



Sample Information

Patient Name: 鄭美綺
Gender: Female
ID No.: A225516879
History No.: 46832466
Age: 32

Ordering Doctor: DOC3697E 陳怡仁
Ordering REQ.: 0BEXFEK
Signing in Date: 2021/04/21

Path No.: S110-98620
MP No.: BR21016
Assay: Oncomine BRCA1/2 Assay
Sample Type: FFPE
Block No.: S110-65636S
Percentage of tumor cells: 70%
Note:

Sample Cancer Type: Ovarian Cancer

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Variant Details

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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 Frequency	Transcript	Variant Effect	ClinVar ¹	Coverage
BRCA2	p.(?)	c.-26G>A	chr13:32890572	99.70%	NM_000059.3	unknown	Benign	1998
BRCA2	p.(E350K)	c.1048G>A	chr13:32906663	2.47%	NM_000059.3	missense		1987
BRCA2	p.(T582P)	c.1744A>C	chr13:32907359	61.99%	NM_000059.3	missense	Benign	1997

¹ Based on Clinvar version 20180225

Variant Details (continued)

DNA Sequence Variants (continued)

Gene	Amino Acid Change	Coding	Locus	Allele Frequency	Transcript	Variant Effect	ClinVar ¹	Coverage
BRCA2	p.(=)	c.3396A>G	chr13:32911888	99.60%	NM_000059.3	synonymous	Benign	1994
BRCA2	p.(=)	c.4563A>G	chr13:32913055	100.00%	NM_000059.3	synonymous	Benign	1340
BRCA2	p.(=)	c.6513G>C	chr13:32915005	99.90%	NM_000059.3	synonymous	Benign	1992
BRCA2	p.(=)	c.7242A>G	chr13:32929232	99.60%	NM_000059.3	synonymous	Benign	1999
BRCA2	p.(V2466A)	c.7397T>C	chr13:32929387	99.95%	NM_000059.3	missense	Conflicting interpretations of pathogenicity	2000
BRCA1	p.(=)	c.114G>A	chr17:41267763	47.10%	NM_007300.3	synonymous	Benign	1998

¹ Based on Clinvar version 20180225

Signatures

Testing Personnel:

Laboratory Supervisor:

Pathologist: