



Patient Report

Specimen ID: 072-319-0005-0
Control ID: B0056006444

Acct #: 0000000000

Phone: (813) 445-7342

Rte: 00

Defy Medical, LLC
4809 N. Armenia Ave. Ste 220
Tampa FL 33603



Patient Details

DOB: 03/01/1982
Age(y/m/d): 035/01/13
Gender: M SSN:
Patient ID:

Specimen Details

Date collected: 03/13/2017 0753 Local
Date entered: 03/13/2017
Date reported: 03/17/2017 0820 ET

Physician Details

Ordering: J SAYA
Referring:
ID: 12040542
NPI: 1093940041

General Comments & Additional Information

Alternate Control Number: B0056006444
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 and Total; Thyroxine (T4) Free, Direct, S; DHEA-Sulfate; TSH; Prostate-Specific Ag, Serum; Estradiol, Sensitive; Triiodothyronine, Free, Serum; Sex Horm Binding Glob, Serum; Drawing Fee; Cardiovascular Report

TESTS	RESULT	FLAG	UNITS	REFERENCE	INTERVAL	LAB
CBC With Differential/Platelet						
WBC	5.04		thous/uL	4.80 - 10.80		01
RBC	4.98		mill/uL	4.70 - 6.10		01
Hemoglobin	15.8		gm/dL	14.0 - 18.0		01
Hematocrit	47.1		%	42.0 - 52.0		01
MCV	94.6		fL	83.0 - 97.0		01
MCH	31.7	High	pg	27.0 - 31.0		01
MCHC	33.5		gm/dL	32.3 - 36.5		01
RDW	12.1		%	11.5 - 15.5		01
Platelets	273		thous/uL	130 - 450		01
Neutrophils	65.7		%			01
Lymphs	25.2		%			01
Monocytes	6.5		%			01
Eos	1.4		%			01
Basos	1.0		%			01
Neutrophils (Absolute)	3.31		thous/uL	1.50 - 7.50		01
Lymphs (Absolute)	1.27		thous/uL	1.10 - 3.00		01
Monocytes (Absolute)	0.33		thous/uL	0.10 - 0.75		01
Eos (Absolute)	0.07		thous/uL	0.00 - 0.50		01
Baso (Absolute)	0.05		thous/uL	0.00 - 0.10		01
Immature Granulocytes	0.2		%			01
Immature Grans (Abs)	0.01		thous/uL	0.00 - 0.10		01
Comp. Metabolic Panel (14)						
Glucose, Serum	79		mg/dL	65 - 99		01
BUN	20		mg/dL	6 - 20		01
Creatinine, Serum	1.09		mg/dL	0.76 - 1.27		01
eGFR If NonAfricn Am	81.82		mL/min/1.73 m2			01

Date Issued: 03/17/17 0824 ET

FINAL REPORT

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Patient ID: [REDACTED]
DOB: 01/30/1982 Patient ID:

Control ID: B0056006444

Specimen ID: 072-319-0005-0
Date collected: 03/13/2017 0753 Local

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
eGFR If Africn Am	99.00		mL/min/1.73 m2		01
BUN/Creatinine Ratio	18		Ratio	6 - 20	01
Sodium, Serum	140		mmol/L	134 - 144	01
Potassium, Serum	4.4		mmol/L	3.5 - 5.2	01
Chloride, Serum	100		mmol/L	97 - 108	01
Carbon Dioxide, Total	27		mmol/L	18 - 29	01
Calcium, Serum	9.3		mg/dL	8.7 - 10.2	01
Protein, Total, Serum	6.8		gm/dL	6.0 - 8.5	01
Albumin, Serum	4.5		gm/dL	3.5 - 5.5	01
Globulin, Total	2.3		gm/dL	2.1 - 4.0	01
A/G Ratio	2.0				01
Bilirubin, Total	1.0		mg/dL	<=1.2	01
Alkaline Phosphatase, S	70		Units/L	39 - 117	01
AST (SGOT)	28		IU/L	<=40	01
ALT (SGPT)	30		IU/L	<=44	01
Lipid Panel w/ Chol/HDL Ratio					
Cholesterol, Total	212	High	mg/dL	100 - 199	01
Triglycerides	41		mg/dL	<=149	01
HDL Cholesterol	>121		mg/dL	>=39	01
Risk Factors for CHD					
Major risk <40 mg/dl					
Normal range 40-59 mg/dl					
Negative risk >= 60 mg/dl					
VLDL Cholesterol Cal	8		mg/dL		01
LDL Cholesterol Calc	<83		mg/dL		01
LDL Reference Range					
Optimal <100 mg/dl					
Near optimal 100-129 mg/dl					
Borderline high 130-159 mg/dl					
High 160-189 mg/dl					
Very high >= 190 mg/dl					
T. Chol/HDL Ratio	<1.8		ratio	0.0 - 4.9	01
Testosterone, Free and Total					
Testosterone, Serum	954		ng/dL	348 - 1197	02
Comment:					
Adult male reference interval is based on a population of lean males up to 40 years old.					
Free Testosterone (Direct)	22.0		pg/mL	8.7 - 25.1	03
Thyroxine (T4) Free, Direct, S					
T4, Free (Direct)	0.95		ng/dL	0.82 - 1.77	01
DHEA-Sulfate	379.2		ug/dL	102.6 - 416.3	02

Patient: [REDACTED]
DOB: 01/30/1962 Patient ID:

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TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
TSH	1.950		IU/mL	0.450 - 4.500	01
Prostate-Specific Ag, Serum					
Prostate Specific Ag, Serum	0.8		ng/mL	0.0 - 4.0	02
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.					
Estradiol, Sensitive	26.8		pg/mL	8.0 - 35.0	03
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)					
Triiodothyronine, Free, Serum	2.0		pg/mL	2.0 - 4.4	02
Sex Horm Binding Glob, Serum	52.3		nmol/L	16.5 - 55.9	02
Cardiovascular Report					
Interpretation	Note				04
Supplement report is available.					
PDF Image	.				04

01	GENIL	Genesis Med Ctr Silvas 855 Illini Drive Ste 203, Silvis, IL 61282-2904	Dir: Carolyn Miller, MD
02	DA	LabCorp Dallas 7777 Forest Lane Suite C350, Dallas, TX 75230-2544	Dir: CN Etufugh, MD
03	BN	LabCorp Burlington 1447 York Court, Burlington, NC 27215-3361	Dir: William F Hancock, MD
04	LITIL	Litholink Corporation 2250 West Campbell Park Drive, Chicago, IL 60612-3502	Dir: Mitchell Laks, PhD

For inquiries, the physician may contact **Branch: 800-877-5227 Lab: 309-281-4826**

Accessions: 07231900050

DISCLAIMER: These assessments and treatment suggestions are provided as a convenience in support of the physician-patient relationship and are not intended to replace the physician's clinical judgment. They are derived from the national guidelines in addition to other evidence and expert opinion. The clinician should consider this information within the context of clinical opinion and the individual patient.

SEE GUIDANCE FOR CARDIOVASCULAR REPORT: National Heart, Lung, and Blood Institute's Third Report of the NCEP Expert Panel on Detection, Evaluation and Treatment of High Blood Cholesterol in Adults (ATP III) (2002. NIH publication 02-5215); Brunzell et al. Diabetes Care 2008; 31(4):811-82; Contois et al. Clin Chem 2009; 55(3):407-419; Stone NJ et al. 2013 ACC/AHA guideline on the treatment of blood cholesterol to reduce atherosclerotic cardiovascular risk in adults: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines. Circulation 2014;129(suppl 2):S1-S45.

Note: Please refer to your LabCorp Report for all results as well as any test-specific and specimen-specific comments.

Laboratory Director's Notes

Laboratory test values flagged with an asterisk (*) within this report refer to the following commentary from our physicians and quality assurance staff.

COLLECTION DATE	ITEM	RELATED NOTES
03/13/2017	HDL-C	Risk Factors for CHD Major risk <40 mg/dl Normal range 40-59 mg/dl Negative risk >= 60 mg/dl
03/13/2017	LDL(calc)	LDL Reference Range Optimal <100 mg/dl Near optimal 100-129 mg/dl Borderline high 130-159 mg/dl High 160-189 mg/dl Very high >= 190 mg/dl

Mitchell S. Laks, PhD - Laboratory Director

Current Laboratory Results

Blood Draw Date: 03/13/2017 **Date Received:** 03/13/2017 **Date Completed:** 03/13/2017 **Fasting:** YES

Comp. Metabolic Panel (14)

ANALYTE	REF. INTERVAL	LOW	HIGH	RESULT
Glucose mg/dL	65-99			79
BUN mg/dL	6-20			20
Creatinine mg/dL	0.76-1.27			1.09
Sodium mmol/L	134-144			140
Potassium mmol/L	3.5-5.2			4.4
Chloride mmol/L	97-108			100
Carbon Dioxide mmol/L	18-29			27
Calcium mg/dL	8.7-10.2			9.3
Protein, Total, Serum g/dL	6.0-8.5			6.8
Albumin g/dL	3.5-5.5			4.5
Globulin, Total g/dL	2.1-4			2.3
A/G Ratio	1.1 - 2.5			2.0
Bilirubin, Total mg/dL	<=1.2			1.0
Alkaline Phosphatase, S IU/L	39-117			70
AST IU/L	<=40			28
ALT IU/L	<=44			30
BUN: Creatinine Ratio	6-20			18
Anion Gap mmol/L	8 - 14			13
estimated GFR mL/min/1.73mE2	> 59			82

Albumin testing performed on the Roche Modular using the ALB PLUS assay.

Lipid Panel with Chol/HDL Ratio

ANALYTE	REF. INTERVAL	LOW	HIGH	RESULT
Total Cholesterol mg/dL	100-199			212 H
Triglyceride mg/dL	<=149			41
HDL-C mg/dL	>=39			* >121
VLDL mg/dL	5 - 40			8
LDL(calc) mg/dL	0 - 99			* <83
non-HDL cholesterol mg/dL	0 - 129			91
Total Chol:HDL Ratio ratio units	0.0-4.9			<1.8

TSH

ANALYTE	REF. INTERVAL	LOW	HIGH	RESULT
TSH uIU/mL	0.450-4.5			1.950

Legend for Abnormal Flags:

L - Below Low Normal LL - Alert Low < - Panic Low A - Abnormal (applies to non-numeric results)
H - Above High Normal HH - Alert High > - Panic High AA - Critical Abnormal (applies to non-numeric results)

Cardiovascular Report

Patient Assessment

Current available clinical information suggests the patient's risk is at least LOW. If the patient has two or more major risk factors, the risk category is intermediate. If the patient has CHD or a CHD risk equivalent, the risk category is high. If patient does not have CHD or a CHD risk equivalent, consider use of the Pooled Cohort Equations to estimate 10-year CVD risk, as individuals with greater than 7.5% risk may warrant more intensive therapy. The calculator can be found at: <http://tools.cardiosource.org/ASCVD-Risk-Estimator/>

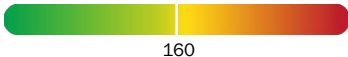


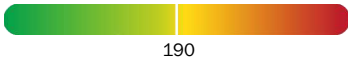


Insulin resistance, obesity, excessive alcohol use, smoking, nephrotic syndrome, liver disease, and certain medications can cause secondary dyslipidemia. Consider evaluation if clinically indicated.

Therapeutic lifestyle changes are always valuable to achieve optimal blood lipid status (diet, exercise, weight management).

Lipid Management

Select one patient risk category based upon medical history and clinical judgment. Additional risk factors such as personal or family history of premature CHD, smoking, and hypertension modify a patient's goals of therapy. In CVD prevention, the intensity of therapy should be adjusted to the level of patient risk. MODERATE intensity statin therapy generally results in an average LDL-C reduction of 30% to less than 50% from the untreated baseline. Examples include (daily doses): atorvastatin 10-20 mg, rosuvastatin 5-10 mg, simvastatin 20-40 mg, pravastatin 40-80 mg, lovastatin 40 mg. HIGH intensity statin therapy generally results in an average LDL-C reduction of 50% or more from the untreated baseline. Examples include (daily doses): atorvastatin 40-80 mg and rosuvastatin 20 mg.

▽ = PATIENT'S RESULT

	Patient Risk Category (select one)		
ANALYTE / RESULT	LOW	INTERMEDIATE	HIGH
LDL-C <83 mg/dL	▽  160	▽  100 130	▽  70 100
non-HDL 91 mg/dL	▽  190	▽  130 160	▽  100 130
Lipid Assessment	LDL-C is optimal, was 93 and now is 83 mg/dL. Non-HDL Cholesterol is optimal, was 100 and now is 91 mg/dL.	LDL-C is optimal, was 93 and now is 83 mg/dL. Non-HDL Cholesterol is optimal, was 100 and now is 91 mg/dL.	LDL-C is normal, was 93 and now is 83 mg/dL. Non-HDL Cholesterol is optimal, was 100 and now is 91 mg/dL.
Treatment Suggestions	Considerations for use of statin therapy include family history of premature atherosclerotic disease, elevated coronary artery calcium score, ankle-brachial index < 0.9, elevated CRP, or elevated lifetime CVD risk.	Consider measurement of LDL particle number or Apo B to adjudicate need for further LDL lowering therapy. Factors that may influence statin use include family history of premature atherosclerotic disease, elevated coronary artery calcium score, ankle-brachial index < 0.9, elevated CRP, or elevated lifetime CVD risk. If statin cannot be tolerated or increased, alternatives include use of an intestinal agent (ezetimibe or bile acid sequestrant) or niacin.	If at least a 50% LDL reduction from baseline has not been achieved, begin or increase statin. Consider measurement of LDL particle number or Apo B to adjudicate need for further LDL lowering therapy. If statin cannot be tolerated or increased, alternatives include use of an intestinal agent (ezetimibe or bile acid sequestrant) or niacin.

