Electrocardiographic findings in 130 hospitalized neonatal calves with diarrhea and associated potassium balance disorders

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Background: Hyperkalemia in neonatal diarrheic calves can potentially result in serious cardiac conduction abnormalities and arrhythmias.

Objectives: To document electrocardiographic (ECG) findings and the sequence of ECG changes that are associated with increasing plasma potassium concentrations (cK\textsuperscript{+}) in a large population of neonatal diarrheic calves.

Animals: One hundred and thirty neonatal diarrheic calves (age \( \leq \) 21 days).

Methods: Prospective observational study involving calves admitted to a veterinary teaching hospital.

Results: Hyperkalemic calves (cK\textsuperscript{+}: 5.8-10.2, blood pH: 6.55-7.47) had significantly (\( P < .05 \)) longer QRS durations as well as deeper S wave, higher T wave, and higher ST segment amplitudes in lead II than calves, which had both venous blood pH and cK\textsuperscript{+} within the reference range. The first ECG changes in response to an increase in cK\textsuperscript{+} were an increase in voltages of P, Ta, S, and T wave amplitudes. Segmented linear regression indicated that P wave amplitude decreased when cK\textsuperscript{+} > 6.5 mmol/L, S wave amplitude voltage decreased when cK\textsuperscript{+} > 7.4 mmol/L, QRS duration increased when cK\textsuperscript{+} > 7.8 mmol/L, J point amplitude increased when cK\textsuperscript{+} > 7.9 mmol/L, and ST segment angle increased when cK\textsuperscript{+} > 9.1 mmol/L. P wave amplitude was characterized by a second common break point at cK\textsuperscript{+} = 8.2 mmol/L, above which value the amplitude was 0.

Conclusions and Clinical Importance: Hyperkalemia in neonatal diarrheic calves is associated with serious cardiac conduction abnormalities. In addition to increased S and T wave amplitude voltages, alterations of P and Ta wave amplitudes are early signs of hyperkalemia, which is consistent with the known sensitivity of atrial myocytes to increased cK\textsuperscript{+}.

KEYWORDS
acidemia, calves, electrocardiography, hyperkalemia, hypokalemia

1 INTRODUCTION

Neonatal diarrhea in calves can result in metabolic derangements including electrolyte and acid-base imbalances. Acidemia and metabolic acidosis are frequently evident in diarrheic calves and are typically characterized by a low strong ion difference as a result of hyponatremia (accompanied by normochloremia or hyperchloremia) and an increase of unmeasured anions such as \( \alpha \)-lactate.\textsuperscript{1-3} Although neonatal diarrheic calves have a negative potassium balance because of intestinal potassium losses and low milk intake,\textsuperscript{4} they usually have normokalemic or hyperkalemic plasma concentrations in the presence of acidemia, with hypokalemia being infrequently observed.\textsuperscript{5,6}

Hyperkalemia is a clinically relevant electrolyte imbalance in diarrheic calves\textsuperscript{7} that has historically been attributed to an acidic state with intracellular buffering of hydrogen ions and impaired Na\textsuperscript{+}/K\textsuperscript{+}-ATPase activity with transcellular movement of potassium ions into the cells.\textsuperscript{8-10}
extracellular space as the proposed underlying mechanism. A potentially viable mechanism for acidemia-induced hyperkalemia is the activation of a cell membrane potassium channel called TREK-1 by low intracellular pH, resulting in potassium efflux from the cell. The presence of a hyperkalemic state in diarrheic calves strictly depends on the nature of an existing acidosis but not on acidemia per se, as D-lactic acidosis is only rarely associated with increased plasma potassium concentrations (cK⁺). More importantly, increased plasma cK⁺ in diarrheic calves is closely associated with severe dehydration, indicating that decreased glomerular filtration rate plays a central pathophysiological role in the development of hyperkalemia.

Abnormal plasma cK⁺ can have a profound effect on excitable tissues because the ratio of the extracellular to intracellular potassium concentration is a major determinant of the resting membrane potential, resulting in skeletal muscle weakness and cardiac conduction abnormalities. Moreover, the cardiotoxic effects of hyperkalemia are exacerbated by the presence of hyponatremia and metabolic acidosis, which are usually present in neonatal diarrheic calves. Electrocardiographic (ECG) manifestations in hyperkalemic human patients typically include tall and symmetric T waves, widening of the QRS complex, progressive flattening and eventually disappearance of P waves, and life-threatening dysrhythmias or ventricular escape rhythms. Similar findings occur in calves with experimentally induced hyperkalemia and experimentally induced or naturally acquired diarrhea. However, knowledge on ECG findings in calves with naturally acquired diarrhea is based on case reports, or case series, or small study populations of calves with marked increases in cK⁺. Moreover, retrospective studies on the presence of ECG changes in hyperkalemic human patients reported a low sensitivity of the ECG for diagnosing the presence of hyperkalemia. Also, ECG abnormalities such as tall and tent-shaped appearing T waves have been observed in acipedic human patients without hyperkalemia. Consequently, the aim of the present prospective observational study was to determine how frequent ECG abnormalities can be found in a large study population of calves with a broad range of cK⁺ values and other metabolic disorders and to assess what kind of ECG findings are associated with certain levels of increased cK⁺. Another aim of our study was to evaluate the association of metabolic imbalances, such as acidemia and dysnatremias, with the presence of ECG abnormalities in neonatal diarrheic calves.

2 | MATERIALS AND METHODS

2.1 | Calves

Between January 2015 and March 2017 a prospective study was conducted involving a convenience sample of 130 calves up to an age of 21 days that were admitted to the Clinic for Ruminants with Ambulatory and Herd Health Services, LMU Munich, with a clinical diagnosis of diarrhea. A subset of included hyperkalemic calves was also used in a recent study focusing on the potassium-lowering effect of different hypertonic infusion solutions. According to previous publications, diarrhea was defined as fecal consistency that permitted feces to run through slightly open fingers. The presence of concurrent problems was not an exclusion criteria for study enrollment; however, calves were not included if diarrhea was not considered to be the main problem on admission. Specifically 2 calves with acute respiratory distress, 4 calves with acute abdominal emergencies (gastrointestinal ileus or peritonitis), 3 calves with neurologic abnormalities and a postmortem diagnosis of meningitis, and 2 calves with complicated navel infections requiring surgical intervention were therefore excluded from the study. Our investigation was approved by the Ethics committee of the Center of Veterinary Clinical Medicine, LMU Munich (permit No. 84-21-09-2016; 29-04-06-2014).

2.2 | Electrocardiographic examinations

Electrocardiographic examinations were performed using a PC based ECG system (PC-EKG 2000, Eickemeyer, Tuttlingen, Germany) after the initial clinical examination was completed on admission to the hospital. Electrocardiograms were recorded in a standardized body position by placing calves in sternal recumbency with the legs positioned parallel to the long axis and folded normally at the carpal and tarsal joints (Figure 1). Electrocardiography electrodes were attached to the skin using alligator clips over the olecranon on the caudal aspect of the forelimbs, over the patellar ligament on the cranial aspect of the hindlimbs, and at the 8th intercostal space of the left thorax close to the costochondral junction (the neutral electrode was placed over the right patellar ligament). For placement of electrodes, a small area of the skin was clipped, cleaned with alcohol (70% solution) and ECG gel applied. An ECG was recorded at least for a 5 minutes period and monitored for the presence of arrhythmias.

The ECG software program digitized the signal at 500 Hz at a band width of 0.05–120 Hz and amplitude error measurement <1%. Digital filters were not applied. The digitization rate and low-frequency cutoff filter of 0.05 Hz met current American Heart Association recommendations. The high-frequency cutoff filter of 120 Hz was less than
The recommended filter of 150 Hz; however, the maximal measured Q, R, or S wave amplitude was 3.6 mV, which at an amplitude error measurement of <1% corresponded to <36 μV. Amplitudes and durations of the ECG complexes were determined using the bipolar limb lead II from a software program that averaged the amplitude and duration of the complexes during a selected 32 seconds period, thereby producing a single template of P, QRS, and T waves.

The software program required placement of markers for the start and end points of the P, QRS, and T waves. The markers were placed as follows. The start of the P wave was identified as the 1st deviation from a stable plateau of the TP line, which was defined as the isoelectrical line. In case of an absent P wave, the TR line was considered to be the isoelectrical line. Because of the presence of Ta waves, the end of the P wave and the start of the QRS complex was identified by means of abrupt slope changes in respect to the PR line. As ST-segment elevations and slope changes were observed in a large proportion of calves, a tangent was drawn from the ST segment centered on the approximate mid part of the QT interval, and the first and second deviations from this tangent defined as the J-point (junction point) and start of the T-wave, respectively.

The end of the T wave was defined as the point where the tangent of the steepest downslope of the T wave intersected with the isoelectrical line. Representative graphical displays of the placement of markers for software based ECG analyses are provided by Figure 2.

Heart rate, durations and amplitudes of P, QRS, and T waves as well as the duration of the PR interval, QT interval, and ST segment were then measured by the ECG analysis software program. All amplitude measurements were performed by referring to the isoelectrical line, except of measurement of the R wave, which was determined as deviation from the PR line.

The amplitude of the T wave was indexed to the modulus of S wave amplitude to provide a relative measure of changes in T wave amplitude. Furthermore, the ratio of T wave amplitude to P wave amplitude was calculated. The angle of the ST segment above the isoelectric line, deviations of the J point and start of T wave from the isoelectric line (J respective ST segment amplitude), as well as the intervals from the J point to the peak of the T wave (JT max) and to the end of the T wave (JT end) were also determined. The value for JT end is equivalent to that of QT interval minus QRS duration, and corrects the QT interval for prolongation of the QRS duration.
The measured QT interval was corrected for heart rate (QTcorrected [QTc]) using the formula according to Fridericia with the RR interval measured in seconds:

\[
\text{QTc interval Fridericia} = \frac{QT}{\sqrt{RR}}
\]

Tall and peak-shaped appearing T waves were defined as the presence of T wave amplitude $\geq 0.75 \text{ mV}$ and a “peakiness index” $\geq 8.0 \mu \text{V}/{\text{ms}}$, which was calculated by dividing T wave amplitude by T wave duration. Arrhythmias were defined as a deviation of 2 consecutive RR intervals of more than 10%. Based on an established reference interval for heart rate of 90–110 beats/min for calves, bradyarrhythmias were defined as the presence of arrhythmias and a heart rate $<90$ beats/min. Analysis of ECG measurements could not be masked as clinical and laboratory examinations were performed by the same investigator.

### 2.3 Determination of repeatability of ECG measurements

Additionally, in a total of 20 consecutively admitted calves, the repeatability of bipolar limb lead II measurements was assessed by taking the electrodes on and off and recording the ECG measurements twice. In those 20 calves the coefficients of variation, amplitudes, and duration from the duplicate measurements of limb lead II were compared to those of limb leads I and III and a base-apex lead. Base-apex leads were recorded by attaching the positive electrode of lead I to the skin of the left thorax in the vicinity of the apex beat at the 5th to 6th intercostal space immediately caudal to the olecranon, and the negative electrode on the skin over the jugular furrow in the caudal 3rd of the right neck.

### 2.4 Laboratory examinations

Within 15 minutes after recording the ECG, a lithium-heparinized blood sample was anaerobically collected from the jugular vein and blood pH, partial pressure of carbon dioxide (pCO$_2$) and oxygen (pO$_2$), sodium (Na$^+$), chloride (Cl$^-$), potassium (K$^+$), and ionized calcium concentrations (Ca$^{2+}$) determined using a blood pH, gas, and electrolyte analyzer with ion-selective electrodes (Rapidpoint 405, Siemens Healthcare Diagnostics, Tarrytown). Further detailed information concerning the determination of acid-base variables can be found elsewhere.

An automated analyzing system (Cobas c 311, Roche Diagnostics, Mannheim, Germany) was used for the biochemical analysis including determination of magnesium (Xyliyl blue), total protein (biuret), albumin (bromocresol green), and creatinine (picric acid) from serum samples and glucose (hexokinase) as well as l-lactate (l-lactate dehydrogenase) and d-Lactate (l-lactate dehydrogenase) from heparinized blood samples containing potassium fluoride as glycolytic agent.

### 2.5 Statistical analysis

Commercially available software programs were used for the statistical analysis of the results (SPSS, version 23, IBM, New York; GraphPad Prism, version 7.01, GraphPad Software, La Jolla; SAS, version 9.4, SAS Inc, Cary North Carolina) and P values of $<0.05$ were considered to be statistically significant. Data are presented as medians and interquartile ranges ($Q_1$/$Q_3$) because a large proportion of the variables were not considered to be normally distributed as indicated by the results of a Shapiro-Wilk test and visual examination of QQ-plots.

A subset of calves with $\text{cK}^+$ and venous blood pH within the reference range was used to compare results of ECG measurements to the remaining calves of our study population, which were assigned to one of the groups according to the measured $\text{cK}^+$ (hypokalemia, normokalemia, and hyperkalemia). For this purpose, a reference range for plasma $\text{cK}^+$ of 3.9 to 5.8 mmol/L and for venous blood pH of 7.35-7.50 was used. Categorized variables were compared using a chi-square test or Fisher’s exact test if the expected frequency in one or more of the cells of the contingency table was $<5$. Comparisons of continuous variables between the defined groups were made using a non-parametric Kruskal-Wallis test. For the subsequent pair-wise comparisons, a Mann-Whitney U test was used and the level of significance adjusted using the Bonferroni method ($P \leq 0.0167$). The same approach was also used to compare lead II measurements with those of limb I and III and a base apex lead. Repeatability for each lead were determined by calculating a variability index which was defined as the ratio between the intraquartile interval and the median.

Spearman’s coefficients of correlation ($r_s$) were also used to characterize associations between ECG measurements and laboratory variables. Using stepwise forward linear regression analysis, models for the P and T amplitudes, QRS duration, T wave peakedness index, and ST segment duration, angle, and amplitude were constructed including variables of clinical pathology significantly correlated to the dependent variable or considered relevant from a biological standpoint of view. To minimize the effects of collinearity, when 2 variables were closely correlated to each other ($r_s > 0.70$), only the variable that had the highest $r_s$ in the preliminary univariable analysis was entered into the model. The relative importance of the included variables was assessed by the order of entry into the model as well as by the change of the model $R^2$ value ($\Delta R^2$). Standardized residual plots of each multivariable model were examined to confirm an approximately normal distribution of residuals. Based on the examination of residual plots, the duration of the QRS interval and ST segment had to be log transformed to the base of 10 and a quadratic transformation of plasma $\text{cK}^+$ was used in all models that incorporated a linear value for $\text{cK}^+$ to prevent a non-hierarchical analysis. Additionally, interaction terms for plasma $\text{cK}^+$ and $\text{cNa}^+$ as well as venous blood pH were calculated and included into the models as centered variables by multiplying the differences from the mean of each variable.

Univariable logistic regression was also used to determine the associations between the presence of arrhythmias ($\text{yes} = 1; \text{no} = 0$), bradyarrhythmias ($\text{yes} = 1; \text{n} = 0$), tall and peak shaped appearing T waves ($\text{yes} = 1; \text{n} = 0$) as well as the absence of P waves ($\text{yes} = 1; \text{n} = 0$) and plasma $\text{cK}^+$, $\text{cNa}^+$, the $\text{K}^+$/Na$^+$-ratio (calculated by dividing $\text{cK}^+$ with $\text{cNa}^+$ and multiplying values with 100), venous blood pH, plasma glucose and serum magnesium concentration, and rectal temperature as continuous variables (determined as part of the clinical
examination on hospital admission). This was conducted by calculation of Odds ratios (OR) and associated 95% confidence intervals (95% CI).

