$\mathbf{}$				Page 1
	Lipoprotein	SPECTRA	CELL LABOR	ATORIES
∖ LPP	Particle Profile <sup>™</sup>	10401 Town Park Dr. Houston, TX 77072 Laboratory Director: Lipid Science Director:	Tel: 713-621-3101 80 CLIA ID 45D0710715 Fi John F. Crawford, Ph.D. Jan M. Troup, Ph.D.	0-227-5227 ax: 713-621-3234
Name: Doe, John			Batcl	n: B4452
DOB: February 23, 1946			Accession N	o: K10133
Physician: Dr. Joe E. Lewis			Draw Date	e: October 16, 2010
Reference: Data\b32.2\K22060.8602.5	5.Rpt 0.150		Report Date	e: March 14, 2011
	<b>Lipoprote</b> Green - Normal	In Particle N Yellow - Borderline Reference Value	umbers ed - Abnormal	Patient Results
VLDL Particles	0 42.5	85	127.5 170	112
Total LDL Particles	0 450	900	1350 1800	1031
Non - HDL Particles	0 500	1000	1500 2000	1143
RLP (Remnant Lipoprotein)	, , 0 75	150	225 300	230
Small - Dense LDL III	, , 0 150	300	450 600	339
Small - Dense LDL IV	, , , , , , , , , , , , , , , , , , , ,	100	150 200	126
Total HDL Particles	14000 10500	7000	3500 0	7148
Large - Buoyant HDL 2b	2800 2100	1400	700 0	1268

	Biomarke	rs and Risl	k Factors	<u>P</u>	atient Results
Apo B-100 (mg/dL)	, , , 0 50	100	150	200	115
Lp(a) (mg/dL) <sup>1</sup>	0 15	30	45	60	42.5
Metabolic Syndrome Traits	0 0.75	1.5	2.25	3	1
Lp-PLA2 (ng/mL)	, , 0 100	200	300	400	225
C-Reactive Protein-hs (mg/L)	0 1.5	] <b>♦</b>	4.5		3.2
<b>Insulin</b> (μιυ/mL)	0 17.5	35	52.5	70	15.5
Homocysteine (µmol/L)	, , 0 5.5	↓ 11	16.5	22	10.2



## Step 1. Primary Risk Assessment

- Lipoprotein non-HDL particle numbers and other lipid and non-lipid risk factors may show a greater patient risk than a standard lipid panel and therefore, a greater LDL reduction than indicated by a standard lipid panel. Non-HDL lipoprotein particle numbers and/or Apo B-100 are measures of the number of atherogenic lipoprotein particles and are compliant with the recent consensus statement from the American Diabetes Association and the American College of Cardiology<sup>1</sup> stating that lipoprotein particle numbers are more predictive of CVD risk than cholesterol.
- Moderate to elevated triglycerides can cause the lipoproteins to be cholesterol depleted or triglyceride enriched and these
  patients will show a greater CVD risk from non-HDL particle numbers or Apo B-100 than from a standard lipid panel.
  This occurs in about 30% of the population.

## Step 2. Modify Risk Using Metabolic Syndrome Traits, Lp(a) and Inflammation Risk Markers

- Evaluate possible Metabolic Syndrome by combining the lipid traits from the LPP<sup>™</sup> test with possible hypertension, obesity and high glucose. Three total traits is a diagnosis of Metabolic Syndrome and raises the CVD risk to the next level. Also check for insulin resistance using the LPP<sup>™</sup> fasting insulin value.
- Take into account additional risk from elevated Lp(a) or inflammation markers such as hs-CRP or Lp-PLA2 if ordered, in determining the final treatment goals. Also consider non-lipid risk factors such as hypertension, obesity, high glucose, smoking, family history and other medical history.
- The risk assessment and treatment goal from **Step 1** should be adjusted in light of the presence of these additional Biomarkers and Risk Factors.
- A standard directly measured cholesterol **Lipid Panel** is presented at the bottom of the report for comparison to previously determined lipid results.

## Step 3. Determine Therapeutic Approach Based On the Lipid Subgroup Distribution, Lp(a) and Therapeutic Guidelines

- Using the risk level established in **Step 2** and treatment goals from the NCEP guidelines, determine if VLDL, LDL subgroups, HDL subgroups and/or Lp(a) should be therapeutic targets.
- The LPP<sup>™</sup> particle numbers by subgroup and Lp(a) each have a specific therapeutic approach that is most effective. Often, combination therapy is needed to address the different risk areas.
- A special HDL species enriched in Apo C-I is atherogenic but displays health attributes<sup>2</sup>. It is generally identified by high HDL>70mg/dL, high HDL 2b > 4000 nmol/L with a high HDL peak or hump extending into LDL IV region and low TG's < 70 mg/dL. Check for CVD development with a CIMT, a Coronary Calcium Score or other method to confirm.</li>
- Refer to the **LPP<sup>TM</sup> Therapeutic Guidelines** for lipoprotein subgroup specific information.

I.Diabetes Care, Volume 31, Number 4, April 2008 2.Kwiterovich P., et al. JAMA 2005; 293(15): 1891



Lipoprotein Parti	cle Numbers (nmol/L)	
Value	Reference Value	Alert (Notes Page 3)
VLDL Particles 112	<85	Borderline High (12)
Total LDL Particles 1031	<900	High (13)
Non - HDL Particles 1143	<1000	High (19)
RLP (Remnant Lipoprotein) 230	<150	High (14)
Small - Dense LDL III 339	<300	Borderline High (15)
Small - Dense LDL IV 126	<100	High (16)
Total HDL Particles 7148	>7000	Borderline-M, Low-F (17)
Large - Buoyant HDL 2b	>1400	Low (18)

<b>Biomarkers and Risk Factors</b>			
	Value	Reference Value	Alert (Notes Page 3)
Apo B-100 (mg/dL)	115	<100	High (20)
Lp(a) (mg/dL)	42.5	<30.0 <sup>1</sup>	High (6)
Metabolic Syndrome Traits	1	0	Possible (8)
Lp-PLA2 (ng/mL)	225	<200	Borderline (7)
C-Reactive Protein-hs (mg/L)	3.2	<3.0	High (9)
Insulin (uIU/mL)	15.5	<35.0	
Homocysteine (umol/L)	10.2	<11.0	

	Lipid Pan	el (mg/dL)	
	Value	Reference Value	Alert (Notes Page 3)
Total Cholesterol	193	<200	
LDL - Cholesterol	121	<130	Borderline High (2)
HDL - Cholesterol	40	>40	Borderline (3)
Triglycerides	217	<150	High (4)
Non - HDL- Cholesterol	153	<160	Borderline (5)

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1. Reference Value for Blacks is 50.0 mg/dL

