

Office of The Commissioner, Food & Drugs Administration M.S. Bandra – Kurla Complex, Bandra (E), Mumbai – 400 051 Date :-29 Mar 2023

## CERTIFICATE OF GOOD MANUFACTURING PRACTICES

This Certificate conforms to the format recommended by the World Health Organization. (General instructions and explanatory notes attached).

Certificate No.: NEW-WHO-GMP/CERT/AD/122144/2023/11/44632

On the basis of the inspection carried out on 23/01/2023 & 24/01/2023 ,we certify that the site indicated on this Certificate complies with **Good Manufacturing Practices** for the dosage forms, categories and activities listed in Table 1.

1.	Name of the Firm	:	LUPIN LIMITED
	Address	:	A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA
2.	Licence No.	;	633 In Form 25, 499 In Form 28

## Table 1

Sr.No.	Dosage Form(s)	Categor(ies)	Activity(ies)
1	Capsules	General ( Other than Cephalosporins, Penicillin, Cytotoxic, Hormones )	Production, Filling, Packing, labelling, Quality Control, Quality Assurance
2	Liquid Orals	General ( Other than Cephalosporins, Penicillin, Cytotoxic, Hormones )	Production, Filling, Packing, labelling, Quality Control, Quality Assurance
3	Tablets	General ( Other than Cephalosporins, Penicillin, Cytotoxic, Hormones )	Production, Filling, Packing, labelling, Quality Control, Quality Assurance

The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.

This certificate remains valid until 28 Mar 2026. It becomes invalid if the activities and / or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.



Name	certificate : ] of Manufacturing Firm : ] License No : 0	RODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Product	Composition
1	Amlodipine Besylate Tablets 10 mg	Each Tablet Contains Amlodipine Besylate USP equivalent to Amlodipine (In House) 10 mg
2	Amlodipine Besylate Tablets 5 mg	Each Tablet Contains Amlodipine Besylate USP equivalent to Amlodipine (In House) 5 mg
3	Cycloserine Capsules USP	Each Capsule Contains Cycloserine USP 250.0 mg Colour:Approved colours used in capsule shell
4	Frusemide and Spironolactone Tablets	Each film coated tablet contains Frusemide IP 20 mg Spironolactone IP 50 mg Colour:Quinoline Yellow WS, Brilliant Blue FCF & Titanium Dioxide IP,
5	Hydroxychloroquine Sulfate Tablets 200mg USP	Each film coated tablet contains Hydroxychloroquine Sulfate USP 200 mg
6	Rifampicin, Isoniazid and Ethambutol Hydrochloride Tablets IP	Each film coated tablet contains Rifampicin IP 150.0 mg Isoniazid IP 75.0 mg Ethambutol Hydrochloride IP 275.0 mg Colour:Red Oxide of Iron and Titanium Dioxide
7	Rifampicin,Isoniazid, Pyrazinamide and Ethambutol Hydrochloride Tablets IP	Each film coated tablet contains Rifampicin IP 150.0 mg Isoniazid IP 75.0 mg Pyrazinamide IP 400.0 mg Ethambutol Hydrochloride IP 275.0 mg Colour:Red Oxide of Iron and Titanium Dioxide IP
8	Rifampin and Isoniazid Dispersible Tablets 75 mg/ 50 mg	Each uncoated Dispersible tablet contains Rifampin USP 75 mg Isoniazid USP 50 mg

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Address of certifying authority : Food & Drug Administration, M.S. Bandra-kurla Complex, Bandra (E), Mumbai – 400 051. Maharashtra,INDIA. Tel: +91-22-26592363/64 Fax: +91-22-26591959 IPULI8612214420230329 LUPIN LIMTED - NEW-WHO-GMP/CERT /AD/122144/2023/11/44632 Name of the Authorised person : MR BHUSHAN PATIL , J.C

Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

No. of	certificate : ]	RODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026
		/44632 LUPIN LIMITED
vame		A - 28 / 1, M.I.D.C. CHIKALTHANA,
		AURANGABAD AURANGABAD 431210
		MAHARASHTRA STATE, INDIA
Drug	Dicense 110	633 In Form 25, 499 In Form 28
r.No.	. Name of the Product	Composition
9		Each Film Coated Tablet contains
	Rifapentine and Isoniazid Tablets 300 mg/300mg	Rifapentine 300 mg Isoniazid USP 300 mg
10	AKT-3	Colour:Approved colours used. Each strip contains
10	AK1-3	Rifampicin Capsules 450 mg IP 1 Capsule Nos
		Ethambutol Hydrochloride 800 mg + Isoniazid 300 mg 1 Tablet Nos
11	AKT-3	Each strip contains
		Rifampicin Capsules 450 mg BP 1 Capsule 0
		Ethambutol Hydrochloride 800 mg + Isoniazid 300 mg IH 1 Tablet 0
12	AKT-4	Each strip contains
		Pyrazinamide Tablets 750 mg IP 2 Tablet 0
		Ethambutol Hydrochloride 800 mg + Isoniazid 300 mg 1 Tablet 0 Rifampicin Capsules 450 mg IP 1 Capsule 0
13	AKT-4	Each strip contains
		Pyrazinamide Tablets 750 mg USP 2 Tablets 0
		Ethambutol Hydrochloride 800 mg + Isoniazid 300 mg IH 1 Tablet 0 Rifampicin Capsules 450 mg BP 1 Capsule 0
1.1		DOD AND DRUM
14	AkuriT	Each film coated tablet contains
	Rifampicin and Isoniazid Tablet IP	Rifampicin IP 150.0 mg
		Isoniazid IP 75.0 mg
		Colours Red Ouide of Iren and Titanium Digwitter
15	AKuriT	Colour:Red Oxide of Iron and Titanium Dioxide
	Rifampin and Isoniazid Tablets	Rifampin USP 150 mg
		Isoniazid USP 75 mg
		12 July
		Colour: Iron Oxide of Red and Titanium Dioxide
16	AKurit - 3	Each film coated tablet Contains
		Rifampin USP 150 mg
	Hydrochloride Tablets	Isoniazid USP 75 mg
		Ethambutol Hydrochloride USP 275 mg
22	45678010	Colour:Red Oxide of Iron and Titanium Dioxide
	4 5 6 7 8 9 10	ame of the Authorised person : MR BHUSHAN PATIL , J.C
	s of certifying authority : Na Drug Administration, M.S.	ame of the Authonseu person . With Briddhait FATE, V.C
andra-	-kurla Complex,	Signature :
	(E), Mumbai – 400 051. shtra,INDIA.	Stamp and Date : Joint Commissioner (HQ) & Controlling Authority
	-22-26592363/64	Food & Drug Administration, M.S.
x: +9	91-22-26591959	Bandra (E), Mumbaí.
	12214420230329 NITED - NEW-WHO-GMP/CERT	Maharashtra State, India Date:29 Mar 2023

	certificate : of Manufacturing Firm :	PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED
		A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA
Drug	License No :	633 In Form 25, 499 In Form 28
Sr.No	. Name of the Product	Composition
17	AKurit – Z	Each film coated tablet Contains
	Rifampin, Isoniazid and	Rifampin USP 150 mg
	Pyrazinamide Tablets USP	Isoniazid USP 75 mg
		Pyrazinamide USP 400 mg
10	1 12 - 12 - 12 - 1	Colour:Red Oxide of Iron, Yellow Oxide of Iron & Titanium Dioxide
18	AKurit Kid Rifempia and Iconiazid Tableta	Each uncoated dispersible tablet Contains
	Rifampin and Isoniazid Tablets	Rifampin USP 60 mg
		Isoniazid USP 30 mg
		Colour:Ponceau 4R
19	AKuriT-4	Each film coated tablet contains
	Rifampin, Isoniazid, Pyrazinamide	Rifampin USP 150 mg
	and Ethambutol Hydrochloride	Isoniazid USP 75 mg
	Tablets USP	Pyrazinamide USP 400 mg
		Ethambutol Hydrochloride USP 275 mg
		Colour:Red Oxide of Iron & Titanium Dioxide
20	Bedaquiline Tablets 100 mg	Each Uncoated Tablet contains:
		- Bedaquiline fumarate equivalent to Bedaquiline (100 mg)
21	BILOCOR 10 mg Tablet	Each Tablet Contains
	Bisoprolol Fumarate Tablet	Bisoprolol Fumarate PhEur 10.0 mg
		100
22	BILOCOR 5 mg Tablet	Each Tablet Contains
	Bisoprolol Fumarate Tablet	Bisoprolol Fumarate PhEur 5.0 mg
23	Bisoprolol Fumarate and	Each film coated tablet Contains
	Hydrochlorothiazide Tablets	Bisoprolol Fumarate PhEur 10.0 mg
	Bisoprolol Fumarate and	Hydrochlorothiazide PhEur 6.25 mg
	Hydrochlorothiazide Tablets	I de la
		Colour:Titanium Dioxide
24	Bisoprolol Fumarate and	Each film coated tablet Contains
	Hydrochlorothiazide Tablets	Bisoprolol Fumarate PhEur 5.0 mg
	Bisoprolol Fumarate and	Hydrochlorothiazide PhEur 6.25 mg
	Hydrochlorothiazide Tablets	Colour: Titanium Dioxide and Iron Oxide Red
123	45678910	al-e
ddress	s of certifying authority : N	Name of the Authorised person : MR BHUSHAN PATIL, J.C
ood &	Drug Administration, M.S.	
	kurla Complex, (E), Mumbai – 400 051.	Signature :
	shtra,INDIA.	Stamp and Date : Joint Commissioner (HQ) & Controlling Authority
el: +91	-22-26592363/64	Food & Drug Administration, M.S.
	01-22-26591959 2214420230329	Bandra (E), Mumbai. Maharashtra State, India

Name	certificate : of Manufacturing Firm :	PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 622 In Franz 25, 400 In
Drug I		633 In Form 25, 499 In Form 28
Sr.No.	Name of the Product	Composition
25	Bisoprolol Fumarate and Hydrochlorothiazide Tablets Bisoprolol Fumarate and Hydrochlorothiazide Tablets	Each film coated tablet Contains Bisoprolol Fumarate PhEur 2.5 mg Hydrochlorothiazide PhEur 6.25 mg
26	Combunex	Colour:Titanium Dioxide and Iron Oxide Yellow
20	Ethambutol and Isoniazid Tablet	Each uncoated tablet contains Ethambutol Hydrochloride BP 400.0 mg Isoniazid BP 150.0 mg
		Colour:Sunset Yellow FCF
1000	Combunex 800 Ethambutol and Isoniazid Tablets	Each uncoated tablet contains Ethambutol Hydrochloride BP 800.0 mg Isoniazid BP 300.0 mg
20	c	Colour:Sunset Yellow FCF
10.2	Combunex 800 Ethambutol Hydrochloride & Isoniazid Tablets IP	Each uncoated tablet contains Ethambutol Hydrochloride IP 800.0 mg Isoniazid IP 300.0 mg Colour:Sunset Yellow FCF
29	Combunex HS	Each film coated tablet Contains
	Ethambutol Hydrochloride 400 mg and Isoniazid 150 mg Tablets	Ethambutol Hydrochloride BP 400 mg Isoniazid BP 150 mg Colour:Titanium Dioxide
30	Cameburgal	Each uncoated tablet contains
	Combutol Ethambutol Hydrochloride Tablet IP	Ethambutol Hydrochloride IP 200.0 mg
	Combutol 100 Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 100.0 mg
101	Combutol 1000 Ethambutol Hydrochloride Tablets IP	Each uncoated tablet contains Ethambutol Hydrochloride IP 1000.0 mg
1234	5678910	
Address Food & I Bandra-I Bandra ( Maharas Fel: +91- Fax: +9 PUL18612 UPIN LIMI		ame of the Authorised person : MR BHUSHAN PATIL, J.C. Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

