Polyunsaturated fatty acids promote the rapid fusion of lipid droplets in Caenorhabditis elegans

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Lipid droplets (LDs) are intracellular organelles that dynamically regulate lipids and energy homeostasis in the cell. LDs can grow through either local lipid synthesis or LD fusion. However, how lipids involving in LD fusion for LD growth is largely unknown. Here, we show that genetic mutation of acox-3 (acyl-CoA oxidase), maac-1 (enoyl-CoA hydratase), dhs-28 (3-hydroxylacyl-CoA dehydrogenase), and daf-22 (3-ketoacyl-CoA thiolase), all involved in the peroxisomal β-oxidation pathway in Caenorhabditis elegans, led to rapid fusion of adjacent LDs to form giant LDs (gLDs). Mechanistically, we show that dysfunction of peroxisomal β-oxidation results in the accumulation of long-chain fatty acid-CoA and phosphocholine, which may activate the sterol-binding protein 1/sterol regulatory element-binding protein to promote gLD formation. Furthermore, we found that inactivation of either FAT-2 (delta-12 desaturase) or FAT-3 and FAT-1 (delta-15 desaturase and delta-6 desaturase, respectively) to block the biosynthesis of polyunsaturated fatty acids (PUFAs) with three or more double bonds (n≥3-PUFAs) fully repressed the formation of gLDs; in contrast, dietary supplementation of n≥3-PUFAs or phosphocholine bearing these PUFAs allowed the formation of gLDs in peroxisomal β-oxidation-defective worms lacking PUFA biosynthesis. Thus, we conclude that n≥3-PUFAs, distinct from other well-known lipids and proteins, promote rapid LD fusion leading to LD growth.

Lipid droplets (LDs) are intracellular organelles present in some prokaryotes and almost all eukaryotic cells from yeast to humans. Distinguished from other organelles, LDs have a unique architecture consisting of a neutral lipids core, mainly enriched in triacylglycerols (TAGs) and sterol esters, which is wrapped by a phospholipid (PL) monolayer decorated with LD proteins (1–3). In cells, the main function of LDs is to dynamically regulate energy homeostasis through a fascinating cycle of biogenesis and consumption in response to energy surplus or requirement (4). Disturbance of this homeostasis often leads to many human metabolic disorders, including insulin resistance, type 2 diabetes, fatty liver disease, cardiovascular diseases, and so on, which are prevalent diseases threatening human health worldwide.

The lipid storage capacity of LDs is controlled by its growth via local lipid synthesis or by LD fusion (5). It is widely accepted that the LDs are initiated and nascent LDs are formed from the endoplasmic reticulum (ER) (6–8). The nascent LDs may grow to mature ones by acquiring neutral lipids from the ER through a continuous association with the ER (9–12). Alternatively, after the nascent LDs bud from the ER, LD-associated enzymes can directly localize to LDs and synthesize TAG at the surfaces of LDs (13–15), which is then incorporated to produce LD growth.

Fusion or coalescence of adjacent LDs can also lead to expanded LDs or giant LDs (gLDs). So far, three main forms of LD fusions have been reported. The first is a slow fusion (often takes around 60 min) of LDs regulated by CIDE (cell death–inducing DNA fragmentation factor alpha–like effector) family proteins that enrich at the LD–LD contact site (16), mediating the directional lipid transfer from a small to a large LD in various tissues including adipose tissues and liver. Second, when the phosphatidylcholine (PC) level is reduced in Drosophila, and/or the phosphatidic acid (PA) level is increased in yeast (9, 17, 18), the LDs in close proximity to each other contact and mix their content within seconds. Third, in Caenorhabditis elegans, temperature shifting from 20 to 30 °C can induce LD fusion, which happens within 30 s (19). However, the underlying mechanisms for both the PL-mediated and the thermosensitive LD fusion are still unclear. For example, the carbon length or saturation level of

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n≥3-PUFAs promote lipid droplet fusion

the fatty acid chain in PA/PC-mediated LD fusion is completely unknown. In addition, it is also unknown whether there are other distinct mechanisms leading to LD fusion/growth.

In cells, the peroxisomes, similar to LDs, regulate lipid and energy metabolism. Peroxisomes and LDs share biogenesis steps (20, 21), and membrane contact sites between LDs and peroxisomes have also been described (22, 23). Mice lacking peroxisome biogenesis genes, such as PEX5 (24), single peroxisomal enzyme deficiencies in ACOX (acyl-CoA oxidase) (25), or multifunctional enzyme type (multifunctional protein) (26), display enlarged LDs in the liver, testis, or retina. Consistently, dysfunction of ACOX homologs (CG4586 and CG9527) in Drosophila (27), or PRX-10 and peroxisomal β-oxidation in C. elegans (28–30), also leads to the enlarged LDs. These conserved observations across evolutionary organisms demonstrate a connection between peroxisome and LD growth. However, the role of peroxisomes in LD growth/ expansion and the underlying mechanism remains largely unknown.

Using the genetically tractable animal model C. elegans, we showed that the formation of gLDs in animals with a dysfunction in peroxisomal β-oxidation was actually because of the fusion of adjacent LDs in less than 30 s. Notably, we discovered that n≥3-PUFAs might trigger this rapid fusion process.

