Focus on Lipids: High-Density Lipoprotein Cholesterol and Its Associated Lipoproteins in Cardiac and Renal Disease

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**Introduction**

Elevated high-density lipoprotein cholesterol (HDL-C) has been suggested to be strongly related to decreased atherosclerotic events in large population-based studies \cite{1}. Each 1 mg/dl increase in HDL-C was associated with a decrease of 2–3\% in the risk of coronary heart disease events \cite{1}. In observational studies and early randomized trials, largely in statin-naive patients, with nicotinic acid, fibrates, and lifestyle changes, an increase in HDL-C was associated with a modest reduction in cardiovascular events (fig. 1). Recently completed studies have shed light on the relationships between the plasma pool HDL-C concentration and cardiovascular events as well as the unique effects of the apoproteins carried by the HDL particle on end-organ disease.

**Trials on Raising HDL-C and Clinical Outcomes**

There have been many clinical trials testing the effect of raising HDL-C. We summarize here the results of modern cholesteryl ester transfer protein (CETP) inhibitors.

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**Key Words**

Lipids · High-density lipoprotein cholesterol · Lipoproteins · ApoL1

**Abstract**

High-density lipoprotein cholesterol (HDL-C) contains dozens of apoproteins that participate in normal cholesterol metabolism with a reliance on renal catabolism for clearance from the body. The plasma pool of HDL-C has been an excellent inverse predictor of cardiovascular events. However, when HDL-C concentrations have been manipulated with the use of niacin, fibric acid derivatives, and cholesteryl ester transferase protein inhibitors, there has been no improvement in outcomes in patients where the low-density lipoprotein cholesterol has been well treated with statins. Apolipoprotein L1 (APOL1) is one of the minor apoproteins of HDL-C, newly discovered in 1997. Circulating APOL1 is a 43-kDa protein mainly found in the HDL3 subfraction. In patients with chronic kidney disease (CKD), mutant forms of APOL1 have been associated with rapidly progressive CKD and end-stage renal disease (ESRD). Because mutant forms of APOL1 are more prevalent in African Americans compared to Caucasians, it may explain some of the racial disparities seen in the pool of patients with ESRD in the United States. Thus, HDL-C is an important lipoprotein carrying apoproteins that play roles in vascular and kidney disease.
tor trials and niacin trials which have been largely carried out under a more modern standard of care where patients have control of low-density lipoprotein cholesterol (LDL-C) levels through an improved diet, statins, ezetimibe, bile acid sequestrants, and other agents. These trials give a more accurate picture of the impact of raising HDL-C as an intervention on top of good clinical practice in primary or specialty care.

**CETP Inhibitor Trials**

CETP works to shuttle cholesterol and triglycerides between HDL and very low-density lipoprotein (VLDL) and other apolipoprotein B100-containing particles (fig. 2). Early studies demonstrated that inhibition of this protein markedly raised the HDL-C concentration; however, it was unclear whether this improved or potentially worsened the reverse cholesterol transport. The Investigation of Lipid Level Management to Understand Its Impact in Atherosclerotic Events (ILLUMINATE) trial is a phase 3 randomized double-blind clinical trial that investigated the role of the first CETP inhibitor, torcetrapib, on cardiovascular events and death in 15,067 patients at high cardiovascular risk. The patients in the torcetrapib group showed an increase of 72% in HDL cholesterol after 12 months compared to baseline [2]. However, the total mortality (HR 1.58; 95% CI 1.14–2.19) and cardiovascular events (HR 1.25; 95% CI 1.09–1.44) also increased significantly [2]. The off-target effect of torcetrapib of increasing the systolic blood pressure, sodium, and aldosterone was blamed as a possible mechanism that resulted in increased mortality rates and cardiovascular events [2, 3]. In this study, despite the overall findings, those who achieved the highest HDL-C concentrations had the lowest overall event rates.

The dal-OPTIMISE trial is a phase 3 randomized double-blind clinical trial that investigated the role of the second CETP inhibitor, dalcetrapib, on the composite primary endpoint of death from coronary heart disease, cardiac arrest with resuscitation, nonfatal myocardial infarction, unstable angina, or ischemic stroke in 15,871 patients with recent acute coronary syndrome [3]. The patients in the dalcetrapib group showed an increase of 31–40% in HDL cholesterol compared to baseline [3]. Dalcetrapib did not lower the risk of the primary endpoint (HR 1.04; 95% CI 0.93–1.56) or total mortality [3].

