

Client: Example Client ABC123  
123 Test Drive  
Salt Lake City, UT 84108  
UNITED STATES

Physician: Doctor, Example

**Patient: Patient, Example**

**DOB:** 6/28/1953  
**Gender:** Male  
**Patient Identifiers:** 01234567890ABCD, 012345  
**Visit Number (FIN):** 01234567890ABCD  
**Collection Date:** 00/00/0000 00:00

**Prostate Specific Antigen, Ultrasensitive**

ARUP test code 0098581

PSA, Ultrasensitive

**5.00 ng/mL H (Ref Interval: 0.00-4.00)**

INTERPRETIVE INFORMATION: PSA Ultra Sensitive  
After radical prostatectomy, the reference interval is less than 0.05 ng/mL if there is no residual disease. In healthy males without prostatectomy, the reference interval is 4.00 ng/mL or less. The lower limit of detection is 0.01 ng/mL.

The Roche PSA electrochemiluminescent immunoassay was used. Results obtained with different assay methods or kits cannot be used interchangeably. The Roche PSA method is approved for use as an aid in the detection of prostate cancer when used in conjunction with a digital rectal exam in men age 50 and older. The Roche PSA method is also indicated for the serial measurement of PSA to aid in the prognosis and management of prostate cancer patients. Elevated PSA concentrations can only suggest the presence of prostate cancer until biopsy is performed. PSA concentrations can also be elevated in benign prostatic hyperplasia or inflammatory conditions of the prostate. PSA is generally not elevated in healthy men or men with nonprostatic carcinoma.

VERIFIED/REPORTED DATES

Procedure	Accession	Collected	Received	Verified/Reported
PSA, Ultrasensitive	20-351-117431	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART

**H=High, L=Low, \*=Abnormal, C=Critical**

Unless otherwise indicated, testing performed at: