

In vitro and *in vivo* performance of methacrylated gellan gum hydrogel formulations for cartilage repair*

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Abstract: Methacrylated gellan gum (GGMA) formulation is proposed as a second-generation hydrogel for controlled delivery of cartilage-forming cells into focal chondral lesions, allowing immediate *in situ* retention of cells and 3D filling of lesion volume, such approach deemed compatible with an arthroscopic procedure. Formulation optimization was carried out *in vitro* using chondrocytes and adipose mesenchymal stromal/stem cells (ASCs). A proof-of-concept *in vivo* study was conducted using a rabbit model with induced chondral lesions. Outcomes were compared with microfracture or non-treated control. Three grading scores were used to evaluate tissue repair after 8 weeks by macroscopic, histological and immunohistochemical analysis. Intense collagen type II and low collagen type I gene and protein expression were achieved *in vitro* by the

ASC + GGMA formulation, in light with development of healthy chondral tissue. *In vivo*, this formulation promoted significantly superior *de novo* cartilage formation compared with the non-treated group. Maintenance of chondral height and integration with native tissue was further accomplished. The physicochemical properties of the proposed GGMA hydrogel exhibited highly favorable characteristics and biological performance both *in vitro* and *in vivo*, positioning itself as an attractive xeno-free biomaterial to be used with chondrogenic cells for a cost-effective treatment of focal chondral lesions. © 2018 Wiley Periodicals, Inc. J Biomed Mater Res Part A: 00: 000–000, 2018

Key Words: cartilage repair, methacrylated gellan gum, hydrogel, adipose stromal/stem cells, chondrocytes

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INTRODUCTION

Complementary efforts for improving cartilage repair¹ have addressed optimization of combination strategies involving predominantly autologous articular chondrocytes^{2,3} for which decades of performance history have been collected.^{4–7} Robust long-term outcomes is among its major advantages,^{8,9} yet suboptimal cell retention within cartilage lesion sites has been a concern,^{10–12} which has driven the development and marketing of novel scaffolds or matrices to enhance efficacy of these procedures.^{12–14} Nonetheless, most surgical protocols intervene directly on the

subchondral bone either for recruitment of cells or for fixation of the scaffold, involve additional fixation systems, or require invasive procedures, such as arthrotomy, to be effectively implanted.^{15,16}

In previous works^{17–19} gellan gum (GG) has been proposed as new biomaterial for cartilage tissue engineering applications. Its versatility and efficacy has been demonstrated for cartilage repair strategies involving both subchondral stimulation and cell transplantation using a rabbit model.¹⁸ Both histological and gene expression outcomes confirmed the potential of this approach for cartilage repair

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but limitations concerning usability and crosslinking kinetics have been identified, which could limit its translation into a clinical setting. Subsequent work explored alternative synthetic routes to enhance performance of GG hydrogels, namely the methacrylation of the molecule (GGMA)^{20–22} for other biomedical applications. Rational design modification of GG yielded a second-generation GGMA polymer endowed with improved physicochemical characteristics, including better solubilization, liquid formulation prior to injection at room temperature, improved gelification kinetics, and more robust mechanical properties of the hydrogel^{20–22} the latter being greatly dependant on the crosslinking mechanism. In an applied perspective, the adoption of an injectable formulation based in GGMA in the context of cartilage repair is highly attractive, as its solution-state properties make it potentially compatible with minimally invasive procedures.

Given the positive track-record of the parent GG molecule on what regards safety and performance, it becomes mandatory to quantify the actual benefits of GGMA in the cartilage repair application context. Understanding physicochemical performance of GGMA could be explored to simplify the surgical protocol, to improve delivery and functional commitment of cells employed, as well as to minimize damage of the subchondral compartment during the surgical protocol. In this regard, this study aims to comparatively assess the safety and performance of GG and GGMA by *in vitro* methods, as well as to characterize the performance of GGMA hydrogel as vehicle for delivery and retention of chondrogenic cells within chondral lesions, by assessing functional development of hyaline cartilage tissue in a rabbit model.

On the perspective of functional performance of cells, the risk of chondrocyte de-differentiation,²³ or lack of potency of the autologous chondrocytes^{24–26} along the need for double surgery and prolonged surgical pre-planning, has inspired the study of alternative cell sources, including mesenchymal stromal/stem cells (MSCs) in general, and adipose-derived stromal/stem cells (ASCs) in particular.^{17,18,27} As compared with other MSC sources, adipose tissue can be harvested with reduced morbidity at the donor site and yields of ASC are considerably high.²⁸ The immunomodulatory and anti-inflammatory properties of ASC makes them an especially attractive cell source for development of off the shelf regenerative medicine treatments.^{27,29,30}

Herein, preliminary screening demonstrated improved cell viability of ASC within ionic-crosslinked GGMA as compared with photo-crosslinked, therefore favoring further experimentation with ionically crosslinked GGMA. The best performing combination was further evaluated for the treatment of focal chondral lesions in a rabbit model, by adopting a physiologically inspired crosslinking approach devoid of toxic photo-initiators and electromagnetic radiation sources, which is highly desirable from both regulatory and surgical protocol perspectives.

