

PROKERA®

U.S. Patent No. 7,494,802 B2

Product Insert

Description:

PROKERA® is a self-retaining biologic corneal bandage. It is comprised of a cryopreserved amniotic membrane graft fastened to an ophthalmic conformer. This product is aseptically processed from tissues obtained from donated human tissue (placentas) according to current Good Tissue Practices (cGTP) and Good Manufacturing Practices (cGMP) regulations established by the US Food & Drug Administration (FDA). PROKERA is a Class II medical device cleared by the FDA.

Donor Eligibility and Summary of Records:

- This tissue was procured from a donor determined to be eligible based on the results of screening and testing. Human Cells, Tissues, and Cellular and Tissue-based Products (HCT/P) donor eligibility and placenta suitability, which is based on the results of donor screening at delivery for infectious, malignant, neurological & auto-immune diseases and for other exposures or social habits, has been determined and documented by TissueTech, Inc.
- A blood specimen, drawn within ± 7 days of donation, has undergone serological testing by a laboratory registered with the 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, or that has met equivalent requirements as determined by the Centers for Medicare and Medicaid Services (CMS). The donor has been tested for the following infectious diseases and results were non-reactive:
 - HIV-1 & HIV-2 Antibody
 - HIV-1 (RNA-NAT)
 - Hepatitis B Surface Antigen (HBsAg)
 - Hepatitis B Core Antibody (HBcAb)
 - Hepatitis B Virus (HBV, DNA-NAT)
 - Hepatitis C Antibody (HCVAb)
 - Hepatitis C Virus (HCV, RNA-NAT)
 - Syphilis (RPR)
 - HTLV I & II Antibody (HTLV I/II Ab)
 - West Nile Virus (WNV, RNA-NAT)
- Microbial testing has also been performed on the final product to ensure no growth of aerobic, anaerobic or fungal cultures.
- The cell activity in the tissue has been inactivated using our CryoTek® cryopreservation process to reduce the possibility of graft rejection, while retaining the natural biologic properties.
- PROKERA is stored in a medium of Dulbecco's Modified Eagle Medium/Glycerol (1:1) containing 20 µg/ml Ciprofloxacin and 1.25 µg/ml Amphotericin B.
- This tissue has been deemed eligible for transplantation based on acceptable screening, serological and microbial test results.
- A Certificate of Compliance regarding the manufacturing, storage, shipping and tracking controls for Bio-Tissue, Inc. products is available upon request.

Indications:

- PROKERA is intended for use in eyes in which ocular surface cells are damaged or underlying stroma is inflamed or scarred and best suited to prevent adhesion of the eyelid to the ocular surface with the large ophthalmic conformer. Acting as a self-retaining biologic corneal bandage, PROKERA effectively treats superficial corneal surface diseases by suppressing inflammation and related pain, promoting epithelial healing, and avoiding haze.
- PROKERA is inserted between the eyeball and the eyelid to maintain space in the orbital cavity and to prevent closure or adhesions. Placement of the conformer also enables application of the cryopreserved amniotic membrane to the ocular surface without the need for sutures.
- PROKERA is for single-use only in one patient by an ophthalmologist or optometrist.

Contraindications:

- PROKERA should not be used in eyes with glaucoma drainage devices or filtering bleb.

Precautions:

- Do not use PROKERA if the device or packaging is damaged – contact Bio-Tissue immediately if there is any abnormality observed (e.g. device, labeling, shipping, missing information, etc.). Refer to the Customer Feedback section for reporting.
- PROKERA exposed to controlled room temperature (20°C - 25°C) for up to 6 hours may be returned to cold temperature storage in accordance with the Storage section as long as the packaging remains unopened and intact.
- Once the packaging is opened, PROKERA shall either be transplanted or otherwise discarded.
- Do not sterilize or autoclave the product before use.
- Handle PROKERA with care during insertion.

Warnings:

- Do not use on patients with a history of drug reactions to Ciprofloxacin, Amphotericin B, Glycerol, and/or DMEM.
- As with the use of any human tissue, the possibility of infectious agent transmission cannot be completely eliminated although all screening and microbial testing results for this donor were satisfactory for transplantation.
- It is imperative that the device is stored properly until use. For proper storage instructions, refer to the Storage section.
- If the patient cannot close the eyelid, has an incomplete blink or demonstrates any other signs of exposure, address these clinical issues before or at the same time of placing PROKERA.
- If the patient cannot tolerate wearing PROKERA, the device should be removed.

Storage:

It is the responsibility of the Tissue Dispensing Service, Tissue Distribution Intermediary, and or End-User Clinician to maintain tissue intended for transplant in appropriate storage conditions prior to further distribution or transplantation.

Upon receipt, ensure the validated time on the shipper has not expired.
Remove the product and store accordingly until use.

Location & Temperature	Use After Receipt
<p>-80°C → 4°C (-112°F → 39.2°F) Example: ultra-low temperature freezer, standard freezer, or standard refrigerator</p>	<p>Within the expiration date printed on product packaging (shelf-life is 2 years from date of manufacture)</p>

Insertion and Removal Instructions:

- If frozen, allow PROKERA to sit at controlled room temperature (20°C - 25°C) in its original unopened packaging for at least 5 minutes.
- Peel open the lid and handle the tray using aseptic techniques.
- Discard the storage media contained within the tray.
- Thoroughly rinse PROKERA with sterile saline solution while inside the tray to remove the storage media and reduce the potential stinging sensation.
- Remove the center retainer piece from the tray to access the PROKERA.
- Patient may experience a temporary foreign body sensation upon insertion.
- Advise patient to minimize contact with eye once PROKERA is placed on the ocular surface.

Insertion



1. Apply topical anesthesia



2. Hold the upper eyelid



3. Ask the patient to look down



4. Insert the PROKERA into the superior fornix



5. Pull lower eyelid down and slide PROKERA under the lower eyelid



6. Check centration under the slit lamp

7. Apply a tape-tarsorrhaphy over the lid crease (as needed)
- Apply appropriate medications (as needed)



Removal



1. Apply topical anesthesia



2. Pull the lower eyelid down



3. Lift lower edge of PROKERA using a cotton swab or blunt forceps



4. Ask the patient to look down



5. Apply gentle pressure on the upper eyelid



6. Slide the PROKERA out

- Upon dissolution of membrane or completion of treatment, remove PROKERA.
- Do not leave in the eye longer than 30 days.

Recipient Records:

It is the responsibility of the Tissue Dispensing Service, Tissue Distribution Intermediary, and/or End-User Clinician to maintain records for the purpose of tracing tissue post-transplantation. The responsible entity should document the disposition (transplanted or discarded) on the Donor and Recipient Information card (DRI), attach one of the provided PROKERA tracking labels to the DRI and mail to Bio-Tissue, Inc. Attach the remaining labels in patient and hospital records.

Customer Feedback:

Within the United States: Report any customer feedback, including complaint, error or accident notification promptly to Bio-Tissue at (888) 296-8858 or (305) 412-4430.

Outside of the United States: Report feedback to your local tissue provider.

Adverse Events:

The FDA requires that information be supplied to the product manufacturer for mandatory reporting of adverse events. Possible significant adverse events include microbial infection and transmission of viral disease. The doctor is responsible for immediately reporting any adverse event potentially attributable to the use of PROKERA to Bio-Tissue.

For Adverse Events, complete the following section. Notify via:

Phone: (888) 296-8858 or (305) 412-4430

Fax: (305) 675-3262

Email: Customerfeedback@biotissue.com

Serial Number: _____

Expiration Date: ____/____/____

Doctor Name: _____

Facility Name: _____

Transplant Date: ____/____/____

Diagnosis/Procedure: _____

Site of Use: _____

Point of Contact's Name: _____

Point of Contact's Phone Number: _____

Date Adverse Event was Reported: ____/____/____

Type of Adverse Event: Microbial Infection Transmission of Viral Disease Other

Describe the Adverse Event: _____

Puede conseguir la versión en español de este prospecto del producto en www.biotissue.com.

**Manufactured for Bio-Tissue, Inc. by
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