

Integrated care pathway for the management of children and young people with

# Diabetic Ketoacidosis

If you are not experienced in managing children in DKA, ask for senior help now.



Affix sticker or complete patient demographics below

Name
Date of Birth
Hospital / NHS Number

**DKA protocol started at:**

hh:mm
dd/mm/yyyy



#### **IMPORTANT SAFETY NOTES:**

These are general guidelines for management. Treatment may need modification to suit the individual patient and these guidelines do not remove the need for frequent detailed reassessments of the individual patient's requirements and specific treatment tailored to those requirements.

This integrated care pathway (ICP) is designed to be used by, or under the supervision of, clinicians experienced in the management of paediatric DKA. It should be used in conjunction with the full BSPED DKA 2021 guideline on which it is based which can be found at: <https://www.bsped.org.uk/clinical-resources/bsped-dka-guidelines/>

This is part of the official patient care record and should be filed in the patient's notes. All professionals involved must document any intervention carried out. When filling out a flow chart, you must complete the box in the lower right corner of the chart with your signature, name, and the date and time. Any variation from the care plan must be documented.

[\*\*dka-calculator.co.uk\*\*](http://dka-calculator.co.uk)

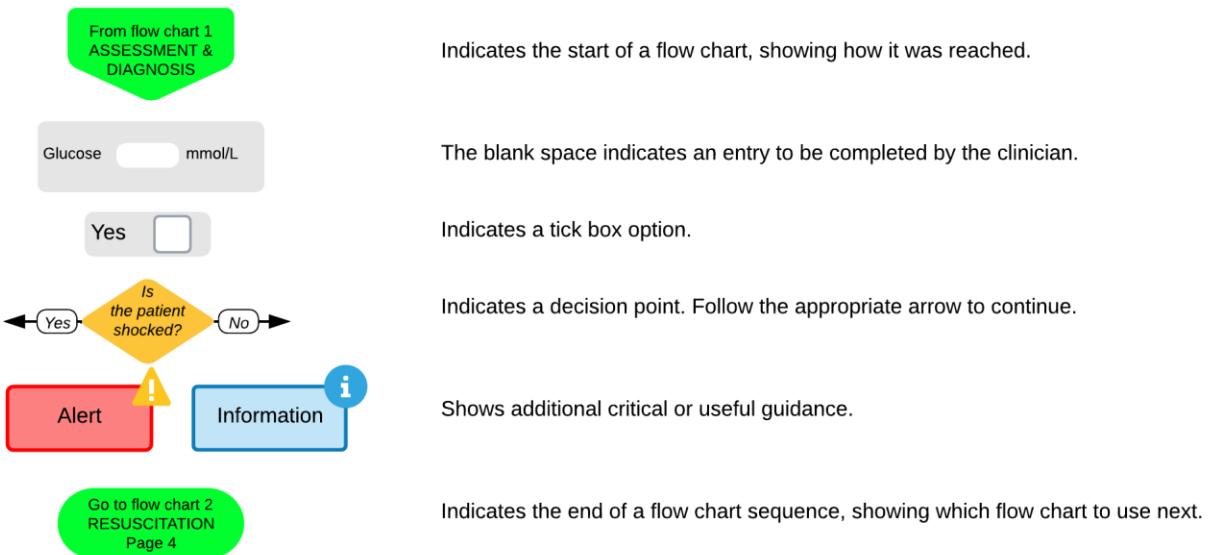
This ICP is designed in conjunction with an online calculator that will pre-fill elements, for example patient demographics and fluid calculations. While the ICP can be used without this step, use of the calculator is strongly advised as it reduces the risk of calculation errors. The calculator is also important for the national DKA audit programme. No patient identifiable data is transmitted or stored when using the online calculator. Access the calculator at the web address above.



## INTRODUCTORY NOTES

This ICP is designed to be worked through and completed to aid with management decisions and to record important events. You should start with flow chart 1 - ASSESSMENT & DIAGNOSIS - on page 3, and proceed as shown in the guidance below. Remember to refer to the additional guidance in the appendices if you are not already familiar with it.

The flow charts are structured in a systematic way as follows:



The ICP is divided into sections which are identified by coloured borders at the side of each page.

### MAIN PROTOCOL SECTION

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- Page 5 – Flow Chart 3 – SECONDARY REVIEW
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### COMPLICATIONS SECTION

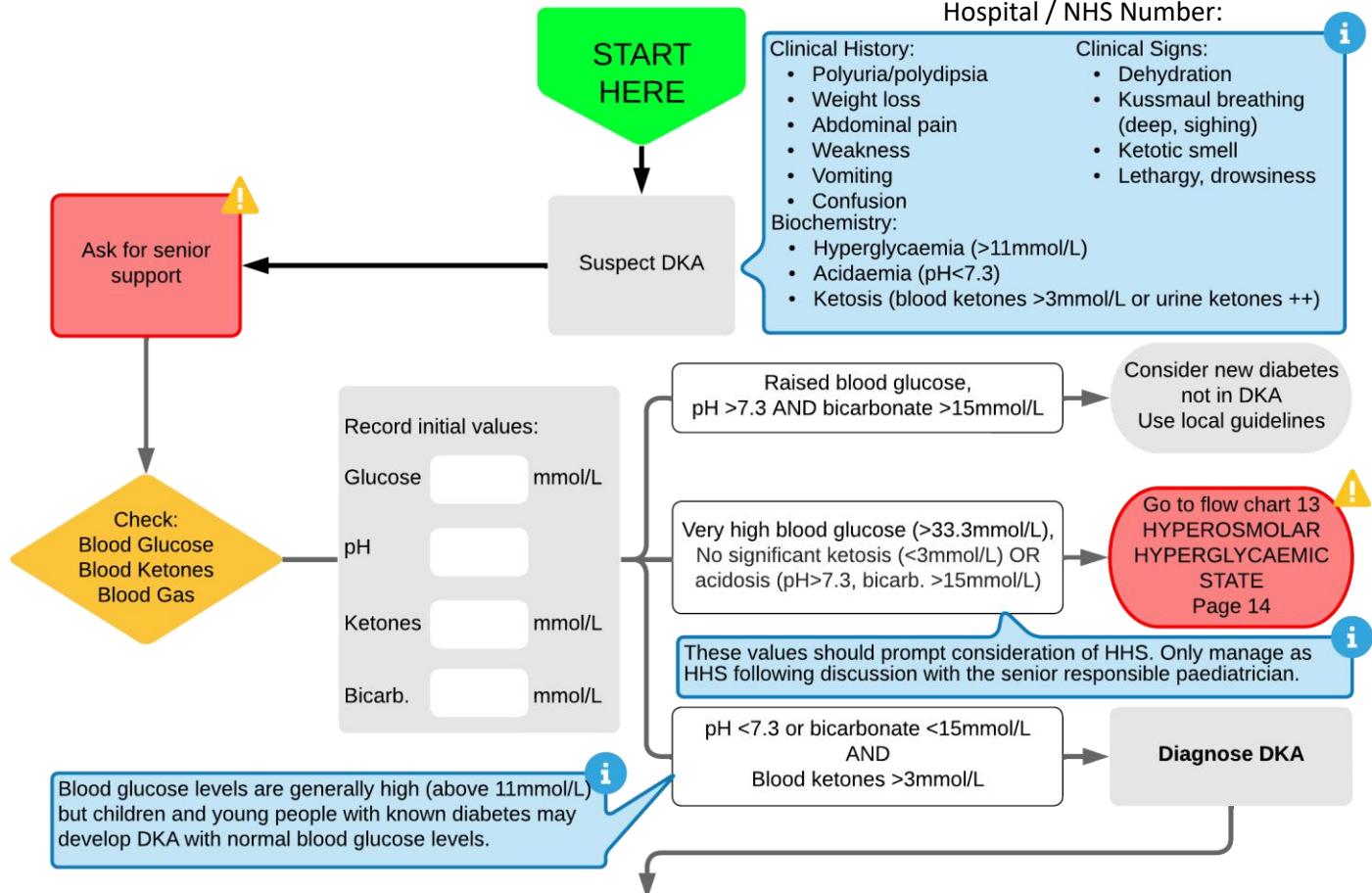
- Page 12 – Flow Chart 9 – CEREBRAL OEDEMA
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- Page 16 – Appendix 3 – MAKING UP IV FLUIDS
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## FLOW CHART 1 – ASSESSMENT &amp; DIAGNOSIS