No. of certificate		NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2020 /44632
Name of Manufacturing Firm	1	LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA
Drug License No	;	633 In Form 25, 499 In Form 28

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Sr.No.	Name of the Product	Composition
33	Combutol 1000 Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 1000.0 mg
34	Combutol 200 Ethambutol Hydrochloride Tablets IP	Each uncoated tablet contains Ethambutol Hydrochloride IP 200.0 mg
35	Combutol 200 Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 200.0 mg
36	Combutol 400 Ethambutol Hydrochloride Tablets IP	Each uncoated tablet contains Ethambutol Hydrochloride IP 400.0 mg
37	Combutol 400 Ethambutol Tablets BP	Each film coated tablet Contains Ethambutol Hydrochloride BP 400 mg Colour:Sunset Yellow FCF & Titanium Dioxide
38	Combutol 400 Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 400.0 mg
39	Combutol 600 Ethambutol Hydrochloride Tablets IP	Each uncoated tablet contains Ethambutol Hydrochloride IP 600.0 mg
40	Combutol 600 Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 600 0 mg
1234	45678910	i alle
Food & Bandra- Bandra Mahara	s of certifying authority : N Drug Administration, M.S. kurla Complex, (E), Mumbai – 400 051. shtra,INDIA. -22-26592363/64	lame of the Authorised person : MR BHUSHAN PATIL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S.

Tel: +91-22-26592363/64 Fax: +91-22-26591959 IPUL18612214420230329 LUPIN LIMITED - NEW-WHO-GMP/CERT /AD/122144/2023/11/44632

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Bandra (E), Mumbai.

Maharashtra State, India Date:29 Mar 2023

Name	certificate : of Manufacturing Firm : License No :	PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Product	Composition
41	Combutol 800 Ethambutol Hydrochloride Tablets IP	Each uncoated tablet contains Ethambutol Hydrochloride IP 800.0 mg
42	Combutol 800 Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 800.0 mg
43	Ethambutol 400 mg Tablets BP	Each film coated tablet contains Ethambutol Hydrochloride BP 400.0 mg Colour:Erythrosine & Titanium Dioxide
44	Ethambutol Hydrochloride & Isoniazid Tablets Ethambutol Hydrochloride & Isoniazid Tablets	Each uncoated tablet contains Ethambutol Hydrochloride IP 800.0 mg Isoniazid IP 300.0 mg Colour:Sunset Yellow FCF
45	Ethambutol Hydrochloride 400 & Isoniazid 150 TabletS	Each tablet contains Ethambutol Hydrochloride BP 400.0 mg Isoniazid BP 150.0 mg Colour:Sunset Yellow FCF
46	Ethambutol Hydrochloride 400 mg & Isoniazid 150 mg Tablets	Each film coated tablet contains Ethambutol Hydrochloride BP 400.0 mg Isoniazid BP 150.0 mg Colour:Yellow Oxide Of Iron & Titanium Dioxide
47	Ethambutol Hydrochloride 800 mg & Isoniazid 300 mg Tablets	Each uncoated tablet contains Ethambutol Hydrochloride BP 800.0 mg Isoniazid BP 300.0 mg
	Ethambutol Hydrochloride and Isoniazid Tablets Ethambutol Hydrochloride and Isoniazid Tablets	Colour:Sunset Yellow FCF Each uncoated tablet contains Ethambutol Hydrochloride IP 800.0 mg Isoniazid IP 300.0 mg Colour:Sunset Yellow FCF
1234	45678910	- ALL
Food & Bandra- Bandra Maharas Tel: +91 Fax: +9 IPUL1861 LUPIN LIM	of certifying authority : N Drug Administration, M.S. kurla Complex, (E), Mumbai – 400 051. shtra,INDIA. -22-26592363/64 11-22-26591959 2214420230329 ITED - NEW-WHO-GMP/CERT 14/2023/11/44632	lame of the Authorised person : MR BHUSHAN PATIL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

