Anatomical Preparation Method for Achilles Tendon Allograft in Anterior Cruciate Ligament Reconstruction

Hechtman, K<sup>1</sup>; Sussman, M<sup>2</sup>; Latta, L.L<sup>3,5</sup>; Delcroix, G.J.-R<sup>3,4</sup>; Kaimrajh, D<sup>5</sup>; Milne, E<sup>5</sup> <sup>1</sup>UHZ Sports Med. Inst.; <sup>2</sup>Florida International Univ., Miami, FL; <sup>3</sup>Univ. of Miami,Dept. Orthop., Miami, FL, <sup>4</sup>Research Service & GREC Center, Bruce W. Carter Veterans Affairs Medical Center, Miami, FL, USA <sup>5</sup>Max Biedermann Inst. for Biomech. (MBI), Mount Sinai Hospital, Miami Beach, FL, USA





Introduction: Achilles tendons are commonly used today as replacements for torn anterior cruciate ligaments. Allografts are used in almost half of primary reconstructions, one-fifth of which are Achilles tendons. Achilles tendons are used because of their strong tensile strength and donor availability. However, in the past, this allograft preparation has been plagued by a high failure rate. This study hypothesize





that the high failure rate may be due in large part to the current preparation method of Achilles tendons.

**Objectives**: Using our anatomical approach, thereby cutting the tendons in a parallel fashion with the fiber orientation, allows the tendon to preserve a larger cross sectional area and possibly high tensile strength, making it a good option for a bone-tendon configuration allograft. The present study aims to determine whether the anatomic preparation of Achilles tendons has a higher tensile strength than the traditional central one-third preparation.

**Methods**: Twelve sets of de-identified cadaver Achilles tendons were procured from the UMTB (Vivex Biomedical Inc. Miami, FL). Tendons were prepared as pairs, with right and left from a single donor. Within a tendon pair, the right and left tendons were randomized to a preparation method. This ensured that both preparation methods were represented in every tendon pair. Tendons would then be prepared using either the central 1/3 method (standard of care) or the novel anatomical preparation method. Tendons were tested on the MTS machine. Results were analyzed with one-tailed paired t-test, using SPSS.

Figure 2 Specimens were mounted with a 4 cm working length, left. The grips were frozen and the specimen pulled at 100% strain/second, right.

**Results**: Twelve pairs of Achilles grafts were tested, for a total of 24 tendons. All of the tendons included in the analysis achieved mid-substance rupture when tested on the MTS machine. A paired t-test was used to evaluate the difference between the two preparation method means. It revealed a statistically significant difference between the two groups, with the anatomical preparation being stronger, Central third 2519.7 N ± 873.8 vs. Anatomic 3171.6 ± 751.4 (95%CI 73.2-1230.6 N, t(11)=2.479, p=0.012). The Anatomic preparations were also significantly stiffer than the central third, 381.0 N/mm ± 114.3, vs. anatomic, 463.7 ± 121.2. The two preparations were not significantly different in ultimate strain, Central third 38.7% ± 57.2 vs. Anatomic 41.8% ± 39.4. The study has a power of 0.79.

Figure 4 The stiffness of the central third of the Achilles tendons vs. the anatomically prepared graft.



**Description of 2 preparation methods** 





Figure 5 Ultimate strain of the central third of the Achilles tendons vs. the anatomically prepared graft.

**Conclusions**: The data from our study indicate that the anatomic preparation method of achilles tendons is significantly stronger than the central one-third method. This challenges the continued use of the central one-third method in ACL reconstructions, given their relatively high failure rate; however, our favorable *ex vivo* results do not necessarily translate to improved clinical outcomes *in vivo*.



Figure 1 Identify the major band (middle), and mark a pair of tendons for preparation, one side for central 1/3<sup>rd</sup>, (right), the other anatomic, (left).

Figure 3 Ultimate strength of the central third of the Achilles tendons vs. the anatomically prepared graft. ACKNOWLEDGEMENTS: This project was supported by UHZ Sports Medicine, Coral Gables, FL and MBI for Biomechanics, Mount Sinai Medical Center, Miami Beach, FL and UMTB, Vivex Biomedical, Inc.