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The Arachidonic Acid Metabolome Serves as a Conserved Regulator of Cholesterol Metabolism

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SUMMARY

Cholesterol metabolism is closely interrelated with cardiovascular disease in humans. Dietary supplementation with omega-6 polyunsaturated fatty acids including arachidonic acid (AA) was shown to favorably affect plasma LDL-C and HDL-C. However, the underlying mechanisms are poorly understood. By combining data from a GWAS screening in >100,000 individuals of European ancestry, mediator lipidomics, and functional validation studies in mice, we identify the AA metabolome as an important regulator of cholesterol homeostasis. Pharmacological modulation of AA metabolism by aspirin induced hepatic generation of leukotrienes (LTs) and lipoxins (LXs), thereby increasing hepatic expression of the bile salt export pump Abcb11. Induction of Abcb11 translated in enhanced reverse cholesterol transport, one key function of HDL. Further characterization of the bioactive AA-derivatives identified LX mimetics to lower plasma LDL-C. Our results define the AA metabolome as conserved regulator of cholesterol metabolism, and identify AA derivatives as promising therapeutics to treat cardiovascular disease in humans.

INTRODUCTION

Atherosclerosis is still the leading cause of death in industrialized countries, and novel therapies to lower low-density lipoprotein...
cholesterol (LDL-C) are urgently needed. Additionally, any approach promoting the transport of excess cholesterol from plaque macrophages back to the liver via plasma high-density lipoprotein (HDL) for biliary and final fecal excretion is expected to prevent atherosclerosis, a mechanistic concept called reverse cholesterol transport (RCT) (Cuchel and Rader, 2006; Degoma and Rader, 2011; Rader and Daugherty, 2008). It is well known that dietary supplementation with omega-6 polyunsaturated fatty acids (omega-6 PUFAs) including arachidonic acid (AA) reduces the risk of cardiovascular disease (CAD) (Harris et al., 2009; Katan, 2009), which is in part attributable to the observation that increased AA plasma levels are associated with beneficial changes in LDL-C and HDL-C.

In humans, AA is metabolized into many potent bioactive compounds, such as (1) prostaglandins (PGs) and thromboxanes (TXs), (2) leukotrienes (LTs), and (3) lipoxins (LXs). Whereas PGs and TXs are formed by cyclooxygenases I and II (COX I/II), LTs are generated through the action of arachidonate 5-lipoxygenase (ALOX5), and LXs—an acronym of lipoxygenase interaction products—by the sequential cell-cell interaction of different lipoxygenases (McMahon and Godson, 2004; Serhan, 2007): LTA₄, the intermediate of LT synthesis, is produced in neutrophils via ALOX5 and can be taken up by platelets and converted into LXs via ALOX12. 15S-Hydroxyeicosatetraenoic acid (15S-HETE) is synthesized in epithelial cells and monocytes via ALOX15, which can be further converted into LXs in leukocytes by ALOX5. Generation of LXs occurs also when 15-HETE accumulates in cell membranes of neutrophils, where it is converted into LXs (McMahon and Godson, 2004; Serhan, 2007). An additional route of LX biosynthesis emerges in cells exposed to aspirin. Aspirin acetylates COX II, changing its activity to a lipoxygenase. This generates 15R-HETE, which is finally converted into 15-epi-lipoxins via ALOX5 (McMahon and Godson, 2004; Serhan, 2007).

To date, the relative pathophysiological roles of lipoxygenases, LTs, and LXs have been extensively studied in inflammation where LTB₄ exerts proinflammatory actions by promoting the recruitment of leukocytes to the site of insult. This is followed by an increase in anti-inflammatory eicosanoids LXA₄ and its regioisomer LXB₄, which mediate resolution of inflammation and failure of its resolution is found in atherosclerosis (Serhan, 2007). One important example of sustained chronic inflammation and failure of its resolution is presented significantly lower levels of HDL-C compared to homozygous carriers of the common TT allele (allele frequency of 0.65) of lead SNP rs12765320 within the ALOX5 gene showed a dose-dependent decrease in plasma HDL-C levels (ES = −0.429 mg dl⁻¹ per copy of T allele; Figure 1B). The reported association of rs12765320 with HDL-C in the GWAS study was independently replicated in the smaller LUDwigshafens Risk and Cardiovascular Health (LURIC) (Winkelmann et al., 2001) cohort comprising 2,095 individuals (HDL-C 37.23 ± 10.69 mg dl⁻¹ versus 36.32 ± 9.42 mg dl⁻¹, male homozygous carriers of the CC and the TT alleles, respectively; p < 0.05).

To further support the association between ALOX5 and HDL-C, a second SNP (rs7068039) belonging to the haplotype block (Table S1) was genotyped in 2,141 individuals from the PLIC (Progressione Della Lesione Intimale Carotidea) study (Norata et al., 2010). Homozygous carriers of the common TT allele presented significantly lower levels of HDL-C compared to homozygous CC carriers (55.11 ± 14.87 mg dl⁻¹ versus 58.78 ± 13.88 mg dl⁻¹, respectively; p < 0.05) (Figure 1C), thus replicating this association in a second independent cohort. Next, we characterized whether the differences in HDL-C levels also translate into alterations of HDL function. Therefore, cholesterol efflux capacity of apoB-depleted serum from age- and independent studies is, however, warranted to fully support the genetic association between ALOX5 and CAD.

