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Human chondrocyte migration behaviour to guide the development of engineered cartilage

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Abstract

Tissue-engineering techniques have been successful in developing cartilage-like tissues *in vitro* using cells from animal sources. The successful translation of these strategies to the clinic will likely require cell expansion to achieve sufficient cell numbers. Using a two-dimensional (2D) cell migration assay to first identify the passage at which chondrocytes exhibited their greatest chondrogenic potential, the objective of this study was to determine a more optimal culture medium for developing three-dimensional (3D) cartilage-like tissues using human cells. We evaluated combinations of commonly used growth factors that have been shown to promote chondrogenic growth and development. Human articular chondrocytes (AC) from osteoarthritic (OA) joints were cultured in 3D environments, either in pellets or encapsulated in agarose. The effect of growth factor supplementation was dependent on the environment, such that matrix deposition differed between the two culture systems. ACs in pellet culture were more responsive to bone morphogenetic protein (BMP2) alone or combinations containing BMP2 (i.e. BMP2 with PDGF or FGF). However, engineered cartilage development within agarose was better for constructs cultured with TGF β 3. These results with agarose and pellet culture studies set the stage for the development of conditions appropriate for culturing 3D functional engineered cartilage for eventual use in human therapies.

Keywords

osteoarthritis; chondrocytes; pellet culture; agarose; tissue engineering; articular cartilage

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1. Introduction

Adult articular cartilage is an avascular, load-bearing tissue that has limited capacity to heal following injury or degeneration. Osteoarthritis is the leading cause of disability in Americans and is associated with medical costs > ~\$130 billion (Kotlarz *et al.*, 2009; Luo *et al.*, 2004). Successful biological repair of osteoarthritic (OA) or injured cartilage may reduce the need for artificial joint replacement by providing long-term pain relief and maintaining the mechanical function of the healthy native tissue. Cultivating engineered tissue *in vitro* has shown promising results for development of engineered cartilage using animal chondrocytes and stem cells (O'Connell *et al.*, 2012a, 2012b; Pei *et al.*, 2008, 2009; Rodrigues *et al.*, 2011; Sampat *et al.*, 2011). For clinical applications, autologous chondrocytes arguably represent the ideal cell source. Aside from the usual concerns of donor site morbidity, however, developing functional engineered cartilage using cells isolated from diseased (e.g. arthritic) tissue is not trivial (Selmi *et al.*, 2008). The successful translation of engineered cartilage to the clinic will necessitate cell expansion to achieve sufficient cell numbers. To date, the culture conditions required for expanding and cultivating engineered cartilage with human cells that produce native properties remain elusive. Current methods for human cells reported in the literature employ various passage conditions and culture media formulae to support growth of chondrocytes (e.g. (Adesida *et al.*, 2012; Alves da Silva *et al.*, 2011; Bian *et al.*, 2011; de Mara *et al.*, 2013; Ghone and Grayson, 2012; Handorf and Li, 2011; Matsumoto *et al.*, 2012; Mendelson *et al.*, 2011; Park *et al.*, 2011; Perrier *et al.*, 2011; Wu *et al.*, 2012; Yin *et al.*, 2011; Yoo *et al.*, 2011)).

In this study, we first evaluated the effect of passaging on cell migration and movement in two-dimensional (2D) culture to guide an appropriate condition (i.e. passage number) for developing three-dimensional (3D) engineered cartilage. The use of direct current (DC) electric fields (EFs) of strengths in the range 1–10 V/cm is known to induce directed movement (galvanotaxis) and shape change (galvanotropism) in a number of terminally differentiated musculoskeletal cells, including chondrocytes, fibroblasts, osteoblasts, osteoclasts and meniscal fibrochondrocytes (Chao *et al.*, 2000; Ferrier *et al.*, 1986; Gunja *et al.*, 2010, 2012; Sillman *et al.*, 2003). Endogenously generated electric field gradients of this strength have been shown to guide cell migration in developing embryos (Robinson, 1985) and at the cut surfaces of wounds (Soong *et al.*, 1990). Using the well-established bovine model for cartilage research (Hung *et al.*, 2004; O'Connell *et al.*, 2012a, 2012b; Sampat *et al.*, 2011), our laboratory has recently demonstrated that calf synovial-derived stem cells (SDSCs), expanded in the presence of a growth factor cocktail comprised of TGF β -1, FGF-2 and PDGF- $\beta\beta$, display passage-dependent cell surface marker expression and migration behaviour in response to applied EFs (Tan *et al.*, 2014). Specifically, late-passage cells (e.g. passages 3 and 4), with sufficient chemical priming, responded to DC EFs with cathodal migration, which is the same direction in which chondrocytes travel (Chao *et al.*, 2000), and were capable of creating cartilaginous-like tissue with elevated amounts of collagen II and GAG deposition (Tan *et al.*, 2014). Importantly, we have also observed similar findings for human MSCs with passage (unpublished). In this manner, we believe that galvanotaxis provides a useful 2D assay for identifying cell populations that exhibit a high propensity to produce cartilage-like tissue when cultured in 3D conditions.

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Monolayer and pellet cultures with human chondrocytes have demonstrated that chondrogenesis can be improved by using various growth factors and their combinations. Bone morphogenetic protein-2 (BMP2) has been suggested to increase aggrecan and collagen type II expression and to prevent dedifferentiation of chondrocytes (Luyten *et al.*, 1992; Sailor *et al.*, 1996; Stewart *et al.*, 2000; Theodoropoulos *et al.*, 2011). Previous studies demonstrated beneficial effects of TGF β on collagen type II production and gene expression (Chiang *et al.*, 2011). Furthermore, combining growth factors, such as TGF β with BMP2, has shown potential to further enhance aggrecan and collagen gene expression (Mendelson *et al.*, 2011; Perrier *et al.*, 2011). While these studies provide valuable information about the cellular response to chemical stimuli, the culture conditions of these relatively short duration studies (i.e. 7–14 days) that monitor gene expression and matrix production in pellet culture may not be indicative of conditions that will lead to long-term functional cartilage development. A successful biological repair strategy for damaged or injured cartilage will need to exhibit functional tissue properties that can withstand and distribute the high compressive joint loads experienced by the native tissue.

With the growing success in developing functional engineered cartilage *in vitro* using chondrocytes and stem cells derived from various animal models (Farrell *et al.*, 2013; Huey *et al.*, 2012; Johnstone *et al.*, 2013; Natoli *et al.*, 2009; O'Connell *et al.*, 2012b), there has been a shift in attention toward translating these findings to developing engineered cartilage with human cells (Naumann *et al.*, 2004). Adopting a suitable cell passage from the galvanotaxis response of the OA chondrocytes, as described above, we then cultured human cells to determine whether stimulation with growth factors that have been used to improve cartilage production in human articular chondrocyte pellet or self-aggregating constructs (Bian *et al.*, 2011; Matsumoto *et al.*, 2012; Yin *et al.*, 2011) is translatable to their culture in agarose, a clinically-relevant hydrogel scaffold for cartilage repair strategies (Selmi *et al.*, 2008). While incorporation of agarose hydrogel would affect the nature of cell–cell interactions indicative of cell-only constructs, fewer cells would be needed to generate an engineered tissue of a specified volume. Moreover, the ability to use human cells successfully with the agarose system would permit direct translation of strategies such as applied deformational loading (Bian *et al.*, 2012; Mauck *et al.*, 2002) and osmotic loading, which have been shown to promote *in vitro* functional cartilage tissue development of non-human mammalian cells encapsulated in agarose (Sampat *et al.*, 2011). Thus, in this study, we examined matrix production and mechanical properties of 3D engineered cartilage in pellet and agarose culture under several different growth factor conditions.

2. Materials and methods

Articular cartilage was acquired according to an IRB-approved protocol from two patients, aged 45 and 51 years, receiving total knee replacement (New York University School of Medicine). Chondrocytes were acquired as described by Attur *et al.* (2012). Cells were expanded in Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% serum, 1% 100 U/ml penicillin, 100 mg/ml streptomycin and amphotericin B (Invitrogen, Carlsbad, CA, USA), 1 ng/ml TGF β 1, 10 ng/ml PDGF $\beta\beta$ and 5 ng/ml FGF2. Cells were passaged four times for galvanotaxis studies, and the results from the galvanotaxis study (study 1)

informed which passage to use for 3D culture (study 2; Figure 1). The same donor cells were expanded for 3D cultures in pellets and agarose scaffolds.

2.1. Galvanotaxis

At each passage, when confluence had been reached, cells were trypsinized and counted. One subset was replated for further expansion and another subset was resuspended at 50×10^3 cells/ml in DMEM containing 5% serum, amino acids (0.5× minimal essential amino acids, 1× non-essential amino acids), buffers (10 mM HEPES, 10 mM sodium bicarbonate, 10 mM TES, 10 mM BES) and antibiotics (100 U/ml penicillin, 100 mg/ml streptomycin) and allowed to equilibrate for 1 h. Then the cells were plated at 2.65×10^4 cells/cm² onto sterile glass slides (Fisher Scientific, Pittsburgh, PA, USA), using removable silicone wells, and allowed to attach for 1 h in a 5% CO₂ incubator at 37 °C. The slides were rinsed with medium to remove any non-adherent cells and placed into a custom galvanotaxis chamber under aseptic conditions (Figure 1). A modified parallel-plate flow chamber was used, as described previously (Chao *et al.*, 2000). Briefly, two Plexiglas blocks with glass windows were separated by a uniform thickness rectangular Silastic spacer (McMaster-Carr, New Brunswick, NJ, USA) and held together with screws to create a sealed rectangular channel with defined geometry (5 cm length, 1.3 cm width, 250 µm height). A power supply (Keithley Instruments) delivered a current of 3.3 mA (6V/cm electric field strength) through the chamber, and experiments were performed at room temperature for 3 h. Control slides were treated similarly, except that no electric field was applied.

Cell migration patterns were captured using a digital camera connected to EOS Utility (Canon Rebel T3i, Canon, USA). Using a × 10 objective (1 pixel = 0.17 µm), approximately 30–40 cells/slide were observed; images were acquired at 10 min intervals. The incremental change in position for each cell was automatically tracked with a custom MATLAB algorithm. Overall cell migration speed was calculated as the net displacement divided by the 3 h time span. Migration direction was quantified as sin φ, where φ is the angle between the x coordinate axis and the migration vector, such that sin φ was defined as the value -1 when φ = 4.71 rad (270°), the direction of the cathode. The directional velocity, defined as the component of the speed directed toward the negative pole (Chao *et al.*, 2000), was obtained by multiplying the cell's speed by sin φ. Each data point represented the mean and standard error (SE; 60–80 samples). Two-way ANOVA was performed on cell speed, directed velocity and total displacement for factors of treatment (applied EFs vs control) and passage number. Tukey's *post hoc* analysis was performed to assess significance between each treatment and passage group; significance was set at $p < 0.05$.

To correlate chondrogenic phenotype with EF-induced migration direction, cells from passages that were directed to the cathode (P2) and anode (P3) were placed in pellet culture, and aggrecan expression was measured after 7 days (see section on 3D cultures, below, for culture details). Pellets were stored in Trizol reagent to perform RNA extraction and total sample cDNA generation. Real-time PCR was then run for human aggrecan and β-actin, a known housekeeping gene. Unpaired *t*-test was performed for aggrecan gene expression normalized to β-actin expression, to compare P2 and P3 OA chondrocytes.

2.2. 3D cultures

P2 human OA chondrocytes exhibited cathodal migration that yielded cell pellets with increased aggrecan expression over other passages examined (see Figure 3 and detailed results below). Previously, P2 bovine chondrocytes were found to display excellent capabilities in developing functional engineered cartilage in agarose cultures (O'Connell *et al.*, 2014). Therefore, cells were passaged twice before using them in 3D culture studies for subsequent studies (Figure 2) (O'Connell *et al.*, 2014).

Once confluence was reached during the second passage, cells were trypsinized and pellets were formed for each cell type by centrifugation. The pellets were cultured for 28 days in a chemically defined medium [CM; DMEM with 0.1 µM dexamethasone, 40 mg/ml L-proline, 50 mg/ml ascorbate 2-phosphate, 100 mg/ml sodium pyruvate, 1× ITS⁺ premix, 100 U/ml penicillin and 100 mg/ml streptomycin and amphotericin B (Invitrogen), supplemented with growth factors]. Based on previously reported values for growth factor concentrations and uses in 3D tissue-culture studies, pellets were cultured with CM supplemented with the following growth factor combinations: 50 ng/ml BMP2 with 10 ng/ml FGF2; and 50 ng/ml BMP2 with 10 ng/ml PDGF $\beta\beta$ (Balakrishnan *et al.*, 2014; Barbero *et al.*, 2004; Chuang *et al.*, 2012; de Mara *et al.*, 2013; Handorf and Li, 2011; Johns and Athanasiou, 2008; Mayer-Wagner *et al.*, 2011; Perrier *et al.*, 2011). Pellets cultured with 10 ng/ml TGF β 3 only or 50 ng/ml BMP2 only served as single growth factor controls (Figure 2). TGF β 3 only was chosen as a study group, as it is directly comparable to our previous work with animal cell sources and one of the most commonly used growth factors in cartilage tissue-engineering studies (Huang *et al.*, 2009; O'Connell *et al.*, 2012b).