Additionally segmented linear regression\(^3\)\(^8\)\(^3\)\(^9\) was used to characterize the relationship between each ECG variable of interest and cK\(^+\) by sequentially modeling the ECG variable-plasma cK\(^+\) relationship using 3 models. Because the focus of the study was on the ECG changes in hyperkalemia, data from calves with hypokalemia (plasma cK\(^+\) < 3.9 mmol/L)\(^6\) were not included in this statistical analysis.

The first model used ordinary linear regression to apply a linear regression equation to the data, such that \(y = b_0 + b_1 \times x\). The second model used nonlinear regression to apply 2 sequential linear regression equations with different slopes and a common break point ([K\(_1\)]) to the data, such that if \(x \leq [K_1]\) then \(y = b_0 + b_1 \times x\); if \(x > [K_1]\) then \(y = b_0 + b_1 \times [K_1 + b_2 \times (x - [K_1])].\) The 3rd model used nonlinear regression to apply 3 sequential linear regression equations with different slopes (the 3rd was constrained to a slope = 0) and 2 common break points ([K\(_1\)], [K\(_2\)]) to the data, such that if \(x \leq [K_1]\) then \(y = b_0 + b_2 \times x\); if \([K_1] < x \leq [K_2]\) then \(y = b_0 + b_1 \times [K_1 + b_2 \times (x - [K_1])],\) and if \(x > [K_2]\) then \(y = b_0 + b_1 \times [K_1 + b_2 \times ([K_2] - [K_1])].\) The last equation assumes that \(y = 0\) when plasma [K] > [K\(_2\)]. When 1 or 2 break points were identified, the estimated cK\(^+\) for the break point and the 95% confidence interval for the estimate were calculated to characterize the sequence of ECG changes as plasma cK\(^+\) increased. Model fit was evaluated using residual plots and summarized as \(R^2\) from ordinary linear regression or pseudo \(R^2\) from nonlinear regression, where pseudo \(R^2 = \frac{\text{Model SS} - \text{Mean SS}}{\text{Total SS} - \text{Mean SS}},\) where Mean SS is the sum of squares due to the mean, Model SS is the sum of squares due to the model, and Total SS is the total (uncorrected) sum of squares of Y (the dependent variable).

3 RESULTS

3.1 General conditions

The median value (and interquartile range) for age was 9 (7–12) days. Because of regional preferences, 90% (n = 117) of calves belonged to the Simmenthal breed (German Fleckvieh). Figure 3 illustrates individual values for venous blood pH and plasma cK\(^+\) of calves of the present study population. Hyperkalemia was present in 44 calves (34%), normokalemia in 73 calves (56%), and hypokalemia in 13 calves (10%). A total of 16 out of 73 normokalemic calves had venous blood pH values in the range of 7.35–7.50 and were therefore used for comparison of ECG measurements. Selected laboratory findings in the defined groups of calves are given in Table 1.

3.2 Specific ECG findings

Table 2 compares the results of lead II ECG measurements between the 4 defined groups of calves with statistically significant differences being found for the majority of ECG variables. Selected ECG findings of calves are shown in Figure 4. Ta waves were detectable in a total of 121 calves and Q waves in 9 calves. P waves were not detectable in 8 markedly hyperkalemic calves (Figure 4H-J). These calves had a median plasma cK\(^+\) of 8.9 mmol/L (range: 7.8–10.2) and a median cNa\(^+\) of 124 mmol/L (range: 120–147 mmol/L).

FIGURE 3 Venous blood pH and plasma potassium concentrations of calves of our study population (n = 130). Gray-shaded areas represent the reference ranges for plasma potassium concentration (3.9–5.8 mmol/L) and venous blood pH (7.35–7.50)

Bradycardia (< 90 beats/min) was observed in 24 calves of which 4 calves suffered from hypokalemia and 9 calves from hyperkalemia. Arrhythmias were seen in 26 calves of which 12 were presented in a hyperkalemic state with a median cK\(^+\) of 7.5 mmol/L (range: 6.0–9.9 mmol/L). An irregular sinus rhythm was evident in all 14 nonhyperkalemic calves of which 6 calves were also bradycardic. In 1 hyperkalemic calf with 1st degree A-V block, arrhythmias were related to supraventricular or junctional premature complexes (Figure 4C). Six out of the 12 hyperkalemic calves with arrhythmias had an irregular sinus rhythm, which was related to premature ventricular complexes in one of them (Figure 4E). The remaining 6 hyperkalemic calves had a slow irregular escape rhythm with missing P waves (Figure 4H).

Of the 8 calves with missing P waves, 2 had ventricular tachycardia (Figure 4I) and 1 calf a R-on-T configuration of ECG complexes (Figure 4J).

Tall (>0.75 mV) and peak-shaped appearing T waves (Figure 4D,E, H,I) were observed in 37 calves (28%) of which 1 calf was hypokalemic, 14 calves normokalemic, and 22 calves hyperkalemic, respectively. Those 37 calves had a median venous blood pH of 7.01 (range: 6.55–7.43).

3.3 Associations between ECG findings and variables of clinical pathology

Results of a univariable logistic regression analysis for the prediction of the presence of arrhythmias, bradyarrhythmias, tall and peak-shaped appearing T waves and absence of P waves are presented in Table 3. The K\(^+\)/Na\(^+\)-ratio was a statistically significant predictor in all models, but the absence of P waves and the presence of arrhythmias was not significantly associated with cK\(^+\).

As shown in Figure 5, no statistically significant association \((r_s = 0.14, p = .12)\) was found between plasma cK\(^+\) and heart rate. Also no or weak to moderate correlations were found between other ECG measurements and laboratory variables (Supporting Information Table S1). Multivariable stepwise linear regression analysis indicated that a
4 | DISCUSSION

Our study confirms that hyperkalemia is a clinically relevant electrolyte imbalance in neonatal diarrheic calves that is associated with typical ECG manifestations such as increased QRS duration, tall and peaked T waves, and findings of atrial standstill and life-threatening arrhythmias, that have also been previously described in Table 1. Selected findings of acid-base and electrolyte status in 130 neonatal diarrheic calves

<table>
<thead>
<tr>
<th>Variable</th>
<th>Group I Normokalemia</th>
<th>Group II Hypokalemia</th>
<th>Group III Normokalemia</th>
<th>Group IV Hyperkalemia</th>
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<tbody>
<tr>
<td>pH &gt; 7.35</td>
<td>n = 16</td>
<td>pH &lt; 7.35</td>
<td>n = 13</td>
<td>pH &lt; 7.35</td>
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<tr>
<td>K⁺ (mmol/L)</td>
<td>4.62 (4.40/4.82)</td>
<td>3.36 (3.12/3.66)*</td>
<td>4.85 (4.25/5.37)</td>
<td>6.75 (6.26/7.68)*</td>
</tr>
<tr>
<td>Na⁺ (mmol/L)</td>
<td>137.9 (135.9/142.6)</td>
<td>148.6 (141.8/158.2)*</td>
<td>136.1 (131.1/144.9)</td>
<td>135.6 (125.8/143.2)</td>
</tr>
<tr>
<td>Cl⁻ (mmol/L)</td>
<td>98 (95/102)</td>
<td>118 (111/124)*</td>
<td>105 (98/112)*</td>
<td>96 (91/103)</td>
</tr>
<tr>
<td>Ca²⁺ (mmol/L)</td>
<td>1.19 (1.14/1.24)</td>
<td>1.43 (1.35/1.49)*</td>
<td>1.30 (1.22/1.40)*</td>
<td>1.23 (1.14/1.35)</td>
</tr>
<tr>
<td>Mg²⁺ (mmol/L)</td>
<td>0.80 (0.74/0.85)</td>
<td>0.88 (0.86/1.27)*</td>
<td>0.93 (0.83/1.14)*</td>
<td>1.31 (1.10/1.51)*</td>
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<tr>
<td>Venous blood pH</td>
<td>7.389 (7.359/7.410)</td>
<td>6.957 (6.920/7.053)*</td>
<td>7.144 (6.988/7.288)*</td>
<td>7.136 (7.005/7.249)*</td>
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<tr>
<td>pCO₂ (mm Hg)</td>
<td>51.9 (48.6/56.0)</td>
<td>35.7 (28.6/41.0)*</td>
<td>44.7 (34.3/51.5)*</td>
<td>49.9 (40.8/61.1)</td>
</tr>
<tr>
<td>pO₂ (mm Hg)</td>
<td>32.7 (30.8/38.9)</td>
<td>47.1 (38.1/53.8)*</td>
<td>38.8 (33.8/43.9)</td>
<td>30.7 (25.3/37.2)</td>
</tr>
<tr>
<td>HCO₃⁻ (mmol/L)</td>
<td>31.4 (26.4/34.7)</td>
<td>8.4 (5.5/10.4)*</td>
<td>13.0 (7.6/23.6)*</td>
<td>16.1 (10.8/22.9)*</td>
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<tr>
<td>Base Excess (mmol/L)</td>
<td>5.6 (1.5/8.7)</td>
<td>-22.3 (-26.5/-19.1)*</td>
<td>-15.6 (-22.7/-2.5)*</td>
<td>-13.2 (-19.3/-5.1)*</td>
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<tr>
<td>Anion gap (mEq/L)</td>
<td>12.8 (11.8/16.6)</td>
<td>26.1 (22.9/28.0)*</td>
<td>22.1 (16.1/27.1)*</td>
<td>26.4 (22.7/31.7)*</td>
</tr>
<tr>
<td>L-lactate (mmol/L)</td>
<td>0.8 (0.3/1.7)</td>
<td>9.7 (7.8/11.6)*</td>
<td>5.2 (1.5/9.2)*</td>
<td>1.9 (1.3/6.1)*</td>
</tr>
<tr>
<td>D-lactate (mmol/L)</td>
<td>1.9 (1.4/2.7)</td>
<td>0.9 (0.7/1.1)*</td>
<td>1.4 (1.0/2.4)</td>
<td>4.4 (2.4/7.1)*</td>
</tr>
</tbody>
</table>

A subset of 16 calves with venous blood pH and plasma potassium concentrations in the reference range (group I) was used to compare respective findings to the remaining calves of our study population, which were assigned to one of the groups II-IV according to categories of plasma potassium concentrations. Values are reported as median and interquartile ranges. P values indicate a statistically significant difference between groups and asterisks indicate values, which are significantly different from group I (P < .0167).

4.5 | Outcome of therapy

A total of 90% of calves (n = 118) of our study population recovered and were discharged after a median duration of 11 days of hospitalization.

A comparative analysis of bipolar limb lead and base-apex lead measurements in neonatal diarrheic calves was given in Supporting Information Table S2. The values for K⁺ and venous blood pH in those 20 calves ranged from 3.3 to 8.0 mmol/L and 6.55 to 7.40, respectively. Statistically significant differences were found for the P, T, S, and ST segment amplitudes, T duration as well as for values of the ST angle and T to S amplitude ratio. However, pair-wise comparisons did not reveal a statistically significant difference between lead II and base-apex lead measurements. Coefficients of variation indicated a higher variation and poorer repeatability of lead I and III measurements when compared to lead II and base-apex lead measurements. Similar coefficients of variation were however observed for base-apex and lead II measurements, but values indicated a smaller variation for S and T amplitudes and the ST angle in lead II.

3.4 | Comparison of ECG measurements between bipolar limb leads and base apex lead

A comparative analysis of bipolar limb lead and base-apex lead measurements is given in Supporting Information Table S2. The values for K⁺ and venous blood pH in those 20 calves ranged from 3.3 to 8.0 mmol/L and 6.55 to 7.40, respectively. Statistically significant differences were found for the P, T, S, and ST segment amplitudes, T duration as well as for values of the ST angle and T to S amplitude ratio. However, pair-wise comparisons did not reveal a statistically significant difference between lead II and base-apex lead measurements. Coefficients of variation indicated a higher variation and poorer repeatability of lead I and III measurements when compared to lead II and base-apex lead measurements. Similar coefficients of variation were however observed for base-apex and lead II measurements, but values indicated a smaller variation for S and T amplitudes and the ST angle in lead II.
single reports of affected neonatal diarrheic calves.19,20,22 However, to the author’s knowledge, this is the first study to statistically determine the sequence of ECG changes as plasma cK+ increases in domestic animals or humans with naturally acquired hyperkalemia. All previous studies have been descriptive studies and have not employed objective quantitative analytical techniques.

We applied segmented regression instead of quantile regression or restricted cubic splines regression for 4 reasons. First, the focus of the study was to identify and statistically prioritize cK+ break points for commonly observed ECG abnormalities in hyperkalemia. Segmented regression provides 95% confidence intervals for an objectively identified break point,40 whereas quantile regression and cubic splines fail to provide this desired information.41 In fact, the researcher typically assigns values for the knots in the last 2 statistical methods.41 Second, segmented regression typically requires fewer data points than quantile regression.41 Third, the use of polynomial modelling is appropriately criticized in biology because a polynomial model is not relevant for many biological systems. Finally, examination of the scatterplots with

A subset of 16 calves with venous blood pH and plasma potassium concentrations in the reference range (group I) was used to compare respective findings to the remaining calves of our study population, which were assigned to one of the groups II-IV according to categories of plasma potassium concentrations.

Values are reported as ratios or median and interquartile ranges. P values indicate a statistically significant difference between groups and asterisks indicate values, which are significantly different from group I (P < 0.0167).

### Table 2: Lead II ECG findings and measurements in 130 neonatal diarrheic calves

<table>
<thead>
<tr>
<th>Variable</th>
<th>Group I Normokalemia pH &gt; 7.35 n = 16</th>
<th>Group II Hypokalemia pH &lt; 7.35 n = 13</th>
<th>Group III Normokalemia pH &lt; 7.35 n = 57</th>
<th>Group IV Hyperkalemia* n = 44</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heart rate (beats/min)</td>
<td>115 (100/144)</td>
<td>106 (83/121)</td>
<td>107 (96/126)</td>
<td>122 (95/148)</td>
<td>.19</td>
</tr>
<tr>
<td>Bradycardia (&lt;90 beats/min)</td>
<td>2/16</td>
<td>4/13</td>
<td>9/57</td>
<td>9/44</td>
<td>.55</td>
</tr>
<tr>
<td>Arrhythmiasb</td>
<td>1/16</td>
<td>2/13</td>
<td>11/57</td>
<td>12/44</td>
<td>.35</td>
</tr>
<tr>
<td>Bradycarrhythmiasc</td>
<td>1/16</td>
<td>0/13</td>
<td>5/57</td>
<td>7/44</td>
<td>.44</td>
</tr>
<tr>
<td>Absence of P-waves</td>
<td>0/16</td>
<td>0/13</td>
<td>0/57</td>
<td>8/44</td>
<td>.001</td>
</tr>
<tr>
<td>Tall and peaked T waves</td>
<td>1/16</td>
<td>1/13</td>
<td>13/57</td>
<td>22/44*</td>
<td>.001</td>
</tr>
<tr>
<td>P amplitude (mV)</td>
<td>0.17 (0.15/0.24)</td>
<td>0.24 (0.16/0.29)</td>
<td>0.19 (0.15/0.25)</td>
<td>0.21 (0.14/0.25)</td>
<td>.60</td>
</tr>
<tr>
<td>R amplitude (mV)</td>
<td>0.06 (0.02/0.20)</td>
<td>0.05 (0.02/0.09)</td>
<td>0.04 (0.02/0.17)</td>
<td>0.03 (0.02/0.06)</td>
<td>.12</td>
</tr>
<tr>
<td>S amplitude (mV)</td>
<td>−1.31 (−1.53/−1.14)</td>
<td>−1.90 (−2.29/−1.62)*</td>
<td>−1.58 (−1.88/−1.31)*</td>
<td>−1.93 (−2.18/−1.72)*</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>T amplitude (mV)</td>
<td>0.53 (0.40/0.61)</td>
<td>0.68 (0.51/0.95)</td>
<td>0.63 (0.48/0.83)</td>
<td>0.91 (0.64/1.16)*</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>J point amplitude (mV)</td>
<td>0.00 (−0.02/0.01)</td>
<td>0.03 (0.02/0.05)*</td>
<td>0.00 (−0.01/0.02)</td>
<td>0.03 (−0.01/0.06)*</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Ta amplitude (mV)</td>
<td>−0.05 (−0.08/−0.03)</td>
<td>−0.05 (−0.07/−0.04)</td>
<td>−0.06 (−0.07/−0.04)</td>
<td>−0.05 (−0.07/−0.03)</td>
<td>.78</td>
</tr>
<tr>
<td>P wave durationd (ms)</td>
<td>58 (56/76)</td>
<td>74 (66/83)</td>
<td>66 (61/74)</td>
<td>66 (60/78)</td>
<td>.098</td>
</tr>
<tr>
<td>PR intervald (ms)</td>
<td>116 (108/138)</td>
<td>136 (129/157)*</td>
<td>120 (113/136)</td>
<td>115 (102/134)</td>
<td>.003</td>
</tr>
<tr>
<td>QRS duration (ms)</td>
<td>55 (52/64)</td>
<td>76 (69/84)*</td>
<td>66 (61/72)*</td>
<td>74 (67/92)*</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>QT interval (ms)</td>
<td>242 (212/274)</td>
<td>292 (258/329)*</td>
<td>260 (230/288)</td>
<td>247 (212/279)</td>
<td>.009</td>
</tr>
<tr>
<td>QTc interval Fridericia (ms)</td>
<td>301 (285/330)</td>
<td>350 (324/369)*</td>
<td>316 (290/331)</td>
<td>311 (283/327)</td>
<td>.002</td>
</tr>
<tr>
<td>ST segment duration (ms)</td>
<td>90 (72/125)</td>
<td>96 (70/134)</td>
<td>92 (67/111)</td>
<td>64 (39/78)*</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>ST segment amplitude (mV)</td>
<td>0.05 (0.02/0.06)</td>
<td>0.09 (0.05/0.14)*</td>
<td>0.05 (0.04/0.08)</td>
<td>0.07 (0.04/0.12)*</td>
<td>.003</td>
</tr>
<tr>
<td>ST angle (°)</td>
<td>2.5 (0.5/4.0)</td>
<td>4.0 (2.0/5.5)</td>
<td>3.0 (2.0/4.5)</td>
<td>5.5 (3.0/9.0)*</td>
<td>.004</td>
</tr>
<tr>
<td>JT max (ms)</td>
<td>152 (131/185)</td>
<td>180 (144/201)</td>
<td>160 (132/187)</td>
<td>126 (100/152)*</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>JT end (ms)</td>
<td>181 (158/219)</td>
<td>222 (183/249)</td>
<td>192 (163/221)</td>
<td>163 (137/193)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>T wave peakedness index (µV/mS)</td>
<td>5.2 (3.6/6.6)</td>
<td>6.3 (4.2/7.3)</td>
<td>6.0 (4.6/7.9)</td>
<td>8.9 (5.7/11.4)*</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>T to P amplitude ratiod</td>
<td>3.0 (1.8/3.6)</td>
<td>3.3 (2.9/4.3)</td>
<td>3.2 (2.1/4.4)</td>
<td>3.5 (2.4/5.0)</td>
<td>.27</td>
</tr>
<tr>
<td>T to S amplitude ratio</td>
<td>0.41 (0.31/0.44)</td>
<td>0.36 (0.30/0.48)</td>
<td>0.39 (0.27/0.52)</td>
<td>0.44 (0.34/0.58)</td>
<td>.21</td>
</tr>
</tbody>
</table>

A subset of 16 calves with venous blood pH and plasma potassium concentrations in the reference range (group I) was used to compare respective findings to the remaining calves of our study population, which were assigned to one of the groups II-IV according to categories of plasma potassium concentrations.
the fitted segmented regression model indicated good model fit to the data.

Segmented regression analysis indicated that the first ECG evidence of hyperkalemia were increased voltages of P, Ta, S, and T wave amplitudes. Although increased T wave amplitudes have been well documented in hyperkalemia,17,42,43 we are unaware of any studies documenting an increase of voltages of P, Ta, and S wave amplitudes associated with early stages of hyperkalemia. These ECG changes are consistent with the well documented higher sensitivity of atrial myocytes to increased cK⁺ when compared with ventricular myocytes.14,44 This also explains the absence of P waves, as an ECG manifestation of atrial standstill, which was seen in 8 calves of the present study. Missing atrial activity was also reported in markedly hyperkalemic (and hyponatremic) diarrheic calves.19,20,22 and in markedly hyperkalemic human patients.15,17,42 In our study, the plasma potassium to sodium ratio, was significantly associated with the absence of P waves in the ECG. Low sodium concentrations also decrease the amplitude and the rate of rise of phase 0 of the cardiac action potential and therefore probably exaggerate the cardiotoxic effect of hyperkalemia.14 This interrelation between increased cK⁺ and low cNa⁺ is of some therapeutic usefulness because the cardiotoxic effects of hyperkalemia can be rapidly reversed by increasing plasma cNa⁺, which has been demonstrated for hypertonic saline in hyperkalemic humans, dogs, and a diarrheic calf.15,20,45

Hyperkalemia results in a lower transmembrane potassium gradient and therefore a less negative resting membrane potential. This magnitude of the transmembrane resting potential is a major determinant of upstroke velocity of the cardiac action potential (phase 0) and therefore the speed of cardiac conduction. Consequently hyperkalemia

FIGURE 4 Selected lead II ECG findings in calves with diarrhea (paper speed 25 mm/s, sensitivity 5 mm/mV). A, Normal appearance of an ECG in a diarrheic calf (cK⁺: 4.5 mmol/L, cNa⁺: 137 mmol/L, pH 7.41) with a Ta wave present (arrows). B, ECG of a severely acidic and hypokalemic diarrheic calf (cK⁺: 2.9 mmol/L, cNa⁺: 147 mmol/L, blood pH 6.74; rectal temperature: 36.7 °C; plasma glucose concentration: 4.2 mmol/L) showing sinus bradycardia (atrial and ventricular rate, 76 beats/min), deep and prominent S waves (arrows), and increased T wave amplitudes (asterisks). C, Prolonged PR intervals (asterisks) and a supraventricular or junctional premature complex (arrow) in a hypokalemic diarrheic calf (cK⁺: 3.1 mmol/L, cNa⁺: 157 mmol/L, blood pH: 6.75). D, Sinus tachycardia (atrial and ventricular rate, 170 beats/min), Ta waves (arrows) and increased amplitude and slightly peaked T waves (asterisks) in a hyperkalemic diarrheic calf (cK⁺: 6.3 mmol/L; cNa⁺: 150 mmol/L, blood pH: 7.01). E, Large amplitude “tented” symmetric T waves and prominent S waves in a hyperkalemic diarrheic calf (cK⁺: 6.8 mmol/L; cNa⁺: 135 mmol/L, blood pH: 7.04). Also note the increased QRS duration (asterisks) and ST amplitude (arrows) when compared with A. F, Unifocal premature ventricular complexes (arrows) in a hyperkalemic diarrheic calf (cK⁺: 6.3 mmol/L; cNa⁺: 126 mmol/L, blood pH: 7.10). G, Almost disappeared P waves (asterisks), marked ST segment elevation (arrows) and prominent S wave amplitudes in a hyperkalemic diarrheic calf (cK⁺: 7.8 mmol/L; cNa⁺: 118 mmol/L, blood pH 7.38). H, Bradycardia (ventricular rate, 56 beats/min) and missing P waves (arrows) and arrhythmias in a severely hyperkalemic diarrheic calf (cK⁺: 9.4 mmol/L, cNa⁺: 124 mmol/L, blood pH: 7.08). I, Ventricular tachycardia (ventricular rate, 133 beats/min) in a severely hyperkalemic diarrheic calf (cK⁺: 10.2 mmol/L; cNa⁺: 147, blood pH 7.15). J, Irregular wide-QRS rhythm in a severely hyperkalemic diarrheic calf (cK⁺: 9.9 mmol/L, cNa⁺: 139 mmol/L, blood pH 7.25)

### TABLE 3 Univariable binary logistic regression analysis for the prediction of specific ECG findings in a study population of 130 neonatal diarrheic calves

<table>
<thead>
<tr>
<th>Variable</th>
<th>OR</th>
<th>95% CI for OR</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Presence of arrhythmias (n = 26)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>cK⁺/cNa⁺-ratio</td>
<td>1.52</td>
<td>1.08-2.135</td>
<td>.016</td>
</tr>
<tr>
<td>Presence of bradyarrhythmias (n = 13)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>cK⁺/cNa⁺-ratio</td>
<td>2.03</td>
<td>1.31-3.16</td>
<td>.002</td>
</tr>
<tr>
<td>Presence of tall and peaked T waves (n = 37)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>cK⁺/cNa⁺-ratio</td>
<td>2.0</td>
<td>1.43-3.70</td>
<td>.001</td>
</tr>
<tr>
<td>Mg²⁺/cK⁺-ratio</td>
<td>2.03</td>
<td>1.42-2.90</td>
<td>.001</td>
</tr>
<tr>
<td>Venous blood pH</td>
<td>20.4</td>
<td>4.7-8.78</td>
<td>.001</td>
</tr>
<tr>
<td>Absence of P waves (n = 8)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>cK⁺/cNa⁺-ratio</td>
<td>66.5</td>
<td>3.0-1501.0</td>
<td>.008</td>
</tr>
<tr>
<td>Mg²⁺/cK⁺-ratio</td>
<td>9.82</td>
<td>1.15-83.84</td>
<td>.037</td>
</tr>
</tbody>
</table>

Abbreviations: OR, odds ratio; 95% CI for OR, 95% confidence interval for odds ratio. Plasma concentrations of potassium and sodium as well as venous blood pH, the K⁺/Na⁺-ratio, plasma glucose and serum magnesium concentrations, and rectal temperature were used as independent variables. Only statistically significant associations are shown.
can slow depolarization of the myocardium which can manifest in the ECG as prolongation of the PR interval (representing the time interval between the onset of atrial and ventricular depolarization) and more commonly as an increased QRS duration (representing the duration of ventricular depolarization).14,43,46

Another effect of hyperkalemia is an increased membrane permeability of potassium ions and an increase in the velocity of phase 3 of the cardiac action potential and therefore shortening of the action potential and acceleration of repolarization.32,46 The latter produces the typical changes of T wave morphology (representing ventricular repolarization) with tall, spiked (symmetric) or peak-shaped appearance which is considered to be the earliest ECG sign of hyperkalemia in humans.17,42,47 More pronounced hyperkalemia increases the risk for a further depression of sinoatrial and atrioventricular conduction which has been described to manifest in the appearance of escape rhythms and complexes57 as also seen in calves of the study reported here (Figure 4H and J). Along with a further prolongation of QRS duration, a fusion of the QRS complex with the T wave can occur,16,17 leading to a sine wave appearance of the ECG, which was also observed in a calf of our study population (Figure 4J).

Increased cK+ was also associated with alterations of Ta wave amplitude of calves in the study reported here. Ta waves represent the atrial repolarization phase and are usually broader, of lower amplitude, and in the opposite direction to the P wave. Studies about Ta waves are scarce as they are difficult to identify because the Ta amplitude is usually small and they merge with the QRS complex under normal conditions.47,48 Therefore studies about Ta wave characteristics are usually based on patients with atrioventricular blocks.47,48 A small increase in Ta wave amplitude was reported as an early sign of experimentally induced hyperkalemia in mice,49 which was similar to the findings of the study reported here.

Increased voltages of S wave amplitude (manifest as increased QRS peak to peak amplitude) was an early sign of hyperkalemia in calves of the present study population. Our finding was consistent with previous observations in calves with diarrhea21 and critically ill humans

### TABLE 4 Results of a stepwise forward linear regression analysis for predicting the log10 transformed duration of the QRS interval, P, T, S, and ST segment amplitude as well as a T wave peakedness index, ST angle and the log10 transformed duration of the ST segment by means of variables of clinical pathology in calves of the study population

<table>
<thead>
<tr>
<th>Order of entry</th>
<th>Variable</th>
<th>Coefficient</th>
<th>± SE</th>
<th>P value</th>
<th>ΔR²</th>
<th>Model R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>K × K</td>
<td>0.015</td>
<td>0.002</td>
<td>&lt;.001</td>
<td>.402</td>
<td>.402</td>
</tr>
<tr>
<td>2</td>
<td>K × K</td>
<td>−0.154</td>
<td>0.022</td>
<td>&lt;.001</td>
<td>.141</td>
<td>.543</td>
</tr>
<tr>
<td>3</td>
<td>pH</td>
<td>−0.127</td>
<td>0.030</td>
<td>&lt;.001</td>
<td>.055</td>
<td>.598</td>
</tr>
<tr>
<td>4</td>
<td>Creatinine</td>
<td>0.0001</td>
<td>0.00004</td>
<td>.001</td>
<td>.028</td>
<td>.626</td>
</tr>
<tr>
<td>5</td>
<td>Na</td>
<td>−0.001</td>
<td>0.001</td>
<td>.005</td>
<td>.023</td>
<td>.649</td>
</tr>
</tbody>
</table>

**P amplitude**

<table>
<thead>
<tr>
<th>Order of entry</th>
<th>Variable</th>
<th>Coefficient</th>
<th>± SE</th>
<th>P value</th>
<th>ΔR²</th>
<th>Model R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>K × K</td>
<td>−0.011</td>
<td>0.002</td>
<td>&lt;.001</td>
<td>.122</td>
<td>.122</td>
</tr>
<tr>
<td>2</td>
<td>Log Mg</td>
<td>0.348</td>
<td>0.062</td>
<td>&lt;.001</td>
<td>.200</td>
<td>.322</td>
</tr>
<tr>
<td>3</td>
<td>K</td>
<td>0.095</td>
<td>0.023</td>
<td>&lt;.001</td>
<td>.081</td>
<td>.403</td>
</tr>
</tbody>
</table>

**T amplitude**

<table>
<thead>
<tr>
<th>Order of entry</th>
<th>Variable</th>
<th>Coefficient</th>
<th>± SE</th>
<th>P value</th>
<th>ΔR²</th>
<th>Model R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>K × K</td>
<td>5.725</td>
<td>0.883</td>
<td>&lt;.001</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>pH</td>
<td>−0.663</td>
<td>0.122</td>
<td>&lt;.001</td>
<td>.152</td>
<td>.402</td>
</tr>
<tr>
<td>3</td>
<td>K</td>
<td>−0.198</td>
<td>0.094</td>
<td>.037</td>
<td>.021</td>
<td>.422</td>
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</table>

**S amplitude**

<table>
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<tr>
<th>Order of entry</th>
<th>Variable</th>
<th>Coefficient</th>
<th>± SE</th>
<th>P value</th>
<th>ΔR²</th>
<th>Model R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Log Mg</td>
<td>−1.840</td>
<td>0.383</td>
<td>&lt;.001</td>
<td>.298</td>
<td>.298</td>
</tr>
<tr>
<td>2</td>
<td>pH</td>
<td>0.595</td>
<td>0.224</td>
<td>.009</td>
<td>.037</td>
<td>.335</td>
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</table>

**T peakedness index**

<table>
<thead>
<tr>
<th>Order of entry</th>
<th>Variable</th>
<th>Coefficient</th>
<th>± SE</th>
<th>P value</th>
<th>ΔR²</th>
<th>Model R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>AG</td>
<td>0.102</td>
<td>0.050</td>
<td>&lt;.043</td>
<td>.240</td>
<td>.240</td>
</tr>
<tr>
<td>2</td>
<td>K</td>
<td>0.867</td>
<td>0.196</td>
<td>&lt;.001</td>
<td>.078</td>
<td>.318</td>
</tr>
<tr>
<td>3</td>
<td>pH</td>
<td>−4.580</td>
<td>1.808</td>
<td>.013</td>
<td>.033</td>
<td>.352</td>
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</table>

**Log10 transformed ST segment duration**

<table>
<thead>
<tr>
<th>Order of entry</th>
<th>Variable</th>
<th>Coefficient</th>
<th>± SE</th>
<th>P value</th>
<th>ΔR²</th>
<th>Model R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>K × K</td>
<td>0.006</td>
<td>0.001</td>
<td>&lt;.001</td>
<td>.266</td>
<td>.266</td>
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</tbody>
</table>

**ST segment amplitude**

<table>
<thead>
<tr>
<th>Order of entry</th>
<th>Variable</th>
<th>Coefficient</th>
<th>± SE</th>
<th>P value</th>
<th>ΔR²</th>
<th>Model R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>K × K</td>
<td>0.049</td>
<td>0.049</td>
<td>&lt;.001</td>
<td>.341</td>
<td>.341</td>
</tr>
<tr>
<td>2</td>
<td>K × K</td>
<td>−0.175</td>
<td>0.017</td>
<td>&lt;.001</td>
<td>.302</td>
<td>.644</td>
</tr>
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</table>

**ST segment angle**

<table>
<thead>
<tr>
<th>Order of entry</th>
<th>Variable</th>
<th>Coefficient</th>
<th>± SE</th>
<th>P value</th>
<th>ΔR²</th>
<th>Model R²</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>K × K</td>
<td>22.516</td>
<td>3.625</td>
<td>&lt;.001</td>
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<td></td>
</tr>
<tr>
<td>2</td>
<td>K × K</td>
<td>−8.639</td>
<td>1.194</td>
<td>&lt;.001</td>
<td>.168</td>
<td>.509</td>
</tr>
<tr>
<td>3</td>
<td>K × Na</td>
<td>0.054</td>
<td>0.019</td>
<td>.005</td>
<td>.028</td>
<td>.537</td>
</tr>
<tr>
<td>4</td>
<td>AG</td>
<td>0.107</td>
<td>0.041</td>
<td>.010</td>
<td>.024</td>
<td>.561</td>
</tr>
</tbody>
</table>

The variables K × pH, pH, and pCO2 were not retained in the final model.
The variables Na and K × Na, K × pH, and pCO2 were not retained in the final model.
The variables Na, K × Na, K × pH, Creatinine, and AG were not retained in the final model.
The variables Na, K × Na, K × pH, Creatinine, and AG were not retained in the final model.
The variables Na, K × Na, K × pH, Creatinine, and AG were not retained in the final model.
The variables Na, K × Na, K × pH, Creatinine, and AG were not retained in the final model.
The variables Na, K × Na, K × pH, Creatinine, and AG were not retained in the final model.
The variables pH, Na, K × pH, and Creatinine were not retained in the final model.
with hyperkalemia, acidemia, and diarrhea because of cholera. Dehydration and hemoconcentration are common in hyperkalemic diarrheic calves and the resultant hypovolemia decreases ventricular chamber volumes. The effect of heart chamber volume on the amplitude of Q, R, and S waves is called the Brody phenomenon, whereby the intracavitary blood mass augments the radial components and reduces the tangential components of the cardiac dipole. The Brody phenomenon explains why decreased ventricular chamber volumes result in alterations of Q, R, and S waves.

However, the Brody phenomenon would predict that Q, R, and S wave amplitude voltages in diarrheic dehydrated calves would decrease as plasma potassium concentration increased because of the decrease in plasma volume and consequently chamber volume, which is opposite to the relationship identified in the study reported here and elsewhere. We found a negative association between hypermagnesemia and S wave amplitude, which at $r_s = -0.59$ represented the strongest association between a serum biochemical analyte and an ECG variable in our study. Stepwise multivariable regression identified that the 2 independent predictors of S wave amplitude were log$_{10}$serum magnesium concentration and blood pH. Marked hypermagnesemia has been associated with ECG abnormalities such as tall T waves, prolonged PR intervals, and increased QRS duration and amplitude and consequently hypermagnesemia might contribute to the observed ECG abnormalities in neonatal diarrheic calves. Similarly, low blood pH can produce ECG abnormalities similar to those seen with hyperkalemia and consequently acidemia might also contribute to the observed ECG abnormalities in neonatal diarrheic calves.

FIGURE 6 Scatterplots illustrating the association between plasma potassium concentration and P amplitude, Ta amplitude, S amplitude, T amplitude, ratio of T amplitude to P amplitude and the ratio of T amplitude to S amplitude. The lines represent the result of simple linear regression (T amplitude) and segmented regression analysis (P amplitude, Ta amplitude, S amplitude, ratios of T amplitude to P amplitude and T amplitude to S amplitude). Gray-shaded areas represent the reference range for plasma potassium concentration (3.9–5.8 mmol/L).
with hyperkalemia, acidemia, and diarrhea because of cholera. Dehydration and hemoconcentration are common in hyperkalemic diarrheic calves and the resultant hypovolemia decreases ventricular chamber volumes. The effect of heart chamber volume on the amplitude of Q, R, and S waves is called the Brody phenomenon, whereby the intracavitary blood mass augments the radial components and reduces the tangential components of the cardiac dipole. The Brody phenomenon explains why decreased ventricular chamber volumes result in alterations of Q, R, and S wave amplitudes. However, the Brody phenomenon would predict that Q, R, and S wave amplitude voltages in diarrheic dehydrated calves would decrease as plasma K increased because of the decrease in plasma volume and consequently chamber volume, which is opposite to the relationship identified in the study reported here and elsewhere. We found a negative association between hypermagnesemia and S wave amplitude, which at represented the strongest association between a serum biochemical analyte and an ECG variable in our study. Stepwise multivariable regression identified that the 2 independent predictors of S wave amplitude were log10(serum magnesium concentration) and blood pH. Marked hypermagnesemia has been associated with ECG abnormalities such as tall T waves, prolonged PR intervals, and increased QRS duration and amplitude and consequently hypermagnesemia might contribute to the observed ECG abnormalities in neonatal diarrheic calves. Similarly, low blood pH can produce ECG abnormalities similar to those seen with hyperkalemia, and consequently acidemia might also contribute to the observed ECG abnormalities in neonatal diarrheic calves.

A recent study reported ST segment elevation in a high proportion of neonatal diarrheic calves with hyperkalemia, which was also commonly seen in calves of the present study (Table 2, Figure 4C-H). ST segment elevation was also described as a rare manifestation of hyperkalemia in humans (most frequently in patients with diabetic ketoacidosis) where the ECG findings can resemble acute myocardial infarction and have therefore been described as pseudoinfarct or pseudoinjury ECG pattern. The mechanisms for hyperkalemia associated ST segment elevation in humans are not well understood, but repolarization abnormalities and other concurrent metabolic abnormalities such as acidosis have been discussed.

Although typical ECG abnormalities were seen in hyperkalemic calves of our study population, bradycardia, or bradyarrhythmia was a rare finding in calves with increases in cK+. There was also no association between heart rate and cK+, which is in agreement to previous clinical studies in diarrheic calves as well as dogs and cats who also found no significant relationship or even a slight positive association. Arrhythmias were detectable in 26 calves of the present study population of which 20 calves had an irregular sinus rhythm. However, it should be considered that the presence of sinus arrhythmias does not necessarily represent a pathologic alteration as it was reported to be detectable in 10% of healthy Holstein calves in a recent study using 24-h ECG Holter monitoring. Nonsignificant or only weak to moderately significant coefficients of correlation between cK+ and specific ECG measurements such as QRS duration or T amplitude were found in our study (Supporting Information Table S1). Studies on the frequency of ECG changes in hyperkalemic dogs and cats and human patients reported a high variability of ECG manifestations and a low sensitivity of the ECG for diagnosing a hyperkalemic state. Also reports on cases of severe
hyperkalemia with minimal or no ECG manifestations in humans have
been published.66,67 This indicates that an increase of extracellular
potassium concentration per se is not the single most important factor
for the development of ECG manifestations associated with hyperkale-
mia. There is some evidence that the rate of rise of cK\(^+\) is also playing
a role in this respect\(^6\) and it is conceivable that some calves of our
study experienced a slow increase of cK\(^+\) such that compensatory
mechanisms probably allowed to stabilize the transmembrane resting
potential. However, an important finding of our study was that ECG
abnormalities such as tall and peak-shaped appearing T waves, deep S
waves, widening of QRS complex or ST segment elevation were also
seen in acidemic calves that were normo- or even hypokalemic. Similar
findings were reported in a study\(^25\) where tall and tent shaped appear-
ing T-waves in acidemic humans without hyperkalemia have been
documented. Experimental and in-vitro studies additionally suggested a
direct effect of acidemia on cardiac conduction. In dogs, ECG abnor-
malities were observed during different types of experimentally
induced acidosis that were similar to those of induced hyperkalemia,
even in the absence of a hyperkalemic response.\(^5\)\(^6\) Also a delayed atrio-
ventricular conduction was reported after exposing heart preparations
in an acidic extracellular solution.\(^5\)\(^9\)\(^,\)\(^70\)

However, in calves of the present study another issue needs to be
considered, as the transmembrane resting potential is not only depend-
et from alterations of extracellular but also from alterations of intracellular
cK\(^+\). Neonatal diarrheic calves have a negative potassium balance because of
intestinal losses and low milk intake,\(^4\) which is likely enhanced by alter-
ations of internal potassium balance because of an acidic state and
resulting intracellular losses of potassium ions. It is therefore conceivable
that in some calves an intracellular depletion of potassium stores and con-
comitant alterations of internal potassium balance because of acidemia
(leading to normokalemia) resulted in a similar extracellular to intracellular
potassium ratio than in calves with marked increases in cK\(^+\) but normal or
only slightly decreased intracellular K\(^+\). This would explain the presence
of ECG abnormalities in acidemic but non-hyperkalemic calves.

Although our study documented ECG findings in a large study
population of diarrheic calves with a broad range of acid-base and
potassium balance disorders, our analysis has also some limitations. A
potential limitation of our study represents the choice of bipolar limb
lead II for analysis, which was selected as it is a frequently used ECG
lead in small animal medicine. A previous study on ECG parameters in
healthy lactating Holstein cows, however, revealed lower amplitudes
and greater wave form variability of limb lead measurements (including
lead II) when compared to standard base-apex lead analyses.\(^7\)\(^1\)\(^-\)\(^7\)\(^4\) Those
differences can be explained by the positions of the electrodes relative to
the position of the heart in the thorax of the bovine species, a higher
susceptibility of limb leads to movement artefacts, and species specific
characteristics of the distribution of the conduction system within the
myocardium, which may cause cancellation of wave fronts in limb lead
measurements.\(^7\)\(^1\)\(^-\)\(^7\)\(^4\) However, in our study limb lead II measurements
produced clear waveforms and a comparative analysis of limb lead II
and base-apex lead did not reveal major differences in respect to ampli-
tudes, durations and variability of ECG parameters (Supporting Infor-
mation Table S2). In addition, analysis of ECG measurements in a
nonblinded manner might be considered as a limitation although the
use of a software program provided a standardized method of meas-
uring the amplitudes and durations of selected ECG complexes.

Another limitation of our study was the lack of a healthy control group.
Although calves of the reference groups had venous blood pH and
plasma cK\(^+\) in the reference range it is possible that other metabolic
conditions might have affected the ECG measurements in those calves.

5 CONCLUSIONS

Results of our study confirm that hyperkalemia is a clinically relevant
and potentially cardiotoxic electrolyte imbalance in neonatal diarrheic
calves. However, increased plasma cK\(^+\) only partly explained the
observed ECG abnormalities in our study population. Especially the
findings that venous blood pH was significantly associated with the
presence of specific ECG abnormalities such as tall and peak-shaped
appearing T waves, and that those abnormalities were also seen in
acidemic but non-hyperkalemic calves, indicate that acidemia is a con-
tributing factor that predisposes to cardiac conduction abnormalities in

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**TABLE 5** Results of a segmented linear regression analysis for modeling the relationship between 8 ECG variables and plasma potassium concentrations in calves of our study population

<table>
<thead>
<tr>
<th>Variable</th>
<th>n</th>
<th>Intercept</th>
<th>Slope 1</th>
<th>Break point [K]₁</th>
<th>95% CI for [K]₁</th>
<th>Slope 2</th>
<th>Break point [K]₂</th>
<th>95% CI for [K]₂</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ta wave amplitude</td>
<td>117</td>
<td>0.02</td>
<td>−0.02</td>
<td>5.7</td>
<td>5.0–6.4</td>
<td>0.03</td>
<td>8.4</td>
<td>6.7–10.1</td>
</tr>
<tr>
<td>P wave amplitude</td>
<td>117</td>
<td>0.01</td>
<td>0.04</td>
<td>6.5</td>
<td>6.1–6.9</td>
<td>−0.16</td>
<td>8.2</td>
<td>7.5–8.9</td>
</tr>
<tr>
<td>T to P amplitude ratio</td>
<td>109</td>
<td>3.18</td>
<td>0</td>
<td>6.6</td>
<td>6.2–6.9</td>
<td>5.53</td>
<td>−</td>
<td>−</td>
</tr>
<tr>
<td>S wave amplitude</td>
<td>117</td>
<td>−0.34</td>
<td>−0.25</td>
<td>7.4</td>
<td>6.0–8.9</td>
<td>0.34</td>
<td>−</td>
<td>−</td>
</tr>
<tr>
<td>QRS duration</td>
<td>117</td>
<td>38.1</td>
<td>5.44</td>
<td>7.8</td>
<td>7.4–8.1</td>
<td>38.27</td>
<td>−</td>
<td>−</td>
</tr>
<tr>
<td>J point amplitude</td>
<td>117</td>
<td>−0.05</td>
<td>0.01</td>
<td>7.9</td>
<td>7.5–8.3</td>
<td>0.12</td>
<td>−</td>
<td>−</td>
</tr>
<tr>
<td>T to S amplitude ratio</td>
<td>117</td>
<td>0.40</td>
<td>0</td>
<td>7.8</td>
<td>6.8–8.7</td>
<td>0.16</td>
<td>−</td>
<td>−</td>
</tr>
<tr>
<td>ST segment angle</td>
<td>117</td>
<td>−0.74</td>
<td>0.84</td>
<td>9.1</td>
<td>8.8–9.3</td>
<td>24.9</td>
<td>−</td>
<td>−</td>
</tr>
</tbody>
</table>

The slope in the 3rd model for P and Ta wave amplitude was 0.
Data from 13 hypokalemic calves were not included in the analysis.
diarrheic calves. We anticipate that application of the segmented regression approach used in the critically ill calves in the study reported here will provide additional insight into the sequence of ECG changes and facilitate the diagnosis and treatment of hyperkalemia in critically ill domestic animals and humans.

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CONFLICT OF INTEREST DECLARATION

Peter Constable has received funding from Boehringer-Ingelheim related to the treatment of hypokalemia in adult dairy cattle.

OFF-LABEL ANTIMICROBIAL DECLARATION

Authors declare no off-label use of antimicrobials.

INSTITUTIONAL ANIMAL CARE AND USE COMMITTEE (IACUC) OR OTHER APPROVAL DECLARATION

This study was approved by the Ethics committee of the Center of Veterinary Clinical Medicine, LMU Munich.

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SUPPORTING INFORMATION
Additional Supporting Information may be found online in the supporting information tab for this article.

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