Bacterial Communities of the Uterus and Rumen During Heifer Development With Protein Supplementation

Taylor B. Ault-Seay1, Kiernan J. Brandt†, Madison T. Henniger1, Rebecca R. Payton1, Daniel J. Mathew1, Sarah E. Moorey1, F. Neal Schrick1, Ky G. Pohler2, Timothy P. L. Smith3, Justin D. Rhinehart1, Liesel G. Schneider1, Kyle J. McLean1* and Phillip R. Myer1*

1 Department of Animal Science, University of Tennessee, Knoxville, TN, United States, 2 Department of Animal Science, Texas A&M University, College Station, TX, United States, 3 United States Meat Animal Research Center, Agricultural Research Service, United States Department of Agriculture, Clay Center, NE, United States

Bacterial communities play major roles in rumen and uterine function toward optimal animal performance and may be affected by changes occurring during heifer development such as nutritional supplementation for optimal growth and the attainment of puberty. The effect of different levels of protein supplementation on ruminal and uterine bacterial communities following weaning was examined through first breeding of heifers. Angus heifers (n = 39) were blocked by initial body weight (BW) and randomly assigned to one of three 163-day (d) crude protein (CP) supplementation diets including control (10% CP, n = 14), 20% CP (n = 11), or 40% CP (n = 14) treatment groups. Growth and development were monitored by body weight, with blood progesterone concentration determined every 14 d to determine pubertal status. Uterine flush and rumen fluid were collected on d 56, 112, and 163 relative to the start of supplementation. Bacterial DNA was extracted from fluid samples, the V1–V3 hypervariable region of the 16S rRNA gene was amplified, and amplicons were sequenced then processed in R 4.1. Statistical analyses were performed in SAS 9.4 with a GLIMMIX procedure utilizing fixed effects of protein, month, pubertal status, and interactions, with random effects including BW, interaction of BW and protein, and heifer within the interaction, and repeated measures of day. In the uterus, pubertal status and day of supplementation affected the observed amplicon sequence variants (ASVs) and led to clustering of samples in a principal coordinate analysis (PCoA; P < 0.05), but no effect of protein supplementation was observed. Ruminal samples clustered in PCoA (P = 0.001), and observed ASVs were impacted over time (P < 0.0001), but no effect of protein supplementation was detected. In contrast, protein supplementation, pubertal status, and day of supplementation affected the abundance of multiple phyla and genera in the uterus and rumen (P < 0.05). Temporal and pubertal status effects on the
heifer’s uterine bacterial communities potentially indicate a maturing uterine microbiome. Protein supplementation did not impact microbial diversity measures but did affect the abundance of individual bacterial phyla and genera that may provide future opportunities to manipulate bacterial community composition and maximize productivity.

**Keywords:** bacteria, beef heifer, development, nutrition, protein, rumen, uterus

**INTRODUCTION**

The development of replacement heifers is crucial for their incorporation into a breeding cow herd. The nutrition provided to heifers during the time of development can affect reproductive efficiency as it influences age at puberty, future success as a breeding cow, and longevity in the herd (Bellows and Short, 1971). Therefore, to support future reproductive performance, it is crucial to provide the required nutrition to meet the demands of the growing heifer. Heifers should ideally reach puberty early, attain approximately 60% of their mature body weight by the first breeding, and achieve a successful pregnancy with as few inseminations as possible to calve by 2 years of age (Larson, 2007; Kuehn et al., 2011; Perry, 2016). Weaned heifers are often developed during a time of dormant forages with lower quality in operations that employ defined breeding seasons. Protein is one of the most vital nutrients for development and reproduction but is often the first limiting nutrient in dormant forages (Marston et al., 1995). Previous studies have shown that supplementing heifers with protein increases their breeding success, muscle growth, and weight gain (Lalman et al., 1993; Martin et al., 2007; Dickinson et al., 2019). Therefore, supplemental feeds high in crude protein are often provided to heifers for continued growth and development for future reproductive success.

The rumen microbiome has been well established as the major producer of energy for the ruminant through the conversions of indigestible feed stuffs to provide volatile fatty acids (VFA), amino acids, metabolites, vitamins, and other useful nutrients (Bergman, 1990). Ruminal bacterial communities can be affected by different feedstuffs that vary in amount and quality of protein, and other components such as fiber and starches (Belanche et al., 2012). Shifts in the rumen bacterial community composition may alter the nutrients metabolized and absorbed for use by the ruminant, which has been shown to have effects systemically and in the reproductive tract through immune system responses (Zebeli et al., 2012; Bilal et al., 2016). Providing supplemented protein in excess has been shown to potentially impact the pH of the reproductive tract by increased plasma urea and ammonia reaching the uterus through histotrophic secretions of the endometrium (Elrod and Butler, 1993; Dawuda et al., 2002). Histotroph contains a variety of nutrients and other factors that contribute to overall uterine health and embryo development (Bazer et al., 2015). Therefore, differences in the nutrient profiles of variable feedstuffs, which impact the rumen microbial communities, could impact future reproductive success by altering the composition of endometrial secretions and the uterine environment via nutrient transporters.

The uterine microbiome of ruminants has been recently determined to be an important contributor to the uterine environment. The reproductive tract was originally thought to be essentially sterile, with the presence of bacteria indicating infection. The emergence of next-generation sequencing has provided the opportunity to recognize the reproductive tract as harboring its own distinct microbiome (Swartz et al., 2014; Laguardia-Nascimento et al., 2015; Clemmons et al., 2017). The composition of the uterine bacterial communities prior to breeding has been associated with differences in breeding outcomes (Ault et al., 2019a; Ault et al., 2019b). For maintenance of healthy bacterial communities in the uterus, the environment must be favorable for microorganisms to perform their metabolic processes, which can benefit reproductive success. However, there has been limited research on the factors that can impact the composition of the uterine bacterial community, potentially compromising the establishment and maintenance of pregnancy. Previous studies have suggested that excess protein and non-protein nitrogen in the diet can result in increased circulating urea and ammonia that can reach the uterus and decrease pH (Elrod and Butler, 1993; Dawuda et al., 2002; Ocon and Hansen, 2003). The observed impacts of diet on the uterine environment in cattle suggests that nutrition may influence the bacterial communities in the uterus. Additionally, differences in uterine bacterial communities have been detected between different phases of the estrous cycle and throughout estrus synchronization protocols (Quereda et al., 2020; Ault et al., 2019a; Ault et al., 2019b). Thus, the physiological changes that occur through the attainment of puberty and changes in hormone response in heifers may affect the uterine bacterial communities present. Through heifer development, the impact of nutrition on the uterine environment may provide the opportunity to influence uterine bacterial communities toward optimal reproductive health and efficiency.

The objective of the current study was to evaluate the effects of different levels of protein supplement on ruminal and uterine bacterial communities following weaning (pre-pubertal) through the first breeding (pubertal) of beef heifers. We hypothesized that 1) protein supplementation will affect developing heifer’s ruminal and uterine bacterial communities during development, and 2) bacterial communities will differ between prepubertal and pubertal heifers.
MATERIALS AND METHODS

All animal experimental procedures for the current study were approved by the Institutional Animal Care and Use Committee at the University of Tennessee, Knoxville.

Experimental Design

Commercial Angus heifers (n = 60) housed at the Middle Tennessee Research and Education Center were enrolled in the study 23 days (d) following weaning. Figure 1 depicts the following study timeline with sampling and estrus synchronization days. On d 0, heifers were evaluated for baseline measurements including body weight (BW), body condition scores (BCS), and blood samples via the jugular vein for determination of pubertal status. Heifers were blocked by BW and assigned to a treatment group (n = 20 heifers per treatment): control supplement (CON; 1.59 kg corn per day), 20% crude protein (CP) supplement (P20; 1.59 kg per day: 75% dried distillers’ grains 25% corn), or 40% CP supplement (P40; 1.59 kg per day: 75% soybean meal 25% dried distillers’ grains). Heifers were maintained in 1.21-hectare pens with five heifers per pen resulting in four pens per treatment group. Supplements were provided four times each week for 140 d, beginning in early October through late March. Body weight, BCS, and blood samples were collected once every 2 weeks to monitor development and pubertal onset. Blood samples were centrifuged at 5,000 × g and 4°C for 20 min to collect serum. Serum samples were stored at -80°C until further analysis.

Circulating progesterone (P4) samples were quantified in serum using the Double Antibody RIA Kit (MP Biomedicals, Irvine, CA) validated previously by Pohler et al. (2016). The progesterone concentration of ≥1 ng/ml in two consecutive samplings that corresponded with a normal estrous cycle indicated a functional corpus luteum, representing puberty (Polat et al., 2009). The heifer was considered pubertal at the time of the second sample of ≥1 ng/ml P4. Serum P4 concentrations at puberty, age at puberty, BW, and BCS for all heifers throughout development, with supplement and total diet feed analyses, were reported in Brandt (2020). Cytokine concentrations of the uterine flush leading up to breeding for all heifers were reported in Ault-Seay et al. (2021).

Only prepubertal heifers (n = 39) were selected to continue through the remainder of the study based on serum P4 concentrations from the four blood collections between d 0 through 42 (Figure 1), resulting in 14 heifers from the CON and P40 treatment groups and 11 heifers from the P20 treatment group. Uterine flushes and rumen fluid were collected every other month on d 56, 112, and 163, relative to the start of protein supplementation, to evaluate the bacterial communities present. For uterine flushes, a Foley catheter was inserted through the vagina and cervix into the uterus to deliver 20 ml of sterile saline, manipulated throughout the uterus by rectal massage, then collected by syringe (Clemmons et al., 2017). Rumen fluid (~50 ml) was collected by oroesophageal tubing into the rumen attached to a hand pump with resulting fluid collected into a flask (Guan et al., 2008). Foley catheters and oroesophageal tubing were disinfected and washed between each heifer to eliminate contamination. Uterine flush and rumen fluid were immediately placed on ice for transport then stored at -80°C until bacterial community analysis.

Prior to the final uterine flush and rumen fluid collections on d 163, heifers were subjected to estrus synchronization utilizing an industry standard 7-Day Co-Synch + CIDR protocol beginning on d 140. Controlled intravaginal drug release devices (CIDR; 1.38 g P4; Zoetis Animal Health, Florham Park, NJ) were inserted, and gonadotropin-releasing hormone (100 µg; Cystorelin, Boehringer Ingelheim, Duluth, GA) was administered i.m. on d 140. An injection of prostaglandin F2α (500 µg; cloprostenol sodium; Synchsure, Boehringer Ingelheim, Duluth, GA) was administered i.m. at CIDR removal on d 147. Approximately 52 h after CIDR removal, heifers were administered GnRH, followed by artificial insemination on d 149.
**Bacterial DNA Extraction, Library Preparation, and Sequencing**

Uterine flush bacterial DNA extraction was performed according to methods from Clemmons et al. (2017) and recommendations for low-biomass samples (Weinroth et al., 2022). Samples were thawed to room temperature (22°C), and 5-mL aliquots were placed in 15-mL conical tubes for centrifugation at 5,000 × g and 4°C for 10 min. The supernatant was removed, and the resulting pellet was resuspended in enzymatic lysis buffer (4% (w/v) sodium dodecyl sulfate (SDS), 500 mM NaCl, and 50 mM EDTA) for incubation at 37°C. Bacterial DNA was extracted using the DNEasy Blood and Tissue kit (Qiagen, Hilden, Germany) according to manufacturer protocol. Extracted DNA concentrations were determined using a spectrophotometer (DeNovix, Wilmington, DE, USA) then stored at -20°C.

Bacterial DNA was extracted from rumen fluid using a modified method validated by Yu and Morrison (2004). Samples were thawed to room temperature (22°C), and approximately 0.2 g of rumen fluid was transferred to a ZR BashingBead Lysis Tube (Zymo Research Corp., Santa Ana, CA, USA) with lysis buffer for chemical lysis. Samples were placed in the TissueLyser II system (Qiagen, Hilden, Germany) for mechanical lysis at 21 Hz for 3 min. Cell debris was removed using 10 M ammonium acetate, then nucleic acids were precipitated using isopropanol. Proteinase K and RNase were used to remove all proteins and RNA. Lastly, DNA samples were purified by centrifugation in QIAamp columns from the QIAamp DNA Stool Mini Kit (Qiagen, Hilden, Germany). Resulting DNA concentrations were quantified via using a spectrophotometer (DeNovix, Wilmington, Delaware) then stored at -20°C.

Library preparation and sequencing were performed at the US Meat Animal Research Center (Clay Center, NE, USA). Bacterial DNA was amplified using polymerase chain reaction (PCR) with AccuPrime Taq high-fidelity DNA polymerase (Life Technologies, Carlsbad, CA, USA) for library preparation. Primers 27F (5′-Adapter/Index/AGATGTGTAGCTCTTCCGCTCACG TGAGT) and 519R (5′-Adapter/Index/GTATTACCCTGCCTCCTGCTGCTG) with TruSeq indices were used to target the V1–V3 hypervariable region of the bacterial 16S rRNA gene (Myer et al., 2015). The resulting rumen and uterine bacterial DNA libraries were sequenced using the 2 × 300, v3 600-cycle kit on the Illumina MiSeq platform (Illumina, Inc., San Diego, CA, USA).

**Sequence Processing and Statistical Analyses**

Sequence quality filtering, trimming, and taxonomic assignment were performed in R 4.1 using the dada2 pipeline (Callahan et al., 2016). Forward sequences were filtered with standard dada2 parameters and a truncation length of 270. Resulting sequences were used to construct an amplicon sequence variant (ASV) table. Chimeric sequences were identified and removed, then taxa were assigned using the SILVA database version 138 (Quast et al., 2013). Metadata were imported and merged with the taxa table for alpha and beta diversity analyses using the phyloseq package (McMurdie and Holmes, 2013) in R 4.1. Unassigned reads were classified to their last known taxonomic classification, and sequences assigned as Eukaryota or Cyanobacteria were removed for all downstream analyses. Alpha diversity indices were calculated for observed species by observed ASVs, and richness and evenness using the Shannon Diversity Index. Beta diversity analyses utilized the principle coordinate analysis (PCoA) method with Bray Curtis distances. Visualizations were generated with ggplot2 including ellipses representing a 95% confidence interval for each cluster. Relative abundance plots were generated for taxa using ggplot2 for the top 10 phyla and genera with a relative abundance ≥1%. Genera with a relative abundance <1% were combined to one group for visualization.

For beta diversity, the vegan package (Oksanen et al., 2017) with the adonis function in R 4.1 was used to perform a PERMANOVA with 999 permutations to determine the significance of each PCoA. Statistical analyses to determine differences in alpha diversity indices and bacterial taxa abundances were performed in SAS 9.4 (SAS Institute, Cary, NC, USA). Normality of observed ASVs and bacterial taxa raw abundance values were evaluated by the univariate procedure. Data were determined to be non-normal with a Shapiro–Wilk value of W < 0.80. Non-normal data were normalized by log transformation to achieve a normal distribution and Shapiro–Wilk value of W > 0.85. A completely randomized block design with repeated measures was utilized with the GLIMMIX procedure including fixed effects of day of sampling, protein supplementation treatment, and pubertal status, and interactions of day × protein and protein × pubertal status. Random effects included the block of BW, interaction of BW × protein, and heifer within the interaction, with repeated measures of day. For bacterial taxa abundance data, means are reported as the mean raw abundance value × 100 to indicate percent relative abundance out of 100%. Significance was determined by P ≤ 0.05 and trends by 0.05 < P ≤ 0.10 for all analyses.

**RESULTS**

Cluster analysis of 16S rRNA amplicon sequences by PCoA among all samples indicated distinct separation in beta diversity of microbial communities between the uterus and rumen (R2 = 0.13, P = 0.001, Figure 2). Rumen samples clustered tightly indicating a high degree of similarity of bacterial community composition among all rumen samples. Uterine samples clustered loosely indicating higher dissimilarity of microbial communities among uterine samples (Figure 2). A total of 39 phyla and 1,633 genera were detected among all uterine and ruminal samples. Supplementary Figure 1 illustrates the overall relative abundance of the top 10 phyla detected among all samples between the uterus and rumen. Firmicutes and Bacteroidota were the most abundant phyla in the uterus and rumen, respectively, independent of protein supplementation treatment, pubertal status, or day of sampling. Supplementary Figure 2 depicts genera with a relative abundance ≥1% in the uterus and rumen, with all genera present in <1% grouped.
Twenty-three genera in the uterus and 25 genera in the rumen were detected with relative abundances ≥ 1%. The total average relative abundance of genera with <1% abundance was 30.27% in the uterus, and 20.01% in the rumen. *Prevotella* was the most abundant genus in both uterine and ruminal samples across all groups.

**Protein Supplementation Level**

The level of protein supplementation did not affect alpha diversity metrics, observed ASVs, or Shannon’s Diversity Index, in the uterus or rumen (Table 1). Similarly, no clear clustering pattern by protein supplementation level was observed by the beta diversity PCoA in uterine (R² = 0.02, P = 0.06, [Figure 3A](#)) or ruminal (R² = 0.02, P < 0.01, [Figure 3B](#)) samples, despite calculated significance in the rumen. However, multiple phylum and genus relative abundances were affected in both the uterus and rumen by protein supplementation level.

In the uterus, Firmicutes was the most abundant phyla and was affected by protein treatment, with P20 having the greatest and P40 having the least abundance (P = 0.05; Table 2). Fibrobacterota and Proteobacteria abundances were affected by protein supplementation (P ≤ 0.01) and tended to affect the abundance of Chloroflexi (P = 0.09; Table 2). In the rumen, Bacteroidota abundance increased with increasing crude protein as P40 heifers had the greatest abundance, and control heifers with the least (P = 0.02; Table 2). Actinobacteriota abundance in the rumen was also affected by protein supplementation, with P20 heifers having the greatest abundance (P = 0.02; Table 2). Additionally, Firmicutes, Proteobacteria, and Spirochaetota abundances tended to be affected by protein supplementation (P < 0.10; Table 2).