MATERIALS AND METHODS

In vitro chondrogenesis

Preparation of purified GG and GGMA. Commercial GG (GGc) (Gelzan, Sigma-Aldrich) was purified according to the

method described by Doner³¹ with several modifications. Briefly, GGc was suspended in distilled water (1% w/V) and warmed to 60°C with stirring. To this solution was added Amberlite IR-120 (H⁺ form) (Sigma-Aldrich) until pH 2.5. The suspension was filtered and aqueous sodium hydroxide (NaOH, 1 N) was added until pH 8, while stirring. The resulting solution was filtered and the filtrate poured onto absolute ethanol (1 L), forming a thick fibrous precipitate. After 1 h, the precipitate was filtered, washed with absolute ethanol and dissolved in distilled water. The resulting solution was transferred to a cellulose membrane (Molecular weight cut-off (MWCO) 12 kDa) and dialyzed against distilled water for 3 days. After freezing (–20°C) and lyophilization, the purified GG (GGp) was obtained. GGMA, with a degree of substitution with methacrylate groups between 1.5 and 5% was prepared as follows: GGc was dissolved in water to give a solution of 1% w/V concentration. Heating was stopped and the solution pH was adjusted to 8.5 by NaOH (1 N). Thereupon, excess glycidyl methacrylate was added in one portion and the methacrylation reaction was allowed to proceed for 24 h whilst maintaining the solution pH close to 8.5. Acetone was then added to the reaction mixture which was allowed to stand for 2 h. The precipitate was recovered by filtration, dissolved in distilled water and then placed in a cellulose dialysis membrane (MWCO 12 kDa) and dialyzed against distilled water for 7 days. The dialyzed solution was then frozen at –20°C and subsequently freeze-dried to give GGMA as an amorphous white solid. For hydrogel preparation, GGp and GGMA powder were dissolved in deionized water to achieve solutions at 1.25 and 2.5% w/V, respectively. Dissolution was effected at 37°C in a water-bath with 100 rpm agitation.

***In vitro* culture of human cells.** Human nasal cartilage (hNC) was obtained with informed consent, as surgical waste from a local hospital and further processed for isolation of chondrocytes as described elsewhere.¹⁷ Chondrocytes were thawed and expanded in DMEM:F12 supplemented with 10% v/v FBS and 1% v/v antibiotic-antimycotic (Gibco, USA) until passage 3. Human adipose tissue (hAT) was obtained from liposuction procedures, after informed consent and medical questionnaire according to European directives. Collection of adipose samples was approved by the Institutional Review Board of “Centro Hospitalar de São João”, Portugal. Briefly, hAT was washed with a decontamination solution (Base-128 Alchimia, Italy) and digested with collagenase (0.4 U/mL, NB6, SERVA, Germany) for 1 h at 37°C with agitation. The stromal vascular fraction (SVF) was collected after purification steps that include washing, centrifugation and lysis of red blood cells. Human adipose mesenchymal stromal/stem cells (hASC) were obtained from SVF by plating and further expansion in low serum media (MesenPro, Gibco, USA) or xeno-free media (Fibrolife, Lifeline, USA) until passage 2 or 4. Quality control included validation of MSC immunophenotype (CD31, CD34, CD45, CD73, CD90, and CD105, BD Biosciences, USA) characterized by flow cytometry analysis (FACS Canto, FACSDiva software, antibodies BD Biosciences, USA), and trilineage

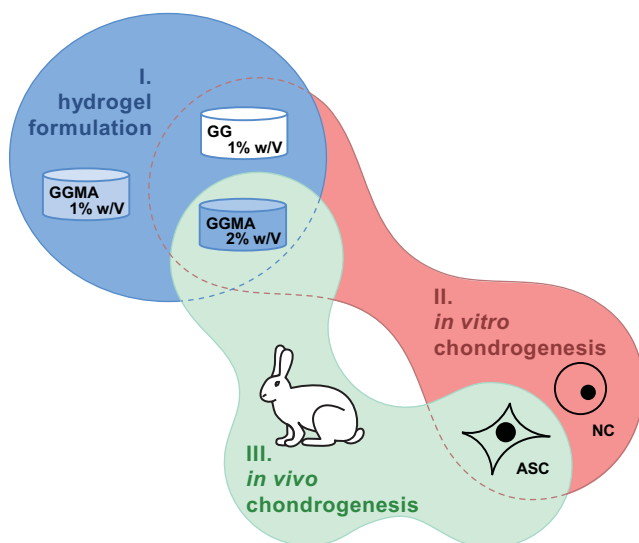


FIGURE 1. Schematic representation of experimental design. Hydrogel formulations based on GG and its methacrylated derivative (GGMA) were tested for gelification and cell encapsulation (stage I). Two formulations were selected for *in vitro* assessment of chondrogenesis (stage II) using nasal chondrocytes (NC) and ASC. A final formulation was applied for treatment of focal chondral lesions in an induced rabbit model (stage III).

differentiation (StemPro, Gibco, USA) identified by alizarin red, oil red O and alcian blue stainings for osteogenesis, adipogenesis and chondrogenesis, respectively.

Preparation of cell-encapsulated hydrogels for *in vitro* culture. An initial comparison study was performed to evaluate comparative performance of GG and GGMA with respect to hydrogel formation and metabolic activity of encapsulated hASC. Selected formulations were further used for assessment of chondrogenesis by hASC or hNC (Fig. 1). Cell suspensions were prepared in cell culture media and mixed with GG or GGMA solution in a 2:8 ratio in order to yield a final cell density of 5×10^6 cells/mL and hydrogel concentrations of 1 and 2% w/V, respectively. Cellular hydrogels of 20 μ L volume were pipetted (with aid of a positive displacement pipette) into wells of non-adherent cell culture well-plates and covered with culture media for crosslinking. To induce chondrogenesis of hASC, serum-free chondrogenic media (StemPro, Gibco, USA) was used. At specific time-points (i.e., 0, 7, and 21 days), individual hydrogels were collected for analysis. The hASC–GGMA combination was further scaled to hydrogels of 50 μ L volume containing 10×10^6 cells/mL and tested for chondrogenesis.

Assessment of *in vitro* cell viability and chondrogenesis. Cell metabolic activity was determined at each time-point by MTS assay (Promega USA) and cell viability was further microscopically assessed by Live/Dead assay (calcein AM and propidium iodide [Invitrogen, USA 1 mg/mL]). For histology and immunohistochemistry (IHC), hydrogels were fixed (10% formalin), followed paraffin processing. Histochemical staining of glycosaminoglycans (GAGs) by Safranin O/Fast green and Alcian Blue were performed as

previously described in.¹⁷ For IHC, reagents from Vector Laboratories (UK) were used. Sections were incubated into recommended antigen retrieval solutions, followed by inhibition of endogenous peroxidases with 0.3% H_2O_2 and blocking with normal horse serum. Thereafter, sections were stained with primary antibody Mouse anti-human Anti-Collagen I or Mouse anti-human Anti-Collagen II (Abcam, UK) for 1 h, RT and a diluted biotinylated secondary antibody solution (VECTASTAIN Elite ABC kit) for 30 min, RT. Signal development was performed with the DAB substrate kit. For gene expression analysis, hydrogels were collected into TRI Reagent (Sigma-Aldrich, USA) and recommended protocol for RNA extraction from tissues was followed. Complementary cDNA was obtained by using the High Capacity cDNA Reverse Transcription Kit. Gene amplification was conducted using TaqMan Fast Advanced Master Mix and TaqMan Gene Expression Assays for Collagen type II (Hs00264051_m1) and Collagen type I (Hs00164004_m1). Glyceraldehyde 3-phosphate dehydrogenase (GAPDH, Hs99999905_m1) was chosen as an invariant standard (housekeeping gene). Quantitative reverse transcription (RT-qPCR) analysis was carried out with the StepOnePlus Real-Time PCR System and software (all reagents and equipment from Applied Biosystems, USA). Results were normalized to GAPDH and expressed as relative gene expression using the $\Delta\Delta C_t$ method. The expression data were presented as average values for each group ($n = 3 \pm SD$).

***In vivo* chondrogenesis**

Chondral lesion induction and repair in a rabbit model.

The ICRS and ASTM guidelines were followed for a proof-of-concept (PoC) study in rabbits to assess *in vivo* cartilage tissue repair.^{32,33} All animal procedures were based upon the “3R’s” policy, approved by the Institutional Ethical Committee, according to the National authority Guide for the Care and Use of Laboratory Animals. Skeletally mature (12- to 14-weeks old) New Zealand white rabbits (2.5 ± 0.25 kg; Charles-River, France, $n = 6$) were used for harvesting adipose tissue and subsequent autologous treatment of focal chondral lesions. Interscapular adipose tissue (~ 10 g) was collected under anesthesia with a mixture of ketamine hydrochloride (Imalgene, 25 mg/kg i.m.) and medetomidine hydrochloride (Domitor, 0.3 mg/kg i.m.). The obtained adipose tissue samples were digested for 1 h at 37°C, 100 rpm with Collagenase NB4 Standard Grade 0.2 U/mL (Serva, Germany). After complete digestion, cells were cultured in complete media based on alpha MEM supplemented with 10% v/v FBS and 1% v/v antibiotic-antimycotic (Gibco, USA), until passage 2. One week after adipose tissue harvest, surgery was conducted to create critical chondral defects in the knee for immediate administration of treatment. Rabbits were anesthetized as described above and both knees were shaved and disinfected. An internal para-patellar incision was made to expose the knee. The patella was dislocated and two 4-mm diameter lesions were made in the trochlear groove of each knee using a biopsy punch. Lesion sites were carefully cleaned with a curette to not affect the sub-chondral

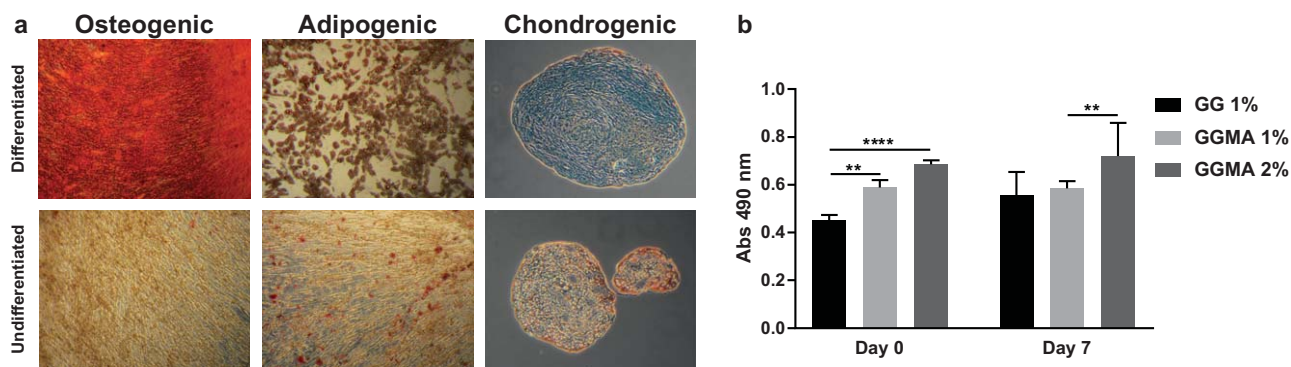


FIGURE 2. Preliminary *in vitro* studies with hASC. (a) Trilineage differentiation of hASC identified by alizarin red, oil red O and alcian blue stainings for osteogenesis, adipogenesis, and chondrogenesis, respectively. (b) Metabolic activity of hASC encapsulated in GG and GGMA hydrogel formulations (1 and 2% w/v) upon encapsulation (day 0) and after 1 week *in vitro* culture. ** $p < 0.01$, **** $p < 0.0001$.

bone. Defects were randomly allocated to one of the following experimental conditions: (1) rabbit autologous ASC encapsulated in GGMA hydrogel (GGMA + rASC); (2) microfracture (MFX) (positive control) and (iii) empty lesion (negative control). Autologous rabbit ASC were encapsulated in GGMA hydrogels as described earlier (10×10^6 cells/mL, 2% w/v) immediately before delivery into the chondral defect. In situ crosslinking was promoted with PBS and a setting time of 10 min was allowed before closure. MFXs were made with a 0.8-mm Kirshner wire (six holes per defect) with 1- to 2-mm depth from which bleeding was observed. Finally, the patella was reduced and the wound was closed. After recovery from surgery, animals were placed in individual cages and fed *ad libitum*.

Assessment of *in vivo* cartilage tissue repair. Cartilage regeneration was allowed for 8 weeks, after which animals were anesthetized as described earlier and euthanized (Eutasil, 200 mg/kg). In each knee, an internal para-patellar incision was made and the patella carefully dislocated. Macroscopic pictures were taken and explant tissue was harvested with a 6-mm diameter punch in order to collect native tissue surrounding the lesion site, as well as subchondral bone. Explants were paraffin-processed after fixation (10% formalin) and decalcification (Biodec R, Bio-Optica, Italy). For IHC, sections were processed as described above, followed by incubation with primary antibody mouse anti-rabbit anti-collagen I (Abcam, UK) or mouse anti-rabbit anti-collagen II (Merck Millipore, USA) for 1 h, RT. Histochemical staining of GAGs was performed by Safranin O/fast green and three scoring systems were used to assess the quality of cartilage repair, namely O'Driscoll, Pineda and Wakitani.³⁴

Statistical analysis

Results are summarized by mean or median and corresponding standard deviation or interquartile range. For *in vitro* studies, Student's *t* test and two-way analysis of variance (ANOVA) were used to evaluate differences among groups. Normality was evaluated by the Shapiro-Wilk test.

When normality or homogeneity of variances was not verified, non-parametric tests were used. For *in vivo* studies, the histological scores for each specimen were evaluated independently by three observers at three different times. For evaluation, the observers were blinded for the type of treatment and the specimens were randomly allocated to each observer. The comparisons between treatment groups were performed by two-way ANOVA. Since there were no statistical differences between the observers' evaluations, the results were analyzed using one-way ANOVA. In cases where homogeneity of variances was not observed, the Kruskal-Wallis nonparametric test was adopted. Multiple comparisons were based on the Tukey HSD test or the Mann-Whitney test, with the corresponding significance level and Bonferroni correction. Statistical analysis was performed using the GraphPad Prism 4.0c software or IBM SPSS Statistics, version 23. Statistical significance was defined for $p < 0.05$.

RESULTS

In vitro chondrogenesis

Trilineage differentiation capacity was confirmed for CD73+/CD90+/CD105+/CD31-/CD34-/CD45- hASC, with expressive mineralization, lipid formation, and GAG deposition upon osteogenesis, adipogenesis and chondrogenesis, respectively [Fig. 2(a)]. Upon encapsulation within GG-based hydrogel formulations, higher metabolic activity was observed for hASC encapsulated in GGMA 1 and 2% w/v as compared with GG at 1% w/v ($p < 0.01$ and 0.001, respectively) [Fig. 2(b)]. GG at 2% w/v provided inadequate sol-gel transition time for cell encapsulation studies. After 7 days of culture, highest metabolic activity was observed by cells encapsulated in GGMA 2% w/v ($p < 0.01$) therefore this formulation was selected for further *in vitro* cell encapsulation studies concurrently with GG 1% w/v (Fig. 3).

After 21 days of *in vitro* culture, the viability of chondrocytes and hASC, assessed microscopically by live/dead assay, was comparable between GGp and GGMA hydrogels [Fig. 3(a), top and middle rows]. On the other hand, hASC chondrogenically differentiated within both hydrogel groups

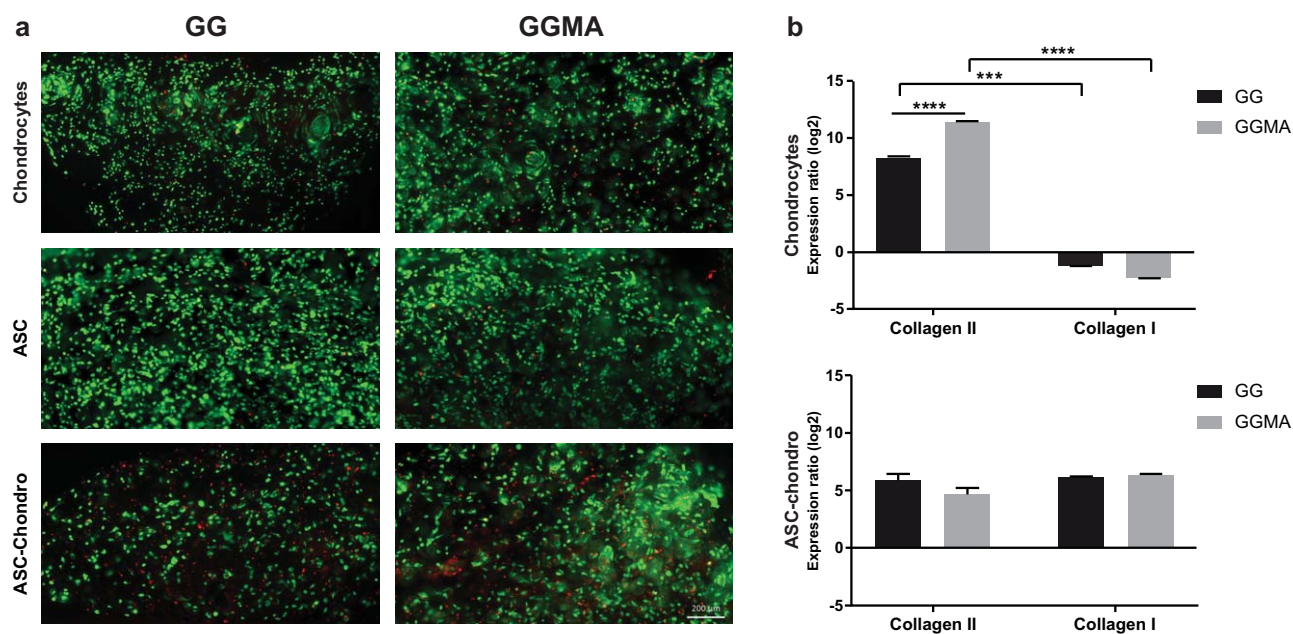


FIGURE 3. *In vitro* chondrogenesis. (a) Microscopic imaging of encapsulated cells stained by calcein AM (live, green) and propidium iodide (dead, red) upon 21 days *in vitro* culture in 1% w/v GG and 2% w/v GGMA hydrogels. (b) Normalized gene expression ratio (days 21–0) of GG/GGMA encapsulated chondrocytes (top) and chondrogenically induced hASC (bottom). *** $p < 0.001$, **** $p < 0.0001$.

(hASC-chondro) demonstrated increased viability within the GGMA hydrogel [Fig. 3(a), bottom row]. On what regards expression of chondrogenic markers, both formulations favored maintenance of healthy chondrocytes as evidenced by significantly increased expression of collagen type II relative to collagen type I. Such response was superior by the GGMA formulation ($p < 0.0001$) as compared with the GG ($p < 0.001$). Furthermore, chondrocytes cultured within the GGMA hydrogel presented higher collagen type II expression ratio as compared with the parent GGp hydrogel [Fig. 3(b), top] ($p < 0.0001$). On what regards hASC [Fig. 3(b), bottom], such cells effectively expressed collagen type II upon chondrogenic stimuli (21 days) when cultured within either hydrogel formulation ($p > 0.05$). Concurrently, collagen type I expression was not superior to collagen type II at this time-point ($p > 0.05$). When doubling hASC concentration within the GGMA hydrogel up to 10×10^6 cells/mL, increasing expression of collagen type II was obtained in the course of chondrogenic differentiation [Fig. 4(a)]. Simultaneously, very low expression of collagen type I was obtained along culture ($p < 0.001$ and 0.05 at days 14 and 21, respectively). Samples were further collected for histological analysis and subsequent identification of extracellular matrix (ECM) components [Fig. 4(b)]. Progressive deposition of healthy chondrogenic ECM was observed as evidenced by safranin O/fast green staining of cartilage matrix, alcian blue detection of sulfated GAG and IHC of human collagen type II. The absence of collagen type I deposition also indicates development of non-fibrous cartilaginous tissue. Macroscopically, transparent hydrogels at the beginning of culture showed reduction in transparency (not quantified) into an off-white opaque appearance after 3 weeks *in vitro* culture [Fig. 4(b), bottom row].

***In vivo* cartilage repair**

At surgical treatment day, expanded rASC were mixed with GGMA solution at time of surgery so as to form a homogeneous suspension. Upon injection into the lesion, the viscosity of the suspension allowed spatial control of delivery within the lesion volume, without spill over at the edges of the defect. Gelification was allowed to occur during 10 min as to assure maintenance of the hydrogel in the lesion site, allowing immediate retention of cells *in situ*. Rabbits remained healthy during all experimentation period, presenting normal weight gain and absence of signs of infection or disease. From macroscopic observation at time of explant surgery, no apparent abnormalities of the patella position were observed; neither signs of inflammation, abnormalities of the synovium, loose bodies, osteophytes or degenerative process were found. Macroscopic observation showed native cartilage near the defect site as well as the opposing cartilage to be bright and white without visible degenerative signs. In all defects, tissue formation was observed (Fig. 5), and the margin between the defect and the surrounding cartilage were visible, which was more evident for the empty control group. The defects treated with the GGMA + rASC combination showed compact bright tissue filling, despite macroscopic variability observed between defects. The lesions treated by MFX presented an irregular filling of the lesion site, with tissue of a dim appearance. A similar outcome was observed for untreated lesions (empty defects). Tissue explants were further harvested for histological analysis. The quality of cartilage repair was assessed by three scoring systems (Fig. 5), which have inverse scales for indication of cartilage quality and outcome: according to O'Driscoll, high point values indicate enhanced cartilage while according to Pineda and Wakitani, low total point values

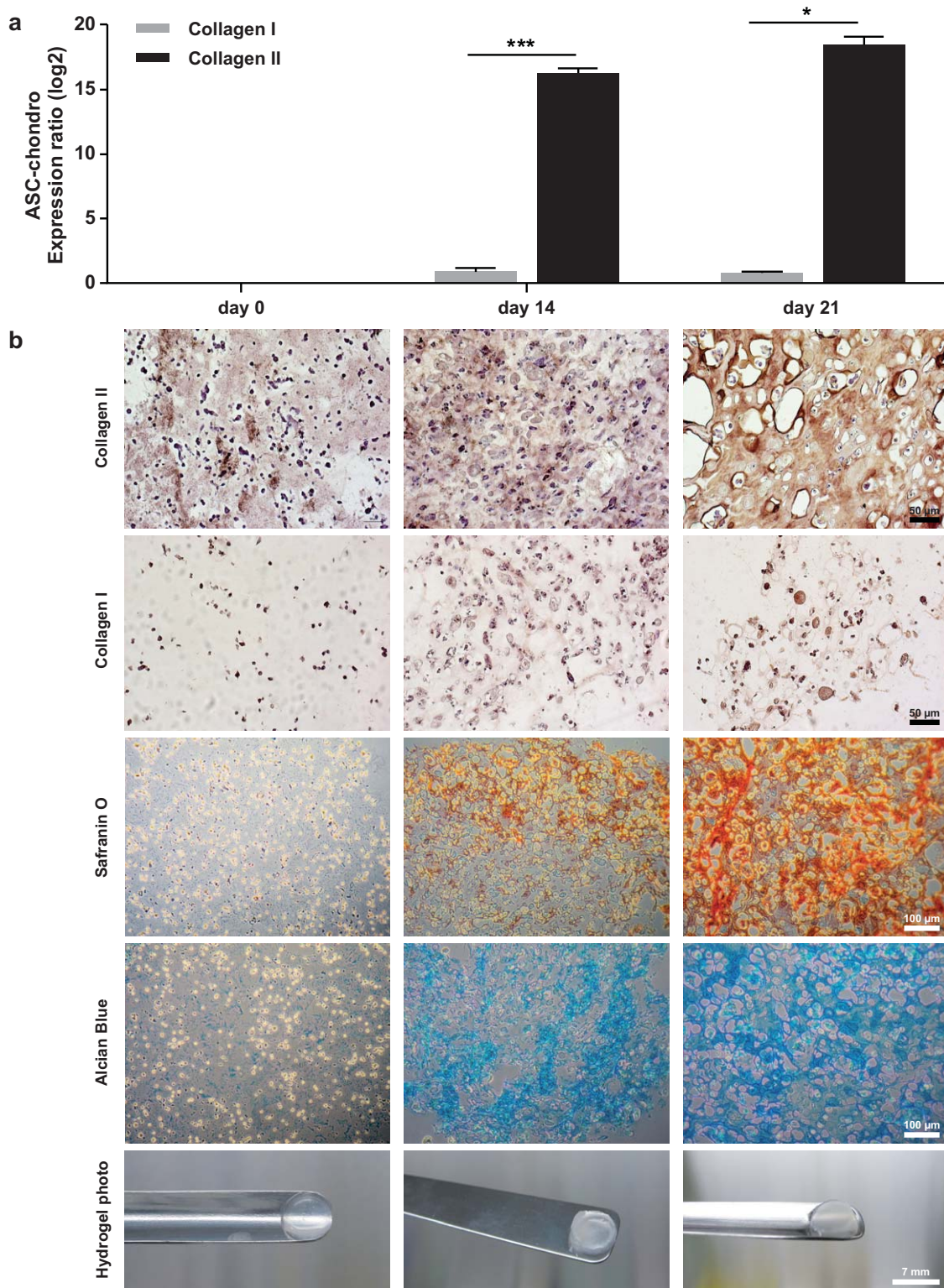


FIGURE 4. *In vitro* chondrogenesis of hASC encapsulated in GGMA 2% w/V. (a) Gene expression ratio normalized to day 0. * $p < 0.05$, *** $p < 0.001$. (b) Histological analysis and macroscopic imaging of hydrogel along *in vitro* culture.

