Potent peroxisome proliferator-activated receptor-\(\alpha\) agonist treatment increases cholesterol efflux capacity in humans with the metabolic syndrome

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Aims
Fibrate medications weakly stimulate the nuclear receptor peroxisome proliferator-activated receptor-\(\alpha\) (PPAR-\(\alpha\)) and are currently employed clinically in patients with dyslipidaemia. The potent and selective agonist of PPAR-\(\alpha\) LYS18674 is known to substantially increase apolipoprotein A-I (apoA-I) turnover without major impact on steady-state levels of apoA-I or high-density lipoprotein-cholesterol (HDL-C). We sought to determine whether therapy with a PPAR-\(\alpha\) agonist impacts cholesterol efflux capacity, a marker of HDL function.

Methods and results
Cholesterol efflux capacity was measured at baseline and after 8 weeks of therapy in a randomized, placebo-controlled trial involving participants with metabolic syndrome treated with either LYS18674 100 \(\mu\)g daily \((n = 13)\) or placebo \((n = 15)\). Efflux capacity assessment was quantified using a previously validated \textit{ex vivo} assay that measures the ability of apolipoprotein-B depleted plasma to mobilize cholesterol from macrophages. LYS18674 led to a 15.7% increase from baseline (95% CI 3.3–28.1%; \(P = 0.02\), \(P\) vs. placebo = 0.01) in efflux capacity. The change in apoA-I production rate in the active treatment arm was strongly linked to change in cholesterol efflux capacity \((r = 0.67, P = 0.01)\).

Conclusions
Potent stimulation of PPAR-\(\alpha\) leads to accelerated turnover of apoA-I and an increase in cholesterol efflux capacity in metabolic syndrome patients despite no change in HDL-C or apoA-I levels. This finding reinforces the notion that changes in HDL-C levels may poorly predict impact on functionality and thus has implications for ongoing pharmaco-logic efforts to enhance apoA-I metabolism.

Keywords
Cholesterol efflux capacity • HDL-cholesterol • Lipid metabolism • PPAR-\(\alpha\)

Translational perspective
The complexities of high-density lipoprotein cholesterol (HDL-C) metabolism have led to interest in moving beyond static assessments of HDL-C levels to assessing functionality. One such measure, cholesterol efflux capacity, measures a key step in the reverse cholesterol transport pathway and is associated with both the prevalence and incidence of coronary disease. Here, we demonstrate that treatment of metabolic syndrome patients with a peroxisome proliferator-activated receptor-\(\alpha\) agonist is associated with improvement in cholesterol efflux capacity despite no change in HDL-C levels. This increase was closely linked to an increase in the rate of apolipoprotein A-I production rate. This finding may inform ongoing efforts to evaluate and develop efficacious therapeutics targeting HDL metabolism.

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Introduction

Therapeutic targeting of high-density lipoprotein cholesterol (HDL-C) metabolism has proven challenging with multiple clinical trial setbacks in recent years involving niacin or cholesteryl ester transfer protein inhibition. Steady-state assessment of circulating HDL concentrations may incompletely reflect in vivo functionality. Cholesterol efflux capacity quantifies the ability of HDL lipoproteins to mobilize cholesterol from macrophages, a critical first step in the anti-atherogenic reverse cholesterol transport pathway. This metric has been shown to be inversely related to both atherosclerotic burden and, more recently, incident cardiovascular events in multiple cohorts independent of circulating levels of HDL-cholesterol.1,2

Peroxisome proliferator-activated receptors (PPARs) are a family of nuclear receptors that modulate both lipid and glucose metabolism. Fibrate therapies serve as weak activators of PPAR-α and are in frequent clinical use in patients with elevated triglycerides. Subsequent efforts have led to more potent and specific PPAR-α ligands, including LY518674. Previous studies with LY518674 in patients with atherogenic dyslipidaemia or the metabolic syndrome has noted decreased triglycerides but increased LDL-C levels and minimal impact on HDL-C or apoA-I levels.3,4 However, a balanced >30% increase in both the production and catabolic rate was noted, reflective of enhanced apoA-I turnover.

Prior efforts to document a change in cholesterol efflux capacity with LY518674 using murine bone marrow-derived macrophages did not show a significant impact. The present study reassessed efflux capacity with a more recently validated assay using the J774 macrophage cell line that may be better suited to clinical samples.

Materials and methods

The study population was derived from a previously described randomized controlled trial that investigated the impact of LY518674 on HDL metabolism (ClinicalTrials.gov NCT00327002). All subjects had low HDL-C levels as well as at least two additional components of the metabolic syndrome. Exclusion criteria included treatment with fibrates, thiazolidinediones, ezetimibe, or niacin (>250 mg/day) as well as a history of cardiovascular disease or diabetes. Participants were randomized in a double-blind fashion to receive LY518674, 100 µg daily, or placebo for 8 weeks. Apolipoprotein kinetics were measured using a deuterated leucine tracer to quantify rate of apoA-I production (i.e. the amount of newly synthesized apoA-I entering plasma).4

Cholesterol efflux capacity was assessed using an assay that quantifies the ability of apolipoprotein B-depleted plasma to accept 3H-radiolabeled cholesterol from J774 macrophages ex vivo as previously reported.1 Efflux capacity assays were performed in duplicate in a paired fashion on 24-well plates.

