

## TEST CHANGE

### 17-Hydroxyprogesterone 30-Min Quantitative by HPLC-MS/MS, Serum or Plasma 2009478, OHPRGSTN30

#### 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-~~5~~4 days

##### Note:

**CPT Codes:** 83498

**New York DOH Approval Status:** This test is New York DOH approved.

##### Interpretive Data:

Reference ranges for 17-~~h~~Hydroxyprogesterone 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:

