



LipoPortfolio

Powered by Vantera®

Comprehensive, Innovative Panels
to **Help Identify Cardiometabolic Risk**

Cardiovascular disease and diabetes

Two of the leading chronic diseases in the US and drivers of morbidity, mortality and rising health care costs.^{1,2}



Guidelines suggest evaluating and controlling cardiometabolic risk,³ a condition in which the risk of developing atherosclerotic cardiovascular disease (ASCVD) and type 2 diabetes mellitus (T2D) is significantly enhanced by the presence of insulin resistance and atherogenic dyslipidemia.⁴



Lipid panels and glycemia assays, such as HbA1c and glucose, have traditionally been used to assess a patient's risk of developing ASCVD and T2D. These assays continue to be the foundation of risk assessment and primary prevention guidelines³ but in some cases they may not reflect the full risk status of a patient. Additional information, such as the presence of insulin resistance, systemic inflammation, and lipoprotein particle levels and size can provide a more detailed—and in some cases, a more accurate—depiction of a patient's cardiometabolic risk.⁵⁻¹³

The inclusion of measurements of apolipoprotein B (ApoB), GlycA (systemic inflammation), and Diabetes Risk Index (insulin resistance and metabolic dysregulation), along with standard lipid testing, can provide actionable clinical data and a more in-depth risk assessment for individualized treatment, as well as prevention and intervention strategies.

ApoB

What is it:

Low-density lipoprotein (LDL) particles are a leading cause of ASCVD.¹⁴ Apolipoprotein B (ApoB) is the primary protein component of LDL and triglyceride-rich lipoprotein (TRL) particles.¹⁵

How will this information enable physicians to help their patients?

The cholesterol content of these atherogenic lipoproteins can vary widely among patients, but each contains one ApoB protein. Measured ApoB thus provides an accurate assessment of atherogenic particle number and its associated ASCVD risk, whereas LDL cholesterol (LDL-C) does not.¹⁵

Although LDL-C, non-HDL-C, and ApoB are highly correlated, “discordance analyses” have demonstrated that ASCVD risk tracks with ApoB, not the cholesterol measures, when levels differ.¹⁵

GlycA

What is it:

GlycA is a composite biomarker that integrates the protein levels and glycosylation states of several of the most abundant acute phase proteins in serum.¹⁶ Data suggest that GlycA has clinical utility similar to high sensitivity C-reactive protein (hsCRP), but has the advantage of having much lower intra-individual day-to-day variability.¹⁶

How will this information enable physicians to help their patients?

GlycA, used as an alternative to or in conjunction with hsCRP, may aid in the identification and stratification of individuals at risk for future ASCVD events. It may also aid as an independent prognostic marker for recurrent cardiovascular events in patients with stable coronary disease or acute coronary syndrome.

Diabetes Risk Index (DRI)

What is it:

LabCorp developed the Diabetes Risk Index (DRI) to help clinicians distinguish, among patients with similar glucose levels, those that have a greater versus lesser likelihood of developing T2D. Since >80 million U.S. adults qualify by glycemic criteria as “prediabetic”, a need has been recognized for a more refined and cost-effective approach to diabetes prevention that directs treatment to the subset of patients at highest risk.¹⁷ Waiting until the onset of prediabetes before initiating preventive measures may also be suboptimal, since many individuals with normal glucose levels progress to T2D in a relatively short time period.¹⁸

The DRI score (values 1-100) is derived from the patient’s measured Lipoprotein Insulin Resistance Index (LP-IR)¹⁹ and selected branched-chain amino acid (BCAA) levels. Both LP-IR and BCAA levels are independently associated with insulin resistance and both have been shown in multiple prospective clinical studies to predict the development of T2D independent of the level of glycemia.⁸⁻¹³

How will this information enable physicians to help their patients?

The LP-IR score, the main determinant of DRI, has been shown to be modifiable by drug and lifestyle interventions that produce weight loss and increase insulin sensitivity.²⁰⁻²² Reductions of DRI and LP-IR are thus clinically achievable and likely to reflect a corresponding reduction of the risk of developing diabetes.



LabCorp developed the Diabetes Risk Index (DRI) to assist clinicians with identifying patients who could be at risk of developing T2D.

Through its efficient, proprietary Vantera® testing platform, LabCorp offers comprehensive, innovative panels to help identify cardiometabolic risk—at cost-effective pricing.

Test Name	Number
Lipid Panel with GlycA (Inflammation)	123510
Lipid Panel with Diabetes Risk Index (DRI)	123525
Lipid Panel with Apolipoprotein B (ApoB)	123544
Lipid Panel with GlycA (inflammation) and Diabetes Risk Index (DRI)	123559
Lipid Panel with Apolipoprotein B (ApoB), GlycA (Inflammation), Diabetes Risk Index (DRI)	123567

