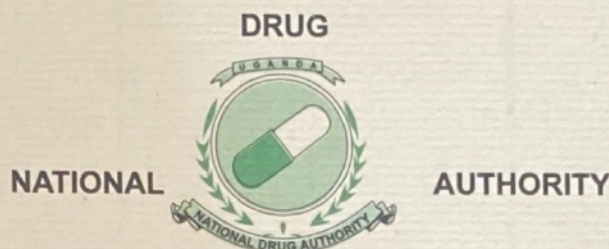


274821



**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.	Laparoscope system, re-usable	Class A	Complete Manufacture of Surgical Instruments & Appliances
2.	Respiratory oxygen therapy system		
3.	Externally - anchored laparoscopic retractor		

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.

