



FARID ALFAUZI

+6281216702077 | apt.faridalfauzi@gmail.com

A highly motivated Pharmacist from Surabaya University with a robust background spanning the pharmaceutical industry, distribution channels, and patient-facing services, I am driven by a strong passion for industrial pharmacy and eager to contribute to the growth of the pharmaceutical industry. I am dedicated, responsible, and adaptable, with a proven commitment to hard work, continuous learning, and delivering exceptional results in every challenge I take on

Education Level

Universitas Surabaya - Surabaya, East Java

Oct 2020 - Oct 2021

Apothecary Professional Program, 3.66/4.00

- President of the Apothecary Professional Program (PSPA) Class of LX

Universitas Surabaya - Surabaya, East Java

Aug 2016 - Aug 2020

Bachelor of Pharmacy, 3.27/4.00

- Undergraduate Thesis: Article Review Analysis of Sildenafil Citrate in Solid Sample of Stamina Herbal Medicine With Thin Layer Chromatography-Densitometry Methods

Work Experiences

PT. Sanbe Farma - Cimareme, West Bandung

Feb 2026 - Present

Sterile Large Volume Parenteral (LVP) Production Manager I

- LVP Manager I responsible for planning, controlling, and evaluating the performance of 10 sterile production lines.
- Responsible for the implementation of the LVP product manufacturing process starting from the initial material collection and weighing, production process, packaging process until the finished product is sent to the finished medicine warehouse.
- Coordinate with related parties so that the batch production and packaging process can run according to the schedule that has been determined and approved.
- Responsible for the ongoing review periodically, updating, and controlling any changes of Contamination Control Strategy (CCS) in the LVP Existing production area and initiates discussion on potential adjustments CCS.
- Provide instructions and direction to subordinate staff on both technical and administrative matters so that it can be ensured that there is a common view on everything that needs to be done.
- Responsible in training to subordinate and other related department.
- Prepare a production schedule for each batch to be made and ensure that the established schedule is implemented properly with PPIC Manager.
- Responsibilities for checking processing and packaging records for each batch that has been produced and ensuring that the manufacture of drugs has been carried out in accordance with established procedures and recorded in the batch record.
- Responsibilities with the QA Manager for an evaluation and determine possible solutions, that the product can meet the specified specifications for every failure or deviation in the drug manufacturing process.
- Responsibilities for conducting evaluations and supervision to ensure that every machine and tools used in the drug manufacturing process has been qualified and validated and used correctly.
- Assist in the implementation of CPOB inspections and be responsible for follow-up actions in accordance with the recommendations from the inspection results.
- Ensure that all employees involved in the drug manufacturing process have received CPOB training and other appropriate training so that employees have the required qualifications.
- Responsibilities for maintaining high performance and discipline of staff and employees, evaluating their performance and taking necessary corrective actions.
- Making reports, conducting inspections and evaluations reports from subordinate staff.
- Coordinating, evaluating and planning necessary corrective actions so that the drug manufacturing process can be more efficient and effective in accordance with the CPOB principles.

PT. Sanbe Farma - Cimareme, West Bandung

Nov 2024 - Feb 2026

Sterile Large Volume Parenteral (LVP) Production Assistant Manager I

- Assisted the LVP manager in planning, controlling, and performance evaluation of 10 sterile productions lines
- Coordination with LVP Production Manager I for the implementation of the LVP Expansion product manufacturing process starting from taking raw materials and weighing, production process, packaging process until the finished product is sent to the finished drug warehouse.
- Coordinate with related parties so that the batch production and packaging process can run according to the schedule that has been determined and mutually agreed upon.
- Responsibilities for the ongoing periodic review, updating, and controlling any changes of Contamination Control Strategy (CCS) in the

LVP Expansion production area and initiates discussions on potential CCS adjustments.

- Provide instructions and direction to staff below him on both technical and administrative matters so that there can be a shared view regarding everything that has to be done.
- Prepare a production schedule for each batch to be made and ensure that the schedule is implemented properly together with PPIC Staff.
- Responsible in training to subordinates and other related department.
- Responsible for checking processing and packaging records for each batch that has been produced and ensuring that drug manufacturing has been carried out in accordance with established procedures and recorded in the batch record.
- Coordination with LVP Production Manager I and QA Manager for an evaluation and determine possible solutions, so that the product produced can meet the specified specifications for every failure or deviation in the drug manufacturing process.
- Responsibilities for conducting evaluations and supervision to ensure that every machine and tools used in the drug manufacturing process has been qualified and validated and used correctly.
- Assist in the implementation of CPOB inspections and be responsible for follow-up actions in accordance with the recommendations from the inspection results.
- Ensure that all employees involved in the drug manufacturing process have received CPOB training and other appropriate training so that employees have the required qualifications.
- Responsibilities for maintaining high performance and discipline of staff and employees, evaluating their performance and taking necessary corrective actions.
- Making reports, conducting inspections and evaluations reports from subordinate staff.
- Coordinating, evaluating and planning necessary corrective actions so that the drug manufacturing process can be more efficient and effective in accordance with the CPOB principles.

PT. Sanbe Farma - Cimareme, West Bandung

Nov 2021 - Sep 2024

Sterile Large Volume Parenteral (LVP) Production Supervisor

- Supervised daily operations of 10 sterile production lines in compliance with GMP standards and production targets.
- Supervised the LVP (Large Volume Parenteral) production process, including mixing and product transfer, primary packaging material preparation, form-fill-seal operations, product sterilization, sterile garment preparation, and sanitation/disinfection of grey and white areas in the LVP plant, ensuring full compliance with SOPs and production schedules.
- Coordinated routinely with production staff to ensure batch production activities were completed according to the approved schedule.
- Provided clear technical and administrative instructions and guidance to subordinates to ensure consistent understanding and execution of assigned tasks.
- Reviewed and verified batch manufacturing records prior to consolidation with batch packaging records, and examined production logbooks in the LVP processing areas.
- Reported process deviations or product failures to the LVP Production Manager and collaborated with the QA Manager to conduct evaluations, determine corrective actions, and prevent recurrence.
- Ensured all required equipment was available, properly cleaned, sanitized, and sterilized in accordance with established procedures.
- Conducted reconciliation of primary packaging materials and LVP products to ensure accuracy and compliance.
- Implemented controls to prevent mix-ups and cross-contamination risks.
- Coordinated with the Calibration and Engineering departments to perform calibration of production equipment and instruments, ensuring calibration labels were properly affixed and within validity periods.
- Ensured cleanliness of LVP production areas and the effective implementation of sanitation programs in accordance with SOPs.
- Prepared purchase requisition forms for equipment and materials supporting smooth LVP production operations.
- Participated in evaluations to ensure all personnel involved in drug manufacturing received GMP (CPOB) and other required training, meeting established qualification standards.
- Prepared required production reports and reviewed and evaluated reports submitted by subordinate staff.
- Contributed to the evaluation and planning of corrective and improvement actions to enhance efficiency and effectiveness of drug manufacturing processes in compliance with GMP principles.
- Analyzed potential occupational safety risks within the plant and proposed corrective actions to reduce workplace accidents, such as electrical repairs, slip hazard mitigation, improved lighting, and use of safety equipment.
- Ensured all subordinates consistently used appropriate personal protective equipment (PPE) such as safety goggles, helmets, and other required safety gear.
- Participated in planning and evaluation of occupational health and safety programs.

PT. Sanbe Farma - Cimareme, West Bandung

Nov 2021 - Sep 2024

Halal Supervisor for Sterile Products

- Coordinate with Production, Quality Assurance, and related departments to ensure the implementation of halal requirements in sterile manufacturing and packaging processes.
- Ensure that all raw materials, packaging materials, processing aids, cleaning agents, and sterilization-related materials used in sterile products are halal-approved and supported by valid halal documentation.
- Supervise the implementation and ongoing review of the Halal Assurance System (HAS) in sterile production areas, including integration with Contamination Control Strategy (CCS).
- Review and approve halal-related documents, including SOPs, batch records, deviation reports, change controls, and risk assessments

related to halal compliance.

- Coordinate with Validation and Engineering teams to ensure that sterilization processes, cleaning validation, and equipment qualification comply with halal requirements.
- Investigate halal deviations, non-conformities, or complaints and ensure the implementation of corrective and preventive actions (CAPA).
- Provide halal training and awareness programs to employees involved in sterile production and supporting activities.
- Prepare reports related to halal compliance, audit results, and halal assurance system performance.
- Coordinate with halal certification bodies and regulatory authorities regarding halal certification, surveillance, and renewal processes for sterile products.
- Continuously evaluate and improve halal compliance to ensure alignment with applicable halal standards (e.g. HAS 23000) and company policies.

PT. Satoria Pharma - Pasuruan, East Java

Apr 2021 - May 2021

Pharmacist Professional Internship

- Involved in New Product Development of Lidocaine HCl 2% from formulation study to production readiness
- Learned sterile product manufacturing processes, including aseptic processing and terminal sterilization
- Gained hands-on understanding of GMP/CPOB implementation in sterile manufacturing

Organisational Experience

Apoteker Berbagi 2023 - Padalarang, West Bandung

Mar 2023 - Apr 2023

Public Relations & Fundraising Coordinator

- Coordinated public relations strategies to enhance the visibility and public image of the Apoteker Berbagi program.
- Established and maintained communication with sponsors, donors, and external partners.
- Prepared and managed fundraising proposal and financial utilization reports.
- Coordinated public relation and fundraising teams to ensure funding and publicity target were achieved
- Managed event publications through social media and official communication channels.

Teaching Assistant at University of Surabaya - Surabaya, East Java

Aug 2019 - Aug 2020

Participant

- Assisted lecturers in course preparation and delivery course materials
- Supported students during practical sessions and discussions
- Helped evaluate assignments and academic activities

Student Executive Board (BEM), Faculty of Pharmacy - University of Surabaya - Surabaya, East Java

Sep 2018 - Sep 2019

Coordinator of Student Resource Development

- Planned and coordinated student development programs
- Led and supervised team members in leadership and soft-skill activities
- Evaluated student performance and organizational effectiveness

ISMAFARSI Regional Conference - East Java - Bali

Jan 2018 - Jan 2019

Participant

- Participated in regional meetings of pharmacy student associations
- Contributed to discussions on organizational development and leadership
- Expanded Professional networking among pharmacy student

Student Executive Board (BEM), Faculty of Pharmacy - University of Surabaya - Surabaya, East Java

Sep 2017 - Sep 2018

Member of Student Resource Development Division

- Assisted in organizing student development and leadership activities
- Supported monitoring and training programs for students
- Collaborated with team members to improve student competencies

Leadership Character Development & Basic Leadership Training (LKTD) - Trawas, East Java

Feb 2017 - Feb 2018

Chairperson

- Led the overall planning and execution of leadership training programs
- Coordinated committees and managed event timelines
- Developed leadership, teamwork, and communication programs for participant

Skills, Achievements & Other Experience

- **Achievements** (2018): Faculty of Pharmacy Student Activist Scholarship - University of Surabaya
- **Achievements** (2019): Faculty of Pharmacy Student Activist Scholarship - University of Surabaya
- **Projects** (2025): Implemented an auto labelling machine project on the packaging line to improve process efficiency and reduce manual

manpower requirements.

- **Projects** (2024): Implemented an automatic check weigher project on an existing packaging line to improve weighing accuracy, process efficiency, and reduce manual manpower requirements through automation
- **Projects** (2023): Managed Sterile Product Expansion 2 involving the addition of two new production lines.
- **Projects** (2023): Implemented a 2D barcode system for sterile product to enhance traceability, regulatory compliance, and production system integration.
- **Forum Discussion** (2025): Sartorius Lunch & Learn : Validate with Confidence
- **Familiar With:** CPOB, CDOB, CPAKB, CDAKB, SJH