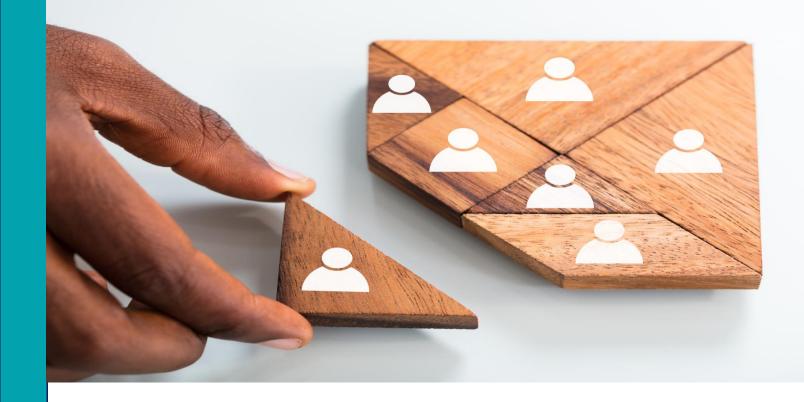
Changing the Way We Work Community of Practice for Ontario Family Physicians

May 2, 2025

Dr. Daniel Warshafsky Dr. Rachita Gurtu



Infectious Disease and Management of STIs





Infectious Disease and Management of STIs

Moderator:

• Dr. Eleanor Colledge, 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.

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.



Home
Vouth 10km Relay Yukon Your Way
Vouth 2023 Photo Gallery Reconciliation
Volunteers



Message from the Director

In 2023 we started our journey of Reconciliation. Here are some of the steps we have taken to improve our event:

- We renamed our event and worked with local artists and storytellers to give our race a stronger link to the traditional knowledge of this region.
- We obtained a land use permit from the First Nation Land's office as a portion of our race takes place in an area with significant environmental and cultural sensitivity.
- We offer complimentary race entries for any Indigenous registrant.
- A portion of race registration fees and donations help local groups to encourage more youth to walk, hike, ski, and run on Yukon trails fo health and wellness. In 2024 this support was directed towards the Kwanlin Koyotes.
- We have added a 4x 2.5km relay to encourage more youth to get involved in the sport of running.
- We shifted the purchasing of our event to include more Indigenous vendors for food, awards, and other event supplies.
- We work year round with our First Nations partners to bring you a unique racing experience.



You'll reach alpine where you've got a 360 view of Bonneville Lakes on one side, and Łu Zil Män o on the other. These are settlement lands, which means the land is owned and managed by the First Nations. The Bonneville Lakes were used by Indigenous people as a regular camp for fall hunting and spring fishing. Fish Lake, or Łu Zil Män in Southern Tutchone, is named for the whitefish that spawn here in the fall. The lake is significant to the Kwanlin Dun First Nation, whose people have gathered, hunted and held potlatches here since the end of the last ice age.

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

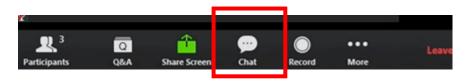
• All questions should be asked using the Q&A function at the bottom of your screen.



• 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?		
6			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

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

Public Health Update

Changing the Way We Work CoP

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

May 2, 2025

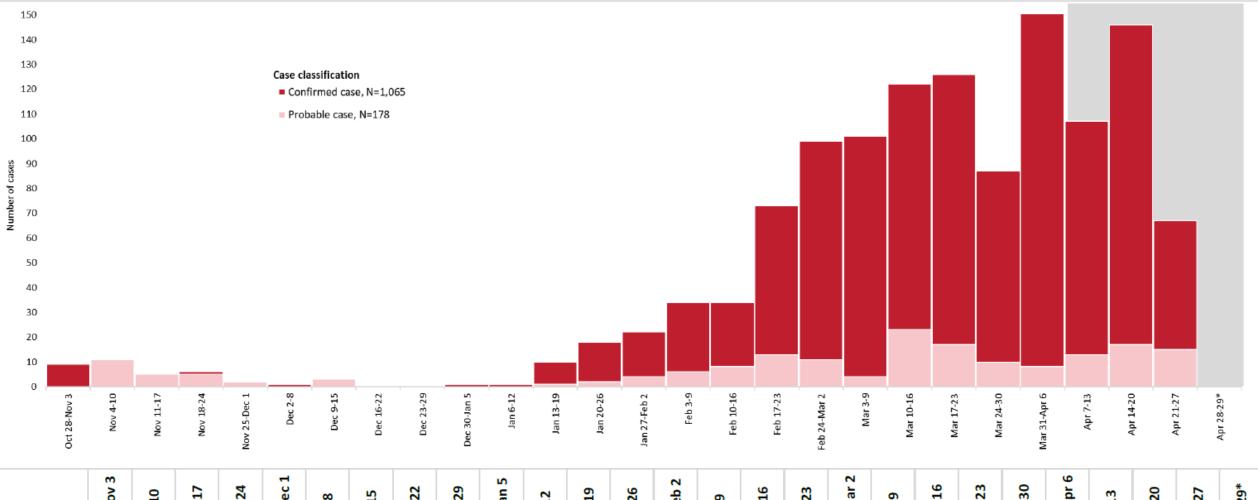


Ontario Measles Outbreak Update

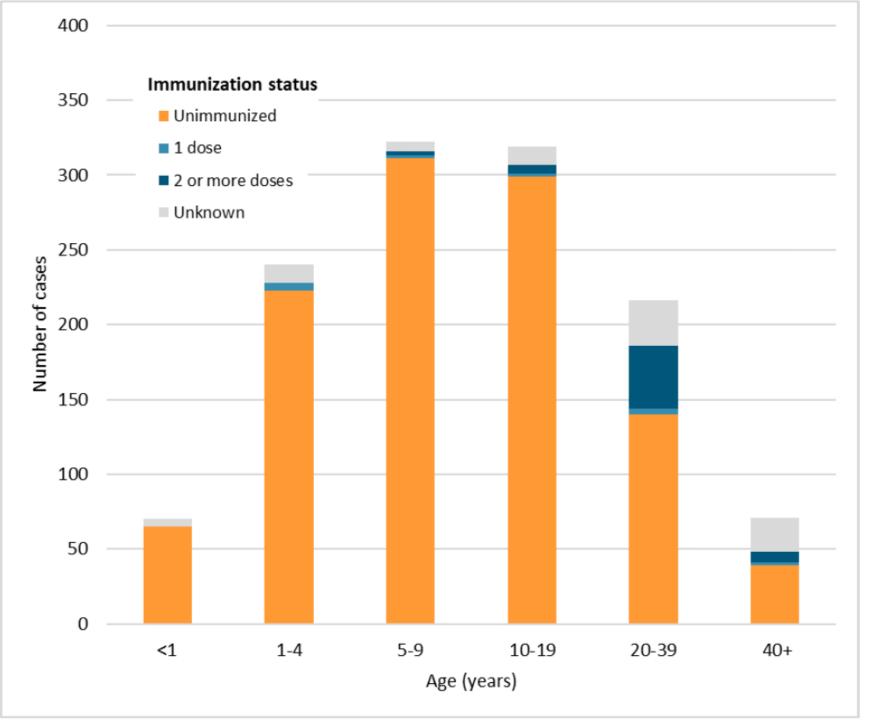
- Measles continues to spread in Ontario due to ongoing exposures and transmission among susceptible individuals
- As of April 29, 2025,
 - 1,243 cases in 17 public health units have been reported in association with this outbreak
 - This represents an increase of 223 cases and 2 new public health units since April 23, 2025
 - 84 hospitalizations
 - 8 ICU admissions
 - 25 pregnant persons
- For more information on place and dates of exposure to measles in Ontario, please visit Public Health Ontario's <u>website</u>



Number of Measles Outbreak Cases by Week of Rash Onset and Case Classification: Ontario, October 28, 2024 – April 29, 2025



Week of rash onset	Oct 28-Nov	Nov 4-10	Nov 11-17	Nov 18-24	Nov 25-Dec	Dec 2-8	Dec 9-15	Dec 16-22	Dec 23-29	Dec 30-Jan	Jan 6-12	Jan 13-19	Jan 20-26	Jan 27-Feb	Feb 3-9	Feb 10-16	Feb 17-23	Feb 24-Mai	Mar 3-9	Mar 10-1(Mar 17-23	Mar 24-3(Mar 31-Apı	Apr 7-13	Apr 14-20	Apr 21-27	Apr 28-29
Confirmed case	9	0	0	1	0	1	0	0	0	1	1	9	16	18	28	26	60	88	97	99	109	77	150	94	129	52	0
Probable case	0	11	5	5	2	0	3	0	0	0	0	1	2	4	6	8	13	11	4	23	17	10	8	13	17	15	0
Total	9	11	5	6	2	1	3	0	0	1	1	10	18	22	34	34	73	99	101	122	126	87	158	107	146	67	0



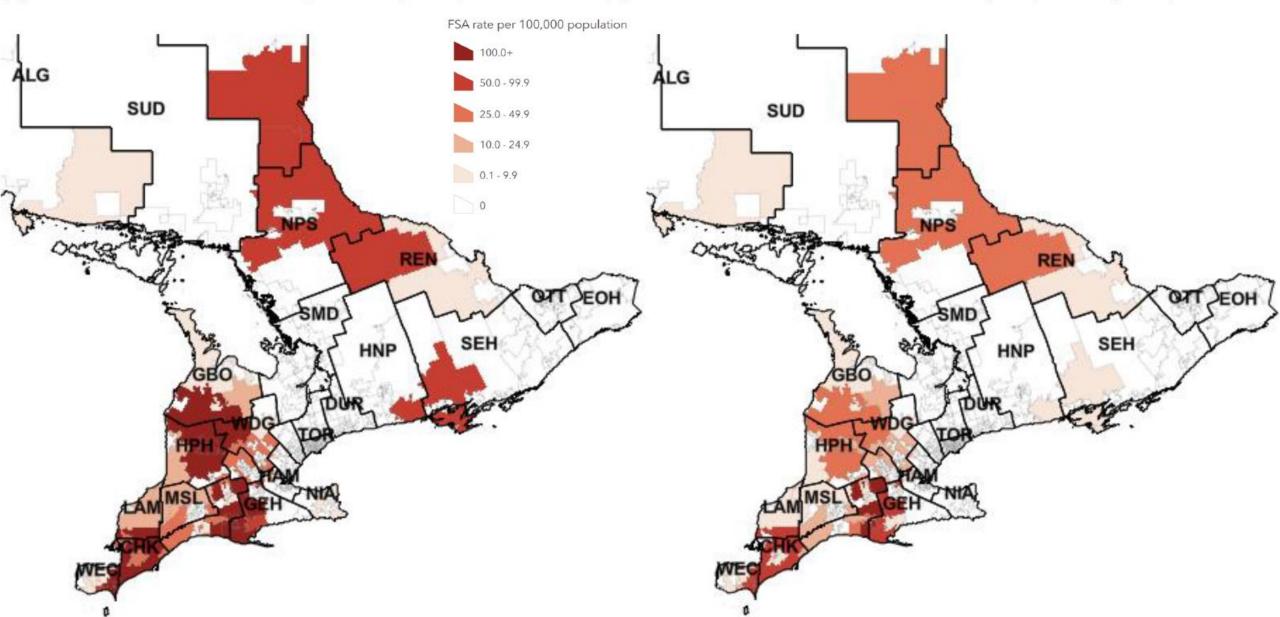
94% of cases in the current measles outbreak were unimmunized or had unknown immunization status



Figure 2: Geographic Distribution of the Rate of Measles Outbreak Cases Per 100,000 Population by Forward Sortation Area (FSA) Among Regions in Ontario with Cases:

(A) Cumulative cases: October 28, 2024 – April 29, 2025

(B) Cases with rash onset in the last 21 days: April 8 – April 29, 2025



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

- For all suspected measles cases, collect NP/throat swab and urine and serology (acute and convalescent)
 - Multiple specimen sites increases diagnostic sensitivity

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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.
<u>Measles serology</u> (diagnosis) *	Whole Blood (minimum 5mL whole blood or 1mL serum)	Blood, clotted serum separator tube (SST)	 Acute: within 7 days of rash onset Convalescent: 7 to 10 days after acute; preferably 10 to 30 days after acute

- 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)

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 and where the risk of exposure in the community is higher are recommended to receive:

Age group	Recommendation
Infants (six to 11 months)	Should receive one dose of the measles, mumps, rubella (MMR) vaccine. Note: Two 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

* Affected regions refer to southwestern Ontario, specifically Southwestern, Grand Erie and Huron Perth

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 Indicatio	n(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)

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
 - Additional PPE such as gloves, gown and eye protection may be added as required based on a point of care risk assessment (PCRA) per Routine Practices
- 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>

Testing for STIs in Primary Care

Dr. Rachita Gurtu MD, CCFP

Objectives

Gain comfort in taking a sexual health history

Review guidelines and practical considerations for chlamydia, gonorrhea, syphilis and herpes testing in a primary care setting

Review treatment guidelines

History taking

Approach should always be respectful, non-judgmental and one based on establishing trust

- Respecting privacy
- Explain the purpose of the questions
- Allow space for the patient to decline to answer if not comfortable

Avoid making assumptions

- Social and/or cultural context
- Sexuality & Gender
- Partners
- Consensual sex

The details are important and can be asked in a respectful way

- Who is/are their partner(s)
- What type of sex are they engaging in
- Are they using condoms

Chlamydia and Gonorrhea

Asymptomatic screening

Consider asymptomatic screening in the following groups:

- All sexually active persons under the age of 30 years
- Adults and adolescents with multiple partners or a new partner
- Pregnant individuals
- Populations or communities experiencing higher prevalence
 - \circ gbMSM
 - \circ HIV+
 - $\circ~$ Individuals who are or have been incarcerated
 - $\circ~$ Use of substances or access addiction services
 - Some Indigenous communities

Available tests

- Nucleic acid amplification tests (NAAT) most sensitive and preferred
 - Urine
 - Vaginal
 - Cervical
 - Pharynx
 - Rectal

Culture

- Gonorrhea not as sensitive, but provides antimicrobial susceptibilities, which is important in monitoring of antimicrobial resistance (AMR) patterns and trend
- Chlamydia available but not used routinely

Which tests should I do?

Female or *trans* male (anyone with a vagina)

Site	NAAT	GC Culture
Urogenital	Vaginal (preferred over urine or cervix) or Urine or Cervix	 Consider if any of the following: Has symptoms at that site Asymptomatic individual notified as a contact of gonorrhea
Pharynx	Consider if history of giving unprotected oral sex and any of the following:Has symptoms of sore throatNew or multiple partners	 When sexual abuse/sexual assault is suspected If the infection might have been acquired in countries or areas with high rates of AMR
Rectal	Consider if history of receiving anal sex	

Which tests should I do?

Male or *trans* female (anyone with a penis)

Site	NAAT	GC Culture
Urogenital	Urine (preferred) or Urethral	Consider if any of the following: • Has symptoms at that site
Pharynx	 Consider if history of performing unprotected oral sex and any of the following: Has symptoms gbMSM Multiple sexual partners or sex with a partner who is at high risk of infection 	 Asymptomatic individual notified as a contact of gonorrhea When sexual abuse/sexual assault is suspected If the infection might have been acquired in countries or areas with high rates of AMR
Rectal	Consider if history of receiving anal sex	



DYNACARE AND LIFELABS

Chlamydia and Gonorrhea NAAT

Hologic Aptima Multitest Swab

- Vaginal (including self-collected)
- Urethral
- *Pharynx
- *Rectal

*preferred for these sites over Unisex swab (Dynacare)

Laborat	r of Health lory Requisition loning Clinician / Practitioner	Labora	tary Use Only									
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Address												

