REPUBLIC OF KENYA

MINISTRY OF HEALTH

PHARMACY AND POISONS BOARD

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Pharmacy and Poisons Board Building
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P.O Box 27663-00506
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GMP CERTIFICATE No: [redacted]

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

PART 1

Issued in accordance with Section 35B of the Pharmacy and Poisons Act (Cap 244) of the Laws of Kenya.

The Pharmacy and Poisons Board, The National Medicines Regulatory Authority of Kenya, confirms the following:

The manufacturer: [redacted]

Site address: [redacted]

From the knowledge gained during the inspection of this manufacturer, the latest of which was conducted on [redacted], GMP Report No. [redacted], the site complies with the prescribed Good Manufacturing Practices as per the relevant WHO Technical Report Series and other internationally acceptable guidelines.

This certificate reflects the compliance status of the manufacturing site as at the time of the inspection above and will be deemed to be valid until [redacted] after which time the Pharmacy and Poisons Board should be consulted.

PART 2

Human Medicinal Products

1. Manufacturing operations authorised/subject to inspection*

1.1 Sterile Products: N/A

1.1.1 Aseptically prepared: N/A
1.1.2 Terminally sterilised: N/A
1.1.3 Testing or batch release only: N/A

1.2 Non-sterile products

DOSAGE FORM	CATEGORY	PRODUCT TYPE	ACTIVITIES
(a) Oral Solids	General	Tablets, Capsules, Dry Syrups	Processing operations, batch certification, packaging, quality control testing
(b) Oral liquids	General	Syrups, Suspensions	
(c) External Preparations	General	Creams, Ointments, Gels, Solutions	

1.2.1 Testing and batch release only: N/A

1.3 Biological medicinal products: N/A

1.4 Other products or manufacturing activity: N/A

1.5 Packaging only: N/A

1.1.1 Primary packing: N/A
1.1.2 Secondary packing: N/A

1.6 Quality Control testing: N/A

1.7 Blinding: N/A

The compliance status shall be deemed valid unless it is invalidated under any of the following conditions;

1. The activities and/or categories certified herewith are changed.
2. The site is no longer considered to be in compliance with GMP.
3. The manufacturing site is changed.

Any restrictions or clarifying remarks related to the scope of this certificate. YES/**NO**

This certificate is valid only when presented with all pages and both parts 1 and 2.

The authenticity of this certificate may be verified with the Kenya Pharmacy and Poisons Board.

[redacted]

Date: 24th March 2025

[redacted]

REGISTRAR/CHIEF EXECUTIVE OFFICER
PHARMACY AND POISONS BOARD