Patient Name: \_\_\_\_\_  
 Date of Birth: \_\_\_\_\_  
 Hospital / NHS Number: \_\_\_\_\_

**Perform rapid emergency assessment**

Record your initial assessment here and use this to guide your management on the following page

<b>Airway</b>	Maintaining own airway? Yes <input type="checkbox"/> No <input type="checkbox"/>	<b>Breathing</b>	RR <input type="text"/> /min SpO <sub>2</sub> <input type="text"/> %	Acidotic pattern? Yes <input type="checkbox"/> No <input type="checkbox"/>
<b>Circulation</b>	HR <input type="text"/> /min CRT <input type="text"/> secs BP <input type="text"/> / <input type="text"/>	Clinically shocked? Yes <input type="checkbox"/> No <input type="checkbox"/>	<b>Disability</b>	GCS <input type="text"/> /15 M: <input type="text"/> /6 V: <input type="text"/> /5 E: <input type="text"/> /4 See also appendix 1 page 15 <span style="color: blue;">i</span>

Further details:

Children who are alert, not clinically dehydrated, not nauseated or vomiting, do not always require IV fluids, even if their ketone levels are high. They usually tolerate oral rehydration and subcutaneous insulin but do require monitoring regularly to ensure that they are improving and their ketone levels are falling. This decision should be made in consultation with the responsible paediatrician.

**Go to flow chart 2 RESUSCITATION Page 4**

Chart completed by: \_\_\_\_\_  
 GMC number: \_\_\_\_\_  
 Signature: \_\_\_\_\_  
 Time / Date: \_\_\_\_\_

## FLOW CHART 2 – RESUSCITATION

Patient Name:  
Date of Birth:  
Hospital / NHS Number:

From flow chart 1  
ASSESSMENT &  
DIAGNOSIS

**A:** Establish airway: Seek urgent anaesthetic review if unable to protect airway.  
If child comatose: Insert NG tube on free drainage.

**B:** Give O<sub>2</sub> 100% via face mask with reservoir bag (only omit if child very well).

**C:** Establish IV access (consider 2<sup>nd</sup> cannula for later blood samples), take bloods (see box).  
Commence cardiac monitoring (peaked T waves may indicate hyperkalaemia).

For estimated weight:  
• Refer to appendix 2, page 15  
• Ensure an accurate weight is obtained before starting maintenance fluids

Weight:  kg

Actual / Estimated / Recent

Recommended bloods:  
• Blood ketones  
• Blood gas  
• HbA1c  
• FBC, U+Es, CRP  
• Lab glucose

For patients newly diagnosed:  
• TFTs  
• TTG  
• Additional bloods as per your local policy

Shocked patients should be discussed with the most senior paediatrician or intensivist at the earliest opportunity.

Is the patient shocked?

• Tachycardia  
• Prolonged central capillary refill  
• Poor peripheral pulses  
• Hypotension (late sign)

Yes

No

Shocked patients: 10 ml/kg bolus of 0.9% saline or plasmalyte over 15 minutes

Volume:  ml

Started: hh:mm dd/mm/yyyy

Reassess: if still shocked further boluses of 10ml/kg (up to total of 40ml/kg) may be given

Volume:  ml hh:mm dd/mm/yyyy

Volume:  ml hh:mm dd/mm/yyyy

Volume:  ml hh:mm dd/mm/yyyy

If still shocked consider inotropes and critical care escalation

All non-shocked children with mild, moderate or severe DKA should receive a 10ml/kg bolus of 0.9% saline over 30 minutes

Volume:  ml

Started: hh:mm dd/mm/yyyy

Whilst excessive fluid should be avoided because of the risk of cerebral oedema, it is important to ensure that the circulation is adequate and fluid should be given to support this. Cerebral perfusion is dependent on both perfusion pressure **and** intracranial pressure, and hypotension will exacerbate the risk of brain injury.

A bolus given on this arm is later subtracted from the calculated fluid deficit, whereas boluses for shocked patients are not. See page 6 for details.

Do NOT give IV sodium bicarbonate to patients with DKA. See appendix 4, page 17, for more information.

Consider placing NG tube to reduce the risk of aspiration in patients with reduced conscious level

**D:** Consider if cerebral oedema may be present

Early manifestations: headache, agitation/irritability, unexpected fall in heart rate, rise in blood pressure  
Additional manifestations: deterioration in conscious level, abnormal breathing pattern, oculomotor palsies, abnormal posturing, pupil inequalities or dilatation

Go to flow chart 3  
SECONDARY REVIEW  
Page 5

Features of cerebral oedema?

Go to flow chart 9  
CEREBRAL OEDEMA  
Page 12

Chart completed by: \_\_\_\_\_  
GMC number: \_\_\_\_\_  
Signature: \_\_\_\_\_  
Time / Date: \_\_\_\_\_

## FLOW CHART 3 – SECONDARY REVIEW

From flow chart 2  
RESUSCITATIONPatient Name:  
Date of Birth:  
Hospital / NHS Number:**History:**

Consider features including:

- Polyuria/polydipsia/wetting
- Weight loss
- Vomiting/abdominal pain
- Headache
- Recent infection

**Past medical history:**

If pre-existing diabetes ask about previous DKA episodes.

**Drug history:**

If pre-existing diabetes include usual insulin regimen details, adherence.

**Allergies:****Family and social history:**

Ask about family history of diabetes, thyroid disease, coeliac disease and other auto-immune conditions.

**Examination:**

Including general status, cardiovascular, abdomen, respiratory/ENT, neurology... Consider signs as shown on ASSESSMENT & DIAGNOSIS flow chart 1

DKA may be precipitated by sepsis or intercurrent infection, and fever is not part of DKA. Infection may co-exist with DKA. Suspect sepsis if there is fever or hypothermia, hypotension, refractory acidosis or lactic acidosis. A high lactate should increase concern about possible infection or sepsis.

Go to flow chart 4  
FLUIDS  
Page 6

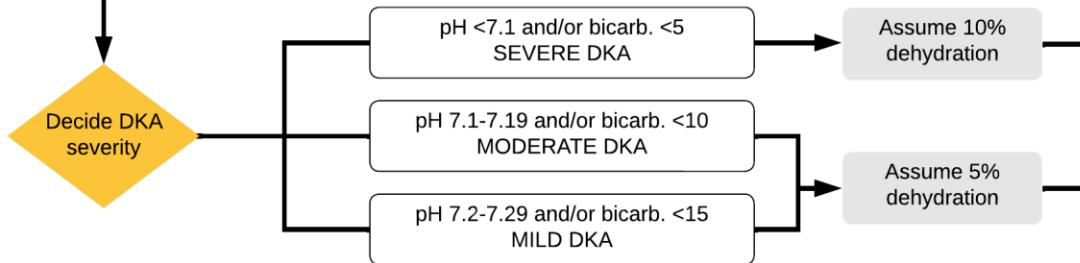
Chart completed by: \_\_\_\_\_  
GMC number: \_\_\_\_\_  
Signature: \_\_\_\_\_  
Time / Date: \_\_\_\_\_

## FLOW CHART 4 - FLUIDS

Patient Name: \_\_\_\_\_  
 Date of Birth: \_\_\_\_\_  
 Hospital / NHS Number: \_\_\_\_\_

From flow chart 3  
 SECONDARY  
 REVIEW

To avoid excessive amounts of fluid in overweight and obese children it is recommended that consideration be given to using a **maximum weight of 75kg or 98<sup>th</sup> centile weight for age** (whichever is lower) when calculating both deficit and maintenance requirements. Please refer to the full BSPED guidelines for further information.



## Fluid calculations

$$\text{Fluid deficit} = \boxed{\text{Patient weight}} \text{ kg} \times \boxed{\% \text{ Dehydration}} \times 10 = \boxed{\quad} \text{ mL}$$

e.g. 24kg × 5% × 10 = 1200mL

Subtract ONLY the 10mL/kg bolus given over 30 minutes to non-shocked patients.  
 DO NOT subtract rapid resuscitation boluses given to shocked patients.

$$\text{Fluid deficit} = \boxed{\text{Fluid deficit}} \text{ mL} - \boxed{10\text{mL/kg bolus volume}} \text{ mL} = \boxed{\quad} \text{ mL}$$

e.g. 1200mL - 240mL = 960mL

$$\text{Deficit replacement rate} = \boxed{\text{Fluid deficit (less bolus volume)}} \text{ mL} \div 48 \text{ hours} = \boxed{\quad} \text{ mL/hour}$$

e.g. 960mL ÷ 48hours = 20.0mL/hour

Use Holliday-Segar formula: i.e. 100mL/kg for first 10kg;  
 50mL/kg for next 10kg; 20mL/kg thereafter.

Note: deficit is replaced over 48 hours, maintenance rate is calculated over 24 hours.

$$\text{Maintenance rate} = \boxed{\text{Daily fluid requirement}} \text{ mL} \div 24 \text{ hours} = \boxed{\quad} \text{ mL/hour}$$

e.g. (for 24kg) (1000mL+500mL+80mL) ÷ 24hours = 65.8mL/hour

$$\text{STARTING FLUID RATE (after bolus complete)} = \boxed{\text{Maintenance rate}} \text{ mL/hour} + \boxed{\text{Deficit replacement rate}} \text{ mL/hour} = \boxed{\quad} \text{ mL/hour}$$

e.g. 65.8mL/hour + 20.0mL/hour = 85.8mL/hour

Plasmalyte 148 can be used as an alternative to 0.9% Sodium Chloride but must have added potassium.

If potassium is above normal range add potassium to fluids only after the patient has passed urine or after the Potassium has fallen to within the normal range.

Once initial bolus is complete:  
 Start 0.9% Sodium Chloride + 20mmol Potassium Chloride in 500mL at STARTING FLUID RATE as above

Fluid start time / date hh:mm dd/mm/yyyy

Go to flow chart 5  
 INSULIN  
 Page 7

Chart completed by: \_\_\_\_\_  
 GMC number: \_\_\_\_\_  
 Signature: \_\_\_\_\_  
 Time / Date: \_\_\_\_\_

## FLOW CHART 5 - INSULIN

Patient Name:  
Date of Birth:  
Hospital / NHS Number:

Starting insulin early may increase the risk of cerebral oedema

Wait for 1-2 hours after starting IV fluid treatment before starting insulin

If potassium <3.0mmol/L defer insulin until potassium >3.0mmol/L. If expected to cause significant delay discuss with intensivist regarding options including central access or oral potassium.

$$\text{Insulin hourly rate} = \boxed{0.05-0.1} \text{ Units/kg} \times \boxed{\text{Patient weight}} \text{ kg} = \boxed{\text{Units/hour}}$$

Starting insulin rates of 0.05 and 0.1 Units/kg/hr are typically suggested. A rate of 0.05 Units/kg/hr should be used in most cases unless your local policy dictates otherwise, or in specific cases (such as in adolescents or severe DKA) as directed by a senior paediatrician or intensivist.

Use pre-filled syringes containing 50 Units of soluble insulin in 50mL 0.9% Sodium Chloride where available. If pre-filled syringes are not available, add 50 Units of soluble insulin (e.g. Actrapid) to 49.5mL 0.9% Sodium Chloride.

Start at INSULIN HOURLY RATE as calculated above:

Insulin start time / date:

hh:mm dd/mm/yyyy

Pre-existing diabetes?

Yes

No

Patients on insulin pumps (CSII) should have their pump stopped once IV insulin is started.

Pump stopped?

Yes   
N/A

For patients already on long-acting insulin consider continuing at the usual dose and time throughout the DKA treatment, in addition to the IV insulin infusion, in order to shorten length of stay after recovery from DKA.

Long-acting insulin continued?

Yes   
No   
N/A

If supported by your local guidelines, consider starting an appropriate dose of long acting background insulin alongside the intravenous infusion.

Long-acting insulin started? Yes  No

Go to flow chart 6  
MONITORING & REVIEWS  
Page 8

Chart completed by:

GMC number:

Signature:

Time / Date:

## FLOW CHART 6 – MONITORING &amp; REVIEWS

From flow chart 5  
INSULINFrom flow chart 7  
ONGOING  
MANAGEMENT**Consider where the child or young person should be nursed:**

Patients with DKA should be cared for with one-to-one nursing if:

- they are younger than 2 years or
- they have severe DKA (blood pH below 7.1)

If one-to-one nursing cannot be provided on HDU/general paediatric ward, consider transfer to PICU.

Patient Name:

Date of Birth:

Hospital / NHS Number:

N.B. Where PICU or HDU do not exist within the admitting hospital, transfer to another hospital may not be appropriate (unless ventilatory support becomes necessary).

However, ALL children with DKA are high-dependency patients and require a high level of nursing care.

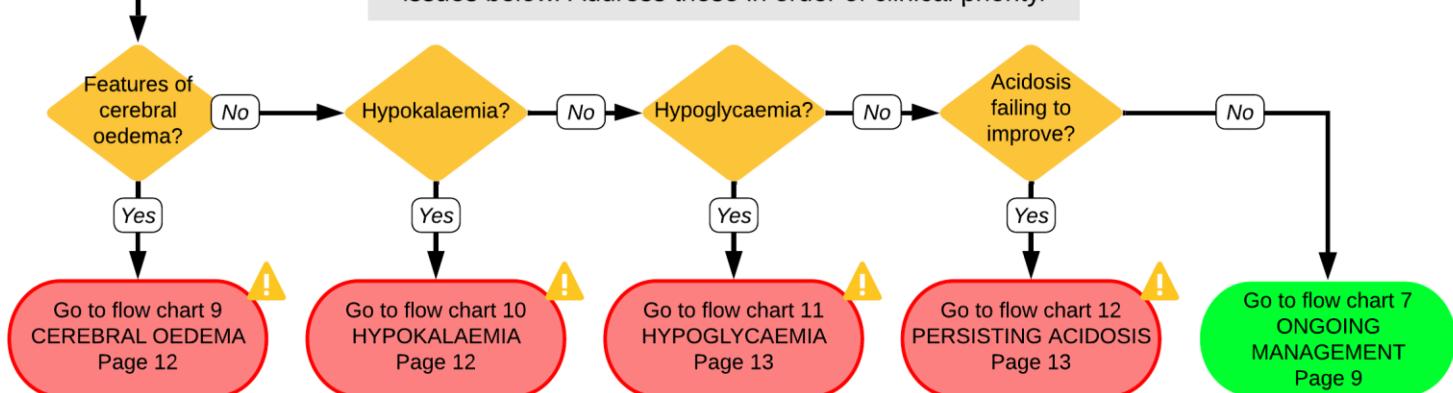
**Nursing Observations** - ensure full instructions are given to nurse responsible including:

- Strict fluid balance including oral fluids and urine output, using fluid balance charts (urinary catheterisation may be necessary in young/sick children)
- Hourly capillary blood glucose measurements
- Capillary blood ketone levels every 1-2 hours
- Hourly BP and basic observations
- Hourly level of consciousness initially, using the modified Glasgow Coma Score
- In children < 2 years of age and in those with a pH < 7.1 (at increased risk of cerebral oedema): Half-hourly neurological observations including the modified Glasgow Coma Score and heart rate
- Report immediately to medical staff:
  - symptoms of headache, or slowing of heart rate, or any change in either conscious level or behaviour
  - any changes in the ECG trace, especially signs of hypokalaemia, including ST-segment depression and prominent U-waves
- Twice daily weight; can be helpful in assessing fluid balance

**Medical Reviews**

- At 2 hours after starting treatment and then at least every 4 hours carry out and record the results of the following blood tests on the SERIAL DATA SHEET (page 11):
  - Glucose (Laboratory measurement)
  - Blood gas (for pH and pCO<sub>2</sub>)
  - Plasma U+Es - ensure samples are sent urgently to the lab
  - Blood ketones
- A doctor (or equivalent practitioner) should carry out a face-to-face review at the beginning of treatment, at 2 hours after starting treatment, and then at least every 4 hours and more frequently if:
  - child is aged under 2 years
  - has severe DKA (pH < 7.1)
  - there are any other reasons for special concern
- At each face-to-face review, provide an update on progress to the child or young person and their family and carers (as appropriate), and assess the following:
  - Clinical status, including vital signs and neurological status
  - Results of blood investigations
  - ECG trace (especially signs of hypokalaemia, including S-T segment depression and prominent U-waves)
  - Cumulative fluid balance record
- Ensure that each review is documented in the patient's medical notes, including the components described above.
- Consider adjusting the total fluid rate using corrected sodium (Na<sub>corr</sub>) (see also appendix 4, page 16) taking into account the circulation and patient's general condition and state of hydration:
  - If the rise in Na<sub>corr</sub> is >5mmol/L in 4-8 hrs it suggests too much fluid loss or insufficient replacement. Consider increasing the fluid rate
  - If there is a fall in Na<sub>corr</sub> by more than 5mmol/L in 4-8 hrs it suggests too much fluid gain or too rapid replacement. Consider reducing the fluid rate

At each review confirm monitoring is compliant with the requirements above. Give specific consideration to the issues below. Address these in order of clinical priority.



## FLOW CHART 7 – ONGOING MANAGEMENT

Patient Name:  
Date of Birth:  
Hospital / NHS Number:

