

Fig. 4. Changes in the P1 peak latencies of the mfERG obtained from 4 mfERG areas in 14 diabetic eyes before and after PRP. The diabetic eyes were derived from 9 patients with preproliferative or early proliferative diabetic retinopathy showing no clinically significant macular edema in this series. All values were expressed as mean \pm SEM. Each bar indicates SEM. ** indicates P < 0.01 by Mann-Whitney's U test. PRP, panretinal photocoagulation; mo, month(s). Control: 15 normal control eyes of 14 healthy volunteers.

Fig. 5. Changes in the P1 – N1 amplitudes of the mfERG obtained from 4 mfERG areas in 14 diabetic eyes before and after PRP. All values were expressed as mean \pm SEM. Each bar indicates SEM. * indicates *P* < 0.05, and ** indicates *P* < 0.01 by Mann-Whitney's *U* test. PRP, panretinal photocoagulation; mo, month(s). Control: 15 normal control eyes of 14 healthy volunteers.

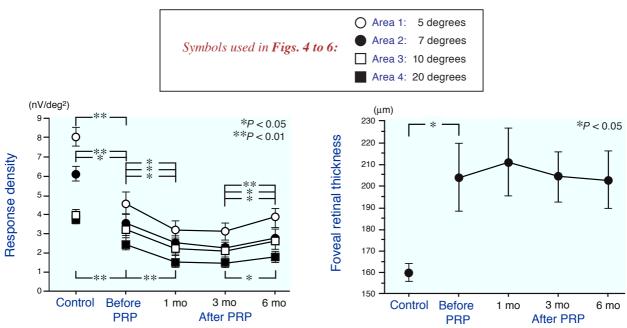


Fig. 6. Changes in the mean response density levels of the mfERG obtained from 4 mfERG areas in 14 diabetic eyes before and after PRP. All values were expressed as mean \pm SEM. Each bar indicates SEM. * indicates *P* < 0.05, and ** indicates *P* < 0.01 by Mann-Whitney's *U* test. PRP, panretinal photocoagulation; mo, month(s). Control: 15 normal control eyes of 14 healthy volunteers.

Fig. 7. Changes in the mean foveal retinal thickness within 5 degrees on OCT in 14 diabetic eyes before and after PRP. All values were expressed as mean \pm SEM. Each bar indicates SEM. * indicates *P* < 0.05 by Mann-Whitney's U test. PRP, panretinal photocoagulation; mo, month(s). Control: 16 normal control eyes of 12 healthy volunteers.

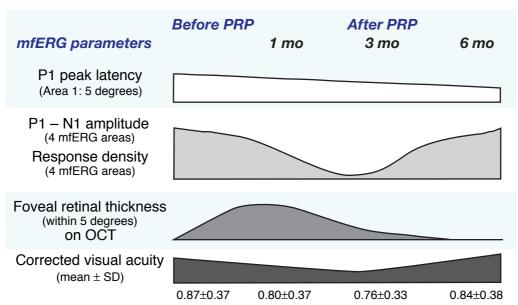


Fig. 8. Summarized data obtained in the present study before and after PRP in 14 diabetic eyes. In the mfERG parameters, the P1 peak latency from the area 1 (within a central radius of 5 degrees), the P1 – N1 amplitudes and the mean response density levels from the 4 mfERG areas are presented at 1, 3 and 6 months after PRP following each data before PRP in addition to the mean foveal retinal thickness within 5 degrees on OCT. The mean corrected visual acuity in the 14 diabetic eyes is also presented throughout the course. PRP, panretinal photocoagulation; mo, months(s).

same tendency as in the P1 – N1 amplitudes, as shown in Fig. 6. That is, the mean response density levels from the 4 areas were remarkably decreased in the 14 diabetic eyes before PRP as compared with those in the 15 control eyes at the 1% or 5% level, followed by a maximum decrease in the parameter at 3 months after PRP. However, remarkable recoveries were detected in the mean response density levels from the 4 areas at 6 months after PRP at the 1% or 5% level.

The mean foveal retinal thickness within 5 degrees on OCT was significantly increased in the 14 diabetic eyes before PRP as compared with the thickness in the 16 control eyes of the 12 healthy volunteers at the 5% level. Most remarkably, a transient increase in the thickness was detected in the diabetic eyes at 1 month after PRP, followed by a tendency for recovery at 3 to 6 months after PRP (Fig. 7).

Discussion

As summarized in Fig. 8, in the mfERG parameters, the P1 peak latency from the area 1 (within a central radius of 5 degrees) was markedly prolonged in the 14 diabetic eyes before PRP as compared with that in the 15 control eyes, but a tendency for recovery was detected throughout the course after the procedure (Fig. 4).

The P1 – N1 amplitudes and the mean response density levels from the 4 mfERG areas were remarkably decreased in the diabetic eyes before PRP as compared with those in the control eyes, followed by a maximum decrease in both parameters at 3 months after PRP. However, remarkable recoveries were detected in both decreased parameters from the 4 areas at 6 months after PRP (Figs. 5, 6 and 8).

The mean foveal retinal thickness within 5 degrees on OCT was remarkably increased in the diabetic eyes before PRP as compared with the

thickness in the 16 control eyes. Most remarkably, a transient increase in the thickness was detected in the diabetic eyes at 1 month after PRP, followed by a tendency of recovery at 3 to 6 months after the procedure (Figs. 7 and 8).

This time, statistical analysis was not undertaken on the correlation between the corrected visual acuity and each data from the mfERG or OCT examinations before and after PRP in each diabetic eye due to the relatively small number of samples. However, as demonstrated in Fig. 8, it is of note that the mean corrected visual acuity before PRP in the 14 diabetic eyes showed a gradual decrease at 1 month, followed by a maximum decrease at 3 months and a tendency of gradual recovery at 6 months after the procedure in each mean corrected visual acuity in those eyes, as in the changes in the P1 – N1 amplitudes and the mean response density levels of the mfERG obtained from the 4 mfERG areas in the 14 diabetic eyes before and after PRP.

The OCT findings showed no coincidental changes between mean foveal retinal thickness and mean corrected visual acuity in the 14 diabetic eyes throughout the course after PRP. However, tendency of recovery in mean foveal retinal thickness at 3 to 6 months after PRP may reflect a reversible foveal function resulting in a gradual recovery of the mean corrected visual acuity at 6 months after the procedure in diabetic eyes (Fig. 8), although we need to do further investigation on this point in more subject patients.

The mfERG derived from the fundus area, especially within 5 degrees of the central portion (Area 1) is supposed to reflect a cone-dominated response, which originates predominantly in the outer 70% of the retina, as in the full-field flash ERG (Hood et al., 1997; Palmowski et al., 1997; Fortune et al., 1999).

Optical coherence tomography provides noninvasively a cross- sectional tomographic image of the retina as described above and is more sensitive to small changes in retinal thickness than slit-lamp biomicroscopy (Hee et al., 1995, 1998; Kang et al., 2004).

Thus, the results obtained in the present survey indicate that the mfERG and OCT examina-

tions are useful for assessment of macular function before and after PRP in diabetic retinopathy, especially within 5 degrees of central portion, and that the effects of PRP on the macular function in this entity seem to be reversible at the foveal region, although we need a further investigation in relation to the outcome of visual acuity.

Acknowledgments: We express our sincere thanks to Professor Yoshitsugu Inoue, Division of Ophthalmology and Visual Science, Department of Medicine of Sensory and Motor Organs, School of Medicine, Tottori University Faculty of Medicine, for his generous help.

References

- Bresnick GH, Palta M. Oscillatory potential amplitudes. Relation to severity of diabetic retinopathy. Arch Ophthalmol 1987;105:929–933.
- 2 Chee CKL, Flanagan DW. Visual field loss with capillary non-perfusion in preproliferative and early proliferative diabetic retinopathy. Br J Ophthalmol 1993;73:726–730.
- 3 Diabetic Retinopathy Study Research Group. Photocoagulation of proliferative diabetic retinopathy. Clinical applications of diabetic retinopathy study (DRS) findings. DRS report number 8. Ophthalmology 1981;88:583–600.
- 4 Early Treatment Diabetic Retinopathy Study Research Group. Early photocoagulation for diabetic retinopathy. ETDRS report number 9. Ophthalmology 1991;98:766–785.
- 5 Fortune B, Schneck ME, Adams AJ. Multifocal electroretinogram delays reveal local retinal dysfunction in early diabetic retinopathy. Invest Ophthalmol Vis Sci 1999;40:2638–2651.
- 6 Frank RN. Visual fields and electroretinography following extensive photocoagulation. Arch Oph-thalmol 1975;93:591–598.
- 7 Hee MR, Puliafito CA, Duker JS, Reichel E, Coker JG, Wilkins JR, et al. Topography of diabetic macular edema with optical coherence tomographpy. Ophthalmology 1998;105:360–370.
- 8 Hee MR, Puliafito CA, Wong C, Duker JS, Reichel E, Rutledge B, et al. Quantitative assessment of macular edema with optical coherence tomography. Arch Ophthalmol 1995;113:1019–1029.
- 9 Hood DC, Seiple W, Holopigian K, Greenstein V. A comparison of the components of the multi-focal and

full-field ERGs. Vis Neurosci 1997;14:533-544.

- 10 Kang SW, Park CY, Ham DI. The correlation between fluorescein angiographic and optical coherence tomographic features in clinically significant diabetic macular edema. Am J Ophthalmol 2004;137:313–322.
- 11 Lewis H, Abrams GW, Blumenkranz MS, Campo RV, Vitrectomy for diabetic macular traction and edema associated with posterior hyaloidal traction. Ophthalmology 1992;99:753–759.
- 12 Massin P, Duguid G, Erginay A, Hauchine B, Gaudric A. Optical coherence tomography for evaluating diabetic macular edema before and after vitrectomy. Am J Ophthalmol 2003;135:169–177.
- 13 Palmowski AM, Sutter EE, Bearse MA Jr, Fung W.

Mapping of retinal function in diabetic retinopathy using the multifocal electroretinogram. Invest Oph-thalmol Vis Sci 1997;38:2586–2596.

- 14 Sutter EE, Tran D. The field topography of ERG components in man. I. The photopic luminance response. Vision Res 1992;32:433–446.
- 15 Yoon HY, Lee J, Kim YJ. Preservation of retinal sensitivity in central visual field after panretinal photocoagulation in diabetics. Korean J Ophthalmol 1996;10:48–54.

Received August 27, 2004; accepted September 24, 2004

Corresponding author: Akihiko Tamai, MD