represents superior repair. Inter observer differences were assessed and no statistical differences were observed. Immunolocalization of rabbit collagen type II and collagen type I (Fig. 5) was performed to further characterize

cartilage formed within the lesion site. Treatment of chondral lesions with GGMA + rASC (Fig. 5, top) allowed restoration of cartilage thickness, integration/bonding with native cartilage, as well as intense and reasonably homogenous

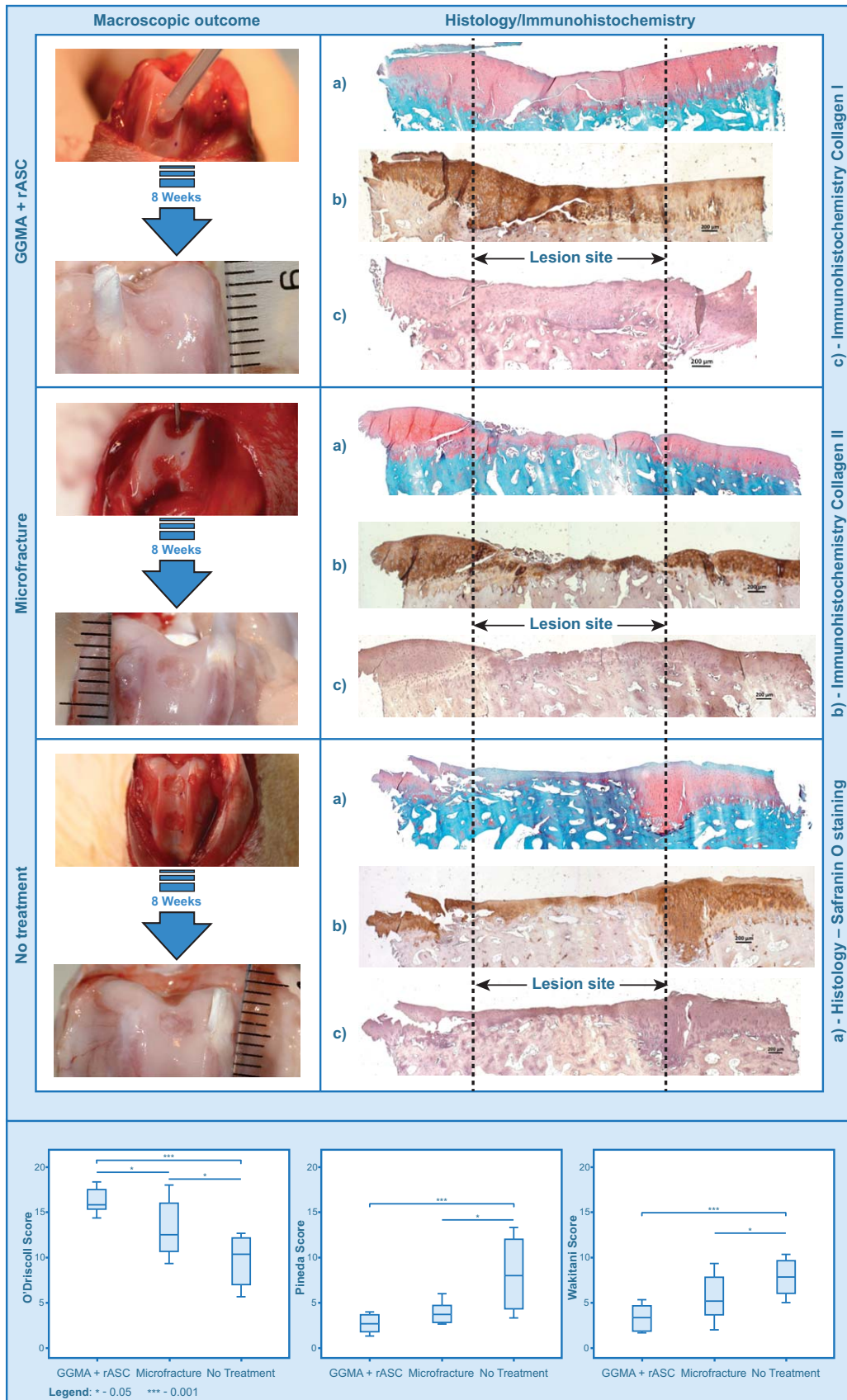


FIGURE 5. *In vivo* chondrogenesis of hASC encapsulated in GGMA 2% w/v. Top panel. Histological analysis and macroscopic imaging of experimental groups after 8 weeks of implantation. Bottom panel. Histological scoring according to O’Driscoll, Pineda, and Wakitani scores. * $p < 0.05$, *** $p < 0.001$.

staining of ECM throughout the lesion site. Quantitative assessment of repair by all three scoring systems indicates significant improvement in cartilage repair as compared with the untreated lesions ($p < 0.001$). According to the O'Driscoll score, GGMA + rASC treatment also outperformed MFX ($p < 0.05$). Lesions treated with MFX (Fig. 5, Middle) demonstrated overgrowth of the subchondral bone into the lesion site which was covered by a thin layer of cartilaginous matrix stained by safranin O and collagen type II. This layer is irregular and bonding with adjacent native cartilage is incomplete. Nevertheless, the extent of cartilage repair by MFX was superior to untreated lesions, independently of the scoring system used ($p < 0.05$). The bottom image represents histological assessment of untreated lesions, whereas overgrowth of subchondral bone was evident and covered by a thin regular tissue. Herein, very limited cartilaginous matrix was formed, as indicated by the lack of safranin O/fast green and reduced collagen type II staining at the top layer of the tissue. Collagen type I deposition was negligible in all groups, yet expressed slightly higher in the untreated defects.

