

Pt Name: AARON,JONATHAN, DOB: 11/3/1961, Task Created: 2/2/2013 4:12:41 PM

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Pt. Home Phone: (310)497-7693, Cell: (310)699-5435, Work:

Frances-Call Rabbi Aaron ASAP and tell him to get his butt in here-haven't seen him forever

Task was un-assigned, now assigned to Frances Mendoza-Kaden

Date/Time: 2/2/2013 4:14:29 PM Task Note Edited by Michael F. Richman, MD

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Called patient message to call us and set up appointment

Date/Time: 2/13/2013 4:51:08 PM Task Note Edited by Frances Mendoza-Kaden

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Called patient, message to call us and set up appointment.

Date/Time: 3/20/2013 1:45:57 PM Task Note Edited by Frances Mendoza-Kaden

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Called patient for follow up visit message on cell phone.

Date/Time: 5/22/2013 11:43:49 AM Task Note Edited by Frances Mendoza-Kaden

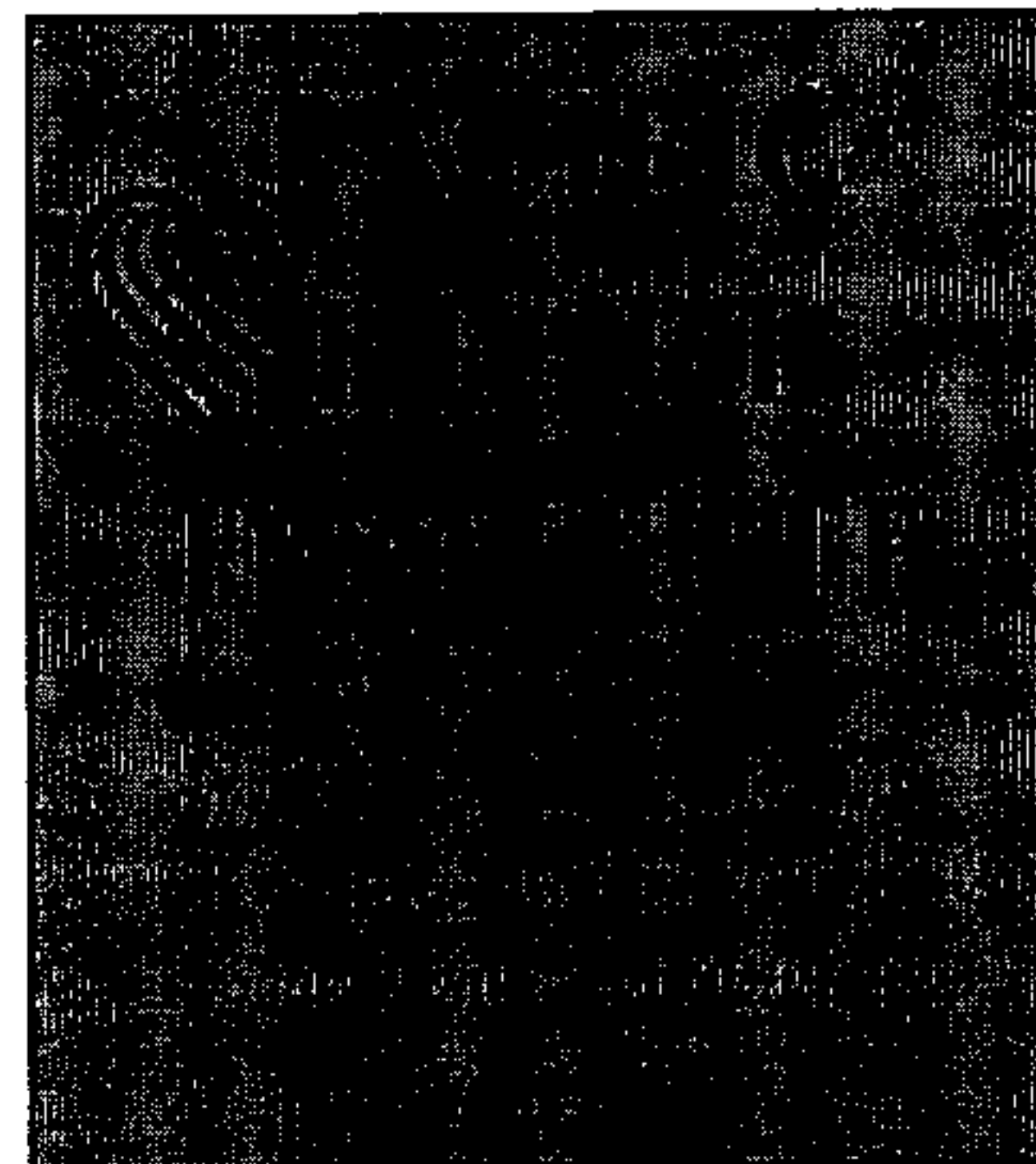
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Date/Time: 8/19/2013 8:51:09 AM Task completed by Frances Mendoza-Kaden

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# TREADMILL STRESS TEST

Patient Name <b>AARON, JONATHAN</b>						Date <b>03/31/11</b>	
Age <b>49</b>	Sex <b>M</b>	Ht. <b>74</b>	In. <b>192</b>	Wt. <b>192</b>	Lbs. <b>192</b>	Referring Physician <b>Richman / Roston, Warren</b>	
Cardiac Medications <b>Crestor, Trilipix, ASA.</b>						Exercise Protocol <b>Bruce</b>	



Lying		
Standing	65	126/73
Hyperventilation		

MIN	SPEED	GRADE	HR	BP	Symptoms / Arrhythmias
1	1.7	10%	84		
2	1.7	10	91		
3	1.7	10	90	132/62	
4	2.5	12	92		
5	2.5	12	101		
6	2.5	12	107	147/76	
7	3.4	14	120		
8	3.4	14	127		
9	3.4	14	133	159/80	
10	4.2	16	149		
11	4.2	16	160		
12	4.2	16			
13	5.0	18			
14	5.0	18			
15	5.0	18			
16	5.5	20			
17	5.5	20			
18	5.5	20			

MIN	HR	BP	Symptoms / Arrhythmias
0	161	159/80	
1	128	146/76	
2	98	196/76	
3	90	183/81	
5			
7			
9			

EXERCISE DURATION <b>10:19</b>	REASON STOPPED <b>fatigue.</b>
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PEAK HR <b>161</b>	PEAK BP <b>159/80</b>	RP PRDCT <b>25,599</b>
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ST SEGMENT RESPONSE <b>Normal</b>
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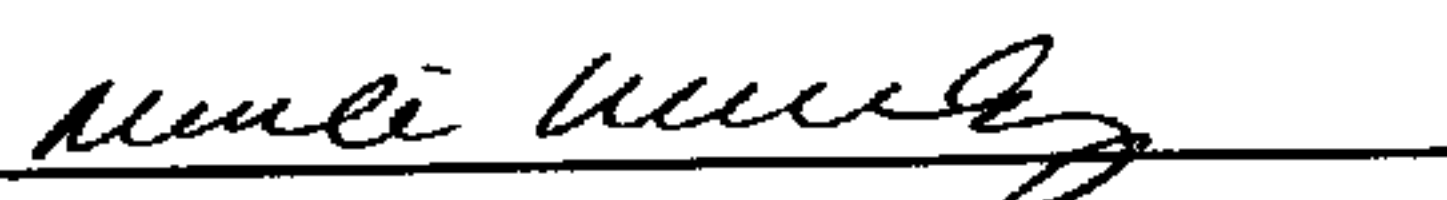
**BASELINE EKG:** Normal sinus rhythm, rate 89, normal resting conduction and repolarization with no arrhythmias.

## INTERPRETATION:

The patient exercised a total of 10:19 achieving a heart rate of 161, which is 94 percent of maximum predicted for age. The test was stopped due to fatigue.

1. There was a normal blood pressure response to stress.
2. The patient experienced no symptoms during stress.
3. Exercise tolerance is excellent.
4. There were no significant arrhythmias.
5. There were no significant ST segment changes.
6. This constitutes a normal electrocardiographic response to stress.

**CONCLUSION:** The overall likelihood of angiographically significant coronary artery disease is low.

  
Nicole M. Weinberg, M.D.

PRESCRIPTION HAS AN ENCRYPTED MICROPRINT™ BACKGROUND - NANOCOPY™

002473

Michael F. Richman, M.D., F.A.C.S.  
Cardiothoracic, General and Vein Surgery  
LIC# G74625 • DEA# BR3315567  
1950 Sawtelle Boulevard, # 150  
Los Angeles, CA 90025  
(310)481-3939 • Fax (310)481-3949



NOTE: SECURITY BACKPRINT • NUMBERING • SAFETY PAPER

Rx

Name

*Jonathan Aaron*

D.O.B.

☐ Female  
☐ Male

Address

Phone

1)

2)

3)

Instant #50041874

Prescription is VOID if the number of drugs prescribed is not noted.

☐ 1 ☐ 2 ☐ 3

TouchSafe®

SP 15

TOUCH OR BREATHE ON TOUCHSAFE® FINGERPRINT TO VALIDATE

Quantity: ☐ 1-24 ☐ 25-49 ☐ 50-74  
☐ 75-100 ☐ 101-150 ☐ 151 and over  
Units \_\_\_\_\_ Refills \_\_\_\_\_ ☐ NR ☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5  
☐ Do not substitute Initial \_\_\_\_\_

Quantity: ☐ 1-24 ☐ 25-49 ☐ 50-74  
☐ 75-100 ☐ 101-150 ☐ 151 and over  
Units \_\_\_\_\_ Refills \_\_\_\_\_ ☐ NR ☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5  
☐ Do not substitute Initial \_\_\_\_\_

Quantity: ☐ 1-24 ☐ 25-49 ☐ 50-74  
☐ 75-100 ☐ 101-150 ☐ 151 and over  
Units \_\_\_\_\_ Refills \_\_\_\_\_ ☐ NR ☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5  
☐ Do not substitute Initial \_\_\_\_\_

VOID APPEARS WHEN COPIED

Date

*2/25/11*

GF92FE



## The Center for Cholesterol Management

PATIENT NAME: Jonathan Aaron

DATE: 6/27/11

DOB: 11/3/1961

### HISTORY OF PRESENT ILLNESS:

<p>Jon comes in today for follow up. He is doing very well and has no <sup>of</sup> what's over. We discussed his lipid panel. 1 CK level</p>
---

### PMHx:

<p>① Met syndrome</p>
<p>② MH</p>

### METABOLIC SYNDROME CRITERION:

(Presence of 3 or more criterion define Metabolic Syndrome)

1. Waist > 40"/35" (M/F)  
Waist > 35"/32" (M/F Asian)
2. HDL-C < 40/50mg/dL (M/F)\*
3. TG > 150 mg/dL\*
4. FBS > 100 mg/dL or DM
5. BP > 130/ > 85\* (\* or on medication)

### LIPOPROFILE RESULTS:

<p>LDL-P: 1029</p>
<p>CK: 1196</p>
<p>6/13/11</p>

MEDICATION:

NAME: Jonathan Aaron

atorvastatin 10mg qd  
Zetia 10mg qd  
Lopraz 50mg qd  
Vit. D 50,000 IU a week  
Benecol chews qd  
Glucophage 1000mg qd  
Baby Aspirin 81mg qd  
trilipix 135mg qd

PHYSICAL EXAM:

BP: 112/79

P: 72

RR: 16

T: 98

WEIGHT:

WAIST CIRCUMFERENCE:

GENERAL: WNL WNL ♂ in NAD

CHEST:

clear

HEART:

PPP

ABDOMEN:

soft, NT

EXTREMITIES:

warm.

ASSESSMENT:

49 y.o. W ♂ c 1/2 M H and metabolic syndrome but ↑ CK c stable and optimal LAb P: TG.

PLAN:

① cont meds.

② CK in 1 month







# The Center for Cholesterol Management

PATIENT NAME: Jonathan Aaron

DATE: 4-22-11

DOB: 11-3-1961

## HISTORY OF PRESENT ILLNESS:

Jonathan comes in today for follow up  
He is doing well and is taking his med.  
He feels well. we discussed his lipid  
treadmill stress <sup>length</sup>  
3/31/11 → low risk for CVD

## PMHX:

① Met. syndrome  
② M/H

## METABOLIC SYNDROME CRITERION:

(Presence of 3 or more criterion define Metabolic Syndrome)

1. Waist > 40"/35" (M/F)  
Waist > 35"/32" (M/F Asian)
2. HDL-C < 40/50mg/dL (M/F)\*
3. TG > 150 mg/dL\*
4. FBS > 100 mg/dL or DM
5. BP > 130/ > 85\* (\* or on medication)

## LIPOPROFILE RESULTS:

LDL-C - 1390	TG - 189	} 4/27/11
HDL-B - 71	HDL-A/C - 5.2	
HDL-C - 33	met D - 28	

NAME: Jonathan Aaron

Crestor 10mg qd  
trilipix 135 qd  
lovaza 4 qd  
Vit D 5000 IU qd  
Glucophage 1000 mg qd

BP 111 / 74

P 68

RR 16

T 98

GENERAL: *WCAW & on NED*

CHEST: Clear

HEART: *H. P. H.*

ABDOMEN: *empty*

EXTREMITIES: *same*

**ASSESSMENT:**

4960 W02 @ 9/10 MH: 111 TGS;  
11 LDL-P now improving

PLAN:

- ① cont needs
- ② add benefit chgs 29¢
- ③ then 10mg
- ④ ↑ to 5mg
- ⑤ 50,000 a week
- ⑥ 1/2 in 2 months

<sup>2</sup>  
**Center for Cholesterol Management** A Medical Corporation

NAME Jonathan Aaron

DATE

2-25-11

DOB 11-3-61

HISTORY OF PRESENT ILLNESS

Jon comes in today for follow up. He  
ran out of meds 2 weeks ago  
we discussed length his current risk;  
tx plan

PMHx

- ① MH
- ② met syndrome

Metabolic Syndrome Criteria  
(Presence of 3 or more criteria define Metabolic Syndrome)

- 1. Waist > 40"/35" (M/F)  
Waist > 35"/32" (M/F Asian)
- 2. HDL-C < 40mg/dL (M/F)\*
- 3. TG > 100 mg/dL\*
- 4. FBS > 100 mg/dL or DM
- 5. BP > 130 / > 85\* (\* or on medication)

Laboratory Results

LDL-C = 1164 on 11/9/10  
F6 - 304  
HbA1c = 29



MEDS

Crestor 10mg qd  
lovaza 400mg qd  
trilipix 135mg qd  
Vit-D 4000 IU qd  
Glucophage 1000mg qd

NAME  
Jonathan  
Aaron

PHYSICAL EXAM

BP 130/79 P 72 RR 16 T 98

GENERAL W/W/N 0? in NAD

CHEST clear

HEART

RR

ABDOMEN

soft, NT

EXTREMITIES

warm

LABS - 49 y/o W 0? 4% M.H.  
met. syndrome  
non-compliant i med

PLAN

*[Signature]*

- ① treadmill stress test
- ② H/L follow up panel
- ③ F/U 2 weeks

The  
**Center for Cholesterol Management** A Medical Corporation

NAME Jonathan Aaron

DATE

DOB

11/03/1961

11/9/10

HISTORY OF PRESENT ILLNESS:

Jon comes in today for follow up. He is doing well but has not been taking his O3A due to nausea when taking

PMHx:

- ① MI
- ② Met. Syndrome

Metabolic Syndrome Criterion

(Presence of 3 or more criterion define Metabolic Syndrome)

- 1. Waist > 40"/35" (M/F)  
Waist > 35"/32" (M/F Asian)
- 2. HDL-C < 40/50mg/dL (M/F)\*
- 3. TG > 150 mg/dL\*
- 4. FBS > 100 mg/dL or DM
- 5. BP > 130 / > 85\* (\* or on medication)

Lipoprofile Results

LDL-P: 1764      unruhi 57  
APOB - 119  
TG - 571  
VLDL-P = 12

8/24/10

MEDS

NAME

Crestor 10mg qd

adam

trilipix 135mg qd

Lovaza 4gams (not taking)

Vit. D 4000 IU qd

PHYSICAL EXAM

Glucophage XR ~~1000~~ mg qd

BP 128/80 P 75 RR 16 T 98

GENERAL: WDNW 8' in NA /

CHEST: clear

HEART: RRR

ABDOMEN: soft, NT

EXTREMITIES warm.

Assessment

49 y.o. w/ 4/0 M/H  
; met. syndrome; ↓ vit D

PLAN:

V

- ① HDL follow up panel
- ② FU 2 weeks

The

# Center for Cholesterol Management

A Medical Corporation

NAME Jonathan Aaron

DATE

9/14/10

DOB 11/03/1961

HISTORY OF PRESENT ILLNESS:

*Jonathan comes in today for follow up  
we discussed his lipid panel & length*

PMHX:

*① ↑TG*

Metabolic Syndrome Criterion

(Presence of 3 or more criterion define Metabolic Syndrome)

1. Waist > 40"/35" (M/F)  
Waist > 35"/32" (M/F Asian)
2. HDL-C < 40/50mg/dL (M/F)\*
3. TG > 150 mg/dL\*
4. FBS > 100 mg/dL or DM
5. BP > 130 / > 85\* (\* or on medication)

Lipoprofile Results

*see results from 8/24/10*

MEDS

NAME

Adam

Ø

PHYSICAL EXAM

BP 118/70 P 72 RR 16 T 98.6

GENERAL: WDMW ♂ in WAD

CHEST: clear

HEART: RRR

ABDOMEN: soft, NT

EXTREMITIES WNM

Assessment

48 y o w ♂ with paroxysmal combined  
hypertension & IR.

PLAN:

- ① Bag ASA<sub>II</sub>
- ② Uterolone 10mg qd
- ③ tramadol
- ④ Lorazepam 4mg qd
- ⑤ streptococcal
- ⑥ Furosemide 125mg qd
- ⑦ Multiple consults
- ⑧ Lisinopril 20mg 1000mg qd
- ⑨ HbA<sub>1c</sub> follow up
- ⑩ vit D 4000 u qd



The

# Center for Cholesterol Management

A Medical Corporation

NAME Jonathan Aaron

DATE

DOB 11/03/1961

8/24/10

CHIEF COMPLAINT: -

48yo w o<sup>2</sup> c 4/0 ↑TGS presents  
for lipid evaluation

HISTORY OF PRESENT ILLNESS:

pt has a had a fluttering feeling in his  
heart but denies chest pressure, denies  
SOB.  $\phi$  MI  $\phi$  CVA. pt does not exercise  
never had a stress test.

PMHx:

- ① osteoarthritis of ② spine  
4/0 numbness ② leg post thigh  
partially.
- ② ↑TGS

PSHx:

- ① 3x microdissection - L<sub>3</sub>-S<sub>1</sub>,

NAME

aaron

MEDS:

Ø

ALLERGIES: NKA

SOCIAL HISTORY: - ⑦ tobacco x 28 yrs.

FAMILY HISTORY: -

- ① cancer
- ② MI

REVIEW OF SYSTEMS: -

see intake form

NAME

norm

PHYSICAL EXAM

BP 112/70 P 72 RR 16 T. 98

GENERAL: WDNW ♂ in NAD

HEENT: NL/AT

NECK: φ/mult

CHEST: clear

HEART: MMR

ABDOMEN: soft, NT

BACK:

EXTREMITIES: WNM

NEURO:

Assessment - 48 y o w o ♂ c H/o ↑ T6s  
72000.

PLAN:

- ① HDL comprehensive panel
- ② F/U 2 weeks

Lu

<b>Patient</b>	Name:	Jonathan Aaron	Phone #:		Patient ID #:	10-237-0263
	Fasting Status:	Non-Fasting	Gender:	Male	Birthdate:	11/3/1961
	Height:		Weight:		BMI:	49
<b>Specimen</b>	Collection Time:	1:50 pm	Specimen ID:	11061400670	Collection Date:	6/13/2011
	Report Type:	Complete	Received Date:	6/14/2011	Report Date:	6/16/2011
<b>Provider</b>	Requesting Provider:					
	Michael Richman The Center for Cholesterol Management 1950 Sawtelle Blvd #150 Los Angeles, CA 90025					
Client ID:						06-90025-18-0000383

Laboratory Test	Notes	High Risk	Intermediate Risk	Optimal	High Risk Range	Intermediate Risk Range	Optimal Range	Previous Results 4/7/2011
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Lipids	Total Cholesterol (mg/dL)			102	≥ 240	200 - 239	< 200	126
	LDL-C Direct (mg/dL)			49	≥ 130 CHD & CHD risk eq. > 100	100 - 129 CHD & CHD risk eq. 70 - 100	< 100 CHD & CHD risk eq. < 70	63
	HDL-C (mg/dL)	38			< 40		≥ 40	33
	Triglycerides (mg/dL)			109	≥ 200	150 - 199	< 150	189
	Non-HDL-C (mg/dL) (calculated)			64	≥ 160	130 - 159	< 130	93

Lipoprotein Particles and Apolipoproteins	Apo B (mg/dL)			55	≥ 80	60 - 79	< 60	71
	LDL-P (nmol/L)		1029		≥ 1300	1000 - 1299	< 1000	1390
	sdLDL (mg/dL)*			14	≥ 31	21 - 30	≤ 20	22
	% sdLDL (calculated)	28			≥ 23	14 - 22	≤ 13	35
	Apo A-I (mg/dL)		124		< 114	114 - 131	≥ 132	117
	HDL-P (μmol/L)		31.8		< 28.0	28.0 - 34.0	≥ 35.0	29.8
	HDL2 (mg/dL)*	8			≤ 8	9 - 11	≥ 12	6
	Apo B:Apo A-I Ratio (calculated)			0.45	≥ 0.81	0.61 - 0.81	≤ 0.6	0.61
	Lp(a) Mass (mg/dL)			3	≥ 30		< 30	5
	Lp(a) Cholesterol (mg/dL)				≥ 6	3 - 5	< 3	

Inflammation/ Oxidation	Myeloperoxidase (pmol/L)				≥ 550	400 - 549	< 400	
	Lp-PLA <sub>2</sub> (ng/mL)			143	> 235	200 - 235	< 200	126
	hs-CRP (mg/L)		1.24		≥ 3.0	1.0 - 2.9	< 1.0	0.66
	Fibrinogen (mg/dL)			334	≥ 465	391 - 464	≤ 390	340

Myocardial Stress	NT-proBNP (pg/mL)			62	≥ 450	125 - 449	< 125	15

Platelets	AspirinWorks® (urine) (pg/mg of creatinine)				> 1500		≤ 1500	

Lab Notes:

Provider Notes:

*optimal LDL-P*  
*T6*  
*T↑ CK*  
*repeat CK*  
*repeat HDL follow up*  
*6/27/11*

Patient	Name:	Jonathan Aaron	Phone #:		Patient ID #:	10-237-0263		
	Fasting Status:	Non-Fasting	Gender:	Male	Birthdate:	11/3/1961	Age:	49
	Height:		Weight:		BMI:		Prev. BMI:	
Specimen	Collection Time:	1:50 pm	Specimen ID:	11061400670	Collection Date:	6/13/2011	Report Type:	Complete
	Received Date:	6/14/2011	Report Date:	6/16/2011				
Provider	Requesting Provider:				Michael Richman			
	The Center for Cholesterol Management				1950 Sawtelle Blvd #150 Los Angeles, CA 90025			
				Client ID:				06-90025-18-0000383

Laboratory Test		Notes	High Risk	Intermediate Risk	Optimal	High Risk Range	Intermediate Risk Range	Optimal Range	Previous Results 4/7/2011
Lipoprotein Genetics	Apolipoprotein E Genotype*					Estimated Genotype Frequency: 2/2 (~1-2%), 2/3 (~15%), 2/4 (~1-2%), 3/3 (~55%), 3/4 (~25%), 4/4 (~1-2%)			3/3
Platelet Genetics	CYP2C19*2*					*1/*1 = optimal, *1/*2 = intermediate, *2/*2 = poor			*1/*1
	CYP2C19*3*					*1/*1 = optimal, *1/*3 = intermediate, *3/*3 = poor			*1/*1
	CYP2C19*17*					*1/*1 = optimal, *1/*17 = rapid, *17/*17 = ultra rapid			Not Tested
Coagulation Genetics	Factor V Leiden					Optimal=Non-carrier (Arg/Arg); At Risk=(Arg/Gln or Gln/Gln)			Arg/Arg
	Prothrombin Mutation					Optimal=Non-carrier (G/G); At Risk=(G/A or A/A)			G/G
Metabolic	Insulin (µU/mL)				8	≥ 12	10 - 11	3 - 9	10
	Free Fatty Acid (mmol/L)			0.66		> 0.7	0.6 - 0.7	≤ 0.59	0.27
	Glucose (mg/dL)				94	≤ 55 or > 125	56-69 or 100-125	70 - 99	84
	HbA1c (%)				5.3	≥ 6.5	5.7 - 6.4	≤ 5.6	5.2
	Estimated Average Glucose (mg/dL) (calculated)				105.9	≥ 139.9	116.9 - 139.8	≤ 116.8	102.5
	25-hydroxy-Vitamin D (ng/mL)				41	≤ 14	15 - 29	30 - 100	28
	Homocysteine (µmol/L)		14			> 13	11 - 13	≤ 10	13

Health Coach, please call 1-877-4HDLABS (1-877-443-5227) or visit us online at www.myhdl.com

Lab Notes:



<b>Provider</b>	<b>Requesting Provider:</b> <b>Michael Richman</b> The Center for Cholesterol Management 1950 Sawtelle Blvd #150 Los Angeles, CA 90025
	<b>Client ID:</b> <b>06-90025-18-0000383</b>

Albumin (g/dl)	4.9		3.5 - 5.2
ALP (U/L)	38	L	40 - 129
ALT / GPT (U/L)	32		Up to 41
AST / GOT (U/L)	46	H	Up to 40
CK (U/L)	1196	H	39 - 308
Direct Bilirubin (mg/dL)	0.2		0.1 - 0.3
Total Bilirubin (mg/dL)	0.5		Up to 1.2
Total Protein (g/dL)	6.6		6.4 - 8.3

**Lab Notes:**

\*This test was developed and its performance characteristics determined by HDL, Inc. It has not been cleared or approved by the U.S. Food & Drug Administration (FDA). The FDA has determined that such clearance or approval is not necessary. This test is used for clinical purposes. It should not be regarded as investigational or for research. This laboratory is certified under CLIA-88 as qualified to perform high complexity clinical laboratory testing.

Patient

Name:

Jonathan Aaron

Phone #:

Patient ID #:

10-237-0263

Fasting Status:

Non-Fasting

Gender:

Male

Birthdate:

11/3/1961

Age:

49

Height:

Weight:

BMI:

Prev. BMI:

Specimen

Collection Time:

1:50 pm

Specimen ID:

11061400670

Collection Date:

6/13/2011

Report Type:

Complete

Received Date:

6/14/2011

Report Date:

6/16/2011

Provider

Requesting Provider:

Michael Richman

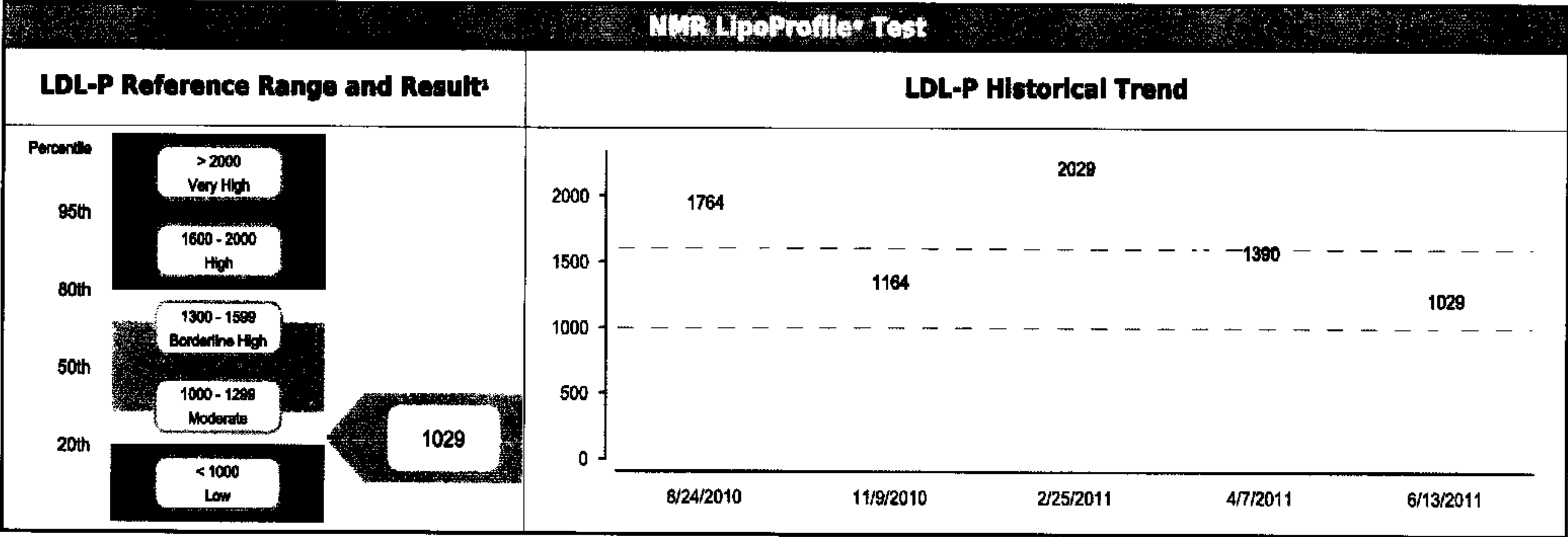
The Center for Cholesterol Management

1950 Sawtelle Blvd #150

Los Angeles, CA 90025

Client ID:

06-90025-18-0000383



Particle Concentration and Size								
Laboratory Test		Result	Percentile in Reference Population <sup>2</sup>					Previous Results
			<div>Higher CVD Risk<div></div>Lower CVD Risk<div></div></div>					
HDL Particles	HDL-P (total)	31.8	low	25th (26.7)	50th (30.5)	75th (34.9)	high	29.8
	μmol/L		<div><div></div><div></div><div>31.8</div><div></div><div></div></div>					
Small LDL-P and LDL Size are associated with CVD risk, but not after LDL-P is taken into account.								
			<div>Insulin Resistance<div></div>Insulin Sensitive<div></div></div>					
Lipoprotein Markers Associated with Insulin Resistance and Diabetes Risk <sup>3</sup>	LARGE VLDL-P	< 0.7	high	75th (6.9)	50th (2.7)	25th (0.9)	low	6.6
	nmol/L		<div><div></div><div></div><div>0.7</div><div></div><div></div></div>					
	SMALL LDL-P	937	high	75th (839)	50th (527)	25th (117)	low	668
	nmol/L		<div><div></div><div>937</div><div></div><div></div><div></div></div>					
	LARGE HDL-P	< 0.7	low	25th (3.1)	50th (4.8)	75th (7.3)	high	< 0.7
	μmol/L		<div><div>0.7</div><div></div><div></div><div></div><div></div></div>					
	VLDL SIZE	42.0	large	75th (52.5)	50th (46.6)	25th (42.4)	small	48.2
nm		<div><div></div><div></div><div>42.0</div><div></div><div></div></div>						
LDL SIZE	19.8	small	25th (20.4)	50th (20.8)	75th (21.2)	large	20.4	
nm		<div><div></div><div>19.8</div><div></div><div></div><div></div></div>						
HDL SIZE	8.4	small	25th (8.9)	50th (9.2)	75th (9.6)	large	< 8.3	
nm		<div><div>8.4</div><div></div><div></div><div></div><div></div></div>						
LP-IR SCORE*	46	insulin resistant	75th (63)	50th (46)	25th (27)	insulin sensitive	73	
0 - 100		<div><div></div><div></div><div>46</div><div></div><div></div></div>						

LP-IR Score is inaccurate if a patient is non-fasting.  
\* The LP-IR Score combines the information from the 6 markers above it to give improved assessment of insulin resistance and diabetes risk.  
These laboratory assays, validated by LipoScience, have not been cleared by the US Food and Drug Administration. The clinical utility of these laboratory values has not been fully established.  
1. Reference population comprises 5,362 men and women not on lipid medication enrolled in the Multi-Ethnic Study of Atherosclerosis (MESA). Mora, et al. Atherosclerosis 2007.  
2. LipoScience reference population comprises 4,588 men and women without known CVD or diabetes and not on lipid medication.  
3. Garvey WT, et al. Diabetes. 2003; 53:453-462. 4. Goff DC et al. Metabolism. 2005; 54:264-270.

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Patient	Name:		Phone #:		Patient ID #:	
	Jonathan Aaron				10-237-0263	
	Fasting Status:		Gender:		Birthdate:	
	Non-Fasting		Male		11/3/1961	
	Height:		Weight:		BMI:	
					Prev. BMI:	

Specimen	Collection Time:		Specimen ID:	
	1:50 pm		11061400670	
	Collection Date:		Report Type:	
	6/13/2011		Complete	
	Received Date:		Report Date:	
	6/14/2011		6/16/2011	

Provider	Requesting Provider:	
	Michael Richman	
	The Center for Cholesterol Management	
	1950 Sawtelle Blvd #150	
	Los Angeles, CA 90025	
	Client ID:	
	06-90025-18-0000383	

Comments:

<p>HDL cholesterol, HDL 2 (large HDL particle) cholesterol, and apolipoprotein A-I are decreased or in the intermediate range. Low HDL cholesterol and low apolipoprotein A-I have been associated with increased risk for cardiovascular disease and events. Low amounts of large HDL particles suggest that the HDL present may not be in its most protective form. HDL cholesterol, large HDL particle, and apolipoprotein A-I concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have been demonstrated to increase large HDL particle concentration.</p>
<p>%SDLDL is elevated in this sample consistent with the presence of small dense LDL particles in this sample. The clinical significance of this, in the setting of optimal values of LDL cholesterol and Apo B is not well characterized. Niacin, fibric acids, and combination therapy (statin + niacin) have been shown to increase LDL particle size.</p>
<p>Although the LDL cholesterol concentration is optimal, LDL particle concentration is borderline high in this sample. Studies have shown that elevated LDL particle concentration is associated with increased risk for coronary heart disease, even in the presence of optimal LDL cholesterol values. Small LDL particles may be observed in association with the metabolic syndrome and pre-diabetes. Statins effectively reduce the number of LDL particles, but do not generally influence the size distribution of the LDL particles. Niacin, fibrates, and combination therapy (statin +niacin) have been shown to increase LDL particle size.</p>
<p>C-reactive protein is in the intermediate range. CRP is an acute phase reactant. Data from prospective studies indicates that increased concentration of CRP is associated with an increased risk for the development of ischemic cardiovascular events. Consider repeat analysis of CRP in 2-4 weeks to establish baseline value. If CRP remains elevated, then lifestyle changes, including weight reduction, low-fat diet, smoking cessation and regular exercise, should be the initial approach. A diet rich in plant sterols, soy protein, viscous fiber, and almonds has been shown to have CRP-lowering effects comparable to that of lovastatin 20 mg/day. Medications that may lower CRP include statins, fibrates, aspirin, and fish oil. Reducing global CHD risk by aggressive treatment of the traditional risk factors by established therapies may also be beneficial.</p>
<p>Increased homocysteine. Most, but not all prospective studies of homocysteine and cardiovascular risk show homocysteine to be associated with cardiovascular events. Levels &gt;13 umol/L are considered elevated. Such increases in homocysteine levels can occur with aging, menopause, hypothyroidism, low plasma levels of vitamin cofactors (B6, B12 and folate), certain drugs, and chronic renal insufficiency. Genetic variation in enzymes involved in homocysteine metabolism contributes to inter-individual differences in plasma homocysteine levels.</p>
<p>Non-Esterified "Free" Fatty Acid concentration is in the intermediate range. Elevated free fatty acids have been associated with the metabolic syndrome and increased risk for the development of type 2 diabetes.</p>
<p>Total HDL particle concentration is in the intermediate range in this sample. Decreased HDL particles have been associated with increased risk for cardiovascular disease. HDL particle concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have been demonstrated to increase HDL particle concentration.</p>

End of Report

ATTN PATIENT: Please contact HDL, Inc. at 1-877-4HDLABS (1-877-443-5227) to set an appointment with your personal health coach to discuss your diet and exercise needs at no charge. You can also visit us online at [www.myhdl.com](http://www.myhdl.com) and schedule an appointment through our web portal.



INSTRUCTIONS

- 1) Please fill in all of the highlighted areas.
- 2) Have patient sign Release and Assignment of Benefits below.
- 3) Copy BOTH sides of patient's insurance card(s).
- 4) Indicate primary and secondary diagnoses below.

NOTE: Physicians (or other individuals authorized to order tests) should only order tests that are medically necessary and reasonable.

PATIENT INFORMATION

Form fields for Patient Information including Last Name, First, Middle Initial, Client Patient ID #, Address, City, State, Zip Code, Home Phone, Sex, Date of Birth, Work Phone, Social Security #, Height, and Weight.

SPECIMEN INFORMATION

Form fields for Specimen Information including Drawing Lab, Phone, Collection Date, Time, Fasting status, and Initials.

HDL, INC. USE ONLY

Form fields for HDL, Inc. Use Only including Received Date, Time, Initials, and various tube counts.

BILLING INFORMATION

Billing Information section with checkboxes for Insurance, Medicare, Check, and Credit Card, along with fields for Name, Total Amount, and Card Issuer.

RELEASE AND ASSIGNMENT OF BENEFITS

As a courtesy, HDL, Inc. will make every reasonable effort to obtain reimbursement for ordered tests. I authorize HDL, Inc. to release to Medicare or its Carriers & any insurance carrier providing medical benefits to me and any health plan of which I am a member...

Bill to my insurance: I understand that if my insurance company pays me directly for services rendered by HDL, Inc., I am responsible for forwarding such payment to HDL, Inc. I also understand, I am responsible for any deductibles/copayment, as required by my plan.

Important: Insurance regulations require HDL, Inc. to seek payment. I permit a copy of this authorization to be in place of the original.

\*Patient Signature

Date

COMMENTS

REQUESTING PROVIDER/LAB/INSTITUTION

Dr. Richman
The Center for Cholesterol Management
1950 Sawtelle Blvd #150
Los Angeles, CA 90025
Client ID: 06-90025-18-0000383
Phone: (310) 481-3839
Fax: (310) 481-3949

\*Physician or Authorized Signature

Date

CUSTOM PANELS

- Baseline Assessment
Lipid Panel
Apo A1
Apo B
LDL P
sdLDL
HDL 2 subclass
Lp(a) mass w/ reflex
Apo E Genotype
Factor V Leiden
Prothrombin Mutation
CYP2C19
Lp-PLA2
CRP-hs
Fibrinogen
Homocysteine
FFA (NEFA)
Insulin
NT pro BNP
Vitamin D
HEP
HbA1C

INDIVIDUAL TESTS

Individual Tests section with checkboxes for Routine Panels, Lipoprotein Particles, Genetic Assays, Inflammation & Platelets, Metabolic, Hormones, Renal, and Thyroid Function.

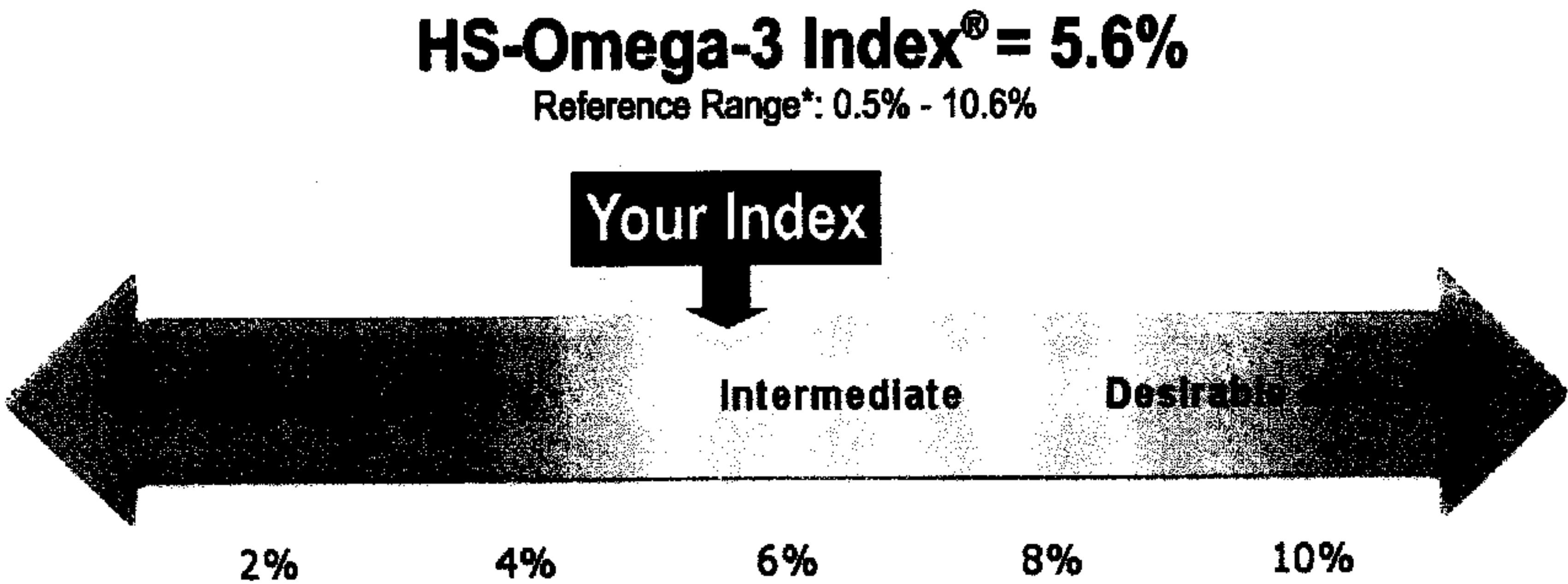
DIAGNOSIS (ICD-9 CODE)

NOTE: The below ICD-9 codes are listed as a convenience. Ordering physicians should report the code that best describes the reason for ordering the tests & are not required to use the codes provided below.

Diagnosis section with checkboxes for Diabetes, Pure Hypercholesterolemia, Pure Hyperglyceridemia, Mixed Hyperlipidemia, Unspecified Hyperlipidemia, Hypertension, Hypertensive Heart Disease, Hypertension w/Renal Disease, Hypertension w/Renal Disease & Heart Failure, Intermediate Coronary Syndrome, Coronary Atherosclerosis Unspecified Vessel, Native or Graft, Congestive Heart Failure, Cirrhosis of liver without mention of alcohol, Other chronic non-alcoholic liver disease, Unspecified chronic liver disease without mention of alcohol, and OTHER.

Patient Name: Aaron, Jonathan  
Date of Birth: 11/3/1961  
Specimen ID: 10-237-0263\_11040800203  
Practice: The Center for Cholesterol Management

Date of Collection: 04/07/2011  
Date Reported: 04/11/2011  
Provider: Michael Richman



Dried Blood Spot Fatty Acid Profile

Omega-3 Fatty Acids			cis-Monounsaturated Fatty Acids		
Alpha-Linolenic	(18:3n3)	0.3 %	Palmitoleic	(16:1n7)	0.3 %
Eicosapentaenoic	(EPA, 20:5n3)	1.2 %	Oleic	(18:1n9)	18.9 %
Docosapentaenoic-n3	(22:5n3)	1.6 %	Eicosenoic	(20:1n9)	0.1 %
Docosahexaenoic	(DHA, 22:6n3)	3.9 %	Nervonic	(24:1n9)	0.3 %
Range*: 0.0% - 12.8%		Total: 7.0 %	Range*: 11.6% - 30.3%		Total: 19.6 %
Omega-6 Fatty Acids			Saturated Fatty Acids		
Linoleic	(18:2n6)	15.7 %	Myristic	(14:0)	0.7 %
Gamma-Linolenic	(18:3n6)	0.3 %	Palmitic	(16:0)	22.5 %
Eicosadienoic	(20:2n6)	0.4 %	Stearic	(18:0)	15.1 %
Dihomo-γ-linolenic	(20:3n6)	1.7 %	Arachidic	(20:0)	0.1 %
Arachidonic	(AA, 20:4n6)	13.0 %	Behenic	(22:0)	0.2 %
Docosatetraenoic	(22:4n6)	2.1 %	Lignoceric	(24:0)	0.2 %
Docosapentaenoic-n6	(22:5n6)	0.4 %	Range*: 26.0% - 38.5%		Total: 38.8 %
Range*: 26.1% - 51.2%		Total: 33.6 %	Trans Fatty Acids		
Fatty Acids Ratios			Trans Palmitoleic	(16:1n7t)	0.1 %
Omega-6:Omega-3	(0.0 – 14.9)*	4.9	Trans Oleic	(18:1t)	0.7 %
AA:EPA	(0.0 – 59.1)*	10.8	Trans Linoleic	(18:2n6t)	0.4 %
			Range*: 0.0% - 4.8%		Total: 1.2 %

\* Reference range is derived from 992 subjects (mean age 67 years).



<b>Patient</b>	Name:		Phone #:		Patient ID #:	
	Jonathan Aaron				10-237-0263	
	Fasting Status:		Gender:		Birthdate:	
	Non-Fasting		Male		11/3/1961	
	Height:		Weight:		BMI:	

<b>Specimen</b>	Collection Time:		Specimen ID:	
	11:30 am		11040800203	
	Collection Date:		Report Type:	
	4/7/2011		Complete	
	Received Date:		Report Date:	
	4/8/2011		4/11/2011	

<b>Provider</b>	Requesting Provider:	
	Michael Richman	
	The Center for Cholesterol Management	
	1950 Sawtelle Blvd #150	
	Los Angeles, CA 90025	
	Client ID:	
	06-90025-18-0000383	


Laboratory Test	Notes	High Risk	Intermediate Risk	Optimal	High Risk Range	Intermediate Risk Range	Optimal Range	Previous Results 2/25/2011
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<b>Lipids</b>	Total Cholesterol (mg/dL)				126	≥ 240	200 - 239	< 200	241
	LDL-C Direct (mg/dL)				63	≥ 130 CHD & CHD risk eq. > 100	100 - 129 CHD & CHD risk eq. 70 - 100	< 100 CHD & CHD risk eq. < 70	99
	HDL-C (mg/dL)		33			< 40		≥ 40	39
	Triglycerides (mg/dL)			189		≥ 200	150 - 199	< 150	494
	Non-HDL-C (mg/dL) (calculated)				93	≥ 160	130 - 159	< 130	202

Lipoprotein Particles and Apolipoproteins	Apo B (mg/dL)			71		≥ 80	60 - 79	< 60	
	LDL-P (nmol/L)		1390			≥ 1300	1000 - 1299	< 1000	2029
	sdLDL (mg/dL)*			22		≥ 31	21 - 30	≤ 20	
	% sdLDL (calculated)		35			≥ 23	14 - 22	≤ 13	
	Apo A-I (mg/dL)			117		< 114	114 - 131	≥ 132	
	HDL-P (μmol/L)			29.8		< 28.0	28.0 - 34.0	≥ 35.0	24.9
	HDL2 (mg/dL)*		6			≤ 8	9 - 11	≥ 12	
	Apo B:Apo A-I Ratio (calculated)			0.61		≥ 0.81	0.61 - 0.81	≤ 0.6	
	Lp(a) Mass (mg/dL)				5	≥ 30		< 30	
	Lp(a) Cholesterol (mg/dL)					≥ 6	3 - 5	< 3	

<b>Inflammation/ Oxidation</b>	Myeloperoxidase (pmol/L)					≥ 550	400 - 549	< 400	
	Lp-PLA <sub>2</sub> (ng/mL)				126	> 235	200 - 235	< 200	
	hs-CRP (mg/L)				0.66	≥ 3.0	1.0 - 2.9	< 1.0	
	Fibrinogen (mg/dL)				340	≥ 465	391 - 464	≤ 390	

Myocardial Stress	NT-proBNP (pg/mL)				15	$\geq 450$	125 - 449	$< 125$	

<b>Platelets</b>	AspirinWorks® (urine) (pg/mg of creatinine)				> 1500		≤ 1500	
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**Lab Notes:****Provider Notes:**

Dr. Joseph P. McConnell | Laboratory Director | CLIA No. 49D1100708 | CAP No. 7224971 | NPI No. 1629209853

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HDL 20.0

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<b>Provider</b>	Requesting Provider: <b>Michael Richman</b> The Center for Cholesterol Management 1950 Sawtelle Blvd #150 Los Angeles, CA 90025
	Client ID: <b>06-90025-18-0000383</b>

<b>Metabolic</b>	Insulin ( $\mu\text{U/mL}$ )			10		$\geq 12$	10 - 11	3 - 9	
	Free Fatty Acid (mmol/L)				0.27	$> 0.7$	0.6 - 0.7	$\leq 0.59$	
	Glucose (mg/dL)				84	$\leq 55$ or $> 125$	56-69 or 100-125	70 - 99	
	HbA1c (%)				5.2	$\geq 6.5$	6.0 - 6.4	4.8 - 5.9	
	25-hydroxy-Vitamin D (ng/mL)			28		$\leq 14$	15 - 29	30 - 100	
	Homocysteine ( $\mu\text{mol/L}$ )			13		$> 13$	11 - 13	$\leq 10$	
	Vitamin B <sub>12</sub> (pg/mL)				639	$< 211$	211 - 299	$\geq 300$	

**Lab Notes:**

\*This test was developed and its performance characteristics determined by HDL, Inc. It has not been cleared or approved by the U.S. Food & Drug Administration (FDA). The FDA has determined that such clearance or approval is not necessary. This test is used for clinical purposes. It should not be regarded as investigational or for research. This laboratory is certified under CLIA-88 as qualified to perform high complexity clinical laboratory testing.

