

**7-5/2013/EU/WC-0124**  
Government of India  
Directorate General of Health Services  
Central Drugs Standard Control Organisation  
(International Cell)

FDA Bhawan, Kotla Road,  
New Delhi-110002

Dated: **19 AUG 2025**

To,

**M/s. Aragen Life Sciences Private Limited**  
**28A, IDA Nacharam, Hyderabad, India,**  
**Nacharam (V), Uppal (M), Medchal – Malkajgiri (D),**  
**Telangana, India**

**SUB:-** Written Confirmation of **M/s. Aragen Life Sciences Private Limited, 28A, IDA Nacharam, Hyderabad, India, Nacharam (V), Uppal (M), Medchal – Malkajgiri (D), Telangana, 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/2025/9969** submitted to CDSCO, DDC(I), Hyderabad zone office, and the recommendation received from DDC(I), Hyderabad zone office, 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 The Written Confirmation will be withdrawn in the events of non-compliance of Standards.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
--	--	19 AUG 2025	02.07.2028
01	07	19 AUG 2025	02.07.2028
02	01	19 AUG 2025	02.07.2028

Yours faithfully,

  
19/08/25

**Ranga Chandrashekar**  
**Joint Drugs Controller (India)**

चंद्रशेखर रंगा/Chandrashekar Ranga  
जॉइंट ड्रग्स कंट्रोलर (भारत) / Joint Drugs Controller (India)  
जॉइंट ड्रग्स कंट्रोलर नियंत्रण, संगठन (मुख्यालय), स्वास्थ्य सेवा महाविद्यालय  
C.D.S.C.(HQ), Dte. General of Health Services  
स्वास्थ्य और परिवार कल्याण मंत्रालय / Ministry of Health and Family Welfare  
एफ.डी.ए. भवन, कोटा 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. Aragen Life Sciences Private Limited  
28A, IDA Nacharam, Hyderabad, India, Nacharam (V),  
Uppal (M), Medchal – Malkajgiri (D), Telangana, India

**2. Manufacturer's licence number:** 120/HD/AP/97/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:** 23.11.2024 & 24.11.2024

**The Written Confirmation remains valid until:** 02.07.2028

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 Chandrashekar,  
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

*Chandrashekar*  
Signature  
चंद्रशेखर रंगा/Chandrashekar Ranga  
संयुक्त औषधि नियंत्रक (भारत) / Joint Drugs Controller (India)  
औषधि और स्वास्थ्य सेवाएं / Ministry of Health and Family Welfare  
C.D.S.C. (HQ), Dte. General Services  
औषधि और स्वास्थ्य सेवाएं / Ministry of Health and Family Welfare  
फिरोज़ रोड, कोटला रोड, नई दिल्ली-110002 / FDA/PS/110002, Kotla Road, New Delhi-110002

Stamp of the authority and date  
  
19 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. Aragen Life Sciences Private Limited**  
**28A, IDA Nacharam, Hyderabad, India, Nacharam (V),**  
**Uppal (M), Medchal – Malkajgiri (D), Telangana, India**

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1.	Dobutamine Hydrochloride BP/USP/Ph.Eur.	Manufacturing & Packing
2.	Ezetimibe USP	Manufacturing & Packing
3.	Lamotrigine USP/Ph.Eur.	Manufacturing & Packing
4.	Trimetazidine Di Hydrochloride Ph.Eur.	Manufacturing & Packing
5.	Pentosan Polysulfate Sodium IH	Manufacturing & Packing
6.	Pralidoxime Chloride USP	Manufacturing & Packing
7.	Moxifloxacin Hydrochloride BP/Ph.Eur.	Manufacturing & Packing

Item(s) Seven (07) Only

The Written Confirmation remains valid until: 02.07.2028

*Chandrasekhar*  
Signature  
चंद्रशेखर रेणा/Chandrasekhar Rana  
संयुक्त औषधि नियंत्रक (भारत) / Joint Drugs Controller (India)  
केंद्रीय औषधि मानक नियंत्रण संगठन (मुख्यालय), स्वास्थ्य विभाग, भारत सरकार, नई दिल्ली-110002  
C.D.S.C.O.(HQ), Dte. General of Health Services, Ministry of Health & Family Welfare  
स्वास्थ्य और परिवार कल्याण विभाग / Ministry of Health & Family Welfare  
एन सी ए भवन, कोटला रोड, नई दिल्ली-110002 / FDA Bhawan, Kotla Road, Delhi-110002

Stamp of the authority and date  
  
19 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

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Uppal (M), Medchal – Malkajgiri (D), Telangana, India

List of APIs:

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

Item(s) One (01) Only

The Written Confirmation remains valid until: 02.07.2028

*Chandrashekar*  
Signature चंद्रशेखर रंगा/Chandrashekar Ranga  
संयुक्त औषधि नियंत्रक (भारत) / Joint Controller (India)  
केंद्रीय औषधि मानक नियंत्रण संगठन (भारत), स्वास्थ्य सेवा महानिदेशालय  
C.D.S.C.O.(HQ), Dte. of Health Services  
स्वास्थ्य और परिवार कल्याण मंत्रालय / Ministry of Health and Family Welfare  
एफ.डी.ए. भवन, कोटला रोड, नई दिल्ली-110002 / F.D.A. Bhawan, Kotla Road, New Delhi-110002

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19 AUG 2025