PH: 0832-2459230, 2459226
Fax.: 0832-2459223
Web Site; www.dfda.goa.gov.in

No.815/MFG/ WHO-GMP/DFDA/2019/ 1472
Government of Goa,
Directorate of Food & Drugs Administration,
'DHANWANTARI' Opp. Shrine of the Holy Cross
Bambolim, Goa-403 202

Dated: 17/7/2019

A copy of this document/CERTIFICATE has been recorded with the Chamber
Authorised Signatory
Bombay Chamber of Commerce and Industry
Regn. No. 128.0 Date 29 JUL 2019
MR. SAMIR PAUL PINTO
ASSISTANT MANAGER

CERTIFICATE

This is to certify that **M/s. Lupin Limited**, situated at Plot No. B-15, Phase IA, Verna Industrial Area, Verna, Goa – 403 722, INDIA, are holding Drug Manufacturing License in Form 25 bearing No. 654 dated 09/03/2004 and in Form 28 bearing No. 833 dated 15/10/2009, issued by this Administration under the provisions of Drugs and Cosmetics Act 1940 and Rules made there under, to manufacture drug formulations at the premises situated at the above address.

It is further certified that the said firm is following **Good Manufacturing Practices** as per World Health Organization (WHO) specification in the manufacturing and testing of the said categories of the products.

It is further certified that the said firm is holding Certificates of Pharmaceutical products granted under **WHO-GMP** certification schemes in the respect of drug products stated in Annexure-I, and the same is valid till **12/03/2022**.

The Firm is also subjected to inspection at regular intervals.

This certificate is issued at the request of M/s. Lupin Limited, situated at Plot No. B-15, Phase IA, Verna Industrial Area, Verna, Goa – 403 722, INDIA.

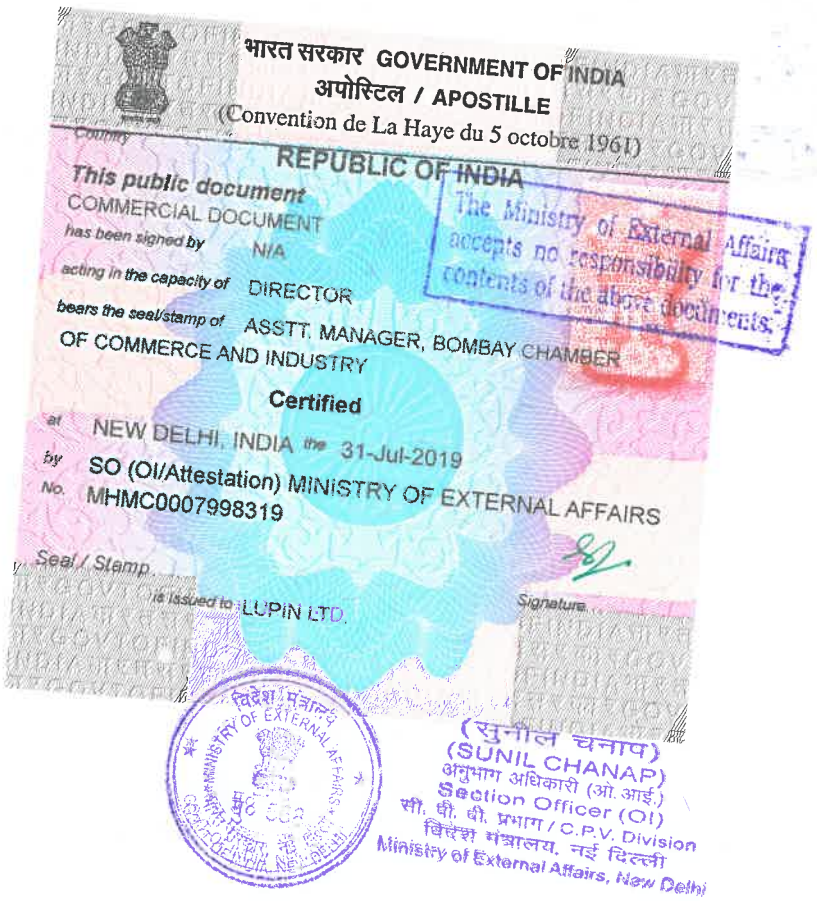
DIRECTORATE OF FOOD & DRUGS ADMN. BAMBOLIM-GOA

Jyoti J. Sardesai
Director, Food & Drugs Admin.

