Differences in Circulating microRNAs between Grazing and Grain-Fed Wagyu Cattle Are Associated with Altered Expression of Intramuscular microRNA, the Potential Target PTEN, and Lipogenic Genes

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Abstract

We aimed to understand the roles of miRNAs in the muscle tissue maturation and those of circulating microRNAs (c-miRNAs) in beef production of Japanese Black (JB) cattle (Wagyu), a breed with genetically background of superior intermuscular fat depot, by comparing different feeding conditions (indoor grain-feeding vs. grazing on pasture). The cattle at 18 months old were assigned to pasture feeding or conventional indoor grain feeding conditions for 5 months. Microarray analysis of c-miRNAs from the plasma extracellular vesicles led to the detection of a total of 202 bovine miRNAs in the plasma, including 15 miRNAs that differed between the feeding conditions. Validation of the microarray results by qPCR showed that the circulating miR-10b level in the grazing cattle was upregulated compared to that of the grain-fed cattle. In contrast, the levels of miR-17-5p, miR-19b, miR-29b, miR-30b-5p, miR-98, miR-142-5p, miR-301a, miR-374b, miR-425-5p, and miR-652 were lower in the grazing cattle than in the grain-fed cattle. Bioinformatic analysis indicated that the predicted target genes of those c-miRNAs were enriched in gene ontology terms associated with blood vessel morphogenesis, plasma membrane, focal adhesion, endocytosis, collagen, ECM-receptor interaction, and phosphorylation. In the grazing cattle, the elevation of miR-10b expression in the plasma was coincident with its elevation in the longissimus lumborum (LL) muscle. Expression of bovine-specific miR-2478, the most plasma-enriched miRNA, tended to be also upregulated in the muscle but not in the plasma. Furthermore, grazing caused the downregulated mRNA expression of predicted miR-10b and or miR-2478 target genes, such as DNAJB2, PTEN, and SCD1. Thus, the feeding system used for JB cattle affected the c-miRNAs that could be indicators of grain feeding. Among these, miR-10b expression was especially associated with feeding-induced changes and with the expression of the potential target genes responsible for glucose homeostasis and intramuscular fat depot in the LL muscle of JB cattle.
Introduction

MicroRNAs (miRNAs) are highly conserved, noncoding small RNAs that regulate the expression of target genes in various biological processes in plants and animals [1]. The mature miRNA is generated by processing from the initial transcript (pri-miRNA) via pre-miRNA and recognizes its target genes as a component of the RNA-induced silencing complex (RISC), resulting in mRNA degradation or destabilization. Approximately 30% of protein-coding genes are under the regulation of miRNAs [2].

To better understand the meat quality traits of domestic animals, skeletal muscle miRNAs have been profiled in investigations into how distinct muscle properties are specified. Some muscle miRNAs are differently expressed between muscle types [3,4] or tissues [5,6] in cattle and pigs. Notably, we have demonstrated that miR-196a/b and miR-885 in cattle are expressed exclusively in fast-glycolytic semitendinosus (ST) muscle but not in slow-oxidative masseter muscle [4]. In addition, skeletal muscle miR-206 and miR-208b expression is temporally upregulated, accompanying the downregulation of myosin heavy chain 2x expression, in grazing Japanese Shorthorn (JSH) cattle whereas the expression did not change in housed [7]. Since muscle physiological property determined by the muscle type have impacts on meat quality [8–10], potential association of miRNAs with muscle gene expression and mat quality have been focused recently [11–13]. Besides skeletal muscle, transcriptomic miRNA analyses in mammalian adipose tissue have unveiled their roles in fat accumulation. The relevance of miRNAs has also been pronounced in subcutaneous and/or visceral fat accumulation in response to growth and dietary nutrition in mice [14–16], sheep [17], and cattle [18]. When primary cultured porcine adipocytes are exposed to miR-130b-enriched micro-vesicles, peroxisome proliferator-activated receptor γ (PPARG) expression is downregulated [19].

Cattle feeding systems and dietary nutrients influence the physiological and histological muscle properties as well as growth performance. Compared to the muscles of housed cattle, those of grazing cattle potentially have distinct physiological properties and chemical compositions, including fatty acids and muscle proteins, due to the intermittent movements involved in grazing, which results in changes in subsequent beef quality. Japanese Black (JB; original cattle breed of Wagyu) cattle are conventionally fed a grain-rich concentrate in cattle houses in order to promote their genetically superior intramuscular fat (IMF) depot, which makes beef tender and gives JB beef the high marbling score that is so prized. Thus, differences in muscle growth and IMF deposition resulting from different feeding systems have been progressively investigated.

In particular, the molecular mechanisms underlying IMF depot in different feeding conditions, including diet composition, have been focused on for adipogenic and myogenic gene expression [20,21]. The longissimus lumborum (LL) and ST muscles of pasture-feeding JB cattle show lower extractable lipid content and expression of adipogenic PPARG and CCAAT-enhancer binding protein a (CEBPA) than those of housed JB cattle [22]. The expression patterns of PPARG and CEBPA in intramuscular adipose tissue are higher in cattle fed a high concentrate than those fed a low concentrate [21], indicating that the dietary roughage/concentrate ratio affects IMF fat depot-specific differences in adipogenic gene expression. It is therefore tempting to hypothesize that influences of dietary and feeding conditions on myogenic and adipogenic gene expression are regulated by miRNA, as suggested by differences in miRNAs between subcutaneous and visceral fat [18]. Understanding potential roles of miRNAs in muscle and adipose tissue development will provide new insights into the molecular basis of beef production and meat quality performance.

