

RIFDA SATYA AZWANA

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Depok, West Java

Fresh graduate Pharmacist from University of Indonesia with hands on experience in batch documentation, quality monitoring product, and production workflows. Involved in pharmaceutical quality evaluation and laboratory testing of formulated products as a research assistant with exposure to quality assessment procedures and data documentation. Familiar with operating analytical instrument and conducting data processing for pharmaceutical analysis including HPLC and GC-MS. Proactive, highly motivated, and recognized for honesty, responsibility, and integrity, with exceptional attention to detail and discipline. Seeking opportunities to contribute and grow in pharmaceutical industry-related roles.

EDUCATION

University of Indonesia – Depok, West Java Jan 2025 – Apr 2026

Professional Pharmacist Program, GPA: 3.68/4.00

- Completed Pharmacist Professional Practice (PKPA) in pharmaceutical industry/hospital/pharmacy community
- Have an active STRA (Surat Tanda Registrasi Apoteker)
- Elective Course : Registration and Pharmaceutical Product Monitoring

Syiah Kuala University – Banda Aceh Jul 2020 – Jan 2024

Bachelor of Pharmacy, GPA: 3.41/4.00

- Participated in Merdeka Belajar Kampus Merdeka (MBKM) Research Program and was awarded a research grant for a study on microbial contamination in intravenous medications used for pediatric patients at a hospital in Aceh
- Served as a Pharmacology mentor and Analytical Chemistry Laboratory Assistant, assisting in teaching activities and laboratory practices

WORK EXPERIENCES

PT Kalbio Global Medika – West Java Jul 2025 – Aug 2025

Pharmacist Professional Internship

Pharmaceutical Company, Part of Kalbe Group

- Assisted in evaluating 30+ packaging personnel performance and operational practices to support process improvements and production efficiency
- Created a training module for newly assigned packaging personnel, covering GMP principles, production procedures, and quality requirements to support operational readiness and compliance with pharmaceutical manufacturing standards
- Identified, documented, and reported defective products during secondary packaging processes in accordance with quality standards.
- Observed Manual Visual Inspection (MVI) procedures for 3 types of sterile products following primary packaging according to standard operating procedures (SOP)
- Participating in department induction including Production Drug Substances and Production Product Workflows, Personal Hygiene, Handling and Control of Documents and Records, Halal Management System, Good Documentation Practice for cross-functional understanding

PT SamMarie Tramedifa – East Jakarta Jun 2025 – Jul 2025

Pharmacist Professional Internship

Pharmaceutical Distributor, Part of SamMarie Group

- Gained exposure to 30+ pharmaceutical supply chain operations, including inventory control, product traceability, documentation management, and distribution quality systems
- Assisted in verification, receipt and distribution of 100+ pharmaceutical products and vaccines by ensuring alignment between physical inventory, invoices, and purchase orders.
- Documented the end-to-end recall process, covering regulatory instructions, distribution monitoring, and product return to suppliers with complete records.
- Supported Regulatory and Quality Assurance activities by developing Cold Chain Product (CCP) handling materials and ensuring proper receiving and cold storage products.
- Prepared and verified electronic regulatory reports (e-reports) based on actual drug sales data, ensuring data accuracy and compliance with applicable regulatory requirements.
- Conducted analysis and evaluation of pharmaceutical distribution activities to assess compliance with Good Distribution Practice (GDP/CDOB) requirements and identify opportunities for process improvement.

- Managed pharmaceutical inventory through planning, procurement, receiving, storage (FIFO/FEFO).
- Analyzed purchasing patterns of 50+ loyal and royal patients to support strategies at improving pharmacy profitability
- Conducted inventory control activities including storage monitoring, stock opname to maintain stock accuracy and availability
- Assisted in ordering pharmaceutical products from distributors to ensure stock availability
- Maintained archiving, documentation, and reporting of prescriptions, narcotics, psychotropics, and pharmaceuticals services in compliance with regulatory requirements, including uploading narcotics and psychotropics reports to SIPNAP under supervision.

- Conducted microbiological contamination testing of three types of intravenous preparations compounded under sterile conditions using Laminar Air Flow (LAF) and open-room environments under sterile compounding standards in open rooms
- Evaluated contamination rates across different preparation methods using Total Plate Count (TPC) method, including comparison between LAF and open-area compounding practices
- Identified and analyzed types of contaminating microorganisms present in intravenous preparations.
- Assessed potential procedural errors during aseptic compounding processes using LAF to identify factors contributing to microbial contamination and support quality improvement in sterile preparation workflows.
- Prepared laboratory documentation and research reports in compliance with Good Laboratory Practice (GLP)

- Trained and assisted 20+ students in operating HPLC, GC-MS, pH Meters and other related instruments.
- Guided users in reading, interpreting, and evaluating analytical results.
- Supervised and guided 20+ students in performing qualitative and quantitative chemical analysis to identify specific compounds in laboratory samples.

ORGANIZATIONAL EXPERIENCE

- Initiated and established three study clubs (Pharmacology, Pharmacokinetics, and Pharmaceutics) as platforms for the academic development of pharmacy students at Syiah Kuala University
- Organized training programs, scientific seminars, and scientific writing competitions for students

- Established partnerships with external parties outside FMIPA USK, including Aceh media outlets and government institutions, particularly in Banda Aceh and surrounding areas

SKILLS

Technical Skills : HPLC, GC-MS, pH Meter, Microbial testing, Laminar Air Flow Operation, Good Manufacturing Practice (GMP), Good Laboratory Practice (GLP), Canva, Microsoft Office, SPSS Statistic

Personal Skills : Detail-oriented, Communicative, Strong teamwork, Collaboration skills, Adaptable, Eager to learn new things