BMJ Open

Hyperuricemia is independently associated with endothelial dysfunction in postmenopausal women but not in premenopausal women

Tatsuya Maruhashi, Ayumu Nakashima, Junko Soga, Noritaka Fujimura, Naomi Idei, Shinsuke Mikami, Yumiko Iwamoto, Masato Kajikawa, Takeshi Matsumoto, Takayuki Hidaka, Yasuki Kihara, Kazuaki Chayama, Chikara Goto, Kensuke Noma, Hirofumi Tomiyama, Bonpei Takase, Akira Yamashina, Yukihito Higashi

ABSTRACT

Objectives: The purpose of this study was to determine the relationships between uric acid, endothelial function and cardiovascular risk factors and to investigate whether menopausal status was associated with the relationship between uric acid and endothelial function in women.

Design: Cross-sectional study.

Setting: 3 general hospitals in Japan.

Participants: 749 Japanese women aged 30–74 years recruited from people who underwent health-screening examinations with agreement for measurement of vascular function.

Measures: We measured serum concentrations of uric acid and flow-mediated vasodilation (FMD). Percentage of FMD (peak diameter/baseline diameter) was used for analysis. Endothelial dysfunction was defined as FMD ≤4.90%, division point for the lowest tertile and the middle tertile of FMD. Menopause women were defined as participants without menstruation for over 1 year or participants with a history of hysterectomy or bilateral oophorectomy.

Results: Of the 749 participants, 368 (49.1%) were premenopausal women and 381 (50.9%) were postmenopausal women. Age, body mass index, systolic blood pressure, total cholesterol, triglycerides, glucose, estimated glomerular filtration rate and Framingham risk score were significantly correlated with serum uric acid level. FMD showed a gradual decrease in accordance with the serum uric acid level. FMD showed a significant decrease in accordance with the serum uric acid level (≤4 mg/dL, 6.85±3.65%; 4 to <5 mg/dL, 6.79±3.60%; 5 to <6 mg/dL, 6.24±3.58%; ≥6 mg/dL, 5.27±3.18%; p=0.01). Multivariate analysis revealed that uric acid was a significantly independent risk factor for endothelial dysfunction in postmenopausal women (OR 1.23, 95% CI 1.01 to 1.50), but not in premenopausal women.

Conclusions: These findings suggest that uric acid can be used as a risk marker of endothelial dysfunction in a female population, and particularly as an independent risk factor in postmenopausal women but not in premenopausal women.

Registration number of the study: UMIN000003409.
understood, several lines of evidence suggest that elevated uric acid impairs endothelial function by inducing intracellular oxidative stress and inflammation through activation of the local renin-angiotensin system, particularly angiotensin II, and the pro-oxidant effect of uric acid per se, once absorbed into endothelial cells.16–18

Endothelial dysfunction is established in the initial step of atherosclerosis and plays an important role in the development of atherosclerotic conditions, leading to cardiovascular outcomes.19 Recently, the measurement of flow-mediated vasodilation (FMD) as an index of endothelium-dependent vasodilation has been widely used as a method for assessing endothelial function.20–23 In addition, growing evidence has shown that endothelial function assessed by FMD can serve as an independent predictor of cardiovascular events.24–27 Recently, several investigators, including us, have shown a relationship between uric acid and endothelial function as assessed by FMD.28–34 As for the relationship between uric acid and FMD in men, previous studies are consistent in demonstrating that uric acid is a significantly independent risk factor for FMD.29–34 However, there is little information on whether uric acid is an independent risk factor for endothelial dysfunction in women. In addition, it remains unclear whether menopausal status is associated with the relationship between uric acid and endothelial function. It is clinically important to confirm the role of uric acid per se in atherosclerosis. We therefore investigated the relationships between uric acid, FMD and cardiovascular risk factors to see whether menopausal status was associated with the relationship between uric acid and endothelial function in women who underwent health-screening examinations.

