

July 17, 2026

apt. Indra Novian Rukmana, S.Farm.

Bandung, Jawa Barat
(+62)85314692924
indranovianr@gmail.com
linkedin.com/in/indranovianrukmana

Dear Hiring Manager,

I am writing to express my interest in the Analytical Development (Andev) Staff position at PT Mersifarma TM. As a licensed pharmacist with professional internship experience in the pharmaceutical industry, I am eager to contribute my analytical skills, scientific knowledge, and commitment to quality in supporting your Analytical Development team.

During my professional internship (PKPA) at PT Bio Farma (Persero), I was assigned to the Transfer Project Division, where I gained exposure to pharmaceutical manufacturing processes in a GMP/CPOB-compliant environment. My responsibilities included assisting in IQ/OQ qualification activities, reviewing Batch Production Records (BPR), performing Water for Injection (WFI) and Pure Steam sampling, and conducting regulatory research on 10 QA findings by benchmarking CPOB, WHO TRS, EudraLex, and EU GMP guidelines. These experiences strengthened my analytical mindset, attention to detail, and understanding of quality and regulatory compliance in pharmaceutical manufacturing.

In addition, my undergraduate research focused on the development of a natural lip cream formulation, where I conducted physical characterization and stability evaluations using UV-Vis Spectrophotometry. Through this research, I developed hands-on experience in experimental design, data analysis, scientific documentation, and interpretation of analytical results. This experience reinforced my interest in analytical work and inspired me to pursue a career in Analytical Development.

I am a detail-oriented, adaptable, and proactive individual with strong analytical thinking, problem-solving, and communication skills. I enjoy working with scientific data, conducting investigations, and continuously learning new analytical techniques. I am confident that my educational background, internship experience, and enthusiasm for analytical science would allow me to contribute positively to PT Mersifarma TM.

Attached is my Curriculum Vitae (CV) for your consideration. I would welcome the opportunity to discuss how my qualifications, skills, and enthusiasm align with the Analytical Development (Andev) Staff position at PT Mersifarma TM.

Thank you for your time and consideration. I look forward to contributing to PT Mersifarma TM.

Sincerely,

apt. Indra Novian Rukmana, S.Farm.



INDRA NOVIAN RUKMANA

+6285314692924 | indranovianr@gmail.com | linkedin.com/in/indranovianrukmana

Bandung, Jawa Barat

Licensed pharmacist with professional internship experience and a strong interest in the pharmaceutical industry, with knowledge of pharmaceutical manufacturing, GMP/CPOB principles, and pharmaceutical regulations. Detail-oriented, adaptable, and proactive with strong analytical thinking, problem solving, and communication skills. Eager to apply my knowledge and skills to support quality, innovation, and continuous improvement in the pharmaceutical industry.

Education

Universitas Islam Bandung – Bandung, Indonesia

Feb 2025 – Apr 2026

Pharmacist Professional Program, 3.88/4.00

- Graduated with Cum Laude honors, completing the Professional Pharmacist Program within 1 year.
- Contributed to multiple projects during the Professional Pharmacist Internship.

Universitas Islam Bandung – Bandung, Indonesia

Sep 2020 – Jul 2024

Bachelor's Degree in Pharmacy, 3.52/4.00

- Graduated with Cum Laude honors, completing the Bachelor's Degree within 4 years.
- Final Project: "Color Stability Test of Lip Cream Formulation with Purple Sweet Potato (*Ipomoea batatas* (L.) Lam.) Ethanol Extract and Sodium Caseinate as Copigment"

Work Experiences

Rumah Sakit Umum Pusat Dr. Hasan Sadikin Bandung – Bandung, Indonesia

Oct 2025 – Nov 2025

Pharmacist Intern

- Provided hospital pharmacy services, including dispensing, sterile compounding, ADR monitoring, and patient counseling.
- Managed 100+ outpatient or 50+ inpatient prescriptions daily, conducted ward rounds, and reviewed 150 prescriptions.
- Performed TDM for 2 patients, evaluated hospital formulary compliance, and promoted rational antibiotic use.

Badan Pengawas Obat dan Makanan RI – Jakarta, Indonesia

Sep 2025

Directorate of Standardization Intern

- Conducted scientific and regulatory assessments of pharmaceutical substances (erythrosine and etomidate).
- Benchmarked international regulatory guidelines, including CPOB, CDOB, GMP, GDP, ICH, WHO, FDA, and EMA.
- Prepared a CAPA plan based on CPOB and CDOB non-conformity cases, developed a GCP gap analysis matrix to support the revision of the previous CUKB guideline, and participated in cross-functional regulatory meetings.

PT. Kuda Mas Jaya Makmur Sentosa – Bandung, Indonesia

Aug 2025

Pharmacist Intern

- Participated in pharmaceutical distribution activities in compliance with CDOB standards, supporting product quality, traceability, inventory management, and FEFO implementation.
- Verified 50+ orders daily, reviewed 50 invoices and 50 purchase orders, and performed final order validation and e- Report documentation.

Apotek Medika Antapani – Bandung, Indonesia

Jul 2025

Pharmacist Intern

- Delivered pharmaceutical care, including dispensing, drug information services, patient counseling, and self-medication support.
- Managed 30+ prescriptions daily, reviewed 75 prescriptions, conducted a pharmacy feasibility study, and performed pareto based inventory analysis.

PT. Bio Farma (Persero) – Bandung, Indonesia

May 2025 – Jun 2025

Research & Development (R&D) – Transfer Project Intern

- Contributed to cross-functional Transfer Project activities involving Production, QC, QA, and R&D within a GMP/CPOB-compliant pharmaceutical manufacturing environment.
- Assisted in IQ/OQ qualification, SOP development, batch production record review for 3 production batches, and WFI & pure steam sampling to support product quality and regulatory compliance.
- Conducted regulatory assessments of 15 QA findings by benchmarking CPOB, WHO TRS, EudraLex, and EU GMP requirements to support continuous quality improvement.
- Conducted technical studies on pharmaceutical manufacturing and regulatory topics, including QbD, vaccine manufacturing, cleanroom operations, drug registration, Supply Chain Management, and HSE (K3L).

Skills

• Hard Skills:

- **Laboratory Instruments:** UV-Vis Spectrophotometer (Advanced), HPLC (Basic), FT-IR Spectroscopy (Basic)
- **Pharmaceutical Knowledge:** GMP/ CPOB Knowledge | Qualification (IQ/ OQ) | Pharmaceutical Manufacturing | Pharmaceutical Regulatory | Documentation & Technical Reporting | Batch Production Record Review | SOP Development | Basic Data Analysis | Inventory Management
- **Software:** Microsoft Office (Word, PowerPoint, Excel), Google Workspace (Docs, Slides, Sheets), Canva

- **Soft Skills:** Detail- oriented, Responsible, Persistent, and Focused | Critical & Analytical Thinking | Problem Solving | Time Management | Teamwork & Independent Working | Adaptability | Quick Learner | Willingness to Learn

- **Languages:** Indonesian (Native), English (Intermediate)