Diabetes in Clinical Practice

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Author biographies

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Dr Fonseca previously served as Chair for the American Diabetes Association (ADA) Clinical Practice Committee and the joint ADA/American College of Cardiology (ACC) "Make the Link" program. He is Editorin-Chief of Diabetes Care, and was Associate Editor of the Journal of the Metabolic Syndrome and Related Disorders and is also on the editorial board of the Journal of Clinical Endocrinology and Metabolism. He is an ad hoc reviewer for several other journals, including Diabetes, Diabetic Medicine, Kidney International, American Journal of Clinical Nutrition, Journal of Clinical Endocrinology and Metabolism, British Medical Journal, JAMA, and Metabolism.

Dr Fonseca is a fellow of the American Association of Clinical Endocrinologists, the Royal College of Physicians (London), and the American College of Physicians. He is a member of the Endocrine Society, the American Diabetes Association, and the International Diabetes Federation. He has lectured in the USA and abroad, serves on several national and international committees, and has published more than 100 papers, review articles, and book chapters.

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Dr Pendergrass is a reviewer for a number of journals including *Diabetes* Care, Metabolic Syndrome and Related Disorders, and New England Journal

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Chapter 1

1

Type 2 diabetes: the modern epidemic

Type 2 diabetes is a major clinical and public health problem. It is estimated that in the year 2000, 171 million people worldwide had type 2 diabetes, including about 18 million Americans, and it is estimated that these numbers will grow to 366 million people worldwide and 30 million Americans by the year 2030.

This high prevalence of diabetes leads to a high global burden of the condition and its complications. Diabetes is the leading cause of blindness below the age of 65 years, and it is responsible for almost half the cases needing dialysis. It is responsible for most non-traumatic amputations. The financial cost of this condition is staggering, with one in seven healthcare dollars in the USA spent on treating the condition or its complications. Indeed, despite the fact that only about 10–15% of the Medicare population has diabetes, about 25% of the American Medicare budget is spent on this condition. In addition, there is a considerable expenditure on the social costs involved with people who suffer with long-term complications, including disability and premature death.

The purpose of this handbook is to give the practitioner a quick overview of the subject, along with practical suggestions for the management of this condition.

Diagnosing diabetes

Criteria for the diagnosis of diabetes

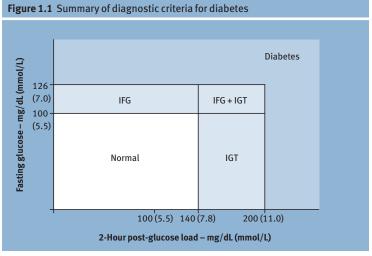
- Symptoms of diabetes (polyuria, polydipsia, unexplained weight loss) plus random plasma glucose concentration >200 mg/dL (11 mmol/L).
- Fasting plasma glucose (FPG) >126 mg/dL (7 mmol/L). (Fasting = no caloric intake for at least 8 h.)
- Two-hour plasma glucose >200 mg/dL (11 mmol/L) during an oral glucose tolerance test (OGTT) (75 g).

In the absence of unequivocal hyperglycemia with acute metabolic decompensation, these criteria should be confirmed by repeat testing on a different day. The OGTT is not recommended for routine clinical use.

These diagnostic criteria are summarized in Figure 1.1.

The diagnosis of diabetes is currently made on the basis of several diagnostic criteria (summarized above) [1]. In patients who have classic symptoms, such as polyuria, polydipsia, and weight loss, a random plasma glucose greater than 200 mg/dL (11 mmol/L) is diagnostic. However, most patients are asymptomatic and those suspected of having diabetes should be screened with either a fasting glucose or a 75 g glucose tolerance test. An FPG greater than or equal to 126 mg/dL (7 mmol/L) is considered diagnostic of diabetes, but should be confirmed on another occasion in asymptomatic patients. Following an OGTT, a value greater than 200 mg/dL (11 mmol/L) is considered diagnostic. However, values below these figures are not entirely normal, as a normal fasting glucose is less than 100 mg/dL (5.5 mmol/L), and therefore fasting glucose between $100\,mg/dL$ (5.5 mmol/L) and $125\,mg/dL$ (7 mmol/L) is diagnostic of impaired fasting glucose (IFG), and a 2-hour post-glucose load of 140-199 mg/dL (7.8-11.0 mmol/L) is called impaired glucose tolerance (IGT). Both of these conditions predict increased risk of subsequent progression to diabetes, and also of macrovascular complications, even in the absence of progression to diabetes. The latter is particularly true in patients with IGT.

It is also relatively easy to identify people who are at increased risk of diabetes, who should therefore be screened early even if asymptomatic in order to start treatment early, or take preventive steps described below should they have IFG or IGT.



IFG, impaired fasting glucose; IGT, impaired glucose tolerance