

Analysis of the Acellular Matrix, Growth Factors, and Cytokines Present in ArthroFlex[®]

OBJECTIVE

Identify the extracellular matrix (ECM) components, growth factors, and cytokines present in ArthroFlex, a decellularized, sterile human dermal allograft.

INTRODUCTION

Surgically reattached tendons often heal improperly by forming weaker fibrous connections between the muscle and bone; they do not recover full mechanical strength and display a high rate of re-tear¹⁻⁴. Acellular human dermis can aid in the repair of torn tendons by providing supplemental strength and structural integrity to the reattachment site. Human dermis is a complex tissue containing various extracellular matrix molecules, growth factors, and cytokines⁵. The purpose of this study was to ensure that ArthroFlex, a minimally manipulated human dermis product, retains the biological components that provide supplemental strength and can structurally support the repair of reattached tendons.

METHODOLOGY

Protein Analysis[†]:

Samples were solubilized in a detergent solution assisted by mechanical homogenization followed by protein separation based on molecular size. Subsequently, the separated proteins were enzymatically fragmented, after which the amino acid sequence of each fragment was determined by liquid chromatography with tandem mass spectrometry (LC-MS/MS)[‡]. The amino acid sequences of each fragment were then compared against a database containing the sequences of known proteins to determine the corresponding protein for each fragment. From this comparison, a list of the proteins that corresponded to each fragment was generated[†]. This list was mined for extracellular matrix (ECM) components, growth factors, and cytokines to create a table of proteins whose fragments were found in the ArthroFlex sample. Additionally, some components were further verified or identified by immunohistochemical staining and by enzyme-linked immunosorbent assay (data on file at LifeNet Health).

CONCLUSION

The results of this study indicate that ArthroFlex retains ECM components, matrikines, growth factors, and cytokines consistent with minimally manipulated human tissue and relevant to the structural support of damaged soft tissue. ArthroFlex provides the collagens that supplement structural integrity and mechanical strength to surgically reattached tendons, aiding in the prevention of a re-tear.

FINDINGS

Fragments of the following proteins were found in ArthroFlex by LC-MS/MS:

| Collagens | GF-binding ECM | Additional ECM | Matrikines | Growth Factors | Cytokines |
|-------------|--|--------------------|-------------|----------------|-----------|
| Type I | Heparan Sulfate Proteoglycan (HSPG) | Elastin | Tenascin-C | BMP6 | IL1a |
| Type III | Chondroitin Sulfate Proteoglycan (CSPG) | Nidogen (Entactin) | Laminins | CTGF | IL1b |
| Type IV | Perlecan (HSPG2) | Keratin | Decorin | EGF | IL2 |
| Type V | Aggrecan | | Endostatin | HGF | IL5 |
| Type VI | Lumican | | Pentastatin | PDGFD | IL9 |
| Type VII | Versican | | Tumstatin | TGFB1 | IL22b |
| Type VIII | Glypican | | Elastokines | VEGFA | IL25 |
| Type XII | Syndecan | | | VEGFD (FIGF) | IL27 |
| Type XIV | Tenascin (C & N) | | | | IL32 |
| Type XVII | Thrombospondin 2 | | | | TNF |
| Type XVIII | Dermatopontin | | | | |
| Type XX | Decorin | | | | |
| Type XXI | Vitronectin | | | | |
| Type XXIII | Laminin (α 1-5, β 1-3, γ 1&3) | | | | |
| Type XXVII | Fibrinogen (Fibrin precursor) | | | | |
| Type XXVIII | | | | | |

DISCUSSION

Torn tendons surgically reattached to bone do not tend to regain their original mechanical strength and typically form an inferior scar tissue at the tendon-bone interface^{1, 2}. Tendon retear rates for rotator cuff repair surgeries range from 50-90% depending on the severity of the original tear and are correlated with the functional outcome following repair^{3, 4}. ArthroFlex, a minimally manipulated acellular human dermal allograft, aids in the repair of torn tendons by providing additional strength to the tendon-bone integration site, potentially preventing retear. In an independent study, ArthroFlex was demonstrated to increase the ultimate strength-to-failure of a reattached tendon compared to reattachment without the use of ArthroFlex⁶.

The natural tendon repair process is characterized by the deposition of fibrous tissue at the tendon-bone interface resulting in a tendon-bone attachment site that is weaker than the native insertion site⁴. When grafted on top of a tendon reattachment, ArthroFlex provides structural support to the repair site and a multitude of *human-derived* structural extracellular matrix protein including elastin and many types of collagen. Collagen and elastin provide strength and flexibility that are not properly recapitulated in the natural tendon healing process.

Chronic tendon injury and tendon retear is believed to be a byproduct of the proteases released following the apoptotic and autophagic cell death that occurs during injury². The protease release results in a loop whereby a tendon injury causes cell death, the dead cells release proteases, the proteases weaken the tendon, and the tendon is left more susceptible to reinjury. Using LC-MS/MS, ArthroFlex was found to contain ECM components present in the native dermis ECM including collagens, proteoglycans, and elastin. ECM can modify the wound environment by providing the substrates for the proteases known to weaken the healing tendon and believed to be responsible for chronic tendon injury.

ArthroFlex Provides Human ECM Support with a 10⁻⁶ Sterility Assurance Level

These findings suggest that ArthroFlex retains a broad array of extracellular matrix components, matrikines, growth factors, and cytokines present in healthy human skin and provides structural ECM components that can help prevent retearing of surgically reattached tendons. ArthroFlex is the only bio-implant for augmentation of tendon reattachment composed of natural *human* ECM with greater than 97% of the DNA removed and a minimal risk of infection with a 10⁻⁶ sterility assurance level.

References

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Notes

- † This analysis was performed by Dr. Qishan Lin at the University of Albany proteomics facility.
- ‡ LC-MS/MS is an analysis whereby the fragmented proteins present in a solution are separated by molecular weight. Then, each separated fragment is further broken into smaller components and the molecular weights of those smaller components are determined. From the molecular weights of the smaller components, the amino acid sequence of the original fragment can be resolved.

What is ArthroFlex?

- ArthroFlex is biocompatible decellularized human dermal allograft with an intact acellular framework.
- ArthroFlex provides natural strength and support for injured tendons with *human* collagen.
- ArthroFlex retains native ECM components, matrikines, growth factors, and cytokines, while providing a scaffold for recipient cell proliferation and migration.