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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. 069/GMP/2023

This is to certify that the drug manufacturing facility:

Name of facility: Bafna Pharmaceuticals Limited, Unit II.

Physical address of facility: 147 Madhavaram Red Hills Road, Grantlyon Village, Vadakarai, Chennai, In 600052, India.

License number of the manufacturer: TN00002269 valid 31.12.2024 TN00002270 valid 31.12.2024.

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

On the basis of the inspection carried out on **25th and 26th October 2022**, 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.	Tablets	Non-Beta Lactam	Manufacture of Finished Pharmaceutical
2.	Hard gelatin capsules		(medicinal) Product- Human

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 **26th October 2025.** 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: 16th June 2023 NATIONAL DRUG AUTHORIT PLOT 19 IDEHIS/IVIN/86FOWE P. O. FOR2THE AUTHORIT