METHODS

Subjects

A total of 5321 Japanese adults aged 17–86 years who underwent health-screening examinations with agreement for measurement of vascular function were enrolled in the Flow-mediated Dilation Japan Registry between 1 April 2010 and 31 August 2012 at three general hospitals in Japan. All employees have an obligation to undergo health-screening examinations every year under the regulations of the Society-managed Health Insurance Union in Japan. In accordance with the regulations, we performed health-screening examinations. From the registry, 895 women aged 30–74 years were recruited for this study. Among the 895 participants, information on the serum uric acid levels, menopause and phases of menstrual cycle was available for 797 participants. Participants during menstruation (n=28), participants receiving hormone replacement therapy (n=3), one participant who was pregnant, participants who had been previously diagnosed with cardiovascular diseases (n=15) and one participant who was being treated with an antihyperuricemic drug were excluded. Finally, 749 women without cardiovascular diseases were enrolled in this study. Hypertension was defined as systolic blood pressure of ≥140 mm Hg or diastolic blood pressure ≥90 mm Hg, in a sitting position, on at least three different occasions.35 Patients with secondary forms of hypertension were excluded in all patients with hypertension on the basis of complete history; physical examination; radiological and ultrasound examinations; urinalysis; plasma renin activity; plasma aldosterone and norepinephrine concentrations; serum creatinine, potassium, calcium and free thyroxine concentrations; and 24 h urinary excretion of 17-hydroxycorticosteroids, 17-ketogenic steroids and vanillylmandelic acid. Diabetes was defined according to the American Diabetes Association recommendations.36 Dyslipidemia was defined according to the third report of the National Cholesterol Education Programme (NCEP) Expert Panel on Detection, Evaluation and Treatment of High Blood Cholesterol in Adults (ATP-III).37 We defined smokers as those who were current smokers. Measurement of FMD was performed without withholding medications. Framingham risk score was calculated by points of risk factors: age, total cholesterol level, high-density lipoprotein cholesterol level, systolic blood pressure and smoking status.38 Diagnosis of metabolic syndrome was made according to the criteria of NCEP ATP-III.39 Thus, metabolic syndrome was diagnosed when three or more of the following risk determinants were present (1) waist circumference of ≥88 cm, (2) triglyceride ≥1.7 mmol/L (150 mg/dL), (3) high-density lipoprotein cholesterol <1.29 mmol/L (50 mg/dL), (4) blood pressure ≥130 and/or 85 mm Hg or (5) fasting blood glucose ≥6.11 mmol/L (110 mg/dL). The estimated glomerular filtration rate (eGFR) was calculated using the Japanese eGER equation.39 Chronic kidney disease (CKD) was defined as eGFR <60 mL/min/1.73 m².40 Menopausal women were defined as participants without menstruation for over 1 year or participants with a history of hysterectomy or bilateral oophorectomy.41 The ethical committees of our institutions approved the study protocol. Written informed consent for participation in the study was obtained from all participants.

Study protocol

We measured vascular responses to reactive hyperaemia in the brachial artery in all participants. Participants fasted the previous night for at least 12 h. The study began at 8:30. The participants were kept in a supine position in a quiet, dark, air-conditioned room (constant temperature 22–25°C) throughout the study. After remaining in the supine position for 30 min, blood samples were obtained for measurement of basal fasting serum concentrations of total cholesterol, triglycerides, high-density lipoprotein cholesterol, low-density lipoprotein cholesterol, creatinine, glucose and uric acid. Then FMD was measured. The observers were blind to the form of examination. Serum uric acid levels were measured by the uricase-peroxidase method (JCA-BM6010; JEOL Ltd, Tokyo, Japan). Levels of serum total cholesterol, triglycerides, high-density lipoprotein cholesterol, low-density lipoprotein cholesterol and creatinine were
enzymatically measured (JCA-BM6010). Glucose levels were measured by the glucose oxidase immobilised oxygen electrode method (GA08II; A&T, Yokohama, Japan).

**Measurement of FMD**

The participants remained supine throughout the study. The vascular response to reactive hyperaemia in the brachial artery was used for assessment of endothelium-dependent FMD. A high-resolution linear artery transducer was coupled to computer-assisted analysis software (UNEXEF18G, UNEX Co, Nagoya, Japan) that used an automated edge detection system for the measurement of the brachial artery diameter. A blood pressure cuff was placed around the forearm. The brachial artery was scanned longitudinally 5–10 cm above the elbow. When the clearest B-mode image of the anterior and posterior intimal interfaces between the lumen and vessel wall was obtained, the transducer was held at the same point throughout the scan by a special probe holder (UNEX Co) to ensure consistency of the image. Depth and gain settings were set to optimise the images of the arterial lumen wall interface. When the tracking gate was placed on the intima, the artery diameter was automatically tracked, and the waveform of diameter changes over the cardiac cycle was displayed in real time using the FMD mode of the tracking system. This allowed the ultrasound images to be optimised at the start of the scan and the transducer position to be adjusted immediately for optimal tracking performance throughout the scan. Pulsed Doppler flow was assessed at baseline and during peak hyperaemic flow, which was confirmed to occur within 15 s after cuff deflation. Blood flow velocity was calculated from the Doppler data and was displayed as a waveform in real time. The baseline longitudinal image of the artery was acquired for 30 s, and then the blood pressure cuff was inflated to 50 mm Hg above systolic pressure for 5 min. The longitudinal image of the artery was recorded continuously until 5 min after cuff deflation. Pulsed Doppler velocity signals were obtained for 20 s at baseline and for 10 s immediately after cuff deflation. Changes in brachial artery diameter were immediately expressed as the percentage change relative to the vessel diameter before cuff inflation. FMD was automatically calculated as the percentage change in peak vessel diameter from the baseline value. Percentage of FMD (peak diameter−baseline diameter/baseline diameter) was used for analysis. Blood flow volume was calculated by multiplying the Doppler flow velocity (corrected for the angle) by the heart rate and vessel cross-sectional area (πr²). Reactive hyperaemia was calculated as the maximum percentage increase in flow after cuff deflation compared with baseline flow.

