Partial Tendon Injury at the Tendon-to-Bone Enthesis Activates Skeletal Stem Cells

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Abstract

The tendon enthesis plays a critical role in facilitating movement and reducing stress within joints. Partial enthesis injuries heal in a mechanically inferior manner and never achieve healthy tissue function. The cells responsible for tendon-to-bone healing remain incompletely characterized and their origin is unknown. Here, we evaluated the putative role of mouse skeletal stem cells (mSSCs) in the enthesis after partial-injury. We found that mSSCs were present at elevated levels within the enthesis following injury and that these cells downregulated TGFβ signaling pathway elements at both the RNA and protein levels. Exogenous application of TGFβ post-injury led to a reduced mSSC response and impaired healing, whereas treatment with a TGFβ inhibitor (SB43154) resulted in a more robust mSSC response. Collectively, these data suggest that mSSCs may augment tendon-to-bone healing by dampening the effects of TGFβ signaling within the mSSC niche.

Key words: skeletal stem cell; enthesis; tendon-to-bone interface; Achilles injury

Graphical Abstract

mSSCs Downregulate TGFβ Signaling in order to Permit Their Expansion and Contribution to Tendon Enthesis Regeneration
Significance Statement
Tendon regeneration is of critical importance in achieving full recovery among the millions of patients with mechanical musculoskeletal injuries. Once injured, the tendon enthesis can never fully recover to its pre-injury function. The cells involved in the healing process are not well characterized and their origin is unknown. We found that mouse skeletal stem cells (mSSCs) play a fundamental role in the tendon enthesis by augmenting healing and potentially suppressing inhibitors such as TGFβ, which otherwise preclude a full tendon recovery.

Introduction
The tendon enthesis or tendon-to-bone interface (TBI) is a complex structure that plays a critical role in permitting movement and reducing stress within joints. This fibrocartilaginous enthesis is composed of diverse cell types to permit the transfer of mechanical forces between bone and tendon—2 tissues with dramatically different material and cellular properties. The enthesis is classically described as being composed of 4 zones, representing a gradient of mechanical and cellular properties: the tendon (containing organized collagen fibrils and tenocytes with Young’s modulus of 0.2 GPa), uncalcified fibrocartilage, calcified fibrocartilage, and bone (containing organized lamellar structure with Young’s modulus of 20 GPa). Following enthesis-injury, the enthesis is unable to fully regenerate and never regain its uninjured morphological appearance, which increases the risk of recurrent injury. There is extensive histological data available characterizing tendon healing. However, the underlying cellular and molecular mechanisms contributing to tendon-to-bone pathogenesis and healing remain largely unknown.

Our group has previously isolated and identified mouse skeletal stem cells (mSSCs) from a variety of skeletal tissues including long bones, mandibles, and cranial sutures. They are defined by their ability to self-renew and their multilineage contribution to bone, cartilage, and stroma, but not fat. Its downstream progenitor is bone, cartilage, and stromal progenitors (BCSPs). They have both been shown to be key facilitators in bone development, maintenance of homeostasis, and response to fracture healing. In this study, we evaluated whether mSSCs participated in the physiologic response to tendon injury at the enthesis and whether modulation of their activity may improve healing. This study, to our knowledge, represents the first characterization of the mSSC response to enthesis injury and may have broad implications for similar musculoskeletal pathology specifically partial TBI injuries that go untreated (eg, partial rotator cuff tears undergoing physical therapy in humans).

Methods
Subject Details
We purchased 9-week-old Black 6 (C57BL/6J, stock number: 000664) mice from Jackson Laboratories. Rainbow mice (ROSA26VT2/GK3) were provided as a gift from the Weissman Laboratory, Stanford University School of Medicine. Mice were housed at the Stanford University Comparative Medicine Pavilion. The facilities provided light and temperature-regulated housing for all animals. A minimum sample size of 3 replicates per group was used for all experiments (exact numbers for specific experiments are provided in the figure legends). All experiments were carried out in accordance with the Stanford IACUC standards of care and approved by the institutional ethics committee, Administrative Panel on Laboratory Animal Care, (protocol reference: 9999).

Partial Tenotomy Mouse Model
Mice were anesthetized with inhaled isoflurane (Henry Schein Animal Health) at a concentration of 1-2% in oxygen at 3 L/ minute on top of a heating pad to ensure the mice were kept warm throughout the procedure. To prevent corneal desiccation ophthalmic ointment (Puralube petrolatum, Dechra Veterinary Products) was applied to the cornea. The mice were placed in the prone position on a clean operating surface. For the partial tendon injury at the tendon-to-bone enthesis, the left posterior foot, ankle, and lower leg were sterilized with 3 applications of betadine followed by 70% ethanol. Using the aseptic technique, an incision was made over the calcaneus and extended 1 cm proximally. The distal Achilles was dissected free from the surrounding tissues and then just above the tendon-to-bone interface a partial injury (~50%) of the tendon was transected using micro-scissors (Fine Science Tools, UK). The skin was closed using a 6-0 nylon suture (Ethilon). On the contralateral limb, a sham skin incision procedure was performed and was closed using a 6-0 nylon suture (Ethilon). During the procedure, the respiratory rates of the animals were monitored and the isoflurane was titrated accordingly. All mice were monitored daily until sutures were removed. For the mid-substance injury, the same sterile technique was used as well as the same incision. The Achilles was dissected free from the surrounding tissues and then in the mid aspect of the tendon, a partial injury (~50%) was created using micro-scissors (Fine Science Tools, UK). The skin was closed in a similar fashion as the previous procedure.

Liposomal Tamoxifen Induction for Rainbow Reporter System
Activated tamoxifen liposomes (LiTMX) were prepared as described by Ransom et al. Briefly, a 90:10 mol:mol mixture of 1,2-dimyristoyl-sn-glycero-3-phosphocholine and cholesterol (Avanti Polar Lipids) was desiccated under 5 kPa stream of nitrogen gas for 5 min. This was followed by vacuum desiccation at 25 °C for 3 h, sonication in 1x PBS for 15 s, and then reconstitution by extrusion using a 100-nm pore-diameter polycarbonate membrane (Sigma-Aldrich) at 32 °C. Using NanoBrook Omni dynamic light scattering (DLS) instrument (Brookhaven) the liposome size was confirmed. Liposomal vesicles were then incubated with 4-hydroxytamoxifen (Sigma–Aldrich) under nitrogen gas at 25 °C for 6 h. LiTMX was applied locally to injury sites or areas of interest (tendon-to-bone interface) for induction of Cre Recombinase at the time of surgery for Rainbow mice.

Exogenous Delivery of TGFβ
Two micrograms of exogenous carrier-free TGFβ-1 (10 µL, Biolegend, Cat# 763102) were delivered locally (adjacent to the tendon-to-bone interface) at the time of surgery before
skin closure. The procedure was conducted as described above. Additional doses of 2 µg were given every 3 days via a Hamilton syringe for fluorescence-activated cell sorting (FACS) and every other day for gross and histological analysis (700 Series, Hamilton). Achilles enthesis were harvested at 3 and 7 days after surgery for FACS sorting as described below. Saline administration was used for control.

Exogenous Delivery of SB431542
A total of 10 µM SB431542 formulated in DMSO (10 µL, selleckchem, Cat# S1067) was delivered locally (adjacent to the tendon-to-bone interface) at the time of surgery after the skin closure of mice. The procedure was conducted as described above. Additional doses of 10 µM were given every other day at the injury site via a Hamilton syringe for FACS and every other day for gross and histological analysis (700 Series, Hamilton). Achilles TBlS were harvested at 3 and 7 days after surgery for FACS sorting as described below. Saline administration was used for control.

Tissue Processing and Histology
Mouse injured and control tissues were fixed in 4% paraformaldehyde (Electron Microscopy Sciences) for 20 h at 4 °C. The specimens were decalcified in 19% EDTA in PBS at 4 °C for 4 weeks with a change of EDTA every 48 h. Specimens were then placed in 30% sucrose (Sigma–Aldrich) until saturation at 4 °C following fixation, followed by OCT until saturation at 4 °C, embedded in OCT, and sectioned at 8 µm on Superfrost Plus microscope slides (FisherScientific). Representative tissue specimens were stained with a modified pentachrome using the protocol routinely used in our laboratory.

Immunohistochemistry
Immunohistochemistry of TBI cryosections was performed using vectastain elite ABC Kit (Vector Laboratories) according to the manufacturer’s instructions. In brief, samples previously cut in 8 µm sections on Superfrost Plus microscope slides (Fisher Scientific) were rehydrated and endogenous peroxidases were quenched using 3% H2O2 for 10 min. Sections were then treated with heated Tri-base (Sigma–Aldrich) for 15 min, permeabilized with 0.5% Triton-X-100. 1X Power Block (BioGeneX) in combination with heated Tri-base (Sigma–Aldrich) for 15 min, permeabilized with 0.5% Triton-X-100. 1X Power Block (BioGeneX) in combination with avidin/biotin blocking kit (Vector Laboratories) was applied for 1 h at room temperature. Sections were then incubated with the primary antibody (SP7, Abcam, ab22552, dilution 1:250) at 4 °C overnight. Subsequently, the respective secondary antibody (prediluted RTU Biotinylated Goat Anti-Rabbit IgG Antibody, Vector Laboratories, BP-9100-50) was incubated on the specimen for 90 min. The specimens are rinsed and then incubated for 1 minute using DAB (BD Sciences, Cat#: 550880, dilution 1:500). Specimens were next washed with PBS, cover-slipped, and imaged with a Leica DMI6000B inverted microscope system.

Cell Culture and Immunofluorescence
FACS-isolated mSSCs were plated in chambered slides coated with 0.1% gelatin and cultured in MEM alpha medium with 10% fetal bovine growth serum (FBS), 1% penicillin-streptomycin. After at least 48 h cells were washed twice with phosphate-buffered saline (PBS) and fixed with 50% acetone-50% methanol for 20 min at 4 °C, followed by washing with PBS–0.1% Triton-100 twice. Then, cells were incubated in a blocking solution with 1% horse serum in PBS–0.05% Tween-20 for 1 h at RT followed by incubation with a primary rabbit Phospho-SMAD2 (Ser465, Ser467) Antibody (Invitrogen, Cat# 701582, dilution 1:250) for 3 h at room temperature. After, cells were washed 3 times with PBS/0.1% Tween-20 and incubated in the blocking solution for 1 h at room temperature followed by incubation with FITC-conjugated Goat Anti-Rabbit secondary antibody (Abcam, Cat# ab6717, dilution 1:500) for 45 min at room temperature. Nuclear counterstaining was performed using Vectashield H-1200 mounting medium with DAPI (Vector Laboratories).

Confocal Imaging and Analysis
The cryopreserved 8-µm sections on Superfrost Plus microscope slides (Fisher Scientific) as well as the chambered cell culture slides were imaged using laser scanning confocal microscopy via the Leica WLL TCS SP8 Confocal Laser Scanning Microscope (Leica Microsystems) located in the Cell Sciences Imaging Facility (Stanford University, Stanford, CA). The 10x objectives were used (10× HC PL APO, oil, N.A. 0.40). Image analysis was performed using Fiji software (ImageJ, NIH). Photoshop was utilized for image intensity analysis using the histogram tool.

Biomechanical Testing
Within 2 h of sacrifice, the Achilles TBI underwent mechanical testing. During the dissection and mechanical testing, samples were kept moist by regular spraying with 10% PBS. Dimensional measurements of the tendon-to-bone interface were performed using a digital micrometer with fine tips and a resolution of 0.01 mm under microscope dissection by one user. Custom grips were used to compressively grip proximally at the Achilles tendon and distally at the calcaneus. The angle of pull was 180°. As per Beason et al, tendons were preloaded to 0.02 N to remove slack and the length of the tendon was measured. The length of the tendon was designated as the distance between the grip ends (ie, the distance of the entire tendon subjected to the displacement). To provide a consistent strain history, the tendons were exposed to 10 cycles of preconditioning from 0.02 N to 0.05 N at a rate of 0.1%/s. After a 300 s hold, tendons were then loaded to 5% strain at a rate of 25%/s and then held for stress relaxation over 600 s. Finally, tendons were unloaded back to 0% strain and then immediately subjected to an extension test to failure at a rate of 2%/s. Young’s modulus was calculated by taking a least-squares regression of the slope at the individual linear portion of the extensional test. Yield force, stress, and strain were taken at the point of the curve which exited linear deformation and entered plastic deformation. Stress was calculated by dividing the force by the cross-sectional area. The strain was measured by dividing the displacement over the original length of the tendon. Stress, strain, and modulus calculations were made using MATLAB (MathWorks).

Sample Preparation and FACS Isolation
The detailed steps listed in Gulati et al were followed for sample preparations and FACS isolation mSSCs. Specimens, including either the TBI or mid-tendon, were dissected and then were gently minced with a dissecting scissor on ice. They were then serially digested in collagenase digestion buffer at 37 °C for 30 minutes under constant agitation in an orbital
shaker at 275 rpm at 37 °C. Digest consisted of serum-free Medium 199/EBs (HyClone) supplemented with 1.6 mg/mL Collagenase Type II (Sigma-Aldrich, Cat# 6885) 2 mg/mL Dispase II (Roche Diagnostics), 1% bovine serum albumin (Sigma-Aldrich), 1% pluronic F-68 (Gibco-Life Technologies), 2% HEPES buffer (Gibco-Life Technologies, Grand Island, NY, USA), 0.4% 2.5 M CaCl2 (Sigma-Aldrich), 100 units/mL deoxyribonuclease I (Worthington Biochemistry). Dissociated cells were filtered through a 70 μm mesh filter, pelleted at 300g at 4 °C, resuspended in FACS buffer which was composed of PBS (Gibco-Life Technologies), 2% fetal bovine serum (Gibco-Life Technologies), 1% penicillin-streptomycin (Gibco-Life Technologies), 1% pluronic F-68 (Gibco-Life Technologies). The digestions were pooled together and total dissociated cells were pelleted at 300g at 4 °C, resuspended in FACS buffer.

