

LGBTQIA2S+ Cancer Screening Considerations

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FINANCIAL DISCLOSURES

I have no financial disclosures to make.



LEARNING OBJECTIVES

1. Define important terms and concepts related to the LGBTQIA2S+ population, understand that these terms change frequently, and articulate how to figure out what terms to use with your patients.
2. Articulate how the medical management recommendations for cancer screenings change between the cisgender and gender diverse population, as well as what areas still need more research.
3. Feel comfortable implementing strategies for providing inclusive care to the LGBTQIA2S+ population in your practice.
4. Find resources for further provider and patient education.

DEFINITIONS

LGBTQIA2S+ = An acronym for different gender identities and sexual orientations; lesbian, gay, bisexual, transgender, questioning or queer, intersex, asexual, and Two-Spirit; the + indicates other identities that aren't explicitly included

- **There are many variations of this acronym!

Sex = A category often assigned at birth based on biological attributes (e.g., the appearance of genitalia or secondary sex characteristics)

- Female, male, intersex / variations of sexual characteristics (VSC)

Gender identity = A person's sense of self and how they fit into the world from the perspective of gender, which may or may not align with sex assigned at birth

- Female, male, transgender, cisgender, non-binary

Gender expression = How someone expresses their gender identity (clothes, hair, name, **pronouns**, etc.)

Sexual orientation = An individual's attraction and identity related to romantic and sexual desire

- Lesbian, gay, bisexual, asexual, pansexual, aromantic, polyromantic, etc. ...

DEFINITIONS



Cisgender / Cis = An individual whose gender identity **aligns** with the sex they were assigned at birth

- Someone assigned male at birth (AMAB) who identifies as male (he/him)
- Someone assigned female at birth (AFAB) who identifies as female (she/her)

Gender diverse (GD) = An umbrella term used to describe gender identities beyond the binary framework

- **Transgender / Trans** = An individual whose gender identity **differs** from the sex assigned to them at birth
 - **Trans woman / Transfeminine** = Someone assigned male at birth who identifies as female (she/her)
 - **Trans man / Transmasculine** = Someone assigned female at birth who identifies as male (he/him)
- **Non-binary / Gender-nonconforming** = Someone who identifies as neither male nor female but somewhere in between
 - They/them, she/they, he/they
- **Intersex / Variations of Sexual Characteristics (VSC)** = Someone born with reproductive or sexual anatomy that does not fit typical definitions of male or female; could identify as any of the above





DEFINITIONS

These definitions are constantly changing!

The best you can do is ask a patient how they would like to be addressed, if it's not already listed in their chart.

HEREDITARY CANCER SYNDROMES (HCS)

When a person has one or more genetic changes or variants that cause them to be at an increased risk for one or more types of cancer.

- Hereditary Breast and Ovarian Cancer syndrome (*BRCA1/2*)
- Lynch syndrome (*MLH1, MSH2, MSH6, PMS2, EPCAM*)
- Li-Fraumeni syndrome (*TP53*)
- Polyposis syndromes
 - Familial Adenomatous Polyposis (*APC*)
 - *MUTYH*-Associated Polyposis (*MUTYH*)
 - Peutz-Jeghers syndrome (*STK11*)
 - Juvenile Polyposis syndrome (*BMPR1A, SMAD4*)
 - Serrated Polyposis syndrome (*RNF43*)
- Cowden syndrome (*PTEN*)
- Hereditary Diffuse Gastric Cancer (*CDH1*)
- *ATM, BRCA1/2, CDKN2A, PALB2, NF1, BLM, NTHL1, POLD1, POLE, etc.*

HEREDITARY CANCER SYNDROMES (HCS)

When a person has one or more genetic changes or variants that cause them to be at an increased risk for one or more types of cancer.

- Increased risk for:
 - Breast
 - Ovarian
 - Uterine
 - Prostate
 - Colon
 - Pancreatic
 - Melanoma
 - Brain
 - Kidney
 - Thyroid
 - Leukemia / Lymphoma
 - Etc.

INTRODUCTION





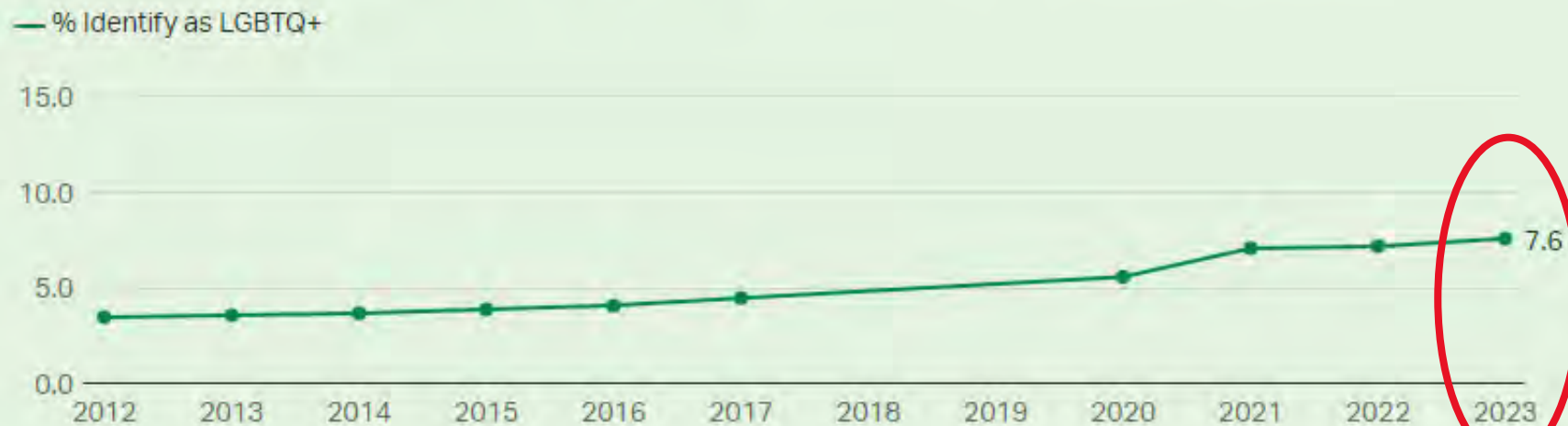
WHY IS THIS IMPORTANT?

- LGBTQIA2S+ community members face significant barriers to accessing quality healthcare
- Healthcare providers need to be aware of community-specific factors so they can provide culturally competent inclusive care
- The LGBTQIA2S+ population in the US is growing and will continue to do so

HOW MANY PEOPLE ARE LGBTQIA2S+?

Americans' Self-Identification as Lesbian, Gay, Bisexual, Transgender, or Something Other Than Heterosexual, 2012-2023

Which of the following do you consider yourself to be? You can select as many as apply. Straight or heterosexual; Lesbian; Gay; Bisexual; Transgender



Respondents who volunteer another identity (e.g., queer; same-gender-loving; pansexual) are recorded as "Other LGBTQ+" by interviewers. These responses are included in the LGBTQ+ estimate.
Data were not collected in 2018 and 2019.
2012-2013 wording: Do you, personally, identify as lesbian, gay, bisexual or transgender?

[Get the data](#) • [Download image](#)

GALLUP

HOW MANY PEOPLE ARE LGBTQIA2S+?

LGBTQ+ Identity, by Generation, 2023

Which of the following do you consider yourself to be? You can select as many as apply. Straight or heterosexual; Lesbian; Gay; Bisexual; Transgender

| | Generation Z | Millennials | Generation X | Baby Boomers | Silent Generation |
|-------------------------------------|--------------|-------------|--------------|--------------|-------------------|
| All adult members of the generation | % | % | % | % | % |
| Total LGBTQ+ | 22.3 | 9.8 | 4.5 | 2.3 | 1.1 |
| Lesbian | 3.0 | 1.3 | 0.7 | 0.7 | 0.2 |
| Gay | 2.6 | 1.6 | 1.3 | 0.9 | 0.4 |
| Bisexual | 15.3 | 5.9 | 1.9 | 0.6 | 0.1 |
| Transgender | 2.8 | 1.1 | 0.5 | 0.2 | 0.4 |
| Other LGBTQ+ | 1.0 | 0.4 | 0.2 | 0.0 | 0.0 |

Sum of categories may exceed the total because respondents can choose multiple identities.

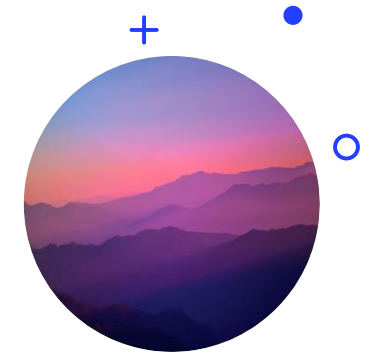
Birth years for each generation: Generation Z (1997-2005), millennials (1981-1996), Generation X (1965-1980), baby boomers (1946-1964), Silent Generation (1945 and earlier).

Based on aggregated data from 2023 Gallup telephone polls.

[Get the data](#) • [Download image](#)

GALLUP

BARRIERS TO INCLUSIVE CARE



- Many GD people don't regularly see a doctor due to stigmatization, discrimination, lack of insurance coverage, living in a rural area, and/or other socioeconomic factors
 - *" 19% [of transgender patients] have reported refusal of care, 28% reported harassment, and 50% were turned off of the healthcare system due to a lack of gender nonconforming providers" (Sterling & Garcia, 2020, p. 2)*
- This population has a lot of general medical mistrust and anxiety, as well as discomfort around organs associated with gender identity
- Many patients may only seek gender-affirming care and not primary care
- GD patients are often unaware of and less likely to complete preventative screenings like colonoscopies, mammograms, and pap smears
- GD people have higher rates of advanced cancer and poorer survival, in addition to other health disparities

BARRIERS TO INCLUSIVE CARE



- Many medical providers know less about GD topics and may feel unprepared to see these patients
- There are very few clinical guidelines for cancer screenings for this population
 - Mostly focused on *BRCA1/2* and Lynch syndrome
 - No set guidelines for any other HCSs, or for cancer screenings for this population in general
- ***We don't know how gender-affirming care and other factors in the GD community may affect cancer risk and risk management***

GENDER-AFFIRMING CARE

- Can include
 - Psychological support,
 - Hormonal treatments,
 - Surgery, and/or
 - Voice therapy
 - **Or some or none** of the above
- Is highly individualized
- Gender-affirming care = GAC
- Gender-affirming surgery = GAS
- Gender-affirming hormone therapy = GAHT





GENDER-AFFIRMING CARE

Transmasculine patients:

- Usual goals:
 - Voice deepening
 - Amenorrhea
 - Increased facial and body hair
 - Redistribution of fat and muscle for more masculine body shape
- Usually achieved through testosterone and/or surgery
- *Some changes are irreversible, some are not*
- *Hormone therapy can be lifelong*
- *Non-binary people might choose lower doses or shorter treatments, ending once desired irreversible results are achieved*

Transfeminine patients:

- Usual goals:
 - Breast growth
 - Decreased libido and erections
 - Reduced muscle mass and body hair
 - Skin softening
 - Body fat redistribution
- Usually achieved through androgen blockers, estrogen, progesterone, and/or surgery



ISSUES TO CONSIDER

- Genetic testing is recommended when results will impact medical decision-making
- Many people with an HCS will have testing in early adulthood long before screenings begin, BUT...
- *GD people often start gender-affirming care long before they consider their future cancer risks*
 - Ex. Puberty blockers, which can be reversible and buy more time for teens to make decisions and provide informed consent
- You might be the first provider who has brought up cancer risk reduction to them; allow for this discussion and/or refer to genetics
 - Having an HCS affects their own care, family members' care, and future family planning considerations

ISSUES TO CONSIDER

- Genetic testing CAN be done for adult-onset HCSs in children / adolescents **if there is an appropriate clinical reason / intervention**, such as gender-affirming care
 - This can ensure patient knows carrier status before transitioning
 - Both ACOG and ASHG agree with this
 - Could also help with insurance coverage for GAS
- If there is a question about whether a patient with an HCS should be on hormones or not...
 - **We have no definitive research to say GAHT increases or decreases cancer risk**
 - Weigh the potential for increased cancer risk versus the intolerable distress of gender dysphoria
 - Keep in mind this population has much higher rates of attempted suicide, poor mental health, drug and alcohol abuse, etc.



CANCER RISK, SURVEILLANCE, AND GAHT FOR GD INDIVIDUALS



GENERAL RECOMMENDATIONS

“There are no prospective data on appropriate cancer risk reduction and/or screening options for transgender, non-binary, or gender diverse individuals who are at average or high risk, regardless of average risk or increased risk.

“Recommendations for risk reduction must be made on a case-by-case basis depending on all of the variables involved.” - NCCN



GENERAL RECOMMENDATIONS

- Healthcare provider needs to know:
 - What decisions / thought has the patient given to GAC?
 - Have they undergone any GAC yet? If so, what?
 - What age did their GAC start, or will start?
 - Use, dosage, and duration of GAHT?
 - Types of GAS done, and with what tissue(s)?
 - Any additional risk factors like family history, smoking, etc.?
 - Have they had genetic testing?



GENERAL RECOMMENDATIONS

- Take an ORGAN INVENTORY
 - “Primary care providers should conduct an organ based routine cancer screening for all transgender patients in accordance with current guidelines as a component of comprehensive primary care. **As a rule, if an individual has a particular body part or organ and otherwise meets criteria for screening based on risk factors or symptoms, screening should proceed regardless of hormone use.** Therefore, an ongoing and thorough medical and surgical history is crucial to determine an individual patient's screening needs.” – UCSF Transgender Cancer Screening Guidelines

CANCER-SPECIFIC RECOMMENDATIONS

Hereditary

- Breast
- Ovarian
- Uterine
- Prostate
- Colon
- Neovagina
- Urothelial

Not Hereditary

- Cervical
- Testicular
- Vulvar
- Penile
- Esophageal
- Anal





NCCN Guidelines Version 3.2025

Genetic/Familial High-Risk Assessment: Breast, Ovarian, Pancreatic, and Prostate

[NCCN Genetic/Familial High-Risk Assessment Panel Members](#)

[Summary of the Guidelines Updates](#)

Principles of Cancer Risk Assessment and Counseling

- [Pre-Test Counseling \(EVAL-A 1 of 11\)](#)
- [Testing Considerations Prior to Testing \(EVAL-A 2 of 11\)](#)
- [Choice of Multigene Testing \(EVAL-A 3 of 11\)](#)
- [Evaluating the Source of Genetic Testing Information \(EVAL-A 4 of 11\)](#)
- [Tumor Genomic Testing: Potential Implications for Germline Testing \(EVAL-A 5 of 11\)](#)
 - ▶ [Circulating Tumor DNA \(ctDNA\)](#)
- [Post-Test Counseling \(EVAL-A 6 of 11\)](#)
 - ▶ [Positive Results](#)
 - ▶ [Negative Results](#)
 - ▶ [Variants of Uncertain Significance](#)
- [Pedigree: First-, Second-, and Third-Degree Relatives of Proband \(EVAL-B\)](#)

Hereditary Testing Criteria

- [General Testing Criteria \(CRIT-1\)](#)
- [Testing Criteria for High-Penetrance Breast Cancer Susceptibility Genes \(CRIT-2\)](#)
- [Testing Criteria for Ovarian Cancer Susceptibility Genes \(CRIT-4\)](#)
- [Testing Criteria for Pancreatic Cancer Susceptibility Genes \(CRIT-5\)](#)
- [Testing Criteria for Prostate Cancer Susceptibility Genes \(CRIT-6\)](#)
- [Testing Criteria for Li-Fraumeni Syndrome \(CRIT-7\)](#)
- [Testing Criteria for Cowden Syndrome/PTEN Hamartoma Tumor Syndrome \(CRIT-8\)](#)

Gene Summary: Risks and Management

- [Testing Criteria Met \(GENE-1\)](#)
- [Cancer Risk Management Based on Genetic Test Results \(GENE-A\)](#)
- [Autosomal Recessive Risk in Cancer Genes – Multigene Panel Testing \(GENE-B\)](#)

Management/Screening

- [BRCA Pathogenic/Likely Pathogenic Variant-Positive Management \(BRCA-A\)](#)
- [Pancreatic Cancer Screening \(PANC-A\)](#)
- [Li-Fraumeni Syndrome Management \(LIFR-A\)](#)
- [Cowden Syndrome/PHTS Management \(COWD-A\)](#)

The NCCN Guidelines® are a statement of evidence and consensus of the authors regarding their views of currently accepted approaches to treatment. Any clinician seeking to apply or consult the NCCN Guidelines is expected to use independent medical judgment in the context of individual clinical circumstances to determine any patient's care or treatment. The National Comprehensive Cancer Network® (NCCN®) makes no representations or warranties of any kind regarding their content, use or application and disclaims any responsibility for their application or use in any way. The NCCN Guidelines are copyrighted by National Comprehensive Cancer Network®. All rights reserved. The NCCN Guidelines and the illustrations herein may not be reproduced in any form without the express written permission of NCCN. ©2025.

