NF 30 STP/FP/40008990		nparative alitative)				
		Dissolution	By HPLC USP 35 NF 30 STP/FP/40008990	D-2	5% to 100%				
		Uniformity of content	By HPLC USP 35 NF 30 STP/FP/40008990	70%	o to 130%				
		Assay	By HPLC USP 35 NF 30 STP/FP/40008990	70%	o to 130%				
99.	Levocetirizine & Montelukast Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv				
		Identification	By HPLC STP/FP/ 40007109		nparative alitative)				
		Uniformity of content	By HPLC STP/FP/ 40007109	70%	o to 130%				
		Assay	By HPLC STP/FP/ 40007109	70%	o to 130%				

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificatio against which tests are performed		ge of Testing / ts of Detection	
100.	Levocetirizine & Ambroxol Hydrochlorido	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test . 1a-i, ii, iii, iv	
Hydrochloride Tablets		Identification	By HPLC STP/FP/ 40001918		parative litative)	
		Uniformity of content for Levocetirizine	By HPLC STP/FP/ 40001918	70%	to 100%	
		Assay	By HPLC STP/FP/ 40001918	70%	to 130%	
101.	Levocetirizine Dihydrochloride	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test . 1a-i, ii, iii	
	Tablets IP	Identification	By HPLC IP 2010 STP/FP/40001179		parative litative)	
		Dissolution	By HPLC IP 2010 STP/FP/40001179	D-25	% to 100%	
		Uniformity of content	By HPLC IP 2010 STP/FP/40001179	70%	to 130%	
		Related substances	By HPLC IP 2010 STP/FP/40001179	0.059	% to 5.0%	
		Assay	By HPLC IP 2010 STP/FP/40001179	70%	to 130%	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificat against which tests are performed		nge of Testing / nits of Detection			
102.	Levofloxacin Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test o. 1a-i, ii, iii			
		Identification	By HPLC IP 2010 STP/FP/ 40001133		nparative alitative)			
		Dissolution	By UV IP 2010 STP/FP/ 40001133	D-2	25% to 100%			
		Related substances	By HPLC IP 2010 STP/FP/ 40001133	0.0	5% to 2.0%			
		Assay	By HPLC IP 2010 STP/FP/ 40001133	709	6 to 130%			
103.	Losartan Potassium & Hydrochlorothiazide	General Tests for Tablets	S.No. 1a-i, ii, iii		general test (o. 1a-i, ii, iii			
	Tablets USP	Identification	By HPLC USP 35 NF 30 STP/FP/40008990		nparative alitative)			
		Dissolution	By HPLC USP 35 NF 30 STP/FP/40008990	D-2	D-25% to 100%			
		Uniformity of content	By HPLC USP 35 NF 30 STP/FP/40008990	70%	6 to 130%			

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests an performed		nge of Testing / hits of Detection	
	Losartan Potassium & Hydrochlorothiazide Tablets USP	Assay	By HPLC USP 35 NF 30 STP/FP/40008990	70%	5 to 130%	
104.	Losartan Potassium Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv	
		Identification	A) By IR B) By UV C) By AAS USP 35 NF 30 STP/FP/40001228		nparative alitative)	
		Water	By KF USP 35 NF 30 STP/FP/40001228	>0.2	2%	
		Related substances	By HPLC USP 35 NF 30 STP/FP/40001228	0.05	5% to 4.0%	
		Assay	By HPLC USP 35 NF 30 STP/FP/40001228	70%	o to 130%	
105.	Metformin Hydrochloride Extended Release	General Tests for Tablets	S.No. 1a-i, ii, iii		general test o. 1a-i, ii, iii	
	Tablets IP	Identification	By UV (Assay) IP-2010 STP/FP/ 40009969		nparative alitative)	
		Dissolution	By UV STP/FP/ 40009969	10%	o to 100%	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection		
	Metformin Hydrochloride Extended Release Tablets IP	Assay	By UV STP/FP/ 40009969	70%	to 130%		
106.	Mefenamic Acid & Paracetamol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	As per Assay STP/FP/ 40005455		nparative alitative)		
		Assay	By UV (For Paracetamol), By Titration (For Mefenamic acid) STP/FP/ 40005455	70%	to 130%		
107.	Meloxicam Tablets BP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	A) TLC B) By HPLC BP 2013 STP/FP/ 40001904		nparative alitative)		
		Related substances	By TLC BP 2013 STP/FP/ 40001904		nparative alitative)		
		Assay	By UV BP 2013 STP/FP/ 40001904	70%	to 130%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection			
108.	Methylergometrine Tablets IP	General Tests for Tablets	S S.No. 1a-i, ii, iii, iv	As general test S.No. 1a-i, ii, iii, iv			
		Identification	 A) By TLC (Related substances) B) & C) By Chemically IP 2010 STP/FP/ 40001351 	Comparative (Qualitative)			
		Related substances	By TLC IP 2010 STP/FP/ 40001351	Comparative (Qualitative)			
		Uniformity of content	By UV IP 2010 STP/FP/ 40001351	70% to 130%			
		Assay	By UV IP 2010 STP/FP/ 40001351	70% to 130%			
109.	Metformin Hcl & Glimepiride Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	As general test S.No. 1a-i, ii, iii, iv			
		Identification	By HPLC STP/FP/40000951	Comparative (Qualitative)			
		Assay	By HPLC STP/FP/40000951	70% to 130%			
110.	Methylergonovine Maleate USP	General Tests for Tablets	S.No. 1a-i, ii, iii	As general test S.No. 1a-i, ii, iii			
		Identification	A) By IR B) By TLC USP 35 NF 30 STP/FP/40008958	Comparative (Qualitative)			

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Disc	ipline	Chemical Testing	Is	sue Date	04.09.2013		
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection		
	Methylergonovine Maleate USP	Dissolution	By HPLC USP 35 NF 30 STP/FP/40008958	D-25	5% to 100%		
		Related Alkaloids	By TLC USP 35 NF 30 STP/FP/40008958	Com	parative		
		Assay	By HPLC USP 35 NF 30 STP/FP/40008958	70%	to 130%		
111.	Moxifloxacin HCl Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	-	eneral test 9. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/40011254		parative litative)		
		Assay	By HPLC STP/FP/40011254	70%	to 130%		
112.	Montelukast Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 9. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/ 40001110		parative litative)		
		Uniformity of content	By HPLC STP/FP/ 40001110	70%	to 130%		
		Assay	By HPLC STP/FP/ 40001110	70%	to 130%		

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Disc	cipline	Chemical Testing		Issue Date	04.09.2013	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests an performed		nge of Testing / its of Detection	
113.	Nifedipine ER Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test o. 1a-i, ii, iii	
		Identification	By HPLC By UV USP35 NF30 STP/FP/40001521		nparative alitative)	
		Dissolution	By UV USP35 NF30 STP/FP/40001521	D-2	5% to 100%	
		Assay	By UV USP35 NF30 STP/FP/40001521	70%	o to 130%	
114.	Nitrofurantoin Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii		general test o. 1a-i, ii, iii	
		Identification	By IR By HPLC USP35 NF30 STP/FP/40002251		nparative alitative)	
		Dissolution	By HPLC USP35 NF30 STP/FP/40002251	D-2	5% to 100%	
		Related substances	By HPLC USP 34 NF 30 STP/FP/40002251	0.05	5% to 4.0%	
		Assay	By HPLC USP35 NF30 STP/FP/40002251	70%	o to 130%	

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Disc	ipline	Chemical Testing	I	Issue Date	04.09.2013	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificat against which tests are performed		ge of Testing / its of Detection	
115.	Nimesulide & Paracetamol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	-	eneral test 5. 1a-i, ii, iii, iv	
		Identification	As per Assay STP/FP/ 40001251		parative llitative)	
		Assay	By UV STP/FP/ 40001251	70%	to 130%	
116.	Nimesulide Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test b. 1a-i, ii, iii, iv	
		Identification	As per Assay STP/FP/ 40006615		parative llitative)	
		Assay	By UV STP/FP/ 40006615	70%	to 130%	
117.	Norfloxacin & Tinidazole Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv	
		Identification	As per Assay STP/FP/ 40011771		parative llitative)	
		Assay	By UV STP/FP/ 40011771	70%	to 130%	
118.	Ofloxacin Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 5. 1a-i, ii, iii	
		Identification	By HPLC IP 2010 STP/FP/ 40000825		parative llitative)	

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Disc	ipline	Chemical Testing		Issue Date	04.09.2013
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificat against which tests are performed		nge of Testing / its of Detection
		Dissolution	By UV IP 2010 STP/FP/ 40000825	D-2	5% to 100%
		Related Substances	By HPLC IP 2010 STP/FP/ 40000825	0.05	% to 4.0%
		Assay	By HPLC IP 2010 STP/FP/ 40000825	70%	o to 130%
119.	Ofloxacin and Ornidazole Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii		general test o. 1a-i, ii, iii
		Identification	As per Assay STP/FP/ 40000829		nparative alitative)
		Dissolution	By HPLC STP/FP/ 40000829	D-2	5% to 100%
		Assay	By HPLC STP/FP/ 40000829	70%	to 130%
120.	Omeprazole Capsules IP	General Tests for Capsules	S.No. 1b-i, ii, iii		general test o. 1b-i, ii, iii
		Identification	A) By UV B) By HPLC IP2010 STP/FP/ 40001635	(Qu	nparative alitative)
		Dissolution in acid & phosphate buffer medium	By HPLC IP2010 STP/FP/ 40001635	50%	d medium-1% to b, Buffer medium- to 150%

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Acc	reditation Standard	ISO/IEC 17025: 2005				
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificati against which tests are performed		ge of Testing / ts of Detection	
	Omeprazole Capsules IP	Uniformity of content	By HPLC IP2010 STP/FP/ 40001635	70%	to 130%	
	Omeprazole Magnesium & Domperidone Tablets	Loss on Drying	IP2010 STP/FP/ 40001635	>0.29	%	
		Assay	By HPLC IP2010 STP/FP/ 40001635	70%	to 130%	
121.		General Tests for Tablets	S.No. 1a-i, ii, iii, iv	-	eneral test . 1a-i, ii, iii, iv	
		Identification	By HPLC STP/FP/ 40001643		parative litative)	
		Uniformity of content	By HPLC STP/FP/ 40001643	70%	to 130%	
		Assay	By HPLC STP/FP/ 40001643	70%	to 130%	
122.	Omeprazole & Domperidone (SR) Capsules	General Tests for Tablets	S.No. 1b-i, ii, iii	-	eneral test . 1b-i, ii, iii	
	F	Identification			parative litative)	
		Dissolution	By HPLC STPFP/40001638	10% to 100%		
		Assay	By HPLC STPFP/40001638	70%	to 130%	
123.	Ofloxacin & Tinidazole Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test . 1a-i, ii, iii, iv	

