# Table of Contents

<table>
<thead>
<tr>
<th>Section</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Summary</strong></td>
<td>3</td>
</tr>
<tr>
<td><strong>Basics</strong></td>
<td>4</td>
</tr>
<tr>
<td>Definition</td>
<td>4</td>
</tr>
<tr>
<td>Epidemiology</td>
<td>4</td>
</tr>
<tr>
<td>Etiology</td>
<td>4</td>
</tr>
<tr>
<td>Pathophysiology</td>
<td>4</td>
</tr>
<tr>
<td>Classification</td>
<td>5</td>
</tr>
<tr>
<td><strong>Prevention</strong></td>
<td>6</td>
</tr>
<tr>
<td>Screening</td>
<td>6</td>
</tr>
<tr>
<td><strong>Diagnosis</strong></td>
<td>7</td>
</tr>
<tr>
<td>Case history</td>
<td>7</td>
</tr>
<tr>
<td>Step-by-step diagnostic approach</td>
<td>7</td>
</tr>
<tr>
<td>Risk factors</td>
<td>8</td>
</tr>
<tr>
<td>History &amp; examination factors</td>
<td>8</td>
</tr>
<tr>
<td>Diagnostic tests</td>
<td>9</td>
</tr>
<tr>
<td>Differential diagnosis</td>
<td>10</td>
</tr>
<tr>
<td>Diagnostic criteria</td>
<td>10</td>
</tr>
<tr>
<td><strong>Treatment</strong></td>
<td>12</td>
</tr>
<tr>
<td>Step-by-step treatment approach</td>
<td>12</td>
</tr>
<tr>
<td>Treatment details overview</td>
<td>17</td>
</tr>
<tr>
<td>Treatment options</td>
<td>18</td>
</tr>
<tr>
<td>Emerging</td>
<td>24</td>
</tr>
<tr>
<td><strong>Follow up</strong></td>
<td>26</td>
</tr>
<tr>
<td>Recommendations</td>
<td>26</td>
</tr>
<tr>
<td>Complications</td>
<td>27</td>
</tr>
<tr>
<td>Prognosis</td>
<td>31</td>
</tr>
<tr>
<td><strong>Guidelines</strong></td>
<td>33</td>
</tr>
<tr>
<td>Diagnostic guidelines</td>
<td>33</td>
</tr>
<tr>
<td>Treatment guidelines</td>
<td>33</td>
</tr>
<tr>
<td><strong>Online resources</strong></td>
<td>35</td>
</tr>
<tr>
<td><strong>References</strong></td>
<td>36</td>
</tr>
<tr>
<td><strong>Disclaimer</strong></td>
<td>45</td>
</tr>
</tbody>
</table>
Type 1 diabetes mellitus is a metabolic disorder characterized by hyperglycemia due to absolute insulin deficiency.

Patients most often present with a few days or weeks of polyuria, polydipsia, weight loss, and weakness.

Some patients may present with diabetic ketoacidosis.

Intensive glycemic control has been shown to decrease the incidence of microvascular and macrovascular complications.

Microvascular complications include retinopathy, nephropathy, and neuropathy.

Macrovascular complications include coronary artery, cerebrovascular, and peripheral vascular disease.
Definition
Type 1 diabetes mellitus is a metabolic disorder characterized by hyperglycemia due to absolute insulin deficiency. The condition develops due to destruction of pancreatic beta cells, mostly by immune-mediated mechanisms. In some patients there may be no evidence of autoimmune destruction of pancreatic beta cells; this is called idiopathic type 1 diabetes.

Epidemiology
Type 1 diabetes accounts for about 5% to 10% of all patients with diabetes. It is estimated that 1,106,500 people ages 0 to 19 years have type 1 diabetes worldwide, with 132,600 newly diagnosed cases each year.[3]

In the US from 2011 to 2012, over 17,900 people younger than 20 years were newly diagnosed with type 1 diabetes annually (annual rate for new cases about 21 per 100,000).[4] In addition, 10% of adults who have been classified as having type 2 diabetes may have circulating islet cell antibodies or antibodies to glutamic acid decarboxylase, indicating autoimmune destruction of beta cells.[5]

There is significant geographical variation in the incidence of type 1 diabetes. It is more common in Europeans and less common in Asians. Thus, a child in Finland is 40 times more likely to develop type 1 diabetes than a child in Japan and almost 100 times more likely to get the disease than a child in the Zunyi region of China.[6] Worldwide, the incidence of type 1 diabetes is increasing by 3% every year, although the reasons for this are unclear.[7] One 2017 report showed a more rapid increase in nonwhite racial and ethnic groups.[8]

Etiology
Certain human leukocyte antigen (HLA)-DR/DQ gene polymorphisms, particularly HLA-DR and HLA-DQ alleles, increase susceptibility to, or provide protection from, the disease.[9] In susceptible individuals, environmental factors may trigger the immune-mediated destruction of pancreatic beta cells. Although the geographic variation in disease prevalence and increasing worldwide incidence of type 1 diabetes argue for a major environmental contribution to pathogenesis, the specific factors involved remain unknown. Among viruses, the strongest associations have been found with congenital rubella syndrome and human enteroviruses.[10][11][12] Among dietary factors, infant supplementation with vitamin D may be protective.[13] Further research is required to determine whether cow’s milk, early introduction of cereals, or maternal vitamin D ingestion increase type 1 diabetes risk.[14][15][16][17][18][19] Celiac disease shares the HLA-DQ2 genotype with type 1 diabetes, and is more common among those with type 1 diabetes.[20] The incidence of type 1 diabetes may also be higher among those with celiac disease, although a causal relationship is not suggested.[21]

Pathophysiology
Type 1 diabetes usually develops as a result of autoimmune pancreatic beta-cell destruction in genetically susceptible individuals. Up to 90% of patients will have autoantibodies to at least one of 3 antigens: glutamic acid decarboxylase; insulin; and a tyrosine-phosphatase-like molecule, islet autoantigen-2 (IA-2).[22] Over 25% of individuals without one of these or islet cytoplasmic autoantibodies will have positive antibodies to ZnT8, a pancreatic beta-cell-specific zinc transporter.[23]
Beta-cell destruction proceeds subclinically for months to years as insulitis (inflammation of the beta cell). When 80% to 90% of beta cells have been destroyed, hyperglycemia develops. Insulin resistance has no role in the pathophysiology of type 1 diabetes. However, with increasing prevalence of obesity, some patients with type 1 diabetes may be insulin resistant in addition to being insulin deficient.

Patients with insulin deficiency are unable to utilize glucose in peripheral muscle and adipose tissues. This stimulates the secretion of counter-regulatory hormones such as glucagon, epinephrine, cortisol, and growth hormone. These counter-regulatory hormones, especially glucagon, promote gluconeogenesis, glycogenolysis, and ketogenesis in the liver. As a result, patients present with hyperglycemia and anion gap metabolic acidosis.

Long-term hyperglycemia leads to vascular complications due to a combination of factors that include glycosylation of proteins in tissue and serum, production of sorbitol, and free radical damage. Microvascular complications include retinopathy, neuropathy, and nephropathy. Macrovascular complications include cardiovascular, cerebrovascular, and peripheral vascular disease. Hyperglycemia is known to induce oxidative stress and inflammation. Oxidative stress can cause endothelial dysfunction by neutralizing nitric oxide. Dysfunctional endothelium allows entry of low-density lipoprotein into the vessel wall, which induces a slow inflammatory process and leads to atheroma formation.[24]

**Classification**

**Types of type 1 diabetes**

**Autoimmune or classical**

- Characterized by absolute insulin deficiency and the presence of antibodies to pancreatic beta cells.

**Idiopathic**

- Uncommon form that is characterized by absence of antibodies.
- Increased likelihood in patients of African or Asian ancestry and has a strong genetic component.

Presentation of idiopathic type 1 diabetes does not differ from that of autoimmune type 1 diabetes.

The American Diabetes Association has produced a staging system for type 1 diabetes based on clinical features and the presence of autoantibodies. The persistent presence of two or more autoantibodies is an almost certain predictor of clinical hyperglycemia and diabetes, and the rate of progression is dependent on the age at first detection of antibody, the number of antibodies, the antibody specificity, and the antibody titer.[1] Glucose and A1C (glycosylated hemoglobin) levels rise well before the clinical onset of diabetes, making early diagnosis possible. This staging can serve as a framework for future research and screening.
**Screening**

Routine screening is not recommended for type 1 diabetes due to a low population prevalence. Screening for antibodies that confer high risk is also not recommended because animal and human studies have not confirmed the utility of treatment (e.g., with nicotinamide; or oral, parenteral, or nasal insulin) to prevent or delay type 1 diabetes in high-risk individuals. Screening for related antibodies is only recommended in the context of a clinical research study.[32]
**Case history**

**Case history #1**

A 12-year-old white girl is brought to the emergency room by her parents due to 12 hours of rapidly worsening nausea, vomiting, abdominal pain, and lethargy. Over the last week she has felt excessively thirsty and has been urinating a lot. Physical examination reveals a lean, dehydrated girl with deep rapid respirations, tachycardia, and no response to verbal commands.

