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DRUG

NATIONAL **AUTHORITY**

CERTIFICATE OF COMPLIANCE WITH GOOD MANUFACTURING PRACTICE GUIDELINES

THE NATIONAL DRUG POLICY AND AUTHORITY ACT, CAP 206

Issued under Regulation 19(5) of the National Drug Policy and Authority (Licensing) Regulations, 2014

Certificate No. 104/GMP/2023

This is to certify that the drug manufacturing facility:

Name of facility: World Medicine İlaç Sanayi Ve Ticaret Anonim Şirketi

Physical address of facility: Çerkezköy Organize Sanayi Bölgesi, Gazi Osmanpasa Mahallesi 6. Cadde, No: 30, Çerkezköy, Tekirdağ, Turkiye

License number of the manufacturer: TR/UY/2020/29-9

Has been inspected by the Authority for compliance with the Good Manufacturing Practice Guidelines.

On the basis of the inspection carried out on **10th, 11th, 12th, 13th and 14th April 2023,** it is certified that the facility indicated on this certificate complies with Good Manufacturing Practice for dosage forms listed in Table 1 below.

Table 1: Approved lines

No.	Dosage form	Category	Activities
1.	Tablets (Film coated & Uncoated)	Non-Beta Lactam	Manufacture of Finished Pharmaceutical (medicinal) Product
2.	Hard Gelatin Capsules		
3.	Tablet Lozenges		
4.	Liquids for internal use		
5.	Liquids for external use (Nasal sprays)		
6.	Suppositories		
7.	Powders in sachets		
8.	Small Volume Liquids in glass vials (Aseptically prepared)		
9.	Lyophilizates		
10.	Small Volume Liquids in ampoules (Aseptically prepared)		
11.	Small Volume Liquids in ampoules (Terminally sterilized)		
12.	Large Volume Liquids (Aseptically filled)		
13.	Large Volume Liquids (Terminally sterilized)		
14.	Sterile Eye/Ear drops (Aseptically prepared)		
15.	Small Volume Liquids in plastic ampoules (Aseptically prepared)		
16.	Semi-solids (Creams, Ointments & Gels)	Corticosteroids & Non-corticosteroids	
17.	Powders for oral suspension	Non-Beta Lactam	Manufacture of bulk product

The responsibility for the quality of the individual batches of the drugs manufactured through this process lies with the manufacturer.

This certificate remains valid until **14th April 2026.** It becomes invalid if the activities or the categories certified change or if the facility is no longer considered to be in compliance with GMP.

Issue Date: 10th October 2023.

Denis Mwesigwa
FOR THE AUTHORITY

NATIONAL DRUG AUTHORITY
10 OCT 2023
PLOT 19 LUMUMBA AVENUE
P. O. BOX 23096, KAMPALA

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