The evaluation of corneal endothelial permeability in PERK study patients*

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SUMMARY  Sixteen patients enrolled in the PERK study were evaluated by fluorophotometry 24 hours or six months after radial keratotomy. A comparison of eyes operated and not operated upon showed that endothelial permeability was not significantly altered 24 hours and six months after surgery. Aqueous humour flow rates and anterior chamber elimination coefficients were significantly higher 24 hours after surgery in the eyes operated on than in those not operated on. Six months after surgery there was no longer a significant difference in these factors. The increase in aqueous humour flow rates 24 hours after surgery may represent a subclinical breakdown in the blood-aqueous barrier.

Anterior radial keratotomy, a surgical procedure for the correction of myopia, is at present being evaluated by the PERK (Prospective Evaluation of Radial Keratotomy) study, to determine the predictability, safety, and short and long term effects of the surgery. The operation, as originally performed by Sato,1 consisted of anterior and posterior radial corneal incisions. Unfortunately, as many as 75% of his patients developed bullous keratopathy 10–15 years after surgery.23 Because of this severe complication, the surgery was altered by Fyodorov and Durnev1 to consist of only anterior radial corneal incisions. We carried out a study to evaluate the effect of anterior radial keratotomy, performed according to the PERK surgical protocol,5 on endothelial cell function, measured by fluorophotometry.

Subjects and methods

Seventeen of the 38 patients enrolled in the PERK study1 at the Mount Sinai Medical Center between December 1982 and October 1983 agreed to undergo fluorophotometry. One patient was excluded from the study owing to difficulty in data interpretation.

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Eleven patients were examined by fluorophotometry 24 hours after surgery and five patients six months after surgery.

Six of the patients examined 24 hours after surgery were male and five were female. Six had the right eye operated on, five the left eye. The patients were between 22 and 46 years of age, average 29 (SD 8) years. Preoperative spherical equivalent varied between −2.25 dioptres and −4.75 dioptres.

Five patients underwent fluorophotometry six months after radial keratotomy, three male and two female. Two patients had the right eye operated on, three the left eye. These patients were between 22 and 38 years of age, average 32 (SD 7) years. The preoperative spherical equivalent varied between −2.00 dioptres and −8.00 dioptres.

Radial keratotomy was performed by one of us (SAO) according to the PERK surgical protocol.1 It consisted of eight radial anterior corneal incisions to 100% of the depth of the thinnest paracentral area, measured with an ultrasonic pachymeter. A central 3 to 4 mm optical clear zone was left untouched, depending on the preoperative refraction, and the incisions did not extend past the limbus. In no case was the anterior chamber entered during surgery.

Fluorophotometry was performed with a slit-lamp fluorophotometer. Fluorescein sodium 0·25% and 0·4% benoxinate hydrochloride (Fluress, Barnes-Hind) was applied topically to both eyes, one drop every three minutes for 10 doses. Measurements of
fluorescence in the cornea and in the anterior chamber were begun two and a half hours after fluorescein application and continued every 20 minutes for four hours. The unoperated eye served as the control eye, and was compared with the operated eye in these experiments.

The data were analysed by a computerised model according to the method of Jones and Maurice on the assumption of an anterior chamber volume of 175 μl and a corneal volume of 70 μl. Kc, the anterior chamber elimination coefficient; Kcag, the cornea-to-aqueous transfer coefficient; Kcag, the corneal transfer coefficient referred to the volume of the anterior chamber; and F, the aqueous humour flow rate, were calculated. Corneal endothelial permeability was calculated on the assumption of a corneal stromal thickness of 0.47 mm. The unoperated control eyes were compared with the operated eyes for statistical analysis.

Gross outflow facility was measured on four patients at 24 hours and at one year after the initial radial keratotomy on the second eye, and on one patient at 24 hours and at one year after the second procedure on the second eye. Outflow facility was measured for four minutes on each eye with an electronic Schiötz tonometer. In three of these five patients the intraocular pressure was measured with a Goldmann applanation tonometer 24 hours and one year after surgery. The scleral rigidity in these three patients was calculated 24 hours and one year after surgery by the Friedenwald nomogram, the applanation intraocular pressure, and the initial intraocular pressure from the tonographic tracing. For statistical analysis the most recently operated eye was compared with the eye which had undergone surgery one year previously.

The paired t test was used to compare the treated and contralateral control eyes.

Results

Twenty-four hours after radial keratotomy was performed on 11 patients there was no significant difference in Kcag, Kcag, and the endothelial permeability in a comparison of operated and unoperated eyes (Table 1). Kc, was significantly (p<0.001) higher in the operated eye, 2.2±0.6 min⁻¹ (mean±SD), compared with the unoperated eye, 1.5±0.6 min⁻¹. F was significantly (p<0.001) higher in the operated eye, 3.9±1.0 μl/min, compared with the unoperated eye, 2.6±1.1 μl/min.

Six months after radial keratotomy there was no significant difference in Kcag, Kcag, endothelial permeability, Kc, or F, in a comparison of operated and unoperated eyes in five patients (Table 2).

Gross outflow facility was significantly (p<0.05) higher in five patients in the newly operated eye, 0.55±0.09 (mean μl/min mmHg±SD), compared with the eye which had been operated on the year before, 0.39±0.10 μl/min mmHg. Intraocular pressure, obtained from the tonograms, was not significantly different from the recently operated and the previously operated eyes respectively, 12.5±3.5 (mean mmHg±SD) and 14.8±3.2 mmHg. One year after surgery on the second eye the outflow facility was similar, 0.30±0.09 μl/min mmHg±SD in the most recently operated eye, and 0.31±0.13 μl/min mmHg in the contralateral eye. The intraocular pressure measured with a Goldmann applanation tonometer was not significantly different in the treated eyes 12.0±1.2 mmHg and the control eyes 13.0±3.7 mmHg one year after surgery. The ocular rigidity in three of these patients was similar, 0.031±0.015 (±SD) in the recently operated eye and 0.028±0.007 in the contralateral eye 24 hours postoperatively, and 0.028±0.004 and 0.026±0.003 at one year in the most recently operated eye and in the contralateral eye, respectively.

Discussion

Endothelial permeability, as measured by fluorophotometry, was unaltered at 24 hours and six months after radial keratotomy. Our results on endothelial permeability are similar to those of Hull et al in the rabbit. This group assessed endothelial permeability by measuring the flux of tritiated inulin
and carbon-14 labelled dextran, and by fluorophotometry. They did not demonstrate a significant alteration in endothelial permeability to any of these three substances between 24 hours and 10 weeks after radial keratotomy. Rabbit corneal endothelium responds to injury by proliferation, in contrast to human corneal endothelium. Hull used the same inulin and dextran radioactive flux technique in the owl monkey and demonstrated a transient increase in endothelial permeability to inulin in the operated eyes two days after radial keratotomy, with a return to levels comparable to those of the unoperated eyes four weeks after surgery. Dextran permeability was unaltered two days and four weeks following the procedure. The monkey corneal endothelium showed a transient increase in permeability to the smaller 1-4 nm inulin molecule, but not to the larger 3-8 nm dextran molecule. The fluorescein molecule measures 0-5 nm, but fluorophotometry was not performed on these monkeys. Recently Beatty and Smith found a persistent increase in corneal endothelial permeability to fluorescein during the first three months following radial keratotomy in the owl monkey. Corneal endothelial permeability in these animals was not increased from six months to two years following surgery.

$K_0$, the transfer coefficient of aqueous out of the eye, was significantly ($p<0.001$) increased 24 hours after surgery in our studies. This transient increase in $K_0$ may be due to a temporary alteration in the configuration of the anterior chamber, resulting in a temporary increase in outflow facility. Zimmerman et al. demonstrated a decrease in outflow facility following removal of through-and-through sutures from penetrating keratoplasting in eye bank eyes. Gross outflow facility was significantly ($p<0.05$) higher in five patients in the newly operated eye, $0.55 \pm 0.09$ (mean $\mu l/min mmHg \pm SD$), compared with the eye which had been operated on the year before, $0.39 \pm 0.10$ $\mu l/min mmHg \pm SD$. Intraocular pressure, obtained from the tonograms, was not significantly different in comparisons of the recently operated and the previously operated eyes, $12.5 \pm 3.5$ (mean mmHg $\pm SD$) and $14.8 \pm 3.2$ mmHg respectively. The coefficient of ocular rigidity was similar in the eyes examined 24 hours after surgery and the contralateral eyes. One year after surgery the outflow facility was no longer significantly different in treated and control eyes. The intraocular pressure and the coefficient of ocular rigidity were also similar one year after surgery.

The increase in aqueous humour flow rates in the operated eye 24 hours after surgery may represent a real increase. More probably it represents a subclinical breakdown in the blood-aqueous barrier. Clinically these patients did not show cells or flare on slit-lamp examination postoperatively. A real increase in aqueous humour flow is difficult to explain after radial keratotomy. Hull et al. reported a decrease in aqueous humour flow rates, measured by fluorophotometry, in rabbits one and 10 weeks after radial keratotomy.

The stability of endothelial permeability to fluorescein at 24 hours and six months after radial keratotomy in humans is encouraging. Two factors must be emphasised. Firstly, fluorophotometry is a mass measurement of corneal endothelial function; very small variations in permeability, or localised alterations in endothelial cell function, may not become apparent with this technique. Secondly, six months is a relatively short-term evaluation of endothelial function. The corneas Sato operated on did not show bullous keratopathy until 10 or more years after surgery; and he was directly injuring the endothelial cells with the posterior incisions.

Other methods of evaluating corneal endothelium after radial keratotomy have included specular microscopy. Studies in animal models (monkeys) suggest cell loss of 10–15% in the early postoperative period, which does not appear to progress over a 1–2 year follow-up period. Several groups have looked at endothelial cell counts in humans and have found 0 to 10% cell loss in the central cornea, the amount depending on the study. Studies of the peripheral cornea, directly under the incisions, have also not demonstrated significant cell loss or alteration in the first year after surgery.

Longer periods of endothelial evaluation postoperatively and more sensitive methods of evaluation would be helpful in determining alterations in corneal endothelial cell function after radial keratotomy.

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