Dendritic cell derived exosomes loaded with immunoregulatory cargo reprogram local immune responses and inhibit degenerative bone disease in vivo

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ABSTRACT

Chronic bone degenerative diseases represent a major threat to the health and well-being of the population, particularly those with advanced age. This study isolated exosomes (EXO), natural nano-particles, from dendritic cells, the “directors” of the immune response, to examine the immunobiology of DC EXO in mice, and their ability to reprogram immune cells responsible for experimental alveolar bone loss in vivo. Distinct DC EXO subtypes including immuno-regulatory (regDC EXO), loaded with TGFβ1 and IL10 after purification, along with immune stimulatory (stimDC EXO) and immune “null” immature (iDCs EXO) unmodified after purification, were delivered via I.V. route or locally into the soft tissues overlying the alveolar bone. Locally administrated regDC EXO showed high affinity for inflamed sites, and were taken up by both DCs and T cells in situ. RegDC EXO-encapsulated immunoregulatory cargo (TGFβ1 and IL10) was protected from proteolytic degradation. Moreover, maturation of recipient DCs and induction of Th17 effectors was suppressed by regDC EXO, while T-regulatory cell recruitment was promoted, resulting in inhibition of bone resorptive cytokines and reduction in osteoclastic bone loss. This work is the first demonstration of DC exosome-based therapy for a degenerative alveolar bone disease and provides the basis for a novel treatment strategy.

Introduction

Periodontitis (PD) is a chronic bone disease that affects over 50% of the U.S. population [1]. PD has been linked to increased risk of a number of other diseases with more serious mortality and morbidity profiles [2,3]. Histopathological studies of PD lesions implicate neutrophils as the first line of defense, which eventually turn against the host [4]. Severe PD lesions are infiltrated with B cells, macrophages and dendritic cell (DC) clusters with CD4+ T cells [5–8], the equivalent of ectopic lymphoid follicles [9,10].

The immune response can be shaped based on the maturation status of DCs [11]. Immature DCs (iDCs) can promote immune tolerance by induction of T cell anergy and regulatory T cell (Treg) response to terminate inflammation [12–17]. Whilst mature DCs can direct a T-helper cell 17 (Th17) response to eradicate invading bacteria. Th17 effector T cells appear to play a bone degenerative role [13,18,19], while Tregs reportedly attenuate inflammatory bone loss [20,21]. Restoration of Treg stability in PD patients by implementation of an anti-infective protocol correlates closely with clinical improvement [22], but as yet no effective immunomodulatory agent for PD has been identified.

The marked ability of DCs to direct the immune response has led to use of whole DCs, modified to be...
Table 1. Cytokine and DCs maturation marker CD86 profile in DCs derived exosomes subsets.

|                  | regDCs EXO | iDCs EXO | stimDCs EXO |
|------------------|------------|----------|-------------|
| TGFβ1            | +          | -        | -           |
| IL10             | -          | -        | +           |
| IL6              | -          | -        | +           |
| TNF              | -          | -        | +           |
| IL18             | -          | -        | +           |
| CD86             | -          | +        | ++          |

+ Detectable
- Below Detectable limit

 tolerogenic genetically or with immunosuppressive cytokines, to suppress inflammatory disease in animals. However, immunomodulation using DCs or other immunosuppressive cells like Treg has some limitations, including rarity and phenotypic instability of whole cells in vivo [23].

Targeting proinflammatory cytokines using inhibitors of TNF and IL1B has shown therapeutic efficacy for treating inflammatory diseases. However, effects are short lived due to rapid clearance and proteolytic degradation in challenging inflammatory environment [24].

Extracellular vesicles (EV) of endosomal origin, called exosomes (EXO), can be isolated from DCs, and their small size (30–150nm) can optimize the delivery of this cargo to recipient cells. Moreover, DC EXO harbour unique proteins that shield them from attack by the complement system [25] and promote binding to tissue integrins [26]. DC EXO can also be loaded with cytokines or peptides using direct or indirect approaches [27–29]. Immunostimulatory cargo-loaded mature DCs EXO have been touted for anti-cancer benefits [30], while tolerogenic DC-derived EXO equipped with immunoregulatory cargo offer promise for treatment of autoimmune and inflammatory diseases [31].

The goal of the current study was to characterise the immunobiology of DC EXO subtypes in vitro and in vivo and their ability to reprogram immune cells responsible for inflammatory bone loss. After purification, reg DC EXO were loaded with immunoregulatory cytokines TGFβ1 and IL10. These cytokines appeared localized on the EXO transmembrane domain and within the EXO lumen, where they were protected from proteolytic degradation. Both in vitro and in vivo studies show important role for EXO-encapsulated TGFβ and IL-10 in prevention of DC maturation; moreover, TGFβ1 was required for increased induction of CD25+Foxp3+T cells (Treg). Moreover, regDC EXO inhibited Th17 and decreased bone loss, further evidenced by decrease in Trap+ osteoclasts.

We conclude that DC EXO loaded with molecular cargo to modulate Th17/Treg balance is an effective immunotherapeutic approach to regulate degenerative bone disease in vivo in this model.

Material and methods

Ethics statement

The Institutional Animal Care and Use Committee (IACUC) of Augusta University (protocol # 2013–0586) approved all experimental procedures.

Generation and culture of iDCs, regDCs and stimDCs

Bone marrow was isolated from tibias and femurs of 6 – to 8-week-old mice as previously described [32]. ACK cell lysis buffer was used to lyse contaminating erythrocytes (Invitrogen, Thermofisher scientific, and Columbia, SC, USA). Cells were cultured for 24 h in complete media (RPMI 1640 containing 10% FBS and 100 IU/ml penicillin/streptomycin) to remove adherent macrophages. Non-adherent cells were then cultured in growth media, containing 20ng/ml of murine GM-CSF and IL-4 (Peprotech, Rocky Hill, NJ, USA). Culture media was changed every 2days and cells were harvested on day 6 and incubated for 2days in EXO depleted complete media (by using EXO free FBS) to generate iDCs. To generate stimDCs, part of the harvested cells were cultured in fresh EXO depleted complete medium containing 1ug/ml LPS (Sigma, St. Louis, M. O., USA) for 48h, and cells were harvested on day 8. To generate regDCs, DCs were cultured for 4days and harvested on day 5 for TGFβ1/IL10 recombinant cytokines treatment in which, 1x10⁷ DCs were incubated for 2hours with 1ug/ml TGFβ1 (R&D Systems, Inc. Minneapolis, MN) and 1ug/mL of the recombinant murine IL-10 (Cell Sciences, Canton, Massachusetts) in total volume of 1mL serum-free media, then diluted 1:10 in fresh complete media for further incubation. On day 6, regDCs were harvested, washed and cultured for 48h in EXO depleted growth media and isolated on day 8. On day 8, culture supernatants were collected for EXO purification. Cultured iDCs, regDCs and stimDCs were defined by expression level of CD11c+ (N418) (Invitrogen), MHCII (M5/114.15.2) (Milteny biotech Auburn, CA,USA) and CD86 (GL1) (Invitrogen), by flow cytometry (Milteny biotech) and by level of pro/anti-inflammatory cytokine mRNA by
PCR, including IL6 (Mm00446190_m1), IL12 (Mm01288998_m1), IL23 (Mm00518984_m1), TGFB1: Mm01178820_m1 and TNF: Mm00443258_m1, (Thermofisher Scientific).

**EXO isolation, purification and cytokine loading of regDC EXO**

EXO isolation was performed as previously described [24]. Briefly, the DC culture supernatants was subjected to three successive centrifugations at 500g for (5 min), 2000g for (20 min), and 10,000g for (30 min) to eliminate cells and debris, followed by ultrafiltration 3× with 0.2 um and 3× with 100 kDa filters (to remove free proteins) and ultracentrifugation for 1.5h at 120,000g. To further remove excess free proteins, EXO pellets were washed with a large volume of PBS and ultra-centrifuged 2× at 120,000g for 1.5h, and finally re-suspended in 100 ul of PBS for further studies. In the case of regDC EXO, 1×10⁹ particles were additionally actively loaded by sonication [27] with 5ug TGFB1 and 5 ug IL10 in 500 ul of PBS then filtered 3x by ultrafiltration with 100KDA filter to remove free proteins and washed 3× with large volume of PBS and ultra-centrifugation at 120,000g for 1.5h to further purify EXO from free molecules, and finally re-suspended in 100 ul of PBS. Supernatant of which regDCs EXO were suspending was isolated and checked for any contaminants by ELISA. The integrity of EXO proteins and loaded cytokines was assessed after sonication using Western blot analysis (Fig. S1A). The size and zeta potential of sonicated exosomes was measured using Nano particle tracking analysis (Fig. S1B). EXO subtypes were quantified by nano-tracking analysis and a BCA protein assay. The yield of exosomes from their respective DC subtypes are as follows:

- RegDCs EXO: approximately four vesicles per one cell per hour
- ImDCs EXO: approximately five vesicles per one cell per hour
- StimDCs EXO: approximately eight vesicles per one cell per hour

**Characterization of DC-Derived EXO**

**Western blotting**

EXO lysates were extracted to determine EXO markers and pro/anti-inflammatory cytokines by Western blotting analysis using anti-TSG101 (MA1-23,296), anti-Alix (MA1-83,977), anti-CD63 (10628D) and GRP94 (MA3-016) from (Invitrogen, Thermofisher scientific West Columbia, SC, USA), anti-MHCII (MABF33) from (Sigma, St. Louis, M.O.,USA), anti-IL1B (63,124), anti-TNF (11,948), anti-iL6 anti-TGFB1 (3711) and anti-B-Actin (3700) from (Cell Signaling Technology, Danvers, MA, USA) and CD86 (AF-441-NA) and anti-IL10(MAB417) from (R&D Systems, Inc. Minneapolis, MN, USA)

**Electron microscopy**

As described previously [33] EXO samples were loaded onto a copper grid. After precipitation of EXO, the sample liquid was extracted and counter stained with 2% phosphotungstic acid solution for 10min then placed under an incandescent lamp for 5 min. EXO sample was examined with TEM. Immune gold plating was done using anti-CD63, anti-TGFB1 and anti-IL10 primary antibodies.

**Nanoparticle tracking analysis**

Nanoparticle tracking analysis (NTA) was used to visualize and quantitate size and count of nanoparticles in suspension. Briefly, as described previously [33–36], 10 ul of EXO suspension was loaded into the sample chamber of ZetaView PMX 110 (Particle Metrix, Meerbusch, Germany). Data analysis was performed with software (ZetaView 8.02.28).

**EXO miRNA microarrays**

miRNAs were isolated from EXO using Total EXO RNA Kit (4,478,545, Thermofisher Scientific) according to manufacturer’s protocol. The concentration of miRNA was determined using a NanoDrop spectrophotometer (Thermo Scientific) and the quality of miRNA analysed using an Agilent 2100 Bioanalyzer. Analysis of miRNAs was performed using an Affymetrix GeneChip® miRNA 4.0 Array at the Integrated Genomics Core, Augusta University, GA. The P-value cut-off of 0.05 and the miRNAs with a fold change above 1.5 were considered differentially expressed. Bioinformatic analysis was performed using Ingenuity Pathway Analysis (IPA), and TargetScan software was used to identify computationally predicted miRNA-mRNA target relationships [37].

**ELISA**

EXO lysed with ELISA lysis buffer (#7018, Cell signaling) or intact were suspended in PBS and analysed according to the manufacturer, using ELISA kits for TGFB1 (BMS608-4) and IL10 (BMS614NS) both from (Invitrogen, Thermofisher scientific West Columbia, SC, USA)

**EXO resistance to proteolytic digestion assay**

RegDCs EXO and Free TGFB1 and IL10 with equivalent concentration to those loaded to regDCs EXO were divided to trypsin or proteinase k treated at 37°
C with concentration of 25μg/ml for 60min and non-treated groups and subjected to Western blotting for TGFβ1 and IL10 detection.

**T-cell isolation**

CD4+ T cells were isolated from mice spleens using negative selection with Mouse T-cell Enrichment Kit (#19,765) (Stem cell technologies, Cambridge, MA, USA). T-cell purity was assessed by flow cytometry analysis of CD4, CD3 and CD8 markers and were typically >95% pure.

**EXO uptake invivo**

For EXO uptake study in vitro, EXO labelled with Dil (D282, Thermofisher Scientific) were co-cultured with DCs or CD4+T cells for 24h. Cells were fixed and stained on glass slides with phalloidin (A12379) and DAPI (D1306) (Invitrogen, Thermofisher scientific West Columbia, SC, USA). The images were obtained by scanning confocal fluorescence microscopy.

