Osteonecrosis of the Jaw Developed in Mice

DISEASE VARIANTS REGULATED BY γδ T CELLS IN ORAL MUCOSAL BARRIER IMMUNITY

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Sil Park,†§‡‡‡, Keichi Kanayama,†§‡, Kausaljit Kaur†, Han-Ching Helen Tseng§, Sina Banankhah‡, Davood Talebi Quje§, James W. Sayre§, Anahid Jewett‡, and Ichiro Nishimura†‡§

From the †Weintraub Center for Reconstructive Biotechnology, Division of Advanced Prosthodontics and ‡Division of Oral Biology and Medicine, UCLA School of Dentistry, Los Angeles, California 90095, the §Department of Periodontology, Asahi University School of Dentistry, Gifu 501-0296, Japan, and the †Department of Biostatistics, UCLA Fielding School of Public Health, Los Angeles, California 90095

Osteonecrosis of the jaw (ONJ), an uncommon co-morbidity in patients treated with bisphosphonates (BP), occurs in the segment of jawbone interfacing oral mucosa. This study aimed to investigate a role of oral mucosal barrier γδ T cells in the pathogenesis of ONJ. Female C57Bl/6J (B6) mice received a bolus zoledronate intravenous injection (ZOL, 540 μg/kg), and their maxillary left first molars were extracted 1 week later. ZOL-treated mice (WT ZOL) delayed oral wound healing with patent open wounds 4 weeks after tooth extraction with characteristic oral epithelial hyperplasia. γδ T cells appeared within the tooth extraction site and hyperplastic epithelium in WT ZOL mice. In ZOL-treated γδ T cell null (Tcrd−/− ZOL) mice, the tooth extraction open wound progressively closed; however, histological ONJ-like lesions were identified in 75 and 60% of WT ZOL and Tcrd−/− ZOL mice, respectively. Although the bone exposure phenotype of ONJ was predominantly observed in WT ZOL mice, Tcrd−/− ZOL mice developed the pustule/fistula disease phenotype. We further addressed the role of γδ T cells from human peripheral blood (h-γδ T cells). When co-cultured with ZOL-pretreated human osteoclasts in vitro, h-γδ T cells exhibited rapid expansion and robust IFN-γ secretion. When h-γδ T cells were injected into ZOL-treated immunodeficient (Rag2−/− ZOL) mice, the oral epithelial hyperplasia developed. However, Rag2−/− ZOL mice did not develop osteonecrosis. The results indicate that γδ T cells are unlikely to influence the core osteonecrosis mechanism; however, they may serve as a critical modifier contributing to the different oral mucosal disease variations of ONJ.

Osteonecrosis of the jaw (ONJ) in the maxilla and mandible has emerged as an uncommon and occasionally severe co-morbidity among patients treated with a group of anti-resorptive agents such as amino-bisphosphonates (BP) (1) and denosumab, a humanized anti-RANKL monoclonal antibody (2). By contrast, ONJ has not been reported in patients treated with hormone replacement therapy or selective estrogen receptor modulators. ONJ is initially defined as an exposed jawbone in the oral cavity for more than 8 weeks even after appropriate intervention therapies in patients with current or past BP treatment (3, 4). An increasing number of case reports suggest that considerable variations exist in the clinical manifestations of ONJ, ranging from radiographic bone pathology without ulcerative oral mucosal lesions to localized swelling with persistent fistula formation (5–7). ONJ is most frequently reported in zoledronate (ZOL)-treated patients with a primary diagnosis of multiple myeloma or bone metastatic tumors such as breast cancer (8, 9). These high risk patients are generally middle-aged to elderly, and they are often treated with immunosuppressive and chemotherapeutic agents, glucocorticoids, or both (10, 11). Therefore, other studies have addressed the combinatory effect of BP with multiple medications on the development of ONJ-like lesions in mice (Table 1). Although additional systemic and poly-pharmaceutical conditions might act as important cofactors that increase the prevalence of ONJ and contribute to its clinical variations (12, 13), the complex conditions have made it difficult to determine its core pathological mechanism.

ONJ primarily occurs in the oral segment of the jawbone that is tightly covered by keratinized oral mucosa. The oral mucosa, i.e. gingiva or palatal tissue, is composed of stratified epithelium and thin connective tissue; furthermore, it is considered to be

**Background:** The pathological mechanism of osteonecrosis of the jaw (ONJ) is unknown.

**Results:** Mouse ONJ-like lesions exhibited epithelial hyperplasia associated with γδ T cells of mouse or human origin.

**Conclusion:** γδ T cells may modify the oral disease phenotypes of ONJ.

**Significance:** ONJ pathogenesis may involve multiple mechanisms separately leading to the development of osteonecrosis or oral epithelial abnormality.

John W. Sayre

1 To whom correspondence should be addressed: The Weintraub Center for Reconstructive Biotechnology, UCLA School of Dentistry, Box 951668, CHS B3-087, Los Angeles, CA 90095. Tel.: 310-794-7612; Fax: 310-825-6345; E-mail: inishimura@dentistry.ucla.edu.

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one of the most protective tissue barriers against physical and chemical damage, bacterial infection, and environmental stress (14). Case control studies of patients with ONJ have indicated an increased risk of developing this condition with tooth extraction or the use of ill-fitting removable dental prostheses (15, 16). These “event-related” oral conditions among BP-treated patients can lead to inflammation in the oral mucosa that likely activates oral barrier immunity. Thus, we hypothesized that the close proximity of the jawbone to the oral mucosa enables the involvement of abnormally stimulated oral barrier immunity during ONJ pathogenesis.

T cells expressing canonical γδ T cell receptors represent a small subset of circulating immune cells and account for 2–5% of peripheral blood T cells in humans. A deficiency in circulating γδ T cells has been reported in patients with long term and repeated BP administrations (17, 18), and BP-induced γδ T cell deficiency was postulated to promote an underlying susceptibility to the development of ONJ (17). Because γδ T cells are preferentially involved in barrier immunity (19, 20), we hypothesized that the γδ T cells in the oral barrier tissue play an important role in the development of ONJ. This study developed a mouse model exhibiting ONJ-like lesions. The role of γδ T cells was addressed in the γδ T cell-deficient Tcrd−/− mice and human γδ T cell-injected Rag2−/− mice. This study suggests that γδ T cells may not be centrally involved in its core osteonecrosis mechanism; however, their presence or absence in oral barrier immunity appears to contribute to the epithelial disease variations of ONJ.

Materials and Methods

Ethics Statement—The UCLA Animal Research Committee reviewed and approved all experimental protocols involving animals (ARC 1997-136). The UCLA Institutional Review Board reviewed and approved all protocols involving human subjects (IRB 12-001176).

ZOL Injection to Mice—This study sought to establish a mouse model exhibiting ONJ-like lesions. Seven-week-old female C57Bl/6j mice (the Jackson Laboratory, Bar Harbor, ME) were subjected to a bolus injection of 540 μg/kg ZOL (Reclast, Novartis, East Hanover, NJ) through the retro-orbital venous plexus (WT ZOL) (21). In the control group, 0.9% NaCl vehicle solution was injected through the retro-orbital venous plexus (WT NaCl). The ZOL dose, which is pharmacologically relevant to oncological doses in humans, was estimated via allometric scaling (22). Each mouse was administered only a single ZOL injection. In the initial study, mouse renal toxicity was evaluated using a creatinine clearance test for serum or urine samples obtained 1 day after ZOL (n = 6) or NaCl (n = 6) injection.

Maxillary First Molar Extraction—One week after the ZOL or NaCl injection, the maxillary left first molar was extracted (23). Mice were anesthetized via isoflurane inhalation and placed on a custom-made surgical table in a supine position using the fixed positioner on the maxillary incisors. A nasal tube was used for the continuous inhalation of 2–4% isoflurane mixed with oxygen during the surgical manipulations in the oral cavity. After the supraborony circumferential periodontal ligament of the attached gingiva was dissected with a dental explorer, the maxillary left first molar was laterally luxated by inserting the tip of a dental explorer between the first and second molars. The luxated molar was then gently removed using surgical forceps. Surgical complications such as tooth fracture occurred and appeared to cause confounding problems. As such, those mice were eliminated from further evaluation. Immediately prior to tooth extraction, 5.0 mg/kg carprofen was administered to alleviate postsurgical pain and to minimize bleeding. Euthanasia by 100% CO2 inhalation was performed on day 4 (WT NaCl, n = 6; WT ZOL, n = 7), week 1 (WT NaCl, n = 8; WT ZOL, n = 9), week 2 (WT NaCl, n = 11; WT ZOL, n = 11), or week 4 (WT NaCl, n = 8; WT ZOL, n = 12) after tooth extraction. The maxilla containing the tooth extraction wound was removed at the time of euthanasia, and each specimen was placed in a plastic bag with 10% buffered formalin for histopathological examination.

Maxillary Tissue, Femur, and Whole Blood Collection—Euthanasia by 100% CO2 inhalation was performed on day 4 (WT NaCl, n = 6; WT ZOL, n = 7), week 1 (WT NaCl, n = 8; WT ZOL, n = 9), week 2 (WT NaCl, n = 11; WT ZOL, n = 11), or week 4 (WT NaCl, n = 8; WT ZOL, n = 12) after tooth extraction. The maxilla, including the tooth extraction wound, was removed at the time of euthanasia and the maxillary tissues were fixed in 10% formalin (pH 7.4) for 24 h. One TOH 10, 2015
and femur were harvested. The maxillary tissue was subjected to standardized digital photo recording. The clinical photograph was enlarged and examined for tooth extraction wound healing.

The harvested maxillary tissue and femurs were fixed in 10% buffered formalin and used for imaging by micro-computed tomography (micro-CT: μCT40, Scanco Medical, Bassersdorf, Switzerland) at an x-ray energy level of 55 peak kV with an intensity of 145 μA. The voxel size was 20 μm with a slice increment of 20 μm. The fixed maxillary tissues were further treated with a formic acid-based decalcifying solution (Immunocal, Unmunotec, Swanton, VT) or 10% EDTA for 7 days for histological section preparation as described below.

Separately, whole blood samples were obtained at the time of euthanasia via cardiac puncture using a 23-gauge needle. Serum chemistry was determined for alkaline phosphatase, calcium, and phosphorus (24).

**Characterization of γδ T Cells in Mouse Oral Mucosal Tissue**—To evaluate γδ T cells in the oral mucosa barrier tissue, a cell dissociation study was performed. Two weeks after molar extraction, the entire gingival/palatal oral mucosa tissue, including the wound area over the tooth extraction socket, was harvested from WT ZOL (n = 3) and WT NaCl (n = 3) mice. The gingival/palatal tissue was cut into small pieces, incubated with the premixed enzymes of a commercially available cell dissociation kit (Tumor Dissociation Kit, Miltenyi Biotec, Auburn, CA), and subjected to repeated mechanical agitations at room temperature and incubation at 37 °C. Dissociated gingival/palatal tissue cells were washed and incubated with FITC-conjugated monoclonal antibody against CD45 and PE-conjugated monoclonal antibodies against CD3, γδTCR (GL3), or DX5 (BioLegend, San Diego). IgG2b was used as the isotype control. After 15 min of incubation on ice, cells were analyzed by flow cytometry (EPICS XL-MCL, Coulter, Miami, FL) (25, 26). The data were presented using the lymphocyte gate.

To further investigate the presence of γδ T cells in the tooth extraction socket histologically, Tcrd-H3BEGFP mice were used. Because of a knock-in mutation of an internal ribosome entry site-controlled histone 2B-enhanced GFP inserted into the 3' end of the T cell receptor δ constant gene, γδ T cells of Tcrd-H2BEGFP mice were highlighted by GFP fluorescence (Tcrd- GFP+) (25). Tcrd-H2BEGFP mice were subjected to ZOL or NaCl injection followed by maxillary first molar extraction as described above. At day 4 (n = 3 in each group) and week 2 (n = 3 in each group) of tooth extraction, the maxillary tissue was harvested. After being fixed in 10% buffered formalin and decalcified, maxillary tissues were embedded in paraffin. Deparaffinized frontal sections, including the tooth extraction site, were subjected to antigen retrieval via microwave irradiation and incubated with an anti-GFP antibody (Santa Cruz Biotechnology, Dallas). GFP+ cells were visualized using the diaminobenzidine substrate, and the sections were counterstained with hematoxylin. Representative tooth extraction sockets were identified, and the number of GFP+ cells within the socket was determined. The data are presented as the number of GFP+ cells per extraction socket area (mm²).

**ONJ Development in γδ T Cell Null (Tcrd−/−) Mice**—γδ T cell null (Tcrd−/−) mice carry T cell receptor δ that was disrupted by gene targeting, resulting in the loss of T cells with γδTCR expression in the lymphoid and epithelial tissues (27). To address the role of γδ T cells in the pathological development of ONJ, female 7-week-old Tcrd−/− mice (B6.129P2-Tcrd<sup>em1Mom</sup>/J, The Jackson Laboratory) were treated with a ZOL injection (Tcrd−/− ZOL) followed by maxillary first molar extraction as described above.

Because WT ZOL mice exhibited delayed tooth extraction healing and developed ONJ-like lesions at week 4, we primarily investigated the wound healing phenotype of Tcrd−/− ZOL mice at week 4 (n = 15). In addition, this study examined the early wound healing process of Tcrd−/− ZOL mice at day 4 (n = 3), week 1 (n = 3), and week 2 (n = 6). The harvested maxillary tissues were photo-recorded, fixed with 10% buffered formalin, and subjected to micro-CT imaging. The specimens were further subjected to decalcification and paraffin embedding for histological specimen preparation. In a parallel experiment, Tcrd−/− mice injected with 0.9% NaCl vehicle solution via the retro-orbital venous plexus were subject to left maxillary first molar extraction (n = 6). Maxillary tissue was harvested on week 2.

**Evaluation of Oral Mucosa Inflammation**—The digital photo-recordings from WT NaCl, WT ZOL, and Tcrd−/− ZOL mice were used to identify the presence or absence of clinical alveolar bone exposure of WT and Tcrd−/− mice. The swelling area of gingival/palatal tissue was measured using a Java-based image-processing program (ImageJ, National Institutes of Health, Bethesda), which was standardized using the circumferential crown area of the remaining first molar.

To assess the inflammation degree of the oral mucosal tissue, cells dissociated from gingival/palatal tissues harvested at day 4 and week 2 (n = 3 in each group) were incubated with a PE-conjugated CD45 monoclonal antibody (BioLegend) and analyzed using flow cytometry. The number of CD45+ lymphocytes among the total cells dissociated from gingival/palatal tissues was determined.

**Histological Examination of ONJ-like Lesions**—The fixed and decalcified maxillary tissues of WT NaCl, WT ZOL, and Tcrd−/− ZOL mice were subjected to a conventional paraffin-embedded histological preparation. A series of frontal sections (8 μm thick) of maxilla were prepared by bisecting the tooth extraction site and the contralateral remaining first molar. Histological sections were stained with hematoxylin and eosin. Histological images were digitized using a high throughput imaging system (Ariol SL-50, Applied Imaging, Grand Rapids, MI) and archived. ONJ-like lesions were determined based on the following criteria following the updated ONJ definition by the American Association of Oral and Maxillofacial Surgeons (28): 1) jawbone exposure associated with abnormal epithelial hyperplasia communicating to the palatal/alveolar bone, or 2) the development of a pustule directly on the surface of the alveolar bone associated with epithelial fistula. The jawbone exposure exhibited food and debris impaction, whereas the pustule only contained inflammatory cells. The prevalence was expressed as the number of animals with ONJ-like oral mucosal abnormalities over the number of animals at week 4.

For alveolar bone characterization, the histological images of palatal/alveolar bone containing the first molar extraction site...
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were horizontally bisected to divide the oral and nasal sides. The primary focus was placed on the oral side of palatal/alveolar bone, which encompassed the buccal border of the alveolar bone to the mid-palatine suture. The degree of osteonecrosis was defined as the ratio between the number of osteoclasts and the number of nonvital osteocytes determined using osteocytic lacunae without a cellular component (i.e., empty lacunae) or abnormally condensed small nuclei (pyknotic osteocytes). An operator blind to condition performed the histological evaluation.

The number of osteoclasts (OCs) within the oral half of the palatal/alveolar bone containing the first molar extraction site was counted in the histological specimens under ×20 magnification. OCs were defined as large cells with multiple nuclei (≥2 nuclei) on the bone surface. Specific histological sections were stained with tartrate-resistant acid phosphatase using a commercially available kit (Sigma). The area of palatal/alveolar bone was measured using an image-processing program (ImageJ, National Institutes of Health). The OC number was normalized to the palatal/alveolar bone area.

Micro-CT Examination—Micro-CT data were reconstructed to generate three-dimensional images at a threshold of 220 using the software provided by the manufacturer of the micro-CT scanner.

Maxillary micro-CT three-dimensional images of WT NaCl, WT ZOL, and Tcrd−/− ZOL mice were used to identify alveolar bone abnormalities such as periosteal reaction, bone sequestra associated with ONJ, and bone remodeling. A blinded operator reviewed the micro-CT three-dimensional images, and the degrees of bone formation and bone resorption at the tooth extraction site were separately rated from 0 to 2. The bone remodeling index was calculated by combining the bone formation and resorption indices. Various micro-CT images exhibited severe periosteal reactions at the external surface of the alveolar bone, which were not considered in the quantitative evaluation.

Bone morphometry of femurs harvested 2 weeks after tooth extraction or 3 weeks after ZOL injection was characterized in WT NaCl, WT ZOL, and Tcrd−/− ZOL mice (n = 6 in each group) by micro-CT. Volume of interest was set 600 µm below the end point of the growth plate, which was used as an anatomical landmark. One hundred slices (1.2 mm) were evaluated in the distal femur metaphysis at a threshold of 220. The volume of interest only included secondary spongiosa. Trabecular bone morphometric measurements in three-dimensions included the following: bone volume normalized to tissue volume, trabecular number, trabecular thickness, and trabecular separation.

Human γδ T Cells and CD3+ T Cells Co-cultured with ZOL-pretreated Human Osteoclasts—For preparing human osteoclasts (h-OCs), peripheral blood was collected from healthy donors, and peripheral blood mononuclear cells were separated by Ficoll-Hypaque gradient centrifugation. Peripheral blood mononuclear cells were cultured onto the tissue culture plate for 1 h, after which the adherent subpopulation of peripheral blood mononuclear cells was detached from the tissue culture plates, and the CD14+ monocytes were purified (EasyStep Human Monocyte Isolation Kit, Stem Cell Technologies, Vancouver, Canada). Approximately 94% purity was achieved based on flow cytometric analysis of CD14. Monocytes (10.3 × 10^6 cells) were cultured in α-minimal essential medium containing 25 ng/ml macrophage colony-stimulating factor (M-CSF) and receptor activator of nuclear factor-κB ligand (RANKL) (25 ng/ml) (29). Medium was refreshed every 3 days. Development of h-OCs was monitored by tartrate-resistant acid phosphatase staining (Primary Cell Co., Sapporo, Japan) and resorption pit formation on synthetic carbonate apatite pre-coated plate (Cosmo Bio USA, Carlsbad, CA) following the manufacturer’s protocols. Mature h-OCs were generated after 14–21 days of incubation. Once developed, h-OCs were treated with 100 nM ZOL for 6 days.

For preparing human γδ T cells and CD3+ T cells (γδ T cells and h-CD3+ T cells, respectively), peripheral blood samples of healthy donors were used to purify the corresponding cells by commercially available selection kits (EasyStep Human Gamma/Delta T Cell Isolation kit and EasyStep Human CD3 Positive Selection kit, StemCell Technologies). As the result, h-γδ T cells (3.0 × 10^9) and h-CD3+ T cells (51 × 10^9) cells, respectively, were obtained. h-γδ T and h-CD3+ T cells were activated in RPMI 1640 media containing anti-CD3 antibody (1 µg/ml), anti-CD28 antibody (7 µg/ml), and rh-IL-2 (100 units/ml) overnight. Then, h-γδ T cells (3.0 × 10^9 cells per plate) and h-CD3+ T cells (16.3 × 10^9 cells per plate) were co-cultured with ZOL-pretreated h-OCs (0.6 × 10^9 and 2.0 × 10^9 cells per plate, respectively).

The number of h-γδ T and h-CD3+ T cells in each plate was counted at co-culture days 1, 3, and 6. The co-culture medium of h-OC/h-γδ T cells and h-OC/h-CD3+ T cells was harvested at day 6, and IFN-γ concentration was determined by ELISA. h-γδ T cells were then collected, washed with PBS, and prepared for injection to the Rag2−/− mouse as described below. On the day of mouse injection, surface markers of h-γδ T cells were analyzed by flow cytometry using PE-conjugated monoclonal anti-human CD3, CD4, CD19, and CD69 antibodies (BioLegend) (25, 26).

ONJ Development in B/T Cell-deficient (Rag2−/−) Mice and the Effect of h-γδ T Cell Repopulation—Rag2−/− mice (B6.Cg-Rag2tm1.1Cgn/J, The Jackson Laboratory) carry null mutation of B and T lymphocytes (30). Female WT mice and Rag2−/− mice received the single injection of 500 µg/kg ZOL (WT ZOL and Rag2−/− ZOL, respectively) or 0.9% NaCl vehicle solution (WT NaCl and Rag2−/− NaCl, respectively) via tail vein. In this experiment, mice were housed with autoclaved cellulose-based bedding (Cell-Sorb Plus, Fangman Specialties Inc., Cincinnati, OH).

Maxillary tissue containing the tooth extraction site was photographed and harvested. After being fixed and decalcified, the conventional histological preparation as described above was stained with H&E.

Six days after the ZOL injection, a group of Rag2−/− ZOL mice (n = 4) was injected via tail vein with h-γδ T cells (6.75 × 10^9 per mouse) that had been activated by anti-CD3 and anti-CD28 antibodies and rh-IL-2 followed by co-culturing with ZOL-pretreated h-OCs as described above (Rag2−/− ZOL h-γδ T cell). One day after the h-γδ T cell injection (Rag2−/− ZOL h-γδ T cell, n = 4) or 1 week after ZOL injection (WT ZOL, n =
5; Rag2−/− ZOL, n = 4; WT NaCl, n = 4; Rag2−/− NaCl, n = 4), the maxillary first molar was extracted.

All mice were euthanized 2 weeks after tooth extraction. Spleens and bone marrow cells were harvested from Rag2−/− ZOL h-γδ T cell mice. Splenocytes and bone marrow cells were incubated with FITC-conjugated monoclonal antibody against mouse CD45 and PE-conjugated monoclonal antibodies against mouse γδ TCR (GL3), CD19, DX5, or F4/80 (BioLegend). Furthermore, splenocytes and bone marrow cells were incubated with FITC-conjugated monoclonal antibodies against human CD45 or CD3 and PE-conjugated monoclonal antibodies against human γδ TCR, CD19, or NkP46 (BioLegend) and analyzed by flow cytometry. IgG2b served as the isotype control.

Using histological sections stained with H&E, osteonecrosis was assessed by the cluster area of ≥5 necrotic osteocytes over the alveolar bone area. Epithelial abnormality was assessed for the presence or absence of hyperplastic epithelium.

**Standardized Osteoclast Number of WT and Rag2−/− Mice**—The histological sections through the distal root area of the first molar from the above experiment were stained by tartrate-resistant acid phosphatase, and the number of OCs on the surface of palatal bone and in the tooth extraction socket was separately counted. The number of OC was standardized by the bone surface linear length.

**Effect of m-OC on h-γδ T Cell and Mouse (m)-γδ T Cells in Vitro**—m-OCs were generated from mouse femur flow-through cells with supplementation with 25 ng/ml M-CSF and 25 ng/ml RANKL for 7 days. h-γδ T cells were isolated from peripheral blood of healthy subjects as described above. m-γδ T cells were isolated from mouse spleen (TCRγδ T cell isolation kit, mouse, Miltenyi Biotec). After the activation with rh-IL2 (100 units/ml) and lipopolysaccharide (LPS) (100 ng/ml), h-γδ T cells and m-γδ T cells were co-cultured with m-OCs that were exposed to 100 nM ZOL for overnight. h-γδ T cells and m-γδ T cells were also co-cultured with m-OCs without ZOL pretreatment. At 1, 4, and 8 days of co-culture, the medium was collected and assayed for IFN-γ by ELISA.

**Statistical Analyses**—Student’s t tests were used to compare means. A one-way analysis of variance followed by a Bonferroni correction was used when more than two groups were compared. Fisher’s exact test was used to describe the prevalence of ONJ-like lesions in contingency tables. The correlation between osteonecrosis and OC numbers was assessed by Pearson correlation and Spearman’s rank correlation tests.

**Results**

**ONJ-like Lesions in WT Mice**—In control WT mice injected with 0.9% NaCl vehicle solution (WT NaCl), the maxillary first molar crown, which remained open after surgery. Four days after tooth extraction, all WT NaCl mice continued to exhibit an open wound in the gingival tissue, which was surrounded by localized swelling (Fig. 1A). The open wound of the maxillary gingival tissue progressively closed and was completely covered by epithelial tissue between weeks 2 and 4. In the ZOL-treated WT mice (WT ZOL), the closure of gingival open wounds was slow. In total, 50% (6/12) of the animals exhibited clinically open wounds at week 4, and the soft tissue swelling continued to surround the wound opening (Fig. 1A). The open wound was more common at the large mesial root, which was observed in all WT ZOL mice.

Histological sections revealed that the tooth extraction wounds of WT NaCl mice healed with new bone formation in the bony socket, and the oral epithelial integrity was re-established between weeks 2 and 4 (Fig. 1B). Mild inflammatory cell infiltrates were observed in the palatal oral mucosa. By contrast, WT ZOL mice exhibited alveolar bone exposure at the mesial root extraction site with food and debris impaction. At the small distal root areas, where the oral mucosa wound was closed based on gross observation, histological evaluations revealed definitive bone exposure facilitated by the abnormal migration of oral epithelial cells. The abnormal proliferation and migration of epithelial cells resembling pseudoeipithelio-matous hyperplasia (24, 31) resulted in direct epithelial contact with the partially necrotic alveolar bone. Reduced new bone formation was noted in the extraction socket, and densely localized inflammatory cell infiltration was observed on the surface of the palatal bone adjacent to the tooth extraction site (Fig. 1B).

WT NaCl and WT ZOL mice showed serum levels of alkaline phosphatase (125.7 ± 29.95 and 97.0 ± 28.36 units/liter, respectively), calcium (8.9 ± 0.61 and 9.1 ± 0.88 mg/dl, respectively), and phosphorus (8.4 ± 1.44 and 7.0 ± 2.59 mg/dl, respectively). These serum chemistry data did not indicate any differences between groups and were within the normal range, i.e. alkaline phosphatase, 13–291 units/liter; calcium, 6.8–11.9 mg/dl; and phosphorus, 5.3–11.3 mg/dl. Furthermore, WT NaCl and WT ZOL mice exhibited similar renal function as measured by creatinine clearance (0.35 ± 0.07 and 0.33 ± 0.01 mg/dl, respectively), which was within the normal range of 0.1–2.1 mg/dl. No signs of toxicity related to ZOL injection were observed.

**Oral γδ T Cells in the Tooth Extraction Wound**—The cellular component was dissociated from gingival/palatal oral mucosa tissues harvested from WT NaCl and WT ZOL mice 2 weeks after the tooth extraction. Cells in the lymphocyte gate in the flow cytometry were predominantly composed of CD45+ cells (Fig. 2A). Within the CD45+ population, the WT NaCl and WT ZOL samples contained 7.1 and 5.5% CD3+ T cells, respectively. GL3+ γδ T cells were 0.4 and 1.3%, and DX5+ natural killer cells were 1.1 and 2.3% in the WT NaCl and WT ZOL samples, respectively. It was estimated that the fraction of γδ T cells in the infiltrated T cell population was 5.6 and 23.6% in the WT NaCl and WT ZOL oral wound samples at week 2 of tooth extraction.

Using Tcrd-H2BEGFP mice, GFP+ γδ T cells were examined in the extraction socket (Fig. 2B). At day 4 after tooth extraction, the number of GFP+ γδ T cells did not differ between Tcrd-H2BEGFP NaCl and ZOL mice (Fig. 2C). However, at week 2 of wound healing, the number of GFP+ γδ T cells significantly decreased in the control mice. By contrast, Tcrd-H2BEGFP ZOL mice exhibited an increase of GFP+ γδ T cells in the tooth extraction socket (Fig. 2C). A few GFP+ γδ T cells were identified in the unwounded junctional epithelium of the free gingiva adjacent to the tooth (Fig. 2D). However, in the oral epithelial hyperplasia in Tcrd-H2BEGFP ZOL mice, an
increased density of GFP\(^+\) γδ T cells was observed (Fig. 2D). Combining the data of GL3\(^+\) and GFP\(^+\) γδ T cells, ZOL treatment appeared to associate with the sustained appearance of γδ T cells in the oral mucosa at the tooth extraction wound site.

**Different Oral Mucosa Wound Healing in Tcrd\(^{-/-}\) ZOL Mice**—We addressed the possible role of γδ T cells in the development of ONJ using γδ T cell-deficient (Tcrd\(^{-/-}\)) mice. The healing patterns of the tooth extraction wound were indistinguishable between the WT ZOL and Tcrd\(^{-/-}\) ZOL mice until week 1. At week 2, the tooth extraction wound area was visibly reduced in Tcrd\(^{-/-}\) ZOL mice compared with WT ZOL mice. At week 4, the tooth extraction wound was closed in 86.7% of Tcrd\(^{-/-}\) ZOL mice (13/15); however, gingival/palatal tissue swelling remained (Fig. 3A). The standardized oral mucosal...
swelling area was significantly larger in the Tcrd−/− ZOL mice compared with the WT ZOL mice at weeks 2 and 4 (Fig. 3B).

Pustule/Fistula Development on the Necrotic Bone Surface of Tcrd−/− ZOL Mice—Histological cross-sections of Tcrd−/− ZOL mice revealed well healed extraction sockets and oral mucosa wounds at week 4 (Fig. 3C). In 26.7% of the Tcrd−/− ZOL mice, small sequestrations of necrotic bone were observed in the connective tissue of the oral mucosa. Unlike the ONJ-like lesions of the WT ZOL mice, the development of large pustules on the surface of necrotic palatal bones was observed in Tcrd−/− ZOL mice (Fig. 3C). The associated epithelium at the pustule was consistent with fistula formation.

Prevalence of ONJ-like Lesions in WT NaCl, WT ZOL, and Tcrd−/− ZOL Mice—Histological jawbone exposure was observed in both the WT ZOL (66.7%) and Tcrd−/− ZOL (20.0%) groups (Fig. 3D). By contrast, pustule/fistula formation was predominantly observed in Tcrd−/− ZOL mice (40.0%) compared with WT ZOL mice (8.3%), although this was not significant (p = 0.06). When the bone exposure phenotype and pustule/fistula phenotype were combined as ONJ, the WT ZOL (75.0%) and Tcrd−/− ZOL (60.0%) mice suggested an equal likelihood of developing ONJ-like lesions (Fig. 3D). The prevalence of oral epithelial abnormalities was 0% in WT NaCl control mice at week 4.
The effect of ZOL on the femurs was characterized in WT ZOL and Tcrd−/− ZOL mice by micro-CT, which demonstrated the similar increases in bone volume normalized to tissue volume and trabecular number and decreases in trabecular separation as compared with the WT NaCl control group (Fig. 3E). No significant difference was noted between WT ZOL and Tcrd−/− ZOL femurs.

Severe Periosteal Reactions Occurred in WT ZOL Mice but Not in Tcrd−/− ZOL Mice—Radiographic and CT image examinations of patients with ONJ have revealed that periosteal reac-
tion predominantly occurs during advanced disease stages (32, 33). This study used micro-CT images to determine the presence of abnormal alveolar bone in mouse maxilla. WT NaCl mice revealed evidence of bone resorption in the palatal alveolar bone at week 2 of tooth extraction as well as bone formation in the extraction socket at week 4 (Fig. 4, A and B). WT NaCl mice did not exhibit abnormal alveolar bone wound healing. ZOL treatment of WT and Tcrd−/− mice appeared to decrease osteolytic signs at the extraction site, whereas bone formation in the extraction socket was less affected (Fig. 4, osteolytic signs at the extraction site, whereas bone formation exhibited periosteal reactions, including severe forms, at the total of 53% of WT ZOL mouse specimens from weeks 2 and 4 exhibited periosteo reactions, including severe forms, at the tooth extraction wound healing site or the contralateral periodontal disease area (Fig. 4A). Tcrd−/− ZOL mice demonstrated a small "sun-ray" type of calcified spikes on the surface of the alveolar bone; however, the severity of the periosteal reaction was attenuated considerably (Fig. 4A).

Chronic Oral Mucosa Inflammation in WT ZOL and Tcrd−/− ZOL Mice—Oral epithelial hyperplasia in WT ZOL mice migrated toward the inflammatory lesion and established direct contact between the basal cells and the surface of partially necrotic alveolar bone (Fig. 5A). These characteristic features were less apparent in Tcrd−/− ZOL mice, and the epithelial tissue did not reach the bone surface except at the periphery of the pustule (Fig. 5A). Despite the different oral mucosal disease phenotypes of ONJ-like lesions in WT ZOL and Tcrd−/− ZOL mice, both models exhibited sustained inflammation. WT ZOL mice exhibited localized and dense inflammatory cell infiltrates near the bone surface, whereas inflammation in the gingival/palatal mucosa tissues was more diffuse and often exhibited cellulitis-like inflammatory cell infiltration in Tcrd−/− ZOL mice (Fig. 5A).

Dissociated cells from the gingival/palatal mucosa tissues contained over 50% CD45+ leukocytes in both the WT NaCl and WT ZOL mice at day 4 (Fig. 5B). Control WT NaCl mice exhibited a significant reduction in CD45+ cells at week 2, whereas WT ZOL mice maintained high levels of CD45+ cells at week 2 (Fig. 5B). Similarly, Tcrd−/− ZOL mice exhibited an increase in CD45+ cells at week 2 (Fig. 5B). Two weeks after the maxillary first molar extraction, Tcrd−/− NaCl mice also sustained oral mucosa swelling and CD45+ cell infiltration (data not shown).

