

## **TEST CHANGE**

| Specimen Requirements:   Patient Preparation:   Collect: Serum separator tube. Also acceptable: Plain red, lavender (EDTA), pink (K2EDTA), or green (sodium or lithium heparin).   Specimen Preparation: Separate serum or plasma from cells ASAP or within 2 hours of |
|--|
| Collect: Serum separator tube. Also acceptable: Plain red, lavender<br>(EDTA), pink (K2EDTA), or green (sodium or lithium heparin).  |
| (EDTA), pink (K2EDTA), or green (sodium or lithium heparin).   |
| Specimen Preparation: Separate serum or plasma from cells ASAP or within 2 hours of  |
| collection. Transfer 0.5 mL serum or plasma to an ARUP<br>Standard Transport Tube and freeze immediately. (Min: 0.25<br>mL)  |
| Transport Temperature: CRITICAL FROZEN. Separate specimens must be submitted when multiple tests are ordered.  |
| Unacceptable Conditions: Refrigerated or room temperature specimens.   |
| Remarks:   |
| Stability:After separation from cells: Ambient: Unacceptable;Refrigerated: Unacceptable; Frozen: 6 months  |
| Methodology: Quantitative High Performance Liquid Chromatography-<br>Tandem Mass Spectrometry  |
| Performed: Mon-Fri   |
| Reported: 1- <u>5</u> 4 days   |
| Note:  |
| CPT Codes: 84143   |
| New York DOH Approval Status: This test is New York DOH approved.  |

Interpretive Data:

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

Reference Interval:

By report