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

x	Biochemistry			x	Hematology		x	Viral Hepatitis (check one only)
	Glucose Random		Fasting		CBC			Acute Hepatitis
	HbA1C				Prothrombin Time (N	R)		Chronic Hepetitis
	Creatinine (sGFR)				Immunology			Immune Status / Previous Exposure
	Uric Acid				Pregnancy Test (Urin	e)	1	Specify: Hepstitis A
	Sodium				Mononucleosis Scree	m		Hepatitis B Hepatitis C
	Potassium				Rubella		11	or order individual hepstitis tests in the
	ALT				Prenatal: ABO, RhD,	Antibody Screen	Ĩ.	"Other Tests" section below
	Alk. Phosphatase				(titre and ident. if posi	itivo)	P	rostate Specific Antigen (PSA)
	Bilirubin				Repeat Prenatal Antit	odies	10	Total PSA
	Albumin				Microbiology ID 8 (if warranted)	Sensitivities	100	icify one below: insured – Meets OHIP eligibility criteria
	Lipid Assessment (includes Choleste calculated LDL-C & Chol/HDL-C ratio	rol, HDL c individ	L-C, Trighycerides, fusi lipid tests may		Cervical		100	Uninsured - Screening: Patient responsible for payment
	be ordered in the "Other Tests" section	n of this	s form)		Vaginal		V	itamin D (25-Hydroxy)
	Albumin / Creatinine Ratio, Urine				Vaginal / Rectal - Gro	oup 8 Strep	1000	risured - Meets OHIP eligibility criteria:
	Urinalysis (Chemical)			~	Chlamydia (specify so	ource): vagina	-	osteopenia; osteoporosis; rickets;
	Neonatal Bilirubin:			x	GC (specify source):	vagina	1	renal disease; malabsorption syndromes; medications affecting vitamin D metabolism
	Child's Age: days		hours		Sputum			Uninsured - Patient responsible for payment
	Clinician/Practitioner's tel. no. ()	010000		Throat		0	ther Tests - one test per line
	Patient's 24 hr telephone no. (and the second			Wound (specify source	:o):		
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	Name of Drug #1	rug #1			Stool Culture			
	Name of Drug #2				Stool Ova & Parasited			
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DYNACARE AND LIFELABS

Chlamydia and Gonorrhea NAAT

Hologic Aptima Unisex Swab

- Endocervical
- Urethral
- Pharynx (Dynacare only)
- Rectal (Dynacare only)

Ontario 😵	Ministry of Healt Laboratory Requ Requisitioning Clin	itioner	Labora	tory Us	e Only																	
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Note: Separate requisitions are required for cytology, Ontario Cervical Screening Program HPV and cytology tests, histology/pathology, ColonCancerCheck Fil test, and tests performed for Public Health Laboratory.

20	, and tests performed for Public Healt	in Laboratory.					
5	Biochemistry		x	Hematology		х	Viral Hepatitis (check one only)
	Glucose Random	Fasting		CBC			Acute Hepatitis
	HbA1C			Prothrombin Time (INR))		Chronic Hepetitis
	Creatinine (eGFR)			Immunology			Immune Status / Previous Exposure
	Uric Acid			Pregnancy Test (Urine)			Specify: Hepatitis A
	Sodium			Mononucleosis Screen		1	Hepatitis B Hepatitis C
	Potassium			Rubella		1	or order individual hepatitis tests in the
	ALT			Prenatal: ABO, RhD, Ar	ntibody Screen	1	"Other Tests" section below
	Alk. Phosphatase			(titre and ident. if positiv	/e)	P	rostate Specific Antigen (PSA)
	Bilirubin			Repeat Prenatal Antibo	dies		Total PSA Free PSA
	Albumin			Microbiology ID & S	Sensitivities	Spe	acify one below:
	Lipid Assessment (includes Cholesterol, H	DL-C Triphenetidae	1	(if warranted)			nsured – Meets OHIP eligibility ortteria
	calculated LDL-C & Chol/HDL-C ratio; ind	ividual lipid tests may		Cervica			Uninsured - Screening: Patient responsible for payment
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	Urinalysis (Chemical)		×	Chlamydia (specify sou	rce): cervix		osteopenia; osteoporosis; rickets; renal disease; malabsorption syndromes;
	Neonatal Bilirubin:		X	GC (specify source): C	ervix		medications affecting vitamin D metabolism
	Child's Age: days	hours		Sputum			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			
	Time Collected #1 hr	#2 hr.		Other Swabs / Pus (spe	cify source):		
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PHO LAB

Chlamydia and Gonorrhea NAAT

Woven swab (clinician or self-collected):

- Vaginal •
- Rectal ٠
- Pharyngeal •

Flocked swab:

Endocervical •

Note: Urethral and penile meatal swabs are not included and will not be accepted.

Submitter / He	alth Care Prov	vider (HCP)	nformation	Patient Info	ormation		
Licence No.;	Lab / Hospital o	or Facility Name	:	Health Card No.	:		
						Se	x: Male
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Address:				Medical Record	No.:		Female
		Postal		Last Name (per health card):			
City:		Code:	Province:	First Name (per health card):			
Tel:		Fax:		Address:		Posta	
Copy to Other La	b / Health Unit / A	uthorized Healt	h Care Provider (HCP)			Code:	
Licence No.:	Other Lab / Hea	alth Unit / Facilit	y Name:	City:		Tel:	
				Investigation / PHO or Health	Outbreak No. from Unit (if applicable):		
HCP Full Name:					nformation		
Address:				A Date Colle	ected 2024-03-2	20 Submitter	
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Other Specify):					Acute Infection (HBsAg + total and	HBc Scr	-Chemotherapy eening (anti-HBs +
	-1/->				+ IgM if total is por	sitive) HBs	Ag + total anti-HBc)
Relevant Trav	ei(s)				Current / Past In		

General Test Requisition

The personal health information is collected under the authority of the Personal Health information Protection Act, 2004, s.36 (1)(c)(ii) for the purposes specified in the Ontario Agency for Nealth Protection and Promotion Act, 2007, s. 1 inducing clinical laboratory testing and public health purposes. If you have questions about the collection of this personal health information please contact the PHO's Laboratory Customer Service at 416-235-6558 or toll fire 1-877-604-4567. F-SD-SCG-1000, version 0004-2 (August 2024)



Ontario 😚



PHO LAB

Gonorrhea culture

- Endocervical
- Urethral
- Pharyngeal
- Rectal

Note: Store and ship specimens at room temperature. Culture specimens must be received at the testing laboratory within 72 hours of collection.

General Test Requisition



Ontario 🕅

ALL sections of the form must be completed by <u>authorized</u> health care providers for each specimen submitted, or testing may be delayed or cancelle 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: publicheal thoritario-ca/requisitions

Submitter / Hea	alth Care Provide	er (HCP) Info	rmation		Patient In	formatio	n			
Licence No.;	Lab / Hospital or Fa	cility Name:			Health Card N	o.:				
					Date of Birth ()	vvvv-mm-dd	0:	s	ex:	Male
HCP Full Name:					Medical Recor		a.	_		Female
Address:					Last Name					
City:	Pos		Province:		(per health card)	c				
	Co				First Name (per health card)	c				
Tel:	Pa	BX:			Address:			Posta Code		
	/ Health Unit / Autho			(HCP)	City:			Tel:		
Licence No.:	Other Lab / Health	Unit / Facility Na	ime:		Investigation	/ Outbreak	No. from			
HCP Full					PHO or Healt	h Unit (if ap	plicable):			
Name:					Specimen	Informa	tion			
Address:					Date Col (yyyy-mr)	m-dd): 20	25-03-24	Submitte Lab No.:		
City:	Pos		Province:		Whole B	lood	Serum		Plas	sma
Tel:	Fi	ax:			Bone Ma	wom	Cerebrospir Fluid (CSF)		Nas	opharyngea ab (NPS)
Patient Setting	I				Orophan / Throat	yngeal Sweb	Sputum		Bro	nchoalveola age (BAL)
Clinic / Community	ER (Not Admi Not Yet Deten		ER (Ad	mitted)	Endocer	_	Vaginal Swa	ab	_	thral Swab
Inpatient (Non-ICU)	ICU / CCU	,	Congre		Urine	Ī	Rectal Swa	ь	Fae	ces
	tion(s) / Criteria				Other (Specify	twne				
Diagnosis	Screening	Immune	Follow	up /	AND body loca	ation):				
Pregnancy /		Status Post-	Conval	escent	Test(s) Re	quested				
Perinatal	Immunity	mortem			Enter each as	say as per t	he publichealtho	ontario cal	testdire	actory:
Other (Specify):					¹ Gonorr	hea cult	ure			
Signs / Sympto	oms				2.					
No Signs / Symptoms	Onset Date (yyyy-mm-dd)	n/a			3					
 Symptoms 	Fever	Rash	STI		4.					
Gastrointestinal		Hepatitis	Meni	ingitis /	5.					
Other	Respiratory	Hepaulus	Ence	phalitis	6.					
(Specify):					For routine he	patitis A, B	or C serology, c	complete t	his sec	tion instead
Relevant Expo	sure(s)						ne Status	Ac	ute Infe	ection
None / Not Applicable	Most Recent Date (yyy-mm-dd):	8			Hepatitis A	(HAVI		sy	AV IgM, mptoms	info)
Occ	cupational Exposure / edlestick Injury (Specif	y): So	urce E	xposed	Hepatitis B	(anti-H	ne Status IBs)	(H	BsAg + t	nfection total anti-HBc)
Other (Specify):						(HBsA	Infection g + total anti-HBc	Sc	reening	notherapy g (anti-HBs + otal anti-HBc)
Relevant Trave	l(s)						iftotal is positive) nt / Past Infectio			
None / Not Most Recent Date Applicable (yyyy-mm-dd):			Hepatitis C		nune status test fo					
Travel Detais:										
The personal health info	rmation is collected unde	r the authority of th	ne Personal He	aith Infor	metion Protection A	of 2004, s.36	5 (1)(c)(ii)			

The personal health information is collected under the authority of the Personal Health Information Protection Act, 2004, s.35 (1)(c)(ii) for the purposes specified in the Ontario/Agency for / health Protection and Promotion Act, 2007, s.1 including clinical laboratory testing and public health purposes. If you have guestions about the collection of this personal health information please contact the PHO's Laboratory Customer Service at 416-235-6556 or to lifere 1477-604-1567. F-SD-SCG-1000, version 004-2 (August 2024).

Chlamydia treatment

Uncomplicated gonorrhea infections	Treatment
Preferred treatment	Doxycycline 100 mg PO BID for 7 days (contraindicated in pregnancy) OR Azithromycin 1 g PO in a single dose Note: Azithromycin may be preferred when poor compliance is anticipated.
Alternative treatment for anogenital infections	Levofloxacin 500 mg PO once a day for 7 days
Alternative for Pregnant or lactating individuals	Amoxicillin 500 mg PO TID for 7 days OR Erythromycin 2 g/day PO in divided doses for 7 days OR Erythromycin 1g/day PO in divided doses for 14 days

Gonorrhea treatment

Uncomplicated gonorrhea infections	Treatment	
Preferred treatment	Ceftriaxone 500 mg IM as a single dose (monotherapy)	
Alternative treatment for anogenital infections	Cefixime 800 mg PO in a single dose PLUS Doxycycline 100 mg PO BID x 7 days	
Alternative treatment for pharyngeal infections	Cefixime 800 mg PO in a single dose PLUS Azithromycin 1 g PO in a single dose	N re
Cephalosporin allergy or resistance or severe non-IgE-mediated reaction to penicillins	Azithromycin 2 g PO in a single dose PLUS Gentamicin 240 mg IM in a single dose	•
Contraindications to macrolides and cephalosporins	Gentamicin 240 mg IM in a single dose PLUS Doxycycline 100 mg orally twice daily for 7 days	

Note: Alternative treatment regimens are not recommended in pregnancy.

