Accumulation of Regulatory T Cells and Chronic Inflammation in the Middle Ear in a Mouse Model of Chronic Otitis Media with Effusion Induced by Combined Eustachian Tube Blockage and Nontypeable Haemophilus influenzae Infection

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Nontypeable Haemophilus influenzae (NTHi) is associated with chronic otitis media (COM). In this study, we generated a murine model of COM by using eustachian tube (ET) obstruction and NTHi (10^7 CFU) inoculation into the tympanic bulla, and we investigated the relationship between regulatory T cells (Treg) and chronic inflammation in the middle ear. Middle ear effusions (MEEs) and middle ear mucosae (MEM) were collected at days 3 and 14 and at 1 and 2 months after inoculation. Untreated mice served as controls. MEEs were used for bacterial counts and to measure the concentrations of cytokines. MEM were collected for histological evaluation and flow cytometric analysis. Inflammation of the MEM was prolonged throughout this study, and the incidence of NTHi culture-positive MEE was 38% at 2 months after inoculation. The levels of interleukin-1β (IL-1β), tumor necrosis factor alpha, IL-10, and transforming growth factor β were increased in the middle ear for up to 2 months after inoculation. CD4+ CD25+ FoxP3+ Treg accumulated in the middle ear, and the percentage of Treg in the MEM increased for up to 2 months after inoculation. Treg depletion induced a 99.9% reduction of bacterial counts in MEEs and also significantly reduced the ratio of NTHi culture-positive MEE. The levels of these cytokines were also reduced in MEEs. In summary, we developed a murine model of COM, and our findings indicate that Treg confer infectious tolerance to NTHi in the middle ear.

Chronic otitis media (COM), including OM with effusion (OME) and recurrent OM, is characterized by clinical evidence of OM and resolution of middle ear effusion (MEE) between episodes. OME represents a spectrum of chronic disease states, ranging from serous to mucoid OM, and is associated with hearing loss, delayed speech development, permanent middle ear damage, and mucosal changes. Although COM remains a common problem in pediatric populations, its etiology and pathogenesis are not fully understood. Eustachian tube (ET) dysfunction is considered to be the underlying pathophysiologic event leading to most cases of OME in children. The most frequent causes of ET dysfunction include upper respiratory infections, adenoid tissue hypertrophy, and cleft palate. In bacterial infection of COM, the majority of cases are caused by Gram-negative bacteria, whereas in acute OM, Gram-positive bacteria are also frequently isolated. Lipopolysaccharides are a component of Gram-negative bacteria that have been detected in human MEEs, and their levels are significantly higher in children with chronic OME than in children with acute OME. Lipopolysaccharides alone have also been shown to induce mucosal inflammation with the accumulation of effusion in the middle ears of animal models. Only 30% of MEEs from COM children yielded an unequivocally positive culture for aerobic bacteria. According to previous reports, components from Gram-negative bacteria are thought to induce COM in humans. However, recently, COM was reportedly associated with a persistent bacterial infection, and biofilm structures were identified in 92% of middle ear mucosa (MEM) biopsy specimens from children with COM. On the basis of these reports, ET dysfunction and a persistent bacterial infection are closely associated with the pathogenesis of COM. Previously, we established a murine model of COM with effusion by ET blockage and endotoxin inoculation into the bulla. COM mice produced by ET blockage showed persistent severe MEE with mild inflammatory responses. COM mice produced by ET blockage and endotoxin inoculation were associated with moderate inflammation and the production of mucoid MEE accompanied by histological changes with inflammatory cell infiltration, especially lymphocytes, and cytokine production in the middle ear cavity. However, this mouse model of COM does not reflect the recent findings of COM in children. Therefore, we need to examine a COM model with ET dysfunction and persistent bacterial infection to investigate the pathogenesis of COM. Nontypeable Haemophilus influenzae is thought to be the chief pathogen in both acute OM and COM. In addition, NTHi is an important pulmonary bacterial pathogen associated with recurrent and persistent lower respiratory tract infections in patients with chronic obstructive pulmonary disease, which affects 16 million people and is the fourth leading cause of death in the United States. NTHi is a commensal to opportunistic pathogen that is highly adapted to the human airway.
strains can persist within the airway for lengthy periods, during which their carriage is mostly asymptomatic in healthy individuals (15). However, when host mucosal clearance mechanisms are compromised or impaired, NTHi can cause an array of airway infections (14). Recently, biofilm formation by NTHi has been defined for persistence and pathogenicity in chronic airway infections to some degree (16). However, in cases of persisting NTHi infection in the upper and lower airways, little is known about the immunological responses from the viewpoint of the host immune reaction against NTHi.

Regulatory T cells (Treg), also known as suppressor T cells, consist of a specific subpopulation of cells that functionally suppress the activation of the immune system and maintain immune tolerance to self-antigens (17). The gene coding for the Treg cell transcription factor forkhead box P3 (FoxP3) is a master gene for the differentiation and function of Treg (18,19). Treg consist of 2 major subsets: natural Treg and adaptive or induced Treg. Natural Treg derived from the thymus are considered the classic Treg, while inducible Treg develop in the periphery in response to self or tumor antigens by converting naive CD4+ T cells into Treg (20). Normally, when these two subsets of Treg are depleted, various organ-specific autoimmune diseases occur spontaneously (17). Recent studies, however, indicate that Treg can also be activated and expanded against bacterial, viral, and parasite antigens in vivo (21). Such pathogen-specific Treg are known to prevent infection-induced immunopathology but may also increase the load of infection and prolong pathogen persistence by suppressing protective immune responses. Clarifying whether Treg play a protective or detrimental role in the immune cellular response during upper respiratory tract infection with NTHi could help establish a treatment strategy for COM.

In this study, we explored a murine model of COM with persistent NTHi infection by using ET blockage and NTHi inoculation and investigated the immune cellular response, specifically Treg, and also the effect of the depletion of Treg on chronic middle ear infection.

MATERIALS AND METHODS

Animals. BALB/c mice were purchased from Kyudo Japan (Fukuoka, Japan). All mice were maintained in a pathogen-free facility until they were 3 weeks old, at which time they were used for the experiments. A total of 136 animals were used in this study. All experiments were approved by the Committee on Animal Experiments of Oita University.

Middle ear challenge with live NTHi and induction of experimental COM. Strain 76 of NTHi, which was isolated from the nasopharynx of a patient with OME at Oita University, was used for the middle ear challenge, and the bacterial suspension was injected into the right tympanic cavity, as described previously (11). NTHi was grown on chocolate agar at 37°C under 5% CO2, for 16 h. The bacterial concentration was determined by optical density at a wavelength of 600 nm, and then a bacterial suspension was prepared to a concentration of 1.0 × 107 CFU/ml in phosphate-buffered saline (PBS) and stored on ice. The bacterial concentration was confirmed by colony counting after an overnight incubation. In brief, with the animal under anesthesia induced with 0.6 mg/ml ketamine hydrochloride, an incision was made in the submandibular skin, the right inferior bulla was exposed, the pharyngeal part of the ET near the tympanic bulla was cut, and a gelatin sponge was inserted into the ET (ET blockage). Subsequently, 2 microholes were made in the bulla with a 27-gauge needle. A micropipette was inserted into one of the holes, and 10 μl (1.0 × 106 CFU) of live NTHi suspension was inoculated slowly. Mice without both ET blockage and NTHi inoculation served as controls based on our previous study (11). The mice were monitored otomicroscopically to confirm the presence of MEE and tympanic membrane changes. Untreated mice were used as controls according to the same procedure for these experiments. Another set of model mice was made for flow cytometric analysis and real-time reverse transcription-PCR (RT-PCR).

Collection of samples and treatment. On day 3, day 14, 1 month, and 2 months after infection, 7 to 10 mice from each group were killed under deep anesthesia by intraperitoneal administration of a pentobarbital solution. Untreated mice were also killed as controls. The mice were monitored otomicroscopically to confirm the presence of MEE and tympanic membrane changes. The characteristics of MEE were recorded and graded by increasing order of inflammation according to the severity of COM, as follows: 0, gray and translucent TM without MEE (normal); 1, gray and opaque TM with serous MEE or mucoid MEE; 2, yellow and opaque TM with purulent MEE; and 3, opaque TM with pultaceous MEE. These grading scales are referred to and modified from Giebink et al. (22). MEE samples were obtained by myringotomy at the time of decapitation, and the middle ears were washed with 200 μl physiologic saline. MEEs were diluted serially, and the diluted samples were plated on chocolate agar. Bacterial colonies were counted after an overnight incubation to measure the rate of NTHi clearance from the middle ear. NTHi was identified by standard bacteriologic techniques, including Gram staining and determination of their V and X growth factor requirements. MEEs were centrifuged, and the supernatants were collected and then stored at −80°C for cytokine analysis by enzyme-linked immunosorbent assay (ELISA; described below).

Histological preparation. After the mice were euthanized and MEE samples were collected, they were fixed by intracardiac perfusion with 10% neutral buffered formalin or periodate lysine paraformaldehyde for histologic evaluation. The mice were then decapitated, and their heads were immersed in the same fixative used for perfusion for 6 h and decalcified in 0.12 M EDTA (pH 7.4) for 2 weeks at 4°C. For hematoxylin–eosin (H&E) and immunohistochemical staining, tissues were dehydrated through a graded series of ethanol, cleared in xylene, and embedded in paraffin.

Histological evaluation. Serial paraffin sections (6 μm) that contained the tympanic bulla were prepared. Mucosal thickness and the number of inflammatory cells in the MEM were measured using ImageJ, a public domain, Java-based image processing program developed at the National Institutes of Health (Bethesda, MD; http://rsweb.nih.gov/ij/). Sections from the largest middle ear bulla around the tympanic orifice were observed by light microscopy, and histological images (magnification ×400) were captured; mucosal thickness and the number of inflammatory cells in the MEM were measured and analyzed at the anterior, posterior, medial, and lateral parts of the bulla. Three mice were used for this experiment.

Immunohistochemistry for CD4+ and FoxP3+ cells. Paraffin sections of the middle ear cavity around the tympanic orifice in the ET were used for the detection of CD4+ and FoxP3+ cells. After deparaffinization, the specimens were treated with 3% H2O2 for 20 min at room temperature. Sections were exposed to 5% normal goat serum in PBS for 30 min and then incubated with a rabbit anti-mouse CD4 antibody (Novus Biologicals, Littleton, CO) and a rabbit anti-mouse FoxP3 antibody (Novus Biologicals) diluted 1:500 in 1% bovine serum albumin (BSA)–PBS for 12 h at room temperature. After being rinsed with PBS, the sections were incubated with a biotinylated goat anti-rabbit antibody (1:2,000) in 1% BSA–PBS for 6 h at room temperature. After being rinsed with PBS, the sections were incubated with ABC reagent (Vector Laboratories, Burlingame, CA) for 1 h and developed in 0.05% 3’,3’-diaminobenzidine–0.01% H2O2 substrate medium in 0.1 M phosphate buffer for 8 min.

Detection of cytokines in MEE by ELISA. Assays for tumor necrosis factor alpha (TNF-α), interleukin-1β (IL-1β), IL-10, and transforming growth factor β (TGF-β) were performed with a commercially available mouse enzyme-linked immunosorbent assay (ELISA) kit (R&D Systems, Inc., Minneapolis, MN), according to the manufacturer’s instructions. All samples were run with negative and positive controls. Standard curves

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Infection and Immunity

iai.asm.org 357

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were generated from known concentrations of cytokines provided by the manufacturer. Optical densities were read with a microtiter ELISA plate reader (Sanko Junyaku Co., Ltd., Tokyo, Japan).

**Flow cytometric analysis of T cells in mononuclear cells infiltrating the MEM.** To analyze the cellular phenotypes of lymphocytes in the middle ear responding to chronic inflammation, mononuclear cells (MNCs) were isolated and stained with antigen-specific antibodies conjugated with a fluorescent label. Another set of COM mice was made, and MNCs from the middle ear were harvested from groups of 5 mice at each time point. For the isolation of MNCs from the MEM, we established a dissociation method (23). Briefly, the skin was removed carefully from the head region, and the bulla was isolated from the temporal bone. Then, the bulla was dissected, and the MEM was detached from the bone under microscopy. MNCs were isolated from the MEM after enzymatic digestion at 37°C for 10 min with 0.5 mg/ml collagenase type IV (Sigma, St. Louis, MO). Approximately 0.5 × 10^6 to 1.0 × 10^6 cells were used for flow cytometry. To detect Treg in the MEM, a fluorescein isothiocyanate (FITC)-conjugated anti-CD4 monoclonal antibody (MAB) (17A2; BD Bioscience, San Diego, CA), phycoerythrin (PE)-conjugated anti-CD8 MAB (53-6.7; BD Bioscience), and peridinin chlorophyll protein (PerCP)-conjugated anti-CD3 MAB (145-2C11; BD Bioscience) were used to analyze T cell subsets in 3-color analysis, and an FITC-conjugated anti-CD4 MAB, phycoerythrin-conjugated anti-FoxP3 MAB (Milenyi Biotec, Auburn, CA), and PerCP-conjugated anti-CD25 MAB (PC61; BD Bioscience) were used to analyze Treg in 3-color analysis. Intracellular FoxP3 staining was performed by using a FoxP3 staining buffer set (Milenyi Biotec). The corresponding isotype control antibodies were used as negative controls of background staining. The immunofluorescent intensities of Treg were analyzed using CellQuest software (Becton Dickinson, Mountain View, CA), where a lymphocyte gate was defined by forward and side scatter parameters. This experiment was performed independently 3 times, each from a pool of 5 mice.

**Detection of FoxP3 mRNA and cytokine mRNA in MNCs from MEM by real-time RT-PCR.** To elucidate the inflammatory responses in the middle ear after bacterial infection, real-time RT-PCR was performed to assess the expression of FoxP3 and cytokine mRNA in the MNCs that had infiltrated the MEM. These MNCs were isolated from 5 mice at the same time as those used for flow cytometry. For the detection of FoxP3, IL-10, TGF-β, IL-1β, and TNF-α mRNAs, total RNA was extracted from the MNCs that had infiltrated the MEM. The primers and probes for FoxP3 mRNA used in this study were obtained from Applied Biosystems (Foster City, CA). Total cellular RNA extraction and first-strand cDNA synthesis were performed using an RNA extraction and reverse transcription kit (Qiagen, Tokyo, Japan). The total concentration of RNA used for this assay in each group was adjusted to 0.25 μg. Each cDNA sample was then used as a template in a PCR amplification mixture containing forward and reverse primers for FoxP3, IL-10, TGF-β, IL-1β, and TNF-α, forward and reverse primers for 18S rRNA (internal control), and TaqMan Universal PCR master mix (Applied Biosystems). PCR amplifications for the target cytokines and internal control 18S rRNA were performed in a single well of a capped 96-well optical plate. Reaction mixtures were subjected to the following amplification scheme: 1 cycle at 50°C for 2 min and 1 cycle at 95°C for 10 min, followed by 40 cycles at 95°C for 15 s (denaturation) and 60°C for 1 min (annealing and extension). Real-time RT-PCR data were analyzed using a StepOnePlus real-time PCR system (Applied Biosystems). Final mRNA quantities were derived using the comparative threshold cycle (Ct) method. These quantities are expressed as the fold difference relative to a calibrator cDNA (from mice that had no treatment as a control) prepared in parallel with the experimental cDNAs. Data are expressed as the arithmetic means of the percentage of positive cells from 3 independent experiments (± standard deviation [SD]), each from a pool of 5 mice.

**In vivo depletion of Treg by administration of an anti-CD25 MAB.** In vivo depletion of Treg is a means of studying the role of these subpopulations in the effector phases of particular in vivo immune responses in COM. To deplete Treg, an anti-CD25 MAB (PC61) was used. Onizuka et al. reported that the administration of anti-CD25 MAB (PC61) caused a reduction in the number of CD4+ CD25+ cells in peripheral lymphoid tissues, and they also reported that 125 μg was an effective dose to reduce Treg (24).

Experimental COM mice were made as described above. The mice were injected intraperitoneally with 125 μl anti-CD25 MAB (PC61.5 at 125 μg/mouse; eBioscience, San Diego, CA) or PBS as a control at 2, 4, and 6 weeks after ET blockage and NTHi inoculation. In a preliminary study, we administered the anti-CD25 antibody once, twice, and three times, and administration of the antibody three times was found to be more effective and stable than administration once and twice (data not shown). At 2 months after treatment, 8 mice from each group were killed under deep anesthesia by intraperitoneal administration of a pentobarbital solution. The same evaluations as in our earlier experiment were performed. To investigate the changes of Treg after treatment with the anti-CD25 MAB, the spleen (SPL) and cervical lymph nodes (CLNs) were collected for flow cytometric analysis of T cells in mononuclear cells infiltrating the tissues, and they also reported that 125 μg was an effective dose to reduce Treg (24).

**RESULTS**

**Middle ear findings of COM.** In the control group, no mice had MEE. In the COM group, MEE was induced in all mice that received an NTHi injection and ET obstruction (A). All MEEs were purulent at day 3 after inoculation, and 71% were purulent at 2 weeks. At 1 month (1M) after inoculation, the ratio of mucoid effusions increased to 70%. At 2 months (2M) after inoculation, 50% of MEEs had become serous and 38% of effusions had become pultaceous (B). ND, not detected.
effusions was increased to 70% (7/10). At 2 months after inoculation, 50% (4/8) of MEEs had become serous, and 38% (3/8) of effusions had become purulent. The severity score of COM at all time points examined showed a significant difference compared to the control group (Table 1).

