

Definition, Classification and Diagnosis of Diabetes Mellitus

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ABSTRACT

Aim of recommendations like this one issued by the German Diabetes Association is to provide the GP and diabetologist and his team with information he needs for his daily practice. These recommendations are updated annually. They are written by a group of experts, but they are not evidence based guidelines. This specific recommendation for diabetes diagnosis briefly describes the diabetes types and the different options for diagnosis. Also the caveats and the practical procedure are presented.

Definition of Diabetes mellitus

Diabetes mellitus is a general term for a group of metabolic disorders with the main feature of chronic hyperglycaemia. It results from either impaired insulin secretion or impaired insulin efficacy or, most often, both.

Gestational Diabetes

A glucose utilization disorder which occurs and is diagnosed during pregnancy for the first time.

Classification

1. Type 1 diabetes
 - β cell destruction which leads to an absolute insulin deficiency in most cases resulting from immunological factors
 - LADA (latent autoimmune diabetes in adults)
2. Type 2 diabetes
 - can range from a predominant insulin resistance with relative insulin deficiency to an extensive secretory defect with insulin resistance.

- is often associated with other disorders (e. g. metabolic syndrome).
3. Specific diabetes types with known causes
 - Diseases involving a pancreatic exocrine deficiency (e. g. pancreatitis, cystic fibrosis, hemochromatosis)
 - Endocrinopathies (e. g. Cushing syndrome, acromegaly, pheochromocytoma)
 - Drug or chemically-induced (e. g. glucocorticoids, neuroleptics, interferon alpha, pentamidine),
 - Genetic defects of the β cell function (e. g. MODY types)
 - Genetic defects of insulin action
 - Other genetic syndromes which can be associated with diabetes
 - Infections
 - Rare forms of autoimmune-mediated diabetes
 4. Gestational diabetes

Diagnostic criteria for Diabetes Mellitus

- Random plasma glucose value of ≥ 200 mg/dl (≥ 11.1 mmol/l) or

- Fasting plasma glucose value of ≥ 126 mg/dl (≥ 7.0 mmol/l) or
- 2-h oral glucose tolerance test (oGTT) value in venous plasma ≥ 200 mg/dl (≥ 11.1 mmol/l)
- HbA1c $\geq 6.5\%$ (≥ 48 mmol/mol Hb)

Impaired fasting glucose values

IFG (impaired fasting glucose) for the range of fasting glucose from 100–125 mg/dl (5.6–6.9 mmol/l) in venous plasma.

Impaired glucose tolerance

IGT (impaired glucose tolerance) corresponds to a 2-h plasma glucose value in oGTT in the range from 140–199 mg/dl (7.8–11.0 mmol/l) with fasting glucose values < 126 mg/dl (< 7.0 mmol/l).

IFG and IGT are also present in many people with a glucose utilization disorder.

