Effect of Freezing Treatment on Colostrum to Prevent the Transmission of Bovine Leukemia Virus

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ABSTRACT. Here, we used a sheep bioassay to determine the effect of freezing colostrum to prevent the transmission of bovine leukemia virus (BLV) among neonatal calves. Leukocytes were isolated from the colostrum of a BLV-infected Holstein cow and were then either left untreated (control) or freeze-thawed. A sheep inoculated intraperitoneally with the untreated leukocytes was infected with BLV at 3 weeks after inoculation, whereas the sheep inoculated with treated leukocytes did not become infected. The uninfected sheep was inoculated again with leukocytes isolated from the colostrum of another BLV-infected Holstein cow after freezing treatment, and again it did not become infected with BLV. Finally, this sheep was inoculated with the leukocytes isolated from the colostrum of another virus-infected cow without freezing treatment, and it became infected with BLV at 4 weeks after inoculation. The results indicate that colostrum should be frozen as a useful means of inactivating the infectivity of BLV-infected lymphocytes.

KEY WORDS: bovine leukemia virus, colostrum, freezing treatment, sheep bioassay.

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Enzootic bovine leukosis (EBL) is a contagious disease of cattle caused by the bovine leukemia virus (BLV), which belongs to the genus *Deltaretrovirus* in the family *Retroviridae* [17]. Most BLV infections are asymptomatic with approximately 30 to 40% of infected cattle developing persistent lymphocytosis (PL), while less than 5% develop malignant lymphosarcoma [16]. BLV infection has a worldwide distribution [16]; however, several countries in Europe have successfully eradicated BLV through changes in management procedures, such as test-and-slaughter control programs [1, 14].

The first large-scale epidemiological survey of BLV in cattle was carried out in 1982 in Japan, and the second survey was conducted during 2007, evaluating a total of 5,420 cattle from 209 farms in seven prefectures [12]. The survey showed that the prevalence of BLV infection was 28.6%, a 4-fold increase compared with the previous study [8]. Although the reason for the spread of BLV infection in Japan was not clarified, it is essential to provide preventive measures to minimize the risk of BLV infection, as well as

to conduct a surveillance study to identify major risk factors.

BLV infects lymphocytes and integrates a DNA intermediate as a provirus into the genome of the cells. The cell-free virus is rarely detected *in vivo* and is thought to appear during acute infection stages before neutralizing antibody production and the period of pregnancy that cause *in utero* infection. [4]. Therefore, most BLV transmission occurs through the transfer of BLV-infected lymphocytes among cattle. Many potential routes of BLV transmission exist; however, the role of colostrum ingestion during natural transmission remains unclear. Previous studies have suggested reciprocal roles. Colostrum and milk from BLV-infected cattle contain BLV-infected cells that may cause infection in neonatal calves from uninfected dams [6, 11, 15]. In contrast, antibodies in the colostrum protect against neonatal infection [10, 13, 15, 18]. In several studies of dairy herds, the transmission rate of colostrum and milk in calves by 6–12 months of age was approximately 6–16% in dairy herds [7]. Feeding management for neonatal calves influenced the risk of colostrum ingestion for the transmission of BLV, although the significance of this risk is uncertain.

An *in vitro* infectivity assay for BLV has been developed using several monolayer cultures, such as fetal bovine lung cells and CC81 cells [9], in which syncytiums are induced. However, because of frequent bacterial content in colostrum, this assay is not considered appropriate for use in studies investigating the inactivation of BLV. On the other hand, sheep are highly susceptible to experimental inoculation

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and are more prone to tumor development at a younger age compared with cattle, in addition to appearing resistant to the bacteria present in colostrum. Therefore, previous studies have successfully demonstrated the utility of using a sheep bioassay to study the frequency with which BLV and BLV infected cells may be detected in colostrum and milk from cows naturally infected with the virus [6, 11].

Both heating and freezing treatments have been considered as techniques to inactivate BLV-infected lymphocytes in colostrum. The effect of heat treatment has been elucidated [2]; however, the effect of freezing treatment has not been elucidated *in vivo* so far. Therefore, here, we used a sheep bioassay to elucidate the effect of freezing colostrum to prevent the transmission of BLV among neonatal calves.

Recipient sheep were born at the Hokkaido Research Station, National Institute of Animal Health. The Animal Care and Use Committee of the National Institute of Animal Health approved all animal procedures prior to initiation of this study. At the start of the study, none of the sheep were positive for BLV antibodies based on an agar gel immunodiffusion (AGID) test (Kitsato Daiichi Sankyo Vaccine, Saitama, Japan) using the virion glycoprotein antigen. The sheep were housed in an insect-proof isolation facility during the experiments. To obtain colostrum, we conducted a serum survey of BLV infection in cattle at several farms and monitored the BLV copy number in the lymphocytes of peripheral blood of pregnant cows at regular intervals. Real-time PCR was performed to determine the BLV copy number by using a CycleavePCR bovine leukemia virus detection kit (TaKaRa, Tokyo, Japan). Colostrum was obtained immediately after parturition from a BLV-infected Holstein cow that had a high BLV copy number in its peripheral blood mononuclear cells (PBMCs) (Table 1). For the inoculum, we used cells isolated from colostrum, which is a more efficient and rapid technique compared with whole milk inoculum [6].

Two liters of colostrum was obtained from cow A, which were separated for the two inoculation treatments, i.e., with and without freezing. For the freezing treatment, colostrum was frozen at -25°C for one night and then thawed. One liter of colostrum from each of the two treatments was centrifuged at $2,800 \times g$ for 15 min to remove butterfat. Then, the aqueous phase and pellets were suspended and centrifuged at $2,800 \times g$ for a further 15 min. The pellets were suspended with phosphate-buffered saline (PBS) and washed once via centrifugation. The pellet cells were resuspended with PBS and were then used to inoculate the sheep intraperitoneally.

Blood samples were obtained from the sheep at weekly intervals, and PCR was used to detect BLV in the lymphocytes [5], while an AGID and passive hemagglutination assay (PHA) (Nisseiken, Tokyo, Japan) were used to detect antibodies to BLV in serum.

Sheep No. 163 was inoculated with cells (approximately 3×10^8 cells) obtained from the 1 l of untreated colostrum of cow A, which had 1,398 copies per 10 ng of DNA of PBMCs. This sheep tested positive for BLV at 3 weeks after inoculation using PCR and at 5 weeks using the AGID and PHA tests. In comparison, sheep No. 164 was inoculated with cells (approximately 4.8×10^8 cells) from 1 l of freeze-thaw-

Table 1. BLV copy number of the peripheral blood mononuclear cells

Pregnant cow	BLV copy number (per 10 ng of DNA) ^{a)}
A	1,398
B	611
C	925

a) BLV copy numbers were determined by using a Cycleave-PCR bovine leukemia virus detection kit (TaKaRa, Tokyo, Japan), which quantified the copy number per 10 ng of DNA in total DNA isolated from peripheral blood mononuclear cells.

Table 2. Results of the genomic and serologic tests of sheep inoculated with cells from the colostrum of cow A

Weeks after inoculation	163 ^{a)}			164 ^{b)}		
	PCR	AGID	PHA	PCR	AGID	PHA
1	—	—	—	—	—	—
2	—	—	—	—	—	—
3	+	—	—	—	—	—
4	+	—	—	—	—	—
5	+	+	+	—	—	—
6	+	+	+	—	—	—
7	+	+	+	—	—	—
8	+	+	+	—	—	—
9	+	+	+	—	—	—

a) No. 163 was inoculated with cells (approximately 3×10^8 cells) from untreated colostrum. b) No. 164 was inoculated with cells (approximately 4.8×10^8 cells) from freeze-thaw-treated colostrum.

treated colostrum from cow A and remained BLV negative in all tests until 9 weeks after inoculation (Table 2). The BLV copy numbers in the untreated and freeze-thaw-treated colostrum of cow A were determined by real-time PCR assay. The assay showed 1.5×10^6 and 1.4×10^6 copies per liter, respectively (data not shown). These results suggest that BLV proviral DNA remained in the inoculum after freeze-thaw treatment. The sheep was inoculated again with cells (approximately 8.9×10^8 cells) from 1 l of freeze-thaw-treated colostrum from cow B, which had 611 copies per 10 ng of DNA of PBMCs, and remained BLV negative in all tests until 29 weeks after inoculation (data not shown). Finally, the sheep, No. 164, was inoculated with cells (approximately 4.8×10^9 cells) from 1.7 liters of untreated colostrum from cow C, which had 925 copies per 10 ng of DNA of PBMCs. The sheep tested positive for BLV at 4 weeks after inoculation using PCR and at 6 weeks for the AGID and PHA tests (Table 3).

A recent seroepidemiological study in Japan confirmed that BLV is more widespread in dairy cattle compared with beef breeding cattle [12]. This difference is probably due to the common practice of feeding bulk milk to calves that are not protected by maternal antibodies at dairy farms [3]. For instance, in Brazilian dairy herds, 27.3% of calves born from BLV-negative dams that were fed pooled milk from BLV-infected cows became BLV positive, whereas similar calves fed pooled milk from BLV-free cows remained negative. In comparison, 16.7% of calves born from BLV-infected

Table 3. Results of the genomic and serologic tests of sheep inoculated with cells from the colostrum of cow C

Weeks after inoculation	164 ^{a)}		
	PCR	AGID	PHA
1	–	–	–
2	–	–	–
3	–	–	–
4	+	–	–
5	+	–	–
6	+	+	+
7	+	+	+

a) No. 164 was inoculated with cells (approximately 4.8×10^9 cells) from untreated colostrum.

dams that received colostrum from their dams and were fed pooled milk from BLV-infected cows became BLV positive, whereas the susceptibility of similar calves fed pooled milk from BLV-free cows was 8.7% [15]. It has been suggested that colostrum and pooled milk increase the risk of BLV transmission, whereas the antibodies in colostrum decrease the risk of infection. Therefore, experiments to reduce the risk of infection and identify the optimal conditions for the treatment of pooled colostrum are required. The results of the current study demonstrated that the freezing treatment for colostrum provides a viable means of inactivating the infectivity of BLV-infected lymphocytes. These data could be used in development of an effective BLV control program by veterinarians and farmers.

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