Diabetic ketoacidosis: Should current management include subcutaneous insulin injections?

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**ABSTRACT**

Diabetic ketoacidosis is a well-known acute complication in patients with both type 1 and type 2 diabetes mellitus. Although mortality has decreased considerably, it remains an important cause for admission to intensive care units. Medical management includes intravenous fluid therapy, insulin, correction of electrolyte abnormalities, and addressing the precipitating factor which in most cases is infection or non-compliance with insulin therapy. Usually patients with diabetic ketoacidosis are admitted to the intensive care unit for continuous infusion of insulin; however, the development of rapid acting insulin analogues has made it possible to treat mild to moderate diabetic ketoacidosis with subcutaneous insulin. Although studies using subcutaneous insulin include only a small number of patients, this approach seems as effective as intravenous insulin infusions in patients with mild to moderate diabetic ketoacidosis. Diabetic education and close follow-up for patients admitted for diabetic ketoacidosis remain essential to avoid recurrence and readmissions.

**Keywords:** Diabetic ketoacidosis, acute complication in diabetes, rapid acting insulin analogues, subcutaneous insulin in diabetic ketoacidosis

**INTRODUCTION**

Diabetic ketoacidosis (DKA) is a well-known acute complication in patients with both type 1 and type 2 diabetes. This condition results from a relative or absolute insulin deficiency combined with counter-regulatory hormone excess: glucagon, catecholamines, cortisol, and growth hormone. Diabetic ketoacidosis can be life threatening, but mortality rates have fallen since 1980, according to the National Diabetes Surveillance Program of the CDC. Mortality is usually related to associated comorbidities rather than hyperglycemia and ketoacidosis.

Diabetic ketoacidosis is defined by the presence of acidosis (serum bicarbonate <15 and/or pH <7.3), ketosis (ketonemia >3.0 mmol/L or ketonuria), and hyperglycemia with blood glucose >250mg/dl or known diabetes mellitus. Although hyperglycemia is usually present in DKA, euglycemic DKA has been reported in patients with type 1 diabetes who were vomiting, were fasting, were pregnant, or had been treated with insulin prior to presentation. Euglycemic diabetic ketoacidosis has also been recently reported in patients using sodium-glucose cotransporter 2 inhibitors, particularly in patients with type 1 DM.

DKA can be classified depending on severity of presentation into mild, moderate, or severe categories.

**PATHOPHYSIOLOGY**

Diabetic ketoacidosis is precipitated by inadequate levels of plasma insulin due to insulin deficiency.
Diabetic Ketoacidosis: Should Current Management Include Subcutaneous Insulin Injections? Quezada et al.

Diabetic Ketoacidosis (DKA) is a metabolic disorder that occurs when there is a lack of insulin, leading to an excess of glucose and ketones in the blood. This condition is characterized by high blood glucose levels, high levels of ketones, and low levels of bicarbonate. DKA is a medical emergency that requires prompt treatment to prevent complications such as brain damage and death.

**Medical Management**

The principles in the management of DKA are:

- IV fluids and correction of electrolyte abnormalities,
- Insulin therapy,
- Correction of the underlying precipitating factor

**IV Fluids**

Initial fluid therapy should expand intravascular and extravascular volume and restore renal perfusion. In the absence of heart failure, isotonic saline (0.9% NaCl) should be infused prior to insulin administration at a rate of 15–20 ml/kg/hr or greater during the 1st hour (≈1–1.5 L in the average adult). Fluid administration in the first hour of therapy before insulin administration has the following advantages: 1) it allows time to obtain a serum potassium level on presentation, 2) it corrects hypotension, which may increase if insulin is used without hydration, 3) it improves insulin action and may reduce the concentration of counter regulatory hormones and hyperglycemia.

ADA guidelines recommend that the subsequent choice for fluid replacement depends on the state of hydration, serum electrolyte levels, and urinary output. In general, 0.45% NaCl infused at 4–14 ml/kg/hr is appropriate if the corrected serum sodium is normal or elevated; 0.9% NaCl at a similar rate is appropriate if corrected serum sodium is low. Martin et al prospectively studied the effects of hypotonic, isotonic, and hypertonic fluids in patients with severe DKA. This study reported no significant difference in the volume of fluid retained with the different solutions; however, hypertonic fluids increased the frequency of hypertonicity, hypernatremia, and hyperchloremia.

In addition, some patients treated with hypotonic fluids developed diuresis; hence, rapid repletion of the plasma and extracellular volume with isotonic fluids is indicated in patients with DKA. In patients with renal or cardiac dysfunction serum osmolality, mental status, and cardiac and renal status should be monitored frequently during IV fluid resuscitation to avoid iatrogenic fluid overload. After the blood glucose level returns to normal, insulin therapy should be started, and the rate of fluid administration should be adjusted accordingly.

**Table DKA classification**

<table>
<thead>
<tr>
<th></th>
<th>Mild</th>
<th>Moderate</th>
<th>Severe</th>
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</thead>
<tbody>
<tr>
<td>Plasma glucose (mg/dL)</td>
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<td>&gt;250</td>
<td>&gt;250</td>
</tr>
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<td>7.00-7.24</td>
<td>&lt;7.00</td>
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<td>Urine ketones</td>
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<tr>
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<tr>
<td>Anion gap</td>
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<td>&gt;12</td>
</tr>
<tr>
<td>Mental status</td>
<td>Alert</td>
<td>Alert/drowsy</td>
<td>Stupor/coma</td>
</tr>
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</table>

Adapted from ADA guidelines for DKA management.
reaches < 250 mg/dL, IV fluids should be switched to 5% dextrose with 0.45-0.75% NaCl.  

**Insulin Therapy**

ADA guidelines recommend starting regular insulin with an initial IV bolus at 0.15U/kg followed by a continuous infusion at a dose of 0.1U/kg/hr. Glucose should fall by 50 mg/dl/hr. If the blood glucose does not decrease at this rate, hydration status should be rechecked and insulin infusion can be increased to reach a steady decline in blood glucose of 50-75mg/h. When the plasma glucose falls to 250 mg/dl, the insulin infusion rate can be decreased to 0.05–0.1U/kg/hr. The rate of insulin administration may need to be adjusted to maintain the above glucose values until acidosis resolves.

A new approach of insulin therapy has been recently studied. The fact that patients in diabetic ketoacidosis are admitted to ICUs for continuous insulin infusions and monitoring has led investigators to look for alternative strategies for treatment. A prospective randomized controlled trial by Doshi et al assigned 20 patients to insulin infusion at 0.1U/kg/hr at a set rate and 20 patients to IV insulin 0.1/kg/hr plus insulin Glargine at 0.3U/kg at the time of diagnosis. The primary end point was time to closure of the anion gap (TCAG). Upon closure of the anion gap, the control group received subcutaneous long acting insulin and IV insulin was continued for 2 more hours. In the experimental group, the insulin infusion was discontinued and the patients received long acting insulin 24 hrs after the initial dose. The estimated mean TCAG was 10.2 hrs in the experimental group and 11.6 hrs in the control group (p=0.63). Hospital length of stay was also slightly shorter in the experimental group (3.9 days) vs. the control group (4.6 days) (p = 0.66). There were no significant differences in outcomes between these two study arms.

Researchers have also started to study rapid acting insulin analogues as an alternative to the standard IV regular insulin continuous infusions in patients with diabetic ketoacidosis. Umpierrez et al studied 45 patients with DKA and randomly assigned them to 3 groups: 1) initial subcutaneous injection of 0.3U/kg of insulin aspart, followed by 0.1 U/kg every hour until resolution of DKA, 2) initial SC injection of 0.3 U/kg of aspart followed by a dose of 0.2U/kg 1 hour later and every 2 hours thereafter until blood glucose was 250mg/dl. The dose was then reduced to 0.1U/kg every 2 hours until resolution of DKA, 3) initial bolus of 0.1 U/kg of regular insulin followed by a continuous infusion at 0.1 U/kg/hr until blood glucose was 250 mg/dL. The infusion rate was then decreased to 0.05U/kg/hr until resolution of DKA. After comparing the three groups, the study found that the duration of treatment until resolution of hyperglycemia and DKA were similar among the three groups. The first group took on average 6.9 hrs of therapy until blood glucose was below 250 compared to 6.1 hrs in the second group and 7.1 hrs in the IV continuous insulin infusion (third group). Duration of therapy until resolution of DKA was 10 hours in the first group, 10.7 hours in the second group, and 11 hours in the third group.

Erso’z et al compared subcutaneous Lispro to regular insulin in 20 patients with mild to moderate DKA. Patients were randomized into two groups: Both groups received an initial bolus of 0.15U/kg of IV regular insulin. Group L was then given 0.075 U/kg of subcutaneous Lispro every hour; Group R was treated with a continuous infusion of IV regular insulin. Time to achieve a glucose level of less than 200 mg/dL was 9.4 hours in the L group and 12.7 hours in the R group. B-hydroxybutyrate was less than 0.6 mEq in 11.2 hrs in the L group vs. 15.3 hrs in the R group.

