

Department of Pathology and Laboratory Medicine No.201, Sec. 2, Shipai Rd., Beitou District, Taipei City, Taiwan 11217, R.O.C.

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Date: 30 Nov 2021 1 of 3

Sample Information

Patient Name: 陳心潔 Gender: Female ID No.: A225776857 History No.: 47885416

Age: 35

Ordering Doctor: DOC2366J 施沐姍 Ordering REQ.: 0BNUWWP Signing in Date: 2021/11/29

Path No.: S110-94637 **MP No.:** BR21051

Assay: Oncomine BRCA1/2 Assay

Sample Type: Blood

Date of blood drawing: 2021/11/17

Reporting Doctor: DOC5452C 周德盈 (Phone: 8#5452)

Note:

Sample Cancer Type: Breast Cancer

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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 | ClinVar1 | Coverage |
|-------|-------------------|-----------|----------------|---------------------|-------------|----------------|----------|----------|
| BRCA2 | p.(?) | c26G>A | chr13:32890572 | 51.64% | NM_000059.3 | unknown | Benign | 3997 |
| BRCA2 | p.(N372H) | c.1114A>C | chr13:32906729 | 50.66% | NM_000059.3 | missense | Benign | 3995 |
| BRCA2 | p.(K1132=) | c.3396A>G | chr13:32911888 | 51.28% | NM_000059.3 | synonymous | Benign | 3998 |
| BRCA2 | p.(L1521=) | c.4563A>G | chr13:32913055 | 99.89% | NM_000059.3 | synonymous | Benign | 3570 |
| BRCA2 | p.(V2171=) | c.6513G>C | chr13:32915005 | 99.87% | NM_000059.3 | synonymous | Benign | 3998 |
| BRCA2 | p.(S2414=) | c.7242A>G | chr13:32929232 | 48.29% | NM_000059.3 | synonymous | Benign | 3999 |
| BRCA2 | p.(V2466A) | c.7397T>C | chr13:32929387 | 99.77% | NM_000059.3 | 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.(S1613G) | c.4837A>G | chr17:41223094 | 99.82% | NM_007294.4 | missense | Benign | 3997 |
| BRCA1 | p.(S1436=) | c.4308T>C | chr17:41234470 | 99.78% | NM_007294.4 | synonymous | Benign | 4000 |
| BRCA1 | p.(K1183R) | c.3548A>G | chr17:41244000 | 99.20% | NM_007294.4 | missense | Benign | 4000 |
| BRCA1 | p.(E1038G) | c.3113A>G | chr17:41244435 | 99.60% | NM_007294.4 | missense | Benign | 3999 |
| BRCA1 | p.(P871L) | c.2612C>T | chr17:41244936 | 99.85% | NM_007294.4 | missense | Benign | 3997 |
| BRCA1 | p.(L771=) | c.2311T>C | chr17:41245237 | 99.90% | NM_007294.4 | synonymous | Benign | 3996 |
| BRCA1 | p.(S694=) | c.2082C>T | chr17:41245466 | 99.70% | NM_007294.4 | synonymous | Benign | 3997 |

¹ Based on Clinvar version 20200329

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