Changing the Way We Work Community of Practice for Ontario Family Physicians

#### June 6, 2025

Dr. Daniel Warshafsky Dr. Rachita Gurtu



#### Infectious Disease and Management of STIs Part II





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Moderator:

• Dr. Eleanor Colledge, CPD Program Director, University of Toronto and Family Physician, South East Toronto Family Health Team, Toronto, ON

Panelists:

- Dr. Daniel Warshafsky, Toronto, ON
- Dr. Rachita Gurtu, Mississauga, ON

Host:

• Dr. Jobin Varughese, Brampton, ON

The Changing the Way We Work Community of Practice for Ontario Family Physicians is a one-credit-per-hour Group Learning program that has been certified for up to a total of 32 credits.

Please note that due to changes to the Cert+ platform, there will be delays in credits being applied to your account.

#### Self-learning program

The session materials, including recordings, tools, and resources are available as self-learning modules.

This one-credit-per-hour Group Learning program has been certified by the College of Family Physicians of Canada and the Ontario Chapter for up to 80 credits.

To participate in this self-learning:

- Select the dates/sessions you wish to participate in. You are welcome to complete as many sessions as you wish.
- Watch the video recording of the live session.
- Review the session tools and resources.
- Complete the self-learning post-session activity, click the button below.

Complete self-learning activity 🗷

Self-Learning Activity and Evaluation: COVID-19 Community of Practice for Ontario Family Physicians

By completing this Self-Learning Activity for the COVID-19 Community of Practice for Ontario Family Physicians, you are confirming that you have completed this activity.

\* 1. Attestation: I confirm that I have completed the COVID-19 CoP self-learning activity (video and resources). (If completing multiple session dates, please enter all that apply below ENTER DATE AS Month-Day-Year i.e. December 10, 2021)

Session Date(s):	
Name:	
Email:	

\* 2. After reviewing this COVID-19 session material (video and resources ), I have a question (s) regarding the content that needs clarifying.

I have no questions

Question:

### Missed a session and want to earn credits?

The Self-learning Program lets you earn credits for watching past sessions. Just click the link and fill out a 60 second survey!

#### Land Acknowledgement

We acknowledge that the lands on which we are hosting this meeting include the traditional territories of many nations.

The OCFP and DFCM recognizes that the many injustices experienced by the Indigenous Peoples of what we now call Canada continue to affect their health and well-being. The OCFP and DFCM respects that Indigenous people have rich cultural and traditional practices that have been known to improve health outcomes.

I invite all of us to reflect on the territories you are calling in from as we commit ourselves to gaining knowledge; forging a new, culturally safe relationship; and contributing to reconciliation.

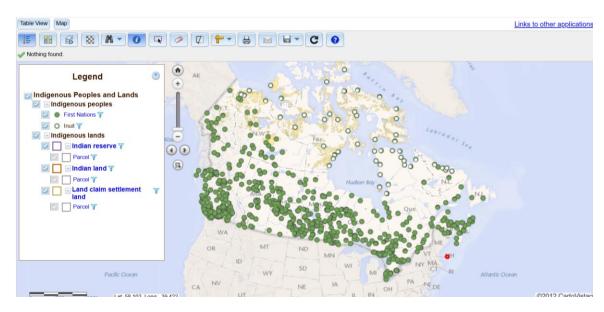


#### **Discover stories and traditions**

National Indigenous History Month

# <image><complex-block><complex-block>

#### Learn more about present day communities



### Explore the past and honour the truth



Reconciliation:

A Starting Point

A product of a proud partnership between the Canada School of Public Service and the Department of National Defence

<sup>(i)</sup> Before You Begin <sup>></sup>

Get Started

Version 3.0 (May 2025)



### Changing the way we work

#### A community of practice for family physicians

At the conclusion of this <u>series</u> participants will be able to:

- Identify the current best practices for delivery of primary care and how to incorporate into practice.
- Describe point-of-care resources and tools available to guide decision making and plan of care.
- Connect with a community of family physicians to identify practical solutions for their primary care practice under current conditions.

#### **Disclosure of Financial Support**

This CPD program has received in-kind support from the Ontario College of Family Physicians and the Department of Family and Community Medicine, University of Toronto in the form of logistical and promotional support.

#### **Potential for conflict(s) of interest:** N/A

#### **Mitigating Potential Bias**

- The Scientific Planning Committee has full control over the choice of topics/speakers.
- Content has been developed according to the standards and expectations of the Mainpro+ certification program.
- The program content was reviewed by a three-member national/scientific planning committee.

*Planning Committee*: Dr. Jobin Varughese (OCFP), Dr. Ali Damji (DFCM), Dr. Eleanor Colledge (DFCM), Dr. Harry O'Halloran, Julia Galbraith (OCFP), Pavethra Yogeswaran (OCFP), Marisa Schwartz (DFCM)

#### Previous webinars & related resources:

https://www.dfcm.utoronto.ca/covid-19-community-practice/past-sessions



#### Dr. Daniel Warshafsky – Panelist

Associate Chief Medical Officer of Health at the Office of the Chief Medical Officer of Health



#### Dr. Rachita Gurtu – Panelist

Family Physician & Medical Director, Healthy Sexuality Clinics Region of Peel Public Health

### **Speaker Disclosure**

- Faculty Name: **Dr. Daniel Warshafsky**
- Relationships with financial sponsors:
  - Grants/Research Support: N/A
  - Speakers Bureau/Honoraria: N/A
  - Others: N/A

- Faculty Name: Dr. Rachita Gurtu
- Relationships with financial sponsors:
  - Grants/Research Support: N/A
  - Speakers Bureau/Honoraria: Ontario College of Family Physicians, UofT Family Medicine Program - Trillium site
  - Others: N/A

#### **Speaker Disclosure**

- Faculty Name: **Dr. Jobin Varughese**
- Relationships with financial sponsors:
  - Grants/Research Support: N/A
  - Speakers Bureau/Honoraria: Ontario College of Family Physicians
  - Others: Toronto Metropolitan University, School of Medicine (Interim Assistant Dean of Primary Care Education), William Osler Health System (Associate Vice President of Academics)

- Faculty Name: **Dr. Eleanor Colledge**
- Relationships with financial sponsors:
  - Grants/Research Support: N/A
  - Speakers Bureau/Honoraria: Ontario College of Family Physicians
  - Others: The Foundation for Medical Practice Education (McMaster University)

### **How to Participate**

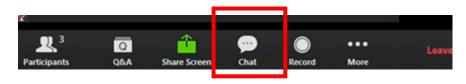
• All questions should be asked using the Q&A function at the bottom of your screen.



• Press the thumbs up button to upvote another guest's questions. Upvote a question if you want to ask a similar question or want to see a guest's question go to the top and catch the panels attention.

🗢 Q&A			
	All questions (1)	My questions	
Lee 01:54 PM			
Will there be a foll	ow-up session?		
1 <b>4</b>			Comment

• Please use the chat box for networking purposes only.





#### Dr. Daniel Warshafsky – Panelist

Associate Chief Medical Officer of Health at the Office of the Chief Medical Officer of Health



#### Dr. Rachita Gurtu – Panelist

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### **Public Health Update**

Changing the Way We Work CoP

Dr. Daniel Warshafsky Office of the Chief Medical Officer of Health

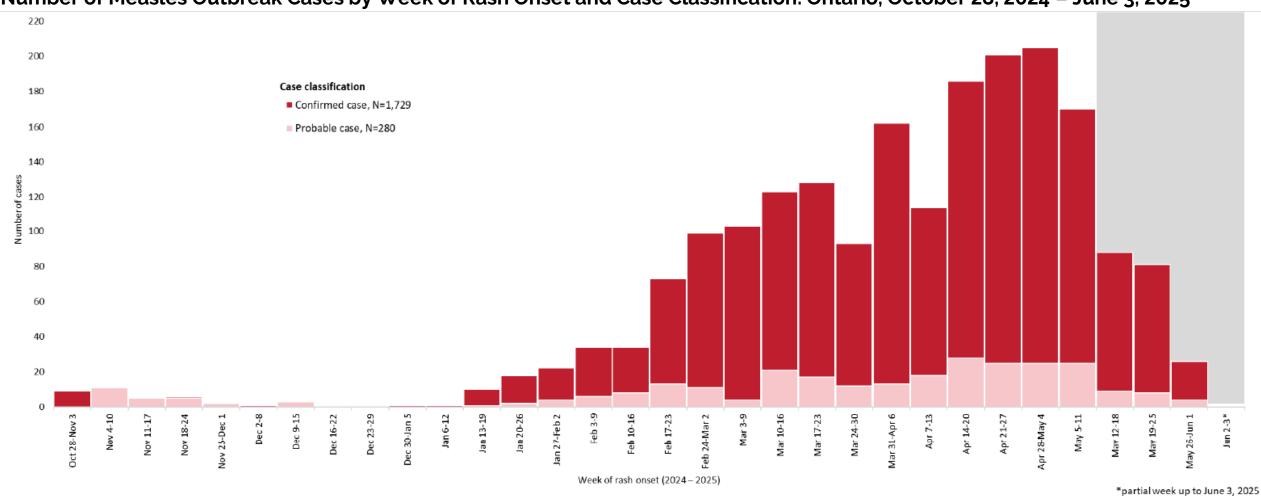
June 6, 2025



### **Ontario Measles Outbreak Update**

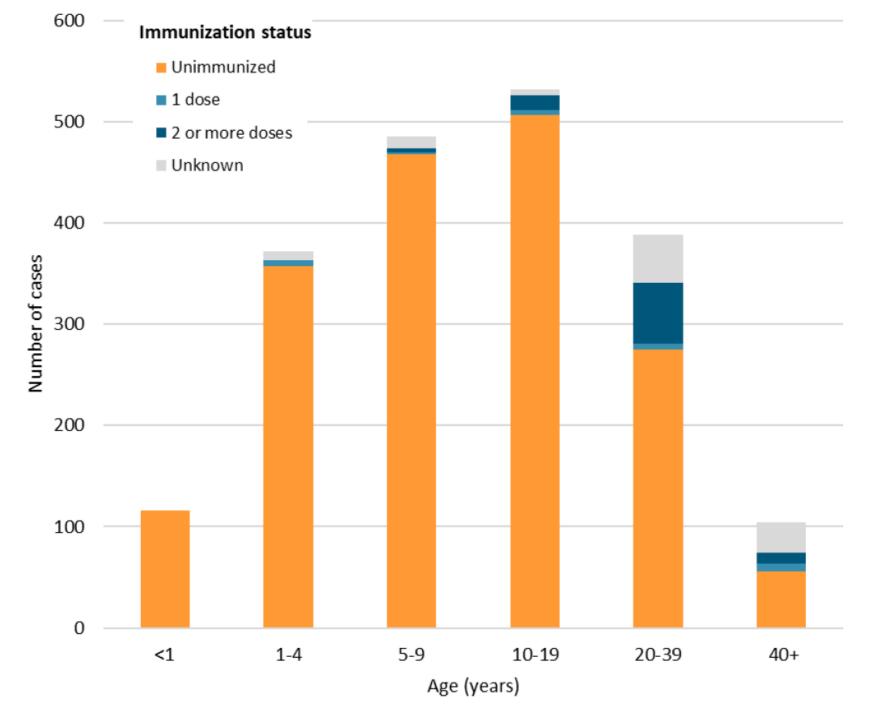
- Measles continues to spread in Ontario due to ongoing exposures and transmission among susceptible individuals
- As of June 3, 2025,
  - 2,009 cases in 19 public health units have been reported in association with this outbreak
    - This represents an increase of 121 cases since May 27, 2025
  - 140 hospitalizations (7.0%)
  - 9 ICU admissions (0.4%)
  - 39 pregnant persons
  - 6 cases of congenital measles
- There was one death that occurred in a congenital case of measles who was born pre-term and had other underlying medical conditions
- For more information on place and dates of exposure to measles in Ontario, please visit Public Health Ontario's <u>website</u>





