



REPUBLIC OF TURKEY
MINISTRY OF HEALTH
TURKISH MEDICINES AND
MEDICAL DEVICES AGENCY

TURKISH MINISTRY OF HEALTH
Turkish Medicines and Medical Devices Agency

Certificate No: TR/GMP/2021/281

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 : WORLD MEDICINE İLAÇ SAN. VE TİC. A.Ş.
Head Office / Correspondence Address : 15 Temmuz Mahallesi Cami Yolu Caddesi No:50 Güneşli
Bağcılar/İSTANBUL
Site Address : 15 Temmuz Mahallesi Cami Yolu Caddesi No:50 Güneşli
Bağcılar/İSTANBUL
Manufacturing Authorization Date : 17/11/2020
Manufacturing Authorization Number : TR/ÜY/2019/12-2

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 18-20/02/2019 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.*

16/12/2021

PhD Pharm. Sevil AZAK SUNGUR
Vice President of the Agency

TR/GMP/2021/281

Address: Söğütözü Mahallesi 2176. Sok. No:5 06520 Çankaya/ANKARA
Tel: (0312) 218 30 00 Fax: (0312) 218 34 60

Part 2

■ **Human Medicinal Products ***

1. MANUFACTURING OPERATIONS - MEDICINAL PRODUCTS*	
<i>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.</i>	
1.1	Sterile Products
	1.1.1 Aseptically prepared (processing operations for the following dosage forms) 1.1.1.4 Small volume liquids 1.1.3 Batch certification
1.2	Non-sterile products
	1.2.1 Non-sterile products (processing operations for the following dosage forms) 1.2.1.1 Capsules, hard shell 1.2.1.2 Capsules, soft shell 1.2.1.5 Liquids for external use 1.2.1.6 Liquids for internal use 1.2.1.8 Other solid dosage forms 1.2.1.11 Semi-solids 1.2.1.12 Suppositories 1.2.1.13 Tablets 1.2.1.15 Other non-sterile medicinal products (...free text)
	1.2.2 Batch certification
1.5	Packaging
	1.5.1 Primary Packaging 1.5.1.1 Capsules, hard shell 1.5.1.2 Capsules, soft shell 1.5.1.5 Liquids for external use 1.5.1.6 Liquids for internal use 1.5.1.8 Other solid dosage forms 1.5.1.11 Semi-solids 1.5.1.12 Suppositories 1.5.1.13 Tablets
	1.5.2 Secondary packaging
1.6	Quality control testing
	1.6.1 Microbiological (sterility)
	1.6.2 Microbiological (non-sterility)
	1.6.3 Chemical/Physical
	1.6.4 Biological testing

16/12/2021



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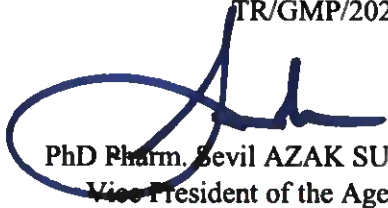
Any restrictions or clarifying remarks related to the scope of this certificate *:

- 1.1.1.4: Valid for “Eye drops, solution”, “eye drops,suspension”
- 1.2.1.1: Valid for “Capsule, hard”
- 1.2.1.2: Valid for “Capsule, soft”
- 1.2.1.5: Valid for “Nasal drops, solution”, “Nasal spray, solution”, “ Ear drops, solution”, “Cutaneous spray, solution”
- 1.2.1.6: Valid for “Oral solution”, “Oral drops, solution”, “ Oral suspension”, “Oral spray”, “Syrup”
- 1.2.1.8: Valid for “Granules”, “Pastille”, “Inhalation powder”, “Oral powder”, “Micropellet capsule”
- 1.2.1.11: Valid for “ Cream”, “Gel”, “Pomade”
- 1.2.1.12: Valid for “ Suppository”
- 1.2.1.13: Valid for “Tablet”, “Film coated tablet”, “Coated tablet”
- 1.2.1.15: Valid for “ Metered dose inhaler”, “Granules in sachet”

This certificate is valid until 31/12/2022.

16/12/2021

TR/GMP/2021/281



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