**Osteoarthritis and Cartilage**

Muscle cell-derived factors inhibit inflammatory stimuli-induced damage in hMSC-derived chondrocytes

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**SUMMARY**

**Objective:** Pro-inflammatory cytokines play an important role in inducing cartilage degradation during osteoarthritis pathogenesis. Muscle is a tissue that lies near cartilage in situ. However, muscle’s non-loading biochemical effect on cartilage has been largely unexplored. Here, we tested the hypothesis that muscle cells can regulate the response to pro-inflammatory cytokine-mediated damage in chondrocytes derived from human bone marrow-derived mesenchymal stem cells (hMSCs).

**Method:** hMSCs were allowed to undergo chondrogenic differentiation in porous silk scaffolds in the typical chondrogenic medium for 12 days. For the next 9 days, the cells were cultured in chondrogenic medium containing 50% conditioned medium derived from C2C12 muscle cells or fibroblast control cells, and were subject to treatments of pro-inflammatory cytokines IL-1β or TNFα.

**Results:** Both IL-1β and TNFα-induced strong expression of multiple MMPs and hypertrophic markers Runx2 and type X collagen. Strikingly, culturing hMSC-derived chondrocytes in C2C12 muscle cell-conditioned medium strongly inhibited the expression of all these genes, a result further confirmed by GAG content and histological evaluation of matrix protein. To determine whether these effects were due to altered chondrocyte growth and survival, we assayed the expression of cell proliferation marker Ki67, cell cycle arrest markers p21 and p53, and apoptosis marker caspase 3. Muscle cell-conditioned medium promoted proliferation and inhibited apoptosis, thereby suggesting a possible decrease in the cellular aging and death that typically accompanies cartilage inflammation.

**Conclusion:** Our findings suggest the role of muscle in cartilage homeostasis and provide insight into designing strategies for promoting resistance to pro-inflammatory cytokines in hMSC-derived chondrocytes.

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**Introduction**

Articular cartilage serves a load-bearing role in diarthrodial joints1,2. However, degenerative erosion of this tissue can lead to osteoarthritis (OA) which is estimated to affect 71 million North Americans by the year 20303. Due to the narrow reparative potential of cartilage, along with its avascular and aneural nature4, current treatments are mainly restricted to alleviating symptoms and pain5. Cartilage tissue engineering has emerged as an attractive therapeutic option for restoring and regenerating cartilage in situ.

Such strategies rely on the combination of repair cells with the appropriate three-dimensional (3D) biomaterial scaffolding and biochemical cues necessary to create a regenerative microenvironment. Engineered constructs can then be implanted at the site of injury to facilitate healing and restoration of tissue function.

For adequate healing and regeneration, engineered cartilage requires appropriate coupling of progenitor repair cells with biocompatible scaffolding that facilitates the generation of a tissue construct similar to native cartilage. With respect to progenitor repair cells, matrix-secreting primary chondrocytes are limited in quantity and capacity for expansion, and are susceptible to de-differentiation *in vitro*4,5. In comparison, human bone marrow-derived mesenchymal stem cells (hMSCs) have emerged as an attractive cell source for cartilage tissue engineering4,5. These multipotent cells can be obtained in relatively large amounts, rapidly expanded *ex vivo*, and...
differentiated to adopt the chondrogenic cell fate. In terms of 3D scaffolding, multiple materials have been explored for engineering cartilage tissue. Among them, silk, derived from the *Bombyx mori* (silkworm), stands out as a biomaterial whose biocompatibility, slow degradation, and robust mechanical properties make it suitable for cartilage tissue engineering.

The hostile pathological environment in the arthritic joint imposes a major challenge on cartilage tissue engineering. For successful cartilage repair, the engineered tissue construct must also be able to maintain its stability in the inflammatory microenvironment in vivo during OA pathogenesis. In the joint, pro-inflammatory cytokines, most notably IL-1β and TNF-α, cause cartilage damage in OA by stimulating the expression of a number of proteolytic enzymes, including matrix metalloproteinases (MMPs) and aggrecanases. Additionally, to further drive the degradation cascade, IL-1β and TNF-α also promote chondrocyte hypertrophy, cellular senescence, and cell death, thereby reducing the proliferation and viability potential for chondrocytes.

Muscle lies in close proximity to cartilage during development and throughout life. This tissue provides cartilage with a biomechanical stimulation that promotes nutrient distribution and maintains homeostasis. For example, age-associated muscle loss is distance to the progression of OA in the elderly and reduced muscle strength has been shown to be a risk factor for knee OA. As a result, muscle strengthening has been explored as an intervention to prevent or delay the onset of OA. Furthermore, muscle paralysed, which leads to muscle atrophy, induces joint abnormality and cartilage degradation. On the other hand, bone morphogenetic protein 2 (BMP-2)-expressing muscle tissue was found to serve as a bridge in enhancing and accelerating femur fracture healing in a rat model. While these studies have demonstrated the significance of muscle tissue on skeletal development, it is unclear whether this regulation involves merely biomechanical stimuli, biochemical mediators, or both.

Recently, our group investigated the non-loading biochemical effect of muscle cells on cartilage gene expression. Our results showed that a rat chondrocyte cell line co-cultured with muscle cells or cultured in muscle cell-conditioned media as monolayers had significantly higher expression levels of cartilage matrix proteins in the presence of IL-1β and TNF-α. However, for the purpose of tissue engineering, it is critical to evaluate the effect of muscle cell-derived factors on primary cells in a 3D microenvironment under such inflammatory stimuli.

In this study, we subjected hMSCs-derived differentiating chondrocytes grown in 3D silk scaffolds to C2C12 muscle cell-conditioned media as well as pro-inflammatory cytokines IL-1β and TNF-α. Administration of IL-1β and TNF-α strongly induced the gene expression of matrix degrading enzymes (MMPs) and hypertrophic markers such as Runx2 and type X collagen (Col X) in hMSC-derived chondrocytes. In contrast, culturing in conditioned medium derived from C2C12 muscle cells resulted in strong inhibition of the expression of these genes, a result further confirmed by quantification of proteoglycan content and histological assessment. To determine whether muscle cell-derived factors altered chondrocyte growth and survival, we assayed the expression of cell proliferation and cell cycle arrest markers as well as apoptosis markers. We found that culturing in muscle cell-conditioned medium promoted chondrocyte proliferation and inhibited cell death under inflammatory stimuli. Together, our findings demonstrate that muscle cell-derived factors, or myokines, provide enhanced resistance to pro-inflammatory cytokine-mediated cartilage degradation, hypertrophy, and cell growth arrest in hMSC-derived chondrocytes and suggest that such factors may be utilized to improve the stability of bioengineered cartilage in the hostile microenvironment of arthritic joints.

### Methods

#### Biomaterial scaffolds

Silk fibroin scaffolds were fabricated as previously described. Briefly, *B. mori* cocoons were boiled in 0.02 M Na2CO3 to extract sericin proteins. Silk fibroin solution was then lyophilized and resuspended in 1,1,3,3,3-hexafluoro-2-propanol (HFIP). NaCl particles (size: 106–212 μm) were added to 12% silk-HFIP solution (w/v) to generate porous scaffolds with a pore size of 106–212 μm. Scaffolds were then punched into 5 mm × 3 mm (diameter × height) disks for our experiments. Scaffolds of the same dimensions, made of 50/50 polyactic acid (PLA) and bovine type I and type III collagen, were obtained from BD Biosciences (Franklin Lakes, NJ).

#### Cell culture

hMSCs were purchased from Lonza Walkersville, Inc (Walkersville, MD). The lot numbers purchased were 1-OFG3285 (donor #1), 8F3520 (donor #2), and 3F4287 (donor #3). Cells were expanded in Dulbecco’s Modified Eagle Medium (DMEM) (Gibco, Grand Island, NY) containing 10% fetal bovine serum (FBS) (Hyclone, Rockford, IL), 1% antibiotic-antimycotic (Gibco), 0.1 mM non-essential amino acids (Gibco), and 1 ng/ml basic fibroblast growth factor (Peprotech, Rocky Hill, NJ). For chondrogenic differentiation, cells were cultured DMEM containing 1% antibiotic-antimycotic, ITS + supplement (BD Biosciences), 0.1 mM ascorbic acid 2-phosphate (Sigma Aldrich, St. Louis MO), 1.25 mg/ml human serum albumin (Sigma), 10-7 M dexamethasone (Sigma), and 10 ng/ml transforming growth factor beta 3 (TGF-β3; R&D Systems).

Murine myoblast cell line (C2C12) and murine fibroblast cell line (NIH3T3) were purchased from ATCC (Rockville, MD). Cell lines were cultured in DMEM with 10% FBS and 1% antibiotic-
antimycotic. Cells were maintained at a 75–80% confluence for 24 h. Conditioned medium was sterile-filtered using a 0.22-μm syringe filter for subsequent use.

#### hMSC seeding and differentiation

Prior to cell seeding, scaffolds were immersed in sterile DMEM. 0.5 × 10^5 hMSCs in 15 μl of chondrogenic media were seeded into scaffolds and allowed to attach for 2 h. hMSC-seeded scaffolds were then transferred into 1 ml of fresh media in 24-well tissue culture plates at 37°C under static culturing conditions for 2 days before being placed on an incubated dynamic shaker (5–6 RPM) for remainder of study.

For the study of hMSC differentiation on different scaffolds, hMSCs (passage 4) were seeded into PLA, collagen, or silk scaffolds, and maintained in chondrogenic media for 21 days. For hMSC differentiation in the presence of pro-inflammatory cytokines, hMSCs (passage 5) were differentiated in silk scaffolds for 12 days using chondrogenic media. At this time, scaffolds were divided into six groups [Fig. 1(A)] and cultured for an additional 9 days in: 1:1 chondrogenic media and C2C12 conditioned media, 5 ng/ml IL-1β in a 1:1 mixture of chondrogenic media and C2C12 conditioned media, and 5 ng/ml TNF-α in a 1:1 mixture of chondrogenic media and C2C12 conditioned media. A control group of scaffolds were cultured with NIH3T3, rather than C2C12, conditioned media. In all 21-day studies, media was changed every 3 days. Unless stated otherwise, data shown represents a sample size of n ≥ 3 independent samples (cell-seeded scaffolds) from one experiment using one donor.
RT-PCR analysis

Total RNA was isolated from scaffolds using the RNeasy Mini kit (Qiagen; Valencia, CA) as per the manufacturer's protocol. RNA was then transcribed to cDNA using random primers (Invitrogen), M-MLV reverse transcriptase kit (Invitrogen), and dNTPs (New England BioLabs; Ipswich, MA) and stored at −20°C until RT-PCR analyses. For RT-PCR, cDNA was mixed with gene-specific primers (sequences

Fig. 1. Study design and mRNA expression of chondrogenic genes. hMSCs were differentiated for 21 days into chondrocytes using chondrogenic media, with treatment of pro-inflammatory cytokines and muscle cell- or non-muscle control cell-conditioned media after Day 12 (A). Relative expression of markers for cartilage matrix production at Day 12 included transcription factor SOX9 (B), collagen II (Col II) (C), and ACAN (D), as compared to undifferentiated hMSCs at Day 0. Matrix markers at Day 21 included collagen II (E) and ACAN (F), while hypertrophic markers included Runx2 (G) and collagen X (Col X) (H). Expression was normalized to GAPDH. Data are presented as mean with 95% confidence intervals (n = 3 independent samples (cell-seeded scaffolds) from one experiment completed using one donor's cells).
available upon request) and SYBR Green Supermix (Quanta Bio- 
sciences, Inc., Gaithersburg, MD), quantified using BioRad’s iQ5 Real 
Time PCR Detection System and optical software (Hercules, CA), and 
normalized against glyceraldehyde-3-phosphate-dehydrogenase 
(GAPDH). For all analyses, at least three independent samples (cell- 
seeded scaffolds) from one experiment were analyzed \( (n \geq 3) \).

**Histology**

Constructs were fixed in formalin, dehydrated in graded ethanol 
solutions, cleared with xylene, embedded in paraffin, and sectioned 
into 5- or 10-μm thick sections. After deparaffinization, sections 
were stained with hematoxylin and 1% eosin (H&E) and toluidine 
blue. Diaminobenzidine (DAB) immunocytochemistry was carried 
out using the DAB staining kit (Vector Laboratories; Burlingame 
CA). Primary antibodies for type II collagen (Abcam; Cambridge, 
MA), caspase 3 (Abcam), and Ki67 (Vector Laboratories) were used. 
When appropriate, sections were counterstained with methyl 
green, prior to imaging and counting.

**Sulfated glycosaminoglycans (GAG) and DNA content quantification**

Constructs were cut into pieces and digested in papain solution 
(10 U/mg in PBS containing 5 mM cysteine-HCl and 5 mM EDTA) at 
60°C overnight, sonicated, and then stored at ~20°C. Sulfated 
glycosaminoglycan (GAG) content was quantified based on the 
absorbance at 525 nm using dimethylmethylene blue (DMMB) dye 
and a shark chondroitin sulfate standard\(^{32}\). DNA content was 
quantified using Picogreen (Invitrogen). At least three independent 
samples were analyzed for each condition.

**Statistical analysis**

Each experiment was prepared from a single donor’s cells. Data 
was calculated from at least three independent samples (cell-
seeded scaffolds) from one experiment using one donor’s cells and 
is reported as mean with 95% confidence intervals. Data was further 
analyzed using Prism 4 (Graphpad; La Jolla, CA) and a student’s 
t-test or one-way analysis of variance (ANOVA) with Tukey post hoc 
assessment. Statistical significance was assigned where appropriate 
with \( P < 0.05 \).

**Results**

**Muscle cell-derived factors inhibit hypertrophy in hMSC-derived 
chondrocytes under inflammatory stimuli**

To assess the effects of muscle cell-derived factors on the 
response of hMSC-derived chondrocytes to inflammatory cyto-
kines, hMSCs were seeded onto silk scaffolds and allowed to un-
dergo chondrogenic differentiation for 12 days using established 
chondrogenic medium\(^ {11}\). We chose silk scaffolds for hMSC seeding 
based on our comparison study on the chondrogenic potentials of 
hMSCs seeded into silk, collagen, or PLA scaffolds, where we found 
that hMSCs in silk scaffolds produced the highest content of 
GAG and DNA, as well as higher levels of aggrecan (ACAN) mRNA 
and lower levels of mRNA for caspase 3 (CASP3) \( (\text{Fig. S1}) \). For 
studying hMSC differentiation in the presence of pro-inflammatory 
cytokines and muscle-derived factors, we first assessed the 
expression of cartilage-specific genes at Day 12 using RT-PCR and 
compared to undifferentiated cells at Day 0. While chondrogenesis 
is still ongoing at Day 12, a level of chondrogenic differentiation has 
been achieved prior to the addition of cytokines and conditioned 
media. This finding is signified by the induction of SOX 9, a tran-
scription factor that acts as a master regulator in chondrogenic 
differentiation, as well as cartilage matrix genes type II collagen 
(Col II) and ACAN \( (\text{Fig. 1(B–D)}) \).

Over the next 9 days, hMSC-derived chondrocytes were subject to 
IL-1β or TNFα treatment. In the presence of pro-inflammatory 
cytokines, the expression of cartilage matrix genes Col II and 
ACAN was significantly reduced while expression of hypertrophic 
cartilage markers Col X and Runx2 was induced, as compared with 
controls \( (\text{Fig. 1(E–H)}) \). It was also noted that TNFα had a stronger 
effect in inducing chondrocyte hypertrophy than IL-1β \( (\text{Fig. 1(G–H)}) \). Culturing with C2C12 muscle cell-conditioned medium did not 
rescue IL-1β- and TNFα-induced Col II and ACAN reduction 
\( (\text{Fig. 1(E–F)}) \). Strikingly however, culturing with muscle cell-
conditioned medium led to a significant decrease in the expres-
sion of hypertrophic markers Col X and Runx2, suggesting that 
muscle cell-derived factors inhibited pro-inflammatory cytokine-
induced chondrocyte hypertrophy \( (\text{Fig. 1(G–H)}) \).

**Muscle cell-derived factors strongly inhibit the induction of multiple 
MMPs by inflammatory stimuli in hMSC-derived chondrocytes**

Matrix metalloproteinases MMP1, MMP8, and MMP13 are 
involved in the proteolytic degradation of cartilage matrix protein 
Col II, while MMP3 activates MMP1 as well as degrades matrix 
proteoglycans\(^ {33}\). Both IL-1β and TNFα induced a significant 
and consistent upregulation of MMP1 \( (\text{Fig. 2(A)}) \), MMP3 \( (\text{Fig. 2(B)}) \), 
MMP8 \( (\text{Fig. 2(C)}) \), and MMP13 \( (\text{Fig. 2(D)}) \) in hMSC-derived 
chondrocytes cultured in the control condition. Strikingly, culturing in 
C2C12 muscle cell-conditioned medium essentially abolished the 
induction of all these MMPs. Even in the absence of pro-
inflammatory cytokines, muscle cell-conditioned medium signifi-
cantly reduced expression of MMP3 and MMP13.