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Ассі	reditation Standard	ISO/IEC 17025: 2005					
Disc	ipline	Chemical Testing		Issue Date	04.09.2013		
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests are performed		ge of Testing / its of Detection		
		Identification	As per Assay STP/FP/ 40000830		nparative alitative)		
		Assay	By UV STP/FP/ 40000830	70%	to 130%		
124.	Ondansetron Mouth Dissolving Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test 5. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/ 40000238		nparative alitative)		
		Uniformity of content	By HPLC STP/FP/ 40000238	70%	to 130%		
		Assay	By HPLC STP/FP/ 40000238	70%	to 130%		
125.	Pantoprazole & Domperidone Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/ 40003940		nparative alitative)		
		Uniformity of content for Domperidone	By HPLC STP/FP/ 40003940	70%	to 130%		
		Assay	By HPLC STP/FP/ 40003940	70%	to 130%		
126.	Paracetamol, Caffeine,	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	S.No	general test 5. 1a-i, ii, iii, iv		
	Chlorpheniramine maleare & Phenylephrine Hcl	Identification	As per Assay STP/FP/40000687		nparative alitative)		
	Phenylephrine Hcl Tablets	Assay	By HPLC STP/FP/40000687	70%	to 130%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests ar performed		nge of Testing / nits of Detection		
127.	Paracetamol, Caffeine & Chlorpheniramine maleate Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii		general test Io. 1a-i, ii, iii		
	maleate Tablets	Identification	As per Assay STP/FP/40009600		mparative ualitative)		
		Assay	By HPLC STP/FP/40009600	709	% to 130%		
128.	Pantoprazole Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test Io. 1a-i, ii, iii		
		Identification	A-By HPLC B-By UV IP 2010 STP/FP/ 40010183		mparative aalitative)		
		Dissolution	By HPLC & UV IP 2010 STP/FP/ 40010183	D-2	25% to 100%		
		Assay	By HPLC IP 2010 STP/FP/ 40010183	709	% to 130%		
129.	Piroxicam Capsules USP	General Tests for Capsules	S.No. 1a-i, ii, iii, iv		general test Io. 1a-i, ii, iii, iv		
		Identification	By TLC USP 35 NF 30 STP/FP/40001774		mparative aalitative)		
		Water	By KF USP 35 NF 30 STP/FP/40001774	>0.	2%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificatior against which tests are performed			
		Uniformity of Dosage units	By HPLC USP 35 NF 30 STP/FP/40001774	70%	to 130%	
		Dissolution	By UV USP 35 NF 30 STP/FP/40001774	10%	to 100%	
		Assay	By HPLC USP 35 NF 30 STP/FP/40001774	70%	to 130%	
130.	Prednisolone Tablets BP	General Tests for Tablets	S.No. 1a-i, ii, iii	-	general test p. 1a-i, ii, iii	
		Identification	A) By IR B) By TLC IP 2010 STP/FP/ 40001792		nparative alitative)	
		Related substances	By HPLC IP 2010 STP/FP/ 40001792	0.05	% to 6.0%	
		Uniformity of content	By UV IP 2010 STP/FP/ 40001792	70% to 130%		
		Dissolution	By UV IP 2010 STP/FP/ 40001792	D-2	5% to 100%	
		Assay	By HPLC IP 2010 STP/FP/ 40001792	70%	to 130%	
131.	Pseudoephedrine HCl & Triprolidine	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test 5. 1a-i, ii, iii, iv	
	Tablets	Identification	As per Assay STP/FP/40008986		nparative alitative)	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection		
		Assay	By HPLC STP/FP/40008986	70%	to 130%		
132.	Pregabalin Capsules	General Tests for Capsules	S.No. 1b-i, ii, iii, iv		eneral test b. 1b-i, ii, iii, iv		
		Identification	As per Assay STP/FP/40012252		parative llitative)		
		Assay	By HPLC STP/FP/40012252	70%	to 130%		
133.	Ranitidine Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	-	eneral test b. 1a-i, ii, iii, iv		
		Identification	 A) By TLC B) By HPLC C) By Chemically USP 35 NF30 STP/FP/40001881 		parative litative)		
		Dissolution	By UV USP35 NF30 STP/FP/40001881	D-25	5% to 100%		
		Related substances	By TLC USP35 NF30 STP/FP/40001881		parative llitative)		
		Assay	By HPLC USP35 NF30 STP/FP/40001881	70%	to 130%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
134.	Rabeprazole & Domperidone Capsules	General Tests for Capsules	S.No. 1b-i, ii, iii	As general test S.No. 1b-i, ii, iii
	[Rabeprazole Sodium 20 mg (As enteric coated), Domperidone	Identification	By UV STP/FP/ 40001832	Comparative (Qualitative)
	10 mg (As immediate release tablets) &	Dissolution	By UV STP/FP/ 40001832	10% to 100%
	Domperidone 20 mg (As sustained release tablet) Capsules]	Assay	By UV STP/FP/ 40001832	70% to 130%
135.	Rabeprazole Sodium & Domperidone Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	As general test S.No. 1a-i, ii, iii, iv
		Identification	By HPLC STP-AKUMS/STP/117	Comparative (Qualitative)
		Uniformity of content for Domperidone	By HPLC STP/FP/ 40001841	70% to 130%
		Assay	By HPLC STP/FP/ 40001841	70% to 130%
136.	Ranitidine Hydrochloride & Domnoridano Toklata	General Tests for Tablets	S.No. 1a-i, ii, iii, iv	As general test S.No. 1a-i, ii, iii, iv
	Domperidone Tablets	Identification	As per Assay STP/FP/ 40007558	Comparative (Qualitative)
		Uniformity of content for Domperidone	By HPLC STP/FP/ 40007558	70% to 130%
		Assay	By HPLC STP/FP/ 40007558	70% to 130%

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection		
137.	Roxithromycin Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 5. 1a-i, ii, iii		
		Identification	By HPLC IP 2010 STP/FP/ 40001946		parative llitative)		
		Dissolution	By HPLC IP 2010 STP/FP/ 40001946	D-25	5% to 100%		
		Assay	By HPLC IP 2010 STP/FP/ 40001946	70%	to 130%		
138.	Rosuvastatin Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 5. 1a-i, ii, iii		
		Identification	By HPLC IP 2010 STP/FP/ 40005472		parative llitative)		
		Dissolution	By HPLC IP 2010 STP/FP/ 40005472	D-25	5% to 100%		
		Uniformity of content	By HPLC IP 2010 STP/FP/ 40005472	70%	to 130%		
		Related substances	By HPLC IP 2010 STP/FP/ 40005472	0.05	% to 6.0%		
		Assay	By HPLC IP 2010 STP/FP/ 40005472	70%	to 130%		
139.	Simvastatin Tablets USP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 5. 1a-i, ii, iii		

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Acc	reditation Standard	ISO/IEC 17025: 2005					
Disc	cipline	Chemical Testing		Issue Date	04.09.2013		
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests ar performed		nge of Testing / nits of Detection		
		Identification	By HPLC USP 32 STP/FP/ 40002027		mparative nalitative)		
		Dissolution	By UV USP 32 STP/FP/ 40002027	D-2	25% to 100%		
		Assay	By HPLC USP 32 STP/FP/ 40002027	709	6 to 130%		
140.	Sildenafil Citrate Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test Io. 1a-i, ii, iii		
		Identification	By HPLC IP 2010 STP/FP/ 4000078		mparative alitative)		
		Dissolution	By UV IP 2010 STP/FP/ 4000078		25% to 100%		
		Related Substances	By HPLC IP 2010 STP/FP/ 4000078		% to 6.0%		
		Assay	By HPLC IP 2010 STP/FP/ 4000078		6 to 130%		
141.	Sodium Valproate & Valproic Acid	General Tests for Tablets	S.No. 1a-i, ii, iii	S.No. 1a-i, ii, iii As genera S.No. 1a-i			
	controlled release Tablets	Identification	A) By HPLC B) By test of Sodium STP/FP/ 40002266		mparative aalitative)		

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Disc	cipline	Chemical Testing	lss	sue Date	04.09.2013	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection	
		Dissolution	By HPLC STP/FP/ 40002266	D-25	% to 100%	
		Assay	By HPLC (For Valproate Ions) By Titration (For Sodium Valproate STP/FP/ 40002266) 70%	to 130%	
142.	S (-) Amlodipine Besilate & Atenolol Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii	-	eneral test 9. 1a-i, ii, iii	
		Identification	By HPLC STP/FP/ 40005336		parative litative)	
		Uniformity of content	By HPLC STP/FP/ 40005336	70%	to 130%	
		Dissolution	By HPLC STP/FP/ 40005336	D-25	%% to 130%	
		Assay	By HPLC STP/FP/ 40005336	70%	to 130%	
143.	S(-) Amlodipine Besilate Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 9. 1a-i, ii, iii	
		Identification	By HPLC IP 2010 STP/FP/ 40002030	Comparative (Qualitative)		
		Uniformity of content	By HPLC IP 2010 STP/FP/ 40002030	70%	to 130%	
		Dissolution	By -UV IP 2010 STP/FP/ 40002030	D-25	5% to 100%	

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Acci	reditation Standard	ISO/IEC 17025: 2005				
Disc	ipline	Chemical Testing	lss	sue Date	04.09.2013	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / ts of Detection	
		Assay	By HPLC IP 2010 STP/FP/ 40002030	70%	to 130%	
144.	Serratiopeptidase Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 1a-i, ii, iii, iv	
		Identification	By UV (Assay) IP 2010 STP/FP/ 40001244		parative litative)	
		Assay	By UV IP 2010 STP/FP/ 40001244	70%	to 130%	
145.	Sodium Feredetate, Folic acid & Zinc sulphate Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 1a-i, ii, iii, iv	
	sulphate Tablets	Identification	As per Assay STP/FP/40005446		parative litative)	
		Assay	By HPLC STP/FP/40005446	70%	to 130%	
146.	Tramadol Hydrochloride & Barrassármal Tablata	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 1a-i, ii, iii, iv	
	Paracetamol Tablets	Identification	By HPLC STP/FP/ 40011830		parative litative)	
		Uniformity of content for Tramadol Hydrochloride	By HPLC STP/FP/ 40011830	70%	to 130%	
		Assay	By HPLC STP/FP/ 40011830	70%	to 130%	

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Acci	reditation Standard	ISO/IEC 17025: 2005	ISO/IEC 17025: 2005				
Disc	ipline	Chemical Testing	Iss	ue Date	04.09.2013		
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection		
147.	Tramadol Hydrochloride	General Tests for Tablets	S.No. 1a-i, ii, iii		eneral test 5. 1a-i, ii, iii		
	Sustained Release Tablets	Identification	By UV (Assay) STP/FP/40005779		parative llitative)		
		Dissolution at different stages	By UV STP/FP/40005779	10%	to 100%		
		Assay	By UV STP/FP/40005779	70%	to 130%		
148.	Tadalafil Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		eneral test 5. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/ 40011212		parative litative)		
		Assay	By HPLC STP/FP/ 40011212	70% to 130%			
149.	Telmisartan Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii	As general test S.No. 1a-i, ii, i			
		Identification	By HPLC IP 2010 STP/FP/ 40005882		parative llitative)		
		Dissolution	By HPLC IP 2010 STP/FP/ 40005882	D-25	5% to 100%		
		Related substances	By HPLC IP 2010 STP/FP/ 40005882	0.5 %	6 to 6.0%		
		Assay	By HPLC IP 2010 STP/FP/ 40005882	70%	to 130%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests are performed		nge of Testing / nits of Detection	
150.	Telmisartan & Hydrochlorothiazide Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test o. 1a-i, ii, iii, iv	
		Identification	By HPLC STP/FP/ 40000353		nparative alitative)	
		Assay	By HPLC STP/FP/ 40000353	70%	6 to 130%	
151.	Terbinafine Dispersible Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv, v		general test o. 1a-i, ii, iii, iv, v	
		Identification	By IR STP/FP/ 40008670		nparative alitative)	
		Dissolution	By UV STP/FP/ 40008670	D-2	25% to 100%	
		Assay	By Titration STP/FP/ 40008670	70%	6 to 130%	
152.	Tinidazole Tablets IP	General Tests for Tablets	S.No. 1a-i, ii, iii		general test 'o. 1a-i, ii, iii	
		Identification	 A) By UV (Assay) B) By Chemically C) By Melting range IP 2010 STP/FP/ 40001012 		nparative alitative)	
		Dissolution	By UV IP 2010 STP/FP/ 40001012	D-2	25% to 100%	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection		
	Tinidazole Tablets IP	Assay	By UV IP 2010 STP/FP/ 40001012	70%	to 130%		
153.	Tamsulosin HCl Capsules BP	General Tests for Capsules	S.No. 1b-i, ii, iii, iv	-	general test 5. 1b-i, ii, iii, iv		
		Identification	 A) By UV B) By HPLC BP 2013 STP/FP/40009730 		nparative alitative)		
		Uniformity of content for	By HPLC BP 2013 STP/FP/40009730	70%	to 130%		
		Related substances	By HPLC BP 2013 STP/FP/40009730	0.5 0	% to 6.0%		
		Assay	By HPLC BP 2013 STP/FP/40009730	70%	to 130%		
154.	Voglibose Tablets	General Tests for Tablets	S.No. 1a-i, ii, iii, iv		general test p. 1a-i, ii, iii, iv		
		Identification	By HPLC STP/FP/ 40001094		nparative alitative)		
		Uniformity of content	By HPLC STP/FP/ 40001094	70%	to 130%		
		Assay	By HPLC STP/FP/ 40001094	70%	to 130%		