Anacetrapib is the third CETP inhibitor currently under evaluation in a phase 3 clinical trial (Randomized Evaluation of the Effects of Anacetrapib through Lipid Modification; REVEAL). The trial has an enrollment of 30,000 patients with atherosclerotic disease on statin treatment (http://clinicaltrials.gov/ct2/show/NCT01252953?term%BCreveal&rank%BC1). The results of the trial on the effects of anacetrapib on the risk of myocardial infarction, coronary death, and coronary revascularization should be available in January 2017 [4].

Evacetrapib is the fourth CETP inhibitor that is now also being evaluated in a phase 3 clinical trial (Assessment of Clinical Effects of Cholesteryl Ester Transfer Protein Inhibition with Evacetrapib in Patients at a High-Risk for Vascular Outcomes; ACCELERATE) enrolling 12,000 patients with high-risk vascular disease on statin treatment (http://clinicaltrials.gov/ct2/show/record/
NCT01687998). The results of this trial on the effects of evacetrapib on the composite endpoint of risk of myocardial infarction, cardiovascular death, stroke, hospitalization for unstable angina, and coronary revascularization will be available in January 2016 [4]. Figure 3 displays the CTEP inhibitors and summarizes their effects on HDL-C and LDL-C.

**Niacin Trials**

The Atherothrombosis Intervention in Metabolic Syndrome with Low HDL/High Triglycerides: Impact on Global Health Outcomes (AIM-HIGH) trial is a randomized double-blind clinical trial that investigated the effect of the addition of extended-release niacin to intensive statin therapy on the composite primary endpoint of death from coronary heart disease, nonfatal myocardial infarction, ischemic stroke, hospitalization for an acute coronary syndrome, or symptom-driven coronary or cerebral revascularization in 3,414 patients with cardiovascular disease [5]. Extended-release niacin increased the HDL cholesterol from 35 to 42 mg/dl after 2 years compared to baseline [5]. Extended-release niacin did not change the risk of the primary endpoint (HR 1.02; 95% CI 0.87–1.12), and the trial was stopped prematurely due to a lack of benefit and because there was a higher rate of ischemic stroke in patients in the niacin group (fig. 4) [5].

The Heart Protection Study 2-Treatment of HDL to Reduce the Incidence of Vascular Event (HPS-2-THRIVE) trial is a randomized placebo-controlled trial that investigated the effect of the addition of extended-release niacin plus laropiprant (ERN/LRPT) to intensive statin therapy on the composite primary endpoint of cardiovascular death, myocardial infarction, stroke, or revascularization in 25,673 patients with occlusive arterial disease [6]. ERN/LRPT increased the HDL-C by 16.9%
compared to baseline [6]. ERN/LRPT did not change the risk of the primary endpoint (14.5% of the ERN/LRPT group vs. 15% of the placebo group, p = 0.29) after a mean of 3.9 years of follow-up [5]. Twenty-five percent of patients in the ERN/LRPT group compared to 17% of the placebo group stopped taking the study medications because of diverse side effects including skin problems and gastrointestinal, musculoskeletal, and diabetes-related side effects and myopathy [7].

There are several possible explanations for the failure of pharmacologic elevation of HDL-C to improve the clinical outcomes in modern trials. First, it may be difficult to show an incremental benefit of raising HDL-C for the prevention of cardiovascular events when patients are currently receiving high-quality medical treatment, including intensive statin, antiplatelet, β-blocker, and angiotensin-converting enzyme inhibitor drugs [3, 5]. Secondly, increased HDL-C may not induce the desired reverse cholesterol transport or that mechanism may be counterbalanced by another proatherosclerotic force. Only 40% of the total variance of the cholesterol efflux capacity from macrophages (a metric of HDL reflecting a reverse cholesterol transport) was explained by the HDL-C level, so quantitative levels of HDL-C may not reflect the desired reverse cholesterol transport function adequately [8]. Also, under oxidative stress, HDL-C can

**Fig. 3.** Chemical structures of CETP inhibitors and their approximate effects on HDL-C and LDL-C. Adapted from von Eckardstein. http://www.medscape.org/noscan/slideshow/760297#25.

**Fig. 4.** Main results of the AIM-HIGH trial. Extended-release niacin did not change the risk of the primary endpoint compared to the placebo group. Adapted from Boden et al. [7]. MI = Myocardial infarction. With permission from the New England Journal of Medicine.
function as a proatherogenic, proinflammatory force rather than as a protective molecule, inhibiting or preventing atherogenesis [9].

**Primer on Apolipoprotein L1**

Apolipoprotein L1 (APOL1) is one of the minor apo-proteins of HDL-C, newly discovered in 1997 [10]. Circulating APOL1 is a 43-kD protein mainly found in the HDL3 subfraction and it was assumed that APOL1 might play a role in cholesterol metabolism, with no convincing evidence [10, 11]. APOL1 is synthesized in a number of tissues, including those of the pancreas and kidney [11]. In a normal kidney, APOL1 is localized in podocytes, proximal tubules, medium-sized arteries, and arteriolar endothelial cells, but it is not certain whether localization of APOL1 in the kidney reflects de novo synthesis or uptake from circulation [11].

Circulating APOL1 functions as a trypanosome lytic factor, creating pores in *Trypanosoma brucei* that confer immunity against this parasite [12]. *T. b. rhodesiense*, which causes African sleeping sickness, can create a resistance factor called the SRA protein that can bind to the C-terminal portion of APOL1 and inhibits APOL1-mediated lysis (fig. 5) [12]. APOL1 renal risk variants such as G1 and G2 variants affect a joint portion between APOL1 and the SRA protein, prohibiting SRA binding of APOL1 and conferring resistance to infection with *T. b. rhodesiense*. This selective pressure was hypothesized to increase G1 and G2 variants among people of African descent and could explain the disproportionally high prevalence of nondiabetic chronic kidney disease (CKD) in people of African descent [12].

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**Fig. 5.** Trypanolytic properties of APOL1. a Circulating APOL1 functions as a trypanosome lytic factor, forming pores in *T. brucei* that confer immunity against this parasite. b *T. b. rhodesiense*, which causes African sleeping sickness, can create a resistance factor called the SRA protein that can bind to the C-terminal portion of APOL1 and inhibits APOL1-mediated lysis. c APOL1 renal risk variants such as G1 and G2 variants affect a joint portion between APOL1 and the SRA protein, prohibiting SRA binding of APOL1 and conferring resistance to infection with *T. b. rhodesiense*. Adapted from Friedman and Pollak [12] and Pays et al. [18] with permission from *Journal of Clinical Investigation* and *Current Opinion in Immunology*. WT = Wild type; Hb-Hpr = hemoglobin and haptoglobin related protein.
Apoprotein L1 and the Progression of CKD

APOL1 renal risk variants may explain the racial differences noted extensively in the literature between African-Americans and Caucasians with respect to the progression of CKD [12, 13]. APOL1 renal risk variants confer 10- to 17-fold higher odds of development of focal segmental glomerulosclerosis (FSGS) [14], 29-fold higher odds of human immunodeficiency virus-associated nephropathy (HIVAN) [15], and 7.3-fold higher odds of end-stage renal disease attributed to hypertension [14].

Whether APOL1 renal risk variants affect kidney function in people with diabetes has been the subject of debate [13]. However, in the Chronic Renal Insufficiency Cohort (CRIC) study, the APOL1 high-risk variant was associated with a rapid decline in the glomerular filtration rate in black patients, regardless of the presence of diabetes mellitus [13]. In summary, analogous to how hemoglobin S in sickle cell disease provides protection from malaria yet causes hematologic disease over the course of a lifetime, APOL1 mutants protect against African sleeping sickness due to trypanosomiasis yet they appear to provide susceptibility to rapidly progressive CKD in those with hypertensive kidney disease, diabetes, and HIVAN and may be directly pathogenic in FSGS (fig. 6). Future research on the renal tubular catabolism of HDL and its apoproteins including APOL1 is likely to lead to new diagnostic and therapeutic targets for patients with rapidly progressive CKD.

Conclusions

The HDL-C concentration is epidemiologically associated with a reduced risk of atherosclerotic events. However, pharmacological increases in this concentration have not resulted in a change in the risk, implying that the reverse cholesterol transport has either not been altered or has been counterbalanced by some other proatherosclerotic force. APOL1 is a minor apolipoprotein which, when in mutant form, protects against trypanosome infections but which has been consistently associated with the progression of CKD in a variety of diseases. APOL1 polymorphisms may have specific relevance for explaining the racial differences noted extensively in the literature between African-Americans and Caucasians with respect to the progression of CKD and the development of end-stage renal disease.
References