Results
Disruption of peroxisomal β-oxidation triggers the rapid fusion of adjacent LDs to form gLDs

Peroxisomes are subcellular organelles that function in multiple anabolic and catabolic processes, in particular the β-oxidation of long-chain fatty acids (LCFAs). In C. elegans, similar to other organisms, the peroxisomal β-oxidation cycle involves four steps catalyzed by four distinct enzymes, ACOX, MAOC-1 (enoyl-CoA hydratase), DHS-28 (3-hydroxyacyl-CoA dehydrogenase), and DAF-22 (3-ketoacyl-CoA thiolase) (28, 30, 31) (Fig. 1A). Both Nile Red and LipidTOX have been widely used to visualize LDs in fixed C. elegans through staining the neutral lipid TAGs and cholesterol esters (32, 33). DHS-3 encodes a short-chain dehydrogenase, which is almost exclusively localized on C. elegans LDs. Therefore, DHS-3::GFP [N2, ldrIs1[Pdhs3::dhs-3::gfp+unc-76(+)] is also used as an LD marker in C. elegans (33). In the intestine of C. elegans, the major site of lipid synthesis and storage, all the aforementioned three methods consistently revealed that the acox-3(tm4033), maoc-1(hj13), dhs-28(tm2581), and daf-22(ok693) mutants displayed a few gLDs (5–6% of total LDs larger than 5 μm), which were undetectable in the wildtype N2 worms (Fig. 1, B and C). These results demonstrate that disruption of any step of the peroxisomal β-oxidation pathway leads to gLDs.

C. elegans contains seven paralogs of ACOXs including ACOX-3 (Fig. S1A), which is a close homolog of CG4586 in Drosophila, and ACOX1 in mice and humans (Fig. S1B) (34, 35). We therefore tested whether other members of the acox family also regulated the LD size. Nile Red staining of fixed worms showed that none of the other acox mutant worms displayed an altered LD size, with the exception of the acox-3(tm4033) mutant worms showing gLDs and the acox-1.5(ok2619) mutant worms showing a slightly reduced LD size (Fig. S1, C and D). Next, to test if the gLDs in the acox-3(tm4033) mutant was specifically because of the dysfunction of ACOX-3, we generated a transgenic worm strain acox-3::gfp [N2;unc-119(ed3);kunEx233[Pacoxx-3::acox-3::gfp]]. Through GFP fluorescence analysis, ACOX-3::GFP is mainly expressed in the intestine of worms and had no obvious effect on LD size (Fig. S1, E–G). We then crossed the acox-3::gfp strain into the acox-3(tm4033) mutant background. As expected, the gLDs completely disappeared in the acox-3(tm4033);acox-3::gfp worms compared with the acox-3(tm4033) mutant worms (Fig. S1, F and G), confirming that gLD formation is indeed because of the loss of function of acox-3.

The gLDs may originate from the growth of LD per se or by the fusion of adjacent LDs (Fig. 1D). The gLDs in the worms with a peroxisomal β-oxidation defect display an unevenly/ nonasymmetric distribution, which looks comparable to the LD phenotype of the reported cct-1 mutant in the Drosophila S2 cell line (17) and the fld-1 mutant in yeast (9), in which the adjacent LDs fuse to form a gLD. We therefore hypothesized that the formation of gLDs in worms with a peroxisomal β-oxidation defect may be a result of adjacent LD fusion. Using time-lapse video microscopy on living worms with the LD marker DHS-3::GFP uncovered that two adjacent LDs closed, contacted, and then fused into a gLD in 10 to 30 s in all acox-3(tm4033), maoc-1(hj13), dhs-28(tm2581), and daf-22(ok693) mutant worms but not in the wildtype N2 worms (Fig. 1E) (Movies S1–S5). In order to avoid the possible unknown role of DHS-3 overexpression on the LD fusion process, we also performed the time-lapse video microscopy on living worms using bright-field mode with differential interference contrast and found the same effects on LD fusion in the peroxisomal β-oxidation defect worms (Fig. 1F) (Movies S6–S10). Thus, dysfunction of peroxisomal β-oxidation can trigger LD fusion that leads to gLDs. Since this process of LD fusion is distinctly very fast (less than 30 s), we named this process as the rapid fusion of LDs.

Dysfunction of peroxisomal β-oxidation causes fatty acid-CoA accumulation

C. elegans contains a series of unsaturated fatty acids including n-3 and n-6 polyunsaturated fatty acids (PUFAs), which can be de novo synthesized from acetyl-CoA through many steps of elongations and desaturations (31, 36, 37) (Fig. 2A). These unsaturated fatty acids, also called LCFAs, can be used as substrates for the biosynthesis of complex lipids or be broken down into medium-chain fatty acids/short-chain fatty acids (SCFAs) through peroxisomal β-oxidation (Fig. 1A). We wondered if dysfunction of peroxisomal β-oxidation might affect fatty acid metabolism. We first examined the fatty acid compositions in these peroxisomal β-oxidation mutants. However, GC analysis did not show a
Figure 1. Dysfunction of peroxisomal β-oxidation pathway led to the formation of giant LDs (gLDS) in Caenorhabditis elegans. A, diagram of fatty acid catabolism by the ACOX-3/MAOC-1/DHS-28/DAF-22 peroxisomal β-oxidation pathway in C. elegans. B, lipid droplets (LDs) of L4 worms including the wildtype strain N2, acox-3(tm4033), maoc-1(hj13), dhs-28(tm2581), and daf-22(ok693) mutants were visualized by postfixed Nile Red staining, postfixed

$n\geq3$-PUFAs promote lipid droplet fusion

J. Biol. Chem. (2022) 298(8) 102179 3
significant change of the fatty acid compositions in these mutants compared with the wildtype N2 (Table S1).

