Result Status: Final

# Labcorp

Patient Name:	Account Number:
Patient Phone:	
Date of Birth (Age):	Account Name:
Sex:	
Referring Dr (NPI #):	Collection Date/Time:
Patient ID:	Received Date/Time:
Specimen ID:	Reported Date/Time:

General Comments and Additional Information

Total Vol:	Source:
Total Vol:	Source:

Result Name		Flag	Result	Range/Units	Status	Lab
002139 CEA						
CEA			2.5	0.0-4.7 / ng/mL	Final	01
			onsmokers	<3.9 <5.6		
	Roche Diagnostic	s Electrochemi	luminescence Immu	unoassay		
	(ECLIA)					
Values obtained with different assay methods or kits						
	cannot be used i	nterchangeably	. Results cannot	: be		
	interpreted as a	bsolute eviden	ce of the present	ee or		
	absence of malig	nant disease.				
002253 AFP, Serum, Tu	umor Marker					

#### 002253 AFP, Serum, Tumor Marker

AFP, Serum, Tumor Marker <1.8 0.0-8.4 / ng/mL Final 01

Roche Diagnostics Electrochemiluminescence Immunoassay (ECLIA)

Values obtained with different assay methods or kits cannot be

Result Status: Final

## Labcorp

Patient Name:	Account Number:
Patient Phone:	
Date of Birth (Age):	Account Name:
Sex:	
Referring Dr (NPI #):	Collection Date/Time:
Patient ID:	Received Date/Time:
Specimen ID:	Reported Date/Time:

Result Name Flag Result Range/Units Status Lab

used interchangeably. Results cannot be interpreted as absolute

evidence of the presence or absence of malignant disease.

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This test is not interpretable in pregnant females.

#### 002261 CA 19-9

CA 19-9 12 0-35 / U/mL Final 01

Roche Diagnostics Electrochemiluminescence Immunoassay (ECLIA)

.

Values obtained with different assay methods or kits cannot be

used interchangeably. Results cannot be interpreted as absolute

evidence of the presence or absence of malignant disease.

### 002303 Cancer Antigen (CA) 125

Cancer Antigen (CA) 125 14.0 Not Estab. / U/mL Final 01

Roche Diagnostics Electrochemiluminescence Immunoassay (ECLIA)

.

Values obtained with different assay methods or kits cannot be used interchangeably. Results cannot be interpreted as absolute

evidence of the presence or absence of malignant disease.

### 010322 Prostate-Specific Ag

Result Status: Final

# Labcorp

Patient Name:

Patient Phone:

Date of Birth (Age):

Sex:

Referring Dr (NPI #):

Patient ID:

Specimen ID:

Account Number:

Account Name:

Collection Date/Time:

Received Date/Time:

Reported Date/Time:

Result Name	Flag	Result	Range/Units	Status	Lab
Prostate Specific Ag		1.3	0.0-4.0 / ng/mL	Final	01

Roche ECLIA methodology.

According to the American Urological Association, Serum PSA should decrease and remain at undetectable levels after radical prostatectomy. The AUA defines biochemical recurrence as an initial PSA value 0.2 ng/mL or greater followed by a subsequent confirmatory PSA value 0.2 ng/mL or greater.

Values obtained with different assay methods or kits cannot be used interchangeably. Results cannot be interpreted as absolute evidence of the presence or absence of malignant disease.

#### **Performing Lab**

01 - Labcorp San Diego, 13112 Evening Creek Dr So Ste 200, San Diego, CA 92128-4108, (858) 668-3700, Galloway, Jenny R MD For Inquiries, the physician may contact the performing lab.

#### **END OF REPORT**