Anisotropic mineralized collagen scaffolds accelerate osteogenic response in a glycosaminoglycan-dependent fashion†

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Regeneration of critically-sized craniofacial bone defects requires a template to promote cell activity and bone remodeling. However, induced regeneration becomes more challenging with increasing defect size. Methods of repair using allografts and autografts have inconsistent results, attributed to age-related regenerative capabilities of bone. We are developing a mineralized collagen scaffold to promote craniofacial bone regeneration as an alternative to repair. Here, we hypothesize modifying the pore anisotropy and glycosaminoglycan content of the scaffold will improve cell migration, viability, and subsequent bone formation. Using anisotropic and isotropic scaffold variants, we test the role of pore orientation on human mesenchymal stem cell (MSC) activity. We subsequently explore the role of glycosaminoglycan content, notably chondroitin-6-sulfate, chondroitin-4-sulfate, and heparin sulfate on mineralization. We find that while short term MSC migration and activity was not affected by pore orientation, increased bone mineral synthesis was observed in anisotropic scaffolds. Further, while scaffold glycosaminoglycan content did not impact cell viability, heparin sulfate and chondroitin-6-sulfate containing variants increased mineral formation at the late stage of in vitro culture, respectively. Overall, these findings show scaffold microstructural and proteoglycan modifications represent a powerful tool to improve MSC osteogenic activity.

1. Introduction

Osseous defects of the skull occur secondary to trauma, congenital abnormalities, or after resection to treat stroke, cerebral aneurysms, or cancer. 1-4 While bone has the capacity to regenerate over time, the large scale of craniofacial (CMF) defects typically preclude effective regeneration. Thus, solutions to regenerate bone in these defects involve the use of autografts, allografts, and other biomaterial implants. Autografts and allografts are the typical standard for repair, using bone from a patient’s own body or from a donor. These methods of repair have variable success rates, due to age-related regenerative property changes in bone grafts. 5-9 Other biomaterial strategies seek to overcome these limitations, and one such material of interest is mineralized collagen. Mineralized collagen scaffolds are composed of the natural components of bone (type I collagen, calcium and phosphate ions, glycosaminoglycans), are highly porous, and have been shown to promote mineral formation in vitro without osteogenic supplements and produce bone in vivo. 10-19 The scale of critical sized defects is a challenge to these and other biomaterial approaches, which demand designs to improve cell invasion, proliferation, and mineral biosynthesis activity. Current clinical strategies often use large exogenous doses of growth factors (e.g., bone morphogenic protein 2, BMP-2), 18 that have significant complications such as ectopic bone formation, resorption of adjacent bone, and maxillary growth disorders. 20-24 These drawbacks highlight the need for a new paradigm.

We are developing a material-only biomaterial approach to actively promote, rather than passively permit, MSC recruitment, osteogenic differentiation, and eventually bone regeneration. Pore size and architecture remain an important factor in guiding tissue regeneration. Pore sizes in the range of 100–350 μm have been shown to promote the formation of bone. 25 Further, the native anisotropy of trabecular bone is increasingly being explored as a design paradigm. 26 Recently, Klaumünzer et al. 26 suggested anisotropic pores in a collagen biomaterial can better promote intramembranous ossification required in CMF bone repair. 26 We have previously developed an approach to fabricate non-mineralized collagen scaffolds with anisotropic pores to increase the metabolic activity of tenocytes and provide directional growth of these cells for tendon repair. 27-31 Further,
glycosaminoglycans (GAGs) in the extracellular matrix play an important role in bone homeostasis and potentially regeneration.\textsuperscript{24} GAGs linked to bone repair include chondroitin-6-sulfate (CS6), chondroitin-4-sulfate (CS4), and heparin sulfate (Heparin). CS6 has been shown to be the main GAG present in articular cartilage, while CS4 is hypothesized to be more important to calcification\textsuperscript{23} and collagen-CS4 biomaterials can promote in vitro bone mineralization.\textsuperscript{24} Alternatively, non-covalent interactions with sulfated GAGs such as heparin sulfate can enhance retention and activity of osteogenic growth factors such as BMP-2 to promote osteogenic activity.\textsuperscript{25–28} We know the GAG chemistry of non-mineralized collagen scaffolds can be exploited to enhance growth factor sequestration and resultant cell activity.\textsuperscript{29} The mineralized collagen scaffolds under development in our lab possess isotropic pores and contain CS6 glycosaminoglycans. In this work we seek to modify the pore anisotropy and GAG content of these scaffolds to increase MSC invasion and osteogenic activity. We hypothesize: (1) anisotropic pores within the mineralized scaffolds will increase cell invasion and osteogenic activity; and, (2) substituting heparin sulfate or chondroitin-4-sulfate for chondroitin-6-sulfate in these scaffolds will increase bone formation. We report the porosity and pore orientation of a family of anisotropic and isotropic scaffolds, examine MSC invasion into the scaffold using a modified transwell membrane assay, and evaluate human MSC osteogenic activity and mineral biosynthesis over 28 days of in vitro culture.

