

apt. Sarah Marissa,S.Farm



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sarahmarissaa



PROFESIONAL STATEMENT

Pharmacists who never stop learning, has good experience in R&D, Quality, and Regulatory. Good leadership, good communication, have a hight motivation, and I can do my best to give positive contribution toward the company.

EDUCATION

Pharmacist Industry	2018
Pancasila University, Jakarta	
Bachelor Degree Pharmaceutical Industry	2016
Pancasila University, Jakarta	

WORK EXPERIENCES

PT. Setia Kawan Abadi As QA & MR Manager (Traditional Medicine & Cosmetics)	October 2024- Current
- Review and approved batch record - Control & Monitoring closure of CAPA, deviation, Chang control, Internal audit, Customer Complaint,COA - Prepare, review and approved validation documents, qualifications, procedures from all departments, QC specifications, supplier complaint management, and all documents related to Quality - Create Weekly report, monthly reports, and present management review to top Management - Register Halal Products - Share facility licensing - Monitoring implementation of ISO 9001 - Ensure that the calibration is carried out in accordance with the schedule - Create Quality Risk Management	
PT. Adev Natural Indonesia As Quality & Regulatory Manager(Skincare and Cosmetics)	July 2023 -June 2024
- Review batch records and approve finished products - Control, supervise, and participate in the registration process for new products and existing products - Receive finished product complaint reports, identify and investigate root cause analysis, and recommend CAPA investigation results - Monitoring the achievement of KPIs for the QA, QC, & Regulatory divitions - Create weekly report, monthly reports, and present management reviews to top Management - Conduct regular internal and external quality control audits to ensure compliance with established standards - Develop quality systems such as CAPA, CPKB, deviation, risk assessment, and implementation of ISO 9001 - Prepare, review, and approve validation documents, qualifications, procedures from all departments, QC specifications, supplier complaint management, and all documents related to quality	

PT. LF Beauty Manufacturing Indonesia, Jakarta**April 2021 -July 2023**

As QA Release Supervisor(Skincare and Cosmetics)

- Ensure batch record and QC analysis result that have been filled by related parties and compliance with the GMP conditions
- Ensure all changes, deviation, concession request and process validation which related with the batch have been dealt with related procedures
- Check result of Critical Process Parameter & Critical Quality Attribute
- Verify the correctness CoA finished Product
- Lead investigation, create complaint report, Follow up all complaints related to product quality complaint,
- Monitoring CAPA implementation as a result from investigation, and make Review and trend of complaint monthly by showing year to month complaint data, receipt and explain in detail the investigation result to complainant
- Create Product Quality Review / Annual Product Quality Review
- Create Monthly report for related Customer
- Part of Continuous Improvement Leader
- Handling Mock Recall & Recall Product

PT. Taisho Pharmaceutical Indonesia, Jakarta**July 2020-December 2020**

As QA Compliance Staff (Pharmaceutical Industry)

- Register change control from all departments, hold meetings with relevant departments according to the matrix to discuss change control and its CAPA provide an assessment of change control from a Quality point of view
- Ensure fulfilment of CAPA from internal audit (SIP deviation, change
- Validation/qualification APQR. Risk Assessment, etc.) and External Audit
- Audit held third party and authority bodies) in timely manner
- Coordinate Change Request Meeting and Assessment. Ensure fulfilment of Change Request CAPA and closure in timely manner
- Coordinate monthly report quality council
- Coordinate training fundamental

PT. Medifarma Laboratories, Jakarta**April 2018 - April 2020**

As Formulator Staff(Pharmaceutical Industry)

- Design formula of new product development
- Reformulation existing product -Monitoring lab and pilot trial
- Monitoring alternate source development Monitoring stability of project, and investigate if any issue
- Prepare supporting doc for lab & pilot trial (batch record, weighing record, flowchart, etc.)
- Create ACTD Dossier and Protocol Report Validation for registration

SKILLS

- Supervisory skills
- English
- Microsoft word
- Hard worker
- Discipline
- Able to work under pressure
- Able to work in team and individual
- Good communication
- Managerial skill