Week of rash onset	Oct 28-Nov 3	Nov 4-10	Nov 11-17	Nov 18-24	Nov 25-Dec 1	Dec 2-8	Dec 9-15	Dec 16-22	Dec 23-29	Dec 30-Jan 5	Jan 6-12	Jan 13-19	Jan 20-26	Jan 27-Feb 2	Feb 3-9	Feb 10-16	Feb 17-23	Feb 24-Mar 2	Mar 3-9	Mar 10-16	Mar 17-23	Mar 24-30	Mar 31-Apr 6	Apr 7-13	Apr 14-20	Apr 21-27	Apr 28-May 4	May 5-11	May 12-18	May 19-25	May 26-Jun 1	Jun 2-3*
Confirmed case	9	0	0	1	0	1	0	0	0	1	1	9	16	18	28	26	60	88	99	102	111	81	149	96	158	176	180	145	79	73	22	0
Probable case	0	11	5	5	2	0	3	0	0	0	0	1	2	4	6	8	13	11	4	21	17	12	13	18	28	25	25	25	9	8	4	0
Total cases	9	11	5	6	2	1	3	0	0	1	1	10	18	22	34	34	73	99	103	123	128	93	162	114	186	201	205	170	88	81	26	0

#### Number of Measles Outbreak Cases by Week of Rash Onset and Case Classification: Ontario, October 28, 2024 – June 3, 2025



94% of cases in the current measles outbreak were unimmunized or had unknown immunization status



#### (A) Cumulative cases: October 28, 2024 – June 3, 2025

100.0+ 50.0 - 99.9 SUD 25.0 - 49.9 10.0 - 24.9 0.1 - 9.9 0 EOH GBO Code Name Code Name ALG Algoma Public Health Unit CHK Chatham-Kent Health Unit Durham Region Health Department DUR OES EOH Eastern Ontario Health Unit GEH Grand Erie Health Unit GBO Grey Bruce Health Unit WEC HNP Haliburton Kawartha Northumberland Peterborough Health Unit HAL Halton Region Health Department HAM Hamilton Public Health Services HPH Huron Perth District Health Unit LAM Lambton Public Health MSL Middlesex-London Health Unit

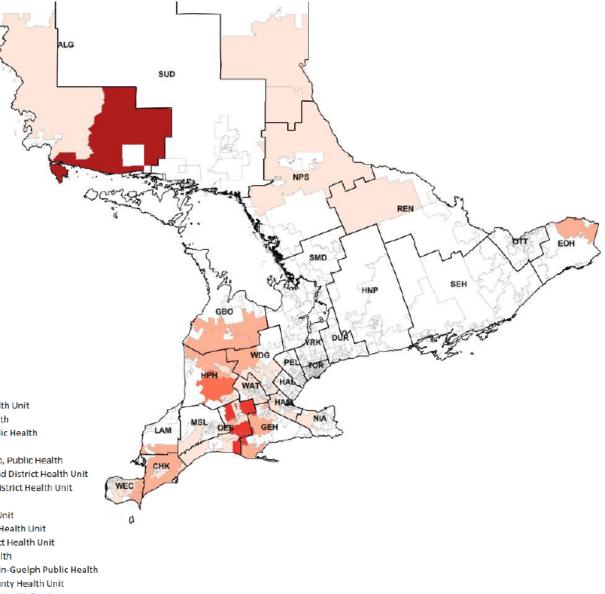
NIA

FSA rate per 100,000 population

#### **Public Health Unit**

- NWR Northwestern Health Unit
- OTT Ottawa Public Health
- Southwestern Public Health
- PEL Peel Public Health
- WAT Region of Waterloo, Public Health
- REN Renfrew County and District Health Unit
- SMD Simcoe Muskoka District Health Unit
- SEH South East Health Unit
- SUD Sudbury & District Health Unit
- THB Thunder Bay District Health Unit
- TOR Toronto Public Health
- WDG Wellington-Dufferin-Guelph Public Health
- Niagara Region Public Health Department WEC Windsor-Essex County Health Unit
- NPS North Bay Parry Sound District Health Unit YRK York Region Public Health Services
- NEH Northeastern Health Unit

#### (B) Recent cases (rash onset in the last 21 days): May 13, 2025 - June 3, 2025



# Testing: Collect specimen from multiple sites and within 7 days of rash onset

For all suspected measles cases, collect NP/throat swab <u>and</u> urine

Test	Specimen Type & Volume	Collection Kit	Timing of Collection
Measles virus detection (PCR)	Nasopharyngeal Swab	Virus Respiratory Kit# 390082	• Within 7 days of rash onset.
Measles virus detection (PCR)	Throat Swab	Virus Culture Kit# 390081	• Within 7 days of rash onset.
Measles virus detection (PCR)	Urine (minimum 10mL)	Sterile container	• Within 14 days of rash onset.

- For laboratory/testing related questions and support, call PHO's Laboratory Customer Service Centre
  - Business hours: 416-235-6556 or toll Free: 1-877-604-4567
  - After hours: 416-605-3113 (duty officer)

### Don't forget clinical information on your requisition!

- Use PHO Laboratory Requisition form BOTH virus detection (PCR) and diagnostic serology
- Check "diagnosis" box and clearly mark "Suspect case of measles" in the Testing Indications

Testing Indication(s) / Criteria								
Jiagnosis	Screening	Immune Status	Follow-up / Convalescent					
Pregnancy / Perinatal	Impaired Immunity	Post- mortem						
Other (Specify): Suspect case of Measles								

- Include the following information on each requisition:
  - Patient's symptoms and onset date
  - Exposure history, travel history (if applicable), MMR/V vaccination history
  - Outbreak or investigation number (if applicable)

### Reporting and Public Health Unit (PHU) Response

All suspect cases of measles should immediately be reported to the local PHU – **do not wait for lab confirmation** 

- Individuals with suspected measles should be advised to isolate while lab results are pending
- For confirmed measles cases, the local PHU will perform contact identification and management, which may include recommendations for measles PEP and/or exclusion from work, school, or other high-risk settings for susceptible contacts
  - Local PHU may contact HCPs to request assistance in determining a patient's susceptibility to measles, and providing measles vaccine to known contacts for PEP for patients in their practices
- For more information on measles PEP, please refer to Public Health Ontario's <u>document</u>

