1. Safe Work Practices - 1.01, 1.02, 1.03, 1.04, 1.05, 1.06, 1.07, 1.08, 1.09, 1.10, 1.11, 1.12, 1.14 2. Data and Specimen Collection and Handling - 2.01, 2.02, 2.03, 2.04, 2.05, 2.07,

2.08, 2.09 3. Specimen Preparation and Pre-analytical Processing - 3.02, 3.03, 3.03.01, 3.04, This course is linked to the following

3.04.01, 3.05, 3.05.01, 3.06, 3.06.01, 3.07, 3.07.01, 3.10, 3.10.01 categories and competencies from the

4. Equipment, Instruments and Reagents - 4.01, 4.02, 4.02.01, 4.04, 4.05, 4.06 5. Analysis - 5.01, 5.02, 5.03, 5.04, 5.10, 5.11, 5.12, 5.13

6. Recording and Reporting - 6.01, 6.02, 6.03, 6.05, 6.07, 6.08

7. Quality Management - 7.02, 7.03, 7.04, 7.06, 7.07, 7.08, 7.09, 7.10, 7.11, 7.12 8. Resource Management - 8.04

9. Communication and Interaction - 9.01, 9.03 10. Professional Practice - 10.06, 10.10

Canadian Society for Medical Laboratory Sciences (CSMLS) Clinical Genetics Competency Profile (2016):



Instructor Name: xxxxxxxxx Email: xxxxxxxxxx Phone: xxxxxxxxxxx Office Location: xxxxxxxx Office Hours: By Appointment Course Website: Blackboard

Textbooks Arsham, M.S., Barch, M.J., & Lawce, H.J. (2017). The AGT Cytogenetics Laboratory Manual (4th Edition). Hoboken: Wiley Blackwell. ISBN 9781119061229. Required. Ronald, C., Scherer, S. & Hamosh, A. (2023). Thompson & hompson Genetics and Genomics in Medicine (9th Edition). Philadelphia: Elsevier. ISBN 9780323547628:

McGowan-Jordan, J., Hastings, R.J. & Moore, S. (2020). ISCN 2020: An International System for Human Cytogenomic Nomenclature. Basel: Karger. ISBN: 978-3-318-06704-0. Required.

Learner Supplies Michener Branded Labcoat

Personal computer with minimum technology requirements as per

Pre/Corequisites Prerequisite Entry to Program

Course Start Date: September 5, 2023 Total Credits: 17 credits Total Hours: 218.4 There are 167.5 instructional hours in this course, including lectures, tutorials, and laboratory sessions. The remaining hours are for self-study and self-assessment of karyotype and nomenclature packages which are completed independently. Number of Weeks: 13 weeks

**Course Information** 

This course will provide the foundational theory and hands-on experience of laboratory techniques currently used in diagnostic Cytogenetics laboratories. The student will acquire the knowledge, skills and techniques required to process Cytogenetic specimens from receipt to chromosome analysis and interpretation. The course is comprised of Laboratory, Tutorial and Lecture components.

% PAT Title Week Due

2 Cell Culture, Equipment & Aseptic Techniques Quiz 2

2 Harvest Theory Quiz 4 2 Accessioning Assessment 5

10 Karyotype Test #1 6

2 Blood Setup Technical Performance 7 10 Midterm 8

10 Karyotype Test #2 9

2 Slide Making & Staining Quiz 11

4 Onscreen Chromosome Analysis 11

3 Microscope-Based Chromosome Analysis 12 10 Karyotype Test #3 12

2 Long-Term Cultures Quiz 13

10 Unknowns 14

1 Preventative Maintenance/Weekly Duty Responsibilities Completed Throughout Semester 30 Final Exam Fall Exam Period Final Exam Week (TBA)

Grading Scale – Performance Assessment Tasks (PATs)

In order to pass the course, students must complete all the following: Obtain a minimum grade of 60% on all karyotype tests o Should a student be unsuccessful on one karyotype test, they will be offered remediation and one further attempt to achieve the 60% pass rate on that karyotype test. The student must then obtain a minimum grade of 60% on all subsequent karyotype tests the first time Obtain a minimum grade of 70% for the final exam Obtain a minimum grade of 60% overall Note: A penalty of 10% is deducted for each 24-hour period (or part thereof) that an assignment is late

# **Program Outcom** in This Course

## The Michener Institute of Education at UHN Cytogenetics Theory & Techniques I

Course Description 

ourse ading

	This course supports the students' attainment of the knowledge, skills and attitudes expected of a Genetics Technology Program Graduate as per the CSMLS. Graduates will be highly competent professionals who:	<ol> <li>Have developed a broad scientific/genetic knowledge base and practice</li> <li>Apply critical thinking skills to constructively solve problems.</li> <li>Practice and promote the principles of continuous quality improveme using personal initiative to improve laboratory practices.</li> <li>Practice to ensure the safety of patients, colleagues, self, and the envious 5. Contribute to the health care and education of the public, promote padignity and confidentiality.</li> <li>Are integral members of the health care team that share knowledge t treatment of disease, promote learning, and collaborate with other profeare.</li> <li>Are responsible and accountable for professional acts and practices are legislation and regulations governing the profession.</li> <li>Abide by the CSMLS Code of Professional Conduct.</li> <li>Use effective interpersonal skills to maintain a professional relationsh professionals.</li> <li>Use all available resources to provide service in a timely, accurate, ar</li> </ol>		
			1. Demonstrates safe practices according to established protocols, safety guidelines, and existing legislature	Learning Objectives 1.a. Uses the principles o 1.b. Demonstrates the us 1.c. Uses infection contro 1.d. Understands the pot 1.e. Demonstrates the us 1.f. Demonstrates the us 1.g. Demonstrates the us 1.h. Applies universal pre 1.i. Demonstrates the ap 1.09, 1.10) 1.j. Understands and app 1.k. Understands and app
			2. Demonstrates accurate data and specimen collection and handling according to established protocols	Learning Objectives 2.a. Understands the req 2.b. Verifies that all requ 2.c. Assesses the suitabil 2.d. Verifies that the requ 2.e. Verifies the suitabilit 2.f. Demonstrates appro 2.g. Identifies discrepance 2.h. States the CCMG gui
			3. Evaluates the completion of pre-analytical processing on a variety of specimen types while following established protocols	Learning Objectives 3.a. Uses resources efficient 3.b. Completes culture set 3.c. Completes culturing 3.d. Completes culture h 3.e. Assesses the quality 3.f. Prepares slides for cy 3.g. Assesses the quality 3.h. Completes staining 3.i. Assesses the quality 3.j. Completes culture m 3.k. Assesses the quality
es Covered			4. Uses laboratory equipment and prepares reagents according to established protocols Learning Objectives	Learning Objectives 4.a. Applies the principles 4.b. Uses computer softw 4.c. Uses and maintains s 4.d. Understands the fun 4.e. Understands the pre
	Course Competencies a Learning Objectives	and	5. Evaluates patient data using scientific knowledge and skills to provide accurate genetic information according to established protocols	Learning Objectives 5.a. Understands the rela 5.b. Analyzes GTG stained 5.c. Analyzes GTG stained 5.d. Analyzes special stain 5.e. Recognizes variant o 5.f. Understands the nee 5.g. Determines the clinic 5.h. Assesses the quality
			6. Uses appropriate terminology, worksheets, reports and established protocols to effectively communicate accurate laboratory results	Learning Objectives 6.a. Records procedure 6.b. Records pertinent of 6.02) 6.c. Uses the current IS 6.d. Demonstrates critic 6.e. States CCMG guide 6.f. Demonstrates the n
			7. Uses principles of quality management to ensure the generation of high quality laboratory results	Learning Objectives 7.a. Completes laborate 7.b. Records and assess 7.c. Uses appropriate co 7.d. Records data in acc 7.e. Identifies deficienc 7.f. Completes assigned 7.g. Understands reage 7.h. Applies concepts of 7.i. Applies reflection fo
			8. Applies critical thinking skills towards the efficient use of materials in the lab	Learning Objectives 8.a. Demonstrates the k
			9. Applies effective communication and teamwork skills	Learning Objectives 9.a. Understands cond communication, writte 9.b. Demonstrates eff
			10. Applies the standards of professional practices when working on specimen to protect the patient's	Learning Objective 10.a. Understands the dignity, values,

right to a reasonable standard of care

such activities (CSMLS 10.10)

Please refer to Michener's Academic Policies located at: .