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HDL 20.0

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<b>Patient</b>	<b>Name:</b>		<b>Phone #:</b>		<b>Patient ID #:</b>	
	Jonathan Aaron				10-237-0263	
	<b>Fasting Status:</b>		<b>Gender:</b>		<b>Birthdate:</b>	
	Non-Fasting		Male		11/3/1961	
	<b>Height:</b>	<b>Weight:</b>	<b>BMI:</b>	<b>Prev. BMI:</b>		

<b>Specimen</b>	Collection Time:	Specimen ID:
	11:30 am	11040800203
	Collection Date:	Report Type:
	4/7/2011	Complete
	Received Date:	Report Date:
	4/8/2011	4/11/2011

<b>Provider</b>	<b>Requesting Provider:</b> <b>Michael Richman</b> The Center for Cholesterol Management 1950 Sawtelle Blvd #150 Los Angeles, CA 90025
	<b>Client ID:</b> <b>06-90025-18-0000383</b>

Other Biomarkers	Result	Flag	Reference Interval
Albumin (g/dl)	4.5		3.5 - 5.2
ALP (U/L)	45		40 - 129
ALT / GPT (U/L)	28		Up to 41
AST / GOT (U/L)	21		Up to 40
Direct Bilirubin (mg/dL)	0.1		0.0 - 0.3
Total Bilirubin (mg/dL)	0.3		Up to 1.2
Total Protein (g/dL)	6.4		6.4 - 8.3

**Lab Notes:**

\*This test was developed and its performance characteristics determined by HDL, Inc. It has not been cleared or approved by the U.S. Food & Drug Administration (FDA). The FDA has determined that such clearance or approval is not necessary. This test is used for clinical purposes. It should not be regarded as investigational or for research. This laboratory is certified under CLIA-88 as qualified to perform high complexity clinical laboratory testing.

**Dr. Joseph P. McConnell | Laboratory Director | CLIA No. 49D1100708 | CAP No. 7224971 | NPI No. 1629209853**

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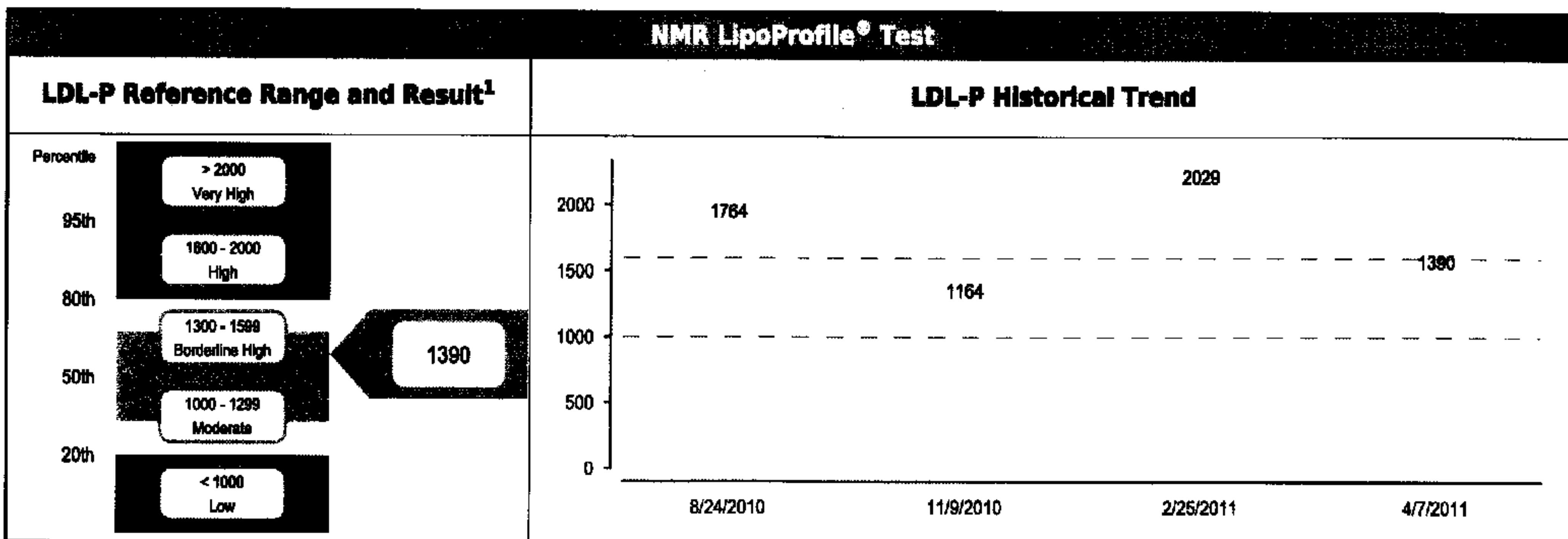
HDL 20.0

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<b>Patient</b>	<b>Name:</b>		<b>Phone #:</b>		<b>Patient ID #:</b>	
	Jonathan Aaron				10-237-0263	
	<b>Fasting Status:</b>		<b>Gender:</b>		<b>Birthdate:</b>	
	Non-Fasting		Male		11/3/1961	
	<b>Height:</b>		<b>Weight:</b>		<b>BMI:</b>	

<b>Specimen</b>	<b>Collection Time:</b>	<b>Specimen ID:</b>
	<b>11:30 am</b>	<b>11040800203</b>
	<b>Collection Date:</b>	<b>Report Type:</b>
	<b>4/7/2011</b>	<b>Complete</b>
	<b>Received Date:</b>	<b>Report Date:</b>
	<b>4/8/2011</b>	<b>4/11/2011</b>

<b>Provider</b>	Requesting Provider: <b>Michael Richman</b> The Center for Cholesterol Management 1950 Sawtelle Blvd #150 Los Angeles, CA 90025
	Client ID:
	<b>06-90025-18-0000383</b>



Particle Concentration and Size							
Laboratory Test		Result	Percentile in Reference Population <sup>2</sup>				Previous Results
			Higher CVD Risk			Lower CVD Risk	
HDL Particles	<b>HDL-P (total)</b> μmol/L	29.8	low	25th (26.7)	50th (30.5)	75th (34.9)	high
							24.9
Small LDL-P and LDL Size are associated with CVD risk, but not after LDL-P is taken into account.			Insulin Resistance			Insulin Sensitive	
Lipoprotein Markers Associated with Insulin Resistance and Diabetes Risk <sup>3,4</sup>	<b>LARGE VLDL-P</b> nmol/L	6.6	high	75th (6.9)	50th (2.7)	25th (0.9)	low
	<b>SMALL LDL-P</b> nmol/L	668	high	75th (839)	50th (527)	25th (117)	low
	<b>LARGE HDL-P</b> μmol/L	< 0.7	low	25th (3.1)	50th (4.8)	75th (7.3)	high
	<b>VLDL SIZE</b> nm	48.2	large	75th (52.5)	50th (46.6)	25th (42.4)	small
<b>LDL SIZE</b> nm	20.4	small	25th (20.4)	50th (20.8)	75th (21.2)	large	
<b>HDL SIZE</b> nm	< 8.3	small	25th (8.9)	50th (9.2)	75th (9.8)	large	
<b>LP-IR SCORE*</b> 0 - 100	73	insulin resistant	75th (83)	50th (45)	25th (27)	insulin sensitive	

**LP-IR Score is Inaccurate if a patient is non-fasting.**

\* The LP-IR Score combines the information from the 6 markers above it to give improved assessment of insulin resistance and diabetes risk.

**These laboratory assays, validated by LipoScience, have not been cleared by the US Food and Drug Administration. The clinical utility of these laboratory values has not been fully established.**

1. Reference population comprises 5,362 men and women not on lipid medication enrolled in the Multi-Ethnic Study of Atherosclerosis (MESA). Mora, et al. *Atherosclerosis* 2007.

1. LipidScience population comprises 3,500 men and women not on lipid medication enrolled in the multi-center Study of Cardiovascular
2. LipScience reference population comprises 4,588 men and women without known CVD or diabetes and not on lipid medication.

3. Garvey WT, et al. *Diabetes*. 2003; 532:463-462. 4. Goff DC et al. *Metabolism*. 2005; 54:264-270.

**Dr. Joseph P. McConnell | Laboratory Director | CLIA No. 49D1100708 | CAP No. 7224971 | NPI No. 1629209853**

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HDL 20.0



Patient	Name:		Phone #:		Patient ID #:	
	Jonathan Aaron				10-237-0263	
	Fasting Status:		Gender:		Birthdate:	
	Non-Fasting		Male		11/3/1961	
	Age:				49	
	Height:		Weight:		BMI:	
					Prev. BMI:	

Specimen	Collection Time:		Specimen ID:	
	11:30 am		11040800203	
	Collection Date:		Report Type:	
	4/7/2011		Complete	
	Received Date:		Report Date:	
	4/8/2011		4/11/2011	

Provider	Requesting Provider:	
	Michael Richman	
	The Center for Cholesterol Management	
	1950 Sawtelle Blvd #150	
	Los Angeles, CA 90025	
	Client ID:	
	06-90025-18-0000383	

Comments:

<p>Decreased HDL cholesterol and increased triglyceride concentrations raise the suspicion of the presence of the metabolic syndrome. According to ATP III guidelines, a diagnosis of metabolic syndrome can be made if 3 or more of the following criteria are met: Triglycerides equal to or great than 150 mg/dl, HDL of equal to or less than 50 mg/dl for women and 40 mg/dl for men, fasting glucose equal to or greater than 110 mg/dl, systolic blood pressure greater than or equal to 130mm HG and diastolic blood pressure equal to or greater than 85 mm HG, and waist circumference for men greater than 40 inches (102 cm) and for women greater than 35 inches (88 cm).</p>
<p>HDL cholesterol, HDL 2 (large HDL particle) cholesterol, and apolipoprotein A-I are decreased or in the intermediate range. Low HDL cholesterol and low apolipoprotein A-I have been associated with increased risk for cardiovascular disease and events. Low amounts of large HDL particles suggest that the HDL present may not be in its most protective form. HDL cholesterol, large HDL particle, and apolipoprotein A-I concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have been demonstrated to increase large HDL particle concentration.</p>
<p>Although LDL cholesterol is optimal or near optimal, small dense LDL cholesterol, %sdLDL, and Apo B are increased or in the intermediate range in this sample, consistent with the presence of small dense LDL particles. Studies have shown that elevated small dense LDL particle concentration is associated with increased risk for coronary heart disease even in the presence of optimal LDL cholesterol values. Small LDL particles may be observed in association with the metabolic syndrome and pre-diabetes. Statins effectively reduce the number of LDL particles, but do not generally influence the size distribution of the LDL particles. Fibrates and niacin have been shown to increase LDL particle size.</p>
<p>Although the LDL cholesterol concentration is optimal, LDL particle concentration is increased in this sample. Studies have shown that elevated LDL particle concentration is associated with increased risk for coronary heart disease, even in the presence of optimal LDL cholesterol values. Small LDL particles may be observed in association with the metabolic syndrome and pre-diabetes. Statins effectively reduce the number of LDL particles, but do not generally influence the size distribution of the LDL particles. Niacin, fibrates, and combination therapy (statin +niacin) have been shown to increase LDL particle size.</p>
<p>The Apo B:Apo A-I ratio was increased. Recently large case control studies have demonstrated that the Apo B:Apo A-I ratio is superior to cholesterol measures and cholesterol ratios for predicting risk for myocardial infarction. In the Interheart study, comparing 12,461 myocardial infarction cases to 14,637 age and gender matched controls in 52 countries, the Apo B:Apo A-I ratio was vastly superior to any of the cholesterol parameters measured including the LDL cholesterol: HDL cholesterol ratio and the total cholesterol to HDL cholesterol ratio in all ethnic groups, in both sexes, and at all ages. Decreasing the Apo B:Apo A-I ratio can be achieved by lowering Apo B and/or by increasing Apo A-I. Statins effectively reduce Apo B as do fibrates and niacin. Combination therapy (statin + niacin) is particularly effective at reducing Apo B, especially when small dense LDL particles are present. Apo A-I concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have also been demonstrated to increase Apo A-I.</p>
<p>Homocysteine is in the intermediate range. Increases in homocysteine concentration can occur with aging, menopause, hypothyroidism, low plasma levels of vitamin cofactors (B6, B12 and folate), certain drugs, and chronic renal insufficiency. Genetic variation in enzymes involved in homocysteine metabolism contributes to inter-individual differences in plasma homocysteine levels.</p>
<p>Elevated fasting insulin. If a fasting insulin level is elevated, it reflects hyperinsulinemia but fasting levels can be normal when levels following a glucose load are elevated. Insulin is elevated postprandially in proportion to the carbohydrate content in the meal. Elevated fasting insulin levels have been related to atherosclerosis risk. The combination of elevated fasting insulin, apolipoprotein B levels, and small LDL size identifies a very high-risk group for the development of ischemic heart disease.</p>
<p>Vitamin D concentration is in the intermediate range. Decreased vitamin D has been associated with hypertension, inflammation, and the metabolic syndrome. More recently, low serum 25(OH)D has been associated with increased incidence of cardiovascular events and all cause mortality.</p>



Patient	Name:		Phone #:		Patient ID #:	
	Jonathan Aaron				10-237-0263	
	Fasting Status:		Gender:		Birthdate:	
	Non-Fasting		Male		11/3/1961	
	Age:		Height:		Weight:	
	49				BMI:	
	Prev. BMI:					
Specimen	Collection Time:		Specimen ID:			
	11:30 am		11040800203			
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	4/7/2011		Complete			
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	4/8/2011		4/11/2011			
Provider	Requesting Provider:					
	Michael Richman		The Center for Cholesterol Management			
	1950 Sawtelle Blvd #150		Los Angeles, CA 90025			
	Client ID:					
	06-90025-18-0000383					

Comments:

Total HDL particle concentration is in the intermediate range in this sample. Decreased HDL particles have been associated with increased risk for cardiovascular disease. HDL particle concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have been demonstrated to increase HDL particle concentration.

End of Report

ATTN PATIENT: Please contact HDL, Inc. at 1-877-4HDLABS (1-877-443-5227) to set an appointment with your personal health coach to discuss your diet and exercise needs at no charge. You can also visit us online at [www.myhdl.com](http://www.myhdl.com) and schedule an appointment through our web portal.

Joseph P. McConnell, Laboratory Director

CLIA No. 49D1100708 | CAP No. 7224971 | NPI No. 1629209853

0000383-0183353

0000383-0183353

0000383-0183353

INSTRUCTIONS

- 1) Please fill in all of the highlighted areas.  
2) Have patient sign Release and Assignment of Benefits below.  
3) Copy BOTH sides of patient's insurance card(s).  
4) Indicate primary and secondary diagnoses below.
- NOTE: Physicians (or other individuals authorized to order tests) should only order tests that are medically necessary and reasonable.

0000383-0183353

0000383-0183353

0000383-0183353

PATIENT INFORMATION

HDL, Inc. will accept the following Demographic Sheet as substitute for Patient Information, provided that it contains all required info.

\*Last Name: First: Middle Initial: Client Patient ID #:

Address:

City: State: Zip Code:

Home Phone: Sex: M. F. Date of Birth:

Work Phone: Social Security #:

\*Height: feet inches \*Weight: pounds

SPECIMEN INFORMATION

Drawing Lab: Phone:

Collection Date: / / Time: am/pm Fasting: ☐ Yes ☐ No

Initials: Insulin: time of last dose: am/pm date: / /

HDL, INC. USE ONLY

Received Date: / / Time: am/pm Initials:

# Serum SST™ 8.5mL tubes: # Plasma PPT™ "Pearl" 5mL tubes:

# Serum "Bumble Bee" 5mL tubes: # Urine 8mL tubes:

# Whole Blood EDTA 4mL tubes: # Other tubes:

BILLING INFORMATION

☐ INSURANCE: Please attach a copy of both sides of patient's insurance card

☐ MEDICARE No.: Is coverage secondary? ☐ Yes ☐ No

☐ CHECK: Please make check payable to Health Diagnostic Laboratory, Inc.

☐ CREDIT CARD No.: Exp. Date: /

Name as it appears on card:

Total Amount: \$ Card Issuer: ☐ Visa ☐ MC ☐ Amex

RELEASE AND ASSIGNMENT OF BENEFITS

As a courtesy, HDL, Inc. will make every reasonable effort to obtain reimbursement for ordered tests. I authorize HDL, Inc. to release to Medicare or its Carriers & any insurance carrier providing medical benefits to me and any health plan of which I am a member, any medical or other information needed for the claim purposes. I authorize payment of Medical Insurance Benefits to the party who bills for this claim & accepts assignment.

Bill to my insurance: I understand that if my insurance company pays me directly for services rendered by HDL, Inc., I am responsible for forwarding such payment to HDL, Inc. I also understand, I am responsible for any deductibles/copayment, as required by my plan.

Important: Insurance regulations require HDL, Inc. to seek payment. I permit a copy of this authorization to be in place of the original.

\*Patient Signature Date

COMMENTS

REQUESTING PROVIDER/LAB/INSTITUTION

☐ M. Richman  
The Center for Cholesterol Management  
1950 Sawtelle Blvd #150  
Los Angeles, CA 90025

Client ID: 06-80025-18-0000383  
Phone: (310) 481-3939  
Fax: (310) 481-3949 V. 4

\*Physician or Authorized Signature Date

CUSTOM PANELS

☐ Baseline Assessment  
Lipid Panel  
Apo A1  
Apo B  
LDL P  
sdLDL  
HDL 2 subclass  
Lp(a) mass w/reflex  
Apo E Genotype  
Factor V Leiden  
Prothrombin Mutation  
CYP2C19  
Lp-PLA2  
CRP-hs  
Fibrinogen  
Homocysteine  
FFA (NEFA)  
Insulin

NT pro BNP  
Vitamin D  
HbA1C  
CMP

☒ Follow-Up Profile  
Lipid Panel  
Apo A1  
Apo B  
LDL P  
sdLDL  
HDL 2 subclass  
Lp(a) mass w/reflex  
Lp-PLA2  
CRP-hs  
Fibrinogen  
Homocysteine  
FFA (NEFA)  
Insulin  
NT pro BNP  
Vitamin D  
HEP  
HbA1C

INDIVIDUAL TESTS

Routine Panels (see reverse side for details)

☐ Basic Metabolic Panel 80048  
☐ Comp Metabolic Panel 80053  
☐ Electrolyte Panel 80051  
☐ Hepatic Function Panel 80076  
☐ Lipid Panel 80061  
☐ Renal Panel 80069  
☐ Thyroid Cascade Panel 84480, 84436, 84439

Lipoprotein Particles & Apolipoproteins

☐ Apolipoprotein A-I 82172  
☐ Apolipoprotein B 82172  
☐ HDL2 Subclass 82664  
☐ LDL-P & HDL-P 83704  
☐ Lp(a) cholesterol 82664  
☐ Lp(a) mass 83695  
☐ sdLDL 83701

Metabolic

☐ C-peptide 84681  
☐ FFA/NEFA 82726  
☒ Glucose 82947  
☐ Hemoglobin A1c 83036  
☐ Homocysteine 83090  
☐ Insulin 83525  
☐ RBC Folate 82747, 85014  
☐ TSH 84443  
☐ Uric Acid 84550  
☒ Vitamin B12 82607  
☐ 25-hydroxy-Vitamin D 82652

Inflammation & Platelets

☐ AspirinWorks® (urine) 83520 & 82565  
☐ F2-Isoprostanes 83789, 82570  
☐ Fibrinogen 85384  
☐ hs-CRP 86141  
☐ Lp-PLA2 83698  
☐ MPO 83516  
☐ NT-proBNP 83880

Hormones

☐ FSH 83001  
☐ LH 83002  
☐ Testosterone 84403

Genetic Assays

☐ Apo E Genotype  
☐ CYP2C19  
☐ Factor V Leiden  
☐ MTHFR  
☐ Prothrombin Mutation  
☐ Warfarin Sensitivity (CYP2C9 & VKORC1)

CPT Codes used for all Genetic Assays: 83891, 83892, 83896, 83903, 83908, 83912

Thyroid Function

☐ T4 84436  
☐ TSH 84443  
☐ T3 84480  
☐ T4, free 84439  
☐ T Uptake 84482

Other

☒ Omega Guard

Renal

☐ Creatinine (serum) 82565  
☐ Cystatin C 82610  
☐ Microalbumin (urine) 82043

DIAGNOSIS (ICD-9 CODE)

NOTE: The below ICD-9 codes are listed as a convenience. Ordering physicians should report the code that best describes the reason for ordering the tests & are not required to use the codes provided below.

Please write in Diagnosis Codes below or clearly mark to the right. PLEASE USE FOURTH & FIFTH DIGIT MODIFIERS.

Primary: ☒ Mixed Hyperlipidemia 272.2  
Secondary: ☐ Hypertension 401.  
Other: ☐ Hypertensive Heart Disease 402.  
☐ Hypertension w/Renal Disease 403.

☐ Diabetes 250.  
☐ Pure Hypercholesterolemia 272.0  
☐ Pure Hyperglyceridemia 272.1  
☐ Unspecified Hyperlipidemia 272.4  
☐ Hypertension 401.  
☐ Hypertensive Heart Disease 402.  
☐ Hypertension w/Renal Disease 403.