So far, neither the relative role of different AA-metabolizing enzymes, including lipoxygenases, nor the potential impact of their main metabolites, i.e., LTs and LXs, on cholesterol metabolism has been systematically investigated. Here, we elucidate the relative roles of LTs and LXs on cholesterol homeostasis by combining data from GWAS analysis in humans, mediator lipidomics, and loss-of-function studies in mice. We identify LX mimetics as promising therapeutics to lower plasma LDL-C and to treat atherosclerosis.
sex-matched TT and CC rs7068039 carriers was measured. Homozygous TT allele carriers showed significantly reduced cholesterol efflux capacity compared to CC carriers (Figure 1D). In summary, GWAS screening in humans helped to identify associations between variants within ALOX5 not only with HDL-C mass but importantly also with HDL function.

Aspirin Treatment Promotes RCT

From the GWAS data it cannot be inferred whether the newly identified variants may regulate expression of arachidonate lipoxigenases and whether differential activation of arachidonate lipoxigenases may impact plasma cholesterol levels or affect functional properties of lipoproteins such as RCT. Therefore, we performed functional validation studies in mice using a systematic approach: (1) simultaneous activation of Alox5 and Alox12/15, (2) selective knockout of Alox5, and (3) knockout of Alox12/15. In contrast to humans, who express at least four isoforms of ALOX15, mice do not have separate 12- and 15-arachidonate lipoxigenases, but rather a combined 12/15-lipoxigenase with variable positional specificity for both the 12-position and the 15-position of AA (Kühn and O’Donnell, 2006).

In our first in vivo experiments, we simultaneously induced the processing of AA by Alox5 and Alox12/15 through pharmacological inhibition of Cox I/II, which shifts the biosynthetic pathways of the AA metabolome toward the formation of LXs and LTs in murine systems (Figure 2A) (Brink et al., 2003; Spite and Serhan, 2010). Inhibition of Cox I/II in mice was achieved by systemic treatment with aspirin (Tancevski et al., 2006), and in vivo macrophage-to-feces RCT studies were performed as described (Tancevski et al., 2010; Zhang et al., 2003); after intraperitoneal injection of [3H]-cholesterol-labeled J774 macrophages, the tracer was measured in plasma and feces (Figure 2B). Aspirin-treated mice had significantly decreased plasma [3H]-cholesterol levels 24 hr postinjection (Figure 2C), which was caused either by enhanced uptake of [3H]-HDL-C into liver and/or by increased biliary transport of sterols. Hepatic protein expression of the HDL receptor (scavenger receptor BI, SR-BI) and of the LDL receptor (LDLr) were unaffected in aspirin-treated mice (Figure S3A), making the hypothesis of enhanced cholesterol clearance from plasma rather unlikely. Accordingly, plasma total cholesterol levels as well as HDL-C levels were unchanged in aspirin-treated mice (Figures S3B and S3C), which was further confirmed by lipoprotein separation analysis via fast protein liquid chromatography (FPLC) (Figure 2E).

Aspirin Promotes Excretion of Bile Acids

In the liver, a major part of cholesterol is converted into bile acids (Lefebvre et al., 2009), which are secreted into bile for eventual removal via feces (Lefebvre et al., 2009; Stieger, 2009; Zollner and Trauner, 2009). To better define the type of sterols found to be increased in feces of aspirin-treated mice, a subsequent RCT study was performed, and fecal sterols were extracted into neutral and acidic sterols. We found a marked increase in [3H]-acidic sterols, i.e., bile acids, underlying the observed increase in total counts (relative content of fecal [3H]-acidic sterols, 78% versus 94%, control versus aspirin-treated mice, p < 0.001). Accordingly, in an independent in vivo experiment, Cox I/II inhibition by aspirin was shown to increase the content of fecal bile acid mass per se, as measured by an enzymatic assay (Figure 2F). Plasma levels of 7a-hydroxy-4-cholesten-3-one (C4), a stable plasma marker of bile acid synthesis (Gálman et al., 2003), were not different between aspirin-treated and control mice (Figure S4). In line with unaffected C4 plasma levels, there was no change in hepatic Cyp7a1 mRNA expression, the rate-limiting enzyme for conversion of cholesterol into bile acids (Figure 2G). Moreover, quantitative real-time PCR with reverse transcription (qRT-PCR) analysis revealed no changes in mRNA levels of hepatic neutral sterol transporters Abcg5 and

Figure 1. Identification of ALOX5 as Regulator of HDL-C in Humans

Common variants in the human ALOX5 gene were evaluated in a GWAS meta-analysis comprising >100,000 individuals of European ancestry. (A) Manhattan plot for GWA to HDL-C in the ALOX5 gene, with values of –log _p_ plotted against position on chromosome 10; colors indicate amount of linkage disequilibrium between SNPs; diamonds indicate the identified lead SNPs rs12765320 and rs7068039 within ALOX5. Plots were generated using LocusZoom. (B) Per allele HDL-C relative change in carriers of the lead ALOX5 SNP rs12765320 as derived from the GLGC data set. (C) Per allele HDL-C relative change in carriers of the lead ALOX5 SNP rs7068039 in the PLIC replication cohort. (D) Cholesterol efflux using apoB-depleted serum from 30 CC and 30 TT age- and sex-matched rs7068039. Graphs show mean ± SEM, *p < 0.05.

