



Sample Information

Patient Name: 謝惠如  
Gender: Female  
ID No.: Q220393093  
History No.: 49568130  
Age: 49  
  
Ordering Doctor: DOC3584C 吳孟芹  
Ordering REQ.: OCRBVWL  
Signing in Date: 2023/10/04

Path No.: M112-00262  
MP No.: BR23072  
Assay: Oncomine BRCA1/2 Assay  
Sample Type: FFPE  
Block No.: N23-19263F  
Percentage of tumor cells: 50%

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

Note:

Sample Cancer Type: Ovarian Cancer

|                   |      |  |
|-------------------|------|--|
| Table of Contents | Page | Report Highlights                          |
| Variant Details   | 1    | 0 Therapies Available<br>0 Clinical Trials |

Relevant Ovarian 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 <sup>1</sup> | Coverage |
| BRCA2                 | p.(N289H)         | c.865A>C | chr13:32906480 | 67.25%           | NM_000059.3 | missense       | Benign               | 1997     |

<sup>1</sup> Based on Clinvar version 20200329

## Variant Details (continued)

### DNA Sequence Variants (continued)

| Gene  | Amino Acid Change | Coding    | Locus          | Allele Frequency | Transcript  | Variant Effect | ClinVar <sup>1</sup> | Coverage |
|-------|-------------------|-----------|----------------|------------------|-------------|----------------|----------------------|----------|
| BRCA2 | p.(S455=)         | c.1365A>G | chr13:32906980 | 65.70%           | NM_000059.3 | synonymous     | Benign               | 2000     |
| BRCA2 | p.(H743=)         | c.2229T>C | chr13:32910721 | 69.37%           | NM_000059.3 | synonymous     | Benign               | 1998     |
| BRCA2 | p.(N991D)         | c.2971A>G | chr13:32911463 | 71.02%           | NM_000059.3 | missense       | Benign               | 1991     |
| BRCA2 | p.(V1269=)        | c.3807T>C | chr13:32912299 | 30.80%           | NM_000059.3 | synonymous     | Benign               | 2000     |
| BRCA2 | p.(L1521=)        | c.4563A>G | chr13:32913055 | 99.80%           | NM_000059.3 | synonymous     | Benign               | 1993     |
| BRCA2 | p.(V2171=)        | c.6513G>C | chr13:32915005 | 100.00%          | NM_000059.3 | synonymous     | Benign               | 1998     |
| BRCA2 | p.(V2466A)        | c.7397T>C | chr13:32929387 | 99.95%           | NM_000059.3 | missense       | Benign               | 1999     |

<sup>1</sup> Based on Clinvar version 20200329