Application of non-HDL cholesterol for population-based cardiovascular risk stratification: results from the Multinational Cardiovascular Risk Consortium


Summary

Background The relevance of blood lipid concentrations to long-term incidence of cardiovascular disease and the relevance of lipid-lowering therapy for cardiovascular disease outcomes is unclear. We investigated the cardiovascular disease risk associated with the full spectrum of bloodstream non-HDL cholesterol concentrations. We also created an easy-to-use tool to estimate the long-term probabilities for a cardiovascular disease event associated with non-HDL cholesterol and modelled its risk reduction by lipid-lowering treatment.

Methods In this risk-evaluation and risk-modelling study, we used Multinational Cardiovascular Risk Consortium data from 19 countries across Europe, Australia, and North America. Individuals without prevalent cardiovascular disease at baseline and with robust available data on cardiovascular disease outcomes were included. The primary composite endpoint of atherosclerotic cardiovascular disease was defined as the occurrence of the coronary heart disease event or ischaemic stroke. Sex-specific multivariable analyses were computed using non-HDL cholesterol categories according to the European guideline thresholds, adjusted for age, cohort, and classical modifiable cardiovascular risk factors. In a derivation and validation design, we created a tool to estimate the probabilities of a cardiovascular disease event by the age of 75 years, dependent on age, sex, and risk factors, and the associated modelled risk reduction, assuming a 50% reduction of non-HDL cholesterol.

Findings Of the 524,444 individuals in the 44 cohorts in the Consortium database, we identified 398,846 individuals belonging to 38 cohorts (184,655 [48.7%] women; median age 51.0 years [IQR 40.7–59.7]), 199,415 individuals were included in the derivation cohort (91,786 [48.4%] women) and 199,431 (92,269 [49.1%] women) in the validation cohort. During a maximum follow-up of 43.6 years (median 13.5 years, IQR 7.0–20.1), 54,542 cardiovascular endpoints occurred. Incidence curve analyses showed progressively higher 30-year cardiovascular disease event rates for increasing non-HDL cholesterol categories (from 7.7% for non-HDL cholesterol <2.6 mmol/L to 33.7% for ≥5.7 mmol/L in women and from 12.8% to 43.6% in men; p<0.0001). Multivariable adjusted Cox models with non-HDL cholesterol lower than 2.6 mmol/L as reference showed an increase in the association between non-HDL cholesterol concentration and cardiovascular disease for both sexes (from hazard ratio 1.1, 95% CI 1.0–1.3 for non-HDL cholesterol 2.6 to <3.7 mmol/L to 1.9, 1.6–2.2 for ≥5.7 mmol/L in women and from 1.1, 1.0–1.3 to 2.3, 2.0–2.5 in men). The derived tool allowed the estimation of cardiovascular disease event probabilities specific for the age of 75 years, dependent on age, sex, and risk factors, and the associated modelled risk reduction, assuming a 50% reduction of non-HDL cholesterol.

Interpretation Non-HDL cholesterol concentrations in blood are strongly associated with long-term risk of atherosclerotic cardiovascular disease. We provide a simple tool for individual long-term risk assessment and the potential benefit of early lipid-lowering intervention. These data could be useful for physician–patient communication about primary prevention strategies.

Funding EU Framework Programme, UK Medical Research Council, and German Centre for Cardiovascular Research.

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Evidence before this study
The blood concentrations of non-HDL and LDL cholesterol are accepted as causal cardiovascular risk factors and constitute a cornerstone of cardiovascular disease risk prediction in the general population. The benefit of lipid-lowering interventions for cardiovascular disease risk reduction is proven by mendelian randomisation studies, prospective epidemiological cohort studies, and randomised trials. For the present study, we assessed recommendations made by the European and US guidelines and their underlying references for lipid-lowering strategies in primary prevention. Furthermore, we reviewed the cardiovascular risk calculators as recommended by these guidelines (Systematic Coronary Risk Evaluation [SCORE] and Pooled Cohort Equations). We found very little data describing the effect of baseline non-HDL and LDL cholesterol on long-term or even lifetime outcomes. Cardiovascular risk assessment was mainly based on 10-year medium-term follow-up. All identified randomised trials for lipid-lowering intervention in cardiovascular primary prevention did not exceed a follow-up of 7 years and, moreover, especially young individuals were underrepresented in those studies.

Added value of this study
To our knowledge, our study provides the most comprehensive analysis of long-term risk for cardiovascular disease related to non-HDL cholesterol and offers an easily applicable tool to assess the long-term probabilities for cardiovascular disease events associated with non-HDL cholesterol. On the basis of a derivation and validation approach to risk prediction, our data provide robust multinational information. By using an up-to-date multinational population-based pooled cohort dataset, we provide a model that calculates the potential benefit of an early lipid-lowering strategy in individuals without prevalent cardiovascular disease across a range of non-HDL cholesterol categories.

Implications of all the available evidence
There is evidence for the positive outcomes in lipid-lowering interventions in individuals with cardiovascular disease or those with very high cholesterol concentrations. Our data extend current knowledge and highlight the impact of non-HDL cholesterol on very long-term cardiovascular disease outcomes. A simulation of early reduction of non-HDL cholesterol across a range of lipid concentration categories estimates the risk reduction in the general population. Meeting targets for global reduction of cardiovascular disease morbidity and mortality will require increased awareness of the value of early cholesterol determination and careful evaluation of potential strategies for reduction of lipid concentrations across the lifespan.

Introduction
Numerous studies have provided consistent evidence for a causal relationship between blood cholesterol concentrations and cardiovascular disease. Calculating the concentration of non-HDL cholesterol offers a simple way to analyse the total amount of proatherogenic lipoproteins containing apolipoprotein B (apoB). Such proteins include very low-density lipoproteins and their metabolic remnants, intermediate-density lipoproteins, lipoprotein(a), and low-density lipoproteins. Besides estimating cholesterol concentrations contained in LDL particles (LDL cholesterol), the assessment of non-HDL cholesterol calculated as total cholesterol minus HDL cholesterol is therefore recommended by current US and European guidelines for cardiovascular risk estimation. The indication for lipid-lowering therapy to prevent cardiovascular disease events in high-risk individuals or for secondary prevention is unequivocal. However, the decision on implementing a lipid-lowering intervention in the primary prevention setting is a major challenge in clinical practice for several reasons. Data on the association between the concentrations of the entire range of bloodstream lipids and very long-term cardiovascular outcomes in the general population are rather sparse. In addition, conventional primary prevention guidelines recommend lipid-lowering intervention on the basis of lipid concentration thresholds and the person’s individual 10-year cardiovascular risk. This risk assessment might underestimate or fail to take proper account of the cumulative lifetime risk of cardiovascular disease and lipid-related cardiovascular disease risk, particularly in young adults. Furthermore, increased non-HDL cholesterol blood concentrations early in life seem to be stable over the life course and are predictive for incident cardiovascular disease. Therefore, in this study we aimed to (1) evaluate long-term risk for cardiovascular disease in the population related to non-HDL cholesterol on the basis of existing thresholds of blood lipid concentrations; (2) establish an easily applicable tool to assess the long-term probabilities for cardiovascular disease events associated with non-HDL cholesterol, using a derivation and validation approach; and (3) provide a model indicating the potential benefit of an early lipid-lowering strategy in individuals without prevalent cardiovascular disease.

Methods

Study design and participants
In this risk-evaluation and risk-modelling study, we used individual-level data from the Multinational Cardiovascular Risk Consortium (appendix pp 2–11). Currently, the Consortium comprises data on 524 444 individuals from 44 population-based cohorts across Europe, Australia, and the USA. These include 23 harmonised cohorts from the MONICA Risk Genetics Archiving and Monograph (MORGAM) and Biomarker for Cardiovascular Risk Assessment in Europe (BiomarCaRE) studies and another 21 harmonised, population-based cohorts with the same
specifications as defined for the BiomarCaRE and MORGAM projects (appendix pp 15–18). Of these, we used cohorts that had available data on cardiovascular disease endpoints (appendix pp 16–18). We excluded data on participants with prevalent cardiovascular disease, defined as a history of myocardial infarction, coronary artery bypass grafting, percutaneous transluminal coronary angioplasty, or ischaemic or haemorrhagic stroke. Detailed information about the study cohorts including the number of individuals with available information for each variable are provided in the appendix (pp 19–27). Informed consent was obtained if necessary (depending on survey year and local requirement). Detailed information for each cohort is given in the study descriptions (appendix pp 2–11).

Procedures
Our outcome measure was the first occurrence of a major cardiovascular event (appendix pp 16–18). This was assessed in all participating individuals and death of non-cardiovascular-disease cause was used as a competing risk if appropriate. We defined it as a composite endpoint of the first non-fatal or fatal coronary heart disease or ischaemic stroke event. Coronary heart disease was defined as a composite of non-fatal or fatal myocardial infarction including unstable angina, coronary death, and coronary revascularisation. Outcomes were measured in each prospective cohort study by standardised follow-up (visit, report, registry information, etc; details are given for each cohort, appendix pp 2–11). Composite endpoint information for current analyses is provided in the appendix (pp 16–18). There were no secondary outcomes.

We based our cholesterol threshold concentrations on the treatment-determining cholesterol categories of the 2016 European guidelines on cardiovascular disease prevention. As recommended by the guidelines, we calculated non-HDL cholesterol thresholds by adding 30 mg/dL to the LDL cholesterol thresholds defined in our cohorts. Our analyses were based on baseline information available for concentrations of non-HDL cholesterol and LDL cholesterol. We report threshold concentrations in mmol/L, assuming that 1 mmol/L contains 38-67 mg/dL of LDL cholesterol and non-HDL cholesterol for non-HDL cholesterol, the following categories were used: <100 mg/dL [<2.6 mmol/L], 100 to <145 mg/dL [2.6 to <3.7 mmol/L], 145 to <185 mg/dL [3.7 to <4.8 mmol/L], 185 to <220 mg/dL [4.8 to <5.7 mmol/L], and ≥220 mg/dL [≥5.7 mmol/L]. The cutoff between the second and third category differs slightly from the European guidelines. We adapted this cutoff from 130 mg/dL (3.4 mmol/L) to 145 mg/dL (3.7 mmol/L), since the latter is the goal for lipid-lowering therapy recommended for individuals at low risk, who comprise most of our study population.

The following study variables were considered for analysis (but were not endpoints): age, sex, year of examination, body-mass index (BMI), systolic blood pressure, smoking status, use of cholesterol-lowering medication, total cholesterol, HDL cholesterol, and non-HDL cholesterol. History of arterial hypertension, diabetes, stroke, and myocardial infarction were defined as they had been documented or self-reported. Anti-hypertensive medication, lipid-lowering medication, and smoking status were self-reported. For individuals receiving lipid-lowering therapy, baseline concentrations of non-HDL cholesterol and LDL cholesterol were inflated by 30% for the analyses, assuming that treatment with statins had a moderate effect on lipid reduction and was initiated late during lifetime. Information about individuals receiving lipid-lowering therapy stratified by survey decades and age is shown in the appendix (p 28).

Statistical analysis
For each analysis, only individuals with all data available on the relevant variables were used. To investigate the association of non-HDL cholesterol concentrations and time to cardiovascular disease, sex-specific cumulative incidence curves were produced according to the defined non-HDL cholesterol threshold concentrations, with death of non-cardiovascular-disease-causes as competing risk, using the Aalen-Johansen estimator. Multivariable Cox proportional hazards models were then computed, stratified for cohort and sex. Non-HDL cholesterol was coded as a categorical variable, with the help of dummy variables, using our predefined thresholds and included an interaction term with sex, to obtain sex-specific associations. To examine the association of non-HDL cholesterol across different ages, the models were expanded by adding an interaction between age category (<45, 45–59, and ≥60 years at baseline) and non-HDL cholesterol. Additional Cox regression analyses were done to model the concentrations of non-HDL cholesterol as a continuous variable via cubic splines. All Cox models were adjusted for age (timescale), sex (strata), cohort (strata), and classical cardiovascular risk factors (smoking, diabetes, BMI, systolic blood pressure, and antihypertensive medication). In a sensitivity analysis, sex-specific hazard ratios (HRs) for non-HDL cholesterol as a categorical variable were estimated in each country using models similar to the primary analysis, and the results were pooled using a random-effects multivariate meta-analysis. Further sensitivity analyses include the computation of time-dependent HRs for non-HDL cholesterol to examine deviations from the proportional hazards assumption (appendix p 29).

