ABDALLA EMADALDEEN ALOLABI

Technical Affairs Coordinator at CityPharm Pharmaceutical Industry

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Email Address: oulabi.abdullah@gmail.com
Location: Sudan – Khartoum, Al-Amarat st.31

Education: Bachelor of chemistry & Higher diploma, Faculty of Science, Damascus university
Experience: 14 years

PERSONAL SUMMARY

Creative professional chemist who has been consistently praised as cooperative by my subordinate, management and colleagues.

Over the course of my **fourteen years** of experience career, I've developed a skill set directly relevant to the pharmaceutical industry roles you are hiring for, including Risk management system, Formulation, Time management, Complex Problem Solving, ISO standard, Understanding of cross-cultural issues, Cost accounting, Validation and Ms Office programs.

My experience has spread over a broad spectrum of pharmaceutical industry in the veterinary and human fields for a variety of pharmaceutical dosage forms such as solid, liquid, semi-solid.

Overall, I have consistently developed quality management system abilities in every aspect of my technical coordinator role in addition to Contribution to improve productivity in many production areas in CityPharm pharmaceutical factory, I aspire to provide exceptional and valuable addition to your organization and I invite you to review my achievements in details.

PERSONAL INFORMATION

Birth date: 12 January 1983 (Age: 37)
Nationality: Syria, Sudan
Residence Country: Sudan – Khartoum
Marital status: Married
Driving license issued from: Syria, Sudan

PREFERRED JOB

Job title (s):, Research and Development, Production, Quality Assurance, Supply chain management, Quality Control
Preferred industry: Pharmaceutical, Cosmeceutical

WORK EXPERIENCE

Total Years of Experience: 14 years, 3 months

Coordinator for Technical Affairs at July 2017 - Present

CityPharm Pharmaceutical Industry

Sudan - Khartoum Bahri

- Coordination between technical departments among itself and between technical department and upper management.
- Provide technical assessments for supply chain changes, highlight technical risks, and recommend the needs to support the change.
- Coordination between production and marketing heads to get delivery on time.
- Identify opportunities to develop technical capabilities within Operations sites, and Provide training to meet agreed capability targets.
- Responsible for collaboration and communication with various teams as needed
- Conduct interviews, hire, and train new technical staff as necessary
- Provide guidance and insight to upper management & report progress, including any changes made to plans and production
- Contribute to product design and establishment of requirements
- Coordination with all departments to deliver products consistently, on time and on budget
- Studying the technical problems that obstacle the work and trying to solve it in

  Coordination with all company's head departments.
- following up implementation of monthly and annually plan of the technical departments previously approved in advance
- Performing periodic inspections to the company's technical departments with the head departments to verify the work performance.
- Ensuring that the quality system is implemented and followed up at all times.
- Reviewing reports of the annual review of final products in coordination with the technical committee members.
- Contribute to the preparation of technically complex responses to regulatory questions.
- Reporting to the upper management on the performance of the adopted management systems and any need for improvement
- participate in establishing quality risk management system.
-Responsible for meeting production requirements, delivering quality products to customers in a timely fashion and cost-efficient manner.
-participate in Responsibility for driving operational excellence and flawless execution in the areas of cost reduction, discard reduction, annual productivity improvements, inventory turnover, on time delivery and lead time reduction,
-Coordinates interdepartmental activities to ensure the arrival materials within scheduled frame time.
-Monitoring raw material and finished product inventory levels to maintain proper replenishment plans.

WORK EXPERIENCE

Research and development supervisor April 2017 to July 2017

CityPharm Pharmaceutical Industry

Sudan - Khartoum Bahri

- Performing analytical method development, validation and project support.
- Identifying, offering, and implementing contingency plans to prevent and overcome issues and perform method /process troubleshooting.
-Maintain stability programs in compliance with regulatory requirements
- Initiate, review, update, and/or complete GMP documentation as need
-Perform laboratory and manufacturing audits as required (cGMP, safety, housekeeping, etc.). Make necessary corrections as needed.
-Execute cleaning validation.
- Participate in the training of technical staff including QC, QA, R&D, production, maintenance, warehouse, (Within the Continuing Training Program) In cooperation with other colleagues.
-Participate in functions involving teams, which impact production, increase efficiency, solve problems, generate cost savings and improve quality.

