

Impact of obesity on incident hypertension independent of weight gain among nonhypertensive Japanese: the Ibaraki Prefectural Health Study (IPHS)

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1 **Title page**

2 Title

3 Impact of Obesity on Incident Hypertension Independent of Weight Gain among
4 Non-hypertensive Japanese: The Ibaraki Prefectural Health Study (IPHS)

5

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29

30 **Abstract**

31 Objective: The aim of this study was to examine the association between body mass index
32 (BMI) and risk of incident hypertension among Japanese men and women who are
33 middle-aged and older.

34 Design: Prospective, population-based cohort study.

35 Subjects: A total of 68 205 non-hypertensive adults (18 336 men and 49 869 women) aged
36 40-79 years who completed health check-ups in the Ibaraki prefecture, Japan, in 1993 were
37 followed-up through 2006. To exclude the impact of BMI change during the follow-up period,
38 a time-dependent covariate Cox proportional hazards model was used to compute the hazard
39 ratios (HRs) of incident hypertension according to BMI categories. Incident hypertension was
40 defined as a systolic blood pressure (BP) of ≥ 140 mmHg, a diastolic BP of ≥ 90 mmHg,
41 and/or hypertensive medication use.

42 Results: A total of 30 982 adults (45.4%) developed hypertension (9 331 men and 21 651
43 women) during a mean of 3.9 years of follow-up. Compared with a BMI of < 19.0 ,
44 time-dependent covariates adjusted HRs (95% confidence interval [CI]) for hypertension
45 among participants with a BMI of ≥ 25.0 were 1.42 (1.17-1.73) for men aged 40 to 59 years,
46 1.34 (1.19-1.51) for men aged 60 to 79 years, 1.47 (1.33-1.62) for women aged 40 to 59 years,
47 and 1.29 (1.18-1.41) for women aged 60 to 79 years.

48 Conclusion: The baseline BMI is associated with future risk for incident hypertension even
49 after accounting for weight change during the follow-up period. Weight loss may be
50 recommended to non-hypertensive obese adults to prevent the development of hypertension.

51

52 **Introduction**

53 Hypertension is a major cause of cardiovascular disease [1]. Currently, approximately 45% of
54 Japanese adults are considered hypertensive [2], defined as systolic blood pressure (SBP) \geq
55 140 mmHg, diastolic blood pressure (DBP) \geq 90 mmHg or treatment with an
56 antihypertensive drug. Many prospective cohort studies have consistently shown that the
57 higher the BMI, the greater the likelihood of developing hypertension [3-11]. Some
58 prospective cohort studies have reported that long-term weight gain was associated with a
59 risk of hypertension among hypotensive or high-normal blood pressure patients [12-15].
60 Moreover, many intervention studies have also shown that weight loss reduced blood
61 pressure [16-18]. These results could lead to a hypothesis that the association between
62 baseline BMI and the risk of hypertension are due to weight gain during the follow-up period.
63 If this hypothesis were true, non-hypertensive obese individuals would not be required to lose
64 weight to prevent incident hypertension. To clarify whether weight loss should be
65 recommended for non-hypertensive obese people to prevent incident hypertension, a large
66 cohort study that accounts for weight change is warranted. The purpose of this study was to
67 clarify the association between long-term weight stability and the risk of incident
68 hypertension in a large community-based cohort of men and women who are middle-aged
69 and older.

70

71 **Methods**

72 **Study population**

73 The study population comprised 194 333 individuals (63 865 men and 130 468 women) aged
74 40-79 years living in the Ibaraki Prefecture, Japan. The population participated in
75 community-based annual health checkups beginning in 1993 (Ibaraki Prefectural Health
76 Study). These health checkups were conducted by local governments in accordance with the
77 Japan Health Laws. Data were collected by the Ibaraki prefectural government from local
78 governments after depersonalizing them to ensure anonymity. We excluded 3 080 individuals
79 (738 men and 2 342 women) with incomplete data and 94 153 individuals (35 647 men and
80 58 506 women) with a SBP of ≥ 140 mmHg and/or a DBP of ≥ 90 mmHg or antihypertensive
81 medication use. We further excluded 28 895 individuals (9 144 men and 19 751 women) who
82 did not participate in the 1994 survey, thereby ensuring that the participants were followed
83 for at least one year.

84 Ultimately, the study subjects consisted of 68 205 individuals (18 336 men and 49 869
85 women). These participants were followed up by annual examinations until hypertension had
86 been diagnosed or until the end of 2006. Blood pressure was measured at annual follow-up
87 examinations. Participants who did not undergo annual checkups during the follow-up
88 periods were censored on the date of their latest checkup. The protocol for this cohort study
89 was approved by the Ibaraki Epidemiology Study Union Ethics Review Committee.