☐ Hypertension w/Renal Disease & Heart Failure 404.  
☐ Intermediate Coronary Syndrome 411.1  
☐ Coronary Atherosclerosis Unspecified Vessel, Native or Graft 414.00  
☐ Congestive Heart Failure 428.  
☐ Cirrhosis of liver without mention of alcohol 571.5  
☐ Other chronic non-alcoholic liver disease 571.8  
☐ Unspecified chronic liver disease without mention of alcohol 571.9

OTHER 277.89



LIPOSCIENCE

LipoScience, Inc.  
2500 Sumner Boulevard  
Raleigh, NC 27616  
877-547-6837  
www.liposcience.com

Page 1 of 1

Patient Name		Sex	Age	Clinician	
Aaron, Jonathan		M	49		
Patient ID	Birth Date	Accession Number		Client Name and Address	
11022600165	11/03/1961	S0372344		HDL-Health Diagnostic Lab Inc 17730/ 737 North 5th Street Suite 103 Richmond, VA 23219 Phone: (804)343-2718 Fax: (804)343-0929	
Date Collected	Date Received	Report Date and Time		Requisition Number	Fasting Status
02/25/2011	02/28/2011	02/28/2011 18:52		11022600165	Not Specified

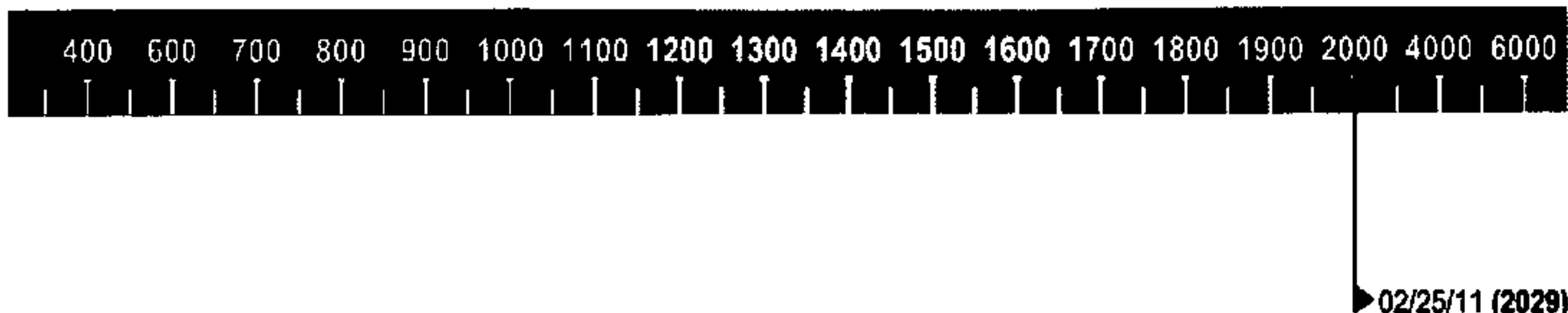
NMR LipoProfile® test

Reference Range<sup>1</sup>

	Percentile <sup>1</sup>	20th	50th	80th	95th	
	nmol/L	Low	Moderate	Borderline-High	High	Very High
<b>LDL-P</b> (LDL Particle Number)	<b>2029</b>	<b>&lt; 1000</b>	<b>1000-1299</b>	<b>1300-1599</b>	<b>1600-2000</b>	<b>&gt; 2000</b>

Historical Reporting

LDL-P



LDL-C



↑↑ LDL-P ~ 2029  
TG - 494  
HDL-C - 39

- restart all  
- meds  
- F/U 2 months

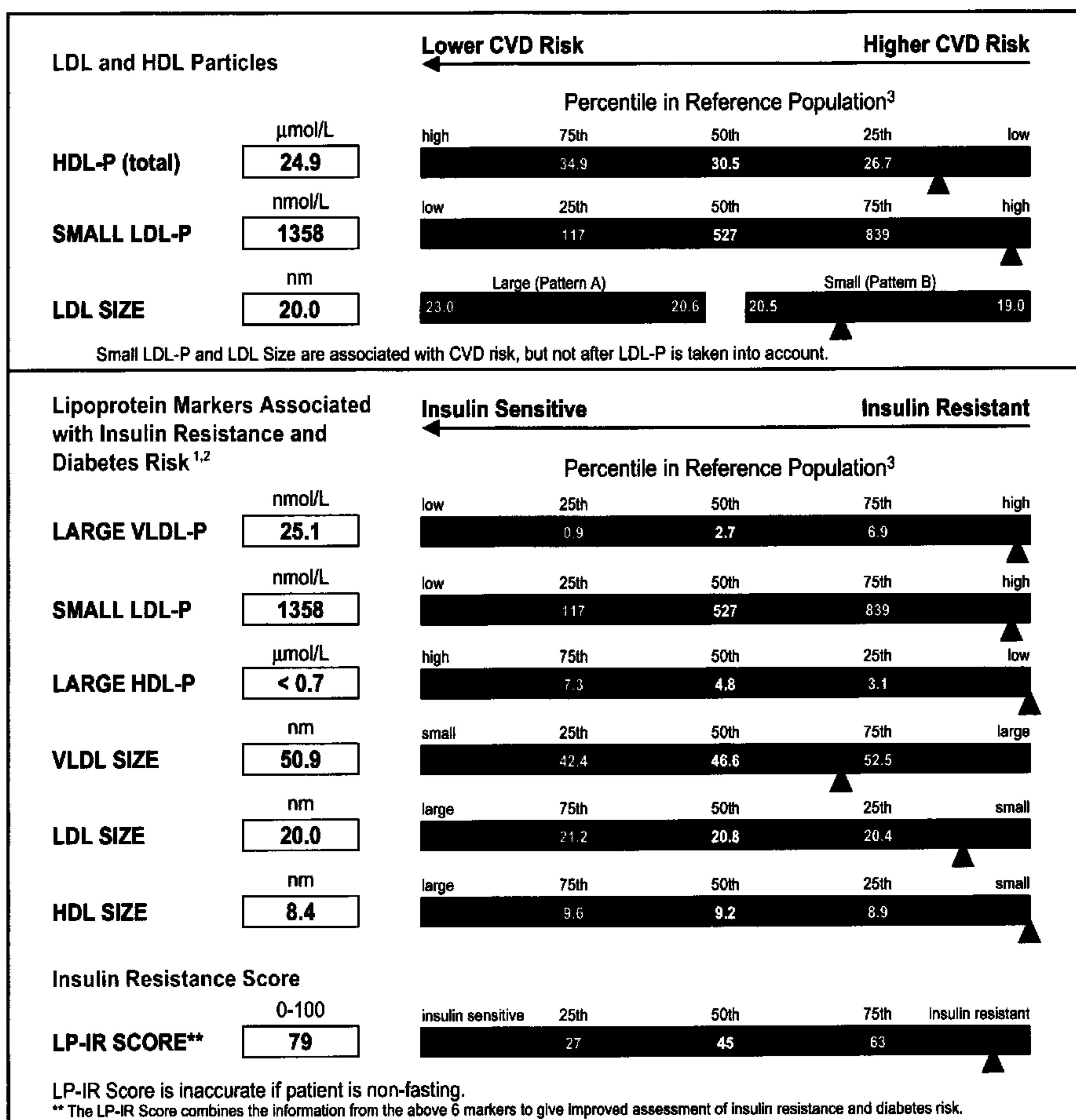
1. Reference population comprises 5,362 men and women not on lipid medication enrolled in the Multi-Ethnic Study of Atherosclerosis (MESA). Mora, et al. Atherosclerosis 2007.

3/10/11



Patient Name			Sex	Age	Clinician	
Aaron, Jonathan			M	49		
Patient ID	Birth Date	Accession Number		Client Name and Address		
11022600165	11/03/1961	S0372344		HDL-Health Diagnostic Lab Inc 17730/ 737 North 5th Street Suite 103 Richmond, VA 23219 Phone: (804)343-2718 Fax: (804)343-0929		
Date Collected	Date Received	Report Date and Time		Requisition Number	Fasting Status	
02/25/2011	02/28/2011	02/28/2011 18:52		11022600165	Not Specified	

## PARTICLE CONCENTRATION AND SIZE



These laboratory assays, validated by LipoScience, have not been cleared by the US Food and Drug Administration. The clinical utility of these laboratory values has not been fully established.

Patient	Name:	Jonathan Aaron	Phone #:		Patient ID #:	10-237-0263		
	Fasting Status:	Non-Fasting	Gender:	Male	Birthdate:	11/3/1961	Age:	49
	Height:		Weight:		BMI:		Prev. BMI:	
Specimen	Collection Time:	10:30 am	Specimen ID:	11022600165	Collection Date:	2/25/2011	Report Type:	Complete
	Received Date:	2/26/2011	Report Date:	2/28/2011				
Provider	Requesting Provider: Michael Richman The Center for Cholesterol Management 1950 Sawtelle Blvd #150 Los Angeles, CA 90025							
	Client ID: 06-90025-18-0000383							

Laboratory Test		Notes	High Risk	Intermediate Risk	Optimal	High Risk Range	Intermediate Risk Range	Optimal Range	Previous Results 11/9/2010
Lipids	Total Cholesterol (mg/dL)		241			≥ 240	200 - 239	< 200	175
	LDL-C Direct (mg/dL)				99	≥ 130 CHD & CHD risk eq. > 100	100 - 129 CHD & CHD risk eq. 70 - 100	< 100 CHD & CHD risk eq. < 70	78
	HDL-C (mg/dL)		39			< 40		≥ 40	35
	Triglycerides (mg/dL)		494			≥ 200	150 - 199	< 150	304
	Non-HDL-C (mg/dL) (calculated)		202			≥ 160	130 - 159	< 130	140
Lipoprotein Particles and Apolipoproteins	Apo B (mg/dL)					≥ 80	60 - 79	< 60	94
	LDL-P (nmol/L)		2029			≥ 1300	1000 - 1299	< 1000	1164
	sdLDL (mg/dL)*					≥ 31	21 - 30	≤ 20	50
	% sdLDL (calculated)					≥ 23	14 - 22	≤ 13	64
	Apo A-I (mg/dL)					< 114	114 - 131	≥ 132	130
	HDL-P (μmol/L)		24.9			< 28.0	28.0 - 34.0	≥ 35.0	31.4
	HDL2 (mg/dL)*					≤ 8	9 - 11	≥ 12	6
	Apo B:Apo A-I Ratio (calculated)					≥ 0.81	0.61 - 0.81	≤ 0.6	0.73
	Lp(a) Mass (mg/dL)					≥ 30		< 30	3
	Lp(a) Cholesterol (mg/dL)					≥ 6	3 - 5	< 3	
Inflammation/ Oxidation	Myeloperoxidase (pmol/L)					≥ 550	400 - 549	< 400	
	Lp-PLA <sub>2</sub> (ng/mL)					> 235	200 - 235	< 200	163
	hs-CRP (mg/L)					≥ 3.0	1.0 - 2.9	< 1.0	1.15
	Fibrinogen (mg/dL)					≥ 465	391 - 464	≤ 390	313
Myocardial Stress	NT-proBNP (pg/mL)					≥ 450	125 - 449	< 125	54
Platelets	AspirinWorks® (urine) (pg/mg of creatinine)					> 1500		≤ 1500	

Lab Notes:

Provider Notes:



Patient	Name: Jonathan Aaron		Phone #:		Patient ID #: 10-237-0263	
	Fasting Status: Non-Fasting		Gender: Male		Birthdate: 11/3/1961    Age: 49	
	Height:		Weight:		BMI:    Prev. BMI:	
Specimen	Collection Time: 10:30 am		Specimen ID: 11022600165			
	Collection Date: 2/25/2011		Report Type: Complete			
	Received Date: 2/26/2011		Report Date: 2/28/2011			
Provider	Requesting Provider: Michael Richman The Center for Cholesterol Management 1950 Sawtelle Blvd #150 Los Angeles, CA 90025					
	Client ID: 06-90025-18-0000383					

Laboratory Test		Notes	High Risk	Intermediate Risk	Optimal	High Risk Range	Intermediate Risk Range	Optimal Range	Previous Results 11/9/2010
Lipoprotein Genetics	Apolipoprotein E Genotype*					Estimated Genotype Frequency: 2/2 (~1-2%), 2/3 (~15%), 2/4 (~1-2%), 3/3 (~55%), 3/4 (~25%), 4/4 (~1-2%)			3/3
Platelet Genetics	CYP2C19*					Estimated Genotype Frequency: *1/*1 (~60-70%), *1/*2 (~20-30%), *2/*2 (~2-3%), *1/*3 (<1%), *2/*3 (<0.05%), *3/*3 (<0.01%)			*1/*1
Coagulation Genetics	Factor V Leiden					Optimal=Non-carrier (Arg/Arg); At Risk=(Arg/Gln or Gln/Gln)			Arg/Arg
	Prothrombin Mutation					Optimal=Non-carrier (G/G); At Risk=(G/A or A/A)			G/G
Metabolic	Insulin (µU/mL)					≥ 12	10 - 11	3 - 9	44
	Free Fatty Acid (mmol/L)					> 0.7	0.6 - 0.7	≤ 0.59	0.45
	Glucose (mg/dL)					≤ 55 or > 125	56-69 or 100-125	70 - 99	
	HbA1c (%)					≥ 6.5	6.0 - 6.4	4.8 - 5.9	
	25-hydroxy-Vitamin D (ng/mL)					≤ 14	15 - 29	30 - 100	29
	Homocysteine (µmol/L)					> 13	11 - 13	≤ 10	12

se call 1-877-4HDLABS (1-877-443-5227) or visit us online at www.myhdl.com

Lab Notes:

<b>Patient</b>	Name:		Phone #:	Patient ID #:	
	Jonathan Aaron			10-237-0263	
	Fasting Status:		Gender:	Birthdate:	Age:
	Non-Fasting		Male	11/3/1961	49
	Height:	Weight:	BMI:	Prev. BMI:	

<b>Specimen</b>	Collection Time:	Specimen ID:
	10:30 am	11022600165
	Collection Date:	Report Type:
	2/25/2011	Complete
	Received Date:	Report Date:
	2/26/2011	2/28/2011

**Provider**

**Requesting Provider:**  
**Michael Richman**  
**The Center for Cholesterol Management**  
**1950 Sawtelle Blvd #150**  
**Los Angeles, CA 90025**

---

**Client ID:**  
**06-90025-18-0000383**

Other Biomarkers	Result	Flag	Reference Interval
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CK (U/L)	147		20 - 200
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**Lab Notes:**

Patient	Name:		Phone #:		Patient ID #:	
	Jonathan Aaron				10-237-0263	
	Fasting Status:		Gender:		Birthdate:	
	Non-Fasting		Male		11/3/1961	
	Age:				49	
	Height:		Weight:		BMI:	
					Prev. BMI:	

Specimen	Collection Time:		Specimen ID:	
	10:30 am		11022600165	
	Collection Date:		Report Type:	
	2/25/2011		Complete	
	Received Date:		Report Date:	
	2/26/2011		2/28/2011	

Provider	Requesting Provider:	
	Michael Richman The Center for Cholesterol Management 1950 Sawtelle Blvd #150 Los Angeles, CA 90025	
	Client ID:	
	06-90025-18-0000383	

Comments:

<p>Decreased HDL cholesterol and increased triglyceride concentrations raise the suspicion of the presence of the metabolic syndrome. According to ATP III guidelines, a diagnosis of metabolic syndrome can be made if 3 or more of the following criteria are met: Triglycerides equal to or greater than 150 mg/dl, HDL of equal to or less than 50 mg/dl for women and 40 mg/dl for men, fasting glucose equal to or greater than 110 mg/dl, systolic blood pressure greater than or equal to 130mm HG and diastolic blood pressure equal to or greater than 85 mm HG, and waist circumference for men greater than 40 inches (102 cm) and for women greater than 35 inches (88 cm).</p>
<p>Although the LDL cholesterol concentration is optimal, LDL particle concentration is increased in this sample. Studies have shown that elevated LDL particle concentration is associated with increased risk for coronary heart disease, even in the presence of optimal LDL cholesterol values. Small LDL particles may be observed in association with the metabolic syndrome and pre-diabetes. Statins effectively reduce the number of LDL particles, but do not generally influence the size distribution of the LDL particles. Niacin, fibrates, and combination therapy (statin + niacin) have been shown to increase LDL particle size.</p>
<p>Total HDL particle concentration is decreased in this sample. Decreased HDL particles have been associated with increased risk for cardiovascular disease. HDL particle concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have been demonstrated to increase HDL particle concentration.</p>

End of Report

ATTN PATIENT: Please contact HDL, Inc. at 1-877-4HDLABS (1-877-443-5227) to set an appointment with your personal health coach to discuss your diet and exercise needs at no charge. You can also visit us online at <a href="http://www.myhdl.com">www.myhdl.com</a> and schedule an appointment through our web portal.
---



INSTRUCTIONS

- 1) Please fill in all of the highlighted areas.
  - 2) Have patient sign Release and Assignment of Benefits below.
  - 3) Copy BOTH sides of patient's insurance card(s).
  - 4) Indicate primary and secondary diagnoses below.
- NOTE: Physicians (or other individuals authorized to order tests) should only order tests that are medically necessary and reasonable.

PATIENT INFORMATION

*Last Name:	First:	Middle Initial:	Client Patient ID #:
Address:			
City:	State:	Zip Code:	
Home Phone:	Sex: M F	Date of Birth:	
Work Phone:	Social Security #:		
*Height:	*Weight:		
feet inches	pounds		

SPECIMEN INFORMATION

Drawing Lab:	Phone:
Collection Date: / /	Time: am/pm
Fasting: <input type="checkbox"/> Yes <input type="checkbox"/> No	Hrs <input type="checkbox"/> No
Initials:	Insulin: time of last dose: am/pm date: / /

HDL, INC. USE ONLY

Plasma PFT 5ml tubes:	Plasma PFT 5ml tubes:
Serum 5ml tubes:	Urine 5ml tubes:
Whole blood EDTA 4ml tubes:	Other tubes:

BILLING INFORMATION

<input type="checkbox"/> INSURANCE: Please attach a copy of both sides of patient's insurance card
<input type="checkbox"/> MEDICARE No.: Is coverage secondary? <input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> CHECK: Please make check payable to Health Diagnostic Laboratory, Inc.
<input type="checkbox"/> CREDIT CARD No.: Exp. Date: / /
Name as it appears on card:
Total Amount: \$ Card issuer: <input type="checkbox"/> Visa <input type="checkbox"/> MC <input type="checkbox"/> Amex

RELEASE AND ASSIGNMENT OF BENEFITS

As a courtesy, HDL, Inc. will make every reasonable effort to obtain reimbursement for ordered tests. I authorize HDL, Inc. to release to Medicare or its Carriers & any insurance carrier providing medical benefits to me and any health plan of which I am a member, any medical or other information needed for the claim purposes. I authorize payment of Medical Insurance Benefits to the party who bills for this claim & accepts assignment.

Bill to my insurance: I understand that if my insurance company pays me directly for services rendered by HDL, Inc., I am responsible for forwarding such payment to HDL, Inc. I also understand, I am responsible for any deductibles/copayment, as required by my plan.

Important: Insurance regulations require HDL, Inc. to seek payment. I permit a copy of this authorization to be in place of the original.

\*Patient Signature Date

COMMENTS

Primary:
Secondary:
Other:

Please write in Diagnosis Codes below or clearly mark to the right. PLEASE USE FOURTH & FIFTH DIGIT MODIFIERS.

Primary:

Secondary:

Other:

- |  |  |   |   |   |  |  |  |
|--|--|---|---|---|--|--|--|
| <input type="checkbox"/> Diabetes 250. | <input type="checkbox"/> Pure Hypercholesterolemia 272.0 | <input type="checkbox"/> Pure Hyperglyceridemia 272.1 | <input type="checkbox"/> Mixed Hyperlipidemia 272.2 | <input type="checkbox"/> Unspecified Hyperlipidemia 272.4 | <input type="checkbox"/> Hypertension 401. | <input type="checkbox"/> Hypertensive Heart Disease 402. | <input type="checkbox"/> Hypertension w/Renal Disease 403. |
|--|--|---|---|---|--|--|--|

- |  |   |  |  |  |  |   |                                |
|--|---|--|--|--|--|---|--------------------------------|
| <input type="checkbox"/> Hypertension w/Renal Disease & Heart Failure 404. | <input type="checkbox"/> Intermediate Coronary Syndrome 411.1 | <input type="checkbox"/> Coronary Atherosclerosis Unspecified Vessel, Native or Graft 414.00 | <input type="checkbox"/> Congestive Heart Failure 428. | <input type="checkbox"/> Cirrhosis of liver without mention of alcohol 571.5 | <input type="checkbox"/> Other chronic non-alcoholic liver disease 571.8 | <input type="checkbox"/> Unspecified chronic liver disease without mention of alcohol 571.9 | <input type="checkbox"/> OTHER |
|--|---|--|--|--|--|---|--------------------------------|

REQUESTING PROVIDER/LAB/INSTITUTION

<input type="checkbox"/> M. Richman The Center for Cholesterol Management 1950 Sawtelle Blvd #150 Los Angeles, CA 90025	Client ID: 06-90025-18-0000383 Phone: (310) 481-3939 Fax: (310) 481-3949
*Physician or Authorized Signature	Date

CUSTOM PANELS

- |  |   |
|--|---|
| <input type="checkbox"/> Baseline Assessment | <input checked="" type="checkbox"/> Follow-Up Profile |
| Lipid Panel                                  | Lipid Panel   |
| Apo A1                                       | Apo A1  |
| Apo B  | Apo B   |
| LDL P  | LDL P   |
| sdLDL  | sdLDL   |
| HDL 2 subclass                               | HDL 2 subclass  |
| Lp(a) mass w/reflex                          | Lp(a) mass w/reflex                                   |
| Apo E Genotype                               | Lp-PLA2   |
| Factor V Leiden                              | CRP-hs  |
| Prothrombin Mutation                         | Fibrinogen  |
| CYP2C19                                      | Homocysteine  |
| Lp-PLA2                                      | FFA (NEFA)  |
| CRP-hs                                       | Insulin   |
| Fibrinogen                                   | NT pro BNP  |
| Homocysteine                                 | Vitamin D   |
| FFA (NEFA)                                   | HEP   |
| Insulin                                      | HbA1C   |

INDIVIDUAL TESTS

Routine Panels (see reverse side for details)	Lipoprotein Particles & Apolipoproteins	Genetic Assays
<input type="checkbox"/> Basic Metabolic Panel 80048	<input type="checkbox"/> Apolipoprotein A-I 82172	<input type="checkbox"/> Apo E Genotype
<input type="checkbox"/> Comp Metabolic Panel 80053	<input type="checkbox"/> Apolipoprotein B 82172	<input type="checkbox"/> CYP2C19
<input type="checkbox"/> Electrolyte Panel 80051	<input type="checkbox"/> HDL2 Subclass 82664	<input type="checkbox"/> Factor V Leiden
<input type="checkbox"/> Hepatic Function Panel 80076	<input type="checkbox"/> LDL-P & HDL-P 83704	<input type="checkbox"/> MTHFR
<input type="checkbox"/> Lipid Panel 80061	<input type="checkbox"/> Lp(a) cholesterol 82664	<input type="checkbox"/> Prothrombin Mutation
<input type="checkbox"/> Renal Panel 80069	<input type="checkbox"/> Lp(a) mass 83695	<input type="checkbox"/> Warfarin Sensitivity (CYP2C9 & VKORC1)
<input type="checkbox"/> Thyroid Cascade Panel 84480, 84436, 84439	<input type="checkbox"/> sdLDL 83701	CPT Codes used for all Genetic Assays: 83891, 83892, 83896, 83903, 83908, 83912
Inflammation & Platelets	Metabolic	Thyroid Function
<input type="checkbox"/> AspirinWorks* (urine) 83520 & 82565	<input type="checkbox"/> C-peptide 84681	<input type="checkbox"/> T4 84436
<input type="checkbox"/> F <sub>2</sub> -Isoprostanes 83789, 82570	<input type="checkbox"/> FFA/NEFA 82726	<input type="checkbox"/> TSH 84443
<input type="checkbox"/> Fibrinogen 85384	<input checked="" type="checkbox"/> Glucose 82947	<input type="checkbox"/> T3 84480
<input type="checkbox"/> hs-CRP 86141	<input type="checkbox"/> Hemoglobin A1c 83036	<input type="checkbox"/> T4, free 84439
<input type="checkbox"/> Lp-PLA <sub>2</sub> 83698	<input type="checkbox"/> Homocysteine 83090	<input type="checkbox"/> T Uptake 84482
<input type="checkbox"/> MPO 83516	<input type="checkbox"/> Insulin 83525	
<input type="checkbox"/> NT-proBNP 83880	<input type="checkbox"/> RBC Folate 82747, 85014	Other
(may require additional ICD-9 coding)	<input type="checkbox"/> TSH 84443	<input type="checkbox"/>
Hormones	<input type="checkbox"/> Uric Acid 84550	<input type="checkbox"/>
<input type="checkbox"/> FSH 83001	<input type="checkbox"/> Vitamin B <sub>12</sub> 82607	<input type="checkbox"/>
<input type="checkbox"/> LH 83002	<input type="checkbox"/> 25-hydroxy-Vitamin D 82652	<input type="checkbox"/>
<input type="checkbox"/> Testosterone 84403	Renal	<input type="checkbox"/>
	<input type="checkbox"/> Creatinine (serum) 82565	<input type="checkbox"/>
	<input type="checkbox"/> Cystatin C 82610	<input type="checkbox"/>
	<input type="checkbox"/> Microalbumin (urine) 82043	<input type="checkbox"/>

DIAGNOSIS (ICD-9 CODE)

NOTE: The below ICD-9 codes are listed as a convenience. Ordering physicians should report the code that best describes the reason for ordering the tests & are not required to use the codes provided below.





Page 1 of 1

Clinician

Patient Name	Sex	Age
Aaron, Jonathan	M	49

Patient ID	Birth Date	Accession Number
10111000598	11/03/1961	W0759272

Client Name and Address	
HDL-Health Diagnostic Lab Inc 737 North 5th Street Suite 103 Richmond, VA 23219 Phone: (804)343-2718	17730/ Fax: (804)343-0929

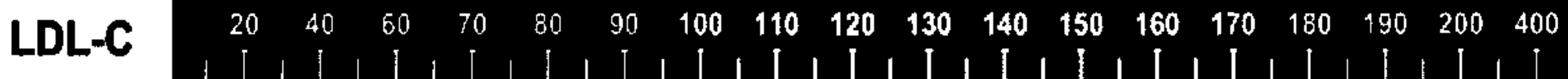
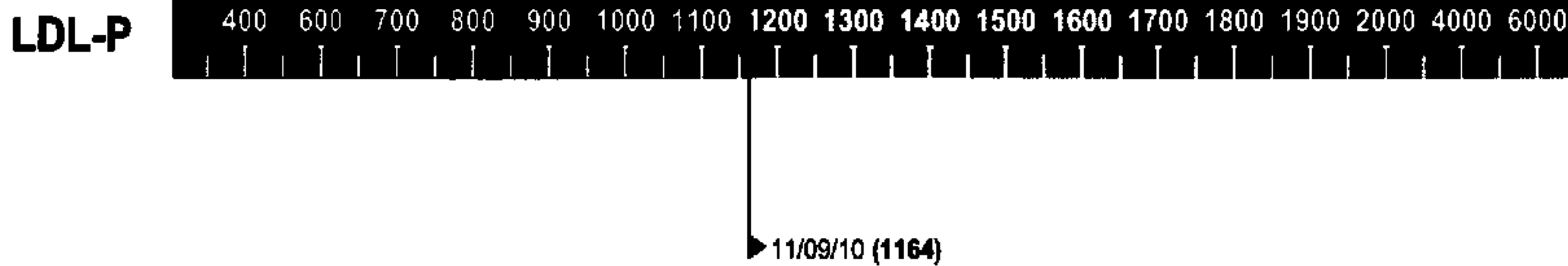
Date Collected	Date Received	Report Date and Time	Requisition Number	Fasting Status
11/09/2010	11/11/2010	11/12/2010 12:09	10111000598	Not Specified

## NMR LipoProfile® test

### Reference Range<sup>1</sup>

	Percentile <sup>1</sup>	20th	50th	80th	95th	
	nmol/L	Low	Moderate	Borderline-High	High	Very High
<b>LDL-P</b> (LDL Particle Number)	<b>1164</b>	<b>&lt; 1000</b>	<b>1000-1299</b>	<b>1300-1599</b>	<b>1600-2000</b>	<b>&gt; 2000</b>

## Historical Reporting



- near optimal LDL-P  
- TT TG but not taking  
① - cont meds  
② - verapamil 2gras BID  
③ - verapamil 5000ug  
④ - nutrition consult  
⑤ - F/u 2 months

1. Reference population comprises 5,362 men and women not on lipid medication enrolled in the Multi-Ethnic Study of Atherosclerosis (MESA). Mora, et al. Atherosclerosis 2007.

*[Handwritten signature]* 11/17/10



Patient Name	Sex	Age
Aaron, Jonathan	M	49

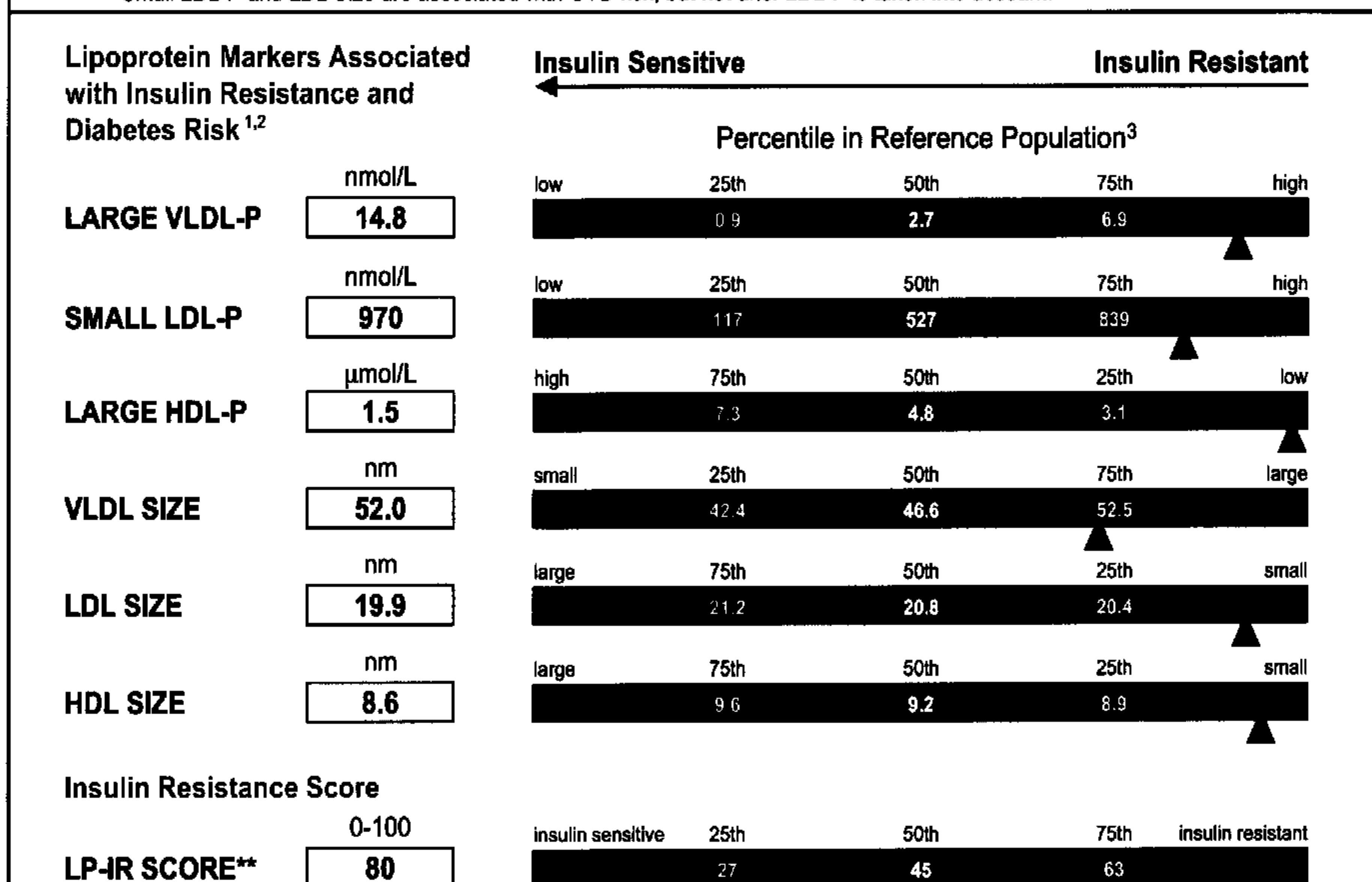
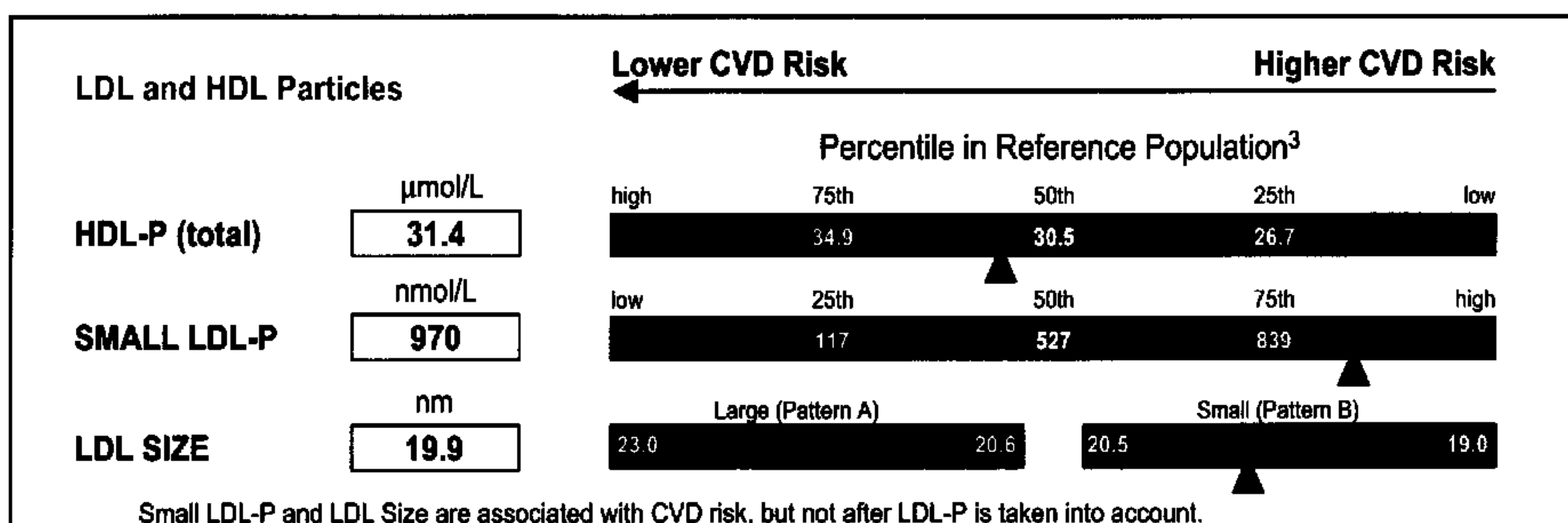
Patient ID	Birth Date	Accession Number
10111000598	11/03/1961	W0759272

Client Name and Address
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HDL-Health Diagnostic Lab Inc 737 North 5th Street Suite 103 Richmond, VA 23219 Phone: (804)343-2718	17730/ Fax: (804)343-0929
--	------------------------------

Date Collected	Date Received	Report Date and Time	Requisition Number	Fasting Status
11/09/2010	11/11/2010	11/12/2010 12:09	10111000598	Not Specified

## PARTICLE CONCENTRATION AND SIZE



LP-IR Score is inaccurate if patient is non-fasting.

\*\* The LP-IR Score combines the information from the above 6 markers to give improved assessment of insulin resistance and diabetes risk.

These laboratory assays, validated by LipoScience, have not been cleared by the US Food and Drug Administration. The clinical utility of these laboratory values has not been fully established.

Patient: Name: Jonathan Aaron, Gender: Male, Date of Birth: 11/3/1961, Patient ID #: 10-237-0263, Fasting Status: Non-Fasting, Age: 49, BMI:
Specimen: Collection Time: 10:40 am, Specimen ID: 10111000598, Collection Date: 11/9/2010, Report Type: Complete, Received Date: 11/10/2010, Report Date: 11/14/2010
Provider: Requesting Provider: Michael Richman, The Center for Cholesterol Management, 1950 Sawtelle Blvd #150, Los Angeles, CA 90025, Client ID: 06-90025-18-0000383

Laboratory Test		Notes	High Risk	Intermediate Risk	Optimal	High Risk Range	Intermediate Risk Range	Optimal Range	Previous Results
Lipids	Total Cholesterol (mg/dL)				175	≥ 240	200 - 239	< 200	263
	LDL-C Direct (mg/dL)				78	≥ 130 CHD & CHD risk eq. > 100	100 - 129 CHD & CHD risk eq. 70 - 100	< 100 CHD & CHD risk eq. < 70	92
	HDL-C (mg/dL)		35			< 40		≥ 40	32
	Triglycerides (mg/dL)		304			≥ 200	150 - 199	< 150	571
	Non-HDL-C (mg/dL) (calculated)			140		≥ 160	130 - 159	< 130	
Lipoprotein Particles and Apolipoproteins	Apo B (mg/dL)		94			≥ 80	60 - 79	< 60	119
	LDL-P (nmol/L)			1164		≥ 1300	1000 - 1299	< 1000	1764
	sdLDL (mg/dL)*		50			≥ 31	21 - 30	≤ 20	59
	% sdLDL (calculated)		64			≥ 23	14 - 22	≤ 13	65
	Apo A-I (mg/dL)			130		< 114	114 - 131	≥ 132	123
	HDL-P (μmol/L)			31.4		< 28.0	28.0 - 34.0	≥ 35.0	23.2
	HDL2 (mg/dL)*		6			≤ 8	9 - 11	≥ 12	12
	Apo B:Apo A-I Ratio (calculated)			0.73		≥ 0.81	0.61 - 0.81	≤ 0.6	0.97
	Lp(a) Mass (mg/dL)				3	≥ 30		< 30	2
	Lp(a) Cholesterol (mg/dL)					≥ 6	3 - 5	≤ 2	
Inflammation/Oxidation	Myeloperoxidase (pmol/L)					> 350	223 - 350	< 223	
	Lp-PLA <sub>2</sub> (ng/mL)				163	> 235	200 - 235	< 200	204
	hs-CRP (mg/L)			1.15		≥ 3.0	1.0 - 2.9	< 1.0	2.06
	Fibrinogen (mg/dL)				313	≥ 465	391 - 464	≤ 390	388
Myocardial Stress	NT-proBNP (pg/mL)				54	≥ 450	125 - 449	< 125	19
Platelets	AspirinWorks® (urine) (pg/mg of creatinine)					> 1500		≤ 1500	

Provider Notes:

To schedule time with a Personal Health Coach, please call 1-877-4HDLABS (1-877-443-5227) or visit us online at www.myhdi.com

Patient	Name:	Gender:	Date of Birth:	Specimen	Collection Time:	Specimen ID:	Provider	Requesting Provider:
	Jonathan Aaron	Male	11/3/1961		10:40 am	10111000598		Michael Richman
	Patient ID #:	Height:	Weight:		Collection Date:	Report Type:		The Center for Cholesterol Management
	10-237-0263				11/9/2010	Complete		1950 Sawtelle Blvd #150
	Fasting Status:	Age:	BMI:		Received Date:	Report Date:		Los Angeles, CA 90025
	Non-Fasting	49			11/10/2010	11/14/2010		Client ID:
								06-90025-18-0000383

Laboratory Test		Notes	High Risk	Intermediate Risk	Optimal	High Risk Range	Intermediate Risk Range	Optimal Range	Previous Results
Lipoprotein Genetics	Apolipoprotein E Genotype*					Estimated Genotype Frequency: 2/2 (~1-2%), 2/3 (~15%), 2/4 (~1-2%), 3/3 (~55%), 3/4 (~25%), 4/4 (~1-2%)			3/3
Platelet Genetics	CYP2C19*					Estimated Genotype Frequency: *1/*1 (~60-70%), *1/*2 (~20-30%), *2/*2 (~2-3%), *1/*3 (<1%), *2/*3 (<0.05%), *3/*3 (<0.01%)			*1/*1
Coagulation Genetics	Factor V Leiden					Optimal=Non-carrier (Arg/Arg); At Risk=(Arg/Gln or Gln/Gln)			Arg/Arg
	Prothrombin Mutation					Optimal=Non-carrier (G/G); At Risk=(G/A or A/A)			G/G
Metabolic	Insulin (µU/mL)		44			≥ 12	10 - 11	3 - 9	57
	C-peptide (ng/mL)					≤ 1.0 or > 4.4		1.1 - 4.4	
	Free Fatty Acid (mmol/L)				0.45	> 0.7	0.6 - 0.7	≤ 0.59	0.50
	Glucose (mg/dL)					≤ 55 or > 125	56-69 or 100-125	70 - 99	107
	HbA1c (%)					≥ 6.5	6.0 - 6.4	4.8 - 5.9	4.0
	25-hydroxy-Vitamin D (ng/mL)			29		≤ 14	15 - 29	30 - 100	12
	Uric Acid (mg/dL)					≥ 8	7.0 - 7.9	3.4 - 6.9	
	TSH (µIU/mL)					< 0.27 or > 4.20		0.27 - 4.20	
	Homocysteine (µmol/L)			12		> 13	11 - 13	≤ 10	12
	Vitamin B <sub>12</sub> (pg/mL)					< 211	211 - 299	300 - 946	
Renal	Microalbumin (urine) (mg albumin/g of creatinine)					≥ 30		< 30	
	Creatinine, serum (mg/dl)					> 1.2		0.7 - 1.2	0.9

Lab Notes:



Patient	Name:	Gender:	Date of Birth:	Specimen	Collection Time:	Specimen ID:	Provider	Requesting Provider:
	Jonathan Aaron	Male	11/3/1961		10:40 am	10111000598		Michael Richman
	Patient ID #:	Height:	Weight:		Collection Date:	Report Type:		The Center for Cholesterol Management
	10-237-0263				11/9/2010	Complete		1950 Sawtelle Blvd #150
	Fasting Status:	Age:	BMI:		Received Date:	Report Date:		Los Angeles, CA 90025
	Non-Fasting	49			11/10/2010	11/14/2010		Client ID:
								06-90025-18-0000383

Other Biomarkers	Result	Flag	Reference Range	CBC with Differential / Platelet	Result	Flag	Units	Reference Interval
Adiponectin (ug/mL)			2 - 20	WBC	7.2		x10E3/ $\mu$ L	4.0 - 10.5
Albumin (g/dl)	4.4		3.5 - 5.2	RBC	4.72		x10E6/ $\mu$ L	4.1 - 5.6
ALP (U/L)	53		40 - 129	Hemoglobin	14.8		g/dL	12.5 - 17.0
ALT / GPT (U/L)	31		Up to 41	Hematocrit	43.9		%	36 - 50
AST / GOT (U/L)	28		Up to 40	MCV	93		fL	80 - 98
BUN (mg/dl)			6 - 20	MCH	31		pg	27 - 34
Calcium (mg/dL)			8.4 - 10.2	MCHC	34		g/dL	32 - 36
CK (U/L)	338	H	20 - 200	RDW	14.1		%	11.7 - 15
Cl- (mmol/L)			96 - 108	Platelets	208		x10E3/ $\mu$ L	140 - 415
CO <sub>2</sub> (mmol/L)			22 - 29	Neutrophils	57		%	40 - 74
CREAT urine (mg/dl)			39 - 259	Lymphs	34		%	14 - 46
D-Dimer ( $\mu$ g/mL)			< 0.463	Monocytes	7		%	4 - 13
Direct Bilirubin (mg/dL)	0.1		0.0 - 0.3	Eos	2.0		%	0 - 7
Ferritin (ng/mL)			30 - 400	Basos	0		%	0 - 3
Fructosamine ( $\mu$ mol/L)			205 - 285	Neutrophils (absolute)	4.1		x10E3/ $\mu$ L	1.8 - 7.8
FSH (mIU/mL)			1.5 - 12.4	Lymphs (absolute)	2.5		x10E3/ $\mu$ L	0.7 - 4.5
GGT (U/L)			8 - 61	Monocytes (absolute)	0.5		x10E3/ $\mu$ L	0.1 - 1.0
Iron ( $\mu$ g/dL)			59 - 158	Eos (absolute)	0.1		x10E3/ $\mu$ L	0.0 - 0.4
K+ (mmol/L)			3.3 - 5.1	Basos (absolute)	0.0		x10E3/ $\mu$ L	0.0 - 0.2
LDH (U/L)			135 - 214	Other				
LH (mIU/mL)			1.7 - 8.6	Estradiol (pg/mL)			Men: <12 - 42.6 Boys (1-10 Years): <12 - 20	
PSA (ng/mL)			< 4	Progesterone (ng/mL)			0.20 - 1.4	
Magnesium (mg/dl)			1.8 - 2.5					
Na+ (mmol/L)			133 - 145					
Phosphorus (mg/dL)			2.7 - 4.5					
Total Bilirubin (mg/dL)	0.3		Up to 1.2					
Total Protein (g/dL)	6.3	L	6.4 - 8.3					
T4 ( $\mu$ g/dL)			5.1 - 14.1					
T4, free (ng/dL)			0.93 - 1.7					
T uptake (TBI)			0.8 - 1.3					
T3 (ng/dL)			84.6 - 201.8					
Testosterone (ng/dL)			Men: 280 - 800 Boys: < 1 year 12 - 21 1 - 6 years 3 - 32 7 - 12 years 3 - 68 13 - 17 years 28 - 1110					

Patient	Name:	Gender:	Date of Birth:	Specimen	Collection Time:	Specimen ID:	Provider	Requesting Provider:
	Jonathan Aaron	Male	11/3/1961		10:40 am	10111000598		Michael Richman
	Patient ID #:	Height:	Weight:		Collection Date:	Report Type:		The Center for Cholesterol Management
	10-237-0263				11/9/2010	Complete		1950 Sawtelle Blvd #150
	Fasting Status:	Age:	BMI:		Received Date:	Report Date:		Los Angeles, CA 90025
	Non-Fasting	49			11/10/2010	11/14/2010		Client ID:
								06-90025-18-0000383

Comments:

<p>Decreased HDL cholesterol and increased triglyceride concentrations raise the suspicion of the presence of the metabolic syndrome. According to ATP III guidelines, a diagnosis of metabolic syndrome can be made if 3 or more of the following criteria are met: Triglycerides equal to or great than 150 mg/dl, HDL of equal to or less than 50 mg/dl for women and 40 mg/dl for men, fasting glucose equal to or greater than 110 mg/dl, systolic blood pressure greater than or equal to 130mm HG and diastolic blood pressure equal to or greater than 85 mm HG, and waist circumference for men greater than 40 inches (102 cm) and for women greater than 35 inches (88 cm).</p>
<p>HDL cholesterol, HDL 2 (large HDL particle) cholesterol, and apolipoprotein A-I are decreased or in the intermediate range. Low HDL cholesterol and low apolipoprotein A-I have been associated with increased risk for cardiovascular disease and events. Low amounts of large HDL particles suggest that the HDL present may not be in its most protective form. HDL cholesterol, large HDL particle, and apolipoprotein A-I concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have been demonstrated to increase large HDL particle concentration.</p>
<p>Although LDL cholesterol is optimal or near optimal, small dense LDL cholesterol, %sdLDL, and Apo B are increased or in the intermediate range in this sample, consistent with the presence of small dense LDL particles. Studies have shown that elevated small dense LDL particle concentration is associated with increased risk for coronary heart disease even in the presence of optimal LDL cholesterol values. Small LDL particles may be observed in association with the metabolic syndrome and pre-diabetes. Statins effectively reduce the number of LDL particles, but do not generally influence the size distribution of the LDL particles. Fibrates and niacin have been shown to increase LDL particle size.</p>
<p>Although the LDL cholesterol concentration is optimal, LDL particle concentration is borderline high in this sample. Studies have shown that elevated LDL particle concentration is associated with increased risk for coronary heart disease, even in the presence of optimal LDL cholesterol values. Small LDL particles may be observed in association with the metabolic syndrome and pre-diabetes. Statins effectively reduce the number of LDL particles, but do not generally influence the size distribution of the LDL particles. Niacin, fibrates, and combination therapy (statin +niacin) have been shown to increase LDL particle size.</p>
<p>The Apo B:Apo A-I ratio was increased. Recently large case control studies have demonstrated that the Apo B:Apo A-I ratio is superior to cholesterol measures and cholesterol ratios for predicting risk for myocardial infarction. In the Interheart study, comparing 12,461 myocardial infarction cases to 14,637 age and gender matched controls in 52 countries, the Apo B:Apo A-I ratio was vastly superior to any of the cholesterol parameters measured including the LDL cholesterol: HDL cholesterol ratio and the total cholesterol to HDL cholesterol ratio in all ethnic groups, in both sexes, and at all ages. Decreasing the Apo B:Apo A-I ratio can be achieved by lowering Apo B and/or by increasing Apo A-I. Statins effectively reduce Apo B as do fibrates and niacin. Combination therapy (statin + niacin) is particularly effective at reducing Apo B, especially when small dense LDL particles are present. Apo A-I concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have also been demonstrated to increase Apo A-I.</p>
<p>C-reactive protein is in the intermediate range. CRP is an acute phase reactant. Data from prospective studies indicates that increased concentration of CRP is associated with an increased risk for the development of ischemic cardiovascular events. Consider repeat analysis of CRP in 2-4 weeks to establish baseline value. If CRP remains elevated, then lifestyle changes, including weight reduction, low-fat diet, smoking cessation and regular exercise, should be the initial approach. A diet rich in plant sterols, soy protein, viscous fiber, and almonds has been shown to have CRP-lowering effects comparable to that of lovastatin 20 mg/day. Medications that may lower CRP include statins, fibrates, aspirin, and fish oil. Reducing global CHD risk by aggressive treatment of the traditional risk factors by established therapies may also be beneficial.</p>
<p>Homocysteine is in the intermediate range. Increases in homocysteine concentration can occur with aging, menopause, hypothyroidism, low plasma levels of vitamin cofactors (B6, B12 and folate), certain drugs, and chronic renal insufficiency. Genetic variation in enzymes involved in homocysteine metabolism contributes to inter-individual differences in plasma homocysteine levels.</p>

