Peyer’s patches are nodules that play a central role in intestinal immunity. Few studies demonstrate the relationship between the number of Peyer’s patches and intestinal polyps. Here we identify a statistically significant inverse correlation between the quantity of Peyer’s patches and the development of intestinal polyps in Apc^Min/+ mice, which are a useful model to clarify the role of Peyer’s patches in intestinal tumorigenesis. Using this model, we increased the number of Peyer’s patches using 0.1% and 1% corn husk arabinoxylan through feed. Intestinal polyp formation increased the number of Peyer’s patches using 0.1% and 1% corn husk arabinoxylan (CHAX) through feed. Intestinal polyp formation increased the number of Peyer’s patches using 0.1% and 1% corn husk arabinoxylan (CHAX) as an adjuvant. We examined the effects of PP development on intestinal polyps following ingestion of CHAX in Apc^Min/+ and Aly^−/−Apc^Min/+ mice, which have no PP in the intestine. Furthermore, we tried to clarify the molecular mechanism for the prevention of polyp formation in the mice.

**Materials and Methods**

**Mice and feed.** Wild type (WT; C57B1/6J) mice and alymphoplasia knockout (Aly^−/−, C57B1/6J) mice, were obtained from Clea Japan, Inc. (Tokyo, Japan). The Apc^Min/+ (C57B1/6J) mice were supplied by Jackson Laboratories (Bar Harbor, Maine, State). The mice were maintained under specific pathogen-free conditions at the Animal Center of Nagasaki International University. (Nagasaki, Japan) Female Apc^Min/+ and male Aly^−/− mice were mated to obtain Aly^−/−Apc^Min/+ mice. The mouse feed was compounded with two different doses of CHAX (NIHON SHOKUKIN KAKO CO., LTD, Tokyo, Japan): 0.1% and 1% (mixed (i.e., 0.2 and 2 g/kg (body weight)/day each). Preparation of CHAX was described elsewhere. According to standard dietary intake guidelines provided by The Ministry of Health, Labour and Welfare, we calculated the appropriate dose for mice. We analyzed 10–12 mice in each feeding group: CHAX-free, 0.1, and 1% CHAX for each genotype (WT, Apc^Min/+ and Aly^−/−Apc^Min/+). Approximately 100 mice were used in the ingestion experiment. Mice were sacrificed at 12, 24, and 32 weeks of age for the analysis of intestinal polyps and PP. All animal experiments were conducted according to the Guidelines for Animal Experiments in the Faculty of Pharmaceutical Sciences, Nagasaki International University.

**Genotyping.** Mouse tails were genotyped using a Genomic DNA Purification Kit (QIAGEN, Hilden, Germany) according to the manufacturer’s protocol. Allele-specific polymerase chain reaction (PCR) primers for the Apc gene were produced according to previously published methods. Forward and reverse primers for Apc^Min/+ mice had complimentary annealing temperatures and a product length of 302 base pairs. Genomic DNA was used as a template for PCR to amplify fragments containing forward Aly Wild 5’-CTGACATCCCGAGCTACTTCAACG, forward Aly Mutant 5’-CTGACATCCCGAGCTACTTCAACA, and reverse 5’-GCTTAGGATCGGCCATTTTCTTCC primer using the AmpliTaq Gold 360 Master Mix (Applied Biosystems, California, CA). Touchdown PCR consisted of one cycle of 94°C for 9 min for the initial denaturation step. This was followed by 10 cycles each of denaturation at 94°C for 1 min, varying annealing conditions for 1 min, and extension at 72°C for 1 min. Annealing temperatures for the touchdown portion were as follows: starting annealing temperatures of 72°C decreasing by 0.5°C decrements.
per PCR cycle down to 67°C. Further 25 cycles consisted of the following: 94°C for 1 min, 68°C for 1 min, and 72°C for 1 min.

Counts of intestinal polyps and statistical analysis. Intestinal polyps and statistical analysis were examined according to previously published methods.(16) All statistical analyses were carried out using the GraphPad Prism 5 program. (GraphPad Software Inc., San Diego, CA) The correlation coefficient of the number of PP and intestinal polyps was analyzed using Pearson’s correlation coefficient. We analyzed two-group by using Mann–Whitney t test, and three or more groups were analyzed simultaneously by using Dunnett comparison test.

Cytopene array. Serum was collected from the heart when mice were dissected. We pooled serum samples from four WT mice fed with or without CHAX, respectively. Similarly, we pooled serum samples from 7-ApcMin/− or 4-Aly−/ApcMin/− mice and we analyzed using Multiplex Suspension Array (Genetic Lab Corp., Hokkaido, Japan) The total number of samples was six, and the experiment was repeated two times using new samples.

Enzyme-linked immunosorbent assay (ELISA). All serum samples were stored in a –80°C deep freezer until use. The levels of interleukin (IL)-17 were measured by IL-17 Mouse ELISA Kit (Abcam, Cambridge, UK), according to the manufacturer’s protocol. We measured fluorescence using a Model 680 Microplate Reader (Bio-Rad, Hercules, CA).

Quantitative real-time PCR analysis. All tissue samples from intestinal polyps of mice fed with or without CHAX were rapidly soaked in RNA later solution (Qiagen). Total RNA was isolated from tissue using the RNeasy Mini Kit (Qiagen). Complementary DNA (from reverse transcribed total RNA) and real-time PCR were examined according to previously published methods.(17) Primers for mouse cyclin D1 (5’primer-CCATGG-AACACCGCTCTTG and 3’primer-CCGTCAGGGTAGTCCATGCC) and glyceraldehyde 3-phosphate dehydrogenase (GAPDH; 5’primer-TGTCAGCAATGCTGCA and 3’primer-TTACTCCTTGGAGGCCATGT) were employed. The expression levels of cyclin D1 were normalized based on GAPDH levels.

Histology and immunohistochemistry. Small intestines were fixed, embedded, and sectioned as Swiss rolls for further immunohistochemical examination with the avidin–biotin complex immunoperoxidase technique and monoclonal mouse anti-β-catenin antibody (Ab; BD Transduction Laboratories, Franklin Lakes, NJ) at 100× dilution. The secondary Ab, biotinylated antibody (Vector Laboratories, California, CA), was used at a 200× dilution. Staining was performed using avidin–biotin reagents (Vectastain ABC reagents; Vector Laboratories), 3,3′-diaminobenzidine and hydrogen peroxide, and the sections were counterstained with hematoxylin to facilitate orientation. As a negative control, consecutive sections were immunostained without exposure to the primary Ab.

Reporter assays. Caco-2 cells were plated at a concentration of 5 × 104 cells/well in 12-well plates and cultivated for 24 h. Furthermore, cells were transfected with pTOP or pTOP-Flash (Merck Millipore, Darmstadt, Germany) luciferase reporter plasmids (1.0 μg each) and Simian virus 40 (SV40)-RenillaLuc (pRL-SV40; 15:1 ratio) for 24 h. All samples were normalized for transfection efficiency using the SV40-RenillaLuc expression plasmid (Promega, Madison, WI) as a transfection control. Cells were harvested and lysed in 200 μl of lysis buffer 48 h after transfection. We measured four points using 20 μl of sample, and luciferase and Renilla luciferase activity were assayed using the Bright-Glo Luciferase Assay System and Renilla-Glo Luciferase Assay System respectively (Promega) using a GENios Microplate Reader (Tecan, Männedorf, Switzerland). Transfections were repeated three times.

After correction of measurements, we calculated the average of four measurements, and transcription factor/lymphoid enhancer-binding factor 4 (TCF/LEF) activity was calculated by TOP divided by FOP. The relative transcriptional activity was calculated for the aforementioned value was divided by the measurement of the control cell.

Results

The number of PP correlates significantly with the number of intestinal polyps in ApcMin/+ mice over 24 weeks in age. We counted the number of intestinal polyps and PP in ApcMin/+ mice >24 weeks in age. Mice that developed many intestinal polyps tended to have been fewer PP in the intestine. A statistically significant inverse relationship existed between the number of intestinal polyps and PP (r = −0.62, n = 20, p = 0.0035; Fig. 1). In addition, the amount of PP in the intestines decreased in older mice groups (Fig. 2). The quantity of PP in the intestines was significantly reduced in ApcMin/+ mice >21 weeks of age compared with 12-week-old ApcMin/+ mice. Moreover, the number of polyps significantly increased in older mice groups.

CHAX increases the number of PP in ApcMin/+ mice. It remains unclear whether or not PP suppresses polyp formation, and what factors affect the development of each condition. However,
Furthermore, we fed 0.1% and 1% CHAX to 1% concentrations of CHAX as optimal doses for PP formation. CHAX has been reported to increase the number of PP; therefore, we used this adjuvant to elucidate the effects of PP on polyp formation.

Mice were fed a diet of 0.1, 0.5, 1, or 4% CHAX to determine the optimal concentration that leads to the greatest volume of PP in WT mice. There were 7.7 ± 0.5 (n = 14) PP in 12-week-old WT mice administered 0.1% CHAX. The number significantly increased to 8.9 ± 0.5 when the mice were fed 0.5% CHAX and higher. (*p = 0.0246, Mann–Whitney test) We chose the 0.1% and 1% CHAX concentrations of CHAX as optimal doses for PP formation. Furthermore, we fed 0.1% and 1% CHAX to ApcMin/+ mice for 4 weeks. The number of PP significantly increased in ApcMin/+ mice compared to WT mice (Table 1). The number of intestinal polyps in 12-week-old WT mice was significantly decreased in ApcMin/+ mice fed or not fed CHAX based on the number of PP in the small intestine. IL-17 levels gradually decreased with increasing number of PP (Fig. 3).

**CHAX leads to a decrease in polyp development through the β-catenin pathway.** Furthermore, we examined immuno-histochemical staining of paraffin-embedded intestinal polyps for β-catenin and proliferation cell nuclear antigen (PCNA). Although β-catenin is well known for accumulating in the nucleus of intestinal polyps in ApcMin/+ mice, this accretion was not observed in mice fed a 1% CHAX diet (Fig. 4a and b). In addition, the staining intensity of PCNA was weaker in mice fed 1% CHAX (Fig. 4c and d). Furthermore, we measured the messenger RNA expression level of cyclin D1, which is downstream of Wnt signaling, in intestinal polyps by quantitative real-time PCR. We found that the expression of cyclin D1 tended to decrease as CHAX concentration increased (Fig. 5).

**IL-17 promotes TCF/LEF transcriptional activation in vitro.** We examined TCF/LEF-dependent transcriptional activity after 24-h treatment with IL-17 in Caco-2 cell transfected with pTOP-Flash or pFOP-Flash and SV40-RenillaLuc, and differentiated with 5 mM sodium butyrate (Fig. 6). Caco-2 cells are an established intestinal cell model. We found that the transcriptional activity of TCF/LEF was increased by IL-17 treatment in Caco-2 cells. These results suggest that IL-17 promotes TCF/LEF transcriptional activation in vitro.

| Table 1. The number of PP in ApcMin/+ mice fed CHAX |
|----------------------------------|------------------|------------------|------------------|------------------|
| **CHAX dosage** (%)             | **Small intestine** |                  |                  |                  |
|                                 | **Proximal**     | **Middle**       | **Distal**       | **Total**        |
| 0                               | 2.8 ± 0.3        | 2.1 ± 0.2        | 3.4 ± 0.2        | 7.3 ± 0.3        |
| 0.1                             | 3.0 ± 0.2        | 2.3 ± 0.2*       | 3.9 ± 0.3        | 9.3 ± 0.3**      |
| 1                               | 2.9 ± 0.2        | 2.8 ± 0.3**      | 3.8 ± 0.3        | 9.5 ± 0.2***     |

Values are mean ± SEM (n = 12/group). *p<0.05, **p<0.01, ***p<0.001; Mann–Whitney t test.

| Table 2. The number of intestinal polyps in 12-week-old ApcMin/+ mice |
|----------------------------------|------------------|------------------|------------------|------------------|
| **CHAX dosage** (%)             | **Small intestine** |                  | **Large intestine** | **Total**        |
|                                 | **Proximal**     | **Middle**       | **Distal**       | **Total**        |
| 0                               | 3.9 ± 0.7        | 22.3 ± 2.2       | 27.2 ± 2.7       | 53.4 ± 4.4       |
| 0.1                             | 2.1 ± 0.5        | 11.0 ± 1.6**     | 16.2 ± 2.3*      | 29.3 ± 3.8**     |
| 1                               | 3.0 ± 0.4        | 10.0 ± 1.7**     | 17.9 ± 2.5*      | 30.9 ± 3.3**     |

Values are mean ± SEM (n = 12/group). *p<0.05, **p<0.01, ***p<0.001; Mann–Whitney t test.

| Table 3. The number of intestinal polyps in 12-week-old Aly−/−ApcMin/+ mice (negative control group) |
|----------------------------------|------------------|------------------|------------------|------------------|
| **CHAX dosage** (%)             | **Small intestine** |                  | **Large intestine** | **Total**        |
|                                 | **Proximal**     | **Middle**       | **Distal**       | **Total**        |
| 0                               | 4.3 ± 2.9        | 13.7 ± 1.6       | 11.3 ± 1.4       | 29.3 ± 2.9       |
| 0.1                             | 2.0 ± 0.3        | 8.4 ± 1.6        | 10.6 ± 2.8       | 21.1 ± 3.7       |
| 1                               | 3.4 ± 0.9        | 11.1 ± 3.2       | 11.3 ± 2.7       | 25.8 ± 5.9       |

Values are mean ± SEM (n = 10/group).

| Table 4. Serum cytokine levels in WT, ApcMin/+ and Aly−/−ApcMin/+ mice fed with or without CHAX |
|----------------------------------|------------------|------------------|------------------|------------------|
|                                 | **WT mice**     | **Rate of change** | **ApcMin/+ mice** | **Rate of change** |
|                                 | − (without CHAX) | + (with CHAX)    | − (without CHAX)  | + (with CHAX)    |
| IL-1β (pg/ml)                   | <0.64           | <0.64            | 4.38            | <0.64            |
| IL-5 (pg/ml)                    | 10.68           | 4.37             | 2.44            | 8.58             |
| IL-10 (pg/ml)                   | 3.47            | 7.55             | 2.18            | 4.87             |
| IL-17 (pg/ml)                   | 3.86            | 1.05             | 3.68            | 3.24             |

Values are the averages of extracted cytokines from double experiments of the cytokine array.

CHAX has been reported to increase the number of PP; therefore, we used this adjuvant to elucidate the effects of PP on polyp formation.
lished cell line derived from colorectal tumors with loss of APC gene function.

There was no significant increase in TCF/LEF-dependent transcriptional activity with 0.2 ng/ml of IL-17, but this was observed at 0.5 ng/ml ($p<0.001$, Dunnett’s multiple comparison test). TCF/LEF activity was activated by IL-17 in a dose-dependent manner. The control, 0.2, and 0.5 ng/ml IL-17 values for TOP/FOP were $1 \pm 0.15$, $1.13 \pm 0.03$, and $1.40 \pm 0.12$, respectively.

**Discussion**

With continuing observation, we noticed that the number of intestinal polyps in $Apc^{Min/+}$ mice varied. Usually, it is assumed that intestinal polyps appear in $Apc^{Min/+}$ mice at 8 weeks of age, and we confirmed several polyps at this age. Furthermore, it is known that the number of polyps will increase with age; the life span of $Apc^{Min/+}$ mouse is approximately 24 weeks because of bleeding caused anemia from intestinal polyps and other factors.\(^{18–20}\) We
Our findings show that intestinal polyp formation in Aly−/−ApcMuMin−/− mice is lower in comparison with ApcMuMin−/− mice. Aly−/− mice are created by a point mutation in the tumor necrosis factor receptor associated factor (TRAF) binding domain, which is in the nuclear factor-xB (NF-xB)-inducing kinase (Nik) gene.10 NF-xB is involved in many physiological phenomena such as acute and a chronic inflammation reactions, cell growth, and apoptosis. Furthermore, in many cases, NF-xB is activated in malignant tumors. Therefore, the decreased number of intestinal polyps in Aly−/−ApcMuMin−/− mice compared to ApcMuMin−/− mice may be because of the TRAF family, which have tumor promoting properties, not binding Nik.21,22

We examined changes in immunoassociated cytokines using cytokine array, because PP are critical organs in intestinal immunity and may be involved in intestinal polyp formation. In this study, we focused on IL-17, which increased the number of PP in intestine; the levels of IL-1β remained unchanged in WT mice. IL-5 and IL-10 were excluded from analysis because the changes in concentration were only observed in Aly−/−ApcMuMin−/− mice that do not develop PP. IL-17 is known to be produced by T helper 17 cells (Th17) cells and plays an important role in allergic responses, such as contact hypersensitivity, delayed hypersensitivity, and airway hyper-reactivity, as determined by IL-17 gene knockout mouse studies.23 We found an inverse correlation existed between IL-17 concentration and the number of PP, suggesting a relationship between IL-17 and the number of PP. Although the reason of this phenomenon is not known in detail, Hirota et al.24 reported that expression of IL-17 decreases when Th17 cells induce B cell formation in PP.

Short chain fatty acids, such as sodium butyrate, are mainly produced by intestinal bacteria, and are largely involved in differentiation and nutrition of intestinal epithelial cells.27–29 Our cytokine array revealed that IL-17 levels were low (Table 4). Moreover, our results demonstrate a role for IL-17 in the enhancement of Wnt signal in the intestine. There are several reports highlighting that IL-17 is involved in intestinal polyp formation in ApcMuMin−/− mice or it can aggravate inflammatory bowel disease.30–32 These findings support the hypothesis that the number of intestinal polyps is decreased due to an increase in PP in ApcMuMin−/− mice fed CHAX. Moreover, reduced expression of IL-17 in serum possibly causes destabilization of Wnt signaling. In the future, we propose that quantification of intestinal PP will be a useful tool for the prevention of tumorigenesis.

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Conflict of interest

No potential conflicts of interest were disclosed.

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Fig. 5. Quantitative real-time PCR of cyclin D1. The p value is * = 0.0426 (Mann–Whitney t test). The number of polyps used: n = 6–8, respectively. cyclin D1 mRNA levels were corrected using GAPDH as a control.

Fig. 6. TCF/LEF transcripational activity was induced by IL-17 in Caco-2 cells. The cells were stimulated with 5 mM sodium butyrate. Data are representive of at least there independent experiments. The TCF/LEF transcripational activity was significantly increased with 0.5 ng/ml of IL-17 (*1 = p<0.001, Dunnett’s multiple comparison test).

noticed the presence of numerous PP when the mice have few polyps in the small intestine, we sought to examine the correlation between the number of polyps and PP. Previous reports have shown that the number of PP in the intestines is increased when mice are fed CHAX and/or short chain fructooligosaccharide diet.21,22 However, there were no reports that described this change in PP levels in detail.

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