

Comparison of Retinal Nerve Fiber Layer Thickness Measurements Using Time Domain and Spectral Domain Optical Coherence Tomography, and Visual Field Sensitivity

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Purpose: To investigate the relationship between the peripapillary retinal nerve fiber layer (RNFL) thickness, as measured by Stratus time domain optical coherence tomography (OCT) (Stratus OCT) and Cirrus spectral domain OCT (Cirrus HD-OCT), and the severity indices of the visual field (VF) defects in glaucoma patients.

Methods: This was a prospective, cross-sectional study. Correlations between the individual VF sensitivity at 52 test points and the Stratus OCT and Cirrus HD-OCT, which determined peripapillary RNFL thicknesses from 6 sectors, were calculated in 54 eyes with open-angle glaucoma and 22 normal control eyes. The association between the RNFL thickness and VF sensitivity was evaluated using a second-order regression model.

Results: A significant correlation was found for the RNFL thicknesses determined by the 2 OCT devices ($r=0.51$ to 0.95 ; $P<0.001$). VF sensitivities at each test point were also significantly correlated with the sectoral RNFL thicknesses. The inferotemporal RNFL sector exhibited the highest coefficient of determination, whereas the superotemporal test point had the highest VF sensitivity (Stratus, 0.70; Cirrus, 0.62).

Conclusions: The structure-function relationship was comparable between Cirrus HD-OCT and Stratus OCT RNFL measurements.

Key Words: peripapillary retinal nerve fiber layer thickness, optical coherence tomography, severity indices of the visual field

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After the introduction of optical coherence tomography (OCT) in clinical practice in 1995, it became possible to carry out routine measurements of retinal thickness. The first commercially available model, OCT 1 (Carl Zeiss Meditec; Humphrey Division, Dublin, CA) had an axial resolution of about $15\ \mu\text{m}$ and a scan velocity of 100 axial scans per second. In 2002, the OCT 3000 model (Stratus OCT; Carl Zeiss Meditec) was introduced. This device used time domain technology, which initially had an axial resolution of $10\ \mu\text{m}$ and a scan velocity of 400 axial scans per second. Recently, the Cirrus HD-OCT (Carl Zeiss Meditec) has become commercially available, with a spectral domain platform that has an axial resolution of $5\ \mu\text{m}$ and a scan velocity of 27,000 axial scans per second. Although the

Stratus OCT and Cirrus HD-OCT have a similar retinal nerve fiber layer (RNFL) measurement locations, they use different technologies. The Stratus OCT measures the RNFL thickness along a 3.46-mm-diameter circle around the optic disc in the fast RNFL mode, whereas the Cirrus HD-OCT resample data along a circle with a 1.73-mm radius around the optic disc.

Glaucoma is a progressive optic neuropathy in which there is loss of retinal ganglion cells that leads to visual field abnormalities. To diagnose and manage glaucomatous optic neuropathy, it is of critical importance to be able to detect the relationship between structural and functional damage. Sensory tests and various imaging techniques have been used to measure this relationship,^{1–3} with significant correlations between perimetric tests and optic disc parameters found for the area of the neuroretinal rim or RNFL thickness. The perimetric mean defect of the whole visual field along with the mean RNFL thickness, as measured by Stratus OCT and Cirrus HD-OCT, have been used to determine the quantitative structure-function association.⁴ However, local correspondence may be of greater importance in being able to judge the progression of glaucoma.

The purpose of this study was to investigate and compare the local structure-function associations obtained when using the Cirrus HD-OCT and the Stratus OCT devices.

PATIENTS AND METHODS

A total of 54 glaucoma patients and 22 normal participants were enrolled in the study and followed up during the period from October 2008 through January 2009 at Kagawa University Hospital. Among these patients, 34 had primary open-angle glaucoma and 20 had normal-tension glaucoma. All eligible participants received a detailed explanation of the study and signed an informed consent form in accordance with the principles embodied in the Declaration of Helsinki. All the participants underwent a complete ophthalmic examination that included visual acuity testing with refraction, intraocular pressure, and dilated fundus examination with stereoscopic biomicroscopy of the optic nerve head using slit lamp and indirect ophthalmoscopy. To be included in the study, all participants had to have a best corrected visual acuity of 20/40 or better, a spherical error within a range between 4.0 and -6.0 diopters, and a cylinder within ± 2.0 diopters. Exclusion criteria included history of any kind of retinal pathology, retinal laser procedure, retinal surgery, or neurologic disease. One eye in each participant was randomly chosen for inclusion in the study. The glaucomatous eyes were defined as the eyes exhibiting structural

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glaucomatous changes (vertical cup-disc asymmetry between fellow eyes of more than equal to 0.2, a cup-disc ratio of more than equal to 0.6, and a neuroretinal rim narrowing, notches, localized pallor, or RNFL defects with glaucomatous visual field (VF) loss in the corresponding hemifield). The glaucomatous VF defects were defined as those with consecutive, repeated abnormal standard automated perimetry (SAP) results (≥ 2 contiguous points with a sensitivity loss of $P < 0.05$ in the superior or inferior arcuate areas, a 10-dB difference across the nasal horizontal midline at greater than 2 adjacent locations, or an abnormal glaucoma hemifield test result). The eyes with preperimetric glaucomatous optic neuropathy were excluded from this study.

Stratus OCT Retinal Nerve Fiber Layer Measurement

The fast RNFL 3.4 scan protocol was used to measure the peripapillary RNFL (software version 4.0.1). The RNFL thickness was determined at 256 circumpapillary points on a circle with a default diameter of 3.4 mm around the center of the optic disc. We excluded all poor-quality scans, which were defined as those with a single strength of < 6 ; the presence of overt misalignment of the surface detection algorithm on $\geq 15\%$ of consecutive A-scans or 20% of cumulative A-scans; or overdecentration of the measurement circle location, as subjectively assessed.

Cirrus HD-OCT Retinal Nerve Fiber Layer Measurement

The Cirrus HD-OCT uses spectral domain technology of an optic disc cube obtained from a 3-dimensional data set composed of 200 A-scans from each of 200 B-scans that cover a 6-mm² area centered on the optic disc. After creating an RNFL thickness map from the cube data set, the software automatically determined the center of the disc and then extracted a circumpapillary circle (1.73-mm radius) from this data set. The same signal strength cutoff value and exclusion criteria used for the Stratus OCT were also used for these measurements.

All images were acquired during a single visit by 1 well-trained technician. Right-hand orientation was used for documentation of the clock hour measurements in both the Cirrus HD-OCT and Stratus OCT.

TABLE 1. Clinical Characteristics of the Study Population

	Glaucoma	Normal	P
Age (y)	61.7 ± 11.8	63.1 ± 9.0	0.62
Sex (male/female)	19/35	14/8	0.19
Diagnosis			
POAG	33		
NTG	21		
Refraction (D)	-1.51 ± 2.47	-1.18 ± 2.25	0.59
Visual field MD (dB)	-7.7 ± 7.5	-1.4 ± 1.6	< 0.01

MD indicates mean deviation; NTG, normal-tension glaucoma; POAG, primary open-angle glaucoma.

Visual Field Examination

Standard visual field testing was done using static automated white-on-white threshold perimetry (Humphrey Field Analyzer II; Carl Zeiss Meditec). A visual field was defined as reliable when fixation losses, false-positive errors, and false-negative errors were less than 20%.

Statistical Analysis

The relationship between the 2 OCT measurements was assessed by Pearson correlation analysis. The structure-function relationships from the 6 measured regions were adapted from the map earlier created by Garway-Heath et al (Fig. 1).⁵ The relationship between RNFL thickness and visual field mean deviation (MD) was evaluated using a second-order regression analysis, as it has been shown that the second-order regression analysis yields a stronger structure-function association than the first-order and other nonlinear regression analyses.^{6,7} Comparisons of the strength of the structure-function association were evaluated by paired *t* tests on the log-transformed absolute value of the studentized residuals from the 2 second-order regression models. A Bland-Altman plot was graphed to assess the agreement between 2 instruments. All statistical values are presented as mean ± standard deviation (SD). In all statistical analyses, *P* values < 0.05 were considered statistically significant (Fig. 1).

RESULTS

The demographic characteristics are detailed in Table 1. As the normal control participants were selected

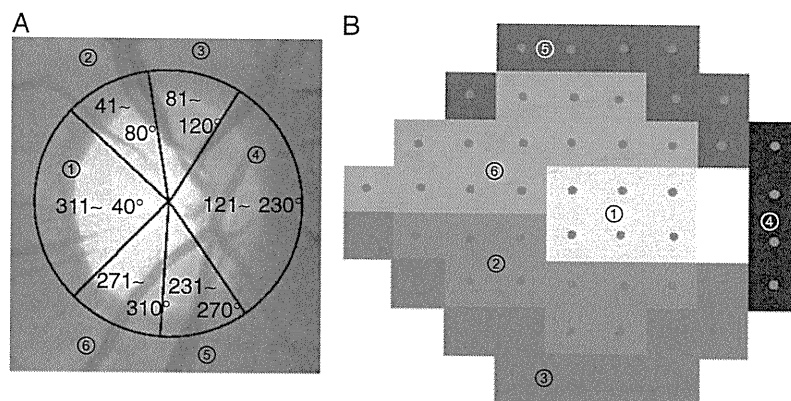


FIGURE 1. A topographic map generated using the methodology of Garway-Heath et al.⁵ Areas of the peripapillary retinal nerve fiber layer are divided into 6 sectors (A), and the corresponding visual field regions are observed in standard automated perimetry (B).

TABLE 2. Correlation of RNFLT Measurement on Average and in Each of 6 Sectors as Determined by Stratus and Cirrus OCT

	RNFLT by Stratus (µm)	RNFLT by Cirrus (µm)	Correlation of Determination (R)	P
Average	74.3 ± 17.9	72.5 ± 13.6	0.91	< 0.001
Temporal, 311 degrees-40 degrees	62.6 ± 17.7	60.3 ± 12.5	0.83	< 0.001
Superotemporal, 41 degrees-80 degrees	86.5 ± 33.4	87.1 ± 28.7	0.88	< 0.001
Superonasal, 81 degrees-120 degrees	90.1 ± 30.8	85.8 ± 24.4	0.80	< 0.001
Nasal, 121 degrees-230 degrees	62.7 ± 14.4	65.5 ± 9.5	0.51	< 0.001
Inferonasal, 231 degrees-270 degrees	87.0 ± 25.1	77.1 ± 20.0	0.85	< 0.001
Inferotemporal, 271 degrees-310 degrees	92.8 ± 38.1	86.9 ± 33.6	0.95	< 0.001

OCT indicates optical coherent tomography; RNFLT, retinal nerve fiber layer.

with age-matching and refractive error-matching, there was no significant difference in the mean age between the normal control participants (63.1 ± 9.0 y; range, 49 to 74) and those with glaucoma (61.7 ± 11.8; range, 35 to 85). The mean deviation (MD) in SAP ranged from -28.73 to -0.78 dB with a mean of -7.7 ± 7.5 dB in patients with glaucoma. Many glaucomatous eyes had early to moderate diseases based on the standard visual field severity grading,⁸ of the 54 patient's eyes, 28 (52%) were classified as early, 13 (24%) were classified as moderate, and 13 (24%) were classified as severe.

The average RNFLT thickness, as measured by Stratus OCT, was 74.3 ± 17.9 µm, whereas the average RNFLT, as measured by Cirrus HD-OCT, was 72.5 ± 13.6 µm. Only the inferonasal Cirrus HD-OCT RNFLT thickness measurement (77.1 ± 20.0 µm) was thinner than those measured by the Stratus OCT (88.0 ± 25.1 µm) (*P* < 0.05, unpaired *t* test). The Pearson correlation coefficient for mean RNFLT thickness in each sector between the 2 instruments was 0.51 to 0.95 (Table 2), with the weakest correlation found in the nasal sector (*r* = 0.51; *P* < 0.001). Figure 2 shows the Bland-Altman plots for the agreement between Cirrus HD-OCT and Stratus OCT for the average RNFLT thickness. For thinner RNFLT thicknesses, Stratus measurements tend to be thinner than Cirrus, whereas for thicker RNFLT thicknesses, Stratus measurements tend to be thicker than Cirrus.

The 52 VF test points were clustered into regional groups of VFs that corresponded to the RNFLT sections. Coefficients of determination between these 6 VF sensitivity (VFS) clusters and the corresponding 6 sectors are listed in Table 3. Except for the nasal sector, there were significant correlations between the VFS at individual test points and in several of the RNFLT sectors. The highest calculated coefficient of determination was for the superotemporal RNFLT sector. There were no significant differences noted in the strength of the structure-function association between the Cirrus HD-OCT and Stratus OCT.

DISCUSSION

The results of this study show that Cirrus RNFLT measurements correlate well with those from Stratus OCT, with comparable strength of the structure-function association between Cirrus HD-OCT and Stratus OCT. However, although we noted a correlation of the RNFLT thickness measurements between Stratus OCT and those from Cirrus OCT, other studies have earlier reported systematic differences in the measurement values between Cirrus HD-OCT and Stratus OCT.⁹⁻¹¹ Although our results showed a high correlation between the Cirrus HD-OCT and Stratus OCT thickness measurements, the RNFLT thickness measured by the Cirrus HD-OCT was only thinner than that measured

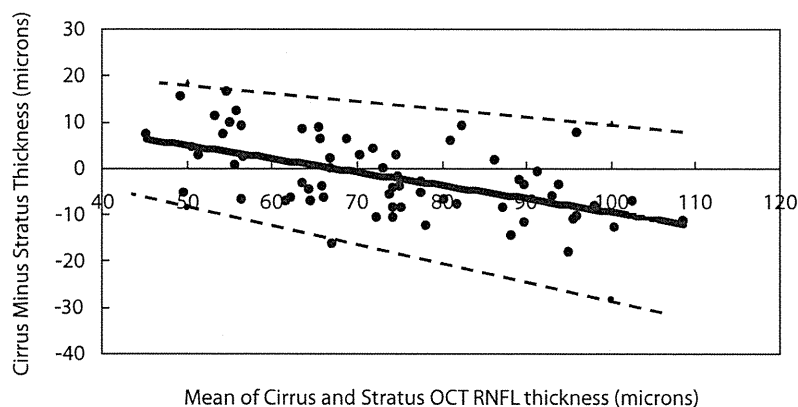


FIGURE 2. Bland-Altman plot of the agreement of mean retinal nerve fiber layer (RNFLT) thickness between Cirrus OCT and Stratus OCT. The difference (Cirrus OCT RNFLT thickness—Stratus OCT RNFLT thickness) between both measurements is plotted against the average of both measurements (*r* = 0.55, *P* < 0.01). The dotted lines of equality are plotted the 95% limits of agreement. OCT indicates optical coherence tomography.

TABLE 3. Comparison of Structure-function Relationship using Stratus or Cirrus OCT by Garway-Heath et al

Sector in RNFL	Correlation of Determination (R)		P
	Stratus OCT	Cirrus OCT	
Temporal, 311 degrees-40 degrees	0.39	0.32	0.31
Superotemporal, 41 degrees-80 degrees	0.70	0.62	0.07
Superonasal, 81 degrees-120 degrees	0.54	0.57	0.63
Nasal, 121 degrees-270 degrees	0.03	-0.06	0.48
Inferonasal, 231 degrees-270 degrees	0.46	0.37	0.17
Inferotemporal, 271 degrees-310 degrees	0.53	0.51	0.65

OCT indicates optical coherent tomography; RNFL, retinal nerve fiber layer.

by Stratus OCT in the inferonasal sector. Although a good agreement was found for all parameters, Bland-Altman plots showed that for thinner RNFL thicknesses, Stratus measurements tend to be thinner than Cirrus, whereas for thicker RNFL thicknesses, Stratus measurements tend to be thicker than Cirrus. We suggest that the RNFL thickness measured by the Stratus OCT can not be transferred to Cirrus HD-OCT thickness measurements because of this discrepancy.

In this study, glaucoma was defined on the basis of structure and VF loss. We wanted to investigate the local structure-function associations. To facilitate this procedure, we included patients with glaucomatous optic neuropathy with VF loss and normal participants with a perfectly normal RNFL without VF loss. The patients with preperimetric glaucoma were excluded from this study. Therefore, we could not find any superiority Cirrus HD-OCT than Stratus OCT in finding RNFL defects.

In our study, 1 examiner scanned each patient on the same 2 instruments during a single day. As eye position relative to the scanning circle can affect the RNFL profile, it is important to keep the scan circle within a consistent location to make serial assessments using the Stratus OCT.¹² As the Cirrus HD-OCT uses a different data-acquisition process that has a considerably higher scan speed, it can be used to obtain 3-dimensional data sets. The Cirrus HD-OCT uses this data set to locate the center of the optic disc and then draws a circle with a 1.73-mm radius. The Cirrus HD-OCT has a theoretical advantage over Stratus OCT in terms of scan registration. The scan registration process carried out by the Cirrus algorithms is fully automated, reducing the likelihood of operator error. We assumed that there was good eye fixation during the RNFL measurements using the Stratus OCT device in all of our participants. However, as we excluded participants with scans that had signal strengths less than 6, we cannot definitively draw any generalized conclusions that apply to all participants, including those with poor signal strengths.

Kanamori et al¹³ reported finding significant correlations between the VFS at individual test points and in several of the RNFL sectors when using the Stratus OCT for measurements. Leung et al⁴ recently reported that there was no significant difference in the strength of the structure-function association between Cirrus HD-OCT and Stratus OCT (average RNFL thickness and visual field MD). With the exception of the nasal sector, we found that the strength of the structure-function association was similar between Cirrus HD-OCT and Stratus OCT in our study. Both instruments measure RNFL thickness at a diameter of

3.46 mm. Glaucomatous structural changes of the RNFL usually occur first in the inferotemporal and superotemporal areas, with corresponding functional VF losses occurring first in the so-called Bjerrum areas of the upper and lower hemifields. Therefore, VFS might be more closely related to the inferotemporal and superotemporal RNFL thickness as opposed to the temporal and nasal RNFL thickness. However, it is very difficult to estimate the structure-function relationship at the nasal area adjacent to the optic disc owing to the paucity of test points on SAP at the temporal regions adjacent to the blind spot.

In conclusion, a significant association with the VFS was noted for both Cirrus HD-OCT and Stratus OCT when used to measure sectoral RNFL thickness. Further longitudinal studies will be needed to determine the applicability of using the structure-function association to describe glaucoma progression.

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Success Rates of Trabeculotomy for Steroid-Induced Glaucoma: A Comparative, Multicenter, Retrospective Cohort Study

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- **PURPOSE:** To evaluate the surgical outcomes of trabeculotomy for steroid-induced glaucoma.
- **DESIGN:** Multicenter, retrospective cohort study.
- **METHODS:** At 17 Japanese clinical centers, 121 steroid-induced glaucoma patients who underwent trabeculotomy between 1997 and 2006 were reviewed. Surgical failure was defined by the need for additional glaucoma surgery, deterioration of visual acuity to no light perception, or intraocular pressure ≥ 21 mm Hg (criterion A) and ≥ 18 mm Hg (criterion B). Surgical outcomes were compared with those of 108 primary open-angle glaucoma (POAG) patients who underwent trabeculotomy and 42 steroid-induced glaucoma patients who underwent trabeculectomy. Prognostic factors for failure were evaluated using the Cox proportional hazards model.
- **RESULTS:** The probabilities of success at 3 years for trabeculotomy for steroid-induced glaucoma vs trabeculotomy for POAG was 78.1% vs 55.8% for criterion A ($P = .0008$) and 56.4% vs 30.6% for criterion B ($P < .0001$), respectively. At 3 years, the success of trabeculotomy for steroid-induced glaucoma was comparable to trabeculectomy for steroid-induced glaucoma for criterion A (83.8%; $P = .3636$), but lower for criterion B (71.6%; $P = .0352$). Prognostic factors for failure of trabeculotomy for steroid-induced glaucoma were previous vitrectomy (relative risk [RR] = 5.340; $P = .0452$ on criterion A, RR = 3.898; $P = .0360$ for criterion B) and corticosteroid administration other than ocular instillation (RR = 2.752; $P = .0352$ for criterion B).
- **CONCLUSIONS:** Trabeculotomy is effective for controlling intraocular pressure < 21 mm Hg in steroid-induced glaucoma eyes. (*Am J Ophthalmol* 2011;151:1047–1056. © 2011 by Elsevier Inc. All rights reserved.)

STEROID-INDUCED GLAUCOMA IS A FORM OF OPEN-angle glaucoma associated with various modalities of corticosteroid administration such as oral, intravenous, inhaled, ocular instilled, intravitreal, and periocular.^{1–6} Some histologic studies have reported the accumulation of extracellular matrices including basement membrane-like material,^{7–9} fine fibrillar-like material,⁸ or proteoglycans⁹ in the trabecular meshwork of steroid-induced glaucoma patients. These observations suggest that such accumulation could lead to an increased resistance to aqueous outflow in the trabecular meshwork of steroid-induced glaucoma patients.

Surgical procedures for intraocular pressure (IOP) reduction in eyes with steroid-induced glaucoma include trabeculectomy,^{2,10,11} trabeculotomy,^{4,12} viscocanalostomy,¹³ and laser trabeculoplasty.^{14–18} Although several case series have shown that these surgeries are effective for IOP reduction, surgical outcomes for steroid-induced glaucoma are not fully understood due to lack of large case-control studies aiming to investigate the success rates of trabeculotomy in steroid-induced glaucoma eyes. It has previously been reported that trabeculotomy more effectively reduces IOP in adult Japanese patients with exfoliative glaucoma than primary open-angle glaucoma (POAG).¹⁹ This IOP-lowering effect in eyes with exfoliative glaucoma is thought to be attributable to the relief of abnormally increased outflow resistance that was induced by the accumulation of exfoliative material within the trabecular meshwork.

For the same reason, trabeculotomy has been the surgical procedure of choice for adult patients with steroid-induced glaucoma among Japanese surgeons.²⁰ We previously showed that trabeculotomy helped to reduce IOPs to 21 mm Hg or less in 14 Japanese patients with steroid-induced glaucoma.¹² However, large-scale, comparative clinical data remain elusive on, for example, whether trabeculotomy is more effective for steroid-induced glaucoma than POAG, whether trabeculotomy for steroid-induced glaucoma offers better IOP management than other surgeries such as trabeculectomy with mitomycin C (MMC), or which characteristics of patients with steroid-induced glaucoma exhibit better prognosis after trabeculotomy. To evaluate the surgical outcomes of trabeculotomy for steroid-induced

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TABLE 1. Patients With Steroid-Induced Glaucoma and Primary Open-Angle Glaucoma who Underwent Trabeculotomy

	SIG-LOT, n (%) (n = 121)	SIG-LET, n (%) (n = 42)	P Value	POAG-LOT, n (%) (n = 108)	P Value
Female	62 (51.2)	26 (61.9)	.232 ^a	38 (35.2)	.014 ^a
Right eye	62 (51.2)	22 (52.4)	.899 ^a	50 (46.3)	.455 ^a
Age (years), mean ± SD	38.4 ± 17.6	42.3 ± 17.9	.153 ^b	45.2 ± 15.0	.001 ^b
Preoperative IOP (mm Hg), mean ± SD	38.1 ± 10.0	35.6 ± 8.3	.169 ^b	28.9 ± 8.4	<.001 ^b
Combined sinusotomy	20 (16.5)	—	—	33 (30.6)	.012 ^a
Previous cataract surgery	17 (14.0)	4 (9.5)	.626 ^c	5 (4.6)	.029 ^c
Previous vitrectomy	6 (5.0)	0 (0.0)	.320 ^c	0 (0.0)	.054 ^c
Diabetic mellitus	13 (10.7)	6 (14.3)	.736 ^c	10 (9.3)	.709 ^a
Hypertension	18 (14.9)	8 (19.0)	.695 ^c	15 (13.9)	.832 ^a
Cause of corticosteroid use					
Atopic dermatitis	21 (17.4)	4 (9.5)	.335 ^c		
Uveitis	25 (20.7)	11 (26.2)	.457 ^a		
Collagen disease	37 (30.6)	17 (40.5)	.240 ^a		
Route of administration					
Ocular instillation only	17 (14.0)	12 (28.6)	.591 ^a		
Posterior sub-Tenon's injection of TA	13 (10.7)	1 (2.4)	.178 ^c		
Intravitreal injection of TA	10 (8.3)	0 (0.0)	.121 ^c		
Oral administration	72 (59.5)	26 (61.9)	.784 ^a		
Intravenous administration	3 (2.5)	2 (4.8)	.826 ^c		
Corticosteroid administration for more than 3 months after surgery	68 (56.2)	25 (59.5)	.708 ^a		

IOP = intraocular pressure; POAG-LOT = primary open-angle glaucoma patients who underwent trabeculotomy; SD = standard deviation; SIG-LET = steroid-induced glaucoma patients who underwent trabeculectomy with mitomycin C; SIG-LOT = steroid-induced glaucoma patients who underwent trabeculotomy; TA = triamcinolone acetonide.

^aP values are based on the χ^2 for independence test.

^bP values are based on Mann-Whitney U test.

^cP values are based on the χ^2 for independence test with Yates' correction.

glaucoma, we retrospectively reviewed clinical charts at 17 clinical centers in Japan.

METHODS

• **PATIENT SELECTION AND SURGICAL PROCEDURES:** We retrospectively reviewed the medical records of patients with steroid-induced glaucoma who underwent trabeculotomy or trabeculectomy with MMC and those with POAG who underwent trabeculotomy between January 1, 1997, and December 31, 2006, at the following 17 clinical centers in Japan: Kumamoto University Hospital (Kumamoto), Niigata University Medical and Dental Hospital (Niigata), University of Tokyo Hospital (Tokyo), Kanazawa University Hospital (Kanazawa), Gifu University Hospital (Gifu), Kagawa University Hospital (Miki), University of Yamanashi Hospital (Chuo), Tohoku University Hospital (Sendai), Ryukyu University Hospital (Nishihara), Kyoto Prefectural University Hospital (Kyoto), Kagoshima University Medical and Dental Hospital (Kagoshima), Kyoto University Hospital (Kyoto), Nagoya City University Hospital (Nagoya), Saga University Hospital (Saga), Kobe University Hospital (Kobe), Hiroshima University Hos-

pital (Hiroshima), and NTT West Kyushu Hospital (Kumamoto).

Eyes that presented with an IOP ≥ 22 mm Hg while on ocular hypotensive medications before surgery were included in this study. Steroid-induced glaucoma eyes were defined as open-angle eyes with an IOP elevation ≥ 22 mm Hg after the administration of corticosteroid. If both eyes underwent glaucoma surgeries, the eye that was treated first was investigated. Exclusion criteria were as follows: eyes with a history of previous glaucoma surgery, eyes that had undergone intraocular surgery up to 3 months before trabeculotomy or trabeculectomy, steroid-induced glaucoma eyes in the active phase of uveitis, eyes associated with IOP ≥ 22 mm Hg before corticosteroid administration in the medical records, and eyes that underwent combined glaucoma and cataract surgeries.

The technique of trabeculotomy performed in this study has been described previously.¹⁹ In brief, after conjunctival incision, a 4 × 4-mm square or triangular scleral flap at four-fifths thickness was created at the corneal limbus. After identification of the Schlemm's canal, its outer wall was cut with a razor blade and excised with fine scissors. U-shaped probes were then inserted into both ends of the opened canal and rotated 90 degrees against the trabecular

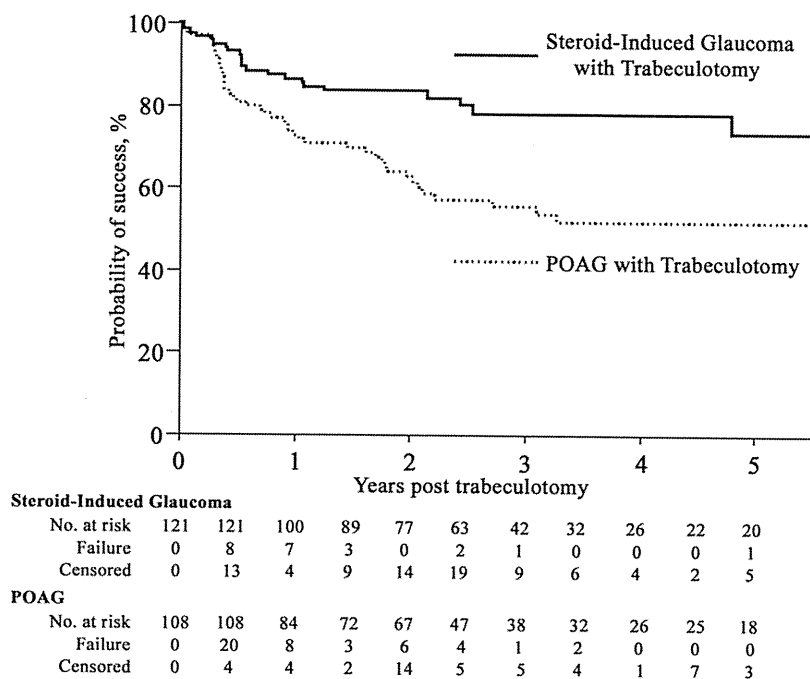


FIGURE 1. Criterion A–based Kaplan-Meier survival curves of surgical outcomes in patients with steroid-induced glaucoma (solid line) vs primary open-angle glaucoma (POAG; dotted line) that underwent trabeculotomy. The steroid-induced glaucoma eyes had a significantly higher cumulative probability of success than the POAG eyes ($P = .0008$).

meshwork. Rotation of these probes achieved 120-degree opening of the trabecular meshwork. The scleral flap was closed with 1 to 7 10-0 nylon sutures until the wound became watertight.

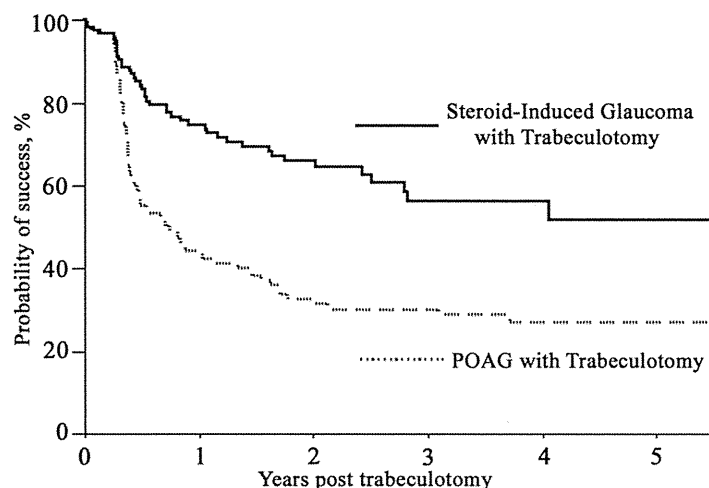
During trabeculotomy, some cases were combined with a sinusotomy, based upon the procedure of Mizoguchi and associates,²¹ which made 1 or 2 sites of 1-mm-diameter sclerotomy with a punch through the scleral flap before closure with 10-0 nylon sutures. Trabeculectomy was performed according to a modification of the technique developed by Cairns.²² Conjunctiva incisions included limbal-based and fornix-based procedures. After the creation of a scleral flap, sponges soaked with MMC (0.4 mg/mL) were applied to the posterior surface of the conjunctiva, Tenon's capsule, the adjacent episcleral tissue, and the scleral flap for 2 to 5 minutes, followed by irrigation with balanced salt solution. A trabecular block was excised to create a fistula in the anterior chamber, and peripheral iridectomy was then performed. The scleral flap was closed with 10-0 nylon sutures while the conjunctival flap was sutured with 10-0 nylon or 7-0 silk. All patients were required to sign informed consent forms before surgery.

• **MAIN OUTCOME MEASURE:** The main outcome measure was the probability of success in the Kaplan-Meier survival-curve analysis. Before data analysis, surgical failure was defined by the following IOP levels, with or without ocular hypotensive medications, which were verified at the next visit: criterion A, IOP ≥ 21 mm Hg; criterion B, IOP

≥ 18 mm Hg. IOP data that were examined using a Goldmann applanation tonometer were collected from patients' medical records. IOPs that corresponded to criteria A and B up to 3 months after surgery were not considered a surgical failure because of the occurrence of postoperative IOP fluctuations after trabeculotomy.¹⁹ If additional glaucoma surgery was performed, or visual acuity deteriorated to an absence of light perception, the eye was regarded as a surgical failure for both criteria.

We compared the surgical outcomes between the steroid-induced glaucoma with trabeculotomy group and the POAG with trabeculotomy group, and between the steroid-induced glaucoma with trabeculotomy group and the steroid-induced glaucoma with trabeculectomy group. To determine potential risk factors for surgical failure of steroid-induced glaucoma after trabeculotomy, the following variables were assessed: gender, age, pseudophakia, previous vitrectomy, route of corticosteroid administration (ocular instillation, intravitreal injection, posterior sub-Tenon's injection, or systemic administration), duration of corticosteroid administration after glaucoma surgery, reason for corticosteroid use (collagen disease, atopic dermatitis, or uveitis), sinusotomy, previous cataract surgery, and baseline IOP. These factors were analyzed statistically in the steroid-induced glaucoma with trabeculotomy group with criteria A and B. Data on postoperative complications were also collected from the medical records.

• **STATISTICAL ANALYSIS:** Data analysis was performed using the JMP version 8 statistical package program (SAS



Steroid-Induced Glaucoma											
No. at risk	121	121	89	76	64	48	33	22	16	13	10
Failure	0	19	9	5	3	2	3	0	0	1	0
Censored	0	13	4	7	13	13	8	6	3	2	2
POAG											
No. at risk	108	108	57	45	37	28	24	22	17	16	12
Failure	0	47	11	6	5	2	0	1	1	0	0
Censored	0	4	1	2	4	2	2	4	0	4	3

FIGURE 2. Criterion B–based Kaplan-Meier survival curves of surgical outcomes in patients with steroid-induced glaucoma (solid line) vs primary open-angle glaucoma (POAG; dotted line) that underwent trabeculectomy. The steroid-induced glaucoma eyes had a significantly higher cumulative probability of success than the POAG eyes ($P < .0001$).

TABLE 2. Cox Proportional Hazards Model Determining Likelihood of Surgical Outcomes for Patients With Steroid-Induced Glaucoma and Primary Open-Angle Glaucoma who Underwent Trabeculectomy

Variable	Criterion A			Criterion B		
	RR	95% CI	P Value	RR	95% CI	P Value
Steroid-induced glaucoma	0.409	0.223–0.735	.0027	0.451	0.286–0.706	.0005
Age (per year)	0.999	0.982–1.015	.8917	1.007	0.995–1.019	.2408
Preoperative IOP (per mm Hg)	1.004	0.977–1.029	.7557	0.993	0.972–1.013	.5115
Female	0.761	0.451–1.260	.2911	0.695	0.468–1.021	.0639
Previous cataract surgery	2.105	0.823–4.681	.1132	1.627	0.784–3.084	.1804
Combined sinusotomy	1.054	0.575–1.847	.8600	0.839	0.526–1.300	.4399

CI = confidence interval; IOP = intraocular pressure; RR = relative risk.

Institute, Cary, North Carolina, USA). Comparisons of the outcomes between the steroid-induced glaucoma with trabeculectomy group and the POAG with trabeculectomy group, as well as between the steroid-induced glaucoma with trabeculectomy group and the steroid-induced glaucoma with trabeculectomy group, were analyzed by the Kaplan-Meier survival curve and the log-rank test. To assess prognostic factors of steroid-induced glaucoma with trabeculectomy in univariate analysis, Kaplan-Meier survival-curve analysis and the log-rank test were used. To confirm the effects of prognostic factors and to identify the relative risk (RR) of surgical failure, multivariate prognostic factor analysis was performed with the Cox proportional hazards model. Multivariate factors were selected from variants with a probability (P) value of less than .15 shown by

univariate analysis. A P value less than .05 was considered statistically significant.

RESULTS

• **PATIENT CHARACTERISTICS:** In total, 163 patients (163 eyes) with steroid-induced glaucoma and 108 patients (108 eyes) with POAG satisfied the study criteria. All eligible patients were Japanese. Of the 163 eyes with steroid-induced glaucoma, 121 were included in the steroid-induced glaucoma with trabeculectomy group and 42 were included in the steroid-induced glaucoma with trabeculectomy group. Table 1 lists the characteristics of the enrolled patients.

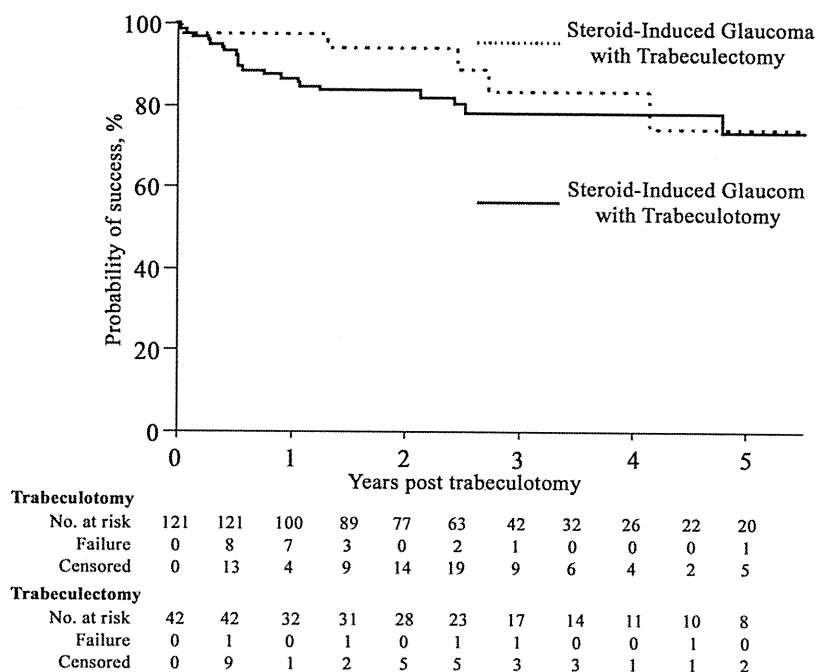


FIGURE 3. Criterion A-based Kaplan-Meier survival curves of surgical outcomes in eyes with trabeculotomy (solid line) vs trabeculectomy (dotted line) for steroid-induced glaucoma. There was no significant difference in the cumulative probability of success between the eyes with trabeculotomy and trabeculectomy ($P = .3636$).

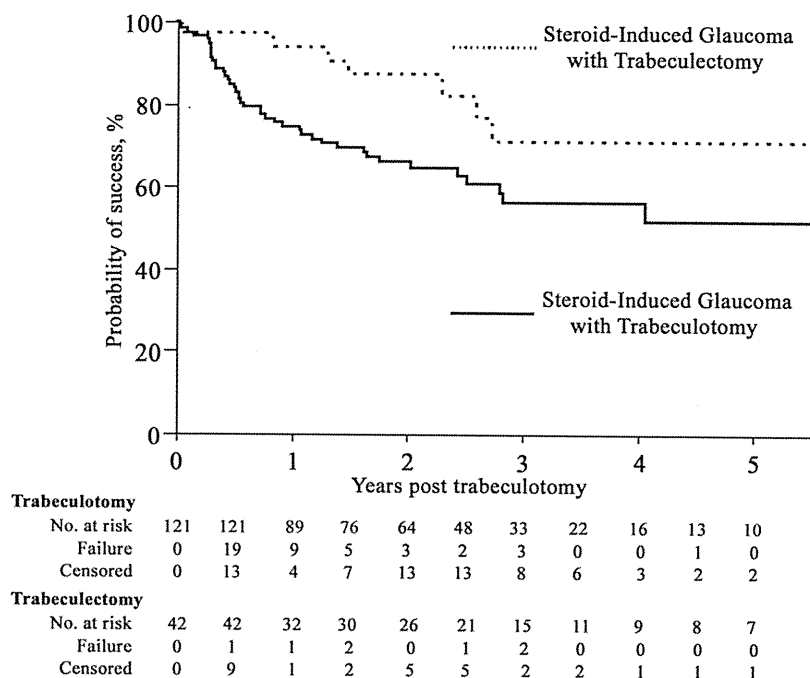


FIGURE 4. Criterion B-based Kaplan-Meier survival curves of surgical outcomes in eyes with trabeculotomy (solid line) vs trabeculectomy (dotted line) for steroid-induced glaucoma. The trabeculectomy group showed a significantly higher cumulative probability of success than the trabeculotomy group ($P = .0352$).

The steroid-induced glaucoma with trabeculotomy group was significantly younger ($P = .001$) and had a higher preoperative IOP ($P < .001$), a higher number of

female patients ($P = .014$), a higher number of previous cataract surgeries ($P = .029$), and a lower number of combined sinusotomies ($P = .012$) than the POAG with

TABLE 3. Influence of Prognostic Factors on Survival Time of Steroid-Induced Glaucoma Patients who Underwent Trabeculotomy

Variable	Number of Patients	Criterion A		Criterion B	
		80% Survival Time (Days)	P Value ^a	80% Survival Time (Days)	P Value ^a
Gender			.1779		.0422
Female	62	>3850		305	
Male	59	385		146	
Age (years)			.9040		.2458
<30	51	852		206	
≥30	70	916		181	
Preoperative IOP (mm Hg)			.7443		.4387
<40	79	181		181	
≥40	42	840		200	
Diabetes mellitus			.1936		.0848
Yes	13	146		96	
No	108	816		291	
Hypertension			.9394		.9915
Yes	18	>2148		291	
No	103	852		195	
Combined sinusotomy			.2416		.9270
Yes	20	146		120	
No	101	916		260	
Previous cataract surgery			.0055		.1829
Yes	17	49		49	
No	104	1732		272	
Previous vitrectomy			<0.0001		.0050
Yes	6	10		10	
No	115	1742		260	
Cause of corticosteroid use					
Collagen disease			.3397		.9248
Yes	37	750		195	
No	84	1732		200	
Atopic dermatitis			.9929		.2449
Yes	21	278		162	
No	100	852		195	
Uveitis			.4674		.7942
Yes	25	>2519		177	
No	96	840		200	
Route of steroid administration					
Ocular instillation only			.6968		.1204
Yes	17	1732		1286	
No	104	852		181	
Posterior sub-Tenon's injection of TA			.9546		.3239
Yes	13	>1339		>1339	
No	108	916		200	
Intravitreal injection of TA			.1843		.4379
Yes	10	49		49	
No	111	916		206	
Oral administration			.7412		.6920
Yes	72	840		181	
No	49	1732		272	
Intravenous administration			.4050		.1883
Yes	3	>1580		>1580	
No	118	852		195	

Continued on next page

TABLE 3. Influence of Prognostic Factors on Survival Time of Steroid-Induced Glaucoma Patients who Underwent Trabeculotomy (*Continued*)

Variable	Number of Patients	Criterion A		Criterion B	
		80% Survival Time (Days)	P Value ^a	80% Survival Time (Days)	P Value ^a
Postoperative corticosteroid administration			.1987		.7335
>3 months	68	<3850		195	
≤3 months	53	372		200	

IOP = intraocular pressure; TA = triamcinolone acetonide.
^aThe P values are based on the log-rank test.

trabeculotomy group. However, there were no significant differences in patient characteristics between the steroid-induced glaucoma with trabeculotomy group and the steroid-induced glaucoma with trabeculectomy group.

• **STEROID-INDUCED GLAUCOMA VS POAG:** The mean follow-up periods were 38.4 ± 28.7 months in the steroid-induced glaucoma with trabeculotomy group and 49.8 ± 37.2 months in the POAG with trabeculotomy group ($P = 0.010$). Kaplan-Meier survival-curve analyses of the steroid-induced glaucoma with trabeculotomy group and the POAG with trabeculotomy group for criteria A and B are presented in Figures 1 and 2, respectively. The steroid-induced glaucoma with trabeculotomy group had a significantly higher cumulative probability of success for criteria A ($P = .0008$) and B ($P < .0001$). For criterion A, the probabilities of success 1, 2, 3, and 5 years after trabeculotomy in the steroid-induced glaucoma with trabeculotomy group and the POAG with trabeculotomy group were as follows: 86.5% vs 73.2%, 83.5% vs 63.0%, 78.1% vs 55.8%, and 73.5% vs 52.2%, respectively. For criterion B, the probabilities of success 1, 2, 3, and 5 years after trabeculotomy in the steroid-induced glaucoma with trabeculotomy group and the POAG with trabeculotomy group were as follows: 74.6% vs 44.7%, 66.1% vs 33.0%, 56.4% vs 30.6%, and 51.7% vs 27.5%, respectively. The number of eyes classified as surgical failures in the steroid-induced glaucoma with trabeculotomy group and the POAG with trabeculotomy group were 22/121 (18.2%) vs 45/108 (41.7%) for criterion A and 42/121 (34.7%) vs 73/108 (67.6%) for criterion B, respectively.

Since there were significant differences between the preoperative data of the steroid-induced glaucoma with trabeculotomy group and the POAG with trabeculotomy group, a Cox proportional hazards model including age, preoperative IOP, gender, previous cataract surgery, and combined sinusotomy was performed (Table 2). The multivariate model suggested that trabeculotomy in steroid-induced glaucoma eyes was independently associated with a better prognosis when compared with the same procedure in POAG eyes, even after adjusting for confounding

factors (criterion A, RR = 0.409, $P = .0027$; criterion B, RR = 0.451, $P = .0005$).

• **TRABECULOTOMY VS TRABECULECTOMY:** The mean follow-up period in the steroid-induced glaucoma with trabeculectomy group was 37.1 ± 31.8 months (38.4 ± 28.7 months in the steroid-induced glaucoma with trabeculotomy group, $P = .808$). The Kaplan-Meier survival-curve analysis between the steroid-induced glaucoma with trabeculotomy group and the steroid-induced glaucoma with trabeculectomy group for criteria A and B are presented in Figures 3 and 4, respectively. No significant difference was found between the 2 groups for criterion A ($P = .3636$). The probabilities of success 1, 2, 3, and 5 years after surgery in the steroid-induced glaucoma with trabeculectomy group for criterion A were 97.6%, 94.3%, 83.8%, and 74.5%, respectively. The number of eyes classified as surgical failures in the steroid-induced glaucoma with trabeculectomy group was 5 (11.9%) for criterion A. However, the steroid-induced glaucoma with trabeculectomy group showed a significantly higher cumulative probability of success for criterion B ($P = .0352$). The probabilities of success 1, 2, 3, and 5 years after surgery in the steroid-induced glaucoma with trabeculectomy group for criterion B were 94.5%, 87.7%, 71.6%, and 71.6%, respectively. The number of eyes classified as surgical failures in the steroid-induced glaucoma with trabeculectomy group was 7 (16.7%) for criterion B.

• **PROGNOSTIC FACTORS FOR FAILURE OF TRABECULOTOMY FOR STEROID-INDUCED GLAUCOMA EYES:** The potential prognostic factors influencing survival time are listed in Table 3. Univariate analysis showed previous cataract surgery ($P = .0055$) and previous vitrectomy ($P < .0001$) to be significant prognostic factors for criterion A, and male gender ($P = .0422$) and previous vitrectomy ($P = .0050$) for criterion B. Diabetes mellitus ($P = .0848$) and ocular instillation of corticosteroid ($P = .1204$) were the factors with a P value of less than .15 for criterion B. The Cox proportional hazards model including these variables revealed that prognostic factors for surgical failure were

TABLE 4. Cox Proportional Hazards Model on Criteria A and B, Determining Likelihood of Surgical Outcomes for All 121 Patients With Steroid-Induced Glaucoma who Underwent Trabeculotomy

Variable	Criterion A			Criterion B		
	RR	95% CI	P Value	RR	95% CI	P Value
Previous cataract surgery	1.614	0.255–5.688	.5488	—	—	—
Previous vitrectomy	5.340	1.037–38.655	.0452	3.898	1.108–10.688	.0360
Male	—	—	—	1.783	0.938–3.414	.0774
Diabetes mellitus	—	—	—	1.871	0.754–4.018	.1632
Corticosteroid administration other than ocular instillation	—	—	—	2.752	1.065–9.426	.0352

CI = confidence interval; IOP = intraocular pressure; RR = relative risk.

previous vitrectomy (RR = 5.340, $P = .0452$ for criterion A; RR = 3.898, $P = .0360$ for criterion B) and corticosteroid administration other than ocular instillation (RR = 2.752, $P = .0352$ for criterion B) (Table 4).

• **POSTOPERATIVE COMPLICATIONS:** In the steroid-induced glaucoma with trabeculectomy group, choroidal detachment occurred in 2 eyes (4.8%), flat anterior chamber requiring anterior chamber reformation occurred in 1 eye (2.4%), and hypotony maculopathy occurred in 7 eyes (16.7%). None of these complications occurred in either the steroid-induced glaucoma with trabeculotomy group or the POAG with trabeculotomy group. The progression of postoperative cataracts was observed in 9 eyes (7.4%) of the steroid-induced glaucoma with trabeculotomy group, 1 eye (0.9%) of the POAG with trabeculotomy group, and 4 eyes (9.5%) of the steroid-induced glaucoma with trabeculectomy group. No eyes encountered postoperative infectious blebitis or endophthalmitis.

DISCUSSION

THIS STUDY COMPARED THE SUCCESS RATES OF TRABECULOTOMY for steroid-induced glaucomatous eyes with those for trabeculectomy for steroid-induced glaucoma eyes, and for trabeculotomy for POAG eyes. Trabeculotomy showed a significantly higher cumulative probability of success in steroid-induced glaucoma patients than POAG patients for both criterion A ($P = .0008$) and criterion B ($P < .0001$). The probability of success in steroid-induced glaucoma eyes treated with trabeculotomy was comparable to that in steroid-induced glaucoma eyes treated with MMC trabeculectomy for criterion A ($P = .3636$), but was significantly lower for criterion B ($P = .0352$). Significant prognostic factors for surgical failure of trabeculotomy in steroid-induced glaucoma patients were previous vitrectomy for criteria A (RR = 5.340, $P = .0452$) and B (RR = 3.898, $P = .0360$), and corticosteroid treatment other than ocular instillation for criterion B (RR = 2.752, $P = .0352$).

Several studies have demonstrated surgical results for steroid-induced glaucoma patients.^{10,11,13–15,17,23–27} For example, Sihota and associates¹⁰ reported that 9 eyes with steroid-induced glaucoma that required trabeculectomy with MMC showed normal IOP levels after surgery. Several reports^{11,23–27} demonstrated that filtering surgery was successful for IOP management in steroid-induced glaucomatous eyes after intravitreal injection of triamcinolone acetonide. Krishnan and associates¹³ found that all of 3 eyes with triamcinolone-induced IOP elevation were successfully treated with viscocanalostomy. For laser trabeculectomy, Ricci and associates¹⁴ and Viola and associates¹⁵ reported that argon laser trabeculectomy was effective in all cases in their studies, and selective laser trabeculectomy was shown to lower IOP in 5 of 7 eyes.¹⁷ However, these reports on surgical treatment for steroid-induced glaucoma included only a small number of cases and lacked control groups and details of long-term prognosis. Our previous study lacked control groups but showed that trabeculotomy reduced IOPs to 21 mm Hg or less in 14 eyes with steroid-induced glaucoma.¹² To our knowledge, our present multicenter study reports on the largest number of steroid-induced glaucoma patients.

Trabeculotomy showed a better prognosis in steroid-induced glaucoma eyes than POAG eyes in the present study. The accumulation of extracellular matrices in trabecular meshwork has been believed to cause increased outflow resistance of the aqueous humor in steroid-induced glaucoma patients. This is because histochemical data demonstrate abnormally accumulated extracellular matrices such as type IV collagen, heparin sulfate, proteoglycan, and fibronectin in the trabecular meshwork of steroid-induced glaucoma patients.⁹ Because the main target of trabeculotomy for IOP reduction is the relief of outflow resistance in the trabecular meshwork, the consistency between the surgical target and the pathologic lesion might explain the effectiveness of surgery for steroid-induced glaucoma eyes.

The surgical success of trabeculotomy for steroid-induced glaucoma was comparable to the success of MMC trabeculectomy for steroid-induced glaucoma for criterion

A. Trabeculectomy has a potential risk of late-onset infection of the filtering bleb.²⁸⁻³¹ A previous multicenter case-control study suggested that the use of systemic corticosteroid and juvenile-onset glaucoma should be included among the risk factors for late-onset infection after filtering surgery.³² Moreover, younger patients are more susceptible to steroid-induced IOP elevation.^{4,33,34} Trabeculectomy might be more beneficial for younger patients with steroid-induced glaucoma, from the viewpoint of late-onset infection, than trabeculectomy because of the nonfiltering surgery.

Trabeculectomy had a significantly higher probability of success than trabeculectomy using criterion B in the present study. Thus, many steroid-induced glaucoma eyes treated with trabeculectomy had postoperative IOPs of 18 to 20 mm Hg, which might be too high to prevent progressive visual field changes for glaucomatous eyes with advanced progressive visual field defects. The Advanced Glaucoma Intervention Study³⁵ found that visual field loss in eyes with advanced open-angle glaucoma progresses further if postoperative IOP of 18 mm Hg or higher is more frequent. These findings imply that trabeculectomy rather than trabeculectomy is more favorable for controlling IOP in steroid-induced glaucoma eyes with advanced visual field loss.

Even when other types of glaucoma are included, the prognostic factors for the surgical failure of trabeculectomy have not been sufficiently identified. Our previous report indicated that higher preoperative IOP results in poorer prognosis in eyes with POAG or exfoliative glaucoma.¹⁹ Higher preoperative IOP might reflect the severity of glaucoma, resulting in a poorer response to reduce IOP. The present study showed that higher preoperative IOP was not a prognostic factor for surgical failure of trabeculectomy for steroid-induced glaucoma, while corticosteroid administration other than ocular instillation was shown to be a prognostic factor in criterion B. These data might

reflect the fact that the severity of steroid-induced glaucoma depends on the route of corticosteroid administration rather than the preoperative IOP levels. In addition, previous vitrectomy was a prognostic factor for surgical failure for both criteria. Although we have no conclusive explanation, it is conceivable that vitrectomy causes the elevation of inflammatory factors or growth factors in the aqueous humor. Vitrectomized eyes might lead to recurrent fibrosis in the outflow pathway of the trabecular meshwork that was created by the trabeculectomy.

This study had some limitations caused by the retrospective design. First, the selection bias of the type of surgery performed for steroid-induced glaucoma eyes might have affected the surgical result. In fact, all 6 vitrectomized eyes were treated with trabeculectomy. Because of conjunctival scarring after vitrectomy, trabeculectomy rather than trabeculectomy might have been the surgery of choice for vitrectomized eyes. Second, as higher IOP is a prognostic factor for failure of trabeculectomy for POAG eyes,¹⁹ POAG patients with higher IOP might have been treated with trabeculectomy rather than trabeculectomy; this might have increased the cumulative probability of success in POAG with trabeculectomy.

In conclusion, this study demonstrates that trabeculectomy might be more effective for steroid-induced glaucoma eyes than POAG eyes. Moreover, the surgical success in steroid-induced glaucoma eyes is comparable to the outcome of trabeculectomy unless more substantial IOP reduction is necessary, in which case trabeculectomy would be a better option. IOP reduction of steroid-induced glaucoma patients with previous vitrectomy or with corticosteroid administration other than ocular instillation might be more resistant to trabeculectomy. Trabeculectomy should be considered as an option for the surgical management of steroid-induced glaucoma, although future prospective studies are necessary to validate our findings.

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Biosketch

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D-Allose as ischemic retina injury inhibitor during rabbit vitrectomy

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Abstract

Purpose To investigate the protective effects of D-allose, a rare sugar, on pressure-induced ischemia during vitrectomy in the rabbit eye.

Methods The rabbits underwent pars plana vitrectomy, and continuous intraocular irrigation at a perfusion pressure of 140 mmHg was performed for 45 min. Intraocular pressure was regulated by adjusting the height of a bottle of balanced saline solution containing D-allose. Morphometric studies were performed to study the effects of D-allose on the histological changes induced by ischemia in the rabbit retina. Electroretinograms (ERGs) were taken before and 1 and 7 days after vitrectomy. Nitroblue tetrazolium was used as an index of superoxide anion (O_2^-) generation. Data were analyzed by use of the unpaired Student's *t* test. **Results** Seven days after ischemia, significant reductions in both number of ganglion cells and the thickness of the inner plexiform layer were observed. D-Allose significantly inhibited ischemic injury of the inner retina ($P < 0.05$). On postoperative day 7, amplitudes of ERG b-waves were significantly lower in the control group than in the D-allose

group ($P < 0.05$). D-Allose suppressed the production of O_2^- .

Conclusions Intraocular irrigation with D-allose during vitrectomy may protect the retina against ischemia-induced damage.

Keywords D-Allose · Neuroprotection · Retinal ischemia · Superoxide anion · Vitrectomy

Introduction

Rare sugars are monosaccharides that exist in nature in limited quantities only. Whereas naturally abundant monosaccharides such as D-glucose and D-fructose are few in number, more than 50 kinds of rare sugars have been identified. D-Allose is a rare aldo-hexose sugar produced from D-psicose by enzymes from microorganisms [1]. Murata et al. [2] examined the scavenging activity of D-allose using electron spin resonance. They showed that D-allose inhibits the production of reactive oxygen species (ROS) in a dose-dependent manner. Furthermore, we recently reported that D-allose might protect neurons against retinal ischemia–reperfusion injury by reducing extracellular glutamate and attenuating oxidative stress [3].

Improvements in the techniques of pars plana vitrectomy (PPV) have resulted in better prognosis for vision-threatening eye diseases such as macular hole, epiretinal membrane, retinal detachment, and proliferative diabetic retinopathy (PDR). However, optic nerve atrophy after successful PPV for diabetic retinopathy is one of the most serious complications and can lead to blindness [4]. Because the extent of autoregulation in the retina and optic nerve head is limited, high infusion pressures during

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vitreous surgery may cause retinal ischemia and lead to damage of the retina and optic nerve.

During or after ischemia, ROS such as hydrogen peroxide (H_2O_2), superoxide anion (O_2^-) and hydroxyl radical (OH^-) can be produced in large quantities and act as cytotoxic metabolites. ROS have been implicated in the process of apoptosis, because treatment of cells with ROS can result in this form of cell death, whereas application of antioxidants can prevent it under some conditions [5, 6]. ROS scavengers have been shown to be neuroprotective against ischemia in acute experimental models [7].

The purpose of this study was to investigate the protective effects of D-allose on pressure-induced ischemia during vitrectomy in the rabbit eye.

Materials and methods

Animals

Female New Zealand White rabbits, weighing 2.0–2.4 kg, were obtained from Kitayama Labs KK (Nagano, Japan). The rabbits were anesthetized by intraperitoneal injection of ketamine (50 mg/kg) and xylazine (10 mg/kg). Animal care and experiments followed the standard guidelines for animal experimentation of Kagawa University Faculty of Medicine and adhered to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research. The animals were divided into two groups according to the intraocular irrigating solutions they received: the vehicle group, using a balanced saline solution (BSS) alone, and the D-allose group, using BSS + D-allose (2%).

Vitrectomy and subsequent vitreous perfusion

A 250-ml bottle of intraocular irrigating solution (BSS; Alcon Laboratories, Fort Worth, TX, USA) was suspended approximately 100 cm above the eye level of each rabbit, and connected to a 20-gauge infusion cannula through a 200-cm-long tube. The infusion cannula was inserted through the sclerotomy in the inferonasal quadrant 3 mm posterior to the limbus, and sutured in place. A vitreous cutter (MVS XX; Alcon) was placed through the sclerotomy in the supratemporal quadrant, and vitrectomy was then performed for 10 min. The vitreous cavity was irrigated during vitrectomy either with BSS alone or with BSS + D-allose. Adequate illumination was provided by a paraxial light operation microscope without intraocular fiberoptic illumination. If retinal tear or retinal detachment occurred during vitrectomy, those animals were excluded from the experiment. After completion of the vitrectomy, the cutter was

removed and the sclerotomy wound was tightly sutured using a 7-0 vicryl. Intraocular pressure (IOP) was raised to 140 mmHg for 45 min by elevating the solution bottle. Retinal ischemia was confirmed by whitening of the fundus. Rectal temperature was monitored throughout the surgery. Because body temperature during vitrectomy in acute ischemic eyes may affect ischemia-induced retinal damage [8, 9], rectal and tympanic temperature was maintained at approximately 37°C by use of a feedback-controlled heating pad (BRC, Nagoya, Japan) during the operation. After restoration of blood flow, the temperature was continuously maintained at 37°C. Eyes were treated topically with levofloxacin (Santen Pharmaceutical, Osaka, Japan) after surgery to prevent postoperative infection.

Electroretinograms (ERGs)

Before surgery, ERG responses were measured after 20 min of dark adaptation using a recording device (Mayo, Aichi, Japan). The procedure was repeated on postoperative days 1 and 7. ERGs were recorded by positioning the rabbit in a box and placing a xenon lamp 15 cm in front of the eye. A flash of maximum intensity 20 J provided photostimulation. A contact lens electrode was placed on the cornea, and a reference electrode subcutaneously at the vertex. ERGs were taken of both eyes for each animal. The a- and b-wave amplitudes of each operated eye were shown as a percentage of those in the pre-operated eye.

Histological examination

For histological examination, the rabbits were anesthetized by intraperitoneal injection of ketamine and xylazine 1 week after ischemia and perfused intracardially with phosphate-buffered saline (PBS), followed by perfusion with 4% paraformaldehyde in PBS. Eyes were removed and embedded in paraffin, and thin sections (5- μm thickness) were cut using a microtome. Each eye was mounted on a silane-coated glass slide and then stained with hematoxylin and eosin (HE).

Morphometric analysis was performed to quantify ischemic injury. The sections for analysis were selected randomly for each eye. A microscopic image of each section within 0.5–1 mm of the optic disc was scanned. In each computer image, the thickness of the inner plexiform layer (IPL) and inner nuclear layer (INL) at the tire frame were measured. Finally, in each eye, the thicknesses of the IPL and INL were obtained as the mean values of five measurements. For each animal, these values for the right eye were normalized to those for the intact left eye and shown as a percentage.

Retrograde labeling of retinal ganglion cells (RGCs)

Seven days before the rabbits were killed, the optic nerves were exposed by lateral orbitotomy. Using a Hamilton syringe, 10 μ l of a 0.1% fluorescent dye (Fast Blue; Polysciences, Warrington, PA, USA) was injected into the optic nerve 2 mm behind the eyeball. Care was taken not to injure any blood vessels, especially the ophthalmic artery that enters the sclera from the ventral margin of the optic nerve. After the surgery the eyes were treated topically with levofloxacin.

Assessment of RGC survival

Animals were killed by an overdose of ketamine and xylazine 1 week after the fast blue application. Whole, flat-mounted retinas were then assayed for the retinal ganglion cell density. The rabbit eyes were enucleated and fixed in 4% paraformaldehyde for 10 h at room temperature. After removal of the anterior segments, the resulting posterior eyecups were left in place. Subsequently, 4 radial cuts were made in the periphery of each eyecup, with the retina then carefully separated from the retinal pigment epithelium. To prepare the flat mounts, the retina was dissociated from the underlying structures, flattened by making 4 radial cuts, and then spread on a gelatin-coated glass slide. Labeled RGCs were visualized under a fluorescence microscope (Olympus BX-51/DP70; Olympus, Tokyo, Japan) by using a filter set (excitation filter 330–385 nm; barrier filter 420 nm; WU; Olympus). Fluorescence-labeled RGCs were counted in 12 microscopic fields of retinal tissue from 2 regions in each quadrant at 2 different eccentricities, central and peripheral. We counted the RGCs in each eye by using Image-Pro Plus software (Version 4.0; Media Cybernetics, The Imaging Expert, Bethesda, MD, USA). Cell counts were conducted by the same person in a

masked fashion, with the identity of the original retinas unknown to the investigator until all cell counts from all the different groups were completed. Changes in the densities of the RGCs were expressed as the RGC survival percentage, which was based on comparison of the surgical and contralateral control eyes. The specimens that were compared came from different retinal regions of the same animal.

O₂⁻ analysis

O₂⁻ generation was determined by measuring the reduction of nitroblue tetrazolium (NBT) to a diformazan precipitate as previously described [10]. NBT (50 mg/ml; Research Organics, Cleveland, OH, USA) was dissolved in dimethylformamide (DMF) and distilled BSS. The final DMF concentration was 10%. Reduction was detected by intravitreal injection of 5- μ l NBT solution in the period after vitrectomy before ischemia. For observation of the diformazan precipitate, a microscope (S5; Carl Zeiss, München, Germany) was used with a 3CCD digital camera (MKC-507; Ikegami Tsushinki, Tokyo, Japan) and recorded using a DVD recorder.

Statistical analysis

Image analysis was performed with Image-Pro Plus software (The Imaging Expert) to assess O₂⁻ generation area density. We determined the total blue-stained area indicated O₂⁻ generation area. Evaluation by Image-Pro Plus analysis of photographs was performed blind by two researchers not directly involved in this study and the generation area was automatically produced by the software. All statistical values are presented as mean \pm standard deviation (SD). Data were analyzed by use of the unpaired Student's *t* test. Values of *p* < 0.05 were considered statistically significant.

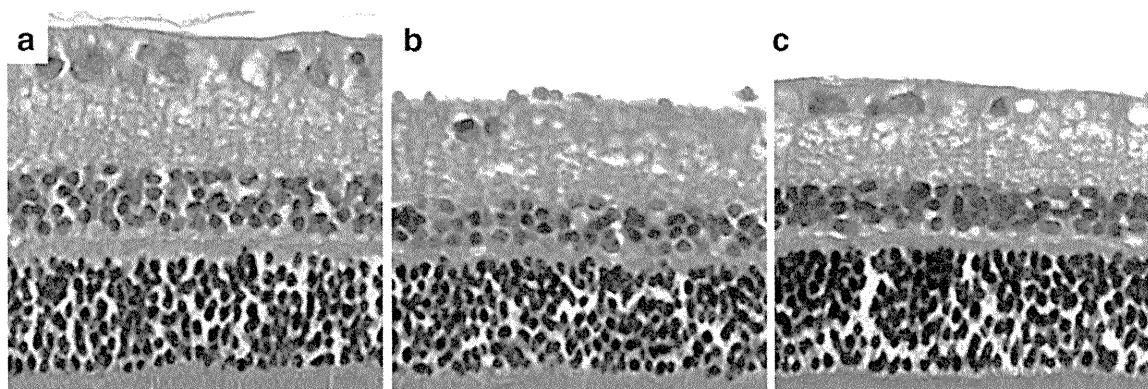


Fig. 1 Light micrographs of a cross-section of **a** hematoxylin and eosin histology through normal rabbit retina and **b** 7 days after ischemia with balanced salt solution (BSS) alone, or **c** BSS +

D-allose. Cell loss in the ganglion cell layer (GCL) and reduced inner nuclear layer (INL) thickness were ameliorated in the D-allose group. Bar 10 μ m

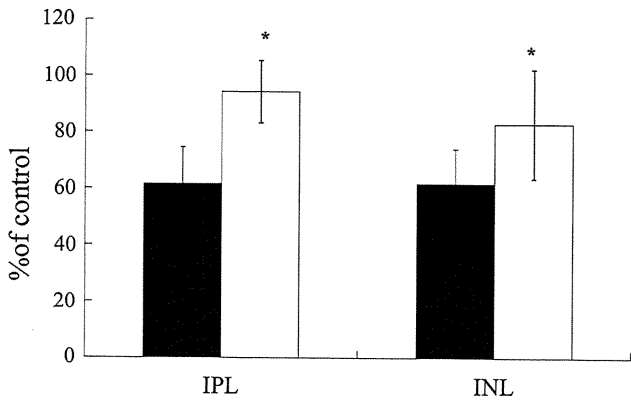


Fig. 2 Percentage changes relative to control values in the thicknesses of the IPL and INL 7 days after ischemia with BSS alone and BSS + D-allose. *Black bar* D-allose group, *white bar* vehicle group. Administration of 2% D-allose significantly prevented the reduction in the number of cells in the GCL and the thickness of the INL. Results are expressed as mean ± standard deviation (SD). *P < 0.05

Results

Histological change in the retina after ischemia with and without D-allose

Figure 1a shows a normal retina. Light-microscopic photographs were taken 7 days after ischemia (Fig. 1b, c). The retina of the untreated eye in the animals, was used as control. In the vehicle group, significant reductions in the thickness of the INL were observed. The thickness of the IPL was 61.6 ± 13.0% that of the control and the thickness of the INL was reduced to 61.8 ± 12.2% that of the control (n = 6; Fig. 2). In the D-allose group, the thickness of the IPL was 94.2 ± 11.1% that of the control and the thickness of the INL was 82.9 ± 19.4% that of the control (n = 5; Fig. 2). Reduced INL thickness was ameliorated in the D-allose group.

Fig. 3 Effects of D-allose on ischemia-induced retinal ganglion cell (RGC) death. **a** Retrograde labeling of RGCs in nonischemic eyes, and **b** 7 days after ischemic injury treated with vehicle or D-allose. *Bar* 100 μm. RGCs were counted in the central and peripheral areas. *Gray bar* D-allose group, *black bar* vehicle group. Graph depicts the mean ± SD for three rabbits treated with vehicle and three rabbits treated with D-allose. *P < 0.05

