Identification of a Pair of Phospholipid:Diacylglycerol Acyltransferases from Developing Flax (Linum usitatissimum L.) Seed Catalyzing the Selective Production of Trilinolenin

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*Running Title: α-linolenic acid-specific PDATs from flax

Keywords: Lipid Metabolism; Triacylglycerol; Molecular Genetics; Metabolic Engineering; Plant Molecular Biology; Phospholipid:Diacylglycerol Acyltransferases; Flax; α-Linolenic Acid

**Background:** Triacylglycerol (TAG) can be formed via an acyl-CoA-dependent or acyl-CoA-independent pathway.

**Results:** Overexpressing particular flax phospholipid:diacylglycerol acyltransferase (PDAT) genes in yeast and Arabidopsis resulted in an enhanced proportion of α-linolenic acid (ALA) in TAG.

**Conclusion:** Certain PDATs have the unique ability to efficiently channel ALA into TAG.

**Significance:** The identified PDATs will benefit the future projects aimed at producing oils with enhanced polyunsaturated fatty acid content.

**ABSTRACT**

The oil from flax (Linum usitatissimum L.) has high amounts of α-linolenic acid (ALA; 18:3 cis9,12,15) and is one of the richest sources of omega-3 polyunsaturated fatty acids (ω-3 PUFAs). To produce approximately 57% ALA in triacylglycerol (TAG), it is likely that flax contains enzymes that can efficiently transfer ALA to TAG. To test this hypothesis, we conducted a systematic characterization of TAG-synthesizing enzymes from flax. We identified several genes encoding acyl CoA:diacylglycerol acyltransferases (DGATs) and phospholipid:diacylglycerol acyltransferases (PDATs) from the flax genome database. Due to recent genome duplication, duplicated gene pairs have been identified for all genes except DGAT2-2. Analysis of gene expression indicated that two DGAT1, two DGAT2 and four PDAT genes were preferentially expressed in flax embryos. Yeast functional analysis showed that a DGAT1, a DGAT2, and two PDAT enzymes restored TAG synthesis when produced recombinantly in yeast H1246 strain. The activity of particular PDAT enzymes (LuPDAT1 and LuPDAT2) was stimulated by the presence of ALA. Further seed-specific expression of flax genes in Arabidopsis thaliana indicated that DGAT1, PDAT1 and PDAT2 had significant effects on seed oil phenotype. Overall, this study indicated the existence of unique PDAT enzymes from flax that are able to preferentially catalyze the synthesis of TAG containing ALA acyl moieties. The identified LuPDATs may have practical applications for increasing the accumulation of ALA and other polyunsaturated fatty acids in oilseeds for food and industrial applications.
makes flax oil suitable for domestic and industrial coatings such as linoleum, varnishes and paints (1). In addition, ALA is an essential fatty acid for human and a precursor for nutritionally beneficial very long chain omega-3 polyunsaturated fatty acids (VLC-α-3-PUFA, ≥ 20 carbons) (2, 3).

Efforts to characterize the molecular basis of high ALA content in flax have focused largely on the characterization of fatty acid desaturases (FAD). Two genes encoding Δ15 desaturases (LuFAD3A and LuFAD3B) have been identified in flax (4). Ethyl methanesulfonate (EMS)-generated point mutations in LuFAD3A and LuFAD3B led to a reduction in ALA content to approximately 1-2%, suggesting that these two genes encoded the main desaturases responsible for the synthesis of ALA in flax (4-6). Generally, there is a good correlation between the expression of FAD3 genes with the accumulation of ALA in flax and other plant species (7). However, along with desaturases, many other enzymes can contribute to the flux of ALA to storage lipids. Essentially, ALA has to be efficiently transferred from the desaturation product (sn-2-linolenolyl-phosphatidylcholine, PC) to the substrates for triacylglycerol (TAG) synthesis and this process can take different biochemical routes. ALA synthesized on PC can enter the acyl-CoA pool by either the reverse action of acyl-CoA:lyso phosphatidylcholine acyltransferase (LPCAT) (8) or the combined action of phospholipase A2 (PLA2) and long-chain acyl-CoA synthetase. 2-Linolenolyl-sn-PC can also be potentially converted into 2-linolenolyl-sn-diacylglycerol (DAG) by the catalytic action of phospholipase C or phospholipase D (9) together with phosphatidic acid phosphatase. In addition, the enzyme phosphatidylglycerol cholinephosphotransferase (PDCT) catalyzes the transfer of a phosphocholine head group from PC to DAG in a symmetrical reaction (10) which could potentially produce DAG enriched in ALA for the synthesis of TAG.

TAG can be formed via an acyl-CoA-dependent or acyl-CoA-independent process. The final step of the acyl-CoA dependent pathway, also known as Kennedy pathway (11), is catalyzed by acyl-CoA:diacylglycerol acyltransferase (DGAT) which uses acyl-CoA as acyl donor to convert DAG to TAG. At least four different types of DGAT have been identified in plants (12-15). DGAT1 has been proposed to exert dominant control over TAG accumulation in many oilseeds (16-19). DGAT2 was first identified in the oleaginous fungus Umbelopsis (formerly Mortierella) ramanniana and shows no sequence homology to DGAT1 (15). DGAT1 and DGAT2 appear to localize to different subdomains of endoplasmic reticulum (ER) and have been suggested to have nonredundant functions in the production of TAG (20). The third type of DGAT (DGAT3) is a soluble enzyme isolated from developing peanut cotyledons and it differs from DGAT1 and DGAT2, which are membrane-bound (13). Another soluble enzyme with DGAT activity known as defective in cuticular ridge (DCR) has been identified in Arabidopsis thaliana (21).

The acyl-CoA independent pathway of TAG synthesis is characterized by the enzymatic action of phospholipid:diacylglycerol acyltransferase (PDAT) that transfers the fatty acyl moiety from sn-2-position of a phospholipid to the sn-3-position of sn-1, 2-DAG (22). Genes encoding PDAT were first reported in yeast (22). Two homologs of yeast PDAT have been identified in Arabidopsis named PDAT1 (At5g13640) and PDAT2 (At3g4480) (23) and three putative PDAT genes have been identified in the castor bean genome (24).

The existence of specialized acyltransferases has been observed in other plant species. DGAT2 from tung tree (Vernicia fordii) and castor bean (Ricinus communis) display substrate preference for unusual fatty acids and are predominantly involved in the incorporation of these fatty acids into seed oils (20, 25). Also, a ricinoleate-specific PDAT from castor bean (RePDAT) has been reported (24). Co-expression of RePDAT with castor fatty acid hydroxylase in Arabidopsis resulted in a relative increase of 58% hydroxy fatty acids in seeds. Considering the natural occurrence of evolved forms of acyltransferases that are selective for unusual fatty acids, we hypothesized that similar mechanisms take place in flax. In spite of the wide range of applications for flax oil, many components involved in TAG biosynthesis have not been characterized at the molecular genetic level. Here we provide this characterization and demonstrate that two pairs of PDAT genes encoding enzymes utilize
preferentially substrates containing ALA, and more importantly, one of the pairs (LuPDAT1 and LuPDAT5) has embryo-preferred expression pattern and appears to contribute mainly to the synthesis of trilinolenin in flax seeds.

EXPERIMENTAL PROCEDURES

Plant Material and Growth conditions—Flax (Linum usitatissimum L.) was grown in the growth chamber with 16 h light at 21°C and 8 h dark at 18°C. Seeds were planted in 1 Gallon pots containing Metro Mix soil (Greenleaf Products Inc., CA). Plants were fertilized with 17-5-19 (125g/100L) and 12-2-14 (83g/100L) once a week before flowering. From the flowering stage, plants were fertilized weekly with 17-5-19 (280g/100L) till harvest. Individual flowers were tagged at anthesis, and embryos at different stages of development were collected in liquid N2 and stored at -80°C. Arabidopsis wild-type Columbia and mutant line AS11 were obtained from the Salk Institute via the ABRC (Ohio State University, Columbus, OH). Arabidopsis seeds in plots or plates were cold-treated at 4°C in the dark for 3 days and then placed into a controlled growth chamber with a constant temperature of 20°C and 16 h photoperiod.

Identification and isolation of candidate genes—In general, all primers are summarized in Supplemental Table 1. Cloning integrity was confirmed by sequencing at each step.

A BLAST analysis (26) was conducted against the flax genomic database (27) by using AtDGAT1, AtPDAT1, AtDCR and tung tree VfDGAT2 as the protein query. The theoretical molecular weight and isoelectric point values were calculated using the algorithm provided in http://web.expasy.org/compute_pi/. To isolate the target genes, total RNA was extracted from the embryo of flax (cultivar AC Emerson) 12 days post anthesis (DPA) using Plant RNeasy plant mini kit (Qiagen, CA) as described by the manufacturer. The First stand of the cDNA was synthesized using SuperScript III reverse transcriptase (Invitrogen, CA) according to the protocol provided by the supplier. The target genes were amplified from the resulting cDNA as the template for 30 cycles of PCR amplification using Platinum Taq DNA Polymerase High Fidelity (Invitrogen, CA). PCR was performed under the following temperature cycle program: 95°C for 2 min; 30 cycles of denaturation (95°C, 20s), annealing (55°C, 15s), and extension (68°C, 2.5 min); and a final extension at 68°C for 2 min. To amplify LuDGAT2-1, LuDGAT2-3, LuPDAT2, and LuPDAT6, the forward primers contained a specific restriction site (underlined) and a Kozak translation initiation sequence (italic) in order to improve the translation of the protein (Supplemental Table 1). A specific restriction site was also introduced in the reverse primer (underlined). The PCR products were cloned into the pYES vector collinear to the GAL1 promoter inducible by galactose. The pYES is a modified pYES2.1/V5-HIS vector (Invitrogen, CA) constructed in our lab, which contains more restriction sites in its multiple cloning site (MCS). For LuDCR1 and LuPDAT1 that could not be amplified by these specially designed primers, the internal primers were used for amplification. For LuDGAT2-2 and LuPDAT5, an internal forward primer and a reverse primer spanning the 3'-untranslated region (3'UTR) were used for amplification. The PCR products were subcloned into pYES2.1/V5-HIS vector using pYES2.1 TOPO kit (Invitrogen, CA) as demonstrated by the supplier. To construct the co-expression vectors, the LuFAD2-1 (28) and LuFAD3B (4) genes were first amplified using PCR with appropriate primers that allowed to add specific restriction sites (underlined, Supplemental Table1) to the ends of amplified products and then inserted into MCS1 and MSC2 of the pESC-URA expression vector (Agilent Technologies, CA), yielding LuFAD2-1-FAD3B/pESC plasmid. The ADH1 terminator: LuFAD2-1: ProGAL10: ProGAL1: LuFAD3B: CYC1 terminator expression cassette of LuFAD2-1-FAD3B/pESC was then excised and subcloned into the recombinant pYES plastids containing LuPDAT1, LuPDAT2 or LuDGAT1 through one-step, isothermal assembly method described by Gibson (29). The resulting plasmids were referred to as LuFAD2-1-FAD3B-PDAT1/pYES, LuFAD2-1-FAD3B-PDAT2/pYES and LuFAD2-1-FAD3B-DGAT1-1/pYES.

Real-Time PCR quantification—Total RNA was isolated from vegetative tissues (stems, leaves, apexes, roots), reproductive tissues (flowers), developing embryos (8, 11, 14, 16, and 20 DPA) and seeds (4, 25 and 40 DPA) of flax cultivar CDC Bethune using RNeasy plant mini kits (Qiagen, CA). Due to technical difficulties in
separating embryos from other seed components at 4, 25 and 40 DPA, the whole seeds at these three stages were used for gene expression analysis. The cDNA was synthesized using SuperScript III reverse transcriptase (Invitrogen, CA). Gene specific primers and probes were designed using Primer Express software (Applied Biosystems) and the sequences are listed in Supplemental Table1. 5’FAM/3’quencher-labelled probes were used to assay the genes involved in TAG synthesis, and probes for reference genes were labelled with 5’VIC/3’quencher. TaqMan-based qRT-PCR assays were performed using Fast Advance Master mix (Invitrogen, CA) on the ABI PRISM 7900HT Real-Time PCR system (Applied Biosystems). PCR efficiency for each amplicon was calculated using a dilution series of a single cDNA sample over several log concentrations. According to the recommendation of Huis et al. (30), two internal reference genes, encoding glyceraldehyde 3-phosphate dehydrogenase (GAPDH) and ubiquitin extension protein (UBI2), were used. The relative expression level was calculated using the comparative Ct method after normalizing to controls using the reference genes.

**Heterologous expression in yeast mutant H1246**—In general, the constructed recombinant plasmids were transformed into the quadruple mutant strain *S. cerevisiae* H1246 by using the LiAc/SS carrier DNA/PEG method (31) and transformants were selected on minimal medium plates lacking uracil. The recombinant yeast cells were first grown in liquid minimal medium with 2% [w/v] raffinose overnight and then inoculated at a starting OD of 0.1 in minimum medium containing 2% [w/v] galactose and 1% [w/v] raffinose (referred to as induction medium) to induce gene expression.