Name	LIST OF certificate : e of Manufacturing Firm : License No :	F PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
sr.No	Name of the Product	Composition
49	Ethambutol Hydrochloride Tablet IP Ethambutol Hydrochloride Tablet IP	Ethambutol Hydrochloride IP 200.0 mg
50	Ethambutol Hydrochloride Tablet IP Ethambutol Hydrochloride Tablet IP	Ethambutol Hydrochloride IP 400.0 mg
51	Ethambutol Hydrochloride Tablet IP Ethambutol Hydrochloride Tablet IP	Each uncoated tablet contains
52	Ethambutol Hydrochloride Tablet IP Ethambutol Hydrochloride Tablet IP	Ethambutol Hydrochloride IP 400.0 mg
53	Ethambutol Hydrochloride Tablet IP Ethambutol Hydrochloride Tablet IP	Ethambutal Hydrochloride IP 600.0 mg
54	Ethambutol Hydrochloride Tablet IP Ethambutol Hydrochloride Tablet IP	Ethambutol Hydrochloride IP 800.0 mg
55	Ethambutol Tablets BP	Each film coated tablet contains Ethambutol Hydrchloride BP 100.0 mg Colour:Sunset Yellow FCF and Titanium Dioxide
56	Ethambutol Tablets BP	Each film coated tablet contains Ethambutol Hydrochloride BP 400.0 mg Colour: Titanium Dioxide
123	<u>4 5 6 7 8 9 10</u>	1 Call
Food & Bandra Bandra Mahan Tel: +9 Fax: + IPUL186 LUPIN LI	ss of certifying authority : & Drug Administration, M.S. a-kurla Complex, a (E), Mumbai – 400 051. rashtra,INDIA. 01-22-26592363/64 +91-22-26591959 612214420230329 MITED - NEW-WHO-GMP/CERT 1144/2023/11/44632	Name of the Authorised person : MR BHUSHAN PATIL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

Name	certificate of Manufacturing Firm License No	LIST OF PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> : NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 : LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA : 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Pr	oduct Composition
57	Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 600.0 mg
58	Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 800.0 mg
59	Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 1000.0 mg
60	Ethambutol Tablets BP Ethambutol Tablets BP	Each film coated tablet contains Ethambutol Hydrochloride BP 400.0 mg Colour:Sunset Yellow FCF & Titanium Dioxide
61	Ethambutol Tablets BP Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 100.0 mg
62	Ethambutol Tablets BP Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 200.0 mg
63	Ethambutol Tablets BP Ethambutol Tablets BP	Each uncoated tablet contains Ethambutol Hydrochloride BP 400.0 mg
64	Ethide Ethionamide Tablets IP	Each film coated tablet contains Ethionamide IP 250.0 mg Colour:Yellow Oxide of Iron & Titanium Dioxide IP
1234	4567 <b>8</b> 910	
Address Food & I Bandra- Bandra I Maharas Tel: +91- Fax: +9 IPUL1861: LUPIN LIMI	of certifying authority : Drug Administration, M.S. kurla Complex, (E), Mumbai – 400 051. shtra,INDIA. -22-26592363/64 1-22-26591959 2214420230329 ITED - NEW-WHO-GMP/CERT 4/2023/11/44632	Name of the Authorised person : MR BHOSHAN PATIL, J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

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No. of	certificate	IST OF PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> : NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632
Name	e of Manufacturing Firm	: LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA
Drug	License No	: 633 In Form 25, 499 In Form 28
Sr.No	. Name of the Pro	luct Composition
65	Ethide	Each film coated tablet Contains
	Ethionamide Tablets USF	Ethionamide USP 250.0 mg
_		Colour: Titanium Dioxide & Yellow Oxide Of Iron
66	Ethide P	Each film coated tablet contains
	Prothionamide Tablets	Prothionamide IH 250.0 mg
		Colour: Quinoline Yellow and Titanium Dioxide
67	Ethionamide Tablets USF	Each film coated tablet contains Ethionamide USP 250.0 mg
		Colour: Yellow Oxide of Iron and Titanium Dioxide
68	Gatispan 400	Each Film coated tablet contains
00	Gatifloxacin Tablets	Gatifloxacin sesquihydrate equivalent to Gatifloxacin IH 1900 mg
69	Isoniazid Tablets BP	Each uncoated tablet contains
		Isoniazid BP 50.0 mg
70	Isoniazid Tablets BP	Each uncoated tablet Contains
	Isoniazid Tablets BP	Isoniazid BP 100 mg
71	Isoniazid Tablets BP	Each uncoated tablet Contains
	Isoniazid Tablets BP	Isoniazid BP 300 mg
72	Isoniazid Tablets IP	Each uncoated tablet contains
	Isoniazid Tablets IP	Isoniazid IP 300.0 mg
122	45678910	( / -0
		Name of the Authorised person : MR BHUSHAN PATIL, J.C
	s of certifying authority : Drug Administration, M.S.	Name of the Autorised person . Internet and the store store store
landra	-kurla Complex,	Signature :
	(E), Mumbai – 400 051.	Stamp and Date : Joint Commissioner (HQ) & Controlling Author
	ishtra,INDIA. 1-22-26592363/64	Food & Drug Administration, M.S.
ax: +9	91-22-26591959	Bandra (E), Mumbai.
UPIN LIN	12214420230329 MITED - NEW-WHO-GMP/CERT	Maharashtra State, India Date:29 Mar 2023
	44/2023/11/44632	Date:29 War 2023

