The role of point-of-care blood testing for ketones in the diagnosis of diabetic ketoacidosis

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Background. Urine dipstick testing for ketones is widely used when diabetic ketoacidosis (DKA) is suspected in patients with hyperglycaemia. If urinary ketones are positive, patients are referred for further management – often inappropriately, as the test is a poor surrogate for plasma ketones. Plasma beta-hydroxybutyrate (β-OHB) levels >3 mmol/L are diagnostic of DKA, while levels <1 mmol/L are insignificant. Objectives. To evaluate a hand-held electrochemical (point-of-care testing; POCT) ketone monitor and compare it with the gold-standard manual enzymatic method (MEM) for detection of plasma ketones.

Methods. In a prospective and comparative study, we evaluated the measurement of β-OHB by means of POCT and the MEM in 61 consecutive samples from patients with suspected DKA at Tygerberg and Karl Bremer hospitals, Cape Town, South Africa. Capillary (for POCT) and plasma samples (for the MEM) were obtained simultaneously and compared for accuracy. Precision was assessed with control samples.

Results. The POCT method was precise (coefficient of variation <4.5%), and there was a good correlation between the two methods (r=0.95). Regression analysis showed a proportional bias, with POCT reading higher than the MEM. However, when assessed at the relevant medical decision limits (β-OHB >3 mmol/L and <1 mmol/L), the total allowable error (bias + imprecision) was not exceeded. Patients will therefore still be classified correctly. The POCT method had a sensitivity of 100% and specificity of 89% for DKA (β-OHB >3 mmol/L), while at levels <1 mmol/L sensitivity was 100% and specificity 87.5%.

Conclusion. The POCT device provides an accurate and precise result and can be used as an alternative to the MEM in the diagnosis of DKA.


Diabetic ketoacidosis (DKA) is a common and severe acute complication of diabetes mellitus. The annual incidence is 46 - 50 episodes per 10 000 diabetic patients. DKA accounts for 14% of hospital admissions of diabetic patients and 10% of all deaths from diabetes in the Western world.10 In sub-Saharan Africa the picture is bleaker, with mortality rates up to 30%.11 It has been estimated that the annual cost of treating DKA in the USA exceeds 1 billion USD12 and that the average cost per DKA episode is USD6 500 - 7 500. This represented 25% of the total spent on diabetes facilities for further management. This results in large numbers of patients being referred inappropriately as having DKA. Uncontrolled hyperglycaemia in the presence of a β-OHB level of >3.0 mmol/L indicates unequivocal DKA, whereas a level of <1 mmol/L excludes any significant ketosis.13

Quantitative β-OHB measurement is unfortunately not offered by the routine laboratory at Tygerberg Hospital (TBH), Cape Town, SA (a tertiary academic training hospital). Samples are referred across town to the Red Cross War Memorial Children’s Hospital (RCWMCH) laboratory, and results are reported within 2 - 3 days. As a result, values cannot be used for decision-making in a medical emergency setting. Quantitative serum β-OHB measurement on a point-of-care testing (POCT) device has the potential to negate inappropriate referrals, which may translate into large cost savings if incorporated in initial patient assessment. It would also identify the ‘false negatives’ missed on Ketostix testing. POCT provides an attractive alternative to our current practice. Prior to its clinical use, it is essential that the POCT be evaluated to determine whether the device is accurate and precise and therefore fit for purpose. The
Clinical and Laboratory Standards Institute (CLSI) has published guidelines to facilitate this evaluation process.

Objectives
(i) To evaluate a hand-held electrochemical POCT ketone monitor (Medisense/Abbott Optium Xceed) and compare it with the manual enzymatic method (MEM), the current gold standard, to assess its accuracy and precision; (ii) to assess diagnostic performance of the POCT v. the MEM; and (iii) to calculate possible cost implications of utilizing POCT as an alternative to the MEM.

Methods
Study design and setting
The study was prospective and comparative, evaluating the measurement of β-OHB by means of electrochemical POCT and the MEM in consecutive patients referred from primary levels of care with suspected DKA. Referral was based on hyperglycaemia (capillary glucose >13.9 mmol/L) and the presence of urinary ketones on dipstick testing.

The study was conducted over a 4-month period in the acute medical admission ward and the high-care ward at TBH as well as in the acute medical admission ward at Karl Bremer Hospital (KBH), also in Cape Town. Venous blood samples were collected at admission and during treatment to ensure that a wide range of ketone values would be obtained. Patients were assessed clinically and attempts were made to identify the precipitant of the DKA episode.

Analytical performance
Method validations are performed to assess the degree of error inherent in a method and to determine whether the inaccuracy/bias and imprecision of the method will affect the interpretation of a test. CLSI documents EP09<sup>910</sup> ('Method comparison and bias estimation using patient samples') and EP15<sup>11</sup> ('User verification of performance for precision and trueness') ([www.clsi.org](http://www.clsi.org)) were followed. These documents state that precision can be determined using quality control samples, and that accuracy can be assessed by analysing at least 40 samples spanning the measurement range.

POCT was performed using the Medisense/Abbott Optium Xceed β-OHB meter, and results were expressed in mmol/L. Laboratory testing was performed by means of a MEM, results being expressed in μmol/L and then converted to mmol/L. The investigators were blinded to the results.

For precision studies, quality control samples at two levels close to the medical decision limits were provided by the manufacturer. Samples were analysed in triplicate over a 5-day period on the POCT device. The mean, standard deviation (SD) and coefficient of variation (CV, %) (mean/SD × 100) were calculated and compared with the manufacturer’s published results as per the package insert. The manufacturer claims a total imprecision of 8.8% at a β-OHB level of 0.7 mmol/L and a total imprecision of 3.1% at a β-OHB level of 4.3 mmol/L.<sup>12</sup>

Accuracy studies were performed on 61 samples collected from the 41 patients enrolled in the study. Samples from consecutively presenting patients were used with ketone levels that spanned the measurement range and included clinical decision-making levels. Capillary and serum samples were obtained at the same time. Capillary blood was used for POCT. Venous whole blood was collected in lithium heparin tubes (BD Vacutainer) and immediately transported on ice to the TBH National Health Laboratory Service (NHLS) laboratory. The transit time ranged from 10 to 20 minutes. At the laboratory, samples were immediately deproteinised with perchloric acid as per the RCWMCH protocol, stored at −70°C and transported to RCWMCH within 48 hours. At the RCWMCH laboratory the samples were stored at −20°C and analysed between 1 and 7 days of arrival. According to the literature, plasma samples are stable for several weeks at −20°C.<sup>13</sup>

Linear regression analysis and a Bland-Altman plot were used for analytical comparison and for depicting allowable bias and total allowable error. Method comparison statistics was performed with the Analyse-It version 2.3 data package for Excel.

Diagnostic performance
Sensitivity and specificity were calculated at the clinical decision-making limits (1 mmol/L and 3 mmol/L) to assess diagnostic performance.

Cost
The cost of reagent strips for POCT was borne by the manufacturer. The cost of the MEM was borne by the NHLS.

The cost for POCT is currently ZAR22 per strip, and the POCT device costs ZAR249. The cost of MEM for detection of serum ketones (including β-OHB + acetacetate) is ZAR228.88.

Results
Demographics and descriptive parameters
Most patients seen at TBH and KBH are of mixed ancestry, the gender and ethnic distribution being representative of the general population served by these hospitals. In total, 61 samples were collected from 41 patients, of whom 24 were females and 17 males. The mean age was 33 years (range 17 - 52). The majority had type 1 diabetes (n=29, 70.7%), seven being newly diagnosed. All the patients were found to have ketones present on dipstick testing of the urine, ranging from 1+ to 3+. Of the 41 referred patients with suspected DKA, only 15 were true DKA cases when using MEM ketones (>3 mmol/L) and hyperglycaemia (fingerprick blood glucose >13.9 mmol/L). The majority of the patients (n=26, 63.4%) were therefore inappropriately referred on the basis of positive urinary

![Fig. 1. Precipitants of DKA.](http://example.com/fig1.png)
ketones and hyperglycaemia. When evaluating ketosis (β-OHB >1 mmol/L and <3 mmol/L), an additional five patients had ketosis but not ketoacidosis. Even if these five patients are added to the 15 with DKA, more than half of all the patients (n=21, 51.2%) were inappropriately referred.

The factors that precipitated the DKA episodes are shown in Fig. 1. Although our study was not designed to investigate the causes thereof, it was clear that lack of adherence to insulin therapy and sepsis seemed to be the major contributors.

**Analytical performance**

**Precision**

Precision of POCT was assessed by adhering to guidelines of the CLSI. A summary of the results is presented in Table 1. At both low and high control levels, the total imprecision of the POCT device did not exceed the imprecision claimed by the manufacturer. Manufacturer claims were therefore verified at both control levels.

**Accuracy**

As the data were not normally distributed, non-parametric statistics were used. Spearman rank order correlation, performed with regard to actual values in mmol/L of both results obtained from POCT and the MEM (Fig. 2), demonstrated a very high correlation between the two methods (r=0.95). Since high correlation coefficients do not imply agreement of methods, we examined the agreement using the Bland-Altman difference plot and depicting allowable bias (18.4%) and total allowable error (68.7%). The Bland-Altman plot demonstrated a proportional bias, with the POCT device reading higher than the MEM. The bias exceeded the total allowable bias, but when combined with the precision, did not exceed the total allowable error (Fig. 3). When assessed at medical decision limits, the positive bias was 0.22 mmol/L at 1 mmol/L (p=0.77) and 0.68 mmol/L at 3 mmol/L (p=0.85).

**Diagnostic performance**

When compared with the MEM, the POCT device demonstrated a sensitivity of 100% and a specificity of 89.5% for diagnosing DKA (β-OHB >3 mmol/L) and a sensitivity of 100% and specificity of 87.5% for excluding DKA (β-OHB <1 mmol/L) (Table 2). Other performance criteria are summarised in Table 2.

**Discussion**

DKA is a common complication in our diabetic population. It is clear that the majority of patients were inappropriately referred with suspected DKA, and that this might have been avoided had the means to detect serum ketones been available at referral centres.

Only a minority of patients had pH measurements before being referred. Is it then correct to diagnose DKA at a certain level of β-OHB (in an uncontrolled diabetic), without knowing the pH level? Ketone body anion concentrations directly reflect the rate of ketone body production, which is accompanied by equimolar production of

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**Table 1. Manufacturer claims for precision of the POCT device validated using control samples**

<table>
<thead>
<tr>
<th>β-OHB, mmol/L</th>
<th>Claimed CV, %</th>
<th>Obtained CV, %</th>
<th>Claim verified</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.7</td>
<td>8.8</td>
<td>4.3</td>
<td>Yes</td>
</tr>
<tr>
<td>4.3</td>
<td>3.1</td>
<td>2.3</td>
<td>Yes</td>
</tr>
</tbody>
</table>

**Fig. 2. Scatter plot showing correlation between the POCT method and the MEM.**

**Fig. 3. Bland-Altman plot depicting allowable bias and total allowable error.**
hydrogen ions. Over the years numerous proposed markers and cut-offs have confirmed that any diagnostic criterion for DKA is arbitrary. This led the ADA to issue a consensus statement in 2006 that included set levels of pH, HCO₃, ketones, anion gap and glucose. While all of these criteria have serious limitations, the most robust criterion is regarded to be a serum bicarbonate level of <18 mEq/L. However, in a 2008 Mayo Clinic study comparing this cut-off level with appropriate criteria, there was a statistically strong correlation overall, but a one-third in the normal range, and one-third in the high-abnormal range could not be followed.

Conclusions

The Abbot OptumXceed POCT device provides an accurate and precise β-OHB result. This device offers a cost-effective practical alternative to laboratory measurement of ketones in patients with suspected DKA. Its adoption would bypass the use of urine dipstick testing to diagnose DKA, leading to more appropriate referrals and major cost savings. Our findings may potentially lead to better management of patients with DKA. However, appropriately designed studies will be needed to confirm this.

References


Table 2. Ability of the POCT device to diagnose DKA and ketosis

<table>
<thead>
<tr>
<th>DKA (β-OHB &gt;3 mmol/L)</th>
<th>n/N (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensitivity</td>
<td>23/23 (100.0)</td>
</tr>
<tr>
<td>Specificity</td>
<td>34/38 (89.5)</td>
</tr>
<tr>
<td>Positive predictive value</td>
<td>23/27 (85.2)</td>
</tr>
<tr>
<td>Negative predictive value</td>
<td>34/34 (100.0)</td>
</tr>
<tr>
<td>Ketosis excluded (β-OHB &lt;1 mmol/L)</td>
<td>n/N (%)</td>
</tr>
<tr>
<td>Sensitivity</td>
<td>33/33 (100.0)</td>
</tr>
<tr>
<td>Specificity</td>
<td>28/32 (87.5)</td>
</tr>
<tr>
<td>Positive predictive value</td>
<td>33/37 (89.2)</td>
</tr>
<tr>
<td>Negative predictive value</td>
<td>28/28 (100.0)</td>
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