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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. 205/GMP/2025

This is to certify that the drug manufacturing facility:

Name of facility: **Shishi International Limited**

Physical address of facility: Plot 2388/110 Kyadondo Block 211. Mawato Namanve Tirupati park view complex, Kampala - Uganda.

License number of the manufacturer: NDA/PRE/PM/0078.

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

On the basis of the inspection carried out on **23rd August 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. 2. 3.	Laparoscope system, re-usable Respiratory oxygen therapy system Externally - anchored laparoscopic retractor	Class A	Complete Manufacture of Surgical Instruments & Appliances

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 **22nd August 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: 6th October 2025.

NATIONAL DRUG AUTHORITY

06 OCT 2025

PLOT 93 BUGANDA ROAD

P. O. BOX 23090 KAMPALA

Denis Mwesigwa

FOR THE AUTHORITY