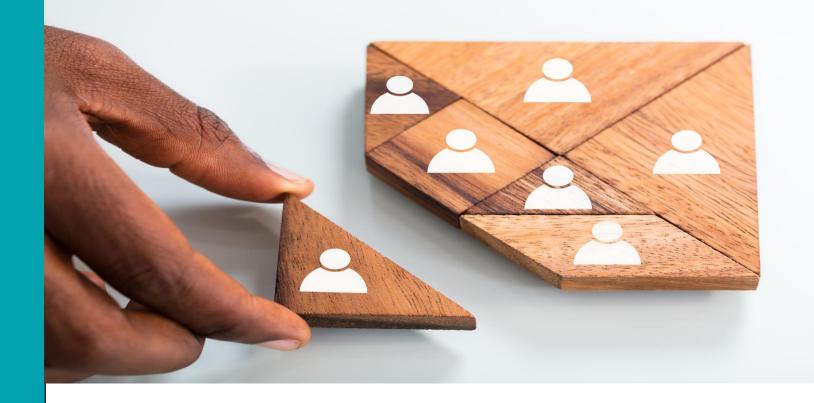
Changing the
Way We Work
Community of
Practice for Ontario
Family Physicians

January 17, 2025

Dr. Daniel Warshafsky Dr. Tehmina Ahmad



Infectious Disease & Gender Affirming Care





Infectious Disease & Gender Affirming Care

Moderator:

 Dr. Ali Damji, Division Head, Primary Care, Trillium Health Partners and Family Physician, Credit Valley Family Health Team, Mississauga, ON

Panelists:

- Dr. Daniel Warshafsky, Toronto, ON
- Dr. Tehmina Ahmad, Toronto, ON

Host:

Dr. Jobin Varughese, Brampton, ON

The Changing the Way We Work Community of Practice for Ontario Family Physicians is a one-credit-perhour 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.

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:



Dr. Daniel Warshafsky – PanelistAssociate Chief Medical Officer of Health at the Office of the Chief Medical Officer of Health



Dr. Tehmina Ahmad – PanelistMedical Consultant – Endocrinology & Metabolism, Toronto Western Hospital, UHN

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. Tehmina Ahmad
- Relationships with financial sponsors:
 - Grants/Research Support: N/A
 - Speakers Bureau/Honoraria: Shoppers Drug Mart (hired consultant for Transgender Module development for Pharmacists), Ontario College of Family Physicians
 - 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)

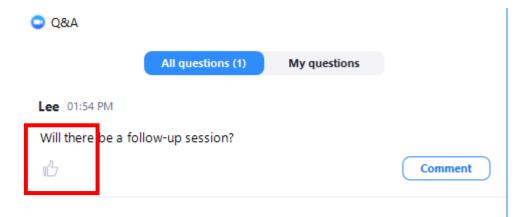
- Name: Dr. Ali Damji
- Relationships with financial sponsors:
 - Grants/Research Support: N/A
 - Speakers Bureau/Honoraria: Ontario Medical Association Section of General & Family Practice, Trillium Health Partners, Canadian Mental Health Association Peel Dufferin, Center for Effective Practice, GSK
 - Advisory boards: Medical Post Advisory Board, Foundation for Advancing Family Medicine, Center for Effective Practice
 - Others: N/A

How to Participate

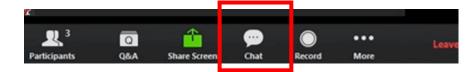
All questions should be asked using the Q&A function at the bottom of your screen.



• 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.



Please use the chat box for networking purposes only.





Dr. Daniel Warshafsky – PanelistAssociate Chief Medical Officer of Health at the Office of the Chief Medical Officer of Health



Dr. Tehmina Ahmad – PanelistMedical Consultant – Endocrinology & Metabolism, Toronto Western Hospital, UHN

Respiratory Season Updates

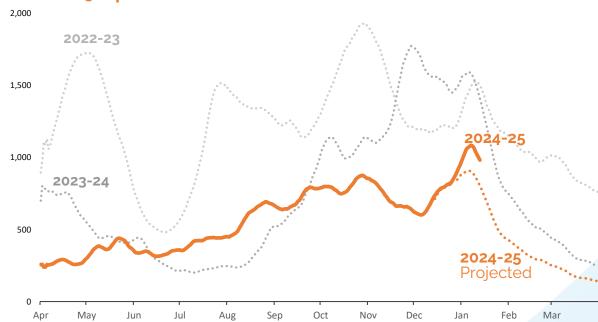
Office of the Chief Medical Officer of Health January 2025



Ontario 2024-25 respiratory season hospitalizations to date vs. previous years

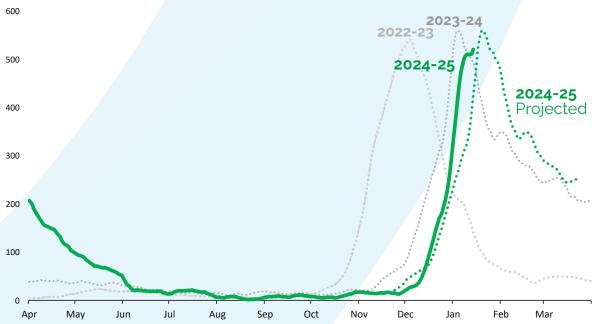
- RSV total hospitalizations have surpassed peak levels in 2022-23 and 2023-24 driven by seniors, although seem to be plateauing. Pediatric RSV hospitalizations peaked in mid-December and are slowly declining.
- Influenza hospitalizations are increasing at a steeper rate than the previous 2 years, currently running at about 1 week behind last year's wave. Hospitalizations are expected to be nearing peak levels.
- COVID-19 hospitalizations had expected post-holiday surge, but are decreasing already and at lower levels than prior years.

COVID-19 inpatients





Influenza inpatients

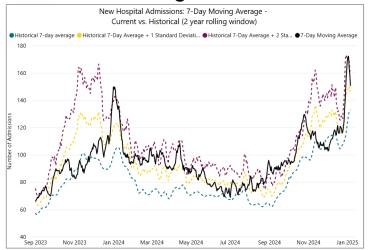


11

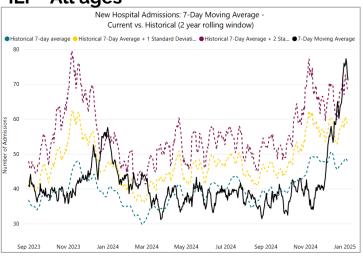
Data: 19 hospital census data as of January 13

ACES hospital admissions tracker

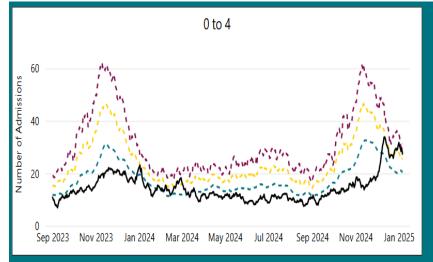
Pneumonia - All ages

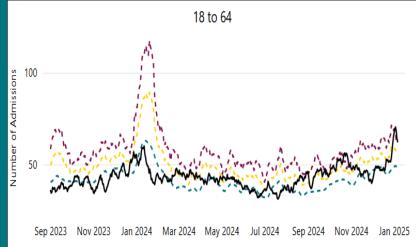


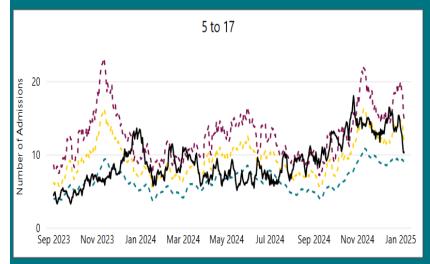
ILI - All ages

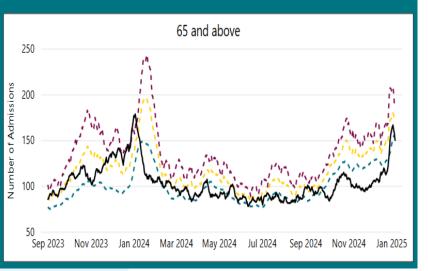


Pneumonia, ILI and COVID-19 - By age group





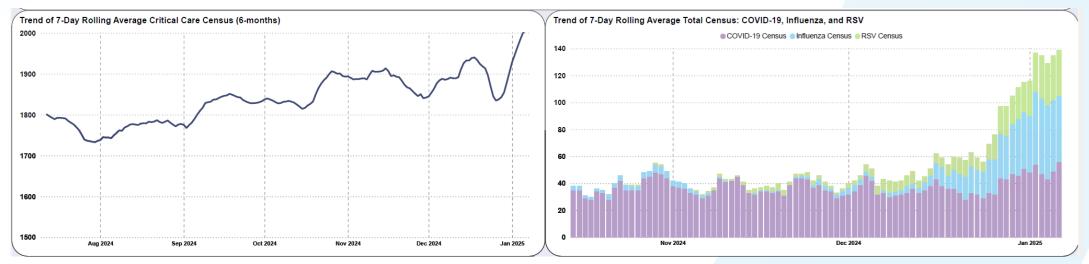




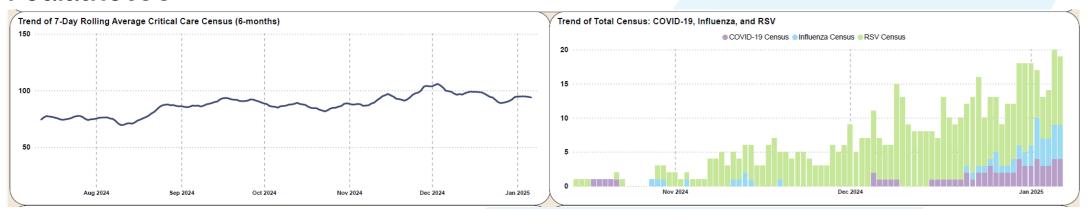
Data: KFLA PHU ACES surveillance data as of January 7

ICU census

Adult ICU

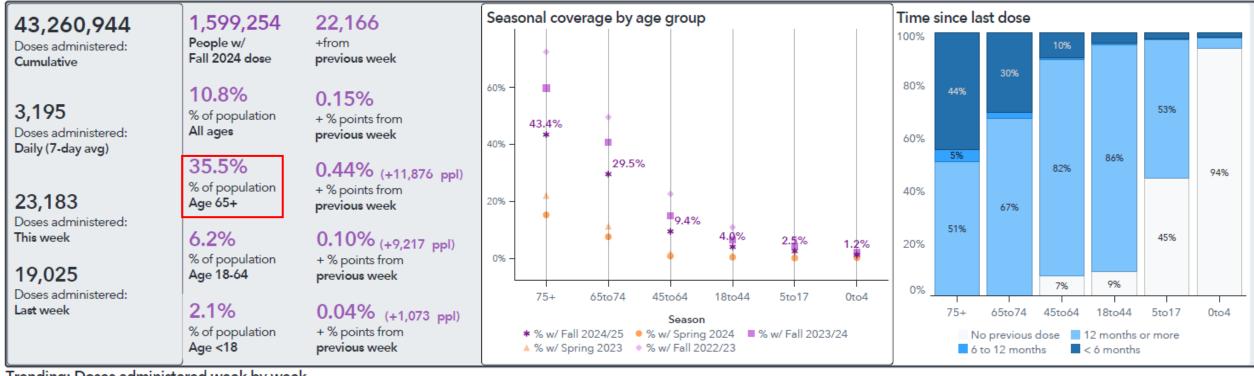


Pediatric ICU

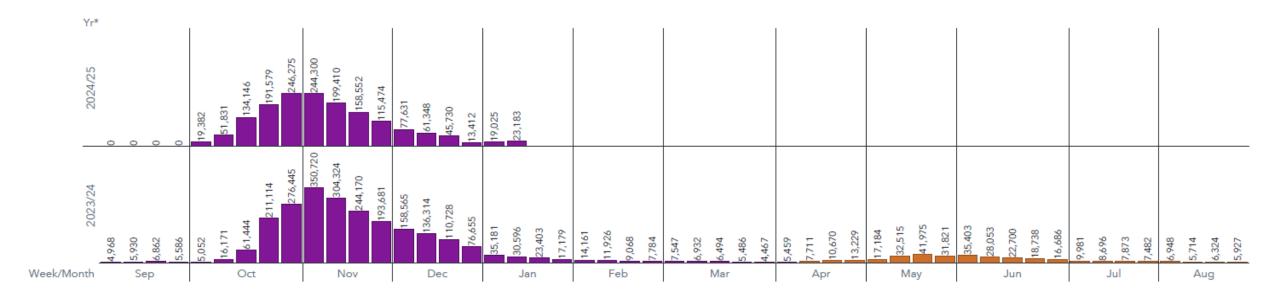


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1. Weekly Summary: Ontario's COVID-19 vaccination program



Trending: Doses administered week by week



Long-Term Care (LTC) Home Residents

Total LTC residents vaccinated with fall dose

35,846

% all LTC residents with fall dose

50.3%

Change from previous report:

592 residents

1.5% points

LTC residents who received a Fall 2023/24 dose (From September 12, 2023 to March 31, 2024): 45,290 (61.6%)

% LTC residents with a dose in the last 6 months

50.5%

Retirement Home (RH) Residents

Total RH residents vaccinated with fall dose

28,624

% all RH residents with fall dose

48.3%

Change from previous report:

363 residents

0.6% points

RH residents who received a Fall 2023/24 (From September 12, 2023 to March 31, 2024): 37,563 (63.4%)

% RH residents with a dose in the last 6 months

48.8%

Flu Distribution in LTCH/RH

As of January 13, 2025:

144,267 (+223 since January 6) doses of publicly funded flu vaccine have been distributed to **587** (no change since January 6) LTCHs via PHUs.

36,615 (+30 since January 6) doses of publicly funded flu vaccine have been distributed to **268** (no change since January 6) retirement homes via PHUs.

Please note:

 Many LTCHs and retirement homes are partnering with pharmacies – these homes and doses are not captured

Flu Administration in Pharmacy

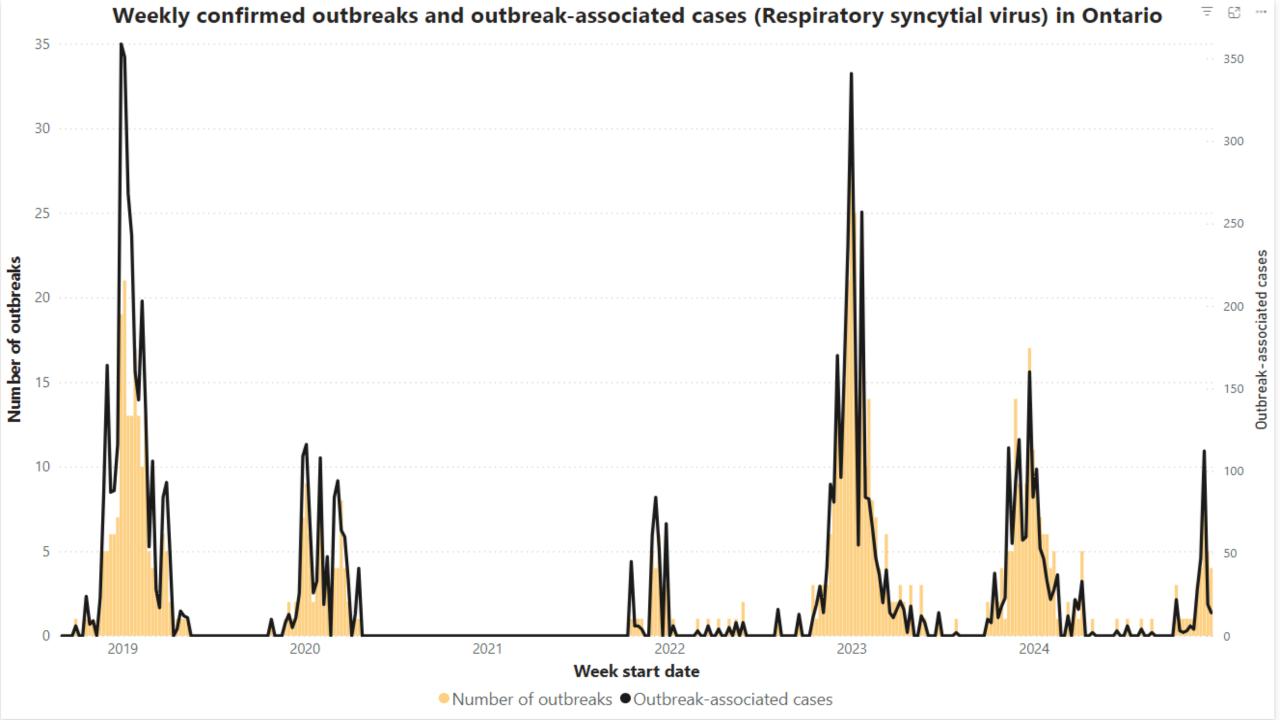
| | 2021/22 | 2022/23 | 2023/24 | 2024/25 |
|--|--|--|--|--|
| Week | Number of doses administered by week |
| Week 1 (Oct 3-9) | 13,798 | 3,171 | 25,584 | 191 |
| Week 2 (Oct 10-16) | 232,232 | 119,093 | 209,677 | 130,191 |
| Week 3 (Oct 17-23) | 198,502 | 238,289 | 258,305 | 236,677 |
| Week 4 (Oct 24-30) | 253,526 | 287,796 | 286,758 | 278,476 |
| Week 5 (Oct 31-Nov 6) | 235,578 | 357,353 | 297,034 | 299,635 |
| Week 6 (Nov 7-13) | 256,211 | 326,372 | 226,309 | 247,091 |
| Week 7 (Nov 14-20) | 161,638 | 261,190 | 167,801 | 189,944 |
| Week 8 (Nov 21-27) | 109,201 | 178,436 | 129,908 | 138,268 |
| Week 9 (Nov 28-Dec 4) | 76,491 | 133,519 | 107,329 | 98,707 |
| Week 10 (Dec 5-11) | 53,341 | 101,039 | 88,126 | 58,253 |
| Week 11 (Dec 12-18) | 34,391 | 73,304 | 62,559 | 52,682 |
| Week 12 (Dec 19-25) | 15,226 | 27,390 | 28,789 | 27,315 |
| Week 13 (Dec 26-Jan 1) | 14,413 | 26,201 | 25,334 | 25,368 |
| Week 14 (Jan 2-8) | 12,635 | 22,290 | 26,866 | 23,879 |
| Total | 1,667,183 | 2,155,443 | 1,940,379 | 1,806,677 |
| Total # of doses administered for season | 1,693,001 | 2,192,776 | 1,992,817 | |

| Age | 2 to 4 years old | 5 to 11 years old | 12 to 18 years old | 19 to 64 years old | 65 years and older | Total |
|----------------------------|------------------------|-------------------------|--------------------------|--------------------------|--------------------------|-----------|
| # of doses administered | 17,462 | 59,983 | 56,449 | 766,070 | 906,713 | 1,806,677 |
| Percentage | 0.97% | 3.32% | 3.12% | 42.40% | 50.19% | 100.00% |

Older Adult RSV Distribution

As of Jan. 10, 2025 (data pulled Jan. 13, 2025):

- 88,232 doses (∆ +682 doses) distributed to Public Health Units (PHU)
 - o 30,752 doses (△ o doses) of Arexvy
 - o 57,480 doses (△ +682 doses) of Abrysvo
- 73,012 doses (Δ +1,067 doses) distributed to Health Care Providers (HCP)
 - \circ 19,445 doses (\triangle +324 doses) distributed to 507 (\triangle +5) Long-Term Care Homes (LTCH) via PHUs
 - \circ 12,800 doses (\triangle +103 doses) distributed to 219 (\triangle +1) Retirement Homes (RH) via PHUs



Infant RSV Distribution

As of Jan. 10, 2025 (data pulled Jan. 13, 2025):

- 121,832 doses (Δ +1,371 doses) distributed to Public Health Units (PHU)
 - 50mg: 53,780 doses (Δ -46 doses)
 - 100mg: 68,052 doses (∆ +1,417 doses)
- 106,558 doses (△ +4,120 doses) distributed to Health Care Providers (HCP)
 - Hospitals: 38,798 doses
 - o 50mg: 28,907 doses
 - o 100mg: 9,891 doses
 - Primary care providers: 67,760 doses
 - o 50mg: 16,712 doses
 - o 100mg: 51,048 doses

Early indications of RSV uptake before discharge



Nov 1 – Dec 31, 2024
BORN captured RSV prevention information on 11,438 newborns
(~47% of expected birth volumes)



70% received Beyfortus (nirsevimab)
Increase from 67% in November



If not given, then why?

