

DHITA GUSTIEN ANDRIANI, S.Farm., Apt.

Head of Formulation | Pharmaceutical R&D

Cimahi, Indonesia

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PROFESSIONAL SUMMARY

Experienced pharmaceutical R&D professional with more than 6 years of experience in formulation development and existing product improvement within the pharmaceutical industry. Skilled in formulation trial, validation, documentation, technology transfer, and cross-functional coordination with QA, Regulatory Affairs, Production, QC, and Purchasing teams. Experienced in handling multiple dosage forms including sterile, liquid, semisolid, and health supplement products. Strong understanding of GMP/CPOB compliance, pharmaceutical documentation, and project coordination.

PROFESSIONAL EXPERIENCE

PT Sanbe Farma — Cimahi, Indonesia

Head of Formulation for Existing Product

August 2024 – Present

Key Responsibilities:

- Lead formulation activities for existing pharmaceutical products in accordance with updated regulations and product requirements.
- Review and approve formulation-related documents including:
 - Batch Analysis
 - Quality Overall Summary (QOS)
 - Process Validation Protocol & Report
 - Pharmaceutical Development Report
 - Master Production Document
 - Technology Transfer Document
- Analyze trial and production data for process improvement and formulation optimization.
- Coordinate formulation staff and pharmacy assistants to improve team efficiency and workflow.
- Collaborate with QA, Regulatory Affairs (local and export), and Quality teams to support renewal and variation registration processes.
- Coordinate with Purchasing for comparator product sourcing and raw material procurement.

Formulation Staff

December 2019 – February 2024

Key Responsibilities:

- Conduct formulation trials for existing products including sterile, liquid, semisolid, and health supplement dosage forms.
- Perform product reformulation and raw material substitution based on updated regulations and pharmacopeial requirements.
- Conduct simple physical evaluations including pH, viscosity, and homogeneity testing.
- Prepare and maintain formulation documentation and pilot batch records.

- Review production batch records and production work instructions.
 - Coordinate with QC, Production, and Analytical R&D teams for stability evaluation and formula improvement.
 - Ensure compliance with GMP/CPOB standards and internal procedures.
 - Prepare technical and validation documents including:
 - Batch Analysis
 - Quality Overall Summary (QOS)
 - Process Validation Protocol & Report
 - Pharmaceutical Development Report
 - Master Production Document
 - Technology Transfer Document
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EDUCATION

Institut Teknologi Bandung (ITB) — Bandung, Indonesia

Apothecary Profession Program (Apoteker)

2019

Bachelor of Pharmacy (S.Farm.)

Major in Pharmaceutical Science and Technology

2018

CORE SKILLS

Pharmaceutical & Technical Skills

- Formulation Development
- Existing Product Improvement
- Technology Transfer
- Process Validation
- GMP / CPOB Compliance
- Pharmaceutical Documentation
- Regulatory Support Documentation
- Stability Evaluation
- Cross-functional Coordination

- Spectrophotometer
- pH Meter
- Viscometer
- Osmometer
- Freeze Drying

Software Skills

- Microsoft Word
- Microsoft Excel
- Microsoft PowerPoint

Laboratory & Analytical Skills

- HPLC
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LANGUAGES

- Indonesian (Native)
 - English (Professional Working Proficiency)
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INTERNSHIP EXPERIENCE

Apotek Medika Antapani — Bandung

PKPA Internship

April 2018

- Provided drug information services to patients.
- Prepared and dispensed medicines according to prescriptions.
- Assisted stock opname and inventory monitoring activities.
- Compounded prescriptions in accordance with pharmacy procedures.

Directorate of Drug and Food Investigation — BPOM RI, Jakarta

PKPA Internship

March 2019

- Assisted in preparing infographic materials regarding illegal cosmetics for public awareness.
- Organized and documented evidence data from BPOM regional offices to support investigation administration.

PT Corsa Laboratories — Tangerang

Quality Assurance Intern

January 2019 – February 2019

- Assisted QA team in preparing Product Quality Review (PQR) documentation.
- Compiled statistical product data for quality evaluation and reporting purposes.

PT Kimia Farma — Bandung

Internship

June 2016 – August 2016

- Compiled raw material testing procedures from various pharmacopeias.
- Monitored reagent availability for laboratory testing.
- Assisted in documenting analytical testing methods and product improvement activities.

ADDITIONAL INFORMATION

- Nationality: Indonesian
- Marital Status: Married