- Participate in improving/developing formulas.

WORK EXPERIENCE

Production Manager May 2015 to January 2017

TechnoPharm Company for Pharmaceutical Industry (Veterinary).

Syria – Dara'a

-Responsible to ensure production output is on time and in full

-Responsible to monitor and ensure the flow of raw materials to finished product is being accomplished in the most efficient manner.

- Responsible for the overall organization and cleanliness of the production area.

- Troubleshooting any issues arising during the manufacturing process.

--Ensures that the operation during the shifts are meeting to the objectives for quality and productivity .

-Ensures that department complies with all safety rules and procedures, as well as good housekeeping within the plant, in accordance with SOPs, Work Instructions and GMPs.

-Responsible to all production area staff including motivating, disciplining, training and maintaining appropriate documentation

-Work Closely with Maintenance and Engineering to ensure proper functioning of production equipment and to minimize downtime.

-Follow up the technical aspects of the expansion project of the factory to ensure that everything meets to cGMP requirements (layout, HVAC, machine urs, etc...).

WORK EXPERIENCE

R&D specialist June 2014 to April 2015

TechnoPharm Company for Pharmaceutical Industry (Veterinary).

Syria – Dara'a

- Development of new products.

- Managing formulation projects from laboratory to full-scale manufacture.

- Troubleshooting any issues arising during the manufacturing process.

- Analysis of current/future markets for development in participation with marketing team.

- Competitor product analysis.

- Day to day follow-up of the problems occurring in the production departments and ensure that robust and effective solutions are found.

- Source, test and approve raw materials for new and existing products.

- Liaising with suppliers as required, to evaluating the new materials to be used.

- Work to develop and improve manufacturing process.

- Designing of most appropriate, economical and feasible formula for given target product

- Assisting when required with day to day production duties.

- provide technical guidance to the technical staff as needed.

- Develop and review source documents for regulatory submissions.

- Participates in internal audits.

WORK EXPERIENCE

Quality Control Supervisor at April 2012 to April 2014

Alfares Pharmaceutical Company

Syria - Damascus

- Supervise quality laboratory operations and personnel

-Assist with resolution of customer complaints and questions regarding product quality.

-participate in Maintain the plant Quality Management System in compliance with outside certifications and internal quality standards.

-Support operations through in-process and product conformance testing.

-Participate in troubleshooting, technology transfers, and process improvement activities.

- Oversee activities in the production departments which involve Quality and QC Inspectors

-Create weekly and weekend QC schedules limiting overtime as much as possible.

-Review quality control standards, specifications, and procedures, and evaluate effectiveness of quality control program

- Supervise the Validation Activities.

-Provide training to the technical staff regarding new SOPs and GMPs.

- Assist in investigation of nonconformance and customer complaints

-Author or amend stability protocols, and author interim and final stability reports.

-Coordinate with Supply Chain to ensure availability of general lab supplies and reagents

-Coordinate the stability testing schedule, supervise QC analysts in the writing of stability related quality system records.

- Troubleshooting of chromatography and other related issues.

- Performing laboratory investigations for any Out-of-Trend, Out-of-Specification or Atypical results to ensure that the root cause is identified and remedial action is taken.

- Evaluation of the measurement uncertainty as well as use statistical techniques for analysis of data

- Participation in the annual program of Proficiency Testing Study (PTS) organized by the European Directorate for the Quality of Medicines (EDQM).

- Implementing the the administrative and technical requirements of ISO 17025:2005.

- Ensuring that all working copies of documents are stored appropriately and that no unauthorized changes to the document are made after validation in the laboratory.