Moreover, circulating miRNAs (c-miRNAs), which are considered as potential indicators of disease and physiological conditions, also are potentially associated with muscle physiology.
and gene expression as suggested previously [23,24]. Likewise, use of c-miRNA as a monitoring marker could contribute to management and improvement of better beef production. Nevertheless, influences of dietary and feeding conditions on c-miRNAs in beef cattle have never been investigated except for effect of grazing of cattle [24].

In the present study, we therefore aimed to determine (1) the c-miRNAs that are affected by differences between grazing and indoor grain feeding of JB cattle, using an extracellular vesicle (EV)-preparation method from plasma [24], and (2) the muscle miRNAs involved in the feeding-induced regulation of beef quality-related gene expression. The grain-feeding condition employed here leads to higher IMF depot in the JB cattle muscle than the grazing condition [22]. Furthermore we discuss how the c-miRNA changes are associated with skeletal muscle miRNA expression, to understand potential impact of both muscle and circulating miRNAs on beef quality-related gene expression of JB cattle.

Materials and Methods

Animals

The animals were cared for as outlined in the Animal Experimental Guidelines of the NARO Western Region Agricultural Research Center (NARO/WARC) established by the Animal Care Committee, NARO/WARC, and this committee approved the study. All efforts were made to minimize suffering. Eight 10-month-old JB steers, bred at NARO/WARC, were grain-fed separately in a stall barn with no weight difference among the cattle until 10 months of age. Then they were fed a concentrate diet (flaked corn, flaked barley, wheat bran, and soybean meal; 73% total digestible nutrients (TDN) and 11.0% crude protein (CP) on a dry matter (DM) basis) ad libitum and 1.5 kg grass hay (43% TDN and 6.3% CP on a dry matter basis) until 16 months of age. Beginning at 18 months of age, the steers were divided into two groups for 5 months: 4 were continually fed the concentrate diet and grass hay in house and the other 4 were allocated to graze on Italian ryegrass pasture. The health of the animals was monitored every day, which satisfied the criteria of animal welfare defined in the Animal Experimental Guidelines of NARO/WARC. The grass mass was measured during the grazing period and their chemical composition was analyzed (Agricultural Chemistry Research Institute, Hokkaido, Japan) (Table 1). Body weight after the 5-month grazing period differed between the grain-fed (675.5 kg) and the grazing (571.5 kg) cattle (\(P = 0.030\)).

Sample collection

A blood sample was drawn from the jugular vein of each animal at the age of 22 months, and the plasma was prepared with 0.1% EDTA followed by storage at -80°C until use. The LL samples were taken from by biopsy at 22 months. The biopsy procedure was as follows: the animal was locally anesthetized by an intramuscular injection of xylazine (Bayer, Tokyo, Japan) and a subcutaneous injection of lidocaine (AstraZeneca, Osaka, Japan). An incision was subsequently made in the skin overlying the LL muscle [25]. All samples were rapidly frozen in liquid nitrogen and stored at -80°C until RNA extraction.

| Grass mass (kg of DM/ha) | TDN (% of DM) | CP (% of DM) | ADF (% of DM) | NDF (% of DM) |
|-------------------------|---------------|--------------|--------------|--------------|
| 3.787                   | 72.7          | 14.9         | 22.7         | 48           |

DM: dry matter, TDN: total digestible nutrients, CP: crude protein, ADF: acidic detergent fiber, NDF: neutral detergent fiber.

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Plasma sample processing

We previously succeeded in preparing miRNA-enriched EVs as exosomes [24]. In the present study we used the same method to prepare miRNA-containing EVs from plasma. In brief, 10 ml of the plasma sample was mixed with 20 ml of PBS and centrifuged at 1,200 g, 4°C for 20 min. The supernatant was centrifuged at 12,000 g, 4°C for 45 min and further centrifuged at 110,000 g, 4°C for 120 min. The precipitation was suspended in PBS and centrifuged at 110,000 g, 4°C for 70 min. The final precipitation was resuspended in PBS, stored at 4°C for a few days, and then processed for RNA preparation.

RNA preparation

Total RNA including miRNA was extracted from muscle or plasma precipitate using the mirVana microRNA isolation kit (Life Technologies Japan Ltd., Tokyo, Japan) for microarray analysis of plasma samples according to the manufacturer’s protocols. The quantity and quality of the RNA were confirmed by an Agilent Bioanalyzer 2100 with an RNA 6000 Pico Kit (Agilent Technologies, Santa Clara, CA, USA) before the RNA was applied to microarray analysis. Muscle total RNAs for qPCR analysis of muscular mRNA expression were prepared using ISOGEN (Nippon Gene, Tokyo, Japan).

Microarray analysis

Of the four plasma precipitate samples, we used the three samples that satisfied the quantitative requirement as microarray samples for each cattle group. The plasma EV RNA samples for each feeding treatment (grazing or indoor grain-fed) were applied to a custom SurePrint G3 8x60K microarray (Agilent Technologies) that was designed using miRBase ver. 21 (Design ID: 077173). The miRNA Microarray System with miRNA Complete Labeling and Hybridization Kit (Agilent Technologies) was used according to the manufacturer’s protocol. The Agilent microRNA Spike-In Kit was used for in-process control to measure labeling and hybridization efficiency. The labeled samples were hybridized with the probes on the custom microarray, and then the arrays were scanned on a G4900DA SureScan scanner (Agilent Technologies) at 3 μm resolution. Microarray signals were extracted using Agilent Feature Extraction software v11.0 (Agilent Technologies), and the signal values were globally normalized to the 90 percentile using GeneSpring GX (Agilent Technologies), as we did previously to successfully analyze plasma exosomal miRNA samples [24].

To determine significantly different miRNAs between the groups, the normalized signal values were further applied to statistical analysis by significance analysis of microarrays (SAM) [26]. The data were computed with R 2.13.1 (http://cran.r-project.org). The miRNAs were considered significantly different between the groups if the resulting P-value < 0.05. Array data were deposited in the National Center for Biotechnology Information (NCBI) Gene Expression Omnibus (GEO) database, and are accessible through GEO Series accession number GSE81946 (http://www.ncbi.nlm.nih.gov/geo).

cDNA synthesis

cDNA was synthesized from 250 ng of total RNA for muscle samples or 9 μl of the final product of RNA preparation for EV samples, using the miScript II RT kit (Qiagen, Tokyo, Japan) at 37°C for 60 min, and then the enzyme was inactivated at 95°C for 5 min. The cDNAs for muscle samples were each synthesized from 1,000 ng of total RNA by the ReverTra Ace qPCR RT kit (Toyobo, Tokyo, Japan).
Quantitative PCR (qPCR) analysis

qPCR was performed using the CFX96 thermal cycler (Bio-Rad, Hercules, CA, USA) under the following program: 15 min at 95°C, followed by 40 cycles of 15 s at 95°C and 30 s at 60°C. For both plasma and muscle samples, miRNA qPCR was performed using the Thunderbird SYBR qPCR kit (Toyobo) in combination with the miScript Primer Assay for let-7g, RNU6-6P RNA (RNU6-6p), miR-10b, miR-19a, miR-21-5p, miR-27b, miR-29b, miR-30b-5p, miR-30d-5p, miR-92a, miR-103, miR-142-5p, miR-144, miR-196a, miR-197, miR-199-3p, miR-204, miR-206, miR-208b, miR-301a, miR-338, miR-345-5p, miR-374b, miR-378, miR-381, miR-425-5p, miR-451, miR-486, miR-499, miR-652, miR-885, miR-2412, and miR-2478 (Qiagen) according to the manufacturer’s protocol. The qPCR analysis of elongase of very long chain fatty acids 6 (ELOVL6), stearoyl-Coenzyme A desaturase 1 (SCD1), phosphatase and tensin homolog (PTEN), profilin 2 (PFN2), DnaJ heat shock protein family (Hsp40) member B2 (DNAJB2), and PPARG2 was performed using the Thunderbird SYBR qPCR kit (Toyobo), followed by normalization with the expression of ribosomal protein L7 (RPL7; an internal control) [4]. The sequences of the primers used are shown in Table 2. Differences between the groups in the expression ratios of the target miRNA/let-7g for plasma samples, the target miRNA/RNU6 for muscle samples, and that of mRNA/RPL7 were compared. Melting curve analysis was used to confirm the specificity of the amplification reactions.

Prediction and functional annotation of miRNA target genes

The miRNA target genes were predicted using the TargetScan system (Release 7.0, http://www.targetscan.org/). The predicted genes were extracted and ranked by cumulative weighted context++ score [27] at the maximum of 1000 targets for each miRNA. To classify the target genes according to functional annotation, both gene ontology (GO) and pathway analyses were performed on the target genes of c-miRNA differentially expressed in the two groups based on the qRT-PCR results. In this study, the Database for Annotation, Visualization, and Integrated Discovery (DAVID) bioinformatic resources (version 6.7, http://david.abcc.ncifcrf.gov) [28] were applied to the potential target genes with Bos taurus set as the background species, to enrich characteristic GO terms and KEGG pathway defined by Kyoto Encyclopedia of Genes and Genomes (KEGG, http://www.genome.jp/kegg/) for the respective miRNA-mediated biological processes. Extraction of the terms was considered significant when the Benjamini P-value was < 0.10.

Statistical analysis

The expression data are shown as means ± SDs and were compared between the feeding groups by the one-sided or two-sided Student’s t-test. It was expected that grazing would shift the

| Gene  | Accession (Gene) | Sequence (Bases) | Product size (bp) |
|-------|-----------------|------------------|------------------|
| COL3A1 | NM_001076831.1 | GGGACCTGGTTACTTTTCGCTC | 212 |
| DNAJB2 | NM_001034592.1 | TAGACGTACCGCGAAGTTCTCACGCT | 169 |
| ELOVL6 | BC148954.1 | AGAACCCGAACTAGGAGATACAAT | 102 |
| PFN2 | NM_001128197.1 | GGGCGTCTTCCAGAGGATTAA | 139 |
| PPARG2 | NM_181024.2 | TATTTCTCAGTGAGGCCAACC | 181 |
| PTEN | NM_001319898.1 | TCTTCGCAGAGAACAGAAGT | 146 |
| SCD1 | NM_173959.4 | ACCGGCTGGTTGAATAGTGC | 176 |

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muscle properties toward slow-oxidative type and decreased adipogenicity, which is associated with expression of miR-206 [29,30], miR-208b [31], and adipogenic genes such as PPARG2 [20,21,25]. Based on this hypothesis, including presumptive declines in the expression of miR-196a, miR-885, PPARG2, SCD1, and ELOVL6 by grazing, the significance of the differences in the t-test results for those targets was analyzed at one side. The effects of feeding condition were considered as significant if \( P < 0.05 \). Significance levels of \( P \leq 0.10 \) were considered a statistical trend.

