ABSTRACT

Forty-four Holstein calves (19 male and 25 female) were used in this study of the relationships among age at first colostrum feeding, IgG intake, and apparent efficiency of IgG absorption. Time of birth was recorded for each calf and the calves were fed colostrum ad libitum after birth at either 0930 or 1630 h. Blood samples were collected immediately before and 24 h after colostrum feeding. Data from calves were then categorized into 4 groups representing time from birth to colostrum feeding: A = fed within 1 h (n = 5); B = fed from 1 to 6 h (n = 10); C = fed from 6 to 12 h (n = 21); and D = fed from 12 to 18 h (n = 8) after birth. Average total intake of colostrum was 3.6 ± 0.1 L. Over 80% of the calves consumed ≥3 L of colostrum. Apparent efficiency of IgG absorption declined remarkably 12 h after birth. Mean apparent efficiency of absorption of IgG in group D (15.8 ± 3.0%) was lower than that in groups A (30.5 ± 3.9%) and B (27.4 ± 2.8%). Serum IgG concentration in calves was positively correlated with IgG intake in all groups. The relationship between mass of IgG consumed and calf serum IgG at 24 h was different for each time of colostrum feeding, with only limited differences observed between groups A and B. We concluded that failure of transfer of passive immunity in newborn calves may be avoided if calves consume ≥3 L of colostrum within 6 h after birth. These findings help define the opportunity to minimize failure of transfer of passive immunity to newborn calves under management programs similar to those used on commercial dairy farms.

Key words: immunoglobulin G intake, age at first colostrum feeding, apparent efficiency of absorption

INTRODUCTION

Calves are defined as experiencing failure of passive transfer (FPT) when serum IgG concentration is <10 mg/mL at 24 and 48 h after birth (Weaver et al., 2000). Calves with low serum IgG concentrations are at greater risk for preweaning morbidity and mortality than calves with higher concentrations of IgG (National Animal Health Monitoring System, 1996). Mortality rate of calves with serum IgG concentration <10 mg/mL was more than twice as high as that of calves with higher IgG concentrations (Wells et al., 1996).

The primary factors that influence transfer of IgG from the colostrum are the age of the calf at the first colostrum feeding and the mass of IgG intake (Stott et al., 1979a,b). Staley and Bush (1985) reported that intestinal epithelial cells in calves lost their ability to absorb intact macromolecules after about 24 h. Apparent efficiency of IgG absorption (AEA) from the small intestine of calves declines soon after birth (Abel Francisco and Quigley, 1993; Rajala and Castrén, 1995). Hopkins and Quigley (1997) showed that serum IgG concentration was positively correlated with IgG intake. Therefore, it is recommended that neonatal calves be supplied adequate high-quality colostrum mass as soon as possible after birth (NRC, 2001).

Several researchers suggest that calves should be provided approximately 4 L of colostrum (Roy, 1980; Besser et al., 1991; Hopkins and Quigley, 1997). However, additional labor is often needed when feeding more colostrum with an esophageal feeder than calves may consume ad libitum or when feeding colostrum more frequently. Limited data (Hopkins and Quigley, 1997) are available that show the mass of colostrum that calves are able to consume ad libitum at the first feeding.

Todd and Whyte (1995) reported that serum IgG concentration of calves was not significantly different when the age at first colostrum feeding occurred between 1 and 8 h after birth. They considered that the high IgG concentration of the colostrum they fed to calves may have compensated for any loss of absorption efficiency associated with delays in the first feeding. In that controlled study, IgG intake was held constant. However, the amount of colostrum and the mass of IgG that calves consume is often not easily controlled in on-farm environments without a significant labor cost. The
goal of this study was to better define the relationships between calf age at colostrum feeding, voluntary intake of colostrum, and apparent efficiency of IgG absorption in calves under management schemes typical of dairy farms. Results from this study may be used in development of newborn calf management protocols.

