

Specimen Details

Date collected: 10/20/2017 1028 Local
Date received: 10/20/2017
Date entered: 10/20/2017
Date reported: 10/25/2017 0715 ET

General Comments & Additional Information

Total Volume: Not Provided

Fasting: Yes

Ordered Items

NMR LipoProfile; Vitamin A and E; CBC With Differential/Platelet; Comp. Metabolic Panel (14); Lipid Panel; Iron and TIBC; Heavy Metals Profile I, Blood; Lp-PLA2 Activity; Hemoglobin A1c; Cortisol; IGF-1; Zinc, RBC; Reverse T3, Serum; Vitamin D, 25-Hydroxy; C-Reactive Protein, Cardiac; Thyroid Cascade Profile; Homocyst(e)ine, Plasma; Uric Acid, Serum; GGT; Insulin; Ferritin, Serum; Triiodothyronine, Free, Serum; Apolipoprotein A-1; Fatty Acids, Free (Nonester); Apolipoprotein B; Venipuncture

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
NMR LipoProfile					
LDL Particle Number					01
LDL-P	1279	High	nmol/L	<1000	01
		Low		< 1000	
		Moderate		1000 - 1299	
		Borderline-High		1300 - 1599	
		High		1600 - 2000	
		Very High		> 2000	
Lipids					01
LDL-C	107	High	mg/dL	0 - 99	
		Optimal		< 100	
		Above optimal		100 - 129	
		Borderline		130 - 159	
		High		160 - 189	
		Very high		> 189	
Comment:					01
LDL-C is inaccurate if patient is non-fasting.					
HDL-C	54		mg/dL	>39	01
Triglycerides	95		mg/dL	0 - 149	01
Cholesterol, Total	180		mg/dL	100 - 199	01
LDL and HDL Particles					01
HDL-P (Total)	35.0		umol/L	>=30.5	01
Small LDL-P	581	High	nmol/L	<=527	01
LDL Size	21.2		nm	>20.5	01

**** INTERPRETATIVE INFORMATION****

PARTICLE CONCENTRATION AND SIZE

<--Lower CVD Risk Higher CVD Risk-->

LDL AND HDL PARTICLES	Percentile	in Reference Population
HDL-P (total)	High	75th 50th 25th Low
	>34.9	34.9 30.5 26.7 <26.7
Small LDL-P	Low	25th 50th 75th High

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
LDL Size	<-Large (Pattern A)-> 23.0	117	527	839	>839
					<-Small (Pattern B)-> 20.6
					20.5
					19.0

Comment: 01
 Small LDL-P and LDL Size are associated with CVD risk, but not after LDL-P is taken into account. These assays were developed and their performance characteristics determined by LipoScience. These assays have not been cleared by the US Food and Drug Administration. The clinical utility of these laboratory values have not been fully established.

Insulin Resistance Score 01
LP-IR Score 55 High <=45 01

INSULIN RESISTANCE MARKER
 <--Insulin Sensitive Insulin Resistant-->
 Percentile in Reference Population
Insulin Resistance Score
 LP-IR Score Low 25th 50th 75th High
 <27 27 45 63 >63

Comment: 01
 LP-IR Score is inaccurate if patient is non-fasting. The LP-IR score is a laboratory developed index that has been associated with insulin resistance and diabetes risk and should be used as one component of a physician's clinical assessment. The LP-IR score listed above has not been cleared by the US Food and Drug Administration.

Vitamin A and E

Vitamin A, Serum	61	ug/dL	24 - 85	01
Vitamin E (Alpha Tocopherol)	13.2	mg/L	5.3 - 17.5	01

CBC With Differential/Platelet

WBC	5.6	x10E3/uL	3.4 - 10.8	02
RBC	5.47	x10E6/uL	4.14 - 5.80	02
Hemoglobin	16.5	g/dL	12.6 - 17.7	02
Hematocrit	47.5	%	37.5 - 51.0	02
MCV	87	fL	79 - 97	02
MCH	30.2	pg	26.6 - 33.0	02
MCHC	34.7	g/dL	31.5 - 35.7	02
RDW	14.2	%	12.3 - 15.4	02
Platelets	239	x10E3/uL	150 - 379	02
Neutrophils	49	%	Not Estab.	02
Lymphs	40	%	Not Estab.	02
Monocytes	6	%	Not Estab.	02
Eos	4	%	Not Estab.	02
Basos	1	%	Not Estab.	02
Neutrophils (Absolute)	2.7	x10E3/uL	1.4 - 7.0	02

