Ecto-Nucleoside Triphosphate Diphosphohydrolase 7 Controls Th17 Cell Responses through Regulation of Luminal ATP in the Small Intestine

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Ecto-Nucleoside Triphosphate Diphosphohydrolase 7 Controls Th17 Cell Responses through Regulation of Luminal ATP in the Small Intestine


Extracellular ATP is released from live cells in controlled conditions, as well as dying cells in inflammatory conditions, and, thereby, regulates T cell responses, including Th17 cell induction. The level of extracellular ATP is closely regulated by ATP hydrolyzing enzymes, such as ecto-nucleoside triphosphate diphosphohydrolases (ENTPDases). ENTPDase1/CD39, which is expressed in immune cells, was shown to regulate immune responses by downregulating the ATP level. In this study, we analyzed the immunomodulatory function of ENTPDase7, which is preferentially expressed in epithelial cells in the small intestine. The targeted deletion of Entpd7 encoding ENTPDase7 in mice resulted in increased ATP levels in the small intestinal lumen. The number of Th17 cells was selectively increased in the small intestinal lamina propria in Entpd7−/− mice. Th17 cells were decreased by oral administration of antibiotics or the ATP antagonist in rodentium infected mice. Th17 cell responses were modulated in the small intestinal lamina propria of Entpd7−/− mice. Th17 cells were decreased by oral administration of antibiotics or the ATP antagonist in rodentium infected mice, indicating that commensal microbiota-dependent ATP release mediates the enhanced Th17 cell development in the small intestinal lamina propria of Entpd7−/− mice. In accordance with the increased number of small intestinal Th17 cells, Entpd7−/− mice were resistant to experimental autoimmune encephalomyelitis, which was associated with increased numbers of CD4+ T cells producing both IL-17 and IFN-γ. Taken together, these findings demonstrate that ENTPDase7 controls the luminal ATP level and, thereby, regulates Th17 cell development in the small intestine. The Journal of Immunology, 2013, 190: 000–000.

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Abbreviations used in this article: DC, dendritic cell; EAE, experimental autoimmune encephalomyelitis; EC, epithelial cell; ENTICase, ecto-nucleoside triphosphate diphosphohydrolase; MLN, mesenteric lymph node; MOG, myelin oligodendrocyte glycoprotein; oATP, oxidized ATP; SFB, segmented filamentous bacteria.

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expressed in immune cells, such as T cells, B cells, NK cells, DCs, and monocytes/macrophages (19, 20), was shown to possess anti-inflammatory activities through ATP hydrolysis. Indeed, severe inflammation was induced in mice lacking ENTPDase1/CD39 in several inflammatory models, including inflammatory bowel disease (21–24). Combinational activity of ENTPDases such as CD39 with CD73 ecto-5′-nucleotidase, which hydrolyzes AMP to adenosine, was also demonstrated in regulatory T cells and intestinal ECs (11, 20, 25). Thus, the immune-modulatory functions of ENTPDase1/CD39 have been well characterized. However, it remains unclear whether other ENTPDase family members are involved in the regulation of immune responses.

In this study, we analyzed the role of ENTPDase7, which was selectively expressed in ECs in the small intestine. Deletion of ENTPDase7 in mice resulted in increased ATP concentrations in the small intestinal lumen and increased numbers of IL-17–producing Th17 cells in the small intestinal lamina propria. Blockade of ATP action decreased the number of Th17 cells in the small intestine of ENTPDase7–deficient mice. In accordance with the increased Th17 cell number, ENTPDase7–deficient mice showed high resistance to the intestinal pathogen Citrobacter rodentium. These findings demonstrate that intestinal ECs participate in the regulation of Th17 cell responses by controlling intestinal ATP levels.

Materials and Methods

Real-time RT-PCR

RNA samples were prepared from various organs, epithelial layer, and lamina propria of C57BL/6 mice (CLEA Japan) using TRIzol reagent (Invitrogen), from single-cell suspensions using an RNaseq Mini Kit (QIAGEN), or from laser-microdissected tissue sections using an RNaseq Micro Kit (QIAGEN). Total RNA was reverse transcribed using Moleceny murine leukemia virus reverse transcriptase (Promega) and random primers (Toyobo) before treatment with RQ1 DNase I (Promega). cDNA was analyzed by real-time RT-PCR using GoTaq qPCR Master Mix (Promega) in an ABI 7300 real-time PCR system (Applied Biosystems). Values were then normalized to the expression of Gapdh, and the fold difference in expression relative to that of Gapdh is shown. The following primer sets were used: Entpd1, 5′-TGGTGCAGCAGTTAGAGGAA-3′ and 5′-GTC-CTTCACCTGTCCCGACTT-3′; Entpd7, 5′-CCCCTTACATCTCTGCAC-3′ and 5′-GTC-AAACCTCAAGCGCAAT-3′; Muc2, 5′-ACATACCTCTGGCGACCTTC-3′ and 5′-GAGGAAAGGCTACTGGTCT-3′; Kr7, 5′-ACGGCTGCT-GAGAATGATT-3′ and 5′-CTGGAAGGCTCTGGAGAAG-3′; and Gapdh, 5′-CCCTGGCCGATGACAAATG-3′ and 5′-TCTCCATTT-GCACTGGCA-3′.

Isolation of epithelium and lamina propria

Intestines were opened longitudinally, washed to remove fecal content, and incubated in PBS containing 30 mM EDTA for 5 min. Epithelial layer was peeled off from intestines and used as epithelium. For isolation of lamina propria, after removing the epithelial layer, fat tissue was also removed and overlaid on 2.5 ml 80% Percoll in a 15-ml Falcon tube. Percoll-gradient separation was performed by centrifugation at 780 × g for 20 min at 37°C. The intraepithelial lymphocytes were collected at the interface of the Percoll gradient and washed with RPMI 1640 containing 10% FBS. For isolation of lamina propria lymphocytes, intestines were opened, washed to remove fecal content, shaken in HBSS containing 5 mM EDTA for 20 min at 37°C. After filtration through nylon mesh, the EC fraction was washed with RPMI 1640 containing 4% FBS, resuspended in 5 ml 40% Percoll (GE Healthcare), and overlaid on 2.5 ml 80% Percoll in a 15-ml Falcon tube. Percoll-gradient separation was performed by centrifugation at 780 × g for 20 min at 37°C. The intraepithelial lymphocytes were collected at the interface of the Percoll gradient and washed with RPMI 1640 containing 10% FBS.

Intracellular cytokine staining

Intracellular expression of IL-17, IFN-γ, and IL-10 in CD4+ T cells was analyzed using a Cytosift/Cytoperm Kit Plus (with GolgiStop; BD Biosciences), according to the manufacturer’s instructions. In brief, lymphocytes obtained from the intestinal lamina propria, spleens, MLNs, or Peyer’s patches were incubated with 50 ng/ml PMA (Sigma), 5 μM calcium ionophore A23187 (Sigma), and GolgiStop at 37°C for 4 h. Surface staining was performed with anti–CD4–PerCP/Cy5.5 (BioLegend) for 20 min at 4°C, the cells were permeabilized with Cytosift/Cytoperm solution for 20 min at 4°C, and intracellular cytokine staining was performed with anti–IL-17A–Alexa Fluor 647 (BD Biosciences), anti–IL-10–PE (BD Biosciences), and anti–IFN-γ–FITC (BioLegend) for 20 min. For intracellular staining of Foxp3, cells were stained using the Foxp3 Staining Buffer set (eBiosciences).

Flow cytometry

The following Abs were used for flow cytometry: anti–CD4–PerCP/Cy5.5, anti–CD8α–Pacific Blue, anti–CD3–FITC, anti–TCRβ–PE, anti–CD8β–Alexa Fluor 647, and anti–CD4–PE/Cy7 (all from BioLegend); anti–B220–PE, anti–CD3–PE/Cy7, and anti–CD8–PE (all from BD Biosciences); and anti–CD8β–FITC (eBiosciences). Anti–Foxp3–Alexa Fluor 647 (eBiosciences) was also used, according to the manufacturer’s instructions. Data were acquired using a FACScanto II (BD Biosciences) and analyzed using FlowJo software (Tree Star).

Establishment of small intestinal EC lines

H-2Kb-tsA58–transgenic mice (26) were backcrossed to C57BL/6 mice for six generations. To establish the small intestinal EC lines from wild-type and Entpd7−/− mice, the mice were crossed with H-2Kb-tsA58–transgenic mice. Small intestinal ECs were isolated, as previously described (27), before incubation at 33°C. To confirm that they were intestinal ECs, a single-cell suspension was prepared and cytopsin onto the glass slides. After fixation, the cells were incubated with polyclonal anti-cytokeratin Ab (1:500; Dako) and then treated with a ChemMate EnVision kit (Dako). DAB (Dako) was used as a chromogen. Images were taken using a BZ-9000 fluorescence microscope (Keyence).

Measurement of ATP

Feces from individual mice were collected, weighed, and gently suspended in PBS containing 0.01% NaN3. After centrifugation, the supernatants were collected, and the levels of ATP were determined with a luciferin-
luciferase assay using the ATP assay kit (Toyo Ink), according to the manufacturer’s instructions. To analyze ATP levels in the small intestinal tissues, the small intestine was isolated and cut into quarters longitudinally. Each piece was weighed and lysed to measure ATP with a luciferin-luciferase assay. To analyze ATP levels in the EC lines, single-cell suspensions of the indicated cell lines were prepared. The cells were counted and lysed to measure ATP with a luciferin-luciferase assay. For determination of luminal ATP levels, the mice were fasted overnight and analyzed as described with 500 mM pentobarbital sodium (Dainippon Sumitomo Pharma). The peritoneal cavity was opened, and the small intestine was ligated with nylon threads at 1.5 and 4.5 cm distal from the Treitz ligament (for the proximal region of the small intestine) or at 3 and 6 cm proximal from the ileum end (for the distal region of the small intestine) to make a closed intestinal loop. A total of 300 ml PBS or 1.5 mM ATP solution was applied luminally with a 29-G needle. The luminal fluid was recovered 15 min later using a 29-G needle and suspended in PBS. After centrifugation, the supernatants were collected, and the levels of ATP were determined with a luciferin-luciferase assay.

Measurement of NTP hydrolyzing activity

NTP (ATP, GTP, UTP, and CTP) hydrolyzing activity was measured in crude membranes from wild-type and Entpd7−/− small intestinal ECs, as previously described (28). Briefly, ECs were homogenized; after removing nuclei, the crude membrane fraction was separated from the cytosol by centrifugation at 100,000 × g for 30 min. To assay for NTP hydrolyzing activity, the membrane fraction containing 10 μg total protein was suspended in reaction buffer (20 mM HEPES [pH 7.4], 120 mM NaCl, 5 mM KCl, 0.2 mM EDTA, 1 mM NaN₃, and 0.5 mM Na₃VO₄, with or without 5 mM CaCl₂). After incubation for 5 min at 37°C, 5 μl the reaction buffer containing 10 mM NTP was added and incubated for 30 min. NTP hydrolyzing activity was determined by measuring the inorganic phosphate, as described previously (28).

In vitro naive T cell differentiation

Naive T cells were grown for 4 d at 5 × 10⁶ cells/ml with plate-bound anti-CD3 (2 mg/ml) in DME supplemented with 10% FBS, penicillin, and streptomycin under Th17-polarizing conditions (2 ng/ml TGF-β, 20 ng/ml IL-6, 5 μg/ml anti-IFN-γ, 5 μg/ml anti-IL-4) or Th2 conditions (5 μg/ml anti-IFN-γ, 5 μg/ml anti-IL-4). Then, cells were incubated with 50 ng/ml anti-IL-4 (Sigma), 5 μM calcium ionophore A23187 (Sigma), and GolgiStop at 37°C for 4 h for flow cytometry analysis.

Treatment with antibiotics

Mice were given a combination of antibiotics containing 500 μg/ml vancomycin (Wako), 1 mg/ml metronidazole, 1 mg/ml ampicillin, and 1 mg/ ml neomycin sulfate (all from Nacalai Tesque) in drinking water from birth. Mice were assigned scores of 1 to 5 as follows: 0, no clinical signs; 1, paralyzation; 2, loss of coordinated movement; 3, both hind limbs paralyzed; 4, forelimbs paralyzed; and 5, moribund.

Statistical analysis

Differences between control and experimental groups were evaluated by the Student t test.

Results

Selective expression of Entpd7 in small intestinal epithelia

ENTPDase1/CD39 encoded by Entpd1 was shown to modulate inflammatory responses in addition to thrombopoiesis (24, 33, 34). Because the ENTPDase family consists of eight members, we analyzed tissue expression of Entpd gene family members. Entpd1 was preferentially expressed in lymphoid organs, such as the spleen and lymph nodes (Fig. 1A). Of the other Entpd genes, we focused on those that showed selective tissue-expression patterns. Entpd7 was highly expressed in the small intestine (Fig. 1B). The highest Entpd7 expression was observed in the proximal region of the small intestine, and its expression gradually decreased as the small intestines descended (Fig. 1C). We then analyzed expression of Entpd7 in the epithelial layers and lamina propria of the small intestine (Fig. 1D). Entpd7 was predominantly expressed in the ECs of the small intestine. We further analyzed which types of intestinal ECs (i.e., goblet cells or absorptive enterocytes) highly expressed Entpd7. Goblet cell–enriched, absorptive enterocyte–enriched, and lamina propria cell–enriched regions were isolated by laser microdissection, and expression of Entpd7 was analyzed (Fig. 1E). Entpd7 was highly expressed in absorptive enterocytes, as well as goblet cells characterized by high expression of Muc2.
Thus, Entpd7 is highly expressed in all types of ECs of the small intestine. Expression of Entpd7 in the small intestine was not altered in mice treated with oral antibiotics, indicating that Entpd7 expression is not influenced by microbiota (Fig. 1F).

To assess the physiological function of ENTPDase encoded by Entpd7, we generated Entpd7−/− mice by gene targeting (Supplemental Fig. 1A, 1B). Entpd7−/− mice were born at the normal Mendelian ratios and grew healthily until 16 wk of age (Supplemental Fig. 1C). Normal lymphocyte development was observed in Entpd7−/− mice (Supplemental Fig. 1D). The composition of lymphocytes in the small and large intestine was not altered in Entpd7−/− mice (Supplemental Fig. 2).

Elevated ATP level in the small intestinal lumen of Entpd7−/− mice

Because ENTPDase is an enzyme that hydrolyzes nucleoside triphosphates, and Entpd7 was selectively expressed in the small intestinal epithelia, we analyzed concentrations of ATP in the intestine. First, the small intestines were cut into four regions, and their lysates were analyzed for ATP concentration (Fig. 2A). The ATP level was not dramatically altered in any region of the small intestinal tissues between wild-type and Entpd7−/− mice. Because Entpd7 is highly expressed in ECs of the small intestine, we established intestinal EC lines from wild-type and Entpd7−/− mice using transgenic mice harboring a temperature-sensitive mutation of the SV40 large tumor Ag gene under the control of an IFN-γ-inducible H-2Kb promoter element to analyze ATP levels in the ECs (26, 35, 36). ECs from wild-type and Entpd7−/− mice expressed keratin proteins equally as well as Krt7 mRNA, indicating that these cells are ECs (Supplemental Fig. 3). Entpd7 was highly expressed in wild-type ECs but not in Entpd7−/− ECs (Supplemental Fig. 3). Intracellular ATP levels were not altered between wild-type and Entpd7−/− ECs (Fig. 2B). Fecal concentrations of ATP were not different in Entpd7−/− mice compared with wild-type mice (Fig. 2C). However, ATP levels in the luminal contents of the small intestine were substantially increased in Entpd7−/− mice (Fig. 2D). We then created a ligated intestinal loop model to analyze alterations in luminal ATP levels. The proximal regions of the small intestine were ligated to make a loop in wild-type and Entpd7−/− mice. Then, ATP or PBS was injected into the
Increased number of Th17 cells in the small intestinal lamina propria in Entpd7–/– mice

A previous study showed that luminal ATP in the intestine mediates Th17 cell development (10). In addition, extracellular ATP was shown to induce Th17 cell development via the inhibition of regulatory T cell functions (15). Therefore, we analyzed the number of CD4⁺ T cells expressing IL-17, IFN-γ, IL-10, and Foxp3 in the lamina propria of the small and large intestines. The numbers of IL-17⁺, IFN-γ⁺, IL-10⁺, and Foxp3⁺-expressing CD4⁺ T cells were not altered in the large intestinal lamina propria of Entpd7–/– mice (Fig. 3A, 3C). In contrast, the number of IL-17⁺-producing CD4⁺ T cells in the small intestinal lamina propria was markedly increased in Entpd7–/– mice compared with wild-type mice, although the numbers of IFN-γ⁺, IL-10⁺, and Foxp3⁺ T cells were not affected (Fig. 3B, 3C). The number of IL-17⁺-producing CD4⁺ T cells was not increased in other lymphoid organs, such as the spleen, MLNs, and Peyer’s patches of Entpd7–/– mice (Supplemental Fig. 4). Thus, Entpd7–/– mice showed elevation of Th17 cells in the small intestinal lamina propria. Consistent with Entpd7-expression patterns in the small intestine, the level of luminal ATP was higher in the distal region than in the proximal region of the small intestine of wild-type mice (Fig. 3D); accordingly, the number of IL-17⁺-producing CD4⁺ T cells was higher in the distal region (Fig. 3E).
Commensal microbiota-dependent, ATP-dependent increase in Th17 cells in Entpd7<sup>−/−</sup> mice

We analyzed whether increased Th17 cell development in the small intestine was intrinsic to the T cell itself or caused by extrinsic environmental factors. We first induced in vitro differentiation of splenic naive CD4<sup>+</sup> T cells into Th17 cells. Naive CD4<sup>+</sup> T cells were cultured in Th17 cell–skewing conditions and analyzed for IL-17 production (Fig. 4A). In vitro–differentiated CD4<sup>+</sup> T cells from wild-type and Entpd7<sup>−/−</sup> mice produced almost equal amounts of IL-17, indicating that Entpd7<sup>−/−</sup> T cells were not intrinsically programmed to preferentially differentiate into Th17 cells. We then treated Entpd7<sup>−/−</sup> mice orally with combinations of four antibiotics (i.e., vancomycin, streptomycin, metronidazole, and ampicillin) from birth (Fig. 4B). In antibiotic-treated wild-type and Entpd7<sup>−/−</sup> mice, the number of IFN-γ- and IL-10-producing T cells, as well as IFN-γ– and IL-10–producing T cells, in the small intestinal lamina propria was dramatically reduced. These findings indicate that the augmentation of Th17 cells in Entpd7<sup>−/−</sup> mice was caused by altered environmental factors influenced by commensal microbiota. Recent data demonstrate that a specific microbiota, such as SFB, induces Th17 cell differentiation in the small intestine (37, 38). Therefore, we analyzed the number of intestinal bacteria in the luminal contents and epithelial layers of the small intestine of wild-type and Entpd7<sup>−/−</sup> mice (Fig. 4C). The number of intestinal bacteria was not altered in Entpd7<sup>−/−</sup> mice, indicating that Entpd7 deficiency did not cause alteration of microbiota.

Because commensal microbiota were shown to influence luminal ATP level (10), we analyzed the effect of the blockade of ATP action. αATP, which antagonizes P2X receptors, was shown to be effective in modulating T cell responses in mice, especially Th17 cell responses (15). Therefore, Entpd7<sup>−/−</sup> mice were treated with αATP; however, the total number of CD4<sup>+</sup> cells in the small intestinal lamina propria was not altered (Fig. 5A). In accordance with the previous finding that αATP inhibits T cell responses, such as cytokine production (14), the number of IFN-γ<sup>+</sup> and IL-10<sup>+</sup> CD4<sup>+</sup> T cells was moderately reduced (Fig. 5B). Notably, the number of IL-17–producing CD4<sup>+</sup> T cells was severely reduced in αATP-treated Entpd7<sup>−/−</sup> mice (Fig. 5C, 5D). In an intestinal inflammation model of immunocompromised Cdb<sup>−/−</sup> mice transferred with conventional T cells, αATP treatment increased the number of Foxp3<sup>+</sup> CD4<sup>+</sup> T cells in MLNs of the diseased mice (14). However, Entpd7<sup>−/−</sup> mice treated with αATP did not show any increase in the number of Foxp3<sup>+</sup> T cells in MLNs of the small intestinal lamina propria (Fig. 5E). Thus, the ATP antagonist severely decreased Th17 cells and moderately reduced IFN-γ<sup>+</sup>– and IL-10–producing T cells. These findings indicate that the increased ATP level is responsible for enhanced Th17 cell development in the small intestine of Entpd7<sup>−/−</sup> mice.
Resistant to intestinal C. rodentium infection in Entpd7−/− mice

A previous study showed that development of Th17 cells in the small intestine provides the resistance to oral infection with C. rodentium (38). Therefore, we orally infected wild-type and Entpd7−/− mice with C. rodentium. The CFU titers of bacteria in the spleen were measured at day 14 after the infection (Fig. 6A, 6B). The number of spleens that was invaded with C. rodentium was dramatically decreased in Entpd7−/− mice. Accordingly, Entpd7−/− mice had decreased numbers of C. rodentium in the spleen compared with wild-type mice. In addition, although some wild-type mice died after the oral C. rodentium infection, none of the Entpd7−/− mice died (Fig. 6C). Thus, Entpd7−/− mice are resistant to the intestinal bacterium C. rodentium.

