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Residual cardiovascular risk in individuals on lipid-lowering treatment: quantifying absolute and relative risk in the community

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ABSTRACT

Objective The residual cardiovascular disease (CVD) risk in individuals on long-term lipid-lowering treatment (LLT) in the general population is not well described.

Methods We estimated absolute CVD risks by age and sex for different categories of low-density lipoprotein cholesterol (LDL-C) levels, stratified by LLT status, and assessed subclinical carotid atherosclerosis in 3012 Framingham Study participants (mean age, 58.4 years; 55% women) free of CVD. Individuals were categorised into five groups: (1) LDL-C <100 mg/dL without LLT; (2) LDL-C ≥100 mg/dL to <130 mg/dL without LLT; (3) LDL-C <130 mg/dL on LLT; (4) LDL-C ≥130 mg/dL without LLT; and (5) LDL-C ≥130 mg/dL on LLT.

Results Individuals in groups 3–5 had significantly more carotid atherosclerosis compared with group 1. During follow-up (median, 13.7 years), 548 CVD events occurred. Individuals on LLT (groups 3 and 5) had substantial residual CVD risk (26.7 (95% CI 19.5 to 34.0) and 24.1 (95% CI 16.2 to 31.9) per 1000 person-years, respectively), representing approximately three times the risk for untreated individuals with LDL <100 mg/dL (group 1: 9.0 (95% CI 6.8 to 11.3) per 1000 person-years). Absolute CVD risks rose with age and were slightly greater in men than in women. After adjustment for traditional risk factors, groups 3–5 displayed increased hazards for CVD (HR=1.47, 1.42 and 1.54, respectively) compared with group 1. Further adjustment for carotid atherosclerosis modestly attenuated these results.

Conclusions There is substantial residual CVD risk in individuals on LLT, compared with participants with optimal LDL-C (<100 mg/dL), even when LDL-C levels <130 mg/dL are reached.

Key questions

What is already known about this subject?

- Lipid-lowering treatment (LLT) reduces cardiovascular disease (CVD) risk and is, therefore, a fundamental pillar of preventive cardiology.
- The residual CVD risk in individuals on long-term LLT in the general population is not well described.

What does this study add?

- In a large community-based sample, we quantified subclinical disease burden at the carotid arteries and provide absolute CVD risk estimates for different categories of low-density lipoprotein cholesterol (LDL-C) levels, representing different treatment intensities of LLT.
- Individuals on LLT had greater subclinical disease burden and displayed a substantial residual CVD risk, approximately tripling the risk of untreated individuals with LDL-C <100 mg/dL.
- This residual risk in individuals on LLT in the general population is partly explained by an adverse profile of other CVD risk factors and in part by carotid subclinical atherosclerosis in these people.

How might this impact on clinical practice?

- Our observations emphasise the necessity to closely monitor all standard CVD risk factors in individuals on LLT, even when LDL levels are lowered effectively.

INTRODUCTION

Based on mounting evidence from randomised controlled trials in both primary and secondary prevention settings,[1–4] lipid-lowering treatment (LLT; pharmacological and lifestyle) is a fundamental pillar of preventive cardiology.

Relatively little evidence exists regarding the remaining absolute disease risk in individuals on LLT, and there is currently an active debate regarding the broad application of statins.[5–7] While most people would agree that there *is* a residual risk, the magnitude of this risk and its variation with age, sex and lipid level categories on LLT (representing different intensities of LLT) is not well quantified, particularly not in the community. This is because most randomised controlled trials have strict inclusion and exclusion criteria and might, therefore, not ideally mirror the situation in the community. Furthermore, the extent to what the residual cardiovascular risk is attributable to subclinical vascular disease

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or to alterations in the neurohormonal or inflammatory milieu is also not well described. A detailed quantification of the residual cardiovascular disease (CVD) risk in the community and a better understanding of its biology would be essential to lower the burden of CVD. Part of the mechanisms by which dyslipidaemia promotes CVD risk is by enhancing the substrate of subclinical atherosclerosis and by altering the neurohormonal and inflammatory milieu.

We hypothesised that (1) although LLT lowers CVD risk, individuals on LLT experience substantial residual CVD risk that exceeds the risk observed in untreated individuals with similar levels of low-density lipoprotein cholesterol (LDL-C), and (2) this residual CVD risk is in part due to the burden of subclinical atherosclerosis and due to systemic inflammation and neurohormonal activation.

Accordingly, we assessed (1) the burden of subclinical disease at the carotid arteries cross-sectionally and (2) the risk for incident CVD prospectively, associated with different levels of LDL-C in a large community-based sample, stratified by the use of LLT.

METHODS

Study sample

The present analyses were based on data from attendees at the sixth examination cycle of the Framingham Offspring cohort (1995–1998).[8] A total of 520 individuals were excluded for reasons detailed in the online supplementary material, resulting in a sample size of n=3012. For 95 individuals, information regarding the subclinical disease burden of the carotid artery (detailed below) was missing, resulting in a sample size of n=2917 for the analyses related to subclinical carotid disease burden. Circulating C reactive protein (CRP) concentrations and plasma levels of B-type natriuretic peptide (BNP) were available in 2811 participants. All participants provided written informed consent.

Laboratory measurements

Lipid traits, including blood levels of total cholesterol, high-density lipoprotein (HDL) cholesterol and triglycerides, were measured in fasting participants using standardised assays. LDL-C was calculated using the Friedewald formula in individuals with triglyceride levels <400 mg/dL.[9] Measurements of CRP and BNP are described in the online supplementary material.

Ultrasound of the carotid artery

Ultrasonographic measurements of the carotid arteries on both sides were performed using a 7.5 MHz transducer and a 5.0 MHz transducer for images of the common and of the internal carotid arteries, respectively, on a Toshiba SSH-140 A machine, following a standardised protocol.[10–12] More details regarding the ultrasound measurements are provided in the online supplementary material.

'Carotid ultrasound abnormality' was defined as (1) increased (≥80th sex-specific percentile) intima media thickness (IMT), a combined standardised measure including information from the internal and common carotid arteries; (2) an extreme increase of the common carotid IMT ≥1 mm; or (3) significant stenosis (≥25% narrowing) of the common or internal carotid arteries, consistent with prior publications.[13 14] We chose this combined parameter in order to reflect both alterations in IMT (a measure of overall atherosclerotic plaque burden) and the presence of arterial stenoses (a measure of obstructive atherosclerosis).

Adjudication of incident CVD events

Framingham participants are under regular surveillance for new-onset CVD events. More details regarding the adjudication of events are provided in the online supplementary material.

Statistical analyses

Based on the intake of lipid-lowering medications and the blood LDL levels determined at the sixth examination cycle, participants were assigned to one of five mutually exclusive groups.

1. usual LDL-C <100 mg/dL, not on LLT
2. usual LDL-C ≥100 mg/dL to <130 mg/dL, not on LLT
3. treated LDL-C <130 mg/dL on LLT
4. usual LDL-C ≥130 mg/dL, not on LLT
5. treated LDL-C ≥130 mg/dL on LLT.

There were very few individuals (n=58) with LDL-C <100 mg/dL on LLT; therefore no separate category was created for these individuals. We chose these cut points taking into account different target lipid levels and the presence versus absence of LLT, thereby ensuring that the strata yielded an adequate number of events to enable analyses with reasonable statistical power.

To quantify the baseline risks in the different LDL groups, we calculated the 10-year probability of a CVD event[15] for each participant and displayed these event probabilities as boxplots according to LDL group. Treated individuals (groups 3 and 5) were compared versus untreated individuals (groups 1, 2 and 4) with respect to their 10-year CVD event probabilities using analysis of variance.

Subclinical disease burden at the carotid arteries by LDL group

First, we calculated the ORs for the binary trait 'carotid ultrasound abnormality' (defined as detailed above, including information on IMT and on stenoses) for each LDL group, using group 1 as the referent, in age-and sex-adjusted as well as multivariable-adjusted logistic regression models. The multivariable models included age, sex, systolic blood pressure, antihypertensive medication, smoking and diabetes mellitus. Second, we compared the odds for 'carotid ultrasound abnormality' in individuals on LLT (group 5 and group 3) with the odds for 'carotid ultrasound abnormality' in untreated

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individuals with similar LDL levels (group 4 and groups 1+2 combined, respectively) using age- and sex-adjusted as well as multivariable-adjusted logistic regression models. We focused on 'carotid ultrasound abnormality' as the primary subclinical disease trait because subclinical alterations of the carotid artery are reported to be strongly associated with increased LDL levels[16] and the selected trait 'carotid ultrasound abnormality' includes both information from IMT and of relevant stenoses of the carotid arteries.[13 14]

Association of LDL category with incident CVD

We performed several analyses to describe the residual cardiovascular risk associated with different levels of LDL-C, stratified by LLT status, age group and sex. First, the unadjusted survival free of CVD by 'LDL category' was graphically displayed using a Kaplan-Meier curve. Second, we report absolute disease risks (events per 1000 person-years) for each LDL category for the entire sample, and stratified by sex and by age group (<60 and ≥60 years). Third, Cox proportional hazard models were used to relate each LDL group to incident CVD, thereby adjusting for age, sex, systolic blood pressure, antihypertensive medication, smoking and diabetes mellitus. In secondary analyses, we additionally adjusted for presence versus absence of carotid ultrasound abnormality (defined as above) and for circulating CRP and BNP levels. In these models, we compared LDL categories 2–5 (each category considered separately) with LDL category 1 (usual LDL-C <100 mg/dL, no LLT). Fourth, we specifically compared treated with untreated individuals with similar LDL levels. Thus, we calculated multivariable-adjusted HR for incident CVD for group 5 (using group 4 as the referent) and for group 3 (using groups 1+2 combined as the referent).

RESULTS

The baseline characteristics of the overall study sample and stratified by LDL group and LLT status are displayed in table 1. Overall, 302 participants (groups 3 and 5; 10% of the overall sample) were on lipid-lowering medications, and 170 treated individuals (group 3; 56% of those on treatment) were controlled to a blood LDL level below 130 mg/dL. Clinical, biochemical and ultrasound measures of the carotid arteries in the sample with available carotid ultrasound information (n=2917) are provided in online supplementary table 1, and the CVD baseline risk (10-year probability of a CVD event)[15] by LDL group is displayed in online supplementary figure 1. The 10-year probabilities for a CVD event in treated individuals (groups 3 and 5) were statistically significantly greater than in untreated individuals (groups 1, 2 and 4; P<0.0001). More detailed information regarding individuals on LLT, including information on the average LDL-C level across exams 1–6 and the average duration of LLT, is provided in online supplementary table 2. As expected, individuals in group 5 (LDL ≥130 mg/dL on treatment) had the highest LDL levels across exams 1–6 and across all exams before LLT was initiated (175 mg/dL; online supplementary table 2). In group 3, more than 80% of participants were on statins, whereas the proportion of statin users was smaller (~64%) in group 5.

Association of LDL group with subclinical disease in the carotid arteries

In age- and sex-adjusted and in multivariable-adjusted models, individuals on lipid-lowering medication (groups 3 and 5) and individuals with an untreated LDL above 130 mg/dL (group 4) had significantly higher ORs for carotid ultrasound abnormalities as compared with individuals in the referent group (usual LDL <100 mg/dL; figure 1A,B). Furthermore, individuals on treatment who reach LDL levels <130 mg/dL (group 3) had statistically significantly increased odds for subclinical carotid ultrasound abnormalities as compared with untreated individuals with similar LDL levels (groups 1+2 combined; OR: 1.95, 95% CI 1.34 to 2.84, P=0.0005). No statistically significant differences between treated individuals with LDL ≥130 (group 5) and untreated individuals with comparable LDL levels (group 4; OR: 1.04, 95% CI 0.69 to 1.57, P=0.86) could be observed.

Residual risk for incident CVD in individuals on LLT

During a median follow-up of 13.7 years (Q1=11.5 years, Q3=14.9 years), 548 (255 women) out of 3012 participants developed an incident CVD event. The Kaplan-Meier curves for survival free of CVD by LDL groups are displayed in figure 2 (log-rank P<0.0001). Table 2 displays the absolute disease risks by LDL group for the entire sample and stratified by sex and age group (<60 and ≥60 years). Individuals on LLT (group 3 and group 5) had substantial residual cardiovascular risk for incident CVD that is, in the overall sample, approximately three times the absolute disease risk observed in group 1 (LDL below 100 mg/dL, no LLT; table 2 'crude event rates'). In individuals below 60 years of age, the absolute disease risks in groups 3 and 5 (on LLT) were almost sixfold and fourfold, respectively, the risk in the referent group (table 2). In the age group ≥60 years, the absolute cardiovascular risk rose for groups on LLT, but also the absolute risk in the referent group 1 increased, so that the relative risks in groups 2–5, compared with group 1, diminished with age. Furthermore, absolute CVD risks were slightly higher in men as compared with women for most LDL groups (table 2).

Impact of adjusting for potential covariates, subclinical atherosclerosis of the carotid artery and biomarker levels on residual CVD risk

Compared with reference group 1, LDL groups 3–5 displayed a statistically significantly increased relative hazard for incident CVD in multivariable-adjusted models (figure 3). Adjustment for subclinical atherosclerosis in the carotid arteries only modestly reduced

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Table 1 Baseline characteristics of the study sample

LDL (in mg/dL) group	Overall sample (n=3012)	Group 1 LDL <100 No treatment (n=546)	Group 2 100≤ LDL <130 No treatment (n=931)	Group 3 LDL <130 On treatment (n=170)	Group 4 LDL ≥130 No treatment (n=1233)	Group 5 LDL ≥130 On treatment (n=132)
Clinical and biochemical features						
Age, years; mean±SD	58.4±9.6	56.2±9.7	57.9±9.7	63.7±8.5	58.4±9.3	62.7±7.6
Women, n (%)	1671 (55.5)	348 (63.7)	506 (54.4)	75 (44.1)	667 (54.1)	75 (56.8)
LDL cholesterol, mg/dL; mean±SD	127.9±33.7	83.5±13.1	115.7±8.4	104.1±18.9	156.9±23.1	157.4±23.8
Total cholesterol, mg/dL; mean±SD	205.8±37.3	163.0±22.6	192.7±18.4	182.2±26.1	234.6±27.6	238.2±30.6
HDL cholesterol, mg/dL; mean±SD	52.1±16.0	58.0±19.9	52.3±16.1	45.0±13.3	50.8±13.9	47.4±11.5
Lipid-lowering treatment, n (%)	302 (10.0)	0	0	170 (100.0)	0	132 (100.0)
Statin treatment, n (%)	224 (7.4)	0	0	140 (82.4)	0	84 (63.6)
Systolic BP, mm Hg; mean±SD	127.7±18.6	123.1±18.2	126.6±18.3	136.8±19.5	128.8±18.3	133.4±18.0
Diastolic BP, mm Hg; mean±SD	75.6±9.4	73.6±9.6	75.2±9.2	76.4±9.2	76.5±9.5	77.7±9.2
Blood pressure-lowering treatment, n (%)	726 (24.1)	88 (16.1)	227 (24.4)	100 (58.8)	256 (20.8)	55 (41.7)
Body mass index, kg/m^2; mean±SD	27.8±5.2	26.6±5.2	27.8±5.4	29.4±4.9	28.1±5.1	28.7±4.5
Smoking, n (%)	437 (14.5)	75 (13.7)	125 (13.4)	14 (8.2)	201 (16.3)	22 (16.7)
Diabetes, n (%)	227 (7.5)	37 (6.8)	64 (6.9)	37 (21.8)	69 (5.6)	20 (15.2)
BNP, pg/mL, median (Q1, Q3)	7.7 (4.0, 16.8)	8.6 (4.0, 19.6)	8.3 (4.0, 18.4)	10.3 (4.0, 22.5)	6.7 (4.0, 14.5)	8.8 (4.0, 16.5)
CRP, mg/L, median (Q1, Q3)	2.0 (0.9, 4.5)	1.8 (0.7, 5.0)	1.8 (0.8, 3.9)	2.4 (1.0, 5.0)	2.1 (1.0, 4.5)	2.8 (1.2, 4.8)

Sample size with available BNP levels, n=2997; sample size with available CRP levels, n=2909.
BNP, B-type natriuretic peptide; BP, blood pressure; CRP, C reactive protein; HDL, high-density lipoprotein; LDL, low-density lipoprotein.

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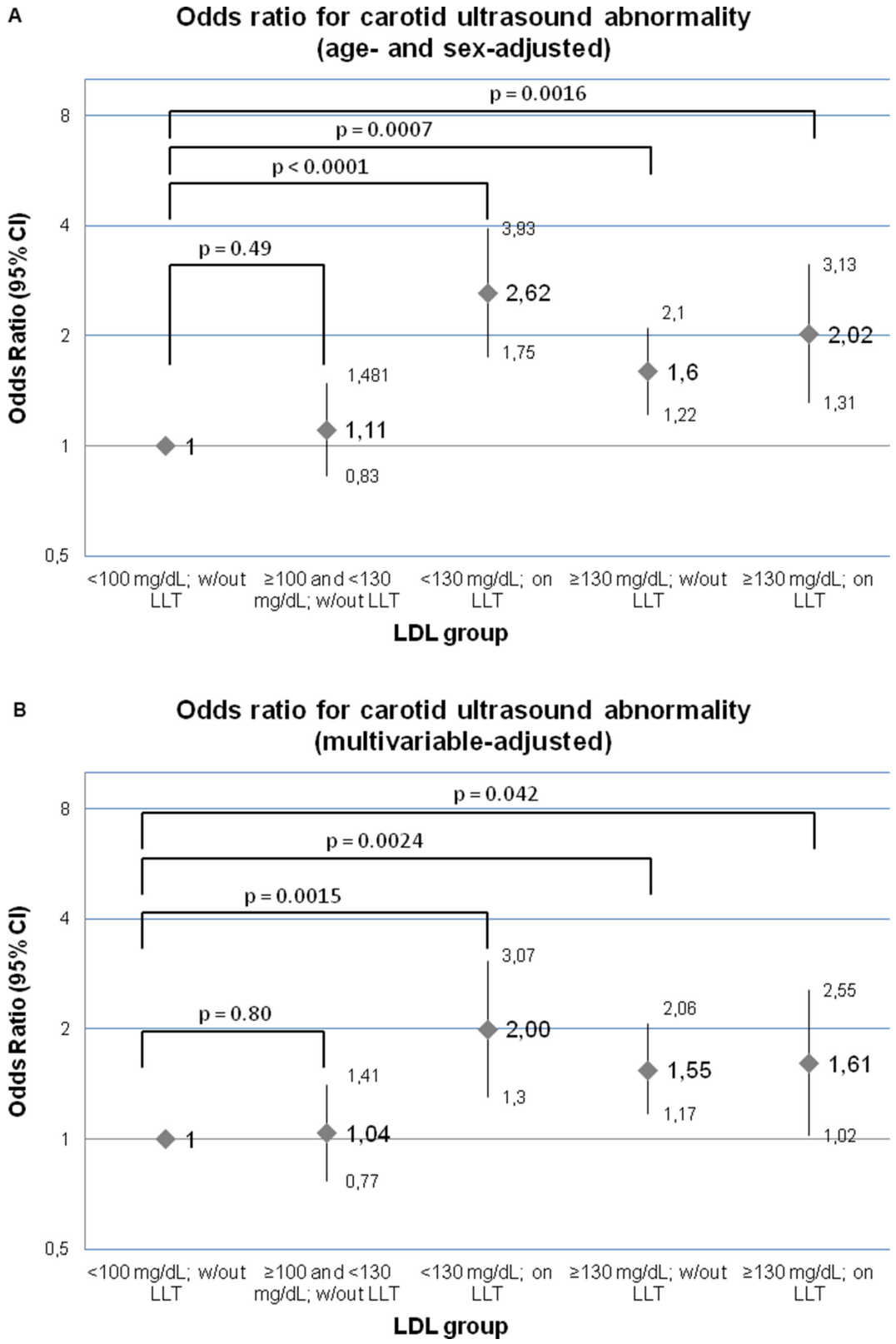
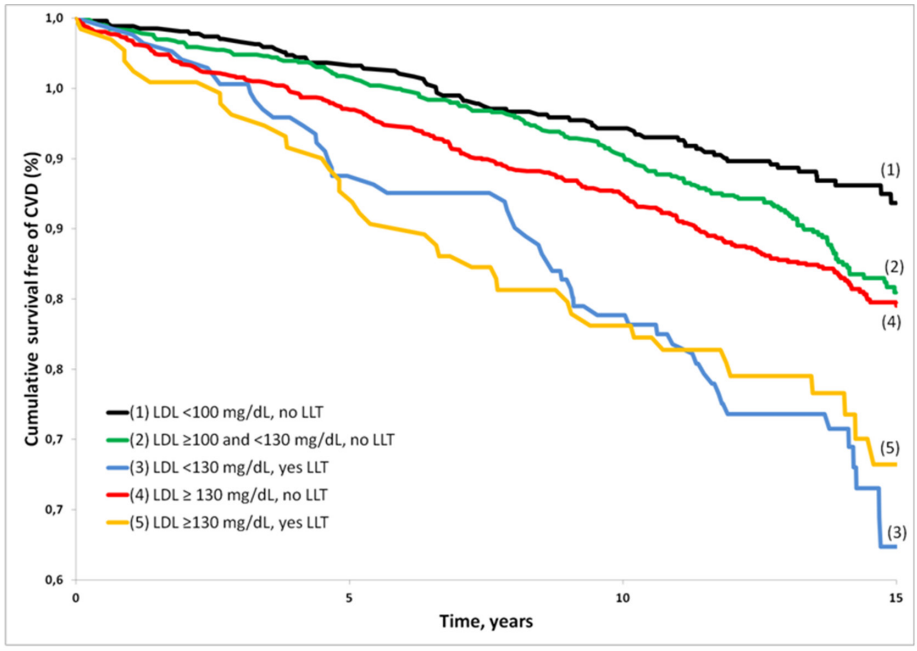


