SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	1 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
Permanent Facility				
1	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Gabapentine and Methylcobalamine tablets	Assay of Methylcobalamine	STP/FP/40037440-00
2	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Lactic Acid bacillus Tablets	Assay	STP/FP/40030217-00
3	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Pharmaceutical Dosage Like Tablets, Capsules	Test for specific organism (P.aeruginosa)	IP 2018 Vol.-I Appendix 2.2.9, Ph. Eur. 10th Edition General chapter 2.6.13, USP NF- 2022 chapter 62 STP/RM/10009691-02 20/06/19
4	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Pharmaceutical Dosage Like Tablets, Capsules	Test for specific organism (Clostridia)	IP 2018, Vol.-I Appendix 2.2.9, USP NF- 2022 chapter 62, Ph. Eur. 10th Edition General chapter 2.6.13,STP/RM/10009691-02 20/06/19
5	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Pharmaceutical Dosage Like Tablets, Capsules	Test for specific organism (Candida albicans)	IP 2018 Vol.-I Appendix 2.2.9, USP NF- 2022 chapter 62 EP volume 01 ,STP/RM/10009691-02
6	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Pharmaceutical Dosage Like Tablets, Capsules	Test for specific organism (Escherichia coli)	IP 2018, Vol.-I Appendix 2.2.9, USP NF- 2022chapter 62, Ph. Eur. 10th Edition General chapter 2.6.13 STP/RM/10009691-02 20/06/19
7	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Pharmaceutical Dosage Like Tablets, Capsules	Test for specific organism (S.aureus)	IP 2018 Vol.-I Appendix 2.2.9, Ph. Eur. 10th Edition General chapter 2.6.13, USP NF- 2022chapter 62 STP/RM/10009691-02 20/06/19
8	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Pharmaceutical Dosage Like Tablets, Capsules	Test for specific organism (Salmonella)	IP 2018 Vol-I Appendix 2.2.9 , Ph. Eur. 10th Edition General chapter 2.6.13, USP NF- 2022 chapter 62 STP/RM/10009691-02 20/06/19
9	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Pharmaceutical Dosage Like Tablets, Capsules	Test for specific organism (Shigella boydii)	IP 2018, Vol.-I Appendix 2.2.9 STP/RM/10009691-02 20/06/19
10	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Pharmaceutical Dosage Like Tablets, Capsules	Total aerobic microbial count/ Viable bacterial count (By Pour plate method)	IP 2018 Vol.- I, Appendix 2.2.9, Ph. Eur. 10th Edition General chapter 2.6.12, USP NF- 2022chapter 61

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	2 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
11	BIOLOGICAL- DRUGS AND PHARMACEUTICALS	Soft gelatin capsule containing omega 3 fatty acid ,chromium picolinate ,wheat germ oil ,menadione sodium ,bisulphite, biotine ,potasium iodide , vitamin B12,sodium molybdate ,dihydrate , sodium selenite pentahydrate ,potassium chloride , sodium borate ,nickle sulphate ,vitamine b6 , vitamin D3 , Copper sulphate pentahydrate ,folic acid , choline bitartrate , colloidal silicon dioxide , stannous chloride dihydrate , sodium metavenadate ,green tea extract , genseng extract , ginkgo biloba extract,grap seed extract , magnesium oxide , Benfotiamine ,citrus bioflavinoids ,natural mixed carotine oil. lutein , lactic acid baccilus ,piperine, ferrous fumerate ,zinc oxide , niacinamide , magnese sulphate , calcium ascorbate , dibasic calium phosphate .	Assay of vitamin B12	STP/FP/40031656-11
12	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	Assay (For Aceclofenac)	By HPLC STP/FP/40030672-01 Effective Date : 10/11/21
13	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	Assay (For paracetamol)	By HPLC STP/FP/40030672-01 Effective Date : 10/11/21
14	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	General Tests for Tablets (Average Weight)	STP/FP/40030672-01 Effective Date : 10/11/21
15	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	General Tests for Tablets (Disintegration Time)	STP/FP/40030672-01 Effective Date : 10/11/21
16	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	General Tests for Tablets (Friability)	STP/FP/40030672-01 Effective Date : 10/11/21
17	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	General Tests for Tablets (Hardness)	STP/FP/40030672-01 Effective Date : 10/11/21
18	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	General Tests for Tablets (Length)	STP/FP/40030672-01 Effective Date : 10/11/21
19	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	General Tests for Tablets (Thickness)	STP/FP/40030672-01 Effective Date : 10/11/21
20	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/40030672-01 Effective Date : 10/11/21
21	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	General Tests for Tablets (Width)	STP/FP/40030672-01 Effective Date : 10/11/21
22	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac & Paracetamol Tablets	Identification By HPLC	STP/FP/40030672-01 Effective Date : 10/11/21
23	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac IP	Assay	IP 2018 STP No.- STP/RM/10000046-06 Effective Date- 10/04/20
24	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac IP	General Test (Heavy Metals)	IP 2018 STP No.- STP/RM/10000046-06 Effective Date- 10/04/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	3 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
25	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac IP	General Test (Identification By IR)	IP 2018 STP No.- STP/RM/10000046-06 Effective Date- 10/04/20
26	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac IP	General Test (Loss on Drying)	IP 2018 STP No.- STP/RM/10000046-06 Effective Date- 10/04/20
27	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000046-06 Effective Date- 10/04/20
28	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac IP	Related substances (Single Impurity) (By HPLC)	IP 2018 STP No.- STP/RM/10000046-06 Effective Date- 10/04/20
29	CHEMICAL- DRUGS & PHARMACEUTICALS	Aceclofenac IP	Related substances (Total Impurities) (By HPLC)	IP 2018 STP No.- STP/RM/10000046-06 Effective Date- 10/04/20
30	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam IP	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000013-07 Effective Date- 28/10/21
31	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam IP	General Test (Heavy Metals)	IP 2018 STP No.- STP/RM/10000013-07 Effective Date- 28/10/21
32	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam IP	General Test (Identification By IR)	IP 2018 STP No.- STP/RM/10000013-07 Effective Date- 28/10/21
33	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam IP	General Test (Loss on Drying)	IP 2018 STP No.- STP/RM/10000013-07 Effective Date- 28/10/21
34	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000013-07 Effective Date- 28/10/21
35	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	Assay By HPLC	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19
36	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	General Tests for Tablets (Average Weight)	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19
37	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	General Tests for Tablets (Diameter)	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19
38	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	General Tests for Tablets (Disintegration Time)	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19
39	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	General Tests for Tablets (Friability)	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19

SCOPE OF ACCREDITATION

Laboratory Name : AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA

Accreditation Standard ISO/IEC 17025:2017

Certificate Number TC-6742 **Page No** 4 of 43

Validity 29/12/2022 to 28/12/2024 **Last Amended on** 24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
40	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	General Tests for Tablets (Hardness)	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19:
41	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	General Tests for Tablets (Thickness)	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19
42	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	General Tests for Tablets (Uniformity of Weight)	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19
43	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	Identification	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19
44	CHEMICAL- DRUGS & PHARMACEUTICALS	Alprazolam Tablets IP	Uniformity of content By HPLC	IP 2018 STP/FP/40037434-00 Effective Date : 09/08/19
45	CHEMICAL- DRUGS & PHARMACEUTICALS	Amlodipine Besylate IP	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000028-05 Effective Date- 04/06/20
46	CHEMICAL- DRUGS & PHARMACEUTICALS	Amlodipine Besylate IP	General Test (Identification By IR)	IP 2018 STP No.- STP/RM/10000028-05 Effective Date- 04/06/20
47	CHEMICAL- DRUGS & PHARMACEUTICALS	Amlodipine Besylate IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000028-05 Effective Date- 04/06/20:
48	CHEMICAL- DRUGS & PHARMACEUTICALS	Amlodipine Besylate IP	General Test (Water)	IP 2018 STP No.- STP/RM/10000028-05 Effective Date- 04/06/20
49	CHEMICAL- DRUGS & PHARMACEUTICALS	Amlodipine Besylate IP	Related substances (Single Impurity) (BY HPLC)	IP 2018 STP No.- STP/RM/10000028-05 Effective Date-04/06/20
50	CHEMICAL- DRUGS & PHARMACEUTICALS	Amlodipine Besylate IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No.- STP/RM/10000028-05 Effective Date- 04/06/20
51	CHEMICAL- DRUGS & PHARMACEUTICALS	Amlodipine Besylate IP	Related substances (By TLC)	IP 2018 STP No.- STP/RM/10000028-05 Effective Date- 04/06/20
52	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium clavulanate tablets	Assay (For Potassium Clavulanate) By HPLC	IP 2018 STP/FP/40037144 Effective Date- 31/07/19
53	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium clavulanate tablets	Assay (For Amoxycillin) By HPLC	IP 2018 STP/FP/40037144 Effective Date- 31/07/19
54	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium clavulanate tablets	Dissolution (For Amoxycillin) By HPLC	IP 2018 STP/FP/40037144 Effective Date- 31/07/19
55	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium clavulanate tablets	Dissolution (For Potassium Clavulanate) By HPLC	IP 2018 STP/FP/40037144 Effective Date- 31/07/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	5 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
56	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium Clavulanate Tablets	General Tests for Tablets (Average Weight)	IP 2018 STP/FP/40037144 Issue No- 00 Effective Date- 31/07/19
57	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium clavulanate tablets	General Tests for Tablets (Length)	IP 2018 STP/FP/40037144 Issue No- 00 Effective Date- 31/07/19
58	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium Clavulanate Tablets	General Tests for Tablets (Thickness)	IP 2018 STP/FP/40037144 Issue No- 00 Effective Date- 31/07/19
59	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium Clavulanate Tablets	General Tests for Tablets (Uniformity of Weight)	IP 2018 STP/FP/40037144 Issue No- 00 Effective Date- 31/07/19
60	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium Clavulanate Tablets	General Tests for Tablets (Width)	IP 2018 STP/FP/40037144 Issue No- 00 Effective Date- 31/07/19
61	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium clavulanate tablets	Identification By HPLC	IP 2018 STP/FP/40037144 Issue No- 00 Effective Date- 31/07/19
62	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin & Potassium clavulanate tablets	Uniformity of Content (For Potassium Clavulanate) By HPLC	IP 2018 STP/FP/40037144 Effective Date- 31/07/19
63	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin Trihydrate IP	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000006-06 Effective Date- 08/04/20
64	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin Trihydrate IP	General Test (Identification By IR)	IP 2018 STP No.- STP/RM/10000006-06 Effective Date- 08/04/20
65	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin Trihydrate IP	General Test (pH)	IP 2018 STP No.- STP/RM/10000006-06 Effective Date- 08/04/20
66	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin Trihydrate IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000006-06 Effective Date- 08/04/20
67	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin Trihydrate IP	General Test (Water)	IP 2018 STP No.- STP/RM/10000006-06 Effective Date- 08/04/20
68	CHEMICAL- DRUGS & PHARMACEUTICALS	Amoxycillin Trihydrate IP	N,N-Dimethylaniline	IP 2018 STP No.- STP/RM/10000006-06 Effective Date- 08/04/20
69	CHEMICAL- DRUGS & PHARMACEUTICALS	Ampicillin Trihydrate IP	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000008-06 Effective Date- 07/12/20
70	CHEMICAL- DRUGS & PHARMACEUTICALS	Ampicillin Trihydrate IP	General Test (Heavy Metals)	IP 2018 STP No.- STP/RM/10000008-06 Effective Date- 07/12/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	6 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
71	CHEMICAL- DRUGS & PHARMACEUTICALS	Ampicillin Trihydrate IP	General Test (Identification By IR)	IP 2018 STP No.- STP/RM/10000008-06 Effective Date- 07/12/20
72	CHEMICAL- DRUGS & PHARMACEUTICALS	Ampicillin Trihydrate IP	General Test (pH)	IP 2018 STP No.- STP/RM/10000008-06 Effective Date- 07/12/20
73	CHEMICAL- DRUGS & PHARMACEUTICALS	Ampicillin Trihydrate IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000008-06 Effective Date- 07/12/20
74	CHEMICAL- DRUGS & PHARMACEUTICALS	Ampicillin Trihydrate IP	General Test (Water)	IP 2018 STP No.- STP/RM/10000008-06 Effective Date- 07/12/20
75	CHEMICAL- DRUGS & PHARMACEUTICALS	Ampicillin Trihydrate IP	N,N-Dimethylaniline	IP 2018 STP No.- STP/RM/10000008-06 Effective Date- 07/12/20
76	CHEMICAL- DRUGS & PHARMACEUTICALS	Atenolol IP	Assay (By Titration)	IP 2018 STP No.- STP/RM/10000039-03 Effective Date- 26/02/19
77	CHEMICAL- DRUGS & PHARMACEUTICALS	Atenolol IP	General Test (Identification By IR)	IP 2018 STP No.- STP/RM/10000039-03 Effective Date- 26/02/19
78	CHEMICAL- DRUGS & PHARMACEUTICALS	Atenolol IP	General Test (Loss on Drying)	IP 2018 STP No.- STP/RM/10000039-03 Effective Date- 26/02/19
79	CHEMICAL- DRUGS & PHARMACEUTICALS	Atenolol IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000039-03 Effective Date- 26/02/19
80	CHEMICAL- DRUGS & PHARMACEUTICALS	Atenolol IP	Related substances (Single Impurity) (By HPLC)	IP 2018 STP No.- STP/RM/10000039-03 Effective Date- 26/02/19
81	CHEMICAL- DRUGS & PHARMACEUTICALS	Atenolol IP	Related substances (Total Impurities) (By HPLC)	IP 2018 STP No.- STP/RM/10000039-03 Effective Date- 26/02/19
82	CHEMICAL- DRUGS & PHARMACEUTICALS	Atorvastatin Calcium IP	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000051-06 Effective Date- 11/04/20
83	CHEMICAL- DRUGS & PHARMACEUTICALS	Atorvastatin Calcium IP	General Test (Heavy Metals)	IP 2018 STP No.- STP/RM/10000051-06 Effective Date- 11/04/20
84	CHEMICAL- DRUGS & PHARMACEUTICALS	Atorvastatin Calcium IP	General Test (Water)	IP 2018 STP No.- STP/RM/10000051-06 Effective Date- 11/04/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	7 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
85	CHEMICAL- DRUGS & PHARMACEUTICALS	Atorvastatin Calcium IP	Related substances (Single Impurity) (By HPLC)	IP 2018 STP No.- STP/RM/10000051-06 Effective Date- 11/04/20
86	CHEMICAL- DRUGS & PHARMACEUTICALS	Atorvastatin Calcium IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No.- STP/RM/10000051-06 Effective Date- 11/04/20
87	CHEMICAL- DRUGS & PHARMACEUTICALS	Atorvastatin Tablets	Assay	STP/FP/40031129 IP2018
88	CHEMICAL- DRUGS & PHARMACEUTICALS	Atorvastatin Tablets	Dissolution	STP/FP/40031129 IP2018
89	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin IP	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000047-07 Effective Date- 11/04/20
90	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin IP	General Test (Heavy Metals)	IP 2018 STP No.- STP/RM/10000047-07 Effective Date- 11/04/20
91	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin IP	General Test (pH)	IP 2018 STP No.- STP/RM/10000047-07 Effective Date- 11/04/20
92	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000047-07 Effective Date- 11/04/20
93	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin IP	General Test (Water)	IP 2018 STP No.- STP/RM/10000047-07 Effective Date- 11/04/20
94	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin IP	Related substances (Impurity B) (By HPLC)	IP 2018 STP No.- STP/RM/10000047-07 Effective Date- 11/04/20
95	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin IP	Related substances (Total Impurities) (By HPLC)	IP 2018 STP No.- STP/RM/10000047-07 Effective Date- 11/04/20
96	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin Tablets IP	Assay	By HPLC IP 2018 STP/FP/ 40004108-01 Effective Date: 01/07/18
97	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin Tablets IP	Dissolution	By HPLC IP 2018 STP/FP/ 40004108-01 Effective Date: 01/07/18
98	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin Tablets IP	General Tests for Tablets (Average Weight)	IP 2018 STP/FP/ 40004108-01 Effective Date: 01/07/18
99	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin Tablets IP	General Tests for Tablets (Disintegration Test)	IP 2018 STP/FP/ 40004108-01 Effective Date: 01/07/18

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	8 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
100	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin Tablets IP	General Tests for Tablets (Length)	IP 2018 STP/FP/ 40004108-01 Effective Date: 01/07/18
101	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin Tablets IP	General Tests for Tablets (Thickness)	IP 2018 STP/FP/ 40004108-01 Effective Date: 01/07/18
102	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin Tablets IP	General Tests for Tablets (Uniformity of Weight)	IP 2018 STP/FP/ 40004108-01 Effective Date: 01/07/18
103	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin Tablets IP	General Tests for Tablets (Width)	IP 2018 STP/FP/ 40004108-01 Effective Date: 01/07/18
104	CHEMICAL- DRUGS & PHARMACEUTICALS	Azithromycin Tablets IP	Identification	By HPLC IP 2018 STP/FP/ 40004108-01 Effective Date: 01/07/18
105	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium and Vitamin D3 Tablet	Assay By HPLC (For Vitamin D3)	STP/FP/40025265 Effective Date-08/05/19
106	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium and Vitamin D3 Tablet	Assay By Titration (For Calcium)	STP/FP/40025265 Effective Date-08/05/19
107	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium and Vitamin D3 Tablet	General Tests for Tablets (Average Weight)	STP/FP/40025265 Effective Date-08/05/19
108	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium and Vitamin D3 Tablet	General Tests for Tablets (Disintegration Test)	STP/FP/40025265 Effective Date-08/05/19
109	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium and Vitamin D3 Tablet	General Tests for Tablets (Length)	STP/FP/40025265 Effective Date-08/05/19
110	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium and Vitamin D3 Tablet	General Tests for Tablets (Thickness)	STP/FP/40025265 Effective Date-08/05/19
111	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium and Vitamin D3 Tablet	General Tests for Tablets (Uniformity of Weight)	STP/FP/40025265 Effective Date-08/05/19
112	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium and Vitamin D3 Tablet	General Tests for Tablets (Width)	STP/FP/40025265 Effective Date-08/05/19
113	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium and Vitamin D3 Tablet	Identification	As per Assay STP/FP/40025265 Effective Date-08/05/19
114	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium Carbonate IP	Assay (By Titration)	IP 2018 STP No.- STP/RM/10000152-07 Effective Date- 29/08/20
115	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium Carbonate IP	General Test (Arsenic)	IP 2018 STP No.- STP/RM/10000152-07 Effective Date- 29/08/20
116	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium Carbonate IP	General Test (Heavy Metals)	IP 2018 STP No.- STP/RM/10000152-07 Effective Date- 29/08/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	9 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
117	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium Carbonate IP	General Test (Lead)	IP 2018 STP No.- STP/RM/10000152-07 Effective Date- 29/08/20
118	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium Carbonate IP	General Test (Loss on Drying)	IP 2018 STP No.- STP/RM/10000152-07 Effective Date- 29/08/20
119	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium Citrate Maleate	Assay (By Titration)	STP No.- STP/RM/10000194-04 Effective Date- 04/05/17
120	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium Citrate Maleate	General Test (Arsenic)	STP No.- STP/RM/10000194-04 Effective Date- 04/05/17
121	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium Citrate Maleate	General Test (Heavy Metals)	STP No.- STP/RM/10000194-04 Effective Date- 04/05/17
122	CHEMICAL- DRUGS & PHARMACEUTICALS	Calcium Citrate Maleate	General Test (Loss on Drying)	STP No.- STP/RM/10000194-04 Effective Date- 04/05/17
123	CHEMICAL- DRUGS & PHARMACEUTICALS	Carbamazepine IP	Assay (By HPLC)	IP 2018 STP/RM/150002509-04 Effective Date- 24/04/20
124	CHEMICAL- DRUGS & PHARMACEUTICALS	Carbamazepine IP	General Test (Chloride)	IP 2018 STP/RM/150002509-04 Effective Date- 24/04/20
125	CHEMICAL- DRUGS & PHARMACEUTICALS	Carbamazepine IP	General Test (Heavy Metals)	IP 2018 STP/RM/150002509-04 Effective Date- 24/04/20
126	CHEMICAL- DRUGS & PHARMACEUTICALS	Carbamazepine IP	General Test (Identification By IR)	IP 2018 STP/RM/150002509-04 Effective Date- 24/04/20
127	CHEMICAL- DRUGS & PHARMACEUTICALS	Carbamazepine IP	General Test (Loss on Drying)	IP 2018 STP/RM/150002509-04 Effective Date- 24/04/20
128	CHEMICAL- DRUGS & PHARMACEUTICALS	Carbamazepine IP	General Test (Sulphated Ash)	IP 2018 STP/RM/150002509-04 Effective Date- 24/04/20
129	CHEMICAL- DRUGS & PHARMACEUTICALS	Carbamazepine IP	Related Substances (Single Impurity) (By HPLC)	IP 2018 STP/RM/150002509-04 Effective Date- 24/04/20
130	CHEMICAL- DRUGS & PHARMACEUTICALS	Carbamazepine IP	Related Substances (Total Impurities) (By HPLC)	IP 2018 STP/RM/150002509-04 Effective Date- 24/04/20
131	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefdinir USP	Assay (By HPLC)	USP45 NF40 STP No.: STS/RM/10007345-04 Effective Date- 28/12/20
132	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefdinir USP	General Test (Identification By IR)	USP NF- 2022 STP No.: STS/RM/10007345-04 Effective Date- 28/12/20
133	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefdinir USP	General Test (Residue on Ignition)	USP NF- 2022 STP No.: STS/RM/10007345-04 Effective Date- 28/12/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	10 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
134	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefdinir USP	General Test (Water)	USP NF- 2022 STP No.: STS/RM/10007345-04 Effective Date- 28/12/20
135	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefdinir USP	Related substances (Individual unknown Impurity) By HPLC	USP45 NF40 STP No.: STS/RM/10007345-04 Effective Date- 28/12/20
136	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefdinir USP	Related substances (Total Impurities) By HPLC	USP45 NF40 STP No.: STS/RM/10007345-04 Effective Date- 28/12/20
137	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefixime Trihydrate IP	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000132-04 Effective Date- 12/03/19
138	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefixime Trihydrate IP	General Test (Identification)	IP 2018 STP No.- STP/RM/10000132-04 Effective Date- 12/03/19
139	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefixime Trihydrate IP	General Test (pH)	IP 2018 STP No.- STP/RM/10000132-04 Effective Date- 12/03/19
140	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefixime Trihydrate IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000132-04 Effective Date- 12/03/19
141	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefixime Trihydrate IP	General Test (Water)	IP 2018 STP No.- STP/RM/10000132-04 Effective Date- 12/03/19
142	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefixime Trihydrate IP	Related substances (Single Impurity) (By HPLC)	IP 2018 STP No.- STP/RM/10000132-04 Effective Date- 12/03/19
143	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefixime Trihydrate IP	Related substances (Total Impurities) (By HPLC)	IP 2018 STP No.- STP/RM/10000132-04 Effective Date- 12/03/19
144	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefpodoxime tablets	Assay By HPLC	STP/FP/40032091-00
145	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefpodoxime Tablets	General Tests for Tablets (Average Weight)	STP/FP/40032091-00: 2019
146	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefpodoxime tablets	General Tests for Tablets (Diameter)	STP/FP/40032091-00: 2019
147	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefpodoxime Tablets	General Tests for Tablets (Disintegration Time)	STP/FP/40032091-00: 2019
148	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefpodoxime Tablets	General Tests for Tablets (Thickness)	STP/FP/40032091-00: 2019