For heterologous expression of single flax genes, yeast cells were grown at 30°C and yeast transformed with pYES LacZ was used as a control. For the feeding experiments, free fatty acids, including oleic (OA; 18:1\textsuperscript{cis9}), linoleic acid (LA; 18:2\textsuperscript{cis9,12}), ALA, stearidonic acid (SDA; 18:4\textsuperscript{cis6,9,12,15}), dihomo-\(\gamma\)-linolenic acid (DGLA; 20:3\textsuperscript{cis8,11,14}), \(\gamma\)-linolenic acid (GLA; 18:3\textsuperscript{cis6,9,12}), arachidonic acid (AA; 20:4\textsuperscript{cis5,8,11,14}), eicosatrienoic acid (ETA; 20:3\textsuperscript{cis9,11,14}), eicosapentaenoic acid (EPA; 20:5\textsuperscript{cis5,8,11,14,17}) and docosahexaenoic acid (DHA; 22:6\textsuperscript{cis4,7,10,13,16,19}), were dissolved at 0.5 M in ethanol. The fatty acid solutions were first dissolved in 0.06% [v/v] tyloxapol (Sigma, CA) and then mixed with induction medium. Tyloxapol is a non-ionic surfactant which was used to disperse the fatty acids into the medium. Yeast was induced in induction medium with 100 μM fatty acid supplementation.

To evaluate the concentration effect of exogenously provided ALA on total TAG content and composition in yeast expressing LuPDAT1 and LuPDAT2, the recombinant yeast cells were cultivated in induction medium supplemented with different concentration of ALA (0 to 300 μM) and harvested at the same growth stage (OD\textsubscript{600nm} = 6.5±0.05). Approximately 30 mg of dry yeast were weighed out and used for lipid analysis after 16 h freeze-drying. One hundred micrograms of the TAG internal standard, triheptadecanoin (C17:0 TAG), were added to each sample before lipid extraction.

For growth curve construction, yeast cells were grown first in minimum medium containing 2% [w/v] raffinose, and then in induction medium supplemented with three different ALA concentrations (0, 100 μM and 1 mM). Aliquots (100 μl) were withdrawn from the culture at intervals of 6 or 12 h and diluted 1:10 with sterile water. One hundred microliters of the diluted cells were transferred to a 96-well clear, flat bottom, polystyrene microplate (UNIPLATE, Whatman). The yeast growth (OD\textsubscript{600nm}) was measured using the Synergy H4 Hybrid multi-mode microplate reader (Biotek Instrument, Inc., USA).

Growth phenotype were further analyzed by first culturing recombinant yeast cells in minimum medium containing 2% [w/v] raffinose, and then spotting 2.5 μl of cell suspension serially diluted to OD\textsubscript{600nm} values of 1, 10\textsuperscript{-1}, 10\textsuperscript{-2}, 10\textsuperscript{-3}, and 10\textsuperscript{-4}, onto agar plates containing induction medium supplemented with different concentration of ALA (0, 100 μM and 1 mM). ALA was first dissolved at 0.5 M in ethanol, and then diluted in a warm medium containing 0.01% [v/v] tyloxapol immediately before plating. The same amount of ethanol and tyloxapol was added to the plates without fatty acid supplementation. Yeast strains BY4742 (wild-type) and H1246 transformed with empty pYES vector were used as positive and negative controls, respectively. Plates were incubated at 30°C for 2 days.
For the co-expression study, yeast cells transformed with LuFAD2-1-FAD3B-PDAT1/pYES, LuFAD2-1-FAD3B-PDAT2/pYES or LuFAD2-1-FAD3B-DGAT1-1/pYES plastid were inoculated in the same induction medium for 3 days at 20°C before cell harvest. Yeast cells transformed with LuFAD2-1-FAD3B/pYES were used as a control.

Western blot analysis – To produce N-terminal-tagged constructs, the complete ORFs of three LuPDATs (LuPDAT1, LuPDAT2 and LuPDAT6) were first amplified by the primers listed in Table1 and then subcloned into the pYES2/NT-C expression vector (Invitrogen). For the generation of C-terminal-tagged constructs, the stop codons of the LuPDATs in the recombinant pYES plasmids were mutated by using the Quickchange mutagenesis kit (Stratagene). Yeast microsomes were prepared as previously described (32). Briefly, the recombinant yeast cells were first cultivated in liquid minimal medium with 2% [w/v] raffinose and then inoculated at a starting OD of 0.4 for 12h. The cells were harvested, washed, and then resuspended in the homogenization buffer (20mM Tris-HCl pH 7.9, 10mM MgCl₂, 1mM EDTA, 5% (v/v) glycerol, 300mM ammonium sulphate, 2mM dithiotreitol). In the same buffer, the cells were broken mechanically with 1/3 volume of 0.5 mm glass beads by a bead beater (Biospec Bartlesvile, USA). To remove the cell debris and glass beads, the homogenate was centrifuged at 10,000 g at 4°C for 20 min. The recovery supernatant was further centrifuged at 48,000 g at 4°C for 1 h. The resulting microsomal pellets were dissolved in 3mM imidazole buffer (pH 7.4) containing 125mM sucrose. Equivalent amount of microsomal proteins (15μg, as determined by Bio-Rad Bradford assay) were separated by 4-12% gradient tris-glycine gels and then blotted onto nitrocellulose membranes. The target N-terminal-tagged proteins were detected using anti-HisG-HRP antibody (Invitrogen). The bounded antibodies were detected using SuperSignal Fermo Chemiluminescent Substrate (Pierce). Equal protein loading was ensured by determining the level of constitutively expressed chaperone Kar2p using a rabbit polyclonal anti-Kar2p (Santa Cruz Biotechnology) as the primary antibody followed by incubation with HRP-goat anti-rabbit IgG(H+L) secondary antibody (Invitrogen).

Nile red fluorescence detection of neutral lipids in yeast – The Nile red fluorescence assay was conducted as described (32, 33). Briefly, the recombinant yeast cells were cultivated in induction medium with or without ALA supplementation and harvested at the stationary growth stage. Cells with ALA feeding were washed three times with deionized water before Nile red measurement. One hundred microliters aliquots of induced yeast cell culture were transferred to a 96-well dark flat-bottom plate (UNIPLATE, whatman). The background fluorescence was measured using the Synergy H4 Hydrid multi-mode microplate reader (Biotek Instument, Inc., USA) with the emission and excitation filter setting at 485 and 538 nm, respectively. Five microliters of freshly prepared methanolic Nile red solution (0.1 mg/ml) was added first and mixed with yeast culture. The second fluorescence intensity value was measured under the same conditions. The final Nile red value was calculated by subtracting the first measurement from the second measurement (AF), and then dividing by the corresponding OD₆₀₀nm value.

Yeast lipid extraction and analysis – Yeast lipid extraction was performed using the method described previously (32, 34). Briefly, induced yeast cultures at the stationary growth stage were harvested by centrifuge, washed and resuspended in 1 ml of 0.9% [w/v] sodium chloride (NaCl). Cells were homogenized with an equal volume of glass beads (0.5 mm) by vigorously vortexing for 2 min. Lipids were extracted by the chloroform:methanol:0.9% aqueous NaCl (2:1:0.9, v/v/v) method. The chloroform phase (lower phase) was collected, dried under nitrogen and resuspended in 30 μl of chloroform. The extracted lipids were resolved on the thin layer chromatography (TLC) plates (SIL G25, 0.25 mm, Macherey–Nagel, Germany) with the solvent system hexane/diethyl ether/acetic acid (80:20:1). The developed plate was visualized with 3% cupric acetate [w/v] and 8% phosphoric acid [v/v] followed by charring at 280°C for 20 min. The TAG bands were identified according to triolein and trilinolenin standards.

Fatty acid analysis was performed by gas chromatography (GC)/mass spectrometry (MS) as described previously (35). Briefly, the extracted
lipids were developed on TLC plate by using the same solvent system but visualized under UV after spraying with 0.05% primuline solution. The bands corresponding to TAG were scraped and transmethylated with 5% [w/v] sodium methoxide (NaOMe) at room temperature for 30 min. The resulting fatty acid methyl esters (FAMEs) were extracted with two portions (2 ml each) of hexane. The hexane phases were pooled together and dried under a stream of nitrogen and immediately resuspended in 250 μl of iso-octane with 0.1 mg/ml C21:0 methyl ester standard. FAMEs were subjected to an Agilent 6890N GC equipped with DB-23 capillary column (30 m x 0.25 mm x 0.25 μm) and a 5975 inert XL Mass Selective Detector. The following temperature program was applied: 165°C hold for 4 min, 10°C/min to 180°C, hold 5 min and 10°C/min to 230°C hold 5 min.

