

Title: Exploring Amniotic Membrane in the Medical Managed Practice

CE Hours: 1

Category: Treatment of Anterior Segment

Description: This one-hour course is focused on amniotic membranes as a therapeutic treatment option for various corneal diseases. We will discuss the history, types, indications, clinical application, and proper billing and coding of amniotic membranes.

Objectives:

- Discuss clinical applications of amniotic membranes in eye care
- Describe the history and indications of amniotic membranes in the optometric office
- Discuss appropriate billing and coding of amniotic membranes

### **Exploring Amniotic Membrane in the Medical Managed Practice**

**Jessilin Quint, OD, MBA, MS, FAAO**

**Steven Potwin, OD, FAAO**

1. What is the amnion?
  - a. Innermost, avascular lining of the fetal membrane & placenta
  - b. Shares the same cell origin as the fetus
  - c. Contains anti-inflammatory cytokines and growth factors
  - d. Harvested in a sterile environment from placental tissue obtained during elective cesarean sections
  - e. Donors are screened for transmissible disease
  - f. Benefits<sup>1</sup>
    - i. Promotes epithelialization
    - ii. Suppresses inflammation
    - iii. Inhibits scarring
    - iv. Inhibits angiogenesis
    - v. Anti-microbial agent
2. History & Evolution
  - a. 1910=1<sup>st</sup> documented use as skin graft <sup>2</sup>
  - b. 1940=1<sup>st</sup> documented use in eye (chemical burn) <sup>3</sup>
  - c. Modern uses in healthcare
3. Amniotic Membrane Components
  - a. Key components:
    - i. Heavy chain hyaluronic acid
    - ii. Proteoglycans
    - iii. Growth factors

- iv. Collagens
    - v. Fibronectin
    - vi. laminin
  - b. Key Functions:
    - i. Promotes epithelial growth through cell migration, adhesion, & differentiation<sup>4,5</sup>
    - ii. Inhibits fibroblast growth (reduces scarring)
    - iii. Inhibits pro-inflammatory cells
      - 1. decreased expression of cytokines
      - 2. Suppresses t-cell activation
      - 3. Inhibits giant cell formation
      - 4. Controls MMP production
      - 5. Blocks angiogenesis
    - iv. Universally tolerated due to lack of histocompatibility antigens (HLA-A, B, or DR)
4. Options
- a. Cryopreserved
    - i. Pros: no lost nutrients
    - ii. Cons: patient comfort, cost
  - b. Dehydrated
    - i. Pros: patient comfort, variety of sizes, reduced cost/greater ROI, easy to stock/store
    - ii. Requires BCL for retention
5. Ocular Indications
- a. Band Keratopathy
  - b. Bullous Keratopathy
  - c. Corneal & conjunctival burns and corrosions
  - d. Filamentary keratitis
  - e. Corneal ulcers
  - f. Corneal abrasions
  - g. Keratopathy defects
  - h. Post-infection keratitis
  - i. Mooren's ulcer
  - j. Neurotrophic keratoconjunctivitis
  - k. Recurrent corneal erosion
  - l. Stevens-Johnson syndrome
6. Ease of Use
- a. Video demonstrating placement of both cryopreserved and dehydrated amniotic membranes on eye
  - b. Clinical "Pearls" on placement of amnio membrane
  - c. Common Pitfalls
    - i. Eye rubbing, slipped membranes, second placement
7. Case Studies
8. Billing & Coding

- a. CPT 65778: Placement of amniotic membrane on ocular surface without sutures
  - b. Bilateral 150% rule applies
  - c. Do not bill CPT 92071 (fitting of contact lens for treatment of ocular surface disease) concurrently
  - d. Do not bill CPT 9921x (E/M office visit) concurrently
  - e. Global period is ZERO days, bill any follow-ups
  - f. No frequency of limitations
9. Patient Benefits & Outcomes
10. Practice Benefits
- a. Advance patient care
  - b. Reimbursement
  - c. Expansion of the types of patients retained or brought into the practice
  - d. Practice viewed as a progressive, medically oriented optometric care clinic

References:

- 1) Dua HS, Gomes JA, King AJ, Maharajan VS. The amniotic membrane in ophthalmology. *Surv Ophthalmol.* 2004; 49(1): 51-77.
- 2) Davis JW. Skin transplantation with a review of 550 cases at the JohnsHopkins Hospital. *Johns Hopkins Med J Hosp Rep* 1910;15:307-96.
- 3) De Rotth A. Plastic repair of conjunctival defects with fetal membranes. *Arch Ophthalmol* 1940;23:522-5.
- 4) Meller D, Tseng SC. *Invest Ophthalmol Vis Sci.* 1999;40(5):878-886.
- 5) Boudreau N et al. *Proc Natl Acad Sci USA.* 1996;93(8):3509-3513.