Hypothermia: prevention and management in adults having surgery 3

NICE guideline: short version

Draft for consultation, September 2016

This guideline covers preventing and managing inadvertent hypothermia in adults having surgery. Inadvertent hypothermia is a common complication of surgery and is associated with poor outcomes for patients. This guideline offers advice on assessing patients' risk of hypothermia, measuring and monitoring temperature, and devices for keeping patients warm before, during and after surgery. The guideline does not cover deliberate induction of hypothermia for medical reasons.

Who is it for?

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- Healthcare professionals
- Adults having surgery, their families and carers

This guideline will update NICE guideline CG65 (published April 2008).

We have added new recommendations on <u>measuring temperature</u>, <u>warming patients</u> <u>before induction of anaesthesia</u> and <u>warming patients after induction of anaesthesia</u>.

You are invited to comment on the new recommendations in this guideline. These are marked as [new 2016].

You are also invited to comment on recommendations that NICE proposes to delete from the 2008 guideline. These are shown in <u>recommendations that have been</u> <u>deleted or changed</u>.

We have not updated recommendations shaded in grey, and cannot accept comments on them. In some cases, we have made minor wording changes for

clarification.

See <u>Update information</u> for a full explanation of what is being updated.

This version of the guideline contains the draft recommendations, context and recommendations for research. The supporting information and evidence for the 2016 recommendations is contained in the <u>addendum</u>. Evidence for the 2008 recommendations is in the <u>full version</u> of the 2008 guideline.

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1 Recommendations

People have the right to be involved in discussions and make informed decisions about their care, as described in <u>your care</u>.

Making decisions using NICE guidelines explains how we use words to show the strength (or certainty) of our recommendations, and has information about prescribing medicines (including off-label use), professional guidelines, standards and laws (including on consent and mental capacity), and safeguarding.

Throughout the guidance 'temperature' is used to denote core temperature. The phrase 'comfortably warm' is used in recommendations relating to both the preoperative and postoperative phases, and refers to the expected normal temperature range of adult patients, which is between 36.5°C and 37.5°C.

2	1.1	Perioperative care
3	1.1.1	Patients (and their families and carers) should be informed that:
4 5 7 8 9		 staying warm before surgery will lower the risk of postoperative complications the hospital environment may be colder than their own home they should bring additional clothing, such as a dressing gown, a vest, warm clothing and slippers, to help them keep comfortably warm they should tell staff if they feel cold at any time during their hospital stay. [2008]
11 12	1.1.2	When using any temperature recording or warming device, healthcare professionals should:
13 14 15 16		 be trained in their use maintain them in accordance with manufacturers' and suppliers' instructions comply with local infection control policies. [2008]

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1	1.1.3	When using any device to measure patient temperature, healthcare		
2		professionals should:		
3		• be aware of, and carry out, any adjustments that need to be made in		
4		order to obtain an estimate of core temperature from that recorded at		
5		the site of measurement		
6		• be aware of any such adjustments that are made automatically by the		
7		device used. [2008]		
8	1.1.4	Measure the patient's core temperature directly, using 1 of the following		
9		sites and basing the choice of site on its suitability for the patient, the type		
10		of surgery and the anaesthetic:		
11		bladder		
12		oesophagus		
13		 pulmonary artery catheter. [new 2016] 		
14	1.1.5	If direct core temperature measurement is not suitable, assess core		
15		temperature indirectly, using a site or device that produces a		
16		measurement accurate to within 0.5°C of the true core temperature. At the		
17		time of consultation these are:		
18		deep forehead		
19		infrared temporal		
20		infrared tympanic		
21		rectal		
22		sublingual		
23		 thermocouple forehead with a +2°C correction factor. [new 2016] 		
24	1.1.6	Do not use any site or device to indirectly assess core temperature in		
25		adults having surgery that has not been shown in research studies to		
26		produce a measurement accurate to within 0.5°C of true core		
27		temperature. [new 2016]		

1 **1.2 Preoperative phase**

2 The preoperative phase is defined as the hour before induction of anaesthesia,

3 during which the patient is prepared for surgery on the ward or in the emergency

4 department, including possible use of premedication.