**Other presentations**

The rate of beta-cell destruction varies in type 1 diabetes. In some patients, there may be a slow destruction leading to gradual onset of symptoms that is clinically indistinguishable from type 2 diabetes. When the initial presentation of type 1 diabetes occurs in adulthood, some refer to it as latent autoimmune diabetes in adults (LADA). It is useful to distinguish LADA from type 2 diabetes, because patients with LADA usually require insulin therapy. Features that suggest the presence of LADA rather than type 2 diabetes include 2 or more of the following: age of onset less than 50 years, acute symptoms, BMI less than 25 kg/m^2, and personal or family history of autoimmune disease.[2]

**Step-by-step diagnostic approach**

**Clinical presentation**

Type 1 diabetes presents with polyuria, polydipsia, weight loss, generalized weakness, and blurred vision. Some patients present with diabetic ketoacidosis, the acute complication of type 1 diabetes. These patients have symptoms of dehydration and acidosis such as nausea, vomiting, abdominal pain, tachypnea, tachycardia, and lethargy. Rarely, a patient is diagnosed with type 1 diabetes during routine blood tests. The condition is diagnosed long before its chronic complications have developed.

**Diagnosis**

Diagnosis can be made on the basis of any of the following: 1) in a symptomatic patient, random plasma glucose of >200 mg/dL (>11 mmol/L); 2) fasting plasma glucose >126 mg/dL (>6.9 mmol/L); 3) plasma glucose ≥200 mg/dL ≥11 mmol/L) 2 hours after 75 g oral glucose; 4) A1C (glycosylated hemoglobin) ≥6.5% (≥48 mmol/mol). In an asymptomatic patient, results should be confirmed by repeating the test. In symptomatic patients, blood glucose rather than A1C is more useful for diagnosing acute onset of type 1 diabetes.[1] Diabetes is the overall diagnostic term applied to people satisfying these criteria, with type 1 and type 2 being further subclasses based on clinical and/or laboratory criteria.[30]

Elevated plasma or urine ketones in the presence of hyperglycemia suggests type 1 diabetes, but is occasionally seen at presentation in a patient with type 2 diabetes.

The diagnosis of type 1 diabetes is often obvious from the clinical presentation, but can be confirmed through additional testing. Low C-peptide levels and presence of one or more autoimmune markers are consistent with a diagnosis of type 1 diabetes. Autoimmune markers include autoantibodies to glutamic acid decarboxylase (GAD), insulin, islet cells, islet antigens (IA2 and IA2-beta), and the zinc transporter ZnT8. For example, when an obese teenager with a positive family history of type 2 diabetes is found to...
Type 1 diabetes mellitus

have high plasma glucose levels on routine blood tests, the diagnosis of type 1 versus type 2 diabetes may not be clear. If C-peptide levels are very low or undetectable relative to the plasma glucose and anti-GAD antibodies are positive in such a patient, a diagnosis of type 1 diabetes can be made.

[VIDEO: Venepuncture and phlebotomy: animated demonstration]

Risk factors

Strong

geographic region

- Human leukocyte antigen (HLA) risk profile for type 1 diabetes is widening over time, which may reflect increased environmental influence on susceptible genotypes.[27]
- Geographic variation ranges from 1/100,000 in regions of China to 38/100,000 in Finland.[6]

Weak

genetic predisposition

- In one study, concordance for type 1 diabetes was 27.3% in monozygotic twins and 3.8% in dizygotic twins.[25]
- HLA on chromosome 6 thought to contribute to half of the familial basis.[26]
- DR4-DQ8 and DR3-DQ2 present in 90% of children with type 1 diabetes; considered susceptibility genes.[26]
- DR15-DQ6 considered protective.[26]
- Insulin gene on chromosome 11 thought to be second most important susceptibility gene, contributing 10% of genetic susceptibility.[26]
- Several other loci associated with type 1 diabetes under study.[26]

infectious agents

- Strongest evidence to date is for congenital rubella and human enteroviruses.[10] [11] [12]

dietary factors

- Among dietary factors, infant supplementation with vitamin D may be protective.[13] Further research is required to determine whether cow’s milk, early introduction of cereals, or maternal vitamin D ingestion increase type 1 diabetes risk.[14] [15] [16] [17] [18] [19] There is no consensus about the effect of breastfeeding on risk for type 1 diabetes.[28] [29]

History & examination factors

Key diagnostic factors

polyuria (common)

- Getting up at night to urinate is typical.

polydipsia (common)

- Getting up at night to drink water is typical.
Type 1 diabetes mellitus

Diagnosis

Other diagnostic factors

young age (common)
- Usually presents in childhood or adolescence. Typical age 5 to 15 years.
- Average age varies in different studies. Incidence increasing in children <5 years old.[31]

weight loss (common)
- Weight loss occurs at onset.

blurred vision (common)
- Occurs with high or fluctuating blood sugar levels.

nausea and vomiting (common)
- Suggest diabetic ketoacidosis (DKA).

abdominal pain (common)
- Suggest DKA.

tachypnea (common)
- Suggest DKA.

lethargy (common)
- Suggest DKA.

coma (uncommon)
- Suggest DKA.

Diagnostic tests

1st test to order

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td>random plasma glucose</td>
<td>≥200 mg/dL (≥11 mmol/L)</td>
</tr>
<tr>
<td>- Confirms diagnosis in presence of symptoms of polyuria, polydipsia, and unexplained weight loss.[1]</td>
<td></td>
</tr>
<tr>
<td>fasting plasma glucose</td>
<td>≥126 mg/dL (≥6.9 mmol/L)</td>
</tr>
<tr>
<td>- Fasting is defined as no caloric intake for at least 8 hours.</td>
<td></td>
</tr>
<tr>
<td>2-hour plasma glucose</td>
<td>≥200 mg/dL (≥11 mmol/L)</td>
</tr>
<tr>
<td>- Plasma glucose is measured 2 hours after 75 g oral glucose load.</td>
<td></td>
</tr>
<tr>
<td>plasma or urine ketones</td>
<td>medium or high quantity</td>
</tr>
<tr>
<td>- In the presence of hyperglycemia suggest type 1 diabetes.</td>
<td></td>
</tr>
<tr>
<td>A1C (glycosylated hemoglobin)</td>
<td>≥6.5% (≥48 mmol/L)</td>
</tr>
<tr>
<td>- Reflects degree of hyperglycemia over the preceding 3 months.</td>
<td></td>
</tr>
</tbody>
</table>
Other tests to consider

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td>fasting C-peptide</td>
<td>low or undetectable</td>
</tr>
<tr>
<td>• C-peptide is a byproduct formed when proinsulin is processed to insulin. Therefore, its levels reflect insulin production. Half life of C-peptide is 3 to 4 times longer than that of insulin. • Low or undetectable C-peptide level indicates absence of insulin secretion from pancreatic beta cells.</td>
<td></td>
</tr>
<tr>
<td>autoimmune markers</td>
<td>positive</td>
</tr>
<tr>
<td>• These include autoantibodies to glutamic acid decarboxylase, insulin, islet cells, islet antigens (IA2 and IA2-beta), and the zinc transporter ZnT8. • Presence indicates autoimmune beta-cell destruction.</td>
<td></td>
</tr>
</tbody>
</table>

Differential diagnosis

<table>
<thead>
<tr>
<th>Condition</th>
<th>Differentiating signs / symptoms</th>
<th>Differentiating tests</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maturity onset diabetes of the young</td>
<td>• Strong family history. • Slow onset. • Absence of ketoacidosis. • Response to sulfonylurea drugs.</td>
<td>• C-peptide present. • Autoantibodies absent.</td>
</tr>
<tr>
<td>Type 2 diabetes</td>
<td>• Typically, signs of insulin resistance (such as acanthosis nigricans) should be sought and in their absence clinical suspicion of type 1 diabetes is greater. • Signs of more marked insulin deficiency (for example, glycemic lability as well as susceptibility to ketosis) raise suspicion of type 1 diabetes. • Older age and slow onset, obesity, a strong family history, absence of ketoacidosis, and initial response to oral antihyperglycemic drugs are typical of type 2 diabetes.</td>
<td>• C-peptide present. • Autoantibodies absent. • Testing for C-peptide and autoantibodies usually not required.</td>
</tr>
</tbody>
</table>

Diagnostic criteria

American Diabetes Association: criteria for diagnosis of diabetes[1]

In the absence of unequivocal hyperglycemia, any of the tests should be confirmed on a subsequent day by repeat testing. Screening tests are generally most applicable to type 2 diabetes.
Type 1 diabetes mellitus

Diagnosis

- Random plasma glucose level $\geq 200$ mg/dL ($\geq 11$ mmol/L) in the presence of symptoms of hyperglycemia; OR
- Fasting plasma glucose $\geq 126$ mg/dL ($\geq 6.9$ mmol/L); OR
- Plasma glucose level $\geq 200$ mg/dL ($\geq 11$ mmol/L) 2 hours after a 75 g oral glucose load; OR
- A1C (glycosylated hemoglobin) $\geq 6.5\%$ ($\geq 48$ mmol/L).
Step-by-step treatment approach

In the short term, insulin is life-saving because it prevents diabetic ketoacidosis, a potentially life-threatening condition. The long-term goal of insulin treatment is the prevention of chronic complications by maintaining blood glucose levels as close to normal as possible. Generally, A1C (glycosylated hemoglobin) goals determine the aggressiveness of therapy, which is in turn individualized. The American Diabetes Association (ADA) recommends a target A1C goal of <7.5% for patients <18 years with type 1 diabetes and <7% for adult patients. Less stringent goals may be appropriate for very young children, older adults, people with a history of severe hypoglycemia, and those with limited life expectancies, advanced microvascular or macrovascular complications, or comorbid conditions.

