



REPUBLIC OF TÜRKİYE
MINISTRY OF HEALTH
MEDICINES AND MEDICAL
DEVICES AGENCY OF TÜRKİYE

Certificate No: TR/GMP/2026/59

CERTIFICATE OF GMP COMPLIANCE OF MANUFACTURER

Part 1

Issued following an inspection in accordance with current Good Manufacturing Practice Guidelines, and the Regulation on Manufacturing Plants of Medicinal Products for Human Use* and the Law No 1262 on Pharmaceutical and Medicinal Preparations. These regulations are in line with the requirements of Pharmaceutical Inspection Co-operation Scheme (PIC/s) and the Directives of the European Commission.

Turkish Medicines and Medical Devices Agency confirms the following:

Manufacturer's Name : PHARMA CARE SAĞLIK HİZMETLERİ LOJİSTİK
ANONİM ŞİRKETİ
Head Office / Correspondence Address : ORUÇREİS MAH. VADİ CAD. İSTANBUL TİCARET
SARAYI NO: 108 İÇ KAPI NO: 185 ESENLER/İSTANBUL
Esenler İstanbul
Site Address : Gebze Dilovası OSB Mahallesi D-4012 Sok. No:3 Gebze
Kocaeli
Manufacturing Authorization Date : 11.02.2026
Manufacturing Authorization Number : TR/SAY/2025/11-1

Has been inspected in accordance with current Good Manufacturing Practice Guidelines, the Regulation on Manufacturing Plants of Medicinal Products for Human Use, the Law No 1262 on Pharmaceutical and Medicinal Products.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 30.09.2025, it is considered that it complies with the requirements of Good Manufacturing Practice (GMP).

This certificate reflects the status of the manufacturing site at the time of the inspection, and Turkish Medicines and Medical Devices Agency should be consulted to verify compliance of the manufacturing site with GMP requirements if more than 3 years have elapsed since the date of inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field.

This certificate is valid only when presented with all pages and both Parts 1 and 2.

The authenticity of this certificate may be verified by Turkish Medicines and Medical Devices Agency upon request.

**This regulation is aligned with European Union Directive Directive 2003/94/EC laying down the principles and guidelines of good manufacturing practice for medicinal products for human use, and Directive 2001/83/EC on the Community code relating to medicinal products for human use.*

Prof. Dr. Ahmet AYAR
President of the Agency

Part 2

■ Human Medicinal Products *

1 MANUFACTURING OPERATIONS - MEDICINAL PRODUCTS*

If the company is engaged in manufacture of products with special requirements, e.g. radiopharmaceuticals or products containing penicillin, sulphanomides, cytotoxics, cephalosporins, substances with hormonal activity or other potentially hazardous active ingredients, this should be stated under the relevant product type and dosage form.

1.5 Packaging

- 1.5.2 Secondary packaging
 - 1.5.2.1 - QR coding operations
 - 1.5.2.2 - Box KT label etc. operations
 - 1.5.2.3 - Ink-jet etc. printing processes

Any restrictions or clarifying remarks related to the scope of this certificate *:

1.5.2: Also applicable for the storage of active substances/excipients.

2 IMPORTATION OF MEDICINAL PRODUCTS*

2.3 Other importation activities

- 2.3.1 Site of physical importation

Any restrictions or clarifying remarks related to the scope of this certificate *:

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26.03.2026

Prof. Dr. Ahmet AYAR
President of the Agency

