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Date: 22 Sep 2022 1 of 3

Sample Information

Patient Name: 鄒秀絨 Gender: Female ID No.: F203310092 History No.: 45421656

Age: 67

Ordering Doctor: DOC3707G 洪煥程

Ordering REQ.: 0BZYMBP Signing in Date: 2022/09/22

Path No.: S111-97901 **MP No.:** BR22054

Assay: Oncomine BRCA1/2 Assay

Sample Type: Blood

Date of blood drawing: 2022/09/13

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

Note:

Sample Cancer Type: Other Solid Tumor

Table of ContentsPageVariant Details1

Report Highlights 0 Therapies Available 0 Clinical Trials

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.(N372H)	c.1114A>C	chr13:32906729	50.67%	NM_000059.3	missense	Benign	3274
BRCA2	p.(K454=)	c.1362A>G	chr13:32906977	53.04%	NM_000059.3	synonymous	Benign	3997
BRCA2	p.(V1269=)	c.3807T>C	chr13:32912299	47.35%	NM_000059.3	synonymous	Benign	3996
BRCA2	p.(L1521=)	c.4563A>G	chr13:32913055	99.97%	NM_000059.3	synonymous	Benign	3989
BRCA2	p.(V2171=)	c.6513G>C	chr13:32915005	99.91%	NM_000059.3	synonymous	Benign	3490
BRCA2	p.(V2466A)	c.7397T>C	chr13:32929387	99.93%	NM_000059.3	missense	Benign	3999
BRCA1	p.(S1613G)	c.4837A>G	chr17:41223094	48.93%	NM_007294.4	missense	Benign	3983

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.(S1436=)	c.4308T>C	chr17:41234470	50.69%	NM_007294.4	synonymous	Benign	3490
BRCA1	p.(K1183R)	c.3548A>G	chr17:41244000	48.80%	NM_007294.4	missense	Benign	4000
BRCA1	p.(E1038G)	c.3113A>G	chr17:41244435	51.63%	NM_007294.4	missense	Benign	4000
BRCA1	p.(P871L)	c.2612C>T	chr17:41244936	49.31%	NM_007294.4	missense	Benign	3995
BRCA1	p.(L771=)	c.2311T>C	chr17:41245237	48.25%	NM_007294.4	synonymous	Benign	3996
BRCA1	p.(S694=)	c.2082C>T	chr17:41245466	49.79%	NM_007294.4	synonymous	Benign	3995

¹ Based on Clinvar version 20200329

Date: 22 Sep 2022 3 of 3

Signatures Testing Personnel:

Pathologist:

Laboratory Supervisor: