

7-5/2013/EU/WC-0136  
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 2025

To

M/s. Dishman Carbogen Amcis Limited  
Plot No. 1216/11, 12 & 1216/20 TO 26,  
G.I.D.C. Estate, Phase IV, Naroda,  
Ahmedabad -382330, Gujarat, India

**Subject:** - Written Confirmation of M/s. Dishman Carbogen Amcis Limited, Plot No. 1216/11, 12 & 1216/20 TO 26, G.I.D.C. Estate, Phase IV, Naroda, Ahmedabad -382330, 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 online application no. WC/RE/2025/9991 submitted to CDSCO, Ahmedabad Zone office on CDSCO Sugam Portal, 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 The Written Confirmation will be withdrawn in the events of non-compliance of Standards.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.
9. The manufacturer is required to comply with the provisions of GSR 20(E), dated 18.01.2022

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
--	--	09 SEP 2025	22.07.2028
1	07	09 SEP 2025	22.07.2028

Yours faithfully,

*Chandrashekhar*  
08/09/2025

Ranga Chandrashekhar  
Joint Drugs Controller (India)

चंद्रशेखर रंगा/Chandrashekhar Ranga

संघरक औषधि नियंत्रक (भारत) / Joint Drugs Controller (India)

केंद्रीय औषधि मानक नियंत्रण संगठन (मुख्यालय), स्वास्थ्य सेवा महानियंत्रण

C.D.S.C. (HQ), Dte. General of Health Services

स्वास्थ्य और परिवार कल्यान मंत्रालय / Ministry of Health and Family Welfare

एफ. हो.ए. भवन, कोटा रोड, नई दिल्ली-110002 / FDA Bhawan, Koda Road, New Delhi-110002



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

CERTIFICATE NO. :

WC-0136

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. Dishman Carbogen Amcis Limited  
Plot No. 1216/11, 12 & 1216/20 TO 26,  
G.I.D.C. Estate, Phase IV, Naroda,  
Ahmedabad -382330, Gujarat, India

2. Manufacturer's licence number: G/25/799

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

**List of API(s):**

as per list enclosed as Annexures

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:** 13.05.2024 and 14.05.2024

The Written Confirmation remains valid until: **22.07.2028**

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

**Name and function of responsible person:** Ranga Chandrashekhar,  
Joint Drugs Controller (India)

**E-mail:** [ranga.cs@cdsco.nic.in](mailto:ranga.cs@cdsco.nic.in);

**Telephone no.:** +91-11-23236965

**Fax no.:** +91-11-23236973

*Ranga Chandrashekhar*  
रांगा चंद्रशेखर संगम/Chandrashekhar Ranga  
संयुक्त ड्रग्स कंट्रोलर (भारत) / Joint Drugs Controller (India)  
केंद्रीय औषधि प्रायोगिक संशोधन संसाधन (मुख्यालय), स्वास्थ्य सेवा महानिदेशालय  
C.D.S.C.O(HQ), Dir. General of Health Services  
स्वास्थ्य और परिवार कल्याण मंत्रालय / Ministry of Health and Family Welfare  
एम.डी.ए.भवन, कोटला रोड, नई दिल्ली-110002 / FDA Bhawan, Kotla Road, New Delhi-110002

Stamp of the authority and date



09 SEP 2025



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

CERTIFICATE NO. : Annexure-01  
WC-0136

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. Dishman Carbogen Amcis Limited  
Plot No. 1216/11, 12 & 1216/20 TO 26,  
G.I.D.C. Estate, Phase IV, Naroda,  
Ahmedabad -382330, Gujarat, India

List of APIs:

S. No.	Active Substance(s)	Activity(ies)
1.	Benzalkonium Chloride Solution BP/USP/JP/Ph.Eur.	Manufacturing & Packing
2.	Benzethonium Chloride BP/USP/JP/Ph.Eur.	Manufacturing & Packing
3.	Bisacodyl BP/USP/JP/Ph.Eur.	Manufacturing & Packing
4.	Cetrimide BP/Ph.Eur.	Manufacturing & Packing
5.	Cetylpyridinium Chloride BP/USP/JP/Ph.Eur.	Manufacturing & Packing
6.	Sodium Picosulfate BP/USP/JP/Ph.Eur.	Manufacturing & Packing
7.	Strong Cetrimide Solution BP	Manufacturing & Packing

ITEM(S) Seven (07) ONLY

The Written Confirmation remains valid until: 22.07.2028

*chandrashekhar*  
Signature 08/09/25  
चंद्रशेखर राणा/Chandrashekhar Ranjan  
जोड़ी औषधि नियंत्रक (भारत) / Joint Drugs Controller(India)  
C.D.S.C.OHQ, Dir. General of Health Services  
स्वास्थ्य और पर्यावरण मंत्रालय / Ministry of Health and Family Welfare  
एफडीए भवन, लोटपोहा, नई दिल्ली-110002 / FDA Bhawan, Lohapura, New Delhi-110002



09 SEP 2025