

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 ionizing 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 another 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.