5	1.2.1	Each patient should be assessed for their risk of inadvertent perioperative
6		hypothermia and potential adverse consequences before transfer to the
7		theatre suite. Patients should be managed as higher risk (see
8		recommendation 1.3.7) if any two of the following apply:
0		Assertions Operate of Asserthesister (AOA) and the Mitch Mitch end
9		American Society of Anestnesiologists (ASA) grade II to V (the higher
10		the grade, the greater the risk)
11		 preoperative temperature below 36.0°C (and preoperative warming is
12		not possible because of clinical urgency)
13		 undergoing combined general and regional anaesthesia
14		undergoing major or intermediate surgery
15		• at risk of cardiovascular complications. [2008]
16	1.2.2	Offer active warming for at least 30 minutes before induction of
17		anaesthesia to all patients having general anaesthesia or central neural
18		blockade for surgery, unless this will delay emergency surgery. [new
19		2016]
20	1.2.3	Pay particular attention to the comfort of patients with communication
21		difficulties during the preoperative phase. [new 2016]
22	1.2.4	Special care should be taken to keep patients comfortably warm when
23		they are given premedication (for example, nefopam, tramadol,
24		midazolam or opioids). [2008]
25	1.2.5	The patient's temperature should be measured and documented in the
26		hour before they leave the ward or emergency department. [2008]
27	1.2.6	If the patient's temperature is below 36.0°C:

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1		 active warming should be started preoperatively on the ward or in the
2		emergency department (unless there is a need to expedite surgery
3		because of clinical urgency, for example bleeding or critical limb
4		ischaemia)
5		active warming should be maintained throughout the intraoperative
6		phase. [2008, amended 2016]
7	1.2.7	The patient's temperature should be 36.0°C or above before they are
8		transferred from the ward or emergency department (unless there is a
9		need to expedite surgery because of clinical urgency, for example
10		bleeding or critical limb ischaemia). [2008]
11	1 2 8	On transfor to the theatre suite:
11	1.2.0	
12		 the patient should be kept comfortably warm
13		• the patient should be encouraged to walk to theatre where appropriate.
14		[2008]
15	1.3	Intraoperative phase
15 16	1.3 The intrao	<i>Intraoperative phase</i> perative phase is defined as total anaesthesia time, from the first
15 16 17	1.3 The intrao anaesthet	<i>Intraoperative phase</i> operative phase is defined as total anaesthesia time, from the first ic intervention through to patient transfer to the recovery area of the
15 16 17 18	1.3 The intrao anaesthet theatre su	<i>Intraoperative phase</i> operative phase is defined as total anaesthesia time, from the first ic intervention through to patient transfer to the recovery area of the ite.
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15 16 17 18 19 20	1.3 The intrao anaesthet theatre su 1.3.1	Intraoperative phase operative phase is defined as total anaesthesia time, from the first ic intervention through to patient transfer to the recovery area of the ite. The patient's temperature should be measured and documented before induction of anaesthesia and then every 30 minutes until the end of
 15 16 17 18 19 20 21 	1.3 The intrao anaesthet theatre su 1.3.1	Intraoperative phase operative phase is defined as total anaesthesia time, from the first ic intervention through to patient transfer to the recovery area of the ite. The patient's temperature should be measured and documented before induction of anaesthesia and then every 30 minutes until the end of surgery. [2008]
 15 16 17 18 19 20 21 22 	1.3 The intrao anaesthet theatre su 1.3.1 1.3.2	Intraoperative phase perative phase is defined as total anaesthesia time, from the first ic intervention through to patient transfer to the recovery area of the ite. The patient's temperature should be measured and documented before induction of anaesthesia and then every 30 minutes until the end of surgery. [2008] Standard critical incident reporting should be considered for any patient
 15 16 17 18 19 20 21 22 23 	1.3 The intrao anaesthet theatre su 1.3.1 1.3.2	Intraoperative phase perative phase is defined as total anaesthesia time, from the first ic intervention through to patient transfer to the recovery area of the ite. The patient's temperature should be measured and documented before induction of anaesthesia and then every 30 minutes until the end of surgery. [2008] Standard critical incident reporting should be considered for any patient arriving at the theatre suite with a temperature below 36.0°C. [2008]
 15 16 17 18 19 20 21 22 23 24 	 1.3 The intrao anaesthet theatre su 1.3.1 1.3.2 1.3.3 	Intraoperative phase perative phase is defined as total anaesthesia time, from the first ic intervention through to patient transfer to the recovery area of the ite. The patient's temperature should be measured and documented before induction of anaesthesia and then every 30 minutes until the end of surgery. [2008] Standard critical incident reporting should be considered for any patient arriving at the theatre suite with a temperature below 36.0°C. [2008] Induction of anaesthesia should not begin unless the patient's
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 15 16 17 18 19 20 21 22 23 24 25 26 	 1.3 The intrao anaesthet theatre su 1.3.1 1.3.2 1.3.3 	Intraoperative phase perative phase is defined as total anaesthesia time, from the first ic intervention through to patient transfer to the recovery area of the ite. The patient's temperature should be measured and documented before induction of anaesthesia and then every 30 minutes until the end of surgery. [2008] Standard critical incident reporting should be considered for any patient arriving at the theatre suite with a temperature below 36.0°C. [2008] Induction of anaesthesia should not begin unless the patient's temperature is 36.0°C or above (unless there is a need to expedite surgery because of clinical urgency, for example bleeding or critical limb
 15 16 17 18 19 20 21 22 23 24 25 26 27 	 1.3 The intrao anaesthet theatre su 1.3.1 1.3.2 1.3.3 	Intraoperative phase perative phase is defined as total anaesthesia time, from the first ic intervention through to patient transfer to the recovery area of the ite. The patient's temperature should be measured and documented before induction of anaesthesia and then every 30 minutes until the end of surgery. [2008] Standard critical incident reporting should be considered for any patient arriving at the theatre suite with a temperature below 36.0°C. [2008] Induction of anaesthesia should not begin unless the patient's temperature is 36.0°C or above (unless there is a need to expedite surgery because of clinical urgency, for example bleeding or critical limb ischaemia). [2008]

