Germline Cancer Predisposition Panel

Test Description

The MolQ Germline Cancer Predisposition Panel include genes associated with both common and rare hereditary cancers.

Patient Demographic

Name: Ms. Poonam Goel
Sex: Female
Date of Birth/Age: 60 years
Disease: Carcinoma Breast

Clinician

Clinician Name: Dr Amit Verma
Medical Facility: Max Hospital
Pathologist: Not Provided

Specimen

Booking ID: 012104200141
Site: NA
Sample Type: Blood
Date of Collection: 20-04-2021
Date of Booking: 20-04-2021

CLINICAL SYNOPSIS

Poonam Goel, is a case of stage-III carcinoma breast. She has a family history of cancer with her mother diagnosed with carcinoma breast at the age of 55 years. She has been evaluated for pathogenic variations in the genes listed in Appendix 2.

RESULTS

Variants of uncertain significance related to the given phenotype were detected

Gene (Transcript) #	Location	Variant	Zygosity	Disease (OMIM/Ref)	Inheritance	Classification
RAD51C (+) (ENST00000337432.9)	Exon 3	C451G>A (p.Val151Met)	Heterozygous	Breast-Ovarian Cancer, Familial, susceptibility to, 3	-	Uncertain Significance
FANCI (+) (ENST00000310775.12)	Exon 25	c.2672C>T (p.Ser891Leu)	Heterozygous	Breast Cancer[1-3]	-	Uncertain Significance

$Genetic test results are reported based on the recommendations of American College of Medical Genetics.[4]

No other variant that warrants to be reported was detected. Variations with high minor allele frequencies which are likely to be benign will be given upon request.

CLINICAL CORRELATION AND VARIANT INTERPRETATION

RAD51C **p.Val151Met**
Overall depth: 75X

Variant description: A heterozygous missense variation in exon 3 of the ***RAD51C*** gene (**chr17:g.58696739G>A; Depth: 75x**) that results in the amino acid substitution of Methionine for Valine at codon 151 (**p.Val151Met; ENST00000337432.9**) was detected (Table). The observed variation is documented as variant of uncertain significance in hereditary cancer-predisposing syndrome in the ClinVar database[5] and it lies in the protein phosphatase 2C domain of the RAD51C_HUMAN protein[6]. The **p.Val151Met** variant has not been reported in the 1000 genomes database and has a minor allele frequency of 0.002% and 0.03% in the gnomAD and our internal database, respectively. The in-silico predictions# of the variant are possibly damaging by PolyPhen-2 (HumDiv), damaging by SIFT, LRT and Mutation Taster 2 tools. The reference codon is conserved across mammals.

OMIM phenotype: Susceptibility to familial breast-ovarian cancer-3 (OMIM#613399) is caused by mutations in the ***RAD51C*** gene (OMIM*602774)[7].

Based on the above evidence[$], this ***RAD51C*** variation is classified as a variant of uncertain significance and has to be carefully correlated with the clinical symptoms.

FANCI **p.Ser891Leu**
Overall depth: 27X

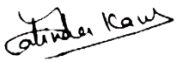
Variant description: A heterozygous missense variation in exon 25 of the ***FANCI*** gene (**chr15:g.89299835C>T; Depth: 27x**) that results in the amino acid substitution of Leucine for Serine at codon 891 (**p.Ser891Leu; ENST00000310775.12**) was detected (Table). The variant lies in the FANCI solenoid-3 domain of the FANCI_HUMAN protein[8]. The **p.Ser891Leu** variant has not been reported in the 1000 genomes, gnomAD and our internal databases. The in-silico predictions# of the variant are benign by PolyPhen-2 (HumDiv), SIFT, LRT and Mutation Taster2 tools. The reference codon is conserved across mammals.

Phenotype: Heterozygous germline loss of function variations in the FANCI gene have been reported in breast cancer patients[1-3].

Based on the above evidence[$], ***this FANCI variation is classified as a variant of uncertain significance and has to be carefully correlated with the clinical symptoms***.

RECOMMENDATIONS

- Validation of the variant(s) by Sanger sequencing is recommended to rule out false positives.
- Sequencing the variant(s) in the other affected and unaffected members of the family is recommended to confirm the significance.
- Genetic counselling is advised for interpretation on the consequences of the variant(s).
- If results obtained do not match the clinical findings, additional testing should be considered as per referring clinician's recommendations.



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APPENDIX 1: TEST METHODOLOGY

Method

Targeted gene sequencing: Selective capture and sequencing of the protein coding regions of the genome/genes is performed. Mutations identified in the exonic regions are generally actionable compared to variations that occur in non-coding regions. Targeted sequencing represents a cost-effective approach to detect variants present in multiple/large genes in an individual.

