

This comprehensive postgraduate diploma program has been developed to meet the skill requirements for effectively performing duties in clinical research studies and for those who want to upgrade existing skills or obtain new skills. Each area of the study will be covered in an in-depth manner for healthcare, pharmaceutical, cannabis and related sectors.

PROGRAM HIGHLIGHTS:

- Hands-on Training
- Optional Job Assistance for co-op, internship and full-time opportunities within pharmaceutical, cannabis and biopharmaceutical and related sectors



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ADMISSION REQUIREMENTS:

- Any Degree or Diploma/ Equivalent
- **In addition,** international students with the exception of students from English speaking countries require IELTS score 6 or equivalent.

PROGRAM FEE:

Tuition Fee: \$9,800, Books/Material Fee: 1,500, International Student Fee: \$2,450 (if applicable).

PROGRAM OUTLINE:

- Clinical Science
- Introduction to Clinical Studies (GCP)
- Regulations and Standards in Clinical Research
- Clinical Research in Medical Device
- Ethics and human protection
- Development of Clinical Trial Protocols
- Clinical Trial Monitoring
- Quality Assurance
- Development of SOP
- Preparing and Hosting Audits and Inspections

- Good Documentation Practices
- Clinical Data Management and Biostatistics
- Principles and Practices in Pharmacovigilance
- Introduction to Regulatory Affairs
- Professional Regulatory Affairs
- Electronic Records and Computer Validation
- Project Management I
- Project Management II
- Privacy

JOB PROSPECTS IN CRO, HOSPITALS AND ACADEMIC RESEARCH SECTORS:

- Clinical Research Assistants
- Clinical Research Coordinators
- Clinical Research Associates

- Project Managers
- Principal Investigators

GET IN TOUCH - Mississauga Campus
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