INVITED REVIEW

The Trials and Tribulations of CETP Inhibitors

Alan R Tall1, Daniel J Rader2

1Division of Molecular Medicine, Department of Medicine, New York, NY 10024; 2Departments of Genetics and Medicine, Perelman School of Medicine, University of Pennsylvania, Philadelphia, PA 19104

Running title: CETP Inhibition and Coronary Disease

Subject Terms:
Atherosclerosis
Coronary Artery Disease

Address correspondence to:
Dr. Alan Tall
Columbia University
Medicine
630 W 168th St
P&S 8-401
New York, New York 10032
UNITED STATES
Tel: 212-305-4899
Fax: 212-305-5052
art1@columbia.edu

DOI: 10.1161/CIRCRESAHA.117.311978
ABSTRACT

The development of CETP inhibitors has had a long and difficult course with three compounds failing in phase III clinical trials. Finally, the REVEAL trial has shown that the CETP inhibitor anacetrapib decreased coronary heart disease when added to statin therapy. While the result is different to earlier studies, this is likely related to the size and duration of the trial. The benefit of anacetrapib appears to be largely explained by lowering of non-HDL cholesterol, rather than increases in HDL cholesterol. Although the magnitude of benefit for CHD appeared to be moderate, in part this may have reflected aspects of the trial design. Anacetrapib treatment was associated with a small increase in BP, but was devoid of major side effects and was also associated with a small reduction in diabetes. Treatment with CETP inhibitors, either alone or in combination with statins, could provide another option for patients with coronary disease who require further reduction in LDL and/or non-HDL cholesterol.

Keywords:
Cholesteryl ester, transfer protein, atherosclerosis, coronary heart disease, LDL, HDL.

Nonstandard Abbreviations and Acronyms:

CETP cholesteryl ester transfer protein
REVEAL Randomized Evaluation of the Effects of Anacetrapib through Lipid modification
FOURIER Further Cardiovascular Outcomes Research with PCSK9 Inhibition in Subjects with Elevated Risk
ACCELERATE Assessment of Clinical Effects of Cholesteryl Ester Transfer Protein Inhibition with Evacetrapib in Patients with a High Risk for Vascular Outcomes
IMPROVE-IT The Improved Reduction of Outcomes: Vytorin Efficacy International Trial
DEFINE Determining the Efficacy and tolerability of CETP inhibition with anacetrapib

INTRODUCTION

The development of CETP inhibitors was motivated by the discovery that humans with genetic CETP deficiency have markedly elevated levels of HDL cholesterol (HDL-C), as well as reduced levels of LDL cholesterol (LDL-C), a profile that is typically associated with reduced atherosclerosis. CETP inhibitors were subsequently shown to raise HDL-C levels, in some cases quite impressively; in addition the more potent CETP inhibitors lowered LDL-C levels. Based on epidemiological observations, it was expected that this marked increase in HDL would deliver a powerful anti-atherogenic effect. This promise has not been realized in cardiovascular clinical outcome trials of CETP inhibitors. In fact, in the first large trial the CETP inhibitor torcetrapib caused an excess of deaths and cardiovascular disease (Table), leading many to conclude that the elevated HDL itself was harmful. The identification of off-target toxic side-effects of torcetrapib led to sufficient clinical equipoise to allow further evaluation of this class of drugs. Subsequent trials with the relatively ineffective CETP inhibitor dalcetrapib and with the potent inhibitor evacetrapib were stopped early for futility (lack of efficacy in reducing CV events). Now results from the largest and longest running trial of a CETP inhibitor, in this case the potent inhibitor anacetrapib, have been published, showing that this drug significantly reduced major coronary events. Although the magnitude of risk reduction was moderate, anacetrapib could find a place in the armamentarium of approved non-statin lipid-targeted agents. However, this result leaves many questions unanswered, a few of which include: 1) Why did this trial show benefit when other trials with CETP inhibitors did not? 2) Given the reductions in LDL and non-HDL cholesterol seen with anacetrapib, did the increase in HDL cholesterol contribute to the...
benefit? This review will attempt to address these questions, while providing a background on the role of CETP in lipoprotein metabolism, emphasizing genetic and human metabolic studies. The reader is referred to earlier reviews for additional background on CETP.6-9

<table>
<thead>
<tr>
<th>TRIAL (drug)</th>
<th>Patients</th>
<th>Lipoprotein Changes</th>
<th>Duration</th>
<th>Outcome</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>ILLUMINATE (Torcetrapib)</td>
<td>15,067 Hi CV Risk</td>
<td>HDL-C↑72% LDL-C ↓*</td>
<td>1-2 years</td>
<td>↑CV Events ↑Death ↑SBP (5mm)</td>
<td>Electrolyte disturbances, hyeraldosteronism identified as off-target effects</td>
</tr>
<tr>
<td>dal-OUTCOMES (Dalcetrapib)</td>
<td>15,871 Post ACS</td>
<td>HDL-C↑~30% LDL-C→</td>
<td>31 months</td>
<td>CV Events→SBP(0.6mm)</td>
<td>Trial stopped early for futility. Possible benefit in a genetic subgroup.</td>
</tr>
<tr>
<td>ACCELERATE (Evacetrapib)</td>
<td>12,092 Hi risk vascular disease</td>
<td>HDL-C↑133% LDL-C↓*</td>
<td>26 months</td>
<td>CV Events→SBP(1.2mm)</td>
<td>Trial stopped early for futility. Possible benefit in a genetic subgroup.</td>
</tr>
<tr>
<td>REVEAL (Anacetrapib)</td>
<td>30,449 Hi risk vascular disease</td>
<td>HDL-C↑104% LDL-C↓17%</td>
<td>4.1 years</td>
<td>↓Coronary Events SBP (0.7mm)</td>
<td>Trial went to planned completion ↓ new onset diabetes</td>
</tr>
</tbody>
</table>

A reduction in coronary heart disease with CETP inhibition is revealed.

The REVEAL study involved 30,449 patients with atherosclerotic cardiovascular disease who were randomized to receive anacetrapib 100 mg daily or placebo on top of effective statin therapy and followed for a median of 4.1 years. After the failure of CETP inhibitors in three successive clinical trials, expectations were low that anacetrapib, a CETP inhibitor developed by Merck, would meet with success. However, REVEAL demonstrated a highly significant reduction (rate ratio = 0.91, p<.004) in the composite primary endpoint of coronary death, myocardial infarction (MI) or coronary revascularization.5 The individual components of the primary endpoint showed similar rate ratios but were only significant for MI and revascularization. The incidence of the pre-specified outcome of coronary death or MI was significantly lower in the anacetrapib group (rate ratio 0.89, p=.008). The secondary endpoint of major coronary event (MI, coronary death or ischemic stroke) just missed significance (rate ratio = 0.93, p=.052). There was also a significant difference for the secondary outcome of major vascular events favoring anacetrapib. At the trial midpoint, anacetrapib raised HDL-C from 42 to 86 mg/dl (104%) and apoA-I by 36%; in addition, it lowered LDL-C from 63 mg/dl to 53 mg/dl (-17%, as determined by ultracentrifugation), non-HDL-C from 96 to 79 mg/dl (-18%) and Lp(a) by -25%. While the outcome of REVEAL appears inconsistent with the previous negative trials, it is important to assess the differences in CETP inhibitors and trial design (Table). Dalcetrapib is a much less potent CETP inhibitor that raised HDL cholesterol less and did not reduce LDL cholesterol; furthermore, its trial dal-OUTCOMES was stopped early.3 The ACCELERATE trial was performed with the potent CETP inhibitor evacetrapib which results in lipid changes similar to anacetrapib. However, ACCELERATE involved less than half the number of patients and CVD events, and was stopped early after about 2.2 years for futility.4 In contrast, REVEAL was continued through its planned duration with a median follow-up of 4.1 years. The longer trial duration was likely of key importance because, similar to trials of other lipid lowering drugs, the beneficial effects on
CVD were greater after the first year of treatment with anacetrapib. Moreover, an exploratory analysis suggested that the benefit of anacetrapib increased with time as the trial proceeded. Thus, it is highly plausible that anacetrapib succeeded where other CETP inhibitors failed because of relative safety (compared with torcetrapib), potency (compared with dalcetrapib) and a study design that included adequate statistical power and sufficiently long duration to uncover the benefit (compared with evacetrapib).

The demonstration of an overall anti-atherogenic effect of CETP inhibition is buttressed by the majority of animal studies which have demonstrated a pro-atherogenic effect of CETP expression. While studies in CETP transgenic mice have produced mixed results in atherosclerosis experiments, inhibition of CETP in rabbits, a species that naturally expresses CETP, have consistently shown reduced atherosclerosis, including in a CETP knockout rabbit which showed reduced aortic and coronary atherosclerosis when fed a high cholesterol diet. Moreover, multiple large human genetic studies have shown that SNPs in the CETP gene that are associated with increased HDL and reduced LDL cholesterol are associated with reduced CHD. This includes SNPs that likely reduce the function of the promoter region upstream of the CETP gene, and most importantly CETP protein truncating mutations that abrogate the function of CETP.

How does CETP inhibition affect plasma lipoprotein levels?

Before considering whether the benefit of anacetrapib was related to changes in HDL, LDL or both, it is worth reviewing the mechanisms underling the effects of CETP inhibition on plasma lipoprotein metabolism. The primary effect of CETP inhibition is a reduced rate of transfer of CE from HDL into triglyceride rich lipoproteins (TRL). This leads to an increased content of CE in HDL and the formation of larger HDL particles that are more slowly catabolized than normal. On the other side of the coin, CE is depleted in the TRL including in VLDL, chylomicrons and their remnants. There is also a depletion of CE in LDL likely reflecting both diminished direct transfer from HDL, reduced amounts of CE in VLDL being converted into LDL CE and increased LDL particle catabolism (Fig 1). Thus, the major impact of CETP inhibition is an increase in HDL cholesterol and a decrease in non-HDL cholesterol (encompassing both cholesterol in TRL and LDL). Less obviously, the reduced transfer of CE from HDL to TRL also leads to a decrease in VLDL and LDL apoB levels. Careful metabolic studies in mildly hypercholesterolemic subjects treated with anacetrapib have shown that the reductions in VLDL and LDL apoB, as in genetic CETP deficiency, result from an increase in catabolism. In contrast, lowering of Lp(a) reflects a decrease in the production rate of apo(a). On a background of statin therapy, anacetrapib modestly lowered plasma triglyceride levels, reflecting an increase in VLDL TG catabolism.

Since LDL catabolism is primarily mediated by LDL receptor mediated clearance in the liver, increased LDL catabolism in anacetrapib treated subjects likely reflects an increase in the clearance of LDL particles via the LDLR. This could result from a relative depletion of the regulatory cholesterol pool in the liver resulting from CETP inhibition and leading to an increase in LDLR mRNA (Fig 1B). However, there was no increase in plasma lathosterol, which likely would have been increased as a result of increased sterol biosynthesis, if regulatory cholesterol pools in the liver were depleted. Thus, the authors speculated that the increased clearance of apoB could be caused by an increase in the affinity of LDL for its receptor, caused by changes in the properties of LDL particles, such as an increase in the LDL TG/CE ratio or an increase in the size or polydispersity of LDL particles (Fig 1B). However, an increase in the numbers of hepatic LDL receptors cannot be completely excluded since they were not directly measured. Studies in CETP transgenic mice have shown that CETP activity increases the cholesterol content in the liver and lowers the levels of the LDL receptor mRNA and protein, as well as HMGCoA reductase mRNA, consistent with an increased content of the regulatory pool of cholesterol in the liver. This may reflect an increased

DOI: 10.1161/CIRCRESAHA.117.311978
efficiency of cholesterol transfer into hepatocytes resulting from transfer of CE from slowly turning over HDL particles into more rapidly cleared TRL remnants, or decreased channeling of HDL cholesterol into biliary cholesterol bypassing the regulatory cholesterol pool. Whatever the precise mechanism, these studies strongly suggest that increased clearance of apoB lipoproteins via the LDL receptor pathway is occurring in subjects treated with potent CETP inhibitors, an effect that ultimately reflects the reduced rate of transfer of CE from HDL to TRL.

**Is the CHD benefit related to increased HDL or decreased LDL/non-HDL cholesterol?**

Formally, the only firm conclusion from REVEAL is that CETP inhibition resulted in a reduction in CV events as captured by the primary endpoint,\(^5\) and the mechanism of benefit is the subject of speculation. However, the mean reduction of non-HDL cholesterol of 18% in REVEAL fell only slightly above the regression line relating the % reduction in CVD events to the absolute decrease in non-HDL-C, based on earlier studies with LDL lowering drugs, suggesting that the benefit may be largely explained by the reduction in non-HDL cholesterol. However, these point estimates have substantial confidence intervals and also are based on an extrapolation of data that were mostly obtained at much higher levels of non-HDL cholesterol. Moreover, there may have been a small adverse effect related to BP elevation in REVEAL, perhaps offsetting a beneficial effect of HDL changes. Overall, the conservative interpretation is that the benefit of anacetrapib was mostly or solely due to reduction in atherogenic lipoproteins; however, a contribution of HDL raising cannot be firmly excluded.

After the failure of torcetrapib, studies of HDL functionality were undertaken to exclude an adverse effect of CETP inhibition. These studies showed that in fact HDL from subjects treated with anacetrapib had enhanced ability to promote cholesterol efflux from cholesterol loaded macrophages and preserved anti-inflammatory effects.\(^22\) Studies using an assay of HDL cholesterol efflux capacity that had been validated as predicting coronary atherosclerosis and CHD in sizable population studies\(^23,\)\(^24\) also showed that HDL from subjects treated with evacetrapib had increased cholesterol efflux capacity.\(^25\) However, for unknown reasons the magnitude of this increase in subjects concomitantly treated with statins was surprisingly small (total cholesterol efflux capacity was increased by 21%) versus a larger increase in those receiving evacetrapib monotherapy (34%). Another possibility is that statin use reduces the expression of the ATP binding cassette transporters (ABCA1 and ABCG1) in macrophages, as result of decreased activity of the sterol-activated transcription factor LXR, thereby offsetting the effects of increased HDL levels on macrophage cholesterol efflux. Further studies of HDL functionality in the context of CETP inhibition are warranted.

**Is the outcome of REVEAL consistent with insights from human genetics studies?**

Mendelian randomization studies of SNPs in multiple genes that are associated with changes in only HDL-C (but not triglycerides or LDL-C), for example in endothelial lipase (LIPG), have shown no association with CHD.\(^26\) This has been interpreted as indicating that HDL-C is not in the causal pathway of atherosclerosis,\(^27\) and suggests that decreased CHD associated with SNPs that reduce CETP expression or function are likely acting through changes in LDL-C (or non-HDL-C). Consistent with this interpretation, in the study of CETP truncating mutations, the magnitude of the benefit on CHD correlated well with the degree of LDL cholesterol lowering.\(^12\) However, variants in HDL-associated genes jointly account for very little of the variance in HDL-C levels and could have pleiotropic effects, weakening the general conclusion that HDL is not in the causal pathway of atherosclerosis. Adding to this complexity, there could be epigenetic effects masking a possible benefit of HDL raising genes. In this regard, a post-hoc analysis of the dal-OUTCOMES trial has identified SNPs in the gene encoding Adenylate Cyclase 9 (ADCY9) as being associated with cardiovascular benefit in a subgroup of patients; a clinical trial (dal-
GENE) in which dalcetrapib is being administered to high CV risk patients carrying the putative protective SNPs is ongoing. It will be of substantial interest to determine whether carriers of these variants had even better outcomes with anacetrapib in the REVEAL trial. Further analysis of CETP inhibitor clinical trial data, new assays of HDL functionality, as well as clinical outcomes studies based on infusion of reconstituted HDL particles that are highly active in promoting cholesterol efflux may provide additional insights into the complex relationship of HDL to atherosclerosis.

Is the glass half empty or half full?

Despite achieving its primary endpoint, there may be concern that the benefit demonstrated by REVEAL was moderate and that the reduction in cardiovascular death was not significant. It is worth considering whether CETP inhibition is a mechanism with intrinsically limited benefit, or whether the result may have reflected aspects of the trial design. In general, the relationship between LDL lowering and its impact on CHD has shown that for every 40 mg/dl decrease in LDL cholesterol, there is about a 25% reduction in CHD risk. This implies that in a trial of LDL lowering, if patients in both placebo and active treatment groups are treated to very low LDL levels prior to randomization, the % reduction in CHD and the absolute benefit in the active treatment group will be less than they would have been if LDL levels were higher at randomization. This is because the % reduction in LDL-C (or non-HDL-C) will likely be similar at lower or higher starting LDL-C, so that the absolute reduction in LDL-C is greater when the starting LDL-C is higher and the risk reduction is accordingly larger. This supposition is supported by studies performed in a subset of patients in the DEFINE trial which showed a larger absolute reduction in LDL-C by anacetrapib in patients with higher LDL-C at baseline compared to those with lower levels of LDL-C at baseline. In REVEAL the mean LDL cholesterol at randomization was very low – 61 mg/dl. In part this reflected the trial goal of achieving a pre-randomization LDL cholesterol < 77 mg/dl by effective use of statins, but also the fact that people with total cholesterol l>155 mg/dl after the statin run-in were intentionally excluded from the study. Notably, patients with non-HDL cholesterol > 101 mg/dl at randomization appeared to benefit more from anacetrapib than patients with non-HDL cholesterol < 85 mg/dl, with reductions in the rate ratio of MI + coronary death of 0.83 versus 0.96. In the clinical setting, patients with persistently elevated LDL or non-HDL cholesterol on maximally tolerated statin therapy would be precisely those most likely to be treated with additional lipid-lowering agents. The implication could be that for people who would likely derive the greatest benefit, i.e. with non-HDL cholesterol>100 mg/dl after statin treatment, the clinical impact of anacetrapib could be considerably greater.

It is of interest to consider the results of REVEAL in the light of other recent trials of LDL lowering therapies that met their primary endpoints. There was a 6% reduction (p<.02) in the primary cardiovascular endpoint in IMPROVE-IT after 6 years of treatment with ezetimibe on top of statins. In this trial the mean LDL cholesterol level in the statin only arm was slightly higher than in REVEAL (69.5 mg/dl) and reductions in LDL cholesterol, non-HDL cholesterol and apoB were comparable to those obtained with atorvastatin. In contrast to evacetrapib, ezetimibe did not substantially increase HDL cholesterol or lower Lp(a) levels. In the FOURIER study in which evolocumab (a PCSK9 mAb) was added to LDL lowering therapy with statins, the reduction in the primary endpoint was about 15%. Compared to anacetrapib, there was a much more dramatic incremental LDL cholesterol lowering of 61%. However, it also should be noted that in FOURIER the LDL cholesterol was 92 mg/dl at entry reflecting the fact that the trial protocol stipulated that patients with already low LDL cholesterol on statins (< 70 mg/dl) were excluded from the study. Thus, the trial was more specifically designed to show a benefit of LDL lowering on CHD. Evolocumab lowered Lp(a) by about 27% and raised HDL cholesterol by 8%. FOURIER was terminated after a mean of only 2.5 years and very likely the beneficial impact on CV events would have been larger had the study been continued for a longer time.
The devil is in the details: Safety and side-effects.

In REVEAL, anacetrapib treatment was devoid of major side effects i.e. involving cancer, infectious diseases, cognitive changes, depression etc. Although not observed in a smaller preliminary safety trial, there was a 0.7 mm increase in SBP and 0.3 mm in DBP in anacetrapib treated subjects in REVEAL, similar in magnitude to what was observed with evacetrapib and dalcetrapib, but much less than for torcetrapib. Increased BP has not been reported in genetic CETP deficiency. However, this effect has been seen in multiple trials with different structural classes of CETP inhibitors strongly suggesting that it is mechanism-based. The nature of this mechanism is unknown. For torcetrapib, hyperaldosteronism and increased responses to endothelin in the vasculature were shown, however this occurred even in species that lack a CETP gene. Thus, relevance to subsequent CETP inhibitors is unlikely. Anacetrapib use was associated with a small increase in the proportion of patients with eGFR<60ml/min at the end of the study, but there was no increase in albuminuria or serious adverse events attributed to renal failure. An increase in hemorrhagic stroke, as might be expected from increased BP and also as seen in a recent genetic study of CETP polymorphisms, was not observed. The beneficial effect of CETP inhibition was observed across multiple pre-specified subgroups. However, patients taking ACE inhibitors or ARBs appeared to benefit less (p<.01, unadjusted for multiple comparisons). While this could represent a chance finding, in a small study an adverse impact of genetic CETP deficiency on CHD was reduced after adjustment for treatment with non-diuretic anti-hypertensive drugs. It is possible that treatment with ACE inhibitors marks a subgroup that had a more marked hypertensive response to CETP inhibition, or that inhibition of the renin-angiotensin system uncovers an adverse effect of CETP inhibition. Further investigation of a possible interaction between the use anti-hypertensive drugs and CETP inhibitors is warranted.

Although several genetic studies have shown an association between CETP polymorphisms and age-related macular degeneration, there was no evidence for an increase in the onset or progression of retinal disease in REVEAL. This might reflect the differences between several years of pharmacologic CETP inhibition vs lifelong genetic reduction in CETP deficiency; longer term monitoring will be needed to exclude macular degeneration as a possible adverse effect related to CETP inhibition. Anacetrapib has a very prolonged half-life due to accumulation in adipose tissue, a property that appears to be specific to anacetrapib and has not been seen with some other potent CETP inhibitors. Plasma levels of anacetrapib fall substantially after cessation of anacetrapib treatment, but the drug is persistent in adipose tissue for at least several years. While no adverse effect has so far been linked to this property, it will be important to continue to monitor patients in REVEAL for potential consequences.

In REVEAL the incidence of new onset diabetes was reduced by about 10%, and there was a small reduction in HbA1c levels amongst non-diabetics. These beneficial effects on diabetes are consistent with previous reports of improvements in glucose/insulin ratios (HOMA-IR) and in HbA1c reported with torcetrapib or evacetrapib (but not dalcetrapib), as well as the reductions in glucose levels seen in a study of subjects with CETP deficiency. The mechanisms underlying the beneficial effects of CETP inhibitors on diabetes are poorly understood, but they contrast with the slight increase in diabetes and HbA1c associated with other mechanisms that lower LDL cholesterol by up-regulating the hepatic LDL receptor pathway, such as statins or genetic factors that reduce PCSK9. Since these effects on diabetes are the opposite to those observed with other drugs increasing LDL clearance by the LDLR pathway, it is tempting to speculate that they may be related to the distinctively increased HDL levels resulting from CETP inhibition. Cholesterol accumulation in islet beta-cells is associated with reduced insulin secretion in mice with knockouts of ABCA1/G1 in pancreatic beta-cells likely reflecting decreased HDL-mediated cholesterol efflux. However, statins would likely reduce beta cell cholesterol accumulation, so this is not an adequate explanation. There is high expression of CETP in insulin target tissues such as adipose and
muscle, raising the possibility of a local effect related to insulin sensitization. Finally, anacetrapib treatment on a background of statin therapy causes an increase in TG/apoB ratio of newly secreted TRL, reflecting decreased TG-CE interchange between TRL and HDL, and this TG enrichment of large TRL may increase susceptibility to lipoprotein lipase-mediated lipolysis. Enhanced activity of lipoprotein lipase through various mechanisms is associated with decreased risk of both CHD and diabetes and this could contribute to the decrease in diabetes associated with use of anacetrapib.

Is there a light at the end of the tunnel?

The results of the REVEAL trial suggest that CETP inhibitors could represent a useful addition to the armamentarium of drugs currently being used to treat high risk subjects intolerant to or not adequately treated with statins. The clinical use of evolocumab in patients who might benefit from this treatment has so far proven to be less than anticipated, in large part reflecting resistance from third party payers, and possibly from patients due to the need for subcutaneous injections. Thus, cost and convenience will likely be a major factor in the uptake of lipid lowering therapies with incremental benefits over generic statins. As noted above, the side effect profile for anacetrapib is distinct from statins and PCSK9 deficiency, in particular the decrease in diabetes risk (although small) may be viewed beneficially by patients and physicians. Although the increase in mean BP is small, in individual patients it may be larger and BP would need to be closely monitored. Interestingly, CETP inhibitors appear to be more effective as monotherapy than when used with statins. The impact on both reducing apoB levels and on increasing cholesterol efflux capacity is substantially more (about 1.5-2.0 fold) when potent CETP inhibitors are used alone versus when they are added to statins. In earlier trials (Table) the degree of LDL-C lowering by CETP inhibitors was overestimated likely reflecting changes in apoB particle composition and as a practical matter non-HDL-C should be used in the future to assess the effects of CETP inhibition. As monotherapy, the decreases in LDL-C and apoB are similar to the large effects on centrifugally separated LDL cholesterol (-40%) and apoB (-35%) that were observed in complete genetic CETP deficiency. This likely reflects the fact that CETP inhibition and statins both act in the same pathway to lower LDL apoB, i.e. by increasing activity of the LDLR, limiting the incremental benefit of the CETP inhibitor. Thus, CETP inhibitors should be further evaluated for use as monotherapy, for example in statin intolerant individuals. In summary, in a trial much larger and more than twice as long as previous trials, the CETP inhibitor anacetrapib was convincingly found to reduce coronary events. While ongoing monitoring and a careful review of safety is essential, anacetrapib could find its way to clinical use as another option for patients with coronary disease who require further reduction in LDL and/or non-HDL cholesterol.

SOURCES OF FUNDING
ART was supported by NIH grant HL107653 and DJR by NIH grant HL111398.

DISCLOSURES
Alan Tall: Consulting: Dalcor, Amgen, CSL, Equity: Staten Biotechnology.
REFERENCES


Figure Legend. CETP inhibition reduces LDL-C levels by three mechanisms, 1) decreased transfer of HDL CE into triglyceride-rich lipoproteins which are converted into LDL; 2) decreased transfer of HDL CE into LDL; and 3) increased uptake of LDL particles by the hepatic LDL receptor.
Liver

LDL particle uptake

TRL

CE

HDL

LDL

CE

FIGURE 1
The Trials and Tribulations of CETP Inhibitors
Alan R Tall and Daniel J Rader

_Circ Res._ published online October 10, 2017;
_Circulation Research_ is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2017 American Heart Association, Inc. All rights reserved.
Print ISSN: 0009-7330. Online ISSN: 1524-4571

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://circres.ahajournals.org/content/early/2017/10/09/CIRCRESAHA.117.311978

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in _Circulation Research_ can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to _Circulation Research_ is online at:
http://circres.ahajournals.org//subscriptions/