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# Socioeconomic deprivation: an important largely unrecognized risk factor in primary prevention of cardiovascular disease

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# Abstract

**Background**—Socioeconomic deprivation is associated with higher cardiovascular morbidity and mortality. Whether deprivation status should be incorporated in more cardiovascular risk estimation scores remains unclear. This study evaluates how socioeconomic deprivation status affects the performance of three primary prevention cardiovascular risk scores.

Methods—The Generation Scotland Scottish Family Health Study was used to evaluate the performance of three cardiovascular risk scores with (ASSIGN) and without (SCORE2, PCE) socioeconomic deprivation as a covariate in the risk prediction model. Deprivation was defined by Scottish Index of Multiple Deprivation score. The predicted 10-year risk was evaluated against the observed event rate for the cardiovascular outcome of each risk score. The comparison was made across three groups defined by the deprivation index score consisting of group 1 defined as most deprived, group 3 defined as least deprived and group 2 which consisted of individuals in the middle deprivation categories.

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The other authors report no conflicts.

**Results**—The study population consisted of 15,506 individuals (60.0% female, median age of 51). Across the population 1,808 (12%) individuals were assigned to group 1 (most deprived), 8,119 (52%) to group 2, 4,708 (30%) to group 3 (least deprived), and 871 individuals (6%) had missing deprivation score. Risk scores based on models that did not include deprivation status significantly under predicted risk in the most deprived (6.43% observed *versus* 4.63% predicted for SCORE2 [P=0.001] and 6.69% observed *versus* 4.66% predicted for PCE [p<0.001]). Both risk scores also significantly overpredicted the risk in the least deprived group (3.97% observed *versus* 4.72% predicted for SCORE2, P=0.007 and 4.22% observed *versus* 4.85% predicted for PCE, P=0.028). In contrast, no significant difference was demonstrated in the observed versus predicted risk when using the ASSIGN risk score, which included socioeconomic deprivation status in the risk model.

**Conclusions**—Socioeconomic status is a largely unrecognized risk factor in primary prevention of cardiovascular disease. Risk scores that exclude socioeconomic deprivation as a covariate under- and overestimate the risk in the most and least deprived individuals, respectively. This study highlights the importance of incorporating socioeconomic deprivation status in risk estimation systems to ultimately reduce inequalities in health care provision for cardiovascular disease.

#### **Keywords**

Cardiovascular disease; Risk prediction; Socioeconomic deprivation; Calibration

# Introduction

Socioeconomic deprivation is closely associated with cardiovascular morbidity and mortality.<sup>1–5</sup> Whilst the burden from cardiovascular disease has decreased over time, healthcare inequity by deprivation status has persisted.<sup>6</sup> Individuals from poorer backgrounds are less likely to receive evidence-based therapy and more likely to experience higher cardiovascular mortality and morbidity.<sup>7</sup> Whilst previous research has extensively studied provision and management of therapy in the primary prevention of cardiovascular disease, the incorporation of deprivation status in risk estimation systems in primary care is less clear.

Cardiovascular risk estimation is the cornerstone for primary prevention of cardiovascular disease. Despite socioeconomic status being closely associated with cardiovascular mortality and morbidity, most cardiovascular risk estimation systems do not incorporate deprivation status in prediction modelling.<sup>8, 9</sup> In the United Kingdom, the ASSessing cardiovascular risk using SIGN (ASSIGN) and QRISK3 risk scores incorporate deprivation status as a covariate.<sup>10, 11</sup> However, whilst international guidelines do make reference to deprivation as important risk modifier, key European and United States-based cardiovascular risk estimation systems do not incorporate deprivation status in prediction modelling.<sup>8, 9, 12–18</sup> Risk estimation systems may vary in performance by deprivation status,<sup>19–21</sup> highlighting the need for specific research to evaluate whether its inclusion in cardiovascular risk estimation systems is warranted.

We compare the predictive ability of risk scores that include (ASSIGN)<sup>10</sup> or exclude (Systematic COronary Risk Evaluation 2 algorithm [SCORE2]<sup>16</sup>, Pooled Cohort Equations[PCE]<sup>13</sup>) socioeconomic deprivation status in prediction modelling using a large contemporary cohort with over 10 years of follow up.

# **Methods**

Because of the sensitive nature of the data collected for this study, requests to access the dataset from qualified researchers trained in human subject confidentiality protocols should be sent to Generation Scotland management team at access@generationscotland.org.

# Study population

We used data from the Generation Scotland Scottish Family Health Study (GS:SFHS). GS:SFHS is a well-phenotyped family-based contemporary cohort that enrolled 24,090 participants aged between 18 and 98 years as previously described.<sup>22, 23</sup> Briefly, individuals between 35 and 65 years old were identified at random from participating general medical practices in Scotland between February 2006 and March 2011. Participants were then asked to identify 1 first-degree relatives aged 18 years who would also be able to participate.

For this study, participants below 30 years of age, or who had cardiovascular disease at baseline, or who did not attend the clinical survey, were excluded. Participants completed a health questionnaire, and clinical characteristics were measured using a standardized protocol. Data was collected on age, sex, diabetes mellitus, systolic blood pressure, body mass index, family history of cardiovascular disease, smoking status and rheumatoid arthritis. Total cholesterol, high-density lipoprotein cholesterol, and serum creatinine, were measured at the time of collection. Ethical approval for the GS:SFHS study was obtained from the NHS Tayside Research Ethics Committee (REC reference number 05/S1401/89). Study participants provided written informed consent, including linkage to their medical records. The study was conducted according to principles of the Declaration of Helsinki.

#### Outcomes

We used the Information Services Division National Health Service record linkage for Scotland to collect non-fatal cardiovascular events and cause-specific death data for each individual from the date of inclusion in the study until the end of August 2021. Information on cause of death was obtained using the National Health Service Central Register. Nonfatal cardiovascular events and cause-specific deaths were classified using the 10<sup>th</sup> revision of the International Classification of Diseases (ICD-10).

#### Socioeconomic deprivation status

Socioeconomic deprivation status was determined using the Scottish Index of Multiple Deprivation (SIMD) 2009 score which is derived from participants' postcodes and compiled using seven domains of deprivation (income, employment, education, health, access to services, crime and housing).<sup>24</sup> The SIMD score is recalculated every few years and the scores used are from 2009, at the mid point of GS:SFHS recruitment. SIMD scores range from 0.94 (least deprived) to 89.89 (most deprived), with quintiles based on the full

derivation cohort, reflecting the wider Scottish population. The cut offs are as follows: SIMD < 7.94 (quintile 5), 7.94 SIMD < 13.67 (quintile 4), 13.67 SIMD < 20.98 (quintile 3), 20.98 SIMD < 33.81 (quintile 2), and SIMD 33.81 (quintile 1).<sup>24, 25</sup> For this study, we classified patients into three groups using these quintile-based cut-offs: group 1 (most deprived based on quintile 1), group 2 (based on quintiles 2-4) and group 3 (least deprived based on quintile 5).

# Cardiovascular risk scores

The performance of ASSIGN, SCORE2 and the PCE risk scores were evaluated.<sup>10, 13, 16</sup> Of these only the ASSIGN risk score includes socioeconomic deprivation as a covariate, and the ASSIGN risk score uses the Scottish Index of Multiple Deprivation score divided by 10 as a covariate in the risk equation. Beta coefficients, centring values and 10-year baseline survival for each risk score were extracted. Outcomes were based on ICD-10 diagnostic classification, and outcomes in the GS:SFHS cohort were mapped to the outcomes of each risk score using ICD-10 codes. For ASSIGN, this is cardiovascular death (I00-I99), the first occurrence of hospitalisation with coronary disease (I20-I25) or stroke (G45 and I60-I69), and Office of Population Censuses and Surveys: Classification of Interventions and Procedures version 4 (OPCS-4) procedure codes (L29.5, L31.1, K40-46, K49, and K75 [procedures comprising carotid endarterectomy, carotid angioplasty, coronary artery bypass graft, and percutaneous transluminal coronary angioplasty]). The SCORE2 outcome was defined more narrowly than ASSIGN as cardiovascular death (I10-I16, I20-I25, I46-I52, I60-I69, I70-I73, R96.0-R96.1 [excluding I51.4, I60, I62, I67.1, I68.2, I67.5]), the first occurrence of non-fatal stroke (G45 and I60-I69) and non-fatal myocardial infarction (I21-I22). For PCE, the outcome was defined as the first occurrence of non-fatal myocardial infarction (I21-I22), fatal coronary heart disease (I20-I25), or fatal or non-fatal stroke (G45 and I60-I69), again narrower than the ASSIGN outcome. More information on the derivation cohorts, outcomes, covariates, statistical approach and model equations for each risk score are provided in Table S1.