Paired t-tests were used to assess the effect of LY518674 and placebo on HDL metabolic parameters. These changes were compared across treatment arms via an analysis of covariance, which included the patient’s baseline value and the treatment group as covariates.

Results

The study population included 28 patients, with 13 randomized to LY518674 and 15 to placebo. Baseline characteristics were well-balanced across randomization groups as previously reported. The cohort included 14 males (50%) with a mean age of 49 years, body mass index of 37 kg/m², and blood pressure of 130/78 mmHg. Baseline laboratory values, expressed as mean ± SD, were total cholesterol 189 ± 43 mg/dL, HDL-C 37 ± 6 mg/dL, LDL-C 117 ± 36 mg/dL, and apoA-I 108 ± 15 mg/dL. Median (IQR) triglyceride value was 167 (104–211) mg/dL.

Cholesterol efflux capacity at baseline was similar across the two treatment arms with mean ± SD of 1.08 ± 0.15 and 1.07 ± 0.22 for the LY518674 and placebo groups, respectively. Minimal relationship (r = 0.17; P = 0.38) was noted between this efflux assessment and previously reported total efflux capacity, likely reflective of differences in assay technique. Eight weeks of therapy with LY518674 were associated with a 15.7% (95% CI 3.3–28.1%) increase from baseline in cholesterol efflux capacity and a 31.1% (95% CI 15.3–46.9%) increase from baseline in the production rate of apoA-I despite no change in HDL-C or apoA-I levels (Table 1). Neither change in HDL-C (r = 0.17; P = 0.66) nor change in apoA-I (r = 0.17; P = 0.66) was predictive of change in efflux capacity with PPAR-α agonist treatment. However, change in apoA-I production rate strongly predicted increased cholesterol efflux capacity (r = 0.67; P = 0.01) as displayed in Figure 1.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>LY518674</th>
<th>Placebo</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>HDL-C</td>
<td>–0.8 (–13.2 to 11.7)</td>
<td>–3.2 (–7.5 to 1.2)</td>
<td>0.31</td>
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<tr>
<td></td>
<td>P = 0.96</td>
<td>P = 0.08</td>
<td></td>
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<tr>
<td>ApoA-I</td>
<td>0.7 (–9.6 to 8.2)</td>
<td>5.6 (20.0 to 9.3)</td>
<td>0.26</td>
</tr>
<tr>
<td></td>
<td>P = 0.81</td>
<td>P = 0.01</td>
<td></td>
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<tr>
<td>Apolipoprotein A-I production rate</td>
<td>31.1 (15.3 to 46.9)</td>
<td>–0.4 (–6.5 to 5.7)</td>
<td>0.0001</td>
</tr>
<tr>
<td></td>
<td>P = 0.001</td>
<td>P = 0.80</td>
<td></td>
</tr>
<tr>
<td>Cholesterol efflux capacity</td>
<td>15.7 (3.3 to 28.1)</td>
<td>–0.2 (–5.1 to 4.8)</td>
<td>0.01</td>
</tr>
<tr>
<td></td>
<td>P = 0.02</td>
<td>P = 0.87</td>
<td></td>
</tr>
</tbody>
</table>

Values represent mean % change (95% CI) for each parameter. P-values are presented for the significance of both change from baseline and difference between treatment arms.
Discussion

Potent and specific stimulation of the PPAR-α receptor leads to an increase in cholesterol efflux capacity, an effect that is associated with the enhanced production of apoA-I, in patients with metabolic syndrome. This finding likely reflects an increase in hepatic generation of lipid poor apoA-I particles that serve as efficient acceptors of cholesterol. Our results are well aligned with a murine study that noted enhanced macrophage reverse cholesterol transport after treatment with potent PPAR-α activation. Although the clinical development of novel PPAR agonists with an acceptable safety profile has proven challenging, several molecules remain in clinical development.

The discrepancy between the current findings and previously reported cholesterol efflux values likely reflects differences in design of the cell-based assay. The prior study quantified efflux to whole plasma from bone marrow-derived murine macrophages loaded with acetylated LDL and treated with a liver X receptor agonist. Our present assay implemented apolipoprotein-B depleted serum with acetylated LDL and treated with a liver X receptor agonist. The prior study quantified efflux to whole plasma from bone marrow-derived murine macrophages loaded with acetylated LDL and treated with a liver X receptor agonist.

Figure 1  Relationship between change in apoA-I production rate and change in cholesterol efflux capacity after 8 weeks of therapy with either LY518674 or placebo.

degree of correlation ($r = 0.89$) between baseline and on-treatment values, again confirming longitudinal stability.

The current findings represent another example of discordance between changes in HDL-C levels and functionality with pharmacologic therapy. For example, the addition of niacin to statin therapy had no impact on efflux capacity despite resulting in a 29% increase in HDL-C. The cholesteryl ester protein inhibitor dalcetrapib increased HDL-C levels by 34% with only a 10% increase in total efflux capacity. These assessments may prove fruitful in offering additional mechanistic understanding to the multiple HDL and apoA-I centric therapeutic candidates currently in development.

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References