



Changes in the predicted function of the rumen bacterial community of Japanese Black beef cattle during the fattening stages according to Kyoto Encyclopedia of Genes and Genomes (KEGG) analyses

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ABSTRACT. We investigated changes in the predicted functions of the rumen bacterial community in Japanese Black beef cattle during fattening. Nine cattle were fed a high-concentrate diet during the early, middle, and late fattening stages consecutively (10–14, 15–22, and 23–30 months of age, respectively). The rumen fluid and solid samples collected at each stage were subjected to sequencing analyses. The sequencing results were clustered and classified into operational taxonomic units (OTUs). Representative sequences and a raw counting table for each OTU were submitted to the Piphillin website. The predicted functions were revealed by the Kyoto Encyclopedia of Genes and Genomes database as the ratio of the total sequence. In the early stage, “Biosynthesis of secondary metabolites” was significantly higher in the fluid fraction than in the solid fraction. “Two-component system” in the middle stage was significantly lower and “Purine metabolism” in the late stage was significantly higher in the fluid fraction than those in the solid fraction. The fluid fraction was significantly correlated with acetic acid, propionic acid, and bacterial metabolism, such as “Biosynthesis of secondary metabolites” and “Sugar metabolism.” Moreover, the solid fraction was correlated with “Purine metabolism” and “Biosynthesis of secondary metabolites.” These results suggest that the rumen bacterial community in Japanese Black beef cattle adapts to changes in rumen conditions by altering their functions in response to a long-term high-grain diet.

KEY WORDS: bacterial function, Japanese Black beef cattle, Kyoto Encyclopedia of Genes and Genomes (KEGG) analysis, rumen fluid and solid bacteria

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Japanese Black cattle are characterized by the ability to deposit a large amount of intramuscular fat and typically consume a large amount of concentrated diet from the age of 10–30 months [22, 23]. This, in addition to the lack of roughage, can lead to excessive generation and accumulation of volatile fatty acids (VFAs), decreasing ruminal pH, such as subacute ruminal acidosis (SARA) [10, 18, 22, 23]. Long-term high-grain diet feeding does not acutely alter bacterial diversity but causes bacteria to adapt and remain stable against the decrease in pH and changes in fermentation to enhance lactic acid production in Japanese Black beef cattle [23]. Furthermore, acute hepatitis related to SARA is a major problem in beef farms [18]. Meanwhile, the ruminal bacterial community and ruminal pH can adapt to and influence each other [19], and the effects of short- (days) and mid-term (weeks) SARA and RA challenges have been explored previously [12, 19, 30]. In general, grain-based SARA challenge induces death or lysis of gram-negative bacteria, such as Bacteroidetes and Proteobacteria, and eventually the proportion of Firmicutes increases with severely low ruminal pH [12, 19].

Large-scale sequencing has been used to detect large amounts of genomic DNA or RNA sequences, and the number of gene expression studies has increased. Ruminal pH, VFAs, and bacterial community structure change dramatically during the fattening

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stage in Japanese Black beef cattle [23]. Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway analyses enable the prediction of the meaning of screened DNA or RNA sequences from large-scale sequencing results. A previous KEGG pathway analysis revealed that live yeast contributes to the rumen microbiota of beef cattle [24]. Furthermore, the rumen microbiome, as a reservoir of antimicrobial resistance, is directly affected by diet [1] and by dietary energy sources and levels that shift the multi-kingdom microbiota and functions in the rumen of dairy cows [26]. However, no study to date has conducted predicted functional analyses of the rumen bacterial community in Japanese Black beef cattle during the fattening stages.

MATERIALS AND METHODS

We re-analyzed our previously published data regarding rumen fermentation parameters and sequencing data [22] to evaluate the relationship between rumen measurements and the predicted functions of the bacterial community in Japanese Black beef cattle. All experimental procedures were approved by the Iwate University Laboratory Animal Care and Use Committee (A201720) and the Hyogo Prefectural Technology Center for Agriculture, Forestry, and Fisheries.

Animals, experimental design, sampling, and measurements

Experimental animals, design, and measurements have been previously described [23]. Briefly, Japanese Black beef cattle ($n=9$) were castrated and equipped with a rumen cannula. The fattening period was subdivided into early, middle, and late stages (10–14, 15–22, and 23–30 months of age, respectively) according to a general agreement in Japan [25]. The cattle were fed concentrate and forage (rice straw) diets during all three stages, and the amount of the concentrate diet was increased gradually throughout the experimental period. The forage diet was supplied daily at 0930 and 1530 hr in two equal portions, and the concentrate was supplied 1-hr after rice straw. A calculated amount of roughage and the concentrate diet was given for a daily gain of 0.8 kg/day during all stages. The forage-to-concentrate ratios were 26:74, 13:87, and 14:86 during the early, middle, and late stages, respectively. The adequacy rate of the diet was calculated based on the nutrient requirements of the Japanese feeding standard for beef cattle [21].

Samples were collected and measurements were made as previously described [23]. Briefly, ruminal pH was measured continuously every 10 min during the final 7 days of the early, middle, and late fattening stages using a radio transmission system (YCOW-S; DKK-TOA, Yamagata, Japan), as previously described [31]. Rumen samples were collected through the rumen cannula every fourth day during the final 7 days of the pH measurements at each stage to analyze the rumen bacterial community, VFAs, lactic acid concentration, and lipopolysaccharide (LPS) activity. The rumen samples were filtered through two layers of cheesecloth, the fluid was treated as the fluid fraction, and the remaining sample on the cheesecloth was collected as the solid fraction.

The ruminal concentrations of VFAs (i.e., acetic acid, propionic acid, and butyric acid) were quantified by gas chromatography (GC-2014; Shimadzu, Kyoto, Japan). The lactic acid concentration was determined using a commercially available kit (F-kit [d-lactate/l-lactate], J.K. International Co., Tokyo, Japan). Ruminal LPS activity was assayed using a kinetic *Limulus* ameocyte lysate assay (Pyrochrome with Glucashield; Seikagaku Corp., Tokyo, Japan).

DNA extraction and sequencing

DNA extraction and Illumina sequencing analyses were performed as previously described [22]. Briefly, total bacterial DNA was extracted from the fluid and solid portions, as described by Kim *et al.* [15]. Sequencing libraries were prepared according to the Illumina 16S Metagenomic Sequencing Library preparation guide (2013). The bacterial 16S rRNA gene was amplified using the barcoded universal primers 515F (5-TGYCAGCMGCCGCGGTAA-3) and 805R (5-GACTACHVGGGTATCTAATCC-3) spanning the V4 hypervariable region. The sequence data were deposited into the Sequence Read Archive (SRA) of the National Center for Biotechnology Information database and can be accessed via SRA accession number PRJNA548210 (<https://submit.ncbi.nlm.nih.gov/subs/sra/>).

Sequencing data analyses

We used the sequencing data presented by Ogata *et al.* [22], analyzed all sequencing results using the MOTHUR program (version 1.41.1; University of Michigan; <http://www.mothur.org/wiki/Schloss>) [32], and mapped sequence data with the SILVA reference database (SSURef release 128; [27]). Then, candidate sequences were screened and filtered, as follows: unique sequences were determined; candidate sequences were pre-clustered to eliminate outliers; chimeras were identified using the “chimera.vsearch” command; and sequence comparison was performed using the Mothur Ribosomal Database Project training set (version 16). Sequences were clustered and classified into operational taxonomic units (OTUs) using a cutoff value of 97% similarity. All samples were standardized by random subsampling (6,741 sequences/sample) using the “sub.sample” command.

Representative sequences for each OTU were set using the BLASTn program (<https://blast.ncbi.nlm.nih.gov/Blast.cgi>) and compared to the 16S ribosomal RNA sequence database (Bacteria and Archaea; May 2019). Furthermore, representative sequences and tabulated raw count data were submitted to the Piphillin website (<http://piphillin.secondgenome.com/>; [13]) and analyzed. A sequence identity cutoff of 97% was applied to analyze the functional categories, and metagenomic functions were assigned using the KEGG database (November 2019).

Statistical analyses

The normality of the data distribution was assessed using the Shapiro-Wilk test. Significant differences (between groups) in variables during the early, middle, and late stages were detected using Student’s *t*-test for normal variables and Welch’s *t*-test for

non-normal variables. Significant changes (within groups) in ruminal pH, VFAs, lactic acid concentration, and ruminal LPS activity during the early, middle, and late stages were assessed using the Friedman test, and the relative abundances of bacteria and their functions were assessed using the Kruskal-Wallis test. Correlation coefficients between the predicted function of rumen bacteria and rumen measurements were calculated. Pearson's correlation coefficient (r) and significance levels (P -values) were used to determine the relationships between rumen measurements and the predicted functions of the bacterial community. All numerical data were analyzed using Prism version 8.10 (GraphPad Software Inc., La Jolla, CA, USA). Statistical significance was set at $P < 0.05$.

RESULTS

pH and volatile fatty acids

The 24-hr mean pH, total VFAs, and acetic acid concentration decreased as the fattening stages proceeded. Propionic acid concentration increased during the middle stage. LPS activity increased throughout the study period (Table 1).

Bacterial abundance

The top 10 bacterial families comprised more than 80% of all sequences during each stage and in each fraction (fluid or solid) (Table 2). Four families (Lachnospiraceae, Ruminococcaceae, Coriobacteriaceae, and Prevotella) accounted for more than 60% of all bacteria in the fluid fraction throughout the fattening stages. Four of the five families (the above four families and Firmicutes-unclassified family) were the same in the solid fraction as in the fluid fraction, which included more than 60% of all bacteria during the early and middle stages, and six families (the above five families and the Acidaminococcaceae family) accounted for more than 60% of all sequences during the late stage.

Predicted functions of rumen bacteria

The predicted functions of the sequence data were revealed against the KEGG database (November 2019) as the ratio of the total sequences. Comparisons between the fluid and solid fractions during each stage are presented in Table 3. "Metabolic pathway" (KEGG pathway object identifier; ko01100) and "Biosynthesis of secondary metabolites" (ko01110) were significantly higher

Table 1. Twenty four-hr mean pH, total volatile fatty acid (VFA), individual VFA proportions, acetic acid-to-propionic acid (A/P) ratio, lactic acid concentrations, and lipopolysaccharide (LPS) activity in Japanese Black beef cattle ($n=9$) during the fattening stages

Item	Stage			P -value ^a
	Early	Middle	Late	
24-hr mean pH	6.22 ± 0.06	6.06 ± 0.08	5.73 ± 0.13	0.031
Total VFA (mmol/dl)	13.1 ± 0.59	12.3 ± 0.80	9.77 ± 0.76	0.057
Acetate (%)	62.6 ± 2.04	57.1 ± 1.53	58.6 ± 1.50	0.329
Propionate (%)	21.4 ± 1.81	27.1 ± 2.23	27.1 ± 2.21	0.187
Butyrate (%)	11.9 ± 1.04	12.7 ± 0.86	11.2 ± 0.85	0.154
A/P ratio	3.05 ± 0.22	2.24 ± 0.21	2.34 ± 0.28	0.154
Lactic acid (mg/l)	67.9 ± 0.01	25.0 ± 0.00	141.4 ± 0.01	<0.001
LPS (EU×10 ⁴ /ml)	1.34 ± 0.39	4.29 ± 1.82	6.62 ± 2.24	0.048

^aFriedman test was used to determine within-group differences.

Table 2. Relative abundance of major bacterial families during each fattening stage (early, middle, and late stages) in the fluid and solid fractions of Japanese Black beef cattle ($n=9$)

Item	Stage					
	Early		Middle		Late	
	Fluid	Solid	Fluid	Solid	Fluid	Solid
<i>Lachnospiraceae</i>	19.6 ± 0.53	21.3 ± 0.51	15.4 ± 0.57	22.8 ± 0.50	20.8 ± 0.57	23.7 ± 0.56
<i>Ruminococcaceae</i>	25.4 ± 0.56	10.8 ± 0.54 ^a	14.9 ± 0.60	6.55 ± 0.52	16.6 ± 0.59	8.74 ± 0.58
<i>Coriobacteriaceae</i>	6.68 ± 0.94	11.8 ± 0.93	9.41 ± 1.00	16.2 ± 0.90 ^a	13.8 ± 1.02	11.1 ± 1.01
<i>Prevotellaceae</i>	8.72 ± 0.84	4.49 ± 0.84	23.7 ± 0.93	10.4 ± 0.81	9.36 ± 0.91	4.33 ± 0.90
<i>Firmicutes_unclassified</i>	9.39 ± 0.52	18.0 ± 0.51 ^a	4.66 ± 0.56	7.76 ± 0.49	5.38 ± 0.56	10.5 ± 0.56
<i>Acidaminococcaceae</i>	3.46 ± 0.18	1.04 ± 0.17	4.26 ± 0.21	5.14 ± 0.16	5.27 ± 0.19	14.8 ± 0.19 ^a
<i>Clostridiales_unclassified</i>	4.10 ± 0.48	8.15 ± 0.47 ^a	2.55 ± 0.52	7.46 ± 0.45 ^a	2.33 ± 0.51	4.48 ± 0.51 ^a
<i>Clostridiales_Incertae_Sedis_XIII</i>	1.72 ± 0.48	7.95 ± 0.47 ^a	2.10 ± 0.52	7.64 ± 0.45 ^a	1.61 ± 0.51	3.90 ± 0.51 ^a
<i>Veillonellaceae</i>	1.50 ± 0.67	2.23 ± 0.66	2.14 ± 0.74	4.88 ± 0.65 ^a	2.04 ± 0.73	4.02 ± 0.72
<i>Bacteria_unclassified</i>	3.00 ± 1.21	2.51 ± 1.21	3.98 ± 1.30	1.97 ± 1.16	2.77 ± 1.30	1.97 ± 1.30

^aDenotes a significant difference ($P < 0.05$) between the solid and fluid fractions.

Table 3. Relative abundance of the bacterial predicted functions during each fattening stage (early, middle, and late stages) in the fluid and solid fractions of Japanese Black beef cattle (n=9)

Item	Stage					
	Early		Middle		Late	
	Fluid	Solid	Fluid	Solid	Fluid	Solid
Metabolic pathways	16.3 ± 0.15	15.7 ± 0.14 ^a	16.5 ± 0.22	16.0 ± 0.03	16.1 ± 0.11	16.0 ± 0.10
Biosynthesis of secondary metabolites	7.44 ± 0.06	7.12 ± 0.04 ^a	7.57 ± 0.07	7.32 ± 0.04	7.38 ± 0.06	7.31 ± 0.03
Biosynthesis of antibiotics	5.65 ± 0.03	5.61 ± 0.04	5.77 ± 0.04	5.77 ± 0.04	5.78 ± 0.04	5.94 ± 0.05
Biosynthesis of amino acids	4.20 ± 0.05	4.08 ± 0.07	4.34 ± 0.08	4.24 ± 0.06	4.29 ± 0.05	4.30 ± 0.05
Microbial metabolism in diverse environments	4.05 ± 0.05	4.29 ± 0.07 ^a	4.00 ± 0.04	4.19 ± 0.07 ^a	4.13 ± 0.04	4.18 ± 0.04
Aminoacyl-tRNA biosynthesis	2.85 ± 0.15	2.69 ± 0.13	3.10 ± 0.19	2.63 ± 0.13	2.65 ± 0.10	2.55 ± 0.13
ABC transporters	2.58 ± 0.18	3.07 ± 0.11	2.26 ± 0.29	2.86 ± 0.11	2.81 ± 0.14	2.73 ± 0.20
Ribosome	2.67 ± 0.05	2.53 ± 0.04	2.78 ± 0.06	2.53 ± 0.04 ^a	2.62 ± 0.03	2.53 ± 0.02 ^a
Carbon metabolism	2.33 ± 0.01	2.35 ± 0.02	2.31 ± 0.02	2.34 ± 0.02	2.33 ± 0.01	2.36 ± 0.01
Purine metabolism	2.07 ± 0.02	2.04 ± 0.02	2.07 ± 0.03	1.99 ± 0.02	2.03 ± 0.02	2.00 ± 0.01 ^a
Pyrimidine metabolism	1.72 ± 0.01	1.70 ± 0.02	1.69 ± 0.02	1.64 ± 0.02	1.68 ± 0.02	1.69 ± 0.02
Quorum sensing	1.51 ± 0.10	1.70 ± 0.02	1.55 ± 0.16	1.62 ± 0.02	1.55 ± 0.06	1.60 ± 0.07
Amino sugar and nucleotide sugar metabolism	1.55 ± 0.04	1.61 ± 0.03	1.36 ± 0.06	1.44 ± 0.03	1.46 ± 0.04	1.43 ± 0.04
Starch and sucrose metabolism	1.34 ± 0.04	1.41 ± 0.04	1.27 ± 0.06	1.39 ± 0.04	1.42 ± 0.04	1.46 ± 0.06
Fructose and mannose metabolism	1.02 ± 0.07	1.27 ± 0.08 ^a	0.83 ± 0.09	1.10 ± 0.08	1.07 ± 0.07	1.11 ± 0.06
Two-component system	1.05 ± 0.08	1.14 ± 0.11	0.90 ± 0.03	1.07 ± 0.10 ^a	0.97 ± 0.06	0.96 ± 0.02
Glycolysis/Gluconeogenesis	1.04 ± 0.02	1.07 ± 0.03	0.99 ± 0.03	1.02 ± 0.03	1.03 ± 0.02	1.04 ± 0.01
Phosphotransferase system (PTS)	0.85 ± 0.17	1.44 ± 0.18	0.64 ± 0.17	1.16 ± 0.18	1.09 ± 0.16	1.14 ± 0.15
Homologous recombination	0.94 ± 0.03	0.94 ± 0.04	0.99 ± 0.04	1.02 ± 0.04	1.04 ± 0.03	1.08 ± 0.04
Pyruvate metabolism	0.92 ± 0.03	1.00 ± 0.02	0.88 ± 0.05	1.01 ± 0.02	1.01 ± 0.02	1.07 ± 0.03

^aDenotes a significant difference ($P < 0.05$) between the solid and fluid fractions.

($P < 0.05$) in the fluid fraction than in the solid fraction during the early stage. “Microbial metabolism in diverse environments” (ko01120) was significantly higher ($P < 0.05$) in the solid fraction than in the fluid fraction. “Ribosome” (ko03010) was significantly higher ($P < 0.05$) in the fluid fraction than in the solid fraction during the middle stage, and “Microbial metabolism in diverse environments” (ko01120) and “Two-component system” (ko02020) were higher in the solid fraction than in the fluid fraction. “Ribosome” (ko03010) and “Purine metabolism” (ko00230) were significantly higher ($P < 0.05$) in the fluid fraction than in the solid fraction during the late stage.

Comparisons of the fluid and solid fractions among the fattening stages are shown in Table 4. In the solid fraction during the late fattening stage, “Biosynthesis of secondary metabolites” (ko01110) and “Biosynthesis of antibiotics” (ko01130) significantly increased ($P < 0.05$), and “Amino sugar and nucleotide sugar metabolism” (ko00520) significantly decreased ($P < 0.05$).

Predicted functional analyses of the Lachnospiraceae and Ruminococcaceae families

The predicted functions of the Lachnospiraceae family are shown in Fig. 1. The Lachnospiraceae family was present in the highest proportion in the solid fraction than in the fluid fraction during the early stage. “ABC transporters” (ko02010), “Two-component system” (ko02020), and “Quorum sensing” (ko02024) were significantly higher in the fluid fraction than in the solid fraction during the early stage ($P < 0.05$). In contrast, “Biosynthesis of amino acids” (ko01230) was significantly higher in the solid fraction than in the fluid fraction ($P < 0.05$). No significant differences were detected between the middle and late stages.

The predicted functions of the Ruminococcaceae family are shown in Fig. 2. The Ruminococcaceae family was more abundant in the solid fraction than in the fluid fraction during the early stage. “Microbial metabolism in diverse environments” (ko01120) was significantly higher in the fluid fraction than the solid fraction during the late stage ($P < 0.05$). No significant differences were observed between the early and middle stages.

Correlation coefficients between the predicted functions of the bacterial community and ruminal LPS activity and the proportion of acetic and propionic acids

Table 5 summarizes the significant correlation coefficients between the predicted functions of the bacterial community and ruminal LPS activity and the proportions of acetic and propionic acids. As a result, 9, 10, and 15 KEGG pathways related to LPS, acetic acid, and propionic acid, respectively, were identified in the fluid fraction (Table 5). Significant and strong correlations were observed in pathways related to LPS, including “Metabolism”, “Environmental Information Processing”, and “Cellular Processes”. Furthermore, among the pathways showing strong correlations with acetic acid, six pathways belonged to “Metabolism” or “Environmental Information Processing” and one belonged to “Genetic Information Processing”. The pathways strongly correlated with propionic acid and acetic acid mostly overlapped. Two propionic acid pathways belonged to “Metabolism” and one belonged to “Cellular Processes.”

Table 4. Relative abundance of the bacterial predicted functions in each fraction (fluid and solid fractions) of Japanese Black beef cattle (n=9) during the fattening stages

Items	Fluid			P-value ^a	Solid			P-value
	Early	Middle	Late		Early	Middle	Late	
Metabolic pathways	16.3 ± 0.15	16.5 ± 0.22	16.1 ± 0.11	0.382	15.7 ± 0.14	16.0 ± 0.03	16.0 ± 0.10	0.362
Biosynthesis of secondary metabolites	7.44 ± 0.06	7.57 ± 0.07	7.38 ± 0.06	0.290	7.12 ± 0.04	7.32 ± 0.04	7.31 ± 0.03	0.049
Biosynthesis of antibiotics	5.65 ± 0.03	5.77 ± 0.04	5.78 ± 0.04	0.154	5.61 ± 0.04	5.77 ± 0.04	5.94 ± 0.05	0.007
Biosynthesis of amino acids	4.20 ± 0.05	4.34 ± 0.08	4.29 ± 0.05	0.884	4.08 ± 0.07	4.24 ± 0.06	4.30 ± 0.05	0.213
Microbial metabolism in diverse environments	4.05 ± 0.05	4.00 ± 0.04	4.13 ± 0.04	0.220	4.29 ± 0.07	4.19 ± 0.07	4.18 ± 0.04	0.629
Aminoacyl-tRNA biosynthesis	2.85 ± 0.15	3.10 ± 0.19	2.65 ± 0.10	0.359	2.69 ± 0.13	2.63 ± 0.13	2.55 ± 0.13	0.711
ABC transporters	2.58 ± 0.18	2.26 ± 0.29	2.81 ± 0.14	0.299	3.07 ± 0.11	2.86 ± 0.11	2.73 ± 0.20	0.715
Ribosome	2.67 ± 0.05	2.78 ± 0.06	2.62 ± 0.03	0.469	2.53 ± 0.04	2.53 ± 0.04	2.53 ± 0.02	0.801
Carbon metabolism	2.33 ± 0.01	2.31 ± 0.02	2.33 ± 0.01	0.333	2.35 ± 0.02	2.34 ± 0.02	2.36 ± 0.01	0.717
Purine metabolism	2.07 ± 0.02	2.07 ± 0.03	2.03 ± 0.02	0.348	2.04 ± 0.02	1.99 ± 0.02	2.00 ± 0.01	0.071
Pyrimidine metabolism	1.72 ± 0.01	1.69 ± 0.02	1.68 ± 0.02	0.260	1.70 ± 0.02	1.64 ± 0.02	1.69 ± 0.02	0.211
Quorum sensing	1.51 ± 0.10	1.55 ± 0.16	1.55 ± 0.06	0.854	1.70 ± 0.02	1.62 ± 0.02	1.60 ± 0.07	0.778
Amino sugar and nucleotide sugar metabolism	1.55 ± 0.04	1.36 ± 0.06	1.46 ± 0.04	0.148	1.61 ± 0.03	1.44 ± 0.03	1.43 ± 0.04	0.003
Starch and sucrose metabolism	1.34 ± 0.04	1.27 ± 0.06	1.42 ± 0.04	0.308	1.41 ± 0.04	1.39 ± 0.04	1.46 ± 0.06	0.919
Fructose and mannose metabolism	1.02 ± 0.07	0.83 ± 0.09	1.07 ± 0.07	0.470	1.27 ± 0.08	1.10 ± 0.08	1.11 ± 0.06	0.326
Two-component system	1.05 ± 0.08	0.90 ± 0.03	0.97 ± 0.06	0.389	1.14 ± 0.11	1.07 ± 0.10	0.96 ± 0.02	0.206
Glycolysis/Gluconeogenesis	1.04 ± 0.02	0.99 ± 0.03	1.03 ± 0.02	0.841	1.07 ± 0.03	1.02 ± 0.03	1.04 ± 0.01	0.330
Phosphotransferase system (PTS)	0.85 ± 0.17	0.64 ± 0.17	1.09 ± 0.16	0.463	1.44 ± 0.18	1.16 ± 0.18	1.14 ± 0.15	0.422
Homologous recombination	0.94 ± 0.03	0.99 ± 0.04	1.04 ± 0.03	0.333	0.94 ± 0.04	1.02 ± 0.04	1.08 ± 0.04	0.292
Pyruvate metabolism	0.92 ± 0.03	0.88 ± 0.05	1.01 ± 0.02	0.126	1.00 ± 0.02	1.01 ± 0.02	1.07 ± 0.03	0.418

^aKruskal-Wallis test was used to determine within-group differences.

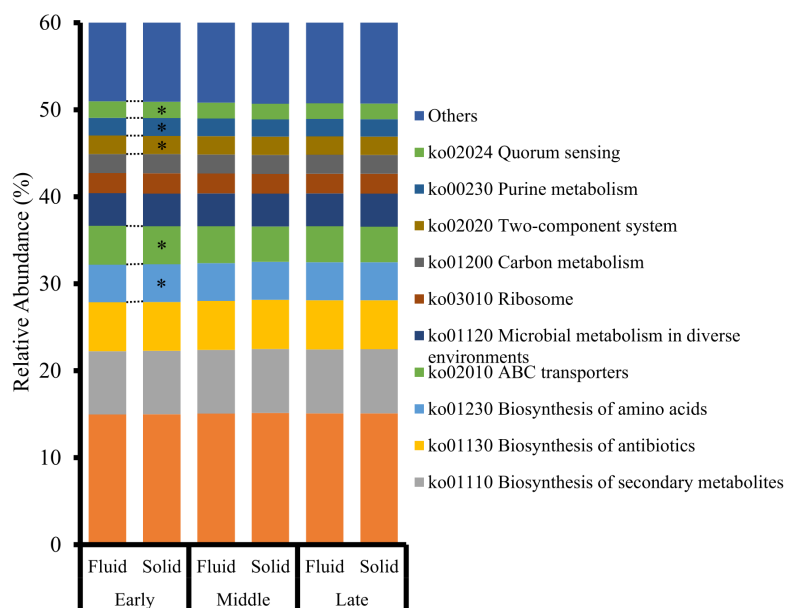


Fig. 1. Relative abundance of Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways inferred from the Lachnospiraceae family in the rumen fluid and solid fractions of Japanese Black beef cattle (n=9) during the fattening stages.

*Indicates a significant difference ($P < 0.05$) between the fluid and solid fractions.

Two, six, and six KEGG pathways ($P < 0.05$, correlation coefficients > 0.5) were related to LPS, acetic acid, and propionic acid, respectively, in the solid fraction (Table 5). Among the pathways related to acetic acid, three belonged to “Metabolism”, and one belonged to “Cellular Processes”. “C5-Branched dibasic acid metabolism” (ko00660) overlapped with both acetic and propionic acid. The pathways strongly related to propionic acid included four that belonged to “Metabolism” and one that belonged to “Secondary metabolites”.

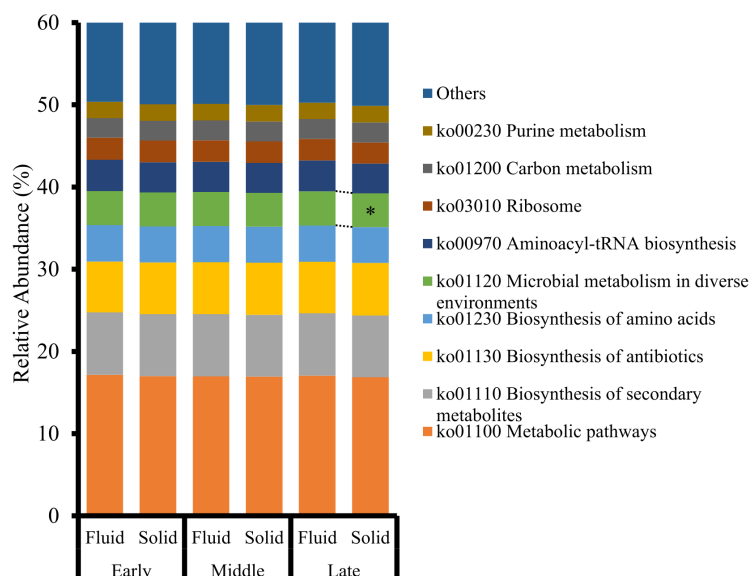


Fig. 2. Relative abundance of Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways inferred from the Ruminococcaceae family in the rumen fluid and solid fractions of Japanese Black beef cattle (n=9) during the fattening stages. *Indicates a significant difference ($P<0.05$) between the fluid and solid fractions.