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1 2		 the ambient temperature should be at least 21°C while the patient is exposed 	
3		 once active warming is established, the ambient temperature may be reduced to allow better working conditions 	
5 6		 using equipment to cool the surgical team should also be considered. [2008, amended 2016] 	
7 8 9	1.3.5	The patient should be adequately covered throughout the intraoperative phase to conserve heat, and exposed only during surgical preparation. [2008]	
10 11	1.3.6	Intravenous fluids (500 ml or more) and blood products should be warmed to 37°C using a fluid warming device. [2008]	
12 13	1.3.7	Warm patients intraoperatively from induction of anaesthesia, using a forced air warming device, if they are:	
14 15 16		 having anaesthesia for more than 30 minutes or having anaesthesia for less than 30 minutes and are at higher risk of inadvertent perioperative hypothermia (see <u>recommendation 1.2.1</u>). 	
17 18		Consider a resistive heating mattress or resistive heating blanket if a forced air warming device is unsuitable. [new 2016]	
19 20 21	1.3.8	The temperature setting on forced air warming devices should be set at maximum and then adjusted to maintain a patient temperature of at least 36.5°C. [2008]	
22 23	1.3.9	All irrigation fluids used intraoperatively should be warmed in a thermostatically controlled cabinet to a temperature of 38–40°C. [2008]	
24	1.4	Postoperative phase	
25 26	The postoperative phase is defined as the 24 hours after the patient has entered the recovery area of the theatre suite.		

271.4.1The patient's temperature should be measured and documented on28admission to the recovery room and then every 15 minutes. [2008]

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1		• Ward transfer should not be arranged unless the patient's temperature
2		is 36.0°C or above.
3		• If the patient's temperature is below 36.0°C, they should be actively
4		warmed using forced air warming until they are discharged from the
5		recovery room or until they are comfortably warm. [2008]
6	1.4.2	Patients should be kept comfortably warm when back on the ward.
7		• Their temperature should be measured and documented on arrival at
8		the ward.
9		• Their temperature should then be measured and documented as part of
10		routine 4-hourly observations.
11		They should be provided with at least one cotton sheet plus two
12		blankets, or a duvet. [2008]
13	1.4.3	If the patient's temperature falls below 36.0°C while on the ward:
14		 they should be warmed using forced air warming until they are
15		comfortably warm
16		• their temperature should be measured and documented at least every
17		30 minutes during warming. [2008]

Putting this guideline into practice

- 19 NICE has produced <u>tools and resources</u> to help you put this guideline into practice.
- 20 [Optional paragraph if issues raised] Some issues were highlighted that might need
- 21 specific thought when implementing the recommendations. These were raised during
- 22 the development of this guideline. They are:
- 23 [add any issues specific to guideline here]
- [Use 'Bullet left 1 last' style for the final item in this list.]
- 25 Putting recommendations into practice can take time. How long may vary from
- 26 guideline to guideline, and depends on how much change in practice or services is
- 27 needed. Implementing change is most effective when aligned with local priorities.

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1 Changes recommended for clinical practice that can be done quickly – like changes

- 2 in prescribing practice should be shared quickly. This is because healthcare
- 3 professionals should use guidelines to guide their work as is required by
- 4 professional regulating bodies such as the General Medical and Nursing and
- 5 Midwifery Councils.

6 Changes should be implemented as soon as possible, unless there is a good reason

7 for not doing so (for example, if it would be better value for money if a package of

8 recommendations were all implemented at once).

9 Different organisations may need different approaches to implementation, depending

10 on their size and function. Sometimes individual practitioners may be able to respond

- 11 to recommendations to improve their practice more quickly than large organisations
- 12 Here are some pointers to help put NICE guidelines into practice:

13 **1. Raise awareness** through routine communication channels, such as email or

- 14 newsletters, regular meetings, internal staff briefings and other communications with
- 15 all relevant partner organisations. Identify things staff can include in their own

16 practice straight away.

17 2. Identify a lead with an interest in the topic to champion the guideline and motivate
18 others to support its use and make service changes, and to find out any significant
19 issues locally.

3. Carry out a baseline assessment against the recommendations to find whether
there are gaps in current service provision.

4. Think about what data you need to measure improvement and plan how you
will collect it. You may want to work with other health and social care organisations
and specialist groups to compare current practice with the recommendations. This
may also help identify local issues that will slow or prevent implementation.

- 26 5. **Develop an action plan**, with the steps needed to put the guideline into practice,
- 27 and make sure it is ready as soon as possible. Big, complex changes may take
- longer to implement, but some may be quick and easy to do. An action plan will help
- in both cases.

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6. **For very big changes** include milestones and a business case, which will set out additional costs, savings and possible areas for disinvestment. A small project group could develop the action plan. The group might include the guideline champion, a senior organisational sponsor, staff involved in the associated services, finance and information professionals.

7. Implement the action plan with oversight from the lead and the project group.
Big projects may also need project management support.

8. Review and monitor how well the guideline is being implemented through the
9 project group. Share progress with those involved in making improvements, as well
10 as relevant boards and local partners.

11 NICE provides a comprehensive programme of support and resources to maximise

12 uptake and use of evidence and guidance. See our <u>into practice</u> pages for more

13 information.

Also see Leng G, Moore V, Abraham S, editors (2014) Achieving high quality care –
 practical experience from NICE. Chichester: Wiley.

16 **Context**

17 Inadvertent perioperative hypothermia is a common but preventable complication of

18 perioperative procedures, which is associated with poor outcomes for patients.

19 Inadvertent perioperative hypothermia should be distinguished from the deliberate

20 induction of hypothermia for medical reasons, which is not covered by this guideline.

21 In this guideline, hypothermia is defined as a patient core temperature of below 22 36.0°C. Hereafter, 'temperature' is used to denote core temperature. Adult surgical 23 patients are at risk of developing hypothermia at any stage of the perioperative 24 pathway. In the guideline, the perioperative pathway is divided into three phases: the 25 preoperative phase is defined as the hour before induction of anaesthesia (when the 26 patient is prepared for surgery on the ward or in the emergency department), the 27 intraoperative phase is defined as total anaesthesia time, and the postoperative phase is defined as the 24 hours after entry into the recovery area in the theatre 28 29 suite (which will include transfer to and time spent on the ward). The phrase

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'comfortably warm' is used in recommendations relating to both the preoperative and
 postoperative phases, and refers to the expected normal temperature range of adult

- postoperative phases, and refers to the expected normal temperature
 patients (between 36.5°C and 37.5°C).
- 4 During the first 30 to 40 minutes of anaesthesia, a patient's temperature can drop to
- 5 below 35.0°C. Reasons for this include loss of the behavioural response to cold and
- 6 the impairment of thermoregulatory heat-preserving mechanisms under general or
- 7 regional anaesthesia, anaesthesia-induced peripheral vasodilation (with associated
- 8 heat loss), and the patient getting cold while waiting for surgery on the ward or in the
- 9 emergency department.
- 10 In 2016 we updated the guideline to take account of new evidence on active
- 11 warming devices. We also added new recommendations on the site and method of
- 12 measuring temperature, which had been identified as an area where guidance would
- 13 be clinically useful.

