

**Wanbury Limited**

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12th May, 2025

National Stock Exchange of India Ltd.
Exchange Plaza, Plot no. C/1, G Block,
Bandra-Kurla Complex, Bandra (E)
Mumbai - 400 051
Security Symbol: WANBURY

BSE Limited
Phiroze Jeejeebhoy Towers,
Dalal Street
Mumbai- 400 001
Security Code: 524212

Dear Sirs,

Sub.: Grant of Good Manufacturing Practices Certificate (CBPF) by ANVISA for its Tanuku facility, Andhra Pradesh.

We are pleased to inform you that, pursuant to Regulation 30 (6) of SEBI (Listing Obligations and Disclosure Requirements) Regulations, Wanbury Limited's, Andhra Pradesh Tanuku facility has successfully completed the ANVISA (Brazil FDA) inspection with zero observations.

The inspection conducted from December 16, 2024 to December 20, 2024 by the Brazil Health Regulatory Agency (ANVISA) and confirmed strict adherence to cGMP (Current Good Manufacturing Practices).

As a result, ANVISA has granted the Good Manufacturing Practices (GMP) Certificate for the facility and the Good Manufacturing Practices Certificate (CBPF) issued by ANVISA in English and Portuguese Languages are attached.

Thanking you.

Yours truly,
For Wanbury Limited

Jitendra J. Gandhi
Company Secretary





MINISTRY OF HEALTH

NATIONAL HEALTH SURVEILLANCE AGENCY

CERTIFICATE OF GOOD MANUFACTURING PRACTICES PHARMACEUTICAL SUPPLIES

The National Health Surveillance Agency - ANVISA, through Resolution RE No. **295**, of **01/23/2025**, published in the Official Gazette of the Union (DOU) on the date of **01/27/2025**, certifies that the company below is periodically inspected and monitored by the National Health Surveillance System and that it complies with the Good Manufacturing Practices guidelines given by Brazilian legislation, which is in line with the recommendations of the World Health Organization.

Manufacturer: Wanbury Limited

Address: Doctors Organic Chemicals Division, K. Illindalaparru - 534 217, Iragavaram Mandal, West Godavari District, Andhra Pradesh

Country: India Unique Code: B.000306 File

Number(s): 4764962/22-5, 0237969/25-4

Certificate of Good Manufacturing Practices for Active Pharmaceutical Ingredients:

Active pharmaceutical ingredients obtained by chemical synthesis:
diphenhydramine hydrochloride, metformin hydrochloride, paroxetine hydrochloride hemihydrate, sertraline hydrochloride, tramadol hydrochloride.

Certificate already including the changes made by the correction published in the DOU of the day **05/05/2025**.

This certification is valid until the day **01/27/2027** and may be cancelled if the competent health authority proves that the requirements set out in current Good Practice standards have not been met.



Document signed electronically by **Marcus Aurelio Miranda de Araujo, General Manager of Sanitary Inspection and Supervision**, on 05/05/2025, at 2:48 pm, according to Brasília official time, pursuant to § 3 of art. 4 of Decree No. 10,543, of November 13, 2020 http://www.planalto.gov.br/ccivil_03/_ato2019-2022/2020/decreto/D10543.htm .



The authenticity of this document can be verified on the website <https://sei.anvisa.gov.br/authenticity> , informing the verification code **3572342** and the CRC code **FEFC7BB0**.



MINISTÉRIO DA SAÚDE

AGÊNCIA NACIONAL DE VIGILÂNCIA SANITÁRIA

CERTIFICADO DE BOAS PRÁTICAS DE FABRICAÇÃO DE INSUMOS FARMACÊUTICOS

A Agência Nacional de Vigilância Sanitária - ANVISA, por meio da Resolução RE nº **295**, de **23/01/2025**, publicada em Diário Oficial da União (DOU) na data de **27/01/2025**, certifica que a empresa abaixo é periodicamente inspecionada e monitorada pelo Sistema Nacional de Vigilância Sanitária e que cumpre com as diretrizes de Boas Práticas de Fabricação dadas pela legislação brasileira, a qual está em consonância com as recomendações da Organização Mundial de Saúde.

Fabricante: Wanbury Limited

Endereço: Doctors Organic Chemicals Division, K. Illindalaparru - 534 217, Iragavaram Mandal, West Godavari District, Andhra Pradesh

País: Índia Código único: B.000306

Expediente(s): 4764962/22-5, 0237969/25-4

Certificado de Boas Práticas de Fabricação de Insumos Farmacêuticos Ativos:

Insumos farmacêuticos ativos obtidos por síntese química: cloridrato de difenidramina, cloridrato de metformina, cloridrato de paroxetina hemi-hidratado, cloridrato de sertralina, cloridrato de tramadol.

Certificado já constando as alterações realizadas pela retificação publicada no DOU do dia **05/05/2025**.

A presente certificação é válida até o dia **27/01/2027** e poderá ser cancelada, caso seja comprovado, pela autoridade sanitária competente, o não cumprimento dos requisitos preconizados pelas normas vigentes de Boas Práticas.



Documento assinado eletronicamente por **Marcus Aurelio Miranda de Araujo, Gerente-Geral de Inspeção e Fiscalização Sanitária**, em 05/05/2025, às 14:48, conforme horário oficial de Brasília, com fundamento no § 3º do art. 4º do Decreto nº 10.543, de 13 de novembro de 2020
http://www.planalto.gov.br/ccivil_03/_ato2019-2022/2020/decreto/D10543.htm.



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