

Projet Leader, Regulatory Development

Status: Temporary**Beginning of contract: March 2010****End of the contract: June 2011****Key responsibilities:**

Accountable for the attainment of development products and marketed products objectives.

Main responsibilities :

- To provide regulatory strategic input to local and global teams for product in development and marketed products throughout their life-cycle
- To develop and execute filing and approval strategies
- To organize and coordinate meetings at Health Canada
- To assure the quality and timeliness of regulatory submissions and approvals
- To ensure maintenance of compliance for marketed products

Characteristics of the ideal candidate:

- Master's degree in pharmacy or any health-related field
- Minimum of 2 years previous experience in the preparation of regulatory Submissions
- Good understanding of the current Health Canada regulations and guidelines regarding pre-clinical, clinical or chemistry and manufacturing issues.
Action-oriented, dealing with ambiguity, command skills, negotiating, priority setting, process management, standing alone, strategic agility
- Bilingual (French & English)
- Computer knowledge: Outlook (Microsoft), PowerPoint, Word, Excel, WordPerfect, MS Project and Adobe

If this opportunity interests you, please send your resume to:
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