By Regd. Post

K. Dis. No.013/D1/4/2019

Office of the Director of Drugs Control,
Tamil Nadu, Chennai – 600 006.
Dated: 22.01.2020.

Sub: Drugs – Drugs & Cosmetics act 1940 rules made there under- M/s. Micro Lab Limited, Unit – 3, No. 92, SIPCOT Industrial Complex, Hosur – 635 126, Krishnagiri District - Application for revalidation of WHO-GMP Certificate on WHO Norms under in Form 25 and Form 28 for a period from 2019-2021 – Issued - Reg.

Ref:
1. Application Dated: 28.12.2018 from M/s. Micro Lab Limited, Unit – 3, No. 92, SIPCOT Industrial Complex, Hosur – 635 126, Krishnagiri District.
2. Letter No. 50-3(201)/SZ/2019/533 Dated: 14.05.2019 of the Deputy Drugs Controller (I), CDSCO South Zone, Chennai – 600 006.
3. Ref. No. 358/D3/2019 dated: 13.05.2019, 21.08.2019 & 01.11.2019 of the Assistant Director of Drugs Control, Salem Zone.

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M/s. Micro Lab Limited, Unit – 3, No. 92, SIPCOT Industrial Complex, Hosur – 635 126, Krishnagiri District are informed that issuance of Good Manufacturing Practice Certificate under WHO norms in Form 25 & 28, bearing Nos. TN00003934 & TN00003935 both dated: 19.10.2015 are issued for the period 2019-2021.

The receipt of this certificate shall be acknowledged early

DIRECTOR OF DRUG CONTROL

22/1/2020

To
M/s. Micro Lab Limited,
Unit – 3, No. 92, SIPCOT Industrial Complex,
Hosur – 635 126, Krishnagiri District.

Copy to: The Assistant Director of Drug Control,
Salem Zone.

Digitally signed by Chirtoacă Iurie
Date: 2021.11.30 15:27:50 EET
Reason: MoldSign Signature
Location: Moldova

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: 013/D1/4/2019, Dated: 22.01.2020**

On the basis of the inspection carried out on 11.03.2019, 12.03.2019 and 24.10.2019 we certify 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. Micro Lab Limited,
Unit – 3, No. 92, SIPCOT Industrial Complex,
Hosur – 635 126, Krishnagiri District.

2 Manufacturer's licence number: Form 25 Bearing No: TN00003934, Dated: 19.10.2015.
Form 28 Bearing No. TN00003935, Dated: 19.10.2015.

3 Table 1:

Dosage form(s)	Category(ies)	Activity(ies)
Tablets and Hard Gelatin 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 31.12.2021 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: K.Sivabalan, B.Pharm.,
Director of Drugs Control
Tamil Nadu.

K. SIVABALAN B.Pharm
Director of Drugs Control
359, Anna Salai, Chennai - 600 006.

22/1/2020

DIRECTORATE OF DRUGS CONTROL CHENNAI-6

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

***This certificate is issued as per WHO norms**

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.