

MINIREVIEW

Physiological Consequences of Compartmentalized Acyl-CoA Metabolism*

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Daniel E. Cooper[‡], Pamela A. Young[‡], Eric L. Klett^{‡§},
and Rosalind A. Coleman^{‡1}

From the Departments of [‡]Nutrition and [§]Medicine, University of North Carolina, Chapel Hill, North Carolina 27599

Meeting the complex physiological demands of mammalian life requires strict control of the metabolism of long-chain fatty acyl-CoAs because of the multiplicity of their cellular functions. Acyl-CoAs are substrates for energy production; stored within lipid droplets as triacylglycerol, cholesterol esters, and retinol esters; esterified to form membrane phospholipids; or used to activate transcriptional and signaling pathways. Indirect evidence suggests that acyl-CoAs do not wander freely within cells, but instead, are channeled into specific pathways. In this review, we will discuss the evidence for acyl-CoA compartmentalization, highlight the key modes of acyl-CoA regulation, and diagram potential mechanisms for controlling acyl-CoA partitioning.

Two prevailing views of metabolic pathways within the cytosol are a) as sequential steps within a largely empty space with substrates and products traveling from one enzyme to the next, and b), as embedded within a dense network of proteins and metabolites, all jockeying for position. In contrast to these two views, it is likely that synthetic and degradative pathways are composed of enzymes and their regulators in highly organized multi-enzyme assemblies designed to enhance efficiency and regulate steady-state flux (1). Proteins within these assemblies might interact via their transmembrane and extra-membrane domains or be tethered to scaffolds, in a manner that is regulated by allosteric effectors and post-translational modifications (2, 3). Such interactions may be transient, as occurs with the enzymes that comprise purinosomes (4), or they may be semi-permanent as occurs within glycogen granules that contain enzymes that synthesize and degrade glycogen, and their regulatory kinases and phosphatases (5, 6). Assemblies of proteins are advantageous because even without a physical tunnel, they can increase reaction rates by enhancing local substrate concentrations, restricting intermediates from entering competing reactions, and stabilizing chemically unstable intermediates.

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¹ To whom correspondence should be addressed: 2301 Michael Hooker Research Center, Dauer Dr., Dept. of Nutrition, University of North Carolina, Chapel Hill, NC 27599, rcoleman@unc.edu.

Because the dysregulation of metabolic homeostasis has been implicated in a multitude of human diseases, acyl-CoA metabolism represents a critical node for understanding whole-body pathophysiology. Apart from eicosanoid synthesis, the first step in the metabolism of long-chain fatty acids (FAs)² is their thioesterification. The resulting acyl-CoA is then metabolized by one of six major enzyme families: elongases and desaturases (7), dehydrogenases (8, 9), acyl-CoA thioesterases (10, 11), carnitine palmitoyltransferases (CPT) (12), and lipid and protein acyltransferases (13) (Fig. 1). These competing pathways, often present within a single subcellular compartment and coupled with the highly specialized metabolism of individual cells and tissues, suggest a level of organization extending beyond enzymatic function. In this review, we will use a physiological lens to focus on long-chain mammalian fatty acyl-CoAs and the evidence for compartmentalized acyl-CoA metabolism.

Metabolism of Fatty Acids and Acyl-CoAs

Many pathways rely on enzymes that contain membrane-spanning domains or are tightly associated with membranes. Plasma membrane, endoplasmic reticulum (ER), and Golgi are sites of protein acylation (14), and the ER provides the site for FA ω -oxidation, for the elongation and desaturation of FAs, for the synthesis of retinol esters, cholesterol esters, and most glycerol- and sphingophospholipids, and for the synthesis of triacylglycerol (TAG). The initial substrate for each of these pathways is a long-chain acyl-CoA, the product of one of 13 long-chain or very-long-chain acyl-CoA synthetases (ACSL, ACSVL/FATP, ACS bubblegum (ACSBg)) that activate the FAs of 14–26 carbons (15). The metabolism of acyl-CoAs within cells appears to be highly controlled, and the movement of the acyl-CoAs themselves seems to be compartmentalized. How this occurs remains obscure.

Acyl-CoAs are amphipathic molecules with critical micellar concentrations of 5–42 μM , depending on chain length and degree of unsaturation (16, 17). However, acyl-CoAs are unlikely to self-aggregate because, within cells, they are bound to proteins and membranes. Because the effective free concentration of acyl-CoAs within the cell is estimated to be substantially lower than 200 nM, and because most proteins that have been found to interact with acyl-CoAs in *in vitro* experiments have specific binding affinities in the μM range, it has been suggested that acyl-CoAs may not be effective modulators of these proteins (18). However, the intracellular concentration of acyl-CoAs, which is measured as between 5 and 160 μM (18), is probably an inaccurate representation of the concentrations

² The abbreviations used are: FA, fatty acid; FATP, FA transport protein; FABP, FA-binding protein; ACS, acyl-CoA synthetase; ACSL, ACS long-chain; ACSVL, ACS very-long-chain; ATGL, adipose triglyceride lipase; CPT, carnitine palmitoyltransferase; DGAT, diacylglycerol acyltransferase; ER, endoplasmic reticulum; GPAT, glycerol-3-phosphate acyltransferase; TAG, triacylglycerol; SREBP, sterol regulatory element-binding protein; PPAR, peroxisome proliferator-activated receptor; PC, phosphatidylcholine; ACAT, acyl-CoA acyltransferase.

Acyl-CoA Metabolic Reactions

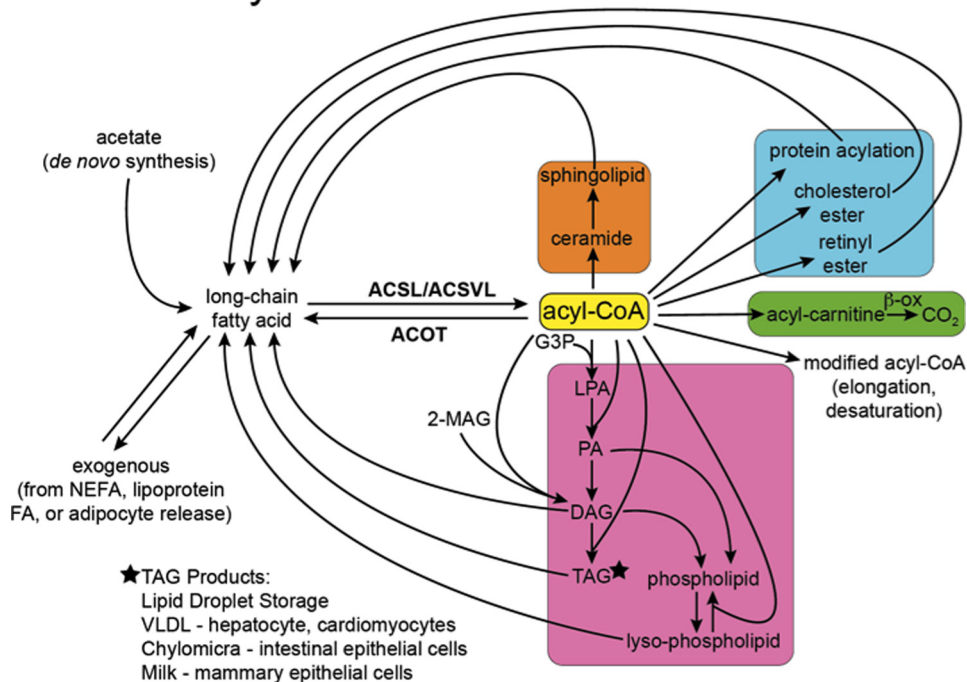


FIGURE 1. **Metabolic reactions of acyl-CoAs.** Long-chain FAs are synthesized *de novo* from acetate or enter cells from the plasma. They are converted to acyl-CoAs by ACSL and ACSVL. The reaction is reversed by acyl-CoA thioesterases (ACOT). Acyl-CoAs can be elongated and desaturated, converted to acyl-carnitines, and metabolized to CO₂ via mitochondrial and peroxisomal enzymes, esterified to glycerol-3-phosphate to form lysophosphatidic acid (LPA), phosphatidic acid (PA), and TAG, and esterified to monoacylglycerol (MAG) to form diacylglycerol (DAG). Both phosphatidic acid and diacylglycerol are precursors for all the glycerophospholipids. Acyl-CoAs are also esterified to retinol and cholesterol, acylated to proteins, and incorporated into ceramide to form sphingolipids. Lipolysis of these products releases fatty acids back into cellular pools. Triacylglycerol, cholesterol esters, and retinol esters are stored in lipid droplets within cells or secreted from specialized cells as lipoproteins or milk constituents. NEFA, non-esterified fatty acid.

within specific membrane domains or near the active sites of metabolic enzymes.

Is Acyl-CoA Metabolism Compartmentalized?

Several independent lines of data support the hypothesis that FAs and acyl-CoAs are compartmentalized within cells and trafficked into specific pathways. Although each is indirect, taken as a whole, the concept is compelling.

Vectorial Acylation

Exogenous FA uptake into cells is enhanced by vectorial acylation, a process that traps entering FAs by converting them into acyl-CoAs, which cannot exit the cell (19, 20), and by downstream metabolic pathways that release the CoA and allow it to be recycled for additional acyl-CoA synthesis (21). The cellular locations of the acyl-CoA synthetases are not critical for FA uptake. Thus, ACSLs and ACSs designated as FA transport proteins (FATPs) enhance FA uptake even when they are located on intracellular membranes (22). An example is provided by similar uptake of arachidonate by each of two differentially spliced variants of ACSL4, one located on the plasma membrane and in the cytosol and the other located on the ER and lipid droplets (23). Overexpressing ACSL4 at the ER generated 50% more phosphatidylinositol than when it was targeted to the mitochondria, compatible with a specific interaction with ER enzymes that synthesize or remodel phosphatidylinositol. These data provide compelling evidence for compartmentalization, as subcellular location of a specific ACSL

isoform may be important for the interaction of FA substrates with a specific metabolic pathway.

Many downstream proteins that metabolize acyl-CoAs have a well defined cellular distribution. CPT1, for example, is expressed on the outer mitochondrial membrane as part of the machinery to transport activated long-chain acyl-CoAs into the mitochondrial matrix for β -oxidation (24), and the major esterification pathways that use acyl-CoAs are predominantly located on the ER. However, the location of several putative intrinsic membrane enzymes may not be static. For example, incubating *Drosophila* S2 cells with oleate causes the ER glycerol-3-phosphate acyltransferase (GPAT) to translocate to newly formed lipid droplets (25), suggesting that GPAT domains formerly believed to constitute transmembrane sequences, can instead integrate within the phospholipid monolayer that surrounds the lipid droplet. With few exceptions, FA channeling has not been rigorously evaluated. ACSL1 and CPT1 interact at the mitochondrial outer membrane, an association that ensures compartmentalized FA channeling into the mitochondrial matrix (26); the interaction of diacylglycerol acyltransferase (DGAT)-2 and monoacylglycerol acyltransferase-2 within a large protein complex via the transmembrane domains of DGAT2 should enable cells to efficiently process acyl-CoAs within the cytosol (27); and recent co-immunoprecipitation studies have identified proteins in the pathway of TAG synthesis that interact with the ER protein seipin (28).