Osteonecrosis in WT ZOL and Tcrd−/− ZOL Mice—The degree of osteonecrosis in the palatal/alveolar bone was defined as the proportion of nonvital osteocyte lacunae in the oral and nasal sides. The nasal side of palatal bone contained 10–15% nonvital osteocyte lacunae in all groups at all of the tested time points (data not shown). The oral side of the palatal bone exhibited a cluster of ~20% nonvital osteocyte lacunae immediately after tooth extraction, which gradually decreased over time in control WT NaCl mice (Fig. 5C). In WT ZOL mice, the area of osteonecrosis was observed in the bone exposure site as well as the palatal bone that interfaced the oral mucosa and dense inflammatory infiltrates (Fig. 5A). The area of osteonecrosis was not reduced over time but rather increased at week 4, which resulted in persistent osteonecrosis (Fig. 5C). The osteonecrosis areas of the Tcrd−/− ZOL mice were associated with bone sequestra and pustules (Fig. 5A). The Tcrd−/− ZOL group exhibited increased osteonecrosis development at week 2, followed by a mild decrease at week 4 (Fig. 5C). The reference Tcrd−/− NaCl group developed a significantly smaller necrotic bone area at week 2, despite the remaining open wound (data not shown).

The number of OCs in the palatal/alveolar bone of WT NaCl control mice demonstrated an early increase at day 4 and week 1 after tooth extraction, followed by a gradual decrease (Fig. 5D). By contrast, OCs in the palatal/alveolar bone of WT ZOL and Tcrd−/− ZOL mice remained relatively unchanged throughout the experiment, resulting in a significantly increased number compared with the WT NaCl mice at week 4 (Fig. 5D). Active bone resorption was observed at the pustule periphery (Fig. 5A). The removal of necrotic bone sequestra might contribute to the late decrease of osteonecrosis area in Tcrd−/− ZOL mice.

The correlation between the area of osteonecrosis and the number of osteoclasts in the combined data of weeks 1, 2, and 4 was evaluated by Pearson correlation and Spearman’s rank correlation tests. The WT NaCl group revealed the significant correlation in both tests, whereas the WT ZOL and Tcrd−/− ZOL groups showed a similar correlation pattern with no statistical significance (Fig. 5E).

In summary, ZOL-treated Tcrd−/− mice developed the different oral mucosa disease phenotype characterized by the pustule and fistula formation; however, the osteonecrosis occurred similarly to WT ZOL mice.

Activation of Human Peripheral Blood γδ T Cells by ZOL-pretreated h-OC—h-γδ T cells contain Vγ9Vδ2 T cells. BP-pretreated macrophages accumulate phosphorylated metabolites such as isopentenyl pyrophosphate (IPP) (34), which have been shown to selectively activate h-γδ T cells but not mouse γδ T cells. In the context of ONJ pathogenesis, we investigated the role of ZOL-pretreated h-OCs, rather than peripheral blood macrophages, on the activation of h-γδ T cells in this project.

Three million h-γδ T cells and 51 million h-CD3+ T cells were isolated from approximately equal volumes of peripheral blood of different healthy donors. The estimated purity of h-γδ
T cells and h-CD3

The number of h-CD3+ T cells decreased (Fig. 6B).

After 6 days of co-culture, the secretion of IFN-γ was significantly increased during the co-culture period, whereas the number of h-CD3+ T cells decreased (Fig. 6B).

FIGURE 4. Alveolar bone healing assessment by micro-CT imaging. A, WT NaCl control mice exhibited signs of osteolysis at week 2 as roughened alveolar and palatal bone surfaces (white arrowheads). Bone formation within the extraction socket (black arrowhead) appeared to complete the wound healing process at week 4. We noted that one WT NaCl mouse exhibited periodontal disease at the nonextraction side (white arrowhead), in WT ZOL and Tcrd−/− ZOL mice, the evidence of bone resorption was not clearly observed, and the extraction socket was not completely filled by new bone formation (black arrowheads). Severe periosteal reaction (black arrows) was observed at the molar extraction site in 53% of the WT ZOL mice. A periosteal reaction was also noted in one WT ZOL mouse at the contralateral nonextraction side with periodontal disease (confirmed with histology). Although the severity of periosteal reaction in Tcrd−/− ZOL mice was much attenuated (arrows), the sun-ray type of calcified spikes was observed both in WT ZOL and Tcrd−/− ZOL mice (SR with black arrow). B, micro-CT images were used to rate bone formation (0–2) and bone resorption (0–2) appearance. The bone-remodeling rate was expressed by combining the rates of bone formation and resorption in each animal. Active bone remodeling was suggested in WT NaCl control mice. Both WT ZOL and Tcrd−/− ZOL mice showed significantly decreased bone resorption. *, p < 0.05.

T cells and h-CD3+ T cells was 76.9 and 96.4%, respectively (Fig. 6A). After overnight activation by anti-CD3 antibody, anti-CD28 antibody, and rh-IL-2, T cells were co-cultured with ZOL-pretreated h-OC. The number of h-γδ T cells robustly increased during the co-culture period, whereas the number of h-CD3+ T cells decreased (Fig. 6B).

The large representation of CD69-positive staining indicated the lymphocyte activation (Fig. 6D). As such, this study suggested that h-γδ T cells were activated by ZOL-pretreated h-OCs.

Rag2−/− Mice Engrafted with h-γδ T Cells—One day after injection of h-γδ T cells to ZOL-treated Rag2−/− mice, the maxillary first molar was extracted. At the time of euthanasia 2 weeks after tooth extraction, spleen and bone marrow cells were harvested. The profile of endogenous mouse immune effectors showed the lack of CD3+ T cells and CD19+ B cells
FIGURE 5. Characterization of osteonecrosis in WT NaCl, WT ZOL, and Tcrd−/− ZOL mice. A, localized inflammation (Inf) was evident near the surface of alveolar bone in WT NaCl mice. In WT ZOL mice, the intense inflammatory reaction was demonstrated in the oral mucosa (double-headed arrow; Inf). Abnormal epithelial tissue adhesion (Epi) was evident on the surface of partially necrotic alveolar bone (Ost. Nec.). In Tcrd−/− ZOL mice, inflammation was more diffused, and epithelial tissue did not reach the necrotic bone surface except at the pustule periphery. In both lesions, clusters of osteoclasts (arrowheads) were observed at and adjacent to the osteonecrosis site. B, CD45+ lymphocytes accounted for 30–45% of the dissociated cells from gingival/palatal tissue at day 4. The fraction of CD45+ cells decreased in WT NaCl mice at week 2. However, WT ZOL and Tcrd−/− ZOL mice exhibited similar fractions of CD45+ lymphocytes in the oral mucosa. C, area of osteonecrosis is presented as the percent of nonvital osteocytes over the total number of osteocytes on the oral side of palatal/alveolar bone. At week 4, the area of osteonecrosis was significantly larger in WT ZOL and Tcrd−/− ZOL mice than WT NaCl control mice. D, number of osteoclasts in the palatal/alveolar bone area suggested that the early increase in the tooth extraction wound of WT NaCl control mice followed by progressive decrease. Both WT ZOL and Tcrd−/− ZOL mice indicated a consistent appearance of osteoclasts throughout the experimental period, resulting in the elevated osteoclast number at week 4. *, p < 0.05; †, p = 0.06 compared with the WT NaCl control group. E, correlation between the osteonecrosis area and the number of OCs in week 1, week 2, and week 4 specimens. The WT NaCl group demonstrated the significant correlation between the osteonecrosis area and osteoclast number. By contrast, correlation pattern was similar in the WT ZOL and Tcrd−/− ZOL groups, which did not show statistical significance.
but the presence of DX5+ natural killer cells in spleen (data not shown) and F4/80+ monocytes in bone marrow (Fig. 6E). The profile of human immune effectors indicated the presence of CD45+γδTCR+ and CD3+γδTCR+ cells (Fig. 6E), suggesting the successful engrafting of h-γδ T cells in Rag2−/− mice, which survived for the experimental period of 2 weeks.

