

Diabetes Risk Index (DRI): Stratifying Risk Independent of Glycemic Status

The Diabetes Risk Index (DRI) measured on LabCorp's proprietary Vantera® platform, an automated nuclear magnetic resonance (NMR) clinical analyzer, combines selected lipoprotein and branched-chain amino acid (BCAA) parameters associated with insulin resistance into a clinically-actionable score (values 1-100) to help identify individuals with similar levels of glucose who differ in their risk of developing type 2 diabetes (T2D).

To help stem the growing epidemic of obesity and T2D, clinical practice guidelines recommend structured lifestyle modification and/or pharmacological intervention for patients with a high-risk glycemic status (ie, prediabetes as defined usually by fasting glucose = 100-125 mg/dL or HbA1c = 5.7-6.4%).^{1,2} Since >80 million U.S. adults qualify as "high-risk" by glycemic criteria, a need has been recognized for a more refined approach to risk stratification, to improve cost-effectiveness by directing treatment to the subset of prediabetes patients at highest risk.³ Waiting until the onset of prediabetes before initiating preventive measures may also be suboptimal, since many individuals with normal glucose levels progress to diabetes in a relatively short time period.⁴ The DRI test assesses a patient's degree of insulin resistance, the core pathophysiologic defect that with time can lead to hyperglycemia caused by impaired insulin secretion resulting from loss of pancreatic β -cell function and mass.^{5,6} By the time a patient reaches the threshold of prediabetes, up to 80% of β -cell function may already have been lost.^{5,6}

The DRI score is calculated from the patient's measured Lipoprotein

Insulin Resistance Index (LP-IR)⁷ plus the concentrations of two branched-chain amino acids, valine and leucine.⁸ LP-IR and BCAA values both have been shown in multiple prospective clinical studies to predict the development of T2D independent of the level of glycemia.⁹⁻¹⁴

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.¹⁵⁻¹⁷ Reductions of DRI and LP-IR are thus clinically achievable and likely to reflect a corresponding reduction of the risk of developing diabetes.

LabCorp offers the Diabetes Risk Index to aid clinicians with therapeutic decision-making based on a patient's risk of developing T2D independent of glycemic status.

Test Name	Test No.	
Diabetes Risk Index (DRI)	123855	
Cut Points	Men	Women
Low risk	< 50	< 40
Moderate risk	50 - 65	40 - 55
High risk	> 65	> 55
Methodology: Nuclear magnetic resonance (NMR)		
Platform: Vantera		

For the most current information regarding test options, including specimen requirements and CPT codes, please consult the online Test Menu at www.LabCorp.com.

References

- (3.) Prevention or delay of type 2 diabetes: Standards of Medical Care in Diabetes – 2019. *Diabetes Care*. 2019 Jan;42(Suppl 1):S29-S33.
- Garber AJ, Abrahamson MJ, Barzilay JI, et al. Consensus Statement by the American Association of Clinical Endocrinologists and American College of Endocrinology on the Comprehensive Type 2 Diabetes Management Algorithm – 2019 Executive Summary. *Endocr Pract*. 2019 Jan;25(1):69-100.
- 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.
- DeFronzo RA, Abdul-Ghani MA. Preservation of β -cell function: the key to diabetes prevention. *J Clin Endocrinol Metab*. 2011 Aug;96(8):2354-2366.
- DeFronzo RA. Banting Lecture. From the triumvirate to the ominous octet: a new paradigm for the treatment of type 2 diabetes mellitus. *Diabetes*. 2009 Apr;58(4):773-795.
- Shalaurova I, Connelly MA, Garvey WT, Ottvos JD. Lipoprotein insulin resistance index: a lipoprotein particle-derived measure of insulin resistance. *Metab Syndr Relat Disord*. 2014 Oct;12(8):422-429.
- Wolak-Dinsmore J, Gruppen EG, Shalaurova I, et al. A novel NMR-based assay to measure circulating concentrations of branched-chain amino acids: elevation in subjects with type 2 diabetes mellitus and association with carotid intima media thickness. *Clin Biochem*. 2018 Apr;54:92-99.
- 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, Shalaurova 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, Osté MCJ, Kiener 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.
- 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.
- Fernández-Castillejo S, Valls RM, Castañer 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.