In fact, fatty acids must first be activated by acyl-CoA synthase to form fatty acid-CoA in order for anabolism or catabolism to proceed. Therefore, we detected the fatty acids-CoA using LC–MS in these peroxisomal β-oxidation mutants. In total, the level of acyl-CoA was consistently significantly increased in three tested mutant worms, acox-3(tm4033), dhs-28(tm2581), and daf-22(ok693), compared with the wildtype N2 worms (Fig. 2B). Of which, the levels of SCFAs-CoA C10:0-CoA and C12:0-CoA were decreased (Fig. 2, D and E), whereas the levels of seven LCFA-CoA including C18:0-CoA, C18:1-CoA, C18:2-CoA, C18:3-CoA, C20:4-CoA, C20:5-CoA, and C22:6-CoA were increased (Fig. 2, H–N), in the aforementioned mutant worms. Taken together, these results suggest that the C. elegans peroxisomal β-oxidation catalyzes the conversion of LCFA-CoA to SCFAs-CoA.

Figure 2. Accumulation of PUFAs-CoA in the peroxisomal β-oxidation defect worms. A, diagram of the biosynthesis of PUFAs in Caenorhabditis elegans. B–N, the levels of acyl-CoA (B), C6:0-CoA (C), C10:0-CoA (D), C12:0-CoA (E), C16:0-CoA (F), C16:1-CoA (G), C18:0-CoA (H), C18:1-CoA (I), C18:2-CoA (J), C18:3-CoA (K), C20:4-CoA (L), C20:5-CoA (M), and C22:6-CoA (N), in the wildtype N2, acox-3(tm4033), dhs-28(tm2581), and daf-22(ok693) mutants. All data are presented from three independent biological replicates. Significant difference between a specific mutant and wildtype N2, *p < 0.05, **p < 0.01, and ***p < 0.001. PUFA, polyunsaturated fatty acid.

Inactivation of PUFA biosynthesis eliminates the gLDs in peroxisomal β-oxidation mutants

As mentioned previously, dysfunction of peroxisomal β-oxidation causes not only gLDs but also altered fatty acid metabolism, raising the question of whether they had a direct connection. The gLDs might be caused by either the reduction of SCFAs-CoA or the accumulation of LC/very long-chain fatty acids-CoA in the peroxisomal β-oxidation mutants. Since the four peroxisomal β-oxidation mutant worms displayed consistent gLDs and altered fatty acids-CoA, we chose the dhs-28(tm2581) and daf-22(ok693) mutants for most of the following experiments for convenience.

To distinguish both possibilities, we first fed dietary fatty acids C10:0 or C12:0 to dhs-28(tm2581) and daf-22(ok693) mutant worms. However, dietary supplementation of C10:0 or C12:0 could not repress the gLD formation in these mutant worms (Fig. S2, A and B), suggesting that the gLDs...
might not be caused by the reduction of SCFAs-CoA. In *C. elegans*, the desaturases FAT-1, FAT-2, and FAT-3 are involved in the biosynthesis of PUFAs containing two or more double bonds (Fig. 3A) (31, 37). Quantitative PCR (qPCR) analysis revealed that the mRNA levels of *fat-1*, *fat-2*, and *fat-3* were significantly upregulated in both *dhs-28(tm2581)* and *daf-22(ok693)* mutant worms compared with the wildtype N2 worms (Fig. 3B), implying an upregulated desaturation of PUFAs in peroxisomal β-oxidation mutants.

Figure 3. Inactivation of n≥3-PUFAs biosynthesis eliminates gLD formation in peroxisomal β-oxidation defect worms. A, diagram of the structure and biosynthesis pathway of PUFAs in *Caenorhabditis elegans*. B, the relative mRNA level of *fat-1*, *fat-2*, and *fat-3* in the wildtype N2, *dhs-28(tm2581)*, and *daf-22(ok693)* mutant worms. C, Nile Red staining of fixed L4 worms. For all representative animals, anterior is left and posterior is right. The scale bar represents 10 μm. The dashed circles and arrows indicated representative gLDS. D, percentage (%) of gLDS with diameters >5 μm quantified from (C), n = 10 for each worm strain. All data are presented from three independent biological replicates. Significant difference between a specific mutant and wildtype N2, *p < 0.05, **p < 0.01, and ***p < 0.001. gLD, giant LD; PUFA, polyunsaturated fatty acid.
To determine which specific LCFA was required for the formation of gLDs, we then crossed fat-1(wa9), fat-2(wa17), and fat-3(ok1126) mutants into the dhs-28(tm2581) and daf-22(ok693) mutants, respectively. Inactivation of FAT-2 results in a loss of all PUFAs (Figs. 3A and S2C). As well, the gLDs also disappeared in the fat-2(wa17);daf-22(ok693) and fat-2(wa17);dhs-28(tm2581) double mutants compared with their corresponding daf-22(ok693) and dhs-28(tm2581) single mutant, respectively (Fig. 3, C and D). In addition, the fat-2(wa17) mutation also suppressed the gLDs in the acox-3(tm4033) and maoc-1(h13) mutant worms (Fig. S2, D and E). These results indicate that PUFAs are necessary for the formation of gLDs in the peroxisomal β-oxidation mutants.

Inactivation of both FAT-1 and FAT-3 blocks the biosynthesis of PUFAs with three or more double bonds (Figs. 3A and S2C). Similar to the fat-2(wa17);daf-22(ok693) and fat-2(wa17);dhs-28(tm2581) double mutants, the fat-3(ok1126);fat-1(wa9);daf-22(ok693) or fat-3(ok1126);fat-1(wa9);dhs-28(tm2581) triple mutants also had undetectable gLDs (Fig. S2, C and D), suggesting that PUFAs bearing three or more double bonds (simply named as n≥3-PUFAs), but not C18:2(n-6), are necessary for the formation of gLDs in the peroxisomal β-oxidation mutants. Inactivation of FAT-3 alone blocks the biosynthesis of C18:3(n-6), C18:4(n-3), and their downstream products, C20-PUFAs (Figs. 3A and S2C). However, the fat-3(ok1126);daf-22(ok693) and fat-3(ok1126);dhs-28(tm2581) double mutants still had gLDs (Fig. 3, C and D), which might be due to the presence of C18:3(n-3). Meanwhile, inactivation of FAT-1 impairs the conversion of omega-6 (n-6) PUFAs to omega-3 (n-3) PUFAs (Figs. 3A and S2C), whereas the fat-1(wa9);daf-22(ok693) and fat-1(wa9);dhs-28(tm2581) double mutants still displayed gLDs (Fig. 3, C and D), suggesting that the C18:3(n-6), C20:3(n-6), and C20:4(n-6) in these mutants may still be involved in the formation of gLDs. Altogether, these lines of evidences uncover that n≥3-PUFAs (LCFAs) are necessary for the formation of gLDs in the peroxisomal β-oxidation mutants.

**Dietary n≥3-PUFAs are sufficient to promote the formation of gLDs in the peroxisomal β-oxidation mutants**

As mentioned previously, the loss of n≥3-PUFAs (LCFAs) by mutations of desaturases could abolish the formation of gLDs in the peroxisomal β-oxidation mutants. We next asked whether supplementation of n≥3-PUFAs could restore the gLDs in these mutants. Since either the fat-3(ok1126);fat-1(wa9) double mutations or the fat-2(wa17) single mutation could completely abolish the gLDs in the daf-22(ok693) and dhs-28(tm2581) mutant worms, we therefore tested their response to dietary supplementation of n≥3-PUFAs. We first confirmed the successful uptake of each dietary PUFA by worms using GC (Fig. S3).

Inactivation of FAT-2 blocks the biosynthesis of C18:2(n-6) and its downstream fatty acids (Figs. 3A and S2C). Dietary C18:2(n-6), C18:3(n-6), C18:3(n-3), C20:4(n-6), and C20:5(n-3) had no effect to promote the formation of gLDs in the wildtype N2, fat-2(wa17) single mutant, and fat-3(ok1126);fat-1(wa9) double mutant worms (Fig. 4, A and B). However, remarkably, these dietary fatty acids could fully restore the gLDs in the fat-2(wa17);daf-22(ok693) (Fig. 4, A and B) and fat-2(wa17);dhs-28(tm2581) mutant worms (Fig. S4, A and B), suggesting that PUFAs are essential to promote gLD formation in the peroxisomal β-oxidation mutants. Inactivation of both FAT-1 and FAT-3 impairs the conversion of C18:2(n-6) to n≥3-PUFAs (Figs. 3A and S2C). Dietary C18:3(n-3), C18:3(n-6), C20:4(n-6), and C20:5(n-3), but not C18:2(n-6), also successfully restored the gLDs in the fat-3(ok1126);fat-1(wa9);daf-22(ok693) and fat-3(ok1126);fat-1(wa9);dhs-28(tm2581) triple mutant worms (Fig. S4, A and B) triple mutant worms. Thus, the recovery of the gLDs in the fat-3(ok1126);fat-1(wa9);daf-22(ok693) and fat-3(ok1126);fat-1(wa9);dhs-28(tm2581) triple mutant worms by individual dietary C18:3(n-3), C18:3(n-6), C20:4(n-6), or C20:5(n-3), but not C18:2(n-6), clearly demonstrates that PUFAs with three or more double bonds are sufficient to trigger the formation of gLDs in the peroxisomal β-oxidation mutants.

**n≥3-PUFAs–PC promotes gLD formation in peroxisomal β-oxidation mutants**

PUFAs can be incorporated into complex lipids such as TAGs and PLs, including PC, phosphatidylethanolamine (PE), phosphatidylserine (PS), which composes the lipids core and the enclosed PL monolayer of LDs, respectively. Since n≥3-PUFAs are required for the formation of gLDs in the peroxisomal β-oxidation mutants, we thereby asked which complex lipid containing n≥3-PUFAs is involved in this process.

