Characterization of a Novel Intestinal Glycerol-3-phosphate Acyltransferase Pathway and Its Role in Lipid Homeostasis*

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Dietary triglycerides (TG) are absorbed by the enterocytes of the small intestine after luminal hydrolysis into monoaclglycerol and fatty acids. Before secretion on chylomicrons, these lipids are reesterified into TG, primarily through the monoacylglycerol pathway. However, targeted deletion of the primary murine monoaclglycerol acyltransferase does not quantitatively affect lipid absorption, suggesting the existence of alternative pathways. Therefore, we investigated the role of the glycerol 3-phosphate pathway in dietary lipid absorption. The expression of glycerol-3-phosphate acyltransferase (GPAT3) was examined throughout the small intestine. To evaluate the role for GPAT3 in lipid absorption, mice harboring a disrupted Gpat3 gene (Gpat3−/−) were subjected to an oral lipid challenge and fed a Western-type diet to characterize the role in lipid and cholesterol homeostasis. Additional mechanistic studies were performed in primary enterocytes. GPAT3 was abundantly expressed in the apical surface of enterocytes in the small intestine. After an oral lipid bolus, Gpat3−/− mice exhibited attenuated plasma TG excursion and accumulated lipid in the enterocytes. Electron microscopy studies revealed a lack of lipids in the lamina propria and intercellular space in Gpat3−/− mice. Gpat3−/− enterocytes displayed a compensatory increase in the synthesis of phospholipid and cholesteryl ester. When fed a Western-type diet, hepatic TG and cholesteryl ester accumulation was significantly higher in Gpat3−/− mice compared with the wild-type mice accompanied by elevated levels of alanine aminotransferase, a marker of liver injury. Dysregulation of bile acid metabolism was also evident in Gpat3-null mice. These studies identify GPAT3 as a novel enzyme involved in intestinal lipid metabolism.

The ability of an organism to efficiently extract energy-rich nutrients from its diet is critical for survival. Triglycerides (TG) are energy-dense lipid molecules; however, their biological properties preclude direct absorption in the gut. The primary site for the absorption of TG and other dietary lipids is the small intestine. Lipid absorption is a multistep process that begins with the intraluminal emulsification of fat by bile acids, which in turn facilitates hydrolysis by multiple lipases. TG hydrolysis leads to the liberation of monoacylglycerol (MAG) and free fatty acid (FFA). Absorption of these hydrolyzed products occurs in the epithelial monolayer lining the villi and involves both passive diffusion and facilitated transport. After absorption, FFAs are primarily reesterified into TG and either temporarily stored as cytosolic lipid droplets (CLDs) or packaged into chylomicrons and secreted into the lymphatic vessels of the lamina propria through the basolateral surface of enterocytes (1).

There are two primary routes for TG synthesis, namely, the MAG and glycerol 3-phosphate (G3P) pathways. Both pathways produce diacylglycerol (DAG) as a common intermediate. In the MAG pathway DAG is directly synthesized from sn-2 MAG through the action of monoacylglycerol acyltransferases (MGAT). In the G3P pathway DAG is produced by three reactions. Initially, G3P is acylated by the members of the glycerol-3-phosphate acyltransferase (GPAT) family. The resultant lysophosphatic acid is further acylated by 1-acylglycerol-3-phosphate O-acyltransferases and then dephosphorylated by phosphatidic acid phosphatase enzymes to produce DAG. In most cell types producing TG, the G3P pathway is the primary route for TG synthesis (2). In the intestine MAG is the preferred acceptor of FFAs, and the MAG pathway accounts for ~80% of TG synthesis (3–7). Importantly, targeted disruption of the murine Mogat2 gene, which encodes the predominant MGAT in the small intestine, does not quantitatively affect lipid absorption. Rather, loss of MGAT2 activity in the gut results in a shift in the spatiotemporal profile of lipid absorption with the distal small intestine assuming a more prominent role in this process (8). Although the enzymes responsible for TG absorption in Mogat2−/− mice have not been delineated, it is possible that a hitherto unidentified MGAT compensates for the loss of MGAT2 or that the MAG is further hydrolyzed to glycerol and FFA and enters the G3P pathway (9–11).

To date, four mammalian GPAT genes (GPAT1–4) have been identified. GPATs catalyze the first committed step in TG acyltransferase pathway and then dephosphorylated by phosphatidic acid phosphatase enzymes to produce DAG.
GPAT enzymes are broadly categorized based on their sensitivity to inhibition by sulphydryl group-modifying reagents like N-ethylmaleimide (NEM) and subcellular location. Thus, GPAT1 and GPAT2 localize to mitochondria, whereas GPAT3 and GPAT4 are nominally microsomal. GPAT1 is resistant to inhibition by NEM, whereas GPAT2, 3, and 4 are NEM-sensitive (13–17). In mice, the gene encoding GPAT4 is highly expressed in the liver, whereas GPAT3 is predominantly expressed in the small intestine and adipose tissue (18). Gpat3<sup>−/−</sup> mice have reduced NEM-sensitive GPAT activity in white adipose depots and liver and, when maintained on standard chow, do not exhibit any obvious metabolic abnormalities aside from increased hepatic cholesterol and cholesteryl ester levels. When fed a high fat diet, modest metabolic perturbations, including decreased body weight gain and elevated physical activity, were evident in female mice (18). Given the abundant expression of GPAT3 in the small intestine, we investigated a potential role for this enzyme in intestinal lipid absorption. We observed a distinct regional localization for GPAT3 protein to the jejunum, the primary site for lipid absorption. We demonstrate for the first time that GPAT3 plays a role in intestinal lipid absorption and that loss of GPAT3 expression results in abnormalities in dietary lipid absorption and a disruption of enteric and hepatic lipid homeostasis.

**Experimental Procedures**

**Animal Studies**—GPAT3 knock-out (Gpat3<sup>−/−</sup>) mice were generated as described previously (18). Animals were bred at Taconic (Germantown, NY) and transferred to a Pfizer facility at least 14–21 days before the start of experiments. Mice were housed under standard laboratory conditions on a 12-h light, 12-h dark cycle (lights on at 06:00 a.m.) and allowed free access to food and water. All studies were performed according to Pfizer internal animal use protocols and approved by Pfizer's Institutional Animal Care and Use Committee. All studies used age-matched (14–16 weeks) male wild-type and Gpat3<sup>−/−</sup> mice. For studies using fed and fasted cohorts, mice were either allowed standard laboratory rodent chow 5001 ad libitum or fasted for 6 h from 06:00 a.m. to 12:00 p.m. In high fat diet studies, mice were either allowed standard laboratory rodent chow (5001) or fed a Western-type diet (WTD, TD88137, Harlan Teklad, 42% kcal from fat, 21) containing [14C]triolein (0.5 mCi/mmol) and 50 μM oleoyl-CoA. Samples were incubated with the buffer and the substrate for 15 min at room temperature. For NEM-resistant activity, the total cell lysates were preincubated with 0.4 mM NEM for 15 min on ice before the initiation of the GPAT reaction. Products were extracted using chloroform and methanol and separated by thin-layer chromatography (TLC) using chloroform:methanol:water (65:25:4), bands were scraped, and 14C counts were quantitated using scintillation counter. MGAT activity assays in the intestinal mucosal tissue were determined by measuring incorporation of the [14C]oleoyl moiety into diacylglycerol. Reactions were initiated by incubating microsomal proteins with 4 μM [14C]decanoyl-CoA (PerkinElmer Life Sciences) in the presence or absence (control) of 35 μM 2-oleoylgllycerol in 50 mM HEPES (pH 7.4), 10 mM MgCl<sub>2</sub>, 0.001% Triton X-100, and 2.5% v/v acetone. A non-selective lipase inhibitor MAPF (methyl arachidonoyl fluorophosphates) was added to block the hydrolysis of both product and substrate as previously described (24). After 30 min reactions were terminated by adding 50 μl of 1% phosphoric acid. Lipids were extracted using CHCl<sub>3</sub>/methanol (2:1, v/v) and separated by TLC, and DAG bands were quantified.

**Immunohistology, Immunohistochemistry, and Electron Microscopy**—The entire small intestine was removed for histological analysis and cut into ~5.0-cm sections and flushed with physiological saline. From each 5.0-cm segment cross-sections were fixed with 10% neutral-buffered formalin and processed to paraffin blocks. Individual small intestine segments or liver tissues were embedded in paraffin blocks so that areas of could be evaluated on one slide per animal. Five-micron sections were cut using a rotary microtome and mounted on slides for routine

**Western Immunoblotting**—Microsomes were isolated as described elsewhere (22). Briefly, intestinal segments were homogenized in buffer containing 250 mM sucrose, 20 mM Tris-HCl, pH 7.4, 1 mM EDTA. Cellular debris was removed by centrifugation at 5000 x g for 10 min, and crude mitochondrial pellets were obtained by centrifugation of the 5000 x g supernatant at 15,000 x g for 10 min. The supernatant was centrifuged at 106,000 x g for 1 h, and the resultant pellet, corresponding to the microsomal fraction, was resuspended in 20 mM Tris-HCl, 0.5 n NaCl (pH 7.4), lysed, and centrifuged at 106,000 x g for 1 h to obtain the luminal contents. Protease inhibitor mixture (Sigma) was included in all of the solutions used in this isolation procedure. Proteins (50 μg) were separated on 4–20% Novex Tris-glycine gels (Life Technologies), transferred onto 0.45-μm nitrocellulose, and developed for GPAT3 expression using (LPAAT-θ), T-17 (Santa Cruz Biotechnology, Dallas, TX), and secondary antibody, anti-goat IgG, horseradish peroxidase conjugate (Promega, Sunnyvale, CA).

**GPAT and MGAT Activity Assays**—GPAT activity assays in the murine small intestinal segments were performed ostensibly as described elsewhere (17, 18, 23). Intestinal mucosal tissues were scraped and homogenized in ice-cold PBS, and enzyme reactions were initiated by the addition of 6 μg of microsomal protein in 75 mM Tris-HCl (pH 7.5), 4 mM MgCl<sub>2</sub>, 1 mg/ml fatty acid free BSA, 150 μM [14C]glycerol 3-phosphate (55 mCi/mmol) and 50 μM oleoyl-CoA. Samples were incubated with the buffer and the substrate for 15 min at room temperature. For NEM-resistant activity, the total cell lysates were preincubated with 0.4 mM NEM for 15 min on ice before the initiation of the GPAT reaction. Products were extracted using chloroform and methanol and separated by thin-layer chromatography (TLC) using chloroform:methanol:water (65:25:4), bands were scraped, and 14C counts were quantitated using scintillation counter. MGAT activity assays in the intestinal mucosal tissue were determined by measuring incorporation of the [14C]oleoyl moiety into diacylglycerol. Reactions were initiated by incubating microsomal proteins with 4 μM [14C]decanoyl-CoA (PerkinElmer Life Sciences) in the presence or absence (control) of 35 μM 2-oleoylgllycerol in 50 mM HEPES (pH 7.4), 10 mM MgCl<sub>2</sub>, 0.001% Triton X-100, and 2.5% v/v acetone. A non-selective lipase inhibitor MAPF (methyl arachidonoyl fluorophosphates) was added to block the hydrolysis of both product and substrate as previously described (24). After 30 min reactions were terminated by adding 50 μl of 1% phosphoric acid. Lipids were extracted using CHCl<sub>3</sub>/methanol (2:1, v/v) and separated by TLC, and DAG bands were quantified.
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Hematoxylin and eosin (H&E) staining and immunohistochemistry. For Brightfield immunohistochemistry, 5-μm sections were deparaffinized, hydrated, and treated with Biogenex Citra (Biogenex, Fremont, CA) using the Decloaking Chamber (Biocare Medical, Concord, CA) to expose the antigenic sites. Endogenous peroxidase was blocked by incubations with 3.0% solution of hydrogen peroxide (ThermoFisher, Waltham, MA) for 10 min, and nonspecific sites were blocked using Dako serum-free protein block (Dako, Carpinteria, CA) for 20 min. The following antibodies were all incubated for 60 min at room temperature: goat anti-GPAT3 1:2000 (Santa Cruz Biotechnology, Dallas, TX) and guinea pig anti-TIP47 1:2000 (Progen, Heidelberg, Germany). GPAT3 was detected using Biocare goat probe horseradish peroxidase followed by visualization with Dako Liquid diaminobenzidine (DAB+). All slides were counterstained with Mayer’s hematoxylin (Dako) and mounted using permanent mounting medium (Sakura, Torrance, CA). For immunofluorescence immunohistochemistry, 5-μm sections were prepared as described above. Goat anti-GPAT3 and guinea pig anti-TIP47 were used at a dilution of 1:500 for 60 min at room temperature. Detection was accomplished using Alexa fluor secondary antibodies, donkey anti-goat IgG Alexa 488, and donkey anti-guinea pig IgG Alexa 594 secondary antibodies (Life Technologies) for 45 min at room temperature. All slides were mounted with Prolong Gold plus DAPI (Life Technologies). For electron microscopy, ~1 mm of jejunum was fixed in 0.1 M phosphate-buffered fixative containing 4% formaldehyde (methanol-free) and 1% glutaraldehyde for a minimum of 24 h. Specimens were rinsed thoroughly and dehydrated through graded alcohols (methanol-water, 80:20 v/v) and heptadecanoic acid (200 nM) as an intercalating agent. 90 nm thin sections were mounted with Prolong Gold plus DAPI (Life Technologies) and visualized with a Hitachi H-7100 transmission EM, and representative images were digitally captured (advanced microscopy techniques). For secretion studies, primary enterocytes were isolated and incubated with 0.5 μCi/ml [3H]oleic acid or [14C]cholesterol (PerkinElmer Life Sciences) for 1 h at 37 °C with constant shaking, and cell suspensions were gassed with 95% O2 and 5% CO2. Cellular uptake and secretion assays were performed as previously described (26, 27). Briefly, isolated enterocytes were incubated with 0.5 μCi/ml [3H]oleic acid/[14C]cholesterol. Total lipids were extracted to measure total counts and separated on TLC. Lipids were extracted from the remaining cell lysate using modified a Bligh and Dyer method in a 2:1 ratio of methanol:chloroform. Lipid counts were determined on an aliquot of the extract by liquid scintillation, and incorporation of radioisotope into different intracellular lipids was monitored by TLC separation.

For secretion studies, primary enterocytes were isolated and incubated with 0.5 μCi/ml of [3H]oleic acid or [14C]cholesterol for 1 h, and cell pellets were collected and washed as described above after incubation. Cells were resuspended in modified DMEM (27). Media were collected after 2 h, and lipids were extracted to measure total counts and separated on TLC. For cholesterol secretion, media were further subjected to density gradient ultracentrifugation, and counts were measured (26, 27).

High performance Liquid Chromatography (HPLC), Tissue Lipid and Bile Acid Analysis—Liver or intestinal tissues were weighed, and lipids were extracted using a modified Bligh and Dyer method. Before lipid determination, the intestine was thoroughly flushed with saline to ensure no contamination with residual luminal lipids. Phospholipid C, triglycerides M, and total and free cholesterol assays were purchased from Wako (Richmond, VA). HPLC of frozen liver samples to determine...
hepatic lipid levels was performed exactly as described previously (18). For determination of hepatic bile acid levels, liver samples were homogenized in 75% ethanol and incubated at 50 °C in a water bath. Samples were briefly spun, and supernatant containing bile acids was collected. Bile acid concentrations were determined using a commercially available kit (Diazyme, Poway, CA).

Plasma Clinical Chemistry and Incretin Analysis—Plasma analytes were analyzed using clinical analyzer, Hitachi Cobas (Hoffmann-La Roche). Plasma total glucagon-like peptide-1 (GLP-1) and gastric inhibitory polypeptide (GIP) were determined using commercial assays (EMD Millipore, Darmstadt, Germany).

RNA Isolation and Gene Expression—Tissue-specific expression of the Gpat3 gene was determined on RNA obtained from Clonect (Mountain View, CA). For all other gene expression assays, RNA was extracted from aliquots of instantly frozen tissue samples using Qiazol according to the manufacturer’s instructions (Qiagen, Germantown, MD). Gene expression was analyzed using real-time PCR system. All primer-probe assays were from LifeTechnologies with assay numbers as follows: DGAT1 (diacylglycerol O-acyltransferase 1), Mm00515643_m1; DGAT2 (diacylglycerol O-acyltransferase 2), Mm00499536_m1; MGAT2 (monoacylglycerol O-acyltransferase 2), Mm00624192_m1; CPT1A (carnitine palmitoyltransferase 1a, liver), Mm01231183_m1; ACADL (acyl-coenzyme A dehydrogenase, long-chain), Mm00599660_m1; ACADM (acyl-coenzyme A dehydrogenase, medium chain), Mm01323360_g1; ACACA (acytelyl-CoA acyltransferase 1), Mm00624282_m1; ACOT1 (acyl-CoA thioesterase 1), Mm01622471_s1; ACOT2 (acyl-CoA thioesterase 2) Mm01622461_s1; ACAC (acetyl-coenzyme A acyltransferase 1B), Mm00728805_s1; ACOX1 (acyl-coenzyme A oxidase 1, palmitoyl), Mm01246834_m1; SCARB1 (scavenger receptor class B, member 1), Mm00450234_m1; FGF15 (fibroblast growth factor 15), Mm00433278_m1; SHP (nucleolar receptor subfamily 0, member B, member 2), Mm00442278_m1; PPARA (peroxisome proliferator-activated receptor α), Mm00440939_m1; FXR (NR1H4, nuclear receptor subfamily 1, group H, member 4), Mm00436425_m1; APOA4 (apolipoprotein A-IV), Mm00431814_m1; GCG (glucagon), Mm00735956_m1; PPARC (peroxisome proliferator-activated receptor gamma), Mm00309687_m1; TAXOD (transcribed at high levels in diabetic kidney disease), Mm00464771_m1; ELOVL2 (elongation of very long chain fatty acids 2), Mm01253490_m1; ELOVL3 (elongation of very long chain fatty acids 3), Mm00419850_m1; ELOVL4 (elongation of very long chain fatty acids 4), Mm00419851_m1; ELOVL5 (elongation of very long chain fatty acids 5), Mm00419852_m1; ELOVL6 (elongation of very long chain fatty acids 6), Mm00419853_m1; HSL (hormone sensitive lipase), Mm00754840_m1; LPL (lipoprotein lipase), Mm00446857_m1; PSMD12 (proteasome subunit, non-ATPase 12), Mm00472611_m1; PPARA (peroxisome proliferator-activated receptor α), Mm00440939_m1; FXR (NR1H4, nuclear receptor subfamily 1, group H, member 4), Mm00436425_m1; APOA4 (apolipoprotein A-IV), Mm00431814_m1; GCG (glucagon), Mm00735956_m1; PPARC (peroxisome proliferator-activated receptor gamma), Mm00309687_m1; TAXOD (transcribed at high levels in diabetic kidney disease), Mm00464771_m1; ELOVL2 (elongation of very long chain fatty acids 2), Mm01253490_m1; ELOVL3 (elongation of very long chain fatty acids 3), Mm00419850_m1; ELOVL4 (elongation of very long chain fatty acids 4), Mm00419851_m1; ELOVL5 (elongation of very long chain fatty acids 5), Mm00419852_m1; ELOVL6 (elongation of very long chain fatty acids 6), Mm00419853_m1; HSL (hormone sensitive lipase), Mm00754840_m1; LPL (lipoprotein lipase), Mm00446857_m1; PSMD12 (proteasome subunit, non-ATPase 12), Mm00472611_m1.

Statistical Analysis—Statistical tests were performed (GraphPad Prism software, version 6.0, GraphPad, La Jolla, CA).

Results

GPAT3 Was Abundantly Expressed in the Absorptive Epithelial Cells of the Small Intestine—Previous studies have documented a role for GPAT3 in the synthesis of triacylglycerol in adipocytes (17, 18, 28, 29). To further investigate the physiological role of GPAT3, the level of GPAT3 mRNA was determined in a panel of tissues. In line with previous reports, the murine Gpat3 gene was abundantly expressed in the adipose, heart, and small intestine (Fig. 1A). GPAT3 protein was detectable in microsomes isolated from the jejunum. A distribution profile of GPAT3 along the small intestine exhibited expression confined to the central (jejunum) portion of the small intestine that was strongly induced after a 6-h fast (Fig. 1B). As expected, immunoreactive GPAT3 was absent in Gpat3−/− mice (Fig. 1B). Activity assays demonstrated an ~40% decrease in total GPAT activity in the Gpat3−/− jejunum (6.44 ± 0.35 and 4.05 ± 0.45 nmol/min/mg in wild-type and Gpat3−/−, respectively) mice. The decrease in jejunal total GPAT activity was primarily attributable to a reduction in NEM-sensitive activity. No significant changes were observed in jejunal NEM-resistant activity as well as total and NEM-sensitive GPAT activities in the duodenum and ileum of wild-type and Gpat3−/− mice (Fig. 1C). Taken together these data indicate that GPAT3 is abundantly expressed in the jejunal region of the mouse small intestine.

Immunohistochemistry further localized GPAT3 to the absorptive epithelium of the villus (Fig. 2A). Immunoreactive staining was completely absent in Gpat3−/− mice. During lipid absorption, enterocytes transiently store TGs in CLD before packaging and secretion in chylomicron particles (30). To determine if GPAT3 localizes to lipid droplets during the absorptive period, wild-type and Gpat3−/− mice were administered an oral bolus of olive oil after a 6-h fast. GPAT3 was exclusively expressed on the apical region of the wild-type enterocytes (Fig. 2B, left panel). We further monitored location of the PAT (perilipin) family protein TIP47 (PLIN3), which has been reported to be highly expressed in the enterocytes and localizes to CLD after acute high fat feeding (30, 31). In similarity to GPAT3, TIP47 was readily detectable in the apical region of the enterocytes with dense staining around the surface of the lipid droplets (Fig. 2B, right panel). In these lipid-laden villi, intense TIP47 staining was evident on the surface of the lipid droplet, but although we could identify GPAT3 immunofluorescence in the proximity to lipid droplets (inset, left panel, Fig. 2C). It was not possible to definitively colocalize GPAT3 and TIP47 to CLD Fig. 2C, left panel). The pattern of TIP47 staining did change in Gpat3-null enterocytes (Fig. 2C, right panel). These data demonstrate that GPAT3 and TIP47 localize to the apical surface of the enterocyte.

Gpat3−/− Mice Exhibited Dysregulated Intestinal Lipid Absorption—Given the abundant expression of GPAT3 in the small intestine, we next examined whether loss of GPAT3 function modulated intestinal lipid handling. In fasted animals, there were no differences in intestinal TG levels between wild-type and Gpat3−/− mice. However, TG content was significantly elevated in the ileum of the Gpat3−/− mice (Fig. 3A). Titration of the lipid bolus administered an oral bolus of olive oil after a 6-h fast. GPAT3 was significantly elevated in the ileum of the Gpat3−/− mice (Fig. 3A). Titration of the lipid bolus administered an oral bolus of olive oil after a 6-h fast. GPAT3 was significantly elevated in the ileum of the Gpat3−/− mice (Fig. 3A).
As such, these findings may be most relevant to conditions where large amounts of fat are ingested. To further characterize this phenotype, plasma lipoprotein clearance was blocked by administering poloxamer 407, and tracer studies were performed in vivo using [3H]triolein combined with an oral olive oil bolus. As expected, wild-type mice displayed accumulation of radiolabeled lipid in the plasma; however, the rate of appearance was significantly reduced in Gpat3−/− mice (Fig. 3C). Plasma FPLC fractionation showed reduced radiolabel associated with TG-rich lipoproteins (fractions 5–10, Fig. 3D). There was no difference in fecal tracer counts between the groups (data not shown). Distribution of radiolabeled lipid throughout the small intestine showed increased 3H-labeled lipids in the jejunal portion of the small intestine accompanied by decreased lipid in the proximal segments (Fig. 3E).

To further examine the effects on the intestinal function, immunohistochemistry and electron microscopy were performed on the jejunum. Histology revealed extensive lipid droplet accumulation within the enterocytes of both wild-type and Gpat3−/− mice. Although wild-type mice showed marked lipid staining in the lamina propria, Gpat3−/− mice exhibited a striking reduction in transient lipids (Fig. 4A). Histological scoring confirmed that the Gpat3−/− mice, compared with the wild-type animals, exhibited significantly lower lipids in the lamina propria and intercellular space versus total amount of lipids present in the jejunal segments (Fig. 4B). Electron microscopy using TEM for single cell analysis revealed reduced trafficking of lipids, presumably chylomicrons in the basolateral surface of the enterocytes in the Gpat3−/− mice compared with the wild-type mice (Fig. 4C). Collectively, these observations suggest that Gpat3 deficiency results in the temporal accumu-
FIGURE 2. **GPAT3 is expressed in the absorptive epithelial cells of the small intestine.** Animals were maintained on standard chow and fasted for 6 h before administration of an oral olive oil bolus. Small intestines were collected 2 h post-dose as described in Fig. 1. **A,** immunohistochemistry of a representative jejunal segment showing GPAT3 protein expression and localization in WT (left panel) and **Gpat3**-null (right panel) enterocytes. Scale bars = 50 μm. **B,** individual immunofluorescence staining for GPAT3 (green, left panel) and TIP47 (red, right panel) in wild-type intestinal epithelium. **C,** merged staining for GPAT3 (green) and TIP47 (red) in wild-type (left panel) and **Gpat3**-null (right panel) intestinal epithelium. Nuclei are stained blue with DAPI (4',6-diamidino-2-phenylindole). Scale bars = 20 μm.
lation of lipids within the enterocytes during increased dietary lipid loaded states.

GPAT3 Deficiency Results in Alterations in Lipid Metabolism and Enteroendocrine Hormone Production—In the Gpat3/H11002/H11002 mice, the blunted postprandial plasma TG excursion after olive oil bolus was associated with increases in duodenal, jejunal, and ileal lipid content (Fig. 5A). Similar to mice maintained on standard chow, Gpat3/H11002/H11002 mice administered an olive oil bolus exhibited elevated TG levels in the distal region of the small intestine (Figs. 3A and 5A). In line with GPAT3 protein expression, the Gpat3 gene was most abundantly expressed in jejunum (Fig. 5B). The loss of Gpat3 expression was accompanied by a compensatory increase in Gpat4 mRNA (Fig. 5C). Small intestinal levels of FFAs were modestly but significantly elevated in Gpat3/H11002/H11002 compared with the wild-type animals (Table 1). Interestingly, although the relative abundance of long chain polyunsaturated fatty acids such as eicosapentaenoic (C20:5), docosapentaenoic (C22:5), and docosahexaenoic acid (C22:6) was low throughout the small-intestine, Gpat3−/− mice had reduced levels of these lipids compared with the wild type in the jejunum and ileum (Table 1). The changes in TG and FFA were accompanied by significant increases in intestinal long chain acylcarnitines in Gpat3−/− mice (Table 2). There were no changes in short chain acylcarnitines (data not shown). These data accompanied with the increases in total FFA levels suggest increased intestinal fatty acid oxidation in the Gpat3−/− mice.

To explore any potential adaptive changes that take place in the intestinal mucosa of Gpat3−/− mice, we evaluated the expression of multiple genes encoding proteins involved in lipid metabolism. Compared with the wild-type mice, significant decreases in the expression of Dgat1, Dgat2, and Mgat2 were evident in Gpat3−/− jejunum (Fig. 5D). In line with increased FFA and long-chain acyl carnitine levels, genes encoding proteins involved in fatty acid oxidation, such as carnitine palmitoyltransferase 1 (Cpt1a, ~1.3-fold), acyl-CoA thioesterase (Aco1, ~4-fold), and acyl-CoA oxidase 1 (Acox1, ~3.5-fold) were significantly induced in Gpat3−/− mice (Fig. 5D). The level of mRNA encoding fibroblast growth factor 15 (Fgf15) and...
small heterodimer partner (Shp), which are both central components in bile acid homeostasis, were elevated by ~2.5–3-fold. Expression of the nuclear receptor peroxisome proliferator-activated receptor α (Ppara) was significantly decreased in Gpat3<sup>−/−</sup> mice compared with the wild-type animals (Fig. 5D).

Because we observed deregulated lipid secretion and accumulation of TG in the enterocytes of Gpat3<sup>−/−</sup> mice after an oral lipid bolus, we examined the effect of GPAT3 deletion on the expression of apolipoprotein A4 (Apoa4) and glucose-dependent insulino tropic polypeptide (Gip). The Gip and Apoa4 genes are preferentially expressed in the small intestine where the synthesis and secretion of these proteins is reported to be modulated by dietary lipids (33–35). Expression of the Apoa4 and Gip genes was significantly induced throughout the small intestine in Gpat3<sup>−/−</sup> mice compared with the wild-type animals (Fig. 5E). Interestingly, expression of the glucagon gene (Gcg), which encodes glucagon-like peptide-1 (GLP-1) in the gut, was also increased in the intestine of Gpat3<sup>−/−</sup> mice (Fig. 5E). Gpat3<sup>−/−</sup> plasma GIP levels revealed a ~2-fold increase compared with the wild-type animals when either fasted for 6 h (100 ± 17 and 201 ± 38 pg/ml in wild-type and Gpat3<sup>−/−</sup> mice, respectively, p < 0.05) or allowed food ad libitum (105 ± 14 and 216 ± 27 pg/ml in wild-type and Gpat3<sup>−/−</sup> mice, respectively, p < 0.01) (Fig. 5F). When mice were gavaged with olive oil there was a marked increase in circulating GIP levels; however, the genotype effect apparent in mice maintained on standard chow was lost (Fig. 5G). We were unable to detect any changes in total GLP in either setting (data not shown).

**Attenuation of Postprandial Triglyceride Excursion Is Secondary to Decreased Secretion and Not Differential Absorption from Intestinal Lumen**—To better understand the effects of GPAT3 deletion on lipid absorption and secretion, we performed a series of pulse-chase studies using [3H]oleate and [14C]cholesterol in ex vivo cultures of murine enterocytes derived from wild-type and Gpat3<sup>−/−</sup> mice (Fig. 6A). Radiolabeled oleate uptake into primary enterocytes, as determined by [3H]oleate incorporation into the cellular lipid fraction, was not significantly affected by loss of GPAT3 (Fig. 6B, inset). However, when the distribution of [3H]oleate into cellular lipids was examined, a marked redistribution was apparent. Thus, in Gpat3<sup>−/−</sup> enterocytes, incorporation of [3H]oleate into TG was reduced with a parallel increase in the production of 3H-labeled

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**FIGURE 4. Lipid uptake and distribution in the intercellular space of the intestinal villi.** Mice were fasted (6 h) before dosing with a single oral bolus of olive oil. Small intestinal segments were collected 2 h post-dose. A, H&E staining (top panel, 10×) and osmium paraphenylenediamine staining (bottom panel, 10×) of the jejunal segments after olive oil feeding in wild-type and Gpat3<sup>−/−</sup> mice. Insets (40X, scale = 25 μm) show magnified images of lipid traffic in the lamina propria (red arrows) in the wild-type but not in the Gpat3<sup>−/−</sup> mice with respect to lipids retained within the enterocytes (blue arrow), B, lipid scores in the lamina propria and intercellular space (top panel); ratio of lipids in the intercellular space with respect to the total lipids in the intestinal epithelium (bottom panel). Scores were assigned based on whole slide analysis for n = 10 animal/group. C, representative TEM electron micrographs of epithelium in the jejunal intestinal villi. Extracellular chylomicrons (arrows) are prominent in WT (A, left panel) compared with Gpat3<sup>−/−</sup> (B, right panel). Accumulation of microvesicular bodies (asterisk (*)) were present in Gpat3<sup>−/−</sup>. Scale bars = 4 μm.
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FIGURE 5. Effects of GPAT3 deletion on intestinal lipid profile, gene expression, and gut nutrient sensing. Fasted mice were gavaged with olive oil and sacrificed 2 h later. The small intestine was harvested, and mucosa was collected from the indicated segments for quantification of triglyceride levels or gene expression. Data are expressed as fold change relative to jejunal expression.

TABLE 1

Intestinal FFA profile determined by LC-MS analysis

Animals were administered an oral olive oil bolus (15 ml/kg) after a 6-h fast. Before the study wild type and animals were maintained on standard laboratory chow. Data are the mean ± S.D.

| Species | Duodenum | Jejunum | Ileum |
|---------|----------|---------|-------|
|         | Wild type (n = 6) | Gpat3−/− (n = 7) | Wild type (n = 6) | Gpat3−/− (n = 7) | Wild type (n = 6) | Gpat3−/− (n = 6) |
| 14:0    | 230 ± 41 | 359 ± 98 | 413 ± 90 | 584 ± 140^a | 623 ± 105 | 949 ± 82^a |
| 16:0    | 12,462 ± 2,286 | 14,151 ± 1,906 | 12,517 ± 1,362 | 14,505 ± 2,812 | 12,428 ± 2,513 | 14,961 ± 2,259 |
| 16:1    | 1,319 ± 183 | 1,689 ± 321^a | 1,345 ± 147 | 1,916 ± 590^a | 1,685 ± 58.0 | 2,542 ± 393^b |
| 18:0    | 6,083 ± 434 | 6,038 ± 540 | 6,433 ± 400 | 6,103 ± 331 | 5,829 ± 624 | 5,033 ± 654 |
| 18:1    | 36,157 ± 9,571 | 37,852 ± 8,491 | 24,428 ± 5431 | 27,015 ± 6128 | 17,849 ± 1,649 | 15,762 ± 1,665 |
| 18:2    | 8,142 ± 964 | 8,587 ± 1144 | 6,529 ± 632 | 7,226 ± 1446 | 6,244.5 ± 579 | 7,451 ± 1028 |
| 20:4    | 1,520 ± 572 | 1,266 ± 623 | 1,113 ± 206 | 907 ± 168 | 1,114 ± 325 | 701 ± 120 |
| 20:5    | 179 ± 58 | 122 ± 91 | 132 ± 11 | 90 ± 19^a | 132 ± 30 | 92 ± 17^a |
| 22:5    | 115 ± 38 | 95 ± 48 | 126 ± 23 | 110 ± 334 | 146 ± 34 | 108 ± 26 |
| 22:6    | 763 ± 252 | 517 ± 318 | 640 ± 179 | 432 ± 134^a | 640 ± 165 | 368 ± 136^a |

*p < 0.05, as determined by Student’s t test.
*p < 0.001, as determined by Student’s t test.
*p < 0.01, as determined by Student’s t test.
TABLE 2
Intestinal long chain acylcarnitine profile
Animals were administered an oral olive oil bolus (15 ml/kg) after a 6-h fast. Before the study wild-type and animals were maintained on standard laboratory chow. Data are the mean ± S.D.

| Species | Duodenum | Jejunum | Ileum |
|---------|----------|---------|-------|
|         | Wild type | Gpat3−/− | Wild type | Gpat3−/− | Wild type | Gpat3−/− |
|         | (n = 6)   | (n = 7)  | (n = 7)   | (n = 6)  | (n = 7)   | (n = 6)  |
| 14:0    | 75.6 ± 21 | 73.6 ± 15 | 45.2 ± 12 | 73.5 ± 33a | 54.5 ± 18 | 83.4 ± 23a |
| 16:0    | 124 ± 25  | 111 ± 28  | 92.9 ± 18 | 112 ± 30a | 1219 ± 19 | 126 ± 13  |
| 16:1    | 6.2 ± 1.1 | 5.2 ± 1.0  | 4.4 ± 0.61 | 4.8 ± 0.98 | 5.4 ± 0.81 | 5.1 ± 0.46 |
| 18:2    | 31.1 ± 7.2 | 23.0 ± 7.9a | 18.5 ± 5.6 | 22.4 ± 8.0a | 30.5 ± 7.8 | 25.6 ± 5.7 |
| 18:1    | 105 ± 19   | 92.8 ± 30  | 85.3 ± 26  | 118 ± 45a | 98.2 ± 19  | 105 ± 21  |
| 18:0    | 106 ± 25   | 119 ± 20   | 98.6 ± 34  | 150 ± 65a | 94.0 ± 31  | 161 ± 28b |
| 20:6    | 3.9 ± 0.71 | 3.3 ± 0.42 | 4.4 ± 0.28 | 4.0 ± 1.01 | 4.3 ± 0.61 | 3.5 ± 0.45 |
| 20:5    | 2.4 ± 0.56 | 1.8 ± 0.36a | 1.99 ± 0.25 | 1.9 ± 0.45 | 2.2 ± 0.71 | 1.6 ± 0.30 |
| 20:4    | 6.1 ± 0.54 | 5.4 ± 1.3  | 5.3 ± 0.57  | 5.9 ± 0.75 | 7.5 ± 1.1  | 6.5 ± 1.1  |
| 20:3    | 1.4 ± 0.25 | 1.3 ± 0.31 | 1.4 ± 0.18  | 1.3 ± 0.27 | 2.1 ± 0.41 | 1.7 ± 0.30 |
| 20:2    | 5.6 ± 0.85 | 5.6 ± 1.2  | 6.3 ± 1.7   | 6.9 ± 2.0  | 5.9 ± 1.6  | 6.5 ± 1.1  |
| 20:1    | 81.0 ± 8.2 | 84.3 ± 22  | 81.8 ± 24   | 85.3 ± 28  | 67.5 ± 16  | 74.1 ± 8.0 |
| 20:0    | 17.6 ± 4.5 | 16.9 ± 4.7 | 14.4 ± 4.1  | 15.2 ± 4.3 | 12.3 ± 3.8 | 10.5 ± 1.6 |
| 22:0    | 6.2 ± 1.9  | 5.9 ± 1.9  | 5.00 ± 0.90 | 4.7 ± 1.3  | 4.9 ± 1.5  | 3.5 ± 0.45 |

* p < 0.05, as determined by Student’s t test.
** p < 0.01, as determined by Student’s t test.

**FIGURE 6.** Ex vivo flux analysis in primary enterocytes from wild-type and Gpat3−/− mice. A, flowchart of ex vivo uptake or secretion studies using dual radioisotopes. B, distribution of [3H]oleic acid (OA) in the intracellular lipid pool. PC, phosphatidylcholine; PE, phosphatidylethanolamine. The inset shows total cellular lipid counts. C, total lipids secreted in the media (inset) and incorporation of tracer in intracellular CE. Total cellular lipids were determined by liquid scintillation counting, whereas the incorporation into TG and phospholipid was monitored by TLC. ns, not significant. E, [14C]cholesterol counts in media fractions subject to density gradient ultracentrifugation. Large chylomicrons, CM (Sf > 400); small chylomicrons, CM (Sf = 60–400); CM (Sf = 20–60); fraction 4–10 representing HDL. Data are the mean ± S.E. (n = 4). *, p < 0.05; ***, p < 0.001 by either one-way analysis of variance followed by Newman-Keuls multiple comparison test or Student’s t test.

cholesteryl ester (Fig. 6B). Secretion of 3H-labeled lipid was examined during the chase period. There were no significant differences in total lipid secretion (Fig. 6C, inset). In enterocytes derived from the wild-type mice, TG was the primary lipid species secreted into the media; however, in Gpat3−/− enterocytes TG secretion was reduced, and there was a compensatory increase in the secretion of labeled cholesteryl ester (CE) and phosphatidylethanolamine (PE) (Fig. 6C). When [14C]cholesterol was used as the tracer, a significant increase in the production of intracellular [14C]CE in Gpat3−/− cells was observed.
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(Fig. 6D). Finally, media were collected, and lipoprotein fractions were separated by density gradient fractionation. [14C]Cholesterol secretion in the large (floatation coefficient, Sf > 400) and small (Sf = 60–400) chylomicron (CM1, CM2) population was significantly increased in media derived from Gpat3−/− enterocytes. CE secretion onto HDL (fractions 4–10) was unaffected by disruption of GPAT3 (Fig. 6E). Taken together these data suggest that the reduced appearance of TG in the plasma after an oral lipid bolus in Gpat3−/− mice was as a result of reduced TG secretion, whereas the uptake of FFA from the intestinal lumen remains largely intact. In the absence of GPAT3, excess FFA derived either directly from luminal lipid absorption or liberated within the cell after hydrolysis of stored lipid is esterified with cholesterol, thereby protecting the cell from the accumulation of potentially harmful FFA. Ultimately, the excess CE produced in GPAT3-null mice is secreted on CM particles resulting in the production of cholesterol-rich CM.

GPAT3 Abrogation Attenuated Plasma Cholesterol, Exacerbated Hepatic Steatosis, and Altered Cholesterol-Bile Acid Homeostasis—Based on the findings that loss of GPAT3 in the intestine results in altered lipid secretion and the dysregulation of multiple genes involved in enteric fatty acid and bile acid homeostasis, we further evaluated the physiological role of GPAT3 in whole-body lipid metabolism in mice fed a high fat diet. We found that the loss of GPAT3 activity led to pronounced significant but comparable body weight gain in both wild-type and Gpat3−/− mice (Fig. 7A). No changes in fasted plasma TG were observed (Fig. 7B); however, Gpat3−/− mice showed decreased total plasma cholesterol under both chow and WTD-fed conditions (Fig. 7B). Size fractionation of plasma lipoproteins by gel filtration to analyze lipid distribution in different lipoprotein classes showed no differences in VLDL-TG (data not shown); however, lower HDL-cholesterol was observed in the Gpat3−/− mice (Fig. 7D). Plasma alanine transaminase (ALT) levels, a marker of liver damage, were increased in Gpat3−/− mice compared with wild-type mice (Fig. 7E). Both wild-type and Gpat3−/− mice fed on WTD exhibited pronounced hepatic steatosis compared with the chow-fed wild-type and Gpat3−/− mice; however, the livers of WTD-fed Gpat3-null mice accumulated significantly more lipid compared with the WTD-fed wild-type livers. HPLC analysis showed increases in TG, DAG, and CE in WTD-fed Gpat3−/− cohorts compared with the wild-type animals (Fig. 7, F and G). The livers of Gpat3−/− mice exhibited significantly high levels of cholesteryl ester when maintained on a WTD (Fig. 7H).

In the liver biliary cholesterol secretion or catabolism to bile acids represent the primary routes for elimination for excess hepatic cholesterol. Given the elevated hepatic CE (Fig. 7H) and dysregulation of the Fgf15 and Shp genes in the small intestine of Gpat3−/− mice (Fig. 5D), we examined the effects of GPAT3 deletion on bile acid homeostasis. Wild-type and Gpat3−/− mice exhibited similar levels of plasma total bile acids (TBAs) when maintained on standard chow (Fig. 7I). WTD feeding resulted in a ∼2.6-fold reduction in plasma TBAs in wild-type mice; however, plasma TBAs in Gpat3−/− cohorts were maintained at a comparable level with that observed in chow-fed animals (Fig. 7I). In contrast, hepatic TBA was elevated after WTD feeding, and the level was significantly (∼1.9-fold) lower in Gpat3−/− mice (Fig. 7J). These data suggest that disruption of the Gpat3 gene results in alterations of entero-hepatic bile acid homeostasis.

Discussion

For more than 40 years it has been recognized that the monoglycerol pathway is the primary route for the resynthesis of dietary triacylglycerol in the enterocyte. However, MGAT2-deficient mice exhibit minimal quantitative effects on lipid absorption (8). These data combined with earlier studies suggest the existence of alternative, MGAT-independent pathways for lipid absorption (3–7). In this study we utilized Gpat3−/− mice to characterize an intestinal glycerol 3-phosphate pathway.