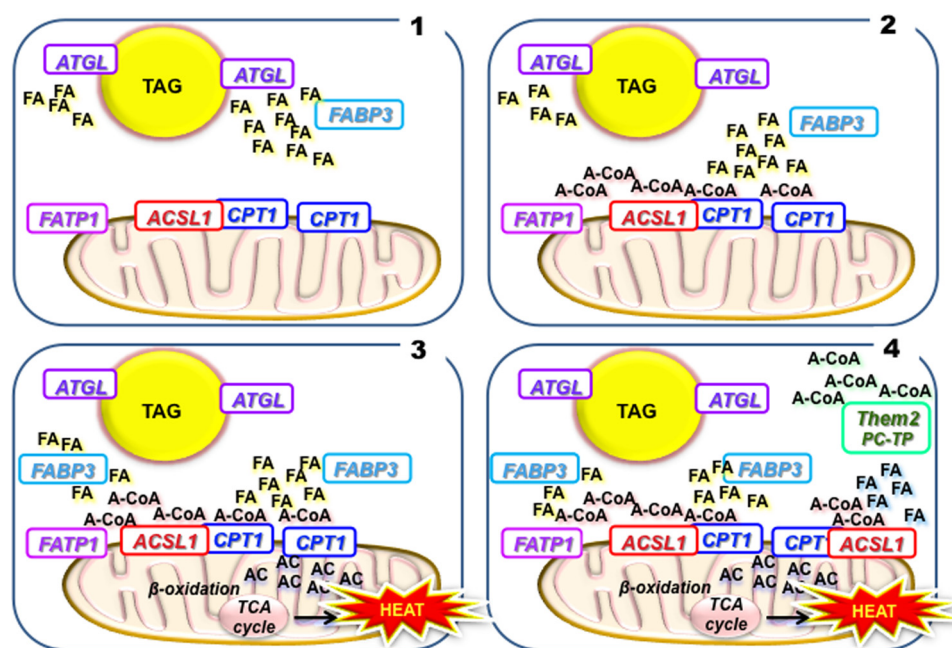


FIGURE 2. **A hypothetical assembly of proteins, substrates, and products that are critical for thermogenesis in brown adipocytes.** Mice are unable to maintain a normal body temperature if they lack ATGL, FABP3, FATP1, and ACSL1 or are heterozygotes for CPT1. **1)** ATGL hydrolyzes FA from TAG in lipid droplets. **2)** FABP3 binds these released FAs and transports them to FATP1 and ACSL1, which convert them to acyl-CoAs (A-CoA). **3)** These acyl-CoAs are substrates for CPT1, which converts them to acylcarnitines (AC) that enter the mitochondrial matrix and are metabolized by β -oxidation to produce heat. **4)** Thioesterase superfamily member-2 (*Them2*) and its partner phosphatidylcholine transfer protein (*PC-TP*) improve the thermogenic process by converting acyl-CoAs, perhaps those formed in other locations, back to fatty acids that can be reactivated by FATP1 and ACSL1 located at the mitochondrial membrane. TCA, tricarboxylic acid cycle.

Evidence from Thermogenesis

FA oxidation in brown adipose is impaired by deficiencies in several proteins and enzymes that provide a view of what a multi-protein metabolic cluster might encompass. In addition to the mitochondrial enzymes of β -oxidation, thermogenesis requires proteins involved in TAG lipolysis and FA transport and activation (Fig. 2). These proteins include adipose triglyceride lipase (*ATGL*), a lipid droplet-associated lipase that releases FAs (29), palmitoyl-protein thioesterase-1, which diminishes TAG lipolysis (30), FA-binding protein (FABP)-3, a FA carrier (31, 32), and ACSL1 (33) and FATP1/ACSLV4 (34), which are FA activators. The block in thermogenesis caused by the absence of any one of these proteins strongly suggests that they form an assembly in brown adipose that metabolizes a pool of FAs that originate in lipid droplets and are specifically targeted to CPT1 for β -oxidation. In addition, the thioesterase, *Them2*, and its binding partner phosphatidylcholine transfer protein, enhance thermogenesis, suggesting that these, too, are members of the thermogenic cluster (35).

Metabolism of de Novo Synthesized FA

Studies of mice deficient in acyl-CoA synthetases suggest that acyl-CoAs may be sequentially synthesized, hydrolyzed, and resynthesized, perhaps as they are targeted to different organelles within cells. For example, we can construct a sequence beginning with the insulin-induced *de novo* synthesis of 16:0 within the cytosol, followed by conversion to 16:0-CoA by an ACSL and then esterification to form lysophosphatidic acid by GPAT1, which is embedded in the outer mitochondrial membrane and prefers saturated FAs that have been synthe-

sized *de novo* (36). In GPAT1-deficient mice, FA oxidation increases, suggesting that GPAT1 and CPT1 compete at the mitochondrial membrane for a specific pool of acyl-CoAs (37). Regulation occurs via SREBP1- and carbohydrate-responsive element-binding protein (ChREBP)-mediated up-regulation of enzymes of FA synthesis and GPAT1 (13), so that when acyl-CoAs are the product of *de novo* biosynthesis, their capture by GPAT1 prevents newly synthesized FAs from subsequent β -oxidation.

In contrast to this evidence for intracellular FA compartmentalization, dietary triolein supplementation restores lipid accumulation in stearoyl-CoA desaturase 1 (*SCD1*)-deficient liver (38). However, because dietary triolein does not correct the phenotype in global *SCD1* deficiency, exogenous and endogenous oleate pools may be metabolized differently in extrahepatic tissues.

ACSL1 Knock-out Mice

Studies of ACSL1 knock-out mice provide the most persuasive argument for compartmentalized acyl-CoA metabolism. In liver, ACSL1 is equally distributed on the ER and the mitochondrial outer membrane. Because it comprises only ~50% of the total ACS activity in either location, its absence results in minor decreases in TAG synthesis and FA oxidation (39). In contrast, in highly oxidative tissues, including heart, skeletal muscle, and brown adipose, ACSL1 constitutes 90% of total ACS activity and its absence severely diminishes FA oxidation, although the remaining ACSL isoforms are sufficient for normal TAG and membrane phospholipid biosynthesis (33, 40, 41). One might reason that when ACSL1 is absent, FA oxidation would be neg-

ligible, merely because less exogenous FA enters the cells. Arguing against this interpretation, however, are data from mice deficient in skeletal muscle ACSL1 (41). Because FAs cannot be used for fuel, these mice can run only half as far as their wild type littermates before becoming hypoglycemic. Surprisingly however, their muscle content of long-chain acyl-CoA increases 10-fold during exercise and at exhaustion is 89% higher than controls. The low muscle content of long-chain acylcarnitines confirms that the large amount of acyl-CoA synthesized by other ACSL isoforms in ACSL1-deficient muscle is metabolically unavailable for β -oxidation.

Similar to skeletal muscle, ACSL1 is the predominant ACS in heart. Genetic deletion of ACSL1 results in a 97% decrease in total long-chain acyl-CoAs (40). This decrease results in a nearly complete inhibition of mitochondrial FA oxidation, although membrane phospholipid and TAG synthesis remain intact. Again, it appears that mitochondrial ACSL1 is essential in the heart to partition FA toward oxidation. ACSL1 functions similarly in adipose tissue (33). When mice are challenged with cold exposure, brown adipose FAs cannot be mobilized from the TAG stored in lipid droplets and are unavailable for mitochondrial oxidation and thermogenesis, and even in white adipose, the reduced β -oxidation results in preferential partitioning of FA into TAG stores (33).

ACSL Subcellular Location

Intracellular FAs originate from three primary sources, exogenous FAs that enter cells from the blood or from the gut lumen, FAs that arise via *de novo* synthesis from acetate, and FAs that are released within the cell by the hydrolysis of acylated proteins, phospholipids, and TAG. Indirect data suggest that FAs originating from each source comprise distinct intracellular pools and are preferentially activated by specific acyl-CoA synthetases and directed into specific metabolic pathways. For example, FAs derived from *de novo* synthesis versus exogenous sources are metabolized differently by GPAT isoforms located on the mitochondria or the ER (36) and even by the two DGAT isoforms, located on the ER or on lipid droplets (42–44).

Although subcellular location ought to critically define aspects of acyl-CoA partitioning, studies of the ACSL isoforms have been problematic. Subcellular fractionation frequently results in contamination of specific organelles with other cellular constituents. For example, several ACSL isoforms have been identified by proteomic studies of lipid droplets, but it remains unclear as to whether discrepancies in these reports reflect stable or transient associations of a particular ACSL on the lipid droplet or, instead, contamination with fragments of associated ER (45). Confocal studies are superior in that the normal cell structure is not disrupted, but they depend on highly specific antibodies to native proteins, and may not identify proteins expressed at low amounts in some organelles. Overexpression studies, particularly with tagged proteins, may not reflect the endogenous location or interactions with other proteins. The reported interaction of FATP1 and ACSL1 (46), for example, should be re-examined using a confocal method. Massively overexpressing a protein may result in ER retention and may not mirror the location of the native protein.

In addition to technical limitations in assigning location, the individual ACSL isoforms appear to move in response to changes in physiologic stimuli. Similar to studies of GPAT4 (25), treatment with exogenous FA causes tagged ACSL3 to move from the ER to newly forming lipid droplets (47), and a knockdown of *Acs13* in hepatocytes reduces *de novo* FA synthesis (48, 49), suggesting that ACSL3 is involved in the flux of acyl-CoAs toward storage. Treatment with insulin stimulates FATP1 translocation from the ER to the plasma membrane and increases the intracellular content of long-chain FA.