Overexpression of LuDGATs and LuPDATs in Arabidopsis—Agrobacterium tumefaciens strain GV3101 and pGreen/pSoup based dual binary vectors (36) were used for Arabidopsis transformation. The coding regions of target genes were amplified using recombinant pYES plasmids as template with specially designed primers for each gene (see Supplemental Table 1). The amplicons were cloned into the pGreen vector under the control of the seed specific napin promoter. The resulting construct and the helper plasmid pSoup were co-transformed into A. tumefaciens GV3101 by electroporation (37). Agrobacterium strains containing the pGreen/pSoup dual binary vectors were used to transform the Arabidopsis wild-type (Columbia) and mutant line AS11 by the floral dipping method (37). Plants transformed with an empty vector pGreen were used as controls. T1 seeds of transgenic plants were selected on half-strength Murashige and Skoog (MS) agar plates supplemented with 80μM herbicide phosphinothricin. Transformants were then transferred to soil and grown to maturity to produce T2 seeds. The presence of the target genes was confirmed by gene-specific PCR analysis using DNA extracted from T2 young leaf tissue as template. T2 seeds were collected and used for total lipid and fatty acid analysis.

Analysis of Arabidopsis seed oil—Total lipid content and the fatty acid composition of T2 seeds were determined by GC. Approximately 10 mg seeds per replicate were weighted out after stabilizing seed moisture content in desiccators for 72h. One hundred micrograms triheptadecanoin (C17:0 TAG) were used as a TAG internal standard. Seeds were subjected to treatment with 2 ml of 3N methanolic-HCL and heated at 80°C for 16 h. The extracted FAMEs were suspended in 1.5 ml of iso-octane with 0.1 mg/ml C21:0 methyl ester standard and analyzed by GC-MS using the same column and temperature gradient. Total lipid content was determined by multiplying the peak-area ratio of the total fatty acid and the internal standard by the initial internal standard amount.

Flax seed oil and fatty acid composition analysis—Flax embryos of cultivar CDC Bethune at different developmental stages (8, 11, 14, 16 and 20 DPA) and seeds (25, 30 and 40 DPA) were collected, measured for fresh weight, immersed in liquid nitrogen, and then placed at -80°C for storage. Frozen embryos and seeds (approximately 15 to 75 mg fresh weight per biological replicate) were freeze-dried for four days and then homogenized in a mortar and pestle in the presence of liquid nitrogen. Homogenates were preceded for lipid extraction and fatty acid analysis as described in Arabidopsis seed oil analysis. The oil and ALA content were calculated on a fresh weight basis.

Phylogenetic analysis—The multiple sequence alignments of PDAT, DCR, DGAT1 and DGAT2-like family proteins were generated using the ClustalW module (38) within MEGA5 (http://www.megasoftware.net/mega.html) (39) with the default parameters (gap penalty, 10.0; gap length penalty, 0.2; Gonnet matrix). The phylogenetic trees were constructed using the same software with the following parameters: neighbor-joining method, Poisson model, complete deletion and bootstrap (1000 replicates). Numbers above branches indicate the percentage of bootstrap values.

Accession number—Accession numbers of protein sequences from the Arabidopsis Genome Initiative or EMBL/GenBank database or flax genome database (http://www.phytozome.net/flax) used in the current study are given in Supplemental Table 2.

