**PharmEng Technology** is currently seeking a **Process Validation Specialist f**or a project in **Maryland, US. T**he position is available immediately conditional of a successful interview

**Job Description:**

* Provide support to Manufacturing and Packaging Operations by identifying process improvement opportunities and developing proposals for problem solving, improvement or optimization
* Support process improvement projects that have the objectives of developing more cost efficient and quality enhanced processes
* Evaluate data generated through studies and develop/review production standards and Manufacturing/Packaging process to assure utilization of best process
* Optimize process output and throughput, equipment utilization, equipment downtime, waste reduction/yield improvements
* Provide technical support to the production areas in evaluating process deficiencies, process changes, modifications and equipment failure
* Define, analyze, review and evaluate problems for prompt solutions. Determine, adapt, and modify methods and standards to control all aspects of assigned projects
* Perform investigation and write deviation reports following the established timeline. Route and follow up the deviation reports through the evaluation, review and approval
* Provide recommendations and assist in CAPA implementation efforts
* Support process transfers, contract manufacturing, and other projects in an assigned engineering specialization for the development, manufacture, installation, operation and maintenance of products, production processes and/or equipment, packaging and other related activities
* Write and revise manufacturing and packaging batch records and Standard Operating Procedures
* All other duties as require

**Qualifications:**

* University Engineering degree (i.e. Mechanical, Chemical Process Engineering) with minimum one (1) year relevant experience is preferred.
* Proficient in Microsoft Office (Word, Excel, PowerPoint, Outlook)
* Proficient in Trackwise and MasterControl
* Experience in Active Pharmaceutical Ingredient (API) Manufacturing
* Experience working with potent compounds
* Applied knowledge of pharmaceutical GMPs, FDA guidelines, and industry standards
* Ability to apply GMP to company specific processes and products
* Excellent professional documentation skills
* Ability to objectively, accurately, and thoroughly convey complex issues in writing
* Ability to produce a large volume of written materials independently
* Ability to interact with other departments effectively
* Ability to review work performed by other personnel, communicate problems and deficiencies, elicit corrections, and enforce company policy and procedures as they relate to manufacturing requirements
* Ability to handle confidential company data, projects, information, etc.

If you are interested in this position, please submit your resume to [**careers@pharmeng.com**](mailto:careers@pharmeng.com)**.**

Thank you for your consideration and application! We review all resumes and submissions, however, due to the sheer volume of requests that we receive, only successful candidates will be contacted.