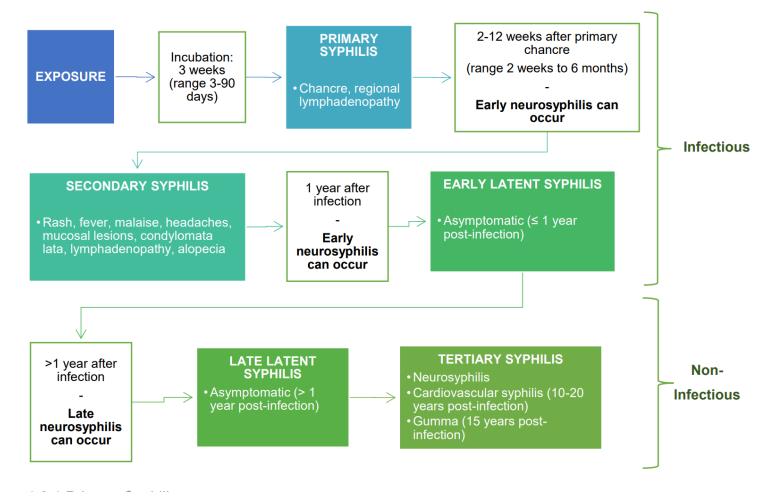
- In cases of cephalosporin allergy or other contraindications, consult with an infectious disease specialist.
- Doxycycline is contraindicated in pregnant and lactating individuals.
- Combination therapy containing gentamycin is not recommended in pregnancy

Test of Cure (TOC)

Infection	ТОС	Follow up		
	NAAT	GC Culture	screening	
Chlamydia	 Recommended 3-4 weeks after completion of treatment if: compliance to treatment is suboptimal an alternative treatment regimen was used persisting signs of symptoms post treatment the person is prepubertal or pregnant 	n/a	3 months	
Gonorrhea	 Recommended for all positive cases 3-4 weeks after treatment 	 If need to complete TOC earlier than 3 weeks, then perform gonorrhea culture only (too early for NAAT) If treatment failure is suspected, and more than 3 weeks after treatment – perform both NAAT and culture 	6 months	

Syphilis

Figure 1. Summary of the natural history of untreated syphilis and its associated clinical manifestations



Syphilis Stages

Source: Syphilis in Canada: Technical report on epidemiological trends, determinants and interventions

Primary Syphilis



Source: <u>https://thestdproject.wpenginepowered.com/wp-content/uploads/2012/07/std_pictures_of_syphilis_pictures.pdf</u>

Secondary Syphilis





Multiple erosions (mucous patches) are present on the tongue in this patient with secondary syphilis.

Source: UpToDate- Syphilis: Epidemiology, pathophysiology, and clinical manifestations in patients without HIV

Secondary Syphilis





Sources:

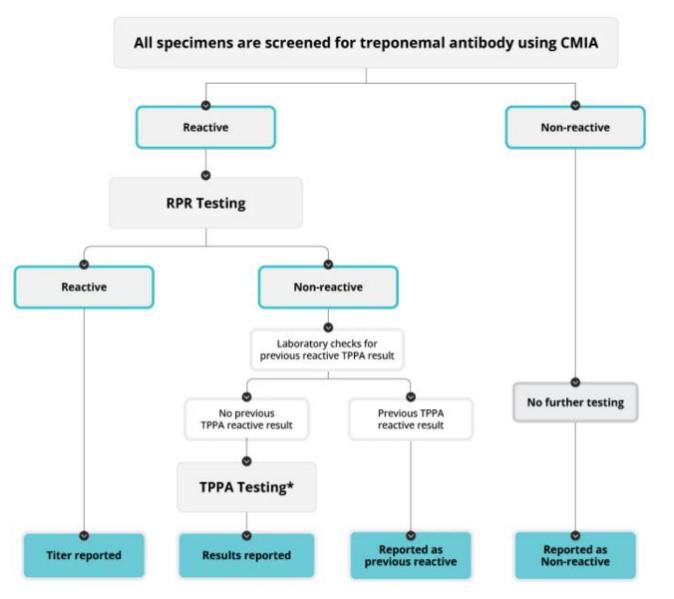
UpToDate- Syphilis: Epidemiology, pathophysiology, and clinical manifestations in patients without HIV https://ijdvl.com/scrotal-plaques-as-a-predominant-presentation-in-a-case-of-secondary-syphilis/

Serological tests for Syphilis

CATEGORY	TEST	COMMENTS
Treponemal	Treponema pallidum particle agglutination [TP-PA] Chemiluminescent immunoassay [CLIA]	 Detect antibodies specific to <i>T. pallidum</i> Usually will remain reactive for the remainder of patient's life, regardless of adequate treatment or disease activity.
Non-treponemal	Rapid plasma regain [RPR]	 Not specific for <i>T. pallidum</i> False-positive results can be associated with multiple medical conditions and factors unrelated to syphilis, including other infections (e.g., HIV), autoimmune conditions, vaccinations, injecting drug use, pregnancy, and older age Titers might correlate with disease activity and are used for monitoring treatment response

