



J. Dairy Sci. 98:1–4
<http://dx.doi.org/10.3168/jds.2014-8972>
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Short communication: Characterization of the serologic response induced by vaccination of late-gestation cows with a *Salmonella* Dublin vaccine

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ABSTRACT

Diarrhea due to *Salmonella* infection is an important cause of neonatal calf diarrhea. The acquisition of passive immunity in the calf by vaccinating the dam has shown some success in previous studies; however, no data exists on the use of currently licensed vaccines in the United States. Therefore, the purpose of this study was to determine whether vaccinating cows in late gestation with a commercially available *Salmonella* Dublin vaccine would stimulate *Salmonella*-specific antibodies in the colostrum of cows at calving and whether these antibodies would be transferred to the calf. Thirty Holstein cows were vaccinated 3 wk before the end of lactation with a *Salmonella enterica* serovar Dublin vaccine, with a second dose given at dry-off. An additional 30 cows received only saline. Calves had a blood sample collected immediately after birth and were then fed fresh colostrum from their dam within 2 h of calving. A postcolostrum blood sample was collected 24 to 48 h later. *Salmonella* Dublin antibodies in colostrum as well as serum from the cows and calves were measured using an ELISA technique. Results of this study showed that vaccinated cattle had elevated *Salmonella* Dublin antibody titers at the time of calving (40.3 ± 9.1) as compared with control cows (-9.4 ± 1.1). Calves that received colostrum from vaccinated cattle also had a significant increase in *Salmonella* Dublin antibodies (88.5 ± 8.9) as compared with calves born to unvaccinated cows (-3.2 ± 1.2). This study demonstrated that the use of a commercially available *Salmonella* Dublin vaccine can stimulate antibodies that are passed on to the calf via colostral transfer. Further studies need to be done to determine whether these antibodies will offer protection against *Salmonella* challenge.

Key words: calf, colostrum, immunoglobulin G, *Salmonella* Dublin, passive immunity

Short Communication

Salmonella infections are an economically important disease of cattle and also represent a worldwide public health concern. Although cattle of all ages can be infected with *Salmonella* bacteria, death is most often reported in calves less than 8 wk of age (Smith et al., 1980; Mohler et al., 2009). The science behind the vaccination of cattle for different serotypes of *Salmonella* has progressed in the past 20 yr (Mohler et al., 2009), and 2 commercially available vaccines are currently on the market in the United States for use in cattle. However, neither of these vaccines are approved for calves less than 2 wk of age, and diarrhea due to *Salmonella* in neonatal calves still remains a problem. This has led some veterinarians to use vaccines in young calves by unapproved routes of administration (Habing et al., 2011). Another potential approach to achieving at least partial immunity in young calves is through vaccination of the dam. This has long been a common practice for other causes of neonatal calf diarrhea including *Escherichia coli*, rotavirus, and coronavirus with established efficacy (Snodgrass et al., 1982; Crouch et al., 2000). Previous studies suggested that the use of passive immunity through vaccination of the dam or via the use of *Salmonella*-specific antibodies derived from egg yolks could offer at least partial protection against clinical disease in calves (Royal et al., 1968; Weaver, 1976; Smith et al., 1980; Jones et al., 1988; Yokoyama et al., 1998). Therefore, the purpose of this study was to determine whether vaccinating cows at dry-off with a commercially available *Salmonella* Dublin vaccine would result in the presence of *Salmonella*-specific IgG antibodies in the colostrum of cows at calving and whether these colostral antibodies would be transferred to the calf.

This study was approved by the Institutional Animal Care and Use Committee at North Carolina State University. Sixty lactating Holstein cattle ranging from 3 to 9 yr of age were identified for this study according to computer records (Table 1). All cattle were housed on a single dairy farm in North Carolina. This farm had

Received October 13, 2014.

Accepted December 16, 2014.

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never used an autogenous or commercially available *Salmonella* vaccine, and there had not been a diagnosis of clinical salmonellosis on the farm for at least 10 yr. Although extensive testing had not been done to rule out the presence of subclinical salmonellosis in the herd, the farm had extensive diagnostic testing records of both calves and adult cattle. This data included fecal cultures from healthy animals as well as those with diarrhea, along with numerous necropsy reports spanning a period of 10 yr. During this period there had never been a positive *Salmonella* culture on the farm. A search of the farm records identified the next 60 cows scheduled for dry-off. These cows were randomly allocated to serve in either vaccinated or control groups by assigning every other cow in the computer-generated list to opposite groups. Thirty cows were allocated to a group that was vaccinated 3 wk before the scheduled dry-off date. A total of 10 mL of blood was collected, and each cow was vaccinated with 2 mL of a commercially available modified live *Salmonella enterica* serotype Dublin bacterin (Entervene-d, Boehringer Ingelheim Vetmedica, St. Joseph, MO) given subcutaneously in the neck. The vaccine is approved in the United States for cattle greater than 2 wk of age. An additional 30 cows were allocated to the control group and received 2 mL of saline subcutaneously in the neck. Cows were identified with neck bands by group at the time of initial vaccination. Rectal temperature was monitored before vaccination as well as 2, 4, 12, and 24 h after vaccination. In addition, daily milk production from both groups was recorded for 2 wk after the initial vaccination, and any abortions or stillbirths throughout the study period were noted. Three weeks after the initial injection (on the actual dry-off day), all cows in the vaccinated group received a second injection dose of (2 mL) vaccine, and the control group again received saline. Cows were housed together in a grass calving lot during the 8-wk dry period.

At the time of calving, colostrum was harvested within 2 h of parturition, and a 10-mL blood sample was collected from all cows. A 50-mL sample of colostrum was saved from each cow, and then all calves were fed 4 L of colostrum from their dam within 2 h

of birth. A presuckle blood sample was collected from each calf before colostrum feeding, with a follow-up blood sample collected from each calf 24 to 48 h after colostrum administration.

Serum and colostrum antibody titers for *Salmonella* Dublin were determined using an ELISA. The assay was performed by a commercial diagnostic laboratory (Animal Health Diagnostic Center, Cornell College of Veterinary Medicine, Ithaca, NY), and personnel performing the ELISA assays were not aware of group assignment for any of the animals. *Salmonella* Dublin titers were determined using a commercially available ELISA kit (PrioCHECK *Salmonella* Ab Bovine Dublin, Prionics Life Technologies, Schlieren, Switzerland) that has been validated for plasma, serum, and milk. This ELISA assay was developed in cooperation with the Danish Veterinary Institute and is based on the same principle as the Danish *Salmonella* Dublin ELISA that has been used extensively in previous studies (Nielsen and Ersbøll, 2004a,b, 2005). Results of the ELISA assays were reported as percent positivity.

Data are presented as mean \pm standard error. Rectal-temperature and milk-production data along with serologic titers for cows and calves born to cows from each group were compared using one-way ANOVA. Values of $P < 0.05$ were considered significant. A statistical software package was used to conduct the data analysis (SAS, version 9.1; SAS Institute Inc., Cary, NC).

Rectal temperature did not significantly differ in any time point after vaccination as compared with baseline (prevaccination). Although milk production in both groups declined toward the end of the lactation period, vaccinated and control groups did not differ at any point during the 2-wk period where milk production was monitored. For the first 5 d following vaccination, daily milk production did not change from its baseline value (Table 1) in either group. A total of 59 of the 60 cows calved successfully and had colostrum samples collected (one cow from the control group aborted during the dry period and was removed from the study). *Salmonella* titers did not differ between groups of cows at the time of initial vaccination (Table 2). Cows in the vaccinated group had a significant ($P = 0.007$) increase

Table 1. Summary of age and production history for cows included in this study¹

Item	Control group	Vaccinated group
Age of cow (yr)	5.8 \pm 2.4	5.4 \pm 2.3
Lactation number	2.7 \pm 1.4	2.5 \pm 1.3
DIM (current lactation)	341 \pm 24	352 \pm 29
305-d mature equivalent yield (kg)	10,400 \pm 1,410	10,310 \pm 1,620
Milk production—first day of study (kg)	19.3 \pm 3.3	17.9 \pm 3.1

¹The data were collected from farm records on the first day of the experimental period (3 wk before dry-off).

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