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Interactions of alfalfa hay and sodium propionate on dairy calf performance and rumen development

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ABSTRACT

The objective of this experiment was to investigate the effects of different levels of alfalfa hay (AH) and sodium propionate (Pro) added to starter diets of Holstein calves on growth performance, rumen fermentation characteristics, and rumen development. Forty-two male Holstein calves (40 ± 2 kg of birth weight) were used in a complete randomized design with a 3 × 2 factorial arrangement of treatments. Dietary treatments were as follows: (1) control = concentrate only; (2) Pro = concentrate with 5% sodium propionate [dry matter (DM) basis]; (3) 5% AH = concentrate + 5% alfalfa hay (DM basis); (4) 5% AH + Pro = concentrate + 5% alfalfa hay + 5% sodium propionate (DM basis); (5) 10% AH = concentrate + 10% alfalfa hay (DM basis); and (6) 10% AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate (DM basis). All calves were housed in individual pens bedded with sawdust until 10 wk of age. They were given ad libitum access to water and starter throughout the experiment and were fed 2 L of milk twice daily. Dry matter intake was recorded daily and body weight weekly. Calves from the control, 10% AH, and 10% AH + Pro treatments were euthanized after wk 10, and rumen wall samples were collected. Feeding of forage was found to increase overall dry matter intake, average daily gain, and final weight; supplementing sodium propionate had no effect on these parameters. Calves consuming forage had lower feed efficiency than those on the Pro diet. Rumen fluid in calves consuming forage had higher pH and greater concentrations of total volatile fatty acids and molar acetate. Morphometric parameters of the rumen wall substantiated the effect of AH supplementation, as plaque formation decreased macroscopically. Overall, the interaction between forage and sodium propionate did not affect calf performance parameters measured at the end of the experiment. Furthermore, inclusion of AH in starter diets positively enhanced the growth performance of male Holstein calves and influenced both the macroscopic and microscopic appearances of the rumen wall. These benefits, however, were small when only sodium propionate was offered.
Key words: calf, forage, rumen development, sodium propionate

INTRODUCTION

Early weaning strategies in dairy calves depend on the consumption of a starter diet and associated development of the rumen. There is disagreement regarding the chemical and physical characteristics of starter diets and the optimal rate of forage provision to preruminant calves (Coverdale et al., 2004; Suarez et al., 2007). Feed concentrates have been formulated for calves to maximize not only DMI and ADG, but also VFA production (Suarez et al., 2006). Ruminal fermentation of dietary concentrates and their end products (especially butyrate and propionate) play a central role in ruminal epithelium differentiation and papilla development (Flatt et al., 1958; Mentschel et al., 2001). In addition, increasing ruminal propionate production by feeding a high-concentrate diet may be associated with improved animal performance and rumen development via its indirect effect on endocrine hormones (Peiris et al., 1998; Liu et al., 2010). However, overloading butyrate and propionate may promote keratinization of papillae by increasing the mitogenic rate and decreasing the apoptosis rate of the epithelium (Flatt et al., 1958; Mentschel et al., 2001).

Because of the lower digestibility of forage in the rumen of calves, the VFA produced are insufficient for optimal growth of papillae and thus, rumen development is slowed (Coverdale et al., 2004). Moreover, an increased forage level in calf diets is reportedly accompanied by decreased ruminal propionate (Suarez et al., 2007; van Ackerena et al., 2009). Research has shown that forage

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has the potential to encourage early rumination and enhance rumen pH and rumen muscle strength, while also reducing cancerous proliferation and keratinization of rumen papilla (Tamate et al., 1962; Klein et al., 1987; Beharka et al, 1998). On the other hand, large amounts of forage in the feed leads to decreased DMI and ADG in calves (Hill et al., 2008, 2009, 2010). A number of studies, however, have reported that feeding forage increases starter intake by the calf and enhances muscular development of the rumen (Coverdale et al., 2004; Khan et al., 2011; Castells et al., 2012).

Based on the above considerations, we hypothesized that adding propionate to diets with alfalfa hay could enhance calf performance and improve rumen conditions. We further postulated that simultaneous use of propionate and forage would be more effective than either offered separately. The objective of current experiment was to examine the effects of, and interactions between, different levels of alfalfa hay and sodium propionate on performance, weaning age, and ruminal fermentation characteristics in calves. Furthermore, macroscopic and microscopic observations in selected treatments were used to determine the influence of the interaction between forage (as the physical factor) and propionate (as the chemical factor) on rumen development.

MATERIALS AND METHODS

The experiment was conducted on a local dairy farm (FKA Agri-Animal Production Co., Isfahan, Iran). Calves were cared for according to the guidelines of the Iranian Council of Animal Care (1995).

Animals, Housing, and Diets

Forty-two male Holstein calves (40 ± 2 kg; mean ± SD) were separated from their mothers, weighed, fed colostrum within 6-h after birth for the first 3 d of age, and then moved to individual pens (1.2 m × 2.4 m) bedded with sawdust. A complete randomized design with a 3 × 2 factorial arrangement was used in the experiments, with different levels of alfalfa hay (**AH**) and propionate (**Pro**) as the experimental factors. The dietary treatments included concentrate and variable ratios of chopped AH with or without sodium propionate salt (143473, Panrac Co., Barcelona, Spain). Dietary treatments were as follows: (1) Control = concentrate only; (2) Pro = concentrate + 5% sodium propionate (DM basis); (3) 5%AH = concentrate + 5% alfalfa hay (DM basis); (4) 5%AH + Pro = concentrate + 5% alfalfa hay + 5% sodium propionate (DM basis); (5) 10%AH = concentrate + 10% alfalfa hay (DM basis); and (6) 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate (DM basis). Starter diets and fresh water were provided ad libitum throughout the experimental period. Concentrate was offered in meal form and AH was chopped by a machine (Golchin Trasher Hay Co., Isfahan, Iran) with a geometric mean particle size of 2.6 mm on the basis of Penn State Particle Separator box values. All diets were formulated to be isonitrogenous and isocaloric and met NRC (2001) requirements for calf nutrients. The ingredient and nutrient compositions of the starters are presented in Table 1. The starter diets were high in grain content and the AH contained 16% CP and 40% NDF. All calves were fed pasteurized waste milk at a rate of 10% of their initial BW (4 kg/head per day). Milk (2.7% protein, 3.4% fat, and SCC of 801 × 1,000 cells/mL) was approximately 38°C when fed via mobile metal bottles (2-L capacity) twice daily at 0800 and 1500 h.

Data Collection, Sampling, and Laboratory Analyses

Starter offered and refused were recorded daily for each calf to determine DMI. Calves were initially weighed at approximately 1200 h on d 0 of the experiment and every 7 d until the end of the experiment period (d 70). Feed efficiency (**FE**) was calculated as kilograms of BW gain per kilogram of total DMI. The structural growth indices considered included body height (distance from base of the rear feet to shoulder bones), hip width (distance between hip bone), and hip height, all measured according to Lesmeister and Heinrichs (2005). These parameters were recorded at birth, weaning, and at the end of the experimental period. Calves were weaned when they consumed 1 kg of their allocated starter for 3 consecutive days.

Rumen Fluid Sampling and VFA Analyses

On d 35 and 70 of age, ruminal contents were collected approximately 3 h postfeeding using a stomach tube. Sample pH was determined immediately using a pH meter (HI 8314 membrane pH meter, Hanna Instruments, Villafranca, Italy). The ruminal contents were subsequently squeezed through 4 layers of cheesecloth to collect rumen fluid samples (10 mL) in tubes, which were then placed on ice and transferred to the laboratory where they were acidified with 3 mL of 25% metaphosphoric acid and stored (–20°C) until analyzed for VFA by gas chromatography (0.25 × 0.32 m, 0.3 μm i.d. fused-silica capillary, model no. CP-9002 Vulcanusweg 259 a.m., Chrompack, Delft, the Netherlands).

Rumen Tissue Sampling and Analyses

On d 70, 9 calves from selected treatments (control, 10%AH, and 10%AH + Pro; 3 calves per treatment)

Table 1. Ingredients and chemical composition (% of DM unless otherwise noted) of experimental starter feeds

Item	Treatment[1]					
	Control	Pro	5%AH	5%AH + Pro	10%AH	10%AH + Pro
Ingredient						
Alfalfa hay	—	—	5.0	5.0	10.0	10.0
Corn grain, ground	61.2	60.0	57.2	56.2	54.2	54.2
Barley grain, ground	7.0	2.0	7.0	2.0	5.0	—
Soybean meal	22.0	23.0	19.0	20.0	19.0	19.0
Extruded soybean	8.0	8.0	10.0	10.0	10.0	10.0
Calcium carbonate	0.9	0.9	0.9	0.9	0.9	0.9
Dicalcium phosphate	0.6	0.8	0.6	0.6	0.6	0.6
Vitamin and mineral premix[2]	0.3	0.3	0.3	0.3	0.3	0.3
Sodium propionate	—	5.0	—	5.0	—	5.0
Composition						
DM, %	89.00	88.00	89.00	89.00	89.00	89.00
CP	20.10	20.30	20.30	19.80	20.30	19.70
NDF	11.60	10.70	13.30	12.30	14.80	14.00
NFC	61.20	62.00	59.10	60.40	57.10	58.80
Fat	4.30	4.20	4.60	4.50	4.60	4.50
Ca	0.64	0.64	0.71	0.70	0.77	0.77
P	0.51	0.51	0.51	0.49	0.50	0.49
ME,[3] Mcal/kg	3.28	3.31	3.20	3.24	3.42	3.43
NE_G,[3] Mcal/kg	1.57	1.58	1.58	1.57	1.60	1.60

[1]Control = concentrate only with no forage and sodium propionate supplemented; Pro = concentrate + 5% sodium propionate; 5%AH = concentrate + 5% alfalfa hay; 5%AH + Pro = concentrate + 5% alfalfa hay + 5% sodium propionate; 10%AH = concentrate + 10% alfalfa hay; 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate.

[2]Contained per kilogram of supplement: 250,000 IU of vitamin A, 50,000 IU of vitamin D, 1,500 IU of vitamin E, 2.25 g of Mn, 120 g of Ca, 7.7 g of Zn, 20 g of P, 20.5 g of Mg, 186 g of Na, 1.25 g of Fe, 3 g of S, 14 mg of Co, 1.25 g of Cu, 56 mg of I, and 10 mg of Se.

[3]Calculated from NRC (2001).

were euthanized to collect digestive tract samples. After removing the digestive tract contents, samples were taken for histological examination from different parts of the rumen including the caudal and ventral portions of the caudoventral blind sac, the cranial and caudal parts of the dorsal sac, and the cranial part of the ventral sac (Figures 1 and 2), according to Lesmeister et al. (2004). The samples were then flushed with normal saline and fixed in 10% buffered formalin (Merck, Darmstadt, Germany) for 96 h. Tissue samples were then dehydrated and cleared by a series of graded alcohols and xylene before being embedded in paraffin. Sections (6 μm) were stained with hematoxylin and eosin for observation under a CX21 light microscope (Olympus, Tokyo, Japan). An Olympus (U-TVO 63XC) camera mounted on the microscope was used to take microphotographs. The length and width of the papillae and the thicknesses of the epithelium, keratin layer, muscle layers, and rumen wall were measured using image analyses computer software (DP2-BSW, version 1.3; Olympus).

Statistical Analyses

Dry matter intake, ADG, and FE were analyzed separately for wk 1 to 6, wk 7 to 10, and over the entire experimental period as a complete randomized design in a 3 × 2 factorial arrangement with 3 levels of AH (0, 5, and 10%) and 2 levels of sodium propionate (0 and 5%). Dry matter intake, ADG, FE, and ruminal fermentation characteristics were statistically analyzed using the following model:

$$Y_{ijkl} = \mu + F_i + P_j + W_k + \left(F \times W\right)_{ik} + \left(P \times W\right)_{jk} + \left(F \times P\right)_{ij} + (F \times P \times W)_{ijk} + \beta\left(X_i - \bar{X}\right) + \varepsilon_{ijkl},$$

where Y_{ijkl} is the dependent variable; μ is the average experimental value; F_i is the effect of forage i (i = 0, 5, and 10%); P_j represents the effect of propionate j (j = diet with or without propionate); W_k is the effect of week k (k = number of weeks); $(F \times P)_{ij}$ designates the effect of the interaction between forage and propionate; $(F \times W)_{ik}$ designates the effect of the interaction between forage and week; $(P \times W)_{jk}$ represents the effect of the interaction between propionate and week; $(F \times P \times W)_{ijk}$ is the tripartite effect of forage, propionate, and week; $\beta\left(X_i - \bar{X}\right)$ designates the covariate variable, where β is the regression coefficient relating the covariate factor to the variable measured, X_i is the covariate factor for the ith subject, and $\bar{X}$ is the overall mean of covariate factor; and ε_{ijkl} is the error term.

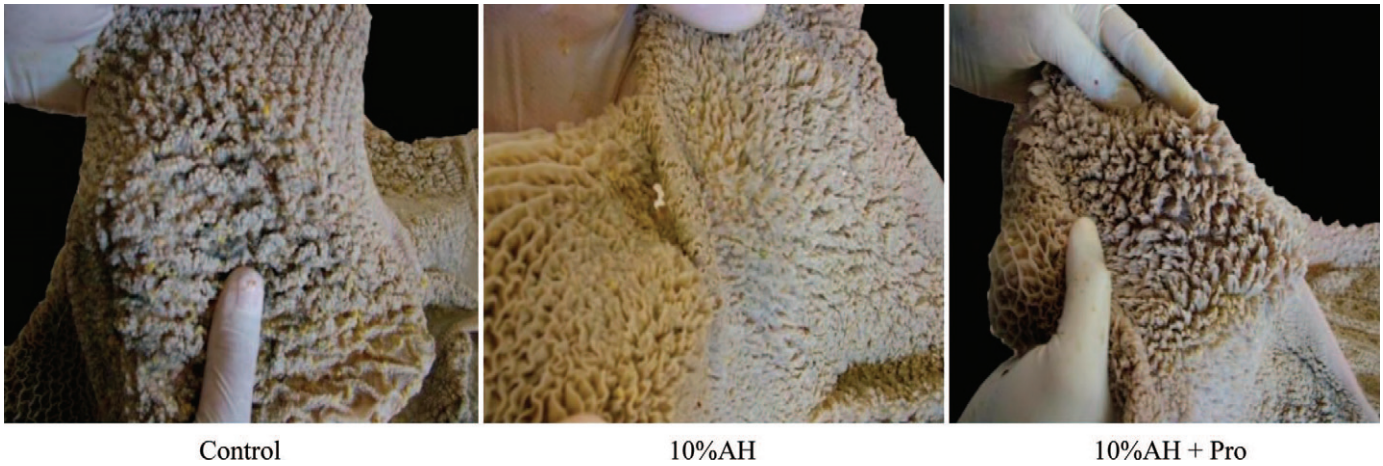

Figure 1. Interior of rumen of selected treatments. Control = concentrate only with no forage and sodium propionate supplemented; 10%AH = concentrate + 10% alfalfa hay; 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate. Color version available in the online PDF.

Calf within treatment was included as a random effect to investigate the main effect of the treatment. Week was modeled as a repeated measurement by using an autoregressive type 1 covariance structure. The lowest level of the Bayesian information criterion (fit statistic) was used to select the covariance structure of the model for each parameter. Weaning day and initial weight were included in the model as covariates for weaning weight. In addition, weaning weight and initial weight were included as covariates for final weight. For skeletal growth parameters, weaning age was included in the model as a covariate. The differences among treatment means were determined using the Tukey multiple range test. Significance was declared at $P \leq 0.05$ unless otherwise noted, and trends declared at $0.05 < P \leq 10$.

Rumen tissue parameters were analyzed as a complete randomized design. Continuous data were subjected to ANOVA according to the following model:

$$Y_{ij} = \mu + \text{diet}_i + \varepsilon_{ij},$$

where Y_{ij} is the dependent variable; μ is the average experimental value; diet is the effect of dietary treatment i (i represents the control diet, 10% AH, and 10% AH + Pro; $i = 1, 2, 3$, respectively); and ε_{ij} is the error term.

RESULTS AND DISCUSSION

Effects of AH on Performance

The values for DMI, ADG, FE, and weaning day, as well as weaning and final weights, are presented in Table 2. Starter intake and total DMI increased with the age of the calves and were affected by AH additions to the diets. Calves fed 10%AH and 10%AH + Pro exhibited greater ($P < 0.01$) DMI values than those fed the control, Pro, and 5%AH + Pro diets. In this study, the higher values of DMI in calves fed the AH diets might have resulted from the higher pH values in the rumen fluid. Previous studies have shown that decreased ruminal pH is associated with decreased starter consumption in calves (Suarez et al., 2006; Khan et al., 2011).

Average daily gain in calves fed the 10%AH + Pro diet was greater ($P < 0.05$) than that for calves fed the Pro and 5%AH diets. Final weight was greater ($P < 0.01$) in calves fed the 10%AH and 10%AH + Pro diets than in those in other treatments. Weaning weight was not affected by treatment.

Forage inclusion in diets has been reported to have mixed effects on ADG: Hill and colleagues (2008, 2010) reported a decreasing effect, whereas others (Coverdale et al., 2004; Khan et al., 2011; Castells et al., 2012) reported an increasing effect on ADG in calves. The differences in forage source, quantity, and particle size might have influenced these differences in ADG and final BW compared with the current study.

Consistent with the linear decline in FE with increasing percentage of hay inclusion reported by Hill et al. (2008), FE decreased when AH was added ($P = 0.003$) in the current research. However, Castells et al. (2012) did not find any differences in FE in dairy calves fed different forage types. In the current research, calves fed the 5%AH, 10%AH, and 10%AH + Pro diets exhibited decreased ($P < 0.01$) values of FE compared with those fed the control and Pro diets. In addition, calves fed 5%AH + Pro showed greater ($P < 0.05$) FE than those fed the 5%AH or 10%AH diets.

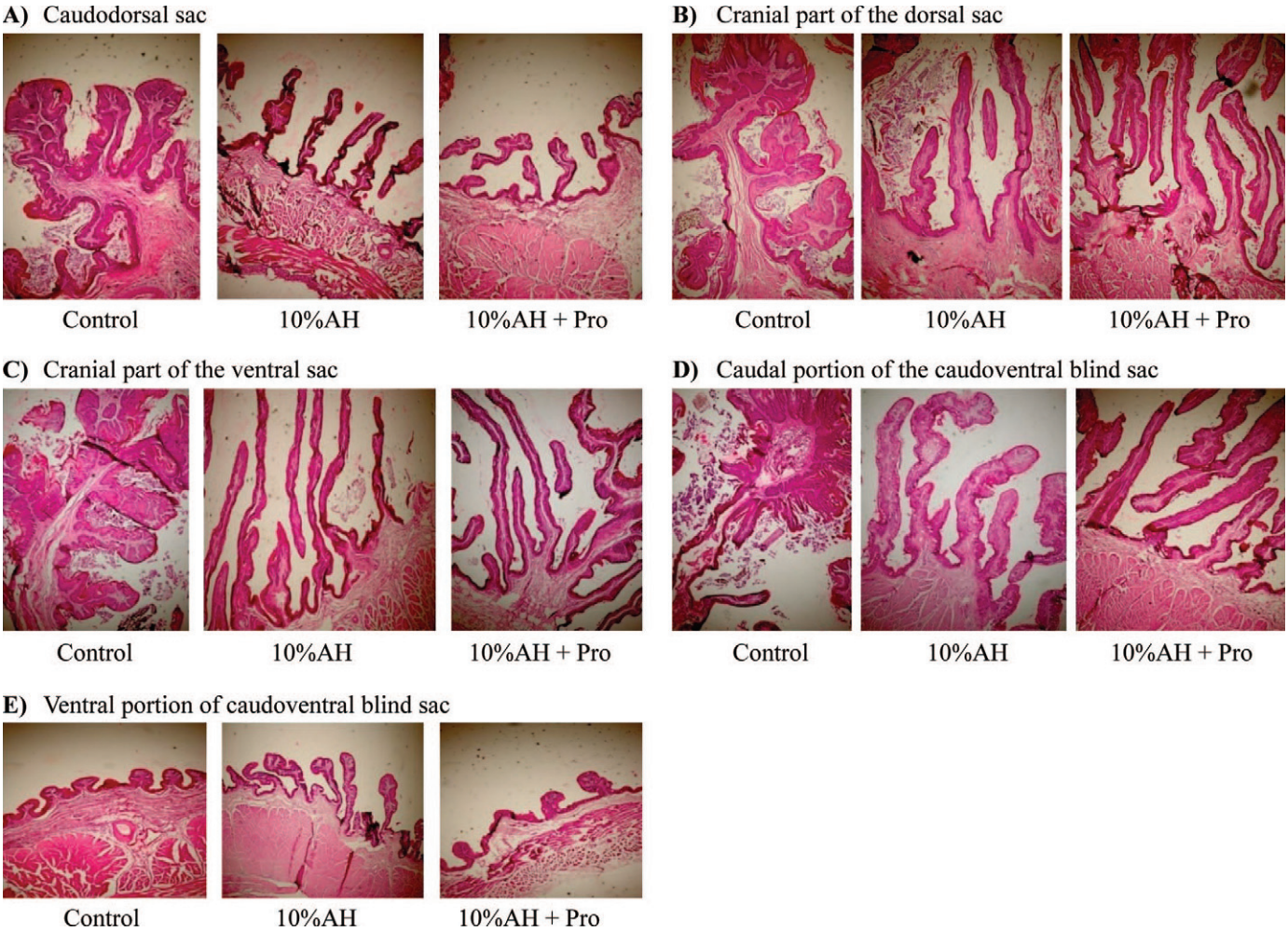

Figure 2. Photomicrographs from all part of rumen bags of selected treatments. A) Caudodorsal sac; B) cranial part of dorsal sac; C) cranial part of ventral sac; D) caudal portion of caudoventral blind sac; E) ventral portion of caudoventral blind sac. Control = concentrate only with no forage or sodium propionate supplemented; 10%AH = concentrate + 10% alfalfa hay; 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate. All sections of the ruminal papillae were stained with hematoxylin and eosin (160×). Color version available in the online PDF.

Weaning age was significantly affected ($P < 0.01$) by AH, with earlier weaning days observed for 5%AH, 10%AH, and 10%AH + Pro treatments compared with control, Pro, and 5%AH + Pro treatments. In contrast to these results, Coverdale et al. (2004) and Porter et al. (2007) reported no differences in the weaning age of dairy calves fed either low or high levels of grass hay and beet pulp. Klein et al. (1987) concluded that consumption of concentrate and roughage together with water at an early age was fundamentally important for both rumen maturation and weaning. In addition, Quigley (1996) stated that the 3 factors involved in weaning were age, BW, and DMI. Despite the positive effect of AH on DMI, not only in wk 1 to 6 but also during the whole 70 d of the current experiment, the reduced weaning age in AH-fed calves in our study might have been due to the earlier and greater DMI allowing early establishment of rumen fermentation and supporting greater amounts of specific VFA for developing rumen epithelium.

The structural growth measures are presented in Table 3. Clearly, alfalfa had no effect on body length, hip height, or hip width; this finding is not consistent with that reported by Hill et al. (2008), who found a linear decline in hip width as hay percentage increased in their experimental diets.

Effects of AH on Ruminal pH and Fermentation Characteristics

According to the ruminal fermentation characteristics presented in Table 4, pH values of sampled rumen fluid were within the range of those previously reported (Khan et al., 2008, 2011). Ruminal pH on d 35 was less ($P < 0.05$) in calves fed the Pro diet than in those fed the 10%AH, 10%AH + Pro, and 5%AH + Pro diets

Table 2. Dry matter intake, ADG, feed efficiency, weaning day, weaning weight, and final weights as influenced by dietary alfalfa hay and sodium propionate supplementation

	Treatment[1]							*P*-value		
Parameter	Control	Pro	5%AH	5%AH + Pro	10%AH	10%AH + Pro	SEM	Forage (F)	Propionate (P)	F × P
Starter intake, kg/d										
wk 1 to 6	0.34[bc]	0.29[c]	0.38[abc]	0.35[bc]	0.50[a]	0.48[ab]	0.05	<0.01	0.49	0.95
wk 7 to 10	1.72[bcd]	1.58[d]	2.04[abc]	1.70[cd]	2.07[ab]	2.44[a]	0.13	<0.01	0.74	0.04
wk 1 to 10	0.89[b]	0.81[b]	1.08[ab]	0.90[b]	1.12[a]	1.25[a]	0.07	<0.01	0.46	0.13
ADG, kg/d										
wk 1 to 6	0.44[a]	0.39[ab]	0.33[b]	0.37[ab]	0.38[ab]	0.40[ab]	0.02	0.06	0.87	0.28
wk 7 to 10	0.74[b]	0.72[b]	0.77[ab]	0.78[ab]	0.85[ab]	0.90[a]	0.05	0.03	0.72	0.76
wk 1 to 10	0.56[ab]	0.52[b]	0.50[b]	0.53[ab]	0.57[ab]	0.60[a]	0.02	0.05	0.72	0.32
Feed efficiency										
wk 1 to 6	0.48[a]	0.47[ab]	0.36[bc]	0.42[ab]	0.32[c]	0.41[ab]	0.03	0.01	0.09	0.30
wk 7 to 10	0.35[ab]	0.38[a]	0.29[b]	0.39[a]	0.35[ab]	0.27[b]	0.02	0.19	0.36	<0.01
wk 1 to 10	0.42[a]	0.43[a]	0.33[c]	0.41[ab]	0.33[c]	0.35[bc]	0.02	<0.01	0.08	0.25
Weaning day	57[a]	61[a]	46[b]	59[a]	45[b]	47[b]	2.70	<0.01	0.01	0.09
Weaning weight	65.58	66.27	64.08	65.03	61.37	63.72	2.91	0.65	0.62	0.93
Final weight	79.85[b]	82.22[b]	80.35[b]	80.20[b]	88.74[a]	87.78[a]	1.69	<0.01	0.76	0.53

[a–d]Means within a row with different superscripts differ ($P < 0.05$).

[1]Control = concentrate only with no forage and sodium propionate supplemented; Pro = concentrate + 5% sodium propionate; 5%AH = concentrate + 5% alfalfa hay; 5%AH + Pro = concentrate + 5% alfalfa hay + 5% sodium propionate; 10%AH = concentrate + 10% alfalfa hay; 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate.

(5.36 vs. 5.79, 5.79, and 5.89, respectively). Moreover, ruminal pH values on d 70 were less ($P < 0.05$) in calves fed control, 5%AH, and 5%AH + Pro diets (5.66, 5.85, and 5.85, respectively) than for those fed the 10%AH diet (6.27). Higher ruminal pH in calves fed the 10%AH diet could be attributed to better rumen conditions, which contribute to increased levels of mastication and salivary flow and suitable VFA blends with lower propionate levels (Khan et al., 2008, 2011). Total VFA concentration was not affected by AH provision on d 35, but was greater ($P < 0.01$) on d 70. Alfalfa hay inclusion increased ($P < 0.05$) acetate concentration on d 35 and 70 of the experimental period and tended to increase ($P = 0.09$) the molar proportion of butyrate on d 35 but not on d 70. We observed no significant effects of AH on the molar proportion of propionate or on the acetate:propionate ratio (Table 4). The higher ruminal total VFA concentrations in calves fed the 10%AH diet compared with those fed the control, Pro, or 5%AH + Pro diets might have been associated with the greater solid feed consumption during the experimental period, the (likely) earlier establishment of ruminal microbes, and better fermentation of OM (Baldwin et al., 2004; Khan et al., 2008; Li et al., 2012).

Table 3. Structural growth parameters in calves fed the different diets[1]

	Treatment[2]							*P*-value		
Parameter	Control	Pro	5%AH	5%AH + Pro	10%AH	10%AH + Pro	SEM	Forage (F)	Propionate (P)	F × P
Body height										
at birth	79.42	80.85	79.35	81.50	81.00	80.57	0.76	0.70	0.10	0.23
at weaning	88.19	89.73	89.58	88.43	89.29	92.05	1.56	0.57	0.45	0.56
at end	92.67	96.05	92.30	93.87	90.84	93.17	1.88	0.61	0.19	0.86
Hip height										
at birth	83.85	84.14	82.78	85.42	84.57	84.14	0.95	0.92	0.29	0.25
at weaning	92.33	92.45	94.49	89.89	95.01	96.44	1.74	0.24	0.54	0.38
at end	101.38	100.81	96.42	98.57	95.40	97.37	1.83	0.14	0.49	0.66
Hip width										
at birth	18.28	19.00	18.71	19.42	19.42	19.43	0.45	0.23	0.20	0.66
at weaning	22.07	21.97	21.96	21.95	21.42	22.78	0.76	0.97	0.53	0.53
at end	25.60	23.40	23.12	22.90	22.62	23.63	1.25	0.49	0.79	0.63

[1]Statistical significance was declared at probabilities $P < 0.05$.

[2]Control = concentrate only with no forage and sodium propionate supplemented; Pro = concentrate + 5% sodium propionate; 5%AH = concentrate + 5% alfalfa hay; 5%AH + Pro = concentrate + 5% alfalfa hay + 5% sodium propionate; 10%AH = concentrate + 10% alfalfa hay; 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate.

Table 4. Ruminal pH and VFA concentration (m*M*) measured at d 35 and 70 in calves receiving different starter diets

	Treatment[1]							*P*-value		
Parameter	Control	Pro	5%AH	5%AH + Pro	10%AH	10%AH + Pro	SEM	Forage (F)	Propionate (P)	F × P
pH										
d 35	5.50[ab]	5.36[b]	5.75[ab]	5.89[a]	5.79[a]	5.79[a]	0.14	0.03	0.98	0.66
d 70	5.66[b]	5.91[ab]	5.85[b]	5.85[b]	6.27[a]	6.13[ab]	0.14	0.03	0.76	0.44
Total VFA, m*M*										
d 35	90.11	86.81	88.18	93.36	91.32	97.82	4.84	0.41	0.49	0.69
d 70	103.91[bc]	96.15[c]	113.70[abc]	106.62[bc]	122.81[a]	113.95[ab]	5.70	0.01	0.11	0.98
Acetate, m*M*										
d 35	50.38[b]	49.16[b]	49.08[b]	50.04[b]	51.62[b]	59.27[a]	2.36	0.03	0.22	0.15
d 70	53.82[ab]	51.92[ab]	50.11[b]	50.36[b]	64.84[a]	63.70[a]	4.73	0.02	0.81	0.97
Propionate, m*M*										
d 35	28.87	32.97	31.65	31.93	30.13	35.02	2.27	0.70	0.12	0.64
d 70	27.39[c]	42.62[a]	29.18[bc]	35.08[abc]	38.57[ab]	35.62[abc]	3.34	0.36	0.04	0.04
Butyrate, m*M*										
d 35	9.62[a]	3.48[b]	12.10[a]	9.51[a]	9.30[a]	7.94[ab]	1.71	0.09	0.03	0.33
d 70	13.87[a]	6.03[b]	13.31[a]	10.12[ab]	11.51[a]	12.80[a]	1.42	0.31	0.01	0.02
A:P ratio										
d 35	1.66	1.55	1.62	1.58	1.58	1.77	0.14	0.86	0.92	0.54
d 70	1.91	1.45	1.68	1.60	1.72	1.98	0.20	0.60	0.61	0.27

[a–c]Means within a row with different superscripts differ ($P < 0.05$).

[1]Control = concentrate only with no forage and sodium propionate supplemented; Pro = concentrate + 5% sodium propionate; 5%AH = concentrate + 5% alfalfa hay; 5%AH + Pro = concentrate + 5% alfalfa hay + 5% sodium propionate; 10%AH = concentrate + 10% alfalfa hay; 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate.

Effects of Pro on Performance

Inclusion of sodium propionate in diets had no effect ($P > 0.10$) on DMI, ADG, or weaning and final weights during the experimental period (Table 2). Our results were in agreement with those obtained by Ferreira and Bittar (2011), who reported that inclusion of calcium propionate in the diet of dairy calves had no effect on DMI, ADG, or weaning and final weights. Feed efficiency tended to increase ($P = 0.08$) and was greater in calves fed the control, Pro, and 5%AH + Pro diets than in calves fed the other treatments. Lee-Rangel et al. (2012) reported that inclusion of calcium propionate in the diets of lambs had no effect on their feed conversion. These observations suggest that the ME supplied by VFA is more efficiently utilized than the same amount of ME supplied by the diet (Poole and Allen, 1970). Inclusion of sodium propionate in diets increased ($P = 0.01$) weaning age in the current study. In addition, weaning and final measures of skeletal growth were not affected ($P > 0.10$) by sodium propionate inclusion in the diets (Tables 2 and 3). These results are consistent with those of Ferreira and Bittar (2011), who reported that calves fed calcium propionate and sodium butyrate exhibited skeletal growth measures similar to those of the control group.

Effects of Pro on Ruminal pH and Fermentation Characteristics

Inclusion of sodium propionate in diets had no effect ($P > 0.10$) on ruminal pH, total VFA, acetate, or

Table 5. Effects of the 3 selected diets on rumen tissue parameters

	Selected treatment[1]				
Parameter (μm)	Control	10%AH	10%AH + Pro	SEM	*P*-value
Papillae length	1,762.6	1,923.5	1,771.1	133.5	0.62
Papillae width	508.3[b]	329.8[a]	374.5[a]	23.3	<0.01
Epithelium thickness	132.9[b]	88.8[a]	96.5[a]	4.9	<0.01
Keratin layer thickness	16.1[b]	11.2[a]	13.0[a]	1.1	<0.01
Muscles layer thickness	1,434.9[b]	1,904.1[a]	1,827.9[a]	118.5	0.01
Rumen wall thickness	1,992.5[b]	2,553.8[a]	2,537.2[a]	141.7	<0.01

[a,b]Means within a row with different superscripts differ ($P < 0.05$).

[1]Control = concentrate only with no forage and sodium propionate supplemented; 10%AH = concentrate + 10% alfalfa hay; 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate.

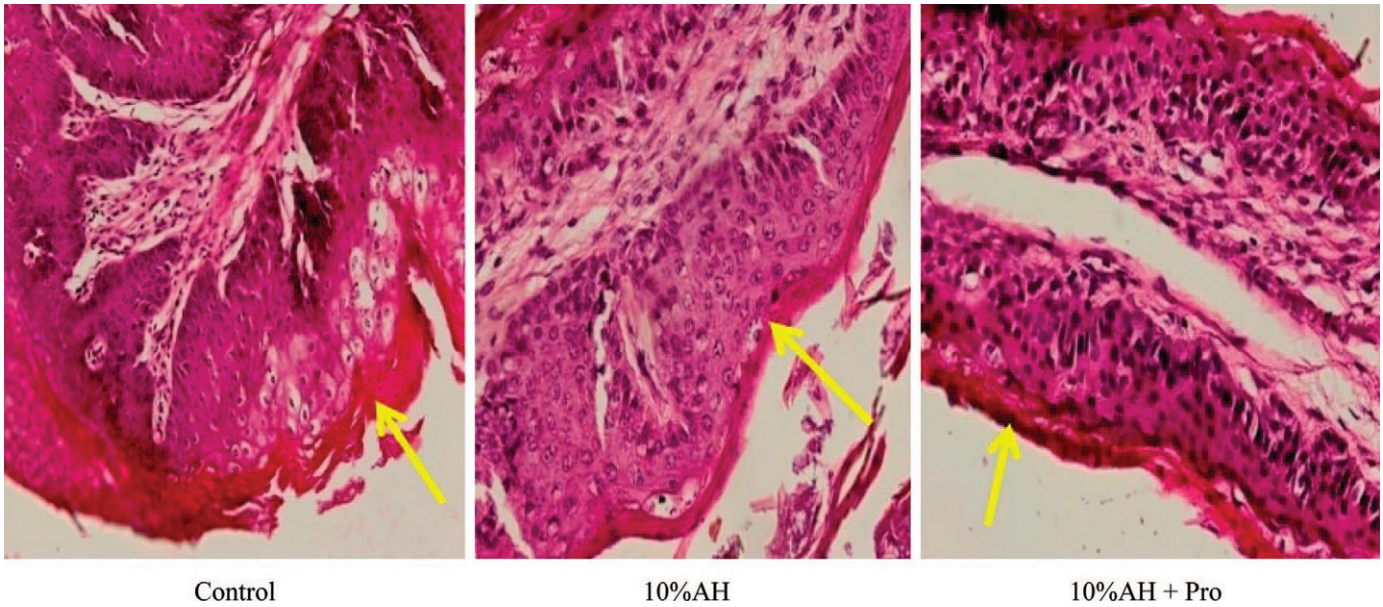

Figure 3. Papilla keratin layer (arrows) from rumen of selected treatments. Control = concentrate only with no forage or sodium propionate supplemented; 10%AH = concentrate + 10% alfalfa hay; 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate. Color version available in the online PDF.

acetate:propionate ratio (Table 4). The molar proportion of ruminal propionate on d 35 was also similar across the treatments. However, the highest ruminal propionate concentrations were observed for the Pro treatment on d 70, which was similar to the level observed by Suárez et al. (2006). They reported that the molar proportion of propionate was >27% when veal calves were fed various concentrates. The increased propionate concentration observed in the rumen of calves fed the Pro diet may be attributed to the supplemental sodium propionate in the diet (Majdoub et al., 2003). Inclusion of sodium propionate in diets decreased ($P < 0.05$) the molar proportion of ruminal butyrate on d 35 and 70 of the experimental period (Table 4). The lowest ruminal butyrate concentrations among the treatments were observed for the Pro diet. These results are in agreement with those of Majdoub et al. (2003), who observed lower butyrate concentrations when propionate was infused into the rumen of growing lambs, perhaps because of a small decline in the protozoa population in the rumen.

Effects of AH and Pro Interaction on Performance

An interaction was observed between AH and Pro for DMI (kg/d) from wk 7 to 10 ($P = 0.04$). Dry matter intake increased ($P < 0.05$) when propionate was added to the high-forage diets (10%AH + Pro) but decreased ($P < 0.05$) when propionate was added to the low-forage diets or those without forage (Pro and 5%AH + Pro; Table 2). Increased DMI has been reported for forage mixed diets (Coverdale et al., 2004; Khan et al., 2011; Castells et al., 2012) but not for VFA salts (specifically propionate salts; Lane and Jesse, 1997; Ferreira and Bittar, 2011; Lee-Rangel et al., 2012). The AH × Pro interaction in calves offered the 10%AH + Pro diet may have improved the rumen environment, which, in turn, may have contributed to the stimulation of starter intake. Simultaneous administration of AH with Pro did not affect ($P > 0.10$) ADG or weaning and final weights, but decreased ($P < 0.01$) FE from wk 7 to 10. Moreover, weaning day tended to increase (P =0.08) as a result of the AH × Pro interaction. Previous studies have shown that higher DMI levels increase passage rate (Dijkstra et al., 2002), which could decrease digestibility of OM (Castells et al., 2012). Therefore, the higher DMI in calves fed the AH diets may be associated with the lower FE. In addition, we observed no interactions during the experimental period between AH and Pro with respect to structural growth parameters (Table 3).

Effects of AH and Pro Interaction on Rumen Fermentation Characteristics

We observed no interactions between AH and Pro on ruminal pH, total VFA, molar proportions of acetate or propionate, or acetate:propionate ratio (Table 4). We observed an interaction of the molar proportion of butyrate on d 70 ($P = 0.02$) with AH and Pro. Changes in ruminal butyrate concentration were observed when

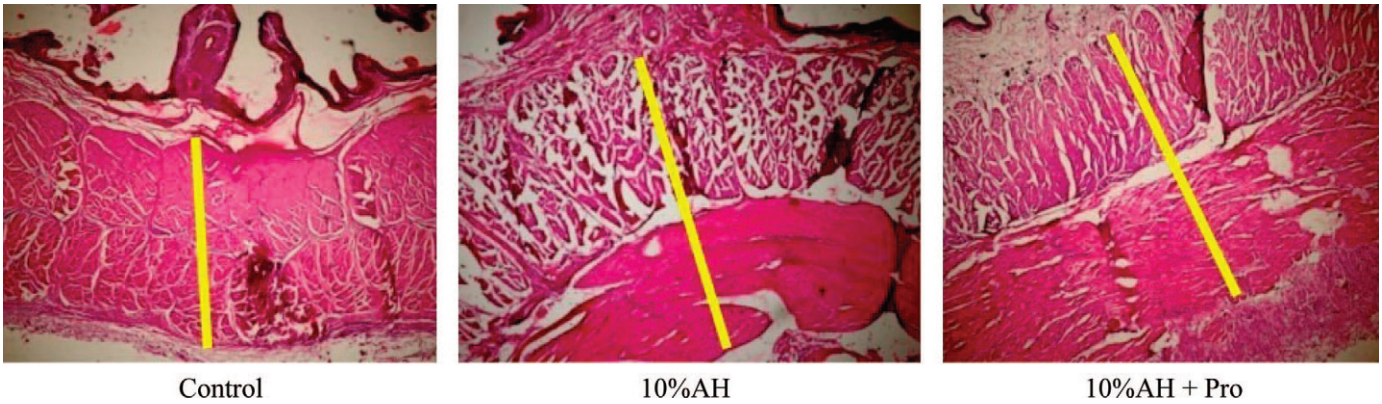

Figure 4. Rumen muscles layer from cranial ventral sac of rumen of selected treatments. Control = concentrate only with no forage or sodium propionate supplemented; 10%AH = concentrate + 10% alfalfa hay; 10%AH + Pro = concentrate + 10% alfalfa hay + 5% sodium propionate. Color version available in the online PDF.

different forage-to-concentrate ratios were offered to young calves (Žitnan et al, 1998). However, no similar published data are available for comparison.

