

7-5/2013/EU/WC-0024
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organisation
(International Cell)

FDA Bhawan, Kotla Road,
New Delhi-110002
Dated: 08 AUG 2025

To,

**M/s. Granules India Limited (Unit-IV),
Plot No. 8, Jawaharlal Nehru Pharma City,
Tadi (V), Parawada Mandal, Anakapalli District -531019,
Andhra Pradesh, India**

SUB:- Written Confirmation of **M/s. Granules India Limited (Unit-IV), Plot No. 8, Jawaharlal Nehru Pharma City, Tadi (V), Parawada Mandal, Anakapalli District -531019, Andhra Pradesh, 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/2025/11175 submitted to CDSCO, ADC(I), Sub-Zone Visakhapatnam, and the recommendation received from ADC(I), Sub-Zone Visakhapatnam, 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.

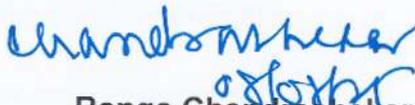
5. The Written Confirmation will be withdrawn in the events of non compliance of Standards.
6. This Written Confirmation, unless it is sooner suspended or cancelled, shall be valid for a period of three years.
7. In the event of any Non Compliance observed during inspections conducted by Local or International 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
--	--	0 8 AUG 2025	26.06.2028
01	21	0 8 AUG 2025	26.06.2028

Yours faithfully,


Ranga Chandrashekar
 Joint Drugs Controller (India)

चंद्रशेखर रंगा/Chandrashekar Ranga
 सचिव औषधि नियंत्रण, भारत (Joint Drugs Controller(India))
 केंद्रीय औषधि मानक नियंत्रण (सुखवाड़ा), स्वास्थ्य सेवा महानिदेशालय
 C.D.S.C.(I/O), Dte. General of Health Services
 स्वास्थ्य और परिवार कल्याण मंत्रालय / Ministry of Health and Family Welfare
 एन.डी.ए. भवन, कोटला रोड, नई दिल्ली-110002 / FDA Bhawan, Kotla Road, New Delhi-110002



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. Granules India Limited (Unit-IV),
Plot No. 8, Jawaharlal Nehru Pharma City,
Tadi (V), Parawada Mandal, Anakapalli District -531019,
Andhra Pradesh, India**

2. Manufacturer's licence number: **26/VP/API/2007/B/CC/R & 19/VP/API/2013/B/CC/R**

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: 12.09.2023 & 13.09.2023

The Written Confirmation remains valid until: **26.06.2028**

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 Chandrashekar,
Joint Drugs Controller (India)**

E-mail: ranga.cs@cdsco.nic.in;

Telephone no.: +91-11-23236965

Fax no.: +91-11-23236973

Chandrashekar
08/08/25

Signature: **चंद्रशेखर रंगा/Chandrashekar Ranga**
केन्द्रीय औषधि मानक नियंत्रण संगठन (भारत) / Joint Drugs Controller (India)
C.D.S.C.O.(HQ), Dte. General of Health Services
स्वास्थ्य और परिवार कल्याण विभाग / Ministry of Health and Family Welfare
एच.डी.ए. भवन, कोटला रोड, नई दिल्ली-110002 / FDA Bhawan, Kotla Road, New Delhi-110002

Stamp of the authority and date



08 AUG 2025



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. Granules India Limited (Unit-IV),
Plot No. 8, Jawaharlal Nehru Pharma City, Tadi (V),
Parawada Mandal, Anakapalli District -531019,
Andhra Pradesh, India

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1.	Atazanavir Sulphate IH/Ph.Eur.	Manufacturing & Packing
2.	Azathioprine USP/Ph.Eur.	Manufacturing & Packing
3.	Cetirizine Dihydrochloride Ph.Eur.	Manufacturing & Packing
4.	Dimethyl Fumarate IH	Manufacturing & Packing
5.	Brinzolamide IH/USP	Manufacturing & Packing
6.	Fexofenadine Hydrochloride USP	Manufacturing & Packing
7.	Gliclazide IP/IH/BP/Ph.Eur.	Manufacturing & Packing
8.	Darunavir Amorphous IH	Manufacturing & Packing
9.	Levocetirizine Dihydrochloride IH	Manufacturing & Packing
10.	Losartan Potassium Ph.Eur.	Manufacturing & Packing
11.	Metformin Hydrochloride IP/IH/BP/USP	Manufacturing & Packing
12.	Omeprazole Magnesium IH/USP	Manufacturing & Packing
13.	Pencillamine USP/Ph.Eur.	Manufacturing & Packing
14.	Pirfenidone IH/Ph.Eur.	Manufacturing & Packing
15.	Rifaximin Ph.Eur.	Manufacturing & Packing
16.	Tenofovir Disoproxil Succinate IH	Manufacturing & Packing
17.	Tizanidine Hydrochloride IP/USP	Manufacturing & Packing
18.	Methocarbamol USP	Manufacturing & Packing
19.	Dapagliflozin IH	Manufacturing & Packing
20.	Empagliflozin IH	Manufacturing & Packing
21.	Trazadone Hydrochloride IH/USP	Manufacturing & Packing

ITEM(S) TWENTY-ONE (21) ONLY

The Written Confirmation remains valid until: 26.06.2028


Signature of Chandrashekar Ranga

संयुक्त औषधि नियंत्रक (भारत) / Joint Drugs Controller (India)
केन्द्रीय औषधि मानक निबंधन संगठन (मुंबई/कोलकाता), स्वास्थ्य सेवा महानिदेशालय
C.D.S.C.O.(HQ), Dte. General of Health Services
स्वास्थ्य और परिवार कल्याण विभाग / Ministry of Health and Family Welfare
एन.डी.ए. भवन, मोहन रोड, अंधाबाद-110002 / FDA Bhawan, Kola Road, New Delhi-110002



08 AUG 2025