PROVIDER Points

Comparing Tests for Diabetes and Prediabetes:
A Quick Reference Guide

This fact sheet compares the following tests:
- A1C test
- fasting plasma glucose (FPG) test
- oral glucose tolerance test (OGTT)
- random plasma glucose (RPG) test

In addition, the National Diabetes Education Program (NDEP) offers a pocket guide, Diabetes Numbers At-a-Glance, which can be ordered at www.ndep.nih.gov. Both resources utilize current American Diabetes Association (ADA) clinical recommendations for diagnosing and managing diabetes and prediabetes.

CONFIRMING DIAGNOSIS OF TYPE 2 DIABETES AND PREDIABETES
Diagnosis must be confirmed unless symptoms are present. Repeat the test using one of the following methods:
- Repeat the same test on a different day—preferred.
- If two different tests are used—e.g., FPG and A1C—and both indicate diabetes, consider the diagnosis confirmed.
- If the two different tests are discordant, repeat the test that is above the diagnostic cut point.

If diagnosis cannot be confirmed using the results of two tests, but at least one test indicates high risk, health care providers may wish to follow the patient closely and retest in 3 to 6 months.

INTERPRETING LABORATORY RESULTS
When interpreting laboratory results health care providers should:
- consider that all laboratory test results represent a range, rather than an exact number.
- be informed about the A1C assay methods used by their laboratory.
- send blood samples for diagnosis to a laboratory that uses an NGSP-certified method for A1C analysis to ensure the results are standardized.
- consider the possibility of interference in the A1C test when a result is above 15% or is at odds with other diabetes test results.
- consider each patient’s profile, including risk factors and history, and individualize diagnosis and treatment decisions in discussion with the patient.
### COMPARING DIABETES BLOOD TESTS‡

<table>
<thead>
<tr>
<th>Test</th>
<th>Uses</th>
<th>Technical Features</th>
<th>PROS</th>
<th>CONS</th>
</tr>
</thead>
</table>
| **A1C Test** | • Screening and diagnosis of prediabetes  
   - ≥ 5.7–6.4%†  
   • Screening and diagnosis of type 2 diabetes  
   - ≥ 6.5%†  
   • Monitoring of diabetes | • Diagnosis requires a laboratory test certified by the NGSP; not meter—point-of-care A1C tests are only suitable for monitoring  
   • Sample any time of day, no fasting  
   • Sample: unanticoagulated whole blood  
   • Sample stability: ≤ 7 days  
   • Sensitivity: ≤ 8.0%  
   • Coefficient of variation: assay variability, see www.ngsp.org | • Reflects long-term blood glucose concentration  
   • Unaffected by acute changes in glucose levels due to stress or illness  
   • Highly correlated with risks for complications such as retinopathy and cardiovascular disease (CVD)  
   • Convenient for patient and health care providers  
   • Most stable sample after collection  
   • Low within-patient variability  
   • Established international standardization of lab tests  
   • Accuracy of test is monitored | • Lower sensitivity: identifies fewer cases of diabetes than the glucose tests  
   • Interference resulting in falsely increased or lowered results due to*  
   - generic variants including HbS, HbC, HbE, and HbH traits and HbF  
   - affects people of African, Mediterranean, and Southeast Asian heritage  
   - kidney disease  
   - liver disease  
   - iron deficiency anemia  
   - heavy bleeding  
   • Not recommended for rapidly progressing diabetes, e.g., type 1 diabetes in children  
   • May not be available in some laboratories/areas of the world  
   • Expensive  
   *See www.ngsp.org for information on A1C interference and recommended testing overlays.  
   **See the NDDG publication The A1C Test and Diabetes at www.diabetes.niddk.nih.gov.  
| **FPG Test** | • Screening and diagnosis of prediabetes or impaired fasting glucose (IFG)  
   - 100–125 mg/dL†  
   • Screening and diagnosis of diabetes  
   - ≥ 126 mg/dL†  
   • repeat for confirmation of diagnosis | • Diagnosis requires a laboratory test, not meter  
   • Sample in morning, after 8-hour fast  
   • Sample: sodium fluoride plasma preferred  
   • Sample stability: low—processing within 30 minutes  
   • Sensitivity: greater than the A1C test, less than the OGTT  
   • Coefficient of variation: assay variability: | • Low cost  
   • Assay is widely available  
   • Assay is automated | • Indicates single-point blood glucose level  
   • Affected by short-term lifestyle changes: stress or illness  
   • Less tightly linked to diabetes complications than A1C  
   • Not convenient for patient or health care provider: requires fasting and scheduling a morning appointment or return visit  
   • Diurnal variation  
   • Sample not stable after collection  
   • High within-patient variability  
   • Many laboratories measure serum, which is not recommended  
   • Inadequate standardization of assays  
| **OGTT** | • Screening and diagnosis of prediabetes or impaired fasting glucose tolerance (IGT)  
   - 140–199 mg/dL at 2 hr†  
   • Screening and diagnosis of diabetes  
   - ≥ 200 mg/dL at 2 hr†  
   • repeat for confirmation of diagnosis  
   • Screening and diagnosis of gestational diabetes mellitus (GDM)* | • Sample in morning, after 8-hr. fast and 2 hrs. after glucose load  
   • Sample stability: low—requires processing within 30 minutes  
   • Patients should ingest at least 150 g/day of carbohydrates for 3 days prior  
   • Sensitivity: greater than the A1C or the FPG tests | • Sensitive indicator of risk of developing diabetes  
   • Early marker of impaired glucose balance | • Affected by short-term lifestyle changes: stress, illness, and medications  
   • Not convenient for patient or health care provider: requires fasting and scheduling a morning appointment or return visit  
   • Extensive patient preparation  
   • Sample not stable after collection  
   • High within-patient variability  
   • Low reproducibility  
   • Expensive  
| **RPG Test** | • Diagnosis of diabetes—used only with classic symptoms of hyperglycemia or hyperglycemic crisis:  
   - polyuria, polydipsia, and unexplained weight loss  
   - 200 mg/dL† | • Sample any time, no fasting  
   • Sample stability: low—requires processing in fewer than 2 hours | • Convenient  
   • Part of basic metabolic panel screen | • Indicates single-point blood glucose level  
   • Used only in symptomatic patients, not recommended for screening  
   • Insensitive measurement  
   • Greater within-patient variability  
   • Affected by short-term lifestyle changes and prandial state  

*Testing for GDM is not covered in this publication.  

Source: Courtesy of David Aron, M.D., Louis Stokes Department of Veterans Affairs Medical Center
COMPARING DIAGNOSES

In some people, a blood glucose test may indicate a diagnosis of diabetes even though an A1C test does not. The reverse can also occur—an A1C test may indicate a diagnosis of diabetes even though a blood glucose test does not.

Because of these variations in test results, health care providers should repeat tests before making a diagnosis. People with differing test results may be in an early stage of the disease, where blood glucose levels have not risen high enough to show on every test.

REFERENCES

3. See www.ngsp.org for information on A1C test interference and recommended testing methods.

ACKNOWLEDGMENTS

Publications produced by the Clearinghouse are carefully reviewed by both NIDDK scientists and outside experts. This publication was originally reviewed by Randie Little, Ph.D., University of Missouri-Columbia and David B. Sacks, M.B., Ch.B., FRCPATH, NIH Clinical Center. The updated version of this publication was reviewed by David Aeron, M.D., Louis Stokes Department of Veterans Affairs Medical Center. This publication is published by the NIDDK in collaboration with the NDEP. This publication is available at www.diabetes.niddk.nih.gov.

Another quick-reference tool, Diabetes Numbers At-a-Glance 2012, is offered by the NDEP and is available at www.ndep.nih.gov.