## SpectraCell Clinical Suggestions for Alert References

1	Elevated Total Cholesterol (TC): Borderline 20	0-240 mg/dl consider treatment when patient has 2 of	r more risk factors. High >240 mg/dl consider		
	treatment after assessing secondary dyslipidemias	s. >300 mg/dl suggest higher likelihood of genetic ca	usation.		
2	Elevated LDL-Cholesterol (LDL-C): Follow ATPIII Guidelines for treatment goals and strategy: See online at				
	http://www.nhlbi.nih.gov/guidelines/cholester	ol/atglance.htm			
	0-1 Risk factors: Goal <160 mg/dl	2 Risk factors: Goal <130 mg/dl	High Risk/ CHD or CHD equivalent Goal: <100 mg/dl (or 70 mg/dl)		
3	Low HDL-Cholesterol (HDL-C): Follow ATF	III Guidelines for treatment goals and strategy:	See online at		
	http://www.nhlbi.nih.gov/guidelines/cholester	ol/atglance.htm			
	Males: <40 mg/dl (is 1 of 5 traits for Metabolic	Females: <50 mg/dl (is 1 of 5 traits for Metabolic			
	Syndrome)	Syndrome)			
4	Elevated Triglycerides (TG) : TG >150 mg/dl i	s 1 of 5 traits for Metabolic Sydnrome			
	Borderline: 150-199 mg/dl	High: 200-499 mg/dl	Very High: 500 mg/dl Consider Genetic disorders.		
	Follow ATPIII Guidelines for treatment goals	and strategy: See online at http://www.nhlbi.nih.	gov/guidelines/cholesterol/atglance.htm		
	*Triglyceride levels can be elevated if patient wa	s not fasted- confirm that patient was properly fasted	before setting goals.		
5	Non-HDL-C: A cholesterol measurement that predicts risk better than LDL-C				
	Optimal: <130 mg/dL	Borderline Risk : 130-160 mg/dL	High Risk: >160 mg/dL		
6	High Lp(a). (Lp(a) >30.0): Lp(a) is highly ass	ociated with cardiovascular disease. Lp(a) is an inhe	rited trait and does not respond to diet, exercise or		
	statin drugs. Treatment for high Lp(a) is typicall	y niacin and aggressive LDL treatment.			
7	High Lp-PLA2. (200 to 235 borderline, > 235	high): Is associated with a two fold increased car	diovascular risk.		
8	Metabolic Syndrome Traits: This test reports of	nly 2 of the 5 traits associated with the ATPIII Metab	oolic Sydrome Definition: Elevated TG (>150 mg/dl),		
	Low HDL-C (<50 mg/dl in men; <40 mg/dl in wo	omen). Additionally, this number adds a third feature	(elevat		
9	C-Reactive Protein (hs-CRP): Marker of Inflan	imation tied to increased cardiovascular risk			
	Low Risk: 0-1 mg/L	Borderline: 1-3 mg/L	High: >3 mg/L		
10	<b><u>Elevated Insulin:</u></b> (Insulin > 35.0 uIU/mL): Hig	h fasting insulin is associated with increased cardiov	ascular risk and/or metabolic syndrome. Test units		
	may not correlate to other labs using different me	thods.			
11	* Insulin levels can be elevated if patient was no	Tasted- confirm that patient was properly fasted being	bre setting goals.		
11	Elevated Homocysteine: Intermediate in methyl	ation pathways- risk factor for CVD, stroke, Alznein			
	Ideal: <11 µmol/L	Borderline: 11-15 µmol/L: Consider dietary changes and/or multivitamins/B-complex	High: Above 15 µmol/L Consider dietary changes and/or multivitamins/B-complex supplements.		
		supplements.	1 11		
12	High VLDL Particle Number (VLDL > 85 nm	<b><u>ol/L</u></b> ): No reported clinical guidance by NCEP, howe	ever this correlates to triglyceride values of over 200		
	mg/dL, high RLP and possible metabolic syndror	ne.			
13	Borderline, High to Very High LDL Particle N	umber (LDL > 700, 900, 1100 nmol/L): Patients	with 2 or more risk factors are recommended to		
	initiate therapeutic lifestyle changes and/or drug	herapy to lower $LDL < 900$ nmol/L. Patients with Cl	HD or CHD equi		
14	Flowated Domnant Linenvetein Doutiele Numb	or (DID >150 nmol/L). This now NCED risk factor	has been shown to be highly correlated with CHD		
14	and should be monitored along with other risk fac	tors during lifestyle supplement and/or drug treatme	nas been shown to be nightly correlated with CITD		
	and should be monitored along with other fisk fac	tors during mestyle, supplement and/or drug reading	nt. Onlega-5 ta		
15	Borderline to High LDL III Particle Number (	LDL III>300 nmol/L): Indicates an abundance of	small-dense atherogenic LDL-particles. Management		
16	should be considered depending on LDL-C goals	and risk factors. Reducing LDL-C and TG often will	lower dense-LDL		
	should be considered depending on LDL-C goals Elevated LDL IV (LDL IV> 100 nmol/L): Ind	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and		
	should be considered depending on LDL-C goals <u>Elevated LDL IV (LDL IV&gt; 100 nmol/L)</u> : Ind d=1.08 and often is located in the range for LDL	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar,		
	should be considered depending on LDL-C goals <u>Elevated LDL IV (LDL IV&gt; 100 nmol/L):</u> Ind d=1.08 and often is located in the range for LDL	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar,		
17	should be considered depending on LDL-C goals <u>Elevated LDL IV (LDL IV&gt; 100 nmol/L):</u> Ind d=1.08 and often is located in the range for LDL <u>Low HDL particle count &lt;7000 nmol/L, 7000</u>	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s - 8500 nmol/L is Borderline for Males and Low for	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar, or Females: Indicates potential for atherogenic		
17	should be considered depending on LDL-C goals Elevated LDL IV (LDL IV> 100 nmol/L): Ind d=1.08 and often is located in the range for LDL Low HDL particle count <7000 nmol/L, 7000 dyslipidemia. Beneficial therapies similar to thos	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s - 8500 nmol/L is Borderline for Males and Low for e which raise HDL-C and reducing elevated TG (Die	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar, <b>r Females:</b> Indicates potential for atherogenic t, Exercise, Niaci		