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
Lymphs (Absolute)	2.2		x10E3/uL	0.7 - 3.1	02
Monocytes (Absolute)	0.4		x10E3/uL	0.1 - 0.9	02
Eos (Absolute)	0.2		x10E3/uL	0.0 - 0.4	02
Baso (Absolute)	0.0		x10E3/uL	0.0 - 0.2	02
Immature Granulocytes	0		%	Not Estab.	02
Immature Grans (Abs)	0.0		x10E3/uL	0.0 - 0.1	02
Comp. Metabolic Panel (14)					
Glucose, Serum	87		mg/dL	65 - 99	02
BUN	17		mg/dL	6 - 24	02
Creatinine, Serum	0.92		mg/dL	0.76 - 1.27	02
eGFR If NonAfricn Am	104		mL/min/1.73	>59	
eGFR If Africn Am	120		mL/min/1.73	>59	
BUN/Creatinine Ratio	18			9 - 20	
Sodium, Serum	139		mmol/L	134 - 144	02
Potassium, Serum	4.4		mmol/L	3.5 - 5.2	02
Chloride, Serum	100		mmol/L	96 - 106	02
Carbon Dioxide, Total	25		mmol/L	18 - 29	02
Calcium, Serum	9.2		mg/dL	8.7 - 10.2	02
Protein, Total, Serum	6.7		g/dL	6.0 - 8.5	02
Albumin, Serum	4.1		g/dL	3.5 - 5.5	02
Globulin, Total	2.6		g/dL	1.5 - 4.5	
A/G Ratio	1.6			1.2 - 2.2	
Bilirubin, Total	0.6		mg/dL	0.0 - 1.2	02
Alkaline Phosphatase, S	48		IU/L	39 - 117	02
AST (SGOT)	26		IU/L	0 - 40	02
ALT (SGPT)	36		IU/L	0 - 44	02
Lipid Panel					
Cholesterol, Total	179		mg/dL	100 - 199	02
Triglycerides	88		mg/dL	0 - 149	02
HDL Cholesterol	53		mg/dL	>39	02
VLDL Cholesterol Calc	18		mg/dL	5 - 40	
LDL Cholesterol Calc	108	High	mg/dL	0 - 99	
Iron and TIBC					
Iron Bind.Cap. (TIBC)	358		ug/dL	250 - 450	
UIBC	259		ug/dL	111 - 343	02
Iron, Serum	99		ug/dL	38 - 169	02
Iron Saturation	28		%	15 - 55	
Heavy Metals Profile I, Blood					
Lead, Blood	None Detected		ug/dL	0 - 19	01
Environmental Exposure:					

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
				WHO Recommendation <20	
				Occupational Exposure:	
				OSHA Lead Std 40	
				BEI 30	
				Detection Limit = 1	
Arsenic, Blood	10		ug/L	2 - 23	01
				Detection Limit = 1	
Mercury, Blood	None Detected		ug/L	0.0 - 14.9	01
				Environmental Exposure: <15.0	
				Occupational Exposure:	
				BEI - Inorganic Mercury: 15.0	
				Detection Limit = 1.0	
Lp-PLA2 Activity					
Lp-PLA2 Activity	63		nmol/min/mL	<75	03
	Relative Risk: LOW				
	Based on the documented clinical utility of Lp-PLA2 Activity to assess risk of CHD (1), the following cut-off has been defined for Lp-PLA2 Activity: A cut-off of >=75 nmol/min/mL defines a population with increased relative risk of developing CHD. (Reference: 1-The Lp-PLA2 Studies Collaboration. Lancet. 2010; 375: 1536-1544).				
	This test is performed by a Liquid Chromatography-Tandem Mass Spectrometry (LC/MS/MS) method. This test was developed and its performance characteristics determined by the Cleveland HeartLab, Inc. It has not been cleared or approved by the U.S. FDA. The Cleveland HeartLab is regulated under Clinical Laboratory Improvement Amendments (CLIA) as qualified to perform high-complexity testing. This test is used for clinical purposes. It should not be regarded as investigational or for research.				
PDF Image	.				03
Hemoglobin A1c					
Hemoglobin A1c	5.1		%	4.8 - 5.6	02
Please Note:					02
	Pre-diabetes: 5.7 - 6.4				
	Diabetes: >6.4				
	Glycemic control for adults with diabetes: <7.0				
Cortisol					
	8.2		ug/dL		02
			Cortisol AM	6.2 - 19.4	
			Cortisol PM	2.3 - 11.9	
IGF-1					
Insulin-Like Growth Factor I	172		ng/mL	83 - 233	01
Zinc, RBC					
	1326		ug/dL	878 - 1660	01
	Please note reference interval change				

