Macular circulation in patients with diabetes mellitus with and without arterial hypertension

O Arend, M Rüffer, A Remky

Abstract

Background—Previous fluorescein angiographic studies have shown alterations in the macular microcirculation in patients with diabetes mellitus and arterial hypertension. In both diseases capillary blood velocity was reduced and capillary density decreased. These changes were more pronounced in diabetic patients. We have examined the influence of arterial hypertension in combination with diabetes mellitus.

Methods—62 patients with diabetes mellitus and arterial hypertension (group 1) were matched with patients with diabetes mellitus but without arterial hypertension (group 2, match criteria: ETDRS stage of retinopathy). In all subjects fluorescein angiograms were performed with a scanning laser ophthalmoscope. Macular capillary blood velocity (CBV), perifoveal intercapillary area (PIA), the coefficient of variation of both parameters, the area of the foveal avascular zone (FAZ), and the arteriovenous passage time (AVP) were assessed by digital image analysis.

Results—Systolic and diastolic blood pressures were significantly increased in the patients with arterial hypertension (systolic p=0.0008; diastolic p=0.03). Neither dynamic measures (AVP: 1.64 (0.49) seconds (group 1), 1.72 (0.58) seconds (group 2); CBV: 1.98 (0.39) mm/s (group 1), 2.09 (0.43) mm/s (group 2)) nor morphological measures (PIA: 7985 (3137) μm² (group 1), 8338 (3376) μm² (group 2); FAZ: 0.319 (0.206) mm² (group 1), 0.363 (0.237) mm² (group 2)) were significantly different between the two groups of diabetic patients.

Conclusion—Arterial hypertension did not result in more severe macular capillary dropout than diabetes without hypertension. This might be explained by the fact that most of the patients were being treated with antihypertensive drugs.

Retinal changes associated with arterial hypertension and diabetes mellitus have been well documented. Considering the frequent coincidence of both diseases, it is important for both the prognosis and treatment to determine whether arterial hypertension aggravates diabetic retinopathy. A number of studies have shown an additive effect which varies with the type of diabetes and the systolic or diastolic blood pressure (BP). With the introduction of the scanning laser technology, angiograms of high spatial and temporal resolution can measure simultaneously both dynamic and morphological parameters. In the past, fluorescein angiographic studies in patients with diabetes or hypertension have shown reduced capillary blood velocity and decreased capillary density. Enlargement of the area of the foveal avascular zone and the perifoveal intercapillary area, as measures of capillary density, are related to decreased visual acuity and contrast sensitivity. Furthermore, in early diabetic angiopathy capillary blood velocities are reduced and the perifoveal intercapillary area is increased before microaneurysm formation occurs. Thus, digital fluorescein angiography identifies passage of dynamic and morphological changes in the retinal microcirculation of diabetic and hypertensive patients.

Several studies have examined circulatory changes in diabetic patients but the effect on the ocular circulation of diabetes mellitus combined with arterial hypertension has not been studied to date. From epidemiological data one might expect that capillary perfusion is further decreased and capillary density reduced in these patients. Previous microcirculatory studies have shown that measurement of capillary density is a valuable diagnostic tool for differentiating capillary loss. In this study we have used digital fluorescein angiograms to quantify the retinal microcirculation in diabetic patients with and without arterial hypertension. By matching for stage of retinopathy, the effect of diabetes mellitus induced capillary closure has been eliminated and an attempt has been made to define the influence of obstructive hypertensive microangiopathy.

Materials and methods

Digital recordings of scanning laser videofluorescein angiograms (Scanning Laser Ophthalmoscope; Rodenstock Instruments, Munich, Germany) and image analysing technology (PC) allow measurements of arteriovenous passage times (AVP), mean capillary blood velocity (CBV), the perifoveal intercapillary area (PIA), and the foveal avascular zone (FAZ). In brief, the CBV is determined by measuring the velocities of hypofluorescent particles through the parafoveal vasculature (PC, self-developed software). These particles are presumed to be rouleaux formations of red blood cells. The AVP time characterises the shortest passage from the dye arriving in the artery, passing through the capillary vasculature, and arriving in the corresponding vein. The AVP time is correlated with the macular
Circulation in patients with diabetes and arterial hypertension

1393

refractive error using axial length (A scan patient. All data were corrected for individual distribution of PIA and CBV in a single classification15 retinopathy. On the basis of the ETDRS classification15 the patients were assigned by stand-

ardised fundus photography to the following categories:16 Forty three (70%) patients in group 1 were treated with an antihypertensive drug, 31 (72%) with a single drug (β blocker, n=3; diuretics, n=11; calcium channel blocker, n=11; ACE inhibitor, n=6), eight (19%) with a combination of two drugs, and four (9%) with three or more agents. A total of 19 patients (30%) had no antihypertensive drug therapy. To determine the influence of antihypertensive medication these 19 patients (group A) were compared with matched patients without arter-

ial hypertension (group B). The match criterion again was the stage of diabetic retinopathy.

Informed consent was obtained from all patients before participation in the study. The protocol was approved by the human study committee of the Technical University of Aachen and followed the tenets of the Helsinki declaration. Exclusion criteria included lenticular or corneal diseases precluding detailed angiographic study. In addition, patients with allergic disease or history of sensitivity to fluorescein were excluded.

Best corrected visual acuity was determined by an ophthalmologist followed by a complete ophthalmological examination. The studied eye was selected at random if both qualified for the study. These eyes were then studied by fluorescein angiography using the scanning laser ophthalmoscope. Blood pressure (BP) and heart rate were measured before examina-

tion with an automatic device (Criticare Vital Dation Monitor, Criticon Inc, Tampa, FL, USA) in the sitting position after 5 minutes of rest. The mean arterial pressure was calculated as one third pulse pressure plus the diastolic (diast) pressure (BPdiast + (BPsys−BPdiast)/3)).17

DATA ANALYSIS

Mean values and standard deviations are given for all samples with normal distribution (Kolmogorov-Smirnov test). The Student’s t test was used for paired and unpaired samples with normal distribution. Findings with an error probability of <0.05 were considered to be statistically significant. Pearson correlation coefficients were calculated to evaluate the relationship between the parameters. p Values were obtained after carrying out Fisher’s r to z transformations.

Results

The clinical and demographic data of the two groups of patients are shown in Table 1. The patients in group 1 had a higher mean age (p<0.0001; 24%), higher systolic (p=0.0008; 12%) and diastolic BP (p=0.03; 10%), and higher mean arterial pressure (p=0.002; 8%) capillary velocities as demonstrated in healthy subjects.13

The PIA provides an estimate of capillary density in the network around the FAZ (5° circle centred over the FAZ).5 The borders of these intercapillary areas are marked interactively by drawing the surrounding capillaries with the cursor in the digital image. The area described by the cursor is calculated with the picture analysing system (Matrox Inspector 2.1, Matrox Electronic Systems Ltd, Quebec, Canada). One hundred randomly selected areas surrounded by capillaries are marked and the mean area is calculated from these measurements for each patient. The size of the FAZ is assessed using the same procedure.5

All coefficients of variation (CV=SD/mean) for CBV and PIA measurements for each sub-
ject were calculated. The mean coefficients of variation (CV(CBV), CV(PIA)) characterise the homogeneity of the perifoveal microcircu-
larisation and morphological features of each group.4 Figures 1A and B illustrate the distribution of PIA and CBV in a single patient. All data were corrected for individual refractive error using axial length (A scan ultrasonography) and keratometry.14

PATIENTS

Fluorescein angiograms were performed in 62 diabetic patients with arterial hypertension (group 1; 11 with insulin dependent diabetes mellitus (IDDM) and 51 with non-insulin dependent diabetes mellitus (NIDDM)) and in 62 diabetic patients without arterial hyper-
tension (group 2, 20 IDDM, 42 NIDDM). The two groups were matched for stage of retinopathy. On the basis of the ETDRS classification the patients were assigned by standardised fundus photography to the following groups: S10 (no retinopathy), n=4; S20 (microaneurysms only), n=6; S35 (mild non-

proliferative diabetic retinopathy (NPDR)), n=8; S43 (moderate NPDR), n=18; S53 (severe NPDR), n=9; S61 (mild proliferative retinopathy), n=8; S71 (proliferative retinopathy with high risk characteristics), n=9. The patients with severe non-proliferative or proliferative retinopathy underwent the angiographic study before pan-retinal photocoagula-

tion. The diagnosis of arterial hypertension was based on repeated readings of >160 mm Hg systolic BP and >95 mm Hg diastolic BP. The patients with arterial hyper-
tension exhibited only mild vascular alterations corresponding to stages 0 to II described by Scheie.14 A total of 19 patients (30%) had no antihypertensive drug therapy. To determine the influence of antihypertensive medication these 19 patients (group A) were compared with matched patients without arter-

ial hypertension (group B). The match criterion again was the stage of diabetic retinopathy.

www.bjophthalmol.com
Table 1. Mean (SD) clinical and demographic data of diabetic patients with (group 1) and without arterial hypertension (group 2).

<table>
<thead>
<tr>
<th></th>
<th>Group 1</th>
<th>Group 2</th>
<th>Significance*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>45.4 (10)</td>
<td>41.4 (13)</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Sex (M/F)</td>
<td>36/26</td>
<td>36/26</td>
<td>NS</td>
</tr>
<tr>
<td>Systolic BP (mm Hg)</td>
<td>150 (21)</td>
<td>132 (13)</td>
<td>p=0.03</td>
</tr>
<tr>
<td>Diastolic BP (mm Hg)</td>
<td>87 (13)</td>
<td>108 (13)</td>
<td>p=0.0002</td>
</tr>
<tr>
<td>Mean arterial pressure (mm Hg)</td>
<td>15 (8)</td>
<td>16 (6)</td>
<td>NS</td>
</tr>
<tr>
<td>Duration of diabetes (years)</td>
<td>8.4 (1.6)</td>
<td>8.6 (1.6)</td>
<td>NS</td>
</tr>
</tbody>
</table>

AVP = arteriovenous passage time; CBV = capillary blood velocity; CV(CBV) = mean coefficient of variation of capillary blood velocity; CV(CBV) = mean coefficient of variation of perifoveal intercapillary area; CV(CBV) = mean coefficient of variation of perifoveal intercapillary area; FAZ = area of foveal avascular zone; PIA = perifoveal intercapillary area; CV(PIA) = mean coefficient of variation of perifoveal intercapillary area; FAZ = area of foveal avascular zone.

AVP = arteriovenous passage time; CBV = capillary blood velocity; CV(CBV) = mean coefficient of variation of capillary blood velocity; CV(CBV) = mean coefficient of variation of perifoveal intercapillary area; CV(CBV) = mean coefficient of variation of perifoveal intercapillary area; FAZ = area of foveal avascular zone; PIA = perifoveal intercapillary area; CV(PIA) = mean coefficient of variation of perifoveal intercapillary area; FAZ = area of foveal avascular zone.

AVP = arteriovenous passage time; CBV = capillary blood velocity; CV(CBV) = mean coefficient of variation of capillary blood velocity; CV(CBV) = mean coefficient of variation of perifoveal intercapillary area; CV(CBV) = mean coefficient of variation of perifoveal intercapillary area; FAZ = area of foveal avascular zone; PIA = perifoveal intercapillary area; CV(PIA) = mean coefficient of variation of perifoveal intercapillary area; FAZ = area of foveal avascular zone.

Table 2. Microcirculatory results of diabetic patients with untreated arterial hypertension (n=19; group A) and matched normotensive diabetic patients (n=19; group B).

<table>
<thead>
<tr>
<th></th>
<th>Group A</th>
<th>Group B</th>
<th>Significance*</th>
</tr>
</thead>
<tbody>
<tr>
<td>FAZ (mm²)</td>
<td>0.302 (0.11)</td>
<td>0.267 (0.09)</td>
<td>NS</td>
</tr>
<tr>
<td>CV(CBV) (%)</td>
<td>15.9 (6.3)</td>
<td>15.6 (5.3)</td>
<td>NS</td>
</tr>
<tr>
<td>CBV (mm/s)</td>
<td>2.0 (0.41)</td>
<td>2.0 (0.37)</td>
<td>NS</td>
</tr>
<tr>
<td>PIA (µm²)</td>
<td>7015 (1726)</td>
<td>6522 (2596)</td>
<td>NS</td>
</tr>
<tr>
<td>CV(PIA) (%)</td>
<td>66.8 (43.1)</td>
<td>64.3 (27.0)</td>
<td>NS</td>
</tr>
</tbody>
</table>

The effect of antihypertensive medication on retinal circulation was studied in 19 diabetic patients with arterial hypertension receiving no antihypertensive medication (group A) and a group of 19 normotensive diabetic patients matched for stage of retinopathy (group B). There were no significant differences between the two groups for age (54 (9) v 47 (18) years), duration of diabetes (9.6 (5.7) v 13.1 (7.7) years), mean glycosylated haemoglobin (HbA1c) (8.0 (2.1) v 8.3 (1.4)%), and systolic BP (148 (24) v 143 (23) mm Hg) but significant differences were found in diastolic BP (93 (16) v 81 (17) mm Hg; p=0.04). There were no significant differences in the microcirculatory data tested between the patients in groups A and B (Table 2).

Discussion

The increased risk of cerebral, cardiovascular, and renal disease resulting from arterial hypertension in patients with diabetes mellitus has been described in various studies. However, these studies reached different conclusions with regard to the effect of reducing systolic and diastolic BP and the influence of the type of diabetes. The impact of arterial hypertension on diabetic retinopathy is either an increased rate of progression or it has little or no effect.

Past studies using scanning laser fluorescein angiography have shown decreased capillary density in patients with arterial hypertension and diabetic retinopathy. Furthermore, capillary perfusion in the macula region is decreased in diabetic and in hypertensive patients. A decrease in perfusion occurs early in diabetes when no microaneurysms are visible and is associated with increased perifoveal intercapillary area as a sign of decreased capillary density. Various studies have found hyperfusion or hypoperfusion to be pathogenic in patients with diabetes mellitus. The effect of decreased perfusion can be explained by progressive capillary closure with increased resistance and decreased perfusion. Capillary closure is a well established angiographic and histopathological sign of diabetic retinopathy. Capacil lass, as reflected by the...
Circulation in patients with diabetes and arterial hypertension

1395

...diabetic hypertension include a range of changes in the vessel wall and induction of humoral factors.

Vascular changes resulting from arterial hypertension include a range of changes in the vessel wall and can result in capillary rarefaction. In a 2-year angiographic follow-up study of hypertensive patients, the capillary density remained unchanged.

The present study was designed to examine whether progressive retinal capillary closure with attenuated circulation occurs in patients with diabetes mellitus and arterial hypertension. Both patient groups had prolonged AVP times with decreased CBV and increased FAZ and PIA compared with reference data from healthy subjects. With increasing stage of diabetic retinopathy there was a significant increase in PIA and FAZ but no interaction was seen between the presence or absence of arterial hypertension and any of the morphological or dynamic parameters measured. This suggests that the capillary closure associated with diabetes mellitus is so dominant that the arterial hypertension results in no additional detectable capillary closure. Furthermore, the mean coefficients of variation (CV(CBV), CV(PIA)) showed no further heterogeneity in perfusion or capillary density resulting from the presence of arterial hypertension.

Correlation analysis revealed a decrease in FAZ with increasing systolic BP and mean arterial pressure in diabetic patients with arterial hypertension. The significance of this finding was low and further studies are needed to confirm this trend, but it supports the similar findings in diabetic patients with unaffected visual acuity. Invest Ophthalmol Vis Sci 1997;38:1819–24.