17	should be considered depending on LDL-C goals <u>Elevated LDL IV (LDL IV&gt; 100 nmol/L)</u> : Ind d=1.08 and often is located in the range for LDL <u>Low HDL particle count &lt;7000 nmol/L, 7000</u> dyslipidemia. Beneficial therapies similar to thos <u>Risk Factor for HDL2b between 1400 and 180</u>	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s - 8500 nmol/L is Borderline for Males and Low for e which raise HDL-C and reducing elevated TG (Die 0 nmol/L is borderline for males and risk factor f	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar, <b>r Females:</b> Indicates potential for atherogenic t, Exercise, Niaci <b>or females.</b> Values less than 1400 is a risk factor		
17 18	should be considered depending on LDL-C goals <u>Elevated LDL IV (LDL IV&gt; 100 nmol/L)</u> : Ind d=1.08 and often is located in the range for LDL <u>Low HDL particle count &lt;7000 nmol/L, 7000</u> dyslipidemia. Beneficial therapies similar to thos <u>Risk Factor for HDL2b between 1400 and 180</u> <u>for males</u> : Indicates that the HDL reverse transp	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s - 8500 nmol/L is Borderline for Males and Low for e which raise HDL-C and reducing elevated TG (Die 0 nmol/L is borderline for males and risk factor for ort system is not working well to remove excess cho	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar, or Females: Indicates potential for atherogenic t, Exercise, Niaci or females. Values less than 1400 is a risk factor esterol.		
17 18 19	should be considered depending on LDL-C goals <u>Elevated LDL IV (LDL IV&gt; 100 nmol/L):</u> Ind d=1.08 and often is located in the range for LDL <u>Low HDL particle count &lt;7000 nmol/L, 7000</u> dyslipidemia. Beneficial therapies similar to thos <u>Risk Factor for HDL2b between 1400 and 180</u> <u>for males:</u> Indicates that the HDL reverse transp <u>Non-HDL Particle Numbers: Non-HDL particle</u>	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s - 8500 nmol/L is Borderline for Males and Low for e which raise HDL-C and reducing elevated TG (Die 0 nmol/L is borderline for males and risk factor f ort system is not working well to remove excess choic e numbers is the best overall indicator of CVD risk.	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar, or Females: Indicates potential for atherogenic t, Exercise, Niaci or females. Values less than 1400 is a risk factor esterol.		
17 18 19	should be considered depending on LDL-C goals Elevated LDL IV (LDL IV> 100 nmol/L): Ind d=1.08 and often is located in the range for LDL Low HDL particle count <7000 nmol/L, 7000 dyslipidemia. Beneficial therapies similar to thos Risk Factor for HDL2b between 1400 and 180 for males: Indicates that the HDL reverse transp Non-HDL Particle Numbers: Non-HDL particle Optimal: <800 nmol/L	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s - 8500 nmol/L is Borderline for Males and Low for e which raise HDL-C and reducing elevated TG (Die 0 nmol/L is borderline for males and risk factor f ort system is not working well to remove excess choic e numbers is the best overall indicator of CVD risk. Borderline Risk : 800-1000 nmol/L	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar, <b>r Females:</b> Indicates potential for atherogenic t, Exercise, Niaci <b>or females.</b> Values less than 1400 is a risk factor esterol. High Risk: >1000 nmol/L		
17 18 19 20	should be considered depending on LDL-C goals Elevated LDL IV (LDL IV> 100 nmol/L): Ind d=1.08 and often is located in the range for LDL Low HDL particle count <7000 nmol/L, 7000 dyslipidemia. Beneficial therapies similar to thos Risk Factor for HDL2b between 1400 and 180 for males: Indicates that the HDL reverse transp Non-HDL Particle Numbers: Non-HDL particle Optimal: <800 nmol/L ApoB: A measure of all non-HDL particle number	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s - 8500 nmol/L is Borderline for Males and Low for e which raise HDL-C and reducing elevated TG (Die 0 nmol/L is borderline for males and risk factor f ort system is not working well to remove excess choice numbers is the best overall indicator of CVD risk. Borderline Risk : 800-1000 nmol/L ers.	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar, <b>r Females:</b> Indicates potential for atherogenic t, Exercise, Niaci <b>or females.</b> Values less than 1400 is a risk factor esterol. High Risk: >1000 nmol/L		
17 18 19 20	should be considered depending on LDL-C goals Elevated LDL IV (LDL IV> 100 nmol/L): Ind d=1.08 and often is located in the range for LDL Low HDL particle count <7000 nmol/L, 7000 dyslipidemia. Beneficial therapies similar to thos Risk Factor for HDL2b between 1400 and 180 for males: Indicates that the HDL reverse transp Non-HDL Particle Numbers: Non-HDL particle Optimal: <800 nmol/L ApoB: A measure of all non-HDL particle number Optimal: <80 mg/dL (goal for very high risk	and risk factors. Reducing LDL-C and TG often will icated abundance of very small-dense atherogenic pa IV. Treatment for high LDL IV and Lp(a) are very s - 8500 nmol/L is Borderline for Males and Low for e which raise HDL-C and reducing elevated TG (Die 0 nmol/L is borderline for males and risk factor f ort system is not working well to remove excess choic e numbers is the best overall indicator of CVD risk. Borderline Risk : 800-1000 nmol/L ers. Borderline Risk: 80-100 mg/dL	lower dense-LDL rticles. Lp(a) is found typically between d=1.05 and imilar, <b>r Females:</b> Indicates potential for atherogenic t, Exercise, Niaci <b>or females.</b> Values less than 1400 is a risk factor esterol. High Risk: >1000 nmol/L High Risk: >100 mg/dL		