Peroxisome Protein Transportation Affects Metabolism of Branched-Chain Fatty Acids That Critically Impact Growth and Development of C. elegans

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Abstract

The impact of specific lipid molecules, including fatty acid variants, on cellular and developmental regulation is an important research subject that remains under studied. Monomethyl branched-chain fatty acids (mmBCFAs) are commonly present in multiple organisms including mammals, however our understanding of mmBCFA functions is very limited. C. elegans has been the premier model system to study the functions of mmBCFAs and their derived lipids, as mmBCFAs have been shown to play essential roles in post-embryonic development in this organism. To understand more about the metabolism of mmBCFAs in C. elegans, we performed a genetic screen for suppressors of the L1 developmental arrest phenotype caused by mmBCFA depletion. Extensive characterization of one suppressor mutation identified prx-5, which encodes an ortholog of the human receptor for the type-1 peroxisomal targeting signal protein. Our study showed that inactivating prx-5 function compromised the peroxisome protein import, resulting in an increased level of branched-chain fatty acid C17ISO in animals lacking normal mmBCFA synthesis, thereby restoring wild-type growth and development. This work reveals a novel connection between peroxisomal functions and mmBCFA metabolism.

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Introduction

Lipids are known for their basic functions in energy storage, structural support of cell membranes, and adaptation to different environmental conditions [1,2]. Extensive research has indicated that they also play critical signaling and regulatory roles in various cellular and developmental events [3-8]. Like proteins and noncoding RNAs, lipid molecules are extremely diverse in their structures, but the functional specificities associated with these structural variations are poorly understood. Fatty acids (FAs), the basic components of lipids, are highly variable in their structure with varying carbon chain length, straight- or branched-chain, and saturated or unsaturated characteristics [9,10]. In particular, very little is known about the roles of monomethyl branched-chain fatty acids (mmBCFAs) that are present in animals including mammals [11-13].

mmBCFAs C15ISO and C17ISO are saturated tetradecanoic and hexadecanoic fatty acids, respectively, that have a single methyl group added on the carbon next to the terminal carbon (Figure 1A). ELO-5 is a FA elongation enzyme, or elongase, specifically required for biosynthesis of mmBCFAs in C. elegans; disrupting elo-5 function by a deletion mutation or by RNAi treatment causes dramatic and specific reduction of the levels of C15ISO and C17ISO in C. elegans (in the absence of dietary mmBCFAs) [11]. Eliminating elo-5 function causes several developmental defects including robust developmental arrest at the first larval stage (L1 arrest), all of which can be overcome by dietary supplementation of chemically synthesized C15ISO or C17ISO (Figure 1B-D) [11,14]. Further study showed that mmBCFAs are incorporated into specific phospholipids by an acyl-CoA synthetase (ACS-5), which has a profound influence on IP3 signaling and membrane dynamics during early embryogenesis [15]. Recent work from our lab discovered that an mmBCFA-containing sphingolipid, d17iso-glucosylceramide, mediates the role of mmBCFAs in promoting postembryonic growth and development by activating the TORC1 signaling pathway in the intestine [16].

The peroxisome is an important organelle in all eukaryotic cells. It contains enzymes that are essential for multiple...
Figure 1. The growth defect of elo-5(If) is suppressed by mutations in prx-5 and prx-6. (A) Structure of mmBCFA C17ISO (15-methyl hexadecanoic acid) and C15ISO (13-methyl tetradecanoic acid). ELO-5 and ELO-6 are responsible for synthesis of C15ISO and C17ISO. (B-E) Microscopic images of C. elegans of indicated genotypes and treatments. Unlike WT (B) elo-5(If) mutants depleted for mmBCFA after hatching display a robust L1 growth arrest phenotype (indicated by arrows) (C) that can be overcome by dietary supplementation of C17ISO (D). The growth arrest phenotype of elo-5(If) is suppressed by the ku517 allele (E). (F) Schematic representation of the suppressor screen. The extrachromosomal array is composed of two transgenic markers: GFP and rol-6 (dr), but only GFP is shown in the cartoon for simplicity. (G) Bar graph showing percentage of animals that reached adulthood for strains with indicated genotypes. Three loss-of-function (If) mutations in the prx-5 and prx-6 genes significantly suppressed the L1 arrest phenotype of elo-5(If).

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metabolic pathways, including amino acid metabolism, hydrogen peroxide regulation and lipid metabolism. Peroxisomal proteins are synthesized in the cytosol and require an import mechanism to execute their metabolic functions in the peroxisome. The peroxisomal import mechanism involves multiple proteins, including the peroxisomal membrane proteins and PTS receptor. In human, receptor PEX5, which has an ortholog PRX-5 in C. elegans, interacts directly with proteins containing the type 1 peroxisomal targeting signal (PTS-1) [17]. Peroxisomal matrix proteins containing PTS-1 on their C-terminus are synthesized on free polyribosomes in the cytosol, and then recognized by PEX5 through its interaction with the PTS-1. PEX5 then transports proteins with PTS-1 into the peroxisome matrix [18]. The peroxisomal import mechanism is essential for peroxisomal functions as it has been shown that inactivating the peroxisomal import mechanism compromises peroxisomal metabolism and causes accumulation of metabolites in humans [19].

In this study, we identified a connection between mmBCFA metabolism and peroxisomal protein transport through a search for suppressor mutations of the L1 arrest phenotype caused by a loss-of-function (lf) mutation in elo-5. Here we describe the identification and characterization of mutations in prx-5 that disrupt the peroxisomal import mechanism and alter mmBCFA metabolism in C. elegans. This change, mostly likely through reducing peroxisome-involved degradation of mmBCFAs, is able to compensate for the loss of the major C17ISO biosynthesis enzyme.

Materials and Methods

General C. elegans maintenance and strains

All C. elegans strains were maintained at 20°C on nematode growth media (NGM) with OP 50 bacteria according to the standard protocol. The wild-type strain used was the N2 Bristol. The following mutant strains were used in this study: elo-5(gk208)V, prx-5(ka517)II, prx-5(tm4948)II, prx-5(ok984)IV, dpy-10(e128); unc-4(e120)II, maoc-1(ok2645)II, dhs-28(tm2581)X and daf-22(ok693)III. elo-5(gk208) was originally obtained from the C. elegans Gene Knockout Consortium. The prx-5(tm4948)II and dhs-28(tm2581)X deletion strains were obtained from the National Bioresource Project in Japan. The Hawaiian strain (CB4856) was used for SNP mapping [20]. The strain containing [P:GFP-SKL] was kindly provided by M. Driscoll. The elo-5(gk208) strain and mutants containing the elo-5(gk208) allele were maintained on OP 50 bacteria supplemented with the mmBCFA-producing *Stenotrophomonas maltophilia* bacteria.

elo-5(gk208) suppressor screen

An elo-5(gk208) mutant containing an elo5(+) rescuing transgene and visible markers, [elo-5(gk208);Ex[elo-5(+);rol-6 (su1006)]II, Pelo-5:GFP], was first constructed for the screen. L4-staged worms (generation P0) grown in the presence of 1 mM C13ISO were mutagenized by EMS. Their first and second generation progeny, F1 and F2, were maintained as a group until the F2 animals grew to the L4 stage. Among F2 animals, non-roller and GFP-negative worms, presumably not containing...
the rescuing transgene, were singled out and plated on OP 50 plates without mmBCFA supplementation. Suppressor candidates were determined by the ability of their progeny to survive on these plates. From ~9000 haploid genomes, four suppressor mutants were isolated. Mapping analyses indicated that they are mutations in at least three different genes. ku517 was mapped to chromosomal II and outcrossed five times before further analysis.

**Gas chromatography (GC) to analyze fatty acid composition**

A mixed population of worms was collected from plates with M9 buffer, rinsed 3-4 times, pelleted, freeze-dried, and weighed, in preparation for lipid extraction. Collected samples were mixed with methanol with 2M KOH for 5 min at room temperature for methylation; FA methyl esters were extracted using hexane and subjected to GC analysis as described previously [11]. The abundance of mmBCFAs was analyzed by calculating the area of their peaks on the GC profile. For each sample, three replicates were assayed. The average and standard deviation (s.d.) were calculated using Excel. Statistical significance was determined using the student's T-test.

**Position cloning of prx-5**

A SNP mapping method [21] was used to locate ku517 to the middle of chromosome II. The mutation was further mapped to a small region between dpy-10 and unc-4 using the three-point mapping method (Figure 2A). Recombinants were generated by crossing ku517; elo-5(gk208) to the dpy-10(e128); unc-4(e120); elo-5(gk208) strain. In F3 worms, Unc non-Dpy and Dpy non-Unc recombinants were isolated. Of these recombinants, 13/16 Unc non-Dpy and 1/8 Dpy non-Unc recombinants had the ku517 suppressor mutation.

By applying whole genome sequencing [22] and subsequent data analysis [23], multiple DNA lesions were identified within the region. Between dpy-10 and unc-4, only four homozygous DNA lesions in the coding region were identified and only a C to T lesion in the coding region of the prx-5 gene is consistent with the three-point mapping data. In addition, only this prx-5 mutation is present in the ku517; elo-5(gk208) double mutant but not in the elo-5(gk208) single mutant. This substitution is predicted to change a codon for sur-5::GFP -3'. Expression construct of prx-5 was generated by PCR amplification of the genomic fragment of prx-5, including 3.1kb upstream of initiation codon of prx-5 and 0.4kb downstream of its stop codon. Primers: F – BamHI 5' - GAGGGGCTCAAGGAGTTGGTC -3', R - KpnI 5' - TCTGTAAAAATTTGAAGAATTCGAGAG -3'. The rescue construct of prx-5 was generated by PCR amplification of the genomic fragment of prx-5, following the procedures described in [17].

**DNA constructs for transgenes**

The prx-5::GFP transcriptional reporter construct was generated by PCR amplification of 3 kb upstream of the prx-5 initiation codon from N2 genomic DNA and cloned into GFP vector pPD95.77 (gift from A. Fire). Primers: F- BamHI 5'-GAGGGGCTCAAGGAGTTGGTC -3'; R - KpnI 5' - TCTGTAAAAATTTGAAGAATTCGAGAG -3'. The rescue construct of prx-5 was generated by PCR amplification of the genomic fragment of prx-5, following the procedures described in [17].

**Microscopy**

Fluorescent and Nomarski optics of a Zeiss Axioplan2 microscope and Zeiss AxioCam MRm camera were used to analyze GFP expression and abnormalities in phenotypes. Phenotypes on plates were observed through a Leica MZ16F dissecting microscope and pictures were taken with a C4742-95 CCD camera.

**Examining mRNA level by real-time quantitative PCR (qPCR)**

Total RNA was isolated from asynchronous, well-fed worms with TRizol reagent (Invitrogen) following the manufacturer’s procedure. Total RNA was reverse transcribed with an oligo-dT primer, and subjected to quantitative PCR using Rotor-Gene RG-3000 (Qiagen). All qPCR data was normalized to the average Ct of a housekeeping gene, rpl-26.

**Results**

Isolation of suppressor mutations of the growth defect associated with elo-5(f)

Previous work in our lab has shown that mmBCFAs C15ISO and C17ISO are required for normal C. elegans development. When the fatty acid elongase gene elo-5 is mutated in parental animals, the resulting hatched worms uniformly arrest at the
first larval (L1) stage. This robust larval growth and developmental defect can be overcome by dietary supplementation of mmBCFAs [11,14] (Figure 1A-D). To identify genes that may be involved in the metabolism of mmBCFAs or in a cellular pathway regulating mmBCFA-mediated functions in development, we performed a genetic

Figure 2. Positional cloning and analysis of prx-5 defined by the ku517 mutation. (A) Schematic representation of 2 centimorgan (cM) region of chromosome II containing the ku517 mutation. The position of ku517 relative to the dpy-10 and unc-4 genes was determined by three-point mapping. The location of a 7 kb genomic DNA clone able to rescue the ku517 allele (3 out of 3 transgenic lines) is also indicated. (B) Structure of the prx-5 gene. There are two potential cDNA isoforms differing by 6 nucleotides (wormbase.org). The C to T substitution in prx-5(ku517) results in a premature stop codon terminating the encoded protein at amino acid residue 475. The double-arrow indicates the 437 bp deletion in prx-5(tm4948) allele. (C-F) Fluorescence images showing the broad expression pattern of a P<sub>prx-5</sub>:GFP fusion protein in C. elegans. The GFP reporter was observed in the embryo (C), intestine (D), hypodermis (E) and neurons (F). (G-J) Images showing the distribution of the GFP-SKL reporter that is the readout of peroxisomal import activity [17]. GFP was localized in peroxisomes as indicated by the punctate pattern in both wild type and elo-5(1f) mutants (G and H). In prx-5(tm4948), GFP is dispersed throughout the cytoplasm, indicating a defect in peroxisomal import (I and J). Bars, 25 μm.

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suppressor screen, modified from our previous study [20]. We looked for mutations that would rescue the growth of animals containing a loss-of-function (lf) mutation as a result of deletion in elo-5 gene, gk208, without mmBCFA supplementation. Hereafter, elo-5(gk208) is referred to as elo-5(lf).

For this screen, we generated a strain that was homozygous for elo-5(lf) and carried an extrachromosomal array containing copies of wild-type elo-5 and two markers (Figure 1F; Materials and Methods). Progeny that did not inherit the extrachromosomal array and were maintained without C17ISO supplement uniformly arrested their development at L1 stage. The worms carrying the array were mutagenized by EMS and their F2 progeny that did not contain the extrachromosomal array but developed to fertile adults were singled out. These animals potentially contained suppressor mutations that permitted the elo-5(lf) animals to overcome the L1-arrest phenotype (Figure 1F). After screening ~9000 haploid genomes, we isolated 4 alleles, among which ku517 was extensively characterized. The ku517; elo-5(lf) double mutant was able to grow indefinitely without dietary supplementation of C17ISO (Figure 1E and 1G). Thus, ku517 may carry a mutation that alters either mmBCFA metabolism, or a regulatory pathway, that bypasses the requirement for these mmBCFAs in development.

**ku517 is a loss-of-function mutation in prx-5, an ortholog of the human receptor for the type-1 peroxisomal targeting signal proteins**

Using single nucleotide polymorphism (SNP) mapping [21], conventional three-point mapping methods and the whole genome sequence analysis, we determined that ku517 is a C-T substitution that changes CGA coding for Arginine 475 to a CAA coding for Lysine 475. Since this amino acid change is hypothesized to alter peroxisomal import function due to the ineffectiveness of RNAi on this gene. The worms carrying the array were mutagenized by EMS and their F2 progeny that did not contain the extrachromosomal array and were maintained without C17ISO (Figure 1E and 1G). Thus, ku517 may carry a mutation that alters either mmBCFA metabolism, or a regulatory pathway, that bypasses the requirement for these mmBCFAs in development.

To examine the expression pattern of prx-5, we generated a transcriptional GFP reporter (Materials and Methods) and observed that the Pprx-5::GFP is expressed at all developmental stages in many tissues including the intestine, hypodermis and neurons (Figure 2C-F).

**prx-5(lf) mutation compromises peroxisomal import function**

To examine the impact of prx-5(lf) on peroxisomal import function, we analyzed the distribution of a reporter protein containing a GFP fused with a type-1 peroxisomal targeting signal SKL (Serine-Lysine-Leucine) at the C-terminus (GFP-SKL), which interacts directly with receptor PRX-5 [17]. This reporter was expressed in a transgenic strain carrying the P_HSP16/2::GFP-SKL construct behind a heat-shock promoter (Figure 2G-J). After a heat-shock and recovery, GFP-SKL was found to be imported from the cytosol into peroxisomes showing clear, punctate fluorescence in multiple tissues in wild-type and elo-5(lf) animals (Figure 2G-H). In contrast, the GFP signal lost the punctate pattern and was distributed throughout the cytoplasm in prx-5(tm4948) and prx-5(ok984); elo-5(lf) mutants (Figure 2I-J). This observation indicated that peroxisomal proteins failed to be imported into peroxisomes and were thus retained in the cytoplasm.

This defect appears to be more severe than that from the reported RNAi analysis [17], supporting our suggestion that prx-5(RNAi) has a weak loss of function effect. The results from both studies support the conclusion that prx-5 is required for peroxisomal import of peroxisomal matrix protein containing PTS-1 signal in *C. elegans*.

**Compromising peroxisomal import functions changes mmBCFA metabolism**

Because prx-5(lf) mutations disrupt peroxisome transport that is critical for basic peroxisomal functions, we asked whether suppression of the L1 arrest phenotype of elo-5(lf) is a result of interrupting general peroxisomal function rather than an undefined specific function of prx-5. We thus tested the effect of inactivating prx-6 in elo-5(lf) mutants. PRX-6 is predicted to be an ortholog of PEX6 in human, which is an AAA-ATPase responsible for recycling the peroxisomal matrix protein receptor back into the cytosol [27]. We found that a deletion allele of prx-6, prx-6(ok984), also suppressed the L1 arrest phenotype of elo-5(lf). About 60% of prx-6(ok984); elo-5(lf) double mutants were able to grow to fertile adults without dietary supplementation of C17ISO (Figure 1G). The suppression level is comparable to that of prx-5(ku517); elo-5(lf) and prx-5(tm4948); elo-5(lf). This result supports the idea that disrupting the functions of peroxisome import mechanism leads to the suppression of the elo-5(lf) phenotype. The absence of genetic mutations and ineffectiveness of RNAi prevented us from examining more peroxisomal genes.

To determine whether compromising peroxisomal function causes changes in mmBCFA metabolism or influences a signal pathway regulating mmBCFA-mediated functions in development, we examined the fatty acid composition of these suppressor mutants by gas chromatography (GC) (Materials and Methods). We found that both prx-5(ku517); elo-5(lf) and
prx-5 (tm4948); elo-5 (lf) double mutants had a higher level of mmBCFA C17ISO when compared to that in the elo-5 (lf) single mutant, albeit that the C17ISO level was still significantly below that observed in wild-type (Figure 3A). This result suggests that the suppression may be due to changes in mmBCFA metabolism.

Suppression of elo-5 (lf) by prx-5 (lf) depends on another fatty acid elongase, elo-6

Our previous studies showed that elo-6 is also involved in mmBCFA biosynthesis through its role in elongating C13ISO FA to C17ISO FA [11,14]. Therefore, it is possible that ELO-6 permits a low level of synthesis of C17ISO in the absence of ELO-5. Although the ELO-6 dependent synthesis of C17ISO is not sufficient to rescue the developmental defects in elo-5 (lf), it could be sufficient in the prx-5 (lf); elo-5 (lf) double mutants. If that is the case, and if the suppression of the L1 arrest phenotype of elo-5 (lf) is mainly attributed to the increase in C17ISO level, then the suppression effect is expected to depend on intact ELO-6 activity. We thus applied elo-6 RNAi to prx-5 (ku517); elo-5 (lf) double mutants and observed that the suppressor effect of prx-5 (ku517) was largely eliminated. More than 99% of F1 progeny from elo-6 (RNAi) treated prx-5 (ku517); elo-5 (lf) double mutants were arrested at the first or second larval stages without dietary supplementation of C17ISO (Figure 3B-D). Furthermore, GC analysis showed that elo-6 (RNAi) caused a decrease of the C17ISO level in prx-5 (ku517); elo-5 (lf) double mutants (Figure 3A). These results indicate that disruption of peroxisomal import functions suppresses elo-5 (lf) growth defect by increasing the endogenous level of mmBCFA C17ISO. This may be achieved either by enhancing ELO-6-dependent biosynthesis of C17ISO or by decreasing the degradation of the ELO-6 dependent
C17ISO that may be synthesized at a low level in the elo-5(If) mutant.

**prx-5(If) likely decreases mmBCFA degradation**

The increased level of C17ISO in the prx-5(If); elo-5(If) double mutant, compared to the elo-5(If) single mutant, could conceivably be caused by an increase in FA synthesis. Logically, it may be due to either a general increase in the expression of FA synthesis enzymes or a specific increase in the expression of one FA synthesis enzyme that has biochemical function similar to ELO-5. We thus examined the mRNA level of several FA elongases involved in the elongation of fatty acids with different efficiency and specificity [6,11,28-30]. Using quantitative RT-PCR (qRT-PCR), we examined the mRNA level of elo-1, elo-2, elo-3, elo-5 and elo-6 genes in wild-type, elo-5(If), prx-5(ku517); elo-5(If), prx-5(tm4948); elo-5(If) and prx-6(ok1984); elo-5(If) animals. The data presented in Figure 4B indicate that the suppression mutations in prx-5 and prx-6 do not cause a significant and general increase in the mRNA levels of FA elongases. Interestingly, the mRNA level of elo-3 is significantly decreased in two of the three suppressor mutants we observed, and the two prx-5 mutants (ku517 and tm4948) behave differently. It is not clear what is the cause of the potential down-regulation of elo-3 by these peroxisome defective mutations. The difference between the two prx-5 mutations in this regard, possibly caused by differences in the genetic backgrounds of these two strains, does suggest that the change in elo-3 mRNA level is not specifically linked to the suppression effects of these mutations.

We found that, when compared to wild type, the mRNA level of elo-6 was significantly higher in the three double mutants, as well as in the elo-5(If) single mutant (Figure 4A). This increase in elo-5(If) is consistent with the previous finding that elo-6 may be subjected to feedback regulation by C17ISO [14]. Although the elo-6 mRNA level in three prx(If); elo-5(If) double mutants appeared to be higher than in the elo-5(If) single mutant, the difference was not statistically significant. In this analysis, we did not observe consistent changes in mRNA levels of other tested FA elongation enzyme genes (Figure 4B), consistent with the idea that the expression of FA elongases is not prominently affected by compromising peroxisomal import functions.

Since ELO-6 plays an essential role in synthesizing a low level of mmBCFA in prx-5(If) in the absence of ELO-5, it is conceivable that the increase in C17ISO level in the prx-5(If) mutant is due to increased elo-6 activity. If that were the case, over expressing elo-6 alone would likely to suppress elo-5(If) L1 arrest. We thus generated transgenic animals that carried an extrachromosomal array containing multiple copies of wild type elo-6 with its native promoter. By qPCR, we confirmed that the elo-6 transgene was transcribed at a level significantly higher than that in the suppressor mutants (Figure 4A). Expression of this transgene failed to overcome the growth defect of elo-5(If); 100% of elo-5(If) animals carrying this transgene were arrested at L1 stage without dietary supplementation of C17ISO (n>400). Therefore, the suppression of elo-5(If) by prx-5(If) is not likely due to an increased expression of elo-6.

We then asked whether a decrease in mmBCFA degradation could be the key cause of the suppression effect of prx-5(If). β-oxidation of fatty acids is one of the many conserved metabolic functions associated with the peroxisome [31]. In *C. elegans*, MAOC-1/ hydratase, DHS-28/dehydrogenase and DAF-22/thiolase were reported to play important roles in the peroxisomal β-oxidation pathway [32]. If the peroxisome is responsible for the degradation of mmBCFA C17ISO, disruption of peroxisomal fatty acid β-oxidation may increase the C17ISO level and suppress the phenotype of elo-5(If). We then applied RNAi to knock down maoc-1, dbs-28 and daf-22 and found the treatments were not able to suppress the developmental defect of elo-5(If). Further analysis using single, double and triple mutants of these genes also generated negative results, indicating that mutating these genes is not sufficient to rescue the phenotype of elo-5(If).

We thus analyzed the fatty acid profiles of these single mutants. We found that a deletion mutant of maoc-1 displayed a significant increase in the level of C15ISO (~55%) and a small but still significant increase in the level of C17ISO (~17%) in total lipid extracts, over that in samples from wild type (Figure 4C). This increase suggests that the MAOC-1-involved β-oxidation pathway contributes to the degradation of mmBCFA, but that the increase is not sufficient to suppress the defect of elo-5(If).

GC analyses of deletion mutants of dbs-28 and daf-22 showed that eliminating the activity of either gene did not cause significant increase in C17ISO level, suggesting that mutating these two genes has a weaker effect on the β-oxidation pathway than the maoc-1 mutant. One possible scenario is that there is genetic redundancy associated with all of these genes so that none of these individual mutations completely abolish peroxisomal β-oxidation function and thus cannot sufficiently raise the mmBCFA level for the suppression of elo-5(If). Consistent with this idea, none of three mutants displayed robust morphological defects (see further discussion in Discussion). An RNAi-based search for suppression of the elo-5(If) phenotype failed to identify factors in additional pathways potentially involved in FA degradation, although it is possible that RNAi may not be sufficient to disrupt the function of some peroxisomal genes.

Together, these results suggest that the increase of mmBCFA level in the prx-5(If); elo-5(If) suppressor mutant is not likely caused by a defect in a single peroxisomal-involved mechanism that impacts mmBCFA degradation or biosynthesis. While β-oxidation may contribute to the change in mmBCFA level, β-oxidation-independent mechanisms are also likely to be involved (see Discussion).

**Discussion**

Despite the broad presence of mmBCFAs in multiple organisms, little is known about their metabolism and roles in regulating development. In our previous studies, we discovered that inactivating the elo-5 gene depleted mmBCFAs in *C. elegans*, causing robust arrest in post-embryonic development.
A mmBCFA-containing sphingolipid mediates this function by activating the TORC1 signaling pathway in the intestine [16,33]. In this study, we identified a new physiological role for peroxisomal functions in regulating mmBCFA metabolism that critically affects animal development and behaviors. Compromising the peroxisomal import function leads to an increase in mmBCFA levels in animals defective in mmBCFA biosynthesis, and likely does so by affecting multiple mechanisms. While over-expression of elo-6 is not sufficient to suppress elo-5(lf), we cannot exclude the possibility that

Figure 4. prx-5/6 mutations likely cause a decrease in mmBCFA degradation. (A) qRT-PCR data showing mRNA levels of the elo-6 gene in worms of indicated genotypes. elo-5(lf); Ex[elo-6] is a transgenic strain that significantly overexpressed the elo-6 gene and was not sufficient to suppress the L1 arrest phenotype of elo-5(lf). The data suggest that a higher level of elo-6 expression is not likely the cause of the suppression. (B) qRT-PCR data showing mRNA levels corresponding to several other FA elongation enzymes in worms of indicated genotypes. Since the elo-5 mRNA being measured is from the elo-5(gk208, deletion) allele, the significant decrease in elo-5 mRNA level is likely due to mutation-induced mRNA degradation. (C) Comparison of mmBCFA composition in the indicated strains. Percentage of C15ISO and C17ISO has a statistically significant increase in the maoc-1(lf) mutant.

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prx-5(lf) increases the synthesis of mmBCFA in the elo-5(lf) mutant even though it is unlikely given the known roles associated with the peroxisome.

Fatty acid β-oxidation is one of the important peroxisomal functions in all eukaryotic organisms. Compromising peroxisomal FA β-oxidation function has been shown to cause acetyl-CoA accumulation [34]. The results from this study suggest that prx-5 mutants have a reduced rate of mmBCFA degradation. In the elo-5(lf) mutant, although mmBCFA biosynthesis is dramatically compromised, it still remains at a low level, which depends on the function of elo-6. Under such a condition, decreasing mmBCFA degradation through compromising peroxisomal FA β-oxidation might increase the impermissibly low concentration of mmBCFAs and thus rescue elo-5(lf) developmental defect phenotype. However, we failed to phenocopy the suppression effect of elo-5(lf) by mutating or knocking down genes responsible for peroxisomal fatty acid β-oxidation. The possible explanation is that an inactivating prx-5 compromises multiple pathways involved in fatty acid degradation, including the peroxisomal FA β-oxidation pathway and other lipid synthesis pathway. Compromising peroxisomal FA β-oxidation alone may contribute to, but may not be sufficient, to cause the adequate increase in mmBCFA level that suppresses the developmental arrest of elo-5(lf) observed in the prx-5(lf); elo-5(lf) double mutant. Consistent with this idea, the level of mmBCFA C17ISO has mild increase in a deletion mutant of a key gene in the peroxisomal FA β-oxidation pathway (Figure 4C). It is possible that the RNAi and compromising peroxisomal FA β-oxidation might increase the impermissibly low concentration of mmBCFAs and thus rescue elo-5(lf) developmental defect phenotype.

Additionally, peroxisomal FA β-oxidation has been shown to be involved in the synthesis of certain lipids such as components of active dauer pheromone [34,35]. We could thus also speculate that mmBCFAs might be partially degraded or chain-shortened in the peroxisome and integrated into certain types of complex lipids that negatively regulate post-embryonic development. An alternative hypothesis is that mmBCFA could be incorporated into other lipids by peroxisomal enzymes during lipid anabolism. This would reduce mmBCFA levels and compete with their developmental roles. Therefore, compromising peroxisomal function may decrease the efflux of mmBCFA and lead to increasing the level of mmBCFAs.

Thus, our results raised several interesting and important questions regarding peroxisome-mmBCFA relations and functions. For example, what other peroxisomal pathways are involved in mmBCFA metabolism, and are such mechanisms involved in feedback regulation of mmBCFA levels. It is certainly important for us to understand whether the findings in C. elegans are conserved in mammals and if so what are physiological effects of de novo biosynthesis and dietary uptake of mmBCFA in humans.

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Author Contributions

Conceived and designed the experiments: RW MK. Performed the experiments: RW. Analyzed the data: RW MK MH. Contributed reagents/materials/analysis tools: MK. Wrote the manuscript: RW MH. Supervised the study: MH.

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