Macroscopic and Microscopic Evaluation of the Rumen Wall

Representative images of the rumen wall are shown in Figure 3, and the associated morphometric parameters are presented in Table 5. The rumen mucosa of calves fed the control diet show focal or multifocal patches with coalescing and adhering papillae covered by a sticky mass of feed, hair, and cell debris. Rumen images from calves in the 10%AH and 10%AH + Pro treatments showed excellent rumen mucosa development with a healthy coloration associated with proper microbial fermentation and increased mucosa development up to 10 wk of age. Limited pitches are also observed in the rumen wall of calves fed the 10%AH + Pro diet.

Although rumen wall papillae length (varying between 1,763 and 1,924 μm) was not affected ($P > 0.10$) by the dietary treatments, papillae width decreased ($P < 0.01$) in calves fed the 10%AH and 10%AH + Pro diets compared with those fed the control diet. Calves fed the 10%AH and 10%AH + Pro diets also had thinner ($P < 0.01$) rumen epithelium and keratin layer thicknesses than those fed the control diet (Figure 3). Moreover, calves fed the control diet had thinner ($P < 0.01$) muscle layers and decreased ($P < 0.01$) rumen wall thicknesses than those fed the other treatments (Table 5, Figure 4). Microscopic views of papillae from different rumen bags of calves fed the selected diets are shown in Figure 2 and those from calves fed the control diet are shown in Figure 5. Similar to our findings, Nocek (1997) and Suárez et al. (2007) reported that calves fed a starter diet without forage exhibited greater levels of plaque formation when judged macroscopically than did control calves. Gäbel et al. (1987) observed that the number of cell layers present in the stratum corneum (keratin layer) was highly dependent on the dietary composition, noting as many as 15 cell layers in thickness when concentrate diets were fed or

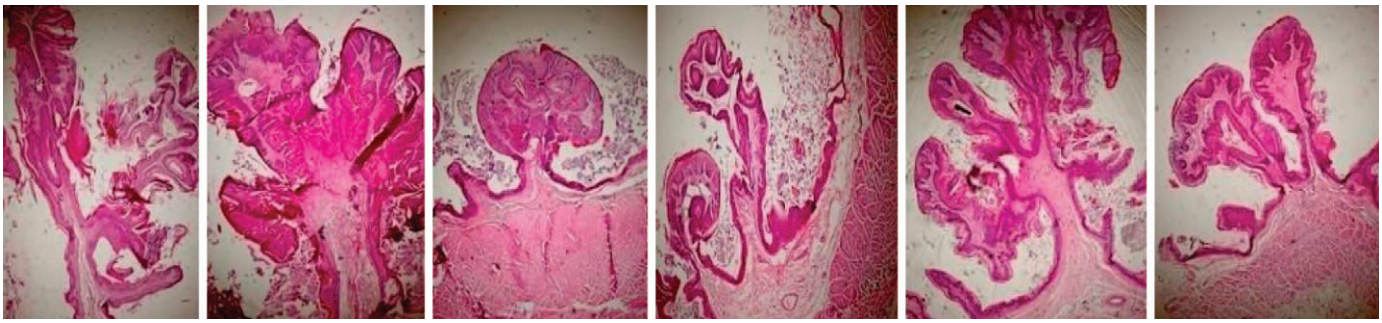

Figure 5. Deformation papillae from different bags of rumen tissue of calves fed the control diet (concentrate only with no forage or sodium propionate supplemented). Sections of ruminal papillae were stained with hematoxylin and eosin (160×). Color version available in the online PDF.

fewer than 4 layers when roughage diets were fed. The thinner keratin layer in the papilla of calves fed the forage mixed starter could be attributed to the abrasive effects of forage (Figure 3). The higher values of rumen wall and muscle layer thicknesses observed in calves fed the starter diet with AH could be attributed to the higher DMI, which can induce greater motility and contractions of the rumen wall due to the forage portion of the starter, thereby giving rise to bulkier and stronger rumen muscles. These results are in agreement with those of Beharka et al. (1998), who reported that physical stimulation by solid feed was needed, not only for the proper development of muscle layers but also for ruminal motility. Similarly, Harrison et al. (1960) demonstrated that calves fed sawdust had 168% greater muscle development in the rumen than those fed only milk. In our experiment, calves fed the 10%AH and 10%AH + Pro diets consumed sufficient DMI to achieve normal rumen development. It is well established that deformed or branching papillae occur due to the effect of uncontrolled ruminal butyrate and propionate on cell proliferations. Our results are in agreement with those of Beharka et al. (1998) and Mentschel et al. (2001). They reported that the greater mitotic index of papilla epithelium in the rumen of calves fed ground diets or diets containing butyrate and propionate might be caused by induced cell proliferation as a strategy to increase the branching or surface area. As shown in calves fed the control diet in the current study, starter diets lacking in components with abrasive properties could result in aggregation of dead cells on the papillae and the formation of keratinized layers, which would eventually lead to parakeratosis.

CONCLUSIONS

Feeding AH increased overall DMI, ADG, and final weight, thereby shortening the time to weaning. This effect of forage was attributable, in part, to its physical property (abrasion effect), which leads to reduced plaque formation in the rumen wall and improves the macroscopic and microscopic appearance of the rumen. However, regardless of the level of AH included in the diet, FE was reduced over the entire experimental period. Supplementing propionate had no effect on performance but did increase weaning age; this might be attributed to the reduced molar proportion of butyrate. Except for the molar proportion of butyrate, we observed no interactions between dietary forage and sodium propionate on performance, skeletal growth, or rumen fermentation characteristics at the end of the experimental period. Forage (physical factor) had a more effective role than sodium propionate (chemical factor) on calf performance and rumen development under current feeding conditions.

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