RESULTS

Identification and isolation of genes encoding TAG-synthesizing enzymes from flax—The recently
completed flax genome sequence database (27) provided the starting point for identifying flax genes homologous to known enzymes involved in the final step of TAG synthesis. Using basic local alignment search tool (BLAST), we identified three DGAT2s (LuDGAT1-1, LuDGAT2-2 and LuDGAT2-3), two DCRs (LuDCR1 and LuDCR2) and six PDATs (LuPDAT1, LuPDAT2, LuPDAT3, LuPDAT4, LuPDAT5 and LuPDAT6) in the flax genome. In addition, we had previously reported a DGAT1 (here referred to as LuDGAT1-1) (32) and now identified a second gene (LuDGAT1-2) in the flax genome. General information about the identified genes is listed in Table 1 while the gene structures were represented in Figure 1. Both LuDGAT1 genes contain 15 exons and like DGAT1 from other plant species, the first exon is the longest and encodes a hydrophilic domain (40). LuDGAT2-1 and LuDGAT2-3 are composed of 8 exons each and share a more similar gene structure than with LuDGAT2-2, which contains only 6 exons. Interestingly, LuDGAT2-3 is located adjacent to LuDGAT2-2 in the genome (Supplementary Figure 1). Six LuPDATs can be divided into three groups based on polypeptide sequences (Supplementary Figure 2A). The genes within each group have a similar gene structure, where LuPDAT3 and LuPDAT6 genes contain 5 exons, one less than in LuPDATs of the other two groups. Phylogenetic analyses of the predicted polypeptides for each class indicate a close relationship with known TAG-synthesizing enzymes from other species (Supplementary Figures 2A to 2D). Therefore, we conclude that the identified genes comprise the most probable candidates for encoding TAG-synthesizing enzymes from flax. In addition, the identification of gene pairs for all genes, except to LuDGAT2-2, is consistent with the findings that suggest whole-genome duplication in flax (27). The two genes in a gene pair have a very high degree of sequence identity at both nucleotide and amino acid levels (Table 2).

Further genomic analysis revealed that LuDGAT1-1 and LuPDAT5 are located close together on the same chromosome and the other member of the gene pair (LuDGAT1-2 and LuPDAT1) are similarly arranged on a different chromosome (Supplementary Figure 3).

Several DGAT1, DGAT2, PDAT genes are expressed preferentially in seeds and correlate with oil accumulation—To investigate the potential physiological roles of all candidate genes, we studied the expression patterns of LuDGAT1, LuDGAT2, LuDCR and LuPDAT in a range of organs including vegetative tissues, reproductive tissues and mature seeds (40 DPA). Genes encoding LuDGAT1 or LuDGAT2 had significantly higher expression in seeds than in other vegetative tissues (Figures 2A to 2C). In the case of LuDCR, the expression level was significantly higher in vegetative tissues than that in seeds, suggesting that this gene might play a minor role in TAG biosynthesis (Figure 2D). LuPDAT genes displayed two distinct expression profiles. Transcripts of LuPDAT1/LuPDAT5, LuPDAT3 and LuPDAT6 (Figures 3A to 3C) were detected preferentially in seeds and to a lesser extent in other tissues, and the opposite was observed for LuPDAT2 and LuPDAT4 (Figures 3D and 3E). This result suggests that two groups of PDATs might have unrelated physiological function roles in flax. An attempt to individually detect the transcript of LuPDAT5 failed, but we were able to isolate the cDNA of LuPDAT5 using the reverse primer targeting 3’UTR. Due to the very high level of sequence identity between LuDGAT1 and LuPDAT5 (97%), the transcripts of LuPDAT1 and LuPDAT5 were detected together. The transcripts for LuDGAT2-2 could not be detected in any of the tested plant tissues. Attempts to isolate the corresponding cDNAs were also unsuccessful, suggesting that LuDGAT2-2 may not be expressed in the tested tissues.

These experiments indicated that LuDGAT1, LuDGAT2-1, LuDGAT2-3, LuPDAT1/PDAT5, LuPDAT3 and LuPDAT6 are preferentially expressed in seeds than other tissues. To further decipher the role of these seed-preferred genes in oil synthesis, we analyzed the relationship between gene expression and seed oil and ALA accumulation patterns throughout seed development (Figures 4A to 4I). During seed development, the oil content on a fresh weight basis fit a sigmoidal curve ($R^2 = 0.969$) with the rapid phase of oil accumulation occurring between 8 and 20 DPA (Figure 4B). The ALA content on a fresh weight basis increased steadily until about 16 DPA (Figure 4C). The expression of DGATs and PDATs displayed dissimilar temporal regulation. The level of DGATs transcripts (Figures 4D to 4F) peaked during the late stages of seed development,
when the rate of oil and ALA accumulation already reached a plateau. However, the highest expression of PDATs (Figures 4G to 4I) was found in the early stages of seed development, during the period of active seed oil and ALA accumulation. This result indicated that DGATs and PDATs might contribute differently to TAG synthesis during flax seed development.