Changes to course outlines are governed by The Michener Institute's Course Management Policy located here: https://michener.ca/discover-michener/policies/course-management-

Please note that the Michener Institute of Education at UHN may be required to modify content and/or mode of delivery and/or mode of assessment of any course in order to comply with institutional directives or emergency orders from government officials in relation to the ongoing situation with the COVID-19 virus. All changes will be reviewed programmatically to ensure all competencies are covered and program requirements satisfied.

Turnitin

Academic

Policies

policy/

This course may leverage the use of Turnitin, a software platform that helps to support students in their writing by ensuring submitted works meet Michener's standards for academic integrity. Normally, students will be required to submit their written assessments to Turnitin for a review of originality. In doing so, students will allow their work to be included as source documents in the tool's reference database, where they will be used solely for the purpose of verifying originality. Students may opt out of this process by contacting their course instructor at the beginning of the course, in which case the course instructor will arrange alternative methods to verify originality. Refer to Michener's Learning Resource Centre for guidelines on how to properly reference in academic writing.

Minimum Technology Requirements

Refer to Michener's current minimum technology requirements. For Helpdesk support and FAQs please click here.

### Accessibility and Accommodations

Students with diverse learning needs are welcome in this course. If you have a disability that may require accommodations, please visit Michener's Accessibility & Accommodations Services, or contact accessibility@michener.ca.

The Institute provides reasonable accommodation of the needs of students who observe religious holy days other than those already accommodated by ordinary scheduling and statutory holidays. Students have a responsibility to alert members of their faculty in a timely fashion to upcoming religious observances and anticipated absences and instructors will make every reasonable effort to avoid scheduling tests, examinations or other compulsory activities at these times. Please see Michener's Religious or Spiritual Observance Policy for more information.

#### Library Resources for Students

The Learning Resource Collaboratory Virtual Portal is your gateway to information: From this website, find your one-stop-shop for all your program information needs under our Program Resource Guides. You have questions? We have answers! Chat with a Librarian at lrc@michener.ca.

### Learning Supports

The Student Success Network provides learning supports to help students achieve academic success. Workshops are available to be tailored to meet your learning needs by a learning facilitator. Peer tutoring is also available to support your academic progress. Please contact success@michener.ca for more information.

tical skills.

ent including professional development and

/ironment. patient welfare, and respect the patient's

that is essential to the diagnosis and

ofessionals in providing effective patient

according to standards of practice as well as

hip with clients and health care

nd cost-effective manner.