#### Statistical analysis

Continuous variables are presented as median and interquartile range [IQR], and categorical variables are presented as absolute number (%). Ten-year estimated cardiovascular disease risks for each patient were calculated using the published risk models for ASSIGN, SCORE2 and PCE. The observed 10-year event rates were derived using Kaplan-Meier estimates to account for differing follow-up times among individuals.

Recalibration of the baseline survival was conducted to diminish over- or underestimation of risk. This was done by replacing the original 10-year baseline survival with the updated 10-year baseline survival derived from the GS:SFHS cohort. Recalibration was done in the whole cohort, not within each socioeconomic strata. We evaluated the performance of the recalibrated and non-recalibrated risk scores by assessing measures of calibration and discrimination, stratified by socioeconomic deprivation status. Calibration refers to how closely the predicted 10-year risk agrees with the observed 10-year risk. Calibration plots were constructed using deciles of predicted risk scores. We evaluated the calibration intercept and slope of each plot. Furthermore, we calculated an observed *versus* predicted ratio

by dividing the predicted risk by observed risk and evaluated whether the ratios differed between socioeconomic deprivation groups. We conducted a Z-test to evaluate differences between predicted and observed risks, taking account of the uncertainty in the observed risk, and a P-value <0.05 was considered statistically significant. Discrimination is the ability of the risk score to differentiate between patients who do and do not experience an event during the study period, and discrimination was assessed using the C-statistic. To further explore whether socioeconomic depreviation should be incorporated in cardiovascular risk scores, we fitted sex-specific Cox regression models to the GS:SFHS cohort using the same outcomes, model structure and covariates as were used to derive the original ASSIGN, SCORE2 and PCE risk scores, but with socioeconomic deprivation status as an additional risk factor. The present manuscript follows the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) guidelines.<sup>26</sup> For the primary analysis. multiple imputation by chained equations (MICE) was used to impute missing values of covariate data using fully-conditional models including clinical characteristics and outcomes, and a single imputed dataset was selected. A complete-case sensitivity analysis was also performed. All statistical analyses were carried out in R (version 3.6.1). Key packages used were 'survival' to calculate Kaplan-Meier survival probabilities, fit relevant Cox proportional hazards models and 'ggplot2' to produce calibration plots. The R code is available on GitHub (https://github.com/leahpirondini/risk-score-calibration).

# Results

#### Study population

A total of 15,506 individuals (60.0% female, median 51 years of age) were included in our study. Individuals in the most deprived group were more likely to be female (group 1, 63.8% *versus* group 3, 57.9%), current smokers (group 1, 30.1% *versus* group 3, 8.5%), and to have diabetes (group 1, 4.0% *versus* group 3, 2.0%) (Table 1). Despite being younger on average, individuals in the most deprived group had the highest incident risk of future cardiovascular events (Table 2). At 10 years, the cumulative incidence of cardiovascular death was 2.2%, 1.6%, and 1.4% for socioeconomic deprivation groups 1 (most deprived) to 3 (least deprived), respectively.

# Performance of cardiovascular risk scores by socioeconomic deprivation status

In the most deprived individuals, no statically significant difference was demonstrated between the observed *versus* predicted risk when using the recalibrated ASSIGN risk score (9.13% observed *versus* 8.39% predicted, P=0.256, Figure 1, Table 3). In contrast, risk scores based on the recalibrated SCORE2 and PCE models that did not include deprivation status, significantly under predicted risk in the most deprived (Figure 2, Table 3). For SCORE2, the observed risk was 6.43% in the most deprived individuals whereas the predicted risk was 4.63% (P=0.001). Similarly, for PCE, the observed risk was higher at 6.69% than the predicted risk of 4.66% in those who were most deprived (P<0;0.001). A minimal and non-significant difference was demonstrated in the observed *versus* 6.45% predicted, P=0.478, Figure 1, Table 3). However, both SCORE2 and PCE risk scores significantly over predicted the risk in the least deprived group (3.97% observed *versus* 

4.72% predicted for SCORE2, P=0.007 and 4.22% observed *versus* 4.85% predicted for PCE, P=0.028, Figure 2, Table 3).