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S.No.	Product /	Specific Test Performed	Test Method Specification	Range of Testing /
	Material of Test		against which tests are	Limits of Detection
			performed	

3. General Tests of Chemicals & Pharmaceutical Ingredients

A-Chemicals & Pharmaceutical Ingredients	i. Acid Value	IP 2010 As per respective STPs	1 – 100
	ii. Acetyl value	IP 2010 As per respective STPs	NLT 100
	iii. Acidity or Alkalinity	IP 2010 BP 2013 USP35 NF30 As per respective STPs	Comparative (Qualitative)
	iv. Aluminium	IP 2010 As per respective STPs	Comparative (Qualitative)
	v.Aluminium and calcium	IP 2010 As per respective STPs	Comparative (Qualitative)
	vi. Ammonium	IP 2010 As per respective STPs	Comparative (Qualitative)
	vii. Angle of repose	IP 2010 As per respective STPs	0° - 90°
	viii. Arsenic	IP 2010 As per respective STPs	Comparative (Qualitative)
	ix. Barium	IP 2010 As per respective STPs	Comparative (Qualitative)
	x. Bromide	IP 2010 As per respective STPs	Comparative (Qualitative)

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specific against which tests a performed		Range of Testing / Limits of Detection		
		xi. Melting Range/ Melting Point	IP 2010 As per respective STPs		30°C to 300°C		
		xii. Magnesium	IP 2010 As per respective STPs		Comparative (Qualitative)		
		xiii.Nitrogen Content	IP 2010 As per respective STPs		More than 2 mg		
		xiv.Nitrate	IP 2010 As per respective STPs		Comparative (Qualitative)		
		xv.Chloride	IP 2010 As per respective STPs		Comparative (Qualitative)		
		xvi.Calcium	IP 2010 As per respective STPs		Comparative (Qualitative)		
		xvii.Clarity And Color of Solution/ Appearance of solution	IP 2010 As per respective STPs		Comparative (Qualitative)		
		xviii. Optical Rotation and Specific Rotation	IP 2010 As per respective STPs		Up to 360°		
		xix. Peroxide Value	IP 2010 As per respective STPs		1 - 200		
		xx. pH	As per respective STPs		1 to 14		
		xxi.Residue on Ignition	IP 2010 As per respective STPs		More than 0.01%		
		xxii.Defoaming Activity	IP 2010 As per respective STPs		Comparative (Qualitative)		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specific against which tests a performed		nge of Testing / its of Detection		
		xxiii.Description	As per respective STPs	Ası	per specification		
		xxiv.Ester Value	IP 2010 As per respective STPs	1 –	100		
		xxv.Solubility	IP 2010 BP 2013 USP35 NF30 As per respective STPs		nparative alitative)		
		xxvi.Sulphate	IP 2010 As per respective STPs		nparative alitative)		
		xxvii.Sulphated Ash/ Residue On Ignition	IP 2010 As per respective STPs	Mor	re than 0.01%		
		xxviii.Sodium	IP 2010 As per respective STPs		nparative alitative)		
		xxix.Total Dye Content	As per respective STPs		nparative alitative)		
		xxx.Hydroxyl Value	IP 2010 As per respective STPs	1 –	1000		
		xxxi.Heavy Metals	IP 2010 As per respective STPs		nparative alitative)		

IP 2010

IP 2010

As per respective STPs

As per respective STPs

xxxii.Halogens

xxxiii.Iron

Comparative (Qualitative)

Comparative (Qualitative)

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	editation Standard	ISO/IEC 17025: 2005			- / / -
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests ar performed		nge of Testing / its of Detection
		xxxiv.Iodine Value	IP 2010 As per respective STPs	1 - 2	200
		xxxv.Identification A) By Chemical	IP 2010 BP 2013 USP35 NF30		nparative alitative)
		B) By UV	As per respective STPs IP 2010 BP 2013 USP35 NF30		nparative alitative)
		C) By HPLC	As per respective STPs IP 2010 BP 2013 USP35 NF30		nparative alitative)
		D) By IR	As per respective STPs IP 2010 BP 2013 USP35 NF30 As per respective STPs		nparative alitative)
		xxxvi.Loss On Drying	IP 2010 As per respective STPs	Mor	re than 0.01%
		xxxvii.Loss On Ignition	IP 2010 As per respective STPs	Mor	re than 0.01%
		xxxviii.Lead	IP 2010 As per respective STPs		nparative alitative)
		xxxix.Water by KF	IP 2010 As per respective STPs	0.19	6 - 100%

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection

4. Drug & Pharmaceutical Ingradients Having Specific Tests

1.	Activated Dimethicone IP	Assay	IP 2010 STP NoSTP/RM/10000011	80% to 120%
		General Test	S. No.3 iii, xxii, xxiii, xxxi, xxxv	As per specification
2.	Aceclofenac IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000046	80% to 120%
		Related substances (By HPLC)	IP 2010 Single Impurity: < 0.5% Total Impurities: < 2.0%	0.05% to 4.0%
		General Test	S.No.3-xxiii, xxxi, xxxv, xxvii, xxxvi	As per specification
3.	Albendazole IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000012	80% to 120%
		Related substances (By TLC)	IP 2010	Comparative (Qualitative)
		General Test	S.No.3-xxiii, xxxi, xxxv, xxvii, xxxvi	As per specification
4.	Alprazolam IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000013	80% to 120%
		Related substances (By TLC)	IP 2010	Comparative (Qualitative)
		General Test	S.No. 3-xxiii, xxx23, 31, 35, 27, 36	As per specification

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection
5.	Ambroxol HCl IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000010	80%	to 120%
		Related substances (By HPLC)	IP 2010 Single Impurity: < 0.5% Total Impurities: < 1.0%	0.059	% to 2.0%
		General Test	S.No.3-xxiii, xxxv, xxv, xxvi, xxxi, xxix, xi	As p	er specification
6.	Amitriptyline HCl IP	Assay (By Titration)	IP 2010 STP No STP/RM/10005394	80%	to 120%
		Related substances (By TLC)	Comparative (Qualitative)	As p	er specification
		General Test	S.No.3-xxiii, xxxv, xxv, xxvi, xvii, xxxi, xxvii, xxxvi	As p	er specification
7.	Amlodipine Besilate IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000028	80%	to 120%
		Related substances (By TLC & HPLC)	IP 2010 For TLC: Single Impurity: < 0.3% Total Impurities: < 0.1%		FLC: parative litative)
			For HPLC: Single Impurity: < 0.3% Total Impurities: < 0.3%		HPLC: % to 2.0%
		General Test	S.No.3-xxiii, xxxv, xxv, xxvi, xvii, xxxi, xxvii, xxxvi	As p	er specification
8.	Aspirin IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000024	80%	to 120%

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Salicylic acid	Comparative (Qualitative)	As per specification
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, viii, xxxi, xxvii, xxxvi	As per specification
9.	Ascorbic Acid IP (Coated)	Assay (By Titration)	IP 2010 STP No STP/RM/10004626	80% to 120%
		Light absorption	About 0.56	As per specification
		Oxalic Acid	Comparative (Qualitative)	As per specification
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xviii, xxxi, xxvii	As per specification
10.	Atenolol IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000039	80% to 120%
		Related substances (By HPLC)	Single Impurity: 0.25% Total Impurities: 0.5%	0.05% to 1.0%
		General Test	S.No.3-xxiii, xxxv, xvii, xv, xxxvi	As per specification
11.	Atorvastatin Calcium IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000051	80% to 120%
		Related substances (By HPLC)	Single Impurity: 0.5% Total Impurities: 2.0%	0.05% to 4.0%
		General Test	S.No.3-xxiii, xxxv, xxv, xviii, xxxi	As per specification
12.	Azithromycin IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000047	80% to 120%

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection	
		Related substances (By HPLC)	Impurity B: 2.0% Single Impurity: 1.0% Total Impurities: 5.0%	0.05	% to 10.0%	
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xviii, xxxi, xxvii, xxxix	As p	er specification	
13.	Amoxicillin Trihydrate IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000006	80%	80% to 120%	
		N,N-Dimethylaniline	NMT 20 ppm (Comparative (Qualitative))	As p	er specification	
		General Test	S.No.3- xxiii, xxxv, xvii, xx, xviii, xxxi, xxvii, xxxix	As p	er specification	
14.	Ampicillin Trihydrate IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000008	80%	to 120%	
		N,N-Dimethylaniline	NMT 20 ppm (Comparative (Qualitative))	As p	er specification	
		General Test	S.No. 3- xxiii, xxxv, xxv, xvii, xx xviii, xxxi, xxvii, xxxix	, As p	er specification	
15.	Bromhexine hydrochloride IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000085	80%	to 120%	
		Related Substances (By TLC)	Comparative (Qualitative)	As p	er Specification	
		General Test	S.No.3-xxiii, xxxv, xxv, xxvii, xxxvi	As p	er specification	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection			
16.	Betamethasone sodium Phosphate IP	Assay (By UV)	IP 2010 STP No STP/RM/10000086	80% to 120%			
		Light absorption	1.7 to 1.9	As per specification			
		Inorganic phosphate (By UV)	NMT 0.5%	0.25% to 1.0%			
		Free Betamethasone and other derivatives (By TLC)	Comparative (Qualitative)	As per specification			
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xviii, xxxix	As per specification			
17.	Biotin USP	Assay (By Titration)	USP35 NF30 STP No STP/RM/10000108	80% to 120%			
		General Test	S.No.3- xxiii, xxxv, xxv, xviii	As per specification			
18.	Bisacodyl IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000088	80% to 120%			
		General Test	S.No.3-xxiii, xxxv, xxv, iii, xxvii, xxxvi	As per specification			
19.	Betacarotene 30%	Assay (By UV)	STP No STP/RM/10000091	80% to 120%			
		Related substances (By UV)	NLT 1.5 (Comparative (Qualitative))	As per specification			
		General Test	S.No.3-xxiii, xxxv, xxv, xxxi, xxxvi, xxvii	As per specification			

