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Systematic Evaluation of Pleiotropy Identifies 6 Further Loci Associated With Coronary Artery Disease

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ABSTRACT

BACKGROUND Genome-wide association studies have so far identified 56 loci associated with risk of coronary artery disease (CAD). Many CAD loci show pleiotropy; that is, they are also associated with other diseases or traits.

OBJECTIVES This study sought to systematically test if genetic variants identified for non-CAD diseases/traits also associate with CAD and to undertake a comprehensive analysis of the extent of pleiotropy of all CAD loci.

METHODS In discovery analyses involving 42,335 CAD cases and 78,240 control subjects we tested the association of 29,383 common (minor allele frequency >5%) single nucleotide polymorphisms available on the exome array, which included a substantial proportion of known or suspected single nucleotide polymorphisms associated with common diseases or traits as of 2011. Suggestive association signals were replicated in an additional 30,533 cases and 42,530 control subjects. To evaluate pleiotropy, we tested CAD loci for association with cardiovascular risk factors (lipid traits, blood pressure phenotypes, body mass index, diabetes, and smoking behavior), as well as with other diseases/traits through interrogation of currently available genome-wide association study catalogs.

RESULTS We identified 6 new loci associated with CAD at genome-wide significance: on 2q37 (*KCNJ13-GIGYF2*), 6p21 (*C2*), 11p15 (*MRVI1-CTR9*), 12q13 (*LRP1*), 12q24 (*SCARB1*), and 16q13 (*CETP*). Risk allele frequencies ranged from 0.15 to 0.86, and odds ratio per copy of the risk allele ranged from 1.04 to 1.09. Of 62 new and known CAD loci, 24 (38.7%) showed statistical association with a traditional cardiovascular risk factor, with some showing multiple associations, and 29 (47%) showed associations at $p < 1 \times 10^{-4}$ with a range of other diseases/traits.

CONCLUSIONS We identified 6 loci associated with CAD at genome-wide significance. Several CAD loci show substantial pleiotropy, which may help us understand the mechanisms by which these loci affect CAD risk. (J Am Coll Cardiol 2017;69:823-36) © 2017 The Authors. Published by Elsevier on behalf of the American College of Cardiology Foundation. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

Over the past decade, genome-wide association studies (GWAS) have identified several thousand robust associations ($p < 5 \times 10^{-8}$) for a range of human traits and diseases. For coronary artery disease (CAD), 56 such loci have been identified so far, explaining ~15% of the disease's heritability (1,2). Approximately one-third of the CAD loci also show association with a known or putative

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cardiovascular risk factor, particularly blood pressure and lipid traits (2). Furthermore, several loci show association with other diseases; for example, the CAD-associated variants in the chromosome 9p21 locus also associate with risk of stroke as well as abdominal, aortic, and intracranial aneurysms (3,4). These observations suggest that a comprehensive analysis of variants associated with other diseases and traits might not only identify additional loci associated with risk of CAD, but also provide important insights into genetic mechanisms shared by different diseases.

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Here, we leveraged the HumanExome BeadChip (Illumina, Inc., San Diego, California) to test the contribution of 29,393 common variants of single nucleotide polymorphisms (SNPs) (minor-allele frequency >5%) for association with CAD. The variants included the majority of reported trait/disease-associated lead SNPs in the National Human Genome Research Institute GWAS catalogue as of August 2011, as well as a number of associations for complex diseases unpublished at that time, variants in the human leukocyte antigen (HLA) region, and a scaffold of approximately 5,000 SNPs placed

**ABBREVIATIONS
 AND ACRONYMS**

- BMI** = body mass index
- CAD** = coronary artery disease
- CETP** = cholesteryl ester transfer protein
- eQTL** = expression quantitative trait locus
- GWAS** = genome-wide association study
- HDL** = high-density lipoprotein
- LD** = linkage disequilibrium
- LDL** = low-density lipoprotein
- SNP** = single nucleotide polymorphism

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on the array for identity by descent testing. The results of an analysis of rare (minor-allele frequency <5%) coding sequence (“exome”) variants on this array with CAD were recently reported (5).

We identified 6 new loci associated at genome-wide significance with CAD, annotated these, and undertook a detailed examination of the extent of pleiotropy of these loci as well the previously known CAD loci.

METHODS

The study consisted of discovery and replication phases and has been described in more detail elsewhere (5). Briefly, the discovery cohort included 42,335 cases and 78,240 control subjects from 20 individual studies (Online Table 1); the replication cohort, which was separately assembled and

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Drs. Akinsanya, Wu, Yin, and Reilly are employees of Merck Sharp & Dohme; and Dr. Vogt was an employee of Merck when aspects of this research was conducted, but is now retired from Merck. A cholesteryl ester transfer protein inhibitor, Anacetrapib (MK-0859), is currently undergoing clinical investigation in the REVEAL outcome trial sponsored by Merck Sharp & Dohme. Dr. Schick is an employee of Recombin. Dr. Dube has equity in DalCor Pharmaceuticals. Dr. McCarthy is a member of advisory boards for Pfizer and Novo Nordisk; has received honoraria from Pfizer, Novo Nordisk, and Eli Lilly; and has received research funding provided by Pfizer, Novo Nordisk, Eli Lilly, Servier, Sanofi-Aventis, Janssen, Roche, Boehringer-Ingelheim, Takeda, Merck, and AstraZeneca. Dr. Ferrieres has received grants from Merck Sharp & Dohme, Amgen, and Sanofi. Dr. Sattar has served as a consultant for Amgen and Sanofi. Dr. Butterworth has received grants from Pfizer and Merck. Dr. Danesh has served as a consultant for Takeda; has served on the Novartis Cardiovascular & Metabolic Advisory Board and International Cardiovascular and Metabolism Research and Development Portfolio Committee of Novartis; has served on the UK Atherosclerosis Advisory Board of Merck Sharp & Dohme; has served on the advisory board of Sanofi; has served on the Pfizer Population Research Advisory Panel; and has financial relationships with the British Heart Foundation, BUPA Foundation, diaDexus, European Research Council, European Union, Evelyn Trust, Fogarty International Centre, GlaxoSmithKline, Merck, National Heart, Lung, and Blood Institute, National Health Service Blood and Transplant, National Institute for Health Research, National Institute of Neurological Disorders and Stroke, Novartis, Pfizer, Roche, Sanofi, Takeda, The Wellcome Trust, UK Biobank, University of British Columbia, and UK Medical Research Council. 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ascertained to have no sample overlap with the discovery cohorts, included 30,533 cases and 42,530 control subjects from 8 studies (Online Table 2). With the exception of participants from 2 studies in the replication cohort who were of South Asian ancestry, all participants were of European ancestry (Online Table 2).

