

TEST CHANGE

Maternal Screening, Sequential, Specimen #2, Alpha Fetoprotein, hCG, Estriol, and Inhibin A

3000148, MS SEQ2

Specimen Requirements:

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| Patient Preparation: | Specimen must be drawn between 14 weeks, 0 days and 24 weeks, 6 days gestation. The recommended time for maternal serum screening is 16 to 18 weeks gestation. |
| Collect: | Serum separator tube (SST) or plain red. |
| Specimen Preparation: | Separate from cells ASAP or within 2 hours of collection. Transfer 3 mL serum to an ARUP standard transport tube. (Min: 1 mL) |
| Transport Temperature: | Refrigerated. |
| Unacceptable Conditions: | Plasma. Hemolyzed specimens. |
| Remarks: | Requires that a previous first trimester specimen, Maternal Screening, Sequential, Specimen #1, hCG, PAPP-A, NT (ARUP test code 3000146), has been performed. |
| Stability: | After separation from cells: Ambient: 72 hours; Refrigerated: 2 weeks; Frozen: 1 year (Avoid repeated freeze/thaw cycles.) |
| Methodology: | Quantitative Chemiluminescent Immunoassay (CLIA) |
| Note: | This test is used to screen for fetal risk of Down syndrome (trisomy 21), trisomy 18, and open neural tube defect (ONTD, spina bifida). |
| CPT Codes: | 81511 |
| New York DOH Approval Status: | This test is New York DOH approved. |
| Interpretive Data: | |
| Refer to report. | |
| Reference Interval: | |
| By report | |

HOTLINE NOTE: There is a component change associated with this test. One or more components have been added or removed. Refer to the Hotline Test Mix for interface build information.