



Jason Eldwin

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Degree : Pharmacy (S.Farm.), Apothecary (apt.)

ABOUT ME

Industrial Pharmacist with **years of experience in pharmaceutical manufacturing**, specializing in **Quality Management Systems, GMP/CPOB compliance, Product Quality Review (PQR), data integrity, and audit readiness**. Proven capability to operate at **managerial level**, leading cross-functional quality initiatives, supporting regulatory inspections, and driving **digital transformation (eQMS, EBR, document digitalization)**. Experienced in both **domestic onsite audits and international remote audits**, with strong scientific and risk-based decision-making skills.

Skills

- Quality Management System (QMS)
- GMP / CPOB Compliance
- Internal & External Audits (Onsite & Remote)
- Data Integrity (ALCOA++)
- Document Management
- Cross-Functional Leadership
- eQMS & Digital Quality Transformation
- Statistical Data Analysis

Experience

- QA Senior Supervisor at PT. Interbat (February 2024 – present)
- QA Staff at PT. Interbat (September 2023 – January 2024)
- Quality Control & RnD (Apoteker Penanggung Jawab) at PT. Gra Parianom Sejati (July 2022 – July 2023)
- Stability and Cleaning Validation Supervisor at PT. Guardian Pharmatama (July 2021 – July 2022)
- Assistant Lecturer at Widya Mandala Catholic University of Surabaya, Faculty of Pharmacy (February 2018 – January 2020)
 - Analytical Chemistry Laboratory
 - Formulation and Development of Semisolid and Liquid Dosage Form Laboratory
 - Pharmacognosy and Phytocemistry Laboratory

Education

- Apothecary at Widya Mandala Catholic University of Surabaya (Graduated in 2021 with 3.93 GPA)
- Bachelor of Pharmacy at Widya Mandala Catholic University of Surabaya (Graduated in 2020 with 3.63 GPA)
- High School at Petra 2 Christian High School (Graduated in 2015)

Achievements

- Patent of "Teknologi Produksi Granul Ekstrak Sinom". Patent number: 000178264 (December 2019 – December 2069)
- Patent of "Teknologi Produksi Granul Effervescent Sinom". Patent number: 000178253 (December 2019 – December 2069)
- The Best Apothecary at Widya Mandala Catholic University of Surabaya (55th Period of Apothecary, March 2021)
- Dexa Award from PT. Dexa Medica (April 2021)



PROFESSIONAL EXPERIENCE

QA Senior Supervisor

PT. Interbat | Indonesia

February 2024 – Present

- Lead and oversee Quality Assurance activities to ensure consistent compliance with GMP/CPOB and continuous readiness for internal audits and regulatory inspections.
 - Manage and control key elements of the Quality Management System (QMS), including document control and data integrity governance.
 - Act as the Quality Assurance representative in cross-functional discussions (in registration topics) with Production, Quality Control, Validation, and Regulatory Affairs, supporting risk-based decision making.
 - Lead the execution of Product Quality Reviews (PQR), including trend analysis and risk-based evaluation, to support continuous improvement and regulatory compliance.
 - Provide scientific and statistical justification for quality-related activities, particularly in process validation, data evaluation, and deviation management.
 - Ensure the implementation and sustainability of Data Integrity principles (ALCOA++) across all quality documentation and records, and promote data integrity as an integral part of the company's quality culture.
 - Support and participate in internal audits and regulatory inspections, including document preparation, audit accompaniment, and follow-up of findings (CAPA).
 - Conduct external audits of suppliers and partners, including domestic onsite audits and international remote audits.
 - Initiate and coordinate quality document digitalization, including the integration of data integrity principles into electronic systems.
 - Serve as a core team member in the development and implementation of Electronic Batch Record (EBR) and internally developed Electronic Quality Management System (e-QMS).
 - Support regulatory registration dossiers and contribute to the preparation of the Pharmaceutical Site Master File (SMF) for PT Interbat.
 - Manage and evaluate monthly temperature and humidity monitoring data for all storage and warehouse areas, including trend analysis.
 - Perform semi-annual environmental trend reviews of sterile manufacturing areas, including temperature, humidity, differential pressure, and microbiological data, as part of ongoing quality control and risk assessment.
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QA Staff

PT. Interbat | Indonesia

September 2023 – January 2024

- Prepare and compile Product Quality Review (PQR) documents for various pharmaceutical dosage forms in accordance with GMP/CPOB requirements.
 - Review and verify process validation documentation to ensure compliance with approved protocols and regulatory expectations.
 - Updating approved supplier and vendor lists for all materials and products used in manufacturing.
 - Support the organization and documentation of change control activities, including coordination with cross-functional teams.
 - Monitor and update applicable pharmaceutical regulations to ensure continuous compliance with current regulatory requirements.
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Quality Control (Apoteker Penanggung Jawab)

PT. Gra Parianom Sejati | Indonesia

July 2022 – July 2023

- Led the establishment of a new pharmaceutical manufacturing facility in compliance with GMP/CPOB 2018 requirements.
 - Developed and implemented company quality systems, standard operating procedures, and working instructions.
 - Oversaw Quality Control operations and ensured alignment with regulatory and internal quality standards.
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Stability & Cleaning Validation Supervisor

PT. Guardian Pharmatama | Indonesia

July 2021 – July 2022

- Supervised stability study programs for more than 150 pharmaceutical products to meet regulatory and product safety requirements.
- Managed cleaning validation activities, including protocol preparation, execution oversight, and report review.
- Coordinated analytical testing and ensured data accuracy, compliance, and traceability.
- Supported capacity planning and analytical workload management within the laboratory environment.