**MENDELIAM RANDOMIZATION**

**Relationship Between Blood Pressure and Incident Cardiovascular Disease**

Linear and Nonlinear Mendelian Randomization Analyses

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**ABSTRACT:** Observational studies exploring whether there is a nonlinear effect of blood pressure on cardiovascular disease (CVD) risk are hindered by confounding. This limitation can be overcome by leveraging randomly allocated genetic variants in nonlinear Mendelian randomization analyses. Based on their association with blood pressure traits in a genome-wide association study of 299,024 European ancestry individuals, we selected 253 genetic variants to proxy the effect of modifying systolic and diastolic blood pressure. Considering the outcomes of incident coronary artery disease, stroke and the combined outcome of CVD, linear and nonlinear Mendelian randomization analyses were performed on 255,714 European ancestry participants without a history of CVD or antihypertensive medication use. There was no evidence favoring nonlinear relationships of genetically proxied systolic and diastolic blood pressure with the cardiovascular outcomes over linear relationships. For every 10-mm Hg increase in genetically proxied systolic blood pressure, risk of incident CVD increased by 49% (hazard ratio, 1.49 [95% CI, 1.38–1.61]), with similar estimates obtained for coronary artery disease (hazard ratio, 1.50 [95% CI, 1.38–1.63]) and stroke (hazard ratio, 1.44 [95% CI, 1.22–1.70]). Genetically proxied blood pressure had a similar relationship with CVD in men and women. These findings provide evidence to support that even for individuals who do not have elevated blood pressure, public health interventions achieving persistent blood pressure reduction will be of considerable benefit in the primary prevention of CVD. *(Hypertension. 2021;77:2004–2013. DOI: 10.1161/HYPERTENSIONAHA.120.16534.)*

**Key Words:** blood pressure ◼ coronary artery disease ◼ hypertension ◼ primary prevention ◼ stroke

More than 1 billion people worldwide experience hypertension, which is estimated to account for >20% of cardiovascular disease (CVD). Meta-analyses of randomized controlled trials have shown that a 10-mm Hg reduction in systolic blood pressure (SBP) is associated with a 15% to 20% reduction in the risk of coronary artery disease (CAD) and a 25% to 30% reduction in the risk of stroke. As such, blood pressure lowering is one of the most effective strategies for reducing the burden of CVD.

Large observational studies have previously explored the relationship between blood pressure and cardiovascular risk, potentially identifying linear associations in individuals free of CVD at baseline but J-shaped associations both in the general population and in patients with a history of CAD and stroke. However, it is difficult to make causal conclusions about the effects of altering blood pressure from such data because any identified associations may be susceptible to confounding from unknown or unmeasured factors. For the patients with elevated cardiovascular risk recruited to the SPRINT (Systolic Blood Pressure Intervention Trial), SBP lowering to <120 mm Hg as compared with 140 mm Hg resulted...
in fewer major cardiovascular events. However, no high-quality clinical trials have investigated the effect of blood pressure lowering below this level. Excessive blood pressure reduction in patients with atherosclerotic disease can reduce organ perfusion and increase CVD risk. Insight into the shape of the relationship between blood pressure and CVD risk is, therefore, critical for informing optimal prevention strategies.

In the Mendelian randomization (MR) paradigm, genetic variants can be used as proxies for studying the effect of varying blood pressure. In the same way as treatment allocation in a randomized controlled trial setting, random allocation of genetic variants means that they are unlikely to be affected by confounding from environmental factors. Recent methodological developments have allowed for MR investigation into the shape of the relationship between risk factors and outcomes. In this study, we use MR to investigate the shape of the relationship between genetically proxied blood pressure and incident CVD in a general population without a history of CVD or antihypertensive medication use. Our analyses aim to provide novel insight that can be used to inform public health strategies toward the primary prevention of CVD.
incident CAD and incident stroke), the first event recorded was used. Related individuals (kinship coefficient \(>0.0884\)) and those with prevalent CVD (identified through hospitalization codes and self-report) were excluded from the analyses. Individuals taking antihypertensive medications at baseline (UK Biobank field 200003) were also excluded from the analyses because their observed blood pressure is not reflective of their genetically predicted blood pressure, thus introducing bias into the nonlinear MR estimates.\(^{15,17}\)

