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CD39 and CD161 Modulate Th17 Responses in Crohn’s Disease

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CD39 (ENTPD1) is expressed by subsets of pathogenic human CD4+ T cells, such as Th17 cells. These Th17 cells are considered important in intestinal inflammation, such as seen in Crohn’s disease (CD). Recently, CD161 (NKR-PIA) was shown to be a phenotypic marker of human Th17 cells. In this study, we report that coexpression of CD161 and CD39 not only identifies these cells but also promotes Th17 generation. We note that human CD4+CD39+CD161+ T cells can be induced under stimulatory conditions that promote Th17 in vitro. Furthermore, CD4+CD39+CD161+ cells purified from blood and intestinal tissues, from both healthy controls and patients with CD, are of the Th17 phenotype and exhibit proinflammatory functions. CD39 is coexpressed with CD161, and this association augments acid sphingomyelinase (ASM) activity upon stimulation of CD4+ T cells. These pathways regulate mammalian target of rapamycin and STAT3 signaling, thereby limiting IL-17 production in CD4+ T cells obtained from both controls and patients with active CD. Increased levels of CD39+CD161+ CD4+ T cells in blood or lamina propria are noted in patients with CD, and levels directly correlate with clinical disease activity. Hence, coexpression of CD39 and CD161 by CD4+ T cells might serve as a biomarker to monitor Th17 responsiveness. Collectively, CD39 and CD161 modulate human Th17 responses in CD through alterations in purinergic nucleotide–mediated responses and ASM catalytic bioactivity, respectively. The Journal of Immunology, 2014, 193: 000–000.

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Abbreviations used in this article: ADA, adenosine deaminase; ADO, adenosine; AMPK, AMP-activated protein kinase; ASM, acid sphingomyelinase; CD, Crohn’s disease; mTOR, mammalian target of rapamycin; qRT-PCR, quantitative real-time PCR; sh-C, shRNA vector control; shRNA, short hairpin RNA; TLC, thin layer chromatography; Treg, regulatory T cell.
Acid sphingomyelinase (ASM) is a hydrolase enzyme located at the plasma membrane, which plays an important role in mediating cell signaling by catalyzing sphingomyelin into ceramide (20). ASM was shown to promote LPS-induced proinflammatory cytokine release from macrophages. Furthermore, ASM inhibition protects mice against dextran sulfate sodium–induced colitis (21). However, it remains unclear whether ASM has the potential to modulate Th17 cell differentiation by specifically impacting ceramide-mediated cellular signaling pathways.

In this study, we observe substantial increases in IL-17–producing CD39+CD161+ CD4+ T cell subpopulations in both peripheral blood and lamina propria of patients with active CD. Coexpression of CD39 and CD161 modulates human CD4+ Th17 responses by interactions that are inclusive of links to ASM. Upon activation, this putative molecular association boosts ASM enzymatic activity, which enhances cellular ceramide production to activate downstream signaling components via mammalian target of rapamycin (mTOR) and STAT3. These signaling pathways are crucial elements that control Th17 generation (22–24). Importantly, we also show that blockade of ASM bioactivity, with imipramine or by knockdown of ASM, limits CD4+ Th17 generation from T cells derived from healthy volunteers, as well as from patients with CD.

We suggest that CD39 and CD161 coexpression may serve as a reliable surface marker of Th17 cells in CD. We further note and describe novel and possibly linked elements of both extracellular nucleotide and sphingolipid homeostasis that are pivotal in the control of Th17 cells and which might be exploited therapeutically.

Materials and Methods

General reagents

All chemicals were purchased from Sigma-Aldrich (St. Louis, MO), unless otherwise stated. All cell culture media and reagents were from Life Technologies (Grand Island, NY), and all cytokines were from R&D Systems (Minneapolis, MN).

Abs

FACS studies were performed using FITC-, PE-, PE-Cy5-, PE-Cy7-, Pacific blue–, or allophycocyanin-conjugated anti-human Abs to CD4 (clone ORT4), CD3 (HIT3a), CD161 (HP-3G10), IL-17A (BL168), IFN-γ (4S.B3), IL-10 (JES5-19H7), and CCR6 (T/G7) (all from BioLegend, San Diego, CA); CD39 (BU61) (Ancell, Bayport, MN); CD45RO (UCHL1) (eBioscience, San Diego, CA); IL-23R (218213) (R&D Systems); phospho-mTOR (Ser2448) (#2971), mTOR (#2972), phospho-p70 S6 kinase (AMPK) (ab125452), IL-10 (JES3-19F1), and CCR6 (TG7) (all from BioLegend, San Diego, CA); phospho-mTOR (Ser2448) (#2971) (Cell Signaling Technology, Danvers, MA). Isotype-control Abs (IgG1) were from Ancell or BioLegend. Abs used for Western blot included phospho-Akt (Ser473) (#9271), phospho-mTOR (Ser2448) (#2971), mTOR (#2972), phospho-p70 S6 kinase (Ser473) (#208), phospho-STAT3 (Ty#705) (#9145), STAT3 (#9132), phospho-PKM2 (Ty#136) (#8327), phospho–AMP-activated protein kinase (AMPKα) (Thr#172) (#2535), phospho-AMPKβ1 (Ser#188) (#1481), and phospho-ACC (Ser#79) (#3661) (all from Cell Signaling Technology); ASM (SMPD1) (Cell Signaling Technology and R&D Systems); CD161 (DX12) and CD4 (RPA-T4) (BD Biosciences); CD39 (BU61) (Ancell); or β-actin (AC-15, #ab8276) (Abcam, Cambridge, MA).

Study populations

PBMCs were isolated from platelet-depleted blood (leukofilters) obtained from healthy blood donors (Blood Donor Center at Children’s Hospital, Boston, MA) or from patients with CD (active disease or in remission). A total of 49 patients with CD was recruited in the study (22 men and 27 women; age range, 24–71 y; ileocolonic or colonic disease). Among those patients, 11 had received or were still receiving anti-TNF treatment, I was being treated with oral steroids, and the other 37 patients were studied at either the initial presentation or while not on active immunomodulatory treatments (inclusive of anti-TNF, steroids, or thiopurines). All diagnoses were confirmed by clinical, radiologic, endoscopic, and histologic criteria. All human studies were conducted in accordance with Helsinki principles and were approved by the Beth Israel Deaconess Medical Center Institutional Review Committee (No. 2011-P-000228). Written informed consent was obtained from all study participants.

Cell culture

Peripheral blood CD4+ T cells from healthy volunteers or patients with CD were isolated using a Human CD4+ T cell Enrichment Cocktail Kit (STEMCELL Technologies, Vancouver, BC, Canada) with slight modifications. Subtypes of CD3+CD4+ T cells were positively selected using a FACSaria cell sorter (BD Biosciences, San Jose, CA) with enrichment to >99% cell population. Cells were cultured in complete RPMI 1640 medium supplemented with 2 mM l-glutamine, 100 U/ml penicillin, 100 μg/ml streptomycin, 1% nonessential amino acids, and 10% FCS.

Lamina propria mononuclear cells

Lamina propria mononuclear cells were isolated from freshly biopsied colonic mucosal tissues using a previously described method with minor modifications (25). Briefly, tissues were incubated in HBSS containing EDTA (0.75 mM) and DTT (1 mM) at 37°C to remove the epithelium, followed by incubation with digestion cocktails (RPMI 1640 medium containing collagenase IV [400 U/ml] and Dnase I [0.01 mg/ml]) at 37°C for 1–2 h. The digested tissues were filtered through a 100-μm cell strainer and centrifuged at 1500 rpm for 5 min. Cell pellets were resuspended in complete RPMI 1640 media and overlaid on a 40–100% Ficoll gradient (GE Healthcare Life Sciences, Pittsburgh, PA). After centrifugation, the interphase (lamina propria mononuclear cells) was collected and washed once for subsequent experimentations.

Flow cytometric analysis

Flow cytometric analysis was performed as previously described (17) with slight modifications. The relevant fluorescein-labeled anti-CD3 or anti-CD5 Abs were used to set up the required compensations for flow cytometric analyses. For surface marker analyses, cells were stained with Abs diluted in PBS containing 0.1% BSA. For adenosine deaminase (ADA) expression assessment, cells were incubated with rabbit anti-human ADA Ab (Sigma-Aldrich; #HPA001399), followed by fluorescein-conjugated anti-rabbit Ab (Santa Cruz Biotechnology). For intracellular staining, cells were treated for 5 h with PMA (50 ng/ml), ionomycin (500 ng/ml), and brefeldin A (10 μg/ml). After surface staining, cells were permeabilized with 0.5% saponin in PBS for 20 min, followed by incubation with fluorescein-conjugated Abs. To determine intracellular phospho-mTOR levels, cells were incubated with phospho-mTOR (Ser2448) Ab after three washes with PBS containing 0.5% saponin, followed by fluorescein-conjugated anti-rabbit Ab. Fluorescein-labeled isotype IgG was used as a negative control in all cases. FACS data were acquired on a multicolor LSR II (BD Biosciences) and analyzed with FlowJo software (TreeStar, Ashland, OR).

In vitro T cell differentiation

Sorted CD4+ T cells were cultured in complete RPMI 1640 media. A total of 5 × 10^5 T cells was stimulated with anti-CD3/CD28 T-activation Dynabeads (Life Technologies) at a 1:1 cell/bead ratio. For Th17 polarization, cytokine cocktails including IL-6 (30 ng/ml), IL-1β (10 ng/ml), IL-23 (10 ng/ml), and TGF-β1 (1 ng/ml) were added to the cultures.

Cell cultures supplemented with IL-12 (10 ng/ml), IL-4 (10 ng/ml), or IL-2 (10 ng/ml), together with TGF-β1 (10 ng/ml), were used for deviation toward Th1 cells, Th2 cells, or Tregs, respectively. For agonist or antagonist studies, chemical compounds or control vehicles were introduced to cell cultures at the beginning of stimulation.

Quantitative real-time PCR

Total RNA was extracted from cells using the RNeasy Kit (QiAGEN, Valencia, CA) and reverse transcribed into cDNA using ABI Prism TaqMan reverse transcription reagents (cat. no. 204054; Applied Biosystems, Foster City, CA). Specific primers for quantitative real-time PCR (qRT-PCR) were obtained from Life Technologies; the sequences are shown in Supplemental Table I. Then, qRT-PCR was performed using a QuantFast SYBR Green PCR Kit (cat. no 204054; QiAGEN) on a Stratagene Mx3005P (Agilent Technologies, Santa Clara, CA). Relative expression was calculated using the ΔΔCT algorithm and GAPDH as an internal control.

Confocal microscopy

Cells were stained with Abs to CD39 (FITC; Ancell) and CD161 (PE-Cy5; BioLegend) and viewed with a confocal microscope (ZEISS LSM 510 Meta; Carl Zeiss Microscopy, Thornwood, NY).
**Commmunoprecipitation and immunoblotting**

Cells were washed with ice-cold PBS three times and lysed on ice in modified RIPA buffer (50 mM Tris-HCl [pH 7.4], 1% Nonidet P-40, 0.25% sodium deoxycholate, 150 mM NaCl) supplemented with Complete Protease Inhibitor Cocktails (Roche Diagnostics, Indianapolis, IN) and Phosphatase Inhibitor Cocktails (Sigma-Aldrich). The lysates were spun at 10,000 × g for 5 min at 4˚C.

Protein concentrations were determined by Bio-Rad DC protein assay reagents (Bio-Rad, Hercules, CA).

For immunoprecipitation, 300 μg total lysates from each sample was incubated with mouse anti-human Ab to CD39 (BU61; Ancell), ASM (SMPD1, R&D Systems), or control IgG (BD Biosciences), with end-over-end rotation at 4˚C for 3 h. A total of 30 μg protein G-Sepharose 4B (Sigma-Aldrich) was added to each mixture and incubated at 4˚C overnight. The mixtures were centrifuged at 4˚C for 5 min at 2500 rpm, washed five times with RIPA buffer, and eluted with 2× XT Sample Buffer (cat. no. 161-0791; Bio-Rad), followed by Western blot analysis.

For Western blotting, proteins were separated on 4–12% Criterion XT Bis-Tris SDS-PAGE gels (Bio-Rad) and transferred to a polyvinylidene difluoride membrane (cat. no. IPVH00010; EMD Millipore, Billerica, MA) by semidyry electroblotting. The membranes were then probed with specific Abs against proteins of interest. Bands were visualized using HRP-conjugated goat anti-mouse, donkey anti-rabbit, or donkey anti-sheep IgG (Thermo Scientific, Rockford, IL) and SuperSignal West Femto Maximum Sensitivity Substrate reagents (cat. no. PI-34096; Thermo Scientific), according to the manufacturer’s instructions.

**Ectonucleotidase enzymatic activity analysis**

Cell-associated nucleoside triphosphate diphosphohydrolase activity was analyzed by thin layer chromatography (TLC), a technique fully established in the laboratory (26). Freshly sorted CD4+ T cells (2 × 10^6) were incubated with 2 μCi/ml [3^14]ADP (GE Healthcare Life Sciences) in 10 mM Ca^2+ and 5 mM Mg^2+ for the indicated times from 5 to 30 min. Then 5-μl aliquots were removed at each time point and analyzed for the presence of [3^14]ADP hydrolysis products by TLC. [3^14]ADP, [3^14]AMP, and [3^14]adenosine (ADO) incubated in PBS served as standards. The TLC glass plate was exposed in a storage phosphor cassette for 24 h, followed by visualization of phosphor release captured by a Storm Scanner. ADO uptake was inhibited by the use of the Ca^2+ channel blocker dipryridamole (10 μM).

**ASM activity assay**

Enzymatic activity of ASM of CD4+ T cells was measured, as previously reported, with minor modifications (27). Levels of derived ceramide, the lipid cleavage product of ASM, were used as a marker indicator for ASM bioactivity. Briefly, cells (1 × 10^6) were processed in lysis buffer (50 mM sodium acetate [pH 5], 0.5% Nonidet P-40). A total of 10 μg cell lysates was incubated with 0.2 μg sphingomyelin (dissolved in a 1:1 mixture of isopropanol and methanol (3:2). Ceramide production was visualized in iodine followed by chromatographic separation using a solvent system of chloroform and methanol (9:1). The mixtures were centrifuged at 4˚C for 5 min at 2500 rpm, washed five times with RIPA buffer, and eluted with 2× XT Sample Buffer (cat. no. 161-0791; Bio-Rad), followed by Western blot analysis.

**Statistical analysis**

Results in this study are expressed as the mean ± SEM of values. Differences between experimental groups were assessed by one-way ANOVA. The two-tailed Student t test was used to compare two groups, and the Tukey–Kramer multiple-comparison test was used to compare multiple groups. Correlations between subtype cell population and clinical activity was calculated by the Spearman test. Significance was defined as p < 0.05.

**Results**

**CD39^+CD161^+ CD4^+ T cells exhibit molecular signatures of Th17 cells**

CD4^+ T cells were isolated from peripheral blood of healthy volunteers and analyzed by FACS. These cells were grouped into four distinct subpopulations, based on the differential expression of CD39 and CD161 (Fig. 1A). To determine the exact nature of these subpopulations, we examined the expression of Th17 phenotypic markers by FACS and/or qRT-PCR analyses. CD39^+CD161^+ cells exhibited robust expression of Th17 markers, such as chemokine receptors inclusive of CCR5, CCR6, and CXCR3, and cytokine receptors, including TGFRI, IL-23R, and IL-6R (30), compared with the other three subpopulations (Fig. 1B, 1C). Interestingly, the vast majority of CD39^+CD161^+ cells appeared to be of the memory cell phenotype (CD45RO^+), with prototypic Th17 features (5, 31) (Fig. 1B). The mRNA expression levels of IL-17 and RORγt were considered comparable among the four CD4^+ T cell subsets in the quiescent state, prior to cell activation (data not shown).

To determine whether dual expression of CD39/CD161 correlates with Th17 differentiation, we next directly stimulated quiescent peripheral blood CD4^+ T cells with various cytokines and under different T cell–polarizing conditions for 18 h. We noted that the frequencies of CD39^+CD161^+ populations increased markedly under these Th17-differentiation conditions (with or without IL-1β + IL-23), in contrast to parallel studies with conditions skewing differentiation toward Tregs (anti-CD3/28 + TGF-β1 + IL-2) (Fig. 1D) or Th1/2 cells (data not shown). Moreover, in the absence of Th17-skewing cytokines, the 72-h stimulation with anti-CD3/CD28 T-activation Dynabeads alone had minimal effects on the induction of IL-17 expression in any of the four CD4^+ T cell subsets (data not shown). This induction of CD39 expression and Th17 cells was further linked to STAT3 signaling (23).

We then evaluated the proinflammatory potential and metabolic characteristics of CD4^+CD39^+CD161^- T cells. The four subpopulations were FACS sorted and then cultured under the same Th17-differentiation conditions. After 72 h, the CD4^+CD39^+CD161^- population produced substantial levels of intracellular IL-17 (Fig. 2A, Supplemental Fig. 1A). This activity was concordant with other typical aspects of Th17 cells (22-24, 30, 32, 33): robust gene expression of Th17 markers (IL-17, IL-22, and RORγt), induction of glycolysis modulators (HIF1α and LDH1) (Fig. 2B), and enhanced signaling via Akt-mTOR and STAT3, together with other expected metabolic activation (PKM2 and AMPK-ACC) (34, 35) (Fig. 2C, 2D).

