

7-5/2013/EU/WC-0119
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organisation
(International Cell)

FDA Bhawan, Kotla Road,
New Delhi-110002
Dated

To

**M/s. Aurore Pharmaceutical Private Limited,
Plot No. 35,36,38,39,40,49,50 & 51,
Phase –IV, IDA Jeedimetla, Medchal-Malkajgiri District,
Telangana State, India**

07 AUG 2020

SUB:- Written Confirmation of M/s. Aurore Pharmaceutical Private Limited, Plot No. 35,36,38,39,40,49,50 & 51, Phase –IV, IDA Jeedimetla, Medchal-Malkajgiri District, Telangana State, 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, Hyderabad Zone office, and the recommendation received from DDC (I), Hyderabad 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 conform 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 The Written Confirmation will be withdrawn in the events of non compliance of Standards.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.

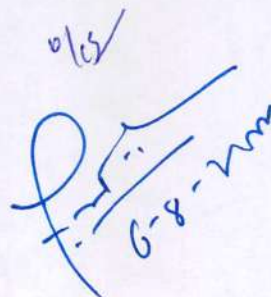
Please acknowledge the receipt.

Annexure No.	No. of Products	Date of Issue	Valid Upto
1	20	09.03.2017	02.07.2022
2	01	09.03.2017	02.07.2022
3	06	12.06.2020	02.07.2022
4	01	12.06.2020	02.07.2022
5	08	07 AUG 2020	02.07.2022
6	01	07 AUG 2020	02.07.2022

Yours faithfully,



(Dr. V.G.Somani)
Drugs Controller General (India)

01/05

 6-8-2020



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. Aurore Pharmaceutical Private Limited,
Plot No. 35,36,38,39,40,49,50 & 51,
Phase -IV, IDA Jeedimetla, Medchal-Malkajgiri
District, Telangana State, India**

List of APIs:

Sr. No.	Active substance (s)	Activity(ies)
1.	Amlodipine Besilate Ph.Eur/BP	Manufacturing & Packing
2.	Amlodipine Besylate USP	Manufacturing & Packing
3.	Levetiracetam USP/Ph.Eur/BP	Manufacturing & Packing
4.	Tolvaptan IH	Manufacturing & Packing
5.	Valaciclovir Hydrochloride Ph.Eur/BP	Manufacturing & Packing
6.	Valacyclovir Hydrochloride USP	Manufacturing & Packing
7.	Valaciclovir Hydrochloride Hydrated Ph.Eur/BP	Manufacturing & Packing
8.	Levothyroxine Sodium USP/EP	Manufacturing & Packing

ITEM(S) EIGHT (08) ONLY

The Written Confirmation remains valid until: 02.07.2022

Signature

[Handwritten signature]
0102
6-8-2020

Stamp of the authority and date



07 AUG 2020



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. Aurore Pharmaceutical Private Limited,
Plot No. 35,36,38,39,40,49,50 & 51,
Phase -IV, IDA Jeedimetla, Medchal-Malkajgiri
District, Telangana State, India**

List of APIs:

Sr. No.	Active substance (s)	Activity(ies)
1.	Desvenlafaxine Oxalate IH	Manufacturing & Packing

ITEM(S) ONE (01) ONLY

This certificate is being issued subject to condition that the firm shall obtained NOC from the Competent Authority, on case to case basis, to manufacture the above mentioned active substance for the purpose of export only, as the above mentioned active substance is not approved for manufacture for sale in India

The Written Confirmation remains valid until: 02.07.2022

Signature

[Handwritten signature]
6-8-2020

Stamp of the authority and date



07 AUG 2020