### Candidate Instrumental Variables

For our primary analysis, we selected 253 uncorrelated \(\left( \text{r}^2<0.1 \right) \) single-nucleotide polymorphisms as candidate instrumental variables for SBP and DBP based on their previously published associations with blood pressure traits.\(^{25}\) Their associations with SBP and DBP were estimated in a genome-wide association study of 299,024 European ancestry individuals performed by the International Consortium of Blood Pressure study, which did not include UK Biobank participants.\(^{20}\) Using the coefficients for association with SBP and DBP (Table S2), a weighted allele score for each participant was created by multiplying the blood pressure–increasing allele dosage with the variant’s association with SBP or DBP, respectively, and summing across the 253 variants. The above genetic association estimates were taken from a study that adjusted for body mass index. As this study of 299,024 European ancestry individuals performed by the International Consortium of Blood Pressure study, which did not include UK Biobank participants,\(^{20}\) we further performed a sensitivity analysis that selected variants from a genome-wide association study meta-analysis of 2 non-UK Biobank cohorts that did not adjust for body mass index \((n=122,361; \text{Methods in the Data Supplement}).\) Fixed-effects meta-analysis was performed using METAL,\(^{22}\) and variants reaching genome-wide significance \((P<5\times10^{-8})\) were clumped to correlation codes and self-report) were excluded from the analyses. Related individuals (kinship coefficient, \(>0.0884\)) and those with prevalent CVD (identified through hospitalization codes and self-report) were excluded from the analyses. Individuals taking antihypertensive medications at baseline (UK Biobank field 200003) were also excluded from the analyses because their observed blood pressure is not reflective of their genetically predicted blood pressure, thus introducing bias into the nonlinear MR estimates.\(^{15,17}\)

### Statistical Analyses

All statistical analyses were performed using R (version 3.6.2). Differences in characteristics between UK Biobank population subgroups were assessed using a Student t test, Wilcoxon rank-sum test, Fisher exact test, or \(\chi^2\) test as appropriate. We performed MR analyses investigating the association between genetically proxied blood pressure (either SBP or DBP) and incident CVD, CAD, and stroke risk. Analyses were performed by modeling a linear relationship between genetically proxied blood pressure and the outcomes (linear MR)\(^{14,24}\) and also using the fractional polynomial method to test for a nonlinear relationship between genetically proxied blood pressure and the outcomes (nonlinear MR).\(^{15,17}\)

### Linear MR

We used the ratio of coefficients method to perform MR analyses that assumed a linear association of genetically proxied blood pressure with the risk of incident CVD, CAD, and stroke.\(^{25}\) This represents the association of the allele score with the cardiovascular outcome (incident CVD, CAD, or stroke) divided by the association of the allele score with the blood pressure trait (either SBP or DBP).\(^{20}\) Linear regression was used to estimate the association of the allele score with blood pressure, incorporating age, sex, principal components 1 to 10 of genetic ancestry, genotyping chip, and assessment center as covariates. The proportion of blood pressure variance explained by the allele score and its F statistic were calculated to estimate \(r^2<0.01\) using PLINK.\(^{23}\) We extracted 22 uncorrelated variants as instrumental variables for SBP and 27 uncorrelated variants as instrumental variables for DBP in this sensitivity analysis (Tables S3 and S4).

### Table 1. Distribution of Risk Factors for Individuals in the Analyzed Population That Had a Weighted Allele Score for SBP and DBP Above and Below the Population Median in the Main and Sensitivity Analyses

| Variable | Main analysis (allele score adjusted for BMI) | Sensitivity analysis (allele score not adjusted for BMI) |
|----------|---------------------------------------------|--------------------------------------------------------|
|          | SBP-weighted allele score | DBP-weighted allele score | SBP-weighted allele score | DBP-weighted allele score |
|          | Below median | Above median | Below median | Above median | Below median | Above median | Below median | Above median |
| Age, y; mean (SD) | 55.9 (8.0) | 55.4 (8.1) | 55.8 (8.0) | 55.4 (8.1) | 55.7 (8.0) | 55.6 (8.0) | 55.7 (8.0) | 55.5 (8.0) |
| Sex, n (%) | 54,642 (43.0) | 53,959 (42.4) | 53,842 (42.5) | 53,533 (42.3) | 54,549 (42.9) | 54,052 (42.5) | 53,770 (42.5) | 53,605 (42.3) |
| BMI, kg/m²; mean (SD) | 26.9 (4.5) | 26.7 (4.4) | 26.9 (4.5) | 26.7 (4.4) | 26.9 (4.5) | 26.9 (4.4) | 26.9 (4.5) | 26.9 (4.4) |
| Smoking index, mean (SD)\(\text{t}\) | 0.4 (0.7) | 0.4 (0.7) | 0.4 (0.7) | 0.4 (0.7) | 0.4 (0.7) | 0.4 (0.7) | 0.4 (0.7) | 0.4 (0.7) |
| DBP, mm Hg; mean (SD) | 134.6 (17.8) | 133.8 (18.6) | 135.0 (18.0) | 137.9 (18.5) | 135.5 (18.0) | 137.4 (18.5) | 135.7 (18.2) | 137.3 (18.4) |
| LDL-C, mmol/L; mean (SD) | 3.7 (0.8) | 3.7 (0.8) | 3.7 (0.8) | 3.7 (0.8) | 3.7 (0.8) | 3.7 (0.8) | 3.7 (0.8) | 3.7 (0.8) |

**BMI** indicates body mass index; **DBP**, diastolic blood pressure; **LDL-C**, low-density lipoprotein cholesterol; and **SBP**, systolic blood pressure.

* Socioeconomic status quintiles according to the Townsend deprivation index combining information on social class, employment, car availability, and housing.

†Lifetime smoking index, as detailed by Wootton et al.\(^{22}\)
instrument strength. Cox proportional hazard regression was used to estimate the association of the allele score with the outcomes, incorporating age, sex, principal components 1 to 10 of genetic ancestry, genotyping chip, and assessment center as covariates. As sensitivity analyses, we considered each variant in the allele score separately and performed MR methods that differ in their requisite assumptions regarding the inclusion of pleiotropic variants: random-effects inverse-variance weighted MR, MR-Egger, weighted median MR, and MR-PRESSO. An intercept term in MR-Egger differing from zero can be used to evidence the presence of directional pleiotropy, and MR-PRESSO is able to identify variants with outlying estimates that may in turn be excluded from analyses.

Nonlinear MR

We applied the fractional polynomial method to investigate for evidence of a nonlinear relationship between genetically proxied blood pressure and risk of incident CVD, CAD, and stroke. This approach has been described previously in detail and is outlined in Methods in the Data Supplement. Briefly, we stratified the population into centiles based on residual blood pressure, defined as a participant's blood pressure minus the genetic contribution to blood pressure from the allele score. By doing this, we aimed to compare individuals in the population who would have similar blood pressure values (values in the same centile) if they had the same genetic predisposition. Stratifying on blood pressure directly would introduce collider bias to distort estimates, as blood pressure is on the causal pathway from the genetic variants to CVD. For each centile, we calculated a linear MR estimate for the association of genetically proxied blood pressure with the outcome using the ratio of coefficients method, as described above. Using a flexible semiparametric framework, we then performed a meta-regression of the linear MR estimates obtained for each centile against the mean blood pressure in that centile. A fractional polynomial test was used to investigate whether a nonlinear model fit this meta-regression better than a linear model (further detailed in Methods in the Data Supplement). A Bonferroni correction was applied to account for multiple testing of the 2 blood pressure traits and 3 outcomes, with \( P < 8 \times 10^{-3} \) representing statistical significance. We further conducted a priori–specified subgroup analyses considering...
men and women separately to investigate potential sex-specific effects.

Individuals with elevated blood pressure are more likely to be prescribed antihypertensive medications, and, therefore, exclusion of these individuals from the main analysis could potentially distort MR estimates due to selection effects and introduction of collider bias. Inverse probability weighting was, therefore, performed in a sensitivity analysis to investigate this, as described in Methods in the Data Supplement.

RESULTS

A total of 255,714 participants were included in analyses, after excluding 66,011 individuals with a history of antihypertensive medication use and 6,506 individuals with a history of CVD (but not on antihypertensive medications). There were 10,606 incident CVD events, including 8,430 incident CAD events (68.1% International Classification of Diseases, Tenth Revision, based) and 2,176 incident stroke events. The allele score explained 4.8% and 4.5% of the variance for SBP and DBP, respectively, corresponding to F statistics of 58.6 and 54.1 and low risk of substantial weak instrument bias. The distribution of CVD risk factors for individuals in the analyzed population that had a weighted allele score for SBP and DBP above and below the population median in the main and sensitivity analyses is provided in Table 1. Table S5 provides these data for individuals in the top and bottom deciles of residual blood pressure in the main analysis.

Linear MR

Linear MR analyses identified a strong association of both genetically proxied SBP and DBP with the cardiovascular outcomes. For a 10-mm Hg increase in genetically proxied SBP, the hazard ratio (HR) of incident CVD was 1.49 ([95% CI, 1.38–1.61] \( P=7 \times 10^{-25} \)), incident CAD was 1.50 ([95% CI, 1.38–1.63] \( P=2 \times 10^{-21} \)), and incident stroke was 1.44 ([95% CI, 1.22–1.70] \( P=1 \times 10^{-5} \)). For a 5-mm Hg increase in genetically proxied DBP, the HR of incident CVD was 1.35 ([95% CI, 1.29–1.42] \( P=5 \times 10^{-34} \)), incident CAD was 1.36 ([95% CI, 1.26–1.47] \( P=1 \times 10^{-15} \)), and incident stroke was 1.39 ([95% CI, 1.20–1.62] \( P=2 \times 10^{-5} \)). The MR-Egger test did not detect significant directional

Figure 2. Nonlinear Mendelian randomization considering genetically proxied diastolic blood pressure (DBP) and incident cardiovascular outcomes.

Nonlinear Mendelian randomization considering genetically proxied diastolic blood pressure (DBP) and incident cardiovascular outcomes: (A) all incident cardiovascular disease (CVD) events, (B) incident coronary artery disease (CAD), and (C) incident stroke. Displayed on the x axis are DBP values in mm Hg. The y axis shows the hazard ratio for the respective incident cardiovascular event. Reference is set to a population mean DBP value of 81.8 mm Hg. Gray lines depict the 95% CI.
pleiotropy (Table S6), and MR-PRESSO only identified 16 single-nucleotide polymorphisms as outliers in the analysis of genetically proxied SBP and CAD (Table S2). Similar MR estimates were obtained in sensitivity analyses (Table S6; Figures S1 and S2).

Nonlinear MR
While in some cases the best-fitting fractional polynomial was a nonlinear function, we observed no evidence favoring a nonlinear relationship between genetically proxied blood pressure and the cardiovascular outcomes over a linear one (Figures 1 and 2). This means that any departure from linearity was no greater than would be expected by chance due to random variability. Compared with the population mean SBP of 137 mm Hg, individuals with a genetically proxied SBP of 120 mm Hg had a 47% lower risk of incident CVD (HR, 0.53 [95% CI, 0.49–0.58]; Table 2). Compared with the population mean DBP of 82 mm Hg, individuals with a genetically proxied DBP of 70 mm Hg had a 53% lower risk of incident CVD (HR, 0.47 [95% CI, 0.41–0.53]; Table 2). MR estimates for population subgroups based on stratification into

| Blood pressure | CVD | | CAD | | Stroke |
|----------------|-----|-----|-----|-----|-----|-----|
|                | Hazard ratio 95% CI | Hazard ratio 95% CI | Hazard ratio 95% CI |
| SBP, mm Hg     | Min (102.9) 0.28 0.23–0.33 0.21 0.15–0.29 0.42 0.29–0.63 | 110 0.36 0.32–0.42 0.3 0.24–0.39 0.49 0.36–0.68 | 120 0.53 0.49–0.58 0.49 0.43–0.57 0.62 0.50–0.77 | 130 0.78 0.75–0.81 0.77 0.73–0.81 0.82 0.75–0.90 | 140 1.14 1.12–1.17 1.16 1.12–1.20 1.13 1.07–1.19 | 150 1.67 1.56–1.80 1.7 1.52–1.90 1.63 1.30–2.05 | 160 2.45 2.16–2.78 2.43 2.02–2.93 2.49 1.64–3.79 | 170 3.59 2.99–4.30 3.41 2.64–4.39 4.02 2.12–7.61 | 180 5.25 4.16–6.64 4.68 3.40–6.44 6.87 2.84–16.63 | 190 7.69 5.76–10.26 6.32 4.31–9.26 12.55 3.93–40.10 | Max (194.6) 9.15 6.69–12.52 7.16 4.76–10.76 16.65 4.58–60.56 |
| DBP, mm Hg     | Min (58.6) 0.2 0.15–0.25 0.14 0.09–0.23 0.3 0.16–0.52 | 60 0.22 0.17–0.28 0.17 0.11–0.26 0.31 0.18–0.54 | 65 0.32 0.27–0.39 0.28 0.21–0.38 0.39 0.25–0.62 | 70 0.47 0.41–0.53 0.44 0.36–0.54 0.51 0.37–0.70 | 75 0.65 0.61–0.70 0.64 0.58–0.72 0.67 0.55–0.81 | 80 0.9 0.88–0.91 0.9 0.87–0.92 0.9 0.85–0.95 | 85 1.21 1.17–1.24 1.21 1.15–1.26 1.23 1.11–1.35 | 90 1.6 1.48–1.72 1.57 1.41–1.75 1.71 1.32–2.21 | 95 2.08 1.85–2.35 1.99 1.68–2.35 2.42 1.58–3.70 | 100 2.68 2.28–3.14 2.45 1.97–3.06 3.5 1.92–6.39 | 105 3.4 2.78–4.15 2.97 2.28–3.88 5.16 2.35–11.34 | Max (111.7) 4.6 3.58–5.89 3.72 2.70–5.14 8.88 3.11–25.27 |

**DISCUSSION**
By applying nonlinear MR methods in the UK Biobank, we were able to examine the shape of the relationship between genetically proxied blood pressure and incident CVD in a population without a history of CVD or antihypertensive medication use. We found no evidence favoring nonlinear relationships between genetically proxied SBP or DBP and risk of the cardiovascular outcomes.
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over linear ones. Similar results were obtained when considering males and females separately.

Blood pressure control represents a global health challenge, and hypertension thresholds have been lowered in recent consensus guidelines. The MR estimates obtained in this study may be used to quantify the effect of a persistent, lifelong reduction in blood pressure on the primary prevention of CVD and highlight the potential gains of clinical and public health interventions that achieve this. Importantly, they support the notion that for a population without a history of CVD or antihypertensive medication use, a similar relative reduction in CVD risk will be observed irrespective of baseline blood pressure, including for individuals who have normal blood pressure. This means that fixed changes in blood pressure will lead to similar changes in CVD risk on the HR scale. On the absolute scale, risk reduction will be greater for those with a higher baseline blood pressure. This finding is consistent with previous large-scale observational analyses performed in individuals free of CVD at baseline. In contrast, excessive blood pressure reduction in patients with atherosclerotic disease can reduce organ perfusion and increase CVD risk, and it is, therefore, important that our findings are not extrapolated to infer the effect of blood pressure lowering in individuals with preexisting CVD. It is also important to appreciate that absolute risk reduction conferred from blood pressure lowering will remain greatest for those with the highest blood pressure. Our current data support the concept that risk factor targeting in low- and medium-risk individuals on a population-wide level is likely to also substantially contribute to reducing the burden of CVD. Dietary modification and reduced sodium consumption represent examples of public health strategies that can be adopted to achieve this.

We found no evidence for a J-shaped association of either genetically proxied SBP or DBP with any of the outcomes. This contrasts the findings of a recent observational study using data from 1.3 million general outpatients with a low prevalence of CAD, which identified a J-shaped association of blood pressure with the

Figure 3. Nonlinear Mendelian randomization considering genetically proxied systolic blood pressure (SBP) and incident cardiovascular outcomes split by sex.

Nonlinear Mendelian randomization considering genetically proxied systolic blood pressure (SBP) and incident cardiovascular outcomes split by sex: (A) all incident cardiovascular disease (CVD) events in men, (B) incident coronary artery disease (CAD) in males, and (C) incident stroke in men. D–F, Equivalent analyses in women. Displayed on the x-axis are SBP values in mm Hg. The y-axis shows the hazard ratio for the respective incident cardiovascular event. Reference is set to a mean SBP value of 136.5 mm Hg. Gray lines depict the 95% CI.
A composite outcome of myocardial infarction and stroke. This J shape was only partially attenuated after adjusting for age, ethnicity, and comorbidities, and there remains the possibility that residual unknown or unmeasured confounding factors are responsible for the discrepancy with our findings. A systematic review and meta-analysis of blood pressure–lowering trials considering 613,815 participants from 123 studies found no trend for CVD risk reduction per 10 mm Hg lower SBP when stratifying trials by mean baseline SBP. In the SPRINT trial, SBP lowering to <120 mmHg as compared with 140 mm Hg resulted in fewer major cardiovascular events. The findings from our current MR study additionally support a relative CVD risk reduction from blood pressure lowering below this level in patients without a history of CVD.

Our study has a number of strengths. By employing randomly allocated genetic variants as proxies for the effect of modifying blood pressure, we were able to use the MR paradigm to overcome the environmental confounding bias that can limit causal inference in observational association studies. The implementation of both linear and nonlinear MR methods within the comprehensive UK Biobank resource enabled us to efficiently study the relationships of genetically proxied SBP and DBP with incident CVD, CAD, and stroke, including in sex-stratified analyses. Importantly, the fractional polynomial method allowed us to investigate for evidence of nonlinear associations.

Our study also has limitations. This work only considered participants without a history of CVD or antihypertensive medication use, and its findings should not be extrapolated to populations with established CVD. Individuals that reported taking antihypertensive medications were excluded to allow for meaningful stratification into blood pressure quantiles, and as such, there is the possibility that ascertainment bias may have been introduced. Reassuringly, similar findings were obtained in inverse probability weighting sensitivity analyses, suggesting that any such bias is unlikely to be affecting our conclusions. The employed MR approach assumes that the genetic variants utilized as proxies for blood pressure
do not affect CVD risk through alternative (pleiotropic) pathways—an assumption that cannot be tested and if violated could introduce bias to the obtained estimates. Our used MR method also explores the effects of lifetime changes in blood pressure, and its estimates should, therefore, not be extrapolated to quantify the effect of blood pressure modification in adult life, such as through use of antihypertensive medications. Finally, there were differences in the distribution of risk factors between individuals in the highest and lowest deciles of residual blood pressure (Table S5), suggesting that this MR analysis may still be vulnerable to environmental confounding.

**PERSPECTIVES**

For a population without a history of CVD or antihypertensive medication use, genetically proxied blood pressure reduction was associated with lower CVD risk at all levels of blood pressure. These findings provide evidence to support that public health interventions achieving persistent, population-wide blood pressure reduction will be of considerable benefit in the primary prevention of CVD.

**ARTICLE INFORMATION**

Received October 16, 2020; accepted February 16, 2021.

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**Acknowledgments**

This research has been conducted using the UK Biobank Resource (UK Biobank application No. 2532). The UK Biobank data are available on application at https://www.ukbiobank.ac.uk/register-apply. The Trndelag Health Study is a collaboration between the HUNT Research Centre (Faculty of Medicine and Health Sciences, Norwegian University of Science and Technology), Nord-Trndelag County Council, Central Norway Regional Health Authority, and the Norwegian Institute of Public Health. The genotype quality control and imputation in HUNT has been conducted by the K.G. Jebsen Center for Genetic Epidemiology, Department of Public Health and Nursing, Faculty of Medicine and Health Sciences, NTNU, Norwegian University of Science and Technology. D. Gill, R. Malik, M. Vujkovic, S. Burgess, and M.K. Georgakis designed the study. R. Malik, M. Vujkovic, J.N. Hellwege, MM. Shuey, TL. Edwards, B.M. Brumpton, T. Rogne, C. Fava, and A. Giuntella analyzed the data. R. Malik, D. Gill, J.N. Hellwege, B.M. Brumpton, and M.K. Georgakis drafted the manuscript. All authors interpreted the results and critically revised the manuscript for intellectual content.

**Sources of Funding**

This work was supported by the UK National Institute for Health Research Cambridge Biomedical Research Centre. M.K. Georgakis is funded by a scholarship from the Onassis Foundation. S.M. Damrauer was supported by the Department of Veterans Affairs Office of Research and Development (IK2-CX001780). P. Elliott acknowledges support from the British Heart Foundation (REF:18/4/43215), the Medical Research Council (MR/S019669/1), the National Institute for Health Research Imperial Biomedical Research Centre, Imperial College London (RF/03), the UK Dementia Research Institute (DRI) at Imperial College London funded by UK DRI, Ltd (funded by the Medical Research Council, Alzheimer’s Society, Alzheimer’s Research UK), and Health Data Research (HDR) UK London funded by HDR UK Ltd (funded by a consortium led by the Medical Research Council Grant: MR/K020283/1). J.N. Hellwege is supported by K12 HD04348. MM. Shuey is funded by the National Institutes of Health (DK108444). S. Burgess is supported by a Sir Henry Dale Fellowship jointly funded by the Wellcome Trust and the Royal Society (204623/Z/16/Z). M. Dicksang acknowledges funding from the European Union Horizon 2020 research and innovation programme (666881), SVSDs@target (667375), CoSTREAM, Synergy (EXC 2145 Synergy, ID 390957198), the CRC 1123 (BS and project DI 722/13-1), the Corona Foundation, the LMU-excellent Fund, the eMed program (eAtheroSymMed), and the FP7/2007-2013 European Union project CVgenes@target (Health-F2-2013-601456). D. Gill is supported by the Wellcome Trust 4i Programme (203928/Z/16/Z) and British Heart Foundation Centre of Research Excellence (REF:18/4/43215) at Imperial College London and, a National Institute for Health Research Clinical Lecture Ship at St. George’s, University of London (CL-2020-16-001). The BioVU data set used for the analyses described was obtained from the Vanderbilt University Medical Center’s BioVU, which is supported by numerous sources: institutional funding, private agencies, and federal grants. These include the National Institutes of Health (NIH)–funded Shared Instrumentation Grant S10RR025141 and CTSA grants UL1TR000243, UL1TR000445, and UL1TR0024975. Genomic data are also supported by investigator-led projects that include U01HG004798, R01NS052830, RC2GM092618, P50GM151350, U01HG006378, U19HL14962, and R01HD074711 and additional funding sources listed at https://victr.vumc.org/biovu-funding/. The genotyping in HUNT was financed by the NIH, University of Michigan, the Research Council of Norway; the Liaison Committee for Education, Research and Innovation in Central Norway, and the Joint Research Committee between St. Olav’s Hospital and the Faculty of Medicine and Health Sciences, NTNU.

**Disclosures**

D. Gill is employed part-time by Novo Nordisk and has received consultancy fees from Abbott Laboratories. S.M. Damrauer has received grants from the US Department of Veterans Affairs, Calico Labs, and Renalytix AI plc outside the submitted work. The other authors report no conflicts.

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