Cells were stained with fluorochrome-conjugated antibodies CD45 (Biolegend, Cat# 103110), Ter119 (ThermoFisher, Cat# 15-5921-81, dilution 1:200), CD202/TIE2 (eScience, Cat# 14-5987-81, dilution 1:20), Thy1.1 (ThermoFisher Cat# 47-0900-82, dilution 1:100), Thy 1.2 (ThermoFisher Cat# 47-0902-82, dilution 1:100), CD105/6C3 (ThermoFisher, Cat# 13-1051-85, dilution 1:100), CD51 (BD Sciences, Cat# 551187, dilution 1:50), CD 200 (BD Sciences, Cat# 745255, dilution 1:50), goat anti-rat IgG secondary antibody (ThermoFisher, Cat# Q-11601MP, dilution 1:50), and Alexa Fluor 680 Fluorophore Antibody (ThermoFisher, Cat# A20188, dilution 1:200). Stained cells were pelleted at 300g at 4 °C, resuspended in 300 μL of FACS buffer 70 μm filter for flow cytometry. For information regarding antibodies used for FACS isolation of skeletal stem/progenitor cells refer to Supplementary Table S1.

Flow cytometry was performed on the FACS Aria II in the Lorye Lokey Stem Cell Institute Shared FACS Facility. First, CD45+ and dead cells (Pi+) were fractionated out and the remaining population (P3) was fractionated based upon previously described antigens by Chan et al13 and Gulati et al14. The mSSC profile was CD45+, Ter119−, Thy1.1−, Thy1.2−, 6C3−, CD105−, CD51+, and CD200−. The mBCSP profile was CD45+, Ter119−, Thy1.1−, Thy1.2−, 6C3−, CD51−, and CD105+. The sequential FACS gating strategy is shown in Supplementary Fig. S2. Cells were sorted directly into TRIzol Reagent (Ambion-Life Technologies) for RNA isolation and extraction or alpha-MEM supplemented with 20% fetal bovine serum, and 1% penicillin-streptomycin (Gibco-Life Technologies) for cell culture and differentiation assays.

Bulk mRNA Sequencing

mSSCs, freshly isolated from 3 groups of 5 pooled injured or control TBLs of C57BL/6J mice were sorted directly into TRIzol Reagent (Ambion-Life Technologies). All cells collected were used for RNA extraction. The average number of cells collected for each pooled group was 2140 ± 3487 cells. RNA extraction was performed using Qiagen miRNaseasy kit with one column DNAse treatment per the manufacturer’s recommendations. To generate cDNA from 1 ng total RNA, the Clontech Smarter Ultra Low Input RNA kit (Takara Bio) was used per the manufacturer’s recommendations. Amplified cDNA was purified using SPRI Ampure Beads (Beckman Coulter) and the quality and quantity were measured using a high sensitivity DNA chip on the Agilent 2100 Bioanalyzer (Agilent Technologies). The cDNA was enzymatically fragmented to an average of 300 base pairs and libraries were generated with the Nextra-Xt kit (Illumina). The samples were uniquely barcoded, pooled, and sequenced on a single lane of the HiSeq (Illumina). A total of 263 million paired-end reads were obtained, with a minimum threshold of 37.5 million reads per sample.

Bulk mRNA Sequencing Data Analysis

A total of 6 samples (3 injured and 3 control samples containing pooled 5 specimens each) were profiled by bulk RNA sequencing at post-injury day 7 as described above. Raw FASTQ reads were aligned to GENCODE vM20 reference transcripts (GRCh38.p6) for a mouse with Salmon v0.12.0 using the -seqBias, -gcBias, -posBias, -useVBOpt, -rangeFactorizationBins 4, and -validateMappings flags and otherwise default parameters for single-end mapping. Count normalization and differential gene expression analysis were performed using the DESeq2 v1.22.2 package in R. Counts were size-factor normalized using the “DESeq” function and log2-transformed. Pairwise differential gene expression analysis was performed using the lfcShrink function with “type = apeglm,” which applies an adaptive t prior shrinkage estimator. Principal component analysis (PCA) was used to evaluate aggregate separation between injury and control (sham-surgery) transcriptomes. Gene enrichment analysis was performed on ranked lists of differentially expressed genes (n = 500 genes), ordered by adjusted P-value, using the enrichR toolkit against the KEGG, WikiPathways, and Gene Ontology databases.26,27 Additional analysis was performed using subsets of these genes that were specifically upregulated (n = 247) and downregulated (n = 253).

