Running title: Mitochondrial fatty acid synthase

Dual-localized enzymatic components constitute the fatty acid synthase systems in mitochondria and plastids

Xin Guan,a,b Yozo Okazaki,c,1 Rwisdom Zhang,b,d Kazuki Saito,c,e and Basil J Nikolaua,b,f,*

a Roy J. Carver Department of Biochemistry, Biophysics, and Molecular Biology, Iowa State University, Ames, Iowa 50011, USA; b The Engineering Research Center for Biorenewable Chemicals (CBiRC), Iowa State University, Ames, Iowa 50011, USA; c Metabolomics Research Group, RIKEN Center for Sustainable Resource Science, Yokohama 230-0045, Japan; d Department of Chemical Engineering, University of Southern California, Los Angeles, CA 90007, USA; e Graduate School of Pharmaceutical Sciences, Chiba University, Chiba 260-8675, Japan; f Center for Metabolic Biology, Iowa State University, Ames, Iowa 50011, USA; 1 Current address: Graduate School of Bioresources, Mie University, Tsu City, Mie 514-8507, Japan

* For correspondence (e-mail dimmas@iastate.edu)

One-sentence summary: Mitochondrial- and plastid-localized fatty acid synthase systems share 3 enzymatic components, products of dual-targeted gene expression.

Author contributions

X.G. and B.J.N. designed the research; Y.O. and K.S. analyzed the lipidome; X.G. and R.Z. performed the in vitro enzymatic assays; X.G. and B.J.N. carried out all other
experiments; X.G. and B.J.N. coordinated the preparation of the article; all authors contributed to the analysis of the collected data and writing of the article.
Plant fatty acid biosynthesis occurs in both plastids and mitochondria. Here, we report the identification and characterization of Arabidopsis (*Arabidopsis thaliana*) genes encoding three enzymes shared between the mitochondria- and plastid-localized Type II fatty acid synthase systems (mtFAS and ptFAS, respectively). Two of these enzymes, β-ketoacyl-acyl carrier protein (ACP) reductase (pt/mtKR) and enoyl-ACP reductase (pt/mtER) catalyze two of the reactions that constitute the core four-reaction cycle of the FAS system, which iteratively elongates the acyl-chain by two carbon atoms per cycle. The third enzyme, malonyl-CoA:ACP transacylase (pt/mtMCAT) catalyzes the reaction that loads the mtFAS system with substrate by malonylating the phosphopantetheinyl cofactor of ACP. Green fluorescent protein (GFP) fusion experiments revealed that these enzymes localize to both chloroplasts and mitochondria. This localization was validated by characterization of mutant alleles, which were rescued by transgenes expressing enzyme variants that were retargeted only to plastids or only mitochondria. The singular retargeting of these proteins to plastids rescued the embryo-lethality associated with disruption of the essential ptFAS system, but these rescued plants displayed phenotypes typical of the lack of mtFAS function, including reduced lipoxygenation of the H subunit of the glycine decarboxylase complex, hyperaccumulation of glycine, and reduced growth. However, these latter traits were reversible in an elevated CO₂ atmosphere, which suppresses mtFAS-associated photorespiration-dependent chemotypes. Sharing enzymatic components between mtFAS and ptFAS systems constrains the evolution of these non-redundant fatty acid biosynthetic machineries.
INTRODUCTION

In plants, *de novo* fatty acid biosynthesis occurs in two distinct subcellular compartments, the plastids and mitochondria (Ohlrogge & Jaworski, 1997, Wada et al., 1997). These two fatty acid biosynthetic systems utilize an acyl carrier protein (ACP)-dependent, multi-component Type II fatty acid synthase (FAS) to catalyze the assembly of these essential cellular components. Furthermore, genetic studies indicate that these two FAS systems are not redundant and have been evolutionarily retained (Guan et al., 2017). The plastid-produced fatty acids serve as the acyl building blocks for the assembly of the majority of the lipids in plant cells, including membrane lipids, signaling lipids and storage lipids (Li-Beisson et al., 2013). In comparison, the primary role of mitochondrial FAS (mtFAS) is to synthesize the acyl-precursor for the biosynthesis of lipoic acid (Yasuno & Wada, 1998, Wada et al., 2001a), which is the cofactor essential for the catalytic competence of several key metabolic enzymes, including the glycine decarboxylase complex (GDC), mitochondrial pyruvate dehydrogenase (mtPDH), α-ketoglutarate dehydrogenase (KGDH), branched-chain α-ketoacid dehydrogenase (BCKDH) (Taylor et al., 2004), and plastidial PDH (ptPDH) (Yasuno & Wada, 2002, Wada et al., 2001b). In addition, mtFAS appears to be involved in remodeling mitochondrial cardiolipins (Frentzen & Griebau, 1994, Griebau & Frentzen, 1994) and in detoxifying free malonic acid (Guan & Nikolau, 2016), a competitive inhibitor of succinate dehydrogenase of the tricarboxylic acid cycle (Quastel & Wooldridge, 1928, Greene & Greenamyre, 1995).

Detailed biochemical characterizations, going back to the 1980’s, have led to the identification of the four enzymatic components of the core plastidic FAS (ptFAS) system, i.e., three isozymes of 3-ketoacyl-ACP synthases (ptKAS), a 3-ketoacyl-ACP reductase (ptKAR), a 3-hydroxyacyl-ACP dehydrase (ptHD) and an enoyl-ACP
reductase (ptER). In contrast, the enzymatic components of the mtFAS system are less well characterized. To date, only two of the core enzymatic components of the plant mtFAS system have been characterized, β-ketoacyl-ACP synthase (mtKAS) (Yasuno et al., 2004, Olsen et al., 2004, Ewald et al., 2007) and 3-hydroxyacyl-ACP dehydratase (mtHD) (Guan et al., 2017), along with the three mtACP isoforms that carry the intermediates of the mtFAS system (Fu et al., 2020). Other characterized supportive components of the mtFAS system include the mitochondrial phosphopantetheinylation transferase (mtPPT) that activates mtACP by phosphopantetheinylation (Guan et al., 2015), and malonyl-CoA synthetase (mtMCS) that generates the malonyl-CoA precursor for mtFAS (Guan & Nikolau, 2016). These more recent characterizations of the mtFAS system establish that whereas the ptFAS system uses acetyl-CoA, generated by the ptPDH, as the precursor for fatty acid biosynthesis, mtFAS uses malonyl-CoA as the precursor, which is generated by mtMCS (Guan & Nikolau, 2016).

Here in, we report the identification and characterization of three additional enzymatic components of mtFAS that have not previously been identified. Two of these catalyze core reactions of the mtFAS cycle, namely β-ketoacyl-ACP reductase (KR), and enoyl-ACP reductase (ER), and the third is the mitochondrial malonyl-CoA:ACP transacylase (MCAT), which loads the mtFAS system by malonylating the phosphopantetheinyloxyl cofactor of ACP. Unexpectedly, genetic and transgenic expression of fluorescently tagged proteins indicate that these three enzymatic components are shared between both the mtFAS and ptFAS systems.
RESULTS

Computational identification of candidate genes encoding mtFAS catalytic components

BLAST analysis of the Arabidopsis (*Arabidopsis thaliana*) genome using the sequences of the yeast (*Saccharomyces cerevisiae*) mtFAS components and *Escherichia coli* FAS components as queries identified putative Arabidopsis ORFs coding for mitochondrial MCAT, KR and ER catalytic components (Supplemental Figure 1). These analyses identified a single candidate gene each for mitochondrial MCAT (AT2G30200) (Bryant et al., 2011) and KR (AT1G24360) (Bryant et al., 2011), and two potential candidate genes for the mitochondrial ER (AT3G45770 and AT2G05990 (Mou et al., 2000, Wu et al., 2015)). The sequences of these candidate proteins were the most similar in the Arabidopsis proteome identified in TAIR 10 (www.arabidopsis.org), as indicated by BLAST e-value scores of between $10^{-15}$ and $10^{-68}$, and these proteins share between 20% and 45% sequence identities with the query sequences (Supplemental Figure 1). Prior characterizations have indicated that all but AT3G45770 encode components of the ptFAS system (Bryant et al., 2011, Mou et al., 2000, Wu et al., 2015), and computational analyses suggest that AT3G45770 encodes a mitochondrial-localized ER component (Li-Beisson et al., 2013, Li-Beisson et al., 2010).

Organelle targeting of candidate mtFAS components

Because prior studies of these candidate mtFAS genes have indicated that they may be components of the ptFAS system (Bryant et al., 2011, Mou et al., 2000, Wu et al., 2015), initial characterizations directly evaluated whether these putative mtFAS component enzymes are mitochondrially located. Transgenic experiments were conducted with GFP-fusion proteins, expressed under the transcriptional regulation of the CaMV 35S
promoter (Figure 1). Three types of GFP-fusion transgenes were evaluated for each of
the candidate mtFAS component proteins: a) GFP was translationally fused at the C-
terminus of each candidate mtFAS component protein; b) the N-terminal segment from
each candidate mtFAS component protein that was computationally predicted to be an
organelle-targeting pre-sequence was translationally fused to the N-terminus of GFP;
and c) GFP was translationally fused at the C-terminus of each candidate mtFAS
component protein lacking the putative organelle-targeting pre-sequence. Individual
GFP-fusion transgenes were stably integrated into the Arabidopsis genome, and
confocal micrographs of the resulting transgenic roots and leaves visualized the
subcellular location of the GFP-fusion proteins.

In these experiments, organelles were identified by a combination of two fluorescence
markers: a) MitoTracker Orange for mitochondria and chlorophyll auto-fluorescence for
plastids; and b) the fluorescence signals revealed by the expression of control GFP-
tagged markers, the p35S::mtPPT1-240-GFP transgene (Figure 1A) that we previously
identified as being targeted to mitochondria (Guan et al., 2015), and the p35S::UGP31-
600-GFP transgene that is plastid targeted (Okazaki et al., 2009) (Figure 1A). These
control markers show distinct patterns that are consistent with mitochondrial or plastid
localization, and these are distinct from the GFP-signal observed with the non-targeted
GFP control, which as previously characterized (Li et al., 2011a), is located in the cytosol
and nucleus (Figure 1A).

The fluorescence observed in transgenic plants carrying C-terminal GFP-fusions with
each full-length candidate mtFAS component-protein showed an assortment of organelle
localizations. The interpretation of these results can be simplified if one considers that
subcellular targeting information is segregated between both the N-terminal, potential
organelle-targeting segment, and the mature segment of each protein. This was
particularly the case for the proteins encoded by AT2G05990, AT2G30200 and AT1G24360. The full-length proteins encoded by these three genes guide the expression of the fused GFP to both mitochondria and plastids (Figure 1C to 1E). In the case of AT2G05990-encoded protein, the N-terminal pre-sequence directs GFP to plastids (row 2 of Figure 1E), but the segment which lacks this N-terminal pre-sequence guides GFP to mitochondria (row 3 of Figure 1E). In contrast, the N-terminal pre-sequences of AT2G30200 and AT1G24360 direct the GFP-fusions to mitochondria (row 2 of Figures 1C and 1D), whereas upon removal of these N-terminal pre-sequences from each protein, the remaining mature segments direct the accumulation of the fused GFP protein to plastids (row 3 of Figures 1C and 1D).

Subcellular targeting information of the AT3G45770-encoded protein is simpler to interpret, with the full-length AT3G45770-protein (rows 1 of Figure 1B) and the N-terminal pre-sequence (rows 2 of Figure 1B) directing GFP to mitochondria, whereas removal of the N-terminal pre-sequence from the AT3G45770-encoded protein guides GFP to the cytosol (rows 3 of Figure 1B). In summary therefore, in contrast to the AT3G45770-encoded protein, proteins encoded by AT2G30200, AT1G24360 and AT2G05990 encode dual localization signals, for both mitochondria and plastids. One of these signals resides in the N-terminal pre-sequence and the other in the mature portion of these proteins.

**Experimental authentication of candidate mtFAS component enzymes**

The catalytic functions of the individual putative FAS component proteins were explored via two independent strategies. In the first series of experiments, the catalytic function of the putative mtFAS component proteins was evaluated by expressing each protein in *S. cerevisiae* mutant strains that lack a functional copy of a mtFAS component and testing...
for genetic complementation. The expression of each candidate Arabidopsis protein was accurately targeted to yeast mitochondria, by genetically fusing the mitochondrial pre-sequence (MP) of the yeast COQ3 protein (Hsu et al., 1996) to the N-terminus of the mature Arabidopsis proteins, and their expression in yeast was under the transcriptional control of the constitutive PGK promoter (de Moraes et al., 1995).

The yeast mutant strains that lack mtFAS functions cannot utilize glycerol as a sole carbon source because they are deficient in respiration (Torkko et al., 2001). On media that use glycerol as the sole carbon source, the yeast mutant strains lacking individual mtFAS components (i.e., mct1, oar1 or etr1) showed no growth unless they expressed the mitochondrially-targeted Arabidopsis proteins encoded by AT2G30200, AT1G24360, AT2G05990 or AT3G45770 (Figure 2A). In each case these results mirrored the results obtained by similarly expressing the yeast MCT1, OAR, or ETR1 proteins (Figure 2A). In contrast, neither the control empty expression plasmid or the expression plasmid carrying only the COQ3 mitochondrial targeting element rescued the growth deficiency of these yeast strains in glycerol media (Figure 2A).

In the case of the ER components of the mtFAS system, recombinant purified proteins encoded by AT2G05990 or AT3G45770 (expressed in E. coli) were also evaluated in vitro for their ability to catalyze the expected chemical reaction. Because enoyl-ACP reductases are active with both enoyl-ACP (their native substrate) and enoyl-CoA (Chen et al., 2008), in these experiments each protein was tested for the ability to reduce enoyl-CoA substrates. These assays were conducted with Δ2-trans-10:1-CoA and Δ2-trans-16:1-CoA, and activity was monitored by the decrease in A340 due to the coupled oxidation of the pyrimidine nucleotides (NADH or NADPH). Both AT2G05990 and AT3G45770 proteins were capable of reducing the enoyl-CoA substrates, and they exhibited comparable K_m, V_max and catalytic efficiency (k_cat/K_m) with both tested
substrates (Figure 2B). Moreover, these assays established that AT2G05990 is an
NADH-dependent reductase, and its activity with NADPH was undetectable. In contrast,
AT3G45770 is an NADPH-dependent reductase, and its activity with NADH was
undetectable.

In combination therefore, the GFP-transgenic fluorescence data, the yeast genetic
complementation experiments and the biochemical characterizations of purified proteins
indicated that three Arabidopsis genes (AT2G30200, AT1G24360 and AT2G05990)
encode proteins that are dual targeted to plastids and mitochondria, and they have the
ability to catalyze the MCAT, KR and ER reactions, respectively. We therefore labeled
these proteins as pt/mtMCAT (AT2G30200), pt/mtKR (AT1G24360) and pt/mtER
(AT2G05990) indicating their dual localizations. In contrast, AT3G45770 encoded a
mitochondrially-localized ER enzyme, which we labeled as mtER, indicating its
functionality in the sole organelle.

The in planta roles of pt/mtMCAT and pt/mtKR in mtFAS

The role of pt/mtMCAT (AT2G30200) and pt/mtKR (AT1G24360) as enzymatic
components of the mtFAS system was further evaluated by characterizing Arabidopsis
plants carrying T-DNA-tagged mutant alleles at each locus (details in the Materials and
Methods section). As previously described (Bryant et al., 2011), mutant plants
homozygous for the T-DNA allele at the AT2G30200 locus are not recoverable as they
are embryo lethal. Similarly, mutant plants homozygous for the T-DNA allele at
AT1G24360 also display an embryo lethal phenotype. This embryo-lethality is
associated with the fact that these two gene products are components of the ptFAS
system (Bryant et al., 2011), which prior genetic studies have established as being
essential; these genetic conclusions are exemplified by mutations in other ptFAS
components, such as the heteromeric acetyl-CoA carboxylase hACC\textsubscript{ase} (Li et al., 2011b) and 3-ketoacyl-ACP synthase I KASI (Wu & Xue, 2010).

Therefore, we designed transgenic complementation experiments to further confirm that these dual localized gene products are also components of the mtFAS system. Specifically, plants that were heterozygous mutants at the AT2G30200 locus (pt/mtMCAT) or the AT1G24360 locus (pt/mtKR) were transformed with transgenes that express two versions of the pt/mtMCAT or pt/mtKR proteins, respectively. One version of these transgenes expressed the ORF that encodes the full-length proteins, and as indicated by the GFP transgenic localization experiments, these full-length proteins contained both an N-terminal, mitochondrial targeting pre-sequence and an internal plastid-targeting signal. These proteins would therefore be expected to be dual-targeted to both plastids and mitochondria. The other version of these proteins expressed ORFs missing the N-terminal targeting pre-sequence (removing the first 68 and 71 amino acids, respectively, from the full-length proteins) and thereby deleting the mitochondrial-targeting information from each protein. These truncated proteins contained only the internal plastid-targeting signal and thereby would be expected to express these catalytic functions only in plastids, but not in mitochondria.

Compared to the non-transgenic siblings, which failed to generate homozygous mutant plants (due to missing ptFAS function), all four recovered transgenic lines generated homozygous mutant progeny plants. At 16 day after imbibition (DAI), the transgenic plants, which expressed the full-length pt/mtMCAT or pt/mtKR proteins, were indistinguishable from wild-type plants, indicating that these transgenically expressed proteins complemented the deficiency in both mtFAS and ptFAS function.
In contrast, the transgenic plants expressing the N-terminally truncated pt/mtMCAT or pt/mtKR proteins (i.e., these proteins were predicted to be plastid localized but not targeted to mitochondria) exhibited reduced size (Figure 3A). Most significantly, when these plants were grown in a 1% CO2 atmosphere, where photorespiration deficiency is suppressed, the stunted growth morphology was reversed (Figure 3A). Further biochemical analyses of these plants established that the lipoylation status of the H protein subunit of glycine decarboxylase (GDC) was reduced to less than 10% of the control levels (Figure 3B). Analogous immunoblot analyses revealed the lipoylation states of additional three lipoylated proteins, namely the E2 subunits of mitochondrial PDH and KGDH and the E2 subunit of the plastidial PDH (Supplemental Figure 2); the E2 subunit of BCKDH was not detectable (Ewald et al., 2007).

As a consequence of the reduced lipoylation of the H protein, glycine levels were induced by about 100-fold in these plants (Figure 3C). Growing these transgenic mutant plants in the 1% CO2 atmosphere restored the accumulation of glycine to near control levels (Figure 3C). Changes in the levels of other amino acids were barely detectable (Supplemental Figure 3).

The reduced lipoylation of the H protein of GDC is an attribute previously characterized with mutations in other mtFAS components (e.g., mtKAS (Ewald et al., 2007), mtHD (Guan et al., 2017), mtPPT (Guan et al., 2015), mtMCS (Guan & Nikolau, 2016), and mtACP (Fu et al., 2020). These latter mutants exhibit a growth-stunting that is reversible when plants are grown in an elevated CO2 atmosphere. Such characteristics have been attributed to the fact that mtFAS generates the lipoic acid cofactor for GDC, and this deficiency blocks photorespiration, leading to the growth deficiency, and hyperaccumulation of glycine, traits that are all reversed when mtFAS mutant plants are grown in an elevated CO2 atmosphere that suppresses photorespiration (Guan et al.,
Collectively therefore, these findings demonstrate that the MCAT and KR catalytic functions of the mtFAS and ptFAS systems are genetically encoded by two respective genes that each encode dual localized proteins, pt/mtMCAT and pt/mtKR.

**Two enoyl-ACP reductase isozymes for mtFAS**

Two genetic loci appear to encode proteins that catalyze the enoyl-ACP reduction reaction of mtFAS, the AT2G05990 locus, which encodes an NADH-dependent reductase, and the AT3G45770 locus, which encodes an NADPH-dependent reductase. The former protein was dual targeted (i.e., pt/mtER), and the latter was targeted to mitochondria (i.e., mtER). We genetically dissected the roles of these two enoyl-ACP reductase genes by characterizing three mutant lines: 1) two T-DNA-tagged mutant alleles at the mtER locus (mter-1 and mter-2), both of which eliminated the expression of mtER; 2) RNAi knockdown lines of the pt/mtER locus (pt/mter-rnai-1 and pt/mter-rnai-2), which reduced the expression of pt/mtER to ~2% of wild-type levels; and 3) double mutant lines (i.e., mter-1-pt/mter-rnai-1 and mter-1-pt/mter-rnai-2), which eliminated the expression of mtER and reduced the expression of pt/mtER to ~2% of wild-type levels (Figure 4).

As indicated by the exemplary data gathered from plants at 16 DAI, plants homozygous for the mutant mtER alleles were morphologically and metabolically (i.e., amino acids, fatty acids, and lipids) indistinguishable from wild-type plants (Supplemental table 1). In contrast, the aerial organs of the pt/mter-rnai mutant and mter-1-pt/mter-rnai double mutant lines were considerably reduced in size (Figure 4A). This growth phenotype was equally expressed independent of whether these plants were grown in ambient air or a 1% CO₂ atmosphere (Figure 4A); the expectation being that in the latter conditions any potential mtFAS-associated photorespiration phenotype would be suppressed (Ewald et
Therefore, these observations suggest that these mutations do not affect mtFAS, and they are consistent with prior characterization of an ethyl methanesulfonate-generated pt/mter mutant (Mou et al., 2000, Wu et al., 2015) that identified this gene product as a component of the ptFAS system.

In contrast, biochemical analyses that evaluated the metabolic status of these mutant plants indicated that both mtER and pt/mter contribute to mtFAS. The evidence that supports this conclusion includes: a) immunoblot analysis that indicated the lipoylation states of GDC in mter and pt/mter-rnai single mutants was reduced to between 60% to 80% of the wild-type levels, and this protein was even further under-lipoylated to about 10% of the wild-type level in the double mutant plants (Figure 4B); b) accompanied with the reduction in lipoylation status of the H protein, glycine accumulation was increased by about 15-fold in these double mutant lines; and c) this latter attribute was completely reversed when these double mutant plants were grown in the 1% CO2 atmosphere, which suppresses photorespiration (Figure 4C). Other changes in amino acid levels were detected in these double mutants (Supplemental Table 1). Collectively therefore, we conclude that the mtFAS system appears to be redundantly enabled by two enoyl-ACP reductases, a dual localized enoyl-ACP reductase (pt/mtER; AT2G05990) and a mitochondrially localized enoyl-ACP reductase (mtER; AT3G45770).
As a consequence of the evolutionary origin of mitochondria and plastids as eubacterial symbionts in eukaryotic cells (Gray, 2004), plant cells have maintained two Type II FAS systems that are located in these organelles. Whereas the genetic and enzymatic components of the ptFAS system have received considerable attention (Ohlrogge & Jaworski, 1997), the analogous components of the mtFAS system are only just being characterized (Fu et al., 2020, Guan et al., 2015, Guan & Nikolau, 2016, Guan et al., 2017). This complexity associated with organelle-specificity of the FAS systems is further complicated by the genetic redundancy that has been revealed by the genomic-based bioinformatic analysis of plant genomes, as revealed by Li-Beisson et al (Li-Beisson et al., 2013). This study completes the identification and characterization of the core catalytic components of the mtFAS system. These characterizations unexpectedly find that some of these components are shared between the ptFAS and mtFAS system, which was not predictable based solely on bioinformatic analysis of genomic data (Beisson et al., 2003).

Dual-localized MCAT, KR and ER components of mtFAS

The plant mtFAS system is a pathway that contributes acyl substrates required for a series of essential metabolic processes, including photorespiration and biosynthesis of lipid A-like molecules (Guan et al., 2017). To date four enzymatic components of the mtFAS system have been experimentally characterized, mtKAS (Yasuno et al., 2004, Olsen et al., 2004, Ewald et al., 2007), mtHD (Guan et al., 2017), mtPPT (Guan et al., 2015) and mtMCS (Guan & Nikolau, 2016). In this study we identified the additional three components that are required to complete the mtFAS cycle. These three components (pt/mtMCAT, pt/mtKR and pt/mtER) localized to both plastids and
mitochondria, referred to as “dual-targeted”, and thus they contribute to both mtFAS and
ptFAS capabilities. In addition, the mitochondrial ER reaction appears to be catalyzed by redundant enzymes, an NADPH-dependent mtER and a dual-targeted NADH-dependent pt/mtER. The pt/mtER and mtER belong two distinct families of enzymes; pt/mtER belongs to the short chain dehydrogenase/reductase (SDR) family, whereas mtER belongs to the medium chain dehydrogenase/reductase (MDR) family (Hiltunen et al., 2010). Despite the distinct preference for different reducing cofactors (NADH versus NADPH), both pt/mtER and mtER display a similar chain length preference for the 2-enoyl substrates, being able to almost equally utilize both medium (10 carbon atoms) and long-chain (16 carbon atoms) substrates. This latter finding is consistent with the ability of the plant mtFAS system to not only generate octanoic acid for lipoic acid biosynthesis, but also longer chain fatty acids that are used in the assembly of lipid A-like molecules (Guan et al., 2017).

The dual localized mtFAS components (i.e., pt/mtMCAT and pt/mtKR) had previously been characterized as components of the ptFAS system (Bryant et al., 2011, Wu & Xue, 2010, Mou et al., 2000), and these characterizations had established the essentiality of these components and the ptFAS system during embryogenesis. Thus, deducing that these are also components of the mtFAS system required a combination of reverse genetic and transgenic strategies. Specifically, we generated plants that were deficient in mitochondrial MCAT or KR functions but normally expressed plastidial-localized MCAT or KR functions. This could be achieved by the fact that for these two proteins, the plastid-targeting information is encompassed within the mature protein sequence, whereas the mitochondrial-targeting information is encompassed in the N-terminal signal-peptide extension sequence. Thus, the transgenically expressed, N-terminal truncated MCAT or KR proteins were singly targeted to plastids, rather than to both the
plastids and mitochondria. Therefore, these transgenically plastid-only retargeted MCAT or KR alleles rescued the embryo lethal phenotypes associated with the deficiency in ptFAS, but the recovered transgenic plants displayed phenotypes that are typical of plants lacking mtFAS functionality. These phenotypes include reduced H-protein lipoylation, elevated glycine accumulation and reduced growth; the latter two attributes being reversed by growing plants in a CO₂-enriched atmosphere. Because these biochemical traits are commonly expressed with other mutations that affect mtFAS, these findings are consistent with the role these two gene products have in providing the MCAT or KR enzymatic functions of the mtFAS system, respectively.

The characterization of the mitochondrial ER component was more complex because two loci can provide this functionality, a dual-localized pt/mtER (AT2G05990) and a singly targeted mtER (AT3G45770). Plants deficient in mitochondrial ER function were generated by combining mutants that affect both loci, using a combination of T-DNA-tagged null alleles at AT3G45770 and RNAi-knock-down alleles of AT2G05990. These plants exhibited traits that are typical of a deficiency in mtFAS; namely, depleted lipoylation of the H-protein, hyperaccumulation of glycine and miniature aerial organs. All these traits appear when these plants are grown in ambient air, and glycine hyperaccumulation is reversed when they are grown in an elevated CO₂ atmosphere.

Dual-localization of proteins to plastids and mitochondria

Proteomic analyses of isolated organelles have indicated that a number of proteins (>100) generated from a single gene locus may be dual-targeted to both mitochondria and plastids (Carrie et al., 2009b). Although this conclusion is based on experimental outcomes that are fraught with technical challenges associated with the ability to purify distinct organelle preparations (Rao et al., 2017, van Wijk & Baginsky, 2011), the use of
fluorescently tagged proteins substantiate the occurrence of this phenomenon, although even these experiments need to be carefully considered (Sharma et al., 2018b, Sharma et al., 2018a). In this study, we presented GFP localization studies that substantiate the global proteomics evidence concerning dual plastid-mitochondrial localization conclusions. Additional molecular genetic studies used transgenic constructs to retarget the dual-localized proteins to either plastids or mitochondria. These experiments confirmed that the plastid re-targeted constructs only affected the ptFAS-associated outcomes, and the mitochondrial re-targeted constructs only affected the mtFAS-associated outcomes.

An obvious evolutionary advantage of such dual localization of products from a single gene locus would be the apparent energy and resource economy of maintaining only one gene for both organelles instead of a gene for each organelle. However, from an evolutionary point of view, concentrating two functions in a single gene locus is unusual, as the more common tendency is for neo-specialization following gene duplications (Carrie & Small, 2013). In addition, there are clear disadvantages for dual targeted proteins. For example, the protein may not function optimally in both organelles, in terms of pH optima, concentration of substrates and co-factors, and protein-protein interactions. Moreover, a single crucial mutation could lead to the loss of this function in both organelles.

Insights may be provided by considering the functions of the proteins that are known to be dual-targeted. Dual-targeted proteins appear to be enriched in a few specific functional groups (Carrie & Small, 2013), such as DNA replication, tRNA biogenesis, protein translation and metabolic processes (e.g. fatty acid biosynthesis as demonstrated in the current study). This potential bias may reflect mechanistic
constraints on dual-targeting and selection pressure that could favor dual-targeting over evolutionary timescales.

Mechanistically, for dual-targeting between mitochondria and plastids, the targeted protein has to be recognized by the import machinery of both organelles (Carrie et al., 2009a). Evolutionarily, the protein import machineries in these organelles arose independently, are non-homologous, and therefore would normally be expected to recognize different targeting signals (Schleiff & Becker, 2011). Indeed, in the case of pt/mtMCAT, pt/mtKR and pt/mtER, these proteins appear to utilize bi-partite targeting signals, an N-terminal signal unique for one organelle and an internal signal that specifies import to the other organelle. Deletion of the N-terminal signal of pt/mtMCAT and pt/mtKR abolishes import into mitochondria and enhances import into plastids, whereas the contrary was observed for pt/mtER. Similar situations have been reported with many other such dual targeted proteins (e.g. the amino acyl-tRNA synthetases, AspRS, LysRS and ProRS (Berglund et al., 2009a, Berglund et al., 2009b), mitochondrial carrier protein, BT1 (Bahaji et al., 2011), and phosphatidyl glycerophosphate synthase 1 (Babiychuk et al., 2003)); deletion of their N-terminal targeting sequences only affects localization to one of these two organelles. In the case of the mtER isozyme, deletion of the N-terminal targeting sequence resulted in the cytosolic localization of the protein, indicating the presence of an additional signal that is needed for dual-localization.

Dual localization of pt/mtMCAT, pt/mtKR and pt/mtER, a trait acquired during the co-evolution of plastids and mitochondria, suggests that further investigations of protein sorting mechanisms and re-evaluation of organelle proteomes may be useful to reveal how generalizable this phenomenon is in plant cells. Moreover, the general persistence of such genetic and biochemical redundancy in plant metabolism may indicate
evolutionary biological advantages to the generation of such complexity in metabolism, a characteristic that was initially surmised from the study of specialized metabolism (Pichersky & Lewinsohn, 2011).

**MATERIALS AND METHODS**

**Yeast strains and genetic complementation**

The yeast (*Saccharomyces cerevisiae*) strain deficient in the *MCT1*, *OAR1* and *ETR1* genes (YBR026C; BY4741 background; and MATa) was obtained from Thermo Scientific. The yeast *MCT1* (primers M1 and M2, see Supplemental Table 2 for primer sequences), *OAR1* (primers M3 and M4) and *ETR1* genes (primers M5 and M6) of wild-type BY4741 strain were cloned into YEp351 vector (PGK promoter-driven gene expression) (de Moraes et al., 1995). Arabidopsis (*Arabidopsis thaliana*) *pt/mtMCAT* (AT2G30200, primers M7 and M8), *pt/mtKR* (AT1G24360, primers M9 and M10), *pt/mtER* (AT2G05990, primers M11 and M12) and *mtER* (AT3G45770, primers M13 and M14) genes of the Col-0 strain were cloned into YEp351M (Guan et al., 2017) (5’ plant targeting signals were replaced by the yeast COQ3 mitochondrial presequence CDS). Complementation tests were performed as previously described (Torkko et al., 2001).

**Protein overexpression and in vitro kinetic assays**

The *mt/ptER* (primers M15 and M16) and *mtER* (primers M17 and M18) genes were cloned into pBE522 vector (Zhu et al., 2011). These constructs express recombinant proteins with a His-tag located at the N-terminus. Recombinant proteins were expressed in the *Escherichia coli* BL21* strain (Invitrogen, Carlsbad, CA) and further purified using Probond Nickel-Chelating Resin (Invitrogen). The kinetic constants of *mt/ptER* and *mtER*
were determined spectrophotometrically as previously described (Chen et al., 2008) with modification. Specifically, the enoyl-CoA substrates (i.e., trans-Δ\(^2\)-10:1 and trans-Δ\(^2\)-16:1) were synthesized as previously described (Guan et al., 2017). Concentrations of three ingredients were optimized (i.e., 30 mM potassium phosphate (pH = 7.8), 2-200 μM enoyl-CoA, 300 μM NADPH or NADH, and 3 ug/mL recombinant proteins). The consumption of NADPH or NADH at 340 nm (for 0, 5, 10, 15 and 20 min at 22°C) was measured to monitor the enoyl-CoA reduction reaction. Kinetic values for mt/ptER and mtER were calculated using Prism version 5.0 (GraphPad Software).

**Plant strains and genetic transformations**

The Arabidopsis genetic strains used in this study included mcat-1 (Ler background, GT_5_100190), kr-1 (Col-3 background, SAIL_165_A11), mter-1 (Col-0 background, SALK_056770) and mter-2 (Col-0 background, SALK_085297), which were obtained from the Arabidopsis Biological Resource Center (Columbus, OH; http://abrc.osu.edu).

In the GFP transgene experiments, PCR inserts were obtained using the following primer pairs: pt/mtMCAT1-1179 (primers M19 and M20), pt/mtMCAT1-216 (primers M19 and M21), pt/mtMCAT205-1179 (primers M22 and M20), pt/mtKR1-957 (primers M23 and M24), pt/mtKR1-234 (primers M23 and M25), pt/mtKR214-957 (primers M26 and M24), pt/mtER1-1167 (primers M27 and M28), pt/mtER1-261 (primers M27 and M29), pt/mtER262-1167 (primers M30 and M28), mtER1-1122 (primers M31 and M32), mtER1-300 (primers M31 and M33), mtER301-1122 (primers M34 and M32), UGP3-1-600 (primers M35 and M36). PCR inserts were cloned into pENTR/D-TOPO vector (Invitrogen), and subcloned into pEarleyGate103 (Earley et al., 2006) using Gateway LR Clonase II Enzyme Mix (Invitrogen).
In the complementation transgene experiment, \( pt/mtMCAT_{1\text{-}1182} \) (primers M19-M37), \( pt/mtMCAT_{205\text{-}1182} \) (primers M22-M37), \( pt/mtKR_{1\text{-}960} \) (primers M23-M38) and \( pt/mtKR_{214\text{-}960} \) (primers M26-M38) were cloned into pENTR/D-TOPO vector and subcloned into pEarleyGate100 (Earley et al., 2006). Destination vectors were used to transform the Arabidopsis heterozygous mutant plants deficient in \( pt/mtMCAT \) and \( pt/mtKR \).

In the RNAi experiment, \( pt/mtER_{202\text{-}380} \) (primers M39-M40) was cloned into pENTR/D-TOPO and subcloned into pB7GWIWG2(II) (Karimi et al., 2002). Destination vectors were used to transform the Arabidopsis wild-type plants or \( mter-1 \) mutant plants as previously described.

Plant seedlings were grown on Murashige and Skoog agar medium at 22 °C under continuous illumination (photosynthetic photon flux density 100 \( \mu \)mol m\(^{-2}\) s\(^{-1}\)) as previously described (Jin et al., 2012). The atmospheric CO\(_2\) condition was maintained in ambient level or in the 1% CO\(_2\) level (in a growth chamber).

**Immunoblot analyses**

Total protein was extracted from 200 mg fresh aerial organs of plants at 16 DAI as described previously (Che et al., 2002). Immunoblot analysis was carried out with anti-lipoic acid antibodies (EMD Millipore, Billerica, MA) or with anti-H protein antibodies (a gift from Dr. David Oliver at Iowa State University) on 50 \( \mu \)g total protein as described previously (Ewald et al., 2007, Guan et al., 2015).

**Metabolomic analyses**

Metabolites (3-6 replicates) were extracted from 50 mg fresh aerial organs of plants (grown in a completely randomized design) at 16 DAI and analyzed using multiple analytical platforms: Agilent 7890 GC-MS system for fatty acids (Lu et al., 2008); Agilent
1200 HPLC system equipped with a fluorescence detector for amino acids (Guan et al., 2015); and Waters Xevo G2 Q-TOF MS equipped with Waters ACQUITY UPLC system for glycerolipids and chlorophylls (Okazaki et al., 2015). Log2-ratio and standard error (SE) were calculated as described previously (Quanbeck et al., 2012).

**ACCESSION NUMBERS**

AT2G30200, mt/ptMCAT; AT1G24360, mt/ptKR; AT3G45770, mtER; and AT2G05990, pt/mtER.

**SUPPLEMENTAL DATA**

Supplemental Figure S1. Arabidopsis gene candidates.

Supplemental Figure S2. Western blot analysis of the lipoylation status of indicated lipoylated proteins detected with anti-lipoic acid antibodies.

Supplemental Figure S3. Amino acid accumulation in pt/mtmcat and pt/mtkr mutants.

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FIGURE LEGENDS

Figure 1. Subcellular localization of potential mtFAS proteins. Confocal fluorescence micrographs of roots and leaf mesophyll cells, imaging the emission of GFP, MitoTracker Orange, chlorophyll auto-fluorescence, and the merged images. Fluorescence micrographs are from non-transgenic wild-type control plants (WT), transgenic plants carrying the p35S::GFP control, p35S::mtPPT1-240-GFP, or p35S::UGP3-600-GFP transgenes (A); p35S::mtER1-1122-GFP, p35S::mtER1-300-GFP, or p35S::mtER301-1122-GFP transgenes (B); p35S::pt/mtMCAT1-1179-GFP, p35S::pt/mtMCAT1-216-GFP, or p35S::pt/mtMCAT205-1179-GFP transgenes (C); p35S::pt/mtKR1-957-GFP, p35S::pt/mtKR1-234-GFP, or p35S::pt/mtKR214-957-GFP transgenes (D); p35S::pt/mtER1-1167-GFP, p35S::pt/mtER1-261-GFP, or p35S::pt/mtER262-1167-GFP (E).

Figure 2. Genetic and biochemical characterization of mtFAS gene candidates. (A) Genetic complementation of yeast mtFAS mutants (mct1, oar1 and etr1) by expression of Arabidopsis mtFAS candidate genes (AT2G30200, AT1G24360, AT3G45770, and AT2G05990); expression of the WT yeast homolog (MCT1, OAR1, and ETR1) served as a positive control. Gene expression was
transcriptionally controlled by the phosphoglycerate kinase promoter (pPGK) and terminator (tPGK). Mitochondrial pre-sequence of yeast COQ3 protein was fused to N-terminus of each protein to ensure the mitochondrial localization. All yeast strains, carrying the indicated expression cassettes were grown on media containing either glycerol or glucose as the sole carbon source, and a dilution series served as the inoculum for each strain.

(B) In vitro characterization of the catalytic capability of purified recombinant Arabidopsis mtER and pt/mtER enzymes. Substrate concentration dependence of the enoyl reductase activity was assayed with increasing concentrations of either trans-Δ^2-10:1-CoA or trans-Δ^2-16:1-CoA as substrates. The tabulated Michaelis-Menten kinetic parameters were calculated from 3 to 6 replicates for each substrate concentration.

Figure 3. In vivo physiological characterization of the pt/mtMCAT and pt/mtKR genes.

(A) Morphological phenotype of the pt/mtMCAT and pt/mtKR mutants complemented by full-length or truncated transgenes. Plants were grown in either ambient air or in a 1% CO₂ atmosphere.