- Developed new analytical techniques for the analysis of drugs and improvement of analysis procedures.

- participate in developing new products / renovate existing formulas as deemed necessary in collaboration with formulation Department.

Makes decisions concerning quality in the absence of the QC Manager

WORK EXPERIENCE

Research & Development Chemist at February 2011 to January 2013

Alfares Pharmaceutical Company

Syria – Damascus

- Existing product evaluation and formulation modification as required.

- Designing of most appropriate, economical and feasible formula for given target product

-improve and troubleshooting problem submitted by customer from another factory (human-veterinary-cosmetic).

- Preparing samples of products for customer approval.

- Conduct analysis and experimentation on formula, for such purposes as product and process development and improvement of analytical methodologies.

-Develop standard operating procedures (SOPs).

-Conduct compatibility studies for the newly developed formulations of various dosage forms like tablets, capsules, suspensions, creams, ointments and topical solutions to study the characteristic changes, drug interaction between the API and the excipients.

Writes and reviews analytical methods, deviations, protocols, SOPs , BMR and any other technical documents.

-Analysis of finished products, in-process materials, raw materials, and cleaning verification and validation samples according to the assigned specifications, methods and protocols.

-Provides project support for new product development which requires a close working relationship directly with Customers as well as suppliers.

WORK EXPERIENCE

Quality Control Senior Analyst at August 2008 to January 2011

Alfares Pharmaceutical Company

Syria - Damascus

- Testing stability samples by following the testing protocols and validated methods.

- Development and validation of analytical methods for new and existing products.
- Preparation of validation protocols and reports.
- Routine testing of samples, products and raw materials according to approved procedures
- Performing audits, in-process checks and housekeeping duties.
- Implementing the administrative and technical requirements of ISO 17025:2005
- Participation in the annual program of Proficiency Testing Study (PTS) organized by the European Directorate for the Quality of Medicines (EDQM).
- Execute the reference material procedure to fulfilling the requirements of ISO 17034 for the preparation and handling of internal reference materials.
- checking the performance of water treatment units (waste water treatment units & purified water unit) in the factory.
- Documenting all results of analysis in the records and/or logbooks.
- Applying quality control mechanisms as per SOPs and managing control charts.
- Assuring the active implementation of the implemented management systems and ISO/IEC 17025:2005.
- Testing the primary and secondary packaging materials
- Checking physical specification of dosage forms
- Monitoring the manufacturing & packaging process as per QA procedures

WORK EXPERIENCE

Quality Control Chemist at April 2006 to July 2008

Alsham Company for detergents

Syria - Damascus

- Routine testing of products and raw materials.
-Sampling and analysis of raw materials, packing materials, in process and finished products according to approved QC/QA procedures.
-Implementation of QA & QC procedures
- Contribute to the review and preparation of the test / operational procedures.

WORK EXPERIENCE

QA/QC Analyst at August 2005 to April 2006

Alazmah Company for Paints industry

Syria - Damascus

- Sampling and analysis of raw materials, packing materials, in process and finished products according to approved procedures.
- testing new coloring material and Developing color ratio
-Implement the approved QA/QC procedures
- Contribution to Quality related Investigations. Investigate out of specifications and Recommend CAPA for final decision.
- Complaint Handling: Evaluate & investigate product complaints. Recommend CAPA.

EDUCATION

Bachelor's degree, Faculity of science – Applied Chemistry
Damascus University
Syria – Damascus
June 2006

Higher diploma, Faculty of science
Damascus University
Syria – Damascus
June 2008
Techniques and methods of teaching physics and chemistry

