Central Drugs Testing Laboratory, 37, Naval Hospital Road, Periamet, Chennai, Tamil Nadu	
ISO/IEC 17025: 2005	
Page 1 of 16	
19 Last Amended on	

SI.	Product / Material	Specific Test Performed	Test Method Specification	Range of Testing /
	of Test		against which tests are	Limits of Detection
			performed	

CHEMICAL TESTING

I.	DRUGS & PHARM	IACEUTICALS		
1.	Aceclofenac Tablets	Identification by HPLC	IP 2014, Vol II, Page No. 982	Qualitative
		Average Weight	IP 2014, Vol II, Page No. 982	Qualitative
		Uniformity of Weight	IP 2014, Vol II, Page No. 982	Qualitative
		Test for Dissolution	IP 2014, Vol II, Page No. 982	5.0 % to 120.0 % of Labelled Claim
		Assay	IP 2014, Vol II, Page No. 982	70.0 % to 120.0 % of Labelled Claim
2.	Albendazole Tablets	Identification by TLC & UV	IP 2014, Vol II, Page No. 1006	Qualitative
		Average Weight	IP 2014, Vol II, Page No. 1006	Qualitative
		Uniformity of Weight	IP 2014, Vol II, Page No. 1006	Qualitative
		Disintegration Test	IP 2014, Vol II, Page No. 1006	Qualitative
		Assay	IP 2014, Vol II, Page No. 1006	70.0 % to 120.0 % of Labelled Claim
3.	Bromhexine Hydrochloride	Identification by UV & Chemical Test	IP 2014, Vol II, Page No. 1203	Qualitative
	Tablets	Average Weight	IP 2014, Vol II, Page No. 1203	Qualitative
		Uniformity of Content	IP 2014, Vol II, Page No. 1203	70.0 % to 120.0 % of Labelled Claim
		Disintegration Test	IP 2014, Vol II, Page No. 1203	Qualitative
		Assay	IP 2014, Vol II, Page No. 1203	70.0 % to 120.0 % of Labelled Claim

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 2 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on
SI Product / Material	Specific Test Performed Test Meth	od Specification Range of Testing /

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
4.	Co-trimoxazole Tablets	Identification by IR & TLC	IP 2014, Vol III, Page No.2924	Qualitative
		Average Weight	IP 2014, Vol III, Page No.2924	Qualitative
		Uniformity of Weight	IP 2014, Vol III, Page No.2924	Qualitative
		Disintegration Test	IP 2014, Vol III, Page No.2924	Qualitative
		Assay 1)Sulphamethoxazole	IP 2014, Vol III, Page No.2924	70.0 % to 120.0 % of Labelled Claim
		2) Trimethoprim	IP 2014, Vol III, Page No.2924	70.0 % to 120.0 % of Labelled Claim
5.	Furazolidone Tablets	Identification by Chemical Test	IP 2014, Vol II, Page No. 1837	Qualitative
		Average Weight	IP 2014, Vol II, Page No. 1837	Qualitative
		Uniformity of Weight	IP 2014, Vol II, Page No. 1837	Qualitative
		Disintegration Test	IP 2014, Vol II, Page No. 1837	Qualitative
		Assay	IP 2014, Vol II, Page No. 1837	70.0 % to 120.0 % of Labelled Claim
6.	Glipizide Tablets	Identification by IR & UV	IP 2014, Vol II, Page No. 1868	Qualitative
		Average Weight	IP 2014, Vol II, Page No. 1868	Qualitative
		Uniformity of Content	IP 2014, Vol II, Page No. 1868	70.0 % to 120.0 % of Labelled Claim
		Disintegration Test	IP 2014, Vol II, Page No. 1868	Qualitative
		Assay	IP 2014, Vol II, Page No. 1868	70.0 % to 120.0 % of Labelled Claim
7.	Metformin Hydrochloride Tablets	Identification by IR, Chemical Test & Chloride Test	IP 2014, Vol II, Page No. 2187	Qualitative

Laboratory	Central Drugs Testing Laboratory, 37, Naval Hospital Road, Periamet,
	Chennai, Tamil Nadu

