



M. YERICHO

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PROFESSIONAL SUMMARY

A licensed pharmacist and graduate of the Professional Pharmacy Program with a strong interest in the pharmaceutical industry. Experienced as a Technical Person in Charge (PIC), with a solid understanding of Good Manufacturing Practices (GMP), Indonesian Good Manufacturing Practices (CPOB), and pharmaceutical regulatory compliance. A highly adaptable, results-oriented professional with strong analytical and problem-solving skills, capable of working effectively both independently and as part of a team. Committed to contributing to product quality assurance and ensuring compliance with applicable regulatory standards.

WORK EXPERIENCE

Technical Person in Charge, PT Arfia Megah Cikarang

July 2025- May 2026

- Prepared product registration dossiers and managed the registration process for veterinary pharmaceutical products with the Indonesian Ministry of Agriculture.
- Conducted training programs for all personnel involved to ensure regulatory compliance and operational readiness.
- Supervised daily production activities to ensure smooth operations and on-time completion according to production schedules.
- Oversaw quality control of in-process and finished products to ensure compliance with quality standards.
- Managed and monitored raw material and packaging material inventory to support uninterrupted production.
- Ensured the implementation and compliance of Good Manufacturing Practices for Veterinary Medicinal Products (CPOHB/GMP).
- Served as the primary liaison between the company and government authorities on regulatory and compliance matters.

Beta-Lactam Production Pharmacist (Intern) PT Sanbe Farma Unit 2, Cimahi Aug 2024 – Sep 2024

- Gained a solid understanding of Good Manufacturing Practices (GMP/CPOB) in pharmaceutical manufacturing.
 - Monitored production processes to ensure compliance with quality standards and regulatory requirements.
 - Developed an understanding of pharmaceutical documentation systems and controlled documentation practices.
 - Acquired knowledge of quality control testing for raw materials, in-process materials, and finished products to ensure compliance with product specifications.
 - Developed an understanding of the Indonesian National Agency of Drug and Food Control (BPOM) regulations related to pharmaceutical product registration.
 - Gained knowledge of Supply Chain Management processes, from marketing order planning through product distribution to end customers.
 - Performed trend analysis of rubber stopper and aluminum cap consumption data using process capability indices (Cp and Cpk) to support process performance evaluation.
 - Conducted a shelf-life study for pharmaceutical vials to support product stability assessment.
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EDUCATION

Pharmacist Professional Program	Mar 2024 - Apr 2025
Universitas Islam Sultan Agung Semarang	
• Graduated as a licensed pharmacist with a GPA of 3.50/4.00.	
Bachelor of Pharmacy	Sep 2019 - Sep 2023
Univesitas Sari Mulia Banjarmasin	
• Graduated with a Bachelor of Pharmacy with a GPA of 3.56/4.00..	

SKILLS

- **Good Manufacturing Practices (GMP/CPOB/CPOHB):** Strong understanding of the principles and implementation of Good Manufacturing Practices in compliance with BPOM regulations.
- **Pharmaceutical Manufacturing Processes:** Knowledge of pharmaceutical manufacturing processes for solid, liquid, semi-solid, sterile, and non-sterile dosage forms.
- **Production Documentation:** Familiar with batch records, logbooks, Standard Operating Procedures (SOPs), and Good Documentation Practices (GDP).
- **Teamwork & Effective Communication:** Able to collaborate effectively in cross-functional teams and communicate information clearly and professionally.
- **Attention to Detail & Procedural Compliance:** Detail-oriented with a strong commitment to following SOPs, maintaining accuracy, and meeting deadlines.
- **Problem-Solving:** Capable of identifying process-related issues, analyzing root causes, and supporting the implementation of practical solutions.

CERTIFICATIONS

Understanding and Implementation of CPOB, CPOTB, and CPOBAB in the Pharmaceutical Industry	May 2026
JadiPraktisi.com	
Completed an intensive training program covering Good Manufacturing Practices (CPOB), including the 12 Key Aspects of CPOB, CPOB 2025 updates, strategies for effective CPOB implementation, and Quality Management System (QMS) integration in pharmaceutical manufacturing.	
Veterinary Drug Registration Procedures Training	Aug 2025
Asosiasi Obat Hewan Indonesia	
Completed intensive training on veterinary drug registration procedures, including the registration process, submission workflow, and preparation of regulatory documentation in accordance with the standards of the Indonesian Center for Veterinary Drug Testing (BBPMSOH) and the Ministry of Agriculture.	
Good Manufacturing Practices (GMP/CPOB) Training	Mar 2025
www.makinahli.com	
Completed intensive training on the principles and implementation of Good Manufacturing Practices (GMP/CPOB) in the pharmaceutical industry, covering Quality Management Systems (QMS), documentation practices, process validation, production control, and preparation for regulatory audits and inspections in accordance with BPOM regulations and international GMP standards.	
Quality Assurance (QA) In Pharmaceutical Industry	Mar 2025
www.makinahli.com	
Completed training on the roles and responsibilities of Quality Assurance (QA) in the pharmaceutical industry, covering Quality Management Systems (QMS), documentation practices, change control, deviation investigations, Corrective and Preventive Actions (CAPA), and preparation for internal and external audits in compliance with Good Manufacturing Practices (GMP/CPOB).	