TNSWP Case No. FSDAD-14403-2025
Application ID : 344814

Office of the Director of Drugs Control,
Tamil Nadu, Chennai - 600 006.
Dated: 02/04/2026

Sub: Drugs - Drugs & Cosmetics act 1940 rules made there under M/s Sai Sarvaa Biotech Private Limited, Plot No. 8 and 9, Auroville Post, Balaji Nagar, Pattanur, 605101, VANUR Taluk, Villupuram District, Tamil Nadu Application for Issuance of WHO-GMP Certificate on WHO Norms under licence in Form 25 bearing No. TN00003900 dated: 02/09/2015 & Form 28 bearing No. TN00003901 dated: 02/09/2015 are approved for the period from 02/04/2026 to 01/04/2029 - Issued - Reg.

Ref: Application Dated:17/11/2025 of M/s.Sai Sarvaa Biotech Private Limited, Plot No. 8 and 9, Auroville Post, Balaji Nagar, Pattanur, 605101, VANUR Taluk, Villupuram District, Tamil Nadu.

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M/s.Sai Sarvaa Biotech Private Limited, Plot No. 8 and 9, Auroville Post, Balaji Nagar, Pattanur, 605101, VANUR Taluk, Villupuram District, Tamil Nadu are informed that issuance of Good Manufacturing Practice Certificate under WHO Norms under licence in Form 25 bearing No. TN00003900 dated: 02/09/2015 & Form 28 bearing No. TN00003901 dated: 02/09/2015 are approved for the period from 02/04/2026 to 01/04/2029

The receipt of this certificate shall be acknowledged early.

Signature Not Verified

Signed by: G.PERUBHARATHI.S
State Licensing Authority,
Food Safety and Drug Administration
Date:02-Apr-2026 16:09:13

Joint Director of Drugs Control and
Controlling Authority cum Licensing Authority

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

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/00049 **Dated:**02/04/2026

On the basis of the inspection carried out on 05.02.2026 to 06.02.2026 and 12.03.2026. 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: Sai Sarvaa Biotech Private Limited,
Plot No. 8 and 9, Auroville Post, Balaji Nagar, Pattanur, 605101, VANUR Taluk,
Villupuram District, Tamil Nadu

2 Manufacturer"s licence number: Form 25 bearing No. TN00003900 dated: 02/09/2015 &
Form 28 bearing No. TN00003901 dated: 02/09/2015

3 Table 1:

Dosage form(s)	Category(ies)	Activity(ies)
Tablets	General (Other than Betalactum, Sex Hormones & Cytotoxic)	Manufacturer
Hard gelatin capsules	General (Other than Betalactum, Sex Hormones & Cytotoxic)	Manufacturer
Tablets	Betalactum	Manufacturer
Hard gelatin capsules	Betalactum	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 01/04/2029 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 Not Verified

Joint Director of Drugs Control and
Controlling Authority cum Licensing Authority

Signed by: GURUBHARATHI S,
State Licensing Authority,
Food Safety and Drug Administration
Date:02-Apr-2026 16:09:13

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*** 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 â€œnot a pplicableâ€ 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 Not Verified

Signed by: GURUBHARATHI S,
State Licensing Authority,
Food Safety and Drug Administration
Date:02-Apr-2026 16:09:13

List of already permitted drugs in Form 25 bearing No. TN00003900 dated: 02/09/2015 of M/s.Sai Sarvaa Biotech Private Limited, Plot No. 8 and 9, Auroville Post, Balaji Nagar, Pattanur, 605101, VANUR Taluk, Villupuram District, Tamil Nadu, are included under WHO GMP certificate which is valid upto 01/04/2029.

WHO GMP Certificate No TN/WHO/00049

Sr. No.	Details of Product
	Tablets
1	Proper Name of Product : Aceclofenac 100mg and Paracetamol 325mg tablets IP Composition : Each Flim Coated Tablet Contains : Aceclofenac IP 100mg, Paracetamol IP 325mg. Colour : Approved colour Used Brand/Trade Name : - Dosage Forms : Tablets
	Capsules
2	Proper Name of Product : Acebrophylline Capsules 100mg Composition : Each Hard Gelatin Capsules Contains : Acebrophylline 100mg Colour : Approved Colour Used In Empty Capsules Shells Brand/Trade Name : - Dosage Forms : capsule

Joint Director of Drugs Control and
Controlling Authority cum Licensing Authority

Signature Not Verified
Signed by: GURUBHARATHI S,
State Licensing Authority,
Food Safety and Drug Administration
Date:02-Apr-2026 16:09:13

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List of already permitted drugs in Form 28 bearing No. TN00003901 dated: 02/09/2015 of M/s.Sai Sarvaa Biotech Private Limited, Plot No. 8 and 9, Auroville Post, Balaji Nagar, Pattanur, 605101, VANUR Taluk, Villupuram District, Tamil Nadu, are included under WHO GMP certificate which is valid upto 01/04/2029.

WHO GMP Certificate No TN/WHO/00049

Sr. No.	Details of Product
	Tablets
1	Proper Name of Product : Cefixime Dispersible Tablets 200mg Composition : Each Uncoated Dispersible Tablets Contains : Cefixime Trihydrate IP Equivalent to Anhydrous Cefixime 200mg, colour : Approved Colour Used Brand/Trade Name : - Dosage Forms : Tablets
	Capsules
2	Proper Name of Product : Amoxycillin Capsules IP 500mg Composition : Each Hard Gelatin Capsules Contains : Amoxycillin Trihydrate IP Equivalent to Amoxycillin 500mg, Colour : Approved Colour Used in Empty Capsules Shells Brand/Trade Name : - Dosage Forms : Capsules

Signature Not Verified

Signed by: GURUBHARATHI S,
State Licensing Authority,
Food Safety and Drug Administration
Date:02-Apr-2026 16:09:13

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