LYCAT, a homologue of *C. elegans acl-8, acl-9, and acl-10*, determines the fatty acid composition of phosphatidylinositol in mice

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Abstract Mammalian phosphatidylinositol (PI) has a unique fatty acid composition in that 1-stearoyl-2-arachidonoyl species is predominant. This fatty acid composition is formed through fatty acid remodeling by sequential deacylation and reacylation. We recently identified three *Caenorhabditis elegans* acyltransferases (ACL-8, ACL-9, and ACL-10) that incorporate stearic acid into the sn-1 position of PI. Mammalian LYCAT, which is the closest homolog of ACL-8, ACL-9, and ACL-10, was originally identified as a lysocardiolipin acyltransferase by an in vitro assay and was subsequently reported to possess acyltransferase activity toward various anionic lysophospholipids. However, the in vivo role of mammalian LYCAT in phospholipid fatty acid metabolism has not been well elucidated. In this study, we generated LYCAT-deficient mice and demonstrated that LYCAT determined the fatty acid composition of PI in vivo. LYCAT-deficient mice were outwardly healthy and fertile. In the mice, stearoyl-CoA acyltransferase activity toward the sn-1 position of PI was reduced, and the fatty acid composition of PI, but not those of other major phospholipids, was altered. Furthermore, expression of mouse LYCAT rescued the phenotype of *C. elegans acl-8 acl-9 acl-10* triple mutants. Our data indicate that LYCAT is a determinant of PI molecular species and its function is conserved in *C. elegans* and mammals.—Imae, R., T. Inoue, Y. Nakasaki, Y. Uchida, Y. Ohba, N. Kono, H. Nakanishi, T. Sasaki, S. Mitani, and H. Arai. LYCAT, a homologue of *C. elegans acl-8, acl-9, and acl-10*, determines the fatty acid composition of phosphatidylinositol in mice. *J. Lipid Res.* 2012, 53: 335–347.

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Phosphatidylinositol (PI) is a relatively minor component of membrane phospholipids but plays important roles in signal transduction through distinct phosphorylated derivatives of the inositol head group (1, 2). Four-fourths or more of membrane PI obtained from mammalian tissues consist of the 1-stearoyl-2-arachidonoyl (18:0/20:4) species (3, 4), which is thought to be formed by a fatty acid remodeling reaction after the de novo synthesis of PI (5–10). The remodeling reaction involves the hydrolysis of a fatty acyl ester bond at the sn-1 or sn-2 position of the newly synthesized PI and subsequent incorporation of the appropriate fatty acid into the position. In an RNA interference (RNAi)-based genetic screen using *Caenorhabditis elegans*, we identified *mboa-7/LPIAT1* as an acyltransferase that selectively incorporates arachidonic acid into the sn-2 position of PI (11). More recently, we demonstrated that *C. elegans acl-8, acl-9, and acl-10*, which show significant sequence homology to each other, encode acyltransferases that incorporate stearic acid (18:0) into the sn-1 position of PI (12). Stearic acid attached at the sn-1 position of PI

Abbreviations: AGPAT, 1-acylglycerol-3-phosphate Oacyltransferase; A-P axis, anterior-posterior axis; CL, cardiolipin; ER, endoplasmic reticulum; LPCAT, lysophosphatidylcholine acyltransferase; LPEAT, lysophosphatidylethanolamine acyltransferase; LPGAT, lysophosphatidylglycerol acyltransferase; LPIAT, lysophosphatidylinositol acyltransferase; LPSAT, lysophosphatidylsersine acyltransferase; LYCAT, lysocardiolipin acyltransferase; MEF, mouse embryonic fibroblast; PC, phosphatidylcholine; PE, phosphatidylethanolamine; PG, phosphatidylglycerol; PI, phosphatidylinositol; PIP1, phosphatidylinositol monophosphates; PIP2, phosphatidylinositol bisphosphates; PS, phosphatidylinerine; X-Yn-Z, fatty acid chain of X carbon atoms and Y methyl-ene-interrupted αx bonds (Z indicates the position of the terminal double bond relative to the methyl end of the molecules).

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5 The online version of this article (available at http://www.jlr.org) contains supplementary data in the form of one table and three figures.
was replaced with cis-vaccenic acid (18:1n-7) in *acl-8 acl-9 acl-10* triple mutants. The gene product of *acl-10*, the predominant acyltransferase among *acl-8*, *acl-9*, and *acl-10*, incorporates stearic acid into the sn-1 position of PI in vitro. *acl-8 acl-9 acl-10* triple mutants were defective in the asymmetric cell division of epithelial cells. We also showed that *acl-8*, *acl-9*, and *acl-10* function in the same pathway with *ipla-1* (13), a phospholipase A₁ that hydrolyzes the fatty acyl chain of PI (12). *C. elegans ipla-1* mutants have fatty acid compositions of PI similar to *acl-8 acl-9 acl-10* triple mutants. *ipla-1* mutants show epithelial cell defects similar to *acl-8 acl-9 acl-10* triple mutants. No synergism was observed between the *ipla-1* and *acl-8 acl-9 acl-10* mutations. These data support a model in which IPLA-1, the gene product of *ipla-1*, produces the sn-2 acyl lysoPI, which is subsequently reacylated by ACL-8, ACL-9, and ACL-10, gene products of *acl-8*, *acl-9*, and *acl-10*, respectively, in the fatty acid remodeling of the sn-1 position of PI.

ACL-8, ACL-9, and ACL-10 belong to the 1-acylglycerol-3-phosphate O-acyltransferase (AGPAT) family (14) (supplementary Table I), in which the members share four conserved AGPAT motifs that are involved in substrate binding and catalysis (15, 16) (supplementary Fig. I, motif I-IV). In mammals, the AGPAT family consists of at least 16 members. Among these members, lysocardiolipin acyltransferase (LYCAT), also known as LCLAT1 and ALCAT1, is the closest homolog of *C. elegans* ACL-8, ACL-9, and ACL-10 (supplementary Table I) (14). ACL-8, ACL-9, and ACL-10 in mammals possess highly conserved amino acids in the AGPAT motifs, which are unique in *LYCAT*/ACL-8,-9,-10 subfamily members, but not in other AGPAT family members (supplementary Fig. 1B, amino acids indicated in blue). Aciyltransferases with these highly conserved amino acids are evolutionarily conserved in various species including human, zebrafish, and *C. elegans*, but not in yeast.

Mammalian LYCAT was originally identified as a lysocardiolipin (lysoCL) acyltransferase by an in vitro enzyme assay (17). Thereafter, it has been shown that mammalian LYCAT possesses acyltransferase activity toward the sn-2 position of other anionic lysophospholipids including...
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Materials and Methods

Materials

PI and lysoPI from bovine liver, dioleoyl phosphatidylcholine (PC), dioleoyl phosphatidylethanolamine (PE), and 1-palmitoyl-2-oleoyl phosphatidylserine (PS) were purchased from Avanti Polar Lipids (Alabaster, AL). Phosphatidylglycerol (PG) from egg yolk was purchased from Sigma-Aldrich (St. Louis, MO). 1,2-Dipalmitoyl PI was purchased from Serdary Research Laboratories (London, ON, Canada). 1,2-Dipalmitoyl phosphatidylinositol monophosphates (PIP1), 1,2-dipalmitoyl phosphatidylinositol bisphosphates (PIP2), and 1,2-dioleoyl PIP2 were purchased from Cayman Chemical (Ann Arbor, MI). [1-14C]stearoyl-CoA and [1,13C]arachidonoyl-CoA were purchased from American Radio-labeled Chemicals (St. Louis, MO). Rhizopus arrhizus lipase and phospholipase A2 from honey bee venom were purchased from Sigma-Aldrich. DEAE column was purchased from Wako Pure Chemical Industries (Osaka, Japan).

Worm strains

General methods for maintaining C. elegans are described by Brenner (21). The orientation of seam cell division and seam cell lineages were analyzed as previously described (13). The following mutations and transgenes were used: acl-8 acl-9(tm2290), acl-10(tm1045), wIs51[scm::gfp], xhEx3521[dpy-7p::mouse LYCAT; Pges-1::dsREDm] (12).

Preparation of sn-2-acyl lysophospholipids

Each sn-2-acyl-1-lysophospholipid (sn-2-acyl lysophospholipids) was prepared as follows using dioleoyl PC, dioleoyl PE, 1-palmitoyl-2-oleoyl PS, bovine liver PI, or egg yolk PG. Each phospholipid (1 pmol) dissolved in 1 ml of diethyl ether was incubated with 100 µl of 100 mM CaCl2, 600 µl of 50 mM Tris-malate, pH 5.7, and 400 µl of enzyme solution containing 15 mg of Rhizopus arrhizus lipase for 1 h at room temperature while stirring vigorously. After the incubation, the reaction was terminated by adding 1 ml of methanol. Remaining phospholipids and liberated fatty acids were removed by three extractions with 4 ml of diethyl ether-petroleum ether (1:1, v/v). sn-2-acyl lysophospholipid present in the lower layer was extracted by the method of Bligh and Dyer (22) and further purified by TLC on silica gel 60 plates (Merck Biosciences, Darmstadt, Germany) in chloroform-methanol-acetic acid (65:25:13, v/v). The area of silica gel corresponding to sn-2-acyl

Fig. 2. Generation of LYCAT-deficient (LYCAT+/−) mice. A: Schematic representation of the LYCAT gene-targeting strategy. Homologous recombination of the targeting vector with the wild-type allele results in the replacement of 0.5-kilobase of the LYCAT genomic sequence containing exon 4 with the neo cassette, deleting the AGPAT motifs II and III that are essential for acyltransferase activity in AGPAT family members (15, 16). A thymidine kinase gene (TK) was used for negative selection of clones with random integration of the targeting vector. B: PCR-based genotyping of genomic DNA from LYCAT+/+, LYCAT+/-, and LYCAT−/− mice. The orientation and location of PCR primers are indicated with arrows in (A). C: Western blot analysis using an anti-mouse LYCAT antibody, YN1. Total protein extracts of liver from LYCAT+/+ and LYCAT−/− mice were subjected to SDS-PAGE, blotted onto a PVDF membrane, and stained with YN1.
**TABLE 1. Acylation of sn-2-acetyl lysophospholipid preparations by [14C]stearoyl-CoA in mice liver**

| Acyl acceptors | sn-1 position | sn-2 position |
|----------------|---------------|---------------|
| lysoPC         | 9.87 ± 0.94   | 0.74 ± 0.01   |
| Lycat+/+       | 10.06 ± 0.78  | 0.85 ± 0.05   |
| lysoPE         | 48.42 ± 1.74  | 2.27 ± 0.03   |
| Lycat+/-       | 41.54 ± 0.09  | 2.15 ± 0.01   |
| lysoPI         | 8.10 ± 0.11   | 0.15 ± 0.00   |
| Lycat+/-       | 2.92 ± 0.22   | 0.10 ± 0.00   |
| lysoPS         | 8.02 ± 0.13   | 0.22 ± 0.04   |
| Lycat+/-       | 7.80 ± 0.27   | 0.19 ± 0.00   |
| lysoPG         | 10.45 ± 0.05  | 0.60 ± 0.07   |
| Lycat+/-       | 4.57 ± 0.17   | 0.26 ± 0.03   |

The acyltransferase reaction mixtures contained 80 µM sn-2-acetyl lysophospholipid, 12.5 µM [14C]stearoyl-CoA, and 10 µg of liver microsomal protein in a total volume of 0.8 ml assay buffer. Each 14C-labeled phospholipid produced after the incubation was treated again with phospholipase A2. The distribution of radioactivity between the resultant lysophospholipid and free fatty acid was taken as a measure of acylation at the sn-1 position (sn-2-acetyl lysophospholipid acyltransferase activity) (a) and the sn-2 position (sn-1-lacyllysophospholipid acyltransferase activity) (b), respectively. Data represent the mean ± SEM of triplicate measurements. *P < 0.01 versus Lycat+/-.

Lysophospholipid was scraped off the plates. sn-2-acetyl lysophospholipid was reextracted by methanol and immediately used for acyltransferase assays.

**Acylation assay**

Mouse tissues were pulverized under liquid nitrogen and homogenized in quadruple volumes (w/v) of SET buffer (10 mM Tris-HCl, pH 7.4, 1 mM EDTA, 250 mM sucrose) with protease inhibitors (5 µg/ml pepstatin, leupeptin, and apropin). After centrifugation at 2,500 g for 20 min at 4°C, the resulting supernatant was further centrifuged at 105,000 g for 60 min. The resulting pellet (microsomal fraction) was resuspended in homogenizing buffer (50 mM potassium phosphate buffer (pH 7.0) containing 0.15 M KCl, 0.25 M sucrose) and used for the enzyme assay. Acyl-CoA:sn-2-acetyl lysophospholipid acyltransferase assay and acyl-CoA:sn-1-lacyl lysophospholipid acyltransferase assay were performed essentially as described previously, except that the amount of microsomal protein was 10 µg and the incubation temperature was 37°C. Because sn-2-acetyl lysophospholipid is known to easily isomerize to sn-1-lacyl lysophospholipid, it is possible that the acyl donor is incorporated into sn-1-lacyl lysophospholipid and sn-2-acetyl lysophospholipid. Thus, an accurate measure of sn-2-acetyl lysophospholipid acyltransferase activity could only be obtained by determining the position that had been acylated. To check the positional specificity, the radiolabeled product was treated with bee venom phospholipase A2 (supplementary Fig. II). The distribution of radioactivity between the resultant sn-1-lacyl lysophospholipid and free fatty acid was assessed after TLC.

**Generation of Lycat-deficient mice**

Mouse Lycat genomic DNA fragments were cloned from a C57BL/6-derived genomic DNA and subcloned into the pPNT vector. The targeting vector contains a PGK-neo cassette with 2.3 kb of Lycat homologous regions upstream of exon 4 and with 5.9 kb homologous regions downstream of exon 4. The targeting vector was linearized and electroporated into RENKA ES cell lines (TransGenic Inc., Kumamoto, Japan) derived from the C57BL/6 mouse substrain. Stable clones were selected for G418 resistance. Out of 569 G418-resistant clones, three clones were correctly targeted, as confirmed homologous recombination by Southern blot analysis with 5′, 3′, and neo probes. Blastoycists obtained by aggregation methods of ICR morula embryos and three different ES cells to recipient uterus as from 3 days of pseudopregnancy. Chimeric males were mated to C57BL/6 females and offspring animals (F1) were genotyped by Southern blot analysis. F2 homozygous mutant mice were then generated by intercrossing F1 heterozygous males and females. Mice were genotyped by PCR analyses of genomic DNA isolated from tail biopsies. The following primers were used for genotyping: primer 1, 5′-GTG GTA CTT CAC ACC TTT GAT GGC TCT TAT CAT CCT TGC-3′; primer 2, 5′-CAA AAT GGC TCT TAT CAT CCT TGC-3′; and primer 3, 5′-GAG CAG AGT ACC TAC ATT TTG AAT GG-3′. The wild-type allele is detected with primers 1 and 2, yielding a 356 bp product, whereas the mutant allele is detected with primers 1 and 3, yielding a 498 bp fragment. The animals used were Lycat+/+ (wild-type) and Lycat−/- (homozygous) mice in the same C57BL/6 background between 13 and 20 weeks old. The mice were kept in a temperature-controlled environment with a 12 h light and 12 h dark cycle and received a standard diet and water ad libitum.

**Generation of monoclonal antibodies against mouse Lycat**

A recombinant mouse Lycat (amino acid numbers 78–208 of mouse Lycat, GenBank accession number NM_001081071) that was expressed and purified by Escherichia coli pCold TF expression system (TaKaRa, Japan) was injected into the hind foot pads of WKY/Izm rat strain by using Freund’s complete adjuvant. The enlarged mediatal iliac lymph nodes were used for cell fusion with mouse myeloma cells, PAI. In the present study, the established monoclonal antibody, named YN1, was used for Western blotting and immunocytochemistry at 1:2,000 and 1:100 dilutions, respectively.

**Western blot**

Murine tissues were homogenized in quadruple volumes (w/v) of SET buffer with protease inhibitors (0.5 mM phenylmethylsulfonil fluoride, 2 µg/ml pepstatin, 2 µg/ml leupeptin, 2 µg/ml aprotinin). After centrifugation at 1,000 g for 10 min at 4°C, the supernatants were used as the total protein extracts. The protein concentrations of samples were determined by the bicinchoninic acid method. The protein samples were run on a 15% SDS-PAGE gel, transferred to a nitrocellulose membrane, and probed with the monoclonal antibody YN1.
LYCAT determines the fatty acid composition of PI in mice

Fig. 4. Fatty acid compositions of PC, PE, PI, PS, PG and CL in liver, heart, skeletal muscle, and brain. After methyl esterification, fatty acids of each phospholipid were quantified by GC-MS. LYCAT+/−, closed bars; LYCAT−/−, open bars. FAs, fatty acids. *P<0.01, **P<0.001. Data represent the mean ± SEM of triplicate measurements.
Fig. 4. Continued.
Phospholipid analysis

Lipids of each tissue were extracted by the method of Bligh and Dyer (22). Phospholipids were separated from total lipids by one-dimensional TLC on silica gel 60 plates in chloroform-methanol-acetic acid (65:25:13, v/v). The area of silica gel corresponding to each phospholipid (PC, PE, PG, and PI+PS) was scraped off the plates. The PI+PS fraction was reextracted, separated by TLC in chloroform-methanol-formic acid-water (60:30:7:3, v/v), and the areas of silica gel corresponding to PI and PS were scraped off the plates. Isolated phospholipids were methylated with 2.5 % H2SO4 in methanol. The resulting fatty acid methyl esters were then extracted with hexane and subjected to GC-MS analysis as described previously (23). Liquid chromatography-electrospray ionization mass spectrometry (LC-ESIMS) analysis was performed as described previously (12) using 1,2-dipalmitoyl PI (16:0/16:0-PI), as internal standards. Phospholipids in the PI+PS fraction were purified by TLC and the chromatography was performed at room temperature (25). Column-bound lipids were washed with chloroform-methanol (1:1, v/v) and chloroform-methanol-28% aqueous ammonia-acetic acid (200:100:3:9.9, v/v) (3 ml) in series. Phosphoinositides were then eluted with chloroform-methanol-HCl-water (12:12:1:1, v/v) (1.5 ml). After addition of 0.75 ml of water and 0.1 ml of 1 M NaCl, the solution was shaken and centrifuged to collect the lower layer. After addition of 50 µl of 2 M TMS-diazomethane in hexane, samples were incubated for 10 min at room temperature (25). Then, 10 µl of glacial acetic acid and 0.7 ml of wash solution [chloroform-methanol-water (48:49:3, v/v)] were added to each solution. The solution was shaken and centrifuged to collect the lower layer. Each sample was dried under nitrogen gas and was redissolved in 24 µl of methanol-70% ethylamine (100:0.065, v/v), followed by addition of 8 µl of 1 M ammonium bicarbonate.

The LC-ESIMS/MS analysis was performed by using a TSQ Vantage (Thermo-Fisher Scientific) with an UltraMate 3000 LC system (Thermo-Fisher Scientific, Waltham, MA) combined with an HTC PAL autosampler (CTC Analytics, Zwingen, Switzerland). The phosphoinositides fractions were separated by a step gradient with mobile phase A (methanol-water-70% ethylamine (20:80:0.13, v/v))/mobile phase B [methanol-water-isopropanol-70% ethylamine (5:5:90:0.13, v/v)] ratios of 90%/10% (0 min), 70%/30% (0–1 min), 10%/90% (1–3 min), 10%/90% (3–15 min), 90%/10% (15–16 min), and 90%/10% (16–22 min). Flow rate was 30 µl/min and the chromatography was performed at room temperature using Waters X-Bridge C8 (3.5 µm, 150 mm × 1.0 mm i.d.) column. Phosphoinositides molecular species were measured by the selected reaction monitoring (SRM) in positive ion mode. The characteristic fragmentation patterns of individual phospholipids were determined by the product ion scan (MS/MS mode). Each molecular species was identified by the LC retention time.

RESULTS

Mouse LYCAT is a functional homolog of C. elegans acl-8, acl-9, and acl-10

We previously reported that C. elegans acl-8 acl-9 acl-10 triple mutants are defective in the divisions of specialized epithelial cells called seam cells (12). In the wild-type, 10 seam cells (H0-2, V1-6 and T in Fig. 1A) are present on each side of a newly hatched worm and divide asymmetrically in a stem cell-like manner during the larval stage. Both daughter cells just after the division express a seam cell marker, scm::gfp (Fig. 1B). Then, an anterior daughter
LYCAT deficiency causes reduced acyltransferase activity toward the sn-1 position of PI and PG in mouse liver

We generated LYCAT-deficient mice by gene targeting. A LYCAT-targeting construct was designed to replace exon dpy-7 promoter in acl-8 acl-9 acl-10 triple mutants. Expression of mouse LYCAT restored the parallel orientation of seam cell divisions in acl-8 acl-9 acl-10 triple mutants (Fig. 1D). The angle between the A-P axis and the direction of cell division fell within 10 degrees in most seam cells in LYCAT-expressing mutants. Expression of mouse LYCAT also effectively rescued the defects in cell-fate determination of the daughter cells in acl-8 acl-9 acl-10 triple mutants (Fig. 1G). These results indicate that LYCAT is a functional homolog of C. elegans acl-8, acl-9, and acl-10.
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4 encoding the acyltransferase motifs II and III that are conserved in the AGPAT family with a neomycin cassette (Fig. 2A, B) (15, 16). LYCAT protein was not detected in the liver of LYCAT−/− mice (Fig. 2C). LYCAT−/− mice were viable and fertile. They were outwardly healthy and had normal litter sizes. No histological abnormalities were detected in various organs from LYCAT−/− mice by light microscopy. There was no significant change in plasma triglyceride or cholesterol levels (supplementary Fig. III A) or hepatic triglyceride content (supplementary Fig. III B) in LYCAT−/− mice. Moreover, no significant changes were observed in serum markers of liver injury (AST and ALT) (supplementary Fig. III C) or the number of blood cells (leukocytes, platelets, and erythrocytes) (data not shown).

C. elegans acl-10 shows acyltransferase activity that incorporates stearic acid into the sn-1 position of PI in an in vitro assay (12). To assess the contribution of LYCAT to the acyltransferase activity toward the sn-1 position of phospholipids, we measured [14C]stearoyl-CoA:sn-2-acyl lysophospholipid acyltransferase activity using the liver membrane fraction of LYCAT−/− mice. Because sn-2-acyl lysophospholipid is known to easily isomerize to sn-1-acyl lysophospholipid, it is possible that the [14C]acyl donor is incorporated into both sn-1-acyl lysophospholipid and sn-2-acyl lysophospholipid (supplementary Fig. II). To check the positional specificity, [14C]phospholipid produced after the incubation was treated again with phospholipase A2 (for details, see Materials and Methods). The amount of radioactivity of the resultant lysophospholipid was taken as a measure of acylation at the sn-1 position (sn-2-acyl lysophospholipid acyltransferase activity) and the amount of radioactivity of the free fatty acid was taken as a measure of acylation at the sn-2 position (sn-1-acyl lysophospholipid acyltransferase activity). The sn-1- and sn-2-acyl lysophospholipid acyltransferase activities toward various lysophospholipids in the liver microsomes from wild-type and LYCAT−/− mice are summarized in Table 1. At least 90% of [14C] stearoyl-CoA was incorporated into the sn-1 position of all the lysophospholipids tested (Table 1), indicating that sn-2-acyl lysophospholipid acyltransferase activity was predominant under the present assay conditions. In the liver from LYCAT−/− mice, sn-2-acyl lysoPI acyltransferase (LPIAT) activity was reduced by 64% (from 8.10 ± 0.11 to 2.92 ± 0.22 nmol/min/mg protein) and lysoPG acyltransferase (LPGAT) activity was reduced by 56% (from 10.45 ± 0.05 to 4.57 ± 0.17 nmol/min/mg protein). The sn-2-acyl lysoPE acyltransferase (LPEAT) activity was also reduced, but only slightly. In contrast, sn-2-acyl lysoPC acyltransferase (LPCAT) and lysoPS acyltransferase (LPSAT) activities were unchanged in the liver from LYCAT−/− mice. These results indicate that LYCAT significantly contributes to acyltransferase activity toward the sn-1 position of PI and PG in mouse liver.

Furthermore, sn-1-acyl LPIAT activity was reduced by 33% (from 0.15 ± 0.00 to 0.10 ± 0.00 nmol/min/mg protein) and LPGAT activity was reduced by 57% (from...
Ferases activities toward the sn-2 position of PI and PG (20). sn-1-acyl LPCAT, LPEAT, and LPSAT activities were not altered significantly in the LYCAT−/− liver (Table 1).

Reduction of acyltransferase activity toward the sn-1 position of PI in various tissues of LYCAT−/− mice

As described above, LYCAT was largely responsible for the acyltransferase activity that incorporates stearic acid into the sn-1 position of PI in the liver (Table 1). We also examined acyltransferase activity toward the sn-1 position of PI in other tissues such as heart, skeletal muscle, and brain. As shown in Fig. 3, a significant portion of the activity was decreased in these tissues from LYCAT−/− mice. Especially in the heart and skeletal muscle, LYCAT was the main acyltransferase that incorporates stearic acid into the sn-1 position of PI.

LYCAT deficiency causes a major change in the fatty acid composition of PI

To determine the role of LYCAT in phospholipid metabolism in vivo, we analyzed the fatty acid composition of phospholipids in tissues from LYCAT−/− mice. GC-MS analysis revealed that LYCAT deficiency significantly altered the fatty acid composition of PI in all the tissues examined including liver, heart, skeletal muscle, and brain (Fig. 4). In LYCAT−/− mice, the amounts of stearic acid in PI of the liver, heart, and skeletal muscle were reduced by 42%, 36%, and 66% compared with the amounts in wild-type mice, respectively (Fig. 4A–C). The amount of arachidonic acid, which is a major fatty acid in the sn-2 position of PI, was also reduced in these tissues, though the degree of reduction was less than that of stearic acid. Conversely, the amounts of palmitic acid (16:0), oleic acid (18:1n-9), cis-vaccenic acid (18:1n-7), and linoleic acid (18:2n-6) in PI in these tissues increased. In the brains of LYCAT−/− mice, the change in the fatty acid composition of PI was

0.60 ± 0.07 to 0.26 ± 0.03 nmol/min/mg protein) (Table 1). This observation is consistent with the previous data that small interfering RNA (siRNA)-mediated knockdown of LYCAT in HeLa cells results in reductions of acyltransferase activities toward the sn-2 position of PI and PG (20). sn-1-acyl LPCAT, LPEAT, and LPSAT activities were not altered significantly in the LYCAT−/− liver (Table 1).
milder than that in the liver, heart, and skeletal muscle, and the amount of arachidonic acid in PI was not affected (Fig. 4D). Similarly, the amount of arachidonic acid in PI was not affected in the testis of LYCAT−/− mice (Fig. 5). Meanwhile, the amount of stearic acid in PI of the testis was reduced by 66% and the amounts of palmitic acid (16:0) and cis-vaccenic acid (18:1n-7) in PI increased. In contrast to the remarkable change in the fatty acyl chains of PI, no drastic changes were observed in the fatty acid composition of other phospholipids such as PC, PE, PS, PG, and CL (Fig. 4). One exception was a relatively large change in the fatty acid composition of PG in skeletal muscle; i.e., the amount of cis-vaccenic acid (18:1n-7) was reduced and instead, the amount of oleic acid (18:1n-9) increased considerably in LYCAT−/− mice (Fig. 4C). These results indicate that LYCAT mainly determines the fatty acid composition of PI in vivo.