Figure 1 OR for the binary trait 'carotid ultrasound abnormality' in an age- and sex-adjusted model (A), as well as in a multivariable-adjusted model (B), including age, sex, systolic blood pressure, antihypertensive medication, smoking and diabetes mellitus. 'Carotid ultrasound abnormality' was defined as (1) increased (≥80th sex-specific percentile) carotid IMT, a combined standardised measure including information from the internal and common carotid arteries, (2) an extreme increase of the common carotid IMT ≥1 mm, or (3) significant stenosis (≥25% narrowing) of the common or internal carotid arteries, consistent with prior publications.[13 14] IMT, intima media thickness; LDL, low-density lipoprotein; LLT, lipid-lowering treatment; w/out, without.

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	Cumulative No. of Events/No. at Risk	Cumulative No. of Events/No. at Risk	Cumulative No. of Events/No. at Risk	Cumulative No. of Events/No. at Risk
LDL group 1	0/546	17/506	41/450	60/122
LDL group 2	0/931	38/863	87/768	148/214
LDL group 3	0/170	19/147	35/122	51/24
LDL group 4	0/1233	79/1129	151/998	222/307
LDL group 5	0/132	16/113	28/92	36/26

Figure 2 Survival free of CVD, stratified by LDL group. CVD, cardiovascular disease; LDL, low-density lipoprotein; LLT, lipid-lowering treatment.

these HRs by up to ~5%, and further adjustment for CRP and BNP had no relevant effect on the effect estimates (table 3).

Compared with untreated individuals with similar LDL (groups 1+2 combined), individuals on treatment who reach LDL <130 mg/dL (group 3) had statistically increased hazards for incident CVD in an age- and sex-adjusted model (HR: 1.44, 95% CI 1.06 to 1.96, P=0.02), but this difference was no longer statistically significant on multivariable adjustment (HR: 1.20, 95% CI 0.88 to 1.64, P=0.25). With respect to individuals with LDL ≥130 mg/dL, treated individuals (group 5) were not different from untreated individuals (group 4) with respect to new-onset CVD in age- and sex-adjusted (HR: 1.31, 95% CI 0.92 to 1.87, P=0.13) and in multivariable-adjusted (HR: 1.09, 95% CI 0.76 to 1.55, P=0.65) models.

DISCUSSION

In a large community-based sample, we provide absolute CVD risk estimates for the entire sample and stratified by age and by sex for different categories of LDL levels, representing different treatment intensities of LLT. Furthermore, we assessed potential mechanisms by which the residual risk could be mediated: subclinical carotid disease, neurohormonal activation or chronic inflammation. In addition, we describe the subclinical disease burden at the carotid arteries associated with different lipid levels in the community, stratified by treatment status.