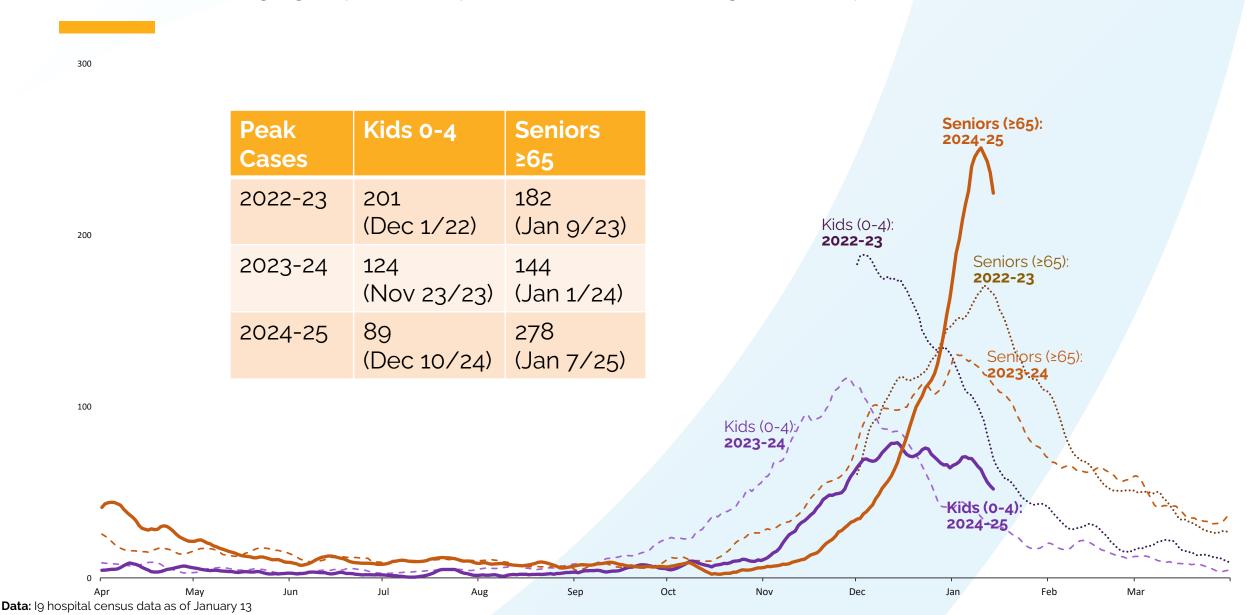
- 1) 19% parent declining
- 2) 4.2% prenatal RSV protection via maternal vaccine
- 3) <3% other reasons (including preference to delay, baby unwell, etc)
- 4) <2% discharge prior to provision (including to NICU),
- 5) <2% in the early stages of the roll-out no supply available or program not yet initiated at

Anecdotally, clinicians share that pre-admission awareness of the program reduces hesitancy.



Comparing RSV hospitalizations by age: Kids (0-4) and seniors (≥65)

The 0-4 and ≥65 age groups make up 82% of historical average RSV hospital census.



23

Mycoplasma pneumoniae

Diagnosis and treatment for paediatric care providers





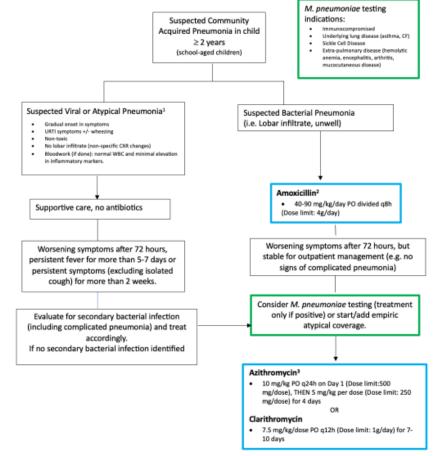






Mycoplasma pneumoniae

Community Acquired Pneumonia – Outpatient Management Algorithm



Key Points:

- Most patients with a mild M. pneumoniae infection will recover without antibiotics.
- Either low dose amoxicillin (40-50 mg/kg/day) or high dose amoxicillin (80-90 mg/kg/day) <u>divided 3 times a day</u> can be used in most jurisdictions as Streptococcus pneumoniae susceptibility to penicillin remains high. Alternatively, if twice a day dosing is prescribed, higher dose amoxicillin should be used (45 mg/kg/dose 90-q 12 k, maximum 4 g/day)
- 3. Macrolides alone should not be used alone if Streptococcus pneumoniae is suspected due to high resistance rates (up to 50%)

Additional Treatment Considerations

Levofloxacin and Doxycycline should be reserved for patients with confirmed M. pneumoniae infections where IV therapy is needed (levofloxacin), there is severe disease, or lack of response after 48-72 hours of macrolide therapy (levofloxacin or doxycycline)

Amoxicillin/clavulanate would add coverage for infections with beta-lactamase producing organisms (e.g. Haemophilus influenzae, Moraxella catarrhalis) as well as methicillin susceptible Staphylococcus aureus, but these are less common causes of pneumonia in children.

Human Metapneumovirus (HMPV)

- HMPV is common, on average 10% of seasonal respiratory illness in children is caused by HMPV — most people get it before they turn 5
- Causes predominantly upper respiratory symptoms
- High risk groups:
 - Are younger than 5 (especially premature infants) or older than 65.
 - Immunocompromised.
 - Have asthma or COPD.
- Recent surge in cases of human metapneumovirus in northern China
 - Within the expected range for the winter season with no unusual outbreak patterns reported

COVID-19

Percent positivity

in the most recent week

13.5%

Outbreaks

reported in the most recent week

78

Hospital bed occupancy

reported in the most recent week

865

Deaths

reported in the most recent week

8

Episodes

reported in the most recent week

1,655

Influenza (all types)

Percent positivity

in the most recent week

11.8%

Outbreaks

reported in the most recent week

11

Hospital bed occupancy

reported in the most recent week

225

Deaths

reported in the most recent week

Not available

Cases

reported in the most recent week

1,181

RSV

Percent positivity

in the most recent week

10.8%

Outbreaks

reported in the most recent week

4

Hospital bed occupancy

reported in the most recent week

239

Deaths

reported in the most recent week

Not available

Cases

reported in the most recent week

Not available

Other respiratory virus activity in the most recent week

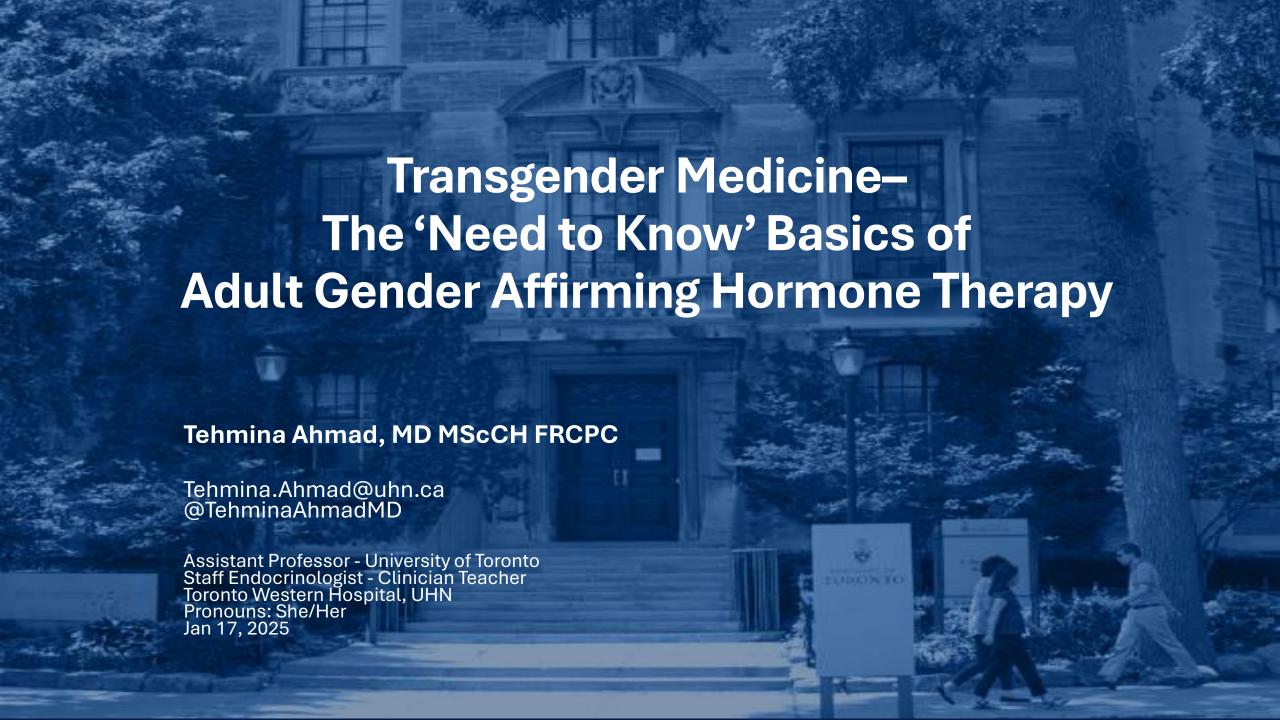
| Virus | Percent positivity (%) |
|----------------------------|------------------------|
| Adenovirus | 1.0 |
| Entero/Rhinovirus | 5.9 |
| Human metapneumovirus | 0.7 |
| Parainfluenza (all types) | 1.8 |
| Seasonal human coronavirus | 1.8 |

Download data for the previous and current surveillance period

Cases Outcomes

Lab testing / percent positivity Outbreaks

Historical activity assessment PHU influenza activity/positivity



Learning Objectives

1. Understand terminology related to sex, gender identity, gender expression, and sexual orientation.

- 2. Build an approach to the diagnosis of gender dysphoria/gender incongruence, gender health history, and informed consent.
- 3. Review common adult gender affirming hormone therapies (GAHT), dosing, monitoring, and side effects.

Outline

1. Conceptualizing gender identity, gender expression, and sex

- 2. Understanding 'transitioning'
- 3. The 'What, Who, How, and Why of gender affirming care
- 4. How to provide (adult) gender affirming hormone therapy
- 5. The one-liner on gender affirming surgeries & regret



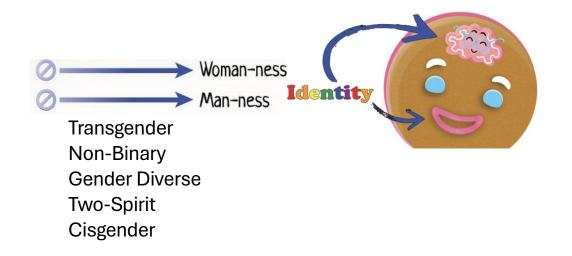
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- Transgender: A person whose gender identity differs from the sex that was assigned at birth
- Sex (AFAB/AMAB): The sex assigned at birth based on genotype or phenotypic external genitalia

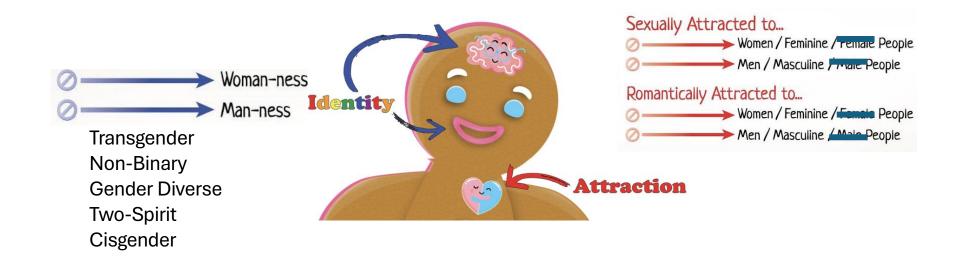
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- **Gender identity**: A person's deeply felt, internal, intrinsic sense of their own gender

- Transgender: A person whose gender identity differs from the sex that was assigned at birth
- Sex (AFAB/AMAB): The sex assigned at birth based on genotype or phenotypic external genitalia
- **Gender identity**: A person's deeply felt, internal, intrinsic sense of their own gender
- Transsexual: A clinical term which had historically been used to describe those transgender people who sought medical intervention (hormones, surgery) for gender affirmation (term is no longer used!)

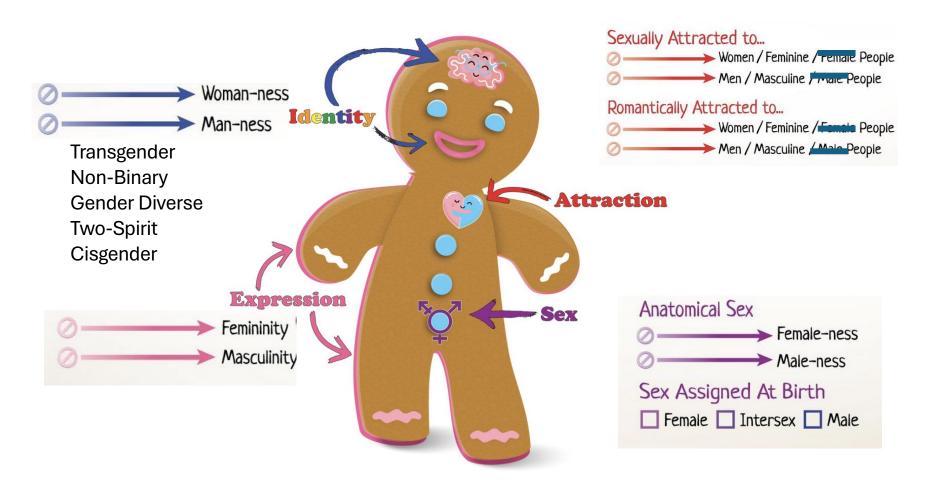
Gender Identity ≠ Sexual Orientation ≠ Sex ≠ Gender Expression



Gender Identity ≠ Sexual Orientation ≠ Sex ≠ Gender Expression



Gender Identity ≠ Sexual Orientation ≠ Sex ≠ Gender Expression



Genderbread Person

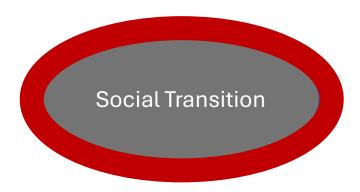
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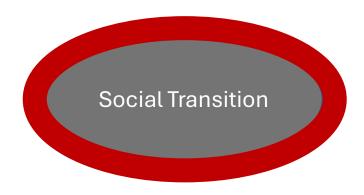


What Does it Mean to 'Transition'?



- Adopting new name and pronouns
- Altering physical appearance to align with gender identity
 - Hairstyle
 - Dress + Cosmetics
 - Body Hair Changes
- Adopting mannerisms and behaviors aligned with preferred gender role

What Does it Mean to 'Transition'?

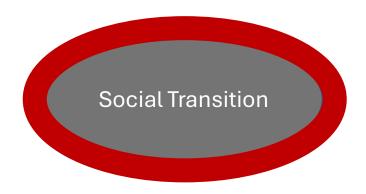


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- Puberty blockers to prevent or stall secondary sex changes
- Gender affirming hormones to induce changes associated with gender identity

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- Puberty blockers to prevent or stall secondary sex changes
- Gender affirming hormones to induce changes associated with gender identity



- Surgical removal of morphologic indicators of birth sex
- Surgical creation of morphologic features of sex associated with gender identity

Outline

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Research Letter | Surgery

June 27, 2024

Prevalence of Gender-Affirming Surgical Procedures Among Minors and Adults in the US

Dannie Dai, BS¹; Brittany M. Charlton, ScD²; Elizabeth R. Boskey, PhD³; et al

Author Affiliations | Article Information

JAMA Netw Open. 2024;7(6):e2418814. doi:10.1001/jamanetworkopen.2024.18814

Cross-sectional study

Calculated the rate of people who received GAS with a trans-related dx per 100,000 people categorized as:

- adults (18yo or older)
- minors (15-17yo, 13-14yo, <12yo)

And compared this to the proportion of gender affirming breast reduction surgeries in **cis-males** for gynecomastia

Recall, if you were born male, and identify as a man; your breast reduction was done to affirm your male gender identity

Research Letter | Surgery

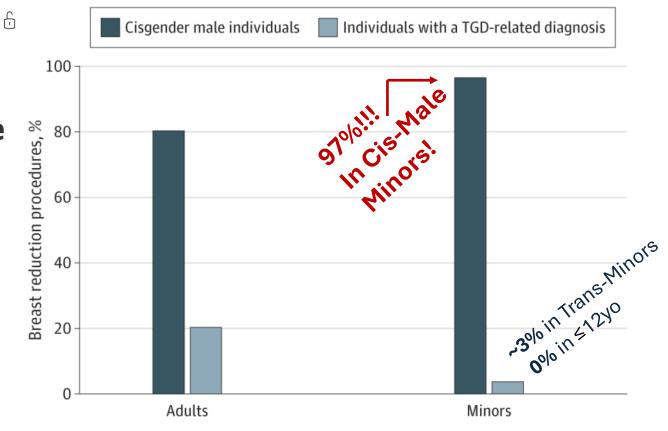
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Research Letter

January 6, 2025

Gender-Affirming Medications Among Transgender Adolescents in the US, 2018-2022

Landon D. Hughes, PhD¹; Brittany M. Charlton, ScD²; Isa Berzansky, MSc²; et al

Author Affiliations

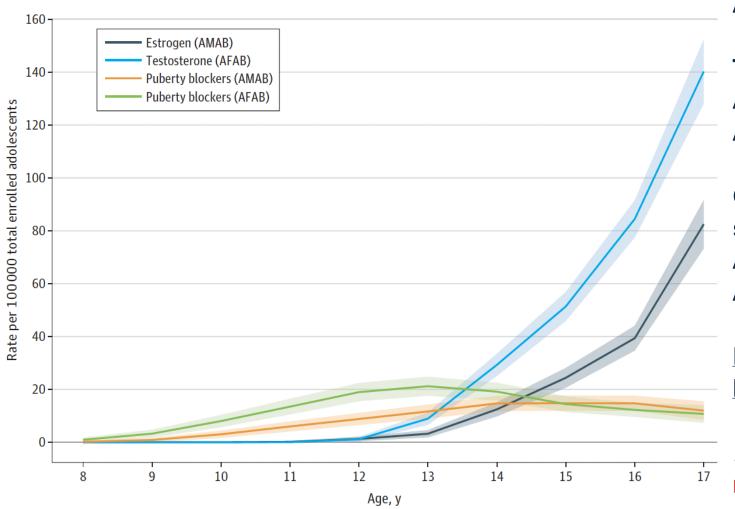
JAMA Pediatr. Published online January 6, 2025. doi:10.1001/jamapediatrics.2024.6081

Cross-sectional study

Calculated the rate of adolescents with a TGD-related diagnosis who received hormonal care per 100 000 total adolescents enrolled in insurance by age (8-17yo) and sex assigned at birth.