# IPAC Considerations for Measles When Providing Care to Patients with Suspect/Confirmed Measles

- Only health care workers with presumptive immunity to measles should provide care to patients with suspected/confirmed measles
  - Evidence of presumptive immunity = at least two doses of measles-containing vaccine (MMR/V) after 1 year of age OR laboratory evidence of immunity
  - Consider obtaining staff's evidence of immunity on file to avoid staff exclusion in the event of a measles exposure
- Health care workers should wear a fit-tested, seal-checked N95 respirator when providing care
  - **Droplet and Contact Precautions** (gloves, gown, eye protection) are also recommended due to risk of exposure to rash and respiratory secretions
- If referring patients to other health care settings (e.g., lab, hospital), call ahead prior to patient's arrival so that appropriate IPAC precautions can be implemented to avoid exposures (i.e., mask upon arrival, arrange for patient to be placed immediately in an appropriate isolation room)
- For more information on IPAC practices, please refer to Public Health Ontario's <u>webpage</u>

### **Outbreak Immunization Strategy in Impacted Areas**

As part of the outbreak management strategy, individuals who live, work, travel (e.g., family visit), worship, or spend time in affected regions and communities\* with active measles cases are recommended to receive MMR vaccine as follows:

Age group	Recommendation
Infants (six to 11 months)	Should receive one dose of the measles, mumps, rubella (MMR) vaccine. Note: <b>Two</b> additional doses are required after the age of one year
Children (one to four years)	Children who have received their first dose of MMR vaccine are encouraged to receive a second dose as soon as possible (at a minimum of four weeks from the first dose)
Adults (18+ years) born on or after 1970	A second dose of MMR vaccine is recommended

 These recommendations also apply to those who are travelling to areas where measles is of concern, either domestically or internationally

\* Affected regions refer to southwestern Ontario, specifically Southwestern, Grand Erie, Huron Perth, Chatham-Kent, Windsor-Essex, and other regions

### **General Immunization Considerations**

- Serologic testing is not recommended before or after receiving measles-containing vaccine
- If an individual's immunization records are unavailable, immunization with measlescontaining vaccine is generally preferred to ordering serology to determine immune status
  - There is no harm in giving measles-containing vaccine to an individual who is already immune

### **Measles in Pregnancy**

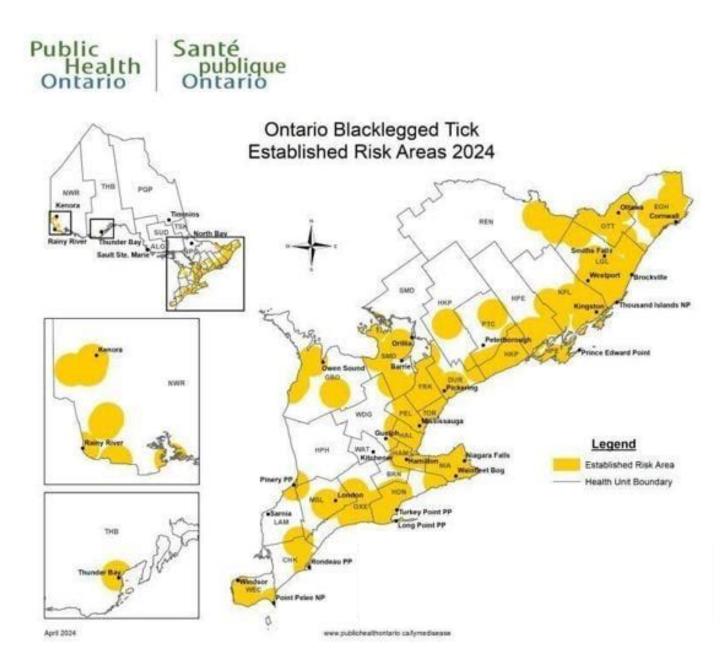
- What are the risks of maternal measles infection during pregnancy?
  - Increased risk of maternal complications
  - Pregnancy loss
  - Preterm birth
  - Low birth weight
  - Congenital measles infection in the infant
- Actions for HCPs:
  - Assess measles immunity status of pregnant patients, ideally prior to pregnancy
    - If non-immune, MMR vaccination should be given at least 4 weeks prior to becoming pregnant

### Lyme Disease

- Tick-borne illness caused by *Borrelia burgdorferi*
- In **2024**, there were **2,337 cases** of Lyme disease reported across Ontario (149.8 cases per 1,000,000 population)
  - Higher than 5-year average of 97.2 cases per 1,000,000 population
- In **2025**, there are **85 cases** year-to-date (data available for January to March)
- However, ticks are most active in the **spring**, **summer**, **and fall**

### **Established Risk Areas**

- Despite established risk areas, it is possible to encounter an infective blacklegged tick almost anywhere in Ontario, given that the habitat is suitable (e.g., wooded or brushy areas)
- Refer to PHO's <u>Ontario Vector-</u> <u>Borne Disease Tool</u> for up-todate surveillance data



### **Tick Submission and Testing**

- Tick submission is **no longer required for surveillance in most areas in Ontario** 
  - Primary purpose of tick identification is to monitor new and emerging tick populations
  - Submitting ticks from areas with established tick populations does not provide additional information
- Ticks can be submitted for identification within 48 hours through the <u>eTick website</u>

### Lyme Disease Prevention

#### • Before going into wooded or brushy areas:

- Wear closed-toe shoes
- Tuck shirt into pants, and pants into socks
- Wear light-coloured, long-sleeved clothes to spot ticks easily
- Apply insect repellant to clothing and exposed skin

#### • While outdoors:

- Walk on cleared paths/trails
- Avoid using trails created by animals (such as deer and moose), as ticks are often found on the grass and plants along these trails

#### • Before returning or when you are indoors:

- Do a full-body tick check on yourself, children, and pets
- Shower as soon as possible to wash off unattached ticks
- If you find an attached tick, remove it by the head as soon as possible



Explore geographic trends in vector-borne diseases dating back to 2014 up until the current week. Use the tabs at the bottom of the tool to navigate between vectorspecies and surveillance topics. Data will be updated weekly for all VBD human cases. Mosquito data will be updated weekly during the mosquito season (May-October).