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
Reverse T3, Serum	15.6		ng/dL	9.2 - 24.1	01
Vitamin D, 25-Hydroxy	42.7		ng/mL	30.0 - 100.0	02
<p>Vitamin D deficiency has been defined by the Institute of Medicine and an Endocrine Society practice guideline as a level of serum 25-OH vitamin D less than 20 ng/mL (1,2). The Endocrine Society went on to further define vitamin D insufficiency as a level between 21 and 29 ng/mL (2).</p> <p>1. IOM (Institute of Medicine). 2010. Dietary reference intakes for calcium and D. Washington DC: The National Academies Press.</p> <p>2. Holick MF, Binkley NC, Bischoff-Ferrari HA, et al. Evaluation, treatment, and prevention of vitamin D deficiency: an Endocrine Society clinical practice guideline. JCEM. 2011 Jul; 96(7):1911-30.</p>					
C-Reactive Protein, Cardiac	1.24		mg/L	0.00 - 3.00	02
<p>Relative Risk for Future Cardiovascular Event</p> <p>Low <1.00</p> <p>Average 1.00 - 3.00</p> <p>High >3.00</p>					
Thyroid Cascade Profile					
TSH	2.430		uIU/mL	0.450 - 4.500	02
Homocyst(e)ine, Plasma	8.2		umol/L	0.0 - 15.0	02
Uric Acid, Serum					
Uric Acid, Serum	4.7		mg/dL	3.7 - 8.6	02
Please Note:					02
Therapeutic target for gout patients: <6.0					
GGT	17		IU/L	0 - 65	02
Insulin	6.7		uIU/mL	2.6 - 24.9	02
Ferritin, Serum	159		ng/mL	30 - 400	02
Triiodothyronine, Free, Serum	3.1		pg/mL	2.0 - 4.4	02
Apolipoprotein A-1	158		mg/dL	101 - 178	01
Fatty Acids, Free (Nonester)	0.4		mEq/L	0.1 - 0.6	01
Apolipoprotein B	85		mg/dL	52 - 135	01

01	BN	LabCorp Burlington 1447 York Court, Burlington, NC 27215-3361	Dir: William F Hancock, MD
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02	PDLCA	LabCorp Phoenix 5005 S 40th Street Ste 1200, Phoenix, AZ 85040-2969	Dir: Brian Poirier, MD
03	CLHRT	Cleveland Heartlab Inc 6701 Carnegie Avenue Ste 500, Cleveland, OH 44103-4623	Dir: Deborah Sun, PhD

For inquiries, the physician may contact **Branch: 888-522-2677 Lab: 800-762-4344**

COMPLETE REPORT

PATIENT INFORMATION		SPECIMEN INFORMATION		PRACTITIONER INFORMATION	
Age		Order ID		Name	LABCORP LABCORP
Patient ID	Gender Male	Collection Date/Time 10/20/2017, 10:28 AM		Address	5005 SOUTH 40TH STREET SUITE 1200 PHOENIX, AZ 85040
Fasting Status Fasting	DOB	Received Date/Time 10/24/2017, 12:02 PM			
Ethnicity	BMI	Report Date/Time 10/25/2017, 06:42 AM			

INFLAMMATION							Previous Result	Date
	In Range	Out of Range	Flag**	Relative Risk	Reference Range	Units		
Lp-PLA ₂ Activity ⁽¹⁾	63			LOW	<75	nmol/min/mL		
Based on the documented clinical utility of Lp-PLA ₂ Activity to assess risk of CHD (1), the following cut-off has been defined for Lp-PLA ₂ Activity: A cut-off of ≥ 75 nmol/min/mL defines a population with increased relative risk of developing CHD. (Reference: 1-The Lp-PLA ₂ Studies Collaboration. Lancet. 2010; 375: 1536-1544).								

Comments

(1) This test is performed by a Liquid Chromatography-Tandem Mass Spectrometry (LC/MS/MS) method. This test was developed and its performance characteristics determined by the Cleveland HeartLab, Inc. It has not been cleared or approved by the U.S. FDA. The Cleveland HeartLab is regulated under Clinical Laboratory Improvement Amendments (CLIA) as qualified to perform high-complexity testing. This test is used for clinical purposes. It should not be regarded as investigational or for research.

**Flags: H = Out of Range High; L = Out of Range Low; CH = Critical High; CL = Critical Low