Patient	Name:	Gender:	Date of Birth:	Specimen	Collection Time:	Specimen ID:	Provider	Requesting Provider:
	Jonathan Aaron	Male	11/3/1961		10:40 am	10111000598		Michael Richman
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	Non-Fasting	49			11/10/2010	11/14/2010		Client ID:
								06-90025-18-0000383

Comments:

<p>Elevated fasting insulin. If a fasting insulin level is elevated, it reflects hyperinsulinemia but fasting levels can be normal when levels following a glucose load are elevated. Insulin is elevated postprandially in proportion to the carbohydrate content in the meal. Elevated fasting insulin levels have been related to atherosclerosis risk. The combination of elevated fasting insulin, apolipoprotein B levels, and small LDL size identifies a very high-risk group for the development of ischemic heart disease.</p>
<p>Vitamin D concentration is in the intermediate range. Decreased vitamin D has been associated with hypertension, inflammation, and the metabolic syndrome. More recently, low serum 25(OH)D has been associated with increased incidence of cardiovascular events and all cause mortality.</p>
<p>Total HDL particle concentration is in the intermediate range in this sample. Decreased HDL particles have been associated with increased risk for cardiovascular disease. HDL particle concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have been demonstrated to increase HDL particle concentration.</p>

End of Report

<p>ATTN PATIENT: Please contact HDL, Inc. at 1-877-4HDLABS (1-877-443-5227) to set an appointment with your personal health coach to discuss your diet and exercise needs at no charge. You can also visit us online at <a href="http://www.myhdl.com">www.myhdl.com</a> and schedule an appointment through our web portal.</p>
--



Joseph P. McConnell, Laboratory Director

CLIA No. 49D1100708 | CAP No. 7224971 | NPI No. 1629209553

INSTRUCTIONS

- 1) Please fill in all of the highlighted areas.
  - 2) Have patient sign Release and Assignment of Benefits below.
  - 3) Copy BOTH sides of patient's insurance card(s).
  - 4) Indicate primary and secondary diagnoses below.
- NOTE: Physicians (or other individuals authorized to order tests) should only order tests that are medically necessary and reasonable.

PATIENT INFORMATION

*Last Name:	First:	Middle Initial:	Client Patient ID #:
Address:			
City:		State:	Zip Code:
Home Phone:		Sex: M. F.	Date of Birth:
Work Phone:		Social Security #:	
*Height: feet inches		*Weight: pounds	

SPECIMEN INFORMATION

Drawing Lab:	Phone:
Collection Date: / /	Time: am/pm
Fasting: <input type="checkbox"/> Yes <input type="checkbox"/> No	Hrs <input type="checkbox"/> No
Initials:	Insulin: time of last dose: am/pm date: / /

HDL, INC. USE ONLY

Received Date: / /	Time: am/pm	Initials:
# Serum SST™ 8.5mL tubes:	# Plasma PPT™ "Pearl" 5mL tubes:	
# Serum "Bumble Bee" 5mL tubes:	# Urine 8mL tubes:	
# Whole Blood EDTA 4mL tubes:	# Other tubes:	

BILLING INFORMATION

☐ INSURANCE: Please attach a copy of both sides of patient's insurance card

☐ MEDICARE No.: Is coverage secondary? ☐ Yes ☐ No

☐ CHECK: Please make check payable to Health Diagnostic Laboratory, Inc.

☐ CREDIT CARD No.: Exp. Date: / /

Name as it appears on card:

RELEASE AND ASSIGNMENT OF BENEFITS

As a courtesy, HDL, Inc. will make every reasonable effort to obtain reimbursement for ordered tests. I authorize HDL, Inc. to release to Medicare or its Carriers & any insurance carrier providing medical benefits to me and any health plan of which I am a member, any medical or other information needed for the claim purposes. I authorize payment of Medical insurance Benefits to the party who bills for this claim & accepts assignment.

Bill to my insurance: I understand that if my insurance company pays me directly for services rendered by HDL, Inc., I am responsible for forwarding such payment to HDL, Inc. I also understand, I am responsible for any deductibles/copayment, as required by my plan.

Important: Insurance regulations require HDL, Inc. to seek payment. I permit a copy of this authorization to be in place of the original.

\*Patient Signature

Date

COMMENTS

REQUESTING PROVIDER/LAB/INSTITUTION

Dr. Friedman  
The Center for Cholesterol Management  
1950 Sawdell Blvd #150  
Los Angeles, CA 90025  
Client ID: 06-90025-12-0000382  
Phone: (310) 451-3936  
Fax: (310) 451-3948

\*Physician or Authorized Signature

Date

CUSTOM PANELS

<input type="checkbox"/> Baseline Assessment	<input checked="" type="checkbox"/> Follow-Up Profile
Lipid Panel	Lipid Panel
Apo A1	Apo A1
Apo B	Apo B
LDL P	LDL P
sdLDL	sdLDL
HDL 2 subclass	HDL 2 subclass
Lp(a) mass w/reflect	Lp(a) mass w/reflect
Apo E Genotype	Lp-PLA2
Factor V Leiden	CRP-hs
Prothrombin Mutation	Fibrinogen
CYP2C9	Homocysteine
Lp-PLA2	FFA (NEFA)
CRP-hs	Insulin
Fibrinogen	NT pro BNP
Homocysteine	Vitamin D
FFA (NEFA)	HEP
Insulin	

INDIVIDUAL TESTS

<b>Routine Panels</b> (see reverse side for details)	<b>Lipoprotein Particles &amp; Apolipoproteins</b>	<b>Genetic Assays</b>
<input type="checkbox"/> Basic Metabolic Panel 80048	<input type="checkbox"/> Apolipoprotein A-I 82172	<input type="checkbox"/> Apo E Genotype
<input type="checkbox"/> Comp Metabolic Panel 80053	<input type="checkbox"/> Apolipoprotein B 82172	<input type="checkbox"/> CYP2C19
<input type="checkbox"/> Electrolyte Panel 80051	<input type="checkbox"/> HDL2 Subclass 82664	<input type="checkbox"/> Factor V Leiden
<input type="checkbox"/> Hepatic Function Panel 80076	<input type="checkbox"/> LDL-P & HDL-P 83704	<input type="checkbox"/> MTHFR
<input type="checkbox"/> Lipid Panel 80061	<input type="checkbox"/> Lp(a) cholesterol 82664	<input type="checkbox"/> Prothrombin Mutation
<input type="checkbox"/> Renal Panel 80069	<input type="checkbox"/> Lp(a) mass 83695	<input type="checkbox"/> Warfarin Sensitivity (CYP2C9 & VKORC1)
<input type="checkbox"/> Thyroid Cascade Panel 84480, 84436, 84439	<input type="checkbox"/> sdLDL 83701	CPT Codes used for all Genetic Assays: 83891, 83892, 83896, 83903, 83908, 83912
<b>Inflammation &amp; Platelets</b>	<b>Metabolic</b>	<b>Thyroid Function</b>
<input type="checkbox"/> AspirinWorks® (urine) 83520 & 82565	<input type="checkbox"/> C-peptide 84681	<input type="checkbox"/> T4 84436
<input type="checkbox"/> F <sub>2</sub> -Isoprostanes 83789, 82570	<input type="checkbox"/> FFA/NEFA 82726	<input type="checkbox"/> TSH 84443
<input type="checkbox"/> Fibrinogen 85384	<input type="checkbox"/> Glucose 82947	<input type="checkbox"/> T3 84480
<input type="checkbox"/> hs-CRP 86141	<input type="checkbox"/> Hemoglobin A1c 83036	<input type="checkbox"/> T4, free 84439
<input type="checkbox"/> Lp-PLA <sub>2</sub> 83698	<input type="checkbox"/> Homocysteine 83090	<input type="checkbox"/> T Uptake 84482
<input type="checkbox"/> MPO 83516	<input type="checkbox"/> Insulin 83525	
<input type="checkbox"/> NT-proBNP 83880	<input type="checkbox"/> RBC Folate 82747, 85014	
(may require additional ICD-9 coding)	<input type="checkbox"/> TSH 84443	
<b>Hormones</b>	<input type="checkbox"/> Uric Acid 84550	
<input type="checkbox"/> FSH 83001	<input type="checkbox"/> Vitamin B <sub>12</sub> 82607	
<input type="checkbox"/> LH 83002	<input type="checkbox"/> 25-hydroxy-Vitamin D 82652	
<input type="checkbox"/> Testosterone 84403	<b>Renal</b>	
	<input type="checkbox"/> Creatinine (serum) 82565	
	<input type="checkbox"/> Cystatin C 82610	
	<input type="checkbox"/> Microalbumin (urine) 82043	

DIAGNOSIS (ICD-9 CODE)

NOTE: The below ICD-9 codes are listed as a convenience. Ordering physicians should report the code that best describes the reason for ordering the tests & are not required to use the codes provided below.

Please write in Diagnosis Codes below or clearly mark to the right. PLEASE USE FOURTH & FIFTH DIGIT MODIFIERS.

Primary:

Secondary:

Other:

<input type="checkbox"/> Diabetes 250.	<input type="checkbox"/> Hypertension w/Renal Disease & Heart Failure 404.
<input checked="" type="checkbox"/> Pure Hypercholesterolemia 272.0	<input type="checkbox"/> Intermediate Coronary Syndrome 411.1
<input type="checkbox"/> Pure Hyperglyceridemia 272.1	<input type="checkbox"/> Coronary Atherosclerosis Unspecified Vessel, Native or Graft 414.00
<input type="checkbox"/> Mixed Hyperlipidemia 272.2	<input type="checkbox"/> Congestive Heart Failure 428.
<input type="checkbox"/> Unspecified Hyperlipidemia 272.4	<input type="checkbox"/> Cirrhosis of liver without mention of alcohol 571.5
<input type="checkbox"/> Hypertension 401.	<input type="checkbox"/> Other chronic non-alcoholic liver disease 571.8
<input type="checkbox"/> Hypertensive Heart Disease 402.	<input type="checkbox"/> Unspecified chronic liver disease without mention of alcohol 571.9
<input type="checkbox"/> Hypertension w/Renal Disease 403.	<input type="checkbox"/> OTHER



The NMR LipoProfile® test may be covered by one or more issued or pending patents, including U.S. Patent Nos. 5,343,389; 6,518,069; 6,576,471; 6,653,140; and 7,243,030. CLIA: 34D0952253



**LIPOSCIENCE**

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2500 Sumner Boulevard  
Raleigh, NC 27616  
877-547-6837  
www.liposcience.com

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Patient Name		Sex	Age	Clinician	
Aaron, Jonathan		U	48		
Patient ID	Birth Date	Accession Number			
10082500324	11/03/1961	W0706502			
Client Name and Address					
HDL-Health Diagnostic Lab Inc 17730/ 737 North 5th Street Suite 103 Richmond, VA 23219 Phone: (804)343-2718 Fax: (804)343-0929					
Date Collected	Date Received	Report Date and Time		Requisition Number	Fasting Status
08/24/2010	08/26/2010	08/27/2010 07:14		10082500324	Not Specified

**NMR LipoProfile® test**

**Reference Range<sup>1</sup>**

	Percentile <sup>1</sup>	20th	50th	80th	95th	
	nmol/L	Low	Moderate	Borderline-High	High	Very High
<b>LDL-P</b> (LDL Particle Number)	<b>1764</b>	<b>&lt; 1000</b>	<b>1000-1299</b>	<b>1300-1599</b>	<b>1600-2000</b>	<b>&gt; 2000</b>

**Historical Reporting**

**LDL-P** 400 600 700 800 900 1000 1100 1200 1300 1400 1500 1600 1700 1800 1900 2000 4000 6000

08/24/10 (1764)

**LDL-C** 20 40 60 70 80 90 100 110 120 130 140 150 160 170 180 190 200 400

↑ LDL-P ~ 1764  
↑ Apo B - 119

↓ HDL-P ~ 23.2  
↑ TG - 571  
↑ VLDL-P - 24.9  
vitamin D ~ 12  
insulin 57 uU/mL

*[Handwritten signature]*

1. Reference population comprises 5,362 men and women not on lipid medication enrolled in the Multi-Ethnic Study of Atherosclerosis (MESA). MESA is a National Heart, Lung, and Blood Institute (NHLBI) study.

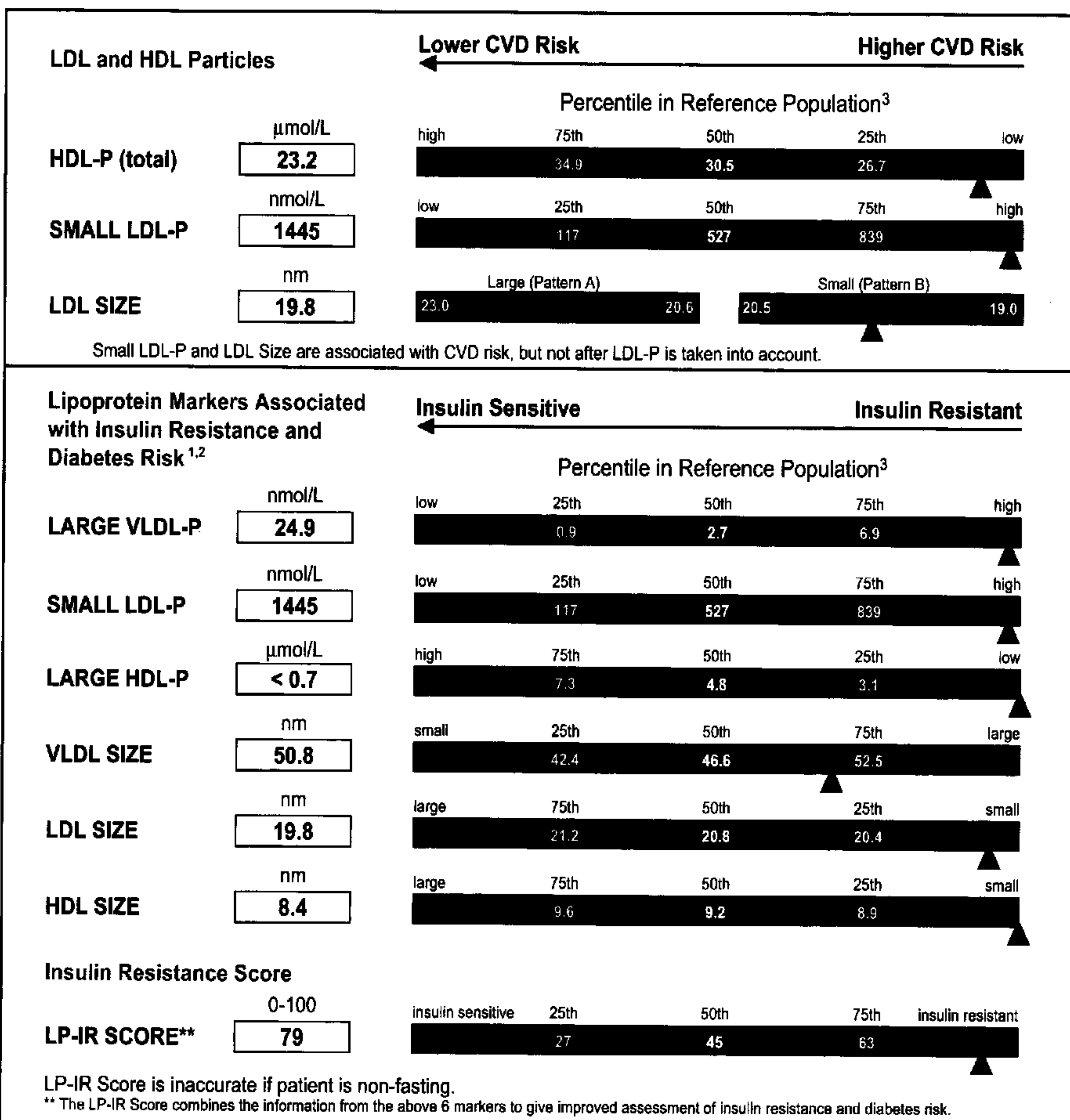
*[Handwritten notes and signatures]*  
① LipoScience 10mg  
② LipoScience 10mg  
③ LipoScience 135mg



Page 1 of 1

Patient Name			Sex	Age	Clinician	
Aaron, Jonathan			U	48		
Patient ID			Birth Date		Accession Number	
10082500324			11/03/1961		W0706502	
Client Name and Address						
HDL-Health Diagnostic Lab Inc 17730/ 737 North 5th Street Suite 103 Richmond, VA 23219 Phone: (804)343-2718 Fax: (804)343-0929						
Date Collected		Date Received		Report Date and Time		Fasting Status
08/24/2010		08/26/2010		08/27/2010 07:14		Not Specified
Requisition Number						
10082500324						

## PARTICLE CONCENTRATION AND SIZE



These laboratory assays, validated by LipoScience, have not been cleared by the US Food and Drug Administration. The clinical utility of these laboratory values has not been fully established.

Requesting Physician:

**Michael Richman**

The Center for Cholesterol

1950 Sawtelle Blvd #150

Los Angeles, CA 90025

Client Number:

06-90025-18-0000383

Patient Name:

**Jonathan Aaron**

Patient ID:

10-237-0263

Specimen ID:

10082500324

Fasting Status:

Unknown

Report Type:

Complete

Birth Date:

11/3/1961

Height:

Weight (pounds):

Report Date:

Gender:

**Male**

BMI:

Report Date:

8/31/2010

Age:

**48**

Collection Date: Collection Time:

8/24/2010 9:25 am

Received Date:

8/25/2010

## Laboratory Results

### NCEP ATP III Lipid Tests

For descriptions of results, see reverse side of this document (or separate sheet if faxed).

	At Risk	Intermediate	Optimal	High Risk	Intermediate Risk	NCEP ATP-III Goal	Previous Result
TCHOL (mg/dL)	263			≥ 240	200 - 239	< 200	
HDL-C (mg/dL)	32			< 40		≥ 40	
LDL-C (mg/dL)			92	≥ 130 CHD & CHD risk eq. > 100	100 - 129 CHD & CHD risk eq. 70 - 100	< 100 CHD & CHD risk eq. < 70	
TRIG (mg/dL)	571			≥ 200	150 - 199	< 150	

### Advanced Cardiovascular Risk

	At Risk	Intermediate	Optimal	High Risk	Intermediate Risk	Optimal Value	Previous Result
Apo B (mg/dL)	119			≥ 90 CHD & CHD risk eq. > 80	60 - 89 CHD & CHD risk eq. 55 - 79	≤ 60 CHD & CHD risk eq. < 55	
LDL-P (nmol/L)	1764			≥ 1300	1000 - 1299	< 1000	
sdLDL (mg/dl) *	59			≥ 31	21 - 30	≤ 20	
% small dense LDL	65			≥ 23	14 - 22	≤ 13	
Apo A1 (mg/dL)		123		< 114	114 - 131	≥ 132	
HDL-P (μmol/L)	23.2			< 28	28 - 34	≥ 35	
HDL2 (mg/dL) *			12	≤ 8	9 - 11	≥ 12	
Apo B:Apo A1 Ratio	0.97			≥ 0.81	0.61 - 0.81	≤ 0.6	
Lp(a) mass (mg/dL)			2	≥ 30		< 30	
Lp(a) Cholesterol (mg/dL)				≥ 6	3 - 5	< 2	
Apolipoprotein E Genotype *			3/3	Estimated Genotype Frequency in Humans: 2/2 (~1-2%), 2/3 (~15%), 2/4 (~1-2%), 3/3 (~55%), 3/4 (~25%), 4/4 (~1-2%)			
Factor V leiden *			Arg/Arg	Optimal=Non-carrier (Arg/Arg); At Risk (Arg/Gln or Gln/Gln)			
Prothrombin Mutation *			G/G	Optimal=Non-carrier (G/G); At Risk (G/A or A/A)			
CYP2C19			*1/*1	Estimated Genotype frequency: *1/*1 (~60-70%), *1/*2 (~20-30%), *2/*2 (~2-3%), *1/*3 (<1%), *2/*3 (<0.05%), *3/*3 (<0.01%)			
Lp-PLA2 (ng/mL)		204		> 235	200 - 235	< 200	
hsCRP (mg/L)		2.06		≥ 3.00	1.00 - 2.99	< 1.00	
Fibrinogen (mg/dL)			388	≥ 465	391 - 464	≤ 390	
Homocysteine (μmol/L)		12		> 13	11 - 13	≤ 10	
NTproBNP (pg/mL)			19	≥ 450	125 - 449	< 125	
Insulin (μU/mL)	57			≥ 12	10 - 11	3 - 9	
Free Fatty Acid (mmol/L)			0.50	> 0.7	0.6 - 0.7	≤ 0.59	
Vitamin D (ng/mL) (25 hydroxy)	12			≤ 14	15 - 29	30 - 100	
Aspirinworks				> 1500		≤ 1500	

### Lab notes:

\*This test was developed and its performance characteristics determined by HDL. It has not been cleared or approved by the U.S. Food & Drug Administration (FDA). The FDA has determined that such clearance or approval is not necessary. This test is used for clinical purposes. It should not be regarded as investigational or for research. This laboratory is certified under CLIA-88 as qualified to perform high complexity clinical laboratory testing.

To schedule time with a Personal Health Coach, please call 1-877-4HDLABS (1-877-443-5227)



<b>Requesting Physician:</b>	<b>Pat Name:</b>	<b>Birth Date:</b>	<b>Sex:</b>	<b>Age:</b>	
<b>Michael Richman</b>	<b>Jonatnan Aaron</b>	<b>11/3/1961</b>	<b>Male</b>	<b>48</b>	
The Center for Cholesterol	<b>Patient ID:</b>	<b>Fasting Status:</b>	<b>Height:</b>	<b>BMI:</b>	<b>Collection Date: Collection Time:</b>
1950 Sawtelle Blvd #150	<b>10-237-0263</b>	<b>Unknown</b>			<b>8/24/2010 9:25 am</b>
Los Angeles, CA 90025	<b>Specimen ID:</b>	<b>Report Type:</b>	<b>Weight (pounds):</b>	<b>Report Date:</b>	<b>Received Date:</b>
<b>Client Number:</b>	<b>10082500324</b>	<b>Complete</b>		<b>8/31/2010</b>	<b>8/25/2010</b>
<b>06-90025-18-0000383</b>					

**Comments:**

Decreased HDL cholesterol and increased triglyceride concentrations raise the suspicion of the presence of the metabolic syndrome. According to ATP III guidelines, a diagnosis of metabolic syndrome can be made if 3 or more of the following criteria are met: Triglycerides equal to or great than 150 mg/dl, HDL of equal to or less than 50 mg/dl for women and 40 mg/dl for men, fasting glucose equal to or greater than 110 mg/dl, systolic blood pressure greater than or equal to 130mm HG and diastolic blood pressure equal to or greater than 85 mm HG, and waist circumference for men greater than 40 inches (102 cm) and for women greater than 35 inches (88 cm).

