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Sequential sequestrations increase the incorporation and retention of multiple growth factors in mineralized collagen scaffolds†

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Trauma induced injuries of the mouth, jaw, face, and related structures present unique clinical challenges due to their large size and complex geometry. Growth factor signaling coordinates the behavior of multiple cell types following an injury, and effective coordination of growth factor availability within a biomaterial can be critical for accelerating bone healing. Mineralized collagen scaffolds are a class of degradable biomaterial whose biophysical and compositional parameters can be adjusted to facilitate cell invasion and tissue remodeling. Here we describe the use of modified simulated body fluid treatments to enable sequential sequestration of bone morphogenetic protein 2 and vascular endothelial growth factor into mineralized collagen scaffolds for bone repair. We report the capability of these scaffolds to sequester 60–90% of growth factor from solution without additional crosslinking treatments and show high levels of retention for individual (>94%) and multiple growth factors (>88%) that can be layered into the material via sequential sequestration steps. Sequentially sequestering growth factors allows prolonged release of growth factors *in vitro* (>94%) and suggests the potential to improve healing of large-scale bone injury models *in vivo*. Future work will utilize this sequestration method to induce cellular activities critical to bone healing such as vessel formation and cell migration.

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1. Introduction

Craniofacial injuries – injuries of the mouth, jaw, face, and related structures – can be caused by a wide range of congenital abnormalities, oral cancer treatments, and traumatic injuries. Trauma related injuries experienced by civilians^{1–5} and high-energy impact injuries experienced by warfighters^{6–8} present unique clinical challenges. These injuries are often large, complex in geometry, and cannot be repaired with external fixtures alone.^{9,10} Most clinical treatments use autografts, bone taken from a secondary site in the patient with the injury, or allografts, bone taken from a human donor.^{11–13} While autografts are considered the gold-standard and maintain osteo-conductive and osteo-inductive abilities, allografts are a popular alternative because of their availability.^{14,15} However, there are limitations to the use of these bone grafts. Autografts are limited by the size of the injury site, and

allografts raise concerns about disease transmission, transplant rejection, and their purification methods are not uniform, resulting in variability between grafts.^{16–18} Thus, there is a clinical need for alternative solutions to address critically sized craniofacial injuries.

Regenerative medicine solutions commonly seek to combine a biomaterial template with strategies to accelerate healing such as the incorporation of biomolecule stimuli.^{19,20} Bone healing is a multistep process with multiple cell types and is coordinated through growth factor signaling.^{21–25} Briefly, inflammatory cytokines such as tumor necrosis factor alpha are released following injury and lead to the formation of a hematoma.^{26,27} These cytokines recruit macrophages and other immune cells to the injured site and initiate new vessel formation.²⁷ Vascular endothelial growth factor (VEGF) and angiopoietin-1 and 2 are critical growth factors for promoting and maintaining angiogenesis.^{26,28–30} Immune cells then release factors such as bone morphogenetic protein 7 to recruit mesenchymal stem cells to the injury site.³¹ Mesenchymal stem cells start depositing extracellular matrix components such as collagen that lead to callous formation at the injured site.^{22,24} Bone morphogenetic protein 2 (BMP2) and other signaling molecules induce the differentiation of mesenchymal stem cells into osteoblasts (bone depositing cells) and mineralization of the callous occurs.^{31,32} As the injury continues to heal, growth factors such as osteoprotegerin recruit osteoblasts and osteoclasts to form mechanically weak bone.³³ This weak bone will

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continue to be remodeled and will eventually be fully replaced by mechanically robust bone.²⁵

Due to the complexity of native bone healing, a wide range of efforts have explored the use of these and other factors to accelerate cell recruitment and regenerative activity. For example, stromal derived factor 1 and platelet derived growth factor have been delivered from bone-mimetic scaffolds to increase cell migration *in vitro*³⁴ and improve bone healing *in vivo*.^{35,36} Incorporating BMP2 into scaffolds for bone repair is very common and has shown to improve osteogenesis in multiple systems.^{36–42} However, the need for doses of soluble BMP2 larger than what's normally found in the body has led to complications such as abnormal bone formation.^{43,44} Prolonged release of VEGF from bone-mimetic scaffolds in small concentrations has been shown to promote vascularization of the injury site and improve bone healing *in vivo*.⁴⁵ However, delivery strategies that result in quick release of high concentrations of VEGF increase vascularization at the expense of bone quantity *in vivo*.⁴⁶ Thus, it is critical to determine the optimal dose for growth factor delivery, understanding that this dose will vary based on the desired results, and develop strategies to control the delivery and release of factors from biomaterial substrates to accelerate healing.

Our lab has developed a mineralized collagen-glycosaminoglycan scaffold capable of inducing osteogenesis without the addition of exogenous factors,⁴⁷ and these materials have been used to heal sub-critical injuries *in vivo*.^{48,49} However, as we move into larger injury models, we may require biomolecular supplements to improve implant-bone integration, cell recruitment, and vascular remodeling. Thus, we are interested in exploring how growth factor supplementation can be used to improve *in vivo* healing and integration with host tissue. Our group has previously used strategies such as photopatterning,^{50,51} covalent immobilization,^{52–55} and sequestration modulated by glycosaminoglycan content⁵⁶ and small molecules⁵⁷ to add growth factors to non-mineralized and mineralized collagen scaffolds. Recently, the Murphy lab has described the use of modified simulated body fluid (mSBF) to create mineral coatings on material surfaces to deliver growth factors,^{58–62} plasmid DNA lipoplexes,⁶³ and condensed mRNA.⁶⁴ We were particularly interested in the work by Clements *et al.* in which they performed sequential sequestrations to layer interleukin-1 receptor antagonist on nanoparticles to prolong *in vivo* activity.⁶²

In this manuscript, we describe the use of mSBF and sequential sequestration to incorporate and retain BMP2 and VEGF in three-dimensional mineralized collagen scaffolds. First, we hypothesized that soaking mineralized collagen scaffolds in mSBF prior to sequestration would increase incorporation and retention of growth factors within our scaffolds. Next, we hypothesized that sequential sequestrations would increase incorporation and retention of growth factors within our scaffolds compared to a single sequestration. We examined the capacity for mineralized collagen scaffolds to sequester growth factors from solution without additional crosslinking treatments and the capability to increase incorporation and

extend retention of single growth factors or multiple growth factors *via* sequential sequestrations.

2. Materials and methods

2.1. Mineralized collagen scaffold fabrication and hydration

Mineralized collagen-glycosaminoglycan scaffolds were fabricated *via* lyophilization from a mineralized collagen precursor suspension as described before.^{47,65,66} The precursor suspension was created by homogenizing type I collagen (1.9 weight per volume, Sigma Aldrich, St. Louis, Missouri USA), chondroitin-6-sulfate (0.84 weight per volume, Sigma Aldrich), and calcium salts (calcium hydroxide and calcium nitrate, Sigma Aldrich) in a mineral buffer solution (0.1456 M phosphoric acid/0.037 M calcium hydroxide). The precursor suspension was stored at 4 °C and degassed prior to lyophilization.

Mineralized collagen scaffolds were fabricated *via* lyophilization using a Genesis freeze-dryer (VirTis, Gardener, New York USA) as described before.⁶⁶ Briefly, 100 µL of precursor suspension was pipetted into a custom 144-well polysulfone mold (6 mm diameter, 7 mm tall wells). The precursor solution was frozen by cooling from 20 °C to –10 °C at a constant rate of 1 °C per minute followed by a temperature hold at –10 °C for 2 hours. The frozen suspension was then sublimated at 0 °C and 0.2 Torr, resulting in a porous scaffold network.

All scaffolds were hydrated for 2 hours in ethanol, cross-linked for 2 hours in EDC-NHS, and washed in phosphate buffered saline (PBS) for 48 hours prior to use in experiments.

2.2. Modified simulated body fluid preparation

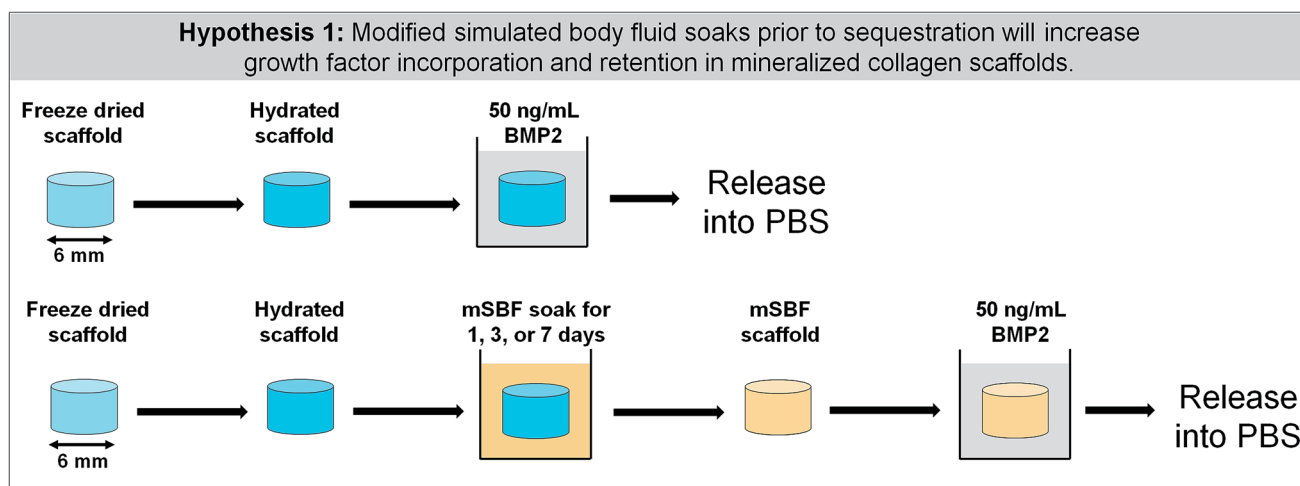
Modified simulated body fluid (mSBF) was made according to previous recipes.^{58–64} Briefly, 1.41 mM sodium chloride (NaCl, Sigma Aldrich), 4.0 mM potassium chloride (KCl, Sigma Aldrich), 0.5 mM magnesium sulfate (MgSO₄, Sigma Aldrich), 1.0 mM magnesium chloride (MgCl₂, Sigma Aldrich), 5.0 mM calcium chloride (CaCl₂, Sigma Aldrich), 1.0 mM potassium phosphate (KH₂PO₄, Sigma Aldrich), and 4.2 mM sodium carbonate (NaHCO₃, Sigma Aldrich) were dissolved in deionized water and sterile filtered through a MilliporeSigma Stericup with a 0.22 µm filter (Fisher Scientific, Hampton, New Hampshire USA). mSBF was stored at 4 °C until use.

2.3. Sequestration of bone morphogenic protein 2 following modified simulated body fluid treatments

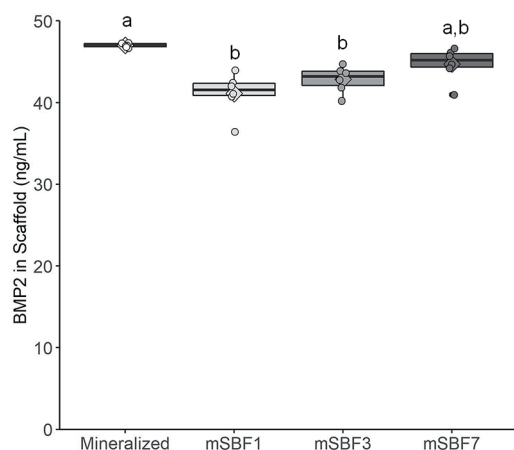
Following hydration, scaffolds were soaked in 50 ng mL^{–1} bone morphogenic protein 2 (BMP2) or treated in modified simulated body fluid (mSBF) for 1, 3, or 7 days and then soaked in 50 ng mL^{–1} BMP2 (Fig. 1A). mSBF soaks were done at room temperature under mild shaking and mSBF solution was changed each day. BMP2 was diluted in 1% bovine serum albumin in phosphate buffered saline (1% BSA in PBS), and sequestration was done for 1 hour at room temperature under mild shaking. After sequestration, the BMP2 solution was saved and stored at –20 °C and scaffolds were placed in PBS for 7 days at 37 °C. PBS was replaced each day and stored at –20 °C. An enzyme-linked immunosorbent assay (ELISA) (R&D Systems, Minneapolis,



A



B



C

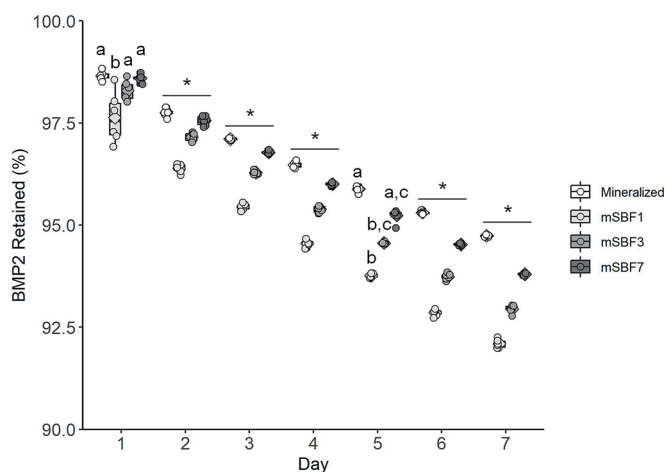


Fig. 1 Growth factor sequestration with varying modified simulated body fluid (mSBF) treatments. (A) Schematic of experiment. Hydrated scaffolds and scaffolds soaked in mSBF for 1, 3, or 7 days were used to sequester BMP2. (B) Sequestration of BMP2 after varying treatment times in mSBF. Mineralized collagen scaffolds are capable of sequestering BMP2 without mSBF treatments. Groups that share a letter are not significantly different ($p < 0.05$). (C) Retention of BMP2 within scaffolds for 7 days. Mineralized collagen scaffolds had the highest retention compared to all mSBF treated groups by day 7. Groups that share a letter within a day are not significantly different ($p < 0.05$). * indicates significant differences between all groups within a day. Note: the y-axis starts at 90% to better show differences between groups. To see plots with the y-axis 0–100% please see ESI Fig. 1.† DATA: boxplots overlaid with individual data points are used to represent data.

Minnesota, USA) was used to quantify the amount of BMP2 sequestered and retained within the scaffolds. Retention is reported as the percent of BMP2 remaining in the scaffold to BMP2 initially sequestered into the scaffold.

2.4. Compression testing

Stress–strain curves of hydrated scaffolds, scaffolds soaked in phosphate-buffered saline (PBS) for 7 days, and scaffolds soaked in modified simulated body fluid (mSBF) for 7 days were generated with the Instron 5943 mechanical tester (Instron, Norwood, Massachusetts USA) using a 5 N load cell; hydrated scaffolds were not submerged in liquid during testing. Samples were compressed at a rate of 1 mm min^{-1} with the Young's modulus determined from the stress–strain curves using

conventional analysis methods for low-density open-cell foam structures such as the mineralized collagen scaffolds.^{67,68}

2.5. Scanning electron microscopy

Critical point drying was done using a Samdri-PVT-3D (Tousimis, Rockville, Maryland USA) to prepare hydrated scaffolds (Mineralized), and scaffolds soaked in modified simulated body fluid (mSBF) for 1, 3, and 7 days (mSBF1, mSBF3, mSBF7) for scanning electron microscopy (SEM) analysis of scaffold morphology using previously described methods.^{57,69} Briefly, scaffolds were placed in formalin for 24 hours and then the aqueous solution in the scaffolds was replaced with ethanol and then liquid carbon dioxide. The specimens were then held above 6.895 kPa and 31°C to remove the carbon dioxide as a gas



with minimal structural deformation. Dried scaffolds and scaffolds cut into semi-cylinders were placed on carbon tape, sputter coated with gold/palladium, and imaged using a Philips XL30 ESEM-FEG (FEI Company) at 5 kV with a secondary electron detector.

2.6. Sequential sequestration of single growth factors

2.6.1. One treatment. Following hydration, scaffolds were soaked in 80 ng mL^{-1} bone morphogenic protein 2 (BMP2) or vascular endothelial growth factor (VEGF) (Fig. 3A). Growth factors were diluted in 1% bovine serum albumin in phosphate buffered saline (1% BSA in PBS), and sequestration was done for 1 hour at room temperature under mild shaking. After sequestration, the growth factor solution was saved and stored at -20°C and scaffolds were placed in PBS for 7 days at 37°C .

2.6.2. Sequential treatments. Following hydration, scaffolds were soaked in 10 ng mL^{-1} BMP2 or VEGF and put into modified simulated body fluid (mSBF) overnight at room temperature under mild shaking. This was repeated for 7 consecutive days (8 total sequestrations) (Fig. 3A). Growth factors were diluted in 1% BSA in PBS, and sequestration was

done for 1 hour at room temperature under mild shaking. After each sequestration, the growth factor solution was saved and stored at -20°C . The solution from mSBF soaks were saved and stored at -20°C . After the final sequestration, scaffolds were placed in PBS for 7 days at 37°C .

2.7. Sequential sequestration of multiple growth factors

2.7.1. One treatment. Following hydration, scaffolds were soaked in 80 ng mL^{-1} bone morphogenic protein 2 (BMP2) and 10 ng mL^{-1} vascular endothelial growth factor (VEGF) (Fig. 4A). Growth factors were diluted in 1% bovine serum albumin in phosphate buffered saline (1% BSA in PBS), and sequestration was done for 1 hour at room temperature under mild shaking. After sequestration, the growth factor solution was saved and stored at -20°C and scaffolds were placed in PBS for 7 days at 37°C .

2.7.2. Sequential treatments. Following hydration, scaffolds were soaked twice in 40 ng mL^{-1} BMP2 and once in 10 ng mL^{-1} VEGF with overnight modified simulated body fluid (mSBF) soaks following each sequestration step (Fig. 4A). mSBF soaks were done at room temperature under mild shaking.

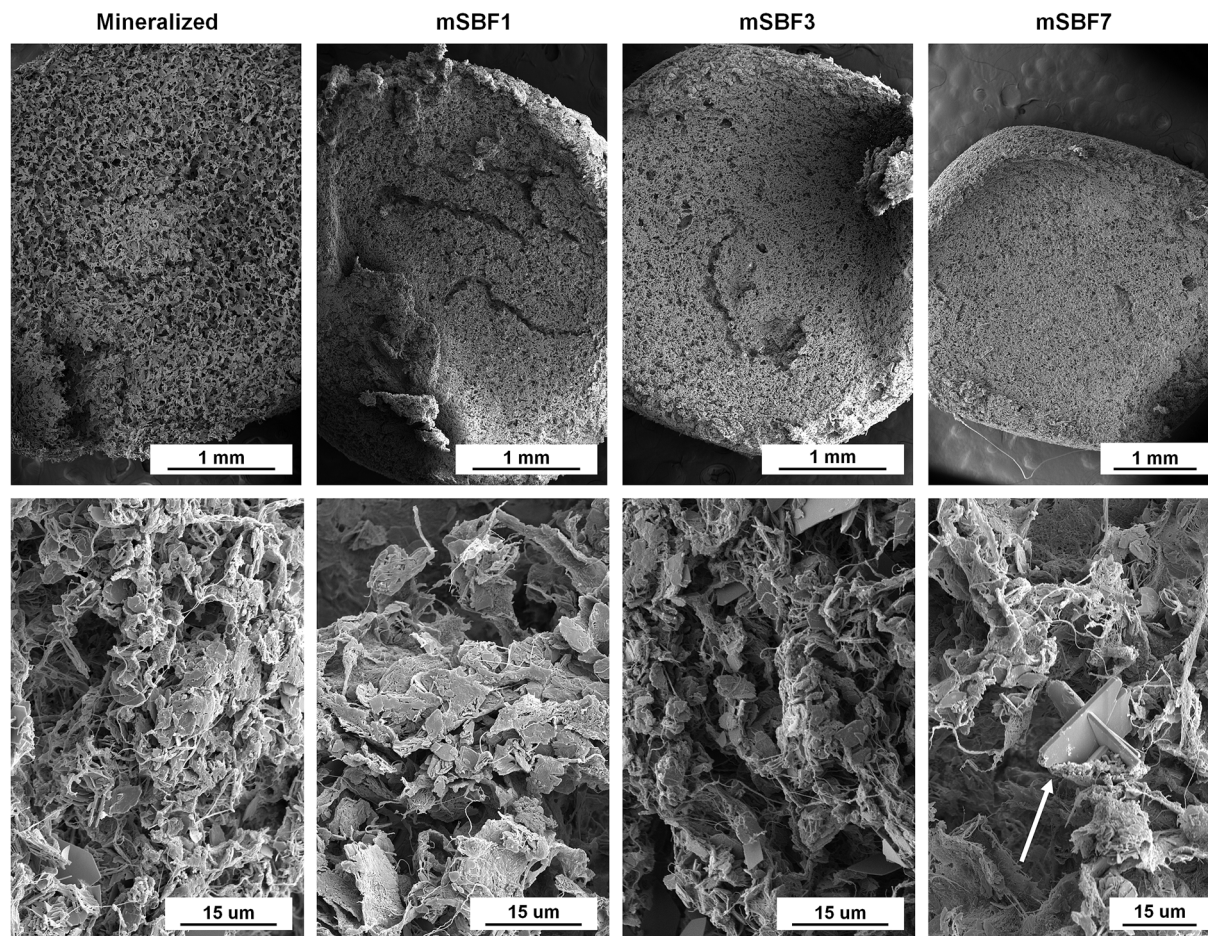


Fig. 2 Scanning electron micrographs of unseeded mineralized collagen scaffolds after exposure to modified simulated body fluid treatments. White arrow indicates angular structures observed during imaging. Magnifications top: $50\times$, $50\times$, $50\times$, $40\times$. Magnifications bottom: $3000\times$, $3000\times$, $3000\times$, $2500\times$.

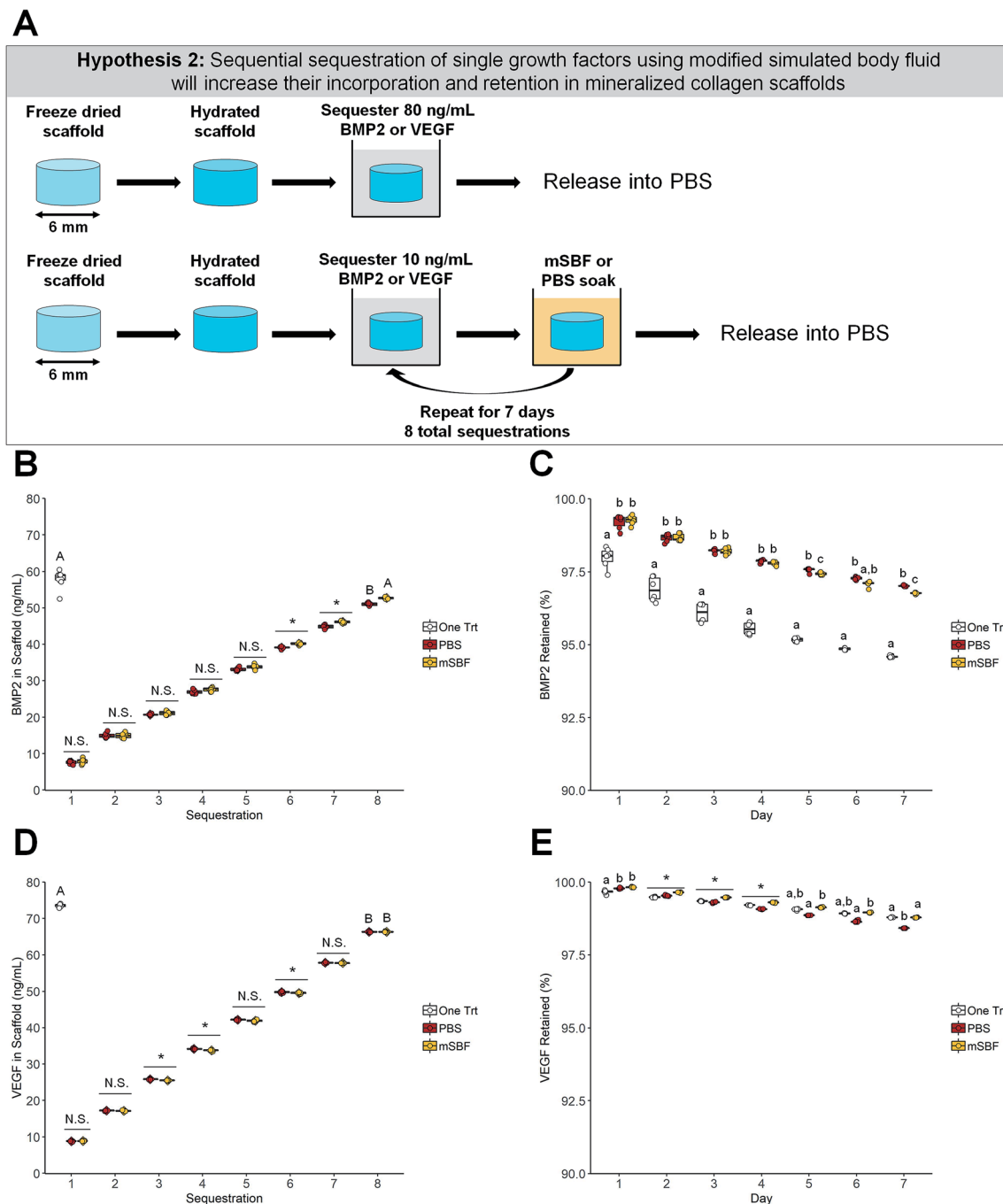


Fig. 3 Sequential sequestration of individual growth factors. (A) Schematic of experiment. Scaffolds soaked once in BMP2 or VEGF were compared to sequentially sequestered BMP2 or VEGF groups. (B) BMP2 sequestration. The one-time BMP2 treatment and the modified simulated body fluid (mSBF) sequentially sequestered groups had the same amount of BMP2 sequestered ($p < 0.05$). * indicates significance between all groups within a day ($p < 0.05$). N.S. indicates no significance between groups within a day ($p < 0.05$). Groups that share a letter are not significantly different ($p < 0.05$). (C) BMP2 retention. Scaffolds treated once with BMP2 had lower retention of the growth factor in the scaffold compared to both sequentially sequestered groups. Groups that share a letter within a day are not significantly different ($p < 0.05$). Note: the y-axis starts at 90% to better show differences between groups. To see plots with the y-axis 0–100% please see ESI Fig. 3.† (D) VEGF sequestration. The one-time VEGF treatment had more VEGF sequestered than both sequentially sequestered groups ($p < 0.05$). * indicates significance between groups within a day ($p < 0.05$). N.S. indicates no significance between groups within a day ($p < 0.05$). Groups that share a letter are not significantly different ($p < 0.05$). (E) VEGF retention. Scaffolds soaked once in VEGF and scaffolds sequentially sequestered with mSBF treatments have the highest retention by day 7. * indicates significance between all groups within a day ($p < 0.05$). Groups that share a letter within a day are not significantly different ($p < 0.05$). Note: the y-axis starts at 90% to better show differences between groups. To see plots with the y-axis 0–100% please see ESI Fig. 3.† DATA: boxplots overlaid with individual data points are used to represent data.



Growth factors were diluted in 1% BSA in PBS, and sequestration was done for 1 hour at room temperature under mild shaking. After each sequestration, the growth factor solution was saved and stored at -20°C . The solution from mSBF soaks were saved and stored at -20°C . After the final sequestration, scaffolds were placed in PBS for 7 days at 37°C .

2.8. Enzyme-linked immunosorbent assays

2.8.1. One-time treatments. PBS was replaced each day and stored at -20°C during growth factor release. An enzyme-linked immunosorbent assay (ELISA) (R&D Systems) was used to quantify the amount of BMP2 and VEGF sequestered and retained within the scaffolds. Retention is reported as the percent of BMP2 remaining in the scaffold to BMP2 initially sequestered into the scaffold.

2.8.2. One-time treatments. PBS was replaced each day and stored at -20°C during growth factor release. An enzyme-linked immunosorbent assay (ELISA) (R&D Systems) was used to quantify the amount of BMP2 and VEGF sequestered and retained within the scaffolds. Growth factor released during mSBF soaks was factored into the final concentration of BMP2 or VEGF sequestered into mineralized collagen scaffolds. Retention is reported as the percent of BMP2/VEGF remaining in the scaffold to BMP2/VEGF initially sequestered into the scaffold.

2.9. Human umbilical vein endothelial cell culture and preliminary transwell experiment

Human umbilical vein endothelial cells (HUVECs) (Lonza, Basel, Switzerland) were expanded in T75 flasks (Fisher Scientific) and cultured in endothelial cell growth media (Lonza) at 37°C and 5% carbon dioxide until confluent. Once confluent, passage 4 HUVECs were seeded (62 000 cells per well) into the lower chamber of the transwell plate and scaffolds were loaded into the transwell insert. 300 μL of endothelial cell growth media was used in the lower chamber and 700 μL of endothelial cell growth media was used in the transwell insert. Cells were cultured in endothelial cell growth media at 37°C and 5% carbon dioxide for 7 days.

Four groups were used for this preliminary transwell experiment: (1) scaffolds soaked in PBS with no VEGF added into the transwell insert (blank); (2) scaffolds soaked in PBS with 10 ng mL^{-1} VEGF added into the transwell insert (soluble); (3) scaffolds that had been soaked in 5 ng mL^{-1} of VEGF, soaked in mSBF overnight, and soaked in another 5 ng mL^{-1} of VEGF (mSBF); (4) scaffolds soaked once in 10 ng mL^{-1} VEGF (one Trt). The soluble groups had no additional VEGF added after the first media change at day 3.

2.10. Human umbilical vein endothelial cell metabolic activity

The metabolic activity of human umbilical vein endothelial cells (HUVECs) seeded in transwells was measured using alamarBlue *via* fluorescent spectrophotometer (Tecan Infinite F200 Pro, Männedorf, Switzerland). HUVECs seeded in the bottom chamber of a transwell were incubated in a 10% alamarBlue

solution (Invitrogen, Carlsbad, California USA) for 90 minutes at 37°C under moderate shaking. The relative cell metabolic activity was determined from a standard curve generated with known cell concentrations. An experimental value of 1 indicates the metabolic activity of the number of cells originally seeded into the transwell.

2.11. Human umbilical vein endothelial cell gene expression

RNA was isolated from cell seeded scaffolds using TRIzol Reagent (Invitrogen) following the provided protocol. Isolated RNA was quantified using the NanoDrop Lite Spectrophotometer (ThermoFisher, Waltham, Massachusetts USA) and reverse transcribed using a QuantiTect Reverse Transcription Kit (Qiagen, Hilden, Germany) and a BioRad S1000 thermal cycler (BioRad, Hercules, California USA).

Each real-time polymerase chain reaction (PCR) was carried out in duplicate, using 10 ng cDNA and Taqman primers (ThermoFisher). A Taqman PCR kit (ThermoFisher) along with an Applied Biosystems 7900HT Fast Real-Time PCR System was used to perform the real-time PCR. 18S ribosomal RNA (18S) and peptidylprolyl isomerase A (PPIA) were used as house-keeping genes. Gene expression profiles were obtained for platelet derived growth factor (PDGF), hypoxia inducible factor 1 α (Hif1a), angiopoietin 1 (Ang1), and angiopoietin 2 (Ang2) (details: ESI Table 1†).

2.12. Statistics

RStudio was used for all plotting and statistical analyses. Sample size was six ($n = 6$) for all experiments except compression testing ($n = 12$) and polymerase chain reaction (PCR, $n = 8$). PCR data have some groups with sample size less than 8 due to undetermined threshold cycle values, but 90% of groups have $n \geq 6$. See ESI Tables 2 and 3† for details on sample size for PCR data. Data in tables are presented as average $\pm$ standard deviation.

2.12.1. Plotting. The Rstudio data visualization package “ggplot2” was used to plot all data. The graphs in this manuscript display the data using boxplots overlaid with individual observations.

2.12.2. Statistics on data with two experimental groups. A t -test was used for data that were normally distributed and had equal variance. A Welch's t -test was used for data that were normally distributed and did not have equal variance. The Mann-Whitney U test was used for data that were not normally distributed and had equal variance. Experimental groups were tested for normality using the Shapiro-Wilk test. Homogeneity of variance was tested using the Levene test.

2.12.3. Statistics on data with three or more experimental groups. The Kruskal-Wallis test (one experimental factor) was used for data that were not normally distributed and had equal variance, and significance was determined using Dunn's post-hoc test. One-way ANOVA (one experimental factor) or two-way ANOVA (two experimental factors) was run for data that were normally distributed and had equal variance, and significance was determined using Tukey's post-hoc test. Welch's one-way ANOVA was used for data that were normally distributed and



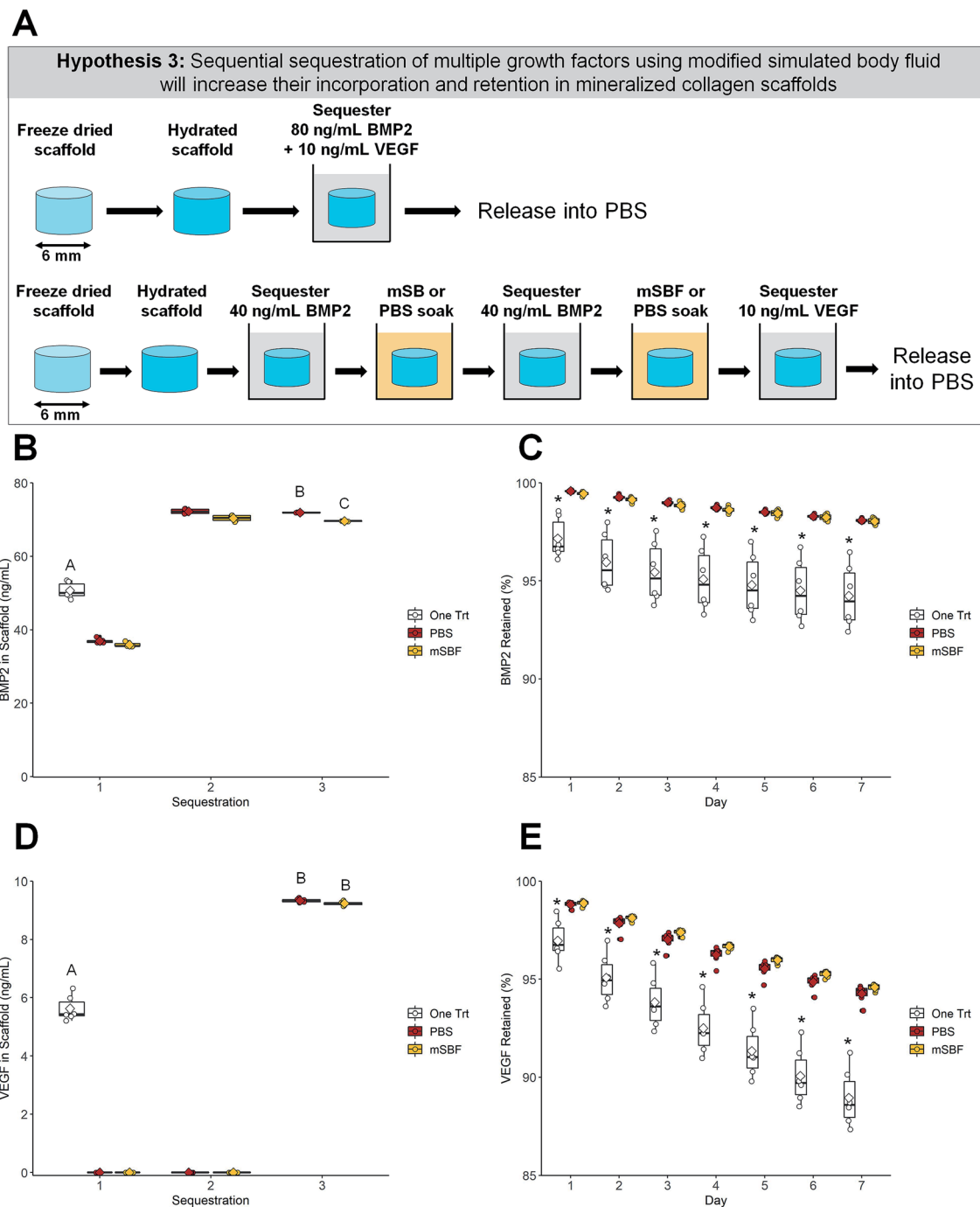
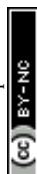


Fig. 4 Sequential sequestration of multiple growth factors. (A) Schematic of experiment. Scaffolds soaked once in a solution of BMP2 and VEGF were compared to sequentially sequestered BMP2 then VEGF groups. (B) BMP2 sequestration. Both sequentially sequestered scaffolds had higher concentrations of BMP2 compared to the one-treatment scaffolds. Groups that share a letter are not significantly different ($p < 0.05$). (C) BMP2 retention. Scaffolds treated once in BMP2 have the lowest retention compared to both sequentially sequestered groups ($p < 0.05$). * indicates significant differences compared to the other two groups within a day ($p < 0.05$). Note: the y-axis starts at 85% to better show differences between groups. To see plots with the y-axis 0–100% please see ESI Fig. 4.† (D) VEGF sequestration. Both sequentially sequestered scaffolds had higher concentrations of VEGF compared to the one-treatment scaffolds. Groups that share a letter are not significantly different ($p < 0.05$). (E) VEGF retention. Scaffolds treated once in VEGF have the lowest retention compared to both sequentially sequestered scaffolds. * indicates significant differences compared to the other two groups within a day ($p < 0.05$). Note: the y-axis starts at 85% to better show differences between groups. To see plots with the y-axis 0–100% please see ESI Fig. 4.† DATA: boxplots overlaid with individual data points are used to represent data.

did not have equal variance, and significance was determined using Tukey's post-hoc test. Mood's median test (*via* a Monte Carlo simulation, one experimental factor) was used for data

that were not normally distributed and did not have equal variance, and significance was determined using a pairwise median test. Residuals were tested for normality using the



Shapiro–Wilk test. Homogeneity of variance was tested using the Levene test.

3. Results

3.1. Mineralized collagen scaffolds can sequester growth factors without additional treatments

We first determined whether extended soaking of mineralized collagen scaffolds in modified simulated fluid (mSBF) prior to growth factor sequestration would increase incorporation and retention of growth factors within our scaffolds. Our mineralized collagen scaffold natively sequesters bone morphogenic protein 2 (BMP2) better than mineralized collagen scaffolds soaked in mSBF for 1 or 3 days (Fig. 1B). However, extended exposure to mSBF (7 days) recovers this sequestration ability, with no significant difference in BMP2 sequestration compared to the native mineralized scaffolds (Fig. 1B). All groups sequestered greater than 80% of BMP2 out of solution (Fig. 1B). The mineralized collagen scaffolds also retained the highest percentage of sequestered BMP2 over 7 days compared to all mSBF soaked scaffold groups, with the difference becoming significant by day 2 (Fig. 1C). Overall, mineralized collagen scaffolds had the highest final concentration of BMP2 at day 7 than scaffolds with extended mSBF treatments prior to sequestration (Table 1). Scaffolds that were soaked in mSBF for 7 days both sequestered and retained BMP2 better than shorter mSBF exposures (1 and 3 days); however, subsequent experiments did not include mSBF pre-soaks because there was no improvement in BMP2 sequestration or retention compared to native mineralized collagen scaffolds. Summary statistics (average $\pm$ standard deviation) and full-scale retention data for this experiment: Table 1, ESI Fig. 1.†

The scaffold microstructure was not noticeably affected by mSBF exposure (Fig. 2). The porous network and collagen fibers are unchanged when viewed using scanning electron microscopy (Fig. 2). While we normally observe plate-like brushite crystals in our mineralized scaffolds,⁶⁹ we observed additional deposits in scaffolds soaked in mSBF for 7 days (Fig. 2, white arrow). We believe this may have been a result of the high salt concentrations in the mSBF soaks, and tested the mechanics of these scaffolds to define mechanical consequences of mSBF exposure. Mechanical analysis of the different scaffolds showed the elastic modulus of scaffolds soaked in mSBF or PBS for 7 days is significantly lower than immediately after hydration (ESI Fig. 2†). However, we observed no significant differences in elastic modulus between scaffolds soaked in mSBF or PBS for 7

days (ESI Fig. 2†), suggesting the loss of mechanical performance is not a specific effect of mSBF exposure.

