NICE National Institute for Health and Care Excellence



Intravenous fluid therapy in adults in hospital

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Your responsibility

The recommendations in this guideline represent the view of NICE, arrived at after careful consideration of the evidence available. When exercising their judgement, professionals and practitioners are expected to take this guideline fully into account, alongside the individual needs, preferences and values of their patients or the people using their service. It is not mandatory to apply the recommendations, and the guideline does not override the responsibility to make decisions appropriate to the circumstances of the individual, in consultation with them and their families and carers or guardian.

All problems (adverse events) related to a medicine or medical device used for treatment or in a procedure should be reported to the Medicines and Healthcare products Regulatory Agency using the <u>Yellow Card Scheme</u>.

Local commissioners and providers of healthcare have a responsibility to enable the guideline to be applied when individual professionals and people using services wish to use it. They should do so in the context of local and national priorities for funding and developing services, and in light of their duties to have due regard to the need to eliminate unlawful discrimination, to advance equality of opportunity and to reduce health inequalities. Nothing in this guideline should be interpreted in a way that would be inconsistent with complying with those duties.

Commissioners and providers have a responsibility to promote an environmentally sustainable health and care system and should <u>assess and reduce the environmental</u> <u>impact of implementing NICE recommendations</u> wherever possible.

Contents

Overview	4			
Who is it for?	4			
Introduction	5			
Key priorities for implementation	8			
Principles and protocols for intravenous fluid therapy	8			
Assessment and monitoring	8			
Resuscitation	10			
Routine maintenance	10			
Training and education	11			
Recommendations	12			
1.1 Principles and protocols for intravenous fluid therapy	12			
1.2 Assessment and monitoring	14			
1.3 Resuscitation	17			
1.4 Routine maintenance	17			
1.5 Replacement and redistribution	19			
1.6 Training and education				
Terms used in this guideline	26			
Recommendations for research	27			
1 Assessment and monitoring	27			
Finding more information and committee details	28			
Update information	29			

This guideline is the basis of QS66.

Overview

This guideline covers the general principles for managing intravenous (IV) fluid therapy in hospital inpatients aged 16 and over with a range of conditions. It aims to help prescribers understand the optimal amount and composition of IV fluids to be administered and the best rate at which to give them, to improve fluid prescribing and outcomes among people in hospital. It does not cover pregnant women, and those with severe liver or renal disease, diabetes or burns.

Who is it for?

- Healthcare professionals
- People who receive IV fluid therapy in hospital and their families and carers

Introduction

Many adult hospital inpatients need intravenous (IV) fluid therapy to prevent or correct problems with their fluid and/or electrolyte status. Deciding on the optimal amount and composition of IV fluids to be administered and the best rate at which to give them can be a difficult and complex task, and decisions must be based on careful assessment of the patient's individual needs.

Errors in prescribing IV fluids and electrolytes are particularly likely in emergency departments, acute admission units, and general medical and surgical wards rather than in operating theatres and critical care units. Surveys have shown that many staff who prescribe IV fluids know neither the likely fluid and electrolyte needs of individual patients, nor the specific composition of the many choices of IV fluids available to them. Standards of recording and monitoring IV fluid and electrolyte therapy may also be poor in these settings. IV fluid management in hospital is often delegated to the most junior medical staff who frequently lack the relevant experience and may have received little or no specific training on the subject.

The <u>National Confidential Enquiry into Perioperative Deaths report Extremes of age</u> in 1999 highlighted that a significant number of hospitalised patients were dying as a result of infusion of too much or too little fluid. The report recommended that fluid prescribing should be given the same status as drug prescribing. Although mismanagement of fluid therapy is rarely reported as being responsible for patient harm, it is likely that as many as 1 in 5 patients on IV fluids and electrolytes suffer complications or morbidity due to their inappropriate administration.

There is also considerable debate about the best IV fluids to use (particularly for more seriously ill or injured patients), resulting in wide variation in clinical practice. Many reasons underlie the ongoing debate, but most revolve around difficulties in interpretation of both trial evidence and clinical experience, including the following factors:

- Many accepted practices of IV fluid prescribing were developed for historical reasons rather than through clinical trials.
- Trials cannot easily be included in meta-analyses because they examine varied outcome measures in heterogeneous groups, comparing not only different types of fluid with different electrolyte content, but also different volumes and rates of

administration and, in some cases, the additional use of inotropes or vasopressors.

- Most trials have been undertaken in operating theatres and critical care units rather than admission units or general and elderly care settings.
- Trials claiming to examine best early therapy for fluid resuscitation have actually evaluated therapy choices made after initial fluid resuscitation, with patients already in critical care or operating theatres.
- Many trials inferring best therapy for fluid resuscitation after acute fluid loss have actually examined situations of hypovolaemia induced by anaesthesia.

There is a clear need for guidance on IV fluid therapy for general areas of hospital practice, covering both the prescription and monitoring of IV fluid and electrolyte therapy, and the training and educational needs of all hospital staff involved in IV fluid management.

The aim of this NICE guideline is to help prescribers understand the:

- physiological principles that underpin fluid prescribing
- pathophysiological changes that affect fluid balance in disease states
- indications for IV fluid therapy
- reasons for the choice of the various fluids available and
- principles of assessing fluid balance.

In developing the guideline, it was necessary to limit the scope by excluding patient groups with more specialised fluid prescribing needs. It is important to emphasise that the recommendations do not apply to patients under 16 years, pregnant women, and those with severe liver or renal disease, diabetes or burns. They also do not apply to patients needing inotropes and those on intensive monitoring, and so they have less relevance to intensive care settings and patients during surgical anaesthesia. Patients with traumatic brain injury (including patients needing neurosurgery) are also excluded. The scope of the guideline does not cover the practical aspects of administration (as opposed to the prescription) of IV fluids.

It is hoped that this guideline will lead to better fluid prescribing in hospitalised patients, reduce morbidity and mortality, and lead to better patient outcomes.

Strategies for further research into the subject have also been proposed.

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

Key priorities for implementation

The following recommendations have been identified as priorities for implementation.

Principles and protocols for intravenous fluid therapy

- When prescribing intravenous (IV) fluids, remember the 5 Rs: Resuscitation, Routine maintenance, Replacement, Redistribution and Reassessment.
- Offer IV fluid therapy as part of a protocol (see the <u>algorithms for IV fluid therapy</u>):
 - Assess patients' fluid and electrolyte needs following algorithm 1: assessment.
 - If patients need IV fluids for fluid resuscitation, follow algorithm 2: fluid resuscitation.
 - If patients need IV fluids for routine maintenance, follow algorithm 3: routine maintenance.
 - If patients need IV fluids to address existing deficits or excesses, ongoing abnormal losses or abnormal fluid distribution, follow algorithm 4: replacement and redistribution.
- Patients should have an IV fluid management plan, which should include details of:
 - the fluid and electrolyte prescription over the next 24 hours
 - the assessment and monitoring plan.

Initially, the IV fluid management plan should be reviewed by an <u>expert</u> daily. IV fluid management plans for patients on longer-term IV fluid therapy whose condition is stable may be reviewed less frequently.

Assessment and monitoring

• Assess the patient's likely fluid and electrolyte needs from their history, clinical examination, current medications, clinical monitoring and laboratory investigations:

- History should include any previous limited intake, thirst, the quantity and composition of abnormal losses (see the <u>diagram of ongoing losses</u>), and any comorbidities, including patients who are malnourished and at risk of refeeding syndrome (see the <u>NICE guideline on nutrition support for adults</u>).
- Clinical examination should include an assessment of the patient's fluid status, including:
 - \diamond pulse, blood pressure, capillary refill and jugular venous pressure
 - \diamond presence of pulmonary or peripheral oedema
 - \Diamond presence of postural hypotension.
- Clinical monitoring should include current status and trends in:
 - ◇ National Early Warning Score (NEWS)
 - \diamondsuit fluid balance charts
 - \diamond weight.
- Laboratory investigations should include current status and trends in:
 - \diamondsuit full blood count
 - \diamond urea, creatinine and electrolytes.
- All patients continuing to receive IV fluids need regular monitoring. This should initially include at least daily reassessments of clinical fluid status, laboratory values (urea, creatinine and electrolytes) and fluid balance charts, along with weight measurement twice weekly. Be aware that:
 - Patients receiving IV fluid therapy to address replacement or redistribution problems may need more frequent monitoring.
 - Additional monitoring of urinary sodium may be helpful in patients with high-volume gastrointestinal losses. (Reduced urinary sodium excretion [less than 30 mmol/l] may indicate total body sodium depletion even if plasma sodium levels are normal. Urinary sodium may also indicate the cause of hyponatraemia, and guide the achievement of a negative sodium balance in patients with oedema. However, urinary sodium values may be misleading in the presence of renal impairment or diuretic therapy.)

- Patients on longer-term IV fluid therapy whose condition is stable may be monitored less frequently, although decisions to reduce monitoring frequency should be detailed in their IV fluid management plan.
- Clear incidents of fluid mismanagement (for example, unnecessarily prolonged dehydration or inadvertent fluid overload due to IV fluid therapy) should be reported through standard critical incident reporting to encourage improved training and practice (see the <u>table on consequences of fluid mismanagement to be reported as</u> <u>critical incidents</u>).

Resuscitation

 If patients need IV fluid resuscitation, use crystalloids that contain sodium in the range 130 to 154 mmol/l, with a bolus of 500 ml over less than 15 minutes. (For more information, see the <u>table on composition of commonly used crystalloids on the</u> <u>guideline's tools and resources page</u>.)

Routine maintenance

- If patients need IV fluids for routine maintenance alone, restrict the initial prescription to:
 - 25 to 30 ml/kg/day of water and
 - approximately 1 mmol/kg/day of potassium, sodium and chloride **and**
 - approximately 50 to 100 g/day of glucose to limit starvation ketosis. (This quantity will not address patients' nutritional needs; see the <u>NICE guideline on nutrition</u> <u>support in adults</u>.)

Weight-based potassium prescriptions should be rounded to the nearest common fluids available (for example, a 67 kg person should have fluids containing 20 mmol and 40 mmol of potassium in a 24-hour period). Potassium should not be added to IV fluid bags as this is dangerous. For more information, see the <u>table on</u> <u>IV fluid prescription for routine maintenance over a 24-hour period</u>.

Training and education

- Hospitals should establish systems to ensure that all healthcare professionals involved in prescribing and delivering IV fluid therapy are trained on the principles covered in this guideline, and are then formally assessed and reassessed at regular intervals to demonstrate competence in:
 - understanding the physiology of fluid and electrolyte balance in patients with normal physiology and during illness
 - assessing patients' fluid and electrolyte needs (the 5 Rs: Resuscitation, Routine maintenance, Replacement, Redistribution and Reassessment)
 - assessing the risks, benefits and harms of IV fluids
 - prescribing and administering IV fluids
 - monitoring the patient response
 - evaluating and documenting changes and
 - taking appropriate action as required.
- Hospitals should have an IV fluids lead, responsible for training, clinical governance, audit and review of IV fluid prescribing and patient outcomes.

Recommendations

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

<u>Making decisions using NICE guidelines</u> 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.

1.1 Principles and protocols for intravenous fluid therapy

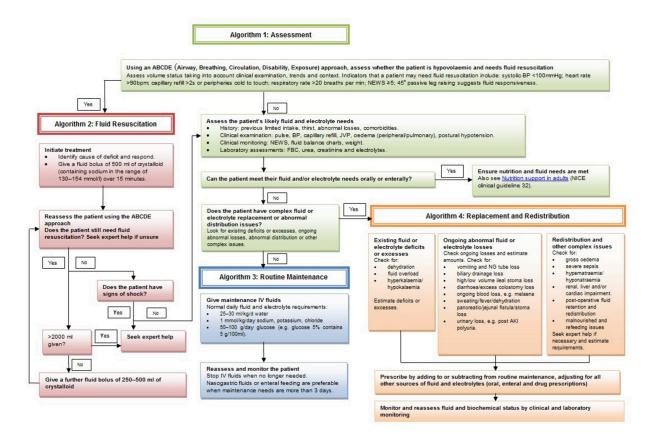
The assessment and management of patients' fluid and electrolyte needs is fundamental to good patient care.

- 1.1.1 Assess and manage patients' fluid and electrolyte needs as part of every ward review. Provide intravenous (IV) fluid therapy only for patients whose needs cannot be met by oral or enteral routes, and stop as soon as possible.
- 1.1.2 Skilled and competent healthcare professionals should prescribe and administer IV fluids, and assess and monitor patients receiving IV fluids (see the recommendations in the section on training and education).
- 1.1.3 When prescribing IV fluids, remember the 5 Rs: Resuscitation, Routine maintenance, Replacement, Redistribution and Reassessment.
- 1.1.4 Offer IV fluid therapy as part of a protocol (see the algorithms for IV fluid therapy):
 - Assess patients' fluid and electrolyte needs following algorithm 1: assessment.

- If patients need IV fluids for fluid resuscitation, follow algorithm 2: fluid resuscitation.
- If patients need IV fluids for routine maintenance, follow algorithm 3: routine maintenance.
- If patients need IV fluids to address existing deficits or excesses, ongoing abnormal losses or abnormal fluid distribution, follow algorithm 4: replacement and redistribution.

Algorithms for IV fluid therapy

See the downloadable PDF version of the algorithm poster set.



- 1.1.5 Include the following information in IV fluid prescriptions:
 - the type of fluid to be administered
 - the rate and volume of fluid to be administered.

- 1.1.6 Patients should have an IV fluid management plan, which should include details of:
 - the fluid and electrolyte prescription over the next 24 hours
 - the assessment and monitoring plan.

Initially, the IV fluid management plan should be reviewed by an <u>expert</u> daily. IV fluid management plans for patients on longer-term IV fluid therapy whose condition is stable may be reviewed less frequently.

- 1.1.7 When prescribing IV fluids and electrolytes, take into account all other sources of fluid and electrolyte intake, including any oral or enteral intake, and intake from drugs, IV nutrition, blood and blood products.
- 1.1.8 Patients have a valuable contribution to make to their fluid balance. If a patient needs IV fluids, explain the decision, and discuss the signs and symptoms they need to look out for if their fluid balance needs adjusting. If possible or when asked, provide written information (for example, <u>NICE's information for the public</u>), and involve the patient's family members or carers (as appropriate).

1.2 Assessment and monitoring

Initial assessment

1.2.1 Assess whether the patient is hypovolaemic. Indicators that a patient may need urgent fluid resuscitation include:

- systolic blood pressure is less than 100 mmHg
- heart rate is more than 90 beats per minute
- capillary refill time is more than 2 seconds, or peripheries are cold to touch
- respiratory rate is more than 20 breaths per minute
- National Early Warning Score (NEWS) is 5 or more

• passive leg raising suggests fluid responsiveness.

Passive leg raising is a bedside method to assess fluid responsiveness in a patient. It is best undertaken with the patient initially semi-recumbent and then tilting the entire bed through 45°. Alternatively, it can be done by lying the patient flat and passively raising their legs to greater than 45°. If, at 30 to 90 seconds, the patient shows signs of haemodynamic improvement, it indicates that volume replacement may be required. If the condition of the patient deteriorates, in particular breathlessness, it indicates that the patient may be fluid overloaded.

- 1.2.2 Assess the patient's likely fluid and electrolyte needs from their history, clinical examination, current medications, clinical monitoring and laboratory investigations:
 - History should include any previous limited intake, thirst, the quantity and composition of abnormal losses (see the <u>diagram of ongoing losses</u>), and any comorbidities, including patients who are malnourished and at risk of refeeding syndrome (see the <u>NICE guideline on nutrition support in adults</u>).
 - Clinical examination should include an assessment of the patient's fluid status, including:
 - pulse, blood pressure, capillary refill and jugular venous pressure
 - presence of pulmonary or peripheral oedema
 - presence of postural hypotension.
 - Clinical monitoring should include current status and trends in:
 - NEWS
 - fluid balance charts
 - weight.
 - Laboratory investigations should include current status and trends in:
 - full blood count

- urea, creatinine and electrolytes.

Reassessment

- 1.2.3 If patients are receiving IV fluids for resuscitation, reassess the patient using the ABCDE approach (Airway, Breathing, Circulation, Disability, Exposure), monitor their respiratory rate, pulse, blood pressure and perfusion continuously, and measure their venous lactate levels and/or arterial pH and base excess according to guidance on advanced life support (Resuscitation Council UK, 2011).
- 1.2.4 All patients continuing to receive IV fluids need regular monitoring. This should initially include at least daily reassessments of clinical fluid status, laboratory values (urea, creatinine and electrolytes) and fluid balance charts, along with weight measurement twice weekly. Be aware that:
 - Patients receiving IV fluid therapy to address replacement or redistribution problems may need more frequent monitoring.
 - Additional monitoring of urinary sodium may be helpful in patients with high-volume gastrointestinal losses. (Reduced urinary sodium excretion [less than 30 mmol/l] may indicate total body sodium depletion even if plasma sodium levels are normal. Urinary sodium may also indicate the cause of hyponatraemia, and guide the achievement of a negative sodium balance in patients with oedema. However, urinary sodium values may be misleading in the presence of renal impairment or diuretic therapy.)
 - Patients on longer-term IV fluid therapy whose condition is stable may be monitored less frequently, although decisions to reduce monitoring frequency should be detailed in their IV fluid management plan.
- 1.2.5 If patients have received IV fluids containing chloride concentrations greater than 120 mmol/l (for example, sodium chloride 0.9%), monitor their serum chloride concentration daily. If patients develop hyperchloraemia or acidaemia, reassess their IV fluid prescription and assess their acid–base status. Consider less frequent monitoring for patients who are stable.
- 1.2.6 Clear incidents of fluid mismanagement (for example, unnecessarily prolonged

dehydration or inadvertent fluid overload due to IV fluid therapy) should be reported through standard critical incident reporting to encourage improved training and practice (see the <u>table on consequences of fluid mismanagement to</u> <u>be reported as critical incidents</u>).

1.2.7 If patients are transferred to a different location, reassess their fluid status and IV fluid management plan on arrival in the new setting.

1.3 Resuscitation

- 1.3.1 If patients need IV fluid resuscitation, use crystalloids that contain sodium in the range 130 to 154 mmol/I, with a bolus of 500 ml over less than 15 minutes. (For more information, see the <u>table on composition of commonly used crystalloids on the guideline's tools and resources page</u>.)
- 1.3.2 Do not use tetrastarch for fluid resuscitation.
- 1.3.3 Consider human albumin solution 4% to 5% for fluid resuscitation only in patients with severe sepsis.

1.4 Routine maintenance

- 1.4.1 If patients need IV fluids for routine maintenance alone, restrict the initial prescription to:
 - 25 to 30 ml/kg/day of water and
 - approximately 1 mmol/kg/day of potassium, sodium and chloride and
 - approximately 50 to 100 g/day of glucose to limit starvation ketosis. (This quantity will not address patients' nutritional needs; see the <u>NICE guideline</u> on nutrition support in adults.)

Weight-based potassium prescriptions should be rounded to the nearest common fluids available (for example, a 67 kg person should have fluids

containing 20 mmol and 40 mmol of potassium in a 24-hour period). Potassium should not be added to IV fluid bags as this is dangerous. For more information, see the <u>table on IV fluid prescription for routine</u> <u>maintenance over a 24-hour period</u>.

- 1.4.2 For patients who are obese, adjust the IV fluid prescription to their ideal body weight. Use lower range volumes per kg (patients rarely need more than a total of 3 litres of fluid per day) and seek <u>expert</u> help if their body mass index (BMI) is more than 40 kg/m².
- 1.4.3 Consider prescribing less fluid (for example, 20 to 25 ml/kg/day fluid) for patients who:
 - are older or frail
 - have renal impairment or cardiac failure
 - are malnourished and at risk of refeeding syndrome (see the <u>NICE guideline</u> <u>on nutrition support in adults</u>).
- 1.4.4 When prescribing for routine maintenance alone, consider using 25 to 30 ml/kg/ day sodium chloride 0.18% in 4% glucose with 27 mmol/l potassium on day 1 (there are other regimens to achieve this). Prescribing more than 2.5 litres per day increases the risk of hyponatraemia. These are initial prescriptions and further prescriptions should be guided by monitoring.