**Immune modulatory effect of EXO in vitro**

**DCs activation:** The immune modulatory influence of EXO subtypes on acceptor DC was investigated by incubating 10^7/ml EXO in DC culture on day 5, followed by harvesting on day 8 and measuring the expression of antigen presenting and maturation markers MHCII and CD86 by flow cytometry and mRNA level of proinflammatory cytokines including IL6, IL12 and IL23. Analyses were performed in the presence or absence of DC stimulators/inhibitors including LPS and neutralizing antibodies against TGFβ1 (MAB1835, R and D) and IL10 (JES5-2A5, Thermo fisher scientific). Phosphorylation of TGB1 and IL10 transcription factors and MHCII was assayed by Western blot using anti PSMAD2/3 (D6G10), anti SMAD2/3 (D7G7), anti PSTAT3 (D3A7), antiSTAT3 (D3Z2G) and MHCII (MABF33) antibodies and GAPDH (D16H11) (Cell Signaling Technology, Danvers, MA, USA).

**Antigen-presentation**

T-cells specific for OVA antigen peptides were isolated from spleen of OT-II transgenic mice and stained with carboxyfluoresceine succinimidyl ester (CFSE) (C1157, Thermo fisher scientific) followed by washing to remove excess free CFSE.10^8/ml treated – (EXO added to DCs culture on day 5) or untreated (control) – DCs were incubated with OVA peptide (Q1641, Sigma Aldrich) 1μg/ml for 24hours, then washed and cocultured with CD4+T-cells, at a ratio of 1:10 DCs to T-cells in a 96 well round bottom plate and complete RPMI 1640, and incubated at 37° and 5% CO2 for 5days. T-cell proliferation was assessed by % loss of CFSE in each generation by flow cytometry.

**T-cell activation and differentiation**

To stimulate Tregs or effector CD4+ T-cell proliferation, 96 round-bottom plate were coated and incubated overnight at 4°C with 10μg/ml anti CD3 antibody. CFSE-stained T-cells were added with or without EXO 10^8/ml to the plate in complete RPMI 1640 media containing 10μg/ml anti CD28 antibody and incubated at 37° and 5% CO2 for 5days. In some experiments, TGFβ1 and IL10 neutralizing antibodies were added. T-cell proliferation was assessed by CFSE dilution using flow cytometry analysis by gating on the CD4+ cells. Regulatory T-cells (Tregs) were identified by gating on double positive cells for FOXP3 (FJK-16s, Thermo fisher Scientific) and CD25 (PC61.5, Thermo fisher Scientific), in CD4+ (GK1.5, Thermo fisher Scientific), population while T-helper 17 induction was identified by measuring IL17 median fluorescent intensity in CD4 population by using anti IL7 Antibody (eBio17B7, Thermofisher scientific), followed by flow cytometry. Supernatants were analysed for IL17 by ELISA (BMS6001, Thermo fisher Scientific). Western blot was used to assess the expression of FOXP3 using antiFOXP3 (FJK-16s, Thermo fisher Scientific).

**In vivo imaging of EXO biodistribution in murine periodontitis model**

**Induction of experimental periodontitis (PD) in mice model**

This model was developed by ligation of the upper right second molar with black silk suture to accumulate bacteria and induce inflammation induced alveolar bone loss as described previously [38,39]. This model does not induce inflammatory bone loss in germ free animals and is thus dependent on the endogenous flora.

**In vivo imaging:** 1.5 to 2 mCi of Indium-111-oxine (Anazao Health Corporation, Tampa, FL, USA) in PBS was added to 200μl of exosomes particles (~2 × 10^9 particles) and incubated at 37°C for 20minutes. Free indium was removed by repeated PBS washes through an Amicon ultrafiltration device. Collected In-111-labelled EXO were diluted to 200 μCi of radioactivity per dose and injected intravenously in tail vein or locally in palatal tissue in periodontitis mice model on day 3 relative to ligature placement. Control...
animals received injection of equivalent activity of free Indium-111-oxine. Whole body and head single photon emission spectroscopy (SPECT) images were acquired by Mediso’s nanoScan microSPECT/CT system ((Mediso, USA) at 24h after injection, and images were reconstructed to determine radioactivity in maxilla and whole body. Radioactivity in maxilla was expressed in percent of activity in the whole body (total radioactivity). Maxilla were excised and ex vivo radioactivity measurements were performed by gamma counter (Perkin Elmer Packard Cobra II Auto-Gamma) [40].

**EXO uptake in vivo in murine periodontitis models**

For EXO uptake *in vivo*, Dil-labelled EXO were locally injected through palatal gingiva on day 3 relative to ligature placement. After 24 h, mice were sacrificed and gingival tissues from maxilla were removed and treated with Collagenase type II (2mg/mL) and DNase I (1mg/mL) (both from Sigma Aldrich) solution in PBS plus 2% FCS for 30min at 37°C in a shaker bath. 0.25μL of 0.5M EDTA per 2mL sample were added and incubated for an additional 15minutes, followed by centrifugation at 4°C, 400 × g for 8min in PBS with 2% FBS. Supernatants were removed, cells were suspended in PBS with 2% FBS and filtered through a 70-μm cell strainer and stained on glass slide with primary antibodies including antiCD11c (N418), antiCD4 (OK1.5) and Secondary antibodies including Goat anti-Rat IgG, DyLight 488 (SA5-10,018) and Goat anti-Hamster IgG, Alexa Fluor 488 (A-21110), all from thermofisher Scientific. Then slides were mounted with DAPI and images were captured by scanning confocal fluorescence microscopy.

**Immune modulatory effect of EXO in vivo**

The immunomodulatory functions of EXO in six experimental groups were analysed. **Group 1** received locally 10 ul of phosphate buffer saline (PBS) (n=5). The other five groups of animals were subjected to ligation with 5–0 silk to the upper left second molar to induce inflammatory bone loss. The contralateral molar tooth in each mouse was left un-ligated to serve as baseline control for alveolar bone volume measurements. The ligatures remained in place in all mice throughout the experimental period (nine days). **Group 2** received ligation plus local administration of 10 ul of PBS (n=5), **Group 3** received ligation plus local delivery of 10⁸ particles of regDCs EXO suspended in 10 ul PBS (n=5), **Group 4** received ligation plus local delivery of 10⁸ particles of iDCs EXO suspended in 10 ul PBS (n=5), **Group 5** received ligation plus local delivery of 10⁸ particles of stimDCs EXO suspended in 10 ul PBS (n=5), and **group 6** received ligation plus 60 ng TGFβ1 and 10 ng IL10 which are equivalent concentrations to TGFβ1 and IL10 in regDCs EXO as determined by ELISA. The delivery of control PBS or EXO in all groups was performed at days −2, 0 and 2 relative to ligature placement. After 9 days of ligature placement maxilla and gingival tissue were harvested, and gingival cells were isolated. Cells were stained with antibodies against CD3 (17A2), CD86 (GL1), CD11c (N418), CD25 (PC61.5), CD4 (GK1.5), Foxp3: (FJK-16s), IL17A (eBio17B7), all from Thermofisher Scientific and MHCII (M5/114.15.2, Milteny biotech), in order to identify DCs and their activation markers and T cells subsets infiltrating gingival tissue. Draining lymph nodes were excised and cells were isolated and stained with anti-CD4 antibody to determine frequency of CD4+T cells by flow cytometry.

**Immunohistochemical (IHC) staining of gingival tissue**

Tissue blocks were de-paraffinized, cut into 5 μ sections and mounted on slides. Antigen retrieval was done in water bath heating with citrate buffer PH6. IHC staining with DAB/peroxidase was performed using the following antibodies against the proteins indicated: MHCII (ab180779, Abcam, Cambridge, MA), CD86 (GL1), FOXP3 (FJK-16s), IL17 (PA5-79,470), RANKL (PA5-87,147), all from Thermofisher Scientific and TNF (SAB4502982, Sigma Aldrich). Multiple random microphotographs were taken for each gingival tissue in lamina propria region, at 40× objective lens using Zeiss microscope (Zeiss AxioIma, Carl Zeiss Microscopy GmbH, Jena, Germany). Image-J software was used and a threshold was adjusted based on the negative control.

**Micro-CT imaging and bone parameter analysis**

Micro-CT imaging of maxillae was performed using a SkyScan 1272 microfocus X-ray system (SkyScan®, Kontich, Belgium) with software including NRecon reconstruction*, CTAn 1.8*, and CTvol. In this study, the X-ray source set at 70 kV and 100μA with the pixel size of 6μm, a 0.5-mm filter, and a tomographic rotation of 180° (rotation step of 0.2°). Residual bone volume (BV, μm³) around second upper molars was quantified and 3-D models were generated with teeth auto traced in yellow while bone in white [39,41].

**Periodontal tissue histology**

Formalin-fixed maxillae specimens were decalcified with EDTA and embedded in paraffin. Sections were
stained with haematoxylin and eosin, to evaluate alveolar bone loss at microscopic level.

**TRAP staining**

As described previously [42], TRAP staining for periodontal tissue sections used Acid Phosphatase, Leukocyte (TRAP) Kit (387A-1KT, Sigma Aldrich) to quantify osteoclastic cells, revealed by light microscopy. Multinucleated osteoclasts in interproximal bone and furcation were counted blindly and osteoclast density was expressed as number of osteoclasts per square micrometre of bone using computer software (ImageJ) [42].

**In vitro osteoclastogenesis assay**

1x $10^5$ EXO were co-cultured with murine CD4T cells in osteoclast assay 96 well plates (CLS3988, Sigma Aldrich) with 1x$10^5$ bone marrow cells and supplemented with 50ng/ml of RANKL (ab129136, Cambridge, MA) and 50ng/ml M-CSF (315-03, Rocky Hill, NJ, USA). The culture media was changed every 2 days. On day 5 TRAP staining was performed to detect multinucleated osteoclasts using Acid Phosphatase Leukocyte (TRAP) Kit (387A-1KT, Sigma Aldrich).

**Flow cytometry and antibodies**

Staining was performed on ice, with Flow Cytometry Staining Buffer (Thermofisher Scientific). Blocking of FC receptors (FCR) was achieved with mouse FcR blocking reagent (Miltenyi Biotec) for 15 minutes on ice and protected from light, followed by addition of fluorophore conjugated antibody at recommended concentration on ice for 30 minutes, after which cells were washed, re-suspended in flow cytometry buffer and data was acquired using Milteny biotech machine and software.

**Real time PCR**

Total RNA was isolated from DCs *in vitro* and from gingival tissue of the experimental groups used for *in vivo* studies using QIAGEN RNAeasy mini kit (Qiagen, Inc., Valencia, CA, and USA). RNA concentration and purity were assessed with Nanodrop (NanoDrop 1000 UV-VIS Spectrophotometer Software Ver.3.8.1, Thermofisher Scientific). Ratio of 260/280 of 2.0 was deemed acceptable for further analysis, and was reverse transcribed to cDNA. Amplification by PCR was performed using the High-Capacity cDNA Reverse Transcription Kit and PCR in total reaction of 20μL. Quantitative real-time PCR was performed using TaqMan gene expression primers specific for IL6 (Mm00446190_m1), IL12 (Mm01288989_m1), IL23 (Mm00518984_m1), TGFB1 (Mm01178820_m1) and TNF (Mm00443258_m1) IL10(Mm01288386_m1), FOXP3 (Mm00475162_m1), CTLA4 (Mm00486849_m1), Rankl (Mm00441906_m1) and Beta Actin (Mm02619580_g1), all from Thermofisher Scientific. RT-PCR was run in StepOnePlus Real-Time PCR System. Relative gene expression was calculated using delta-delta CT and plotted as relative fold change.

**Western blotting**

Cells or EXO lysates were extracted by addition of RIPA buffer supplemented by protease/phosphatase inhibitor cocktail and incubation for 20 minutes on ice. Proteins (10μg) were separated using 14% MINI-PROTEAN TGX Precast Protein Gel (Bio-Rad Laboratories, Hercules, CA), and transferred onto PVDF membranes (Sigma-Aldrich). After blocking with 5% non-fat dry milk in PBS, the membrane was incubated with primary antibodies, washed with TBST and incubated with HRP-conjugated secondary antibodies for 1 hour at room temperature. The membranes were developed using ECL kit and imaged using ChemiDoc MP Imaging Gel (Bio-Rad Laboratories, Hercules, CA).

**Statistical analysis**

Data analysis was performed by one-way ANOVA followed by Tukey’s multiple-comparisons test or student T-test using GraphPad Prism 6 (GraphPad Software, La Jolla, CA). Values are expressed as mean ± standard deviation (SD) and experiments were done in triplicates.

**Results**

**Generation and characterization of DC EXO subtypes prior to in vitro and in vivo functional testing.**

EXO donor DC subtypes including immunoregulatory DCs (regDCs), immature DCs (iDCs), and immunostimulatory (stimDCs) were generated from bone marrow derived non-adherent cells of C57BL/6 mice, and phenotype and cytokine profiles characterized as follows: regDCs: CD11c+, low MHCII+, low CD86+, low CD80+ and low CD40+, iDCs: CD11c+, intermediate MHCII+, intermediate CD86+, intermediate CD80+, intermediate IL6+, intermediate IL12+ and intermediate IL23+ and stimDCs: CD11c+, high MHCII+, high CD86+, high
CD80+, high IL6+, high IL12+ and high IL23+. From these DC subtypes, EXO were purified, quantitated and correct size distribution (30–150nm) confirmed by nanoparticle tracking analysis (Figure 1A). Further characterization indicates predominant population of EXO, as determined by immunogold-CD63 TEM (Figure 1B), immunoblot for Tsg101 (component of Escort1 complex), CD81, B-actin, Alix (Escort Associated Protein) and EXO negative marker GRP94 (Figure 1C), as previously reported [43].

Additional active loading by sonication of regDC EXO with TGFβ1 and IL10 was achieved, residual proteins removed by further processing, and all DCs EXO subtypes analysed by immunoblot (Figure 1C) and ELISA (Figure 1D) revealing distinct molecular content of EXO subtypes as follows: regDCs (high TGFβ1/IL-10, undetectable IL-6/TNF/IL-1B and CD86), iDCs (undetectable TGFβ1/IL-10, IL-6/TNF/IL-1B, low CD86) and stimDCs (undetectable TGFβ1/IL-10, high IL-6/TNF/IL-1B and CD86) (Table S1). The relative levels of TGFβ1 and IL-10 in both lumen and transmembrane domain of regDC EXO was determined as in materials and methods, and as shown in (Figure 1D and Figure 1E). Neither TGFβ1 nor IL10 were detectable in iDC EXO and stimDC EXO, nor in the PBS supernatant of regDC EXO after isolation, ruling out soluble carry over. Higher magnification TEM of regDC EXO confirms ELISA data, with TGFβ1 inside and attached to outer EXO membrane (Figure 1F). Sonication of the exosomes did not affect the stability of exosomal protein markers such as Alix, TSG101 and tetraspanin CD81 as shown by Western blot analysis (Fig. S1A). Furthermore, the sonication did not influence the size or zeta potential of regDCs EXO as shown in Fig. S1B.

Figure 1. RegDCs EXO protect encapsulated immunoregulatory cargo (TGFβ1 and IL10) from proteolytic degradation. (A) Nano-tracking analysis to determine EXO number and size distribution in nm. (B) Transmission electron microscopy (TEM) to visualize EXO shape and immuno-gold TEM to detect EXO marker tetraspanin CD63, showing unstained (left) and positive staining (right)(arrows). (C) Western blotting to detect other EXO-related markers including CD81, TSG101, GRP94 and B-actin in donor DCs and EXO (left) and anti/pro-inflammatory cytokines including TGFβ1, IL10, IL6, IL1B and TNF and the costimulatory molecule CD86 as well as EXO associated proteins ALIX and TSG101 in DCs EXO subsets (right). (D) TGFβ1 and IL10 content of lysed EXO and in regDCs EXO supernatant by ELISA. (E) TGFβ1 and IL10 content of non-lysed EXO (transmembrane domain) by ELISA. (F) Immunogold TEM to detect luminal and transmembrane TGFβ1 in regDC EXO. (G) regDCs EXO (left) or equivalent concentration of free TGFβ1 and IL10 (right) were treated with trypsin (upper panel) or proteinase-K (lower panel) and incubated in control buffer (1hour at 37°C) and analysed by western blotting to detect the levels of TGFβ1, IL10 and exosomal markers TSG101 (upper panel) and CD81 (lower panel).

RegDC EXO induce an immune regulatory effect on acceptor DCs mediated by TGFβ1 and IL10 in vitro

Co-culture of Dil-labelled regDC EXO with iDCs confirmed EXO internalization by acceptor DCs in vitro as shown by confocal microscopy analysis. Images show Dil-EXO (red) internalized by recipient DCs.
counterstained with phalloidin (green) (Figure 3A). Expression of co-stimulatory molecules required for T cell stimulation or maturation markers of acceptor iDCs were modulated by different EXO subtypes. This was evidenced by differential MHCII and CD86 expression on DCs shown by flow cytometry analysis. While reg DCs EXO significantly inhibited maturation markers of acceptor DCs, as evidenced by decrease in % CD11c+ MHCII+ and CD11c+ CD86+ expression, compared to control, stimDC EXO had the opposite effect, increasing %CD11c+ MHCII+ and CD11c+ CD86+ expression, with iDC EXO-treated DCs intermediate of the two. (Figure 3B and Figure 3C). Cytokine profiling for IL-6, IL-12 and IL-23 mRNA further confirmed immuno-regulatory, - stimulatory or null phenotype of respective EXO (Figure 3D). iDCs co-cultured with DC EXO subtypes were analysed by immunoblot for activated transcription factors of IL-10 (pSTAT3) and TGFβ1 (pSMAD2/3) signalling and total STAT3 and SMAD 2/3. RegDC EXO induced highest level of pSTAT3 and pSMAD2/3, consistent with TGFβ1 and IL-10-mediated signalling, respectively. RegDC EXO induced lowest level of MHCII (Figure 3E).

To assess the integrity and stability of EXO during incubation for multiple days in culture media, we performed immunoblot analysis of exosomal protein markers and loaded cytokines (TGFβ and IL-10) in EXO after incubation in serum free culture media for 0, 1, 3, 5 and 7 days. Results indicate that the EXO marker, Alix and the anti-inflammatory cytokine IL10, were stable until day 1 followed by a subtle decrease over time. However, stability of CD81 did not seem to be influenced by incubation for longer times. TGFβ-1 showed stability at all time points (Fig. S2A). The size and zeta potential showed minimal changes (Fig S2B).

To determine the role of TGFβ and IL-10 in inhibition of DC maturation by regDC EXO neutralizing antibodies to TGFβ1 and IL10 were added. Complete restoration of MHCII expression was achieved by neutralization of both TGFβ-1 and IL-10 while either alone was not sufficient for restoration (Figure 3F). These results suggest the crucial role of both cytokines in the inhibition of acceptor DC maturation by regDC EXO.

**RegDC EXO increase acceptor DC resistance to LPS mediated maturation and lower antigen presenting ability**

RegDC EXO had the added capability to prevent *Escherichia coli* LPS-mediated maturation of iDCs, as evident by a significant decrease in MHCII (Figure 4A),
CD86 (Figure 4B) expression shown by flow cytometry analysis and downregulation of mRNA level of IL-6 (Figure 4C). The ability of DC EXO to regulate Ag-presentation in acceptor DCs was examined by proliferation of splenic OVA-specific T cells in response to OVA-iDCs. OVA-specific Ag-presentation by DCs was significantly reduced by regDC EXO and increased by stimDCs EXO-treated OVA-DCs (Figure 4D).

RegDC EXO is also taken up by CD4T-cells inhibiting proliferation and promoting TGFb1 mediated Tregs induction in vitro

Dil-labelled DC exo were co-cultured with splenic T cells to determine their ability to internalize DC exo. Scanning confocal microscopy revealed internalized regDC EXO inside Tcells (Figure 5A). Only regDC EXO significantly increased TCR-driven Tregs (CD25/FOXP3+) induction, an effect that was abrogated by blocking TGFb-1 and not IL-10 using neutralizing antibody as shown by flow cytometry (Figure 5B and Figure 5C) and Western blotting (Figure 5D). These results suggest that TGFb-1 has a more dominant role in regulating Treg induction. RegDCs EXO also suppressed anti-CD3/CD28 antibodies mediated T cell proliferation while stimDC EXO increased proliferation in vitro (Figure 5E). StimDCs EXO were uniquely able to induce T-helper 17 cells, as measured by flow cytometry (Figure 5F) and ELISA (Figure 5G).

Figure 4, RegDCs EXO increase acceptor DC resistance to LPS mediated maturation and lower antigen presenting ability. Flow cytometry histograms showing in vitro influence of EXO subtypes on MHCII (A) and CD86 (B) expression on acceptor DCs challenged by LPS (left). Results presented as median fluorescent intensity measurements in representative bar graph (right). (C) IL-6 mRNA expression in acceptor DCs by PCR analysis. (D) Flow cytometry histograms showing proliferation of splenic ovalbumin specific CD4+T – cells, labelled with CFSE, after coculture with ovalbumin – pulsed DCs treated with or without EXO subtypes. Results shown are representative of three independent experiments (* P<0.05 by one-way ANOVA followed by Tukeys multiple comparisons).

To determine effects of T cells incubated with regDCs EXO on osteoclastogenesis, regDCs EXO-treated T cells were co-cultured with pre-osteoclasts, inhibiting osteoclastogenesis relative to non-EXO-treated control T cells, as demonstrated by TRAP staining (Fig. S3).

In vivo retention of DC EXO in gingival site of disease induction.

Initial in vivo studies focused on the bio distribution of EXO, 24hrs after IV or local palatal injection at the site of ligature-induced periodontitis (PD). Whole-body imaging in live mice of radiolabeled (In-111) DC EXO was performed using SPECT-CT. Control mice were administered equivalent activity of free In-111 by both routes. Free In-111 was rapidly cleared after IV administration, while In-111-labelled EXO were cleared slowly and bio distributed to spleen and other organs, but not maxilla (Figure 6A). In-111-labelled EXO administered in palatal maxillary tissue persisted at the site of experimental PD in maxilla, ostensibly from EXO adhesion to inflammatory integrins as reported [48], but also were observed in local lymph nodes. In contrast, free In-111 injected in the gingiva redistributed rapidly to other non-target extraoral body sites (Figure 6B). It should be noted that gingival injection of In-111-labelled EXO in inflamed diseased site yielded 10x higher radioactivity than injection of free In-111 (Figure 6C). Analysis of excised maxilla post-mortem also showed significant increase in radio-activity in In-111-labelled EXO injected group compared to free In-111, confirming in vivo SPECT-CT data (Figure 6D). Taken together, these data suggest higher retention of reg DC EXO in inflamed gingival tissue, where it persists exerting its action on resident immune cells leading to their reprogramming and modulation of immune response. 3D-video of In-111-labelled EXO biodistribution data are shown in Fig.S5.

Uptake of injected EXO by gingival DCs/T cells, with subtypes differentially regulating DC maturation and Treg-Th17 infiltration in vivo

Mice were injected in the palatal gingiva with buffer (no EXO), Dil-labelled regDC EXO, iDC EXO, stimDC EXO, or TGFb/IL-10 (at equivalent concentrations to those carried in regDC EXO) in buffer on days –2, 0 and 2. Ligatures were placed on day 0 to promote bacterial accumulation and induce inflammatory alveolar bone loss, as detailed in materials and methods. Negative control group received neither ligatures nor EXO, while positive control group received ligatures but no EXO. On day 9, animals were sacrificed and disaggregated gingival cells were analysed by confocal microscopy for association of labelled EXO with...
regDCs EXO uptake by CD4+T-cells, promoting TGFβ1 dependent Tregs induction while stim DCs EXO induce T-helper17 response in vitro. (A) Uptake of Dil labelled EXO (red) by splenic CD4+T-cells, showing DAPI-labelled nuclei, and phalloidin (green) labelled cell membrane, as visualized under confocal microscopy. (B) Flow cytometry scattergrams showing in vitro influence of EXO on induction of T-regulatory cells, as measured by % of double positive CD25 and Foxp3 cells in gated CD4+T-cell population, stimulated with antiCD3/CD28 antibodies. (C) Summary bar graph of flow cytometry data. (D) Western blotting analysis of Foxp3 expression in acceptor CD4+T-cells. (E) Flow cytometry analysis showing proliferation % of CFSE-labelled splenic CD4+T-cells, as expressed by histograms (left panel) and bar graphs (right panel), after stimulation with anti-CD3/CD28 in presence or absence of EXO subtypes. (F) Flow cytometry analysis of CD4 +IL-17+T cells %, as expressed by histograms (left) and bar graph (right). (G) ELISA of IL17 expression in supernatant of T cells treated with EXO subtypes. Results shown are representative of three independent experiments (* P<0.05 by one-way ANOVA followed by Tukeys multiple comparisons).

CD11c+DCs and CD4+T cells. Both DCs (Figure 7A) and T-cells (Figure 8A) are observed colocalized with EXO in situ. The influence of EXO on gingival DC maturation status was determined by flow cytometry analysis of MHCII and CD86 expression on isolated CD11c+DCs from 6 experimental groups. As observed in vitro, regDC EXO injected in vivo induced significant decrease in MHCII and CD86 expression (Figure 7B, C) on gingival DCs, further confirmed by immunohistochemical staining for CD86 (Figure 7D and Figure 7E). Furthermore, only in the regDC EXO-treated group was a significant increase in Tregs and decrease in %CD4+IL-17+T cells observed (Figure 8B, C). Analysis of FOXP3, CTLA4 and IL10 mRNA further confirms the immunoregulatory influence of regDs EXO in situ (Figure 8C). Immunohistochemical analysis of gingival tissue in situ revealed higher positive staining for Foxp3 (Figure 8D) and lower positive staining for IL17 (Figure 8E) in regDC EXO + ligature-treated group in comparison to no EXO +ligature group. Due to observed bio distribution of labelled EXO to local lymph nodes, CD4+T cells were quantified in draining lymph nodes, revealing a significant decrease in frequency of CD4+T-cells in regDC EXO-treated group (Fig. S4A and Fig. S4B). These data show that both in vitro and in vivo, regDC EXO-mediated reprogramming of DCs and T cells towards an immune modulatory phenotype.

Differential regulation of local bone resorbing mediators and osteoclast density by DC EXO subtypes

Analysis of mRNA in gingival tissues from experimental groups reveals a significant decrease in RANKL (Figure 9A) and TNF (Figure 9B) in regDC EXO-injected group
Figure 7. RegDCs EXO co-localize with gingival acceptor DCs, inhibiting maturation in alveolar bone loss model. DIL (Red) labelled EXO were injected into the palatal gingiva of the upper right second molar of inflammatory alveolar bone loss model at days −2, 0 and 2 relative to ligature placement. At day 9 gingival tissues were harvested and dissected for cell isolation. Immunofluorescence labelling of cells was performed using Alexa Flour 488-labelled anti CD11c to identify dendritic cells, counterstained with nuclear stain DAPI and visualized under confocal microscopy. (A) Co-localization of EXO (red) with nucleus (DAPI) and dendritic cells (green). (B) Flow cytometry analysis of MHCII+ cells % in CD11c+DCs gate (top histograms) and CD86+cells % in MHCII+ CD11c+DCs gate (bottom histograms) in gingival cells isolated from six tested groups. (C) Bar graphs of flow cytometry data from (B) showing statistically significant differences in the groups. (D) Representative immunohistochemical staining of CD86+presumptive DCs (arrows) in lamina propria of gingival tissue sections of (left) no EXO-treated +ligature and (right) regDCs EXO-treated +ligature groups. (E) Bar graph of immunohistochemical staining data from (D) showing statistically significant differences in the groups. N=5 in each group; * P<0.05 by one-way ANOVA, followed by Tukeys multiple comparisons.

Regulation of ligature-induced alveolar bone loss by DC EXO subtypes
De-fleshed maxillae from EXO-treated and untreated mice were analysed for bone loss, volumetrically, using microCT. 3-D images of bone volume are shown (Figure 10A). Quantitative analysis confirms 35% bone loss induced by ligature-induced inflammation in the absence of EXO. Injection of regDC EXO reduced alveolar bone loss significantly (p<0.05), while stimDC EXO increased bone loss, with control immune null iDCs EXO and free TGFβ/IL-10 intermediate of the two (Figure 10B). The lack of efficacy of free TGFβ/IL-10 emphasizes the importance of EXO encapsulation of cargo to protect it from proteolytic degradation (Figure 1G) and promote its persistence at diseased site (Figure 6). Histological sections confirm bone changes observed in volumetric microCT data (Figure 10C). These results further substantiate the hypothesis that regDCs EXO, loaded with TGFβ and IL-10 reprogram the immune response and inhibit inflammatory bone loss.

Discussion
The role of Tregs in dampening inflammatory Th17 responses, inhibiting autoimmune disorders, graft
rejection and alveolar bone loss is previously reported [21,23,49-55]. However, there are no published reports of DC-derived EXO being used to reprogram the Th17 mediated immune response to attenuate alveolar bone loss. EXO are naturally released from cells, and endowed with surface adhesion proteins that promote higher interaction and cellular uptake capacity, in comparison to other synthetic agents like liposomes and other nanoparticles which may be toxic [27,29,56,57]. Clinical trials have shown the feasibility and safety of DC-derived EVs administration in melanoma [58], non-small lung cancer [59] and advanced non-small lung cancer [60]. Other therapeutic applications include colon cancer [61] cutaneous ulcers, and mesenchymal stem cell derived EXO for type 1 diabetes mellitus [26]. Despite excitement of EXO therapy for a variety of human diseases, no published study to our knowledge has examined DC EXO as an immunotherapeutic agent for inflammatory alveolar bone loss. Our data suggest that TGFB1- and IL10-loaded DC EXO shield cytokines from proteolytic attack, are retained in periodontal tissues and taken up by DCs and T-cells in gingiva, modulating their phenotype and functions in vivo, resulting in inhibition of alveolar bone loss.

TGFB1 is an immunoregulatory cytokine with differential effects on immune cells, including direct inhibition of effector T-cell activation and proliferation, inhibition of maturation of DCs and macrophages and induction of T-regulatory cells [62,63]. IL-10 is also an immunoregulatory cytokine that inhibits immune cell activation and pro-inflammatory cytokine release that contribute to inflammatory bone loss such as TNF, IL6 and IL17 and RANKL [21]. Moreover, TGFB1 and IL-10 have synergistic regulatory effects [64,65]. Using antibody inhibition studies, we have shown the crucial role of TGFB-1 and IL-10 in the induction of regulatory immune response by regDCs EXO in vitro, with TGFB1 the dominant driving force for Treg differentiation (Figure 5(B, C)) while both cytokines are required for inhibition of DC maturation (Figure 3F). However, the exact mechanism of TGFB-1 interaction with target cells remains to be fully understood.

