

PUTI APIFAH CAHYANA

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PROFESSIONAL SUMMARY

Pharmaceutical and Food Analysis graduate from Poltekkes Kemenkes Jakarta II, with hands-on experience in method validation, verification, and instrumental analysis within pharmaceutical industry settings. Knowledge of GMP, GLP, and laboratory documentation. Highly detail-oriented with hands-on experience in HPLC, AAS, UV-Vis spectrophotometry, dissolution testing, TLC-spectrophotodensitometry, Tablet Compression, Tablet Coating, Granulation, and Homogenization. Seeking an entry-level R&D Analytical Development or Chemical Analyst or Formulation Development role to contribute to robust and regulatory-compliant method development.

EDUCATION

Diploma III in Pharmaceutical and Food Analysis – Poltekkes Kemenkes Jakarta II (2022–2025)
GPA: 3.63

RESEARCH EXPERIENCE

Final Project: Method Validation for Identification of Ursolic Acid in Marasai Leaf Extract Using TLC-Spectrophotodensitometry

- Validated analytical method including accuracy, precision, linearity, LOD and LOQ.
- Performed calibration curve preparation and statistical data evaluation.
- Conducted method optimization to ensure reproducibility and reliability.
- Prepared structured scientific documentation and reporting.

PROFESSIONAL EXPERIENCE

Analytical Development Intern – PT Novell Pharmaceutical Laboratories (Feb 2025 – Mar 2025)

- Performed method validation and method verification for finished pharmaceutical products.
- Conducted assay and quantitative analysis of raw materials and finished products.
- Prepared working standards and calibration curves.
- Prepared mobile phase and conducted sample preparation.
- Performed data processing and documentation in compliance with GMP and SOP.
- Prepared analytical reports and Lembar Kerja Analisis (LKA).

R&D Formulation Development – PT Pharos Indonesia (Sep 2025 – Present)

- Assisted in formulation trials including weighing, mixing, granulation, compression, sealing and coating.
- Performed IPC and physical evaluation of trial batches.
- Reviewed CoA and pre-formulation data.
- Prepared samples for Analytical Development testing.
- Assisted in development documentation and reporting.
- Operated tablet press machinery to manufacture tablets and caplets in strict compliance with quality and weight specifications.
- Executed tablet coating processes to enhance product stability, aesthetic appearance, and active pharmaceutical ingredient (API) release profiles.
- Conducted formulation trials for semi-solid dosages (creams, emulgels) and liquid dosages (syrups) during the product development phases.

TECHNICAL SKILLS

- Formulation & Development: Solid (Tablets, Caplets), Semi-Solid (Creams, Emulgels), and Liquid (Syrups) Dosages.
- Pharmaceutical Technology: Tablet Compression, Tablet Coating, Granulation, and Homogenization.
- Analytical Instrumentation: HPLC, UV-Vis Spectrophotometry, Polarimetry, AAS, Karl Fischer.
- Laboratory Testing: Dissolution, Friability, Hardness, Viscosity, and pH Testing.
- Regulatory Compliance: GMP, GLP, and R&D Documentation.
- Software: Microsoft Office (Excel, Word, PowerPoint)

SOFT SKILLS

- Analytical Thinking
- Attention to Detail
- Problem Solving
- Time Management
- Team Collaboration