

7-5/2016/EU/WC-0370  
Government of India  
Directorate General of Health Services  
Central Drugs Standard Control Organisation  
(International Cell)

FDA Bhawan, Kotla Road  
New Delhi-110002  
Dated:

28 JAN 2020

To

M/s. Alkem Laboratories Ltd., Plot No. 288/A, 289-290, G.I.D.C., At & Post-Ankleshwar, Dist.-Bharuch, Gujarat, India.

**SUB: Written Confirmation of M/s. Alkem Laboratories Ltd., Plot No. 288/A, 289-290, G.I.D.C., At & Post- Ankleshwar, Dist.-Bharuch, 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 office 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 The Written Confirmation will be withdrawn in the events of non-compliance of Standards.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	Validity
1	06	18.11.2019	05.06.2022
2	02	18.11.2019	05.06.2022
3	02	28 JAN 2020	05.06.2022

Yours faithfully,

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





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

Annexure- 03

CERTIFICATE NO. : WC 0370

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

Name and Address of site: M/s. Alkem Laboratories Ltd., Plot No. 288/A, 289-290, G.I.D.C., At & Post- Ankleshwar, Dist- Bharuch, Gujarat, India.

List of APIs:

Sl. No.	Name of the Active Substance (S)	Activity(ies)
1.	Rizatriptan Benzoate USP	Manufacturing & Packing
2.	Riluzole USP	Manufacturing & Packing

ITEM(S) Two (2) Only

The Written Confirmation remains valid until: 05.06.2022

Signature

Stamp of the authority and date



28 JAN 2020