

Ethnicity    White    Black    Hispanic    Asian    Other    Age    Sex    M    F

**CONDITION BEING TREATED WITH AMNIOTIC MEMBRANE THERAPY**

- |   |   |
|---|---|
| <input type="checkbox"/> Anterior basement membrane dystrophy | <input type="checkbox"/> Disruption of surgical wound   |
| <input type="checkbox"/> Band-shaped keratopathy              | <input type="checkbox"/> Dry eye syndrome               |
| <input type="checkbox"/> Blepharitis                          | <input type="checkbox"/> Entropion                      |
| <input type="checkbox"/> Chemical burn                        | <input type="checkbox"/> Herpes zoster / herpes simplex |
| <input type="checkbox"/> Thermal burn                         | <input type="checkbox"/> Hyperlacrimation               |
| <input type="checkbox"/> Chronic conjunctivitis               | <input type="checkbox"/> Keratoconjunctivitis           |
| <input type="checkbox"/> Conjunctivochalasis                  | <input type="checkbox"/> Keratoconus                    |
| <input type="checkbox"/> Corneal abrasion                     | <input type="checkbox"/> Peripheral degeneration        |
| <input type="checkbox"/> Corneal edema                        | <input type="checkbox"/> Pterygium                      |
| <input type="checkbox"/> Corneal ulcer                        | <input type="checkbox"/> Recurrent corneal erosion      |
|   | <input type="checkbox"/> Stevens-Johnson Syndrome       |

ICD-10 Code \_\_\_\_\_

**ALLOGRAFT ID NUMBER**

Right Eye

STICKER

Left Eye

STICKER

Date of placement procedure \_\_\_\_\_

Date of allograft response \_\_\_\_\_

**Surgical events, including but not limited to:**

- Surgery performed on a wrong body part inconsistent with the documented informed consent
- Surgery performed on the wrong patient
- Wrong surgical procedure performed on a patient
- Retention of foreign object in a patient after surgery or other procedure
- Discontinuation of therapy session prior to demonstrable improvement

**Product or device events, including but not limited to:**

- A patient death or serious disability associated with the use of a contaminated drug, device, or biologic provided by the office
- A patient death or serious disability associated with the use or function of a device in patient care in which the device is used or functions other than as intended

**Care management events, including but not limited to:**

- A Type II allergic reaction that involves the interaction of immunoglobulins with foreign substances or autoantigens closely associated with cell membranes and/or proteins, no loss of vision or serious visual disability after treatment
- A Type IV hypersensitivity reaction mediated by the adaptive immune system, no loss of vision or serious visual disability after treatment
- A loss of vision or serious visual disability associated with a cytotoxic hypersensitivity reaction and subsequent stromal cell lysis
- A loss of vision or serious disability by infectious means secondary to overnight wear of the amniotic membrane allograft and/or the retaining contact lens

Doctor's Signature \_\_\_\_\_

**POSTOPERATIVE MEDS**

- None
- Artificial tears
- Unpreserved artificial tears
- Antibiotic eyedrops
- Steroid eyedrops
- Steroid/antibiotic combination
- NSAID eyedrops

**RISK FACTORS FOR ALLOGRAFT RESPONSE**

- History of previous rejections
- Erosion of corneal privilege in diseased corneas in which inflammation has occurred
- Enhanced sensitization in diseased corneas secondary to chronic inflammation from ongoing ocular surface disease
- Diabetes and other inflammatory diseases
- Healing from previous surgeries or trauma
- Deep neovascularization of the recipient cornea
- Severe atopic dermatitis
- Limbal stem cell deficiency