DISCUSSION

Bacterial adhesions for feeding particles are characterized by a complex and diverse population [16], and the free-living bacterial species in the fluid fraction do not contribute much to metabolic activity. Instead, species in the fluid fraction contribute to the development of solid-adherent biofilms to newly ingested feed [17]. Therefore, bacterial communities in the fluid and solid fractions continuously interact with each other [6], but with potentially different functions [30]. Hackmann *et al.* [10] discovered that the recognized pathways for metabolizing pentose and hexose sugars into short-chain fatty acids do not adequately explain the fermentation products generated by a wide range of rumen bacteria. The Piphillin algorithm used in the present study for bacterial community analyses strongly relies on the amount and quality of reference genomes and can be used to obtain accurate results in animals [1, 24, 26].

The predicted functions of the bacterial community changed during the fattening stages following the altered rumen fermentation measurements described above. Several cellulolytic bacteria were found in the solid fraction during the early stage, and the higher proportion of free-living species in the fluid plays a role in metabolizing the end products of carbohydrate fermentation [6]. For example, secondary metabolites are synthesized via multistep pathways, and *Fusarium avenaceum*, a widespread pathogen of important crops, can produce many bioactive secondary metabolites using different carbon sources [35]. In addition, bacterial secondary metabolites often contain carbohydrate attachments that play a significant role in conferring biological activity [37]. However, the investigation of “Biosynthesis of secondary metabolites” in the rumen bacterial community is still limited. In contrast, the most abundant family in the fluid fraction during the middle fattening stage, Prevotellaceae (accounting for 24%), was engaged in various types of glycation and proteolysis [2]; thus, “Two-component systems” may be responsible for the signaling mechanism to respond to diverse environmental changes [4]. Furthermore, “Purine metabolism”, which is a metabolic pathway of the purine bases in nucleic acids, was identified as a significantly enriched pathway during the late stage. This pathway has been reported in carnivore fecal gut bacteria that predominantly have purine metabolic function [42] and in lactic acid-producing species of goats (*Lactococcus lactis*) [14] and humans (*Lactobacillus gaasseri PA-3*) [40], which might be consistent with the increase in lactic acid concentration during the late stage (Table 1) in the present study. The solid fraction bacteria also changed to digest cellulose materials during the early and middle stages and to respond to ruminal environment changes during the late stage. These results suggest that adaptation by the bacteria was identified simultaneously with rumen measurements, such as pH, VFA, and lactic acid, as well as the predicted functions of the bacterial community.

When fermentable fibers are lacking, microorganisms switch their energy metabolism to less energetically desirable sources such as amino acids, endogenous proteins, or dietary fats [5, 39]. As a result, fermentation activity decreases and VFAs as end products decrease [29]. Bacterial amino acid fermentation compensates for the decrease in VFAs [34]. Purine metabolism significantly increased in the fluid fraction during the late fattening stage, as shown in Table 3, when VFA concentrations tended to be low. Therefore, the significantly lower proportion of “Purine metabolism” in the solid fraction than in the fluid fraction suggested that fermentation did not proceed at a high rate during the late fattening stage.

During the fattening stages, a significant change in the predicted bacterial function was identified in the solid fraction. The glycosylated secondary metabolites in bacteria continue to serve as an important source for antibiotic discovery [37], and

Table 5. Pearson correlation coefficient analysis between inferred Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways in fluid and solid fractions and rumen lipopolysaccharide (LPS) activity and proportions of acetic and propionic acids (%) in Japanese Black beef cattle (n=9)

KEGG pathways	KEGG identifier	Pearson's correlation coefficient ¹		
		LPS	Acetic Acid	Propionic Acid
Fluid fraction				
AGE-RAGE signaling pathway in diabetic complications	ko00061	0.737		
ECM-receptor interaction	ko00524	0.738		
Focal adhesion	ko00950	0.738		
Isoflavonoid biosynthesis	ko02060	0.745		
Melanogenesis	ko00510	0.754		
Monoterpenoid biosynthesis	ko04964	0.733		
Platelet activation	ko00791	0.737		
Relaxin signaling pathway	ko05164	0.737		
Type I polyketide structures	ko01056	0.738		
Microbial metabolism in diverse environments	ko01120		-0.545	0.650
Pyruvate metabolism	ko00620		-0.554	0.654
Biosynthesis of secondary metabolites	ko01110		-0.515	0.595
Sphingolipid signaling pathway	ko04071		-0.612	0.691
Retinol metabolism	ko00830		0.536	-0.506
Taurine and hypotaurine metabolism	ko00430		-0.522	0.530
Acarbose and validamycin biosynthesis	ko00525		-0.582	0.581
RNA transport	ko03013		-0.549	0.545
Peptidoglycan biosynthesis	ko00550		0.569	-0.627
Carbon fixation pathways in prokaryotes	ko00720		0.543	-0.623
Bisphenol degradation	ko00363			0.556
Renin-angiotensin system	ko04614			0.552
Riboflavin metabolism	ko00740			0.528
MicroRNAs in cancer	ko05206			0.518
Meiosis –yeast	ko04113			-0.530
Solid fraction				
Hypertrophic cardiomyopathy (HCM)	ko05410	0.607		
AGE-RAGE signaling pathway in diabetic complications	ko04933	0.524		
Purine metabolism	ko00230		0.636	
Necroptosis	ko04217		0.608	
Tuberculosis	ko05152		0.580	
Sulfur metabolism	ko00920		-0.599	
Vancomycin resistance	ko01502		-0.524	
C5-Branched dibasic acid metabolism	ko00660		-0.745	0.552
Inositol phosphate metabolism	ko00562			0.564
Flavone and flavonol biosynthesis	ko00944			0.551
Amino sugar and nucleotide sugar metabolism	ko00520			-0.542
2-Oxocarboxylic acid metabolism	ko01210			-0.518

¹Pathways showed significant correlation coefficients (>|0.5|) between the predicted functions of the bacterial community and ruminal LPS activity and the proportions of acetic and propionic acids.

“Biosynthesis of antibiotics” by *Bacillus amyloliquefaciens* play an important role in its bacterial activity to suppress *Rhizoctonia solani* and other fungal plant pathogens as biocontrol agents [41]. Furthermore, the “Amino sugar and nucleotide sugar metabolism” pathway may be involved in carbohydrate transport and metabolism [9], and altered nucleotide sugar metabolism influences exopolysaccharide-producing strains of lactic acid bacteria such as *Streptococcus thermophilus* LY03 [36]. Collectively, significant changes in the bacterial predicted function of the solid fraction might imply the adaptation of rumen bacteria to altered feeding management and rumen fermentation during the fattening stages. However, the investigation of the predicted bacterial function of rumen bacterial communities is still limited. Therefore, further studies are required to elucidate the relationships between changes in rumen fermentation and the relative abundance of bacteria and their metagenomic functions.

Lachnospiraceae and Ruminococcaceae had the highest and second highest relative abundances among bacterial families in the present study, respectively. The Lachnospiraceae family mainly hydrolyzes starch and other sugars to butyric acid and other VFAs [3]. Metagenomic analyses have revealed the ability of the Lachnospiraceae family to decompose complex plant structures and transport decomposed products of various sizes [7]. This is achieved by a secondary product of an “ABC transporter” protein encoded by the Lachnospiraceae family [38]. In addition, the Ruminococcaceae family made up a larger proportion of the fluid

fraction than the solid fraction, which was related to the significantly higher proportion of “Microbial metabolic pathways in diverse environments” in the fluid compared to the solid fraction during the late fattening stage (Fig. 2); in contrast, there was no difference in the functional proportion between the fractions during the same stage (Table 4). Therefore, Lachnospiraceae had a relatively high abundance in the solid fraction, which was probably composed of species that use sugars from decomposed products to synthesize amino acids. Furthermore, the adaptation of Japanese Black beef cattle to long-term ruminal acidic conditions using a high-grain diet to increase intramuscular fat content [22] suggests that the Ruminococcaceae family in the fluid fraction adapted to the late ruminal environment under conditions of significantly higher “Microbial metabolic pathways in diverse environments”.

Pearson’s correlation coefficient analyses revealed that all pathways correlated with LPS were positively correlated, which was influenced by the increase in LPS during the fattening stages. Cell adhesion points and platelet activation were correlated with “ECM-receptor interactions” in the KEGG pathway analyses; ECM-receptors are important in neuronal information processing [8]. In the present study, LPS activity increased throughout the fattening period, and information processing in the ECM receptors and surrounding pathways may have been more active during the latter fattening stages due to an increase in ruminal LPS activity. Furthermore, acetic acid is produced from pyruvate by intestinal bacteria [28], and propionic acid is produced by the conversion of succinate to methylmalonyl-CoA in the TCA circuit from the acrylate pathway [11] and via the propanediol pathway [33]. Therefore, acetic and propionic acids were strongly correlated with pyruvate metabolism, the carbon fixation pathway, and secondary metabolites during glycolysis [20] to produce or decompose acetic and propionic acids in the present study.

In conclusion, although it was difficult to apply the KEGG pathway results directly to rumen bacterial functions, our results suggest that the overall functional structure of the rumen bacterial community is affected by long-term feeding of a high-grain diet. Combined with a dietary transition and changes in ruminal pH, our results suggest underlying differences in fermentation or functional adaptation of Japanese Black beef cattle during the fattening stages.

CONFLICT OF INTEREST. The authors have nothing to disclose.

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