(B) Western blot analysis of the H subunit of GDC detected with either anti H-protein antibodies, or anti-lipoic acid antibodies, detecting the lipoylation status of the H-protein.

(C) Glycine accumulation. Plants of the indicated genotype were grown in either ambient air or in a 1% CO₂ atmosphere.
Figure 4. Physiological characterization of the *pt/mtER* and *mtER* genes.

(A) Morphological phenotypes of single or double mutants of the *pt/mtER* and *mtER* genes. Plants were grown in either ambient air or in a 1% CO₂ atmosphere.

(B) Western blot analysis of the H subunit of GDC detected with anti H-protein antibodies, and the lipoylation status of the H-protein detected with anti-lipoic acid antibodies. Plants were grown in a 1% CO₂ atmosphere.

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Supplemental Figure S1. Arabidopsis gene candidates

Supplemental Figure S2. Western blot analysis of the lipoylation status of indicated lipoylated proteins detected with anti-lipoic acid antibodies. PDH, pyruvate dehydrogenase; KGDH, a-ketoglutarate dehydrogenase.

Supplemental Figure S3. Amino acid accumulation in *pt/mtmcat* and *pt/mtkr* mutants.
Figure 1. Confocal fluorescence micrographs

Confocal fluorescence micrographs of roots and leaf mesophyll cells, imaging the emission of GFP, MitoTracker Orange, chlorophyll auto-fluorescence, and the merged images. Fluorescence micrographs are from non-transgenic wild-type control plants (WT), transgenic plants carrying the p35S::GFP control, p35S::mtPPT1-240-GFP, or p35S::UGP31-600-GFP transgenes (A); p35S::mtER1-1122-GFP, p35S::mtER1-300-GFP, or p35S::mtER301-1122-GFP transgenes (B); p35S::pt/mtMCAT1-1179-GFP, p35S::pt/mtMCAT1-216-GFP, or p35S::pt/mtMCAT205-1179-GFP transgenes (C); p35S::pt/mtKR1-957-GFP, p35S::pt/mtKR1-234-GFP, or p35S::pt/mtKR214-957-GFP transgenes (D); p35S::pt/mtER1-1167-GFP, p35S::pt/mtER1-261-GFP, or p35S::pt/mtER262-1167-GFP (E).

(A) p35S::GFP control, p35S::mtPPT1-240-GFP, or p35S::UGP31-600-GFP transgenes
**Figure 1.** Confocal fluorescence micrographs

(B) p35S::mtER1-1122-GFP, p35S::mtER1-300-GFP, or p35S::mtER301-1122-GFP transgenes

| Roots | Leaf mesophyll cells |
|-------|----------------------|
| GFP   | GFP                  |
| MitoTracker Orange | Chlorophyll auto-fluorescence |
| Merged | Merged |

- **Roots:**
  - GFP
  - MitoTracker Orange
  - Merged

- **Leaf mesophyll cells:**
  - GFP
  - Chlorophyll auto-fluorescence
  - Merged

*Images show fluorescence micrographs of roots and leaf mesophyll cells with GFP, MitoTracker Orange, and merged images. The images depict the localization of GFP under different mitochondrial and cytosolic conditions.*
Figure 1. Confocal fluorescence micrographs

(C) p35S::pt/mtMCAT$_{1-1179}$-GFP, p35S::pt/mtMCAT$_{1-216}$-GFP, or p35S::pt/mtMCAT$_{205-1179}$-GFP transgenes
Figure 1. Confocal fluorescence micrographs

(D) p35S::pt/mtKR<sub>1</sub>-957-GFP, p35S::pt/mtKR<sub>1</sub>-234-GFP, or p35S::pt/mtKR<sub>2</sub>-957-GFP transgenes
**Figure 1.** Confocal fluorescence micrographs

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(A) Genetic complementation of yeast mtFAS mutants (mct1, oar1 and etr1) by expression of Arabidopsis mtFAS candidate genes (AT2G30200, AT1G24360, AT3G45770, and AT2G05990); expression of the WT yeast homolog (MCT1, OAR1, and ETR1) served as a positive control. Gene expression was transcriptionally controlled by the phosphoglycerate kinase promoter (pPGK) and terminator (tPGK). Mitochondrial pre-sequence of yeast COQ3 protein was fused to N-terminus of each protein to ensure the mitochondrial localization. All yeast strains, carrying the indicated expression cassettes were grown on media containing either glycerol or glucose as the sole carbon source, and a dilution series served as the inoculum for each strain.
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**Figure 2.** Genetic and biochemical characterization of mtFAS gene candidates

| Enzyme     | Enyol-CoA substrate | Km (µM)  | Vmax (µmol min$^{-1}$ mg$^{-1}$) | kcat/Km (µM$^{-1}$ min$^{-1}$) |
|------------|---------------------|----------|---------------------------------|---------------------------------|
| mtER       | trans-$\Delta^2$-10:1 | 8.8 ± 0.9 | 4.8 ± 0.1                       | 0.87 ± 0.09                     |
| mtER       | trans-$\Delta^2$-16:1 | 12.1 ± 2.4| 4.4 ± 0.2                       | 0.58 ± 0.12                     |
| pt/mtER    | trans-$\Delta^2$-10:1 | 10.9 ± 1.6| 6.8 ± 0.3                       | 1.11 ± 0.16                     |
| pt/mtER    | trans-$\Delta^2$-16:1 | 8.3 ± 2.2 | 8.8 ± 0.6                       | 1.88 ± 0.67                     |
Figure 3. *In vivo* physiological characterization of the *pt/mtMCAT* and *pt/mtKR* genes

(A) Morphological phenotype of the *pt/mtMCAT* and *pt/mtKR* mutants complemented by full-length or truncated transgenes. Plants were grown in either ambient air or in a 1% CO₂ atmosphere.

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