Miguel A. Gonzalez Quero

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Objective

- To obtain a challenging position in a high-quality engineering environment where my resourceful experience and academic skills will add value to organizational operations.

Education

POST-GRADE – SPECIALIZATION IN BUSINESS ADMINISTRATION, QUALITY AND PRODUCTIVITY | DECEMBER 2012 | UNIVERSIDAD TECNOLOGICA DEL CENTRO - VALENCIA

- Specialization Thesis: entitled "Improvement of solids packaging processes in a pharmaceutical plant by applying assessment tools of productivity and efficiency".

BACHELOR OF SCIENCES IN CHEMICAL ENGINEERING | OCTOBER 2007 | UNIVERSIDAD DE CARABOBO - VALENCIA

- Merit Scholar II-2002 and I-2004 (Faculty of Engineering, Universidad de Carabobo).
- Credentials evaluated and accredited on July 2015 by Josef Silny & Associates, Inc. USA.

Core Qualifications

- Comprehensive knowledge of methods and standards of chemical/process manufacturing unit.
- Strong knowledge of chemistry fundamentals.
- Able to identify and implement strategies that reduce costs, increase revenue, improve business processes and drive profitable growth.
- High experience on managing teams, supporting staff and fostering safety along the organization.
- Knowledge of cGMP in the pharmaceutical industry, quality regulations (international), and lean manufacturing.
- High proficiency in Microsoft Office (Word, Excel, PowerPoint, Publisher, Outlook), MS Dynamics, and Lotus Notes.
- Excellent conflict management, problem solving, and time management skills.
- Languages: Spanish, English and Portuguese (Excellent reading, writing and speaking skills). Credentials available.
- Enjoy cross-cultural work environments and willing to relocate according to business needs.

Professional Experience

PACKAGING SUPERVISOR | SOLARA LABS - MIAMI, FL | JULY 2017 TO PRESENT

- Responsible of an area with at least 6 employees.
- Team work and focus on results in a weekly planning.
- Lead deviation and issues investigations and solutions to improve packaging processes.
- Monitor of daily productivity and provide daily reports to management team.

ADVISOR ENGINEER | FREELANCE – VALENCIA, VENEZUELA | SEPTEMBER 2015 TO AUGUST 2017

- Assessment of manufacturing processes as contractor in manufacturing sites.
- Time and motion studies.
- Re-engineering of tasks and activities in manufacturing plants.
- Business strategies in internal and relationship marketing.
- Audits and Quality Trainings and Lecturer.

- Documentation Standard Plans.

QUALITY SYSTEM MANAGER | TEVA PHARMACEUTICAL, INC. – GUACARA, VENEZUELA | AUGUST 2014 – JUNE 2015

- Managed a new manufacturing facility in Venezuela and selected plant and equipment layout technology in coordination with management.
- Established lab procedures; ensured compliance with environmental regulations, and emission standards.
- Responsible for ensuring healthy, safety, and environmental concerns for all projects.
- Conducted manufacturing deviation investigations.
- Prepared, evaluated, and implemented modifications to improve productivity and quality.
- Provided quality and technical support to multidisciplinary teams.

VALIDATION MANAGER | TEVA PHARMACEUTICAL, INC. – GUACARA, VENEZUELA | AUGUST 2013 – AUGUST 2014

- Led area to maintain the arrangement of activities related to validation of manufacturing and cleaning, qualification of machines, critical systems (HVAC, PW, CAS), facilities and computerized system.
- Responsible for validation master plan and risk assessments.
- Participated in multiple projects for process improvement.

LIQUID-LOZENGES AREA SUPERVISOR (PRODUCTION) | TEVA PHARMACEUTICAL, INC. – GUACARA, VENEZUELA | SEPTEMBER 2011– AUGUST 2013

- Shared a team jointly with a other supervisor to achieve integration of two different teams (34 people) as plant strategy.
- Led project to improve instructions and trainings for processes of mixing and primary packaging of liquids, ointments, suppositories, lozenges and cleaning instructions.
- Conducted deviation investigations and solutions to mitigate and succeed in manufacturing and packaging processes.
- Contributed to plant equipment layout design.

SOLID AREA SUPERVISOR (PRODUCTION) | TEVA PHARMACEUTICAL, INC. – GUACARA, VENEZUELA | MARCH 2010 – SEPTEMBER 2011

- Responsible for a team of 15 employees of the OSD packaging area.
- Monitored daily productivity and provided reports to management team.
- Provided safety, technical and operational guidelines to staff.
- Evaluated business requirements for maintaining product security and resolved any ongoing issues.

VALIDATION SPECIALIST (QUALITY) | TEVA PHARMACEUTICAL, INC. – GUACARA, VENEZUELA | MARCH 2007 – MARCH 2010

- Responsible for validation of protocols of manufacturing process, of production machines, critical systems (HVAC, PW, CAS) and facilities in order to achieve compliance on GMP standards and procedure.
- Member of internal audits team and member of staff support for "Quality Excellence Program" with the intention of raising awareness, information and capacity to all personnel in recognize flaws in operations.
- Achieved excellent performance that resulted in considerable savings and deviation identifications on operations.

Additional Information

- Venezuelan Citizen/Married and without children.
- Amateur swimmer open water and long course pool.