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Spine

# AlloCraft<sup>®</sup> CA Corticocancellous Cervical Graft

The AlloCraft<sup>™</sup> CA Corticocancellous Cervical Graft is a proprietary assembled, freeze-dried, bone graft with 5° lordosis.



Porous matrix of cancellous bone promotes bone ingrowth

V-Groove pattern helps prevent graft movement and expulsion once implanted

Proprietary corticocancellous assembly provides load-bearing strength and stability and helps promote bone ingrowth

Biomechanically tested to assure strength and stability

- **5**° lordosis supports proper alignment of the cervical spine
- The patented BioCleanse® Tissue Sterilization Process assures sterility to SAL 10<sup>-6</sup>

## **Mechanical Testing**



Extensive biomechanical testing was performed on the AlloCraft<sup>™</sup> CA. Test methods were designed to replicate the amount of force and load applied in the cervical spine.

#### **Impact Testing**

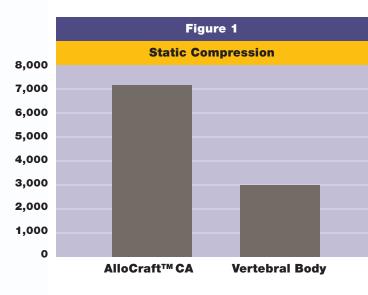
Impact testing was performed to simulate the amount of force applied to insert and tamp the AlloCraft<sup>™</sup> CA into the cervical disc space. Successful testing survived a force of 700lbs exerted for a repetition of 10 cycles.

#### **Static Compression**

Successful static compression was conducted to assure the AlloCraft<sup>™</sup> CA could withstand the natural load applied in the cervical spine (3,028 N of compression). All tested samples surpassed 7,000 N of compression (see Figure 1).

#### **Dynamic Compression**

Dynamic compression testing was performed to measure the fatigue point of AlloCraft<sup>™</sup> CA. To assure strength and stability, successful testing was completed with a minimum force of 250 N applied for five million cycles.



AlloCraft<sup>™</sup> CA Corticocancellous Cervical Graft Specifications

Product Code	Height (±25)	Width (±0.4)	Depth (±0.4)	Lordosis
6183-7-005	5.25mm	14mm	11mm	5°
6183-7-006	6.25mm	14mm	11mm	5°
6183-7-007	7.25mm	14mm	11mm	5°
6183-7-008	8.25mm	14mm	11mm	5°
6183-7-009	9.25mm	14mm	11mm	5°
6183-7-010	10.25mm	14mm	11mm	5°



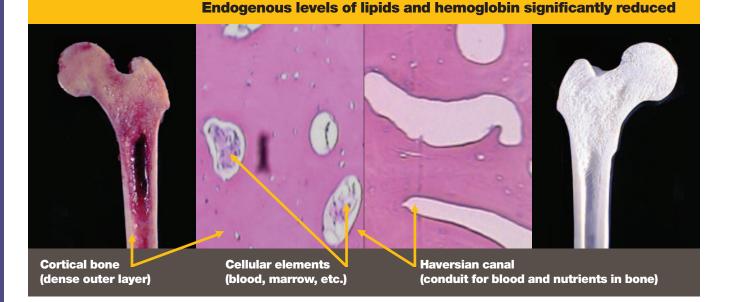
## The BioCleanse®

Difference

The BioCleanse<sup>®</sup> Tissue Sterilization Process is a proprietary, low temperature, pharmaceutical-grade chemical sterilization process for musculoskeletal bone and soft tissue implants.

- Penetrates inner matrices of the bone. Removes cellular elements such as blood, lipids and marrow.
- Validated to eliminate resistant organisms viruses, fungi, bacteria and spores (HIV, hepatitis, E.coli, Clostridium, etc.).
- Preserves biomechanical integrity and biocompatibility.

### **Removal of Blood Elements**



The BioCleanse<sup>®</sup> Tissue Sterilization Process is a validated technology that sterilizes tissue, has been scientifically and clinically proven to eliminate donor-to-recipient disease transmission risk, and preserves tissue strength and biocompatibility.





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Stryker Spine 2 Pearl Court Allendale, NJ 07401 t: 201-760-8000 www.stryker.com

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to their own clinical judgment when deciding which treatments and procedures to use with patients. Products may not be

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