Pellets were cultured for 28 days and extracellular matrix production was assessed bi-weekly (i.e. days 0, 14 and 28; $n = 5$ /group at each time point). Media aliquots were saved to measure GAG release into the medium and pellets were prepared for biochemical analysis. The pellet culture was repeated using chondrocytes from a second donor to demonstrate robustness of the results. DNA content was measured using the PicoGreen Kit (Invitrogen). GAGs were measured using DMMB assay, and collagen content was measured using hydroxyproline assay. Two-way ANOVA was performed on the biochemical properties, with the treatment group and time as factors. Bonferroni *post hoc* analysis was used when significance was found ($p < 0.05$).

Following the pellet study, cells were passaged for 3D culture in 2% w/v low-gelling agarose slab (Type VII, Sigma-Aldrich) at a final concentration of 30×10^6 cells/ml (Mauck *et al.*, 2002). Constructs were punched from the slab (2.34 mm thickness) using a 4 mm diameter biopsy punch and cultured in CM supplemented with growth factors, based on the results from the pellet culture. Due to the number of cells required for agarose culture, chondrocyte-laden constructs were cultured with either 10 ng/ml TGF β 3 (control) or 50 ng/ml BMP2.

At day 28, cell viability in agarose constructs was assessed with the Live/Dead Kit to determine cell distribution throughout the construct ($n = 1$ /group; Invitrogen). At day 56, mechanical, biochemical and histological analyses were performed ($n = 6$ –8/group/donor). The equilibrium Young's modulus was determined from a stress relaxation test in unconfined compression at 10% strain. A sinusoidal compressive strain ($\pm 1\%$, 0.5 Hz) was

then applied to determine the dynamic modulus at that frequency. Wet weights (ww) were measured following mechanical testing and the samples were dried to obtain dry weights (dw). DNA content was determined using the PicoGreen Kit (Invitrogen). The GAG and collagen content were determined using 1,9-dimethylmethylen blue (DMMB) and hydroxyproline assay, respectively. The ratio of hydroxyproline to collagen was assumed to be 7.64, based on the molecular weights provided in Hollander *et al.* (1994). The GAG and collagen contents were normalized by wet weight, dry weight and DNA content. Student's *t*-test was performed to compare mechanical and biochemical properties assessed from 3D agarose cultures. Biochemical composition was normalized by wet and dry weight as a measure of matrix composition in the *de novo* cartilage, and for comparison with previously reported values of native cartilage. Normalization by DNA content was used as a measurement of cell activity in matrix production.

At day 56, extra constructs were preserved for histology to evaluate the distribution of cells, collagen and GAGs. Briefly, histological samples were embedded in paraffin wax and slides were prepared (8 µm sections). The sections were stained with safranin-O for GAG distribution and picrosirius red for collagen distribution. Sections were also stained for collagen types I, II and VI. Briefly, the sections were digested in 0.5 mg/ml testicular hyaluronidase, swollen in 0.5 M acetic acid, blocked in 10% normal goat serum (NGS) and labelled with 10% NGS containing monoclonal primary antibody for collagen types II and VI (Abcam). AlexaFluor 488-conjugated goat anti-mouse secondary antibody labelling (Invitrogen) and TOTO-3 nuclear counterstaining (Invitrogen) were performed to visualize the collagen networks and cells, respectively. After staining, slides were coverslipped with Fluoroshield (Sigma) and the sections were analysed using an inverted microscope with an Olympus Fluoview confocal system, with dual wavelength excitation at 488 and 640 nm.

3. Results

At all passages, chondrocytes exposed to an applied direct current electric field travelled faster and farther than cells not exposed to a field ($p < 0.0001$; Figure 3A). By late passage (P4), EF-exposed cells were more responsive to the applied field, showing increased migration speeds compared to early passage (P2) cells (11.6 ± 0.84 vs 8.34 ± 0.76 µm/h; $p < 0.05$; Figure 3A). This change in overall speed through passage number was mirrored with a change in migration direction travelled by EF-exposed cells. Control cells exhibited no preferential direction (i.e. directed velocity = 0 µm/h; $p > 0.05$); however, EF-exposed cells travelled significantly further toward a specific pole, depending on the passage. P2 cells expressed preferential migration toward the cathode (negative pole, -6.21 ± 0.95 µm/h; $p < 0.0001$); later passage cells began changing their direction of migration, turning notably toward the anode (positive pole) by P3 (Figure 3B; 1.21 ± 0.93 µm/h; $p < 0.05$) and increasingly continuing for P4 (Figure 3B; 2.39 ± 1.08 µm/h; $p < 0.05$). Therefore, P2 cells were used for 3D pellet and agarose culture studies.

Subsequently, a relationship between galvanotaxis response and chondrogenic phenotype was investigated for pellet cultures established from P2 and P3 OA chondrocytes (pooled donors). Cathodal-directed P2 cells exhibited a nearly five-fold increase in aggrecan expression compared to those derived from anodal-directed P3 cells (P2, $4.72 \pm 2.52 \times 10^{-5}$

vs P3, $0.97 \pm 0.77 \times 10^{-5}$; $p < 0.05$; $n = 5\text{--}6$ samples/group). Together, these findings led us to adopt P2 OA chondrocytes for our subsequent studies comparing the influence of growth factors on pellet and agarose cultures.

Pellets cultured in CM supplemented with BMP2 were significantly larger than the control (Figure 4, inset). Human ACs maintained cell viability throughout the culture period, as denoted by vital staining as well as DNA content (Figure 4A). Cell proliferation was observed for pellets cultured with BMP2 + PDGF $\beta\beta$, where the DNA content at day 14 was almost 2 \times the day 0 values ($p < 0.0001$; Figure 4A, ^). However, there was a decrease in DNA content from day 14 to day 28 (Figure 4A, b), resulting in a final DNA content that was not significantly different from day 0. By day 28, the GAG content in pellets cultured with BMP2 only and BMP2 + FGF2 was 80% greater than the control ($p = 0.015$; Figure 4B). The GAGs measured in the media aliquots accounted for 5.0–7.4% of the total GAG content measured (i.e. pellet GAG + medium GAG). The GAG content in the medium normalized by DNA was greatest in the BMP2-only group and was 70% greater than the control (medium GAG/DNA for the BMP2 group = 4.9 ± 0.7 mg/mg; Figure 4B, z). At day 28, the collagen content normalized by DNA content was 56.6 ± 26.4 mg/mg for the control group, and there were no significant differences with growth factor treatment ($p = 0.14$; Figure 4C). The ratio of total GAG content: collagen content was not significantly different across the experimental groups ($p = 0.12$).

Cell viability in agarose was maintained throughout the culture period (Figure 5A, Figure 6A top row) and cell proliferation was observed in constructs cultured with TGF β 3 (DNA content at day 56, 6.85 ± 0.44 μ g; vs 1.18 ± 0.16 μ g for the BMP2 group; $p < 0.001$; Figure 6A). Young's modulus of the agarose constructs was 9.3 ± 1.2 kPa after casting (day 0). At day 56, Young's modulus of the TGF β 3 group was 96.0 ± 16.5 kPa and was 6.5 times greater than in the BMP2 group ($p = 0.0001$; Figure 5B). Similarly, the dynamic modulus of the TGF β 3 group was 1120 ± 190 kPa and was six times greater than in the BMP2 group (175 ± 35 kPa; $p < 0.0001$). This finding was repeated independently using chondrocytes derived from another donor, where the Young's modulus of engineered cartilage developed was approximately 2.5 times greater for the TGF β group than the BMP2 group (data not shown).

Agarose constructs increased in opacity with culture, demonstrating increased matrix deposition throughout the construct (Figure 5, inset). GAG content of engineered constructs cultured with TGF β 3 was two to eight times greater than the BMP2 group for GAG normalized by dry weight (TGF β 3 group = $23.4 \pm 5.4\%$ /dw), wet weight ($2.16 \pm 0.51\%$ ww) and DNA content (116.3 ± 22.3 μ g/ μ g; $p < 0.003$; Figure 5C). The intensity of safranin-O staining for GAGs was much greater for the TGF β 3 group and was more uniformly distributed throughout the construct than the BMP2 group (Figure 6, middle row). The collagen content of the TGF β 3 group was $18.03 \pm 6.21\%$ dw, or 91.3 ± 35.8 μ g/ μ g when normalized by DNA content, and was not significantly different from the collagen content of the BMP2 group ($p > 0.3$; Figure 5D). The collagen content normalized by wet weight was $1.66 \pm 0.57\%$ ww for the TGF β 3 group and was significantly greater than in the BMP2 group ($0.48 \pm 0.59\%$ /ww; $p = 0.01$). Collagen was uniformly distributed in both the TGF β 3 and BMP2 groups, but staining was greater for the TGF β 3 group, which compared with the

quantitative assay data (Figure 6, bottom row). Immunohistochemistry stained positive for collagen II and VI for the TGF β 3 and BMP2 groups. Collagen VI in the BMP2 group remained closer to the cell edge than in the TGF β 3 group (Figure 6, bottom row inset).

4. Discussion

Engineering of articular cartilage may provide a biological repair strategy for injured or arthritic cartilage. Extensive work has been performed using animal models and cells from animal donors (Hung *et al.*, 2004; Johnstone *et al.*, 2013; Kock *et al.*, 2012; O'Connell *et al.*, 2012b). However, direct translation of culture techniques and conditions appropriate for growing *de novo* animal cartilage tissue to human tissue has been challenging. Our laboratory, and others, have demonstrated that supplementation of expansion medium with a growth factor cocktail of FGF, PDGF and TGF β 1 is crucial for developing functional engineered cartilage in 3D cultures (Pei *et al.*, 2008; Sampat *et al.*, 2011). In this study, an applied DC EF was applied to human chondrocytes from OA cartilage to determine a passaging limit for 2D expansion, in order to maintain the chondrogenic phenotype. Galvanotaxis to the cathode is exhibited by normal chondrocytes and stem cells expanded with growth factors for defined passages (Chao *et al.*, 2000; Tan *et al.*, 2014). Our results suggest that using human chondrocytes at P2 may improve the matrix production and mechanical properties of 3D engineered functional cartilage. It was observed that matrix production by P2 human chondrocytes was dependent on the 3D culture environment (pellet vs agarose hydrogel) and growth factor supplementation. The findings from this study further support the notion that functional engineered cartilage can be developed from chondrocytes from OA tissues (Bian *et al.*, 2011).

Autologous chondrocytes are the ideal cell source for clinically relevant engineered cartilage implants; however, these cells need to be expanded to obtain sufficient cell numbers for developing neocartilage. With each subsequent passage, the chondrocytes dedifferentiate towards a fibroblastic lineage, which is noted by a change in morphology and increased cell migration (Figure 3) (Chao *et al.*, 2000, 2007). We observed that cell migration behaviour of OA P2 chondrocytes was similar to that of juvenile bovine chondrocytes, which is a robust cell source for developing neocartilage *in vitro* (O'Connell *et al.*, 2012b). Although the current study was limited by the number of human donors ($n=2$), we suggest that the high repeatability of our experiments across donors is indicative of a robust observation. As such, our findings reported here suggest that adult chondrocytes can be passaged twice for expansion, while maintaining the chondrogenic phenotype, for developing neocartilage.

For the conditions of the present study, large differences were observed in matrix production between pellet and agarose culture conditions, which agrees with previous studies reporting gene expression of 3D cultures (Dehne *et al.*, 2009). In pellet culture, BMP2 increased the swelling and physical size of the *de novo* tissue with human chondrocytes (Figure 4, inset). Furthermore, BMP2 demonstrated increased GAG production under specific conditions. However, matrix production and mechanical properties were higher for chondrocyte-seeded agarose constructs cultured with TGF β 3. It is possible that these differences in cell behaviour and matrix production are due to differences in cell-cell interactions, or in the diffusion of nutrients or growth factors (Bian *et al.*, 2013; Pei *et al.*, 2008). Although the

sizes of TGF β 3 (13 kDa) and BMP2 (26 kDa) differ only by a factor of two; this difference may be sufficient to restrict diffusion of nutrients, such as growth factors, through the construct (Albro *et al.*, 2009; Chahine *et al.*, 2009).

Gene expression and, in turn, matrix production potential decreases significantly with age (Liu *et al.*, 2013). Therefore, the results from this study, which used chondrocytes from OA knee joints, demonstrates promising results towards developing functional engineered cartilage from patient tissues. The GAG deposition observed in the agarose constructs was comparable to native values (2.2% ww at day 56; native cartilage 2–3% ww) (Treppo *et al.*, 2000). Collagen deposition was significantly lower than native values (native cartilage = 10–15% ww) (O'Connell *et al.*, 2012b; Treppo *et al.*, 2000), which represents a limitation of the hydrogel scaffold system despite the fact that it maintains the round chondrocyte morphology.

The compressive Young's modulus of engineered cartilage constructs cultured with TGF β 3 was approximately 100 kPa (native cartilage = 400–600 kPa) (Jurvelin *et al.*, 2003; Treppo *et al.*, 2000). These results are encouraging, as they suggest that human chondrocytes from OA tissues can be used to develop larger engineered cartilage tissues (i.e. millimeter-scale). Moreover, recent work has demonstrated better integration of biological repair tissues when the mechanical properties of the repair tissue are lower than those of the surrounding native tissue (Miot *et al.*, 2012). Miot *et al.* (2012) demonstrated that a shorter *in vitro* culture for engineered cartilage provided a balance between integration with the native cartilage and matrix deposition. Therefore, these results suggest that engineered cartilage constructs may be developed *in vitro* with mechanical integrity that may be sufficient for initial repair and integration into native tissue.

While we demonstrate that neocartilage produced by chondrocytes from OA cartilage resembles native cartilage, it remains to be seen whether these tissues function as normal cartilage after implantation. Further assessment of the tissue response to applied stimuli, including proinflammatory cytokines, will help to determine whether the cells maintain an osteoarthritic phenotype. In this context, these engineered tissues could alternatively serve as a platform to study pathological cartilage (Musumeci *et al.*, 2011).

There are many studies on human cells that have evaluated gene expression in monolayer culture in media supplemented with serum. Our preliminary findings in pellet culture with serum demonstrated an increase in cell proliferation; however, matrix production of serum-supplemented pellets was significantly lower than that in pellets cultured without serum (data not shown). These findings are comparable to the findings of Stewart *et al.* (2000), in which serum supplementation decreased gene expression of aggrecan and collagen in pellet culture when compared to a serum-free culture condition. Similarly, our pellets cultured with BMP2 + PDGF demonstrated that increases in cell proliferation resulted in a relative decrease in GAG content when normalized by DNA (Figure 4). These results demonstrate that culture conditions need to balance cell proliferation and matrix production, as some media components may cause cells to focus more energy on proliferation at the expense of synthesizing cartilage with mechanical and biochemical properties comparable to native values.

5. Conclusion

The duration of the studies performed here was significantly longer than most studies that evaluate gene expression or matrix production of monolayer and pellet cultures. Our findings suggest that culture periods > 8 weeks will be necessary to develop mature engineered cartilage in free swelling conditions prior to implantation (chondrocyte-seeded scaffolds cultured with TGF β 3 was approximately 100 kPa at day 56; Figure 5). While these findings are encouraging, long culture times are a significant limitation for clinical applications, due to the increased time between initial tissue harvest and final implantation surgeries. Future work will focus on improving the rate of matrix deposition by applying dynamic loading in culture, which have been shown to upregulate BMP2 and TGF β 3 expression (Nam *et al.*, 2013; Tenney and Discher, 2009) and increase extracellular matrix deposition rates of agarose constructs seeded with bovine cells (Kelly *et al.*, 2013; Sampat *et al.*, 2011). Additionally, these efforts would likely benefit from incorporation of strategies that can provide further refinement of the cell population to those geared to produce tissue in 3D.

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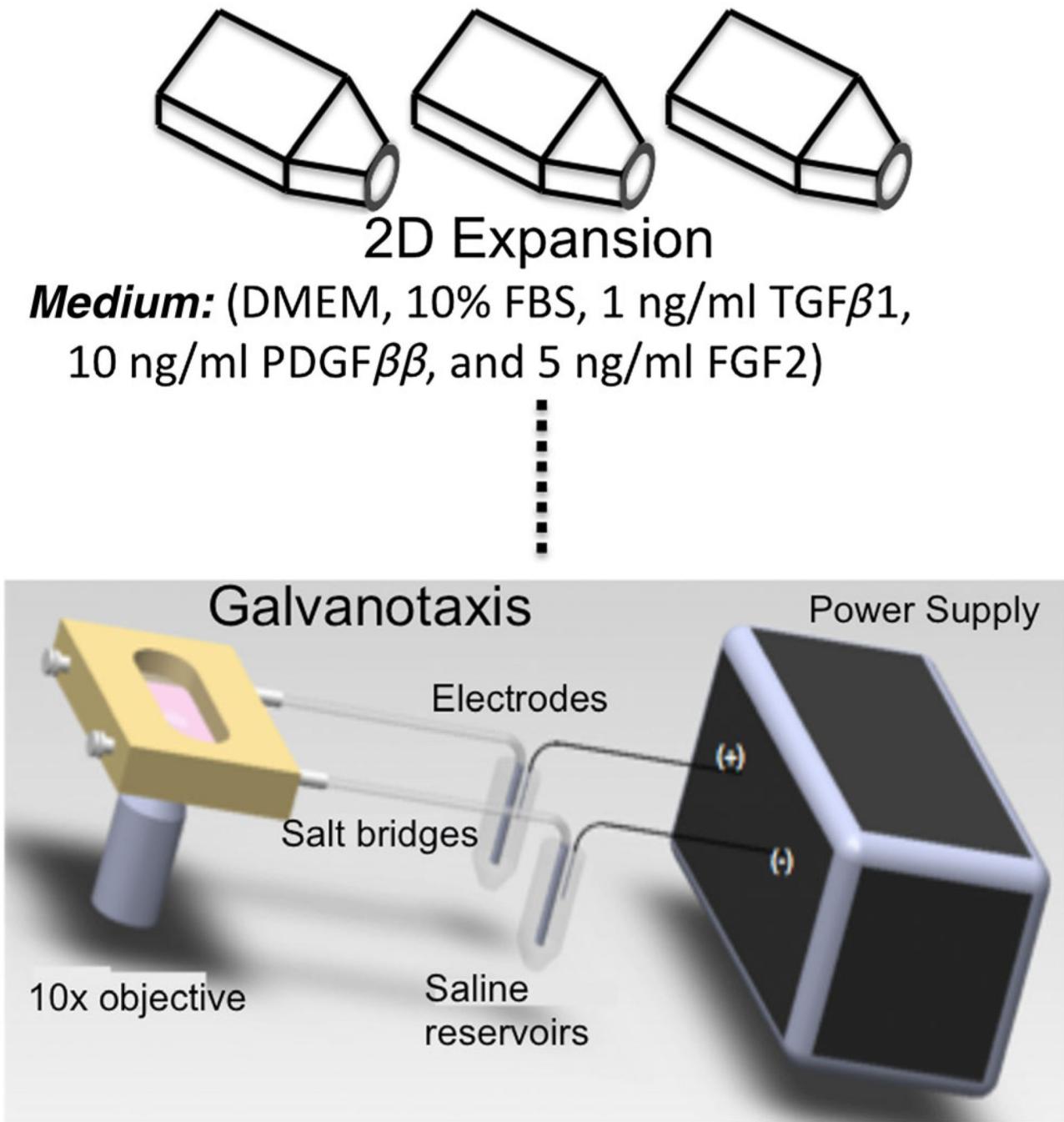


Figure 1.

Schematic of galvanotaxis studies. Human chondrocytes were cultured in expansion medium containing 1 ng/ml TGF β 1, 10 ng/ml PDGF $\beta\beta$ and 5 ng/ml FGF2 to prime the cells for 3D cultures. Galvanotaxis studies were performed for four passages

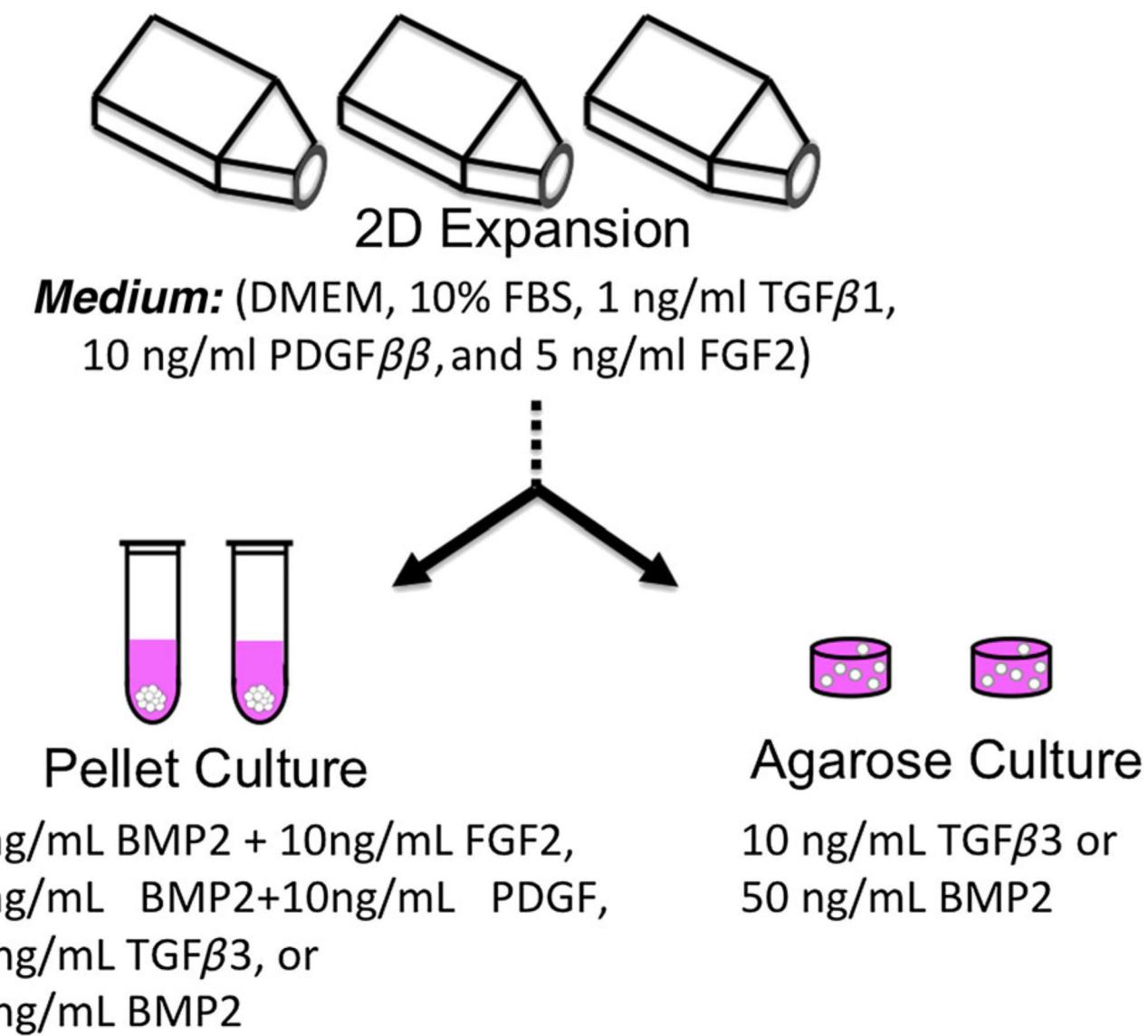
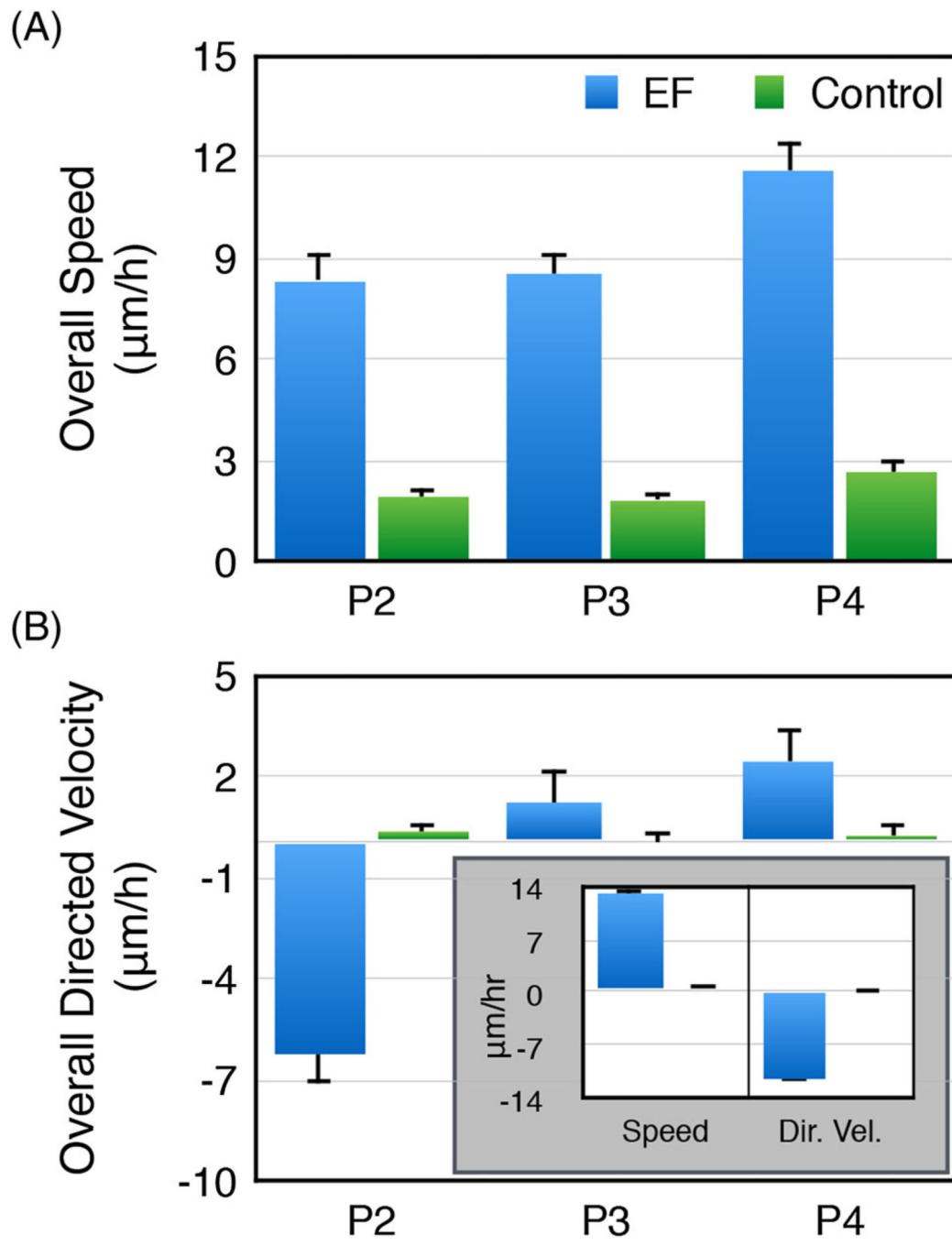


Figure 2.

Schematic of tissue-engineering study for human chondrocytes acquired from osteoarthritic joints; four growth factor combinations were evaluated for pellet cultures and agarose constructs were cultured with either BMP2 or TGF β 3

**Figure 3.**

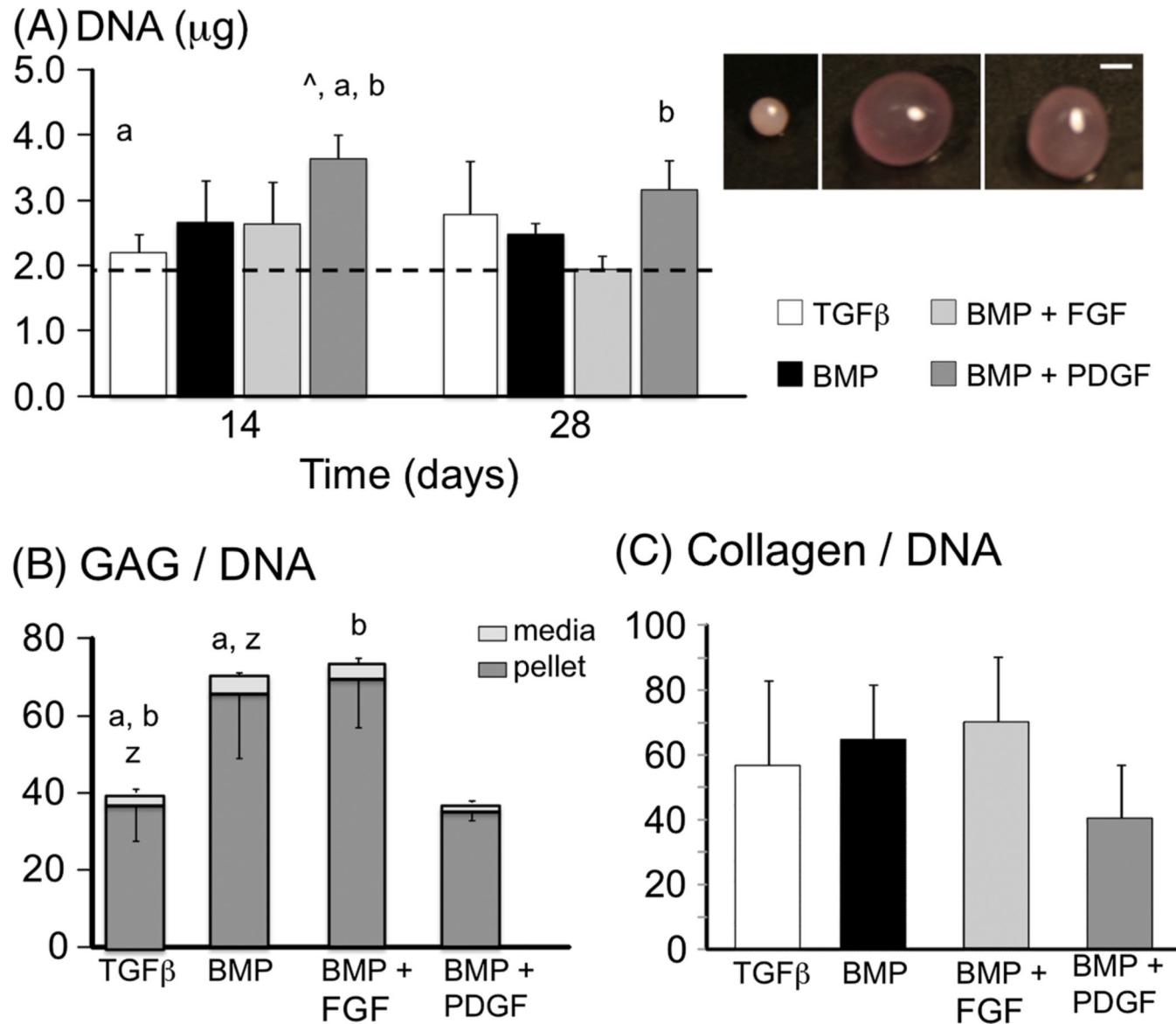
Cell migration parameters for passaged human chondrocytes: (A) overall speed for EF-treated cells increased with passage number; (B) directed velocity for those cells is orientated toward the cathode at P2 before changing direction toward the anode for later passages; (inset) speed and directed velocity of juvenile bovine chondrocytes at P2; * $p < 0.05$ vs control at each passage

