Haptoglobin 2-2 Genotype is Not Associated With Cardiovascular Risk in Subjects With Elevated Glycohemoglobin—Results From the Bruneck Study

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Background—Haptoglobin (Hp) is an abundant plasma protein with antioxidant properties. The Hp 2-2 genotype has previously been linked to coronary heart disease risk in individuals with elevated glycosylated hemoglobin (HbA1c). We investigated the association of Hp and HbA1c with cardiovascular disease (CVD) in the longitudinal, population-based Bruneck Study.

Methods and Results—Hp genotype was determined by polymerase chain reaction according to standard procedures and HbA1c concentration by a Diabetes Control and Complications Trial-aligned assay. HbA1c was measured in 1995, 2000, and 2005. Occurrence of the combined CVD endpoint of myocardial infarction or stroke was recorded between 1995 and 2010. Outcome analyses employed the Cox proportional hazards model with HbA1c category as time-varying covariate. At baseline in 1995, 806 subjects (male sex, 49.3%; age, mean±standard deviation, 62.70±11.08 years) were included. During follow-up, 123 subjects experienced at least 1 CVD event (48 suffered myocardial infarction, 68 stroke, and 7 both). Among subjects with HbA1c≥6.5% (≥48 mmol/mol), those with the Hp 2-2 genotype did not show an elevated risk of incident CVD compared with those with other genotypes (age- and sex-adjusted hazard ratio [95% CI], 0.47 [0.19, 1.13], P=0.092) and a null association was also observed in subjects with HbA1c<6.5% (1.10 [0.75, 1.62], P=0.629) (P for interaction=0.082).

Conclusions—Subjects with the Hp 2-2 genotype and elevated HbA1c compared with subjects with other Hp genotypes and elevated HbA1c did not show increased CVD risk. (J Am Heart Assoc. 2014;3:e000732 doi: 10.1161/JAHA.113.000732)

Key Words: cardiovascular diseases • diabetes mellitus • genetics
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featured by a high level of oxidative stress, relevance of the antioxidant Hp may be considerably higher in diabetic patients.\textsuperscript{11} Actually, in a meta-analysis of 5 studies involving 1829 patients with diabetes, the pooled odds ratio (95% CI) for CVD was 2.03 (1.46, 2.81) in a comparison of the Hp 2-2 genotype with other genotypes.\textsuperscript{12} In line with this finding, Hp 2-2 was associated with incident CVD in type 2 diabetes mellitus patients (odds ratio [95% CI], 4.96 [1.85, 13.33]). In conflict with these findings, the Framingham Offspring Study reported that Hp 2-2 was associated with significantly lower CAD prevalence among diabetic patients than Hp 1-1 (odds ratio [95% CI], 0.43 [0.21, 0.90]).\textsuperscript{13}

As a potential explanation for the substantial heterogeneity in the Hp - CVD association between the various studies effect modification by levels of HbA1c has been proposed.\textsuperscript{14} Indeed, the reported dysfunction of Hp 2-2 in preventing Hb-mediated oxidative damage was found to be accentuated with glycosylated Hb (HbA1c) in cell culture experiments.\textsuperscript{4} Testing this hypothesis in humans, Cahill and colleagues\textsuperscript{14} reported the Hp 2-2 genotype to be associated with elevated CAD risk among subjects with elevated HbA1c in the Nurses’ Health Study (odds ratio [95% CI], 10.12 [1.08, 94.97]) and validated their finding in a cohort of type 2 diabetic subjects (hazard ratio [95% CI], 7.55 [2.79, 20.47]). Based on this and earlier studies, Hp genotyping among diabetic patients and antioxidant treatment in diabetic patients with the Hp 2-2 genotype has been propagated\textsuperscript{2,12} but further confirmatory data are required to change clinical routine. Given its potentially large impact on diabetes management, we investigated the association of Hp genotype with CVD conditional on elevated HbA1c in the prospective, population-based Bruneck Study.

Methods

Study Population and Data Collection

The Bruneck Study is a prospective, population-based survey on the epidemiology and pathogenesis of atherosclerosis and cardiovascular disease.\textsuperscript{15-18} At baseline in 1990 the study population comprised an age- and sex-stratified random sample of all inhabitants of Bruneck (125 men and 125 women from each of the fifth through eighth decades of age, for an age- and sex-stratified random sample of n=1000, all of western European descent). No subjects were enrolled after study initiation. In 1995, 826 subjects participated in the first quinquennial re-examination and DNA samples for genotyping were available in 816 individuals. During follow-up from 1995 to 2010, detailed information about fatal and nonfatal new-onset CVD was carefully collected. Follow-up was 100% complete for clinical endpoints, which was made possible by the extremely low population mobility of 0.2% in the Bruneck area. The study protocol was approved by the ethics committees of Bolzano and Verona and conformed to the Declaration of Helsinki. All study subjects provided written informed consent. Risk factors were assessed by means of validated standard procedures as described previously.\textsuperscript{15,19}

At the baseline and follow-up examinations in 1995, 2000, 2005, and 2010, venous blood was sampled in the morning after an overnight fast for laboratory measurements, including fasting plasma glucose and HbA1c (Diabetes Control and Complications Trial-aligned assay; equipment and reagents from BioRad, Milan, Italy, at both baseline and follow-up examinations). In accordance with the study of Cahill et al, HbA1c was dichotomized according to a cut-off of 6.5% (48 mmol/mol, International Federation of Clinical Chemistry units).

Diabetes mellitus was coded present for subjects with fasting glucose levels $\geq$7 mmol/L ($\geq$126 mg/dL) or a medical record confirmed prediagnosis of definite disease status. Based on the literature\textsuperscript{14} we anticipated few subjects with the Hp 1-1 genotype and, to maximize statistical power, decided prior to analysis to pool the Hp 1-1 and 2-1 genotypes, forming a group of Hp1 allele carriers, which is a common approach.\textsuperscript{14}

We ascertained leisure time-related physical activity by a standardized questionnaire\textsuperscript{20} rating the intensity of activities according to the compendium of physical activities\textsuperscript{21} and calculated average metabolic equivalent hours per week to estimate long-term physical activity. We assessed food intake by a standardized food-frequency questionnaire (FFQ) based on the gold standard Harvard FFQ by Willett and colleagues and adapted the FFQ to the dietary peculiarities in the survey area. We validated our FFQ with a dietician-supervised short-term assessment of food intake. Validity was high and similar to that previously found for the same FFQ in other populations.\textsuperscript{22} The Alternative Healthy Eating Index (AHEI) was calculated from these data as a quantitative measure of healthy dietary behavior.\textsuperscript{23} Body mass index (BMI) was calculated as weight in kilograms over height in meters squared.

Haptoglobin Genotyping

The Haptoglobin genotype was determined by PCR as described previously\textsuperscript{24} with slight modifications. In brief, 20 $\mu$L reactions contained 20 ng genomic DNA, 2 units Qiagen Taq Polymerase (Qiagen), 1 $\times$ Qiagen PCR buffer (Qiagen, Hilden, Germany), 1 $\times$ Q solution (Qiagen), 200 $\mu$mol/L of each dNTP (Peqlab) and 0.25 $\mu$mol/L of each primer (A/B or C/D). Oligonucleotide primers A and B were used for amplification of a 1757-bp Hp 1 allele-specific sequence and a 3481-bp Hp 2 allele-specific sequence. Primers C and D were used to amplify a 349-bp Hp 2 allele-specific sequence. The primers A (5’-GAGGGAGCTTGCCCTTTQ-3’), B (5’-GGCTTTGGAATGGCCGAAG-3’), C (5’-GGCTTTGGAATGGCCGAAG-3’), D (5’-GGCTTTGGAATGGCCGAAG-3’) (48 mmol/mol, International Federation of Clinical Chemistry units).

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CCATTG-3'), B (5'-GAGATTITGAGGCCCTGCTGT-3'), C (5'-CCTGCCCCGATTAATCGCCATT-3') and D (5'-CCGAGTGCTC-CACCATGCGCTGT-3') were purchased from Microsynth. The amplification reactions were conducted on a DNA Engine Cycler (BioRad) under the following conditions: initial denaturation 3 minutes 94°C; 94°C 30 seconds, 57°C (primers A/B) and 62°C (primers C/D) 30 seconds, 72°C 2 minutes, 35 cycles; final extension 10 minutes 72°C. After amplification 8 μL PCR product A/B and 2 μL PCR product C/D were mixed and separated together on a 1% agarose gel.

Endpoints

The composite CVD endpoint included incident fatal and non-fatal myocardial infarction and stroke. Presence of myocardial infarction was assessed by World Health Organization criteria (definite disease status), while stroke was classified according to the criteria of the National Survey of Stroke. Events were ascertained by careful review of medical records provided by general practitioners, death certificates, and Bruneck Hospital files. A major advantage of the Bruneck Study is that virtually all inhabitants of Bruneck are referred to 1 local hospital that cooperates closely with the general practitioners. This allows retrieval of complete medical information.

Subjects who had experienced CVD events before the study baseline in 1995 were included in the main analysis and observed for recurring incident CVD between 1995 and 2010, and excluded in a sensitivity analysis.

Statistical Analysis

Hardy-Weinberg equilibrium was tested by a permutation-based chi-squared test. Associations with incident CVD were assessed by Cox proportional hazards regression with HbA1c or diabetes status (present/absent) as time-varying covariates in a granularity of 5 years. This approach relates the most current measure of glycemic exposure to incident CVD and avoids potential confounding due to reliance on a single baseline measurement. Effect modification of Hp 2-2 by HbA1c category was tested using an appropriate interaction term. To simultaneously obtain marginal effects of Hp genotype for both HbA1c groups as well as the interaction effect, a linear combination of model parameters was made. The proportional hazards assumption was tested by computing the correlation coefficient of survival time with scaled Schoenfeld residuals and was met. Base models were adjusted for age and sex (model 1) and 2 multivariable models with progressive adjustment were employed (model 2: additionally for current smoking, systolic blood pressure, low-density lipoprotein (LDL) cholesterol, and high-density lipoprotein (HDL) cholesterol; model 3: additionally for metabolic equivalent hours, alternative healthy eating index, statin use, and body mass index). Last observation carried forward or next observation carried backward imputation reduced the proportion of missing HbA1c values from 5.9% to 0.4%.

Sensitivity analyses excluded subjects with missing HbA1c values, used age as the time scale and focused on individual disease endpoints. All tests were 2-sided and P values smaller than 0.05 were considered significant. All analyses were performed using the R statistical package.

Results

Characteristics of the study population are shown in Table 1. Hp genotyping resulted in unambiguous results for 810 of 816 subjects for which DNA samples were available (Call rate, 99.3%). Of these, 4 had missing values in HbA1c concentration, which resulted in a baseline study size of 806 subjects. Duplicate measurement of 24 DNA samples yielded concordant findings in all cases. Genotypes were distributed as follows: Hp 1-1, 10.3% (n=83), Hp 2-1, 41.7% (n=336), Hp 2-2, 48.0% (n=387) and were in Hardy-Weinberg equilibrium overall (P=0.46), in subgroups of subjects with diabetes (P=0.13) and without (P=0.79), and subjects younger than 65 (P=0.52) and subjects at least 65 years old (P=0.80). There were no significant differences in prevalent diabetes (1995 baseline) or in incident diabetes (follow-up 1995-2010) between dichotomized Hp genotypes (P=0.71 and 0.92, respectively). For repeated measurements of HbA1c concentration we found an intra-class correlation coefficient [95% CI] of 0.60 [0.56, 0.63].

The analyses reported in the following were adjusted for age and sex, unless specified otherwise, and utilize updates of HbA1c levels during follow-up (time-varying covariate). From 1995 to 2010, 123 subjects experienced at least 1 CVD event (48 suffered myocardial infarction(s), 68 stroke(s), and 7 both). Only the first CVD event was considered in the main analysis. No significant differences emerged when examining the risk of CVD by Hp genotype (hazard ratio [HR] [95% CI] for Hp 2-2 versus Hp 2-1/1-1: 0.98 [0.69, 1.40]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure). This result did not materially change upon adjustment for conventional risk factors (age and sex, unless specified otherwise). There were no significant differences among subjects with HbA1c≥6.5% (HR [95% CI], 0.47 [0.19, 1.13]; P=0.092) (Figure).
Table 1. Characteristics of the Study Population According to Haptoglobin Genotype

| Measurements taken in 1995 | 1-1 | 2-1 | 2-2 | P Value |
|---------------------------|-----|-----|-----|---------|
| n (%)                     | 83 (10.3) | 336 (41.7) | 387 (48.0) | 0.417 |
| Age, y                    | 63.6±12.6 | 62.1±10.3 | 63.0±11.4 | 0.342 |
| Male sex, n (%)           | 40 (48.2) | 175 (21.2) | 180 (46.5) | 0.496 |
| HbA1c, %                  | 5.3 (5.1 to 5.7) | 5.4 (5.1 to 5.8) | 5.4 (5.1 to 5.8) | 0.575 |
| HbA1c≥6.5%, n (%)         | 3 (3.6) | 15 (4.5) | 23 (5.9) | 0.481 |
| Diabetes, n (%)           | 12 (14.5) | 31 (9.2) | 44 (11.4) | 0.122 |
| Hemoglobin, g/dL          | 14.1±1.3 | 14.3±1.2 | 14.3±1.3 | 0.140 |
| Glucose, mg/dL            | 98.0 (91.0 to 109.5) | 97.0 (90.0 to 104.0) | 98.0 (91.0 to 107.0) | 0.067 |
| Current smoking, n (%)    | 13 (15.7) | 68 (20.7) | 74 (19.5) | 0.714 |
| Total cholesterol, mg/dL  | 227.7±41.3 | 225.6±38.9 | 234.4±45.5 | 0.036 |
| HDL cholesterol, mg/dL    | 57.4±13.9 | 58.5±15.7 | 59.3±17.0 | 0.623 |
| Systolic BP, mm Hg        | 145.3±21.4 | 148.1±20.3 | 149.00±20.9 | 0.225 |
| Current smoking, n (%)    | 70 (84.3) | 268 (79.8) | 311 (80.4) | 0.707 |
| Body mass index, kg/m²    | 26.0±4.4 | 25.9±3.6 | 25.4±3.9 | 0.214 |
| AHEI, score               | 40.6±9.00 | 39.2±8.8 | 39.2±8.6 | 0.317 |
| MET, h/wk                 | 42.0 (30.4) | 42.0 (35.6) | 42.0 (38.3) | 0.782 |
| Statin use, n (%)         | 1 (1.2) | 7 (2.1) | 16 (4.2) | 0.170 |
| Prior CVD, n (%)          | 8 (9.6) | 21 (6.2) | 19 (4.9) | 0.304 |
| Measurements taken in 2000 |     |       |       |         |
| n (%)                     | 68 (9.7) | 300 (42.8) | 333 (47.5) | 0.895 |
| Age, y                    | 60.7±11.5 | 60.7±9.7 | 61.1±10.6 | 0.067 |
| Male sex, n (%)           | 27 (39.7) | 154 (51.3) | 144 (43.2) | 0.155 |
| HbA1c, %                  | 5.7 (5.5 to 5.9) | 5.7 (5.5 to 6.0) | 5.7 (5.5 to 6.0) | 0.064 |
| HbA1c≥6.5%, n (%)         | 3 (4.4) | 17 (5.7) | 34 (10.2) | 0.198 |

Values are presented as mean±standard deviation, median (interquartile range), or count (percentage). P values are for tests of any difference between the 3 groups under adjustment for age and sex. AHEI indicates alternative healthy eating index; BP, blood pressure; CVD, cardiovascular disease; HbA1c, glycosylated hemoglobin; HDL, high-density lipoprotein; MET, metabolic equivalent.

Finally, Hp 2-2 was not associated with CVD risk in subjects with diabetes (P=0.944) nor in those without (P=0.950), and no statistical interaction existed between diabetes and Hp genotype (P=0.927).

Sensitivity analyses excluding subjects with missing HbA1c values, excluding subjects with prior CVD or using age as the time scale (instead of time-on-study) yielded similar findings (Table 2) as did analyses on individual disease endpoints (HR [95% CI], 0.97 [0.25, 3.76], P=0.950). These results are shown in Table 3.

Finally, among those with HbA1c≥6.5%, risk estimates were almost identical for Hp 2-1 compared with Hp 1-1 subjects (HR [95% CI] for stroke: 0.48 [0.16, 1.47], P=0.198; HR [95% CI] for myocardial infarction: 0.59 [0.16, 2.20], P=0.433; Hp 2-2 versus other). Subgroup analyses according to sex and age (Table 2) should be interpreted cautiously given limited sample sizes in these groups.

Finally, among those with HbA1c≥6.5%, risk estimates were almost identical for Hp 2-1 compared with Hp 1-1 subjects (HR [95% CI], 0.97 [0.25, 3.76], P=0.950) providing a post-hoc justification for the dichotomization of Hp genotypes applied in the current study (Hp 2-2 versus 2-1/1-1).

We also investigated differences in blood lipids by Hp genotype. Total cholesterol and LDL cholesterol were higher in Hp 2-2 subjects than in those with other genotypes when adjusting for age, sex, and body mass index (P=0.008 and P=0.016, respectively). These results are shown in Table 3.

Discussion

Our data do not confirm the recent report of Cahill and colleagues that identified the Hp 2-2 genotype as a cardiovascular risk factor among subjects with elevated HbA1c.14

Table 1. Continued

| Measurements taken in 2005 | 1-1 | 2-1 | 2-2 | P Value |
|---------------------------|-----|-----|-----|---------|
| n (%)                     | 61 (10.1) | 256 (42.6) | 284 (47.3) | 0.086 |
| Age, y                    | 59.6±10.9 | 59.7±9.5 | 59.3±9.7 | 0.327 |
| Male sex, n (%)           | 25 (41.0) | 124 (48.4) | 122 (43.0) | 0.359 |
| HbA1c, %                  | 5.5 (5.4 to 5.7) | 5.6 (5.4 to 5.9) | 5.6 (5.4 to 5.9) | 0.359 |
| HbA1c≥6.5%, n (%)         | 3 (4.9) | 20 (7.8) | 28 (9.9) | 0.328 |
| Diabetes, n (%)           | 6 (9.8) | 34 (13.3) | 36 (12.7) | 0.787 |
Part of this discrepancy may derive from differences in study design and population. Cahill used a nested case-control design with 1:1 matching in female nurses with 14-year follow-up for their main analysis, and data from a clinical trial with 18 months of follow-up performed in type 2 diabetic patients for confirmation, whereas we used a 15-year prospective observational design on a random sample of the general population. Importantly, Cahill excluded participants with prior CVD, which we did not, and used a coronary heart disease (CHD) endpoint, while we used a compound of myocardial infarction and stroke in the main analysis. Another potential explanation for the discrepancy between our and Cahill’s results is survival bias. However, we observed a higher frequency of Hp 2-2 genotype (48%) than most previous studies in Caucasians did (36 to 40%), which is the opposite of what would be expected in the presence of a survival disadvantage. Furthermore, genotypes were in Hardy-Weinberg equilibrium overall as well as in the younger and older half of our sample separately, which argues against the existence of significant survival bias.

Our results are unexpected because Hp 2-2 as a risk genotype among diabetic subjects would be backed by a biologically plausible rationale based on its weaker antioxidant properties. However, not all aspects of this rationale are fully consistent. In particular, the Hp 2-2-Hb compared with the Hp 1-1-Hb complex has been reported to be taken up into macrophages via the CD163 scavenger receptor at higher rates as well as at lower rates. No differences in Hb binding and antioxidant potency between purified human Hp 1-1 and Hp 2-2 were recently found in an animal model of hemolysis. Hp has numerous functions apart from Hb clearance, taking part in inflammatory pathways (prostaglandin synthesis, cathepsin B activity, endothelium-dependent vasodilation) and interfering with the functions of immune cells. Hp phenotypes differ in some of these functions. Notably, Hp 2-2 is the most angiogenic phenotype, which was discussed as an explanation for longer walking distance in Hp 2-2 peripheral vascular disease patients, and Hp 1-1 was linked to decreased endothelial repair potential in lacunar stroke patients.

In this study, subjects with the Hp 2-2 genotype had elevated total and LDL cholesterol (Table 3), which has been reported previously. This finding may be explained by the close genetic linkage between Hp genotype and a single nucleotide polymorphism in the gene encoding haptoglobin-related protein, which is associated with levels of total but not HDL cholesterol via apolipoprotein L.
Table 2. Subgroup Analyses, Endpoint-Specific Analyses, and Sensitivity Analyses

| HbA1c Group | Model 1 | Model 2 | Model 3 |
|-------------|---------|---------|---------|
|             | HR [95% CI] | P Value | P Interaction | HR [95% CI] | P Value | P Interaction | HR [95% CI] | P Value | P Interaction |
| Full sample | <6.5%     | 1.10 [0.75, 1.62] | 0.629 | 0.082 | 1.11 [0.74, 1.66] | 0.623 | 0.117 | 1.13 [0.75, 1.69] | 0.563 | 0.149 |
|            | ≥6.5%     | 0.47 [0.19, 1.13] | 0.092 | 0.48 [0.18, 1.26] | 0.136 | 0.52 [0.19, 1.39] | 0.189 |

**Subgroups**

**Age <75 y**

- <6.5%: 0.98 [0.59, 1.63] (0.949) 0.092
- ≥6.5%: 0.79 [0.26, 2.35] (0.666)

**Age ≥75 y**

- <6.5%: 1.22 [0.65, 2.27] (0.536) 0.035
- ≥6.5%: 0.11 [0.01, 0.95] (0.044)

**Men**

- <6.5%: 0.98 [0.59, 1.62] (0.938) 0.033
- ≥6.5%: 0.97 [0.24, 3.92] (0.963) 0.044

**Women**

- <6.5%: 1.36 [0.72, 2.55] (0.039) 0.018
- ≥6.5%: 0.27 [0.08, 0.88] (0.030)

**Individual endpoints**

**Stroke**

- <6.5%: 0.97 [0.53, 1.61] (0.601) 0.037
- ≥6.5%: 0.87 [0.46, 1.65] (0.198)

**Myocardial infarction**

- <6.5%: 1.41 [0.84, 2.31] (0.165) 0.021
- ≥6.5%: 1.51 [0.94, 2.45] (0.198)

**Sensitivity analyses**

**Age as time scale**

- <6.5%: 1.09 [0.74, 1.61] (0.659) 0.014
- ≥6.5%: 0.97 [0.50, 1.82] (0.125)

**No HbA1c imputation**

- <6.5%: 1.08 [0.72, 1.61] (0.708) 0.025
- ≥6.5%: 0.60 [0.24, 1.52] (0.281)

**Subjects with prior CVD excluded**

- <6.5%: 1.24 [0.80, 1.90] (0.332) 0.030
- ≥6.5%: 0.55 [0.21, 1.44] (0.222)

**HR [95% CI] and P value are for the comparison of Hp 2-2 versus other genotypes among subjects with glycosylated hemoglobin (HbA1c) <6.5% or HbA1c ≥6.5%, respectively. PInteraction is for a difference in the effect of dichotomized Hp genotype between those with HbA1c <6.5% and those with HbA1c ≥6.5%. Model 1: adjustment for age and sex; Model 2: as model 1, with additional adjustment for current smoking, systolic blood pressure, LDL cholesterol, and HDL cholesterol; Model 3: as model 2, with additional adjustment for metabolic equivalent hours, alternative healthy eating index, statin use, and body mass index. CVD indicates cardiovascular disease; HbA1c, glycosylated hemoglobin; HR, hazard ratio.**
adjusting for LDL levels did not appreciably change risk estimates (Table 2), which argues against the possibility that CVD risk differences due to Hp genotype are mediated by LDL levels.

Strengths of the present study include the virtually complete, long-term follow-up, detailed characterization of the study population and repeated measurements of HbA1c. Another major strength of the Bruneck Study is its high representativity for the general population. It comprises predominantly low- and medium-risk individuals, which are of the foremost public health interest since most CVD events happen in such individuals. One downside of this is a comparatively low prevalence of elevated HbA1c, which precludes more complex analyses like 3-way interactions between HbA1c, diabetes, and Hp genotype. It merits attention that while we found a trend towards protective effects of the Hp 2-2 genotype among subjects with elevated glycohemoglobin, this result was statistically not significant, and hazardous effects would also be compatible with our data. The largest hazardous effect that we cannot refute at the α=0.05 level is a risk elevation of ≈26% (multivariable model 2, Figure). Larger studies are required to draw definite conclusions.

In summary, this study does not confirm that the Hp 2-2 genotype is associated with a higher CVD risk in subjects with elevated HbA1c.

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Disclosures

None.

Table 3. Blood Lipid Levels (mg/dL) by Haptoglobin Genotype

| Lipid parameter     | Hp 1-1 | Hp 2-1 | Hp 2-2 | P_{Hp2-2 vs other} |
|---------------------|--------|--------|--------|-------------------|
| Total cholesterol   | 227.7±41.3 | 225.6±38.8 | 234.4±45.5 | 0.028             |
| HDL                 | 57.4±13.9  | 58.5±15.7  | 59.3±17.0  | 0.788             |
| LDL                 | 144.1±35.6  | 141.6±35.7  | 148.5±40.3  | 0.053             |
| Triglycerides       | 143.8±116.1 | 126.7±66.7  | 133.3±80.6  | 0.075 | 0.189 |

Values are presented as mean±standard deviation. P values are adjusted for age, sex, and body mass index. HDL indicates high-density lipoprotein cholesterol; Hp, Haptoglobin; LDL, low-density lipoprotein cholesterol.

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