No. of certificate : Name of Manufacturing Firm : Drug License No :		PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28	
Sr.No.	Name of the Product	Composition	
73	Isoniazid Tablets IP Isoniazid Tablets IP	Each uncoated tablet contains Isoniazid IP 100.0 mg	
74	Isoniazid Tablets IP Isoniazid Tablets IP	Each uncoated tablet contains Isoniazid IP 300.0 mg	
75	L-flox Levofloxacin Hemihydrate Tablets 250 mg	Each film coated tablet contains Levofloxacin Hemihydrate equivalent to Levofloxacin IH 250.0 mg Colour:Yellow Oxide of Iron and Titanium Dioxide	
76	L-flox Levofloxacin Hemihydrate Tablets 500 mg	Each film coated Tablet contains Levofloxacin hemihydrate equivalent to Levofloxacin IH 500.0 mg Colour:Yellow Oxide Of Iron & Titanium Dioxide	
77	Lipril 10 Lisinopril Tablets USP	Each uncoated Tablet Conatins: Lisinopril USP Equivalent to anhydrous Lisinopril 10 mg	
78	Lipril 5 Lisinopril Tablets USP	Each uncoated tablet contains Lisinopril USP equivalent to anhydrous Lisinopril 5.0 mg	
79	Lisinopril Tablets BP 10 mg	Each uncoated tablet contains Lisinopril Dihydrate Ph.Eur equivalent to anhydrous Lisinopril 10.0 mg	
80	Lisinopril Tablets BP 5.0 mg	Each uncoated tablet contains Lisinopril Dihydrate Ph.Eur equivalent to anhydrous Lisinopril 5.0 mg	
123/	45678910		
Food & I Bandra- Bandra I Maharas Tel: +91- Fax: +9 IPUL1861: LUPIN LIMI	Drug Administration, M.S. kurla Complex, (E), Mumbai – 400 051. shtra,INDIA. -22-26592363/64 1-22-26591959 2214420230329 ITED - NEW-WHO-GMP/CER 4/2023/11/44632	Name of the Authorised person : MR-BHUSHAN PATIL , J.C Signature : AND On Gamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023	

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	of Manufacturing Firm : License No :	/44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Product	Composition
81	LISORETIC Tablet 10 / 12.5 mg Lisinopril Dihydrate & Hydrochlorothiazide Tablets	Each uncoated tablet contains Lisinopril Dihydrate equivalent to Lisinopril Ph.Eur 10.0 mg Hydrochlorothiazide Ph.Eur 12.5 mg
82	LISORETIC Tablet 20 / 12.5 mg Lisinopril Dihydrate & Hydrochlorothiazide Tablets	Each uncoated tablet contains Lisinopril Dihydrate equivalent to Lisinopril Ph.Eur 20.0 mg Hydrochlorothiazide Ph.Eur 12.5 mg
83	LupiTB-4	Each Film coated Tablet contains Rifampin USP 150 mg Isoniazid USP 75 mg Pyrazinamide USP 400 mg Ethambutol Hydrochloride USP 275 mg
84	M-Cin Moxifloxacin Hydrochloride Tablet	Each film coated tablet contains Moxifloxacin Hydrochloride equivalent to Moxifloxacin BP 400.0 mg Colour:Red Oxide Of Iron & Titanium Dioxide
85	Mofloxin Lupin Moxifloxacin Hydrochloride Tablets	Each film coated tablet contains Moxifloxacin Hydrochloride BP equivalent to Moxifloxacin 400.0 mg
86	Moksispan Moxifloxacin Hydrochloride Tablets	Each film coated tablet contains Moxifloxacin Hydrochloride BP equivalent to Moxifloxacin 400.0 mg
	Moxyspan Moxifloxacin Hydrochloride Tablets	Each film coated tablet contains Moxifloxacin Hydrochloride BP equivalent to Moxifloxacin 400 mg
	Oseltamivir Phosphate for Oral Suspension 6mg / ml	Colour:Iron Oxide Red Each ml contains Oseltamivir Phosphate USP equivalent to Oseltamivir base (constituted to final concentration) IH 6.0 mg
11 1	2 13 14 15 16 17 18 19 20	
Food & I Bandra-I Bandra ( Maharas	of certifying authority : Na Drug Administration, M.S. Kurla Complex, (E), Mumbai – 400 051. Shtra,INDIA. 22-26592363/64 1-22-26591959 2214420230329 TED - NEW-WHO-GMP/LED 4/2023/11/44632	ame of the Authorised person : MR BHUSHAN PATIL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

Name	LIST OF certificate : of Manufacturing Firm : License No :	PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Product	Composition
89	Prothionamide Tablets Prothionamide Tablets	Each film coated Tablet contains Prothionamide IH 250.0 mg Colour:Quinoline Yellow and Titanium Dioxide
90	Pyrazinamide 500 mg	Each uncoated tablet contains Pyrazinamide BP 500.0 mg
91	Pyrazinamide 500 mg Tablets BP	Each uncoated tablet contains Pyrazinamide BP 500.0 mg Colour:Quinoline Yellow WS and Brillant Blue FCF
92	Pyrazinamide Tablets BP	Each uncoated tablet contains Pyrazinamide BP 150.0 mg
93	Pyrazinamide Tablets BP	Each uncoated tablet contains Pyrazinamide BP 400.0 mg
94	Pyrazinamide Tablets BP	Each uncoated tablet contains Pyrazinamide BP 750.0 mg
95	Pyrazinamide Tablets IP Pyrazinamide Tablets IP	Each uncoated tablet contains Pyrazinamide IP 500.0 mg
96	Pyrazinamide Tablets IP Pyrazinamide Tablets IP	Each uncoated tablet contains Pyrazinamide IP 750:0 mg
11 1	12 13 14 15 16 17 18 19 20	6 6 2
Food & Bandra- Bandra Maharas Fel: +91 Fax: +9 PUL1861 UPIN LIM	e of certifying authority : Drug Administration, M.S. kurla Complex, (E), Mumbai – 400 051. shtra,INDIA. -22-26592363/64 1-22-26591959 2214420230329 ITED - NEW-WHO-GMP//PO5 H4/2023/11/44632	Name of the Authorised person : MR BHUSHAN PATIL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