**Statistical analysis**

Results are presented as mean±SD. All reported probability values were two-sided, and a probability value of <0.05 was considered to be statistically significant. Categorical variables were compared by means of a χ² test. Comparisons between the groups categorised according to the serum uric acid level were carried out using P for trend analysis with Bonferroni’s test for post hoc comparisons. Relations between variables were determined by Spearman’s correlation coefficients analysis. Stepwise multivariate logistic regression analyses were performed to identify factors associated with endothelial dysfunction among potential confounders (p<0.20) in univariate analysis. Using variables associated with endothelial dysfunction in stepwise multiple regression analyses, we performed the final logistic regression analyses for which results are presented in the results section. The data were processed using the software package Stata V.9 (Stata Co, College Station, Texas, USA).

**RESULTS**

**Baseline clinical characteristics**

The baseline clinical characteristics are summarised in table 1. Of the 749 participants, 112 (15.0%) had hypertension, 262 (35.0%) had dyslipidemia, 34 (4.5%) had diabetes mellitus, 59 (7.9%) were current smokers, 65 (8.7%) had metabolic syndrome and 56 (8.4%) had chronic kidney disease. The mean Framingham risk score was 3.94±3.62%. The mean value of the serum uric acid level was 4.44±1.09 mg/dL (median 4.30 mg/dL; IQR 3.80–5.00 mg/dL; range 0.80–10.0 mg/dL). The mean value of FMD was 6.59±3.60% (range –4.70 to

| Variables                  | Total (n=749) |
|---------------------------|---------------|
| Age (years)               | 50.2±10.6     |
| Body mass index (kg/m²)   | 22.0±3.4      |
| Systolic blood pressure (mm Hg) | 121.1±19.0    |
| Diastolic blood pressure (mm Hg) | 73.9±12.4     |
| Heart rate (bpm)          | 64.4±10.2     |
| Total cholesterol (mmol/L) | 5.37±0.88     |
| Triglycerides (mmol/L)    | 1.07±0.71     |
| HDL-cholesterol (mmol/L)  | 1.80±0.42     |
| LDL-cholesterol (mmol/L)  | 3.11±0.77     |
| Glucose (mmol/L)          | 5.32±1.47     |
| Uric acid (mg/dL)         | 4.44±1.09     |
| eGFR (mL/min/1.73 m²)     | 77.1±13.3     |
| Framingham risk score (%) | 3.94±3.62     |
| Hypertension (n (%))      | 112 (15.0)    |
| Dyslipidemia (n (%))      | 262 (35.0)    |
| Diabetes mellitus (n (%)) | 34 (4.5)      |
| Smoking (n (%))           | 59 (7.9)      |
| Metabolic syndrome (n (%)) | 65 (8.7)      |
| Chronic kidney disease (n (%)) | 56 (8.4)     |
| Menopause (n (%))         | 381 (50.9)    |
| Flow-mediated vasodilation (%) | 6.59±3.60    |
| Baseline brachial artery diameter (mm) | 3.40±0.47    |

eGFR, estimated-glomerular filtration rate; HDL, high-density lipoprotein; LDL, low-density lipoprotein.
Relationships between the serum uric acid level and cardiovascular risk factors

Univariate regression analysis revealed that the serum uric acid level correlated significantly with age (r=0.21, p<0.001), body mass index (r=0.26, p<0.001), systolic blood pressure (r=0.17, p<0.001), diastolic blood pressure (r=0.17, p<0.001), total cholesterol (r=0.23, p<0.001), triglycerides (r=0.25, p<0.001), high-density lipoprotein cholesterol (r=−0.13, p=0.002), low-density lipoprotein cholesterol (r=0.22, p<0.001), glucose (r=0.12, p=0.004) and eGFR (r=−0.27, p=0.001). Participants were categorised according to the serum uric acid levels (table 2). Age, body mass index, systolic blood pressure, total cholesterol, triglycerides, glucose and Framingham risk score were significantly increased and eGFR was significantly decreased with an increase in the serum uric acid level. As for the prevalence of cardiovascular risk factors, there were significant increases in the prevalence of hypertension, dyslipidemia, current smoking, metabolic syndrome, chronic kidney disease and menopause in relation to an increase in the serum uric acid level.

