

# BIOLAB Membrane Wrap - Hydro™

## amniotic allograft membrane

### Instructions for Use

#### PRODUCT DESCRIPTION

**BioLab Membrane Wrap - Hydro™** is a human tissue allograft derived from the amniotic membrane that provides structural tissue for use as a wound and protectant covering.

**Membrane Wrap - Hydro™** is an Amniotic fluid-hydrated human tissue allograft [Human Cellular and Tissue Based Product (HCT/P)] for transplantation regulated by the US Food and Drug Administration under 21 CFR Part 1271.

#### PACKAGE CONTENTS

The product package contains the following items:

- One tissue graft, double packaged in sealed pouches
- Instructions for Use insert (this document)
- One set of supplemental Tracking Labels
- One allograft Tissue Tracking Record (TTR) card

If any of these items are missing, please contact BioLab Sciences.

#### PREPARATION & APPLICATION

**Step 1:** Open product box and remove the product pouch.

**Step 2:** Using aseptic technique, peel open the outerpouch and place the inner pouch into the sterile field.

**Step 3:** When ready to use, either tear open the inner pouch at the notch or cut the pouch at the notch to expose the graft.

**Step 4:** Remove the top layer of mesh and apply the membrane onto the wound using the remaining mesh.

**Step 5:** Use the forceps to grasp the membrane and then peel the top mesh from the membrane, leaving the membrane in place

**Step 6:** Pour the remaining fluid onto the graft which is now covering the wound

**Step 7:** The excess membrane can be folded into the wound. Then the covering dressing with a non-adherent layer is placed, followed by the outer dressing as pre the clinician.

**Step 8:** Graft may be cut with scissors to apply over multiple sites.

**Step 9:** If desired, graft may be hydrated prior to application with sterile saline for tight or hard to reach areas.

**Step 10:** Use forceps to apply the graft over the intended site. Achieve full contact and remove the packaging mesh.

**Step 11:** Ensure the HCT/P is secured in place by the Physician's choice of fixation.

#### STORAGE AND HANDLING

- Store product at ambient temperature (15-25°C, 59-77°F).
- Handle using aseptic techniques.

#### WARNINGS

- For single patient use only.
- To be used under the supervision of a qualified healthcare provider.
- Do not use if the product sterile barrier system or its packaging is compromised.
- Cannot be re-sterilized.

#### PRECAUTIONS

- The graft should not be applied in the presence of a live infection.
- BioLab Sciences makes no claims concerning the biological properties of this allograft tissue. All tissues have been collected, processed, stored, and distributed in compliance with US Food and Drug Administration (FDA) regulations governing human cells, tissues, and cellular and tissuebased products (HCT/Ps) to prevent the transmission of communicable diseases listed on page two. Current technologies may not preclude the transmission of communicable diseases.

## HCTP RECORD TRACKING

Recipient records must be maintained for the purpose of tracking tissue post-transplant in accordance with The Joint Commission standards and the FDA requirements under 21 CFR Part 1271. Supplemental labels, which indicate the tissue ID number, are contained in this package for tracking processes. The allograft ID number must be recorded in the operative record. The provided Tissue Tracking Record (TTR) must be completed and returned to BioLab Sciences.

## PROCESSING

The HCT/Ps are processed in accordance with FDA's Good Tissue Practice regulations in a controlled cleanroom environment, using processes designed to prevent contamination of the tissue, and to prevent the introduction, transmission, or spread of communicable diseases.

The tissue products are sterilized using electron-beam irradiation for a Sterility Assurance Level of SAL<sup>10<sup>-6</sup></sup>.

## DONOR SCREENING, TESTING AND ELIGIBILITY

The donated human birth tissue has been determined to be eligible for transplantation by a licensed physician, the Medical Director of BioLab Sciences. In accordance with FDA regulations under 21 CFR Part 1271, the donor has been deemed to be free from risk factors for, and clinical evidence of, infection due to relevant communicable diseases and other exclusionary disease conditions through review of donor records, including a medical/behavior risk assessment, medical records, and a recent physical examination. Additionally, testing of a qualified blood sample indicates that the donor is nonreactive or negative for the following communicable disease markers:

- Antibody to human immunodeficiency virus (HIV) types 1 & 2
- Hepatitis B surface antigen (HBsAg)
- Hepatitis B core (HBc total)
- Antibody to hepatitis C (HCV)
- Syphilis (RPR)\*
- WNV NAT
- HCV NAT
- HIV NAT
- HBV NAT

\* Tissues from a donor whose blood specimen is initially reactive for the non-treponemal screening assay, are cleared for transplantation use **only** when the confirmatory result from the treponemal specific assay is non-reactive.

All laboratories performing donor screening tests for this product are registered with the FDA and certified to perform testing on human specimens under the Clinical Laboratory Improvement Amendments of 1988 (CLIA) and 42 CFR part 493, or equivalent requirements. Test kits used are approved and cleared by the FDA for screening blood specimens collected from living donors. A copy of the relevant medical records can be obtained from BioLab Sciences upon request.

## ADVERSE REACTIONS

No adverse clinical reactions to this tissue product have been reported. Adverse reactions or outcomes that potentially involve the use of this tissue product must be reported immediately to BioLab Sciences at [QA@biolabsciences.net](mailto:QA@biolabsciences.net) or by calling 480-656-5746, ext. 200.

## ALLOGRAFT TISSUE PROCESSED BY:

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FDA FEI (FDA Establishment Identifier)#: 3014573577

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