Field trial to evaluate the effect of an intranasal respiratory vaccine protocol on calf health, ultrasonographic lung consolidation, and growth in Holstein dairy calves

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ABSTRACT

The objective of this field trial was to evaluate the effect of a vaccine protocol using a commercially available trivalent vaccine designed for intranasal use. Experimental challenge studies have demonstrated varying efficacies of vaccines administered via the intranasal route. A total of 468 calves from 3 herds were enrolled and randomized into 3 treatment groups (positive control, PC, $n = 211$; intranasal vaccine, IN, $n = 215$; negative control, NC, $n = 42$) and followed for 8 to 12 wk. The PC consisted of one dose of commercially available multivalent injectable vaccine against bovine respiratory syncytial virus, infectious bovine rhinotracheitis, parainfluenza 3, and bovine viral diarrhea administered subcutaneously at 6 wk of age. The IN was administered at enrollment and 6 wk of age, and contained antigen against bovine respiratory syncytial virus, infectious bovine rhinotracheitis, and parainfluenza 3. The NC was sterile saline administered intranasally and subcutaneously at enrollment and 6 wk of age. Clinical illness was assessed using systematic respiratory scoring, and thoracic ultrasonography was used to identify the lung consolidation associated with pneumonia. Rib fractures were identified in 6% of calves, and an association was observed between rib fractures and calving ease. Overall, 54% of the calves had at least one episode of an abnormal respiratory score (ILL). Vaccination protocol did not affect the occurrence of ILL. Similarly, 54% of the calves had at least one episode of lung consolidation ≥ 3 cm (CON). Vaccine protocol affected the odds of CON. The odds of CON in PC were 1.63 (95% confi-

dence interval: 1.04–2.56) times the odds of CON in IN, and 0.38 (95% confidence interval: 0.16–0.93) times the odds of CON in NC. The odds of CON in IN were 0.23 (95% confidence interval: 0.09–0.59) times the odds of CON in NC. The outcomes ILL and CON were associated; however, the measure of agreement was only fair ($\kappa = 0.38$). Multivariable linear regression revealed an interaction between vaccine protocol and herd on average daily gain (ADG); therefore, these data were stratified. In herd 1, IN (0.53 ± 0.03 kg/d) decreased ADG compared with PC (0.63 ± 0.03 kg/d). In herd 2, IN increased ADG (0.41 ± 0.03 kg/d) compared with PC (0.38 ± 0.03 kg/d). In contrast, none of the protocols affected ADG at herd 3. In conclusion, this commercially available trivalent IN vaccine protocol did not alter the incidence of ILL, reduced the risk of lung lesions associated with pneumonia, and improved the ADG of the calves in one of the commercial study herds.

Key words: bovine respiratory disease, intranasal vaccine, thoracic ultrasound

INTRODUCTION

Calves are born agammaglobulinemic and must ingest maternal colostrum for immune support. Although maternal transfer of antibody to the newborn calf provides many great benefits (Faber et al., 2005), high levels of maternal antibodies are associated with delayed antibody production by the neonate, as well as selective inhibition of lymphocyte responses (Tizard, 2013). The potential for maternal blockade has caused concern regarding the practice of early life vaccination to prevent bovine respiratory disease (BRD), as maternal antibodies can be present for up to 6 mo of age (Menanteau-Horta et al., 1985). Vaccination to reduce the incidence of BRD would be beneficial, as BRD is a

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leading cause of morbidity and mortality in preweaned dairy calves (Windeyer et al., 2014).

It has been established that 3- to 8-d-old Holstein calves are capable of mounting a mucosal immune response, even when maternal antibodies are present (Hill et al., 2012). Over the last 10 yr, reports have been inconsistent regarding the potential for intranasal vaccination to protect young dairy calves, regardless of maternal antibody status, from infection with bovine respiratory syncytial virus (**BRSV**) and parainfluenza 3 (**PI₃**). In one study, calves between 3 and 8 d of age were vaccinated intranasally with a product intended for subcutaneous administration, and challenged with BRSV 21 d later, or at 4.5 mo of age (Ellis et al., 2010). After the delayed challenge at 4.5 mo, seropositive calves had wider temperature fluctuations compared with seronegative calves; however, no differences were detected in other health parameters. Seronegative calves that were vaccinated intranasally with a low dose of BRSV antigen, and challenged with BRSV 21 d later, had less extensive lung lesions at postmortem examination, even though no differences in clinical scores or mortality were noted (Ellis et al., 2010). Intranasal vaccination in 3- to 8-d-old calves resulted in improved partial pressure of oxygen, fewer lung lesions, and lower mortality rate than in unvaccinated calves, when they were challenged with BRSV at 9 wk, but not 14 wk, indicating that the duration of immunity is short (Ellis et al., 2013). Additionally, 2 doses of monovalent injectable BRSV product used intranasally resulted in complete protection from clinical disease, one dose resulted in minimal clinical signs, whereas unvaccinated calves experienced severe clinical signs (Ellis et al., 2007).

The previously mentioned challenge studies often found that IN calves had less extensive lesions associated with lung consolidation than control calves despite the lack of observable clinical changes. In the beef industry, several reports suggest that evaluation of lung lesions at harvest may be a more accurate means of documenting BRD than clinical observations (Wittum et al., 1996; Thompson et al., 2006; White and Renter, 2009). Unfortunately, documenting lung lesions directly requires euthanasia, which often limits the size of study populations. As an alternative, thoracic ultrasonography (**TUS**) can be performed quickly and provides an accurate ante-mortem assessment of lung health (Rabeling et al., 1998; Ollivett et al., 2013, 2015). When lung consolidation identified by TUS was confirmed by the gold standard postmortem examination, TUS manifested a sensitivity of 85 and 94% in chronic clinical (Rabeling et al., 1998) and acute subclinical (Ollivett et al., 2015) cases, respectively; and specificity was 98 and 100%, in chronic clinical and acute subclinical cases, respectively. Additionally, Buczinski et al. (2015) using

Bayesian latent class analysis determined the sensitivity and specificity to be 79.4 and 93.9%, respectively.

The primary objective of this randomized controlled field trial was to evaluate the effect of an intranasal vaccine protocol on the health of young Holstein dairy calves. The secondary objectives were to evaluate the effect of this vaccine on ultrasonographic lung consolidation and growth.

MATERIALS AND METHODS

Animals and Facilities

This study was carried out on 3 dairies in southwestern Ontario, Canada, between January and December 2012. Two of these herds were the Elora (herd 1) and Ponsonby (herd 3) Dairy Research Centres associated with the University of Guelph. The third dairy was a privately owned commercial herd (herd 2). Each herd was visited twice a week (all herds) or 3 times a week during periods of high enrollment (herd 2 only) to enroll calves twice a week. Male and female Holstein calves were enrolled between 3 to 6 d of age into 3 groups according to vaccine protocol (positive control, **PC**; intranasal vaccine, **IN**; and negative control, **NC**) and followed for 8 to 12 wk. A birth record was filled out by the dairy producer after the birth of each calf. The PC (Bovi-Shield Gold 5, Zoetis, New York, NY) consisted of one dose of commercially available multivalent injectable vaccine against BRSV, infectious bovine rhinotracheitis, **PI₃**, and bovine viral diarrhea administered by subcutaneous injection at 6 wk of age. The IN (Inforce 3, Zoetis) was administered twice (first dose: 3 to 6 d of age; second dose: 6 wk of age) and contained BRSV, infectious bovine rhinotracheitis, and **PI₃** antigens. The NC, sterile saline, was administered both intranasally and subcutaneously twice (first dose: 3 to 6 d of age; second dose: 6 wk of age). All doses of IN were administered via a single-use plastic nasal cannula (Zoetis) into one nostril. All injections were administered subcutaneously in the neck. Treatments were administered by members of the research team not involved in respiratory scoring or TUS. Different randomization methods were used to assign calves to the 3 treatment groups at each herd because of differences in barn design. Treatment groups were housed in separate areas to prevent contamination of PC and NC calves from the potential nasal virus shedding by IN calves following vaccination. Weight, respiratory score (**RS**), and TUS observations were performed by the principal investigator (TLO) who was blinded to treatment throughout all data collection. This study was conducted with the approval of the University of Guelph's Animal Care Committee (AUP #11R110).

Herd-Specific Calf Management

In herd 1 (lactating cows, $n = 150$), calves were fed 4 L of single source colostrum within 24 h of birth. Calves were housed in either individual stalls within an enclosed room with a mechanical ventilation system, or outside tethered to individual plastic hutches. Calves were fed 6 L of whole unpasteurized milk per day until approximately 6 wk of age, at which point they were gradually weaned and moved to group housing by 8 wk of age. Free choice water and calf starter were available beginning at 3 d of age. Prior to the start of the study, rotating treatment groups were assigned to each room (enclosed nurseries, $n = 3$; outdoor "hutch room," $n = 1$) by drawing the protocol name from a hat with replacement. Four cycles were drawn for each room to accommodate the number of calves for the anticipated duration of the study. Each room was filled with 8 to 10 calves over a 2-wk period. Calves were housed together according to birth order regardless of sex. Each room was cleaned, disinfected, and allowed to sit empty for approximately 1 wk before new calves were added.

In herd 2 (lactating cows, $n = 650$), single source colostrum (2 L) was offered by bottle within 30 min of birth. An additional 4 L was offered in 2 separate feedings over the 24 h following birth. Calves were fed 6 L of unpasteurized whole milk twice daily while housed in individual pens until approximately 3 wk of age. Calves were then moved as a group of 20 animals per pen. Eight liters of unpasteurized whole milk was offered via an automated system in the group pens (Forster Technik, DeLaval, Peterborough, ON, Canada). Calves were allowed 3 L per feeding within a 3-h period. Free choice starter was offered within the first 3 d of life. Water was not available until the calves reached the group pen. Individual and group pens were bedded with a sawdust base covered by a top layer of straw. Calves were removed from the group pen at approximately 8 wk of age. Calves spent the first 24 to 36 h of life in a straw-bedded room adjacent to the maternity area before moving to 1 of 2 identical recently built barns. Each barn held 40 calves in individual stalls and 80 calves split between 4 group pens for a total of 120 calves per barn. Each barn was curtain sided and used both natural and positive pressure ventilation systems. As calves were born, individual stalls were filled in one barn, followed by the second barn. The owners of this commercial facility were not willing to have a NC group; therefore, calves were only enrolled into PC and IN groups. Treatment groups were randomly assigned at the barn level before the start of the study by picking the protocol from a hat.

In herd 3 (lactating cows, $n = 55$), calves were fed 4 L of single source colostrum within 24 h of birth, fol-

lowed by 2 L of unpasteurized whole milk 3 times daily until abrupt weaning at approximately 8 wk of age. All calves were housed outside tethered to individual plastic hutches until weaning, at which point they were moved to group housing. Free choice water and calf starter were available beginning at 3 d of age. Calves were blocked by sex, and treatment was assigned to each hutch by drawing protocol names from a hat in sets of 3 without replacement.

Sample Collection, Weighing, Health Scoring, and Ultrasonographic Data Collection

For assessing passive transfer of maternal antibodies, whole blood was collected from 3- to 6-d-old calves at enrollment, 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), into sterile, plastic, commercial blood collection tubes without anticoagulant. Blood tubes were stored on ice and serum was separated by centrifugation at $1,500 \times g$ for 15 min at $\sim 20^\circ\text{C}$, within 4 to 6 h of collection. Analysis of serum total protein (**STP**) was performed by a research assistant using a digital refractometer (Misco PA202X-003-105, Cleveland, OH). Also at enrollment, a 1- to 2-cm skin sample (ear notch) was collected from the ventral edge of the pinna to determine the bovine viral diarrhea persistent infection status for each calf using antigen-capture ELISA. Ear notch samples were placed in sterile, plastic, commercial blood collection tubes without anticoagulant, stored on ice, and refrigerated within 4 to 6 h of collection before submission to the University of Guelph's Animal Health Laboratory.

At each examination, calves were weighed 3 times using a weigh tape (Coburn Company, Whitewater, WI), and these weights were averaged to provide the weight for each examination. Average daily gain during the study period was calculated by dividing the difference in the weight recorded 56 d after enrollment and enrollment by 56. Calves that died before 56 d of age were excluded from the ADG analysis.

Respiratory scoring (Lago et al., 2006; McGuirk and Peek, 2014) and TUS were performed at each visit. Briefly, the RS assigned 0 to 3 points for each of the following categories: rectal temperature, nasal discharge, cough, and ocular discharge or ear position. Respiratory scores could range from 0 to 12 and any calf with $\text{RS} > 4$ was considered sick (McGuirk and Peek, 2014). Fecal scores (**FS**) were obtained after each RS by direct visualization of fresh manure. Digital examination of the rectum was used to stimulate defecation. Fecal scores were based on a 3-point scale: 1 = sample is in "patty" form; minimal water content, does not flow across or

down a surface; 2 = sample is more of a puddle, some water content, flows slowly across or down a surface; 3 = sample is watery, flows across or down a surface while leaving some to no adherent material (Ollivett et al., 2009). Regarding TUS, both lungs were scanned and images were interpreted as previously described (Ollivett et al., 2015). The depth and dorsal to ventral extent of consolidated lung were measured within each intercostal space using the 1-cm grid lines on the ultrasound screen. Observations and measurements were spoken and recorded using a digital voice recorder and later manually transcribed into a database (Microsoft Access 2010, Microsoft Corp., Redmond, WA).

Statistical Analyses

In addition to the 3 treatment groups, several potential explanatory covariates were investigated. Continuous variables included age at enrollment in days (**EAGE**), age at first TUS examination in days, and weight at first ultrasound in kilograms (**W1**). Nondichotomous categorical variables included treatment (1 = PC; 2 = IN; 3 = NC), herd (1 = herd 1; 2 = herd 2; 3 = herd 3), season of enrollment (winter = January through March; spring = April through June; summer = July through September), birth weight category (0 = $W1 < 40$ kg; 1 = $40 \text{ kg} \leq W1 < 46$ kg; 2 = $W1 \geq 46$ kg), and exam (chronologically ordered number of TUS exam). Dichotomous variables included **ILL** (all $RS \leq 4 = 0$; 1 or more $RS > 4 = 1$); **CON** (< 3 cm TUS lung consolidation on at all examinations = 0; 1 or more TUS examinations with ≥ 3 cm TUS lung consolidation present = 1); sex (male = 0; female = 1); twin (no = 0; yes = 1); dystocia (**DYST**; calving with no assistance or easy pull = 0, calving with hard pull or surgical delivery = 1); failure of passive transfer (**FPT**; $STP \geq 5.2$ g/dL = 0; $STP < 5.2$ g/dL = 1, Tyler et al., 1996); housing (outdoors = 0; indoors = 1); scour ($FS < 3$ within 21 d of enrollment = 0; $FS = 3$ during at least one exam within 21 d of enrollment = 1); and rib fracture (**RIBF**; rib fracture not palpable or visible on TUS = 0; rib fracture palpable and visible on TUS = 1). The 3-cm cut-off value for CON was determined by selecting the value between the 90th and 95th percentiles (2.25 and 3.5 cm, respectively) of TUS lung consolidation in those calves with any amount of lung consolidation during their first TUS examination. The scour variable was intended to capture those calves having at least one bout of severe diarrhea during the high-risk period of the first 21 d of life.

A standard statistical package was used for all analyses (SAS version 9.4, SAS Institute Inc., Cary, NC), except for the sample size calculation (Stata 12.1, Stata Corp. LP, College Station, TX). This study was designed as a

superiority trial where IN was expected to outperform NC and PC. Initial expected prevalence of respiratory disease was 15 to 20%. Sample size of 270 calves per treatment was estimated initially to provide a power of 80% and detect a 10% difference in clinical disease with $\alpha = 0.05$. Sample size was re-estimated and adjusted to 220 calves per treatment after recognizing the higher than anticipated level of disease and presumed greater effect of the intervention. All continuous variables were assessed for normality using the Shapiro-Wilk test. Measures of central tendency are presented as mean (SD) for ADG, and median (interquartile range, **IQR**) for EAGE. Raw ADG were compared using the *t*-test. Enrollment ages were compared by Wilcoxon rank sum test. Categorical variables (birth weight, **DYST**, twin, sex, **FPT**, **RIBF**, **ILL**, and **CON**) were assessed with contingency tables and chi-squared analysis, or Fisher's exact test when individual cell counts were < 5 . Relative risk was calculated to estimate the association between **RIBF** and **DYST** and the association between **ILL** and **CON**. A kappa was calculated using **ILL** and **CON** as variables in the **FREQ** procedure with the **TEST KAPPA** statement. The purpose of this kappa test was to determine the agreement between **CON** and **ILL** in their ability to identify calves with BRD during the preweaning period. When a calf died or was euthanized due to severe disease, a field-based postmortem examination was performed by a veterinarian. Overall mortality risk was calculated by dividing the number of calves that died during the follow-up period by the total number of enrolled calves. The overall risk of death from BRD was calculated by dividing the number of calves that died as the result of BRD during the follow-up period by the total number of enrolled calves. The BRD case fatality rate was calculated by dividing the number of calves that died due to BRD by the total number of calves positive for BRD. Calves positive for **CON** or **ILL** were considered positive for BRD.

Logistic regression models were fit for the outcomes **ILL** and **CON** using the **GLIMMIX** procedure. The explanatory variable of interest was treatment. All variables that had $P < 0.20$ after univariable analysis were offered into a multivariable model as fixed effects, including herd. Treatment was forced into each model. All 2-way interactions between treatment and fixed effects were tested. Manual backward stepwise elimination was used to refine the model to include variables that had $\alpha \leq 0.05$ level. Additionally, a change in estimate criterion of $\geq 25\%$ was used to assess for confounding before the final elimination of a variable. Predicted means for **CON** and ADG were assessed using the **LSMEANS** statement and these were adjusted using the **DIFF** option. Type 3 tests of fixed effects were used to determine significance which was set at

$\alpha < 0.05$. Collinearity (type II tolerance <0.10) was assessed with the GLM procedure.

Average daily gain was modeled using the MIXED procedure. The primary explanatory variable of interest was treatment. Variables were offered similar to the logistic regression models. Knowing that male calves typically grow faster than heifer calves (Koch et al., 1959), sex was forced into the model even though it was not significant in the univariable analysis. Predictions based on categorical variables were assessed using the LSMEANS statement. Pearson correlation coefficients were calculated to assess the correlations between model predictions and actual observations. Each model was assessed graphically for outliers and the normality of residuals was tested using the Shapiro-Wilk, Anderson-Darling, Kolmogorov-Smirnov, and Cramer-von Mises tests. Significance was set at $\alpha < 0.05$.

RESULTS

A total of 468 calves from 3 herds were enrolled and randomized into 3 treatment groups (PC, $n = 211$; IN, $n = 215$; NC, $n = 42$). All calves had a negative bovine viral diarrhea virus persistent infection status at enroll-

ment. Distributions of variables potentially affected by incomplete randomization are summarized in Table 1. A difference was observed in treatment group size between herds ($P < 0.0001$) due to the lack of NC at herd 2. However, no difference ($P = 0.48$) was present in the proportions of calves enrolled in PC and IN. Calves were younger at enrollment in IN (median age, IQR: 4, 3–5 d) compared with PC (median age, IQR: 4, 4–5 d; $P = 0.02$) and NC (median age, IQR: 5, 4–6 d; $P = 0.02$). However, no difference ($P = 0.22$) was present in EAGE between PC and NC. Serum protein data were missing from some calves in PC ($n = 3$), IN ($n = 5$), and NC ($n = 2$). The risk of FPT was greater for NC (15%) compared with PC (4%) and IN (7%; $P = 0.01$ and $P = 0.08$, respectively). The risk of FPT was not different between PC and IN ($P = 0.29$). Rib fractures were observed in 6% ($n = 28$) of calves. Rib fractures were associated with dystocia (RR 3.78, 95% CI: 1.83, 7.78). However, no association was observed between rib fractures and treatment ($P = 1.0$).

Overall, 54% ($n = 251$) calves were ILL positive. Before controlling for potentially confounding explanatory variables, NC had a lower proportion of ILL positive calves than PC (NC = 33% vs. PC = 59%; $P = 0.003$),

Table 1. Distribution of variables in 468 calves enrolled in a randomized blinded field study by calf characteristic and treatment group

Variable	Category	Treatment group, no. (% across category)			P-value
		PC ¹	IN ²	NC ³	
Herd	Herd 1	46 (40.4)	37 (32.5)	31 (27.2)	<0.0001
	Herd 2	156 (48.0)	169 (52.0)	—	
	Herd 3	9 (31.0)	9 (31.0)	11 (37.9)	
Birth weight (kg)	<40 kg	50 (46.7)	49 (45.8)	8 (7.5)	0.42
	40–46 kg	117 (48.0)	106 (43.4)	21 (8.6)	
	>46 kg	44 (37.6)	60 (51.3)	13 (11.1)	
Calving ease	Hard pull, no	183 (45.6)	182 (45.4)	36 (9.0)	0.74
	Hard pull, yes	25 (42.4)	30 (50.9)	4 (6.8)	
Twin	No	184 (44.1)	193 (46.3)	40 (9.6)	0.58
	Yes	21 (50.0)	19 (45.2)	2 (4.8)	
FPT ⁴	No	200 (46.5)	196 (45.6)	34 (7.9)	0.04
	Yes	9 (31.0)	14 (48.3)	6 (20.7)	
Sex	Male	112 (44.3)	124 (42.3)	17 (6.7)	0.11
	Female	99 (46.1)	91 (42.3)	25 (11.6)	
First RS ⁵	<5	208 (45.3)	211 (46.0)	40 (8.7)	0.33
	≥5	3 (33.3)	4 (44.4)	2 (22.2)	
First US ⁶	<3 cm	207 (45.0)	211 (45.9)	42 (9.1)	1.00
	≥3 cm	4 (50.0)	4 (50.0)	0	

¹Positive control (PC): 2 mL of commercially available multivalent injectable vaccine against bovine respiratory syncytial virus (BRSV), infectious bovine rhinotracheitis (IBR), parainfluenza 3 (PI₃), and bovine viral diarrhea administered subcutaneously at 6 wk of age.

²Intranasal vaccine (IN): 2 mL of commercially available trivalent injectable vaccine against BRSV, IBR, and PI₃ administered intranasally at 3 to 6 d of age.

³Negative control (NC): 2 mL of sterile saline administered both intranasally and subcutaneously at 3 to 6 d of age and 6 wk of age.

⁴Failure of passive transfer; cut-off serum total protein <5.2 mg/dL.

⁵Respiratory score, Wisconsin Calf Scoring Chart.

⁶Ultrasound examination.

Table 2. Multivariable logistic regression model for the prediction of clinical illness (ILL) in 468 Holstein dairy calves from 3 herds in southwestern Ontario randomly assigned to receive 1 of 3 preweaning vaccination protocols¹

Variable	Estimate	SE	P-value	Odds ratio (95% CI)
Intercept	-1.41	0.51	—	—
Treatment				
PC ²	0.15	0.41	0.71	1.16 (0.52–2.61)
IN ³	-0.16	0.42	0.71	0.85 (0.38–1.93)
NC ⁴	Referent	—	—	—
Sex				
Female	0.41	0.20	0.04	1.51 (1.02–2.23)
Male	Referent	—	—	—
Scours				
Yes	0.65	0.22	<0.01	1.91 (1.25–2.93)
No	Referent	—	—	—
Herd				
1	0.23	0.47	0.62	1.26 (0.50–3.16)
2	1.58	0.46	<0.001	4.85 (1.95–12.06)
3	Referent	—	—	—

¹ILL calves had a respiratory score >4 on at least one occasion during the study period.

²Positive control (PC): 2 mL of commercially available multivalent injectable vaccine against bovine respiratory syncytial virus (BRSV), infectious bovine rhinotracheitis (IBR), parainfluenza 3 (PI₃), and bovine viral diarrhoea administered subcutaneously at 6 wk of age.

³Intranasal vaccine (IN): 2 mL of commercially available trivalent injectable vaccine against BRSV, IBR, and PI₃ administered intranasally at 3 to 6 d and 6 wk of age.

⁴Negative control (NC): 2 mL of sterile saline administered both intranasally and subcutaneously at 3 to 6 d of age and 6 wk of age.

and IN (IN = 53%; $P = 0.02$). No difference was observed in the proportion of ILL calves between PC and IN ($P = 0.20$). After controlling for the effects of herd, sex, and scour, treatment was not associated with the odds of ILL (Table 2; $P = 0.32$).

Overall, 54% ($n = 253$) of the calves were CON positive. Before controlling for potentially confounding explanatory variables, NC had a lower proportion of CON positive calves than PC (NC = 33% vs. PC = 60%; $P = 0.002$), and IN (IN = 53%; $P = 0.02$). There was no difference in proportion of CON positive calves between PC and IN ($P = 0.14$). However, after controlling for herd, DYST, and RIBF, treatment was significantly associated with the odds of CON (Table 3; $P = 0.005$). The predicted probabilities of CON for each treatment are shown in Figure 1. The odds of CON in PC were 1.63 (95% CI: 1.04–2.56) times the odds of CON in IN ($P = 0.03$) and 0.38 (95% CI: 0.16–0.93) times the odds of CON in NC ($P = 0.03$). The odds of CON in IN were 0.23 (95% CI: 0.09–0.59) times the odds of CON in NC (0.002). A 2×2 contingency table compares the overall distribution of CON and ILL (Table 4). The outcomes CON and ILL were associated (RR 2.20, 95% CI: 1.79, 2.70; $P < 0.0001$), but their ability to identify the same animals as affected with respiratory disease during the preweaning period was only fair ($\kappa = 0.38$).

The ADG for all calves was 0.56 kg/d (SD = 0.17). Before controlling for potentially confounding explanatory variables, the ADG for NC was 0.69 kg/d (SD =

0.14), which was greater ($P < 0.0001$) than both PC and IN; however, PC and IN were not different [PC: 0.55 kg/d (SD = 0.17) vs. IN: 0.55 kg/d (SD = 0.17; $P = 0.56$]. Multivariable linear regression revealed a significant interaction between treatment and herd on the outcome ADG ($P < 0.01$; Table 5, Figure 2). In herd 1, IN decreased ADG by 0.10 kg/d (SE = 0.03) compared with PC ($P < 0.01$). In contrast, at herd 2, IN increased ADG by 0.03 kg/d (SE = 0.02; $P = 0.04$). Treatment did not influence ADG at herd 3 ($P > 0.25$). Thirty-nine (8%) calves died or were euthanized during the study. The overall risk of death from respiratory disease was 1.9% ($n = 9$). The case fatality rate for calves that were positive for CON or ILL was 3% (9/301). No difference was observed in overall mortality rates between vaccine protocols ($P = 0.58$).

DISCUSSION

The purpose of this study was to compare the effect of an intranasal vaccine, given to 3- to 6-d-old calves, with a subcutaneous vaccine, given later in life, and a placebo vaccine. Two different classification methods were used to identify diseased calves, and they included RS and TUS. Growth was assessed during the first 56 d of life. Two doses of a commercially available trivalent IN respiratory vaccine (experimental treatment, IN, first dose: 3 to 6 d of age; second dose: 6 wk of age) had no effect on the odds of having at least one episode of

Table 3. Multivariable logistic regression model for the prediction of CON in 451 Holstein dairy calves from 3 herds in southwestern Ontario randomly assigned to receive 1 of 3 preweaning vaccination protocols¹

Variable	Estimate	SE	P-value	Odds ratio (95% CI)
Intercept	−0.79	0.54	—	—
Treatment				
PC ²	−0.96	0.46	0.03	0.38 (0.16–0.93)
IN ³	−1.45	0.47	<0.01	0.23 (0.09–0.59)
NC ⁴	Referent	—	—	—
Dystocia				
Yes	0.73	0.35	0.04	2.08 (1.05–4.15)
No	Referent	—	—	—
Rib fracture				
Yes	1.31	0.56	0.02	3.71 (1.23–11.17)
No	Referent	—	—	—
Herd				
1	0.16	0.57	0.77	1.17 (0.39–3.56)
2	2.64	0.58	<0.0001	14 (4.4–43.82)
3	Referent	—	—	—

¹CON calves had ≥ 3 cm lung consolidation on at least one occasion throughout the study period.

²Positive control (PC): 2 mL of commercially available multivalent injectable vaccine against bovine respiratory syncytial virus (BRSV), infectious bovine rhinotracheitis (IBR), parainfluenza 3 (PI₃), and bovine viral diarrhea administered subcutaneously at 6 wk of age.

³Intranasal vaccine (IN): 2 mL of commercially available trivalent injectable vaccine against BRSV, IBR, and PI₃ administered intranasally at 3 to 6 d and 6 wk of age.

⁴Negative control (NC): 2 mL of sterile saline administered both intranasally and subcutaneously at 3 to 6 d of age and 6 wk of age.

ILL during the study period, compared with a commercially available subcutaneous product (PC, only dose: 6 wk of age) or sterile saline (NC) after accounting for potentially confounding variables. In contrast, IN significantly reduced the odds of having at least one episode of CON compared with both positive and negative controls. The effect of IN on calf growth was herd dependent. This interaction resulted in a greater ADG in herd 2 in IN treated calves compared with PC. In

Figure 1. Predicted probability of CON by vaccine protocol after controlling for herd, dystocia, and rib fractures. Error bars represent SEM. CON = occurrence of ≥ 3 cm lung consolidation at least once in the study period. PC = white; IN = dark gray; NC = light gray. PC = positive control: 2 mL of commercially available multivalent injectable vaccine against bovine respiratory syncytial virus (BRSV), infectious bovine rhinotracheitis (IBR), parainfluenza 3 (PI₃), and bovine viral diarrhea administered subcutaneously at 6 wk of age. IN = intranasal treatment: 2 mL of commercially available trivalent injectable vaccine against BRSV, IBR, and PI₃ administered intranasally at 3 to 6 d and 6 wk of age. NC = negative control: 2 mL of sterile saline administered both intranasally and subcutaneously at 3 to 6 d and 6 wk of age.

herd 1, IN was associated with either a lower ADG or no change in ADG compared with the PC and NC, respectively. No differences in ADG were observed in herd 3.

Clinical observation is used commonly to detect respiratory disease (Amrine et al., 2013). However, compared with TUS, respiratory scoring did not detect the disease sparing effects of IN in the current study. As previously mentioned, lung lesions typically affect more calves at slaughter than indicated by observation-based treatment records (Wittum et al., 1996; Thompson et al., 2006), suggesting the presence of subclinical disease. A study done in a single herd demonstrated that clinical signs alone failed to identify lung lesions, which were seen in more than 60% of preweaned calves (Ollivett and Buczinski, 2016). Recent findings demonstrate that lung consolidation has long-term implications on dairy calf survival and reproduction (Adams and Buczinski, 2016; Teixeira et al., 2017). This highlights

Table 4. A 2×2 contingency table comparing the distribution of CON and ILL in Holstein dairy calves in Southwestern Ontario, enrolled in a randomized vaccine field trial (n = 468)

ILL ¹	CON ²	
	1	0
1	181	70
0	72	145

¹At least one respiratory score >4 .

²At least one ultrasound exam with ≥ 3 cm lung consolidation.

Table 5. Multivariable linear regression model for ADG on 421 Holstein dairy calves from 3 herds in southwestern Ontario, randomly assigned to receive 1 of 3 preweaning vaccination protocols¹

Variable	Estimate	SE	95% CI	P-value
Intercept	0.513	0.0761	0.364, 0.663	<0.0001
Treatment				
PC ²	0.00439	0.0638	-0.121, 0.130	0.95
IN ³	-0.0426	0.0655	-0.171, 0.0862	0.52
NC ⁴	Referent	—	—	—
Twin				
Yes	-0.1380	0.0253	-0.188, -0.0882	<0.0001
No	Referent	—	—	—
Dystocia				
Yes	-0.0467	0.0212	-0.0884, -0.00497	0.028
No	Referent	—	—	—
House				
Outdoors	-0.0285	0.0378	-0.103, 0.0459	0.45
Indoors	Referent	—	—	—
Rib fracture				
Yes	-0.0565	0.0285	-0.113, -0.00062	0.048
No	Referent	—	—	—
Sex				
Male	0.0447	0.0143	0.0166, 0.0728	<0.01
Female	Referent	—	—	—
Birth weight				
<40 kg	-0.0645	0.0212	-0.106, -0.0228	<0.01
40–46 kg	-0.0217	0.0173	-0.0557, 0.0123	0.21
>46 kg	Referent	—	—	—
Enrollment age (d)	0.00332	0.0060	-0.0086, 0.0152	0.58
Protocol × herd interaction				
PC × 1	0.0494	0.0733	-0.0947, 0.193	0.50
PC × 2	-0.0804	0.0687	-0.216, 0.0547	0.4
IN × 1	0.00023	0.0754	-0.149, 0.148	0.99
Herd				
1	-0.0549	0.0606	-0.174, 0.0642	0.37
2	-0.173	0.0625	-0.295, -0.0497	0.006
3	Referent	—	—	—

¹Calves dying before 56 d of age were not included in the analysis.

²Positive control (PC): 2 mL of commercially available multivalent injectable vaccine against bovine respiratory syncytial virus (BRSV), infectious bovine rhinotracheitis (IBR), parainfluenza 3 (PI₃), and bovine viral diarrhea administered subcutaneously at 6 wk of age.

³Intranasal vaccine (IN): 2 mL of commercially available trivalent injectable vaccine against BRSV, IBR, and PI₃ administered intranasally at 3 to 6 d and 6 wk of age.

⁴Negative control (NC): 2 mL of sterile saline administered both intranasally and subcutaneously at 3 to 6 d of age and 6 wk of age.

the importance of minimizing the incidence of CON in preweaned dairy calves. The use of IN may prove beneficial in minimizing the risk of CON, given the findings of the present study.

The fact that agreement between diagnostic methods was just fair indicates that different populations of calves were identified by each predictor. It is possible that a clinical difference did not exist, the cut-points used to create ILL and CON were inappropriate, or the subjective nature and variability inherent to clinical scoring precluded finding small differences in a limited study population. The disagreement between the 2 variables rested evenly between CON positive calves that were ILL negative and ILL positive calves that were CON negative. This could be the result of clinical scoring systems that are not highly specific to the disease they are supposed to be detecting. As can be

seen in the logistic regression model, the significant association between scour and ILL highlights the fact diseases other than BRD can evoke a positive test when respiratory scoring. The effect of these variables on objective measures, such as calf growth, will help researchers understand which variable most accurately identifies the population of calves affected with BRD.

Rib fractures are common in neonatal foals (Jean et al., 1999); however, their presence and effect on ADG was an unexpected finding in this study in dairy calves. Rib fractures were typically located near the costochondral junction of the cranial thorax, similar to previous reports (Jean et al., 1999), and were easily imaged via TUS. Six percent of calves in the current study had obvious rib fractures, which is much lower than previous reports of 23% in calves that die during the perinatal period (Schuijt, 1990) and 40% of calves

Figure 2. Differences in the LSM of ADG (kg/d) showing the interaction between treatment and herd. The linear mixed model controlled for vaccine type, herd, twin, housing, dystocia, rib fractures, birth weight, and age at enrollment. Different letters (a,b) within a herd category are statistically different ($P < 0.05$). PC = white; IN = dark gray; NC = light gray. PC = positive control: 2 mL of commercially available multivalent injectable vaccine against bovine respiratory syncytial virus (BRSV), infectious bovine rhinotracheitis (IBR), parainfluenza 3 (PI₃), and bovine viral diarrhoea administered subcutaneously at 6 wk of age. IN = intranasal treatment: 2 mL of commercially available trivalent injectable vaccine against BRSV, IBR, and PI₃ administered intranasally at 3 to 6 d and 6 wk of age. NC = negative control: 2 mL of sterile saline administered both intranasally and subcutaneously at 3 to 6 d and 6 wk of age.

born with veterinary assistance (Mee, 2008). This likely reflects different study populations, although an equine study reported a 20% prevalence of rib fractures in foals in one breeding herd, twice as many resulting from dystocia than from normal parturitions (Jean et al., 1999). Despite the fact that none of the observed fractures were severely displaced or causing internal thoracic abnormalities, such as hemothorax or pneumothorax, rib fractures did contribute significantly to the variation in ADG. Interestingly, recent work has shown that anti-inflammatory doses of meloxicam given immediately after a difficult birth results in greater calf growth during the early weeks of life (Murray, 2014). One reason for this finding could be treatment of the pain and inflammation associated with rib fractures, which allows for improved utilization of liquid feed and greater growth. In the current study, an association was present between dystocia and rib fractures; however, the correlation was very low and highlights the fact that a particularly difficult calving is not a prerequisite for rib fractures.

Although herd differences were noted in this study, the fact that only 3 herds were included is a limitation to the ability to draw inferences about herd specific factors regarding the efficacy of IN. Inclusion of a large number of herds varying in locale and management practices would be necessary to determine exactly which situations IN would be most effective. An additional limitation is the potential confounding effect of barn on

the relationship between the measured outcomes and treatment at herd 2. This design was intentional as IN and PC calves needed to be housed separately to prevent exposure of PC to IN vaccine virus. The authors do acknowledge that switching the treatment status of each barn would have been ideal. However, the feasibility and additional time requirements necessary for incorporating washout periods when transitioning between groups were considered impractical. The 2 barns were close in proximity to each other, being separated by a milk preparation room; and barn age, design, flow of animals, cleaning, and labor were identical. These factors should have reduced the risk of a barn effect on calf health and performance. The authors also acknowledge that our study design was not perfectly balanced due to incomplete randomization of calves in herd 2, resulting in a quasi-randomized clinical trial.

This study differed from previous work on the efficacy of intranasal vaccination in several ways. The calves in the current study were conventionally raised and likely exposed to chronic, low level natural challenges from the whole spectrum of respiratory pathogens (Gorden and Plummer, 2010). In contrast, previous studies incorporated controlled experimental BRSV challenges (Woolums et al., 2004; Ellis et al., 2007, 2010, 2013; Vangeel et al., 2007) and PI₃ (Vangeel et al., 2007) intended to replicate natural disease in small group of animals. Despite the high prevalence of disease in the study population, the risk of death from respiratory disease was low. This suggests that the natural challenge in this study might not be as acutely aggressive as those demonstrated by single pathogen challenge models in past studies, as several calves from each BRSV challenge died or required euthanasia due to the severity of disease (Woolums et al., 2004; Ellis et al., 2007, 2013). Acknowledging that previous study designs were in part due to federal regulations regarding licensing procedures for new vaccines (Ellis et al., 2013), data from this field study may be more relevant to dairy producers and bovine practitioners.

CONCLUSIONS

A commercially available trivalent IN vaccine has the potential to reduce the lung lesions associated with BRD and improve growth in young dairy cattle. Although these findings were meaningful, herd factors played a role in determining whether or not significant changes in ADG were seen. Also, IN vaccination did not eliminate the risk of disease in the current study; therefore, this practice should not be viewed as a panacea. Best management practices regarding colostrum management, calf nutrition, housing strategies, ventila-

tion, and appropriate vaccination protocols should be integrated to provide the optimal environment for the growing dairy calf.

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