Methylation at CpG sites related to growth differentiation factor-15 was not prospectively associated with cardiovascular death in discordant monozygotic twins

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Myocardial infarction patients had decreased methylation at four growth differentiating factor-15 (GDF-15) related CpG sites (cg13033858, cg16936953, cg17150809, and cg18608055). These sites had not been studied for their association with cardiovascular disease (CVD) deaths. Thus, we aimed to assess the associations independent of genes, shared environment, and traditional CVD risk factors. Nineteen white, male, monozygotic twin pairs discordant for CVD deaths were included from the National Heart, Lung and Blood Institute Twin Study (NHLBI) initiated in 1969. Data on vital status was collected through December 31, 2014. Methylation of buffy coat DNA at exam 3 (1986–87) was measured using the Illumina HumanMethylation450 BeadChip. Principal component analysis was used to generate a score representing blood leukocyte composition and baseline CVD risk factors and predominated with natural killer cells, CD4+ T cells, and Framingham risk score. Conditional logistic regression demonstrated that methylation at the four CpG sites was not associated with CVD deaths before (all p > 0.05, bootstrapped p > 0.05) and after adjustment for the score (all p > 0.05). Joint influences of cg16936953 and the score were statistically significant (p < 0.05). In conclusion, joint influences of methylation at the site cg16936953 and the score are prospectively associated with CVD deaths independent of germline and common environment.

ClinicalTrials.gov Identifier for NHLBI Twin Study: NCT00005124.

Abbreviations

NHLBI  National Heart, Lung, Blood Institute
CVD  Cardiovascular disease
MZ  Monozygotic twins
GDF-15  Growth differentiating factor-15
HR  Hazard ratio
CI  Confidence interval

Cardiovascular disease (CVD) is the leading cause of death worldwide, with most associated deaths resulting from atherosclerotic CVD such as heart attacks or heart failure in the U.S.1. Increasing our understanding of the contributions of gene-environment interactions to the risk of death from CVD remains of high importance to the treatment and prevention of this disease. Epigenetic changes constitute molecular modifications to the

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Epigenetic changes to the genome are stable across mitotic divisions. They involve chemical modifications to DNA or associated proteins, such as the methylation of 5-cytosine in CpG (cytosine-phosphate-guanine) DNA dinucleotide pairs in the genome via the transfer of a methyl group to the pyrimidine ring of the cytosine. Methylation of promoter sequences is associated with repression of expression of the associated gene(s) and establishment of longer-term gene silencing. While most variation in DNA methylation patterns between individuals is due to environmental factors, genetic influences on DNA methylation are high at certain genomic sites, especially those exhibiting high levels of variability in methylation.

In a genome-wide methylation association study, after adjusting for white blood cell composition, methylation levels at four CpG sites (cg17150809, cg13033858, cg16936953, and cg18608055) were associated with circulating GDF-15 levels in discovery and validation populations and were potentially associated with prior myocardial infarction history. Differential DNA methylation has been associated with expression levels of growth differentiation factor-15 (GDF-15), a biomarker for CVD risk. GDF-15 is a member of the transforming growth factor-β family involved in regulating inflammatory processes. A recent meta-analysis of 31 prospective studies demonstrated that higher levels of circulating GDF-15 were associated with a higher risk of cardiovascular mortality [hazard ratio (HR) 2.11; 95% confidence interval (CI) 1.57–2.66 per log-unit ng/L increment] among 53,706 individuals who had an average age of 60–75 years.

Although the associations of methylation at the four GDF-15 related CpG sites with the prior myocardial infarction history became statistically insignificant after multiple-testing adjustment, the sites are of interest for further studies because potential residual confounding, such as potential genetic factors, could mask the association. Epigenetic association studies for CVD in the general population are potentially confounded by genetic factors, as genetic variation impacts human DNA methylation patterns and CVD. Prior studies on the influence of human DNA methylation at these GDF-15 related CpG sites on CVD hard outcomes are rare. Importantly, previous few studies demonstrate inconsistent findings of methylation at the GDF-15 related CpG sites in relation to CVD risk factors among humans, suggesting potential genetic confounding. For example, a prior study of blood levels of C-reactive protein (CRP) as a biomarker for chronic low-grade inflammation towards CVD reported inconsistent directions of association of the site cg16936953 but not cg18608055 with serum CRP between individuals of European ancestry and those of African Americans. Inconsistent associations were also reported between methylation at the CpG site cg18608055 and obesity as a CVD risk factor. A negative association was found among middle-aged African Americans from the Atherosclerosis Risk in Communities (ARIC) study and among young Norwegian women; however, the negative association was unable to be replicated in Finnish monozygotic twins discordant for body mass index, implying that the observed association might be due to potential confounding from germline and shared environment.

Discordant monozygotic (MZ) twin pair studies, in which one member of the twin pair (i.e., one co-twin) is affected by a disease and the other is not, provide an optimal, natural experiment to control for genetic confounding because of the identical genomic sequence between co-twins of a monozygotic twin pair. In addition, twin studies also control for age and shared environmental factors, such as the prenatal environment, and thus can provide significant power to detect associations between epigenetic patterns and complex diseases. This study used a discordant MZ twin design to include MZ twins discordant for CVD death from the 41-year follow-up National Heart, Lung, and Blood Institute (NHLBI) Twin Study. We aimed to assess whether methylation at the four genomic CpG sites linked to circulating GDF-15 levels was prospectively associated with CVD death risk independent of germline, shared environmental factors, blood leukocyte composition, and known traditional CVD risk factors.

Results

Characteristics of study participants. All 19 control co-twins outlived their case co-twin brothers in this study, of which 18 control co-twins died before the date of the end of follow up (i.e., December 31, 2014) and one control co-twin was still alive on December 31, 2014. The diseased twins died significantly earlier than their co-twin brothers (74.7 versus 83.7 years old at death on average) with the within-pair differences (i.e., pair-wise differences) in age at death or the last follow-up date (years) ranging from 3.5 to 23.8 years [mean ± SD 10.3 ± 5.7 years; median (IQR) 10.0 (9.3) years] for control twins relative to their diseased identical twin brothers (Table 1 and Supplementary Fig. S1). Diseased twins had significantly higher systolic blood pressure (p = 0.03) and lower levels of low-density lipoprotein cholesterol (p = 0.008) at baseline than their control twin brothers.

Associations of methylation at the four CpG sites with risk for CVD death. Table 2 shows that methylation levels at the 4 CpG sites, cg13033858, cg16936953, cg17150809, and cg18608055, were not statistically significantly associated with the risk of death from total CVD (all p > 0.05) before and after controlling for the principal component score as a surrogate score with wide 95% CIs. However, the joint influence of cg16936953 and the score was statistically significantly associated with CVD death (p < 0.05), and the adjusted hazard ratio (HR) was marginally statistically significant for cg16936953 [HR 4.38, 95% confidence intervals (CI) 0.79–24.3, p = 0.09] and the score (p = 0.06), respectively. The information gain from the joint influence of CpG sites and the surrogate score above 10% was 13% and 10% for cg16936953 and cg17150809, respectively, suggesting the biological, clinical, or scientific importance of the joint influences associated with CVD death risk (Table 2). The varimax-rotated weight for the score was −0.78, 0.67, 0.65, −0.40, −0.25, 0.05, −0.097, and 0.06 for natural killer cells, CD4+ T cells, Framingham risk score, CD8+ T cells, body mass index, B cells, monocytes,
and years of education, respectively, suggesting the biological and clinical importance of natural killer cells, CD4+ T cells, and Framingham risk score. Statistical insignificant results were found among stringently defined twin pairs discordant for CVD, primarily and stringently defined twin pairs discordant for coronary heart death (Supplementary Table S1).

**Bootstrap analysis.** Methylation levels at the 4 CpG sites were not statistically significantly associated with the risk of death from total CVD (all bootstrapped p > 0.05) without controlling for the surrogate score in the bootstrap analysis of 10,000 bootstrapped samples. The bootstrapped 95% CI was (0.37–1.47), (0.82–1.81), (0.55–3.58), and (0.57–2.01) after natural adjustment for germline and shared environment for sites cg13033858, cg16936953, cg17150809, and cg18608055, respectively.

**Discussion**

After natural adjustment for germline and common environment, methylation levels at the 4 CpG sites, cg13033858, cg16936953, cg17150809, and cg180608055, were not statistically significantly associated with death from total CVD before and after additional controlling for potential confounding from blood leukocyte composition and traditional known CVD risk factors in the original sample as well as after natural controlling for germline and shared environment only in 10,000 bootstrapped samples. Unadjusted bootstrapped 95% CIs indicated the precision of estimates at sites cg17150809 and cg18608055 were very poor compared to that at sites cg13033858 and cg16936953, implying the sites cg13033858 and cg16936953 might be more clinically useful, particularly the site cg16936953 with the narrowest 95% CI among the 4 CpG sites. From a clinical view, influences of 10% increment in methylation β values on the CVD survival ranged from 18% protective to 81%
Controversial cardiovascular functions might depend on specific cell subtypes and the pathophysiological stages in blood monocytes might contribute to atherosclerosis. As reviewed by Kapnick et al., the reduction in FBXL4 miRNA-21 biogenesis by methylation at the site cg16936953 could be detrimental to cardiac function recovery.

DNA were not associated with incident acute coronary syndrome among Chinese. We did not find previously published research on methylation at sites cg17150809, cg13033858, or cg16936953 related to other hard CVD outcomes in humans. Therefore, our study provides unique evidence on cardiovascular death as a CVD hard outcome independent of germline, common environment, blood leukocyte composition, and traditional CVD risk factors, in particular, evidence of the potential joint influence of the CpG sites and the overall effect of blood leukocyte subtypes. *p < 0.05 likelihood ratio test for the adjusted model in which the score had an adjusted p-value of 0.06.

Table 2. Hazard ratio per 10% unit increase in methylation beta value derived from the regression coefficient from a conditional logistic model among monozygotic twins discordant for total cardiovascular death. CI confidence interval. Adjusted hazard ratios were obtained after control for a principal component score representing age, years of education, smoking status, body mass index, systolic blood pressure, low-density lipoprotein cholesterol, high-density lipoprotein cholesterol, use of antihypertensives, presence of diabetes, and leukocyte subtypes. *p < 0.05 likelihood ratio test for the adjusted model in which the score had an adjusted p-value of 0.06.

| CpG site | Hazard ratio (95% CI) | p value | Information gain |
|----------|----------------------|---------|------------------|
| cg13033858 |                      |         |                  |
| Crude     | 0.64 (0.18, 2.22)    | 0.48    | 1.4%             |
| Adjusted  | 1.23 (0.25, 6.04)    | 0.80    | 5.3%             |
| cg16936953 |                      |         |                  |
| Crude     | 2.14 (0.64, 7.22)    | 0.22    | 4.7%             |
| Adjusted* | 4.38 (0.79, 24.3)    | 0.09    | 12.8%            |
| cg17150809 |                      |         |                  |
| Crude     | 1.60 (0.35, 7.21)    | 0.54    | 1.0%             |
| Adjusted  | 3.48 (0.46, 26.5)    | 0.23    | 10.3%            |
| cg18608055 |                      |         |                  |
| Crude     | 1.08 (0.28, 4.07)    | 0.91    | 0.03%            |
| Adjusted  | 2.28 (0.42,12.3)     | 0.34    | 9.1%             |

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Hazard ratio per 10% unit increase in methylation beta value derived from the regression coefficient from a conditional logistic model among monozygotic twins discordant for total cardiovascular death. CI confidence interval. Adjusted hazard ratios were obtained after control for a principal component score representing age, years of education, smoking status, body mass index, systolic blood pressure, low-density lipoprotein cholesterol, high-density lipoprotein cholesterol, use of antihypertensives, presence of diabetes, and leukocyte subtypes. *p < 0.05 likelihood ratio test for the adjusted model in which the score had an adjusted p-value of 0.06.

The site cg13033858 is located on chromosome 12 within the gene body coding for protein phosphatase slingshot homolog-1 (SSH1) and related to the CpG shelf, 2 to 4 kb from the upstream of the CpG island. SSH1 may reduce fibrosis by suppressing angiotensin II-induced remodeling in the vasculature through regulating actin filament dynamics and SSH1 activity. Methylation at the site cg13033858 would thus enhance fibrosis by downregulating SSH1 biosynthesis, leading to heart failure.

The site cg17150809 is located on chromosome 6 and upstream of the gene coding for F-Box and Leucine Rich Repeat Protein 4 (FBXL4). FBXL4 protein suppresses mitochondrial degradation through its function related to E3 ubiquitin ligase complex activity. Bioinformatics analysis revealed that the FBXL4 gene in peripheral blood monocytes might contribute to atherosclerosis. As reviewed by Kapnick et al., the reduction in FBXL4 protein contributed to immune dysfunction, including neutropenia, lymphopenia, and frequent infections, and reductions in natural killer cells, total CD8+ T cells, and CD8+ memory T cells. Our findings suggested the potential biological and clinical important role of natural killer cells and CD4+ T cells, to a lesser extent CD8+ T cells, along with this site cg17150809 in cardiovascular death contributing to the pathophysiological process.

Sites cg16936953 and cg18608055 appear to have bi-directional roles in CVD. First, the site cg16936953 was located on chromosome 17 within the Vacuole membrane protein 1 (VMP1)/Micro-RNA 21 (MIR21)/promoter to regulate miRNA-21 biogenesis. MiRNA-21 plays crucial but controversial roles in CVD. Its controversial cardiovascular functions might depend on specific cell subtypes and the pathophysiological stages in cardiovascular impairment. MiRNA-21 could protect cardiac function by preventing excessive inflammation and cardiac dysfunction after myocardial infarction through targeting KBTBD7. Thus, downregulation of miRNA-21 biogenesis by methylation at the site cg16936953 could be detrimental to cardiac function recovery.
On the other hand, miRNA-21 could promote the progress of cardiac hypertrophy to heart failure by mediating cardiac fibrosis. Therefore, downregulation of miRNA-21 biogenesis by methylthion at this site could prevent the hypertrophic heart from heart failure. Our findings suggested the potential statistical and clinical importance of the surrogates score, particularly natural killer cells, CD4+ T cells, and Framingham risk score, in their joint effect with the site cg16936953 on the cardiovascular death contributing to the pathophysiological process.

Second, the CpG site cg18608055 is located on chromosome 19 within the body of the gene “Strawberry notch homologue 2 (SBNO2).” Evidence from laboratory experiments suggested that SBNO2 was involved in both inflammation and anti-inflammation. SBNO2 was an acute inflammatory response gene upregulated by IL-6, while SBNO2 protein had strong repressive activity for NF-κB and was a component of the pathway regulating the downstream anti-inflammatory effects of interleukin-10.

Overall, the pathophysiological mechanisms are putative as methylation at the four sites could regulate the biogenesis of factors in the regulatory network contributing to atherosclerosis and fibrosis. Thus, more laboratory research is needed to elucidate pathophysiological mechanisms linking methylation at these sites to CVD disease.

There are limitations to this study. Our discordant MZ twin sample size seems small. As described in our previous study, the twin birth was less than 1.5 in 100 births during 1917 to 1927 when twins in the NHLBI Twin Study were born. Further considering male MZ twin live birth rate and live births during the same period in the US, we estimated that the 254 MZ pairs initially enrolled in the NHLBI Twin Study were comparable to 110,535 male singletons. Since the Framingham Heart Study of cardiovascular disease enrolled 2344 men in its initial cohort, our sample size was large when we took twinning birth rate and male monozygotic twin live birth rate into consideration. The extremely long follow-up in the NHLBI Twin Study provided a unique opportunity to prospectively study cardiovascular death in the large discordant twin cohort. However, we cannot exclude the potential statistical insignificance because of insufficient statistical power resulting from the small sample size. Our results could be used to estimate the sample size for future large-scale studies. Second, we used the first principal analysis component as a surrogate for the overall effect of blood leukocytes and known CVD risk factors to minimize overfitting. The first component explained less than 70% of the variation in blood leukocytes and known CVD risk factors. Although we could not exclude residual confounding, the most parsimonious model was the first principal component adjusted model for the four CpG sites (Supplementary Table S2). We did not measure messenger RNA (mRNA) to evaluate GDF-15 expression since the technique to preserve mRNA was not used in the mid-1980s when biospecimens were collected in the NHLBI Twin Study. We were unable to evaluate the between-pair effects because of the use of the discordant twin pair design and the resultant conditional logistic regression analysis.

Our study has several advantages. The use of MZ discordant twin pairs allowed for an assessment of associations in the absence of genetic confounding, as MZ twin pairs are the same germline. In addition to adjustment for age-cohort-period influences in our discordant twin design, we controlled for traditional known CVD risk factors to a certain degree, including years of education representing socioeconomic status, body mass index, age, systolic blood pressure, low-density lipoprotein cholesterol, high-density lipoprotein cholesterol, current cigarette smoking status, use of antihypertensives, and the presence of type 2 diabetes. We provided the unadjusted 95% CIs from 10,000 bootstrap resamples as an alternative to minimize potentially biased results because of the relatively small sample size. To further reduce the bias, a larger sample of discordant identical twin pairs would be needed. Our extended longitudinal design demonstrated that methylation temporally occurred before cardiovascular death. This temporal order was critical to explain the causal role of methylation at the studied 4 CpG sites in cardiovascular death.

In conclusion, the joint influences of methylation at cg16936953 and the overall effect of blood leukocyte composition (particularly, natural killer cells and CD4+ T cells) and known CVD risk factors (particularly, Framingham risk score) are prospectively associated with the risk of death from total CVD, independent of genetic and shared environmental factors.

Methods

Study population. The NHLBI Twin Study was a longitudinal cohort study designed to study genetic and environmental causes of cardiovascular disease in the United States. The original cohort was initiated in 1969 and included 514 middle-aged white male veteran twin pairs (1028 men, 254 monozygotic and 260 dizygotic twin pairs) from the National Academy of Sciences-National Research Council Veteran Twin registry who lived within 200 miles of five research centers: Framingham, Massachusetts; San Francisco, California; Davis, California; Los Angeles, California; and Indianapolis, Indiana. Members of the cohort were born between 1917 and 1927 and were 42–55 years of age at the baseline examination (Exam 1, 1969–1973). The baseline and subsequent examinations were conducted using the reputable Framingham Heart Study protocol to ensure uniform physical examination by experienced cardiovascular epidemiologists. Five active follow-up studies were performed during 1981–1982, 1986–1987, 1995–1997, 1999–2001, and 2001–2003. Both biochemical and questionnaire data were collected at baseline and exams 2 and 3. The last three examinations (exams 4–6) collected structural brain MRI and questionnaire data. A total of 792, 622, 395, 438, and 174 twins participated in exams 2–6, respectively. Zygosity was identified in the 1960s using eight red blood cell antigen groups and in the 1980s using the variable number of tandem repeat DNA markers.

This reported study used a discordant monozygotic twin pair design that included identical twin pairs discordant for CVD death. As described in our previous study, this approach was a specific type of the 1:1 individually matched, nested case–control design, in which one co-twin within a twin pair was the case and his co-twin brother was the control. Thus, this prospective design could be called a nested case-co-twin-control design. In the nested case–control design, a control had not developed the disease by the time of disease occurrence in the case (index date) and might later become a case. The advantage of this definition was to make inference
under a proportional hazards model from the conditional logistic approach. Thus, this design could provide unbiased relative risk estimates. We used the following definitions to identify twin pairs discordant for CVD. The primary definition of a twin pair discordant for total CVD death was one where a co-twin died from CVD (case co-twin) and his co-twin brother who did not die from CVD on the index date or died from CVD at least 1 year later (control co-twin). A more stringent definition was one where a co-twin died from CVD (case) and his co-twin brother did not die from CVD by the end of the follow-up (control). Inclusion criteria included: (1) availableuffy coat DNA ≥ 400 µg on the DNA inventory of Veteran Twin Samples after exam 5 for each co-twin of a twin pair, (2) available data on vital status, dates of death, and causes of death through December 31, 2010, and (3) methylation and hydroxymethylation were measured. We performedzygosity-specific stratified random sampling by twins discordant for coronary heart death and non-coronary heart CVD death. Vital data were updated in 2014. The primarily-defined discordant twin pairs using 2010 and 2014 data were identical. Thus, we finally included 19 monozygotic twin pairs (CVD-dMZ) discordant for CVD death. This study was approved by the Institutional Review Board (IRB) of Vanderbilt University as a non-human subject study due to the use of de-identified information (IRB #141163, July 25, 2014), and informed consent was also waived by this IRB. In addition, the approval from the IRB of Des Moines University was waived due to the prior approval at Vanderbilt University. All methods were performed in accordance with the relevant guidelines and regulations.

DNA sample collection. As previously published, buffy coat DNA samples were collected at exam 3 (1986–1987) in the NHLBI Twin Study. Whole blood was drawn from the antecubital vein into EDTA tubes after an overnight fast and immediately placed on ice. Buffy coats were obtained and used to extract DNA. Spectral analysis was used to determine the quantity and quality of the extracted DNA. DNA samples were stored at −70 °C. All samples were labeled by the study number only.

Genome-wide methylation (5mC) measures. Genome-wide methylation was measured with the Illumina Infinium HumanMethylation450 (450 K) BeadChip array following the established Illumina protocol. Co-twin samples were processed in the same analytical run without knowing disease status to minimize measurement error. For quality control, beta values with associated detection p-values greater than 0.01 were set to missing. All samples passed quality control (missing percentage < 1.5%). CpGs with greater than 10% missing data were removed (452 CpGs removed).

Other known cardiovascular risk factors at baseline. All major cardiovascular risk factors were recorded on a questionnaire through in-person interviewing and physical examination at baseline (exam 1, 1969–1973). Participants were assessed on demographic, lifestyle, familial, socioeconomic, biochemical, and clinical factors. Age, years of education, marital status, and smoking status were recorded, and physical exam measurements included height, weight, diastolic and systolic blood pressures, and a 12-lead electrocardiogram. Lab measurements included 9-h fasting glucose levels and a glucose load tolerance test. Low-density lipoprotein cholesterol levels were calculated using the Friedewald equation. Physicians assessed the history of heart and other cardiovascular diseases and cardiovascular events and procedures.

Follow-up and assessment of endpoints. Vital status and the cause and date of death were ascertained through medical records in the five active follow-up examinations and later using death certificates or the National Death Index through December 31, 2014. As previously described, physicians assigned corresponding International Classification of Diseases, Ninth Revision codes (ICD-9) for morbidity outcomes. Death certificates or the National Death Index coded to the ninth revision codes were obtained for decedents. The endpoint was death from all cardiovascular diseases (390–398, 402, 404, 410–438). Subjects were considered lost to follow-up if a death certificate or coding from the National Death Index could not be traced. The follow-up was terminated at the date of death, end of follow-up, or loss to follow-up, whichever occurred first.

Statistical analysis. The post-normalized beta values were calculated using the 450 K array methylation data following the published method. Briefly, data was batch normalized using the Combat function in R using a set of 12 samples on a single array as a batch. To carry out the normalization, parallel processing was used on random subsets of 20,000 CpGs. Probes of different chemistries (Infinium I or Infinium II) were normalized separately. After normalization, a chemistry correction was applied. This correction was based on applying a second-order polynomial fit using CpGs of different chemistries within 50 bp of each other (correlations > 0.99 for pairs of CpGs within this distance). The missing value for a co-twin was replaced with his co-twin brother's value. Only one participant had missing values for lipid profile and Framingham risk score, while another participant had a missing value for the beta value at cg17150809.

Principal component analysis with varimax rotation was performed on blood leukocytes and known CVD risk factors. Five of the six types of blood leukocytes (i.e., CD4+ T-cells, CD8+ T-cells, natural killer (NK) cells, B-cells, monocytes, and granulocytes), except for granulocytes, were selected a priori as the sum of the composition of six types of blood leukocyte composition was equal to 100%. CVD risk factors included years of education (continuous), body mass index (continuous) and the calculated Framingham risk score, which incorporated factors including age, systolic blood pressure (continuous), total cholesterol (continuous), high-density lipoprotein cholesterol (continuous), use of blood pressure medication (yes/no), cigarette smoker (yes/no), and presence of diabetes (yes/no). To avoid overfitting, we used the first principal component score as the surrogate measure to reflect the potential confounding from the leukocyte and CVD risk factors. We used conditional logistic regression to calculate hazard ratios for evaluating the association of the continuous methylation level in β values with CVD death. We first evaluated the association without adjustment.
for any covariates. Then, we adjusted for the first principal component score. The information gain from adding predictors such as CpG site and the surrogate score into the model was calculated as the difference between the negative of twice the log likelihood (−2LogL) from a model without predictors and −2LogL from the model with the predictors (CpG in the unadjusted model or CpG and the score in the adjusted model) divided by twice the number of twin pairs, and then multiplied by 100. This sample size adjusted information gain reflects the scientific and practical importance of the predictor.

Lastly, a bootstrap resampling analysis was performed as previously described. Bootstrapping randomly sampled individual twin pairs to create a new data set of equal size to the original dataset and had several statistical advantages. The simulated samples came from the actual data, making no assumptions about the parent population parameters. This method also allowed for bypassing the large sample size assumption needed for most other statistical analyses. We generated 10,000 resampled data sets using the nonparametric bootstrap method. Then, unadjusted models were repeatedly fitted to each bootstrapped data set. The 95% confidence intervals (95% CIs) for hazard ratios were computed for each CpG site from the 10,000 bootstrap samples.

R and SAS version 9.4 statistical software was used for the analysis. Blood leukocyte composition was estimated for six blood leukocyte types from the 450 K array methylation data using the “estimateCellCounts” function in the R package “minfi”. The “clogit” and “prcomp” functions were used for conditional logistic regression and principal component analysis, respectively. Bootstrap analysis was performed using the R package “bootstrap.” We also performed the sensitivity analysis among a subset of 13 stringently defined CVD-dMZ pairs and another subset of 13 MZ pairs primarily defined discordant for coronary heart death out of the primarily defined 19 CVD-dMZ pairs. Both subsets had an overlap of the 8 MZ pairs. As results were materially similar to those from primarily defined pairs, we reported the results from the primarily-defined twin pairs discordant for CVD here and sensitivity analysis results in the supplementary file. The statistical significance level was set at the alpha level of 0.05.

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**Author contributions**

J.D. designed, directed and oversaw the project and maintained the data. S.S.M., P.M., M.L., and J.D. performed data analysis. P.M. identified the literature-based CpG sites. S.S.M., M.M.M., C.E.V., and J.D. wrote the manuscript. All authors critically read and revised the manuscript. All authors read and approved the final manuscript.

**Competing interests**

The authors declare no competing interests.

**Additional information**

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