Lowering triglycerides to modify cardiovascular risk: will icosapent deliver?

Abstract: Despite the clinical benefits of lowering levels of low-density lipoprotein cholesterol, many patients continue to experience cardiovascular events. This residual risk suggests that additional risk factors require aggressive modification to result in more effective prevention of cardiovascular disease. Hypertriglyceridemia has presented a considerable challenge with regard to understanding its role in the promotion of cardiovascular risk. Increasing evidence has established a clear causal role for elevated triglyceride levels in vascular risk. As a result, there is increasing interest in the development of specific therapeutic strategies that directly target hypertriglyceridemia. This has seen a resurgence in the use of omega-3 fatty acids for the therapeutic lowering of triglyceride levels. The role of these agents and other emerging strategies to reduce triglyceride levels in order to decrease vascular risk are reviewed.

Keywords: hypertriglyceridemia, omega-3 fatty acid, fish oil, cardiovascular risk, lipids

Introduction

The role of low-density lipoprotein cholesterol (LDL-C) in cardiovascular disease is well established, with LDL-C lowering being the cornerstone of preventive strategies. This is evidenced from statin trials, in which the degree of LDL-C lowering directly associated with their clinical benefit. However, many cardiovascular events continue to occur even in patients who achieve low LDL-C levels, reflecting a substantial residual risk. In the search to identify factors associated with persistent cardiovascular risk, it is clear that LDL-C alone does not fully reflect the circulating burden of atherogenic lipoproteins. In particular, triglyceride-rich particles (very low-density lipoproteins [VLDL], intermediate density lipoproteins, chylomicrons) play an important role in the promotion of the atherosclerotic disease process and are poorly reflected by conventional LDL-C measures. It is therefore not surprising that levels of non-high-density lipoprotein cholesterol (non-HDL-C), which reflect the entire burden of atherogenic lipid particles, associate with vascular risk and have become increasingly used as a secondary target for modification in treatment guidelines. As the incidence of abdominal obesity, metabolic syndrome, and type 2 diabetes mellitus rises, associated triglyceride-rich lipoproteins are likely to have a greater impact on promoting vascular risk.

Population studies of hypertriglyceridemia and cardiovascular risk

Elevated triglyceride levels have become increasingly prevalent in the community. Follow-up of the National Health and Nutrition Examination Survey has found that
32.2% of individuals have elevated triglyceride levels, including 1.7% with levels considered severe (500–2,000 mg/dL). While early studies failed to convincingly establish an association between triglyceride levels and cardiovascular disease, more recent data, including cohort studies and meta-analyses, have shown that increases in both fasting and nonfasting triglyceride levels directly associate with cardiovascular risk.

Further analysis has revealed that the strength of the association between triglyceride levels and cardiovascular risk may differ between males and females, with evidence that each 1 mmol/L increase in triglyceride associates with a 32% and 76% increase in risk, respectively. While the degree of this association is attenuated after controlling for other metabolic risk factors, a greater impact in females continues to be observed. In clinical trials of statins, elevated triglyceride levels identify patients at a higher risk of cardiovascular events, despite use of high-intensity statin therapy.

Generation of triglyceride-rich lipoproteins

Hypertriglyceridemia results from a number of sources, including dietary fat consumption and generation of chylomicron and remnant particles, hepatic synthesis of VLDL, and impaired metabolic clearance of these particles from the circulation. Triglyceride-rich lipoproteins undergo hydrolysis by lipoprotein lipase within capillary beds in adipose, cardiac, and skeletal muscle tissue beds to release nonesterified fatty acids for local use, generating remnant particles within the systemic circulation. These remnant particles, specifically VLDL remnants, contain substantial cholesterol due to increased activity of cholesteryl ester transfer protein in hepatic triglyceridemia favoring transfer to VLDL rather than LDL particles. Reduced size of these remnant particles renders them more likely to enter and be retained within the artery wall. In this location, they play a significant role in formation of foam cells, the major cellular component of the evolving atherosclerotic plaque.

Apolipoprotein C (apoC)-III resides on VLDL, LDL, chylomicrons, and HDL particles within the circulation. Increased apoC-III production is a characteristic feature of patients with insulin resistance and hypertriglyceridemia, by virtue of its role in the inhibition of lipolytic activity and hepatic uptake of triglyceride-rich lipoproteins. Increasing evidence implicates apoC-III in the promotion of inflammatory pathways within the artery wall, suggesting that it may influence atherogenesis via lipid-mediated and lipid-independent processes. Further evidence that these factors involved in the metabolism of triglyceride-rich lipoproteins play an important role in atherosclerosis comes from genomic and biomarker studies demonstrating their association with cardiovascular events.