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 3 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Average Weight	IP 2014, Vol II, Page No. 2187	Qualitative
		Uniformity of Weight	IP 2014, Vol II, Page No. 2187	Qualitative
		Test for Dissolution	IP 2014, Vol II, Page No. 2187	5.0 % to 120.0 % of Labelled Claim
		Assay	IP 2014, Vol II, Page No. 2187	70.0 % to 120.0 % of Labelled Claim
8.	Nalidixic Acid Tablets	Identification by IR	IP 2014, Vol III, Page No.2293	Qualitative
		Average Weight	IP 2014, Vol III, Page No.2293	Qualitative
		Uniformity of Weight	IP 2014, Vol III, Page No.2293	Qualitative
		Disintegration Test	IP 2014, Vol III, Page No.2293	Qualitative
		Assay	IP 2014, Vol III, Page No.2293	70.0 % to 120.0 % of Labelled Claim
9.	Paracetamol Tablets	Identification by IR & Chemical Test	IP 2014, Vol III, Page No. 2434	Qualitative
		Average Weight	IP 2014, Vol III, Page No. 2434	Qualitative
		Uniformity of Weight	IP 2014, Vol III, Page No. 2434	Qualitative
		Test for Dissolution	IP 2014, Vol III, Page No. 2434	5.0 % to 120.0 % of Labelled Claim
		Assay	IP 2014, Vol III, Page No. 2434	70.0 % to 120.0 % of Labelled Claim
10.	Ibuprofen Tablets	Description	IP 2014, Vol II, Page No. 1945	Qualitative
		Identification by IR and Melting Point	IP 2014, Vol II, Page No. 1945	Qualitative
		Average Weight	IP 2014, Vol II, Page No. 1945	Qualitative

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 4 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Uniformity of Weight	IP 2014, Vol II, Page No. 1945	Qualitative
		Test for Dissolution	IP 2014, Vol II, Page No. 1945	5.0 % to 120.0 % of Labelled Claim
		Related Substances	IP 2014, Vol II, Page No. 1945	Qualitative
		Assay	IP 2014, Vol II, Page No. 1945	70.0 % to 120.0 % of Labelled Claim
11.	Riboflavin Tablets	Identification by Chemical Test	IP 2014, Vol III, Page No. 2658	Qualitative
		Average Weight	IP 2014, Vol III, Page No. 2658	Qualitative
		Uniformity of Content	IP 2014, Vol III, Page No. 2658	70.0 % to 120.0 % of Labelled Claim
		Disintegration Test	IP 2014, Vol III, Page No. 2658	Qualitative
		Assay	IP 2014, Vol III, Page No. 2658	70.0 % to 120.0 % of Labelled Claim
12.	Metronidazole Tablets	Description	IP 2014, Vol II, Page No. 2219	Qualitative
		Identification by IR, Chemical Test & Melting Point	IP 2014, Vol II, Page No. 2219	Qualitative
		Average Weight	IP 2014, Vol II, Page No. 2219	Qualitative
		Uniformity of Weight	IP 2014, Vol II, Page No. 2219	Qualitative
		Test for Dissolution	IP 2014, Vol II, Page No. 2219	5.0 % to 120.0 % of Labelled Claim
		Related Substances	IP 2014, Vol II, Page No. 2219	Qualitative
		Assay	IP 2014, Vol II, Page No. 2219	70.0 % to 120.0 % of Labelled Claim

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 5 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on
SL Product / Material	Specific Test Performed Test Me	thed Specification Bange of Test