14 More information

To find out what NICE has said on topics related to this guideline, see our web page on surgical care.

15

Recommendations for research

- 17 The guideline committee has made the following recommendations for research.
- 18 As part of the 2016 update, the standing committee made an additional research
- 19 recommendation on combined methods of intraoperative active warming compared
- 20 with a single method, and removed 3 research recommendations, on preoperative
- 21 insulation and warming, comparison of intraoperative warming devices, and use of
- both preoperative and intraoperative warming. Details can be found in the
- addendum.

1 1 Combined methods of intraoperative active warming compared

2 with a single method

- 3 What is the clinical and cost effectiveness of combined methods of intraoperative
- 4 active warming compared with a single method in preventing inadvertent
- 5 perioperative hypothermia?

6 Why this is important

7 A combination of active warming devices, such as forced air warming together with a 8 resistive heating mattress, is usually used to warm patients during surgery. However, 9 there is not enough evidence to show whether this is more clinically effective than a 10 single active warming device, such forced air warming on its own. Large randomised controlled trials with at least 100 patients in each arm should be carried out to 11 12 compare combined methods of intraoperative active warming (such as forced air 13 warming together with a resistive heating mattress, or a resistive heating mattress 14 together with a resistive heating blanket) with a single method of active warming 15 (such as forced air warming). All intravenous fluids should be warmed to 37°C. 16 Primary outcomes should be core temperature at the end of surgery and incidence of 17 hypothermia. Patients should be stratified by anaesthesia duration and type of surgery. Adverse effects and numbers of patients with complications of hypothermia 18 19 (for example, morbid cardiac events or wound infections) should be recorded. [new 20 2016]

21 **2 Temperature thresholds for preoperative warming**

22 What is the optimum temperature target when warming patients preoperatively?

23 Why this is important

- 24 Preoperative warming is intended to minimise the impact of redistribution
- 25 hypothermia by reducing the temperature difference between the patient's core
- 26 temperature and peripheral temperature. There is a lack of evidence for the optimum
- 27 preoperative temperature for preventing intraoperative hypothermia. Large RCTs
- 28 (with at least 100 patients in each arm) should be conducted in adults undergoing
- surgery to compare warming patients to 36.5°C and 37.0°C in the preoperative
- 30 phase. Warming should be continued intraoperatively in all patients. All intravenous

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- 1 fluids given should be warmed to 37°C. Primary outcomes should be the incidence of
- 2 hypothermia, and patient temperature intraoperatively (at 15, 30, 60 and
- 3 120 minutes) and in recovery. The duration of warming required to achieve the target
- 4 preoperative temperature should be recorded. Adverse effects (including patient
- 5 discomfort) and numbers of patients with complications of hypothermia (for example,
- 6 morbid cardiac events, wound infection) should be recorded. [2008]

7 **3 Effects of nutritional solutions**

- 8 Does the infusion of nutritional solutions such as amino acids and fructose further
- 9 reduce the incidence of inadvertent perioperative hypothermia in patients receiving
- 10 intraoperative warming?

11 Why this is important

- 12 Limited evidence suggests that infusion of amino acids or fructose in the
- 13 preoperative and intraoperative phases may prevent hypothermia. Such infusions
- 14 may also have additional benefits in fasted patients. A large RCT (with at least 100
- 15 patients in each arm) comparing infusions of amino acids, fructose and saline should
- 16 be conducted in adults undergoing surgery. These infusions should be started before
- 17 the induction of anaesthesia and continued throughout the intraoperative phase. All
- 18 patients should receive forced air warming intraoperatively and all intravenous fluids
- 19 given should be warmed to 37°C. Primary outcomes should be the incidence of
- 20 hypothermia, and patient temperature intraoperatively (at 15, 30, 60 and
- 21 120 minutes) and in recovery. Adverse effects and numbers of patients with
- 22 complications of hypothermia (for example, morbid cardiac events, wound infections)
- 23 should be recorded. [2008]

24 Update information

- 25 September 2016
- 26 New recommendations have been added on measuring temperature, warming
- 27 patients before induction of anaesthesia and warming patients after induction of
- 28 <u>anaesthesia</u>. These are marked as **[new 2016].**

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- 1 NICE proposes to delete some recommendations from the 2008 guideline, because
- 2 either the evidence has been reviewed and the recommendations have been
- 3 updated, or NICE has updated other relevant guidance and has replaced the original
- 4 recommendations. <u>Recommendations that have been deleted</u> sets out these
- 5 recommendations and includes details of replacement recommendations. Where
- 6 there is no replacement recommendation, an explanation for the proposed deletion is
- 7 given.
- 8 Where recommendations are shaded in grey and end **[2008]**, the evidence has not
- 9 been reviewed since the original guideline.
- 10 Where recommendations are shaded in grey and end [2008, amended 2016], the
- 11 evidence has not been reviewed but changes have been made to the
- 12 recommendation wording that change the meaning (for example, because of
- 13 equalities duties or a change in the availability of medicines, or incorporated
- 14 guidance has been updated). These changes are marked with yellow shading, and
- 15 explanations of the reasons for the changes are given in 'Recommendations that
- 16 have been deleted or changed' for information.
- 17 See also the <u>original NICE guideline and supporting documents</u>.

18 **Recommendations that have been deleted or changed**

19 Recommendations that have been deleted

20

Recommendation in 2008 guideline	Comment
Healthcare professionals should ensure that patients are kept comfortably warm while waiting for surgery by giving them at least one cotton sheet plus two blankets, or a duvet. [1.2.2]	Replaced by: Offer active warming for at least 30 minutes before induction of anaesthesia to all patients having general anaesthesia or central neural blockade for surgery. [new 2016] [1.2.2]
Patients who are at higher risk of inadvertent perioperative hypothermia (see section 1.2.1) and who are having anaesthesia for less than 30 minutes should be warmed intraoperatively from induction of anaesthesia using a forced air warming device. [1.3.7]	Replaced by: Warm patients intraoperatively from induction of anaesthesia, using a forced air warming device or a resistive heating blanket, if they are:

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	30 minutes or	
	 having anaesthesia for less than 30 minutes and are at higher risk of inadvertent perioperative hypothermia (see recommendation 1.2.1). 	
	Consider a resistive heating mattress or resistive heating blanket if a forced air warming device is unsuitable. [new 2016] [1.3.7]	
All patients who are having anaesthesia	Replaced by:	
for longer than 30 minutes should be warmed intraoperatively from induction of anaesthesia using a forced air warming device. [1.3.8].	Warm patients intraoperatively from induction of anaesthesia, using a forced air warming device or a resistive heating blanket, if they are:	
	 having anaesthesia for more than 30 minutes or 	
	 having anaesthesia for less than 30 minutes and are at higher risk of inadvertent perioperative hypothermia (see recommendation 1.2.1). 	
	Consider a resistive heating mattress or resistive heating blanket if a forced air warming device is unsuitable. [new 2016] [1.3.7]	

1

2

1 Amended recommendation wording (change to meaning)

2

Recommendation in 2008	Recommendation in current	Reason for change	
guideline	guideline		
If the patient's temperature is below 36.0°C:	If the patient's temperature is below 36.0°C:	'Forced air warming' has been amended to 'active warming' to include	
 forced air warming should be started preoperatively on the ward or in the emergency department (unless there is a need to expedite surgery because of clinical urgency, for example bleeding or critical limb ischaemia) forced air warming should be maintained throughout the intraoperative phase. [1.2.5] 	 active warming should be started preoperatively on the ward or in the emergency department (unless there is a need to expedite surgery because of clinical urgency, for example bleeding or critical limb ischaemia) active warming should be maintained throughout the intraoperative phase. [2008] [1.2.6] 	other types of active warming.	
1.3.4 In the theatre suite:	1.3.4 In the theatre suite:	'Forced air warming' has	
 the ambient temperature should be at least 21°C while the patient is exposed 	 the ambient temperature should be at least 21°C while the patient is exposed 	warming' to include other types of active warming.	
 once forced air warming is established, the ambient temperature may be reduced to allow better working conditions using equipment to cool the surgical team should also be considered. [1.3.4] 	 once active warming is established, the ambient temperature may be reduced to allow better working conditions. using equipment to cool the surgical team should also be considered. [2008] [1.3.4] 		

3

4 Appendix: The algorithm

5 There is a care pathway for the management of perioperative hypothermia in adults

- 6 on page 27 of the <u>full guideline</u>.
- 7 ISBN:

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