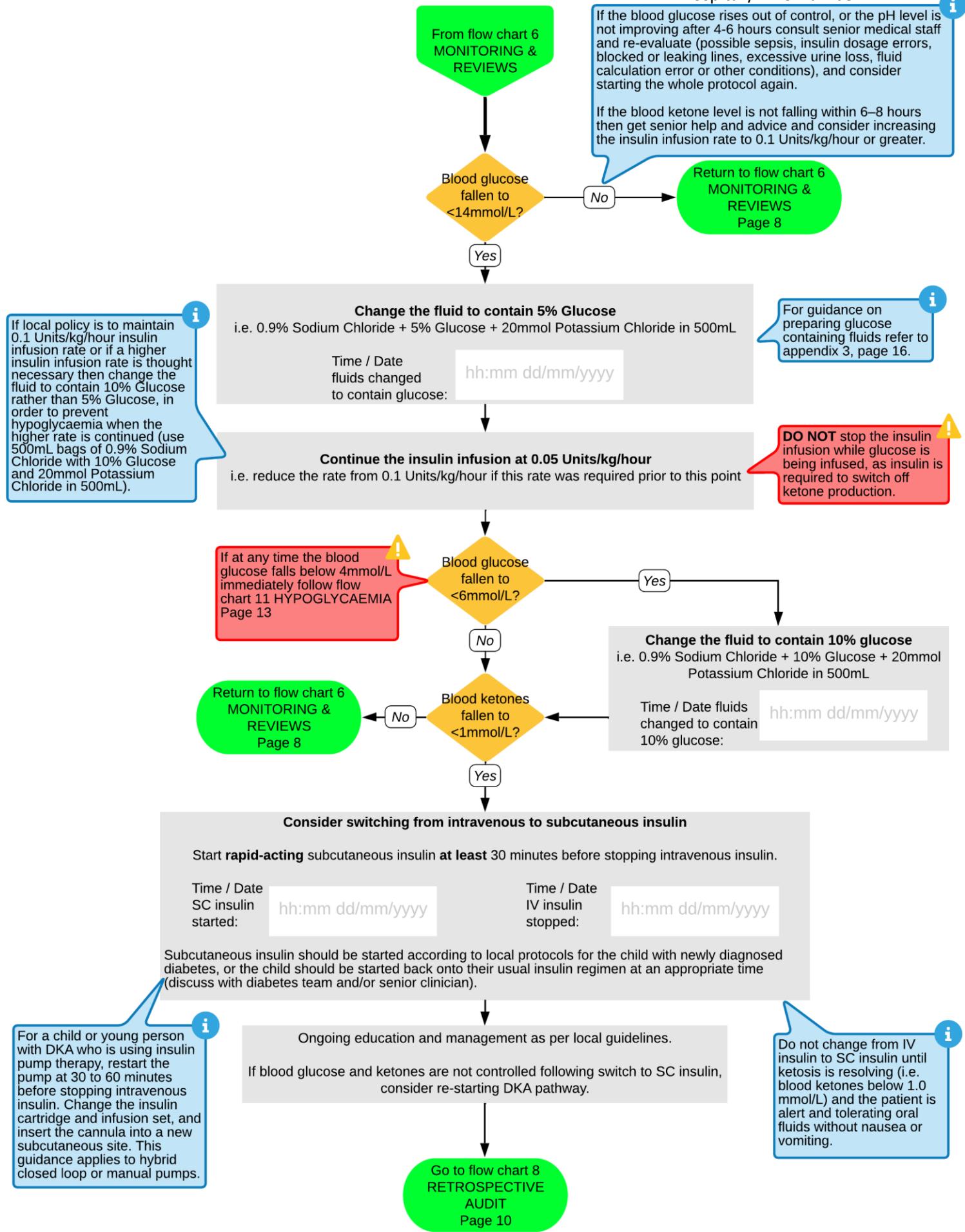


Chart completed by: \_\_\_\_\_

GMC number: \_\_\_\_\_

Signature: \_\_\_\_\_

Time / Date: \_\_\_\_\_

## FLOW CHART 8 – RETROSPECTIVE AUDIT

Patient Name:  
Date of Birth:  
Hospital / NHS Number:

From flow chart 7  
ONGOING  
MANAGEMENT

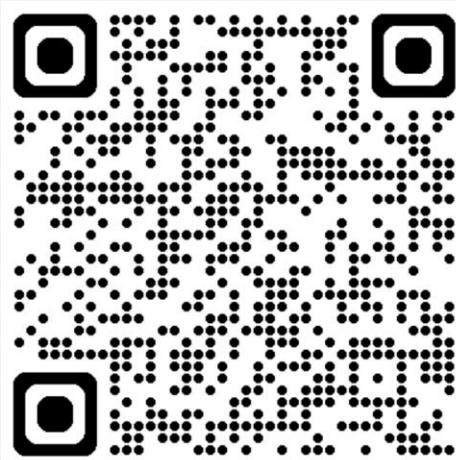
Please provide audit data for this DKA episode:

Go to [dka-calculator.co.uk](http://dka-calculator.co.uk) and select  
'Provide audit data for a DKA episode'.

When asked 'Do you have an Audit ID?'  
select 'No' to create a retrospective episode.

Or

Scan the QR code below.



Tick when audit complete

Were there any  
preventable factors which may  
have contributed to this  
episode of DKA?

Yes

Consider if any patient level or  
system level actions are required.

The following preventable/modifiable factors can be selected  
on the DKA Calculator:

- Missed/delayed diagnosis
  - Lack of or delayed access to primary care appointment
  - Missed or delayed diagnosis in primary care
  - Suboptimal or incorrect investigation/referral by primary care
  - Missed or delayed diagnosis in secondary/tertiary care
  - Suboptimal or incorrect management by secondary/tertiary care
  - Other diagnosis issue
- Diabetes technology issue
  - Concern of insulin pump malfunction
  - Concern insulin pump used incorrectly
  - Concern of glucose sensor malfunction
  - Other technology issue
- Lack of adherence
  - Concern insulin pump used incorrectly
  - Concern of inadequate supervision by parent/carer
  - Concern of lack of adherence to usual insulin therapy by child/young person
  - Suboptimal monitoring of glucose or ketones
  - Sick day rules not followed optimally
  - Other adherence issue
- Social factors
  - Concern of inadequate supervision by parent/carer
  - Language barrier
  - Other social factor

End of DKA Pathway

Chart completed by:	
GMC number:	
Signature:	
Time / Date:	

TABLE 1 - SERIAL DATA SHEET

Patient Name:

Date of Birth:

Hospital / NHS Number:

Time since protocol start (hrs)	Date/time (hh:mm dd/mm/yyyy)	Blood glucose (mmol/L)	Blood ketones (mmol/L)	pH	Base Excess	Bicarbonate (mmol/L)	Sodium (mmol/L)	Corrected sodium (mmol/L)	Potassium (mmol/L)	Urea (mmol/L)	Fluid balance (±mL)	Initial
0												
+2												
Changes:												
+6												
Changes:												
+10												
Changes:												
+14												
Changes:												
											Weight:	
+18												
Changes:												
+22												
Changes:												
+26												
Changes:												
											Weight:	
+30												
Changes:												
+34												
Changes:												
+38												
Changes:												
											Weight:	
+42												
Changes:												

After entering data values at each timeslot record any changes made on the following line. Record your clinical review and detailed plans in the patient notes. Remember to initial after completing each timeslot entry. Corrected sodium levels should typically rise as blood glucose levels fall during treatment. Corrected sodium levels may give an indication of the risk of cerebral oedema with a falling corrected sodium indicating an excess of free water and an increased risk of cerebral oedema. If corrected sodium levels fall during treatment, discuss with the consultant on call. See appendix 4, page 16.