Good glycemic control in type 1 diabetes requires attention to diet, exercise, and insulin therapy. All 3 components should be coordinated for ideal control. Self-monitoring of blood glucose (SMBG) is a core component of good glycemic control. Patients on multiple injections daily should consider SMBG before meals, occasionally after meals and at bedtime, and before exercising, to assess presence and adequate treatment of hypoglycemia, and before any task during which hypoglycemia could have particularly dangerous consequences. Some patients will need to check their blood glucose 6 to 10 times daily. As continuous glucose monitoring (CGM) technology continues to improve, the indications for its use are likely to expand. Currently, it is considered to improve glycemic control in patients >18 years old. However, the most cost-effective or appropriate use of CGM is when targeted at people with type 1 diabetes who have hypoglycemic unawareness, frequent hypoglycemia, or continued poor control during intensified insulin therapy, and those who are willing to use CGM frequently. The limiting factor for tight glycemic control in type 1 diabetes is hypoglycemia. Well-controlled blood pressure and lipids, and avoidance of smoking, are essential components of cardiovascular risk reduction.

The ADA recommends including technology-based methods, along with individual and group settings, for the delivery of effective diabetes self-management education and support. This approach can be used for adults as well as children and adolescents.

Diet and exercise

There is no standardized dietary advice that is suitable for all individuals with diabetes. Individualized nutrition advice should be based on personal and cultural preferences, health literacy and numeracy, access to healthful food choices, and willingness and ability to make behavioral changes. It should also address barriers to change. All patients with diabetes should receive individualized medical nutrition therapy, preferably provided by a registered dietitian who is experienced in providing this type of therapy to diabetes patients. Carbohydrate counting or consistent carbohydrate intake with respect to time and amount may improve glycemic control. Rapid-acting insulins may make timing of meals less crucial than in the past, but regular meals are still important.

The ADA recommends 150 minutes/week of moderate-intensity aerobic exercise (to 50-70% of max heart rate) spread over at least 3 days per week with no more than 2 consecutive days without exercise. Pre-exercise carbohydrate intake and insulin doses can be effectively managed to avoid hypoglycemia during exercise and sports. Clinical judgment should be used in determining whether to screen asymptomatic individuals for coronary artery disease prior to recommending an exercise program. For example, an exercise stress test prior to starting a program is advised if more than brisk walking is planned for sedentary people whose 10-year risk of a coronary event is 10% or greater by the Framingham Risk Score.
Type 1 diabetes mellitus

The following should be assessed prior to starting an exercise program: age; physical condition; blood pressure; and presence or absence of autonomic neuropathy or peripheral neuropathy, preproliferative or proliferative retinopathy, or macular edema. Vigorous exercise may be contraindicated with proliferative or severe preproliferative diabetic retinopathy. Nonweight-bearing exercise may be advisable in patients with severe peripheral neuropathy. Insulin should be adjusted to avoid hypoglycemia, which can occur up to 24 hours after exercise. This may require reducing insulin dosage on days of planned exercise. A carbohydrate snack should be given at the start of exercise if the blood sugar is <100 mg/dL (<5.6 mmol/L).

Prolonged sitting should be interrupted every 30 minutes with short bouts of physical activity.[1]

One consensus statement provides guidance on how patients with type 1 diabetes can safely exercise and manage their glucose levels.[40]

**Initiating insulin**

Intensive therapy with insulin should be started as soon as possible after diagnosis. Unlike older regimens that used nonphysiologic insulin dosing, intensive therapy aims to mimic physiologic insulin release by combining basal insulin with bolus dosing at mealtime. Both continuous infusion with an insulin pump and a regimen of multiple daily injections (MDI) can provide intensive therapy.[41] The choice between pump and MDI is based on patient interest and self-management skills as well as physician preference, as outcomes are generally similar.[42] The insulin pump uses regular or rapid-acting insulin, and provides a basal rate of insulin and delivers mealtime bolus dosing. However, the patient or parent must still measure blood glucose frequently in order to adjust the pump to deliver the appropriate amount of insulin. Insulin pumps may reduce hypoglycemia, especially when combined with continuous glucose monitoring systems (CGMS) and threshold suspend features,[43] and improve A1C, while providing greater flexibility.[44] [45] [46] Use of a pump requires a motivated patient with strong family support (for children), and access to practitioners trained in pump therapy.[47]

Using a combination of long- (insulins glargine or detemir) or intermediate- (NPH) acting insulin for basal dosing, and rapid- (insulins lispro, aspart, or glulisine) or short- (regular) acting insulin for bolus dosing, MDI regimens can be designed based on physician and patient preference and modified based on fingerstick data. There is no consensus as to whether insulin analogs are superior to conventional insulins for short-term glycemic control or reductions in complications.[48]

In the past, many patients were managed with twice-daily injections of a mixture of rapid-acting and intermediate-acting insulin. This regimen may be used if patients are unable to comply with MDI, but it is no longer a first-line recommendation for management because of its lack of flexibility.

**Designing a regimen**

An initial total daily dose of insulin in adults can be 0.2 to 0.4 units/kg/day. In children an initial daily dose will be 0.5 to 1 units/kg/day, and during puberty the requirements may increase to as much as 1.5 units/kg/day. Often, when first started on insulin, patients with type 1 diabetes will experience a honeymoon period, during which they may require only 10 or 15 units/day. One half of the total dose is given as basal insulin and one half as bolus dosing. The bolus dosing is divided and given before meals. Patients need to self-monitor their blood glucose levels. The insulin doses can be adjusted every 2 to 3 days to maintain target blood glucose. To achieve an A1C <7% (<53 mmol/mol), the premeal blood glucose goal is 80 to 130 mg/dL (4.4 to 7.2 mmol/L) and the postmeal blood glucose goal (1-2 hours after starting the meal) is less than 180 mg/dL (10.0 mmol/L).[1]
The simplest approach to covering mealtime insulin requirements is to suggest a range of doses, such as 4 units for a small meal, 6 units for a medium-sized meal, and 8 units for a larger meal. For greater flexibility of carbohydrate content of meals, premeal insulin can be calculated based on the estimated amount of carbohydrate in the meal and the patient’s individual insulin-to-carbohydrate ratio. A simple beginning approach is to use one unit of mealtime insulin for every 15 g of carbohydrate in the meal. Patients can use the carbohydrate content per serving listed on food packaging to assess the number of grams in their anticipated meal, but carbohydrate counting is best learned with the help of a nutritionist. Using a food diary and 2-hour postprandial blood glucose measurements, the insulin-to-carbohydrate ratio can be adjusted.

A correction dose may be added to the bolus insulin based on the premeal blood glucose level. Correction dosing may be calculated as follows when the patient’s total daily dose of insulin (TDD) and food intake is stable: \(1800/TDD = \text{the predicted point drop in blood glucose per unit of rapid acting insulin. For example, if the TDD is 40 units of insulin,} \frac{1800}{40} = 45 \text{ point drop per unit of insulin.}\)

Example of correction dosing based on premeal glucose and above calculation:

- 45 to 90 mg/dL (2.2 to 4.9 mmol/L): subtract 1 unit from mealtime insulin
- 91 to 135 mg/dL (5 to 7.4 mmol/L): add 0 units of correction insulin
- 136 to 180 mg/dL (7.5 to 9.9 mmol/L): add 1 unit of correction insulin
- 181 to 225 mg/dL (9.9 to 12.4 mmol/L): add 2 units of correction insulin
- 226 to 270 mg/dL (12.4 to 14.5 mmol/L): add 3 units of correction insulin
- 271 to 315 mg/dL (14.5 to 17.3 mmol/L): add 4 units of correction insulin
- 316 to 360 mg/dL (17.4 to 19.8 mmol/L): add 5 units of correction insulin
- 361 to 405 mg/dL (19.8 to 22.3 mmol/L): add 6 units of correction insulin
- >405 mg/dL (>22.3 mmol/L): add 7 units of correction insulin; call healthcare provider.