**Muscle cell-derived factors promote GAG production and Col II 
protein expression in the presence of inflammatory stimuli**

H&E histological analysis showed that the cells were distributed 
uniformly throughout scaffolds \( (\text{Fig. 3(A–F)}) \). To determine the 
levels of cartilage matrix deposition in the cartilage constructs, we 
performed toluidine blue staining \( (\text{Fig. 3(G–L)}) \), as well as immu-
nochemistry staining for Col II \( (\text{Fig. 3(M–R)}) \). While both 
toluidine blue staining and Col II staining were diminished with the 
addition of pro-inflammatory cytokines in cells grown in control 
conditioned media, staining remained consistent for all cells 
cultured with muscle cell-conditioned media, regardless of in-
flammatory cytokine treatment.

To further assess whether muscle cell-derived factors promoted 
cartilage matrix production, we quantified the amount of sulfated 
GAG within these constructs using the DMMB assay\(^ {32}\). While there 
were no observable differences in GAG \( (\text{Fig. 4(A)}) \) and DNA 
\( (\text{Fig. 4(B)}) \) content among the samples in the absence of cytokine 
treatment, constructs cultured in the presence of muscle cell-
derived factors yielded significantly higher levels of GAG and 
DNA content, as well as GAG/DNA ratio \( (\text{Fig. 4(C)}) \) under IL-1β, but 
not TNFα treatment as compared to the control. This result suggests 
that under the inflammatory condition of IL-1β, muscle cell-derived 
factors promoted the maintenance of cartilage matrix.

**Muscle cell-derived factors promote cell proliferation and inhibit 
apoptosis under IL-1β treatment**

Because we observed that cells cultured in muscle cell-
conditioned medium showed increased DNA content under IL-1β 
treatments, we next investigated whether this increase was due to 
increased cell proliferation or decreased cell death. To examine cell 
proliferation status, we used immunocytochemistry to quantify the
number of cells that were stained positive for Ki67, a marker for cell proliferation [Fig. 5(A–B)]. Muscle cell-conditioned medium did not enhance cell proliferation under non-inflammatory conditions, which is consistent with the quantification of total DNA level. IL-1β treatment completely abolished Ki67 staining, suggesting a strong inhibition of cell proliferation. Remarkably, under IL-1β treatment, muscle cell-derived factors induced a stark increase in the percentage of Ki67-positive cells. Unlike IL-1β, TNFα did not cause a significant reduction in cell proliferation as compared to the control condition. Furthermore, no changes in cell proliferation were observed for TNFα treatment in the presence of muscle cell-derived factors.

To assay if the change of cell proliferation was caused by altered expression of cell cycle regulators, we then assessed the mRNA expression of cell cycle inhibitors p21 [Fig. 5(C)] and p53 [Fig. 5(D)]. Culturing with muscle cell-conditioned medium did not significantly alter the expression of these two markers under the non-inflammatory conditions. On the other hand, both IL-1β and TNFα induced the expression of p21, while TNFα also significantly induced p53 expression. Significantly, culturing with muscle cell-derived factors led to a marked decrease in the expression of these inhibitors, indicating a possible mechanism for the observed changes in cell proliferation.

Fig. 2. Relative expression of matrix degradation markers in differentiated hMSCs at Day 21 after treatment of pro-inflammatory cytokines and muscle cell-derived factors as compared with controls. Matrix degradation markers included MMP1 (A), MMP3 (B), MMP8 (C), and MMP13 (D). Expression was normalized to GAPDH. Data are presented as mean with 95% confidence intervals (n = 3 independent samples (cell-seeded scaffolds) from one experiment completed using one donor’s cells).

Fig. 3. Representative histological evaluation of hMSC-derived chondrocytes within silk scaffolds after Day 21 of culture in the presence of pro-inflammatory cytokines and muscle cell-derived factors as compared to controls. H&E staining (A–F). Toluidine blue staining for proteoglycans (G–L) and immunocytochemical staining for type II collagen (M–R). (Scale bars: 250 μm).
Data are presented as mean with 95% confidence intervals (n = 6 independent samples (cell-seeded scaffolds) using one donor’s cells).

**Discussion**

Catabolic mediators, in particular pro-inflammatory cytokines IL-1β and TNFα, have been demonstrated to inhibit chondrogenesis and promote chondrocyte hypertrophy and degradation in OA joints, thus leading to cartilage instability. This is especially important for immature engineered tissues composed of progenitor stem cells actively undergoing chondrogenesis as such constructs may not have generated enough protective extracellular matrix (ECM) at the time of construct implantation and consequently, may be especially more susceptible to degradation.

Indeed, the therapeutic viability of engineered cartilage relies on methods that promote resistance to these catabolic inducers. Here, we addressed this challenge by investigating the effects of muscle cell-derived factors on the stability of cartilage constructs under inflammatory conditions.

Our findings demonstrated that muscle cell-derived factors inhibited the mRNA expression of MMPs as well as hypertrophic markers in bioengineered cartilage. Pro-inflammatory cytokines IL-1β and TNFα cause cartilage destruction by inducing the expression of MMPs, promoting chondrocyte hypertrophy, and reducing the synthesis of cartilage matrix genes. Although the induction of hypertrophic markers by IL-1β and TNFα were about 2–4 fold, we believe this is biologically meaningful as OA chondrocytes are known to exhibit higher levels of hypertrophy. For all the MMPs we examined (MMP1, MMP3, MMP8, and MMP13), muscle cell-derived factors abolished IL-1β- and TNFα-induced expression at the mRNA level, indicating a strong impairment of the cartilage-destructive ability of these pro-inflammatory cytokines. Furthermore, we found a significant inhibition of hypertrophic markers Col X and Runx2 in constructs cultured with muscle cell-conditioned medium, suggesting that muscle cell-derived factors inhibited the terminal differentiation of hMSCs, which is one of the major obstacles for the translational application of hMSC chondrogenesis. These results were consistent among all three donors tested (Figs. 1 and 2 for donor #1; Fig. S2 for donors #2 and #3). Although there were no differences in the mRNA expression of Col II and ACAN in hMSC-derived chondrocytes cultured with muscle cell-conditioned medium, the dramatic reduction of MMPs and the inhibition of chondrocyte hypertrophy could result in the maintenance of protein expression of cartilage matrix components under inflammatory conditions.

Thus, to substantiate our RT-PCR gene expression analyses, we performed biochemical and histological studies to quantitatively and qualitatively evaluate the expression of cartilage matrix proteins. Our immunocytochemistry and toluidine blue staining results showed that cartilage matrix proteins, mainly Col II and total proteoglycan, produced by constructs exposed to muscle cell-conditioned medium were maintained as compared to non-muscle cell-conditioned medium under IL-1β or TNFα stimulation. Quantification of sulfated GAG content revealed only statistically significant differences in GAG levels between muscle-treated samples and controls in the case of IL-1β stimulation. For TNFα treatments, it is not clear why there are no quantifiable significant differences in GAG content as compared to the proteoglycan histological staining. Such difference in results may be likely due to the fact that while the DMMB dye, which was used for GAG quantification, only binds to sulfated proteoglycans, toluidine blue stains both sulfated and non-sulfated proteoglycans, such as hyaluronan. Further investigation of individual proteoglycan components may help to understand whether muscle cell-derived factors promote additional cartilage matrix components in maintaining the stability of hMSC-derived chondrocytes under inflammatory conditions. It should be noted that TNFα led to a higher level of cartilage destruction as evidenced by its stronger induction of chondrocyte hypertrophic formation.
markers Col X and Runx2, as well as all MMPs, as compared to IL-1β. Consequently, to overcome the inflammatory damage induced by TNFα, a larger amount of muscle cell-derived factors may be required. Finally, these results are consistent with our prior study using rat chondrosarcoma (RCS) chondrocytes in 2D cultures, which showed that muscle cell-secreted factors promoted cartilage matrix protein expression under inflammatory cytokine treatments\textsuperscript{30,31}.

To further explore the mechanisms by which muscle cell-derived factors regulate chondrocyte stability under inflammatory conditions, we analyzed cell proliferation and cell death. Muscle cell-derived factors strongly rescued IL-1β-induced growth arrest, which

\begin{figure}
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\includegraphics[width=\textwidth]{fig5.png}
\caption{Biochemical quantification and relative gene expression analysis of cell proliferation, cell cycle regulation, and apoptosis. Cells were stained via immunocytochemistry for proliferative marker Ki67 (A) and the percentage of Ki67 + stained cells were quantified (B). mRNA expression for markers for cell cycle regulation and inhibition included p21 (C) and p53 (D). Cells were stained via immunocytochemistry for active caspase 3 (CASP3) (E) And the percentage of positively stained cells were quantified (F). mRNA expression for CASP3 was also assessed (G). All mRNA expression was normalized to GAPDH. For immunocytochemistry, sections were counterstained with methyl green. Data are presented as mean with 95% confidence intervals (n = 4 sections for quantification of Ki67 + cells, n = 3 independent samples (cell-seeded scaffolds) for RT-PCR, and n = 6 sections for quantification of caspase 3 + cells).}
\end{figure}
may be mediated in part by downregulating the expression of cell cycle inhibitor p21. Although p53 mRNA expression was not induced by IL-1β or inhibited by muscle cell-conditioned medium, it is not clear whether the level of p53 activation was altered in these conditions. Interestingly, TNFα did not change the proliferation level in these cells as assayed by Ki67 staining; however, it did significantly induce p21 and p53 expression. It has been established that these two transcription factors, in addition to their function as cell cycle inhibitors, also play a significant role in cellular aging and senescence, and it will be interesting to determine the effect of muscle cell-derived factors on these pro-inflammatory cytokine-induced events.

We also observed that muscle cell-derived factors inhibited IL-1β-induced CASP3 mRNA expression but did not inhibit TNFα-induced expression. As noted previously, higher levels of muscle cell-conditioned medium might be required to overcome TNFα-induced CASP3 expression, as TNFα typically is a stronger inducer of apoptosis. Furthermore, it is worth noting that while cytokine-induced increases in CASP3 mRNA expression were observed, there was no similar trend observed in cytokine-induced protein expression of active CASP3, a cleaved product of pro-CASP3. Here, it is not clear whether the enzymatic activity of CASP3 was altered by IL-1β and TNFα. What our data clearly shows is that muscle cell-conditioned medium significantly inhibited the protein expression of activated CASP3 with respect to IL-1β treatment. These findings parallel the observed increase in DNA content by muscle cell-derived factors during IL-1β treatment and further propose the role of muscle in promoting cell survival under the condition of IL-1β treatment.

Several future studies can stem from the results presented here. It is important to determine the effective component(s) in the muscle cell-conditioned medium that are contributing to the observed prevention of matrix degradation. Recently, muscle has been recognized as an endocrine organ and is found to secrete multiple myokines. Although muscle lies outside the joint capsule, it is in the position to influence cartilage gene expression locally and systemically, as these myokines can potentially pass through this capsule. In fact, it was reported that synovial fluid assayed by Ki67 staining; however, it did significantly induce p21 and p53 expression. It has been established that these two transcription factors, in addition to their function as cell cycle inhibitors, also play a significant role in cellular aging and senescence, and it will be interesting to determine the effect of muscle cell-derived factors on these pro-inflammatory cytokine-induced events.

In summary, our findings reveal an important role of muscle in regulating cartilage homeostasis and offer a mechanism to promote stability and resistance in engineered cartilage to pro-inflammatory cytokines. We hope this study will lead to a deeper understanding of the control of cartilage development, which may eventually be harnessed to improve tissue engineering strategies.

Author contributions

Conception and design: Rainbow, Zeng.
Acquisition of data: Rainbow, Kwon, Foote.
Provision of study materials: Preda, Kaplan.
Analysis and interpretation of data: Rainbow, Zeng.
Drafting of article: all authors.
Final approval of article: all authors.

Conflict of interest

None of the authors have any conflict of interest related to this work.

Role of the funding source

The NSF, NIH, and NIGMS had no role in the design, analysis or interpretation of the data or in the writing of the manuscript.

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Supplementary data

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