

Prospective cohort study of retinal vessel diameters and risk of hypertension

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Abstract

Objective To examine the relation between diameters of the retinal arterioles and 10 year incidence of hypertension.

Design Population based prospective cohort study.

Setting Beaver Dam eye study.

Participants 2451 normotensive people aged 43 to 84 years.

Main outcome measures Diameters of retinal arterioles and venules measured from digitised photographs of the retina taken at baseline. Measurements summarised as the arteriole:venule ratio, with a lower ratio indicating smaller arteriolar diameters. Incident hypertension, defined as systolic blood pressure ≥ 140 mm Hg, diastolic blood pressure ≥ 90 mm Hg, or use of antihypertensive drugs during follow up.

Results 721 participants developed hypertension over a 10 year period. Those with lower arteriole:venule ratios had a higher cumulative incidence of hypertension (incidences of 17.4%, 24.1%, 31.0%, and 45.1%, respectively, for decreasing quarters of distribution of arteriole:venule ratio). After adjustment for age and sex, participants with arteriole:venule ratios in the lowest quarter had a threefold higher risk of hypertension (odds ratio 2.95, 95% confidence interval 2.77 to 3.88) than those with ratios in the highest quarter. This association remained significant after further adjustment for baseline systolic and diastolic blood pressure and other risk factors (1.82, 1.39 to 2.40, for lowest versus highest ratio quarters).

Conclusions Narrowed retinal arterioles are associated with long term risk of hypertension, suggesting that structural alterations of the microvasculature may be linked to the development of hypertension.

Introduction

There is continuing uncertainty about the basic pathophysiological mechanisms of essential hypertension.¹ A key pathological characteristic of hypertension is the presence of small vessel disease, specifically vasoconstriction and narrowing of the peripheral small arteries and arterioles.²⁻⁶ Narrowing of these arterioles may contribute to the pathogenesis of hypertension by increasing peripheral vascular resistance. However, it is unclear whether arteriolar narrowing precedes development of hypertension or is merely a physiological response to raised blood pressure.⁷

Measurement of the retinal arterioles offers a non-invasive means to evaluate systemic associations of the human microcirculation in vivo. We have recently developed a computer based method to measure retinal arterioles and venules from photographic images.⁸ We examined the prospective association between retinal arteriolar diameters and the 10 year incidence

of hypertension in a population based cohort of normotensive people.

Methods

The Beaver Dam eye study has been described in detail elsewhere.⁹⁻¹¹ In brief, a private census of the population of Beaver Dam, Wisconsin, USA, was performed in 1987-8. Of 5924 eligible individuals, 4926 participated in the baseline examination in 1988-90,⁹ 3684 (81.1% of survivors) participated in the five year follow up examination in 1993-5,¹⁰ and 2764 (82.9%) participated in the 10 year follow up examination in 1998-2000.¹¹

To study the association between retinal vessel diameters and incident hypertension we excluded those with pre-existing hypertension at baseline ($n=1820$), missing systolic or diastolic blood pressure values ($n=13$), missing important covariate information (cigarette smoking ($n=11$), glycated haemoglobin concentrations ($n=14$), physical activity ($n=16$), total or high density lipoprotein cholesterol concentrations ($n=34$)), or ungradeable retinal photographs ($n=295$), leaving 2451 normotensive individuals who contributed data to the current analysis.

Measurement of retinal vessels

Methods used to measure and summarise retinal vessel diameters from digitised photographs followed a standardised protocol described elsewhere.^{12,13} In brief, 30° colour retinal photographs of the right eye taken at baseline were digitised by a high resolution scanner with standard settings. Graders masked to the characteristics of participants used a computer program to measure diameters of all arterioles and venules in a specified zone surrounding the optic disc. These measurements were combined into summary indices representing the average arteriolar and venular diameters of that eye.¹³ These were finally expressed as the retinal arteriole:venule ratio. The ratio compensated for possible magnification differences between eyes, and an arteriole:venule ratio of 1 indicated that arteriolar diameters were, on average, the same as venular diameters in that eye, while a smaller ratio suggested narrower arterioles.¹³ Examples of smaller and higher ratios are shown in the figure.

Definitions of hypertension

A standardised interview and examination was performed at each follow up visit.¹⁴ Blood pressure was measured with a random zero sphygmomanometer according to the hypertension detection and follow up programme protocol at each examination.¹⁵ We used the average of two measurements for analysis.

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Retinal photographs showing normal retinal arterioles with arteriole:venule ratio of 1.08 (left) and narrowed retinal arterioles with arteriole:venule ratio of 0.64 (right)

We defined hypertension as systolic blood pressure ≥ 140 mm Hg, diastolic blood pressure ≥ 90 mm Hg, or the combination of self reported diagnosis of high blood pressure and use of antihypertensive drugs. Participants who met these criteria at either the five year or 10 year follow up examination were defined as having incident hypertension.

Definitions of other variables

The interview and examination included measurements of body mass index and questions about family income, education, physical activity, cigarette smoking, and alcohol consumption. Non-fasting blood specimens were obtained from all participants.¹⁴ From them we obtained plasma concentrations of glycated haemoglobin, and serum concentrations of total cholesterol and high density lipoprotein cholesterol. We defined participants as having diabetes mellitus if they had a history of diabetes or diabetes had been newly diagnosed (no medical history of diabetes mellitus in the presence of raised glycated haemoglobin concentration and a random blood sugar > 11 nmol/l).¹⁴

Statistical analysis

We categorised retinal vessel data (retinal arteriole:venule ratio, retinal arteriolar, and venular diameters) into quarters, with lower quarters indicating smaller ratios and vessel diameters. We also analysed data on retinal vessels as continuous variables (for example, per SD change in arteriole:venule ratio). We used logistic regression models to determine the odds ratio for incident hypertension, controlling simultaneously for possible confounders including age, sex, diabetes, body mass index, physical activity (number of times a week), total and high density lipoprotein cholesterol concentrations, cigarette smoking, alcohol consumption (see bmj.com for further details); and baseline blood pressure.

Results

Table 1 shows baseline characteristics of participants, according to arteriole:venule ratio. Lower arteriole:venule ratios were associated with older age, male sex, diabetes, higher glycated haemoglobin concentration, higher systolic and diastolic blood pressure, higher

Table 1 Baseline characteristics of study population in relation to retinal arteriole:venule ratio. Figures are means unless stated otherwise

	Quarter of retinal arteriole:venule ratio*				P value†
	1(n=612)	2(n=612)	3(n=613)	4(n=614)	
Age (years)	62.1	58.9	59.9	59.6	<0.001
No (%) of men	295 (48.2)	288 (47.1)	280 (45.7)	265 (43.1)	<0.001
Family income (\$10 000‡)	5.0	5.1	5.0	5.1	0.07
Length of schooling (years)	12.0	12.2	12.9	12.2	0.08
No (%) with diabetes	43 (7.1)	38 (6.2)	33 (5.4)	31 (5.1)	<0.001
Glycated haemoglobin (%)	6.7	5.9	5.7	5.4	<0.001
Systolic blood pressure (mm Hg)	129.2	127.1	122.8	120.2	<0.001
Diastolic blood pressure (mm Hg)	77.3	75.4	75.2	72.0	<0.001
Body mass index (kg/m ²)	28.4	28.2	28.1	26.2	0.05
Total cholesterol (mmol/l)	6.0	5.9	5.9	5.9	0.05
HDL cholesterol (mmol/l)	1.3	1.4	1.4	1.4	0.06
Cigarette smoking (pack years)	20.8	18.9	18.1	14.2	<0.001
Alcohol intake (ounces/week)§	4.8	4.4	4.1	3.4	0.05

* Ranges: 1st=0.50-0.67, 2nd=0.68-0.71, 3rd=0.72-0.75, 4th=0.76-1.04.

† For test of difference in characteristics by arteriole:venule ratio quarters based on analysis of variance or χ^2 tests as appropriate.

‡ £5684 (€8413).

§ See bmj.com for further explanation.

body mass index, higher total cholesterol concentration, lower high density lipoprotein cholesterol concentration, cigarette smoking, and alcohol consumption.

Over the 10 year period 721 people developed hypertension. After adjustment for age and sex, those who developed incident hypertension had significantly lower mean baseline arteriole:venule ratio (0.69 *v* 0.72, $P < 0.001$) and smaller retinal arteriolar diameter (166.0 μm *v* 172.5 μm , $P < 0.001$) compared with those who did not develop hypertension. In contrast, baseline retinal venular diameter was similar in participants who did and did not develop hypertension (241.0 μm *v* 241.7 μm , $P = 0.17$).

Table 2 shows the incidence and odds ratio for incident hypertension in relation to the retinal arteriole:venule ratio. After we controlled for age and sex, participants with an arteriole:venule ratio in the first (lowest) quarter were three times as likely to develop hypertension than those with a ratio in the fourth quarter (odds ratio 2.95, 95% confidence interval 2.27 to 3.85). Further adjustment for risk factors associated with hypertension had minimal impact on this association (2.72, 2.08 to 3.54). This association was attenuated but remained significant even after further adjustment for baseline systolic and diastolic blood pressure and pulse pressure (1.82, 1.39 to 2.40). The pattern of association for retinal arteriolar diameter was essentially similar to that of the arteriole:venule ratio, but retinal venular diameter was unrelated to incident hypertension (see table A on [bmj.com](#)).

When we analysed the arteriole:venule ratio as a continuous variable, each SD decrease (a decrease of 0.07) was associated with a 30% increase in odds of hypertension. There were no significant interactions with sex or other risk factors for hypertension (diabetes, body mass index, waist:hip ratio, cigarette smoking, and alcohol consumption).

Finally, we constructed multiple linear regression models of retinal arteriole:venule ratio and change in blood pressure between baseline and the 10 year follow up. After we excluded participants taking antihypertensive medications at the follow up examinations, baseline retinal arteriole:venule ratio was inversely associated with a 10 year change in blood pressure (see table B on [bmj.com](#)). When we included the blood pressures of those who were taking antihypertensive medications at the follow up examinations, the magnitude of this inverse association was attenuated, as expected, but remained significant ($P < 0.05$).

Discussion

In this prospective cohort study we showed that smaller retinal arteriolar diameter is associated with increased risk of hypertension over a 10 year period. After we controlled for age, sex, and other risk factors for hypertension, people with an arteriole:venule ratio in the lowest quarter were about 80% more likely to develop hypertension than those with a ratio in the highest quarter. Furthermore, each SD reduction in the arteriole:venule ratio was associated with a 30% increased risk of hypertension, independent of baseline blood pressure and other risk factors.

Experimental studies have long indicated that peripheral arteriolar constriction and increased vascular resistance are classic features of essential

Table 2 Odds ratio (95% confidence interval) for incident hypertension in relation to retinal arteriole:venule ratio

Ratio	No at risk	No of cases	Incident hypertension		
			Adjusted for age and sex	Multivariable*	Multivariable†
1st quarter	612	276	2.95 (2.27 to 3.85)	2.72 (2.08 to 3.54)	1.82 (1.39 to 2.40)
2nd quarter	612	190	2.05 (1.57 to 2.69)	1.91 (1.46 to 2.51)	1.47 (1.11 to 1.95)
3rd quarter	613	148	1.17 (0.87 to 1.55)	1.13 (0.85 to 1.51)	0.98 (0.73 to 1.32)
4th quarter	614	107	1‡	1‡	1‡
1 SD decrease	2451	721	1.54 (1.38 to 1.72)	1.50 (1.34 to 1.68)	1.31 (1.18 to 1.45)

*Further adjusted for diabetes, glycated haemoglobin, body mass index, physical activity, total and high density lipoprotein cholesterol concentrations, cigarette smoking, and alcohol consumption.

†Further adjusted for systolic and diastolic blood pressure and pulse pressure.

‡Reference.

hypertension. Narrowed small arterioles may cause hypertension by increasing peripheral vascular resistance and haemodynamic load. However, partly because of a lack of prospective studies, there is no clear consensus as to whether reduced arteriolar size arises as an adaptive consequence of raised blood pressure or is a pathogenic factor in the development of hypertension.²⁻⁸ Our study now provides longitudinal data in a large unselected cohort of normotensive people to support the concept that arteriolar narrowing precedes the subsequent development of hypertension.

Narrowing of the retinal arterioles seen clinically reflects intimal thickening and medial hyperplasia, hyalinisation, and sclerosis seen histopathologically.¹⁶ We have previously shown that such narrowing is not only associated with raised blood pressure but also with systemic markers of inflammation and possibly endothelial dysfunction.¹⁷ It is therefore possible that microvascular narrowing due to these mechanisms is involved in the pathogenesis of hypertension.

There are few studies for direct comparison. The atherosclerosis risk in communities study also found that narrowed retinal arteriolar diameter was associated with an increased risk of hypertension developing over a three year period in 5628 normotensive people (multivariable adjusted relative risk 1.62, 95% confidence interval 1.21 to 2.18, comparing lowest to highest fifths of arteriole:venule ratio).¹⁸ That study, however, was limited by a relatively short interval to follow up. We had a considerably longer follow up, and our results provide stronger evidence that reduced retinal arteriolar diameter precedes the long term risk of hypertension.

Limitations

Several limitations of this study should be highlighted. Firstly, blood pressure measurements taken once during each five year interval may have resulted in some misclassification of hypertension. However, we have no reason to believe that potential misclassification error would substantially bias our results (except towards the null). Secondly, residual confounding from pre-existing hypertension or other unmeasured factors may explain some of these associations. For example, the strength of the association between arteriolar narrowing and risk of hypertension was substantially weakened after we adjusted for baseline blood pressure, suggesting that baseline blood pressure was an important confounder. Finally, selective mortality may have masked some associations. If people with reduced arteriolar diameters who were at risk of developing hypertension were more likely to die before the follow up examinations, these associations could be falsely

What is already known on this topic

Narrowing of the small blood vessels may have a role in the pathogenesis of hypertension, though few prospective trials support such a hypothesis

What this study adds

People with smaller retinal arteriolar diameters are more likely to develop hypertension than people with larger arteriolar diameters, independent of known risk factors for hypertension

This finding suggests that narrowed arterioles are associated with the development of hypertension and therefore that small vessel disease may be a target for antihypertensive treatment

attenuated. Strengths of our study include its population based nature, long follow up, and quantitative evaluation of retinal arteriolar diameters.

Clinical implications

Our results may have potential clinical implications. There is increasing recognition of the value of specifically targeting the microcirculation in the prevention and treatment of hypertension and its complications.^{19,20} For example, there is some evidence that angiotensin converting enzyme inhibitors have direct beneficial effects on microvascular structure and may have therapeutic value in preventing cardiovascular morbidity beyond that of lowering blood pressure.²¹ Our study indicates that further research to evaluate approaches to treatment focused on the microcirculation may be important.

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Crunching the numbers

A 70 year old man with giant cell arteritis complains of pains in his shoulders. What percentage of patients with giant cell arteritis also have polymyalgia rheumatica?

- A: Less than 25%
- B: 25-40%
- C: Greater than 60%
- D: 80%

The use of imprecise terms in examination questions causes considerable confusion. A common error when writing such questions is to use inconsistent numeric data.¹ When you use numeric data, you should list the options in numeric order and also in a single format (that is, as single terms or as ranges).

In the example above, options A, B, and C are expressed as ranges whereas option D is a specific percentage. Mixing the formats like this is confusing, and readers could easily mistake option C (greater than 60%) with option D (80%).

BMJ Learning offers online learning modules, throughout which we ask questions, but we avoid imprecise and ambiguous questions. Our new learning module on giant cell arteritis outlines how to diagnose and treat this condition. It also gives an update on recent advances such as the role of colour flow Doppler in diagnosis and how best to avoid corticosteroid induced osteoporosis. You will also find out what percentage of patients with giant cell arteritis also have polymyalgia rheumatica.

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