Furthermore, blocking glycolysis (by 2-DG), STAT3 (by WP1066), or AMPK (by compound C) in this population markedly impaired the generation of Th17 cells, whereas inhibiting fatty acid oxidation (by etomoxir) substantially augmented IL-17 expression (Fig. 2E, 2F).
Supplemental Fig. 1B). These data suggest that CD39+CD161+ Th17 cells are preferentially active metabolically with respect to glycolytic pathways (32, 33).

Taken together, the data clearly show that coexpression of CD39 and CD161 defines a CD4+ T cell population that is more highly enriched in Th17 cells and has distinct phenotypic features.

CD39 and CD161 coexpression denotes Th17 populations in CD

To further investigate the physiological relevance of CD39 and CD161 coexpression in CD, we next contrasted patterns of CD39 and CD161 expression in CD4+ T cells isolated from peripheral blood and lamina propria of healthy volunteers and from patients with CD.

We noted that levels of CD4+ T cells expressing CD39+, CD161+, or the double-positive label of CD39+CD161+ in blood or lamina propria were consistently higher in patients with active CD compared with healthy controls (Fig. 3A, 3B). Concurrently, levels of CD39+CD4+ and/or CD39+CD161+ CD4+ T cells in patients with active CD were significantly higher than those in patients with quiescent disease. In contrast, levels of CD4+ CD161+ T cells without the additional CD39 expression marker in these lamina propria cells did not allow differentiation between patients with active and inactive CD (Fig. 3B).

FIGURE 1. CD39 and CD161 expression patterns denote Th17 cell phenotypes in human CD4 T cell compartments. (A) Representative FACS analysis of CD39 and CD161 surface expression on CD4+ T cells of peripheral blood from healthy volunteers (n = 36). Surface and mRNA expression of Th17-related molecules in four CD4+ T cell subsets (n = 5), as determined by FACS (B) and qRT-PCR (C), respectively. (D) Fresh isolated CD4+ T cells (n = 4) were stimulated with different cytokine cocktails in the presence of anti-CD3/CD28 T-activation Dynabeads for 18 h. Frequencies of CD39+CD161+ CD4+ T cells were evaluated by FACS, and changes are expressed relative to unstimulated controls. Data are mean ± SEM of the indicated number of experiments. Percentages of gated cells are shown. *p < 0.05, **p < 0.01, ***p < 0.001.
Given the above findings, we next evaluated relationships between CD4⁺ T cell expression of CD39/CD161 and clinical disease activity scores. The varied levels of CD39⁺ and CD39⁺CD161⁺ CD4⁺ T cells in peripheral blood correlated positively with disease activity using the Harvey Bradshaw Index (36) (Supplemental Fig. 2A). In contrast, levels of lamina propria CD161⁺CD4⁺ T cells, not considering levels of CD39 expression, did not show any direct statistical correlation with disease activity (Supplemental Fig. 2A).

It was noted previously that lamina propria CD4⁺ T cells from patients with active CD express higher levels of intracellular IL-17