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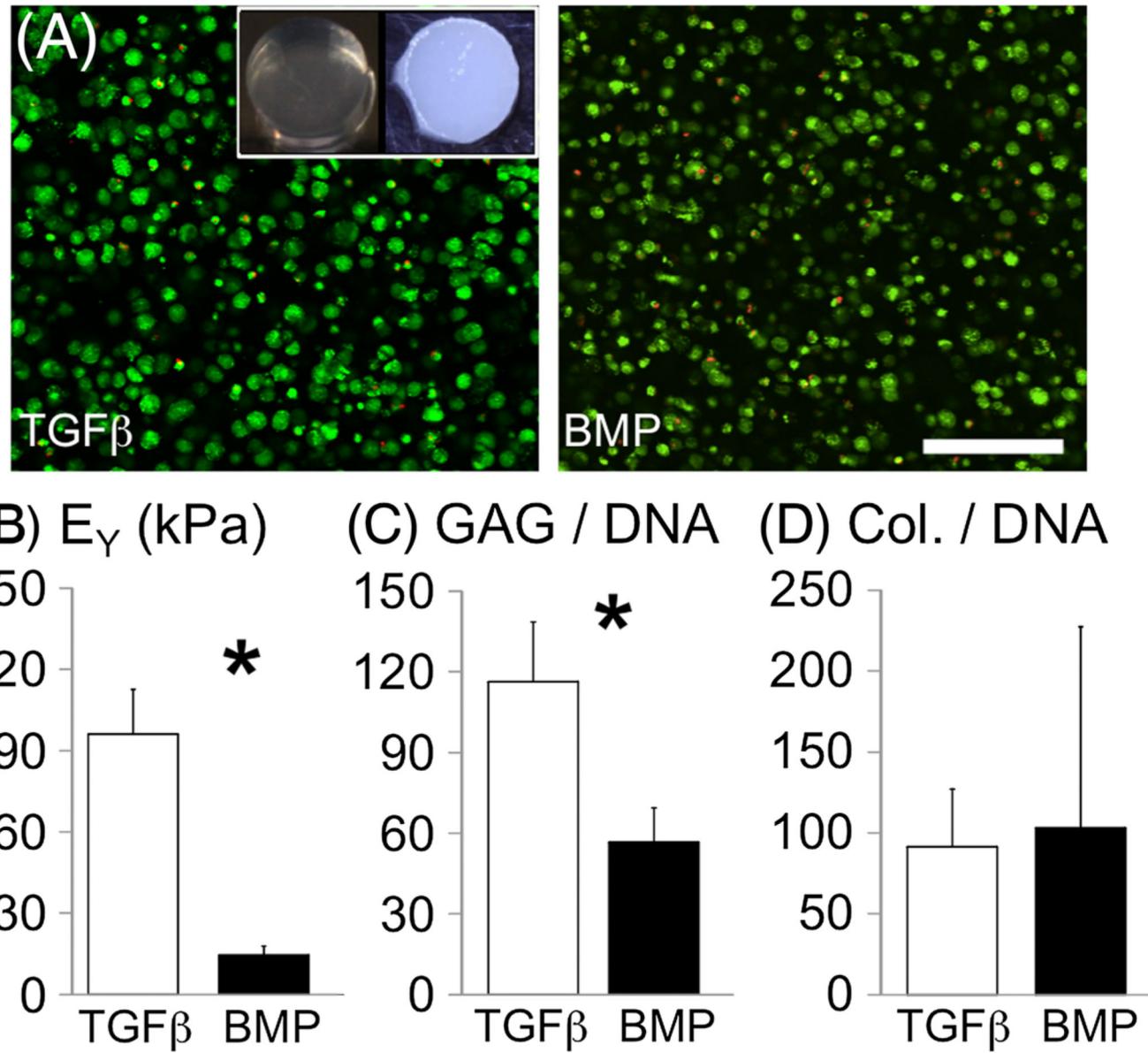
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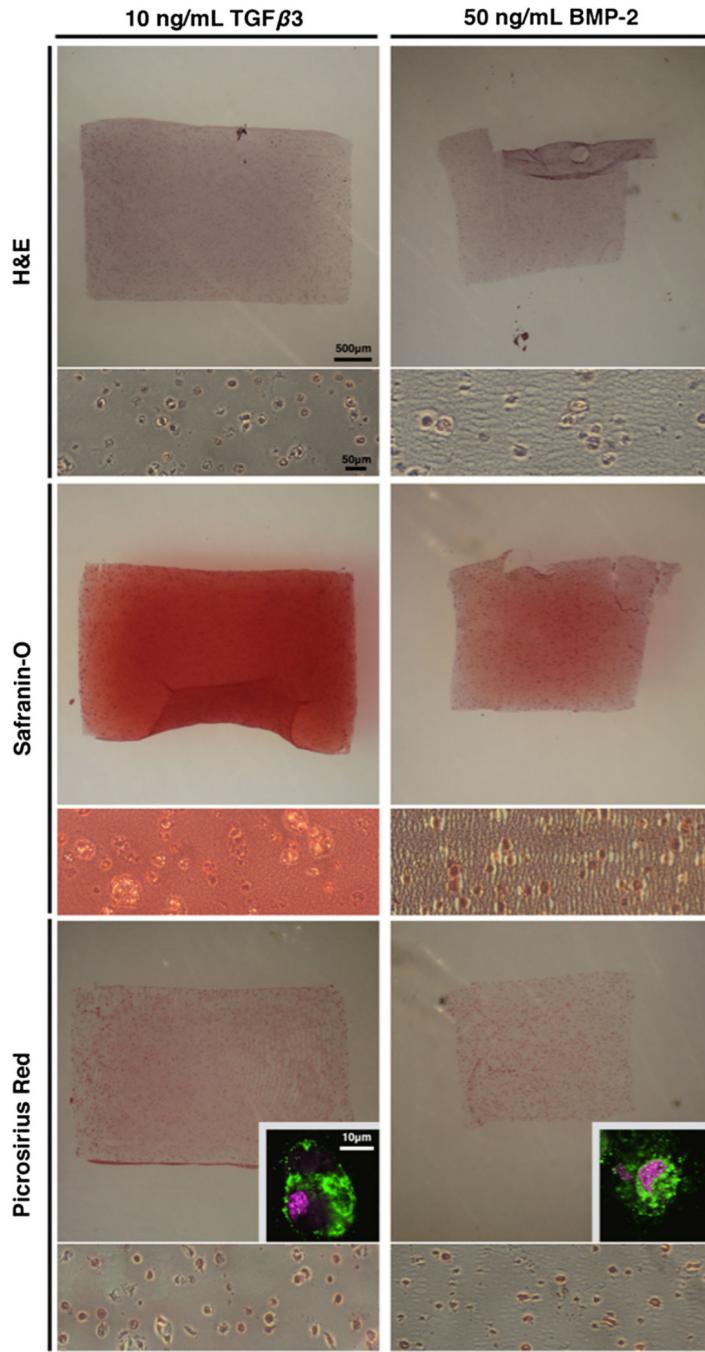
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**Figure 4.**

(A) DNA content of pellets over the 28 day culture period; dotted line represents day 0 values; groups with the same letter were found to be significantly different from one another, i.e. $p < 0.05$ a vs a); $^{\wedge}$ represents $p < 0.05$ vs day 0 values. (B) GAG content normalized by DNA content at day 28; a and b represent differences in pellet GAG content, and z represents differences in medium GAG content. (C) Collagen content normalized by DNA content at day 28; no significant differences ($p = 0.14$): (inset) representative images of pellets at day 28; pellets were cultured with TGF β 3-only (left), BMP2 with FGF (middle) or BMP2 with PDGF $\beta\beta$; note that the BMP2-only group is not shown here; bar = 1 mm

**Figure 5.**

(A) Cell viability of chondrocyte-seeded scaffolds: living cells shown in green and dead cells in red; bar = 150 μ m; (inset) representative images of a construct at day 0 (left) and a construct cultured with TGF β 3 (right). (B) Equilibrium Young's modulus, (C) GAG and (D) collagen contents normalized by dry weight for culture with TGF β 3 only or BMP2 only (day 56 values); *significant differences between groups

**Figure 6.**

Representative histology for agarose constructs cultured with TGF β 3 or BMP2. Samples were stained with haematoxylin and eosin (H&E) for cell distribution (top row), safranin-O for GAG distribution (middle row) and picrosirius red for collagen distribution (bottom row); (inset) representative image of collagen VI staining from immunohistochemistry; note that the darker region at the bottom of the safranin-O stain for the TGF β 3 group is due to a fold in the sample