Our data has shown that TGFB-1 is localized on the outer surface of the EXO as well as inside the EXO as shown by TEM and ELISA (Figure 1(D, E, F). The EXO-associated TGFB-1 was protected from degradation by trypsin and partially protected from proteinase-
Moreover, in TGFB1 inhibited or in exosome TGFB-1 [66].

The interactions between reg DC EXO and acceptor cells are under intensive investigation in our laboratory. Two principal mechanisms, not mutually exclusive, are proposed, involving EXO binding to cell surface receptors, stimulating downstream signalling of SMAD2 phosphorylation, as well as EXO internalization by target cell where it unloads its cargo. A third mechanism could be through the slow release of TGFB-1 from the EXO to the extracellular matrix promoting a paracrine effect on the recipient cell [66–69].

A recent study has shown that neutralizing TGFB-1 on the surface of EXO as well as blocking of TGFBR-1 inhibited SMAD2 signalling [66], alluding to the importance of surface TGFB-1 in the function of EXO . This suggests that an initial interaction of EXO-associated TGFB with TGF-B1 receptor complex is paramount. The same study showed that the EXO facilitates the internalization of EXO-TGFB-1 receptor complex. Within the recipient cell, trafficking of the exosome occurs through endosomal translocation, rather than lysosomal degradation [66,69,70]. A recent study has localized EXO-associated TGFB1 bound to its receptor within endosomal compartments [66] and not in lysosomal compartments, suggesting that EXO protect TGFB-1 from lysosomal degradation. The acidic endosomal compartments would then lead to the release of TGFB-1 bound to its receptor resulting in prolonged sustained SMAD2 signalling [66].

Another suggested mechanism is that after EXO internalization into the recipient cell, TGFB-1 released inside the cell is recycled and shuttled back to the cell membrane where it is secreted to act on the cell surface receptor in an autocrine manner [68]. These results collectively suggest that the mechanism of action of TGFB-1 bearing EXO on the recipient cells requires an initial interaction with the cell surface receptor. However, the internalization of EXO and its endosomal translocation are essential for a sustainable prolonged signalling. Further studies are required to unravel the exact mechanism of communication between EXO and recipient cell.

Unencapsulated TGFB1 and IL-10, injected into gingiva in buffer, did not induce Treg recruitment (Figure 8B, C) or inhibit inflammatory bone loss (Figure 10A, B, C). Free TGFB1 and IL-10 are likely cleared and degraded rapidly in vivo at the disease site. DC EXO prevent proteolytic degradation of EXO encapsulated cytokines (Figure 1G). Moreover, DC EXO persist at the disease site (Figure 6). DC EXO express a variety of adhesion molecules, tetraspanins, integrins and lectins that could promote inflammatory cell binding persistence at the inflamed site [70]. In the present study, three injections of regDC EXO were sufficient to inhibit in situ DC maturation (Figure 7), increasing Tregs and decreasing Th17 effectors locally (Figure 8) and attenuate inflammatory bone loss (Figure 10). Evidence further documents the physical uptake of DC EXO by DCs and T-cells in vitro (Figure 3A and Figure 5A) and in vivo in gingival tissues (Figure 7A and Figure 8A). The mechanisms of EXO uptake by recipient DCs and T cells are critical determinants of the ensuing cellular response [45], and are thus being studied by our group outside the purviews of this work.

In a previous elegant approach to attenuate bone loss, CCL22-loaded synthetic micro-particles were injected to recruit gingival Tregs [21]. Distinct differences in our study included the use of DC-derived nanoparticles [45] to deliver either anti-, or pro-inflammatory cytokine cargo to the site of disease, and assess its potential for predictable disease resolution or exacerbation. Moreover, alveolar bone loss volume in 3-D, was measured by microCT. Our study further traced the anatomical fate of DC EXO in whole live animals, revealing EXO distribution to spleen and other organs when injected by IV route, versus its persistence in the gingiva upon local administration (Figure 6). Our findings are consistent with promising reports of DC EXO therapy for other inflammatory disease models, such as colitis andencephalitis, although adenovirus transfected DC EXO, that secrete TGFB1 were used. While mitigation of Th17 responses were commensurate with clinical improvement [71,72], use of adenovirus raises immunogenicity, toxicity and transformation concerns. The present study used recombinant TGFB1 and IL10, isolated from non-genetically modified DCs to passively and actively load EXO. StimDC EXO were not actively loaded with pro-inflammatory cytokines, but nonetheless enhanced Th17 induction and inflammatory bone loss. The role of Th17 effectors in bone degeneration is well established [13,22,73,74]. Here we show the mechanism involves downregulation of bone resorbing cytokines RANKL and TNF and reduction of TRAP+ osteoclasts in vivo (Figure 9). We also showed that regDC EXO-treated T-cells could inhibit osteoclastogenesis in vitro (Fig. S3), suggesting their bone protective effect [75,76].

DC EXO contain other cargo including mRNA, miRNA and additional proteins that could influence recipient cell functions in vitro and in vivo. We therefore examined distinct miRNA profiles in DC EXO subtypes (Figure 2B). Worthy of emphasis is miR155-5p, downregulated in regDC EXO, upregulated in stimDC EXO. Reported functions of miR155-5p in recipient immune cells include increased induction of pro-inflammatory mediators and decrease in programmed death ligand-1 (PD-L1) [47].
EXO-bound miRNAs are recognized as novel endogenous targets for therapeutic treatments [77]. Determining functions of specific miRNA cargo in these DC EXO, including generation of mimics/anti-mimics is in planning stages.

The DC EXO subtypes used here contain a constellation of other surface bound and encapsulated proteins coding and non-coding RNA that need to be definitively characterized so that we can better understand the immunobiology of DC EXO. Efforts in our laboratory are focused on LC/MS/MS characterization of DC EXO proteins, identifying shared and unique pathways, biological processes and metabolic functions.

In summary, the feasibility and efficacy of DCs EXO loaded with anti-inflammatory cargo as a therapeutic approach to experimental degenerative bone disease was established. The immunologic mechanisms responsible for inhibition (or acceleration) of bone loss in this model was further delineated by using immuno-stimulatory and immune null DC EXO. A more targeted engineering approach, with defined surface proteins and molecular cargo into null DC EXO is in progress.

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