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	11 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
149	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefpodoxime Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/40032091-00: 2019
150	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefuroxime Axetil IP	Acetone	IP 2018 STP No.- STP/RM/10000196-04 Effective Date- 02/08/20
151	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefuroxime Axetil IP	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000196-04 Effective Date- 02/08/20
152	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefuroxime Axetil IP	Diastereoisomer ratio	IP 2018 STP No.- STP/RM/10000196-04 Effective Date- 02/08/20
153	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefuroxime Axetil IP	General Test (Identification by IR)	IP 2018 STP No.- STP/RM/10000196-04 Effective Date- 02/08/20
154	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefuroxime Axetil IP	General Test (Water)	IP 2018 STP No.- STP/RM/10000196-04 Effective Date- 02/08/20
155	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefuroxime Axetil IP	Related substances (D3-isomer) (By HPLC)	IP 2018 STP No.- STP/RM/10000196-04 Effective Date- 02/08/20
156	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefuroxime Axetil IP	Related substances (E-isomer) (By HPLC)	IP 2018 STP No.- STP/RM/10000196-04 Effective Date- 02/08/20
157	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefuroxime Axetil IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No.- STP/RM/10000196-04 Effective Date- 02/08/20
158	CHEMICAL- DRUGS & PHARMACEUTICALS	Cefuroxime Axetil IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No.- STP/RM/10000196-04 Effective Date- 02/08/20
159	CHEMICAL- DRUGS & PHARMACEUTICALS	Cetirizine Hydrochloride IP	Assay (By Titration)	IP 2018 STP No.- STP/RM/10000125-07 Effective Date- 11/03/20
160	CHEMICAL- DRUGS & PHARMACEUTICALS	Cetirizine Hydrochloride IP	General Test (Identification by IR)	IP 2018 STP No.- STP/RM/10000125-07 Effective Date- 11/03/20
161	CHEMICAL- DRUGS & PHARMACEUTICALS	Cetirizine Hydrochloride IP	General Test (Loss on Drying)	IP 2018 STP No.- STP/RM/10000125-07 Effective Date- 11/03/20
162	CHEMICAL- DRUGS & PHARMACEUTICALS	Cetirizine Hydrochloride IP	General Test (pH)	IP 2018 STP No.- STP/RM/10000125-07 Effective Date- 11/03/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	12 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
163	CHEMICAL- DRUGS & PHARMACEUTICALS	Cetirizine Hydrochloride IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000125-07 Effective Date- 11/03/20
164	CHEMICAL- DRUGS & PHARMACEUTICALS	Ciprofloxacin HCl IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10000119-06 Effective Date- 20/04/20
165	CHEMICAL- DRUGS & PHARMACEUTICALS	Ciprofloxacin HCl IP	Fluoroquinolonic acid (By TLC)	IP 2018 STP No. STP/RM/10000119-06 Effective Date- 20/04/20
166	CHEMICAL- DRUGS & PHARMACEUTICALS	Ciprofloxacin HCl IP	General Tests (Heavy Metals)	General Tests (Heavy Metals)
167	CHEMICAL- DRUGS & PHARMACEUTICALS	Ciprofloxacin HCl IP	General Tests (Identification by IR)	IP 2018 STP No. STP/RM/10000119-06 Effective Date- 20/04/20
168	CHEMICAL- DRUGS & PHARMACEUTICALS	Ciprofloxacin HCl IP	General Tests (Sulphated Ash)	IP 2018 STP No. STP/RM/10000119-06 Effective Date- 20/04/20
169	CHEMICAL- DRUGS & PHARMACEUTICALS	Ciprofloxacin HCl IP	General Tests (Sulphates)	IP 2018 STP No. STP/RM/10000119-06 Effective Date- 20/04/20
170	CHEMICAL- DRUGS & PHARMACEUTICALS	Ciprofloxacin HCl IP	General Tests (Water)	IP 2018 STP No. STP/RM/10000119-06 Effective Date- 20/04/20
171	CHEMICAL- DRUGS & PHARMACEUTICALS	Ciprofloxacin HCl IP	Related substances (Single Impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000119-06 Effective Date- 20/04/20
172	CHEMICAL- DRUGS & PHARMACEUTICALS	Ciprofloxacin HCl IP	Related substances (Total Impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000119-06 Effective Date- 20/04/20
173	CHEMICAL- DRUGS & PHARMACEUTICALS	Citicoline Sodium	Assay	STP No. STP/RM/10000120-05 Effective Date- 28/01/20
174	CHEMICAL- DRUGS & PHARMACEUTICALS	Citicoline Sodium	General Test (Chlorides)	STP No. STP/RM/10000120-05 Effective Date- 28/01/20
175	CHEMICAL- DRUGS & PHARMACEUTICALS	Citicoline Sodium	General Test (Heavy Metals)	STP No. STP/RM/10000120-05 Effective Date- 28/01/20
176	CHEMICAL- DRUGS & PHARMACEUTICALS	Citicoline Sodium	General Test (Identification by IR)	STP No. STP/RM/10000120-05 Effective Date- 28/01/20
177	CHEMICAL- DRUGS & PHARMACEUTICALS	Citicoline Sodium	General Test (Iron)	STP No. STP/RM/10000120-05 Effective Date- 28/01/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	13 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
178	CHEMICAL- DRUGS & PHARMACEUTICALS	Citicoline Sodium	General Test (pH)	STP No. STP/RM/10000120-05 Effective Date- 28/01/20
179	CHEMICAL- DRUGS & PHARMACEUTICALS	Citicoline Sodium	General Test (Water)	STP No. STP/RM/10000120-05 Effective Date- 28/01/20
180	CHEMICAL- DRUGS & PHARMACEUTICALS	Clarithromycin	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000227-04 Effective Date- 09/07/20
181	CHEMICAL- DRUGS & PHARMACEUTICALS	Clarithromycin	General Test (Heavy Metals)	IP 2018 STP No.- STP/RM/10000227-04 Effective Date- 09/07/20
182	CHEMICAL- DRUGS & PHARMACEUTICALS	Clarithromycin	General Test (Identification by IR)	IP 2018 STP No.- STP/RM/10000227-04 Effective Date- 09/07/20
183	CHEMICAL- DRUGS & PHARMACEUTICALS	Clarithromycin	General Test (Water)	IP 2018 STP No.- STP/RM/10000227-04 Effective Date- 09/07/20
184	CHEMICAL- DRUGS & PHARMACEUTICALS	Clarithromycin	Related substances (Single Impurity) (By HPLC)	IP 2018 STP No.- STP/RM/10000227-04 Effective Date- 09/07/20
185	CHEMICAL- DRUGS & PHARMACEUTICALS	Clarithromycin	Related substances (Total Impurities) (By HPLC	IP 2018 STP No.- STP/RM/10000227-04 Effective Date- 09/07/20
186	CHEMICAL- DRUGS & PHARMACEUTICALS	Clonazepam IP	Assay	IP 2018 STP No.- STP/RM/10000206-04 Effective Date- 15/03/19
187	CHEMICAL- DRUGS & PHARMACEUTICALS	Clonazepam IP	General Test (Identification by IR)	IP 2018 STP No.- STP/RM/10000206-04 Effective Date- 15/03/19
188	CHEMICAL- DRUGS & PHARMACEUTICALS	Clonazepam IP	General Test (Loss on Drying)	IP 2018 STP No.- STP/RM/10000206-04 Effective Date- 15/03/19
189	CHEMICAL- DRUGS & PHARMACEUTICALS	Clonazepam IP	General Test (Sulphated Ash)	IP 2018 STP No.- STP/RM/10000206-04 Effective Date- 15/03/19
190	CHEMICAL- DRUGS & PHARMACEUTICALS	Clonazepam IP	Related substances (Single Impurity) (By HPLC)	IP 2018 STP No.- STP/RM/10000206-04 Effective Date- 15/03/19
191	CHEMICAL- DRUGS & PHARMACEUTICALS	Clonazepam IP	Related substances (Total Impurities) (By HPLC)	IP 2018 STP No.- STP/RM/10000206-04 Effective Date- 15/03/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	14 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
192	CHEMICAL- DRUGS & PHARMACEUTICALS	Clopidogrel Bisulfate IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10000200-03 Effective Date- 11/01/19
193	CHEMICAL- DRUGS & PHARMACEUTICALS	Clopidogrel Bisulfate IP	General Test (Identification by IR)	IP 2018 STP No. STP/RM/10000200-03 Effective Date- 11/01/19
194	CHEMICAL- DRUGS & PHARMACEUTICALS	Clopidogrel Bisulfate IP	General Test (Loss on Drying)	IP 2018 STP No. STP/RM/10000200-03 Effective Date- 11/01/19
195	CHEMICAL- DRUGS & PHARMACEUTICALS	Clopidogrel Bisulfate IP	General Test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000200-03 Effective Date- 11/01/19
196	CHEMICAL- DRUGS & PHARMACEUTICALS	Clopidogrel Bisulfate IP	Related substances (Total Impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000200-03 Effective Date- 11/01/19
197	CHEMICAL- DRUGS & PHARMACEUTICALS	Clopidogrel Bisulfate IP	Related substances (Single Impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000200-03 Effective Date- 11/01/19
198	CHEMICAL- DRUGS & PHARMACEUTICALS	Cloxacillin Sodium IP	Assay (By HPLC)	IP 2018 STP No.- STP/RM/10000110-05 Effective Date- 20/08/19
199	CHEMICAL- DRUGS & PHARMACEUTICALS	Cloxacillin Sodium IP	General Test (Identification by IR)	IP 2018 STP No.- STP/RM/10000110-05 Effective Date- 20/08/19
200	CHEMICAL- DRUGS & PHARMACEUTICALS	Cloxacillin Sodium IP	General Test (pH)	IP 2018 STP No.- STP/RM/10000110-05 Effective Date- 20/08/19
201	CHEMICAL- DRUGS & PHARMACEUTICALS	Cloxacillin Sodium IP	General Test (Water)	IP 2018 STP No.- STP/RM/10000110-05 Effective Date- 20/08/19
202	CHEMICAL- DRUGS & PHARMACEUTICALS	Cloxacillin Sodium IP	N,N-Dimethylaniline	IP 2018 STP No.- STP/RM/10000110-05 Effective Date- 20/08/19
203	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	Assay (Diclofenac Sodium)	By HPLC STP/FP/40030677 Effective Date-20/11/19
204	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	Assay (Paracetamol)	By HPLC STP/FP/40030677 Effective Date-20/11/19
205	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	General Tests for Tablets (Average Weight)	STP/FP/40030677 Effective Date-20/11/19
206	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	General Tests for Tablets (Disintegration Time)	STP/FP/40030677 Effective Date-20/11/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	15 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
207	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	General Tests for Tablets (Friability)	STP/FP/40030677 Effective Date-20/11/19
208	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	General Tests for Tablets (Hardness)	STP/FP/40030677 Effective Date-20/11/19
209	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	General Tests for Tablets (Length)	STP/FP/40030677 Effective Date-20/11/19
210	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	General Tests for Tablets (Thickness)	STP/FP/40030677 Effective Date-20/11/19
211	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/40030677 Effective Date-20/11/19
212	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	General Tests for Tablets (Width)	STP/FP/40030677 Effective Date-20/11/19
213	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	Identification	By HPLC STP/FP/40030677 Effective Date-20/11/19
214	CHEMICAL- DRUGS & PHARMACEUTICALS	Diclofenac Sodium & Paracetamol Tablets	Uniformity of content for Diclofenac	By HPLC STP/FP/40030677 Effective Date-20/11/19
215	CHEMICAL- DRUGS & PHARMACEUTICALS	Dicyclomine & Paracetamol Tablets	Assay (For Paracetamol)	STP/FP/ 40031218
216	CHEMICAL- DRUGS & PHARMACEUTICALS	Dicyclomine & Paracetamol Tablets	General Tests for Tablets (Average Weight)	STP/FP/ 40031218: 2019
217	CHEMICAL- DRUGS & PHARMACEUTICALS	Dicyclomine & Paracetamol Tablets	General Tests for Tablets (Diameter)	General Tests for Tablets (Diameter)
218	CHEMICAL- DRUGS & PHARMACEUTICALS	Dicyclomine & Paracetamol Tablets	General Tests for Tablets (Disintegration Time)	STP/FP/ 40031218: 2019
219	CHEMICAL- DRUGS & PHARMACEUTICALS	Dicyclomine & Paracetamol Tablets	General Tests for Tablets (Friability)	STP/FP/ 40031218: 2019
220	CHEMICAL- DRUGS & PHARMACEUTICALS	Dicyclomine & Paracetamol Tablets	General Tests for Tablets (Hardness)	STP/FP/ 40031218: 2019
221	CHEMICAL- DRUGS & PHARMACEUTICALS	Dicyclomine & Paracetamol Tablets	General Tests for Tablets (Thickness)	General Tests for Tablets (Thickness)
222	CHEMICAL- DRUGS & PHARMACEUTICALS	Dicyclomine & Paracetamol Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/ 40031218: 2019
223	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium Delayed Release Tablets USP	Assay By HPLC	STP/FP/40023402 Effective Date- 10/11/17
224	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium Delayed Release Tablets USP	Dissolution By HPLC	STP/FP/40023402 Effective Date- 10/11/17
225	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium Delayed Release Tablets USP	General Tests for Tablets (Average Weight)	STP/FP/40023402 Effective Date- 10/11/17

