106284-CHE-5OQ-E-2010
Senior QC Specialist

Holds a B. Sc. in Chemistry and has over 7 years hands-on experience working in QC field.

PERSONAL DATA

Nationality	:	Egyptian
Gender	:	Male
Marital Status	:	Single
Residence	:	Assiut

EDUCATION

: B. Sc. in Chemistry, Assiut University, 2010

LANGUAGES

Arabic	:	Native Language
English	:	Very Good
German	:	Good

COMPUTER SKILLS

: Windows, MS Office, Internet

TRAINING COURSES AND CERTIFICATIONS

: Successfully completed ASEC Cement Manufacturing Course (ACMC) in ASEC Academy for training (Nov. 2018).
: Cement chemistry course & X-Ray spectrometry course in Misr Cement Company (Qena) - ASEC (Mar. 2017).
: Course use of radiation sources / materials and radiation protection against lionizing radiation organized by the Egyptian Atomic Energy Authority (Apr. 2016).
: Successfully completed ASEC Basic Cement Manufacturing Course (ABCMC) in ASEC Academy for training (Dec. 2015).
: Occupational health and safety course in National Institute of occupational safety & health (Apr. 2014).
: Occupational health and safety assessment series (OHSAS 18001) in Egyptian Syndicate of scientific professions (Mar. 2013).
: European good manufacture practice (Euro. GMP) in T3A Pharmaceutical Industrial Factory (Sep. 2011).
: Successfully passed all modules required of the granting of the International Computer Driving License (ICDL) (Apr. 2011).
: Computer's maintenance (software & hardware) course in Smart Group for training (Oct. 2010).

: Training course in Organization of the new and renewable energy (Aug. 2009).
: Training course in Assiut Oil Refining Company (ASORC) (Aug. 2008).
: English and computer maintenance course in Assiut University (Jul. 2008).

CHRONOLOGICAL EXPERIENCE RECORD

Dates : From Mar. 2015 till now
Employer : Arab Swiss Engineering Company (ASEC)
Project : Assiut Project
Job title : Senior Quality Control Specialist
Job Description :

- Ensure that all laboratory are maintained to a high level of safety and risk assessments and Operating procedures.
- Maintain product quality to the market and standards requirements.
- Development, implement and maintain the quality control department procedures to manage quality control of raw materials, raw mix, clinker and cement.
- Responsible for the assessment of raw mix composition and pilling formation according to the type of Mix (SRC OR OPC) using mix design.
- Responsible for dry cement line (raw mix, raw mills and cement mills) by using gamma matrix and x-ray fluorescence spectrometry (ARL 9900 series) to monitor their complete chemical analysis.
- Responsible for the quality control of plant quarries (limestone and clay).
- Check up & standardizations for the x-ray apparatus by the chemical analysis of the samples and compare the result with the x-ray result.
- Create all the database for laboratory (document, reports, and maintenance documents).
- Responsible for all of chemical analysis for cement and raw materials.
- Using LECO CS230SH to control sulphur and carbonate content in raw material and cement.
- Provide training for the new staff.
- Suggesting new solutions for quality problems.
- Monitor the physical and chemical test results of raw materials, in process products and finished cement for conformance to regulatory specifications.
- Auditing materials provided by suppliers to assure quality before incorporation into the product.
- Coordinate between lab results and production teams to ensure that product meet required the national and international standard specifications.
- Supervision the different types of lab. Testing (x-ray, chemical and physical testing).
- Creating all analytical programs on OXSAS software
- Achieved both daily quality target and quantity of production target.
- Performing inspection and calibration monthly of x-ray instrument as well as anther lab instrumentation and maintaining its condition.

Dates : From Aug. 2011 till Mar. 2015
Employer : T3A Pharmaceutical Industrial Factory

Job title : Senior Quality Assurance Specialist

Job Description :

- Follow up the new requirements of ISO 9001.
- Responsible for preparing the annual product review as a half yearly basis.
- Follow up and perform all the required documents for release of finished products.
- Follow up and assure that the production processes are implemented according to GMP, ISO 9001:2000, ISO 22000:2005 in pharma and ceph. Building and sent a daily report to QA manager.
- Improve the QA systems implementation to reduce the market complaint.
- Issuance and conducting the self inspection plan regarding to ISO 9001:2000 requirements.
- Responsible for assurance of that all quality management systems comply with ISO 9001:2000 standards.
- Reviewing for all processing and packaging batch records.
- Ensuring the release of finished products at its predetermined time schedules.
- Controlling the whole types of returned products till its final destination.
- Controlling the rejected / destructed of materials / products till their final destination.
- Responsible for repackaging procedure.