**Table 1** Severity scores of OM

<table>
<thead>
<tr>
<th>Group</th>
<th>Severity score of OM</th>
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<tbody>
<tr>
<td>Control</td>
<td>0 ± 0</td>
</tr>
<tr>
<td>Day 3</td>
<td>2 ± 0*</td>
</tr>
<tr>
<td>Day 14</td>
<td>1.7 ± 0.5*</td>
</tr>
<tr>
<td>1 mo</td>
<td>1.2 ± 0.4*</td>
</tr>
<tr>
<td>2 mo</td>
<td>1.8 ± 1.0*</td>
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</table>

* Seven to 10 mice from each group were monitored otomicroscopically at day 3, day 14, 1 month, and 2 months after injection with NTHi. The severity score of OM was measured according to tympanic membrane (TM) color and opacity and the presence of middle ear effusion (MEE), as follows: 0, gray and translucent TM without MEE (normal); 1, gray and opaque TM with serous or mucoid MEE; 2, yellow and opaque TM with purulent MEE; and 3, opaque TM with pultaceous MEE. The severity score is expressed as the mean ± SD. * P < 0.05 compared to the control by Mann-Whitney test.

Bacterial clearance and inflammatory cell infiltration. Bacterial counts in MEEs decreased gradually during the course of COM. NTHi concentrations were highest at day 3 after inoculation and decreased gradually but were sustained for up to 2 months after inoculation. Inflammatory cell infiltration contained polymorphonuclear cells, and these inflammatory responses were strong up to day 14. From 1 to 2 months after inoculation, mucosal thickening and inflammatory cell infiltration continued with lymphocyte migration in the MEM. Mucosal thickness (Fig. 3B) and the number of inflammatory cells (Fig. 3C) in the middle ear were analyzed using ImageJ software. Compared to the control group, mucosal thickness of the middle ear increased significantly throughout the experiments (mean thickness of MEM ± SD in control versus day 3, day 14, 1 month, and 2 months, 15 ± 9 versus 35 ± 11, 59 ± 20, 48 ± 31, and 41 ± 8 nm, respectively; P < 0.05), peaking at day 14. Inflammatory cell infiltration also increased (mean number of inflammatory cells in the MEM ± SD in control versus day 3, day 14, 1 month, and 2 months, 19 ± 11 versus 161 ± 58, 104 ± 54, 124 ± 93, and 72 ± 27 cells/10,000 nm², respectively; P < 0.05). The inflammatory responses due to NTHi inoculation continued for up to 2 months in the middle ear.

**Table 2** Range of NTHi infection and rate of MEE

<table>
<thead>
<tr>
<th>Group</th>
<th>NTHi infection, CFU/mouse</th>
<th>Rate of culture positive MEE, no. positive/total (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0</td>
<td>0/10 (0)</td>
</tr>
<tr>
<td>Day 3</td>
<td>2.4 × 10³–1.1 × 10⁶</td>
<td>10/10 (100)</td>
</tr>
<tr>
<td>Day 14</td>
<td>0 × 10⁶–1.3 × 10⁶</td>
<td>4/7 (57)</td>
</tr>
<tr>
<td>1 mo</td>
<td>0 × 10⁶–1.6 × 10⁶</td>
<td>4/10 (40)</td>
</tr>
<tr>
<td>2 mo</td>
<td>0 × 10⁶–5.4 × 10⁶</td>
<td>3/8 (38)</td>
</tr>
</tbody>
</table>

MEEs were collected from 7 to 10 mice at day 3, day 14, 1 month, and 2 months after injection with NTHi and ET blockage.
Cytokine levels in MEE and cytokine mRNA expression in MNCs in the middle ear. Figure 6 shows the cytokine levels in MEE. According to ELISA, in the MEE samples, the levels of IL-1β and TNF-α in the COM group were higher than in the control group. The differences in IL-1β concentration reached statistical significance throughout the experiments (mean concentration of IL-1β SD in control versus day 3, day 14, 1 month, and 2 months, 4.5 ± 10 versus 297.6 ± 163.8, 118.9 ± 59.5, 106.3 ± 79.6, and 75.6 ± 87.7 pg/ml, respectively; P < 0.05). The TNF-α concentration was also significantly higher in the COM group at day 3, day 14, and 1 month after inoculation (mean concentration of TNF-α SD in control versus day 3, day 14, and 1 month, 0 ± 0 versus 143.7 ± 79.6, 52.3 ± 64.6, and 32.7 ± 43.3 pg/ml, respectively; P < 0.05). Similarly, the IL-10 concentration in the MEE samples was significantly higher in the COM group at day 3, day 14, and 2 months after inoculation (mean concentration of IL-10 SD in control versus day 3, day 14, and 2 months, 0 ± 0 versus 30 ± 26.5, 21 ± 15.2, and 16 ± 10 pg/ml, respectively; P < 0.05), as was the TGF-β concentration at day 3, day 14, 1 month, and 2 months after inoculation (mean concentration of TGF-β SD in control versus day 3, day 14, 1 month, and 2 months, 0 ± 0 versus 186.5 ± 260.5, 457.6 ± 421, 418.6 ± 367.2, and 359.3 ± 333.8 pg/ml, respectively; P < 0.05).

To define the induction of cytokine transcription in MNCs from the MEM, the expression of the IL-1β, TNF-α, IL-10, and TGF-β genes was evaluated by real-time RT-PCR analysis (Fig. 7). The FoxP3, IL-1β, TNF-α, IL-10, and TGF-β genes were significantly more induced by inoculation with NTHi than in the controls.

Effect of Treg depletion on immune responses in COM. Figure 8A shows the results of flow cytometry for CD4+ CD25+ T cells. There were a large number of positive cells at 3 and 2 months after inoculation (1 M and 2 M, respectively). (B) Flow cytometric analysis for T cell subsets in the MEM. The percentage of CD4+ CD8+ T cells was increased in the middle ear from day 3 to 2 months after inoculation. However, the percentage of CD4+ T cells was always greater than that of CD8+ T cells in the middle ear throughout the experiment.
FoxP3⁺ T cells with anti-CD25 MAb or PBS treatment. Compared with the PBS-treated mice, in which the percentages of Treg were 7.0% in the SPL, 8.5% in the CLNs, and 32.4% in the MEM, anti-CD25 MAb-treated mice had a lower percentage of Treg in each tissue (1.3% in the SPL, 1.3% in the CLNs, and 18.2% in the MEM). In terms of bacterial counts in MEE, PBS-treated mice had higher counts than the anti-CD25 MAb-treated mice; a 99.9% reduction was seen in the anti-CD25 MAb-treated mice (Fig. 8B). Figure 8B also shows that there was a significantly higher percentage of NTHi culture-positive MEE in the PBS-treated mice than in the anti-CD25 MAb-treated mice (rate of NTHi culture-positive MEE in PBS-treated versus anti-CD25 MAb-treated mice, 62.5% [5/8] versus 12.5% [1/8], respectively; chi-square test, \( P < 0.05 \)).

The cytokine levels in MEE, the levels of all cytokines tested, including proinflammatory and immunosuppressive cytokines, were reduced by anti-CD25 MAb treatment. Especially, IL-1 was reduced significantly between the two groups (Fig. 8C).

### DISCUSSION

In this study, we developed an experimental mouse model of COM by inoculating NTHi into the middle ear bulla and using ET blockage and then studied the local immune responses of the middle ear against NTHi. We found that middle ear inflammation, such as mucosal thickening and inflammatory cell infiltration, continued for at least 2 months after inoculation of the middle ear, and NTHi was detected using a bacterial culture method in the middle ear cavity in 40% of our COM model mice for up to 2 months after the challenge. This is the first report showing chronic inflammation can be induced in the middle ear with NTHi culture-positive MEEs. To our knowledge, these are also the first detailed findings regarding the inflammatory responses of the middle ear in a murine COM model. This is in contrast to other animal models using NTHi, such as the chinchilla and rat. Green et al. (25) reported that inoculation of viable NTHi induced acute

### TABLE 3

Average frequencies of CD4, CD8, and regulatory T cells in the middle ear

<table>
<thead>
<tr>
<th>Group</th>
<th>% CD4 ± SD</th>
<th>% CD8 ± SD</th>
<th>CD4/CD8 ratio</th>
<th>% FoxP3⁺</th>
<th>% CD25⁺ FoxP3⁺</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>42.8 ± 15.6</td>
<td>17.7 ± 11</td>
<td>2.7 ± 1.2</td>
<td>18.5 ± 1.3</td>
<td>12.2 ± 2.4</td>
</tr>
<tr>
<td>Day 3</td>
<td>40.2 ± 8.3</td>
<td>10.0 ± 2.1</td>
<td>4.0 ± 0.2</td>
<td>15.4 ± 4.5</td>
<td>10.6 ± 1.8</td>
</tr>
<tr>
<td>Day 14</td>
<td>30.8 ± 8.6</td>
<td>12.8 ± 1.5</td>
<td>2.4 ± 0.6</td>
<td>45.0 ± 4.0⁺</td>
<td>38.3 ± 4.9⁺</td>
</tr>
<tr>
<td>1 mo</td>
<td>33.7 ± 11.0</td>
<td>8.4 ± 4.7</td>
<td>4.6 ± 1.6</td>
<td>40.3 ± 1.8⁺</td>
<td>34.7 ± 2.0⁺</td>
</tr>
<tr>
<td>2 mo</td>
<td>33.6 ± 8.6</td>
<td>7.3 ± 2.3</td>
<td>4.8 ± 1.2</td>
<td>34.4 ± 2.5⁺</td>
<td>27.8 ± 2.3⁺</td>
</tr>
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</table>

- MNCs were collected from 5 mice. This experiment was performed independently 3 times.
- *⁺, \( P < 0.05 \) compared to control by Mann-Whitney test.

**FIG 5** (A) Immunohistochemistry for FoxP3⁺ cells in MEM. A rabbit polyclonal anti-mouse FoxP3 MAb was used. FoxP3⁺ cells were also seen at day 14. The distribution of positive cells was sustained at 1 and 2 months (1 M and 2 M, respectively). (B) Percentage of CD4⁺ FoxP3⁺ T cells in the MEM. The percentage of CD4⁺ FoxP3⁺ T cells increased in the middle ear from day 14 to 2 months after inoculation compared to in the control group. (C) Cell fraction of FoxP3⁺ T cells. The number of CD25⁺ T cells also increased after inoculation, and the number of CD25⁺ FoxP3⁺ T cells in the MEM was greater than that of CD25⁺ FoxP3⁺ T cells among the CD4⁺ T cells at all time points examined. One representative experiment out of 3 is shown.
by the inhibition of IL-2 gene expression, modulation of costimulatory molecules on antigen-presenting cells, interaction of lymphocyte activation gene 3 with major histocompatibility complex (MHC) class II molecules, secretion of immunosuppressive cytokines such as IL-10 and TGF-β, induction of tryptophan catabolism through cytotoxic T lymphocyte-associated antigen 4 (CTLA-4), and cytotoxicity (28). Previously, CD4+CD25+FoxP3+ Treg were considered natural Treg, which are physiologically produced by the normal thymus as a functionally mature and distinct population, and their development and function depend on the expression of the transcription factor FoxP3 (17, 18, 28). However, CD4+CD25−naïve T cells can be converted to CD4+CD25+Treg by the TGF-β-induced expression of FoxP3 (29).

Adaptive or induced CD4+ Treg include two types of cells: T regulatory type 1 (Tr1) cells and Th3 Treg. Tr1 cells suppress antigen-specific immune responses by producing high levels of IL-10, and Tr1 cells can transiently express FoxP3 upon activation (30, 31). Th3 cells secrete TGF-β and IL-10 and express FoxP3 (31, 32). Th3 cells play a crucial role in inducing and maintaining peripheral tolerance by driving the differentiation of antigen-specific FoxP3 regulatory cells in the periphery by producing high levels of TGF-β (33). In this study, not only inflammatory cytokines (IL-1β and TNF-α) but also immunosuppressive cytokines (IL-10 and TGF-β) increased in the middle ear at the protein and mRNA level after inoculation compared to the control group. Although we did not distinguish natural Treg from adaptive or induced Treg in the CD4+FoxP3+ Treg, these Treg are considered to be related to the persistent inflammation in NTHi infection.

Treg are also known to play an important role in controlling infectious disease, including viruses, parasites, and bacteria, and Treg play two different key immunological roles—protective or detrimental—in host defense.