In addition to physiologic changes, the location of ACSL isoforms may be tissue-dependent, but few studies have reported on both location and function. Thus, the presence of ACSL5 on liver mitochondria (50) suggested that, like ACSL1, ACSL5 would target FA toward β -oxidation. However, although overexpressed ACSL5 in McArdle-RH7777 cells was located on both mitochondria and ER, oleate incorporation increased only into TAG and not into oxidation products (51). In support of these data, the siRNA knockdown of ACSL5 in rat primary hepatocytes decreased oleate-mediated TAG synthesis and lipid droplet formation (52).

Nutritional Regulation

To maintain energy homeostasis, cells must adapt rapidly to the available nutrients. Switching between glucose and FAs (53) is regulated by the cellular level of malonyl-CoA, which inhibits CPT-I, thereby blocking the mitochondrial entry and oxidation of long-chain fatty acyl-CoAs, which are then available for esterification into glycerolipids. During fasting, rising cellular AMP levels activate AMP-activated protein kinase, which enhances ATP-generating pathways that diminish malonyl-CoA production, thereby relieving the inhibition of CPT-I. In addition, fasting induces TAG lipolysis, and the released FAs activate the deacetylase, SIRT1, and the PPAR co-activator, PGC1 α , to increase mitochondrial biogenesis (54). Thus, cytosolic metabolites control the metabolic fate of the acyl-CoAs.

Nutrient status similarly controls the transcription of ACSL isoforms. In rat liver, fasting increases and refeeding decreases the mRNA abundance of *Acs11* and *Acs14*, whereas the expression of *Acs15* mRNA responds in the opposite manner (55). However, even these changes can be altered by dietary nutrients, so that fasted rats that are refed with a high-fat (20% soybean oil) diet increase *Acs11* mRNA expression in liver (56).

Transcriptional regulation of ACSL isoform mRNA expression depends on activated PPARs. In heart, liver, and skeletal muscle, PPAR α activation increases the transcription of *Acs11* (57, 58), but in adipocytes, the transcription of *Acs11* is enhanced by PPAR γ (59), *Acs14* transcription in liver is also increased by PPAR β/δ activation (60). Although synthetic ligands were used to activate the PPAR isoforms, these studies suggest that, as PPAR ligands, endogenous and exogenous FAs or acyl-CoAs may themselves alter ACSL isoform expression and activity.

Although *in vitro* experiments must be interpreted cautiously, FAs appear to control ACSL expression, perhaps via PPAR activation. Exogenous oleate added to rat cardiomyocytes increases *Acs11* expression (57), and in rat insulinoma cells or mouse hepatocytes, polyunsaturated FAs, but not satu-

rated or monounsaturated FAs, reduce both *Acs14* mRNA and protein expression (61, 62). Polyunsaturated FAs also increase ACSL4 protein degradation via a ubiquitination-dependent, proteasomal degradative pathway (62). In addition to promoting changes in subcellular location, insulin increases *Acs11* and *Acs15* expression in cultured hepatoma cells (63), *Acs11* and *Acs13* expression in rat cardiomyocytes, and *Acs16* expression in cardiomyocytes (57). It is not known whether the inductive effect of insulin on the ACSL isoforms is mediated by SREBP. These insulin exposure studies appear to be at odds with fasting studies in which one would expect that insulin levels would be relatively low.

Polyunsaturated FAs are natural ligands for nuclear transcription factors such as the PPARs, liver X receptors (LXRs), HNF-4 α , and SREBPs 1 and 2, but their chain length, degrees of unsaturation, and source (exogenous *versus de novo*) have differential effects on the gene regulation. For example, hepatic PPAR α binds oleate, arachidonate, and docosahexaenoic acid with near equal affinity (64); however, in rat hepatocytes, only docosahexaenoic acid activates PPAR α (65). These observations suggest that the metabolism of specific FA species is highly organized, supporting the hypothesis that acyl-CoAs are compartmentalized.