RT-PCR Analysis for Genes Expression

To analyze the expression level of genes of interest, RNAs were isolated from cells using the Trizol (Ambion-Life Technologies, Carlsbad, CA) procedure according to the manufacturers’ instructions. Isolated RNAs were submitted to RT-PCR procedure as previously described.24 Primer-sequences for Col1a, PAI1, and Gapdh genes, and annealing temperatures ions are reported in Supplementary Table S2.

Statistical Analysis

Statistical analyses were performed using the software GraphPad Prism v.6. Results are expressed as absolute numbers, percentages, or means ± SD. Data analysis was performed using an unpaired t-test assuming 2-tailed distribution or one-way analysis of variance (ANOVA) and post hoc Tukey correction were used to compare groups where relevant. Fisher’s exact test was used for gene ontology analysis.26,27 Statistical significance was assigned for P ≤ .05.

Results

Mouse Achilles TBI Injury Model

The cellular mechanisms responsible for TBI healing after injury remain poorly characterized. We aimed to evaluate whether new tissue formation within the injury site involved tissue-resident skeletal stem cells. We used a mouse partial Achilles TBI tenotomy model known to alter the mechanical loading of the enthesis, resulting in impaired healing that never fully recapitulates normal morphology (Fig. 1A; Supplementary Fig. S1A). Using this model, we confirmed that the biomechanical properties of the TBI were significantly altered. The linear region of the stress–strain curve was significantly reduced at
post-injury day 7 (control 97.4 ± 30.2 MPa vs. post-injury day 7 30.4 ± 22.8 MPa; \( P = .005 \)); the elastic modulus was also significantly reduced 7 days after injury (control 610.1 ± 479.2 MPa vs. post-injury day 7 128.2 MPa ± 140.9 \( P = .02 \); Fig. 1B-1D; Supplementary Fig. S1B, S1C).

Polyclonal Expansion After Partial Achilles TBI-Injury

We began by assessing the extent of the clonal proliferation of local cells in Achilles TBI after injury. We created a partial Achilles TBI tenotomy to mimic tendon TBI injury in rainbow mice (Actin\textsubscript{cre}\textsuperscript{ERT2}; Rosa\textsubscript{R26\textscript{S}}), given that no existing genetic drivers are known to delineate mSSC clonal proliferation. These rainbow mice express a Cre-inducible fluorescent reporter under the ubiquitous Actin promoter following tamoxifen induction. After recombination, cells express one of 4 colors (eGFP, mCerulean, mCherry, and mOrange). The resulting progeny are marked with the same color as the parent cell.\textsuperscript{29,30} To achieve tissue-specific rainbow induction, we applied liposomes at the region of interest immediately following injury.\textsuperscript{23} Local administration of 4-hydroxytamoxifen liposomes instead of systemic tamoxifen injection was performed to ensure that

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**Figure 1.** Tissue and cell clonality characterization of partial Achilles tenotomy model. (A) Illustration and gross imaging of partial Achilles tenotomy model. (B) Elastic modulus of the Achilles TBI is significantly decreased 7 days post-injury (\( n = 6 \) per group). (C) The linear region of the stress–strain curve of an injured Achilles TBI is significantly less than the sham control at 7 days (\( n = 6 \) per group). (D) Representative stress–strain curve (\( n = 6 \) per group). (E-H) Gross imaging, pentachrome staining, confocal micrographs, and SP7 immunohistochemistry of transverse TBI sections from Actin\textsubscript{cre}\textsuperscript{ERT2}; Rosa\textsubscript{R26\textscript{S}} mice which were locally induced with activated tamoxifen liposomes using a published protocol at the time of surgery.\textsuperscript{23} Harvested at 3, 7, and 14 days post-surgery, clonal proliferation were visualized within the calcaneus at day 3, and by day 7 there was clonal proliferation within the calcaneus and along with the injury at the enthesis (noted by an arrow). (I) Quantification of the average number of clones at the injury site after injury (POD3-14, \( n = 3 \) per group). Representative samples calcaneus and tendon are labeled respectively as “C” and “T” an arrow marks the injury (\( n = 3 \) per group). Scale bars, 100 \( \mu \)m. Data and error bars are shown as mean ± SD. \(* P < .05, \) unpaired 2-tailed \( t \)-test.
only tissue-resident cells would be labeled. This would avoid labeling any cells outside of the local environment.