We next analyzed the molecular species of PI in LYCAT−/− mice by LC-ESIMS. In the liver of LYCAT−/− mice, 18:0/20:4-PI, the major molecular species of PI, was significantly decreased, and conversely, 16:0/20:4-PI, 16:0/18:2-PI, and 18:1/18:2-PI were increased compared with those in the liver of wild-type mice (Fig. 6). The ratio of PI to total phospholipids was significantly reduced by 35% compared with the ratio in wild-type mice as judged by quantification of phospholipid phosphorus (Fig. 7). On the other hand, the content and molecular species of PC and PE were not affected in LYCAT−/− mice (Fig. 7 and data not shown). These results confirm the notion that LYCAT plays an important role in determining the molecular species of PI in mice. A deficiency in the reacetylation reaction of lysoPI may reduce the PI content.

**LYCAT deficiency affects the molecular species of phosphoinositides**

Because PI is a precursor for phosphoinositides (2), we also examined the molecular species of phosphoinositides in the liver of LYCAT−/− mice. For this, we used LC-ES-IMS/MS with the selected reaction monitoring (SRM) technique (see Materials and Methods). The amounts of each molecular species in PI, phosphatidylinositol monophosphates (PIP1), or phosphatidylinositol bisphosphates (PIP2) in the liver from wild-type and LYCAT−/− mice are shown in Fig. 8. In PI of the LYCAT−/− liver, the amount of 18:0/20:4 species was reduced by 27% and the amounts of 16:0/18:2, 18:0/18:2, 18:1/18:2, and 16:0/20:4 species increased (Fig. 8A). Similarly, in PIP1 and PIP2, the amounts of 18:0/20:4 species were reduced by 41% and 50%, respectively, and the amounts of 16:0/18:2, 18:0/18:2, 18:1/18:2, 16:0/20:4, and 16:0/22:6 species increased (Fig. 8B, C). These data indicate that LYCAT deficiency affects the molecular species of PIP1 and PIP2 as well as PI.

**LYCAT deficiency does not affect acyltransferase activity that incorporates arachidonic acid into the sn-2 position of PI**

In the liver, heart, and skeletal muscle of LYCAT−/− mice, the amount of arachidonic acid in PI was also reduced, though the degree of reduction was less than that of stearic acid (Fig. 4A–C). Arachidonic acid is the predominant fatty acid at the sn-2 position of PI in mammals and is incorporated into PI by LPIAT after de novo synthesis (5–8). We first investigated whether LYCAT contributes to acyltransferase activity that incorporates arachidonic acid into the sn-2 position of PI in liver, heart, skeletal muscle, and brain. Acyltransferase assays were performed using [13C]arachidonoyl-CoA and sn-1-acyl lysoPI as substrates. As shown in Fig. 9A, [13C]arachidonoyl-CoA: sn-1-acyl LPIAT activity was not altered significantly in all the LYCAT−/− tissues examined. Furthermore, the expression level of LPIAT1, a major acyltransferase that incorporates arachidonic acid into the sn-2 position of PI (11), was not changed in LYCAT−/− liver (Fig. 9B). These results indicate that LYCAT does not contribute to acyltransferase activity that incorporates arachidonic acid into the sn-2 position of PI in mice.

**LYCAT is localized to the ER**

Cao et al. (17) reported that flag-tagged mouse LYCAT transfected into COS-7 cells displayed a perinuclear and punctate pattern that colocalized well with an ER marker but not with a mitochondrial marker. However, using the same type of cells, another study recovered recombinant flag-tagged mouse LYCAT from mitochondria and mitochondrial-associated membranes but not from the microsomes (26). In the present study, we established a mouse LYCAT-specific monoclonal antibody YN1 and used it to analyze the subcellular localization of endogenous LYCAT. YN1 did not stain LYCAT−/− MEFs, indicating that YN1 specifically recognized mouse LYCAT (Fig. 10). Wild-type MEFs were costained with MitoTracker Red dye (a mitochondrial selective probe) or were costained with an anticalnexin antibody that recognizes the calnexin ER-specific protein and the LYCAT antibody. We found that anti-LYCAT staining overlapped substantially with anti-calnexin but not with MitoTracker (Fig. 10), suggesting that endogenous LYCAT is localized to the ER but not to the mitochondria.

