

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) Table 1
List the dosage forms, starting materials, categories and activities.
Examples give below.

Pharmaceutical	Category(ies)	Activity(ies)
Dosage form(s):		
Tablets	Cytotoxic	Packaging
	Hormone	Production, packaging, quality control
	Penicillin	Repackaging and labelling
Injectables	Cefalosporin	Aseptic preparation, packaging, labelling

Example 2

Pharmaceutical Product(s) ²	Category(ies)	Activity(ies)
Starting material(s) ³		
Paracetamol	Analgesic	Synthesis, purification, packing, labelling

2 Pharmaceutical Products: Any medicine intended for human use or veterinary product administered to food-producing animals, presented in its finished dosage form or as a starting material for use in such a dosage form, that is subject to control by pharmaceutical legislation in both the exporting state and the importing state.

3 Starting Materials: Any substance of a defined quality used in the production of pharmaceutical product, but excluding packaging materials. Use, whenever available, International Nonproprietary Names (INNs) or otherwise national nonproprietary names.

- (5) 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.
- (6) 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.



**GOVERNMENT OF KARNATAKA
DRUGS CONTROL DEPARTMENT**

No : FDA/SPL.CELL-I/CR-587/2025-26
GSC No: DD008S25000087

Office of the Drugs Controller
For the State of Karnataka
P.B. No. 5377 ,Palace Road,
Bangalore - 560001
Date: 08/08/2025

WHO GMP CERTIFICATE

This one page certificate conforms to the format recommended by the World Health Organization

(General Instructions and explanatory notes are attached)

On the basis of the inspection carried out on M/s : Suraj Laboratories Private Limited we certify that the site indicated on this certificate complies with Good Manufacturing Practices(GMP) for the dosage form categories and activities listed on

Table 1

01.Name and Address of the site : Suraj Laboratories Private Limited

Plot No:540/3P1,540/4 and 540/5, KIADB Kadechur Industrial Area, Kadechur, Yadgir District, Karnataka India-585221

02.Manufacturers Licence Numbers : KTK/25/0861/2024

03.Table 1:

S. No.	Pharmaceutical Products / Dosage forms	Category(ies)	Activity(ies)
1.	Fenofibrate Ph.Eur	Antilipemic	Manufacturing, Quality Control, Packing, Labeling
2.	Fenofibrate BP	Antilipemic	Manufacturing, Quality Control, Packing, Labeling
3.	Fenofibrate IP	Antilipemic	Manufacturing, Quality Control, Packing, Labeling
4.	Fenofibrate USP	Antilipemic	Manufacturing, Quality Control, Packing, Labeling

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

This certificate remains valid upto 28/05/2028 . It become invalid if the activities and or categories certified herewith are changed or if the site is on longer considered to be in compliance with GMP.

This certificate is issued as per WHO TRS No.908 of 2003.

Address of Certifying authority :Office of the Drugs Controller,For the State of Karnataka ,Palace Road,Bangalore - 560001,INDIA.

Name of the Authorized
Person: Dr.Umesh.S

Email:dckarnataka@gmail.com



Signature Not Verified

Digitally signed by Umesh S
Date: 2025.08.08 18:36:09 IST

The Drugs Controller &
Licensing Authority