90

91 **Assessment of body mass index, blood pressure, and other covariates**

92 Height in stockinged feet and weight in light clothing were measured at baseline. BMI was
93 calculated as weight in kilograms divided by the height in meters squared. BP was measured
94 from the right arm of seated participants who had rested for more than 5 minutes by trained
95 observers using standard mercury sphygmomanometers during examinations that occurred

96 between 1993 and 2004 and by an automated sphygmomanometer between 2005 and 2006.
97 When the SBP was greater than 150 mmHg or DBP was greater than 90 mmHg, a second
98 measurement was performed after several deep breaths, and the lower values, which were
99 almost always observed after the second measurement, were used for analyses.

100 Blood samples were drawn from seated participants into two polyethylene terephthalate
101 tubes: one with an accelerator and the other with sodium fluoride and
102 ethylenediaminetetraacetic acid. Overnight fasting (≥ 8 hours) was not required. The blood
103 glucose level was measured by the glucose oxidase electrode method with a GA1140 device
104 (Kyoto Daiichi Kagaku, Kyoto, Japan) during examinations that occurred between 1993 and
105 1996, by the enzyme method with a H7170 device (Hitachi, Tokyo, Japan) between 1997 and
106 2003, and with a H7700 device (Hitachi, Tokyo, Japan) between 1994 and 2006. Participants
107 were considered diabetic if they had a fasting plasma glucose level of at least 6.1 mmol/L or a
108 non-fasting glucose level of at least 7.8 mmol/L, or if they were being treated for diabetes
109 mellitus. Serum total cholesterol and serum triglyceride levels were measured by the enzyme
110 method with a RX-30 device (Nihon Denshi, Tokyo, Japan) between 1993 and 1995, with a
111 H7350 device (Hitachi, Tokyo, Japan) between 1996 and 2003, and with a H7700 device
112 between 2004 and 2006. High-density lipoprotein (HDL) cholesterol levels were measured by
113 the phosphotungstic acid magnesium method with a MTP-32 device (Corona Electric, Ibaraki,
114 Japan) between 1993 and 1995, by the selective inhibition method with a H7350 device
115 between 1996 and 2003, and with a H7700 device between 2004 and 2006. The laboratory
116 participated in external standardization and successfully met the criteria for precision
117 accuracy for the measurement of blood samples by the Japan Medical Association, the
118 Japanese Association of Medical Technologists, and the Japan Society of Health Evaluation
119 and Promotion. An interview was conducted to ascertain medical history, smoking status
120 (never, ex-, current < 20 cigarettes/day, and ≥ 20 cigarettes/day), and alcohol intake (non-,

121 sometimes, < 66 g/day, and \geq 66 g/day).

122

123 **Endpoint determination**

124 Incident hypertension was defined as SBP of \geq 140 mmHg, DBP of \geq 90 mmHg, and/or when
125 treatment for hypertension was initiated.

126

127 **Statistical analysis**

128 Participants were classified into the following 7 categories with regard to their BMI: < 19.0,
129 19.0-20.9, 21.0-22.9, 23.0-24.9, 25.0-26.9, 27.0-29.9, or \geq 30.0 kg/m². Baseline participant
130 characteristics were compared according to BMI categories using chi-squared tests for
131 categorical variables and analysis of variance for continuous variables. Hazard ratios (HRs)
132 with 95% confidence intervals (CIs) for incident hypertension relative to BMI categories
133 were calculated with a reference of < 19.0 kg/m² using a Cox proportional hazards regression
134 model [19]. The analyses were stratified by sex and age groups (40-59 and 60-79 years).

135 Three multivariable-adjusted models were used. In Model 1, covariates included age
136 (years), SBP (mmHg), DBP (mmHg), fasting status (yes or no), total cholesterol level
137 (mmol/L), HDL cholesterol level (mmol/L), log-transformed triglyceride level (mmol/L),
138 lipid medication use (yes or no), blood glucose status (normal: < 6.1 mmol/L during fasting
139 or < 7.8 mmol/L during non-fasting; border: 6.1-7.0 mmol/L during fasting or 7.8-11.1
140 mmol/L during non-fasting; hyperglycemia: \geq 7.0 mmol/L during fasting or \geq 11.1 mmol/L
141 during non-fasting), smoking status (never, ex-, current < 20 cigarettes/day, or \geq 20
142 cigarettes/day), and alcohol intake (non-, sometimes, < 66 g/day, or \geq 66 g/day). In Model 2,
143 the changes in BMI, total cholesterol level (mmol/L), HDL cholesterol level (mmol/L), and
144 log-transformed triglyceride level (mmol/L) from baseline to the end of follow-up were
145 added as covariates to Model 1. In Model 3, time-dependent variables of BMI and covariates