Histologically, we observed both gradual TBI repair and bone remodeling of the calcaneus (Fig. 1E–1H, Supplementary Fig. 1D, 1F). On confocal microscopy, we observed an increased number of single-colored clones at the TBI after injury (Fig. 1E–1I). On immunohistochemistry, we noted increased expression of the SP7 protein, a known regulator of osteoblast differentiation and associated with bone formation, at post-injury day 7 compared to sham control (Fig. 1E–1H). Taken together, these findings suggest that cells of skeletal origin proliferate clonally in response to TBI-injury.

mSSCs Are Present Within the TBI at Homeostasis and Proliferate in Response to Injury

Since we had previously observed that mSSCs are enriched in mouse skeletal tissues after injury, we evaluated TBI cells using a similar strategy with FACS (Fig. 2A, Supplementary Fig. S2). We collected tissues from sham control Achilles TBI and tendon mid-substance (mid-tendon). We identified the presence of mSSCs and their downstream bone-cartilage-stromal progenitors (mBCSPs) within both tissues and found a significantly larger representation of mSSCs and mBCSPs within the TBI compared to the mid-substance (Fig. 2B).

Subsequently, we sought to determine how mSSCs respond to TBI and/or mid-substance injury. We performed partial Achilles tenotomies at either site, utilizing the contralateral leg as a sham surgery control. This experiment uncovered a significant mSSCs expansion 7 days post-injury in response to TBI-injury, but not after mid-substance injury (Fig. 2C, 2D). Similarly, TBI injury elicited an expansion of mBCSPs, suggesting that both progenitor populations in this tissue are injury-responsive.

TBI-Injury-Specific mSSC Gene Regulation

To characterize gene expression changes associated with mSSC in response to injury, we FACS-isolated injured and sham control mSSCs for bulk RNAseq analysis. Clear differences were observed between injury and control transcriptomes (Fig. 3A). Using enrichment analysis, we evaluated the top 500 genes with the greatest difference in expression between injury vs. sham mSSCs (Fig. 3B). Genes upregulated in injured mSSCs suggest these cells were locally activated. Specifically, *Pannx3*, associated with cellular activation of differentiation pathways including osteogenesis and chondrogenesis, was significantly upregulated in injured mSSCs. Additional pathways upregulated in injured mSSCs suggesting that the mSSCs are actively playing a role in TBI healing were the Indian hedgehog (Ihh) pathway linked to TBI fibrocartilage development and mineralization, glycosaminoglycan biosynthesis (data not shown, see the Gene Expression Omnibus for bulk RNA data), known to be an important part of the extracellular matrix in managing TBI compressive forces, and in cell cycling suggesting that mSSCs are being activated to replicate and possibly differentiate (Fig. 3C, 3D; Supplementary Fig. S3A).

Interestingly, we found an overall downregulation of TGFβ pathway-components, indicated by downregulation of *Tgfbi1, Tgfbi2, Smad3*, and upregulation of *Fbn1*, known to control the bioavailability of TGFβ (Fig. 3C). Gene set enrichment analysis also highlighted the downregulation of TGFβ receptor binding pathways in injured mSSCs (Fig. 3D). BMP and FGF signaling have been shown to facilitate differentiation into enthesis components; however, there were no significant changes in the expression of genes associated with these 2 signaling pathways in mSSCs following injury, suggesting that these mechanisms may not play a significant role in mSSC activation during TBI healing.

To confirm our RNAseq findings that TGFβ signaling is downregulated in injured mSSCs, we performed immunofluorescence of injured and control mSSCs investigating the expression of pSMAD2. SMAD is a major receptor-activated downstream effector of TGFβ signaling. There was a decrease in the intensity of expression and in the number of cells that had positive pSMAD2 nuclear staining thus, suggesting that there may be the downregulation of TGFβ signaling in mSSCs after injury (Fig. 3E, 3F). Further confirmation of the differential activation of TGFβ signaling between uninjured and injured mSSCs was obtained from the PCR expression analysis of 2 specific targets of activated TGFβ signaling, *Col1a1*, and *PaI1*. These 2 genes were found to be upregulated in sham control mSSCs as compared to injured mSSCs (Fig. 3G). The above findings strongly indicate that there is a difference in TGFβ signaling between injured and uninjured mSSCs.

**Artificially Elevated Levels of Active TGFβ-1 After Injury Reduce the mSSC Response and Worsen TBI Healing**

Over-activation of TGFβ signaling in the bone after partial mouse Achilles tendon injury is known to contribute to enthesisopathy, whereas inhibition of TGFβ via an exogenous TGFβ-neutralizing antibody improves tendon-to-bone healing. The response to TGFβ signaling relies on precise spatial and temporal activation; if altered, as seen in TBI healing, the alignment of enthesis fibrocartilage cells and proteoglycan synthesis is disturbed. Given the downregulation of gene expression of several components of the TGFβ signaling pathway within the mSSC niche after partial-injury, we sought to investigate the effects of modulating TGFβ signaling on the mSSC response. We applied exogenous TGFβ-1 to TBI sites and found that TGFβ-1 treatment led to a significant inhibition of the normal mSSC and mBCSP expansion at 7 days post-injury (Fig. 4A). Histologically, there was increased tissue disorganization and enthesisopathy-like morphology seen in TGFβ1 treated TBIs (Supplementary Fig. S4). Grossly, we saw delays in healing with a profound inflammatory and fibrotic response in injuries treated with exogenous TGFβ-1 (Fig. 4B). These results further support prior research that the presence of excess TGFβ after TBI injury results in poor TBI healing and the possible inhibition of mSSC expansion (Fig. 4C).