**FIGURE 2.** CD4⁺CD39⁺CD161⁺ T cells exhibit an activated Th17 molecular signature. Subsets of CD4⁺ T cells were sorted by CD39 and CD161 expression by FACS and cultured under Th17-differentiation conditions for 48 or 72 h. Representative flow cytometry analysis of intracellular levels of IL-17 and IFN-γ at 72 h (n = 16) (A) and gene expression at 48 h (n = 4) (B). (C and D) Components of intracellular signaling transduction were measured at 48 h by Western blot using probes as indicated (n = 4). (E) CD4⁺CD39⁺CD161⁺ T cells were expanded under Th17 condition for 72 h in the presence of vehicle, 2-deoxy-D-glucose (2-DG, a prototypical inhibitor of glycolysis), or etomoxir (an inhibitor of carnitine palmitoyltransferase 1, the rate-limiting enzyme in β-oxidation of fatty acids). Representative flow cytometry analysis of intracellular IL-17 and IL-10 (n = 4). *p < 0.05, **p < 0.01 versus the other three groups.
FIGURE 3. CD4⁺CD39⁺CD161⁺ T cells are indicative of Th17 responses in CD. (A and B) CD4⁺ T cells were isolated from healthy volunteers and patients with active or inactive CD using peripheral blood (n = 36, 16, or 12, respectively) or lamina propria (n = 17, 15, or 6, respectively) cells. Representative FACS analysis of subpopulations sorted by CD39 and CD161 (A) are expressed as a percentage of total CD4⁺ T cells (B). (C-E) CD4⁺ T cells were purified from lamina propria of patients with active CD (n = 8) and treated with PMA (50 nM) and ionomycin (500 nM) for 5 h. This was followed by flow cytometry analysis of intracellular IL-17 and IFN-γ (C and D) and CD39 and CD161 expression patterns in gated IL-17⁺CD4⁺ T cells (E). *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001.
and IFN-γ (37, 38). In this study, we demonstrated that lamina propria CD4+CD39+CD161+ T cells in patients with active inflammation also exhibited substantial levels of intracellular IL-17 and IFN-γ (Fig. 3C, 3D), in association with high levels of mTOR phosphorylation (Supplemental Fig. 2B, 2C) and STAT3 (data not shown).

We also noted that most IL-17+ CD4+ T cells express both CD39 and CD161 (Fig. 3E). Therefore, dual expression of high levels of CD39 and CD161 by CD4+ T cells appears indicative of the Th17 phenotype within both the gut and the peripheral blood of patients with CD.

**CD39 associates with CD161 on CD4+ T cells**

Next, we investigated how dual expression of CD39 and CD161 might impact Th17 cell phenotype and functions. Because both molecules are cell surface markers, we first examined their localization on CD4+ T cell membranes. We noted that CD39 potentially colocalized with CD161 in CD4+ T cells, as shown by confocal imaging and coimmunoprecipitation studies (Fig. 4A, 4B). Such associations were noted regardless of the state of cell activation (resting or after activation with anti-CD3/CD28 beads).

In addition to CD39, differential expression and activity of other purinergic ecto-enzymes in sorted CD4+ T cell subsets were examined. Our specific focus was on the ecto-enzymes CD73 [ecto-5′-nucleotidase, which catalyzes the hydrolysis of AMP to ADO (39)] and ADA [which catalyzes deamination of ADO to form inosine (40)].

The two CD39+ subpopulations exhibit substantial CD39 activities, whereas CD39+CD161+ cells contained higher ADA expression and possible specific activities (inferred by higher inosine levels) (Fig. 4C–E). Hence, ADO is readily cleaved by ADA on Th17 cells to form inosine (Fig. 4C), which has no effect on human Th17 generation (data not shown). These functional data were concordant with the protein expression levels of these molecules, in that CD39+CD161+ were shown to express high levels of ADA (Fig. 4D, 4E).

We previously showed that extracellular purine metabolism, with ADO generation as modulated by CD39 and CD73, impacts murine adaptive immune responses (16, 17). We then tested the relevance of this system in human CD4+ Th17 cells. In contrast to mouse cells (16, 17, 23), purines (i.e., ATP, AMP, or ADO) exogenously added to CD39+CD161+ cells had no substantial impact on human Th17 generation (Fig. 4F). These data infer that rapid scavenging of ATP and ADO by the high levels of CD39 and ADA expressed by CD4+CD39+CD161+ cells results in this particular Th17 phenotype. These cells are refractory to both ATP (because of CD39 activity) and ADO (because of ADA associations, which rapidly deaminates this nucleoside).

**CD39–CD161 association involves ASM activation and impacts STAT3/mTOR signaling in vitro**

Human CD161 was shown to interact with, and then activate, ASM to generate ceramide, a second messenger with prominent roles in NK cell biology (41). In this study, we evaluated the role of ASM in human Th17 development. Production of ceramide, a lipid cleavage product of ASM and an indicator of bioactivity, was dramatically elevated by 2 h and reached a plateau after 24 h under Th17-polarization conditions (data not shown).

This change was noted to occur with increased activation of Akt-mTOR and STAT3 signaling. Conversely, inhibition of ASM activity in CD4+ T cells by imipramine substantially blocked ceramide generation and concurrently inhibited Akt-mTOR and STAT3 activation (Fig. 5A), which differentially limited IL-17+ cell induction (Fig. 5B). The results suggest that ASM bioactivity might be an integral component, at least in part, of Th17 generation.

We then examined the properties of possible CD39/CD161 colocalization, as might concern ASM activation. By performing coimmunoprecipitation, we also noted that ASM associates with CD39 and CD161 in human CD4+ T cells (data not shown), and in CD4+CD39+CD161+ T cells. ASM has the potential to interact in the same manner with both CD39 and CD161 in quiescent cells, as well as in cells stimulated to drive Th17 differentiation (Fig. 5C).

Next, we detailed profiles of ASM enzymatic activity among the four CD4+ T cell subpopulations. At 48 h post-Th17 differentiation, robust ceramide generation was observed only in CD4+CD39+CD161+ cells (Fig. 5D). No differences in ASM protein levels were noted among the subsets (data not shown).

We then evaluated the impact of manipulating cells, by cross-linking Abs to CD39 and/or CD161, on ASM activity in the four freshly isolated CD4 T subsets. Ceramide production indicating ASM bioactivity was stimulated by TNF (27) or by these cross-linked Abs to CD161 (41). These processes appeared regulated albeit shown to be transient. We measured ceramide production at various time points after the addition of cross-linked Abs. Peak ASM activity was observed at or before 5 min post-stimulation. As shown in Fig. 5E, cross-linking Abs to either CD39 or CD161 appeared to augment ASM activity in CD39+CD161+ cells, reaching the highest level at 5 min, but not in CD39– or CD39+CD161– cells (data not shown). Additionally, additive effects were seen when both Abs were applied (Fig. 5E).

Lastly, blocking ASM activity (by imipramine) or mTOR signaling (by rapamycin) markedly decreased Th17 generation of CD4+CD39+CD161+ T cells, in contrast to cells treated with vehicle only (Fig. 5F). Moreover, inhibition of ASM, mTOR, or STAT3 (by WP1066) also downregulated CD39 expression in Th17-deviated CD4+ T cells (Supplemental Fig. 3) compared with vehicle-treated cells. These data show reciprocal interactions between ASM-STAT3/mTOR signals and CD39 expression in deviating CD4+ T cells toward the Th17 destination.

Taken together, these data indicate that CD39 and CD161 pudently exhibit associations that further enhance ASM activity, thereby ultimately controlling Th17 generation of CD4+ T cells that is mediated by mTOR- and STAT3-signaling cascades.

**Knockdown of ASM or ASM inhibition abrogates Th17 generation**

Because of the pivotal role of ASM in Th17 generation, as suggested above with the use of the pharmacological inhibitor imipramine, we next tested shRNA treatment to “knockdown” ASM levels in healthy blood CD4+ T cells (Fig. 6A). In these studies, two shRNAs, termed sh-1 and sh-3, were used to assess the effect of direct downregulation of ASM on downstream STAT3 signals and Th17 generation.

Compared with sh-C, knockdown of ASM by sh-1 or sh-3 significantly inhibited ASM bioactivity, as determined by ceramide production (Fig. 6B), STAT3 signaling (Fig. 6C), and decreased IL-17 levels (Fig. 6D). These data further suggest ASM links to STAT3 signaling that might be associated with Th17 generation.

Next, CD4+ T cells from lamina propria of patients with active CD were treated with imipramine, followed by evaluation of all aspects of Th17 differentiation, as above. Ceramide production was substantially abrogated by imipramine (Fig. 7A), concurrently with decreased activation of Akt-mTOR and STAT3 signaling (data not shown) and diminished levels of intracellular IL-17 and IFN-γ (Fig. 7B, 7C). These data infer that blockade of ASM bioactivity might have therapeutic potential in Th17-dominant inflammatory states, such as in CD.
**Discussion**

Th17 cells are novel and important immunomodulatory cells in the mammalian immune system. These proinflammatory lymphoid cells are crucial in mediating host defense against both intracellular and extracellular pathogens. Moreover, Th17 cells were implicated in the pathogenesis of CD and other inflammatory conditions (5, 6). We (10) and other investigators (42, 43) showed that the likely inhibition of Th17 responses can lead to amelioration of inflammation in experimental models of intestinal disease and other inflammatory disorders. Hence, more detailed molecular insights into Th17 generation are of paramount importance in developing new therapies to supplement the current clinical approaches in CD.

Specifically identifying Th17 cells in patients has remained problematic, which is due, in part, to the lack of reliable surface...
markers. Recently, CD161 was identified as a relatively specific surface marker for human Th17 cells, with application specifically to CD (14, 15). However, CD161 expression is also associated with NK cells and, as we demonstrate in this study, CD161 alone is insufficient for depicting and characterizing these Th17 cells and associated responses.

We note in this article that dual expression of CD39 and CD161 more clearly defines Th17 lineages of human CD4+ T cells. Furthermore, we also show that blood and lamina propria levels of IL-17 and IFN-γ (72 h) were evaluated by TLC (A) and FACS (B), respectively. CD39+CD161+ and CD39−CD161− CD4+ T cells were stimulated with Th17-driving conditions for 12 h or were left unstimulated (0 h). Cell lysates were used for coimmunoprecipitation against anti-ASM Ab, followed by Western blot using the probes indicated. TLC analysis of ceramide production by 48-h Th17-deviated CD4+ T cell subsets (D) or CD4+CD39+CD161+ T cells stimulated with cross-linked anti-CD39 Ab (4 μg/ml), anti-CD161 Ab (4 μg/ml), or both for different times (E). Mouse anti-human IgG (4 μg/ml) served as control. (F) Effects of imipramine or rapamycin on IL-17 and IFN-γ production by CD4+CD39+CD161+ T cells at 72 h post-Th17 polarization. Data are mean ± SEM of three or four independent experiments. *p < 0.05 versus the other three groups.

FIGURE 5. Direct interaction of ASM with CD39 and CD161 mediates Th17 generation through mTOR/STAT3 signaling. (A and B) CD4+ T cells were expanded by Th17-differentiation conditions, either in the presence of vehicle or imipramine (20 μM). Production of ceramide (48 h) and intracellular levels of IL-17 and IFN-γ (72 h) were evaluated by TLC (A) and FACS (B), respectively. (C) CD39+CD161+ and CD39−CD161− CD4+ T cells were stimulated with Th17-driving conditions for 12 h or were left unstimulated (0 h). Cell lysates were used for coimmunoprecipitation against anti-ASM Ab, followed by Western blot using the probes indicated. TLC analysis of ceramide production by 48-h Th17-deviated CD4+ T cell subsets (D) or CD4+CD39+CD161+ T cells stimulated with cross-linked anti-CD39 Ab (4 μg/ml), anti-CD161 Ab (4 μg/ml), or both for different times (E). Mouse anti-human IgG (4 μg/ml) served as control. (F) Effects of imipramine or rapamycin on IL-17 and IFN-γ production by CD4+CD39+CD161+ T cells at 72 h post-Th17 polarization. Data are mean ± SEM of three or four independent experiments. *p < 0.05 versus the other three groups.
In this study, we note that human CD39\(^{+}\)CD161\(^{+}\) CD4\(^{+}\) T cells express high levels of ecto-NTPDase as well and that extracellular ADO is extremely short lived, as a result of higher rates of ADA-mediated catabolism. Supplementing purine products ATP, AMP, or ADO did not affect expansion of CD39\(^{+}\) CD161\(^{+}\) CD4\(^{+}\) Th17 cells in these models, given the rapid

**FIGURE 6.** Knockdown of ASM blocks STAT3 signals and Th17 generation. (A) Western blotting of ASM expression in healthy blood CD4\(^{+}\) T cells after inhibition of ASM using lentiviral shRNAs. Sh-C and ASM knockdown (sh-1 and sh-3) healthy peripheral blood CD4\(^{+}\) T cells were stimulated with anti-CD3/CD28 beads under the defined Th17 differentiation conditions, followed by representative TLC analyses of ceramide at 2 h (B), flow cytometry analysis of intracellular p-STAT3 at 3 h (C), and intracellular IL-17 and IFN-\(\gamma\) expression at 96 h (D). Data are mean ± SEM of three independent experiments. *\(p < 0.05\) versus the sh-C group.

**FIGURE 7.** ASM inhibition abrogates in vitro generation of Th17 cells using lamina propria CD4\(^{+}\) T cells purified from patients with CD. Lamina propria CD4\(^{+}\) T cells isolated from patients with active disease were stimulated with anti-CD3/CD28 activation beads in the presence of vehicle or imipramine (20 \(\mu\)M), followed by TLC analysis of ceramide levels at 2 h (\(n = 3\)) (A) and flow cytometry determination of intracellular IL-17 and IFN-\(\gamma\) expression at 5 h (\(n = 6\)) (B and C). *\(p < 0.05\), **\(p < 0.01\) versus vehicle.


