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### The association of ideal cardiovascular health with incident type 2 diabetes mellitus: the Multi-Ethnic Study of Atherosclerosis

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#### Abstract

**Aims/hypothesis**—Levels of ideal cardiovascular health (ICH) and incident type 2 diabetes mellitus have not been examined in a multiethnic population. We assessed the total and race/ ethnicity-specific incidence of diabetes based on American Heart Association (AHA) ICH components.

**Methods**—Incident diabetes was assessed among 5,341 participants in the Multi-Ethnic Study of Atherosclerosis without prevalent diabetes between 2002 and 2012. ICH components (total cholesterol, BP, dietary intake, tobacco use, physical activity and BMI) were assessed at baseline and participants were categorised as having ideal, intermediate or poor cardiovascular health, as defined by the AHA 2020 impact goals. We developed a scoring system based on the number of

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ICH components (0–1 'poor', 2–3 'intermediate', and 4 'ideal'). HRs were calculated using Cox models.

**Results**—During a median follow-up of 11.1 years, we identified 587 cases of incident diabetes. After multivariable adjustment, participants with 2-3 and 4 ICH components vs 0-1 components had a 34% lower (HR 0.66; 95% CI 0.54, 0.80) and a 75% lower (HR 0.25; 95% CI 0.18, 0.35) diabetes incidence, respectively. There were significant differences by race/ethnicity: African-American and Hispanic-American participants with 4 ICH components had diabetes incidence rates per 1,000 person-years of 5.6 (95% CI 3.1, 10.1) and 10.5 (95% CI 6.7, 16.4), respectively, compared with 2.2 (95% CI 1.3, 3.7) among non-Hispanic white Americans.

**Conclusions/interpretation**—Meeting an increasing number of AHA 2020 impact goals for dietary intake, physical activity, smoking, BP, cholesterol and BMI was associated with a dose-dependent lower risk of diabetes with significant variation by race/ethnicity.

#### Keywords

Adiposity; Blood pressure; Cholesterol; Diabetes; Dietary intake; Glucose; Ideal cardiovascular health; Physical activity; Race/ethnicity; Smoking

#### Introduction

In 2010, the American Heart Association (AHA) published 2020 impact goals for cardiovascular health promotion with the aim of improving cardiovascular health and reducing deaths from cardiovascular disease (CVD) and stroke by 20% by the year 2020 [1]. As part of this initiative, the AHA defined the concept of ideal cardiovascular health (ICH), identifying seven health factors or behaviours that have been associated with healthy ageing without cardiovascular and other chronic diseases. These include total cholesterol, BP, fasting plasma glucose, dietary intake, tobacco use, physical activity and BMI (Table 1) [1]. Many cardiovascular risk factors also confer a high risk of incident type 2 diabetes mellitus, including physical inactivity, obesity, unhealthy dietary habits, and, to a lesser extent, elevated BP and dyslipidaemia [2]. CVD is the leading cause of morbidity and mortality among individuals with diabetes. Individuals with diabetes have a threefold increase in cardiovascular mortality compared with those without diabetes [3]. Given the shared risk factors and burden of both CVD and diabetes in the USA, it is important to determine whether the 2020 impact goals for ICH will impact diabetes incidence.

Adherence to the various components of ICH varies by ethnicity [4–6]. While one study among American Indians suggested that individuals who met a greater number of ICH goals had a reduced risk of incident diabetes [6], to our knowledge this hypothesis has not been assessed in a multiethnic population.

We tested the hypothesis that participants with higher levels of cardiovascular health are less likely to develop diabetes based on the AHA metric in four racial/ethnic groups in the Multi-Ethnic Study of Atherosclerosis (MESA). We additionally examined whether the association of individual or overall cardiovascular health components with incident diabetes varies by race/ethnicity.

#### Methods

MESA is a population-based sample of 6,814 men and women from four ethnic groups: non-Hispanic whites (NHW; 38%), African-Americans (AA; 28%), Chinese-Americans (CA; 12%) and Hispanic-Americans (HA; 22%). Participants were aged 45–84 years at baseline; those who reported a medical history of heart attack, angina, coronary revascularisation, pacemaker or defibrillator implantation, valve replacement, heart failure or cerebrovascular disease were excluded. Details of sampling and recruitment have been published elsewhere [7]. The study was approved by the institutional review boards of the six participating institutions.

During the baseline examination (2000–2002), standardised questionnaires and calibrated devices were used to obtain participant data including: demographics, occupation, level of education, tobacco use, alcohol consumption, medical conditions and current prescription medication. Race/ethnicity was assessed by self-report and categorised as NHW, non-Hispanic Black (AA), Hispanic (HA) or Chinese (CA). Current occupational status was classified into ten categories: homemaker, employed (full time, part time, on leave due to ill health, or on leave due to non-health related issues), unemployed (<6 or >6 months) and retired (not working, working or volunteering). Education was classified into one of the following five categories: less than high school, completed high school, some college/ technical school certificate or associate degree, bachelor's degree, and graduate or professional school. Smoking history was obtained using a questionnaire developed from the third National Health and Nutrition Examination Survey (NHANES III), the National Health Interview Survey (NHIS) and the Atherosclerosis Risk in Communities Study (ARIC). Current alcohol use was assessed as a binary variable by asking participants whether they presently drank alcoholic beverages. Calibrated devices were used to measure participants' weight, waist circumference and height; BMI was calculated as weight in kilograms divided by the square of height in metres. Resting seated BP was measured three times using a Dinamap Pro 100, automated oscillometric sphygmomanometer (Critikon, Tampa, FL, USA); the last two measurements were averaged for analysis.

#### Physical activity

The MESA Typical Week Physical Activity Survey, adapted from the Cross-Cultural Activity Participation Study, was used to assess physical activity [8]. We used the intentional exercise variable (sum of walking for exercise, sports/dancing, and conditioning in metabolic equivalent of task [MET] min/week) with the following categorisation: 'poor' <3 MET min/week, 'intermediate' 3–449 MET min/week, 'ideal' 450 MET min/week. Given that 3 METs and 6 METs are the lower limits for moderate and vigorous intensity physical activity, our MET min/week categorisation is consistent with the minutes per week measure in the AHA guidelines [1].

#### **Dietary assessment**

Dietary intake was assessed using the MESA food frequency questionnaire, a selfadministered modified-Block-style 120-item questionnaire adapted from the Insulin Resistance Atherosclerosis Study instrument [7]. The MESA questionnaire had some slight

differences compared with the 2020 guidelines regarding units of servings, requiring modification of the metrics. Components of the modified ideal diet score are: fruits and vegetables, 4.5 cups (equivalent to 1.08 l)/day; fish, two or more 98 g (3.5 ounce) servings per week (non-fried); fibre-rich whole grains, three or more 28 g (1 ounce)-equivalent servings/day; sodium, <1,500 mg/day; sugar-sweetened beverages, 1,884 kJ (1008 g)/ week. Participants were given 1 point per dietary component at goal for a total score ranging from 0 to 5. Participants were classified as ideal (4-5 of 5 metrics), intermediate (2-3 of 5 metrics) or poor (0-1 of 5 metrics).

#### Laboratory assessment

Fasting blood samples were drawn and processed using a standardised protocol and sent to central laboratories at the Collaborative Studies Clinical Laboratory at Fairview University of Minnesota Medical Center (Minneapolis, MN, USA) for measurement of glucose, cholesterol and estimated glomerular filtration rate [9]. Serum glucose was measured by rate reflectance spectrophotometry using thin-film adaptation of the glucose oxidase method on a Vitros analyser (Ortho Clinical Diagnostics, Rochester, NY, USA). Total cholesterol was measured using a cholesterol oxidase method (Roche Diagnostics, Indianapolis, IN, USA) on the Roche Cobas Fara centrifugal analyser. Serum creatinine was measured by rate reflectance spectrophotometry using thin-film adaptation of the creatinine amidinohydrolase method on the Vitros analyser (Ortho Clinical Diagnostics) and calibrated to Cleveland Clinic. Creatinine was used to estimate glomerular filtration rate based on the Chronic Kidney Disease Epidemiology Collaboration equation [10].

#### Cardiovascular health

Cardiovascular health was assessed among the seven baseline metrics: smoking status, diet, physical activity, BMI, serum cholesterol and BP, with the addition of fasting plasma glucose in the sensitivity analyses [6]. Each baseline metric was scored and categorised as poor, intermediate or ideal, as specified by AHA recommendations, with consideration of medication use (i.e. antihypertensive, lipid-lowering, glucose-lowering) where appropriate [1]. Points were allocated for each of the seven baseline metrics with scores of 0 (poor or intermediate) or 1 (ideal) for each health behaviour (diet, smoking, physical activity, BMI) and health factor (BP, blood sugar, total cholesterol) (Table 1). We summed the metrics into the following categories: poor (0–1), intermediate (2–3) and ideal (4) levels of cardiovascular health. Second, we used a previously developed scoring system [4] in which each cardiovascular health (Table 1), respectively. A total cardiovascular health score (excluding glucose) ranging from 0 to 12 was calculated as the sum of the individual cardiovascular health component scores. This score was classified into three levels as inadequate (0–4), average (5–8) or optimal (9–12) cardiovascular health.

