TANZANIA MEDICINES AND MEDICAL DEVICES AUTHORITY

TMDA
Tanzania Medicines & Medical Devices Authority

CERTIFICATE OF COMPLIANCE TO GOOD MANUFACTURING PRACTICES (GMP)

(For Medicines Manufacturing Facilities)
(Made under Section 20 (2a) of Tanzania Medicines and Medical Devices Act, Cap 219)

GMP Certificate No. TMDA0019/GMP/O/D/0110

This is to certify that M/S BDA HEALTHCARE PRIVATE LIMITED located at PLOT NO.B-1, B-2, B-3, NEAR ITI, M.I.D.C PARSEONI 441105, DIST NAGPUR, MAHARASHTRA, INDIA has been found to comply with current Good Manufacturing Practice requirements for dosage forms and categories of medicines listed below:

S/N	Dosage Form(s)	Categories of Medicines	Manufacturing operations
1.	Tablets and Capsules	General Pharmaceutical Formulations	Manufacturing and Packaging
2.	Sachets	General Pharmaceutical Formulations	Manufacturing and Packaging

The responsibility for the quality of the individual batches of pharmaceutical products manufactured lies with the manufacturer and /or local agent.

Issued On: December 19, 2019

Expires On: December 18, 2022

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A. M. FIMBO

ACTING DIRECTOR GENERAL

Note: This certificate shall be invalid if the forms and operations herewith are changed or if the site is no longer considered to be in compliance with current GMP requirements.