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DRUG

NATIONAL AUTHORITY

CERTIFICATE OF COMPLIANCE WITH GOOD MANUFACTURING PRACTICE GUIDELINES

THE NATIONAL DRUG POLICY AND AUTHORITY ACT, CAP 206

Issued under Regulation 19(5) of the National Drug Policy and Authority (Licensing) Regulations, 2014

Certificate No. 063/GMP/2025

This is to certify that the drug manufacturing facility:

Name of facility: **Ahlcon Parenteral (India) Limited**

Physical address of facility: SP-917 & 918, Phase III, RIICO Industrial Area, Bhiwadi, Dist.Alwar (Rajasthan) India.

License number of the manufacturer: Raj. 1594 on Form 28, issued by Drugs Control Organization, Rajasthan with a validity of up to 31/12/2026.

Has been inspected by the Authority for compliance with the Good Manufacturing Practice Guidelines.

On the basis of the inspection carried out on **3rd and 4th October 2024,** it is certified that the facility indicated on this certificate complies with Good Manufacturing Practice for dosage forms listed in Table 1 below.

Table 1: Approved lines.

No.	Dosage Form	Category	Activities
1.	Small volume parenteral products (Aseptically prepared)	Non-Beta Lactam	Manufacture of Finished Pharmaceutical (medicinal) Product
2.	Small volume parenteral products (Terminally sterilized)		
3.	Large volume parenteral products (Terminally sterilized)		
4.	Eye/Ear drops (Aseptically prepared)		

The responsibility for the quality of the individual batches of the drugs manufactured through this process lies with the manufacturer.

This certificate remains valid until **4th October 2027.** It becomes invalid if the activities or the categories certified change or if the facility is no longer considered to be in compliance with GMP.

Issue Date: 16th April 2025.

NATIONAL DRUG AUTHORITY
16 APR 2025
PLOT 6A BUGANDA ROAD
P. O. BOX 23052 KAMPALA

Solomon Orem
FOR THE AUTHORITY

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