Amniotic Membrane Grafts Following Excision of Corneal and Conjunctival Intraepithelial Neoplasia

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Background: We evaluated the efficiency of amniotic membrane grafts (AMGs) for reconstructing the conjunctival surface following excision of corneal and conjunctival intraepithelial neoplasia (CIN).

Methods: This was a retrospective, noncomparative, interventional study. Five eyes in 5 patients were treated between April 1996 and September 2002 with the same procedure to apply amniotic membrane grafts after excising CIN. According to a standard protocol, the amniotic membrane was harvested and processed under sterile conditions. The amniotic membrane graft was applied over the excised bare scleral area and anchored with 10-0 nylon interrupted sutures. Patient data and clinical photographs were reviewed and analyzed.

Results: The mean follow-up period was 27 (range, 6-69) months. Ocular surface healing was rapid and complete in all cases. No recurrence was found.

Conclusion: Amniotic membrane graft is an effective and safe alternative adjunctive treatment for primary CIN. Further studies with longer follow-up are recommended to evaluate the risk of recurrence and other adverse effects.

Key words: corneal and conjunctival intraepithelial neoplasia, amniotic membrane graft, recurrence.

Corneal and conjunctival intraepithelial neoplasias (CINs) are uncommon tumors of the ocular surface. Traditionally, these lesions were regularly treated with complete surgical excision with or without adjunctive therapy in an effort to reduce the rate of recurrence. However, the recurrent rate may reach 50% of cases when the pathology of the excised margin is not confirmed. Many adjunctive treatments such as cryotherapy, beta irradiation, Mohs' technique, mitomycin-C, 5-fluorouracil, and interferon have been introduced to try to reduce the recurrence rate. However, potential complications resulting from adjunctive therapies cannot be ignored. In this report, we present patients with pathologically proven CIN treated with amniotic membrane grafts following primary surgical excision of their lesions. Our results show satisfactory outcomes for conjunctival surface reconstruction by amniotic membrane grafts.

METHODS

Patients

Five patients were included in this study between April 1996 and September 2002. Informed consent from all patients was obtained prior to the
study. One eye in each patient was afflicted with CIN. All patients underwent the same procedure of lesion excision followed by an amniotic membrane (AM) graft. The operations were performed by two of the authors (DHKM and HCL) in Chang Gung Memorial Hospital, Linkou, Taiwan, using standard surgical procedures.

Tumor excision and amniotic membrane grafting

Under retrobulbar anesthesia, lesions were removed using the technique of a superficial keratectomy with a 3-4 mm tumor-free margin at the corneal and conjunctival side. The AM was prepared and preserved as described previously. After 1999, to meet new US FDA requirements, AM donors were examined for human immunodeficiency virus (HIV) 6 months after delivery; the AMs were purchased from Bio-Tissue (Miami, FL, USA). The AM was sutured to the adjacent conjunctiva and episcleral tissue using interrupted 10-0 nylon sutures with the basement membrane side facing up. Following AM grafting, Maxitrol ointment (Alcon), 2 to 4 times a day, and 25 mg oral indomethacin and antacid, 4 times a day, were administered during the first week. During the second week, only Maxitrol was administered, but if there was excessive irritation due to the stitches, lubricants such as Tear Naturale or Balanced Salt Solution (Alcon) were also administered. Essentially all the stitches were removed at the end of 2 weeks. Thereafter, the eye drops were changed to 0.1% fluorometholone (CIBA Vision, Hettlingen, Switzerland) q.i.d. for 1 month, then decreased to b.i.d. for another month, and then changed to 0.02% fluorometholone (Santen, Osaka, Japan) q.i.d. to b.i.d. for the following 3 months.

Pathology

All excised tumors were sent for a pathological examination at the Department of Pathology, Chang Gung Memorial Hospital. All tumors were confirmed to be CIN.

RESULTS

We studied 5 patients with pathologically proven primary CIN. A representative lesion is shown in Fig. 1A,B (case 1). Patients were treated with an AM graft following excision of their tumors. During a follow-up period of 6 to 69 (mean, 27 ± 24.87) months, no clinical evidence of recurrence was observed. In 2 cases, short-term hyperemia and redness occurred within the first 2 weeks after surgery. Clinical data obtained from the 5 patients are listed in Table 1. Histopathological results showed that cases 1 and 2 had carcinoma in situ (CIN, grade III) while cases 3, 4, and 5 had mild dysplasia (CIN, grade I).

Case 1

A 76-year old man was referred to our clinic with a hyperemic mass on his right eye, which he had had for months. His medical history was unremarkable. Uncorrected visual acuity was 20/100 in his right eye and 20/800 in his left eye. Biomicroscopy of his right eye showed a papilloma-like lesion with a neovascular ingrowth. The corneal lesion was 10×4 mm in size and extended from the 4- to the 8-o’clock position (Fig. 2A). He received superficial keratectomy, cryotherapy, and then AM grafting. Postoperative complications were minimal except for red eye and hyperemia within the first 2 weeks following surgery (Fig. 2B). No recurrence occurred during 69 months of follow-up.

Case 2

A 71-year-old man, with no special medical conditions, had experienced blurred vision and foreign body sensation in his right eye for 10 months. Examination showed that his visual acuity was 20/70 in his right eye and 20/50 in his left eye. Biomicroscopy showed a leukoplakic-like lesion which extended from the 7- to the 2-o’clock position (Fig. 2A). He received superficial keratectomy, cryotherapy, and then AM grafting. Postoperative complications were minimal except for red eye and hyperemia within the first 2 weeks following surgery (Fig. 2B). No recurrence occurred during 69 months of follow-up.

Case 3

A 71-year-old woman with a history of red left eye with bloody discharge (OS) for 2 weeks was referred to our institute from a local clinic for further treatment. The corrected visual acuity in her affected eye had deteriorated to 20/100. Slit lamp biomicroscopy showed a leukoplakic pannus-like lesion.
with engorged vessels spanning the area from the 8- to the 4-o’clock position with a 4-mm-wide extension into the corneal side (Fig. 4A). She received superficial keratectomy, cryotherapy, and AM grafting. Complete corneal and conjunctival reepithelialization over the AM was observed in 2 weeks. There was no recurrence after 2 years of follow-up (Fig. 4B).

**Case 4**

A 74-year-old man had redness and irritation in his right eye for 1 week. His medical history was unremarkable. Uncorrected visual acuity was 20/50 in his right eye and 20/70 in his left eye. Slit lamp examination showed an elevated nodular lesion over the temporal side of his right cornea, extending 2 mm from the limbus. After lesion excision followed by AM grafting, no recurrence was observed during a follow-up period of 24.5 months.

**Case 5**

A 31-year-old young woman had suffered from periodic red eye and foreign body sensation (OD) in her eye for 1 year. Slit lamp biomicroscopy showed a papilloma-like lesion with engorged vessels spanning the area from the 7- to the 9 o’clock position, with a 2-mm extension into the corneal side. After excision of this tumor, the area was soaked with mitomycin-C (0.02%) for 1 min followed by AM grafting to cover the bare scleral area. No recurrence occurred after 6 months of follow-up, but occasional redness of the eye occurred within the first 2 weeks following surgery.
Fig. 2 (A) Pretreatment view of the left eye showing a papilloma-like lesion with neovascular ingrowth. (B) Mild hyperemia and congestion within 2 weeks after the operation.

Fig. 3 (A) Case 2 presenting a leukoplakic-like lesion with extension from the 7- to the 2-o'clock position. (B) The corneal and conjunctival lesion measuring about 10×6 mm with a 2-mm extension onto the corneal side.

Fig. 4 (A) Case 3 showing a leukoplakic pannus-like lesion with engorged vessels involving the area from the 8- to the 4-o'clock position with extension onto the corneal side with a 4-mm width. (B) Complete corneal and conjunctival reepithelialization over the amniotic membrane which was completed in 2 weeks. The cosmetic appearance was quite good.
DISCUSSION

CIN, a precancerous lesion of squamous cell carcinoma on the ocular surface, is slowly progressive with a low malignant potential. CIN usually involves the limbal area. It is characterized by nodular, gelatinous, papilliform, flat and superficial or elevated leukoplakic lesions in a diffuse invasive fashion combined with tufts of engorged blood vessels. Traditionally, the treatment modality for CIN involved surgical excision with or without adjunctive treatment such as cryotherapy, radiation, topical mitomycin, 5-FU, interferon (by a topical or intralesional injection), and phototherapeutic keratectomy by excimer laser, which have been used in an effort to reduce the recurrence rate. In spite of this, the recurrence rate of CIN and conjunctival squamous cell carcinoma ranges from 15% to 52%. In addition, complications such as corneal edema, fibrosis, iris atrophy, and intraocular inflammation have been reported to be associated with cryotherapy. Side effects of radiation therapy include scleral necrosis, cataracts, dry painful eye, and even visual loss as a result of radiation vasculopathy. Mitomycin-C has been introduced to treat CIN. However, serious complications such as scleral melting, infection, cataract formation, and limbal deficiency are possible with this treatment. Acute transient toxic keratoconjunctivitis was observed in 5-FU-treated cases. Side effects of interferon treatment include scleral necrosis, cataracts, dry painful eye, and even visual loss as a result of radiation vasculopathy. Mitomycin-C has been introduced to treat CIN. However, serious complications such as scleral melting, infection, cataract formation, and limbal deficiency are possible with this treatment. Acute transient toxic keratoconjunctivitis was observed in 5-FU-treated cases. Side effects of interferon treatment include scleral necrosis, cataracts, dry painful eye, and even visual loss as a result of radiation vasculopathy.

Cryopreserved human AM contains a thick natural basement membrane and an avascular stroma, which may provide an optimal microenvironment to allow epithelial cell proliferation and differentiation. AM has recently been proposed as a new substrate for ocular surface reconstruction surgeries such as in pterygium, symblepharon, conjunctivochalasis, scarring, neoplasia, advanced chemical burns, Stevens-Johnson syndrome, ocular cicatricial pemphigoid, persistent epithelial defects, and corneal ulcers. AM grafting promotes epithelialization and restores ocular surface integrity without inflammation or scarring. The AM stromal matrix suppresses signal activity of transforming growth factor β in human corneal and conjunctival fibroblasts and can also induce apoptosis of leucocytes as shown in an animal study. The AM contains protease indicators indicating that it can reduce inflammation-associated proteolytic activity after application.

In order to prevent subsequent conjunctival ingrowth and corneal vascularization following extensive tumor excision, both autograft and allograft limbal transplantations have been proposed. An AM graft alone has been demonstrated for reconstruction of the ocular surface in eyes with partial limbal stem cell deficiency. Moreover, the advantages of AM grafts over traditional buccal mucosal autografts are the excellent cosmetic outcome and a normal conjunctival epithelial phenotype as demonstrated by impression cytology.

In this study, all cases had good results with no special complications. Only limited short-term complications such as redness and hyperemia occurred. Complete epithelialization occurred within 2 weeks. After a mean following-up period of 27 months, no recurrence was noted, and the cosmetic outcome was good. This study demonstrates the usefulness of AM grafts for the treatment of corneal and conjunctival CIN. The application of AM grafts is considered safe as long as the AM is prepared according to the standard protocol. The transparency of the AM allows for a good cosmetic appearance and ease of monitoring tumor recurrence.

Nevertheless the immunogenic characteristics of AM remain unclear. Gabler and Messmer showed the presence of hypopyon after AM transplantation. In the present study, no one was found to have hypopyon after AM grafts. Also, severe com-
lications such as bacterial contamination of the AM were not observed in this study.

In conclusion, an AM graft is an effective adjunctive therapy for reducing the recurrence of CIN following primary excision of tumors. Larger population studies with longer follow-up periods are recommended to further assessing the risk of recurrence and other possible side effects.

Acknowledgements

The authors thank Professor Ray Jui-Fang Tsai for his effort to introduce and promote the clinical application of amniotic membranes, and for serving as the inspiration for the present study.

REFERENCES

切除角結膜表皮細胞內腫瘤後以羊膜移植之效果

葉龍坤 林信璋 馬惠康

背 景：評估角結膜表皮細胞內腫瘤切除後以羊膜移植重建眼表層之效果。

方 法：收集5個角結膜表皮細胞內腫瘤病例，於1996年4月至2000年9月由醫師施行相同手
術切除後，再施以羊膜移植手術。羊膜以無菌標準程序處理後，覆蓋於術後裸露之
基膜上，以10-0尼龍線縫合，之後追蹤評估其效果。

結 果：平均追蹤期27月(6月至69個月)，所有病例眼表層表皮生長及癒合快速，無特別副作
用且無一復發病例。

結 論：羊膜移植手術對角結膜表皮細胞內腫瘤是一有效且安全之輔助治療。對於眼表層表
皮生長及重建，有良好的效果。

(長庚醫誌 2003;26:737-44)

關鍵字：角結膜表皮細胞內腫瘤，羊膜移植，復發。