

RAHMA AZIZAH

082118356215 | raazizah0811@gmail.com | linkedin.com/in/rahma-azizah-131a77216

Pharmacist with hands-on exposure to pharmaceutical manufacturing, sterile environment monitoring, and Quality Assurance through industrial pharmacist internship. Experienced in GMP/CPOB compliance, SOP monitoring, batch documentation review, manufacturing process monitoring, IPC, quality documentation, and audit readiness. Familiar with deviation investigation, Root Cause Analysis, CAPA, QRM, FMEA, Change Control, qualification, and validation. Experienced in pharmaceutical laboratory testing and instrumental analysis with strong attention to detail and data traceability.

Professional Experience

PT. Enseval Putera Megatrading Tbk. Apr 2026

- Pharmacist Intern*
- Verified completeness, accuracy, and integrity of 40+ distribution documents weekly, including invoices, delivery orders, shipping lists, and product return records, with >98% accuracy.
 - Reviewed 30+ receiving and distribution records for CDOB/GDP compliance, product traceability, batch/expiry information, and documentation completeness.
 - Reconciled product and documentation data across 20+ transactions, identifying and reporting discrepancies before distribution processing.
 - Supported product return documentation and traceability checks for 10+ returned items in accordance with applicable procedures.

Lembaga Farmasi Angkatan Udara Feb 2026 - Mar 2026

- Pharmacist Intern*
- Verified GMP/CPOB and SOP compliance across pharmaceutical manufacturing activities, ensuring process adherence and documentation integrity.
 - Performed In-Process Control (IPC) and process compliance checks during manufacturing; monitored operations in controlled manufacturing environments, documented observations, and escalated discrepancies to supervisors.
 - Reviewed 20+ Batch Production Records (BPR) and supporting documents for completeness, accuracy, data integrity, and traceability before quality evaluation.
 - Supported deviation investigation and Root Cause Analysis (RCA) for 3+ quality issues; contributed to CAPA recommendations, Quality Risk Management (QRM), and FMEA assessments.
 - Inspected facility and critical equipment parameters, including HVAC, HEPA H13 filtration, and pressure cascade, and documented results according to applicable procedures.
 - Supported Change Control, qualification, validation, and audit readiness activities by compiling evidence and verifying required documentation.
 - Conducted sampling and laboratory testing of pharmaceutical samples with complete and traceable documentation.

Bandung Islamic University Feb 2025 - Aug 2025

- Research & Laboratory Assistant*
- Conducted 50+ physicochemical tests and instrumental analyses under SOP/GLP, maintaining accurate and traceable documentation.
 - Performed 30+ laboratory activities including equipment preparation, calibration checks, and performance verification.
 - Analyzed and documented 100+ test results with accurate calculations, data traceability, and systematic record verification.

Organizational

Himpunan Mahasiswa Farmasi Jun 2022 - Aug 2024

- Coordinated administrative activities across 12 divisions and managed documentation for 15+ organizational programs
- Standardized record-keeping workflows, ensuring timely documentation, proper filing, and cross-departmental communication
- Prepared meeting minutes, official correspondence, and activity reports; improved document traceability and retrieval efficiency

Education

Universitas Islam Bandung Sep 2025 - Aug 2026

- Professional Pharmacist Program, 3.99/4.00*
- Practical exposure to pharmaceutical manufacturing, GMP/CPOB, quality systems, IPC, batch documentation, deviation, CAPA, quality risk management, validation, and qualification.

Universitas Islam Bandung Aug 2021 - Feb 2025

Undergraduate Pharmacy, 3.75/4.00

Core Competencies

- **Quality Assurance & GMP:** GMP/CPOB | SOP Compliance | In-Process Control (IPC) | Batch Record Review | Batch Documentation | Manufacturing Process Monitoring | Deviation | Root Cause Analysis (RCA) | CAPA | QRM | FMEA | Change Control | Validation & Qualification | Audit Readiness | Sampling | Product Traceability
- **Facility & Manufacturing:** HVAC | HEPA H13 Filtration | Pressure Cascade | Controlled Manufacturing Environment | Facility & Equipment Inspection
- **Laboratory & Analytical:** Physicochemical Testing | Instrumental Analysis | HPLC | UV-Vis | Dissolution | Stability Testing | pH | Viscosity | Particle Size | GLP | Calibration Checks | Sampling
- **Software:** Microsoft Excel | Microsoft Word | Microsoft PowerPoint
- **Languages:** Bahasa Indonesia — Native | English — Intermediate / Professional Working Proficiency

MAKIN AHLI



CERTIFICATE

479/AF-0F8/MA/VIII-2026

This Certificate is Proudly Presented to

Rahma Azizah

Has successfully completed training

**Quality Assurance (QA)
in Pharmaceutical Industry**

On August 29, 2026



Ariyani Wastika Mukti, S. Si

Head of Learning and Development

apt. Herdyna Gita Violleni, S.Farm.

Trainer

www.makinahli.com