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	16 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
226	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium Delayed Release Tablets USP	General Tests for Tablets (Length)	STP/FP/40023402 Effective Date- 10/11/17
227	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium Delayed Release Tablets USP	General Tests for Tablets (Thickness)	STP/FP/40023402 Effective Date- 10/11/17
228	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium Delayed Release Tablets USP	General Tests for Tablets (Uniformity of Weight)	STP/FP/40023402 Effective Date- 10/11/17
229	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium Delayed Release Tablets USP	General Tests for Tablets (Width)	STP/FP/40023402 Effective Date- 10/11/17
230	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium Delayed Release Tablets USP	Identification By HPLC	STP/FP/40023402 Effective Date- 10/11/17
231	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium IP	Assay (By HPLC)	IP2018 STP No. STP/RM/10004212-01 Effective Date- 03/10/18
232	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium IP	General Test (Heavy Metals)	IP2018 STP No. STP/RM/10004212-01 Effective Date- 03/10/18
233	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium IP	General Test (Identification by IR)	IP2018 STP No. STP/RM/10004212-01 Effective Date- 03/10/18
234	CHEMICAL- DRUGS & PHARMACEUTICALS	Divalproex Sodium IP	General Test (Water)	IP2018 STP No. STP/RM/10004212-01 Effective Date- 03/10/18
235	CHEMICAL- DRUGS & PHARMACEUTICALS	Domperidone Maleate IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000253-05 Effective Date- 04/03/20
236	CHEMICAL- DRUGS & PHARMACEUTICALS	Domperidone Maleate IP	General Test (Heavy Metals)	IP 2018 STP No. STP/RM/10000253-05 Effective Date- 04/03/20
237	CHEMICAL- DRUGS & PHARMACEUTICALS	Domperidone Maleate IP	General Test (Identification by IR)	IP 2018 STP No. STP/RM/10000253-05 Effective Date- 04/03/20
238	CHEMICAL- DRUGS & PHARMACEUTICALS	Domperidone Maleate IP	General Test (Loss on Drying)	IP 2018 STP No. STP/RM/10000253-05 Effective Date- 04/03/20
239	CHEMICAL- DRUGS & PHARMACEUTICALS	Domperidone Maleate IP	General Test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000253-05 Effective Date- 04/03/20
240	CHEMICAL- DRUGS & PHARMACEUTICALS	Domperidone Maleate IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000253-05 Effective Date- 04/03/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	17 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
241	CHEMICAL- DRUGS & PHARMACEUTICALS	Domperidone Maleate IP	Related substances (Total Impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000253-05 Effective Date- 04/03/20
242	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine HCl	Assay (By Titration)	STP No. STP/RM/10000277-02 Effective Date- 23/03/18
243	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine HCl	General Test (Heavy Metals)	STP No. STP/RM/10000277-02 Effective Date- 23/03/18: 2018
244	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine HCl	General Test (Identification by IR)	STP No. STP/RM/10000277-02 Effective Date- 23/03/18
245	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine HCl	General Test (Loss on Drying)	STP No. STP/RM/10000277-02 Effective Date- 23/03/18
246	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine HCl	General Test (Sulphated Ash)	STP No. STP/RM/10000277-02 Effective Date- 23/03/18
247	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine Hydrochloride Tablets	Assay By HPLC	STP/FP/40031098 Effective Date- 20/04/18
248	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine Hydrochloride Tablets	General Tests for Tablets (Average Weight)	STP/FP/40031098 Effective Date- 20/04/18
249	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine Hydrochloride Tablets	General Tests for Tablets (Disintegration Time)	General Tests for Tablets (Disintegration Time)
250	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine Hydrochloride Tablets	General Tests for Tablets (Uniformity of Weight)	General Tests for Tablets (Uniformity of Weight)
251	CHEMICAL- DRUGS & PHARMACEUTICALS	Duloxetine Hydrochloride Tablets	Identification By HPLC	STP/FP/40031098 Effective Date- 20/04/18
252	CHEMICAL- DRUGS & PHARMACEUTICALS	Enalapril Maleate IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10000305-03 Effective Date- 22/01/19
253	CHEMICAL- DRUGS & PHARMACEUTICALS	Enalapril Maleate IP	General Test (Heavy Metals)	IP 2018 STP No. STP/RM/10000305-03 Effective Date- 22/01/19
254	CHEMICAL- DRUGS & PHARMACEUTICALS	Enalapril Maleate IP	General Test (Identification by IR)	IP 2018 STP No. STP/RM/10000305-03 Effective Date- 22/01/19
255	CHEMICAL- DRUGS & PHARMACEUTICALS	Enalapril Maleate IP	General Test (Loss on Drying)	IP 2018 STP No. STP/RM/10000305-03 Effective Date- 22/01/19
256	CHEMICAL- DRUGS & PHARMACEUTICALS	Enalapril Maleate IP	General Test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000305-03 Effective Date- 22/01/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	18 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
257	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole 40 mg (As enteric coated pellets) & Domperidone 30 mg (As sustained release pellets) Capsules	Dissolution By HPLC	STP/FP/40031214 Issue No- 00 Effective Date- 12/04/18
258	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole 40 mg (As enteric coated pellets) & Domperidone 30 mg (As sustained release pellets) Capsules	General Tests for Capsule: (Disintegration Test)	STP/FP/40031214 Issue No- 00 Effective Date- 12/04/18
259	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole 40 mg (As enteric coated pellets) & Domperidone 30 mg (As sustained release pellets) Capsules	Identification By HPLC	STP/FP/40031214 Issue No- 00 Effective Date- 12/04/18
260	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole 40 mg (As enteric coated pellets) &Domperidone 30 mg (As sustained release pellets) Capsule	General Tests for Capsule: (Average Fill)	STP/FP/40031214 Issue No- 00 Effective Date- 12/04/18
261	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole 40 mg (As enteric coated pellets) &Domperidone 30 mg (As sustained release pellets) Capsule	General Tests for Capsule: (Uniformity of Weight)	STP/FP/40031214 Issue No- 00 Effective Date- 12/04/18
262	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole 40mg (As enteric coated pelletes) &Domperidone 30 mg (As sustained release pellets)	Assay of Domperidone By HPLC	STP/FP/40031214 Issue No-00 Effective Date-12/04/18
263	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole 40mg (As enteric coated pelletes) &Domperidone 30 mg (As sustained release pellets)	Assay of Esomeprazole By HPLC-	STP/FP/40031214 Issue No -00Effective Date -12/04/18
264	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole Magnesium IP	Assay (By HPLC)	IP2018 STP No. STP/RM/10000316-04 Effective Date- 29/04/20
265	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole Magnesium IP	General Test (Heavy Metals)	IP2018 STP No. STP/RM/10000316-04 Effective Date- 29/04/20:
266	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole Magnesium IP	General Test (Identification by IR)	IP2018 STP No. STP/RM/10000316-04 Effective Date- 29/04/20
267	CHEMICAL- DRUGS & PHARMACEUTICALS	Esomeprazole Magnesium IP	General test (Water)	IP2018 STP No. STP/RM/10000316-04 Effective Date- 29/04/20
268	CHEMICAL- DRUGS & PHARMACEUTICALS	Fexofenadine Hydrochloride USP	Assay (By HPLC)	USP45 NF40 STP No. STP/RM/10003891-03 Effective Date- 12/05/18
269	CHEMICAL- DRUGS & PHARMACEUTICALS	Fexofenadine Hydrochloride USP	Content of chloride	USP45 NF40 STP No. STP/RM/10003891-03 Effective Date- 12/05/18
270	CHEMICAL- DRUGS & PHARMACEUTICALS	Fexofenadine Hydrochloride USP	General test (Identification by IR)	USP NF- 2022 STP No. STP/RM/10003891-03 Effective Date- 12/05/18

SCOPE OF ACCREDITATION

Laboratory Name : AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA

Accreditation Standard ISO/IEC 17025:2017

Certificate Number TC-6742 **Page No** 19 of 43

Validity 29/12/2022 to 28/12/2024 **Last Amended on** 24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
271	CHEMICAL- DRUGS & PHARMACEUTICALS	Fexofenadine Hydrochloride USP	General test (Residue on Ignition)	USP NF- 2022 STP No. STP/RM/10003891-03 Effective Date- 12/05/18
272	CHEMICAL- DRUGS & PHARMACEUTICALS	Fexofenadine Hydrochloride USP	General test (Water)	General test (Water)
273	CHEMICAL- DRUGS & PHARMACEUTICALS	Fexofenadine Hydrochloride USP	Limit of Fexofenadine related compound B (By HPLC)	USP45 NF40 STP No. STP/RM/10003891-03 Effective Date- 12/05/18
274	CHEMICAL- DRUGS & PHARMACEUTICALS	Fexofenadine Hydrochloride USP	Related compounds (By HPLC)	USP45 NF40 STP No. STP/RM/10003891-03 Effective Date- 12/05/18
275	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole Tablet IP	Assay By HPLC	IP 2018 STP/FP/40026792-01 Effective Date-11/09/20
276	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole Tablet IP	General Tests for Tablets (Average Weight)	IP 2018 STP/FP/40026792-01 Effective Date-11/09/20
277	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole Tablet IP	General Tests for Tablets (Disintegration Time)	IP 2018 STP/FP/40026792-01 Effective Date-11/09/20
278	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole Tablet IP	General Tests for Tablets (Friability)	IP 2018 STP/FP/40026792-01 Effective Date-11/09/20
279	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole Tablet IP	General Tests for Tablets (Hardness)	IP 2018 STP/FP/40026792-01 Effective Date-11/09/20
280	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole Tablet IP	General Tests for Tablets (Thickness)	IP 2018 STP/FP/40026792-01 Effective Date-11/09/20
281	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole Tablet IP	General Tests for Tablets (Uniformity of Weight)	IP 2018 STP/FP/40026792-01 Effective Date-11/09/20
282	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole Tablet IP	General Tests for Tablets (Width)	IP 2018 STP/FP/40026792-01 Effective Date-11/09/20
283	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole Tablet IP	Identification	IP 2018 STP/FP/40026792-01 Effective Date-11/09/20
284	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole USP	Assay (By Titration)	USP45 NF40 STP No. STP/RM/10000332-03 Effective Date- 20/08/18
285	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole USP	General test (Identification by IR)	USP NF- 2022 STP No. STP/RM/10000332-03 Effective Date- 20/08/18
286	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole USP	General test (Loss on Drying)	USP NF- 2022 STP No. STP/RM/10000332-03 Effective Date- 20/08/18