**DISCUSSION**

Mammalian LYCAT is reported to possess acyltransferase activity toward the sn-2 position of anionic lysophospholipids including lysoPI, lysoPG, and lysoCL in vitro (17–20). In this study, we examined the fatty acid composition of each phospholipid from various tissues of LYCAT−/− mice, and showed that LYCAT determines the fatty acid composition of PI in vivo. 1-Stearoyl-2-arachidonoyl (18:0/20:4) PI is a major molecular species of PI in mammals. The amount of stearic acid in PI was greatly reduced in all the tissues examined from LYCAT−/− mice. Acyltransferase activity that incorporates stearic acid into the sn-1 position of PI was also reduced in the tissues from LYCAT−/− mice, whereas acyltransferase activity that incorporates arachidonic acid into the sn-2 position of PI was not altered. 1-Stearoyl-2-arachidonoyl PI is thought to be formed through fatty acid remodeling that is regulated by phospholipase A and lysoPI acyltransferase (5–10, 27,
Thus, LYCAT should serve as a lysoPI acyltransferase that catalyzes the incorporation of stearic acid into the sn-1 position of PI in the fatty acid remodeling pathway in vivo. Furthermore, mammalian LYCAT can substitute for the C. elegans homologs acl-8, acl-9, and acl-10 because expression of mouse LYCAT efficiently rescued the seam cell defects observed in acl-8 acl-9 acl-10 triple mutants, in which the fatty acid composition of the sn-1 position of PI is selectively altered (12). Our immunocytochemical analysis using MEFs revealed that endogenous LYCAT is mainly localized to the ER, suggesting that the fatty acid remodeling of PI by LYCAT occurs in the ER membrane.

In the membrane fraction of LYCAT−/− mouse liver, acyltransferase activity that incorporates stearic acid into the sn-1 position of PI was reduced to one-third of that in wild-type mice, whereas acyltransferase activities that incorporate stearic acid into the sn-1 position of other phospholipids such as PC, PE, and PS, were not reduced appreciably in the liver of LYCAT−/− mice, which suggests that one or more enzymes other than LYCAT contribute to the activities. In fact, the amounts of stearic acid in phospholipids such as PE and PS were not changed in LYCAT−/− mice. In addition, acyltransferase activity that incorporates stearic acid into the sn-1 position of PI, acyltransferase activity toward lysoPG was reduced significantly in the membrane fraction of LYCAT−/− mouse liver. Indeed, we observed a relatively large change in the fatty acid composition of PG in the skeletal muscle of LYCAT−/− mice. These results indicate that LYCAT partially determines the fatty acid composition of PG, at least in certain organs.

In this study, we found that the amount of arachidonic acid in PI was also reduced in the liver, heart, and skeletal muscle of LYCAT−/− mice. Acyltransferase activity that incorporates arachidonic acid into the sn-2 position of PI was not reduced in these tissues. Moreover, the expression level of LPIAT1, a major acyltransferase that incorporates arachidonic acid into the sn-2 position of PI (11), was not changed. Thus, the reduction of the amount of arachidonic acid in PI of LYCAT−/− mice may be secondary to the reduction of stearic acid at the sn-1 position of PI. One possible explanation for the reduced arachidonic acid in PI is that LPIAT1 preferentially incorporates arachidonic acid when stearic acid is attached at the sn-1 position of lysoPI. In LYCAT−/− mice, stearic acid attached to PI is replaced by other fatty acids, such as palmitic acid and oleic acid, which may lead to reduced incorporation of arachidonic acid into PI.

Unlike the liver, heart, and skeletal muscle, the brain and testis of LYCAT−/− mice did not show any significant changes in the amount of arachidonic acid in PI. It is interesting to note that these tissues possess CDS1 (29, 30), one isofrom of CDP-diacylglycerol synthase, which catalyzes the synthesis of CDP-diacylglycerol, a direct precursor of PI, from phosphatidic acid (PA). CDS1 prefers arachidonic acid-containing PA as a substrate (29) and therefore, CDS1 could have a role in increasing the arachidonic acid content of PI to compensate for the LYCAT deficiency in these tissues.

Recently, Li et al. (26) reported that the fatty acid composition of CL was altered in LYCAT−/− heart under high-fat diet conditions. In their study, however, the fatty acid composition of PI was not presented. The present results show that the fatty acid composition of PI was altered significantly in LYCAT−/− mice, whereas that of CL was less affected under normal diet conditions. So far, we have not observed any apparent abnormalities in LYCAT−/− mice under normal diet conditions. However, it is interesting that insulin-stimulated Akt phosphorylation was significantly enhanced in tissues from LYCAT−/− mice under high-fat diet conditions (26). PI plays important roles in intracellular signaling cascades, such as the PI3K/Akt pathway through distinct phosphorylated derivatives of the inositol head group (1, 2). Further analysis of LYCAT−/− mice should reveal the biological significance of PI molecular species and/or the fatty acid remodeling of PI in mammals

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