

Diabetes-Associated Autoantibodies

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Diabetes mellitus (DM) refers to a group of metabolic disorders characterized by hyperglycemia that results from defects in insulin secretion, insulin action, or both. Type 1 DM (T1DM) is less common than type 2 DM (T2DM) and is characterized by insulin deficiency, which often results from the autoimmune-mediated destruction of insulin-producing cells. The detection of diabetes-associated autoantibodies confirms an autoimmune etiology for that individual.

Indications for Insulin Antibody Testing

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- Do not order tests individually; order tests for ≥2 antibodies if pursuing testing¹
- For most cases, order GAD in combination with ≥1 of the following antibodies: IA-2, IAA, ICA, ZnT8

T1DM	 Patient should have been previously diagnosed with DM Antibody testing is not useful for the diagnosis of DM Patients should be receiving insulin ≤2 wks, ideally Testing is not recommended for patients receiving insulin >2 wks, as insulin antibody formation may occur (false-positive result possible) Most useful in children or in adults without traditional risk factors for T2DM Traditional risk factors include excess weight/obesity (BMI ≥25 kg/m² or ≥23 kg/m² in Asian individuals), a first-degree relative with diabetes, being a high-risk race/ethnicity, physical inactivity, etc. For a full list of traditional risk factors, see Table 2.3 in the Standards of Medical Care in Diabetes² It may be useful in difficult adult cases to help differentiate between T1DM or T2DM ³
T2DM	No indication for routine evaluation or management ¹
Screening	Acceptable only for first-degree relatives of a proband with T1DM or in research settings
Limited use	 Differentiate LADA from T2DM⁴ Rule out autoantibodies as a cause of DM in patients with suspected genetic DM types (eg, monogenic DM, MODY)

GAD, glutamic acid decarboxylase antibody; IA-2, islet antigen-2; IAA, insulin antibody; ICA, islet cell cytoplasmic antibody; LADA, latent autoimmune diabetes of the adult; MODY, maturity-onset diabetes of the young; ZnT8, zinc transporter 8 antibody

Diabetes Mellitus Type 1 Overview

Prevalence

1.25 million in the United States

Featured ARUP Testing

Glutamic Acid Decarboxylase Antibody 2001771

Method: Semi-quantitative Enzyme-Linked Immunosorbent Assay

Use in combination with another insulin antibody test to determine autoimmune DM

Islet Antigen-2 (IA-2) Autoantibody, Serum 3001499

Method: Quantitative Enzyme-Linked Immunosorbent Assay

Useful to establish autoimmune etiology in previously diagnosed T1DM

Insulin Antibody 0099228

Method: Semi-Quantitative Radioimmunoassay

- Use to determine presence of antibodies to endogenous or exogenous insulin analogues
- Testing is not recommended for patients receiving insulin >2 weeks, as insulin antibody formation may occur

Islet Cell Cytoplasmic Antibody, IgG 0050138

Method: Semi-Quantitative Indirect Fluorescent Antibody

Useful to establish autoimmune etiology in previously diagnosed T1DM

Zinc Transporter 8 Antibody 2006196

Method: Semi-Quantitative Enzyme-Linked Immunosorbent Assay

Useful to establish autoimmune etiology in previously diagnosed $\mathsf{T1DM}$

Age of Onset

Most common in children but can develop in individuals of any age, especially in late 30s or early 40s

Symptoms

- Excessive thirst, hunger, and urination
- Fatigue, nausea, blurred vision
- Unexplained weight loss (obesity is rare at initial diagnosis)
- Possible co-occurring autoimmune disorders

Physiology

- · Caused by autoimmune-mediated destruction of insulin-producing beta cells of the islets of Langerhans in the pancreas
- · Five major autoantibodies of diagnostic interest:
 - GAD
 - IAA
 - IA-2
 - ICA
 - ZnT8
- · Antibodies may be present in individuals years before the onset of clinical symptoms.
- A presence in individuals with diabetes confirms an autoimmune etiology.

Test Interpretation

Sensitivity/Specificity

Moderate sensitivity, high specificity in newly diagnosed T1DM

- The presence of antibodies may decrease with long-term disease.
- Insulin antibody testing loses specificity once the patient has been on exogenous insulin for >2 weeks.

Results

- The presence of multiple insulin antibodies (GAD, IA-2, IAA, ICA, and ZnT8) is predictive of T1DM.
- If one autoantibody is found, others should be assayed; the risk of T1DM increases (>90%) if an individual tests positive for two or more autoantibodies.
- For further risk stratification, HLA-DR or HLA-DQ genotyping may be helpful.

Limitations

- Negative test results do not rule out autoimmune diabetes; autoantibody response varies by individuals.
- Presence of a single autoantibody in the absence of clinical symptoms has low predictive value (1-2% in healthy individuals).
- Not all individuals with antibodies will develop T1DM.
- Do not use to monitor or diagnose T1DM.
- IAA testing does not differentiate between antibodies specific for endogenous and exogenous forms of insulin.

References

- 1. Insel RA, Dunne JL, Atkinson MA, et al. Staging presymptomatic type 1 diabetes: a scientific statement of JDRF, the Endocrine Society, and the American Diabetes Association. *Diabetes Care*. 2015;38(10):1964-1974.
- 2. American Diabetes Association. Standards of care in diabetes-2023. Published Jan 2023; accessed Apr 2023.
- 3. Blonde L, Umpierrez GE, Reddy SS, et al. American Association of Clinical Endocrinology clinical practice guideline: developing a diabetes mellitus comprehensive care plan-2022 update. Endocr Pract. 2022;28(10):923-1049.
- 4. Pieralice S, Pozzilli P. Latent autoimmune diabetes in adults: a review on clinical implications and management. Diabetes Metab J. 2018;42(6):451-464.

Related Information

Diabetes Mellitus - Type 1, Type 2, and Gestational

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