2. Materials and methods

2.1. Experimental design

The aim of this study was to determine mineral formation and cellular activity changes of human mesenchymal stem cells (hMSCs) seeded on mineralized collagen scaffolds as a function of pore orientation and glycosaminoglycan content. Firstly, we compared cell invasion into, subsequent bioactivity, and mineral formation as a function of pore anisotropy for a family of mineralized collagen scaffolds with a conventional glycosaminoglycan content (chondroitin-6-sulfate). Secondly, we compared cell activity and mineral formation of mineralized collagen scaffolds with anisotropic pores as a function of glycosaminoglycan content (chondroitin-6-sulfate, chondroitin-4-sulfate, heparin sulfate). All comparisons involved testing cell viability, gene expression, cytokine release, and mineral formation via alkaline phosphatase, microcomputed tomography (Micro-CT), and Inductively Coupled Plasma (ICP) assays (Fig. 1).

2.2. Fabrication of isotropic and anisotropic mineralized collagen scaffolds

Mineralized collagen scaffolds were fabricated using previously described procedures.\textsuperscript{10,13,14,27,46–48} In a cooled, jacketed vessel, 1.9 w/v% type I bovine collagen (Sigma-Aldrich, Missouri, USA), 0.84 w/v% chondroitin-6-sulfate (Sigma-Aldrich), and Ca(OH)\textsubscript{2}, H\textsubscript{3}PO\textsubscript{4}, and Ca(NO\textsubscript{3})\textsubscript{2}·4H\textsubscript{2}O were thoroughly homogenized, making sure to prevent collagen clumping. The mineralized collagen suspension was transferred to 7.62 × 7.62 cm aluminum molds to create isotropic scaffolds. Alternatively, the suspension was pipetted into Teflon molds with a copper base to create anisotropic scaffolds.\textsuperscript{49} The suspensions were cooled at a constant rate of 1 °C min\textsuperscript{−1} from 20 °C to −10 °C. Once frozen, the suspension was held at this temperature for 2 hours then lyophilized (0.2 torr pressure, 0 °C) in a VirTis Genesis 25XL Lyophillizer (SP Industries, Inc., Pennsylvania, USA).

Anisotropic mineralized collagen scaffolds with disparate GAG content were fabricated following the same method. However initial mineralized collagen suspensions were created using one of three potential glycosaminoglycans: chondroitin-6-sulfate (chondroitin sulfate sodium salt from shark cartilage, Sigma-Aldrich), chondroitin-4-sulfate (sodium chondroitin sulfate A, Toronto Research Chemicals Inc., Ontario, Canada), or heparin sulfate (Heparin). CS6 has been shown to be the main GAG present in cartilage mucosa, Sigma-Aldrich).

2.3. SEM imaging and pore size analysis of mineralized collagen scaffolds

A scanning electron microscope (SEM) was used to visualize pore orientation and mineral crystals in isotropic, anisotropic, and various glycosaminoglycan-containing mineralized collagen scaffolds. Scaffolds were cut in half to expose the interior, and then sputter coated with Au/Pd prior to visualizing with a FEI Quanta FEG 450 ESEM (FEI, Oregon, USA). The pore size of isotropic and anisotropic mineralized collagen scaffolds was analyzed following a JB-4 (Polysciences, Inc., Pennsylvania, USA) embedding procedure.\textsuperscript{36–37} After hydrating scaffolds in ethanol and subsequently JB-4, the scaffolds were placed into wells with three samples of each type placed flat into the mold and three placed on their side to create transverse and longitudinal sections. Polymerization was completed overnight at 4 °C. Embedded scaffolds were further embedded in paraffin to fit molds for microtome sectioning, and 5 μm sections were cut using a RM2255 microtome (Leica, Wetzlar, Germany) with a tungsten carbide blade. Sectioned scaffolds were placed onto glass slides and stained with an aniline blue solution (Thermo Fisher Scientific, Massachusetts, USA) to stain collagen fibers. Slides were then imaged with a NanoZoomer Digital Pathology System (Hamamatsu, Japan) and pore structure was analyzed with a custom Matlab pore size code to acquire an average pore size and pore aspect ratio for each scaffold.\textsuperscript{30,41}

2.4. Sterilization, hydration, and crosslinking of mineralized collagen scaffolds

All scaffolds were sterilized using a dehydrothermal crosslinking and sterilization procedure.\textsuperscript{44} Scaffolds were placed under vacuum in a vacuum oven set at 105 °C for 24 h to complete crosslinking and sterilization.\textsuperscript{42} All subsequent methods followed sterile techniques. Scaffolds were hydrated and further crosslinked before cell seeding. Briefly, scaffolds were soaked in 100% ethanol, followed by multiple washes in PBS, and then crosslinked with EDC–NHS carbodiimide chemistry. Scaffolds were then washed in Phosphate Buffered
Saline (PBS) and soaked in normal growth media for 48 hours before cell seeding.

2.5. hMSC cell culture and scaffold seeding

Passage 4 human mesenchymal stem cells (Lonza, Switzerland) were expanded to passage 5 in an incubator set at 37 °C and 5% CO₂ and fed with low glucose DMEM and glutamine, 10% mesenchymal stem cell fetal bovine serum (Gemini, California, USA), and 1% antibiotic-antimycotic (Gibco, Massachusetts, USA). Before seeding cells onto scaffolds, mycoplasma contamination was tested with a MycoAlert™ Mycoplasma Detection Kit (Lonza); all experiments used cells that tested negative for mycoplasma. hMSCs were seeded on hydrated scaffolds at a density of 150,000 cells per scaffold, with 7500 cells per mL on the bottom and top of each scaffold using a previously defined static seeding method.27,48 Seeded scaffolds were placed in an incubator for an hour and a half to allow cell attachment before basal media was added without osteogenic supplements. Media was replaced every third day and the study proceeded for 28 days.

2.6. Metabolic activity quantification of cell-seeded scaffolds

An alamarBlue® assay was used to quantify the metabolic activity of all scaffold variants.27 A standard curve was generated with known cell numbers to correlate fluorescent readings with fold change of metabolic activity. In reported data, a metabolic activity of 1 represents the metabolic activity of the initial scaffold cell seeding density (150,000 cells). Scaffolds were washed in PBS at every timepoint (day 1, 4, 7, 14, 28) and placed in a solution of alamarBlue® (Invitrogen, California, USA) in an incubator at 37 °C on a shaker. After incubation for 1.5 hours, the alamarBlue® solution was collected and fluorescence of resorufin (540(52) nm excitation, 580(20) nm emission) was measured with a F200 spectrophotometer (Tecan, Switzerland) \( (n = 6) \).

2.7. Gene expression quantification of cell-seeded scaffolds

Osteogenic (RUNX2, COL1A2, Osterix, BMP2), chondrogenic (SOX9), and adipogenic (PPARγ) gene expression was monitored over a 28 day in vitro experiment.38 RNA was isolated from cell-seeded scaffolds using a RNaseq Plant Mini Kit (Qiagen, California, USA). RNA was reverse transcribed to cDNA using a QuantiTect Reverse Transcription kit (Qiagen) and a S100 thermal cycler (Bio-Rad, California, USA). 10 ng of cDNA was

| Gene       | Primer information          |
|------------|-----------------------------|
| GAPDH      | Hs02786624_g1               |
| COL1A2     | Hs00264051_m1               |
| Osterix    | Hs01866874_s1               |
| BMP2       | Hs00154192_m1               |
| PPARγ      | Hs01115513_m1               |
| SOX9       | Hs00165814_m1               |
| RUNX2      | Hs01047973_m1               |

Table 1 Primers used for RT-PCR

Fig. 1 Experimental Outline. Hypothesis 1: anisotropic pores will have greater mineral formation and cell activity than isotropic pores. The different molds used during freeze drying to fabricate isotropic and anisotropic mineralized collagen scaffolds is demonstrated. Hypothesis 2: chondroitin-4-sulfate and heparin sulfate will have greater mineral formation and cell activity than scaffolds containing chondroitin-6-sulfate. Anisotropic scaffolds containing the same amount of chondroitin-6-sulfate, chondroitin-4-sulfate, or heparin sulfate are compared.
used in each well of a real-time PCR reaction, using Taqman primers and a Taqman fast advanced master mix (Applied Biosystems, California, USA). Primers can be found listed in Table 1. PCR plates were read using a QuantstudioTM 7 Flex Real-Time PCR System (Thermo Fisher Scientific). Results \((n = 5)\) were analyzed using a delta-delta CT method and expressed as fold change normalized to expression levels prior to seeding on scaffolds. \(GAPDH\) was used as a housekeeping gene.

