ABSTRACT

Bovine respiratory disease (BRD) is a complex disease process and many reports emphasize the negative implications of clinical BRD in dairy calves. Early diagnosis can be difficult because of inconsistent or absent clinical signs; however, the use thoracic ultrasonography has the potential to improve detection of respiratory disease. Earlier detection of BRD may result in actions to improve calf welfare and production. The objective of this prospective cohort study was to determine if lung consolidation (LC) in young dairy calves influenced age at first calving (AFC), first-lactation milk production, and survival to the end of first lactation. A total of 215 female calves from 3 dairy herds in southwestern Ontario were enrolled and assessed weekly during their first 8 wk of life for evidence of LC using thoracic ultrasonography (Ibex Pro, Loveland, CO). Consolidation was measured, using gridlines on the screen of the ultrasound, in the first 10 intercostal spaces on both sides of the thorax. Calves were considered LC positive if ≥3 cm of consolidated lung was present. Multivariable linear regression models were used to identify risk factors associated with AFC and first-lactation 305-d milk production. A survival analysis was conducted to determine differences in survival from enrolment until the end of first lactation between calves with and without consolidation. In the study population, the following calfhood conditions were detected: twins (4%; n = 8), diarrhea in the first 21 d of life (31%; n = 66), rib fractures (7%; n = 14), lung abscesses (3%; n = 6), and at least one diagnosis of LC (57%; n = 123). Overall, 7% (n = 15) of calves died, and 18% (n = 38) of animals were sold before the end of first lactation. The presence of LC, at least once in the first 8 wk of life, did not influence AFC, but did result in a 525 kg (95% confidence interval: −992.81 to −60.25) decrease in first-lactation 305-d milk production. No difference in survival was detected between LC groups. These results indicate that LC during the first 56 d of life has a long-term effect on dairy calves, manifested as reduced milk production during first lactation.

Key words: bovine respiratory disease, milk production, thoracic ultrasound

INTRODUCTION

Bovine respiratory disease (BRD) is a complex, multifactorial disease process and early diagnosis can be difficult because of inconsistent or absent clinical signs (McGuirk, 2008). A wide range of bacteria and viruses interact with management, environment, and calf factors to cause BRD. It has been reported that 22% of dairy calves are treated for BRD before weaning, and almost 20% of those calves are treated more than once (Windeyer et al., 2014). Bovine respiratory disease in early life is associated with increased risk of mortality; decreased ADG, resulting in calves with BRD being 11 kg lighter at 6 mo compared with calves that did not have BRD; increased age at first calving (AFC); and reduced first-lactation milk production (van Der Fels-Klerx et al., 2002; Stanton et al., 2012).

Clinical scoring systems have been developed to assist in the detection of BRD in dairy calves, but no gold standard currently exists. Specifically, the Wisconsin Calf Scoring Chart partitions respiratory disease signs into 5 categories: rectal temperature, cough, nasal discharge, ocular discharge, and ear position (McGuirk, 2008). Each category is assigned a score ranging from 0 to 3 based on specific criteria, and an overall score is tabulated; a calf with a score ≥5 is considered clinically ill (McGuirk, 2008). In addition to the Wisconsin Calf Scoring System, a dichotomous scoring system, measuring similar clinical signs, has been developed (Love et al., 2014). Drawbacks related to these approaches in respiratory disease diagnosis include both the subjectivity of the scoring, and dependence on the presence of clinical signs.

Noninvasive diagnostic imaging techniques are available for determining the presence of BRD. Formal evalu-
Production, and survival to the end of first lactation. Young dairy calves influenced AFC, first-lactation milk productivity cohort follow-up study was to determine if LC in (Ollivett, 2014), the objective of the present prospective approach for the diagnoses of subclinical BRD (Ollivett and Buczinski, 2016). Comparisons have been made between TUS and clinical scoring (Buczinski et al., 2015). Although both methods are imperfect, TUS has demonstrated a greater sensitivity and specificity compared with clinical scoring (Buczinski et al., 2015).

Research evaluating lasting effects of lung consolidation (LC) on productivity of dairy animals is limited. Adams and Buczinski (2016) used a single TUS assessment on 3-mo-old Jersey heifers to determine the presence of LC. Calves with extensive LC or a lung abscess had a greater risk of being culled or dying, but no association was found between degree of LC and AFC. Teixeira et al. (2017) measured LC by TUS dichotomously in female calves at 60 d of age, and found no association with mortality in the first 350 d of life. However, calves that had LC were more likely to be removed from the herd between 305 d of age and first calving, and had reduced reproductive performance following first calving. No difference was detected in weekly average milk production in the first 3 mo of lactation.

A gap in knowledge currently exists in the relationship between lung lesions in calves and medium- to long-term performance outcomes. We hypothesized that LC will negatively affect the dairy calf and have lasting effects, which will be detected in delayed calving age and decreased first-lactation milk production. Although the initial purpose of this research was to determine the efficacy of a novel BRD vaccine in calves (Ollivett, 2014), the objective of the present prospective cohort follow-up study was to determine if LC in young dairy calves influenced AFC, first-lactation milk production, and survival to the end of first lactation.

MATERIALS AND METHODS

Preweaning Management

Calves were raised on 3 separate dairy farms in southwestern Ontario, Canada, and enrolled between January and December 2012. Elora Dairy Research Center [herd 1, n = 150 lactating cows, average first-lactation 305-d milk production (305M) = 8,567 ± 1,314 kg (mean ± SD)], a privately owned commercial dairy farm (herd 2, n = 650 lactating cows, average 305M = 8,595 ± 1,541 kg), and Ponsonby Dairy Research Center (herd 3, n = 55 lactating cows, average 305M = 9,028 ± 1,053 kg) were included in the study.

In herd 1, calves were fed 4 L of single-source colostrum within 12 h of birth. Calves were housed either within individual stalls in an enclosed room or outside in individual hutches. Feeding management included feeding 6 L of whole unpasteurized milk daily until approximately 6 wk of age, when calves were gradually weaned and moved to group housing by 8 wk of age. Bull and heifer calves were housed together. Calves had access to calf starter and water beginning at 3 d of age.

In herd 2, 2 L of single-source colostrum was offered to calves within 30 min of birth, with an additional 4 L offered in 2 feedings over the 24 h after birth. Calves were fed 6 L of whole unpasteurized milk twice daily, and housed in individual pens. At 3 wk of age, calves were moved as a group into a pen containing 20 calves. Here, calves were fed 8 L/d of whole unpasteurized milk via an automated feeder (DeLaval Canada, Peterborough, ON, Canada), and had access to free choice water. Calves had access to free choice starter from 3 d of age.

In herd 3, 4 L of single-source colostrum was fed to calves within 12 h of birth, followed by 2 L of whole unpasteurized milk fed twice daily. At approximately 8 wk of age, all calves were weaned. All calves were housed outdoors in individual hutches until weaning, when they were moved to group housing. Both water and calf starter were offered free choice beginning at 3 d of age.

Herd 2 calves were enrolled twice weekly. Both male and female Holstein calves were enrolled between 3 and 6 d of age in a vaccine trial (Ollivett, 2014). Briefly, calves were assigned to 1 of 3 vaccine groups: an intranasal vaccine (Inforce 3, Zoetis, Florham Park, NJ), subcutaneous vaccine (Bovi-Shield Gold 5, Zoetis), or intranasal and subcutaneous saline (placebo). All calves were followed for 8 to 12 wk after enrolment; only female calves were followed postweaning.

Ultrasonography and Health Data Collection

Whole blood samples were collected to assess passive transfer of maternal antibodies, in 3- to 6-d-old calves by jugular venipuncture, using a 20-gauge, 1-inch (2.54 cm) hypodermic needle (BD Vacutainer Precision Glide, Becton Dickinson and Co., Franklin Lakes, NJ). Samples were collected into a sterile, plastic, commercial blood collection tube without anticoagulant (BD Vacutainer, Becton Dickinson and Co.) and stored on ice. Within 4 to 6 h of collection, serum was separated by centrifugation at 970 × g for 15 min at approximately 20°C. Serum total protein (STP) was analyzed with
a digital Brix refractometer (Misco PA202X-003–105, Cleveland, OH).

A portable linear ultrasound set at a depth of 9 cm, frequency of 6.2 MHz, and gain of 16 dB (Near 13 dB; Far 36 dB; Ibex Pro; E. I. Medical, Loveland, CO) was used to perform TUS on calves weekly (Ollivett et al., 2015). A single researcher performed all TUS exams at each visit. The thorax of calves was not shaved or clipped before performing ultrasound examinations. Approximately 300 mL of 70% isopropyl alcohol was used as a transducing agent and was applied to the hair of the thorax (Ollivett et al., 2015). Both sides of the thorax were scanned, beginning dorsally at the level of the scapula in the 10th intercostal space (ICS), advancing cranially toward the right 1st ICS or the left 2nd ICS (Ollivett et al., 2015). This assessment was completed by holding the ultrasound probe parallel to the rib and advancing the probe from the dorsal to ventral aspect of the ICS, allowing for visualization of specific ultrasound landmarks (Ollivett et al., 2015). To view the lung adjacent to the right 4th through 1st ICS and left 4th through 2nd ICS, the probe was placed between the forelimb and the body wall (Ollivett et al., 2015).

Clinical respiratory scoring (McGuirk, 2008) was performed at each visit. The respiratory scoring system assigned a score of 0–3 to 5 different health parameters, including rectal temperature, nasal discharge, cough, and ocular discharge or ear position. After the individual parameters were assessed, the scores were totaled and could range from 0 to 12. When both ocular discharge and ear position scored ≥1, only the highest score was included (McGuirk, 2008). Any calf with a score >4 was considered clinically ill (McGuirk, 2008).

For lung tissue to be considered normal (LC−), a hyperechoic line and reverberation artifact were required. These characteristics are indicative of a contrast between the high impedance of the thoracic body wall and the low impedance of normal lung tissue (Blond and Buczinski, 2009). In contrast, when consolidation was present (LC+) the lung appeared hypoechoic and both the bright white band at the pleural interface and reverberation artifact were absent. To measure the extent of LC, the 1-cm gridlines on the ultrasound screen were used. A digital voice recorder was used to record all observations, which were later manually transcribed into a database (Microsoft Access 2010, Microsoft Corp., Redmond, WA).

Postweaning

In herd 1, approximately 2 wk after weaning, calves were group housed in a naturally ventilated bedded pack barn and offered hay in addition to calf starter. Breeding began at approximately 13 mo of age using AI exclusively. After calving, cows were milked twice daily in a parlor and fed a TMR diet.

Calves in herd 2 were housed in a combination of freestall and bedded pack pens following weaning, and the calves’ diet was composed of calf starter and hay. Beginning at 6 mo of age, heifers were fed the same diet as the lactating cows. Breeding began at approximately 12 mo of age, and heifers were bred using a combination of sexed semen and a herd bull. After calving, heifers were milked twice daily in a rotary parlor; at 10 DIM cows were milked 4 times daily in the same parlor for the remainder of the lactation.

In herd 3, calves were moved into a bedded pack barn, attached to the lactating cow facility, 2 wk after weaning. Here, the calves were raised in groups and fed hay and calf starter. Heifers were bred, beginning at 13 mo of age, using AI. Following calving, cows were housed in a tiestall and milked twice daily using a pipeline milking system and fed a TMR diet.

Statistical Analyses

Farm records were obtained from the herds’ computer software (DairyComp 305, VAS, Tulare, CA). Data retrieved from DairyComp included the date of first calving, postweaning diseases, postpartum diseases (metritis, retained placenta, ketosis, and mastitis), first-lactation milk production, and removal from the herd (date of sale or death). Age at removal and age at parturition were back calculated using birthdate and date of the event.

Continuous outcomes included AFC and completed 305M. Dichotomous variables for calf events included twin (yes = 1, no = 0), scour (fecal score in the first 21 d <3 = 0, fecal score = 3 during at least one exam in first 21 d = 1), LC [<3 cm of consolidated lung on all ultrasound examinations = 0 (LC−), 1 or more examinations with ≥3 cm consolidated lung = 1 (LC+)], dystocia (no assistance or easy pull = 0, hard pull or surgical delivery = 1), failure of passive transfer (STP ≥5.2 g/dL = 0, STP <5.2 g/dL = 1), rib (rib fracture not palpable or visible on ultrasound = 0, rib fracture palpable or visible on ultrasound = 1), abscess (regions in the lung that contained both fluid and well-defined hypoechoic areas; no = 0, yes = 1), and illness (ILL; respiratory score of ≤4 = 0 or >4 = 1). The cutoff of 3 cm for LC was determined by selecting the values between the 90th and 95th percentiles of ultrasound consolidation in calves with any amount of consolidation in their first ultrasound examination.

All statistical analyses were conducted using a standard statistical package (Stata 14, StataCorp LLC, College Station, TX). Sample size was originally calcu-
lated to satisfy requirements for the primary purpose of this study, which was to determine efficacy of a novel vaccine, and address short-term implications of respiratory disease (Ollivett, 2014). As the present study was completed as a follow-up to the original study, all female calves that underwent TUS in the original study were enrolled, and no further sample size calculations were made. The primary outcome of interest was 305M, and all variables that resulted in \( P < 0.20 \) in univari-able linear regression were entered as main effects in a multivariable linear regression model. The variable ILL was not entered as a predictor variable in the models because LC was the predictor variable of interest, and ILL was considered to be an intervening variable. Man-ual backward stepwise elimination was used to refine the model to include only variables that were significant at the \( P < 0.05 \) level. Prior to the final elimination of a variable, a change of \( >20\% \) of the estimates of all other variables was used to assess for confounding. A Shapiro-Wilk test and a Cook-Weisberg test were completed to test for normality and homoscedasticity of residuals, respectively. All plausible interactions were tested. Linearity of continuous predictor variables were tested using locally weighted regression. A model was built in the same manner to address the outcome AFC, and LC was forced into the model, regardless of significance, as this was the predictor variable of interest.

To address the secondary objective, a survival analysis was competed using a Cox proportional hazard ratio and a Kaplan-Meier survival estimate to assess the effect of LC on survival. Animals were censored when they left the herd regardless of reason (animals who were sold or died), or right censored when they completed their 305-d first lactation. To standardize first lactation production, animals were considered to have finished their first lactation at 305 DIM. Schoenfeld residuals were used to statistically test the assumption of proportional hazards.

**RESULTS**

A total of 215 female Holstein dairy calves were enrolled and followed forward in time. In the study population, the following calfhood conditions were detected: twins (4%; \( n = 8 \)), diarrhea in the first 21 d of life (31%; \( n = 66 \)), rib fractures (7%; \( n = 14 \)), lung abscesses (3%; \( n = 6 \)), at least one diagnosis of ILL (58%, \( n = 124 \)), and at least one diagnosis of LC (57%; \( n = 123 \)). Distribution of consolidation detection by week of assessment is summarized in Figure 1. Consolidation was detected as early as the second assessment, indicating calves were developing consolidation as early as 12 d of age. Both consolidation and a positive clinical score was detected in 43% (\( n = 92 \)) of calves. Death occurred in 2% (\( n = 4 \)) of calves before 56 d of age; therefore, 211 calves remained for postweaning follow-up.

An AFC was retrieved for 88% (\( n = 183 \)) of the animals, and was measured in months. No difference was observed in the odds of calving between LC+ and LC− calves [odds ratio: 0.65 (95% CI: 0.35 to 1.23)]. In the final multivariable model, failure of passive transfer, vaccine type, and herd were significant and were therefore retained in the model (Table 1). No significant association was observed between LC and AFC (\( P = 0.553 \)).

Complete lactation data were obtained for 66% (\( n = 140 \)) of animals, and average milk production for the study population was 8,626 ± 1,427 L (mean ± SD). Age at first calving and LC were significant in the final multivariable model of 305M (Table 2). An increase in 305M of nearly 140 kg (95% CI: 33 to 241 kg; \( P = 0.01 \)) was associated with a 1-mo increase in AFC. In contrast, LC+ was associated with a 525 kg (95% CI: −993 to −60 kg; \( P = 0.03 \)) decrease in first-lactation milk production.

Overall, 7% (\( n = 15 \)) of heifers died, and 18% (\( n = 38 \)) of animals were sold before the end of first lactation. In the LC− group 5% of calves died, whereas 9% of LC+ calves died; however, this difference was not statistically significant (\( P = 0.33 \)). There was no difference (\( P = 0.42 \)) in the proportion of calves sold, with 15% of LC− and 20% of LC+ calves being sold before the end of first lactation. The hazard of leaving the herd before the end of first lactation was not significantly different between LC+ and LC− calves [hazard ratio = 1.3; 95% CI (0.7 to 2); \( P = 0.368 \); Figure 2].

**DISCUSSION**

The purpose of this study was to assess the effects of LC in young Holstein dairy heifers on their survival and first-lactation milk production. Thoracic ultrasonography was used to diagnose calves as having consolidation (LC+), using a cutoff of 3 cm of consolidated lung tissue. No difference was observed in AFC or survival until the end of first lactation between consolidation groups (Table 1; Figure 2). However, milk production was significantly lower in LC+ calves.

Clinical respiratory scoring is a more common practice among veterinarians and dairy producers, and is less expensive than TUS because ultrasound equipment is not required. Using clinical signs alone may fail to identify lung lesions, which was shown to affect more than 60% of calves during the preweaning period in one herd (Ollivett and Buczinski, 2016). Given that only 43% of calves had both a clinical respiratory score and consolidation supplies further evidence that clinical scoring alone fails to identify all calves with consolida-
Figure 1. Frequency distribution of lung consolidation by assessment week in 123 female Holstein calves diagnosed with lung consolidation by thoracic ultrasound in the first 56 d of life. Lung scores were dichotomized to no consolidation (<3 cm consolidation) or consolidation (≥3 cm consolidation).

Table 1. Multivariable linear regression model of the assessment of lung consolidation with age at first calving (mo of age) for 183 Holstein heifer calves in 3 herds in southwestern Ontario

<table>
<thead>
<tr>
<th>Variable</th>
<th>Coefficient</th>
<th>SE</th>
<th>P-value</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Herd 1(^1)</td>
<td>Referent</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Herd 2(^2)</td>
<td>0.47</td>
<td>0.46</td>
<td>0.30</td>
<td>−0.43, 1.38</td>
</tr>
<tr>
<td>Herd 3(^3)</td>
<td>−1.16</td>
<td>0.67</td>
<td>0.09</td>
<td>−2.49, 0.16</td>
</tr>
<tr>
<td>FPT(^4)</td>
<td>−0.97</td>
<td>0.69</td>
<td>0.17</td>
<td>−2.34, 0.40</td>
</tr>
<tr>
<td>Injectable vaccine(^5)</td>
<td>Referent</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Intranasal vaccine(^6)</td>
<td>0.14</td>
<td>0.35</td>
<td>0.68</td>
<td>−0.54, 0.82</td>
</tr>
<tr>
<td>Placebo vaccine(^7)</td>
<td>−0.43</td>
<td>0.58</td>
<td>0.48</td>
<td>−1.57, 0.74</td>
</tr>
<tr>
<td>Consolidation(^8)</td>
<td>0.20</td>
<td>0.34</td>
<td>0.55</td>
<td>−0.47, 0.88</td>
</tr>
<tr>
<td>Intercept</td>
<td>25.59</td>
<td>0.41</td>
<td>&lt;0.01</td>
<td>24.78, 26.40</td>
</tr>
</tbody>
</table>

\(^1\)Elora Dairy Research Center, Ontario, Canada.
\(^2\)Commercial dairy in southwestern Ontario, Canada.
\(^3\)Ponsonby Dairy Research Center, Ontario, Canada.
\(^4\)Failure of passive transfer as identified by serum total protein <5.2 mg/dL.
\(^5\)Injectable vaccine (Bovi-Shield Gold, Zoetis Canada, Kirkland, Quebec, Canada) = 2 mL of commercial multivalent vaccine against bovine respiratory syncytial virus, infectious bovine rhinotracheitis, parainfluenza virus-3, and bovine viral diarrhea administered to calves subcutaneously at 6 wk of age.
\(^6\)Intranasal vaccine (Inforce 3, Zoetis Canada, Kirkland, Quebec, Canada) = 2 mL of commercial trivalent vaccine against bovine respiratory syncytial virus, infectious bovine rhinotracheitis, and parainfluenza virus-3 administered to calves 3 to 6 d of age.
\(^7\)Placebo vaccine = 2 mL of sterile saline administered both intranasally and subcutaneously at 3 to 6 d of age and 6 wk of age.
\(^8\)≥3 cm of lung consolidation in at least one ultrasound exam during the first 8 wk of life.
tion. Results from the present study indicate that ultrasonographic lung lesions develop as early as 12 d of age (Figure 1) and have a long-term effect on the dairy calf. Early interventions may be warranted to try to mitigate the associated milk loss.

Compared with previous work, the prevalence of LC detected in this study is higher, potentially due to more frequent ultrasound examinations performed throughout the preweaning period. Performing only one TUS exam at weaning, or when calves leave custom rearing (Adams and Buczinski, 2016; Teixeira et al., 2017), is likely to underestimate the number of calves affected with LC during the preweaning period.

Contrary to expectations, LC in the first 8 wk of life was not associated with an increase in AFC among those that calved. This finding agrees with that of Adams and Buczinski (2016), who also reported no association between LC and AFC. The inclusion of calves from only 3 herds may have influences observed results, and is a limitation to this study.

No differences were noted in survival between LC+ animals and LC− animals; however, milk production in the first lactation differed between the 2 groups. This contrasts with previous results reported by Adams and Buczinski (2016) and Teixeira et al. (2017). Study design may have played a role in observed results because previous studies only took one measurement of LC around weaning. Another reason that milk production differences in this study may differ from previous work is complete first-lactation milk production data were used, and a small number of animals left the herds before the end of first lactation. Teixeira et al. (2017) used partial lactation data and found no difference in production. These apparently conflicting results may in fact be different manifestations of the same phenomenon. It is possible that in the Teixeira et al. (2017)

<table>
<thead>
<tr>
<th>Variable</th>
<th>Coefficient</th>
<th>SE</th>
<th>P-value</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age at first calving (mo)</td>
<td>137.12</td>
<td>52.53</td>
<td>0.01</td>
<td>33.25, 240.98</td>
</tr>
<tr>
<td>Consolidation1</td>
<td>−526.53</td>
<td>235.80</td>
<td>0.03</td>
<td>−992.81, 0.25</td>
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<td>Intercept</td>
<td>5,325.62</td>
<td>1,358.28</td>
<td>&lt;0.01</td>
<td>2,639.72, 8,011.52</td>
</tr>
</tbody>
</table>

1≥3 cm of lung consolidation in at least one ultrasound exam during the first 8 wk of life.
study, the animals exhibiting effects related to early LC were culled from the herd early in first lactation, and therefore not able to express their decreased milk production, whereas a management decision made to retain all first-lactation animals until the end of their lactation allowed the animals in the current study to express the production loss. It is also important to note that no difference in milk production seen in past research may be related to differential misclassification bias. Because past research only took one measurement of LC, it is possible that some truly diseased calves were classified as healthy. This would have biased results toward the null, supporting the finding of no difference in milk production.

Currently, no gold standard exists for the diagnosis of respiratory disease. However, LC can be rapidly detected using the same portable ultrasound unit and linear probe that is used routinely for bovine reproductive examinations. The use of regular TUS exams by veterinarians should be considered, given the long-term implications seen in this study. This practice would allow for earlier and more accurate detection of BRD. Additionally, TUS may prove useful for troubleshooting poor heifer performance and to identify risk factors for respiratory complications. Because little is currently known about which treatments and management practices may be effective in reducing the incidence, severity, or duration of LC, it may be beneficial to conduct intervention studies in the future to determine if the lasting effects of LC can be mitigated.

CONCLUSIONS

Diagnosing LC in the first 56 d of life has a significant long-term effect on the calf, manifested as reduced milk production in first lactation. The regular use of calf-side TUS for diagnosis of LC should be considered, given the prevalence and lasting effects seen. Further research, with increased sample sizes and variation in housing and management strategies, should be conducted to confirm the findings of the present study. In addition, future research looking at methods to mitigate the effects of LC may be warranted.

ACKNOWLEDGMENTS

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REFERENCES