Arachidonic Acid Metabolome and Cholesterol

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Abcg8 (Figure 2G). Immunoblot analysis showed increased Abcb11 protein levels in livers of aspirin-treated mice (Figure 2H), which constitutes a major determinant of bile flow. On the apical membrane of hepatocytes resides not only Abcb11 but also the multidrug resistance-associated protein 2 (Mrp2 or Abcc2), capable of promoting bile acid excretion (Zollner and Trauner, 2009). Abcb11 mediates excretion of monovalent bile acids, whereas divalent bile acids are exported by Abcc2, which is a conjugate export pump and which has also been discussed as an alternative canalicular bile acid export system in mice (Zollner...
and Trauner, 2009). In contrast to Abcb11, Abcc2 protein levels were found strongly decreased in livers of aspirin-treated mice (Figure 2I). Under physiological conditions, Abcb11 constitutes the rate-limiting step in bile acid transport from the liver into the bile and subsequently intestine (Figge et al., 2004; Stieger, 2009; Stieger and Beuers, 2011; Zollner and Trauner, 2009), and its transgenic overexpression in mice increases bile flow by 30% and fecal bile acid content by more than 40% (Wang et al., 2010). To firmly establish that Cox I/II inhibition by aspirin increases bile acid excretion, [14C]-glycocholic acid was injected into the tail vein of mice, and after 30 min the tracer was quantified in gall bladders of control and aspirin-treated animals (Wang et al., 2001). In confirmation of our data, Cox I/II inhibition by aspirin increased the amount of tracer in bile/C24/2-fold (Figure 2J).

Finally, treatment of Abcb11/C0/C0/C0 mice with aspirin had no effect on macrophage-to-feces RCT (Figures 2K and 2L), proving the hypothesis of an Abcb11-dependent mechanism. Thus, we conclude that aspirin treatment of Abcb11/C0/C0/C0 mice has no effect on macrophage-to-feces RCT (Figures 2K and 2L), proving the hypothesis of an Abcb11-dependent mechanism. Thus, we conclude that aspirin induces regression of atherosclerosis in LDLr/C0/C0/C0 mice.

**Aspirin Induces Regression of Atherosclerosis in LDLr/C0/C0/C0 Mice**

Aspirin is one of the most widely used drugs for primary and secondary prevention of CAD, prescribed to patients at high cardiovascular risk (i.e., diagnosed with CAD, diabetics, etc.). So far, the atheroprotective effect of aspirin was related to its antithrombotic and anti-inflammatory/proresolving properties (Spite and Serhan, 2010). However, the identification of aspirin as a RCT-promoting drug also raised the question of whether aspirin could confer regression of established atherosclerosis. For this purpose, we performed a study in male LDLr/C0/C0/C0 mice fed a western-type diet for 14 weeks, then switched to normal chow and divided into two groups, one receiving placebo and the other receiving aspirin in the drinking water for another 6 weeks. Aspirin treatment led to a 50% reduction in atherosclerotic lesion size, showing that aspirin can lead to the regression of atherosclerosis (Figure 2M).

**Lipidomic Profiling of Aspirin-Treated Mice**

To verify adequate Cox I/II inhibition by aspirin and to analyze changes in intrahepatic levels of LTs and LXs, we next performed qRT-PCR analysis of the genes involved in the biogenesis of these lipids followed by mediator lipidomic analysis (Arita, 2012; Morita et al., 2013). As shown in Figure 3A, and in line with a previous report (Xu et al., 1999), livers of aspirin-treated C57BL/6 mice had reduced Cox2 expression, whereas the expression of arachidonate lipoxigenases was unaffected by aspirin treatment. Aspirin is expected to inhibit Cox I activity, thereby blocking the formation...
of prothrombotic TXA2. This was confirmed by lipidomic profiling, as levels of the metabolite of TXA2, namely TXB2, were dramatically reduced in livers of aspirin-treated mice (Figure 3B). Finally, lipidomic profiling revealed that inhibition of the Cox I/II pathway led to enhanced generation of both LTs (LTB4) and LXs (LXA4) in livers of aspirin-treated animals (Figure 3B). We thus reasoned that the observed increase in hepatic Abcb11 protein expression in mice treated with aspirin may have been conferred by either LTs or LXs, or both.

Leukotrienes and Lipoxins Regulate Abcb11, SR-BI, and LDLr in Hepatocytes

Using primary murine hepatocytes derived from C57BL/6 mice, we confirmed our in vivo studies by demonstrating that both aspirin and its active metabolite salicylic acid induce Abcb11 protein expression. Moreover, aspirin and salicylic acid induced SR-BI protein expression, whereas they downregulated the expression of the LDLr (Figure 4A). Presently, little is known about lipoxygenase pathways within hepatocytes. One main difference between hepatocytes and other cell types is...
that hepatocytes do not express Alox5. In this regard, Claria’s lab showed that Kupffer cells, which are of myeloid origin and express high levels of Alox5, are the major source of LTB4 and LXs in rat liver exposed to aspirin (Planagumà et al., 2002). Accordingly, when comparing mediator lipidomic profiles of mouse liver and of isolated murine hepatocytes, we found that (1) in both settings, aspirin dramatically reduced TXB2 levels as result of efficient Cox I inhibition, (2) in liver extracts, aspirin treatment increased both LTs and LXs, and (3) in isolated mouse hepatocytes, aspirin treatment increased only LX formation (Figure 4B). When analyzing the relative effects of different LX isomers on Abcb11 expression, we found that LXA4, 15-epi-LXA4, and LXB4 increased Abcb11 at 10 nM, with LXB4 showing the strongest induction (Figure 4C). Dose-titration studies as well as coinubcation experiments with LTB4 and LXB4 in primary hepatocytes derived from C57BL/6 mice revealed that both lipid mediators increase Abcb11 expression. Intriguingly, LXB4 decreased Abcb11 protein expression at high dosages. On the other hand, SR-BI and LDLr protein expression was induced mainly by LTB4 (Figures 4D and 4E).

Leukotrienes and Lipoxins Regulate Abcb11 in a Posttranslational Fashion

To further decipher the molecular mechanisms underlying the induction of Abcb11 by aspirin, we performed further studies in primary hepatocytes. The main positive regulator of Abcb11 expression is the transcription factor farnesoid X receptor (Fxr) (Steiger, 2009; Zollner and Trauner, 2009). However, FXR reporter studies ruled out any direct activation of this nuclear receptor by both LTB4 and LXB4 (Figure 4F). In line with this finding, LTs and LXs neither induced the transcription of Abcb11 mRNA (Figure 4G), nor did they influence Abcb11 mRNA degradation (Figure 4H). In addition, blocking protein translation by cycloheximide did not abolish Abcb11 protein expression induced by eicosanoids, suggesting that neosynthesis of protein was not required and that a direct regulation was taking place at a posttranslational level (Figure 4I).

Abcb11 protein expression is known to be stabilized by the mitogen-activated protein kinase (MAPK) p38 (Kubitz et al., 2004), and aspirin, in turn, is known to activate p38 in different cell types, including hepatocytes (Oshima et al., 2008; Trujillo-Murillo et al., 2008). As shown in Figure 4J, blockage of p38 by the selective inhibitor SB 203580 abolished the induction of Abcb11 protein expression by LTB4 and LXB4. To summarize, LTs and LXs increase the expression of Abcb11 by a posttranscriptional and posttranslational mechanism, involving the activity of MAPK p38.

The Role of Alox5 in Cholesterol Metabolism and RCT

Next, we wondered whether knocking out Alox5 would affect hepatic Abcb11 expression, cholesterol homeostasis, and RCT in mice (Figure 5A). As shown in Figure 5B, no marked difference in plasma LDL-C and HDL-C between Alox5+/+ and Alox5−/− mice was observed, whereas VLDL-C levels increased in the knockouts. In macrophage-to-feces RCT experiments, Alox5−/− mice showed reduced plasma tracer levels over 48 hr (Figure 5C) but no significant change in fecal excretion of [3H]-sterols, when compared to Alox5+/+ mice (Figure 5D). Immunoblot analysis revealed an ∼2-fold induction of hepatic Abcb11 and a moderate increase in SR-BI, but no effect on LDLr protein expression (Figure 5E). Finally, lipidomic profiling of livers from Alox5+/+ and Alox5−/− mice showed no changes in LX levels, whereas LTB4 levels were drastically reduced (Figure 5F).

The Role of Alox12/15 in Cholesterol Metabolism and RCT

Lack of Alox12/15 in Alox12/15−/− mice neither affected plasma cholesterol levels nor influenced macrophage-to-feces RCT, when compared to Alox12/15+/+ mice (Figures 6A–6D). Accordingly, no change in expression of hepatic Abcb11, LDLr, and SR-BI was observed in Alox12/15−/− mice (Figure 6E). Mediator lipidomic analysis showed unchanged levels of LXs in livers of Alox12/15−/− mice compared to Alox12/15+/+ mice, whereas LTB4 levels were increased in the knockouts (Figure 6F).

Treatment with Lipoxin Mimetics Lowers Plasma LDL-C

Finally, in an approach to translate our combined findings from GWAS analysis in humans, mediator lipidomics and functional studies in mice into the identification of novel compounds which beneficially influence plasma cholesterol levels and thus putatively the course of CAD, we reasoned to study the impact of systemic treatment with LXB4 on cholesterol metabolism in mice (Figure 7A). Because LXB4 is unstable and rapidly inactivated within the circulation, we performed daily i.v. injections into mice with synthetic stable analogs of LXB4, including 5-(R/S)-methyl-LXB4 and 8,9-acetylenic-LXB4, the latter being a log order of magnitude less potent than LXB4 (Maddox et al., 1998). After 4 days of treatment, 8,9-acetylenic-LXB4 caused a nonsignificant ∼15% decrease in plasma total cholesterol, whereas 5-(R/S)-methyl-LXB4 significantly reduced plasma cholesterol levels by ∼30% (Figure 7B). FPLC analysis showed a decrease in LDL-C in 5-(R/S)-methyl-LXB4-treated mice, whereas HDL-C remained unchanged (Figure 7S). Accordingly, treatment with 5-(R/S)-methyl-LXB4 was associated with an ∼3-fold increase in the protein expression of hepatic LDLr and a moderate increase in hepatic protein levels of SR-BI (Figure 7C). Abcb11 protein expression was decreased in animals treated with LX mimetics compared to controls.

DISCUSSION

Lipoxygenases are lipid peroxidizing enzymes belonging to the nonheme iron dioxygenases family and are conserved across species including bacteria, algae, plants, fish, amphibia, reptilia, and mammals. In animals, the conventional nomenclature classifies lipoxygenases according to their positional specificity of AA oxygenation (Ivanov et al., 2010). Human arachidonate 5-lipoxygenase thus catalyzes oxidation of AA at the 5-position, leading to the formation of hydroxyeicosatetraenoic acids, which can be further metabolized into LTs and LXs through cell-cell interaction. Leukocytes—well-appreciated players in atherosclerosis and myocardial infarction (Drechsler et al., 2010; Dutta et al., 2012)—represent a main source of ALOX5-derived lipid mediators under pathophysiological conditions such as sustained inflammation in atherosclerosis (Spite and Serhan, 2010). Thus ALOX5 is believed to potentially accelerate atherosclerosis by promoting the inflammatory process within the arterial wall.
through increased synthesis of LTs by leukocytes (Peters-Golden and Henderson, 2007). Here, we extend this knowledge by showing that ALOX5 SNPs associate with HDL-C mass and function in human plasma. During the last decade, the old paradigm focusing solely on HDL-C levels as predictor of atherosclerosis changed to a more comprehensive, functional view of HDL particles. In this regard, it was shown that one key function of HDL particles, namely cholesterol efflux capacity from macrophages, has a strong inverse association with atherosclerosis that is independent of HDL-C levels (Khera et al., 2011). Simultaneously, macrophage-to-feces RCT became a recognized key function of HDL important for regression of atherosclerosis (Cuchel and Rader, 2006). Rader and others identified a multitude of novel approaches to promote RCT, including transgenic overexpression of SR-BI or apo-AI, apo-AI-directed therapeutics, and Lxr and Pparα activation (Rosenson et al., 2012). Moreover, biliary sterol secretion was recognized to be required for functional macrophage-to-feces RCT (Nijstad et al., 2011). Here, we show that...
aspirin, one of the most widely used drugs for prevention of atherosclerosis, promotes macrophage-to-feces RCT by increasing fecal excretion of bile acids. Moreover, we found that aspirin induces regression of established atherosclerosis in mice, which may—besides its well-known anti-inflammatory and platelet-inhibitory effects—be at least in part due to promotion of RCT.

We identify Abcb11 to be upregulated upon aspirin treatment. Analysis of bile acid kinetics, together with macrophage-to-feces RCT measurement in Abcb11−/− mice, showed that promotion of bile flow is the driving force of RCT in mice treated with aspirin. Failure of aspirin to increase RCT in Abcb11−/− mice is conceivably due to the lack of Abcb11, which, however, cannot be definitively ruled out, because there may be other hypothetic compensatory mechanisms. Intriguingly, by the help of lipid mediator profiling we found that both Alox5-dependent classes of lipids, namely LTs and LXs, induce protein expression of Abcb11 in a posttranslational fashion, presumably by
stabilizing MAPK p38-dependent trafficking of Abcb11 from the Golgi to the cytosol and plasma membrane (Kubitz et al., 2004). Depletion of Kupffer cells by injection of mice with clodronate liposomes abrogated the effect of aspirin on hepatic Abcb11 expression, indicating that these cells serve as a major source for LT and LX formation in vivo (Figure S6). So far, Alox5 was thought to be critical for LT and LX generation. However, our lipidomic analyses in primary murine hepatocytes and in livers of Alox5<sup>−/−</sup> mice indicate that in the absence of Alox5 other enzymes may compensate the lack of this lipoxigenase. Interestingly, this alternative pathway remains aspirin responsive, since treatment of Alox5<sup>−/−</sup> mice with aspirin led to a further decrease in LT levels (Figure S7), thus indicating the existence of a regulated compensatory mechanism of LT and LX synthesis in liver of mice.