Treatment of hypertriglyceridemia

Lifestyle modification

The initial effort to lower triglyceride levels should begin with lifestyle modification and improved glycemic control in diabetes, with evidence that these measures can have rapid and profound beneficial effects. Weight loss is associated with consistent reductions in triglyceride levels across multiple studies and is an essential part of treatment for hypertriglyceridemia. The relative impact of specific diets and exercise regimens on achieving weight loss has had variable reports with regard to their effects on triglyceride levels.

High-fat content diets lead to obesity and increased visceral adiposity, which is associated with insulin resistance. While a low-fat diet would intuitively seem the best dietary approach, there are some reports that comparisons of moderate-fat content diets with low-fat diets actually show that moderate fat intake elicits a more favorable effect on triglyceride levels. High-carbohydrate diets are associated with increased fatty liver changes, which are associated with increased hepatic triglyceride-rich lipoprotein production in proportion to the degree of liver fat. Low-carbohydrate diets consistently lower triglycerides, more so in those with higher baseline levels, and when integrated within Mediterranean diets appear to be superior in their triglyceride-lowering effect than low-fat diets.

Exercise increases lipoprotein lipase activity with modest triglyceride lowering. While smoking cessation can often be associated with weight gain, its benefit in terms of triglyceride lowering is still evident, given the association of insulin resistance and postprandial hypertriglyceridemia in smokers. There appears to be a U-shaped relationship between amount of alcohol intake and triglyceride levels, so hypertriglyceridemic patients should be encouraged to only drink alcohol in moderation. For patients presenting with marked hypertriglyceridemia, cessation of alcohol consumption, restriction of dietary fat intake, and more aggressive glycemic control in diabetic patients can each have a rapid and profound effect on triglyceride levels.

Statins

Statins lower triglycerides in a dose-dependent manner, particularly in the hypertriglyceridemia patient. Given their
benefit in terms of cardiovascular event rate reduction, treatment guidelines recommend intensification of LDL-C lowering with statins as the first pharmacological treatment choice in hypertriglyceridemic patients. The persistence of elevated triglyceride levels identifies patients with a residual clinical risk in statin trials, suggesting an ongoing need to develop additional therapeutic strategies for these patients.

**Fibrates**

Fibrates are modest pharmacological agonists of peroxisome proliferator-activated receptor-α (PPAR-α), a transcription factor and major regulator of lipid metabolism. Their major influences on plasma lipids include reducing triglycerides and increasing HDL cholesterol. Their lipid effects appear to be mediated by a reduction in apoC-III expression and increases in lipoprotein lipase activity, hepatic synthesis of the major apolipoproteins (apoA-I, apoA-II) carried on HDL particles, and oxidation of fatty acids, which in turn lead to a decrease in the rate of hepatic lipogenesis. While studies have shown that fibrates retard progression of angiographic disease, the major clinical outcome trials of fibrates have demonstrated variable effects on cardiovascular events. While studies of gemfibrozil have demonstrated a favorable effect, it can be difficult to administer in combination with statin therapy. More recent studies of fenofibrate in statin-treated patients with diabetes failed to demonstrate a benefit in terms of the primary endpoint, although subsequent analyses have shown a favorable impact on hard clinical endpoints (death, myocardial infarction, stroke) in patients with a prior myocardial infarction and on the microvascular complications of diabetes. While triglyceride lowering does not appear to contribute to any potential benefit of fibrate therapy, meta-analyses have demonstrated that it is the patient with elevated triglyceride levels at baseline in whom they are most likely to be protective. Recent attempts to develop more potent PPAR-α and dual PPAR-α/γ agonists have failed to demonstrate incremental efficacy and have raised additional safety concerns.

**Niacin**

Niacin has a range of lipid-modifying effects, including substantial raising of HDL-C in addition to lowering of triglycerides, LDL-C, and lipoprotein(a). While the mechanism underlying its lipid effects remains to fully elucidated, triglyceride lowering is likely to result from inhibition of diacylglycerol acyltransferase 2. Early trials prior to the use of statins demonstrated reductions in cardiovascular events in survivors of myocardial infarction with immediate-release formulations of niacin administered as monotherapy and in combination with clofibrate. Subsequent arterial wall imaging studies have demonstrated that niacin has a favorable impact on progression of atherosclerotic disease within the coronary and carotid vasculature. However, niacin use is limited by near ubiquitous experience of flushing mediated by prostaglandin-mediated vasodilation. This prevents appropriate dose escalation required to achieve lipid effects. Multiple efforts have been undertaken to improve tolerance of niacin, including use of extended-release formulations and administration in combination with a selective prostaglandin-2 receptor inhibitor, laroniprant. However, while studies of these approaches in statin-treated patients have not demonstrated clinical efficacy, suggesting that niacin is not likely to be a widely used approach for lipid modification moving forward.

**Omega-3 fatty acids in hypertriglyceridemia**

Considerable data, ranging from studies of fish consumption in populations through to supplementation studies have consistently demonstrated that omega-3 (n-3) fatty acids lower triglyceride levels by 10%–30% in patients with hypertriglyceridemia, depending on the amount of n-3 fatty acid administered and the baseline triglyceride level. The major n-3 fatty acids, eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA), modulate each of the major nuclear receptors that regulate triglyceride levels, including liver X receptor (LXR), farnesol X receptor (FXR), hepatocyte nuclear factor-4α, and PPAR α, β, and γ.

These receptors play an important role in regulation of triglyceride-rich lipoproteins and therefore present a target for therapeutic modification by n-3 fatty acids. LXR regulates expression of sterol regulatory element-binding protein-1c (SREBP-1c), which plays a major role in transcription of fatty acid-synthesizing enzymes. The n-3 fatty acids inhibit LXR from binding to the LXR response element and reducing SREBP-1c transcription and act directly on SREBP-1c, inhibiting its maturation. FXR plays a regulatory role in the enterohepatic circulation of lipids, with it binding and being activated by several bile acids, with stimulation increasing expression of short heterodimer protein, which via heterodimerization has multiple effects on bile acids, reducing transcription of SREBP-1c, upregulating expression of PPAR-α, and decreasing hepatic secretion of...
triglyceride-containing lipoproteins. The n-3 fatty acids, via their FXR interaction, can differentially regulate expression of FXR targets. While n-3 fatty acids interact with all PPAR species, they have a clear preference for the α-subtype, decreasing triglycerides in a similar fashion to fibrates. Hepatocyte nuclear factor-α has multiple important roles in hepatic regulation of lipid metabolism, carbohydrate metabolism, hematopoiesis, and blood coagulation. It plays an important role in the regulation of hepatic expression of apolipoproteins (A-I, A-II, B, C-II, C-III) and microsomal transfer protein. n-3 fatty acid stimulation of hepatocyte nuclear factor-α leads to a reduction of apoC-III expression. EPA, but not DHA, inhibits activity of diacylglycerol acyltransferase, an enzyme that regulates the terminal and rate-limiting step of hepatic triglyceride synthesis. Accordingly, n-3 fatty acids play an important and diverse role in influencing a range of factors implicated in the regulation of triglyceride levels.

**Prescription n-3 fatty acid formulations**

A range of over-the-counter preparations of n-3 fatty acids, often under the label of fish oil, are widely used in population for a range of indications, including lowering of plasma triglyceride levels. However, the finding that large doses of EPA and DHA, in the range of 3–4 g daily, are often required to produce meaningful triglyceride lowering in hypertriglyceridemic patients, presents a challenge for the use of many of these preparations. The actual content of EPA and DHA in over-the-counter preparations is often low, requiring individuals to consume large numbers of capsules on a daily basis (often up to 12 daily) to achieve effective triglyceride lowering. Such findings have led to the production of pharmaceutical grade n-3 fatty acids with much higher concentrations (85% of their content were ethyl esters of EPA and DHA in comparison with most standard fish oil preparations containing 18% EPA and 12% DHA). The first prescription n-3 fatty acid preparation (Lovaza®, initially marketed as Omacor®) approved for use as an adjunct to diet in the treatment of severe hypertriglyceridemia contained more than 90% of n-3 fatty acid ethyl esters (465 mg EPA, 375 mg DHA). Administration for up to 24 weeks in patients with severe hypertriglyceridemia demonstrated reductions in triglyceride levels by 19%–47%. In parallel, early observations of increases in LDL-C levels were reported, the extent of which depended on the baseline triglyceride level. It remains unknown whether this increase in LDL-C might have an adverse effect on the artery wall. It has been postulated by some that DHA plays a more prominent role in LDL-C elevation, with meta-analyses of both individual and comparative studies revealing increases in LDL-C in 75% of DHA-treated and 40% of EPA-treated patients. While the mechanism for the LDL-C increase remains to be fully elucidated, it has been proposed that this is likely to involve accelerated conversion of VLDL to intermediate density lipoprotein and LDL particles.

**Icosapent**

Icosapent is the first EPA only prescription n-3 fatty acid (containing 96% of the ethyl ester of EPA) to receive regulatory approval. It undergoes de-esterification during intestinal absorption and enters the systemic circulation via the thoracic duct, achieving its peak plasma concentration 5 hours following ingestion. Icosapent is >99% protein bound in the plasma, has a volume of distribution of 88 liters, and displays predictable linear pharmacokinetics. It is currently approved for use as an adjunct to diet to reduce triglyceride levels in adult patients with severe (≥500 mg/dL) hypertriglyceridemia. There are ongoing studies assessing its efficacy in reducing the risk of cardiovascular events in high-risk patients with mixed dyslipidemia.

A multicenter, placebo-controlled, randomized, double-blind 12-week study with an open-label extension (MARINE) was conducted in patients with very high triglyceride levels, with randomization to either 2 g or 4 g daily of icosapent or placebo for 12 weeks. The treatment period was preceded by a 4–6-week run-in period, where subjects were counseled on a lifestyle changes diet to comply with throughout the trial. Subjects with triglyceride levels of between 500 mg/dL and 2,000 mg/dL following the run-in period qualified for randomization. The major exclusion criteria included superimposed obesity (body mass index >45 kg/m²), poorly controlled diabetes, uncontrolled thyroid disease, nephritic syndrome range proteinuria (>3 g/day), and heavy alcohol intake. Twenty-five percent of the subjects included in the randomization were treated with a statin. Significant placebo-adjusted reductions in triglycerides were observed in both the 4 g (−33.1%, $P<0.0001$) and 2 g (−19.7%, $P=0.005$) groups. In those individuals with baseline triglyceride levels ≥750 mg/dL, greater reductions were observed in the 4 g (−45.4%, $P=0.0001$) and 2 g (−32.9%, $P=0.002$) groups. Associated reductions in non-HDL-C, VLDL, and lipoprotein phospholipase A2 were also observed, with greater effects noted in the 4 g group. Significant LDL-C elevations were not observed in this study, although the exploratory analysis was underpowered. Icosapent was well tolerated, with similar rates of adverse events (most commonly nausea, diarrhea, and eructation) observed in both groups.
The ANCHOR study was a randomized, double-blind, placebo-controlled study of patients with a high cardiovascular disease risk and adequate LDL-C control (40–115 mg/dL) with statins and ezetimibe, but with high triglyceride levels (185–500 mg/dL). A total of 702 patients were randomized to 2 g or 4 g of icosapent or placebo daily for a 12-week treatment period, following a run-in period similar to that of the MARINE study. The study included a prespecified analysis to demonstrate noninferiority of LDL-C change compared with placebo. The key exclusion criteria for the trial were supermorbid obese patients with a body mass index > 45 kg/m², poorly controlled diabetics, and subjects treated with fribrates or niacin. Significant reductions in triglyceride levels were observed in both the 4 g (~21.5%, P < 0.0001) and 2 g (~10.1%, P = 0.0005) groups. Similar reductions in associated lipoproteins were observed as reported in MARINE, with the additional finding of a significant reduction in C-reactive protein in the 4 g group. Noninferiority compared with placebo with regards to LDL-C changes was observed in both active treatment groups. Icosapent was well tolerated, with similar adverse event and discontinuation rates reported in all three treatment groups.

Outcome trials with EPA
JELIS (the Japan EPA Lipid Intervention Study) was a prospective, randomized, open-label, blinded endpoint trial of 18,645 patients with elevated serum total cholesterol (~250 mg/dL), but notably relatively normal triglyceride levels, treated with EPA 1.8 g or placebo daily in addition to statin therapy. Treatment with EPA was associated with a greater lowering of triglycerides (~9% versus ~4%, P < 0.0001) and a 19% reduction in the composite primary endpoint of any major coronary event, including sudden cardiac death, fatal or nonfatal myocardial infarction and nonfatal events, unstable angina determined to be caused by myocardial ischemia on invasive or noninvasive testing and requiring emergent hospitalization. This will determine whether high-dose administration of EPA has a cardiovascular benefit in high-risk statin-treated patients with modest hypertriglyceridemia.

Conclusion
Hypertriglyceridemia at severe levels is an important risk factor for pancreatitis, and persistently high triglyceride levels despite optimal LDL-C control conveys increased residual risk of cardiovascular disease. There is a clear need for a therapeutic option in hypertriglyceridemia that can be used in addition to statin therapy to further reduce the cardiovascular disease risk and burden. Icosapent shows a clear benefit in the treatment of severe hypertriglyceridemia and has also shown efficacy in the treatment of patients with more moderate elevations in serum triglyceride levels already treated with statins. However, favorable effects on lipid biomarkers, particularly those beyond LDL-C, have not been easily demonstrated to translate to a beneficial effect on cardiovascular events. Accordingly, the result of REDUCE-IT will be eagerly awaited to determine whether icosapent will be widely used or remain as a treatment for severe hypertriglyceridemia alone.

Disclosure
The authors report no conflicts of interest in this work.

References


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