#### Key features:

Overview - ON: View provincial-level most recent week (MRW) and year-to-date (YTD) case counts for all reportable vector-borne diseases (VBD), including West Nile virus illness, Lyme disease, anaplasmosis, babesiosis, Powassan virus infection. Overview - PHU: Explore data at the public health unit level, including most recent week (MRW) and year-to-date (YTD) case counts, as well as percent positivity (PP) for West Nile virus (WNV) and eastern equine encephalitis virus (EEEV) in mosquito pools. Maps: To change VBD dimensions, turn on/off the layer of interest from the layer drop-down button within each map tab.

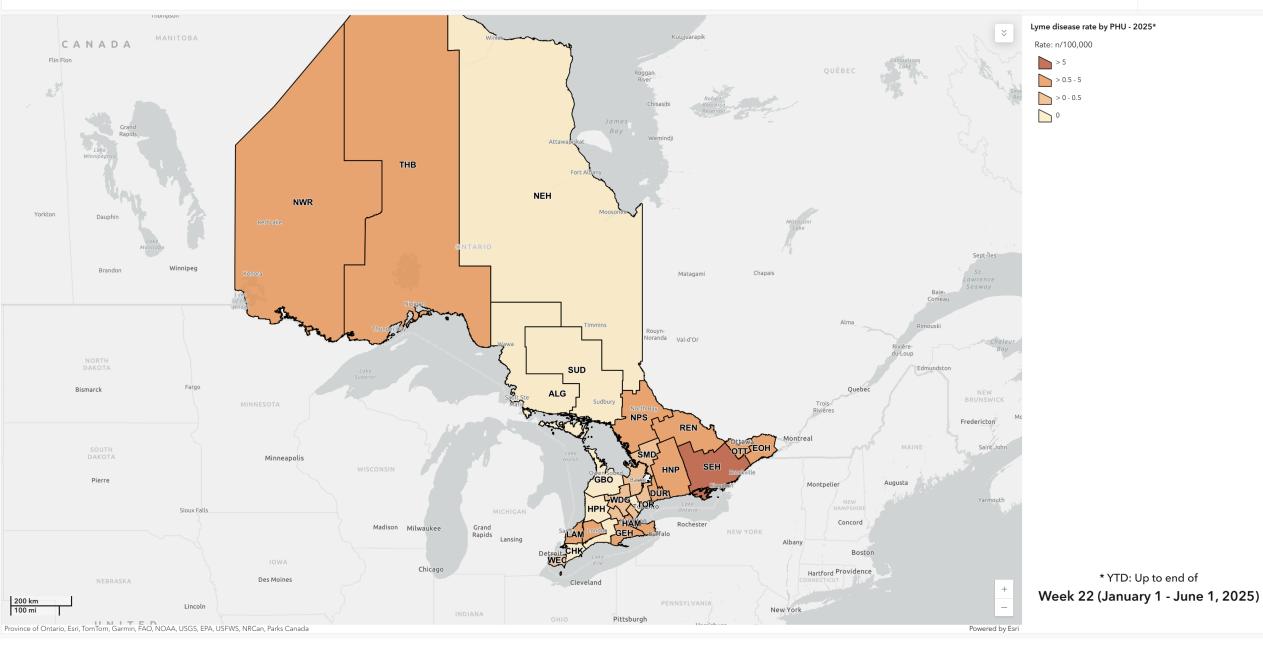
Need help using the tool? Technical notes and a user guide are available in the menu at the top right corner of the tool to support interpretation and navigation.

Contact communicable.diseasecontrol@oahpp.ca for more information.

in West Nile Virus	<b>O</b> Cases - Most Recent Week	<b>O</b> Cases - YTD				
🕷 Lyme Disease	<b>5</b> Cases - Most Recent Week	<b>165</b> Cases - YTD				
Anaplasmosis	<b>O</b> Cases - Most Recent Week	<b>5</b> Cases - YTD				
Babesiosis	<b>O</b> Cases - Most Recent Week	<b>O</b> Cases - YTD				
🕷 Powassan Virus Disease	<b>O</b> Cases - Most Recent Week	<b>O</b> Cases - YTD				
Most Recent Week: Week 22	(May 26 - June 1, 2025); YTD: Up to end of Week 22 (January 1 - June 1, 2025)					
Overview - Ontario Overview - PHU BLT Risk Areas Mosquito Traps Mosquito Species Anaplasmosis Babe	esiosis EEEV LD Powassan WNV					

Ontario Vector-Borne Disease Tool

#### Lyme Disease Human Cases, Positive Ticks by Public Health Unit



### Testing for STIs in Primary Care

Dr. Rachita Gurtu MD, CCFP

### Objectives

- Testing indications for genital herpes and Trichomonas
- Available tests for genital herpes and Trichomonas in Ontario
- Treatment recommendations for genital herpes and Trichomonas

### Genital Herpes

### Genital Herpes- Counselling key points

- Devastating diagnosis due attached social stigma
- Common condition
  - ~14% individuals 14-59 years old in Canada have HSV-2
- Recurrent, chronic infection; however, it is a manageable condition
- Infection can live in your body for a long time undetected or unrecognized
- Most people who have genital herpes don't know they have it because they have mild, short-lived or no symptoms, or they think the symptoms are due to another condition
- Condoms, if used consistently and correctly will reduce, but will not eliminate the risk of HSV transmission or acquisition

### **Genital Herpes-Testing**

#### • HSV NAAT

• Sample from lesions-approaches sensitivity and specificity of 100%

#### • Type Specific Serology (TSS)

- Not recommended for screening in asymptomatic individuals
- When to consider:
  - Signs and symptoms of HSV are present but NAAT is negative or not feasible
    - Note: repeat viral testing of fresh lesions is preferred over TSS
  - To identify the need for preventative measures when sexual partners are suspected to be serodifferent/serodiscordant

### **Genital Herpes-Testing**



#### **General Test Requisition**

ALL sections of the form must be completed by authorized health care providers for each specimen submitted, or testing may be delayed or cancelled. Verify that all testing requirements are met before collecting a specimen. For HIV, respiratory viruses, or culture isolate requests, use the dedicated requisitions available at: publichealthontario.ca/requisitions

