



Demonstrating collagen tendon fibril segments involvement in intrinsic tendon repair

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ABSTRACT

Severed tendons can undergo regenerative healing, intrinsic tendon repair. Fibrillogenesis of chick tendon involves “collagen fibril segments” (CFS), which are the building blocks of collagen fibers that make up tendon fascicles. The CFS are 10.5 micron in length, composed of tropocollagen monomers arranged in parallel arrays. Rather than incorporating single tropocollagen molecules into growing collagen fibers, incorporating large CFS units is the mechanism for generating collagen fibers. Is intrinsic tendon repair through the reestablishment of tendon embryogenesis? Gentamicin treated 10-day-old chick embryo tendons released CFS were fluorescently tagged with Rhodamine (Rh). Organ cultured severed 14-day-old embryo tendon explants received Rh tagged CFS. At day 4 auto fluorescent tagged CFS were identified at the severed tendon ends by fluorescent microscopy. Accumulation of fluorescent tagged CFS was exclusively localized to the severed ends of tendon explants. Parallels between collagen fiber growth during embryonic fibrillogenesis and tendon repair reveal CFS incorporation is responsible for collagen fibers growth. CFS incorporation into frayed collagen fibers from severed tendons is the proposed mechanism for intrinsic tendon repair, which is an example of regenerative repair.

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Introduction

Tendons are rigid cables that connect bone to muscle. The parallel arrangement of collagen fibers within fascicles is accountable for the rigidity of these cables (Jozsa et al., 1991). The major collagen type within tendon fibers is type I collagen (Birk et al., 1989a). Robert Trelstad and coworkers studies of chick embryo tendon development showed banded collagen structures within the chick embryonic tendon fibroblast cytoplasmic-like vacuoles (Trelstad and Hayashi, 1979). These banded collagen fibril structures are not ingested fibrils destined for breakdown, rather they are newly synthesized collagen fibrils that will be utilized for generating new collagen fibers. The banded collagen fibrils are not in typical cytoplasmic vacuole, such as Golgi vesicles or lysosomes, but are specialized cell plasma membrane clefts that are specialized extracellular compartments. Newly synthesized procollagen released from the rough endoplasmic reticulum (RER) is cleaved into tropocollagen within these clefts (Birk and Trelstad, 1986). Tropocollagen is not directly incorporated into collagen fibers, but is initially incorporated into Collagen Fibril Segments (CFS) that are incorporated into expanding collagen fibers (Birk et al., 1989a; Franchi et al., 2007).

Damaged flexor tendon repair can terminate in scar, like other soft tissues, referred to as extrinsic tendon repair (Manske et al., 1985). Another option for tendon repair is intrinsic tendon repair, where regenerative healing occurs by a process that simulates embryonic fibrillogenesis. The termination of tendon repair by intrinsic tendon repair is a regenerated tendon. Extrinsic tendon repair is typical wound healing, characterized by the initial intrusion of inflammatory cells followed by fibroblasts derived from neighboring tissues, terminating in the deposition of a new disorganized connective tissue matrix, a scar. Such a repaired tendon may have deficiencies in strength as well as the ability to glide within its sheath. A major difference separating extrinsic and intrinsic tendon repair is the organization of the newly deposited connective tissue within tendon fascicles. In intrinsic tendon repair the fascicle collagen fibers run in parallel arrays, indistinguishable from the orientation of collagen fibers in uninjured tendon. The outcome of extrinsic tendon repair is disrupted fascicles and newly deposited connective tissue organized in random arrays showing variations in the collagen fiber diameters (Ehrlich et al., 2005).

Success in studying repair in isolated organs such as tendon is advanced by utilizing embryonic derived tissues. Because these tissues flourish *in vivo* in an environment deficient in oxygen, they can retain their viability in stationary organ culture. Embryonic chick explants remain viable in stationary organ culture for long periods of time in ambient air and 5% CO₂ (Fell, 1956). Unlike metabolically active fibroblasts typical of the proliferative phase of wound repair, intrinsic tendon repair is completed in the absence of angiogenesis and developing a new blood supply. The restoration of normal

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function to a severed tendon by intrinsic tendon repair requires endogenous tendon fibroblasts bridging the gap between the severed tendon ends with the deposition of new connective tissue matrix. It is proposed that released CFS from severed collagen fibers generate a pool of CFS that supply a portion of the building blocks utilized at the healing site (Ingraham et al., 2003). Here the study of intrinsic tendon repair employs the introduction of fluorescent tagged CFS to 14 day old severed chick tendon explants maintained in organ culture. The pattern of fluorescence that appears within severed tendon explants is evaluated 4 days later by fluorescent microscopy.

Method

Isolated tendons, harvested from fertilized chicken eggs, were obtained from the GemWillow Farm (Grantville, PA). Eggs were incubated in a poultry incubator. Tendons from 10 day old chick embryos were isolated by pulling on the chick's toes. The freed tendons were suspended in 5 ml of phosphate buffered saline (PBS) with gentamicin at 1 mg/mL (Sigma Chemical Co., St. Louis, MO) within a sealed 12 mL centrifuge tube for 90 days at 4 °C with agitation. The tendon-antibiotic suspension was vortexed, then centrifuged at 10,000 $\times$ g for 2 min and the supernatant saved. The supernatant fraction contained the CFS and the pellet fraction contained insoluble tissues that were discarded.

Transmission electron microscopy (TEM) confirmed the identity of CFS. Briefly a drop of supernatant was placed on a formvar coated copper grid for 10 min, followed by staining with 2% phosphotungstic acid and then viewed with a Philips 400 Electron Microscope. The CFS from the supernatant were fluorescently tagged with Lissamine rhodamine (Rh) B-200 on celite (USB Corp., Cleveland, OH). To obtain a minimal number of Rh covalently bound to the CFS so that steric hindrance was minimized, the following procedure was followed. The CFS suspension was placed on ice and the pH adjusted to 7.0. About 10 mg of Lissamine Rh on celite was added, the tube briefly vortexed then immediately centrifuged at 5000 $\times$ g for 1 min, the supernatant saved and the celite rich pellet discarded. The minimally tagged CFS are referred to as Rh-FS. A high density of Rh bound onto CFS would be expected to interfere with the association of Rh-FS with cells as well as with growing collagen fibers at tendon wound edges.

Chick embryos at 14 days were removed from their shells and their leg tendons harvested by pulling on their toes with a forceps. Isolated tendons were placed in Dulbecco's modification of Eagles medium (DMEM) with 10% new born bovine serum and 5 μ g gentamicin/mL, referred to as complete DMEM. Tendons were cut into 5 mm explants. Four tendon explants were placed on a Gelman Sciences filter membrane (Pall Corp. Port Washington, NY), which was centered on a stainless steel mesh triangle suspended over the central well of a 60 $\times$ 15 mm Falcon #3037 Organ Culture Dish (BD Biosciences, Lincoln Park, NJ). The central well was filled with 1.0 ml complete DMEM, and the surrounding outer well received 2.5 ml of sterile water. Surface tension kept the tendon explants covered with thin film of culture medium, allowing a gas exchange between the ambient air and tendon explants. Each tendon explant was severed completely with a scalpel and the cut ends re-approximated by carefully bringing them together. The adherence of the tendon explant to the underlying membrane surface was sufficient for maintaining the 2 severed explants lengths in a fixed position. The fixation of the tendon lengths, which reestablished the continuity of the severed tendon, mimicked the stabilization of a suture repaired severed tendon. Rh-FS were introduced to wounded tendons by directly pipetting a 1 μ L suspension of Rh-FS onto tendon explants. Tendons were maintained in an incubator set at 37 °C with 5% CO₂ in a water saturated atmosphere.

Wounded 14 day chick embryo tendon explants maintained in organ culture with Rh-FS for 4 days were prepared for fluorescence microscopy. At 4 days tendon explants were carefully harvested by leaving the explant attached to the underlying membrane so as not to

disrupt the repair region between explants. The healing tendon explants with associated membrane were fixed with 4% paraformaldehyde for 5 min and then rinsed with PBS. A pair of tendon explants was incubated with DAPI (Invitrogen, Carlsbad, CA) for 10 min to fluorescently stain nuclei blue. Unstained and DAPI blue stained tendon explants attached to membrane were transferred to glass slides, a drop of GelMount (Biomedica, Foster City, CA) places on top, followed with a coverslip. Slides were viewed with a Zeiss Inverted Fluorescent microscope equipped with epifluorescence and images recorded with a digital camera.

Results

CFS were released by incubating isolated 10-day chick embryo tendons with a high concentration of gentamicin in the cold for 3 months. A centrifugal force of 10,000 $\times$ g for 2 min was not sufficient to pellet the CFS, but sufficient to remove tendons and other debris. As shown in Fig. 1, a CFS identified by TEM imaging, showed a long CFS had a tapered end, which is the expected morphology of a CFS (Kadler et al., 1996). The inset in Fig. 2 taken at higher magnification showed the typical collagen like banding pattern along the length of the isolated CFS.

A 1 μ L Rh-FS suspension was added to organ cultured severed tendon explants and viewed at 4 days with a fluorescent microscope. The accumulation of Rh-FS with the tendon explants occurred in a specific manner. Minimal red fluorescence was localized along the uninjured intact length of tendon explants. Red fluorescence appeared at the distal cut ends of the explants (not shown) as well as within the gap between wound edges (Fig. 2). The red auto-fluorescence within the gap between the approximated wound edges of tendon explants presented in Fig. 2 demonstrated the specificity of that fluorescence accumulation. In Fig. 2, panels A and B, the typical pattern of red auto-fluorescence that filled the gap between the severed edges, showed a collagen fiber pattern that had organization. The accumulation of Rh-FS was limited to the gap and cut ends of the tendon explants. In addition to red auto-fluorescent fibers, blue fluorescent stained nuclei from tendon fibroblasts are presented in Fig. 2, panels C and D. DAPI stained tendon fibroblast nuclei identified the localization of tendon fibroblasts in the repair region of severed tendon explants. The blue fluorescent tendon fibroblasts were associated with red fluorescent Rh-FS fibers located at the wound edges. At the center of the gap between the severed tendon lengths, where there was the absence of red auto-fluorescent Rh-FS collagen fibers that was the deficiency of tendon fibroblasts. The absence of blue fluorescent nuclei in the central gap region confirmed the absence of tendon fibroblasts.

Discussion

When viewed at 4 days, introducing fluorescent tagged isolated collagen CFS to wounded tendon explants in organ culture produces a localized intense red fluorescence in the wound gap. The concept is the accumulation of fluorescent CFS at the wound gap results from their translocation by tendon fibroblasts to their eventual entrapment within the gap between a severed tendon explant. CFS, the building

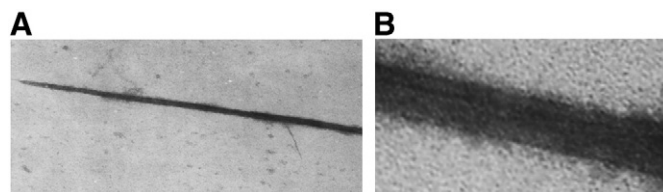


Fig. 1. A TEM of a fibril segment isolated from the supernatant of a long term incubation of a 10 day chick embryo tendon with 1 mg/mL gentamicin antibiotic. Released fibril segments were isolated by centrifugation and an aliquot subjected to TEM. The inset shows a higher power image with a banding pattern associated with a fibril segment.

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