Lack of ONJ-like Legions in ZOL-treated Rag2−/− Mice—Rag2−/− mice with NaCl or ZOL injection showed accelerated tooth extraction wound healing as compared with WT mice. All Rag2−/− NaCl mice showed the closure of the tooth extraction wound at week 2 (Fig. 7A). Although Rag2−/− ZOL mice showed small food and debris impaction at the wound site in three out of five mice in the same healing period, histological examination revealed well healed extraction sockets (Fig. 7A).

Oral Epithelium Hyperplasia in Rag2−/− ZOL Mice Engrafted with h-γδTCR—Rag2−/− ZOL mice engrafted with h-γδ T cells showed an unusual roughened oral soft tissue healing at week 2 (Fig. 7B). Histological examination revealed vari-
ous degrees of oral epithelial hyperplasia in all mice. Oral epithelium exhibited irregular rete pegs and extensions of the basal cells. Where the hyperplastic oral epithelium embedded in the deep connective tissue, small food and debris were enclosed, and the tooth extraction socket showed the delay in bone healing (Fig. 7B).

In this project, we employed the autoclaved cellulose bedding for immunodeficient $\text{Rag2}^{-/-}$ mice, which might, in turn, change the wound-healing environment. WT NaCl mice housed in the same bedding environment showed the accelerated tooth extraction wound closure, whereas WT ZOL mice housed in autoclaved cellulose bedding demonstrated the ONJ.
phenotype and oral wound was open at week 2 in all mice. WT ZOL mice showed the significantly larger osteonecrosis area than WT NaCl mice (Fig. 7C). The osteonecrosis area of Rag2<sup>-/-</sup> NaCl and Rag2<sup>-/-</sup> ZOL mice remained small at the level of WT NaCl mice. Despite the development of oral epithelial hyperplasia in h-γδ T cell-engrafted Rag2<sup>-/-</sup> ZOL mice, the osteonecrosis area remained unaffected (Fig. 7C).

**Reduced Osteoclastogenesis in Rag2<sup>-/-</sup> Mice—Rag2<sup>-/-</sup>** ZOL mice with or without h-γδ T cell engraftment demonstrated significantly less osteoclasts on the palatal bone surface than WT NaCl and WT ZOL mice (Fig. 8A). Dense inflammatory cell infiltration was observed in the palatal/gingival tissue of WT ZOL mice, which interfacial palatal bone lined by a number of osteoclasts. By contrast, no inflammatory infiltration was observed in Rag2<sup>-/-</sup> ZOL mice (Fig. 8A). Because the reduced osteoclast number was similarly observed in Rag2<sup>-/-</sup> NaCl mice (Fig. 8A), it was postulated that osteoclastogenesis on the palatal bone surface required the mediation by T cells, whereas the normalized osteoclast number in the extraction socket of Rag2<sup>-/-</sup> mice could be associated with the accelerated bone wound healing and remodeling.

**m-OC Induced the Secretion of IFN-γ of h-γδ T Cells**—The potential interaction between m-OC and h-γδ T cells was addressed by in vitro co-culture study. When h-γδ T cells were co-cultured with ZOL pretreated m-OC, IFN-γ secretion was significantly increased (Fig. 8B). Because m-OC alone did not secrete IFN-γ, activated h-γδ T cells were the source of IFN-γ. Surprisingly, IFN-γ secretion by h-γδ T cells was also increased when co-cultured with m-OC without ZOL pretreatment, albeit with a transient fashion (Fig. 8B). In the case of m-γδ T cells, nonspecific activation by m-OC with or without ZOL pretreatment was observed resulting in the higher levels of IFN-γ secretion (Fig. 8B).

**Discussion**

This study demonstrated the ONJ-like lesions with severe bone exposure (Fig. 1) and the nonexposure variants (Fig. 3) generated in mice that differed in the presence or absence of γδ T cells, respectively. WT ZOL mice exhibited extended exposure of partially necrotic alveolar bone associated with abnormal oral epithelial hyperplasia. By contrast, in the absence of γδ T cells, Tcrd<sup>-/-</sup> ZOL mice developed pustules on the surface of necrotic alveolar bone, which appeared to form fistulae with the oral epithelium. The exposure of necrotic bone has been a clinical hallmark of a fully developed ONJ lesion. However, the current position of the American Association of Oral and Maxillofacial Surgeons now includes intraoral or extraoral fistulae in the maxillofacial region as an additional clinical definition of medication-related ONJ (28). This revised ONJ definition underscores large clinical variations in the development of oral mucosa abnormalities in this disorder.

The area of osteonecrosis in WT ZOL mice progressively increased, whereas the osteonecrosis area in Tcrd<sup>-/-</sup> ZOL mice peaked at week 2 and then declined (Fig. 5C). The number of osteoclasts on the surface of palatal bone was equivalent in the WT ZOL and Tcrd<sup>-/-</sup> ZOL mice (Fig. 5D). However, histologically, osteoclasts appeared to cluster immediately outside the pustule in Tcrd<sup>-/-</sup> ZOL, resulting in bone sequestration (Fig. 3C). The bone sequestra in Tcrd<sup>-/-</sup> ZOL mice appeared to move toward the oral epithelium and was moved out to the oral cavity, which potentially contributed to the late reduction of the osteonecrotic area. In the presence of γδ T cells, osteoclasts in WT ZOL mice did not cluster but distributed on the palatal bone surface interfacing oral mucosa with severe inflammatory reaction (Fig. 8A). It was further noted that Rag2<sup>-/-</sup> mice lacking T and B cells showed the decreased osteoclast number on the palatal bone surface without inflammatory cell infiltration in the oral mucosa (Fig. 8A). Therefore, we postulate that tooth extraction-induced oral mucosa inflammation containing γδ T cells may be responsible, in part, for the induction of osteoclasts distributed on the palatal bone surface, although the severe inflammatory reaction in the form of pustule formation may be required for the localized osteoclast clustering in Tcrd<sup>-/-</sup> ZOL mice.

A recent retrospective analysis of BP-treated patients in Copenhagen concluded that the severe bone exposure and the nonexposure variants of ONJ belonged to the same disease condition. However, although the severe bone exposure variant was observed from the early BP treatment stage, the nonexposure variant was not reported until patients received repeated BP injections (35). ONJ patients who had received multiple injections of BP have been shown to develop deficiency of circulating γδ T cells (17). Human γδ T cells are largely composed of Vγ9Vδ2 T cells, which can react to phosphoantigens (pAg) such as bacteria or parasite-derived (E)-4-hydroxy-3-methylbut-2-enyl-2-phosphophosphate from the non-mevalonate pathway (36). Vγ9Vδ2 T cells can also react to mevalonate pathway-related pAg's such as IPP and isomer dimethylallyl pyrophosphate, albeit at much lower potencies (37, 38). Macrophages...
exposed to BP such as ZOL become an activator of Vγ9Vδ2 T cells, likely in consequence of intracellular IPP accumulation. The repeated BP injections are postulated to result in the repeated activation of Vγ9Vδ2 T cells leading to eventual γδ T cell depletion (17, 18). It is tempting to speculate that the decreased circulating Vγ9Vδ2 T cells may causally associate with the development of nonexposure and fistula phenotype of ONJ in humans.

