



PharmEng Company Profile

PharmEng International Inc., (trading symbol TSXV: PII), is a full service consulting firm that serves the pharmaceutical, medical devices and biotechnology industries in North America and internationally. The company provides industry recognized services on consulting from drug discovery to commercialization, contract manufacturing and testing, as well as cGMP training through the PharmEng Learning Institute.

PharmEng Innovations (a division of PharmEng International Inc.) provides comprehensive pharmaceutical research & development services including formulation and analytical development, clinical material manufacturing and scale-up services for the global markets.

Validation Scientist

PRINCIPAL DUTIES:

- Instrument qualifications (IQ/OQ/PQ), repair and maintenance of laboratory instrumentations
- Initiate, plan and implement work schedule according to project timelines
- Participate in internal/external audits. Present and discuss laboratory instrument quality systems with external clients or auditors.
- Identify and implement new instrumentation services to provide total solutions to clients
- Produce, review and approve technical document according to cGMP

SKILLS/EXPERIENCE REQUIRED:

- B.Sc., M.Sc. with 3+ years of pharmaceutical laboratory instrument qualification experience
- Extensive instrumentation validation experience for HPLC, GC, Dissolution and Spectroscopy (e.g., Agilent, Waters) in the pharmaceutical industry.
- Exceptional cGMP, GCP, GLP and FDA compliance knowledge.
- Superior interpersonal and communication skills (both oral and written), leadership ability and organizational skills.

Please forward your resume in confidence, to:

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ONLY CANDIDATES SELECTED FOR INTERVIEW WILL BE CONTACTED