

308063



**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. 043/GMP/2025**

This is to certify that the drug manufacturing facility:

**Name of facility:** West Coast Pharmaceutical Works Limited.

**Physical address of facility:** Nr. Bhagwati Glass Factory, Opp. Jogni Mata Temple,  
Denis Chem Lab, Vadaswami Side Road,  
Chattral-Vadaswami Road, Dist Gandhinagar, Gujarat-  
382740, India.

**License number of the manufacturer:** G/28/1894 and G/25/2608  
Issue date 28.12.22 valid till 27.12.2027.

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

On the basis of the inspection carried out on **27<sup>th</sup>, 28<sup>th</sup> and 29<sup>th</sup> May 2024**, it is certified that the facility indicated on this certificate complies with Good Manufacturing Practice for dosage forms listed in Table 1 below.

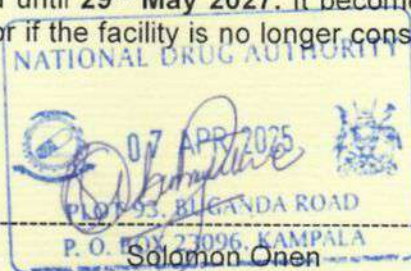
**Table 1:** Approved lines.

| No. | Dosage Form                 | Category                        | Activities   |
|-----|-----------------------------|---------------------------------|--|
| 1.  | Tablets (Coated & Uncoated) | Non-Beta<br>Lactam<br>(General) | Manufacture of<br>Controlled/Modified<br>Finished Pharmaceutical<br>(medicinal) Products |
| 2.  | Capsules                    |                                 |  |
| 3.  | Creams, Ointments and Gels  |                                 |  |
| 4.  | External powders            |                                 |  |
| 5.  | Liquids for internal use    |                                 |  |
| 6.  | Liquids for external use    |                                 |  |
| 7.  | Tablets (Coated & Uncoated) | Hormonal                        |  |
| 8.  | Creams, Ointments and Gels  |                                 |  |

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 **29<sup>th</sup> May 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:** 7<sup>th</sup> April 2025.



**FOR THE AUTHORITY**

