

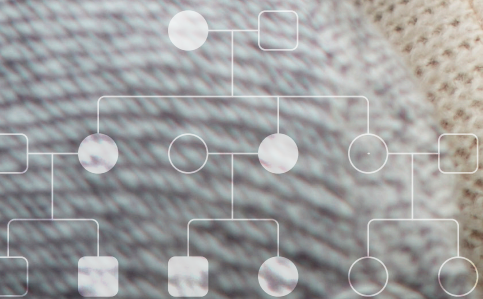


Empower[™]
Hereditary cancer test



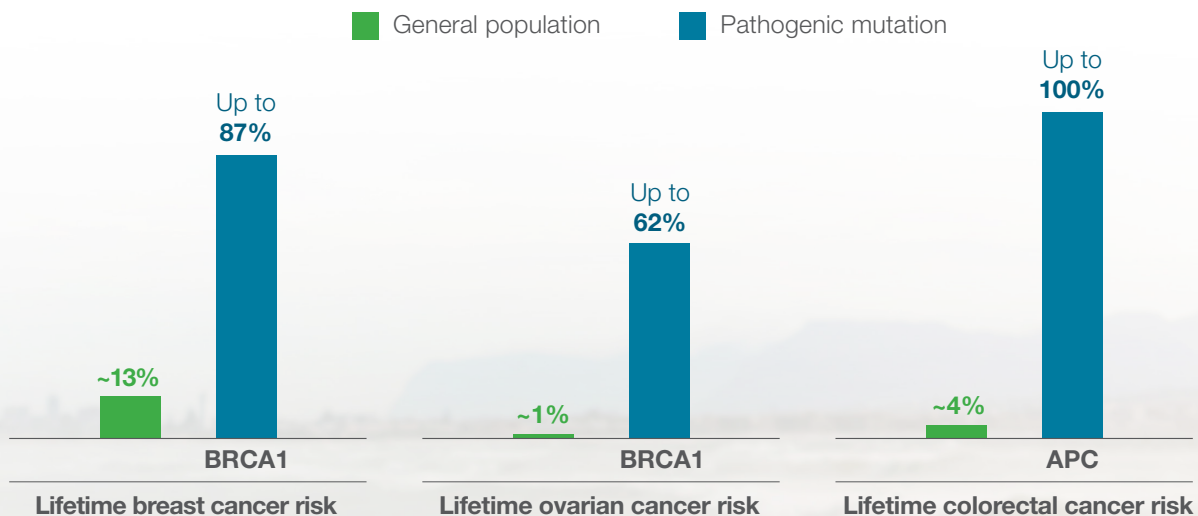
Meaningful insights,
within reach.

High-quality hereditary cancer testing made accessible



Talk about hereditary cancer risk with your patients

Inherited mutations can significantly increase lifetime risk for developing cancer¹



Many high risk patients are not tested



of women at “high risk” for hereditary breast or ovarian cancer never discussed genetic testing with their health care provider²



of individuals at risk for Lynch syndrome were never advised by their health care provider to undergo genetic testing³

A central role for women’s healthcare providers

“Evaluating a patient’s risk of hereditary breast and ovarian cancer should be a routine part of obstetric and gynecologic practice”

— American College of Obstetricians and Gynecologists practice bulletin 182 2017 (reaffirmed 2019)⁴

Introducing Empower™

High-quality hereditary cancer testing made accessible

1

Advanced technology

Next generation sequencing used to analyze up to 53 genes associated with an increased risk for common hereditary cancers

2

Clear answers you can act on

Reports include detailed patient management recommendations based on the latest medical guidelines

3

Commitment to affordability

In-network with a majority of insurance plans, and comprehensive programs to ensure patient access to testing

4

Practice support

Services to simplify testing workflow at every step, including streamlined patient education, ordering and sample collection, billing, counseling and documentation.