Name	LIST O certificate : of Manufacturing Firm : License No :	F PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Product	Composition
97	Pyrazinamide Tablets USP	Each uncoated tablet contains Pyrazinamide USP 750.0 mg
98	Pyrazinamide Tablets USP Pyrazinamide Tablets USP	Each uncoated tablet contains Pyrazinamide USP 500.0 mg
99	Pyrazinamide Tablets USP Pyrazinamide Tablets USP	Each uncoated tablet contains Pyrazinamide USP 250.0 mg
100	Pyrazinamide Tablets USP Pyrazinamide Tablets USP	Each uncoated tablet contains Pyrazinamide USP 400.0 mg
101	Pyzina Pyrazinamide Tablets BP	Each uncoated tablet Contains Pyrazinamide BP 500 mg
102	Pyzina Pyrazinamide Tablets IP	Each uncoated tablet contains Pyrazinamide IP 500.0 mg
103	Pyzina Pyrazinamide Tablets USP	Each uncoated tablet contains Pyrazinamide USP 500.0 mg
104	Pyzina 1000 Pyrazinamide Tablets IP	Each uncoated tablet contains Pyrazinamide IP 1000 0 mg
11	12 13 14 15 16 17 18 19 20	i l'all
Food & Bandra- Bandra Maharas Fel: +91 Fax: +9 PUL1861 UPIN LIM	of certifying authority : Drug Administration, M.S. kurla Complex, (E), Mumbai – 400 051. shtra,INDIA. -22-26592363/64 01-22-26591959 2214420230329 ITED - NEW-WHO-GMP/CERT 14/2023/11/44632	Name of the Authorised person : MR BHUSHAN PATIL, J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

	certificate of Manufacturing Firm	LIST OF PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> : NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 : LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210
Drug l	License No	MAHARASHTRA STATE, INDIA : 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Pro	oduct Composition
105	Pyzina 750 Pyrazinamide Tablets BP	Each uncoated tablet contains Pyrazinamide BP 750.0 mg
106	Pyzina 750 Pyrazinamide Tablets IP	Each uncoated tablet contains Pyrazinamide IP 750.0 mg
107	R-Cin Rifampicin Capsule BP	Each Capsule contains Rifampicin BP 150.0 mg
108	R-Cin Rifampicin Capsules IP	Each capsule contains Rifampicin IP 150.0 mg Colour:Approved colours used in capsule shell
109	R-Cin Rifampin Capsule USP	Each capsule contain Rifampin USP 150.0 mg Colour:Approved colours used in capsule shell
110	R-Cin 300 Rifampicin Capsules BP	Each Capsule Contains Rifampicin BP 300 mg Colour:Approved colours used in capsule shell
111	R-Cin 300 Rifampicin Capsules IP	Each capsule contains Rifampicin IP 300.0 mg Colour:Approved colours used in capsule shell
112	R-Cin 300 Rifampin Capsules USP	Each capsule contains Rifampin USP 300.0 mg Colour:Approved colours-used in capsule shell
11	12 13 14 15 16 17 18	
Food & Bandra- Bandra Mahara Tel: +91 Fax: +9 Fax: +9 LPUL1861	s of certifying authority : Drug Administration, M.S. -kurla Complex, (E), Mumbai – 400 051. shtra,INDIA. -22-26592363/64 91-22-26591959 12214420230329 MTED - NEW-WHO-GMP/CERT 44/2023/11/44632	Name of the Authorised person : MR BHUSHAN PATIL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

Name	LIST O certificate : of Manufacturing Firm : License No :	F PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Product	Composition
113	R-Cin 450 Rifampicin Capsules BP	Each Capsule Contains Rifampicin BP 450.0 mg
114	R-Cin 450 Rifampicin Capsules IP	Each Capsule contains Rifampicin IP 450.0 mg
115	R-Cin 450 Rifampin Capsule USP	Colour: Approved colours used in capsule shell Each capsule contains Rifampin USP 450.0 mg Colour: Approved colours used in capsule shell
116	R-Cin 600 Rifampicin Capsules BP	Each Capsule Contains Rifampicin BP 600.0 mg
117	R-Cin 600 Rifampicin Capsules IP	Each Capsule contains Rifampicin IP 600.0 mg Colour: Approved colours used in capsule shell
118	R-Cin Rifampicin BP 300 Rifampicin Capsules BP	Each capsule contains Rifampicin BP 300.0 mg Colour: Approved colours used in capsule shell
119	R-Cinex Rifampin and Isoniazid Capsules USP	Each capsule contains Rifampin USP 450.0 mg Isoniazid USP 300.0 mg Colour:Approved colours used in capsule shell
120	R-Cinex Rifampin and Isoniazid Capsules USP)	Each capsule contains Rifampin USP 450.0 mg Isoniazid USP 300.0 mg Colour: Approved colours used in capsule shell
Address ood & Bandra- Bandra Maharas fel: +91 Fax: +9 PUL1861 UPIN LIM	12 13 14 15 16 17 18 19 20 s of certifying authority : Drug Administration, M.S. kurla Complex, (E), Mumbai – 400 051. shtra,INDIA. -22-26592363/64 01-22-26591959 2214420230329 JITED - NEW-WHO-GMP/CERT 14/2023/11/44632	Name of the Authorised person : MR BHUSHAN PATHL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