Principal observations

First, subclinical disease burden and risk for incident CVD rose with increasing LDL category. Second, individuals who reach LDL levels <130 mg/dL on treatment (group 3) as well as individuals with LDL ≥130 mg/dL (groups 4 and 5) displayed greater carotid atherosclerotic disease burden, as compared with untreated individuals with usual LDL <100 mg/dL (group 1). Third, individuals on lipid-lowering medication (groups 5 and 3) had a substantial residual absolute risk for incident CVD that was about three times the risk observed in individuals with LDL <100 mg/dL who were not on treatment. Residual CVD risk rose substantially with age, and men had slightly higher absolute CVD risk than women in most LDL categories. Fourth, part of this increased risk was due to worse cardiovascular risk profiles in groups 3–5 (table 1). However, on adjustment for traditional CVD risk factors, individuals on LLT who reach LDL <130 mg/dL still had about 50% greater hazards for CVD as compared with the referent group. Similarly, groups 4 and 5 displayed statistically significant HRs for CVD (compared with group 1), taking potential confounders into account. Fifth, adjustment for subclinical carotid disease burden only moderately attenuated these HRs (by about 5%) consistent with partial mediation of the residual CVD risk by the burden of subclinical atherosclerosis. Further

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Table 2 Absolute CVD risk estimates in the largest possible sample (n=3012) and stratified by sex and age group (<60 and ≥60 years of age) for different categories of LDL levels, stratified by use of lipid-lowering medication

			Overall sample		Women		Men	
	LDL group	LDL treatment	Events (n)/At risk (n)	Crude CVD event rates (95% CI)*	Events (n)/At risk (n)	Crude CVD event rates (95% CI)*	Events (n)/At risk (n)	Crude CVD event rates (95% CI)*
1	<100 mg/dL	No	62/546	9.0 (6.8 to 11.3)	36/348	8.0 (5.4 to 10.6)	26/198	10.9 (6.7 to 15.1)
2	≥100 to <130 mg/dL	No	164/931	14.0 (11.9 to 16.2)	74/506	11.4 (8.8 to 14.1)	90/425	17.2 (13.7 to 20.8)
3	<130 mg/dL	Yes	52/170	26.7 (19.5 to 34.0)	24/75	26.8 (16.1 to 37.5)	28/95	26.7 (16.8 to 36.5)
4	≥130 mg/dL	No	234/1233	15.3 (13.3 to 17.2)	103/667	11.9 (9.6 to 14.3)	131/566	19.5 (16.2 to 22.9)
5	≥130 mg/dL	Yes	36/132	24.1 (16.2 to 31.9)	18/75	20.4 (10.9 to 29.8)	18/57	29.5 (15.9 to 43.1)

			<60 years of age		≥60 years of age	
	LDL group	LDL treatment	Events (n)/At risk (n)	Crude CVD event rates (95% CI)*	Events (n)/At risk (n)	Crude CVD event rates (95% CI)*
1	<100 mg/dL	No	15/359	3.2 (1.6 to 4.8)	47/187	22.2 (15.8 to 28.5)
2	≥100 to <130 mg/dL	No	63/551	8.6 (6.5 to 10.7)	101/380	23.2 (18.6 to 27.7)
3	<130 mg/dL	Yes	15/63	19.3 (9.6 to 29.2)	37/107	31.6 (21.4 to 41.8)
4	≥130 mg/dL	No	104/726	11.0 (8.9 to 13.1)	130/507	22.1 (18.3 to 25.9)
5	≥130 mg/dL	Yes	7/45	11.6 (3.0 to 20.2)	29/87	32.5 (20.7 to 44.3)

*Events per 1000 person-years.
CVD, cardiovascular disease; LDL, low-density lipoprotein.