References

- Chapel JM, Ritchey MD, Zhang D, Wang G. Prevalence and medical costs of chronic diseases among adult Medicaid beneficiaries. *Am J Prev Med.* 2017;53(6S2):S143-S154. doi:10.1016/j.amepre.2017.07.019
- Centers for Disease Control and Prevention. Chronic diseases in America. <https://www.cdc.gov/chronicdisease/resources/infographic/chronic-diseases.htm>. Accessed December 12, 2019.
- Arnett DK, Blumenthal RS, Albert MA, et al. 2019 ACC/AHA Guideline on the Primary Prevention of Cardiovascular Disease: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. *Circulation.* 2019;140(11):e596-e646. doi:10.1161/CIR.0000000000000678
- Ruilope LM, de la Sierra A, Segura J, Garcia-Donaire JA. The meaning of cardiometabolic risk in hypertensive patients. *US Endocrinology.* 2007;(1):60-3. <https://www.touchendocrinology.com/the-meaning-of-cardiometabolic-risk-in-hypertensive-patients/>. Accessed December 12, 2019. DOI: <http://doi.org/10.17925/USE.2007.00.1.60>
- Choy E, Ganeshalingam K, Semb AG, Szekanecz Z, Nurmohamed M. Cardiovascular risk in rheumatoid arthritis: recent advances in the understanding of the pivotal role of inflammation, risk predictors and the impact of treatment. *Rheumatology (Oxford).* 2014 Dec;53(12):2143-2154.
- Myasoedova E, Crowson CS, Kremers HM, et al. Lipid paradox in rheumatoid arthritis: the impact of serum lipid measures and systemic inflammation on the risk of cardiovascular disease. *Ann Rheum Dis.* 2011 Mar;70(3):482-487.
- Bag-Ozbek A, Giles JT. Inflammation, adiposity, and atherogenic dyslipidemia in rheumatoid arthritis: is there a paradoxical relationship? *Curr Allergy Asthma Rep.* 2015 Feb;15(2):497.
- Mackey RH, Mora S, Bertoni AG, et al. Lipoprotein particles and incident type 2 diabetes in the multi-ethnic study of atherosclerosis. *Diabetes Care.* 2015 Apr;38(4):628-636.
- Dugani SB, Akinkuolie AO, Paynter N, Glynn RJ, Ridker PM, Mora S. Association of lipoproteins, insulin resistance, and rosuvastatin with incident type 2 diabetes mellitus: Secondary analysis of a randomized clinical trial. *JAMA Cardiol.* 2016 May 1;1(2):136-145.
- Harada PHN, Demler OV, Dugani SB, et al. Lipoprotein insulin resistance score and risk of incident diabetes during extended follow-up of 20 years: The Women's Health Study. *J Clin Lipidol.* 2017 Sep-Oct;11(5):1257-1267.
- Flores-Guerrero JL, Connelly MA, Shalauova I, et al. Lipoprotein insulin resistance Index, a high-throughput measure of insulin resistance, is associated with incident type II diabetes in the Prevention of Renal and Vascular End-Stage Disease Study. *J Clin Lipidol.* 2019 Jan-Feb;13(1):129-137.e1.
- Flores-Guerrero JL, Oste MJC, Kieneker LM, et al. Plasma branched-chain amino acids and risk of incident type 2 diabetes: Results from the PREVENT Prospective Cohort Study. *J Clin Med.* 2018 Dec 4;7(12). pii: E513.
- Tobias DK, Mora S, Verma S, Lawler PR. Altered branched chain amino acid metabolism: toward a unifying cardiometabolic hypothesis. *Curr Opin Cardiol.* 2018 Sep;33(5):558-564.
- Ference BA, Ginsburg HN, Graham I, et al. Low-density lipoproteins cause atherosclerotic cardiovascular disease. 1. Evidence from genetic, epidemiologic, and clinical studies. A consensus statement from the European Atherosclerosis Society Consensus Panel. *Eur Heart J.* 2017 Aug;38(32):2459-2472.
- Cantey EP, Wilkins JT. Discordance between lipoprotein particle number and cholesterol content: an update. *Curr Opin Endocrinol Diabetes Obes.* 2018 Apr;25(2):130-136.
- Otvos JD, Shalauova I, Wolak-Dinsmore J, et al. GlycA: A composite nuclear magnetic resonance biomarker of systemic inflammation. *Clin Chem.* 2015 May;61(5):714-723.
- Ackerman RT. From programs to policy and back again: The push and pull of realizing type 2 diabetes prevention on a national scale. *Diabetes Care.* 2017 Oct;40(10):1298-1301.
- Nichols GA, Hillier TA, Brown JB. Normal fasting plasma glucose and risk of type 2 diabetes diagnosis. *Am J Med.* 2008 Jun;121(6):519-524.
- Shalauova I, Connelly MA, Garvey WT, Otvos JD. Lipoprotein insulin resistance index: a lipoprotein particle-derived measure of insulin resistance. *Metab Syndr Relat Disord.* 2014;12(8):422-429. doi: 10.1089/met.2014.0050
- Ellsworth DL, Costantino NS, Blackburn HL, Engler RJ, Kashani M, Vernalis MN. Lifestyle modification interventions differing in intensity and dietary stringency improve insulin resistance through changes in lipoprotein profiles. *Obes Sci Pract.* 2016 Sep;2(3):282-292.
- Fernandez-Castillejo S, Valls RM, Castaner O, et al. Polyphenol rich olive oils improve lipoprotein particle atherogenic ratios and subclasses profile: A randomized, crossover, controlled trial. *Mol Nutr Food Res.* 2016 Jul;60(7):1544-1554.
- Tuccinardi D, Farr OM, Upadhyay J, et al. Lorcaserin treatment decreases body weight and reduces cardiometabolic risk factors in obese adults: A six month, randomized, placebo controlled double-blind clinical trial. *Diabetes Obes Metab.* 2019 Jun;21(6):1487-1492.