Aerocyst urethral catheter insertion compared to expansion sponges application in external dacryocystorhinostomy

Yi Shao¹, Yao Yu², Chong-Gang Pei¹, Qiong Zhou¹, Wen-Jia Dong¹, Yangluowa Qu³, Lu Yang¹, Gui-Ping Gao¹

INTRODUCTION

Reconstruction of the lacrimal passages can be achieved by several surgical techniques, although external dacryocystorhinostomy (EXT-DCR), proposed for the first time by Ohm [1] and Onerci [2] in 1921, is still the most popular and successful operation. Recently, there were so many filling material applications in that operation. Plastic or rubber drainage tube, cosmolin or iodiform gauze strip and compressed gelatin are routinely used as antifibroblast proliferation materials in EXT-DCR in earlier period. But some of these materials have complications such as hemorrhagic after extubation, scar proliferation and nasal cavity erhythesis and so on. This study was to report the experience and the results of our service with the application of aerocyst urethral cathete in EXT-DCR, discuss the advantages and disadvantages of this technique comparing to the expansion sponges. This modification of EXT-DCR has not been previously published to the authors' knowledge.
MATERIALS AND METHODS

Patients This study was based on the retrospective analysis of 180 medical charts of patients (240 eyes) submitted to EXT-DCR in the First Affiliated Hospital of Nanchang University in the period from April 1, 2000 to April 1, 2005. All patients had severe epiphora, and most of those cases were accompanying with lacrimal sac bulge or pyogenic. The diagnosis of the chronic dacryocystitis disease was confirmed by the dripping test or infusion of fluorescein in the conjunctiva or lachrymal canaliculus (Jones test) without stained nasal drainage. The level of obstruction was defined by dacryocystography. Out of 180 patients, 120 presented unilateral obstruction and 60 had bilateral obstruction. The age at diagnosis ranged from 18 to 76 years old, with mean age of 46 years old and standard deviation of 18.8 years old (Table 1). This was a prospective interventional study, performed with Institutional Review Board approval. The study was also approved by the ethical Committee for Human Research of Nanchang University. Informed consent was obtained from all patients. According to the different filling material in the EXT-DCR, we divided the patients (180 patients, 240 eyes) into three groups randomly: negative control group (group 1), expansion sponges group (group 2) and acrocyst urethral catheter group (group 3, acrocyst urethral catheter: Unomedical Sdn, Bhd, Kedah, Malaysia; expansion sponges: Medtronic XOMED, Jacksonville, FL, USA). Patients were assessed according to surgical technique performed; intraoperative filling material and duration of insertion; number of surgeries necessary to correct; some aspects of postoperative follow-up, such as the use of antibiotics and the observed complications. Postoperative success was evaluated by DCR patency to irrigation, a positive dye test, hemorrhage conditions after extubation and subjective resolution of epiphora and liquor puris.

Methods To prevent nasal lateral wall, stypsis inferior nasal meatus and anesthetize nasal mucosa, we used several gauze strips packing soaked with 5g/L tetracaine or 50g/L cocaine HCl drops and 1g/L adrenaline drops which were inserted into the anodic of inferior nasal meatus before surgery. Group 1: A straight 15-18 mm skin incision placed 5-8 mm nasal to the medial canthus and tangential to the inferonasal rim of the orbit is made. Next the orbicularis muscle was bluntly dissected, and the anterior limb of the medial canthal tendon and the perioseum were exposed. The medial flap was forwards for about 10mm and raised from the bone with traction sutures. The lateral flap is elevated with the medial canthal tendon toward the anterior lacrimal crest and the lacrimal sac was then reflected laterally from the floor of the lacrimal fossa. Then, an osteotomy, 12mm×15mm wide, in the lateral nasal wall was created and the nasal mucosa was exposed. If an anterior ethmoidal cell was entered, the bony aperture is extended in order to expose the nasal mucosa [3]. Next, anterior and posterior flaps of lacrimal sac and nasal mucosa are created which look like the letter "H", as big and mobile as possible. 6/0 polyglycolic acid sutures are passed through the superior and inferior angles in anterior and posterior flaps of lacrimal sac and nasal mucosa and closed respectively. Group 2: interrupted suture and tie the posterior flaps, put expansion sponges group on, and interrupted suture anterior flaps, inject 1-2mL water into acrocyst, make the expansion sponges touch with and compress on the anterior and posterior flaps, and interrupted suture subcutaneous tissue. Group 3: take out of the plugging gauze strips in nasal cavity, interrupted suture the posterior and anterior flaps, stretch the double-cavity acrocyst urethral catheter into nasal mucosa, inject 1-2mL water into acrocyst, make the acrocyst located in and compress on the anterior and posterior flaps, and interrupted suture subcutaneous tissue. The skin wound is closed with a continuous intradermal suture of 6/0 silk. Meticulous haemostasis is carried out during all phases of the operation. A consultant ophthalmologist or an associate specialist performed the surgery. The persistenctime of filling material is recommended for 7 days after surgery. They were followed up a week later and the skin sutures were removed. The use of steroid drops and systemic antibiotics is recommended for a week after surgery. The patency of the dacryocystorhinostomy was examined after 7, 14, 21, 30 days and later every 4 months for the first 6 years. During the first examinations, if the dacryocystorhinostomy was found to be blocked, probing was performed, clots were removed, and steroid drops and systemic antibiotics were applied for one more week. Totally 170 patients (224 eyes) were asked to complete a telephone questionnaire to

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<th>Table 1</th>
<th>General condition of 180 patients (240 eyes) in EXT-DCR</th>
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<td>Group</td>
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evaluate the long-term postoperative improvement of their epiphora. The patients were asked to quantify their symptoms as follows: (A) no improvement, tearing is the same or worse; (B) slight improvement, but still tearing both indoors and outdoors; (C) significant improvement, but persistent slight tearing outdoors; (D) complete resolution of tearing, both indoors and outdoors. Patients self-reporting as A or B were defined as “failures,” C or D as “successes”.

Statistical Analysis Summary data were reported as mean±SD. SPSS 15.0 statistical software was used for data analysis. P <0.05 was considered statistically significant.

RESULTS Totally 180 patients attended the final review. Slight conjunctival hyperemia, eyelid congestion and edema with good skin could be seen in all patients at the week post-operation. Gauze and bandage was clean without any bleeding. Two weeks after surgery, conjunctival hyperemia and eyelid congestion and edema were relieved with ocular condition established well. In group 1 and group 2, there was no marked inflammatory reaction and subjective resolution of epiphora and liquor puris observed after 2 weeks. Postoperative pain was mild, and analgesics were applied. We did not observe complications such as epistaxis, healing retardation, infection and suppurition, subcutaneous emphysema, damage to orbital structures or visual affections. Patients presented mild periorbital ecchymosis and dorsum nasi collapse, without visual impairment and progressed with spontaneous resolution. On post-operative day 7, three cases needed probing and clot removal (one of them was a patient with severe coagulation problems due to thrombocytopenia) while five cases needed only probing only. Systemic antibiotics, such as first generation cephalosporins and topical ophthalmic drugs were used in the postoperative period, as well as frequent nasal lavage with sterile solution. Symptomatic outcomes at an average of 60 months follow up are summarized in Table 2 and Figure 1. The percentage of success in three groups were 73.7% (group 1), 86.5% (group 2), 98.7% (group 3), respectively. There was significant statistical difference in the surgical success rate and the operative complications (P <0.05). Compared to other groups, aecryocyst urethral catheter application had less complications, higher success ratio and no hemorrhage cases were found after extubation. At the end of follow up period, in group 3, only one patient with dacryocyst mucocele had epiphora symptoms of tearing and abnormal Jones I dye test.

DISCUSSION The history of lacrimal pathway surgery dates back from Hamurabi (2200 B.C.) [4]. Since then, techniques for treatment of lacrimal pathways have been developed, fighting against infections and restoring the transit of tears through the lacrimal system. The endonasal approach was described for the first time by Caldwell in 1893 [5], but it was forgotten for decades by the limited vision and the assessment of nasosinus anatomy. Currently, endoscopic DCR is a well-accepted and established technique in the treatment of the lacrimal sac and nasolacrimal duct disease. It is effective but with limited application due to its high cost and specification in underdeveloped countries and some developing countries.

To overcome the limits of traditional EXT-DCR, several authors have suggested modifications [6-10]. In this study we placed different filling material in the anastomotic stoma. There are some respective disadvantages in each surgery group, for instance, no filling material surgery always lead to anastomotic stoma collapse and healing retard; expansion sponges filled surgery always induce the difficulty of exelcymosis. Potential complications when the inflammation spreads include extension around the nasosinusitis or to the orbita (orbital cellulitis). In worst, an orbital abscess might develop that is potentially life threatening. Our previous
study found that aerocyst urethral cathete may as a new antifibroblast proliferation materials regulate the scar formation in dacryocyst mucocele. In this study, we used the aerocyst urethral cathete as the filling material in EXT-DCR, and tried to examine its effectiveness and mechanism after the long-term observation [11]. Aerocyst urethral cathete as the filling material in EXT-DCR have the following characteristics. Firstly, aerocyst urethral, a silica gel stuff, has been confirmed to have effects on the inhibition of fibroblast proliferation and scar formation without immunogenicity. Secondly, it has a stable chemical property and no tissue reaction. Thirdly, after water flooding, the water sac adjustly in the top can be satisfied with different patients. Fourthly, it may also be helpful for wound protection, in addition, it can act as a barrier to expansion space and to separate the surfaces which have potential possibility of adherence. Fifthly, aerocyst urethral catheter also plays an important role in stopping hemorrhage by oppression, preventing adhesion of anterior and posterior flaps by strong support and inhabiting fibrous tissue growth and reducing contraction of anastomosis. Lastly, it can prevent some postoperative complications, such anastomotic stoma collapse and healing retard, infection by exodus easily, the mechanical injury of anastomotic stoma, hemorrhage by difficulty of excycnosis without visual impairment and of spontaneous resolution. The expansion sponges, has been confirmed to have also effects on the inhibition of fibroblast proliferation and scar formation without immunogenicity, but the water absorption and the adhesiveness can cause the errophy and hemorrhage after the extubation.

Aerocyst urethral catheter plays an important role in stopping hemorrhage, preventing adhesion, inhabiting fibrous tissue growth and reducing contraction of anastomosis. Aerocyst urethral catheter is a safe, effective and ideal filling material for treatment of the chronic dacryocystis disease. Additional economic benefits could also be realised by removing the need for readmission for EXT-DCR. We believe that this modified EXT-DCR procedure is easy and quick to perform. It may be suitable not only for oculoplastic surgeons but also for ophthalmologists whose main area of interest is not lacrimal surgery. The results of this study demonstrate the reliability of our modification although a larger series of cases would be more conclusive.

Acknowledgements: We thank Dr. Hua Wang (Department of Ophthalmology, Xiangya Hospital of Zhongnan University), Gang Tan (Department of Ophthalmology, The First Affiliated Hospital of Nanhua University), Juan Peng (Department of Ophthalmology, the Second Affiliated Hospital of Guangzhou Medical College), for their generous support in clinic study.

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