

Department of Pathology and Laboratory Medicine No.201, Sec. 2, Shipai Rd., Beitou District, Taipei City, Taiwan 11217, R.O.C. Tel: 02-2875-7449

Date: 03 Jun 2021

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## **Sample Information**

Patient Name: 鄭肇心 Gender: Female ID No.: F227854724 History No.: 22963924

**Age:** 45

Ordering Doctor: DOC2589K 曾令民

Ordering REQ.: 0BGNYJJ Signing in Date: 2021/06/02

**Path No.**: S110-98911 **MP No.**: BR21026

Assay: Oncomine BRCA1/2 Assay

Sample Type: Blood

Date of blood drawing: 2021/05/26

Note:

### Sample Cancer Type: Breast Cancer

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Report Highlights
0 Therapies Available
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#### **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.(?)	c26G>A	chr13:32890572	48.70%	NM_000059.3	unknown	Benign	3992				
BRCA2	p.(=)	c.3396A>G	chr13:32911888	52.20%	NM_000059.3	synonymous	Benign	3979				
BRCA2	p.(=)	c.3807T>C	chr13:32912299	51.01%	NM_000059.3	synonymous	Benign	3991				
BRCA2	p.(=)	c.4563A>G	chr13:32913055	100.00%	NM_000059.3	synonymous	Benign	2880				
BRCA2	p.(=)	c.6513G>C	chr13:32915005	99.92%	NM_000059.3	synonymous	Benign	3968				
BRCA2	p.(=)	c.7242A>G	chr13:32929232	50.83%	NM_000059.3	synonymous	Benign	3996				
BRCA2	p.(V2466A)	c.7397T>C	chr13:32929387	99.72%	NM_000059.3	missense	Conflicting interpretations of pathogenicity	3999				
BRCA1	p.(S1634G)	c.4900A>G	chr17:41223094	49.71%	NM_007300.3	missense	Benign	3989				

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

# **Variant Details (continued)**

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

Gene	Amino Acid Change	Codina	Locus	Allele Frequency	Transcript	Variant Effect	ClinVar1	Coverage
Gene	Allillo Acid Olidlige	Odding	Locus	rrequericy	Transcript	Variant Enect	Ollif Val	Ouverage
BRCA1	p.(=)	c.4308T>C	chr17:41234470	48.25%	NM_007300.3	synonymous	Benign	3998
BRCA1	p.(K1183R)	c.3548A>G	chr17:41244000	53.32%	NM_007300.3	missense	Benign	3995
BRCA1	p.(E1038G)	c.3113A>G	chr17:41244435	50.39%	NM_007300.3	missense	Benign	3997
BRCA1	p.(P871L)	c.2612C>T	chr17:41244936	48.16%	NM_007300.3	missense	Benign	3991
BRCA1	p.(=)	c.2311T>C	chr17:41245237	49.37%	NM_007300.3	synonymous	Benign	3992
BRCA1	p.(=)	c.2082C>T	chr17:41245466	50.29%	NM_007300.3	synonymous	Benign	3997

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

Date: 03 Jun 2021

# **Signatures**

**Testing Personnel:** 

**Laboratory Supervisor:** 

Pathologist: