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**Date**: 27 Sep 2023 1 of 2

### **Sample Information**

Patient Name: 陳美齡 Gender: Female ID No.: F225006515 History No.: 46375831

**Age:** 42

Ordering Doctor: DOC1242E 劉峻宇

Ordering REQ.: 0CRHLLZ Signing in Date: 2023/09/27

**Path No.:** M112-00257 **MP No.:** BR23071

Assay: Oncomine BRCA1/2 Assay

Sample Type: Blood

Date of blood drawing: 2023/09/21

Reporting Doctor: DOC5424G 彭昱璟 (Phone: 8#5424)

Note:

## Sample Cancer Type: Breast Cancer

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#### **Relevant Breast Cancer Variants**

Gene	Finding
BRCA1	None detected
BRCA2	None detected

#### **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.(N372H)	c.1114A>C	chr13:32906729	51.75%	NM_000059.3	missense	Benign	3998	
BRCA2	p.(L1521=)	c.4563A>G	chr13:32913055	100.00%	NM_000059.3	synonymous	Benign	2244	

<sup>1</sup> Based on Clinvar version 20200329

# **Variant Details (continued)**

## **DNA Sequence Variants (continued)**

Gene	Amino Acid Change	Coding	Locus	Allele Frequency	Transcript	Variant Effect	ClinVar <sup>1</sup>	Coverage
BRCA2	p.(V2171=)	c.6513G>C	chr13:32915005	99.97%	NM_000059.3	synonymous	Benign	4000
BRCA2	p.(V2466A)	c.7397T>C	chr13:32929387	99.91%	NM_000059.3	missense	Benign	3303
BRCA1	p.(S1613G)	c.4837A>G	chr17:41223094	99.90%	NM_007294.4	missense	Benign	3998
BRCA1	p.(S1436=)	c.4308T>C	chr17:41234470	99.90%	NM_007294.4	synonymous	Benign	3887
BRCA1	p.(K1183R)	c.3548A>G	chr17:41244000	99.70%	NM_007294.4	missense	Benign	4000
BRCA1	p.(E1038G)	c.3113A>G	chr17:41244435	99.55%	NM_007294.4	missense	Benign	4000
BRCA1	p.(P871L)	c.2612C>T	chr17:41244936	99.82%	NM_007294.4	missense	Benign	3996
BRCA1	p.(L771=)	c.2311T>C	chr17:41245237	99.97%	NM_007294.4	synonymous	Benign	3528
BRCA1	p.(S694=)	c.2082C>T	chr17:41245466	99.77%	NM_007294.4	synonymous	Benign	3995

<sup>1</sup> Based on Clinvar version 20200329