Aspirin reduces hypertriglyceridemia by lowering VLDL-triglyceride production in mice fed a high-fat diet

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Systemic inflammation is strongly involved in the pathophysiology of the metabolic syndrome, a cluster of metabolic risk factors that includes hypertriglyceridemia. Aspirin treatment lowers inflammation via inhibition of NF-κB activity but also reduces hypertriglyceridemia in humans. The aim of this study was to investigate the mechanism by which aspirin improves hypertriglyceridemia. Human apolipoprotein CI (apoCI)-expressing mice (APOCI mice), an animal model with elevated plasma triglyceride (TG) levels, as well as normolipidemic wild-type (WT) mice were fed a high-fat diet (HFD) and treated with aspirin. Aspirin treatment reduced hepatic NF-κB activity in HFD-fed APOCI and WT mice, and in addition, aspirin decreased plasma TG levels (−32%, P < 0.05) in hypertriglyceridemic APOCI mice. This TG-lowering effect could not be explained by enhanced VLDL-TG clearance, but aspirin selectively reduced hepatic production of VLDL-TG in both APOCI (−28%, P < 0.05) and WT mice (−33%, P < 0.05) without affecting VLDL-apoB production. Aspirin did not alter hepatic expression of genes involved in FA oxidation, lipogenesis, and VLDL production but decreased the incorporation of plasma-derived FA by the liver into VLDL-TG (−24%, P < 0.05), which was independent of hepatic expression of genes involved in FA uptake and transport. We conclude that aspirin improves hypertriglyceridemia by decreasing VLDL-TG production without affecting VLDL particle production. Therefore, the inhibition of inflammatory pathways by aspirin could be an interesting target for the treatment of hypertriglyceridemia.

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conditions with a 12:12-h light-dark cycle. At the age of 10–12 wk, mice received a HFD (45 energy% derived from palm oil, D12451; Research Diet Services, Wijk bij Duurstede, The Netherlands) for a period of 6 wk. Aspirin treatment (120 mg·kg⁻¹·day⁻¹ in drinking water, pH 6.4) was given during the last 4 wk on HFD, and mice were subsequently used for experiments after an overnight fast at 9 AM. Control mice received the same drinking water of pH 6.4 without the addition of aspirin. Mice were allowed free access to food and water. Animal experiments were approved by the institutional ethics committee on animal care and experimentation at Leiden University Medical Center.

Liver NF-κB activation. Since the most common form of NF-κB is the p50/p65 heterodimer (20), the activity of both the p50 and p65 subunits in liver tissue was determined using electrophoretic mobility shift assay (22). Shortly, tissues were homogenized in ice-cold passive lysis buffer (Promega, Madison, WI) and centrifuged (14,000 rpm, 20 min; 4°C). Protein content of the supernatant was determined using the BCA protein assay kit (Pierce, Rockford, IL). For the electrophoretic mobility shift assay, the gel shift assay system was purchased from Promega. The probe was end-labeled using T4 polynucleotide kinase and [³²P]ATP and purified on a Microspin G-25 column (GE Healthcare, Piscataway, NJ). For each sample, 50 μg of protein was incubated with labeled probe and binding buffer (Promega) for 20 min at room temperature. Specific competition was done by adding unlabeled NF-κB binding probe to the reaction. The mixtures were run on 4.5% polyacrylamide gel electrophoresis in 0.5 × Tris-Borate-EDTA buffer. The gel was vacuum-dried and exposed to radiographic film.

Plasma parameters. Blood was collected from the tail vein into chilled paraxoan-coated capillaries (Sigma, St. Louis, MO) to prevent ongoing lipolysis (37). Capillaries were placed on ice and centrifuged, and plasma was assayed for TG, total cholesterol (TC), and phospholipids (PL) using commercially available enzymatic kits from Roche Molecular Biochemicals (Indianapolis, IN). Free fatty acids (FFA) were measured using NEFA C kit from Wako Diagnostics (Instruchemie, Delfzijl, The Netherlands). β-Hydroxybutyrate (β-HB) was determined using the enzymatic β-HB Assay kit from BioVision (Mountain View, CA).

Liver lipids. Lipids were extracted from livers according to a modified protocol from Bligh and Dyer (3). Shortly, a small piece of liver was homogenized in ice-cold methanol. After centrifugation, lipids were extracted by addition of 1,800 μl of CH₃OH-CHCl₃ (3:1 vol/vol) to 45 μl of homogenate. The CHCl₃ phase was dried and dissolved in 2% Triton X-100. Hepatic TG and TC concentrations were measured using commercial kits, as described in Methods. Lipid classes were determined using the enzymatic methods for lipase, cholesterol esterase, and cholesterol oxidase from BioVision (Mountain View, CA). The quality of each mRNA sample was examined by lab-on-a-chip technology using Experion Sdsens analysis kit (Bio-Rad, Hercules, CA). One microgram of total RNA was reverse-transcribed with iScript cDNA synthesis kit (Bio-Rad), and obtained cDNA was purified with Nucleospin Extract II kit (Macherey-Nagel). Real-Time PCR was carried out on the IQ5 PCR machine (Bio-Rad) using the Sensimix SYBR Green RT-PCR mix (Quantace, London, UK). mRNA levels were normalized to mRNA levels of cyclophilin (Cyclo) and glyceraldehyde-3-phosphate dehydrogenase (Gapdh). Primer sequences are listed in Table 1.

Contribution of plasma FA to VLDL-TG production. To measure the contribution of plasma-derived FA to the VLDL-TG production in vivo, mice were fasted overnight as described above. Mice received a continuous iv infusion of [³¹H]-labeled FA [9,10(n)-³¹H]palmitic acid in PBS with 2% bovine serum albumin] at a rate of 100 μl/h (1.6 μCi/h). After 2 h of [³¹H]-labeled FA infusion, a blood sample was taken (t = 0 min), and Triton WR-1339 (Sigma-Aldrich) was injected iv (0.5 mg/g body wt, 10% solution in PBS) to block serum VLDL clearance. Additional blood samples were drawn 15, 30, 60, and 90 min after injection and used for determination of [³¹H] activity in the TG fraction. Lipids were extracted by adding 10 μl of plasma to 3.25 ml of extraction fluid (heptane-methanol-chloroform, 100:128:137 vol/vol/vol). [³¹H]TGs were subsequently separated from [³¹H]FA; 1 ml of potassium carbonate (0.1 M K₂CO₃, pH 10.5) was added, followed by vortexing and centrifugation (3,600 rpm, 15 min), leading to an upper alkaline-methanol-aqueous phase containing saponified [³¹H]FA and a lower chloroform-organic phase containing [³¹H]TG (1). A fraction (0.5 ml) of the total aqueous phase (2.45 ml) was counted for [³¹H] activity in scintillation fluid. The amount of [³¹H]TG in each sample was calculated by subtracting total [³¹H]FA activity from total [³¹H] activity.

Statistical analysis. Data are presented as means ± SD. Statistical differences were calculated using the Mann-Whitney test for two independent samples with SPSS 16.0 (SPSS, Chicago, IL). P < 0.05 was regarded as statistically significant.

RESULTS

Aspirin reduces hepatic NF-κB activation. To verify that aspirin inhibits hepatic NF-κB activity, the activities of the NF-κB subunits p50 and p65 were measured in livers of APOC1 and WT mice fed a HFD and treated with or without...
Aspirin lowers plasma TG and cholesterol levels in HFD-fed APOC1 mice. To examine whether aspirin could reduce hypertriglyceridemia in APOC1 mice, hyperlipidemic APOC1 mice were fed a HFD for 6 wk and treated with or without aspirin, and plasma lipids were determined (Fig. 2). Treatment of mice with aspirin reduced plasma TG levels by 32% (3.94 ± 0.15 to 2.67 ± 0.59 mmol/L, P < 0.05; Fig. 2A) and plasma TC levels by 33% (4.09 ± 0.52 to 2.76 ± 0.90 mmol/L, P < 0.05; Fig. 2B). Aspirin treatment did not affect plasma PL (Fig. 2C) or FFA levels (Fig. 2D). The reduction in plasma TG and TC levels was not caused by a reduction in body weight, since aspirin did not affect body weight in APOC1 mice (control: 30.5 ± 2.1 g; aspirin: 28.9 ± 3.0 g). In WT mice fed a HFD for 6 wk, aspirin did not affect plasma TG, TC, PL, or FFA levels (Fig. 2, E–H). In addition, aspirin did not affect body weight in WT mice (control: 30.3 ± 2.1 g; aspirin: 30.8 ± 1.9 g).