ACSL Substrate Preference

The Yamamoto group (66), which cloned the five rat ACSL isoforms, used a coupled spectrophotometric assay to identify the FA preferences of purified recombinant rat isoforms. Although all isoforms activated saturated and unsaturated FAs of 16–20 carbons, ACSL6 (originally called ACSL2) had higher activities with 20:4 ω 6, 20:5 ω 3, and 22:6 ω 3, suggesting that it might be involved in the metabolism of brain phospholipids, which have a high content of polyunsaturated FAs, or in eicosanoid function (67, 68). In fact, in differentiating PC12 cells, ACSL6 enhances both the uptake of 22:6 and neurite outgrowth (69, 70). ACSL4 has a reported preference for arachidonate, and as might be expected, the incorporation and the secretion of eicosanoids and arachidonate are diminished when ACSL4 is knocked down in INS832/13 (61), 3Y1 rat fibroblast (71), and activated hepatic stellate cells (HSC) (72). However, the published substrate preference for each ACSL family member is not invariably observed. For example, although the purified recombinant ACSL1 had no preference for arachidonate or linoleate, ACSL1 is critical for the synthesis of arachidonoyl-CoA in macrophages (73) and was inferred to prefer linoleate in the heart (74). Studies performed in different cell lines also reveal disparate effects of ACSL4 deficiency on arachidonic acid metabolism for phospholipid synthesis and prostaglandin secretion. In 3Y1 and HSC cells, siRNA-mediated knockdown of ACSL4 reduces the incorporation of arachidonic acid into phosphatidylcholine and dramatically decreases the secretion of prostaglandins, but in INS cells, the deficiency of ACSL4 has no effect on the incorporation of arachidonic acid into PC. Instead, the knockdown of ACSL4 in INS cells results in increased unesterified epoxyeicosatrienoic acids because of diminished activation and esterification into glycerolipids.

Most members of the closely related FATP family (also called ACSVL for ACS very-long-chain) can use FAs of 16–24 car-

bons (75). Because each tissue expresses several ACSL and FATP/ACSVL isoforms, the location of the isoform might direct a particular chain length FA into a specific pathway. Although not experimentally demonstrated, products synthesized by the same acyltransferase, but that differ in FA composition, suggest that a specific ACSL isoform must provide the esterifying enzyme with larger amounts of specific acyl-CoA substrates. Such products include the overwhelming predominance of cholesteryl oleate in liver and LDL (76) and the predominance of cholesteryl adrenate (22:4 ω 6) and cholesteryl arachidonate (20:4 ω 6) in rat adrenal cortex (77). Because the substrate preference of ACAT1, which is present in rat adrenal cortex (76), is oleate, whereas ACAT2 esterifies 18:1, 20:5, and 22:5 equally (78), the composition of tissue cholesteryl esters seems more aligned with the ACSL isoforms than the ACAT isoforms present in each tissue. Similarly, 77% of cardiolipin species in heart and skeletal muscle, in which ACSL1 contributes more than 90% of total ACSL activity, are tetra-18:2 (79), whereas tetralinoleoyl-cardiolipin is ~50% of the cardiolipin species in liver (69), a tissue in which ACSL1 is responsible for only about 50% of the total ACSL activity (39). In brain, where ACSL1 is minimally expressed (55), the major cardiolipin acyl chains are oleate and arachidonate (80).

PC biosynthesis has also been linked to ACSL3 activity; in human hepatoma Huh7 cells, siRNA-mediated knockdown of *Acs13* diminishes oleate incorporation into PC and reduces VLDL secretion (81). Thus, ACSL3 may activate FAs destined for the synthesis of the phosphatidylcholine that is essential for VLDL secretion (81). Taken together, these findings demonstrate that, depending upon cell type, different ACSL isoforms activate specific FAs that can be incorporated into a variety of glycerolipids. Furthermore, although each ACSL isoform may have a specific substrate preference, the function of each ACSL isoform seems to vary in different cell types.

ACSL Isoforms and Signaling Acyl-CoAs

Lipolysis-derived FAs and acyl-CoAs have been implicated as essential intracellular signaling metabolites (82). As signaling molecules, acyl-CoAs link the nutritional environment to transcription factors capable of reorganizing cellular metabolism to ensure an adequate response to changes in nutrient availability and to maximize metabolic efficiency (83). For example, ACSL3 appears to promote *de novo* lipogenesis and storage of neutral lipid because the siRNA knockdown of *Acs13* in primary rat hepatocytes also diminishes the expression of PPAR γ , carbohydrate-responsive element-binding protein, SREBP1c, and liver X receptor α as well as the expression of the target genes of these transcription factors (48).