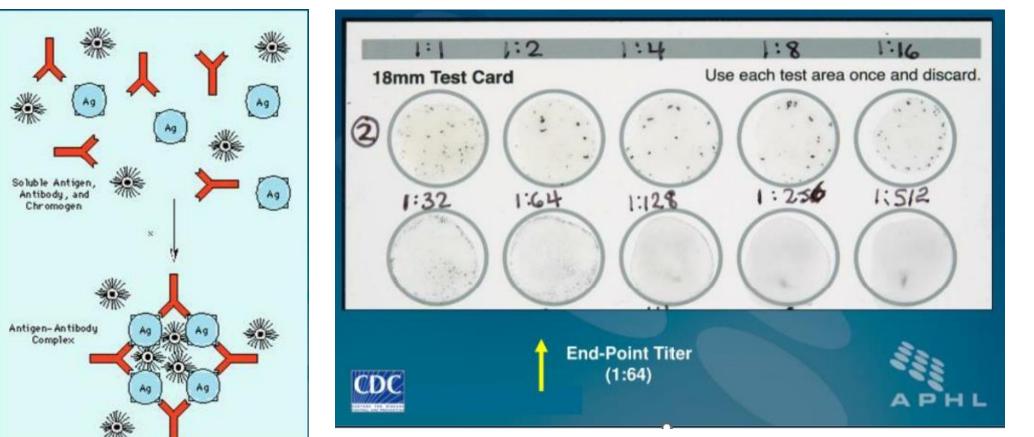
PHO Syphilis serology testing algorithm

PHO follows a reverse algorithm for syphilis testing



*For infants ≤18 months, TPPA testing is completed regardless of RPR result.

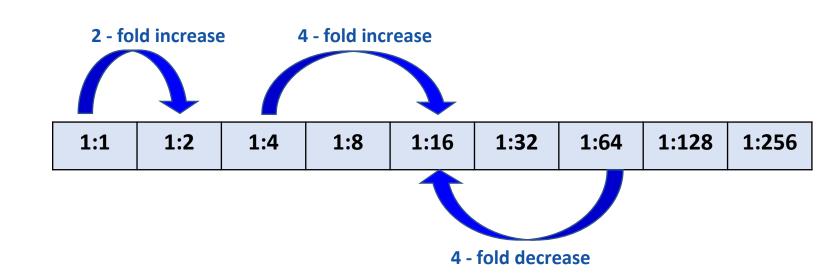
Rapid Plasma Reagin (RPR)



Sources:

- <u>Rapid Plasma Reagin StatPearls NCBI Bookshelf (nih.gov)</u>
- RAPID PLASMA REAGIN 18-MM CIRCLE CARD TEST (cdc.gov)
- <u>PPT STI Update PowerPoint Presentation, free download -</u> ID:960395 (slideserve.com)

Rapid Plasma Reagin (RPR)



*To be considered a significant rise or decrease, there should be at least a 4-fold change.

Positive results

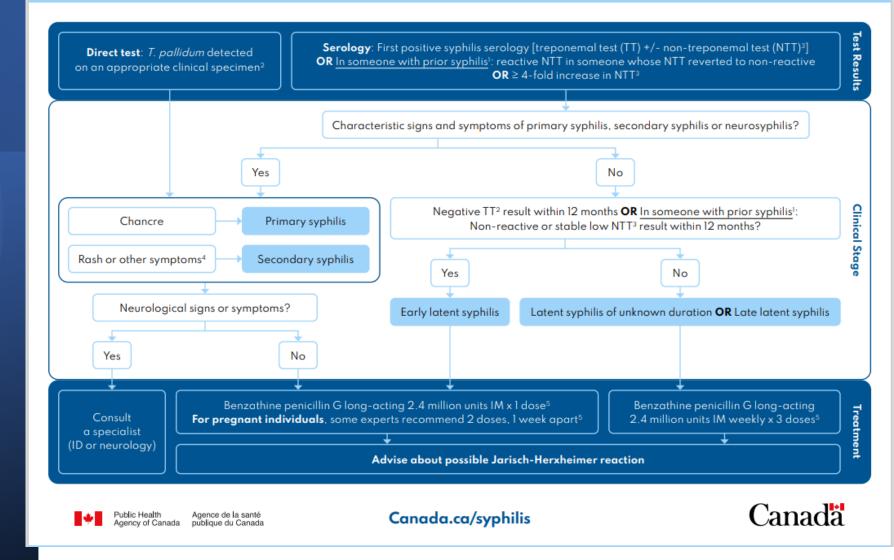
Syphilis Antibody Screen Syphilis RPR Quantitation Syphilis Serology Interpretation

Reactive Reactive 1:8 Consistent with recent or prior syphilis infection.

Syphilis TP.PA Syphilis Serology Interpretation	Reactive Consistent with recent or prior syphills infection.
Syphilis RPR Screen	Non-Reactive
Syphilis Antibody Screen	Reactive

Staging algorithm

Simplified algorithm for clinical syphilis staging and treatment in adolescents and adults



Source: syphilis-staging-algorithm-eng.pdf

March 2024

Treatment and follow up

Stage	1 st Line Treatment	Penicillin allergy	Serology follow up
Primary/Secondary/ Early Latent	Benzathine penicillin G-LA 2.4 million units IM x 1 dose	Doxycycline 100 mg PO BID x 14 days	3, 6, 12 months
Late Latent	Benzathine penicillin G-LA 2.4 million units IM qweekly x 3 doses	Doxycycline 100 mg PO BID x 28 days	12, 24 months

Genital Herpes

Genital Herpes- Counselling

- Can be a devastating diagnosis for many people due to how infection is perceived and the attached social stigma
- Genital herpes is a common condition
 - One analysis estimates that 14% of people between 14 and 59 years of age in Canada have HSV-2
- HSV-1 and HSV-2 can live in your body for a long time undetected or unrecognized
- Genital herpes is a recurrent, chronic infection; however, it is a manageable condition
- Most people who have genital herpes don't know they have the infection because they have mild, short-lived or no symptoms, or they think the symptoms are due to another condition (e.g., yeast infection, boils, bug bites, friction burns).
- Condoms, if used consistently and correctly will reduce, but will not eliminate the risk of HSV transmission or acquisition.

