Regulation of Triglyceride Metabolism
II. Function of mitochondrial GPAT1 in the regulation of triacylglycerol biosynthesis and insulin action

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Consistent with the hydrophobicity analysis, protease-protection experiments and topography studies using epitope-tagged, recombinant GPAT1 show two transmembrane domains (aa 472–493 and aa 576–592) with both the NH2 and COOH termini facing the cytosol and a loop (aa 494–575) facing the intermembrane space (4). The active-site residues (motifs I–IV, Table 1) lie within the NH2-terminal domain, but mutagenesis studies suggest that both the loop and the COOH-terminal domain are required for enzyme activity. The COOH-terminal domain interacts physically with the NH2-terminal domain and probably contributes to substrate binding and catalysis (23).

Members of the pfam 01553 family of glycerolipid acyltransferases contain four conserved motifs (Table 1) that were first identified by a bioinformatics approach in 1997 (4). Subsequently, mutagenesis of invariant residues in these motifs verified that these highly conserved regions were essential for the activity of Escherichia coli GPAT (PisB), mouse GPAT1, and human dihydroxyacetone-phosphate acyltransferase (DHAPAT) (4). Residues important for catalysis are the invariant histidine and aspartate in motif I, the phenylalanine, glycine, and arginine in motif III, and the proline in motif IV (4). Amino acids important for binding glycerol-3-phosphate are the phenylalanine and arginine in motif II and the glutamate and serine in motif III (4). A diverse array of acyltransferases have been assigned to the pfam 01553 family, but biochemical activities have been verified only for GPAT1 and -2, lysophosphatic acid acyltransferase (AGPAT)-1 and -2, DHAPAT, lysophosphatidylcholine acyltransferase, lysophosphatidylglycerol acyltransferase, and monolysocardiolipin acyltransferase (2, 4, 19). Although AGPAT-6 and -8 were originally identified as lysophosphatidic acid acyltransferases, each has GPAT activity, and we will term them GPAT4 and GPAT3, respectively, in this review. Despite its homology to other lipid acyltransferases, tafazzin exhibits a transacylase activity (25). The amino acids important for the catalytic activity of most pfam 01553 family members have yet to be identified.

LysoPhosphatidate Is Transported From Mitochondria to Endoplasmic Reticulum for the Final Steps of Complex Lipid Synthesis

The location of GPAT1 on the outer mitochondrial membrane is surprising because the enzymes that catalyze the final biosynthetic steps that form TAG and the glycerophospholipids are located in the endoplasmic reticulum membrane (4). Although GPAT1 and -2 are located in the mitochondrial outer membrane, their LPA product is clearly used for the synthesis of complex glycerolipids other than mitochondrial cardiolipin. Transport of LPA to the endoplasmic reticulum appears to be...
Acyltransferase motifs in GPAT isoforms

<table>
<thead>
<tr>
<th>Motif P</th>
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<tr>
<td>GPAT1b (NM_008149)</td>
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<td>GPAT2 (XM_130488)</td>
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<td>GPAT3α (NP_766303)</td>
<td>64 HRTRVD</td>
<td>109 IFIHR</td>
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<tr>
<td>GPAT4α (NM_018743)</td>
<td>249 HTSPID</td>
<td>288 VWFER</td>
<td>319 FPETCCIN</td>
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</tbody>
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*Acyltransferase motifs as described in Ref. 4. Boldface residues are highly conserved across all pfam 01553 family members. Underlined amino acids are important for glycerol-3-phosphate acyltransferase 1 (GPAT1) activity as determined by mutagenesis. Previously called mtGPAT or mtGAT. Also identified as xGPAT1 (10). Originally identified as AGPAT8, but has GPAT activity (1). Originally identified as AGPAT6.

GPAT1 is Regulated in a Manner Consistent With a Role in Initiating TAG Synthesis

Although the specific cellular functions of each GPAT isoform are not yet understood, strong experimental evidence supports the participation of GPAT1 in de novo TAG synthesis under conditions of dietary carbohydrate or fat excess. Adenovirus- and plasmid-mediated overexpression of GPAT1 in Chinese hamster ovary and HEK293 cells (4) and in primary rat hepatocytes (13, 15) increases TAG content and oleate incorporation into TAG and variably affects very-low-density lipoprotein (VLDL) secretion. In contrast, in Gpat1 knockout mice, hepatic TAG mass is 40% lower than in wild-type mice (4).

Enzymes required for TAG synthesis are coordinately regulated through the counterregulatory hormones insulin and...
Glucagon. Insulin enhances and both fasting and diabetic states reduce the activities of lipogenic enzymes, including GPAT1. Thus, in rat liver and adipose tissue, a 48-h fast decreases GPAT1 protein expression and activity >30%, and sucrose refeeding of previously fasted rats increases GPAT1 protein expression and activity more than twofold in liver (4). Consistent with nutritional regulation, in perfused rat liver, insulin increases GPAT1 activity 34% (4). In addition, streptozotocin-induced diabetes decreases GPAT1 activity in rat epididymal fat 60% and insulin administration restores GPAT activity (4).

Changes in GPAT1 Activity Are Mediated Transcriptionally

When previously fasted mice are refed a high-carbohydrate diet, hepatic Gpat1 mRNA levels increase >20-fold owing to enhanced transcription rates (4). This refeeding-induced increase in Gpat1 message is not observed in streptozotocin-diabetic mice unless insulin is administered. Insulin regulation of Gpat1 is mediated by induction of sterol regulatory element binding protein-1 (Srebf-1) transcription, the main transcriptional regulator of Gpat1, and glucagon-mediated elevation of cAMP opposes the action of insulin (7). The mouse Gpat1 promoter region contains three sterol regulatory element binding protein (SREBP-1) consensus sites that are responsible for SREBP-1-mediated transactivation (4); thus ectopic expression of SREBP-1c in 3T3-L1 adipocytes causes Gpat1 mRNA expression to increase 6.7-fold. Similarly, in Srebf-1a and -1c transgenic mice, the hepatic Gpat1 message increases 10- and 3-fold, respectively (4). Although LXR agonists have been reported to upregulate Gpat1, LXR does not directly transactivate Gpat1 but rather upregulates Srebf-1c (4). Furthermore, although carbohydrate feeding increases Gpat1 mRNA upregulation proceeds through insulin-mediated transactivation of Srebf-1c rather than through the carbohydrate response element (3).

GPAT1 Is Required for Normal Metabolism of Acyl-CoA and Its Absence Increases Fatty Acid Oxidation and Oxidative Stress

Gpat1−/− mice are healthy and breed normally, indicating that GPAT1 is not essential for embryogenesis, reproduction, or synthesis of milk TAG (4). GPAT1 contributes only 10% of total GPAT activity in every tissue in which it has been reported, except for liver where it provides 30–50% of total GPAT activity. Thus it is not surprising that the major phenotype observed in Gpat1−/− mice involves liver. Because AMPK reciprocally regulates GPAT1 and CPT1 (4), we expected that mice lacking GPAT1 would not be able to counterbalance CPT1 and would therefore increase their oxidation of fatty acids. This expectation was validated in mice fed a diet that was high in fat and sucrose. In Gpat1−/− mice plasma β-hydroxybutyrate and acyl-carnitines (suggesting incomplete fatty acid oxidation) were twice as high as in controls, and hepatic mitochondrial HMG-CoA synthase mRNA was upregulated. In addition, hepatic acyl-CoA content increased threefold in Gpat1−/− fed a low-fat diet, demonstrating the importance of GPAT1 in metabolizing acyl-CoAs, even under conditions in which dietary fat is restricted.

In vitro studies have shown that GPAT1 prefers to esterify saturated fatty acyl-CoAs, particularly palmitoyl-CoA, at the sn-1 position of glycerol-3-phosphate (4). The Gpat1−/− mouse confirmed that the enzyme functions similarly in vivo. In Gpat1−/− mice, less palmitate is present in hepatic TAG, phosphatidylcholine (PC), and phosphatidylethanolamine (PE). The PE and PC contain ~21% less palmitate and 30 and 64% more stearate and oleate, respectively, in the sn-1 position. Surprisingly, PE and PC contain ~40% more arachidonate at the sn-2 position, suggesting that fatty acid species at the sn-1 position control the fatty acid species that are esterified at the sn-2 position. Arachidonate is also 21 and 67% higher in mitochondrial PE and PC, respectively, from Gpat1−/− liver than wild-type controls, and 4-hydroxyxenonenal, a product of arachidonate peroxidation, is also increased (8).

These increases in arachidonate content and in fatty acid oxidation in Gpat1−/− mice result in a 20% increase in the rate of reactive oxygen species production, a marked increase in sensitivity to the induction of the mitochondrial permeability transition, and an increase in both oxidative stress-related hepatocyte apoptosis and in bromodeoxyuridine labeling (8). Thus, in the absence of GPAT1, excess fatty acid oxidation appears to stress the liver, but the resulting increase in apoptosis is balanced by an increase in hepatocyte proliferation. Such changes in cellular viability and proliferation could alter the risk for liver carcinogenesis.

GPAT1 Mediates Hepatic Fatty Acid Content and Insulin Sensitivity

Female Gpat1−/− mice weigh less than controls and have reduced gonadal fat pad weights and lower hepatic TAG content, plasma TAG, and secretion of VLDL-TAG (4). After a high-fat diet for 3 wk, liver from Gpat1−/− mice contains markedly lower amounts of TAG and diacylglycerol (DAG) than wild-type mice. The Gpat1−/− liver is protected from hepatic insulin resistance during a hyperinsulinemic euglycemic clamp, possibly due to reduced DAG-mediated activation of PKCe (20). Although hepatic acyl-CoA had previously been implicated in the pathogenesis of insulin resistance, compared with wild-type mice, Gpat1−/− mice exhibit increased hepatic insulin sensitivity despite an almost twofold elevation in hepatic acyl-CoA content (20). The contribution of GPAT1 to insulin resistance is further highlighted by a study showing that a 90% adenovirus-mediated shRNA knockdown of hepatic NEM-resistant GPAT1 activity in ob/ob mice reduces liver TAG and DAG content 40–50% and reduces plasma glucose and cholesterol concentrations 30–40% (24). These effects suggest that inhibiting GPAT1 might ameliorate hepatic steatosis and insulin resistance.

Because GPAT1 is upregulated by SREBP-1c, we determined what would happen to Gpat1−/− mice fed diets high in sucrose and fat (from coconut oil) or low in sucrose and fat for 4 mo (9). Although hepatic TAG content is 60% lower in Gpat1−/− mice than in control mice fed the high-fat and -sucrose diet, hyperinsulinemia and glucose intolerance are greater in the Gpat1−/− mice, and expression of the gluconeogenic genes glucose-6-phosphatase and PEPCK is not suppressed. These data are opposite those observed with 3 wk of high-fat (safflower oil) feeding (20), possibly because of the length of time on the diets or differences in the fatty acid species fed.

The short-term feeding studies and data from GPAT1 overexpression in hepatocytes suggested that GPAT1 overexpres-
sion would result in a fatty liver. Hepatic steatosis is commonly associated with insulin resistance, but the proximate cause of insulin resistance is not well understood. It has been variously suggested that lipid intermediates or inflammation due to activated NF-κB may play a role. When an adenovirus construct is used to overexpress GPAT1 in mouse or rat liver, hepatic fatty acid oxidation diminishes, the livers become fatty, and plasma becomes hyperlipidemic (Ref. 14; Nagle CA, An J, Shiota M, Torres TP, Cline GW, Liu ZX, Wang S, Catlin RL, Shulman GI, Newgard CB, Coleman RA, unpublished observations). In mice treated with adenovirus-GPAT1, hepatic TAG increases 12-fold, and adipocyte differentiation-related protein and stearoyl-CoA desaturase-1 expression are induced (14). Although body and fat pad weights are similar in rats that overexpress GPAT1 and control rats that express green fluorescent protein, a hyperinsulinemic-euglycemic clamp study shows that hepatic insulin resistance develops, with a glucose output that is 2.5-fold higher than present in control mice (Nagle CA, An J, Shiota M, Torres TP, Cline GW, Liu ZX, Wang S, Catlin RL, Shulman GI, Newgard CB, Coleman RA, unpublished observations). NF-κB and its target genes, IL-1β, and TNF-α are not affected, however, indicating that inflammation does not play a role in this model of hepatic insulin resistance. However, hepatic DAG and LPA are elevated, suggesting a role for these lipid metabolites in the development of hepatic insulin resistance. PKCε is also activated, suggesting that insulin resistance might be a response to the elevated liver DAG. In addition, glycogen synthesis and the uptake of 2-deoxyglucose are reduced in skeletal muscle, suggesting that overexpression of GPAT1 in liver causes mild peripheral insulin resistance: the higher TAG content in skeletal muscle probably occurred because of the hyperlipidemia. These results indicate that an increase in de novo hepatic TAG synthesis can cause hepatic and systemic insulin resistance in the absence of obesity or a lipogenic diet.

**GPAT1 in Tissues Other Than Liver**

Few recent studies have examined the function of GPAT1 in tissues other than liver. In rat splenic T-lymphocytes and in Jurkat cells GPAT1 activity correlates positively with lymphocyte proliferation (5). In several other tissues GPAT1 protein, mRNA expression, and specific activity are discordant (4), suggesting that GPAT1 might be regulated posttranslationally. For example, GPAT1 protein expression is high in heart and adrenal gland, but the specific activity of NEM-resistant GPAT is relatively low in these tissues.

**Questions for the Future**

In pathways of intermediary metabolism such as glycolysis, glycogen synthesis, the urea cycle, and gluconeogenesis, it is unusual to find more than one isofrom catalyzing any enzymatic step, so why do we need at least four GPAT isoforms? Is it because of the multiple glycerolipid products that are produced and the need to regulate different pathways independently? If this were true, it is odd that GPAT1 seems to be the only regulated isoform, at least in liver and adipose tissue. However, we know virtually nothing about the microsomal GPATs. Are the activities of these isoforms regulated or coordinated in any tissue? Are their LPA products destined for different fates? Does LPA synthesized at the outer mitochondrial membrane by GPAT1 enter into the same pathways as the LPA that is synthesized by GPAT-3 and -4 on the endoplasmic reticulum? Questions of different fates would imply that protein-protein interactions exist to channel lipid intermediates toward or away from particular pathways.

Another major question is whether GPAT isoforms alter the production of lipid intermediates that participate in signaling pathways. For many years it has been assumed that the lipid signals LPA, PA, and DAG arise only from phospholipase-mediated hydrolysis of membrane phospholipids. Now studies of deficient or overexpressed GPAT1 are forcing us to rethink lipid signaling. For example, GPAT1 deficiency decreases hepatic DAG content and PKCε activation (20) whereas overexpression does the opposite (Nagle CA, An J, Shiota M, Torres TP, Cline GW, Liu ZX, Wang S, Catlin RL, Shulman GI, Newgard CB, Coleman RA, unpublished observations). These effects suggest that the DAG formed during glycerolipid synthesis is able to interact with and activate PKC at distant membranes. Furthermore, LPA, a ligand for PPARγ (16), is also diminished in liver from GPAT1−/− mice and increases when GPAT1 is overexpressed. Could physiological changes in GPAT1 activity alter the effects of PPARγ on its target genes? Does GPAT1 (or the other GPAT isoforms) alter substrates such as acyl-CoA and glycerol-3-phosphate and/or products such as LPA, PA, and DAG, all of which are signaling molecules, and have effects on intracellular signaling or transcription? These complex questions demand further concerted study of the four GPAT isoforms.

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**REFERENCES**