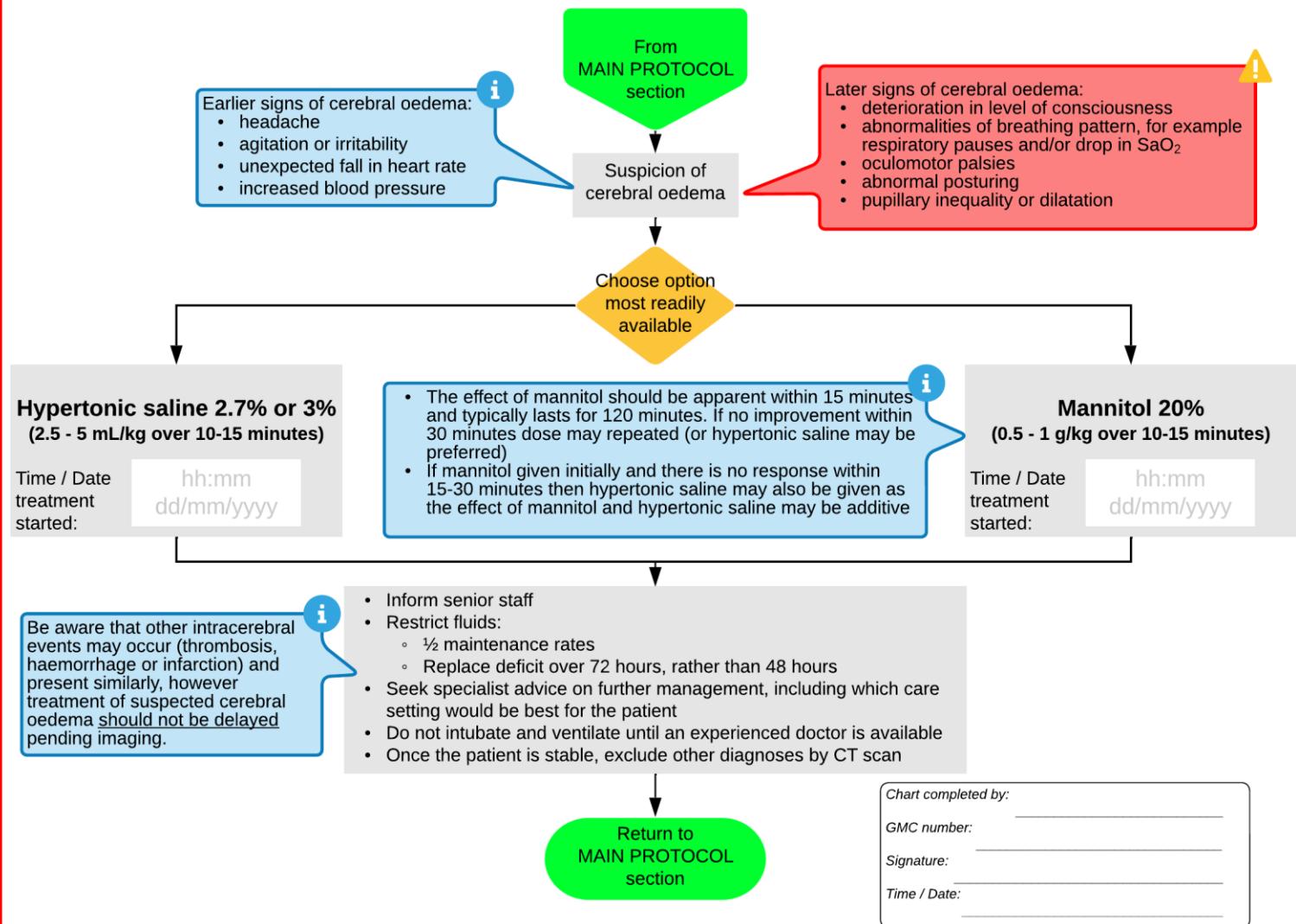
$$Na_{corr} = Na_{measured} + \left( \frac{Glucose - 5.6}{3.5} \right)$$

Or, use the corrected sodium / effective osmolality calculator  
[dka-calculator.co.uk/sodium-osmo](http://dka-calculator.co.uk/sodium-osmo)

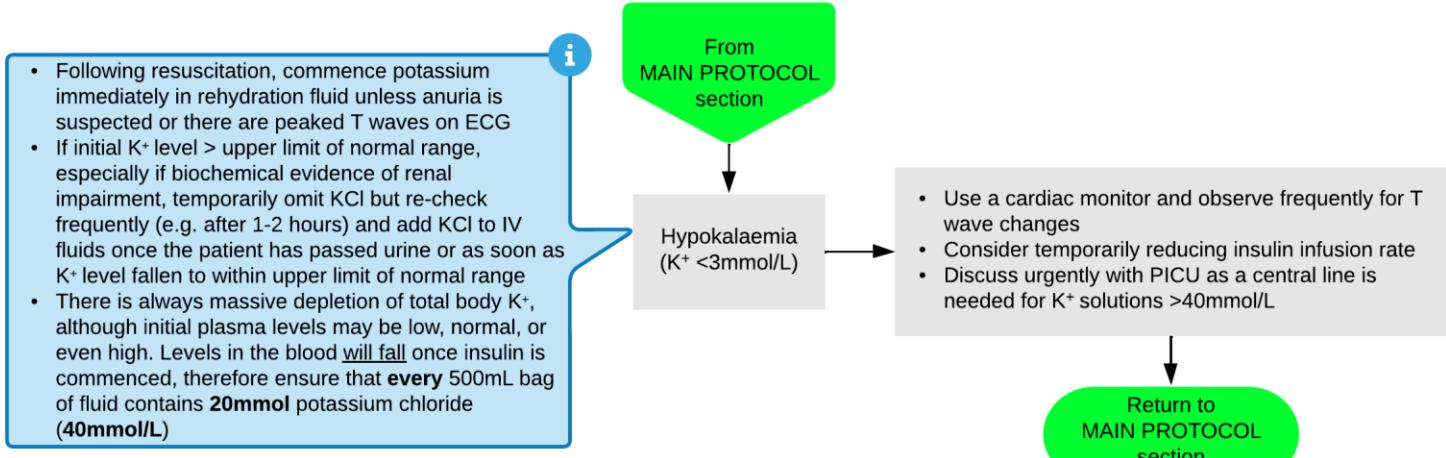


## FLOW CHART 9 – CEREBRAL OEDEMA

Patient Name:  
Date of Birth:  
Hospital / NHS Number:

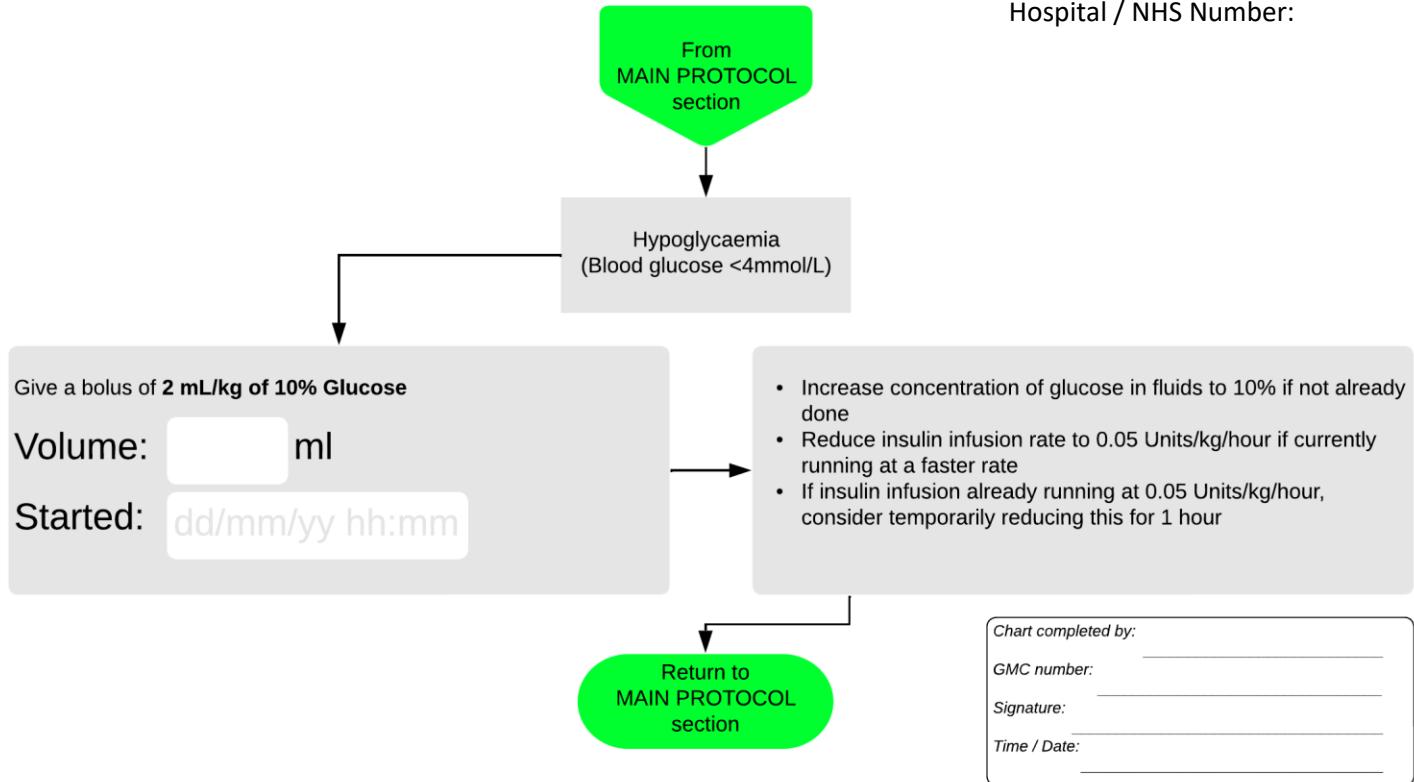


## FLOW CHART 10 – HYPOKALAEMIA



## FLOW CHART 11 – HYPOGLYCAEMIA

Patient Name:  
Date of Birth:  
Hospital / NHS Number:



## FLOW CHART 12 – PERSISTING ACIDOSIS

- i**
- Consideration should be given to calculating the anion gap
  - The anion gap is typically 20-30mmol/L in a patient with ketoacidosis. An anion gap >35mmol/L may suggest concomitant lactic acidosis due to sepsis or poor perfusion and should prompt a review of the overall clinical picture
  - It is not required for routine monitoring but may be helpful if the clinical picture or biochemistry is not improving

Anion gap =  Sodium - (  Chloride +  Bicarbonate ) =  mmol/L

e.g. 130 - (95 + 10) = 25mmol/L

If acidosis is not correcting, consider the following:

- Insufficient insulin to switch off ketones
- Inadequate resuscitation
- Sepsis
- Salicylate or other prescription or recreational drugs

Once all these causes of acidosis have been excluded, and if ketones are falling gradually, then residual acidosis is likely to be due to hyperchloraemia. This can be left to resolve spontaneously, and does not require any treatment. Acidosis due to hyperchloraemia need not delay the transition to oral fluids and subcutaneous insulin. It needs differentiating from ongoing ketosis.

Return to  
MAIN PROTOCOL  
section

Chart completed by:  
GMC number:  
Signature:  
Time / Date:

## FLOW CHART 13 - HYPEROSMOLAR HYPERGLYCAEMIC STATE

Patient Name: \_\_\_\_\_  
 Date of Birth: \_\_\_\_\_  
 Hospital / NHS Number: \_\_\_\_\_

Features which can differentiate HHS from other hyperglycaemic states such as DKA are:

- Hypovolaemia
- Marked hyperglycaemia (33.3mmol/L or more)
- No significant hyperketonaemia (<3mmol/L) or acidosis (pH>7.3, bicarbonate >15mmol/L)
- Osmolality usually 320mosmol/kg or more
- Often altered consciousness

From  
MAIN PROTOCOL  
section

Suspicion of  
Hyperosmolar  
Hyperglycaemic  
State (HHS)

Discuss with the responsible senior paediatrician – these children can be very difficult to manage, often requiring intensive care.

## Fluid therapy

The goal of initial fluid therapy is to expand the intra and extravascular volume and restore normal renal perfusion.

The rate of fluid replacement should be more rapid than is recommended for DKA.

Give an initial bolus of 20 mL/kg of 0.9% Saline

Volume:  mL

Started:  hh:mm dd/mm/yyyy

Additional fluid boluses should be given, if necessary, to restore peripheral perfusion.

Volume:  mL

Started:  hh:mm dd/mm/yyyy

Volume:  mL

Started:  hh:mm dd/mm/yyyy

- The goal is to promote a gradual decline in serum sodium concentration and osmolality
- As isotonic fluids are more effective in maintaining circulatory volume, isotonic saline should be restarted if perfusion and hemodynamic status appear inadequate as serum osmolality declines
- Serum sodium concentrations should be measured frequently and the sodium concentration in fluids adjusted to promote a gradual decline in corrected serum sodium concentration
- Mortality has been associated with failure of the corrected serum sodium concentration to decline with treatment, which may be an indication for haemodialysis
- Although there are no data to indicate an optimal rate of decline in serum sodium, 0.5mmol/L per hour has been recommended for hypernatraemic dehydration

Thereafter, 0.45–0.75% Saline with potassium should be administered to replace the deficit over 24–48 hours. Assume a fluid deficit of approximately 12–15% of body weight.

Rate:  mL/hour

Started:  hh:mm dd/mm/yyyy

## Further management considerations:

- If there is a continued rapid fall in serum glucose (>5mmol/L per hour) after the first few hours, consider adding 2.5 or 5% Glucose to the rehydration fluid. Failure of the expected decrease of plasma glucose concentration should prompt reassessment and evaluation of renal function
- Unlike treatment of DKA, replacement of urinary losses is recommended. The typical urine sodium concentration during an osmotic diuresis approximates 0.45% Saline; however, when there is concern about the adequacy of circulatory volume, urinary losses may be replaced with a fluid containing a higher sodium concentration
- **Insulin therapy**
  - Blood glucose levels will fall with fluid alone and insulin is NOT required early in treatment
  - Insulin administration should be initiated when serum glucose concentration is no longer declining at a rate of at least 3mmol/L per hour with fluid administration alone
- **Potassium**
  - Patients with HHS also have extreme potassium deficits; a rapid insulin-induced shift of potassium to the intracellular space can trigger an arrhythmia. Therefore potassium MUST be included in all fluids
- For further information see ISPAD Guidelines:

Return to  
MAIN PROTOCOL  
section

Chart completed by: _____
GMC number: _____
Signature: _____
Time / Date: _____

**APPENDIX 1 – GLASGOW COMA SCORE**
**Best Motor Response**

- 1 = none
- 2 = extensor response to pain
- 3 = abnormal flexion to pain
- 4 = withdraws from pain
- 5 = localises pain
- 6 = responds to commands

**Eye Opening**

- 1 = none
- 2 = to pain
- 3 = to speech
- 4 = spontaneous

**Best Verbal Response (with modification for younger patients)**
**>5 years**

- 1 = none
- 2 = incomprehensible sounds
- 3 = inappropriate words
- 4 = appropriate words but confused
- 5 = fully orientated

**2-5 years**

- 1 = none
- 2 = grunts
- 3 = cries or screams
- 4 = monosyllables
- 5 = words of any sort

**<2 years**

- 1 = none
- 2 = grunts
- 3 = inappropriate crying or unstimulated screaming
- 4 = cries only
- 5 = appropriate non-verbal responses (coos, smiles, cries)