No. of certificate : Name of Manufacturing Firm :		PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210
Drug	License No :	MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Product	Composition
121	R-Cinex Kid Rifampicin and Isoniazid Tablets	Each uncoated dispersible tablet contains Rifampicin IP 100.0 mg Isoniazid IP 100.0 mg
122	Rcinex Kid NS Rifampin and Isoniazid dispersible Tablets 75 mg/50 mg	Each uncoated dispersible Tablet contains Rifampin USP 75 mg Isoniazid USP 50 mg Colour:Ponceau 4R
123	Ribavin 200 Ribavirin Capsules	Each Capsule contains Ribavirin USP 200.0 mg Colour: Approved colours used in capsule shell
124	Ributin Rifabutin Capsules USP	Each Capsule contains Rifabutin USP 150 mg
125	Ributin Rifabutin Capsules USP	Each Capsule Contains Rifabutin USP 150 mg
126	Rifabutin Capsules USP	Colour: Approved colours used in capsule shell Each capsule contains Rifabutin USP 150.0 mg
127	RIFAMED 150 Rifampicin Tablets	Colour:Approved colours used in capsule shell Each film coated tablet contains Rifampicin Ph.Eur 150.0 mg Colour:Red Oxide of Iron, Yellow Oxide of Iron, Black Oxide of Iron, Quinoline Yellow WS & Titanium Dioxide
128	RIFAMED 300 Rifampicin Tablets	Each film coated tablet contains Rifampicin Ph.Eur 300.0 mg Colour:Red Oxide of Iron, Yellow Oxide of Iron, Ponceau 4R and Titanium Dioxide
11	12 13 14 15 16 17 18 19 20	
Address Food & Bandra- Bandra Maharas Tel: +91 Fax: +9		Name of the Authorised person : MR BHUSHAN PATIL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

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Name	LIST OF certificate : of Manufacturing Firm : License No :	PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
Sr.No.	Name of the Product	Composition
129	Rifampicin & Isoniazid Capsules	Each Capsule contains Rifampicin IP 450.0 mg Isoniazid IP 300.0 mg Colour: Approved colours used in capsule shell
130	Rifampicin and Isoniazid Tablets Rifampicin and Isoniazid Tablets	Each uncoated Dispersible tablet contains Rifampicin IP 100 mg Isoniazid IP 100 mg Colour: Ponceau 4R
131	Rifampicin Capsuels IP	Each Capsule contains Rifampicin IP 300.0 mg Colour:Approved colour used in Capsule shell
132	Rifampicin Capsules BP	Each capsule contains Rifampicin BP 150.0 mg Colour:Approved colours used in capsule shell
133	Rifampicin Capsules BP	Each Capsule contains Rifampicin BP 300.0 mg Colour:Approved colours used in capsule shell
134	Rifampicin Capsules BP	Each capsule contains Rifampicin BP 450.0 mg Colour:Approved colours used in capsule shell
135	Rifampicin Capsules BP	Each Capsule contains Rifampicin BP 600.0 mg Colour:Approved colours used in capsule shell
136	Rifampicin Capsules IP	Each Capsule contains Rifampicin IP 150.0 mg Colour:Approved colour used in Capsule shell
11	12 13 14 15 16 17 18 19 20	L'd'X
Address Food & Bandra Bandra Mahara Tel: +91 Fax: +91 Fax: +91	s of certifying authority : Drug Administration, M.S. -kurla Complex, (E), Mumbai – 400 051. I-22-26592363/64 91-22-26591959 12214420230329 MITED - NEW-WHO-GMP/CERT 44/2023/11/44632	Name of the Authorised person : MR BHUSHAN PATIL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

in la		PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026
No. of a	certificate :	/44632
Name of Manufacturing Firm :		LUPIN LIMITED
		A - 28 / 1, M.I.D.C. CHIKALTHANA,
		AURANGABAD AURANGABAD 431210
	1	MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In
Drug I	License No	Form 28
_		Composition
137		Composition Each Capsule contains
137	Rifampicin Capsules IP	Rifampicin IP 450.0 mg
		Colour:Approved colour used in Capsule shell
138	Rifampicin Capsules IP	Each Capsule contains
150	Rifampicin Capsules IP	Rifampicin IP 600 mg
	Anompient capsules it	
139	Rifampin and Isoniazid Capsules	Each Capsule contains
	USP	Rifampin USP 450.0 mg
		Isoniazid USP 300.0 mg
		Colour: Approved colour used in Capsule shell
140	Rifampin and Isoniazid Capsules	Each Capsule contains
	USP	Rifampicin IP 450 mg
	Rifampin and Isoniazid Capsules	Isoniazid IP 300 mg
	USP	
141	Rifampin and Isoniazid Tablet	Each uncoated dispersible Tablet contains
		Rifampin USP 60.0 mg
		Isoniazid USP 30.0 mg
		al day
-		Colour:Ponceau 4R
142	Rifampin and Isoniazid Tablets	Each film coated tablet contains
		Rifampin USP 150.0 mg
		Isoniazid USP 150.0 mg
		Colourity and a filtran Bad avide of Iron and Titanium diavide
142	Differentia and transformed Tableta	Colour:Yellow oxide of Iron, Red oxide of Iron and Titanium dioxide Each film coated tablet Contains
143	Rifampin and Isoniazid Tablets	
	Rifampin and Isoniazid Tablets	Rifampin USP 150 mg
		Isoniazid USP 75.0 mg
	1	Colour:Red Oxide of Iron and Titanium Dioxide
144	Rifampin Capsules USP	Each Capsule coantains
0.0.0	The second s	Rifampin USP 450.0 mg
		Colour: Approved colours used in capsule shell
11	12 13 14 15 16 17 18 19 20	L Call
ddres	s of certifying authority :	Name of the Authorised person : MR BHUSHAN PATIL , J.C
S boo	Drug Administration, M.S.	
Sandra	-kurla Complex, (E), Mumbai – 400 051.	Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority
Mahara	ashtra,INDIA.	Stamp and Date : Joint Commissioner (HQ) & Controlling Admonty Food & Drug Administration, M.S.
Tel: +91	1-22-26592363/64	Bandra (E), Mumbai.
	91-22-26591959 12214420230329	Maharashtra State, India
PUL18612214420230329 UPIN LIMITED - NEW-WHO-GMP/CERT		Date:29 Mar 2023

