

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: 27 Dec 2023 1 of 2

Sample Information

Patient Name: 張珈瑋 Gender: Female ID No.: N223428162 History No.: 47860991

Age: 44

Ordering Doctor: DOC2589K 曾令民

Ordering REQ.: D79A931 Signing in Date: 2023/12/27

Path No.: M112-00340 **MP No.:** BR23093

Assay: Oncomine BRCA1/2 Assay

Sample Type: Blood

Date of blood drawing: 2023/12/25

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

Note:

Sample Cancer Type: Breast Cancer

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Report Highlights
0 Therapies Available
0 Clinical Trials

Relevant Breast Cancer Variants

| Gene | Finding |
|-------|---------------|
| BRCA1 | None detected |
| BRCA2 | None 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.(N372H) | c.1114A>C | chr13:32906729 | 49.54% | NM_000059.3 | missense | Benign | 3999 | |
| BRCA2 | p.(V1269=) | c.3807T>C | chr13:32912299 | 50.23% | NM_000059.3 | synonymous | Benign | 3992 | |

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 |
|-------|-------------------|-----------|----------------|---------------------|-------------|----------------|----------------------|----------|
| BRCA2 | p.(L1521=) | c.4563A>G | chr13:32913055 | 99.94% | NM_000059.3 | synonymous | Benign | 3386 |
| BRCA2 | p.(V2171=) | c.6513G>C | chr13:32915005 | 99.97% | NM_000059.3 | synonymous | Benign | 4000 |
| BRCA2 | p.(V2466A) | c.7397T>C | chr13:32929387 | 99.85% | NM_000059.3 | missense | Benign | 4000 |
| BRCA1 | p.(S1613G) | c.4837A>G | chr17:41223094 | 49.97% | NM_007294.4 | missense | Benign | 3992 |
| BRCA1 | p.(S1436=) | c.4308T>C | chr17:41234470 | 49.63% | NM_007294.4 | synonymous | Benign | 4000 |
| BRCA1 | p.(K1183R) | c.3548A>G | chr17:41244000 | 49.69% | NM_007294.4 | missense | Benign | 3999 |
| BRCA1 | p.(E1038G) | c.3113A>G | chr17:41244435 | 50.46% | NM_007294.4 | missense | Benign | 3999 |
| BRCA1 | p.(P871L) | c.2612C>T | chr17:41244936 | 49.51% | NM_007294.4 | missense | Benign | 3999 |
| BRCA1 | p.(L771=) | c.2311T>C | chr17:41245237 | 51.65% | NM_007294.4 | synonymous | Benign | 3996 |
| BRCA1 | p.(S694=) | c.2082C>T | chr17:41245466 | 51.19% | NM_007294.4 | synonymous | Benign | 3995 |

¹ Based on Clinvar version 20200329