MATERIALS AND METHODS

Animal management and blood sampling were carried out according to the guidelines for animal experiments of Hokkaido Animal Research Center at Shin-toku (Hokkaido, Japan). The present study included 44 Holstein dairy calves (19 male and 25 female) from the Hokkaido Animal Research Center. The birth of each calf was carefully observed and calves were immediately separated from their dams to avoid suckling. Time of birth was recorded and calves were weighed and placed in a dry, disinfected pen with straw bedding until colostrum feeding. Cows were milked at either 0600 or 1600 h for the first time after calving, and first-milking colostrum was used to feed each cow's own calf. The time of feeding colostrum to calves was fixed at either 0930 or 1630 h. Calves were initially offered 4 L of their dam's colostrum using a nipple pail. If it was apparent that the calf would consume all 4 L of colostrum, an additional 2 L of colostrum was added to the nipple pail. In some cases, dams did not yield sufficient colostrum to meet the intake of the calf or colostrum from the dam was not otherwise available. In those cases, frozen colostrum from other cows (colostrometer reading of >1.050) was thawed and used to supplement the dam's colostrum. Calves were fed their dam's colostrum first before the thawed colostrum source was fed. Frozen colostrum was stored in plastic bags at −30°C. Frozen colostrum was thawed by sealing the bag in a second bag and immersing the bag in water at 40 to 45°C until thawed. The volume of colostrum consumed was determined by the difference between the quantity offered to the calf and the residual colostrum left in the pail after feeding. Blood samples were collected just before and 24 h after colostrum feeding. Data from calves were then categorized into 4 groups representing time from birth to colostrum feeding. The groups were as follows: A = colostrum fed within 1 h (n = 5); B = colostrum fed between 1 and 6 h (n = 10); C = colostrum fed between 6 and 12 h (n = 21); and D = colostrum fed between 12 and 18 h (n = 8) after birth. One calf consumed only 1.5 L at 20 min after birth and was excluded from the data set. The number of calves that received only frozen colostrum was 1, 2, 3, and 0 for groups A, B, C, and D, respectively. Calves were not fed milk for 24 h after the feeding of colostrum. Calves were then fed 2 L of fresh milk from a nipple pail at 0930 and 1630 h.

Samples of colostrum fed to each calf were collected for IgG analysis. Colostrum was centrifuged at 18,000 \( \times g \) for 30 min at 4°C and the supernatant collected and stored at −30°C. Blood samples were collected from calves by jugular venipuncture with a 10-mL syringe and 18-gauge needle, and blood was transferred to a plastic evacuated tube (Venoject II, Terumo Inc., Tokyo, Japan). Blood was allowed to clot at room temperature for 30 min and serum was collected after centrifugation at 2,000 \( \times g \) for 10 min at 20°C. Serum was stored at −30°C. Concentration of IgG in defatted colostrum and serum samples was determined by single radial immunodiffusion (The Institute for Metabolic Ecosystem Inc., Miyagi, Japan), according to the manufacturer’s instructions.

Intake of IgG, serum IgG and AEA of IgG were calculated as per Quigley and Drewry (1998), where

\[
\text{IgG intake (g)} = \text{IgG concentration in milk (mg/mL)} \times \text{milk volume (L)},
\]

\[
\text{serum IgG (g)} = \text{IgG concentration in serum (mg/mL)} \times \text{serum volume (L)},
\]

\[
\text{AEA (%)} = \left( \frac{\text{serum IgG (g)}}{\text{IgG intake (g)}} \right) \times 100.
\]

Serum volume (L) was estimated using the value of 7% of BW at birth, which has been used by others in determining AEA (Quigley and Drewry, 1998).

Statistical analyses were performed using SAS software (version 9.3, SAS Institute Inc., Cary, NC). Descriptive data at the first colostrum feeding were analyzed by 1-way ANOVA using the GLM procedure. Treatment means were separated using the least significant difference test when a significant treatment effect was observed. Statistical differences were considered significant at \( P < 0.05 \). The relationship between AEA and age at first colostrum feeding was analyzed by using the NLIN procedure to perform breakpoint analysis. A regression analysis with the REG procedure was used to assess the association between IgG intake and serum IgG concentrations at 24 h after colostrum intake in each group.