Diagnostic approach

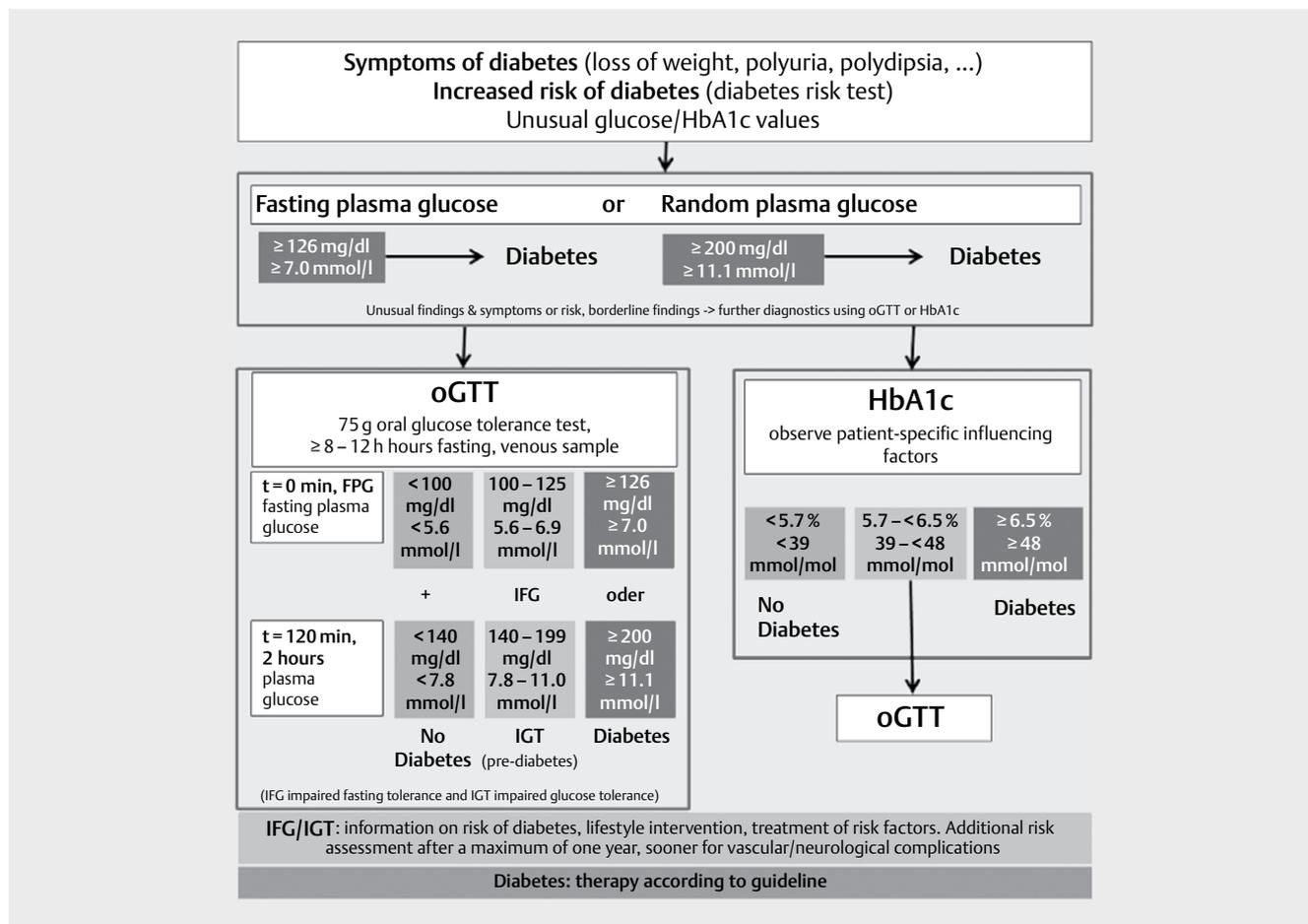
The recommended diagnostic procedure is shown in ► **Fig. 1** and the differential diagnostic approach criteria are shown in ► **Table 1**.

Only quality-assured laboratory methods may be used when measuring venous plasma glucose and HbA1c in diabetes diagnosis. This is defined in the guidelines published by the ‘Richtlinie der

Bundesärztekammer zur Qualitätssicherung laboratoriumsmedizinischer Untersuchungen (Rili-BAEK)’ (Guideline of the German Medical Association on Quality Assurance in Medical Laboratory Examinations) as standard for both central laboratories and point-of-care testing (POCT). Participation in external quality assessment (EQA) schemes has not been compulsory for POCT methods used in practices in Germany. If POCT systems are approved for diagnostic purposes by the manufacturer, the successful participation in EQA schemes, however, is mandatory before they can be used in diagnostics. Currently, the gold standard for the diagnosis of diabetes is the measurement of glucose in venous plasma.

Procedures if measurement results are close to the diagnostic criteria levels

When using the fasting glucose value, a fasting time of 8 to 12 h is essential. The guidelines for performing an oGTT must be followed (► **Table 2**). Using the HbA1c value for diagnosis is currently not recommended, in particular, because the allowed deviation for external quality control tests is $\pm 18\%$ to date in Germany. This value will be reduced to $\pm 8\%$ in the next years in the Rili-BAEK. This will improve the diagnostic value of HbA1c. The sensitivity of the laboratory tests for diagnosing diabetes is specified below in ascending order in relation to the oGTT, i. e. an oGTT should be performed to



► **Fig. 1** Diagnostic approach for diagnosing diabetes.

► **Table 1** Differential diagnostic criteria for patients with type 1 or type 2 diabetes for diagnosing diabetes. (Table modified from the National Medical Guideline for type 2 diabetes; www.versorgungslinien.de).

