| TEST CHANGE |  |
| :---: | :---: |
| Hepatitis C Virus Antibody by CIA with Reflex to HCV by Quantitative NAAT |  |
| 2010784, HCV AB QR |  |
| Specimen Requirements: |  |
| Patient Preparation: |  |
| Collect: | Serum Separator Tube (SST). Also acceptable: Lavender (EDTA) or Pink (K2EDTA). |
| Specimen Preparation: | Separate from cells within 6 hours of collection. Transfer 2.5 mL serum or plasma to an ARUP Standard Transport Tube. <br> (Min: 1.5 mL ) <br> This test requires a dedicated transport tube submitted only for HCV AB QR testing. |
| Transport Temperature: | Frozen. |
| Unacceptable Conditions: | Specimens containing particulate material. Severely hemolyzed, heat-inactivated, or lipemic specimens. Heparinized plasma. |
| Remarks: |  |
| Stability: | After separation from cells: Ambient: $\underline{24}$ hoursUnacceptable; Refrigerated: 5 days; Frozen: 2 months (avoid freeze/thaw cycles) |
| Methodology: | Qualitative Chemiluminescent Immunoassay (CLIA) \&Quantitative Transcription-Mediated Amplification (TMA) |
| Performed: | Sun-Sat |
| Reported: | 1-2 days |
| Note: | If the anti-HCV screening result is low positive or high positive, the Hepatitis C Virus by Quantitative NAAT will be added. Additional charges apply. |
| CPT Codes: | 86803; if reflexed, add 87522 |
| New York DOH Approval Status: | This test is New York DOH approved. |
| Interpretive Data: |  |
| This assay should not be used for screening Human Cell, Tissues, a | blood donor screening, associated re-entry protocols, or for d Cellular and Tissue-Based Products (HCT/P). |


| Components | Interpretation |
| :--- | :--- |
| Hepatitis C | 0.79 IV or less |
| Antibody by CIA | Negative 0.80 to |
| Index | 0.99 IV Equivocal |
|  | 1.00 to 10.99 IV |
|  | Low Positive |
|  | 11.00 IV or |
|  | greater High |
|  | Positive |

Reference Interval:

| Test <br> Number | Components | Reference Interval |
| :--- | :--- | :--- |
|  | Hepatitis C Antibody by CIA Interp | Negative |