Weight-based potassium prescriptions should be rounded to the nearest common fluids available (for example, a 67 kg person should have fluids containing 20 mmol and 40 mmol of potassium in a 24-hour period). Potassium should not be added to IV fluid bags as this is dangerous.

1.4.5 Consider delivering IV fluids for routine maintenance during daytime hours to promote sleep and wellbeing.

1.5 Replacement and redistribution

- 1.5.1 Adjust the IV prescription (add to or subtract from maintenance needs) to account for existing fluid and/or electrolyte deficits or excesses, ongoing losses (see the <u>diagram of ongoing losses</u>) or abnormal distribution.
- 1.5.2 Seek <u>expert</u> help if patients have a complex fluid and/or electrolyte redistribution issue or imbalance, or significant comorbidity, for example:
 - gross oedema
 - severe sepsis
 - hyponatraemia or hypernatraemia
 - renal, liver and/or cardiac impairment
 - post-operative fluid retention and redistribution
 - malnourished and refeeding issues (see the <u>NICE guideline on nutrition</u> <u>support in adults</u>).

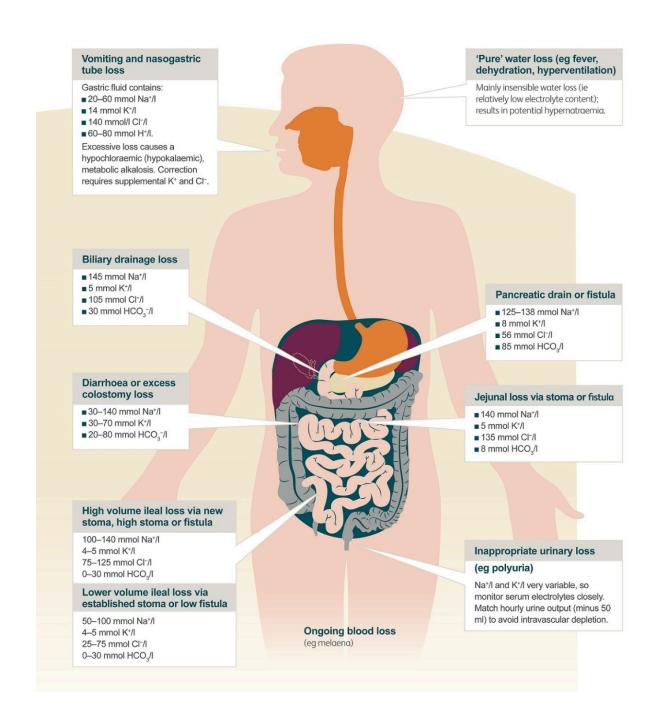
1.6 Training and education

- 1.6.1 Hospitals should establish systems to ensure that all healthcare professionals involved in prescribing and delivering IV fluid therapy are trained on the principles covered in this guideline, and are then formally assessed and reassessed at regular intervals to demonstrate competence in:
 - understanding the physiology of fluid and electrolyte balance in patients with normal physiology and during illness
 - assessing patients' fluid and electrolyte needs (the 5 Rs: Resuscitation, Routine maintenance, Replacement, Redistribution and Reassessment)
 - assessing the risks, benefits and harms of IV fluids
 - prescribing and administering IV fluids

- monitoring the patient response
- evaluating and documenting changes and
- taking appropriate action as required.
- 1.6.2 Healthcare professionals should receive training and education about, and be competent in, recognising, assessing and preventing consequences of mismanaged IV fluid therapy, including:
 - pulmonary oedema
 - peripheral oedema
 - volume depletion and shock.
- 1.6.3 Hospitals should have an IV fluids lead, responsible for training, clinical governance, audit and review of IV fluid prescribing and patient outcomes.

