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Date: 29 Jan 2024

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Sample Information

Patient Name: 陳品樺 Gender: Female ID No.: A220880865 History No.: 48876319

Age: 52

Ordering Doctor: DOC2456B 游暄萱

Ordering REQ.: 0CWEUUW Signing in Date: 2024/1/22

Path No.: M113-00023 **MP No.:** BR24006

Assay: Oncomine BRCA1/2 Assay

Sample Type: Blood

Date of blood drawing: 2024/1/18

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.(?) | c26G>A | chr13:32890572 | 51.65% | NM_000059.3 | unknown | Benign | 3996 | |
| BRCA2 | p.(K1132=) | c.3396A>G | chr13:32911888 | 52.69% | NM_000059.3 | synonymous | 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 | ClinVar1 | Coverage |
|-------|-------------------|-----------|----------------|---------------------|-------------|----------------|----------|----------|
| BRCA2 | p.(V1269=) | c.3807T>C | chr13:32912299 | 50.17% | NM_000059.3 | synonymous | Benign | 3444 |
| BRCA2 | p.(L1521=) | c.4563A>G | chr13:32913055 | 100.00% | NM_000059.3 | synonymous | Benign | 1808 |
| BRCA2 | p.(V2171=) | c.6513G>C | chr13:32915005 | 99.97% | NM_000059.3 | synonymous | Benign | 3997 |
| BRCA2 | p.(S2414=) | c.7242A>G | chr13:32929232 | 50.32% | NM_000059.3 | synonymous | Benign | 2784 |
| BRCA2 | p.(V2466A) | c.7397T>C | chr13:32929387 | 99.84% | NM_000059.3 | missense | Benign | 2519 |
| BRCA1 | p.(S1613G) | c.4837A>G | chr17:41223094 | 48.60% | NM_007294.4 | missense | Benign | 3994 |
| BRCA1 | p.(S1436=) | c.4308T>C | chr17:41234470 | 50.65% | NM_007294.4 | synonymous | Benign | 2529 |
| BRCA1 | p.(K1183R) | c.3548A>G | chr17:41244000 | 50.77% | NM_007294.4 | missense | Benign | 4000 |
| BRCA1 | p.(E1038G) | c.3113A>G | chr17:41244435 | 49.46% | NM_007294.4 | missense | Benign | 3999 |
| BRCA1 | p.(P871L) | c.2612C>T | chr17:41244936 | 49.07% | NM_007294.4 | missense | Benign | 3998 |
| BRCA1 | p.(L771=) | c.2311T>C | chr17:41245237 | 49.78% | NM_007294.4 | synonymous | Benign | 2750 |
| BRCA1 | p.(S694=) | c.2082C>T | chr17:41245466 | 49.59% | NM_007294.4 | synonymous | Benign | 3999 |

¹ Based on Clinvar version 20200329