Anas Zaghal

Contact

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Languages

Arabic – Mother
English – Very good

Summary

R&D/Manager/Principal Scientist specializing in Analytical Research Laboratories. 13 years experienced with all stages of the development, validation, stability cycles and preparing all R&D filing products documentations as per the guidelines for the markets. Registration the projects in US, EU and MENA requirements.

Skill Highlights

- Project management
- Strong decision maker
- Complex problem solver
- Creative design
- Innovative
- Service-focused

Experience

Research Chemist - 09/2009 to 03/2015

Hikma Pharmaceutical, Amman

- As Analyst in formulation support analysis and stability for one-year, Analytical Research chemist method validation for two and half years, Analytical Research chemist method development for about two years in Analytical Research (Research and development department).

R&D/AR Project Team Leader - 04/2017 to 06/2022
Hikma Pharmaceutical, Amman, Portugal, Italy

- As R&D Project Team Leader in Analytical Research Laboratories (Research and development department).

R&D/AR Project Team Leader - 04/2017 to 06/2022
Hikma Pharmaceutical, Amman, Portugal, Italy

- As R&D Project Team Leader in Analytical Research Laboratories (Research and development department).

R&D/Manager/Principal Scientist - 06/2023
Hikma Pharmaceutical, Amman

- As Principal Scientist in (Research and development department).

Education

Bachelor of Science: **Chemical Technology** – 2009
Tafila Technical University

Certifications

- I-Communicate, what is your problem? and Business Etiquette Courses – Hikma Academy in Amman.
- Compendial HPLC Practices – USP Education in Amman.
- English Courses at ESL Center (The American ESL Center).
- English Courses at Berlitz Center.
- Advanced Chromeleon– Thermo Scientific in Germany.
- ICDL Certificate.
- ISO lectures at Hikma Pharmaceutical.
- Good Manufacturing Practice (GMP) and Good Laboratory Practice (GLP).

Responsibilities

- ARI Drug master file evaluation with United states pharmacopeia and European pharmacopeia monographs.
- Creating API specification/certificate of analysis and evaluating the chemistry, chromatography, and microbiology tests per USP, Ph. Eur and ICH.
- Creating drug product release/shelf-life specification/certificate of analysis and evaluating the chemistry, chromatography, and microbiology tests per USP, Ph. Eur and ICH.
- Creating API/drug product release/shelf-life justification of specification reports.
- Evaluating the API/drug tests that need method validation as per the guidelines (USP, Ph. Eur and ICH).
- Troubleshooting and developing the tests that not worked.
- Planning and manage the products with the timeline.
- Creating/reviewing method validation/verification/transfer protocols, report while ensuring compliance to SOP, s, cGMP, and GLP.
- Following up on submission batches analysis as Assay of active & inactive ingredients, Related substances, Cleaning Method, Diluent study Procedure, photostability study procedure, Standardization for Raw Material and Physical Characteristics.
- Following up and reviewing the preventive maintenance, qualifications, and calibration for chemistry equipments and chromatography.
- Training the teams on any updated SOP's, GMP or GLP.
- Following up on the updated USP and Ph. Eur monographs.
- Following up on the updated USP, Ph. Eur. and ICH guidelines.
- Evaluated the elemental impurities, residual solvents, Nitrosamines and extractable and leachable with risk assessments reports as per the ICH.
- Reviewing/approving API/Drug product certificate of analysis.
- Preparing and approving the drug product safety risk assessment elemental impurities, Nitrosamine, residual solvents, etc.
- Preparing and approving the analytical part in the product file before the submission.