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NCCN Categories of Evidence and Consensus: All recommendations are category 2A unless otherwise indicated.

See [NCCN Categories of Evidence and Consensus](#).

• [Breast, Ovarian, Uterine, and Prostate Cancer Risk Reduction Strategies for Transgender, Non-Binary and Gender Diverse People with Hereditary Cancer Syndromes \(TNBGD-1\)](#)

• [Summary of Cancers and/or Syndromes Included/Mentioned in Other NCCN Guidelines \(SUMM-1\)](#)

• [Abbreviations \(ABBR-1\)](#)

• For chemoprevention options, see [NCCN Guidelines for Breast Cancer Risk Reduction](#).

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- Breast, Ovary, Uterine, and Prostate Cancer Risk Reduction Strategies for Transgender, Non-Binary and Gender Diverse People with Hereditary Cancer Syndromes (TNBGD-1)
- Summary of Genes and/or Syndromes Included/ Mentioned in Other NCCN Guidelines (SUMM-1)
- Abbreviations (ABBR-1)
- For chemoprevention options, see NCCN Guidelines for Breast Cancer Risk Reduction.

Creating Breast and Gynecologic Cancer Guidelines for Transgender Patients With *BRCA* Mutations

Bronwyn S. Bedrick, MD MSCI, Timothee F. Fruhauf, MD MPH, Stephen J. Martin, MD, and James S. Ferriss, MD

Bedrick, B. S., Fruhauf, T. F., Martin, S. J., & Ferriss, J. S. (2021). Creating breast and gynecologic cancer guidelines for transgender patients with BRCA mutations. Obstetrics and Gynecology, 138(6), 914. <https://doi.org/10.1097/AOG.0000000000004597>



BREAST CANCER

- General risk to get breast cancer:

Cisgender men < GD people < Cisgender women
Lowest risk Highest risk

- There is insufficient evidence to say whether GAHT alters breast cancer risk
 - *But we also don't have long-term GAHT usage data
- Breast cancer risk models can still be considered for care management regardless of GAHT for trans men / those AFAB; no risk models for GD people at this time

BREAST CANCER – TRANS WOMEN

Table 1. Breast Screening Guidelines for Transgender Individuals

| Population | Gender-Affirming Care | <i>BRCA1</i> or <i>BRCA2</i> Carrier Status | RRM | Breast Cancer Screening Recommendation |
|-------------------|-----------------------------|---|-----|--|
| Transgender women | None | Not carrier | NA | NA |
| | | Carrier | No | Follow NCCN screening for cisgender male <i>BRCA1</i> or <i>BRCA2</i> mutation carriers |
| | | | Yes | Consider yearly clinical examination |
| | Hormonal, surgical, or both | Not carrier | NA | Screening mammography every other year starting at age 50 y and after 5–10 years of hormonal treatment |
| | | Carrier | No | Individualized based on transition age, family history, preexisting gynecomastia |
| | | | Yes | Yearly clinical examinations |

BREAST CANCER – TRANS WOMEN

- GAS could include breast augmentation
 - Bedrick et al (2021) recommends offering trans women who are *BRCA1/2+* risk-reducing mastectomy to remove native breast tissue before breast augmentation
- GAHT with estrogens and anti-androgens increases breast tissue; this should not be described as gynecomastia
- Trans women are more likely to have dense breast tissue, an independent breast cancer risk factor
- Absolute risk for breast cancer is still relatively low
- Estrogen and anti-androgens are not contraindicated even though they may increase breast cancer risk
- Very little data exists about screening trans women on GAHT with an HCS; NCCN says to screen them similarly to cis males with an HCS

BREAST CANCER – TRANS MEN

Table 1. Breast Screening Guidelines for Transgender Individuals

| Population | Gender-Affirming Care | <i>BRCA1</i> or <i>BRCA2</i> Carrier Status | RRM | Breast Cancer Screening Recommendation |
|-----------------|-----------------------------|---|-----|---|
| Transgender men | None | Not carrier | NA | Follow NCCN screening for cisgender women |
| | | Carrier | No | Follow NCCN screening for cisgender female <i>BRCA1</i> or <i>BRCA2</i> mutation carriers |
| | | | Yes | Follow NCCN screening for cisgender female <i>BRCA1</i> or <i>BRCA2</i> mutation carriers |
| | Hormonal, surgical, or both | Not carrier | NA | If postmastectomy, consider annual examination, breast ultrasonogram ⁸ ; otherwise, follow screening for cisgender women |
| | | Carrier | No | Follow NCCN screening for cisgender female <i>BRCA1</i> or <i>BRCA2</i> mutation carriers |
| | | | Yes | Yearly clinical examinations starting at 35 y |

RRM, risk-reducing mastectomy; NA, not applicable; NCCN, National Comprehensive Cancer Network.

BREAST CANCER – TRANS MEN

- Most common GAS is chest surgery of some sort
 - When discussing with the patient, may want to call it “chest cancer” instead of “breast cancer”
- “Top surgery” ≠ Risk-reducing mastectomy (RRM)
- **Top surgery** can include several options, but usually some breast tissue is left for cosmetic purposes
 - Still have potential to get breast cancer in that tissue
 - If patient only has top surgery and then finds out he has a HCS, further surgery may be needed to remove remaining tissue
- **Risk-reducing mastectomy** gets rid of as much of the breast tissue as possible
 - If patient has had RRM, there is very little tissue left to screen; have discussion with patient about unknown residual risks of breast cancer / limitations of screening
- For trans men and non-binary individuals with an HCS who have had top surgery or no surgery at all, NCCN recommends breast screening begin at an earlier age and include mammography and breast MRI
- ER+ tumors have been linked to long-term estrogen exposure in postmenopausal cis women; could also apply to trans men

ACR APPROPRIATENESS CRITERIA

The American College of Radiology has released a transgender breast cancer screening guide that can be used to guide care

Expert Panel on Breast Imaging, Brown, A., Lourenco, A. P., Niell, B. L., Cronin, B., Dibble, E. H., DiNome, M. L., Goel, M. S., Hansen, J., Heller, S. L., Jochelson, M. S., Karrington, B., Klein, K. A., Mehta, T. S., Newell, M. S., Schechter, L., Stuckey, A. R., Swain, M. E., Tseng, J., ... Moy, L. (2021). ACR Appropriateness Criteria® Transgender Breast Cancer Screening. Journal of the American College of Radiology: JACR, 18(11S), S502–S515. <https://doi.org/10.1016/j.jacr.2021.09.005>

Variant 1: Breast cancer screening. Transfeminine (male-to-female) patient, 40 years of age or older with past or current hormone use equal to or greater than 5 years. Average-risk patient.

| Procedure | Appropriateness Category | Relative Radiation Level |
|---|--------------------------|--------------------------|
| Digital breast tomosynthesis screening | May Be Appropriate | ⊕⊕ |
| Mammography screening | May Be Appropriate | ⊕⊕ |
| US breast | Usually Not Appropriate | ○ |
| MRI breast without and with IV contrast | Usually Not Appropriate | ○ |
| MRI breast without IV contrast | Usually Not Appropriate | ○ |

Variant 2: Breast cancer screening. Transfeminine (male-to-female) patient, 25 to 30 years of age or older with past or current hormone use equal to or greater than 5 years. Higher-than-average risk (patient with personal history of breast cancer or chest irradiation at 10 to 30 years of age, patient with genetic predisposition to breast cancer, patient with family history of breast or ovarian cancer, and untested patient with first-degree relative with genetic predisposition to breast cancer).

| Procedure | Appropriateness Category | Relative Radiation Level |
|---|--------------------------|--------------------------|
| Digital breast tomosynthesis screening | Usually Appropriate | ⊕⊕ |
| Mammography screening | Usually Appropriate | ⊕⊕ |
| US breast | Usually Not Appropriate | ○ |
| MRI breast without and with IV contrast | Usually Not Appropriate | ○ |
| MRI breast without IV contrast | Usually Not Appropriate | ○ |

Variant 3: Breast cancer screening. Transfeminine (male-to-female) patient with no hormone use (or hormone use less than 5 years) at any age. Average-risk patient.

| Procedure | Appropriateness Category | Relative Radiation Level |
|---|--------------------------|--------------------------|
| US breast | Usually Not Appropriate | ○ |
| Digital breast tomosynthesis screening | Usually Not Appropriate | ⊕⊕ |
| Mammography screening | Usually Not Appropriate | ⊕⊕ |
| MRI breast without and with IV contrast | Usually Not Appropriate | ○ |
| MRI breast without IV contrast | Usually Not Appropriate | ○ |



UTERINE & OVARIAN CANCER

- ACOG recommends hysterectomy (TAH) with or without bilateral salpingo-oophorectomy (BSO) for those with gender dysphoria who want it; however, many trans men do not do this
 - Very important to know if your patient still has their uterus &/or ovaries or not!
- There are no effective screening guidelines for uterine or ovarian cancer for cis women
 - Transvaginal ultrasound with or without random endometrial biopsies is sometimes done; utility is questionable
- The effect of GAHT on uterine tissue is unknown, BUT additional androgens may increase circulating estrogen levels and “pose a risk” to the uterus, per NCCN (TNBGD-2)
- There is no data on the effect of medical uterine or ovarian suppression on uterine or ovarian cancer risk

Bedrick, B. S., Fruhauf, T. F., Martin, S. J., & Ferriss, J. S. (2021). Creating breast and gynecologic cancer guidelines for transgender patients with BRCA mutations. *Obstetrics and Gynecology*, 138(6), 914. <https://doi.org/10.1097/AOG.0000000000004597>

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Gupta, S., et. al. (2025, April 2). *NCCN Clinical Practice Guidelines in Oncology: Genetic/Familial High-Risk Assessment: Colorectal, Endometrial, and Gastric Cancers, Version 4.2025*. NCCN. <https://www.nccn.org/guidelines/guidelines-detail?category=2&id=1544>



UTERINE & OVARIAN CANCER

- If someone has a HCS that increases their risk for uterine or ovarian cancer, risk-reducing TAH &/or BSO is recommended (ex. *BRCA1/2*) or should be strongly considered (ex. Lynch) depending on the HCS/gene & organ
- If a trans man still has his uterus, he should be counseled on the early warning signs of uterine cancer
 - Endometrial biopsy recommended for bleeding occurring after GAHT-induced amenorrhea
- RR TAH/BSO can be considered at an earlier age to alleviate gender dysphoria
 - Patients should be counseled about adverse effects of RRSO before natural menopause (outlined in NCCN)
 - Would have to be on hormone therapy for life to mitigate negative bone / cardiovascular effects

Bedrick, B. S., Fruhauf, T. F., Martin, S. J., & Ferriss, J. S. (2021). Creating breast and gynecologic cancer guidelines for transgender patients with *BRCA* mutations. *Obstetrics and Gynecology*, 138(6), 914. <https://doi.org/10.1097/AOG.0000000000004597>

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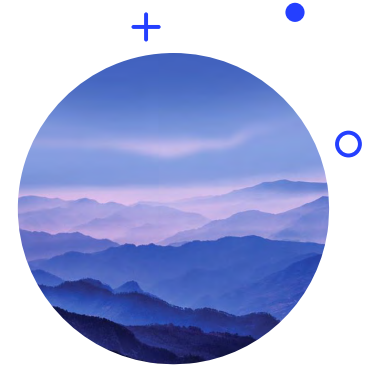
Hodan, R., Rodgers-Fouche, L., Chittenden, A., Dominguez-Valentin, M., Ferriss, J., Gima, L., Hamnvik, O.-P. R., Idos, G. E., Kline, K., Koeller, D. R., Long, J. M., McKenna, D., Muller, C., Thoman, M., Wintner, A., Bedrick, B. S., & on behalf of the Collaborative Group of the Americas on Inherited Gastrointestinal Cancer. (2023). Cancer surveillance for transgender and gender diverse patients with Lynch syndrome: A practice resource of the Collaborative Group of the Americas on Inherited Gastrointestinal Cancer. *Familial Cancer*, 22(4), 437–448. <https://doi.org/10.1007/s10689-023-00341-4>



UTERINE & OVARIAN CANCER

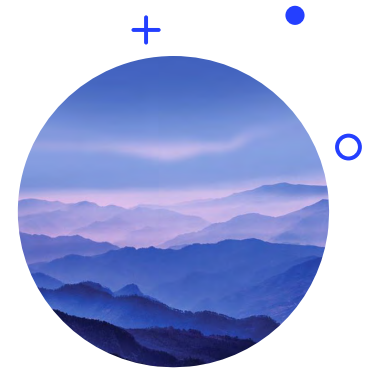
- Screening guidelines for *BRCA1/2*+ cis women after BSO include annual abdominal and pelvic exam and serum CA-125 for 10 years (debatable effectiveness)
 - Offer trans men the same screening
- If transvaginal ultrasound isn't possible due to vaginal atrophy, consider transrectal ultrasound and/or use of topical vaginal estradiol before exam
- Other ovarian cancer risk factors to consider:
 - Trans men often have less childbearing, oral contraceptive pill use
 - Trans men's risk may be similar to nulliparous cis women's risk

PROSTATE CANCER



- Prostate is usually not removed during GAC
 - Internal location can shift due to neovagina
 - Estrogen causes the prostate to shrink
- Cis men are screened via prostate specific antigen (PSA) level & digital rectal exam (DRE)
- NCCN still recommends using PSA even though GAHT can reduce effectiveness
 - The abnormal PSA threshold is lowered to 1.0 ng/ml for trans women
 - This number is somewhat arbitrary; each patient should establish their own baseline PSA to identify future deviations
- Prostate cancer incidence is lower in trans women than in cis men

PROSTATE CANCER



- Follow gene-specific screening guidelines for cis men who have a hereditary cancer risk
 - Ex. Consider MRI for someone with Lynch syndrome with elevated PSA or abnormal prostate exam
- DRE may be harder to do if the prostate is inaccessible; can try to access transvaginally, but transrectally or transperineally may work better
- There are conflicting reports on whether estrogen is protective against prostate cancer or not
- For *BRCA1/2*+ trans women, Crowley et. al (2023) argues that any change in DRE or detectable PSA should prompt a diagnostic prostate biopsy

Crowley, F., Mihalopoulos, M., Gaglani, S., Tewari, A. K., Tsao, C.-K., Djordjevic, M., Kyprianou, N., Purohit, R. S., & London, D. J. (2023). Prostate cancer in transgender women: Considerations for screening, diagnosis and management. *British Journal of Cancer*, 128(2), 177–189. <https://doi.org/10.1038/s41416-022-01989-y>

Daly, M., Pal, T., et. al (2025, March 6). NCCN Clinical Practice Guidelines in Oncology: Genetic/Familial High-Risk Assessment: Breast, Ovarian, Pancreatic, and Prostate—Version 3.2025. NCCN. <https://www.nccn.org/guidelines/guidelines-detail?category=2&id=1545>

Sterling, J., Carbonella, J., Jones, T., Hanchuk, S., Kelly, P., & Garcia, M. M. (2023). Cancer screening for transgender individuals: Guidelines, best practices, and a proposed care model. *The Urologic Clinics of North America*, 50(4), 563–576. <https://doi.org/10.1016/j.ucl.2023.06.014>

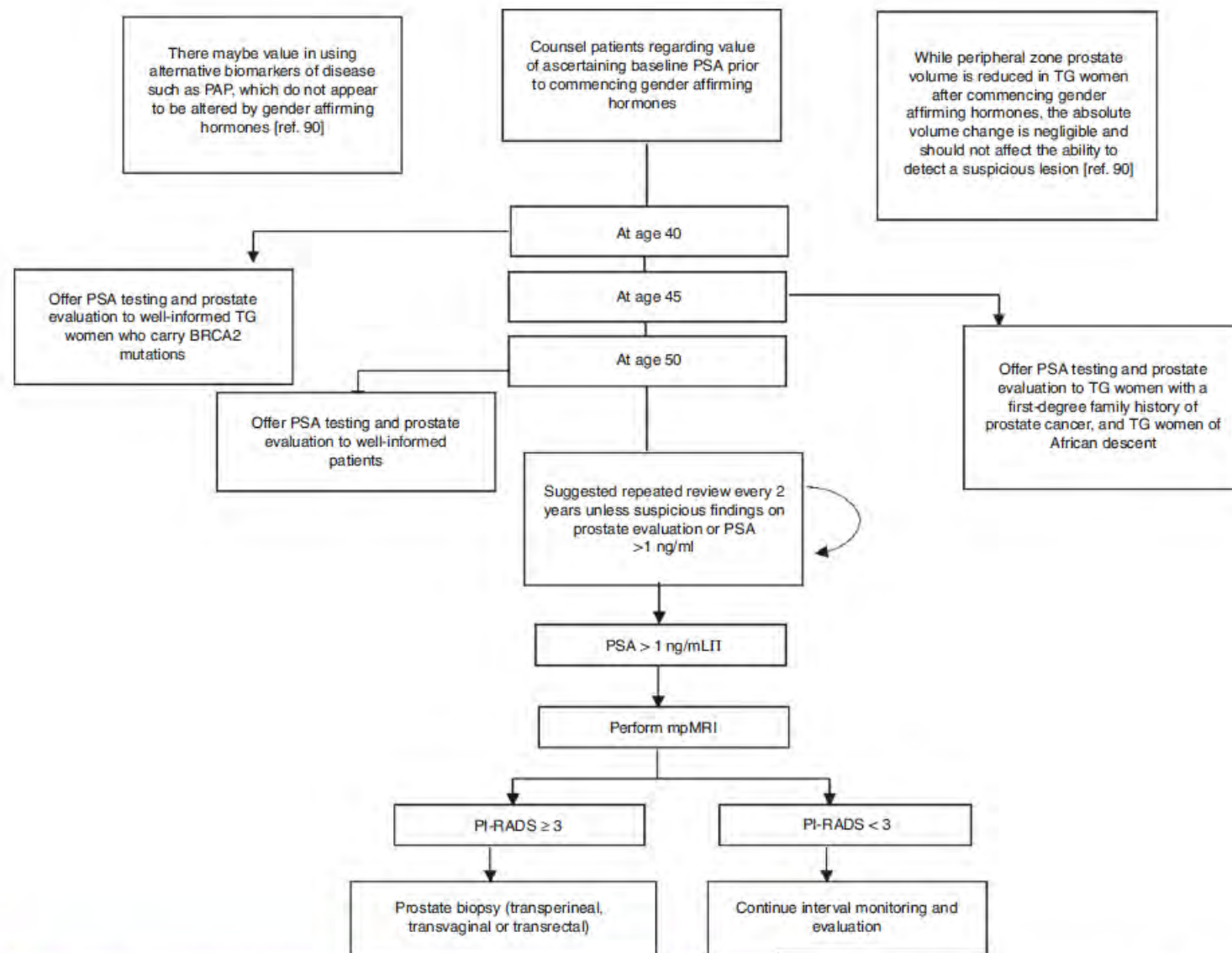


Fig. 2 Proposed approach to prostate cancer screening in transgender women. It = the choice of a PSA threshold of 1.0 ng/ml is an arbitrary one widely in use, and should be revised as a broader evidence base is established. Establishing a baseline PSA in each individual and identifying deviations from baseline, is likely to be of greater utility than an absolute cut-off value applied to everybody.



Cancer surveillance for transgender and gender diverse patients with Lynch syndrome: a practice resource of the Collaborative Group of the Americas on Inherited Gastrointestinal Cancer

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Abstract

Transgender and gender diverse (TGD) populations with hereditary cancer syndromes face unique obstacles to identifying and obtaining appropriate cancer surveillance and risk-reducing procedures. There is a lack of care provider knowledge about TGD health management. Lynch syndrome (LS) is one of the most common hereditary cancer syndromes, affecting an estimated 1 in 279 individuals. There are no clinical guidelines specific for TGD individuals with LS, highlighting a need to improve the quality of care for this population. There is an urgent need for cancer surveillance recommendations for TGD patients. This commentary provides recommendations for cancer surveillance, risk-reducing strategies, and genetic counseling considerations for TGD patients with LS.

Keywords Transgender and gender diverse · Cancer · Lynch syndrome · Surveillance · Management

Table 1 Surveillance and management recommendations for TGD individuals with Lynch Syndrome

| | LS management for CG patients ^a | | Additional considerations for TGD patients | |
|---------------------|--|---|---|--|
| General | Elicit organ inventory before making recommendations | | | |
| | Apply principles of trauma-informed care [127] | | | |
| Site/Organ | <i>MLH1/MSH2/EPCAM</i> | <i>MSH6/PMS2</i> | Transfeminine | Transmasculine |
| CRC | Colonoscopy at age 20-25y or 2-5y prior to earliest CRC and repeat every 1-2y Consider chemoprevention | Colonoscopy at age 30-35y or 2-5y prior to earliest CRC and repeat every 1-3y Consider chemoprevention | Use of sigmoid colon for neovagina creation is contraindicated. For those who are diagnosed with LS after GAS with use of sigmoid colon, pelvic exam and colposcopy recommended at interval paralleling colorectal surveillance | Gynecologic screening can be offered at the time of colonoscopy sedation to minimize pelvic exam anxiety |
| Endometrial | Consider EMB at age 30-35y and repeat every 1-2y Consider post-menopausal transvaginal ultrasound Consider TAH Education on symptoms ^b | | N/A | EMB for bleeding after GAHT induced amenorrhea Consider short-term vaginal estradiol to reduce vaginal atrophy and discomfort |
| Ovarian | Consider BSO ^c | | N/A | If undergoing gender-affirming hysterectomy, consider BSO concurrently |
| Urothelial | Annual urinalysis beginning age 30-35y | | | |
| Gastric/small bowel | EGD at age 30–40 and repeat every 2-4y | | | |

Table 1 Surveillance and management recommendations for TGD individuals with Lynch Syndrome

| | LS management for CG patients ^a | | Additional considerations for TGD patients | |
|------------|---|------------------|---|---|
| General | Elicit organ inventory before making recommendations | | | |
| | Apply principles of trauma-informed care [127] | | | |
| Site/Organ | <i>MLH1/MSH2/EPCAM</i> | <i>MSH6/PMS2</i> | Transfeminine | Transmasculine |
| Pancreas | Alternate annual MRI/MRCP and/or EUS beginning at age 50y if there is a family history of pancreatic cancer in a 1 st or 2 nd degree relative (or 10y young than the youngest age of pancreatic cancer diagnosis in the family) | | | |
| Prostate | Consider annual PSA and DRE beginning at age 40 | | Awareness of vaginoplasty, intestinal interposition, or perineal flap which may affect palpation during DRE Consider MRI for elevated PSA or abnormal prostate exam or finding raising concern for prostate malignancy PSA > 1 ng/mL is considered abnormal with testosterone suppression | N/A |
| Breast | Not enough evidence to support increased screening | | Follow local breast cancer screening guidelines ^d developed for CG women for those who have taken estradiol for ≥ 5 years | No imaging indicated after bilateral mastectomies. Otherwise, follow local breast cancer screening guidelines ^d and/or based on family history |

BSO bilateral salpingo-oophorectomy, *CG* cisgender, *CRC* colorectal cancer, *DRE* digital rectal exam, *EGD* Esophagogastroduodenoscopy, *EMB* endometrial biopsy, *EUS* endoscopic ultrasound, *GAHT* gender affirming hormone therapy, *GAS* gender affirming surgery, *LS* Lynch syndrome, *mpMRI* multiparametric magnetic resonance imaging, *MRCP* Magnetic resonance cholangiopancreatography, *MRI* Magnetic resonance imaging, *N/A* not applicable, *PSA* prostate specific antigen, *TAH* total abdominal hysterectomy, *y* year(s)

^aAdapted from NCCN Genetic/Familial High-Risk Assessment: Colorectal Version 2.2022[21]

^bWarning signs of endometrial cancer include abnormal vaginal bleeding, especially postmenopausal bleeding, bloating, changes to bowel or bladder habits, weight loss, abdominal or pelvic pain, increased satiety, increased abdominal girth

^cThere is not enough evidence to recommend BSO for individuals with *MSH6/PMS2* mutations

^d[102]



COLORECTAL CANCER

- **Transfeminine** patients with LS or increased CRC risk:
 - Don't use the sigmoid colon for **neovaginal creation**
 - If already done, pelvic exam & colposcopy at time of colonoscopy may be necessary
 - Intestinal mucosa tissue may be more susceptible to STIs
 - Persistent HPV infection has been reported in patients with a neovagina
- **Transmasculine** patients:
 - Gynecologic screening can be offered at time of colonoscopy sedation to decrease pelvic exam discomfort / anxiety
- No current data suggest that GAHT increases or decreases risk for CRC



NEOVAGINAL CANCER

- Of note, another possible contribution besides HCSs can be chemical stimulants and irritants in semen
- Some say neovaginas are prone to precancerous lesions and so patients should get appropriate cancer screening
 - Routine gynecological examination to assess for HPV
 - HPV vaccine is recommended for everyone by age 26

UROTHELIAL CANCER

- Patients with *MSH2*-associated Lynch syndrome are at an increased risk for kidney, ureter, and/or bladder cancer
- In general (not related to Lynch syndrome),
 - Trans women have a higher incidence of bladder cancer than cis women, but a similar incidence to cis men
 - GD patients have a poorer bladder cancer survival rate than cis counterparts
- Possible contributors to this could include:
 - Non-genetic factors like tobacco use
 - Androgenic hormones
 - An X-linked gene, *KDM6A*, acting as a protective factor



CERVICAL CANCER

- Not hereditary; usually caused by HPV
 - LGBTQIA2S+ population has higher HPV+, HIV+ rates
- If a trans man still has a uterus, he will usually require cervical cancer screening
- Many trans men will avoid gynecological care such as pap smears as it is dysphoric for them
- Testosterone can cause vaginal atrophy which makes pap smears even more uncomfortable and more likely to fail

Hodan, R., Rodgers-Fouche, L., Chittenden, A., Dominguez-Valentin, M., Ferriss, J., Gima, L., Hamnvik, O.-P. R., Idos, G. E., Kline, K., Koeller, D. R., Long, J. M., McKenna, D., Muller, C., Thoman, M., Wintner, A., Bedrick, B. S., & on behalf of the Collaborative Group of the Americas on Inherited Gastrointestinal Cancer. (2023). Cancer surveillance for transgender and gender diverse patients with Lynch syndrome: A practice resource of the Collaborative Group of the Americas on Inherited Gastrointestinal Cancer. *Familial Cancer*, 22(4), 437–448. <https://doi.org/10.1007/s10689-023-00341-4>

Iwamoto, S. J., Grimstad, F., Irwig, M. S., & Rothman, M. S. (2021). Routine screening for transgender and gender diverse adults taking gender-affirming hormone therapy: A narrative review. *Journal of General Internal Medicine*, 36(5), 1380–1389. <https://doi.org/10.1007/s11606-021-06634-7>

CERVICAL CANCER



- Can offer to do gynecological procedures such as pap smears, pelvic exams, endometrial biopsies, etc. during colonoscopies / endoscopies...
 - Can offer topical vaginal estradiol 2 weeks prior to soften area (won't affect their hormone therapy)
 - Can administer an anxiolytic prior to exam
 - Offer patients use of a mirror and/or to have a support person in the room
- Can offer high-risk HPV DNA primary screening to patients over age 25; patients can self-swab if necessary

Hodan, R., Rodgers-Fouche, L., Chittenden, A., Dominguez-Valentin, M., Ferriss, J., Gima, L., Hamnvik, O.-P. R., Idos, G. E., Kline, K., Koeller, D. R., Long, J. M., McKenna, D., Muller, C., Thoman, M., Wintner, A., Bedrick, B. S., & on behalf of the Collaborative Group of the Americas on Inherited Gastrointestinal Cancer. (2023). Cancer surveillance for transgender and gender diverse patients with Lynch syndrome: A practice resource of the Collaborative Group of the Americas on Inherited Gastrointestinal Cancer. *Familial Cancer*, 22(4), 437–448. <https://doi.org/10.1007/s10689-023-00341-4>

Iwamoto, S. J., Grimstad, F., Irwig, M. S., & Rothman, M. S. (2021). Routine screening for transgender and gender diverse adults taking gender-affirming hormone therapy: A narrative review. *Journal of General Internal Medicine*, 36(5), 1380–1389. <https://doi.org/10.1007/s11606-021-06634-7>



TESTICULAR CANCER

- Not hereditary
- If a trans woman still has testicles, follow the same guidelines as cis men (annual physical exam)

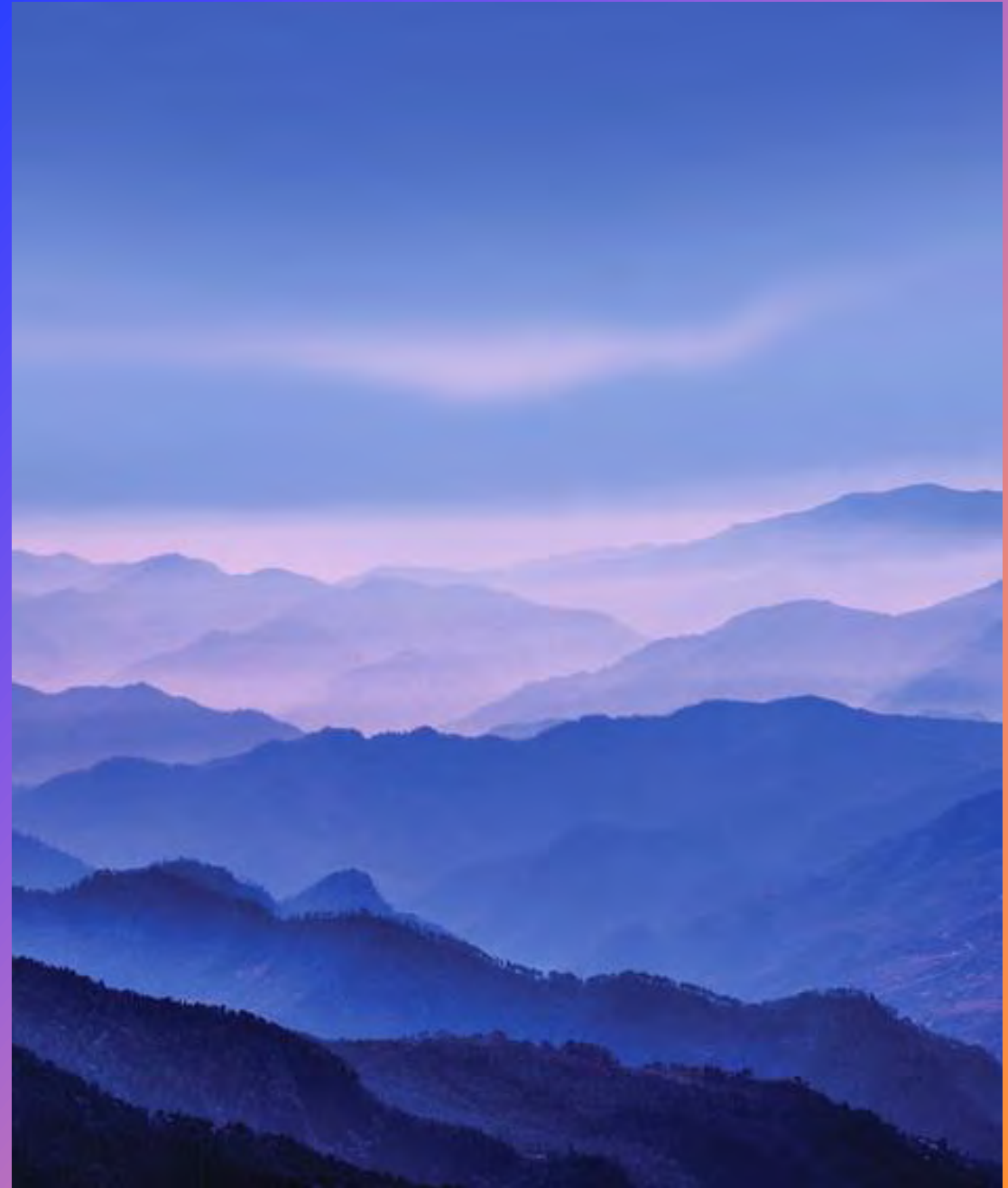
VULVAR / PENILE CANCER

- Not hereditary
- No screening guidelines exist for cis people; only one case of vulvar cancer in a trans man has been reported and none of penile cancer

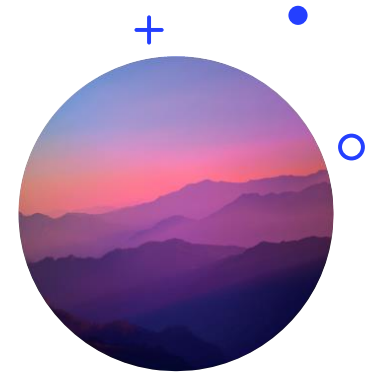


ESOPHAGEAL CANCER

- HPV is a possible mechanism for esophageal squamous cell carcinoma
- Since HPV infection is more prevalent in the LGBTQIA2S+ population, they could be eligible for an esophageal screening program



ANAL CANCER



- Usually caused by HPV / HIV
 - Men who have sex with men, especially if they are HIV+, are at highest risk
 - Also at risk: those with HPV+, higher # sexual partners, genital warts
- No standard guidelines for anal cancer screening, but...
- HPV vaccination is recommended for everyone by age 26
- All HIV+ adults should have an annual digital anorectal exam (DARE)
 - People <35 who show signs of anal cancer during DARE should undergo standard anoscopy
 - Older people should undergo high resolution anoscopy (HRA) if they are...
 - Men who have sex with men
 - Transfeminine people over 35
 - All other HIV+ people over 45
- Can perform surveillance annually for HIV+ males and every 3-6 months for people with low- or high-grade squamous intraepithelial lesions

Coelho, R., Gonçalves, R., Mendes, F., & Macedo, G. (2024). Gastroenterology healthcare in LGBTQ+ individuals. *European Journal of Gastroenterology & Hepatology*, 36(9), 1059–1067. <https://doi.org/10.1097/MEG.0000000000002808>

HIV clinical guidelines now recommend high resolution anoscopy as part of anal cancer screening program for people with HIV. (n.d.). National Institutes of Health. Retrieved October 10, 2024, from <https://oar.nih.gov/update-clinical-guidelines-high-resolution-anoscopy-anal-cancer-screening>



OTHER RISK FACTORS

Other risk factors for cancer can be higher in LGBTQIA2S+ populations:

- Smoking / Tobacco use
 - 16% of LGB adults, 4% of LGB youth, and 5% of trans youth smoke compared to 12% heterosexual adults and 1% heterosexual / cisgender youth
- Excessive alcohol use
 - 14% bi women consume >7 drinks/wk vs. 6% heterosexual women
- Higher BMI
 - 68% of LB women vs 61% of heterosexual women
- HPV or HIV infections

MORE RESEARCH IS NEEDED

- That is all the info we currently have about how to change screenings for LGBTQIA2S+ patients with an increased risk for cancer.
- More equitable research is needed, particularly around hormone therapy!





INCREASING SAFETY AND COMFORT

BEFORE THE APPOINTMENT

- Intake forms should include *“chosen name, pronouns, gender identity, sex assigned at birth, sexual orientation as well as marital, partnership, and family status”* (Hodan et. al, p. 442)

Hereditary Cancer Clinic Family History Questionnaire



Part A: Basic Information

| | | | | |
|--|--|------|---|-----------------------|
| Last Name | First Name | M.I. | Maiden Name | Sex assigned at birth |
| | | | | |
| Date of Birth | Phone Number | | | Pronouns |
| | Home: | | Cell: | |
| Mailing Address | | City | State | Zip Code |
| | | | | |
| E-mail address | Referring Physician | | Hospital or Clinic | |
| | | | | |
| What do you consider to be your race/ethnicity? (check all that apply) | | | | |
| <input type="checkbox"/> African-American/Black | <input type="checkbox"/> Middle Eastern/North African/West Asian | | <input type="checkbox"/> South Asian | |
| <input type="checkbox"/> East/Southeast Asian | <input type="checkbox"/> Native American/Alaska Native/First Nations | | <input type="checkbox"/> White/Caucasian | |
| <input type="checkbox"/> Latinx/Hispanic | <input type="checkbox"/> Native Hawaiian/Pacific Islander | | <input type="checkbox"/> Prefer not to answer | |
| <input type="checkbox"/> Other | | | | |

BEFORE THE APPOINTMENT

Part E: Family History Information

| | |
|---|---|
| What is your mother's ancestry/country of origin? (German, Dominican Republic, Nigerian, Russian, etc.) | What is your father's ancestry/country of origin? (German, Dominican Republic, Nigerian, Russian, etc.) |
| <input type="text"/> | <input type="text"/> |
| Are any of your relatives of Ashkenazi Jewish descent? <input type="checkbox"/> Yes <input type="checkbox"/> No | |
| How many biological children do you have? <input type="text"/> | Number of sons <input type="text"/> Number of daughters <input type="text"/> Number of non-binary children <input type="text"/> |
| Please list the first name, age, and any health concerns/diagnoses for your children: | <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> |
| How many biological siblings do you have? <input type="text"/> Number of brothers <input type="text"/> Number of sisters <input type="text"/> Number of non-binary siblings <input type="text"/> | If any siblings have a different mother or father: How many have the same mother as you? <input type="text"/> <input type="text"/> <input type="text"/> (brothers) (sisters) (non-binary) How many have the same father as you? <input type="text"/> <input type="text"/> <input type="text"/> (brothers) (sisters) (non-binary) |
| How many biological siblings does your mother have? <input type="text"/> | Number of brothers <input type="text"/> Number of sisters <input type="text"/> Number of non-binary siblings <input type="text"/> |
| How many biological siblings does your father have? <input type="text"/> | Number of brothers <input type="text"/> Number of sisters <input type="text"/> Number of non-binary siblings <input type="text"/> |

Part D: For Female Patients and **Patients assigned Female at birth**

BEFORE THE APPOINTMENT

- Use EHRs that allow inclusive documentation of GD patients

Gender Identity

Would you like to provide information about your Gender Identity?

Yes No

Autofill with default responses:

female

male

Can you confirm your sex assigned at birth?

Female

Male

Unknown

Not recorded on birth certificate

Choose not to disclose

Uncertain

Patient's gender identity:

Female

Male

Transgender Female / Male-to-Female

Transgender Male / Female-to-Male

Other

Choose not to disclose

What gender pronoun do you use?



she/her/hers

he/him/his

they/them/theirs

patient's name

decline to answer

unknown

BEFORE THE APPOINTMENT



Gender identity: Male
Legal sex: Male
Sex assigned at birth: Female
It may be possible for this patient to become pregnant



| | | |
|--------------------|-----------------|-----------------------|
| Legal Sex | Gender Identity | Sex Assigned at Birth |
| Male | Non-Binary | Male |
| Sexual Orientation | | |
| Bisexual | | |

Gender identity: Male
Legal sex: Female
Sex assigned at birth: Female
It may be possible for this patient to become pregnant



- GD patients may have **lower insurance coverage** and/or **higher rates of insurance denials** for services (why would a guy need a pap smear? etc.)
 - Additional documentation may be necessary



BEFORE THE APPOINTMENT

- Educate patients about:
 - The importance of sharing their sexual orientation / gender identity (SOGI) data
 - Institutional privacy policies
- Ask if patient has medical records under a different name rather than asking what their name was prior to transitioning (“deadname”)
- Avoid gendered terms / places / items like “women’s health” or “men’s health,” feminine artwork or pink gowns in the OB/GYN office, etc.



DURING THE APPOINTMENT

- Train ALL STAFF who will interact with the patient (schedulers, valet parking, admin, billing, doctors, nurses, etc.) to use inclusive language
- Call a patient from the waiting room using their last name
- Physical things:
 - Have pronouns on your badge
 - Have a pride flag / safe space sticker / some other indication that you're LGBTQIA2S+ friendly
 - Have diverse posters on the walls of the waiting room / exam room



DURING THE APPOINTMENT

- Pronouns: Give them the option to tell you their pronouns but don't force it
 - "My name is _ and my pronouns are _"
- Ask what name they go by
- Don't assume a patient has a partner, or that their partner is of the opposite sex
- If someone else is with them, ask, "and who do you have with you today?" – don't assume it's their partner



DURING THE APPOINTMENT

- Mirror a patient's language when possible; use the organ terminology they prefer
- Use medical terminology rather than gendered terms, i.e. "estrogen" and "testosterone" rather than "female hormone" or "male hormone"
- Acknowledge that you have limited experience working with this population (cultural humility)
- If you do end up misgendering someone in person, correct it and move on; **don't dwell on it**
- Be clear about WHY you're asking a question and how it will help you provide appropriate medical care

DURING THE APPOINTMENT

- Collect an organ inventory, surgical history, and vaccination history
- Explain that it's important to discuss these organs to understand and reduce their cancer risk...
- ...BUT, stick to what's relevant

| Transfeminine patients: | | Transmasculine patients: | |
|-------------------------|-----|--|-----|
| HPV vaccination? | Y/N | HPV vaccination? | Y/N |
| HPV+? | Y/N | HPV+? | Y/N |
| HIV+? | Y/N | HIV+? | Y/N |
| Neovaginal surgery? | Y/N | Would pt want gyn care during colonoscopies? | Y/N |
| Tissue used? | | | |
| Surgery date? | | | |

| | | | |
|----------------------|------------------|---------------------|------------------|
| Estrogen | How long? | Testosterone | How long? |
| Progesterone | How long? | Oophorectomy | Y/N |
| Breast augmentation | Operative report | Salpingectomy | Y/N |
| Removal of testicles | Y/N | Hysterectomy | Y/N |
| Vaginoplasty | Y/N | Mastectomy | Operative report |
| | | Pregnancies | How many? |
| | | Birth control pills | How long? |












DURING THE APPOINTMENT

- Don't forget to discuss non-genetic factors like tobacco and alcohol use
- Be mindful that GD patients are less likely to follow through with screening; ask how you can support them to continue their care?
- Ask permission before touching the patient, describe the exam in detail, and make sure they are comfortable with every step of what's happening as it happens