Deteriorated EAE in Entpd7−/− mice

Enhanced Th17 responses are implicated in the development of several immune disorders, including EAE (39). Th17 cells, which develop in the small intestine, were shown to induce inflammation in extraintestinal tissues, such as arthritis in the ankle joints (40). Furthermore, commensal microbiota were shown to be involved in the pathogenesis of EAE (41). Therefore, we used a MOG peptide–induced model of EAE in Entpd7−/− mice to determine the effect of ENTPDase7-mediated regulation of intestinal Th17 cells in inflammatory conditions in vivo. As shown in Fig. 7A, s.c. immunization of wild-type mice with the MOG peptide, together with pertussis toxin, induced encephalomyelitis associated with rapidly ascending paralysis appearing at approximately day 10–12. MOG peptide–immunized Entpd7−/− mice showed more severe clinical symptoms. We then analyzed cytokine production from CD4+ T cells infiltrated into the CNS of the diseased mice (Fig. 7B). In wild-type and Entpd7−/− mice, infiltration of IL-17– and IFN-γ–producing CD4+ T cells, as well as IL-17/IFN-γ double-producing cells, was observed. IL-17/IFN-γ double-producing CD4+ T cells increased markedly in Entpd7−/− mice compared with diseased wild-type mice. Thus, in the absence of Entpd7, severe EAE developed that was accompanied by an increased infiltration of CD4+ T cells producing both IL-17 and IFN-γ.

Discussion

In the current study, we analyzed the physiological function of ENTPDase7, which is preferentially expressed in ECs of the small intestine. ENTPDase7 deficiency in mice led to increased ATP levels in the small intestinal lumen, indicating that ENTPDase7 is responsible for the maintenance of luminal ATP levels. The number of IL-17–producing Th17 cells in the small intestinal lamina propria was increased in Entpd7−/− mice. The number of Th17

FIGURE 4. Decreased number of Th17 cells in antibiotic-treated Entpd7−/− mice. (A) Splenic naive CD4+ T lymphocytes were cultured for 4 d under Th17-polarizing conditions (TGF-β, IL-6, anti–IFN-γ, and anti–IL-4) or Th0 conditions (anti–IFN-γ and anti–IL-4). Then, lymphocytes were harvested, stimulated, permeabilized, stained for IL-17 and IFN-γ, and analyzed by flow cytometry. Data are representative of three independent experiments. (B) Wild-type (n = 4) and Entpd7−/− (n = 4) mice were administered vancomycin, metronidazole, ampicillin, and neomycin sulfate in drinking water from birth. The small intestinal lamina propria lymphocytes were isolated at 8 wk of age and analyzed for production of IL-17, IFN-γ, and IL-10 from CD4+ T cells by flow cytometry. Representative FACS dot plots and total numbers of cells gated on small intestinal lamina propria CD4+ cells are shown. (C) Intestinal bacteria in the luminal contents and epithelial layers of the small intestines of wild-type and Entpd7−/− mice. DNA isolated from the luminal contents and epithelial layers of the small intestines was analyzed by real-time quantitative PCR using primers for bacterial group–specific 16S rRNA genes. Data are representative of two independent experiments and are mean ± SD of five mice.
cells was decreased in Entpd7−/− mice in the absence of commensal microbiota or after ATP antagonist treatment. Entpd7−/− mice were resistant to infection with C. rodentium, against which Th17-related cytokines play a major role.

A previous report indicated that human ENTPDase7 is expressed in the membrane of intracellular compartments (28). The intracellular ATP level, which was analyzed using total-cell lysates, was not altered in Entpd7−/− ECs. However, given that the ATP concentration in the cytoplasm is >1 mM, whereas the ATP concentration in the extracellular compartment is usually <10 nM, ATP levels within the ENTPDase7-expressing cellular vesicles of ECs would be increased in the absence of ENTPDase7. Indeed, the membrane fraction of intestinal ECs had an enzymatic activity to hydrolyze ATP, and its activity was decreased in the absence of Entpd7. Because Entpd7 was highly expressed in goblet cells, as well as absorptive ECs, it is possible that Entpd7 is

**FIGURE 5.** Decreased number of Th17 cells in oATP-treated Entpd7−/− mice. (A–C) Entpd7−/− mice were administered 100 µl of 6 mM oATP or PBS i.v. daily for 2 wk. The small intestinal lamina propria lymphocytes were then isolated and analyzed for production of IFN-γ, IL-10, and IL-17 from CD4+ T cells by flow cytometry. The percentages and total numbers of CD4+ T cells (A), IFN-γ and IL-10–producing CD4+ T cells (B), and IL-17–producing CD4+ T cells (C) in the small intestinal lamina propria of PBS- or oATP-treated Entpd7−/− mice. Data are representative of two independent experiments and are mean ± SD of four mice. (D) Representative FACS graph showing IL-17 production gated on small intestinal lamina propria CD4+ T cells of the indicated mice. (E) The percentages and total numbers of Foxp3+ CD4+ T cells in the small intestinal lamina propria and MLNs of PBS- or oATP-treated Entpd7−/− mice. Data are representative of two independent experiments and are mean ± SD of three mice. *p < 0.05, **p < 0.01.

**FIGURE 6.** Resistance to intestinal C. rodentium infection in Entpd7−/− mice. (A–C) Wild-type (n = 19) and Entpd7−/− (n = 19) mice were infected orally with C. rodentium. (A) Detection rate of C. rodentium in the spleen on day 14. The pooled data of two independent experiments are shown. (B) Log10 CFU of C. rodentium in spleens. (C) Survival rate of the mice at the indicated time points. The pooled data of two independent experiments are shown. **p < 0.01.

**FIGURE 7.** Severe EAE in Entpd7−/− mice. (A) Wild-type (n = 10) and Entpd7−/− (n = 11) mice were immunized with 100 µg MOG35–55 peptide in CFA; 100 ng of pertussis toxin was injected i.p. on days 0 and 2. The mean clinical score was calculated by averaging the scores of the mice in each group. Data are mean ± SEM at each time point. Experiments were performed twice with similar results. *p < 0.05. (B) Representative FACS dot plots gated on CD4+ cells of the CNS in the indicated mice at day 17 after EAE induction (left and middle panels). CNS lymphocytes were isolated from wild-type and Entpd7−/− mice 17 d after EAE induction and analyzed for the production of IFN-γ and IL-17 from CD4+ T cells by flow cytometry (right panel). Data are representative of five mice analyzed. **p < 0.05.
expressed in the membrane of mucin-containing vacuoles of goblet cells to control ATP levels in the vacuole. Given that goblet cells of the airway were shown to secrete ATP, as well as mucin (42), intestinal goblet cells might be a major source of luminal ATP, the level of which is closely regulated by ENTPDase7.

Human ENTPDase7 was shown to preferentially hydrolyze UTP, GTP, and CTP rather than ATP (28). However, the membrane fraction of mouse intestinal ECs effectively hydrolyzed ATP, and its activity was impaired by Entpd7 deficiency. Thus, mouse ENTPDase7, unlike human ENTPDase7, effectively hydrolyzes ATP. Indeed, apparent differences in amino acid sequences are observed in a domain between the second and third apyrase-conserved regions, supporting that mouse and human ENTPDase7 have different substrate affinities.

Luminal ATP is supposed to be derived from ECs, as discussed above. In addition, commensal microbiota are a source of luminal ATP (10). In this regard, commensal microbiota, especially SFB, mediate Th17 cell development in the small intestine, possibly through ATP-independent mechanisms (37, 43). Therefore, in the small intestine, luminal ATP may mediate Th17 cell development cooperatively with Th17-inducing commensal microbiota. There is controversy as to how luminal ATP is sensed and induces Th17 development. Intestinal CXCR1+ DCs were shown to extrude their dendrites into the lumen to sample intestinal Ags (44, 45). These intestinal DCs might sense luminal ATP via purinergic receptors. Alternatively, as reported in several studies, ECs sense extracellular ATP (11, 25, 46). Therefore, intestinal ECs trigger inflammatory responses to activate T cell development via ATP sensing. Indeed, intestinal ECs were shown to control DC functions (47).

C. rodentium is an enteric bacterium that colonizes the intestine of mice postinfection. Clearance of C. rodentium is shown to be dependent on Th17-related cytokines, such as IL-17 and IL-22 (48, 49). Data showing that mice lacking IL-23, a critical cytokine for Th17 cell development, are highly susceptible to C. rodentium infection also indicate that Th17-related cytokines are critical for the resistance to intestinal C. rodentium infection (50). Consistent with these facts, Entpd7−/− mice showing an increased number of Th17 cells in the small intestine are highly resistant to intestinal infection with C. rodentium. IL-22 and IL-17, which induce production of antibacterial peptides (REGIIIγ and δ) from intestinal ECs (48, 51), are produced from other cell populations, such as innate lymphoid cells and γδT cells (52, 53). Therefore, Th17 cells, together with an innate type of IL-17–producing cells, contribute to intestinal pathogens.

In an EAE model, CD4+ T cells producing both IL-17 and IFN-γ are observed in the CNS (54–56). It is still controversial whether these IL-17/IFN-γ double-producing T cells are Th1 or Th17 cells, but they do contribute to EAE pathogenesis (54). A study using Il23ra−/− mice, which showed a reduced number of Th17 cells, as well as IL-17/IFN-γ double-producing T cells and a normal number of Th1 cells, suggested that IL-17/IFN-γ double-producing T cells are derived from Th17 cells (57). Therefore, increased numbers of IL-17/IFN-γ double-producing T cells in the CNS of Entpd7−/− mice with EAE might be due to enhanced Th17 responses.

It remains unclear how Th17 cells residing in the small intestine mediate EAE. However, several lines of evidence indicate the relevance between gut immune cells and immune disorders in extraintestinal tissues: Th17 cells induced by SFB in the small intestinal lamina propria were shown to be responsible for the induction of autoimmune arthritis (40); alteration of the commensal flora composition influences the severity of EAE (41); and SFB-induced Th17 cells in the small intestine induce EAE (58).

Thus, intestinal effector T cells are responsible for the induction of immune disorders in nongut tissues, including the CNS, and our present study demonstrates that enhanced intestinal Th17 responses can induce severe inflammatory conditions in these disease models.

In this study, we showed that an ENTPDase expressed by intestinal ECs regulates luminal ATP levels and, thereby, controls intestinal immune responses. Another ENTPDase, ENTPDase8, is selectively expressed by the ECs in the large intestine, as well as the small intestine (T. Kusu and K. Takeda, unpublished observations). Characterization of ENTPDase8 functions in terms of regulation of intestinal immune responses will be an interesting issue to be addressed in the future.

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Disclosures
The authors have no financial conflicts of interest.

References


Supplemental Figure 1. Targeted disruption of the gene encoding mouse ENTPDase7

(A) The structures of the Entpd7 gene (top), the targeting vector (middle) and the predicted disrupted gene (bottom). Black box, coding exon. (B) Southern blot analysis of offspring from the heterozygote intercrosses. Genomic DNA was extracted from mouse tail tissue, digested with ScaI, separated by electrophoresis and hybridized with the radiolabeled probe shown in (A). Southern blot analysis detected a single 8.5-kb band for wild-type mice (+/+), a 5.5-kb band for homozygous Entpd7-deficient mice (-/-), and both bands for heterozygous mice (+/-). (C) Northern blot analysis of offspring from the heterozygote intercrosses. Total RNA was extracted from the small intestine, separated by electrophoresis and hybridized with the radiolabeled Entpd7 cDNA probe. Northern blot analysis detected Entpd7 mRNA bands for wild-type mice (+/+) and no Entpd7 mRNA bands for homozygous Entpd7-deficient mice (-/-). Bottom, the same amount of mRNA extracts hybridized with the radiolabeled β-actin (Actb) cDNA probe to monitor mRNA extraction. (D) Representative FACS dot plots of surface markers of splenic lymphocytes. Spleens were collected from wild-type and Entpd7/-/- mice and single-cell suspensions were prepared. The cells were stained for CD4/CD8 or CD3/B220 and analyzed by flow cytometry.
Supplemental Figure 2. Lymphocyte composition in the intestine of Entpd7−/− mice
Intraepithelial lymphocytes and lamina propria lymphocytes were isolated from the small and large intestines of wild-type and Entpd7−/− mice, stained for CD3/TCRβ, TCRβ/CD4/CD8α/CD8β and TCRγδ/CD4/CD8α/CD8β, and analyzed by flow cytometry. Total numbers of CD3+ cells and proportion of each cellular subset were shown. Data are shown as means + SD of four mice.
Supplemental Figure 3

To establish small intestinal epithelial cell lines, wild-type and Entpd7−/− mice were crossed with H-2Kb-tsA58 transgenic mice. Crypts isolated from the small intestine were cultured at 33°C for 3 weeks, and growing cells were cloned and expanded to establish the epithelial cell lines. The cell lines were stained with anti-cytokeratin, and visualized with diaminobenzidine (A). The expression of Krt7, encoding cytokeratin-7, and Entpd7, was analyzed by real-time RT-PCR. ND; not detected (B).
Supplemental Figure 4. Cytokine production of CD4⁺ T cells in several lymphoid organs
Representative FACS dot plots of intracellular cytokine profiles of CD4⁺ T cells and total numbers of IL17-producing CD4⁺ cells in the MLNs, Peyer's patches, and spleen. Cells were isolated from the indicated lymphoid organs of wild-type and Entpd7⁻/⁻ mice, stimulated, and permeabilized to stain intracellular IL-17 and IFN-γ. Data are shown as means ± SD of four mice.