Genital Herpes- Clinical manifestation

Primary Episode		Recurrences
	• Usually have more severe and systemic symptoms	Milder symptoms than with primary episode
	• Extensive, painful, bilateral vesiculo-ulcerative genital or anal lesions (may involve the exocervix)	 Prodromal symptoms lasting 1-2 days (43-53% of cases) including focal burning, itching, tingling,
	• Systemic symptoms: fever, malaise, myalgia and	hyperesthesia or dysesthesia
	headache (about 67% of cases)	Unilateral, localized small erythematous patch,
	Tender inguinal lymphadenopathy (about 80% of	painful genital vesicles and ulcers
	cases)	 Systemic symptoms (5–12% of cases)
	Complications include meningitis (16-26%),	
	extragenital lesions (10-28%)	

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 / Symp	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	 Acyclovir 200 mg PO five times per day for 5-10 days [A-I]- ideally within 7 days after symptom onset or Famciclovir 250 mg PO TID for 5 days [A-I]- ideally within 5 days after symptom onset or Valacyclovir 1000 mg PO BID for 10 days [A-I]- ideally within 3 days after symptom onset 	 Provide antiviral treatment to those experiencing a first episode of genital herpes unless all lesions have already healed.
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]	 Prompt initiation of episodic antiviral therapy at the onset of prodromal symptoms may shorten the severity and duration of lesions
Suppressive Treatment	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] or 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)	 Reduces the length, frequency and severity of recurrences, asymptomatic viral shedding and transmission

Genital Herpes-Treatment (in pregnancy)

Treatment	Instructions for use	Comments
Primary episode	Acyclovir 200 mg PO QID for 5-10 days [A-I]	 Caesarean delivery can reduce the risk of vertical transmission Caesarean delivery is recommended in the case of newly acquired genital HSV in the third trimester
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]	 Shown to be effective in reducing the risk of symptomatic recurrences and asymptomatic viral shedding at the time of delivery and the need for Caesarean section Suppressive therapy should be initiated at 36 weeks and continued until delivery for anyone with a history of HSV-2 and for those who had a recurrence of genital herpes within the previous year. Caesarean section is not necessary unless genital lesions are present during labour

Background: Primary Care Action Team



In January, \$1.4B in new funding announced by PCAT and provincial government. The announcement included commitments to:

- Expand team-based primary care
- Enhance digital tools for providers and patients
- Reduce administrative burden
- Modernize information sharing
- Improve the referral process.



Interprofessional Primary Care Team Expansion



- First round of the proposal process to expand interprofessional primary care teams under PCAT comes to a close today.
- Creating/expanding up to 80 teams in 125 postal codes with lower attachment rates.
- Following last month's announcement, OCFP met with Dr. Philpott.
- We welcome this opportunity for more family physicians to join or form new interprofessional teams.
- Next round of proposals: expected September 2025.

OCFP Focus



Participate in OCFP's upcoming member survey, focused on:

- Innovative team-based approaches to meet the needs of FPs and your patients.
- And how we can best help you navigate through, and participate in, system changes.

New Investments In Northern Ontario





- Part of arbitrated awards under the Year 1 targeted physician investments in the 2024-2028 Physician Services Agreement negotiated between the OMA and MOH.
- The OCFP has long championed the creation a Coordination Centre for Northern Ontario and we look forward to learning more about what this will entail.

OCFP Leadership Academy for Family Physicians

SPOTS AVAILABLE!

Modules	Dates	
Module 1 (In-Person)	 June 12-14, 2025 (2.5 days) June 12 (evening session): 4:30 – 8 p.m. June 13-14 (full day sessions): 8:30 a.m. – 4:30 p.m. 	
Online Module (half day session)	 September 18, October 23, November 19, December 17, January 21 and February 19 8:30 a.m. – noon. 	
Module 2 (In-Person)	 April 16-18, 2026 (2.5 days) April 16 (evening session): 4:30 – 8 p.m. April 17-18 (full day sessions): 8:30 a.m. – 4:30 p.m. 	

- Hosted in Toronto at Rotman
- \$5000 + HST and travel expenses
- Great opportunity if you have funding available from your PCN/OHT
- Open to family physicians and OHT/PCN administrative leaders

Interested family physicians can contact Leigh Anne Butler: <u>labutler@ocfp.on.ca</u> to confirm your participation by May 13!





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:

Supporting patients with ADHD and comorbidities (May 28th) <u>Navigating the Complexities of</u> <u>Opioid Prescribing for Chronic</u> (June 25th)



Peer Connect Mentorship

Join

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

RECENT SESSIONS

January 17	Infectious Disease & Gender Affirming Care	Dr. Daniel Warshafsky Dr. Tehmina Ahmad
February 21	Infectious Disease & Navigating Ontario's Disability Support Program	Dr. Alon Vaisman Dr. Mohamed Alarakhia Norma English
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

Previous webinars & related resources:

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

UPCOMING SESSIONS

Month	Date
May 2025	May 23
June 2025	June 6 June 27
July 2025	July 18

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

Our next Community of Practice: May 23, 2025

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

Visit: <u>https://www.ontariofamilyphysicians.ca/tools-resources/covid-19-</u> <u>resources</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.



