



## Sample Information

**Patient Name:** 張傳穎  
**Gender:** Female  
**ID No.:** A225824821  
**History No.:** 46794223  
**Age:** 35

**Ordering Doctor:** DOC2589K 曾令民  
**Ordering REQ.:** 0BFWFBN  
**Signing in Date:** 2021/05/12

**Path No.:** S110-98765  
**MP No.:** BR21019  
**Assay:** Oncomine BRCA1/2 Assay  
**Sample Type:** Blood  
**Date of blood drawing:** 2021/05/06  
**Note:**

## Sample Cancer Type: Breast Cancer

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Variant Details

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### Report Highlights

0 Therapies Available  
0 Clinical Trials

## Relevant Biomarkers

No clinically significant biomarkers found in this sample.

## Variant Details

### DNA Sequence Variants

Gene	Amino Acid Change	Coding	Locus	Allele Frequency	Transcript	Variant Effect	ClinVar <sup>1</sup>	Coverage
BRCA2	p.(?)	c.-26G>A	chr13:32890572	50.02%	NM_000059.3	unknown	Benign	3998
BRCA2	p.(N372H)	c.1114A>C	chr13:32906729	50.98%	NM_000059.3	missense	Benign	3998
BRCA2	p.(=)	c.3396A>G	chr13:32911888	51.71%	NM_000059.3	synonymous	Benign	3976
BRCA2	p.(=)	c.4563A>G	chr13:32913055	99.92%	NM_000059.3	synonymous	Benign	3741
BRCA2	p.(=)	c.6513G>C	chr13:32915005	99.70%	NM_000059.3	synonymous	Benign	3960
BRCA2	p.(=)	c.7242A>G	chr13:32929232	52.25%	NM_000059.3	synonymous	Benign	3996
BRCA2	p.(V2466A)	c.7397T>C	chr13:32929387	99.72%	NM_000059.3	missense	Conflicting interpretations of pathogenicity	3998
BRCA1	p.(S1634G)	c.4900A>G	chr17:41223094	52.38%	NM_007300.3	missense	Benign	3988

<sup>1</sup> Based on Clinvar version 20180225

## Variant Details (continued)

### DNA Sequence Variants (continued)

Gene	Amino Acid Change	Coding	Locus	Allele Frequency	Transcript	Variant Effect	ClinVar <sup>1</sup>	Coverage
BRCA1	p.(=)	c.4308T>C	chr17:41234470	49.79%	NM_007300.3	synonymous	Benign	3999
BRCA1	p.(K1183R)	c.3548A>G	chr17:41244000	49.57%	NM_007300.3	missense	Benign	3996
BRCA1	p.(E1038G)	c.3113A>G	chr17:41244435	50.16%	NM_007300.3	missense	Benign	3999
BRCA1	p.(P871L)	c.2612C>T	chr17:41244936	49.42%	NM_007300.3	missense	Benign	3990
BRCA1	p.(=)	c.2311T>C	chr17:41245237	49.75%	NM_007300.3	synonymous	Benign	3996
BRCA1	p.(=)	c.2082C>T	chr17:41245466	48.47%	NM_007300.3	synonymous	Benign	3998

<sup>1</sup> Based on Clinvar version 20180225

## Signatures

Testing Personnel:

Laboratory Supervisor:

Pathologist: