Quantification of lacrimal function after D-shaped field irradiation for retinoblastoma

S M Imhof, P Hofman, K E W P Tan

Abstract
To study the quantitative effects of megavoltage external beam irradiation in a D-shaped field in patients with retinoblastoma, biomicroscopy was performed in 61 patients and tear function tests (Schirmer-lactoferrin and lysozyme tests) on 45 eyes in 34 irradiated patients. The results were compared with those obtained in 25 non-irradiated control eyes. The Schirmer test was significantly diminished (p<0.001) in irradiated eyes, as were the lactoferrin and lysozyme values (p<0.001). A mild to severe keratitis was found in 17 of the 61 patients (28%). A significant correlation (p<0.005) was found between the Schirmer and the mean Schirmer values; the mean lactoferrin and lysozyme values were diminished in all patients but did not correlate significantly with the corneal abnormalities. These quantitative data, obtained in patients treated for retinoblastoma, confirm the qualitative data found in patients irradiated for other reasons such as orbital or sinus tumours. Irradiation for retinoblastoma is not a harmless treatment and serious late side effects have to be considered.

Methods
Irradiation technique
Radiation therapy is carried out on a 6 MV or 8 MV linear accelerator using a lateral D-shaped field of 26×32 mm. This D-shaped field is especially contoured to irradiate the entire retina with sparing of the radiosensitive lens as much as possible. Accurate positioning of the eye in the sharply collimated treatment field is easily performed by magnetic fixation of the eye to the beam defining collimator by use of a low vacuum contact lens. The axial dimensions of each eye are determined by ultrasonic biometry before treatment to ascertain the exact volume to be irradiated. The applied radiation dose is normalised at a minimum tumour dose of 45 Gy given in 15 fractions of 3 Gy at three fractions a week.

Statistical methods
To compare the outcome of the tear function tests and the biomicroscopy findings we used the two sample t test, the multivariate analysis, and the Mann-Whitney U test.

Slit-lamp biomicroscopy
Objectively, extensive slit-lamp biomicroscopy provided information on corneal epithelial scarring, punctate keratitis, and filamentosal keratitis. We divided our findings into mild keratitis (mild punctate keratitis and conjunctival redness) and severe keratitis (keratopathy and/or large epithelial punctate and/or filamentosal keratitis).
Tear function tests
All tear function tests were performed without anaesthetic eye drops and with both eyes closed.

Schirmer tear test strips. Those used were from Clement Clarke International Ltd (London), normal values >15 mm/5 minutes. In Schirmer’s blotting paper test one end of a strip of filter paper, 0.5 cm × 3.5 cm, is inserted into the lower conjunctival fornix folded at right angles over the ciliary body of the lid where it is left in position: the normal secretion ought to moisten at least 1.5 cm of the strip as measured from the fold in 5 minutes. The test is rough, but it provides an indication, at least, of excessive lacrimation or of marked hyposecretion or absence of tears. Lactoplate. The one used was from JDC, Culemborg (The Netherlands). Lactoplate contains a tear lactoferrin immunoassay test with rabbit antiserum to human lactoferrin in an agarose gel. It quantifies the lactoferrin concentration in tear fluid by radial immunodiffusion (Mancini test). Normal values are >1.0 mg/ml. A small round filter paper is placed in the lower conjunctival fornix. After 5 minutes the filter paper is taken out and put on the special test agarose gel. The Lactoplate is kept for 48–72 hours at a temperature of 21°C when the diffusion circles are measured. The lactoferrin concentration is measured by extrapolation of the diameter on the Lactoplate.

The lysozyme test. This is performed in the same way as the lactoferrin, with a round testing filter paper in the lower conjunctival fornix. It is then sent to the microbiology department. The patient’s filter paper is put on a DST (diagnostic sensitivity test) agarose gel with a testing microbe (Micrococcus lysodeikticus). The control is done with lysozyme extracted from ‘Huehnereiweiss’ which is put on DST agar gel to make a standard curve. Normal values are 1.425 mg/ml.

Results
SLIT-LAMP BIOMICROSCOPY
This was performed on 61 patients and the results are given in Table 1. Of 61 patients, 44 did not have corneal abnormalities, 10 patients had a mild keratitis, and seven had a severe keratitis.

Table 1 Biomicroscopy findings in irradiated patients

<table>
<thead>
<tr>
<th>Slit-lamp examination</th>
<th>No corneal abnormalities</th>
<th>Mild keratitis</th>
<th>Severe keratitis</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Radiotherapy</td>
<td>44</td>
<td>10</td>
<td>7</td>
<td>61</td>
</tr>
</tbody>
</table>

Table 2 Biomicroscopy findings and mean tear function values

<table>
<thead>
<tr>
<th>Mean values</th>
<th>No corneal abnormalities</th>
<th>Mild keratitis</th>
<th>Severe keratitis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Schirmer (mm)</td>
<td>16</td>
<td>14</td>
<td>4.6</td>
</tr>
<tr>
<td>Lysozyme (mg/ml)</td>
<td>1.185</td>
<td>0.860</td>
<td>0.620</td>
</tr>
<tr>
<td>Lactoferrin (mg/ml)</td>
<td>0.6</td>
<td>0.5</td>
<td>0.5</td>
</tr>
</tbody>
</table>

LACTOFERRIN CONCENTRATION
In 31 patients with 41 irradiated eyes the lactoferrin concentration ranged between 0 and 1.3 mg/ml, mean 0.6 mg/ml. In 18 non-irradiated eyes (14 patients) the lactoferrin concentration ranged from 0.4 to 2.2 mg/ml, mean 1.3 mg/ml. The difference in the lactoferrin concentrations between irradiated and non-irradiated eyes was significant (two sample t test, p<0.001) (Fig 2).

LYSOZYME CONCENTRATION
In 29 patients with 39 irradiated eyes lysozyme concentrations ranged from 0.2 to 2.40 mg/ml, mean 1.02 mg/ml. In 14 patients with 19 normal eyes the lysozyme concentrations were greater, with values between 0.8 and 3.6 mg/ml, mean 2.35 mg/ml.
2.2 mg/ml. With respect to the lysozyme concentration the difference between irradiated and non-irradiated eyes was significant (twosample ttest, p<0.001) (Fig 3).

Discussion
Late effects on the eye and orbit after ocular or orbital irradiation are well known. Complications may arise with all forms of radiotherapy, but vary with the different ocular tissues involved and the form of irradiation used.

Serious sequelae of external beam radiotherapy applied for retinoblastoma in often very young children are: radiation-induced orbital growth retardation, retinopathy, cataract, corneal damage, and dysfunction of the lacrimal gland. L-D-shaped external beam irradiation in the treatment of retinoblastoma reduces cataract formation. In addition, lateral D-shaped external beam irradiation is expected to spare the main and accessory lacrimal glands, because of their position just outside the field of irradiation. To investigate this hypothesis, we evaluated the quantitative effects on tear function of megavoltage external beam radiotherapy with a D-shaped field in patients with retinoblastoma.

The main and accessory lacrimal glands (of Wolfling and Krause) contribute to the formation of the middle aqueous tear film layer, consisting of water, electrolytes, enzymes, and proteins, such as lactoferrin and lysozyme. The watery component, lactoferrin and lysozyme, plays an important role in keeping the precorneal tear film stable and preventing the corneal epithelium drying up. Lacking these components will thus result in instability of the tear film and damage to the corneal epithelium, as can be seen in keratitis sicca.

In our patients, treated with external beam irradiation with a D-shaped field, we found a significant lessening of tear production (as measured with the Schirmer test) and a significant reduction in tear protein production when compared with a control group (Figs 1, 2, and 3). Although the patient group and the control group are not identical or randomised (which is extremely difficult in patients), we can conclude that the quantitative lacrimal function tests are significantly disturbed in children treated with external beam irradiation as a result of irradiation damage to the main lacrimal gland and, possibly, the accessory glands of Klause lying in the upper conjunctival fornix. As a consequence, these children have a diminished stability of the tear film and thus may be prone to (sub)epithelial keratopathies. In almost 30% of them these (sub)epithelial keratopathies were found and these lesions appeared to correlate significantly with decreasing Schirmer values. No such a significant correlation was found between the mean lactoferrin and lysozyme concentrations and the presence and severity of the corneal abnormalities, but this may be explained by the fact that the sample size of the patient group with mild or severe keratitis was too small.

We conclude that, although irradiation in patients with retinoblastoma has proved to be a valuable tool in tumour control with excellent survival rates, side effects are still considerable. In evaluating these late side effects we are convinced that a search for irradiation techniques which are as effective in tumour control, but less damaging, should be stimulated.

We acknowledge the help of Dr M Ph Mourits.