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	20 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
287	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluconazole USP	General test (Residue on Ignition)	USP NF- 2022 STP No. STP/RM/10000332-03 Effective Date- 20/08/18
288	CHEMICAL- DRUGS & PHARMACEUTICALS	Fluoxetine Hydrochloride	Identification (By IR)	BY IR STP/RM/10000349-05 Effective Date: 07/05/20
289	CHEMICAL- DRUGS & PHARMACEUTICALS	Gabapentin USP	Assay (By HPLC)	USP45 NF40 STP/RM/10000378-04 Effective Date- 26/12/20
290	CHEMICAL- DRUGS & PHARMACEUTICALS	Gabapentin USP	General test (Identification by IR)	USP NF- 2022 STP/RM/10000378-04 Effective Date- 26/12/20
291	CHEMICAL- DRUGS & PHARMACEUTICALS	Gabapentin USP	General test (pH)	USP NF- 2022 STP/RM/10000378-04 Effective Date- 26/12/20
292	CHEMICAL- DRUGS & PHARMACEUTICALS	Gabapentin USP	General test (Residue on Ignition)	USP NF- 2022 STP/RM/10000378-04 Effective Date- 26/12/20
293	CHEMICAL- DRUGS & PHARMACEUTICALS	Gabapentin USP	General test (Water)	USP NF- 2022 STP/RM/10000378-04 Effective Date- 26/12/20
294	CHEMICAL- DRUGS & PHARMACEUTICALS	Glibenclamide IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000369-05 Effective Date- 23/03/20
295	CHEMICAL- DRUGS & PHARMACEUTICALS	Glibenclamide IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000369-05 Effective Date- 23/03/20
296	CHEMICAL- DRUGS & PHARMACEUTICALS	Glibenclamide IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000369-05 Effective Date- 23/03/20
297	CHEMICAL- DRUGS & PHARMACEUTICALS	Glibenclamide IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000369-05 Effective Date- 23/03/20
298	CHEMICAL- DRUGS & PHARMACEUTICALS	Glibenclamide IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000369-05 Effective Date- 23/03/20
299	CHEMICAL- DRUGS & PHARMACEUTICALS	Gliclazide BP	Assay (By Titration)	BP 2022 STP No. STP/RM/10000372-04 Effective Date- 14/12/19
300	CHEMICAL- DRUGS & PHARMACEUTICALS	Gliclazide BP	General test (Identification by IR)	BP 2022 STP No. STP/RM/10000372-04 Effective Date- 14/12/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	21 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
301	CHEMICAL- DRUGS & PHARMACEUTICALS	Gliclazide BP	General test (Loss on Drying)	BP 2022 STP No. STP/RM/10000372-04 Effective Date- 14/12/19
302	CHEMICAL- DRUGS & PHARMACEUTICALS	Gliclazide BP	General test (Sulphated Ash)	BP 2022 STP No. STP/RM/10000372-04 Effective Date- 14/12/19
303	CHEMICAL- DRUGS & PHARMACEUTICALS	Gliclazide BP	Related substances (Single Impurity) (By HPLC)	BP 2022 STP No. STP/RM/10000372-04 Effective Date- 14/12/19
304	CHEMICAL- DRUGS & PHARMACEUTICALS	Gliclazide BP	Related substances (Total Impurities) (By HPLC)	BP 2020 STP No. STP/RM/10000372-04 Effective Date- 14/12/19
305	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	Assay By HPLC (Glimepiride),	STP/FP/40037614 Effective Date- 04/12/19
306	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	Assay By UV (Metformin HCl)	STP/FP/40037614 Effective Date- 04/12/19
307	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	General Tests for Tablets (Average Weight)	STP/FP/40037614 Effective Date- 04/12/19
308	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	General Tests for Tablets (Friability)	STP/FP/40037614 Effective Date- 04/12/19
309	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	General Tests for Tablets (Hardness)	Date- 04/12/19
310	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	General Tests for Tablets (Length)	STP/FP/40037614 Effective Date- 04/12/19
311	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	General Tests for Tablets (Thickness)	STP/FP/40037614 Effective Date- 04/12/19
312	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/40037614 Effective Date- 04/12/19
313	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	General Tests for Tablets (Width)	STP/FP/40037614 Effective Date- 04/12/19
314	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	Identification	As per Assay STP/FP/40037614 Effective Date- 04/12/19
315	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride & Metformin Hydrochloride SR Tablets	Uniformity of content for Glimepiride	By HPLC STP/FP/40037614 Effective Date- 04/12/19
316	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride USP	Assay (By HPLC)	USP45 NF40 STP No. STP/RM/10000375-04 Effective Date- 13/01/22
317	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride USP	General test (Identification by IR)	USP NF- 2022STP No. STP/RM/10000375-04 Effective Date- 13/01/22

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	22 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
318	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride USP	General test (Residue on Ignition)	USP NF- 2022STP No. STP/RM/10000375-04 Effective Date- 13/01/22
319	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride USP	General test (Water)	USP NF- 2022STP No. STP/RM/10000375-04 Effective Date- 13/01/22
320	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride USP	Related compounds (Related compound B) (By HPLC)	USP45 NF40 STP No. STP/RM/10000375-04 Effective Date- 13/01/22
321	CHEMICAL- DRUGS & PHARMACEUTICALS	Glimepiride USP	Related compounds (Total Impurities) (By HPLC)	USP45 NF40 STP No. STP/RM/10000375-04 Effective Date- 13/01/22
322	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000400-03 Effective Date-30/04/20
323	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000400-03 Effective Date-30/04/20
324	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000400-03 Effective Date-30/04/20
325	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen IP	General test (LOD)	IP 2018 STP No. STP/RM/10000400-03 Effective Date-30/04/20
326	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000400-03 Effective Date-30/04/20
327	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000400-03 Effective Date-30/04/20
328	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000400-03 Effective Date-30/04/20
329	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen Tablet IP	Assay (By Titration)	IP2018 STP/FP/190002467-02 Effective Date- 20/08/19
330	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen Tablet IP	Dissolution By UV	IP2018 STP/FP/190002467-02 Effective Date- 20/08/19
331	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen Tablet IP	General Tests for Tablets (Average Weight)	IP2018 STP/FP/190002467-02 Effective Date- 20/08/19
332	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen Tablet IP	General Tests for Tablets (Uniformity of Weight)	IP2018 STP/FP/190002467-02 Effective Date- 20/08/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	23 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
333	CHEMICAL- DRUGS & PHARMACEUTICALS	Ibuprofen Tablet IP	Identification A) By IR B) By Melting point	IP2018 STP/FP/190002467-02 Effective Date- 20/08/19
334	CHEMICAL- DRUGS & PHARMACEUTICALS	Levetiracetam USP	Assay (By HPLC)	USP45 NF40 STP No: STP/RM/10003238-04 Effective Date- 26/06/20
335	CHEMICAL- DRUGS & PHARMACEUTICALS	Levetiracetam USP	General test (Identification by IR)	USP NF- 2022 STP No: STP/RM/10003238-04 Effective Date- 26/06/20
336	CHEMICAL- DRUGS & PHARMACEUTICALS	Levetiracetam USP	General test (Residue on Ignition)	USP NF- 2022 STP No: STP/RM/10003238-04 Effective Date- 26/06/20
337	CHEMICAL- DRUGS & PHARMACEUTICALS	Levetiracetam USP	General test (Water)	USP NF- 2022 STP No: STP/RM/10003238-04 Effective Date- 26/06/20: 2020
338	CHEMICAL- DRUGS & PHARMACEUTICALS	Levetiracetam USP	Related substances (Single impurity) (By HPLC)	USP45 NF40 STP No: STP/RM/10003238-04 Effective Date- 26/06/20
339	CHEMICAL- DRUGS & PHARMACEUTICALS	Levetiracetam USP	Related substances (Total impurities) (By HPLC)	USP45 NF40 STP No: STP/RM/10003238-04 Effective Date- 26/06/20
340	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	Assay (For Levocetirizine) By HPLC	STP NO.: STP/FP/40030665-02 Effective Date-20/07/21
341	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	Assay (For Montelukast) By HPLC	STP NO.: STP/FP/40030665-02 Effective Date-20/07/21
342	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	General Tests for Tablets (Average Weight)	STP/FP/40030665-02 Effective Date-20/07/21
343	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	General Tests for Tablets (Diameter)	STP/FP/40030665-02 Effective Date-20/07/21
344	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	General Tests for Tablets (Disintegration Time)	STP/FP/4030665-02 Effective Date-20/07/21
345	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	General Tests for Tablets (Thickness)	STP/FP/40030665-02 Effective Date-20/07/21
346	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/40030665-02 Effective Date-20/07/21
347	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	Identification By HPLC	STP NO.: STP/FP/40030665-02 Effective Date-20/07/21
348	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	Uniformity of content (For Levocetirizine) By HPLC	STP NO.: STP/FP/40030665-02 Effective Date-20/07/21
349	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine & Montelukast Tablets	Uniformity of content (For Montelukast) By HPLC	STP NO.: STP/FP/40030665-02 Effective Date-20/07/21

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	24 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
350	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine Hydrochloride IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000431-05 Effective Date- 26/06/21
351	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine Hydrochloride IP	Enantiomeric purity (By HPLC)	IP 2018 STP No. STP/RM/10000431-05 Effective Date- 26/06/21
352	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine Hydrochloride IP	General test (Heavy Metals)	General test (Heavy Metals)
353	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine Hydrochloride IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000431-05 Effective Date- 26/06/21
354	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine Hydrochloride IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000431-05 Effective Date- 26/06/21
355	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine Hydrochloride IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000431-05 Effective Date- 26/06/21
356	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine Hydrochloride IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000431-05 Effective Date- 26/06/21
357	CHEMICAL- DRUGS & PHARMACEUTICALS	Levocetirizine Hydrochloride IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000431-05 Effective Date- 26/06/21
358	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Hemihydrate IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000430-07 Effective Date- 11/05/20
359	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Hemihydrate IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000430-07 Effective Date- 11/05/20
360	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Hemihydrate IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000430-07 Effective Date- 11/05/20
361	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Hemihydrate IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000430-07 Effective Date- 11/05/20
362	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Hemihydrate IP	General test (Water)	IP 2018 STP No. STP/RM/10000430-07 Effective Date- 11/05/20
363	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin tablets	Assay By HPLC	IP 2018 STP/FP/40030655 Effective Date-17/03/18

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	25 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
364	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin tablets	Dissolution By UV	IP 2018 STP/FP/40030655 Effective Date-17/03/18
365	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Tablets	General Tests for Tablets (Average Weight)	IP 2018 STP/FP/40030655 Effective Date-17/03/18
366	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Tablets	General Tests for Tablets (Disintegration Time)	IP 2018 STP/FP/40030655 Effective Date-17/03/18
367	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin tablets	General Tests for Tablets (Length)	IP 2018 STP/FP/40030655 Effective Date-17/03/18
368	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Tablets	General Tests for Tablets (Thickness)	IP 2018 STP/FP/40030655 Effective Date-17/03/18
369	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Tablets	General Tests for Tablets (Uniformity of Weight)	IP 2018 STP/FP/40030655 Effective Date-17/03/18
370	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin Tablets	General Tests for Tablets (Width)	IP 2018 STP/FP/40030655 Effective Date-17/03/18
371	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin tablets	Identification By HPLC	IP 2018 STP/FP/40030655 Effective Date-17/03/18
372	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin tablets	Related substances (Single Impurity) By HPLC	IP 2018 STP/FP/40030655 Effective Date-17/03/18
373	CHEMICAL- DRUGS & PHARMACEUTICALS	Levofloxacin tablets	Related substances (Total Impurities) By HPLC	IP 2018 STP/FP/40030655 Effective Date-17/03/18
374	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	Assay (For Hydrochlorothiazide) By HPLC	USP STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
375	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	Assay (For Lisinopril) By HPLC	USP STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
376	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	Dissolution (For Hydrochlorothiazide) By HPLC	USP STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
377	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	Dissolution (For Lisinopril) By HPLC	USP STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
378	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Average Weight)	USP NF-2022 STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
379	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Diameter)	USP NF-2022 STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
380	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Disintegration Time)	USP NF-2022 STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	26 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
381	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Friability)	USP NF-2022 STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
382	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Hardness)	USP NF-2022 STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
383	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Thickness)	USP NF-2022 STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
384	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Uniformity of Weight)	USP NF-2022 STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
385	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	Identification By HPLC	USP STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
386	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	Uniformity of Content (For Hydrochlorothiazide) By HPLC	USP STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
387	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril & Hydrochlorothiazide Tablets USP	Uniformity of Content (For Lisinopril) By HPLC	USP STP/FP/40009179 Issue No- 05 Effective Date- 01/06/19
388	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10000433-05 Effective Date- 11/05/20
389	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000433-05 Effective Date- 11/05/20
390	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000433-05 Effective Date- 11/05/20
391	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000433-05 Effective Date- 11/05/20:
392	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril IP	General test (Water)	IP 2018 STP No. STP/RM/10000433-05 Effective Date- 11/05/20
393	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000433-05 Effective Date- 11/05/20
394	CHEMICAL- DRUGS & PHARMACEUTICALS	Lisinopril IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000433-05 Effective Date- 11/05/20
395	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	Assay By HPLC	USP STP/FP/40033271-00 Effective Date-06/10/18

