

Rwanda Food and Drugs Authority

Nyarutarama Plaza, KG 9 Avenue P.O. Box: 1948 Kigali - Rwanda Email: info@rwandafda.gov.rw website: www.rwandafda.gov.rw

QMS Nº: FDISM/FDIC/FMT/001

Revision No: 1

Effective Date: 11/10/2022

CERTIFICATE OF COMPLIANCE WITH GOOD MANUFACTURING PRACTICE

Certificate No: 3705

Issue Date: .05./09/2023

Valid up to: 0.4./09/2026

This is to certify that the pharmaceutical manufacturing facility with following details:

Name of facility: World Medicine Ilac Sanayi ve Ticaret. A.S.

Physical address: Gazi Osman Osb Mahallesi, 6 Cadde No.30 Street Organized Industrial Zone,

Cerkezkoy, 59500

License number: TR/UY/2020/29-9

Country: Turkey

E-mail: alkanat.deniz@worldmedicine.com.tr / mehlika.isk@worldmedicine.com.tr

Telephone: +0 (0282) 758 22 27

Has been assessed by the Rwanda Food and Drugs Authority for compliance with the Good Manufacturing Practice Guidelines. Based on the recognition pathway carried out on 07/08/2023, and 08/08/2023 it certifies that the pharmaceutical manufacturing facility indicated on this certificate complies with Good Manufacturing Practice for dosage forms, categories and activities listed in Table below:

No	Dosage form	Category	Activities
1.	Large Volume liquids, Lyophilizes and	Sterile	Manufacture of Finished
	Small volume liquids	products: Non	Pharmaceutical products: sterile
		beta lactams	and non-sterile products;
2	Capsules-hard shells, tablets (Film coated)	Non sterile	Packaging: Primary packaging,
	Other solid dosage forms: prolonged-	products: non	and secondary package
	release capsule, hard, chewable tablets,	beta lactams	Quality control testing:
	modified-release tablet, gastro-resistant		Microbiological: sterility,
EVA	tablet, film-coated tablets, lozenge/pastille		Microbiological: non-sterility,
	Other: Powder for oral solution in sachet,		Chemical/Physical
	granules in sachet		

The responsibility for the quality of the individual batches of pharmaceutical products manufactured through this process lies with the manufacturer. This certificate becomes invalid if the activities or the categories certified change or if the facility is no longer rated to be in compliance with Good Manufacturing Practice.

Dr. Emile BIENVENU Director General