3.2. Sequential sequestration of individual growth factors increases retention in mineralized collagen scaffolds

We subsequently examined the use of mSBF treatments and sequential sequestrations to increase incorporation and retention of growth factors within our scaffolds compared to a single sequestration (one-time treatment). BMP2 and vascular endothelial growth factor (VEGF) can be sequentially sequestered into mineralized collagen scaffolds, and more VEGF was sequestered into mineralized collagen scaffolds than BMP2 for all treatment groups (Fig. 3B and D). Notably, sequential sequestrations with repeated mSBF soaks resulted in significantly higher final concentrations of incorporated BMP2 than sequential sequestrations with repeated PBS soaks (Fig. 3B). One-time treatments and the sequentially sequestered mSBF groups sequestered $\sim 72\%$ and 66% of BMP2 out of solution, respectively, while sequentially sequestered PBS groups had significantly lower sequestration ($\sim 63\%$ BMP2 sequestered; Fig. 3B). One-time treatments sequestered significantly more VEGF out of solution than both sequentially sequestered groups (mSBF and PBS; Fig. 3D). One-time treatments sequestered $\sim 91\%$ VEGF out of solution while both sequentially sequestered groups (mSBF and PBS) sequestered $\sim 82\%$ VEGF out of solution (Fig. 3D).

Next, we examined whether the mode of growth factor sequestration into the scaffold altered factor release. Notably, mineralized collagen scaffolds have higher retention of sequestered BMP2 when sequentially sequestered compared to a one-time treatment for the entire release period (Fig. 3C). Sequentially sequestered PBS groups have higher BMP2 retention than sequentially sequestered mSBF groups by day 7 (97.02% and 96.77%, respectively; Fig. 3C). Although the sequentially sequestered groups have higher retention of sequestered BMP2, the final concentration of BMP2 in the scaffolds is highest in the one-time treatment group (54.77 ng mL^{-1} ; Table 2). Mineralized collagen scaffolds with a one-time treatment and sequentially sequestered scaffolds with mSBF soaks have the highest retention of sequestered VEGF by day 7 (Fig. 3E), though all groups have greater than 98% of sequestered VEGF retained after 7 days (Fig. 3E). Here, one-time treatments have the highest final concentration of VEGF ($\sim 72 \text{ ng mL}^{-1}$) compared to the sequentially sequestered groups ($\sim 65 \text{ ng mL}^{-1}$ VEGF for mSBF and PBS treatments; Table 2). Summary

Table 1 Sequestration and retention values for Fig. 1 (mSBF treatments)^a

Group	BMP2 sequestered (ng mL^{-1})	BMP2 retained (% , day 7)	BMP2 retained (ng mL^{-1} , day 7)
Mineralized	46.99 ± 0.25	94.74 ± 0.03	45.72 ± 0.01
mSBF1	41.12 ± 2.55	92.09 ± 0.10	39.48 ± 0.02
mSBF3	42.84 ± 1.61	92.94 ± 0.09	41.30 ± 0.02
mSBF7	44.70 ± 2.06	93.80 ± 0.25	43.29 ± 0.01

^a Data is presented as average $\pm$ standard deviation. The sequestration was done out of 50 ng mL^{-1} BMP2.



Table 2 Sequestration and retention values for Fig. 3 (individual growth factors)^a

Group	BMP2			VEGF		
	Sequestered (ng mL ⁻¹)	Retained (%; day 7)	Retained (ng mL ⁻¹ , day 7)	Sequestered (ng mL ⁻¹)	Retained (%; day 7)	Retained (ng mL ⁻¹ , day 7)
One Trt	57.90 ± 2.85	94.59 ± 0.03	54.77 ± 0.02	73.50 ± 0.44	98.79 ± 0.01	72.61 ± 0.01
PBS	51.06 ± 0.46	97.02 ± 0.02	49.54 ± 0.01	66.33 ± 0.16	98.42 ± 0.01	65.28 ± 0.01
mSBF	52.73 ± 0.26	96.77 ± 0.03	51.02 ± 0.01	66.34 ± 0.12	98.79 ± 0.1	65.53 ± 0.01

^a Data is presented as average ± standard deviation. The sequestration was done out of 80 ng mL⁻¹ BMP2 or 80 ng mL⁻¹ VEGF.

statistics (average ± standard deviation) and full-scale retention data for this experiment: Table 2, ESI Fig. 3.†

3.3. Sequential sequestration of multiple growth factors increases sequestration and retention in mineralized collagen scaffolds

We then explored whether the sequential sequestration approach provides an advantage for selectively incorporating multiple growth factors within the scaffold. Scaffolds could be exposed to a large dose of mixed factors (one-time treatments), or scaffolds could be sequentially exposed to one factor, followed by a PBS or mSBF treatment, then exposed to a second factor. Scaffolds that sequentially sequestered BMP2 then VEGF had higher concentrations of both growth factors compared to scaffolds that were soaked once in a solution containing BMP2 and VEGF (Fig. 4B and D). One-time treatments sequestered ~60% of both BMP2 and VEGF while sequentially sequestered groups sequestered ~87% of BMP2 and greater than 90% of VEGF from solution. Data show it's possible to quantify growth factor sequestration following each stage of exposure: (1) exposure to 40 ng mL⁻¹ BMP2; (2) exposure to a second dose of 40 ng mL⁻¹ BMP2; (3) exposure to 10 ng mL⁻¹ VEGF (Fig. 4B and D).

Sequentially sequestered groups (mSBF and PBS) also showed higher retention of both growth factors over 7 days when compared to one-time treatment scaffolds (Fig. 4C and E). Sequentially sequestered scaffolds retained ~98% and ~95% of sequestered BMP2 and VEGF by day 7, respectively. One-time treatments retained ~94% and ~88% BMP2 and VEGF by day 7, respectively. Scaffolds that sequentially sequestered growth factors from solution had the highest final concentration of BMP2 and VEGF by day 7 (~68 ng mL⁻¹ and ~9 ng mL⁻¹, respectively; Table 3). Additionally, scaffolds that sequentially sequestered growth factors were less variable in their retention profiles (indicated by a narrower boxplot). Summary statistics (average ± standard deviation) and full-scale retention data for this experiment: Table 3, ESI Fig. 4.†

3.4. *In vitro* cell activity data suggest large growth factor doses are required to illicit cell response

We subsequently examined the ability for 10 ng mL⁻¹ sequestered VEGF to induce shifts in human umbilical vein endothelial cell (HUVEC) activity using a transwell assay (*i.e.* examining the effect of released VEGF). We compared results for HUVECS in the scaffold with no exposure to VEGF (blank), exposure to 10

ng mL⁻¹ VEGF supplemented in the media (soluble), a single VEGF sequestration (one Trt), or sequential VEGF sequestration with mSBF treatments (mSBF). We did not observe significant differences in HUVEC metabolic activity between treatment groups in response to 10 ng mL⁻¹ VEGF by day 7 (Fig. 5). While we observed some short-term differences (days 1 and 4) in gene expression for platelet derived factor (PDGF), hypoxia induced factor 1 alpha (HIF1A), angiopoietin-1 (ANG1), and angiopoietin-2 (ANG2), there was no significant effect of VEGF delivery method on long-term gene expression (day 7) (Fig. 6).