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
13.	Mifepristone	Description	IP 2014, Vol II,	Qualitative
	Tablets		Page No. 2235	
		Identification by HPLC &	IP 2014, Vol II,	Qualitative
		UV	Page No. 2235	
		Average Weight	IP 2014, Vol II,	Qualitative
			Page No. 2235	
		Uniformity of Weight	IP 2014, Vol II,	Qualitative
			Page No. 2235	
		Test for Dissolution	IP 2014, Vol II,	5.0 % to 120.0 % of
			Page No. 2235	Labelled Claim
		Related Substances	IP 2014, Vol II,	Qualitative
			Page No. 2235	
		Assay	IP 2014, Vol II,	70.0 % to 120.0 % of
			Page No. 2235	Labelled Claim
14.	Chloramphenicol	Identification by IR or	IP 2014, Vol II,	Qualitative
	Capsules	Chemical Test	Page No. 1349	
		Average Fill	IP 2014, Vol II,	Qualitative
			Page No. 1349	
		Uniformity of Fill	IP 2014, Vol II,	Qualitative
			Page No. 1349	
		Test for Dissolution	IP 2014, Vol II,	5.0 % to 125.0 % of
			Page No. 1349	Labelled Claim
		Assay	IP 2014, Vol II,	70.0 % to 120.0 % of
			Page No. 1349	Labelled Claim
15.	Aceclofenac	Description	IP 2014, Vol II,	Qualitative
			Page No. 981	
		Identification by IR	IP 2014, Vol II,	Qualitative
			Page No. 981	
		Loss on Drying	IP 2014, Vol II,	0.01 % to 20.0 % w/w
		<u> </u>	Page No. 981	
		Sulphated Ash	IP 2014, Vol II,	0.01 % to 5.0 % w/w
			Page No. 981	
		Assay	IP 2014, Vol II,	70.0 % to 110.0 % w/w
			Page No. 981	
		<u> </u>	<u> </u>	

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 6 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on
SL Dreduct / Motorial	Specific Test Derformed Test Me	thed Specification Dange of Tee

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
16.	Albendazole	Description	IP 2014, Vol II, Page No. 1004	Qualitative
		Identification by IR & Melting Point	IP 2014, Vol II, Page No. 1004	Qualitative
		Related Substances	IP 2014, Vol II, Page No. 1004	Qualitative
		Loss on Drying	IP 2014, Vol II, Page No. 1004	0.01 % to 20.0 % w/w
		Sulphated Ash	IP 2014, Vol II, Page No. 1004	0.01 % to 5.0 % w/w
		Assay	IP 2014, Vol II, Page No. 1004	70.0 % to 110.0 %w/w
17.	Amoxycillin Trihydrate	Description	IP 2014, Vol II, Page No. 1054	Qualitative
		Identification by IR	IP 2014, Vol II, Page No. 1054	Qualitative
		рН	IP 2014, Vol II, Page No. 1054	0.1 to 14.0
		Sulphated Ash	IP 2014, Vol II, Page No. 1054	0.01 % to 5.0 % w/w
		Water	IP 2014, Vol II, Page No. 1054	0.01 % to 20.0 % w/w
		Specific optical Rotation	IP 2014, Vol II, Page No. 1054	-360 ° to +360 °
		Assay	IP 2014, Vol II, Page No. 1054	70.0 % to 110.0 % w/w
18.	Chloramphenicol	Description	IP 2014, Vol II, Page No. 1348	Qualitative
		Identification by IR	IP 2014, Vol II, Page No. 1348	Qualitative
		рН	IP 2014, Vol II, Page No. 1348	0.1 to 14.0
		Loss on Drying	IP 2014, Vol II, Page No. 1348	0.01 % to 20.0 % w/w

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 7 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Sulphated Ash	IP 2014, Vol II, Page No. 1348	0.01 % to 5.0 % w/w
		Specific optical Rotation	IP 2014, Vol II, Page No. 1348	-360 ° to +360 °
		Assay	IP 2014, Vol II, Page No. 1348	70.0 % to 110.0 % w/w
19.	Diclofenac Sodium	Description	IP 2014, Vol II, Page No. 1550	Qualitative
		Identification by IR & Chemical Test	IP 2014, Vol II, Page No. 1550	Qualitative
		рН	IP 2014, Vol II, Page No. 1550	0.1 to 14.0
		Light Absorption	IP 2014, Vol II, Page No. 1550	Qualitative
		Loss on Drying	IP 2014, Vol II, Page No. 1550	0.01 % to 20.0 % w/w
		Assay	IP 2014, Vol II, Page No. 1550	50.0 % to 120.0 % w/w
20.	Furazolidone	Description	IP 2014, Vol II, Page No. 1836	Qualitative
		Identification by IR & Chemical	IP 2014, Vol II, Page No. 1836	Qualitative
		рН	IP 2014, Vol II, Page No. 1836	0.1 to 14.0
		Loss on Drying	IP 2014, Vol II, Page No. 1836	0.01 % to 20.0 % w/w
		Sulphated Ash	IP 2014, Vol II, Page No. 1836	0.01 % to 5.0 % w/w
		Assay	IP 2014, Vol II, Page No. 1836	70.0 % to 110.0 % w/w
21.	Griseofulvin	Description	IP 2014, Vol II, Page No. 1876	Qualitative
		Identification by IR, Chemical Test & Melting Point	IP 2014, Vol II, Page No. 1876	Qualitative

