

# POST-SECONDARY DIPLOMA PHARMACEUTICAL QUALITY CONTROL & QUALITY ASSURANCE



Inquiry: 416-412-7374

Call / Text: 647-998-7374

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Program Code: PHA.QC.QA.DIP
Program Duration: 26 Weeks of
Training with Optional Job
Assistance program

## Admission Requirements

 Students with Minimum of Grade 12 Diploma with required science and math courses / Equivalent

### **Tuition Fee**

#### Fee Breakdown

• Tuition Fee: \$9,900

Books/Material Fee: \$1,500

• Lab/ PPE Fee: \$1,100

Total Fee - \$12,500\*

\*Please note that the amount shown is in Canadian Dollars.

### Program Outline

#### **SEMESTER 1**

#### **GCCP01 - Good Manufacturing and Good Clinical Practices**

- Concepts and requirements necessary for compliance with Good Manufacturing Practice (GMP) and Good Clinical Practice (GCP).
- GMP controls required to manufacture drug products in Canada, the United States and in many parts of the world.
- Focus on real work industrial scenarios.

#### **ILTP02 - Introduction to Laboratory Testing & GLP**

- Compliance with Good Laboratory Practice (GLP) for pharmaceuticals and biopharmaceuticals.
- Address the most important laboratory technique.
- Exposure and experience with all major pharmaceutical instruments.
- GLP guidelines widely used in pharmaceutical and related sectors.
- Understanding of instrumentations, calibration and the application of various instruments.

#### STW006 - Advanced Scientific-Technical Writing

- Different types of technical writing.
- SOPs, Communication Methods And Issues, Validation Report Writing, Testing Methods, CofAs, CAPA, Change Control, Gap Analysis, OOT, OOS, Deviation.

#### **SEMESTER 2**

#### **HPL004 - High Performance Liquid Chromatography (HPLC)**

- HPLC/DAD fundamentals and functionality, sample preparations and system setting-up.
- Chemstation software, IQ, OQ and PQ, testing and analyzing data.
- Troubleshooting and investigations.

#### **HPD001 - HPLC Method Development and Method Validation**

- Method development and method validation using the HPLC / DAD.
- Major industrial regulations in terms of developing and validating testing methods.

#### **SEMESTER 3**

#### DIS003 - Dissolution and Dissolution Hands-on Training

- Introduction to dissolution current testing practices, testing requirements, critical components, sample prep techniques, media preparation, sample testing, data analysis, report preparation, limitations, IQ / OQ / PQ, calibration, troubleshooting, maintenance.
- Out of specification investigation.
- Dissolution techniques, applications, USP apparatus, dissolution testing, report writing, calibration, troubleshooting.

#### GCT005 - Gas Chromatography (GC)

- Gas chromatography history, concepts, carrier gas.
- Testing requirements, critical components, sample preparation techniques.
- Sample testing, data analysis, report preparation, limitations.
- GC-FID IQ / OQ / PQ, calibration, troubleshooting.
- Maintenance, and out-of-specification investigation.

#### **GCUV002 - GC Method Development and Method Validation**

- Method development and method validation using the GC / FID.
- Major industrial regulations, in terms of developing and validating testing methods.



# For more questions Visit Us at

#### Toronto Campus

9-5310 Finch Ave East
Toronto, ON
M1S 5E8

#### **Brampton Campus**

44 Queen St E, Brampton, ON L6V 1A2

#### Contact Us

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