We then examined whether functional modulation of TGFβ may affect TBI healing and the mSSC response using the small-molecule inhibitor SB431542. SB431542 is a selective inhibitor of the TGFβ superfamily type I activin receptor-like kinase receptors, which have been explored in a variety of fibrotic pathologies. It has been previously shown in a mouse model of massive rotator cuff tear results in decreased fibrosis, fatty infiltration, and muscle weight loss after injury. When this inhibitor was applied at the location of the injury, we found that there was a significant expansion of mSSCs and mBCSPs in both wild-type mice (Fig. 4A). Grossly, in mice treated with the inhibitor we saw a dramatic decrease in both fibrotic and inflammatory responses, compared to mice treated with exogenous TGFβ (Fig. 4B). Unexpectedly, the administration of SB431542 also improved the biomechanical
properties of the injured TBI compared to treatment with PBS, as a control as well as treatment with TGFβ (Fig. 4D, 4E; Supplementary Fig. S4B). This outcome may reflect that TGFβ has a negative impact on the tissues within the TBI.

Discussion

The inability of the enthesis to fully recapitulate its mechanical and morphological properties after a partial injury leaves it at high risk for future rupture. In this study, we identified mSSC as a cellular component of the TBI at homeostasis and observed their activation, in concert with their likely downregulation of TGFβ signaling, following TBI-injury. Furthermore, we found that mSSC representation was dramatically decreased in mice treated with exogenous TGFβ-1 immediately following injury.

TGFβ plays an important role in cartilage and bone homeostasis, the development of tendons and bony eminences/interfaces, and the regulation of embryonic progenitor specification. It is secreted in an inactive, latent form and deposited in the extracellular matrix. Several physiological processes result in the conversion of inactive TGFβ protein to its active form. However, excessive active TGFβ uncouples bone remodeling and contributes to the pathogenesis of osteopenia, Camurati-Engelmann disease, and other skeletal diseases. High concentrations of active TGFβ-1 in the TBI have been shown to cause enthesopathy-like-states and the use of TGFβ-1 inhibitors after tendon injury has been shown to reduce fibrosis. Our transcriptomic profiling identified a downregulation in the expression of several TGFβ signaling components in mSCs isolated from injured-TBI in wild-type mice. In addition, our transcriptomic data suggests TGFβ signaling downregulation is seen within mSCs the population present after TBI injury.

Previous studies have found that TGF-β1 localizes to forming scar tissue after TBI injury and increases the formation of fibrous tissues at the healing site by increasing collagen accumulation without the expansion of tenogenic
TGF-β1 may initially improve an injured TBI’s overall strength due to scarring compared to untreated injuries, but it does not restore the pre-injury TBI morphology and overall biomechanical properties. As a comparison, fetal wound healing, where no scars are seen, is characterized by low expression of TGF-β1, while adult wound healing is characterized by high levels of TGF-β1 and typically excessive scar formation. The administration of TGF-β1 to mSSCs in culture does not result in the expansion of mSSCs, but rather altered the morphology of the cells. Figure 3. Transcriptomic profile demonstrates downregulation of TGFβ signaling components in mSSCs isolated from injured TBI. (A) Principal component analysis plot shows clustering of mSSCs FACs sorted specimens from injured and control tissues (n = 5 pooled, with 3 replicates) at post-injury day 7. (B) Heatmap of bulk RNA-seq data showing differential peaks (log2(normalized reads in peaks)) of injured mSSCs and control mSSCs gene expression from post-injury day 7. Gene enrichment is noted in color key and histogram at the upper left. (C) Heatmap of selected genes of interest from overall heatmap differential peaks with significant upregulation of the Indian hedgehog pathway and downregulation of the TGFβ pathway. GO Terms Molecular Function Significant in injured mSSC. GO terms enriched for genes upregulated (red) and downregulated (blue) in injured versus control mSSCs at post-injury day 7. Identified by bulk RNA-seq. Select GO terms with significant P-values (Fischer exact test) are shown. (E) Injured mSSCs showed decreased image intensity of the nuclear localization of pSMAD2 (green) compared to control mSSCs (P < .001). Immunofluorescence of pSMAD2 (top column), DAPI staining (middle column), and merged images are shown. Original magnification: 20×. The image is representative of 3 experiments. F ) Significant decrease in quantified mSSC pSMAD2 immunofluorescence image intensity after injury which was performed with ImageJ and photoshop software. G, RT-PCR assay confirms Col1a and Pal1 were downregulated in injured mSSCs compared to sham control (n = 15 pooled mice).
of colony; while administration of exogenous BMP-2 resulted in the rapid expansion of mSSCs and has also been shown to have a positive impact on tendon healing, suggesting the need for future investigation. Interestingly, TGF-β1 has been shown to possibly facilitate tenogenic differentiation of bone marrow-derived mesenchymal stem cells, suggesting that TGF-β1 may have different effects on mSSCs versus bone marrow-derived mesenchymal stem cells.

