

**Certificate of Conformance (COC) –  
Surgalign FibreX Demineralized Bone Matrix**



## FORM

Document No.	FORM-0305
Parent SOP	SOP-0075
Revision	00
Effective Date	09 MAR 2021

**DONOR DESCRIPTION**

Donor Number OB21-0013  
 Age 53  
 Gender MALE

**CONSENT**

YES NO NOT DETERMINED

Research/Education  YES  NO  NOT DETERMINED  
 International  YES  NO  NOT DETERMINED

**ELIGIBILITY DETERMINATION**

YES NO NOT DETERMINED

EU criteria per EU Directive 2006/17/EC including HTLV I/II  YES  NO  NOT DETERMINED

**PRODUCT DESCRIPTION**

*Freeze dried, aseptically processed bone (demineralized bone fibers) cleansed with a validated process that utilizes Gentamicin. Final product is terminally sterilized via validated irradiation process. Storage is 15°C to 30°C.*

Product ID Number(s) OB21-0013 Lot No. Range -096 thru -116  
 Customer Part Number FBX-0010  
 Expiration Date 21 MAY 2023

**PRODUCT ANALYSIS**

**MICROBIAL TESTING** Tissue is subjected to microbiological testing at recovery and processed and sterilized under validated conditions to be considerably free of specific aerobic / anaerobic microorganisms and fungal contaminants whose presence would preclude tissue from processing or transplantation.

The above listed product complies with the requirements as set forth by the American Association of Tissue Banks (AATB) and the Food and Drug Administration (FDA) for human tissue for transplantation.

**TEST****RESULT**

Sterility Dose Report PASS  
 Residual Calcium PASS  
 Donor Osteoinductivity Potential PASS  
 Residual Moisture PASS

**DONOR ELIGIBILITY AND SEROLOGICAL TESTING**

Donor eligibility (screening and testing) is performed in accordance with AATB standards and FDA regulations (21 CFR 1271). Donor screening includes assessment of the medical and social history as well as physician assessment of the donor to assure that no conditions exist that may make the tissue unacceptable for transplantation. Donor eligibility is determined by a Board-Certified Physician (Medical Director).

Communicable disease testing was performed by a laboratory registered with FDA to perform donor testing and certified to perform such testing on human specimens in accordance with the Clinical Laboratory Improvement Amendments of 1988 (CLIA) and 42 CFR Part 493. The testing was conducted using FDA licensed, approved, or cleared donor screening tests for cadaveric specimens where applicable. The records of this testing are maintained at Origin Biologics, 6635 S. Eastern Avenue, Suite 100, Las Vegas, NV 89119. The following required testing was performed and found to be negative or non-reactive:

Antibody to Human Immunodeficiency Virus 1 & 2 (HIV 1 & 2)  
 Human Immunodeficiency Virus Type 1 (HIV-1 NAT)  
 Hepatitis B Core IgG/IgM Antibody (HBcAb)  
 Rapid Plasma Reagin or Serologic Test for Syphilis (RPR or STS)

Antibody to Hepatitis C (HCV)  
 Hepatitis C Virus (HCV-NAT)  
 Hepatitis B Surface Antigen (HBsAg)

Additional tests including, but not limited to, Human T-Cell Lymphotropic Virus Type I & II (HTLV I & II) and HBV NAT may have been performed at the time of donor screening and were found to be acceptable for transplantation.

Quality Representative:

Michelle Haider

Date:

29 July 2021