# Management of paediatric diabetic ketoacidosis: a retrospective study

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# Background

- Diabetic ketoacidosis (DKA) is a major and common complication associated with a significant morbidity and mortality in children with diabetes.
- Following the Canadian Diabetes Association (CDA) guidelines, our tertiary care paediatric hospital was provided a DKA management protocol in 2009.

# Objectives

**Primary:** Assess the proportion of DKA cases that are marked by non-adherence to our DKA management protocol.

**Definition**: Non-adherence was arbitrarily defined as the occurrence of ≥1 major deviation and/or ≥2 minor deviations to the protocol during the acute management of DKA.

#### **Secondary:**

Describe the prevalence of complications

- Electrolytes disturbances
- Hyperchloremic acidosis
- Cerebral edema

### Methods

- Retrospective study by chart review, using a standardized data entry form
- Inclusion:
  - Age < 18 years,</li>
  - Pediatric ICU admission in our hospital between March 2009 and September 2013
- Exclusion:
  - Non-type I diabetes
  - Principal diagnosis at presentation other than DKA

Actual clinical practice of physicians in our hospital may not always correspond to our protocol recommendations.

### Results

- 73 patients included; demographic and clinical data (table 1)
- Protocol deviations and non-adherence (table 2)
  - 51% patients' management was marked by ≥ 1 major deviation, mostly:
    - Failure to monitor hourly neurological status
    - Inappropriate administration of 0,9%NaCl IV bolus in well-hydrated patients
  - 90% patient's management was marked by ≥ 2 minor deviations, mostly:
    - Failure to monitor urine ketones level every 4 hours
    - Failure to monitor blood gas and serum electrolytes every 2 hours
  - Overall, non-adherence to DKA protocol occurred in 95% of cases.

Complications and evolution (table 3)

- Rare hypoglycemic episodes and clinically suspected cerebral edema
- 75% cases of hyperchloremic acidosis (HA):
  - Mean time apparition: 4h41 after insulin perfusion beginning, using a normal anion gap definition of ≤14 (figure 1)
  - Perfusion insulin time in patients without HA: 9.2+/-5.6 hours; in patients with HA 17.1+/-9.1 hours (p<0,002)

#### Table 3 Complications

complications		
Hypoglycaemia	2 (3)	
Hyperchloremic acidosis	55 (75.3)	
Cerebral edema Clinically suspected Radiologically suspected	4(5.5) 1(1.4)	
Death	0	
Insulin perfusion length (h)	15.3(9)	

Figure 1

Anion gap and pH evolution during insulin perfusion

Date are expressed as mean(SD) or number(%)

### Demographic and clinical data

Age (y)	9.7 (5.0)
Female sex	42 (53.2)
Weight (kg)	33.9 (18.1)
New-onset diabetes	60 (75.9)
Antecedent of previous DKA	11 (13.9)
Insulin administration (in known diabetic patients)  · Multiple injections  · Insulin pump	12 (63.2) 7 (36.8)
Severity of DKA  · Mild  · Moderate  · Severe	24 (30.4) 30 (38.0) 25 (31.6)
Identified precipitating factor (in known diabetic patients) <ul> <li>Infection</li> <li>Poorly controlled diabetes</li> <li>Insulin pump dysfunction</li> <li>Omission of insulin (voluntary or not)</li> </ul>	1 (1.3) 3 (3.8) 4 (5.1) 10 (12.7)

Date are expressed as mean(SD) or number(%)

#### Table 2

#### Protocol deviations and non-adherence

Major deviations	
Administration of IV fluid bolus in a well-hydrated patient	5 (6.8)
Use of IV fluid other than saline 0,9% for initial rehydration	7 (12.7)
Rate of maintenance IV fluids > 2 times maintenance fluid requirements	4 (5.5)
Administration of IV insulin bolus	3 (4.1)
Initial insulin IV infusion rate > 0,1 U/kg/h	1 (1.4)
Administration of bicarbonates bolus without indication	0
Failure to monitor neurologic signs every hour in at least two occasions in the first 12 hours of acute management*	32 (46.4)
Total ≥ 1 major deviation	37 (50.7)
Minor deviations	
Failure to monitor blood glucose every hour in ≥ 2 occasions **	21(29.2)
Failure to monitor blood gas every 2 hours	33 (45.2)
Failure to monitor serum electrolytes every 2 hours	33 (45.2)
Failure to monitor BUN level every 4 hours	31 (42.5)
Failure to monitor urine ketones level every 4 hours	66 (90.4)
Failure to monitor serum calcium level every 4 hours	13 (17.8)
Failure to add 5% dextrose to IV fluid when blood glucose drops < 17 mmol/L	9 (12.3)
Failure to add 10% dextrose to IV fluid when blood glucose drops < 11 mmol/L	9 (12.3)
Failure to add KCl to IV fluid if serum potassium levels ≤ 5,0 mmol/L *	24 (33.3)
Failure to add $K_2PO_4$ to IV fluid if supplementation in KCl > 40 mEq/L or if serum potatssium level < 3,5 mmol/L	3/25 (12)
Total ≥ 2 minor deviations	66 (90.4)
Non-adherence	69 (94.5)

Date are expressed as number(%)

\*: 4 missing values; \*\*: 1 missing value

Non-adherence was defined as the occurrence of ≥1 major deviation and/or ≥2 minor deviations to the protocol during the acute management of DKA.

#### Mean\_pH ——— Mean\_AG

Temps

15

20

More education is needed about the importance of monitoring neurological status hourly and not giving overly generous intravenous hydration.

Discussion

Relevance to monitor ketonuria:

7.35

7.30

7.25

7.20

Mean\_pH

- No mention in CDA guidelines recommandations.
- ISPAD does not see the usefulness of urine ketones to determine when to stop the insulin infusion.
- Despite poor adherence demonstrated by our study, the number of complications observed did not exceed the complication rate expected.
- Frequently observed hyperchloremic acidosis
  - Probably related to prolonged use of NS
  - Extends the use of the IV fluids, insulin and probably ICU length of stay.

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# References

- 1. Devalia, B. (2010). Adherance to protocol during the acute management of diabetic ketoacidosis: would specialist involvement lead to better outcomes? International Journal of Clinical Practice, 64, 1580-1582.
- 2. Wherrett, D and all.(2013). Type 1 Diabetes in Children and Adolescents, Canadian Journal of Diabetes, Canadian Diabetes Association.
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**Acknowlegments:** M. David Simonyan and Dr Nathalie Laflamme

## Conclusion

- Non-adherence to our DKA management protocol was observed in majority of cases for minor deviations.
- The establishment of a protocol does not guarantee its adherence and surveillance of its application is needed.
- More studies have to be done to understand clinical impact of hyperchloremic acidosis.













