



Sample Information

Patient Name: 林怡君
Gender: Female
ID No.: R222489413
History No.: 37601971
Age: 43

Ordering Doctor: DOC2095G 蔡宜芳
Ordering REQ.: 0CKRLCJ
Signing in Date: 2023/05/18

Path No.: M112-00104
MP No.: BR23029
Assay: Oncomine BRCA1/2 Assay
Sample Type: Blood
Date of blood drawing: 2023/05/08

Reporting Doctor: DOC5424G 彭昱璟 (Phone: 8#5424)

Note:

Sample Cancer Type: Breast Cancer

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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 ¹	Coverage
BRCA2	p.(V1269=)	c.3807T>C	chr13:32912299	99.52%	NM_000059.3	synonymous	Benign	3994
BRCA2	p.(L1521=)	c.4563A>G	chr13:32913055	99.91%	NM_000059.3	synonymous	Benign	3452
BRCA2	p.(V2171=)	c.6513G>C	chr13:32915005	100.00%	NM_000059.3	synonymous	Benign	3997
BRCA2	p.(V2466A)	c.7397T>C	chr13:32929387	99.90%	NM_000059.3	missense	Benign	3999
BRCA1	p.(S1613G)	c.4837A>G	chr17:41223094	49.77%	NM_007294.4	missense	Benign	3982
BRCA1	p.(S1436=)	c.4308T>C	chr17:41234470	49.09%	NM_007294.4	synonymous	Benign	3999
BRCA1	p.(K1183R)	c.3548A>G	chr17:41244000	49.34%	NM_007294.4	missense	Benign	3999

¹ Based on Clinvar version 20200329

Variant Details (continued)

DNA Sequence Variants (continued)

Gene	Amino Acid Change	Coding	Locus	Allele Frequency	Transcript	Variant Effect	ClinVar ¹	Coverage
BRCA1	p.(E1038G)	c.3113A>G	chr17:41244435	52.70%	NM_007294.4	missense	Benign	3998
BRCA1	p.(P871L)	c.2612C>T	chr17:41244936	50.58%	NM_007294.4	missense	Benign	3994
BRCA1	p.(L771=)	c.2311T>C	chr17:41245237	49.70%	NM_007294.4	synonymous	Benign	3998
BRCA1	p.(S694=)	c.2082C>T	chr17:41245466	50.24%	NM_007294.4	synonymous	Benign	3999

¹ Based on Clinvar version 20200329