Ordering Hea	althcare Provider Information	
Licence No.:	Healthcare Provider Full Name:	Health Card No.: 12345678
001234	Dr. R. Gurtu	Date of Birth (yyyy-mm-dd): 1992-01-20 Sex: Male
Org. Name:	Address:	Medical Record No.: Female
City:	Postal Code: Province:	Last Name (per health card): DOC
Tel:	Fax:	First Name (per health card): Jane
Copy to Lab / H	lealth Unit / Other Authorized Healthcare Provider	Address: Postal Code:
Licence No.:	Lab / Health Unit / Other Authorized Provider Name:	City: Tel:
		Investigation / Outbreak No. from
Org. Name:	Address:	PHO or Health Unit (if applicable):
City:	Postal Code: Province:	Specimen Information
Tel:	Fax:	Date Collected (yyyy-mm-dd): 2023-10-02 Submitter Lab No.:
		Whole Blood Serum Plasma
Patient Setti	•	Bone Marrow Cerebrospinal Nasopharyngea Fluid (CSF) Swab (NPS)
Clinic / Community	ER (Not Admitted / Not Yet Determined) ER (Admitted)	Oropharyngeal Bronchoalveola
Inpatient (Non-ICU)	ICU / CCU Congregate Living Setting	/ Throat Swab Sputtin Lavage (BAL)
Testing India	ation(s) / Criteria	Swab
J Diagnosis	Screening Immune Follow-up / Status Convalescent	Urine Rectal Swab Faeces
Pregnancy / Perinatal	Impaired Post- Immunity mortem	Other (Specify type AND body location): LABIA SWAB
Other (Specify):		Test(s) Requested
Signs / Sym	atoms	Enter each assay as per the <u>publichealthontario.ca/testdirectory</u> :
No Signs /	▲ Onset Date	HSV NAAI
Symptoms	★ (yyyy-mm-dd): 2023-09-28	2.
	Fever Rash STI	3.
Gastrointesti	nal Respiratory Hepatitis Meningitis / Encephalitis	4.
Other (Specify):		5.
Relevant Exp	posure(s)	6.
None / Not Applicable	Most Recent Date	For routine hepatitis A, B or C serology, complete this section instead
(	(yyyy-mm-dd): Decupational Exposure / Veedlestick Injury (Specify): Source Exposed	Hepatitis A Immune Status Acute Infection (HAV IgG) Symptoms info)
Other (Specify):		Hepatitis B Immune Status Chronic Infection (anti-HBs) (HBsAg + total anti-HBc
Relevant Tra	vel(s)	Acute Infection (HBsAg + total anti-HBc Screening (anti-HBs +
None / Not Applicable	Most Recent Date (yyyy-mm-dd):	
Travel Details:	()))	Hepatitis C Current / Past Infection (HCV total antibodies) No immune status test for HCV is currently available.
/ciail5.		

The personal health information is collected under the authority of the Personal Health Information Protection Act, s.36 (1)(c)(iii) for the purpose of clinical laboratory testing. If you have questions about the collection of this personal health information please contact the PHO's Laboratory Customer Service at 416-235-6556 or toll free 1-877-604-4567. F-SD-SCG-1000, version 004 (September 2023).

Santé publique Ontario Public Health Ontario For Public Health Ontario's laboratory use only:

PHO Lab No .:

Ontario 🕅

Date Received

(yyyy-mm-dd):

Patient Information

### Genital Herpes-Treatment (non-pregnant)

Treatment	Instructions for use	Comments
Primary episode	<ul> <li>Acyclovir 200 mg PO five times per day for 5-10 days [A-I]- ideally within 7 days after symptom onset</li> <li>or</li> <li>Famciclovir 250 mg PO TID for 5 days [A-I]- ideally within 5 days after symptom onset</li> <li>or</li> <li>Valacyclovir 1000 mg PO BID for 10 days [A-I]- ideally within 3 days after symptom onset</li> </ul>	<ul> <li>Provide treatment to those experiencing a first episode unless all lesions have already healed.</li> </ul>
Episodic treatment	Valacyclovir 500 mg PO BID OR 1 g PO once daily for 3 days [B-I] or Famciclovir 125 mg PO BID for 5 days [B-I] or Acyclovir 200 mg PO 5 times per day for 5 days [C-I]	<ul> <li>Prompt initiation at the onset of prodromal symptoms may shorten the severity and duration of lesions</li> </ul>
Suppressive Treatment	Valacyclovir 500 mg PO OD [A-I] (for people with ≤ 9 recurrences per year) or Valacyclovir 1000 mg PO OD [A-I] (for people with >9 recurrences per year) or Acyclovir 200 mg PO 3-5 five times per day or 400 mg PO BID [A-I] or Famciclovir 250 mg PO BID [A-I]	<ul> <li>Reduces length, frequency and severity of recurrences, asymptomatic viral shedding and transmission</li> </ul>

### Genital Herpes-Treatment (in pregnancy)

Treatment	Instructions for use	Comments
Primary episode	Acyclovir 200 mg PO QID for 5-10 days [A-I]	<ul> <li>C-section can reduce the risk of vertical transmission</li> <li>C-section is recommended in the case of newly acquired genital HSV in the third trimester</li> </ul>
Suppressive Treatment	Acyclovir 200 mg PO QID [A-I] or Acyclovir 400 mg PO TID [A-I] or Valacyclovir 500 mg PO BID [A-I]	<ul> <li>To reduce the risk of outbreak and asymptomatic viral shedding at the time of delivery and the need for C- section</li> <li>Start at GA 36 weeks and continue until delivery for anyone with a history of HSV-2 and for those who had a recurrence of genital herpes within the previous year.</li> </ul>

# Trichomonas

### Trichomonas vaginalis

- Protozoan infection NOT a reportable infection in Ontario
- Sexually transmitted cause of vaginitis
  - Penis-vagina sex, vagina-vagina, vagina-toys/fomites
- Often asymptomatic, particularly in males/trans females
- Signs and symptoms:
  - Vaginal discharge (yellow-green, thin, large amount, frothy)
  - Erythema of vulva and cervix ("strawberry cervix")
  - Vulvar irritation or itch
  - Dysuria/burning

### Trichomonas - Who to screen?

- No clear recommendations for asymptomatic screening either through PHAC or CDC
  - Data lacking on whether asymptomatic screening reduces any adverse health events or community spread
- May want to consider asymptomatic screening in the following groups:
  - Pregnant individuals
    - May be associated with premature rupture of the membranes, preterm birth and low birth weight. However, not known if treatment will improve pregnancy outcomes.
  - Those at risk for other STIs
    - Trichomonas infection associated with an increased risk of HIV acquisition and transmission
  - Females/trans males living with HIV
    - Trichomonas infection associated with increased risk for PID

#### Trichomonas Which tests should I do?

Test	Site	
NAAT	<ul> <li>Vaginal</li> <li>Endocervical</li> <li>Urine</li> </ul>	<ul> <li>Vaginal or endocervical preferred for female/trans male</li> <li>Urine should be first catch, not midstream</li> <li>Can use same sample for chlamydia or gonorrhea NAAT</li> </ul>
Culture	• Vaginal	<ul> <li>Can be combined with yeast, BV testing</li> </ul>

# Which swab should I use?

Aptima

OIOH



#### Hologic Aptima Unisex Swab

• Endocervical

(can use same swab if also testing for Chlamydia and Gonorrhea NAAT)

	Ontario 🕅	Ministry of Hea Laboratory Req Requisitioning C	quisition	tioner	Labo	orato	ny U	lse Oi	nly																			
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Note: Separate requisitions are required for cytology, Ontario Cervical Screening Program HPV and cytology tests, histology/pathology, ColonCancerCheck Fi lest, and tests performed for Public Health Laboratory.

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# Which swab should I use?



#### **DYNACARE AND LIFELABS**

#### **Hologic Aptima Multitest Swab**

• Vaginal

(can use the same swab if also testing for Chlamydia and Gonorrhea NAAT)

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Biochemistry Glucose Random HbA1C	Fasting	x	Hematology	x	Viral Hepatitis (check one only)
	Fasting	-			
HbA1C			CBC		Acute Hepatitis
			Prothrombin Time (INR)		Chronic Hepatitis
Creatinine (eGFR)			Immunology		Immune Status / Previous Exposure
Uric Acid			Pregnancy Test (Urine)	Τ	Specify: Hepatitis A
Sodium			Mononucleosis Screen	1	Hepatitis B
Potassium			Rubella	1	Hepatitis C     or order individual hepatitis tests in the
ALT			Prenatal: ABO, RhD, Antibody Screen	1	"Other Tests" section below
Alk. Phosphatase		1	(titre and ident. if positive)	P	rostate Specific Antigen (PSA)
Bilirubin			Repeat Prenatal Antibodies		Total PSA Free PSA
Albumin			Microbiology ID & Sensitivities	Spe	ecify one below:
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Albumin / Creatinine Ratio, Urine			Vaginal / Rectal – Group B Strep		Insured - Meets OHIP eligibility criteria:
Urinalysis (Chemical)		×	Chlamydia (specify source): vagina	1	osteopenia; osteoporosis; rickets; renal disease; malabsorption syndromes;
Neonatal Bilirubin:		X	GC (specify source): vagina	1_	medications affecting vitamin D metabolism
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Patient's 24 hr telephone no. ( )			Wound (specify source):	Tri	chomonas NAAT-vagina
Therapeutic Drug Monitoring:			Urine		
Name of Drug #1			Stool Culture		
Name of Drug #2		$\square$	Stool Ova & Parasites	$\square$	
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# Urine NAAT



#### DYNACARE AND LIFELABS

#### **Urine NAAT**

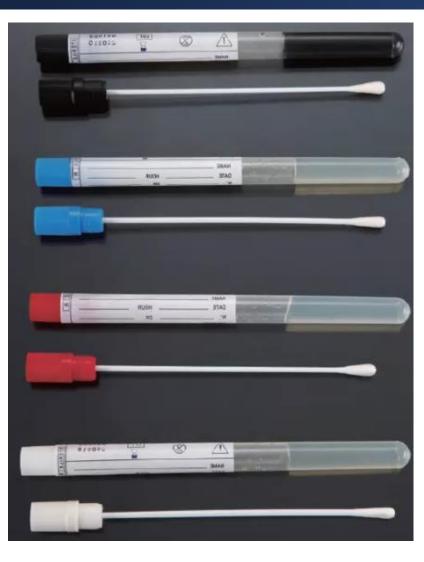
- First-catch urine (20 to 30 mL of initial urine stream)
- Can use same sample for Chlamydia and Gonorrhea NAAT)
- Vaginal or cervical NAAT preferred for female/trans male patients

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Note: Separate requisitions are required for cytology, Ontario Cervical Screening Program HPV and cytology tests, histology/pathology, ColonCancerCheck FI1 test, and tests performed for Public Health Laboratory.

х	Biochemistry		x	Hematology		x	Viral Hepatitis (check one only)
	Glucose Random	Fasting	$\square$	CBC		$\square$	Acute Hepatitis
	HbA1C			Prothrombin Time (INR	)	$\square$	Chronic Hepatitis
	Creatinine (eGFR)			Immunology			Immune Status / Previous Exposure
	Uric Acid			Pregnancy Test (Urine)			Specify: Hepatitis A
	Sodium			Mononucleosis Screen		1	Hepatitis B
	Potassium			Rubella		1	or order individual hepatitis tests in the
	ALT			Prenatal: ABO, RhD, Ar	ntibody Screen	1	"Other Tests" section below
	Alk. Phosphatase		1	(titre and ident. if position	(e)	F	Prostate Specific Antigen (PSA)
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	Neonatal Bilirubin:		X	GC (specify source): U	rine	]_	medications affecting vitamin D metabolism
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	Clinician/Practitioner's tel. no. ( )			Throat		C	Other Tests - one test per line
	Patient's 24 hr telephone no. ( )			Wound (specify source,	k:	Tri	ichomonas NAAT-urine
	Therapeutic Drug Monitoring:			Urine			
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# Which swab should I use?



