Angiogenesis is the major feature in the pathogenesis of proliferative diabetic retinopathy (PDR). In this condition, retinal neovascularisation has a catastrophic effect on vision by causing vitreous haemorrhage, retinal detachment with formation of a fibrovascular membrane, and eventual blindness. The factors that stimulate the growth of retinal blood vessels have not been fully defined, but circumstantial evidence indicates that this not only involves angiogenic cytokines such as vascular endothelial growth factor (VEGF) but also vasoactive hormones such as angiotensin II. Angiotensin II has a proliferative effect and has been reported to regulate the growth of vascular smooth muscle cells and to stimulate the induction of various growth factors. Recent studies have suggested that abnormalities of the renin-angiotensin system (RAS) may also play a part in the progression of diabetic retinopathy. Inhibition of angiotensin converting enzyme (ACE) has been reported to be associated with a reduction of PDR, suggesting that suppression of the RAS may be of value for preventing and treating retinal neovascularisation. The hypothesis that an ocular RAS is involved in the development of PDR is supported by evidence that all components of the RAS are present in the retina and that angiotensin II, the effector molecule of this system, has angiogenic activity. These findings prompted us to examine whether angiotensin II plays a part in the development of PDR in combination with VEGF, which is considered to be the most potent factor in promoting angiogenesis. Therefore, we investigated the relation between the levels of angiotensin II and VEGF in the vitreous fluid of diabetic patients as well as the correlation between these factors and the severity of PDR. The present study revealed that angiotensin II and VEGF levels in the vitreous fluid were correlated with the severity of PDR and that the vitreous levels of these two molecules were also correlated with each other. Furthermore, both angiotensin II and VEGF were elevated in the active stage of PDR. Angiotensin II may induce neovascularisation via a paracrine effect on VEGF in diabetic patients with PDR.

**MATERIALS AND METHODS**

**Patients**

Undiluted vitreous fluid samples were harvested at the start of vitrectomy after informed consent was obtained from each subject following an explanation of the purpose and potential adverse effects of the procedure. This study was performed in accordance with the 1975 Declaration of Helsinki, as revised in 1983. Vitreous fluid samples were obtained from 51 patients with PDR, six diabetic patients without diabetic retinopathy, and 16 non-diabetic patients with ocular disease. Vitrectomy was performed on the 51 patients with PDR for the following reason: 27 had vitreous and/or preretinal haemorrhage, 17 had retinal detachment, and seven had macular heterotropia with proliferative tissues. The cases with macular heterotropia hoped to undergo the surgery because of the disturbed vision. The six diabetic patients without diabetic retinopathy included four with macular hole and two with epiretinal membrane, while the 16 non-diabetic patients included 12 with macular hole and four with epiretinal membrane (none of these 16 patients had proliferative vitreoretinopathy). Exclusion criteria for this study were: (1) treatment with an ACE inhibitor or an angiotensin II receptor antagonist (ARA),
(2) previous ocular surgery, and (3) a history of ocular inflammation. All patients with PDR underwent retinal photocoagulation (mean 922 shots; range 342–1682 shots) before vitreous surgery. Vitrectomy was performed at the Tokyo Women’s Medical University.

**Fundus findings**

Preoperative and operative fundus findings were recorded for each subject. The severity of diabetic retinopathy was assessed by standardised fundus colour photography and fluorescein angiography (FAG), which were performed with a Topcon TRC-50HA fundus camera, an image-net system (Tokyo Optical Co Ltd, Japan), and a preset lens with a slit lamp. Diabetic retinopathy was graded according to the modified Early Treatment Diabetic Retinopathy Study (ETDRS) retinopathy severity scale. 21–23

PDR was classified as active (30 eyes) if there were new preretinal capillaries and as quiescent (21 eyes) if the vasoproliferation only consisted of large vessels within the membrane at the time of surgery. 21–22

**Sample collection**

Samples of vitreous fluid were collected into sterile tubes and were rapidly frozen at −80°C. Those samples were obtained at the time of vitreoretinal surgery, with the protocol for sample collection being approved by the institutional review board and with all patients giving informed consent.

Plasma samples were also collected from the 51 patients with PDR. Blood was immediately placed on ice and subjected to centrifugation at 3000 g for 5 minutes at 4°C, after which the plasma was rapidly frozen at −80°C until assay. The institutional review board also approved the protocol for blood collection.

**Measurement of VEGF, angiotensin II, and ACE levels**

Both VEGF and angiotensin II were measured in vitreous samples from the same eye as well as in the plasma. The VEGF concentration was measured by an enzyme linked immunosorbent assay (ELISA) for human VEGF (R&D Systems, Minneapolis, MN, USA). 24 This VEGF kit was able to detect two of the four VEGF isoforms (VEGF121 and VEGF165), probably because these two shorter VEGF isoforms are secreted and the two longer isoforms are cell associated. The assay was performed according to the manufacturer’s instructions. A standard solution (100 µl) or sample (10 or 100 µl) was added to the wells of a 96 well plate coated with an immobilised monoclonal antibody. After incubation, the plate was washed and the enzyme labelled antibody was added. After further incubation, the plate was washed again and the substrate was added. The reaction was stopped after colour had developed by adding the stop solution, and the optical density was determined at 450 and 620 nm using an absorption spectrophotometer (Titertek Multiscan MCC/340; ICN, Tokyo, Japan). A standard curve was plotted from the measurements made with the standard solution (from 15.6 to 1000 pg/ml), and the concentration of VEGF in each sample was determined from this curve. The VEGF levels in vitreous fluid and plasma were within the detection range of the assay, since the minimum detectable concentration was 15.6 pg/ml (coefficient of variation (CV) intra-assay 3.5%, CV interassay 5.8%).

Vitreous and plasma angiotensin II levels were determined by radioimmunoassay (RIA) 25 and serum ACE levels were determined by the method of Kasahara. 26 For the measurement of angiotensin II, 0.5 ml of vitreous fluid or plasma was mixed with 2.5 ml of ethanol. After centrifugation at 2000 g for 15 minutes at 4°C, the supernatant was dried under nitrogen gas at 37°C. The dried samples were reconstituted with 0.5 ml of assay buffer, and assay was performed with an angiotensin II RIA kit (Nichols, CA, USA). Measurement of ACE activity was performed with an ACE colour kit (Fujirebio, Tokyo, Japan). Serum (0.05 ml) was mixed with 0.5 ml of p-hydroxybenzoyl-glycyl-L-histidyl-L-leucine. After incubation, the reaction was stopped when colour had developed by adding the stop solution and the optical density was determined at 505 nm using an absorption spectrophotometer (AU600; Olympus, Tokyo, Japan). The angiotensin II and ACE levels in vitreous fluid and plasma were within the detection range of these assays, since the minimum detectable concentration was 4.0 pg/ml (CV intra-assay 3.8%, CV interassay 6.0%) and 2.5 pg/ml (CV intra-assay 4.0%, CV interassay 6.2%), respectively.

**Statistical analysis**

All analyses were performed with SAS System 6.12 software (SAS Institute Inc, Cary, NC, USA). 27 Data are presented as the frequency or mean (SD). Data with a skewed distribution were transformed to a logarithmic scale, and the geometric mean was calculated together with 1 SD below and 1 SD above the mean on that scale. Analysis of variance (ANOVA) was used to test for statistically significant differences among the groups and the Turkey-Kramer multiple comparison test was also applied when appropriate. Correlations were tested using Spearman’s rank correlation coefficients. A two tailed p value of less than 0.05 was considered to indicate statistical significance.

**RESULTS**

**Vitreous levels of VEGF and angiotensin II**

The diabetic patients included 31 men and 26 women mean age 60.7 (SD 10.5) years, with a diabetes duration of 16.2 (6.3) years. The non-diabetic patients included 15 men and 11 women, mean age 60.4 (13.4) years. The NDM patients had significantly lower VEGF and angiotensin II levels than the NDR and PDR patients (NDM vs NDR, p<0.0001; NDM vs PDR, p<0.0001).

![Figure 1](http://example.com/figure1.png)

**Figure 1**  [A] VEGF concentrations in the vitreous fluid of non-diabetic patients (NDM), diabetic patients without retinopathy (NDR), and PDR patients (PDR) (*p<0.0001). [B] Angiotensin II concentrations in the vitreous fluid of non-diabetic patients (NDM), diabetic patients without retinopathy (NDR), and PDR patients (PDR) (*p<0.0001).
years and an HbA₁c level of 7.6% (2.3%). The 16 patients with non-diabetic ocular disease included eight men and eight women aged 63.4 (6.5) years. There was no significant difference in age between the diabetic and non-diabetic patients (p=0.2364). Vitreous fluid concentrations of VEGF were significantly elevated in the samples from patients with PDR (1135.2 pg/ml (75.6 to 3280.0)) when compared with the samples from non-diabetic patients (19.3 pg/ml (15.6 to 40.6)) (p<0.0001) or diabetic patients without retinopathy (49.9 pg/ml (23.4 to 105.0)) (p<0.0001) (Fig 1A). Vitreous fluid concentrations of angiotensin II were also significantly elevated in the patients with PDR (25.0 pg/ml (4.0 to 76.0)) when compared with non-diabetic patients (8.9 pg/ml (4.0 to 16.0)) (p<0.0001) or diabetic patients without retinopathy (11.3 pg/ml (6.0 to 22.0)) (p<0.0001) (Fig 1B). There was a significant relation between the vitreous fluid concentration of VEGF and that of angiotensin II (p=0.702, p<0.0001) (Fig 2). Furthermore, the vitreous fluid concentrations of VEGF and angiotensin II in the patients with active PDR (VEGF: 1697.3 pg/ml (107.0 to 3280.0); angiotensin II: 30.4 pg/ml (6.0 to 76.0)) were significantly higher (p<0.0001 and p=0.0005, respectively) than in the patients with quiescent PDR (VEGF: 332.2 pg/ml (75.6 to 968.0); angiotensin II: 17.4 pg/ml (4.0 to 48.0)) (Fig 3A and B).

### Vitreous and plasma levels of VEGF, angiotensin II, and ACE

The vitreous fluid concentration of VEGF was significantly higher than the plasma VEGF level (50.9 pg/ml (15.6 to 396.0)) in the patients with PDR (p<0.0001) (Table 1). The vitreous fluid concentration of angiotensin II was also significantly higher than the plasma angiotensin II level (17.5 pg/ml (4.0 to 46.0)) in the PDR patients (p =0.0106) (Table 1). Plasma angiotensin II levels showed a significant correlation with the vitreous fluid levels of both VEGF and angiotensin II (p=0.596, p<0.0001 and p=0.755, p<0.0001, respectively). The plasma ACE level (14.2 pg/ml (2.5 to 26.0)) was significantly correlated with the vitreous fluid level of angiotensin II (p=0.372, p=0.0013), but was not significantly correlated with that of VEGF (p=0.223, p=0.0594). There was no significant relation between plasma and vitreous fluid VEGF levels (p=0.128, p=0.1490). There was also no significant relation between HbA₁c (7.6 % (4.8 to 12.3)) and the vitreous levels of VEGF or angiotensin II (p=0.220, p=0.1712 and p=0.253, p=0.0626, respectively).

### DISCUSSION

The present study showed that both angiotensin II and VEGF levels were increased in the vitreous fluid of patients with PDR and were correlated with the severity of diabetic retinopathy. In addition, angiotensin II and VEGF showed a statistically significant correlation with each other and the vitreous fluid level of angiotensin II was elevated in the active stage of PDR. We showed that not only the vitreous level of VEGF but also that of angiotensin II was significantly elevated in PDR patients when compared with non-diabetic patients or diabetic patients without retinopathy. Angiotensin II has been shown to promote the growth of capillary vessels in the chorioallantoic membrane and to stimulate new vessel formation in the rabbit cornea. A protective effect of an ACE inhibitor and of an ARA AT1 receptor on hyperoxia induced and normoxia induced neovascularisation has been demonstrated in newborn mice. However, continuous transvitreal infusion of angiotensin II alone produced retinal artery constriction, but not new vessel formation from the retina to the vitreous in the cat eye. Angiotensin II not only has a growth promoting effect, but also stimulates the induction of many cytokines and growth factors. Therefore, angiotensin II may affect neovascularisation in combination with other cytokines or growth factors.

![Figure 2](image-url) Relation between the vitreous fluid levels of VEGF and angiotensin II in PDR patients and diabetic patients without retinopathy (p=0.702, p<0.0001).

![Figure 3](image-url) (A) VEGF levels in the vitreous fluid of patients with active PDR and quiescent PDR (*p<0.0001). (B) Angiotensin II levels in the vitreous fluid of patients with active PDR and quiescent PDR (*p=0.0005).

### Table 1 Comparison of VEGF and angiotensin II concentrations in vitreous fluid and plasma in diabetic patients with PDR

<table>
<thead>
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<th>VEGF</th>
<th>Plasma</th>
<th>p Value</th>
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</thead>
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<td>1135.2 (837.7)</td>
<td>50.9 (38.9)</td>
<td>&lt;0.0001</td>
<td>51</td>
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<tr>
<td>25.0 (14.3)</td>
<td>17.5 (9.6)</td>
<td>0.0106</td>
<td>51</td>
</tr>
</tbody>
</table>

**Table 1** Comparison of VEGF and angiotensin II concentrations in vitreous fluid and plasma in diabetic patients with PDR.
In the present study, the vitreous level of angiotensin II was statistically correlated with that of VEGF. It has been suggested that induction of VEGF mRNA most probably occurs through transcriptional regulation. Receptors for angiotensin II are present on endothelial cells, and angiotensin II acts to stimulate endothelial cell growth and upregulate VEGF mRNA expression. Moreover, angiotensin II may potentiate VEGF induced angiogenic activity in the retina through increased expression of the VEGF receptor Flk-1/KDR. The effect of angiotensin II on VEGF expression was completely inhibited by an AR.

There is a possibility that angiotensin II might influence VEGF elicited signal transduction or post-transcriptional regulation of KDR. The functioning of the ocular RAS is not yet clear. However, our results and previous studies have suggested that an autocrine-paracrine relation may exist between angiotensin II and VEGF in ocular tissues. The capacity of VEGF to act as a potent angiogenic agent suggests that an angiotensin II induced increase of VEGF production could have a key role in the occurrence of neovascularisation in PDR. Further investigations will be needed to clarify the ocular interactions between angiotensin II and VEGF as well as the role of angiotensin II during neo-vascularisation in PDR.

Previous studies suggested that the vitreous fluid levels of angiotensin II and VEGF were significantly higher in active PDR than in quiescent PDR. The levels of both angiotensin II and VEGF in the vitreous fluid seem to increase during active neovascularisation and to decrease in the absence of neovascularisation, because we classified the severity of PDR according to the activity of neo-vascularisation in this study. It was previously reported that the vitreous fluid level of VEGF was higher in active PDR than in quiescent PDR and that VEGF played a major part in mediating intraocular neovascularisation in diabetic retinopathy. However, to our knowledge, the present study provides the first evidence that the vitreous fluid levels of angiotensin II are elevated in the active stage of PDR.

It is still unclear whether production of angiotensin II can occur in ocular tissues. From our results, it cannot be said whether ocular angiotensin II is located intracellularly or extracellularly and it is also impossible to determine whether angiotensin II is synthesised locally in the eye or sequestered from the plasma. Sequestration is not very likely since that would imply a specific uptake process. The local concentration of angiotensin II in the retinal microvasculature is reported to be higher than the serum and vitreous fluid levels. In the present study, the vitreous fluid level of angiotensin II was significantly higher than the plasma level, but the statistical difference was small. Furthermore, the plasma levels of angiotensin II and ACE were significantly correlated with the vitreous fluid level of angiotensin II. The level of angiotensin II in the vitreous fluid from normal porcine eyes is low to undetectable, in contrast with the relatively high levels in surrounding ocular tissues such as the RPE and choroids. Breakdown of the blood-retinal barrier (BRB) may facilitate diffusion of angiotensin II from the blood into the vitreous fluid. Since the vitreous can be considered the repository for products originating from the retina, a high level of angiotensin II might well be explained by its production and secretion from the retina. Accordingly, angiotensin II may be produced locally in ocular tissues, but little of this angiotensin II may leak into the ocular fluid under normal conditions and only when the BRB is disrupted will angiotensin II reach the vitreous fluid in high concentrations. The patients with active PDR in the present study had a hyperfluorescein pattern on FAG just before surgery. These results and previous reports suggest that disruption of the BRB may lead to elevation of the vitreous fluid concentration of angiotensin II. ACE inhibitors have been reported to maintain the BRB in diabetic patients and to have a favourable effect on diabetic retinopathy.

In summary, the present study showed that the concentrations of both angiotensin II and VEGF in the vitreous fluid of patients with PDR were significantly higher than those in non-diabetic patients or diabetic patients without retinopathy. The therapeutic implication is that inhibition of the RAS may be beneficial for the treatment of PDR. Indeed, the beneficial effects of ACE inhibition for patients with PDR have recently been shown by the EUCLID study. Our results support the possibility that the treatment with an ACE inhibitor or AR may effectively prevent the development of PDR. We hope to investigate the possibility that ACE inhibitor or ARB inhibit angiotensin II in eyes and also inhibit the progression of retinopathy in PDR in the next stage.

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Angiotensin II and VEGF in vitreous fluid

Angiotensin II and vascular endothelial growth factor in the vitreous fluid of patients with proliferative diabetic retinopathy
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