



Understanding Test Results

The MyOme® Proactive Health Single-Gene Risk™ test analyzes genes linked to adult onset, actionable conditions, such as cardiac disease and cancer, to support early detection or reduced risk of developing the disease.



Test Result Overview

Positive Result

A pathogenic or likely pathogenic variant was found in one of the genes analyzed, suggesting an increased risk for the associated condition(s).

+ A pathogenic variant was identified in the PALB2 gene.

GENE	RESULT	VARIANT	ZYGOSITY
PALB2	Pathogenic	NM_024675.3:c.2167_2168del	Heterozygous

CLINICAL INFORMATION

This testing found that you have a change in a gene called *PALB2*, which is known to increase the risk of developing certain cancers, primarily breast, ovarian, and pancreatic.

Women with a pathogenic variant in *PALB2* have a 33-58% chance of developing breast cancer (PMID 25099575, 31841383), a 5% chance of developing ovarian cancer and a 1-4% chance of developing pancreatic cancer by age 80. Men have a 2-5% chance of developing pancreatic cancer and a 1% chance of developing male breast cancer by age 80 (PMID 31841383).

GENE INFORMATION

The gene in which the variant was identified.

VARIANT

The specific genetic variant detected.

ZYGOSITY

How many copies of the variant were detected: heterozygous (one) or homozygous (two).

CLINICAL INFORMATION

Brief overview of the disease and risks associated with the genetic variant detected.

Interventions may be necessary to manage the risk, including:



INCREASED SURVEILLANCE

More frequent screening or monitoring for the associated condition.



PREVENTIVE INTERVENTIONS

Lifestyle modifications or medications aimed at reducing risk.



SPECIALIST REFERRAL

Referral to a genetic counselor or specialist for further assessment and personalized risk management.

Negative Result

No known pathogenic or likely pathogenic variants were found in the genes analyzed.

A negative result does not eliminate the risk of the patient developing any of the associated conditions. The test may not cover all genes or genetic variants associated with a particular condition. A patient's family history, lifestyle, and environment should also be considered.*



CONTINUE MONITORING

General population screening is still indicated.

*Increased clinical screening or diagnostic genetic testing may be necessary for patients with a personal or family history of an inherited condition included in the test.



Positive Result Next Steps

- A positive result is medically actionable.
- Family members may be at increased risk for this variant.
- Patients should share results with their family.
- Genetic testing can identify who may benefit from increased screening and care.



PATIENT EXAMPLE

A patient has a positive result with a pathogenic variant in the MYH7 gene associated with hypertrophic cardiomyopathy (HCM). Patients at risk for HCM should be referred to cardiology for screening, typically including echocardiogram and follow-up per cardiology recommendations.



Negative Result Next Steps

- Negative results should be considered alongside clinical findings, family history, and other data.
- A negative result doesn't rule out unique risks for family members.
- Family members may benefit from genetic testing to assess their disease risk.



RESOURCES TO HELP

Genetic counseling can help address any questions you or your patients have about test results and personal or family medical history.



MyOme Gene List

A complete list of the genes in the MyOme® Proactive Health™ screening test is available at www.myome.com/proactive-health.



Make MyOme Proactive Health part of your clinical care.
Contact support@myome.com to get started.