	Type 1 diabetes ¹	Type 2 diabetes	MODYs
Aetiology	Autoimmune, genetic predisposition	Genetic predisposition, multifactorial	Monogenic
Heredity	Variable	Variable	Autosomal dominant; diabetes in ≥ 3 generations
Percentage of all diabetes types	5 – 10%	90 – 95%	ca. 2%
Pathogenesis	Autoantibodies, absolute insulin deficiency	Insulin resistance and insulin secretion disorder up to absolute insulin deficiency	Mutation of genes of transcription factors or glucokinase of the β cells
Typical age of manifestation	Childhood to adulthood	Adulthood	Youth to early adulthood
Clinical manifestation	Acute polyuria, polydipsia, severe hyperglycaemia, ketoacidosis	Gradual onset, secondary complications at diagnosis, moderate to severe hyperglycaemia	Gradual onset, variable hyperglycaemia
Comorbidities	Autoimmune thyroiditis, celiac disease	Visceral obesity, hypertension, dyslipidemia (also called metabolic syndrome)	Kidney cysts according to MODY type
Tendency toward ketosis prone	Yes	No	No
Weight	Normal weight	Overweight, obese	Normal weight
Plasma insulin / C-peptide	Strongly decreased, mostly absolute deficiency	Often elevated at the beginning, then decreased	Mostly decreased
Autoantibodies	Yes	No	No
Insulin resistance	No	Yes	No
Therapy	Insulin	Lifestyle intervention, oral antidiabetics, GLP-1-RA, insulin	Maybe none, OADs, insulin (according to MODY type)

¹LADA (latent autoimmune diabetes in adults) is associated with a slow loss of β cell function. A rapid failure of oral antidiabetics can be expected with LADA. Upon suspicion of LADA: test for GAD-antibodies is recommended

rule-out diabetes if the HbA1c values and fasting glucose values lie closely below the diagnostic criteria.

How should the diagnostic value of a single measurement be assessed in consideration of the variability of measurement results?

For the results of these measurands the question is whether the deviation from the diagnostic criterion (or cut-off value) exceeds this value to such an extent (i. e. larger than the minimal difference (see below)) that this can clearly be considered elevated. If this is the case, one individual measurement is sufficient to confirm diabetes diagnosis. For two values diverging the cut-off value (above and below), the American Diabetes Association (ADA) recommends considering the higher value. This value should be repeated and becomes the deciding factor for the diagnosis. If clinically necessary, verification should be performed in time, e. g. after three to six months. The procedure for diagnosing diabetes (as published by the ADA 2018) is as follows: 'If a parameter is used and then repeated to confirm the diagnosis of diabetes, it must be ensured that each blood sample was standardized and comparably processed.' How well this is realized for plasma glucose is not simply verifiable. If the diabetes diagnosis is based on an HbA1c measurement, the confirmation with the same measurand is not advisable. Due to analytical differences per laboratory the HbA1c measurement is currently not very well reproducible. Additionally, a "false" value may be obtained repeatedly due to the same patient-specific influencing factors. Independent of the measurand used for the

► **Table 2** Oral glucose tolerance test (oGTT).

75 g oGTT in accordance with WHO guidelines
Performing the test in the morning <ul style="list-style-type: none"> – after 8 – 12 h fasting from food, nicotine and fluid including alcohol – after a ≥ 3-day diet rich in carbohydrates (≥ 150 g CHO per day) – sitting or lying (no muscular exertion); no smoking before or during the test
At time 0 drink 75 g glucose (or equivalent amount of hydrolysed starch) in 250 – 300 ml of water within 5 min <ul style="list-style-type: none"> – children 1.75 g/kg (maximal 75 g) – venous blood collection at time 0 and 120 min – proper sample processing and storage
The test is contraindicated for intercurrent diseases, such as status after gastrointestinal resection or gastrointestinal illness with changed resorption or if diabetes mellitus has already been diagnosed.
Preparation of the glucose solution by the physician instead of the manufacturer is rejected by the DDG for reasons of liability and for medical reasons; see statement by KLD and AGDT on the DDG website (www.deutsche-diabetes-gesellschaft.de).

diagnosis of diabetes, the result may be incorrect resulting from patient-specific influencing factors and/or insufficient measurement accuracy. A questionable diabetes diagnosis should therefore be confirmed using the other measurand (i. e. either glucose or HbA1c) in order to reduce interference or influencing factors. A measurement result which is used as a basis for diagnosis should be confirmed so that the diagnosis is made on the basis of confirmed values. It can be confirmed either by analysing a new blood sample using the same measurand (e. g. within 14 days) or by using

► **Table 3** Blood collection tubes commercially available in Germany which completely inhibit glycolysis with the addition of fluoride and citrate. (current status 17.7.2017, see home pages of the respective manufacturers).

Manufacturer	Product name	Correct filling absolutely necessary	Sufficient mixing required	Correction factor
Greiner bio-one	Vacurette® FC-Mix	no	10 times	no (powder)
Kabe	Primavette®, KABEVETTE®	yes	few times	1.16 (liquid additive)
Sarstedt	S-Monovette GlucoEXACT®	yes	few times	1.16 (liquid additive)

The blood collection tubes made by the company Greiner bio-one (Vacurette® FC-Mix) contain granulate. The tubes must be gently inverted ten times to adequately dissolve the additive and mix the sample with the glycolytic inhibitors. Experience has shown that dilution errors occur when using blood collection tubes by the company Sarstedt (S-Monovette® GlucoEXACT) and Kabe (Primavette®, KABEVETTE®) if the tubes have not been filled completely. The laboratory has to identify such types of tubes to, on the one hand, identify tubes that do not meet the filling requirements specified by the manufacturer and exclude them from the analysis, and, on the other hand, take the dilution factor of 1.16 into account.

► **Table 4** Contributing factors leading to an influence or falsification of the HbA1c value.

1. Haemoglobin variants (HbS, HbE, HbF, HbC, HbD etc.) – the extent of the distortion depends on the measurement methods used.
2. Conditions with increased or lower life expectancy of erythrocytes (haemolytic anaemia, iron deficiency anaemia, new blood cell formation as part of anaemia treatment, status after splenectomy or illnesses of the spleen, liver or kidney)
3. Chemical modifications of haemoglobin
4. Uraemia (carbamyated Hb), high dosed long-term therapy with acetylsalicylic acid (acetylated Hb)
5. Inhibition of glycolysis (e. g. long-term therapy with ascorbic acid or vitamin E). The clinical relevance of this phenomenon has not been investigated adequately.
6. Pregnancy
7. Ethnicity and age (HbA1c increases with age, therefore a potential age adjustment of the diagnostic criterion must be discussed and evaluated). Moreover, discussion should be extended to the possible alternative measurands such as fructosamine or glycated albumin)

one of the three measurands described above. If, during a second determination of the same laboratory test, there is a discrepancy in the value with regard to the cut-off value, one of the alternatives should be used for determination. If there are disparities in the results with regards to the diagnostic criterion in two different measurands, the higher value should be confirmed. If the values are close to the limit values, they should be monitored within 3 – 6 months again.

Pre-analytics of glucose measurement

Proper pre-analytic handling of the blood sample is crucial. Suitable blood collection tubes must be used to completely inhibit glycolysis in the collected blood. The addition of citrate and fluoride is necessary since fluoride alone is insufficient. The glycolytic inhibitor tubes for blood collection currently available on the German market exhibit different handling problems (► **Table 3**). One alternative recommendation for tubes without glycolytic inhibitor is to centrifuge the sample as soon as possible after blood collection. If a period of 30 min to centrifugation is exceeded, samples should be discarded due to occurring glycolysis. After centrifugation, the resulting plasma supernatant must be separated from the blood cells. This occurs during centrifugation using a gel (gel tubes) or

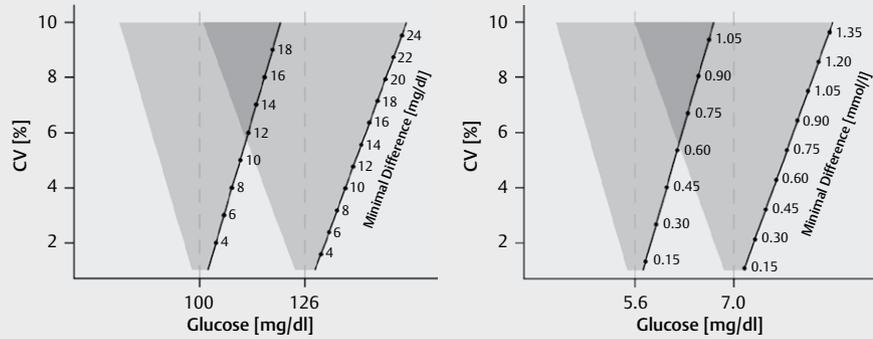
mechanical separators. It is also possible to pipette off the plasma supernatant immediately after centrifugation. Diligent and optimal pre-analytical handling of the blood collection tubes can result in a higher rate of diabetes diagnosis in practice and should not be considered over-diagnosing; however, the cut-off values used below may require adjustment and should be verified with corresponding studies.