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	27 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
396	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	Dissolution By UV	USP STP/FP/40033271-00 Effective Date-06/10/18
397	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	General Tests for Tablets (Average Weight)	USP NF-2022 STP/FP/40033271-00 Effective Date-06/10/18
398	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	General Tests for Tablets (Diameter)	USP NF-2022 STP/FP/40033271-00 Effective Date-06/10/18
399	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	General Tests for Tablets (Disintegration Time)	USP NF-2022 STP/FP/40033271-00 Effective Date-06/10/18
400	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	General Tests for Tablets (Friability)	USP NF-2022 STP/FP/40033271-00 Effective Date-06/10/18
401	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	General Tests for Tablets (Hardness)	USP NF-2022 STP/FP/40033271-00 Effective Date-06/10/18
402	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	General Tests for Tablets (Thickness)	USP NF-2022 STP/FP/40033271-00 Effective Date-06/10/18
403	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	General Tests for Tablets (Uniformity of Weight)	USP NF-2022 STP/FP/40033271-00 Effective Date-06/10/18
404	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine Tablets USP	Identification By HPLC	USP STP/FP/40033271-00 Effective Date-06/10/18
405	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine USP	Assay (By HPLC)	USP45 NF40 STP No. STP/RM/10000457-04 Effective Date- 21/01/21
406	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine USP	General test (Identification by IR)	USP NF- 2022STP No. STP/RM/10000457-04 Effective Date- 21/01/21
407	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine USP	General test (Loss on Drying)	USP NF- 2022STP No. STP/RM/10000457-04 Effective Date- 21/01/21
408	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine USP	General test (Residue on Ignition)	USP NF- 2022STP No. STP/RM/10000457-04 Effective Date- 21/01/21
409	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine USP	Related compounds (Single impurity) (By HPLC)	USP45 NF40 STP No. STP/RM/10000457-04 Effective Date- 21/01/21

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	28 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
410	CHEMICAL- DRUGS & PHARMACEUTICALS	Loratadine USP	Related compounds (Total impurities) (By HPLC)	USP45 NF40 STP No. STP/RM/10000457-04 Effective Date- 21/01/21
411	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	Assay (For Hydrochlorothiazide) By HPLC	STP/FP/40021295-02 Effective Date- 29/05/21
412	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	Assay (For Losartan potassium) By HPLC	STP/FP/40021295-02 Effective Date- 29/05/21
413	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	Dissolution (For Hydrochlorothiazide) By HPLC	STP/FP/40021295-02 Effective Date- 29/05/21
414	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	Dissolution (For Losartan potassium:) By HPLC	STP/FP/40021295-02 Effective Date- 29/05/21
415	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Average Weight)	STP/FP/40021295-02 Effective Date- 29/05/21
416	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Disintegration Time)	STP/FP/40021295-02 Effective Date- 29/05/21
417	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Thickness)	STP/FP/40021295-02 Effective Date- 29/05/21
418	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	General Tests for Tablets (Width)	STP/FP/40021295-02 Effective Date- 29/05/21
419	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	Identification	STP/FP/40021295-02 Effective Date- 29/05/21
420	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	Uniformity of Content (For Hydrochlorothiazide) By HPLC	STP/FP/40021295-02 Effective Date- 29/05/21
421	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium & Hydrochlorothiazide Tablets USP	Uniformity of Content (For Losartan potassium) By HPLC	STP/FP/40021295-02 Effective Date- 29/05/21
422	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10000458-03 Effective Date- 13/05/20
423	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000458-03 Effective Date- 13/05/20
424	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000458-03 Effective Date- 13/05/20
425	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium IP	General test (Water)	IP 2018 STP No. STP/RM/10000458-03 Effective Date- 13/05/20
426	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000458-03 Effective Date- 13/05/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	29 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
427	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000458-03 Effective Date- 13/05/20
428	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan potassium tablets	Assay By HPLC	STP/FP/40037191-00 Effective Date- 20/09/19
429	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium Tablets	General Tests for Tablets (Average Weight)	STP/FP/40037191-00 Effective Date- 20/09/19
430	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan potassium tablets	General Tests for Tablets (Diameter)	STP/FP/40037191-00 Effective Date- 20/09/19
431	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium Tablets	General Tests for Tablets (Disintegration Time)	STP/FP/40037191-00 Effective Date- 20/09/19
432	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium Tablets	General Tests for Tablets (Thickness)	STP/FP/40037191-00 Effective Date- 20/09/19
433	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan Potassium Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/40037191-00 Effective Date- 20/09/19
434	CHEMICAL- DRUGS & PHARMACEUTICALS	Losartan potassium tablets	Related substances By HPLC	STP/FP/40037191-00 Effective Date- 20/09/19
435	CHEMICAL- DRUGS & PHARMACEUTICALS	Magnesium Hydroxide IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000509-04 Effective Date- 25/01/19
436	CHEMICAL- DRUGS & PHARMACEUTICALS	Magnesium Hydroxide IP	General test (Arsenic)	IP 2018 STP No. STP/RM/10000509-04 Effective Date- 25/01/19
437	CHEMICAL- DRUGS & PHARMACEUTICALS	Magnesium Hydroxide IP	General test (Chloride)	IP 2018 STP No. STP/RM/10000509-04 Effective Date- 25/01/19
438	CHEMICAL- DRUGS & PHARMACEUTICALS	Magnesium Hydroxide IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000509-04 Effective Date- 25/01/19
439	CHEMICAL- DRUGS & PHARMACEUTICALS	Magnesium Hydroxide IP	General test (Iron)	IP 2018 STP No. STP/RM/10000509-04 Effective Date- 25/01/19
440	CHEMICAL- DRUGS & PHARMACEUTICALS	Magnesium Hydroxide IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000509-04 Effective Date- 25/01/19
441	CHEMICAL- DRUGS & PHARMACEUTICALS	Magnesium Hydroxide IP	General test (Sulphates)	IP 2018 STP No. STP/RM/10000509-04 Effective Date- 25/01/19

SCOPE OF ACCREDITATION

Laboratory Name : AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA

Accreditation Standard ISO/IEC 17025:2017

Certificate Number TC-6742 **Page No** 30 of 43

Validity 29/12/2022 to 28/12/2024 **Last Amended on** 24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
442	CHEMICAL- DRUGS & PHARMACEUTICALS	Mefenamic Acid IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000506-05 Effective Date- 10/01/19
443	CHEMICAL- DRUGS & PHARMACEUTICALS	Mefenamic Acid IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000506-05 Effective Date- 10/01/19
444	CHEMICAL- DRUGS & PHARMACEUTICALS	Mefenamic Acid IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000506-05 Effective Date- 10/01/19
445	CHEMICAL- DRUGS & PHARMACEUTICALS	Mefenamic Acid IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000506-05 Effective Date- 10/01/19
446	CHEMICAL- DRUGS & PHARMACEUTICALS	Meloxicam BP	Assay (By Titration)	BP 2018 STP No. STP/RM/10000520-03 Effective Date- 24/09/18
447	CHEMICAL- DRUGS & PHARMACEUTICALS	Meloxicam BP	General test (Identification by IR)	BP 2022 STP No.STP/RM/10000520-03 Effective Date- 24/09/18
448	CHEMICAL- DRUGS & PHARMACEUTICALS	Meloxicam BP	General test (Loss on Drying)	BP 2022 STP No. STP/RM/10000520-03 Effective Date- 24/09/18
449	CHEMICAL- DRUGS & PHARMACEUTICALS	Meloxicam BP	General test (Sulfated Ash)	BP 2022 STP No. STP/RM/10000520-03 Effective Date- 24/09/18
450	CHEMICAL- DRUGS & PHARMACEUTICALS	Meloxicam BP	Related substances (Total impurities) (By HPLC)	BP 2018 STP No. STP/RM/10000520-03 Effective Date- 24/09/18
451	CHEMICAL- DRUGS & PHARMACEUTICALS	Meloxicam BP	Related substances (Single impurity) (By HPLC)	BP 2018 STP No. STP/RM/10000520-03 Effective Date- 24/09/18
452	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000521-06 Effective Date- 24/02/22
453	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000521-06 Effective Date- 24/02/22
454	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000521-06 Effective Date- 24/02/22
455	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000521-06 Effective Date- 24/02/22

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	31 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
456	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000521-06 Effective Date- 24/02/22
457	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin hydrochloride sustained-release tablets	Assay By UV	IP 2018 STP/FP/40037008-02 Effective Date- 09/03/22
458	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin hydrochloride sustained-release tablets	Dissolution By UV	IP 2018 STP/FP/40037008-02 Effective Date- 09/03/22
459	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride Sustained-Release Tablets	General Tests for Tablets (Average Weight)	IP 2018 STP/FP/40037008-02 Effective Date- 09/03/22
460	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin hydrochloride sustained-release tablets	General Tests for Tablets (Diameter)	IP 2018 STP/FP/40037008-02 Effective Date- 09/03/22
461	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride Sustained-Release Tablets	General Tests for Tablets (Friability)	IP 2018 STP/FP/40037008-02 Effective Date- 09/03/22
462	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride Sustained-Release Tablets	General Tests for Tablets (Hardness)	IP 2018 STP/FP/40037008-02 Effective Date- 09/03/22
463	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride Sustained-Release Tablets	General Tests for Tablets (Thickness)	IP 2018 STP/FP/40037008-02 Effective Date- 09/03/22
464	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin Hydrochloride Sustained-Release Tablets	General Tests for Tablets (Uniformity of Weight)	IP 2018 STP/FP/40037008-02 Effective Date- 09/03/22
465	CHEMICAL- DRUGS & PHARMACEUTICALS	Metformin hydrochloride sustained-release tablets	Identification By UV	IP 2018 STP/FP/40037008-02 Effective Date- 09/03/22
466	CHEMICAL- DRUGS & PHARMACEUTICALS	Methylcobalamin IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10011475-01 Effective Date- 12/06/18
467	CHEMICAL- DRUGS & PHARMACEUTICALS	Methylcobalamin IP	General test (Identification By UV)	IP 2018 STP No. STP/RM/10011475-01 Effective Date- 12/06/18
468	CHEMICAL- DRUGS & PHARMACEUTICALS	Methylcobalamin IP	General test (Water)	IP 2018 STP No. STP/RM/10011475-01 Effective Date- 12/06/18
469	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole IP	Assay (By Titration)	IP 2018 STP/RM/150002484-02 Effective Date- 29/04/20
470	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole IP	General test (Heavy Metals)	IP 2018 STP/RM/150002484-02 Effective Date- 29/04/20: 2020
471	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole IP	General test (Identification By IR)	IP 2018 STP/RM/150002484-02 Effective Date- 29/04/20
472	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole IP	General test (Loss on Drying)	IP 2018 STP/RM/150002484-02 Effective Date- 29/04/20