Relative abundances of genera *Alistipes*, *Bacteroides*, *Fibrobacter*, *Prevotella*, UCG-005, and unclassified genera from Family Bacteroidales BS11 and Family Lachnospiraceae were

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**TABLE 1** | Impact of protein supplement, pubertal status, and day of supplementation on alpha diversity metrics observed ASVs and Shannon’s Diversity Index in the uterus and rumen

| Fixed effect                  | Uterus                      | Rumen                      |
|-------------------------------|-----------------------------|---------------------------|
|                               | Observed ASVs | Shannon’s | Observed ASVs | Shannon’s |
| Protein supplement            |               |          |               |          |
| CON                           | 6584 ± 373     | 7.28 ± 0.24 | 2682 ± 95     | 7.13 ± 0.04 |
| P20                           | 5921 ± 394     | 7.16 ± 0.26 | 2684 ± 100    | 7.09 ± 0.05 |
| P40                           | 6263 ± 353     | 7.00 ± 0.23 | 2690 ± 89     | 7.12 ± 0.04 |
| P value                       | 0.48           | 0.69     | 0.99          | 0.81     |
| Pubertal status               |               |          |               |          |
| Pre-pubertal                  | 5499 ± 341 b   | 6.75 ± 0.23 | 2709 ± 86     | 7.13 ± 0.04 |
| Pubertal                      | 7013 ± 449 a   | 7.54 ± 0.30 | 2662 ± 111    | 7.10 ± 0.05 |
| P value                       | 0.03           | 0.78     | 0.65          | 0.65     |
| Day of supplementation        |               |          |               |          |
| 56                            | 4738 ± 490 b   | 7.10 ± 0.34 | 2161 ± 117 b  | 6.88 ± 0.05 b |
| 112                           | 7178 ± 381 a   | 7.03 ± 0.26 | 3367 ± 92 a   | 7.30 ± 0.04 a |
| 163                           | 6852 ± 455 a   | 7.31 ± 0.31 | 2529 ± 108 b  | 7.17 ± 0.05 a |
| P value                       | 0.002          | 0.81     | < 0.0001      | <0.0001 |

1. a,b represents P ≤ 0.05.
2. Alpha diversity values presented as mean ± SEM.
affected by protein supplementation in the uterus \((P < 0.05; \text{Table 2})\). The most abundant uterine genus, \textit{Prevotella}, had the greatest abundance in the P40 supplement group, with lesser, similar abundances in the CON and P20 groups \((P = 0.02; \text{Table 2})\). Similarly in the rumen, \textit{Prevotella} was the most abundant genus and increased in abundance with increasing crude protein level of the supplement \((P = 0.05; \text{Table 2})\). The protein supplementation level also affected relative abundances of the unclassified genera from Family Lachnospiraceae and Family UCG-010 \((P \leq 0.01)\) in the rumen and tended to affect UCG-004 \((P = 0.10; \text{Table 2})\).

**Pubertal Status**

In the uterus, observed ASVs and Shannon’s Diversity Index were affected by pubertal status. Pubertal heifers had a greater number of observed ASVs \((P = 0.03)\) and tended to have a higher Shannon Diversity Index value \((P = 0.08)\), than prepubertal heifers \((\text{Table 1})\). Clustering was observed in the PCoA by

**TABLE 2** | Bacterial phyla and genera with relative abundance ≥1% in the uterus and rumen impacted by protein supplementation treatment.

| Environment | Taxonomy | Phyla | Protein supplementation treatment1 | \(P\) |
|-------------|----------|-------|-----------------------------------|------|
| Uterine     | Phyla    |       | CON                               |      |
|             |          |       | P20                               |      |
|             |          |       | P40                               |      |
|             | Genera   |       |                                    |      |
|             | Chloroflexi | 0.12 ± 0.01 | 0.122 ± 0.01 | 0.09 ± 0.01 | 0.09 |
|             | Fibrobacterota | 1.40 ± 0.13 | 1.44 ± 0.14 | 1.02 ± 0.12 | 0.008 |
|             | Firmicutes | 48.92 ± 2.88 | 52.72 ± 2.88 | 45.35 ± 2.61 | 0.05 |
|             | Proteobacteria | 3.20 ± 0.59 | 2.53 ± 0.62 | 4.99 ± 0.56 | 0.01 |
|             | Alistipes | 1.36 ± 0.23 | 1.27 ± 0.25 | 0.74 ± 0.22 | 0.04 |
|             | Bacteroides | 1.51 ± 0.22 | 1.31 ± 0.23 | 0.70 ± 0.21 | 0.006 |
|             | Family Bacteroidales BS11 gut group | 1.04 ± 0.10 | 0.88 ± 0.11 | 1.21 ± 0.10 | 0.03 |
|             | Family Lachnospiraceae | 5.12 ± 0.38 | 4.19 ± 0.40 | 3.22 ± 0.36 | <0.001 |
|             | Fibrobacter | 1.39 ± 0.13 | 1.43 ± 0.14 | 1.91 ± 0.19 | 0.008 |
|             | Prevotella | 12.08 ± 1.22 | 11.06 ± 1.30 | 15.09 ± 1.18 | 0.02 |
|             | UCG-005 | 4.89 ± 0.61 | 4.45 ± 0.64 | 2.29 ± 0.58 | 0.006 |
| Rumen       | Phyla    |       |                                    |      |
|             | Actinobacteriota | 0.07 ± 0.01 | 0.10 ± 0.01 | 0.07 ± 0.01 | 0.02 |
|             | Bacteroidota | 64.26 ± 0.77 | 65.10 ± 0.82 | 67.39 ± 0.74 | 0.02 |
|             | Firmicutes | 28.11 ± 0.77 | 27.32 ± 0.82 | 25.65 ± 0.74 | 0.07 |
|             | Proteobacteria | 1.89 ± 0.14 | 1.46 ± 0.15 | 1.52 ± 0.14 | 0.06 |
|             | Spirochaetaota | 0.52 ± 0.03 | 0.44 ± 0.03 | 0.41 ± 0.03 | 0.07 |
|             | Genera   |       |                                    |      |
|             | Family Lachnospiraceae | 2.43 ± 0.08 | 2.01 ± 0.08 | 1.99 ± 0.08 | <0.001 |
|             | Family UCG-010 | 0.97 ± 0.04 | 1.15 ± 0.04 | 1.00 ± 0.04 | 0.01 |
|             | Prevotella | 28.03 ± 1.18 | 29.38 ± 1.26 | 31.95 ± 1.14 | 0.05 |
|             | UCG-004 | 1.36 ± 0.13 | 1.29 ± 0.14 | 0.95 ± 0.13 | 0.1 |

1 Relative abundances presented as mean (%) ± SEM.

\(ab\) means within row different at indicated \(P\).
pubertal status in uterine samples ($R^2 = 0.03$, $P = 0.001$), depicting high variation in prepubertal heifers and tighter clustering of pubertal heifers (Figure 4A). However, pubertal status did not impact rumen alpha diversity metrics or beta diversity PCoA clustering ($R^2 = 0.02$, $P > 0.05$; Figure 4B).

Pubertal status affected the abundance of the most abundant uterine bacterial phyla, Firmicutes, with prepubertal heifers having the greatest abundance compared to pubertal heifers ($P = 0.04$; Table 3). The abundance of Firmicutes was less in pubertal heifers, although multiple phyla were greater or tended to be greater in pubertal heifers including Bacteroidota ($P = 0.04$), Proteobacteria ($P = 0.04$), and Verrucomicrobiota ($P = 0.07$; Table 3). In the rumen, pubertal status did not affect the abundance of Bacteroidota, but did affect Actinobacteria, Firmicutes, and Proteobacteria ($P < 0.05$; Table 3). Similar to the uterus, Firmicutes was also decreased in the rumen, while Proteobacteria was increased in pubertal heifers compared to prepubertal heifers (Table 3).

The relative abundance of unclassified genera from Family F082 ($P = 0.04$) was affected in the uterus by pubertal status and tended to affect Prevotellaceae-UCG001, Ureaplasma, and unclassified genera from Family Bacteroidales BS11 and Family Prevotellaceae ($P < 0.10$; Table 3). All of these genera had a greater relative abundance in pubertal heifers, except Ureaplasma (Table 3). Puberty affected the ruminal relative abundance of NK4A214 and Succiniclasticum, which were greater in prepupalheifers and unclassified genera from Order Rhodospirillales which was greater in pubertal heifers ($P < 0.05$; Table 3).

**Day of Supplementation**

Observed ASVs in the uterus were affected by day of supplementation. The least number of observed ASVs was

**TABLE 3 | Bacterial phyla and genera with relative abundance ≥1% in the uterus and rumen impacted by pubertal status.**

| Environment | Taxonomy | Pubertal status | $P$ |
|-------------|----------|-----------------|-----|
| Uterus      | Phyla    | Prepubertal     | Pubertal |
|             | Bacteroidota | 37.19 ± 2.19 | 43.13 ± 2.66 | 0.06 |
|             | Elusimicrobiota | 0.10 ± 0.01 | 0.14 ± 0.01 | 0.05 |
|             | Firmicutes | 53.85 ± 2.53 | 44.15 ± 3.21 | 0.04 |
|             | Proteobacteria | 2.51 ± 0.54 | 4.64 ± 0.72 | 0.04 |
|             | Verrucomicrobiota | 0.77 ± 0.06 | 0.98 ± 0.07 | 0.07 |
|             | Genera | Prepubertal | Pubertal |
|             | Family Bacteroidales BS11 gut group | 0.90 ± 0.10 | 1.18 ± 0.11 | 0.07 |
|             | Family F082 | 4.45 ± 0.39 | 5.90 ± 0.49 | 0.04 |
|             | Family Prevotellaceae | 1.35 ± 0.12 | 1.75 ± 0.15 | 0.08 |
|             | Prevotellaceae-UCG001 | 1.48 ± 0.13 | 1.85 ± 0.16 | 0.09 |
|             | Ureaplasma | 13.78 ± 3.19 | 4.46 ± 4.25 | 0.07 |
| Rumen       | Phyla    | Prepubertal | Pubertal |
|             | Actinobacteria | 0.09 ± 0.01 | 0.07 ± 0.01 | 0.03 |
|             | Firmicutes | 28.90 ± 0.73 | 25.15 ± 0.97 | 0.01 |
|             | Proteobacteria | 1.33 ± 0.13 | 1.91 ± 0.17 | 0.02 |
|             | Genera | Prepubertal | Pubertal |
|             | NK4A214 group | 1.04 ± 0.06 | 0.83 ± 0.07 | 0.03 |
|             | Order Rhodospirillales | 0.98 ± 0.10 | 1.53 ± 0.13 | 0.006 |
|             | Succiniclasticum | 1.04 ± 0.07 | 0.76 ± 0.08 | 0.007 |

1 Relative abundances presented as mean (%) ± SEM.

ab means within row different at indicated $P$. 

**FIGURE 4 | Beta diversity by principal coordinate analysis with Bray Curtis distances for pubertal status in the uterus (A; $R^2 = 0.03$, $P = 0.001$) and rumen (B; $R^2 = 0.02$, $P = 0.01$). Significance was determined by PERMANOVA with 999 permutations. Ellipses represent 95% confidence interval.**
detected on d 56 then significantly increased to the greatest observed ASVs by d 112 and remained similar on d 163 ($P < 0.001$; Table 1). The Shannon Diversity Index followed a similar trend but was not significant (Table 1). Ruminal observed ASVs and Shannon’s Diversity were affected by day, with the greatest number detected on d 112 and less observed ASVs on d 56 and d 163 ($P < 0.0001$; Table 1).

Beta diversity analysis by PCoA indicated uterine samples clustered by day with overlap ($R^2 = 0.10$, $P = 0.001$, Figure 5A). Samples on d 56 had the greatest variation overlapping with d 112 and 163 uterine samples. However, d 112 and 163 samples clustered tightly and depicted the greatest separation between days, with slight overlap (Figure 5A). Rumen samples had similar PCoA clustering by day as the uterus by day with overlap ($R^2 = 0.08$, $P = 0.001$, Figure 5B). The greatest separation was observed between d 56 and 112 indicating the greatest difference in overall community composition.

In the uterus, day of supplementation tended to affect the abundance of Firmicutes ($P = 0.09$, Table 4). Multiple additional phyla also shifted over time including Actinobacteria, Chloroflexi, and Patescibacteria ($P < 0.05$; Table 4), with a tendency for Fibrobacterota ($P = 0.07$). All top 10 phyla detected in the rumen were affected by the day of sampling ($P < 0.05$; Table 4), except Fibrobacterota. The most abundant phyla in the rumen, Bacteroidota, had the greatest abundance on d 56, with lesser, similar abundances on d 112 and 163 ($P < 0.0001$; Table 4). The majority of genera in the rumen and uterus had shifts in their abundance over time; therefore, Table 5 indicates the top 5 most abundant genera that shifted over time in each environment, with all other affected genera listed in Supplementary Table 1. There were 23 genera detected in the uterus with a relative abundance ≥1%, of which 16 shifted over time ($P \leq 0.10$). All rumen genera detected with relative abundance ≥1% shifted over time ($P \leq 0.10$), except for unclassified genera from Family Prevotellaceae.

### Interactions of Protein Supplementation, Day of Supplementation, and Pubertal Status

The triple interaction of protein supplementation level, day of supplementation, and pubertal status, and the interaction of day of supplementation × pubertal status, was not able to be analyzed in the current study due to all heifers being prepubertal at the first sampling on d 56. Alpha diversity metrics in the uterus and rumen had no significant interaction of protein supplementation level × day of supplementation, or for any of the top 10 uterine bacterial phyla relative abundances. For uterine genera, however, a protein supplementation × day interaction was observed for UCG-005 ($P = 0.04$) and unclassified genera from Family Lachnospiraceae ($P < 0.01$). In the rumen, a supplement × day interaction was detected for phyla Bacteroidota ($P = 0.01$), Firmicutes ($P = 0.02$), and Spirochaetota ($P = 0.04$), and tendencies for Actinobacteria ($P = 0.10$) and Chloroflexi ($P = 0.09$). Ruminal genera Prevotella, UCG-004, and unclassified genera from Family Lachnospiraceae ($P \leq 0.05$) were affected by the interaction. Neither alpha diversity metrics nor relative abundance of genera showed significant effects for the interaction of protein supplementation level × pubertal status in the rumen or uterus. The only significant interaction detected was the phyla Actinobacteria in the rumen ($P = 0.03$).

### DISCUSSION

Ruminal and uterine bacterial communities have been established as playing important roles in feed efficiency and breeding outcome success, respectively (Myer et al., 2015; Ault et al., 2019a). Therefore, bacterial communities of these environments may be affected during heifer development and potentially influence the long-term productivity of the heifer. Physiological changes associated with attainment of puberty, and nutritional supplementation commonly applied to support heifer growth and development, may influence the composition and effects of these microbial communities. The first objective of the current study was to evaluate the impact of differing levels of supplemented crude protein throughout heifer development on ruminal and uterine bacterial communities. Ruminal bacterial communities are well known to shift with changes in dietary feedstuffs or nutrient...

![FIGURE 5](image-url) | Beta diversity by principal coordinate analysis with Bray Curtis distances for day of supplementation in the uterus (A; $R^2 = 0.10$, $P = 0.001$) and rumen (B; $R^2 = 0.08$, $P = 0.001$). Significance was determined by PERMANOVA with 999 permutations. Ellipses represent 95% confidence interval.
composition (McCann et al., 2014; Clemmons et al., 2019a; Gruninger et al., 2019). As expected, we observed that ruminal bacterial communities were affected as a result of differences in protein supplementation treatment consistent with previous studies (Table 2). *Prevotella*, the most abundant genus detected in the rumen, and its respective phylum Bacteroidota were observed to have greater abundances as supplement crude protein level increased. *Prevotella* spp. are commonly the dominant genus in the rumen, although abundances may differ by the type of diet such as a greater concentrate compared to fiber diet (Stevenson and Weimer, 2007; Pitta et al., 2014). Additionally, abundance of *Prevotella* has been greater in heifers than mature cows (Liu et al., 2017) and different between cattle of differing feed efficiencies (Carberry et al., 2012; Myer et al., 2015). *Prevotella* spp. are functionally diverse but are well established as major proteolytic bacterial species in the rumen and capable of proteolysis via a variety of mechanisms (Wallace, 1996; Griswold et al., 1999). Multiple studies have observed a greater abundance of *Prevotella* spp. in diets with greater crude protein content (Belanche et al., 2012; Pitta et al., 2014), as observed in the current study. Due to the high proteolytic activity of *Prevotella*, increases in its abundance potentially impact nitrogen utilization and amino acids reaching the lower gastrointestinal tract for absorption and use by the host (Bach et al., 2005). However, additional data from the current study did not observe changes in uterine pH based on protein supplementation treatment, and only two amino acid concentrations in the uterine luminal fluid were impacted (Brandt, 2020).

Potential impacts of nutrition on uterine bacterial communities in bovine have not been previously studied. The current study applied a common base diet across all heifer groups with only crude protein level of the supplement modified. With the protein supplement, overall diet crude proteins were 11%, 15%, and 19% among groups. Regardless, the relative abundance of multiple bacterial phyla and genera was affected in the uterus by the level of crude protein supplement (Table 2). Firmicutes was the most abundant bacterial phylum detected in the uterus, but *Prevotella* was

### Table 4: Bacterial phyla in the uterus and rumen impacted by day of supplementation.

| Environment | Phyla              | Day of supplementation | \( P \) |
|-------------|--------------------|------------------------|--------|
| Uterus      | Actinobacteriota   | 3.27 ± 0.35 a          | 2.37 ± 0.26 b | 1.38 ± 0.33 c | 0.004 |
|             | Chloroflexi        | 0.12 ± 0.01          ab | 0.13 ± 0.01 a | 0.09 ± 0.01 b | 0.04 |
|             | Fibrobacterota     | 1.43 ± 0.18          | 1.85 ± 0.14 | 1.49 ± 0.17 | 0.07 |
|             | Firmicutes         | 47.07 ± 3.61         | 45.81 ± 2.82 | 54.11 ± 0.03 | 0.09 |
|             | Patescibacteria    | 1.04 ± 0.13          b | 1.74 ± 0.10 a | 1.23 ± 0.12 b | <0.0001 |
|             | Actinobacteriota   | 0.05 ± 0.01          b | 0.10 ± 0.01 a | 0.10 ± 0.01 a | <0.0001 |
|             | Bacteroidota       | 72.51 ± 1.18         a | 61.22 ± 0.88 b | 63.52 ± 1.10 b | <0.0001 |
|             | Chloroflexi        | 0.16 ± 0.01          b | 0.08 ± 0.01 a | 0.09 ± 0.01 b | 0.01 |
|             | Elusimicrobiota    | 0.10 ± 0.01          b | 0.15 ± 0.01 a | 0.09 ± 0.01 b | 0.0004 |
|             | Firmicutes         | 21.09 ± 1.15         b | 30.07 ± 0.86 a | 29.92 ± 1.07 a | <0.0001 |
|             | Patescibacteria    | 0.57 ± 0.11          c | 1.92 ± 0.09 a | 1.26 ± 0.10 b | <0.0001 |
|             | Proteobacteria     | 1.95 ± 0.19          a | 2.10 ± 0.15 a | 0.81 ± 0.17 b | <0.0001 |
|             | Spirochaetota      | 0.46 ± 0.03          ab | 0.50 ± 0.03 a | 0.41 ± 0.03 b | 0.05 |
|             | Verrucomicrobiota  | 0.98 ± 0.10          | 1.17 ± 0.08 | 0.94 ± 0.10 | 0.06 |

1 Relative abundances presented as Mean (%) ± SEM.
2 ab means within row different at indicated P.

### Table 5: Top 5 most abundant bacterial genera in the uterus and rumen impacted by day of supplementation.

| Environment | Genera                          | Day of supplementation | \( P \) |
|-------------|---------------------------------|------------------------|--------|
| Uterus      | Family F082                     | 4.89 ± 0.55 h          | 6.44 ± 0.43 a | 4.18 ± 0.52 b | 0.001 |
|             | Family Lachnospiraceae          | 3.78 ± 0.56 b          | 2.48 ± 0.42 c | 6.27 ± 0.52 | <0.0001 |
|             | Family UC93010                  | 2.97 ± 0.61 a          | 1.68 ± 0.46 b | 4.35 ± 0.56 a | 0.0009 |
|             | UCG-005                        | 4.26 ± 0.83 h          | 0.83 ± 0.84 h | 6.55 ± 0.78 | <0.0001 |
|             | Unspalasma                      | 6.48 ± 4.81           | 14.00 ± 5.63 | 6.86 ± 4.45 | 0.07 |
| Rumen       | Christensenellaceae group      | 3.30 ± 0.24 b          | 4.26 ± 0.19 a | 4.19 ± 0.23 a | 0.004 |
|             | Family Bacteroidales group      | 6.31 ± 0.72 a          | 5.25 ± 0.55 a | 2.58 ± 0.67 b | 0.004 |
|             | Family F082                     | 5.91 ± 0.57 b          | 6.85 ± 0.47 a | 5.40 ± 0.54 b | 0.04 |
|             | Prevotella                      | 36.45 ± 1.75 a         | 22.63 ± 1.33 c | 30.27 ± 1.65 b | <0.0001 |
|             | Rikenellaceae RC9 gut group     | 6.84 ± 0.50 h          | 10.55 ± 0.38 a | 8.38 ± 0.46 b | <0.0001 |

1 Relative abundances presented as mean (%) ± SEM.
2 ab means within row different at indicated P.
the most abundant genus and increased in abundance with increasing supplemented crude protein, similar to ruminal \textit{Prevotella}. The relative abundance of \textit{Prevotella} spp. in the uterus of healthy cows has been previously observed to range between <2% and 12% on average (Clemmons et al., 2017; Ault et al., 2019a). Interestingly, increased \textit{Prevotella} spp. in the uterus is typically associated with uterine disease in postpartum cows (Sheldon et al., 2009; Galvão et al., 2019) and was one of eight bacteria that differed in abundance between successful (<1% abundance) and failed (>1% abundance) pregnancies in apparently healthy postpartum cows (Ault et al., 2019a), indicating potential to influence breeding success. However, limited research exists on the uterine microbiome of heifers between weaning and breeding. The presence of \textit{Prevotella}, and other uterine disease-associated bacteria, have been found on endometrial biopsies from the uterus of virgin, pubertal heifers (Moore et al., 2017). The current study also found common uterine pathogen \textit{Ureaplasma} to also be highly abundant in the uterus of heifers, but relative abundance was not affected by protein supplementation level and did not shift over time. Other bacteria not associated as uterine pathogens were also affected by protein supplementation, such as \textit{Alistipes} and \textit{Bacteroides}, with the greatest abundance detected in control and P40 groups. The abundance of these bacteria was detected in >1% relative abundance in uterine samples but not rumen samples. \textit{Alistipes} and \textit{Bacteroides} have been previously associated with greater abundances in the lower gastrointestinal tract and feces of ruminants (Holman and Gzyl, 2019). Vaginal bacterial communities commonly share genera with gastrointestinal and fecal bacteria, which can enter into the uterus (Clemmons et al., 2017), or possibly enter the uterine environment through blood (Jeon et al., 2017). \textit{Alistipes} and \textit{Bacteroides} have been recognized as core vaginal bacteria (Rodrigues et al., 2015), and proposed as potential predictors of reproductive success based on abundance in fecal samples (McClure, 2018). Further research is needed to understand the role of potentially pathogenic bacteria from gastrointestinal sources, and their effects on the uterine environment and endometrial tissue. Despite the minimal differences in overall diet crude protein, differences in uterine bacterial communities by protein supplementation treatments were detected. Greater differences in diet such as different forage to concentrate ratios, energy supply, or other overall nutrient levels may have a larger impact on additional bacteria in the uterus. Therefore, the ability of protein supplementation to affect the abundances of bacteria in the uterus may allow for manipulation of the microbiome toward an optimal uterine environment.

The attainment of puberty is a significant milestone in heifer development. The activity of endometrial tissues and the uterine environment changes with the onset of puberty including tissue gene expression, hormonal signal response, and uterine histotrophic secretions (Cánovas et al., 2014; Fortes et al., 2018). Therefore, an additional objective of the current study was to evaluate the impact of pubertal status on bacterial communities, beginning with prepubertal heifers and ending 2 weeks following the first breeding. Alpha diversity metrics indicated differences in the bacterial community of the uterus during pubertal attainment, as observed ASVs increased over time and were greater in pubertal than prepubertal heifers (Table 1). In a study of the vaginal microbiome between cows and heifers, cows had a greater number of observed bacterial species than heifers (Laguardia-Nascimento et al., 2015). Pubertal heifers also had increased similarity of their overall uterine microbiome than prepubertal heifers according to beta diversity analyses of the current study (Figure 4A). When evaluating specific bacterial taxa in the uterus of our study, five phyla, including the two most abundant of Firmicutes and Bacteroidota, were affected by pubertal status (Table 3). Firmicutes was greater in prepubertal than pubertal heifers, but the remaining four were observed to have greater relative abundances in pubertal heifers compared to prepubertal heifers. Similar results were observed previously in cows during estrus synchronization leading up to breeding (Ault et al., 2019a) where Firmicutes was highly abundant (61%–74% abundance) and dominated the uterine bacterial communities at the beginning of estrous synchronization during low P4 concentrations. Firmicutes then decreased to 36% on average, with other phyla significantly increasing, 2 days before breeding during higher P4 concentrations (Ault et al., 2019a). Luteal function and resulting P4 concentrations are the defining parameter of pubertal animals, indicative of their ability to successfully ovulate and establish and maintain a pregnancy (Kinder et al., 1995; Atkins et al., 2013; Fortes et al., 2018). Progesterone influences uterine function through changes in endometrial gene expression related to activities such as immune response, nutrient exchange to the lumen, and other processes related to the establishment of pregnancy (Forde et al., 2010; Forde et al., 2014; Fortes et al., 2018). Companion data from the current study reported by Brandt (2020) found that 18 amino acids, regardless of protein supplementation treatment, reached their highest concentration in the uterus at d 163 indicating changes to the uterine environment through development to puberty attainment. These changes occurring with pubertal onset from P4 stimulation to the uterine environment and endometrial tissue may influence the bacterial communities of the uterus, leading to a more similar overall uterine microbiome of pubertal animals.

Furthermore, there was no difference in observed ASVs and fewer bacterial phyla and genera were affected by pubertal status in the rumen in contrast to the uterus (Tables 1, 3). Steroid hormone receptors associated with reproduction, such as estrogen and androgen receptors, have been reportedly expressed in the rumen, but P4 nuclear receptor expression has not been detected (Pfaill et al., 2003). Therefore, minimal direct impact on the bacterial communities is expected in the rumen due to the changes specifically associated with puberty and increased P4 secretion. Temporal shifts in bacterial diversity and relative abundances of taxa, however, were observed in the current study of both the rumen and uterus (Tables 4, 5). Alpha diversity analyses indicated changes in the number of ASVs, and beta diversity indicated a slight shift in the rumen overall rumen bacterial community composition over time (Table 1). Ruminal
bacterial communities have been shown to undergo shifts according to a change in diet (McCann et al., 2014; Clemmons et al., 2019a; Gruninger et al., 2019). However, previous research has indicated stability of the rumen microbiome by 9 weeks following a diet change (Clemmons et al., 2019b). Sampling of the rumen bacterial communities in the current study did not occur until 56 days following the beginning of supplementation. Therefore, the rumen microbiome had likely reached any diet-induced equilibrium, following the addition of supplements to the heifer’s diet, prior to sampling. The development of the rumen microbiome from birth to adulthood is not an immediate process, with the number of species shown to increase from birth up to 2 years of age (Guo et al., 2020), along with shifts in bacterial taxa composition (Jami et al., 2013). The first few days of life and weaning are likely the two most influential periods on rumen microbiome development (Guo et al., 2020; Amin and Seifert, 2021). Therefore, the rumen microbiome of heifers in the current study was likely continuing to develop following weaning into its mature state.

In conclusion, weaning through breeding during heifer development is a crucial period to evaluate the uterine and ruminal bacterial communities and factors that may influence their composition, such as nutrition and puberty attainment. Despite relatively minor changes in overall diet crude protein, the current study was able to detect differences in the uterine and rumen bacterial communities with varying levels of protein supplementation. Pubertal status had a greater impact on uterine bacterial communities than in the rumen. The effects observed over time and by pubertal status on bacterial communities of the heifer uterus potentially indicates progression of a maturing uterine microbiome through development. Future studies are needed to understand how nutrition and the overall diet may be used to manipulate uterine bacterial communities, and how the changing uterine microbiome through development may impact the uterine environment. Together, such studies would contribute to a greater understanding of the developing uterine bacterial communities and how they might be modified to improve future reproductive success and lifetime reproductive efficiency.

DATA AVAILABILITY STATEMENT

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found below: https://www.ncbi.nlm.nih.gov/, PRJNA819438.

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ETHICS STATEMENT

The animal study was reviewed and approved by Institutional Animal Care and Use Committee at the University of Tennessee, Knoxville.

AUTHOR CONTRIBUTIONS

TA-S was responsible for primary manuscript preparation, sampling, DNA extraction, and data analyses. KB assisted with sample collections and manuscript preparation. MH assisted with data analyses and manuscript preparation. RP assisted with laboratory analyses and manuscript preparation. TS was responsible for DNA library preparation and sequencing. DM, SM, FS, and KP were responsible for experimental design development and manuscript preparation. JR, LS, KM, and PM were responsible for sampling, experimental design development, and manuscript preparation. All authors contributed to the article and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fanim.2022.903909/full#supplementary-material

Ault, T. B., Clemmons, B. A., Reese, S. T., Dantas, F. G., Franco, G. A., Smith, T. P. L., et al. (2019a). Bacterial Taxonomic Composition of the Postpartum Cow Uterus and Vagina Prior to Artificial Insemination. J. Anim. Sci. 97 (10), 4305–4313. doi: 10.1093/jas/skz212

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