



OKIN

Quality Assurance | Production

Contact



Palembang, 11 October 1992



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Griya Indah Serpong,
Gunung Sindur, Kab Bogor, Jabar

Education

- 2016-2017 **Apotechary**
UIN Syarif Hidayatullah JKT
- 2012-2016 **Pharmacy**
UIN Syarif Hidayatullah JKT

Achievements and Training

- Bronze Award BIID Distributor Transition Project
- 2nd Winner Urgent Supply for Covid Support Drugs
- LEAN Manufacturing and Sterile/Aseptic
- SIPOC, PDCA, OEE

Expertise

- Social Perceptiveness
- Critical Thinking
- Development and Building Teams
- Time management
- Hard and smart working, responsible and willing to learn

Profile

A process-oriented and results-driven Quality Assurance and Production who has a passion for excellence. Experience of responsible on the wide variety of manufactured goods (Oral dosage forms, Suppository and Nasal/Inhalation Spray) possesses a successful track of determining whether products meet expected standards. I continuously monitoring and improvement process, handling inventory in the warehouse and documentation as Health Authority regulations.

Professional Experience

Ferring Pharmaceuticals Industry

Teamleader Production (Oct 2023 - Presently)

- Designing SOPs, Work Instructions, Protocol Validation and Batch Records related to the process in Sterile Technologies Facilities.
- Perform Project for Process Validation Activities and Stability Study
- Perform maintenance, repairing, arrangement and lead Inventory WH
- Perform routine reporting process (materials and products) to BPOM and Kemenkes RI

Boehringer Ingelheim Indonesia

QA Inspector, GMP Officer (Dec 2017 - Sep 2023)

- Responsible for checking that a company's products meet the spec determined and no impact on product quality.
- Investigation of complaint, deviation and defect materials after QC release/non-conformity.
- Ensure that the application of SOP is in accordance with GMP requirements in all daily GMP activities in the factory.
- Responsible for ensuring factory audit readiness.
- Review the doc. related process, batch records prior to release.
- Monitor the product release process according to schedule and prepare data for KPI report in QA for weekly and monthly reports.
- Conduct training related to GMP activities in the production and warehouse team, develop operators and analysts to raise their quality awareness and continuously improve their standard of working.
- Coordinate with toll manufacturing related to the third party product batch record review and its disposition timeline.
- Contribute ensure validation activity of specification test for bulk identification is carried out according to cGMP Req and guidelines.

As a Quality and Production Team routinely oversees the overall production process, relating to win-win solutions over quality, compliance, business, and safety.

Additional Skills and Interest

Technical Skills: Ms Programs (Word, Excel, Powerpoint), Adobe

Interpersonal skills: Public Speaking, Team leadership, problem solving & communication.

Interest: Pharma Sterile, Tech Solid, Liq, Semi-solid Dosage Forms, Medical Device, Cosmetics Tech, Logistic/SCM, Digital Pharmacy, Improvement Projects, Quality Management System.