DNA extracted from sample was used to perform targeted gene capture using a custom capture kit. The libraries were sequenced to mean >80- 100X coverage on Illumina sequencing platform. We follow the GATK best practices framework for identification of variants in the sample using Sentieon (v201808.07)[1]. The sequences obtained are aligned to human reference genome (GRCh38.p13) using Sentieon aligner[1,2] and analyzed using Sentieon for removing duplicates, recalibration and re-alignment of indels[1]. Sentieon haplotype caller has been used to identify variants which are relevant to the clinical indication. Gene annotation of the variants is performed using VEP program[3] against the Ensemble release 99 human gene model[4]. In addition to SNVs and small Indels, copy number variants (CNVs) are detected from targeted sequence data using the ExomeDepth (v1.1.10) method[5]. This algorithm detects rare CNVs based on comparison of the read-depths of the test data with the matched aggregate reference dataset.

Clinically relevant mutations were annotated using published variants in literature and a set of diseases databases - ClinVar, OMIM (updated on 11th May 2020), GWAS, HGMD (v2020.2) and SwissVar[6-10]. Common variants are filtered based on allele frequency in 1000Genome Phase 3, gnomAD (v2.1), EVS, dbSNP (v151), 1000 Japanese Genome and reference lab internal Indian population database[11-15]. Non-synonymous variants effect is calculated using multiple algorithms such as PolyPhen-2, SIFT, MutationTaster2 and LRT. Only non-synonymous and splice site variants found in the germline cancer predisposition panel genes were used for clinical interpretation. Silent variations that do not result in any change in amino acid in the coding region are not reported.

Average Sequencing Depth (x)	Average On-Target Sequencing Depth (x)	Percentage Target Base Pairs Covered		
		0x	≥5x	≥20x
137	112.91	1.17	97.85	92.06

Total data generated (Gb)	1.86
Total reads aligned (%)	99.82
Reads that passed alignment (%)	96.35
Data ≥ Q30 (%)	94.77

$The classification of the variations is done based on American College of Medical Genetics as described below[16]

Variant	A change in a gene. This could be disease causing (pathogenic) or not disease causing (benign).
Pathogenic	A disease causing variation in a gene which can explain the patient's symptoms has been detected. This usually means that a suspected disorder for which testing had been requested has been confirmed.
Likely Pathogenic	A variant which is very likely to contribute to the development of disease however, the scientific evidence is currently insufficient to prove this conclusively. Additional evidence is expected to confirm this assertion of pathogenicity.
Variant of Uncertain Significance	A variant has been detected, but it is difficult to classify it as either pathogenic (disease causing) or benign (non-disease causing) based on current available scientific evidence. Further testing of the patient or family members as recommended by your clinician may be needed. It is probable that their significance can be assessed only with time, subject to availability of scientific evidence.

#The transcript used for clinical reporting generally represents the canonical transcript, which is usually the longest coding transcript with strong/multiple supporting evidence. However, clinically relevant variants annotated in alternate complete coding transcripts could also be reported.
Variants annotated on incomplete and nonsense mediated decay transcripts will not be reported.
#The *in-silico* predictions are based on Variant Effect Predictor, Ensembl release 99 (SIFT version - 5.2.2; PolyPhen - 2.2.2); bdNSFPv4.0 (LRT version – December 5, 2019) and Mutation Taster2 (MT2). MutationTaster2 predictions are based on NCBI/Ensembl 66 build (GRCh38 genomic coordinates are converted to hg19 using UCSC LiftOver and mapped to MT2).
For any further technical queries please contact contact@molq.in.

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DISCLAIMER

- Interpretation of variants in this report is performed to the best knowledge of the laboratory based on the information available at the time of reporting. The classification of variants can change over time and MolQ cannot be held responsible for this. Please feel free to contact MolQ Laboratory (contact@molq.in) in the future to determine if there have been any changes in the classification of any variations. Re-analysis of variants in previously issued reports in light of new evidence is not routinely performed, but may be available upon request.
- The sensitivity of this assay to detect large deletions/duplications of more than 10 bp or copy number variations (CNV) is 70-75%. The CNVs detected have to be confirmed by alternate method.
- Due to inherent technology limitations of the assay, not all bases of the exome can be covered by this test. Accordingly, variants in regions of insufficient coverage may not be identified and/or interpreted. Therefore, it is possible that pathogenic variants are present in one or more of the genes analyzed, but have not been detected. The variants not detected by the assay that was performed may impact the phenotype.
- It is also possible that a pathogenic variant is present in a gene that was not selected for analysis and/or interpretation in cases where insufficient phenotypic information is available.
- Genes with pseudogenes, paralog genes and genes with low complexity may have decreased sensitivity and specificity of variant detection and interpretation due to inability of the data and analysis tools to unambiguously determine the origin of the sequence data in such regions.
- The mutations have not been validated/confirmed by Sanger sequencing.
- Incidental or secondary findings (if any) that meet the ACMG guidelines[17] can be given upon request.
- The report shall be generated within turnaround time (TAT), however, such TAT may vary depending upon the complexity of test(s) requested. MolQ Laboratory under no circumstances will be liable for any delay beyond afore mentioned TAT.
- It is hereby clarified that the report(s) generated from the test(s) do not provide any diagnosis or opinion or recommends any cure in any manner. MolQ Laboratory hereby recommends the patient and/or the guardians of the patients, as the case may be, to take assistance of the clinician or a certified physician or doctor, to interpret the report(s) thus generated. MolQ Laboratory hereby disclaims all liability arising in connection with the report(s).
- In a very few cases genetic test may not show the correct results, e.g. because of the quality of the material provided to MolQ Laboratory. In case where any test provided by MolQ Laboratory fails for unforeseeable or unknown reasons that cannot be influenced by MolQ Laboratory in advance, MolQ Laboratory shall not be responsible for the incomplete, potentially misleading or even wrong result of any testing if such could not be recognized by MolQ Laboratory in advance.
- This is a laboratory developed test and the development and the performance characteristics of this test was determined by reference laboratory.

LIMITATIONS

- Genetic testing is an important part of the diagnostic process. However, genetic tests may not always give a definitive answer. In some cases, testing may not identify a genetic variant even though one exists. This may be due to limitations in current medical knowledge or testing technology. Accurate interpretation of test results may require knowing the true biological relationships in a family. Failing to accurately state the biological relationships in {my/my child's} family may result in incorrect interpretation of results, incorrect diagnoses, and/or inconclusive test results.
- Test results are interpreted in the context of clinical findings, family history and other laboratory data. Only variations in genes potentially related to the proband's medical condition are reported. Rare polymorphisms may lead to false negative or positive results. Misinterpretation of results may occur if the information provided is inaccurate or incomplete.
- Specific events like copy number variations, translocations, repeat expansions and chromosomal rearrangements may not be reliably detected with targeted clinical exome sequencing. Variants in untranslated region, promoters and intronic variants are not assessed using this method.
- Genetic testing is highly accurate. Rarely, inaccurate results may occur for various reasons. These reasons include, but are not limited to: mislabelled samples, inaccurate reporting of clinical/medical information, rare technical errors or unusual circumstances such as bone marrow transplantation, blood transfusion; or the presence of change(s) in such a small percentage of cells that may not be detectable by the test (mosaicism).
- The variant population allele frequencies and in silico predictions for GRCh38 version of the Human genome is obtained after lifting over the coordinates from hg19 genome build. The existing population allele frequencies (1000Genome, gnomAD-Exome) are currently available for hg19 genome version only. This might result in some discrepancies in variant

annotation due to the complex changes in some regions of the genome.

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APPENDIX 2: GENE LIST WITH COVERAGE

Gene	Percentage of coding region covered	Gene	Percentage of coding region covered	Gene	Percentage of coding region covered
AIP	100.00	*ALK*	100.00	*APC*	100.00
AR	100.00	*ATM*	100.00	*BAP1*	100.00
BARD1	100.00	*BLM*	100.00	*BMPR1A*	100.00
BRCA1	100.00	*BRCA2*	100.00	*BRIP1*	100.00
BUB1B	100.00	*CD82*	100.00	*CDC73*	100.00
CDH1	100.00	*CDK4*	100.00	*CDKN1C*	100.00
CDKN2A	100.00	*CEBPA*	100.00	*CEP57*	100.00
CHEK2	100.00	*CYLD*	100.00	*DDB2*	100.00
DICER1	100.00	*DIS3L2*	100.00	*EGFR*	100.00
ELAC2	100.00	*ENG*	100.00	*EPCAM*	100.00
ERCC2	100.00	*ERCC3*	100.00	*ERCC4*	100.00
ERCC5	100.00	*EXT1*	100.00	*EXT2*	100.00
EZH2	100.00	*FANCA*	100.00	*FANCB*	100.00
FANCC	100.00	*FANCD2*	100.00	*FANCE*	100.00
FANCF	100.00	*FANCG*	100.00	*FANCI*	100.00
FANCL	100.00	*FANCM*	100.00	*FH*	100.00
FLCN	100.00	*GATA2*	100.00	*GPC3*	100.00
HRAS	98.71	*KIT*	100.00	*MAG*	100.00
MAX	100.00	*MEN1*	100.00	*MET*	100.00
MLH1	100.00	*MLH3*	100.00	*MRE11A*	100.00
MSH2	100.00	*MSH3*	100.00	*MSH6*	100.00
MSR1	100.00	*MUTYH*	100.00	*MXI1*	100.00
NBN	100.00	*NF1*	100.00	*NF2*	100.00
NSD1	100.00	*PALB2*	100.00	*PHOX2B*	100.00
PMS1	100.00	*PMS2*	100.00	*PRF1*	100.00
PRKAR1A	100.00	*PTCH1*	100.00	*PTEN*	100.00
RAD50	100.00	*RAD51C*	100.00	*RAD51D*	100.00
RAD54L	100.00	*RB1*	100.00	*RECQL4*	100.00
RET	100.00	*RHBDF2*	100.00	*RNASEL*	100.00
RUNX1	100.00	*SBDS*	100.00	*SDHAF2*	100.00
SDHB	100.00	*SDHC*	100.00	*SDHD*	100.00
SLX4	100.00	*SMAD4*	100.00	*SMARCB1*	100.00
STK11	100.00	*SUFU*	100.00	*TGFBR2*	100.00
TMEM127	100.00	*TP53*	100.00	*TSC1*	100.00
TSC2	100.00	*VHL*	100.00	*WRN*	100.00
WT1	100.00	*XPA*	100.00	*XPC*	100.00