RESULTS

Birth weight of calves was not different among groups (Table 1). The age at colostrum feeding across all calves ranged from 20 min (calf consumed only 1.5 L of colostrum) to 17.8 h; mean age at colostrum feeding was different for each group (Table 1). Mean colostrum intake volume was 3.6 ± 0.1 L and ranged from 1.5
to 5.2 L for all calves. Birth weight, colostrum intake, and the ratio of colostrum intake to birth weight were not significantly different among groups. Eighty-four percent of all calves (36 of 43 total calves) consumed >3 L of colostrum. Of the calves consuming <3 L of colostrum, the average colostral intake volume was 2.7 L and ranged from 2.4 to 2.9 L.

The relationship between AEA and age at first colostrum feeding is shown in Figure 1. We observed wide variation in AEA; however, AEA tended to decline slowly to an inflection point at 12.26 h, after which time it declined more rapidly. Mean AEA of group D was significantly lower \((P < 0.05)\) than that of group A or B (Table 1).

The relationship between mass of IgG intake and the resulting calf serum IgG concentration at 24 h after colostrum intake is illustrated in Figure 2. Concentration of IgG in calf serum at 0 h was below the detection limits of the assay and therefore regarded as 0 mg/mL. As IgG intake increased, serum IgG concentrations also

Figure 1. Relationship between age at colostrum feeding and apparent efficiency of absorption (AEA) of IgG. Breakpoint analysis indicates that the decline in AEA accelerates 12 h after colostrum feeding.

<table>
<thead>
<tr>
<th>Item</th>
<th>A (n = 4)</th>
<th>B (n = 10)</th>
<th>C (n = 21)</th>
<th>D (n = 8)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth weight (kg)</td>
<td>43.3 ± 0.5</td>
<td>43.3 ± 2.5</td>
<td>42.4 ± 1.2</td>
<td>42.9 ± 2.0</td>
</tr>
<tr>
<td>Age at colostrum feeding (h)</td>
<td>0.9 ± 0.1</td>
<td>3.3 ± 0.4</td>
<td>8.9 ± 0.4</td>
<td>15.5 ± 0.6</td>
</tr>
<tr>
<td>Colostrum intake (L)</td>
<td>3.0 ± 0.2</td>
<td>3.4 ± 0.2</td>
<td>3.7 ± 0.2</td>
<td>3.8 ± 0.3</td>
</tr>
<tr>
<td>Intake (% of birth weight)</td>
<td>7.0 ± 0.5</td>
<td>8.0 ± 0.5</td>
<td>8.8 ± 0.3</td>
<td>8.9 ± 0.7</td>
</tr>
<tr>
<td>Serum IgG at 24 h (mg/mL)</td>
<td>26.8 ± 6.0</td>
<td>22.6 ± 1.2</td>
<td>17.2 ± 1.5</td>
<td>12.5 ± 2.2</td>
</tr>
<tr>
<td>Apparent efficiency of absorption (%)</td>
<td>30.5 ± 3.9</td>
<td>27.4 ± 2.8</td>
<td>23.7 ± 1.7</td>
<td>15.8 ± 3.0</td>
</tr>
</tbody>
</table>

\(^{a, b}\)Means within rows with different superscripts differ \((P < 0.05)\).

\(^{1}\)Calf groups are as follows: A = fed colostrum 0 to 1 h after birth; B = fed colostrum 1 to 6 h after birth; C = fed colostrum 6 to 12 h after birth; and D = fed colostrum 12 to 18 h after birth.
increased in all groups. However, the slope of regression declined with increasing age at colostrum feeding, consistent with the declining AEA.

**DISCUSSION**

The purpose of this study was to more clearly define the relationships between age at first feeding of colostrum, volume of colostrum consumed, and mass of IgG consumed by the calf relative to the resulting calf serum IgG concentrations. These relationships were characterized under management protocols that reflected typical on-farm care of newborn calves. Consumption of high-quality colostrum (high concentrations of IgG) is important to ensure that adequate serum IgG concentrations are attained in the calf (Jaster, 2005). Others have recommended that calves be fed >100 g of IgG in the first colostrum feeding (Davis and Drackley, 1998). Results from the present study demonstrate that calves need to consume 120 g of IgG if fed in the first hour after birth or 125 g of IgG if fed between 1 and 6 h, to achieve 10 mg/mL of serum IgG at 24 h. These data suggest that a serum IgG concentration >10 mg/mL would be achieved if calves consumed 3 L of colostrum containing ≥40 mg/ml IgG in one feeding within 6 h after birth.