### 2.8. Mineral formation analysis via micro-CT and ICP optical emission spectrometry

Final mineral formation in cell-seeded scaffolds was measured by micro-CT and ICP. For micro-CT analysis, scaffolds were fixed in formalin and then stored in PBS at 4 °C prior to imaging. A MicroXCT-400 (Zeiss, Germany) was used to scan samples \((n = 3)\) using a 1× magnitude camera and a power set to 8 W and 40 kV, with the same X-ray source and detector distance, exposure time, and binning for each scaffold. ImageJ was used to evaluate the mineral content from z-stacks of 2D images, and a threshold intensity of 214/255 was set for every scaffold. The particle average intensity was then calculated in ImageJ, following a previously described procedure. To calculate average fill of the scaffolds, the average total area of the particles was divided by the area of the scaffold. Scaffold average fill at day 28 was compared to unseeded scaffolds at day 0.

After micro-CT analysis, scaffolds were analyzed by ICP optical emission spectrometry to define the calcium and phosphorous content. Samples \((n = 3)\) were soaked in 100% ethanol dried with an Autosamdir 931 critical point dryer (Tousimis, Maryland, USA). 0.5 mL to 1 mL of liquid samples and ~5 mg of solid samples were used for elemental analysis. After carefully weighing/pipetting the samples, they were transferred to digestion tubes, and were further treated with concentrated trace metal grade nitric acid (67–70%) purchased from Fisher Scientific. This was followed by automated sequential microwave digestion in a CEM Mars 6 microwave digester (CEM Microwave Technology Ltd., North Carolina, USA). The final digested product was a transparent, aqueous solution which was further diluted to a volume of 50 mL using DI water, so as to make the final concentration of the acid <5%. Matrix matched standards were prepared for the calibration curves. The solution was then introduced to inductively coupled plasma-mass spectrometer (ICP-OES, Optima 8300, PerkinElmer, USA) for the elemental analysis in axial mode. Digestion and ICP-MS analysis parameters are listed in ESI Tables 1 and 2.†

### 2.9. Alkaline phosphatase activity

A colorimetric Alkaline Phosphatase Assay Kit (Abcam, United Kingdom) was used to measure changes in cell-dependent active bone formation in hMSC-seeded scaffolds as a function of pore orientation and glycosaminoglycan content over 28 days \(in vitro\). Day 0 analysis was performed on 2 unseeded scaffolds of each type; subsequent (day 7, 28) analyses were performed on 2 unseeded and 6 MSC-seeded scaffolds of each type. Scaffolds were washed with PBS, collected in tubes, frozen in liquid nitrogen and stored in a −80 °C freezer. Scaffolds were homogenized and centrifuged and 70 μL of sample per scaffold was used in conjunction with assay buffer and \(p\)-nitrophosphophosphate (pNPP) solution for the incubation; after the incubation period, absorbance was measured with a Tecan Infinite M200 Pro (Tecan) at 405 nm. Absorbance values for unseeded day 7 and day 28 scaffolds were subtracted from values for seeded scaffolds of the respective groups, and the normalized values were divided by group-averaged values from day 0 to give a fold change for each scaffold type \((n = 6)\).

### 2.10. Cell migration assay using transwell membranes

Cell migration in anisotropic and isotropic scaffolds containing chondroitin-6-sulfate was evaluated \(via\) a modified transwell membrane invasion assay (ESI Fig. 1†). Briefly, hMSCs were seeded on the top surface of 8 μm pore size adjustable transwell membranes (ThermoFisher Scientific). Scaffolds were added to the bottom of the plate with 600 μL of normal growth media containing 50 ng mL⁻¹ of PDGF-BB (ThermoFisher) to induce chemotactic migration. Transwells were adjusted to rest on the top of the scaffolds and 50 000 hMSCs were seeded onto the transwell membrane in 100 μL of normal growth media. Scaffolds were allowed to sit for 24 h, and after 24 h scaffolds were removed and an alamarBlue® assay was used to quantify the number of cells that had migrated into each scaffold type following the procedure outlined in 2.6 and a standard curve using 50 000 cells.

| Table 2 | Pore size and aspect ratio analysis of chondroitin-6-sulfate isotropic and anisotropic scaffolds cut in the longitudinal and transverse direction. There was no significant difference \((p < 0.05)\) in pore size, however, the pore aspect ratio of the anisotropic scaffolds cut transversely was significantly \((p < 0.05)\) different than the isotropic scaffolds cut longitudinally |
| --- | --- | --- |
| Pore alignment | Orientation | Average pore aspect ratio | Average pore diameter (μm) |
| Isotropic | Transverse | 0.948 ± 0.004 | 177 ± 19.3 |
| | Longitudinal | 0.955 ± 0.003 | 169 ± 5.2 |
| Anisotropic | Transverse | 0.887 ± 0.034 | 170 ± 25.7 |
| | Longitudinal | 0.938 ± 0.007 | 140 ± 9.5 |
2.11. Cytokine release

A RayBio® C-Series Human Cytokine Antibody Array C5 (Ray Biotech, Georgia, USA) was used to measure the relative levels of 80 cytokines eluded from hMSC-seeded scaffolds as a function of pore orientation and glycosaminoglycan content. MSC-seeded scaffolds were incubated for 7 days and media was collected and replaced on the 3rd, 6th, and 7th days. Following collection, media was pooled for analysis. Cell media not exposed to scaffolds was used as a control. The procedure for the assay was performed in triplicate with 1 mL of undiluted sample for each group. Membranes were imaged with an ImageQuant LAS 4010 (GE Healthcare Life Sciences, Massachusetts, USA) for 1 second. Resulting images were analyzed with ImageJ to measure intensity levels of each cytokine; once measurements were obtained one sample was arbitrarily chosen for positive control normalization. Results were then normalized to the media control by dividing to give a fold increase over the control ($n = 3$).

2.12. Statistical analysis

Statistics were performed with an OriginPro software (OriginPro, Massachusetts, USA). Significance set to $p < 0.05$ and tested in accordance with literature.\textsuperscript{23} Firstly, a Shapiro–Wilk test was used to test for normality and then a Grubbs outlier test was used to remove any outliers in non-normal data sets. If data still did not meet the normality assumption, a Kruskal–Wallis test was used to evaluate significance in data. For normal data, a Browne–Forsythe test was used to test the equal variance assumption. A $t$-test with a Welch correction was used between two samples if the equal-variance assumption was not met. A $t$-test was used for all analyses between only two samples. If the assumption of normality and equal variance was met, then an ANOVA with a Tukey post-hoc test was used to determine significance. For powers below 0.8, the data was deemed inconclusive. Sample numbers used for each group was based on previous studies utilizing similar collagen scaffolds:\textsuperscript{27,41–43,32} metabolic activity ($n = 6$), gene expression ($n = 5$), micro-CT ($n = 3$), ICP ($n = 3$), ALP ($n = 6$), cytokine array ($n = 3$). Error bars for all data are represented as mean ± standard deviation.

3. Results

3.1. SEM images and pore size analysis demonstrate anisotropic structure of scaffolds

SEM imaging revealed anisotropic scaffolds fabricated via a directional solidification approach demonstrated elongated pores with alignment in one direction (ESI Fig. 2\textsuperscript{†}). Stereology was subsequently used to quantify pore size and aspect ratio, revealing a significantly smaller aspect ratio in the transverse (vs. longitudinal) plane of anisotropic scaffolds to confirm structural anisotropy (ESI Fig. 3\textsuperscript{†}, Table 2). Averaged across all planes of analysis, there was no significant difference in the mean pore size of anisotropic and isotropic scaffolds.

3.2. All scaffold variants supported increasing MSC metabolic activity over 28 days

Pore orientation did not affect the metabolic activity of hMSCs seeded on scaffolds for 28 days. Isotropic and anisotropic variants both displayed an overall increase in MSC metabolic activity over 28 days in culture (Fig. 2A). Changing the glycosaminoglycan type within scaffolds also had no effect on metabolic activity, and at day 28 the metabolic activity of all scaffolds was greater than day 1 of the study (Fig. 2B).

3.3. Anisotropic scaffolds had an increase in chondrogenic gene, SOX9, at day 7, but no difference in osteogenic gene expression compared to isotropic variants

Osteogenic genes (RUNX2, Osterix, BMP2, and COL1A2), chondrogenic gene (SOX9), and adipogenic gene (PPAR\textgreek{g}) were examined in isotropic and anisotropic scaffolds over the course of 28 days. There was an increase in PPAR\textgreek{g} over 28 days in both scaffold groups, and a spike in SOX9 expression at day 7 in anisotropic scaffolds (Fig. 3). There were no significant differences in all osteogenic gene fold changes between isotropic and anisotropic variants. However, anisotropic scaffolds had an overall increase in gene expression of COL1A2 by day 7 not seen in isotropic scaffolds.

Fig. 2 Metabolic activity of hMSC seeded on mineralized collagen scaffolds with isotropic and anisotropic pores and GAGs chondroitin-6-sulfate (CS6), chondroitin-4-sulfate (CS4), and heparin sulfate (H). Metabolic activity was normalized to the metabolic activity of 150 000 hMSCs (cell seeding activity), indicated by a red dashed line. ^ indicates one group was significantly ($p < 0.05$) greater than the same group on day 1. Data expressed as average ± standard deviation ($n = 6$).
3.4. CS6-containing scaffolds had an increase in chondrogenic gene, SOX9, at day 7, and heparin-containing scaffolds had an increase in Osterix expression at day 4

Osteogenic genes (RUNX2, Osterix, BMP2, and COL1A2), chondrogenic gene (SOX9), and adipogenic gene (PPARγ) were examined in scaffolds containing CS6, CS4, and heparin over the course of 28 days (Fig. 4). There was an increase in PPARγ over 28 days in all scaffold groups, and CS6-containing scaffolds had a greater SOX9 expression than heparin-containing scaffolds at day 7. At day 7, heparin-containing scaffolds had greater expression of COL1A2 compared to day 1 than any other group.

3.5. Greater mineral formation in anisotropic and CS6-containing scaffolds

Mineral formation in scaffolds was evaluated by micro-CT and ICP analysis at day 28. As all scaffolds contained CaP mineral to begin with, results are reported as mineral content in scaffolds seeded with cells versus acellular scaffolds maintained in culture media. Micro-CT revealed anisotropic scaffolds had significantly greater mineral formation than isotropic scaffolds. Micro-CT revealed no differences in mineral formation between the different glycosaminoglycans, and all scaffolds had significantly greater mineral formation than day 0 unseeded scaffolds (Fig. 5A, ESI Fig. 4†). ICP analysis revealed that the CS6-containing scaffolds had significantly greater calcium and phosphorous than CS4- and heparin-containing scaffolds (Fig. 5B). Micro-CT was further validated by a greater calcium and phosphorous percent in the anisotropic variants.

3.6. Greater osteogenic activity in scaffolds containing Heparin

Osteogenic activity was quantified by an Alkaline Phosphatase assay, again with MSC-seeded scaffolds compared to acellular scaffolds with anisotropic and isotropic pores. * indicates one group is significantly (p < 0.05) greater than another group on the same day. ^ indicates one group was significantly (p < 0.05) greater than the same group on day 1. Anisotropic scaffolds had significantly (p < 0.05) greater SOX9, chondrogenic, gene expression on day 7 than isotropic scaffolds. Data expressed as average ± standard deviation (n = 5).

Fig. 3 Osteogenic, chondrogenic, and adipogenic gene expression of mineralized collagen scaffolds with anisotropic and isotropic pores. * indicates one group is significantly (p < 0.05) greater than another group on the same day. ^ indicates one group was significantly (p < 0.05) greater than the same group on day 1. Anisotropic scaffolds had significantly (p < 0.05) greater SOX9, chondrogenic, gene expression on day 7 than isotropic scaffolds. Data expressed as average ± standard deviation (n = 5).
controls. There was no difference in osteogenic activity between the isotropic and anisotropic scaffold variants, however, heparin-containing scaffolds had the greatest active bone formation day 28 (Fig. 6, Table 3). Interestingly, CS4-containing scaffolds had the lowest active bone formation at day 28.

3.7. No difference in migration efficiency between anisotropic and isotropic scaffolds containing CS6

Pore anisotropy did not increase cell invasion as measured via metabolic activity after 24 h. Anisotropic scaffolds had a higher metabolic activity, however, this was not significantly greater than isotropic scaffolds \( (p = 0.37) \) (ESI Fig. 4†).

3.8. Secretome from anisotropic and CS6-containing scaffolds showed increased levels of PDGF-BB and angiogenin, respectively

PDGF-BB was significantly more expressed by cells maintained in anisotropic scaffolds for 7 days (Table 4). Angiogenin was significantly more expressed in CS6-containing scaffolds versus other GAG variants (Table 4). Overall, levels of anti-inflammatory cytokines remained the same, IL-6 and IL-8 pro-inflammatory cytokines had high fold changes. Notably, osteogenic cytokines OPN and OPG were upregulated in all scaffold variants but was not influenced by pore anisotropy or GAG content.

4. Discussion

Craniomaxillofacial defects represent a unique and challenging type of bone defect. Herein, we explored modification of pore anisotropy and glycosaminoglycan content of a mineralized collagen scaffold under development for craniofacial bone repair. We hypothesized that anisotropic pores in the scaffold would enhance cell migration via directional guidance in order to subsequently improve osteogenic activity and new mineral biosynthesis. We further hypothesized that CS4 and Heparin in the scaffold would promote osteogenic activity and new mineral biosynthesis due to their potential to regenerate bone in literature.\(^{33-36}\)
We expected anisotropic pores to promote cell migration due to the aligned orientation of the pores; however, we saw no differences in either of these aspects between our anisotropic and isotropic pores. This contradicted some instances in literature, where accelerated migration could be achieved by anisotropic materials *in vivo* and *in vitro*, especially in aligned tissues. Bone is considered to be an aligned tissue, both the cortical and cancellous portions of bone having anisotropic behavior. It was of notice that a cytokine array revealed significantly greater expression of the chemokine PDGF-BB in anisotropic scaffolds than isotropic ones, thus a transwell study with longer timepoints may be of interest in the future to determine if migration effects are significant over longer periods of time. In addition, PDGF is important in the skeletal system, activating MSCs to differentiate to osteoblasts and activates macrophages. This could indicate that with significantly greater PDGF levels in anisotropic scaffolds, these scaffolds had greater MSC differentiation to osteoblasts. Although migration and activity of hMSCs was unaffected by anisotropy, this pore orientation change effected the ultimate behavior of the cells and mineral formation.

Both scaffold types had upregulated osteogenic genes Runx2, BMP2, Osterix, and COL1A2 but anisotropic variants had a significant increase in chondrogenic gene, SOX9, at day 7. This

![Graphs showing mineral quantification](image)

**Fig. 5** Mineral quantification of unseeded and hMSC seeded scaffolds by micro-CT and ICP. (A) Micro-CT image stacks were processed in ImageJ with a threshold of 214/255 and the white area was compared to the total scaffold area to produce a % area fill. No statistical significance was found between the CS6, CS4, and H scaffold groups, however, all scaffolds had significantly (*p* < 0.05) greater % area fill at day 28 compared to day 0. * indicates there was a significant (*p* < 0.05) difference in % area fill between the isotropic and anisotropic groups seeded with cells (power: 0.78794). (B) ICP analysis of calcium and phosphorous content of scaffolds post-micro-CT scanning and critical point drying compared to dry unseeded scaffolds. * indicates there is significantly (*p* < 0.05) greater % found in at day 28 than day 0 of the same group. Data expressed as average ± standard deviation (*n* = 3).
greater mineral formation did not negatively impact osteogenic activity, as anisotropic scaffolds had significantly greater mineral formation than isotropic variants at the end of the study. RUNX2 is a major transcription factor of bone, involved in the MSC transition to osteoblasts. There was no significant effect of pore orientation on RUNX2 gene expression; however, this was upregulated throughout the study, indicating differentiation of hMSCs in both scaffold variants. The COL1A2 gene is highly expressed in most connective tissues, such as bone, skin, and tendon. This gene was upregulated throughout the study, but expressed the greatest at day 7, indicating cells producing type I collagen, which could be related to bone formation occurring. BMP2 is also involved in osteogenic differentiation, as well as bone formation. BMP2 was only upregulated at days 7 and 14, possibly indicating that the greatest bone formation or differentiation was occurring at these timepoints. Osterix is important for late-stage bone formation, regulating the differentiation of osteoprogenitors to osteoblasts. Osterix was upregulated at the early-stages of the study, possibly indicating hMSCs were beginning to differentiate early-on. Interestingly, the expression of the osteoclast inhibitory factor OPG was upregulated in all scaffold groups, particularly in the anisotropic mineralized scaffold. We have previously shown that mineralized collagen scaffolds containing isotropic pores and CS6 induce MSCs to produce OPG when seeded with MSCs to enhance bone regeneration. Suggesting the anisotropic mineralized scaffold may be particularly interesting for future studies to explore MSC-osteoclast interactions.

Although there were minimal differences in gene expression between isotropic and anisotropic scaffolds, we observed greater mineral formation via ICP and micro-CT analyses in anisotropic scaffolds. At the end of the study, total amounts of calcium and phosphorous were greater in the anisotropic scaffold compared to the isotropic scaffold. Interestingly, a study by Bobbert et al. revealed that aligned fibers in PCL structures had a lower ALP activity than random fibers. This difference in results could be attributed to a difference in metabolic activity of PCL structures; however, our scaffolds had no metabolic activity differences, indicating no bias on ALP activity. Of note, another study by Seong et al. indicated that small channels enhanced proliferation and migration in vitro and pore sizes of 270 µm were best for cell migration in vivo. This could explain the lack of migration and metabolic activity differences in our scaffolds, which had average pore diameters in the range of 150–200 µm.

We expected addition of chondroitin-4-sulfate or heparin sulfate to increase osteogenic activity and bone formation versus chondroitin-6-sulfate-containing scaffolds. This was based on observations in the literature that CS4 is more involved in the bone formation process while CS6 is involved in cartilage formation. We found no differences in metabolic activity between any of the glycosaminoglycans, but all promoted a greater than 2.5-fold change in activity after 28 days. Overall, osteogenic activity was observed throughout all variants. However, heparin-functionalized variants increased relative expression (vs. day 1) of COL1A2 as well as greater expression of Osterix than CS4 or CS6 variants, suggesting more osteoblastic differentiation in heparin scaffolds at this period. CS6-functionalized variants also showed an early (day 7) spike in SOX9 chondrogenic gene expression. However, overall, all osteogenic genes were upregulated for most of the study regardless of glycosaminoglycan type, suggesting future studies to look at long-term functional responses in vitro and in vivo.

Analysis of mineral formation via micro-CT found increased mineral deposition in all variants but was unable to resolve difference in mineral formation at 28 days in culture. Analysis of

![Graph](image_url)  
**Fig. 6** Active bone formation in scaffolds with varying pore orientation and glycosaminoglycan content. A colorimetric ALP assay was used to determine the ALP enzymatic activity in scaffolds after 7 and 28 days, which can be related to active bone formation. Values of the seeded scaffolds at days 7 and 28 were subtracted from unseeded scaffolds at the same days, then normalized to day 0 unseeded controls to achieve a fold change in ALP activity from the start of the experiment (1 represents activity at day 0). Chondroitin-6-sulfate, chondroitin-4-sulfate, and heparin sulfate-containing scaffolds are represented by CS6, CS4, and H (respectively). ^ indicates one scaffold group has significantly (p < 0.05) greater ALP activity at day 28 compared to the same group at day 7. ** indicates the heparin scaffold group had significantly (p < 0.05) greater ALP activity at day 7 compared to both CS6 and CS4. * indicates one scaffold group is significantly (p < 0.05) greater than a different group on the same day. Data expressed as average ± standard deviation (n = 6).
ALP levels indicated that active bone formation was greatest in Heparin scaffolds at day 7 and 28, although, there were no differences in ALP activity between Heparin and CS6 scaffolds at day 28. Overall, CS4-containing scaffolds showed the smallest amount of mineral formation compared to CS6 or Heparin scaffolds. While not examined explicitly here, strategies to improve angiogenesis are essential for large volume bone regeneration; analysis of the secretome of MSCs in all scaffolds revealed CS6 or Heparin-containing anisotropic scaffolds showed significantly greater angiogenin expression than CS4 variants. Overall, our data suggest anisotropic mineralized collagen scaffolds containing either CS6 or Heparin GAGs have the greatest potential for bone regeneration.

Table 3  Alkaline phosphatase concentrations in unseeded and cell-seeded scaffolds with a media control

|             | Isotropic  | Day 0          | Day 7 unseeded | Day 7          | Day 28 unseeded | Day 28          |
|-------------|------------|----------------|----------------|----------------|-----------------|-----------------|
|             | Isotropic  | 67.9 ± 64.5    | 3.1 ± 5.2      | 27.1 ± 13.8    | 11.9 ± 6.3      | 102.6 ± 5.1     |
|             | Anisotropic| 0.5 ± 3.1      | 7.4 ± 3.1      | 20.6 ± 4.6     | 12.3 ± 12.7     | 122.0 ± 39.0    |
| CS4         | 0.5 ± 3.1  | 7.4 ± 3.1      | 20.6 ± 4.6     | 12.3 ± 12.7    | 122.0 ± 39.0    | 87.4 ± 11.1     |
| Heparin     | 5.5 ± 0.9  | 0.5 ± 1.4      | 37.0 ± 6.5     | 4.1 ± 0.6      |                 | 159.5 ± 16.8    |
| Media       | 31.3       |                |                |                |                 |                 |

Table 4  Secreted cytokines of hMSC seeded scaffolds with varying pore orientation. A cytokine assay was conducted on eluted media after 7 days from scaffolds with isotropic and anisotropic pores. Values of array intensity were measured with ImageJ, and a control array (exposed only to media) was used to determine fold change due to cellular activity. 5 key groups are shown – cytokines connected to inflammation, anti-inflammatory cytokines, cytokines connected to cell movement, angiogenic cytokines, bone formation cytokines. Green indicates cytokines with a higher fold change and yellow indicates limited or no change compared to media. * indicates a statistically significant (p < 0.05) increase in cytokine level for anisotropic pores over isotropic pores (power: 0.69176). Data expressed as average ± standard deviation (n = 3)

5. Conclusions

This study examined the role of pore anisotropy and GAG composition on MSC osteogenic activity in a class of mineralized collagen scaffolds under development for CMF bone repair applications. We hypothesized that anisotropic scaffolds would promote greater cell migration, osteogenic activity, and bone formation. We found that anisotropic pores did not increase cell activity or migration but did improve overall bone formation. We also hypothesized that inclusion of chondroitin-4-sulfate and heparin sulfate would increase cell activity and bone formation in mineralized collagen scaffolds. Notably, chondroitin-6-sulfate and heparin sulfate had the greatest effect on improved mineral formation. These efforts represent an important step toward optimizing mineralized collagen scaffolds for bone regeneration. These studies also set the stage to examine the role of anisotropy and GAG content on...
inflammatory and angiogenic responses as well as crosstalk between these axes and MSCs.

**Conflicts of interest**

There are no conflicts to declare.

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