AFTER THE APPOINTMENT

- Avoid misgendering, blaming, or stigmatizing patients in notes
 - This erodes trust and contributes to minority stress and adverse health outcomes
- Be thoughtful when making referrals to other providers; are they LGBTQIA2S+-friendly? Do they know how to treat these patients?
- Use standard inclusive pedigree nomenclature

| Gender | Sex | | |
|---------------------------|---|---|--|
| | Male | Female | Unassigned at Birth |
| Man/Boy |  56y |  AFAB 34y |  U/AAB 28y |
| Woman/Girl |  AMAB 56y |  34y |  U/AAB 28y |
| Non-binary/Gender Diverse |  AMAB 56y |  AFAB 34y |  U/AAB 28y |

Bennett, R. L., French, K. S., Resta, R. G., & Austin, J. (2022). Practice resource-focused revision: Standardized pedigree nomenclature update centered on sex and gender inclusivity: A practice resource of the National Society of Genetic Counselors. Journal of Genetic Counseling, 31(6), 1239. <https://doi.org/10.1002/jgc4.1621>

RESOURCES



RESOURCES FOR GD PEOPLE WITH LS

Table 2 Additional resources for TGD individuals with Lynch Syndrome

| Organization | Resource |
|--|---|
| The Gay, Lesbian, Bisexual and Transgender (GLBT) Health Access Project, a collaborative, community-based program funded by the Massachusetts Department of Public Health (MDPH) | <i>Community standards of practice for provision of quality health care services for Gay, Lesbian, Bisexual and Transgendered Clients</i> http://www.glbthealth.org/documents/SOP.pdf |
| National LGBTQIA +Education Center, a program of the Fenway Institute | <i>10 strategies for creating inclusive health care environments</i> https://www.lgbtqiatheducation.org/wp-content/uploads/2021/05/Ten-Strategies-for-Creating-Inclusive-Health-Care-Environments-for-LGBTQIA-People-Brief.pdf |
| The World Professional Association for Transgender Health (WPATH), formerly known as the (Harry Benjamin International Gender Dysphoria Association (HBIGDA)) | <i>Standard of care version 8</i> https://www.wpath.org/soc8 |

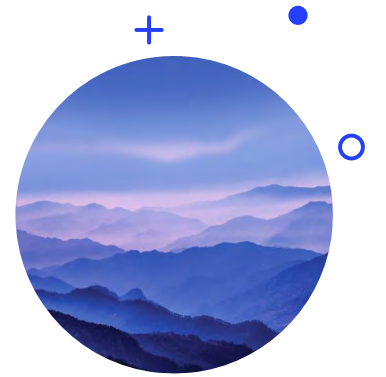
Hodan, R., Rodgers-Fouche, L., Chittenden, A., Dominguez-Valentin, M., Ferriss, J., Gima, L., Hamnvik, O.-P. R., Idos, G. E., Kline, K., Koeller, D. R., Long, J. M., McKenna, D., Muller, C., Thoman, M., Wintner, A., Bedrick, B. S., & on behalf of the Collaborative Group of the Americas on Inherited Gastrointestinal Cancer. (2023). Cancer surveillance for transgender and gender diverse patients with Lynch syndrome: A practice resource of the Collaborative Group of the Americas on Inherited Gastrointestinal Cancer. *Familial Cancer*, 22(4), 437–448. <https://doi.org/10.1007/s10689-023-00341-4>

RESOURCES FOR LGBTQIA2S+ PATIENTS•

- UNMC Gender Care Clinic: <https://www.nebraskamed.com/transgender-care>
- Find an LGBTQIA2S+ welcoming provider:
 - <https://cancer-network.org/providerdatabases/>
 - <https://lgbtqhealthcaredirectory.org/>
- National LGBT Cancer Network: <https://colorectalcancer.cancer-network.org/risk-reduction/>
- The National LGBT Cancer Project: <https://www.lgbtcancer.org/>
- Do Ask, Do Tell: <https://www.lgbtqiahealtheducation.org/publication/dadt/>
- FORCE: <https://www.facingourrisk.org/portal/lgbtq>
- CancerCare: <https://www.cancercare.org/tagged/lgbtq+>
- Patient Handout: How can breast and chest cancer screening help trans and gender diverse people protect themselves? (CONTACT GRACE FOR PDF)
- The Breasties - <https://thebreasties.org/>
- TGD CanScreen Project <http://tgd.dfci.harvard.edu/main>
- Gay & Lesbian Medical Association (GLMA): <https://www.glma.org/resources.php>
- GLAAD Transgender Resources: <https://glaad.org/transgender/resources/>
- Fenway Health: <https://fenwayhealth.org/>
 - National LGBTQIA+ Health Education Center: <https://www.lgbtqiahealtheducation.org/>
- InterACT: <https://interactadvocates.org/resources/>
- Queering Cancer - www.queeringcancer.ca/



RESOURCES FOR PROVIDERS

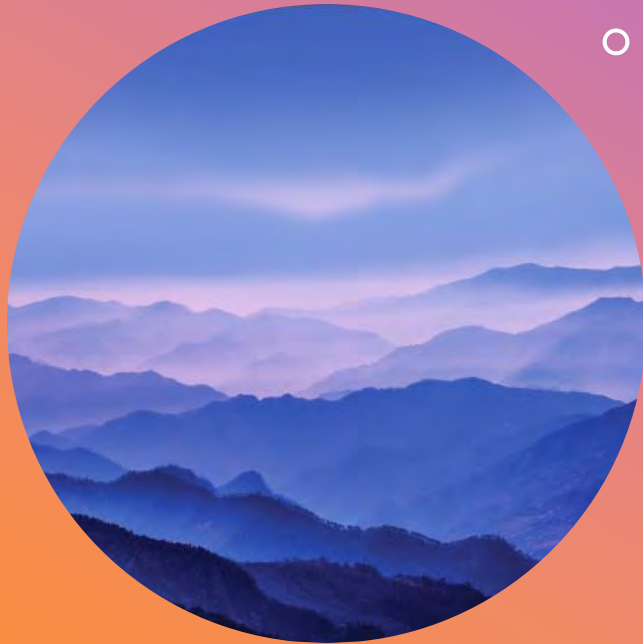


- **(Add yourself)** Find an LGBTQIA2S+ welcoming provider:
 - <https://cancer-network.org/providerdatabases/>
 - <https://lgbtqhealthcaredirectory.org/>
- UNMC Gender Care Clinic: <https://www.nebraskamed.com/transgender-care>
- UNMC SafeSpace Training: <https://unmc.campuslabs.com/engage/organization/gsrc> (UNMC) or on Apollo for Nebraska Medicine colleagues
- Provider Cultural Competency Training: <https://cancer-network.org/cultural-competency-training/>
- **NCCN Guidelines:** Bookmark the WEBPAGE, not the PDF itself (as that is updated often)
 - Genetic/Familial High-Risk Assessment: Breast, Ovarian, Pancreatic, and Prostate (including the GD section): <https://www.nccn.org/guidelines/guidelines-detail?category=2&id=1545>
 - Genetic/Familial High-Risk Assessment: Colorectal, Endometrial, and Gastric: <https://www.nccn.org/guidelines/guidelines-detail?category=2&id=1544>
- Increasing CRC Screening Among LGBTQ+ Communities Brief: <https://nccrt.org/resource/80-in-every-community-lgbt/>
- CGA-IGC LGBTQI+ Toolkit for Providers: <https://www.cgaigc.com/toolkits>
- Variations of Sex Characteristics Cancer Providers Guide: https://www.cgaigc.com/_files/ugd/96b355_37f310d9ef4545a5bd7ce1f2e4e2b7bd.pdf
- LGBTQ+ Cancer Fact Sheet for Providers: <https://www.cancer.org/content/dam/cancer-org/cancer-control/en/booklets-flyers/lgbtq-people-with-cancer-fact-sheet.pdf>
- Collecting SOGI Data: <https://www.whitehouse.gov/wp-content/uploads/2023/01/SOGI-Best-Practices.pdf>
- Guidelines for the Primary and Gender-Affirming Care of Transgender and Gender Nonbinary People: <https://transcare.ucsf.edu/guidelines>
- Fenway Health: <https://fenwayhealth.org/>
- InterACT: <https://interactadvocates.org/resources/>
- The Gay and Lesbian Medical Association (GLMA): <https://www.glma.org/resources.php>

CONCLUSIONS

- The LGBTQIA2S+ population in the US is growing and you are going to see these patients in your practice.
- These patients face health disparities, higher rates of advanced cancer, and lower rates of survival than cis individuals.
- We need more cancer guidelines for this population to provide better care!
- Further studies are needed to determine the impact of hormones and other cancer risks in this population.
- There are many resources and ways you can make this patient population feel welcome and safe.





THANK YOU

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