From Natera, the experienced leader in women's health genetics

2M+

tests performed

100+

clinicians, PhD's, and scientists

CAP

accredited

CLIA

certified

90

countries worldwide

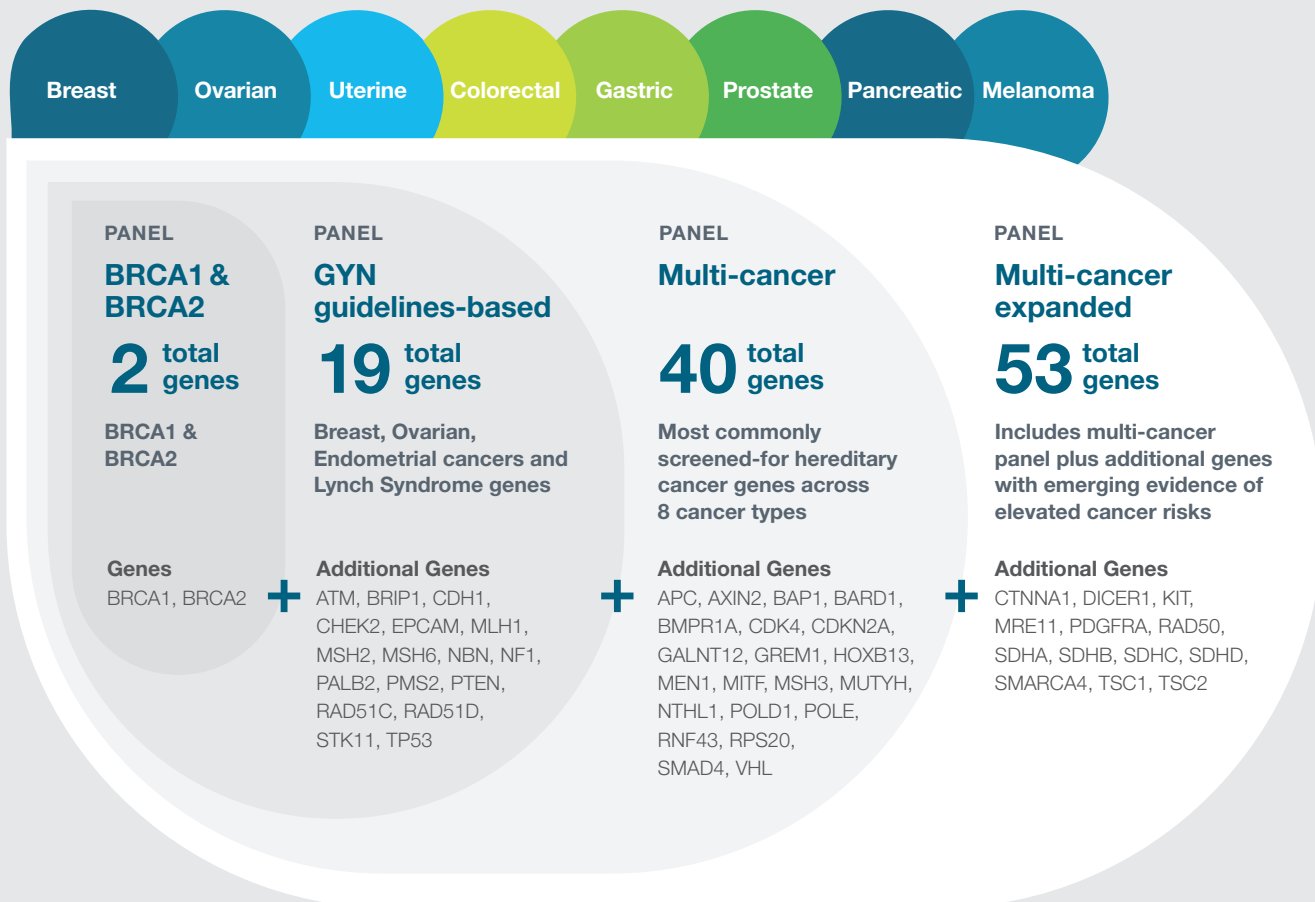
50+

peer-reviewed publications

Hereditary cancer testing simplified

Designed with your practice in mind

Empower panels include genes associated with increased risk of common hereditary cancers, with options to suit your preferred screening strategy.



Vital testing made affordable

Broad, in network coverage

In-network provider with most health plans, including Anthem, Cigna, and UnitedHealthcare.

Check out our growing list at: natera.com/in-network-plans

Price transparency

Personalized estimates help patients understand coverage and cost—so you can focus on care.

