

APT. RIMAWATI S HUSNA, S. FARM

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SUMMARY

Licensed Pharmacist with supervisory experience in pharmaceutical Quality Control covering raw materials and finished goods. Experienced in BPOM audit preparation, vendor qualification audit, and KPI-based performance monitoring. Strong background in GMP compliance, laboratory oversight, and quality system implementation. Demonstrated ability to lead QC operations and support continuous improvement initiatives.

WORK EXPERIENCE

PT. Sampharindo Retroviral Indonesia, Semarang - Central Java, Indonesia

Specializing in the production of antiretroviral (ARV) medicines for HIV treatment, sister company of Macleods Pharmaceuticals Ltd, (India)

Quality Control Supervisor - Raw & Packaging Material

Jan 2025 - Present

- Supervising raw material testing and release for APIs, excipients, and packaging materials (primary, secondary, and tertiary) to ensure compliance with GMP standards
- Reviewing analytical data and ensuring compliance with pharmacopeial specifications
- Interpreted analytical results and ensured accuracy, traceability, and adherence to ALCOA+ principles.
- Conducting external vendor qualification, supplier audits, and regulatory inspections.
- Participating in BPOM audit (2025) and supporting laboratory readiness & documentation review
- Arrange and implementing KPI monitoring system to improve QC performance and operational effectiveness
- Coordinating with QA, PPIC, and Warehouse for material disposition and investigation handling
- Manage sampling, laboratory investigations, CAPA, Change Control, and QRM.
- Develop and revise SOPs, specifications, analytical data sheets and quality documentation.

Quality Control Supervisor - In Process Control & Finish Goods

Jan 2024 - Dec 2024

- Supervised daily laboratory operations for Finished Goods (FG) and In-Process Control (IPC) testing, ensuring compliance with GMP requirements, SOPs, and testing schedules.
- Planned and coordinated weekly analytical workloads, sample priorities, and personnel assignments to ensure timely testing and product release.
- Supervised and evaluated the performance of QC analysts and IPC personnel while providing technical guidance and on-the-job training.
- Reviewed analytical data, Analytical Data Sheets (ADS), Certificates of Analysis (CoA), and laboratory documentation prior to approval.
- Ensured the completeness, accuracy, and compliance of laboratory records, including logbooks, Batch Manufacturing Records (BMR), and analytical documentation.
- Coordinated laboratory resources and overtime activities to maintain testing timelines and operational efficiency.
- Prepared, revised, and implemented Analytical Data Sheets (ADS), Standard Operating Procedures (SOPs), and other quality documents.
- Supported continuous improvement initiatives by maintaining laboratory compliance with internal quality standards and GMP requirements.

EDUCATION

Profession Apothecary, Airlangga University | 2022-2023

GPA: 3.47 / 4.00

Bachelor of Pharmacy, Airlangga University | 2018-2022

GPA: 3.53 / 4.00

ADDITIONAL INFORMATION

- **Technical Skills:** Good Manufacturing Practice (CPOB/GMP), Quality Control Management, Raw Material & Packaging Material Testing, Finished Goods & In-Process Control (IPC), Laboratory Investigation (OOS/OOT), Risk Management (QRM), CAPA & Change Control, Vendor Qualification & Vendor Audit, SOP & Work Instruction Development, Root Cause Analysis (5 Why, Fishbone), Statistical Data Trending & KPI Monitoring
- **Software:** Microsoft Office (Excel, PowerPoint, Word, Outlook), Windows
- **Professional Registration:** Registered Pharmacist (Apothecary), STRA (Active)
- **Training & Certifications:** Internal GMP/CPOB Training, Data Integrity Training, CAPA & Deviation Training, Risk Management Training, HPLC & GC Training, Seminar HISFARIN