The sample included yielded: *11 879 766* person-years of data

Figure. Gender-Affirming Medication Receipt per 100 000 Enrolled Adolescents by Age and Sex Assigned at Birth, 2018-2022



The rate of receiving *Puberty Blockers*:

AFAB – 20.81 per 100 000 adolescents AMAB – 15.22 per 100 000 adolescents

The rate of receiving *Hormone Therapy*:

AFAB – 49.9 per 100 000 adolescents AMAB – 25.34 per 100 000 adolescents

GAHT rates peak at age 17yo, **but** still remain low:

AFAB – 140.16 per 100 000 adolescents AMAB – 82.42 per 100 00 adolescents

NO adolescents received a hormone prescription when ≤12yo

Fewer than 1 in 1,000 adolescents receive hormones or puberty blockers

Is this Care *Really* Supported?

American Academy of Child and Adolescent Psychiatry

American Academy of Dermatology

American Academy of Family Physicians

American Academy of Nursing

American Academy of Pediatrics

American Academy of Physician Assistants

American College Health Association

American College of Nurse-Midwives

American College of Obstetricians and Gynecologists

American College of Physicians

American Counseling Association

American Heart Association

American Medical Association

American Medical Student Association

American Nurses Association

American Osteopathic Association

American Psychiatric Association

American Psychological Association

American Public Health Association

American Society of Plastic Surgeons

Endocrine Society

Federation of Pediatric Organizations

GLMA: Health Professionals Advancing LGBTQ Equality

National Association of Nurse Practitioners in Women's Health

National Association of Social Workers

National Commission on Correctional Health Care

Ontario Medical Association

Pediatric Endocrine Society

Society for Adolescent Health and Medicine

World Medical Association

World Professional Association for Transgender Health

Is this Care *Really* Supported?



"We **stand firm in our support** of gender-affirming care. Transgender and gender-diverse people deserve access to needed and often life-saving medical care.

NHS England's recent report, the Cass Review, **does not contain any new research** that would contradict the recommendations made in our <u>Clinical Practice Guideline</u> on genderaffirming care.

The [Clinical Practice] guideline, which cites more than 260 research studies, recommends a very conservative approach to care, with no medical intervention prior to puberty.

...Banning evidence-based medical care based on misinformation takes away the ability of parents and patients to make informed decisions.

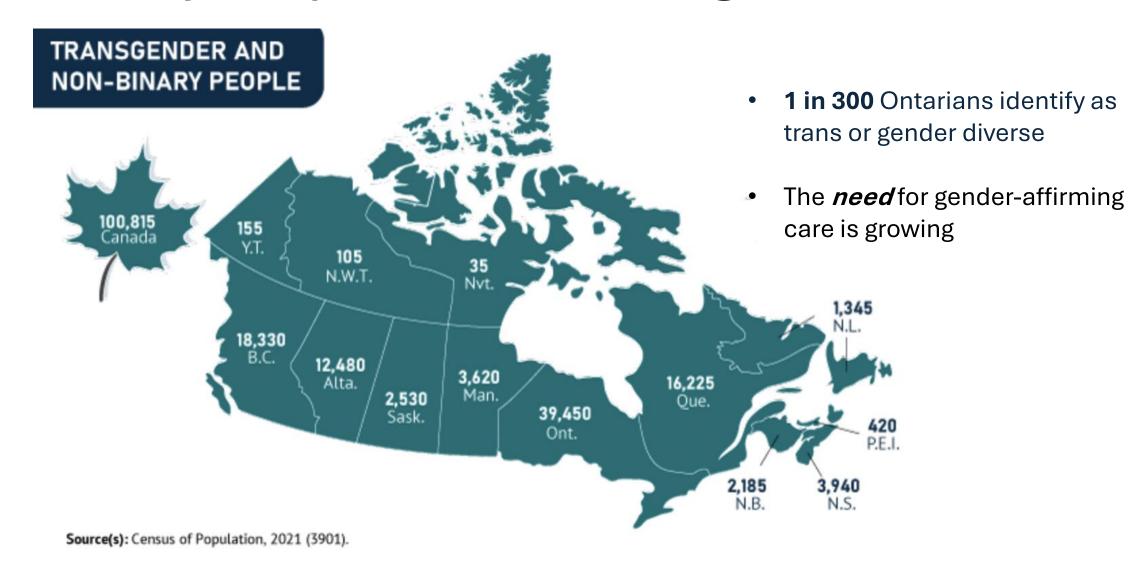
Medical evidence, not politics, should inform treatment decisions."

Outline

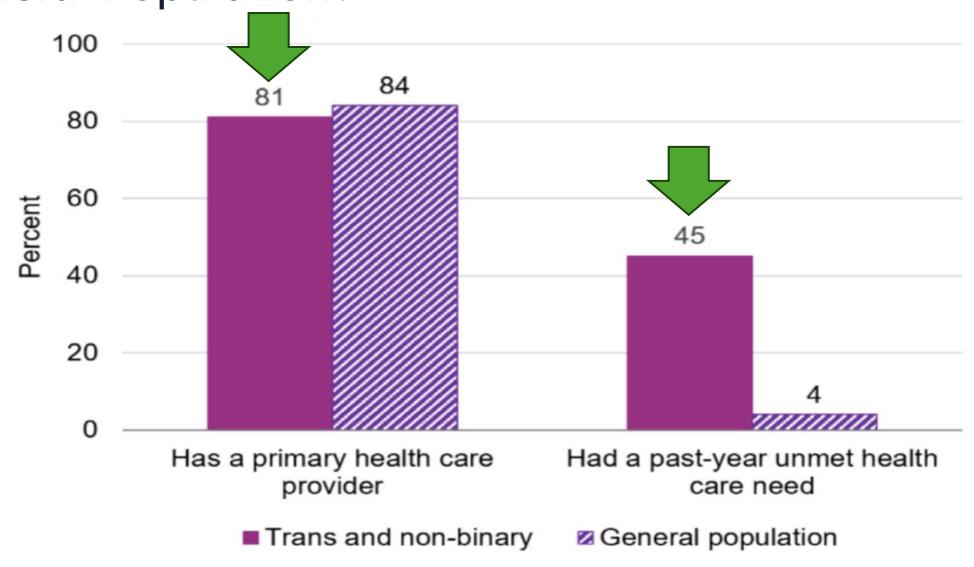
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How Many People Are We Talking About?



How Does Trans Healthcare Access Compare to the General Population?



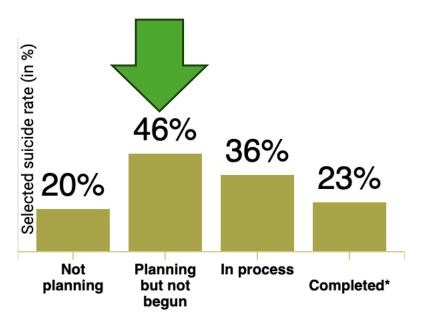
Factors Impacting Discomfort with Family Doc

| Has a family doctor ever | Has a family doctor ever Transmasculine spectrum ^a N = 184 | | Transfeminine spectrum ^b n = 172 | |
|--|--|---------------------|---|---------------------|
| | | | | |
| | % | 95% CI [†] | % | 95% CI [†] |
| Refused to see you or ended care because you are trans | 7.2 | (1.6, 12.9) | 5.0 | (0.0, 10.3) |
| Used hurtful or insulting language about trans identity or experience | 9.0 | (3.5, 14.4) | 12.1 | (4.1, 20.0) |
| Refused to discuss or address trans-related health concerns | 10.6 | (5.3, 15.8) | 13.5 | (5.5, 21.6) |
| Told you that you are not really trans | 8.2 | (2.7, 13.6) | 10.9 | (2.8, 19.0) |
| Discouraged you from exploring gender | 9.1 | (3.4, 14.8) | 6.6 | (0.7, 12.4) |
| Told you they don't know enough about trans-related care to provide it | 24.5 | (15.8, 33.1) | 29.1 | (18.5, 39.7) |
| Belittled or ridiculed you for being trans | 6.8 | (1.2, 12.4) | 8.1 | (1.9, 14.3) |
| Thought the gender listed on your ID or forms was a mistake | 6.2 | (2.3, 10.1) | 3.8 | (0.0, 9.0) |
| Refused to examine parts of your body because you are trans | 4.5 | (1.4, 7.7) | 6.2 | (0.2, 12.2) |
| At least one of the above | 37.2 | (27.2, 47.1) | 38.1 | (26.5, 49.6) |

Why Should We Care?

Suicidal ideation and attempt rates of **trans** people at different stages of medical transition:

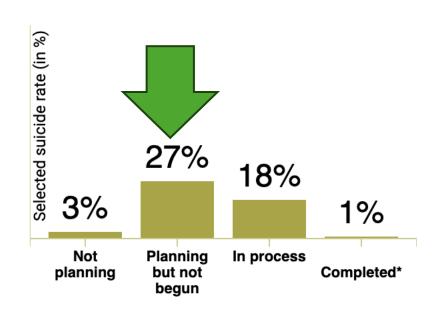




Medical Transition Status

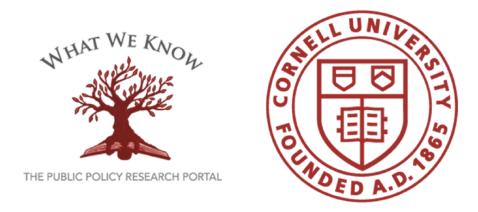
Suicidal ideation and attempt rates of **trans** people at different stages of medical transition:





Medical Transition Status

GAHT Improves Well-Being



www.whatweknow.info

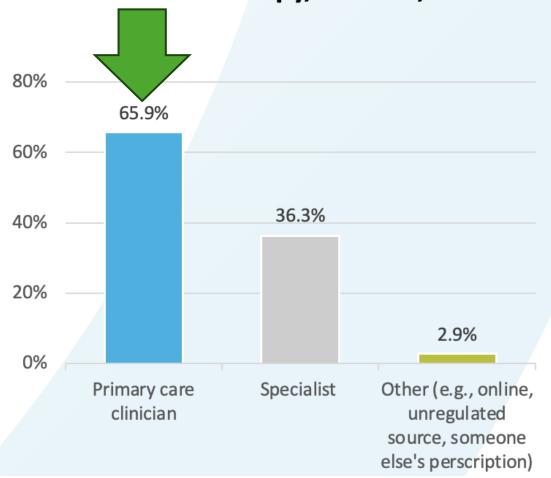
What does the scholarly research say about the effect of gender transition on transgender well-being?