The number used to calculate the correction dose may be as low as 1500 or as high as 2200. There are no specific guidelines to determine this number. In general, a lower number should be used for obese, insulin-resistant patients, and a higher number should be used for lean, insulin-sensitive patients.

This correction dose can be added to the patient’s mealtime insulin requirement (whether based on general meal size or carbohydrate counting) and given as the total bolus dose.

Pump therapy utilizes a similar concept as basal and bolus dosing and does not require multiple injections of insulin. However, patients still need to monitor their blood glucose from 4 to 7 times daily. There is some evidence that insulin pump therapy may be associated with improved glycemic control and lower risk of hypoglycemia, including in children, adolescents, and young adults. Because of the monitoring and dose adjustment required, patients selected for pump therapy must be skilled in diabetes self-management and able to manage and troubleshoot the various pump components.

The insulin pump uses a subcutaneous insulin injection port. The port is changed every 3 days and may reduce anxiety and help achieve better glycemic control in selected patients.

CGMS measure subcutaneous interstitial fluid glucose every 5 minutes. CGMS may be indicated in selected patients with widely fluctuating glucose levels or hypoglycemia unawareness. A 3-day glucose monitoring system using a CGMS may help the physician adjust insulin doses. Real time CGMS, worn by a patient on a regular basis, may help improve glycemic control. The glucose sensors used in CGMS are not reliable at lower ranges of glucose, and thus do not eliminate the need for fingersticks. Development of these systems is ongoing.
Type 1 diabetes mellitus

CGMS are also less accurate than traditional capillary blood glucose monitoring methods. However, they provide information on glucose trends, provide alarms to alert patients to impending hypo- or hyperglycemia, and reduce episodes of hypoglycemia.[43] [59] Insulin pumps with glucose sensors integrated into the same unit are called sensor-augmented insulin pumps. Functionality between sensor and pump has been integrated in one available device; the insulin delivery can be determined automatically based on sensed glucose levels. These integrated devices use a computerized control algorithm to create a closed-loop insulin delivery system, which functions as an artificial pancreas. In clinical trials, such systems have been shown to reduce the risk of nocturnal hypoglycemia and to improve glucose control.[60] Use of sensors and sensor-augmented pumps is increasing and is increasingly reimbursed by insurance providers.

Hypoglycemia is the most common and potentially most serious side effect of insulin therapy, as it can lead to decreased quality of life, confusion, seizures, and coma. Episodes of hypoglycemia should be sought at each visit, and efforts made to determine contributing factors, and the ability of the patient to recognize and treat it appropriately. Dinnertime NPH is a frequent cause of symptomatic and asymptomatic nocturnal hypoglycemia, and can be taken at bedtime so that the peak effect is closer to the early morning increase in cortisol.

Goal not met

If glycemic control is not adequate as measured by the A1C or by episodes of hypoglycemia, the patient's nutrition, exercise, and insulin regimen must be reexamined. Children and adolescents may have erratic eating patterns or snack frequently. Consultation with a nutritionist is an invaluable part of the treatment approach, as patients can learn how to count carbohydrates and adjust their premeal insulin to allow for flexibility in meal content and activity. Consistent hyperglycemia may require an increase in basal insulin. Preprandial and postprandial hyperglycemia may be due to inadequate insulin coverage for the most recent meal, and may be addressed by considering carbohydrate content of meals, the patient’s assessment of their carbohydrate intake, and subsequent premeal insulin dosing. If a patient is getting regular insulin, replacing it with rapid-acting insulin may reduce postprandial glucose excursions.

Other conditions contributing to unstable diabetes and that co-exist most commonly with diabetes include celiac disease, thyroid disease, Addison disease, and psychosocial distress. Celiac disease, thyroid disease, and psychosocial distress should be screened for at diagnosis and on a regular basis, while increased clinical suspicion should prompt screening for Addison disease and pernicious anemia.

Episodes of hypoglycemia occur with different frequency among patients. Patients should check a 3 a.m. blood glucose if there is concern about risk of nocturnal hypoglycemia. Nocturnal hypoglycemia may result in rebound hyperglycemia in the morning. The dose of basal insulin should be decreased to prevent nocturnal hypoglycemia. A bedtime snack is not an effective way of decreasing the risk of nocturnal hypoglycemia.[61] Alcohol may cause acute hypoglycemia, but both alcohol and exercise can cause delayed hypoglycemia (up to 24 hours).

Noninsulin treatments

Pramlintide is indicated as adjunctive treatment in patients with postprandial hyperglycemia that cannot be controlled with premeal insulin alone. For example, it may be useful in a patient with high postprandial glucose, but who develops late hypoglycemia when premeal insulin is increased.

The therapy of people with type 1 diabetes also involves regular eye examinations, foot care, treatment of dyslipidemia, and blood pressure control.
Adults with type 1 diabetes are at 3 times the risk of clinical depression compared with those without type 1 diabetes.[62] The prevalence of depression in diabetes is higher in women (28%) compared with men (18%).[63] The risk may also be higher in adolescents, at diagnosis, or when there is a change in disease status.[64] Psychosocial screening and support can help to ameliorate distress and improve the individual’s and family’s capacity for self-care.

**Pregnancy**

Infants of women with diabetes are at high risk of major congenital malformations and miscarriage. Preconception diabetes care reduces this risk.[65] Preconception counseling should therefore be incorporated in the routine diabetes clinic visit for all women of childbearing potential. Women with type 1 diabetes should use an effective method of contraception until they plan pregnancy. The ADA recommends that A1C should be <6.5% (<48 mmol/mol) before conception if this can be achieved without hypoglycemia.[1] Women should also be evaluated before pregnancy for retinopathy, nephropathy, neuropathy, and possible cardiovascular disease, which may worsen during or complicate pregnancy.

In addition to the complications noted above, infants of mothers with hyperglycemic diabetes are at risk of macrosomia and neonatal distress. Preeclampsia is also more common in diabetic pregnancies. Euglycemia or near-euglycemia reduces the risk of complications. During pregnancy women should be cared for by a multidisciplinary team including a nutritionist, a nurse educator, an endocrinologist, and an obstetrician. All pregnant women require a dilated eye exam soon before or early in pregnancy. Women with diabetes have an increased risk of having infants with neural tube defects, compared with the general population.[66] Statins, angiotensin-converting enzyme inhibitors, and angiotensin-II receptor blockers should be discontinued preconception. Intensive insulin treatment with MDI or insulin pump should be started. Commonly used insulins during pregnancy include NPH, detemir, regular, lispro, and aspart.[67] Use of CGM during pregnancy may help in improving glycemic control and neonatal outcomes.[68]

There are no large randomized trials supporting the safety of insulin glargine in pregnant patients with diabetes. However, insulin glargine has been safely used in many patients during pregnancy, although it is US Food and Drug Administration pregnancy class C. It can be considered second-line to NPH or insulin detemir for basal insulin dosing during pregnancy because there are fewer long-term safety monitoring data. There are few data comparing outcomes for continuous subcutaneous insulin infusion versus multiple daily injections of insulin for pregnant women with diabetes.[69] ADA guidelines recommend the following blood glucose targets in pregnant women with pre-existing type 1 diabetes (the same as for gestational diabetes): <95 mg/dL (<5.3 mmol/L) fasting, and either ≤140 mg/dL (≤7.8 mmol/L) 1 hour postprandially or ≤120 mg/dL (≤6.7 mmol/L) 2 hours postprandially, with A1C goal individualized between <6% and <6.5% (<42 and <48 mmol/mol) or up to <7% (<53 mmol/mol) as necessary to prevent hypoglycemia.[1]

The ADA recommends that all pregnant women with preexisting type 1 diabetes should consider daily low-dose aspirin starting at the end of the first trimester in order to reduce the risk of preeclampsia.[1]

**Comorbidities**

Guidelines emphasize the importance of assessing and managing comorbidities. An expanded list of diabetes comorbidities now includes autoimmune diseases, HIV, anxiety disorders, depression, disordered eating behavior, and serious mental illness.[1]
Treatment details overview

Consult your local pharmaceutical database for comprehensive drug information including contraindications, drug interactions, and alternative dosing. (see Disclaimer)