To establish a tool to assess the probability of cardiovascular disease by the age of 75 years, we split the whole data randomly into two parts of approximately equal size into derivation and validation datasets. In particular, each single cohort was part of both datasets. Using those observations where the required endpoint and covariate information was available, we developed a model that was calculated on the basis of cause-specific Cox models for cardiovascular disease stratified Russian Academy of Sciences, Novosibirsk, Russia (Prof Y P Nikonov PhD); Department of Epidemiology, Cardiovascular Disease Prevention and Health Promotion, National Institute of Cardiology, Warsaw, Poland (Prof W Drygas PhD); Department of Clinical Sciences, Malmö, Lund University, Malmö, Sweden (Prof D Melander MD); Institute for Medical Informatics, Biometry and Epidemiology, University Hospital of Essen, Essen, Germany (Prof K H Jockel PhD); Department of Cardiology, Charité Berlin—University Medicine, Campus Benjamin Franklin, Berlin, Germany (D M Leitner MD, Prof U Landmesser MD); German Centre for Cardiovascular Research, Partner Site Berlin, Berlin, Germany (D M Leitner, Prof U Landmesser); Berlin Institute of Health, Berlin, Germany (Prof U Landmesser); Baker Heart and Diabetes Institute, Melbourne, VIC, Australia (Prof J Shaw MD); Department of Nutrition and Dietetics, School of Health Science and Education, Harokopio University, Athens, Greece (Prof D Pagourtzis MD); University of New South Wales, Sydney, NSW, Australia (Prof A A Simons MD); St Vincent’s Hospital, Sydney, NSW, Australia (Prof A A Simons); Department of Epidemiology, Erasmus Medical Center, University Medical Center Rotterdam, Rotterdam, Netherlands (M Kavousi PhD); Boston University and the National Heart, Lung, and Blood Institute’s Framingham Study, Framingham, MA, USA (Prof S S Vasan MD); Department of Endocrinology, University Medical Center Groningen, University of Groningen, Groningen, Netherlands (R F Dulliaan MD); Department of Primary Care and Population Health, University College London, London, UK (Prof G Wannamethee PhD); Department of Public Health and Caring Sciences, Clinical Nutrition and Metabolism, Uppsala University, Uppsala, Sweden (Prof U Risérus PhD); Departments of Medicine and articles articles www.thelancet.com Vol 394 December 14, 2019 2175
for sex and cohort. Death from non-cardiovascular-disease causes was added as a competing risk, to account for the fact that an individual dying from such a cause before developing cardiovascular disease will not be able to develop cardiovascular disease. These models used age as the timescale and the same variables as our previous Cox models. Non-HDL cholesterol, BMI, and systolic blood pressure were modelled using cubic splines. The models included an interaction between non-HDL cholesterol and sex. The model computation used the full available follow-up periods but was applied only to individuals aged between 35 and 70 years at baseline to estimate their probability of cardiovascular disease by the age of 75 years. These predicted probabilities were averaged within each combination of sex, non-HDL cholesterol category, age category, and according to whether the number of risk factors present (smoking, arterial hypertension, diabetes, and obesity) was greater than or equal to two, or less than two. We then assumed a reduction of the probabilities and their averages through a 50% or 30% reduction of non-HDL cholesterol, on the basis of the equation 1–exp(−0·249 + [number of years of treatment −5]×[−0·0152]) for the expected proportional risk reduction per mmol/L in LDL cholesterol for a given treatment duration. It was assumed that this equation was also valid for non-HDL cholesterol. Following the methods described by Austin, these average probabilities can then be used to estimate risk differences, numbers needed to treat, and differences in relative risk.

To obtain predicted probabilities from the cause-specific Cox models for each combination of sex and cohort, the respective cause-specific baseline hazard function was estimated by fitting a Weibull curve to age cohort, the respective cause-specific baseline hazard specific Cox models for each combination of sex and relative risk. Differences, numbers needed to treat, and differences in average probabilities can then be used to estimate risk through a 50% or 30% cholesterol category, age category, and according to

Risk estimates are calculated using a reduction of baseline risk by 30% or 50% (adapted from the 2018 American College of Cardiology/American Heart Association Guideline on the Management of Blood Cholesterol). On the basis of this reduction, we recalculated the probability of a cardiovascular disease event by the age of 75 years, as in our predictive model. The models and their averaged predictions were calculated in the derivation cohorts and also independently in the validation cohorts, and the discrepancy between the average predictions was compared using the root mean square error. This last step was repeated using the complete dataset, excluding each country in turn and also in the excluded country, thus generating country-specific root mean square errors. The final models and their predictions were computed in the complete dataset. Cumulative incidence curves for specific combinations of variables were produced on the basis of the cause-specific Cox models. Here, we also assumed a 50% reduction in non-HDL cholesterol for the included individuals. Applying this equation for the expected proportional risk reduction for a given treatment duration could yield a non-monotonic curve. Therefore,
the resulting curves were forced to remain constant in the event of a decrease.

To compare the discrimination of a model using non-HDL cholesterol with that of a model using LDL cholesterol, additional cause-specific Cox models were computed using age as the timescale, sex and cohort as strata, adjusting for cardiovascular risk factors (smoking status, diabetes, BMI, systolic blood pressure, and antihypertensive medication), and including a sex–lipid variable interaction. All continuous variables, except age, were modelled using cubic splines. Tenfold cross-validation was used when computing the C-indices. The main analyses for non-HDL cholesterol were repeated for LDL cholesterol. All statistical methods were implemented in R statistical software, version 3.6.1.

Role of the funding source
The funder of the study had no role in study design, data collection, data analysis, data interpretation, or writing of the report. The corresponding author (SB) and the lead statistician (FO) had full access to all the data in the study, and the corresponding author had final responsibility for the decision to submit for publication.

Results
We identified 38 cohorts reporting on 398 846 individuals with 560 4735 person-years of follow-up data available for analysis, of whom 184 055 (48·7%) were women (tables 1 and 2, appendix pp 19–27). Examination years ranged from 1970 to 2013 (years for each cohort are given in the appendix pp 19–27). We included 199 415 individuals in the derivation cohort (91 786 [48·4%] women) and 199 431 (92 269 [49·1%] women) in the validation cohort.

The median age of participants was 51·0 years (IQR 40·7–59·7). 12 311 (4·9%) were receiving lipid-lowering therapy. Median follow-up time was 13·5 years to non-HDL cholesterol concentration categories in women and men. Death from non-cardiovascular-disease causes was used as competing risk. p values are given for Gray’s test comparing cumulative incidence curves.

The median age of participants was 51-0 years (IQR 40-7–59-7). 12 311 (4-9%) were receiving lipid-lowering therapy. Median follow-up time was 13-5 years (IQR 7-0–20-1) with a maximum follow-up of 43-6 years.
10 years, even in the lowest cholesterol categories (figure 1). The cumulative incidence curves showed a stepwise increase of cardiovascular disease events across increasing concentrations of non-HDL cholesterol. 30-year cardiovascular disease event rates were approximately three-to-four times higher in women and men in the highest non-HDL cholesterol category (≥5.7 mmol/L) than those in the lowest category (<2.6 mmol/L; 3253 [33.7%] vs 262 [7.7%] in women and 7689 [43.6%] vs 375 [12.8%] in men). Multivariable analyses confirmed a strong continuous association of concentrations of non-HDL cholesterol with the sex-specific cardiovascular disease risk. The lowest hazard for cardiovascular disease was found in women and men with the lowest non-HDL cholesterol concentrations, whereas there was a continuous and linear increase for higher non-HDL cholesterol concentrations (figure 2; the reference value [HR 1.0] was set as 2.6 mmol/L non-HDL cholesterol in both women and men). The hazard of cardiovascular disease increased progressively with higher non-HDL cholesterol categories compared with the reference category (figure 3). This association remained consistent in a sensitivity analysis using a random-effects multivariate meta-analysis (appendix p 31). The steepest increase of the relative hazard associated with non-HDL cholesterol was found in individuals younger than 45 years at baseline.

**Figure 2:** Sex-specific continuous association of non-HDL cholesterol and cardiovascular disease

Sex-specific linear association of non-HDL cholesterol and cardiovascular disease risk (winsorised at 1.6 and 8.5 mmol/L). The Cox model used is adjusted for age, sex, study cohort, smoking, diabetes, body-mass index, systolic blood pressure, and antihypertensive medication. Non-HDL cholesterol was modelled using cubic splines. An interaction between sex and non-HDL cholesterol was included in the model. Median follow-up was 12.8 (IQR 7.5–18.4) years.

**Figure 3:** Age-specific and sex-specific association of non-HDL cholesterol and cardiovascular disease

Lifetime sex-specific HRs for fatal and non-fatal cardiovascular disease (reference non-HDL cholesterol <2.6 mmol/L) in the overall cohort and according to three age categories (p=0.001) for the interaction of age and non-HDL cholesterol categories in women and in men. The Cox regression models were adjusted for age at baseline, sex, study cohort, smoking, diabetes, body-mass index, systolic blood pressure, and antihypertensive medication. HR=hazard ratio.
baseline (maximum HR 4.3, 95% CI 3.0–6.1 in women and 4.6–6.3, 3.3–6.5 in men, for non-HDL cholesterol ≥5.7 mmol/L). Within the older groups, the association of non-HDL cholesterol with incident cardiovascular disease was attenuated but still detectable in individuals aged 60 years and older (HR 1.4, 95% CI 1.1–1.7 in women and 1.8, 1.5–2.2 in men, for non-HDL cholesterol ≥5.7 mmol/L; p<0.001 for the interaction of age and non-HDL cholesterol categories in women and in men; figure 3). Similar results were found in sensitivity analyses excluding individuals with diabetes (appendix p 38) and those receiving lipid-lowering therapy (appendix p 39) at baseline. The association of non-HDL cholesterol and cardiovascular disease remained consistent after additionally adjusting for HDL cholesterol (appendix p 40).

We found similar results for the association of blood concentrations of LDL cholesterol and incident cardiovascular disease to non-HDL cholesterol (appendix p 32). Cardiovascular disease event rates increased stepwise with increasing LDL cholesterol categories (appendix p 41). Multivariable analyses showed a linear association of LDL cholesterol with cardiovascular disease on the log-hazard scale, with the lowest relative risk for cardiovascular disease in the individuals with the lowest LDL cholesterol concentrations (appendix p 42). The strongest association was detected in individuals younger than 45 years and was attenuated in individuals aged 60 years and older (p<0.001 for the interaction of age and LDL cholesterol category in women and in men; appendix p 43).

To estimate the long-term probability of a cardiovascular disease event associated with non-HDL cholesterol, we established a model for cardiovascular disease risk up to the age of 75 years on the basis of a derivation–validation approach. Besides the non-HDL cholesterol category, sex, age, and the baseline number (≤1 vs ≥2) of classical modifiable cardiovascular risk factors (ie, arterial hypertension, diabetes, obesity, and smoking) were incorporated into this model (figure 4). For example, women with non-HDL cholesterol concentrations between 3.7 and 4.8 mmol/L, younger than 45 years, and with at least two additional cardiovascular risk factors had a 15.6% (95% CI 14.9–16.6) probability of experiencing a non-fatal or fatal cardiovascular disease event by the age of 75 years (28.8%, 28.1–29.5 in men with the same characteristics). The highest long-term risk of cardiovascular disease was seen in individuals younger than 45 years of age. The risk predictions obtained from this model were highly comparable in the derivation and validation datasets with a root mean square error lower than 1% for the estimated probabilities of cardiovascular disease (appendix p 44). Country-specific root mean square errors, as well as C-indices and smooth calibration curves specific for sex and follow-up time are provided in the appendix (pp 33, 34, 45). Further sensitivity analyses using time-dependent HRs did not show relevant differences in the prediction of cardiovascular disease by the age of 75 years (appendix p 46).

We calculated the optimally achievable risk reduction for cardiovascular disease by the age of 75 years...
assuming a 50% reduction of non-HDL cholesterol (figure 4). In the population with non-HDL cholesterol of 3·7–4·8 mmol/L, younger than 45 years, and with at least two risk factors, the long-term risk of cardiovascular disease could hypothetically be reduced from 15·6% (95% CI 14·9–16·6) to 3·6% (3·4–3·8) in women and from 28·8% (28·1–29·5) to 6·4% (6·3–6·6) in men. Further, corresponding numbers of patients needed to treat over the lifespan up to the age of 75 years, and achievable relative risk reduction (RRR) are shown in table 3. For example, in the same individuals, the number needed to treat to reduce one cardiovascular disease event over the lifespan by the age of 75 years was 8·3 in women and 4·5 in men. The model used is assuming a hypothetical 50% reduction of non-HDL cholesterol. Further, we modelled the potentially achievable long-term cardiovascular disease risk, assuming a 50% reduction of non-HDL cholesterol.

### Discussion

Using individual-level data from the Multinational Cardiovascular Risk Consortium of individuals without prevalent cardiovascular disease, we characterised the age-specific and sex-specific long-term association of non-HDL cholesterol with cardiovascular disease. On the basis of this association, we derived and validated a tool specific for age, sex, and cardiovascular risk factors to assess the individual long-term probability of cardiovascular disease by the age of 75 years associated with non-HDL cholesterol. Further, we modelled the potentially achievable long-term cardiovascular disease risk, assuming a 50% reduction of non-HDL cholesterol.

The proatherogenic effects of apoB-containing lipoproteins are largely based on the development and progression of atherosclerotic plaques and by accumulation of these lipoproteins within the arterial intima.\(^\text{30}\) Although the measurement of LDL cholesterol can be influenced by increased triglycerides\(^\text{22}\) in cardiovascular risk assessment, an appropriate way to estimate the amount of apoB-containing lipoproteins is the determination of non-HDL cholesterol.\(^\text{12,24,25}\) Therefore, our main analyses are based on non-HDL cholesterol. In line with previous studies, we found a comparable prognostic relevance for non-HDL cholesterol and LDL cholesterol.\(^\text{12,24,25}\) A large body of evidence including inherited disorders,\(^\text{27}\) genetic\(^\text{28,29}\) and epidemiological studies,\(^\text{10,11}\) and clinical trials\(^\text{7,9}\) leads to the hypothesis that reducing apoB-containing lipoproteins, regardless of the method, should yield a corresponding reduction in cardiovascular events. To intervene early and intensively during the lifespan might even lead to regression of early manifestation of atherosclerosis.\(^\text{10}\) Epidemiological studies have shown the association between cholesterol concentration and cardiovascular mortality during long-term follow-up.\(^\text{7–9}\) Furthermore, repeated non-HDL cholesterol testing in 2516 individuals of the Framingham Offspring Study suggested that stable concentrations over the lifetime\(^\text{10}\) might represent increased risk of cardiovascular disease. Clinical trials, however, mostly address older (age >60 years for most studies) individuals with prevalent cardiovascular disease or at markedly increased cardiovascular risk and report observations over short follow-up intervals, varying between 2 and 7 years.\(^\text{7–9}\)

Considerable uncertainty exists about the extent to which slightly increased or apparently normal cholesterol concentrations affect lifetime cardiovascular risk and about which thresholds should be used to merit a treatment recommendation, particularly in young people. Our study extends current knowledge because it suggests that increasing concentrations of non-HDL cholesterol predict long-term cardiovascular risk, particularly in cases of modest increase at a young age. Only a few studies have reported the age-dependent association of lipids and cardiovascular disease with focus on young individuals.\(^\text{11,13}\) In our analyses, individuals younger than 45 years showed the strongest HRs for the
association of blood lipids with the incidence of cardiovas-
cular disease during long-term follow-up. This effect is
probably related to the following considerations: (1) the
effect of lifetime accumulation of proatherogenic lipids
in the vulnerable period of the fourth and fifth decades
of life,10,11 (2) the shorter timespan that individuals aged
60 years and older have to achieve the cardiovascular
disease endpoint due to their age itself, and (3) the fact
that people with prevalent cardiovascular disease are not
included in our study and thus individuals who reach
the age of 60 without prevalent cardiovascular disease are
a population enriched with protective characteristics
against the proatherogenic effect of cholesterol. However,
although there is a weaker association of non-HDL
cholesterol with cardiovascular disease in older indi-
viduals (>60 years), these people are at higher absolute
risk than young individuals, and so the more modest
relative risks represent large absolute risk differences
across non-HDL cholesterol categories.12

The risk circle tool displays the individual lifetime risk
of non-fatal and fatal cardiovascular disease by the age
of 75 years that is associated with non-HDL cholesterol
and is specific for the burden of age, sex, and risk factor.
It also estimates the corresponding risk reduction from
lipid-lowering therapies. The risk scores currently used
for decision making about lipid-lowering intervention
assess only the 10-year cardiovascular risk10,11 and there-
fore underestimate lifetime risk, particularly in young
individuals.11,12 Pencina and colleagues11 estimated the
30-year risk of cardiovascular disease in the Framingham
Offspring cohort and showed 30-year risks that were up
to ten times higher than the projected 10-year risks in
young people.11 Therefore, in young individuals,
the long-term risk of cardiovascular disease should
constitute the basis of primary prevention. Since the risk
factor burden is incorporated as a categorical variable in
our model, we cannot provide a weighting among the
different risk factors or a quantitative consideration of
every single risk factor as do other risk models. Instead,
because of its simplicity, our risk circle tool provides an
opportunity to estimate lifetime risks based on non-
HDL cholesterol in an accessible and easily understood
way that can improve physician–patient communication
about preventive strategies in clinical practice. The
modelled risk reduction is calculated hypothetically and
is based on a 50% lipid reduction leading to a decrease
of the cumulative burden of non-HDL cholesterol. Since
the calculated numbers needed to treat are based on
different timeframes for different age ranges, they have
to be compared carefully between the different age
categories. They have to be interpreted individually for
a certain spectrum of ages, sexes, and risk factors to
guide decision making on any lipid-lowering inter-
vention. Furthermore, numbers needed to treat might be
higher than those we showed in low-risk countries
with lower than average cardiovascular disease incidence
and mortality. The following limitations merit considera-
tion. First, due to the population-based longitudinal study
design, the endpoint information is mainly based on medical
reports or local registers and we cannot exclude mis-
classification of endpoints for some of the observed
effects. However, endpoint information was harmoni-
ised on the basis of individual data according to the
specification of the MORGAM study, and multivariable
analyses were adjusted for potential confounders.
Therefore, the best achievable data quality can be
assumed for the current analyses. Second, only baseline
data of blood lipids were available for current analyses.
No information about dynamic changes during follow-
up or about the initiation of lipid-lowering therapy,
particularly in those individuals with very high blood
concentrations, could be considered.13 However, non-
HDL cholesterol concentrations in young individuals
are generally stable over the 30-year life course,14 and
not adjusting for changes in risk factors was shown to
have little bearing on lifetime cardiovascular disease
risk.13 Third, our cohorts largely included individuals
with European ancestry from high-income countries in
Europe, North America, and Australia. The general-
siability of our findings to other regions or individuals
from other racial and ethnic groups is unknown. Finally,
the therapeutic benefit of lipid-lowering intervention
investigated in our study is based on a hypothetical
model that assumes a stable reduction of non-HDL
cholesterol. We recognise that making such an assump-
tion posits that treatment effects are sustained over
a much longer term than has been studied in clinical
trials and that real-world benefits of taking statins are
probably lower than those shown in trials because of
adherence and side-effects.16 However, since clinical
trials investigating the benefit of lipid-lowering therapy
in individuals younger than 45 years during a follow-
up of 30 years are not available, our study provides
unique insights into the benefits of a potential early
intervention in primary prevention. We agree with
Hippisley-Cox and colleagues17 that future research
needs to address many related questions, including
whether intervention in young people with a high
lifetime risk but low 10-year risk would yield greater
benefits than late intervention.

