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

EGAC Accredited
MDQMS Certification
CAB # 012226

MEMBER OF MULTILATERAL RECOGNITION ARRANGEMENT IAF

(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.*

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