<table>
<thead>
<tr>
<th>Ongoing</th>
<th>(summary)</th>
<th>1st</th>
<th>basal-bolus insulin</th>
</tr>
</thead>
<tbody>
<tr>
<td>nonpregnant</td>
<td></td>
<td>adjunct</td>
<td>premeal insulin correction dose</td>
</tr>
<tr>
<td></td>
<td></td>
<td>adjunct</td>
<td>amylin analog</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2nd</td>
<td>fixed-dose insulin</td>
</tr>
<tr>
<td>pregnant</td>
<td></td>
<td>1st</td>
<td>basal-bolus insulin</td>
</tr>
<tr>
<td></td>
<td></td>
<td>plus</td>
<td>low-dose aspirin</td>
</tr>
</tbody>
</table>
# Treatment options

<table>
<thead>
<tr>
<th>Ongoing nonpregnant</th>
<th>1st</th>
<th>basal-bolus insulin</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td><strong>Primary options</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td>» insulin glargine: injected subcutaneously once daily</td>
</tr>
<tr>
<td></td>
<td></td>
<td>» insulin NPH: injected subcutaneously twice daily</td>
</tr>
<tr>
<td></td>
<td></td>
<td>» insulin detemir: injected subcutaneously twice daily</td>
</tr>
<tr>
<td></td>
<td></td>
<td>» insulin degludec: injected subcutaneously once daily</td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>--AND--</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td>» insulin regular: injected subcutaneously two to three times daily</td>
</tr>
<tr>
<td></td>
<td></td>
<td>» insulin lispro: injected subcutaneously premeal</td>
</tr>
<tr>
<td></td>
<td></td>
<td>» insulin aspart: injected subcutaneously premeal</td>
</tr>
<tr>
<td></td>
<td></td>
<td>» insulin glulisine: injected subcutaneously premeal</td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>OR</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td>» pump: uses regular insulin or insulins lispro, aspart or glulisine</td>
</tr>
<tr>
<td></td>
<td></td>
<td>An initial total daily dose of insulin in adults can be 0.2 to 0.4 units/kg/day. In children an initial daily dose will be 0.5 to 1 units/kg/day, and during puberty the requirements may increase to as much as 1.5 units/kg/day. Often, when first started on insulin, patients with type 1 diabetes will experience a honeymoon period, during which they may require only 10 or 15 units/day. One half of the total dose is given as basal insulin and one half as bolus dosing. The bolus dosing is divided and given before meals. Patients need to self-monitor their blood glucose levels. The insulin doses can be adjusted every 2 to 3 days to maintain premeal and postmeal targets.[1]</td>
</tr>
<tr>
<td></td>
<td></td>
<td>The simplest approach to covering mealtime insulin requirements is to suggest a range of doses, such as 4 units for a small meal, 6 units</td>
</tr>
</tbody>
</table>
Type 1 diabetes mellitus

Treatment

Ongoing

for a medium-sized meal, and 8 units for a larger meal. For greater flexibility of carbohydrate content of meals, premeal insulin can be calculated based on the estimated amount of carbohydrate in the meal and the patient’s individual insulin-to-carbohydrate ratio. A simple beginning approach is to use one unit of mealtime insulin for every 15 g of carbohydrate in the meal. Patients can use the carbohydrate content per serving listed on food packaging to assess the number of grams in their anticipated meal, but carbohydrate counting is best learned with the help of a nutritionist. Using a food diary and 2-hour postprandial blood glucose measurements, the insulin-to-carbohydrate ratio can be adjusted.

» Reasonable to begin therapy with 2 to 4 insulin injections daily to cover basal insulin requirements and to cover mealtime insulin needs. Intermediate- or long-acting insulins to cover basal, and short- or rapid-acting to cover mealtime needs, should be used.

» Regular and NPH insulins are less expensive than the insulin analogs. Regular insulin is given about 30 minutes prior to the meal, while rapid-acting insulins (lispro, aspart, or glulisine) can be injected 15 minutes before to shortly after a meal. In children with erratic eating habits, rapid-acting insulins can be given just after the meal. NPH and insulin detemir are injected twice daily while insulin glargine can be injected once daily. The regimen should be individualized to obtain the best possible glycemic control.

» A correction dose may be incorporated into the insulin doses based on premeal glucose levels.

» Patients with interest and good self-management skills may prefer to use an insulin pump.

adjunct  premeal insulin correction dose

» A correction dose may be added to the bolus insulin based on the premeal blood glucose level. Correction dosing may be calculated as follows when the patient’s total daily dose of insulin (TDD) and food intake is stable: 1800/TDD = the predicted point drop in blood glucose per unit of rapid acting insulin. For example, if the TDD is 40 units of insulin, 1800/40 = 45 point drop per unit of insulin.

» Example of correction dosing based on premeal glucose and above calculation:
## Ongoing

- **45 to 90 mg/dL (2.2 to 4.9 mmol/L):** subtract 1 unit from mealtime insulin  
- **91 to 135 mg/dL (5 to 7.4 mmol/L):** add 0 units of correction insulin  
- **136 to 180 mg/dL (7.5 to 9.9 mmol/L):** add 1 unit of correction insulin  
- **181 to 225 mg/dL (9.9 to 12.4 mmol/L):** add 2 units of correction insulin  
- **226 to 270 mg/dL (12.4 to 14.5 mmol/L):** add 3 units of correction insulin  
- **271 to 315 mg/dL (14.5 to 17.3 mmol/L):** add 4 units of correction insulin  
- **316 to 360 mg/dL (17.4 to 22.3 mmol/L):** add 5 units of correction insulin  
- **361 to 405 mg/dL (19.8 to 22.3 mmol/L):** add 6 units of correction insulin  
- **>405 mg/dL (>22.3 mmol/L):** add 7 units of correction insulin; call healthcare provider.

The number used to calculate the correction dose may be as low as 1500 or as high as 2200. There are no specific guidelines to determine this number. In general, a lower number should be used for obese, insulin-resistant patients, and a higher number should be used for lean, insulin-sensitive patients.

This correction dose can be added to the patient's mealtime insulin requirement (whether based on general meal size or carbohydrate counting) and given as the total bolus dose.

### adjunct amylina analog

#### Primary options

- **pramlintide:** 15-60 micrograms subcutaneously before each meal

  Synthetic analog of human amylin, a protein that is co-secreted with insulin by pancreatic beta cells. It reduces postprandial glucose increases by prolonging gastric emptying time, reducing postprandial glucagon secretion, and reducing food intake through centrally mediated appetite suppression.[70]

  May be given as an injection before each meal to get more stable glycemic control. However, insulin treatment must continue in addition to pramlintide.
### Ongoing

» At initiation the current premeal insulin dose should be reduced by about 50% to avoid hypoglycemia, and then titrated up.

» Indicated as adjunctive treatment in patients with postprandial hyperglycemia that cannot be controlled with premeal insulin alone. For example, it may be useful in a patient with high postprandial glucose, but who develops late hypoglycemia when premeal insulin is increased.

» Should not be used in a patient with gastroparesis. The most common side effect is nausea, occurring in 28% to 48% of patients.[70]

### 2nd fixed-dose insulin

**Primary options**

- **insulin NPH/insulin regular:** (50/50, 70/30) injected subcutaneously twice daily

OR

- **insulin aspart protamine/insulin aspart:** (70/30) injected subcutaneously twice daily

OR

- **insulin lispro protamine/insulin lispro:** (50/50, 75/25) injected subcutaneously twice daily

OR

- **insulin degludec/insulin aspart:** (70/30) injected subcutaneously once or twice daily

» Fixed-dose insulin is used when patients are already doing well on a fixed-dose regimen; or cannot manage 3 to 4 insulin injections daily; or have trouble mixing insulin.

### pregnant

### 1st basal-bolus insulin

**Primary options**

- **insulin NPH:** injected subcutaneously twice daily
  - or -
  - **insulin detemir:** injected subcutaneously twice daily

--AND--

- **insulin regular:** injected subcutaneously two to three times daily
  - or -
## Treatment

### Ongoing

- **insulin lispro**: injected subcutaneously premeal  
  -or-  
- **insulin aspart**: injected subcutaneously premeal

### Secondary options

- **insulin glargine**: injected subcutaneously once daily
  -AND--  
- **insulin regular**: injected subcutaneously two to three times daily  
  -or-  
- **insulin lispro**: injected subcutaneously premeal  
  -or-  
- **insulin aspart**: injected subcutaneously premeal

### OR

- **pump**: uses regular insulin or insulins lispro or aspart

Blood sugar goals, if able to be achieved without significant hypoglycemia, are fasting ≤90 mg/dL (≤5 mmol/L), 1-hour postprandial ≤130-140 mg/dL (≤7.2-7.8 mmol/L), and 2-hour postprandial ≤120 mg/dL (≤6.7 mmol/L). If these targets result in hypoglycemia, less stringent targets are appropriate.[1]

An A1C (glycosylated hemoglobin) target of 6% to 6.5% (42 to 48 mmol/mol) is recommended, but in the second and third trimester A1C <6% (<42 mmol/mol) may provide additional benefit, if it can be achieved without hypoglycemia.[1]

A1C in pregnancy can be monitored monthly.[1]

Patients should monitor their blood glucose from 4 to 7 times a day and the pattern should be examined every few weeks early in pregnancy so that nutrition content and timing, exercise patterns, and the insulin doses can be modified to achieve optimal control.

