MR Dacryocystography in the Evaluation of Patients with Obstructive Epiphora Treated by Means of Interventional Radiologic Procedures

B. Coskun, E. Ilgit, B. Onal, O. Konuk and G. Erbas

*AJNR Am J Neuroradiol* 2012, 33 (1) 141-147
doi: https://doi.org/10.3174/ajnr.A2889
http://www.ajnr.org/content/33/1/141
MR Dacryocystography in the Evaluation of Patients with Obstructive Epiphora Treated by Means of Interventional Radiologic Procedures

BACKGROUND AND PURPOSE: Most imaging techniques used for the evaluation of obstructive epiphora, such as DS DCG, rely on undesired ionizing radiation. We evaluated the efficacy of topical contrast-enhanced MR DCG in comparison with DS DCG in patients with obstructive epiphora who underwent balloon DCG or stent placement.

MATERIALS AND METHODS: Thirty-six LDSs of 21 patients treated with balloon DCG (n = 11) or stent placement (n = 11) were examined with MR DCG and DS DCG. Contralateral LDSs (n = 14) were also evaluated in patients with unilateral disease. A sterile 0.9% NaCl solution containing 1:100 diluted gadolinium chelate was instilled into conjunctival sacs. The 3D FSPGR sequence was used with a 1.5T scanner. MR and DS DCG findings were scored and compared according to morphology of the lacrimal sac, junction, and NLD and the presence of contrast media in the nasal cavity.

RESULTS: Comparison of MR DCG and DS DCG findings showed no significant statistical differences in reference to anatomic locations according to the McNemar test (P ≈ .05). Good or very good agreement (κ value > 0.61) was observed according to the κ statistics.

CONCLUSIONS: Topical contrast-enhanced MR DCG is an effective and reliable noninvasive method for evaluation of the LDS in patients treated with IR procedures. This method avoids both cannulation and ionizing radiation and can, therefore, be repeated as often as is necessary in these complex patients.

ABBREVIATIONS: DCG = dacryocystography; DCR = dacryocystorhinostomy; DS = digital subtraction; FSPGR = fast-spoiled gradient-recalled; Gd-DTPA = gadolinium-diethylene-triamine penta-acetic acid; IR = interventional radiology; LDS = lacrimal drainage system; MIP = maximum intensity projection; NaCl = sodium chloride; NLD = nasolacrimal duct

Obstruction of the LDS, resulting in inadequate drainage of tears, can lead to intermittent or constant tearing, which is termed "epiphora." It is an annoying condition representing 3%–5% of clinical consultations in ophthalmology. Most pri-
termed "epiphora." It is an annoying condition representing

in 1993, the MR DCG evaluation of the LDS in patients

required for cannulation of the soft tissues.

The classic treatment of epiphora resulting from LDS obstructions is external or endonasal endoscopic DCR. Transluminal balloon dilation of the LDS has been proposed as an alternative to surgical treatment. Placement of nasolacrimal polyurethane stents in the LDS is another less invasive approach to the treatment of epiphora. The cause of epiphora can be diagnosed by physical examination, diagnostic clinical tests, and imaging procedures. DS DCG is currently considered to be the criterion standard among imaging techniques. It has several drawbacks, however, including its inability to provide a functional evaluation, its use of ionizing radiation, a requirement for cannulation of the nasolacrimal duct. The classic treatment of epiphora resulting from LDS obstructions is external or endonasal endoscopic DCR. Transluminal balloon dilation of the LDS has been proposed as an alternative to surgical treatment. Placement of nasolacrimal polyurethane stents in the LDS is another less invasive approach to the treatment of epiphora.

The cause of epiphora can be diagnosed by physical examination, diagnostic clinical tests, and imaging procedures. DS DCG is currently considered to be the criterion standard among imaging techniques. It has several drawbacks, however, including its inability to provide a functional evaluation, its use of ionizing radiation, a requirement for cannulation of the nasolacrimal duct. The classic treatment of epiphora resulting from LDS obstructions is external or endonasal endoscopic DCR. Transluminal balloon dilation of the LDS has been proposed as an alternative to surgical treatment. Placement of nasolacrimal polyurethane stents in the LDS is another less invasive approach to the treatment of epiphora.

Although MR DCG was first carried out by Goldberg et al in 1993, the MR DCG evaluation of the LDS in patients treated with either balloon DCG or nasolacrimal stent placement has not been previously reported, to our knowledge. In this study, we aimed to compare the findings of topical contrast-enhanced MR DCG with those of DS DCG in patients with LDS treated either with balloon DCG or stent placement and to determine the efficacy of the MR DCG technique in the evaluation of LDS obstructions treated by means of IR procedures.

Materials and Methods

Patients

This study was approved by our institutional review board, and informed consent was obtained from all patients. The patient population included 17 women and 4 men (mean age, 50 years; range, 37–73 years) who were treated by interventional techniques for obstructive epiphora. In all cases, disease-free canaliculi were confirmed. Eleven balloon dilation procedures and 11 stent placements were performed in 22 LDSs of 21 patients. In 5 LDSs, formerly placed stents had been removed before MR DCG. The polyurethane nasolacrimal stent is 35 mm long and has a mushroom proximal tip (5 mm in diameter and length) like a Malecot catheter. The outer diameter of the stent is 2 mm, and the luminal diameter is 1.5 mm. Contralateral LDSs (n = 14) were also evaluated in patients with unilateral disease. In 6 patients, unilateral MR DCG examinations were performed. Grading of epiphora was determined on the basis of the Munk et al classification (0 = no epiphora, 1 = epiphora requiring wiping the eye less than twice a day, 2 = epiphora requiring wiping 2–4 times a day, 3 = epiphora requiring wiping 5–10 times a
day, 4 = epiphora requiring wiping >10 times a day, 5 = constant tearing) at the time of MR DCG (Table 1). All patients had DS DCG 1 day to 3 months (average, 27.90 days) before MR DCG examinations. The interval between the last intervention and MR DCG was at least 1 year and at most 12 years (mean, 6.33 years; median, 5 years). Exclusion criteria included a history of severe allergy, being younger than 18 years of age, pregnancy, and/or contraindications to MR imaging, such as severe claustrophobia and incompatible metallic implants.

**MR Imaging**

Eye drops containing 1:100 diluted gadobutrol (Gd-DO3A-butrol, Gadovist 1.0 mmol/mL; Bayer Healthcare, Berlin, Germany) in a sterile 0.9% NaCl solution were administered to 1 or both conjunctival sacs 4 times at 1-minute intervals while the patient was in a sitting position with his or her head in hyperextension. Then the patient was asked to lie down on the MR imaging table, and another 2 drops were administered to the conjunctival sacs before a 3-inch dual coil was placed on both orbits. After the localizer images were obtained, another 2 drops were administered without the need to reposition the coil.

MR imaging was performed with a 1.5T system (Signa Excite; GE Healthcare, Milwaukee, Wisconsin). The gradient system operates with a maximum gradient strength of 33 mT/m and a slew rate of 120 T/m/s. The 3D FSPGR sequence was used to obtain images in the axial and coronal planes. The 3D FSPGR sequence was used to obtain images in the axial and coronal planes. The total image acquisition time was <15 minutes in all cases, as shown in Table 2.

Following the examination, patients were asked about the presence of discomfort due to contrast media installation or any adverse effects and their comments were noted.

### Table 1: Patients who underwent BD or nasolacrimal stent placement

<table>
<thead>
<tr>
<th>No.</th>
<th>Age (yr)</th>
<th>Sex</th>
<th>Epiphora* (grade)</th>
<th>Intervention</th>
<th>Time Intervala (yr)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>R</td>
<td>L</td>
<td>R</td>
</tr>
<tr>
<td>1</td>
<td>65</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>2</td>
<td>39</td>
<td>F</td>
<td>0</td>
<td>1</td>
<td>–</td>
</tr>
<tr>
<td>3</td>
<td>44</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>4</td>
<td>40</td>
<td>F</td>
<td>1</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>5</td>
<td>59</td>
<td>F</td>
<td>1</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>6</td>
<td>56</td>
<td>M</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>7</td>
<td>44</td>
<td>M</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>8</td>
<td>42</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>9</td>
<td>38</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>10</td>
<td>59</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>11</td>
<td>43</td>
<td>F</td>
<td>0</td>
<td>5</td>
<td>–</td>
</tr>
<tr>
<td>12</td>
<td>60</td>
<td>F</td>
<td>2</td>
<td>2</td>
<td>–</td>
</tr>
<tr>
<td>13</td>
<td>63</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>14</td>
<td>47</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>15</td>
<td>47</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>16</td>
<td>37</td>
<td>F</td>
<td>1</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>17</td>
<td>63</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>18</td>
<td>53</td>
<td>M</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>19</td>
<td>43</td>
<td>F</td>
<td>0</td>
<td>3</td>
<td>–</td>
</tr>
<tr>
<td>20</td>
<td>73</td>
<td>M</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>21</td>
<td>38</td>
<td>F</td>
<td>0</td>
<td>0</td>
<td>–</td>
</tr>
</tbody>
</table>

**Note:** BD indicates balloon DCG; +, present; –, absent; R, right; L, left.

*a At the time of MR DCG.

*b The time interval between the last intervention and MR DCG.

### DS DCG Imaging

All patients had DS DCG examinations 1 day to 3 months before the MR DCG examinations. Before the procedure, a topical anesthetic solution (0.4% benoxinate hydrochloride) was applied to the conjunctival sac. Following the dilation of the lower punctum, a flexible 23-ga lacrimal cannula was placed into the inferior canaliculus. After the acquisition of the mask images, 2–4 mL of nonionic contrast media was injected, and images (1 frame/s) were obtained in a sequence directed by an experienced radiologist blinded to the clinical findings of the patients. The MR DCG evaluation was reviewed in the following order: axial, coronal, and postacral images during imaging of the lacrimal drainage system was complete and the contrast had reached the nasal cavity or when the reflux of the contrast media toward the superior punctum was observed.

### Image Evaluation

MR images were reconstructed with the MIP algorithm by using a computer workstation (Advantage Windows Workstation 4.1, GE Healthcare). The images obtained by MR DCG and DS DCG were assessed by an experienced radiologist blinded to the clinical findings of the patients. The MR DCG evaluation was performed in the following order: axial, coronal, and postacral images during imaging of the lacrimal drainage system was complete and the contrast had reached the nasal cavity or when the reflux of the contrast media toward the superior punctum was observed.

### Statistical Analysis

Statistical analysis was performed by using commercially available statistical software (Statistical Package for the Social Sciences, Version 10.0 for Windows; SPSS, Chicago, Illinois). MR DCG and DS DCG findings were compared by using McNemar test and $\kappa$ statistics. For the McNemar test, a $P$ value of <.05 was considered to indicate a significant difference. $\kappa$ values for the $\kappa$ statistic were interpreted as follows: <0.00 represented poor, between 0.00 and 0.20 represented slight, between 0.21 and 0.40 represented fair, between 0.41 and 0.60 represented moderate, between 0.61 and 0.80 represented good agreement, and between 0.81 and 1.00 represented very good agreement.

### Results

MR DCG was performed after topical contrast administration and diagnostic images were obtained successfully for each patient.

A total of 36 LDSs were evaluated with MR imaging in 21 patients who had undergone DS DCG examinations. Balloon DCG was performed in 11 LDSs (Fig 2), stent placement was performed in 11 LDSs (Fig 3), and contralateral LDSs were evaluated in 14 patients with unilateral diseases. No side effects occurred during or after the instillation of diluted eye drops.

The MR DCG and DS DCG findings were statistically compared with the above-referenced scoring system. In all LDSs ($n = 36$), no significant difference was determined for the morphology of the lacrimal sacs, junctions, NLDs, and presence of contrast media in the nasal cavity according to the McNemar test ($P > .05$). According to the $\kappa$ statistics, MR DCG and DS DCG analysis of the morphology of the lacrimal sacs, junctions, and NLDs showed very good agreement ($\kappa >$...
the observation of contrast media presence in the nasal cavity showed good agreement ($\kappa = 0.786$).

In the intervened (balloon DCG and stent placement) LDSs ($n = 22$), no significant difference was determined in lacrimal sacs, junctions, NLDs, and the presence of the contrast media in the nasal cavity according to the McNemar test ($P > .05$). Very good agreement ($\kappa = 0.842$) was observed in the lacrimal sacs, and good agreement ($\kappa = 0.80$) was observed in the junctions, NLDs, and presence of contrast media in the nasal cavity with regard to the $\kappa$ statistics. The results of the statistical analysis are summarized in Table 4.

In 5 of the LDSs, discrepancies were noted between the findings of the 2 methods. All patients had been treated by balloon DCG, and the discrepancies are summarized in Table 5.

**Discussion**

For many years, the treatment of obstructive epiphora was surgical external DCR. Despite its high success rate, its drawbacks included being an invasive procedure, often requiring general anesthesia, and the development of facial scar tissue.\textsuperscript{2,8-12} In recent years, endonasal endoscopic DCR has been developed as a less invasive treatment. In addition, IR methods such as balloon dilation and nasolacrimal polyurethane stent placement have been effectively used. Both endoscopic and
IR-based methods are more easily tolerated by the patient compared with external DCR.\textsuperscript{4,5,13}

It is important to determine the level of the obstruction before any surgical or radiologic treatment procedure. In addition to DS DCG, numerous imaging modalities have been used to evaluate the site and type of obstructive epiphora; these imaging techniques have a variety of advantages and limitations and are discussed in detail below. The lens of the eye is the most sensitive tissue to ionizing radiation in the region being studied, with a risk of subcapsular opacities and cataracts with a threshold-dependent deterministic effect.\textsuperscript{14,15} Therefore, the absence of ionizing radiation would be a great advantage for an imaging technique.

DS DCG is the criterion standard for the diagnosis of LDS obstruction, given its high spatial resolution, but it is not a functional method of analysis if performed with cannula-
to the lens of the treated side and 38.5 mGy. Wilhelm et al.\textsuperscript{20} reported the mean radiation dose as 5.43 mGy.

Both of these techniques have the disadvantage of exposing the lens to significant doses of ionizing radiation.\textsuperscript{22,23} The lacrimal system includes lacrimal scintigraphy and CT DCG, which are useful imaging techniques for the evaluation of LDS.

We administered nonionic gadolinium-based MR imaging contrast agent (gadobutrol, Gd-DOTA-butrol; Gadovist 1.0 mmol/mL), which is more viscous (viscosity, 4.96 mPa·s at 37°C) and has twice the gadolinium concentration (1.0 mmol/mL) of the other gadolinium-based contrast agents. During the DS DCG examinations, Priebe et al.\textsuperscript{30} administered the same contrast media into the LDSs of 3 patients who had a history of severe allergic reactions to iodinated contrast media.\textsuperscript{30} No side effects or complications were reported in these patients.

Topical application is a more physiologic technique than the cannulation method. Contrast material, like tears, is propelled to the LDS by blinking, muscular contraction, and capillary action after topical administration\textsuperscript{21}; thus, the LDS is not distended, unlike in DS DCG with intracanalicular injection. All 5 of the discrepancies between the 2 methods (illustrated in Figs 4 and 5) can be explained by this difference.

Conclusions
MR DCG is a useful imaging technique for the evaluation of LDS in patients treated with IR procedures, and it compares

<table>
<thead>
<tr>
<th>Study</th>
<th>Year</th>
<th>No. of Patients</th>
<th>Cannulation</th>
<th>Topical</th>
<th>Sequence Technique</th>
</tr>
</thead>
<tbody>
<tr>
<td>Goldberg et al.\textsuperscript{6}</td>
<td>1993</td>
<td>11</td>
<td>Gd-DTPA</td>
<td>Gd-DTPA</td>
<td>T1WI (fat-sat)</td>
</tr>
<tr>
<td>Caldemeyer et al.\textsuperscript{11}</td>
<td>1998</td>
<td>11</td>
<td>–</td>
<td>Saline solution</td>
<td>FSE T2WI</td>
</tr>
<tr>
<td>Kirchhof et al.\textsuperscript{28}</td>
<td>2000</td>
<td>11</td>
<td>–</td>
<td>Gd-DTPA</td>
<td>T1 and T2WI (fat-sat)</td>
</tr>
<tr>
<td>Manfre et al.\textsuperscript{29}</td>
<td>2000</td>
<td>36</td>
<td>Gd-DTPA</td>
<td>Gd-DTPA</td>
<td>SE T1WI (fat-sat)</td>
</tr>
<tr>
<td>Yoshikawa et al.\textsuperscript{25}</td>
<td>2000</td>
<td>18</td>
<td>Gd-DTPA</td>
<td>Saline solution</td>
<td>FSE T2WI, SE T1WI</td>
</tr>
<tr>
<td>Takehara et al.\textsuperscript{21}</td>
<td>2000</td>
<td>8</td>
<td>Saline solution</td>
<td>–</td>
<td>Heavily T2WI</td>
</tr>
<tr>
<td>Karagülle et al.\textsuperscript{24}</td>
<td>2002</td>
<td>19</td>
<td>Gd-DTPA</td>
<td>–</td>
<td>3D FSPGR</td>
</tr>
<tr>
<td>Cubuk et al.\textsuperscript{32}</td>
<td>2010</td>
<td>35</td>
<td>Saline solution</td>
<td>–</td>
<td>Single-shot SE T2WI (fat-sat)</td>
</tr>
<tr>
<td>Our study</td>
<td>2011</td>
<td>21</td>
<td>–</td>
<td>Gd-BT-D03A</td>
<td>3D FSPGR</td>
</tr>
</tbody>
</table>

Note: –fat-sat indicates fat-saturated; SE, spin-echo; –, absent.
favorably with the criterion standard DS DCG. Our study is the first use of MR DCG in the evaluation of the LDS in a group of patients who underwent balloon DCS or stent placement with the topical application of a gadolinium-based contrast media with relatively high concentration. Among its advantages, this method requires no cannulation and avoids exposure of the radiosensitive lens to ionizing radiation.

References

Fig 4. Discrepancy of the MR and DS DCG in the asymptomatic patient who had been treated with balloon DCG. A and B. MR DCG reveals normal right LDS (A), and there is no obvious difference after the sac massage (B). C. MR DCG findings could not be confirmed at the DS DCG showing stenosis of the junction and NLD (arrow) with a dilated lacrimal sac (arrowhead).

Fig 5. Junctional stenosis in patient having grade 1 epiphora. A. MR DCG shows occlusion at the nasolacrimal junction (arrow) with a dilated lacrimal sac on the right. The left LDS is normal. B. After the lacrimal sac massage, there is no difference on the right and drainage of the left LDS with free flow of contrast media. C. DS DCG reveals severe junctional stenosis (black arrow) with dilation of the lacrimal sac (arrowhead). Passage of the contrast media to the nasal cavity is obvious (white arrow).