KHALED AL-SARARI

Quality Assurance

PROFILE SUMMARY

I'm a Quality assurance. I'm a always looking for new challenges. Obtain a challenging position in a pharmaceutical manufacturing company (Sterile area Vials and Non sterile area cephalosporin and Non cephalosporin) CAPA, Deviation, Validation, Review BMR,BPR, SOP,Release of Batch, SAP, int/external audit which offers opportunities for advancement in career.

INFORMATION

Date of Birth

10/November/1983

Nationality

Yemeni

Place of Brith

Tabuk-KSA

ACHIEVEMENTS

EMPLOYEE Of MONTH AWARD

Tabuk pharmaceuticals company

Volunteering Work Certificate

Minister of Health

Quality Ambassador program Team

Saudi Standard,Metrology,Quality Organization (SASO)

CONTACTS WITH ME

Mobile: 0509927275

Email: dr.alsarari@gmail.com

Linkedin.com/in/khaled-alsarari

ACADEMIC BACKGROUND

Master Degree

Al-Isra University : Master Degree in pharmaceutical science

BSc

AL- Zaytoonah Private University

CAREER BACKGROUND

Tabuk pharmaceuticals company
2009-2011 Quality Assurance Officer
2014-2019 Quality Assurance Senior Officer

TRAINING COURSES

PHARMACEUTICALS

Pharmaceuticals Selling Skills Course : At Jordan pharmacist Association.
Courses Duration 100 Hours

CLINICAL PHARMA

Clinical Pharmacy Skills Course : At Jordan pharmacist Association.
Courses Duration 08 Hours

CLINICAL PHARMA

Clinical Pharmacy Practical Course : At Jordan pharmacist Association.
Courses Duration 72 Hours

INJECTION COURSES

Injection Course : At Jordan pharmacist Association.
Courses Duration 08 Hours

CLEANING VALIDATION

Cleaning Validation - Principle & Practices for The 21st century Century : At Oakes Group Global
Courses Duration 100 Hours

LEAN SIX SIGMA

Udemy online
Courses Duration 02 Hours

MANAGEMENT SYSTEMS

The role of ISO 9001 Quality management system saudi standard, Metrology,Quality org.(SASO)
Courses Duration 02 Hours

LEAN SIX SIGMA

Application to Measurement Control the Quality operational performance At: Saudi Society for Quality
Courses Duration 02 Hours

PERSONAL SKILLS

- **Microsoft office Word/ExcelPowerPoint**
- **Web Searching My Challenging Data**
- **Communication skills &The Ability To work Effectively within A Team**
- **Strong Interperson &Hardworking**
- **Good presentation & Communication ExcellentOrganizational &Filing skills**
- **Quality-Oriented With very Good Analytical Skills**
- **Cooperate,Excellent Ability To learn**
- **Effective in Delivering Ideas For All Levels, Solve problems**
- **Self-Motivated**

LANGUAGES

- **ARABIC (NATIVE)**
- **ENGLISH (GOOD) Writing & Speaking**

DRIVING LICENSE

WORK EXPERIENCE

- Review of all master documents related to manufacturing batch and packaging.
- Approval and final Release of batch as per regulations requirement .
- Follow up the preparation & aseptic filling process to insure it's adherence to the SOP & the validated procedure's.
- Support & implement in media fill stimulation test of vials
- Prepare all paper work & filling relevant to the job
- Following up all manufacturing steps (receiving & weighing of raw materials, preparation, filtration, filters integrity test, autoclave loading & de-loading checking, on all charts on spot, filling, leak test
- Coordinating with QC for analytical report review material approval.
- Review of post and per execution deviation, change Control, OOS, CAPA and their assessment
- Ensure that all deviations & problems are covered & solved according to the quality system & GMP requirements
- Check & follow the documentation within manufacturing, filling, packaging batch records.
- Assur the integrity of the manufacturing & packaging products.
- Follow up the in process quality test such as moisture analysis of powder & granules, disintegration, friability, weight variation, thickness & hardness of Tablets.
- Follow up the sampling of stability pack's, filled primary packaging components & semi-finished bulk for stability studies, microbial contamination & quality control testing respectively of validation batches.
- Follow up the leakage test for primary packaging materials such as bottles & blisters.
- Follow up the sampling of semi-finished bulk before packaging for quality control testing.
- Follow up the sampling of filled primary packaging components such as bottles, blisters, vials....etc. for microbiological testing
- Follow up the sampling of retention/ reference of finished of all pharmaceuticals products produced by the industry.
- Follow up the production area to ensure the implementation of GMP & to perform IPC Test.
- Full participation in processing of solid, semi-solid & liquid batches (preparation, compression, filling & packaging.
- Sampling of in process, finished product, Control sample, cleaning validation.
- Follow up the internal/External audit and any inspection notes & requirements.
- Ensure that cleaning activities in the production areas & stores carried out according to approved procedures & cleaning validation studies.
- Follow up the implementation & closing with related documentation of corrective & preventive action system inputs.
- Management of SAP production related activities.