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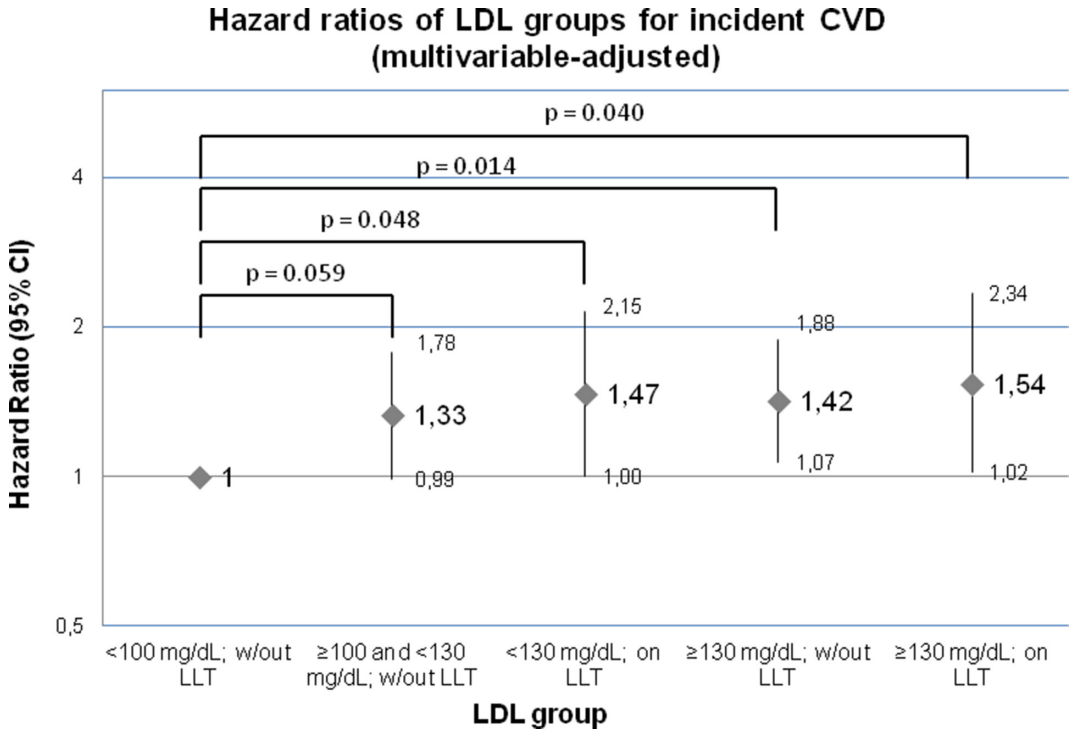


Figure 3 HRs for incident CVD by LDL group, adjusted for age, sex, systolic blood pressure, antihypertensive medication, smoking and diabetes mellitus. CVD, cardiovascular disease; LDL, low-density lipoprotein; LLT, lipid-lowering treatment; w/out, without.

adjustment for CRP and BNP did not significantly alter the risk estimates.

In the context of the published literature

The association between lipid measures and CVD risk is well established in the epidemiological and clinical literature.[17] Numerous clinical trials have demonstrated that lowering of LDL-C through pharmacological or lifestyle interventions reduces the risk for recurrent CVD and mortality in patients with established CVD[1 18] or in the primary prevention setting.[2 19 20] Furthermore, clinical trials have shown that statins—by lowering LDL—also improve subclinical CVD measures. A recent meta-analysis of 11 clinical trials reported beneficial effects of statins on carotid atherosclerosis.[21]

Despite clinically and statistically significant reductions in clinical and subclinical CVD risk in the LLT arm compared with the respective comparison groups, many clinical trials reported a relevant residual cardiovascular risk in individuals receiving LLT, with the magnitude of the residual risk depending on the exact patient sample, the duration of follow-up, the type (generation) and dosage of treatment used, the target LDL levels that have been reached, and the exact endpoint investigated in the study.[1 18 19 22–25] Clinical trials have relatively strict inclusion and exclusion criteria so that it is unclear whether the observations from clinical trials regarding the residual cardiovascular risk on LLT are applicable to the general population.[26]

In our unselected sample from the community, we confirm a substantial residual absolute CVD risk in individuals on LLT (group 3 and group 5). Individuals in these groups had about three times the risk of the referent group 1 (untreated LDL <100 mg/dL) in the overall sample. In individuals below 60 years of age, the ratios of the absolute disease risks in those on LLT and those in the referent group 1 were even more pronounced.

Furthermore, we explored the subclinical disease burden at the carotid arteries associated with various LDL levels, stratified by LLT status. Individuals on LLT and with greater LDL levels displayed statistically significantly greater odds for carotid atherosclerosis on ultrasound, including increased IMT and stenoses,[13] consistent with other observational studies.[27 28] In addition, we evaluated to what extent the increased hazards for CVD in LDL groups 3–5 (figure 3) were attributable to subclinical atherosclerosis at the carotid arteries and to systemic inflammation and neurohormonal activation. Our observations indicate that adjustment for subclinical carotid atherosclerosis only modestly attenuated the relative risk estimates, consistent with the concept that a modest proportion of the increased risk is attributable to subclinical atherosclerosis. Additional adjustment for established biomarkers of systemic inflammation (CRP) and neurohormonal activation (BNP) had no significant effect on the risk estimates.

Important drivers for the elevated subclinical and clinical CVD risk in individuals on LLT with LDL levels <130 mg/dL (group 3) are unfavourable levels for other risk factors, including systolic blood pressure, body mass index, diabetes and HDL in participants in

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Table 3 Impact of subclinical atherosclerosis and of BNP and CRP levels on the association of LDL group with incident CVD

	LDL group	LDL treatment	HR (95% CI)*	HR (95% CI)†	Change in HR‡	HR (95% CI)§	Change in HR¶
1	<100 mg/dL	No	1.00 (referent)	1.00 (referent)		1.00 (referent)	
2	≥100 to <130 mg/dL	No	1.34 (0.99 to 1.81)	1.34 (0.99 to 1.81)	0.0% decrease	1.35 (1.00 to 1.83)	0.1% increase
3	<130 mg/dL	Yes	1.57 (1.07 to 2.31)	1.50 (1.02 to 2.20)	4.5% decrease	1.55 (1.05 to 2.28)	1.3% decrease
4	≥130 mg/dL	No	1.44 (1.08 to 1.93)	1.39 (1.04 to 1.86)	3.5% decrease	1.42 (1.06 to 1.91)	1.4% decrease
5	≥130 mg/dL	Yes	1.51 (0.98 to 2.32)	1.43 (0.93 to 2.20)	5.3% decrease	1.41 (0.92 to 2.17)	6.6% decrease

Analyses were conducted in a subsample with available information on carotid ultrasound abnormality, BNP and CRP (n=2811).
*HRs are adjusted for age, sex, systolic blood pressure, antihypertensive medication, smoking and diabetes mellitus. Note: HRs are slightly different from figure 3 due to differences in sample size.
†Multivariable model additionally adjusted for the presence versus absence of carotid ultrasound abnormality.
‡Change in the HR on additional adjustment for the presence of carotid ultrasound abnormality compared with the HR adjusted for only age, sex, systolic blood pressure, antihypertensive medication, smoking and diabetes mellitus.
§Multivariable model additionally adjusted for the presence versus absence of carotid ultrasound abnormality+adjustment for ln(BNP) and ln(CRP).
¶Change in the HR on additional adjustment for the presence of carotid ultrasound abnormality+ln(BNP)+ln(CRP) compared with the HR adjusted for only age, sex, systolic blood pressure, antihypertensive medication, smoking and diabetes mellitus.
BNP, B-type natriuretic peptide; CVD, cardiovascular disease; CRP, C reactive protein; LDL, low-density lipoprotein.

this group (table 1). The greater CVD risk at baseline in individuals on LLT (also reflected in online supplementary figure 1) may reflect indication bias (confounding by indication; higher risk individuals are more likely to be treated by their physicians), as expected in observational (non-randomised) studies assessing residual risk on treatments.[29] Indeed, one of the factors contributing to the decision whether LLT is initiated in a given patient is the overall CVD risk over a 10-year time horizon, which takes levels of several risk factors into account.[17] On the other hand, and as a clinical consequence, these observations underscore the importance of monitoring all traditional risk factors in individuals on LLT, even if people reach LDL levels below 130 mg/dL.

It also has to be kept in mind that LDL-cholesterol, despite being an established lipid measure and CVD risk factors, might mirror LDL-related residual CVD risk not ideally, particularly in subgroups of the population, where other lipid measures (such as apolipoprotein B (ApoB) concentrations) provide discordant information. In the Women's Health Study, approximately one in five women were discordant for LDL-C and ApoB; and CVD risk in these individuals was underestimated or overestimated, when estimations were solely based on LDL-C.[30] Also in the Framingham Offspring cohort, ApoB levels provided additional information about CVD risk, beyond established lipid measures, in individuals, where the observed ApoB levels were discordant to the expected ApoB concentrations, as predicted based on circulating LDL-C.[31] Furthermore, in Jupiter trial participants (individuals with LDL <130 mg/dL and high-sensitivity C-reactive protein (hsCRP) ≥2.0 mg/L), ApoB levels, but not LDL-C, were associated with incident CVD in the placebo arm.[32] In the on-statin arm, however, ApoB failed to reach significance (HR: 1.20, 95% CI 0.98 to 1.46 for incident CVD).[32]

Strengths and limitations

The strengths of our investigation include the large community-based sample, the careful and comprehensive characterisation of our study participants, including measures of subclinical atherosclerosis at the carotid arteries and of biomarkers of systemic inflammation and neurohormonal activation, and the prospective follow-up for incident CVD (median follow-up, 13.7 years). The following limitations merit consideration. We assigned participants to LDL groups based on a single LDL measurement and a single assessment of the intake of lipid-lowering medication. This might result in some misclassification, which is likely to be non-differential; this would bias us towards the null hypothesis of no difference between the groups in risk of outcome events. In our analyses we did not differentiate between different types of lipid-lowering medication, dosages and durations of treatment. Unfortunately, information on the generation of statin taken or on statin dose was not available in our sample. Furthermore, the prescription of LLT is determined by circulating lipid concentrations and by

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the overall burden of CVD risk factors. Thus, individuals on LLT in the community might a priori be at higher CVD risk as compared with individuals not receiving such medication (confounding by indication).[29] Finally, we have a relatively small number of individuals on statins. However, these individuals were well phenotyped and had a relatively long follow-up period well beyond that of typical clinical trials of LLT.

In conclusion, we observed a substantial residual cardiovascular risk in individuals on lipid-lowering medication in the community, partly explained by an adverse profile of other CVD risk factors and in part (by about 5%) by carotid subclinical atherosclerosis in these people. These observations emphasise the necessity to closely monitor all standard CVD risk factors in individuals on LLT, even when LDL levels are lowered effectively.

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Data sharing statement The data from the Framingham Heart Study used for this project are available in the NHLBI data repository BioLINCC ().

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