Treg play protective roles against certain viral pathogens. Lund et al. (34) showed an accelerated fatal infection with increased herpes simplex virus loads in the vaginal mucosa and central nervous system after ablation of Treg. Similarly, FoxP3+ Treg ablation before West Nile virus infection in mice led to an increase of disease severity, which was associated with higher viral loads in the brain and spinal cord (35). As for bacterial infections, Treg reportedly continue to play a detrimental role in host defense. To investigate the relationship between Treg and pregnancy-associated infection susceptibility, Rowe et al. (36) demonstrated that expan-

OM in the chinchilla and that the bacterial cells existed for 14 days of the challenge. Melhus et al. (26) found that middle ear infections were resolved within 8 days in a rat model. These animal models were not treated with an ET block, and middle ear infections were limited to several weeks. On the basis of these findings and compared with previous studies, this mouse model treated with both an ET block and NTHi inoculation in the middle ear may make a useful model for studying the role of chronic inflammation in a persistent NTHi infection.

Recently, the characterization of T cell subsets has accelerated the understanding of inflammatory and humoral immune responses in part by unveiling the enormous plasticity within these subsets. The balance in T helper subsets as observed in healthy mucosa is changed in inflamed mucosa. According to flow cytometric analysis of the middle ear in our model, the CD4/CD8 ratio was increased from day 3 to 2 months after inoculation compared to the control group, as Derycke et al. (27) reported the CD4/CD8 ratio was increased depending on the severity of inflammation in chronic rhinosinusitis. This result showed that chronic inflammation occurred in the middle ear for a period of 2 months. Furthermore, the number of CD4+FoxP3+ Treg was significantly increased in the COM group from 2 weeks to 2 months after inoculation compared with the control group. In the CD4+FoxP3+ Treg, CD4+CD25+FoxP3+ Treg were mainly increased in the MEM. The immunohistochemistry findings also supported the results of these flow cytometric analyses. CD4+CD25+FoxP3+ Treg have a primary effect on T cells and/or dendritic cells

FIG 6 Levels of cytokines in MEE. In the MEE samples, the levels of IL-1β and TNF-α were higher in the COM group than in the control group. The differences in the concentration of IL-1β were statistically significant throughout the experiments. The TNF-α concentration was also significantly higher in the COM group at day 3, day 14, and 1 month (1 M) after inoculation. Similarly, the concentration of IL-10 and TGF-β in the MEE samples increased significantly in the COM group for up to 2 months (2 M) after inoculation. *, P < 0.05 compared to the control by Mann-Whitney test.

FIG 7 Quantification of the levels of mRNA transcripts. The FoxP3, IL-1β, TNF-α, IL-10, and TGF-β genes were significantly more induced by inoculation with NTHi than the controls throughout the experiment. 1 M, 1 month; 2 M, 2 months; *, P < 0.05 compared to the control by Mann-Whitney test.
viation of immune-suppressive FoxP3+ Treg, which occurs physiologically during pregnancy or when experimentally induced in transgenic mice, caused enhanced susceptibility to prenatal pathogens, including *Listeria* and *Salmonella* species. Reciprocally, infection susceptibility was uniformly reduced with Treg ablation. Johanns et al. (37) also demonstrated that FoxP3+ Treg dictated the natural progression of persistent virulent *Salmonella* infection in mice. In human cases, *Helicobacter pylori*-positive patients revealed a 19- to 25-fold induction of FoxP3 transcript levels in the antrum and cardia, and naturally occurring Treg (CD4+ CD25\textsuperscript{high} FoxP3+) are associated with higher *H. pylori* colonization and increased TGF-β1 gene expression (38). In addition, CD4+ FoxP3+ Treg suppressed host responses to *Haemophilus ducreyi* during experimental skin infection of human volunteers (39). On the basis of these reports, Treg are associated with the pathogenesis of a persistent bacterial infection, but not viral infection, in both animal models and humans. In this study, Treg depletion induced a 99.9% reduction of bacterial counts in MEE and also significantly reduced the ratio of NTHi culture-positive MEE. The levels of proinflammatory cytokines were also reduced in MEEs in children with acute OM is dependent on the number of bacterial species identified, and this supports our findings of the cytokine levels in MEE. On the basis of our observations, Treg may also suppress immune responses and cause persistent infection in the middle ear of a mouse model of COM induced by NTHi inoculation and ET blockade.

In conclusion, we developed a murine model of COM with persistent NTHi infection by using an ET block for 2 months. Treg gathered in the middle ear, and especially CD4+ CD25+ FoxP3+ Treg are assumed to confer infectious tolerance to NTHi in the middle ear. These phenomena may also be at work in the pathogenesis of COM with NTHi infection in other upper and lower airway diseases. A new therapeutic strategy may be developed against chronic inflammatory diseases with NTHi infection by using our mice model.

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