7-5/2013/EU/WC-0082 Government of India Directorate General of Health Services Central Drugs Standard Control Organisation (International Cell)

FDA Bhawan, Kotla Road, New Delhi-110002 Dated

To

0 9 MAY 2022

M/s Alembic Pharmaceuticals Limited, API Division Karakhadi, Plot No. 842-843, At – Karakhdi, Tal. – Padra, Dist – Vadodara, Gujarat

SUB:- Written Confirmation of M/s Alembic Pharmaceuticals Limited, API Division, Karakhadi, Plot No. 842-843, At – Karakhdi, Tal. – Padra, Dist – Vadodara 391 450, Gujarat 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 online application no. WC/FR/2022/2106 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:-

- 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 The Written Confirmation will be withdrawn in the events of non compliance of Standards.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 Upto
01	27	26.06.2019	02.07.2022
02	13	26.06.2019	02.07.2022
03	05	05.07.2019	02.07.2022
04	03	05.07.2019	02.07.2022
05	08	22.10.2019	02.07.2022
06	01	18.02.2020	02.07.2022
07	02	20.03.2020	02.07.2022
08	04	20.04.2021	02.07.2022
09	01	19.09.2021	02.07.2022
10	. 03	05.10.2021	02.07.2022
11	08	05.10.2021	02.07.2022
12	05	02.12.2021	02.07.2022
13	01	02.12.2021	02.07.2022
14	03	0 9 MAY 2022	02.07.2022

Yours faithfully,

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

0/1 5

Annexure-14

CERTIFICATE NO.:

WC-0082

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 Alembic Pharmaceuticals Limited, API Division

Karakhadi, Plot No. 842-843, At - Karakhdi,

Tal. - Padra, Dist - Vadodara, Gujarat

List of APIs:

Sr. No.	Active substance (s)	Activity(ies)
1.	Donepezil Hydrochloride Monohydrate Ph.Eur.	Manufacturing & Packing
2.	Fluphenazine Dihydrochloride Ph. Eur.	Manufacturing & Packing
3.	Theophylline Ph. Eur.	Manufacturing & Packing

ITEM(S) Three (03) ONLY

The Written Confirmation remains valid until: 02.07.2022

Signature Vinc

the authority and date

0 9 MAY 2022