Diagram of ongoing losses

See the downloadable PDF version of the diagram of ongoing losses.



Source: Copyright – National Clinical Guideline Centre

Consequence of fluid mismanagement	Identifying features	Time frame of identification
Hypovolaemia	 Patient's fluid needs not met by oral, enteral or intravenous (IV) intake and: Features of dehydration on clinical examination Low urine output or concentrated urine Biochemical indicators, such as more than 50% increase in urea or creatinine with no other identifiable cause 	Before and during IV fluid therapy
Pulmonary oedema (breathlessness during infusion)	 No other obvious cause identified (for example, pneumonia, pulmonary embolus or asthma) Features of pulmonary oedema on clinical examination Features of pulmonary oedema on X ray 	During IV fluid therapy or within 6 hours of stopping IV fluids
Hyponatraemia	 Serum sodium less than 130 mmol/l No other likely cause of hyponatraemia identified 	During IV fluid therapy or within 24 hours of stopping IV fluids
 Serum sodium 155 mmol/l or more Baseline sodium normal or low IV fluid regimen included 0.9% sodium chloride No other likely cause of hypernatraemia identified 		During IV fluid therapy or within 24 hours of stopping IV fluids

Table 1 Consequences of fluid mismanagement to be reported as critical incidents

Consequence of fluid mismanagement	Identifying features	Time frame of identification
Peripheral oedema	 Pitting oedema in extremities and/or lumbar sacral area No other obvious cause identified (for example, nephrotic syndrome or known cardiac failure) 	During IV fluid therapy or within 24 hours of stopping IV fluids
Hyperkalaemia	 Serum potassium more than 5.5 mmol/l No other obvious cause identified 	During IV fluid therapy or within 24 hours of stopping IV fluids
Hypokalaemia	 Serum potassium less than 3.0 mmol/l likely to be due to infusion of fluids without adequate potassium provision No other obvious cause (for example, potassium-wasting diuretics, refeeding syndrome) 	During IV fluid therapy or within 24 hours of stopping IV fluids

Weight-based potassium prescriptions should be rounded to the nearest common fluids available (for example, a 67 kg person should have fluids containing 20 mmol and 40 mmol of potassium in a 24-hour period). Potassium should not be added to IV fluid bags as this is dangerous.

Source: Table 1 was drafted based on the consensus decision of the members of the guideline development group.

Body weight (in kg)	Water (25 to 30 ml/ kg/day)	Sodium, chloride, potassium (approximately 1 mmol/kg/day of each)
40	1000 to 1200	40
41	1025 to 1230	41
42	1050 to 1260	42

43 1075 to 1290 43 44 1100 to 1320 44 45 1125 to 1350 45 46 1150 to 1380 46 47 1175 to 1410 47 48 1200 to 1440 48 49 1225 to 1470 49 50 1250 to 1500 50 51 1275 to 1530 51 52 1300 to 1560 52 53 1325 to 1590 53 54 1350 to 1620 54 55 1375 to 1650 55 56 1400 to 1680 56 57 1425 to 1710 57 58 1450 to 1740 58 59 1475 to 1770 59 60 1500 to 1800 60 61 1525 to 1830 61 62 1550 to 1860 62 63 1575 to 1890 63 64 1600 to 1920 64 65 1625 to 1950 65 <th></th> <th></th> <th></th>			
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60 1500 to 1800 60 61 1525 to 1830 61 62 1550 to 1860 62 63 1575 to 1890 63 64 1600 to 1920 64 65 1625 to 1950 65 66 1650 to 1980 66 67 1675 to 2010 67	58	1450 to 1740	58
61 1525 to 1830 61 62 1550 to 1860 62 63 1575 to 1890 63 64 1600 to 1920 64 65 1625 to 1950 65 66 1650 to 1980 66 67 1675 to 2010 67	59	1475 to 1770	59
62 1550 to 1860 62 63 1575 to 1890 63 64 1600 to 1920 64 65 1625 to 1950 65 66 1650 to 1980 66 67 1675 to 2010 67	60	1500 to 1800	60
63 1575 to 1890 63 64 1600 to 1920 64 65 1625 to 1950 65 66 1650 to 1980 66 67 1675 to 2010 67	61	1525 to 1830	61
64 1600 to 1920 64 65 1625 to 1950 65 66 1650 to 1980 66 67 1675 to 2010 67	62	1550 to 1860	62
65 1625 to 1950 65 66 1650 to 1980 66 67 1675 to 2010 67	63	1575 to 1890	63
66 1650 to 1980 66 67 1675 to 2010 67	64	1600 to 1920	64
67 1675 to 2010 67	65	1625 to 1950	65
	66	1650 to 1980	66
68 1700 to 2040 68	67	1675 to 2010	67
	68	1700 to 2040	68