Overview: We conducted a systematic literature review of all peer-reviewed articles published in English between 1991 and June 2017 that assess the effect of gender transition on transgender well-being. We identified 56 studies that consist of primary research on this topic, of which 52 (93%) found that gender transition improves the overall well-being of transgender people, while 4 (7%) report mixed or null findings. We found no studies concluding that gender transition causes overall harm. As an added resource, we separately include 17 additional studies that consist of literature reviews and practitioner guidelines.

Bottom Line: This search found a robust international consensus in the peer-reviewed literature that gender transition, including medical treatments such as hormone therapy and surgeries, improves the overall well-being of transgender individuals. The literature also indicates that greater availability of medical and social support for gender transition contributes to better quality of life for those who identify as transgender.

You Are Already at the Forefront of Care!

Self-reported source of gender-affirming hormone therapy, Ontario, 2019



- 2019 Trans PULSE Canada survey found that
 65.9% of respondents on hormone therapy in
 Ontario were prescribed by *primary care*
- There are 560 family medicine residency positions in Ontario
- There are only 11 endocrinology positions each year in Ontario

Outline

- 1. Conceptualizing gender identity, gender expression, and sex
- 2. Understanding 'transitioning'
- 3. The 'What, Who, How, and Why of gender affirming care
- 4. How to provide (adult) gender affirming hormone therapy
- 5. The one-liner on gender affirming surgeries & regret

Diagnostic Terminology

Gender Identity Disorder (DSM4 -1994)

Transgender identity viewed as mental health disorder

Mental Health Provided Gate keeping

No trans input into care framework/pathways

Mental Health involved Consent Gender Dysphoria (DSM5 -2013)

Transgender identity not considered a disorder; distress arising from diverse gender identity is the disorder

Variable practice in mental health provider required input

No trans input into care framework/pathways

Written Consent

Gender Incongruence (ICD 2019)

Greater recognition and acceptance for gender diversity

Mental health provider not needed to provide permission

Trans input into care framework

Verbal Consent

WPATH SOC 7 (2011)



WPATH SOC 8 (2022)

How to Diagnose Gender Dysphoria?

A. Marked incongruence between one's experienced/expressed gender and assigned gender, of at *least 6 months* duration, as manifested by at least TWO of the following:

A marked incongruence between one's experienced/expressed gender and primary and/or secondary sex characteristics (or in young adolescents, the anticipated secondary sex characteristics)

A strong desire to be rid of one's primary and/or secondary sex characteristics because of a marked incongruence with one's experienced/expressed gender

A strong desire for the primary or secondary sex characteristics of the other gender

B. The condition is associated with clinically significant *distress* or *impairment* in social, occupational, or other important areas of functioning

NB: There is added distress from simply living in a transphobic society.

Gender Identity

- In your own words, how would you describe your gender identity?
- When did you first recognize your assigned gender did not match your gender identity?
- What was puberty like? Are there parts of your body associated with gender you like/don't like?

Understanding relevant information about the treatment

Appreciating treatment information in the context of your own life Decision-Making Capacity

Reasoning about available treatment options

Gender Identity

- In your own words, how would you describe your gender identity?
- When did you first recognize your assigned gender did not match your gender identity?
- What was puberty like? Are there parts of your body associated with gender you like/don't like?

Understanding relevant information about the treatment

Gender Expression

 Have you started making changes to your outward appearance to more closely match who you are?

Appreciating treatment information in the context of your own life Decision-Making Capacity

Reasoning about available treatment options

Gender Identity

- In your own words, how would you describe your gender identity?
- When did you first recognize your assigned gender did not match your gender identity?
- What was puberty like? Are there parts of your body associated with gender you like/don't like?

Understanding relevant information about the treatment

Gender Expression

 Have you started making changes to your outward appearance to more closely match who you are?

Appreciating treatment information in the context of your own life Decision-Making Capacity

Reasoning about available treatment options

Perceptions of Others/Support

- How do you wish others would see you?
- How does your gender identity impact your life at work?
 Your relationships? Your family?
- Who are your supports?

Understanding Expectations

- Can you tell me about how hormone therapy will work for you?
- What changes you are looking forward to?
- Are there any changes that you aren't sure about?
- Have you considered any alternatives to hormone therapy?

Understanding relevant information about the treatment

Appreciating treatment information in the context of your own life Decision-Making Capacity

Reasoning about available treatment options

Understanding Expectations

- Can you tell me about how hormone therapy will work for you?
- What changes you are looking forward to?
- Are there any changes that you aren't sure about?
- Have you considered any alternatives to hormone therapy?

Best Practices

- Fertility preservation referral?
- Ask about partners/protection/contraception?
- Risk mitigation: Smoking cessation?

Understanding relevant information about the treatment

Appreciating treatment information in the context of your own life Decision-Making Capacity

Reasoning about available treatment options

Contraindications to Gender Affirming Hormone Therapy (GAHT)

Absolute Contraindications to Estrogen

- unstable ischemic cardiovascular disease
- active, known sex hormone-sensitive cancer (e.g. breast, prostate)
- end-stage chronic liver disease
- psychiatric conditions that limit the ability to provide informed consent
- hypersensitivity to one of the components of the formulation

Absolute Contraindications to Testosterone

- pregnancy or breastfeeding
- active, known sex hormone-sensitive cancer (e.g. breast, endometrial)
- unstable ischemic cardiovascular disease
- poorly controlled psychosis or acute homicidally
- psychiatric conditions that limit the ability to provide informed consent
- hypersensitivity to one of the components of the formulation

Feminizing Hormone Therapy: Estrogen +/- Anti-Androgen

Estrogen often serves as the foundation of feminizing hormone therapy (17 β -estradiol)

Anti-Androgens may be used in conjunction with estrogen to reduce effects of testosterone *Spironolactone*

- Peripheral Androgen receptor antagonist
- Mineralocorticoid antagonist: induces gynecomastia (intended effect)

Cyproterone Acetate

- Peripheral Androgen receptor antagonist + Central Androgen antagonist
- Progesterone activity: inhibits LH and therefore testosterone production

*Leuprolide

 GnRH agonist, reduces GnRH pulsatility causing reversible central down-regulation of LH and FSH signalling

Estrogen +/- Anti-Androgen Formulations & Dosing

| Gender Affirming Therapy | Starting dose | Usual Dose | Maximum dose |
|---------------------------------|---------------------|---------------------------|---------------------------|
| Estradiol (oral) | 0.5mg to 2mg | 4mg daily (or div BID) | 6mg daily (or div BID) |
| Estradiol | 50mcg patch, | 100mcg to 200mcg | 200 to 300mcg |
| (transdermal patch) | change 2x per week | change 2x per week | change 2x per week |
| Estradiol | 2.5g daily of | 3.75g to 5g daily of | 6.75g daily of |
| (transdermal gel) | 0.06% estradiol gel | 0.06% estradiol gel | 0.06% estradiol gel |
| Estradiol valerate (injectable) | 3mg once weekly | 3 to 6mg weekly | 10mg weekly |
| Spironolactone (oral) | 50 to 100mg daily | 200mg daily | 300mg daily |
| | | (or divided BID) | (or divided BID) |
| Cyproterone (oral) | 12.5mg q 2 days | 12.5mg daily | 25mg daily* |

Note: Generic Oral Estradiol is covered by provincial drug coverage plans (Trillium, ODSP, OW, OHIP+)

EAP for 'transdermal' estradiol only applicable if proven that patient cannot swallow pills

Estrogen +/- Anti-Androgen Formulations & Dosing

| Gender Affirming Therapy | Starting dose | Usual Dose | Maximum dose |
|---|--|--|-------------------------------------|
| Estradiol (oral) Do labs 4-6hrs post E2 | 0.5mg to 2mg 2 po or SL | 4mg daily (or div BID) | 6mg daily (or div BID) |
| Estradiol Do labs btwn patch ch (transdermal patch) | 50mcg patch, nange change 2x per week | 100mcg to 200mcg change 2x per week | 200 to 300mcg change 2x per week |
| Estradiol Do labs 4-6hrs post Ez (transdermal gel) | 2.5g daily of 0.06% estradiol gel | 3.75g to 5g daily of 0.06% estradiol gel | 6.75g daily of 0.06% estradiol gel |
| Estradiol valerate Do labs on post inject (injectable) Do labs on post inject | 3mg once weekly ion day 3 or day 4 if q weekly ion day 7 or day 8 if g2 weekly | 3 to 6mg weekly | 10mg weekly |
| Spironolactone (oral) | 50 to 100mg daily | 200mg daily (or divided BID) | 300mg daily (or divided BID) |
| Cyproterone (oral) | 12.5mg q 2 days | 12.5mg daily | 25mg daily* |

*Target estradiol

*Target testosterone

~200-735pmol/L

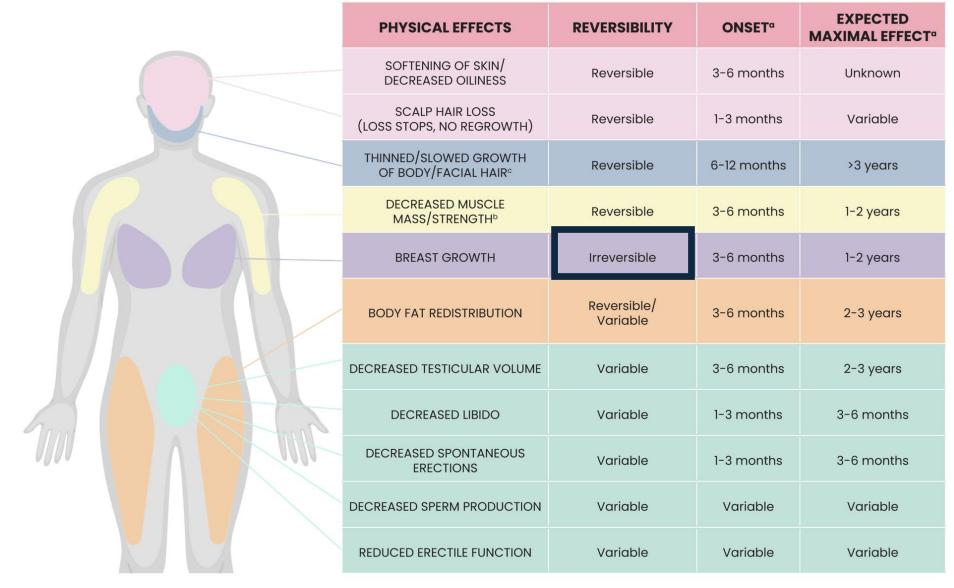
~<2nmol/L

**<735pmol/L

Note: Generic Oral Estradiol is covered by provincial drug coverage plans (Trillium, ODSP, OW, OHIP+)

EAP for 'transdermal' estradiol only applicable if proven that patient cannot swallow pills

Timeline of Expected Effects – Feminizing GAHT



Estrogen & Anti-Androgen Side Effects

| | Estrogen | Spironolactone | Cyproterone |
|--------------------------------|--|---|--|
| Possible side effects | VTE Cholelithiasis Hyper Tg Wt Gain Incr risk of Br Ca | High K, Cr (caution with ACEi/ARB) Polyuria/Dehydration | Liver toxicity, Depression/mood impacts, VTE Black box warning on meningioma risk with >10mg/d doses |
| Effects on testosterone levels | Decrease in testosterone levels | Mild/moderate decrease in testosterone levels | Effective testosterone suppression |
| Strength/potency | Moderate/strong anti-androgen | Weaker anti-androgen Peripheral androgen blocker | Stronger anti-androgen Peripheral and central androgen blocker |
| Effects on HDL cholesterol | Variable impact on HDL | HDL increases | HDL decreases |
| Effects on prolactin | May increase prolactin levels | Little to no increase in prolactin levels | Significant increase in prolactin levels is common |

Lab Monitoring for Feminizing GAHT

| | BASELINE | MONTH 3 | MONTH 6 | MONTH 12° | YEARLY | ACCORDING TO GUIDELINES FOR CIS PATIENTS, OR PROVIDER DISCRETION |
|-----------------------------|--|----------------|--|------------------|---------------|---|
| EXAM/ INVESTIGATIONS | Focused Physical Exam. Include: height, weight, BP, +/- breast inspection/ measurement(s)* | BP, we meas | ight, +/- breast inspe surement(s) at 12 mo | ection/ nths* | transfeminine | entive care checklist for patients and accompanying ons in the full Guidelines. |
| BLOODWORK | | | | | | |
| СВСа | \checkmark | √ | √ | \checkmark | ✓ | |
| ALT ^b | \checkmark | \checkmark | | \checkmark | √ | \checkmark |
| CREATININE/ LYTESC | \checkmark | \checkmark | \checkmark | ✓ | ✓ | |
| HbA1c OR FASTING GLUCOSE | \checkmark | | | \checkmark | | \checkmark |
| LIPID PROFILE | \checkmark | | | \checkmark | | \checkmark |
| TOTAL TESTOSTERONE | \checkmark | \checkmark | \checkmark | \checkmark | \checkmark | |
| ESTRADIOL | \checkmark | \checkmark | \checkmark | \checkmark | \checkmark | |
| PROLACTIN | \checkmark | | | √ | √ | ✓ |

Masculinizing Hormone Therapy: Testosterone

Testosterone serves as the foundation of masculinizing hormone therapy

Testosterone enanthate (IM/SC)

- 'double concentrated' 200mg/mL (less volume required for same dose)
- formulated in sesame seed oil

Testosterone cypionate (IM/SC)

- -'single concentrated' 100mg/mL
- -formulated in cotton seed oil

Testosterone 1% gel (daily topical application)

Can apply for exceptional access program (EAP) for funding through provincial drug coverage plans (Trillium, ODSP, OW, OHIP+)

Testosterone Formulations & Dosing

| Gender Affirming Therapy | Starting dose | Usual Dose | Maximum dose |
|---------------------------------------|------------------------|------------------------|--------------|
| Testosterone enanthate (injectable) | 20mg to 40mg weekly | 40mg to 80mg weekly | 100mg weekly |
| or | | | |
| Testosterone cypionate (injectable) | | | |
| Testosterone gel 1% (transdermal gel) | 2.5g daily | 5g to 10g daily | 10g daily |

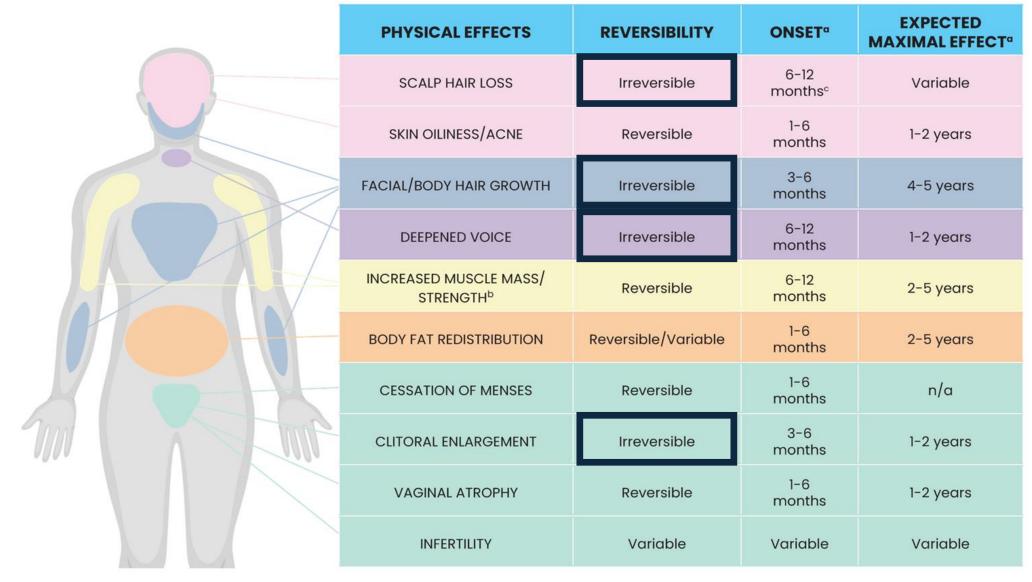
Testosterone Formulations & Dosing

| Gender Affirming Therapy | Starting dose | Usual Dose | Maximum dose | |
|---------------------------------------|--|------------------------|------------------|--|
| Testosterone enanthate (injectable) | 20mg to 40mg weekly | 40mg to 80mg weekly | 100mg weekly | |
| or | Do labs on post inje | ection day 3 or day | y 4 if q weekly | |
| Testosterone cypionate (injectable) | Do labs on post inje | ection day 7 or day | y 8 if q2 weekly | |
| Testosterone gel 1% (transdermal gel) | 2.5g daily | 5g to 10g daily | 10g daily | |
| | Do labs ~ 4-6 hours post gel application | | | |

^{*}Target testosterone

~8.5-30nmol/L

Timeline of Expected Effects – Masculinizing GAHT



Testosterone Side Effects

| | Testosterone |
|-----------------------------|--|
| Possible side effects | Polycythemia (Hb/Hct) Acne Androgenic alopecia (hereditary) Hypertension Sleep apnea Weight gain |
| Effects on estradiol levels | Decrease in estradiol levels *re-assess T-dosing if estradiol is elevated* |
| Strength/potency | Strong anti-estrogen |
| Effects on HDL cholesterol | HDL decreases; LDL increases |
| Effects on prolactin | No impact on prolactin levels |

Lab Monitoring for Masculinizing GAHT

| | BASELINE | MONTH 3 | MONTH 6 | MONTH 12 ^{b,c} | YEARLY | ACCORDING TO GUIDELINES FOR CIS PATIENTS, OR PROVIDER DISCRETION |
|-----------------------------|---|--------------|--------------|-------------------------|---|---|
| EXAM/ INVESTIGATIONS | Focused Physical Exam with PAP if indicated. Include: height, weight, BP. | BP, weight | | | See Preventive Care Checklist for Transmasculine Patients and accompanying explanations in the Guidelines for Gender-Affirming Primary Care with Trans and Non-Binary Patients. | |
| BLOODWORK | | | | | | |
| СВС | \checkmark | \checkmark | \checkmark | \checkmark | \checkmark | |
| ALT | \checkmark | √° | | | \checkmark | |
| HbA1c OR FASTING GLUCOSE | \checkmark | √° | | | \checkmark | |
| LIPID PROFILE | \checkmark | √° | | | \checkmark | |
| TOTAL TESTOSTERONE | \checkmark | \checkmark | \checkmark | \checkmark | \checkmark | |
| LH° | | | | | | |

Key Summary: Endocrine Society 2017 Guidelines

- 3.1. We recommend that clinicians <u>confirm the diagnostic criteria of GD/gender incongruence</u> and the criteria for the endocrine phase of gender transition before beginning treatment. $(1 | \oplus \oplus \oplus \bigcirc)$
- 3.3. We suggest that clinicians <u>measure hormone levels during treatment</u> to ensure that endogenous sex steroids are suppressed and administered sex steroids are maintained in the normal physiologic range for the affirmed gender. (2 $|\oplus \oplus \bigcirc \bigcirc$)
- 3.4. We suggest that endocrinologists provide education to transgender individuals undergoing treatment about the <u>onset and time course</u> of physical changes induced by sex hormone treatment. (2 $|\oplus \bigcirc\bigcirc\bigcirc$)
- 4.1. We suggest <u>regular clinical evaluation</u> for physical changes and potential adverse changes in response to sex steroid hormones and laboratory monitoring of sex steroid hormone levels <u>every 3 months during the first</u> <u>year of hormone therapy</u> for transgender males and females and then once or twice yearly. (2 $|\oplus \oplus \bigcirc$)

Outline

- 1. Conceptualizing gender identity, gender expression, and sex
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- 5. The one-liner on gender affirming surgeries & regret



What Gender Affirming Surgeries are Available?

