MicroRNA-449a deficiency promotes colon carcinogenesis

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MicroRNAs have broad roles in tumorigenesis and cell differentiation through regulation of target genes. Notch signaling also controls cell differentiation and tumorigenesis. However, the mechanisms through which Notch mediates microRNA expression are still unclear. In this study, we aimed to identify microRNAs regulated by Notch signaling. Our analysis found that microRNA-449a (miR-449a) was indirectly regulated by Notch signaling. Although miR-449a-deficient mice did not show any Notch-dependent defects in immune cell development, treatment of miR-449a-deficient mice with azoxymethane (AOM) or dextran sodium sulfate (DSS) increased the numbers and sizes of colon tumors. These effects were associated with an increase in intestinal epithelial cell proliferation following AOM/DSS treatment. In patients with colon cancer, miR-449a expression was inversely correlated with disease-free survival and histological scores and was positively correlated with the expression of MLH1 for which loss-of function mutations have been shown to be involved in colon cancer. Colon tissues of miR-449a-deficient mice showed reduced Mlh1 expression compared with those of wild-type mice. Thus, these data suggested that miR-449a acted as a key regulator of colon tumorigenesis by controlling the proliferation of intestinal epithelial cells. Additionally, activation of miR-449a may represent an effective therapeutic strategy and prognostic marker in colon cancer.

Mature microRNAs (miRNAs) are noncoding RNAs of approximately 22 nucleotides that regulate gene expression by targeting the 3′-untranslated regions (UTRs) of mRNAs, resulting in inhibition of mRNA translation1,2. MiRNAs regulate various aspects of cell physiology, including proliferation, differentiation, cell death, and development1. Moreover, aberrantly expressed miRNAs are involved in tumorigenesis, as either oncogenes or tumor suppressors3–5. For example, miR-34a-5p suppresses colorectal cancer metastasis, and its expression predicts recurrence in patients with stage II/III colorectal cancer6. Additionally, miR-183 functions as an oncogene and promotes tumor cell migration7. Therefore, miRNAs may be potential targets for cancer therapy.

Notch signaling also has pleiotropic roles in cell differentiation, proliferation, and cell death, and dysregulation of Notch is involved in many types of malignant tumors, including T-cell acute lymphoblastic leukemia (T-ALL) and lymphoma8–10. Notably, we have previously reported that Notch is an essential mediator of T-cell differentiation11, 12 and maintenance of memory CD4 T cells13. Moreover, in studies of Notch and cancer, frequent active mutations in Notch1 have been reported in patients with T-ALL14, 15 and gain-of-function mutations and copy numbers of Notch2 are increased in diffuse large B-cell lymphoma16.

MicroRNA-449a (miR-449a) is a member of the miR-449 family (miR-449a, miR-449b, and miR-449c). The miR-449 cluster contains sequences and secondary structures similar to those of the miR-34 family and has therefore been classified as a single family of miRNAs. The expression of miR-449a is decreased in several cancers, including gastric and bladder cancer17, 18. Furthermore, miR-449a regulated several genes associated with tumorigenesis, including the gene encoding histone deacetylase (HDAC)19 and CDC25A20, suggesting that miR-449a may have oncogenic effects. However, the roles of miR-449a in tumorigenesis in vivo have not yet been determined.

In this study, we investigated whether miRNAs were regulated by Notch signaling. Our results showed that miR-449a was upregulated by Notch signaling. Unexpectedly, miR-449a-deficient mice did not show any defects...
in the development of T cells, marginal zone B cells, and CD8α− splenic dendritic cells, all of which are regulated by Notch signaling. Additionally, miR-449a-deficient mice showed increased susceptibility to azoxymethane (AOM) and dextran sodium sulfate (DSS)-induced colon cancer with increased proliferation of intestinal epithelial cells. Furthermore, the expression of miR-449a was inversely correlated with histological scores and disease-free survival in patients with colon cancer. Deficiency of miR-449a in the colon resulted in downregulation of Mlh1, and expression of miR-449a was positively correlated with that of Mlh1 in patients with colon cancer. These data highlighted the role of miR-449a as a tumor suppressor in colon cancer and suggested that miR-449a may be a therapeutic target in the treatment of colon cancer.

Results
Notch regulated miR-449a expression. We first searched for miRNAs regulated by Notch signaling using an miRNA microarray. Because the interactions between Notch and Notch ligands allow γ-secretase to cleave Notch, resulting in translocation of the intracellular domain of Notch into the nucleus, overexpression of the intracellular domain of Notch in cells can activate Notch signaling. We compared the expression of miRNA between DO11.10 cells infected with control retrovirus or retrovirus carrying the intracellular domain of Notch1 (Fig. 1a). Only miR-449a was upregulated by more than 3 fold in DO11.10 cells infected with the intracellular domain of Notch1, as confirmed by real-time polymerase chain reaction (PCR) analysis (Fig. 1a and b). Additionally, miR-22321, which has been reported to be related to Notch signaling, was not altered in this comparison (Fig. 1a). Because Rbpj is essential for Notch signaling, we compared the expression of miR-449a in Rbpj-deficient and wild-type T cells. Rbpj-deficient T cells from Rbpjfl/fl mice crossed with CD4-Cre transgeni
mice showed substantially reduced expression of miR-449a compared with that in wild-type cells (Fig. 1c), although miR-449a expression was still detected in Rbpj-deficient cells. These data demonstrated that Notch was an upstream regulator of miR-449a.

We next sought to evaluate whether Notch signaling directly controlled miR-449a expression. There were two putative Rbpj binding regions upstream of the miR-449a locus in homologous regions shared between mice and humans (depicted as 11k and 450) (Fig. 1d). We performed chromatin immunoprecipitation assays with anti-Rbpj antibodies and detected binding of Rbpj in Dtx1, a known Notch target gene. The relative increase in PCR products in two regions by anti-Rbpj antibody compared with that of the control antibody was similar to that of the Cnot3 region (Fig. 1d), suggesting the indirect regulation of miR-449a expression by Notch signaling.

Establishment of miR-449a-deficient mice. In order to evaluate the roles of miR-449a in immune cell development and tumorigenesis, we established miR-449a-deficient (miR-449a−/−) mice. The miR-449a locus was replaced with a neo-cassette (Fig. 2a), and homologous recombination was confirmed by Southern blotting and PCR (Fig. 2b and Supplementary Figure 2). T cells from miR-449a+/+ or miR-449a−/− mice showed approximately 50% expression or no expression of miR-449a, respectively, compared with those of wild-type cells (Fig. 2c), indicating that complete deficiency of miR-449a expression was achieved in miR-449a−/− mice. miR-449a−/− mice were born according to Mendelian inheritance rules and did not show any gross body changes. miR-449a−/− mice were viable for up to at least 60 weeks of age (data not shown).

We assessed the expression of miR-449a in various organs by real-time PCR. Our results showed that miR-449a was highly expressed in the thymus and lung, but was expressed at low levels in the brain and colon (Fig. 2d).

MiR-449a deficiency did not affect immune cell development. Because Notch signaling regulates early T-cell development and effector T-cell differentiation, we evaluated changes in lymphocytes and the thymus and lung. We found that miR-449a deficiency did not affect the development of T cells in the thymus and lung.

**Figure 2.** Establishment of miR-449a-deficient mice. (a) Configuration of the targeting vector of miR-449a-deficient mice. (b) Genomic DNA was digested with BclI and subjected to Southern blot analysis. Genomic DNA from wild-type or miR-449a−/− mice was amplified by PCR primers that detected the wild-type or mutant allele. (c) RNA was isolated from T cells of wild-type, miR-449a+/+, and miR-449a−/− mice, and the expression of miR-449a was evaluated by real-time PCR. Expression relative to that of G3pdh is shown. The data are shown as means ± SDs. **p < 0.01. (d) Total RNA was isolated from the indicated organs, and the expression of miR-449a in each organ was evaluated by real-time PCR. Expression relative to G3pdh is shown. The data are representative of three independent experiments.
miR-449a deficiency did not affect immune cell development or differentiation, which are regulated by Notch signaling. Promotion of colon cancer after AOM/DSS treatment in miR-449a−/− mice. Because previous reports have shown that miR-449a is involved in tumorigenesis in prostate, breast, lung, and gastric cancers and because miR-449a is expressed in the colon, we next sought to assess the roles of miR-449a in the tumorigenesis of colon cancer. We first treated wild-type or miR-449a−/− mice with 2% DSS and monitored body weight loss. There were no significant differences in body weight between the two groups, although body weight loss in miR-449a−/− mice tended to be more abundant than that in wild-type mice (Fig. 4a). We then treated wild-type and miR-449a−/− mice with AOM and 2% DSS to induce colon cancer (Fig. 4b) and measured the numbers of mice with tumors, tumor numbers in each mouse, and sizes of tumors at 6, 12, and 18 weeks after AOM treatment. Tumor incidence was 100%, and polyp numbers were comparable in both experimental groups and controls (data not shown). Tumors were found in both wild-type and miR-449a−/− mice at 6 weeks after AOM treatment, and miR-449a−/− mice had significantly more tumors than wild-type mice at 12 and 18 weeks after AOM treatment (Fig. 4c). The sizes of tumors were also larger in miR-449a−/− mice at 18 weeks after AOM treatment, with approximately 50% of miR-449a−/− mice developing tumors larger than 12 mm in diameter (Fig. 4d).

To characterize the nature of the deregulated carcinogenic signals in miR-449a−/− mice, we examined colonic epithelial proliferation in the tumorigenic process in the intestine. Ki67 was mainly expressed in the basal region of the colon in wild-type and miR-449a−/− mice (Fig. 4e). The number of intestinal epithelial cells expressing Ki67 was much larger in miR-449a−/− mice than in wild-type mice (Fig. 4e), demonstrating the increased proliferation of intestinal epithelial cells in miR-449a−/− mice after AOM/DSS treatment. Taken together, these data suggested that miR-449a acted as a tumor suppressor in colon cancer.

Disease-free survival in colon cancer was much longer in patients with higher miR-449a expression. The results obtained in miR-449a−/− mice led us to look for a link between miR-449a expression and pathology or prognosis in patients with colon cancer. We compared the expression of miR-449a in the intact colon and colon cancer in 76 patients with colon cancer. The expression level of miR-449a was similar in colon cancer tissue and normal colon tissue (Fig. 5a). We next assessed the association of pathological findings and expression...
level of miR-449a in colon cancer tissue. Higher expression of miR-449a was correlated positively with depth, differentiation, and size of colon cancer, but was not correlated with lymphatic invasion, venous invasion, or lymph node metastasis (Fig. 5b). We also compared the expression level of miR-449a and the prognosis of patients with colon cancer. Overall survival was not affected by miR-449a expression; however, disease-free survival was much longer in patients with higher miR-449a expression (Fig. 5c). Taken together, these results strongly suggested that there was a negative correlation between miR-449a expression and the severity of human colon cancer.

Expression of Mlh1 was positively correlated with miR-449a. Next, we compared the mRNA expression in the upper and lower colons of wild-type and miR-449a−/− mice to identify miR-449a target genes relevant to colon tumorigenesis (Fig. 6a). Among 16 known genes associated with colon cancer, Mlh1 was downregulated in both the upper and lower colons in miR-449a−/− mice compared with those from wild-type mice. The downregulation of Mlh1 was confirmed by real-time PCR (Fig. 6a). Another 15 genes were not significantly altered by deleting miR-449a.

MLH1 is a known tumor suppressor in colon cancer. Therefore, we evaluated whether there was an association between MLH1 and miR-449a expression in patients with colon cancer. The expression of MLH1 was positively correlated with miR-449a expression in 72 patients with colon cancer (Fig. 6b), suggesting that miR-449a played a role in susceptibility to colon cancer through controlling MLH1 expression.
In this study, we searched for miRNAs that were regulated by Notch signaling and found that miR-449a was indirectly upregulated by Notch signaling. Although Notch signaling is involved in the development or differentiation of various immune cells, miR-449a−/− mice do not show any defects in immune cell development, a process regulated by Notch signaling. Notably, miR-449a−/− mice showed higher susceptibility to AOM/DSS-induced colon tumorigenesis than wild-type mice, and the expression level of miR-449a was inversely correlated with disease severity, including disease-free survival, in patients with colon cancer. Moreover, miR-449a-deficient cells expressed lower levels of Mlh1 than control mice, and the expression of miR-449a was positively correlated with MLH1 in cancerous tissues from patients with colon cancer. These data suggested that miR-449a functioned to suppress colon tumorigenesis, at least partly through regulating MLH1 expression, and highlighted miR-449a as a therapeutic target and prognostic marker in the treatment of colon cancer.

The expression of miR-449a is frequently decreased in malignant tumors, including gastric and bladder cancer. In these cancers, miR-449a may inhibit cell growth or induce senescence and apoptosis by activating the p53 pathway. One recent paper reported that miR-449a was downregulated, while STAB2 expression was upregulated in patients with colorectal cancer. However, it is still unclear whether miR-449a is directly associated with tumorigenesis of various types of cancer because of a lack of data from in vivo models; accordingly, we have established miR-449a−/− mice. In these mice, AOM/DSS treatment increased the incidence of colon cancer and the rate of Ki-67-positive intestinal epithelial cells compared with those in control mice, directly demonstrating that miR-449a suppressed colon tumorigenesis and intestinal epithelial cell proliferation. Additionally, our microarray analysis showed that Mlh1 expression was lower in miR-449a−/− mice than in wild-type mice. MLH1 forms a heterodimer with PMS2 and functions to correct small errors involving mispaired nucleotides during DNA replication. MLH1 is frequently mutated in patients with colon cancer. Furthermore, Mlh1 deficiency in mice also accelerates colon carcinogenesis when combined with inflammation. Notably, miR-449a expression...
was positively correlated with MLH1 in patients with colon cancer in our present study. Thus, although it was unclear how miR-449a affected Mlh1 expression, our data suggested that miR-449a-mediated upregulation of MLH1 regulated the initiation or progression of colon cancer. The associations of miR-449a and MLH1 in terms of tumorigenesis should be analyzed by overexpressing Mlh1 in miR-449a-deficient mice, and it is also essential to assess how miR-449a regulates MLH1 expression. In addition, the miR-449 cluster contains sequences and secondary structures similar to those of the miR-34 family, which was found to be a p53-responsive gene cluster\(^35, 36\). miR-34 targets the histone deacetylase SIRT1\(^37\), leading to the accumulation of acetylated and therefore highly active p53. Moreover, miR-34 also downregulates several cyclin-dependent kinases, cyclins, and E2Fs\(^38, 39\), leading to cell cycle arrest. Therefore, the miR-449 cluster, including miR-449a, may undergo similar regulatory processes, contributing to the suppression of colon tumorigenesis by miR-449a.

Previous papers have reported that overexpression of miR-449a reduced Notch signaling\(^40\) and that blocking of miR-449-binding sites of endogenous human Notch1 or frog Dll1 strongly repressed multiciliogenesis\(^41\). Furthermore, miR-449a reduces cancer cell survival by directly downregulating Notch1\(^142\). These data indicated that miR-449a could suppress Notch signaling. However, our results demonstrated that stimulation of T cell hybridomas with overexpression of the intracellular domain of Notch1 upregulated miR-449a. These data were supported by low miR-449a expression in Rbpj-deficient T cells. However, we did not detect any defects in the development of T cells, marginal zone B cells, or splenic CD8\(^+\) dendritic cells, all of which are regulated by Notch signaling, in miR-449a\(^−/−\) mice. Therefore, miR-449a was not involved in Notch-mediated immune cell development. Notch signaling has also been implicated in tumorigenesis in many cancers, including colon cancer, due to the induction of prosurvival signaling in colonic epithelial cells\(^43\). Therefore, given the upregulation of miR-449a by Notch, Notch-mediated miR-449a expression in the colon may function as a self-guarding signal to suppress Notch-induced expression of tumorigenesis genes in the same cells.

In summary, our results revealed that the expression of miR-449a was inversely correlated with disease-free survival in colon cancer, and deficiency of miR-449a in mice increased susceptibility to AOM/DSS-induced colon cancer. In a previous study, miR-449a expression in carcinoma tissues was found to be inversely correlated with the levels of serum carcinoembryonic antigen\(^18\), supporting our present findings. With the goal of developing novel therapeutic strategies, stimulators of miR-449a may have beneficial effects on suppressing colon cancer and
identification of target genes for miR-449a may yield novel target molecules to suppress colon cancer. Our present data also highlighted miR-449a not only as a therapeutic target of colon cancer but also as a prognosis marker. Furthermore, previous papers have demonstrated that the expression of miR-449a is associated with progression of lung, gastric, and bladder cancers; thus, our miR-449a-deficient mouse model may be a useful tool to address the contribution of miR-449a to tumorigenesis in these cancers.

Methods

Mice. Six- to 8-week-old C57BL/6 mice were purchased from Japan SLC (Hamamatsu, Japan). Rbpj\(^{\text{floox}/\text{floox}}\) crossed with CD4-Cre transgenic mice were previously reported\(^{13,44}\). All mice were maintained under specific pathogen-free conditions in the animal facilities at Tokushima University, and all animal experiments were approved by the animal research committee of Tokushima University and performed in accordance with our institution’s guidelines for animal care and use.

Establishment of miR-449a\(^{\text{−/−}}\) mice. Murine genomic DNA of miR-449a was cloned by PCR. A 377b genomic miR-449a (restriction enzyme: NotI, EcoRV) fragment was replaced with a neo resistance gene cassette. E14 embryonic stem cells (1 × 10\(^5\)) were electroporated with 20μg of linearized targeting vector. G418-resistant colonies were obtained after 10 days. PCR screening for homologous recombination was carried out with a primer specific for the neo resistance gene (5′-CTATCCAGCACATGGTTGGG-3′) and an outside primer specific for miR-449a (5′-CTGTTCGCGTGAATCTAAAGG-3′) upstream of the construct. Homologous recombination was subsequently confirmed by BclI digestion of genomic DNA and hybridization with specific probes. Germline transmission of the miR-449a mutation was confirmed by Southern blot analysis.

Flow cytometry. Cells from the thymus and single-cell suspensions from the spleen or lymph nodes were stained with combinations of following antibodies: anti-mouse CD8α (53–6.7), anti-CD4 (145-2C11), anti-CD11b (M1/70), and anti-Gr1 (RB6-8C5), from Tonbo Biosciences (San Diego, CA, USA); anti-CD4 (GK1.5) and anti-B220 (RA3-6B2) from BD Biosciences (Franklin Lakes, NJ, USA); and anti-TCR\(^{\gamma\delta}\) (GL-3), anti-F4/80 (BM8), and anti-CD21 (8D9), and anti-CD23 (B3B4), from eBiosciences (San Diego, CA, USA). All samples were resuspended in phosphate-buffered saline (PBS) staining buffer containing 2% fetal bovine serum and 0.01% NaN\(_3\) and pre-incubated for 15 min at 4°C with 2.4G2 supernatant to block the Fc receptor. Samples were then washed and stained with specific mAbs for 20 min at 4°C. Data were collected on a FACSChanto II (BD Biosciences) and analyzed using FACS Diva (BD Biosciences) or FlowJo (Tree Star, OR, USA) software.

Purification of T cells. Total T cells from the spleens of C56BL/6 and Rbpj\(^{\text{floox}/\text{floox}}\) mice crossed with CD4-Cre transgenic mice were purified with a T-cell isolation kit (Miltenyi Biotech, Gladbach, Germany). Total RNA was isolated with RNeasy (Qiagen, Hilden, Germany).

Microarray. The intracellular domain of mouse Notch1 was cloned into the pKD004 retrovirus vector encoding green fluorescent protein (GFP)\(^{35}\). The vectors were transfected with Plat-E cells\(^{46}\) with GeneJuice (Merck Millipore, Darmstadt, Germany), and supernatants were collected 2 days after transfection. DO.11.10 cells were infected with retrovirus by centrifuging cells at 2600 rpm for 90 min. The miRNAs were collected using a High Pure miRNA isolation kit (Roche). One hundred nanograms of miRNA was used for the RNA probe. Microarray analyses were performed on a Mouse miRNA microarray 8 × 15 K miRBase 12.0 (Agilent).

Total RNA was isolated from the colons of mice using a Relia RNA Cell Miniprep System (Promega, Madison, WI, USA), and RNA quality was assessed by analysis with an Agilent 2100 BioAnalyzer. Thirty nanograms of RNA was used for the RNA probe. Probe preparation and microarray analyses were performed on a Whole Mouse Genome OligoDNA microarray kit ver2.0 44 K (Agilent Technologies). The resulting data were normalized with GeneSpring (Agilent Technologies) software. Genes showing at least 3.0-fold changes in expression (p < 0.05) between groups were considered to be differentially expressed.

AOM/DSS treatment. Mice were injected intraperitoneally with 10 mg/kg AOM (Sigma, St. Louis, MO, USA). Seven days later, 2% dextran sodium sulfate (ICN, MW 5,000 kDa) was given in the drinking water for 7 days. Body weights were measured every day after DSS treatment.

Histological studies. Mouse colon tissues were fixed in 10% formalin neutral buffer solution (Wako) and then embedded in paraffin. Paraffin-embedded sections were cut to 5 μm thickness and stained with hematoxylin and eosin solution. Paraffin-embedded sections were then stained with anti-Ki67 antibodies (D3B5; Cell Signaling Technology) followed by horseradish peroxidase (HRP)-labeled anti-rabbit antibodies.

Real-time PCR. Total RNA was extracted using RNeasy Plus Mini Kits (Qiagen, Valencia, CA, USA), and cDNA was synthesized using an Omniscript RT Kit (Qiagen). Gene expression was analyzed by qPCR on a Step-One RT PCR system (Applied Biosystems) using SBYR green incorporation. All genes were normalized to Hprt, and relative expression was calculated using the ΔΔCT method. All primer pairs were validated for amplification efficiency. The following primers were used: Mlh1, Fwd-5′-TATCGGACATAGCGGCTGGG-3′; Rev-5′-TTGACGTCCACGTTGAGG-3′; Hprt, Fwd-5′-AGCCTAAGATGAGCGCAAGT-3′, Rev-5′-TTACTAGGCAGATGGCCACA-3′.

Chromatin immunoprecipitation. Cells were incubated with 1% formaldehyde for 5 min at 4°C, and the crosslinked chromatin was then sonicated to shear chromatin fragments (200–1000 bp). The sonicated chromatin was immunoprecipitated with anti-Rbpj antibodies (RbpSUH, Cell Signaling Technology), and
the negative control was immunoprecipitated with control antibodies. The immunocomplexes were recovered using Dynabeads Protein G (Invitrogen). After treatment with proteinase K, DNA was purified by phenol/chloroform extraction. Real-time PCR was performed to quantify Rbpj-binding miR-449a promoter fragments using the following primers: miR-449a (450 bp), forward, 5′-GATGCCTAGGAGCTAATGAC-3′ and reverse, 5′-GCCACATAAACCTCTTCTCCT-3′; miR-449b (11 kb), forward, 5′-CAACGGTAGTGTACGTTGTG-3′ and reverse, 5′-CAGCTAGGCTCCATCTCCATA-3′; Cnot3, forward, 5′-CAAGACATGGTAGCATC-3′ and reverse, 5′-TGGTATCTAACCCTCCTCAAT-3′; and Dtx1, forward, 5′-CACACACCCCTCTCGAGTC-3′ and reverse, 5′-CAAGGGAGAGTGTCGATGC-3′.

Tumor samples. Pairs of primary colon tumor tissues and adjacent non-tumor tissues were collected from 80 patients recruited from Tokushima University. Detailed background information for each tissue donor, including age, sex, clinical stage, tumor location, and survival time after diagnosis, was collected. The patients were ordered based on miR-449a expression levels, and miR-449a low and high groups were designated based on the median value for all patients. This study was approved by the Institutional Review Board for Human Subject Research at Tokushima University. All participants provided written informed consent to participate in this study, and the consent procedure was approved by the Institutional Review Board. All methods were performed in accordance with the relevant guidelines and regulations.

Statistics. The means ± standard deviations (SDs) were calculated for all parameters determined. Statistical significance was evaluated using one-way analysis of variance (ANOVA), followed by Fisher’s protected least significant difference test. P values less than 0.05 were considered statistically significant. For all experiments, the significance of differences between groups was calculated using the Mann-Whitney U test for unpaired data. The Kaplan-Meier method and log rank test were used to estimate overall survival. The correlation coefficients were analyzed by standard Pearson correlation analysis.

References


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Author Contributions

M.N., K.N., D.I., and K.Y. designed the project and planned the experiments. K.N. and J.N. generated the miR-449a-deficient mice. S.W.L. and H.J.P. performed all experiments and quantifications. C.I., S.T., M.S., S.N., and M.N., K.N., D.I., and K.Y. designed the project and planned the experiments. K.N. and J.N. generated the miR-449a-deficient mice. S.W.L. and H.J.P. performed all experiments and quantifications. C.I., S.T., M.S., S.N., and Y.M. analyzed and discussed the results. M.N. and K.Y. wrote the manuscript.

Additional Information

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