4. Discussion

Bone healing is a complex process coordinated through growth factor signaling,^{21–25} and tissue engineers use these factors to improve bone healing in bone-mimetic scaffolds.^{19,20} Bone morphogenic protein 2 (BMP2),^{20,42} vascular endothelial growth factor (VEGF),²⁰ and platelet derived growth factor (PDGF)³⁴ are popular candidates for use in bone biomaterials. In this manuscript, we described the use of modified simulated body fluid (mSBF) and sequential sequestrations to incorporate BMP2 and VEGF into mineralized collagen scaffolds. This work seeks to adapt promising results using mSBF to deposit a mineral layer that can sequester growth factors onto surfaces^{58–64} to selectively incorporate growth factors used for bone repair applications into three-dimensional, porous biomaterials. Our primary goal is to improve growth factor incorporation and retention into mineralized collagen scaffolds *via* sequential sequestration strategies to enhance *in vivo* bone healing.

First, we hypothesized that extended exposure to mSBF prior to sequestration would increase growth factor incorporation and retention in mineralized collagen scaffolds. We found that 1 and 3 day mSBF treatments prior to sequestration reduced the amount of bone morphogenic protein 2 (BMP2) incorporated and retained in mineralized collagen scaffolds (Fig. 1B, C and Table 1). This suggests the native mineral content of the mineralized collagen scaffolds is enough to sequester growth factors without additional treatments and that mSBF treatments prior to sequestration do not improve retention. Our mineralized collagen scaffolds are high in mineral content (40 weight percent⁶⁵) and highly porous (>85%⁶⁶), so it is likely that these two properties allowed for the high sequestration efficiency in our mineralized collagen scaffolds. We are able to



Table 3 Sequestration and retention values for Fig. 4 (multiple growth factors)^a

Group	BMP2			VEGF		
	Sequestered (ng mL ⁻¹)	Retained (% day 7)	Retained (ng mL ⁻¹ , day 7)	Sequestered (ng mL ⁻¹)	Retained (% day 7)	VEGF retained (ng mL ⁻¹ , day 7)
One Trt	50.64 ± 2.16	94.23 ± 1.62	47.72 ± 0.82	5.62 ± 0.43	88.95 ± 1.48	5.00 ± 0.08
PBS	71.91 ± 0.09	98.09 ± 0.08	70.53 ± 0.06	9.34 ± 0.07	94.25 ± 0.46	8.76 ± 0.04
mSBF	69.58 ± 0.06	98.04 ± 0.16	68.22 ± 0.11	9.24 ± 0.07	94.59 ± 0.18	8.75 ± 0.02

^a Data is presented as average ± standard deviation. The sequestration was done out of 80 ng mL⁻¹ BMP2 and 10 ng mL⁻¹ VEGF.

control the mineral content⁶⁵ and pore size⁶⁶ of our materials, and future efforts will explore the effects of these characteristics on the sequestration and retention capabilities of mineralized collagen scaffolds.

Scaffolds soaked in mSBF for 1, 3 and 7 days exhibited no significant changes in the collagen network (Fig. 2). However, scaffolds soaked in either mSBF or PBS for 7 days were significantly softer than hydrated scaffolds (ESI Fig. 2†), consistent with prior data showing that the osteogenic nature of the mineralized collagen scaffold is enhanced *via* release of mineral ions into the media.⁴⁷ While the porous nature of the scaffolds is essential for biological activity (*i.e.* cell penetration, diffusive biotransport), all scaffold variants show sub-optimal mechanical performance. While beyond the scope of this project, we have described methods to increase the mechanical stability of a cell activity optimized scaffolds *via* the inclusion of 3D printed polymeric meshes.^{49,70} Based on these results, we did not use extended mSBF treatments prior to sequestration in subsequent experiments and instead focused on repeated mSBF treatments during sequestration.

Second, we hypothesized that sequential exposure to mSBF during growth factor sequestration would increase the incorporation and retention of growth factors in mineralized collagen scaffolds. Clements *et al.* has previously identified repeated mSBF treatments as a method to increase growth factor incorporation, maintain protein activity,⁶⁰ and prolong the release of growth factors.⁶² One-time treatments and sequentially sequestered mSBF groups showed equal ability to incorporate BMP2 into the mineralized collagen scaffolds, but sequestered BMP2 had higher retention in the sequentially sequestered scaffolds (mSBF and PBS) (Fig. 3B, C and Table 2). This suggests that the mSBF and PBS soaks between sequestrations aid in the retention of BMP2 within mineralized collagen scaffolds. However, one-time treatments sequestered higher concentrations of VEGF out of solution than both sequentially sequestered groups, and sequestered VEGF had the highest retention in scaffolds with a one-time treatment and scaffolds sequentially sequestered with mSBF soaks (Fig. 3D, E and Table 2). This prolonged VEGF release may be useful for *in vivo* implants, as suggested by previous work.⁴⁵

VEGF (20–22 kDa; R&D systems catalog #293-VE) is approximately 37% larger than BMP2 (15–16 kDa; R&D systems catalog #355-BM), and this size difference might play a role in how these two factors are incorporated and maintained inside our mineralized collagen scaffolds. While we observed largely consistent results between mSBF and PBS soaks, we plan to test protein integrity and stability *via* circular dichroism^{71,72} or differential scanning calorimetry.^{73,74} Future efforts will also be needed to confirm protein bioactivity by evaluating the activation of receptors for the growth factor of interest in cells (*e.g.* vascular endothelial growth factor receptor, VEGFR,^{75,76} and platelet derived growth factor receptor, PDGFR⁷⁷). VEGFR activation *via* phosphorylation has been used to evaluate the bioactivity of VEGF,^{75,76} and PDGFR activation and the phosphorylation of Erk1/2 and Akt have been used to evaluate the bioactivity of PDGF isoforms.⁷⁷ Previous work utilizing mSBF treatments during sequestration suggests that the mineral coatings stabilize the growth factors and prolong their activity.⁶² The experiments described above will refine our understanding of whether mSBF soaks are necessary for extended protein activity in our scaffolds or if PBS soaks would be enough due to the already high mineral content in our materials.^{65,66} Regardless, these results demonstrate that strategies to incorporate growth factors within the scaffold may require different

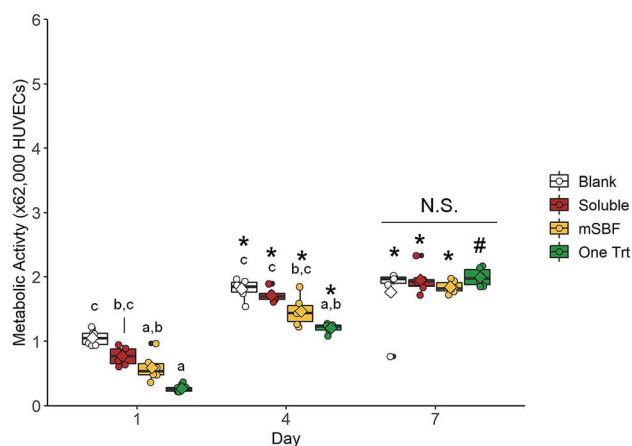


Fig. 5 Cell metabolic activity in response to vascular endothelial growth factor released from scaffolds. Boxplots overlaid with individual data points are used to represent data. There is no significant difference in human umbilical vein endothelial cell metabolic activity between groups by day 7. Groups that share a letter within a day are not significantly different ($p < 0.05$). N.S. indicates no significant difference between indicated groups ($p < 0.05$). * indicates significance compared to the same group at day 1 ($p < 0.05$). # indicates significance compared to the same group at day 1 and day 4 ($p < 0.05$).



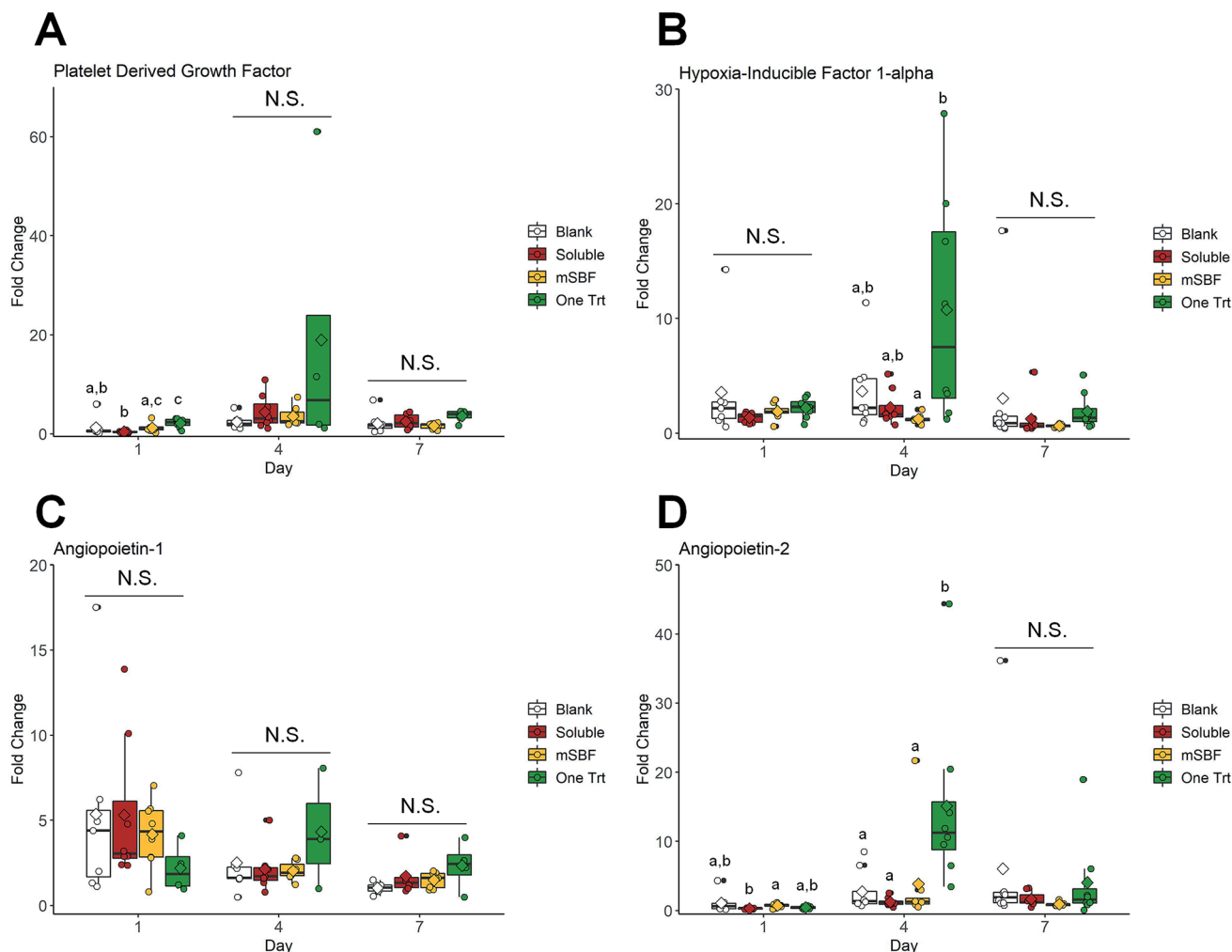


Fig. 6 Gene expression in response to vascular endothelial growth factor released from scaffolds. Boxplots overlaid with individual data points are used to represent data. (A) Platelet derived growth factor fold change. (B) Hypoxia induced factor 1 alpha fold change. (C) Angiopoietin-1 fold change. (D) Angiopoietin-2 fold change. Overall, there are no significant differences in human umbilical vein endothelial cell gene expression between treatments groups by day 7. Groups that share a letter within a day are not significantly different ($p < 0.05$). N.S. indicates no significant difference between indicated groups ($p < 0.05$).

methodologies to maximize incorporation and retention. As a result, it was essential to show the sequestration-based approach reported here could be adapted to sequentially incorporate multiple factors.

Third, we hypothesized that sequestering multiple growth factors *via* sequential sequestration would result in higher incorporation and retention of both factors. These results appear most promising because scaffolds that sequentially sequestered BMP2 then VEGF had significantly higher incorporation of both growth factors than scaffolds treated in a mixed solution of BMP2 and VEGF (Fig. 4B, D and Table 3). Here, the one-time treatment group sequestered ~60% of both BMP2 and VEGF from solution while sequentially sequestered groups sequestered ~90% of BMP2 and VEGF from solution. Retention of both factors and final BMP2 and VEGF concentrations within the scaffolds was higher in the sequentially sequestered groups (Fig. 4C, E and Table 3), meaning our mineralized collagen scaffolds sequester high concentrations of

multiple factors within a single construct and have those factors retained for an extended period. We believe this sustained release would be particularly useful for cell migration and new vessel formation. Prolonged VEGF and platelet derived growth factor (PDGF) presence has been shown to promote migration, proliferation, and angiogenesis.^{45,78}

Lastly, we evaluated cell activity in response to 10 ng mL^{-1} VEGF delivered through a transwell membrane (*i.e.* VEGF released from the scaffolds). These preliminary results suggest larger growth factors doses will be required to illicit extended cell responses (Fig. 5 and 6). There are no significant effects of VEGF (soluble or sequestered) on human umbilical vein endothelial cell metabolic activity (Fig. 5) and gene expression (Fig. 6, ESI Tables 2 and 3†) compared to blank scaffolds (no VEGF) after 7 days. This is not surprising given the wide range of VEGF doses used *in vitro* and *in vivo* for tissue engineering purposes (25 ng mL^{-1} to $250 \text{ } \mu\text{g mL}^{-1}$),^{45,79–82} suggesting future efforts are required to define the correct dose for our mineralized collagen



scaffolds. Immediate next steps are to determine the correct concentration of released growth factor needed to illicit a cellular response. However, we suspect that cells will be most influenced when cultured directly on the scaffolds due to the high concentration of growth factors retained within the material. Therefore, we will also culture cells within the scaffolds and analyze gene and protein expression to see how biologically active retained growth factors are. In both cases, released and retained growth factors, we are interested in dosing our mineralized collagen scaffolds with PDGF to induce cell migration and VEGF to induce vessel formation.

5. Conclusions

We have previously reported the use of covalent immobilization, photopatterning, and supramolecular interactions using cyclodextrins and glycosaminoglycans to incorporate growth factors into non-mineralized and mineralized collagen scaffolds. Here we explored sequential sequestration and the use of modified simulated body fluid to incorporate growth factors into mineralized collagen scaffolds and prolong growth factor release *in vitro*. We report the native mineralized collagen scaffolds can sequester growth factors (>90%) without any additional treatments, and this is a feature of these materials we have not previously explored. We show improved bone morphogenic protein 2 (BMP2) retention in mineralized collagen scaffolds using modified simulated body fluid treatments and sequential sequestrations (97% retention compared to 94% for one-time treatments). Notably, we demonstrate sequential sequestration can significantly increase incorporation and prolong retention for multi-factor cocktails (BMP2 and vascular endothelial growth factor, VEGF). Sequentially sequestering multiple factors led to ~90% of BMP2 and VEGF being incorporated compared to only 60% when they were sequestered in a one-time treatment. Sequentially sequestered groups retained 98% of BMP2 and 94% of VEGF after 7 days while one-time treatments retained 94% of BMP2 and 89% of VEGF after 7 days. These methods provide an additional means to add growth factors into our materials and add complexity to aid in the healing of large bone injuries *in vivo*. Future work will investigate growth factor activity after sequestration and its influence on cellular activities critical to bone healing such as cell migration and vessel formation.

Conflicts of interest

There are no conflicts to declare.

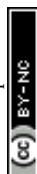
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