



MEYLIANA ADHITYA KUSUMAWARDHANI

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Solo, Central Java

SUMMARY

A Pharmacist graduates from Sebelas Maret University with 6 months of industrial, clinical, and distribution internship experience and 1.5 years in analytical development. Have a high interest in a career in the health and pharmaceutical industry, especially analytical development.

EDUCATION LEVEL

Universitas Sebelas Maret - Surakarta

Aug 2019 - Sept 2023

Bachelor Degree in Pharmacy, 3.62 / 4.00

- Get funding in Program Ilmiah Mahasiswa (PKM) from Kemenristekdikti 2021.
- Get funding in Program Kompetisi Kampus Merdeka (PKKM) from University 2021 and 2022.
- Thesis title: Optimatization of Condition High Performance Liquid Chromatography in Resveratrol Analysis Based on Central Composite Design

Universitas Sebelas Maret - Surakarta

Jan 2024 - April 2025

Pharmacist, 3.84 / 4.00

ACHIEVEMENT

Achievements (2022): Author of scientific articles in the Bulletin of Pharmaceutical Sciences. Assiut.

PROFESSIONAL EXPERIENCES

PT Naturafit Tribunnabawi Internasional - Sragen, Central Java

April 2024

- Learned activities of an apothecary in small business of herbal medicine
- Observed and analyzed the implementation of CPOTB (Cara Pembuatan Obat Tradisional yang Baik) in small business of herbal medicine
- Learned the process of making herbal medicine from the initial ingredients until released product

Universitas Sebelas Maret Hospital - Solo, Central Java

May 2024 - Jun 2024

- Assisted in the preparation of pharmaceuticals, medical devices and BMHP management
- Assisted with administrative activities such as monitoring reports on drug stock storage
- Performed Reconciliation admissions, transfers and discharge
- Verified prescriptions and drugs to be given to patients
- Provided counseling regarding the use of special drugs to patients
- Provided drug information, conduct visits, and educate patients before going home
- Conducted case studies monitoring drug therapy in patients with renal failure
- Learned about drug side effect monitoring (MESO) and Pharmacovigilance

PT Zen Semitra Raya - Solo, Central Java

Jul 2024

- Implemented distribution aspects in pharmacy operations, Good Distribution Practice (GDP), Supply chain management, warehouse management

Ngoresan Health Center - Solo, Central Java

Aug 2024

- Learned some activities of apothecary in public health center such as counseling drug information, drug procurement, and compile LPLPO (Laporan Pemakaian dan Lembar Permintaan Obat)
- Conducted and assisted pharmaceutical and clinical services such as screening prescription, dispensing drug, drug information services, counseling and communicating with patients.

Kimia Farma Pharmacy - Boyolali, Central Java

Sept 2024

- Performed pharmaceutical services such as screening and prescription services, drug dispensing and compounding, drug information services, self-medication, and patient counseling
- Recapitulation of daily use of narcotics, psychotropics and certain drugs (OOT) to be reported to SIPNAP every month
- Stored medicines grouped based on pharmacology, alphabetically, storage temperature, First in first out (FIFO), and First expired first out (FEFO). Over-the-counter (OTC) medicines are placed on the gondola by category.

PT Finusolprima Farma Internasional (a Kalbe Company) - Bekasi, West Java

Sept 2024 - Nov 2024

Quality Assurance (Computer System Validation)

- Learned how to make medicines properly and correctly according to the CPOB (Good Ways to Make Medicines) aspects in the pharmaceutical industry.
- Learned several activities and roles of a pharmacist who must fill the pharmaceutical industry in three positions, namely Quality Assurance (QA), Quality Control (QC), and Production
- Prepared product quality reports, stability reports and performance qualifications
- Analyze and prepare Quality Risk Management for equipment and machine
- Prepared Preventive Maintenance (PM), Corrective Maintenance (CM) and Breakdown Maintenance (BM) reports in machine usage reports
- Analyze and prepare Computer Validation IQ and OQ for data Laboratory and e-registration
- Performed spreadsheet validation for QC product and equipment calibration
- Analyze and prepare Product Quality Review (PQR) and Contamination Control Strategy Document

Research Assistant

Aug 2022 - Des 2022

- Title: Bioavailability Assessment of Mesoporous Incorporated Resveratrol Nanocrystal-Based Preparations Materials and Polyherbal Nanoemulsion as Scientific Evidence of Drug Delivery System
- Conducted biological sample analysis using hplc with several methods optimized with design expert software to produce the most optimal system suitability test

Faculty of Mathematics and Science Laboratory - Surakarta, Central Java

Aug 2022 - Des 2023

Operator HPLC

- Performed analyze several samples using HPLC in the Laboratorium MIPA terpadu

WORK EXPERIENCES

PT Erlangga Edi Laboratories - Semarang, Central Java

Oct 2025 - June 2026

Supervisor QC Raw Material

- Supervise and ensure proper testing and inspection of all incoming raw materials in accordance with approved specifications, analytical methods, and standard operating procedures
- Ensure full compliance with Good Manufacturing Practices (GMP) and internal company quality standards throughout all QC activities
- Review, verify, and approve analytical test results as well as Certificates of Analysis (CoA) to ensure accuracy, completeness, and regulatory compliance
- Plan, organize, and manage sampling activities, testing schedules, and overall laboratory operations to ensure timely and efficient workflow
- Lead investigations of out-of-specification (OOS) results and deviations, identify root causes, and implement appropriate corrective and preventive actions (CAPA)
- Coordinate effectively with warehouse, production, and PPIC departments to ensure proper status, storage, and timely release or rejection of raw materials
- Train, guide, and supervise QC staff to enhance technical competency, ensure adherence to procedures, and maintain high performance standards
- Maintain comprehensive documentation and ensure data integrity in compliance with regulatory requirements, including proper recording, reporting, and archiving of QC data