Aspirin attenuates VLDL-like emulsion particle-TG clearance in HFD-fed APOC1 but not WT mice. A decrease in VLDL-TG clearance can be explained by an increase in VLDL-TG production and/or a decrease in hepatic VLDL-TG production. To determine whether aspirin enhances the clearance of VLDL-TG, the plasma clearance and organ distribution of [3H]TO-labeled, TG-rich, VLDL-like emulsion particles was evaluated in aspirin- and control-treated hypertriglyceridemic APOC1 mice (Fig. 3). Unexpectedly, aspirin inhibited rather than enhanced serum clearance of [3H]TO (t1/2 = 15.9 ± 6.6 vs. 5.6 ± 2.6 min; Fig. 3A) in APOC1 mice. This reduction in [3H]TO clearance upon aspirin was reflected by reduced uptake of [3H]TO-derived radioactivity by the liver by ~60% (123 ± 1 vs. 308 ± 75 nmol/g, P < 0.05), by skeletal muscle by ~66% (11 ± 2 vs. 31 ± 15 nmol/g, P < 0.05), and by white adipose tissue (WAT), which reached statistical significance for gonadal WAT (12 ± 3 vs. 44 ± 22 nmol/g, P < 0.05) (Fig. 3B). Apparently, aspirin reduces rather than enhances TG clearance in APOC1 mice and therefore cannot explain the aspirin-induced reduction in VLDL-TG. In WT mice fed a HFD for 6 wk, aspirin did not affect plasma clearance of [3H]TO (Fig. 3C) or organ-specific uptake of [3H]TO-derived radioactivity (Fig. 3D) in WT mice. Apparently, the decreasing effect of aspirin on TG clearance may be specific for APOC1 mice.

Aspirin lowers VLDL-TG production in HFD-fed APOC1 and WT mice. Because the decrease in plasma TG levels in APOC1 mice upon aspirin treatment was not caused by increased TG clearance, we investigated whether the decreased

Table 1. Primers used for quantitative real-time PCR analysis

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<th>Reverse Primer</th>
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<td>ATGGTTTGAGTTGAGTACAG</td>
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Acox1, acyl-coenzyme A oxidase 1, palmitoyl; Apob, apolipoprotein B; Cd36, fatty acid translocase; Cptla, carnitine palmitoyltransferase 1a; Dgat1, diglyceride acyltransferase 1; Fabp1, fatty acid-binding protein 1; Fasn, fatty acid synthase; Mtpg, microsomal triglyceride transfer protein; Ppara, peroxisome proliferator-activated receptor-α; Slc27a2, fatty acid transport protein 2; Slc27a4, fatty acid transport protein 4; Slc27a5, fatty acid transport protein 5; Srebp1, sterol regulatory element-binding protein.

Aspirin reduces hepatic NF-κB activation. Apolipoprotein (apo)CI (APOCI) and wild-type (WT) mice were fed a high-fat diet (HFD) for 6 wk and treated without or with aspirin. Mice were euthanized after an overnight fast, and hepatic NF-κB activity was measured by electrophoretic mobility shift assay in liver tissue of APOCI (A) and WT mice (B) treated without (open bars) or with aspirin (closed bars). Activities of subunits p65 and p65 were measured. Values are means ± SD (n = 3–4). *P < 0.05.
TG levels could be explained by diminished hepatic VLDL-TG production in *APOC1* mice. The rate of hepatic VLDL-TG production was measured by determining plasma TG levels after intravenous Triton WR-1339 injection (Fig. 4). We found a reduction in hepatic VLDL-TG secretion rate in *APOC1* mice treated with aspirin by $-28\%$ (3.42 ± 0.53 vs. 4.95 ± 1.11 mM/h, $P < 0.05$; Fig. 4A), whereas aspirin did not affect the rate of VLDL-apoB production (Fig. 4B). Interestingly, similar to our observation in *APOC1* mice, aspirin did reduce the hepatic VLDL-TG secretion rate in HFD-fed WT mice by $-33\%$ (2.79 ± 0.47 vs. 4.19 ± 0.48 mM/h, $P < 0.05$; Fig. 4C), whereas VLDL-apoB production rate was also not affected (Fig. 4D). Apparently, aspirin generally reduces the VLDL-TG production in HFD-fed mice independent of the genotype. Furthermore, since each VLDL particle contains a single apoB molecule, this observation shows that aspirin treatment inhibits
hepatic VLDL-TG production without affecting the rate of VLDL particle production.

**Aspirin does not affect liver lipid levels in HFD-fed APOC1 and WT mice.** To determine whether the attenuation in hepatic VLDL-TG production was the result of decreased lipid substrate availability in the liver, the effect of aspirin on hepatic lipid content was measured (Fig. 5). However, aspirin did not affect liver TG (Fig. 5A) or TC levels (Fig. 5B) in APOC1 mice. Also, aspirin did not affect liver TG (Fig. 5C) or TC (Fig. 5D) levels in WT mice.

**Aspirin treatment does not affect hepatic expression of genes involved in FA oxidation, lipogenesis, or VLDL production.** Because changes in hepatic gene expression could underlie the reduction in VLDL-TG production, we determined the effect of aspirin on expression of genes involved in FA oxidation, lipogenesis, and VLDL production (Table 2). In both APOC1 and WT mice, aspirin did not affect expression of peroxisome proliferator-activated receptor-α (Ppara), a transcription factor that regulates genes involved in FA oxidation and ketogenesis, nor did it affect its target genes acyl-coenzyme A oxidase 1 (Acox1) or carnitine palmitoyltransferase 1a (Cpt1a). In line with these results, aspirin did not increase plasma B-HB levels in WT mice (data not shown), which is a plasma marker for hepatic FA oxidation and ketogenesis. This implies that the reduced VLDL-TG production upon aspirin treatment is not caused by increased hepatic FA oxidation. We additionally determined the effect of aspirin on expression of genes involved in lipogenesis. In both APOC1 and WT mice, aspirin did not affect expression of sterol regulatory element-binding protein-1c (Srebp-1c), which regulates genes required for de novo lipogenesis, nor did it affect acyl/diacylglycerol transferase 1 (Dgat1), which catalyzes the final and only committed step in TG synthesis, or FA synthase (Fasn), which plays a key role in FA synthesis. These data suggest that aspirin does not affect genetic regulation of de novo lipogenesis. In addition, although aspirin induced an increase in VLDL-TG secretion, it did not affect hepatic gene expression of microsomal TG transfer protein (Mttp), which is involved in the assembly and secretion of VLDL. Furthermore, aspirin does not affect hepatic gene expression of apolipoprotein B (Apob) in APOC1 mice, which is in line with the observation that aspirin does not affect VLDL-apoB secretion in vivo. However, despite the fact that aspirin did not affect VLDL-apoB secretion in WT mice, gene expression of Apob was increased in WT mice.