Samples were genotyped on the Illumina HumanExome BeadChip versions 1.0 or 1.1, or the Illumina OmniExome (which includes markers from the HumanExome BeadChip) arrays followed by quality control procedures as previously described (5).

STATISTICAL ANALYSIS. In discovery samples that passed quality control procedures, we performed individual tests for association of the selected variants with CAD in each study separately, using logistic regression analysis with principal components of ancestry as covariates (5). We combined evidence across individual studies using an inverse-variance weighted fixed-effects meta-analysis. Heterogeneity was assessed by Cochran's Q statistic (6). In the discovery phase, we defined suggestive novel association as a meta-analysis p value $\leq 1 \times 10^{-6}$.

For variants with suggestive association, we performed association analysis in the replication studies (Online Appendix). We defined significant novel associations as those nominally significant ($p < 0.05$) in the replication study and with an overall (discovery and replication combined) p value $< 5 \times 10^{-8}$.

BIOINFORMATICS ANALYSIS. To identify any association between the novel loci and gene expression traits, we performed a systematic search of cis-expression quantitative trait loci (eQTL) (described in the Online Appendix). To identify candidate causal SNPs at the new loci, we annotated each of the lead variants as well as SNPs in high linkage disequilibrium (LD) ($r^2 > 0.8$) on the basis of position, overlap with regulatory elements, and in silico SNP prioritization tools (Online Appendix).

For both the novel loci and all previously reported CAD loci (1,2), we tested the association of the lead CAD-associated variant (or, if unavailable, a proxy) with traditional cardiovascular risk factors using publicly available GWAS meta-analyses datasets for systolic, diastolic, and pulse pressures (7,8); low-density lipoprotein (LDL) cholesterol level; high-density lipoprotein (HDL) cholesterol level; triglycerides level (9,10); type 2 diabetes mellitus (11); body mass index (BMI) (12); and smoking quantity (13). The maximum size of these datasets ranged from 41,150 to 339,224 individuals. For variants available

on the exome array with a known genome-wide association with a risk factor, we also compared the magnitude of the reported association with the risk factor to the observed association with CAD in our analysis.

To identify any associations with other diseases or traits, we searched version 2 of the GRASP (Genome-Wide Repository of Associations between SNPs and Phenotypes) database (14) and the National Human Genome Research Institute-European Bioinformatics Institute GWAS catalog (15), plus we collected all associations below 1×10^{-4} . For all associations, we identified the lead variant for that trait or disease and calculated pairwise LD with the lead CAD-associated variant using the SNAP web server (16).

RESULTS

In the discovery cohort, 28 variants not located in a known CAD locus (defined as ± 300 kb from the published lead SNP) showed association with CAD at a p value $< 1 \times 10^{-6}$ (Online Table 3). No marked heterogeneity was observed, justifying the use of a fixed-effects model. We then tested these 28 variants for replication, and 6 variants showed both a nominally significant ($p < 0.05$) association in the replication cohort and a combined discovery and replication meta-analyses p value exceeding the threshold for genome-wide significance ($p < 5 \times 10^{-8}$) (Table 1). As typical for GWAS findings, the risk alleles were common (allele frequencies ranging from 15% to 86%), and the risk increase per allele was modest (ranging from 4% to 9%) (Table 1).

ANNOTATION OF NOVEL LOCI. Forest and regional association plots for the 6 novel loci are shown in Online Figures 1 and 2, respectively. Interrogation of the 1000 Genomes Project phase 1 EUR data using Haploreg (BROAD Institute, Massachusetts Institute of Technology and Harvard, Boston, Massachusetts) (17) showed that the number of SNPs in high LD ($r^2 > 0.8$) with the lead variant varied between 1 (*LRP1* locus and *CETP* locus) and 111 (*KCNJ13-GIGYF2* locus) (Online Table 4). Apart from the lead variant at the *KCNJ13-GIGYF2* locus, which is a nonsynonymous SNP, none of the other loci had a variant affecting protein sequence in high LD with the lead variant.

Notable cis-eQTL findings for the new loci are shown in Online Table 5 and functional annotation of the lead variant and variants in high LD appear in Online Figure 3. The main findings from these analyses are discussed here locus by locus.

16q13. The lead variant, rs1800775, also known as -629C>A, is in the promoter of the cholesterol

TABLE 1 Novel Loci Significantly Associated With CAD

| Lead Variant | Locus | Locus Name | A1/2 (Freq) | Discovery | | Replication | | Meta-Analysis | | Known Associations With Lead Variant | | |
|--------------|-------|--------------|-------------|------------------------|------------------|------------------------|------------------------|------------------|-----------------------|--------------------------------------|------------------------|------------------------------------|
| | | | | Cases/Control Subjects | OR (95% CI) | p Value | Cases/Control Subjects | OR (95% CI) | p Value | | OR (CI 95%) | p Value |
| rs1801251 | 2q37 | KCNJ13-GGYF2 | A/G (0.35) | 42,332/78,229 | 1.06 (1.04-1.08) | 1.46×10^{-8} | 30,528/42,521 | 1.03 (1.01-1.06) | 0.007 | 1.05 (1.03-1.06) | 1.48×10^{-9} | — |
| rs3130683 | 6p21 | C2 | T/C (0.86) | 39,494/72,267 | 1.09 (1.06-1.13) | 7.87×10^{-8} | 30,450/42,485 | 1.09 (1.05-1.14) | 2.97×10^{-5} | 1.09 (1.07-1.12) | 1.04×10^{-11} | — |
| rs11042937 | 11p15 | MRVI1-CTR9 | T/G (0.49) | 42,335/78,234 | 1.05 (1.03-1.07) | 3.21×10^{-8} | 30,533/42,527 | 1.03 (1.00-1.05) | 0.019 | 1.04 (1.03-1.06) | 1.18×10^{-8} | Bipolar disorder and schizophrenia |
| rs1172113 | 12q13 | LRP1 | C/T (0.41) | 42,335/78,234 | 1.06 (1.04-1.08) | 1.78×10^{-8} | 28,503/36,433 | 1.06 (1.03-1.08) | 1.16×10^{-6} | 1.06 (1.04-1.07) | 9.25×10^{-14} | Migraine, pulmonary function |
| rs11057830 | 12q24 | SCARB1 | A/G (0.15) | 42,331/78,237 | 1.09 (1.06-1.11) | 3.69×10^{-10} | 20,395/30,592 | 1.07 (1.03-1.11) | 0.0003 | 1.08 (1.06-1.10) | 4.61×10^{-13} | Vitamin E level |
| rs1800775 | 16q13 | CETP | C/A (0.51) | 38,810/62,756 | 1.06 (1.04-1.08) | 2.21×10^{-8} | 22,445/32,148 | 1.03 (1.00-1.05) | 0.032 | 1.04 (1.03-1.06) | 9.83×10^{-9} | HDL cholesterol |

A1/2 = allele 1/allele 2; CAD = coronary artery disease; CI = confidence interval; freq = frequency of allele 1; HDL = high-density lipoprotein; OR = odds ratio for disease for carriers of allele 1.

ester transfer protein (*CETP*) gene, which mediates the transfer of cholesteryl esters from HDL cholesterol to other lipoproteins and was placed on the array because of its association with plasma HDL cholesterol level (9,10). The risk (C) allele is associated with lower HDL cholesterol and modest increases in plasma LDL cholesterol and triglycerides levels (9,10). Previous studies have shown that rs1800775 is itself functional in that the C allele disrupts binding of the Sp1 transcription factor resulting in increased promoter activity (18). This is in agreement with our annotation, which predicts this to be more likely to be a functional SNP than the only other SNP in high LD, rs3816117 (Online Figure 3). Consistent with this, we also found associations between rs1800775 and *CETP* expression (r^2 of 0.77) with the best eSNP (i.e., the lead SNP for the eQTL) in monocytes and liver (Online Table 5), and previous studies have shown that the variant is also associated with plasma *CETP* level (19,20).

12q24. The lead variant, rs11057830, and all 8 variants in high LD are located in a region of approximately 10 kb in intron 1 of *SCARB1*, which encodes SR-B1, a receptor for HDL cholesterol. Other variants at this locus have been associated with HDL cholesterol level (9,10). However, these HDL cholesterol variants are not in high LD with the CAD-associated variants identified here, which only have a modest association with plasma HDL cholesterol level (Online Table 6), but a stronger association with plasma LDL cholesterol and triglycerides levels (Table 2). rs11957830 was included on the array because of an association of the A allele (CAD risk-associated allele) with higher levels of vitamin E (Table 3) (21). Variants in high LD with the CAD risk allele at rs11057830 have also been associated with increased lipoprotein-associated phospholipase A₂ (Lp-PLA₂) activity (22). Analysis of eQTL identified an association between rs11057841 ($r^2 = 0.92$ with the lead variant), and expression of *SCARB1* in the intestine (Online Table 5). Functional annotation of the locus did not identify a strong candidate causal SNP, but rs10846744 ($r^2 = 0.94$ with the lead variant) overlaps a deoxyribonuclease I hypersensitivity peak in a region bound by several transcription factors (Online Figure 3).

12q13. The lead variant, rs11172113, is in intron 1 of *LRP1* (LDL receptor-related protein-1) and only has 1 other adjacent SNP in high LD (Central Illustration, Online Table 4). The risk (C) allele of the lead variant has previously been associated with reduced risk of migraine (23), and there is an association of the alternate (T) allele with reduced lung function (24). There are also associations at this locus for abdominal aortic aneurysm (25) and triglyceride levels (10);

however, these variants are in modest or low LD to the CAD-associated SNP (r^2 of 0.54 and 0.07, respectively). The lead variant overlaps a region containing peaks in deoxyribonuclease I hypersensitivity in several cells and tissues, including aortic smooth muscle cells, within a predicted enhancer element (Online Figure 3). We found associations between the CAD risk allele at rs11172113 and reduced expression of *LRP1* in atherosclerotic and nonatherosclerotic arterial wall, as well as eQTLs in omental and subcutaneous adipose tissue (Online Table 5).

11p15. The lead variant, rs11042937, at this locus lies in an intergenic region between *MRVI1* (murine retrovirus integration site 1 homolog) encoding inositol-trisphosphate receptor-associated cyclic guanosine monophosphate kinase substrate, a mediator of smooth muscle tone and *CTR9* that encodes a component of the PAF1 complex with some SNPs in high LD located within intron 1 of *MRVI1* (Online Figure 3). The lead variant was included on the array because of a suggestive association with bipolar disorder and schizophrenia (26). There was no association of the locus with any cardiovascular risk factors, and we did not identify any eQTLs. Evidence for a regulatory function for either the lead variant or any of the SNPs in high LD was also weak (Online Figure 3).

6p21. The lead variant, rs3130683, lies in the HLA complex in intron 1 of *C2*, which encodes the complement C2 protein. There are just 14 SNPs in high LD with the lead variant (Online Table 4), but the CAD signal spans a region of approximately 300 kb including more than 20 genes (Online Figure 3). Apart from a single synonymous variant in *HSPA1A* (heat shock 70kDa protein 1A), the other high LD variants are noncoding with several of the variants showing evidence for regulatory functionality (Online Figure 3). Although there is a large number of eQTLs in the HLA region, most of these are variants with modest ($r^2 < 0.5$) LD with the CAD-associated variants, and the only eQTL of note was with *CYP21A2* (cytochrome P450, family 21, subfamily A, polypeptide 2) expression in whole blood (Online Table 5). rs3869109, another variant at the HLA locus approximately 700 kb away from the new lead variant, has been reported to be associated with CAD (27). In our discovery cohort, rs3869109 has a p value of association with CAD of 0.23.

2q37. The lead variant, rs1801251, was included on the array for identity by descent testing; rs1801251 causes a threonine to isoleucine amino acid change at position 95 in *KCNJ13*, an inwardly rectifying potassium channel protein. However, this is not predicted to be functionally important. There is

