



Tiska Lestari

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Jl. Jatisari Blok B10 No 124 Jaticempaka Pondok Gede Bekasi

PROFESSIONAL PROFILE

Pharmaceuticals Manufacturing Lead with experience the production of non-sterile and sterile pharmaceuticals while ensuring compliance with SOP, GMP, safety and quality guidelines. Proficient at ensuring manufacturing project with agree time plan and quality standar for product, so it can safe for consumption by public.

CORE SKILLS

- Manufacturing Operations
- GMP Standar
- Standar Operating Procedures
- Batch Manufacturing
- Sterile and Non Sterile Manufacturing
- Aseptic Manufacturing
- Electronic Batch Record

CARRER SUMMARY

Senior Production Supervisor

2023 - Present

PT Ferron Par Pharmaceuticals Cikarang

- Reporting production output and quality aspect to Production Manager.
- Planning, Organizing Actuating and Controlling production Process in order to meet the GMP requirements and target from company.
- Review production capacity and improve production process.
- Collaboration with internal departement and external departement related production process.
- Improvement process :
 1. Developing Process effectivity for Sterile 1 Line (Eyedrop Line).
 2. Investigate and solving problem in production process based on OEE for Sterile 2 and Sterile 3 Line.
 3. CAPA effectiveness.
- FAT for Blistering, Cartoning and Case Packing Ampoule and Cartoning Vials for FPP Extension Sterile 2 and Sterile 3.
- FAT for MG2 Planeta for capsule in Solid 2 Line.

Sterile Line 3 Production Supervisor

2022 - 2023

PT Ferron Par Pharmaceuticals Cikarang

- Reporting production output and quality aspect to Production Operation Manager for target and goals.
- Planning, Organizing Actuating and Controlling production Process in order to meet the GMP requirements and target from company.
- Investigate and solving problem in production process.
- Develops or revises Standar Operational and Working Practices and observes workers to ensure compliance with standards.
- Monitoring Overall Equipment Effectiveness for filling and packaging process.
- Organise the repair of any damaged equipment liasie with different departments, teams, and companies. e.g supplier, managers, client.
- Complete detailed quality system documentation including deviation, Change Control, CAPA.
- Ensure that health and safety guidelines are followed at all times.
- Coordinate the day-to-day activities of the team and provide training for subordinate.

Electronic Batch Record Champion
PT Ferron Par Pharmaceuticals Cikarang

2021 - 2022

- Develop for MBR (Manufacturing Batch Record) design production electronic batch record.
- Training all production supervisor/officer/staff, operator, packer for using electronic batch record.

Solid Production Supervisor

2013 - 2020

PT Ferron Par Pharmaceuticals Cikarang

- Reporting production output and quality aspect to Production Operation Manager for target and goals.
- Planning, Organizing Actuating and Controlling production Process in order to meet the GMP requirements and target from company.
- Investigate and solving problem in production process.
- Develops or revises Standar Operational and Working Practices and observes workers to ensure compliance with standards.
- Organise the repair of any damaged equipment liasie with different departments, teams, and companies. e.g supplier, managers, client.
- Complete detailed quality system documentation including deviation, Change Control, CAPA.
- Ensure that health and safety guidelines are followed at all times.
- Coordinate the day-to-day activities of the team and provide training for subordinate.
- MHRA (UK) certification for Solid 1 Line.

Quality Supervisor

2012 - 2013

PT Mitrapak Eramandiri Cikarang

- Reporting quality aspect to Quality Manager for target and goals.
- Reporting quality aspects to the Quality Manager.
- Training related to PT Unilever's product specification for implementation at PT Mitrapak.
- Collaborating with internal departments, external departments, and external parties (PT Unilever) regarding quality.
- Creating batch procedures for product filling and packaging processes.
- IPC training for quality personnel regarding product specifications.

Production Solid Supervisor For Filling Capsule and Primary Packaging

2010 - 2012

PT Pyridam Farma, Tbk Cipanas

- Reporting production output and quality aspect to Production Manager for target and goals
- Planning, Organizing Actuating and Controlling production Process in order to meet the GMP requirements and target from company.
- Investigate and solving problem in production process.
- Develops or revises Standar Operational and Working Practices and observes workers to ensure compliance with standards.
- Organise the repair of any damaged equipment liasie with different departments, teams, and companies. e.g supplier, managers, client.

Apoteker

2009 - 2010

Apotek Farmarin

- Responsible for all pharmacy operation and performance.
- Planning and creating purchase order for drug procurement.
- Receive and check all drugs coming into pharmacy.
- Prepare medication according to doctor's prescription.
- Manage and supervise pharmacy staff

EDUCATION

University / School	Faculty	Discipline	GPA	Dates From to
SDN Malaka Jaya 05 Pagi Jakarta	-	-	43,18	1991-1997
SMPN 139 Jakarta	-	-	42,37	1997-2000
SMUN 61 Jakarta	-	IPA	85,36	2000-2003
University of Indonesia	FMIPA	Pharmacy (Apothecary)	3,15	2003-2008