To do so, the total lipids (nTL), nTAG, and nPL (containing 50 µg fatty acids), which contains n≥3-PUFAs in different degrees, were isolated from the wildtype N2 worms ("n" stands for lipids isolated from N2) and then were individually added to the nematode growth media (NGM) plates and cultured with different worm strains. The isolated complex lipids with PUFAs were confirmed using GC (Fig. 5A). Supplementation of nTL, nTAG, and nPL, similar to dietary PUFAs (Fig. 4, A and B), did not cause any effect on the LD phenotypes of the N2, daf-22(ok693), fat-3(ok1126);fat-1(wa9) worms. However, supplementation of nTL and nPL completely reproduced the gLDs in the fat-3(ok1126);fat-1(wa9);daf-22(ok693) mutant worms (Fig. 5, A and B). In addition, nTAG, which also contains a very small amount of C20-PUFAs such as 20:4(n-3) and 20:5(n-3) (31), had the same effect but to a much lesser degree for the formation of gLDs (Fig. 5, A and B).

The PLs are divided into several distinct molecular classes, distinguished by the head group attached to the phosphate moiety, such as PC, PE, PI, and PS. To determine which PL type is involved in the formation of gLDs, we further extracted and separated C. elegans nPC, nPE, nPI, and nPS and then individually added them to different worm strains. We used the commercial standard sPC, sPE, sPI, and sPS as their corresponding controls ("s" is standard for commercial lipids). Very consistently, similar to applying nPL, applying nPC and sPC, which contains a small amount of C20:4(n-6) (Fig. 5, A and B), could successfully trigger the formation of the gLDs in the fat-3(ok1126);fat-1(wa9);daf-22(ok693) mutant worms (Fig. 5, A and B). These results indicate that PUFAs with three or more double bonds are necessary for the formation of gLDs in the peroxisomal β-oxidation mutants.
and B). In contrast, supplementation of fPC, which was isolated from fat-3(ok1126);fat-1(wa9) mutant worms that completely lack n≥3-PUFAs, or commercial 16:0-PC bearing C16:0 at the sn-1 and sn-2 positions of PC, failed to produce gLDs in the fat-3(ok1126);fat-1(wa9) mutant worms (Fig. 5, A and B). Meanwhile, the levels of total PC and PC bearing UFAs containing one or more double bonds were significantly increased in the daf-22(ok693) and dhs-28(tm2581) mutant worms compared with the wildtype N2 worms (Fig. 5, C–F). On the other hand, surprisingly, although the nPE, nPI, and nPS contain n≥3-PUFAs, they could not recover the gLDs in the fat-3(ok1126);fat-1(wa9);daf-22(ok693) mutant worms (Fig. S5, A and B). Moreover, the dietary sPE, sPI, and sPS, which lack PUFAs, also had no effect to trigger gLD formation in the fat-3(ok1126);fat-1(wa9);daf-22(ok693) mutants (Fig. S5, A and B).

Altogether, these lines of evidences suggest that the choline
**Figure 5. n≥3-PUFA-PC determines the gLD formation.** 

_A, left panel._ Nile Red staining of fixed L4 worms supplemented with different lipid species. “n” represents lipids isolated from N2 worms. “s” represents standard commercial PL purchased from Sigma. “f” represents lipid isolated from the fat-3(ok1126);fat-1(wa9) worms. For all representative animals, anterior is left and posterior is right. The scale bar represents 10 μm. The dashed circles and arrows indicated the representative gLDs. The right panel showed the GC profile of PUFAs in different lipid species supplemented to worms. “1” indicates C20:3(n-6), “2” indicates C20:4(n-6), “3” indicates C20:4(n-3), and “4” indicates C20:5(n-3). B, percentage (%) of LDs with diameters > 5 μm as quantified from (A), n = 10 for each worm strain. C–F, the levels of total PC (C), n≥3 PC with three or more double bonds on the sn-2 position (D), n = 2 PC with two double bonds (n = 2) on the sn-2 position (E), C18:1 PC with C18:1 on the sn-1 and/or sn-2 position (F) in worms. All data are presented from three independent biological replicates. Significant difference between a specific mutant and wildtype N2 under same treatment, *p < 0.05, **p < 0.01, and ***p < 0.001. gLD, giant lipid droplet; PC, phosphatidylcholine; PL, phospholipid; PUFA, polyunsaturated fatty acid.
headgroup and n≥3-PUFAs as fatty acid chain are two indis-

pensable factors for PC to promote gLD formation in the

eroxisomal β-oxidation mutants.

Dysfunction of peroxisomal β-oxidation activates sterol
regulatory element–binding protein to promote gLD
formation

The sterol regulatory element–binding proteins (SREBPs)
play important roles regulating lipid homeostasis from
C. elegans to mammals (31, 38). The sterol-binding protein 1
(SBP-1), a homolog of mammalian SREBP-1, has been
reported to regulate fatty acid and PL biosynthesis in
C. elegans (39–42). A previous report showed that blocking
SAMe or PC synthesis in C. elegans, mouse liver, and human
cells activates SREBP-1 to promote LD accumulation (40). We
thereby asked whether dysfunction of peroxisomal β-oxid-
ation, which leads to the accumulation of fatty acids-CoA and
PC, could also regulate SREBP-1 to trigger gLD formation.

