307808

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. 228/GMP/2021

This is to certify that the drug manufacturing facility:

Name of facility: Kampala Pharmaceutical Industries (1996) Limited.

Physical address of facility: Stretcher Road, Plot M444B, Ntinda, Kampala.

License number of the manufacturer: NDA/PRE/PMC/2367.

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

On the basis of the assessment carried out on **21st December 2021,** 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. 2. 3.	Tablets (Coated & Uncoated) Capsules (Hard gelatin) Creams & Ointments	Non-Beta Lactam	Manufacture of Finished Pharmaceutical Product
4. 5.	Tablets (Coated & Uncoated) Capsules (Hard gelatin)	Beta-Lactam (Penicillin)	
6.	Secondary packaging (Injectaplan)	Non-Beta Lactam	Secondary Packing

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 **21st December 2024.** 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: 05th January 2022.

NATIONAL DRUG AUTHORITY
05 JAN 2022
PLOT 19 LUMUMBA AVENUE
P.O. BOX 23006 KAMPALA

Denis Mwesigwa
FOR THE AUTHORITY