TABLE 2 Significant Associations of CAD Variants With Selected CV Risk Factors*

| Locus | Locus Name | Lead Variant | Trait | Effect | p Value |
|-------------------|----------------------------|--------------|-------|-------------------|------------------------|
| New Loci | | | | | |
| 6p21 | <i>C2</i> | rs3130683 | T2D | 1.12 [†] | 2.7×10^{-5} |
| | <i>SCARB1</i> | rs11057830 | LDL | 0.006 | 2.6×10^{-5} |
| 16q13 | <i>CETP</i> | rs1800775 | TG | 0.022 | 8.3×10^{-5} |
| | | | LDL | 0.041 | 8.5×10^{-24} |
| | | | HDL | -0.202 | 3.3×10^{-644} |
| | | | TG | 0.04 | 1.3×10^{-26} |
| Known Loci | | | | | |
| 1p32 | <i>PCSK9</i> | rs11206510 | LDL | 0.083 | 2.4×10^{-53} |
| 1p13 | <i>SORT1</i> | rs602633 | LDL | 0.159 | 1.5×10^{-261} |
| | | | HDL | -0.033 | 3.5×10^{-14} |
| 2p24 | <i>APOB</i> | rs515135 | LDL | 0.139 | 1.1×10^{-178} |
| 2p21 | <i>ABCG5-ABCG8</i> | rs6544713 | LDL | 0.081 | 4.84×10^{-83} |
| 4q32 | <i>GUCY1A3</i> | rs7692387 | DBP | 0.326 | 3.4×10^{-5} |
| 5q31 | <i>SLC22A4-SLC22A5</i> | rs273909 | LDL | 0.022 | 2.3×10^{-5} |
| 2q33 | <i>WDR12</i> | rs6725887 | LDL | -0.026 | 1.3×10^{-5} |
| 6q25 | <i>LPA</i> | rs3798220 | LDL | 0.158 | 6.1×10^{-11} |
| | | | LDL | 0.019 | 1.3×10^{-6} |
| 7q32 | <i>ZC3HC1</i> | rs11556924 | DBP | NA | 1.8×10^{-5} |
| | | | HDL | -0.018 | 1.3×10^{-5} |
| 7q36 | <i>NOS3</i> | rs3918226 | SBP | 0.96 | 1.1×10^{-6} |
| | | | DBP | 0.81 | 2.2×10^{-9} |
| 8p21 | <i>LPL</i> | rs264 | HDL | -0.098 | 8×10^{-77} |
| 8q24 | <i>TRIB1</i> | rs2954029 | TG | 0.093 | 2.4×10^{-84} |
| | | | LDL | 0.056 | 2.1×10^{-50} |
| | | | HDL | -0.04 | 2.7×10^{-29} |
| | | | TG | 0.076 | 1×10^{-107} |
| 9q34 | <i>ABO</i> | rs579459 | LDL | 0.067 | 2.4×10^{-44} |
| 10q24 | <i>CYP17A1-CNNM2-NT5C2</i> | rs12413409 | SBP | 1.034 | 2×10^{-9} |
| | | | DBP | 0.483 | 3.4×10^{-5} |
| | | | PP | 0.56 | 5.7×10^{-8} |
| | | | BMI | -0.03 | 2.2×10^{-8} |
| 11q23 | <i>ZNF259-APOA5-APOA1</i> | rs964184 | LDL | 0.086 | 2×10^{-26} |
| | | | HDL | -0.107 | 6.1×10^{-48} |
| | | | TG | 0.234 | 6.6×10^{-244} |
| 12q24 | <i>SH2B3</i> | rs3184504 | LDL | -0.027 | 4.2×10^{-12} |
| | | | HDL | -0.026 | 4.1×10^{-12} |
| | | | SBP | 0.598 | 2×10^{-9} |
| | | | DBP | 0.483 | 8.8×10^{-6} |
| | | | BMI | -0.131 | 9.4×10^{-6} |
| 15q26 | <i>FURIN-FES</i> | rs17514846 | SBP | 0.509 | 1.2×10^{-5} |
| 17p13 | <i>SMG6</i> | rs2281727 | BMI | 0.015 | 3.64×10^{-6} |
| 18q21 | <i>PMAIP1-MC4R</i> | rs663129 | HDL | -0.026 | 5.5×10^{-9} |
| | | | BMI | 0.056 | 8.8×10^{-53} |
| 19p13 | <i>LDLR</i> | rs1122608 | LDL | 0.074 | 8.5×10^{-57} |
| 19q13 | <i>APOE-APOC1</i> | rs2075650 | LDL | 0.177 | 1.7×10^{-214} |
| | | | HDL | -0.055 | 9.7×10^{-26} |
| | | | TG | 0.044 | 2.3×10^{-21} |
| | | | BMI | -0.026 | 1.3×10^{-8} |
| | | | LDL | 0.363 | 6.6×10^{-397} |
| | | | HDL | -0.051 | 1.9×10^{-10} |
| | | rs445925 | LDL | 0.363 | 6.6×10^{-397} |
| | | | HDL | -0.051 | 1.9×10^{-10} |
| | | | TG | 0.101 | 3.6×10^{-39} |

*Effects are either absolute beta estimates of the association of the CAD risk allele on the trait (with a positive association indicating a higher value of the trait per copy of the risk allele) or log odds ratio for diabetes mellitus, tper copy of the risk allele. This table only includes associations that passed Bonferroni correction.

BMI = body mass index; CAD = coronary artery disease; CV = cardiovascular; DBP = diastolic blood pressure; HDL = high-density lipoprotein; LDL = low-density lipoprotein cholesterol level; T2D = type 2 diabetes mellitus; TG = triglycerides.

TABLE 3 Association of CAD Loci With Other Diseases or Traits

| Locus | Locus Name | Disease or Trait | Disease or Trait Lead SNP | p Value | Direction | r ² (CAD Lead and Disease or Trait Lead) |
|-------------------|-------------------------|---|---------------------------|-----------------------|-----------|---|
| New Loci | | | | | | |
| 6p21 | <i>C2</i> | Systemic lupus erythematosus | rs3130342 | 9.3×10^{-7} | + | 0.87 |
| | | Primary biliary cirrhosis | rs3134954 | 1×10^{-5} | + | 0.86 |
| | | Multiple sclerosis | rs3134954 | 3.2×10^{-9} | - | 0.86 |
| 11p15 | <i>MRV11-CTR9</i> | Bipolar disorder and schizophrenia | rs2018368 | 1×10^{-6} | + | 0.96 |
| 12q13 | <i>LRP1</i> | Migraine | rs11172113 | 4.3×10^{-9} | - | Same SNP |
| | | Lung function (FEV1/FVC) | rs11172113 | 1.2×10^{-8} | + | Same SNP |
| | | Cervical artery dissection | rs11172113 | 3×10^{-7} | - | Same SNP |
| 12q24 | <i>SCARB1</i> | Lp-PLA ₂ activity | rs11057841 | 6.1×10^{-14} | + | 0.92 |
| | | Circulating vitamin E levels | rs11057830 | 8.2×10^{-9} | + | Same SNP |
| | | Lp-PLA ₂ mass | rs10846744 | 6.8×10^{-6} | + | 0.94 |
| Known Loci | | | | | | |
| 1p32 | <i>SORT1</i> | Lp-PLA ₂ activity | rs7528419 | 1.3×10^{-17} | + | 0.9 |
| | | Metabolic syndrome domains (Atherogenic dyslipidemia-PC1) | rs12740374 | 8×10^{-16} | + | Same SNP |
| | | Lp-PLA ₂ mass | rs7528419 | 7.1×10^{-5} | + | 0.9 |
| 1q21 | <i>IL6R</i> | C-reactive protein | rs4845625 | 4.2×10^{-7} | + | Same SNP |
| 2p21 | <i>ABCG5-ABCG8</i> | Serum phytosterol | rs4245791 | 2.2×10^{-70} | + | 1 |
| 2p11 | <i>VAMP5-VAMP8-GGCX</i> | Prostate cancer | rs10187424 | 2.7×10^{-15} | - | 0.87 |
| 2q33 | <i>WDR12</i> | Cerebral white matter hyperintensities burden | rs6705330 | 5.7×10^{-5} | + | 1 |
| 3q22 | <i>MRAS</i> | Coronary artery calcification | rs2306374 | 2.7×10^{-5} | + | 1 |
| 4q12 | <i>REST-NOA1</i> | Height | rs17081935 | 6.7×10^{-17} | + | 0.95 |
| 4q31 | <i>EDNRA</i> | Carotid intima media thickness | rs1878406 | 7×10^{-12} | + | Same SNP |
| | | Intracranial aneurysm | rs6842241 | 2.4×10^{-9} | + | 0.94 |
| | | Coronary artery calcification | rs9349379 | 4×10^{-22} | + | Same SNP |
| 6p24 | <i>PHACTR1</i> | Cervical artery dissection | rs9349379 | 1×10^{-11} | - | Same SNP |
| | | Migraine | rs9349379 | 5×10^{-8} | - | Same SNP |
| | | Pulse wave velocity | rs7750679 | 5.4×10^{-5} | + | 1 |
| 6q25 | <i>LPA</i> | Lipoprotein (a) | rs3798220 | 1.6×10^{-49} | + | Same SNP |
| | | Colorectal cancer | rs7758229 | 5.6×10^{-9} | - | 0.85 |
| 7p21 | <i>HDAC9</i> | Stroke (large vessel stroke) | rs11984041 | 1.9×10^{-11} | + | 1 |
| 7q22 | 7q22 | Endometriosis | rs10953541 | 3.2×10^{-5} | + | Same SNP |
| 8q24 | <i>TRIB1</i> | Metabolic syndrome domains (Atherogenic dyslipidemia-PC1) | rs2954021 | 1.2×10^{-11} | + | Same SNP |
| | | Adiponectin levels | rs2954021 | 1.8×10^{-5} | + | Same SNP |
| | | Serum creatinine | rs2954021 | 2.3×10^{-5} | + | Same SNP |
| 9p21 | <i>CDKN2BAS1</i> | Coronary artery calcification | rs1333049 | 3.3×10^{-24} | + | 0.97 |
| | | Abdominal aortic aneurysm | rs2383207 | 1.9×10^{-8} | + | 0.91 |
| | | Ankle brachial index | rs10757269 | 2.7×10^{-9} | + | 0.9 |
| | | Stroke (large vessel stroke) | rs2383207 | 2.4×10^{-6} | + | 0.91 |
| | | Intracranial aneurysm | rs10733376 | 4×10^{-12} | + | 0.94 |
| 9q34 | <i>ABO</i> | Alkaline phosphatase in plasma | rs579459 | 3×10^{-123} | + | Same SNP |
| | | Activated partial thromboplastin time | rs579459 | 1.7×10^{-74} | + | Same SNP |
| | | Soluble P-selectin | rs579459 | 1.9×10^{-41} | + | Same SNP |
| | | Soluble E-selectin | rs579459 | 1.3×10^{-29} | + | Same SNP |
| | | Plasma carcinoembryonic levels | rs579459 | 3×10^{-21} | + | Same SNP |
| | | Red blood cell count | rs579459 | 9.3×10^{-18} | + | Same SNP |
| | | Hemoglobin | rs579459 | 1.4×10^{-15} | + | Same SNP |
| | | Hematocrit | rs579459 | 7.6×10^{-14} | + | Same SNP |
| | | Interleukin-6 levels | rs579459 | 3.6×10^{-13} | + | Same SNP |
| | | Circulating galectin-3 levels | rs579459 | 1.9×10^{-10} | + | Same SNP |
| | | Factor XIII antigen | rs579459 | 2.3×10^{-8} | + | Same SNP |
| | | von Willebrand factor | rs651007 | 1×10^{-161} | + | 1 |
| | | Venous thromboembolism | rs495828 | 2×10^{-17} | + | 1 |

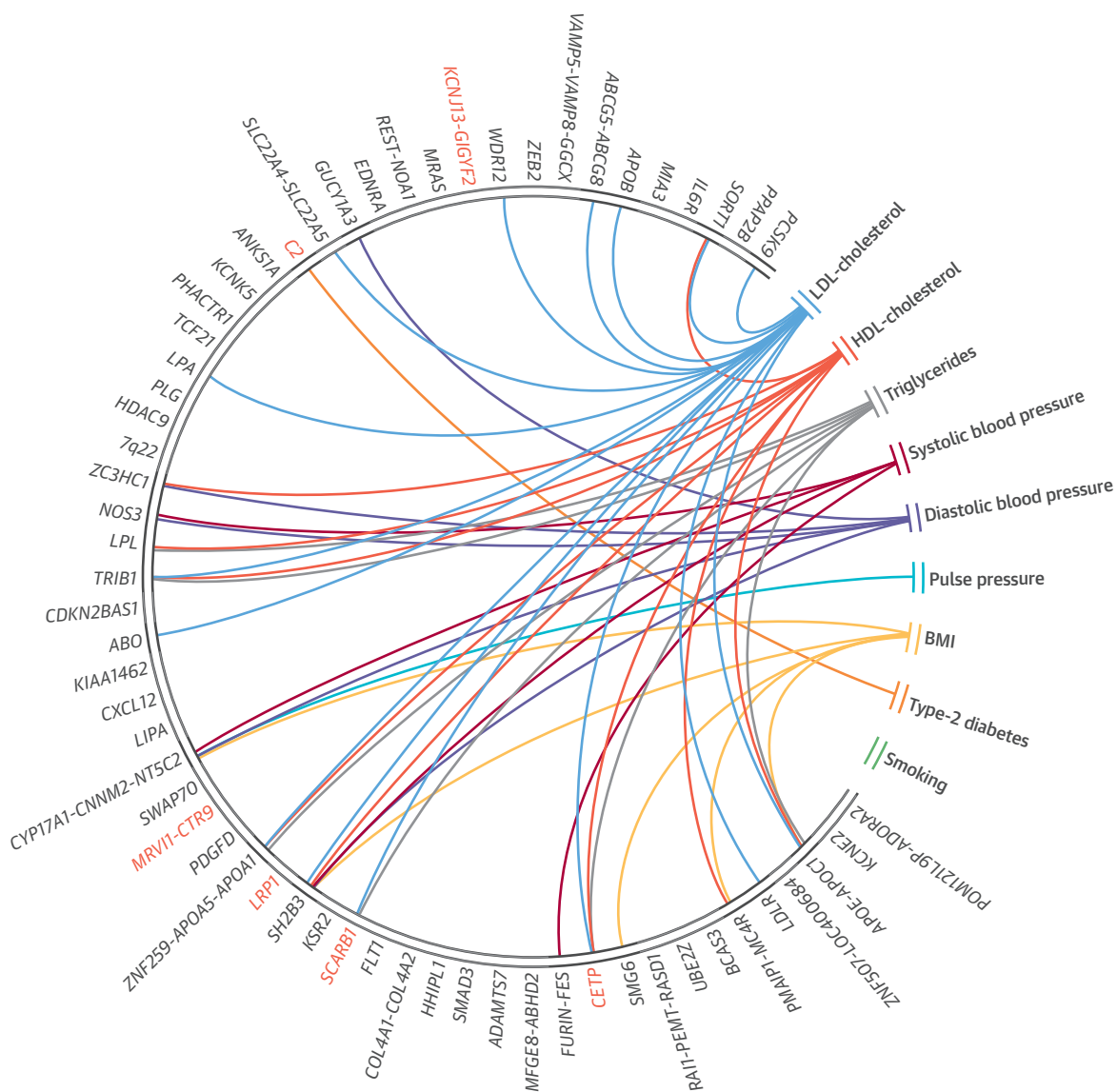
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TABLE 3 Continued

| Known Loci | | | | | | |
|------------|----------------------------|---|------------|-----------------------|---|----------|
| 10q24 | <i>CYP17A1-CNNM2-NT5C2</i> | Serum alkaline phosphatase levels | rs651007 | 1×10^{-56} | + | 1 |
| | | Ferritin levels | rs651007 | 1×10^{-8} | + | 1 |
| | | Soluble ICAM-1 | rs507666 | 3×10^{-91} | + | 0.83 |
| | | Intracranial aneurysm | rs12413409 | 1.2×10^{-9} | + | Same SNP |
| | | Schizophrenia | rs11191580 | 1.7×10^{-9} | + | 1 |
| 11q23 | <i>ZNF259-APOA5-APOA1</i> | Autism spectrum disorder | rs11191454 | 1.4×10^{-8} | + | 1 |
| | | Parkinson's disease | rs17115100 | 7.4×10^{-8} | + | 0.9 |
| | | Metabolic syndrome domains (Atherogenic dyslipidemia-PC2) | rs964184 | 1.8×10^{-12} | + | Same SNP |
| | | Vitamin E levels | rs964184 | 7.8×10^{-12} | + | Same SNP |
| 12q24 | <i>SH2B3</i> | Lp-PLA ₂ activity | rs964184 | 8.4×10^{-11} | + | Same SNP |
| | | Metabolic syndrome domains (Atherogenic dyslipidemia-PC1) | rs964184 | 1.2×10^{-10} | + | Same SNP |
| | | Selective immunoglobulin A deficiency | rs3184504 | 5.6×10^{-31} | + | Same SNP |
| | | Type 1 diabetes | rs3184504 | 2.8×10^{-27} | + | Same SNP |
| | | Celiac disease | rs3184504 | 5.4×10^{-21} | + | Same SNP |
| | | Hemoglobin | rs3184504 | 4.3×10^{-19} | + | Same SNP |
| | | Celiac disease | rs3184504 | 5.4×10^{-21} | + | Same SNP |
| | | Blood eosinophil count | rs3184504 | 6.5×10^{-19} | + | Same SNP |
| | | Generalized vitiligo | rs3184504 | 2.5×10^{-17} | + | 0.97 |
| | | Soluble ICAM-1 | rs3184504 | 2.9×10^{-17} | + | Same SNP |
| | | Hematocrit | rs3184504 | 7.9×10^{-16} | + | Same SNP |
| | | Hypothyroidism | rs3184504 | 2.6×10^{-12} | + | Same SNP |
| | | Rheumatoid arthritis and celiac disease | rs3184504 | 1.4×10^{-11} | + | Same SNP |
| | | Primary sclerosing cholangitis | rs3184504 | 5.9×10^{-11} | + | Same SNP |
| | | Serum urate | rs3184504 | 2.6×10^{-10} | + | Same SNP |
| | | Juvenile idiopathic arthritis | rs3184504 | 2.6×10^{-9} | + | 0.9 |
| | | Lymphocyte count | rs3184504 | 1.1×10^{-8} | + | Same SNP |
| | | Plasma beta-2 microglobulin levels | rs3184504 | 3.1×10^{-8} | + | Same SNP |
| | | White blood cell count | rs3184504 | 6.3×10^{-6} | + | Same SNP |
| | | Rheumatoid arthritis | rs3184504 | 6×10^{-6} | + | Same SNP |
| 13q34 | <i>COL4A1-COL4A2</i> | Tetralogy of Fallot | rs11065987 | 4.6×10^{-8} | + | 0.91 |
| | | Coronary artery calcification | rs3809346 | 8.6×10^{-7} | + | 0.97 |
| 15q22 | <i>SMAD3</i> | Crohn's disease | rs17293632 | 2.7×10^{-19} | - | 0.9 |
| | | Inflammatory bowel disease | rs17293632 | 6×10^{-16} | - | 0.9 |
| | | Ulcerative colitis | rs17293632 | 9.5×10^{-6} | - | 0.9 |
| 15q25 | <i>ADAMTS7</i> | Self-reported allergy | rs17228058 | 1.2×10^{-8} | - | 0.9 |
| | | Coronary artery calcification | rs3825807 | 6.5×10^{-6} | + | Same SNP |
| 17p13 | <i>SMG6</i> | Aortic root size | rs10852932 | 2.3×10^{-11} | + | 0.96 |
| 17q21 | <i>UBE2Z</i> | Height | rs318095 | 1.5×10^{-16} | - | 1 |
| | | Breast size (bra cup size in women) | rs12603969 | 3×10^{-5} | - | 1 |
| 18q21 | <i>PMAIP1-MC4R</i> | Obesity | rs17782313 | 4.8×10^{-15} | + | 0.86 |
| | | Height | rs11152213 | 6.9×10^{-13} | + | 0.91 |
| 19q13 | <i>APOE-APOC1</i> | Antipsychotic drug induced weight gain | rs12967878 | 3.6×10^{-7} | + | 0.9 |
| | | Common carotid artery intima-media thickness | rs445925 | 1.7×10^{-8} | + | Same SNP |
| | | Lp-PLA ₂ activity | rs445925 | 3.3×10^{-10} | + | Same SNP |
| | | Metabolic syndrome domains (Atherogenic dyslipidemia-PC1) | rs445925 | 1.3×10^{-35} | + | Same SNP |
| | | Alzheimer disease | rs2075650 | 1×10^{-295} | + | Same SNP |
| | | Longevity | rs2075650 | 3.4×10^{-17} | - | Same SNP |
| | | Lp-PLA ₂ activity | rs2075650 | 8.1×10^{-15} | + | Same SNP |
| | | Age-related macular degeneration | rs2075650 | 8.4×10^{-8} | - | Same SNP |
| | | C-reactive protein | rs2075650 | 4.2×10^{-8} | + | Same SNP |
| | | Cognitive decline | rs2075650 | 2×10^{-8} | + | Same SNP |

+ indicates increase in disease or trait with the CAD risk allele.

CAD = coronary artery disease; FEV1 = forced expiratory volume 1; FVC = forced vital capacity; ICAM-1 = intercellular adhesion molecule 1; Lp-PLA₂ = lipoprotein-associated phospholipase A₂; PC = principal component; SNP = single nucleotide polymorphism.

CENTRAL ILLUSTRATION Significant Associations of CAD Loci With Cardiovascular Risk Factors

Webb, T.R. et al. *J Am Coll Cardiol.* 2017;69(7):823-36.

This chord diagram depicts associations that passed Bonferroni correction (Table 2). Connections indicate that single nucleotide polymorphisms at respective loci associate with both coronary artery disease (CAD) and the respective risk factor; they do not imply that the risk factor causally explains the association with CAD. **Red** indicates new CAD loci. BMI = body mass index; HDL = high-density lipoprotein; LDL = low-density lipoprotein.

extended linkage at this locus, with more than 100 SNPs in high LD and the lead variant in a region of ~170 kb also spanning *GIGYF2* (GRB10 interacting GYF protein 2) (Online Figure 3). *KCNJ13* is located entirely within *GIGYF2* and transcribed in the opposite direction. A number of the associated variants are in annotated regulatory regions, with the top

scoring candidate by in silico prediction, rs11555646, lying in the 5'-UTR of *GIGYF2* close to the initiating methionine (Online Figure 3). There was no association of the locus with any of the cardiovascular risk factors, but we found eQTLs for the lead variant or a variant in high LD for both *GIGYF2* and *KCNJ13* (Online Table 5).

CAD LOCI AND PLEIOTROPY. We undertook an updated analysis of the association of all 62 CAD loci (56 published and 6 novel in this report) with traditional cardiovascular risk factors (blood pressure traits, lipid traits, BMI, type 2 diabetes, and smoking). The full results are shown in [Online Table 6](#), and the significant associations are summarized in [Table 2](#). Of the 62 CAD loci, 24 (38.7%) showed a statistical association at a Bonferroni corrected p value $< 8.32 \times 10^{-5}$ with a traditional cardiovascular risk factor with some loci showing multiple associations ([Central Illustration](#)). The largest number of associations were with lipid traits (14 with LDL cholesterol, 9 with HDL cholesterol, and 7 with triglycerides), followed by blood pressure traits (5 with diastolic blood pressure, 4 with systolic blood pressure, and 1 with pulse pressure), BMI (5 associations), and type 2 diabetes (1 association). Most associations were in the direction consistent from the epidemiological association of these risk factors with CAD, although a few displayed effects in the opposite direction (the risk variants at 2q33 and 12q24 are associated with reduced plasma LDL cholesterol, and those at 10q24, 12q24, and 19q13 are associated with lower BMI).

To inform the interpretation of these data, we conducted a complementary analysis for variants available on the array with a known genome-wide association with a risk factor; also, we compared the magnitude of the reported association with the risk factor to the observed association with CAD in our data. Except for LDL cholesterol and BMI, the correlations between the 2 effects were either weak or insignificant ([Online Figure 4](#)). In a separate analysis conducted in the 150,000 participants in UK Biobank with currently released genotype data, we confirmed that none of the CAD-associated variants showed a sex difference in allele frequency (data not shown).

We next analyzed the association of the 62 CAD loci with other diseases and traits. When restricted to variants with a high LD ($r^2 > 0.8$) with the lead CAD variant, 29 of 62 (47%) loci showed an association with another disease/trait at a p value $< 1 \times 10^{-4}$. Several loci showed multiple associations ([Table 3](#)). Although in most cases, the CAD-associated risk allele was also associated with an increased risk (or level) of the other disease or trait, this was not always the case. Furthermore, in some loci with multiple associations, the direction of association varied between diseases ([Table 3](#)).

DISCUSSION

This large-scale meta-analysis of common variants, including many with prior evidence for association

with another complex trait, resulted in the identification of 6 new CAD loci at genome-wide significance. We also showed that almost one-half of the CAD loci that have been identified to date demonstrate pleiotropy, an association with another disease or trait. The findings added to our understanding of the genetic basis of CAD and might provide clues to the mechanisms by which such loci affect CAD risk.

Our findings of a genome-wide association with CAD of a functional variant in the promoter of the *CETP* gene that is also associated with its expression and plasma activity (18-20) have added to previous evidence linking genetically determined increased activity of this gene with higher risk of CAD (20). There has been a longstanding interest in CETP inhibition as a therapeutic target, primarily because of the effect on plasma HDL cholesterol level. However, several CETP inhibitors have recently failed to improve cardiovascular outcomes in large randomized clinical trials (28-30) and, in 1 case, caused harm (28), despite markedly increasing plasma HDL cholesterol. Furthermore, Mendelian randomization studies have questioned the causal role of lower plasma HDL cholesterol in increasing CAD risk (31,32). Although previous studies have shown that the CETP genetic variant we report here affects CETP activity, the precise mechanism(s) by which this variant modifies CAD risk remains uncertain.

A notable finding was the association with CAD of common variants located in the *SCARB1* gene. Association of variants at the *SCARB1* locus with CAD was also reported by the CARDIoGRAMplusC4D consortium, but this did not reach genome-wide significance (1). The gene encodes the canonical receptor, SR-BI, responsible for HDL cholesteryl ester uptake in hepatocytes and steroidogenic cells (33). Genetic modulation of SR-BI levels in mice is associated with marked changes in plasma HDL cholesterol (34). Consistent with this, a rare loss of function variant in which leucine replaces proline 376 (P376L) in *SCARB1* was recently identified through sequencing of individuals with high plasma HDL cholesterol (35). Interestingly, despite having higher plasma HDL, 346L carriers had an increased risk of CAD, suggesting that the association of variation at this locus on CAD is not driven primarily through plasma HDL (35). Indeed, there is only a nominal association of the lead CAD variant at this locus (rs11057830) with plasma HDL cholesterol ([Online Table 6](#)). The variant is also modestly associated with plasma LDL cholesterol and serum triglycerides ([Table 2](#)). All 3 of these lipid associations are directionally consistent with epidemiological evidence linking them to CAD risk and could,

in combination, explain the association of the locus with CAD. However, the lead variant is more strongly associated with Lp-PLA₂ activity and mass (Table 3), which could provide an alternative explanation for its association with CAD. Irrespective of the mechanism, our findings, when combined with those of Zanoni et al. (35), suggest that modulating SR-B1 may be therapeutically beneficial.

After adjusting for multiple testing, we found that slightly more than one-third of the CAD loci showed an association with traditional cardiovascular risk factors. Although the vast majority of associations were in the direction consistent with the epidemiological association of these risk factors with CAD, as noted in the previous text with respect to loci affecting the HDL cholesterol level, this should not be interpreted as implying that these loci affect CAD risk through an effect on the specific risk factor. Indeed, for variants available on the array with a known genome-wide association with these risk factors, we found a poor correlation between the magnitudes of their effect of the risk factor and their association with CAD in our dataset except for LDL cholesterol (Online Figure 4). Nonetheless, formal causal inference analyses, using Mendelian randomization, have implicated LDL cholesterol, triglyceride-rich lipoproteins, blood pressure, type 2 diabetes, and BMI as causally involved in CAD (36).

Almost one-half of the CAD loci showed a strong or suggestive association with other diseases or traits with, in many cases, the identical variant being the lead variant reported for the association with these other conditions (Table 3). Some of the associations with other traits—for example, coronary calcification (3q22, 6p24, 9p21, 13q34, and 15q25) or carotid intima-media thickness (4q31 and 19q13)—are not surprising, as these traits are known to be correlated with CAD. Others, such as risk of stroke (7p21 and 9p21), might reflect a shared etiology. However, the mechanism(s) behind most of the observed pleiotropy is not clear, although the findings could provide clues as to how the locus may affect CAD risk. As an example, 5 loci (12q24, 1p13, 6q25, 11q23, and 19q13) show strong associations with plasma activity and/or mass of Lp-PLA₂. Lp-PLA₂ is expressed in atherosclerotic plaques where studies have suggested a role in the production of proinflammatory and proapoptotic mediators, primarily through interaction with oxidized LDL (37,38). A meta-analysis of prospective studies showed an independent and continuous relationship of plasma Lp-PLA₂ with CAD risk (39). However, it should be noted that Mendelian randomization analyses have not supported a causal

role of secreted Lp-PLA₂ in coronary heart disease (40), and phase III trials of darapladib, an Lp-PLA₂ inhibitor, have shown no benefit in patients with stable coronary heart disease (41) or acute coronary syndromes (42) when added to conventional treatments including statins.

Chronic inflammation plays a key role in both the pathogenesis of CAD and of inflammatory bowel disease. It is therefore interesting to note the association of the same locus at 15q22 with CAD as well as Crohn's disease and ulcerative colitis (Table 3). Association of this locus with CAD at genome-wide significance was recently reported by the CARDIOGRAMplusC4D consortium (2) with the lead SNP (rs56062135) showing strong linkage disequilibrium ($r^2 = 0.9$) with the lead SNP (rs17293632) associated with inflammatory bowel disease. Both rs56062135 and rs17293632 lie in a region of ~30 kb within the initial introns of the SMAD family member 3 gene (*SMAD3*), a signal transducer in the transforming growth factor-beta pathway. Indeed, rs17293632 was included on the exome array because of its known association with Crohn's disease and showed a significant association with CAD in our combined dataset ($p = 1.78 \times 10^{-8}$). Farh et al. (43) interrogated ChIP-seq data from ENCODE and found allele-specific binding of the AP-1 transcription factor to the major (C) allele in heterozygous cell lines and suggested that the T allele of rs17293632 increases risk of Crohn's disease by disrupting AP-1 regulation of *SMAD3* expression. Interestingly, the direction of effect on CAD risk observed for this variant was in the opposite direction to that for inflammatory disorders, with the C allele being the risk allele. Recent analysis of this variant in arterial smooth muscle cells confirmed that the CAD risk allele preserves AP-1 transcription factor binding and increases expression of *SMAD3* (44). Further investigation of the discordant effects of *SMAD3* may shed light on the mechanisms of both diseases.

STUDY LIMITATIONS. First, in our discovery study, we were only able to interrogate common variants associated with other diseases and traits that were known at the time of the creation of the exome array in late 2011 and, thus, included on the array. Conversely, our interrogation for pleiotropic associations of the new and known CAD has used the latest data available in the GWAS catalogs and other sources. Second, the common variants tested in our study conferred statistically robust yet quantitatively modest effects on both CAD and potentially related traits. Thus, we may have missed associations with other traits. However, if such traits were considered as intermediary steps in the etiology of CAD,

exploration of our large GWAS sample sets and respective GWAS catalogs should have detected relevant associations. Third, our discovery analysis is largely on the basis of subjects with Western-European ancestry, and any association with CAD of the new loci in other populations needs further evaluation. Finally, although we used relatively stringent criteria (minimal $r^2 > 0.8$ between the CAD SNP and the lead variant associated with the other disease/trait), the limited content of the exome array and the information available in the GWAS catalogs meant that we could not examine the extent of overlap in the loci in detail.

CONCLUSIONS

Through an analysis of selected variants associated with other disease traits, we reported the discovery of 6 further loci associated with CAD. Furthermore, in the most comprehensive analysis to date, we showed that several of the new and previously established loci demonstrated substantial pleiotropy, which may help our understanding of the mechanisms by which these loci affect CAD risk.

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PERSPECTIVES

COMPETENCY IN MEDICAL KNOWLEDGE: Novel genetic loci influence risk of coronary artery disease, but only one-third are associated with conventional cardiovascular risk factors, whereas at least one-half of the loci are associated with other diseases or traits (pleiotropy).

TRANSLATIONAL OUTLOOK: Future studies should investigate the mechanisms that relate the observed pleiotropy to the pathogenesis of atherosclerosis and ischemic events.

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APPENDIX For supplemental tables and figures as well as a full list of acknowledgments and funding sources, please see the online version of this article.