



## Patient Report

Specimen ID: 297-298-1445-0

Acct #: 09018260

Phone: (813) 498-1070

Rte: 00

Control ID: B0082220747

4PMD

4910 E Adamo Dr Unit B

TAMPA FL 33605



## Patient Details

DOB: 02/27/1978

Age(y/m/d): 040/07/27

Gender: M SSN:

Patient ID: [REDACTED]

## Specimen Details

Date collected: 10/24/2018 0918 Local

Date received: 10/25/2018

Date entered: 10/25/2018

Date reported: 11/01/2018 0831 ET

## Physician Details

Ordering: R MCCLAIN

Referring:

ID: 15759893

NPI: 1487804308

## General Comments &amp; Additional Information

Clinical Info: [REDACTED]

Alternate Control Number: B0082220747

Total Volume: Not Provided

Alternate Patient ID: Not Provided

Fasting: Yes

## Ordered Items

CBC With Differential/Platelet; Comp. Metabolic Panel (14); Lipid Panel w/ Chol/HDL Ratio; Testosterone, Free+Total LC/MS; Dihydrotestosterone; Thyroxine (T4) Free, Direct, S; DHEA-Sulfate; TSH; Prolactin; Prostate-Specific Ag, Serum; IGF-1; Estradiol, Sensitive; Triiodothyronine (T3), Free; Sex Horm Binding Glob, Serum; Venipuncture

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
CBC With Differential/Platelet					
WBC	4.1		x10E3/uL	3.4 - 10.8	01
RBC	5.17		x10E6/uL	4.14 - 5.80	01
Hemoglobin	15.8		g/dL	13.0 - 17.7	01
Hematocrit	47.5		%	37.5 - 51.0	01
MCV	92		fL	79 - 97	01
MCH	30.6		pg	26.6 - 33.0	01
MCHC	33.3		g/dL	31.5 - 35.7	01
RDW	14.0		%	12.3 - 15.4	01
Platelets	216		x10E3/uL	150 - 379	01
Neutrophils	55		%	Not Estab.	01
Lymphs	29		%	Not Estab.	01
Monocytes	10		%	Not Estab.	01
Eos	5		%	Not Estab.	01
Basos	1		%	Not Estab.	01
Neutrophils (Absolute)	2.3		x10E3/uL	1.4 - 7.0	01
Lymphs (Absolute)	1.2		x10E3/uL	0.7 - 3.1	01
Monocytes (Absolute)	0.4		x10E3/uL	0.1 - 0.9	01
Eos (Absolute)	0.2		x10E3/uL	0.0 - 0.4	01
Baso (Absolute)	0.1		x10E3/uL	0.0 - 0.2	01
Immature Granulocytes	0		%	Not Estab.	01
Immature Grans (Abs)	0.0		x10E3/uL	0.0 - 0.1	01
Comp. Metabolic Panel (14)					
Glucose	87		mg/dL	65 - 99	01
BUN	13		mg/dL	6 - 24	01
Creatinine	1.02		mg/dL	0.76 - 1.27	01

Date Issued: 11/01/18 1049 ET

## FINAL REPORT

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Patient: [REDACTED]  
DOB: 02/27/1978

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Control ID: B0082220747

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Date collected: 10/24/2018 0918 Local

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
eGFR If NonAfricn Am	92		mL/min/1.73	>59	
eGFR If Africn Am	106		mL/min/1.73	>59	
BUN/Creatinine Ratio	13			9 - 20	
Sodium	139		mmol/L	134 - 144	01
Potassium	4.5		mmol/L	3.5 - 5.2	01
Chloride	100		mmol/L	96 - 106	01
Carbon Dioxide, Total	23		mmol/L	20 - 29	01
Calcium	9.5		mg/dL	8.7 - 10.2	01
Protein, Total	6.2		g/dL	6.0 - 8.5	01
Albumin	4.1		g/dL	3.5 - 5.5	01
Globulin, Total	2.1		g/dL	1.5 - 4.5	
A/G Ratio	2.0			1.2 - 2.2	
Bilirubin, Total	0.6		mg/dL	0.0 - 1.2	01
Alkaline Phosphatase	48		IU/L	39 - 117	01
AST (SGOT)	57	High	IU/L	0 - 40	01
ALT (SGPT)	34		IU/L	0 - 44	01

## Lipid Panel w/ Chol/HDL Ratio

Cholesterol, Total	148		mg/dL	100 - 199	01
Triglycerides	54		mg/dL	0 - 149	01
HDL Cholesterol	53		mg/dL	>39	01
VLDL Cholesterol Cal	11		mg/dL	5 - 40	
LDL Cholesterol Calc	84		mg/dL	0 - 99	
T. Chol/HDL Ratio	2.8		ratio	0.0 - 5.0	
Please Note:					01

### T. Chol/HDL Ratio

	Men	Women
1/2 Avg.Risk	3.4	3.3
Avg.Risk	5.0	4.4
2X Avg.Risk	9.6	7.1
3X Avg.Risk	23.4	11.0

## Testosterone, Free+Total LC/MS

Testosterone, Total, LC/MS	1018.1	High	ng/dL	264.0 - 916.0	02
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This LabCorp LC/MS-MS method is currently certified by the CDC Hormone Standardization Program (HoSt). Adult male reference interval is based on a population of healthy nonobese males (BMI <30) between 19 and 39 years old. Travison, et.al. JCEM 2017,102;1161-1173. PMID: 28324103.

Disclaimer: 02

This test was developed and its performance characteristics determined by LabCorp. It has not been cleared or approved by the Food and Drug Administration.

Free Testosterone (Direct)	21.1		pg/mL	6.8 - 21.5	02
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Dihydrotestosterone	117	High	ng/dL		03
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TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
<b>Reference Range:</b>					
<b>Adult Male: 30 - 85</b>					
<b>Thyroxine (T4) Free, Direct, S</b>					
T4, Free (Direct)	1.50		ng/dL	0.82 - 1.77	01
<b>DHEA-Sulfate</b>	127.0		ug/dL	102.6 - 416.3	01
<b>TSH</b>	1.680		uIU/mL	0.450 - 4.500	01
<b>Prolactin</b>	9.3		ng/mL	4.0 - 15.2	01
<b>Prostate-Specific Ag, Serum</b>					
Prostate Specific Ag, Serum	1.7		ng/mL	0.0 - 4.0	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.					
<b>IGF-1</b>					
Insulin-Like Growth Factor I	250	High	ng/mL	83 - 233	02
<b>Estradiol, Sensitive</b>	13.9		pg/mL	8.0 - 35.0	02
This test was developed and its performance characteristics determined by LabCorp. It has not been cleared by the Food and Drug Administration.					
Methodology: Liquid chromatography tandem mass spectrometry (LC/MS/MS)					
<b>Triiodothyronine (T3), Free</b>	3.6		pg/mL	2.0 - 4.4	01
<b>Sex Horm Binding Glob, Serum</b>	42.8		nmol/L	16.5 - 55.9	01

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02	BN	LabCorp Burlington 1447 York Court, Burlington, NC 27215-3361	Dir: William F Hancock, MD
03	ES	Esoterix Inc 4301 Lost Hills Road, Calabasas Hills, CA 91301-5358	Dir: Samuel Pepkowitz, MD

For inquiries, the physician may contact **Branch: 800-877-5227 Lab: 972-598-6000**