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# **Sample Information**

Patient Name: 何美麗 Gender: Female ID No.: C220183238 History No.: 38607066

**Age:** 64

Ordering Doctor: DOC3559K 屠乃方 Ordering REQ.: 0BDUVUP Signing in Date: 2021/03/24

**Path No.**: S110-98470 **MP No.**: BR21012

Assay: Oncomine BRCA1/2 Assay

Sample Type: FFPE Block No.: \$110-65649\$ Percentage of tumor cells: 95%

Note:

### Sample Cancer Type: Ovarian Cancer

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# **Relevant Ovarian Cancer Variants**

Gene	Finding
BRCA1	Not detected
BRCA2	Not 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.(=)	c.3396A>G	chr13:32911888	11.36%	NM_000059.3	synonymous	Benign	1998	
BRCA2	p.(=)	c.4563A>G	chr13:32913055	99.94%	NM_000059.3	synonymous	Benign	1681	
BRCA2	p.(=)	c.6513G>C	chr13:32915005	99.95%	NM_000059.3	synonymous	Benign	1988	

<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
BRCA2	p.(=)	c.7242A>G	chr13:32929232	11.10%	NM_000059.3	synonymous	Benign	2000
BRCA2	p.(V2466A)	c.7397T>C	chr13:32929387	99.80%	NM_000059.3	missense	Conflicting interpretations of pathogenicity	1998
BRCA1	p.(S1634G)	c.4900A>G	chr17:41223094	90.48%	NM_007300.3	missense	Benign	1996
BRCA1	p.(=)	c.114G>A	chr17:41267763	10.41%	NM_007300.3	synonymous	Benign	1999

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

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# Signatures Testing Personnel: Laboratory Supervisor:

Pathologist: