TEST CHANGE

17-Hydroxyprogesterone 60-Min Quantitative by HPLC-MS/MS, Serum or Plasma

2009480, OHPRGSTN60

Specimen Requirements:	
Patient Preparation:	
Collect:	Serum separator tube. Also acceptable: Plain red, pink (K2EDTA), plasma separator tube, green (sodium heparin), or green (lithium heparin).
Specimen Preparation:	Transfer 1 mL serum or plasma to an ARUP standard transport tube.~~Standard Transport Tube.~~ (Min: 0.3 mL)
Transport Temperature:	Refrigerated. Also acceptable: Frozen.
Unacceptable Conditions:	Grossly hemolyzed specimens.
Remarks:	
Stability:	After separation from cells: Ambient: Unacceptable; Refrigerated: 1 week; Frozen: 6 months
Methodology:	Quantitative High Performance Liquid Chromatography-Tandem Mass Spectrometry
Performed:	Sun-Sat
Reported:	1-54 days
Note:	
CPT Codes:	83498
New York DOH Approval Status:	This test is New York DOH approved.

Interpretive Data:

Reference ranges for 17-h~~H~~ydroxyprogesterone following stimulation are not well defined and are dependent on the stimulation method utilized.

~~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: