Serial plasma glucose changes in dogs suffering from severe dog bite wounds

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ABSTRACT

The objective of this study was to describe the changes in plasma glucose concentration in 20 severely injured dogs suffering from dog bite wounds over a period of 72 hours from the initiation of trauma. Historical, signalment, clinical and haematological factors were investigated for their possible effect on plasma glucose concentration. Haematology was repeated every 24 hours and plasma glucose concentrations were measured at 8-hourly intervals post-trauma. On admission, 1 dog was hypoglycaemic, 8 were normoglycaemic and 11 were hyperglycaemic. No dogs showed hypoglycaemia at any other stage during the study period. The median blood glucose concentrations at each of the 10 collection points, excluding the 56-hour and 64-hour collection points, were in the hyperglycaemic range (5.8–6.2 mmol/l). Puppies and thin dogs had significantly higher median plasma glucose concentrations than adult and fat dogs respectively (P < 0.05 for both). Fifteen dogs survived the 72-hour study period. Overall 13 dogs (81.3%) made a full recovery after treatment. Three of 4 dogs that presented in a collapsed state died, whereas all dogs admitted as merely depressed or alert survived (P = 0.004). The high incidence of hyperglycaemia can possibly be explained by the ‘diabetes of injury’ phenomenon. However, hyperglycaemia in this group of dogs was marginal and potential benefits of insulin therapy are unlikely to outweigh the risk of adverse effects such as hypoglycaemia.

Keywords: dog bite wounds, glucose, SIRS, trauma.


INTRODUCTION

Published reports on the incidence of bite wounds in dogs and cats vary from 10.2% to 14.7% of trauma cases, with a high incidence of severe bite wounds in intact male dogs weighing less than 10 kg. The average dog’s canine tooth can generate a crushing pressure of 150 to 450 pounds per square inch. The shearing and tensile forces involved in the crushing and shaking of especially smaller dogs can cause a large amount of damage to the underlying fascia, muscle, vasculature, nervous tissue, bone and parenchymatous organs, which predisposes these wounds to swelling, ischaemia and necrosis. This type of tissue damage is a result of the penetration of the dog’s teeth from an area with elastic tissue to an area with less elastic tissue.

This phenomenon has been referred to as the ‘iceberg effect’, because the external wounds conceal the severity of the underlying damage. Local and systemic factors can influence the wound healing process. Some of the local factors needed for optimal wound healing include oxygen and nutrients. During the early stages of healing, proteolysis of endogenous proteins mobilises amino acids, which are in turn used for glucose production. Glucose is the primary source of energy for leucocytes and the predominating carbohydrate substrate for fibroblasts in their synthesis of proteoglycan polymers during wound healing. However, excessively high plasma glucose concentrations can potentially inhibit neutrophil function by impairing phagocytosis and diminishing the production of oxygen radicals.

Dogs with dog bite wounds are generally in a state of shock when admitted and are therefore under excessive sympathetic adrenal control. After injury there is a rapid increase in plasma cortisol levels, which peak within 4–6 hours and decrease to resting levels within 24 hours. Changes in blood volume and afferent sensory nerve stimulation of the hypothalamus due to injury result in the release of catecholamines into the bloodstream. These hormones increase cardiac output and raise blood pressure. Furthermore, epinephrine increases basal metabolic rate by as much as 100% and stimulates metabolic activities such as glycogenolysis in the liver as well as glucose, cortisol and free fatty acid release into the blood. Patients with severe injury therefore undergo an accelerated form of starvation. During the hypermetabolic state, a number of alterations in carbohydrate metabolism are induced, these include:

- Enhanced peripheral uptake and utilisation of glucose by the wound and other organs such as the liver and spleen, which are involved in the immune response;
- Increased glucose production stimulated by the release of epinephrine, cortisol, glucagon and growth hormone;
- Hyperlactataemia due to hypoperfusion;
- Decreased glucose production due to the release of gram-negative bacterial endotoxin, interleukin-6, insulin-like factors and decreased hepatic export or increased utilisation;
- Depressed glycogenesis due to the persistent high rate of glycogen breakdown under the influence of epinephrine and glucagon, decreased glycogen synthetase activity or the presence of tumour necrosis factor (TNF);
- A rise in extra-cellular glucose concentrations due to defective suppression of gluconeogenesis, glucose intolerance and resistance to the peripheral action of insulin, also termed ‘diabetes of injury’.

Tissue trauma with or without infection can initiate the systemic inflammatory response syndrome (SIRS), in which multiple inflammatory, immunological, coagulation and fibrinolytic cascades are activated and interact. SIRS is characterised by hypermetabolism, a hyperdynamic cardiovascular state and clinical manifestations of fever or hypothermia, tachycardia, tachypnoea and leucocytosis.
Sepsis has been defined as SIRS with infection and severe sepsis as SIRS with infection and haemodynamic compromise. Septic shock is defined as severe sepsis that requires both volume replenishment and inotropic support to restore tissue perfusion. All these syndromes are commonly seen in dogs with bite wounds.

Alterations in blood glucose concentrations represent one of the most consistent findings in models of experimentally induced sepsis. Sepsis and SIRS can lead to hypoglycaemia due to impaired gluconeogenesis, especially from amino acids and increased peripheral uptake and utilisation of glucose. Lethal models of sepsis in animals demonstrate an initial hyperglycaemia followed by a phase of hypoglycaemia during which glucose production is suppressed.

Hypothermia, anorexia, age, pregnancy and breed have also been shown to influence blood glucose concentrations. A previous veterinary study has shown that blood glucose concentrations were significantly higher in dogs and cats with head trauma than in the control animals, but were not associated with outcome. In contrast, canine babesiosis cases with low blood glucose concentrations were more likely to die. However, dog bite wounds are peracute injuries usually inflicted on otherwise healthy animals and cannot be directly compared with a more chronic illness like babesiosis in which dogs can be presented days after the start of the disease process. Hyperglycaemia has also been reported in dogs with heart failure, a condition in which dogs are often presented days to months after the onset of the disease, rendering a comparison with acute trauma cases problematic.

The aim of this study was to describe the prevalence and prospective incidence of hypo-, normo- or hyperglycaemia in admitted canine bite wound cases over a 72-hour period. Historical, signalment and clinical factors were investigated to determine their possible effect on blood glucose concentrations.

### MATERIALS AND METHODS

Twenty dogs presented to the Onderstepoort Veterinary Academic Hospital (OVAH) that had been bitten within the preceding 24 hours, with at least 1 open wound, and whose clinical condition, in the opinion of the attending outpatients clinician, warranted the use of intravenous fluid therapy, were recruited into the study. The criteria set out above were used in order to collect the most severely affected animals. All the patients recruited for the study were admitted to the Intensive Care Unit (ICU) of the OVAH. Possible Babesia canis infection was excluded by means of a peripheral blood smear, because this disease can result in perturbations in blood glucose concentrations. In order to minimise any confounding influences on plasma glucose concentrations, all patients that received intravenous fluids containing dextrose or drugs that act by antagonising alpha-receptors were excluded from the study. These cases presented over an 8-month period.

The dogs’ owners signed a consent form and were asked to complete a questionnaire regarding the approximate time of the bite incident, time since the last meal, the dog’s pregnancy status, known medical conditions and the administration of any medication prior to presentation. On admission a full clinical examination was performed and habitus (alert, depressed or collapsed), age, sex, breed and body weight (Table 1), body condition (Purina Nine Point Body Condition Score), temperature, pulse and respiration were recorded. Blood was collected after a fasting period of at least 2 hours via the cephalic or jugular vein into a paediatric EDTA tube and a tube containing Sodium Fluoride Oxalate (NaF/Ox). Additional, serial NaF/Ox samples for plasma glucose determination were taken at 8-hourly intervals post-trauma and haematology at 24-hour intervals, measured from the time the bite was inflicted, as indicated by the owners. These blood collection time points were selected with the purpose of comparing the dogs at similar times in the trauma pathway. A full haematological examination was also done on admission. The NaF/Ox samples were centrifuged within minutes of collection at a speed of 1730 g for 8 minutes. The plasma was separated and aliquoted into cryopreservation tubes, labelled and immediately frozen at −20 °C. Glucose analysis was performed in a single batch using the hexokinase method, ensuring that the primary investigator was blinded as to the blood glucose concentrations of the patients for the duration of the study. Hypoglycaemia was defined as plasma glucose concentration below 3.3 mmol/l and hyperglycaemia as plasma glucose concentration above 5.5 mmol/l.

Temperature, pulse and respiration were recorded at each collection point. The study period extended from infliction of trauma to 72 hours, or until euthanasia or death. The study did not interfere with the normal clinical management of the cases.

Habitus at admission, time since last meal, time from infliction of trauma, age (<6 months = puppy; >6 months = adult), sex, body weight, body condition and outcome (died vs survived) were included as predictor variables for plasma glucose concentrations. Patients were classified into discrete, clinically meaningful categories of each predictor variable. The median plasma glucose concentration for each category of a variable was compared using Kruskal-Wallis 1-way ANOVA on ranks for multiple categories and the Wilcoxon rank sum test for 2 categories.

The presence of SIRS was determined from the admission data and other clinical and haematological data collected at 24-hour intervals after the initial trauma. To be considered SIRS positive, patients had to satisfy the following:

- White cell count <6000/mm³ or >16 000/mm³ and/or >3 % band cells, plus at least 1 of the following:
  - Rectal temperature of 38.1 or >39.2 °C;
  - Heart rate of >120 beats per minute;
  - Respiratory rate of >20 breaths per minute.

The plasma glucose concentration of SIRS-positive and SIRS-negative groups were compared at each blood collection time point using the Wilcoxon rank sum test. The association between habitus on admission and outcome was assessed using Fisher’s exact test. Statistical analysis was done using NCSS 2004 (NCSS, Kaysville, UT, USA).

### RESULTS

#### Signalement

The 3 dog breeds that were over-represented in this study included: Jack Russell terriers (5), Dachshunds (3) and Boerboels (3). The mean age of the patients in this study was 41.7 months. These patients had a mean weight of 10.1 kg and most dogs were females (11). Of the female dogs, 5 were intact (Table 1).

#### Outcome

Two dogs were euthanased within 8 hours of being bitten and only admission samples were obtained. One dog was euthanased after 48 hours. All instances of euthanasia were attributable to financial reasons and not prognosis. Another dog was discharged after 16 hours, also because of financial constraints; this dog’s outcome was unknown. This left 16 dogs in the study in which mortality could be assessed. One dog died during the study period and 2 died thereafter. Thus, 15 of 16 dogs survived the 72-hour study period and 13 of 16 dogs made an eventual recovery. All the dogs admitted as either alert or depressed survived (12), whereas 3 of 4 of the dogs admitted as collapsed, died (P = 0.004).
Incidence of glucose abnormalities

The medians and inter-quartile ranges of plasma glucose concentrations and the proportion of dogs that were hypo-, normo- and hyperglycaemic at each given collection time point, are shown in Table 2.

On admission, 1 dog was hypoglycaemic, 8 were normoglycaemic and 11 were hyperglycaemic. None of the dogs showed hypoglycaemia during the remainder of the study period. The median plasma glucose concentration at each of the 10 collection points was consistently in the hyperglycaemic range (5.8–6.2 mmol/l), excluding the 56-hour (5.4 mmol/l) and the 64-hour collection points (5.5 mmol/l), which were normoglycaemic. No dogs were found to be severely hypo- (<2.2 mmol/l) or severely hyperglycaemic (>10 mmol/l) during the study.

Systemic inflammatory response syndrome

The prevalence of SIRS at the 24-hour, 48-hour and 72-hour collection points is shown in Table 3. No significant differences in median plasma glucose concentrations between SIRS-positive and SIRS-negative dogs were detected at any collection point (Table 3).

Putative risk factors/predictor variables

No significant differences (P > 0.05) in the plasma glucose concentration were found, when assessing the groups within the different risk factors/predictor variables, with the exception of:

Habitus

- At the 32-hour collection point (depressed dogs = 5.6 mmol/l; collapsed dogs = 6.3 mmol/l).

Age

- On admission (puppy = 7 mmol/l; adult = 5.4 mmol/l).
- At the 8-hour collection point (puppy = 7.6 mmol/l; adult = 5.7 mmol/l).
- At the 16-hour collection point (puppy = 7 mmol/l; adult = 6 mmol/l).

Body condition

- On admission (thin dogs = 6.8 mmol/l; fat dogs = 4.7 mmol/l).
- At the 16-hour collection point (thin dogs = 6.9 mmol/l; fat dogs = 4.5 mmol/l).

A number of significant correlations were found between clinical or haematological variables and plasma glucose concentration:

- A positive correlation with respiratory rate (r = 0.63; P = 0.0051) only on admission.
severe sepsis in dogs with profound metabolic illness and that hypoglycaemia has been associated with a negative correlation. The study was surprising, considering the fact that the incidence of hypoglycaemia found in this study was lower than expected. The dogs were never in the hypoglycaemic range, and hypoglycaemia was only encountered in the 72-hour collection point.

A negative correlation was found between the degree of hypoglycaemia and the risk of death in human intensive care units, from 20.2% to 10.6%. A positive correlation has been found between dogs with and without SIRS in our study.

A consistent high median plasma glucose concentration was found in all collection points, except at 56 and 64 hours (Table 2). Plasma cortisol levels in healthy hospitalised (stressed) dogs would provide additional information if it can be compared with the plasma cortisol levels in dogs with dog bite wounds. Stress during hospitalisation might thus have contributed to the hyperglycaemia and could perhaps explain why 40% of dogs that presented with normal glucose levels subsequently developed hyperglycaemia.

In contrast with the babesiosis study, no association was found between hypoglycaemia and collapse at presentation. The low prevalence of hypoglycaemia might be attributed to the low case numbers. This study found no association between the degree of hypo- or hyperglycaemia and the eventual outcome, which was similar to a study on head trauma.

Median plasma glucose concentrations were never in the hypoglycaemic range and hypoglycaemia was only encountered in 1 case on admission. The low incidence of hypoglycaemia found in this study was surprising, considering the fact that hypoglycaemia has been associated with profound metabolic illness and severe sepsis in dogs.

A consistently high median plasma glucose concentration was found at all collection points, except at 56 and 64 hours (Table 2). Plasma cortisol levels in healthy hospitalised (stressed) dogs would provide additional information if it can be compared with the plasma cortisol levels in dogs with dog bite wounds. Stress during hospitalisation might thus have contributed to the hyperglycaemia and could perhaps explain why 40% of dogs that presented with normal glucose levels subsequently developed hyperglycaemia.

During the post-traumatic 'ebb phase', the body's initial response is centred on the release of glucagon, the mobilisation of lipid stores and reducing energy expenditure. The metabolic rate declines during this phase, but increases later. The final stage of the ebb phase is recognised by marked sympatho-adrenal discharge, lasts approximately 24 hours and can be associated with hyperglycaemia. If the animal survives this 'ebb phase', it progresses to the 'flow phase', which is characterised by an increased metabolic rate and enhanced breakdown of lean body mass. The afferent signals from the wound via pain and other peripheral receptors, as well as cytokines, interleukin 1 and tumour necrosis factor, initiate the process. These signals are integrated in the hypothalamus and stimulate the secretion of glucagon, cortisol, catecholamines and growth hormone. The net effect of these hormones, coupled with a defective suppression of gluconeogenesis and resistance to the peripheral action of insulin, is hyperglycaemia. This is all part of the hypermetabolic response to trauma, which may last for 7–10 days.

A high prevalence of SIRS was encountered (Table 3) on admission and during the study period. Sepsis has been defined as SIRS with an associated infection of the host by microorganisms. All bite wounds should be considered contaminated or dirty and infected and the local environment of the wound is ideal for bacterial replication and infection, with subsequent bacteraemia and or endotoxaemia. A previous report states that these syndromes are frequently seen in dogs with bite wounds, but no significant statistical differences in the median plasma glucose concentrations could be established between dogs with and without SIRS in our population.

Table 2: Descriptive statistics of plasma glucose concentrations (mmol/l) at admission and at 8-hour intervals since trauma.

<table>
<thead>
<tr>
<th>Category</th>
<th>Admission</th>
<th>8 h</th>
<th>16 h</th>
<th>24 h</th>
<th>32 h</th>
<th>40 h</th>
<th>48 h</th>
<th>56 h</th>
<th>64 h</th>
<th>72 h</th>
</tr>
</thead>
<tbody>
<tr>
<td>Count</td>
<td>20</td>
<td>16</td>
<td>18</td>
<td>17</td>
<td>17</td>
<td>17</td>
<td>17</td>
<td>16</td>
<td>15</td>
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</tr>
<tr>
<td>Median</td>
<td>6.0</td>
<td>6.2</td>
<td>6.2</td>
<td>5.8</td>
<td>5.8</td>
<td>6.2</td>
<td>6.2</td>
<td>5.9</td>
<td>5.4</td>
<td>5.5</td>
</tr>
<tr>
<td>IQR</td>
<td>5–7</td>
<td>5.4–7.3</td>
<td>5.6–6.8</td>
<td>5.2–6.8</td>
<td>5.6–6.3</td>
<td>5.5–6.8</td>
<td>5.1–6.5</td>
<td>4.8–5.9</td>
<td>5.1–6.2</td>
<td>5.1–6.8</td>
</tr>
<tr>
<td>Hypoglycaemic (&lt;3.3 mmol/l)</td>
<td>1 (5%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Normoglycaemic (3.3–5.5 mmol/l)</td>
<td>8 (40%)</td>
<td>5 (31%)</td>
<td>4 (22%)</td>
<td>5 (29%)</td>
<td>4 (24%)</td>
<td>6 (35%)</td>
<td>10 (63%)</td>
<td>9 (60%)</td>
<td>5 (33%)</td>
<td></td>
</tr>
<tr>
<td>Hyperglycaemic (&gt;5.5 mmol/l)</td>
<td>11 (55%)</td>
<td>11 (69%)</td>
<td>14 (78%)</td>
<td>12 (71%)</td>
<td>13 (76%)</td>
<td>11 (65%)</td>
<td>6 (37%)</td>
<td>6 (40%)</td>
<td>10 (67%)</td>
<td></td>
</tr>
</tbody>
</table>

Table 3: Prevalence and median values of plasma glucose concentrations (mmol/l) for SIRS positive and SIRS negative cases at admission and at the 24-hour, 48-hour and 72-hour collection points.

<table>
<thead>
<tr>
<th>Category</th>
<th>Admission</th>
<th>24 h</th>
<th>48 h</th>
<th>72 h</th>
</tr>
</thead>
<tbody>
<tr>
<td>SIRS +</td>
<td>13</td>
<td>6.6</td>
<td>5.8</td>
<td>5.9</td>
</tr>
<tr>
<td>SIRS –</td>
<td>7</td>
<td>5.4</td>
<td>6.8</td>
<td>5.9</td>
</tr>
<tr>
<td>% SIRS +</td>
<td>65</td>
<td>81</td>
<td>76</td>
<td>80</td>
</tr>
</tbody>
</table>

*Complete haematological data for one case was not available.

DISCUSSION
This study has found a significantly lower survival rate in bite wound cases that presented in a collapsed state. The low survival rate of collapsed animals should be investigated further as this may prove to be a relevant measure of severity and predictor of outcome of dog bite wounds, as was found in a recent study in 100 dogs with virulent canine babesiosis.

In contrast with the babesiosis study, no association was found between hypoglycaemia and collapse at presentation. The low prevalence of hypoglycaemia might be attributed to the low case numbers. This study found no association between the degree of hypo- or hyperglycaemia and the eventual outcome, which was similar to a study on head trauma.

Median plasma glucose concentrations were never in the hypoglycaemic range and hypoglycaemia was only encountered in 1 case on admission. The low incidence of hypoglycaemia found in this study was surprising, considering the fact that hypoglycaemia has been associated with profound metabolic illness and severe sepsis in dogs.

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During the post-traumatic ‘ebb phase’, the body’s initial response is centred on the release of glucagon, the mobilisation of lipid stores and reducing energy expenditure. The metabolic rate declines during this phase, but increases later. The final stage of the ebb phase is recognised by marked sympatho-adrenal discharge, lasts approximately 24 hours and can be associated with hyperglycaemia. If the animal survives this ‘ebb phase’, it progresses to the ‘flow phase’, which is characterised by an increased metabolic rate and enhanced breakdown of lean body mass. The afferent signals from the wound via pain and other peripheral receptors, as well as cytokines, interleukin 1 and tumour necrosis factor, initiate the process. These signals are integrated in the hypothalamus and stimulate the secretion of glucagon, cortisol, catecholamines and growth hormone. The net effect of these hormones, coupled with a defective suppression of gluconeogenesis and resistance to the peripheral action of insulin, is hyperglycaemia. This is all part of the hypermetabolic response to trauma, which may last for 7–10 days.

A high prevalence of SIRS was encountered (Table 3) on admission and during the study period. Sepsis has been defined as SIRS with an associated infection of the host by microorganisms. All bite wounds should be considered contaminated or dirty and infected and the local environment of the wound is ideal for bacterial replication and infection, with subsequent bacteraemia and or endotoxaemia. A previous report states that these syndromes are frequently seen in dogs with bite wounds, but no significant statistical differences in the median plasma glucose concentrations could be established between dogs with and without SIRS in our population.
A limited number of significant associations were found between the predictor variables and plasma glucose concentration. It is interesting to note that puppies and thin dogs tended to have higher plasma glucose concentrations than adult or obese dogs, respectively, particularly during the 1st 16 hours of trauma. The authors hypothesise that these dogs might differ in their cortisol responses to traumatic scenarios or have age-related differences in their cortisol feedback mechanisms.

The differences in response to cortisol can also be the result of differing sensitivities of the GLUT 4 transport proteins to cortisol and glucagon. The positive correlation between respiratory rate on admission and the blood glucose levels can potentially be related to stress on admission. Higher cortisol and glucagon levels can result in a decrease in GLUT 4 activity. Increased utilisation of glucose by bacteria and polymorphonuclear leucocytes has been implicated in hypoglycaemia associated with sepsis. Consequently, it is surprising that this study did not find any correlation between glucose concentration and leucocyte counts. The negative correlation between plasma glucose and haematocrit and the total red blood cell count can be the result of post-traumatic bleeding. Haemolysis can also be the result of infection due to an increase in the production of antibodies against red blood cells and the activation of complement- and phagocytic systems. It has also been shown that certain bacteria produce haemolysins during septicaemia (Leptospira spp., Clostridium spp., Streptococcus spp. and Staphylococcus spp.). These bacteria made up 30 % of the aerobic and 36 % of the total bacteria cultured from dog bite wounds in a study conducted in the same hospital as our current study, in which 84 % of bite wounds yielded positive cultures.

Due to the clinical and ethical nature of the study, all animals were maintained on Lactated Ringers solution (Ringers lactate, Fresenius Kabé) as intravenous infusion. Lactate can act as a precursor for glucose and, although unlikely, may have influenced the plasma glucose concentrations.

Food, although standardised where possible, was allowed ad lib, but removed 2 hours before sampling and could have influenced the plasma glucose concentrations. It is believed that this is unlikely, because the dogs had very poor appetites due to the severity of their illness. Furthermore, feeding small amounts was only commenced after 24 hours, by which time the median glucose concentrations were already in the mild hyperglycaemic range and showed no appreciable increase after this time point. It would be unlikely for varying degrees of food intake to influence plasma glucose in such a consistent manner and we suggest that this phenomenon should be more thoroughly investigated in veterinary medicine. A study on normal beagles showed that 1–2 hours of food deprivation was sufficient to discount the intake of food’s influence on blood glucose concentrations. Other potential limitations of the study include:

- Low numbers of patients used with no control group for comparison.
- The differing levels of severity (although an attempt was made to select the most severely bitten animals) and the current unavailability of a meaningful severity score for bite wound cases.
- The influence of stress induced by hospitalisation.
- The low specificity of the SIRS criteria used.

CONCLUSIONS

The low incidence of hypoglycaemia is surprising considering the high incidence of SIRS encountered. There was no significant difference between the plasma glucose concentrations of the SIRS-positive and SIRS-negative cases. The results show an almost uniformly high median plasma glucose concentration over the 72-hour sampling period. Although the study has its limitations regarding other potential causes of the mild hyperglycaemia encountered, this could possibly be ascribed to the ‘diabetes of injury’ response as reported in the human ICU literature. This may warrant exploration in future larger studies, because the use of insulin in these human cases has shown a significant reduction of fatalities resulting from acute injury. Few important associations were made between historical and clinical variables and plasma glucose concentration. However, the high incidence of death in the collapsed group and the higher plasma glucose concentrations found in puppies and thin dogs warrants future investigation with a larger group of animals.

ACKNOWLEDGEMENTS

The authors would like to thank Mss Elsbe Myburgh and Gertie Pretorius of the

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