SCOPE OF ACCREDITATION

Laboratory Name : AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA

Accreditation Standard ISO/IEC 17025:2017

Certificate Number TC-6742 **Page No** 32 of 43

Validity 29/12/2022 to 28/12/2024 **Last Amended on** 24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
473	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole IP	General test (Sulphated Ash)	IP 2018 STP/RM/150002484-02 Effective Date- 29/04/20
474	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole Tablet	Assay By Titration	IP 2018 STP/FP/ 190002214 Effective Date- 23/11/18
475	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole Tablet	Dissolution By UV	IP 2018 STP/FP/ 190002214 Effective Date- 23/11/18
476	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole Tablet	General Tests for Tablets (Diameter)	IP 2018 STP/FP/ 190002214 Effective Date- 23/11/18
477	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole Tablet	Identification By IR	IP 2018 STP/FP/ 190002214 Effective Date- 23/11/18
478	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole Tablet IP	General Tests for Tablets (Average Weight)	IP 2018 STP/FP/ 190002214 Effective Date- 23/11/18
479	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole Tablet IP	General Tests for Tablets (Disintegration Time)	IP 2018 STP/FP/ 190002214 Effective Date- 23/11/18
480	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole Tablet IP	General Tests for Tablets (Thickness)	IP 2018 STP/FP/ 190002214 Effective Date- 23/11/18
481	CHEMICAL- DRUGS & PHARMACEUTICALS	Metronidazole Tablet IP	General Tests for Tablets (Uniformity of Weight)	IP 2018 STP/FP/ 190002214 Effective Date- 23/11/18
482	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl BP	Assay (By HPLC)	BP2022 STP No, STP/RM/10000556-03 Effective Date- 21/08/18
483	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl BP	General test (Identification By IR)	BP2022 STP No, STP/RM/10000556-03 Effective Date- 21/08/18
484	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl BP	General test (Sulfated Ash)	BP2022 STP No, STP/RM/10000556-03 Effective Date- 21/08/18
485	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl BP	General test (Water)	BP2022 STP No, STP/RM/10000556-03 Effective Date- 21/08/18
486	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl Tablets	Assay By HPLC	STP/FP/40031547-00 Effective Date- 20/04/18
487	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl Tablets	General Tests for Tablets (Average Weight)	General Tests for Tablets (Average Weight)
488	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl Tablets	General Tests for Tablets (Disintegration Time)	STP/FP/40031547-00 Effective Date- 20/04/18
489	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl Tablets	General Tests for Tablets (Length)	STP/FP/40031547-00 Effective Date- 20/04/18

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	33 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
490	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl Tablets	General Tests for Tablets (Thickness)	STP/FP/40031547-00 Effective Date- 20/04/18
491	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/40031547-00 Effective Date- 20/04/18
492	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl Tablets	General Tests for Tablets (Width)	STP/FP/40031547-00 Effective Date- 20/04/18
493	CHEMICAL- DRUGS & PHARMACEUTICALS	Moxifloxacin HCl Tablets	Identification By HPLC	STP/FP/40031547-00 Effective Date- 20/04/18
494	CHEMICAL- DRUGS & PHARMACEUTICALS	Ofloxacin IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000632-05 Effective Date- 23/05/20
495	CHEMICAL- DRUGS & PHARMACEUTICALS	Ofloxacin IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000632-05 Effective Date- 23/05/20
496	CHEMICAL- DRUGS & PHARMACEUTICALS	Ofloxacin IP	General test (Identification By IR)	IP 2018 STP No. STP/RM/10000632-05 Effective Date- 23/05/20
497	CHEMICAL- DRUGS & PHARMACEUTICALS	Ofloxacin IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000632-05 Effective Date- 23/05/20
498	CHEMICAL- DRUGS & PHARMACEUTICALS	Ofloxacin IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000632-05 Effective Date- 23/05/20
499	CHEMICAL- DRUGS & PHARMACEUTICALS	Ofloxacin IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000632-05 Effective Date- 23/05/20
500	CHEMICAL- DRUGS & PHARMACEUTICALS	Ofloxacin IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000632-05 Effective Date- 23/05/20
501	CHEMICAL- DRUGS & PHARMACEUTICALS	Olanzapine IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000607-04 Effective Date- 23/05/20
502	CHEMICAL- DRUGS & PHARMACEUTICALS	Olanzapine IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000607-04 Effective Date- 23/05/20
503	CHEMICAL- DRUGS & PHARMACEUTICALS	Olanzapine IP	General test (Identification By IR)	IP 2018 STP No. STP/RM/10000607-04 Effective Date- 23/05/20
504	CHEMICAL- DRUGS & PHARMACEUTICALS	Olanzapine IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000607-04 Effective Date- 23/05/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	34 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
505	CHEMICAL- DRUGS & PHARMACEUTICALS	Olanzapine IP	General test (Water)	IP 2018 STP No. STP/RM/10000607-04 Effective Date- 23/05/20
506	CHEMICAL- DRUGS & PHARMACEUTICALS	Olanzapine IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000607-04 Effective Date- 23/05/20
507	CHEMICAL- DRUGS & PHARMACEUTICALS	Olanzapine IP	Related substances (By HPLC) Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000607-04 Effective Date- 23/05/20
508	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole and Domperidone Capsule	Assay (For Domperidone) By HPLC	STP/FP/40001638-05 Effective Date- 03/08/19
509	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole and Domperidone Capsule	Assay (For Omeprazole) By HPLC	STP/FP/40001638-05 Effective Date- 03/08/19
510	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole and Domperidone Capsule	General Tests for Capsule: (Average Fill)	STP/FP/40001638-05 Effective Date- 03/08/19
511	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole and Domperidone Capsule	General Tests for Capsule: (Disintegration Test)	General Tests for Capsule: (Disintegration Test)
512	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole and Domperidone Capsule	General Tests for Capsule: (Uniformity of Weight)	STP/FP/40001638-05 Effective Date- 03/08/19
513	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole and Domperidone Capsule	Identification By HPLC	STP/FP/40001638-05 Effective Date- 03/08/19
514	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole Capsule IP	Assay By HPLC	IP 2018 STP/FP/ 40030670-01 Effective Date: 15/11/2019
515	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole Capsule IP	General Tests for Capsule: (Average Fill)	IP 2018 STP/FP/ 40030670-01 Effective Date: 15/11/2019
516	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole Capsule IP	General Tests for Capsule: (Disintegration Test)	IP 2018 STP/FP/ 40030670-01 Effective Date: 15/11/2019
517	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole Capsule IP	General Tests for Capsule: (Uniformity of Weight)	IP 2018 STP/FP/ 40030670-01 Effective Date: 15/11/2019
518	CHEMICAL- DRUGS & PHARMACEUTICALS	Omeprazole Capsule IP	Identification	IP 2018 STP/FP/ 40030670-01 Effective Date: 15/11/2019
519	CHEMICAL- DRUGS & PHARMACEUTICALS	Ondansterone Hydrochloride IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10000617-05 Effective Date- 05/04/19
520	CHEMICAL- DRUGS & PHARMACEUTICALS	Ondansterone Hydrochloride IP	General test (Identification By IR)	IP 2018 STP No. STP/RM/10000617-05 Effective Date- 05/04/19
521	CHEMICAL- DRUGS & PHARMACEUTICALS	Ondansterone Hydrochloride IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000617-05 Effective Date- 05/04/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	35 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
522	CHEMICAL- DRUGS & PHARMACEUTICALS	Ondansterone Hydrochloride IP	General test (Water)	IP 2018 STP No. STP/RM/10000617-05 Effective Date- 05/04/19
523	CHEMICAL- DRUGS & PHARMACEUTICALS	Ondansterone Hydrochloride IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000617-05 Effective Date- 05/04/19
524	CHEMICAL- DRUGS & PHARMACEUTICALS	Ondansterone Hydrochloride IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000617-05 Effective Date- 05/04/19
525	CHEMICAL- DRUGS & PHARMACEUTICALS	Paracetamol IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000634-07 Effective Date- 02/05/20
526	CHEMICAL- DRUGS & PHARMACEUTICALS	Paracetamol IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000634-07 Effective Date- 02/05/20
527	CHEMICAL- DRUGS & PHARMACEUTICALS	Paracetamol IP	General test (Identification By IR)	IP 2018 STP No. STP/RM/10000634-07 Effective Date- 02/05/20
528	CHEMICAL- DRUGS & PHARMACEUTICALS	Paracetamol IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000634-07 Effective Date- 02/05/20
529	CHEMICAL- DRUGS & PHARMACEUTICALS	Paracetamol IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000634-07 Effective Date- 02/05/20
530	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam Capsule	Assay By HPLC	STP/FP/40037200-00 Effective Date- 13/09/19
531	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam Capsule	Dissolution By UV	STP/FP/40037200-00 Effective Date- 13/09/19
532	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam Capsule	General Tests for Capsule: (Disintegration Test)	STP/FP/40047740-00 Effective Date-15/08/21
533	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam Capsule	Identification By TLC	STP/FP/40037200-00 Effective Date- 13/09/19
534	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam Capsule	Uniformity of Dosage units By HPLC	STP/FP/40037200-00 Effective Date- 13/09/19
535	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10000643-04 Effective Date- 29/05/20
536	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000643-04 Effective Date- 29/05/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	36 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
537	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam IP	General test (Identification By IR)	IP 2018 STP No. STP/RM/10000643-04 Effective Date- 29/05/20
538	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000643-04 Effective Date- 29/05/20
539	CHEMICAL- DRUGS & PHARMACEUTICALS	Piroxicam IP	General test (Water)	IP 2018 STP No. STP/RM/10000643-04 Effective Date- 29/05/20
540	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
541	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	General test (Iodides)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
542	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	General test (Arsenic)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
543	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	General test (Barium)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
544	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	General test (Bromides)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
545	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	General test (Calcium & Magnesium)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
546	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
547	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	General test (Iron)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
548	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
549	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Chloride IP	General test (Sulphates)	IP 2018 STP No. STP/RM/10000697-05 Effective Date- 30/06/20
550	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Clavulanate Diluted IP	Assay (By HPLC)	IP2018 STP No.- STP/RM/10003108-02 Effective Date- 31/08/18

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	37 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
551	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Clavulanate Diluted IP	General test (Identification by HPLC)	IP2018 STP No.- STP/RM/10003108-02 Effective Date- 31/08/18
552	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Clavulanate Diluted IP	General test (pH)	IP2018 STP No.- STP/RM/10003108-02 Effective Date- 31/08/18
553	CHEMICAL- DRUGS & PHARMACEUTICALS	Potassium Clavulanate Diluted IP	General test (Water)	IP2018 STP No.- STP/RM/10003108-02 Effective Date- 31/08/18
554	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin	Assay	By HPLC STP/FP/40047740-00 Effective Date-15/08/21
555	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin	General Tests for Capsule	STP/FP/40047740-00 Effective Date-15/08/21
556	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin	Identification	As per Assay STP/FP/40047740-00 Effective Date-15/08/21
557	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin Capsule	General Tests for Capsule: (Average Fill)	General Tests for Capsule: (Average Fill)
558	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin Capsule	General Tests for Capsule: (Disintegration Test)	General Tests for Capsule: (Disintegration Test)
559	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin Capsule	General Tests for Capsule: (Uniformity of Weight)	STP/FP/40047740-00 Effective Date-15/08/21
560	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin Capsule	Identification	STP/FP/40047740-00 Effective Date-15/08/21
561	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10009203-02 Effective Date- 15/06/20
562	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10009203-02 Effective Date- 15/06/20
563	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10009203-02 Effective Date- 15/06/20
564	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10009203-02 Effective Date- 15/06/20
565	CHEMICAL- DRUGS & PHARMACEUTICALS	Pregabalin IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10009203-02 Effective Date- 15/06/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	38 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
566	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
567	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	General test (Chlorides)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
568	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
569	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
570	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	General test (Iron)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
571	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
572	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
573	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	General test (Sulphates)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
574	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
575	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	Related substances (Impurity A) (By HPLC)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
576	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyrantel Pamoate IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000690-05 Effective Date- 31/03/20
577	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyridoxine hydrochloride IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000876-04 Effective Date- 31/03/19
578	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyridoxine hydrochloride IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000876-04 Effective Date- 31/03/19:
579	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyridoxine hydrochloride IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000876-04 Effective Date- 31/03/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	39 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
580	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyridoxine hydrochloride IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000876-04 Effective Date- 31/03/19
581	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyridoxine hydrochloride IP	General test (pH)	IP 2018 STP No. STP/RM/10000876-04 Effective Date- 31/03/19
582	CHEMICAL- DRUGS & PHARMACEUTICALS	Pyridoxine hydrochloride IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000876-04 Effective Date- 31/03/19
583	CHEMICAL- DRUGS & PHARMACEUTICALS	Rabeprazole Sodium IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10000731-06 Effective Date- 16/02/22
584	CHEMICAL- DRUGS & PHARMACEUTICALS	Rabeprazole Sodium IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000731-06 Effective Date- 16/02/22
585	CHEMICAL- DRUGS & PHARMACEUTICALS	Rabeprazole Sodium IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000731-06 Effective Date- 16/02/22
586	CHEMICAL- DRUGS & PHARMACEUTICALS	Rabeprazole Sodium IP	General test (Water)	IP 2018 STP No. STP/RM/10000731-06 Effective Date- 16/02/22
587	CHEMICAL- DRUGS & PHARMACEUTICALS	Rabeprazole Sodium IP	Related substances (Single impurity) (By HPLC)	IP 2018 STP No. STP/RM/10000731-06 Effective Date- 16/02/22
588	CHEMICAL- DRUGS & PHARMACEUTICALS	Rabeprazole Sodium IP	Related substances (Total impurities) (By HPLC)	IP 2018 STP No. STP/RM/10000731-06 Effective Date- 16/02/22
589	CHEMICAL- DRUGS & PHARMACEUTICALS	Ranitidine hydrochloride IP	Assay (By HPLC)	IP 2018 STP No. STP/RM/10000728-04 Effective Date- 22/01/19
590	CHEMICAL- DRUGS & PHARMACEUTICALS	Ranitidine hydrochloride IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000728-04 Effective Date- 22/01/19
591	CHEMICAL- DRUGS & PHARMACEUTICALS	Ranitidine hydrochloride IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000728-04 Effective Date- 22/01/19
592	CHEMICAL- DRUGS & PHARMACEUTICALS	Ranitidine hydrochloride IP	General test (pH)	IP 2018 STP No. STP/RM/10000728-04 Effective Date- 22/01/19
593	CHEMICAL- DRUGS & PHARMACEUTICALS	Ranitidine hydrochloride IP	General test (Sulphated Ash)	IP 2018 STP No. STP/RM/10000728-04 Effective Date- 22/01/19