146 in Model 1 were used with a time-dependent Cox hazard model, which was used for adjusting
147 the change in the variable from the baseline to the final year of follow-up [20]. For the
148 secondary analysis, we excluded 34 190 participants with incomplete data and 16,370
149 participants with a SBP of ≥ 140 mmHg and/or a DBP of ≥ 90 mmHg or antihypertensive
150 medication use at a 5-year follow-up. The remaining 17 645 qualified participants were
151 classified into the following 4 categories with regard to their baseline BMI and BMI 5 years
152 later: (a) BMI of less than 25 kg/m^2 at baseline and BMI of less than 25 kg/m^2 at a 5-year
153 follow-up (-/-); (b) BMI of 25 kg/m^2 or more at baseline and BMI of less than 25 kg/m^2 at a
154 5-year follow-up (+/-); (c) BMI of less than 25 kg/m^2 at baseline and BMI of 25 kg/m^2 or
155 more at a 5-year follow-up (-/+); and (d) BMI of 25 kg/m^2 or more at baseline and BMI of 25
156 kg/m^2 or more at a 5-year follow-up (+/+). We calculated HRs for incident hypertension to
157 consider long-term changes of their weights.

158 All statistical analyses were conducted using SAS, version 9.1 (SAS Institute, Inc., Cary,
159 North Carolina).

160

161 **Results**

162 Sex-stratified baseline characteristics according to BMI categories are shown in Table 1. All
163 covariates, except for blood glucose status, diabetic medication use and smoking status in
164 men and lipid medication use in women, were associated with BMI in both sexes. A higher
165 BMI was linked with higher SBP and DBP at baseline in both sexes.

166 Of the 68 205 adults (18 336 men and 49 869 women), 30 982 (45.4%) developed
167 hypertension (9 331 men and 21 651 women) during a mean follow-up of 3.9 years (3.4 years
168 for men and 4.1 years for women). Table 2 shows sex- and age-stratified HRs for
169 hypertension according to the baseline BMI categories. In men aged 40-59 years, compared
170 with a baseline BMI < 19.0 kg/m², the multivariate HRs of BMI ≥ 25.0 kg/m² was
171 significantly higher in Model 1 and Model 2. In men aged 60-79 years, compared with a
172 baseline BMI < 19.0 kg/m², the multivariate HRs of BMI ≥ 19.0 kg/m² was significantly
173 higher in Model 1 and the multivariate HRs of BMI ≥ 23.0 kg/m² was significantly higher in
174 Model 2. In both age groups among men, time-dependent covariates adjustment HRs (BMI ≥
175 25.0 kg/m²: 1.42, ≥ 30.0 kg/m²: 1.77 in aged 40-59 years; BMI ≥ 25.0 kg/m²: 1.34, ≥ 30.0
176 kg/m²: 1.68 in aged 60-79 years) showed a trend similar to Model 1 and Model 2.

177 In women aged 40-59 years, compared with a baseline BMI < 19.0 kg/m², the multivariate
178 HRs of BMI ≥ 21.0 kg/m² was significantly higher in Model 1 and of BMI ≥ 23.0 kg/m² was
179 significantly higher in Model 2.

180 In women aged 60-79 years, compared with a baseline BMI < 19.0 kg/m², the multivariate
181 HRs of BMI ≥ 21.0 kg/m² was significantly higher in Model 1 and 25.0-26.9 kg/m² of BMI
182 was significantly higher in Model 2. In both age groups among women, time-dependent
183 covariates adjustment HRs (BMI ≥ 25.0 kg/m²: 1.53, ≥ 30.0 kg/m²: 2.07 in age 40-59 years;
184 BMI ≥ 25.0 kg/m²: 1.47, ≥ 30.0 kg/m²: 1.91 in age 60-79 years) showed a trend similar to
185 Model 1 and Model 2.

186 Figure 1 represents HRs for hypertension according to the BMI categories at baseline and
187 at a 5-year follow-up. Among all non-hypertensive participants at baseline and at a 5-year
188 follow-up, compared with the -/- group, the age-adjusted HRs (95% CI) of +/-, -/+, and +/+
189 were 1.12 (1.03-1.41), 1.69 (1.47-1.94), and 1.67 (1.53-1.82), respectively. The multivariate
190 HRs of +/-, -/+, and +/+ were 1.12 (1.03-1.41), 1.69 (1.47-1.94), and 1.67 (1.53-1.82),
191 respectively.

192 **Discussion**

193 To the best of our knowledge, the results of the present large prospective study showed that
194 stable obesity in non-hypertensive individuals was significantly associated with an increased
195 risk of incident hypertension among men and women who are middle-aged and older.

196 Previous studies that examined the relationship between BMI and hypertension reported
197 results consistent with the present study. Many cross-sectional studies showed a
198 dose-dependent relationship between BMI and the prevalence of hypertension in Caucasian
199 [5, 21] and Asian [3, 13, 22] populations. Kawada [3] reported that the odds ratios for
200 hypertension against the lowest quartile of BMI (separated by 20, 25 and 28) were 1.7, 3.6,
201 and 7.2, respectively, with 95% CIs of 1.2-2.3, 2.5-5.2 and 4.7-11.0, respectively, among
202 middle-aged men and women (5 346 men and 11 525 women, aged 40-59). Prospective
203 studies [4, 12, 23] also showed similar results to the present study. The Physicians' Health
204 Study (13 563 healthy and non-hypertensive men in U.S.) [4] showed that the HRs (95% CIs)
205 of developing hypertension for men with a BMI of 22.4-23.6, 23.7-24.7, 24.8-26.4, and >26.4
206 kg/m² were 1.20 (1.09-1.32), 1.31 (1.19-1.44), 1.56 (1.42-1.72), and 1.85 (1.69-2.03),
207 respectively (*P* for the trend, < 0.001), compared with participants who had the lowest BMI
208 quintile (< 22.4 kg/m²). A cohort study of U.S. female nurses (82 473 women, aged 30-55)
209 [12] showed that the multivariate HRs (95% CIs) of developing hypertension for women with
210 a BMI of 22.0-22.9, 24.0-24.9, 26.0-27.9, and > 31.0 kg/m² were 1.57 (1.44-1.72), 2.15
211 (1.97-2.35), 3.33 (3.06-3.62), and 6.31 (5.80-6.87), respectively (*P* for the trend, < 0.001),
212 compared with women who had the lowest BMI decile (< 20.0 kg/m²), and an increase in
213 BMI of 1 kg/m² was associated with a 12 % increased risk for incident hypertension. These
214 results suggested a strong association between higher BMI and increased risk of incident
215 hypertension in men and women.

216 We also examined the effect of 5-year changes in BMI category (BMI less than, or equal or

217 more than 25 kg/m²) on incident hypertension. The HRs of non-obese people at a 5-years
218 follow-up, despite obesity at baseline, was not significant compared with non-obese people at
219 both baseline and at a 5-years follow-up. In contrast, the HRs of obese individuals at a 5-year
220 follow-up, despite non-obesity at baseline, was significantly higher compared with non-obese
221 people at both baseline and at a 5-years follow-up. The effect of change in BMI on blood
222 pressure has already been examined in some studies [12-18]. A systematic review [24] of
223 lifestyle intervention studies reported differences ranging from -11 to + 4 kg for weight, -7 to
224 + 2.2 mmHg for DBP and -13 to + 6.1 mmHg for SBP. It is suggested that SBP differences
225 were equal to a ratio of 1 kg to every 1 mmHg. These studies suggest the efficacy of weight
226 loss for the hypertensive obese population. The present study further represented the efficacy
227 of weight loss for the non-hypertensive obese population to prevent incident hypertension.
228 Williams [25] examined the relationship between long-term stable individuals' BMI (change
229 less than ± 0.4 kg/m²) and the cumulative incidence of hypertension and showed that the odds
230 ratio of each kg/m² increment in baseline BMI was 1.19 (95% CIs: 1.14-1.24) in men and
231 1.11 (95% CIs: 1.02-1.20) in women. Matsuo et al [26] reported in a cohort study of 5 201
232 Japanese middle-aged male employees using of a time-dependent Cox proportional hazard
233 model for analysis that the HRs (95% CIs) of stable BMI of 25.0-26.9, 27.0-29.9, and ≥ 30.0
234 kg/m² were 1.11 (0.93-1.32), 1.34 (1.09-1.65) and 1.49 (1.09-2.04), respectively, compared
235 with men who had a BMI of 23.0-24.9 kg/m². These results are consistent with our results,
236 although Williams calculated the BMI with a self-reported weight and cumulative incidence
237 of hypertension at 2 points (baseline and 7 years later), and Matsuo et al. studied middle-aged
238 male employees.

239 The exact mechanism by which stable adiposity raises BP is uncertain. However, a likely
240 explanation is that obesity-associated hypertension appears to be due to inadequate
241 vasodilatation in the setting of increased blood volume and cardiac output. A defect in control

242 of vascular resistance has been attributed to enhanced activity of the sympathetic nervous
243 system, abnormal rennin-angiotensin-aldosterone relations, and insulin resistance [27-31].
244 The peripheral vascular resistance is increased by adipocytokines (e.g., leptin, TNF- α , RBP4)
245 [32-34].

246 The study has several strengths. First, the effect of weight change during the follow-up
247 period was taken into account to investigate the association between long-term weight
248 stability and future incident risk for hypertension, while many previous epidemiological
249 studies did not account for these factors. Second, the present study had a large cohort for
250 which gender, age and BMI stratification analysis was possible. In addition, all blood samples
251 in each year were measured by the same device, reagents, and quality control programs [35].

252 However, there are several limitations. First, the participation rate for the follow-up
253 examinations was approximately 40%, but the mean SBP and DBP did not differ between
254 individuals who were and were not followed up. In this context, the potential selection bias
255 may have been small. Second, the instrument used to measure the blood pressure was
256 switched from the standard mercury sphygmomanometer to an automatic
257 sphygmomanometer following 2005. The BP determined by means of an automated
258 sphygmomanometer method was compared with those determined by a standard mercury
259 sphygmomanometer method, using 18 859 samples (non-medication participants) in 2004
260 and 2005. Comparability between BP measured using the 2 sphygmomanometers was
261 acceptable (SBP: $r = 0.69$, DBP: $r = 0.55$, $P < 0.05$, respectively). Third, we could not assess
262 the potential confounding variables such as fat distribution, physical activity, nutritional
263 status (e.g., sodium intake), and family history of hypertension.

264 In conclusion, the baseline BMI is associated with a future incident risk for hypertension,
265 even after accounting for weight change during the follow-up period. Weight loss may be
266 recommended for obese non-hypertensive individuals to prevention incident hypertension

267 among men and women who are middle-aged and older.

268

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270 We thank all the participants in this study.

271

272 **Conflict of interest**

273 The authors declare no conflict of interest.

274

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371

372 Figure legend

373 Figure 1

374 Abbreviations: HRs, hazard ratios; CIs, confidence intervals. Age-adjusted HRs with 95%
375 confidence intervals were adjusted for age (years) in 1998 and sex. Multivariate HRs with
376 95% confidence intervals were adjusted for possible confounders measured in 1998: age
377 (years), sex, systolic blood pressure (mmHg), diastolic blood pressure (mmHg), fasting status
378 (yes or no), total cholesterol level (mmol/liter), high density lipoprotein cholesterol level
379 (mmol/liter), log-transformed triglyceride level (mmol/liter), lipid medication use (yes or no),
380 blood glucose status (normal, border, hyperglycemia), diabetic medication use (yes or no),
381 smoking status (never, ex-, < 20 cigarettes/day, or \geq 20 cigarettes/day), and alcohol intake
382 (never, sometimes, < 66 g/day, or \geq 66 g/day). Bold values showed statistical significance
383 ($P < 0.05$).

Table 1. Baseline characteristics of participants by body mass index (BMI) categories

Gender and baseline variables	Body mass index, kg/m ²							P-value
	< 19.0	19.0 - 20.9	21.0 - 22.9	23.0 - 24.9	25.0 - 26.9	27.0 - 29.9	> 30.0	
Men (n = 18 336)								
No. of subjects	1 545	3 476	4 934	4 544	2 598	1 085	154	
Age, years	63.5 (9.5)	60.5 (10.2)	58.8 (10.3)	57.7 (10.2)	56.5 (10.0)	56.6 (9.8)	55.3 (10.0)	<.001
Change in body mass index, kg/m ²	0.0 (1.1)	0.1 (1.2)	0.0 (1.2)	-0.1 (1.3)	-0.2 (1.3)	-0.4 (1.5)	-0.7 (1.5)	<.001
Systolic blood pressure, mmHg	119.9 (11.4)	121.7 (10.8)	122.7 (10.2)	124.0 (9.5)	124.4 (9.4)	125.4 (9.2)	126.2 (8.8)	<.001
Diastolic blood pressure, mmHg	71.9 (8.4)	73.1 (8.0)	74.1 (7.9)	75.4 (7.3)	75.9 (7.4)	76.6 (7.1)	78.1 (7.8)	<.001
Total cholesterol, mmol/liter	4.7 (0.8)	4.8 (0.8)	5.0 (0.8)	5.1 (0.9)	5.2 (0.8)	5.2 (0.9)	5.2 (0.9)	<.001
HDL cholesterol, mmol/liter	1.6 (0.4)	1.5 (0.4)	1.4 (0.4)	1.3 (0.3)	1.2 (0.3)	1.1 (0.3)	1.1 (0.3)	<.001
Triacylglycerol, mmol/liter	1.1 (0.5)	1.2 (0.7)	1.5 (0.8)	1.8 (1.0)	2.0 (1.2)	2.2 (1.3)	2.2 (1.0)	<.001
Lipid medication use, %	3.2	2.8	2.6	3.1	3.2	3.1	2.6	<.001
Diabetic medication use, %	0.4	0.8	1.0	1.3	1.2	1.4	2.6	0.177
Blood glucose status, %								0.866
Normal	81.9	83.9	84.3	84.6	81.9	82.0	83.8	
Border	13.6	12.4	12.0	11.4	13.5	12.7	9.7	
Hyperglycemia	4.5	3.7	3.7	4.0	4.5	5.3	6.5	
Smoking status, %								0.116
Never	19.4	20.6	22.4	25.1	24.1	23.3	25.3	
Ex	23.4	23.3	27.0	28.1	29.5	33.9	27.9	
Current								
< 20 cigarettes/day	22.8	18.7	15.3	12.3	11.5	9.8	9.7	
≥ 20 cigarettes/day	34.4	37.4	35.3	34.4	34.9	33.0	37.0	
Alcohol intake, %								<.001
Never	48.5	40.6	37.2	36.6	37.6	38.2	51.3	
Sometimes	11.5	13.2	13.6	15.1	15.9	18.8	17.5	
Everyday								
< 66 g/day	36.6	41.9	44.0	43.4	41.4	37.1	25.3	
≥ 66 g/day	3.4	4.3	5.1	5.0	5.1	5.8	5.8	

Table 1. Baseline characteristics of participants by body mass index (BMI) categories (continued)

Gender and baseline variables	Body mass index, kg/m ²							P-value
	< 19.0	19.0 - 20.9	21.0 - 22.9	23.0 - 24.9	25.0 - 26.9	27.0 - 29.9	> 30.0	
Women (n = 49 869)								
No. of subjects	3 796	9 391	13 824	11 887	6 781	3 434	756	
Age, years	56.8 (10.9)	54.1 (10.1)	54.4 (9.7)	55.1 (9.5)	56.0 (9.4)	56.1 (9.4)	55.4 (9.2)	<.001
Change in body mass index, kg/m ²	0.2 (1.2)	0.1 (1.2)	0.0 (1.3)	-0.1 (1.4)	-0.2 (1.5)	-0.3 (1.6)	-0.6 (2.0)	<.001
Systolic blood pressure, mmHg	117.6 (11.9)	118.5 (11.5)	120.0 (11.1)	121.3 (10.7)	122.7 (10.2)	123.8 (9.8)	124.7 (8.9)	<.001
Diastolic blood pressure, mmHg	70.0 (8.5)	70.9 (8.3)	72.1 (8.1)	73.3 (7.9)	74.3 (7.6)	75.3 (7.5)	76.7 (7.4)	<.001
Total cholesterol, mmol/liter	5.1 (0.9)	5.2 (0.9)	5.3 (0.9)	5.4 (0.9)	5.4 (0.9)	5.5 (0.9)	5.5 (0.9)	<.001
HDL cholesterol, mmol/liter	1.7 (0.4)	1.6 (0.4)	1.5 (0.4)	1.4 (0.3)	1.4 (0.3)	1.3 (0.3)	1.3 (0.3)	<.001
Triacylglycerol, mmol/liter	1.0 (0.5)	1.2 (0.6)	1.3 (0.7)	1.5 (0.9)	1.6 (0.9)	1.8 (1.0)	1.9 (1.0)	<.001
Lipid medication use, %	1.7	1.5	2.3	2.9	3.3	3.1	3.2	0.367
Diabetic medication use, %	0.9	1.0	1.5	1.2	1.6	1.8	2.2	<.001
Blood glucose status, %								<.001
Normal	91.1	92.5	92.6	91.8	90.0	89.1	87.0	
Border	7.5	6.1	6.1	6.3	8.0	8.4	9.7	
Hyperglycemia	1.3	1.4	1.3	1.9	1.9	2.6	3.3	
Smoking status, %								<.001
Never	92.6	94.7	94.7	95.2	94.9	94.5	92.2	
Ex	0.6	0.5	0.7	0.5	0.6	0.9	0.9	
Current								
< 20 cigarettes/day	4.7	3.4	3.2	2.8	2.9	3.0	3.6	
≥ 20 cigarettes/day	2.1	1.5	1.5	1.5	1.6	1.6	3.3	
Alcohol intake, %								<.001
Never	90.9	88.7	89.0	89.6	90.2	90.7	90.7	
Sometimes	5.2	6.9	7.2	6.8	6.5	6.0	5.4	
Everyday								
< 66 g/day	3.9	4.2	3.7	3.5	3.3	3.1	3.4	
≥ 66 g/day	-	0.1	0.1	0.1	0.0	0.2	0.4	

HDL = high-density lipoprotein. The means (SD) are shown for continuous variables: age, fasting and non-fasting blood glucose, systolic and diastolic blood pressure, total cholesterol, HDL cholesterol, and triglycerides. SI conversion factors: blood glucose values were converted to mmol/L by multiply by 0.05551; cholesterol values were converted to mmol/L by multiply by 0.02586; triglycerides values were converted to mmol/L by multiply by 0.01129.

Table 2. The effect of hazard ratios on hypertension within each of seven BMI categories

	Baseline BMI categories (kg/cm ²)						
	< 19.0	19.0 - 20.9	21.0 - 22.9	23.0 - 24.9	25.0 - 26.9	27.0 - 29.9	> 30.0
Men, age 40-59 years (n = 8 540)							
No. of person-years	1 590	5 713	9 042	8 267	4 964	1 952	285
No. of hypertension cases	127	434	882	1 048	740	351	61
Incidence rates per 1 000 person-years	80	76	98	127	149	180	214
Age-adjusted HRs (95% CIs)	1.00	1.01 (0.83-1.23)	1.26 (1.05-1.52)	1.57 (1.31-1.89)	1.81 (1.50-2.19)	2.12 (1.73-2.60)	2.48 (1.83-3.36)
Model 1† HRs (95% CIs)	1.00	0.90 (0.74-1.09)	1.05 (0.87-1.27)	1.19 (0.98-1.44)	1.36 (1.16-1.66)	1.52 (1.23-1.89)	1.75 (1.28-2.40)
Model 2‡ HRs (95% CIs)	1.00	1.01 (0.59-1.73)	0.97 (0.58-1.65)	1.10 (0.64-1.88)	1.82 (1.04-3.16)	1.83 (1.00-3.35)	2.37 (0.91-6.17)
Model 3§ HRs (95% CIs)	1.00	0.95 (0.78-1.15)	1.13 (0.94-1.36)	1.24 (1.03-1.50)	1.42 (1.17-1.73)	1.60 (1.30-1.97)	1.77 (1.29-2.41)
Men, age 60-79 years (n = 9 796)							
No. of person-years	3 831	6 982	8 334	6 582	2 937	1 075	101
No. of hypertension cases	517	1 177	1 566	1 366	710	314	38
Incidence rates per 1 000 person-years	135	169	188	208	242	292	376
Age-adjusted HRs (95% CIs)	1.00	1.25 (1.13-1.38)	1.38 (1.25-1.52)	1.51 (1.36-1.67)	1.67 (1.49-1.87)	1.94 (1.69-2.24)	2.24 (1.61-3.11)
Model 1† HRs (95% CIs)	1.00	1.15 (1.03-1.27)	1.20 (1.09-1.33)	1.26 (1.13-1.41)	1.35 (1.20-1.53)	1.54 (1.32-1.78)	1.68 (1.20-2.35)
Model 2‡ HRs (95% CIs)	1.00	1.01 (0.71-1.44)	1.24 (0.88-1.75)	1.48 (1.03-2.13)	1.52 (0.98-2.37)	2.01 (1.11-3.64)	N/A
Model 3§ HRs (95% CIs)	1.00	1.11 (1.00-1.22)	1.17 (1.06-1.29)	1.26 (1.14-1.40)	1.34 (1.19-1.51)	1.48 (1.28-1.72)	1.68 (1.21-2.35)
Women, age 40-59 years (n = 32 133)							
No. of person-years	10 877	32 781	45 716	34 389	16 754	7 410	1 402
No. of hypertension cases	496	1 682	2 975	2 969	1 940	1 001	265
Incidence rates per 1 000 person-years	46	51	65	86	116	135	189
Age-adjusted HRs (95% CIs)	1.00	1.15 (1.04-1.27)	1.40 (1.27-1.54)	1.76 (1.60-1.94)	2.25 (2.04-2.49)	2.56 (2.30-2.86)	3.32 (2.86-3.86)
Model 1† HRs (95% CIs)	1.00	1.04 (0.94-1.15)	1.15 (1.05-1.27)	1.33 (1.20-1.46)	1.58 (1.43-1.75)	1.65 (1.48-1.84)	1.90 (1.63-2.22)
Model 2‡ HRs (95% CIs)	1.00	1.12 (0.90-1.14)	1.23 (0.99-1.53)	1.59 (1.28-1.98)	1.67 (1.32-2.11)	1.99 (1.52-2.60)	1.86 (1.16-2.98)
Model 3§ HRs (95% CIs)	1.00	0.98 (0.89-1.08)	1.12 (1.02-1.23)	1.27 (1.16-1.39)	1.47 (1.33-1.62)	1.55 (1.39-1.73)	1.91 (1.66-2.21)
Women, age 60-79 years (n = 17 736)							
No. of person-years	5 535	10 525	15 260	12 907	7 673	3 611	598
No. of hypertension cases	790	1 591	2 580	2 569	1 663	940	190
Incidence rates per 1 000 person-years	143	151	169	199	217	260	318
Age-adjusted HRs (95% CIs)	1.00	1.11 (1.02-1.21)	1.24 (1.14-1.34)	1.42 (1.31-1.54)	1.53 (1.40-1.66)	1.77 (1.61-1.95)	2.07 (1.77-2.43)
Model 1† HRs (95% CIs)	1.00	1.04 (0.95-1.13)	1.09 (1.01-1.19)	1.21 (1.11-1.31)	1.25 (1.15-1.37)	1.39 (1.26-1.54)	1.61 (1.37-1.90)
Model 2‡ HRs (95% CIs)	1.00	1.04 (0.80-1.35)	1.16 (0.90-1.49)	1.25 (0.96-1.62)	1.35 (1.02-1.80)	1.20 (0.84-1.70)	1.66 (0.82-3.38)
Model 3§ HRs (95% CIs)	1.00	1.06 (0.98-1.16)	1.10 (1.02-1.19)	1.22 (1.12-1.32)	1.29 (1.18-1.41)	1.40 (1.27-1.54)	1.72 (1.47-2.01)

Abbreviations: HRs, hazard ratios; CIs, confidence intervals; N/A, data was not available.

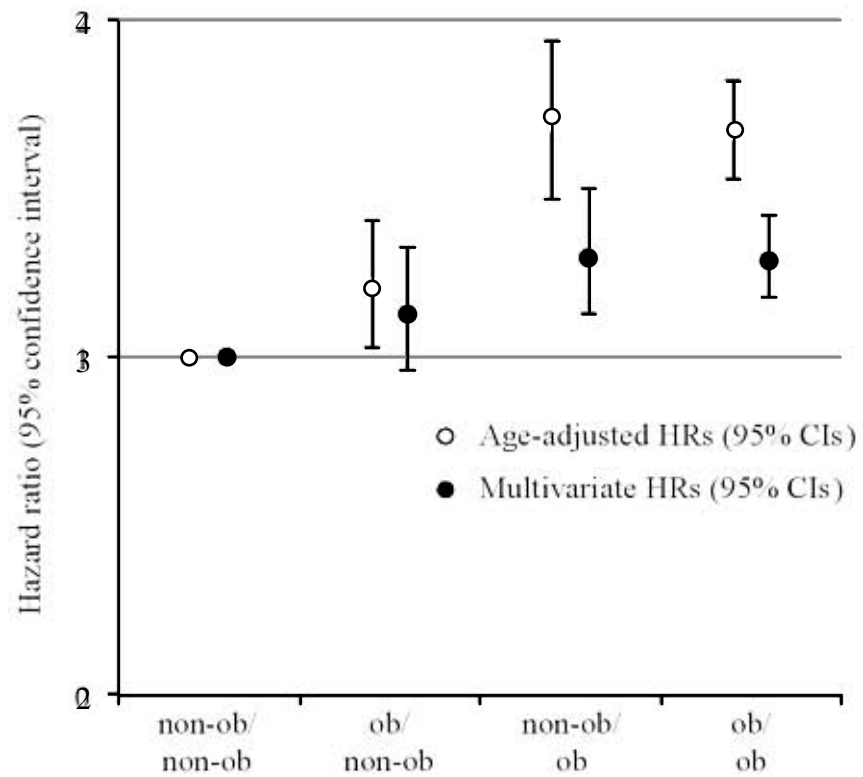
†Adjusted for age (years), systolic blood pressure (mmHg), diastolic blood pressure (mmHg), fasting status (yes or no), total cholesterol level (mmol/liter), high density lipoprotein cholesterol level (mmol/liter), log-transformed triglyceride level (mmol/liter), lipid medication use (yes or no), blood glucose status (normal, border, hyperglycemia), diabetic medication use (yes or no), smoking status (never, ex-, < 20 cigarettes/day, or ≥ 20 cigarettes/day), and alcohol intake (never, sometimes, < 66 g/day, or ≥ 66 g/day).

‡Addition of BMI (kg/m²), total cholesterol level (mmol/liter), high-density lipoprotein cholesterol level (mmol/liter), and log-transformed triglyceride level (mmol/liter) change values (the endpoint minus the baseline) to Model 1 for adjustment.

§Adjusted for time-dependent covariates: age (years), systolic blood pressure (mmHg), diastolic blood pressure (mmHg), fasting status (yes or no), total cholesterol level (mmol/liter), high density lipoprotein cholesterol level (mmol/liter), log-transformed triglyceride level (mmol/liter), lipid medication use (yes or no), blood glucose status (normal, border, hyperglycemia), diabetic medication use (yes or no), smoking status (never, ex-, < 20 cigarettes/day, or ≥ 20 cigarettes/day), and alcohol intake (never, sometimes, < 66 g/day, or ≥ 66 g/day).

Bold values showed statistical significance ($P < 0.05$).

Figure 1



No. of subjects	14 055	740	676	2 174
No. of hypertension cases	2 877	165	165	662