307879

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. 128/GMP/2023

This is to certify that the drug manufacturing facility:

Name of facility: Labor LS SE & Co. KG.

Physical address of facility: Mangelsfeld 4,5,6 97708 Bad Bocklet - Grossenbrach Germany.

License number of the manufacturer: DE_BY_05_MIA_2022_0050/55.2-2678.4-1-52-3.

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

On the basis of the inspection carried out on **17th and 18th October 2023,** 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.	Not Applicable	Not Applicable	Microbiology Testing

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 **18th October 2026.** 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: 30th November 2023.

NATIONAL DRUG AUTHORITY
30 NOV 2023
PLOT 19 LUMUMBA AVENUE
P. O. BOX 23096, KAMPALA

Nasser Lubowa
FOR THE AUTHORITY