SKILLS

QUALITY CONTROL / Level: Expert
ISO 17025:2005 / Level: Expert
ISO 9001:2015 / Level: Expert
HPLC / Level: Expert
CALIBRATION / Level: Intermediate
ISO 14001:2004 / Level: Beginner
ADOBE PHOTOSHOP / Level: Beginner
Ms Office ExceL / Level: Expert
Ms Office Word / Level: Expert
Spss / Level: Beginner
Statistical analysis / Level: Intermediate
Pharmaceutical Formulation / Level: Intermediate
DOCUMENTATION / Level: Expert
Ms Office PowerPoint / Level: Intermediate
OHSAS 18001:2007 / Level: Intermediate
CTD / Level: Beginner
cGMP / Level: Intermediate
Good Storage Practices / Level: Intermediate
Good Laboratory Practices / Level: Intermediate
Validation / Level: Intermediate
Quality Assurance / Level: Expert
Stability Study for finish product / Level: Intermediate
Analytical Instruments / Level: Intermediate
Strategic Planning / Level: Intermediate
Change control / Level: Intermediate
Microbiology / Level: Beginner
bioequivalence / Level: Beginner
Risk Assessment / Level: Beginner
Iso Standards / Level: Intermediate
Analytical chemistry / Level: Intermediate
Complex Problem Solving / Level: Intermediate
Analytical Thinking / Level: Intermediate
Time Management / Level: Intermediate
Persistence / Level: Intermediate
inductive Reasoning / Level: Intermediate
Coordination / Level: Intermediate
Microsoft Visio. / Level: Beginner

LANGUAGES

Arabic / Level: Expert
English / Level: Intermediate

Training and Certifications

The 7 step process for tooling care and maintenance
Training Institute: Alfares Pharmaceutical Company **(Damascus)**
Date attended: November 2008 **(**18 hours**)**

Scale-up & Post approval changes guidelines (SUPAC)
Training Institute: Nara Tech **(Jordan)**
Date attended: June 2012 **(**16 hours**)**

Positive thinking & communication skills
Training Institute: Value Added center **(Damascus)**
Date attended: June 2011 **(**21 hours**)**

ISO specialist Skills
Training Institute: 7vision for training and Development **(Sudan)**
Date attended: January 2017 **(**20 hours**)**

Internal Auditor
Training Institute: government center for developing the productivity & management **(Damascus)**
Date attended: April 2011 **(**20 hours**)**
Duration: 20 hours

Implementation of cGmp
Training Institute: Arab quality makers & QMS **(Damascus)**
Date attended: August 2010 **(**10 hours**)**
Duration: 10 hours

Good Manufacturing Practices
Training Institute: Alfares Pharmaceutical company **(Damascus)**
Date attended: January 2010 **(**18 hours**)**

Internal Auditor ISO 9001:2008
Training Institute: National Company for consultancy **(Damascus)**
Date attended: November 2015 **(**10 hours**)**

Awareness ISO 9001:2008
Training Institute: National Company for consultancy **(Damascus)**
Date attended: November 2015 **(**10 hours**)**

Implementation of ISO 17025:2005
Training Institute: Syrian Scientific Society for Quality **(Damascus)**
Date attended: March 2010 **(**15 hours**)**

Quality management system according to ISO 9001:2015
Training Institute: Syrian Scientific Society for Quality **(Damascus)**
Date attended: May 2016 **(**12 hours**)**

Hobbies and interestes

Developing formulas for cosmetic and pharma products
By virtue of my knowledge of the properties of most of the materials used in the pharmaceutical & cosmeceuticals, I have been able to develop many products in my home or in my work during the previous years and I feel pleasure when I contribute to such things.

Reading references and guidelines relevant to pharmaceutical industry, applied science

I currently own a huge collection of references and books related to the pharmaceutical and cosmetic industry in addition to ISO standard A large part of it has been read in whole or in part and is supported as a reference which can be consulted when making decisions are required

Memberships:

Organization: Syrian Scientific Society for Quality
Membership/Role: fellow
Member since: January 2009

Social Profiles:

Facebook: facebook.com/abdullah.oulabi
Linkedin: linkedin.com/in/abdullah-al-oulabi-89143092

Experience profile:

Kindly see attachments

References

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