NOTARY Ranjeet Singh Gr. Mumbai Reg. No.9136 Exp.Dt. 20-10-2021 GOVERNMENT OF INDIA

TRUE COPY
Attested by me
RANJEET SINGH
NOTARY GOVT OF INDIA
29 JUL 2019

NOTARY
RANJEET SINGH



भारत सरकार GOVERNMENT OF INDIA
अपोस्टिल / APOSTILLE
(Convention de La Haye du 5 octobre 1961)

1. Country: REPUBLIC OF INDIA

This public document
COMMERCIAL DOCUMENT
has been signed by N/A
acting in the capacity of DIRECTOR
bears the seal/stamp of ASSTT. MANAGER, BOMBAY CHAMBER OF COMMERCE AND INDUSTRY

Certified
at NEW DELHI, INDIA the 31-Jul-2019
by SO (OI/Attestation) MINISTRY OF EXTERNAL AFFAIRS
No. MHMC0007998319

Seal / Stamp
is issued to LUPIN LTD.
Signature

The Ministry of External Affairs accepts no responsibility for the contents of the above documents.

विदेश मंत्रालय MINISTRY OF EXTERNAL AFFAIRS NEW DELHI

(सुनील चनाप)
(SUNIL CHANAP)
अनुभाग अधिकारी (ओ. आई.)
Section Officer (OI)
सी. पी. वी. प्रभाग / C.P.V. Division
विदेश मंत्रालय, नई दिल्ली
Ministry of External Affairs, New Delhi

Annexure - I

Attachment to certificate No. 815/MFG/WHO-GMP/DFDA/2019/ dated: / /2019

Sr. No	Name of the Product	Composition	CoPP No. / Dated
01	Abacavir and Lamivudine Tablets USP 600/300mg	Each film coated tablet Contains: Abacavir Sulfate USP equivalent to Abacavir ..... 600 mg Lamivudine USP..... 300 mg	815/MFG/WHO-GMP/DFDA/ 2019/307 dt. 14 JUN 2019
02	Levetiracetam Tablets 250mg Levetiracetam Lupin 250mg	Each film-coated tablet contains: 250mg Levetiracetam	815/MFG/WHO-GMP/DFDA/ 2019/30 dt. 25 JUN 2019
03	Levetiracetam Tablets 500mg Levetiracetam Lupin 500mg	Each film-coated tablet contains: 500mg Levetiracetam	815/MFG/WHO-GMP/DFDA/ 2019/31 dt. 25 JUN 2019
04	Levetiracetam Tablets 750mg Levetiracetam Lupin 750mg	Each film-coated tablet contains: 750mg Levetiracetam	815/MFG/WHO-GMP/DFDA/ 2019/32 dt. 25 JUN 2019
05	Levetiracetam Tablets 1000mg Levetiracetam Lupin 1000mg	Each film-coated tablet contains: 1000mg Levetiracetam	815/MFG/WHO-GMP/DFDA/ 2019/33 dt. 25 JUN 2019
06	Levetiracetam Extended Release Tablets USP 500 mg	Each film coated extended release tablet contains Levetiracetam USP ... 500 mg	815/MFG/WHO-GMP/DFDA/ 2019/34 dt. 25 JUN 2019
07	Levetiracetam Extended Release Tablets USP 1000 mg	Each film coated extended release tablet contains Levetiracetam USP ... 1000 mg	815/MFG/WHO-GMP/DFDA/ 2019/35 dt. 25 JUN 2019
08	Bupropion Hydrochloride Extended-Release Tablets USP 150mg (XL)	Each extended-release tablets contains: Bupropion Hydrochloride USP 150mg	815/MFG/WHO-GMP/DFDA/ 2019/159 dt. 25 JUN 2019
09	Bupropion Hydrochloride Extended-Release Tablets USP 300mg (XL)	Each extended-release tablets contains: Bupropion Hydrochloride USP 300mg	815/MFG/WHO-GMP/DFDA/. 2019/160 dt. 25 JUN 2019
10	Metformin Hydrochloride Extended Release Tablets USP 500mg	Each extended release coated tablets contains: Metformin Hydrochloride USP 500mg	815/MFG/WHO-GMP/DFDA/ 2019/228 dt. 25 JUN 2019
11	Metformin Hydrochloride Extended Release Tablets USP 1000mg	Each extended release coated tablets contains: Metformin Hydrochloride USP 1000mg	815/MFG/WHO-GMP/DFDA/ 2019/229 dt. 25 JUN 2019

Jyoti J. Sardesal
Director, Food & Drugs Admin.

TRUE COPY

Attested by me
RANJEET SINGH
NOTARY GOVT OF INDIA

29 JUL 2019