HbA1c for diagnosis purposes

Since 2010, the German Diabetes Association ('Deutsche Diabetes Gesellschaft', DDG) has recommended the use of the HbA1c value for the diagnosis of diabetes (see the statement on the DDG homepage). This was made possible by improvements in measurement accuracy resulting from international standardization of HbA1c analyses. Simultaneously, epidemiological studies in recent years have shown that the specificity of a HbA1c value of $\geq 6.5\%$ or ≥ 48 mmol/mol is sufficiently high to diagnose diabetes with satisfactory level of certainty. At the same time, the sensitivity of an HbA1c value of $< 5.7\%$ (< 39 mmol/mol Hb) is sufficiently high to rule-out the diagnosis of diabetes. For patients with HbA1c values in the range of 5.7 to $< 6.5\%$ (39 to < 48 mmol/mol Hb) or with a high clinical risk (see Screening), the diagnosis of diabetes and its early stages can only be excluded by measuring plasma glucose in accordance with the usual criteria including an oGTT. The HbA1c value is not suitable for diagnosing diabetes if values can be expected to be influenced or falsified (► **Table 4**). It is also important to note that regardless of successful standardization, measurement accuracy of HbA1c can vary considerably depending on the measurement method. In our opinion, this problem significantly limits the exclusive use of HbA1c for the diagnosis of diabetes ► **Table 4** in the legend to ► **Fig. 1**. In particular, the increase of HbA1c that is associated with increasing age independent of diabetes, which can be up to an absolute value of 0.4 – 0.7% (4 – 8 mmol/mol Hb) constitutes another limitation in addition to differences due to methodology for using HbA1c to diagnose diabetes especially in the range below 7.0% (53 mmol/mol Hb).

Quality control checks

Internal quality control must be performed every working day with suitable control material. Successful participation in external quality assurance is required once quarterly. This applies to all labora-



► **Fig. 2** Minimal Difference, given in the unit of glucose determination (mg/dL or mmol/L) for the considered diagnostic cut-off value depending on the coefficients of variation. If the measured values lie in the overlapping areas of the inverted triangles, the diagnostic limit values cannot be differentiated from each other and the measured values cannot be used for diagnosis.

tory systems as well as to POCT “unit use” systems (single test strips or cuvettes as per the definition of Rili-BAEK) which are designated appropriate for diagnosis by the manufacturer.

Minimal difference

In order to meet the clinical requirements, the analytical variability of the absolute values at decision limits should be specified. The “Minimal Difference (MD)” is a simple tool to demonstrate the significance of random errors to the user and is calculated from the standard deviation (SD) ($MD = 2 \times SD$) (► **Fig. 2**). The MD gives specific concentrations as absolute values to specify if a measurement result differs from a cut-off value with diagnostic relevance. The MD should not be larger than 12.6 mg/dL (0.7 mmol/L) for a fasting glucose limit value of 126 mg/dL (7.0 mmol/L). Accordingly, for the HbA1c cut-off value of 48 mmol/mol Hb the MD should not be larger than 1.9 mmol/mol Hb. A statement on the subject of MD can be found on the DDG homepage and will be published separately soon.

Screening

For primary screening for diabetes either a ‘Diabetes Risk Test’ is recommended (http://www.dife.de/de/presse/Diabetes_Test_Fragebogen.pdf) or occasional measurement results using venous plasma. In case of elevated risk scores detected by this questionnaire, documented cardiovascular disease, presence of excess weight with additional risk factors such as hypertension, dyslipidaemia (increased triglyceride values or low HDL cholesterol values) or a positive family history of type 2 diabetes in first-degree relatives, gestational diabetes or polycystic ovarian (PCO) syndrome or non-alcoholic fatty liver disease, proceed as described in ► **Fig. 1**. Alternatively, the FINDRISK questionnaire can also be used (www.diabetestiftung.de).

► **Table 5** Diagnosis of gestational diabetes. **Diabetes is confirmed if one criterion is met.** For the pre-analytics of glucose determination, reference is made to the guideline for gestational diabetes; a sufficient inhibition of glycolysis is necessary.

	venous plasma	
	mg/dL	mmol/l
fasting	≥ 92	≥ 5.1
60 min	≥ 180	≥ 10.0
120 min	≥ 153	≥ 8.5

Gestational diabetes

The cut-off values for oGTT stated in ► **Table 5** are based on the results of the HAPO Study. They differ only slightly from the previously used decision limits. However, now already one value exceeding the decision limit suffices to confirm the diagnosis while previously two elevated values were required.

Internet addresses

<http://www.deutsche-diabetes-gesellschaft.de>

- Current version of evidence-based guidelines <http://www.deutsche-diabetes-gesellschaft.de>

Initial Publication

This Clinical Practice Guideline is a translation of: Nauck M, Petermann A, Müller-Wieland D, et al. Definition, Klassifikation und Diagnostik des Diabetes mellitus. Diabetologie 2017; 12 (Suppl 2): S94–S100. DOI: 10.1055/s-0043-115953