Markedly increased Triglycerides >500 mg/dL). Secondary causes of hypertriglyceridemia (thyroid disease, diabetes, alcohol intake, drug interactions, etc) should be investigated.

Although LDL cholesterol is optimal or near optimal, small dense LDL cholesterol, %sdLDL, and Apo B are increased or in the intermediate range in this sample, consistent with the presence of small dense LDL particles. Studies have shown that elevated small dense LDL particle concentration is associated with increased risk for coronary heart disease even in the presence of optimal LDL cholesterol values. Small LDL particles may be observed in association with the metabolic syndrome and pre-diabetes. Statins effectively reduce the number of LDL particles, but do not generally influence the size distribution of the LDL particles. Fibrates and niacin have been shown to increase LDL particle size.

Although the LDL cholesterol concentration is optimal, LDL particle concentration is increased in this sample. Studies have shown that elevated LDL particle concentration is associated with increased risk for coronary heart disease, even in the presence of optimal LDL cholesterol values. Small LDL particles may be observed in association with the metabolic syndrome and pre-diabetes. Statins effectively reduce the number of LDL particles, but do not generally influence the size distribution of the LDL particles. Niacin, fibrates, and combination therapy (statin +niacin) have been shown to increase LDL particle size.

The apoB:apoA1 ratio was increased. Recently large case control studies have demonstrated that the apoB:apoA1 ratio is superior to cholesterol measures and cholesterol ratios for predicting risk for myocardial infarction. In the Interheart study, comparing 12,461 myocardial infarction cases to 14,637 age and gender matched controls in 52 countries, the apoB:apoA1 ratio was vastly superior to any of the cholesterol parameters measured including the LDL cholesterol: HDL cholesterol ratio and the total cholesterol to HDL cholesterol ratio in all ethnic groups, in both sexes, and at all ages. Decreasing the apoB:apoA1 ratio can be achieved by lowering apoB and/or by increasing apoA1. Statins effectively reduce apoB as do fibrates and niacin. Combination therapy (statin + niacin) is particularly effective at reducing apoB, especially when small dense LDL particles are present. ApoA1 concentration may be increased by exercise, fish oil, or alcohol consumption in moderation. Niacin, fibric acids, and combination therapy (statin + niacin) have also been demonstrated to increase apoA1.

C-reactive protein is in the intermediate range. CRP is an acute phase reactant. Data from prospective studies indicates that increased concentration of CRP is associated with an increased risk for the development of ischemic cardiovascular events. Consider repeat analysis of CRP in 2-4 weeks to establish baseline value. If CRP remains elevated, then lifestyle changes, including weight reduction, low-fat diet, smoking cessation and regular exercise, should be the initial approach. A diet rich in plant sterols, soy protein, viscous fiber, and almonds has been shown to have CRP-lowering effects comparable to that of lovastatin 20 mg/day. Medications that may lower CRP include statins, fibrates, aspirin, and fish oil. Reducing global CHD risk by aggressive treatment of the traditional risk factors by established therapies may also be beneficial.

Lp-PLA2 is increased in this sample. Lp-PLA2 is an inflammatory risk marker that, unlike hsCRP, is not an acute phase reactant. It is produced by macrophages and is a marker of vascular specific inflammation. Lp-PLA2 circulates in the plasma primarily bound to LDL particles. High plasma Lp-PLA2 is associated with increased risk for cardiovascular disease and events (myocardial infarction and stroke). Increased values have also been associated with endothelial dysfunction and peripheral arterial disease. Lp-PLA2 is the only test that is FDA approved to assess risk for stroke. Patients in the upper tertile for both CRP and Lp-PLA2 are at highest risk. In the Atherosclerosis risks in communities (ARIC) study, patients with both CRP and Lp\_PLA2 in the upper tertile of the population had 5X increased risk for myocardial infarction and 11X increased risk for stroke. Statins, fibric acids, and niacin have been shown to have Lp-PLA2 lowering effects.

Homocysteine is in the intermediate range. Increases in homocysteine concentration can occur with aging, menopause, hypothyroidism, low plasma levels of vitamin cofactors (B6, B12 and folate), certain drugs, and chronic renal insufficiency. Genetic variation in enzymes involved in homocysteine metabolism contributes to inter-individual differences in plasma homocysteine levels.



# HealthDiagnosticLaboratory, Inc.

beyond disease diag

**Health Diagnostic Laboratory, Inc.**  
737 N. 5th St., Suite 103  
Richmond, VA 23219  
Phone: (804) 343-2718  
Fax: (804) 343-2704

**Joseph P. McConnett, Ph.D.**  
Laboratory Director  
CLIA No. 49D1100708  
CAP No. 7224971  
NPI No. 1629209853

0000383-0050107

Last	First	Last
0000383-0050107	0000383-0050107	0000383-0050107
Last	First	Last
0000383-0050107	0000383-0050107	0000383-0050107

Last Name (Please Print)		First	Middle Initial	Client Patient ID No.
Agton, Jonathan				
Address				
7817 S. Bradford St.				
City	State	Zip Code		
LA	CA	90035		
Home Phone	Sex	Date of Birth		
(310) 497-7693	M F	11/13/16		
Work Phone	SSN			
( )				
Height	Weight			
feet inches	pounds			

**Dr. Richman**  
The Center for Cholesterol Management  
1950 Sawtelle Blvd #150  
Los Angeles, CA 90025

Client ID: 06-90025-18-0000383  
Phone: (310) 481-3939  
Fax: (310) 481-3949

Physician or Authorized Signature		Date
[Signature]		11/13/16
Send Addl. Copy to:	Fax: ( )	

Drawing Lab	Phone
	( )
Collection Date	Time
8/24/10	9:25 am / pm
Initials	
SR	

Received into Lab:	<input type="checkbox"/> WB 9mL EDTA / # tubes <input type="checkbox"/> Serum / # tubes <input type="checkbox"/> Plasma / # tubes <input type="checkbox"/> Urine / # tubes
Date	
Time	am / pm
Initials	

As a courtesy, HDL, Inc. will make every reasonable effort to obtain reimbursement for ordered tests. I authorize HDL, Inc. to release to Medicare or its Carriers and any insurance carrier providing medical benefits to me and any health plan of which I am a member, any medical or other information needed for the claim purposes. I authorize payment of Medical insurance Benefits to the party who bills for this claim and accepts assignment.

**Bill to my insurance:** I understand that if my insurance company pays me directly for services rendered by HDL, Inc., I am responsible for forwarding such payment to HDL, Inc. I also understand, I am responsible for any deductibles/co-payment, as required by my plan.

**Important:** Insurance regulations require HDL, Inc. to seek payment. I permit a copy of this authorization to be in place of the original.

Patient's Signature	Date
[Signature]	8-24-10

• Have patient sign Release of Benefits above & Copy BOTH sides of patient's insurance card(s).  
• Indicate primary and secondary diagnoses below.

**NOTE: Physicians (or other individuals authorized to order tests should only order tests that are medically necessary and reasonable.**

Does patient have Medicare coverage? ☐ YES ☐ NO

Is the Medicare coverage secondary? ☐ YES ☐ NO

Medicare No.: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

☐ CHECK: Please make check payable to Health Diagnostic Laboratory, Inc.

Name as it appears on card: \_\_\_\_\_

Credit card number: \_\_\_\_\_ Exp. Date: \_\_\_\_/\_\_\_\_/\_\_\_\_

Card holder Signature: \_\_\_\_\_

Total Amount: \$ \_\_\_\_\_ CREDIT CARD TYPE: ☐ Visa ☐ MC ☐ Amex

## COMPREHENSIVE CVD BASELINE ASSESSMENT

Lipid Panel	FFA (NEFA)
Apo A1	Insulin
Apo B	NT pro BNP
LDL P	Vitamin D
sdLDL	HbA1c
HDL 2 subtypes	CRP
Lp(a) mass w/refer to Lp(a)CHOL	
Apo E Genotype	
Factor V Leiden	
Prothrombin G20210A Mutation	
CYP2C9	
Lp-PLA2	
CRP-hs	
Homocysteine	

## CVD/METABOLIC FOLLOW-UP PROFILE

Lipid Panel	HEP (Hepatic Function Panel)
Apo A1	
Apo B	
LDL P	
sdLDL	
HDL 2 subtypes	
Lp(a) mass w/refer to Lp(a)CHOL	
Lp-PLA2	
CRP-hs	
Fibrinogen	
Homocysteine	
FFA (NEFA)	
Insulin	
NT pro BNP	

## Genetic Assays

<input type="checkbox"/> Apo E Genotype	83891, 83892, 83896, 83903, 83908, 83912
<input type="checkbox"/> Warfarin Sensitivity (CYP2C9*2, *3, & VKORC1)	83891, 83892, 83896, 83903, 83908, 83912
<input type="checkbox"/> Factor V Leiden	83891, 83892, 83896, 83903, 83908, 83912
<input type="checkbox"/> Prothrombin G20210A Mutation	83891, 83892, 83896, 83903, 83908, 83912

## Routine Panels (see reverse side for details)

<input type="checkbox"/> Basic Metabolic Panel	80048
<input type="checkbox"/> Comp Metabolic Panel	80053
<input type="checkbox"/> Electrolyte Panel	80051
<input type="checkbox"/> Hepatic Function Panel	80076
<input type="checkbox"/> Renal Panel	80069
<input type="checkbox"/> Lipid Panel	80061
<input type="checkbox"/> Thyroid Cascade Panel	84480, 84436, 84439

## Miscellaneous Tests

<input type="checkbox"/> Vitamin D (25hydroxy)	82652
<input type="checkbox"/> Insulin	83525
<input type="checkbox"/> Aspirin-works (urine)	83520 & 82565
<input type="checkbox"/> Creatine Kinase	82550
<input type="checkbox"/> Cystatin C	82610
<input type="checkbox"/> y-GT	82977
<input type="checkbox"/> D-Dimer	85362
<input type="checkbox"/> Fibrinogen	85384
<input type="checkbox"/> Micro-albumin (urine)	82043
<input type="checkbox"/> Homocysteine	83090
<input type="checkbox"/> Iron & TIBC	83540
<input type="checkbox"/> PSA, total	84153
<input type="checkbox"/> AFP	82105
<input type="checkbox"/> CEA	82378

## Miscellaneous Lipids

<input type="checkbox"/> sdLDL	83701
<input type="checkbox"/> LDL-C Direct	83721
<input type="checkbox"/> Apolipoprotein B	82172
<input type="checkbox"/> CRP-hs	86141
<input type="checkbox"/> Lp(a) mass	83695
<input type="checkbox"/> Lp(a)-CHOL	82664
<input type="checkbox"/> Lp-PLA2	83698
<input type="checkbox"/> Apolipoprotein A1	82172
<input type="checkbox"/> NT pro BNP *	83880
<input type="checkbox"/> HDL2 Subclass	82664
<input type="checkbox"/> Triglycerides	84478
<input type="checkbox"/> HDL-Direct	83718
<input type="checkbox"/> FFA (NEFA)	82726
<input type="checkbox"/> LDL-P by NMR	83704

\*NT pro BNP may require additional CVD coding

## Hormones

<input type="checkbox"/> FSH	83001
<input type="checkbox"/> Testosterone	84403
<input type="checkbox"/> LH	83002

## Metabolic Syndrome

<input type="checkbox"/> Glucose	82947
<input type="checkbox"/> C-peptide	84681
<input type="checkbox"/> Insulin	83525
<input type="checkbox"/> Hemoglobin A1c	83036
<input type="checkbox"/> FFA (NEFA)	82726

## Thyroid Function

<input type="checkbox"/> T4	84436
<input type="checkbox"/> TSH	84443
<input type="checkbox"/> T3	84480
<input type="checkbox"/> T4, free	84439
<input type="checkbox"/> T Uptake	84482
<input type="checkbox"/> Other	
<input type="checkbox"/> Other	

NOTE: The below ICD-9 codes are listed as a convenience. Ordering physicians should report the code that best describes the reason for ordering the test and are not required to use the codes provided below.

Please write in Diagnosis Codes below or clearly mark to the right. PLEASE USE FOURTH & FIFTH DIGIT MODIFIERS.	<input type="checkbox"/> Diabetes	250.	<input type="checkbox"/> Hypertension w/Renal Disease & Heart Failure	404.
	<input type="checkbox"/> Pure Hypercholesterolemia	272.0	<input type="checkbox"/> Intermediate Coronary Syndrome	411.1
	<input checked="" type="checkbox"/> Pure Hyperglyceridemia	272.1	<input type="checkbox"/> Coronary Atherosclerosis Unspecified Vessel, Native or Graft	414.00
	<input type="checkbox"/> Mixed Hyperlipidemia	272.2	<input type="checkbox"/> Congestive Heart Failure	428.
PRIMARY:	<input type="checkbox"/> Unspecified Hyperlipidemia	272.4	<input type="checkbox"/> Cirrhosis of liver without mention of alcohol	571.5
	<input type="checkbox"/> Hypertension	401.	<input type="checkbox"/> Other chronic non-alcoholic liver disease	571.8
	<input type="checkbox"/> Hypertensive Heart Disease	402.	<input type="checkbox"/> Unspecified chronic liver disease without mention of alcohol	571.9
	<input type="checkbox"/> Hypertension w/Renal Disease	403.	<input type="checkbox"/> OTHER	



The Center for Cholesterol Management

A Medical Corporation

Nutrition and Exercise History

Name: Jonathan Aaron Verified? ✓ Physician Richman Discussed results? ✓

Date: 10.7.10 E-mail address: jmaaron@me.com

Questions regarding labs? —

Height: 6'2" Weight: 190s now 193 Desired Weight: 180 Weight Hx February ↓ 20 on Clean Diet

Exercise? Y/N Average # of days/week — Average # minutes/session — Type: —

BEE — Calorie needs — previously swimming & now.

Circle any family history if any:

Diabetes - Heart Disease - High Blood Pressure - Stroke

What do you hope to gain from meeting with an RD: eat better / med's all new concerned.

Post Menopausal? Y/N If so, since? — On hormone replacement? Y/N Type: —

Smoke: Y/N # of cigarettes a day? 1 pack/day Previously smoked? Y/N (quit in plan - time line? date: NYR?)

Medications/Supplements: —

Allergies: —

DIET HISTORY: Typical Day lactose intolerant?

- 8/9
1. COFFEE whole milk, sugar - cheese danish sometimes
  2. 12/2:00 Forget or Subway tuna 12", turkey mayo/lettuce, cheese vit H2O, coke (occ), baked chips \* not a snack person - 1+2 week/choc. alc 2x/month.
  3. Last night Burrito/lettuce, cheese, gn. beef, rice 1x/wk Soup plantation No water, NO ~~water~~ FISH 2-3x/week red meat Root beer.. breaded chicken, cesar sal. chicken. ⇒





THE CENTER FOR CHOLESTEROL MANAGEMENT

A Medical Corporation  
1950 Sawtelle Blvd. Suite 150  
Los Angeles, CA 90025

NAME Jonathon Aaron

CHART NUMBER \_\_\_\_\_

RD CONSULT-SEE NUTRITION

PROGRESS NOTES 10.7.10

D. ADAIR, M.S., R.D.

last

① D/C Fish oil Th./Fri. GI symptoms, but may have

② b/ct. healthy food, w/ also been flu?

"TREAT" Foods 2x/week (Pizza, Steak, tuna w/ mayo, pasta red cream sauce chicken, etc.)

③ Start swimming 2-3x/week, interval for 30 min.

④ Smoking cess.

⑤ high fiber cereal in am.

⑥ Email sent 10.7., copy in chart.

RD CONSULT-SEE NUTRITION

PROGRESS NOTES 4-6 weeks

D. ADAIR, M.S., R.D. 11.18.10

① Wt. 193 SELF report ② Not obese as much as discussed ③ LDL ↓ ④ INS ~~111~~ (11.57) counseled on importance of exercise re: INS, \*HDL-C, vit<sup>D</sup> improved, HDL-2 ↑ → Trilipix ⑤ Avoid eating refined carbs + any carbs alone.

## **Dominique Adair**

---

**From:** Dominique Adair  
**Sent:** Thursday, October 07, 2010 1:51 PM  
**To:** 'jmaaron@me.com'  
**Subject:** Appointment Summary  
**Attachments:** My Guide to Heart Healthy Eating.pdf

Very nice to meet you today. You had me a little worried when you said "do you really want to know..." but in fact I believe you have a good awareness of eating for your health, and now it is a matter of fine tuning and making sure that your choices match your understanding.

Here is a brief summary of you plan.

- You have stopped taking the Omega 3 supplement because of GI upset and a generally unpleasant reaction. Would you consider taking ½ of the original dose (2), and let me know how that goes? You don't like fish, so the only other way I can increase your Omega 3s is with flax seeds, flaxseed oil, hemp – food form like hemp milk, almonds, tofu and other soy products and walnuts among a few other others, but it would be hard for you to eat the amount that you need to produce health benefits like reduce inflammation, keep your blood from clotting excessively, improve your lipids etc. So, let me know if you're willing to try ½ dose and we'll take it from there.
- I'd like you to change your breakfast – adding one cup of high fiber cereal (Total, Raisin Bran, Cheerios, oatmeal) with 1% milk and a serving of fruit. Coffee is fine, Danishes, muffins, pastries are not.
- I'd like you to try to have lunch before 2:00. That way you will spread your nutrients throughout the day which is generally better for health, and be ready to eat a healthy dinner around 5 – 7:00.
- Many of your food choices are fine (chicken Cesar, bean/beef burrito, turkey sandwich, etc.) but there are some which should be limited (pizza, tuna salad sandwich with mayo, creamy red sauce with pasta and chicken, chicken parmesan, red meats in large portions, etc.). I'd like you to try to limit these types of foods to 2x/week. If you're not sure, email me to ask.
- Limit alcohol, as per your record, 2x/month.
- Limit coke/soda to 1/week, vitamin water OK for now.
- Try to convert all complex carbohydrates to the whole, less refined version (i.e. brown rice, whole wheat bread, whole wheat pasta, etc.).
- Try to eat AT LEAST 3 servings of vegetables and 2 serving of fruit per day – Yes, I just threw this one in.
- Begin swimming for 30 minutes (interval format) 2-3 times a week. If you can't do 3/week, don't give up on the whole idea.

I looked more closely at the Clean Diet website, and while some of principles are for the most sound, I don't think you will eat this way long term. Our project is to come up with a lifestyle program that you can live with, then, if you want to do the 21 days program once a year, that's OK.

I also attached a guide to heart healthy eating which you might like to share with your wife. It is pretty straightforward, and might help answer any of the questions we didn't go over.

Please email if you have any questions. Same for your wife.





THE CENTER FOR CHOLESTEROL MANAGEMENT  
A Medical Corporation  
1950 Sawtelle Blvd, Suite 150  
Los Angeles, CA 90025

\*\*\*Please complete all pages of this form\*\*\*

NAME: JONATHAN AARON DATE: 8.24.10  
SEX: XM   F DOB: 11/31/61 SSN: 040-54-7095 DL#: A1239139  
ADDRESS: 1817 S. BEDFORD ST.  
CITY: CA STATE: CA ZIP: 90035  
FAX:    EMAIL: JMAARON@CME PHONE: 310-497-7693  
EMERGENCY CONTACT: Michelle AARON <sup>LOW</sup> PHONE: 310-968-2726  
ADDRESS: SAME  
CITY:    STATE:    ZIP:     
EMPLOYER: Temple Emanuel of BH PHONE: 310-288-3762  
ADDRESS: 8844 BURTON WAY CITY: BH STATE: CA ZIP: 90211

Please list all of your medications, include non-prescription drugs, dietary supplements, and vitamins.

NAME OF DRUG:	DOSE:	No. TIMES DAILY:
---------------	-------	------------------


Have you ever been diagnosed with?

High Blood Pressure	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No	How long ago? <u>  </u>
Diabetes	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No	How long ago? <u>  </u>
Stroke	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No	When did it occur? <u>  </u>
High Cholesterol	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	What medications do you take for this, if any? <u>DID TAKE - DONT REMEMBER</u>
Lung Disease	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No	What type? <u>  </u>

Heart Disease ☐ Yes ☒ No

How long ago? \_\_\_\_\_

Other Vascular Disease ☐ Yes ☒ No

How long ago? \_\_\_\_\_

List other medical problems you have had. These would include problems for which you have taken medications or been hospitalized. Please include the dates these problems occurred.

98 - Surgery for Back - Microdissection

99 - Back Surgery

01 - " "

Are you allergic to any medications? ☐ Yes ☒ No

List these medications? \_\_\_\_\_

Are you allergic to X-Ray dye? ☐ Yes ☒ No

List all surgeries, both major and minor, you have had:

SURGERY

DATE

HOSPITAL

Have you ever smoked? ☒ Yes ☐ No

How many cigarettes per day? pack

How long (have) did you smoke (d)? 28 yrs

If you quit, when did you quit? \_\_\_\_\_

How many glasses per week do you consume of? WINE        BEER        COCKTAILS 1/4

Has anyone in your family had any of the following illnesses?

	WHICH FAMILY MEMBER	HOW OLD WERE THEY
Cancer	<u>Grandfather</u>	<u>50 s</u>
Heart Attack	<u>Grandfather</u>	<u>60 s</u>
Angina or clogged arteries		

Sudden death \_\_\_\_\_

Hypertension \_\_\_\_\_

Other heart disease \_\_\_\_\_

High cholesterol \_\_\_\_\_

Stroke \_\_\_\_\_

Diabetes \_\_\_\_\_

Are you having or have you ever had? (check all for which the answer is "yes").

- |   |  |
|---|--|
| <input type="checkbox"/> Increasing Breathlessness With Your Usual Activities                                 | <input type="checkbox"/> Recent Cough              |
| <input type="checkbox"/> Unexpected weight gain of more than 5 lbs in the last weeks or months                |  |
| <input checked="" type="checkbox"/> Pain, pressure/discomfort in the chest                                    | <input type="checkbox"/> Passed (ing) out-fainting |
| <input type="checkbox"/> Shortness of breath at rest, laying down   | <input type="checkbox"/> worsening fatigue         |
| <input type="checkbox"/> Any neck, jaw, left arm discomfort   | <input type="checkbox"/> Swelling of the ankles    |
| <input type="checkbox"/> Pain or cramps in leg(s) with walking  | <input type="checkbox"/> Dizzy spells              |
| <input type="checkbox"/> A stroke or temporary stroke   | <input type="checkbox"/> Heart murmur              |
| <input type="checkbox"/> Spells of rapid irregular heartbeat  | <input type="checkbox"/> Heart attack              |
| <input type="checkbox"/> Urination at night   | <input type="checkbox"/> Rheumatic fever           |
| <input type="checkbox"/> Abnormal EKG   | <input type="checkbox"/> Varicose veins            |
| <input type="checkbox"/> Have you ever been hospitalized for your heart, or what they thought was your heart? |  |
| <input type="checkbox"/> Any other cardiac diagnosis? _____   |  |

☐ Any tests done for your heart? What tests? \_\_\_\_\_

When where they done? \_\_\_\_\_

After any problems you wish to address at this visit?

