

ISO 13485:2016

Certificate of Registration

This is to Certify that
Medical Device Quality Management System of

AZ PHARMA LABORATOIRES

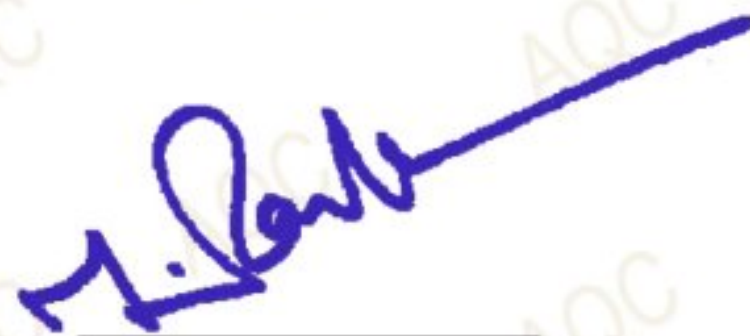
N°44, RDC 1ER ET 2EME ETAGE, ZONE INDUSTRIELLE AIN CHKAF, 30000, FES,
MOROCCO

has been assessed and found to conform to the requirements of
ISO 13485:2016
for the following scope :

MANUFACTURE, STORAGE AND DISTRIBUTION OF CLASS I NON-STERILE,
NON-INJECTABLE AND NON- IMPLANTABLE MEDICAL DEVICES (LAXATIVES
AND RECTAL ENEMAS, ORAL AND DENTAL SOLUTIONS, NASAL SOLUTIONS).

FABRICATION, STOCKAGE ET DISTRIBUTION DES DISPOSITIFS MÉDICAUX NON STÉRILES,
NON INJECTABLES ET NON IMPLANTABLES DE CLASSE I (PRODUITS LAXATIFS ET
LAVEMENT RECTAL, SOLUTIONS BUCCO DENTAIRE, SOLUTIONS NASALES)

Certificate No	: 25EMOV58		
Initial Registration Date	: 26/03/2025	Issuance Date	: 26/03/2025
Date of Expiry	: 25/03/2028		
1st Surv. Due	: 26/02/2026	2nd Surv. Due	: 26/02/2027



Director



(Scan to Verify)

Certificate Verification: The Certification Validity can be checked at www.aqcworld.com at Clients Directory.

Certificate is the property of Assurance Quality Certification LLC (AQC) located at Sharjah Media City, SHAMS, Sharjah, UAE.

Email: info@aqcworld.com and shall be returned immediately when demanded.

*Validity of the Certificate is subject to successful completion of surveillance audit on or before of due date.