



Berliana Rizka Ramadhan

Apothecary

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Summary

Industrial Pharmacist with extensive experience in pharmaceutical and cosmetic manufacturing environments. Skilled in ensuring GMP compliance, batch documentation review, and production process monitoring to maintain product quality standards. Experienced in handling production deviations, preparing investigation reports, and implementing corrective and preventive actions (CAPA). Strong understanding of manufacturing processes combined with quality assurance principles. A detail-oriented professional with strong analytical and problem-solving skills, capable of working collaboratively with cross-functional teams to support continuous improvement and regulatory compliance.

Education

University of Indonesia

(Feb 2021 - Mar 2022)

Bachelor Apothecary Program

GPA: 3.67/4.00

Sebelas Maret University

(Aug 2016 - Jul 2020)

Bachelors Pharmacy

GPA: 3.71/4.00

Assistant Lecturer

- Assist in provided direction when practicing in the laboratory and teach how to operate instrument analyze HPLC, Spectrofotometry, and AAS.
- Evaluated the understanding of each student in every activity that has been carried out for this course; Solid Dosage Forms Formulation and Technology (2020), Physical Pharmacy (2020), Chemistry Analysis (2019), and Steril dosage forms Formulation and Technology (2019).

Work Experience

PT. Sinar Alfa Omega - Periuk, Tangerang

(18 Jan 2026- Jul 2026)

Supervisor Production

- Ensured production processes complied with GMP and company quality standards.
- Reviewed and verified Batch Manufacturing Records (BMR) to ensure completeness and accuracy before submission to QA.
- Investigated production deviations and coordinated with the Quality team to implement Corrective and Preventive Actions (CAPA).
- Ensured all production activities followed approved Standard Operating Procedures (SOPs) and documentation practices.
- Supported deviation investigation, change control, and quality documentation review.
- Conducted operator briefings and training related to GMP compliance and documentation accuracy.
- Participated in internal audits and implemented corrective actions related to production findings.
- Ensured proper line clearance and adherence to hygiene and sanitation standards in the production area.
- Developed and reviewed Batch Manufacturing Records (BMR) for production scale to ensure accuracy and compliance with GMP requirements.

PT. Gondowangi Tradisional Kosmetika - Cikarang, Bekasi

(22 April 2024 - Jan 2026)

Supervisor Production

- Manage and lead a team of production leaders and operators, providing guidance and training.
- Oversee daily manufacturing operations, ensuring compliance with GMP and other regulatory standards.
- Collaborate with cross-functional teams, including R&D, Quality Assurance, and Supply Chain, to streamline production processes.
- Develop and implement production strategies to optimize efficiency and productivity.
- Ensure adherence to quality control measures and safety protocols.
- Investigate and resolve production issues while ensuring minimal downtime.
- Maintaining accurate records of production activities, including raw material usage, production processes, and quality control data
- Maintain and update manufacturing records, sign off completed batch records before handover to Quality Department for release.
- Write and review incident reports, change requests corrective action reports and risk assessments.

Foreman Packaging and Liquid Process

Responsibilities:

- Supervised 80+ employees for herbal process and packaging lines.
 - Participating in the production planning process and ensuring coordination on materials and arrangement of appropriate resources to support the production plan.
 - Reviewing and approving issued and executed batch records and ensure cGMP is maintained in production operations.
 - Performing process investigations, documenting process deviations, initiating Capas and minor equipment change controls.
 - Analyzed and published daily production OEE output results.
 - Made daily production schedules using SAP system when product was released to manufacturing.
 - Ensure that production equipment remains in an appropriate state of maintenance, calibration & validation.
- Created protocols, execution of qualifications and make summary reports for the production equipment.

Mahakam Beta Farma- Jatinegara, Jakarta Timur**(1 July 2021 - 31 August 2021)**

Student Intern

Responsibilities:

- Participated in Performance Qualification System Heating, Ventilation, and Air Conditioning in Sterile Area Production at Non Cephalosporin Site PT Mahakam Beta Farma period 2021.
- Participated in Online Training "Basic Quality Control".
- Helped to write reports about cleaning validation using Microsoft Word and Excel.
- Produced Handsanitizer for internal purposes at Mahakam Beta Farma.

IKOP Sdn Bhd- Kuantan District, Pahang, Malaysia**(20 Jan-31 Jan 2020)**

Student Intern

Responsibilities:

- Learned about Industrial Pharmacy Product with shariah requirement and halal manufacture works with all departement such as QA, QC and Production in office.
- Participated in Quality Control for nished product iGesic Ointment and do In-Process Control for tablet Product.
- Controlled and organize raw material for production in warehouse

Course & Training

• Pelatihan ISO 9001	2025
• Pelatihan CPKB- PT. Gondowangi Tradisional Kosmetika	2024
• Pelatihan CPOB/CPOTB- PT. Supra Ferbindo Farma	2024
• Kenal Profesi PPIC Supervisor- Latih	2022
• Management Skills for Warehouse Supervisor- Latih	2022
• Basic Quality Control- PT. Mahakam Beta Farma	2021
• Kupas Tuntas Proses Produksi di Industri Farmasi- Pharmatech Talkshow	2019

Publication

ERMAWATI, D. E., YUGATAMA, A., RAMADHAN, B. R., PERTIWI, I., ROSIKHOH, A., & NOVACHIRIA, S. R. (2022). STABILITY AND ANTIBACTERIAL ACTIVITY TEST OF NANOSILVER BIOSYNTHETIC HYDROGEL. International Journal of Applied Pharmaceutics, 14(2), 221–226.

Softskill

- Leadership
- Teamwork
- Time Management
- Communication
- Problem Solving

Hardskill

- MS Office
- SAP
- IBM SPSS
- Calibration and Qualification
- Process Validation
- Pharmaceutical Research

Portofolio**Journal: International Journal of Applied Pharmaceutics**<https://bit.ly/3Mm4FYv>**Portofolio & Certificates**<https://bit.ly/3xM8WRj>**References**

apt. Diana Luky, S.Farm.

Foreman Production- PT. Supra Ferbindo Farma (The Tempo Scan Group)<https://www.linkedin.com/in/diana-luky-mirnasari-23a8b8208/>