Antegrade balloon dilatation of nasolacrimal duct obstruction in adults

A Kuchar, F J Steinkogler

Abstract

Aims—To determine the efficacy of antegrade balloon dilatation of postsaccal lacrimal stenosis in adults.

Methods—Balloon dilatation was performed in a series of 30 patients with complete nasolacrimal duct obstructions and epiphora. Obstruction was diagnosed by canalicular irrigation and transcanalicular endoscopic examination of the lacrimal pathway. Except for four cases in which general anaesthesia was applied, the procedure was performed under local anaesthesia. The Lacricath balloon catheter set was used. Silicone intubation was performed simultaneously. The time at which the tubes were removed depended on the findings at postoperative follow up but was, at the earliest, 3 months postoperatively.

Results—Success was objectified by irrigation and was evaluated subjectively at each follow up examination according to Munk’s scale. In all cases the procedure could be performed with subsequent silicone intubation. Three months postoperatively 89.9% of all cases were positive on simple irrigation, and subjective success was also registered (Munk’s grade 0 or 1). At 6 months 70% of all cases were positive on irrigation, again with subjective success (Munk’s grade 0 or 1). One year postoperatively 73.3% of all procedures showed subjective success (two successful redilatations would raise the success rate to 79.9%).

Conclusion—Retrograde as well as antegrade dilatation has been reported to be more or less successful in partial nasolacrimal obstruction. Although the procedure is used as primary treatment in cases of complete obstruction, it can still be performed under local anaesthesia on an outpatient basis. Long term observation will be required to prove the sustained effect of this procedure.

Materials and methods

Between September 1998 and July 1999 balloon catheter dilatation was performed in a series of 30 patients with complete obstructed nasolacrimal systems. Traumatic cases, tough presaccal stenoses, and patients with a history of former lacrimal surgery were excluded from this new procedure. In six patients a bilateral procedure was performed. In order to avoid exceeding sample size over subjects only the first treated obstruction of these patients was included in the study.

Complete obstruction was confirmed by canalicular irrigation and proved by transcanalicular endoscopy. Epiphora was evaluated subjectively according to a scale proposed by Munk and associates, which is graduated as follows: grade 0 = no epiphora; grade 1 = occasional epiphora requiring drying or dabbing less than twice a day; grade 2 = epiphora requiring dabbing two to four times per day; grade 3 = epiphora requiring dabbing five to 10 times per day; grade 4 = epiphora requiring dabbing more than 10 times per day; grade 5 = constant tearing.

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Accepted for publication 22 September 2000

Postsaccal obstruction with consequent epiphora has been treated by procedures involving the creation of a new anastomosis between the lacrimal sac and the nose. Currently the most common surgical procedure is external dacryocystorhinostomy (DCR). This procedure is associated with a high success rate and with few side effects. In an attempt to dispense with general anaesthesia and to avoid external scars, several procedures have been developed, including silicone intubation, and endonasal laser DCR, and endocanalicular laser procedures.

Lacrimal pathway dilatation is most frequently performed as retrograde dilatation, requiring radiographic and endonasal endoscopy, and is associated with a widely varying degree of success. Perry et al reported on antegrade dilatation in adult patients with partial nasolacrimal duct obstruction.

In this study antegrade dilatation in complete postsaccal stenosis after endoscopy of the lacrimal system was performed. In all the cases reported here, a bypass procedure would have been performed in the past.

The efficacy of this new procedure is still to be determined.

Figure 1 The Lacricath will be advanced until the upper mark has reached the superior punctum. This will be the most advanced position of the balloon.
All patients suffered from at least grade 3 epiphora (Munk’s classification). Nineteen female and 11 male patients were operated on by this new procedure after written consent had been obtained. The mean age of the patients was 55.8 (SD 14.7) years and the mean interval between the onset of epiphora and surgery was 14.4 (7.9) months. In four (13.3%) cases a stenosis of the common canaliculus necessitated an additional Er:YAG laser membranotomy in conjunction with postnasal dilatation. In these cases only a thin membranous membrane (less than 1 mm) was present and could be opened by a few laser effects. After this procedure probing and syringing into the lacrimal sac was very easy.

Dilatation was performed under general anaesthesia in the first four cases. All other patients were operated on under local subcutaneous anaesthesia. Lignocaine (lidocaine) 2% was infiltrated into the medial canthus and the perisaccal area subcutaneously. The mucosa of the nose was anaesthetised with lignocaine spray.

The procedure was started with initial syringing and probing of the lacrimal system. Since all patients presented with complete postsaccal obstruction, probing up to the obstruction was started carefully; a mechanical opening of the obstruction either by means of high pressure syringing or mechanical pressure of the probe was performed until fluid reached the nasal cavity. In a next step the balloon catheter was introduced into the superior canaliculus and advanced into the lacrimal sac and further to the nasal cavity passing the area of the stenosis after reopening. The catheter was advanced up to the superior mark on the catheter (Fig 1). The balloon was inflated with a pressure of 9 bar for 90 seconds. Since the length of the catheter is about 3.0 cm the stenotic area was dilated by the balloon. In the first four cases nasal endoscopy showed the inflated

![Figure 2](image1)  [Inflated balloon catheter just reaching the nasal cavity, blue prolene thread behind the catheter.]

![Figure 3](image2)  [View from the lacrimal sac through the reopened nasolacrimal duct into the nose. Blue prolene thread passing through the nasolacrimal duct.]

Table 1  Grade of epiphora according to Munk and the results of saline irrigation preoperatively and at each follow up examination

<table>
<thead>
<tr>
<th>No</th>
<th>Age</th>
<th>Sex</th>
<th>Duration of symptoms</th>
<th>Epiphora grade</th>
<th>Saline irrigation</th>
<th>Remarks</th>
</tr>
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<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Prop 3 months 6 months 12 months</td>
<td>3 months 6 months 12 months</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>37</td>
<td>F</td>
<td>13</td>
<td>4 0 0</td>
<td>O O O</td>
<td>Lost tubes 2 weeks post op</td>
</tr>
<tr>
<td>2</td>
<td>57</td>
<td>F</td>
<td>27</td>
<td>4 1 2</td>
<td>O P P</td>
<td>Massive intrasaccal inflammation</td>
</tr>
<tr>
<td>3</td>
<td>61</td>
<td>F</td>
<td>4</td>
<td>4 0 0</td>
<td>O O O</td>
<td>Er-YAG MT</td>
</tr>
<tr>
<td>4</td>
<td>41</td>
<td>F</td>
<td>23</td>
<td>3 0 0</td>
<td>O O O</td>
<td>Er-YAG MT</td>
</tr>
<tr>
<td>5</td>
<td>72</td>
<td>F</td>
<td>15</td>
<td>4 0 0</td>
<td>O O O</td>
<td>Er-YAG MT</td>
</tr>
<tr>
<td>6</td>
<td>44</td>
<td>F</td>
<td>20</td>
<td>3 1 1</td>
<td>O O O</td>
<td>Er-YAG MT</td>
</tr>
<tr>
<td>7</td>
<td>71</td>
<td>F</td>
<td>23</td>
<td>3 1 1</td>
<td>O P P</td>
<td>Severe allergic reaction</td>
</tr>
<tr>
<td>8</td>
<td>40</td>
<td>M</td>
<td>20</td>
<td>4 2 2</td>
<td>O P C</td>
<td>Lost tubes 1 week post op re op</td>
</tr>
<tr>
<td>9</td>
<td>61</td>
<td>M</td>
<td>15</td>
<td>3 0 3</td>
<td>O C O</td>
<td>Re-op</td>
</tr>
<tr>
<td>10</td>
<td>42</td>
<td>M</td>
<td>26</td>
<td>4 1 1</td>
<td>O P P</td>
<td>DCR</td>
</tr>
<tr>
<td>11</td>
<td>56</td>
<td>F</td>
<td>5</td>
<td>4 1 3</td>
<td>P C C</td>
<td>DCR</td>
</tr>
<tr>
<td>12</td>
<td>59</td>
<td>M</td>
<td>3</td>
<td>3 1 2</td>
<td>P P P</td>
<td>DCR</td>
</tr>
<tr>
<td>13</td>
<td>59</td>
<td>M</td>
<td>29</td>
<td>4 1 2</td>
<td>P P P</td>
<td>DCR</td>
</tr>
<tr>
<td>14</td>
<td>55</td>
<td>M</td>
<td>17</td>
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<td>O O O</td>
<td>Er-YAG MT</td>
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<td>15</td>
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<td>25</td>
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<td>O O O</td>
<td>Er-YAG MT</td>
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<td>16</td>
<td>87</td>
<td>F</td>
<td>15</td>
<td>4 1 0</td>
<td>O O O</td>
<td>DCR</td>
</tr>
<tr>
<td>17</td>
<td>60</td>
<td>M</td>
<td>8</td>
<td>3 1 1</td>
<td>P P P</td>
<td>DCR</td>
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<tr>
<td>18</td>
<td>73</td>
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</tr>
<tr>
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<td>5</td>
<td>4 2 2</td>
<td>P P P</td>
<td>DCR</td>
</tr>
<tr>
<td>20</td>
<td>36</td>
<td>M</td>
<td>4</td>
<td>3 1 1</td>
<td>O P P</td>
<td>DCR</td>
</tr>
<tr>
<td>21</td>
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<td>F</td>
<td>7</td>
<td>3 1 1</td>
<td>O O O</td>
<td>DCR</td>
</tr>
<tr>
<td>22</td>
<td>51</td>
<td>F</td>
<td>9</td>
<td>3 0 0</td>
<td>O O O</td>
<td>DCR</td>
</tr>
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<td>F</td>
<td>12</td>
<td>4 0 2</td>
<td>O P P</td>
<td>DCR</td>
</tr>
<tr>
<td>24</td>
<td>43</td>
<td>M</td>
<td>6</td>
<td>4 3</td>
<td>C C C</td>
<td>DCR</td>
</tr>
<tr>
<td>25</td>
<td>60</td>
<td>F</td>
<td>7</td>
<td>3 1 1</td>
<td>O P P</td>
<td>DCR</td>
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<td>26</td>
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<td>DCR</td>
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<td>O P P</td>
<td>DCR</td>
</tr>
<tr>
<td>30</td>
<td>44</td>
<td>F</td>
<td>14</td>
<td>3 1 1</td>
<td>O P P</td>
<td>DCR</td>
</tr>
</tbody>
</table>

O = open, P = partially closed, C = closed, Er-YAG MT = laser membranotomy.
balloon just reaching the nasal cavity (Fig 2). The stiff catheter could not be moved after inflation indicating the appropriate placing at the stenotic area. It was then deflated and placed at the lower mark of the catheter at the superior punctum. The balloon was inflated again at 9 bar for 60 seconds, and was then completely deflated and pulled out of the superior canaliculus. Immediate irrigation after dilatation demonstrated free passage into the nose. Endoscopic control after dilatation provided evidence of the reopened passage (Fig 3).

For splinting the system a silicone intubation was performed immediately after dilatation. The entire procedure required about 15–20 minutes.

Antibiotic (Nebacetin, neomycin + bacitracin) and steroid (Ultracortenol, prednisolone) drops were instilled into the system. The patients were instructed to use both drops for 2 weeks four times daily.

The silicone tubes were removed between 3 and 6 months postoperatively.

Follow up examinations were performed 1 week, 1 month, 3 months, and then every 3 months postoperatively. At each follow up, Munk's grade was determined, irrigation was performed, and complications were entered in a database.

Results
In all cases balloon dilatation could be performed after opening the obstruction and gaining access to the nose through the nasolacrimal duct. In five cases (16.6%) moderate bleeding occurred during the procedure; in all other cases only slight bleeding occurred and could be controlled by means of gentle pressure on the lids. In all cases a silicone intubation was performed immediately after dilatation. In six cases (20.0%) only a monocanalicular intubation could be performed because of canalicular or communis stenosis. No particular disadvantage due to local anaesthesia was observed. Several patients presented a blood tinged skin as a result of local anaesthesia and bloody nasal discharge for 1–3 days after the procedure. The most uncomfortable sensation was experienced when the tubes were pulled through the nose by means of a tiny blunt hook. In three cases (10.0%) the tubes were lost before the planned removal. In all other cases the tubes were removed 3 months postoperatively except for cases with Munk's grade 2 epiphora. In these cases tubes were removed after 6 months. The procedure was considered successful when epiphora had been completely eliminated or had been reduced to Munk's grade 1.

Three months after the procedure saline irrigation revealed complete free passage in 27 obstructions (90.0%). Eleven cases (36.6%) had grade 0 tearing and 16 cases (53.3%), grade 1 tearing. In three cases (patient nos 8, 19, 24) (10.0%) the grade of epiphora was 2 or 3.

Six month results were available in 29 cases. Twenty one cases revealed free passage after irrigation (70%) with Munk's classification 0 or 1 (eight, grade 0; 13, grade 1). Eight cases (26.6%) had Munk’s grade 2 or grade 3 epiphora. One patient (no 24) required a conventional DCR to cure the postnasal obstruction.

In one patient with a history of allergy the obstruction showed subjective improvement but did not meet the criteria for success (patient no 8). Two cases showed grade 3 epiphora. One patient lost the tubes 1 week after surgery and in one patient no particular cause could be found. In two patients (patient nos 9 and 12) a further dilatation was performed with consecutive silicone intubation and one additional patient (patient no 19) required a DCR after the 6 month follow up.

All cases (including those requiring redilatation) could be followed up for 1 year. In 25
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The course of the mean preoperative Munk's value compared with follow up examinations 3, 6, and 12 months postoperatively.

<table>
<thead>
<tr>
<th>X Labels</th>
<th>Preop</th>
<th>3 months</th>
<th>6 months</th>
<th>12 months</th>
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<tr>
<td>Minimum</td>
<td>3.000</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>25% Percentile</td>
<td>3.000</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Median</td>
<td>4.000</td>
<td>1.000</td>
<td>1.000</td>
<td>1.000</td>
</tr>
<tr>
<td>75% Percentile</td>
<td>4.000</td>
<td>1.000</td>
<td>2.000</td>
<td>1.000</td>
</tr>
<tr>
<td>Maximum</td>
<td>4.000</td>
<td>3.000</td>
<td>3.000</td>
<td>3.000</td>
</tr>
</tbody>
</table>

Antegrade dilatation is a procedure which can be performed on an outpatient basis under local anaesthesia. The entire procedure, including silicone intubation, requires no more than 15–20 minutes. Expensive equipment as in retrograde dilatation is not needed. The procedure can be performed easily, is repeatable and avoids an external approach through the skin.

Discussion

Most cases of acquired nasolacrimal duct obstruction in adults result from age related stenotic changes in the lacrimal drainage system. Stenosis may predispose the sac to stasis and infection, which causes pathological changes in the lacrimal mucous membrane.26–28

Balloon catheter dilatation may reopen a pathological obstruction and can lead to recanalisation of the lacrimal drainage system.

Several published reports describe retrograde dilatation requiring radiographic support.15–25 These procedures are very expensive, because of the radiological equipment, and time consuming. The long term success rates are reported to range between 90% and 25%.16–17 Our own results of retrograde dilatation combined with monocanalicular silicone intubation demonstrated a success rate of 60% in partial obstructed nasolacrimal ducts.18 In most of the previous reports no silicone intubation was performed and a higher rate of recurrent epiphora was registered. Consecutive silicone intubation for splinting the reopened stenosis seems to be very important for permanent patency during the initial scarring process.

Munk and others13 consider substantial improvement to consist of reduction of epiphora to grade 1 or grade 0. Accordingly, we used a reduction of epiphora to grade 1 or grade 0 as the criterion for subjective success. Three months after dilatation a positive irrigation could be achieved in 90.0%. A significant reduction of epiphora according to Munk's scale was regarded as evidence of a patent nasolacrimal duct. The 1 year success rate of 79.9% is high enough to warrant performance of this procedure as primary surgery in all cases with postnasal stenosis except in patients with a history of severe allergy, lacrimal sac tumours, former lacrimal surgery, or traumatic changes in the lacrimal bone. Perry and others reported similar results in patients with partial stenosis using antegrade dilatation.23 As reported in the latter paper we also performed silicone intubation in every case. Compared with previous reports of silicone intubation alone for lacrimal pathway obstruction a reporting success rates of 22%–78%, an additional dilatation in completely obstructed postnasal passages may improve the rate of success. Postoperative endoscopic control of the lacrimal pathway system demonstrated recanalisation and showed the silicone tubes to be in place in all cases. The silicone tubes were removed after 3 months, thus avoiding possible complications of long term silicone intubation.27

No difference was observed between patients with long standing obstructions and those suffering from epiphora of a few months' duration. Patients with a history of allergy did not meet our criterion of success in spite of silicone intubation and local steroid medication. Those patients are not suitable candidates for primary dilatation. In all other cases balloon catheter recanalisation may be an effective primary treatment. The fact that it can be performed under local anaesthesia makes it even more suitable.

Most recurrences of epiphora occurred during the first 6 months after treatment. Owing to persistent epiphora in spite of silicone intubation, four patients required a DCR between 4 and 12 months after initial dilatation. Endoscopy showed intrasaccal and postnasal stenosis, which could not be reopened successfully by dilatation. A further two patients lost their tubes and needed a repeated dilatation procedure within 6 months. Thus, in six cases (20.0%) primary dilatation failed and further surgery was required to cure epiphora.

Antegrade dilatation is a procedure which can be performed on an outpatient basis under local anaesthesia. The entire procedure, including silicone intubation, requires no more than 15–20 minutes. Expensive equipment as in retrograde dilatation is not needed. The procedure can be performed easily, is repeatable and avoids an external approach through the skin.

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and bone. Therefore, it is best suited as an initial approach in patients with acquired postacral stenosis except traumatic cases, lacrimal sac tumours, and patients with severe allergy.

In spite of encouraging 1 year results long term observations will be required to prove the efficacy of the procedure over several years.

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Br J Ophthalmol 2001 85: 200-204
doi: 10.1136/bjo.85.2.200

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