



THE USE OF DCR SILICON STENTS TO CORRECT PTOSIS WITH POOR LEVATOR ACTION DURING FRONTALIS SLING SURGERY

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ABSTRACT

When levator function is poor or absent, frontalis sling is one of the preferred methods of treatment of ptosis. Various materials have been used for frontalis slings with varying results. The current retrospective study was performed in Al-Karama teaching hospital-Baghdad / Iraq during the period from 2014 to 2018, and aimed to find an effective, easily available and lasting frontalis sling material with minimal complications. DCR Silicone stents were used for brow suspension in cases of ptosis with poor levator action (less than 5 mm) in the form of a single rhomboid like in FOX method with slight modification. Retrospectively all the records of these patients were retrieved and the surgical results studied and analyzed. Mean follow up period was 2.6 years. Good long term functional results (assessed in terms of eyelid height, skin crease depth, eyelid contour and patient satisfaction) and achieved in 33 eyelids [90%]. Under correction was noticed in 3 eyelids [8%]. Re-surgery was done for these patients and all of them improved well. Lagophthalmos and mild corneal exposure were noticed in 6 eyelids which improved after the use of artificial tear eye drops and ointments. It can be concluded from this study that Silicone DCR stents tube is good alternative to AFL as a frontalis sling material as its functional results (90%) almost match that of AFL. In addition it has lesser complications compared to many other synthetic sling materials available. Compared to AFL, it has the advantage of avoiding second incision in the leg and can be done under local anesthesia.

KEYWORDS: Frontalis Sling Surgery; Ptosis; Ophthalmoplegia.

INTRODUCTION

Ptosis is an abnormal drooping of upper eyelid in relation to globe.^[1] The normal position of the upper lid is between the superior limbus and upper pupillary margin.^[2] (covering about 1-2 mm of upper cornea). Ptosis may be congenital or acquired. Congenital ptosis is usually unilateral (70%o-75% of ptosis) and is non-hereditary. Rarely congenital ptosis may be due to sympathetic paralysis (Congenital Horner's syndrome).^[3] Acquired ptosis is divided into four main types and these are neurogenic, Apo neurotic, mechanical and myogenic. Ptosis, when present, always produces a cosmetic problem. In children, it may be associated with astigmatism and can result in deprivation amblyopia in cases where it covers the pupil. Treatment is surgical correction. There are various surgical procedures to correct ptosis. In children, surgery should be performed when accurate evaluation can be done and the child is able to cooperate postoperatively. However, when there is danger of amblyopia or chin elevation with resultant

torticollis, it should be performed as early as possible. Primary goal of the surgery is elevation of the lid and maintenance of symmetry. The best results are obtained when the surgeon is able to balance the lids in primary position.^[2] Levator function is the pivotal factor in planning surgery for ptosis. When levator function is poor or absent, frontalis sling is one of the preferred methods.^[4] Various materials have been used for frontalis sling. Smith and Cherubini used 4/0 silk. Frieden Wald, et al. used a similar material but the technique was slightly different. Crawford has described the use of autologous fascia lata (AFL) which continues to be the gold standard for sling surgery.^[4,5] Some surgeons prefer non-autologous material because surgery can be done under local anesthesia in adult patients and a second incision (in leg) can be avoided. It is also preferred when AFL is not available (as in very young children with short legs). The other materials used are skin, silicone band, Silicone rods etc.^[5-8]

METHODS

The study was carried out between 2014 to 2018 and a single surgeon was involved in carrying out all the surgeries. Approval from the ethics committee of the hospital was obtained prior to the procedure. A total of 21 patients (29 eyes) with moderate to severe ptosis were included in the study. All patients had poor (4 mm or less) levator action. One patient (more than 50 yrs old) was having chronic progressive external ophthalmoplegia (CPEO) with bilateral ptosis. The rest were all of congenital type. Two patients, aged less than 4 years with severe congenital ptosis, one had bilateral ptosis with chin elevation and the other with upper lid severe ptosis were also included. All other patients were between 8 to 50 years of age. Most of the congenital cases were done under general anesthesia barring few who cooperated to be operated under local anesthesia. Adult case with CPEO were operated under local anesthesia. In this study, DCR (dacryocystorhinostomy) stents were used for frontalis suspension, which has the advantage of being easily available and at the same time being a relatively non-degradable and less reactive, was expected to have good long term functional benefits.

Fox pentagon procedure with modification was used in all cases. Instead of two direct stab incisions over the lid, we made a superficial transverse incision, including in it skin and superficial fibers of orbicularis muscle at about 4 mm from lid margin and then within this incision, two stab 2 mm cuts, corresponding to medial and lateral limbus were made to the depth of tarsus. Two bone depth incisions along the upper border of eye brows were made, each about one cm from the lateral and medial brow margins. Finally, a one cm bone depth incision was made on the forehead, about one and a half cm from the mid-point of an imaginary line joining the two eye brow incisions. The tube was then hooked or tied to a 3-0 silk suture attached to a 26 mm cutting needle. The needle was first passed from the forehead incision to the lateral brow incision and with little traction; the silicon tube would follow it. The needle was then passed deep to include the orbital septum and then to exit at lateral limbal level incision in the lid incision. From here, the

needle and thereby the tube was made to pass under the orbicularis (at tarsus depth) to exit next in the medial deep cut (corresponding to the medial limbus) on the lid incision. From here, the needle and thereby the silicon tube was passed up deep to orbital septum to come out at medial eyebrow incision. Finally, it was made to exit at the forehead incision where it meets the other end. The two ends are pulled to attain a lid height corresponding to the upper limbus (with expected fall of 1 mm or more postoperatively in sitting or standing position). Then the two ends were tied together making five to six knots together which was then buried deep through a pocket under the frontalis muscle. This pocket over the knot was sutured with 6-0 vicryl. Forehead wound was closed with non-absorbable interrupted sutures and the horizontal lid incision was closed with 6-0 continuous silk sutures. Brow incisions were not sutured. Finally, frost suture was applied to lower lid and the eye patched for 24 hrs. Patient was put on systemic broad spectrum antibiotic and an analgesic for 7 days post operatively. Frost suture was removed next day and patient was put on frequent artificial tear eye drops during the day and ointment during night. Lid and forehead sutures were removed on 5th day. Patients were followed at one month, 3 months and then 6 months for an average of (2-2.5) years.

RESULTS

The records of all these patients were retrieved and studied. The patients were followed for one to four years with a mean follow up period of 2 and half years. Among 21 patients (29 eyes) included in the study, 10 were females and 11 were males, although it does not depict the true gender distribution of the disorder because only the cases with poor levator action with good Bell's phenomenon were enrolled for this surgery. Ages of our patients ranged from 3.5 years to 60 years. The youngest patient in our study was a 3.5 year old child with moderate to severe degree of bilateral congenital ptosis with chin elevation. The next youngest was approximately 3 years and 10 months old with unilateral severe ptosis (pupil covered). Both were subjected to temporary sling surgery using same material to avoid ocular torticollis and deprivation amblyopia (Table 1).

Table (1): Age and laterality.

Age in years	No. of patients	Unilateral	Bilateral	Total no. of eyes involved
0-4	2	1	1	3
5-10	7	5	2	9
11-20	8	5	3	11
21-30	2	2	0	2
31-50	1	0	1	2
51-60	1	0	1	2

Table (2): Complications.

Complications	No. of cases	Percentage
Lagophthalmos with mild Corneal exposure	5	17.2%
Wound infection	2	6.9%
Granuloma formation	1	3.4%

Most of the patients in our series were in the age group of five to twenty years. Also, most of the cases were unilateral. This does not depict the actual laterality of the disorder, since only those patients enrolled for this surgery that had (1) moderate to severe degree of ptosis, (2) poor LPS function but with good preserved Bell's phenomenon. One patient was more than 50 years of age, he was a male and had chronic progressive external ophthalmoplegia (CPEO).

On each follow up visit, patients were assessed to evaluate the functional results and also to look for any complications associated with the surgery. Good functional results were assessed in terms of eyelid height (MRD I), skin crease depth, eyelid contour and patient satisfaction. Marginal reflex distance more than 3 mm in postoperative patients, a good skin crease and a satisfied

patient were each taken as a good result respectively during each follow up visit (Table 2).

Good long term functional results were achieved in most of the cases. In roughly above 86-96% of cases, adequate eyelid height, skin crease depth and proper eyelid contour were achieved. The overall patient satisfaction was also very good. In 2 of the cases where adequate lid height was not obtained in the first surgery, re-surgery was done and all showed satisfactory improvement (Figure 1). In 5 cases (eyes), there was mild to moderate lagophthalmos with mild corneal exposure, which was managed conservatively. Wound infection was found in 2 cases in our series, which was mild and was treated by systemic and topical antibiotics. One patient developed granuloma over the forehead wound which was excised.

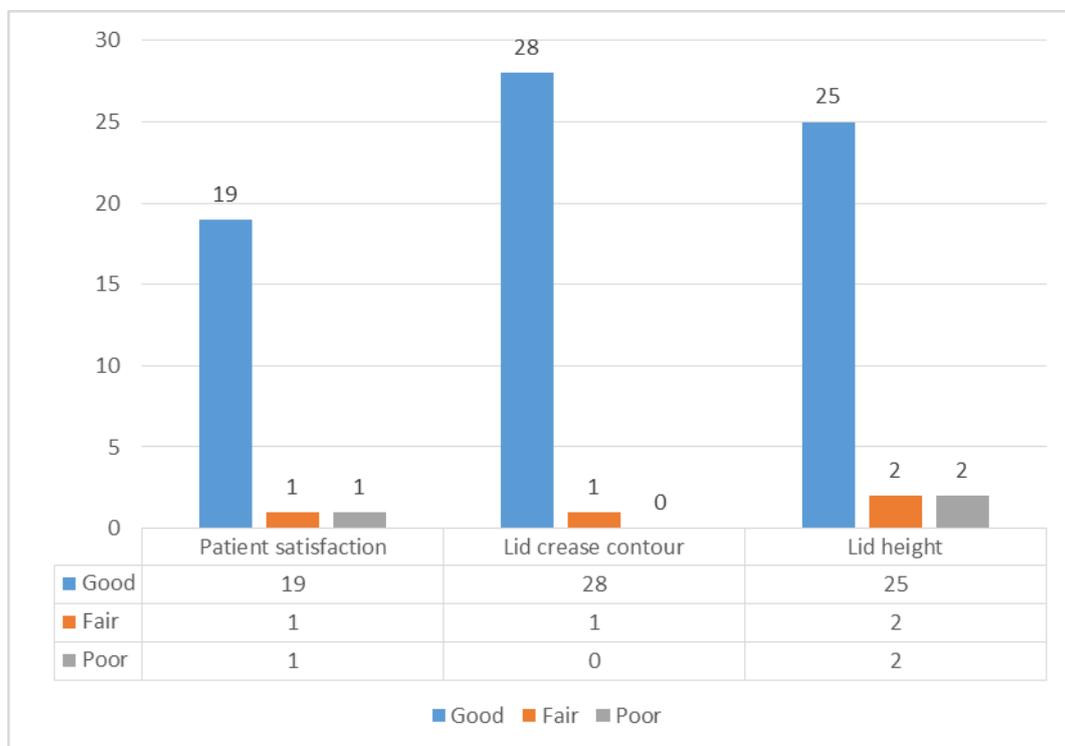


Figure (1): Assessment of functional results after surgery.

DISCUSSION

In ocular cases of poor LPS action, accessory elevators have always been used for elevating the lids.^[8] Frontalis sling surgery is the most commonly performed surgery for this group.^[9] Although none of the surgical procedures gives perfect results, it helps to alleviate the severity of the condition.^[10] AFL is the best material for frontalis suspension. It is non-degradable and allows fibrovascular in-growth leading to bio-integration without significant inflammation.^[11-14] AFL has good long term functional results with maintenance of lid height and well-formed skin crease. There are only occasional reports of post-operative wound infection. El-Toukhy *et al.* using AFL showed one wound infection out of 34 eyelids [3%]. The Wasserman *et al.* showed 2

infections out of 24 eyelids [8%].^[15] Deenstra *et al.* demonstrated 12 cases of exposure keratopathy with no wound infection out of 119 eyelids.^[16] Contrary to this, use of preserved fascia lata has a recurrence rate of 50% after 8-9 years of surgery.^[8]

Different non autologous materials have also been used, including silicone rod, monofilament nylon, polypropylene, polyester and expanded polytetrafluoroethylene. Carter *et al.* using silicon rod showed 3 cases of extrusion and 4 cases of recurrences out of 61 eyelids [11%], while Wasserman, *et al.* using nylon showed 9 cases of recurrences and one infection out of 13 eyelids.^[16-18]

In our study, of 29 eyes, good lid height was achieved in 86% and in other 7% cases fair lid height was achieved (this included the case of elderly patient with CPEO, where only fair lid height was aimed at during surgery to avoid post-operative exposure keratopathy because of associated elevation deficit). Good lid crease and contour with satisfactory cosmesis was attained in more than 96% of cases, which is much better compared to previous studies, probably in part due to the horizontal incision given across the lid. On the whole, good functional results were achieved in 90% of cases which almost matches AFL (autogenous fascia lata) sling (92%).

Silicone stent being relatively non-biodegradable, not much inflammatory reaction is seen since there is better bio-integration with the tissues. In our study, we had only 5 cases with mild corneal exposure keratopathy in the immediate post-operative period due to early lagophthalmos which improved with topical tear drops and ointment. The lagophthalmos, improved within 6-8 weeks, with meticulous use of artificial tear drops during the day and ointment at night (with eye patching during night as per need, for initial one or two weeks) during this time. Wound infection which has been reported in many studies with the use of various synthetic materials was seen in only 2 cases in the forehead wound in our series. It did not seem to have any relation with the sling material, as it was superficial (probably related to external stitches) which improved within few days with local antiseptics and systemic antibiotics.

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