

Comparison of Two Procedures: Photorefractive Keratectomy Versus Laser In Situ Keratomileusis for Low to Moderate Myopia

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Purpose: A prospective study was conducted to compare the effectiveness, safety, and stability of photorefractive keratectomy (PRK) and laser in situ keratomileusis (LASIK) for correction of low to moderate myopia.

Methods: Forty-five patients with a manifest refraction (PRK, -4.54 ± 0.80 ; LASIK, -4.82 ± 1.10) from -1.50 to -6.00 diopters (D) were treated and followed-up for 6 months. In each case, 1 eye received PRK and the other LASIK. The first eye treated, and the surgical method used in the first eye, were randomized. Uncorrected and corrected visual acuity, manifest refraction, corneal haze, and topographic analysis of ablation decentration were examined.

Results: The uncorrected visual acuity was 20/20 or better in 35 PRK eyes (77.8%) and 28 LASIK eyes (62.2%) at 6 months ($P = .107$). At 6 months, 28 eyes (62.2%) that received PRK showed a spherical equivalent of within ± 0.5 D as compared with 24 eyes (53.4%) that received LASIK ($P = .393$). The amount of ablation decentration was 0.37 ± 0.25 mm in PRK eyes and 0.49 ± 0.38 mm in LASIK eyes at 3 months ($P = .36$).

Conclusions: In our study, PRK and LASIK were found to be similarly effective and predictive of correction in low to moderate myopia. PRK has the advantage of less ablation decentration and is safer than LASIK, so we recommend PRK for eyes with low to moderate myopia. **Jpn J Ophthalmol 2001;45:487-491** © 2001 Japanese Ophthalmological Society

Key Words: Ablation decentration, LASIK, low to moderate myopia, PRK.

Introduction

Since the 193-nm excimer laser was introduced to the ophthalmology field, its applications to the treatment of myopia, astigmatism, and hyperopia have been increasing.¹⁻⁴ Photorefractive keratectomy (PRK) is believed to be a very safe corrective procedure for low to moderate myopia. However, postoperative pain, corneal haze, and myopic regression are known problems of PRK.⁵ Laser in situ keratomileusis (LASIK) may be a procedure preferred to PRK, particularly in cases with a high degree of myopia.

However, epithelial ingrowth, corneal flap-related complications, and corneal ectasia are recognized as shortcomings of LASIK.⁶⁻⁸ Some authors have reported promising results with LASIK for the correction of high myopia.^{9,10} However, in low to moderate myopia, it is still a point of contention and its use depends on the individual surgeon's preference. Therefore, to discern the efficiency and safety differences between the procedures, we compared the results of PRK in 1 eye and LASIK in the contralateral eye, performed on patients who had a low to moderate level of myopia ranging from -1.50 D to -6.00 D.

Materials and Methods

Forty-five patients with myopia (90 eyes) were enrolled in this study between January 1999 and September 1999. All the patients had received a full ex-

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planation of the procedures and informed consent was obtained before surgery. Each patient received PRK in 1 eye and LASIK in the other eye by the same surgeon (JBL). The first eye treated, and the surgical method used in the first eye, were randomized. The time interval between the procedures in both eyes was 2 weeks in all patients. The preoperative corrected visual acuity of all patients was 20/20 or better. Each patient received preoperative ophthalmic examinations that included slit-lamp microscopy, fundus examination, cycloplegic and manifest refraction, corneal keratometry, corneal topography, central corneal thickness, and Goldmann tonometry. Patients having systemic or ocular diseases such as diabetes mellitus, connective tissue disease, amblyopia, corneal disease, cataract, glaucoma, and retinal disease were excluded from the study.

Photorefractive Keratectomy Procedure

The procedure was done under topical anesthesia with proparacaine hydrochloride 0.5%. We used an excimer laser (Keratome II®; Coherent-Schwind, Neuostheim, Germany). At the completion of the surgery, a drop of ofloxacin 0.3% (Ofloxacin®; Sam-il Pharmacy, Seoul, Korea) and a drop of diclofenac 0.1% (Optanac®; Sam-il Pharmacy) were administered and a therapeutic contact lens (Hypaday®, diameter 14.2 mm, BC 8.7 mm; Chonan, Korea) was applied to the eye. After the epithelium had healed, Ofloxacin and fluorometholon 0.1% (Fluorometholon®, Sam-il Pharmacy) were administered four times a day and gradually tapered over 4 months.

Laser In Situ Keratomileusis Procedure

In the LASIK procedure, a microkeratome (Automated Corneal Shaper®; Chiron Vision, Claremont, CA, USA) was used. We used a number 160 thickness plate, which produced a cut 160 μm in thickness; the intended stromal bed thickness was at least 250 μm . This cut was followed by a midstromal ablation by the laser. Subsequent to the surgery, a drop of Ofloxacin 0.3% and a drop of diclofenac 0.1% were instilled. Ofloxacin and prednisolone 0.125% (Optilon®; Chonggundang Pharmacy, Seoul, Korea) were administered four times a day beginning 1 day after surgery and continuing for 1 week. We gradually tapered these over a 1-month period.

Following surgery, the uncorrected and corrected visual acuity and manifest refraction were measured at 1, 3, and 6 months after surgery, and the complications in both PRK and LASIK eyes were also re-

corded. From the difference map of topography (Orbscan®; Orbtex, Salt Lake City, UT, USA), ablation decentration from the pupil center was calculated. This was done by placing the cursor at the center of the ablation zone; from the screen, decentration from the pupil center was measured off in millimeters.

Subepithelial corneal haze levels were detected by slit-lamp examination and subjectively graded according to Hanna's method¹¹ at 6 months after PRK and LASIK surgery. Subepithelial haze was graded from 0 to 4 as follows: 0: totally clear; 0.5: a faint corneal opacity seen only by oblique indirect illumination; 1: an opacity of minimal density seen with difficulty with direct and diffuse illumination; 2: an easily visible opacity; 3: a denser opacity that significantly decreased the visualization of intraocular structures such as the iris and retina; and 4: an opaque cornea. Paired *t*-tests were used to compare the pre-operative data. Chi-square tests were used to compare the uncorrected visual acuity 20/20 or better and mean spherical equivalent refractions within ± 0.5 diopter (D) at 6 months after surgery. Values of $P < .05$ were considered statistically significant.

Results

In both the PRK- and LASIK-treated eyes, there was no statistically significant difference in spherical equivalent, average keratometry, intraocular pressure, or central corneal thickness between the 2 eyes before surgery (Table 1). The mean preoperative spherical equivalent refraction was -4.54 ± 0.80 D (range, -1.50 to -6.00 D) in the PRK eyes and -4.82 ± 1.10 D (range, -1.75 to -6.00 D) in the LASIK eyes. The mean preoperative amount of astigmatism was 0.73 ± 1.08 in PRK and 0.87 ± 1.21 in LASIK eyes. At 6 months, it was 0.34 ± 0.83 in PRK and 0.54 ± 0.71 in LASIK eyes, respectively.

At 1 week, 12 PRK eyes (26.7%) and 21 LASIK eyes (46.7%) could see 20/20 or better without correction. At 6 months, the uncorrected visual acuity was 20/20 or better in 35 PRK eyes (77.8%) and 28 LASIK eyes (62.2%) but this data showed no statistical significance ($P = .107$) (Table 2). Table 3 summarizes the refractive results during follow-up. At 6 months after surgery, 28 eyes (62.2%) that had received PRK showed a spherical equivalent of within ± 0.5 D as compared with 24 eyes (53.4%) that received LASIK ($P = .393$). Furthermore, 39 eyes (86.7%) that received PRK showed a spherical equivalent of within ± 1.0 D as compared with 38 eyes (84.5%) that received LASIK.

Table 3. Distribution of Refraction After Photorefractive Keratectomy (PRK) and Laser In Situ Keratomileusis (LASIK) in Same Patient*

Spherical Equivalent Refraction (D)	1 Month		3 Months		6 Months	
	PRK (n = 45)	LASIK (n = 45)	PRK (n = 45)	LASIK (n = 45)	PRK (n = 45)	LASIK (n = 45)
+2.0 to +1.6	1(2.2)	1(2.2)	1(2.2)	0(0.0)	0(0.0)	0(0.0)
+1.5 to +1.1	2(4.4)	1(2.2)	1(2.2)	1(2.2)	1(2.2)	1(2.2)
+1.0 to 0.6	7(15.6)	6(13.3)	4(8.9)	3(6.7)	2(4.4)	2(4.4)
+0.5 to 0.1	8(17.8)	5(11.1)	7(15.6)	4(8.9)	6(13.3)	3(6.7)
0 to -0.5	15(33.3)	15(33.3)	19(42.2)	19(42.2)	22(48.9)	21(46.7)
-0.6 to -1.0	9(20.0)	14(31.1)	9(20.0)	13(28.9)	9(20.0)	12(26.7)
-1.1 to -1.5	3(6.7)	3(6.7)	4(8.9)	5(11.1)	5(11.1)	5(11.1)
-1.6 to -2.0	0(0.0)	0(0.0)	0(0.0)	0(0.0)	0(0.0)	1(2.2)
Mean (D)	0.0	-0.3	-0.4	-0.5	-0.5	-0.6
Range (D)	-1.1 to +1.9	-1.2 to +1.8	-1.4 to +1.6	-1.5 to +1.4	-1.5 to +1.4	-1.7 to +1.3
SD	0.8	0.6	0.7	0.5	0.8	0.5

*Values are numbers of eyes, with percentages in parentheses, except for mean, range, and SD.

†D: Diopters, SD: standard deviation.

of 20/20 or better at 6 months after surgery. The possible causes for these results can be speculated as follows. First of all, the ablation decentration was less following PRK than LASIK. This may be due to the patient having difficulty in seeing the fixation light through an irregular corneal stroma after the corneal flap was flipped during the LASIK procedure.¹⁷ Secondly, there was more irregular astigmatism following LASIK than following PRK. Although we could not measure the irregular astigmatism from the topography, in LASIK, when the flap is placed at its original position following laser ablation, the corneal flap does not precisely return to its exact original position. We are able to observe a small contraction of the corneal flap towards the hinge, producing a small gap along the flap edge after surgery. Thus, irregular astigmatism tends to develop, which cannot be ignored.

Thirdly, prior to surgery, particularly in PRK, preliminary detailed and sufficient patient education was conducted, which included explanations of postoperative check-ups as well as precise application of the steroid eyedrops. That might have reduced the degree of corneal haze and myopic regression after PRK.

In addition, occurrence of complications was lower in PRK than in LASIK. An increase in intraocular pressure was seen in 1 PRK eye, which was controlled with a β -blocker. In an eye that had a +2 grade corneal haze after PRK, the manifest refraction at 6 months was sph -0.50 = cyl -0.50, Axis 180°; the uncorrected visual acuity was 20/30 and the corrected visual acuity was 20/20. In LASIK eyes, in the eye that had an operation for epithelial ingrowth, the manifest refraction at 6 months was sph -0.25 = cyl -1.00, Axis 180°; the uncorrected visual acuity

was 20/50 and the corrected visual acuity was 20/30 at 6 months. The manifest refraction of the free cap eye was sph -0.50 = cyl -2.00, Axis 150°; the uncorrected visual acuity was 20/70 and the corrected visual acuity was 20/50 at 6 months.

Conclusions

Although in this study PRK-treated eyes had a slower visual recovery in the early postoperative period and slightly more corneal haze that was not vision-threatening, the PRK procedure was safer than LASIK in low to moderate myopia cases. Careful selection of patients and sufficient education and understanding before surgery are considered necessary to achieve a good outcome.

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