CellCAN
Regenerative Medicine and Cell Therapy Network

The highly acclaimed Hands-on GMP Workshop is launching in a new **INTERACTIVE E-LEARNING FORMAT!**

This course is designed for people who are working, or plan to work in areas associated with the manufacturing of cell and gene therapies, or want to expand their knowledge on advancing cell and gene therapy projects towards the clinic.

At the end of this course, the participants will be able to:

- Identify the **essentials of GMP** requirements and the relevant Canadian and international regulations;
- Identify **cleanroom facility and gowning requirements** and **define aseptic workflow**;
- Identify the components of a recognized GMP training program;
- Describe how to plan a **validation strategy through Quality by Design and Risk Management**;
- Recall the main components of a **Quality Management Systems (QMS)**, including SOP writing and quality manual development, batch records, environmental monitoring and GMP training program;
- Explain basic manufacturing techniques for cell-based therapies;
- Identify the **quality control** steps and parameters required for **product release**.

MORE INFORMATION & REGISTRATION : www.cellcan.com/training

MORE THAN 12 HOURS OF INTERACTIVE TRAINING

LEARN AT YOUR OWN PACE AND IN THE COMFORT OF YOUR OWN HOME

TEST YOUR KNOWLEDGE THROUGH ASSESSMENTS AND QUIZZES

PASS THE EXAM AND OBTAIN YOUR CERTIFICATE OF COMPLETION

ADD TO YOUR HQP TRAINING FILE FOR REGULATORY COMPLIANCE

INTERACT WITH EXPERTS AND NETWORK WITH OTHER TRAINEES