Specific pools of FAs or acyl-CoAs are critical in the heart. In hearts deficient in ATGL, the impaired release of FAs from stored TAG results in low expression of PPAR α target genes, inadequate FA oxidation, TAG accumulation, and heart failure (84–87). Because treating mice with fibrates increased FA oxidation, reduced TAG stores, and improved heart function, it appeared that the ATGL deficiency had reduced the availability of FAs and/or acyl-CoA ligands required to activate PPAR α ; thus, despite high levels of acyl-CoAs that partitioned into neutral lipids, exogenously derived FAs were not PPAR α ligands

(86). A competing interpretation suggests that the effect of ATGL-derived FAs on PPAR α is mediated indirectly by SIRT1 (54), but the data from both studies are consistent with transcriptional signaling by distinct pools of FA and acyl-CoAs originating from a specific endogenous source.

Mechanisms, Possibilities, and Unanswered Questions

The wide range of functions initiated by each ACSL isoform suggests three major problems. First, the different effects observed in cell lines of similar origin suggest that analyzing function in the relevant primary cells would provide the most reliable and conclusive information. Second, in addition to mislocalizing proteins, overexpression studies can overwhelm downstream metabolic pathways, resulting in a misleading interpretation of function (88). For example, when ACSL1 is overexpressed in primary hepatocytes, the protein localizes to the ER and increases oleate re-esterification into diacylglycerol and phospholipids (89). In contrast, the liver-specific knockout of ACSL1 disrupts TAG synthesis and oxidation only mildly (39, 89). Thirdly, the disparate physiological effects observed in different tissues when ACSL1 is absent suggest that protein-protein interactions between ACSL isoforms and downstream metabolic enzymes vary considerably in tissues that have specialized physiological functions. Thus, differing strongly from the FA storage and degradation pathways affected in ACSL1-deficient liver, adipose, skeletal muscle, or heart, ACSL1 in macrophages predominantly mediates inflammatory effects and channels FAs toward the synthesis of phospholipid (90, 91), whereas ACSL1 in endothelial cells is unrelated to inflammatory changes associated with dietary fat (92). Clearly, a better understanding of the locations of each of the ACSLs and their downstream partners is needed.

Important questions remain concerning the trafficking of FAs and acyl-CoAs. We do not know how exogenous FAs are transported from the cytosolic face of the plasma membrane to ACSL isoforms that are located on intracellular membranes. Interaction with the intracellular acyl-CoA-binding proteins or FABP might direct FAs and acyl-CoAs to specific sites within cells. Although directional transport by these binding proteins has been inferred, it has not been demonstrated (93). For example, the absence of FABP4 diminishes PPAR activation by FA (94), and FABP4 enhances lipase activity by interacting with hormone-sensitive lipase (HSL) to bind released FAs (95). Although the absence of FABP and acyl-CoA-binding proteins alters lipid metabolism, the function of these carrier proteins may be more analogous to that of albumin in allowing FAs and acyl-CoAs to be used by enzymes or to bind to transcription factors in a controlled manner (96).

Although indirect data imply the existence of organized assemblies of enzymes that metabolize acyl-CoAs, definitive proof is absent. Newer methods, both physical and *in silico*, must be sought to identify protein partners and interactions without the potential hazards of overexpression (97–99). We also lack any understanding of why the up-regulation of β -oxidation not only increases the expression of *Acs11*, but, counterproductively, also increases the expression of thioesterases such as Them2 and its partners (35). It is unclear as to how protein assemblies form in response to specific nutrient or

energy signals. If proteins associate with each other or with scaffolding proteins through the interactions of specific domains, what then causes them to disperse? Phosphorylation and acetylation of ACSL1 are modified hormonally on numerous sites (100), but the functions of these post-translational modifications are unknown. The answers to these questions demand new tools to identify the movements of lipids within cells and new methods to capture transient protein assemblies and to manipulate them.

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