Comprehensive patient access solutions

Self-pay pricing and compassionate care options are available for patients without adequate insurance coverage.

Family testing program

Testing for first-degree relatives of patients with a positive result at no additional charge.

Visit natera.com/Empower for more details

Actionable reports to guide patient management

- Screenings that detect cancer at its earliest, most treatable stage
- Risk-reducing medications and surgeries
- Surgical or therapeutic decisions for patients diagnosed with cancer
- Informing family members to help them proactively manage hereditary cancer risk

Cancer Risk Estimates for *BRCA2*

Cancer risk estimates for a positive result are typically based on individuals with a family or personal history of cancer. Your risk may be different if you do not have a personal or family history of cancer.

Female	Cancer type	General Population - Estimated Lifetime Cancer Risk*	Positive Result - Estimated Lifetime Cancer Risk**
♀	Breast	12.80%	Up to 84% risk
	Ovarian	1.3%	Up to 27% risk
	Pancreatic	1.60%	2-7%
	Melanoma	1.80%	Increased

Risk Management and Screening Guidelines for Hereditary Breast and Ovarian Cancer Syndrome *

The following information is a summary of current US guidelines. Please discuss with your healthcare provider as screening recommendations may vary by country and can change often.

Cancer type	Mode of Screening or Risk Reduction	Typical Age to Begin	How often
Female Breast	Breast self-exam	18	Monthly
	Breast exam with clinician	25	Every 6-12 months
	Breast MRI with contrast	25-29 or individualized if family history of breast cancer below age 30	Annually
	Mammogram with consideration of tomosynthesis (3-D Mammogram)	30	Annually
	Risk-reducing medication	Individualized	Discuss with your healthcare provider
	Risk-reducing breast surgery (mastectomy); discuss with your healthcare provider	Individualized; a consideration for those with a personal and/or family history of breast cancer	N/A
Male Breast	Breast self-exam	35	Monthly
	Breast exam with clinician	35	Annually

Report supplements include detailed patient management recommendations based on medical guidelines

Delivered with clinical rigor

Expert variant interpretation

An experienced team of scientists, physicians and genetic counselors rigorously classifies variants according to American College of Medical Genetics (ACMG) guidelines

Sharing knowledge

Baylor Genetics regularly contributes clinically significant variants to ClinVar public database for the benefit of patients and the medical community

DEVELOPED IN PARTNERSHIP WITH

BAYLOR
GENETICS

>4
million
clinical tests
performed

+

>40
years
of
experience

Patients first. Partners always.

Natera supports providers and their patients at every step of the genetic testing process



References

1. Cancer risk estimates for a positive result are typically based on individuals with a family or personal history of cancer. NCCN Clinical Practice Guidelines in Oncology Genetic/Familial High Risk Assessment: Breast, Ovarian and Pancreatic v1.2020
2. Bellcross CA, Peipins LA, McCarty FA, Rodriguez JL, et al. Genet Med. Characteristics associated with genetic counseling referral and BRCA1/2 testing among women in a large integrated health system. 2015 Jan;17(1):43-50.
3. Patel SG, Ahnen DJ, Kinney AY, et al. Am J Gastroenterol. Knowledge and uptake of genetic counseling and colonoscopic screening among individuals at increased risk for lynch syndrome and their endoscopists from the family health promotion project. 2016 Feb;111(2):285-93.
4. Modesitt SC, Lu K, Chen L and Powell CB. Obstet Gynecol. Practice Bulletin No 182: Hereditary Breast and Ovarian Cancer Syndrome. 2017 Sep;130(3):e110-e126. (Reaffirmed 2019)

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The test described has been developed and its performance characteristics determined by the CLIA-certified laboratory performing the test. The test has not been cleared or approved by the US Food and Drug Administration (FDA). Although FDA is exercising enforcement discretion of premarket review and other regulations for laboratory-developed tests in the US, certification of the laboratory is required under CLIA to ensure the quality and validity of the tests. CAP accredited, ISO 13485 certified, and CLIA certified. © 2020 Natera, Inc. All Rights Reserved. EMP_BR_ProviderBrochure_20200817_NAT-8020080

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Hereditary cancer test