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	40 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
594	CHEMICAL- DRUGS & PHARMACEUTICALS	Risperidone BP	Assay (By Titration)	BP 2022 STP No. STP/RM/150000433-03 Effective Date- 22/04/18
595	CHEMICAL- DRUGS & PHARMACEUTICALS	Risperidone BP	General test (Identification by IR)	BP 2022 STP No. STP/RM/150000433-03 Effective Date- 22/04/18
596	CHEMICAL- DRUGS & PHARMACEUTICALS	Risperidone BP	General test (Loss on Drying)	BP 2022 STP No. STP/RM/150000433-03 Effective Date- 22/04/18
597	CHEMICAL- DRUGS & PHARMACEUTICALS	Risperidone BP	General test (Sulfated Ash)	BP 2022 STP No. STP/RM/150000433-03 Effective Date- 22/04/18
598	CHEMICAL- DRUGS & PHARMACEUTICALS	Risperidone BP	Related substances (Single impurity) (By HPLC)	BP 2022 STP No. STP/RM/150000433-03 Effective Date- 22/04/18
599	CHEMICAL- DRUGS & PHARMACEUTICALS	Risperidone BP	Related substances (Total impurities) (By HPLC)	BP 2022 STP No. STP/RM/150000433-03 Effective Date- 22/04/18
600	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Ascorbate IP	Assay (By Titration)	IP 2018 STP No. STP/RM/150000388-03 Effective Date- 07/05/20
601	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Ascorbate IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/150000388-03 Effective Date- 07/05/20
602	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Ascorbate IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/150000388-03 Effective Date- 07/05/20
603	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Ascorbate IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/150000388-03 Effective Date- 07/05/20
604	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Ascorbate IP	General test (pH)	IP 2018 STP No. STP/RM/150000388-03 Effective Date- 07/05/20
605	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Valproate IP	Assay (By Titration)	IP 2018 STP No. STP/RM/10000784-04 Effective Date- 26/06/20
606	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Valproate IP	General test (Chlorides)	IP 2018 STP No. STP/RM/10000784-04 Effective Date- 26/06/20
607	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Valproate IP	General test (Heavy Metals)	IP 2018 STP No. STP/RM/10000784-04 Effective Date- 26/06/20

SCOPE OF ACCREDITATION

Laboratory Name : AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA

Accreditation Standard ISO/IEC 17025:2017

Certificate Number TC-6742 **Page No** 41 of 43

Validity 29/12/2022 to 28/12/2024 **Last Amended on** 24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
608	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Valproate IP	General test (Identification by IR)	IP 2018 STP No. STP/RM/10000784-04 Effective Date- 26/06/20
609	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Valproate IP	General test (Loss on Drying)	IP 2018 STP No. STP/RM/10000784-04 Effective Date- 26/06/20
610	CHEMICAL- DRUGS & PHARMACEUTICALS	Sodium Valproate IP	General test (Sulphates)	IP 2018 STP No. STP/RM/10000784-04 Effective Date- 26/06/20
611	CHEMICAL- DRUGS & PHARMACEUTICALS	Tadalafil	Assay (By HPLC)	STP No. STP/RM/10000842-03 Effective Date- 29/12/18
612	CHEMICAL- DRUGS & PHARMACEUTICALS	Tadalafil	General test (Heavy Metals)	STP No. STP/RM/10000842-03 Effective Date- 29/12/18
613	CHEMICAL- DRUGS & PHARMACEUTICALS	Tadalafil	General test (Identification by IR)	STP No. STP/RM/10000842-03 Effective Date- 29/12/18
614	CHEMICAL- DRUGS & PHARMACEUTICALS	Tadalafil	General test (Loss on Drying)	TP No. STP/R/10000842-03 Effective Date- 29/12/18
615	CHEMICAL- DRUGS & PHARMACEUTICALS	Tadalafil	General test (Sulphated Ash)	STP No. STP/RM/10000842-03 Effective Date- 29/12/18
616	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	Assay (For Hydrochlorothiazide) By HPLC	STP/FP/40030617-01 Effective Date-11/02/20
617	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	Assay (For Telmisartan) By HPLC	STP/FP/40030617-01 Effective Date-11/02/20
618	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	General Tests for Tablets (Average Weight)	STP/FP/40030617-01 Effective Date-11/02/20
619	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	General Tests for Tablets (Friability)	STP/FP/40030617-01 Effective Date-11/02/20
620	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	General Tests for Tablets (Hardness)	STP/FP/40030617-01 Effective Date-11/02/20
621	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	General Tests for Tablets (Length)	STP/FP/40030617-01 Effective Date-11/02/20
622	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	General Tests for Tablets (Thickness)	STP/FP/40030617-01 Effective Date-11/02/20
623	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/40030617-01 Effective Date-11/02/20
624	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	General Tests for Tablets (Width)	STP/FP/40030617-01 Effective Date-11/02/20
625	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan & Hydrochlorothiazide Tablets	Identification By HPLC	STP/FP/40030617-01 Effective Date-11/02/20

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA
Accreditation Standard	ISO/IEC 17025:2017
Certificate Number	TC-6742
Page No	42 of 43
Validity	29/12/2022 to 28/12/2024
Last Amended on	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
626	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	Assay By HPLC	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
627	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	Dissolution By HPLC	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
628	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	General Tests for Tablets (Average Weight)	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
629	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	General Tests for Tablets (Diameter)	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
630	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	General Tests for Tablets (Disintegration Time)	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
631	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	General Tests for Tablets (Friability)	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
632	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	General Tests for Tablets (Hardness)	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
633	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	General Tests for Tablets (Thickness)	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
634	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	General Tests for Tablets (Uniformity of Weight)	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
635	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	Identification By HPLC	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
636	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	Related Substances (Single Impurity) By HPLC	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
637	CHEMICAL- DRUGS & PHARMACEUTICALS	Telmisartan Tablets IP	Related Substances (Total impurities) By HPLC	IP 2018 STP/FP/40030631 Issue No- 01 Effective Date- 02/11/19
638	CHEMICAL- DRUGS & PHARMACEUTICALS	Tetracycline capsule IP	Assay By HPLC	IP 2018 STS/FP/40035540-00 Effective Date-04/04/2019
639	CHEMICAL- DRUGS & PHARMACEUTICALS	Tetracycline capsule IP	Dissolution By -UV	IP 2018 STS/FP/40035540-00 Effective Date-04/04/2019
640	CHEMICAL- DRUGS & PHARMACEUTICALS	Tetracycline capsule IP	General Tests for Capsule: (Average Fill)	IP 2018 STS/FP/40035540-00 Effective Date-04/04/2019
641	CHEMICAL- DRUGS & PHARMACEUTICALS	Tetracycline capsule IP	General Tests for Capsule: (Disintegration Test)	IP 2018 STS/FP/40035540-00 Effective Date-04/04/2019
642	CHEMICAL- DRUGS & PHARMACEUTICALS	Tetracycline capsule IP	General Tests for Capsule: (Uniformity of Weight)	IP 2018 STS/FP/40035540-00 Effective Date-04/04/2019
643	CHEMICAL- DRUGS & PHARMACEUTICALS	Tetracycline capsule IP	Identification	STS/FP/40035540-00 Effective Date-04/04/2019
644	CHEMICAL- DRUGS & PHARMACEUTICALS	Thickness	General Tests for Tablets (Average Weight)	STP/FP/40032091-00: 2019

National Accreditation Board for Testing and Calibration Laboratories

SCOPE OF ACCREDITATION

Laboratory Name :	AKUMS DRUGS AND PHARMACEUTICALS LTD. (PLANT1), QUALITY CONTROL LABORATORY, 19-20-21, SECTOR-6A, I.I.E., SIDCUL, RANIPUR, HARIDWAR, UTTARAKHAND, INDIA		
Accreditation Standard	ISO/IEC 17025:2017		
Certificate Number	TC-6742	**Page No**	43 of 43
Validity	29/12/2022 to 28/12/2024	**Last Amended on**	24/01/2023

S.No	Discipline / Group	Materials or Products tested	Component, parameter or characteristic tested / Specific Test Performed / Tests or type of tests performed	Test Method Specification against which tests are performed and / or the techniques / equipment used
645	CHEMICAL- DRUGS & PHARMACEUTICALS	Vitamin C (Ascorbic acid)Tablets	Assay By Titration	STP/FP/190002364-01 Effective Date-14/08/20
646	CHEMICAL- DRUGS & PHARMACEUTICALS	Vitamin C (Ascorbic acid)Tablets	General Tests for Tablets (Average Weight)	STP/FP/190002364-01 Effective Date-14/08/20
647	CHEMICAL- DRUGS & PHARMACEUTICALS	Vitamin C (Ascorbic acid)Tablets	General Tests for Tablets (Diameter)	STP/FP/190002364-01 Effective Date-14/08/20
648	CHEMICAL- DRUGS & PHARMACEUTICALS	Vitamin C (Ascorbic acid)Tablets	General Tests for Tablets (Friability)	STP/FP/190002364-01 Effective Date-14/08/20
649	CHEMICAL- DRUGS & PHARMACEUTICALS	Vitamin C (Ascorbic acid)Tablets	General Tests for Tablets (Hardness)	STP/FP/190002364-01 Effective Date-14/08/20
650	CHEMICAL- DRUGS & PHARMACEUTICALS	Vitamin C (Ascorbic acid)Tablets	General Tests for Tablets (Thickness)	STP/FP/190002364-01 Effective Date-14/08/20
651	CHEMICAL- DRUGS & PHARMACEUTICALS	Vitamin C (Ascorbic acid)Tablets	General Tests for Tablets (Uniformity of Weight)	STP/FP/190002364-01 Effective Date-14/08/20
652	CHEMICAL- DRUGS & PHARMACEUTICALS	Vitamin C (Ascorbic acid)Tablets	Identification By Chemical	STP/FP/190002364-01 Effective Date-14/08/20