7-5/2013/EU/WC-0144
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organisation
(International Cell)

FDA Bhawan,Kotla Road,
New Delhi-110002
Dated: 09 SEP 2019

To

M/s. Cipla Limited,
D-7, MIDC, Industrial Area, Kurkumbh,
Taluka: Daund, District: Pune -Zone4- 413 802
Maharashtra, India

Subject:- Written Confirmation of M/s.Cipla Limited, D-7, MIDC, Industrial Area, Kurkumbh, Taluka: Daund, District: Pune- Zone4- 413 802, Maharashtra, 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, West Zone office Mumbai and the recommendation received from DDC(I), West Zone Mumbai 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 The Written Confirmation will be withdrawn in the events of non compliance of Standards.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 Upto
1	24	09 SEP 2019	Three (03) years from date of issue
2	04	09 SEP 2019	Three (03) years from date of issue

O/C

Yours faithfully,

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

30/08/19

30-8-19

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

CERTIFICATE NO. : WC-0144

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.Cipla Limited,**
D-7, MIDC, Industrial Area, Kurkumbh,
Taluka: Daund, District: Pune -Zone4- 413 802
Maharashtra, India

2. **Manufacturer's licence number**: 28-PD/42 and 25-PD/46

Regarding the manufacturing plant under (1) of the following Active substance(s) exported to the EU for medicinal products for human use

As per List enclosed as Annexure-1 & 2

The issuing Regulatory Authority hereby confirms that:

The standards of good manufacturing practice applicable to this manufacturing plant are at least equivalent to those laid down in the EU (=GMP of WHO/ICH Q7);

The manufacturing plant 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 at least equivalent to that in the EU; and

In the event of findings relating to non-compliance, information on such findings is supplied by the exporting third country without delay to the EU.

Date of Inspection of the plant: 27.06.2019 & 28.06.2019

The Written Confirmation remains valid until: **Three (03) years from date of issue.**

The authenticity of this written confirmation may be verified with the issuing regulatory authority.

This written confirmation is without prejudice to the responsibilities of the manufacturer to ensure the quality of the medicinal product in accordance with Directive 2001/83/EC.

Address of the issuing regulatory authority: **Central Drugs Standard Control Organisation**
FDA Bhawan, Kotla Road,
New Delhi- 110 002, India

O/C

Name and function of responsible person: Dr. V.G. Somani
Drugs Controller General (India)

E-mail: dci@nic.in,
Telephone no.: +91-11-23236965
Fax no.: +91-11-23236973

Signature

30/08/19

30.8.19

Stamp of the authority and date

Central Drugs Standard Control Organisation
सत्यमेव जयते
Ministry of Health Family Welfare, Government of India

09 SEP 2019

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

CERTIFICATE NO. : **Annexure-1 WC-0144**

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. Cipla Limited,**
D-7, MIDC, Industrial Area, Kurkumbh,
Taluka: Daund, District: Pune- Zone4- 413 802
Maharashtra, India

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1.	Alendronate Sodium Trihydrate USP	Manufacturing & Packing
2.	Carvedilol BP/Ph.Eur/USP	Manufacturing & Packing
3.	Celecoxib BP/Ph.Eur/USP	Manufacturing & Packing
4.	Citalopram Hydrobromide BP/Ph.Eur/USP	Manufacturing & Packing
5.	Darunavir Ethanolate IH	Manufacturing & Packing
6.	Disodium Pamidronate BP/Ph.Eur	Manufacturing & Packing
7.	Famciclovir IH	Manufacturing & Packing
8.	Fluconazole BP/Ph.Eur/USP	Manufacturing & Packing
9.	Formoterol Fumarate Dihydrate BP/Ph.Eur/USP	Manufacturing & Packing
10.	Glatiramer Acetate IH	Manufacturing & Packing
11.	Lamivudine BP/Ph.Eur/USP	Manufacturing & Packing
12.	Meloxicam BP/Ph.Eur/USP	Manufacturing & Packing
13.	Olanzapine BP/Ph.Eur/USP	Manufacturing & Packing
14.	Ondansetron IH	Manufacturing & Packing
15.	Ondansetron Hydrochloride Dihydrate BP/Ph.Eur/USP	Manufacturing & Packing
16.	Pramipexole Dihydrochloride Monohydrate BP/Ph.Eur/USP	Manufacturing & Packing
17.	Ramipril BP/Ph.Eur/USP	Manufacturing & Packing
18.	Sertraline Hydrochloride BP/Ph.Eur/USP	Manufacturing & Packing
19.	Tiotropium Bromide Monohydrate BP/Ph.Eur	Manufacturing & Packing
20.	Zoledronic Acid IH	Manufacturing & Packing
21.	Budesonide BP/Ph.Eur/USP	Manufacturing & Packing
22.	Ciclesonide IH	Manufacturing & Packing
23.	Fluticasone Propinate BP/Ph.Eur/USP	Manufacturing & Packing
24.	Mometasone Furoate BP/Ph.Eur/USP	Manufacturing & Packing

ITEM(S) Twenty Four (24) ONLY

The Written Confirmation remains valid until: Three (03) years from date of issue.

Signature

30/08/19

30-8-19

Stamp of the authority and date

Central Drugs Standard Control Organisation
सत्यमेव जयते
Ministry of Health & Family Welfare, Government of India

09 SEP 2019

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

CERTIFICATE NO. : Annexure-2 WC-0144

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. Cipla Limited,
D-7, MIDC, Industrial Area, Kurkumbh,
Taluka: Daund, District: Pune- Zone4- 413 802
Maharashtra, India

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1.	Risedronate Sodium 2.5 Hydrate Ph.Eur/BP	Manufacturing & Packing
2.	Valganciclovir Hydrochloride USP	Manufacturing & Packing
3.	Beclometasone Dipropionate USP	Manufacturing & Packing
4.	Beclometasone Dipropionate Monohydrate BP/Ph.Eur	Manufacturing & Packing

ITEM(S) Four (04) ONLY

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

01C **The Written Confirmation remains valid until: Three (03) years from date of issue.**

Signature

Stamp of the authority and date

Central Drugs Standard Control Organization
सत्यमेव जयते
Ministry of Health & Family Welfare, Government of India

09 SEP 2019

30/08/19

30-8-19