

MyRisk™ Hereditary Cancer Test  
MyRisk Genetic Result

RECEIVING HEALTHCARE PROVIDER: Test HCP MD, Test Medical Center, 6589 BLANCO RD STE 200, SAN ANTONIO, TX 78216

SPECIMEN: Specimen Type: Blood, Draw Date: Dec 03, 2021, Accession Date: Dec 03, 2021, Report Date: Dec 03, 2021

PATIENT: Name: Pt Last Name, Pt First Name, Date of Birth: Dec 03, 1976, Patient ID: Patient id, Gender: Female, Accession #: 07007253-BLD, Requirement #: 90026822

GENETIC RESULT: POSITIVE - CLINICALLY SIGNIFICANT MUTATION IDENTIFIED

CLINICAL HISTORY ANALYSIS: NO ADDITIONAL MANAGEMENT GUIDELINES IDENTIFIED BASED ON THE CLINICAL HISTORY PROVIDED

Other clinical factors may influence individualized management. This analysis may be incomplete if details about cancer diagnoses, ages, family relationships or other factors were omitted or ambiguous.

GENE	MUTATION	INTERPRETATION
BRCA2	c.1163_1164dupTA (p.Pro387Yfs*11) Heterozygous	High Risk This patient has Hereditary Breast and Ovarian Cancer syndrome (HBOC).

DETAILS ABOUT: BRCA2 c.1163\_1164dupTA (p.Pro387Yfs\*11); NM\_000593.3; (aka: I1392NTA)

Functional Significance: Deleterious - Abnormal Protein Production and/or Function

The heterozygous germline BRCA2 mutation c.1163\_1164dupTA is predicted to result in the premature truncation of the BRCA2 protein at amino acid position 397 (p.Pro387Yfs\*11).

Clinical Significance: High Risk

This mutation is associated with increased cancer risk and should be regarded as clinically significant.

ADDITIONAL FINDINGS: NO VARIANT(S) OF UNCERTAIN SIGNIFICANCE (VUS) IDENTIFIED

Details About Non-Clinically Significant Variants: All individuals carry DNA changes (i.e., variants), and most variants do not increase an individual's risk of cancer or other diseases.

Variant Classification: Myriad's myRisk™ Variant Classification Program performs ongoing evaluations of variant classifications.

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ADDITIONAL INFORMATION

Genes Analyzed: Sequencing (seq) and large rearrangement analyses were performed for all coding exons in the following genes, unless otherwise indicated: APC, ATM, AXIN2, BAP1, BARD1, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN2A, CHEK2, CTNNA1, FANCL, FANCD1, FANCD3, FANCF, FANCG, FANCI, FANCD2, FANCD3, FANCD4, FANCD5, FANCD6, FANCD7, FANCD8, FANCD9, FANCD10, FANCD11, FANCD12, FANCD13, FANCD14, FANCD15, FANCD16, FANCD17, FANCD18, FANCD19, FANCD20, FANCD21, FANCD22, FANCD23, FANCD24, FANCD25, FANCD26, FANCD27, FANCD28, FANCD29, FANCD30, FANCD31, FANCD32, FANCD33, FANCD34, FANCD35, FANCD36, FANCD37, FANCD38, FANCD39, FANCD40, FANCD41, FANCD42, FANCD43, FANCD44, FANCD45, FANCD46, FANCD47, FANCD48, FANCD49, FANCD50, FANCD51, FANCD52, FANCD53, FANCD54, FANCD55, FANCD56, FANCD57, FANCD58, FANCD59, FANCD60, FANCD61, FANCD62, FANCD63, FANCD64, FANCD65, FANCD66, FANCD67, FANCD68, FANCD69, FANCD70, FANCD71, FANCD72, FANCD73, FANCD74, FANCD75, FANCD76, FANCD77, FANCD78, FANCD79, FANCD80, 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