Increasing colostrum intake is one important factor to avoid FPT. Earlier studies suggested that newborn calves should receive approximately 2 L of colostrum in the first few hours after birth (Stott et al., 1979a; Roy, 1980). Others have indicated that colostrum volume is negatively correlated with colostrum IgG1 concentration (Pritchett et al., 1991). Besser et al. (1991) reported that calves often receive an inadequate mass of immunoglobulin when 2 L of first-milking colostrum is fed. To maximize serum immunoglobulin concentrations, therefore, recommended colostrum feeding methods indicate that neonatal calves should be fed greater quantities of colostrum (>2 L) as early as possible after birth. Roy (1980) suggested that calves should consume 4 L of colostrum divided into 2 feedings within 12 h after birth. Besser et al. (1991) and Morin et al. (1997) also reported that serum immunoglobulin concentration was greater when calves were fed 4 L of colostrum at birth versus 2 L at birth or 2 L at 12 h. In the latter study, colostrum was administered to calves by esophageal feeder to achieve the expected colostrum intake. Hopkins and Quigley (1997) reported that mean consumption of colostrum by calves was 3.0 L when calves were offered 3.8 L of colostrum in 1 feeding at a mean of 1.3 h after birth.

In the present study, the colostrum intake measurements indicated that 3 L of colostrum could be consumed without esophageal tube administration for most calves at the first feeding, regardless of time after birth. We observed that colostrum intake by one calf at 20 min after birth was only 1.5 L. That calf could not yet stand on its own and did not show a significant appetite for colostrum. It may be that a calf’s ability to consume colostrum immediately after birth is limited.

Intestinal epithelial cells lose their ability to absorb intact macromolecules as the age of the newborn increases (Staley and Bush, 1985). In the calf, AEA decreases from birth to essentially zero at approximately 24 h (Weaver et al., 2000). In the current study, AEA varied widely within each group (Figure 1). Although Stott and Fellah (1983) indicated that AEA was improved by feeding 1 L of colostrum compared with feeding an equivalent mass of immunoglobulin in 2 L of colostrum, others have observed that increasing the IgG mass by feeding supplemental colostrum replacer decreased the AEA of IgG (Besser et al., 1985; Campbell et al., 2007). Those studies differed in the mass of immunoglobulin fed, with <130 g being fed in the Stott and Fellah (1983) study, 100 to 700 g in Besser et al. (1985), and 130 to 390 g in Campbell et al. (2007). Besser et al. (1985) considered that calves had a physiological limitation for absorption of immunoglobulin. In the present study, wide variation in AEA was observed within each group, which probably reflects the wide range of IgG mass consumed (from 23 to 553 g). Although calves begin to lose the ability to absorb macromolecules such as immunoglobulin (Abel Francisco and Quigley, 1993; Rajala and Castrén, 1995), results from the present study suggest that the decline in AEA occurs at a slower rate early after birth followed by a
much faster rate after about 12 h. These data suggest that AEA in the initial hours after birth is influenced more by the mass of immunoglobulin consumed than by age at the first colostrum feeding, at least within the first 6 h after birth.

The relationship between IgG intake and calf serum IgG was linear and positive (Figure 2), in agreement with observations by Hopkins and Quigley (1997). In the present study, the effect of age of calf at colostrum feeding on the relationship between IgG intake and IgG serum concentration was also evaluated. We observed that the slope of regression lines declined with increasing age at colostrum feeding.

CONCLUSIONS

In this study, AEA and serum IgG concentration in calves at 24 h was significantly influenced by the mass of IgG consumed. Apparent efficiency of absorption of IgG declined by less than 0.3%/h from calving to 12 after birth, and then declined more rapidly at 2.5%/h to at least 18 h after birth. For supplying sufficient IgG to calves, the mass of IgG intake was more important than age at first colostrum feeding within the limited early period of macromolecular transport. Most calves could consume at least 3 L of colostrum in one feeding. Therefore, calves consuming at least 3 L of colostrum containing ≥40 mg/mL of IgG should avoid failure of transfer of passive immunity.

REFERENCES