These studies suggest the following conclusions:

1. Duration of therapy until resolution of DKA in hours is 11-12 hrs ± 2 hrs in the IV insulin infusion groups and 10-12 hrs ± 2 hrs in the groups treated with SC insulin analogues.

2. Although the reported studies involved a small number of patients, treatment with subcutaneous insulin analogues appears to be effective in treating mild to moderate DKA patients.

It is important to understand that this type of treatment for DKA patients requires trained personnel in a step-down or regular floor units and very frequent subcutaneous injections which may not be practical on busy in-patient services. Although the studies
discussed above had a relatively small number of patients, the recommendation for using subcutaneous insulin for carefully selected patients is already included in the most recent publication of the ADA for Diabetes Care in the Hospital. This approach has not been studied in patients with severe DKA and, therefore, is discouraged.

**ELECTROLYTE ABNORMALITIES**

**POTASSIUM**

Serum potassium is frequently elevated in patients with diabetic ketoacidosis. This is caused by the extracellular shift of potassium in exchange for the hydrogen ions accumulated in acidosis and by the release of potassium from cells caused by glycolysis, insulin deficiency, and hyperosmolality. Treatment with insulin will shift potassium into the cell, causing a rapid decrease in potassium levels. ADA guidelines recommend that an insulin infusion should not be started if potassium levels are <3.3 mEq/l. If this is the case, potassium should be replaced before starting insulin treatment. When serum potassium levels are below 5.5 mEq/l, maintenance potassium of 20mEq can be added to each liter of IV fluids to keep concentrations within the normal range and avoid hypokalemia.

**PHOSPHATE**

Serum phosphate is lost by diuresis in diabetic ketoacidosis. Levels may be either normal or high upon presentation due to insulin deficiency. The level will start to decrease as soon as insulin treatment is established. Studies have failed to show any benefit with phosphate replacement in clinical outcomes in DKA. Overcorrection of phosphate can lead to severe hypercalcemia. Careful replacement may be indicated in patients with cardiac dysfunction, anemia, and respiratory depression and in patients with serum concentrations less than 0.32 mmol/L.

**BICARBONATE**

The use of bicarbonate therapy in patients with DKA is controversial. Some experts recommend the administration of 100 mmol of sodium bicarbonate to patients with a pH<6.9. However, several studies have shown no benefit with the administration of bicarbonate. In a recent retrospective study, Duhon et al found no benefit in administering bicarbonate in patients with severe DKA with pH <6.9. This study included 86 patients; 44 received IV bicarbonate therapy and 42 did not. The study concluded that the times to resolution of acidosis were similar in the two groups (8 hrs vs. 8 hrs, p=0.72). Additionally, patients who received IV bicarbonate also received slightly more insulin in the first 12 hrs of hospital stay (48 U vs. 44 U, p= 0.05) and first 24 hrs (100 U vs. 86 U p=0.04). A prospective randomized study of 21 patients failed to show beneficial or adverse effects on morbidity and mortality with bicarbonate therapy in patients with DKA whose admission arterial pH was between 6.9-7.1. Patients with very low pH values may benefit from partial correction of their pH. However, bicarbonate can cause hypokalemia, cerebral edema, and paradoxical central nervous system acidosis.

**CORRECTION OF THE UNDERLYING ETIOLOGY**

The most common precipitating factors are infection and inadequate insulin treatment as a result of non-compliance, especially in young patients. Other factors include alcohol abuse, pancreatitis, myocardial infarction, trauma, drugs, and new onset diabetes, especially in patients with type 1 diabetes. Diabetic ketoacidosis has also been reported with mismanagement of insulin pumps or leakage of the infusion system.

**RESOLUTION**

The resolution of DKA is determined by:

1. Blood glucose <200-250 mg/dL
2. Serum bicarbonate >18 mEq/L
3. Venous pH >7.3

Blood glucose levels alone should not be used as an indicator for the resolution of DKA since ketonemia takes longer to resolve than hyperglycemia. After
Diabetic ketoacidosis resolves, IV insulin and fluid replacement should be continued if the patient is NPO. When the patient is able to eat, a combination of short or rapid acting insulin with intermediate or long acting insulin can be started. Intravenous insulin infusion should be continued 1-2 hrs after starting subcutaneous insulin to avoid rebound ketoacidosis. To transition to subcutaneous insulin it is recommended that 60-80% of the daily infusion should be administered as basal insulin. These patients need close follow-up in outpatient clinics and diabetic education.

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**REFERENCES**