A Genetic Screen for Mutants with Supersized Lipid Droplets in Caenorhabditis elegans

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ABSTRACT To identify genes that regulate the dynamics of lipid droplet (LD) size, we have used the genetically tractable model organism Caenorhabditis elegans, whose wild-type LD population displays a steady state of size with an upper limit of 3 μm in diameter. From a saturated forward genetic screen of 6.7 × 10^6 mutagenized haploid genomes, we isolated 118 mutants with supersized intestinal LDs often reaching 10 μm. These mutants define nine novel complementation groups, in addition to four known genes (maoc-1, dhs-28, daf-22, and prx-10). The nine groups are named drop (lipid droplet abnormal) and categorized into four classes. Class I mutants drop-5 and drop-9, similar to prx-10, are up-regulated in ACS-22-DGAT-2-dependent LD growth, resistant to LD hydrolysis, and defective in peroxisome import. Class II mutants drop-2, drop-3, drop-6, and drop-7 are up-regulated in LD growth, are resistant to LD hydrolysis, but are not defective in peroxisome import. Class III mutants drop-1 and drop-8 are neither up-regulated in LD growth nor resistant to LD hydrolysis, but seemingly up-regulated in LD fusion. Class IV mutant drop-4 is cloned as sams-1 and, different to the other three classes, is ACS-22-independent and hydrolysis-resistant. These four classes of supersized LD mutants should be valuable for mechanistic studies of LD cellular processes including growth, hydrolysis, and fusion.

The fat triacylglycerol (TAG) is stored in lipid droplets (LDs), a type of cellular organelle conserved across eukaryotes (Martin and Parton 2006). While the importance of LD in cellular fat storage and fat mobilization has emerged through several lines of research, mechanisms underlying basic LD cellular processes such as de novo formation, growth, fusion, and hydrolysis are still poorly understood (Ploegh 2007; Walther and Farese 2012; Yang et al. 2012). Changes in the processes of de novo formation, growth, fusion, or hydrolysis may lead to changes in the number or size of LDs. Thus, size change may serve as a convenient morphological entry point to dissect the underlying metabolic pathways that anabolize and catabolize fat, and to uncover the protein-lipid membrane machinery that channels fat into and out of LDs.

Candidate gene approaches based on gene knockouts or proteomics have led to the identification and an initial understanding of the roles of LD surface proteins such as PLINs, ATGL, DGAT, and Cidec in the regulation of LD hydrolysis, growth, and fusion (Walther and Farese 2012). As confirmed by several studies, functional perturbation of LD processes through gene manipulation often leads to changes in LD size, number, or distribution. Thus, genetic screens for morphological LD alterations may be used as a first step to discover genes and to relate gene functions to cognate LD processes. Along this line, whole genome reverse genetic screens have been undertaken to identify genes whose functional loss leads to changes in LD size, number, or spatial distribution at the single-cell level. For example, whole genome RNAi screens in Drosophila S2 cells uncovered the involvement of Arfl-COPI complex in LD morphology and a role of CTP-phosphocholine cytidylyltransferase (CCT) in typical LD fusion (Beller et al. 2008; Guo et al. 2008; Krahmer et al. 2011). Deletion mutant library screens in the single-celled yeast Saccharomyces cerevisae revealed the roles of SEIPIN and phosphatidic acid metabolism enzymes in typical LD fusion (Szymanski et al. 2007; Fei et al. 2008, 2011). Unbiased saturated forward genetic screens for morphological LD mutants in multicellular systems, however, have rarely been reported.
Through a combination of biochemical, histological, and microscopy approaches, Zhang and colleagues recently demonstrated that LDs rather than lysosome-related organelles (LROs) are the fat storage organelles in Caenorhabditis elegans (Zhang et al. 2010b). In C. elegans, vital Nile Red staining labels LROs but not LDs. However, postfix Nile Red staining and postfix Oil-Red-O staining specifically label LDs (Zhang et al. 2010a,b). These results also demonstrated that “fat stores accessible to postfix Nile Red” and “fat vesicles distinct from LROs” defined in two earlier reports (Brooks et al. 2009; O’Rourke et al. 2009) were actually LDs, and explained why staining intensity of postfix Nile Red and postfix Oil-Red-O but not of vital Nile Red may be used as a semiquantitative measurement of relative TAG level (Schroeder et al. 2009; Brooks et al. 2009; O’Rourke et al. 2009). In contrast to vital Nile Red, vital BODIPY labels LDs in addition to LROs. For this reason, BODIPY fluorescence intensity may not be used to quantify relative TAG level. However, vital BODIPY is still qualitatively useful for labeling LDs in wild type and for labeling LDs specifically when LROs are abolished in a glo-4 mutant background (Zhang et al. 2010a,b).

As the only qualitatively reliable vital LD dye currently available, BODIPY should be valuable for the fast identification of mutants with LD size changes in forward genetic screens of a large number of live animals. This argument is supported by the fact that in a screen of $3.6 \times 10^4$ haploid genomes, vital BODIPY labeling allowed the isolation of 11 bona fide supersized LD mutants corresponding to four genes: maoc-1, dhs-28, daf-22, and prx-10 (Butcher et al. 2009; Zhang et al. 2010a). MAOC-1, DHS-28, and DAF-22 are three consecutive enzymes for peroxisomal fatty acid b-oxidation. PRX-10 is responsible for the import of peroxisome matrix enzymes including MAOC-1, DHS-28, and DAF-22. Loss-of-function mutation of these proteins results in a decrease of fatty acid breakdown and an increase of fatty acid storage in the form of TAG. The increase of TAG storage is expressed as an enhancement of LD growth concurrent with an inhibition of LD hydrolysis. The net effect of growth and hydrolysis is accumulated across developmental stages, resulting in supersized LDs 3–10 μm in diameter by larval stage L4 and even larger in adults (Zhang et al. 2010a). The enhanced LD growth is dependent on the TAG-synthesis enzyme complex ACS-22/FATP1-DGAT-2, which is conserved between C. elegans and mammals (Xu et al. 2012). Whether the BODIPY-based forward genetic screen was saturated was not tested.
To fully exploit the BODIPY-based strategy for isolating supersized LD mutants, we have now conducted a saturated genetic screen with several innovations: (1) increasing the haploid genome number 18.6 times to \(6.7 \times 10^6\), (2) using both ethylmethanesulfonate (EMS) and ethynitrosoure (ENU) as mutagens, (3) applying a novel temperature-sensitive mutant selection regimmen, (4) using high-copy transgenic lines expressing wild-type MAOC-1/DSHS-28/DAF-22 as a mutagenesis background in order to target only new genes, and (5) using acs-22 mutant as a mutagenesis background to enrich mutants of novel genes that areACS-22-independent. In this way, we isolated in total 118 mutants. Forty-five are new alleles of four previously reported genes for peroxisome function: maoc-1, dhs-28, daf-22, and prx-10; the other 73 define nine novel complementation groups that are named drop (lipid droplet abnormal) and are categorized into four classes. These four classes of mutants represent valuable models for the regulation of LD growth, hydrolysis, and fusion.

**MATERIALS AND METHODS**

**C. elegans strains and culture conditions**

The wild-type strain was N2 Bristol. The mapping strain was a Hawaiian isolate CB4856. All strains were raised on nematode growth media (NGM) plates seeded with OP50 Escherichia coli. Growth temperature was 20°C if not otherwise indicated. Mutant alleles and transgenic lines published before or obtained from other sources are: maoc-1(hij3) II, daf-22(ok693) II, prx-10(hij21) III, glo-4(ok623) V, acs-22(tm3236) X, dhs-28(tm2581) X, hijIs37[vha-6p::mrfp::pts1], hijIs73[vha-6p::gfp::daf-22], and hijIs65[vha-6p::3xflag::1vpc::gfp::dgat-2] II.

The 13 complementation groups of 118 mutant alleles (reference alleles in bold) isolated in the current study are:

- maoc-1/ssd64, ssd65, ssd66, ssd67, ssd196) II
- dhs-28/ssd39, ssd40, ssd41, ssd42, ssd43, ssd44, ssd45, ssd46, ssd47, ssd48, ssd50, ssd197, ssd207, ssd208, ssd209) X
- daf-22/ssd51, ssd52, ssd53, ssd54, ssd55, ssd56, ssd57, ssd58, ssd59, ssd60, ssd61, ssd62, ssd63, ssd195, ssd198, ssd200, ssd201, ssd202, ssd204, ssd205, ssd210, ssd215, ssd217) II
- prx-10(ssd68, ssd69) III
- drop-1/ssd1, ssd2, ssd3, ssd4, ssd5, ssd6, ssd7, ssd8, ssd9, ssd10, ssd11, ssd12, ssd79, ssd80, ssd81, ssd82, ssd83, ssd84, ssd85, ssd86, ssd87, ssd88, ssd203, ssd212, ssd219, ssd220, ssd221, ssd222, ssd223, ssd224, ssd225, ssd226, ssd227, ssd228, ssd229, ssd230, ssd231, ssd232, ssd233, ssd211) IV
- drop-2/ssd13, ssd14, ssd15, ssd16, ssd17, ssd18, ssd19, ssd20, ssd21, ssd22, ssd23, ssd24, ssd25, ssd26, ssd27, ssd28, ssd29, ssd30, ssd31, ssd32, ssd33, ssd211) IV
- drop-3/ssd34, ssd35, ssd36, ssd37, ssd74) V
- drop-4(ssd206) X
- drop-5(ssd72, ssd216) X
- drop-6(ssd38, ssd73) X
- drop-7(ssd75)
- drop-8(ssd89) III
- drop-9(ssd213) II

Also generated in this study are dgat-2(ssd90) V, ssdIs1[dhs-28; maoc-1; 3xflag::daf-22::SL2::mrfp::pts1] V, ssdIs5[dhs-28; maoc-1; 3xflag::daf-22::SL2::mrfp::pts1] V, ssdIs1 and ssdIs5 were made by UV integration of an extrachromosomal array ssdIs1[X(dhs-28); maoc-1; 3xflag::daf-22::SL2::mrfp::pts1], which was derived from an injection of plasmid vha-6p::dhs-28, plasmid vha-6p::maoc-1, and plasmid vha-6p::3xflag::daf-22::SL2::mrfp::pts1, each at 10 ng/μl.

**Mutagenesis and screening**

N2 Bristol (WT), ssdIs1/ssdIs5 (Is[mdmd]), acs-22(tm3236) were used as mutagenesis backgrounds. Synchronized L4 hermaphrodites of each background were mutagenized with 50 mM EMS (ethylmethanesulfonate, #M880-10G, Sigma-Aldrich) or 0.5 mM ENU (ethylnitrosourea, #N3383-1G, Sigma-Aldrich) for 4 hr. ENU was stored and applied essentially as described in De Stasio and Dorman (2001). Healthy mutagenized P0 animals were picked onto 6 cm NGM/OP50 plates, 3 to 5 P0s per plate. On each plate, P0s were allowed to lay 150–200 F1s. Each plate of F1s was an independent pool of haploid genomes. For each screen series, about 300 pools/plates were set up. F1s on each plate were allowed to lay enough F2 eggs. The plate was then washed, leaving only F2 eggs. F2 eggs were allowed to hatch for 2–4 hr to produce 1500–3000 synchronized stage L1 larvae. Synchronized F2 larvae were transferred onto a BODIPY-layered NGM/OP50 plate to grow for 2–3 d at 15°C or 20°C, and were then transferred to 30°C for 4 hr. F2s were screened under a Nikon SMZ1500 stereo microscope in green fluorescent
protein (GFP) fluorescence and bright field channels. Candidate super-sized LD mutants were picked out by the presence of enlarged BODIPY-positive globular structures in intestinal cells. From each pool only a single isolate was kept. Candidate isolates identified by BODIPY and bright field visualization were further tested by postfix Oil-Red-O staining, postfix Nile Red staining, and GFP::DGAT-2 marking.

Vital BODIPY staining and visualization
A total of 100 μl of 5 μM green BODIPY (#D-3823, Invitrogen) or red BODIPY (#D-3835, Invitrogen) in 1X phosphate buffered saline (PBS, pH 7.2) was added to the ~2 cm-in-diameter OP50 bacteria lawn of a 6 cm NGM plate. The plate was immediately dried in a laminar flow hood to grow and stain animals. For screening, freshly hatched L1s were loaded onto the plate to grow to L4 and adult stages. BODIPY fluorescent staining was examined under a Nikon SMZ1500 stereo microscope. For high power confocal imaging, young adult animals were loaded onto the BODIPY plate to produce progeny and late L4 stage progeny were then mounted in 0.2 mM levamisole on agarose-padded slides for imaging.

Postfix Oil-Red-O staining and bright field imaging
Late L4 stage animals were fixed in 1% paraformaldehyde/PBS for 30 min with rocking. Samples were then immediately frozen on dry ice/ethanol and thawed with running tap water. Samples were frozen and thawed again, three times in total. Samples were washed three times with 1X PBS, dehydrated in 60% isopropanol for 2 min, stained with 0.5 ml 60% Oil-Red-O (#O0625-25G, Sigma-Aldrich) working solution for 30 min with rocking. Samples were then immediately frozen on dry ice and thawed three times in total. Oil-Red-O stock solution was dissolved in isopropanol at a concentration of 0.5 g/100 ml and was equilibrated for several days. Oil-Red-O working solution was prepared fresh by mixing 60% volume of stock with 40% volume of water. The mix was equilibrated for 10 min, and then filtered with a 0.22 μm spin-filter. Stained samples were then washed three times with 1X PBS, rehydrated in 1X PBS, and mounted in 1X PBS onto agarose-padded slides for imaging. Some Oil-Red-O color images were acquired on a Zeiss Axiovert 200M compound microscope using a FLUAR 40X/1.30 Oil Iris objective. The camera was a RETIGA-SRV FAST1394 device (QIMAGING) controlled by Image-Pro Plus 7.0 software. Other Oil-Red-O images were acquired on a Zeiss Imager M1 compound microscope using an EC Plan-Neofluar 40X/1.30 Oil DIC objective. The camera used was a Zeiss AxioCam HRc5 controlled by Zeiss Axiovision LE 4.0 software.

Postfix Nile Red staining
Late L4 animals were fixed, freeze-thawed, washed, and dehydrated in the same way as in postfix Oil-Red-O staining. Animals were then stained with 0.5 ml of 1 μg/ml Nile Red (#N1142, Invitrogen) in 60% isopropanol for 30 min. Stained samples were then washed three times with 1X PBS, rehydrated in 1X PBS, and mounted in 1X PBS onto agarose-padded slides for confocal imaging.

Confocal fluorescence imaging and spectral analysis
All confocal fluorescence images were acquired on a Zeiss LSM780 inverted confocal microscope with a Plan-Apochromat 63X/1.40 Oil DIC M27 or EC Plan-Neofluar 40X/1.30 Oil DIC M27 objective. Channel mode imaging was applied to BODIPY, GFP::DGAT-2, GFP::DAF-22, and mRFP::PTS1 fluorescence. For GFP and green BODIPY, the excitation laser was 488 nm and emission was collected at 493–598 nm; for mRFP and red BODIPY, excitation was 543 nm, and emission 582–728 nm. Time-lapse channel mode imaging of GFP::DGAT-2 was at one frame every 30 sec. Each frame was a 9 μm Z-stack and was exported as an extended focus view. Lambda mode imaging and spectral analysis were conducted essentially in the same way as that described in Zhang et al. (2010b). For lambda mode imaging of GFP::DGAT-2 and autofluorescence, excitation was 488 nm and emission was 495–647 nm at 9 nm per channel. GFP and autofluorescence signals were separated using linear unmixing. For lambda mode imaging of postfix Nile Red fluorescence, excitation was 514 nm and emission was 540–691 nm at 9 nm per channel. LD fluorescence was separated from the diffuse background fluorescence using linear unmixing.
continuously at 20°C or shifted to 30°C for 4 hr to reach late L4 stage. These animals were then fasted in M9 buffer at 20°C for 24 hr. LDs in these animals are almost completely hydrolyzed. In each Oil-Red-O experiment, > 50 stained animals were examined under a stereo scope, and at least 20 animals were imaged on a compound microscope. In other figures, if not otherwise indicated, all staining was conducted on late stage L4 animals, fasting was at 20°C, the same numbers of Oil-Red-O-stained animals were examined and imaged, and images are anterior half of the animal with head to the left.

**Genetic background cleaning**

All strains were crossed at least four times before phenotype analysis. Mutants isolated from N2 Bristol/WT background were backcrossed four times with N2. Mutants isolated from *acs-22(tm3236)* background were backcrossed four times with *acs-22(tm3236)* and then segregated out of *acs-22(tm3236)*. Mutants isolated from *Is[mddm]* were backcrossed four times with *Is[mddm]* and then segregated out of *Is[mddm]*.

**Complementation test**

The 118 mutants were tested for complementation against *maoc-1 (hj13), dhs-28(tm2581), prx-10(hj21)*, and *daf-22(ok693)*. Forty-five mutants were identified as new alleles of these four groups. The other 73 mutants were tested against each other and were sorted into nine groups.

**maoc-1/dhs-28/da f-22/prx-10 genomic DNA sequencing**

The primers used for amplifying and sequencing the four genes’ genomic DNA are listed below. Exons, intron splice donor sites, and intron acceptor sites were covered for each gene.

- *maoc-1*: 0062F: GTTTTTTGTTCCTCTCGGAC/0071R: CAGACTTCTGAAACTCTGG, 0072F: CCAGATAGAGCTCCAGAATGACATCTTAGCAAACTCTGG, 0072F: CCAGATAGAGCTCCAGAACTCTGG, 0073F: CGAAGAGCAGAGGTAGTGC/0067R: GTACCTTGGCCTCGAGCGGAG, 0068F: CAACTACGCTGCTGCCAAAAG/0073R: CGGAGGAATTTGCTACTTAATCTTACCAGTG, 0074F: CAGAGTGCTAGACTTGAGAATATATTGTTGGATGAG

**Genetic mapping**

Genetic mapping was conducted essentially in the same way as that described in Davis *et al.* (2005). Briefly, *drop* mutants were crossed with the Hawaiian strain CB4856. F2 progeny with supersized LDs were picked out and tested for homologous recombination events that brought Hawaiian single nucleotide polymorphisms (SNP) into N2 Bristol background.

**Construction of double mutants of drop with *acs-22 or dgat-2***

We constructed the following double mutants: *drop-1(ssd9); acs-22(tm3236), drop-2(ssd14); acs-22(tm3236), drop-3(ssd36); acs-22(tm3236), acs-22(tm3236), drop-4(ssd206); acs-22(tm3236), drop-5(ssd72), dgat-2(ssd90); drop-6(ssd73), drop-7(ssd75); acs-22(tm3236), drop-8(ssd89); acs-22(tm3236), and drop-9(ssd213); acs-22(tm3236). For constructing double mutants, *drop* mutants were mated with *acs-22(tm3236) or dgat-2(ssd90); at least eight F2s with supersized LDs were picked out and singled to produce F3s. F2s were subjected to *acs-22/dgat-2* genotyping. If F2s were *acs-22* homozygous, *i.e.*, *drop/drop; acs-22/acs-22*, F3s derived from the F2 were kept as double mutants. The *drop* mutant was inferred as *ACS-22/DGAT-2*-independent. If none of the F2s was *acs-22/dgat-2* homozygous but heterozygous, F3s derived from F2 heterozygotes were counted to calculate the ratio of supersized LD animals. If the ratio was three quarters, the *drop* mutant was inferred as *ACS-22/DGAT-2*-dependent. F3s without supersized LDs were singled to produce F4s, and were then subjected to *acs-22/dgat-2* genotyping. F4s of *acs-22/dgat-2* homozygous F3s were kept as double mutants.

**Brood size test**

Five stage L4 animals (P0s) of each *drop* reference allele were singled to lay eggs. Each P0 was transferred onto a new plate every 24 hr to lay eggs. Each P0 was allowed to lay eggs for a total of 144 hr when no more eggs were produced. The number of F1 progeny that eventually developed to L4 stage was counted.
Fasting

Fasting was conducted essentially as in Zhang et al. (2010a).

LD purification and western blot

Purification of LDs was essentially as in Zhang et al. (2010b) and Zhang et al. (2012) with some modifications. Briefly, 2 × 10³ freshly hatched wild-type or hjSi56 animals were grown on 20 15 cm-NGM/OP50 plates. Animals were allowed to grow to the 1-d adult stage, then harvested and washed three times with 1X PBS/0.001% Triton X-100 (PBST). Animals were then washed with 5 ml buffer A [25 mM Tricine, pH 7.6, 250 mM sucrose, one protease inhibitor Complete Tablet (PBST). Animals were then washed with 5 ml buffer A [25 mM Tricine, pH 7.4, 100 mM KCl, and 2 mM MgCl₂] was gently added on top of the homogenate was transferred into a 15 ml centrifuge tube. Buffer A was added to the tube to bring the total volume to 10 ml. Homogenate was centrifuged at 1000 g for 10 min. LD protein were extracted. Briefly, 1 ml acetone was added to the LD or TM sample. Samples were vortexed rigorously and then chilled at −20° for 1 hr. Samples were centrifuged at 16,000 g for 10 min. LD protein and TM protein were spun down as pellets. Pellets were air dried and were dissolved in 30 μl 2X SDS protein sample buffer. Protein concentrations were measured using a BCA kit (#23227, Thermo Scientific). A total of 10 μg of each protein sample was loaded and separated on 8% SDS-PAGE according to standard methods. Proteins separated were subjected to western blot using a mouse anti-FLAG antibody (#F3165-1MG, Sigma-Aldrich) at 1:500 dilution.

Data availability

Strains and data are available upon request. The authors state that all data necessary for confirming the conclusions presented in the article are represented fully within the article.

RESULTS

Design of the screen

The main aim of this study was to conduct a large-scale screen for recessive loss-of-function mutants with supersized LDs in C. elegans. We extended a previous screen by exploiting several innovations. First, in addition to using the wild-type N2 as a mutagenesis background, we used an integrated stable transgenic line (termed Is[mddm]) bearing multiple copies of wild-type maoc-1, dhs-28, daf-22, and a peroxisome import marker mRFP:PTS1 cDNAs driven by an intestine-specific vha-6 promoter. This line was used to bypass the reisolation of maoc-1/dhs-28/daf-22, to immediately identify prx-10 based on the mislocalization of peroxisome marker protein mRFP:PTS1 (Zhang et al. 2010a), and to
enrich new genes. Second, since the supersized LD phenotype of *maoc-1(dhs-28)/daf-28*/*prx-10* is a result of enhanced LD growth mediated by the ACS-22-DGAT-2 enzyme complex (Xu et al. 2012), we used a deletion mutant *acs-22(tm3236)* as a mutagenesis background in hope of enriching new genes representing novel models of supersized LD formation. Third, before phenotype screening we shifted the F2 progeny from a low cultivation temperature of 15°C or 20°C to 30°C for the brief time of 4 hr, in hope of finding temperature-sensitive (ts) mutants with a fast response to temperature shift. We failed to find any ts mutants with a shift to 25°C so we switched to 30°C. Fourth, we used ENU as a mutagen in addition to EMS. ENU has a broader mutagenic spectrum of DNA base substitution and fragment deletion (Anderson 1995; De Stasio et al. 1997; De Stasio and Dorman 2001). ENU may increase the chance of isolating genes with very low allele frequency, especially those rare ts alleles. In total we conducted six series of screens: EMS/WT, ENU/WT, EMS/Is[mddm], ENU/Is[mddm], EMS/acs-22, and ENU/acs-22. Each series allowed both non-ts and ts mutant isolation (Figure 1A). Fifth, in all series except one, we increased the haploid genome number to ~1.0 × 10^5. The total number of haploid genomes screened was more than half a million (Table 1).

We used five labeling and visualization criteria to identify supersized LDs: vital BODIPY staining, bright field visualization, postfix Oil-Red-O staining, postfix Nile Red staining (Zhang et al. 2010a,b), and GFP::DGAT-2 LD marker labeling (Xu et al. 2012). To test whether GFP::DGAT-2 specifically labels LDs, we purified LDs and TM from *hjsi56 [vha-6p::3xflag::gfp::dgat-2]* transgenic animals, and found that DGAT-2 localizes to LD but not to TM (Supplemental Material, Figure S1A). Under lambda mode confocal microscopy, LRO’s autofluorescence appears as a filled globe, while LD’s GFP::DGAT-2 fluorescence appears as an encircled globe. The two kinds of fluorescence display distinct emission spectra and the globular structures labeled by each completely exclude the other (Figure S1, B–E). *glo-4* mutation eliminates LROs, leaving fluorescent structures 100% GFP::DGAT-2 positive (Figure S1, F and G). These data corroborated the specificity of GFP::DGAT-2 as an LD marker. In wild type, 78.9% of vital BODIPY-labeled structures are encircled by GFP::DGAT-2, while 21.1% are not and should be LROs since *glo-4* mutation leaves BODIPY structures 99.9% encircled by GFP::DGAT-2 (Figure S2, A–C).

To screen for recessive mutants, F2 progeny of mutagenized P0s were grown on BODIPY-layered NGM/OP50 media. F2s with supersized LDs were picked out by virtue of BODIPY fluorescence and bright light-refraction under a stereoscope at 180× magnification. Among the isolates picked out from the six screening series, only 118 (about a third of total) survived and proliferated. F3 and further generations of
these 118 isolates were subjected to postfix Oil-Red-O staining, postfix Nile Red staining, and GFP::DGAT-2 labeling (Figure 1A) which unsurprisingly confirmed all 118 as bona fide supersized LD mutants (e.g., Figure 1, B–E), a further validation of each LD labeling and visualization method.

Identification of nine novel complementation groups of supersized LD mutants

The 118 isolates were backcrossed four times, segregated from the original genetic background when necessary, and examined for temperature sensitivity. During crossing, we found that all mutants are recessive loss-of-function, completely penetrant, and with no detectable maternal effect. These isolates were tested for complementation first against known maoc-1, dhs-28, daf-22, and prx-10 alleles. We found five maoc-1 alleles, 15 dhs-28, 23 daf-22, and two prx-10. The other 73 were tested against each other and sorted into nine novel complementation groups named drop (lipid droplet abnormal) (Table 1). The isolation origins of these 13 complementation groups display expected patterns. maoc-1, dhs-28, and daf-22 were bypassed in the EMS/ENU/Isl[mdm] and EMS/ENU/acs-22 series; prx-10 was bypassed in the EMS/ENU/acs-22 series. Interestingly, six of the nine drop groups, drop-2, 3, 5, 6, 7, and 9, were also bypassed in the EMS/ENU/acs-22 series. The temperature regimen proved a success: five of the nine groups, drop-1, 3, 5, 6, and 8, are ts. Curiously, the temperature sensitivity is either all or none: all alleles of drop-1, 3, 5, 6, and 8 are ts; all alleles of drop-2, 3, 5, 6, and 9, maoc-1, dhs-28, daf-22, and prx-10, are non-ts. The restrictive temperature of drop-1, 3, 5, 6, and 8 is 30°C as was screened for, while that of drop-5 (20°C) occurred fortuitously.

Heterozygotes of the only allele of drop-7 segregated 1/16 supersized LD progeny (see later), indicating mutations in two unlinked genetic loci. Heterozygotes of the other eight drop groups segregated 1/4 supersized LD progeny, indicating mutation in a single locus. Seven of the eight single drop loci were positioned by SNP-based chromosome mapping and three-point interval mapping (Davis et al. 2005) (Figure S3); the other locus drop-4 was cloned by direct sequencing. The eight single loci reside in discrete short physical regions of Chr II, III, IV, and X (Figure 2).

Class I (drop-5 and drop-9) is up-regulated in LD growth and defective in peroxisome import

As indicated by isolation origins, the drop mutants may be classified according to whether they are dependent on ACS-22 for supersized LD phenotype and whether they affect peroxisome marker mRFP::PTS1 import. Consistent with previous reports (Zhang et al. 2010a,b), wild-type LDs are usually below 3 μm in diameter at stage L4 and display no obvious size difference when grown continuously at 15°C, 20°C, or shifted to 30°C for a brief period of 4 hr (Figure 3, A–C). At a restrictive temperature of 20°C, the reference allele ssd72-ts of drop-5 accumulates supersized LDs > 3 μm in diameter and in only the intestinal cells throughout the anterior–posterior axis (Figure 4A). The only allele of drop-9, ssd213, is a non-ts mutant with intestinal supersized LDs formed at all three temperatures (Figure 4B; unpublished results). The acs-22 (tm3236) null mutation has no effect on wild-type LDs (Xu et al. 2012) but completely prevents supersized LD formation in drop-5 (ssd72-ts and drop-9(ssd213) (Figure 3D and Figure 4C and D). Supersized LDs that have already formed in drop-5(ssd72-ts and drop-9 (ssd213) are mostly resistant to hydrolysis induced by a 24 hr-fasting
post L4 while LDs in wild type are not (Figure 3, E and F). drop-3(ssd72)p was isolated in the EMS/ls[mdm]p series and was noted for its ts peroxisome import defect: peroxisome matrix protein GFP::DAF-22 and mRFP::PTS1 are correctly imported at 15° but not at 20° (Figure 4, G–I). Similarly, mRFP::PTS1 is not imported in drop-9(ssd213) (Figure 4f). Thus, drop-5 and drop-9 are similar to maoc-1/dls-28/daf-22/prx-10 in supersized LD phenotype, ACS-22-dependence, and resistance to fasting-induced hydrolysis, and similar to prx-10 in peroxisome import defect (Zhang et al. 2010a; Xu et al. 2012). It strongly suggests that drop-5 and drop-9 supersized LDs form by an enhancement of LD growth and an inhibition of LD hydrolysis. drop-5 and drop-9 may encode peroxins and are categorized as class I.

Class II (drop-2, drop-3, drop-6 and drop-7) is up-regulated in LD growth but not defective in peroxisome import

Class II mutants drop-2, drop-3, drop-6, and drop-7 are similar to class I in supersized LD phenotype, ACS-22/DGAT-2-dependence, and resistance to fasting-induced hydrolysis (Figure 5, A–L). But these four groups are not peroxisome import defective. Interesting differences exist between the four groups. All 22 alleles of drop-2 are non-ts and exhibit similarly supersized LDs. It is unlikely that these 22 alleles are a series of partial loss-of-function mutations since drop-2 (ssd29) is an amber mutant suppressible by sup-7(ts5). All five alleles of drop-3 and both alleles of drop-6 are ts with a restrictive temperature of 30°. Furthermore, the supersized LDs formed at 30° for 4 hr are only partially resistant to fasting-induced hydrolysis (Figure 5, K and L). The unusually high restrictive temperature and the partial resistance to fasting-induced hydrolysis may be due to the existence of drop-3 and drop-6 homologs. Consistent with this notion is that ssd75, the only drop-7 allele, is non-ts and is fully resistant to fasting-induced hydrolysis (Figure 5, B and F). drop-7(ssd75) appears mutated in two unlinked genes since its heterozygote segregates 1:1/16 (9/137) supersized LD progeny. However, drop-3 and drop-6 are unlikely to be homologous to each other since the double mutant drop-3(ssd36)p; drop-6(ssd73)p is not stronger than each single mutant in LD phenotype. With a similarity to maoc-1/dls-28/daf-22 in supersized LD phenotype and a lack of peroxisome import defect, drop-2, drop-3, drop-6, and drop-7 may encode additional enzymes of peroxisomal fatty acid β-oxidation and are thus categorized as class II.

Class III (drop-1 and drop-8) forms supersized LDs by fusion

All alleles of class III mutants drop-1 and drop-8 are ts with a restrictive temperature of 30° (Figure 6, A–D). This class differs drastically from class I and II mutants in three aspects. First, the supersized LD phenotype is not abolished by acs-22 mutation (Figure 6, E and F). Second, supersized LDs are completely exhausted by fasting-induced hydrolysis (Figure 6, G and H). Third, the formation of supersized LDs upon a shift to 30° for 4 hr is accompanied by an obvious decrease of small LDs, indicating a process of enhanced LD fusion (Figure 6, A–D). To directly test for LD fusion, we time-lapse imaged GFP::DGAT-2-labeled LDs in drop-1 upon an up-shift to 30°. More than 20 fusion events were recorded. As shown in one example, two LDs directly fuse during a short time window of 30 sec to form a supersized LD with a diameter > 5 μm (Figure 6f).

Class IV (drop-4) forms supersized LDs independent of ACS-22 and is hydrolysis-resistant

Class IV is defined by a single group with a single non-ts allele drop-4 (ssd206). The supersized LD phenotype of drop-4(ssd206) is already pronounced at L4 stage and increases to a phenomenal level at 2-d adult stage (Figure 7, A and B). The extremely large LDs resemble those in a previously reported mutant sams-1 (Li et al. 2011). drop-4(ssd206) does not complement a deletion mutant sams-1(ok2946) and bears a missense mutation Ala42Val at the SAMS-1 protein level. So, drop-4 is sams-1. Unlike in drop-1 and drop-8, the formation of supersized LDs in drop-4/sams-1 is unaccompanied by a detectable clearance of small LDs (Figure 7A), independent of ACS-22, and resistant to fasting-induced hydrolysis (Figure 7, C and D). These features make drop-4/sams-1 distinct from class I, II, and III.

Allele frequency, enrichment of novel genes, and a test of saturation

In the six series of genetic screens we isolated 13 complementation groups. Four are the same peroxisome mutants as previously found. Nine are novel. This collection of mutants showed a drastic difference in allele numbers. Four genes (dls-28, daf-22, drop-1 and drop-2) are defined by > 15 alleles; the other four (drop-4, drop-7, drop-8, and drop-9), by just one (Table 1). To estimate the screening efficiency, we calculated allele frequencies of these genes according to their isolation origins (Table 2).

We first compared the efficiency of the two mutagens. In the EMS/WT screen, allele frequencies of three known genes are comparable to those reported before: 2.5 × 10⁻⁵ vs. 5.6 × 10⁻⁵ of maoc-1, 7.6 × 10⁻⁵ vs. 11.1 × 10⁻⁵ of dls-28, and 15.3 × 10⁻⁵ vs. 11.1 × 10⁻⁵ of daf-22 (Butcher et al. 2009; Zhang et al. 2010a). ENU mutagenesis did not result in a higher allele frequency for these three genes. The cumulative allele frequency of all 13 genes with ENU is 22.5 × 10⁻⁵, lower than 43.5 × 10⁻⁵ with EMS. For every one of the nine genes (maoc-1, dls-28, daf-22, prx-10, drop-1, drop-2, drop-3, drop-5, and drop-9), the frequency with ENU is consistently lower than that with EMS and this...
Table 2 Allele frequencies with two different mutagens and three backgrounds

| Mutagen | EMS       | ENU       | WT       | is[mddm]  | acs-22    |
|---------|-----------|-----------|----------|-----------|-----------|
| maoc-1 II | $2.5 \times 10^{-5}$a | $1.6 \times 10^{-5}$a | $2.0 \times 10^{-5}$ | $6.1 \times 10^{-5}$ | $1.6 \times 10^{-5}$ |
| dhs-28 X | $7.6 \times 10^{-5}$a | $4.7 \times 10^{-5}$a | $6.1 \times 10^{-5}$ | $9.4 \times 10^{-5}$ | $8.4 \times 10^{-5}$ |
| daf-22 II | $15.3 \times 10^{-5}$a | $3.9 \times 10^{-5}$a | $1.2 \times 10^{-5}$ | $3.1 \times 10^{-5}$ | $0.8 \times 10^{-5}$ |
| prx-10 III | $0.9 \times 10^{-5}$b | $0.3 \times 10^{-5}$b | $0.4 \times 10^{-5}$ | $0.4 \times 10^{-5}$ | $0.3 \times 10^{-5}$ |
| drop-1 II | $6.4 \times 10^{-5}$c | $4.9 \times 10^{-5}$c | $3.7 \times 10^{-5}$ | $8.3 \times 10^{-5}$ | $2.8 \times 10^{-5}$ |
| drop-2 IV | $8.0 \times 10^{-5}$c | $3.1 \times 10^{-5}$c | $4.1 \times 10^{-5}$ | $9.4 \times 10^{-5}$ | $1.6 \times 10^{-5}$ |
| drop-3 IV | $1.4 \times 10^{-5}$c | $1.3 \times 10^{-5}$c | $1.2 \times 10^{-5}$ | $3.1 \times 10^{-5}$ | $0.0 \times 10^{-5}$ |
| drop-4 X | $0.9 \times 10^{-5}$c | $0.3 \times 10^{-5}$c | $0.4 \times 10^{-5}$ | $0.4 \times 10^{-5}$ | $0.3 \times 10^{-5}$ |
| drop-5 X | $1.3 \times 10^{-5}$c | $0.6 \times 10^{-5}$c | $0.8 \times 10^{-5}$ | $0.8 \times 10^{-5}$ | $0.8 \times 10^{-5}$ |
| drop-6 X | $0.9 \times 10^{-5}$c | $0.8 \times 10^{-5}$c | $0.9 \times 10^{-5}$ | $0.9 \times 10^{-5}$ | $0.9 \times 10^{-5}$ |
| drop-7 | $0.5 \times 10^{-5}$c | $0.3 \times 10^{-5}$c | $0.3 \times 10^{-5}$ | $0.3 \times 10^{-5}$ | $0.3 \times 10^{-5}$ |
| drop-8 II | $0.5 \times 10^{-5}$c | $0.3 \times 10^{-5}$c | $0.3 \times 10^{-5}$ | $0.3 \times 10^{-5}$ | $0.3 \times 10^{-5}$ |
| Total | $43.5 \times 10^{-5}$ | $22.5 \times 10^{-5}$ | $28.5 \times 10^{-5}$ | $16.5 \times 10^{-5}$ | $9.0 \times 10^{-5}$ |

a The number of alleles divided by the number of haploid genomes of the WT screen since Is[mddm] transgene bypasses the selection of and acs-22 suppresses these mutants. In a similar EMS/WT screen in Zhang et al. (2010) and Butcher et al. (2009), allele frequency of maoc-1 is $5.6 \times 10^{-5}$; dhs-28, 11.1 $\times 10^{-5}$; daf-22, 11.1 $\times 10^{-5}$; prx-10, 2.7 $\times 10^{-5}$.

b The number of alleles divided by the sum of haploid genomes of the WT and is[mddm] screens since acs-22 suppresses these mutants.

c The number of alleles divided by the sum of haploid genomes of the WT, Is[mddm], and acs-22 screens since acs-22 does not suppress these mutants.

DISCUSSION

The fact that TAG is stored in LDs in C. elegans was not established until Zhang and colleagues analyzed wild type, glo mutants, and a class of peroxisomal β-oxidation mutants. They found that: (1) C. elegans LDs as visualized by transmission electron microscopy are electron transparent structures delimited by a phospholipid monolayer; (2) LDs are not abolished in glo mutants and supersized LDs form in peroxisomal β-oxidation mutants; (3) in wild type, glo mutants, and peroxisomal β-oxidation mutants, small and supersized LDs can be labeled both by postfix Oil-Red-O and by postfix Nile Red staining; (4) postfix Nile Red labels LDs and emits fluorescence with a true color of gold; (5) vital Nile Red only labels LROs with a red fluorescence in wild type but labels both LROs and LDs in peroxisomal β-oxidation mutants; (6) vital BODIPY labels LDs in addition to LROs with the same emission fluorescence; and (7) vital BODIPY-labeled LDs can be separated from LROs by density centrifugation-based purification or by genetically ablating LROs with the glo-4 mutation (Zhang et al. 2010a,b). These findings explained previous intriguing results that postfix Nile Red staining and postfix Oil-Red-O staining serve as proxies for relative TAG level better than vital Nile Red does (Hermann et al. 2005; Schroeder et al. 2007; Brooks et al. 2009; O’Rourke et al. 2009), and identified C. elegans fat storage organelles as LDs instead of “vesicles distinct from LROs.”

Postfix Nile Red staining and Oil-Red-O staining, however, can only serve as a semiquantitative measurement for TAG levels for the following reasons. First, Nile Red fluorescence and Oil-Red-O staining intensity are difficult to normalize against body size or protein mass, which usually differ between wild type and various mutants. Second, staining intensities and image acquisition vary to some extent between individuals and between staining trials of the same genotype (this study and our unpublished observations) (Brooks et al. 2009; O’Rourke et al. 2009).
biased forward genetic screen has been reported. Small-scale candidate transgenic marker, post demonstrates the reliability and efficacy of forward genetic screens for LD mutants, and the success of isolating with a much stronger intensity (Zhang et al. 2010b). Interestingly, several mutants with enlarged LDs in oocytes and embryos were identified by DIC microscopy screening of a collection of candidate lethal mutants and animals subjected to RNAi of a small pool of candidate genes. But these mutants did not form supersized LDs at larval or adult stage (Schmokel et al. 2016). In contrast, the vital BODIPY-based large-scale forward genetic screen proved successful in isolating supersized LD mutants here and in our previous, smaller scale study (Zhang et al. 2010a).

Our study was successful in realizing two aims: to further address C. elegans LD labeling methodology and to explore the full potential of C. elegans forward genetic screens to establish models of LD size regulation. When incorporated into supersized LDs, BODIPY emits a strong enough fluorescence which can be readily visualized under a stereo fluorescence microscope, allowing us to rapidly screen millions of F2 animals descended from 0.67 million mutagenized haplod genomes. A complication of vital BODIPY is that it labels LROs in addition to LDs. This problem was obviated by introducing the glo-4 mutation to eliminate LROs (Figure S1 and Figure S2) (Schroeder et al. 2007; Zhang et al. 2010b). Even in the wild-type background, the presence of LROs did not pose a problem since we looked for a phenotype of size increase and supersized LRO mutants were not found in this study or in a previous direct RNAi screen of LRO phenotype (Soukas et al. 2013). As such, our forward genetic screen for supersized BODIPY-positive structures isolated 118 mutants that were all confirmed as supersized LD mutants by postfix Nile Red staining, postfix Oil-Red-O staining, and GFP::DGAT-2 marking (Figure 1). The BODIPY screen allowed the isolation of four mutant complementation groups reported before, and more importantly, nine novel complementation groups representing four models of LD regulation. One group drop-4 was cloned as sans-1, a supersized LD mutant that was originally identified not by direct LD phenotype screening but by candidate gene approaches based on proteomics and gene expression data (Li et al. 2011; Walker et al. 2011; Ehmke et al. 2014). The success of our forward genetic screen justifies the BODIPY approach. We note that the BODIPY approach may not reveal subtle LD phenotypes such as size decrease, small change in number, and change in distribution pattern, for which screening with transgenic markers or postfix staining might also be required. Nevertheless, we suggest that the supersized LD phenotype may serve as a benchmark for testing the reliability and efficacy of any LD labeling and imaging methods in genetic screens.

Have we explored the full potential of the current BODIPY-based forward genetic screens? We think so for the following considerations. The ease with vital BODIPY made it possible to screen a large number of 6.7·10⁵ mutagenized haplod genomes. The use of a novel temperature regimen led to the identification of four genes (drop-1, 3, 6, and 8) that otherwise would have escaped isolation. The use of a gene-class-bypassing strategy permitted the identification of two additional genes (drop-4 and 7). The use of ENU, a mutagen with a broader mutagenic spectrum (Anderson 1995; De Stasio et al. 1997), allowed the identification of four genes (drop-4, 6, 7, and 8). In total, we identified 118 bona fide supersized LD mutants, of which 73 are mutations in nine new genes (Table 1 and Table 2). How frequent is the occurrence of supersized LD mutants? The total allele frequency induced by EMS is 43.5·10⁻⁴, 4.8·10⁻⁵ per gene, 10 times lower than 5·10⁻⁴ of an average gene (Brenner 1974). The average allele frequency of maoc-1/dhs-28/daf-22 is 6.6·10⁻⁵, comparable to 7.6·10⁻⁵ in a previous report (Zhang et al. 2010a). This suggests that the average allele frequency of supersized LD mutants is inherently low. We used the Poisson equation to estimate whether our screen system was saturated, i.e., whether additional genes were unidentified. As shown in Table 3, in the WT series alone, only 0.009 gene was unidentified, and in screens against all three backgrounds together, 0.001. However, it does not necessarily

### Table 3 The extent of saturation for genes that can be mutated to a fertile supersized LD phenotype

| Series       | WT    | EMS/WT | ENU/WT | WT + Is/mdm/acs-22 |
|--------------|-------|--------|--------|-------------------|
| maoc-1 II    | 5     | 3      | 2      | 5                 |
| dhs-28 X     | 15    | 9      | 6      | 15                |
| daf-22 II    | 23    | 18     | 5      | 23                |
| pxe-10 III   | 2     | 2      | 2      | 2                 |
| drop-1 II    | 9     | 5      | 4      | 38                |
| drop-2 IV    | 10    | 5      | 5      | 22                |
| drop-3 IV    | 3     | 3      | 5      | 5                 |
| drop-4 X     | 1     | 1      | 1      | 1                 |
| drop-5 X     | 1     | 1      | 1      | 1                 |
| drop-6 X     | 2     | 2      | 2      | 2                 |
| drop-7 X     | 1     | 1      | 1      | 1                 |
| drop-8 III   | 1     | 1      | 1      | 1                 |
| drop-9 X     | 1     | 1      | 1      | 1                 |

Allele numbers are listed for each gene according to isolation origins. A, mean allele number; P(0), probability of unidentifed genes; N(0), number of unidentified genes.

- Assume that the allele frequency observes Poisson distribution. The probability of unidentified genes, i.e., the category with 0 allele, follows P(k) = λ^k * e^-λ/k!.
- When λ = 0, P(0) = e^-λ.
- N(0) = N * e^(-N/e^λ). N, number of identified genes.

LS(0) = 0.0009 0.0036 0.0281 0.0001

LS(1) = 0.009 0.029 0.202 0.001

LS(2) = 9.0 3.6 5.6 7.0

Typical of phospholipids in addition to the gold fluorescence of LDs (Figure 1D) (Greenspan et al. 1985; Zhang et al. 2010b), making indiscriminate quantification of fluorescence intensity error-prone for TAG levels. The application of vital BODIPY fluorescence intensity to TAG quantification suffers similar problems: in addition to labeling intestinal LDs, BODIPY labels intestinal LROs and hypodermal LDs with a much stronger intensity (Zhang et al. 2010a,b). Thus, while these staining methods are well suited for qualitative LD labeling, a precise quantification of TAG in C. elegans depends on quantitative lipid analytical chemistry with a normalization against total lipid/phospholipid mass or total protein mass (Schroeder et al. 2007; Brooks et al. 2009; O’Rourke et al. 2009; Zhang et al. 2010a).

Although qualitatively specific for LDs, postfix Nile Red and Oil-Red-O are too labor-intensive for large-scale genetic screens. Most transgenic LD protein markers such as ATGLL1::GFP, GFP::DGAT-2, DHS-3::GFP, R01B10.6::GFP; PLIN1::GFP, and DHS-4::GFP are functional lipid metabolism proteins whose transgenic expression levels should be kept low so as not to complicate the LD phenotype (Zhang et al. 2010a; Xu et al. 2012; Zhang et al. 2012; Klemm et al. 2013; Liu et al. 2014; Vrablik et al. 2015). Thus, transgenic LD markers necessitate high-power microscopy which is also labor-intensive. Raman scattering imaging has the same practical limitation in addition to the problem of segregating TAG signal from protein and phospholipid signals (Hellerer et al. 2007; Wang et al. 2014). The reliability and efficacy of labeling methods are key to the applicability of C. elegans to large-scale forward genetic screens for LD mutants, and the success of isolating forward genetic mutants with bona fide mutant LD phenotypes demonstrates the reliability and efficacy of these methods – to date, apart from a forward screen of an unknown scale based on R01B10.6::GFP which identified a mini-sized LD mutant (Klemm et al. 2013), no transgenic marker, postfix staining, or Raman microscopy-based unbiased forward genetic screen has been reported. Small-scale candidate gene and RNAi screens revealed no known or new supersized LD mutants (Liu et al. 2014). Interestingly, several mutants with enlarged
mean that new genes will not be discovered. For instance, the ENU/WT screen identified drop-6 and drop-7 which were predicted nonexistent by Poisson analysis of the EMS/WT screen; the EMS/WT screen identified drop-3, drop-5, and drop-9 predicted nonexistent by the ENU/WT screen; and the acs-22 screen identified drop-4 and drop-8 predicted nonexistent in the WT screen. drop-4, drop-5, drop-7, drop-8, and drop-9 are of very low allele frequency with just one or two alleles. Interestingly, these alleles of drop-4, drop-5, drop-7, and drop-8 have a brood size significantly lower than wild type and other drop mutants (Figure S4) due to either sterility or embryonic/larval lethality. These alleles may be rare partial loss-of-function mutations. Thus, it seems unlikely that major complementation groups with high allele frequency were missed in our system. Any missing ones may involve sterility and or lethality genes, which may be discovered by screening F2 progeny of cloned F1s and by using genetic balancers.

We categorized the nine drop genes into four classes according to their effect on peroxisome matrix protein import, dependence on ACS-22-DGAT-2-mediated LD growth, and occurrence of direct LD fusion. The rate-limiting enzyme complex ACS-22/FATP1-DGAT-2 together with glycerol-3-phosphate O-acyltransferase 4 (GPAT4) synthesizes TAG on the surface of LDs and TAG is directly packaged into existing LDs, leading to an enhancement of LD growth and supersized LD formation in C. elegans, Drosophila, and mammalian cells (Harris et al. 2011; Xu et al. 2012; Wilfling et al. 2013). Similar to peroxisomal fatty acid β-oxidation mutants maao-1/hdls-28(daf-22), class I and class II mutants are ACS-22-DGAT-2-dependent and hydrolysis-resistant. Class I and class II thus represent a genetic model of enhanced LD growth and inhibited LD hydrolysis. Class I drop-5 and drop-9 likely encode PRX proteins involved in peroxisome import. Class II drop-2, drop-3, drop-6, and drop-7 may encode peroxisomal fatty acid β-oxidation enzymes not yet identified in C. elegans (Zhang et al. 2010a). Class III drop-1 and drop-8, being ACS-22-independent and hydrolysis-sensitive, are apt to represent enhanced LD fusion. LD fusion in C. elegans has never been convincingly reported before. Our model of LD fusion is particularly useful because it is controllable by a temperature up-shift and the process is fast, resembling “typical LD fusion” rather than “atypical LD fusion” in yeasts, Drosophila, and mammalian cells (Fei et al. 2011; Gong et al. 2011; Krahmer et al. 2011). It is interesting that a restrictive temperature of 30°C is required for triggering the fusion process even in drop-1 null alleles (unpublished results), suggesting that LD fusion may be a temperature-sensitive process with unusual thermodynamics. The factors contributing to this unusual thermodynamics may include phospholipid composition, fatty acid chain length, degree of unsaturation, and branching in the monolayered membrane of LDs (Fei et al. 2011; de Mendoza 2014). In natural environments, C. elegans frequently encounters extreme temperatures which have been shown to play important roles in several physiological processes including dauer formation, synaptic vesicle endocytosis, and temperature adaptation (Golden and Riddle 1984; Sato et al. 2009; Svensk et al. 2013; Ma et al. 2015). Further studies of our model may yield insights into regulatory mechanisms of LD fusion. Class IV drop-4/sams-1 is ACS-22-independent and hydrolysis-resistant. It was thought that through up-regulating the expression of ACS-22, DGAT-2, fatty acid synthase, desaturases, and elongases, sams-1 mutation decreases phosphatidylcholine (PC) synthesis and increases TAG synthesis (Li et al. 2011; Walker et al. 2011; Ehmeke et al. 2014). But it is still poorly understood why TAG increase results in LD size increase but not number increase. The ACS-22-dependence of drop-4/sams-1 indicates that supersized LDs do not form by enhanced LD growth. However, a possibility of ACS-22-independent LD growth cannot be excluded and evidence for enhanced LD fusion is lacking. Detailed molecular analysis of these four classes of genes in LD size regulation and fat metabolism will be published elsewhere.

ACKNOWLEDGMENTS

We thank Drs. David Weisblat, Gian Garriga, Yikun He, Peng Li, and Xiao Liu for critical reading of and constructive comments on this paper. Some C. elegans strains were kindly provided by Dr. Ho Yi Mak. Some other strains were provided by the Caenorhabditis Genetics Center, which is funded by the National Institutes of Health (NIH) Office of Research Infrastructure Programs (P40 OD010440), and by the MITANI Laboratory through the National Bio-Resource Project of the MEXT, Japan. S.L. was supported in part by an International Exchange and Joint Training Grant of the Graduate School of Capital Normal University (CNU). This work was supported by a startup fund from CNU, a grant from the National Natural Science Foundation of China (grant no. 31370820), and a grant from the Beijing Municipal Education Commission (grant no. KZZ201410028032) to S.O.Z.

Author contributions: S.L. and S.O.Z. designed the experiments; S.L., S.X., Y.M., S.W., Y.F., Q.C., L.C., Z.S., Y.K., X.Z., J.Y., M.W., and S.O.Z. performed the experiments; S.L. and S.O.Z. analyzed the data and wrote the manuscript. All authors read the final version of the manuscript.

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Communicating editor: D. S. Fay