An Autologous Muscle Tissue Expansion Approach for the Treatment of Volumetric Muscle Loss

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Abstract

Volumetric muscle loss (VML) is a hallmark of orthopedic trauma with no current standard of care. As a potential therapy for some VML indications, autologous minced muscle grafts (1 mm³ pieces of muscle) are effective in promoting remarkable de novo fiber regeneration. But they require ample donor muscle tissue and therefore may be limited in their application for large clinical VML. Here, we tested the hypothesis that autologous minced grafts may be volume expanded in a collagen hydrogel, allowing for the use of lesser autologous muscle while maintaining regenerative and functional efficacy. The results of the study indicate that 50% (but not 75%) less minced graft tissue suspended in a collagen hydrogel promoted a functional improvement similar to that of a 100% minced graft repair. However, approximately half of the number of fibers regenerated de novo with 50% graft repair. Moreover, the fibers that regenerated had a smaller cross-sectional area. These findings support the concept of using autologous minced grafts for the regeneration of muscle tissue after VML, but indicate the need to identify optimal carrier materials for expansion.

Key words: physiology; tissue engineering; skeletal muscle; injury

Introduction

Muscle weakness and sub-optimal limb function are commonly a part of the sequelae of orthopedic trauma, even after successful limb salvage.1,2 The volumetric loss of skeletal muscle (volumetric muscle loss [VML]) often underlies persistent functional deficits after trauma, and there is no standard of care for the regeneration of the lost muscle tissue. Autologous free or rotational muscle grafts are commonly used for soft tissue replacement after severe open fracture to support bone regeneration, but not for the purpose of restoring muscle function.3 In some instances, functional whole skeletal muscle grafts have been successfully transplanted.4 And a breadth of animal studies have indicated that whole skeletal muscle grafts with surgically restored tendon attachment and neurovascular supply may be sufficiently large for clinical relevance.5 However, the availability of an adequate size donor muscle and potential donor site morbidity may limit the treatment of large VML injuries using this approach.

As a guide, the gold standard treatment for segmental bone defects involves transplantation of autologous bone grafts.6 However, in recognition of the limited availability of autologous bone for grafting, biomaterials are effective as conductive agents. For example, co-delivery of demineralized bone matrices (e.g., bone allograft) with autologous bone grafts is an effective means to volumetrically expand bone graft-mediated regeneration.7,8 Although differences exist in the mechanisms underlying bone and muscle regeneration, an analogous approach using autologous muscle grafts is reasonable for the repair of VML. As an alternative to replacing muscle tissue with whole muscle grafts, muscle regeneration may be achieved using autologous minced muscle grafts. Whole muscles minced into small pieces (1 mm³) maintain the capability to regenerate a functional muscle, although the regenerated...
fibers may be misaligned.9–12 More recently, minced muscle grafts were effective in promoting de novo muscle fiber regeneration and functional recovery in a rodent VML model,13 and minced grafts were successful in improving urinary incontinence in humans.14 An advantage of autologous minced versus whole muscle grafts is the potential for volume expansion in a myoconductive biomaterial, that is, the potential to regenerate a relatively large volume of muscle with a reduced amount of donor tissue.

The purpose of this study was to investigate the expansion capacity of autologous minced muscle grafts for the treatment of VML. This initial investigation involved the transplantation of a titrated amount of autologous minced skeletal muscle grafts delivered in a standard collagen hydrogel to the site of injury, the tibialis anterior (TA) muscle, in a rat VML model. Collagen hydrogel was chosen as an initial expansion biomaterial, because collagen I is the primary constituent of skeletal muscle extracellular matrix (ECM) and collagen-based gels and ECMs are available for clinical use.15,16 100% minced graft and nonrepaired muscles served as positive and negative treatment controls, respectively, to establish upper- and lower-bound limits of regeneration in this VML model.

Materials and Methods

Experimental design

Male Lewis rats were divided among five experimental groups that each had VML injury in the left TA muscle: (1) no repair (negative treatment control), (2) repair with collagen hydrogel (0% minced graft), and repair with a (3) 25%, (4) 50%, or (5) 100% (positive treatment control) defect volume replacement of autologous minced grafts (e.g., for a 100 mg tissue defect, 100 mg of minced graft was replaced in the 100% group). The 25% and 50% volume graft replacement was suspended in collagen hydrogel; rat tail collagen I (Becton Dickinson) was diluted in 2.5·Dulbecco’s modified Eagle’s medium (Gibco) to create a 3 mg/mL solution and kept on ice while the minced graft was prepared. After mincing, the desired volume (0–50%) of muscle tissue was added to a well of a 48-well culture plate. A volume of collagen hydrogel was added to the well to bring the final volume to 400 μL (i.e., the volume of muscle tissue was calculated based on muscle density), which filled the defect area. The mixture was stirred to create a homogenous distribution of minced grafts within the construct. The collagen constructs were then allowed to crosslink at 37°C for 45 min, after which time the constructs were transplanted to the VML defect. Fascia and skin were closed by suturing and stapling each layer, respectively.

Animals

This work has been conducted in compliance with the Animal Welfare Act, the implementation of Animal Welfare Regulations and in accordance with the principles of the Guide for the Care and Use of Laboratory Animals. All animal procedures were approved by the United States Army Institute of Surgical Research Institutional Animal Care and Use Committee. Adult male Lewis rats (Harlan Laboratories) and GFP-Lewis rats (SD-Tg(GFP)Bal), Rat Resource and Research Center (RRRC), University of Missouri, Columbia, MO were housed in a vivarium accredited by the Association for Assessment and Accreditation of Laboratory Animal Care International, and provided with food and water ad libitum. A total of 34 rats were used in this study.

TA muscle VML injury

VML was surgically created in similar fashion to that previously reported.2,13,17 The TA and underlying extensor digitorum longus (EDL) muscles were separated using blunt dissection. A punch biopsy (6 mm) was performed through the middle third of the TA muscle, and the biopsied tissue was removed (~75 mg). Sustained release buprenorphine (72 h) was delivered (1.2 mg/kg SC) before surgery for pain.

Construct preparation

Autologous minced grafts were derived from the original TA muscle tissue that was excised to make the VML defect. The excised muscle tissue was minced into ~1 mm³ pieces. The minced grafts were then suspended in a collagen hydrogel; rat tail collagen I (Becton Dickinson) was diluted in 2.5× Dulbecco’s modified Eagle’s medium (Gibco) to create a 3 mg/mL solution and kept on ice while the minced graft was prepared. After mincing, the desired volume (0–50%) of muscle tissue was added to a well of a 48-well culture plate. A volume of collagen hydrogel was added to the well to bring the final volume to 400 μL (i.e., the volume of muscle tissue was calculated based on muscle density), which filled the defect area. The mixture was stirred to create a homogenous distribution of minced grafts within the construct. The collagen constructs were then allowed to crosslink at 37°C for ~45 min, after which time the constructs were transplanted to the VML defect. Fascia and skin were closed by suturing and stapling each layer, respectively.

In vivo functional assessment

Isolated TA muscle in vivo functional properties were measured in anesthetized rats (isoflurane 1.5–2.0%) in both legs as previously described.13 A nerve cuff...
with multistranded stainless steel (632; Cooner Wire) wire electrodes was implanted in each leg around the peroneal nerve. Legs were tested separately and in randomized order. The foot was strapped using silk surgical tape to a footplate attached to a dual-mode muscle lever system (Mod. 305b; Aurora Scientific, Inc.). The knee was secured on either side using a custom-made mounting system, and the knee and ankle were positioned at right angles. Optimal voltage (2–5 V) was set with a series of tetanic contractions (5–10 contractions; 150 Hz, 0.1 msec pulse width, 400 msec train). Then, a skin incision was made at the antero-lateral aspect of the ankle and the distal EDL muscle tendon and extensor hallucis longus muscle was isolated and severed above the retinaculum. The TA muscle and tendon, as well as the retinaculum were undisturbed. Four to five tetani were performed with a 1-min rest interval to enable torque stabilization. The contribution of the tenotomized EDL muscle was negligible in this testing system.\(^1\)\(^3\) Peak TA muscle isometric torque was determined with the ankle at a right angle (0°), assuming a moment arm of 3 mm.\(^1\)\(^8\) In vivo isometric torque values were normalized to body weight.\(^1\)\(^8\)

**Histology and immunohistology**

TA muscles were embedded in a talcum-based gel, frozen in 2-methylbutane (Fisher Scientific), and supercooled in liquid nitrogen using standard methodology previously reported.\(^1\)\(^7\) Frozen cross-sections (8 μm) were cut from the middle third of the TA muscle in the area where the original surgical defect was made. Sections were stained with hematoxylin and eosin (H&E), and oil red O, to assess tissue structure and cellularity, and fat deposition. Immunofluorescence was performed on sections probed for collagen I (1:500, AB755P; Millipore), sarcomeric myosin (MF20: 1:10; Hybridoma Bank), and nuclei (DAPI; 1:100; Invitrogen). Corresponding Alexaflour\(^4\)\(^8\) 488 and 596 labeled secondary antibodies (1:200–1:500; Invitrogen) were used for detection.

**Morphological assessment**

Whole muscle H&E cross-sections were analyzed for total fiber number and muscle fiber morphological characteristics. Total fiber number was analyzed by manually counting each fiber in an entire TA muscle cross-section (n = 3 muscles per group). Fiber morphological characteristics that included muscle fiber area, diameter (maximum Feret’s diameter), and roundness were assessed in nonoverlapping images taken from the defect or remaining muscle mass regions of three TA muscles per group. Measurements were made by manually outlining each fiber using ImageJ (NIH), similar to methodology previously described.\(^1\)\(^7\) Only fibers with sizes between 50 and 7000 μm\(^2\) and a circularity between 0.3 and 1.0 were included for analysis to remove the measurement of neurovascular bundles and oblique fibers.\(^2\)\(^0\) These criteria excluded less than 1% of fibers analyzed in the defect region. Fibers on the border of the image were also excluded from analysis.

**Statistical analyses**

Dependent variables were analyzed using one- and two-way analysis of variances (ANOVAs) and linear regression. Fisher’s LSD post-hoc means comparisons testing was performed when a significant ANOVA was observed. The distribution frequencies of muscle fiber areas were compared using a Chi-Square test. Muscle fiber area frequencies were binned to 5 classifications (<1000, 1001–2000, 2001–3000, 3001–4000, and >4000 μm\(^2\)) requiring a \(\chi^2\) of 9.49 to reach significance. Alpha was set at 0.05 for all tests. Values are listed as means±SEM (standard error of the mean). Statistical testing was performed using Microsoft Excel or Prism 6 for Mac (Graphpad).

**Results**

**Capacity of minced grafts to orchestrate de novo muscle fiber regeneration**

To determine whether minced graft-derived myogenic cells (i.e., primarily satellite cells delivered within their niche) contributed to de novo muscle fiber regeneration in a VML defect, grafts (100% replacement of excised tissue) were transplanted from syngeneic GFP donor rat TA muscles. Eight weeks postinjury, a large mass
of GFP+ muscle fibers, which co-labeled for sarcomeric myosin, were observed (Fig. 1). The GFP+ fibers were localized primarily within the defect area, though some GFP+ fibers were observed proximal and distal to the defect area, suggesting migration of transplanted myogenic cells (Fig. 1B–E).

FIG. 1. Contribution of donor-derived myogenic cells to de novo fiber regeneration after VML injury. TA muscle minced grafts derived from donor GFP-Lewis rats were transplanted to the site of VML injury at a 100% tissue replacement. (A) Longitudinal (scale bar = 1.0 cm) and (B–E) cross-sections (scale bars = 50 μm) from the remaining muscle mass (B, C) and defect area (D, E) were probed for sarcomeric myosin and GFP co-localization. TA, tibialis anterior; VML, volumetric muscle loss.

TA muscle strength gains with minced graft volume expansion
A potential limitation of the treatment of large clinical VML defects with autologous muscle is the availability of ample donor tissue and potential donor site complications and morbidity. Therefore, we sought to determine whether autologous minced muscle grafts could be volume expanded in a simple collagen hydrogel to achieve a similar functional recovery as a 100% minced graft replacement. To do so, in vivo neural-evoked torque of isolated TA muscles (no synergist muscle contribution) was measured (Fig. 2). In this model, VML injury without repair resulted in a 27% torque deficit 8 weeks postinjury compared with contralateral uninjured muscles. As a positive treatment control, a 100% minced graft tissue replacement restored 33% of the torque deficit. Volume expansion using 50% of the removed tissue achieved a similar (34%) restoration of the torque deficit; however,
further graft volume expansion to 25% replacement of the tissue lost did not improve torque beyond VML-injured nonrepaired muscles. Repair of the VML defect only with collagen hydrogel (no minced grafts) also did not improve torque compared with nonrepaired muscles.

Histological and immunohistological assessment of tissue regeneration
The remaining portion of the muscle after VML is not capable of endogenously regenerating the lost muscle tissue. This is reflected in TA muscle weights 8 weeks postinjury remaining 80 mg less than

FIG. 2. *In vivo* TA muscle torque production after treatment with autologous minced graft expansion constructs. Minced skeletal muscle grafts were transplanted to a TA muscle VML defect as a % of tissue defect volume. Grafts were delivered in a collagen hydrogel, reaching a similar volume (400 µL) per group. (A) Representative neural-evoked isometric tetanic torque waveforms (150 Hz; 400 msec train; 0.1 msec pulse width) and (B) group means ± SEM are presented. (C) TA muscle wet weight. At the time of functional testing, body weights were similar among groups (414.3 ± 3.3 g; ANOVA p = 0.30). † < Contralateral Control; * > all other VML injured groups, p < 0.05. ANOVA, analysis of variance.
contralateral uninjured muscles, since the initial defect weight was \( \sim 75 \) mg. However, TA muscle weights did not necessarily reflect functional recovery, since 25% graft repair presented fully recovered muscle weights despite no functional improvements (Fig. 2C). Therefore, histological analysis of the tissue was performed. VML-injured nonrepaired muscles presented few regenerated muscle fibers within the defect area, though the fibers in the remaining portion of the muscle appeared normal 8 weeks postinjury (Fig. 3A, B). Within the defect area, the density of muscle fibers appeared to increase with greater % volume graft repair (Fig. 3B). To confirm this observation, the area fraction of myosin and collagen was determined in images from the defect area (Fig. 3C). In comparison to contralateral uninjured muscles, all other groups exhibited lesser myosin and greater collagen I (Fig. 3E). However, the 50% and 100% graft repair promoted significantly greater myosin and lesser collagen I than all other experimental groups. In contrast, the presence of lipid deposition appeared to increase as the % volume graft repair decreased (Fig. 3D) repair with less than 100% graft repair resulting in greater lipid presence in the defect area (Fig. 3E).

**Analysis of de novo muscle fiber regeneration**

Strength gains after repair of VML are often attributed to *de novo* regeneration of muscle fibers throughout the literature, though this relationship has not been conclusively demonstrated. To assess *de novo* muscle fiber regeneration, the total number of muscle fibers in a cross-section from the middle third of the TA muscle was counted (Fig. 4A). Contralateral uninjured muscles comprised \( \sim 13,000 \) muscle fibers, and the

![FIG. 3. *De novo* muscle fiber regeneration within the defect area after treatment with autologous minced graft expansion constructs. Representative images from hematoxylin and eosin-stained cross-sections from the (A) remaining muscle mass area and (B) the defect area are presented. Cross-sections were probed for (C) myosin, collagen I and nuclei, and (D) oil red O. (E) Sections were semi-quantitatated for area fraction of myosin, collagen I, and oil red O. Values are means ± SEM. † ≠ Contralateral Control; * ≠ all other VML injured groups; ‡ > No Repair, \( p < 0.05 \). All scale bars are 50 μm.](http://online.liebertpub.com/doi/10.1089/biores.2015.0009)
VML injury reduced the number of muscle fibers by ~40% (~7800 fibers in VML-injured nonrepaired muscles) 8 weeks postinjury. Though not significant, 0% and 25% graft volume repair resulted in a ~15% and 7% increase in fiber number compared with nonrepaired muscles. Fifty percent graft repair significantly increased fiber number by ~20% compared with no repair. One hundred percent graft volume repair resulted in a significant increase in fiber number compared with all other VML injured muscles (e.g., ~41% > nonrepaired), but did not achieve full fiber restoration (i.e., ~14% < contralateral uninjured muscle). Notably, regression analysis demonstrated a strong relationship between fiber number and isometric torque among groups (Fig. 4B: $R^2 = 0.88$; $p = 0.005$), indicating de novo regeneration of muscle fibers as a primary determinant of recovery of strength with increasing % graft volume repair.

Morphological assessment of regenerated muscle fibers

Although the 50% and 100% graft volume repair groups presented significantly greater fiber number, the morphology of regenerated fibers appeared less organized than uninjured tissue (Fig. 3). Therefore, muscle fiber morphological characteristics in the remaining and defect areas were analyzed (Fig. 5). In the remaining muscle area, there were no differences among contralateral uninjured, 50% and 100% graft groups for average fiber area, fiber diameter, or fiber roundness. In addition, the distribution of fiber area frequencies within the remaining muscle area of 50% and 100% graft groups was similar to uninjured contralateral control muscles ($\chi^2 = 5.70$ and 3.49, respectively). Within the defect area, both the 50% and 100% graft groups presented regenerated fibers with a smaller fiber area and diameter (Fig. 5B, C) and lesser roundness (Fig. 5D), suggesting an increased pennation angle. In addition, the distribution of fiber area frequencies within the defect area of 50% and 100% graft groups was significantly different from uninjured contralateral control muscles ($\chi^2 = 796$ and 544, respectively). Therefore, the fibers regenerated after graft transplantation are smaller and disoriented compared with uninjured muscles.

**Discussion**

The most important finding of this study is that expanded autologous minced muscle grafts (50% less tissue) promote a similar functional recovery as a 100% graft repair. However, regenerative expansion was not fully supported in the collagen hydrogel, since only approximately half the number of fibers regenerated de novo with a 50% graft repair, compared with 100% repair. That being said, we conclusively demonstrate using GFP+ grafts that myogenic cells derived (e.g., satellite cells) from the minced grafts survive and directly contribute to significant de novo fiber regeneration within the defect area. Though the muscle tissue regenerated via minced grafts presented greater collagen content and altered histomorphometry compared with uninjured muscle, the fact that the 50% and 100% graft replacement promoted significant de novo muscle fiber regeneration supports the fact that an autologous muscle tissue-based approach may form a foundation on which restoration of functional muscle mass may be achieved after VML injury.
There was a corresponding decline in the number of fibers regenerated between 100% and 50% graft repair, and a precipitous decline from 50% to 25% graft repair, indicating that the collagen hydrogel did not provide ample support for expansion. One explanation is that the mechanical stability of the gel composition used (3 mg/mL collagen I) was not biomimetic of skeletal muscle. This possibility is in line with a recent paper, indicating that this formulation of collagen hydrogel supported ample vascularization after delivery of microvascular fragments to VML injured muscle, but did not support significant myogenesis. However, the biomaterial may not be the only limiting factor. Previously vital grafts co-delivered with devitalized muscle scaffolds (50% of each), which maintain many properties of muscle tissue, also did not appear to support graft expansion. Therefore, it is possible that de novo muscle tissue regeneration may improve with a myoconductive scaffold that not only provides a more informative matrix both structurally and compositionally (e.g., inclusion of basal lamina) but also may require adjunct therapies such as alternative
myogenic cells, growth factors/small molecules, and/or physical rehabilitation. 

Autologous minced graft expansion and ECM-mediated regeneration may be primarily restricted by a similar mechanism, as both rely on satellite cell migration for de novo fiber regeneration. Previously, we and other labs have demonstrated that ECMs derived from muscle and other tissues (e.g., urinary bladder matrix & small intestine submucosa) do not promote appreciable muscle fiber regeneration beyond ~0.5 mm from the remaining muscle mass, if at all. The collagen hydrogel used in this study was also ineffective in promoting muscle fiber regeneration. Collectively, these findings indicate that providing a matrix alone is not sufficient to promote appreciable muscle fiber regeneration in a VML defect. In a recent investigation, the restricted de novo muscle fiber regeneration mediated by ECMs was partly attributed to an impaired migration of satellite cells (Pax7) from the remaining portion of the host muscle to the scaffolding in the defect area. The findings of this study that (1) minced graft-derived myogenic cells directly mediate de novo fiber regeneration, (2) de novo fiber regeneration is directly related to the amount of minced graft replacement (i.e., myogenic cells), and (3) collagen–hydrogel alone does not support fiber regeneration, as well as previous reports that devitalized scaffolds (which maintain many properties of muscle tissue) and small muscle grafts devoid of functional satellite cells (Pax7 knockout) do not promote regeneration, highlight that inclusion of myogenic cells (satellite cells) is necessary for de novo fiber regeneration. Significant intramuscular satellite cell migration has been observed in mice with focal muscle injury and, to that end, we observed putative satellite cell migration from minced grafts to the remaining muscle mass in this study, which indicates that the inherent ability of satellite cells to migrate is not the limiting factor per se. Instead, the pro-fibrotic environment created by VML likely restricts satellite cell migration during graft expansion and ECM-mediated regeneration, and, therefore, further exploration of approaches to improve myogenic cell migration by attenuating ensuing inflammatory and fibrotic responses is warranted.