qPCR analysis showed that the mRNA expression of sbp-1
was significantly increased in daf-22(ok693) and acox-
3(tm4033) mutant worms compared with the wildtype (N2)
worms (Fig. 6A). In addition, we crossed the gfp::sbp-1
[N2;ftsIs7(Psbp-1::gfp::sbp-1)] strain into daf-22(ok693) and
acox-3(tm4033) mutants and found that the fluorescence
intensity of GFP::SBP-1 was consistently upregulated in nu-
uclear but downregulated in cytosol in daf-22(ok693) and acox-
3(tm4033) mutant worms (Fig. 6, B and C), suggesting an
activated SBP-1 when the peroxisomal β-oxidation is dis-
rupted. Next, to test whether SBP-1 was necessary for gLD
formation, we crossed sbp-1(1ep79) mutant into daf-22(ok693)
and acox-3(tm4033) mutant to generate sbp-1(1ep79);daf-
22(ok693) and sbp-1(1ep79);acox-3(tm4033) double mutant
worms. Although the mRNA expression of fat-1, fat-2, and fat-
3 was significantly increased in both daf-22(ok693) and
acox-3(tm4033) mutant worms, inactivation of SBP-1 (1ep79
mutation) could abolish these effects (Fig. 6A). More im-
portantly, the gLDS disappeared in sbp-1(1ep79);daf-22(ok693) and
sbp-1(1ep79);acox-3(tm4033) double mutant worms compared
with the daf-22(ok693) or acox-3(tm4033) single mutant
worms, respectively (Fig. 6, D and E), suggesting that the gLD
formation depends on the activity of SBP-1. Collectively, these
results demonstrate that peroxisomal β-oxidation dysfunction
activates SBP-1 to promote gLD formation.

Discussion

LDS are dynamic organelles whose number and size un-
dergo constant changes in response to internal and external
energy demands (43). Previous studies have reported that the
LD fusion can be mediated by CIDE family proteins, taking
around an hour, as well as reduction of PC (18) in Drosophila
(17), or an increase of PA in yeast within a minute (9). The
C. elegans peroxisomal β-oxidation pathway consists of four
enzymes: ACOX-3, MAOC-1, DHS-28, and DAF-22 to break
down the LCFAs. In this study, we found that inactivation of
this process leads to a few gLDS, which are actually formed by
the fusion of adjacent LDS within 30 s. Furthermore, we
revealed that inactivation of peroxisomal β-oxidation conse-
quently led to the accumulation of n≥3-PUFAs-CoA, which
may be incorporated into PC to mediate this rapid fusion
process. Thus, we discovered a completely distinct type of
rapid LD fusion because of inactivation of peroxisomal
β-oxidation in the model organism C. elegans.

In C. elegans, the peroxisomal β-oxidation pathway was
previously reported to catalyze the β-oxidation of LCFAs (30).
However, the levels of LCFAs were not changed in the
peroxisomal β-oxidation mutants compared with the wildtype
N2, which is consistent to a previous report (30). In fact, fatty
acids must be activated by acyl-CoA synthase to form fatty
acid-CoA in order for anabolism or catabolism to occur. Our
data showed that the levels of LCFAs-CoA but not LCFAs
were indeed significantly increased, while reversely, the levels
of SCFAs-CoA were significantly decreased in the peroxisomal
β-oxidation mutants. Therefore, our work uncovers that
peroxisomal β-oxidation actually breaks down the LCFAs-
CoA, instead of LCFAs, confirming this biological function
of LCFAs-CoA β-oxidation in the peroxisome in C. elegans.

As the structural lipids in eukaryotic membranes, PLs
mainly contain PC, PE, PS, PI, and PA. The PC amount is
more than 50% of all PLs (44) and is the major PL in
C. elegans (31, 45, 46). PC is cylindrical in shape and provides
excellent coverage of the surface area and lower surface ten-
sion. Therefore, PC is considered crucially important for LD
stability (47). Dysfunction of key synthesis enzyme CCT-1 of
PC in Drosophila S2 cells (17) or artificial droplets generated
in vitro with low PC (18) leads to gLDS or to LD coalescence
via a fusion process. Independent from these findings and
on the other hand, we reveal that excessive accumulation of
specific PC species may also be involved in LD fusion to form
gLDS in the peroxisomal β-oxidation defect mutants in
C. elegans. Thus, these findings, including our current study,
consistently reinforce the idea that the PC regulates LD fusion,
because of either its deficiency or accumulation in distinct
animal or cell models.

PC commonly involves one cis-unsaturated acyl chain, for
instance eicosapentaenoic acid C20:5(n-3), an abundant fatty
acid in C. elegans. Five cis-double bonds create a strong kink
that decreases the packing compactness of the acyl chains and
in turn raises membrane fluidity (48). This may influence
membrane curvature and fusion events. Peroxisomal biogen-
esis disorders and peroxisomal β-oxidation defects (PEX13,
PEX16, and ACOX1) showed elevated levels of PC species with
very long–chain FAs in fibroblasts from patients (49, 50). Very
consistently, our in-depth analysis of lipids revealed that
inactivation of peroxisomal β-oxidation promoted the accu-
mony of both LCFAs-CoA, total PC, and PC containing
LCFAs in C. elegans. In the peroxisomal β-oxidation defect
mutants, inactivation of n≥3-PUFAs biosynthesis by inactiva-
tion of both FAT-1 and FAT-3 blocked the formation of gLDS,
which could be rescued by supplementation of PC only with
n≥3-PUFAs (n≥3-PUFAs-PC) but not PC without PUFAs,
either isolated from C. elegans or by using commercially
available products. In addition, feeding other PLs such as PE,
PS, PI with or without n≥3-PUFAs had no such an effect. Thus,
Figure 6. Dysfunction of peroxisomal β-oxidation activates the transcriptional factor SREBP/SBP-1 to promote gLD formation. A, the relative mRNA level of sbp-1, fat-1, fat-2, and fat-3 in different worm strains. B, the fluorescence intensity of nuclear and cytosolic GFP::SBP-1 (N2;fts7(Psbp-1::gfp::sbp-1)) in different worm strains. The scale bar represents 10 μm. C, quantification of the fluorescence intensity of nucleus and cytosol GFP::SBP-1 in different worm strains from (B). D, Nile Red staining of fixed L4 worms. For all representative animals, anterior is left and posterior is right. The scale bar represents 10 μm. The dashed circles and arrows indicated representative gLDs. E, percentage (%) of LDs with diameters >5 μm as quantified from (D). n = 10 for each worm strain. All data are presented from three independent biological replicates. Significant difference between a specific mutant and wildtype N2, *p < 0.05, **p < 0.01, and ***p < 0.001. Significant difference between two indicated worm strains, *p < 0.05, **p < 0.01, ***p < 0.001. gLD, giant lipid droplet; SBP-1, sterol-binding protein 1; SREBP, sterol regulatory element-binding protein.
we uncovered that it is actually some specific PC species bearing n≥3-PUFAs, but not the general PC, that promotes the rapid LD fusion process. Whether n≥3-PUFAs and n≥3-PUFAs-PC functioning as signal or the membrane constitute of LDs for gLD formation needs future investigation.

The transcription factor SREBP play critical roles to transcriptionally regulate genes involved in fatty acids, triglycerides, and cholesterol metabolism from C. elegans to mammals. Previously, we revealed a zinc-mediated regulation of the SREBP-SCD (stearoyl-CoA desaturase) axis in lipid metabolism in C. elegans (51). Our current work showed that dysfunction of peroxisomal β-oxidation leads to accumulation of PUFA-CoA and PC, which may activate the nuclear translocation of SREBP to promote gLD formation, distinguishing from the negative regulation of SREBP by PC in C. elegans (40). In the peroxisomal β-oxidation defect mutants, whether increased fatty acids-CoA or PCs (or a specific lipid species) activate SREBP directly or via another unknown factor and how activated SREBP thereby regulates gLD formation through its transcriptional activity or unknown function needs to be further investigated.

In summary, we propose a model to explain our current work. Inactivation of peroxisomal β-oxidation leads to the accumulation of n≥3-PUFAs-CoA, which in turn, may be incorporated into the PC to promote the rapid fusion of LDs, consequently resulting in gLD formation (Fig. 7).

**Experimental procedures**

**Nematode strains**

Worm strains used in this study are listed in Table S2. N2 Bristol was used as the wildtype strain. C. elegans worm strains were grown on NGM plates seeded with Escherichia coli OP50. All experiments were conducted at 20 °C. All experiments were conducted on NGM plates containing E. coli OP50 (CGC) as the food source. A single clone of E. coli OP50 was inoculated into LB broth and cultured at 37 °C overnight to stationary phase. Then E. coli was seeded directly onto NGM plates. After 24 h drying, the plates were ready for feeding worms.

**Construction of transgenic strains**

The transgenic strains were created by microinjection following the methods previously described (51). For construction of plasmid Pacox-3:acoX-3:gfp, the acox-3 gene of C. elegans, including its own 2.845 kb promoter sequence, was

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**Figure 7. Proposed model.** Inactivation of peroxisomal β-oxidation leads to the accumulation of n≥3-PUFAs-CoA, which in turn, may be incorporated into the PC to promote the rapid fusion of LDs, consequently resulting in gLD formation. gLD, giant LD; LD, lipid droplet; PC, phosphatidylcholine; PUFA, polyunsaturated fatty acid.
cloned by PCR. The GFP fragment was amplified from the pPD95.75 vector. Meanwhile, the vector pCFJ151 was also fragmented for use. The fragments of Pacox-3:acox-3 and gfp were recombined into the pCFJ151 vector to generate transgenic plasmids. The primers used for construction of transgenes are listed in Table S3. Approximately 50 ng/μl transgenic plasmids, 50 ng/μl pIL43.1, and 3 ng/μl pCF90 were injected into the gonad of EG4322 [ttTi5605;unc-119(ed3)], and progenies expressing GFP were selected (52).

**Fatty acid supplementation**

Fatty acids were purchased from Nuk Prep, Inc. Individual fatty acid was mixed with bacteria in LB at a concentration of 1 mM. Then, 400 μl of the mixed bacteria were seeded onto the NGM plates (6 cm in diameter). Standard PLs, including sPC, sPE, sPI, and sPS, as well as lipids isolated from *C. elegans* N2 worms including nTAG, nPL (PLs), nTL (total lipids), nPC, nPE, nPI, and nPS, each of these lipids containing equal amount of fatty acids (50 μg) for normalization, were mixed with bacteria and seeded on NGM plates. Synchronized L1 worms were transferred to the NGM plate seeded with *E. coli* and fatty acid/TAG/PL. The worms were harvested at the L4 stage and used for Nile Red staining and GC analysis.

**Staining of LDs**

Nile Red and LipidTOX were used as the fluorescent dye for staining of neutral fat. Staining of fixed L4 worms was performed as previously described (32, 53). Images of Nile Red staining were used to measure the diameter of the LDs in the posterior of the intestine with the same area (80 μm × 60 μm) using cellSens Standard software (Olympus). For each worm strain with or without treatment, at least three biological repeats were performed and more than 60 worms were observed, 10 representative worms were picked for the quantitation of LD size (diameters).

