

apt. Renanda Natanael, S.Farm.

Registered Pharmacist | QA Validation Support | Process Validation | Validation
Documentation | GMP/CPOB | Quality Management System (QMS)

Bekasi, West Java, Indonesia | (+62) 822 6959 9340

renanda.natanael@gmail.com | <https://www.linkedin.com/in/renanda-n-2b547a140/>

PROFESSIONAL SUMMARY

Registered Pharmacist with internship experience in Quality Assurance and Quality Control within GMP/CPOB-regulated pharmaceutical manufacturing. Possesses practical knowledge of laboratory analytical testing, quality documentation, Good Manufacturing Practices (GMP), Good Laboratory Practices (GLP), and Good Documentation Practices (GDP). Experienced in supporting quality control activities through laboratory documentation, analytical data verification, instrument operation during laboratory practical sessions, and cross-functional collaboration with Quality Assurance, Production, and Warehouse teams. Familiar with pharmaceutical quality systems, data integrity principles, and regulatory compliance. Detail-oriented, analytical, and committed to ensuring product quality, laboratory compliance, and continuous improvement in pharmaceutical manufacturing.

CORE COMPETENCIES

Quality Control Operations

- Raw Material Testing Awareness
- Finished Product Testing Awareness
- Laboratory Analytical Testing
- Chemical Analysis
- Physical Testing
- Laboratory Documentation
- Sample Preparation
- Analytical Data Verification
- Analytical Instruments
- HPLC (Basic Operation)
- UV-Visible Spectrophotometer
- Atomic Absorption Spectroscopy (AAS)
- Analytical Balance
- Laboratory Equipment Operation
- Instrument Verification Awareness

Quality Systems

- GMP / CPOB
- Good Laboratory Practice (GLP)
- Good Documentation Practice (GDP)
- Data Integrity
- Regulatory Compliance
- Quality Control Documentation
- SOP Compliance

Technical Skills

- Microsoft Excel
- Microsoft Word
- Technical Documentation
- Analytical Thinking
- Attention to Detail
- Problem Solving
- Cross-functional Collaboration
- Continuous Improvement

EXPERIENCE

INTERN PHARMACIST

Feb 2025

PT MILLENIUM PHARMACON INTERNATIONAL – BANDUNG, INDONESIA

- Verified pharmaceutical distribution documentation to ensure compliance with GDP/CDOB requirements and documentation traceability.
- Supported inventory quality control through FIFO and FEFO implementation to maintain product quality and traceability.
- Utilized Microsoft Excel to monitor inventory records, documentation accuracy, and operational reporting.
- Collaborated with Warehouse and Operations teams to maintain compliance with pharmaceutical distribution procedures and quality standards.
- Maintained documentation accuracy and consistency in accordance with Good Documentation Practice (GDP).

INTERN PHARMACIST- QUALITY ASSURANCE UNIT

Aug-Sep 2024

PT. HOLI PHARMA – CIMAHI, INDONESIA

- Verified production documentation and Batch Manufacturing Records (BMR) to ensure compliance with GMP/CPOB requirements.
- Collaborated with Quality Control personnel in reviewing testing documentation for raw materials, intermediate products, and finished products against quality specifications.
- Supported quality compliance monitoring activities to minimize quality deviations throughout manufacturing operations.
- Assisted in quality documentation management to maintain audit readiness, data integrity, and documentation traceability.
- Reviewed operational documents to support Quality Management System (QMS) implementation and regulatory compliance.
- Worked closely with Quality Assurance, Quality Control, Production, and Warehouse teams to support pharmaceutical quality systems.

LABORATORY ASSISTANT – PHARMACEUTICAL ANALYTICAL LAB

2021 – 2023

DEPARTMENT OF PHARMACY, SUMATERA INSTITUTE OF TECHNOLOGY

- Assisted laboratory practical sessions in Pharmaceutical Analysis, Instrumental Analysis, Medicinal Chemistry, Biochemistry, and Basic Chemistry.
- Guided students in laboratory procedures, analytical techniques, and laboratory documentation.
- Ensured laboratory activities complied with Good Laboratory Practice (GLP).
- Prepared laboratory documentation and maintained documentation traceability.
- Supported laboratory operational readiness through accurate documentation and data verification.

EDUCATION

- **Apothecary Profession (Profesi Apoteker), STFI Bandung | GPA: 3.80 | Pharmaceutical Science OSCE Score: 93.33**
- **Bachelor of Pharmacy (Sarjana Farmasi), ITERA | GPA: 3.49**

TRAINING CERTIFICATIONS

- GMP / CPOB Training (2025)
- Quality Assurance in Pharmaceutical Industry (2025)
- ISO 9001:2015 Quality Management System Awareness Training (2026)
- Quality Control in Pharmaceutical Industry (2025)
- Research & Development in Pharmaceutical Industry (2025)