

**7-5/2021/EU/WC-0501**  
Government of India  
Directorate General of Health Services  
Central Drugs Standard Control Organisation  
(International Cell)

FDA Bhawan, Kotla Road,  
New Delhi-110002

Dated:

16 MAY 2024

To,

**M/s. Escientia Advanced Sciences Pvt. Ltd.,  
Plot No. 11, 11A, 12 & 12A, APSEZ,  
Atchutapuram, Anakapalli District -531011,  
Andhra Pradesh, India**

**SUB:-** Written Confirmation of **M/s. Escientia Advanced Sciences Pvt. Ltd., Plot No. 11, 11A, 12 & 12A, APSEZ, Atchutapuram, Anakapalli District -531011, 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/2024/7906 dated 13.01.2024 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 The Written Confirmation will be withdrawn in the events of non compliance of Standards.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
--	--	16 MAY 2024	24.05.2027
01	01	16 MAY 2024	24.05.2027

Yours faithfully,

(Dr. Rajeev Singh Raghuvanshi)  
Drugs Controller General (India)



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. Escientia Advanced Sciences Pvt. Ltd.,  
Plot No. 11, 11A, 12 & 12A, APSEZ,  
Atchutapuram, Anakapalli District -531011,  
Andhra Pradesh, India

2. Manufacturer's licence number: 06/VSP/AP/2018/B/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: 20.06.2023 & 21.06.2023

The Written Confirmation remains valid until: 24.05.2027

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: Dr. Rajeev Singh Raghuvanshi,  
Drugs Controller General (India)

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

  
Signature

16 MAY 2024



Stamp of the authority and date



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. Escientia Advanced Sciences Pvt. Ltd.,  
Plot No. 11, 11A, 12 & 12A, APSEZ,  
Atchutapuram, Anakapalli District -531011,  
Andhra Pradesh, India

List of APIs:

S. No.	Active substance(s)	Activity(ies)
01	(5S,6S,9R)-5-amino-6-(2,3-difluorophenyl)-6,7,8,9-tetrahydro-5H-cyclohepta [b] pyridin-9-yl 4-(2-oxo-2,3-dihydro-1H-imidazo [4,5-b] pyridin-1-yl) piperidine-1-carboxylate. Sulfuric acid salt (Rimegepant Sulfate/ BHV-3000) IH	Manufacturing & Packing

ITEM(S) ONE (01) 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.

The Written Confirmation remains valid until: 24.05.2027

  
Signature

16 MAY 2024

Stamp of the authority and date



**7-5/2021/EU/WC-0501**  
**Government of India**  
**Directorate General of Health Services**  
**Central Drugs Standard Control Organisation**  
**(International Cell)**

FDA Bhawan, Kotla Road,  
New Delhi-110002

Dated: 14 JUN 2024

To,

**M/s. Escientia Advanced Sciences Pvt Ltd.,**  
**Plot No. 11, 11A, 12 and 12A, APSEZ,**  
**Atchutapuram, Anakapalli District-531011,**  
**Andhra Pradesh, India**

**SUB:-** Written Confirmation of **M/s. Escientia Advanced Science Pvt Ltd., Plot No. 11, 11A, 12 and 12A, APSEZ, Atchutapuram, Anakapalli District-531011, 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/2024/7937 dated 19.01.2024 submitted to CDSCO, ADC(I), Visakhapatnam Sub Zone, and the recommendation received from ADC(I), Visakhapatnam Sub 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€, 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
--	--	16.05.2024	24.05.2027
01	01	16.05.2024	24.05.2027
02	01	14 JUN 2024	24.05.2027

Yours faithfully,

(Dr. Rajeev Singh Raghuvanshi)  
Drugs Controller General (India)



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

Annexure-02

WC-0501

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. Escientia Advanced Sciences Pvt Ltd.,  
Plot No. 11, 11A, 12 and 12A, APSEZ,  
Atchutapuram, Anakapalli District-531011,  
Andhra Pradesh, India

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1.	Nicardipine IH	Manufacturing & Packing

ITEM(S) ONE (01) ONLY

The Written Confirmation remains valid until: 24.05.2027

*Signature*

Stamp of the authority and date



14 JUN 2024