DISCUSSION

GG polysaccharide offers attractive features and characteristics for this particular application due to its aqueous solubility and viscous properties at physiological temperature, which makes it appealing for implementation of cell combination and surgical implantation protocols. In addition, the crosslinking by physiological ions leads to formation of a stable 3D structure and subsequent cell retention. In this regard, functionalization of the GG molecule by methacrylation extends solubility, allows control of spatiotemporal crosslinking, which combined extends flexibility of combination and implantation procedures and fine-tuning of hydrogel stiffness. For instance, at 2% w/V, GGMA presents increased storage modulus as compared with the unmodified polysaccharide, 89.5 ± 7.4 and 56.2 ± 1.4 kPa, respectively.²² Matrix stiffness, as a result of increased concentration or biochemical cues, has been reported to influence stem cell fate and particularly chondrogenesis.³⁵ Within the context of this study, cells cultured within the GGMA 2% w/V hydrogel shown improved cell metabolic activity, viability, and healthy expression of chondrogenic markers as compared with the least concentrated (1% w/V) or non-functionalized matrix.

PoC in the rabbit model allowed evaluation of the cartilage repair potential of the GGMA 2% w/V hydrogel as compared with MFX treatment. Currently considered as a gold-standard treatment, MFX still has limitations on what concerns the quality of the regenerated tissue, which can ultimately lead to treatment failure upon recurrence of symptoms.^{36,37} In this study, MFX group outcome showed formation of a thin layer of chondral tissue concurrent with subchondral bone overgrowth and an irregular surface. Such repair outcome is likely to disfavor adequate load bearing as well as smooth, pain-free joint motion.^{14,38,39} A different outcome was observed for lesions treated with

GGMA-rASC—cartilage thickness was maintained equivalently to adjacent native tissue while avoiding bone overgrowth. A smooth chondral surface was obtained following this treatment and ECM staining demonstrated reasonably uniform distribution of collagen type II and GAGs (Fig. 5). This repair outcome is believed to support long-term quality of the tissue as opposed to MFX treatment. At the 8-week time-point, statistical differences were obtained between these groups according to O'Driscoll scoring. Nevertheless, it is important to mention that quantitative scores with a broad numerical range such as the O'Driscoll system may increase the likelihood of finding statistically significant differences.³⁴ Still, the adoption of O'Driscoll score in the context of this study is pertinent as, contrary to alternative scores, it assesses integration of the repair tissue with its surroundings.³⁴ In this study, no additional fixation technique was used to retain the hydrogel within the lesion site and precise volume filling was achieved, avoiding the need for on-site shaping of the scaffolding structure, which could be an advantage as compared with other cell-based and tissue engineered cartilage products currently in clinical development.¹² In addition, viscous and sol-gel transition properties of the tested GGMA 2% w/V hydrogel allowed controlled delivery of the matrix containing autologous ASC,¹⁴ directly to lesion site, which favored delivery and retention of cells *in situ*. This fact is of significant importance as cell retention at lesion site is one of the main indicators of success for lesion repair.^{12,40} Adoption of the rabbit animal model for cartilage repair studies has significant advantages due to availability, ease of handling, low cost and abundance of comparative literature,^{14,41} but poses challenges related to reduced thickness (0.4 ± 0.1 mm in the trochlear groove) and surface area of articular cartilage. This model is adequate for PoC studies during early stage development of new technologies, particularly on what concerns evaluation of fixation of implantable devices.^{32,33} In this regard, the fixation merits of GGMA-rASC combination was demonstrated in a particularly challenging environment. Heterogeneous outcomes were naturally observed yet *bona fide* chondral repair was obtained with this treatment group, which is attributed to GGMA-rASC combination alone. The analysis of non-treated lesions demonstrated a limited self-healing of the induced defect that was statistically inferior than GGMA-rASC treatment. The self-repair ability of cartilage lesions, which is commonly reported in the rabbit model^{42,43} has been minimized by implementation of chondral defects with critical-size for which penetration and damage to the subchondral bone plate was avoided.

The GGMA formulation successfully supported *in vitro* chondrogenesis of both mature and progenitor cartilage-forming cells. In a rabbit model, controlled delivery of cells into chondral lesions was achieved, while adequate spatiotemporal crosslinking supported volumetric filling of cartilage lesions and *in situ* retention of cells. Following 8 weeks of treatment, the combination of GGMA-rASC, supported full thickness regeneration of critical size lesions, good integration/bonding with native cartilage. Such combination

therapy exhibited highly favorable physicochemical characteristics and good biological performance which may support less invasive and complex surgical procedures for cartilage repair.

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