Shaik Azmath

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| linkedin.com/in/shaikazmath | Portfolio

Summary

Dedicated QC Chemist with over 2+ years of experience in pharmaceutical quality control, skilled in analytical techniques including HPLC, UV-Vis, and IR spectroscopy. Proficient in GMP documentation, regulatory compliance, and maintaining instrument calibration. Proven ability to ensure product quality and safety through method validation, stability testing, and precise chemical analysis. Seeking to contribute technical expertise and a compliance-driven mindset to a quality-focused organization.

Experience

Quality Control Chemist Hyderabad, TS

KP Labs *(Pharmaceutical Company)* *Nov 2022-Feb 2025*

- Conducted routine testing and analysis of raw materials, in-process samples, and finished products to ensure compliance with quality standards and regulatory requirements.
- Utilized advanced techniques, including HPLC, UV-Vis spectroscopy, and chromatography, for precise analysis of pharmaceutical and chemical compounds.
- Managed and maintained laboratory instruments and equipment, ensuring proper calibration and adherence to Good Laboratory Practices (GLP) and Standard Operating Procedures (SOPs).
- Assisted in the preparation of quality control documentation, including test reports, batch records, and certificates of analysis, ensuring accuracy and compliance with internal and regulatory standards.
- Collaborated with cross-functional teams to investigate and resolve quality issues, providing technical expertise in root cause analysis and corrective actions.

Skills

- **Analytical Techniques:** HPLC, UV-Visible Spectroscopy, IR Spectroscopy, Mass spectrometry, Basic NMR (Nuclear Magnetic Resonance), Wet Chemistry (classical titrations, Karl Fischer, pH, LOD), Dissolution Testing.
- **Quality Systems:** GMP, GLP, SOP Documentation, Method Validation, Instrument Calibration, Stability Testing, Raw Material and Finished Product Testing.
- **Quality Investigations & Instrument Support:** OOS/OOT Handling, Deviation, CAPA, Maintenance, Troubleshooting.
- **Regulatory Affairs & Compliance:** Knowledge of Regulatory Submission (CTD/eCTD format), Drug Approval Process, Compilation of Dossiers and Product Registration Documents, ICH Guidelines (Q1–Q10).
- **Soft Skills:** Attention to Detail, Problem Solving, Team Collaboration, Time Management.

Education

Osmania University Hyderabad, TS

Master of Science - (Organic Chemistry) *Graduation Date: Sep 2023*

Telangana University Nizamabad, TS

Bachelor of Science - MPC (Maths, Physics, Chemistry) *Graduation Date: Aug 2021*

Certifications & Training

Drug Regulatory Affairs | *S.G Pharma* | Oct 2023-Nov 2023

- Learned preparation of regulatory submission documents (CTD/eCTD) and dossier compilation.
- Gained knowledge of global drug approval processes and ICH quality guidelines (Q1–Q10).
- Strengthened skills in ensuring product quality and compliance with regulatory standards.

HPLC Training | *Udemy |* *Mar 2025-April 2025*

- Acquired foundational knowledge of HPLC instrumentation, including setup and calibration.
- Developed skills in method development, sample preparation, and analysis for various chemical and pharmaceutical samples.
- Gained proficiency in interpreting HPLC data and understanding various detectors and column types.

Languages known

English | Hindi | Telugu | Urdu