Patient name (sign)

Date

Witness

Date

## INSURANCE INFORMATION

Please provide us with your medical insurance information:

### PRIMARY INSURANCE POLICY:

Company: BLUE SHIELD OF CAL Phone: 888-552-5345

Policy #: XECJO4204767 Group: SP2277

Name and SS# of Insured: JONATHAN ADAM - 040547095

### SECONDARY INSURANCE POLICY:

Company: \_\_\_\_\_ Phone: \_\_\_\_\_

Policy #: \_\_\_\_\_  
Group: \_\_\_\_\_

Name and SS# of Insured: \_\_\_\_\_

### OTHER INSURANCE:

company: \_\_\_\_\_ Phone: \_\_\_\_\_

Policy #: \_\_\_\_\_ Group: \_\_\_\_\_

Name and SS# of Insured: \_\_\_\_\_

## ASSIGNMENT BENEFITS

I HEREBY ASSIGN TO MICHAEL RICHMAN M.D., MY RIGHT TO AND INTEREST IN ANY AND ALL HEALTH CARE AND /OR SURGICAL BENEFITS, OTHERWISE PAYABLE TO ME , FOR MEDICAL AND/OR SURGICAL TREATMENT RENDERED BY ANY OF THE ASSIGNEES. I HEREBY DIRECT MY INSURANCE COMPANY TO MAKE PAYMENTS DDIRECTLY TO THE ASSIGNEE AT 1900 SAWTELLE BLVD # 145A LOS ANGELES, CA 90025.  
IN UNDERSTAND THAT I AM RESPONSIBLE FOR ANY CHARGES NOT PAID BY MY INSURANCE COMPANY(DZS), UNLESS SUCH CHARGES ARE LIMITED BY EXISTING CONTRACT AGREEMENTS BETWEEN THE ASSIGNEE AN MY MEDICAL CARRIER, AND THAT FINANCE CHARGES WDL BE ADDED TO ANY OUTSTANDING BALANCE, STARTING THIRTY DAYS FROM THE DATE A BILL IS SUBMITTED TO MY INSURANCE COMPANY, OR FROM THE DATE OF MY FIRST STATEMENT, IF CHARGES ARE NOT COVERED BY MY INSURANCE COMPANY, I AUTHORIZE THE PHYSICIAN LISTED ABOVE TO RELEASE TO MY INSURANCE COMPANY/OR ITS REPRESENTATIVES OR AGENTS, ANY MEDICAL INFORMATION RELATIVE TO THE SERVICES RENDERED TO ME. I ACKNOWLEDGE THAT A PHOTOCOPY OR FAX OF THIS ORIGINAL IS AS VALID AS THE ORIGINAL.

  
Your signature here

8-24-10  
Today's date



## PRIVACY OF MEDICAL RECORDS

Our physicians and staff are fully and acutely aware of the potentially sensitive nature of the information contained in your medical record. Therefore, we ask that you provide us below with a list of those individuals or parties whom you intend to have access to such information in your medical records, and those whom you do not. Unless you request otherwise, it is our policy to share such information with the following individuals or parties:

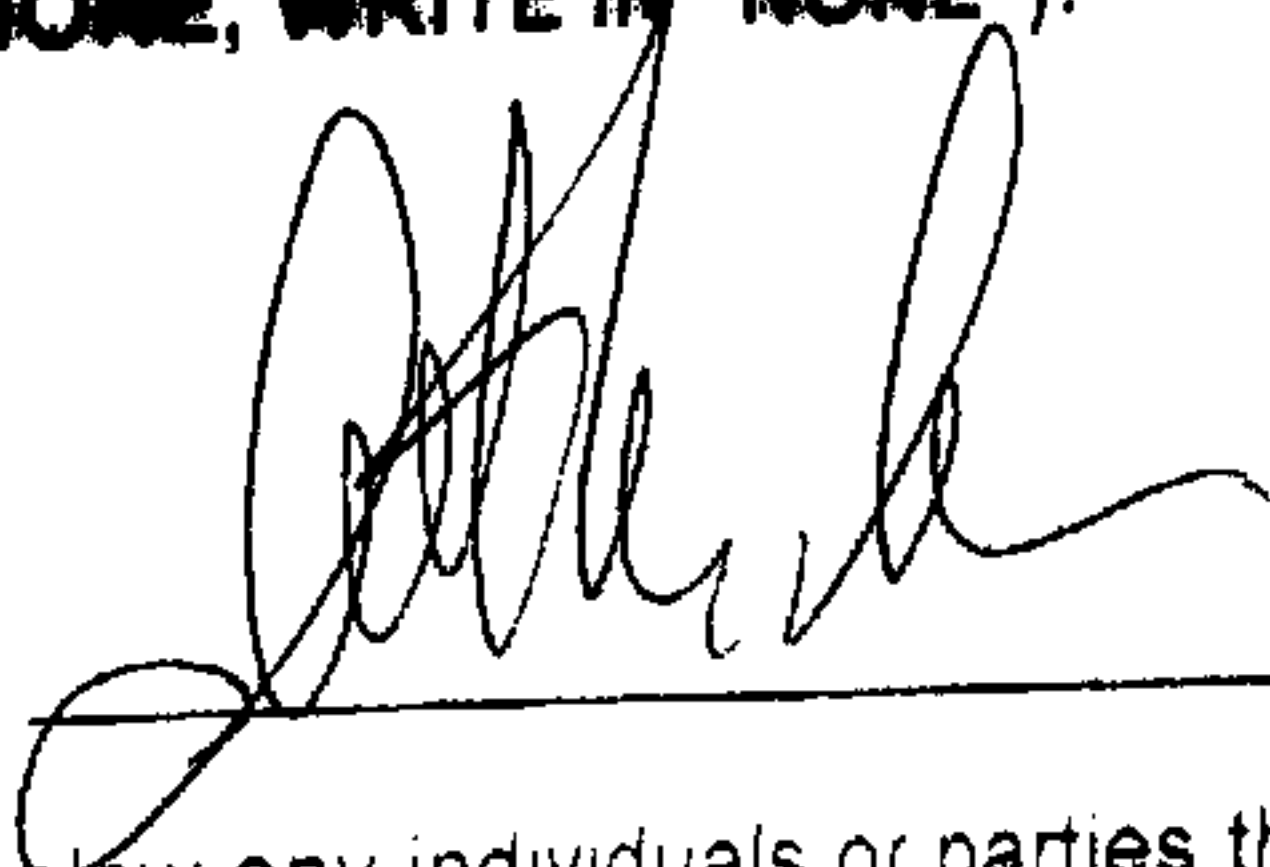
1. Your next of kin, usually identified as the emergency contact and/or the person(s) who accompanies you during your office visit(s), spouse, child (ren), and/or parent(s);
2. Your medical insurance carrier and its agents;
3. Your referring physician and his/her staff;
4. The physicians and professionals to whom we make referrals, including the pathologist, radiologist, and anesthesiologist, and their staff.

We CANNOT bill your insurance company and/or collect any money from them on your behalf unless we have your permission to disclose such information to them. Also, the quality of your medical care might be compromised if our physicians do not have your permission to consider your case fully and frankly with other physicians and professionals who are involved in your medical care.

Please acknowledge below that you permit the foregoing individuals or parties to have access to the information contained in your medical records by signing below, and list additional individuals or parties that you permit access to such information.

**THE FOLLOWING IS A LIST OF ADDITIONAL INDIVIDUALS OR PARTIES WHO HAVE MY PERMISSION TO ACCESS THE INFORMATION CONTAINED IN MY MEDICAL RECORD (IF THERE ARE NONE, WRITE IN "NONE"):**

Your signature (required):



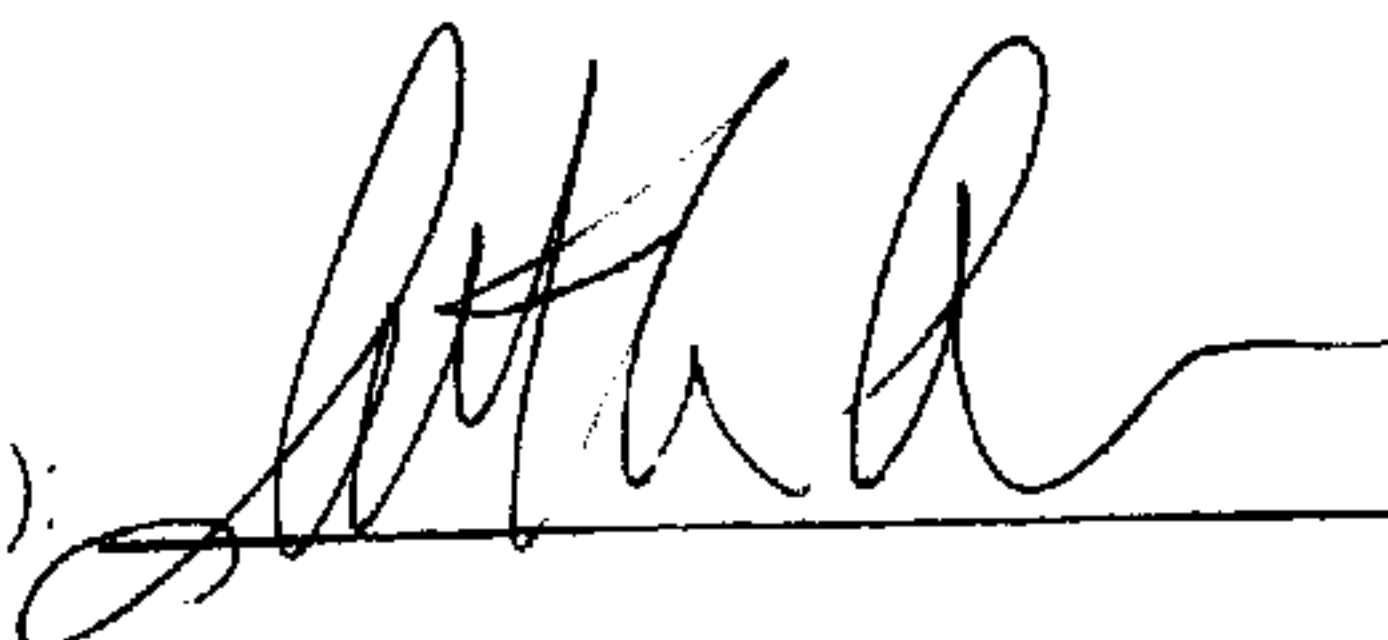
Date:

8-24-10

Please acknowledge below any individuals or parties that you DO NOT authorize access to the information contained in your medical record by signing below.

**THE FOLLOWING IS A LIST OF INDIVIDUALS OR PARTIES WHO DO NOT HAVE MY PERMISSION TO ACCESS THE INFORMATION CONTAINED IN MY MEDICAL RECORD (IF THERE ARE NONE, WRITE IN "NONE"):**

Your signature (required):



Date:

8-24-10



**THE CENTER FOR CHOLESTEROL MANAGEMENT**  
A Medical Corporation

**BILLING POLICY**

We would like to prevent any misunderstanding about our billing financial policies. Please let the office administration know if you would like to discuss any of the following policies in more detail.

If you belong to an HMO, or any other restricted insurance plan, you **MUST** let us know before you are treated. Some of these plans limit your choice of doctor or hospital, and some exclude particular medical conditions. If you need surgery, we will try to select the hospital and doctors from your plan, although this might not always be possible or practical, particularly with the pathologist and the radiologist. Please provide our business office with all of your insurance information before you are treated, and we will help you fulfill the terms of your policy so that you can obtain maximum and timely reimbursement.

We will send you monthly statements until your insurance company has paid, regardless of our provider status. This allows you to verify that your insurance company was billed correctly, and to see how long they take to pay. If you have more than one insurance policy and the benefits are not coordinated, each company will determine benefits separately. In this situation, it might happen that we have different agreements with different companies. We will then collect benefits from each company and reimburse you any amount above billed charges.

Starting January 2001, you will also need to complete a separate form, "Privacy of Medical Records," so that we have a clear understanding of those individuals and parties whom you intend to have access to information contained in your medical record, and those whom you do not.

We accept Visa, MasterCard, and Diner's. There is a \$25 charge for all checks returned by the bank. If you would like us to bill your insurance company on your behalf, please complete the Assignment of Benefits sections below. Please sign below once you have had a chance to review our billing policies.

**I AUTHORIZE MICHAEL RICHMAN M.D., AND STAFF TO PROVIDE ME WITH REASONABLE AND PROPER MEDICAL CARE. I UNDERSTAND THAT I WILL HAVE AN OPPORTUNITY TO ASK QUESTIONS AND TO HAVE MY QUESTIONS ANSWERED, BEFORE I DECIDE TO PROCEED.**

Your signature (required):

Date: 8-29-10

**THE CENTER FOR CHOLESTEROL MANAGEMENT**  
**A Medical Corporation**  
**1950 Sawtelle Blvd, Suite 150**  
**Los Angeles, CA 90025**

Dr. Richman is a participating provider with most major insurance plans.

Should Dr. Richman not be a provider under your plan, he will act as a provider, and accept the ALLOWED amount for any covered charges.

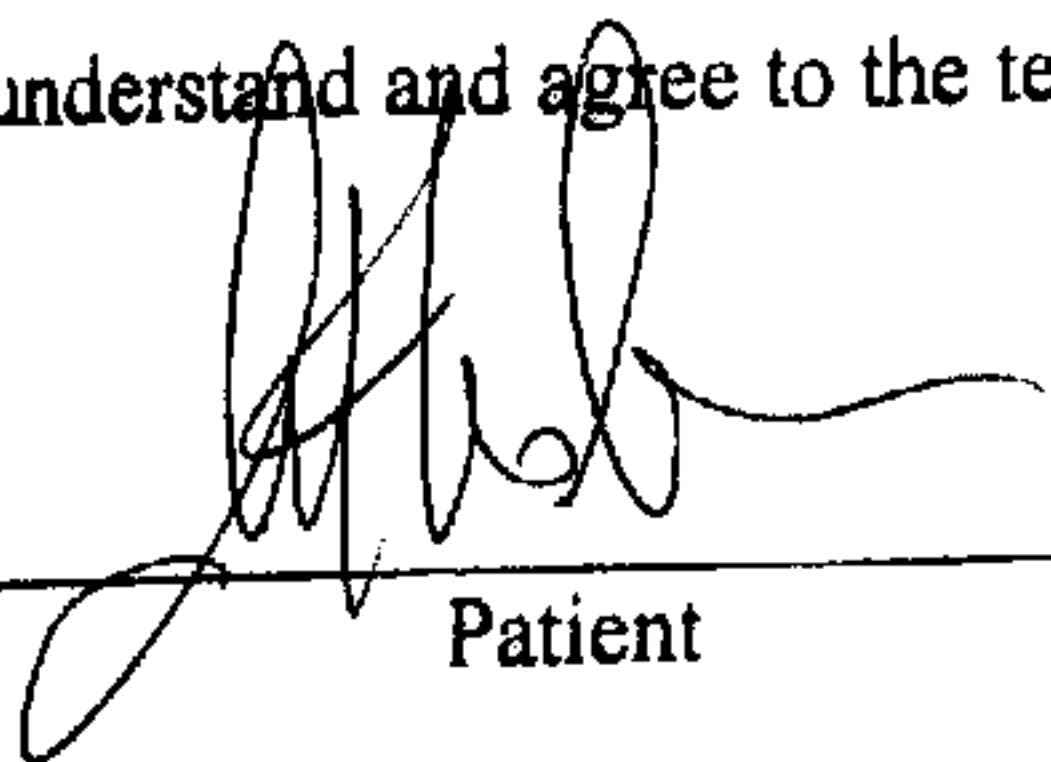
Your insurance may only pay a portion of the allowed amount, therefore leaving you with a balance which you will be billed for.

It is your responsibility to verify with your insurance carrier your individual coverage benefits prior to being seen for your initial consultation.

During your consultation, additional x-rays or ultra sounds may be necessary to fully determine the nature and extent of your condition, as well as determining your optimal course of treatment. These will be billed to your insurance in addition to the consultation fee.

Prior to your consultation, you can contact your carrier to determine what your financial portion will be.

I understand and agree to the terms outlined above

  
\_\_\_\_\_  
Patient

8.24.10  
\_\_\_\_\_  
Date



Lumenos HSA 3000 (100/70)

Member Name:  
**Jonathan Aaron**

Member ID:  
**XDX130A57237**

Group No.: **482287**  
Contract Code: **DY11**  
Effective Date: **09/01/2010**  
BIN: **610575**  
Plan: **040**

Office Visit 0%



[anthem.com/ca](http://anthem.com/ca)

**Members:** Possession or use of this card does not guarantee payment. If a provider does not submit your claim on your behalf, please file the claim to:

Anthem Blue Cross  
PO Box 60007  
Los Angeles, CA 90060

**Providers:** Please submit claims to local Blue plan. If Medicare is primary, file claims with Medicare.

Member Services:	1-866-837-4595
Provider Services:	1-800-677-6669
Pharmacy Services:	1-800-281-4890
Preservice Review	1-800-274-7767
24/7 Nurseline:	1-866-800-8780
Coverage while traveling:	1-800-810-BLUE

This plan is insured by Anthem Blue Cross Life and Health Insurance Company. Blue Cross of California using the trade name Anthem Blue Cross, administers claims on behalf of Anthem Blue Cross, Life and Health Insurance Company but is not financially liable for benefits payable. Independent licensees of the Blue Cross Association.

Rx 09/11/10



**CALIFORNIA**  
**DRIVER LICENSE**  
A123456789  
EXPIRES 11-03-16  
JONATHAN A AARON  
1237 S BEDFORD ST  
W LOS ANGELES CA 90035  
SEX:M WT:6-02 HAIR:GRY WT:200  
DOB:11-03-61  
CLASS:C  
FD/14



Michael F. Richman, M.D., F.A.C.S.  
Cardiothoracic, General and Vein Surgery  
LIC# G74625 • DEA# BR3315567  
1950 Sawtelle Boulevard, # 150  
Los Angeles, CA 90025  
(310)481-3939 • Fax (310)481-3949

00198

R Name Jonathan Larson D.O.B. \_\_\_\_\_ ☐ Female  
☐ Male

Address \_\_\_\_\_ Phone \_\_\_\_\_

1) Zetia 10mg  
Quantity: ☐ 1-24 ☒ 25-49 ☐ 50-74  
☐ 75-100 ☐ 101-150 ☐ 151 and over  
Units \_\_\_\_\_ Refills \_\_\_\_\_ ☐ NR ☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5  
☐ Do not substitute Initial \_\_\_\_\_

2) Tab po qd #30  
Quantity: ☐ 1-24 ☐ 25-49 ☐ 50-74  
☐ 75-100 ☐ 101-150 ☐ 151 and over  
Units \_\_\_\_\_ Refills \_\_\_\_\_ ☐ NR ☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5  
☐ Do not substitute Initial \_\_\_\_\_

3) \_\_\_\_\_  
Quantity: ☐ 1-24 ☐ 25-49 ☐ 50-74  
☐ 75-100 ☐ 101-150 ☐ 151 and over  
Units \_\_\_\_\_ Refills \_\_\_\_\_ ☐ NR ☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5  
☐ Do not substitute Initial \_\_\_\_\_

Prescription is VOID if the number of drugs prescribed is not noted. ☐ 1 ☐ 2 ☐ 3

NON-NEGOTIABLE

Date

7/22/11

GF92PE



Michael F. Richman, M.D., F.A.C.S.  
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1950 Sawtelle Boulevard, # 150  
Los Angeles, CA 90025  
(310)481-3939 • Fax (310)481-3949

00318

R Name Jonathan Aaron D.O.B. \_\_\_\_\_ ☐ Female  
Address \_\_\_\_\_ Phone \_\_\_\_\_ ☐ Male

1) <u>Crestor 10mg</u>	Quantity: <input type="checkbox"/> 1-24 <input type="checkbox"/> 25-49 <input checked="" type="checkbox"/> 50-74 <input type="checkbox"/> 75-100 <input type="checkbox"/> 101-150 <input type="checkbox"/> 151 and over Units _____ Refills _____ <input type="checkbox"/> NR <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/> Do not substitute Initial _____
2) <u>1 tab po qd</u>	Quantity: <input type="checkbox"/> 1-24 <input type="checkbox"/> 25-49 <input type="checkbox"/> 50-74 <input type="checkbox"/> 75-100 <input type="checkbox"/> 101-150 <input type="checkbox"/> 151 and over Units _____ Refills _____ <input type="checkbox"/> NR <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/> Do not substitute Initial _____
3) <u># 30</u>	Quantity: <input type="checkbox"/> 1-24 <input type="checkbox"/> 25-49 <input type="checkbox"/> 50-74 <input type="checkbox"/> 75-100 <input type="checkbox"/> 101-150 <input type="checkbox"/> 151 and over Units _____ Refills _____ <input type="checkbox"/> NR <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/> Do not substitute Initial _____

Prescription is VOID if the number of drugs prescribed is not noted. ☐ 1 ☐ 2 ☐ 3

NON-NEGOTIABLE

[Signature] Date 11/9/10

GC00RP





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1950 Sawtelle Boulevard, # 150  
Los Angeles, CA 90025  
(310)481-3939 • Fax (310)481-3949

00160

R Name Jonathan Carson D.O.B. \_\_\_\_\_ ☐ Female  
Address \_\_\_\_\_ Phone \_\_\_\_\_ ☐ Male

1) <u>levaz 1000mg</u> <u>4 tabs po qd #120</u>	Quantity: <input type="checkbox"/> 1-24 <input type="checkbox"/> 25-49 <input type="checkbox"/> 50-74 <input type="checkbox"/> 75-100 <input type="checkbox"/> 101-150 <input type="checkbox"/> 151 and over Units _____ Refills <u>5</u> <input type="checkbox"/> NR <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input checked="" type="checkbox"/> 5 <input type="checkbox"/> Do not substitute Initial _____
2) <u>glucophage XR</u> <u>500mg</u> <u>2 tabs po qd #60</u>	Quantity: <input type="checkbox"/> 1-24 <input type="checkbox"/> 25-49 <input checked="" type="checkbox"/> 50-74 <input type="checkbox"/> 75-100 <input type="checkbox"/> 101-150 <input type="checkbox"/> 151 and over Units _____ Refills <u>5</u> <input type="checkbox"/> NR <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input checked="" type="checkbox"/> 5 <input type="checkbox"/> Do not substitute Initial _____
3) <u>2 tabs po qd</u>	Quantity: <input type="checkbox"/> 1-24 <input type="checkbox"/> 25-49 <input type="checkbox"/> 50-74 <input type="checkbox"/> 75-100 <input type="checkbox"/> 101-150 <input type="checkbox"/> 151 and over Units _____ Refills _____ <input type="checkbox"/> NR <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/> Do not substitute Initial _____

Prescription is VOID if the number of drugs prescribed is not noted. ☐ 1 ☒ 2 ☐ 3

NON-NEGOTIABLE

Date

9/14/10

GC00RP



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LIC# G74625 • DEA# BR3315567  
1950 Sawtelle Boulevard, # 150  
Los Angeles, CA 90025  
(310)481-3939 • Fax (310)481-3949

00159

R Name Jonathan Aaron D.O.B. \_\_\_\_\_ ☐ Female  
Address \_\_\_\_\_ Phone \_\_\_\_\_ ☐ Male

1) Cresta 10mg #30 1 tab po qd	Quantity: <input type="checkbox"/> 1-24 <input checked="" type="checkbox"/> 25-49 <input type="checkbox"/> 50-74 <input type="checkbox"/> 75-100 <input type="checkbox"/> 101-150 <input type="checkbox"/> 151 and over Units _____ Refills _____ <input type="checkbox"/> NR <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input checked="" type="checkbox"/> 5 <input type="checkbox"/> Do not substitute Initial _____
2) Teclor 435mg 1 tab po qd #30	Quantity: <input type="checkbox"/> 1-24 <input checked="" type="checkbox"/> 25-49 <input type="checkbox"/> 50-74 <input type="checkbox"/> 75-100 <input type="checkbox"/> 101-150 <input type="checkbox"/> 151 and over Units _____ Refills _____ <input type="checkbox"/> NR <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input checked="" type="checkbox"/> 5 <input type="checkbox"/> Do not substitute Initial _____
3)	Quantity: <input type="checkbox"/> 1-24 <input type="checkbox"/> 25-49 <input type="checkbox"/> 50-74 <input type="checkbox"/> 75-100 <input type="checkbox"/> 101-150 <input type="checkbox"/> 151 and over Units _____ Refills _____ <input type="checkbox"/> NR <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/> Do not substitute Initial _____

Prescription is VOID if the number of drugs prescribed is not noted. ☐ 1 ☒ 2 ☐ 3

NON-NEGOTIABLE

Date

9/14/10

GC00RP