Pathologically high concentrations of active TGFβ can impact the recruitment and proliferation of mesenchymal stem cells as well as a subset of bone marrow precursor-cells, defined by Nestin positivity, contributing to the pathogenesis of enthesopathy. This Nestin+ lineage, with known endothelial and mesenchymal lineage-cell specification, is unlikely to promote TBI-regeneration. Given that endogenous activation of TGFβ signaling increases Nestin+ cells in tendons, we sought to investigate whether administration of exogenous TGFβ would impact the prevalence of mSSCs post-injury. We observed a reduced mSSC response with post-injury exogenous TGFβ treatment in wild-type. Furthermore, the mSSC response was augmented beyond the control in mice treated with SB431542, a TGFβ inhibitor. These data indicate that, contrary to previous reports on Nestin+ lineage cells, after TBI injury mSSCs may orchestrate downregulation of TGFβ signaling to permit their expansion and eventual contribution to TBI regeneration. This is demonstrated by the fact that mSCCs expand after acute injury and downregulate TGFβ signaling, while exogenous administration of TGFβ attenuates the mSSC response/activation after TBI injury.

Increased TGFβ-1 is associated not only with enthesopathy and poor TBI healing but also with pathologic bone formation
states, including heterotopic ossification and osteoarthritis. In contrast, inhibiting TGFβ signaling attenuates the progression of such disease processes and promotes the endogenous regenerative potential of the mammalian calvarium after injury. This elevated signaling alters enthesis organ-dogenous regenerative potential of the mammalian calvarium progression of such disease processes and promotes the en-

It is important to note that our findings are specific to acute transection of a normal tendon that was not immobilized. Future research is required to evaluate the role of mSSCs in the setting of degenerative tendon/TBI injury that typically occurs in humans. Further, given that we used only Black 6 and rainbow mice strains in the study, there may be species- or strain-specific differences that limit the scope of the findings. Regarding our statistical methods, we ran the Shapiro–Wilk test on our data and found that for the immunofluorescence where there are 20 replicates, it met the assumptions for parametric testing via the Shapiro–Wilk’s test, however, the sample sizes for our other results were too small (<5) to test for normality. Additionally, the study only covers the early phase of TBI repair as it uses early time points, and as a result future studies with longer time points are needed to investigate the long-term role of mSSCs in TBI healing and tissue remodeling.

Conclusion
Collectively, our results frame a conceptual advance in the understanding of tendon-to-bone regeneration by demonstrating, the involvement of a skeletal stem cell population in tendon-to-bone healing. Furthermore, these findings suggest that mSSCs play a role in repair and regeneration through the downregulation of TGFβ signaling. These mSSC and TGFβ pathway interactions may represent potential targets for medical interventions aimed at mitigating the painful and persistent enthesisopathy that develops after partial enthesis injury.

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Conflicts of Interest
The authors declared no potential conflicts of interest.

Author Contributions
A.L.T.: conception and design, financial support, administrative support, collection of data, data analysis, and interpretation, manuscript writing. M.D.: collection of data, data analysis, and manuscript writing. D.F.: study design, data collection, analysis and interpretation, and final approval of the manuscript. A.S., S.M., K.C.: data collection, analysis and interpretation, and final approval of the manuscript. M.L.: data collection, analysis, and final approval of the manuscript. E.F.: data collection and final approval of the manuscript. R.E.J., P.F.: study design, data interpretation, and manuscript writing. I.B.: data interpretation and manuscript writing. A.B.: data analysis, final approval of the manuscript. M.J.: data analysis and interpretation, manuscript writing. G.G.: data interpretation and final approval of the manuscript. C.C.: study design, data interpretation, and writing of the manuscript. N.Q.: conception and design, collection of data, data analysis and interpretation, and manuscript writing. M.T.L.: conception and design, financial support, administrative support, data interpretation, and manuscript writing.

Data Availability
Data to support the conclusions drawn in this manuscript can be found in the primary figures. All RNA-seq data can be accessed from the Gene Expression Omnibus (GEO, accession number GSE167080, https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE167080).

Supplementary Material
Supplementary material is available at Stem Cells Translational Medicine online.

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