

**7-5/2024/EU/WC-0598**  
Government of India  
Directorate General of Health Services  
Central Drugs Standard Control Organisation  
(International Cell)

FDA Bhawan, Kotla Road,  
New Delhi-110002

Date: 6 DEC 2024

To,

**M/s. Vasudha Pharma Chem Limited,  
Unit-V: Plot No. 24, 24A & 24B,  
APSEZ-DENOTIFIED Area, Rambilli(M),  
Atchutapuram-531 011, Anakapalli District,  
Andhra Pradesh, India**

**SUB:-** Written Confirmation **M/s. Vasudha Pharma Chem Limited, Unit-V: Plot No. 24, 24A & 24B, APSEZ-DENOTIFIED Area, Rambilli(M), Atchutapuram-531 011, Anakapalli District, Andhra Pradesh, India**, as per requirement of the 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/2024/8156 submitted to CDSCO, Visakhapatnam Subzone, and the recommendation received from ADC(I), Visakhapatnam Subzone, 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 The Written Confirmation will be withdrawn in the events of non-compliance of Standards.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
--	--		Three Years from date of issue
01	16	06 DEC 2024	Three Years from date of issue

Yours faithfully,

*Chandrashekar*  
06/12/24  
(Ranga Chandrashekar)  
Joint Drugs Controller (India)

चंद्रशेखर रंगा/Chandrashekar Ranga  
संयुक्त औषधि नियंत्रक (भारत) / Joint Drugs Controller (India)  
केंद्रीय औषधि मानक नियंत्रण संगठन (मुंबई), स्वास्थ्य सेवा महानिदेशालय  
C.D.S.C.O(HQ), Dte. General of Health Services  
स्वास्थ्य और परिवार कल्याण मंत्रालय / Ministry of Health and Family Welfare  
एच.टी.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. Vasudha Pharma Chem Limited,  
Unit-V: Plot No. 24, 24A & 24B,  
APSEZ-DENOTIFIED Area, Rambilli(M),  
Atchutapuram-531 011, Anapapalli District,  
Andhra Pradesh, India

2. Manufacturer's licence number: 14/VSP/AP/2019/B/G

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 Annexure-01

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: 19.02.2024 and 20.02.2024

The Written Confirmation remains valid until: Three 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

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

[ranga.cs@cdsco.nic.in](mailto:ranga.cs@cdsco.nic.in)

E-mail:

Telephone no.:

+91-11-23236965

Fax no.:

+91-11-23236973

Signature

चंद्रशेखर रंगा/Chandrashekar Ranga

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

केन्द्रीय औषधि मानक निरीक्षण संगठन (मुख्यालय), 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

एफ डी ए भवन, कोटला रोड, नई दिल्ली-110002 / FDA Bhawan, Kotla Road, New Delhi-110002

Stamp of the authority and date





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

CERTIFICATE NO. :

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. Vasudha Pharma Chem Limited,  
Unit-V: Plot No. 24, 24A & 24B,  
APSEZ-DENOTIFIED Area, Rambilli(M),  
Atchutapuram-531 011, Anapapalli District,  
Andhra Pradesh, India

List of API(s):

Sr. No.	Active substance (s)	Activity(ies)
1.	Amitriptyline Hydrochloride USP/Ph.Eur./BP/IP	Manufacturing & Packing
2.	Cyproheptadine Hydrochloride USP/Ph.Eur. /BP/IP	Manufacturing & Packing
3.	Domperidone Ph.Eur./BP	Manufacturing & Packing
4.	Domperidone Maleate Ph.Eur./BP	Manufacturing & Packing
5.	Esomeprazole magnesium USP	Manufacturing & Packing
6.	Esomeprazole magnesium trihydrate Ph.Eur. /BP	Manufacturing & Packing
7.	Fexofenadine Hydrochloride USP	Manufacturing & Packing
8.	Flurbiprofen BP/ USP / Ph.Eur.	Manufacturing & Packing
9.	Levetiracetam USP / Ph. Eur.	Manufacturing & Packing
10.	Losartan Potassium USP /Ph.Eur./BP/IP	Manufacturing & Packing
11.	Mebeverine Hydrochloride Ph.Eur. /BP	Manufacturing & Packing
12.	Nortriptyline Hydrochloride USP / Ph.Eur. /BP/IP	Manufacturing & Packing
13.	Olmesartan Medoxomil USP / Ph.Eur./BP	Manufacturing & Packing
14.	Pantaprazole Sodium USP/IP	Manufacturing & Packing
15.	Pantaprazole Sodium Sesquihydrate Ph.Eur. /BP	Manufacturing & Packing
16.	Telmisartan USP / Ph.Eur./ BP/ IP	Manufacturing & Packing

ITEM(S) SIXTEEN (16) ONLY

The Written Confirmation remains valid until: Three years from date of issue

*Chandrashekar Ranga*

Signature

चंद्रशेखर रंगा/Chandrashekar Ranga  
संयुक्त औषधि नियंत्रक (भारत) / Joint Drugs Controller (India)  
केन्द्रीय औषधि मानक नियंत्रण संगठन (मुख्यालय), स्वास्थ्य सेवा महानिदेशालय  
C.D.S.C.(HQ), Dte. General of Health Services  
स्वास्थ्य सेवा महानिदेशालय / Ministry of Health and Family Welfare  
एन.डी.ए. भवन, कोटा रोड, नई दिल्ली-110002



Stamp of the authority and date

06 DEC 2024