**APPENDIX 2 – ESTIMATED WEIGHT TABLE**

Age	Guide weight (kg)	
	Male	Female
<b>6 months</b>	<b>8</b>	<b>7</b>
<b>12 months</b>	<b>9.5</b>	<b>9</b>
<b>18 months</b>	<b>11</b>	<b>10</b>
<b>2 years</b>	<b>12</b>	<b>12</b>
<b>3 years</b>	<b>14</b>	<b>14</b>
<b>4 years</b>	<b>16</b>	<b>16</b>
<b>5 years</b>	<b>18</b>	<b>18</b>
<b>6 years</b>	<b>21</b>	<b>20</b>
<b>7 years</b>	<b>23</b>	<b>22</b>
<b>8 years</b>	<b>25</b>	<b>25</b>
<b>9 years</b>	<b>28</b>	<b>28</b>
<b>10 years</b>	<b>31</b>	<b>32</b>
<b>11 years</b>	<b>35</b>	<b>35</b>
<b>12 years</b>	<b>43</b>	<b>43</b>
<b>14 years</b>	<b>50</b>	<b>50</b>
<b>Adult</b>	<b>70</b>	<b>70</b>

*Adapted from Advanced Paediatric Life Support, version 6, 2016*

## APPENDIX 3 – MAKING UP IV FLUIDS

The following fluids are generally available from Pharmacy. They may not be available on every ward. If you need to make it up, please do so as below, rather than waiting for pharmacy.

### 0.9% Sodium Chloride with 5% Glucose and 20mmol Potassium Chloride in 500mL

1. Remove 50mL from a bag of Sodium Chloride 0.9% with 20mmol Potassium Chloride in 500mL
2. Draw up 50mL of Glucose 5% using a syringe and add to the above bag to make the glucose concentration 5%
3. Mix well before administration

### 0.9% Sodium Chloride with 10% Glucose and 20mmol Potassium Chloride in 500mL

1. Remove 50mL from a bag of Sodium Chloride 0.9% with 5% Glucose and 20mmol Potassium Chloride in 500mL
2. Draw up 50mL of Glucose 5% using a syringe and add to the above bag to make the glucose concentration 10%
3. Mix well before administration

Plasmalyte does not contain enough potassium to be used on its own; discuss with pharmacy/PICU before using as maintenance fluid to ensure adequate potassium replacement is provided.

## APPENDIX 4 – EXPLANATORY NOTES

### Sodium and Corrected Sodium ( $Na_{corr}$ )

Hyponatraemia occurs in DKA as with hyperglycaemia the extracellular osmolality rises resulting in water movement from the intracellular space into extracellular space causing dilution of extracellular sodium and a low serum sodium. However, when glucose begins to fall through hydration and insulin, and the plasma glucose concentration is reduced, water leaves the extracellular space entering intracellular space raising the extracellular sodium concentration again and the serum sodium typically rises. Corrected sodium levels give an indication of the amount of free water in the circulation.

Corrected sodium levels should typically rise as blood glucose levels fall during treatment. It has been suggested that corrected sodium levels give an indication of the risk of cerebral oedema with a falling corrected sodium indicating an excess of free water and an increased risk of cerebral oedema. If corrected sodium levels fall during treatment, discuss with the consultant on call.

The formula for corrected sodium is:

$$Na_{corr} = Na_{measured} + \left( \frac{Glucose - 5.6}{3.5} \right)$$

You can also use the corrected sodium / effective osmolality calculator available at <https://dka-calculator.co.uk/sodium-osmo>.

For worked examples, refer to the full guideline (<https://www.bsped.org.uk/clinical-resources/bsped-dka-guidelines/>).

### Hyperchloraemic metabolic acidosis

Hyperchloraemic metabolic acidosis may occur following the administration of large amounts of chloride containing fluids given during the management of DKA. The preferential renal excretion of ketones instead of chloride can result in hyperchloraemia. The acidifying effect of chloride can mask the resolution of ketoacidosis if base deficit alone is used to monitor progress as there may appear to be a continuing base deficit with a continued low bicarbonate due to the chloride component rather than due to ketosis. Direct monitoring of ketones and calculation of the component of the base deficit due to chloride will help differentiate whether persisting acidosis is due to ongoing ketosis that may need additional treatment (adjustment to insulin infusion or fluids) or due to hyperchloraemia. Acidosis due to hyperchloraemia will correct spontaneously and doesn't need specific treatment. Acidosis due to hyperchloraemia need not delay the transition to oral fluids and subcutaneous insulin. It needs differentiating from ongoing ketosis.

The formula for calculating the component of the base excess due to chloride is:

$$BE_{\text{due to chloride}} = (\text{Sodium} - \text{Chloride}) - 32$$

For worked examples, refer to the full guideline (<https://www.bsped.org.uk/clinical-resources/bsped-dka-guidelines/>).

### Albumin

A low serum albumin can also contribute to a persisting acidosis which may be erroneously attributed to persisting ketosis. Some intensivists also recommend partitioning the component of apparent acidosis due to the reduced albumin to avoid it being inappropriately attributed to persisting ketosis.

The formula for calculating the component of the base excess due to albumin is:

$$BE_{\text{due to albumin}} = 0.25 \times (42 - \text{Albumin})$$

### Bicarbonate

Do not give intravenous sodium bicarbonate to children and young people with DKA. Only consider bicarbonate if there is life threatening hyperkalaemia or in severe acidosis with impaired myocardial contractility. It is anticipated that this would only ever be done following discussion with an intensivist.

### Risk of venous thrombosis

Be aware that there is a significant risk of femoral vein thrombosis in young and very sick children with DKA who have femoral lines inserted. Lines should be in situ as short a time as possible. Thromboembolic prophylaxis should be considered in young people >16 years (in line with NICE guidance), in young women taking the combined oral contraceptive pill and sick patients with femoral lines, following discussion with an intensivist.

### Oral fluids

Do not give oral fluids to a child or young person who is receiving intravenous fluids for DKA until ketosis is resolving and there is no nausea or vomiting.

A nasogastric tube may be necessary in the case of gastric paresis.

If oral fluids are given before the 48 hour rehydration period is completed, the IV infusion needs to be reduced to take account of the oral intake.

### Fluid losses

Do not give additional intravenous fluid to replace urinary losses. Urinary catheterisation should be avoided but may be useful in the child with impaired consciousness.

If a massive diuresis continues for several hours fluid input may need to be increased; this should be isotonic to the urine. If large volumes of gastric aspirate continue, these will need to be replaced with 0.45% saline with Potassium Chloride.

### Other complications

Other associations with DKA require specific management:

Continuing abdominal pain is common and may be due to liver swelling, gastritis, bladder retention, ileus. However, beware of appendicitis and ask for a surgical opinion once DKA is stable. A raised amylase is common in DKA.

Other problems are pneumothorax ± pneumo-mediastinum, interstitial pulmonary oedema, unusual infections (e.g. TB, fungal infections), hyperosmolar hyperglycaemic non-ketotic coma, ketosis in type 2 diabetes.

Discuss these with the consultant on-call.