

Tendon's ultrastructure

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Summary

The structure of a tendon is an important example of complexity of ECM three-dimensional organization. The extracellular matrix (ECM) is a macromolecular network with both structural and regulatory functions. ECM components belong to four major types of macromolecules: the collagens, elastin, proteoglycans, and noncollagenous glycoproteins. Tendons are made by a fibrous, compact connective tissue that connect muscle to bone designed to transmit forces and withstand tension during muscle contraction. Here we show the ultrastructural features of tendon's components.

KEY WORDS: tendon, collagen, ultrastructure, extracellular matrix.

Introduction

The extracellular matrix (ECM) is a macromolecular network with both structural and regulatory functions.

Indeed, ECM furnishes mechanical and biochemical signals that cooperate in the integrated control of cell proliferation, survival, migration, and differentiation¹⁻³. ECM components belong to four major types of macromolecules: the collagens, elastin, proteoglycans, and noncollagenous glycoproteins. The most copious proteins in the ECM are members of the collagen family. The collagens are involved in the formation of ECM fibrillar and microfibrillar networks and play a key role in determining tissue-specific mechanical properties. Collagens can be categorized into fibril forming collagens (types I, II, III, V, XI), fibril-associated collagens (FACITs; types IX, XII, XIV, XIX, XX, XXI), hexagonal network-forming collagens (types VIII and X), microfibril-forming collagen (type VI), basement membrane collagen (type IV), anchoring fibril-forming collagen (type VII), transmembrane collagens (types XIII and XVII), and other collagens with peculiar functions (types XV, XVI, XVIII)⁴⁻⁶. Other ECM components that affects tissue mechanical properties are elastin, which is the main constituent of ECM elastic fibers, and the proteoglycans which are composed of a protein core linked to glycosaminoglycan (GAG) side chains⁷. A different group of ECM components is represented by ECM glycoproteins including laminins and entactin/nidogen which are the major glycoprotein components of the basement membrane^{1,8}, fibronectin which is found as a soluble dimer in plasma and as an insoluble fibrillar polymer of interstitial ECM and basement membranes^{1,8} and tenascins, which has adhesive and anti-adhesive effects on cell-matrix interactions^{1,9}.

It has been supposed that collagens have a significant role in the evolution of metazoans. Indeed, the fibril-forming collagens may correspond to the most ancient type of collagen in the metazoan and the variety of these fibrils has enhanced during the development of the chordates¹⁰.

Tendon's ultrastructure and ECM

The structure of a tendon is an important example of complexity of ECM three-dimensional organization. Tendons are made by a fibrous, compact connective tissue that connect muscle to bone designed to transmit forces and withstand tension during muscle contraction^{11,12}. The apparently simple tendon's structure and composition provide both rigidity and flexibility. This ability is based on tendon non-linear, viscous-elastic, anisotropic and heterogeneous mechanical properties¹¹⁻¹³.

Similarly to other tissues there is a clear relationship between structure and function of the tendon¹³. Indeed, different tissue microenvironments provide specific

characteristics to the different three-dimensional organization of the ECM since embryogenesis^{1,3}. The organization of the ECM in the tendon is peculiar as well as within different tendons. The process of tendon's embryonic structural development differs by location and type of tendon¹⁴. In the first stage, for example, tendon progenitor cells originate from the primary germ layer compartments along with progenitors for cartilage and muscle¹⁴. This process is different for longer limb tendons as opposed to short trunk tendons¹⁴. The dialogues between different cells (fibroblasts, muscle and cartilage cells) or better the dialects between different cells mediated by various signaling cascades dependent on FGFs and TGF-beta are used at decisive stages of development to make specific the phases of induction, organization, aggregation or differentiation of cells¹⁴. The structure of the tendon is so complex that it is very difficult to maintain its function in the healing since this process is reparative rather than regenerative in the adult^{15,16}. A tendon consists of 70% of water and 30% of dry mass, which is composed by 60-80% of type I collagen and 2% of elastin. Among collagens, the most abundant component is collagen type I (95%), while type III and type V collagens represent the remaining 5% of the total collagens¹⁶. Tendon's collagen's is organized in multi-hierarchical structures including fibrils, fibers (primary bundles), fascicles (secondary boundless)^{15,16} (Fig. 1). The fibrillogenesis in the form of small diameter fibrils begins in embryogenesis and continues after birth with the assembly of collagen type I molecules, followed by linear and lateral growth and collagen interactions with proteins such as other collagens and proteoglycans¹⁴. In particular, the linear and lateral growth is determined by a variety of molecules including other collagens (type III, V, XI, XII and XIV) which are expressed in a variable manner. Electron microscopy of longitudinally oriented tendon specimen shows a parallel arrangement of collagen fib-

rils. The ratio of collagen fibril sizes in human tendon is variable from 1,750 to 600 and 100 A diameter^{13,17}. However, it is difficult to link the size of collagen fibrils with different zone of the tendon¹⁷. The collagen type III plays a key role in regulating fibrillogenesis and extensibility of the tendon and its production slowly decreases in the development to reappear in appreciable amount after injury in the healing process¹⁴⁻¹⁶.

Collagen type V and XI are associated to collagen type I and II, respectively and they determine their quantity and quality^{1,2,4,14}. It must be stressed that collagen type V exists in the form of heterotrimers sequestered in the fibrils of abundant collagen type I and in the form of homodimers as thin filament which can act in the ECM as a molecular linker between collagen fibrils or macromolecules depending on their respective distribution in different tissues^{18,19}. The location of collagen type V in the perivascular tissues have made it as a "sequestered antigen" which might be recognized by the immune system in a variety of inflammatory processes and therefore might induce autoimmune diseases^{3,20} (Fig. 2).

Type XII and XIV collagens are fibril-associated collagens with interrupted triple helices (FACIT)⁴⁻⁶. Tendons in additions to collagen contain proteoglycans, glycosaminoglycans and glycoproteins including fibronectin, trombospondin and tenascin-C immersed in different composition of ECM lying cellular elements represented (90/95%) by tenoblasts and tenocytes¹³ (Fig. 3). The tenoblasts show a prominent rough endoplasmic reticulum, as well as a developed Golgi complex. The remaining 5-10% of the cells consists of chondrocytes, synovial and vascular cells¹³.

Tendons are responsible for the connection between bone and muscle at their ends. The junction is called enthesis, which can be divided in fibrous enthesis and fibrocartilaginous enthesis. In tendons, the fibrocartilage is arranged in region compressed against bony or fibrous pulleys²¹. Fibrocartilage is an avascular tissue whose

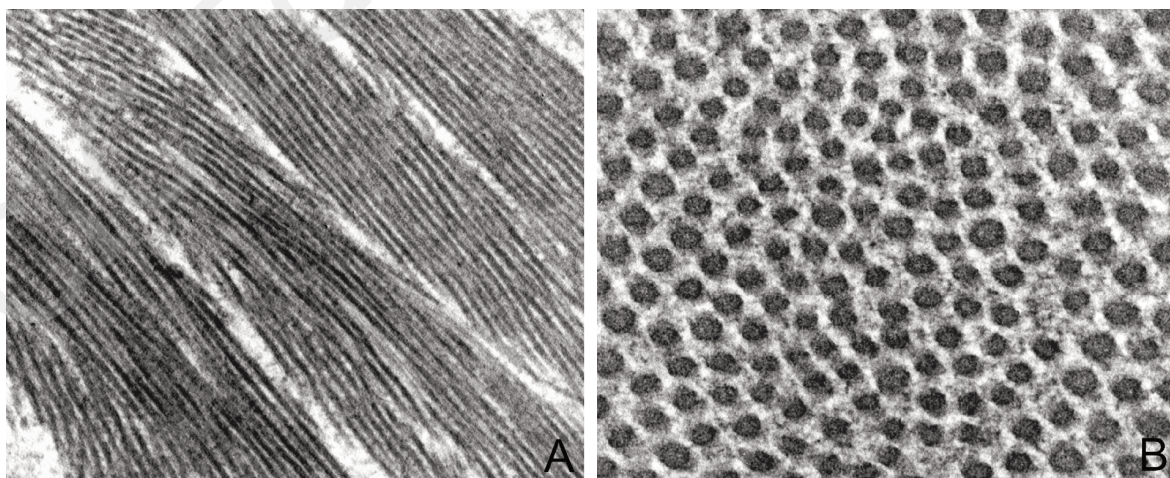


Figure 1. Ultrastructure features of ECM in tendon. A: Collagen I is the mainly constituent of tendons and creates highly aligned fibrils organized as fibers oriented longitudinally, transversally and also crossing each others, thus conferring a great resistance to different forces (O.M. x7900). B: Collagen I fibrils are in contact to thin fibrils of collagen V and to proteoglycans and glycoproteins (O.M. x12500).

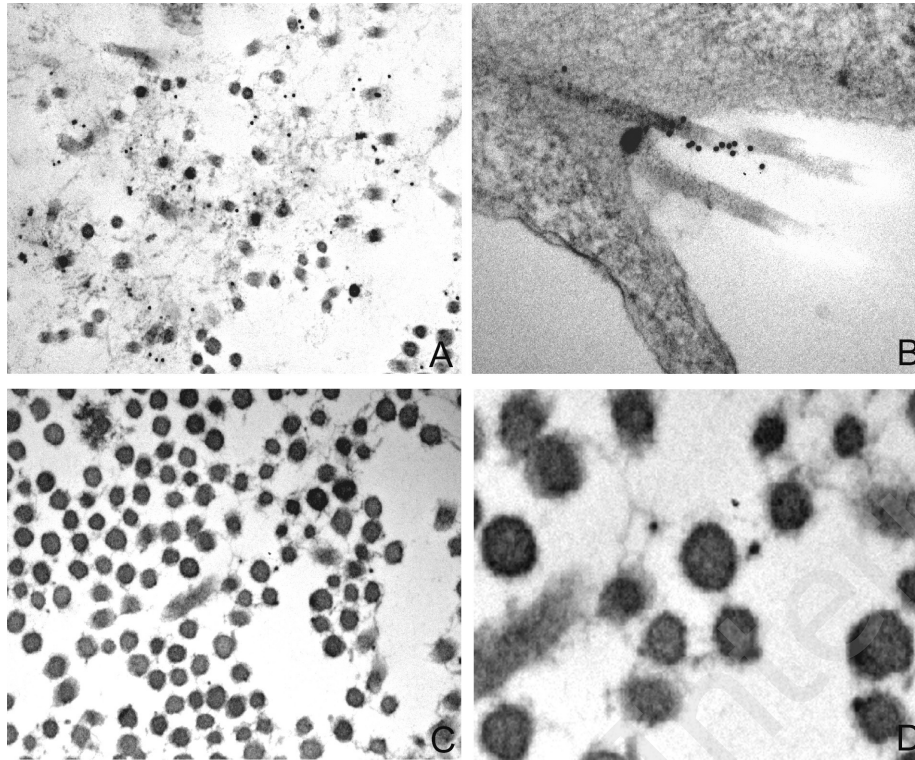


Figure 2. Immunolocalization of type V collagen in different tissues. Immunostaining with a specific monoclonal antibody against collagen type V and colloidal gold conjugated goat anti-mouse (O.M. x 11500, x7900, x18000, x 45000 respectively). A: desmoplasia in breast carcinoma; B: fibroblasts tissue culture; C-D: Degenerative tendon scar

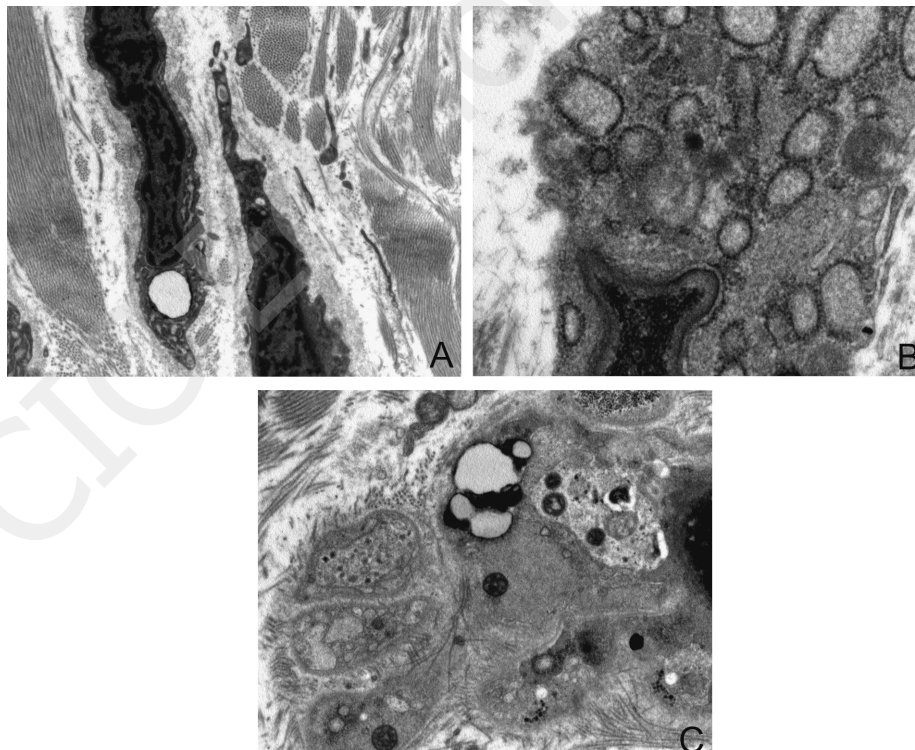


Figure 3. Ultrastructure features of tenocytes and tenoblasts. The cellular population of tendon is mainly represented by tenocytes: tenocytes are very elongated cells with ovoid and very long nuclei, occupying the entire length of the cell, with a high nucleus-to-cytoplasm ratio. The cellular process are long and thin and in direct contact with extracellular matrix components (A). In the cytoplasm of tenoblasts is often visible a well developed rough endoplasmic reticulum with dilated cisternae (B), few mitochondria, primary and secondary lysosomes, vacuoles, phagosomes and glycogen (C) (O.M. x 3900, x7900, x3900).

cells resemble, at the ultrastructural examination, chondrocytes with a prominent rough endoplasmic reticulum, glycogen granules and lipid droplets and intermediate filaments²¹. The cells synthesize collagens (type I, II, III, V, XII, XII), proteoglycans (aggrecan, versican, decorin, biglycan, lumican, fibromodulin), glycoprotein and in par-

ticular tenascin-C by mechanical strain. The tendons is covered by the epitenon, a delicate, loose connective-tissue sheath containing blood and lymphatic vessels, and nerves. The epitenon expands deeply between the tertiary bundles as the endotenon, made by connective tissue surrounding each fiber¹³ (Figs. 4 and 5).