In mammals, four distinct GPAT genes have been identified (12). GPAT3 was initially characterized by Cao et al. (17) and shown to be induced in adipocytes after exposure to the PPARα agonist rosiglitazone. The Gpat3 gene was found to be highly expressed in adipose and small intestine. Our study demonstrates that the GPAT3 protein is abundantly expressed in the enterocytes of the jejunum where it is strongly up-regulated during fasting. Targeted disruption of the Gpat3 gene resulted in ∼40% reduction in total and NEM-sensitive microsomal GPAT activity in the jejunum. In line with the protein expression data, microsomal GPAT activity was preserved in the duodenum and ileum of Gpat3−/− mice. These data suggest that either GPAT4 is the primary contributor to GPAT activity in the duodenum and ileum or that there is a compensatory increase in Gpat4 activity or another unidentified GPAT after loss of GPAT3 activity. Although the majority of GPAT3 staining within the enterocyte appeared to be diffuse, dense punctate staining of GPAT3 was evident in proximity to the surface of the lipid droplet after an oral lipid bolus. Although we did not perform a detailed analysis of the subcellular localization of GPAT3, it is possible that, in analogy to closely related GPAT4, the GPAT3 protein relocates from the endoplasmic reticulum to the surface of the lipid droplet upon cellular lipid loading (36).

The intestinal expression profile of GPAT3 suggests a role in intestinal lipid absorption. In support of this, Gpat3−/− mice exhibit a reduction in plasma TG excursion when administered an oral lipid bolus that was accompanied by increased accumulation of TG in enterocytes. There were no differences in intestinal TG levels between wild-type and Gpat3−/− mice when animals were fasted, and there was no evidence of fecal lipid loss after an acute lipid load. GPAT3-deficient mice displayed a modest increase in enteric free fatty acid levels that was paralleled by induction of multiple genes involved in fatty acid oxidation (e.g. Cpt1a, Acot1, Acox1) and elevations in acylcarnitines, suggesting an adaptive response to protect against the accumulation of non-esterified fatty acids. Although we observed increased TG accumulation in the small intestine of Gpat3−/− mice, there was a marked decrease in lipids in the lamina propria and intercellular space, suggesting that GPAT3 plays a role in chylomicron assembly or subsequent secretion into the lymph via the basolateral surface. This hypothesis was supported by a series of pulse-chase studies performed in...
enterocytes derived from wild-type and Gpat3−/− mice. Taken together, these data suggest that GPAT3 plays a role in multiple facets of intestinal lipid metabolism. Thus, in the face of elevated lipid levels, GPAT3 localizes to the CLD where it contributes to the short term storage of dietary lipids in the enterocyte. In addition, GPAT3 appears to be involved in the appropriate partitioning of lipid for the formation and secretion of chylomicrons, possibly through participation in a lipolysis reesterification process analogous to that proposed for VLDL secretion in the liver.

Overall, the intestinal phenotype observed in the Gpat3−/− is qualitatively similar to that observed in Dgat1−/− mice (8, 32, 37). Thus, Dgat1-null animals exhibited decreased plasma chyomicronemia after an oral lipid bolus and accumulate TG in the gut epithelia (32). Disruption of the Dgat1 locus or pharmacological inhibition of DGAT1 is associated with an improved

FIGURE 7. Gpat3−/− mice exhibit dysregulated hepatic cholesterol and bile acid homeostasis. WT and Gpat3−/− mice were fed a WTD for 4 weeks. Mice were fasted for 6 h on the day of sacrifice. A, body weight change (percent change calculated from initial weight) during the course of the diet. Plasma TG (B) and total plasma cholesterol (C) in chow- and WTD-fed animals. D, distribution of cholesterol in different plasma lipoprotein populations separated by FPLC. Plasma alanine transaminase (ALT) (E) and hepatic TG (F), DAG (G), and hepatic cholesterol (CE, cholesteryl ester; FC, free cholesterol) (H). Plasma (I) and hepatic total bile acids (TBA) (J) levels. All data are the mean ± S.E. (n = 8). *, p < 0.05; **, p < 0.01 by Student’s t test.
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metabolic profile upon high fat diet feeding, including decreased body weight gain, increased insulin sensitivity, and reduced hepatic lipid accumulation (32, 37–40). However, in contrast, when maintained on high fat diets, Gpat3−/− mice exhibit weight gain in line with wild-type animals and accumulate excess hepatic lipid in the form of TG, DAG, and CE. The reason for this difference is unclear; however, it could relate to divergent effects on enteroendocrine hormone release. Thus, although inhibition of DGAT1 is associated with suppression of GIP and robust increases in GLP-1 and the anorectic hormone peptide YY (PYY) (38, 41), loss of Gpat3 results in elevated levels of circulating GIP. We were unable to discern an effect on GLP-1 levels and mice containing a disrupted GIP receptor gene (Gipr−/−) are protected from diet-induced obesity (42). The molecular mechanism underlying this observation could relate to differences in the expression of the Gpat3 and Dgat1 genes throughout the small intestine or the liberation of specific lipids after blockade of these enzymes (43).

In preliminary experiments we determined that an oral lipid bolus of 15 ml/kg olive oil was required to elicit significant differences between wild-type and Gpat3−/− mice in the plasma lipid excursion (data not shown). Although this degree of oral lipid-loading has been previously described (20, 21), it is markedly higher than the dose of olive oil (~4–8 ml/kg) required to delineate the role of MGAT2 and DGAT1 in fat absorption (8, 32). Thus, it appears that under normal physiological conditions, the GPAT3 pathway is a minor pathway for lipid absorption compared with the canonical MGAT2-DGAT1 pathway. It is possible that GPAT3 is a component of a rescue system that evolved to ensure sufficient absorption of ingested lipid after a fat-rich meal. Alternatively, intestinal GPAT3 could play a role in the absorption of specific species. Notably, the acute lipid absorption studies we performed exclusively utilized olive oil, which is predominantly composed of oleate (C18:1) and, to a lesser extent, palmitate (C16:0).

In summary, we have characterized a role for GPAT3 in the regulation of intestinal lipid absorption. Targeted disruption of the Gpat3 locus results in dysregulation of TG secretion from the small intestine.

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