#### **Diabetes definition**

Among those without prevalent baseline diabetes, individuals newly using glucose-lowering medication or having fasting glucose 7 mmol/l (126 mg/dl) at one of the four subsequent examinations (the last follow-up visit occurring in 2010-2012) were considered to have

incident diabetes [11]. We defined time of incident diabetes as the midpoint between the last examination without diabetes and the examination at which diabetes developed.

We were unable to distinguish between type 1 diabetes and type 2 diabetes, but in this age range, incident type 1 diabetes is extremely uncommon, so we assumed a predominance of type 2 diabetes in our population.

#### Statistical analysis

We excluded participants with diabetes at baseline (n=883), missing diabetes status at their last follow-up (n=57) or missing data on baseline covariates (n=526). The 583 participants excluded due to missing diabetes and covariate status had a higher percentage of AA, higher BMI, higher smoking, lower education, less physical activity and a higher rate of incident diabetes (p<0.05 for all comparisons) (electronic supplementary material [ESM] Table 1). Descriptive statistics were used to compare the baseline characteristics of all participants (non-stratified) and by race/ethnicity, using appropriate parametric or non-parametric tests for continuous variables and the  $\chi^2$  test for categorical variables. Among the seven ICH components, we did not consider the blood glucose metric in the main analysis, as diabetes was the endpoint of interest. We used the other six components of the ICH score [6] to develop the two aforementioned scoring systems for ICH and total cardiovascular health. Last, we assessed the association between individual cardiovascular health components and incident diabetes by comparing participants with intermediate or ideal status with those with poor status (reference group) at baseline for each of the six factors. Unadjusted diabetes incidence rates for cardiovascular health scores were calculated using person-time analysis assuming a Poisson distribution. Participants were censored at the last attended follow-up examination. Incidence rate ratios were assessed using the logrank test. Cox proportional hazards modelling was used to estimate HRs associated with the aforementioned classifications. Covariates in adjusted analyses included baseline study site, age, sex, race/ ethnicity, education, occupational status, alcohol use and estimated glomerular filtration rate. The baseline number of cardiovascular health components and cardiovascular health values categorised as poor, intermediate or ideal were evaluated to estimate the proportion of cases in the population that might be attributable to suboptimal levels of cardiovascular health (population-attributable risk). The population-attributable risk was calculated using the formula p(RR-1)/(1+p[RR-1]), where p is the prevalence of individuals not in the low-risk group and RR (risk ratio, HR) is the associated multivariable-adjusted relative risk of those individuals. Upper and lower 95% CIs of the population-attributable risk were derived using this formula and the upper and lower 95% CI estimates of the multivariable-adjusted risk ratio [12]. Given that the association of cardiovascular health attainment with diabetes risk may differ by age, sex and race/ethnicity, we tested for interaction by these factors with each cardiovascular health measure by inserting an interaction term in the model and using the likelihood ratio test. Statistical significance was defined as two-sided  $\alpha$ <0.05. Analyses were performed using Stata 13.1 software (StataCorp, College Station, TX, USA).

#### Results

#### Characteristics of the cohort

The baseline characteristics of participants stratified by race/ethnicity revealed that characteristics were different across race/ethnicity. HA and AA participants had significantly higher BMI, systolic BP and fasting glucose and lower levels of ICH components compared with NHW (Table 2).

#### Incidence of diabetes

During a median follow-up of 11.1 years, 587 participants developed incident diabetes (incidence rate 11.1 per 1,000 person–years). Incidence rates were highest among HA (15.3) and AA (12.3) (Table 2). Participants who developed diabetes had higher baseline BMI (30.8 vs 27.5 kg/m<sup>2</sup>), systolic BP (130 vs 125 mmHg) and fasting plasma glucose (5.61 vs 4.88 mmol/l,[101 vs 88 mg/dl]) (comparisons p<0.0001; ESM Table 2).

Diabetes incidence rates per 1,000 person-years in participants with 0-1, 2-3 or 4 ICH components were 18.9 (95% CI 15.9, 22.4), 12.1 (95% CI 11.0, 13.3) and 4.4 (95% CI 3.4, 5.7), respectively, with a risk ratio of incident diabetes per category of 0.52 (95% CI 0.46, 0.60) (Table 3). Incident diabetes rates decreased for every additional ICH component achieved in the overall cohort and each racial/ethnic group (Table 3).

#### ICH and incident diabetes

The unadjusted and adjusted HRs for incident diabetes associated with baseline ICH are presented in Table 4. After adjustment, the HR for 5-6 vs 0 individual ICH components was 0.17 (95% CI 0.08, 0.38), and the HRs for intermediate or ideal categories compared with the poor cardiovascular health category was 0.66 (95% CI 0.54, 0.80) and 0.25 (95% CI 0.18, 0.35), respectively. Among the individual cardiovascular health components, ideal vs poor status was associated with a decreased risk of incident diabetes for BP (HR 0.53; 95% CI 0.42, 0.68) and BMI (HR 0.25; 95% CI 0.19, 0.33) in adjusted analyses and for physical activity (HR 0.82; 95% CI 0.67, 1.00) in unadjusted analyses (Table 5). We found no consistent evidence of interactions between cardiovascular health components and age or sex (ESM Table 3). Sensitivity analyses conducted using the total cardiovascular health score (ESM Tables 4-6, ESM Figure 1), including participants with missing dietary data (n=5,817) (ESM Table 7), the addition of baseline glucose as a covariate (ESM Table 8), or the inclusion of glucose as a cardiovascular health component (seven components) (ESM Table 9) revealed similar findings and remained significant. Ideal (<5.6 mmol/l [<100 mg/dl]) vs intermediate (5.6-7.0 mmol/l [100-<126 mg/dl]) baseline blood glucose categories were associated with an 87% reduction in risk (ESM Table 10).

#### Cardiovascular health and incident diabetes by race/ethnicity

There were significant differences by race/ethnicity in the association of baseline ICH components with risk of diabetes (all p<0.01; ESM Table 2). In Table 4, ideal vs poor cardiovascular health was associated with a greater reduction in diabetes risk in NHW and CA (87% and 88%) vs AA and HA (66% and 50%) (Fig. 1). Among the individual cardiovascular health components, ideal vs poor status was significant for: BP in NHW, CA

and AA, with the greatest association of risk reduction in NHW (HR 0.36; 95% CI 0.23, 0.55); and BMI in all ethnicities, with no significant difference among the race/ethnicities (Table 5).

#### Population-attributable diabetes risk

We evaluated diabetes risk by baseline number of cardiovascular health components and categories at baseline, compared with all others not in that group (Table 6). The baseline number of ICH components 2, 3, 4 or 5 were associated with a 30%, 23%, 63% and 66% lower diabetes incidence, respectively. Individuals in the ICH category at baseline (23%) had a 65% lower incidence of diabetes, and three in five new cases of diabetes appeared attributable to not being in the ICH group at baseline.

#### Discussion

In this large, contemporary, multiethnic cohort study, the presence of an increasing number of ICH components at baseline was strongly and inversely associated with incident diabetes. The effect of ICH was graded: participants with intermediate or ideal vs poor cardiovascular health had a 34% and 75% lower diabetes incidence, respectively, which is consistent with previous studies [12, 13]. In analyses stratified by race/ethnicity, the greatest reductions in diabetes risk based on ICH components were seen in NHW and CA. Overall, three out of five cases of diabetes in this middle-aged population appeared attributable to not having ICH at baseline. If these associations are causal, diabetes could be prevented by attainment of at least four ICH components. Compared with previous studies examining the combined effects of multiple risk factors on the incidence of diabetes [12, 13], we used a novel AHA concept that may be more adapted for translation of prevention policies that encompass both diabetes and CVD into practice. Public health interventions over the last 30 years have led to plateauing of diabetes incidence among NHW, but among AA and HA there remains a continued ascent in incident diabetes [14]. Overall, these findings support promotion of and adherence to the AHA 2020 impact goals to lower diabetes incidence among all races/ ethnicities, but emphasise the importance of tailoring interventions to prevent diabetes in racial/ethnic minorities given the observed disparities.

#### ICH goals and incident diabetes

Our study is the first multiethnic analysis to assess the association of baseline ICH with incident diabetes. A previous study among American Indians showed that participants who achieved 0–1 ICH goals compared with those who achieved 2–3 or 4 ICH goals had a 60% and 89% lower odds of developing diabetes [6], compared with the 34% (2-3 ICH goals) and 75% (4 ICH goals) lower diabetes incidence compared with 0-1 ICH goals in our study. The observed difference is likely related to the inclusion of glucose among the ICH components in the Fretts et al study [6], whereas we excluded glucose in our main analysis, as it is directly in the causal pathway to diabetes. However, sensitivity analyses including glucose as a component of cardiovascular health revealed a similar order of risk reduction: -61% and -86% lower diabetes risk of participants in the intermediate or ideal category compared with those in the poor category. In the Cardiovascular Health Study [12], among older US adults (age >65 years) low-risk lifestyle groups defined by physical activity, dietary

score, smoking, alcohol use, BMI and waist circumference were associated with an 89% reduction in diabetes risk with five low-risk lifestyle factors, similar to our 89% reduction in risk with 4 ICH components. In the National Institutes of Health–American Association of Retired Persons (NIH-AARP) Diet and Health Study, Reis et al [13] examined the association of low-risk lifestyle factors including BMI, diet, smoking, moderate alcohol consumption and regular physical activity with incident diabetes (self-report). The ORs for incident diabetes over 11 years in those with all five low-risk lifestyle factors at baseline were 0.28 (95% CI 0.23, 0.34) and 0.16 (95% CI 0.10, 0.24) for men and women, respectively, suggesting sex differences. These findings were similar to our ICH category ( 4 ICH components), and the test for effect modification by sex in our analysis was non-significant. We extended these previous findings by assessing a multiethnic population, as the Cardiovascular Health Study is 88.6% NHW and the NIH-AARP Diet and Health Study is 94.3% NHW, thus limiting generalisability of the finding to other racial/ethnic groups.

#### **Racial/ethnic differences**

Consistent with previous US population-based studies [15], AA and HA in the MESA cohort had lower levels of ICH at baseline. We observed significant variation of the association of higher ICH with lower incident diabetes among racial/ethnic groups. NHW and CA had greater magnitudes of risk reduction and better cardiovascular health compared with AA and HA (p<0.01). The lower prevalence of ICH [15], combined with lower magnitude of diabetes reduction with ICH in AA and HA, provide a potential explanation and intervention target for the disparities in diabetes prevalence among these groups.

#### Individual AHA cardiovascular health goals and incident diabetes

Impaired glucose tolerance, impaired fasting glucose and  $HbA_{1c}$  in the prediabetic range (5.7-6.4% [38.8 – 46.4 mmol/mol]) are associated with the greatest risk of diabetes, as they reflect derangements in the pathway from normal glucose tolerance to diabetes [16, 17]. Among risk factors not directly in the causal pathway, BMI was the predominant risk factor for diabetes, with a 48% and 77% lower diabetes risk for overweight BMI and normal BMI, respectively, vs obese BMI. This is consistent with previous studies showing increasing BMI as the primary risk factor increasing diabetes prevalence in the USA over the last 30 years [18]. In the Diabetes Prevention Program (DPP), sustained weight loss was the primary driver of reduced diabetes risk and cardiometabolic improvement [19]. In US populations, the imbalance between caloric intake and energy expenditure is the primary driver of increasing BMI and may have even greater importance in some racial/ethnic groups due to a reduction in baseline energy expenditure [20].

Among the other individual components, normal BP (<120/<80 mmHg) was associated with a 47% diabetic risk reduction in the overall cohort vs elevated BP (>140/90 mmHg), with NHW showing the greatest reduction in diabetes risk compared with the other racial/ethnic groups. In a prior study, BP elevation was associated with incident diabetes in AA and NHW in age-adjusted analyses, but the association was nonsignificant among AA after adjustment for other diabetic risk factors [21]. This corresponded with an earlier study in which mean BP was significantly correlated with fasting plasma insulin and rate of glucose disposal in whites, but not in blacks [22]. Contrary to the prior literature, in our study, BP remained

Physical activity is beneficial in reducing incident diabetes in the majority of NHW observational studies [24]. There is a scarcity of data on physical activity alone and incident diabetes in racial/ethnic minority groups in the USA. In observational studies, AA and HA have lower physical activity levels than NHW, independently of social class [25]. In a study of AA women, vigorous physical activity alone was associated with a dose-dependent reduction in risk of incident diabetes [26]. Further studies to understand the impact of physical activity on diabetes risk are warranted in US racial/ethnic minority groups.

mediated by inflammation and decreased antioxidants, which may partly explain the

association between BP and diabetes [23].

Smoking, dietary intake and total cholesterol were not individually associated with incident diabetes in our analysis. In large multiethnic meta-analyses, smoking increased the risk of diabetes, with a pooled adjusted relative risk of 1.4 (95% CI 1.3, 1.6) vs non-smokers [27]. Data on causal inference between smoking and the development of diabetes are inconsistent: some studies show impairment in insulin sensitivity [28, 29] and glucose tolerance [29, 30], while other studies conclude that causal inference is not likely, after adjusting for confounders including age and BMI [31]. Further complicating the relationship are data from smoking cessation studies showing worsened glucose metabolism in those with diabetes in the first 1–3 years after smoking cessation [32], and an increased risk of incident diabetes in smokers without diabetes who quit, with subsequent improvement to similar risk of that of non-smokers over 12 years [33]. Smokers are also more likely to have unhealthy behaviours and low socioeconomic status, which contribute to the risk of diabetes [27].

While there was no association of dietary intake with incident diabetes, we are unable to draw definitive conclusions because only 1.5% of participants had baseline ideal dietary intake. Components of the AHA diet score including fruits and vegetables, fibre-rich whole grains, decreased sodium and sugar-sweetened beverage intake, and adherence to components of the Dietary Approaches to Stop Hypertension (DASH) and the Mediterranean diet have been associated with improvements in glucose metabolism and decreased incident diabetes [34–38]. A meta-analysis of ten large prospective studies found that dietary patterns similar to the AHA dietary recommendations were associated with a 66% reduction in diabetes [39]. Notably, the AHA diet score does not take into account other forms of dietary intake associated with decreases in diabetes and CVD risk including dairy consumption [40–42] and glycaemic index/load [43–45].

We found no significant overall association between total cholesterol and incident diabetes in our analysis. Components of the lipid profile including triacylglycerols, HDL and nonesterified fatty acids, have been associated with incident diabetes [46].

Among the AHA cardiovascular health components, BMI is well studied in US racial/ethnic minorities, in whom weight loss has been shown to reduce diabetes risk [19]. The association of individual components including BP, physical activity, smoking, total cholesterol and dietary intake with diabetes has previously revealed inconsistent findings and would benefit from further study among US racial/ethnic minorities.

#### Strengths and limitations

Strengths of our analysis include a moderately large, socioeconomically diverse, multiethnic US population with over a decade of follow-up, allowing broad generalisability of our findings. We used validated questionnaires and documentation of diabetes over time with fasting glucose, medication use and self-reported physician diagnosis. Nevertheless, there are several potential limitations. Physical activity and diet were self-reported; thus, misclassification and residual confounding by these variables may have occurred. The time frame queried was the last month prior to the baseline visit; so, depending on the season of examination, the prior month may not be representative of customary physical activity and dietary intake. Sample sizes varied for the racial/ethnic groups, with power implications for detecting significant racial/ethnic interactions in stratified analyses, but the interaction terms for categories of cardiovascular health were significant (p<0.01). As previously mentioned, we were unable to distinguish between type 1 diabetes and type 2 diabetes, so we assumed a predominance of type 2 incident diabetes in our population.

#### Conclusions

Our study showed that increasing levels of ideal and overall cardiovascular health within the guidelines set forth by the AHA 2020 impact goals may reduce the burden of diabetes in the USA. Unfortunately, less than one in four participants in our overall cohort and less than one in six racial/ethnic minorities attained 4 ICH components, which is similar to findings in other studies [4, 5, 15]. Given the racial/ethnic differences in attainment of ICH, the lower magnitude of risk reduction with ICH and the increased burden of diabetes in racial/ethnic minorities, further research on promotion, attainment and ethnic differences of ICH in US racial/ethnic minority groups is of paramount importance to lower the risk of CVD and diabetes.

#### Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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#### References

- Lloyd-Jones DM, Hong Y, Labarthe D, et al. Defining and setting national goals for cardiovascular health promotion and disease reduction: the American Heart Association's strategic impact goal through 2020 and beyond. Circulation. 2010; 121:586–613. [PubMed: 20089546]
- 2. Gress TW, Nieto FJ, Shahar E, et al. Hypertension and antihypertensive therapy as risk factors for type 2 diabetes mellitus. N Engl J Med. 2000; 342:905–912. [PubMed: 10738048]

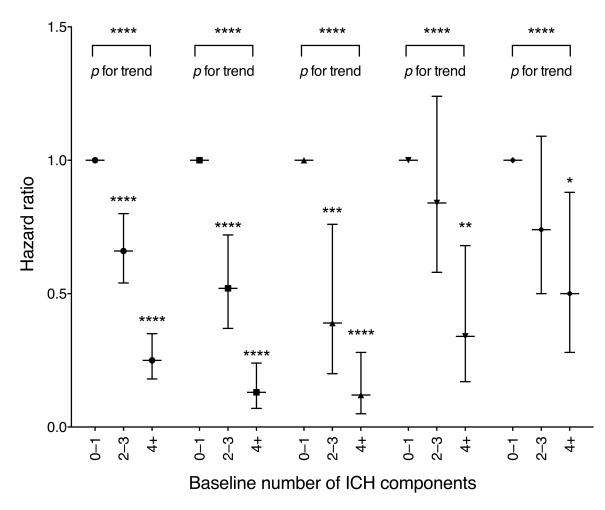
- Taylor KS, Heneghan CJ, Farmer AJ, et al. All-cause and cardiovascular mortality in middle-aged people with type 2 diabetes compared with people without diabetes in a large U.K. primary care database. Diabetes Care. 2013; 36:2366–2371. [PubMed: 23435157]
- Kulshreshtha A, Vaccarino V, Judd SE, et al. Life's simple 7 and risk of incident stroke: the reasons for geographic and racial differences in stroke study. Stroke. 2013; 44:1909–1914. [PubMed: 23743971]
- Folsom AR, Yatsuya H, Nettleton JA, et al. Community prevalence of ideal cardiovascular health, by the American Heart Association definition, and relationship with cardiovascular disease incidence. J Am Coll Cardiol. 2011; 57:1690–1696. [PubMed: 21492767]
- Fretts AM, Howard BV, McKnight B, et al. Life's simple 7 and incidence of diabetes among American Indians: the Strong Heart Family Study. Diabetes Care. 2014; 37:2240–2245. [PubMed: 24804696]
- Bild DE. Multi-Ethnic Study of Atherosclerosis: objectives and design. Am J Epidemiol. 2002; 156:871–881. [PubMed: 12397006]
- Bertoni AG, Whitt-Glover MC, Chung H, et al. The association between physical activity and subclinical atherosclerosis: the Multi-Ethnic Study of Atherosclerosis. Am J Epidemiol. 2008; 169:444–454. [PubMed: 19075250]
- 9. Cushman M, Cornell ES, Howard PR, et al. Laboratory methods and quality assurance in the Cardiovascular Health Study. Clin Chem. 1995; 41:264–270. [PubMed: 7874780]
- Levey AS, Stevens LA, Schmid CH, et al. A new equation to estimate glomerular filtration rate. Ann Intern Med. 2009; 150:604–612. [PubMed: 19414839]
- American Diabetes Association. Diagnosis and classification of diabetes mellitus. Diabetes Care. 2010; 33(Suppl 1):S62–S69. [PubMed: 20042775]
- Mozaffarian D, Kamineni A, Carnethon M, et al. Lifestyle risk factors and new-onset diabetes mellitus in older adults: the Cardiovascular Health Study. Arch Intern Med. 2009; 169:798–807. [PubMed: 19398692]
- Reis JP, Loria CM, Sorlie PD, et al. Lifestyle factors and risk for new-onset diabetes: a populationbased cohort study. Ann Intern Med. 2011; 155:292–299. [PubMed: 21893622]
- Geiss LS, Wang J, Cheng YJ, et al. Prevalence and incidence trends for diagnosed diabetes among adults aged 20 to 79 years, United States, 1980-2012. JAMA. 2014; 312:1218. [PubMed: 25247518]
- Shay CM, Ning H, Allen NB, et al. Status of cardiovascular health in US adults: prevalence estimates from the National Health and Nutrition Examination Surveys (NHANES) 2003–2008. Circulation. 2012; 125:45–56. [PubMed: 22095826]
- Morris DH, Khunti K, Achana F, et al. Progression rates from HbA<sub>1c</sub> 6.0–6.4% and other prediabetes definitions to type 2 diabetes: a meta-analysis. Diabetologia. 2013; 56:1489–1493. [PubMed: 23584433]
- 17. Nathan DM, Davidson MB, DeFronzo RA, et al. Impaired fasting glucose and impaired glucose tolerance: implications for care. Diabetes Care. 2007; 30:753–759. [PubMed: 17327355]
- Menke A, Rust KF, Fradkin J, et al. Associations between trends in race/ethnicity, aging, and body mass index with diabetes prevalence in the United States: a series of cross-sectional studies. Ann Intern Med. 2014; 161:328. [PubMed: 25178569]
- Delahanty LM, Pan Q, Jablonski KA, et al. Effects of weight loss, weight cycling, and weight loss maintenance on diabetes incidence and change in cardiometabolic traits in the Diabetes Prevention Program. Diabetes Care. 2014; 37:2738–2745. [PubMed: 25024396]
- 20. Gannon B, DiPietro L, Poehlman ET. Do African Americans have lower energy expenditure than Caucasians? Int J Obes Relat Metab Disord J Int Assoc Study Obes. 2000; 24:4–13.
- Wei GS, Coady SA, Goff DC, et al. Blood pressure and the risk of developing diabetes in African Americans and whites: ARIC, CARDIA, and the Framingham Heart Study. Diabetes Care. 2011; 34:873–879. [PubMed: 21346180]
- 22. Saad MF, Lillioja S, Nyomba BL, et al. Racial differences in the relation between blood pressure and insulin resistance. N Engl J Med. 1991; 324:733–739. [PubMed: 1997839]
- Das UN. Risk of type 2 diabetes mellitus in those with hypertension. Eur Heart J. 2008; 29:952– 953. [PubMed: 18314422]

- 24. Jeon CY, Lokken RP, Hu FB, van Dam RM. Physical activity of moderate intensity and risk of type 2 diabetes: a systematic review. Diabetes Care. 2007; 30:744–752. [PubMed: 17327354]
- 25. Crespo CJ, Smit E, Andersen RE, et al. Race/ethnicity, social class and their relation to physical inactivity during leisure time: results from the Third National Health and Nutrition Examination Survey, 1988-1994. Am J Prev Med. 2000; 18:46–53. [PubMed: 10808982]
- 26. Krishnan S, Rosenberg L, Palmer JR. Physical activity and television watching in relation to risk of type 2 diabetes: the Black Women's Health Study. Am J Epidemiol. 2008; 169:428–434. [PubMed: 19056835]
- 27. Willi C, Bodenmann P, Ghali WA, et al. Active smoking and the risk of type 2 diabetes: a systematic review and meta-analysis. JAMA. 2007; 298:2654–2664. [PubMed: 18073361]
- Frati AC, Iniestra F, Ariza CR. Acute effect of cigarette smoking on glucose tolerance and other cardiovascular risk factors. Diabetes Care. 1996; 19:112–118. [PubMed: 8718429]
- 29. Piatti P, Setola E, Galluccio E, et al. Smoking is associated with impaired glucose regulation and a decrease in insulin sensitivity and the disposition index in first-degree relatives of type 2 diabetes subjects independently of the presence of metabolic syndrome. Acta Diabetol. 2014; 51:793–799. [PubMed: 24934227]
- Janzon L, Berntorp K, Hanson M, et al. Glucose tolerance and smoking: a population study of oral and intravenous glucose tolerance tests in middle-aged men. Diabetologia. 1983; 25:86–88.
   [PubMed: 6354814]
- Wareham NJ, Ness EM, Byrne CD, et al. Cigarette smoking is not associated with hyperinsulinemia: evidence against a causal relationship between smoking and insulin resistance. Metabolism. 1996; 45:1551–1556. [PubMed: 8969291]
- 32. Lycett D, Nichols L, Ryan R, et al. The association between smoking cessation and glycaemic control in patients with type 2 diabetes: a THIN database cohort study. Lancet Diabetes Endocrinol. 2015; 3:423–430. [PubMed: 25935880]
- Yeh HC, Duncan BB, Schmidt MI, et al. Smoking, smoking cessation, and risk for type 2 diabetes mellitus: a cohort study. Ann Intern Med. 2010; 152:10–17. [PubMed: 20048267]
- Liese AD, Nichols M, Sun X, et al. Adherence to the DASH diet is inversely associated with incidence of type 2 diabetes: the Insulin Resistance Atherosclerosis Study. Diabetes Care. 2009; 32:1434–1436. [PubMed: 19487638]
- 35. Shirani F, Salehi-Abargouei A, Azadbakht L. Effects of Dietary Approaches to Stop Hypertension (DASH) diet on some risk for developing type 2 diabetes: a systematic review and meta-analysis on controlled clinical trials. Nutrition. 2013; 29:939–947. [PubMed: 23473733]
- Martinez-Gonzalez MA, de la Fuente-Arrillaga C, Nunez-Cordoba JM, et al. Adherence to Mediterranean diet and risk of developing diabetes: prospective cohort study. BMJ. 2008; 336:1348–1351. [PubMed: 18511765]
- Salas-Salvado J, Bullo M, Babio N, et al. Reduction in the incidence of type 2 diabetes with the Mediterranean diet: results of the PREDIMED-Reus nutrition intervention randomized trial. Diabetes Care. 2011; 34:14–19. [PubMed: 20929998]
- Schwingshackl L, Missbach B, König J, Hoffmann G. Adherence to a Mediterranean diet and risk of diabetes: a systematic review and meta-analysis. Public Health Nutr. 2015; 18:1292–1299. [PubMed: 25145972]
- Esposito K, Kastorini CM, Panagiotakos DB, Giugliano D. Prevention of type 2 diabetes by dietary patterns: a systematic review of prospective studies and meta-analysis. Metab Syndr Relat Disord. 2010; 8:471–476. [PubMed: 20958207]
- Soedamah-Muthu SS, Ding EL, Al-Delaimy WK, et al. Milk and dairy consumption and incidence of cardiovascular diseases and all-cause mortality: dose-response meta-analysis of prospective cohort studies. Am J Clin Nutr. 2011; 93:158–171. [PubMed: 21068345]
- O'Connor LM, Lentjes MAH, Luben RN, et al. Dietary dairy product intake and incident type 2 diabetes: a prospective study using dietary data from a 7-day food diary. Diabetologia. 2014; 57:909–917. [PubMed: 24510203]
- 42. Yakoob MY, Shi P, Willett WC, et al. Circulating biomarkers of dairy fat and risk of incident diabetes mellitus among men and women in the United States in two large prospective cohorts. Circulation. 2016; 133:1645–1654. [PubMed: 27006479]