Table 2 Clinical characteristics according to the serum uric acid levels†‡§

| Variables                  | Uric acid categories (mg/dL) |      |      |      | p Value |
|----------------------------|------------------------------|------|------|------|---------|
| Uric acid (mg/dL)           | <4.0                         | 3.38±0.54 | 4.42±0.28 | 5.33±0.26 | 6.90±0.90 |
| (n=245)                     |                              | 4.0 to <5.0 | (n=302) | 5.0 to <6.0 | (n=144) | 6.0≤ | (n=58) |
| Age (years)                 | 48.3±10.9                    | 49.4±10.1 | 53.5±9.9† | 54.7±10.0† | <0.001 |
| Body mass index (kg/m²)     | 20.8±2.8                     | 22.0±3.4* | 23.0±3.3* | 24.3±3.9† | <0.001 |
| Systolic blood pressure (mm Hg) | 117.2±17.6                 | 120.4±18.3 | 125.9±20.4* | 129.2±20.5* | <0.001 |
| Diastolic blood pressure (mm Hg) | 71.5±11.4                 | 73.6±12.1 | 77.4±13.4* | 77.0±12.5* | <0.001 |
| Heart rate (bpm)            | 64.6±9.6                     | 65.5±12.4 | 65.8±11.1 | 0.14 |
| Total cholesterol (mmol/L)  | 5.20±0.82                    | 5.33±0.86 | 5.60±0.91† | 5.67±0.88* | <0.001 |
| (mol/L)                     | 5.0 to <6.0                  | 5.23±0.91 | 5.33±0.88* | 5.38±0.85* | <0.001 |
| Triglycerides (mmol/L)      | 0.89±0.51                    | 0.97±0.50 | 1.32±0.81† | 1.69±1.32* § | <0.001 |
| HDL-cholesterol, (mmol/L)   | 1.83±0.38                    | 1.83±0.41 | 1.69±0.45† | 1.69±0.54  | 0.001 |
| (mmol/L)                    | 3.00±0.70                    | 3.10±0.77 | 3.31±0.80* | 3.19±0.96  | 0.006 |
| Glucose (mmol/L)            | 5.09±0.84                    | 5.28±1.38 | 5.52±2.00* | 6.03±2.09* | <0.001 |
| (mg/dL)                     | 4.0 to <5.0                  | 4.24±1.32* | 4.74±1.30* | 6.77±3.13* § | <0.001 |
| Flow-mediated vasodilation (%) | 6.85±3.65                    | 6.79±3.60 | 6.24±3.58 | 5.27±3.18† | 0.01 |
| Baseline brachial artery diameter (mm) | 3.35±0.46 | 3.40±0.48 | 3.43±0.46 | 3.61±0.51† | <0.002 |
| Framingham risk score (%)   | 3.27±3.44                    | 3.49±2.98 | 5.08±3.65† | 6.33±5.42* | <0.001 |
| Hypertension (n (%))        | 25 (10.2)                    | 34 (11.3) | 34 (23.6)* | 19 (32.8)* | <0.001 |
| Dyslipidemia (n (%))        | 59 (24.1)                    | 90 (29.8) | 72 (50.0)* | 41 (70.7)* § | <0.001 |
| Diabetes mellitus (n (%))   | 9 (3.7)                      | 8 (2.7)  | 9 (6.3)  | 8 (13.8)* | 0.008 |
| Smoking (n (%))             | 12 (4.9)                     | 23 (7.6)  | 15 (10.4) | 9 (15.5)  | 0.04 |
| Metabolic syndrome (n (%))  | 12 (4.9)                     | 18 (6.0)  | 19 (13.2)* | 16 (28.1)* | <0.001 |
| Chronic kidney disease (n (%)) | 10 (4.6)          | 22 (7.8)  | 14 (10.9) | 10 (24.4)* | <0.001 |
| Menopause (n (%))           | 103 (42.0)                   | 146 (48.3) | 91 (63.2)* | 41 (70.7)* | <0.001 |
| Endothelial dysfunction (n (%)) | 75 (30.6)         | 87 (28.8) | 50 (34.7) | 32 (55.2)* § | <0.002 |

p Values for comparisons across the uric acid categories were performed with ANOVA for continuous variables and χ² test for categorical variables.

†p<0.05 vs serum uric acid level<4 mg/dL group.
‡p<0.05 vs serum uric acid level 4 to <5 mg/dL group.
§p<0.05 vs serum uric acid level 5 to <6 mg/dL group.

ANOVA, analysis of variance; eGFR, estimated glomerular filtration rate; HDL, high-density lipoprotein; LDL, low-density lipoprotein.
menopause on the relationship between the serum uric acid level and endothelial function (see online supplementary table S1). Of the 749 participants, 368 (49.1%) were premenopausal women and 381 (50.9%) were postmenopausal women. Postmenopausal women were significantly older than premenopausal women (58.6±6.3 vs 41.5±6.1 years, p<0.001). The ages ranged from 30 to 54 years in premenopausal women and 31 to 74 years in postmenopausal women. The serum uric acid level was significantly higher in postmenopausal women than in premenopausal women (4.6±1.11 vs 4.2±1.03 mg/dL, p<0.001). FMD was significantly impaired in postmenopausal women compared to premenopausal women (5.51±3.14 vs 7.77±3.71%, p<0.001), and the prevalence of endothelial dysfunction was significantly higher in postmenopausal women than in premenopausal women (66.4% vs 34.0%, p<0.001). In postmenopausal women, multiple logistic regression analysis revealed that the serum uric acid level was significantly associated with endothelial dysfunction after adjustment for age (OR, 1.26; 95% CI 1.05 to 1.53) (table 4). After adjustment for other risk parameters, the association between the uric acid level and endothelial dysfunction persisted after adjustment for other risk factors (OR 1.23; 95% CI 1.01 to 1.50). In contrast, there was no significant association between the serum uric acid level and endothelial dysfunction in premenopausal women (OR 1.14; 95% CI 0.90 to 1.44; table 4).

**Table 3. Multivariate analysis of the relation between endothelial dysfunction and variables.**

| Model | Variables                  | OR (95% CI)      | p Value |
|-------|----------------------------|------------------|---------|
| 1     | Uric acid (mg/dL)          | 1.30 (1.13 to 1.50) | <0.001  |
| 2     | 1+age                      | 1.20 (1.03 to 1.39) | 0.02    |
| 3     | 2+other variables          | 1.13 (0.97 to 1.32) | 0.11    |

Initial factors included in model 3 were body mass index, systolic blood pressure, total cholesterol, triglycerides, high-density lipoprotein cholesterol, low-density lipoprotein cholesterol, glucose, eGFR and smoking. Using stepwise multiple regression analysis, we selected high-density lipoprotein cholesterol, glucose and smoking as independent factors for endothelial dysfunction (other variables).

**DISCUSSION**

In the present study, we demonstrated that the uric acid level was significantly associated with cardiovascular parameters and prevalence of cardiovascular risk factors. Although FMD showed a graded decrease according to the serum uric acid level, uric acid was not an independent risk factor for endothelial dysfunction after adjustment for other cardiovascular risk parameters. However, when participants were divided into two groups according to menopausal status, uric acid was a significantly independent risk factor for endothelial dysfunction even after adjustment for other cardiovascular risk parameters in postmenopausal women, but not in premenopausal women. To our knowledge, this is the first report showing a difference in the relationship between uric acid and endothelial dysfunction depending on menopausal status in women.

As a marker of atherosclerotic diseases, it has been demonstrated that the serum uric acid level is associated with cardiovascular risk factors and subclinical atherosclerosis, such as hypertension, metabolic syndrome, kidney disease, coronary artery calcification and carotid atherosclerosis. In the present study, we confirmed that the serum uric acid level was significantly associated with age, body mass index, systolic blood pressure, total cholesterol, triglycerides, glucose and eGFR, most of which are components of metabolic syndrome and CKD. Indeed, the prevalence of hypertension, dyslipidemia, metabolic syndrome and CKD was linearly increased in relation to an increase in the serum uric acid level. An association between uric acid and metabolic syndrome has been demonstrated by the fact that the uric acid level is often elevated in participants with metabolic syndrome and that the prevalence of metabolic syndrome increases linearly in relation to the uric acid level. The mechanism behind the association between an increase in the serum uric acid level and metabolic syndrome is thought to be hyperinsulinaemia, mediated by insulin resistance and visceral adiposity, leading to an increase in uric acid absorption in renal tubules. Uric acid levels are also frequently elevated in patients with kidney disease as a result of reduction in GFR and renal urate excretion. In addition, several epidemiological studies have shown a link between the elevated serum uric acid level and subsequent cardiovascular events. The association between the serum uric acid level and cardiovascular disease has been reported.

Figure 1. Bar graphs showing flow-mediated vasodilation in women categorised according to the serum uric acid level.
to be generally stronger in women than in men.9 40–48 These findings suggest that the uric acid level is a biochemical marker of atherosclerotic disease and a useful predictor of the development of cardiovascular diseases.

Despite the association between the elevated uric acid level and cardiovascular conditions, uric acid has not been established as an independent causal risk factor for cardiovascular disease because of the link between uric acid and established cardiovascular risk factors. Another possible explanation for the difficulty in determining the role of uric acid in cardiovascular disease is that uric acid may function as a powerful antioxidant and as a scavenger of singlet oxygen and radicals.49 50 In patients with cardiovascular disease, an increase in the uric acid level might therefore be considered as a compensatory mechanism to counteract the oxidative stress induced in these conditions. Although it remains controversial whether uric acid is a causative factor or merely a marker of cardiovascular disease, some epidemiological studies have demonstrated that the association between uric acid and cardiovascular disease remains significant even after adjustment for concomitant risk factors, suggesting that uric acid per se is an independent risk factor for cardiovascular disease.5 7–10

Although the precise mechanism behind the relationship between uric acid and cardiovascular disease remains to be elucidated, growing evidence indicates that endothelial dysfunction induced by uric acid plays an important role in the development of cardiovascular disease.15 Several lines of evidence suggest that uric acid also has proinflammatory effects on endothelial cells, leading to the reduction of endothelial nitric oxide bioavailability and consequent endothelial dysfunction.17 18 Yu et al.46 demonstrated that uric acid enhances the production of reactive oxygen species by activation of the local renin-angiotensin system, particularly angiotensin II, in human endothelial vascular cells and that probenecid, an inhibitor of urate transporter, inhibits uric acid-induced oxidative stress. Thus, uric acid is thought to mediate endothelial dysfunction by generating oxidative stress once absorbed into endothelial cells. In a clinical setting, the measurement of FMD as an index of endothelium-dependent vasodilation in the brachial artery using high-resolution ultrasound has been widely used as a method for assessing vascular function.20–23

Although several investigators have shown that uric acid correlates inversely with FMD and that uric acid is an independent predictor of FMD, the participants in those studies were limited to a small number of participants or highly selected participants, such as patients with increased cardiovascular risk,30 hyperuricemia32 or non-diabetic CKD.33 In addition, uric acid has been shown to be a significantly independent risk factor for endothelial dysfunction in men in an analysis of a large population,34 but there is little information on the relationship between uric acid and FMD in women. As a risk marker, we demonstrated that the serum uric acid level was significantly associated with FMD in women. FMD was significantly impaired with an increase in the serum uric acid level. FMD is known to be impaired as a consequence of cumulative cardiovascular risk factors.21 51 Therefore, a significant decrease in FMD according to the serum uric acid level may reflect significant associations between uric acid and other cardiovascular risk factors, including hypertension, dyslipidemia, metabolic syndrome and CKD, as a risk marker for cardiovascular disease in women. These findings provide a rationale for the relationships of uric acid with cardiovascular disease and mortality being generally strong in women.

As for the role of uric acid as a causal risk factor in endothelial function, multivariate analysis performed for the entire population indicated that uric acid was not an independent predictor of endothelial dysfunction after adjustment for other risk factors. However, when participants were divided into two groups according to menopausal status, multivariate analyses revealed that uric acid was a significantly independent risk factor for endothelial dysfunction in postmenopausal women but not in premenopausal women. The incidence of cardiovascular diseases in women is lower than that in men until around the age of menopause, after which it increases markedly and becomes equal to that of their male counterparts, suggesting a protective effect of endogenous ovarian hormones on atherosclerosis.52 Ovarian hormones, especially oestradiol, have been suggested to have protective effects on endothelial function.53 54 In addition to its protective effect on endothelial function, oestradiol has been shown to lower the uric acid level through mechanisms involving renal clearance, secretion and reabsorption.55–57 Therefore, endogenous oestradiol may preserve endothelial function and lower the serum uric acid level regardless of the presence or absence of cardiovascular risk factors, resulting in the weak relationship between uric acid and endothelial dysfunction in premenopausal women. Our findings in the

Table 4 Multivariate analysis of the relation between endothelial dysfunction and variables

| Model | Covariates | Postmenopausal OR (95% CI) | p Value | Premenopausal OR (95% CI) | p Value |
|-------|------------|---------------------------|---------|---------------------------|---------|
| 1     | Uric acid (mg/dL) | 1.27 (1.05 to 1.53) | 0.01   | 1.14 (0.90 to 1.44) | 0.28   |
| 2     | 1+age      | 1.26 (1.05 to 1.53) | 0.02   | 1.12 (0.85 to 1.39) | 0.50   |
| 3     | 2+other variables | 1.23 (1.01 to 1.50) | 0.04   | 0.98 (0.75 to 1.26) | 0.85   |

Other variables: high-density lipoprotein cholesterol, glucose and smoking.

Oestradiol has been shown to lower the uric acid level through mechanisms involving renal clearance, secretion and reabsorption.55–57 Therefore, endogenous oestradiol may preserve endothelial function and lower the serum uric acid level regardless of the presence or absence of cardiovascular risk factors, resulting in the weak relationship between uric acid and endothelial dysfunction in premenopausal women. Our findings in the
analysis of premenopausal women are consistent with the results of a previous study demonstrating that uric acid was not an independent risk factor of FMD in young women aged 30–45 years. In contrast to premenopausal women, uric acid remained a significantly independent risk for endothelial dysfunction even after adjustment for other risk factors in postmenopausal women. Although Maxwell et al reported that uric acid was not an independent determinant of FMD in women, the non-independent association between FMD and uric acid observed in their study may be, in part, attributed to inclusion of entire women in the analysis for the relationship between FMD and uric acid without taking into account menopausal status. The increased cardiovascular risk in postmenopausal women has been suggested to be associated with impairment of endothelial function after menopause. There is a possibility that treatment of hyperuricemia in postmenopausal women will improve endothelial function, leading to a decrease in cardiovascular events.

There were some limitations in this study. The cross-sectional design did not allow us to establish a definitive causal relationship between hyperuricemia and endothelial dysfunction. In addition, multiple factors, such as vitamins, phosphate and statin use, that are known to influence FMD and residual unrecognised confounding factors were not taken into account in this study. Future prospective and interventional studies are certainly warranted to obtain more specific conclusions as to whether hyperuricemia should be treated, which subgroup should be treated and whether treatment of hyperuricemia improves endothelial function with a subsequent decrease in cardiovascular events.

In conclusion, uric acid is a useful marker for endothelial function in women. Significant relationships between the serum uric acid level and cardiovascular risk factors, including hypertension, dyslipidemia, metabolic syndrome and CKD, may result in a significant association between FMD and the serum uric acid level as a risk marker of atherosclerotic diseases. Uric acid, as a causal cardiovascular risk factor, may be an independent risk for endothelial dysfunction in postmenopausal women but not in premenopausal women. Further studies are needed to investigate whether treatment for hyperuricemia improves endothelial function in postmenopausal women.

Author affiliations
1Department of Cardiovascular Medicine, Hiroshima University Graduate School of Biomedical Sciences, Hiroshima, Japan
2Division of Reproduction and Medicine, Hiroshima University Hospital, Hiroshima, Japan
3Department of Gastroenterology and Metabolism, Institute of Biomedical and Health Sciences, Graduate School of Biomedical Sciences, Hiroshima University, Hiroshima, Japan
4Hiroshima International University, Hiroshima, Japan
5Department of Cardiovascular Regeneration and Medicine, Research Institute for Radiation Biology and Medicine, Hiroshima University, Hiroshima, Japan
6The Second Department of Internal Medicine, Tokyo Medical University, Tokyo, Japan
7Division of Biomedical Engineering, National Defense Medical College Research Institute, Tokorozawa, Japan

Acknowledgements The authors would like to thank Megumi Wakisaka, Miki Kumiji, Kichihiro Kawano and Satoko Michiyama for their excellent secretarial assistance.

Contributors TM and YH were involved in the drafting of the article and conception of the study. JS, NF, NI, SM, YL, MK, TM, TH, CG, KN, AN, BT and HT were involved in performing the ultrasoundography. KC, YK and AY were involved in revising the manuscript for important intellectual content. All the authors approved the final version of the article to be published.

Funding This study was supported in part by a Grant-in-Aid for Scientific Research from the Ministry of Education, Science and Culture of Japan (1859081500 and 21590898) and a Grant in Aid of Japanese Arteriosclerosis Prevention Fund.

Competing interests None.

Patient consent Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement No additional data are available.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 3.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/3.0/

REFERENCES
1. Reunanen A, Takkunen H, Knelt P, et al. Hyperuricemia as a risk factor for cardiovascular mortality. Acta Med Scand Suppl 1982;688:49–59.
2. Levine W, Dyer AR, Shekelle RB, et al. Serum uric acid and 11.5-year mortality of middle-aged women: findings of the Chicago Heart Association Detection Project in Industry. J Clin Epidemiol 1989;42:257–67.
3. Wannamethee SG, Shaper AG, Whincup PH. Serum urate and the risk of major coronary heart disease events. Heart 1997;78:147–53.
4. Lehto S, Niskanen L, Ronnemaa T, et al. Serum uric acid is a strong predictor of stroke in patients with non-insulin-dependent diabetes mellitus. Stroke 1998;29:635–9.
5. Alderman MH, Cohen H, Madhavan S, et al. Serum uric acid and cardiovascular events in successfully treated hypertensive patients. Hypertension 1999;34:144–50.
6. Culleton BF, Larson MG, Kannel WB, et al. Serum uric acid and risk for cardiovascular disease and death: the Framingham Heart Study. Ann Intern Med 1999;131:7–13.
7. Moriaty JT, Folsom AR, Iribarren C, et al. Serum uric acid and risk of coronary heart disease: Atherosclerosis Risk in Communities (ARIC) Study. Ann Epidemiol 2000;10:136–43.
8. Verdecchia P, Schiacci G, Reboldi G, et al. Relation between serum uric acid and risk of cardiovascular disease in essential hypertension. The PIUMA Study. Hypertension 2000;36:1072–8.
9. Fang J, Alderman MH. Serum uric acid and cardiovascular mortality the NHANES I epidemiologic follow-up study, 1971–1992. National Health and Nutrition Examination Survey, JAMA 2000;283:2404–10.
10. Ioachimescu AG, Brennan DM, Hoar BM, et al. Serum uric acid is an independent predictor of all-cause mortality in patients at high risk of cardiovascular disease: a Preventive Cardiology Information System (PreCIS) Database Cohort Study. Arthritis Rheum 2008;58:823–30.
11. Cannon PJ, Stason WB, Demartini FE, et al. Hyperuricemia in primary and renal hypertension. N Engl J Med 1996;275:457–464.
12. Ford ES, Li C, Cook S, et al. Serum concentrations of uric acid and the metabolic syndrome among US children and adolescents. Circulation 2007;115:2526–32.
13. Siu YP, Leung KT, Tong MK, et al. Use of allopurinol in slowing the progression of renal disease through its ability to lower serum uric acid level. Am J Kidney Dis 2006;47:51–9.
14. Feig DI, Kang DJ, Johnson RJ. Uric acid and cardiovascular risk. N Engl J Med 2006;355:1811–15.
15. Gagliardi AC, Miname MH, Santos RD. Uric acid: a marker of increased cardiovascular risk. Atherosclerosis 2009;202:11–17.
16. Khosla UM, Zharikov S, Finch JL, et al. Hyperuricemia induces endothelial dysfunction. *Kidney Int* 2005;67:1738–42.
17. Kang DH, Park SK, Lee IK, et al. Uric acid-induced C-reactive protein expression: implication on cell proliferation and nitric oxide production of human vascular cells. *J Am Soc Nephrol* 2005;16:3535–62.
18. Yu MA, Sanchez-Lozada LG, Johnson RJ, et al. Oxidative stress with an activation of the renin-angiotensin system in human vascular endothelial cells as a novel mechanism of uric acid-induced endothelial dysfunction. *J Hypertens* 2002;20:242–9.
19. Ross R. Atherosclerosis—an inflammatory disease. *N Engl J Med* 1999;340:115–26.
20. Celermajer DS, Sorensen KE, Gooch VM, et al. Non-invasive detection of endothelial dysfunction in children and adults at risk of coronary artery disease. *Lancet* 1992;340:1111–15.
21. Benjamin EJ, Larson MG, Keyes MJ, et al. Clinical correlates and heritability of flow-mediated dilation in the community: the Framingham Heart Study. *Circulation* 2004;109:613–19.
22. Idei N, Nishioka K, Soga J, et al. Uric acid and cardiovascular risk factors. *Arterioscler Thromb Vasc Biol* 2011;31:2535–9.
23. Soga N, Kama K, Hata T, et al. Rho-associated kinase activity, endothelial function, and cardiovascular risk factors. *Arterioscler Thromb Vasc Biol* 2011;31:2535–9.
24. Fuguiren N, Kama K, Hata T, et al. Mineralocorticoid receptor blocker eplerenone improves endothelial function and inhibits Rho-associated kinase activity in patients with hypertension. *Clin Pharmacol Ther* 2012;91:289–97.
25. Modena MG, Bonetti L, Copp F, et al. Prognostic role of reversible endothelial dysfunction in hypertensive postmenopausal women. *J Am Coll Cardiol* 2002;40:505–10.
26. Gokce N, Keaney JF Jr, Hunter LM, et al. Risk stratification for postoperative cardiovascular events via noninvasive assessment of endothelial function: a prospective study. *Circulation* 2002;106:1059–67.
27. Breveiti G, Silvestro A, Schiano V, et al. Endothelial dysfunction and cardiovascular risk prediction in peripheral arterial disease: additive value of flow-mediated dilation to ankle-brachial pressure index. *Circulation* 2003;108:2092–3.
28. Lerman A, Zeitler AM. Endothelial function: cardiac events. *Circulation* 2005;111:363–8.
29. Maxwell AJ, Bruinsma KA. Uric acid is closely linked to vascular nitric oxide activity. Evidence for mechanism of association with cardiovascular disease. *J Am Coll Cardiol* 2001;38:1850–8.
30. Mercuro G, Vitale C, Cercutani E, et al. Effect of hyperuricemia upon endothelial function in patients at increased cardiovascular risk. *Am J Cardiol* 2004;94:932–5.
31. Kato M, Hisatome I, Tomikura Y, et al. Menopause is associated with endothelial dysfunction in women. *J Hypertens* 1999;17(Suppl 1):S1–4.
32. Wilson PW, Castelli WP, Kannel WB. Coronary risk prediction in adults (the Framingham Heart Study). *Am J Cardiol* 1987;59:91G–6G.