#### DYNACARE AND LIFELABS

• Vaginal C&S (Yeast, BV, Trichomonas)

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Note: Separate requisitions are required for cytology, Ontario Cervical Screening Program HPV and cytology tests, histology/pathology, ColonCancerCheck Fi test, and tests performed for Public Health Laboratory.

tes	t, and tests performed for Public Heal	in Laboratory.					
х	Biochemistry		x	Hematology		х	Viral Hepatitis (check one only)
	Glucose Random	Fasting		CBC			Acute Hepatitis
	HbA1C			Prothrombin Time (INR)			Chronic Hepatitis
	Creatinine (eGFR)			Immunology			Immune Status / Previous Exposure
	Uric Acid			Pregnancy Test (Urine)			Specify: Hepatitis A
	Sodium			Mononucleosis Screen			Hepatitis B
	Potassium			Rubella			or order individual hepatitis tests in the
	ALT			Prenatal: ABO, RhD, Antib	ody Screen		"Other Tests" section below
	Alk. Phosphatase		1	(titre and ident. if positive)		P	rostate Specific Antigen (PSA)
	Bilirubin			Repeat Prenatal Antibodie	8		Total PSA Free PSA
	Albumin			Microbiology ID & Ser	nsitivities	Spe	cify one below:
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	Albumin / Creatinine Ratio, Urine			Vaginal / Rectal – Group B	3 Strep		insured - Meets OHIP eligibility criteria:
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	Child's Age: days	hours		Sputum		Ц	Uninsured - Patient responsible for payment
	Clinician/Practitioner's tel. no.( )			Throat		0	ther Tests - one test per line
	Patient's 24 hr telephone no. ( )			Wound (specify source):			
	Therapeutic Drug Monitoring:			Urine			
	Name of Drug #1			Stool Culture			
	Name of Drug #2			Stool Ova & Parasites			
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### **Trichomonas- Treatment**

	Treatment	Comments
Non-pregnant	Metronidazole 2 g PO in a single dose [A-I] or Metronidazole 500 mg PO BID for 7 days [A-I]	<ul> <li>Intravaginal metronidazole gel is not effective</li> </ul>
Pregnancy	Metronidazole 2 g PO in a single dose for symptom relief [A-I]. Alternate treatment: Metronidazole 500 mg PO BID for 7 days [A-I]	<ul> <li>It is not known whether treatment will improve pregnancy outcomes</li> </ul>

Note: Recommended to treat any current sexual partners with same treatment regimen

### Trichomonas – Follow up

- Repeat testing if symptoms persist
- No recommendations for test of cure (TOC) in Canada
  - CDC suggests retesting all women/trans men 3 months after treatment due to high rates of re-infection.
  - CDC does not recommend retesting in men/trans females (data is insufficient)

### Key take-aways

#### HSV

- Can be a devastating diagnosis- counselling is important
- Testing of lesions with HSV NAAT is the preferred method for diagnosis
- HSV serology not recommended for asymptomatic screening

#### Trichomonas

- STI-associated cause of vaginitis
- Test in those with symptoms using NAAT (cx, vagina, urine) or vagina C&S
- Treatment is mainly for symptom relief
  - Insufficient data on whether it changes health outcomes or community spread
- No clear guidelines for asymptomatic screening for Trichomonas
  - May want to consider in pregnant individuals, those at-risk for other STIs or in women/trans males living with HIV



- <u>Genital herpes guide: Key information and resources Canada.ca</u>
- <u>STI-associated syndromes guide: Vaginitis Canada.ca</u>
- <u>CDC- Trichomoniasis STI Treatment Guidelines</u>

#### **OCFP** supports for Mental Health, Addictions and Chronic Pain

Mental health, addictions and chronic pain are challenging conditions. Find information to support the care you give patients – in a way that also considers your wellbeing.



#### **Community of Practice**

Join upcoming sessions:

Income Benefit Programs for People Living on Low Income and in Poverty-Primary Care Providers' Role (June 19<sup>th</sup>)

Join

<u>Navigating the Complexities of</u> <u>Opioid Prescribing for Chronic</u> (June 25<sup>th</sup> )

Best Practices for Nicotine Cessation (July 23<sup>rd</sup>)



#### **Peer Connect Mentorship**

Receive tailored support to skillfully respond to mental health issues, address substance use disorders, and chronic pain challenges in your practice.

# Health Equity CoP



The OCFP, in partnership with the DFCM, has developed a new community of practice series, focused on enhancing care for marginalized or underserved populations; supporting family physicians in addressing the unique needs of their patients.

#### Launch: June 19

**Topic:** Income Benefit Programs for People Living on Low Income and in Poverty – Primary Care Providers' Role



# **RECENT SESSIONS**

March 7	Infectious Disease & HPV Cervical Screening Implementation	Dr. Daniel Warshafsky Dr. Jonathan Isenberg Dr. Rachel Kupets
March 21	Infectious Disease & Dermatology Treatments	Dr. Gerald Evans Dr. Juthika Thakur
April 4	Infectious Disease, Penicillin Allergy (De)labelling & Newcomer Care Resources	Dr. Daniel Warshafsky Dr. Mariam Hanna Dr. Vanessa Redditt
May 2	Infectious Disease and Management of STIs	Dr. Daniel Warshafsky Dr. Rachita Gurtu
May 23	Infectious Disease and Opportunities for Improving the Way We Work	Dr. Allison McGeer Dr. Tara Kiran

**Previous webinars, Self-Learning & Related Resources:** 

https://dfcm.utoronto.ca/changing-way-we-work-community-practice

# UPCOMING SESSIONS

Month	Date
June 2025	June 27
July 2025	July 18
September 2025	September 5

#### **SAVE THE DATE** Registration link will be emailed to you closer to the date

Family & Community Medicine UNIVERSITY OF TORONTO



### **Questions?**

Webinar recording and curated Q&A will be posted soon

Our next Community of Practice: June 27, 2025

Contact us: <u>ocfpcme@ocfp.on.ca</u>

*Visit*: <u>https://dfcm.utoronto.ca/past-changing-way-we-work-community-practice-sessions</u>

The Changing the Way we Work Community of Practice for Ontario Family Physicians is a one-credit-per-hour Group Learning program that has been certified for up to a total of 32 credits.

Post session survey will be emailed to you. Mainpro+ credits will be entered for you with the information you provided during registration.