Name	of Manufacturing Firm :	PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28	
Sr.No.	Name of the Product	Composition	
145	Rifampin Capsules USP	Each capsule contains Rifampin USP 150.0 mg	
146	Rifampin Capsules USP	Colour: Approved colours used in capsule shell Each Capsule contains Rifampin USP 600.0 mg Colour: Approved colours used in capsule shell Colour: Approved colours used in capsule shell	
147	Rifampin Capsules USP	Colour:Approved colours used in capsule shell Each Capsule contains Rifampin USP 300.0 mg Colour:Approved colours used in capsule shell	DRUGS
148	Rifampin, Isoniazid and Pyrazinamide Dispersible Tablets Rifampin 75 mg + Isoniazid 50 mg + Pyrazinamide 150 mg Dispersible Tablets	Each uncoated dispersible tablet contains: Rifampin USP 75 mg	ر.
	Rifampin, Isoniazid and Pyrazinamide Tablets USP	Each Film coated Tablet contains. Rifampin USP 150.0 mg Isoniazid USP 75.0 mg Pyrazinamide USP 400.0 mg. Colour:Red Oxide of Iron, Yellow oxide of Iron and Titanium dioxide	-
	Rifampin, Isoniazid, Pyrazinamide and Ethambutol Hydrochloride Tablets USP	Each film coated tablet contains Rifampin USP 150.0 mg Isoniazid USP 75.0 mg Pyrazinamide USP 400.0 mg Ethambutol Hydrochloride USP 275.0 mg	
151	Rifampin,Isoniazid and Ethambutol Hydrochloride Tablet	Colour: Red Oxide of Iron and Titanium Dioxide Each Film coated Tablet contains Rifampin USP 150.0 mg Isoniazid USP 75.0 mg Ethambutol Hydrochloride USP 275.0 mg	
152	Rifampin,Isoniazid and Pyrazinamide Tablets	Colour:Red Oxide of Iron and titanium Dioxide Each uncoated dispersible tablet contains Rifampin USP 60.0 mg Isoniazid USP 30.0 mg Pyrazinamide USP 150:0 mg Colour:Ponceau4R	
11	12 13 14 15 16 17 18 19 20	L'AZ	
Address Food & Bandra- Bandra Maharas Fel: +91 Fax: +9 PUL1861 UPIN LIM		ame of the Authorised person : MR BHUSHAN PATIL , J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023	

Name (	LIST OF certificate : of Manufacturing Firm : License No :	PRODUCT APPROVED UNDER WHO GMP <sup>1</sup> NEW-WHO-GMP/CERT/AD/122144/2023/11 VALID UP TO :28 Mar 2026 /44632 LUPIN LIMITED A - 28 / 1, M.I.D.C. CHIKALTHANA, AURANGABAD AURANGABAD 431210 MAHARASHTRA STATE, INDIA 633 In Form 25, 499 In Form 28
sr.No.	Name of the Product	Composition
153	Rifapentine 300 mg	Each Film coated Tablet Contains Rifapentine 300 mg
	Rifapex Rifapentine Tablets	Colour:Approved colour used. Each film coated tablet contains Rifapentine IH 150.0 mg
	RIFAZID 150 Rifampicin and Isoniazid Tablets	Colour:Red Oxide of Iron and Titanium Dioxide Each film coated tablet contains Rifampicin Ph.Eur 150.0 mg Isoniazid Ph.Eur 100.0 mg Colour:Red Oxide of Iron, Yellow Oxide of Iron and Titanium Dioxide
	RIFAZID 300 Rifampicin and Isoniazid Tablets	Each film coated tablet contains Rifampicin Ph.Eur 300.0 mg Isoniazid Ph.Eur 150.0 mg Colour:Red Oxide of Iron,Yellow Oxide of Iron, Black Oxide of Iron and Titanium
157	ZARTAN 100 mg Losartan Potassium Tablets	Dioxide Each film coated tablet contains Losartan Potassium Ph.Eur 100.0 mg Colour:Titanium Dioxide
158	ZARTAN 50 mg Losartan Potassium Tablet	Each film coated tablet contains Losartan Potassium Ph.Eur 50.0 mg Colour: Titanium Dioxide
Address Food & I Bandra- Bandra Maharas Fel: +91 Fax: +9 PUL1861 UPIN LIM	12 13 14 15 16 17 18 19 20 s of certifying authority : Drug Administration, M.S. -kurla Complex, (E), Mumbai – 400 051. shtra,INDIA. I-22-26592363/64 21-22-26591959 12214420230329 ITED - NEW-WHO-GMP/CERT	Name of the Authorised person : MR BHUSHAN PATIL, J.C Signature : Stamp and Date : Joint Commissioner (HQ) & Controlling Authority Food & Drug Administration, M.S. Bandra (E), Mumbai. Maharashtra State, India Date:29 Mar 2023

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