Contributors
SB, FJB, and CW conceived the idea for the study, interpreted the data,
and wrote the paper with input from all authors. FO and KK harmonised,
analysed, and interpreted the data. VS, FK, SSa, BT, SG, PB, HT-P, MM,
Li, GV, GG, EBM, SSo, AL, HB, PA, JF, AT, VPN, WD, OM, K-HJ, DML,
JES, DBP, LAS, MK, RSV, RPFD, SGW, UR, SSB, JAdl, TO, and Ul
provided the data and were involved in the data interpretation and
discussion of the results. All authors provided critical feedback and
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Declaration of interests

FJB and CW report personal grants from the ARSEP Cardiovascular grant award, Pfizer, during the conduct of the study. CW reports personal fees from Amgen and Sanofi, outside the submitted work. V5 reports personal fees from Novo Nordisk and Sanofi and grants from Bayer, outside the submitted work. SSö reports personal fees from Actelion and Amgen, from Novo Nordisk and Sanofi, and grants and personal fees from Total, Genoscreen, and Fondation Alzheimer, outside the submitted work. JF reports personal fees from Merck, Amgen, Servier, and Sanofi, outside the submitted work. JES reports grants from the Australian Commonwealth Department of Health and Aged Care, Abbott Australasia, Alphapharm, AstraZeneca, Aventis Pharma, Bristol-Myers Squibb, Eli Lilly, GlaxoSmithKline, Janssen-Cilag, Merck Lipha, Merck, Novartis, Novo Nordisk, Pharmacia and Upjohn, Pfizer, Sanofi Synthelabo, Servier Laboratories, the Australian Kidney Foundation, and Diabetes Australia, during the conduct of the study; and personal fees from AstraZeneca, Mylan, Boehringer Ingelheim, Sanofi, Merck, Novo Nordisk, Eli Lilly, and Abbott Laboratories, outside the submitted work. RSV is supported in part by the Evans Medical Foundation and the Jay and Louis Coffman Endowment from the Department of Medicine (Boston University School of Medicine, Boston, MA, USA). SSö reports grants from the US National Institutes of Health, outside the submitted work. JAdL reports personal fees from Amgen, Regeneron, and Esperion, outside the submitted work. TO reports grants and personal fees from Roche, Abbott, and Novartis, personal fees from Siemens and Bayer, and grants from Singulys and Somalogic, outside the submitted work. KK reports grants from the EU and the UK Medical Research Council, during the conduct of the study. UL reports personal fees from The Medicines Company, Amgen, Sanofi, Berlin Chemiker, Abbott Laboratories, AstraZeneca, and Novartis, and grants and personal fees from Bayer, outside the submitted work. SB reports grants and personal fees from Abbott Laboratories, Bayer, Siemens, and ThermoFisher Scientific, grants from Singulys, and personal fees from AstraZeneca, Amgen, Medtronic, Pfizer, and Roche, outside the submitted work. All other authors declare no competing interests.

Acknowledgments

This Article was prepared using data from the US National Heart, Lung, and Blood Institute Biologic Specimen and Data Repository Information Coordinating Center for the following cohorts: Cardiovascular Health Study, Atherosclerosis Risk in Communities Study, Jackson Heart Study, Multi-Ethnic Study of Atherosclerosis, and Framingham Heart Study. The Biomarker CaRE Project is funded by the EU Seventh Framework Programme (FP7/2007–2013) under grant agreement HEALTH-F2–2011–278913. The activities of the MORGM data centre have been also sustained by funding from the 9th EU Framework Programme since 1998 (EU FP7 project CHANCES; HEALTH–F3–2010–242244). A part of the biomarker determinations in the population cohorts was funded by the UK Medical Research Council (G0601461, 89083; Biomarkers in the MORGM Populations). Additional funding was provided by the German Centre for Cardiovascular Research (81Z170101). Further detailed funding information is provided in the appendix (pp 48–50).

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