Insulin requirements generally increase early in pregnancy, then decrease from about 8 to 16 weeks before rising throughout the rest of the pregnancy.

Commonly used insulins during pregnancy include NPH, detemir, regular, lispro, and
Use of CGM during pregnancy may help in improving glycemic control and neonatal outcomes.[68] There are no large randomized trials supporting the safety of insulin glargine in pregnant patients with diabetes. However, insulin glargine has been safely used in many patients during pregnancy, although it is US Food and Drug Administration pregnancy class C. It can be considered second-line to NPH or insulin detemir for basal insulin dosing during pregnancy because there are fewer long-term safety monitoring data. There are few data comparing outcomes for continuous subcutaneous insulin infusion versus multiple daily injections of insulin for pregnant women with diabetes.[69]

**Primary options**

» **aspirin**: 60-125 mg orally once daily, usual dose 81 mg/day

» The American Diabetes Association recommends that all pregnant women with preexisting type 1 diabetes should consider daily low-dose aspirin starting at the end of the first trimester in order to reduce the risk of preeclampsia.[1]
Emerging

Implantable continuous insulin infusion pumps

A surgically implanted insulin pump is available in the European Union and is under investigation in the US. Insulin is delivered into the peritoneal cavity or intravascularly. In a trial there were fewer episodes of hypoglycemia with the implantable pump than with multiple subcutaneous insulin injections.[71]

Islet cell transplantation

Islet cells prepared from a donor pancreas are injected into the portal vein. The cells seed in the liver and produce insulin. Patients who undergo this procedure require immunosuppressive therapy afterwards. There is some initial success with this procedure but the long-term results remain disappointing. Even in the best centers, less than 50% patients are free of insulin requirement at 1 year and only 10% at 5 years.[72] [73] The American Diabetes Association recommends that this procedure be performed only within the context of a controlled research study at this time.

Inhaled insulin

In June 2014, the US Food and Drug Administration approved a rapid-acting inhaled insulin. It can be administered before meals and should be used in combination with long-acting insulin. It can cause bronchospasm in patients with asthma and chronic obstructive pulmonary disease, and should not be used if these conditions are present. The most common side effects in a 24-week safety and efficacy trial were hypoglycemia, cough, and throat infection. Long-term safety data are lacking.[74] Moreover, it is available only in fixed doses of 4 or 8 units. Therefore, dose adjustments can be made only in multiples of 4 which may present difficulty in fine-tuning the dose in patients with type 1 diabetes. More experience is needed before inhaled insulin is routinely prescribed in type 1 diabetes.

Immunotherapy

Type 1 diabetes is an autoimmune disease modulated by cytotoxic T cells. Several agents have been studied for treatment of new-onset disease. Nonantigen-specific systemic immunotherapies, including T-cell suppressors (cyclosporine), antiproliferative agents (methotrexate, azathioprine), and antithymocyte globulin have shown a strong tendency to adverse effects. Although cyclosporine use did reduce insulin requirements in the short term, it was associated with nephrotoxicity, and the effect on beta cells waned with treatment cessation. Antigen-specific vaccination with recombinant glutamic acid decarboxylase was shown to increase stimulated C-peptide in patients treated within 3 months of diagnosis.[75] Monoclonal antibodies to CD3 and CD20 have also shown some promise.[76] [77] Other trials are under way to investigate treatment of type 1 diabetes with dendritic cells, mesenchymal stem cells, cord blood transfusion, and immunomodulators currently approved for use in other diseases, such as granulocyte colony stimulating factor or tumor necrosis factor-alpha inhibitors.[78]

Islet cell regeneration

Studies done in mouse models show that from the onset of insulinitis, there is a mass of beta cells within an inflammatory milieu that may be recoverable and serve as a future source of functioning beta cells.[79] Several trials are under way to investigate mono- and combination therapies to arrest inflammation and possibly allow beta-cell regeneration.

Insulin sensitizers

A systematic review suggested that use of metformin in type 1 diabetes reduced insulin requirements but not A1C (glycosylated hemoglobin) levels after 1 year of follow-up.[80] Further research is indicated to better delineate the potential indications and benefits of this treatment in type 1 diabetes.[81] [82]

Glucagon-like peptide-1 (GLP-1) agonists
GLP-1 is a gut peptide that increases insulin secretion and decreases glucagon secretion in a glucose-dependent manner. In patients with type 2 diabetes, GLP-1 receptor agonists increase levels of GLP-1 and lead to more glucose-dependent insulin secretion, less glucagon secretion, delayed gastric emptying, and increased satiety. The specific advantage of GLP-1 agonists is weight loss, which may be desirable in some patients with type 1 diabetes. The GLP-agonist liraglutide added to insulin improved glucose control in clinical trials with type 1 diabetes, but also increased the risk of both hypoglycemia and hyperglycemia with ketosis. Therefore, GLP-1 agonists should not routinely be used in type 1 diabetes.

**Sodium-glucose co-transporter 2 inhibitors**

Sodium-glucose co-transporter 2 (SGLT2) inhibitors reduce glucose in an insulin-independent manner, by inhibiting SGLT2 in the proximal renal tubule and blocking glucose reabsorption. They are associated with modest weight loss and blood pressure reduction. SGLT2 inhibitors are approved for use in individuals with type 2 diabetes. Several reports have highlighted the risk of euglycemic diabetic ketoacidosis in both type 2 and type 1 diabetes. While studies are under way to assess safety and efficacy in type 1 diabetes, the class of medications is not currently recommended for those with type 1 diabetes.
**Monitoring**

- **A1C (glycosylated hemoglobin)** should be checked twice yearly in patients who are meeting treatment goal of <7.5% for patients <18 years with type 1 diabetes and <7% for adult patients. It is recommended to check A1C every 3 months in patients whose therapy is being modified or who are not meeting the goal. In very elderly or very young patients and in those with a history of severe hypoglycemia or limited lifespan, the A1C goal can be less stringent.[1][35]

- **Check blood pressure at each visit and treat to a goal of less than 140/90 mmHg.[1]** In older adults, treating to <130/70 mmHg is not recommended.

- For patients who are not on statins, it is recommended to check a screening lipid profile (total cholesterol, LDL cholesterol, HDL cholesterol, and triglycerides) in adults with diabetes at the time of first diagnosis, at initial medical evaluation, and then every 5 years thereafter. Lifestyle modification should be recommended to all patients with diabetes to improve lipid profile. For patients with atherosclerotic cardiovascular disease, a high-intensity statin should be added to lifestyle therapy. For patients aged 40 to 75 years without additional atherosclerotic cardiovascular disease risk factors, the American Diabetes Association (ADA) recommends adding a moderate-intensity statin. For patients aged 40 to 75 years with additional atherosclerotic cardiovascular disease risk factors, the ADA recommends adding a high-intensity statin. Once a patient is taking a statin, LDL cholesterol testing may be considered on an individual basis (e.g., to monitor adherence and efficacy). If a patient is adherent to statin therapy but not responding, clinical judgment is recommended to determine the need for and timing of lipid panels.[1]

- In the US, initial screening for retinopathy by an ophthalmologist is recommended within 5 years of initial diagnosis of diabetes, and every 2 years after that if no evidence of retinopathy. In the presence of abnormal findings, more frequent follow-up may be indicated (e.g., annually).[1] Recommendations differ in other countries; for example, in the UK, screening for retinopathy is offered at the time of diagnosis and annually to all patients over the age of 12.[103][104] Local guidance should be consulted.

- Yearly screening for increased urinary albumin excretion and serum creatinine to estimate glomerular filtration rate should be done in all patients who have had type 1 diabetes for 5 years or more.[1]

- Screen yearly for distal symmetric polyneuropathy using pinprick sensation, temperature and vibration perception (with 128 Hz tuning fork), and 10 g monofilament pressure sensation at the distal plantar aspect of both great toes and ankle reflexes.

- Symptoms of autonomic neuropathy can be assessed through history (exercise intolerance, constipation, diarrhea, gastroparesis, bladder or sexual dysfunction, hypoglycemic autonomic failure), and physical exam (resting tachycardia, orthostatic hypotension).[1]

- Yearly dental exams are indicated in patients with and without teeth, to control periodontal disease, which both contributes to and exacerbates hyperglycemia.

- Vaccines should be provided in accordance with age-specific guidelines for the general population, including those for influenza and pneumococcal pneumonia. Hepatitis B vaccine should be provided for unvaccinated adults with diabetes aged 19 to 59 years, and should be considered for unvaccinated adults with diabetes aged ≥60 years.[1]

- Patients with autoimmune diabetes are more likely to have thyroid disease, celiac disease, and depression.[35] Physicians should have a low threshold for screening for these conditions.

**Patient instructions**

- The physician should advise the patient and/or caregivers that it is important to eat a healthy diet and regular meals. Referral to a nutritionist or a dietitian can be helpful in planning a diet.

- Patients should be advised on exercise. They should build up exercise slowly. If the weather is very hot or cold, they could walk at an indoor track or mall. Patients may need to take less insulin or eat a snack before exercise. They should also be advised to check their blood glucose before and after
exercising. Any patients with peripheral neuropathy would be wise to perform low-impact exercises such as swimming, bicycling, or arm exercises.

• The physician should help the patient plan how often to check blood glucose. The most likely times would be before each meal and at bedtime. Patients may also check 2 hours after meals and when exercising.

• Patients should usually have a A1C performed every 3 months.

• Patients should be advised that hypoglycemia may occur if they skip a meal, take too much insulin, exercise, or become ill. Alcohol and exercise can cause delayed hypoglycemia that may appear even up to 24 hours later. Symptoms should be described including feeling very hungry, nervous, shaky, sweaty, dizzy, or confused. In order to raise the blood glucose, patients can take glucose tablets or gels, or drink milk or juice, depending on how low the blood sugar falls. Patients should see their physician for adjustment of medication should hypoglycemia occur. A glucagon kit should be prescribed for emergencies in the case of severe hypoglycemia or when the patient is unable to drink or eat. Family members and coworkers should be instructed on how to administer this. For children, school and camp staff and caretakers should be educated on how to deal with low blood sugar.

• The American Diabetes Association (ADA) defines a glucose alert value as <70 mg/dL (<3.9 mmol/L), requiring treatment with fast-acting carbohydrate and dose adjustment of glucose-lowering therapy. Clinically significant hypoglycemia is defined as <54 mg/dL (<3.0 mmol/L), indicating serious, clinically important hypoglycemia.[1] Severe hypoglycemia is defined as any low blood glucose level leading to cognitive impairment requiring assistance from another person for recovery.

• Patients should discuss their insulin requirement with their physician prior to skipping any meals - for example, for a medical test.

• Patients should also be educated on the symptoms of hyperglycemia, including blurred vision, thirst, frequent urination, or tiredness, and should see their physician immediately if these occur. Patients should also seek medical attention if they develop a fever, cough, dysuria, or wounds on the feet. If patients are sick or if they note a fingerstick >250 mg/dL (>13.9 mmol/L) on 2 successive premeal checks, they should check their urine ketones and call their physician if the ketones are positive.

• If patients smoke, they should be strongly advised to quit, and offered appropriate treatments as needed.

• Physicians should check patients’ cholesterol and blood pressure, and assess for neuropathy at office visits.

• Patients should be up to date with their vaccination schedule.

• Patients should be encouraged to discuss any feelings of depression with their physicians so that appropriate treatment can be offered.

• Children can take part in all activities at home or school. Staff members at school or camp should assist with a child's needs, including checking blood sugars, taking insulin as needed, eating regular meals, and treating any low blood sugars.

• Further information is available at the ADA website. [American Diabetes Association]

### Complications

<table>
<thead>
<tr>
<th>Complications</th>
<th>Timeframe</th>
<th>Likelihood</th>
</tr>
</thead>
<tbody>
<tr>
<td>diabetic ketoacidosis (DKA)</td>
<td>short term</td>
<td>high</td>
</tr>
</tbody>
</table>
DKA is the classical acute complication of type 1 diabetes, characterized by hyperglycemia and metabolic acidosis.

The most common precipitants are missed insulin injections or physiologic stresses such as infection or myocardial infarction.

Workup (e.g., ECG, search for infection) is indicated to detect precipitating factors.

In the setting of insulin deficiency, stress hormones including glucagon, cortisol, and catecholamines raise blood glucose levels and stimulate ketogenesis.

Hyperglycemia and ketosis cause osmotic diuresis leading to dehydration.

Symptoms tend to be due to dehydration and metabolic acidosis and include dry mouth, shortness of breath, abdominal pain, nausea, vomiting, and altered sensorium.

Blood glucose and ketone levels are high and there is an anion gap metabolic acidosis.

Treatment involves rapid hydration, insulin infusion, and correction of electrolyte imbalance. Hourly monitoring of blood glucose and 1- to 4-hourly monitoring of electrolytes is required. Insulin infusion must continue until ketosis has resolved and a subcutaneous injection of insulin has been given.

Closure of the anion gap will indicate correction of the ketoacidosis.

Potassium repletion is usually indicated because initially apparently normal serum potassium does not reflect true total body depletion.

Treatment with bicarbonate is not indicated except when arterial blood pH is less than 6.9. Serum phosphorus level is usually low, but does not require replacement unless it is less than 1.0 mg/dL (0.323 mmol/L).[97]

<table>
<thead>
<tr>
<th>Complications</th>
<th>Timeframe</th>
<th>Likelihood</th>
</tr>
</thead>
<tbody>
<tr>
<td>hypoglycemia</td>
<td>short term</td>
<td>high</td>
</tr>
</tbody>
</table>
Type 1 diabetes mellitus

Follow up

Complications | Timeframe | Likelihood
---|---|---
The main complication of insulin treatment is hypoglycemia. The American Diabetes Association defines a glucose alert value as <70 mg/dL (<3.9 mmol/L), requiring treatment with fast-acting carbohydrate and dose adjustment of glucose-lowering therapy. Clinically significant hypoglycemia is defined as <54 mg/dL (<3.0 mmol/L), indicating serious, clinically important hypoglycemia. Severe hypoglycemia is defined as any low blood glucose level leading to cognitive impairment requiring assistance from another person for recovery.

Patients with type 1 diabetes are generally sensitive to insulin. Therefore, even a slightly higher dose of insulin, decreased food intake, or increased physical activity can lead to hypoglycemia. Children <6 to 7 years old may not be aware of hypoglycemia, necessitating less stringent goals for glucose control. Other risk factors for hypoglycemia include a prior episode of hypoglycemia, hypoglycemic unawareness, autonomic neuropathy, and long duration of diabetes. Alcohol and exercise can cause delayed hypoglycemia, up to 24 hours after the event.

If the patient is able to ingest orally, hypoglycemia can be treated with 4 ounces of fruit juice or sweetened fluids or glucose tablets (15-20 g of carbohydrate). Blood sugar should be tested and treatment effect apparent in 15 minutes.

If oral intake is not possible, an injection of glucagon or intravenous dextrose is required.

Patient caregivers and family members of patients with type 1 diabetes should be educated about the signs and symptoms of hypoglycemia and taught how to administer oral glucose, or intramuscular or deep subcutaneous glucagon. Unless hypoglycemia is recurring, the next meal or snack should be eaten and the next dose of basal insulin should be given.

Episodes of hypoglycemia and the possibility of hypoglycemia unawareness should be assessed at each visit. A strict period of several weeks without hypoglycemia may improve hypoglycemic awareness in some patients.

| retinopathy | long term | high |
---|---|---|
Retinopathy is the most common microvascular complication of diabetes and its risk is increased at all levels of A1C (glycosylated hemoglobin) above the nondiabetic range. The incidence is 1 per 100 person-years for a mean A1C value of 5.5% and 9.5 per 100 person-years for a mean A1C value of 10.5%. There is an increased risk of retinopathy in women with pre-existing type 1 diabetes during pregnancy.

Twenty years after diagnosis, most patients have evidence of retinopathy. Patients develop microaneurysms, exudates, hemorrhages, angiogenesis, and glaucoma.

Retinopathy is usually asymptomatic until its late stages, so screening is essential.

Primary prevention includes strict glycemic control. Progression of very mild to moderate nonproliferative retinopathy can be delayed through glycemic, blood pressure, and lipid control. In advanced disease, photocoagulation and vitrectomy can be done to prevent blindness. Intravitreal injections of antivascular endothelial growth factors are given for center-involved macular edema.

| diabetic kidney disease | long term | high |
---|---|---|

Diabetic kidney disease is the most common cause of end-stage renal disease in developed countries. Although albuminuria occurs in 20% to 40% of patients, the prevalence of end-stage renal disease is 2.2% at 20 years and 7.7% at 30 years of onset of type 1 diabetes.[98]

The pathogenesis of diabetic nephropathy involves glomerular mesangial sclerosis leading to proteinuria and progressive decline in glomerular filtration. Increased urinary albumin excretion (>30 mg/day) is the earliest sign of disease and a marker of much increased cardiovascular risk. Test yearly in people who have had type 1 diabetes for 5 years or more.[1]

Glycemic control and blood pressure control with an angiotensin-converting enzyme (ACE) inhibitor, or angiotensin-II receptor blocker delays onset and slows progression of disease.[99]

Dietary protein limitation, if protein intake is high, can be considered in patients whose diabetic kidney disease is progressing despite optimal glucose and blood pressure control and use of an ACE inhibitor or angiotensin-II receptor blocker.[1]

More than 50% of patients will develop neuropathy.[100]

Strict glycemic control prevents onset and delays progression of diabetic neuropathy, which manifests most commonly as distal symmetric polyneuropathy affecting sensory axons.

The duration and extent of hyperglycemia are the greatest risk factors, although other cardiovascular risk factors probably also contribute.

The other most common types of neuropathy include mononeuropathy, mononeuritis multiplex, polyradiculopathies, and autonomic neuropathy.

Once distal symmetric polyneuropathy is diagnosed, simple inspection should be performed at 3- to 6-month intervals, and referral for podiatric care and special footwear should be made. There are several medications that are particularly effective and may be considered. In the US, Food and Drug Administration-approved medications for diabetic neuropathic pain include pregabalin, duloxetine, and tapentadol. Other treatments that are not approved for this indication may also be helpful, including tricyclic antidepressants, anticonvulsants, a 5-hydroxytryptamine and norepinephrine uptake inhibitor, or capsaicin cream.[1]

For autonomic neuropathy, current treatments for this complication are mostly inadequate. However, symptom management can be considered: for example, compressive stockings for postural hypotension.[1]

<table>
<thead>
<tr>
<th>Complications</th>
<th>Timeframe</th>
<th>Likelihood</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diabetic kidney disease</td>
<td>long term</td>
<td>high</td>
</tr>
<tr>
<td>peripheral or autonomic neuropathy</td>
<td>long term</td>
<td>high</td>
</tr>
<tr>
<td>cardiovascular disease</td>
<td>long term</td>
<td>high</td>
</tr>
</tbody>
</table>
Complications | Timeframe | Likelihood
---|---|---
Cardiovascular disease is the major cause of death and a major cause of morbidity for patients with diabetes.

Intensive glycemic control has been shown to decrease the incidence of macrovascular disease in type 1 diabetes.[93]

The cardiovascular disease risk can be further decreased by modification of other cardiovascular risk factors. Lifestyle and behavioral therapy are essential components of treatment.

Hypertension is often secondary to underlying nephropathy in patients with type 1 diabetes. Blood pressure should be treated to less than 140/80 mm Hg with an ACE inhibitor or angiotensin-II receptor blocker; most patients will require 2 or 3 drugs to reach goal.

For patients of all ages with diabetes and overt cardiovascular disease, high-intensity statin therapy should be added to lifestyle therapy. For patients without known cardiovascular disease, individualization of statin therapy according to their cardiovascular disease risk score is recommended. [ASCVD risk estimator] [101]

Intensive lifestyle therapy and optimal glycemic control are recommended to decrease cardiovascular risk in patients with triglycerides ≥150 mg/dL (≥1.7 mmol/L) and/or HDL <40 mg/dL (<1 mmol/L) (<50 mg/dL [<1.3 mmol/L] among women).[1] There is no specific LDL target.

Children should have a fasting lipid profile at age 10 years or soon after the diagnosis of diabetes once adequate glucose control is achieved.[1] Monitoring can be every 3 to 5 years if LDL <100 mg/dL (≤2.6 mmol/L); otherwise annual monitoring is reasonable. The optimal pharmacologic treatment of hyperlipidemia in children has not been clearly defined, although an initial approach to lipid lowering should include modifications to diet and increased exercise. Statins are not approved for children <10 years of age.

All adult patients with diabetes and cardiovascular disease should be treated with aspirin for secondary prevention (75-162 mg/day). Aspirin can be considered for primary prevention for men and women who have a 10-year risk of atherosclerotic cardiovascular disease risk of over 10%. [1] All patients should have smoking-cessation counseling and treatment as needed.

Patients aged >55 years old with or without hypertension, but with cardiovascular disease, dyslipidemia, increased urinary albumin excretion, or smoking, may benefit from an ACE inhibitor to reduce the risk of cardiovascular events.[102]

No evidence-based guidelines exist for screening asymptomatic patients for coronary heart disease.[1]

Prognosis

Untreated type 1 diabetes is a fatal condition due to diabetic ketoacidosis. Poorly controlled type 1 diabetes is a risk factor for chronic complications such as blindness, renal failure, foot amputations, and heart attacks. Intensive glycemic control has been shown to decrease the incidence of microvascular and macrovascular disease in type 1 diabetes,[89] [90] [91] [92] [93] and the decreased incidence of macrovascular disease has been shown to persist for up to 30 years.[94] Even a few years of intensive glucose control translate to reduced rates of microvascular and macrovascular complications 10 years later.[90] [95] The American Diabetes Association recommends maintaining A1C (glycosylated hemoglobin) <7% to prevent complications in most nonpregnant adults with type 1 diabetes, with less stringent goals in children and adolescents.[1]
Overall, cardiovascular disease is the major cause of death and a major cause of morbidity for patients with diabetes. One analysis of patients with type 1 diabetes diagnosed before the age of 15 years found that the leading cause of death before the age of 30 years was acute complications of diabetes. After the age of 30 years cardiovascular disease was predominant, although death attributable to acute complications was still important in this age group.[96]

With careful planning and adequate treatment, most women with type 1 diabetes can have successful pregnancies.
Diagnostic guidelines

**International**


Published by: American Diabetes Association  
Last published: 2018

Type 1 diabetes Centers for Disease Control and Prevention. Type 1 diabetes. October 2017 [internet publication].

Published by: Centers for Disease Control and Prevention  
Last published: 2017


Published by: The Endocrine Society  
Last published: 2016

Treatment guidelines

**International**


Published by: American Diabetes Association  
Last published: 2018

Type 1 diabetes Centers for Disease Control and Prevention. Type 1 diabetes. October 2017 [internet publication].

Published by: Centers for Disease Control and Prevention  
Last published: 2017


Published by: The Endocrine Society  
Last published: 2016
Online resources

1. ASCVD risk estimator (external link)

2. American Diabetes Association (external link)
Key articles


References


<table>
<thead>
<tr>
<th>Reference</th>
<th>Details</th>
</tr>
</thead>
</table>


Type 1 diabetes mellitus

References


74. US Food and Drug Administration. FDA approves Afrezza to treat diabetes. June 2014 [internet publication]. Full text


<table>
<thead>
<tr>
<th>Reference</th>
<th>Title and Details</th>
</tr>
</thead>
</table>
Disclaimer

This content is meant for medical professionals. The BMJ Publishing Group Ltd ("BMJ Group") tries to ensure that the information provided is accurate and up to date, but we do not warrant that it is. The BMJ Group does not advocate or endorse the use of any drug or therapy contained within nor does it diagnose patients. Medical professionals should use their own professional judgement in using this information and caring for their patients and the information herein should not be considered a substitute for that.

This information is not intended to cover all possible diagnosis methods, treatments, follow up, drugs and any contraindications or side effects. We strongly recommend that users independently verify specified diagnosis, treatments and follow up and ensure it is appropriate for your patient. This information is provided on an "as is" basis and to the fullest extent permitted by law the BMJ Group assumes no responsibility for any aspect of healthcare administered with the aid of this information or any other use of this information.

View our full Website Terms and Conditions.

Contact us
+1 855-458-0579 (toll free from USA)
ussupport@bmj.com

BMJ Americas Office
2 Hudson Place, Suite 300
Hoboken, New Jersey 07030
Contributors:

// Authors:

Rajesh K. Garg, MD
Professor of Medicine
University of Miami, Miller School of Medicine, Miami, FL
DISCLOSURES: RKG is an author of a number of references cited in this topic.

Varsha Vimalananda, MD, MPH
Assistant Professor
Section of Endocrinology, Diabetes, Nutrition and Weight Management, Boston University School of Medicine, Boston, Core Investigator, Center for Healthcare Organization and Implementation Research, Edith Nourse Rogers Memorial VA Medical Center, Bedford, MA
DISCLOSURES: VV declares that she has no competing interests.

// Peer Reviewers:

Zachary Bloomgarden, MD
Clinical Professor
Medicine/Endocrinology, Diabetes and Bone Disease, Mount Sinai School of Medicine, New York, NY
DISCLOSURES: ZB declares that he has no competing interests.

Alicia Jenkins, MB, BS, MD, FRACP, FRCP
Associate Professor
Department of Medicine, University of Melbourne, Melbourne, Australia, Professor, Harold Hamm Oklahoma Diabetes Center, University of Oklahoma Health Sciences Center, Oklahoma City, OK
DISCLOSURES: AJ has been a (non-salaried) co-investigator on multi-center clinical trials supported by Novo, Eli Lilly, Sanofi-Aventis, and Medtronic. She does not hold any stocks or shares in these companies. She has received a speaker’s honorarium from Novo.