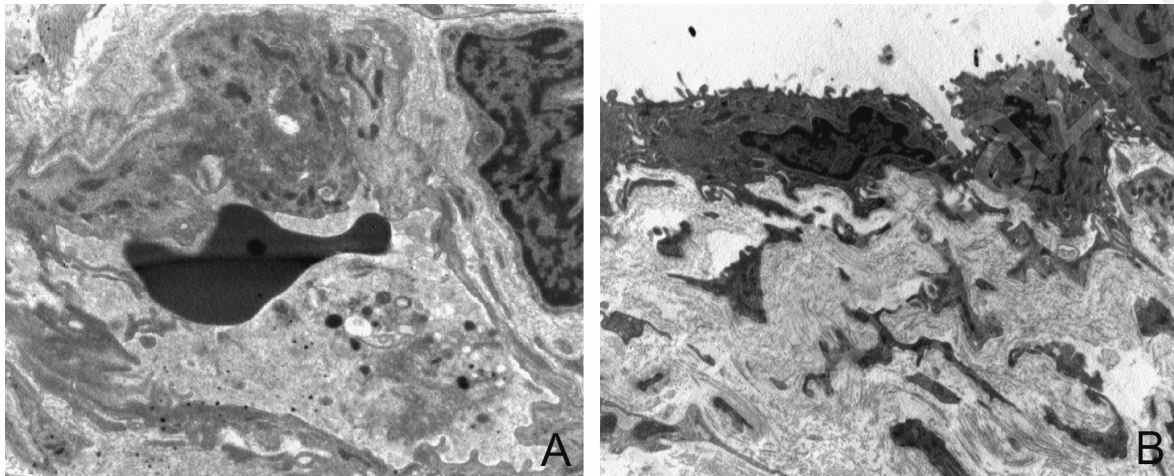


Figure 4. Ultrastructure features of epitenon. Epitenon is a delicate, loose connective-tissue sheath containing blood and lymphatic vessels. A-B: Endothelial cells show cell membrane protrusion and several granules and pinocytic vesicles in the cytoplasm. Endothelial cells are surrounded by pericytes and show a well constituted basal membrane (O.M. x4900, x3900).

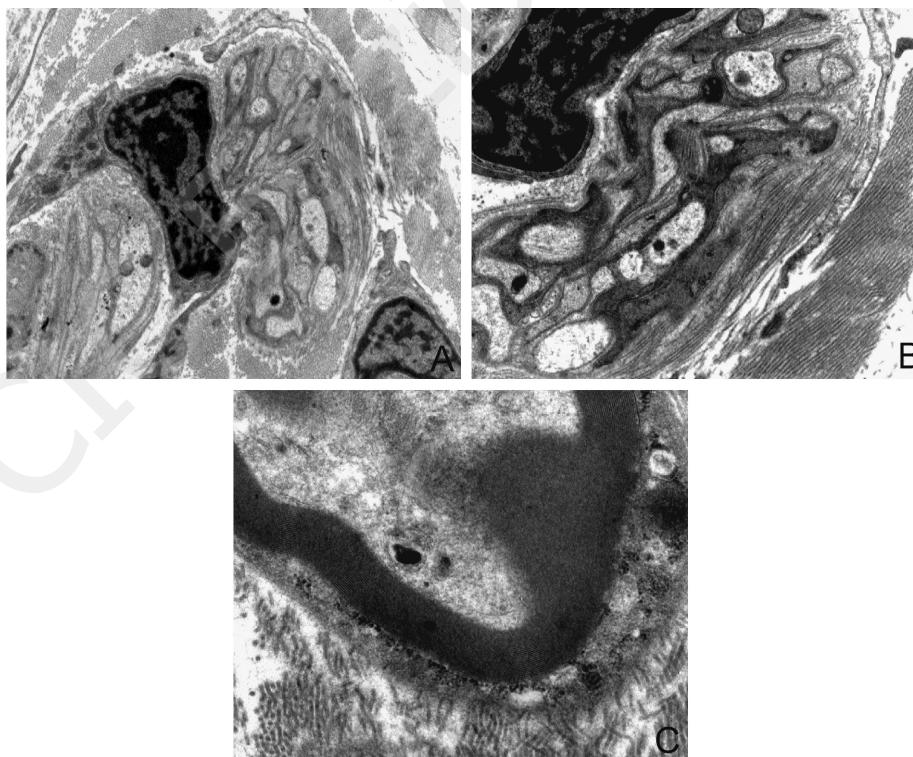


Figure 5. Ultrastructure features of nerves in the epitenon. Unmyelinated (A-B) and myelinated (C) fibers surrounded by cytoplasmic elongation of Schwann cell (O.M. x3900, x5500, x7500).

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