

No.: 7-5/2013/EU/WC-049
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organisation
(International Cell)

FDA Bhawan, Kotla Road
New Delhi-110002

Dated: 29 OCT 2018

To

M/s Lupin Ltd
Block 21, Dabhasa, Tal-Padra,
Dist Vadodara-391440, Gujarat, INDIA.

SUB: - Written Confirmation of M/s Lupin Ltd., Block 21, dabasa, Tal-Padra, Dist Vadodara-391440, Gujarat INDIA as per requirement of EU for import of active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC from India-reg.

Sir,

Please refer to your application submitted to CDSCO, Ahmedabad Zone, and the recommendation received from DDC (I), Ahmedabad Zone, on the above noted subject.

Written Confirmation as required for active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC from India is herewith granted subject to the following conditions:-

1. The Active Pharmaceutical Ingredients shall confirm to Good Manufacturing Practices mentioned in the EU directives or other equivalent (GMP of WHO/ICH Q7).
2. The manufacturer is subject to regular, strict and transparent controls and to the effective enforcement of Good Manufacturing Practice, including repeated and unannounced inspections, so as to ensure a protection of public health equivalent to that in the EU.
3. The manufacturer is required to follow the Guidance document for Issue of Written Confirmation as issued by CDSCO.
4. Written Confirmation shall be produced by the Authorized Exporter as and when required by the Drug Regulatory Authority.
5. The Written Confirmation will be withdrawn in the events of non-compliance of Standards.
6. This Written Confirmation, unless it is sooner suspended or cancelled, shall be valid for a period of three years.

7. In the event of any Non-Compliance observed during inspections conducted by Local or International Drug Authorities, the same shall be forwarded to this office within 7 days of receipt of report.

8. In the event of any drug found not of standard quality, the same shall be reported to this office within 7 days of receipt of report.

Please note that Written Confirmation issued is liable to be suspended / cancelled, if any of the conditions stipulated above are not complied with or in case of violation of the provisions of the Drugs & Cosmetics Act, 1940 and the Rules thereunder as the case may be.

Please acknowledge the receipt.

Annexure No.	No. of Products	Date of Issue	Valid up to
01	12	10.09 2018	05/06/2019
02	03	07.12.2017	05/06/2019
03	01	19.03.2018	05/06/2019
04	03	25.06.2018	05/06/2019
05	02	29 OCT 2018	05/06/2019

Yours faithfully,



(Dr. S. Eswara Reddy)
Drugs Controller General (India)



GOVERNMENT OF INDIA
MINISTRY OF HEALTH & FAMILY WELFARE
Central Drugs Standard Control Organization

Annexure-05
WC-0049

CERTIFICATE NO. :

Written confirmation for active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC

**1. Name and address of site: M/s Lupin Limited,
Block -21,Dabhasa,Tal-Padra,
Dist-Vadodara-391440,Gujarat,India**

List of APIs:

Sl. No.	Active substance(s)	Activity(ies)
1.	Levetriacetam USP	Manufacturing & Packing
2.	Desvenlafaxine Succinate	Manufacturing & Packing

ITEM(S) TWO (02) ONLY

The Written Confirmation remains valid until: 05th June 2019


Signature

Stamp of the authority and date



29 OCT 2018