Results

Circulating miRNA profile of plasma extracellular vesicles in JB cattle

To profile c-miRNAs of JB cattle fed under two different systems, we conducted microarray analysis using plasma EV precipitates prepared by ultracentrifugation. The c-miRNA profile obtained from the averaged raw signal value of microarray results for each group showed that the top 20 miRNAs were 85.7 and 86.8% of the total miRNAs in the grain-fed and grazing cattle, respectively (Fig 1). Most of the top 20 c-miRNAs did not differ between the groups in the order of the amount or the ratio of composition. The only differences were that miR-20a was present at higher levels in the grain-fed cattle than in the grazing cattle, and that miR-29a was present at higher levels in the grazing cattle, although both miR-20a and miR-29a were within the top 25 miRNAs in both cattle groups. In the plasma samples of both groups, miR-2478 and miR-1260b were approximately 46 and 17% of the total miRNAs, respectively (Fig 1). Our previous studies revealed that skeletal muscle-specific miRNAs, namely miR-1, miR-133a/b, miR-206, miR-208a/b, miR-496, and miR-499, were abundant in the muscles of JB cattle [4], whereas none of them was detected in the plasma profiles except for a modest miR-486 content [24]. In contrast, the plasma samples were enriched with moderate concentrations of miRNAs associated with adipose tissue, such as miR-15a, miR-19a/b, miR-21, miR-27b, miR-92a/b, miR-103, miR-106a/b, miR-107, miR-125b, and miR-150 [11,12]. It is especially notable that the bovine-specific miR-1584-5p, miR-2305, miR-2412, and miR-2478, most of which are associated with energy metabolism [32], were highly enriched in the plasma samples of both groups in the present study.

Feeding conditions affect circulating miRNA contents in JB cattle

The statistical SAM analysis led to the extraction of a total of 15 c-miRNAs that differed between the groups (\( P < 0.05 \), Table 3). Of those miRNAs, the contents of miR-652, miR-30d, miR-301a, miR-345-5p, miR-374b, miR-425-5p, miR-23b-3p, miR-30b-5p, miR-17-5p, miR-98, miR-28, and miR-874 were lower in the plasma of the grazing cattle than in that of the grain-fed cattle, whereas the contents of miR-10b, miR-2368-3p, miR-885, and miR-2425-3p were higher in the plasma of the grazing cattle. The contents of miR-2368-3p and miR-2425-3p, putative ruminant-specific miRNAs [31], were very low (less than 0.067%), and no GO term was extracted from the predicted target genes of either miRNA by DAVID analysis.

Therefore, we further focused on the other 13 miRNAs as biologically relevant c-miRNAs. We then conducted qPCR for the contents of c-miRNAs of interest, not only to validate the results of microarray analysis for those 13 miRNAs and the miRNAs below the top 20 in SAM analysis (miR-29b, miR-197), but also to further explore potential feeding-induced miRNAs among adipose-enriched (miR-15a, miR-19a, miR-27b, miR-92a, miR-103, miR-142-5p), plasma-enriched (miR-2412, miR-2478), and bovine-specific (miR-2284x and miR-2295) miRNAs [31]. The results revealed that circulating miR-10b content was higher in grazing cattle than in grain-fed cattle (\( P = 0.005 \), Fig 2). In contrast, the grazing cattle showed significantly lower contents of miR-17-5p (\( P = 0.031 \)), miR-19a (\( P = 0.007 \)), miR-29b (\( P = 0.021 \)), miR-30b-5p (\( P = 0.035 \)), miR-98 (\( P = 0.006 \)), miR-142-5p (\( P = 0.013 \)), miR-301a (\( P = 0.005 \)), miR-374b
Fig 1. Circulating miRNA (c-miRNA) profiles in plasma extracellular vesicles of Japanese Black cattle obtained in microarray analysis. Percentages of c-miRNA contents in grain-fed (red bars) and grazing cattle (green bars) are indicated. The top 20 miRNAs in plasma of grazing cattle are listed.

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miR-652 ($P = 0.010$), miR-30d ($P = 0.012$), miR-301a ($P = 0.016$), miR-345-5p ($P = 0.017$), miR-2368-3p ($P = 0.025$), miR-374b ($P = 0.030$), miR-425-5p ($P = 0.031$), miR-885 ($P = 0.035$), miR-23b-3p ($P = 0.035$), miR-30b-5p ($P = 0.040$), miR-17-5p ($P = 0.040$), miR-98 ($P = 0.042$), miR-2425-3p ($P = 0.045$), miR-28 ($P = 0.046$), miR-874 ($P = 0.048$), miR-17-3p ($P = 0.055$), miR-10b ($P = 0.056$), miR-151-3p ($P = 0.056$), miR-30e-5p ($P = 0.058$), and miR-130b ($P = 0.059$) were all significantly different.

The data was statistically analyzed by SAM method as described in Materials and Methods.

$\log_2(G/H)$: the value of $\log_2$(grazing/grain-fed)

($P = 0.016$), miR-425-5p ($P = 0.010$), and miR-652 ($P = 0.046$). All other miRNAs tested (miR-15a, miR-23b-3p, miR-27b, miR-30d-5p, miR-92a, miR-140, miR-197, miR-345-5p, miR-451, miR-885, miR-2284x, miR-2295, miR-2412, and miR-2478) were not differently expressed.

![Fig 2. Circulating miRNA expression in plasma extracellular vesicles of grain-fed and grazing Japanese Black cattle analyzed by qRT-PCR. ** and * indicate the differences between the feeding conditions at $P < 0.01$ and $< 0.05$, respectively.](doi:10.1371/journal.pone.0162496.g002)
between the groups ($P > 0.10$), except that miR-103 content tended to be higher in the grain-fed cattle than in the grazing cattle ($P = 0.057$).

**Feeding conditions also affect skeletal muscle miRNA expression**

Next, we analyzed the muscle tissue expression of the 11 miRNAs that differed significantly between the groups (miR-10b, miR-17-5p, miR-19a, miR-29b, miR-30b-5p, miR-98, miR-142-5p, miR-301a, miR-374b, miR-425-5p, and miR-652) to determine whether or not c-miRNA expression was associated with that of skeletal muscle tissue miRNAs.

Intriguingly, the miR-10b expression in LL muscle was coincidently higher in the grazing cattle than in the grain-fed cattle ($P = 0.024$, Fig 3). miR-374b and miR-652 also tended toward higher expression in the grazing cattle than in the grain-fed cattle ($P = 0.062$ and $= 0.061$, respectively). We also conducted qPCR of miRNAs that are enriched in muscle or plasma (miR-21-5p, miR-30d-5p, miR-103, miR-206, miR-208b, miR-451, miR-486, miR-499, miR-2412, and miR-2478), some of which are abundant in bovine skeletal muscles [4]. Of those miRNAs, only miR-2478 in LL muscle tended to differ between the feeding conditions ($P = 0.053$).

To predict potential cellular biological events that the significant miRNAs are involved in, a functional annotation of the miRNA target genes predicted by TargetScan was performed by DAVID. The results of the GO and KEGG pathway analyses showed that the potential target genes of uniquely upregulated miR-10b were associated with cell adhesion (Table 4). The potential targets of downregulated c-miRNAs were associated with protein phosphorylation and phosphate metabolism (miR-17-5p), endocytosis (miR-19a, miR-301a), mTOR signaling (miR-19a), MAPK signaling (miR-19a, miR-98), ECM-receptor interaction and adipocytokine signaling (miR-29b), skeletal muscle tissue development (miR-30b-5p), blood vessel development and morphogenesis, and regulation of actin cytoskeleton (miR-301a). The potential targets of the most plasma-enriched miR-2478 were associated with protein localization and

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Fig 3. miRNA expression in longissimus lumborum muscle of grain-fed and grazing Japanese Black cattle analyzed by qRT-PCR. * and + indicate the differences between the feeding conditions at $P < 0.05$ and $< 0.10$, respectively.

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Table 4. Potential cellular biological events predicted from the microRNA target genes.

| miRNA  | Category            | Term                                                                 | Fold enrichment | Benjamin
|--------|---------------------|----------------------------------------------------------------------|----------------|-----------
| miR-10b| GOTERM_BP_FAT       | GO:0007156—homophilic cell adhesion                                   | 4.535          | 0.003     |
|        | GOTERM_BP_FAT       | GO:0016337—cell-cell adhesion                                         | 3.443          | 0.004     |
|        | GOTERM_BP_FAT       | GO:0022610—biological adhesion                                        | 2.203          | 0.062     |
|        | GOTERM_BP_FAT       | GO:0007155—cell adhesion                                              | 2.203          | 0.062     |
| miR-17-5p| GOTERM_BP_FAT  | GO:0006468—protein amino acid phosphorylation                         | 2.161          | 0.010     |
|        | GOTERM_BP_FAT       | GO:0045449—regulation of transcription                                | 1.598          | 0.017     |
|        | GOTERM_BP_FAT       | GO:0006796—phosphate metabolic process                                | 1.836          | 0.021     |
|        | GOTERM_BP_FAT       | GO:0006793—phosphorus metabolic process                               | 1.836          | 0.021     |
|        | GOTERM_BP_FAT       | GO:0016310—phosphorylation                                            | 1.905          | 0.035     |
|        | GOTERM_BP_FAT       | GO:0010604—positive regulation of macromolecule metabolic process     | 2.142          | 0.084     |
|        | KEGG_PATHWAY        | bta05219:Bladder cancer                                               | 5.341          | 0.075     |
| miR-19a| GOTERM_BP_FAT       | GO:0007242—intracellular signaling cascade                           | 2.012          | 0.028     |
|        | KEGG_PATHWAY        | bta04010:MAPK signaling pathway                                       | 3.299          | 0.075     |
|        | KEGG_PATHWAY        | bta04510:Focal adhesion                                               | 2.985          | 0.012     |
|        | KEGG_PATHWAY        | bta04512:ECM-receptor interaction                                     | 3.817          | 0.033     |
|        | KEGG_PATHWAY        | bta04930:Type II diabetes mellitus                                    | 4.297          | 0.049     |
|        | KEGG_PATHWAY        | bta04144:Endocytosis                                                 | 2.268          | 0.058     |
|        | KEGG_PATHWAY        | bta04100:MAPK signaling pathway                                       | 2.019          | 0.066     |
|        | KEGG_PATHWAY        | bta04150:mTOR signaling pathway                                      | 3.981          | 0.071     |
|        | KEGG_PATHWAY        | bta05410:Hyperthrophic cardiomyopathy (HCM)                           | 3.133          | 0.079     |
| miR-29b| KEGG_PATHWAY        | bta04510:Focal adhesion                                               | 3.529          | 0.000     |
|        | KEGG_PATHWAY        | bta04512:ECM-receptor interaction                                     | 4.611          | 0.001     |
|        | KEGG_PATHWAY        | bta05200:Pathways in cancer                                           | 2.201          | 0.008     |
|        | KEGG_PATHWAY        | bta05222:Small cell lung cancer                                       | 3.705          | 0.011     |
|        | KEGG_PATHWAY        | bta04920:Adipocytokine signaling pathway                              | 3.529          | 0.097     |
| miR-30b-5p| GOTERM_BP_FAT   | GO:0007242—intracellular signaling cascade                           | 1.996          | 0.034     |
|        | GOTERM_BP_FAT       | GO:0007519—skeletal muscle tissue development                         | 5.780          | 0.098     |
|        | GOTERM_BP_FAT       | GO:0060538—skeletal muscle organ development                          | 5.780          | 0.098     |
| miR-98 | KEGG_PATHWAY        | bta04010:MAPK signaling pathway                                       | 2.598          | < 0.001   |
|        | KEGG_PATHWAY        | bta05200:Pathways in cancer                                           | 2.203          | 0.005     |
| miR-142-5p| KEGG_PATHWAY   | bta05200:Pathways in cancer                                           | 2.106          | 0.048     |
| miR-301a| GOTERM_BP_FAT      | GO:0007242—intracellular signaling cascade                           | 2.211          | 0.001     |
|        | GOTERM_BP_FAT       | GO:0048514—blood vessel morphogenesis                                | 3.627          | 0.051     |
|        | GOTERM_BP_FAT       | GO:0044087—regulation of cellular component biogenesis                | 4.885          | 0.073     |
|        | GOTERM_BP_FAT       | GO:007264—small GTPase mediated signal transduction                   | 2.675          | 0.074     |
|        | GOTERM_BP_FAT       | GO:0008104—protein localization                                      | 1.932          | 0.076     |
|        | GOTERM_BP_FAT       | GO:0009792—embryonic development ending in birth or egg hatching     | 2.643          | 0.079     |

(Continued)
The genes predicted as targets of miR-374b and miR-652 were not significantly linked with any specific GO terms or KEGG pathways.

Association between miRNA target expression and miRNAs in skeletal muscle

Hypothesizing that the differences in miRNA expression could affect the regulation of miRNA target genes, we further analyzed expression of potential miRNA target genes in LL muscle tissue by qPCR. Although cell adhesion-related genes were employed for extraction of miR-10b-related GO terms in bioinformatic analysis, those genes showed lower cumulative weighted context++ scores [27] than PTEN and ELOVL6 genes. In addition, most of the cell adhesion-related genes were not expressed in JB cattle muscles according to our preliminary analysis. Therefore, based on the result of the target gene prediction by TargetScan analysis, we especially focused on the genes associated with fat and muscle growth, development, and/or metabolism among the predicted genes, to assess the relationships between circulating or muscle miRNAs and the gene expression related to beef quality. One of the lipogenic genes, SCD1, which is predicted as a target of miR-98, miR-142-5p, and miR-2478, showed higher expression in the grain-fed than in the grazing cattle (P = 0.038, Fig 4). The other adipogenic or lipogenic genes, miR-301a target PPARG2 and a target of miR-10b and miR-2478, ELOVL6, also tended to differ between the groups (P = 0.058 and = 0.051, respectively). The expression of PTEN, the potential target of miR-10b, miR-17-5p, miR-19a, miR-29b, miR-30b-5p, miR-142-5p, miR-301a, miR-652, and miR-2478, was lower in the grazing cattle than in the grain-fed cattle (P = 0.011). The mRNA expression of actin-binding protein PPN2, the potential target of miR-17-5p, miR-19a, miR-30b-5p, miR-142-5p, miR-301a, and miR-2478, also tended to be down-regulated in the grazing cattle compared to the grain-fed cattle (P = 0.064), as did the potential miR-29b targets DNAJB2 (P = 0.021) and COL3A1 (P = 0.100). These expression patterns of the miRNA targets indicated that the expressions of these potential miRNA target genes are

Table 4. (Continued)

| miRNA | Category | Term | Fold enrichment | Benjamini |
|-------|----------|------|-----------------|-----------|
| GOTERM_BP_FAT | GOTERM_BP_FAT | GO:0043009–chordate embryonic development | 2.660 | 0.081 |
| GOTERM_BP_FAT | GOTERM_BP_FAT | GO:0001568–blood vessel development | 3.043 | 0.085 |
| GOTERM_BP_FAT | GOTERM_BP_FAT | GO:0051173–positive regulation of nitrogen compound metabolic process | 2.333 | 0.085 |
| GOTERM_BP_FAT | GOTERM_BP_FAT | GO:0001944–vasculature development | 3.165 | 0.086 |
| GOTERM_BP_FAT | GOTERM_BP_FAT | GO:0045935–positive regulation of nucleobase, nucleoside, nucleotide and nucleic acid metabolic process | 2.300 | 0.091 |
| GOTERM_BP_FAT | GO:0010628–positive regulation of gene expression | 2.338 | 0.094 |
| KEGG_PATHWAY | bta04144:Endocytosis | 2.784 | 0.003 |
| KEGG_PATHWAY | bta04115:p53 signaling pathway | 4.262 | 0.006 |
| KEGG_PATHWAY | bta04810:Regulation of actin cytoskeleton | 2.281 | 0.048 |
| miR-2478 | GOTERM_BP_FAT | GO:0045184–establishment of protein localization | 2.096 | 0.055 |
| GOTERM_BP_FAT | GOTERM_BP_FAT | GO:0015031–protein transport | 2.101 | 0.079 |
| GOTERM_BP_FAT | GOTERM_BP_FAT | GO:0008104–protein localization | 2.077 | 0.088 |
| KEGG_PATHWAY | bta04340:Hedgehog signaling pathway | 4.629 | 0.079 |

The data were statistically analyzed by DAVID as described in Materials and Methods.

1Fold enrichment = (genes in list that are annotated with the term)/(total numbers of genes that are annotated with any term in the term category)

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**Fig 4.** miRNA target gene expression in *longissimus lumborum* muscle of grain-fed and grazing Japanese Black cattle analyzed by qRT-PCR. *+* and *+* indicate the differences between the feeding conditions at \( P < 0.05 \) and \( P < 0.10 \), respectively.

**Table 5.** Summary of altered miRNAs and the miRtarget genes of interest in LL muscle of the grazing JB cattle compared to the grain-fed.

| miRNA   | Circulation \(^2\) | LL muscle \(^2\) | DNAJB2 | SCD1    | EVOLV6 | PPARG | PTEN | PI3K signaling | COL3A1 | PFN2 | DNAJB2 |
|---------|-------------------|-----------------|--------|---------|--------|--------|------|----------------|--------|------|--------|
| miR-10b | ⬆⬆               | ⬆⬆             | +      | +       |
| miR-17-5p| ⬇               | +               | +      | +       |
| miR-19a  | ⬇                 | +               | +      | +       |
| miR-29b  | ⬇                 | +               | +      | +       |
| miR-30b-5p| ⬇               | +               | +      | +       |
| miR-98   | ⬇                 | -               | +      | +       |
| miR-142-5p| ⬇               | +               | +      | +       |
| miR-301a | ⬇                 | +               | +      | +       |
| miR-374b | ⬇                 | +               | +      | +       |
| miR-425-5p| ⬇               | +               | +      | +       |
| miR-652  | ⬇                 | +               | +      | +       |
| miR-2478 | ⬇                 | +               | +      | +       |

\(^1\) + indicates the gene targeted by the altered miRNAs in the top row of this table.

\(^2\) \(\uparrow\) or \(\downarrow\): up- or down-regulated at \( P < 0.05 \), \(\uparrow\) or \(\downarrow\): up- or down-regulated at \( P < 0.10 \), \(\cdot\): not changed (\( P \geq 0.10 \)).

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associated with grain-fed housing-induced changes in circulating and muscle miRNAs in JB cattle. These relevant results are summarized in Table 5.

**Discussion**

It has been demonstrated that diet and supplementary nutrients affect miRNA expression in adipose tissues of mice [14–16], sheep [17] and cattle [18]. However, miRNAs in circulation and skeletal muscle have poorly understood in farm animals thus far. Nevertheless, a polyunsaturated fatty acids (PUFAs)-enriched diet might have the potential to alter the expression of c-miRNAs linked to changes in dietary fatty acids (as inferred from plasma fatty acid concentration) [33]. In the present study, we focused on the effects of feeding conditions on c-miRNAs and skeletal muscle miRNAs as well as on muscle target genes in JB cattle, whose beef has high marbling potential, to understand not only the implications of feeding diet and grazing on the expression of the set of miRNAs and their target genes in skeletal muscles, but also the associations between altered muscle gene expression and c-miRNAs. Most of studies examining effects of grain-rich high concentrate feeding compared to high roughage feeding or grazing resulted in higher IMF depot but accompanied higher body weight of JB cattle in the concentrate feeding than in the roughage feeding [22,34], nevertheless a high concentrate feeding that showed higher IMF depot without significant difference in final body weight between cattle fed high and low concentrate indicated a positive effect of grain-rich concentrate [21,35]. In the present study, grain-fed cattle showed significantly higher body weight than grazing cattle, nevertheless the higher lipogenic gene expression in the grain-fed cattle could partly be caused by the difference in the diet.

**Characteristics of plasma extracellular vesicle miRNA profile in JB cattle**

We profiled the plasma EV miRNAs in JB cattle in both the indoor grain-fed and grazing conditions. Previously we successfully prepared EV miRNAs from plasma of cattle by an ultra-centrifugation method [24], and we therefore employed the same method in the present study. The present results revealed no significant difference between the feeding conditions in the profiles of abundant miRNAs in the plasma EV of cattle. Those profiles were enriched with bovine-specific miR-1584-5p, miR-2305, miR-2412, and miR-2478, which are associated with energy metabolism [32]. Although it is unknown why those miRNAs were abundant in the profiles, they may be involved in bovine-specific energy metabolism. Regarding the top 20 c-miRNAs, the c-miRNA profiles of both JB cattle groups were considerably similar to those of JSH cattle [24], except that in the latter, miR-15b, miR-19b, miR-20a, and miR-26a were not in the top 20 but were above 30. The differences between JB and JSH in beef quality traits such as marbling (IMF depot) [36] may not be associated with composition of those major c-miRNAs.

**miR-10b and miR-2478 in circulation and skeletal muscle of JB cattle**

The present results of microarray and qPCR revealed that the grazing JB cattle had a higher plasma content of miR-10b and lower plasma contents of miR-17-5p, miR-23b-3p, miR-28, miR-30b-5p, miR-30d, miR-98, miR-301a, miR-345-5p, miR-374b, miR-425-5p, miR-652, and miR-874 than the grain-fed cattle. Most of the differently expressed c-miRNAs were downregulated in the grazing group. In contrast, the unique miR-10b upregulation in the grazing cattle plasma deserves taking notice. As we hypothesize that c-miRNA changes indicate alterations of skeletal muscle physiology, miR-10b expression in the LL muscle was upregulated coincidently with the change of circulating miR-10b, suggesting involvement of miRNAs in muscle property alteration. The expression levels of muscle miR-374b, miR-652, and miR-2478 also tended to be higher in the LL muscle of the grazing cattle. To date, c-miRNAs in cattle have been reported in...
relation to grazing [24], heat stress [23], pregnancy [37], and viral infection [38], but not in relation to changes in circulating miR-10b. The altered miRNAs could be indicators of feeding conditions such as grain abundance and may be useful noninvasive biomarkers to monitor feeding.

Association of miR-10b with the target PTEN in JB cattle muscle

TargetScan analysis predicted that miR-10b and miR-2478 targeted PTEN. According to a bioinformatic analysis of potential implications of the circulating and muscle miR-10b upregulation, GO terms related to cell adhesion were extracted as biological events associated with the predicted miR-10b target genes. However, most of the cell adhesion-related genes might not be meaningful due to lack of expression in JB cattle muscle in our preliminary result. The PTEN expression in the LL muscle was lower in the grazing JB cattle than in the grain-fed cattle. Indeed, it was demonstrated that miR-10b can target PTEN and cause cell proliferation and/or migration [39,40]. PTEN is the phosphatase that converts phosphatidylinositol 3,4,5-triphosphate (PIP$_3$) to inactivate phosphatidylinositol 4,5-bisphosphate (PI-4,5-P$_2$), and it inhibits phosphatidylinositol-3-kinase/Akt (PI3K/Akt) signaling [41], which is a key determinant of the insulin-dependent increase in glucose uptake into muscle and adipose cells. Overexpression of PTEN inhibits GLUT4 translocation and glucose uptake in 3T3-L1 adipocytes [42]. Skeletal muscle-specific deletion of PTEN protects mice from developing insulin resistance and diabetes induced by a high-fat diet, and thus improves glucose homeostasis [43]. By inhibiting PI3K/Akt signaling, PTEN can regulate muscle cell differentiation [44] and hypertrophy under the reduced availability of insulin-like growth factor binding protein 2 (IGFBP2) [45]. Thus, the reduced PTEN expression in the grazing cattle suggests alterations in both muscle cell growth and glucose homeostasis in the LL muscle, which might be modulated by miR-10b and possibly miR-2478, and might further affect final beef production.

Potential impact of miR-2478 and its targets on JB cattle muscle

miR-2478 potentially targets mRNAs of lipogenic genes ELOVL6 and SCD1. ELOVL6 catalyzes the elongation of saturated and monounsaturated fatty acids with 12-, 14-, and 16-carbons in the endoplasmic reticulum of the adipocyte. SCD1 converts a portion of stearic acid (18:0) into oleic acid (18:1) [46]. The expression of both ELOVL6 and SCD1 increased during IMF deposition and temporally had a very large upregulation in longissimus muscle of Angus steers fed a high-starch diet after an early weaning, compared to normally weaned cattle [47]. This supports our present observation of downregulated ELOVL6 and SCD1 expression in the grazing JB cattle compared to the grain-fed cattle. The downregulated ELOVL6 and SCD1 expression in our study was associated with the increased miR-10b and miR-2478 expression in the LL muscle of grazing JB cattle. Therefore, it is likely that the expression of those lipogenic genes induced by grain-rich feeding was also modulated by miR-10b and/or miR-2478 in the IMF of the LL muscle, and might affect beef quality.

miR-2478 was also predicted to target PFN2 and DNAJB2, whose expression tended to be reduced in the LL muscle of the grazing cattle in the present study. Especially, profilin binds to actin filament at the barbed end and affects actin homeostasis during cell movement [48]. In addition, because profilin can bind to PI-4,5-P$_2$ [49], PFN2 expression might also be associated with the alteration of PTEN expression in the LL muscle, since profilin overexpression induced PTEN upregulation and reduced Akt expression [50]. Expression of DNAJB2, a co-chaperone regulator of heat shock protein Hsp70 that is expressed principally in the nervous system, is induced in human skeletal muscle during its recovery from damage by exercise [51]. In a study using samples of skeletal muscle and the pathologies associated with protein aggregation, it was proposed that DNAJB2 has a role in protein turnover in skeletal muscle [52]. The higher muscle DNAJB2 expression in the grain-fed JB cattle than in the grazing cattle might indicate that
Differences in Circulating and Muscle miRNAs between Grazing and Grain-Fed Wagyu Cattle

DNAJB2 was upregulated via downregulation of miR-2478 by some cellular stress due to difference between the feeding conditions.

Association of circulating miRNA with muscle miRNA and the biological impact on JB cattle

In this study, it is concluded that the differing c-miRNAs between the feeding conditions could be an indicator not only of the grain abundance of a diet but also changes in gene expression associated with PI3K signaling and lipogenesis in the LL muscle of JB cattle. To date, however, the effects of c-miRNAs on tissues including EV recipient cells remain poorly understood.

We concluded that difference in circulating miR-10b between the feeding conditions was associated with the miR-10b upregulation in the LL muscle of cattle. EVs containing miR-210 isolated from metastatic breast cancer cells promote metastasis via the induction of angiogenesis in the tumor, and the addition of miR-210-enriched EVs induced the activation of endothelial cells in vitro [53]. Although roles of c-miRNAs to downregulate the muscle target genes were not demonstrated in this study, at least c-miRNAs, especially miR-10b, might either be taken into the LL muscle, or be secreted from the tissues inside the cattle body including muscles. High-fat diet feeding to cattle raises expression of miRNAs including miR-19a in both the subcutaneous and visceral fat [18], suggesting that high energy diet cause increase in adipose tissue miRNA expression. It is therefore likely that the present grain feeding as a higher energy diet altered miRNA expression in adipose tissue and might promote secretion of exosomes containing miRNAs such as miR-19a into plasma of the grain-fed cattle. It is necessary to clarify the departing and destination tissues of c-miRNAs and the mechanism how the c-miRNAs are balanced between circulation and various tissues of secretion and recipient.

In the present study, muscle PTEN and PFN2 were the predicted targets of most of the altered c-miRNAs (miR-17-5p, miR-19a, miR-30b-5p, miR-142-5p, miR-301a) between the feeding conditions, as well as miR-10b. Moreover, the results of bioinformatic functional annotation indicated that the altered c-miRNAs are associated with cellular biological events involved in phosphorylation, various intracellular signaling pathways, skeletal muscle and blood vessel development, and protein localization. Taking together, c-miRNAs might affect tissue development and maturation in cattle. Further research is needed to elucidate actions of feeding-induced c-miRNAs on the expression of the target genes in potential recipient tissues.

Conclusion

The present study revealed that the grazing of JB cattle on pasture affected the plasma c-miRNAs compared to grain-feeding JB cattle. Especially, miR-10b expression increased in both circulation and the LL muscle in the grazing cattle. The altered expression of miR-10b and miR-2478 in the LL muscle was further associated with the expression of the target lipogenic genes, ELOVL6 and SCD1, respectively. Moreover, expression of the factor regulating glucose homeostasis, PTEN, was also reduced in the grazing cattle. These results indicated that changes in c-miRNAs induced by a grain-rich diet are associated with skeletal muscle miRNA and the potential target gene expression, suggesting in turn that muscle miRNAs are involved in the regulation of muscle and adipogenic genes that determine beef quality.

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Author Contributions
Conceived and designed the experiments: SM.
Performed the experiments: SM MS.
Analyzed the data: SM MO.
Contributed reagents/materials/analysis tools: SM MS MH KO.
Wrote the paper: SM.

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