- 43. Villegas R, Liu S, Gao YT, et al. Prospective study of dietary carbohydrates, glycemic index, glycemic load, and incidence of type 2 diabetes mellitus in middle-aged Chinese women. Arch Intern Med. 2007; 167:2310–2316. [PubMed: 18039989]
- 44. Sluijs I, van der Schouw YT, van der ADL, et al. Carbohydrate quantity and quality and risk of type 2 diabetes in the European Prospective Investigation into Cancer and Nutrition-Netherlands (EPIC-NL) study. Am J Clin Nutr. 2010; 92:905–911. [PubMed: 20685945]
- Burger KNJ, Beulens JWJ, Boer JMA, et al. Dietary glycemic load and glycemic index and risk of coronary heart disease and stroke in Dutch men and women: the EPIC-MORGEN study. PLoS ONE. 2011; 6:e25955. [PubMed: 21998729]
- 46. Pankow JS, Duncan BB, Schmidt MI, et al. Fasting plasma free fatty acids and risk of type 2 diabetes: the Atherosclerosis Risk in Communities Study. Diabetes Care. 2004; 27:77–82. [PubMed: 14693970]

#### Abbreviations

AA	African-American
AHA	American Heart Association
CA	Chinese-American
CVD	Cardiovascular disease
НА	Hispanic-American
ІСН	Ideal cardiovascular health
MESA	Multi-Ethnic Study of Atherosclerosis
MET	Metabolic equivalent of task
NHW	Non-Hispanic white



#### Fig. 1.

Diabetes Incidence HRs for ICH. ICH was classified as poor (0–1 ICH components) (referent group, HR = 1), intermediate (2–3 ICH components) and ideal ( 4 ICH components) cardiovascular health. Circles, all races/ethnicities; squares, NHW participants; up-pointing triangles, CA participants; down-pointing triangles, AA participants; diamonds, HA participants. Cox modelling and logrank test (*p* for trend) with statistical significance for \*p<0.05, \*\*p<0.01, \*\*\*p<0.001 and \*\*\*\*p<0.0001

Table 1
Total cardiovascular health score and definitions of cardiovascular health <sup>a</sup>

Goal/metric	Poor health	Intermediate health	Ideal health
Current smoking, months	Yes	Former 12	Never or quit 12
Total cholesterol, mmol/l (mg/dl)	6.21 ( 240)	5.18-6.18 (200-239), or treated to goal	<5.18 (<200)
BP, mmHg	Systolic 140 or diastolic 90	Systolic 120-139 or diastolic 80-89, or treated to goal	<120 /<80
BMI, kg/m <sup>2</sup>	30	25.0-29.9	<25
Physical activity, MET min/week $^{b}$	<3	3–449	450
Healthy diet score, components <sup><math>C</math></sup>	0-1	2-3	4-5
Fasting plasma glucose, mmol/l (mg/dl) $^d$	7.0 (126)	5.6-6.9 (100-125), or treated to goal	<5.6 (<100)
Points for ICH score per metric	0	0	1
Points for total cardiovascular health score per metric	0	1	2

 $^{a}$ Adapted from the AHA's strategic planning task force and statistical committee 2020 guidelines  $^{1}$ 

<sup>b</sup>Adapted for MESA: MET min/week

<sup>C</sup>Adapted for MESA: fruits and vegetables, 1.081 (4.5 cups)/day; fish, two or more 98 g (3.5 ounce) servings per week (non-fried); fibre-rich whole grains, three or more 28 g (1 ounce)-equivalent servings/day; sodium, <1,500 mg/day; sugar-sweetened beverages, 1,884 kJ (1008 g)/week

 $d_{\text{Fasting plasma glucose was not used in the main analysis but was considered in the sensitivity analyses.}$ 

Characteristic	Overall n=5,348	NHW n=2,277	CA n=676	AA n=1,293	HA n=1,102
Age	61.9 (10.3)	62.4 (10.3)	61.8 (10.3)	62.0 (10.2)	60.7 (10.5)
Female, % <sup>a</sup>	53.5	52.6	52.2	56.8	52.4
Current smoking, %	14.0	13.1	5.3	19.8	14.1
Ideal cardiovascular diet score, $0-5b$	1.5 (0.9)	1.6(0.9)	1.7 (0.8)	1.4 (1.0)	1.3 (0.9)
Current alcohol use, %	58.3	73.4	31.8	52.7	49.9
Education bachelor's degree, %	37.8	51.3	39.9	36.6	11.2
Employed, % full time	39.4	40.5	37.3	39.4	38.5
Exercise, physical activity MET min/week	1,603 (2,410)	1,725 (2,348)	$1,156\ (1,549)$	1,824 (2,992)	1,365 (2,138)
BMI, kg/m <sup>2</sup>	27.9 (5.3)	27.5 (5.0)	23.8 (3.3)	29.7 (5.8)	29.0 (4.7)
Systolic BP, mmHg	125 (21)	123 (20)	124 (22)	131 (21)	126 (22)
Diastolic BP, mmHg	72 (10)	70 (10)	72 (10)	75 (10)	72 (10)
eGFR CKD-EPI, ml min <sup>-1</sup> $1.73 \text{ m}^{-2}$	78 (16)	74 (14)	80 (15)	80 (17)	80 (15)
Total cholesterol, mmol/l (mg/dl)	5.1 (0.9), 195 (35)	5.1 (0.9), 196 (35)	5.0 (0.8), 193 (32)	4.9 (0.9), 190 (36)	5.2 (0.9), 199 (36)
Fasting plasma glucose, mmol/l (mg/dl)	5.00 (0.61), 89 (10)	4.88 (0.56), 88 (10)	5.10 (0.56), 91 (10)	5.00 (0.61), 90 (11)	5.05 (0.61), 91 (11)
Incident diabetes <sup>C</sup>	11.1 (10.3, 12.1)	8.3 (7.2, 9.5)	11.6 (9.3, 14.5)	12.3 (11.0, 15.0)	15.3 (13.1, 17.8)
Baseline no. of ICH components					
Poor (0-1), %	14.2	12.9	5.8	18.4	16.9
Intermediate (2-3), %	62.8	61.1	55.6	66.1	66.8
Ideal (4-6), %	23.0	26.0	38.6	15.5	16.3
Baseline individual cardiovascular health components	omponents				
BP					
Poor, %	24.1	20.3	23.2	30.9	24.8
Intermediate, %	38.6	38.1	32.8	44.9	35.8
Ideal, %	37.3	41.7	43.9	24.2	39.5
BMI					
Poor, %	28.5	25.7	4.1	40.9	34.8
Intermediate, %	40.2	40.5	29.6	38.8	47.4
Ideal, %	31.3	33.8	66.3	20.3	17.8

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Characteristic	Overall n=5,348	Overall n=5,348 NHW n=2,277	CA n=676	AA n=1,293	HA n=1,102
Total cholesterol					
Poor, %	9.6	10.0	7.5	7.8	12.3
Intermediate, %	42.9	46.6	38.8	40.2	40.7
Ideal, %	47.5	43.4	53.7	52.0	47.0
Smoking					
Poor, %	14.0	13.1	5.3	19.8	14.2
Intermediate, %	1.1	1.5	0.9	0.8	1.0
Ideal, %	84.9	85.4	93.8	79.4	84.9
Dietary intake					
Poor, %	50.3	47.8	38.3	53.7	58.6
Intermediate, %	48.2	50.6	60.8	44.6	40.0
Ideal, %	1.5	1.6	0.9	1.8	1.4
Physical activity					
Poor, %	21.8	16.3	25.4	22.4	30.0
Intermediate, %	13.9	14.0	15.4	13.7	13.1
Ideal, %	64.3	69.7	59.2	64.0	56.9