It has been established that the Vγ9Vδ2 T cell receptor as well as butyrophilin-3A1 critical for IPP-induced activation of γδ T cells are conserved only in humans, primates, and a few other placental animals but not in mice (39, 40). Therefore, mouse γδ T cells do not respond to pAgS, and the ZOL injection should not activated the circulating γδ T cells in mice through a similar mechanism as in humans. Thus, this study examined the role of human γδ T cells in the development of oral mucosal phenotypes in ZOL-treated Rag2−/− mice. Like BP-exposed macrophages, this study demonstrated that ZOL-treated h-OCS activated h-γδ T cells. By co-culturing with ZOL-pre-treated h-OC, h-γδ T cells were found to increase the cell

FIGURE 8. Osteoclast behavior in Rag2−/− mice with h-γδ T cell engraftment. A, tartrate-resistant acid phosphatase staining revealed osteoclasts on the surface of palatal bone (arrows) interfacing the area of oral mucosa inflammation (Inf) in WT ZOL mice. Strikingly, osteoclasts on the palatal surface were nearly absent in Rag2−/− ZOL mice, whereas osteoclasts appeared in the bone remodeling area of tooth extraction socket (Soc). The number of osteoclasts normalized by the bone surface was significantly reduced on the palatal bone surface in Rag2−/− regardless of ZOL treatment as well as h-γδ T cell engraftment. The osteoclast number in the extraction socket was normalized in Rag2−/− NaCl and Rag2−/− ZOL mice, whereas h-γδ T cell-engrafted mice remained high. *p < 0.05 compared with the WT NaCl control group. B, stimulatory effect of m-OC on h-γδ T cells and m-γδ T cells was assessed in vitro. ELISA revealed that m-OC with ZOL pretreatment specifically activated the IFN-γ secretion by h-γδ T cells but not by m-γδ T cell. However, nonspecific activation by m-OC was observed in both h-γδ T cells and m-γδ T cells, which secreted higher levels of IFN-γ. *p < 0.05 compared with the h-γδ T cell or m-γδ T cell alone group at each time point.
counts, secrete IFN-γ at a much higher level than the similarly activated h-CD3+ T cells, and express the lymphocyte activation marker CD69 (Fig. 6). When h-γδ T cells that were pre-activated by ZOL-treated h-OCs were engrafted onto Rag2−/− ZOL mice, the tooth extraction wound healed with abnormal oral epithelial hyperplasia (Fig. 7).

The abnormal oral epithelial hyperplasia referred to as pseudoepitheliomatous hyperplasia (PEH) was reported in 60% of human ONJ biopsy specimens (24, 31) and was associated with the more aggressive ONJ phenotype with necrotic bone exposure. Akilov et al. (41) reported the development of PEH in mouse ear skin by the injection of TNF-α and IFN-γ, suggesting that overproduction of Th1 cytokines in the barrier immunity stimulated the unregulated proliferation and migration of epidermal basal cells. Rag2−/− mice used in this study lacked T and B cells, whereas NK cells and monocytes were present (Fig. 6). Therefore, the engrafted h-γδ T cells were likely to be a source of Th1 cytokines such as TNF-α and IFN-γ (42).

This study demonstrated that PEH-like oral epithelial migration was less apparent in Tcrd−/− ZOL mice that nonetheless developed the nonexposure variant-like ONJ lesion (Fig. 4). Conversely, the prolonged retention of γδ T cells in the tooth extraction wound of WT ZOL mice may have contributed to PEH-like oral epithelial hyperplasia leading to the gross exposure of necrotic bone and the severe ONJ phenotype.

To better understand the “γδ T cell humanized” mice, h-γδ T cells and m-γδ T cells were co-cultured with mouse OC (m-OC). As expected, the activation of h-γδ T cells was observed with ZOL pretreated m-OC. However, co-culturing with m-OC without ZOL pretreatment also activated h-γδ T cells. The ZOL treatment of m-OC is expected to accumulate pAg such as IPP, which specifically activate h-γδ T cells. However, the surprising observation of h-γδ T cell activation by ZOL-untreated m-OC, albeit with a transient nature, should suggest a nonspecific activation of immune effectors by OCs. The activation of m-γδ T cells by m-OC with or without ZOL pretreatment further supports the previously unrecognized functions of OCs other than bone resorption, which might include regulatory roles of local immune effectors. Besides the pAg-derived activation, human and mouse γδ T cells are activated by peptide stimulants such as NKG2D ligands (43). It is tempting to speculate that ONJ pathogenesis may involve nonspecific activation of γδ T cells by OCs through the non-pAg activation mechanism in mice as well as in humans.

It is important to report that Rag2−/− ZOL mice with or without h-γδ T cell engraftment did not develop osteonecrosis (Fig. 7), whereas WT ZOL and Tcrd−/− ZOL developed osteonecrosis (Figs. 1, 3, and 5). It appears that the prolonged or unresolved oral inflammation is commonly observed in our mouse models as well as in ONJ patients. The time course profile of OC numbers at the surface of jawbone was similarly sustained in WT ZOL and Tcrd−/− ZOL mice (Fig. 5). By contrast, Rag2−/− mice lacking oral mucosal inflammation significantly reduced the recruitment of osteoclasts (Fig. 8A). Considering the mode of action of BP targeting OCs, it may not be a mere coincidence that osteonecrosis lesions were found to localize with or adjacent to a cluster of OCs (Fig. 3C). Therefore, the lack of OC in Rag2−/− ZOL mice may lead to the lack of osteonecrosis.

We previously proposed that ZOL-affected OCs might mediate the pro-inflammatory environment in oral mucosa based on the unique observation of a group of inflammatory cells closely associated with OCs in rat jawbones treated with ZOL (24). This study further suggested the role of OC on the immune regulation (Fig. 8B). Zhang et al. (44) demonstrated an unusually increased number of IL17-expressing helper T cells and M1 macrophages in the oral mucosa of ONJ patients. The ONJ patient demography with co-morbidities such as diabetes type 2 (45) may suggest the presence of systemic pro-inflammatory susceptibility. As such, the dysregulation of systemic as well as local oral mucosa immunity may provide the underlying core mechanism to the prolonged oral inflammation leading to the development of ONJ, in which γδ T cells may modify the oral epithelial disease phenotype.

**Author Contributions**—I. N., S. P., and A. J. designed this study. S. P., K. K., K. K., H. H. T., S. B., D. T. Q., A. J., and I. N. performed the experiments. S. P., J. W. S., A. J., and I. N. analyzed and interpreted the data. I. N. and A. J. drafted the manuscript. All authors agreed on the content of the manuscript. I. N. accepts responsibility for the integrity of the data analysis.

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