Effect of dehydration on blood tests

In this third article in our ‘Test tips’ series, Dr Muhammad Masood Ashraf and Dr Rustam Rea examine the effects of dehydration on all essential diabetes blood tests, and provide guidance on key practical points to consider.

Introduction
Dehydration is common in patients presenting to the acute admissions ward. The most common reasons include poor oral intake and fluid loss from:

- Gastrointestinal tract (e.g. diarrhoea, vomiting).
- Skin (e.g. fever, burns).
- Urine (e.g. glycosuria, diuretic therapy, diabetes insipidus, diabetic ketoacidosis).

A reduction of the central circulating blood volume due to hypovolaemia accompanying dehydration results in a fall in cardiac filling pressure and stroke volume and, if uncompensated, a fall in cardiac output. The body can compensate by moving water from the extravascular to the intravascular space.¹,² As a result of these fluid shifts, changes in electrolytes and water concentrations in various body compartments occur which are reflected in many blood tests results. This is classically seen in patients with diabetic ketoacidosis and Hyperglycaemic Hyperosmolar State (previously HONK).

The clinical and biochemical features of dehydration³ are summarised in Box 1.

Effect of dehydration on haemoglobin, haematocrit and HbA₁c
Both haemoglobin and haematocrit increase in a dehydrated person.²,⁴ Hiroshi Nose¹ and colleagues induced dehydration in 10 subjects by exercise and checked haemoglobin (Hb), haematocrit (Hct), Na⁺, K⁺, Cl, and plasma osmolality at 0 minutes, 30 minutes and 60 minutes after exercise. Figure 1 shows the change in Hct, Hb, and plasma solids before and after dehydration. Immediately after exercise, these increased from 42.7±0.5% to 44.7±0.5%, 14.8±0.2g/dl to 15.8±0.2g/dl, and 8.4±0.1g/dl to 9.1±0.1g/dl, respectively. The significant differences observed before and after dehydration were maintained for the next 60 minutes.

HbA₁c is the measure of glycaemic status of an individual over the last three months.⁵ It is formed by a non-enzymatic reaction which occurs between glucose and the N-end of the beta chain.⁶ There is very little literature available on data search to suggest that dehydration directly affects HbA₁c. However, a rise in urea level as a result of dehydration can alter the HbA₁c test results depending on the assay.⁶

<table>
<thead>
<tr>
<th>Tests</th>
<th>9 March</th>
<th>11 March</th>
<th>13 March</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hb (g/L)</td>
<td>135</td>
<td>111</td>
<td>113</td>
</tr>
<tr>
<td>WBC (×10⁹)</td>
<td>24.2</td>
<td>16.9</td>
<td>12×10⁹</td>
</tr>
<tr>
<td>Haematocrit</td>
<td>0.443</td>
<td>0.368</td>
<td>0.369</td>
</tr>
<tr>
<td>MCV</td>
<td>96.9</td>
<td>96.1</td>
<td>94.4</td>
</tr>
<tr>
<td>Na⁺ (mmol/L)</td>
<td>160</td>
<td>156</td>
<td>150</td>
</tr>
<tr>
<td>K (mmol/L)</td>
<td>4.4</td>
<td>3.4</td>
<td>3.4</td>
</tr>
<tr>
<td>Urea (mmol/L)</td>
<td>31.3</td>
<td>18.9</td>
<td>9.9</td>
</tr>
<tr>
<td>Creatinine (μmol/L)</td>
<td>263</td>
<td>145</td>
<td>133</td>
</tr>
<tr>
<td>Glucose (mmol/L)</td>
<td>19</td>
<td>11</td>
<td>9</td>
</tr>
<tr>
<td>HbA₁c (mmol/mol)</td>
<td>75</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Total proteins (g/L)</td>
<td>60</td>
<td>–</td>
<td>48</td>
</tr>
<tr>
<td>Albumin (g/L)</td>
<td>29</td>
<td>–</td>
<td>23</td>
</tr>
<tr>
<td>ALT (IU/L)</td>
<td>24</td>
<td>–</td>
<td>23</td>
</tr>
<tr>
<td>ALP (IU/L)</td>
<td>132</td>
<td>–</td>
<td>126</td>
</tr>
<tr>
<td>Total bilirubin (µmol/L)</td>
<td>8</td>
<td>–</td>
<td>10</td>
</tr>
<tr>
<td>Total globulin (g/L)</td>
<td>31</td>
<td>–</td>
<td>28</td>
</tr>
</tbody>
</table>

Vignette
A 75-year-old lady was admitted with history of cough, high-grade fever and reduced oral intake for four days. She also complained of reduced urine output. Her past medical history included type 2 diabetes, hypertension, migraine, gastro-oesophageal reflux disease, and diabetes insipidus. Her medication included metformin and omeprazole. She was found to have dry mucous membranes, temperature 38.2°C, and BP 90/60mmHg. Chest examination revealed L-sided basal crackles.

Her blood tests on admission and after starting IV fluid replacement are summarised in the Table below.
Test tips

Effect of dehydration on blood tests

Clinical features of dehydration

- Dry mucus membrane
- Dry skin
- Reduced skin turgor
- Reduced axillary sweating
- Orthostatic hypotension
- Tachycardia and hypotension (indicates shock)
- Cognitive impairment
- Reduced urinary output (<0.5ml/kg/h is suggestive of acute kidney injury)
- Concentrated urine and high osmolality

Biochemical changes

- Raised serum urea
- Raised creatinine
- Reduced estimated glomerular filtration rate (eGFR)
- Increased urea:creatinine ratio
- Hypernatraemia (loss of water greater than salt loss)
- Raised serum or urine osmolality
- Raised urine specific gravity

Box 1. Summary of the clinical and biochemical features of dehydration

One study showed that in patients with uraemia, HbA1c measured by ion exchange chromatography was significantly elevated, but this was not correlated with the degree of glucose intolerance. This was due to the excessive amount of cyanate derived from the urea, which causes carbamylation at the N-terminal valine residue. This carbamylated haemoglobin (carbHb) results in an increase in the HbA1c (a + b) and, hence, the increased levels of HbA1c. However, newer ion-exchange HPLC assay methods show improved separation of the HbA1c fraction from other haemoglobin fractions and therefore no interference from carbHb.

Effect of dehydration on CBG measurements

Hypotension as a result of dehydration results in decrease in perfusion and increase in glucose utilisation in the local tissue leading to false low results of capillary blood glucose (CBG) tests. One study assessed the validity of the CBG measurements in the hypotensive, critically-ill patients. Capillary glucose values were significantly lower than those obtained from testing venous blood on the reagent strips and also lower than laboratory glucose measurements. Capillary glucose values in the hypotensive group were 33% lower than venous laboratory glucose values, and were significantly lower than the values obtained in the normotensive group.

Effect of dehydration on blood glucose

One study has looked at the effect of dehydration in frogs and demonstrated that dehydration can increase blood glucose levels. The rise in glucose was found to be out of proportion to changes in metabolite concentrations that could be due to passive concentration of the plasma (haemocencentration) as a result of dehydration.

Another study showed an increase in hepatic glucose production, with increased plasma glucose levels during hyperosmolality which can be caused by dehydration. The very high levels of venous glucose seen in patients with Hyperglycaemic Hyperosmolar State often resolve rapidly with rehydration alone without the need for insulin. This would suggest a significant effect of dehydration on venous glucose concentration.

Effect of dehydration on renal function tests

Dehydration has multiple effects on the kidney. The loss of body water leads to an increase in serum osmolality and activation of vasopressin which results in urinary concentration. This can be seen clearly in the above given Vignette.

Nose and colleagues also demonstrated sustained effects on plasma electrolyte concentration before and after dehydration (Figure 2).

However, there is an exception in the case of patients with cranial diabetes insipidus (CDI). Dehydrated patients usually present with an elevated serum urea level, owing in part to increased renal reabsorption of urea mediated by antidiuretic hormone (ADH). Serum urea values fall in patients with ADH deficiency (CDI) and this fact can be used to distinguish patients dehydrated because of CDI from those with usual hypertonic conditions.
dehydration and intact ADH secretion. In one study, the mean serum urea level was 2.9mmol/L in the CDI group and 15.4mmol/L in the patients without CDI, while the mean serum sodium level was 155mmol/L in both groups.13

**Effect of dehydration on lipid profile**

The effect of dehydration on lipid profile has been investigated in fasting subjects.14 Subjects were fasted, initially with no fluid replacement and then with salt and water supplementation. Subjects who had fasted with no fluids had a higher total serum cholesterol, HDL cholesterol, LDL cholesterol, apolipoprotein A-1, and apolipoprotein B, compared to subjects who had fasted with prior fluid and salt replacement.

**Effect of dehydration on liver function tests**

The above given Vignette demonstrates a significant difference in serum total proteins and albumin levels in a dehydrated patient, before and after hydration with intravenous fluids. However, bilirubin and liver enzymes levels remained unchanged, indicating that changes in protein levels were essentially due to hydration status rather than liver abnormality per se.

**Conclusion**

Clinicians should take the hydration status of the patient into account before interpreting the laboratory results. Before routine blood tests, patients should avoid unnecessary physical activity, avoid hot dry environments, ensure adequate intake of water, and avoid diuretic substances such as caffeine.

**References**


**Key points**

- HbA1c values can vary depending on the assay because of rise in urea level as a result of dehydration; however, there is no interference with modern assays
- Capillary blood glucose values in a dehydrated person can be significantly lower than the values obtained by venous reagent strips or laboratory glucose measurements
- Dehydration can increase the blood glucose levels per se
- Dehydration leads to an increase in plasma osmolality and urea levels, seen particularly in Hyperglycaemic Hypersmolar State

---

**Figure 2. Changes in electrolyte concentrations and osmolality (Posmol) in plasma after dehydration.** Significant differences were observed for all variables between control (C) and dehydrated conditions (0, 30, and 60 minutes). 

Dr Muhammad Masood Ashraf, MRCP, ST4, Diabetes and Endocrinology, Stoke Mandeville Hospital, Aylesbury, UK

Dr Rustam Rea, DM, FRCP, Consultant Physician in Diabetes and General Medicine, Oxford Centre for Diabetes, Endocrinology and Metabolism, Oxford University Hospitals NHS Foundation Trust, Oxford, UK

*Correspondence to:* Dr Rustam Rea, email: rustam.rea@nhs.net

---

**PRACTICAL DIABETES VOL. 34 NO. 5**

**COPYRIGHT © 2017 JOHN WILEY & SONS**