69	1725 to 2070	69
70	1750–2100	70
71	1775 to 2130	71
72	1800 to 2160	72
73	1825 to 2190	73
74	1850 to 2220	74
75	1875 to 2250	75
76	1900 to 2280	76
77	1925 to 2310	77
78	1950 to 2340	78
79	1975 to 2370	79
80	2000 to 2400	80
81	2025 to 2430	81
82	2050 to 2460	82
83	2075 to 2490	83
84	2100 to 2520	84
85	2125 to 2550	85
86	2150 to 2580	86
87	2175 to 2610	87
88	2200 to 2640	88
89	2225 to 2670	89
90	2250 to 2700	90
91	2275 to 2730	91
92	2300 to 2760	92
93	2325 to 2790	93
94	2350 to 2820	94

95	2375 to 2850	95
96	2400 to 2880	96
97	2425 to 2910	97
98	2450 to 2940	98
99	2475 to 2970	99
100	2500 to 3000	100
>100	2500 to 3000	100

Add 50 to 100 grams/day glucose (for example, glucose 5% contains 5 g/100 ml).

For special considerations, refer to the recommendations for routine maintenance.

Weight-based potassium prescriptions should be rounded to the nearest common fluids available (for example, a 67 kg person should have fluids containing 20 mmol and 40 mmol of potassium in a 24-hour period). Potassium should not be added to IV fluid bags as this is dangerous.

Source: Table 2 was drafted based on the consensus decision of the members of the guideline development group.

Terms used in this guideline

Expert

In this guideline, the term 'expert' refers to a healthcare professional who has core competencies to diagnose and manage acute illness. These competencies can be delivered by a variety of models at a local level, such as a critical care outreach team, a hospital-at-night team or a specialist trainee in an acute medical or surgical specialty. For more information, see the <u>NICE guideline on acutely ill patients in hospital</u>.

Recommendations for research

The guideline development group has made this recommendation for research.

1 Assessment and monitoring

What is the incidence of complications during, and as a consequence of, intravenous fluid therapy?

Why this is important

This is almost certainly under-reported in the ward setting with significant implications for patients, predominantly morbidity through to mortality. It is probable that complications of fluid therapy are frequent and may be associated with increased clinical needs, such as critical care and, on occasion, may necessitate fluid resuscitation. Lack of a set of clearly defined features of the complications of fluid mismanagement compounds the problem. It is important to define these features and then undertake an observational study in a hospital setting to determine the epidemiology of these complications. Such a study would highlight the prevalence of fluid related complications and inform the development of preventive measures.

Finding more information and committee details

To find NICE guidance on related topics, including guidance in development, see the <u>NICE</u> topic page on medicines management.

For full details of the evidence and the guideline committee's discussion, see the <u>full</u> <u>guideline</u>. You can also find information about <u>how the guideline was developed</u>, including <u>details of the committee</u>.

NICE has produced <u>tools and resources to help you put this guideline into practice</u>. For general help and advice on putting our guidelines into practice, see <u>resources to help you</u> <u>put NICE guidance into practice</u>.

Update information

May 2017: Some research recommendations that have become outdated since original publication have been stood down and deleted.

December 2016: We added information about weight-based potassium prescriptions and clarified that potassium should not be added directly to intravenous fluid bags (as this is dangerous) in recommendations 1.4.1 and 1.4.4, tables 1 and 2 and the accompanying algorithms giving more information on weight-based potassium prescriptions.

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