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
20.	Benfothiamine	Assay (By Titration)	BP 2013 STP No STP/RM/10000090	80% to 120%
		General Test	S.No.3-xxiii, xxxv, xxv	As per specification
21.	Bisoprolol Fumerate USP	Assay (By HPLC)	USP35 NF30 STP No STP/RM/10000090	80% to 120%
		Chromatographic purity	Total impurities: < 0.5%	0.05% to 1.0%
		Content of fumaric acid	14.8% to 15.4%	As per specification
		General Test	S.No.3-xxiii, xxxv, xxv, xviii, xxxix, xxi, xxxi	As per specification
22.	Carbamazepine IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000201	80% to 120%
		Related Substances (By HPLC)	Known impurity: < 0.1% Unknown impurity: < 0.1% Total impurities: < 0.5%	0.05% to 1.0%
		General Test	S.No.3-xxiii, xxxv, xxv, iii, xxxi, xv, xxvii, xxxvi	As per specification
23.	Ciprofloxacin HCl IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000119	80% to 120%
		Related substances (By HPLC)	Individual impurity: < 0.2% Total impurities: < 0.5%	0.05% to 1.0%
		Fluoroquinolonic acid (By TLC)	Comparative (Qualitative)	As per specification

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		General Test	S.No.3-xxiii, xxxv, xxv, xx, xxxi, xxvi, xxvii, xxxix	As per specification
24.	Citicoline Sodium	Assay	STP No. STP/RM/10000120	80% to 120%
		General Test	S.No.3-xxiii, xxxv, xxv	As per specification
25.	Clonazepam IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000206	80% to 120%
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 1.0%
		General Test	S.No.3-xxiii, xxxv, xxv, xxvii, xxxvi	As per specification
26.	Clopidogrel Bisulfate IP	Assay (By HPLC)	IP2010 STP No. STP/RM/10000200	80% to 120%
		Related substances (By HPLC)	Impurity A: $< 0.2\%$ Impurity B: $< 0.3\%$ Impurity C: $< 1.0\%$ Other impurities: $< 0.1\%$ Total impurities: $< 1.5\%$	0.05% to 3.0%
		General Test	S.No. 3- xxiii, xxxv, xxv, xxvii, xxxvi	As per specification
27.	Calcium D Pantothenate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000114	80% to 120%
		β-Alanine (By TLC)	Comparative (Qualitative)	As per specification
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xviii, xxxi, xxxvi	As per specification

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
28.	Chloramphenicol IP	Assay (By UV)	IP 2010 STP No. STP/RM/10000115	80% to 120%		
		Related substances	Comparative (Qualitative)	As per specification		
		(By TLC) General Test	S.No.3-xxiii, xxxv, xxv, xx, xviii, xv, xxvii, xxxvi	As per specification		
29.	Caffeine IP (Anhydrous)	Assay (By Titration)	IP 2010 STP No STP/RM/10000116	80% to 120%		
		Related substances (By TLC)	Comparative (Qualitative)	As per specification		
		General Test	S.No.3-xxii, xxxv, xxv, xvii, iii, viii, xxxi, xxvii, xxxvi	As per specification		
30.	Cinnarizine IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000117	80% to 120%		
		Related substances (By TLC)	Comparative (Qualitative)	As per specification		
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, iii, xxxi, xxvii, xxxvi	As per specification		
31.	Chlorpheniramine Maleate IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000118	80% to 120%		
		Related substances (By TLC)	Comparative (Qualitative)	As per specification		
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xxvii, xxxvi	As per specification		

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32.	Cetirizine HCl IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000125	80%	to 120%		
		Related substances (By HPLC)	Single impurity: < 0.1% Total impurities: < 1.5%	0.059	% to 3.0%		
		General Test	S.No.3- xxiii, xxxv, xxv, xvii, xxvii, xxxvi	xx, As po	er specification		
33.	Chlordiazepoxide IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000134	80%	to 120%		
		Related substances (By TLC)	Comparative (Qualitative)	As pe	er specification		
		General Test	S.No.3-xxiii, xxxv, xxv, xxxi, xxvii, xxxvi	As pe	er specification		
34.	Chloroquine Phosphate IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000142	80%	to 120%		
		Related substances (By TLC)	Comparative (Qualitative)	As pe	er specification		
		General Test	S.No.3- xxiii, xxxv, xxv, xx, x xxxix	xxi As pe	er specification		
35.	Clidinium Bromide USP	Assay (By Titration)	USP35 NF30 STP No STP/RM/10000144	80%	to 120%		
		Related compounds (By TLC)	NMT 0.5% of clidinium bromi related compound A (Comparative (Qualitative))	ide As po	er specification		

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		General Test	S.No.3-xxiii, xxxv, xxv, xxxvi, xxi, xxxi	As per specification			
36.	Chlorzoxazone USP	Assay (By UV)	USP35 NF30 STP No STP/RM/10000148	80% to 120%			
		Chromatographic purity (By TLC)	Comparative (Qualitative)	As per specification			
		Chlorine Content	20.6% and 21.2%	As per specification			
		General Test	S.No.3- xxiii, xxxv, xxv, xi, xxxvi, xxxi, xxi	As per specification			
37.	Calcium Carbonate IP (Oyster Shell)	Assay (By Titration)	IP 2010 STP No STP/RM/10000152	80% to 120%			
		Substances insoluble in acetic acid	Comparative (Qualitative) (< 0.2%)	As per specification			
		General Test	S.No.3- xxiii, xxxv, xxv, viii,xxxi, ix,xxxiii, xv, xxvi, xxxvi	As per specification			
38.	Cyproheptadine Hydrochloride IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000153	80% to 120%			
		Related substances (By TLC)	Comparative (Qualitative)	As per specification			
		General Test	S.No.3- xxiii, xxxv, xxv, xxvii, xxxvi	As per specification			
39.	Calcium Citrate Maleate	Assay (By Titration)	IP 2010 STP No STP/RM/10000194	80% to 120% of LC			

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		General Test	S.No.3- xxiii, xxxv, xxv	As per specification
40.	Calcitriol IP	Assay (By HPLC)	IP 2010 STP NoSTP/RM/10000131	80% to 120%
		Related substances (By HPLC)	Single impurity: < 0.5 Total impurities: < 1.0%	0.05% to 2.0%
		General Test	S.No.3-xxiii, xxxv, xxv	As per specification
41.	Copper Sulphate Pentahydrate BP	Assay (By Titration)	BP 2013 STP No STP/RM/10000203	80% to 120%
		General Test	S.No.3- xxiii, xxxv, xxv, xvii, xv, xxxiii, xxxviii, xxxvi	As per specification
42.	Clotrimazole IP	Assay (By Titration)	IP 2010 STP NoSTP/RM/10000203	80% to 120%
		2-Chlorotritanol	Comparative (Qualitative)	As per specification
		Imidazole (By TLC)	Comparative (Qualitative)	As per specification
		General Test	S.No.3- xxiii, xxxv, xxv, xvii, xxvii, xxxvi	As per specification
43.	Captopril IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000231	80% to 120%
		General Test	S.No.3- xxiii, xxxv, xxv, xviii, xxxi, xxxvi, xxvii	As per specification
44.	Clarithromycin IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000227	80% to 120%

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		Related substances (By HPLC)	Single impurity: < 2.5% Total impurities: < 5.0%	0.05	% to 10.0%	
		General Test	S.No.3- xxiii, xxxv, xxv, xviii, xxxi, xxxix	As p	per specification	
45.	Calcium Ascorbate USP	Assay (By Titration)	USP35 NF30 STP No STP/RM/10000220	80%	to 120%	
		Limit of fluoride	NMT 10 ppm (Comparative (Qualitative))	As p	per specification	
		General Test	S.No.3- xxiii, xxxv, xxv, xviii, xx xxxvi, viii, xxxi	, As p	per specification	
46.	Cefadroxil IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000112	80%	to 120%	
		N,N-Dimethylaniline	Comparative (Qualitative) (NMT 20ppm)	As p	per specification	
		Related substances (By TLC)	Comparative (Qualitative)	As p	per specification	
		General Test	S.No.3- xxiii, xxxv, xxv, xx, xviii xxxix	, As p	per specification	
47.	Cefdinir USP	Assay (By HPLC)	USP35 NF30 STP No STP/RM/10000193	80%	to 120%	
		Related substances	By HPLC (As per specification)	0.05	% to 6.0%	
		General Test	S.No.3- xxiii, xxxv, xxv, xviii, xxxix, xxi, xxxi	As p	per specification	

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48.	Cefixime IP	Assay (By HPLC)	IP2010 STP No STP/RM/10000132	80%	to 120%			
		Related substances (By HPLC)	Single impurity: <0.5% Total impurities: < 3.0%	0.05	% to 6.0%			
		General Test	S.No.3- xxiii, xxxv, xxv, xx, xxxix, xxvii	As p	er specification			
49.	Cefpodoxime Proxetil IP	Assay (By HPLC)	IP2010 STP No STP/RM/10000130	80%	to 120%			
		Related substances (By HPLC)	Any impurity at RRT 0.86: <3.0%; Any impurity at RRT 1. & 1.39: < 1.0%; Other individu impurity at RRT 2.0: <0.5%; Total impurity: 6.0%	.27	% to 6.0%			
		Isomer ratio (By HPLC)	Comparative (Qualitative)	As p	er specification			
		General Test	S.No.3- xxiii, xxxv, xxv, xviii, xxxi, xxvii, xxxix	As p	er specification			
50.	Cefuroxime Axetil IP	Assay (By HPLC)	IP2010 STP No STP/RM/10000196	80%	to 120%			
		Diastereoisomer ratio (By HPLC)	Comparative (Qualitative) (Between 0.48 and 0.55)	As p	er Specification			
		Related substances (By HPLC)	E-isomer: $< 1.0\%$ D ³ -isomer: $< 1.5\%$ Single impurity: $< 0.5\%$ Total impurities: $< 3.0\%$	0.05	% to 6.0%			

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		Acetone	Comparative (Qualitative)	As per specification
		General Test	S.No.3- xxiii, xxxv, xxv, xxxix	As per specification
51.	Cephalexin IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000109	80% to 120%
		Light absorption	0.44 to 0.49	As per specification
		Related substances	D-phenylglycine: <1.0% Total impurities: <3.0%	0.05 to 6.0%
		General Test	S.No.2- xxiii, xxxv, xxv, xx, xviii, xxvii, xxxix	As per specification
52.	Cloxacillin Sodium IP	Assay (By HPLC)	IP 2010 STP No STP/RM/10000110	80% to 120%
		N,N-Dimethylaniline	Comparative (Qualitative) (NMT 20ppm)	As per specification
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xviii	As per specification
53.	Cyanocobalamin IP	Assay (By UV)	IP 2010 STP No STP/RM/10000873	80% to 120%
		Related substances (By HPLC)	Single impurity: < 0.2% Total impurities: < 0.5%	0.05% to 1.0%
		General Test	S.No.3- xxiii, xxxv, xxv, xxxvi	As per specification
54.	Dicloxacillin Sodium USP	Assay (By HPLC)	USP35 NF30 STP No STP/RM/10001988	80% to 120%

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		Crystallinity	Comparative (Qualitative)	As per specification
		Dimethylaniline	Comparative (Qualitative)	As per specification
		General Test	S.No.2-xxiii, xxxv, xxv, xx, xxxix	As per specification
55.	Diazepam IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000238	80% to 120%
		Related substances & decomposition products (By TLC)	Comparative (Qualitative)	As per specification
		General Test	S.No.3-xxiii, xxxv, xxv, xxxi, xxvii, xxvi	As per specification
56.	Diclofenac Sodium IP	Assay (By Titration)	IP 2010 STP No STP/RM/10000236	80% to 120%
		Light absorption	NMT 0.050	As per specification
		Related substances (By HPLC)	Single impurity: < 0.2% Total impurities: < 0.5%	0.05% to 1.0%
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xxxi, xxxvi	As per specification
57.	Diclofenac Potassium BP	Assay (By Titration)	BP 2013 STP No STP/RM/10000279	80% to 120%
		Related substances (By HPLC)	Single impurity: < 0.2% Total impurities: < 0.5%	0.05% to 1.0%
		General Test	S.No.3-xxiii, xxxv, xxv, xvii, xxxi, xxxvi	As per specification

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	Range of Testing / Limits of Detection			
2010 P No. STP/RM/10000269	80% to 120%			
sacetyl diltiazem HCl: <0.5%; lividual impurity: 0.5%; Total purities: 1.0%	0.05% to 2.0%			
Jo.3-xxiii, xxxv, xxv, xviii, ci, xxxvi	As per specification			
2013 P No. STP/RM/10000233	80% to 120%			
purity A, B, C, D, E, F: <0.25%; tal impurities: <0.5%	0.05% to 1.0%			
ło.3-xxiii, xxxv, xxv, xvii, ki, xxxvi, xxvii	As per specification			
2010 P No. STP/RM/10000253	80% to 120%			
gle impurity: <0.25% tal impurities: <0.5%	0.05% to 1.0%			
Jo.3-xxiii, xxxv, xxv, xvii, ki, xxvii, xxxvi	As per specification			
P No. STP/RM/10000277	80% to 120%			
Jo.3-xxiii, xxxv, xxv	As per specification			
2010 P No. STP/RM/10000232	80% to 120%			
	Valid Ur Page St Method Specification anst which tests are formed 2010 2010 2010 2010 2010 2010 2010 2010 2013 2013 2013 2013 2013 2013 2013 2013 2013 2013 2015 2013 2015 2017 2017 2018 2018 2019 2010 20			

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		Related substances (By TLC)	Comparative (Qualitative)	As p	er specification	
		General Test	S.No.3-xiii, xxxv, xxv, xxvi xxxvi	i, As p	er specification	
63.	Dextromethorphan Hydrobromide IP	Assay (By Titration)	IP 2010	80%	to 120%	
		N,N-Dimethylaniline	STP No. STP/RM/10000234 Comparative (Qualitative) (NMT 20ppm)		er specification	
		General Test	S.No.3- xxiii, xxxv, xxv, xvi xxvii, xxxix	ii, iii, As p	er specification	
67.	Dexamethasone IP	Assay (By UV)	IP 2010 STP No. STP/RM/10000237		to 120%	
		Light absorption Related substances (By HPLC)	0.38 to 0.41 Single impurity: <0.5% Total impurities: <1.0%		er specification % to 2.0%	
		General Test	S.No.3- xxiii, xxxv, xxv, xvi xxvii, xxxvi	iii, As p	er specification	
68.	Diphenhydramine HCl IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000239		to 120%	
		Related substances (By TLC)	Comparative (Qualitative)	As p	er specification	
		General Test	S.No.3-xxiii, xxxv, xxv, xvii xxvii, xxxvi	i, As p	er specification	
69.	Doxylamine Succinate USP	Assay (By Titration)	USP35 NF30 STP No. STP/RM/10000260		to 120%	

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		Volatile related compounds	By GC	As per specification
		General test	S.No.3- xxiii, xxxv, xxv, xi, xxxvi, xxxvii	As per specification
70.	Drotaverine HCl	Assay (By Titration)	STP No. STP/RM/10000266	80% to 120%
		General test	S.No.3-xxiii, xxxv, xxv	As per specification
71.	Divalproex Sodium IP	Assay (By HPLC)	STP No. STP/RM/10004212	80% to 120%
		General test	S.No.3-xxiii, xxxv, xxv	As per specification
72.	Diacerein	Assay (By Titration)	STP No. STP/RM/10000276	80% to 120%
		General test	S.No.3-xxiii, xxxv, xxv	As per specification
73.	Duloxetine HCl	Assay (By HPLC)	STP No. STP/RM/10000277	80% to 120%
		General test	S.No.3-xxiii, xxxv, xxv	As per specification
74.	Dexibuprofen	Assay (By HPLC)	STP No. STP/RM/10000281	80% to 120%
		General test	S.No.3-xxiii, xxxv, xxv	As per specification
75.	Escitalopram Oxalate	Assay (By Titration)	STP No. STP/RM/10000309	80% to 120%
		General test	S.No.3-xxiii, xxxv, xxv	As per specification

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76.	Esomeprazole Magnesium	Assay (By HPLC)	STP No. STP/RM/10000325	80% to 120%
		General test	S.No.3-xxii, xxxv, xxv	As per specification
77.	Ergotamine Tartrate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000300	80% to 120%
		Related substances (By TLC)	Comparative (Qualitative)	As per specification
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xviii, xxxvi	As per specification
78.	Etamsylate BP	Assay (By Titration)	BP 2013 STP No. STP/RM/10000301	80% to 120%
		Hydroquinone (By TLC)	Comparative (Qualitative)	As per specification
		Related substances (By HPLC)	Impurity A: 0.1% Unspecified impurities: 0.1% Total impurities: 0.2%	0.05% to 0.5%
		General test	S.Noxxiii, xxxv, xxv, xvii, xx, xxxiii, xxxi, xxxvi, xxvii	As per specification
79.	Ephedrine Hydrochloride IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000302	80% to 120%
		Related substances (By TLC)	Comparative (Qualitative)	As per specification
		General test	S.No.3-xxiii, xxxv, xxv, xvii, iii, xviii, xxvi, xxvii, xxxvi	As per specification
80.	Enalapril Maleate IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000305	80% to 120%

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		General test	S.No.3-xxiii, xxxv, xxv, xxxi, xviii, xxvii, xxxvi	As per specification
81.	Etoricoxib	Assay (By HPLC)	STP No. STP/RM/10000306	80% to 120%
		General test	S.No.3- xxiii, xxxv, xxv	As per specification
82.	Ezetimibe	Assay (By HPLC)	STP No. STP/RM/10000308	80% to 120%
		General test	S.No.3-xxiii, xxxv, xxv	As per specification
83.	Ferrous Fumarate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000335	80% to 120%
		Ferric ion (By titration)	Comparative (Qualitative) (NMT 2.0%)	0.5% to 4.0%
		General test	S.No.3-xxiii, xxxv, xxv, viii, xxxi, xxvi, xxxvi	As per specification
84.	Fluconazole USP	Assay (By Titration)	USP35 NF30 STP No. STP/RM/10000332	80% to 120%
		Related compounds	By HPLC	As per specification
		General test	S.No 3-xxiii, xxxv, xxv, xvii, xxxvi, xxi, xxxiii	As per specification
85.	Folic Acid IP	Assay (By UV)	IP 2010 STP No. STP/RM/10000331	80% to 120%
		Free amines (By UV)	Comparative (Qualitative)	As per specification

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		General test	S.No.3-xxiii,xxxv, xxv, xviii, xxvii, xxxix	As p	er specification	
86.	Famotidine USP	Assay (By Titration)	USP35 NF30 STP No. STP/RM/10000333	80%	to 120%	
		Chromatographic purity	By TLC	As p	er specification	
		General test	S.No.3-xxiii, xxxv, xxv, xxxvi, xxi, xxxi	As p	er specification	
87.	Furazolidone IP	Assay (By UV)	IP 2010 STP No. STP/RM/10005115	80%	to 120%	
		Nitrofurfural diacetate (By TLC)	Comparative (Qualitative)	As p	er specification	
		General test	S.No.3-xxiii, xxxv, xxv, xx, xxvii, xxxvi	As p	er specification	
88.	Ferrous Gluconate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000337	80%	to 120%	
		Ferric iron	NMT 1.0% (Comparative (Qualitative))	As p	er specification	
		Oxalic acid	Comparative (Qualitative)	As p	er specification	
		Reducing sugar	Comparative (Qualitative)	As p	er specification	
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xx, viii, xxxi, xv, xxvi, ix, xxxvi	As p	er specification	

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89.	Fluoxetine Hydrochloride USP	Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10000338	80%	to 120%	
		Related compounds (By HPLC)	Related compound A: <0.15%; α -[2- (methylamino)ethyl]benzenemeth anol: <0.25%; Related compound B: <0.2%; Any other individual impurity: <0.1%; Total impurities: <0.5%	0.05%	6 to 1.0%	
		General test	S.No.3-xxiii, xxxv, xxv, xxxix, xxxi	As pe	er specification	
90.	Ferrous Ascorbate	Ferrous iron (By Titration)	STP No. STP/RM/10000339	Not le	ess than 10%	
		Ascorbic Acid Content (By Titration)	STP No. STP/RM/10000339	15%	to 105%	
		General test	S.No.3-xxiii, xxxv, xxv	As pe	er specification	
91.	Flunarizine Dihydrochloride BP	Assay (By Titration)	BP 2013 STP No. STP/RM/10000340	80%	to 120%	
		Related substances (By HPLC)	Impurity A, D: <0.1%; Impurity B: <0.5%; Impurity C: <0.25%; Any other impurity: <0.1% Total impurities: <1.0%	0.05%	6 to 2.0%	
		General test	S.No.3-xxiii, xxxv, xxv, xxxvi, xxvii	As pe	er specification	

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92.	Fenofibrate BP (MICRONISED)	Assay (By HPLC)	BP 2013 STP No. STP/RM/10000341	80% to 120%			
		Related substances (By HPLC)	Impurity A, B: <0.1%; Impurity G: <0.2%; Unspecified impurities: <0.1%; Total impurities: <0.5%	0.05% to 2.0%			
		General test	S.No.3-xxiii, xxxv, xxv, xvii, iii, xv, xxvi, xxxi, xxxvi, xxvii	As per specification			
93.	Fexofenadine Hydrochloride	Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10003891	80% to 120%			
		Limit of Fexofenadine related compound B	NMT 0.2%	0.05% to 0.4%			
		Related compounds (By HPLC)	Related compound A: <0.2%; Decarboxylated degradant: <0.15%; Any other unknown impurity: <0.1% Total impurities: <0.5%	0.05% to 2.0%			
		Content of chloride	NLT 6.45% and NMT 6.75%	Comparative (Qualitativa)			
		General test	S.No.3-xxiii, xxxv, xxv, xxxix, xxi, xxxi	(Qualitative) As per specification			
94.	Gabapentin USP	Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10000378	80% to 120%			
		Limit of chloride	NMT 0.01%	Comparative (Qualitative)			

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		General test	S.No.3-xxiii, xxxv, xxv, xx, xxxix, xxi, xxxi	As p	er specification	
95.	Gliclazide BP	Assay (By Titration)	BP 2013 STP No.STP/RM/10000372	80%	to 120%	
		Related substances (By HPLC)	Impurity F: 0.1%; Unspecified impurities: 0.1%; Sum of impurities other than F: 0.2%		% to 0.4%	
		Impurity B (By HPLC) General test	NMT 2ppm	As p	er specification	
			S.No.3-xxiii, xxxv, xxv, xxxi, xxxvi, xxvii	As p	er specification	
96.	Glimepiride USP	Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10000375	80%	to 120%	
		Limit of cis-isomer (Glimepiride related compound	NMT 0.8%	As p	er Specification	
		A) Related compounds (By HPLC)	Related compound B: <0.4%; Related compound C: <0.1%; Related compound D: <0.2%; Unspecified individual impurity 0.1%; Total (excluding Related compound B) impurities: 0.5%		% to 1.0%	
		General test	S.No.3-xxiii, xxxv, xxv, xxxix, xxi, xxxi	As p	er specification	
97.	Guaiphensin BP	Assay (By Titration)	BP 2013 STP No.STP/RM/10004306	80%	to 120%	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
		Related substances (By HPLC)	Impurity A: 0.1%; Impurity B: 1.0%; Any other impurity: 0.5%; Total (excluding impurity B) impurities: 1.0%	0.05% to 2.0%		
		Chlorides and monochlorhydrins	NMT 250ppm	Comparative (Qualitative)		
		General test	S.No.3-xxiii, xxxv, xxv, xvii, iii, xxxi, xxvii, xxxvi	As per specification		
98.	Glibenclamide IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000369	80% to 120%		
		Related compounds (By TLC)	Comparative (Qualitative)	As per specification		
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xxxi, xxvii, xxxvi	As per specification		
99.	Gatifloxacin Sesquihydrate	Assay (By Titration)	STP No. STP/RM/10000371	80% to 120%		
		General test	S.No.3-xxiii, xxxv, xxv	As per specification		
100.	Glipizide IP	Assay (By Titration)	IP 2010 STP No.STP/RM/10000370	80% to 120%		
		Related substances (By HPLC)	Single impurity: 0.5% Total impurities: 1.0%	0.05% to 2.0%		
		General test	S.No.3-xxiii, xxxv, xxv, xxvii, xxxvi	As per specification		

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	Material of Test		against which tests are performed	Limits of Detection
101.	Griseofulvin IP	Assay (By UV)	IP 2010 STP No. STP/RM/10000374	80% to 120%
		Matter soluble in light petroleum	Comparative (Qualitative) (NMT 0.2%)	As per specification
		General test	S.No.3-xxiii, xxxv, xxv, xvii, iii, xviii, xxxi, xxvii, xxxvi	As per specification
102.	Ginseng Extract	Assay (By HPLC)	STP No. STP/RM/10000377	50% to 150% of labeled claim
		General test	S.No.3-xxiii, xxxv, xxv	As per specification
103.	Gemifloxacin Mesylate	Assay (By HPLC)	STP No. STP/RM/10000382	80% to 120%
		General test	S.No.3-xxiii, xxxv, xxv, xxvii, xxxvi	As per specification
104.	Haloperidol IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10002007	80% to 120%
		Related substances (By TLC)	Comparative (Qualitative)	As per specification
		General test	S.No.3-xxiii, xxxv, xxv, xxvii, xxxvi	As per specification
105.	Hydrochlorothiazide IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000394	80% to 120%
		Related substances (By TLC)	Comparative (Qualitative)	As per specification

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection	
		General test	S.No.3-xxiii, xxxv, xxv, iii, xv xxvii, xxxvi	, As p	er specification	
106.	Indapamide USP	Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10000408	80%	to 120%	
		Chromatographic purity (By TLC)	Comparative (Qualitative)	As p	er specification	
		General test	S.No.3-xxiii, xxxv, xxv, xxvi, xxi	, As p	er specification	
107.	Ibuprofen IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000400	80%	to 120%	
		Related substances (By HPLC)	Single impurity: < 0.3% Total impurities: < 0.7%	0.059	% to 1.5%	
		General test	S.No.3- xxiii, xxxv, xxv, xvii, xviii	As p	er specification	
108.	Ivermectin BP	Assay (By HPLC)	BP 2013 STP No. STP/RM/10000405	80%	to 120%	
		Related substances (By HPLC)	Impurity with RRT 1.3 to 1.5: < 2.5%; Any other impurity: <1.0%; Total impurities: <5.0%		% to 1.5%	
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xviii, xxxi, xxxix, xxvii	As p	er specification	
109.	Ketorolac Tromethamine IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10002015	80%	to 120%	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
		Light absorption	NMT 0.10	As per specification		
		Related substances (By HPLC)	Single impurity: < 0.1% Total impurities: < 1.0%	0.05% to 2.0%		
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xxxi, xxvii, xxxvi	As per specification		
110.	Lamotrigine IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10003153	80% to 120%		
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 2.0%		
		General test	S.No.3-xxiii, xxxv, xxv, xxxi, xxvii, xxxvi	As per specification		
111.	Levetiracetam USP	Assay (By HPLC)	USP35 NF30 STP No STP/RM/10003238	80% to 120%		
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 2.0%		
		General test	S.No.3-xxiii, xxxv, xxv	As per specification		
112.	Lisinopril IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000433	80% to 120%		
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 2.0%		
		General test	S.No.3-xxiii, xxxv, xxv, xviii, xxxi, xxvii, xxxix	As per specification		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
113.	Lithium Carbonate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10003300	80% to 120%		
		General test	S.No.3-xxiii, xxxv, xxv, xvii, viii xvi, xii, xxxi, xxviii, xv, xxvi	, As per specification		
114.	Loperamide HCl IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000427	80% to 120%		
		Related substances (By HPLC)	Single impurity: < 0.5%	0.05% to 2.0%		
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xxvii, xxxvi	As per specification		
115.	Losartan Potassium IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000458	80% to 120%		
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 2.0%		
		General test	S.No.3-xxiii, xxxv, xxv, xxxi, xxxix	As per specification		
116.	Lansoprazole pellets	Assay (By HPLC)	STP No. STP/RM/10000426	As per Specification		
117.	Loratadine USP	(By HPLC) Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10000457	80% to 120%		
		Related compounds (By HPLC)	Related compounds A: <0.1%; Related compounds B: < 0.1%; Each individual unknown impurity: < 0.1%; Total impurities: < 0.3%	0.05% to 0.6%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
		General test	S.No.3-xxiii, xxxv, xxv, xi, xxxvi, xxi, xxxi	As per specification		
118.	Lysine Hydrochloride USP	Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10000439	80% to 120%		
		Chromatographic purity (By TLC)	Comparative (Qualitative)	As per Specification		
		Content of Chloride	19.0 to 19.6 (Comparative (Qualitative))	As per Specification		
		General test	S.No.3-xxiii, xxxv, xxv, xviii, xxxvi, xxi, xxvi, xxxiii, xxxi	As per specification		
119.	Levofloxacin Hemihydrate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000430	80% to 120%		
		D-ofloxacin (By HPLC)	NMT 2.0%	0.05% to 4.0%		
		General test	S.No.3-xxiii, xxxv, xxv, xxxi, xxvii, xxxix	As per specification		
120.	Levocetirizine hydrochloride IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000431	80% to 120%		
		Enantiomeric purity (By HPLC)	NLT 98.0%	90% to 110%		
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 2.0%		
		General test	S.No.3-xxiii, xxxv, xxv, xviii, xxxi, xxvii, xxxvi	As per specification		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
121.	Lutein 40%	Assay (By UV)	STP No. STP/RM/10000469	80% to 120%
		General test	S.No.3-xxiii, xxxv, xxv	As per specification
122.	Lycopene 10%	Assay (By UV)	STP No. STP/RM/10000464	$\pm20\%$ of label claim
		General test	S.No.3-xxiii, xxxv, xxv	As per specification
123.	Lactitol monohydrate BP	Assay (By HPLC)	BP 2013 STP No. STP/RM/10000466	80% to 120%
		Related substances (By HPLC)	Impurity B: <1.0% Total other impurities: <1.0%	0.05% to 2.0%
		Reducing sugars	NMT 0.2%	As per Specification
		Nickel	NMT 1ppm (Comparative (Qualitative))	As per Specification
		General test	S.No.3-xxiii, xxxv, xxv, xvii, iii, xviii, xxxviii, xxxix, xxvii	As per specification
124.	Lumefantrine	Assay (By HPLC)	STP No. STP/RM/10000467	80% to 120%
125.	Lactulose BP (powder form)	Assay (By HPLC)	BP 2013 STP No. STP/RM/10000476	80% to 120%
		Related substances (By HPLC)	Sum of impurities A, B, C, D, E: <3.0%	0.05% to 6.0%
		Methanol	NMT 50 ppm (Comparative (Qualitative))	As per Specification

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		Boron	NMT 9 ppm (Comparative (Qualitative))	As p	er Specification	
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xx xviii, xxxviii, xxxix, xxvii	, As p	er specification	
126.	Metformin HCl IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000521	80%	to 120%	
		Related substances	Qualitative test (Comparative (Qualitative))	As p	er specification	
		General test	S.No.3-xxiii, xxxv, xxv, xxxi, xxvii, xxxvi	As p	er specification	
127.	Mosapride Citrate Dihydrate	Assay (By Titration)	STP No. STP/RM/10000545	80%	to 120%	
	Dinyurau	Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05	% to 2.0%	
		General test	S.No.3-xxiii, xxxv, xxv, xxxi, xxvii, xxxix	As p	er specification	
128.	Moxifloxacin HCl BP	Assay (By HPLC)	BP2013 STP No, STP/RM/10000556	80%	to 120%	
		General test	S.No.3-xxiii, xxxv, xxv	As p	er specification	
129.	Metoprolol Succinate USP	Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10000549	80%	to 120%	
		Related compounds (By HPLC)	Single impurity: < 0.1% Total impurities: < 0.5%	0.05	% to 1.0%	

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		General test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xxxvi, xxi, xxxi	As per specification			
130.	Metoclopramide Hydrochloride IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000505	80% to 120%			
		Related substances (By TLC)	Comparative (Qualitative)	As per specification			
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xx, xxxi, xxvii	As per specification			
131.	Mefenamic Acid IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000506	80% to 120%			
		Light absorption	0.56 to 0.60	As per specification			
		Copper	Comparative (Qualitative)	As per specification			
		2,3-Dimethylaniline	Comparative (Qualitative)	As per Specification			
		(By TLC) Related substances (By TLC)	Comparative (Qualitative)	As per specification			
		General test	S.No.3-xxiii, xxxv, xxv, xxvii, xxxvi	As per specification			
132.	Methylergometrine Maleate IP	Assay (By UV)	IP 2010 STP No. STP/RM/10000507	80% to 120%			
		Related substances (By TLC)	Comparative (Qualitative)	As per specification			
		General test	S.No.3-xxiii, xxxv, xxv, xx, xviii, xxvii, xxxvi	, As per specification			
133.	Metronidazole IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000559	80% to 120%			

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		Related substances (By TLC)	Comparative (Qualitative)	As per specification	
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xxxi, xxvii, xxxvi	As per specification	
134.	Metronidazole Benzoate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000512	80% to 120%	
		Free Benzoic acid (By Titration)	NMT 0.2%	As per Specification	
		Related substances (By TLC)	Comparative (Qualitative)	As per specification	
		General test	S.No.3-xii, xxxv, xxv, xx, xxvi, xxxvi	As per specification	
135.	Magaldrate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10001826	80% to 120%	
		Soluble chloride	NMT 3.5%	As per Specification	
		Soluble sulphate	NMT 1.9%	As per Specification	
		Aluminium hydroxide	32.1% to 45.9%	As per Specification	
		Magnesium hydroxide	49.2% to 66.6%	As per Specification	
		General test	S.No.3-xxii, xxxv, xxv, viii, xxxi, xxviii, xxvi, xxxvi	As per specification	
136.	Magnesium Hydroxide IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000509	80% to 120%	
		Soluble substances	NMT 1.0%	As per specification	

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		Substances insoluble in acetic	NMT 0.1%	As p	er specification	
		acid General test	S.No.3-xxiii, xxxv, xxv, xvii, viii xxxi, xxxiii, xv, xxvi, xvi, xxxii	· .	er specification	
137.	Mebendazole IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000517	80%	to 120%	
		Related substances (By TLC)	Comparative (Qualitative)	As p	er specification	
		General test	S.No.3-xxiii, xxxv, xxv, xxxi, xxvii, xxxvi	As p	er specification	
138.	Manganese Chloride USP	Assay (By Titration)	USP35 NF30 STP No. STP/RM/10000523	80%	to 120%	
		Insoluble matter	NMT 0.005% (Comparative (Qualitative)	As p	er Specification	
		Substances not participated by ammonium sulfide	NMT 0.2% (Comparative (Qualitative))	As p	er Specification	
		General test	S.No.3-xxiii, xxxv, xxv, xx, xxxvi, xxvi, xxxiii, xxxi	As p	er specification	
139.	Magnesium Chloride IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000532	80%	to 120%	
		General test	S.No.3-xxiii, xxxv, xxv, xvii, iii, viii, xxxi, xxxiii, xvi, xxvi, iv	As p	er specification	
140.	Methylcobalamin	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000542	50%	to 150%	

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		General test	S.No.3-xxiii, xxxv, xxv	As per specification
141.	Methylprednisolone IP	Assay (By UV)	IP 2010 STP No. STP/RM/10000544	80% to 120%
		Light absorption	0.38 to 0.40	As per specification
		Related substances (By TLC)	Comparative (Qualitative)	As per specification
		General test	S.No.3- xxiii, xxxv, xxv, xviii, xxxvi	As per specification
142.	Meloxicam BP	Assay (By Titration)	BP 2013 STP No. STP/RM/10000520	80% to 120%
		Related substances (By HPLC)	Impurity A: <0.1% Impurity C: <0.05% Impurity B: <0.1% Any other impurity: <0.1% Total impurities: < 0.3%	0.02% to 0.6%
		General test	S.No xxiii, xxxv, xxv, xvii, xxxi, xxxvi, xxvii	As per specification
143.	Mycophenolate Sodium	Assay (By HPLC)	STP No. STP/RM/10000560	80% to 120%
		General test	S.No.3- xxiii, xxxv, xxv	As per specification
		Related substances (By HPLC)	Total impurities: <0.3%	0.05% to 0.6%
144.	Nicotinamide IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10002037	80% to 120%

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		Related substances (By TLC)	Comparative (Qualitative)	Asp	per specification	
		General test	S.No.3- xxiii, xxxv, xxv, xvii, xx xxxi, xv, xxvi, xxvii, xxxvi	, As p	per specification	
145.	Norfloxacin IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000577	80%	o to 120%	
		Related substances (By TLC)	Comparative (Qualitative)	Asp	per specification	
		General test	S.No.3- xxiii, xxxv, xxv, xxxi, xxvii, xxxvi	As p	per specification	
146.	Nimesulide BP	Assay (By Titration)	BP 2013 STP No. STP/RM/10000578	80%	o to 120%	
		Absorbance	NMT 0.5	Asp	per specification	
		Related substances (By HPLC)	Any impurity: <0.1% Total impurities: <0.5%	As p	per specification	
		General test	S.No.3- xxiii, xxxv, xxv, xi, xxxi xxxvi, xxvii	, As p	per specification	
147.	Nitrazepam IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000579	80%	o to 120%	
		Related substances & decomposition products (By TLC)	Comparative (Qualitative)	As p	per specification	
		General test	S.No.3- xxiii, xxxv, xxv, xxxi, xxvii, xxxvi	As p	per specification	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
148.	N-Acetylcystein USP	Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10000580	80% to 120%		
		General test	S.No.3- xxiii, xxxv, xxv, xviii, xx, xxxvi, xxi, xxxi	As per specification		
149.	Nitazoxanide	Assay (By HPLC)	STP No. STP/RM/10000582	80% to 120%		
150.	Nifedipine IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10001232	80% to 120%		
		Related substances (By HPLC)	Any impurity: <0.1% Total impurities: <0.5%	As per specification		
		General test	S.No.3-xxiii, xxxv, xxv, xxxvi, xxvii, xxxi	As per specification		
151.	Nitrofurantoin IP	Assay (By UV)	IP 2010 STP No. STP/RM/10000588	80% to 120%		
		Related substances (By TLC)	Comparative (Qualitative)	As per specification		
		General test	S.No.3-xxiii, xxxv, xxv, xxvii, xxxvi	As per specification		
152.	Omeprazole Pellets	Assay (By UV)	STP No. STP/RM/10000603	$\pm 20\%$ of label claim		
		Dissolution	STP No. STP/RM/10000603	\pm 50% of label claim		
		General test	S.No.3-xxiii, xxxv	As per specification		
153.	Omeprazole Magnesium	Assay (By HPLC)	STP No. STP/RM/10000606	80% to 120%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
		General test	S.No.3- xxiii, xxxv, xxv	As per specification		
154.	Ofloxacin IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000632	80% to 120%		
		Light absorption	NMT 0.25	As per specification		
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 3.0%		
		General test	S.No.3- xxiii, xxxv, xxv, xxxi, xxvii, xxxvi	As per specification		
155.	Ornidazole IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000602	80% to 120%		
		Related substances (By HPLC)	2-methyl-5-nitroimidozole: < 0.5% Total impurities: < 1.0%	0.05% to 2.0%		
		General test	S.No.3- xxiii, xxxv, xxv, xxxi, 31 27, 39	, As per specification		
156.	Oxetacaine BP	Assay (By Titration)	BP 2012 STP No. STP/RM/10000605	80% to 120%		

Comparative (Qualitative)

S.No.3- xxiii, xxxv, xxv, xi,

STP No. STP/RM/10000607

xxxvi, xxvii

IP 2010

Related substances (By TLC)

General test

157. Olanzapine IP Assay (By Titration)

80% to 120%

As per specification

As per specification

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection	
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 2.0%	0.05% to 4.0%	
		General test	S.No.3- xxiii, xxxv, xxv, xxxi, xxvii, xxxix	As per specification	
158.	Ondansetron Hydrochloride IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000617	80% to 120%	
		Related substances (By HPLC)	Single impurity: < 0.2% Total impurities: < 1.5%	0.05% to 3.0%	
		General test	S.No.3-xxiii, xxxv, xxv, xxvii, xxxix	As per specification	
159.	Pancreatin IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000699	80% to 120%	
		Fat	NMT 5.0%	As per specification	
		General test	S.No.3- xxiii, xxxv, xxv, xxxvi	As per specification	
160.	Pantoprazole Sodium Sesquihydrate BP	Assay (By HPLC)	STP No. STP/RM/10000637	80% to 120%	
		Related substances (By HPLC)	Impurity A: < 0.2%; Sum of impurities D & F: <0.2%; Impurities B, C, E: <0.1%; Unspecified impurities: <0.1%; Total impurities: <0.5%	0.05% to 1.0%	
		General test	S.No.3-xxiii, xxxv, xxv, xvii, xxviii	As per specification	
161.	Paracetamol IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000634	80% to 120%	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specifica against which tests an performed		ange of Testing / mits of Detection
		4-Aminophenol	50 ppm (Comparative (Qualitative		s per specification
		Related substances (By HPLC)	Comparative (Qualitative) As	s per specification
		General test	S.No. 3-xxiii, xxxv, xxv, xxvii, xxxvi	xxxi, As	s per specification
162.	Piracetam IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000		1% to 120%

		(By HPLC)	STP No. STP/RM/10000659	
		Related substances (By HPLC)	Single impurity: < 0.1% Total impurities: < 0.3%	0.05% to 0.6%
		General test	S.No.3- xxiii, xxxv, xxv, xvii, xxxi, xxvii, xxxvi	As per specification
163.	Pregabalin	Assay (By HPLC)	STP No. STP/RM/10000698	80% to 120%
		General test	S.No. 3- xxiii, xxxv, xxv	As per specification
164.	Prochlorperazine Maleate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000635	80% to 120%
		Related substances (By HPLC)	Comparative (Qualitative)	As per specification
		General test	S.No.3- xxiii, xxxv, xxv, xx, xxvii, xxxvi	As per specification
165.	Pseudoephedrine Hydrochloride IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000636	80% to 120%
		Related substances (By TLC)	Comparative (Qualitative)	As per specification

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection	
		General test	S.No.3- xxiii, xxxv, xxv, xvii, xx, xviii, xxvii, xxxvi	As per specification	
166.	Pheniramine Maleate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10002056	80% to 120%	
		Related substances (By TLC)	Comparative (Qualitative)	As per specification	
		General test	S.No.3- xxiii, xxxv, xxv, xx, xxxi, xxvii, xxxvi	As per specification	
167.	Potassium Clavulanate Diluted IP	Assay (By HPLC)	STP No STP/RM/10000214	80% to 120%	
		Light absorption	NMT 0.40	As per specification	
		Related substances (By HPLC)	Single impurity: <1.0% Total impurities: <2.0%	0.05% to 4.0%	
		General test	S.No.3- xxiii, xxxv, xxv, xx, xxxix	As per specification	
168.	Piroxicam IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000643	80% to 120%	
		General test	S.No. 3-xxiii, xxxv, xxv, xxxi, xxvii, xxxix	As per specification	
169.	Pyridoxine Hydrochloride IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000876	80% to 120%	
		Related substances (By TLC)	Comparative (Qualitative)	As per specification	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection		
		General test	S.No. 3-xxiii, xxxv, xxv, xxxi, xxvii, xxxvi, xvii, xx	As p	er specification		
170.	Pyrimethamine IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000663	80%	to 120%		
		Related substances (By TLC)	Comparative (Qualitative)	As p	er specification		
		General test	S.No. 3-xxiii, xxxv, xxv, xxxi, xvii, iii, xxvi, xxvi	As p	er specification		
171.	Phenylephrine Hydrochloride IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000670	80%	to 120%		
		Phenones	NMT 0.20% (Comparative (Qualitative))	As p	er Specification		
		Related substances (By TLC)	Comparative (Qualitative)	As p	er specification		
		General test	S.No. 3-xxiii, xxxv, xxv, xxxi, xvii, iii, xxvi, xxvi	As p	er specification		
172.	Phenazopyridine Hydrochloride USP	Assay (By UV)	USP35 NF30 STP No. STP/RM/10000686	80%	to 120%		
		Water insoluble substances	NMT 0.1%	As p	er Specification		
		Ordinary Impurities	Comparative (Qualitative)	As p	er Specification		
		General test	S.No. 3-xxiii, xxxv, xxv, xxvi, xxi, xxxi	As p	er specification		
173.	Pyrantel Pamoate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000686	80%	to 120%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection			
		Related substances (By HPLC)	Impurity A: <0.5% Impurity B: <0.2% Single impurity: <0.1% Total impurities: < 0.3%	0.05% to 1.0%			
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, xxvi, xxxiii, xxxi, xxvii, xxxvi	As per specification			
174.	Pioglitazone Hydrochloride IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000694	80% to 120%			
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 2.0%			
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, xxxi, xxvii, xxxvi	As per specification			
175.	Potassium Iodide IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000696	80% to 120%			
		Cyanide	Comparative (Qualitative)	As per specification			
		Iodate	Comparative (Qualitative)	As per specification			
		Thiosulphate	Comparative (Qualitative)	As per specification			
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, iii, viii, xxxi, xxxiii, ix, xxvi, xxxv	As per specification			
176.	Potassium Chloride IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000720	80% to 120%			
		Iodides	Comparative (Qualitative)	As per specification			

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, iii, viii, ix, xxxi, xvi, xii, xxxiii, x, xxvi, xxxvi	As per specification		
177.	Quinine Sulphate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000727	80% to 120%		
		Other cinchona alkaloids (By HPLC)	Content of dihydroquinine: <10%; Content of any related substances: <5%; Content of any other related substances: <2.5%	0.05% to 20%		
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, xx, xvii, xxvii, xxxvi	As per specification		
178.	Rabeprazole Sodium IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000731	80% to 120%		
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.5%	0.05% to 3.0%		
		General test	S.No. 3-xxiii, xxxv, xxv, xxxix,	As per specification		
179.	Ramipril IP`	Assay (By Titration)	IP 2010 STP No. STP/RM/10000732	80% to 120%		
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 2.0%		
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, xvii, xxvii, xxxvi	As per specification		
180.	Ranitidine HCl IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000728	80% to 120%		

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection	
		Related substances (By TLC)	Comparative (Qualitative)	As per specification	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, xx, xxvii, xxxvi	As per specification	
181.	Risperidone BP	Assay (By Titration)	BP 2013 STP No. STP/RM/10000733	80% to 120%	
		Related substances (By HPLC)	Impurity A, B, C,D,E: < 0.2% Any other impurity: 0.1% Total impurities: < 0.3%	0.05% to 0.6%	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, xxxvi, xxvi	As per specification	
182.	Rosiglitazone Maleate IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000739	80% to 120%	
		Related substances (By HPLC)	Single impurity: < 0.5% Total impurities: < 1.0%	0.05% to 2.0%	
		General test	S.No. 3-xxiii, xxxv, xxv, xxxi, xxvii, xxxix	As per specification	
183.	Roxithromycin IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000730	80% to 120%	
		Related substances (By HPLC)	Impurity A: 1.0%; Other individual impurity: <0.5%; Total impurities: < 3.0%;	0.05% to 6.0%	
		General test	S.No xxiii, xxxv, L6, xvii, xxxvi, xxv, xxxi, xxvii, xxxix	As per specification	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection	
184.	Rosuvastatin calcium IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000745	80%	to 120%	
		Related substances (By HPLC)	Impurity A: 1.0%; Other individual impurity: <0.5% Total impurities: < 3.0%;		% to 6.0%	
		General test	S.No xxiii, xxxv, xxv, xxxix	As p	er specification	
185.	S-Amlodipine Besilate	Assay (By HPLC)	STP No. STP/RM/10000769	80%	to 120%	
		General test	S.No 23, 35, 25	As p	er specification	
186.	Sparfloxacin	Assay (By HPLC)	STP No. STP/RM/10000767	80%	to 120%	
		General test	S.No. 3-xxiii, xxxv, xxv	As p	er specification	
187.	Sodium Valproate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000784	80%	to 120%	
		Related substances (By GC)	IP 2010	As p	er specification	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, iii, xv, xxvi, xxxi, xxxvi	As p	er specification	
188.	Salbutamol Sulphate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000762	80%	to 120%	
		Related substances (By TLC)	Comparative (Qualitative)	As p	er specification	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, iii, xiii, xxvii, xxxvi	As p	er specification	

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specificatio against which tests are performed		ge of Testing / its of Detection	
189.	Secnidazole IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000785	80%	to 120%	
		Related substances (By HPLC)	Single impurity: < 0.2% Total impurities: < 0.5%	0.059	% to 1.0%	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii xxxi, xxvii, xxxix	, As p	er specification	
190.	Selenious Acid USP	Assay (By Titration)	USP35 NF30 STP No. STP/RM/10000790	80%	to 120%	
		Insoluble matter	Comparative (Qualitative)	As p	er specification	
		Selenate & Sulfate	Comparative (Qualitative)	As p	er specification	
		General test	S.No. 3-xxiii, xxxv, xxv, xxi	As p	er specification	
191.	Serratiopeptidase IP (Coated)	Assay (By UV)	IP 2010 STP No. STP/RM/10000770	As p	er Specification	
		General test	S.No. 3-xxiii, xxxv, xxv	As p	er specification	
192.	Serratiopeptidase IP (Plain)	Assay (By UV)	IP 2010 STP No. STP/RM/10000771	As p	er Specification	
		General test	S.No. 3-xxiii, xxxv, xxv	As p	er specification	
193.	Sildenafil Citrate IP	Assay (By HPLC)	IP2010 STP No. STP/RM/10000773	80%	to 120%	
		Related substances (By HPLC)	Single impurity: < 0.2% Total impurities: < 0.5%	0.059	% to 1.0%	

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		General test	S.No. 3-xxiii, xxxv, xxv	As per specification
194.	Simethicone Emulsion USP	Assay (By IR)	USP35 NF30 STP No. STP/RM/10001839	50% to 150%
		General test	S.No. 3-xxiii, xxxv, xxv, xxxi	As per specification
195.	Simvastatin USP	Assay (By IR)	USP35 NF30 STP No. STP/RM/10000775	80% to 120%
		Chromatographic purity (By HPLC)	Sum of lovastatin & epilovastatin: <1.0%; Any individual impurity (other than lovastatin & epilovastatin): <0.4%; Total impurities (other than lovastatin & epilovastatin): 1.0%	0.05% to 2.0%
		Limit of lovastatin (By HPLC)	NMT 1.0%	0.05% to 2.0%
		General test	S.No. 3-xxiii, xxxv, xxv, xviii, xxxvi, xxv, xxxi	As per specification
196.	Sodium Ascorbate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000761	80% to 120%
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, xx, xviii, xxxi, xxxvi	As per specification
197.	Sodium Feredetate BP	Assay (By Titration)	BP 2013 STP No. STP/RM/10000779	80% to 120%
		Free sodium edetate	NLT 1.0% (Comparative (Qualitative))	As per Specification

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		Free iron	NMT 500ppm (Comparative (Qualitative))	As per Specification	
		General test	S.No. 3-xxiii, xxxv, xxv, iii, xxxi, xxvi, xxxvi	As per specification	
198.	Spironolactone IP	Assay (By UV/HPLC)	IP 2010 STP No. STP/RM/10000791	80% to 120%	
		Chromium	NMT 50 ppm (Comparative (Qualitative))	As per Specification	
		Free mercapto compounds	Comparative (Qualitative)	As per Specification	
		General test	S.No. 3-xxiii, xxxv, xxv, xviii, xxvii, xxxvi	As per specification	
199.	Sulphamethoxazole IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000763	80% to 120%	
		Related substances	Comparative (Qualitative)	As per Specification	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, iii, xxxi, xxvii, xxxvi	As per specification	
200.	Sulphadoxine IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000765	80% to 120%	
		Related substances	Comparative (Qualitative)	As per Specification	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii,iii, xxxi, xxvii, xxxvi	As per specification	
201.	Telmisartan BP	Assay	BP 2013 STP No. STP/RM/10000841	80% to 120%	

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		General test	S.No. 3-xxiii, xxxv, xxv	As per specification
202.	Tinidazole IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000824	80% to 120%
		Related substances (By TLC)	Comparative (Qualitative)	As per Specification
		General test	S.No. 3-xxiii, xxxv, xxv, xxvii, xxxvi	As per specification
203.	Torsemide USP	Assay (By HPLC)	USP35 NF30 STP No. STP/RM/10000840	80% to 120%
		Related compounds (By HPLC)	Related compound C: <0.2% Related compound B: <0.3% Related compound A: <0.5% Any other impurity: <0.1% Total other impurities: <0.2% Total impurities (including related compound A, B, C): <1.0%	0.05% to 2.0%
		General test	S.No. 3-xxiii, xxxv, xxv, xxxix xxi, xxxi	As per specification
204.	Tranexamic Acid BP	Assay (By Titration)	BP 2013 STP No. STP/RM/10000834	80% to 120%
		Related substances (By HPLC)	Impurity A: <0.1% Impurity B: <0.2% Any other Impurity: <0.1% Total of other impurities: <0.2%	0.05% to 0.5%
		Halides expressed as chlorides	NMT 140 ppm (Comparative (Qualitative)	As per specification

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Disc	cipline	Chemical Testing	Chemical Testing Issue Da		
Cert	ificate Number	T-1997	Valid U	Jntil 03.09.2015	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection	
		General test	S.No. 3-xxiii, xxxv, xxv, xx, xxxi, xxxvi, xxvii	As per specification	
205.	Tadalafil	Assay (By HPLC)	STP No. STP/RM/10000842	80% to 120%	
		General test	S.No. 3-xxiii, xxxv, xxv	As per specification	
206.	Terbutaline Sulphate IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000826	80% to 120%	
		tert-Butylamino-3,5- dihydroxyacetophenone (By TLC)	Comparative (Qualitative)	As per specification	
		Related substances (By TLC)	Comparative (Qualitative)	As per Specification	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, iii, xxxi, xxxvi	As per specification	
207.	Theophylline (Anhydrous) IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000823	80% to 120%	
		Light absorption	NLT 0.53	As per Specification	
		Related substances (By TLC)	Comparative (Qualitative)	As per Specification	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, iii	As per specification	
208.	Thiamine Hydrochloride IP	Assay (By Titration)	IP 2010 STP No. STP/RM/100037451	80% to 120%	
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, xx, xxxi, xiv, xxvi, xxvii, xxxvi	As per specification	

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		T-1997	Valid	Until 03.09.2015	
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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection
209.	Tizanidine Hydrochloride IP	Assay (By HPLC)	IP 2010 STP No. STP/RM/10000828	80%	to 120%
		Total chloride	11.9% to 12.5%	As p	er Specification
		Related substances (By HPLC)	Single impurity: <0.5% Total impurities: <1.0%	0.05	% to 2.0%
		General test	S.No. 3-xxiii, xxxv, xxv, xx, xxxi, xxvii, xxxvi	As p	er specification
210.	Topiramate USP	Assay (By HPLC)	USP 35 USNF 30 STP No. STP/RM/10000850	80%	to 120%
		Related substances (By HPLC)	Impurity A: <0.15% Impurity B: <0.15% Other impurity: <0.1%	0.05	% to 2.0%
		General test	S.No. 3-xxiii, xxxv, xxv, xviii, xxxi, xxvii, xxxix	As p	er specification
211.	Tramadol Hydrochloride BP	Assay (By Titration)	BP 2012 STP No. STP/RM/10000829	80%	to 120%
		Impurity E (By TLC)	Comparative (Qualitative)	As p	er Specification
		Related substances (By HPLC)	Impurity A: <0.2% Any other impurity: <0.1% Total impurities: <0.4%	0.05	% to 0.8%
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, iii, xviii, xxxi, xxxix, xxvii	As p	er specification
212.	Trimethoprim IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10004088	80%	to 120%

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Related substances (By TLC)	Comparative (Qualitative)	As per Specification
		General test	S.No. 3-xxiii, xxxv, xxv, xvii, xxxi, xxvii, xxxvi	As per specification
213.	Triprolidine HCl IP	Assay (By Titration)	IP 2010 STP No. STP/RM/10000825	80% to 120%
		Related substances (By TLC)	Comparative (Qualitative)	As per Specification
		General test	S.No. 3-xxiii, xxxv, xxv, xxxi, xxvii, xxxix	As per specification
214.	Venlafaxine HCl BP	Assay (By Titration)	BP 2013 STP No. STP/RM/10000895	80% to 120%
		General test	S.No. 3-xxiii, xxxv, xxv, i, xix	As per specification
215.	Vitamin A Concentrate Oil IP	Assay	IP 2010 STP No. STP/RM/10005426	80% to 120% of stated no. of units of vitamin
		General test	S.No. 3-xxiii, xxxv, xxv, i, xix	A per gm. As per specification
216.	Vitamin A Concentrate Powder IP	Assay	IP 2010 STP No. STP/RM/10000877	80% to 120% of stated no. of units of vitamin A per gm.
		Related substances & degradation products	NMT 1.054 (Comparative (Qualitative)	A per gni. As per specification
		General test	S.No. 3-xxiii, xxxv, xxv	As per specification

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S.No.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed		ge of Testing / its of Detection
217.	Voglibose	Assay (By HPLC)	STP No. STP/RM/10000892	80%	to 120%
		General test	S.No. 3-xxiii, xxxv, xxv	As p	er specification