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# **Type 1 diabetes: diagnosis and management of type 1 diabetes in adults**

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## **NICE guideline**

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### **Draft for consultation, December 2014**

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If you wish to comment on this version of the guideline, please be aware that all the supporting information and evidence is contained in the full version.

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## 1 **Introduction**

2 Type 1 diabetes affects over 370,000 adults in the UK. It results from  
3 destruction of the cells that normally make insulin. Loss of insulin secretion  
4 results in high blood glucose concentrations and other metabolic and  
5 haematological abnormalities, which have both acute and long-term adverse  
6 effects on health. Over years, type 1 diabetes causes tissue damage which, if  
7 not detected and managed early, can result in disability: blindness, kidney  
8 failure and foot ulceration leading to amputation, as well as premature heart  
9 disease, stroke and death. The risk of all of these complications is greatly  
10 reduced by treatment that re-establishes control of circulating glucose  
11 concentrations to as near normal levels as possible, reducing tissue damage.  
12 Disability from complications that are not avoided by strict diabetes control  
13 can often be prevented by early detection and active management.

14 Treatment of type 1 diabetes is by insulin replacement, supported by active  
15 medical management of other cardiovascular risk factors, such as  
16 hypertension and high circulating lipids. Modern insulin replacement therapy  
17 aims to re-create normal fluctuations in circulating insulin concentrations. This  
18 supports a flexible lifestyle with minimal restrictions and, properly done, can  
19 improve diabetes control, which in turn reduces the risk of structural  
20 complications and of episodes of hypoglycaemia (low blood glucose, a  
21 common complication of insulin therapy caused by overestimation of the dose  
22 required at the time). Flexible insulin therapy usually involves self-injecting  
23 multiple daily doses of insulin, with doses adjusted based on taken or planned  
24 exercise, intended food intake and other factors, including the current blood  
25 glucose concentration, which the insulin user needs to test on a regular basis.  
26 This self-management needs the insulin user to have the skills and confidence  
27 to manage the regimen. One of the most important roles of healthcare  
28 professionals providing diabetes care to adults with type 1 diabetes is to  
29 ensure that systems are in place to provide informed, expert support,  
30 education and training for insulin users, as well as a range of other more  
31 conventional biomedical services and interventions.

1 Although type 1 diabetes in adults is not rare, it is not common enough that all  
2 healthcare professionals who deal with it are able to acquire and maintain all  
3 the necessary skills for its management. The aim of this guideline is to provide  
4 evidence-based, practical advice on the steps necessary to support adults  
5 with type 1 diabetes to live full, largely unrestricted, lives and to avoid the  
6 acute and long-term complications of both the disease and of its treatment.

### 7 ***Reasons for the update***

8 NICE last produced a guideline on type 1 diabetes in 2004. Since then, life  
9 expectancy for adults living with type 1 diabetes has increased but it remains  
10 significantly shorter than for people without diabetes. There remain important  
11 deficiencies in care provision, most adults with type 1 diabetes do not achieve  
12 the target measures of diabetic control, and rates of diabetic ketoacidosis (the  
13 acute complication of insulin deficiency) and of renal failure have increased.  
14 The present guideline is an update of many sections of the 2004 guideline,  
15 focusing on areas where new knowledge and treatment opportunities have  
16 arisen in the last decade. There have been many developments, including  
17 improvements in technology to support better diabetes control, that should  
18 result in improved outcomes for adults with type 1 diabetes. These changes  
19 also present more challenges in terms of the diversity and complexity of the  
20 tools that can now be provided to adults with type 1 diabetes to help them  
21 achieve best outcomes, and this guideline describes evidence-based best  
22 practice for their deployment.

23 Topics updated from the 2004 guideline include:

- 24 • diagnosing type 1 diabetes
- 25 • structured education programmes
- 26 • insulin preparations and regimens associated with improved diabetes  
27 control
- 28 • needle length for insulin injections
- 29 • new technologies for glucose monitoring and insulin delivery
- 30 • managing acute painful neuropathy associated with rapid blood glucose  
31 control, erectile dysfunction in men and gastroparesis

- 1 • primary prevention of cardiovascular disease.

2 The following topics were not included in 2004 and have been added:

- 3 • new insulin formulations  
4 • identifying, quantifying and managing impaired awareness of  
5 hypoglycaemia  
6 • monitoring for thyroid disease  
7 • use of blood ketone measurement in preventing and monitoring diabetic  
8 ketoacidosis  
9 • carbohydrate counting and glycaemic index diets  
10 • referral criteria for transplantation therapies.

11 This guideline is intended to describe the methods for achieving optimal  
12 outcomes for adults with type 1 diabetes and to inform service design and  
13 delivery. Its intended audience therefore includes healthcare professionals  
14 involved in delivering services to adults with type 1 diabetes, service  
15 managers and commissioners, and adults with type 1 diabetes and their  
16 families.

### 17 ***Medicines***

18 The guideline will assume that prescribers will use a medicine's summary of  
19 product characteristics to inform decisions made with individual patients.

20 This guideline recommends a medicine for an indication for which it does not  
21 have a UK marketing authorisation at the date of publication (domperidone for  
22 gastroparesis). The prescriber should follow relevant professional guidance,  
23 taking full responsibility for the decision. The patient (or those with authority to  
24 give consent on their behalf) should provide informed consent, which should  
25 be documented. See the General Medical Council's [Prescribing guidance:  
26 prescribing unlicensed medicines](#) for further information. Where  
27 recommendations have been made for the use of medicines outside their  
28 licensed indications ('off-label use'), these medicines are marked with a  
29 footnote in the recommendations.

30

## 1 **Patient-centred care**

2 This guideline offers best practice advice on the care of adults with type 1  
3 diabetes.

4 Patients and healthcare professionals have rights and responsibilities as set  
5 out in the [NHS Constitution for England](#) – all NICE guidance is written to  
6 reflect these. Treatment and care should take into account individual needs  
7 and preferences. Patients should have the opportunity to make informed  
8 decisions about their care and treatment, in partnership with their healthcare  
9 professionals. If the patient is under 16, their family or carers should also be  
10 given information and support to help the child or young person to make  
11 decisions about their treatment. Healthcare professionals should follow the  
12 [Department of Health's advice on consent](#). If someone does not have capacity  
13 to make decisions, healthcare professionals should follow the [code of practice](#)  
14 [that accompanies the Mental Capacity Act](#) and the supplementary [code of](#)  
15 [practice on deprivation of liberty safeguards](#).

16 NICE has produced guidance on the components of good patient experience  
17 in adult NHS services. All healthcare professionals should follow the  
18 recommendations in [Patient experience in adult NHS services](#).

19

## 1 **Strength of recommendations**

2 Some recommendations can be made with more certainty than others. The  
3 Guideline Development Group makes a recommendation based on the trade-  
4 off between the benefits and harms of an intervention, taking into account the  
5 quality of the underpinning evidence. For some interventions, the Guideline  
6 Development Group is confident that, given the information it has looked at,  
7 most patients would choose the intervention. The wording used in the  
8 recommendations in this guideline denotes the certainty with which the  
9 recommendation is made (the strength of the recommendation).

10 For all recommendations, NICE expects that there is discussion with the  
11 patient about the risks and benefits of the interventions, and their values and  
12 preferences. This discussion aims to help them to reach a fully informed  
13 decision (see also 'Patient-centred care').

### 14 ***Interventions that must (or must not) be used***

15 We usually use 'must' or 'must not' only if there is a legal duty to apply the  
16 recommendation. Occasionally we use 'must' (or 'must not') if the  
17 consequences of not following the recommendation could be extremely  
18 serious or potentially life threatening.

### 19 ***Interventions that should (or should not) be used – a 'strong'*** 20 ***recommendation***

21 We use 'offer' (and similar words such as 'refer' or 'advise') when we are  
22 confident that, for the vast majority of patients, an intervention will do more  
23 good than harm, and be cost effective. We use similar forms of words (for  
24 example, 'Do not offer...') when we are confident that an intervention will not  
25 be of benefit for most patients.

### 26 ***Interventions that could be used***

27 We use 'consider' when we are confident that an intervention will do more  
28 good than harm for most patients, and be cost effective, but other options may  
29 be similarly cost effective. The choice of intervention, and whether or not to

1 have the intervention at all, is more likely to depend on the patient's values  
2 and preferences than for a strong recommendation, and so the healthcare  
3 professional should spend more time considering and discussing the options  
4 with the patient.

5 ***Recommendation wording in guideline updates***

6 NICE began using this approach to denote the strength of recommendations  
7 in guidelines that started development after publication of the 2009 version of  
8 'The guidelines manual' (January 2009). This does not apply to any  
9 recommendations shaded in grey and ending **[2004]** (see 'Update information'  
10 box below for details about how recommendations are labelled). In particular,  
11 for recommendations labelled **[2004]**, the word 'consider' may not necessarily  
12 be used to denote the strength of the recommendation.

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## **Update information**

This guidance is an update of NICE guideline CG15 (published September 2004) and will replace the guidance for adults.

### ***Recommendations with an evidence review***

New recommendations have been added on diagnosis, monitoring and treatment for adults with type 1 diabetes.

You are invited to comment on the new and updated recommendations in this guideline. These are marked as:

- **[new 2015]** if the evidence has been reviewed and the recommendation has been added or updated
- **[2015]** if the evidence has been reviewed but no change has been made to the recommended action.

You are also invited to comment on recommendations that NICE proposes to delete from the 2004 guideline, because either the evidence has been reviewed and the recommendations have been updated, or NICE has updated other relevant guidance and has replaced the original recommendations. Appendix A sets out these recommendations and includes details of replacement recommendations. Where there is no replacement recommendation, an explanation for the proposed deletion is given.

### ***Recommendations without an evidence review***

NICE is piloting a new process for identifying and labelling changes to recommendations that have not undergone an evidence review as part of the update. In this guideline:

- minor editorial changes that do not affect the content of the recommendation have not been highlighted in yellow
- the definition of an 'amended' recommendation has been expanded (see below).

Where recommendations are shaded in grey and end **[2004]**, the evidence has not been reviewed since the original guideline. We will not be able to accept comments on these recommendations.

Where recommendations are shaded in grey and end **[2004, amended 2015]**, the evidence has not been reviewed but either:

- changes have been made to the recommendation wording that change the meaning (for example, because of equalities duties or a change in the availability of drugs, or incorporated guidance has been updated) **or**
- NICE has made editorial changes to the original wording to clarify the action to be taken.

These changes are marked with yellow shading, and explanations of the reasons for the changes are given in appendix A for information. We will not routinely accept comments on these recommendations, but will respond if particular concerns are raised around the proposed amendments.

The original NICE guideline and supporting documents are available [here](#).

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## 1 Key priorities for implementation

2 The following recommendations have been identified as priorities for  
3 implementation. The full list of recommendations is in [section 1](#).

### 4 Education and information

- 5 • Offer all adults with type 1 diabetes a structured education programme of  
6 proven benefit, for example the [DAFNE \(dose adjustment for normal  
7 eating\) programme](#). Offer this programme 6–12 months after diagnosis, at  
8 a time that is clinically appropriate and suitable for the person. **[new 2015]**  
9 **[1.3.1]**

### 10 Blood glucose control

- 11 • Support adults with type 1 diabetes to achieve and maintain a target HbA1c  
12 level of 48 mmol/mol (6.5%) or lower, to minimise the risk of long-term  
13 vascular complications. **[new 2015] [1.6.6]**
- 14 • Agree an individualised HbA1c target with each adult with type 1 diabetes,  
15 taking into account factors such as the person's daily activities, aspirations,  
16 likelihood of complications, comorbidities, occupation and history of  
17 hypoglycaemia. **[new 2015] [1.6.7]**
- 18 • Support adults with type 1 diabetes to test at least 4 times a day, and up to  
19 10 times a day if any of the following apply:
  - 20 – the target for blood glucose control, measured by HbA1c level (see  
21 recommendation 1.6.6), is not achieved
  - 22 – the frequency of hypoglycaemic episodes increases
  - 23 – there is a legal requirement to do so (such as before driving, in line with  
24 DVLA requirements)<sup>1</sup>
  - 25 – during periods of illness
  - 26 – before and after sport
  - 27 – when planning pregnancy, during pregnancy and while breastfeeding  
28 (see the NICE guideline on [diabetes in pregnancy](#)<sup>2</sup>)

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<sup>1</sup> For DVLA guidance on type 1 diabetes, see the [DVLA guidance for people with type 1 diabetes](#).

1 – if there is a need to know blood glucose levels more than 4 times a day  
2 for other reasons (for example, impaired awareness of hypoglycaemia,  
3 high-risk activities). **[new 2015] [1.6.13]**

- 4 • Advise adults with type 1 diabetes to aim for:
  - 5 – a fasting plasma glucose level of 5–7 mmol/litre on waking **and**
  - 6 – a plasma glucose level of 4–7 mmol/litre before meals at other times of
  - 7 the day. **[new 2015] [1.6.15]**

#### 8 **Insulin therapy**

- 9 • Offer multiple daily injection basal–bolus insulin regimens, rather than  
10 twice-daily mixed insulin regimens, as the insulin injection regimen of  
11 choice for all adults with type 1 diabetes. **[new 2015] [1.7.2]**

#### 12 **Awareness and management of hypoglycaemia**

- 13 • Assess awareness of hypoglycaemia in adults with type 1 diabetes at each  
14 annual review. **[new 2015] [1.10.1]**

#### 15 **Care of adults with type 1 diabetes in hospital**

- 16 • Enable adults with type 1 diabetes who are hospital inpatients to self-  
17 administer subcutaneous insulin if they are willing and able and it is safe to  
18 do so. **[new 2015] [1.14.6]**

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<sup>2</sup> This guideline is [currently being updated](#) (publication expected February 2015).

## 1 **1 Recommendations**

2 The following guidance is based on the best available evidence. The [full](#)  
3 [guideline](#) **[hyperlink to be added for final publication]** gives details of the  
4 methods and the evidence used to develop the guidance.

### 5 ***Blood glucose and plasma glucose***

6 This guideline refers frequently to circulating glucose concentrations as 'blood  
7 glucose'. A lot of the evidence linking specific circulating glucose  
8 concentrations with particular outcomes uses 'plasma' rather than 'blood'  
9 glucose. In addition, patient-held glucose meters and monitoring systems are  
10 all calibrated to plasma glucose equivalents. However, the term 'blood  
11 glucose monitoring' is in very common use, so in this guideline we use the  
12 term 'blood glucose', except when referring to specific concentration values.

### 13 **1.1 *Diagnosis and early care plan***

#### 14 **Diagnosis**

15 1.1.1 Diagnose type 1 diabetes on clinical grounds in adults presenting  
16 with hyperglycaemia, bearing in mind that people with type 1  
17 diabetes typically (but not always) have one or more of:

- 18 • ketosis
- 19 • rapid weight loss
- 20 • age of onset below 50 years
- 21 • BMI below 25 kg/m<sup>2</sup>
- 22 • personal and/or family history of autoimmune disease. **[new**  
23 **2015]**

24 1.1.2 Do not discount a diagnosis of type 1 diabetes if a person presents  
25 with a BMI of 25 kg/m<sup>2</sup> or above or is aged 50 years or above.  
26 **[new 2015]**

27 1.1.3 Do not measure C-peptide and/or diabetes-specific autoantibody  
28 titres routinely to confirm type 1 diabetes in adults. **[new 2015]**

- 1 1.1.4 Consider further specialist investigation involving measurement of  
2 C-peptide and/or diabetes-specific autoantibody titres if:
- 3 • type 1 diabetes is suspected but the clinical presentation  
4 includes some atypical features (for example, age 50 years or  
5 above, BMI of 25 kg/m<sup>2</sup> or above, slow evolution of  
6 hyperglycaemia or long prodrome) **or**
  - 7 • type 1 diabetes has been diagnosed and treatment started but  
8 there is a clinical suspicion that the person may have a  
9 monogenic form of diabetes, and C-peptide and/or autoantibody  
10 testing may guide the use of genetic testing **or**
  - 11 • classification is uncertain, and confirming type 1 diabetes would  
12 have implications for availability of therapy (for example,  
13 continuous subcutaneous insulin infusion [CSII or 'insulin pump']  
14 therapy). **[new 2015]**

- 15 1.1.5 When measuring C-peptide and/or diabetes-specific autoantibody  
16 titres, take into account that:
- 17 • autoantibody tests have their lowest false negative rate at the  
18 time of diagnosis, and that the false negative rate rises  
19 thereafter
  - 20 • C-peptide has better discriminative value the longer the test is  
21 done after diagnosis
  - 22 • with autoantibody testing, carrying out tests for 2 different  
23 diabetes-specific autoantibodies reduces the false negative rate.  
24 **[new 2015]**

## 25 **Early care plan**

- 26 1.1.6 At the time of diagnosis (or if necessary after the management of  
27 critically decompensated metabolism), the diabetes professional  
28 team should develop with and explain to the adult with type 1  
29 diabetes a plan for their early care. To agree such a plan will  
30 generally require:

- 1           • medical assessment to:
- 2           – ensure security of diagnosis of type of diabetes
- 3           – ensure appropriate acute care is given when needed
- 4           – review and detect potentially confounding disease and
- 5           medicines
- 6           – detect adverse vascular risk factors
- 7           • environmental assessment to understand:
- 8           – the social, home, work and recreational circumstances of the
- 9           person and carers
- 10          – their preferences in nutrition and physical activity
- 11          – other relevant factors, such as substance use
- 12          • cultural and educational assessment to identify prior knowledge
- 13          and to enable optimal advice and planning about:
- 14          – treatment modalities
- 15          – diabetes education programmes
- 16          • assessment of emotional state to determine the appropriate
- 17          pace of education.
- 18          The results of the assessment should be used to agree a future
- 19          care plan. Some items of the initial diabetes assessment:
- 20          • acute medical history
- 21          • social, cultural and educational history/lifestyle review
- 22          • complications history/symptoms
- 23          • long-term/recent diabetes history
- 24          • other medical history/systems
- 25          • family history of diabetes/arterial disease
- 26          • medication history/current medicines
- 27          • vascular risk factors
- 28          • smoking
- 29          • general examination
- 30          • weight/BMI
- 31          • foot/eye/vision examination

- 1 • urine albumin excretion/urine protein/serum creatinine
- 2 • psychological wellbeing
- 3 • attitudes to medicine and self-care
- 4 • immediate family and social relationships and availability of
- 5 informal support. **[2004]**

6 1.1.7 Elements of an individualised and culturally appropriate plan will  
7 include:

- 8 • sites and timescales of diabetes education, including nutritional
- 9 advice (see sections 1.3 and 1.4)
- 10 • initial treatment modalities, including guidance on insulin
- 11 injection (see sections 1.7 and 1.8)
- 12 • means of self-monitoring and targets (see section 1.6)
- 13 • symptoms, risk and treatment of hypoglycaemia
- 14 • management of special situations, such as driving
- 15 • means and frequency of communication with the diabetes
- 16 professional team
- 17 • management of arterial risk factors (see section 1.13)
- 18 • for women of childbearing potential, implications for pregnancy
- 19 and family planning advice
- 20 • follow-up consultations, including frequency of review of HbA1c
- 21 levels and experience of hypoglycaemia, and surveillance at
- 22 annual review (see section 1.15). **[2004, amended 2015]**

23 1.1.8 After the initial plan is agreed, put arrangements in place to  
24 implement it without inappropriate delay, and to provide for  
25 feedback and modification of the plan over the ensuing weeks.  
26 **[2004]**

## 27 **1.2 Care process and support**

28 1.2.1 Advice to adults with type 1 diabetes should be provided by a range  
29 of professionals with skills in diabetes care working together in a  
30 coordinated approach. A common environment (diabetes centre) is

1 an important resource in allowing a diabetes multidisciplinary team  
2 to work and communicate efficiently while providing consistent  
3 advice. **[2004]**

4 1.2.2 Provide adults with type 1 diabetes with:

- 5 • open-access services on a walk-in and telephone-request basis
- 6 during working hours
- 7 • a helpline staffed by people with specific diabetes expertise on a
- 8 24-hour basis
- 9 • contact information for these services. **[2004]**

10 1.2.3 **Regard** each adult with type 1 diabetes as an individual, rather than  
11 as a member of any cultural, economic or health-affected group  
12 (see also recommendations 1.4.5, 1.4.13 and 1.7.1 about the  
13 cultural preferences of individual adults with type 1 diabetes).  
14 **[2004, amended 2015]**

15 1.2.4 Set up an individual care plan jointly agreed with the adult with type  
16 1 diabetes, review it annually and modify it taking into account  
17 changes in the person's wishes, circumstances and medical  
18 findings, and record the details. The plan should include aspects of:

- 19 • diabetes education, including nutritional advice (see sections 1.3
- 20 and 1.4)
- 21 • insulin therapy, **including dose adjustment** (see sections 1.8 and
- 22 1.9)
- 23 • self-monitoring (see section 1.6)
- 24 • **avoiding hypoglycaemia** and maintaining awareness of
- 25 hypoglycaemia
- 26 • **for women of childbearing potential, family planning,**
- 27 **contraception and pregnancy planning**
- 28 • arterial risk factor surveillance and management (see section
- 29 1.13)
- 30 • **complications** surveillance and management (see section 1.15)

- 1 • means and frequency of communicating with the diabetes  
2 professional team
- 3 • follow-up consultations, including **frequency of review of HbA1c**  
4 **levels and experience of hypoglycaemia, and** next annual  
5 review. **[2004, amended 2015]**

6 1.2.5 Use population, practice-based and clinic diabetes registers (as  
7 specified by the [National service framework for diabetes](#)) to assist  
8 programmed recall for annual review and assessment of  
9 complications and vascular risk. **[2004]**

10 1.2.6 The multidisciplinary team approach should be available to  
11 inpatients with type 1 diabetes, regardless of the reason for  
12 admission (see section 1.14). **[2004]**

13 1.2.7 At the time of diagnosis and periodically thereafter, provide adults  
14 with type 1 diabetes with up-to-date information about diabetes  
15 support groups (local and national), how to contact them and the  
16 benefits of membership. **[2004]**

### 17 **1.3 Education and information**

18 Recommendations in this section update and replace the NICE technology  
19 appraisal guidance on [the use of patient-education models for diabetes](#) for  
20 adults with type 1 diabetes.

21 1.3.1 Offer all adults with type 1 diabetes a structured education  
22 programme of proven benefit, for example the [DAFNE \(dose-](#)  
23 [adjustment for normal eating\) programme](#). Offer this programme 6–  
24 12 months after diagnosis, at a time that is clinically appropriate  
25 and suitable for the person. **[new 2015]**

26 1.3.2 Provide an alternative of equal standard for a person unable or  
27 unwilling to participate in group education. **[new 2015]**

1 1.3.3 Ensure that any structured education programme for adults with  
2 type 1 diabetes includes the following components<sup>3</sup>:

- 3 • It is evidence-based, and suits the needs of the person.
- 4 • It has specific aims and learning objectives, and supports the  
5 person and their family members and carers in developing  
6 attitudes, beliefs, knowledge and skills to self-manage diabetes.
- 7 • It has a structured curriculum that is theory-driven, evidence-  
8 based and resource-effective, has supporting materials, and is  
9 written down.
- 10 • It is delivered by trained educators who have an understanding  
11 of educational theory appropriate to the age and needs of the  
12 person, and who are trained and competent to deliver the  
13 principles and content of the programme.
- 14 • It is quality assured, and reviewed by trained, competent,  
15 independent assessors who measure it against criteria that  
16 ensure consistency.
- 17 • The outcomes are audited regularly. **[new 2015]**

18 1.3.4 Explain to adults with type 1 diabetes that structured education is  
19 an integral part of diabetes care. **[new 2015]**

20 1.3.5 Provide information about type 1 diabetes and its management to  
21 adults with type 1 diabetes at all opportunities from diagnosis  
22 onwards. Follow the principles in the NICE guideline on [patient](#)  
23 [experience in adult NHS services](#). **[new 2015]**

24 1.3.6 **Carry out** more formal review of self-care and needs annually in all  
25 adults with type 1 diabetes. Vary the agenda addressed each year  
26 according to the priorities agreed between the healthcare  
27 professional and the adult with type 1 diabetes. **[2004, amended**  
28 **2015]**

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<sup>3</sup> Components of a structured education programme adapted from the NICE quality standard on [diabetes in adults](#).

- 1 1.3.7 Consider the Blood Glucose Awareness Training (BGAT)  
2 programme for adults with type 1 diabetes who are having  
3 recurrent episodes of hypoglycaemia (see also section 1.10). **[new**  
4 **2015]**

5 **1.4 Dietary management**

6 **Carbohydrate counting**

- 7 1.4.1 Offer carbohydrate-counting training to adults with type 1 diabetes  
8 as part of structured education programmes for self-management  
9 (see section 1.3). **[new 2015]**

- 10 1.4.2 Consider carbohydrate-counting courses for adults with type 1  
11 diabetes who are waiting for a more detailed structured education  
12 programme or are unable to take part in a stand-alone structured  
13 education programme. **[new 2015]**

14 **Glycaemic index diets**

- 15 1.4.3 Do not advise adults with type 1 diabetes to follow a low glycaemic  
16 index diet for blood glucose control. **[new 2015]**

17 **Dietary advice**

- 18 1.4.4 Offer dietary advice to adults with type 1 diabetes about issues  
19 other than blood glucose control, such as weight control and  
20 cardiovascular risk management, as indicated clinically. **[new**  
21 **2015]**

- 22 1.4.5 Provide nutritional information sensitive to personal needs and  
23 culture from the time of diagnosis of type 1 diabetes. **[2004]**

- 24 1.4.6 Provide nutritional information individually and as part of a diabetes  
25 education programme (see section 1.3). Include advice from  
26 professionals with specific and approved training and continuing  
27 accredited education in delivering nutritional advice to people with  
28 health conditions. Offer opportunities to receive nutritional advice at

1 intervals agreed between adults with type 1 diabetes and their  
2 advising professionals. **[2004]**

3 1.4.7 Discuss the hyperglycaemic effects of different foods an adult with  
4 type 1 diabetes wishes to eat in the context of the insulin  
5 preparations chosen to match those food choices. **[2004]**

6 1.4.8 **Make** programmes available to adults with type 1 diabetes to  
7 enable them to make:

- 8 • optimal choices about the variety of foods they wish to consume
- 9 • insulin dose changes appropriate to reduce glucose excursions
- 10 when taking different quantities of those foods. **[2004, amended**
- 11 **2015]**

12 1.4.9 Agree the choice of content, timing and amount of snacks between  
13 meals or at bedtime available to the adult with type 1 diabetes,  
14 based on informed discussion about the extent and duration of the  
15 effects of eating different food types and the insulin preparations  
16 available to match them. Modify those choices based on discussion  
17 of the results of self-monitoring tests. **[2004]**

18 1.4.10 Make information available on:

- 19 • effects of different alcohol-containing drinks on blood glucose
- 20 excursions and calorie intake
- 21 • use of high-calorie and high-sugar **'treats'**. **[2004, amended**
- 22 **2015]**

23 1.4.11 Make information available about the benefits of healthy eating in  
24 reducing arterial risk as part of dietary education in the period after  
25 diagnosis, and according to need and interest at intervals  
26 thereafter. Include information **about fruit** and vegetables, types  
27 and amounts of fat, and ways of making the appropriate nutritional  
28 changes. **[2004, amended 2015]**

1 1.4.12 Modify nutritional recommendations to adults with type 1 diabetes  
2 to take account of associated features of diabetes, including:

- 3 • excess weight and obesity
- 4 • underweight
- 5 • eating disorders
- 6 • hypertension
- 7 • renal failure. **[2004]**

8 1.4.13 Be aware of appropriate nutritional advice on common topics of  
9 concern and interest to adults living with type 1 diabetes, and be  
10 prepared to seek advice from colleagues with more specialised  
11 knowledge. Suggested common topics **include:**

- 12 • **body weight**, energy balance and obesity management
- 13 • cultural and religious diets, feasts and fasts
- 14 • foods sold as 'diabetic'
- 15 • sweeteners
- 16 • dietary fibre intake
- 17 • protein intake
- 18 • vitamin and mineral supplements
- 19 • alcohol
- 20 • matching carbohydrate, insulin and physical activity
- 21 • salt intake in hypertension
- 22 • comorbidities, including nephropathy and renal failure, coeliac  
23 disease, cystic fibrosis or eating disorders
- 24 • use of peer support groups. **[2004, amended 2015]**

## 25 **1.5 Physical activity**

26 1.5.1 Advise adults with type 1 diabetes that physical activity can reduce  
27 their enhanced arterial risk in the medium and longer term. **[2004]**

28 1.5.2 Give adults with type 1 diabetes who choose to integrate increased  
29 physical activity into a more healthy lifestyle information about:

- 1 • appropriate intensity and frequency of physical activity
- 2 • role of self-monitoring of changed insulin and/or nutritional needs
- 3 • effect of activity on blood glucose levels (likely fall) when insulin
- 4 levels are adequate
- 5 • effect of exercise on blood glucose levels when hyperglycaemic
- 6 and hypoinsulinaemic (risk of worsening of hyperglycaemia and
- 7 ketonaemia)
- 8 • appropriate adjustments of insulin dosage and/or nutritional
- 9 intake for exercise and post-exercise periods, and the next 24
- 10 hours
- 11 • interactions of exercise and alcohol
- 12 • further contacts and sources of information. **[2004]**

## 13 **1.6 Blood glucose control**

### 14 **HbA1c measurement and targets**

#### 15 **Measurement**

- 16 1.6.1 Measure HbA1c levels every 3–6 months in adults with type 1  
17 diabetes. **[new 2015]**
- 18 1.6.2 Consider measuring HbA1c levels more often if the person's blood  
19 glucose control is suspected to be changing rapidly; for example, if  
20 the HbA1c level has risen unexpectedly above a previously  
21 sustained target. **[new 2015]**
- 22 1.6.3 Calibrate HbA1c results according to International Federation of  
23 Clinical Chemistry (IFCC) standardisation. **[new 2015]**
- 24 1.6.4 Inform adults with type 1 diabetes of their HbA1c results after each  
25 measurement and ensure that their most recent result is available  
26 at the time of consultation. Follow the principles in the NICE  
27 guideline on [patient experience in adult NHS services](#) about  
28 communication. **[new 2015]**

1 1.6.5 If HbA1c monitoring is invalid (because of disturbed erythrocyte  
2 turnover or abnormal haemoglobin type), estimate trends in blood  
3 glucose control using one of the following:

- 4 • fructosamine estimation
- 5 • quality-controlled blood glucose profiles
- 6 • total glycated haemoglobin estimation (if abnormal  
7 haemoglobins). [2004, amended 2015]

## 8 **Targets**

9 1.6.6 Support adults with type 1 diabetes to achieve and maintain a  
10 target HbA1c level of 48 mmol/mol (6.5%) or lower, to minimise the  
11 risk of long-term vascular complications. [new 2015]

12 1.6.7 Agree an individualised HbA1c target with each adult with type 1  
13 diabetes, taking into account factors such as the person's daily  
14 activities, aspirations, likelihood of complications, comorbidities,  
15 occupation and history of hypoglycaemia. [new 2015]

16 1.6.8 Ensure that achieving, or attempting to achieve, an HbA1c target is  
17 not accompanied by problematic hypoglycaemia. [new 2015]

18 1.6.9 Diabetes services should document the proportion of adults with  
19 type 1 diabetes in a service who achieve an HbA1c level of  
20 53 mmol/mol (7%) or lower. [new 2015]

## 21 **Self-monitoring of blood glucose**

### 22 **Self-monitoring skills**

23 1.6.10 Teach self-monitoring skills at the time of diagnosis and initiation of  
24 insulin therapy. [2004, amended 2015]

25 1.6.11 Carry out a structured assessment annually of self-monitoring  
26 skills, the quality and use made of the results obtained and the  
27 equipment used. Review self-monitoring skills as part of annual

1 review, or more frequently according to need, and reinforce them  
2 where appropriate. **[2004, amended 2015]**

### 3 ***Frequency of self-monitoring of blood glucose***

4 1.6.12 Advise routine self-monitoring of blood glucose levels for all adults  
5 with type 1 diabetes, and recommend testing at least 4 times a day,  
6 including before each meal and before bed. **[new 2015]**

7 1.6.13 Support adults with type 1 diabetes to test at least 4 times a day,  
8 and up to 10 times a day if any of the following apply:

- 9 • the target for blood glucose control, measured by HbA1c level  
10 (see recommendation 1.6.6), is not achieved
- 11 • the frequency of hypoglycaemic episodes increases
- 12 • there is a legal requirement to do so (such as before driving, in  
13 line with DVLA requirements)<sup>4</sup>
- 14 • during periods of illness
- 15 • before and after sport
- 16 • when planning pregnancy, during pregnancy and while  
17 breastfeeding (see the NICE guideline on [diabetes in](#)  
18 [pregnancy](#)<sup>5</sup>)
- 19 • if there is a need to know blood glucose levels more than 4 times  
20 a day for other reasons (for example, impaired awareness of  
21 hypoglycaemia, high-risk activities). **[new 2015]**

22 1.6.14 Enable additional blood glucose testing (more than 10 times a day)  
23 for adults with type 1 diabetes if this is necessary because of the  
24 person's lifestyle (for example, driving for a long period of time<sup>4</sup>,  
25 undertaking high-risk activity or occupation, travel) or if the person  
26 has impaired awareness of hypoglycaemia. **[new 2015]**

### 27 ***Blood glucose targets***

28 1.6.15 Advise adults with type 1 diabetes to aim for:

---

<sup>4</sup> For further details about driving, see the [DVLA guidance for people with type 1 diabetes](#).

<sup>5</sup> This guideline is [currently being updated](#) (publication expected February 2015).

- 1                   • a fasting plasma glucose level of 5–7 mmol/litre on waking **and**  
2                   • a plasma glucose level of 4–7 mmol/litre before meals at other  
3                   times of the day. **[new 2015]**

4   1.6.16       Advise adults with type 1 diabetes who choose to test after meals  
5                   to aim for a plasma glucose level of 5–9 mmol/litre. **[new 2015]**

6   ***Empowering people to self-monitor blood glucose***

7   1.6.17       Educate adults with type 1 diabetes about how to measure their  
8                   blood glucose level, interpret the results and know what action to  
9                   take. **[new 2015]**

10  1.6.18       Support adults with type 1 diabetes to make the best use of data  
11                   from self-monitoring of blood glucose through structured education  
12                   (see recommendations 1.3.1 and 1.3.2). **[new 2015]**

13 ***Sites for self-monitoring of blood glucose***

14  1.6.19       Monitoring **blood glucose** using sites other than **the fingertips**  
15                   **cannot** be recommended as a routine alternative to conventional  
16                   self-monitoring of blood glucose. **[2004, amended 2015]**

17 **Continuous glucose monitoring**

18  1.6.20       Do not offer real-time continuous glucose monitoring routinely to  
19                   adults with type 1 diabetes. **[new 2015]**

20  1.6.21       Consider real-time continuous glucose monitoring for adults with  
21                   type 1 diabetes who are willing to commit to using it at least 70% of  
22                   the time and to calibrate it as needed, and who have any of the  
23                   following that persist despite optimised use of insulin therapy and  
24                   conventional blood glucose monitoring:

- 25                   • more than 1 episode a year of severe hypoglycaemia with no  
26                   obviously preventable precipitating cause  
27                   • complete loss of awareness of hypoglycaemia  
28                   • frequent (more than 2 episodes a week) asymptomatic  
29                   hypoglycaemia that is causing problems with daily activities

- 1 • extreme fear of hypoglycaemia. **[new 2015]**

2 1.6.22 For people who are having continuous glucose monitoring, use the  
3 principles of flexible insulin therapy with either a multiple daily  
4 injection insulin regimen or continuous subcutaneous insulin  
5 infusion (CSII or insulin pump) therapy. **[new 2015]**

6 1.6.23 Continuous glucose monitoring should be provided by a centre with  
7 expertise in its use, as part of strategies to optimise a person's  
8 HbA1c levels and reduce the frequency of hypoglycaemic  
9 episodes. **[new 2015]**

## 10 **1.7 Insulin therapy**

### 11 **Insulin regimens**

12 1.7.1 Discuss and respect cultural preferences in agreeing the insulin  
13 regimen for an adult with type 1 diabetes. **[2004]**

14 1.7.2 Offer multiple daily injection basal–bolus insulin regimens, rather  
15 than twice-daily mixed insulin regimens, as the insulin injection  
16 regimen of choice for all adults with type 1 diabetes. **[new 2015]**

17 1.7.3 Do not offer non-basal–bolus insulin regimens for treating adults  
18 newly diagnosed with type 1 diabetes. **[new 2015]**

### 19 **Long-acting insulin**

20 Recommendations in this section update and replace the NICE technology  
21 appraisal guidance on [the use of long-acting insulin analogues for the](#)  
22 [treatment of diabetes – insulin glargine](#) for adults with type 1 diabetes.

23 1.7.4 Offer twice-daily insulin detemir as basal insulin therapy for adults  
24 with type 1 diabetes. **[new 2015]**

25 1.7.5 Consider, as an alternative basal insulin therapy for adults with type  
26 1 diabetes:

- 1                   • an existing insulin regimen being used by the person that is  
2                   achieving their agreed targets
- 3                   • once-daily insulin glargine if insulin detemir is not tolerated or if  
4                   twice-daily basal insulin injection is not acceptable to the person.

5                   **[new 2015]**

6 1.7.6           For guidance on the use of continuous subcutaneous insulin  
7                   infusion (CSII or insulin pump) therapy for adults with type 1  
8                   diabetes, see [Continuous subcutaneous insulin infusion for the](#)  
9                   [treatment of diabetes mellitus](#) (NICE technology appraisal guidance  
10                  151). **[new 2015]**

### 11 **Rapid-acting insulin**

12 1.7.7           Offer rapid-acting insulin analogues injected before meals, rather  
13                   than rapid-acting soluble human or animal insulins, for mealtime  
14                   insulin replacement for adults with type 1 diabetes. **[new 2015]**

15 1.7.8           Do not advise routine use of rapid-acting insulin analogues after  
16                   meals. **[new 2015]**

17 1.7.9           If an adult with type 1 diabetes has a strong preference for an  
18                   alternative mealtime insulin, respect their wishes and offer the  
19                   preferred insulin. **[new 2015]**

### 20 **Mixed insulin**

21 1.7.10          Consider a twice-daily human mixed insulin regimen for adults with  
22                   type 1 diabetes if a multiple daily injection basal–bolus insulin  
23                   regimen is not possible and a twice-daily mixed insulin regimen is  
24                   chosen. **[new 2015]**

25 1.7.11          Consider a trial of a twice-daily analogue mixed insulin regimen if a  
26                   person using a twice-daily human mixed insulin regimen has  
27                   hypoglycaemia that affects their quality of life. **[new 2015]**

1 **Optimising insulin therapy**

2 1.7.12 For adults with erratic and unpredictable blood glucose control  
3 (hyperglycaemia and hypoglycaemia at no consistent times), rather  
4 than a change in a previously optimised insulin regimen, the  
5 following should be considered:

- 6
- 7 • injection technique
  - 8 • injection sites
  - 9 • self-monitoring skills
  - 10 • knowledge and self-management skills
  - 11 • nature of lifestyle
  - 12 • psychological and psychosocial difficulties
  - 13 • possible organic causes such as gastroparesis. **[2004, amended 2015]**

14 1.7.13 Give clear guidelines and protocols ('sick-day rules') to all adults  
15 with type 1 diabetes to help them to adjust insulin doses  
16 appropriately during periods of illness. **[2004]**

17 **Adjuncts**

18 1.7.14 Consider adding metformin to insulin therapy if an adult with type 1  
19 diabetes and a BMI of 25 kg/m<sup>2</sup> or above wants to improve their  
20 blood glucose control while minimising their effective insulin dose.  
21 **[new 2015]**

22 **1.8 Insulin delivery**

23 1.8.1 Adults with type 1 diabetes who inject insulin should have access to  
24 the insulin injection delivery device they find allows them optimal  
25 wellbeing, often using one or more types of insulin injection pen.  
26 **[2004]**

27 1.8.2 Provide adults with type 1 diabetes who have special visual or  
28 psychological needs with injection devices or needle-free systems  
29 that they can use independently for accurate dosing. **[2004]**

1 1.8.3 Offer needles of different lengths to adults with type 1 diabetes who  
2 are having problems such as pain, local skin reactions and injection  
3 site leakages. **[new 2015]**

4 1.8.4 If possible, choose needles with the lowest acquisition cost to use  
5 with pre-filled and reusable insulin pen injectors. **[new 2015]**

6 1.8.5 Advise adults with type 1 diabetes to rotate insulin injection sites  
7 and avoid repeated injections at the same point within sites. **[new**  
8 **2015]**

9 1.8.6 Provide adults with type 1 diabetes with suitable containers for  
10 collecting used needles. Arrangements should be available for the  
11 suitable disposal of these containers. **[2004]**

12 1.8.7 Check injection site condition **at least** annually and if new problems  
13 with blood glucose control occur. **[2004, amended 2015]**

## 14 **1.9 Referral for islet or pancreas transplantation**

15 1.9.1 Consider referring adults with type 1 diabetes who have recurrent  
16 severe hypoglycaemia that has not responded to other treatments  
17 (see section 1.10) to a centre that assesses people for islet and/or  
18 pancreas transplantation. **[new 2015]**

19 1.9.2 Consider islet or pancreas transplantation for adults with type 1  
20 diabetes with suboptimal diabetes control who have had a renal  
21 transplant and are currently on immunosuppressive therapy. **[new**  
22 **2015]**

## 23 **1.10 Awareness and management of hypoglycaemia**

### 24 **Identifying and quantifying impaired awareness of hypoglycaemia**

25 1.10.1 Assess awareness of hypoglycaemia in adults with type 1 diabetes  
26 at each annual review. **[new 2015]**

1 1.10.2 Use the Gold score or Clarke score to quantify awareness of  
2 hypoglycaemia in adults with type 1 diabetes, checking that the  
3 questionnaire items have been answered correctly. **[new 2015]**

4 1.10.3 Explain to adults with type 1 diabetes that impaired awareness of  
5 the symptoms of plasma glucose levels below 3 mmol/litre is  
6 associated with a significantly increased risk of severe  
7 hypoglycaemia. **[new 2015]**

#### 8 **Strategies for managing impaired awareness of hypoglycaemia**

9 1.10.4 Ensure that adults with type 1 diabetes with impaired awareness of  
10 hypoglycaemia have had structured education in flexible insulin  
11 therapy using basal–bolus regimens and are following its principles  
12 correctly. **[new 2015]**

13 1.10.5 Offer additional education focusing on avoiding and treating  
14 hypoglycaemia to adults with type 1 diabetes who continue to have  
15 impaired awareness of hypoglycaemia after structured education in  
16 flexible insulin therapy. **[new 2015]**

17 1.10.6 Avoid relaxing individualised blood glucose targets as a treatment  
18 for adults with type 1 diabetes with impaired awareness of  
19 hypoglycaemia. **[new 2015]**

20 1.10.7 Review insulin regimens and doses and prioritise strategies to  
21 avoid hypoglycaemia in adults with type 1 diabetes with impaired  
22 awareness of hypoglycaemia, including:

- 23 • reinforcing the principles of structured education
- 24 • offering continuous subcutaneous insulin infusion (CSII or insulin  
25 pump) therapy
- 26 • offering real-time continuous glucose monitoring. **[new 2015]**

27 1.10.8 If impaired awareness of hypoglycaemia is associated with  
28 recurrent severe hypoglycaemia despite these interventions,  
29 consider referring the person to a specialist centre. **[new 2015]**

1 **Preventing and managing hypoglycaemia**

2 1.10.9 Explain to adults with type 1 diabetes that a fast-acting form of  
3 glucose is needed for the management of hypoglycaemic  
4 symptoms or signs in people who are able to swallow. [2004,  
5 amended 2015]

6 1.10.10 Adults with type 1 diabetes with a decreased level of  
7 consciousness as a result of hypoglycaemia and so are unable to  
8 take oral treatment safely should be:

- 9 • given intramuscular glucagon by a family member or friend who  
10 has been shown how to use it (intravenous glucose may be used  
11 by healthcare professionals skilled in obtaining intravenous  
12 access)
- 13 • monitored for response at 10 minutes, and then given  
14 intravenous glucose if their level of consciousness is not  
15 improving significantly
- 16 • then given oral carbohydrate when it is safe to administer it, and  
17 placed under continued observation by a third party who has  
18 been warned of the risk of relapse. [2004, amended 2015]

19 1.10.11 Explain to adults with type 1 diabetes that some hypoglycaemic  
20 episodes are an inevitable consequence of insulin therapy in most  
21 people using any insulin regimen, and that it is advisable that they  
22 should use a regimen that avoids or reduces the frequency of  
23 hypoglycaemic episodes while maintaining as optimal a level of  
24 blood glucose control as is feasible. Make advice available to all  
25 adults with type 1 diabetes to assist in obtaining the best such  
26 balance from any insulin regimen. (See sections 1.7 and 1.8).  
27 [2004]

28 1.10.12 If hypoglycaemia becomes unusually problematic or of increased  
29 frequency, review the following possible contributory causes:

- 1                   • inappropriate insulin regimens (incorrect dose distributions and  
2                   insulin types)  
3                   • meal and activity patterns, including alcohol  
4                   • injection technique and skills, including insulin resuspension  
5                   • injection site problems  
6                   • possible organic causes including gastroparesis  
7                   • changes in insulin sensitivity (including drugs affecting the renin–  
8                   angiotensin system and renal failure)  
9                   • psychological problems  
10                  • previous physical activity  
11                  • lack of appropriate knowledge and skills for self-management.  
12                  **[2004]**

13   1.10.13   Manage nocturnal hypoglycaemia (symptomatic or detected on  
14                  monitoring) by:

- 15                   • reviewing knowledge and self-management skills  
16                   • reviewing current insulin regimen, evening eating habits and  
17                   previous physical activity  
18                   • choosing an insulin type and regimen that is less likely to induce  
19                   low glucose levels **at night. [2004, amended 2015]**

20   1.10.14   Explain to adults with type 1 diabetes that late postprandial  
21                  hypoglycaemia may be managed by eating snacks between meals  
22                  or by using rapid-acting insulin analogues before meals. **[2004]**

23   1.10.15   If early cognitive decline occurs in adults on long-term insulin  
24                  therapy, supplement normal investigations by the consideration or  
25                  investigation of possible brain damage resulting from overt or  
26                  covert hypoglycaemia, and the need to ameliorate this. **[2004]**

1 **1.11 Ketone monitoring and management of diabetic**  
2 **ketoacidosis (DKA)**

3 **Ketone self-monitoring for prevention of DKA**

4 1.11.1 Consider ketone monitoring (blood or urine) as part of 'sick-day  
5 rules' for adults with type 1 diabetes, to facilitate self-management  
6 of an episode of hyperglycaemia. **[new 2015]**

7 **Ketone monitoring in hospital**

8 1.11.2 In adults with type 1 diabetes presenting to emergency services,  
9 consider capillary blood ketone testing if:

- 10 • DKA is suspected **or**  
11 • the person has uncontrolled diabetes with a period of illness,  
12 and urine ketone testing is positive. **[new 2015]**

13 1.11.3 Consider capillary blood ketone testing for inpatient management of  
14 DKA in adults that is incorporated into a formal protocol. **[new**  
15 **2015]**

16 **Management of DKA**

17 1.11.4 Professionals managing DKA in adults should be adequately  
18 trained, including regular updating, and be familiar with all aspects  
19 of its management which are associated with mortality and  
20 morbidity. These topics should include:

- 21 • fluid balance  
22 • acidosis  
23 • cerebral oedema  
24 • electrolyte imbalance  
25 • disturbed interpretation of familiar diagnostic tests (white cell  
26 count, body temperature, ECG)  
27 • respiratory distress syndrome  
28 • cardiac abnormalities  
29 • precipitating causes

- 1 • infection management, including opportunistic infections
- 2 • gastroparesis
- 3 • use of high dependency and intensive care units
- 4 • recommendations 1.11.5 to 1.11.12 in this guideline.

5 Management of DKA in adults should be in line with local clinical  
6 governance. **[2004]**

7 1.11.5 For primary fluid replacement in adults with DKA, **use** isotonic  
8 saline, not given too rapidly except in cases of circulatory collapse.  
9 **[2004]**

10 1.11.6 **Do not** generally use bicarbonate in the management of DKA in  
11 adults. **[2004, amended 2015]**

12 1.11.7 Give intravenous insulin by infusion to adults with DKA. **[2004]**

13 1.11.8 In the management of DKA in adults, once the plasma glucose  
14 concentration has fallen to 10–15 mmol/litre, give glucose-  
15 containing fluids (not more than 2 litres in 24 hours) in **order to**  
16 **allow continued infusion of insulin at a sufficient rate to clear**  
17 **ketones** (for example, 6 units/hour monitored for effect). **[2004,**  
18 **amended 2015]**

19 1.11.9 Begin potassium replacement early in DKA in adults, with frequent  
20 monitoring for the development of hypokalaemia. **[2004]**

21 1.11.10 **Do not** generally use phosphate replacement in the management of  
22 DKA in adults. **[2004, amended 2015]**

23 1.11.11 In adults with DKA whose conscious level is impaired,  
24 consideration should be given to inserting a nasogastric tube,  
25 monitoring urine production using a urinary catheter and giving  
26 heparin. **[2004]**

27 1.11.12 To reduce the risk of catastrophic outcomes in adults with DKA,  
28 **ensure that** monitoring is continuous and that review covers all

1 aspects of clinical management at frequent intervals. [2004,  
2 amended 2015]

### 3 **1.12 Associated illness**

4 1.12.1 In adults with type 1 diabetes who have a low BMI or unexplained  
5 weight loss, assess markers of coeliac disease. [2004]

6 1.12.2 Be alert to the possibility of the development of other autoimmune  
7 disease in adults with type 1 diabetes (including Addison's disease  
8 and pernicious anaemia). For advice on monitoring for thyroid  
9 disease, see recommendation 1.15.5. [2004, amended 2015]

### 10 **1.13 Control of arterial risk**

#### 11 **Aspirin**

12 1.13.1 Do not offer aspirin for the primary prevention of cardiovascular  
13 disease to adults with type 1 diabetes. [new 2015]

#### 14 **Identifying arterial risk**

15 1.13.2 Assess arterial risk factors annually, including:

- 16 • albuminuria
- 17 • smoking
- 18 • blood glucose control
- 19 • blood pressure
- 20 • full lipid profile (including HDL and LDL cholesterol and
- 21 triglycerides)
- 22 • age
- 23 • family history of arterial disease
- 24 • abdominal adiposity. [2004, amended 2015]

25 1.13.3 For guidance on tools for assessing risk of cardiovascular disease  
26 in adults with type 1 diabetes, see recommendation 1.1.9 in the  
27 NICE guideline on [lipid modification](#). [new 2015]

1 **Interventions to reduce risk and manage arterial disease**

2 1.13.4 For guidance on the primary prevention of cardiovascular disease  
3 in adults with type 1 diabetes, see recommendations 1.3.23 to  
4 1.3.25 in the NICE guideline on [lipid modification](#). [new 2015]

5 1.13.5 Give adults with type 1 diabetes who smoke advice on smoking  
6 cessation and use of smoking cessation services, including NICE  
7 guidance-recommended therapies. Reinforce these messages  
8 annually for people who currently do not plan to stop smoking, and  
9 at all clinical contacts if there is a prospect of the person stopping.  
10 **[2004]**

11 1.13.6 Advise young adult non-smokers never to start smoking. **[2004]**

12 1.13.7 Provide intensive management for adults who have had myocardial  
13 infarction or stroke, according to relevant non-diabetes guidelines.  
14 In the presence of angina or other ischaemic heart disease, beta-  
15 adrenergic blockers should be considered. (For use of insulin in  
16 these circumstances, see section 1.14). **For guidance on**  
17 **secondary prevention of myocardial infarction, see the NICE**  
18 **guideline on [MI – secondary prevention](#). [2004, amended 2015]**

19 **Blood pressure management**

20 1.13.8 Intervention levels for recommending blood pressure management  
21 should be 135/85 mmHg unless the adult with type 1 diabetes has  
22 **albuminuria** or 2 or more features of the metabolic syndrome, in  
23 which case it should be 130/80 mmHg. See also recommendations  
24 1.15.19–1.15.21. **[2004]**

25 1.13.9 To allow informed choice by the person with hypertension, discuss  
26 the following with them:

- 27
- reasons for choice of intervention level
  - substantial potential gains from small improvements in blood pressure control
  - possible negative consequences of therapy.
- 28  
29  
30

1 See also recommendations 1.15.19–1.15.20. **[2004, amended**  
2 **2015]**

3 1.13.10 **Start a trial of a renin–angiotensin system blocking drug as first-line**  
4 **therapy for hypertension in adults with type 1 diabetes. [2004,**  
5 **amended 2015]**

6 1.13.11 Provide information to adults with type 1 diabetes on the potential  
7 for lifestyle changes to improve blood pressure control and  
8 associated outcomes, and offer assistance in achieving their aims  
9 in this area. **[2004]**

10 1.13.12 **Do not allow** concerns over potential side effects to inhibit advising  
11 and offering the necessary use of any class of drugs, unless the  
12 side effects become symptomatic or otherwise clinically significant.  
13 In particular:

- 14 • **do not avoid** selective beta-adrenergic blockers **where indicated**  
15 in adults on insulin
- 16 • low-dose thiazides may be combined with beta-blockers
- 17 • when calcium channel antagonists are prescribed, use only long-  
18 acting preparations
- 19 • use direct questioning to detect the potential side effects of  
20 erectile dysfunction, lethargy and orthostatic hypotension with  
21 different drug classes. **[2004, amended 2015]**

22 1.13.13 For guidance on blood pressure management in adults with type 1  
23 diabetes and evidence of renal involvement, see recommendations  
24 1.6.2–1.6.4 in the NICE guideline on [Chronic kidney disease](#). **[new**  
25 **2015]**

## 26 **1.14 Care of adults with type 1 diabetes in hospital**

### 27 **Blood glucose control**

28 1.14.1 Aim for a target plasma glucose level of 5–8 mmol/litre for adults  
29 with type 1 diabetes during surgery or acute illness. **[new 2015]**

- 1 1.14.2 Establish a local protocol for controlling blood glucose levels in  
2 adults with type 1 diabetes during surgery or acute illness to  
3 achieve the target level. **[new 2015]**
- 4 1.14.3 Use intravenous in preference to subcutaneous insulin regimens for  
5 adults with type 1 diabetes:
- 6 • if the person is unable to eat or is predicted to miss more than 1  
7 meal **or**
  - 8 • if an acute situation is expected to result in unpredictable blood  
9 glucose levels – for example, major surgery, high-dose steroid  
10 treatment, inotrope treatment or sepsis **or**
  - 11 • if insulin absorption is expected to be unpredictable, for example  
12 because of circulatory compromise. **[new 2015]**
- 13 1.14.4 Consider continuing the person's existing basal insulin regimen  
14 (including basal rate if they are using continuous subcutaneous  
15 insulin infusion [CSII or insulin pump] therapy) together with  
16 protocol-driven insulin delivery for controlling blood glucose levels  
17 in adults with type 1 diabetes during surgery or acute illness. **[new**  
18 **2015]**
- 19 1.14.5 Use subcutaneous insulin regimens (including rapid-acting insulin  
20 before meals) if an adult with type 1 diabetes and acute illness is  
21 eating. **[new 2015]**
- 22 1.14.6 Enable adults with type 1 diabetes who are hospital inpatients to  
23 self-administer subcutaneous insulin if they are willing and able and  
24 it is safe to do so. **[new 2015]**

#### 25 **Delivery of care**

- 26 1.14.7 From the time of admission, the adult with type 1 diabetes and the  
27 team caring for him or her should receive, on a continuing basis,  
28 advice from a trained multidisciplinary team with expertise in  
29 diabetes. **[2004]**

- 1 1.14.8 Throughout the course of an inpatient admission, respect the  
2 personal expertise of adults with type 1 diabetes (in managing their  
3 own diabetes) and routinely integrate this into ward-based blood  
4 glucose monitoring and **insulin delivery**. **[2004, amended 2015]**
- 5 1.14.9 Throughout the course of an inpatient admission, the personal  
6 knowledge and needs of adults with type 1 diabetes regarding their  
7 dietary requirements should be a major determinant of the food  
8 choices offered to them, except when illness or medical or surgical  
9 intervention significantly disturbs those requirements. **[2004]**
- 10 1.14.10 Members of care teams caring for adults with type 1 diabetes in  
11 institutions, such as nursing homes, residential homes and prisons,  
12 should follow the recommendations in this section. **[2004]**
- 13 1.14.11 Provide optimal insulin therapy, which can be achieved by the use  
14 of intravenous insulin and glucose, to all adults with type 1 diabetes  
15 with threatened or actual stroke. Critical care and emergency  
16 departments should have a protocol for such management. **[2004,**  
17 **amended 2015]**

## 18 **1.15 *Managing complications***

### 19 **Gastroparesis**

- 20 1.15.1 Consider domperidone<sup>6</sup> (in preference to metoclopramide) for  
21 treating gastroparesis<sup>7</sup> in adults with type 1 diabetes. **[new 2015]**
- 22 1.15.2 Consider continuous subcutaneous insulin infusion (CSII or insulin  
23 pump) therapy for adults with type 1 diabetes who have  
24 gastroparesis. **[new 2015]**

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<sup>6</sup> Although this use is common in UK clinical practice, at the time of consultation (December 2014), domperidone did not have a UK marketing authorisation for this indication. The prescriber should follow relevant professional guidance, taking full responsibility for the decision. Informed consent should be obtained and documented. See the General Medical Council's [Prescribing guidance: prescribing unlicensed medicines](#) for further information.

<sup>7</sup> Diagnosis of gastroparesis needing specific therapy can only be made in the absence of hyperglycaemia at the time of testing, because hyperglycaemia induces a physiological delay in gastric emptying.

1 1.15.3 Advise a small-particle-size diet (mashed or pureed food) for  
2 symptomatic relief for adults with type 1 diabetes who have  
3 vomiting caused by gastroparesis. **[new 2015]**

4 1.15.4 Refer adults with type 1 diabetes who have gastroparesis for  
5 specialist advice if the interventions in recommendations 1.15.1,  
6 1.15.2 and 1.15.3 are not beneficial or not appropriate. **[new 2015]**

### 7 **Thyroid disease monitoring**

8 1.15.5 Measure blood thyroid-stimulating hormone (TSH) levels in adults  
9 with type 1 diabetes at annual review. **[new 2015]**

### 10 **Eye disease**

11 1.15.6 Start eye screening for adults newly diagnosed with type 1 diabetes  
12 from diagnosis. **[2004]**

13 1.15.7 Depending on the findings, follow structured eye screening by:

- 14 • routine review in 1 year **or**
- 15 • earlier review **or**
- 16 • referral to an ophthalmologist. **[2004]**

17 1.15.8 Explain the reasons and success of eye screening systems to  
18 adults with type 1 diabetes, so that attendance is not reduced by  
19 lack of knowledge or fear of outcome. **[2004]**

20 1.15.9 Implement digital retinal photography for eye screening  
21 programmes for adults with type 1 diabetes. **[2004]**

22 1.15.10 Use mydriasis with tropicamide when photographing the retina,  
23 after prior agreement with the adult with type 1 diabetes after  
24 discussion of the advantages and disadvantages, including  
25 appropriate precautions for driving. **[2004]**

26 1.15.11 **Make** visual acuity testing a routine part of eye screening  
27 programmes. **[2004, amended 2015]**

1 1.15.12 **Ensure that** emergency review by an ophthalmologist occurs for:

- 2 • sudden loss of vision
- 3 • rubeosis iridis
- 4 • pre-retinal or vitreous haemorrhage
- 5 • retinal detachment. **[2004, amended 2015]**

6 1.15.13 **Ensure that** rapid review by an ophthalmologist occurs for new  
7 vessel formation. **[2004, amended 2015]**

8 1.15.14 Refer to an ophthalmologist for:

- 9 • referable maculopathy:
  - 10 – exudate or retinal thickening within 1 disc diameter of the
  - 11 centre of the fovea
  - 12 – circinate or group of exudates within the macula (the macula
  - 13 is defined here as a circle centred on the fovea, of a diameter
  - 14 the distance between the temporal border of the optic disc
  - 15 and the fovea)
  - 16 – any microaneurysm or haemorrhage within 1 disc diameter of
  - 17 the centre of the fovea, only if associated with a best visual
  - 18 acuity of 6/12 or worse

- 19 • referable pre-proliferative retinopathy:
  - 20 – any venous beading
  - 21 – **any venous reduplication**
  - 22 – any intraretinal microvascular abnormalities (IRMA)
  - 23 – multiple deep, round or blot haemorrhages

24 (If cotton wool spots are present, look carefully for the above  
25 features, but cotton wool spots themselves do not define pre-  
26 proliferative retinopathy)

- 27 • any **large sudden** unexplained drop in visual acuity. **[2004,**  
28 **amended 2015]**

1 **Diabetic kidney disease**

2 1.15.15 For guidance on managing kidney disease in adults with type 1  
3 diabetes, see the NICE guideline on [chronic kidney disease](#). **[new**  
4 **2015]**

5 1.15.16 Ask all adults with type 1 diabetes with or without detected  
6 nephropathy to bring in the first urine sample of the day ('early  
7 morning urine') once a year. Send this for estimation of  
8 albumin:creatinine ratio. Estimation of urine albumin concentration  
9 alone is a poor alternative. Serum creatinine should be measured  
10 at the same time. **[2004]**

11 1.15.17 Suspect other renal disease:

- 12 • in the absence of progressive retinopathy
- 13 • if blood pressure is particularly high
- 14 • if proteinuria develops suddenly
- 15 • if significant haematuria is present
- 16 • in the presence of systemic ill health. **[2004]**

17 1.15.18 Discuss the significance of a finding of **albuminuria** with the person  
18 concerned. **[2004, amended 2015]**

19 1.15.19 Start angiotensin-converting enzyme (ACE) inhibitors and, with the  
20 usual precautions, titrate to full dose in all adults with confirmed  
21 nephropathy (including those with low-level albuminuria  
22 [microalbuminuria] alone) and type 1 diabetes. **[2004]**

23 1.15.20 If ACE inhibitors are not tolerated, substitute angiotensin 2 receptor  
24 antagonists. Combination therapy is not **recommended**. **[2004,**  
25 **amended 2015]**

26 1.15.21 Maintain blood pressure below 130/80 mmHg by addition of other  
27 anti-hypertensive drugs if necessary. **[2004]**

1 1.15.22 Advise adults with type 1 diabetes and nephropathy about the  
2 advantages of not following a high-protein diet. **[2004]**

3 1.15.23 Referral criteria for tertiary care should be agreed between local  
4 diabetes specialists and nephrologists. **[2004]**

5 **Acute painful neuropathy of rapid improvement of blood glucose control**

6 1.15.24 Reassure adults with type 1 diabetes that acute painful neuropathy  
7 resulting from rapid improvement of blood glucose control is a self-  
8 limiting condition that improves symptomatically over time. **[new**  
9 **2015]**

10 1.15.25 Explain to the person that the specific treatments for acute painful  
11 neuropathy resulting from rapid improvement of blood glucose  
12 control:

- 13 • have the aim of making the symptoms tolerable until the
- 14 condition resolves
- 15 • may not relieve pain immediately and may need to be taken
- 16 regularly for several weeks to be effective. **[new 2015]**

17 1.15.26 Use of simple analgesics (paracetamol, aspirin) and local  
18 measures (bed cradles) are recommended as a first step, but if  
19 trials of these measures are ineffective, discontinue them and try  
20 other measures. **[2004]**

21 1.15.27 Do not relax diabetes control to address acute painful neuropathy  
22 resulting from rapid improvement of blood glucose control in adults  
23 with type 1 diabetes. **[new 2015]**

24 1.15.28 If simple analgesia does not provide sufficient pain relief for adults  
25 with type 1 diabetes who have acute painful neuropathy resulting  
26 from rapid improvement of blood glucose control, offer treatment as  
27 described in the NICE guideline on [neuropathic pain –](#)  
28 [pharmacological management](#). Simple analgesia may be continued

1 until the effects of additional treatments have been established.

2 **[new 2015]**

3 1.15.29 When offering medicines for managing acute painful neuropathy  
4 resulting from rapid improvement of blood glucose control to adults  
5 with type 1 diabetes, be aware of the risk of dependency  
6 associated with opioids. **[new 2015]**

### 7 **Erectile dysfunction**

8 1.15.30 Offer men with type 1 diabetes the opportunity to discuss erectile  
9 dysfunction as part of regular review. **[2015]**

10 1.15.31 Offer a phosphodiesterase-5 inhibitor to men with type 1 diabetes  
11 with isolated erectile dysfunction unless contraindicated. Choose  
12 the phosphodiesterase-5 inhibitor with the lowest acquisition cost.  
13 **[new 2015]**

14 1.15.32 Consider referring men to a service offering further assessment  
15 and other medical, surgical or psychological management of  
16 erectile dysfunction if phosphodiesterase-5 inhibitor treatment is  
17 unsuccessful or contraindicated. **[2015]**

### 18 **Diagnosing and managing autonomic neuropathy**

19 1.15.33 In adults with type 1 diabetes who have unexplained diarrhoea,  
20 particularly at night, the possibility of autonomic neuropathy  
21 affecting the gut should be considered. **[2004]**

22 1.15.34 Take care when prescribing antihypertensive medicines not to  
23 expose people to the risks of orthostatic hypotension as a result of  
24 the combined effects of sympathetic autonomic neuropathy and  
25 blood pressure lowering medicines. **[2004]**

26 1.15.35 In adults with type 1 diabetes who have bladder emptying  
27 problems, investigate the possibility of autonomic neuropathy  
28 affecting the bladder, unless other explanations are adequate.  
29 **[2004]**

1 1.15.36 When managing the symptoms of autonomic neuropathy, include  
2 standard interventions for the manifestations encountered (for  
3 example, **for abnormal sweating and postural hypotension**). **[2004,**  
4 **amended 2015]**

5 1.15.37 Anaesthetists should be aware of the possibility of parasympathetic  
6 autonomic neuropathy affecting the heart in adults with type 1  
7 diabetes who are listed for procedures under general anaesthetic  
8 and who have evidence of somatic neuropathy or other  
9 manifestations of autonomic neuropathy. **[2004]**

#### 10 **Managing chronic diabetic neuropathy**

11 1.15.38 For guidance on treating chronic diabetic neuropathy, see the NICE  
12 guideline on [neuropathic pain – pharmacological management](#).  
13 **[new 2015]**

#### 14 **Diabetic foot problems**

15 1.15.39 For guidance on managing foot problems in adults with type 1  
16 diabetes, see the NICE guideline on [diabetic foot problems](#). **[new**  
17 **2015]**

#### 18 **Psychological problems**

19 1.15.40 Members of diabetes professional teams providing care or advice  
20 to adults with type 1 diabetes should be alert to the development or  
21 presence of clinical or subclinical depression and/or anxiety, in  
22 particular if someone reports or appears to be having difficulties  
23 with self-management. **[2004]**

24 1.15.41 Diabetes professionals should:

- 25 • ensure that they have appropriate skills in the detection and
- 26 basic management of non-severe psychological disorders in
- 27 people from different cultural backgrounds
- 28 • be familiar with appropriate counselling techniques and drug
- 29 therapy, while arranging prompt referral to specialists of those

1 people in whom psychological difficulties continue to interfere  
2 significantly with wellbeing or diabetes self-management.

3 See also the NICE guidelines on [common mental health disorders](#),  
4 [generalised anxiety disorder and panic disorder \(with or without](#)  
5 [agoraphobia\) in adults](#) and [depression in adults with a chronic](#)  
6 [health problem](#). [2004, amended 2015]

## 7 Eating disorders

8 1.15.42 Members of diabetes professional teams should be alert to the  
9 possibility of bulimia nervosa, anorexia nervosa and insulin dose  
10 manipulation in adults with type 1 diabetes with:

- 11 • over-concern with body shape and weight
- 12 • low BMI
- 13 • hypoglycaemia
- 14 • poor overall blood glucose control.

15 See also the NICE guideline on [eating disorders](#). [2004, amended  
16 2015]

17 1.15.43 The risk of morbidity from the complications of poor metabolic  
18 control suggests that consideration should be given to early, and  
19 occasionally urgent, referral of adults with type 1 diabetes to local  
20 eating disorder services. [2004]

21 1.15.44 Make provision for high-quality professional team support at regular  
22 intervals with regard to counselling about lifestyle issues and  
23 particularly dietary behaviour for all adults with type 1 diabetes from  
24 the time of diagnosis (see sections 1.3 and 1.4). [2004]

## 25 2 Research recommendations

26 The Guideline Development Group has made the following recommendations  
27 for research, based on its review of evidence, to improve NICE guidance and

1 patient care in the future. The Guideline Development Group's full set of  
2 research recommendations is detailed in the full guideline.

3 **2.1 *Improved methods and interventions for achieving***  
4 ***HbA1c targets in adults with type 1 diabetes***

5 What methods and interventions are effective in increasing the number of  
6 adults with type 1 diabetes who achieve the recommended HbA1c targets  
7 without risking severe hypoglycaemia or weight gain?

8 **Why this is important**

9 The evidence that sustained near-normoglycaemia substantially reduces the  
10 risk of long-term complications in adults with type 1 diabetes is unequivocal.  
11 Current methods for achieving such blood glucose control require skills in  
12 glucose monitoring and insulin dose adjustment, injection technique and site  
13 management, and the ability to use such self-management skills on a day-to-  
14 day basis life-long. Fear of hypoglycaemia and of weight gain are major  
15 barriers to success, as is fitting diabetes self-management into busy lifestyles.  
16 Everyone struggles to meet optimised targets and some are more successful  
17 in achieving them than others. Research into new interventions ranging from  
18 more effective education and support, through improved technologies in terms  
19 of insulin replacement and glucose monitoring, and including use of cell-based  
20 therapies, is urgently needed. It is also important to ensure that adults with  
21 type 1 diabetes are able to engage with such methodologies.

22 **2.2 *Continuous glucose monitoring for adults with type 1***  
23 ***diabetes***

24 In adults with type 1 diabetes who have chronically poor control of blood  
25 glucose levels, what is the clinical and cost effectiveness of continuous  
26 glucose monitoring technologies?

27 **Why this is important**

28 Current CGM systems were found not to be cost-effective in our de-novo  
29 analysis, even in patients who had impaired awareness of hypoglycaemia. In  
30 patients who are poorly controlled, there still may be some value for using

1 CGM systems, and therefore further research is needed to determine whether  
2 newer technologies would prove to be cost-effective, particularly in these  
3 patients..

### 4 **2.3        *Structured education programmes for adults with*** 5 ***type 1 diabetes***

6 In adults with type 1 diabetes, what methods can be used to increase the  
7 uptake of structured education programmes and to improve their clinical  
8 outcomes (particularly achieving and sustaining blood glucose control  
9 targets)?

#### 10 **Why this is important**

11 Structured education programmes in flexible insulin therapy have been shown  
12 to improve diabetes control (lower HbA1c and less hypoglycaemia), but  
13 achieving and sustaining optimal diabetes control for avoidance of  
14 complications remains challenging. Some people fail to achieve ideal targets  
15 for blood glucose control, others achieve but cannot maintain them, and still  
16 others do not access structured education at all. There is therefore a need to  
17 develop and test (1) more effective ways of engaging adults with type 1  
18 diabetes in education; (2) improvements in the delivery of education to  
19 increase the number of people achieving targets for diabetic control and (3)  
20 enhanced support for adults with type 1 diabetes to sustain good diabetic  
21 control over time. If the uptake and delivery of clinically and cost effective  
22 education and support for adults with type 1 diabetes can be improved, it  
23 should be possible to achieve a reduction in the short-term and long-term  
24 complications of the condition.

### 25 **2.4        *Risk stratification tool for HbA1c targets for adults*** 26 ***with type 1 diabetes***

27 Can a risk stratification tool be used to aid the setting of individualised HbA1c  
28 targets for adults with type 1 diabetes?

#### 29 **Why this is important**

1 Strict blood glucose control early in the history of type 1 diabetes has been  
2 shown to reduce the development and progression of long-term  
3 complications, but it is not possible to determine who is at particular risk of  
4 glucose-driven poor outcomes. Furthermore, there is a dearth of evidence of  
5 the risk:benefit ratio of strict blood glucose control in people who already have  
6 diabetes complications. Since achieving and maintaining near-normal blood  
7 glucose concentrations is complicated, a risk stratification tool to calculate the  
8 modifiable individual risk of complications will allow blood glucose targets to  
9 be tailored for each person and appropriate support to be provided.

## 10 **2.5 Technologies for preventing and treating impaired** 11 **awareness of hypoglycaemia in adults with type 1** 12 **diabetes**

13 For adults with type 1 diabetes, what are the optimum technologies (such as  
14 insulin pump therapy and/or continuous glucose monitoring, partially or fully  
15 automated insulin delivery, and behavioural, psychological and educational  
16 interventions) and how are they best used, in terms of clinical and cost  
17 effectiveness, for preventing and treating impaired awareness of  
18 hypoglycaemia?

### 19 **Why this is important**

20 Impaired awareness of hypoglycaemia renders adults with type 1 diabetes  
21 susceptible to sudden unexpected deteriorations of conscious level and  
22 irrational behaviour, and increases their risk of severe hypoglycaemia 6-fold.  
23 Impaired awareness of hypoglycaemia and severe hypoglycaemia creates  
24 barriers to many aspects of daily living, and can cause enormous stress for  
25 family and friends. Severe hypoglycaemia can also cause fear of  
26 hypoglycaemia great enough to prevent a person achieving the glucose  
27 targets that are associated with minimal risk of complications. Impaired  
28 awareness of hypoglycaemia results from overexposure to hypoglycaemia in  
29 daily life, and awareness can be much improved by avoidance of  
30 hypoglycaemia. Developing technologies in glucose monitoring and insulin  
31 delivery have not been rigorously tested in adults with type 1 diabetes and  
32 impaired awareness of hypoglycaemia. Research is needed formally to

1 document the extent to which existing technologies can help the adult with  
2 type 1 diabetes and impaired awareness of hypoglycaemia to avoid  
3 hypoglycaemic episodes and regain awareness for occasional episodes.  
4 Research is also needed to develop new technologies. Research is also  
5 needed into how to engage adults with type 1 diabetes and impaired  
6 awareness of hypoglycaemia with treatment strategies designed to improve  
7 awareness.

## 8 **3 Other information**

### 9 **3.1 Scope and how this guideline was developed**

10 NICE guidelines are developed in accordance with a [scope](#) that defines what  
11 the guideline will and will not cover.

#### **How this guideline was developed**

NICE commissioned the National Clinical Guideline Centre to develop this guideline. The Centre established a Guideline Development Group (see section 4), which reviewed the evidence and developed the recommendations.

The methods and processes for developing NICE clinical guidelines are described in [The guidelines manual](#).

12

### 13 **3.2 Related NICE guidance**

14 Details are correct at the time of consultation on the guideline (December  
15 2014). Further information is available on [the NICE website](#).

#### 16 **Published**

##### 17 **General**

- 18 • [Patient experience in adult NHS services](#) (2012) NICE guideline CG138
- 19 • [Medicines adherence](#) (2009) NICE guideline CG76

- 1 • [Four commonly used methods to increase physical activity](#) (2006) NICE  
2 guideline PH2
- 3 • [Nutrition support in adults](#) (2006) NICE guideline CG32
- 4 • [Brief interventions and referral for smoking cessation](#) (2006) NICE  
5 guideline PH1

6 ***Condition-specific***

- 7 • [Obesity](#) (2014) NICE guideline CG189
- 8 • [Chronic kidney disease](#) (2014) NICE guideline CG182
- 9 • [Gastroelectrical stimulation for gastroparesis](#) (2014) NICE interventional  
10 procedures guidance 489
- 11 • [Lipid modification](#) (2014) NICE guideline CG181
- 12 • [Neuropathic pain – pharmacological management](#) (2013) NICE guideline  
13 CG173
- 14 • [Fluocinolone acetonide intravitreal implant for treating chronic diabetic  
15 macular oedema after an inadequate response to prior therapy](#) (2013)  
16 NICE technology appraisal guidance 301
- 17 • [Lower limb peripheral arterial disease](#) (2012) NICE guideline CG147
- 18 • [Ranibizumab for the treatment of diabetic macular oedema](#) (2011) NICE  
19 technology appraisal guidance 237
- 20 • [Hyperglycaemia in acute coronary syndromes](#) (2011) NICE guideline  
21 CG130
- 22 • [Hypertension](#) (2011) NICE guideline CG127
- 23 • [Dexamethasone intravitreal implant for the treatment of macular oedema  
24 secondary to retinal vein occlusion](#) (2011) NICE technology appraisal  
25 guidance 229
- 26 • [Depression in adults](#) (2009) NICE guideline CG90
- 27 • [Depression with a chronic physical health problem](#) (2009) NICE guideline  
28 CG91
- 29 • [Allogeneic pancreatic islet cell transplantation for type 1 diabetes mellitus](#)  
30 (2008) NICE interventional procedure guidance 257
- 31 • [Continuous subcutaneous insulin infusion for the treatment of diabetes  
32 mellitus](#) (2008) NICE technology appraisal guidance 151

1 **Under development**

2 NICE is [developing](#) the following guidance:

- 3 • Diabetes in pregnancy (update). NICE guideline. Publication expected  
4 February 2015.
- 5 • Diabetic foot problems (update). NICE guideline. Publication expected July  
6 2015.
- 7 • Diabetes in children and young people (update). NICE guideline.  
8 Publication expected August 2015.
- 9 • Type 2 diabetes in adults (update). NICE guideline. Publication expected  
10 August 2015.
- 11 • Buccal insulin for managing type 1 diabetes. NICE technology appraisal  
12 guidance. Publication date to be confirmed.

13

1    **4           The Guideline Development Group, National**  
2                   **Collaborating Centre and NICE project team**

3    **4.1           Guideline Development Group**

4    The Guideline Development Group members listed are those for the 2015  
5    update. For the composition of (the) previous Guideline Development  
6    Group(s), see the full guideline.

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11 **Lyn Knott**

12 Editor

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14

15

1 **Appendix A: Recommendations from NICE guideline**  
 2 **CG15 (2004) that have been deleted or changed**

3 ***Recommendations to be deleted***

4 The table shows recommendations from 2004 that NICE proposes deleting in  
 5 the 2015 update. The right-hand column gives the replacement  
 6 recommendation, or explains the reason for the deletion if there is no  
 7 replacement recommendation.

Recommendation in 2004 guideline	Comment
Diabetes should be confirmed by a single diagnostic laboratory glucose measurement in the presence of classical symptoms, or by further laboratory glucose measurement. The diagnosis may be supported by a raised HbA1c. [1.6.1.1]	This recommendation has been deleted because it is not appropriate for type 1 diabetes. It refers to blood tests for which you need to wait for the result, and therefore implies that it is acceptable to wait for HbA1c or laboratory glucose results. This poses a risk to patients as treatment is delayed.
Where diabetes is diagnosed, but type 2 diabetes is suspected, the diagnosis of type 1 diabetes should be considered if: <ul style="list-style-type: none"> <li>• ketonuria is detected, or</li> <li>• weight loss is marked, or</li> <li>• the person does not have features of the metabolic syndrome or other contributing illness. [1.6.1.2]</li> </ul>	Replaced by: Diagnose type 1 diabetes on clinical grounds in adults presenting with hyperglycaemia, bearing in mind that people with type 1 diabetes typically (but not always) have one or more of: <ul style="list-style-type: none"> <li>• ketosis</li> <li>• rapid weight loss</li> <li>• age of onset below 50 years</li> <li>• BMI below 25 kg/m<sup>2</sup></li> <li>• personal and/or family history of autoimmune disease. <b>[new 2015]</b> [1.1.1]</li> </ul>
When diabetes is diagnosed in a younger person, the possibility that the diabetes is not type 1 diabetes should be considered if they are obese or have a family history of diabetes, particularly if they are of non-white ethnicity. [1.6.1.3]	This recommendation has been deleted because retaining it would be dangerous. Readers are more likely to miss the diagnosis of type 1 diagnosis in a person who is black, has a family history or is obese. This recommendation is more relevant to type 2 diabetes.
Tests to detect specific auto-antibodies or to measure C-peptide deficiency should not be regularly used to confirm the diagnosis of type 1 diabetes. Their use should be considered if predicting the rate of decline of islet B-cell function	Replaced by: Do not measure C-peptide and/or diabetes-specific autoantibody titres routinely to confirm type 1 diabetes in adults. <b>[new 2015]</b> [1.1.3]

<p>would be useful in discriminating type 1 from type 2 diabetes.[1.6.1.4]</p>	
<p>Conventional technology (telephones), or newer technologies for high-density data transmission of images, should be used to improve process and outcomes. [1.7.1.6]</p>	<p>This recommendation has been deleted. It is outdated and no longer valid.</p>
<p>A programme of structured diabetes education covering all major aspects of diabetes self-care and the reasons for it should be made available to all adults with type 1 diabetes in the months after diagnosis, and periodically thereafter according to agreed need following yearly assessment. [1.8.1.1]</p>	<p>Replaced by: Offer all adults with type 1 diabetes a structured education programme of proven benefit, for example the DAFNE (dose-adjustment for normal eating) programme. Offer this programme 6–12 months after diagnosis, at a time that is clinically appropriate and suitable for the person. [1.3.1]</p>
<p>Education programmes for adults with type 1 diabetes should be flexible so that they can be adapted to specific educational, social and cultural needs. These needs should be integrated with individual health needs as dictated by the impact of diabetes and other relevant health conditions on the individual.[1.8.1.2]</p>	<p>Replaced by: Ensure that any structured education programme for adults with type 1 diabetes includes the following components:</p> <ul style="list-style-type: none"> <li>• It is evidence-based, and suits the needs of the person.</li> <li>• It has specific aims and learning objectives, and supports the person and their family members and carers in developing attitudes, beliefs, knowledge and skills to self-manage diabetes.</li> <li>• It has a structured curriculum that is theory-driven, evidence-based and resource-effective, has supporting materials, and is written down.</li> <li>• It is delivered by trained educators who have an understanding of educational theory appropriate to the age and needs of the person, and who are trained and competent to deliver the principles and content of the programme.</li> <li>• It is quality assured, and reviewed by trained, competent, independent assessors who measure it against criteria that ensure consistency.</li> <li>• The outcomes are audited regularly. [1.3.3]</li> </ul>
<p>Education programmes for adults with type 1 diabetes should be designed and delivered by members of the multidisciplinary diabetes team in accordance with the principles of adult</p>	<p>Replaced by: Ensure that any structured education programme for adults with type 1 diabetes includes the following components:</p>

<p>education. [1.8.1.3]</p>	<ul style="list-style-type: none"> <li>• It is evidence-based, and suits the needs of the person.</li> <li>• It has specific aims and learning objectives, and supports the person and their family members and carers in developing attitudes, beliefs, knowledge and skills to self-manage diabetes.</li> <li>• It has a structured curriculum that is theory-driven, evidence-based and resource-effective, has supporting materials, and is written down.</li> <li>• It is delivered by trained educators who have an understanding of educational theory appropriate to the age and needs of the person, and who are trained and competent to deliver the principles and content of the programme.</li> <li>• It is quality assured, and reviewed by trained, competent, independent assessors who measure it against criteria that ensure consistency.</li> <li>• The outcomes are audited regularly. [1.3.3]</li> </ul>
<p>Education programmes for adults with type 1 diabetes should include modules designed to empower adults to participate in their own healthcare through:</p> <ul style="list-style-type: none"> <li>• enabling them to make judgements and choices about how they effect that care</li> <li>• obtaining appropriate input from the professionals available to advise them. [1.8.1.4]</li> </ul>	<p>Replaced by:</p> <p>Ensure that any structured education programme for adults with type 1 diabetes includes the following components:</p> <ul style="list-style-type: none"> <li>• It is evidence-based, and suits the needs of the person.</li> <li>• It has specific aims and learning objectives, and supports the person and their family members and carers in developing attitudes, beliefs, knowledge and skills to self-manage diabetes.</li> <li>• It has a structured curriculum that is theory-driven, evidence-based and resource-effective, has supporting materials, and is written down.</li> <li>• It is delivered by trained educators who have an understanding of educational theory appropriate to the age and needs of the person, and who are trained and competent to deliver the principles and content of the programme.</li> <li>• It is quality assured, and reviewed by trained, competent, independent</li> </ul>

	<p>assessors who measure it against criteria that ensure consistency.</p> <ul style="list-style-type: none"> <li>The outcomes are audited regularly. [1.3.3]</li> </ul>
<p>Professionals engaged in the delivery of diabetes care should consider incorporating educational interchange at all opportunities when in contact with a person with type 1 diabetes. The professional should have the skills and training to make best use of such time.[1.8.1.5]</p>	<p>Replaced by:</p> <p>Provide information about type 1 diabetes and its management to adults with type 1 diabetes at all opportunities from diagnosis onwards. Follow the principles in the NICE guideline on <a href="#">patient experience in adult NHS services</a>. [1.3.5]</p>
<p>Self-monitoring of blood glucose levels should be used as part of an integrated package that includes appropriate insulin regimens and education to help choice and achievement of optimal diabetes outcomes. [1.8.2.1]</p>	<p>Replaced by:</p> <p>Educate adults with type 1 diabetes about how to measure their blood glucose level, interpret the results and know what action to take. [1.6.17]</p> <p>Support adults with type 1 diabetes to make the best use of data from self-monitoring of blood glucose through structured education (see recommendations 1.3.1 and 1.3.2). [1.6.18]</p>
<p>Self-monitoring results should be interpreted in light of clinically significant life events. [1.8.2.3]</p>	<p>Replaced by:</p> <p>Support adults with type 1 diabetes to make the best use of data from self-monitoring of blood glucose through structured education (see recommendations 1.3.1 and 1.3.2). [1.6.18]</p>
<p>Self-monitoring should be performed using meters and strips chosen by adults with diabetes to suit their needs, and usually with low blood requirements, fast analysis times and integral memories.[1.8.2.4]</p>	<p>This recommendation has been deleted because it is no longer relevant. Technology for blood glucose meters has advanced since 2004. All blood glucose meters have integrated memories and fast analysis times.</p>
<p>Adults with type 1 diabetes should be advised that the optimal frequency of self monitoring will depend on:</p> <ul style="list-style-type: none"> <li>the characteristics of an individual's blood glucose control</li> <li>the insulin treatment regimen</li> <li>personal preference in using the results to achieve the desired lifestyle. [1.8.2.6]</li> </ul>	<p>Replaced by:</p> <p>Advise routine self-monitoring of blood glucose levels for all adults with type 1 diabetes, and recommend testing at least 4 times a day, including before each meal and before bed. [1.6.12]</p>
<p>Adults with type 1 diabetes should be advised that the optimal targets for short-</p>	<p>Replaced by:</p> <p>Advise adults with type 1 diabetes to aim</p>

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<p>term glycaemic control are:</p> <ul style="list-style-type: none"> <li>• a pre-prandial blood glucose level of 4.0–7.0 mmol/litre and</li> <li>• a post-prandial blood glucose level of less than 9.0 mmol/litre. [1.8.2.7]</li> </ul>	<p>for:</p> <ul style="list-style-type: none"> <li>• a fasting plasma glucose level of 5–7 mmol/litre on waking and</li> <li>• a plasma glucose level of 4–7 mmol/litre before meals at other times of the day. [1.6.15]</li> </ul>
<p>Clinical monitoring of blood glucose levels by high-precision DCCT-aligned methods of haemoglobin A1c (HbA1c) should be performed every 2-6 months, depending on:</p> <ul style="list-style-type: none"> <li>• achieved level of blood glucose control</li> <li>• stability of blood glucose control</li> <li>• change in insulin dose or regimen. [1.9.1.1]</li> </ul>	<p>Replaced by:</p> <p>Measure HbA1c levels every 3–6 months in adults with type 1 diabetes. [1.6.1]</p>
<p>Site-of-care measurement, or before-clinical-consultation measurement, should be provided. [1.9.1.2]</p>	<p>Replaced by:</p> <p>Inform adults with type 1 diabetes of their HbA1c results after each measurement and ensure that their most recent result is available at the time of consultation. Follow the principles in the NICE guideline on patient experience in adult NHS services about communication. [1.6.4]</p>
<p>HbA1c results should be communicated to the person with type 1 diabetes after each measurement. The term A1c can be used for simplicity.</p>	<p>Replaced by:</p> <p>Inform adults with type 1 diabetes of their HbA1c results after each measurement and ensure that their most recent result is available at the time of consultation. Follow the principles in the NICE guideline on patient experience in adult NHS services about communication. [1.6.4]</p>
<p>Fructosamine should not be used as a routine substitute for HbA1c estimation. [1.9.1.5]</p>	<p>This recommendation has been deleted because it is redundant. Clinical practice has changed. Now it is clinical practice to use HbA1c, whereas in 2004 there was a mixture of both methods.</p>
<p>Continuous glucose monitoring systems have a role in the assessment of glucose profiles in adults with consistent glucose control problems on insulin therapy, notably:</p> <ul style="list-style-type: none"> <li>• repeated hyper- or hypoglycaemia at the same time of day</li> <li>• hypoglycaemia unawareness, unresponsive to conventional insulin and dose adjustment. [1.9.1.6]</li> </ul>	<p>Replaced by:</p> <p>Consider real-time continuous glucose monitoring for adults with type 1 diabetes who are willing to commit to using it at least 70% of the time and to calibrate it as needed, and who have any of the following that persist despite optimised use of insulin therapy and conventional blood glucose monitoring:</p> <ul style="list-style-type: none"> <li>• more than 1 episode a year of severe hypoglycaemia with no obviously</li> </ul>

	<p>preventable precipitating cause</p> <ul style="list-style-type: none"> <li>• complete loss of awareness of hypoglycaemia</li> <li>• frequent (more than 2 episodes a week) asymptomatic hypoglycaemia that is causing problems with daily activities</li> <li>• extreme fear of hypoglycaemia. [1.6.21]</li> </ul>
<p>Adults with type 1 diabetes should be advised that maintaining a DCCT harmonised HbA1c below 7.5% is likely to minimise their risk of developing diabetic eye, kidney or nerve damage in the longer term.[1.9.2.1]</p>	<p>Replaced by:</p> <p>Support adults with type 1 diabetes to achieve and maintain a target HbA1c level of 48 mmol/mol (6.5%) or lower, to minimise the risk of long-term vascular complications. [1.6.6]</p>
<p>Adults with diabetes who want to achieve an HbA1c down to, or towards, 7.5% should be given all appropriate support in their efforts to do so.[1.9.2.2]</p>	<p>Replaced by:</p> <p>Support adults with type 1 diabetes to achieve and maintain a target HbA1c level of 48 mmol/mol (6.5%) or lower, to minimise the risk of long-term vascular complications. [1.6.6]</p>
<p>Where there is evidence of increased arterial risk (identified by a raised albumin excretion rate, features of the metabolic syndrome, or other arterial risk factors), people with type 1 diabetes should be advised that approaching lower HbA1c levels (for example, 6.5% or lower) may be of benefit to them. Support should be given to approaching this target if so wished.[1.9.2.3]</p>	<p>Replaced by:</p> <p>Support adults with type 1 diabetes to achieve and maintain a target HbA1c level of 48 mmol/mol (6.5%) or lower, to minimise the risk of long-term vascular complications. [1.6.6]</p> <p>Agree an individualised HbA1c target with each adult with type 1 diabetes, taking into account factors such as the person's daily activities, aspirations, likelihood of complications, comorbidities, occupation and history of hypoglycaemia.[1.6.7]</p>
<p>Undetected hypoglycaemia and an attendant risk of unexpected disabling hypoglycaemia or of hypoglycaemia unawareness should be suspected in adults with type 1 diabetes who have:</p> <ul style="list-style-type: none"> <li>• lower HbA1c levels, in particular levels in or approaching the normal reference range (DCCT harmonised &lt; 6.1%)</li> <li>• HbA1c levels lower than expected from self-monitoring results.[1.9.2.5]</li> </ul>	<p>Replaced by:</p> <p>Assess awareness of hypoglycaemia in adults with type 1 diabetes at each annual review. [1.10.1]</p> <p>Review insulin regimens and doses and prioritise strategies to avoid hypoglycaemia in adults with type 1 diabetes with impaired awareness of hypoglycaemia, including:</p> <ul style="list-style-type: none"> <li>• reinforcing the principles of structured education</li> <li>• offering continuous subcutaneous insulin infusion (CSII or insulin pump) therapy</li> </ul>

	<ul style="list-style-type: none"> <li>• offering real-time continuous glucose monitoring. [1.10.7]</li> </ul> <p>If impaired awareness of hypoglycaemia is associated with recurrent severe hypoglycaemia despite these interventions, consider referring the person to a specialist centre. [1.10.8]</p>
<p>Where experience or risk of hypoglycaemia is significant to an individual, or the effort needed to achieve target levels severely curtails other quality of life despite optimal use of current diabetes technologies, tighter blood glucose control should not be pursued without balanced discussion of the advantages and disadvantages.[1.9.2.6]</p>	<p>Replaced by:</p> <p>Ensure that adults with type 1 diabetes with impaired awareness of hypoglycaemia have had structured education in flexible insulin therapy using basal–bolus regimens and are following its principles correctly. [1.10.4]</p> <p>Offer additional education focusing on avoiding and treating hypoglycaemia to adults with type 1 diabetes who continue to have impaired awareness of hypoglycaemia after structured education in flexible insulin therapy. [1.10.5]</p> <p>Avoid relaxing individualised blood glucose targets as a treatment for adults with type 1 diabetes with impaired awareness of hypoglycaemia. [1.10.6]</p> <p>Review insulin regimens and doses and prioritise strategies to avoid hypoglycaemia in adults with type 1 diabetes with impaired awareness of hypoglycaemia, including:</p> <ul style="list-style-type: none"> <li>• reinforcing the principles of structured education</li> <li>• offering continuous subcutaneous insulin infusion (CSII or insulin pump) therapy</li> <li>• offering real-time continuous glucose monitoring. [1.10.7]</li> </ul> <p>If impaired awareness of hypoglycaemia is associated with recurrent severe hypoglycaemia despite these interventions, consider referring the person to a specialist centre. [1.10.8]</p>
<p>Adults with type 1 diabetes should have access to the types (preparation and species) of insulin they find allow them</p>	<p>This recommendation has been removed because it has no clear meaning.</p>

optimal well-being.[1.9.3.1]	
Multiple insulin injection regimens, in adults who prefer them, should be used as part of an integrated package of which education, food and skills training should be integral parts.[1.9.3.3]	Replaced by: Offer multiple daily injection basal–bolus insulin regimens, rather than twice-daily mixed insulin regimens, as the insulin injection regimen of choice for all adults with type 1 diabetes. [1.7.2]
Appropriate self-monitoring and education should be used as part of an integrated package to help achieve optimal diabetes outcomes. [1.9.3.4]	Replaced by: Support adults with type 1 diabetes to make the best use of data from self-monitoring of blood glucose through structured education (see recommendations 1.3.1. and 1.3.2). [1.6.18]
Meal-time insulin injections should be provided by injection of unmodified ('soluble') insulin or rapid-acting insulin analogues before main meals. [1.9.3.5]	Replaced by: Offer rapid-acting insulin analogues injected before meals, rather than rapid-acting soluble human or animal insulins, for mealtime insulin replacement for adults with type 1 diabetes. [1.7.7]
Rapid-acting insulin analogues should be used as an alternative to meal-time unmodified insulin: <ul style="list-style-type: none"> <li>• where nocturnal or late inter-prandial hypoglycaemia is a problem</li> <li>• in those whom they allow equivalent blood glucose control without use of snacks between meals and this is needed or desired. [1.9.3.6]</li> </ul>	Replaced by: If an adult with type 1 diabetes has a strong preference for an alternative mealtime insulin, respect their wishes and offer the preferred insulin. [1.7.9]
Basal insulin supply (including nocturnal insulin supply) should be provided by the use of isophane (NPH) insulin or long-acting insulin analogues (insulin glargine). Isophane (NPH) insulin should be given at bedtime. If rapid-acting insulin analogues are given at meal times or the midday insulin dose is small If rapid-acting insulin analogues are given at meal times or the midday insulin dose is small or lacking, the need to give isophane (NPH) insulin twice daily (or more often) should be considered. [1.9.3.7]	Replaced by: Offer twice-daily insulin detemir as basal insulin therapy for adults with type 1 diabetes. [1.7.4]
Long-acting insulin analogues (insulin glargine) should be used when: <ul style="list-style-type: none"> <li>• nocturnal hypoglycaemia is a problem on isophane (NPH) insulin</li> <li>• morning hyperglycaemia on isophane (NPH) insulin results in difficult</li> </ul>	Replaced by: Consider, as an alternative basal insulin therapy for adults with type 1 diabetes: <ul style="list-style-type: none"> <li>• an existing insulin regimen being used by the person that is achieving their agreed targets</li> </ul>

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<p>daytime blood</p> <ul style="list-style-type: none"> <li>• glucose control</li> <li>• rapid-acting insulin analogues are used for meal-time blood glucose control.[1.9.3.8]</li> </ul>	<ul style="list-style-type: none"> <li>• once-daily insulin glargine if insulin detemir is not tolerated or if twice-daily basal insulin injection is not acceptable to the person. [1.7.5]</li> </ul>
<p>Twice-daily insulin regimens should be used by those adults who consider number of daily injections an important issue in quality of life.</p> <ul style="list-style-type: none"> <li>• Biphasic insulin preparations (pre-mixes) are often the preparations of choice in this circumstance.</li> <li>• Biphasic rapid-acting insulin analogue pre-mixes may give an advantage to those prone to hypoglycaemia at night.</li> </ul> <p>Such twice daily regimens may also help:</p> <ul style="list-style-type: none"> <li>• those who find adherence to their agreed lunch-time insulin injection difficult</li> <li>• adults with learning difficulties who may require assistance from others.[1.9.3.9]</li> </ul>	<p>Replaced by:</p> <p>Consider a twice-daily human mixed insulin regimen for adults with type 1 diabetes if a multiple daily injection basal–bolus insulin regimen is not possible and a twice-daily mixed insulin regimen is chosen. [1.7.10]</p>
<p>Adults whose nutritional and physical activity patterns vary considerably from day to day, for vocational or recreational reasons, may need careful and detailed review of their self-monitoring and insulin injection regimen(s). This should include all the appropriate preparations (see sections 1.9.3.6–8), and consideration of unusual patterns and combinations.</p>	<p>This recommendation has been deleted because the content would be covered by structured education programmes.</p>
<p>For adults undergoing periods of fasting or sleep following eating (such as during religious feasts and fasts or after night-shift work), a rapid-acting insulin analogue before the meal (provided the meal is not prolonged) should be considered. [1.9.3.11]</p>	<p>This recommendation has been deleted because a good basal-bolus regimen, as recommended in 1.7.2, should be able to accommodate a period of fasting and feasting.</p>
<p>Continuous subcutaneous insulin infusion (or insulin pump therapy) is recommended as an option for people with type 1 diabetes provided that:</p> <ul style="list-style-type: none"> <li>• multiple-dose insulin therapy (including, where appropriate, the use of insulin</li> <li>• glargine) has failed; and</li> <li>• those receiving the treatment have the commitment and competence to use the</li> </ul>	<p>Replaced by:</p> <p>For guidance on the use of continuous subcutaneous insulin infusion (CSII or insulin pump) therapy for adults with type 1 diabetes, see <a href="#">Continuous subcutaneous insulin infusion for the treatment of diabetes mellitus</a> (NICE technology appraisal guidance 151). [1.7.6]</p>

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<ul style="list-style-type: none"> <li>therapy effectively. [1.9.3.13]</li> </ul>	
<p>Partial insulin replacement to achieve blood glucose control targets (basal insulin only, or just some meal-time insulin) should be considered for adults starting insulin therapy, until such time as islet B-cell deficiency progresses further. [1.9.3.14]</p>	<p>Replaced by:</p> <p>Do not offer non-basal–bolus insulin regimens for treating adults newly diagnosed with type 1 diabetes. [1.7.3]</p>
<p>Oral glucose-lowering drugs should generally not be used in the management of adults with type 1 diabetes. [1.9.3.16]</p>	<p>Replaced by:</p> <p>Consider adding metformin to insulin therapy if an adult with type 1 diabetes and a BMI of 25 kg/m<sup>2</sup> or above wants to improve their blood glucose control while minimising their effective insulin dose. [1.7.14]</p>
<p>Insulin injection should be made into the deep subcutaneous fat. To achieve this, needles of a length appropriate to the individual should be made available.[1.9.4.3]</p>	<p>Replaced by:</p> <p>Offer needles of different lengths to adults with type 1 diabetes who are having problems such as pain, local skin reactions and injection site leakages. [1.8.3]</p>
<p>Adults with type 1 diabetes should be informed that the abdominal wall is the therapeutic choice for meal-time insulin injections.[1.9.4.4]</p>	<p>Replaced by:</p> <p>Advise adults with type 1 diabetes to rotate insulin injection sites and avoid repeated injections at the same point within sites. [1.8.5]</p>
<p>Adults with type 1 diabetes should be informed that extended-acting suspension insulin, for example isophane (NPH) insulin, may give a longer profile of action when injected into the subcutaneous tissue of the thigh rather than the arm or abdominal wall.[1.9.4.5]</p>	<p>This recommendation has been deleted because isophane (NPH) insulin is no longer recommended as first line choice for long-acting insulin.</p>
<p>Adults with diabetes should be recommended to use one anatomical area for the injections given at the same time of day, but to move the precise injection site around in the whole of the available skin within that area.[1.9.4.6]</p>	<p>Replaced by:</p> <p>Advise adults with type 1 diabetes to rotate insulin injection sites and avoid repeated injections at the same point within sites. [1.8.5]</p>
<p>When a more rapid-acting form of glucose is required, purer glucose containing solutions should be given. [1.9.5.3]</p>	<p>This recommendation has been deleted because it is a duplication of recommendation 1.10.9.</p>
<p>Hypoglycaemia unawareness should be assumed to be secondary to undetected periods of hypoglycaemia (&lt; 3.5 mmol/l, often for extended periods, commonly at night) until these are excluded by appropriate monitoring techniques. If present, such periods of hypoglycaemia</p>	<p>This recommendation has been deleted because it is not really a recommendation and does not give clear advice. The definition of hypoglycaemia unawareness is included in the linking evidence to recommendations table for identification of impaired awareness of</p>

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should be ameliorated.[1.9.5.6]	hypoglycaemia and the glossary.
Specific education on the detection and management of hypoglycaemia in adults with problems of hypoglycaemia awareness should be offered. [1.9.5.7]	This recommendation has been deleted because it is redundant.
Aspirin therapy (75mg daily) should be recommended in adults in the highest and moderately-high-risk categories. [1.10.2.3]	Replaced by: Do not offer aspirin for the primary prevention of cardiovascular disease to adults with type 1 diabetes. [1.13.1]
If an abnormal surveillance result is obtained (in the absence of proteinuria / urinary tract infection) the test should be repeated at each clinic visit or at least every 3 - 4 months, and the result taken as confirmed if a further specimen (out of two more) is also abnormal (>2.5 mg/mmol for men, > 3.5 mg/mmol for women). [1.11.2.2]	For guidance on managing kidney disease in adults with type 1 diabetes, see the NICE guideline on <a href="#">chronic kidney disease</a> . [1.15.15]
Men should be asked annually whether erectile dysfunction is an issue. [1.11.4.1]	Replaced by: Offer men with type 1 diabetes the opportunity to discuss erectile dysfunction as part of regular review. [1.15.30]
A PDE5 (phosphodiesterase-5) inhibitor drug, if not contraindicated, should be offered where erectile dysfunction is a problem. [1.11.4.2]	Replaced by: Offer a phosphodiesterase-5 inhibitor to men with type 1 diabetes with isolated erectile dysfunction unless contraindicated. Choose the phosphodiesterase-5 inhibitor with the lowest acquisition cost. [1.15.31]
In adults with type 1 diabetes on insulin therapy who have erratic blood glucose control or unexplained bloating or vomiting, the diagnosis of gastroparesis should be considered. [1.11.4.4]	Replaced by: Consider domperidone <sup>8</sup> (in preference to metoclopramide) for treating gastroparesis <sup>9</sup> in adults with type 1 diabetes. [1.15.1] Consider continuous subcutaneous insulin infusion (CSII or insulin pump) therapy for adults with type 1 diabetes who have gastroparesis. [1.15.2] Advise a small-particle-size diet (mashed or pureed food) for symptomatic relief for adults with type 1 diabetes who have vomiting caused by gastroparesis. [1.5.3]

<sup>8</sup> Although this use is common in UK clinical practice, at the time of consultation (December 2014), domperidone did not have a UK marketing authorisation for this indication. The prescriber should follow relevant professional guidance, taking full responsibility for the decision. Informed consent should be obtained and documented. See the General Medical Council's [Prescribing guidance: prescribing unlicensed medicines](#) for further information.

<sup>9</sup> Diagnosis of gastroparesis needing specific therapy can only be made in the absence of hyperglycaemia at the time of testing, because hyperglycaemia induces a physiological delay in gastric emptying.

	Refer adults with type 1 diabetes who have gastroparesis for specialist advice if the interventions in recommendations 1.15.1, 1.15.2 and 1.15.3 are not beneficial or not appropriate. [1.5.4]
In adults with diabetes who have altered perception of hypoglycaemia, the possibility of sympathetic nervous system damage as a contributory factor should be considered.[1.11.4.5]	Replaced by: Assess awareness of hypoglycaemia in adults with type 1 diabetes at each annual review. [1.10.1] Use the Gold score or Clarke score to quantify awareness of hypoglycaemia in adults with type 1 diabetes, checking that the questionnaire items have been answered correctly. [1.10.2] Explain to adults with type 1 diabetes that impaired awareness of the symptoms of plasma glucose levels below 3 mmol/litre is associated with a significantly increased risk of severe hypoglycaemia. [1.10.3]
For adults with diabetes with diagnosed or suspected gastroparesis, a trial of prokinetic drugs is indicated (metoclopramide or domperidone, with cisapride as third line if necessary). [1.11.4.10]	Replaced by: Consider domperidone <sup>10</sup> (in preference to metoclopramide) for treating gastroparesis <sup>11</sup> in adults with type 1 diabetes. [1.15.1] Consider continuous subcutaneous insulin infusion (CSII or insulin pump) therapy for adults with type 1 diabetes who have gastroparesis. [1.15.2] Advise a small-particle-size diet (mashed or pureed food) for symptomatic relief for adults with type 1 diabetes who have vomiting caused by gastroparesis [1.15.3] Refer adults with type 1 diabetes who have gastroparesis for specialist advice if the interventions in recommendations 1.15.1, 1.15.2 and 1.15.3 are not beneficial or not appropriate. [1.15.4]
Professionals should be alert to the psychological consequences of chronic painful neuropathy, and offer appropriate	Replaced by: For guidance on treating chronic diabetic neuropathy, see the NICE guideline on

<sup>10</sup> Although this use is common in UK clinical practice, at the time of consultation (December 2014), domperidone did not have a UK marketing authorisation for this indication. The prescriber should follow relevant professional guidance, taking full responsibility for the decision. Informed consent should be obtained and documented. See the General Medical Council's [Prescribing guidance: prescribing unlicensed medicines](#) for further information.

<sup>11</sup> Diagnosis of gastroparesis needing specific therapy can only be made in the absence of hyperglycaemia at the time of testing, because hyperglycaemia induces a physiological delay in gastric emptying.

management where they are identified. [1.11.5.7]	<a href="#">neuropathic pain – pharmacological management</a> . [1.15.38]
Where neuropathic symptoms cannot be adequately controlled, it is useful, to help individuals cope , to explain the: <ul style="list-style-type: none"> <li>• reasons for the problem</li> <li>• likelihood of remission in the medium term</li> <li>• role of improved glucose control. [1.11.5.8]</li> </ul>	Replaced by: For guidance on treating chronic diabetic neuropathy, see the NICE guideline on <a href="#">neuropathic pain – pharmacological management</a> . [1.15.38]
Hospitals should ensure the existence and deployment of an approved protocol for inpatient procedures and surgical operations for adults with Type 1 diabetes. This should aim to ensure the maintenance of near-normoglycaemia without risk of acute decompensation, usually by the use of regular quality-assured blood glucose testing driving the adjustment of intravenous insulin delivery. [1.12.3.4]	Replaced by: Aim for a target plasma glucose level of 5–8 mmol/litre for adults with type 1 diabetes during surgery or acute illness. [1.14.1] Establish a local protocol for controlling blood glucose levels in adults with type 1 diabetes during surgery or acute illness to achieve the target level. [1.14.2]
Special management techniques or treatment for non-severe psychological illness should not be commonly used, except where diabetes-related arterial complications give rise to special precautions over drug therapy. [1.12.5.3]	This recommendation has been deleted because it does not give clear advice; leaving it in would cause confusion.

1

2 ***Amended recommendation wording (change to meaning)***

3 Recommendations are labelled **[2004, amended 2015]** if the evidence has  
4 not been reviewed but either:

- 5 • changes have been made to the recommendation wording (indicated by  
6 highlighted text) that change the meaning **or**  
7 • NICE has made editorial changes to the original wording to clarify the  
8 action to be taken.

9 These changes are marked with yellow highlighting.

<b>Recommendation in 2004 guideline</b>	<b>Recommendation in current guideline</b>	<b>Reason for change</b>
Elements of an individualised and culturally appropriate plan will include: <ul style="list-style-type: none"> <li>• sites and timescales of</li> </ul>	Elements of an individualised and culturally appropriate plan will include:	Additional elements have been included to make this recommendation

<p>diabetes education including nutritional advice (see 'Approach to education', Section 1.8.1, and 'Dietary management', Section 1.8.3)</p> <ul style="list-style-type: none"> <li>• initial treatment modalities (see 'Insulin regimens', Section 1.9.3, and 'Insulin delivery', Section 1.9.4)</li> <li>• means of self-monitoring (see 'Self-monitoring of glucose level', Section 1.8.2)</li> <li>• means and frequency of communication with the professional team</li> <li>• follow-up consultations including surveillance at annual review (see individual late complications recommendations)</li> <li>• management of arterial risk factors (see 'Control of arterial risk', Section 1.10). (1.12.1.2)</li> </ul>	<ul style="list-style-type: none"> <li>• sites and timescales of diabetes education, including nutritional advice (see sections 1.3 and 1.4)</li> <li>• initial treatment modalities, including guidance on insulin injection (see sections 1.7 and 1.8)</li> <li>• means of self-monitoring and targets (see section 1.6)</li> <li>• symptoms, risk and treatment of hypoglycaemia</li> <li>• management of special situations, such as driving</li> <li>• means and frequency of communication with the diabetes professional team</li> <li>• management of arterial risk factors (see section 1.13)</li> <li>• for women of childbearing potential, implications for pregnancy and family planning advice</li> <li>• follow-up consultations, including frequency of review of HbA1c levels and experience of hypoglycaemia, and surveillance at annual review (see section 1.15) <b>[2004, amended 2015]</b> (1.1.7)</li> </ul>	<p>comprehensive.</p>
<p>An individual care plan should be set up and reviewed annually, modified according to changes in wishes, circumstances and medical findings, and the details recorded. The plan should include aspects of:</p> <ul style="list-style-type: none"> <li>• diabetes education including nutritional advice (see 'Approach to education', Section 1.8.1, and 'Dietary management', Section</li> </ul>	<p>Set up an individual care plan jointly agreed with the adult with type 1 diabetes, review it annually and modify it taking into account changes in the person's wishes, circumstances and medical findings, and record the details. The plan should include aspects of:</p> <ul style="list-style-type: none"> <li>• diabetes education, including nutritional advice (see sections 1.3</li> </ul>	<p>The word 'late' has been deleted (with respect to complications) because it implies advanced complications and takes the focus away from prevention. Some crucial aspects of a care plan have been added for completeness as they were not covered in</p>

<p>1.8.3)</p> <ul style="list-style-type: none"> <li>insulin therapy (see 'Insulin regimens', Section 1.9.3, and 'Insulin delivery', Section 1.9.4)</li> <li>self-monitoring (see 'Self-monitoring of glucose', Section 1.8.2)</li> <li>arterial risk factor surveillance and management (see 'Control of arterial risk', Section 1.10)</li> <li>late complications surveillance and management (see 'Identification and management of complications', Section 1.11)</li> <li>means and frequency of communication with the professional care team</li> <li> <ul style="list-style-type: none"> <li>follow-up consultations including next annual review. (1.7.1.4)</li> </ul> </li> </ul>	<p>and 1.4)</p> <ul style="list-style-type: none"> <li>insulin therapy, including dose adjustment (see sections 1.8 and 1.9)</li> <li>self-monitoring (see section 1.6)</li> <li>avoiding hypoglycaemia and maintaining awareness of hypoglycaemia</li> <li>for women of childbearing potential, family planning, contraception and pregnancy planning</li> <li>arterial risk factor surveillance and management (see section 1.13)</li> <li>complications surveillance and management (see section 1.14)</li> <li>means and frequency of communicating with the diabetes professional team</li> <li>follow-up consultations, including frequency of review of HbA1c levels and experience of hypoglycaemia, and next annual review. <b>[2004, amended 2015]</b> (1.2.4)</li> </ul>	<p>the 2004 recommendation.</p>
<p>Information should also be made available on:</p> <ul style="list-style-type: none"> <li>effects of different alcohol-containing drinks on blood glucose excursions and calorie intake</li> <li>use of high-calorie and high-sugar 'treats'</li> <li>use of foods of high glycaemic index. (1.8.3.6)</li> </ul>	<p>Make information available on:</p> <ul style="list-style-type: none"> <li>effects of different alcohol-containing drinks on blood glucose excursions and calorie intake</li> <li>use of high-calorie and high-sugar 'treats'. <b>[2004, amended 2015]</b> (1.4.10)</li> </ul>	<p>There is no evidence of benefit for a low glycaemic index diet (see recommendation 1.4.3), so the reference to giving information about foods of high glycaemic index has been deleted.</p>
<p>Information about the benefits of healthy eating in reducing arterial risk should be made available as part of dietary education in the period after diagnosis, and</p>	<p>Make information available about the benefits of healthy eating in reducing arterial risk as part of dietary education in the period after diagnosis, and according to</p>	<p>There is no evidence of benefit for a low glycaemic index diet (see recommendation 1.4.3), so the reference about giving</p>

<p>according to need and interest at intervals thereafter. This should include information about low glycaemic index foods, fruit and vegetables, and types and amounts of fat, and ways of making the appropriate nutritional changes. (1.8.3.7)</p>	<p>need and interest at intervals thereafter. Include information <b>about fruit</b> and vegetables, types and amounts of fat, and ways of making the appropriate nutritional changes. <b>[2004, amended 2015]</b> (1.4.11)</p>	<p>information about foods of low glycaemic index has been deleted.</p>
<p>All healthcare professionals providing advice on the management of type 1 diabetes should be aware of appropriate nutritional advice on common topics of concern and interest to adults living with type 1 diabetes, and should be prepared to seek advice from colleagues with more specialised knowledge. Suggested common topics include:</p> <ul style="list-style-type: none"> <li>• glycaemic index of specific foods</li> <li>• body weight, energy balance and obesity management</li> <li>• cultural and religious diets, feasts and fasts</li> <li>• foods sold as 'diabetic'</li> <li>• sweeteners</li> <li>• dietary fibre intake</li> <li>• protein intake</li> <li>• vitamin and mineral supplements</li> <li>• alcohol</li> <li>• matching carbohydrate, insulin and physical activity</li> <li>• salt intake in hypertension</li> <li>• co-morbidities including nephropathy and renal failure,</li> <li>• coeliac disease, cystic fibrosis or eating disorders</li> <li>• use of peer support groups. (1.8.3.9)</li> </ul>	<p>Be aware of appropriate nutritional advice on common topics of concern and interest to adults living with type 1 diabetes, and be prepared to seek advice from colleagues with more specialised knowledge. Suggested common topics <b>include:</b></p> <ul style="list-style-type: none"> <li>• <b>body weight</b>, energy balance and obesity management</li> <li>• cultural and religious diets, feasts and fasts</li> <li>• foods sold as 'diabetic'</li> <li>• sweeteners</li> <li>• dietary fibre intake</li> <li>• protein intake</li> <li>• vitamin and mineral supplements</li> <li>• alcohol</li> <li>• matching carbohydrate, insulin and physical activity</li> <li>• salt intake in hypertension</li> <li>• comorbidities, including nephropathy and renal failure, coeliac disease, cystic fibrosis or eating disorders</li> <li>• use of peer support groups. <b>[2004, amended 2015]</b> (1.4.13)</li> </ul>	<p>There is no evidence of benefit for a low glycaemic index diet (see recommendation 1.4.3), so the reference about giving information about the glycaemic index of foods has been deleted.</p>
<p>Total glycated haemoglobin estimation, or assessment of glucose profiles, should</p>	<p><b>If HbA1c monitoring is invalid (because of disturbed erythrocyte turnover or</b></p>	<p>The GDG agreed to adopt the recommendation on</p>

<p>be used where haemoglobinopathy or haemoglobin turnover invalidate HbA1c measurement. (1.9.1.4)</p>	<p>abnormal haemoglobin type), estimate trends in blood glucose control using one of the following:</p> <ul style="list-style-type: none"> <li>• fructosamine estimation</li> <li>• quality-controlled blood glucose profiles</li> <li>• total glycated haemoglobin estimation (if abnormal haemoglobins). [2004, amended 2015] (1.6.5)</li> </ul>	<p>this topic from the NICE guideline on type 2 diabetes.</p>
<p>Self-monitoring skills should be taught close to the time of diagnosis and initiation of insulin therapy. (1.8.2.2)</p>	<p>Teach self-monitoring skills at the time of diagnosis and initiation of insulin therapy. [2004, amended 2015] (1.6.10)</p>	<p>The GDG stated that it is important that self-monitoring skills are taught as soon as type 1 diabetes is diagnosed.</p>
<p>Monitoring using sites other than the fingertips (often the forearm, using meters that require small volumes of blood and devices to obtain those small volumes) cannot be recommended as a routine alternative to conventional self-blood glucose monitoring. (1.8.2.8)</p>	<p>Monitoring blood glucose using sites other than the fingertips cannot be recommended as a routine alternative to conventional self-monitoring of blood glucose. [2004, amended 2015] (1.6.19)</p>	<p>Blood glucose has been stated for clarity. The statements about small volumes and special devices for alternative site monitoring have been removed because (1) the 2015 guideline supports the 2004 view that alternative site monitoring is not recommended, so the comment is redundant and (2) all meters now use small volumes.</p>
<p>For adults with erratic and unpredictable blood glucose control (hyper- and hypoglycaemia at no consistent times), rather than a change in a previously optimised insulin regimen, the following should be considered:</p> <ul style="list-style-type: none"> <li>• resuspension of insulin and injection technique</li> <li>• injection sites</li> <li>• self-monitoring skills</li> <li>• knowledge and self-management skills</li> <li>• nature of lifestyle</li> <li>• psychological and</li> </ul>	<p>For adults with erratic and unpredictable blood glucose control (hyperglycaemia and hypoglycaemia at no consistent times), rather than a change in a previously optimised insulin regimen, the following should be considered:</p> <ul style="list-style-type: none"> <li>• injection technique</li> <li>• injection sites</li> <li>• self-monitoring skills</li> <li>• knowledge and self-management skills</li> <li>• nature of lifestyle</li> <li>• psychological and psychosocial difficulties</li> </ul>	<p>Reference to resuspension of insulin is out of date and so has been deleted.</p>

<p>psychosocial difficulties</p> <ul style="list-style-type: none"> <li>• possible organic causes such as gastroparesis. (1.9.3.12)</li> </ul>	<ul style="list-style-type: none"> <li>• possible organic causes such as gastroparesis. <b>[2004, amended 2015]</b> (1.7.12)</li> </ul>	
<p>The injection-site condition should be checked annually and if new problems with blood glucose control occur. (1.9.4.8)</p>	<p>Check injection site condition <b>at least</b> annually and if new problems with blood glucose control occur. <b>[2004, amended 2015]</b> (1.8.7)</p>	<p>The GDG clarified that injection site condition can be checked more frequently than annually if appropriate.</p>
<p>Adults with type 1 diabetes should be informed that any available glucose/sucrose-containing fluid is suitable for the management of hypoglycaemic symptoms or signs in people who are able to swallow. Glucose-containing tablets or gels are also suitable for those able to dissolve or disperse these in the mouth and swallow the products. (1.9.5.1)</p>	<p>Explain to adults with type 1 diabetes that <b>a fast-acting form of glucose</b> is needed for the management of hypoglycaemic symptoms or signs in people who are able to <b>swallow</b>. <b>[2004, amended 2015]</b> (1.10.9)</p>	<p>The GDG clarified that a fast-acting form of glucose can be used for managing hypoglycaemia. The text specifying tablets or gels has been deleted. Glucogel is no longer listed in the BNF. The BNF also advises that other suitable forms of glucose can be used and therefore we did not want to state that only gels and tablets are appropriate..</p>
<p>Adults with decreased level of consciousness due to hypoglycaemia who are unable to take oral treatment safely should be:</p> <ul style="list-style-type: none"> <li>• given intramuscular glucagon by a trained user (intravenous glucose may be used by professionals skilled in obtaining intravenous access)</li> <li>• monitored for response at 10 minutes, and then given intravenous glucose if the level of consciousness is not improving significantly</li> <li>• then given oral carbohydrate when it is safe to administer it, and placed under continued observation by a third party who has been warned of the risk of</li> </ul>	<p>Adults with type 1 diabetes with a decreased level of consciousness as a result of hypoglycaemia <b>and so</b> unable to take oral treatment safely should be:</p> <ul style="list-style-type: none"> <li>• given intramuscular glucagon by <b>a family member or friend who has been shown how to use it</b> (intravenous glucose may be used by <b>healthcare</b> professionals skilled in obtaining intravenous access)</li> <li>• monitored for response at 10 minutes, and then given intravenous glucose if their level of consciousness is not improving significantly</li> <li>• then given oral carbohydrate when it is safe to administer it, and placed under continued observation by a third</li> </ul>	<p>The GDG clarified that this recommendation relates to people who are unable to protect their airway because of a decreased level of consciousness. Glucagon can be administered in an emergency situation. The Human Medicines Regulations 2012 schedule 19 lists glucagon as a medicine that can be administered in an emergency without a prescription. The MHRA states that 'Regulation 238 of the Human Medicines Regulations 2012 allows for certain prescription only medicines to be administered by</p>

<p>relapse. (1.9.5.3)</p>	<p>party who has been warned of the risk of relapse. <b>[2004, amended 2015]</b> (1.10.10)</p>	<p>anyone for the purpose of saving life in an emergency. The medicines this concerns are covered in Schedule 19 and are listed below.’ Therefore the recommendation has been changed to reflect that intramuscular glucagon does not have to be given by a trained user.</p>
<p>Nocturnal hypoglycaemia (symptomatic or detected on monitoring) should be managed by:</p> <ul style="list-style-type: none"> <li>• reviewing knowledge and self-management skills</li> <li>• reviewing current insulin regimen and evening eating habits and previous physical activity.</li> <li>• choosing an insulin type and regimen with less propensity to induce low glucose levels in the night hours, such as: <ul style="list-style-type: none"> <li>– isophane (NPH) insulin at bedtime</li> <li>– rapid-acting analogue with the evening meal</li> <li>– long-acting insulin analogues (insulin glargine)</li> <li>– – insulin pump. (1.9.5.8)</li> </ul> </li> </ul>	<p>Manage nocturnal hypoglycaemia (symptomatic or detected on monitoring) by:</p> <ul style="list-style-type: none"> <li>• reviewing knowledge and self-management skills</li> <li>• reviewing current insulin regimen, evening eating habits and previous physical activity.</li> <li>• choosing an insulin type and regimen that is less likely to induce low glucose levels <b>at night.</b> <b>[2004, amended 2015]</b> (1.10.13)</li> </ul>	<p>Details about insulin types have been deleted because the information is out of date and inconsistent with other recommendations in this guideline.</p>
<p>Healthcare professionals should be alert to the possibility of the development of other autoimmune disease in adults with type 1 diabetes (including Addison’s disease, pernicious anaemia and thyroid disorders). (1.12.4.2)</p>	<p>Be alert to the possibility of the development of other autoimmune disease in adults with type 1 diabetes (including Addison’s disease <b>and pernicious anaemia.</b> <b>For advice on monitoring for thyroid disease, see recommendation 1.15.5.</b> <b>[2004, amended 2015]</b> (1.12.2)</p>	<p>Mention of thyroid disorders has been deleted because thyroid disease is now covered by a separate recommendation to measure TSH levels at annual review.</p>
<p>Adults who have had</p>	<p>Provide intensive</p>	<p>Cross-reference to</p>

<p>myocardial infarction or stroke should be managed intensively, according to relevant non-diabetes guidelines. In the presence of angina or other ischaemic heart disease, beta-adrenergic blockers should be considered. (For use of insulin in these circumstances, see 'Hospital administration and intercurrent disease', Section 1.12.3.) (1.10.2.8)</p>	<p>management for adults who have had myocardial infarction or stroke, according to relevant non-diabetes guidelines. In the presence of angina or other ischaemic heart disease, beta-adrenergic blockers should be considered. (For use of insulin in these circumstances, see section 1.14). <b>For guidance on secondary prevention of myocardial infarction, see the NICE guideline on <a href="#">MI – secondary prevention</a>. [2004, amended 2015]</b> (1.13.7)</p>	<p>relevant NICE guideline added.</p>
<p>Concerns over potential side effects should not be allowed to inhibit advising and offering the necessary use of any class of drugs, unless the side effects become symptomatic or otherwise clinically significant. In particular:</p> <ul style="list-style-type: none"> <li>• selective beta-adrenergic blockers should not be avoided in adults on insulin</li> <li>• low-dose thiazides may be combined with beta-blockers</li> <li>• when calcium channel antagonists are prescribed, only long-acting preparations should be used</li> <li>• direct questioning should be used to detect the potential side effects of erectile dysfunction, lethargy and orthostatic hypotension with different drug classes. [1.10.3.5]</li> </ul>	<p><b>Do not allow</b> concerns over potential side effects to inhibit advising and offering the necessary use of any class of drugs, unless the side effects become symptomatic or otherwise clinically significant. In particular:</p> <ul style="list-style-type: none"> <li>• <b>do not avoid</b> selective beta-adrenergic blockers <b>where indicated</b> in adults on insulin</li> <li>• low-dose thiazides may be combined with beta-blockers</li> <li>• when calcium channel antagonists are prescribed, use only long-acting preparations</li> <li>• use direct questioning to detect the potential side effects of erectile dysfunction, lethargy and orthostatic hypotension with different drug classes. (1.13.12)</li> </ul>	<p>The GDG added 'where indicated' because the indications for beta blockers in pure hypertension are much more reduced now than in 2004.</p>
<p>A trial of a low-dose thiazide diuretic should be started as first-line therapy for raised blood pressure, unless the person with type 1 diabetes is already taking a renin-angiotensin system blocking</p>	<p><b>Start a trial of a renin-angiotensin system blocking drug as first-line therapy for hypertension in adults with type 1 diabetes. [2004, amended 2015]</b> (1.13.10)</p>	<p>The GDG did not review the evidence for this recommendation. However, the NICE guidance on hypertension has</p>

<p>drug for nephropathy (see 'Nephropathy', Section 1.1 1.2). Multiple drug therapy will often be required. (1.10.3.3)</p>		<p>changed since CG15 was published in 2004, and thiazides are no longer first-line therapy for any age group. Thiazides can elevate blood glucose. The GDG recommend renin–angiotensin system blockers as first-line therapy. They are recommended in NICE's <a href="#">hypertension</a> guideline as first-line therapy for people under 55 years, which accounts for most adults with type 1 diabetes and hypertension. For people over 55 years who do not have renal impairment, the NICE hypertension guideline recommends calcium channel blockers. As soon as renal impairment or albuminuria is detected, a renin–angiotensin system blocker is recommended for renal protection. Therefore it is sensible to recommend a renin–angiotensin blocker as first-line therapy for all adults with type 1 diabetes if they have hypertension. Mention of nephropathy has been removed; guidance on nephropathy is given in recommendation 1.15.19.</p>
<p>Throughout the course of an inpatient admission, the personal expertise of adults with type 1 diabetes (in managing their own diabetes) should be</p>	<p>Throughout the course of an inpatient admission, respect the personal expertise of adults with type 1 diabetes (in managing their own diabetes) and routinely</p>	<p>The GDG advised removing 'using the person's own systems', because hospitals increasingly use monitoring</p>

<p>respected and routinely integrated into ward-based blood glucose monitoring and insulin delivery, using the person with type 1 diabetes' own system. This should be incorporated into the nursing care plan. (1.12.3.2)</p>	<p>integrate this into ward-based blood glucose monitoring and <b>insulin delivery</b>. [2004, amended 2015] (1.14.8)</p>	<p>systems that are quality controlled and recorded automatically into electronic patient records that can be reviewed remotely by the diabetes professional team. The updated recommendation does not preclude the person using their own system in addition to the hospital system if they wish to do so. Use of such hospital monitoring systems improves patient care.</p>
<p>1.11.1.10 Referral to an ophthalmologist should occur for:</p> <ul style="list-style-type: none"> <li>• referable maculopathy:             <ul style="list-style-type: none"> <li>– exudate or retinal thickening within 1 disc diameter of the centre of the fovea</li> <li>– circinate or group of exudates within the macula (the macula is defined here as a circle centred on the fovea, of a diameter the distance between the temporal border of the optic disc and the fovea)</li> <li>– any microaneurysm or haemorrhage within 1 disc diameter of the centre of the fovea, only if associated with a best visual acuity of 6/12 or worse</li> </ul> </li> <li>• referable pre-proliferative retinopathy:             <ul style="list-style-type: none"> <li>– any venous beading</li> <li>– any venous loop or reduplication</li> <li>– any intraretinal</li> </ul> </li> </ul>	<p>Refer to an ophthalmologist for:</p> <ul style="list-style-type: none"> <li>• referable maculopathy:             <ul style="list-style-type: none"> <li>– exudate or retinal thickening within 1 disc diameter of the centre of the fovea</li> <li>– circinate or group of exudates within the macula (the macula is defined here as a circle centred on the fovea, of a diameter the distance between the temporal border of the optic disc and the fovea)</li> <li>– any microaneurysm or haemorrhage within 1 disc diameter of the centre of the fovea, only if associated with a best visual acuity of 6/12 or worse</li> </ul> </li> <li>• referable pre-proliferative retinopathy:             <ul style="list-style-type: none"> <li>– any venous beading</li> <li>– <b>any venous reduplication</b></li> <li>– any intraretinal microvascular abnormalities (IRMA)</li> </ul> </li> </ul>	<p>The recommendations on eye disease were reviewed by the National Screening Programme and were amended to make them consistent with the current practice of the diabetes eye screening programme.</p>

<p>microvascular abnormalities (IRMA)</p> <ul style="list-style-type: none"> <li>– multiple deep, round or blot haemorrhages (If cotton wool spots are present, look carefully for the above features, but cotton wool spots themselves do not define pre-proliferative retinopathy)</li> </ul> <ul style="list-style-type: none"> <li>• any unexplained drop in visual acuity. (1.11.1.10)</li> </ul>	<ul style="list-style-type: none"> <li>– multiple deep, round or blot haemorrhages (If cotton wool spots are present, look carefully for the above features, but cotton wool spots themselves do not define pre-proliferative retinopathy)</li> </ul> <ul style="list-style-type: none"> <li>• any <b>large sudden</b> unexplained drop in visual acuity. <b>[2004, amended 2015]</b> (1.15.14)</li> </ul>	
<p>If ACE inhibitors are not tolerated, angiotensin 2 receptor antagonists should be substituted. Combination therapy is not recommended at present. (1.11.2.6)</p>	<p>If ACE inhibitors are not tolerated, substitute angiotensin 2 receptor antagonists. Combination therapy is not <b>recommended</b>. <b>[2004, amended 2015]</b> (1.15.20)</p>	<p>‘at present’ has been removed in view of evidence known to the GDG that the combination can be harmful, increasing risk of hyperkalaemia and acute renal injury.</p>
<p>The management of the symptoms of autonomic neuropathy should include standard interventions for the manifestations encountered (for example, for erectile dysfunction or abnormal sweating). (1.11.4.9)</p>	<p>When managing the symptoms of autonomic neuropathy, include standard interventions for the manifestations encountered (for example, <b>for abnormal sweating and postural hypotension</b>). <b>[2004, amended 2015]</b> (1.15.36)</p>	<p>The GDG added postural hypertension because this is an important manifestation of autonomic neuropathy. There are now separate recommendations about managing erectile dysfunction (1.15.30–1.15.32) and gastroparesis (1.15.1–1.15.4).</p>
<p>Diabetes professionals should ensure that they have appropriate skills in the detection and basic management of non-severe psychological disorders in people from different cultural backgrounds. They should be familiar with appropriate counselling techniques and appropriate drug therapy, while arranging prompt referral to specialists of those people</p>	<p>Diabetes professionals should:</p> <ul style="list-style-type: none"> <li>• ensure that they have appropriate skills in the detection and basic management of non-severe psychological disorders in people from different cultural backgrounds</li> <li>• be familiar with appropriate counselling techniques and drug therapy, while arranging</li> </ul>	<p>Cross-references to relevant NICE guidelines have been added for information.</p>

<p>in whom psychological difficulties continue to interfere significantly with well-being or diabetes self-management. (1.12.5.2)</p>	<p>prompt referral to specialists of those people in whom psychological difficulties continue to interfere significantly with wellbeing or diabetes self-management.</p> <p>See also the NICE guidelines on <a href="#">common mental health disorders, generalised anxiety disorder and panic disorder (with or without agoraphobia) in adults</a> and <a href="#">depression in adults with a chronic health problem</a>. [2004, amended 2015] (1.15.41)</p>	
<p>Members of multidisciplinary professional teams should be alert to the possibility of bulimia nervosa, anorexia nervosa and insulin dose manipulation in adults with type 1 diabetes with:</p> <ul style="list-style-type: none"> <li>• over-concern with body shape and weight</li> <li>• low body mass index</li> <li>• poor overall blood glucose control. (1.12.6.1)</li> </ul>	<p>Members of diabetes professional teams should be alert to the possibility of bulimia nervosa, anorexia nervosa and insulin dose manipulation in adults with type 1 diabetes with:</p> <ul style="list-style-type: none"> <li>• over-concern with body shape and weight</li> <li>• low BMI</li> <li>• hypoglycaemia</li> <li>• poor overall blood glucose control.</li> </ul> <p>See also the NICE guideline on <a href="#">eating disorders</a> [2004, amended 2015] (1.15.42)</p>	<p>The GDG stated that hypoglycaemia is another possible indicator of eating disorders.</p> <p>Cross-reference to the relevant NICE guideline has been added for information.</p>
	<p>1.13.2, 1.13.8, 1.15.18</p>	<p>Change made from '[abnormal] albumin excretion rate' to 'albuminuria' for accuracy.</p>
	<p>1.2.3, 1.3.6, 1.4.8, 1.6.11, 1.11.6, 1.11.8, 1.11.10, 1.11.12, 1.15.11, 1.15.12, 1.15.13</p>	<p>NICE has made editorial changes to the original wording to clarify the action to be taken (no change to meaning): a verb has been added, the verb used has been changed or other wording has changed for clarification.</p>

- 1 ***Changes to recommendation wording for clarification only (no***
- 2 ***change to meaning)***
- 3 Yellow highlighting has not been applied to these changes.

<b>Recommendation numbers in current guideline</b>	<b>Comment</b>
All recommendations except those labelled <b>[new 2015]</b>	Recommendations have been edited into the direct style (in line with current NICE style for recommendations in clinical guidelines) where possible. Yellow highlighting has not been applied to these changes. Where applicable, terminology has been made consistent within the guideline and with terminology that will be used in other updates of NICE guidelines on diabetes (diabetes in pregnancy [publication expected February 2015], type 2 diabetes and diabetes in children and young people [publication expected August 2015]) – for example, 'impaired awareness of hypoglycaemia' rather than 'hypoglycaemia unawareness'; 'blood glucose control' rather than 'glycaemic control'.
1.2.6, 1.2.7, 1.4.5, 1.8.1, 1.8.6, 1.10.10, 1.13.11, 1.14.9, 1.14.11, 1.15.33, 1.15.35, 1.15.37, 1.15.40	Type 1 diabetes is specified for clarity (original wording had 'diabetes' or did not specify diabetes at all).
1.11.4 to 1.11.12	The population is specified as adults with DKA for clarity.
1.15.6, 1.15.7, 1.15.8, 1.15.9, 1.15.11	'eye surveillance' has been changed to 'eye screening', in line with current terminology.

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