



INDAH NUR SAFITRI

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Lampung timur, Lampung, Indonesia

Licensed Pharmacist with strong experience in GMP-compliant pharmaceutical manufacturing and production operations. Proven track record in supporting daily production activities, supervising production operators, handling deviations, CAPA, and change control, and ensuring compliance with CPOB/ GMP requirements. Experienced in cross-functional coordination with QA, QC, Engineering, and Maintenance to maintain efficient, safe, and audit-ready production processes. Detail-oriented, proactive, and ready to take responsibility as a Production Supervisor in a regulated manufacturing environment.

Education Level

Universitas Sriwijaya - Sumatera Selatan

Aug 2018 - Jun 2022

Bachelor of Pharmacy, 3.41/4.00

Universitas Muhammadiyah Purwokerto - Central Java

Feb 2023 - Mar 2024

Profession Pharmacist, 3.81/4.00

PROFESSIONAL EXPERIENCE

PT Pfizer Indonesia - Jakarta

Feb 2026 - Present

Project & Commissioning Engineer

- Responsible for commissioning, qualification and decommissioning activities as per agreed timeline
- Actively involved in equipment and instrument qualification activities (URS, DR, VRA, SLIA/CLIA, IOQ Protocol & Report)
- Developed and ensured successful implementation of change control actions across production areas
- Create and maintain SOP for the projects and after project handover
- Drafting essential project management documentation (i.e project management plan)
- Support in GMP documentation (i.e Drafting URS, change control, CAPA, support investigation if any)
- Support consultant/vendor site survey
- Execute project based on project management plan
- Verification on site material, drawings/as built drawings, invoice delivery by contractors
- Coordinate daily support with maintenance, utility and automation team when needed during construction, if there are any existing facilities, utilities and equipment

PT Pfizer Indonesia - Jakarta

Dec 2024 - Feb 2026

Production Project Specialist

- Maintenance project qualification in the production phase, aligned with CAPEX investment, CAPA implementation, and safety compliance.
- Actively involved in equipment and instrument qualification activities (URS, DR, VRA, SLIA/CLIA, IOQ Protocol & Report)
- Developed and ensured successful implementation of change control actions across production areas
- Responsibilities & Maintained SOP reviews and GMP documentation compliance in production areas
- Ensure instrument calibration compliance through assessment and monitoring of calibration schedules.
- Participate in audit readiness and regulatory inspections including BPOM, internal audits, MSQA, and global audits.
- Ensured workplace safety compliance (K3) through implementation of safety systems, risk assessments (HIRA, MSRA, LOTO, and incident investigation with CAPA)
- Conducted safety training and oversight to ensure operators follow production processes in accordance with approved safety procedures
- Supervised and supported production operators across shifts to ensure GMP compliance and adherence to approved production procedures
- Conducted training and GMP refreshment for production operators.
- Supported continuous improvement initiatives to enhance production compliance, efficiency, and documentation accuracy.
- Acted as liaison between Production, QA, QC, Engineering, and Maintenance during production and compliance activities.
- Assisted production management in decision-making related to quality and safety risks.

PT Pfizer Indonesia - Jakarta

Jun 2024 - Nov 2024

Production Pharmacist Compliance

- Support GMP-compliant pharmaceutical manufacturing with strong focus on quality compliance and documentation.
- Supervised and supported production operators across shifts to ensure GMP compliance and adherence to approved production procedures
- Conducted training and GMP refreshment for production operators.
- Supported continuous improvement initiatives to enhance production compliance, efficiency, and documentation accuracy.

- Acted as liaison between Production, QA, QC, Engineering, and Maintenance during production and compliance activities.
- Assisted production management in decision-making related to quality and safety risks.
- Support CAPA implementation, deviation follow-up, and quality issue resolution in production areas.
- Maintained SOP reviews and GMP documentation compliance in production areas
- Ensure instrument calibration compliance through assessment and monitoring of calibration schedules.
- Participate in audit readiness and regulatory inspections including BPOM, internal audits, MSQA, and global audits.
- Ensured workplace safety compliance (K3) through implementation of safety systems, risk assessments (HIRA, MSRA, LOTO, and incident investigation with CAPA)
- Conducted safety training and oversight to ensure operators follow production processes in accordance with approved safety procedures.
- Conducted annual defect trend analysis per product as part of production quality compliance and continuous improvement

PT Herbaemas Wahidatama - Central Java
QC Compliance Intern

Aug 2023 - Sep 2023

- Supported implementation of GMP, CPOTB, ISO 9001:2015, ISO 45001:2018, and SMK3.
- Prepared Quality Risk Management (QRM) documentation.
- Assisted in preparation and review of OOS/HULS and CAPA reports.
- Supported review of stability study data and cleaning validation documentation.

Organisational Experience

Community of Science at FMIPA UNSRI
Head of Division PKM/PMW

Jan 2021 - Jan 2022

Lampung Students' Association UNSRI (KEMALA)
Secretary of Dept Minat/Bakat

Aug 2020 - Aug 2021

Skills, Project & Achievements

- **Projects** (2026): Lean Six Sigma Yellow Belt – M1 Method
- **Webinars Attended** (2026): Good Cleanroom Practice- Contamination Control, Equipment cleaning & Cleanroom Cleaning Guidance Update by STERIS Corporation
- **Achievements** (2021): Finalist, National Scientific Festival (PIMNAS); Silver Medal Indonesian inventors Day 2021
- **Achievements** (2020): 3rd Place, National Essay Competition ; 3rd Place, National Creative Idea Competition