Effective Date: December 1, 2025

## **TEST CHANGE**

Malignancy Risk Assessment, Pelvic Mass, OVA1 Plus 3002309, OVA1 PLUS

| 0002003, 0 1711 1 200         |   |
|-------------------------------|---|
| Specimen Requirements:        |   |
| Patient Preparation:          | Testing should not be performed on patients 17 years of age or younger.   |
| Collect:                      | Serum <u>separator tube</u> Separator Tube (SST).   |
| Specimen Preparation:         | Transfer 2.2 mL serum to an ARUP <u>standard transport</u> <u>tube.</u> Standard Transport Tube. (Min: 1.1 mL)  |
| Transport Temperature:        | Frozen. Also acceptable: Refrigerated.  |
| Unacceptable Conditions:      |   |
| Remarks:                      | Menopausal <u>s</u> Status required at time of ordering.  |
| Stability:                    | Ambient: Unacceptable; Refrigerated: 8 days; Frozen: 9 weeks  |
| Methodology:                  | Electrochemiluminescent Immunoassay (ECLIA) / Fixed-Rate-Time Nephelometry  |
| Note:                         | OVA1 Biomarkers: CA-125 II, <u>a</u> Apoliproprotein A1 (Apo A-1),<br><u>b</u> Beta-2 <u>m</u> Microglobulin (B2M), <u>t</u> Transferrin, and <u>p</u> Prealbumin.                            |
|                               | OVERA Biomarkers: Apolipoprotein A1 (Apo A-1), HE4 (human epididymisHuman Epididymis protein 4), CA-125 II, FSH (follicle stimulating hormoneFollicle Stimulating Hormone), and taransferrin. |
| CPT Codes:                    | 81503   |
| New York DOH Approval Status: | This test is New York DOH approved.   |
| Interpretive Data:            |   |
| Reference Interval:           |   |
| By Report                     |   |

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