Laboratory	Central Drugs Testing Laboratory, 37, Naval Hospital Road, Periamet,
-	Chennai, Tamil Nadu

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 8 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Loss on Drying	IP 2014, Vol II, Page No. 1876	0.01 % to 20.0 % w/w
		Sulphated Ash	IP 2014, Vol II, Page No. 1876	0.01 % to 5.0 % w/w
		Specific Optical Rotation	IP 2014, Vol II, Page No. 1876	-360 ° to +360 °
		Assay	IP 2014, Vol II, Page No. 1876	70.0 % to 110.0 % w/w
22.	Isoniazid	Description	IP 2014, Vol II, Page No. 2005	Qualitative
		Identification by IR	IP 2014, Vol II, Page No. 2005	Qualitative
		Loss on Drying	IP 2014, Vol II, Page No. 2005	0.01 % to 20.0 % w/w
		Sulphated Ash	IP 2014, Vol II, Page No. 2005	0.01 % to 5.0 % w/w
		рН	IP 2014, Vol II, Page No. 2005	0.1 to 14.0
		Related Substances	IP 2014, Vol II, Page No. 2005	Qualitative
		Assay	IP 2014, Vol II, Page No. 2005	70.0 % to 110.0 % w/w
23.	Levofloxacin Hemihydrate	Description	IP 2014, Vol II, Page No. 2085	Qualitative
		Identification by IR	IP 2014, Vol II, Page No. 2085	Qualitative
		Water Content	IP 2014, Vol II, Page No. 2085	0.01 % to 20.0 % w/w
		Sulphated Ash	IP 2014, Vol II, Page No. 2085	0.01 % to 5.0 % w/w
		Assay	IP 2014, Vol II, Page No. 2085	70.0 % to 110.0 %w/w
24.	Metformin Hydrochloride	Description	IP 2014, Vol II, Page No. 2186	Qualitative

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 9 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Identification by IR &	IP 2014, Vol II,	Qualitative
		Chemical Test	Page No. 2186	
		Loss on Drying	IP 2014, Vol II,	0.01 % to 20.0 % w/w
			Page No. 2186	
		Sulphated Ash	IP 2014, Vol II,	0.01 % to 5.0 % w/w
			Page No. 2186	
		Assay	IP 2014, Vol II,	50.0 % to 120.0 % w/w
		<u> </u>	Page No. 2186	
25.	Metronidazole	Description	IP 2014, Vol II,	Qualitative
			Page No. 2215	
		Identification by IR	IP 2014, Vol II,	Qualitative
			Page No. 2215	
		Loss on Drying	IP 2014, Vol II,	0.01 % to 20.0 % w/w
			Page No. 2215	
		Sulphated Ash	IP 2014, Vol II,	0.01 % to 5.0 % w/w
			Page No. 2215	
		Related Substances	IP 2014, Vol II,	Qualitative
			Page No. 2215	
		Assay	IP 2014, Vol II,	50.0 % to 120.0 % w/w
			Page No. 2215	
26.	Mebendazole	Description	IP 2014, Vol II,	Qualitative
			Page No. 2154	
		Identification by IR,	IP 2014, Vol II,	Qualitative
		Chemical Test & HPLC	Page No. 2154	
		Loss on Drying	IP 2014, Vol II,	0.01 % to 20.0 % w/w
			Page No. 2154	
		Sulphated Ash	IP 2014, Vol II,	0.01 % to 5.0 % w/w
			Page No. 2154	
		Assay	IP 2014, Vol II,	70.0 % to 110.0 % w/w
			Page No. 2154	
27.	Paracetamol	Description	IP 2014, Vol III,	Qualitative
			Page No. 2429	
		Identification by IR	IP 2014, Vol III,	Qualitative
			Page No. 2429	
		<u> </u>		

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 10 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Loss on Drying	IP 2014, Vol III, Page No. 2429	0.01 % to 20.0 % w/w
		Sulphated Ash	IP 2014, Vol III, Page No. 2429	0.01 % to 5.0 % w/w
		Related Substances	IP 2014, Vol III, Page No. 2429	Qualitative
28.	Pyridoxine Hydrochloride	Description	IP 2014, Vol III, Page No. 2600	Qualitative
		Identification by IR or UV, Chemical Test & HPLC	IP 2014, Vol III, Page No. 2600	Qualitative
		Loss on Drying	IP 2014, Vol III, Page No. 2600	0.01 % to 20.0 % w/w
		Sulphated Ash	IP 2014, Vol III, Page No. 2600	0.01 % to 5.0 % w/w
		Assay	IP 2014, Vol III, Page No. 2600	70.0 % to 110.0 % w/w
29.	Riboflavine	Description	IP 2014, Vol III, Page No. 2657	Qualitative
		Identification by IR & Chemical Test	IP 2014, Vol III, Page No. 2657	Qualitative
		рН	IP 2014, Vol III, Page No. 2657	0.1 to 14.0
		Loss on Drying	IP 2014, Vol III, Page No. 2657	0.01 % to 20.0 % w/w
		Sulphated Ash	IP 2014, Vol III, Page No. 2657	0.01 % to 5.0 % w/w
		Specific Optical Rotation	IP 2014, Vol III, Page No. 2657	-360 ° to +360 °
		Light Absorption	IP 2014, Vol III, Page No. 2657	Qualitative
		Assay	IP 2014, Vol III, Page No. 2657	70.0 % to 110.0 % w/w
30.	Rifampicin	Description	IP 2014, Vol III, Page No. 2659	Qualitative

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 11 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		Identification by IR &	IP 2014, Vol III,	Qualitative
		HPLC	Page No. 2659	
		Loss on Drying	IP 2014, Vol III,	0.01 % to 20.0 % w/w
			Page No. 2659	
		Sulphated Ash	IP 2014, Vol III,	0.01 % to 5.0 % w/w
			Page No. 2659	
		рН	IP 2014, Vol III,	0.1 to 14.0
			Page No. 2659	
		Assay	IP 2014, Vol III,	70.0 % to 110.0 % w/w
			Page No. 2659	
31.	Thiamine	Description	IP 2014, Vol III,	Qualitative
	Hydrochloride		Page No. 2856	
		Identification by IR &	IP 2014, Vol III,	Qualitative
		Chemical Test	Page No. 2856	
		Loss on Drying	IP 2014, Vol III,	0.01 % to 20.0 % w/w
			Page No. 2856	
		Sulphated Ash	IP 2014, Vol III,	0.01 % to 5.0 % w/w
			Page No. 2856	
		рН	IP 2014, Vol III,	0.1 to 14.0
			Page No. 2856	
		Assay	IP 2014, Vol III,	50.0 % to 120.0 % w/w
			Page No. 2856	
32.	Thiamine	Description	IP 2014, Vol III,	Qualitative
	Mononitrate		Page No. 2859	
		Identification by IR &	IP 2014, Vol III,	Qualitative
		Chemical Test	Page No. 2859	
		Loss on Drying	IP 2014, Vol III,	0.01 % to 20.0 % w/w
			Page No. 2859	
		Sulphated Ash	IP 2014, Vol III,	0.01 % to 5.0 % w/w
			Page No. 2859	
		рН	IP 2014, Vol III,	0.1 to 14.0
			Page No. 2859	
		Assay	IP 2014, Vol III,	50.0 % to 120.0 % w/w
			Page No. 2859	

Laboratory	Central Drugs Testing Laboratory, 37, Naval Hospital Road, Periamet, Chennai, Tamil Nadu

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 12 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on
SI Product / Material	Specific Test Performed Test Me	whod Specification Bange of Testing /

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
33.	Trimethoprim	Description	IP 2014, Vol III, Page No. 2922	Qualitative
		Identification by IR	IP 2014, Vol III, Page No. 2922	Qualitative
		Loss on Drying	IP 2014, Vol III, Page No. 2922	0.01 % to 20.0 % w/w
		Sulphated Ash	IP 2014, Vol III, Page No. 2922	0.01 % to 5.0 % w/w
		Assay	IP 2014, Vol III, Page No. 2922	50.0 % to 120.0 % w/w
34.	Acyclovir	Description	IP 2014, Vol II, Page No. 988	Qualitative
		Identification by IR	IP 2014, Vol II, Page No. 988	Qualitative
		Sulphated Ash	IP 2014, Vol II, Page No. 988	0.01 % to 5.0 % w/w
		Water Content	IP 2014, Vol II, Page No. 988	0.01 % to 20.0 % w/w
		Assay	IP 2014, Vol II, Page No. 988	70.0 % to 110.0 % w/w
35.	Ceftriaxone Sodium	Description	IP 2014, Vol II, Page No. 1323	Qualitative
		Identification by IR and Chemical Test	IP 2014, Vol II, Page No. 1323	Qualitative
		рН	IP 2014, Vol II, Page No. 1323	0.1 to 14.0
		Water Content	IP 2014, Vol II, Page No. 1323	0.01 % to 20.0 % w/w
		Specific Optical Rotation	IP 2014, Vol II, Page No. 1323	-360 ° to +360 °
36.	Cephalexin	Description	IP 2014, Vol II, Page No. 1333	Qualitative
		Identification by IR & HPLC	IP 2014, Vol II, Page No. 1333	Qualitative

1

Accreditation Standard	ISO/IEC 17025: 2005		
Certificate Number	TC-5674	Page 13 of 16	
Validity	29.11.2017 to 28.11.2019	Last Amended on	

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
		рН	IP 2014, Vol II,	0.1 to 14.0
			Page No. 1333	
		Sulphated Ash	IP 2014, Vol II,	0.01 % to 5.0 % w/w
			Page No. 1333	
		Water Content	IP 2014, Vol II,	0.01 % to 20.0 % w/w
			Page No. 1333	
		Specific Optical Rotation	IP 2014, Vol II,	-360 ° to +360 °
		<u> </u>	Page No. 1333	
		Assay	IP 2014, Vol II,	70.0 % to 110.0 % w/w
			Page No. 1333	
37.	Enalapril Maleate	Description	IP 2014, Vol II,	Qualitative
			Page No. 1655	
		Identification by IR and	IP 2014, Vol II,	Qualitative
		Melting Point	Page No. 1655	
		Sulphated Ash	IP 2014, Vol II,	0.01 % to 5.0 % w/w
			Page No. 1655	
		Loss on Drying	IP 2014, Vol II,	0.01 % to 20.0 % w/w
			Page No. 1655	
		Specific Optical Rotation	IP 2014, Vol II,	-360 ° to +360 °
			Page No. 1655	
		Assay	IP 2014, Vol II,	70.0 % to 110.0 % w/w
			Page No. 1655	
38.	Frusemide	Description	IP 2014, Vol II,	Qualitative
			Page No. 1833	
		Identification by IR	IP 2014, Vol II,	Qualitative
			Page No. 1833	
		Related Substances	IP 2014, Vol II,	Qualitative
			Page No. 1833	
		Sulphated Ash	IP 2014, Vol II,	0.01 % to 5.0 % w/w
			Page No. 1833	
		Loss on Drying	IP 2014, Vol II,	0.01 % to 20.0 % w/w
			Page No. 1833	
•••••		Assay	IP 2014, Vol II,	70.0 % to 110.0 % w/w
			Page No. 1833	

--

Laboratory	Central Drugs Testing Laboratory, 37, Naval Hospital Road, Periamet, Chennai, Tamil Nadu

Accreditation Standard	ISO/IEC 17025: 2005	
Certificate Number	TC-5674	Page 14 of 16
Validity	29.11.2017 to 28.11.2019	Last Amended on
SI Product / Material	Specific Test Performed Test Ma	thad Specification Range of Test

SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection
39.	Ibuprofen	Description	IP 2014, Vol II, Page No. 1941	Qualitative
		Identification by IR	IP 2014, Vol II, Page No. 1941	Qualitative
		Sulphated Ash	IP 2014, Vol II, Page No. 1941	0.01 % to 5.0 % w/w
		Loss on Drying	IP 2014, Vol II, Page No. 1941	0.01 % to 20.0 % w/w
		Optical Rotation	IP 2014, Vol II, Page No. 1941	-360 ° to +360 °
		Assay	IP 2014, Vol II, Page No. 1941	70.0 % to 110.0 % w/w
40.	Methyldopa	Description	IP 2014, Vol II, Page No. 2199	Qualitative
		Identification by IR	IP 2014, Vol II, Page No. 2199	Qualitative
		Sulphated Ash	IP 2014, Vol II, Page No. 2199	0.01 % to 5.0 % w/w
		Water Content	IP 2014, Vol II, Page No. 2199	0.01 % to 20.0 % w/w
		Optical Rotation	IP 2014, Vol II, Page No. 2199	-360 ° to +360 °
		Assay	IP 2014, Vol II, Page No. 2199	70.0 % to 110.0 % w/w
41.	Mifepristone	Description	IP 2014, Vol II, Page No. 2234	Qualitative
		Identification by IR	IP 2014, Vol II, Page No. 2234	Qualitative
		Light Absorption	IP 2014, Vol II, Page No. 2234	Qualitative
		Sulphated Ash	IP 2014, Vol II, Page No. 2234	0.01 % to 5.0 % w/w
		Loss on Drying	IP 2014, Vol II, Page No. 2234	0.01 % to 20.0 % w/w

Laboratory		Central Drugs Testing Laboratory, 37, Naval Hospital Road, Periamet, Chennai, Tamil Nadu				
Acc	reditation Standard	ISO/IEC 17025: 2005	ISO/IEC 17025: 2005			
Cer	tificate Number	TC-5674	TC-5674 Page 15 of 16			
Validity		29.11.2017 to 28.11.20	29.11.2017 to 28.11.2019 Last Amend			
SI.	Product / Material of Test	Specific Test Performed	Test Method Specification against which tests are performed	Range of Testing / Limits of Detection		
		Specific Optical Rotation	IP 2014, Vol II, Page No. 2234	-360 ° to +360 °		
42.	Ofloxacin	Description	IP 2014, Vol II, Page No. 2367	Qualitative		
		Identification by IR	IP 2014, Vol II, Page No. 2367	Qualitative		
		Light Absorption	IP 2014, Vol II, Page No. 2367	Qualitative		
		Related Substances	IP 2014, Vol II, Page No. 2367	Qualitative		
		Sulphated Ash	IP 2014, Vol II, Page No. 2367	0.01 % to 5.0 % w/w		
		Loss on Drying	IP 2014, Vol II, Page No. 2367	0.01 % to 20.0 % w/w		
		Assay	IP 2014, Vol II, Page No. 2367	70.0 % to 110.0 % w/w		

Laboratory		Central Drugs Testing Laboratory, 37, Naval Hospital Road, Periamet, Chennai, Tamil Nadu			
Accreditation Standard		ISO/IEC 17025: 2005			
Certificate Number		TC-5674	Page 16 of	16	
Validity		29.11.2017 to 28.11.20	19 Last Amer	ided on	
SI. Product / Material of Test		Specific Test Performed	Test Method Specification against which tests are		

MECHANICAL TESTING

performed

.....

I.	RUBBER AND RUBBER PRODUCTS			
1.	Rubber Latex Male Condoms and	Dimension a) Length	As per Schedule 'R' of D & C Act/ISO 4074:2002	Upto 300 mm
	related products	b) Width		Upto 60 mm
		c) Thickness		Upto 1 mm
		Burst Volume before & after ageing	As per Schedule 'R' of D & C Act/ISO 4074:2002	0.1 l to 50 l
		Burst Pressure before & after ageing	As per Schedule 'R' of D & C Act/ISO 4074:2002	0.1 Kpa to 3 Kpa
		Water Leakage Test	As per Schedule 'R' of D & C Act/ISO 4074:2002	Qualitative
		Package Integrity Test	As per Schedule 'R' of D & C Act/ISO 4074:2002	Qualitative
		Quantity of Lubricant	As per Schedule 'R' of D & C Act	0.01 g to 100 g
		Colour fastness	As per Schedule 'R' of D & C Act	Qualitative