**Lipid extraction, separation, and analysis**

Worm lipid extraction, separation, and analysis were performed as previously reported (53). In general, about 200,000 L4 worms were collected, and the protein was quantified with a Pierce BCA Protein Assay Kit. About 15 mg protein was used for total lipid extraction. For lipid separation, 40 μl of total lipid was loaded in triplicate on TLC silica plates (Merck) to separate TAG, PE, PI, PS, and PC. The TLC plates were developed in the solvent system chloroform:methanol:water:triethylamine (30:35:7:35) (54) and dried in fume hood. About 0.005% primuline was sprayed evenly on the plate. The spots corresponding to TAG, PE, PI, and PS were visualized under UV light and scraped from TLC plates and trans-esterified (2.5% H2SO4 in methanol). Next, esterified C15:0 was added as an internal standard, and lipids were extracted for GC analysis to determine the relative levels of TAG and PL fractions as previously described (45). Fatty acids were determined with an Agilent 7890 series gas chromatographer equipped with a 30 m × 0.25 mm × 0.25 μm DBWAX column (Agilent), with nitrogen as the carrier gas at 1.4 ml/min, and a flame ionization detector (53).

**Analysis of acyl-CoAs**

Extraction of acyl-CoAs was carried out as previously described with some modifications (55). Briefly, 300 μl of extraction buffer containing isopropanol, 50 mM KH2PO4, and 50 mg/ml bovine serum albumin (25:25:1) acidified with glacial acetic acid was added to the worms. Next, 19:0-CoA was added as an internal standard, and lipids were extracted. To the remaining extracted sample, 5 μl of saturated ammonium sulfate was added followed by 600 μl of chloroform:methanol (1:2). The sample was then incubated on a thermonixer at 450 rpm for 20 min at 25 °C, followed by centrifugation at 12,000 rpm for 5 min at 4 °C. Clean supernatant was transferred to a fresh tube and subsequently dried in a SpeedVac under OH mode (Genevac). Dry extracts were resuspended in an appropriate volume of methanol:water (9:1) prior to LC–MS analyses on a Thermo Fisher U3000 DGLC coupled to Sciex QTRAP 6500 Plus.

**Visualization of GFP**

L4 worms with GFP markers were washed from NGM plates and settled down in 1.5 ml tubes. The worms were then pipetted and plated onto an agarose gel (2%). Worms were visualized using an Olympus BX53 fluorescent microscope or a fluorescent dye for Olympus BX53 fluorescence microscope or ZEISS M2 (Zeiss). Images were captured using the same settings and exposure times for each worm, unless specifically indicated, and were analyzed by the CellSens or Zen 2 software.

**Confocal microscopy**

The LD labeled by DHS-3::GFP was acquired on a confocal microscope LSM880 (Zeiss). For detection of the size of LD, worms with DHS-3::GFP were anesthetized in 1% sodium azide on 2% agarose pads sandwiched between glass microscopic slides and coverslips. About 30 worms of each strain were observed and scored, and the experiment was conducted in three biological replicates. For detection of the LD fusion process, worms were anesthetized in 1 mM levamisole on the lower lid of confocal petri dishes. Then the worms were then covered with 2% agarose and used for microscopy observation. The process of LD fusion was acquired in a λ mode using a 63×, numerical aperture 1.43, with 100 intervals. A z-stack of 7 to 9 sections with 0.5 μm in depth was taken for each animal.

**mRNA isolation and qPCR**

mRNA of L4 worms was isolated as previously described (56). Complementary DNA was synthesized using the PrimeScript RT reagent kit. mRNA levels of each sample were quantified in three biological replicates using SYBR green fluorescence on a real-time PCR instrument 7900HT (ABI). The relative abundance of mRNA transcripts was determined using the ΔΔCt method, and tbb-2 was used as a reference gene. The primers used for construction of transgene qPCR are listed in Table S3.
Quantification and statistical analysis

All experiments were conducted with at least three or four biological replicates. Data were analyzed using the SPSS software (SPSS, Inc), and presented as the means ± SD, unless specifically indicated. The one-way ANOVA statistical analysis was performed for the significant difference among the control group and experimental groups, and statistical analysis by Student’s t tests was used for comparison between the means of two groups. All figures were generated using GraphPad Prism 8 (GraphPad Software, Inc) and Photoshop CS6.

Data availability

All data are contained within the article and supporting information.

Supporting information—This article contains supporting information.

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Conflict of interest—The authors declare that they have no conflicts of interest with the contents of this article.

Abbreviations—The abbreviations used are: ACOX, acyl-CoA oxidase; DAF-22, 3-ketooacyl-CoA thiolase; DHS-28, 3-hydroxylacyl-CoA dehydrogenase; ER, endoplasmic reticulum; gLD, giant LD; LCFA, long-chain fatty acid; LD, lipid droplet; MAOC-1, enoyl-CoA hydratase; NGM, nematode growth media; PA, phosphatidic acid; PC, phosphatidylcholine; PE, phosphatidylethanolamine; PI, phosphatidylinositol; PLC, phospholipid; PS, phosphatidylserine; PUFA, polyunsaturated fatty acid; qPCR, quantitative PCR; SBP-1, sterol-binding protein 1; SCFA, short-chain fatty acid; SREBP, sterol regulatory element–binding protein; TAG, triacylglycerol.

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