A strong correlation existed among muscle fiber number and strength among VML injured and repaired groups, indicating that de novo muscle fiber regeneration is a primary determinant of strength recovery. However, inspection of this relationship between 100% and 50% minced graft groups indicates multiple mechanisms of functional recovery. VML studies conducting functional assessments have reported strength gains related to either appreciable de novo fiber regeneration within the defect (increased force production) or ECM deposition within the defect that improves transmission of force produced by the remaining muscle mass. Given that the 50% graft group regenerated only approximately half of the muscle fibers in the 100% graft group, it is possible that the 50% graft group may be evidence of both mechanisms, that is, improved force production and transmission. However, significant strength improvements in this study were not observed in groups that did not present significant de novo fiber regeneration but presented collagen deposition in the defect (e.g., 25% graft), suggesting that other unidentified mechanisms may underlie the equivalent strength improvements with 50% graft repair.

Minced graft transplantation (50% and 100%) promoted significant de novo fiber regeneration; however, the regenerated tissue composition and fiber morphology was altered compared with uninjured muscle. Notably, the regenerated tissue presented greater collagen and lipid deposition (50% but not 100%) and the fibers had a smaller average cross-sectional area than in uninjured muscle. These characteristics likely contributed to a lesser than expected functional recovery estimated from the number of fibers regenerated. In addition, the roundness of the fibers was reduced, suggesting that the regenerated fibers were pennated more than in uninjured muscle. It is unclear in this study whether this is a positive or negative adaptation. While an increased pennation angle reduces the contribution of a fiber’s force to net muscle torque, increasing pennation angle also allows for more muscle fibers to employ a given area and, therefore, may ultimately result in an increased net torque production.

Despite the altered histomorphology, it is important to note that the muscle tissue regeneration after VML repair using 50% or 100% graft repair appears to be greater than with decellularized ECM-based approaches. Moreover, the re-establishment of a portion of the lost muscle tissue provides the opportunity for further muscle tissue augmentation using physical rehabilitation to improve regenerative and physiological outcomes that would not otherwise be possible.

In addition to efficacy and safety, donor tissue availability is recognized as a potential limitation to clinical
transplantation of minced grafts for the repair of large VML defects in large muscles (e.g., human quadriceps; $\sim 1\,L^{12}$), but not necessarily in smaller muscles (e.g., human TA; $0.15\,L^{43}$). For example, the quadriceps with a 70% VML would require 0.7 (740), 0.35 (370), and 0.18 (190) L (or grams with muscle density of 1.056 g/mL) of donor tissue for a 100%, 50%, and 25% tissue replacement; the TA muscle with a 70% VML would require 0.11 (116), 0.053 (56), and 0.026 (27) L (grams) of donor tissue, respectively. The findings of this study indicate that a standard latissimus dorsi flap, which provides $\sim 0.26\,L$ ($\sim 275$ grams; assuming $1\,cm$ thickness) of muscle tissue,\textsuperscript{44,45} approaches enough tissue to support minced graft expansion for repair of large muscles and fully supports applications in smaller muscles.

The ultimate goal of this line of research is to regenerate muscle tissue and restore functional capacity after clinical VML. Since VML is heterogenous among patients,\textsuperscript{1,13,16,22} clinicians would benefit from a variety of surgical and regenerative treatment options. Xenogeneic decellularized extracellular matrices, as have recently been tested in the clinic,\textsuperscript{16,22} embody an ideal treatment for VML in that no autologous donor tissue is required. Transplantation of ECMs to VML defects in six patients has promoted modest gains in strength with no reported adverse outcomes.\textsuperscript{16,22} However, the histological data reported from different laboratories using a variety of ECMs in various VML animal models do not support ECMs role in promoting appreciable muscle fiber regeneration.\textsuperscript{17,22,30–33,46} On the other hand, minced grafts directly contribute to and orchestrate appreciable \textit{de novo} muscle regeneration,\textsuperscript{10,13,26,47,48} but the amount of autologous donor muscle tissue required to repair large VML may be limited. To address this potential limitation, the capacity of minced grafts for volume expansion in collagen was tested. The results of the study did not support graft expansion in this biomaterial, but indicated that that use of 50% less tissue promoted similar functional improvements as 100% graft repair. Among other barriers to regenerate large volumes of skeletal muscle such as revascularization (e.g., Pilia et al.\textsuperscript{15}), restricted satellite cell migration is likely a significant barrier to minced graft expansion and ECM-mediated regeneration.

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Author Disclosure Statement

No competing financial interests exist.

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