DEPARTMENT OF FOOD SAFETY AND DRUGS CONTROL ADMINISTRATION
GOVERNMENT OF TAMILNADU

359, Anna Salai, Chennai - 600 006, Tamil Nadu.

CERTIFICATE OF GOOD MANUFACTURING PRACTICES*

Certificate No: **K Dis. No: 19014/D1/4/2021, Dated: 14.09.2022**

On the basis of the inspection carried out on 10.02.2022, 11.02.2022 and 27.07.2022, It is certified that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in Table 1.

1 Name and address of site: M/s. Bafna Pharmaceuticals Limited, No.1/15B, New No. 147, Survey No. 30/3, 30/4, 30/5, 30/6, 30/7, & 30/12, Madhavaram Redhills High Road, Grantylon Village, Vadakarai Post, Chennai – 600 052.

2 Manufacturer's licence number: Form 25 Bearing No. TN00002269 Dated: 08.12.2006
Form 28 Bearing No. TN00002270 Dated: 08.12.2006

3 Table 1:

Dosage form(s)	Category(ies)	Activity(ies)
Tablets & Capsules (Vide List Attached)	General (Other Than Betalactum, sex hormones & Cytotoxic)	Manufacturer

The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.

This certificate remains valid until 31.12.2024 It becomes invalid if the activities and/or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

Name and function of responsible person: **P.V. Vijayalakshmi, B.Pharm.,**

14/9/22
Director of Drugs Control (i/c)
14/9/2022 Tamil Nadu.

Email:tndcad@gmail.com **Telephone No**: 91-44-2433 5068 **Fax No.**:91-44-2432 1830

*This certificate is issued as per WHO norms

P.V. VIJAYALAKSHMI,
Director of Drugs Control (i/c)
359, Anna Salai, Chennai - 600 006.

Explanatory notes

1) This certificate, which is in the format recommended by WHO, certifies the status of the Site listed in point 1 of the certificate.
2) The certification number should be traceable within the regulatory authority issuing the certificate.
3) Where the regulatory authority issues a licence for the site this number should be specified. Record "not applicable" in case where there is no legal framework for the issuing of a licence.
4) The certificate remains valid until the specified date. The certificate becomes invalid if the activities and/or categories certified are changed or if the site is no longer considered to be in compliance with GMP.
5) The requirements for good practices in the manufacture and quality control of drugs referred to in the certificate are those included in Quality Assurance of Pharmaceuticals: a compendium of guidelines and related materials. Good manufacturing practices and inspection, Volume 2, 1999. World Health Organization, Geneva and subsequent updates.