

TNSWP Case No.: FSDAD-14381-2025  
Application ID : 343732

Office of the Director of Drugs Control,  
Tamil Nadu, Chennai – 600 006.  
Dated: 23/07/2026

Sub: Drugs – Drugs & Cosmetics act 1940 rules made there under M/s.Quantum Drugs & Chemicals , C-21, SIPCOT Industrial Complex, Manamadurai, 630606, Manamadurai Taluk, Sivagangai District, Tamil Nadu Application for Revalidation of WHO-GMP Certificate on WHO Norms under licence in Form 25 bearing No. 936/2012 dated: 11/06/2015 are approved for the period from 01/01/2026 to 31/12/2028 – Issued - Reg.

Ref: Online Application Dated:15/11/2025 of M/s.Quantum Drugs & Chemicals , C-21, SIPCOT Industrial Complex, Manamadurai, 630606, Manamadurai Taluk, Sivagangai District, Tamil Nadu .

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M/s.Quantum Drugs & Chemicals , C-21, SIPCOT Industrial Complex, Manamadurai, 630606, Manamadurai Taluk, Sivagangai District, Tamil Nadu are informed that Revalidation of Good Manufacturing Practice Certificate under WHO norms in Form 25 bearing No. 936/2012 dated: 11/06/2015 are approved for the period from 01/01/2026 to 31/12/2028.

The receipt of this certificate shall be acknowledged early.

Signature valid

Signed by: GURUBHAKATHI S,  
State Licensing Authority,  
Joint Director of Drugs Control and  
Administration,  
Controlling Authority for Licensing Authority  
Date:23-Jul-2026 15:26:45



**DEPARTMENT OF FOOD SAFETY AND DRUGS CONTROL ADMINISTRATION  
GOVERNMENT OF TAMILNADU**

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

**CERTIFICATE OF GOOD MANUFACTURING PRACTICES**

CertificateNo:TN/WHO/00065

**Dated:23/07/2026**

On the basis of the inspection carried out on 19/03/2026 to 20/03/2026 and 26/05/2026 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.Quantum Drugs & Chemicals , C-21, SIPCOT Industrial Complex, Manamadurai, 630606, Manamadurai Taluk, Sivagangai District, Tamil Nadu
- 2 Manufacturer"s licence number: Form 25 bearing No. 936/2012 dated: 11/06/2015
- 3 Table 1:

| Dosage form(s)                   | Category(ies)                                             | Activity(ies) |
|----------------------------------|-----------------------------------------------------------|---------------|
| Active Pharmaceutical Ingredient | 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 31/12/2028 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:

GURUBHARATHI S

Joint Director of Drugs Control and  
Controlling Authority cum Licensing Authority

**Email:**tndcad@gmail.com

**Telephone No.:** 91-44-2433 5068

**Fax No.:**91-44-2432 1830

Signature valid

**Joint Director of Drugs Control and  
Controlling Authority cum Licensing Authority**  
State Licensing Authority  
Food Safety and Drug  
Administration  
Date:23-Jul-2026 15:26:45

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

List of drugs already permitted in Form 25 bearing No. 936/2012 dated: 11/06/2015 of M/s.Quantum Drugs & Chemicals, C-21, SIPCOT Industrial Complex, Manamadurai, 630606, Manamadurai Taluk, Sivagangai District, Tamil Nadu, are included under WHO GMP certificate which is valid upto 31/12/2028.

**WHO GMP Certificate No.TN/WHO/00065**

| Sr. No. | Details of Product                                                                                                                                   |
|---------|------------------------------------------------------------------------------------------------------------------------------------------------------|
|         | Bulk Drugs                                                                                                                                           |
| 1       | Proper Name of Product : Chlorpropamide BP<br>Brand/Trade Name : -<br>Dosage Forms : Active Pharmaceutical Ingredient<br>For Export Purpose Only     |
|         | Bulk Drugs                                                                                                                                           |
| 2       | Proper Name of Product : Chlorpropamide Ph.Eur<br>Brand/Trade Name : -<br>Dosage Forms : Active Pharmaceutical Ingredient<br>For Export Purpose Only |
|         | Bulk Drugs                                                                                                                                           |
| 3       | Proper Name of Product : Chlorpropamide USP<br>Brand/Trade Name : -<br>Dosage Forms : Active Pharmaceutical Ingredient<br>For Export Purpose Only    |
|         | Bulk Drugs                                                                                                                                           |
| 4       | Proper Name of Product : Tolbutamide USP<br>Brand/Trade Name : -<br>Dosage Forms : Active Pharmaceutical Ingredient<br>For Export Purpose Only       |
|         | Bulk Drugs                                                                                                                                           |
| 5       | Proper Name of Product : Tolbutamide BP<br>Brand/Trade Name : -<br>Dosage Forms : Active Pharmaceutical Ingredient<br>For Export Purpose Only        |
|         | Bulk Drugs                                                                                                                                           |
| 6       | Proper Name of Product : Tolbutamide Ph.Eur<br>Brand/Trade Name : -<br>Dosage Forms : Active Pharmaceutical Ingredient<br>For Export Purpose Only    |

Case No. FSDAD-14381-2025; Application ID-343732 Dated 23/07/2026

Signature valid

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State Licensing Authority,  
Joint Director of Drugs Control and  
Administration,  
Controlling Authority and Licensing Authority  
Date: 23-Jul-2026 15:26:45

Genuineness of this certificate can be verified by using the unique Certificate number/Token ID/application ID in the Website of Tamil Nadu Single Window Portal <https://tnswp.com/DIGIGOV/swp-tnswp.jsp>

## 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.

Signature valid

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State Licensing Authority,  
Food Safety and Drug  
Administration  
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