In group 2, consisting of individuals in the middle deprivation categories, we demonstrated no statistically significant differences in observed *versus* predicted risk across all three risk scores (Table 3). For ASSIGN, the observed and predicted risk was 6.75% and 6.92% (P=0.527), respectively (Figure 1, Table 3). Both SCORE2 and PCE showed good agreement between observed and predicted risk (4.56% observed *versus* 4.72% predicted for SCORE2, P=0.479 and 4.73% observed *versus* 4.88% predicted for PCE, P=0.514, Figure 2, Table 3). The complete-case analysis yielded similar results (data not shown).

We additionally studied the performance of the non-recalibrated cardiovascular risk scores across socioeconomic deprivation groups to evaluate the "raw" cardiovascular risk estimates of each risk score. None of the risk scores performed well before recalibration (Figure S1, Table S2-S3). All three risk scores showed good discrimination with slight differences between socioeconomic deprivation groups (Table S4). We further explored whether socioeconomic deprivation should be incorporated in cardiovascular risk scores, by fitting sex-specific Cox regression models on GS:SFHS using the same framework (outcomes, covariates and model structures) as the risk scores ASSIGN, SCORE2 and PCE model and with socioeconomic deprivation status as an additional risk factor. All refitted risk models showed a significant contribution of the deprivation index score in the male-specific models (HR, 1.148 [95% CI 1.092 to 1.206, ASSIGN], 1.176 [95% CI 1.105 to 1.251, SCORE2], log HR 0.166 [95% CI 0.111 to 0.221, PCE], Table S5-S7).

# Discussion

We evaluated the impact of socioeconomic deprivation on the performance of three primary prevention cardiovascular risk scores that are widely used in clinical practice. The main finding of our study is that socioeconomic deprivation status is an important covariate in cardiovascular risk estimation systems. Risk scores (SCORE2 and PCE) that exclude socioeconomic deprivation in prediction modelling, meaningfully under- and over-estimate the risk in the most and least deprived populations, respectively.<sup>17, 18</sup> As such, a substantial proportion of people living in more deprived communities at higher risk are likely to remain undertreated. The ASSIGN risk score - that includes socioeconomic deprivation in the risk prediction model - shows good performance in individuals living in most and least deprived areas.<sup>27</sup> Our analysis highlights the importance of incorporating a measure of socioeconomic deprivation status in risk estimation systems to reduce inequalities in health care provision for cardiovascular disease.

Low socioeconomic status is associated with worse cardiovascular outcomes<sup>1–5</sup> and our study showed that the magnitude of this association was strengthened after adjusting for sex and age. This is in line with published data showing that the differences in mortality rates between socioeconomic classes increased when age decreased, and observed mortality rates were highest in the youngest age-groups.<sup>28</sup> One in three premature deaths were attributable to socioeconomic inequalities and predominantly driven by cardiovascular disease.<sup>29</sup> Our study shows that socioeconomic status is a largely unrecognized risk factor

in primary prevention of cardiovascular disease. Incorporation of socioeconomic status into cardiovascular risk estimates is important to improving outcomes and closing the gap between the most and least deprived. The ASSIGN risk score was the first cardiovascular risk score developed that included socioeconomic deprivation status as a covariate to achieve equality for deprived individuals, and showed improvement – although marginal – in risk estimation as compared with the Framingham Risk Score that included only traditional risk factors.<sup>10</sup> Similarly, ORISK2 – which also includes socioeconomic deprivation as a covariate - showed higher accuracy when compared to the Framingham Risk Score in the national QRESEARCH database comprised of 2.29 million patients.<sup>30</sup> In addition, incorporation of socioeconomic status to the Framingham Risk Score was evaluated in the Atherosclerosis Risk in Communities study.<sup>31</sup> When socioeconomic status was incorporated as individual-based measures using income and education in the Framingham Risk Score, the bias towards deprived individuals disappeared.<sup>31</sup> In line with these results, our study clearly demonstrates a bias to the most and least deprived areas towards under- and overestimating risk when using risk estimation systems that do not incorporate deprivation status. Furthermore, our analysis also shows that adding a deprivation index score to models using the same covariates as those of SCORE2 and PCE significantly contributed to the prediction of future cardiovascular events for men.

Cardiovascular risk estimation systems that do not incorporate deprivation status as a covariate may falsely classify the most deprived individuals at lower risk, potentially denying them the benefit of pharmacological and non-pharmacological primary prevention therapy. Similarly, the least deprived individuals may be falsely classified as high risk leading to potential overtreatment. A previous study modelling the potential impact of using risk estimation systems incorporating deprivation status showed that such a risk estimation system would result in initiation of lipid lowering therapy in 1 in 7 untreated individuals in the general population.<sup>32</sup>

None of the risk scores performed well when not recalibrated. Although the ASSIGN risk score has been tailored to the Scottish population, the estimated risk was two-fold higher compared to the observed risk across the entire study population. The ASSIGN risk score was derived on a population in the 1980s where the baseline risk was high.<sup>10</sup> This baseline risk has fallen dramatically over the last 25 years,<sup>5</sup> and the overestimation is most likely the result of an inaccurate baseline risk used in the risk equation.<sup>5</sup> Our study shows that cardiovascular risk estimates could be optimized when risk scores are recalibrated using contemporary local data, which is in line with previous studies.<sup>8</sup>, <sup>33–35</sup> Of the three risk scores evaluated, only SCORE2 acknowledged the need for recalibration and has used contemporary data to recalibrate their prediction models during development.<sup>16, 36</sup> However, we feel that the recalibration process should not be a static process. Cardiovascular risk estimation systems can be further optimized when a continuous recalibration system is in place. For example, QRISK3 score is updated on an annual basis using contemporary local data to ensure that the baseline survival and mean of covariates used in risk equation reflects the target population that is being evaluated.<sup>11</sup>

Our study has several strengths. First, we used a contemporary cohort of over 15,000 individuals that enabled us to evaluate the performance in a large group of deprived

individuals and to recalibrate risk scores. Second, we had 10 years of follow-up available that allowed us to report an individual's observed 10-year risk. Third, our cohort had adequate phenotyping at baseline for us to evaluate three cardiovascular risk scores that are commonly applied in clinical practice, and the outcomes for each risk score were matched to those used for derivation of the score as closely as possible.

We also acknowledge several limitations. First, although individuals were randomly invited to participate in GS:SFHS, the response rate was higher in less deprived individuals, and this might have tended to underestimate differences. Second, GS:SFHS predominantly includes individuals of Caucasian background, and we cannot generalize our findings to individuals of other ethnic backgrounds. Third, we acknowledge that the social deprivation score used in our study is an area-based measure rather than the socioeconomic status of the individual introducing ecological bias, whereby an individual who lives in a more deprived area need not necessarily have to experience a high level of deprivation. This also highlights that deprivation based on individual-level data may perform better than those based on geographical location. Furthermore, future work related to individual socioeconomic measures and risk prediction modelling is needed to unravel which component does particularly contribute to our observed findings. Fourth, our analysis showed that social deprivation was positively associated with cardiovascular outcomes for both men and women. However, the associations were stronger for men compared to women, and for women the 95% confidence interval crossed the line of unity. These observed differences in the effect estimate need further evaluation. Finally, we limited our analysis to three cardiovascular risk scores. More risk scores have been developed over the years that are not included in our analysis, but we made the decision to focus particularly on those that are widely applied in practice across North America and Europe. We also did not include other risk estimation systems, including those with deprivation status, due to the lack of availability of model covariates or concordant outcomes.<sup>11</sup>

In conclusion, socioeconomic deprivation status is an important covariate in cardiovascular risk estimation systems. Risk scores that exclude socioeconomic deprivation underand over-estimate risk in the most and least deprived individuals, respectively. Our findings highlight the importance of incorporating socioeconomic deprivation status in risk estimation systems to reduce inequalities in health care provision for cardiovascular disease.

# Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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# Non-standard Abbreviations and Acronyms

ASSIGN	ASSessing cardiovascular risk using SIGN
CI	confidence interval
GS:SFHS	Generation Scotland Scottish Family Health Study
HR	hazard ratio
ICD-10	10 <sup>th</sup> revision of the International Classification of Diseases
IQR	interquartile range
MICE	multiple imputation by chained equations
OPCS-4	Office of Population Censuses and Surveys: Classification of Interventions and Procedures version 4
PCE	Pooled Cohort Equations
SCORE2	Systematic COronary Risk Evaluation 2 algorithm
SIMD	Scottish Index of Multiple Deprivation
STROBE	Strengthening the Reporting of Observational Studies in EpidemiologyClinical perspective

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# **Clinical perspective**

# What is new?

- We report the impact of socioeconomic deprivation on the performance of three primary prevention cardiovascular risk scores that are widely used in practice.
- Socioeconomic deprivation status is an important covariate in cardiovascular risk estimation systems, and risk scores that exclude socioeconomic deprivation under- and over-estimate risk in the most and least deprived individuals, respectively.

#### What are the clinical implications?

- Socioeconomic status is a largely unrecognized risk factor in primary prevention of cardiovascular disease.
- Our findings highlight the importance of socioeconomic deprivation status as a covariate that needs to be considered in addition to the traditional risk factors to promote equitable healthcare, particularly in those most deprived.

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Figure 2. Evaluation of the calibration of the recalibrated SCORE2 (panel A), and PCE (panel B) risk scores using the predicted and observed 10-year risk, stratified by socioeconomic deprivation status.

Each dot represents one decile of risk and is surrounded by 95% confidence interval.

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	All (N=15,506)	Group 1 (most deprived) (N=1,808)	Group 2 (N=8,119)	Group 3 (least deprived) (N=4,708)	Missing (N=871)
Deprivation index score *	11.3 (6.8 to 21.7)	46.0 (39.3 to 54.5)	14.4 (10.5 to 20.8)	5.4 (3.9 to 6.7)	ı
Age (years)	51 (42 to 59)	48 (40 to 57)	51 (41 to 59)	53 (44 to 60)	49 (40 to 58)
Sex (male)	6,202 (40.0%)	654 (36.2%)	3,215 (39.6%)	1,982(42.1%)	351 (40.3%)
Body mass index (kg/m <sup>2</sup> )	26.2 (23.5 to 29.5)	27.2 (24.0 to 31.4)	26.4 (23.6 to 29.6)	25.7 (23.2 to 28.6)	26.7 (23.6 to 29.9)
Systolic blood pressure (mmHg)	131 (120 to 144)	130 (120 to 142)	131 (120 to 144)	131 (121 to 144)	131 (119 to 144)
Total cholesterol (mmol/L)	5.2 (4.6 to 5.9)	5.1 (4.5 to 5.9)	5.2 (4.5 to 5.9)	5.3 (4.6 to 6.0)	5.2 (4.5 to 5.9)
HDL cholesterol (mmol/L)	1.4 (1.2 to 1.7)	1.3 (1.1 to 1.6)	1.4 (1.2 to 1.7)	1.5 (1.2 to 1.8)	1.4 (1.2 to 1.7)
Creatinine (mmol/L)	72 (63 to 82)	71 (63 to 81)	71 (63 to 82)	72 (63 to 83)	73 (64 to 83)
Current smokers (yes)	2,296 (15.3%)	514 (30.1%)	1,248 (15.9%)	390 (8.5%)	144 (17.0%)
Cigarettes per day (smokers)	17 (9 to 21)	17 (12 to 22)	17 (9 to 22)	12 (7 to 20)	14 (7 to 20)
Family history of CVD (yes)	5,942 (39.1%)	665 (37.8%)	3,136 (39.4%)	1,815(39.3%)	326 (38.0%)
Rheumatoid arthritis (yes)	267 (1.7%)	52 (2.9%)	135 (1.7%)	64 (1.4%)	16(1.8%)
Diabetes mellitus (yes)	416 (2.7%)	72 (4.0%)	230 (2.8%)	95 (2.0%)	19 (2.2%)
Lipid-modifying medication (yes)	932 (6.0%)	128 (7.1%)	493 (6.1%)	249 (5.3%)	62 (7.1%)
Antihypertensive medication (yes)	1,268 (8.2%)	145 (8.0%)	696 (8.6%)	360 (7.6%)	67 (7.7%)
* Scottish index of multinle denrivatio	n. Categorical values s	re presented as n (%). and co	ntinuous variables are	presented as median (25 <sup>th</sup> to	75th nercentile), as ann

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15,506 participants, except where number missing (N missing) is indicated in the column. Missing values < 5% if applicable, except for deprivation index score (5.6%). Abbreviations: CVD, cardiovascular disease; HDL, high density lipoprotein.

Table 2

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	ЧI	Group 1 (most deprived)	Group 2	Group 3 (least deprived)	Unadjusted OR (95% CI) Group 1 versus 3	Adjusted OR (95% CI) Group 1 versus 3*	Adjusted OR (95% CI) Group 1 versus 3 <sup>†</sup>
Myocardial	infarction						
At 1 year	20 (0.1%)	2 (0.1%)	14 (0.2%)	3 (0.1%)			
At 5 years	119 (0.8%)	24 (1.4%)	68 (0.8%)	21 (0.4%)			
At 10 years	263 (1.7%)	49 (2.8%)	133 (1.7%)	69 (1.5%)	1.87 (1.29 to 2.70)	2.44 (1.67 to 3.56)	1.96 (1.28 to 2.98)
Stroke							
At 1 year	19 (0.1%)	2 (0.1%)	9 (0.1%)	6~(0.1%)			
At 5 years	121 (0.8%)	23 (1.3%)	58 (0.7%)	30 (0.6%)			
At 10 years	310 (2.1%)	45 (2.6%)	166 (2.1%)	78 (1.7%)	1.52 (1.04 to 2.18)	1.89 (1.29 to 2.75)	1.32 (0.85 to 2.01)
Cardiovascu	ılar death						
At 1 year	12 (0.1%)	3 (0.2%)	5~(0.1%)	3 (0.1%)			
At 5 years	90 (0.6%)	17 (0.9%)	46 (0.6%)	22 (0.5%)			
At 10 years	249 (1.6%)	39 (2.2%)	128 (1.6%)	66 (1.4%)	1.55 (1.03 to 2.30)	2.07 (1.35 to 3.14)	1.14 (0.68 to 1.86)
Composite C	VD outcome	<i>‡</i>					
At 1 year	47 (0.3%)	7 (0.4%)	25 (0.3%)	12 (0.3%)			
At 5 years	293 (1.9%)	57 (3.2%)	154 (1.9%)	64 (1.4%)			
At 10 years	719 (4.8%)	116 (6.7%)	371 (4.7%)	191 (4.2%)	1.62 (1.28 to 2.05)	2.20 (1.71 to 2.82)	1.50 (1.12 to 1.99)
Abbreviations:	CVD, cardiov	ascular disease. N=1	4,635 individu	ıals with available d	eprivation index score	š	
*							
Logistic regre	ssion model. <b>1</b>	The model is adjusted	d for sex and a	ge.			

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ź Logistic regression model. The model is adjusted for sex, age, body mass index, systolic blood pressure, total cholesterol, high-density lipoprotein, cigarettes per day, diabetes mellitus.

 ${}^{\sharp}$ Composite CVD outcome included non-fatal myocardial infarction, non-fatal stroke and cardiovascular death.

# Table 3

# Observed and predicted 10-year cardiovascular risk stratified by socioeconomic deprivation status

	Observed (%)	Predicted (%)	P-value	Ratio*
ASSIGN				
All	6.87%	6.95%	0.676	1.01
Group 1 (most deprived)	9.13%	8.39%	0.256	0.92
Group 2	6.75%	6.92%	0.527	1.03
Group 3 (least deprived)	6.21%	6.45%	0.478	1.04
SCORE2				
All	4.60%	4.71%	0.526	1.02
Group1 (most deprived)	6.43%	4.63%	0.001	0.72
Group 2	4.56%	4.72%	0.479	1.03
Group 3 (least deprived)	3.97%	4.72%	0.007	1.19
PCE				
All	4.81%	4.84%	0.848	1.01
Group 1 (most deprived)	6.69%	4.66%	<0.001	0.70
Group 2	4.73%	4.88%	0.514	1.03
Group 3 (least deprived)	4.22%	4.85%	0.028	1.15

\* Ratio = predicted risk divided by observed risk.

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