of standard precautions to minimize dangers to self and others (CSMLS 1.01) use of appropriate personal protective equipment when required (CSMLS 1.02) rol practices and uses good laboratory hygiene (CSMLS 1.03) tential dangers posed by biological specimens, laboratory supplies and equipment (CSMLS 1.04) sage of laboratory safety equipment in a correct and safe manner (e.g. Fume hoods, biological safety cabinets) (CSMLS 1.05) se of WHMIS and existing legislation for the labelling, handling, storing, and disposing of chemicals, reagents and solutions (CSMLS 1.06) use and disposal of sharps appropriately (CSMLS 1.07) ecautions and existing legislation to handle, transport, store, and dispose of biological and other hazardous materials (CSMLS 1.08) ppropriate method for disinfection/sterilization within the lab, while minimizing the associated risks and hazards associated with such method (CSMLS plies when needed the appropriate measures for spill containment and clean-up for biological and other hazardous materials (CSMLS 1.12) plies when needed the procedure for responding to and reporting incidents related to safety and personal injury within the laboratory (CSMLS 1.11, 1.14) quirements regarding specimen collection, transportation, and storage (CSMLS 2.01) uired data between the specimen and requisition match (CSMLS 2.02) ility of the sample for testing, including sample volume and integrity (CSMLS 2.03) uired clinical information is present on the requisition (CSMLS 2.04) ty of the test requested for the referral reason (CSMLS 2.05) opriate data entry and retrieval from the laboratory information system (CSMLS 2.07) icies during specimen data entry and initiates corrective action as required (CSMLS 2.08) idelines for specimen retention (CSMLS 2.09) ciently during specimen processing (e.g. time, equipment) (CSMLS 3.02) set up for cytogenetic specimens (CSMLS 3.03) g of cytogenetic specimens (CSMLS 3.04) harvest for cytogenetic specimens (CSMLS 3.05) y of short term culture preparations throughout processing and initiates corrective action as required (CSMLS 3.03.01, 3.04.01, 3.05.01) cytogenetic analyses (CSMLS 3.06) y of slide preparations and initiates corrective action as required (CSMLS 3.06.01) g of slides for cytogenetic analyses (CSMLS 3.07) of stained slide preparations and initiates corrective action as required (CSMLS 3.07.01) naintenance for long term cytogenetic specimens (CSMLS 3.10) y of long term culture preparations throughout processing and initiates corrective action as required (CSMLS 3.10.01) es of light microscopy for specimen assessment and analysis (Brightfield, Inverted and Phase Contrast) (CSMLS 4.01) ware to capture, enhance and annotate metaphase images, as well as create karyotypes (CSMLS 4.02, 4.02.01) standard laboratory equipment (e.g. centrifuges, incubators, waterbaths, pipettes) (CSMLS 4.04) nction of standard laboratory equipment and identifies basic malfunctions (CSMLS 4.05) reparation of cytogenetic reagents and culture media (CSMLS 4.06) ationship between clinical information, limits of laboratory analyses, the test result, and their impact on patient care (CSMLS 5.01) ed metaphase chromosomes (CSMLS 5.02) ed karyotypes (CSMLS 5.03) ined chromosomes (CSMLS 5.04) or abnormal genetic findings and states appropriate follow up if required (CSMLS 5.10) eed for further genetic testing and states appropriate follow up if required (CSMLS 5.11) ical significance of cytogenetic findings in relation to the patient's referral reason (CSMLS 5.12) of test results and states corrective action as required (CSMLS 5.13) es and observations throughout specimen processing and analysis according to established protocols (CSMLS 6.01) quality control information according to established protocols (e.g. band resolution, quality of preparations) (CSMLS SCN system to describe cytogenetic results (CSMLS 6.03) itical appraisal of recorded documentation to ensure accurate test results (CSMLS 6.05) elines for retention of test records (CSMLS 6.07) maintenance of patient confidentiality (CSMLS 6.08) tory testing using established policies and protocols (CSMLS 7.02) sses quality control data (e.g. daily QC readings, band resolution) (CSMLS 7.03) controls to validate analyses where required (CSMLS 7.04) cordance with quality assurance procedures (e.g. weekly trends of QC readings) (CSMLS 7.06) cies in the workplace that may affect testing quality (CSMLS 7.07) ed duties in accordance with preventative maintenance program (e.g. weekly maintenance) (CSMLS 7.08) gent quality control (CSMLS 7.09) of quality control with respect to the use of new reagents (CSMLS 7.10) e attention to detail skills to ensure specimens are correctly identified at all times during processing and analysis (CSMLS 7.11) or the purpose of continuous quality improvement and risk management to promote quality clinical laboratory services (CSMLS 7.12) e knowledge of inventory control through conservative use of resources (CSMLS 8.04)

cepts related to effective communication with other health care professionals, including: Active listening, verbal communication, non-verbal ten communication, barriers to effective communication, and use of technology appropriately to facilitate communication (CSMLS 9.01) ffective teamwork skills while completing tasks (CSMLS 9.03)

concepts related to the welfare and confidentiality of the patient, including respect for

, and beliefs of the patient (CSMLS 10.06) 10.b. Recognizes potentially dangerous situations and understands the right to refuse to participate in