In an effort to better understand the relative contributions of the main lipoxigenases involved in LT and LX synthesis to RCT in mice, we performed a series of loss-of-function experiments. Studies in mice lacking Alox5 or Alox12/15 identified Alox5 as the lipoxigenase with RCT-modifying properties. By combining our mechanistic in vitro experiments with mediator lipidomic profiling, our data suggest that in vivo the balance between LXS and LTs modulates the expression of hepatic Abcb11, i.e., when LX levels exceed LT levels. Surprisingly, Alox5<sup>−/−</sup> mice did not show increased excretion of macrophage-derived sterols, despite reduced plasma [3H]-cholesterol levels. This may be due to lower biliary excretion of neutral sterols via Abcg8 and/or increased intestinal absorption of neutral sterols in Alox5<sup>−/−</sup> mice, a prediction which will require further studies to be clarified.

Finally, we tested LX mimetics in vivo for the following reasons: first, LX mimetics are stable in circulation and may thus modulate hepatic expression of sterol receptors/transporters in a favorable way; second, LXS have a proresolving function in atherosclerosis. Taken together, these properties may confer atheroprotective effects on LX mimetics. We found that stable LXB4 analogs strongly induced hepatic expression of LDLR and accordingly reduced plasma cholesterol levels. To better understand how lipoxin mimetics regulate LDLR expression in liver of mice, further studies aimed at investigating the regulation of known LDLR-modifying machineries including HMG-CoAr-, PCSK9-, and IDOL-dependent pathways, together with the analysis of cholesterol-modulating miRNAs in livers of LX-treated mice, are needed. Intriguingly, we found that Abcb11 protein levels were reduced in livers of mice treated with LX mimetics. From our lipidomic analyses we learned that in vivo Abcb11 can be induced when (1) both LTs and LXS are increased, and when (2) LXS remain unchanged and LTs are reduced, whereas Abcb11 expression is unaltered when LXS remain unchanged and LTs increase. These data indicate that in the liver LX generation is tightly regulated and that LXS are probably a more important regulator of Abcb11 than LTs. In fact, our in vitro experiments showed that LXS induce Abcb11 only at low concentrations, whereas they decrease its expression when given at higher dosages. We thus conclude that LXS may regulate Abcb11 with higher specificity than LTs and speculate that exogenously administered LXS could have critically altered lipid mediator homeostasis in the liver, thereby decreasing Abcb11 expression.

Our results support the notion that modulation of the AA metabolome may be used to treat and prevent CAD, and may explain how omega-6 PUFAs influence cholesterol homeostasis. In humans, most PUFAs in the diet consist of the omega-6 type including linoleic acid which is converted into the metabolically important AA after consumption (Harris et al., 2009; Katan, 2009). Higher omega-6 PUFA levels were shown to improve insulin resistance, to reduce the incidence of diabetes mellitus, and to associate with lower blood pressure. Moreover, omega-6 PUFAs were shown to lower plasma LDL-C and plasma total cholesterol-to-HDL-C ratio (summarized in Harris et al., 2009). These effects are believed to confer omega-6 PUFAs and especially AA with atheroprotective properties (Harris et al., 2009; Katan, 2009). Combined data from randomized trials, case-control and cohort studies, and animal experiments indicate that the consumption of at least 5%–10% of energy from omega-6 PUFAs reduces the cardiovascular risk (Harris et al., 2009), which recently prompted the American Heart Association to release a recommendation for dietary supplementation with.
omega-6 PUFAAs (Harris et al., 2009). Using a systematic, interdisciplinary approach, we were able to elucidate the relative role of key players of the AA metabolism in whole-body cholesterol homeostasis in humans and in mice.

To summarize, we show that pharmacological and genetic modulation of the AA metabolome affects one major function of HDL, namely RCT. Moreover, we show that the AA metabolome is a conserved regulator of HDL-C in humans and in mice, and identify LX mimetics (Maddox et al., 1998) as a new biochemical modulator of HDL, namely RCT. Moreover, we show that the AA metabolome plays a physiological role in whole-body cholesterol homeostasis and HDL function in mammals, paving the way for the development of novel lipid-lowering drugs based on the structure of AA metabolites and offering a novel therapeutic strategy to counteract CAD in humans.

**EXPERIMENTAL PROCEDURES**

**Human Genome-Wide Association Data**

Genome-wide association data with HDL-C, LDL-C, and total cholesterol was generated by the Global Lipids Genetics Consortium (GLGC) as previously described (Teslovich et al., 2010); http://www.sph.umich.edu/csg/abecasis/public/lipids2010/. The GLGC 2013 data set was recently published (Willer et al., 2013); http://www.sph.umich.edu/csg/abecasis/public/lipids2013/. Plots used in Figures 1, S1, and S2 were generated using LocusZoom (Pruim et al., 2010).

**Animal Studies**

All animals were handled in strict accordance with good animal practice as defined by the Austrian Authorities, and all work was approved by the Austrian Animal Care and Use Committee (Bundesministerium für Wissenschaft und Forschung—BMWF). Mice were fed a standard chow diet (Sniff). To induce Cox I/II inhibition, mice were treated with drinking water containing aspirin for 7 days, as described previously by our laboratory (Tancevski et al., 2006). On a body-scale-adjusted scale, the amount of aspirin would be equal to ~360 mg per day if the animals weighed 60 kg (6 mg kg \(^{-1}\) per day) (Tancevski et al., 2008).

**Mediator Lipidomics**

LC-MS/MS-based lipidomic analyses were performed using a high-performance liquid chromatography (HPLC) system (Waters UPLC) with a linear ion trap quadrupole mass spectrometer (QTRAP5500; AB SCIEX) equipped with an Acquity UPLC BEH C\(_{18}\) column (Waters) as described (Arta, 2012; Morita et al., 2013). MS/MS analyses were conducted in negative ion mode, and fatty acid metabolites were identified and quantified by multiple reaction monitoring (MRM). Different tissue isolation procedures and/or different age or body weight of mice could cause differences in the basal levels of AA metabolites. Thus, lipid mediator levels shown in Figures 3B, 5F, 6F, and S7 cannot be directly compared.

**Statistical Analysis**

Statistical analysis was carried out with a SPSS statistical package (IBM). We determined significance by unpaired two-tailed Student’s t test, or by one-way ANOVA when more than two groups were compared. \(p < 0.05\) was considered statistically significant.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Supplemental Experimental Procedures, two tables, and seven figures and can be found with this article at http://dx.doi.org/10.1016/j.cmet.2014.09.004.

**AUTHOR CONTRIBUTIONS**

I. Tancevski conceived the study. E.D., A.S., K.A., C.H., J.R.P., P.E., M.T., I. Theurl, M. Theurl, M.S., D.L., U.S., D.H., M.A., S.D., M.N., E.H., M.S., A.R.M., X.L., P.P., H.S., T.S., W.M., M.E.K., K.G., P.U., A.L.C., F.S., M.R., K.K., Y.I., M.A., J.D.S., P.P.P., U.J.F.T., M. Trauner., G.D.N., T.C., A.A.H., G.W., and I. Tancevski performed the experiments and analyzed and interpreted the data. I. Tancevski, P.E., U.J.F.T, G.D.N., A.A.H., T.C., M. Trauner, and G.W. wrote the paper.

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**REFERENCES**

Arta, M. (2012). Mediator lipidomics in acute inflammation and resolution. J. Biochem. 152, 313–319.

Birn, C., Dahlén, S.E., Drazen, J., Evans, J.F., Hay, D.W., Nicosia, S., Serhan, C.N., Shimizu, T., and Yokomizo, T. (2003). International Union of Pharmacology XXXVII. Nomenclature for leukotriene and lipoxin receptors. Pharmacol. Rev. 55, 195–227.

Cuchel, M., and Rader, D.J. (2006). Macrophage reverse cholesterol transport: key to the regression of atherosclerosis? Circulation 113, 2548–2555.

Degoma, E.M., and Rader, D.J. (2011). Novel HDL-directed pharmacotherapeutic strategies. Nat. Rev. Cardiol. 8, 266–277.

Drechsler, M., Megens, R.T., van Zandvoort, M., Weber, C., and Soehnlein, O. (2010). Hyperlipidemia-triggered neutrophilia promotes early atherosclerosis. Circulation 122, 1837–1845.

Dutta, P., Courties, G., Wei, Y., Leuschner, F., Gorbatov, R., Robbins, C.S., Cuchel, M., and Rader, D.J. (2006). Macrophage reverse cholesterol transport: key to the regression of atherosclerosis? Circulation 113, 2548–2555.

Figs, A., Lammert, F., Paigen, B., Henkel, A., Matern, S., Korstanje, R., Strojner, B.L., Chen, F., Stoltenberg, E., Spatz, K., et al. (2004). Hepatic overexpression of murine Abcb11 increases hepatobiliary lipid secretion and reduces hepatic steatosis. J. Biol. Chem. 279, 2790–2799.

Gálman, C., Arvidsson, I., Angelin, B., and Rudling, M. (2003). Monitoring hepatic cholesterol 7alpha-hydroxylation activity by assay of the stable bile acid intermediate 7alpha-hydroxy-4-cholesten-3-one in peripheral blood. J. Lipid Res. 44, 859–866.

Harris, W.S., Mozaffarian, D., Rimm, E., Kris-Etherton, P., Rudel, L.L., Appel, L.J., Engler, M.M., Engler, M.B., and Sacks, F. (2009). Omega-6 fatty acids and risk for cardiovascular disease: a science advisory from the American Heart Association Nutrition Subcommittee of the Council on Nutrition, Physical Activity, and Metabolism; Council on Cardiovascular Nursing; and Council on Epidemiology and Prevention. Circulation 119, 902–907.

Helgadottir, A., Manolescu, A., Thorleifsson, G., Gretarsdottir, S., Jonsdottir, H., Thorsteinsdottir, U., Samani, N.J., Gudmundsson, G., Grant, S.F., Thorgerisson, G., et al. (2004). The gene encoding 5-lipoxygenase activating protein...
protein confers risk of myocardial infarction and stroke. Nat. Genet. 36, 233–239. 
Ivanov, I., Heydeck, D., Hofheinz, K., Roffels, J., O’Donnell, V.B., Kuhn, H., and Walther, M. (2010). Molecular enzymology of lipooxygenases. Arch. Biochem. Biophys. 503, 161–174. 
Katán, M.B. (2009). Omega-6 polyunsaturated fatty acids and coronary heart disease. Am. J. Clin. Nutr. 89, 1283–1284. 
Khera, A.V., Cuchel, M., de la Llera-Moya, M., Rodrigues, A., Burke, M.F., Jafri, K., French, B.C., Phillips, J.A., Muckavage, M.W., Wilensky, R.L., et al. (2011). Cholesterol efflux capacity, high-density lipoprotein function, and atherosclerosis. N. Engl. J. Med. 364, 127–135. 
Kubitz, R., Südfels, G., Kühlkamp, T., Kölling, R., and Häussinger, D. (2004). Trafficking of the bile salt export pump from the Golgi to the canicular membrane is regulated by the p38 MAP kinase. Gastroenterology 126, 541–553. 
Kühn, H., and O’Donnell, V.B. (2006). Inflammation and immune regulation by 12/15-lipoxygenases. Prog. Lipid Res. 45, 334–356. 
Lefebvre, P., Cariou, B., Lien, F., Kuipers, F., and Staels, B. (2009). Role of bile acids and bile acid receptors in metabolic regulation. Physiol. Rev. 89, 147–191. 
Maddox, J.F., Colgan, S.P., Clish, C.B., Petasis, N.A., Fokin, V.V., and Serhan, C.N. (1998). Lipoxin B4 regulates human monocyte/neutrophil adherence and motility: design of stable lipoxin B4 analogs with increased biologic activity. FASEB J. 12, 487–494. 
McMahon, B., and Godson, C. (2004). Lipoxins: endogenous regulators of inflammation. Am. J. Physiol. Renal Physiol. 286, F189–F201. 
Mehrabian, M., Allayee, H., Stockton, J., Lum, P.Y., Drake, T.A., Castellani, L.W., Suh, M., Armour, C., Edwards, S., et al. (2005). Integrating genotypic and expression data in a segregating mouse population to identify inflammation. Am. J. Physiol. 289, 127–135. 
Morata, M., Kuba, K., Ichikawa, A., Nakayama, M., Katabira, J., Iwamoto, R., Watanabe, T., Sakabe, S., Daidoji, T., Nakamura, S., et al. (2013). The lipid mediator protectin D1 inhibits influenza virus replication and improves severe influenza. Cell 153, 112–125. 
Nijstad, N., Gao, Z., Dietrich, J., and Tietje, U.J. (2011). Biliary sterol secretion is required for functional in vivo reverse cholesterol transport in mice. Gastroenterology 140, 1043–1051. 
Norata, G.D., Garlaschelli, K., Grigore, L., Raselli, S., Tramontana, S., Manaresi, A., Rizzitelli, G., Glick, J.M., Rothblat, G.H., and Rader, D.J. (2010). The liver-selective thyromimetic T0681 influences reverse cholesterol transport and atherosclerosis development in mice. PLoS ONE 5, e8722. 
Rosenson, R.S., Brewer, H.B., Jr., Davidson, W.S., Fayad, Z.A., Fuster, V., Goldstein, J., Hailerstein, M., Jiang, X.C., Phillips, M.G., Rader, D.J., et al. (2012). Cholesterol efflux and atheroprotection: advancing the concept of reverse cholesterol transport. Circulation 125, 1905–1919. 
Serhan, C.N. (2007). Resolution phase of inflammation: novel endogenous anti-inflammatory and proresolving lipid mediators and pathways. Annu. Rev. Immunol. 25, 101–137. 
Spité, M., and Serhan, C.N. (2010). Novel lipid mediators promote resolution of acute inflammation: impact of aspirin and statins. Circ. Res. 107, 1170–1184. 
Stieger, B. (2009). Recent insights into the function and regulation of the bile salt export pump (ABC11). Curr. Opin. Lipidol. 20, 176–181. 
Stieger, B., and Beuers, U. (2011). The canalicular bile salt export pump BSEP (ABC11) as a potential therapeutic target. Curr. Drug Targets 12, 661–670. 
Tanevski, I., Wehinger, A., Schoeber, W., Eller, P., Cuzzocrea, S., Foeger, B., Patsch, J.R., and Ritsch, A. (2006). Aspirin regulates expression and function of scavenger receptor-Bl in macrophages: studies in primary human macrophages and in mice. FASEB J. 20, 1328–1335. 
Tanevski, I., Demetz, E., Eller, P., Duvenecke, K., Hoefer, J., Heim, C., Stanzl, U., Wehinger, A., Auer, K., Koller, R., et al. (2010). The liver-selective thromboxenic T-M0811 influences reverse cholesterol transport and atherosclerosis development in mice. PLoS ONE 5, e8722.