



Sample Information

Patient Name: 李沛淳
Gender: Female
ID No.: L221392836
History No.: 46678711
Age: 52

Ordering Doctor: DOC3697E 陳怡仁
Ordering REQ.: 0BDRXYN
Signing in Date: 2021/03/24

Path No.: S110-98469
MP No.: BR21011
Assay: Oncomine BRCA1/2 Assay
Sample Type: Blood
Date of blood drawing: 2021/03/16
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	54.24%	NM_000059.3	unknown	Benign	2301
BRCA2	p.(=)	c.3396A>G	chr13:32911888	50.22%	NM_000059.3	synonymous	Benign	3172
BRCA2	p.(=)	c.3807T>C	chr13:32912299	50.70%	NM_000059.3	synonymous	Benign	2142
BRCA2	p.(=)	c.4563A>G	chr13:32913055	99.71%	NM_000059.3	synonymous	Benign	1048

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.6513G>C	chr13:32915005	99.95%	NM_000059.3	synonymous	Benign	2164
BRCA2	p.(=)	c.7242A>G	chr13:32929232	48.43%	NM_000059.3	synonymous	Benign	1404
BRCA2	p.(V2466A)	c.7397T>C	chr13:32929387	99.86%	NM_000059.3	missense	Conflicting interpretations of pathogenicity	1425
BRCA1	p.(Y856H)	c.2566T>C	chr17:41244982	50.52%	NM_007300.3	missense	Benign	1550

¹ Based on Clinvar version 20180225

Signatures

Testing Personnel:

Laboratory Supervisor:

Pathologist: