

SARAH RIFKA RAHMAWATI

Apothecary (License pharmacist)

Bogor, Jawa Barat. Mobile number [085211652727]. Born on April, 21st 2000.

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SUMMARY

Licensed Pharmacist with experience in pharmaceutical and cosmetic manufacturing, specializing in quality assurance, CAPA management, GMP documentation, and audit support. Knowledgeable in BPOM, CPOB, CPKB, ISO 22716, and ISO 9001 requirements, with a strong focus on regulatory compliance, data integrity, and continuous quality improvement.

EDUCATION

- ❖ Sekolah Tinggi Farmasi Indonesia | 2023-2024
Professional Apothecary | STRA Active
GPA 3.33 of 4.00
- ❖ Sekolah Tinggi Farmasi Indonesia | 2018-2022
Bachelor a degree – Farmasi
GPA 3.47 of 4.00

EXPERIENCE

- ❖ *Toll Manufacture In and Out at PT Inna Pharmaceutical Industry* | January 2025 – Now
 - Ensure toll manufacturing activities comply with GMP / CPOB requirements and internal quality assurance standards.
 - Review and verify batch manufacturing records, material documentation, and regulatory supporting documents to ensure accuracy and compliance.
 - Maintain traceability of raw materials, packaging materials, and finished products within toll manufacturing processes.
 - Collaborate with QA, QC, and Production teams to ensure proper documentation, compliance with quality systems, and timely batch release.
 - Support deviation investigations, CAPA implementation, and corrective actions related to toll manufacturing operations.
 - Participate in Quality Risk Management (QRM) activities by identifying potential risks in manufacturing processes and supporting risk assessment and mitigation actions.
 - Ensure manufacturing processes and documentation are aligned with applicable regulatory and quality standards.
- ❖ *Assistant Distributions Pharmacy at PT Enseval Putera Megatrading* | February 2024 – Des 2024
 - Accurately reviewed and managed all supporting evidence within the CAPA documentation process, ensuring full compliance, traceability, and alignment with BPOM requirements.
 - Proactively assisted and guided clients through the corrective action process, including re-uploading required documents to BPOM, to ensure timely and successful CAPA completion and regulatory approval
- ❖ *Quality Assurance Intern at PT TRIMAN Pharmaceutical Industry* | January 2024
 - Verified and ensured data integrity of PMP documentation in compliance with BPOM requirements, including recall and return traceability
 - Completed PMP database using QA batch records and delivered a raw material supplier audit report.

SKILL

1. CAPA Management (BPOM compliance), PMP, Recall & Return Handling, GMP / CPOB /CPKB, CPOTB Documentation
2. Audit Support, Supplier Evaluation & Documentation Control
3. Deviation & Corrective Action Tracking
4. English (TOEFL 520)
5. Cross-functional Communication
6. Microsoft Office, MS Project
7. Expert to handling Spectro, HPLC, GC instrument lab

ACHIEVEMENT

- Get funding research from DIKTI at PKM (Program Kreatifitas Mahasiswa) 2022 on title “Penentuan Kelarutan Kurkumin dalam Delapan Pelarut Organik Guna Pengembangan Sediaan Farmasi berbahan Dasar Kurkumin menggunakan Spektrofotometri *Visible* dan Gravimetri”. | 2022, March.

ORGANIZATION

- Member of IAI (Ikatan Apoteker Indonesia) | 2024-Now