OHIP-Funded Procedures



Mastectomy*

Clitoroplasty

Augmentation

Scrotoplasty

Mammoplasty**

Metoidioplasty

Phalloplasty

Labiaplasty

Erectile and/or testicular implant

Salpingo-

coordanar impi

oophrectomy

Vaginectomy

Hysterectomy

Orchidectomy

Vaginoplasty

- * Mastectomy includes removal of breast tissue and reconstruction, which involves removal of excess skin, reduction and proper positioning of nipple and areola, and minimization of chest wall scars.
- ** Augmentation mammoplasty is insured when no breast enlargement has occurred, following 12 continuous months of hormone therapy (unless hormones are not appropriate for the person).

Upper Surgery (Specifically chest)

Needs 1 supporting assessment, that has to be from a **qualified** Physician or Nurse Practitioner

Lower Surgery (Genital Surgery)

Needs 2 supporting assessments:

 1 assessment from a qualified Physician or Nurse Practitioner

AND

 A 2nd assessment from a qualified Physician, Nurse Practitioner, Psychologist or Registered Social Worker with a Masters Degree

What Gender Affirming Surgeries are Available?

| Surgery | 1st Letter (MD/NP) | 2nd Letter (NP/ MD/ RN/MSW/ Psychologist) | Hormones | Medical and Mental Health Coniditons | Gender Role Experience |
|--------------------------------------|-----------------------|--|---|--------------------------------------|--|
| Mastectomy | X | | Not a pre-requisite | "controlled" | |
| Augmentation Mammoplasty | X | | 12 continuous months with no breast development | "controlled" | |
| Gonad: Hysterotomy or Orchiectomy | х | X | 12 continuous months | "well controlled" | |
| Vaginoplasty | х | X | 12 continuous months | "well controlled" | 12 continuous months of living in a gender role congruent with gender identity |
| Phalloplasty/ Medtoidioplasty | X | X | 12 continuous months | "well controlled" | 12 continuous months of living in a gender role congruent with gender identity |

What About Regret?

RECONSTRUCTIVE: ORIGINAL ARTICLE

Regret after Gender-affirmation Surgery: A Systematic Review and Meta-analysis of Prevalence

Bustos, Valeria P. MD*; Bustos, Samyd S. MD†; Mascaro, Andres MD‡; Del Corral, Gabriel MD, FACS§; Forte, Antonio J. MD, PhD, MS¶; Ciudad, Pedro MD, PhD¹; Kim, Esther A. MD**; Langstein, Howard N. MD††; Manrique, Oscar J. MD, FACS††

Author Information ⊗

Plastic and Reconstructive Surgery - Global Open 9(3):p e3477, March 2021. | DOI: 10.1097/GOX.000000000003477 €

Systematic review and meta-analysis assessing 27 studies

Of the **7,928** individuals included in the analysis, **1.0** % expressed regret.

The most common reason for postoperative regret was:

"difficulty/dissatisfaction in life with the new gender role."

What About Regret?



The American Journal of Surgery

Volume 234, August 2024, Pages 68-73



Review Article

A systematic review of patient regret after surgery- A common phenomenon in many specialties but rare within genderaffirmation surgery

Sarah M. Thornton, Armin Edalatpour, Katherine M. Gast 🖰 🖾

Percentage of patients reporting regret ranged from:

0 to 47.1 % in breast reconstruction5.1–9.1 % in breast augmentation10.82–33.3 % in body contouring

In other surgical subspecialties:

30 % following prostatectomy19.5 % following bariatric surgery

Percentage of people with regret from other life choices:

16.2 % getting a tattoo7 % having children

Percentage of patients reporting regret from:

~1 % Gender Affirming Surgery

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Take Home Points

- Gender Identity ≠ Sex (assigned at birth)
- Gender Affirming care is well supported by guidelines and clinical practice!
- Feminizing Hormone Therapy = Estrogen +/- Anti-Androgen therapy
- Masculinizing Hormone Therapy = Testosterone therapy
- Desired changes will take 1-6 months to initially take effect
- Baseline labs then follow up at 3, 6, and 12 months then 1-2x annually
 - Do labs at the 'mid-point'
- Gender Affirming Surgeries exist in Ontario many are MoH covered!

Where to Get Additional Training?



- Project ECHO: Trans & Gender Diverse Healthcare
 - https://camh.echoontario.ca/programs-tgdh/

- Rainbow Health Ontario e-learning:
 - https://learn.rainbowhealthontario.ca/learn
 - 'Mentorship Call' (1st and 3rd Wednesday/month)

- National LGBTQIA+ Health Education Center (USA):
 - https://www.lgbtqiahealtheducation.org/resource s/in/transgender-health/type/webinar/

EDUCATION CENTER

NATIONAL LGBTQIA+ HEALTH

A PROGRAM OF THE FENWAY INSTITUTE

Update on Trans-Competent Primary Care

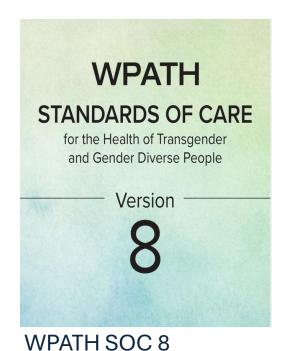
Webinar

Originally presented on 7 November 2022

This course is eligible for CME credit

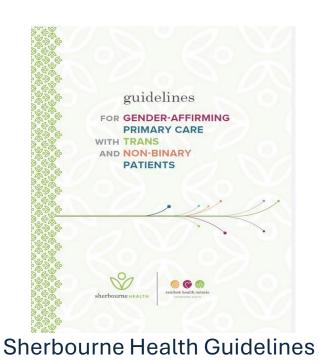
In this talk from the 2022 Advancing Excellence in Transgender Health Care Conference, Dr. Maddie Deutsch provides an update on tran-comptetent primary care.

Thank You for Your Time & Attention!









QUALITY STANDARDS





Gender-Affirming
Care for GenderDiverse People
Care for Adults
Health Quality Ontario



Rainbow Health Ontario (RHO)

UCSF Guidelines Trans Care BC

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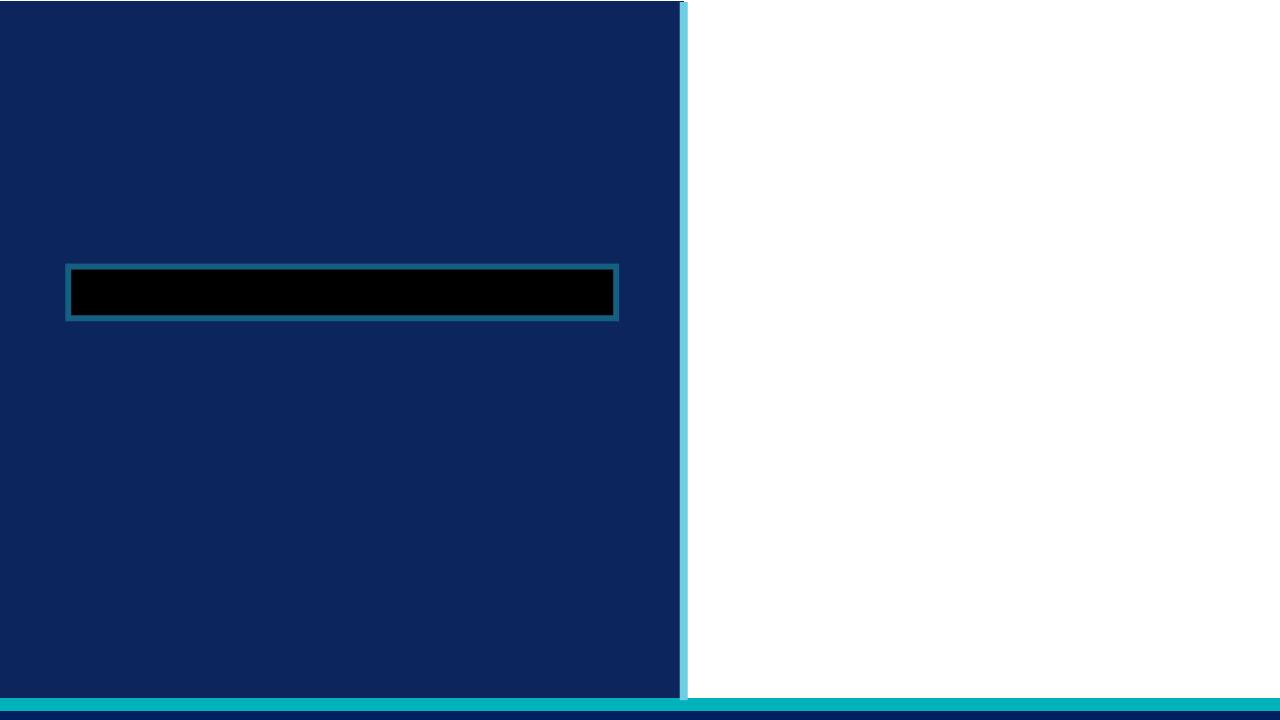
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