Values are mean (SD) or percentages unless otherwise indicated

 $^{a}$ All comparisons p < 0.001 except those that are not significant; p values calculated using  $\chi^{2}$  (categorical variables), ANOVA (continuous variables) and logrank tests (incident diabetes)

b Adapted for MESA, with 1 point given for: fruits and vegetables, 1.081 (4.5 cups)/day; fish, two or more 98 g (3.5 ounce) servings per week (non-fried); fibre-rich whole grains, three or more 28 g (1 ounce)-equivalent servings/day; sodium, <1.500 mg/day; sugar-sweetened beverages, 1,884 kJ (1008 g)/week

 $^{C}$ Rate per 1,000 person-years (95% CI)

eGFR CKD-EPI, estimated glomerular filtration rate based on the Chronic Kidney Disease Epidemiology Collaboration

## Table 3

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<b>Baseline ICH level</b>	Overall	MHM	CA	AA	НА
No. of ICH components					
0	21.8 (12.9, 36.9)	21.8 (12.9, 36.9) 24.9 (11.9, 52.3) $NAb$	$^{q}$ NA	12.8 (4.1, 39.8)	12.8 (4.1, 39.8) 34.8 (13.1, 92.7)
I	18.6 (15.5, 22.2)	17.8 (13.3, 23.8)		35.8 (19.8, 64.7) 16.4 (11.6, 23.2) 19.1 (13.3, 27.3)	19.1 (13.3, 27.3)
2	13.0 (11.4, 14.8)	13.0 (11.4, 14.8) 10.1 (8.0, 12.8) 18.3 (12.2, 27.3) 14.1 (11.0, 18.0) 15.1 (11.6, 19.6)	18.3 (12.2, 27.3)	14.1 (11.0, 18.0)	15.1 (11.6, 19.6)
σ	11.2 (9.7, 12.9)	7.9 (6.1, 10.2)	12.4 (8.7, 17.8)	13.2 (9.9, 17.5) 15.6 (11.8, 20.7)	15.6 (11.8, 20.7)
4	4.7 (3.5, 6.3)	1.9 (1.0, 3.7)	4.7 (2.4, 9.0)	5.8(3.1, 10.8)	12.3 (7.8, 19.6)
5-6	3.6 (2.0, 6.4)	2.9 (1.2, 7.0)	5.2 (1.9, 13.7)	4.2 (0.6. 29.5)	2.8 (0.4, 20.2)
Rate ratio per 1 unit increase	0.71 (0.66, 0.76)	$0.71 \ (0.66, 0.76)  0.61 \ (0.54, 0.69)  0.58 \ (0.47, 0.71)  0.81 \ (0.70, 0.94)  0.84 \ (0.73, 0.98)  0.981 \ (0.73, 0.98) \ (0.73, 0.98) \ (0.73, 0.98) \ (0.73, 0.98) \ (0.73, 0.98) \ (0.73, 0.98$	$0.58\ (0.47,\ 0.71)$	0.81 (0.70, 0.94)	0.84 (0.73, 0.98)
Cardiovascular health category, ICH components					
Poor (0-1)	18.9 (15.9, 22.4)	18.9(15.9,22.4)  18.5(14.1,24.3)  34.5(19.1,62.3)  16.0(11.5,22.3)  20.1(14.4,28.2)	34.5 (19.1, 62.3)	16.0 (11.5, 22.3)	20.1 (14.4, 28.2)
Intermediate (2-3)	12.1 (11.0, 13.3)	9.0 (7.5, 10.7)	14.5 (11.1, 18.9)	$14.5\ (11.1,\ 18.9)  13.7\ (11.4,\ 16.5)  15.3\ (12.7,\ 18.5)$	15.3 (12.7, 18.5)
Ideal (4)	4.4 (3.4, 5.7)	2.2 (1.3, 3.7)	4.8 (2.8, 8.3)	5.6(3.1, 10.1)	10.5 (6.7, 16.4)
Rate ratio per 1 unit increase	$0.52\ (0.46,\ 0.60)$	$0.52\ (0.46, 0.60)  0.39\ (0.31, 0.49)  0.36\ (0.24, 0.53)  0.67\ (0.52, 0.88)  0.73\ (0.56, 0.96)$	$0.36\ (0.24,\ 0.53)$	$0.67\ (0.52,0.88)$	0.73 (0.56, 0.96)

and categorised into poor (0-1 ICH components), intermediate (2-3 ICH components) and ideal ( 4 ICH components)

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b Unable to define incident diabetes rate due to lack of participants meeting criteria for the category NA, not applicable

Table 4

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Baseline ICH level Unadjusted	Unadjusted	Multivariable-adjusted <sup>a</sup>	<i>p</i> MHN	$CA^{a}$	vVa	вAH
No. of ICH components	ts					
0	1 (referent)	1 (referent)	1 (referent)	1 (referent) $b$	1 (referent)	1 (referent)
1	$0.84\ (0.48,1.46)$	0.82 (0.47, 1.43)	0.78 (0.35, 1.73)	$q^{\rm PN}$	1.26 (0.38, 4.12)	0.50(0.17,1.41)
2	0.59 (0.34, 1.01) 0.59 (0.34, 1.01)	$0.59\ (0.34,1.01)$	0.47 (0.21, 1.02) NAb	$q^{\rm PN}$	1.08 (0.34, 3.45)	1.08 (0.34, 3.45) 0.38 (0.14, 1.05)
3	$0.51\ (0.30,0.88)$	$0.51\ (0.30,0.88)$	0.36(0.17,0.80)	$q^{\rm PN}$	1.00 (0.31, 3.21)	$0.39\ (0.14,1.10)$
4	0.21 (0.12, 0.39) 0.22 (0.12, 0.40)	$0.22\ (0.12,0.40)$	$0.09\ (0.03,\ 0.25)$	$q^{\rm PN}$	0.44 (0.12, 1.61)	0.44 (0.12, 1.61) 0.31 (0.10, 0.94)
56	0.16 (0.07, 0.36)	$0.17\ (0.08,\ 0.38)$	0.14~(0.04, 0.45) NAb	$q^{\rm PN}$	0.30 (0.03, 2.90)	$0.07\ (0.01,\ 0.61)$
$p$ for trend $^{\mathcal{C}}$	<0.0001	<0.0001	<0.0001		<0.01	<0.05
Cardiovascular health category, ICH components	category, ICH comp	onents				
Poor (0-1)	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)
Intermediate (2-3)	Intermediate (2-3) 0.64 (0.53, 0.78)	$0.66\ (0.54,0.80)$	0.52 (0.37, 0.72)	0.39 (0.20, 0.76)	$0.52\ (0.37,\ 0.72)  0.39\ (0.20,\ 0.76)  0.85\ (0.58,\ 1.24)$	$0.74\ (0.50,1.09)$
Ideal (4)	0.24 (0.17, 0.32)	0.25 (0.18, 0.35)	0.13 (0.07, 0.24)	$0.12\ (0.05,\ 0.28)$	$0.34\ (0.17,0.68)$	$0.50\ (0.28,\ 0.88)$
$p$ for trend $^{\mathcal{C}}$	<0.0001	<0.0001	<0.0001	<0.0001	<0.01	<0.05

<sup>4</sup> Adjusted for age, education, sex, study site, race/ethnicity, occupational status, alcohol use and estimated glomerular filtration rate; race-stratified analyses were not adjusted for race

 $^{b}$ CA excluded from analysis due to sample size

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 $\stackrel{\mathcal{C}}{p}$  for trend calculated using the logrank test

NA, not applicable

Table 5

$g^d$ 1 (referent)         1 (referent)         1 (referent)         1 (referent)           nediate         0.96 (0.79, 1.16)         0.94 (0.77, 1.13)         0.73 (0.52, 1.03)         1.19 (0.68, 2.09)           0.46 (0.37, 0.58)         0.53 (0.42, 0.68)         0.54 (0.37, 0.58)         0.51 (0.26, 1.00)           nediate         0.52 (0.14, 0.62)         0.54 (0.45, 0.65)         0.56 (0.23, 0.53)         0.51 (0.26, 1.00)           nediate         0.52 (0.17, 0.28)         0.54 (0.45, 0.65)         0.56 (0.41, 0.76)         0.45 (0.53, 0.90)           nobisterol <sup>2</sup> 1 (referent)         1 (referent)         1 (referent)         1 (referent)           nobisterol <sup>2</sup> 1 (referent)         1 (referent)         1 (referent)         1 (referent)           nobisterol <sup>2</sup> 1 (referent)         1 (referent)         1 (referent)         1 (referent)           nodiate         0.90 (0.68, 1.19)         0.87 (0.65, 1.16)         0.72 (0.46, 1.15)         0.67 (0.28, 0.53)           g <sup>a</sup> 1 (referent)         1 (referent)         1 (referent)         1 (referent)           nodiate         0.90 (0.68, 1.16)         0.87 (0.65, 1.18)         0.76 (0.49, 1.18)         0.76 (0.23, 0.28)           g <sup>a</sup> 1 (referent)         1 (referent)         1 (referent) <t< th=""><th>Baseline cardiovascular health component</th><th>Overall</th><th>Overall adjusted<sup>c</sup></th><th>NHWC</th><th><math>CA^{\mathcal{C}}</math></th><th><math>AA^{C}</math></th><th>HA<sup>c</sup></th></t<>	Baseline cardiovascular health component	Overall	Overall adjusted <sup>c</sup>	NHWC	$CA^{\mathcal{C}}$	$AA^{C}$	HA <sup>c</sup>
·       1       (referent)       d       1 (referent)       1 (referent)       1 (referent)         mediate       0.96 (0.79, 1.16)       0.94 (0.77, 1.13)       0.73 (0.52, 1.03)       1.19 (0.68, 2.09)         mediate       0.96 (0.79, 1.16)       0.94 (0.77, 1.13)       0.73 (0.52, 1.03)       0.51 (0.26, 1.00)         rediate       0.46 (0.37, 0.58)       0.53 (0.42, 0.68)       0.36 (0.23, 0.53)       0.51 (0.26, 1.00)         mediate       1 (referent)       1 (referent)       1 (referent)       1 (referent)         mediate       0.22 (0.17, 0.28)       0.25 (0.19, 0.33)       0.24 (0.15, 0.38)       0.16 (0.08, 0.33)         mediate       0.22 (0.17, 0.28)       0.25 (0.19, 0.33)       0.24 (0.15, 0.38)       0.16 (0.08, 0.33)         mediate       0.22 (0.17, 0.28)       0.25 (0.19, 0.33)       0.24 (0.15, 0.38)       0.16 (0.08, 0.33)         mediate       0.22 (0.17, 0.23)       0.25 (0.19, 0.33)       0.24 (0.15, 0.38)       0.16 (0.08, 0.33)         mediate       1 (referent)       1 (referent)       1 (referent)       1 (referent)       1 (referent)         mediate       0.90 (0.68, 1.19)       0.87 (0.55, 1.13)       0.76 (0.40, 1.18)       0.76 (0.32, 1.18)       0.76 (0.32, 1.18)         mediate       0.90 (0.68, 1.10)       0.87 (0.55, 1.23)	$BP^{d}$						
mediate         0.96 (0.79, 1.16)         0.44 (0.77, 1.13)         0.73 (0.52, 1.03)         1.19 (0.68, 2.09)           1         0.46 (0.37, 0.58)         0.53 (0.42, 0.68)         0.56 (0.23, 0.55)         0.51 (0.26, 1.00)          mediate         0.46 (0.37, 0.58)         0.53 (0.42, 0.68)         0.56 (0.21, 0.23)         0.51 (0.26, 1.00)          mediate         0.52 (0.14, 0.62)         0.54 (0.45, 0.65)         0.56 (0.41, 0.76)         0.45 (0.23, 0.33)          mediate         0.22 (0.17, 0.28)         0.24 (0.15, 0.38)         0.16 (0.08, 0.33)           mediate         0.22 (0.17, 0.28)         0.25 (0.19, 0.33)         0.24 (0.15, 0.33)         0.16 (0.08, 0.33)           mediate         0.20 (0.68, 1.19)         0.76 (0.49, 1.18)         1.19 (0.55, 0.53)         0.16 (0.08, 0.33)           mediate         1         1 (referent)         1 (referent)         1 (referent)         1.19 (0.55, 0.53)           mediate         0.90 (0.58, 1.16)         0.37 (0.55, 1.15)         0.76 (0.49, 1.18)         0.76 (0.31, 1.86)           mediate         0.90 (0.68, 1.120)         0.82 (0.55, 1.23)         0.76 (0.31, 1.86)         0.76 (0.31, 1.86)           mediate         0.90 (0.68, 1.120)         0.82 (0.55, 1.23)         0.76 (0.31, 1.86)         0.76 (0.31, 1.86)           mediate <td< td=""><td>Poor</td><td>1 (referent) d</td><td>1 (referent)</td><td>1 (referent)</td><td>1 (referent)</td><td>1 (referent)</td><td>1 (referent)</td></td<>	Poor	1 (referent) d	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)
1 $0.46 (0.37, 0.58)$ $0.53 (0.42, 0.68)$ $0.36 (0.23, 0.55)$ $0.51 (0.26, 1.00)$ mediate $1 (referent)$ $1 (referent)$ $1 (referent)$ $1 (referent)$ mediate $0.52 (0.44, 0.62)$ $0.54 (0.45, 0.65)$ $0.56 (0.41, 0.76)$ $0.45 (0.23, 0.90)$ molesterol <sup>a</sup> $0.22 (0.17, 0.28)$ $0.24 (0.15, 0.38)$ $0.16 (0.08, 0.33)$ holesterol <sup>a</sup> $1 (referent)$ $1 (referent)$ $1 (referent)$ mediate $0.20 (0.58, 1.19)$ $0.27 (0.46, 1.18)$ $1.19 (0.50, 2.83)$ mediate $0.90 (0.68, 1.19)$ $0.87 (0.65, 1.16)$ $0.72 (0.46, 1.18)$ $1.19 (0.50, 2.83)$ mediate $0.90 (0.68, 1.19)$ $0.87 (0.65, 1.16)$ $0.72 (0.46, 1.18)$ $0.67 (0.28, 1.61)$ $mediate$ $0.90 (0.68, 1.19)$ $0.87 (0.65, 1.16)$ $0.72 (0.46, 1.18)$ $0.67 (0.28, 1.61)$ $mediate$ $0.90 (0.68, 1.19)$ $0.87 (0.55, 2.34)$ $0.66 (0.20, 2.83)$ $0.76 (0.31, 1.80)$ $mediate$ $1.30 (0.63, 2.69)$ $1.13 (0.55, 2.34)$ $0.66 (0.20, 2.18)$ $0.76 (0.31, 1.80)$ $mediate$ $0.97 (0.77, 1.22)$ $0.99 (0.78, 1.26)$ $0.82 (0.55, 1.23)$ $0.76 (0.31, 1.80)$ $mediate$ $0.97 (0.77, 1.22)$ $0.99 (0.78, 1.26)$ $0.82 (0.55, 1.23)$ $0.76 (0.3$	Intermediate	0.96 (0.79, 1.16)	0.94 (0.77, 1.13)	0.73 (0.52, 1.03)	1.19 (0.68, 2.09)	0.87 (0.61, 1.23)	$1.26\ (0.84,1.88)$
$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$	Ideal	$0.46\ (0.37,0.58)$	$0.53\ (0.42,0.68)$	$0.36\ (0.23,\ 0.55)$	0.51 (0.26, 1.00)	$0.54\ (0.33,0.89)$	$0.88\ (0.55,1.41)$
Interestion       Interestion <thinterestion< th=""> <thinterestion< th=""></thinterestion<></thinterestion<>	BMI <sup>a</sup>						
inate $0.22 (0.41, 0.62) 0.54 (0.45, 0.65) 0.56 (0.41, 0.76) 0.45 (0.23, 0.90) 0.22 (0.17, 0.28) 0.25 (0.19, 0.33) 0.24 (0.15, 0.38) 0.16 (0.08, 0.33)$ sterol <sup>2</sup> interval 1 (referent)	Poor	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)
	Intermediate	0.52 (0.44, 0.62)	$0.54\ (0.45,0.65)$	$0.56\ (0.41,\ 0.76)$	0.45 (0.23, 0.90)	0.55 (0.39, 0.78)	$0.42\ (0.30,0.59)$
	Ideal	0.22 (0.17, 0.28)	$0.25\ (0.19,0.33)$	$0.24\ (0.15,\ 0.38)$	0.16(0.08,0.33)	0.23 (0.12, 0.43)	0.21 (0.11, 0.39)
Interfacent       Interfacent <thinterfacent< th=""> <thinterfacent< th=""></thinterfacent<></thinterfacent<>	Total cholesterol <sup>a</sup>						
ifate $1.04 (0.75, 1.39) 1.00 (0.75, 1.33) 0.76 (0.49, 1.18) 1.19 (0.50, 2.83) 0.90 (0.68, 1.19) 0.87 (0.65, 1.16) 0.72 (0.46, 1.15) 0.67 (0.28, 1.61) 0.90 (0.68, 1.19) 0.87 (0.65, 1.16) 0.72 (0.46, 1.15) 0.67 (0.28, 1.61) 1 (referent) 1 (referent) 1 (referent) 1 (referent) 1 (referent) 0.97 (0.77, 1.22) 0.99 (0.78, 1.25) 0.66 (0.20, 2.18) NAP 0.97 (0.77, 1.22) 0.99 (0.78, 1.25) 0.82 (0.55, 1.23) 0.76 (0.31, 1.86) 0.97 (0.77, 1.22) 0.99 (0.78, 1.25) 0.82 (0.55, 1.23) 0.76 (0.31, 1.86) 0.97 (0.77, 1.22) 0.99 (0.78, 1.25) 0.82 (0.55, 1.23) 0.76 (0.31, 1.86) 1 (referent) 1$	Poor	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Intermediate	1.04 (0.79, 1.39)	1.00 (0.75, 1.33)	0.76 (0.49, 1.18)	1.19 (0.50, 2.83)	1.19 (0.63, 2.26)	$1.06\ (0.61,1.85)$
I (referent)     I (referent)     I (referent)     I (referent)       iiate     1.30 (0.63, 2.69)     1.13 (0.55, 2.34)     0.66 (0.20, 2.18) $N_A b$ $0.97 (0.77, 1.22)$ 0.99 (0.78, 1.26)     0.82 (0.55, 1.23)     0.76 (0.31, 1.86)       ake <sup>d</sup> 1     (referent)     1 (referent)     1 (referent)       iate     0.97 (0.77, 1.22)     0.99 (0.78, 1.26)     0.82 (0.55, 1.23)     0.76 (0.31, 1.86)       iate     0.98 (0.83, 1.16)     1 (referent)     1 (referent)     1 (referent)       iate     0.98 (0.83, 1.16)     1.15 (0.97, 1.37)     1.13 (0.83, 1.54)     1.21 (0.74, 1.96)       inite     0.98 (0.83, 1.16)     1.15 (0.97, 1.37)     1.49 (0.46, 4.80)     0.64 (0.08, 4.96)       civity <sup>d</sup> 1.53 (0.67, 2.25)     1.70 (0.92, 3.12)     1.49 (0.46, 4.80)     0.54 (0.08, 4.96)	Ideal	0.90 (0.68, 1.19)	0.87 (0.65, 1.16)	0.72 (0.46, 1.15)	0.67 (0.28, 1.61)	$0.84\ (0.44,1.61)$	1.20 (0.69, 2.08)
1 (referent)       1 (referent)       1 (referent)       1 (referent) $1.30 (0.63, 2.69)$ $1.13 (0.55, 2.34)$ $0.66 (0.20, 2.18)$ $N_A b$ $0.97 (0.77, 1.22)$ $0.99 (0.78, 1.26)$ $0.82 (0.55, 1.23)$ $0.76 (0.31, 1.86)$ $0.97 (0.77, 1.22)$ $0.99 (0.78, 1.26)$ $0.82 (0.55, 1.23)$ $0.76 (0.31, 1.86)$ $1 (referent)$ $1 (referent)$ $1 (referent)$ $1 (referent)$ $1 (referent)$ $0.98 (0.83, 1.16)$ $1.15 (0.97, 1.37)$ $1.13 (0.83, 1.54)$ $1.21 (0.74, 1.96)$ $1.23 (0.67, 2.25)$ $1.70 (0.92, 3.12)$ $1.49 (0.46, 4.80)$ $0.64 (0.08, 4.96)$	Smoking <sup>a</sup>						
1.30 (0.63, 2.69)       1.13 (0.55, 2.34) $0.66 (0.20, 2.18)$ $N_A b$ $0.97 (0.77, 1.22)$ $0.99 (0.78, 1.26)$ $0.82 (0.55, 1.23)$ $0.76 (0.31, 1.86)$ 1 (referent)       1 (referent)       1 (referent) $1 (referent)$ 0.98 (0.83, 1.16)       1.15 (0.97, 1.37) $1.13 (0.83, 1.54)$ $1.21 (0.74, 1.96)$ 1.23 (0.67, 2.25) $1.70 (0.92, 3.12)$ $1.49 (0.46, 4.80)$ $0.64 (0.08, 4.96)$	Poor	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)
0.97 (0.77, 1.22)       0.99 (0.78, 1.26)       0.82 (0.55, 1.23)       0.76 (0.31, 1.86)         1 (referent)       1 (referent)       1 (referent)       1 (referent)         0.98 (0.83, 1.16)       1.15 (0.97, 1.37)       1.13 (0.83, 1.54)       1.21 (0.74, 1.96)         1.23 (0.67, 2.25)       1.70 (0.92, 3.12)       1.49 (0.46, 4.80)       0.64 (0.08, 4.96)	Intermediate	1.30 (0.63, 2.69)	1.13 (0.55, 2.34)	$0.66\ (0.20,\ 2.18)$	$^{q}$ NA $^{b}$	1.64 (0.38, 7.05)	1.92 (0.57, 6.46)
1 (referent)       1 (referent)       1 (referent)         0.98 (0.83, 1.16)       1.15 (0.97, 1.37)       1.13 (0.83, 1.54)       1.21 (0.74, 1.96)         1.23 (0.67, 2.25)       1.70 (0.92, 3.12)       1.49 (0.46, 4.80)       0.64 (0.08, 4.96)	Ideal	0.97 (0.77, 1.22)	0.99 (0.78, 1.26)	0.82 (0.55, 1.23)	0.76(0.31,1.86)	1.11 (0.71, 1.74)	1.00 (0.62, 1.62)
1 (referent)     1 (referent)     1 (referent)     1 (referent)       0.98 (0.83, 1.16)     1.15 (0.97, 1.37)     1.13 (0.83, 1.54)     1.21 (0.74, 1.96)       1.23 (0.67, 2.25)     1.70 (0.92, 3.12)     1.49 (0.46, 4.80)     0.64 (0.08, 4.96)	Dietary intake <sup>a</sup>						
0.98 (0.83, 1.16) 1.15 (0.97, 1.37) 1.13 (0.83, 1.54) 1.21 (0.74, 1.96) 1.23 (0.67, 2.25) 1.70 (0.92, 3.12) 1.49 (0.46, 4.80) 0.64 (0.08, 4.96)	Poor	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)
1.23 (0.67, 2.25) 1.70 (0.92, 3.12) 1.49 (0.46, 4.80) 0.64 (0.08, 4.96) 1 (metanovi) 1 (metanovi)	Intermediate	0.98 (0.83, 1.16)	1.15 (0.97, 1.37)	1.13 (0.83, 1.54)	1.21 (0.74, 1.96)	$1.10\ (0.79,1.53)$	1.19 (0.86, 1.66)
1 (metruant) 1 (metruant)	Ideal	1.23 (0.67, 2.25)	1.70 (0.92, 3.12)	$1.49\ (0.46, 4.80)$	0.64 (0.08, 4.96)	2.55 (1.03, 6.47)	1.17 (0.28, 4.89)
1 (metramet) 1 (metramet) 1 (metramet)	Physical activity <sup>a</sup>						
	Poor	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)	1 (referent)
Intermediate 0.95 (0.73, 1.24) 1.08 (0.83, 1.41) 1.30 (0.83, 2.03) 1.48 (0.72, 3.06)	Intermediate	0.95 (0.73, 1.24)	$1.08\ (0.83,\ 1.41)$	$1.30\ (0.83,\ 2.03)$	1.48 (0.72, 3.06)	1.11 (0.63, 1.93)	0.71 (0.40, 1.25)
Ideal 0.82 (0.67, 1.00) 0.98 (0.80, 1.19) 0.83 (0.57, 1.21) 1.23 (0.70, 2.16)	Ideal	$0.82\ (0.67,1.00)$	$0.98\ (0.80,1.19)$	0.83 (0.57, 1.21)	1.23 (0.70, 2.16)	1.26 (0.84, 1.90)	0.95 (0.66, 1.36)

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 $^b\mathrm{CA}$  excluded from analysis due to sample size

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<sup>c</sup>Adjusted for age, education, sex, study site, race/ethnicity, occupational status, alcohol use, estimated glomerular filtration rate and other cardiovascular health components; race-stratified analysis was not adjusted for race

dHRs calculated using Cox proportional hazards modelling with the poor category as the referent group NA, not applicable

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Cardiovascular health	Cardiovascular health Participants, $\%$ ( $n=5,348$ )	HR <sup>a</sup> (95% CI)	HR <sup>b</sup> (95% CI)	$HR^{a} (95\% \text{ CI}) HR^{b} (95\% \text{ CI}) Population-attributable risk, \%^{a} (95\% \text{ CI}) Population-attributable risk, \%^{b} (95\% \text{ CI})$	Population-attributable risk, $\%^b$ (95% CI)
No. of baseline ICH components	ponents				
2	32.0	$0.70\ (0.56,\ 0.87)$		23 (9, 35)	
3	30.7	0.77 (0.64, 0.92)		17 (6, 28)	
4	17.7	0.37 (0.27, 0.50)		58 (45, 69)	
5	5.4	$0.34\ (0.19,0.63)$		64 (36, 80)	
Cardiovascular health category, ICH components	egory, ICH components				
Intermediate (2-3)	62.8		$0.66\ (0.54,\ 0.80)$		16 (9, 24)
Ideal (4)	23.1		0.35 (0.27, 0.46)		59 (47, 68)

to a specified number of baseline ICH components. Analyses were adjusted for age, education, sex, study site, race/ethnicity, occupational status, alcohol use and estimated glomerular filtration rate

b Compared with participants meeting criteria for lower cardiovascular health status. The population-attributable risk is the percentage of new cases of diabetes in the population attributable to not meeting criteria for a specified cardiovascular health category. Analyses were adjusted for age, education, sex, study site, race/ethnicity, occupational status, alcohol use and estimated glomerular filtration rate