



## Sample Information

**Patient Name:** 羅如芳**Gender:** Female**ID No.:** K220843683**History No.:** 14548675**Age:** 55**Ordering Doctor:** DOC3697E 陳怡仁**Ordering REQ.:** 0BAHDX**Signing in Date:** 2020/12/30**Path No.:** S109-96878**MP No.:** BR20007**Assay:** Oncomine BRCA1/2 Assay**Sample Type:** FFPE**Block No.:** S109-69041T**Percentage of tumor cells:** 60%**Note:**

## Sample Cancer Type: Ovarian Cancer

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### Report Highlights

0 Therapies Available

0 Clinical Trials

## 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 Fraction	Transcript	Variant Effect	ClinVar <sup>1</sup>	Coverage
BRCA2	p.(?)	c.-26G>A	chr13:32890572	0.998	NM_000059.3	unknown	Benign	1996
BRCA2	p.(=)	c.3396A>G	chr13:32911888	0.995	NM_000059.3	synonymous	Benign	1996
BRCA2	p.(=)	c.4563A>G	chr13:32913055	0.999	NM_000059.3	synonymous	Benign	1995

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



## Variant Details (continued)

### DNA Sequence Variants (continued)

Gene	Amino Acid Change	Coding	Locus	Allele Fraction	Transcript	Variant Effect	ClinVar <sup>1</sup>	Coverage
BRCA2	p.(=)	c.6513G>C	chr13:32915005	0.997	NM_000059.3	synonymous	Benign	1990
BRCA2	p.(=)	c.7242A>G	chr13:32929232	0.997	NM_000059.3	synonymous	Benign	2000
BRCA2	p.(V2466A)	c.7397T>C	chr13:32929387	0.998	NM_000059.3	missense	Conflicting interpretations of pathogenicity	1996

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



## Signatures

Testing Personnel:

Laboratory Supervisor:

Pathologist: