



[Okky Himawan]

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PROFILE

Cum Laude graduate of the Pharmacist Professional Program at the University of Indonesia with a strong interest in Research & Development (R&D), particularly in formulation development within the pharmaceutical industry. Possesses internship experience in the Quality Control (QC) department of a pharmaceutical company, with hands-on involvement in the development and standardization of quality documentation, stability studies, and analytical activities in accordance with ICH guidelines. Familiar with formulation-related literature studies, stability protocols, and pharmaceutical quality systems, including Good Manufacturing Practice (GMP/CPOB), Good Laboratory Practice (GLP), and Good Documentation Practice (GDP). Supported by academic experience in pharmaceutical formulation and an undergraduate research project entitled “Development of Water Extract of *Costus woodsonii* Maas. Bract as an Antioxidant Shampoo,” which strengthened understanding of dosage form development, product evaluation, and formulation optimization. Detail-oriented, analytical, adaptable, and capable of working effectively in dynamic laboratory and industrial environments with a strong commitment to product quality and continuous improvement.

INTERNSHIP EXPERIENCE (PKPA)

University of Indonesia Hospital (RSUI)

Sep 2025 – Oct 2025

- Supported clinical pharmacy services by dispensing medications accurately based on prescriptions.
- Participated in stock opname activities to ensure inventory accuracy.
- Prepared non-sterile pharmaceutical preparations (ointments and capsules) in accordance with procedures.

PT Sterling Products Indonesia (Haleon Indonesia) – QC Intern

Jul 2025 – Aug 2025

- Developed and streamlined laboratory report templates for product release and stability testing, improving documentation efficiency and supporting GDP implementation.
- Designed and standardized templates for stability assessment and change control to enhance consistency within the Quality Management System (QMS).
- Prepared stability protocol templates in accordance with ICH Guidelines and regulatory requirements.
- Assisted in the development of Standard Operating Procedures (SOP) for attendance systems with structured documentation.

- Compiled halal material documentation in compliance with KMA No. 748/2021 to support regulatory readiness.
- Supported regulatory audit processes by BPOM, ensuring compliance with GMP/CPOB and applicable regulations.

Kimia Farma Pharmacy (Matraman)

Jun 2025 – Jul 2025

- Supported pharmacy services by ensuring accurate dispensing based on prescriptions.
- Provided patient counseling and guidance for appropriate self-medication.
- Participated in Prolanis (Chronic Disease Management Program), including patient education and monitoring.

PT Tri Sapta Jaya

May 2025 – Jun 2025

- Managed and aligned product master data with regulatory databases to ensure compliance.
- Compiled and analyzed distribution (genba) data to support operational monitoring at national level.
- Identified and documented expired marketing authorization (NIE) products as part of regulatory compliance control.

Puskesmas Makasar

Apr 2025 – May 2025

- Supported pharmaceutical services by ensuring accurate dispensing and labeling of medications.
- Provided patient education on HIV therapy, including medication use and adherence.

EDUCATION

Universitas Indonesia (2025-2026)

Pharmacist Professional Program – GPA 3.67

Universitas Pembangunan Nasional “Veteran” Jakarta (2020-2024)

Bachelor of Pharmacy - GPA 3.51

ORGANIZATIONAL EXPERIENCE

Faculty of Medicine Pharmacy Student Association UPN “Veteran” Jakarta (2023)

Head of Student Welfare and Advocacy Division

- Led and coordinated divisional programs effectively.
- Acted as a liaison between students and faculty stakeholders for academic and non-academic matters.

LABORATORY EXPERIENCE

Laboratory Teaching Assistant (Phytochemistry, Pharmaceutical Chemistry, Pharmaceutics) (Feb 2022 - Jun 2023)

- Assisted in laboratory demonstrations and practical sessions.
- Managed laboratory materials and equipment administration.

- Collaborated with laboratory staff to ensure timely preparation of materials.
- Applied Good Laboratory Practice (GLP) in all laboratory activities

SKILLS

- Good Manufacturing Practice (GMP) / CPOB
- Good Documentation Practice (GDP)
- Quality Management System (QMS)
- Stability Study & Stability Protocol
- Change Control & Documentation
- Microsoft Office (Excel, Word, PowerPoint)
- Laboratory Documentation & Administration
- English (Professional Working Proficiency)