



JRF Ortho tendons are aseptically handled throughout the process, never exposed to irradiation.

Some processors market their tissue as processed and packaged aseptically, but up to 65% of their tendons may still be treated with irradiation.¹

An allograft cleanse, soak, and rinse process is used. The formula does not include harsh chemicals – 98% Sterile Water, 2% mixture of isopropyl alcohol and antibiotics. The process does not contain Hydrogen Peroxide, which damages collagen.²

Each tissue has two cultures including a highly sensitive final fluid extraction culture³, which must be negative for any growth.



ASEPTIC TENDONS FROM JRF ORTHO ARE NEVER IRRADIATED, indicated by unique part codes, so that you can be 100% certain of what product you are getting.





FOR THE SURGEON WHO WANTS NON-IRRADIATED TENDON ALLOGRAFTS, JRF ORTHO OFFERS A WIDE VARIETY TO MEET YOUR NEEDS

Bone Tendons

(pre-shaped bone available)

ACHILLES TENDON

PATELLA LIGAMENT

QUADRICEPS TENDON

Soft Tissue Grafts

TIBIALIS TENDON

(anterior and posterior, single and double strand available)

PERONEOUS LONGUS TENDON

SEMITENDINOSUS TENDON

(single & double strand available)

SEMITENDINOSUS/GRACILIS QUAD BUNDLE

GRACILIS TENDON

(single & double strand available)

ENSURINGSafety & Tissue Integrity

ADVANCED CLEANSING TECHNOLOGIES CLEANSE WHILE PRESERVING TISSUE INTEGRITY

Tissue integrity is maintained with consistent processing, monitored temperature and limited reagent exposure.

Dur processors' use a proprietary bioburden reduction step that removes blood and lipids.

MAXIMIZING SAFETY THROUGH DONOR EVALUATION

JRF Ortho and its recovery and processing partners utilize extensive evaluation criteria to identify and qualify suitable donors in accordance with AATB standards including: donor screening, recovery procedures, serological evaluation, microbiological testing and medical records review.

Utilizing highly sensitive NAT testing has resulted in no confirmed incidence of disease transmission.

CONFIRMED ABSENCE OF MICROBES

Tissue is released after no growth culture results from a validated membrane filtration test. Fluid extraction testing is more accurate because contamination is difficult to detect by swabbing the external surface of the graft.⁴

Results clearly demonstrate that the liquid culture method is superior to swab cultures in microbial detection.³

REGULATORY COMPLIANCE

- ✓ FDA
- ✓ AATB
- ✓ CLIA
- ✓ ISO

PICK YOUR SPECIFIC TENDON ONLINE AT JRFORTHO.ORG/ORDER

- "Approximately 65% of all recovered unprocessed tissue is exposed to a low dose (I2 to 18 kGy) of gamma irradiation in a frozen state prior to aseptic processing. The Musculoskeletal Transplant
 Foundation utilizes this low-dose irradiation pretreatment step to decontaminate tissue prior to aseptic processing." Greenberg, D. D., et al. Allograft Compared with Autograft Infection Rates in Primary
 Anterior Cruciate Ligament Reconstruction. The Journal of Bone and Joint Surgery 92.14 (2010): 2402-408.
- 2. Gardner EM, VonderHeide N, Fisher R, Brooker G, Yates PJ.. Effect of hydrogen peroxide on human tendon allograft. Cell Tissue Bank. 2013 Dec;14(4):667-71. doi: 10.1007/s10561-013-9377-x.
- 3. Dennis, Jet al. A comparison of two microbial detection methods used in aseptic processing of musculosketletal allografts tissues. Cell Tissue Bank. 2011:12:45-50
- $4.\ Vehmeyer\ S\ et\ al.\ Bacterial\ contamination\ in\ postmortem\ bone\ donors.\ Acta\ Orthop\ Scand\ 2002; 73(6): 678-683$

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