DISCLAIMER: These assessments and treatment suggestions are provided as a convenience and are neither comprehensive nor intended to replace the physician's clinical judgment. They do not include information such as family history, personal history, or physical findings as would be obtained by the clinician during patient evaluation because LabCorp does not have access to the complete patient medical record.

Patient Results Summary

Cholesterol comes in different forms and has varying effects on your heart health. Some cholesterol is “good” and not known to cause disease, this is HDL. The rest of cholesterol causes disease by clogging your arteries, this is non-HDL. LDL cholesterol is the largest component of the non-HDL cholesterol. Lowering your levels of “bad” cholesterol will lower your risk for disease.

- **LDL cholesterol (LDL-C)** is the largest component of the non-HDL cholesterol (“bad” cholesterol).
- **non-HDL** is composed of many different types of cholesterol (not just LDL-C) and high levels cause disease.

The level to which your LDL must be lowered depends on the risk for developing heart disease or having a heart attack. The higher your risk for heart disease, the lower your LDL goal.

Contributing Risk Factors For Heart Disease

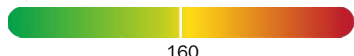
- | | |
|--|---|
| <input type="checkbox"/> Heart and/or vascular disease | <input type="checkbox"/> Cigarette (tobacco) smoking |
| <input type="checkbox"/> High blood pressure | <input type="checkbox"/> Low HDL (men less than 40 mg/dL, women less than 50 mg/dL) |
| <input type="checkbox"/> Diabetes | <input type="checkbox"/> Family history of early onset heart disease |
| <input type="checkbox"/> Chronic kidney disease | <input type="checkbox"/> Man over 45 years or woman over 55 years |
| <input type="checkbox"/> Obesity | <input type="checkbox"/> Familial Hypercholesterolemia |

Your Heart Disease Risk Category

Selected by your physician based upon your risk factors and clinical judgement.

Test /
Your Results

LDL-C
<83 mg/dL



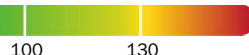
☐ Intermediate



☐ High



non-HDL
91 mg/dL



▽ = Your Result: Left (Green) = Optimal, Center = Acceptable, Right (Red) = High Risk

Your Care Plan (as selected by your physician)

- | | |
|---|---|
| <input type="checkbox"/> Eat less trans fats and saturated fats, red meat, and sugary foods/drinks | <input type="checkbox"/> Control any other medical conditions: such as diabetes, high blood pressure |
| <input type="checkbox"/> Eat more vegetables, fruits, whole grains, low-fat dairy products, poultry, fish, and nuts | <input type="checkbox"/> Visit your doctor as scheduled and obtain all follow-up tests/treatments recommended |
| <input type="checkbox"/> Exercise | <input type="checkbox"/> Take all of your medications your doctor(s) have prescribed |
| <input type="checkbox"/> Lose weight | <input type="checkbox"/> |

DISCLAIMER: You should discuss this information with your physician. Litholink does not have a doctor-patient relationship with you, nor does it have access to a complete medical history or a physical examination that would be necessary for a complete diagnosis and comprehensive treatment plan. Neither you nor your physician should rely solely on this guidance. REFERENCES: National Heart, Lung, and Blood Institute's Third Report of the NCEP Expert Panel on Detection, Evaluation and Treatment of High Blood Cholesterol in Adults (ATP III) (2002. NIH publication 02-5215); National Heart, Lung, and Blood Institute's Your Guide to Lowering Your Cholesterol with TLC (2005. NIH publication 06-5235); Stone NJ et al. 2013 ACC/AHA guideline on the treatment of blood cholesterol to reduce atherosclerotic cardiovascular risk in adults: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines. Circulation. 2013; 00:000-000.