LUND UNIVERSITY

Genetically Determined Height and Coronary Artery Disease

Nelson, C. P.; Hamby, S. E.; Saleheen, D.; Hopewell, J. C.; Zeng, L.; Assimes, T. L.; Kanoni, S.; Willenborg, C.; Burgess, S.; Amouyel, P.; Anand, S.; Blankenberg, S.; Boehm, B. O.; Clarke, R. J.; Collins, R.; Dedoussis, G.; Farrall, M.; Franks, Paul; Groop, Leif; Hall, A. S.; Hamsten, A.; Hengstenberg, C.; Hovingh, G. Kees; Ingelsson, E.; Kathiresan, S.; Kee, F.; Koenig, I. R.; Kooner, J.; Lehtimaeki, T.; Maerz, W.; McPherson, R.; Metspalu, A.; Nieminen, M. S.; O'Donnell, C. J.; Palmer, C. N. A.; Peters, A.; Perola, M.; Reilly, M. P.; Ripatti, S.; Roberts, R.; Salomaa, V.; Shah, S. H.; Schreiber, S.; Siegbahn, A.; Thorsteinsdottir, U.; Veronesi, G.; Wareham, N.; Willer, C. J.; Zalloua, P. A.; Erdmann, J.

Published in:
New England Journal of Medicine

DOI:
10.1056/NEJMoa1404881

2015

Link to publication

Citation for published version (APA):
Nelson, C. P., Hamby, S. E., Saleheen, D., Hopewell, J. C., Zeng, L., Assimes, T. L., Kanoni, S., Willenborg, C., Burgess, S., Amouyel, P., Anand, S., Blankenberg, S., Boehm, B. O., Clarke, R. J., Collins, R., Dedoussis, G., Farrall, M., Franks, P., Groop, L., ... Samani, N. J. (2015). Genetically Determined Height and Coronary Artery Disease. *New England Journal of Medicine*, *372*(17), 1608-1618. https://doi.org/10.1056/NEJMoa1404881

Total number of authors:
56

General rights
Unless other specific re-use rights are stated the following general rights apply:
Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.
• Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
• You may not further distribute the material or use it for any profit-making activity or commercial gain
• You may freely distribute the URL identifying the publication in the public portal

Read more about Creative commons licenses: https://creativecommons.org/licenses/

Take down policy
If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

LUND UNIVERSITY

PO Box 117
221 00 Lund
+46 46-222 00 00

Download date: 12. Oct. 2024

ORIGINAL ARTICLE

Genetically Determined Height and Coronary Artery Disease

C.P. Nelson, S.E. Hamby, D. Saleheen, J.C. Hopewell, L. Zeng, T.L. Assimes, S. Kanoni, C. Willenborg, S. Burgess, P. Amouyel, S. Anand, S. Blankenberg, B.O. Boehm, R.J. Clarke, R. Collins, G. Dedoussis, M. Farrall, P.W. Franks, L. Groop, A.S. Hall, A. Hamsten, C. Hengstenberg, G. Kees Hovingh, E. Ingelsson, S. Kathiresan, F. Kee, I.R. König, J. Kooner, T. Lehtimäki, W. März, R. McPherson, A. Metspalu, M.S. Nieminen, C.J. O'Donnell, C.N.A. Palmer, A. Peters, M. Perola, M.P. Reilly, S. Ripatti, R. Roberts, V. Salomaa, S.H. Shah, S. Schreiber, A. Siegbahn, U. Thorsteinsdottir, G. Veronesi, N. Wareham, C.J. Willer, P.A. Zalloua, J. Erdmann, P. Deloukas, H. Watkins, H. Schunkert, J. Danesh, J.R. Thompson, and N.J. Samani, for the CARDIoGRAM+C4D Consortium*

ABSTRACT

BACKGROUND

The nature and underlying mechanisms of an inverse association between adult height and the risk of coronary artery disease (CAD) are unclear.

METHODS

We used a genetic approach to investigate the association between height and CAD, using 180 height-associated genetic variants. We tested the association between a change in genetically determined height of 1 SD (6.5 cm) with the risk of CAD in 65,066 cases and 128,383 controls. Using individual-level genotype data from 18,249 persons, we also examined the risk of CAD associated with the presence of various numbers of height-associated alleles. To identify putative mechanisms, we analyzed whether genetically determined height was associated with known cardiovascular risk factors and performed a pathway analysis of the height-associated genes.

RESULTS

We observed a relative increase of 13.5% (95% confidence interval [CI], 5.4 to 22.1; P<0.001) in the risk of CAD per 1-SD decrease in genetically determined height. There was a graded relationship between the presence of an increased number of height-raising variants and a reduced risk of CAD (odds ratio for height quartile 4 versus quartile 1, 0.74; 95% CI, 0.68 to 0.84; P<0.001). Of the 12 risk factors that we studied, we observed significant associations only with levels of low-density lipoprotein cholesterol and triglycerides (accounting for approximately 30% of the association). We identified several overlapping pathways involving genes associated with both development and atherosclerosis.

CONCLUSIONS

There is a primary association between a genetically determined shorter height and an increased risk of CAD, a link that is partly explained by the association between shorter height and an adverse lipid profile. Shared biologic processes that determine achieved height and the development of atherosclerosis may explain some of the association. (Funded by the British Heart Foundation and others.)

The authors' full names, academic degrees, and affiliations are listed in the Appendix. Address reprint requests to Dr. Samani at the Department of Cardiovascular Sciences, University of Leicester, BHF Cardiovascular Research Centre, Glenfield Hospital, Groby Rd., Leicester, LE3 9QP, United Kingdom, or at njs@le.ac.uk.

*A complete list of members of the Coronary Artery Disease Genomewide Replication and Meta-Analysis plus the Coronary Artery Disease (CARDIoGRAM+C4D) Consortium is provided in the Supplementary Appendix, available at NEJM.org.

This article was published on April 8, 2015, at NEJM.org.

N Engl J Med 2015;372:1608-18.
DOI: 10.1056/NEJMoa1404881
Copyright © 2015 Massachusetts Medical Society.

The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.

There is a well-established association between a shorter adult height and an increased risk of coronary artery disease (CAD).[1] Shorter stature is also associated with risk factors for CAD, including high blood pressure, high levels of low-density lipoprotein (LDL) cholesterol, and diabetes.[2,3] An individual-level meta-analysis showed that a decrease of 1 SD (approximately 6.5 cm) in height was associated with a relative increase of 8% (95% confidence interval [CI], 6 to 10) in the risk of fatal or nonfatal CAD.[2] The effect was largely unchanged after adjustment for smoking status, systolic blood pressure, history of diabetes, body-mass index, lipid markers, alcohol consumption, education level, and occupation.[2] Therefore, the precise mechanisms linking shorter height with an increased risk of CAD remain unclear.

Genetic variants that affect a trait provide a means of exploring the relationship between the trait and the disease and to identify putative mechanisms. In a genomewide association study, Lango Allen et al.[4] identified a large number of independent genetic variants associated with adult height, which is a highly heritable trait. Large-scale genomewide association studies have also been undertaken to determine genetic variants associated with CAD[5-7] and several cardiovascular risk factors.[8-15] Here, we used the 180 single-nucleotide polymorphisms (SNPs) that explain about 10% of the variation in height, as identified by Lango Allen et al.,[4] and leveraged CAD-association data for the same variants for up to 193,449 persons to examine the association between genetically mediated variation in height and the risk of CAD. We also examined the association between the height-associated variants and several cardiovascular risk factors and performed bioinformatics analyses of the height-associated variants to identify other potential biologic mechanisms that could link a shorter height with an increased risk of CAD.

METHODS

HEIGHT-ASSOCIATED VARIANTS

To identify height-associated genetic variants, Lango Allen et al.[4] (in the Genetic Investigation of Anthropometric Traits [GIANT] Consortium) analyzed 183,727 persons of European descent and observed that variants at 180 loci showed an association with height at a genomewide significance level ($P<5\times10^{-8}$). We used the lead SNP from each locus (i.e., the SNP showing the strongest association) in the current analysis. None of these variants lie in loci implicated by genomewide association studies in susceptibility to CAD.[5-7]

ASSOCIATION BETWEEN HEIGHT-ASSOCIATED VARIANTS AND CAD

To examine the association between height-associated genetic variants and CAD, we extracted summary association statistics for these variants for the cohorts that contributed to the meta-analyses of genomewide association studies of CAD performed by the Coronary Artery Disease Genomewide Replication and Meta-Analysis (CARDIoGRAM) Consortium[5] and the Coronary Artery Disease (C4D) Consortium.[6] Of the 180 SNPs, 112 were also included on the Metabochip array, a customized array containing 200,000 SNP markers.[16] We also extracted data for these 112 SNPs from the Metabochip-array CAD meta-analysis performed by the combined CARDIoGRAM+C4D Consortium for cohorts that were not included in the previous CARDIoGRAM or C4D meta-analyses.[7] Each of the studies that were included in these meta-analyses adhered to a case–control design, including some nested within cohorts.[5-7]

The numbers of cases and controls that were contributed by each consortium are provided in Table S1 in the Supplementary Appendix, available with the full text of this article at NEJM.org. The number of samples and SNPs that were contributed by individual studies within each consortium are provided in Table S2 in the Supplementary Appendix. Details regarding the ascertainment of samples for each study are provided in the primary articles.[5-7] All cases were required to have had a validated history of myocardial infarction, coronary revascularization, or angiographic coronary disease.

HEIGHT-ASSOCIATED VARIANTS AND CARDIOVASCULAR RISK FACTORS

In parallel, to investigate potential explanatory effects of genetically determined height on the risk of CAD through known cardiovascular risk factors, we extracted estimates of effect size for each of the height variants from publicly available meta-analyses of data sets from genome-

The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.

wide association studies for systolic and diastolic blood pressures, mean and pulse pressures,[8,9] LDL cholesterol level, high-density lipoprotein (HDL) cholesterol level, triglyceride level,[10] presence or absence of type 2 diabetes mellitus,[11] body-mass index,[12,13] glucose and insulin levels,[14] and smoking quantity.[15] The maximum sizes of these data sets ranged from 29,182 to 249,796 samples (Table 1).

STATISTICAL ANALYSIS

For each height-associated variant, we calculated β_3 values (the putative association between height and CAD mediated through that variant) from the direct measurements of β_1 (the effect size of the association between the variant and height) and β_2 (the effect size of the association between the variant and CAD), as described previously.[8,17] (A more complete description of β_1, β_2, and β_3 and the relationships among them and how β_3 was calculated is provided in Fig. S1 in the Supplementary Appendix.) The value for β_3 can be interpreted as the odds ratio for CAD per 1-SD increase in genetically determined height. Because the association between each SNP with height and the association with CAD is very small, individual β_3 values are likely to center around 1.0. Combining the β_3 values from all SNPs provides additional power to assess the overall association between height and CAD (i.e., composite association). We used inverse-variance–weighted random-effects meta-analysis to combine individual β_3 estimates. We performed the same analysis in a subgroup of patients with a history of myocar-

Table 1. Association between Genetically Determined Height and Coronary Artery Disease and Cardiovascular Risk Factors.*

Risk Factor	Maximum No. of Samples in Data Set	Estimated Association (95% CI)†	P Value	I^2‡
Coronary artery disease	65,066 cases, 128,383 controls	0.88 (0.82 to 0.95)	<0.001	55.7
Body-mass index§	249,796	0.01 (–0.02 to 0.03)	0.74	14.7
Blood pressure				
Systolic	69,899	0.34 (–0.31 to 1.00)	0.30	41.6
Diastolic	69,909	0.14 (–0.27 to 0.56)	0.50	42.1
Mean arterial pressure¶	29,182	0.20 (–0.19 to 0.60)	0.32	41.6
Pulse pressure‖	74,079	0.23 (–0.06 to 0.52)	0.12	26.9
Cholesterol				
Low-density lipoprotein	95,454	–0.06 (–0.09 to –0.04)	<0.001	31.6
High-density lipoprotein	99,900	–0.02 (–0.05 to 0.02)	0.44	54.0
Triglycerides	96,598	–0.05 (–0.08 to –0.03)	<0.001	29.1
Type 2 diabetes	34,840 cases, 114,981 controls	0.95 (0.83 to 1.07)	0.38	50.0
Glucose	96,496	0.01 (–0.01 to 0.02)	0.48	31.0
Log-transformed plasma insulin	85,573	0.01 (–0.01 to 0.02)	0.29	37.3
Smoking quantity**	41,150	0.04 (–0.01 to 0.09)	0.11	15.8

* Estimates of effect size for each of the height variants were extracted from publicly available meta-analyses of data sets from genomewide association studies.
† The average effect estimates for a 1-SD increase in height are shown as odds ratios for categorical diseases (coronary artery disease and diabetes). For quantitative traits, the β estimates are shown in either absolute values (systolic and diastolic blood pressure, pulse pressure, mean arterial pressure, smoking quantity, glucose, and log insulin) or in SD (body-mass index, high-density lipoprotein and low-density lipoprotein cholesterol, and triglycerides).
‡ I^2 indicates the percentage of total variation in study estimates because of heterogeneity in the meta-analysis.
§ The body-mass index is the weight in kilograms divided by the square of the height in meters.
¶ Mean arterial pressure was defined as two thirds diastolic pressure plus one third systolic pressure.[9]
‖ Pulse pressure was defined as systolic pressure minus diastolic pressure.
** Scores for smoking-quantity levels among smokers (cigarettes smoked per day) were 0 (1 to 10 cigarettes per day), 1 (11 to 20 cigarettes), 2 (21 to 30 cigarettes), and 3 (31 or more cigarettes).[17]

The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.

dial infarction and in men and women separately, using sex-specific estimates of β_1 released by the GIANT Consortium (www.broadinstitute.org/collaboration/giant/index.php/GIANT_consortium_data_files).

For a subgroup of CAD cohorts in which we had access to individual-level genotypes genome-wide (Table S3 in the Supplementary Appendix), we performed a weighted analysis of genetic risk score to evaluate the effect of the presence of an increasing number of height-related variants on the risk of CAD. We calculated a value of 0 to 2 for every SNP for each individual on the basis of the sum of the posterior probabilities for the height-increasing allele and multiplied by the effect size observed for height. We then totaled these values across all SNPs for each individual, and the individuals were then grouped into quartiles. We used logistic regression to assess the quartiles, after adjustment for study, to estimate combined odds ratios for CAD.

To assess the association between height variants and cardiovascular risk factors, we combined the β_3 estimates using a fixed-effects meta-analysis, except in cases in which heterogeneity was high (I^2, >40%), in which case we performed a random-effects meta-analysis. For these analyses, the β_3 values reflect the change in measurement unit of the variable per 1-SD change in height for quantitative variables (with a negative value reflecting an inverse association) or an odds ratio for categorical variables. Because we tested a total of 13 traits (including CAD), we considered a P value of 0.003 to indicate statistical significance (Table 1).

To identify common biologic processes that might explain the association between height and CAD, we performed pathway analysis using Ingenuity Pathway Analysis (IPA) software, version 18488943 (Ingenuity Systems). Such an analysis requires the assignment of each height-associated SNP to a specific gene that is then included in the pathway analysis. (Further details regarding the selection process for the genes are provided in the Supplementary Appendix; the full list of genes that are included in the analysis is provided in Table S4 in the Supplementary Appendix.) The IPA output includes Benjamini–Hochberg Q values for the false discovery rate.[18]

RESULTS

STUDY CASES AND CONTROLS

The maximum number of CAD cases and controls available for analyses were 65,066 and 128,383, respectively (Table S1 in the Supplementary Appendix); 73.8% of the cases and 49.8% of the controls were men. The average age was 57.3 years (range, 42.4 to 75.6), and 65% of the cases reported a history of myocardial infarction.

HEIGHT-ASSOCIATED VARIANTS AND CAD

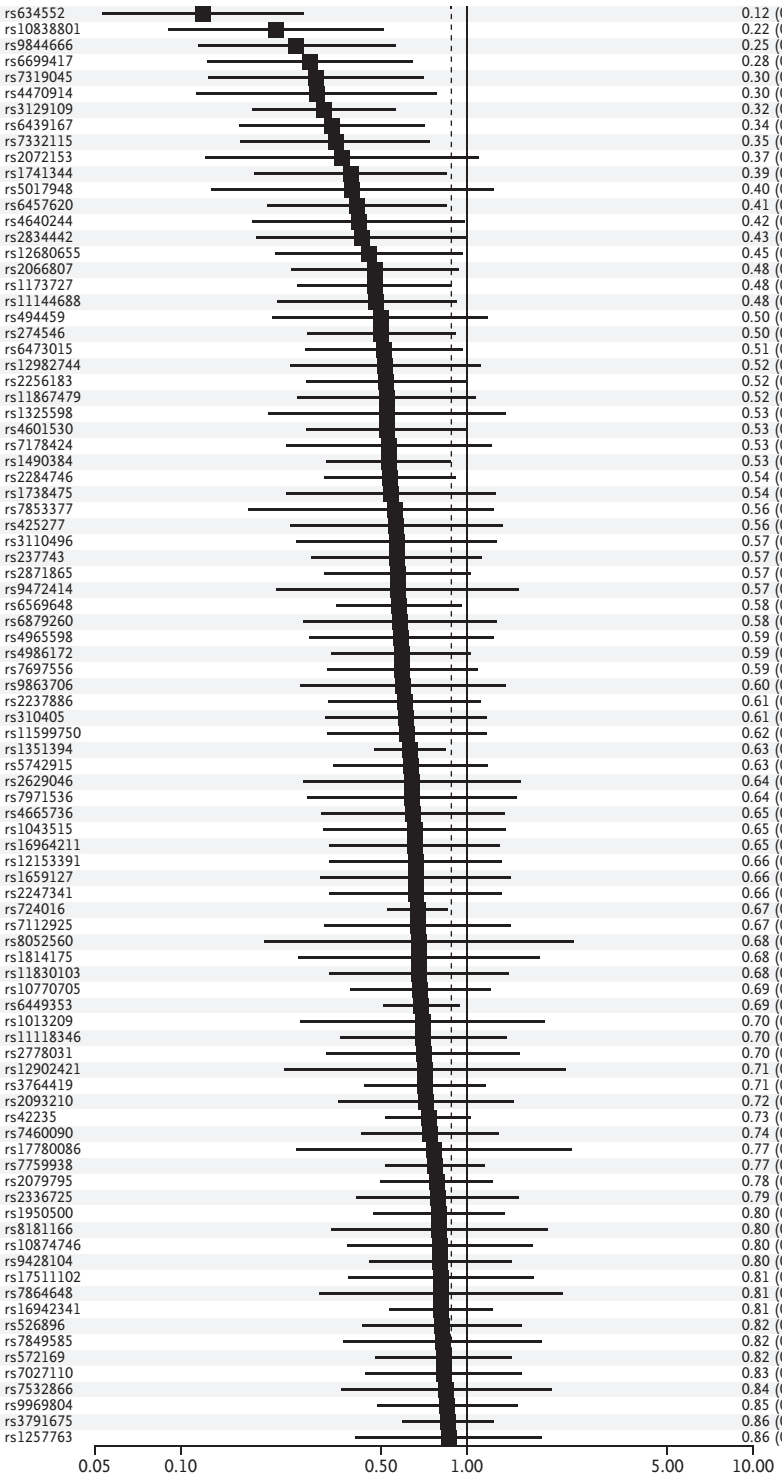
The individual β_3 odds ratios for the 180 SNPs that were analyzed to investigate the association between height and CAD are shown in Figure 1. In a random-effects meta-analysis, there was a significant association between the height-associated SNPs and CAD (odds ratio, 0.88; 95% CI, 0.82 to 0.95; P<0.001) (Table 1). This association translated to a relative increase of 13.5% (95% CI, 5.4 to 22.1) in the risk of CAD per 1-SD decrease in height.

As anticipated, most individual β_3 values centered around 1.0 and were nonsignificant (Fig. 1). However, some values had a nominally significant association (P<0.05) both above and below 1.0. Because 180 variants were tested, some of these associations could reflect chance (only 3 survived Bonferroni correction), but they could also represent pleiotropy — in other words, an effect of these loci on the risk of CAD that was independent of any effect through height. To rule out the possibility that the observed genetic association between height and CAD was being driven by more extreme associations, we repeated the meta-analysis with the exclusion of six SNPS that showed an individual association at P<0.001. The combined association between the remaining

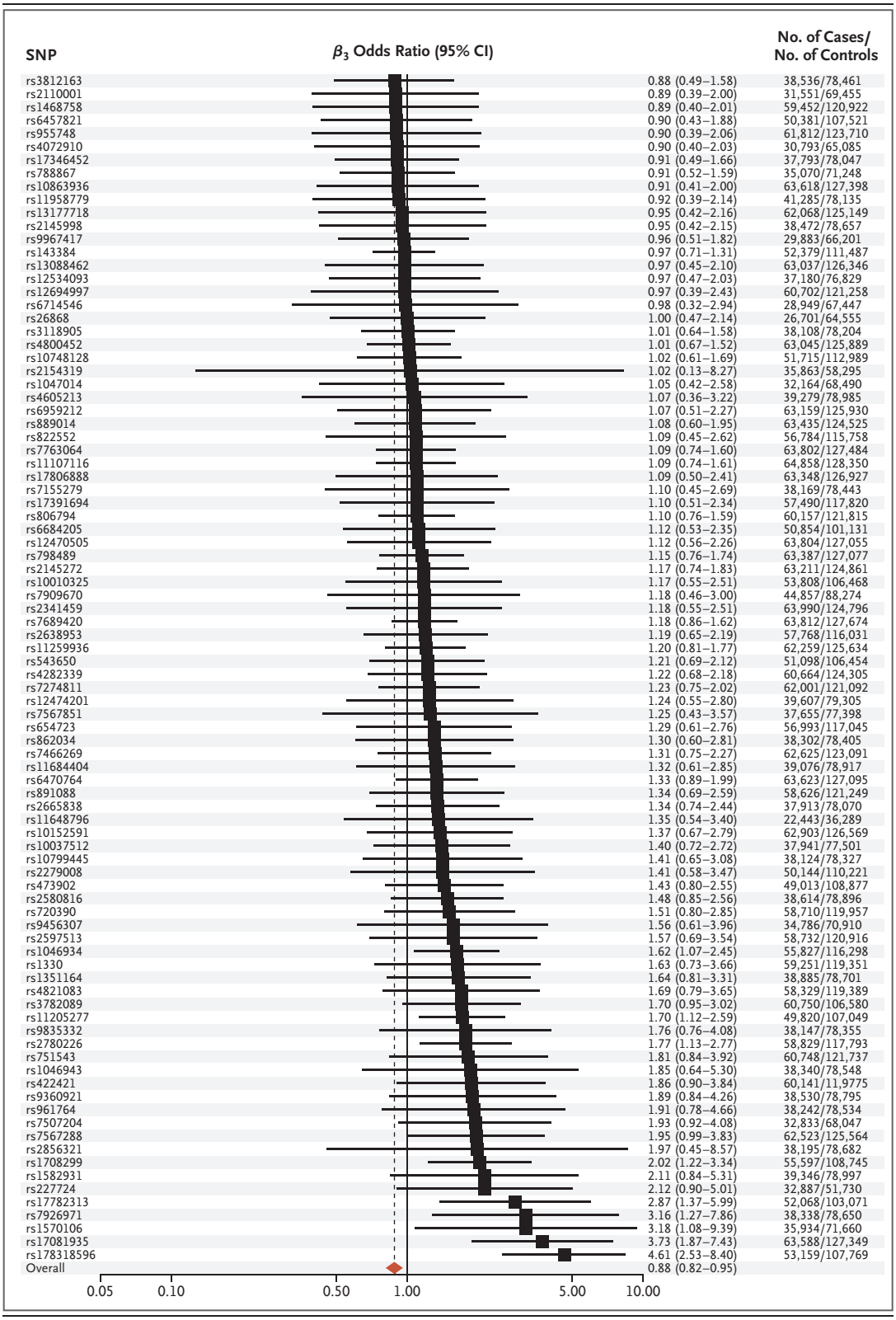
Figure 1 (next pages). Forest Plot Showing the Effect Size of Height on the Risk of Coronary Artery Disease (CAD) for Each Height-Associated Genetic Variant.

Shown are odds ratios for each height-associated single-nucleotide polymorphism (SNP) for β_3 values (i.e., the putative association between height and CAD mediated through that variant). The number of cases and controls that were analyzed for each variant are shown. The β_3 odds ratios are organized in ascending values across two panels for ease of visualization. The overall β_3 estimate (shown in red) is from a random-effects meta-analysis of all SNPs.

The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.



The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.



The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.

variants and CAD was largely unchanged (odds ratio, 0.88; 95% CI, 0.82 to 0.94; P<0.001).

The association between genetically determined height and CAD remained significant in the subgroup of cases with a history of myocardial infarction (odds ratio, 0.88; 95% CI, 0.80 to 0.96; P=0.003). In sex-specific analyses, the association between the variant and CAD was significant in men (odds ratio, 0.88; 95% CI, 0.81 to 0.95; P=0.001) but not in women (odds ratio, 0.96; 95% CI, 0.86 to 1.07; P=0.46). However, in an interaction test, the difference between the sexes was not significant (P=0.19).

GENETIC RISK SCORE AND RISK OF CAD

Individual-level data were available for 18,249 persons (including 8240 cases) from six cohorts (Table S3 in the Supplementary Appendix). The risk of CAD among individuals, as partitioned into quartiles carrying an increasing number of height-raising alleles, is shown in Figure 2. Those with an increased number of height-raising alleles had a reduced risk of CAD, with an odds ratio for quartile 2 vs. quartile 1 of 0.90 (95% CI, 0.83 to 0.98; P=0.02), an odds ratio for quartile 3 vs. quartile 1 of 0.88 (95% CI, 0.81 to 0.96; P=0.003), and an odds ratio for quartile 4 vs. quartile 1 of 0.74 (95% CI, 0.68 to 0.80; P<0.001).

HEIGHT-ASSOCIATED VARIANTS AND CARDIOVASCULAR RISK FACTORS

The findings from the analyses of the composite association between height-associated variants and specific cardiovascular risk factors are provided in Table 1. For most of the risk factors, the analyses did not identify any evidence of an association between genetically determined height and the risk of CAD. The two exceptions were LDL cholesterol and triglyceride levels, for which there were small but significant associations.

For both LDL cholesterol and triglycerides, the associations were in a direction that could have contributed to the observed association between a shorter genetically determined height and an increased risk of CAD. To investigate this finding further, we evaluated the quantitative associations between LDL cholesterol and triglycerides and the risk of CAD that were reported in observational studies,[19] taking into account regression dilution.[20] We determined that for each 1-SD increase, the risk of CAD was raised on average by 45% (log odds ratio, 0.37) for the LDL cholesterol level and by 32% (log odds ratio, 0.28) for the triglyceride level. Then, from the respective changes from a 1-SD change in genetically determined height, we estimated that the risk of CAD would increase by 2.3% (95% CI, 1.9 to 2.8) because of the increase in the LDL cholesterol level and by 1.5% (95% CI, 1.2 to 1.8) because of the increase in the triglyceride level. This suggests that approximately 19% of the observed association between a genetically determined decrease in height and an increased risk of CAD could be explained by the effect of shorter height on LDL cholesterol and approximately 12% by an effect on triglycerides. To confirm that the majority of the genetic association of height with CAD was not mediated by lipid levels, we repeated our analysis of the association between height variants and the risk of

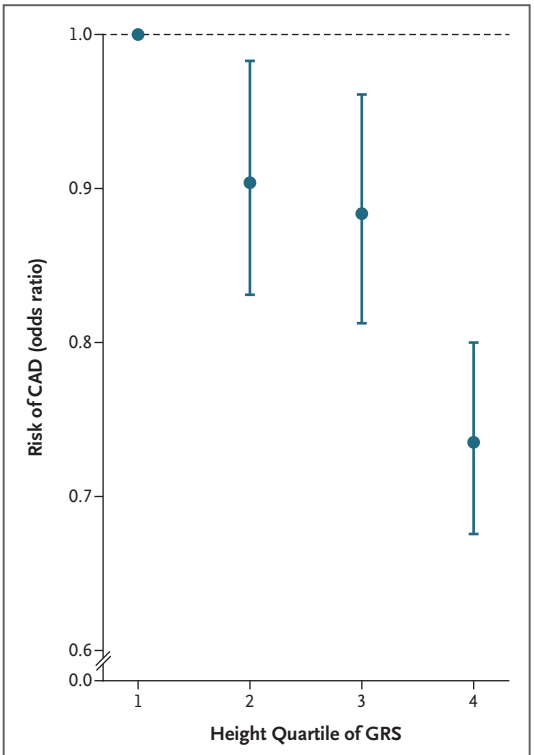


Figure 2. Analysis of the Association between the Presence of an Increasing Number of Height-Related Alleles and the Risk of CAD, According to Quartile of Genetic Risk Score (GRS).

The analysis was performed in 18,249 samples (including 8240 obtained from patients with CAD) with the use of individual-level genotype data. Shown are odds ratios and 95% confidence intervals. Participants were divided into quartiles on the basis of the number of height-increasing alleles that were present, with quartile 1 (reference quartile) carrying the fewest.

The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.

Table 2. Biologic Pathways Identified by Means of IPA of Height-Associated Variants.*

Canonical Pathways in IPA	Q Value†	Ratio‡	Proteins in Pathway
Factors promoting cardiogenesis in vertebrates	0.003	0.07	NKX2–5, BMP2, TGFB2, MEF2C, BMP6, PRKCZ, NOG
Growth hormone signaling	0.03	0.06	SOCS2, IGF1R, GH1, SOCS5, PRKCZ
Axonal guidance signaling	0.03	0.06	FGFR4, SOCS2, IGF1R, INSR, SOCS5
STAT3 pathway	0.03	0.03	SLIT3, PAPPA2, PAPPA, RHOD, ADAM28, GNA12, BMP2, PTCH1, HHIP, NFATC4, BMP6, PRKCZ
BMP signaling pathway	0.03	0.06	NKX2-5, RUNX2, BMP2, BMP6, NOG
TGF-β signaling	0.04	0.05	NKX2-5, AMH, RUNX2, BMP2, TGFB2
IGF-1 signaling	0.049	0.05	SOCS2, IGF1R, IGFBP7, SOCS5, PRKCZ

* BMP denotes bone morphogenetic protein, IGF-1 insulin-like growth factor 1, IPA Ingenuity Pathway Analysis, STAT3 signal transducer and activator of transcription 3, and TGF-β transforming growth factor β.
† The Q value was calculated with the use of the Benjamini–Hochberg method for determining the false discovery rate.
‡ The ratio is the proportion of the genes in the IPA that were part of the input list for the height-related genes.

CAD with the exclusion of 60 SNPs that were associated with a lipid trait at P<0.05. An analysis of the remaining SNPs resulted in an odds ratio of 0.89 (95% CI, 0.81 to 0.98; P=0.01).

PATHWAY ANALYSIS

Biologic pathways (incorporating genes at the height loci) with a Q value of less than 0.05 for the false discovery rate, as identified on the IPA analysis, are provided in Table 2. Also shown are the genes within each pathway that were present on the input list and also the proportion of genes in each pathway formed by them. Pathways in IPA software have a hierarchical organization, and many of the pathways that are identified are overlapping and, in some cases, are subsets of each other. For example, the pathway that is identified as "factors affecting cardiogenesis" is an amalgam of other pathways and overlaps with signaling pathways for bone morphogenetic protein (BMP) and transforming growth factor β (TGF-β), and all three of these pathways share genes with other pathways. Likewise, there is overlap between the signaling pathways for growth hormone and insulin-like growth factor 1 (IGF-1).

DISCUSSION

In this study, we found an association between a genetically determined decrease in height and an increased risk of CAD. Our finding validates the epidemiologic observation of an inverse association between height and CAD.[1,2]

A key advantage of using a genetic approach over a traditional epidemiologic approach to investigate an association such as that between height and CAD is that genotypes (because they are randomly distributed at birth) are unlikely to be confounded by lifestyle or environmental factors. Regardless of whether such factors are known (e.g., poor nutrition or socioeconomic conditions during childhood) or unknown, they can independently affect achieved height and the risk of CAD and lead to a spurious association between them (Fig. 3). It is nonetheless possible that the genetic variants themselves affect height and CAD risk through entirely different mechanisms. However, given the large number of variants that we included in the analysis, all of which were selected only because of their association with height, it is likely that at least some of the processes are shared. This hypothesis is supported by the finding from the individual-level analysis of genetic risk score showing a direct correlation between the presence of an increasing number of height-related alleles and a reduction in the risk of CAD (Fig. 2).

A genetic approach also offers novel methods to explore potential mechanisms linking shorter height with an increased risk of CAD (Fig. 3). In this context, we performed two analyses. First, we applied the same genetic approach to investigate the association between height-related genetic variants and several established and potential cardiovascular risk factors. Notable negative findings here include the lack of an overall effect of height-associated SNPs on body-mass index. This suggests that the association between shorter stature and an increased risk of CAD is not mediated by an effect on obesity. On

The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.

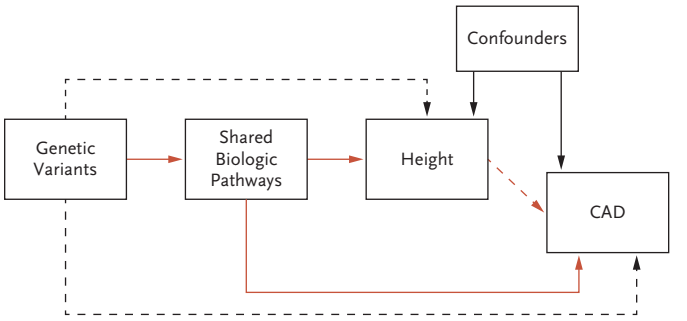


Figure 3. Interpreting the Association between Genetically Determined Shorter Height and Increased Risk of CAD.

The main advantage of the genetic approach is that it reduces the likelihood of known and unknown demographic, lifestyle, socioeconomic, or behavioral confounders that have an independent effect on height and the risk of CAD (solid black lines) and could give rise to a false association between the two factors. It is possible that the association between the studied genetic variants and height and the association with CAD are through completely different mechanisms (dashed black lines). However, the more likely scenario on the basis of our findings is that height variants affect biologic pathways, which on the one hand determine achieved height and on the other hand influence the risk of CAD (solid red lines). It is also possible that genetically determined height itself alters lifestyle or behavior, which then affects the risk of CAD (dashed red line).

the other hand, there was a significant overall association between height SNPs and LDL cholesterol and triglycerides in a direction consistent with their association with CAD. The association between shorter stature and increased plasma LDL cholesterol and triglyceride levels has also been observed in epidemiologic studies.[2] The mechanisms by which height-associated SNPs have an effect on LDL cholesterol and triglyceride levels are not clear. In any case, these effects in combination potentially explain less than one third of the observed association between genetically determined shorter height and an increased risk of CAD.

Second, we performed pathway analysis, which identified a number of overlapping pathways linking height-associated SNPs that could also have an effect on the risk of CAD, including the BMP- and TGF-β–signaling pathways, axon-guidance pathways, and the STAT3 and IGF-I pathways, all of which have experimentally documented roles in the development of atherosclerosis.[21-26] The limitations of pathway analysis included the need to assign a specific gene for each height-associated locus and incomplete knowledge regarding how such pathways are constructed. (A fuller discussion of the pathways is provided in the Supplementary Appendix.) Taken together, these findings suggest that several overlapping and complex biologic pathways on the one hand influence development and height and on the other hand influence the risk of atherosclerosis through an effect on vascular biology and function (Fig. 3).

In contrast to epidemiologic studies in which a similar inverse association between height and CAD was present in both men and women,[2] we did not see a significant association in women. Whether this represents a genuine difference in the effect of genetically determined height on the risk of CAD between men and women or simply reflects the reduced power from the much smaller sample size available for analysis in women is unclear. Notably, the effect sizes that were observed in men and women were not significantly different in an interaction analysis.

Height and other measurements of body size have a positive correlation with the diameter of coronary arteries.[27] Therefore, a potential simple explanation for an increased risk of CAD in shorter persons is that they have proportionally smaller-caliber coronary arteries, so a similar plaque burden could result in greater probability of symptomatic disease. However, women also have smaller-caliber arteries than men, independent of body size and height.[27] Reduced height and female sex would therefore be expected to have an additive effect if this was the mechanism linking shorter height with an increased risk of CAD. In this context, the finding of a weaker association between genetically determined shorter height and CAD in women than in men would argue against a structural explanation on the basis of coronary-vessel caliber as the main explanation for the inverse association between height and CAD.

Although the genetic approach that we used allows us to reduce the possibility of confounding of any observed association by socioeconomic, lifestyle, or environmental factors, it does not rule out the possibility that the association between genetically determined shorter height with an increased risk of CAD is due to lifestyle choices or behavior adopted by such persons as a direct consequence of being shorter (Fig. 3). Indeed, in an exemplar exploration of this possibility, we examined whether the height-related variants showed an association with the quantity

The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.

of cigarettes smoked among smokers but found no evidence for this hypothesis (Table 1). Other relevant behavioral changes that could have an effect on the risk of CAD that could be adopted by persons of short stature include those related to diet, physical activity, and alcohol consumption.

In conclusion, using a genetic approach, we found an association between genetically determined shorter height and an increased risk of CAD. Part of this inverse association may be driven by the association between shorter height and an adverse lipid profile, although the majority of the relationship is likely to be determined by shared biologic processes that determine achieved height and atherosclerosis development. More generally, our findings underscore the complexity underlying the inherited component of CAD.

Supported by the British Heart Foundation, the United Kingdom National Institute for Health Research, the European Union project CVgenes@target, and a grant from the Leducq Foundation.

Disclosure forms provided by the authors are available with the full text of this article at NEJM.org.

We thank the members of the cited consortiums of genome-wide association studies for making their data available.

APPENDIX

The authors are as follows: Christopher P. Nelson, Ph.D., Stephen E. Hamby, Ph.D., Danish Saleheen, M.B., B.S., Ph.D., Jemma C. Hopewell, Ph.D., Lingyao Zeng, M.Sc., Themistocles L. Assimes, M.D., Ph.D., Stavroula Kanoni, Ph.D., Christina Willenborg, Ph.D., Stephen Burgess, Ph.D., Phillipe Amouyel, M.D., Ph.D., Sonia Anand, F.R.C.P.C., Stefan Blankenberg, M.D., Bernhard O. Boehm, M.D., Robert J. Clarke, F.R.C.P., Rory Collins, F.R.C.P.E., George Dedoussis, Ph.D., Martin Farrall, F.R.C.Path., Paul W. Franks, Ph.D., Leif Groop, M.D., Ph.D., Alistair S. Hall, F.R.C.P., Anders Hamsten, M.D., Ph.D., Christian Hengstenberg, M.D., G. Kees Hovingh, M.D., Ph.D., Erik Ingelsson, M.D., Ph.D., Sekar Kathiresan, M.D., Frank Kee, M.D., Inke R. König, Ph.D., Jaspal Kooner, F.R.C.P., Terho Lehtimäki, M.D., Ph.D., Winifred März, Ph.D., Ruth McPherson, F.R.C.P.C., Andres Metspalu, M.D., Ph.D., Markku S. Nieminen, M.D., Christopher J. O'Donnell, M.D., Colin N.A. Palmer, Ph.D., Annette Peters, M.D., Markus Perola, M.D., Ph.D., Muredach P. Reilly, M.B., B.Ch., Samuli Ripatti, Ph.D., Robert Roberts, F.R.C.P.C., Veikko Salomaa, M.D., Ph.D., Svati H. Shah, M.D., Stefan Schreiber, M.D., Agneta Siegbahn, Ph.D., Unnur Thorsteinsdottir, Ph.D., Giovani Veronesi, Ph.D., Nicholas Wareham, M.B., B.S., Ph.D., Cristen J. Willer, Ph.D., Pierre A. Zalloua, Ph.D., Jeanette Erdmann, Ph.D., Panos Deloukas, Ph.D., Hugh Watkins, F.R.C.P., Heribert Schunkert, M.D., John Danesh, F.R.C.P., John R. Thompson, Ph.D., and Nilesh J. Samani, F.R.C.P.

The authors' affiliations are as follows: the Departments of Cardiovascular Sciences (C.P.N., S.E.H., N.J.S.) and Health Sciences (J.R.T.), University of Leicester, and the National Institute for Health Research Leicester Cardiovascular Biomedical Research Unit (C.P.N., S.E.H., N.J.S.), Leicester, the Department of Public Health and Primary Care, University of Cambridge (D.S., S. Burgess, J.D.), Wellcome Trust Sanger Institute (S.R., P.D.), and the Medical Research Council Epidemiology Unit, Institute of Metabolic Science, Addenbrooke's Hospital (N.W.), Cambridge, the Clinical Trial Service Unit and Epidemiological Studies Unit (J.C.H., R.J.C., R.C.), Division of Cardiovascular Medicine, Radcliffe Department of Medicine (M.F., H.W.), and Wellcome Trust Centre for Human Genetics (M.F., H.W.), University of Oxford, Oxford, William Harvey Research Institute, Barts and the London School of Medicine and Dentistry, Queen Mary University of London (S.K., P.D.), Imperial College London (B.O.B.), and the National Heart and Lung Institute, Imperial College (J.K.), London, Leeds Institute of Genetics, Health and Therapeutics, University of Leeds, Leeds (A.S.H.), U.K. Clinical Research Collaboration Centre of Excellence for Public Health (Northern Ireland), Queen's University of Belfast, Belfast (F.K.), and the Medical Research Institute, University of Dundee, Ninewells Hospital and Medical School, Dundee (C.N.A.P.) — all in the United Kingdom; the Center for Non-Communicable Diseases, Karachi, Pakistan (D.S.); the Department of Medicine (D.S.) and Cardiovascular Institute, Perelman School of Medicine (M.P.R.), University of Pennsylvania, Philadelphia; Deutsches Herzzentrum München, Technische Universität München and Deutsches Zentrum für Herz-Kreislauf-Forschung, partner site Munich Heart Alliance (L.Z., R.J.C., R.C., C.H., H.S.), and German Center for Cardiovascular Research partner site Munich (A.P.), Munich, Institut für Integrative und Experimentelle Genomik, Universität zu Lübeck and German Center for Cardiovascular Research, partner site Hamburg–Lübeck–Kiel (C.W., J.E.), and Institut für Medizinische Biometrie und Statistik, Universität zu Lübeck (I.R.K.), Lübeck, Clinic for General and Interventional Cardiology, University Heart Center Hamburg, Hamburg (S. Blankenberg), Ulm University Medical Center, Department of Internal Medicine I, Ulm (B.O.B.), Clinical Institute of Medical and Chemical Laboratory Diagnostics, Medical University of Graz, Graz, Austria (W.M.); Synlab Academy, Synlab Services, Medical Clinic V, Mannheim Medical Faculty, Heidelberg University, Mannheim (W.M.), Institute of Epidemiology II, Helmholtz Zentrum München–German Research Center for Environmental Health, Neuherberg (A.P.), and Institut für Klinische Molekularbiologie, Christian-Albrechts Universität, Kiel (S.S.) — all in Germany; the Department of Medicine, Stanford University School of Medicine, Stanford, CA (T.L.A.); Institut Pasteur de Lille, INSERM Unité 744, Université Lille Nord de France, Lille, France (P.A.); the Population Health Research Institute, McMaster University, Hamilton, ON (S.A.), and the John and Jennifer Ruddy Canadian Cardiovascular Genetics Centre and Atherogenomics Laboratory, University of Ottawa Heart Institute, Ottawa (R.M., R.R.) — both in Canada; Lee Kong Chian School of Medicine, Nanyang Technological University, Singapore, Singapore (B.O.B.); the Department of Dietetics–Nutrition, Harokopio University, Athens (G.D.); the Genetic and Molecular Epidemiology Unit, Department of Clinical Sciences, Lund University Diabetes Center, Skåne University Hospital (P.W.F.), and the Department of Clinical Sciences, Diabetes and Endocrinology, Lund University, University Hospital Malmö (L.G.), Malmö, the Department of Public Health and Clinical Medicine, Umeå University, Umeå (P.W.F.), the Atherosclerosis Research Unit, Department of Medicine, Karolinska Institutet, Stockholm (A.H.), and the Department of Medical Sciences, Molecular Epidemiology and Science for Life Laboratory (E.I.), and Uppsala Clinical Research Center (A.S.), Uppsala University, Uppsala — all in Sweden; the Department of Nutrition, Harvard School of Public Health (P.W.F.), and the Cardiology Division, Center for Human Genetic Research, and Cardiovascular Research Center, Massachusetts General Hospital and Harvard Medical School (S.K.), Boston, Broad Institute of Harvard and MIT, Cambridge (S.K.), and the National Heart, Lung, and Blood Institute Framingham Heart Study, Framingham (C.J.O.) — all in Massachusetts; the Department of Vascular Medicine, Academic Medical Center, Amsterdam (G.K.H.); the Department of Clinical Chemistry, Fimlab Laboratories and School of Medicine, University of Tampere, Tampere (T.L.), and the Division of Cardiology, Department of Medicine, Helsinki University Central Hospital (M.S.N.), Department of Chronic Disease Prevention, National Institute for Health and Welfare (M.P., V.S.), and Hjelt Institute

The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.

and Institute for Molecular Medicine Finland, University of Helsinki (S.R.), Helsinki — all in Finland; the Estonian Genome Center, University of Tartu, Tartu, Estonia (A.M., M.P.); Duke Molecular Physiology Institute, Division of Cardiology, Department of Medicine, Duke University, Durham NC (S.H.S.); deCODE Genetics, Reykjavik, Iceland (U.T.); the Research Center for Epidemiology and Preventive Medicine, Department of Clinical and Experimental Medicine, University of Insubria, Varese, Italy (G.V.); the Departments of Internal Medicine, Human Genetics and Computational Medicine and Bioinformatics, University of Michigan, Ann Arbor (C.J.W.); Lebanese American University, Chouran, Beirut, Lebanon (P.A.Z.); and King Abdulaziz University, Jeddah, Saudi Arabia (P.D.).

REFERENCES

1. Paajanen TA, Oksala NKJ, Kuukasjärvi P, Karhunen PJ. Short stature is associated with coronary heart disease: a systematic review of the literature and a meta-analysis. Eur Heart J 2010;31:1802-9.
2. Emerging Risk Factors Collaboration. Adult height and the risk of cause-specific death and vascular morbidity in 1 million people: individual participant meta-analysis. Int J Epidemiol 2012;41:1419-33.
3. Gunnell D, Whitley E, Upton MN, McConnachie A, Smith GD, Watt GC. Associations of height, leg length, and lung function with cardiovascular risk factors in the Midspan Family Study. J Epidemiol Community Health 2003;57:141-6.
4. Lango Allen H, Estrada K, Lettre G, et al. Hundreds of variants clustered in genomic loci and biological pathways affect human height. Nature 2010;467:832-8.
5. Schunkert H, König IR, Kathiresan S, et al. Large-scale association analysis identifies 13 new susceptibility loci for coronary artery disease. Nat Genet 2011;43:333-8.
6. Coronary Artery Disease (C4D) Genetics Consortium. A genome-wide association study in Europeans and South Asians identifies five new loci for coronary artery disease. Nat Genet 2011;43:339-44.
7. CARDIoGRAMplusC4D Consortium. Large scale association analysis identifies new risk loci for coronary artery disease. Nat Genet 2013;45:25-33.
8. Ehret GB, Munroe PB, Rice KM, et al. Genetic variants in novel pathways influence blood pressure and cardiovascular disease risk. Nature 2011;478:103-9.
9. Wain LV, Verwoert GC, O'Reilly PF, et al. Genome-wide association study identifies six new loci influencing pulse pressure and mean arterial pressure. Nat Genet 2011;43:1005-11.
10. Teslovich TM, Musunuru K, Smith AV, et al. Biological, clinical and population relevance of 95 loci for blood lipids. Nature 2010;466:707-13.
11. Morris AP, Voight BP, Teslovich TM et al. Large-scale association analysis provides insights into the genetic architecture and pathophysiology of type 2 diabetes. Nat Genet 2012;44:981-90.
12. Speliotes EK, Willer CJ, Berndt SI, et al. Association analyses of 249,796 individuals reveal 18 new loci associated with body mass index. Nat Genet 2010;42:937-48.
13. Yang J, Loos RJ, Powell JE, et al. FTO genotype is associated with phenotypic variability of body mass index. Nature 2012;490:267-72.
14. Manning AK, Hivert MF, Scott RA, et al. A genome-wide approach accounting for body mass index identifies genetic variants influencing fasting glycemic traits and insulin resistance. Nat Genet 2012;44:659-69.
15. Liu JZ, Tozzi F, Waterworth DM, et al. Meta-analysis and imputation refines the association of 15q25 with smoking quantity. Nat Genet 2010;42:436-40.
16. Voight BF, Kang HM, Ding J, et al. The metabochip, a custom genotyping array for genetic studies of metabolic, cardiovascular, and anthropometric traits. PLoS Genet 2012;8(8):e1002793.
17. Lieb W, Jansen H, Loley C, et al. Genetic predisposition to higher blood pressure increases coronary artery disease risk. Hypertension 2013;61:995-1001.
18. Benjamini Y, Hochberg Y. Controlling the false discovery rate: a practical and powerful approach to multiple testing. J R Stat Soc B 1995;57:289-300.
19. Emerging Risk Factors Collaboration. Lipid-related markers and cardiovascular disease prediction. JAMA 2012;307:2499-506.
20. Emerging Risk Factors Collaboration. Major lipids, apolipoproteins, and risk of vascular disease. JAMA 2009;302:1993-2000.
21. Toma I, McCaffrey TA. Transforming growth factor-β and atherosclerosis: interwoven atherogenic and atheroprotective aspects. Cell Tissue Res 2012;347:155-75.
22. van Gils JM, Derby MC, Fernandes LR, et al. The neuroimmune guidance cue netrin-1 promotes atherosclerosis by inhibiting the emigration of macrophages from plaques. Nat Immunol 2012;13:136-43.
23. Wanschel A, Seibert T, Hewing B, et al. Neuroimmune guidance cue Semaphorin 3E is expressed in atherosclerotic plaques and regulates macrophage retention. Arterioscler Thromb Vasc Biol 2013;33:886-93.
24. Gharavi NM, Alva JA, Mouillesseaux KP, et al. Role of the Jak/STAT pathway in the regulation of interleukin-8 transcription by oxidized phospholipids *in vitro* and in atherosclerosis *in vivo*. J Biol Chem 2007;282:31460-8.
25. Higashi Y, Sukhanov S, Anwar A, Shai SY, Delafontaine P. IGF-1, oxidative stress and atheroprotection. Trends Endocrinol Metab 2010;21:245-54.
26. von der Thüsen JH, Borensztajn KS, Moimas S, et al. IGF-1 has plaque-stabilizing effects in atherosclerosis by altering vascular smooth muscle cell phenotype. Am J Pathol 2011;178:924-34.
27. O'Connor NJ, Morton JR, Birkmeyer JD, Olmstead EM, O'Connor GT. Effect of coronary artery diameter in patients undergoing coronary bypass surgery. Circulation 1996;93:652-5.

Copyright © 2015 Massachusetts Medical Society.

RECEIVE IMMEDIATE NOTIFICATION WHEN AN ARTICLE IS PUBLISHED ONLINE FIRST

To be notified by e-mail when *Journal* articles are published Online First, sign up at NEJM.org.

The New England Journal of Medicine
Downloaded from nejm.org by Matthias Bank on December 11, 2015. For personal use only. No other uses without permission.
Copyright © 2015 Massachusetts Medical Society. All rights reserved.