**Aspirin treatment decreases the contribution of plasma-derived FA to the VLDL-TG production.** Because the decrease in VLDL-TG production was not caused by a reduced hepatic lipid content or decreased expression of genes involved in de novo lipogenesis that could reduce lipid availability for VLDL-TG secretion, we investigated whether the decreased VLDL-TG production could be explained by a diminished contribution of plasma-derived FA for VLDL-TG secretion in WT mice (Fig. 6). The contribution of plasma-derived FA was measured by determining plasma [3H]TG levels after continuous [3H]FA infusion and intravenous Triton WR-1339 injection. We found that aspirin reduced the hepatic [3H]TG secretion rate in WT mice by −24% (3.1 ± 0.4 vs. 2.4 ± 0.7 × 10^3 disintegrations·min⁻¹·h⁻¹, P < 0.05), which suggests that aspirin reduces VLDL-TG production...
by reducing the incorporation of plasma-derived FA into VLDL-TG. This reduction is not caused by a reduced hepatic expression of genes involved in hepatic FA uptake and transport (Table 2), because aspirin did not affect liver-type FA-binding protein (Fabp1) or FA transport proteins 2, 4, and 5 (Slc27a2, Slc27a4, and Slc27a5, respectively) and even increased expression of FA translocase (Cd36) in APOC1 but not WT mice. These data imply that aspirin reduced the VLDL-TG production independent of changes in hepatic expression of genes involved in FA uptake and transport.

Fig. 4. Aspirin decreases VLDL-TG production in HFD-fed APOC1 and WT mice. APOC1 and WT mice were fed a HFD and treated without or with aspirin. Overnight-fasted mice were injected with Trans35S and Triton WR-1339, and blood samples were drawn at the indicated time points. TG concentrations were determined in APOC1 (A) and WT mice (C) treated without (○) or with aspirin (●) and plotted as the increase in plasma TG relative to t = 0 (A). After 120 min, VLDL was isolated by ultracentrifugation, 35S activity was counted, and the production rate of newly synthesized VLDL-35S-apoB was determined for APOC1 (B) and WT (D). Values are means ± SD (n = 5). *P < 0.05.

Fig. 5. Aspirin does not affect liver lipids in HFD-fed APOC1 and WT mice. Livers were collected from overnight-fasted HFD-fed APOC1 and WT mice treated without or with aspirin. Lipids were extracted, and TG (A and C) and TC (B and D) concentrations were measured and expressed per milligram of protein. Values are means ± SD (n = 6).
elucidate the mechanism by which aspirin reduces the LPL activity in humans (30). It would be interesting to addition to inhibition of inflammation via NF-kB inhibition of inflammation by aspirin is expected to increase (7, 8). If indeed inflammation inhibits clearance of TG, increased TG clearance. Earlier studies report that high-dose of VLDL-TG in HFD-fed mice. Therefore, the decrease in plasma TG levels by aspirin cannot be explained by increased TG clearance. Earlier studies report that high-dose LPS injections reduce the clearance of TG-rich lipoproteins by inhibition of the LPL activity, which is mediated by cytokines (7, 8). If indeed inflammation inhibits clearance of TG, inhibition of inflammation by aspirin is expected to increase TG-rich lipoprotein clearance, which is in contrast to our observation in APOC1 mice. It should be noted that aspirin, in addition to inhibition of inflammation via NF-kB, also inhibits prostaglandin synthesis, which has been demonstrated to restore the LPS-induced inhibition of LPL (5). Moreover, an early report has shown that aspirin treatment inhibits postheparin LPL activity in humans (30). It would be interesting to elucidate the mechanism by which aspirin reduces the VLDL-TG clearance; however, this is beyond the scope of the present study because it does not explain the reduction in hypertriglyceridemia that we observe. Moreover, the observation may be a specific feature of the APOC1 transgenic mouse model, since we did not observe such an effect in WT mice.

Aspirin very effectively reduced hepatic secretion of VLDL-TG in APOC1 mice, explaining the reduction in hypertriglyceridemia upon aspirin treatment. In addition, aspirin equally reduced hepatic secretion of VLDL-TG in WT mice, indicating that the effects of aspirin on the VLDL-TG production do not exclusively occur in hypertriglyceridemic mouse models such as the APOC1 mouse. To our knowledge, we show for the first time that a decrease in inflammation corresponds with a drop in

DISCUSSION

Treatment of obese rodents and patients with type 2 diabetes with high-dose aspirin reduces hypertriglyceridemia (13, 36). However, so far, the mechanistic basis for the relationship between aspirin intake and reduced plasma TG levels has been poorly understood. In the present study we focused on the effects of aspirin on VLDL-TG metabolism in HFD-induced obese hyperlipidemic APOC1 mice and additionally evaluated the effects of aspirin on VLDL-TG metabolism in HFD-fed normolipidemic WT mice. Our results document that aspirin treatment improves hypertriglyceridemia by reducing the hepatic production of VLDL-TG as a result of an attenuated hepatic incorporation of plasma-derived FA into VLDL-TG rather than from increased clearance of VLDL-TG from the circulation.

In the present study, aspirin treatment decreased plasma TG and TC levels in HFD-fed APOC1 mice that display hypertriglyceridemia. This improvement in hyperlipidemia is in accord with earlier studies showing reduced serum TG concentrations upon aspirin or salicylate treatment in patients with type 2 diabetes mellitus (13) and in diabetic rats (36).

Our data show that aspirin treatment attenuated the clearance of VLDL-like TG-rich particles in APOC1 mice. Therefore, the decrease in plasma TG levels by aspirin cannot be explained by increased TG clearance. Earlier studies report that high-dose LPS injections reduce the clearance of TG-rich lipoproteins by inhibition of the LPL activity, which is mediated by cytokines (7, 8). If indeed inflammation inhibits clearance of TG, inhibition of inflammation by aspirin is expected to increase TG-rich lipoprotein clearance, which is in contrast to our observation in APOC1 mice. It should be noted that aspirin, in addition to inhibition of inflammation via NF-kB, also inhibits prostaglandin synthesis, which has been demonstrated to restore the LPS-induced inhibition of LPL (5). Moreover, an early report has shown that aspirin treatment inhibits postheparin LPL activity in humans (30). It would be interesting to elucidate the mechanism by which aspirin reduces the

### Table 2. Aspirin does not generally affect hepatic expression of genes involved in FA uptake and transport, FA oxidation, lipogenesis, or VLDL secretion

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Values are means ± SD (n = 4–5). FA, fatty acid; APOC1, apolipoprotein CI; WT, wild type. Livers were isolated from overnight-fasted APOC1 and WT mice fed a high-fat diet and treated without or with aspirin. mRNA was isolated, and mRNA expression of the indicated genes was quantified by RT-PCR. Genes are grouped as genes involved in FA uptake and transport, FA oxidation, lipogenesis, and VLDL production. Data are calculated as fold difference compared with the control group. *P < 0.05 compared with control group.

Fig. 6. Aspirin reduces the contribution of plasma-derived FA to the VLDL-TG production. WT mice were fed a HFD and treated without or with aspirin. Overnight-fasted mice received a continuous intravenous (iv) infusion of 3H-labeled FA [9,10(n)-3H]palmitic acid for 2 h, followed by an iv injection of Triton WR-1339. Blood samples were drawn at the indicated time points, and 3H activity in the TG fraction was determined in mice treated without (○) or with aspirin (●) and plotted as the increase in plasma [3H]TG relative to t = 0. Values are means ± SD (n = 7). *P < 0.05.
VLDL-TG production. The reduction of VLDL-TG secretion in our study is not paralleled by a reduction in apoB secretion in either APOC1 or WT mice, suggesting that aspirin reduces the lipidation of VLDL particles rather than the number of particles that are secreted by the liver. In contrast to our data on apoB secretion, a recent study by Tsa et al. (32) observed that suppression of IKK with BMS345541 decreased apoB secretion in vitro in primary hamster hepatocytes and HepG2 cells. Although differences between species might explain these conflicting findings, both of these published in vitro studies and our present in vivo study point toward a link between the IKK/NF-κB pathway and the regulation of VLDL production. Moreover, we have recently shown that activation of the hepatic IKK/NF-κB pathway increases VLDL-TG production (33), supporting the hypothesis that the effects of aspirin on the VLDL-TG production are mediated via a reduction in hepatic NF-κB activity. Nevertheless, activation of hepatic IKK/NF-κB increases hepatic Fas expression (33), whereas aspirin in the present study did not change hepatic expression of Fasn, nor did it change expression of other genes involved in TG synthesis, such as Dgat1 and Sreb1c, suggesting that aspirin more likely lowers VLDL-TG production by other mechanisms rather than via its effects on hepatic NF-κB activity. A reduction in hepatic lipid availability by increased lipid oxidation could underlie the mechanism by which aspirin reduces hepatic VLDL-TG production. However, aspirin did not affect expression of genes involved in FA oxidation nor plasma levels of β-HB, a marker of hepatic FA oxidation and ketogenesis. Similarly, aspirin did not affect expression of genes involved in de novo lipogenesis or VLDL production, suggesting that aspirin does not reduce VLDL-TG production by changing expression of genes involved in hepatic lipid metabolism.

It has been suggested that the decrease in plasma TG concentration that occurs upon aspirin treatment might be secondary to the fall in plasma FFA levels (34). A reduction in FFA delivery to the liver could result in a reduced availability of FA for the release of VLDL-TG by the liver (19). Indeed, although aspirin did not change plasma FFA levels, it changed the turnover of FA, as reflected by a ~24% reduction in the incorporation of plasma-derived FA into VLDL-TG, showing that aspirin in fact lowers the availability of plasma-derived FA for VLDL-TG production. This reduction of FA incorporation into VLDL-TG upon aspirin treatment was not caused by a reduced hepatic expression of FA transporter proteins, suggesting that aspirin reduces the FA incorporation via another mechanism. It is possible that aspirin reduces posttranscriptional processing of FA transporters independent of mRNA expression, since expression of FA transporters does not always correlate with changes in protein content or the rate of FA transport (10). Alternatively, aspirin might increase FA uptake and transport via simple diffusion, since FA uptake has been described independent of any FA transporter (10).

The decrease of FA turnover that we observed could be secondary to an increased insulin sensitivity of adipose tissue, thereby decreasing FA mobilization to plasma. Indeed, high-dose salicylates such as aspirin have been shown to increase insulin sensitivity (36), and the reduction in VLDL-TG production in our study is similarly accompanied by an increased insulin sensitivity (van Diepen JA and Voshol PJ, unpublished observations). However, the aspirin-induced reduction in FA utilization and subsequent VLDL-TG secretion in our study were determined under fasting conditions, when the role of insulin is marginal. In fasting conditions, the lipolytic activity of adipocytes is stimulated by catecholamines. Interestingly, aspirin has been reported to reduce catecholamine-stimulated lipolysis, which is therefore a more likely explanation for our findings (29, 31). In addition, it has been shown that aspirin reduces release of FA from adipose tissue directly via inhibition of TNFα-induced lipolysis (38). Therefore, we propose that the fact that aspirin reduces plasma-derived FA utilization by the liver is likely caused via direct inhibition of intracellular lipolysis in adipose tissue, which reduces plasma FA availability. Adipose tissue lipolysis might be further inhibited in the fed state by an increased sensitivity for insulin.

In conclusion, our data show that aspirin inhibits NF-κB and decreases HFD-induced hypertriglyceridemia by reducing hepatic VLDL-TG secretion rather than by accelerating the tissue distribution of VLDL-TG. The reduction in VLDL-TG is not caused by a decreased steatosis, increased FA oxidation, or changes in de novo lipogenesis but by an attenuation of hepatic incorporation of plasma-derived FA into VLDL-TG. In the scope of our findings, aspirin could potentially be a new therapeutic drug in the treatment of hypertriglyceridemia. However, chronic high-dose aspirin is associated with risk for bleeding. On the other hand, salsalate is a nonsteroidal, anti-inflammatory drug with similar structure that is regarded as a safer alternative. High-dose salsalate treatment has been shown recently to reduce TG levels in diabetic patients similarly to high-dose aspirin treatment (11) and could therefore potentially be a new drug for the treatment of hypertriglyceridemia.

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DISCLOSURES
No conflicts of interest, financial or otherwise, are declared by the authors.

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