DGATs and PDATs from flax are functional TAG-synthesizing enzymes—The ability of the polypeptides encoded by LuDCR1, LuDGAT and LuPDAT genes to catalyze the synthesis of TAG was investigated through a functional assay in yeast. Because genes in a gene pair show high sequence similarity, one gene of each pair, including LuDGAT1-1, LuDGAT2-3, LuDCR1, LuPDAT1, LuPDAT2 and LuPDAT6, were chosen for this assay. The cDNAs of selected genes were isolated from 12 DPA flax developing embryos. The respective open reading frames (ORFs) were expressed in the quadruple mutant strain Saccharomyces cerevisiae H1246, which lacks four genes DGA1, LRO1, ARE1 and ARE2 encoding DGAT1, PDAT1, ASAT1 (acyl-CoA:sterol acyltransferase 1) and ASAT2, respectively. With the disruption of these four genes, S. cerevisiae H1246 is unable to synthesize TAG and sterol esters (41), thus highlighting the production of TAG that results from the activity of recombinant flax enzymes. It has been previously demonstrated by our group that yeast H1246 expressing LuDGAT1 has the ability to synthesize TAG (32), thus it was used as a positive control. Yeast transformed with the pYES vector coding for the bacterial protein LacZ was used as a negative control. Production of TAG in recombinant yeast was detected by the Nile red fluorescent assay as well as by TLC. The results of Nile red assay (Figure 5A) showed that besides LuDGAT1-1, LuPDAT2 was the only enzyme that had significant activity in yeast. Because flax oil contains a substantial amount of ALA, which is absent in yeast, we hypothesized that some of these enzymes might be selective for substrates containing ALA. Yeast is capable of importing fatty acids from the environment and converting them to their respective acyl-CoA derivatives (42). Therefore, the expression of the genes was induced in the presence of exogenously added ALA. Nile red results indicated that all enzymes displayed an enhanced activity in the presence of ALA with the exception of LuDCR1 (Figure 5A). The Nile red assay is a high throughput technique for quantitative measurement of neutral lipids; however, it is unable to distinguish different neutral lipid classes. Therefore, TLC was used to further confirm the production of TAG in recombinant H1246 yeast cells. There was a good correlation between the results from Nile red and TLC analysis (Figures 5A to 5D). Two recombinant PDATs (LuPDAT1 and LuPDAT2) restored TAG synthesis in H1246 when culturing yeast in the presence of ALA (Figure 5C). The recombinant LuDGAT2-3 also produced TAG, but at significantly lower levels than those found in LuPDAT1, LuPDAT2 and LuDGAT1-1 (Figure 5C and 5D). This analysis confirmed that most of the flax genes can be functionally expressed in yeast and their encoded enzymes are able to complement the TAG synthesis mutations in H1246.

LuPDAT1 and LuPDAT2 have unique ALA-selectivity - When ALA was exogenously added to induction medium, yeast cells expressing LuPDAT1 and LuPDAT2 produced multiple TAG bands on the TLC plate (Figure 5C). This was not observed in yeast expressing LuDGAT1 (Figure 5D). Comparison of the migration rates of trilinolenin and triolein standards under our TLC conditions (Supplemental Figure 4) suggested that the observed multiple TAG bands could be explained by the different composition of fatty acids in these TAGs. Therefore, it is plausible to assume that in the presence of ALA, LuPDAT1 and LuPDAT2 produce TAG with a specific fatty acid composition. Indeed, GC analysis of TAG bands with different migration rates indicated that the TAG band with lower migration rate was composed uniquely of ALA, while the band with higher migration rate contained 65 mol% and 49 mol% ALA for LuPDAT1 and LuPDAT2, respectively (Figure 6). This result indicated that both LuPDAT1 and LuPDAT2 have the ability to synthesize trilinolenin, which is the major molecular species of TAG in flax oil (43). This experiment also suggested that these two PDATs might be selective for substrates containing ALA, or have a preference for utilizing exogenously imported acyl moieties into yeast. To investigate the origin of this effect, the recombinant yeast strains were cultivated in the presence of OA or...
LA. As shown in Figure 5, yeast cells expressing LuPDAT1 and LuPDAT2 produced much weaker TAG bands in these medium conditions (Figure 5E and 5F) than in medium supplemented with ALA (Figure 5C). It is also worth noting that substantial amount of TAG was produced by LuDGAT1-1 under all conditions (Figure 5D). Overall, these results suggest that both LuPDAT1 and LuPDAT2 have a high preference for ALA-containing substrates. Fatty acids commonly found in yeast, including stearic, palmitic, palmitoleic acids and OA are inefficient substrates for these PDATs.

LuPDAT1 and LuPDAT2 have the similar preference for ALA-containing substrates – The free fatty acid feeding study revealed that both LuPDAT1 and LuPDAT2 are uniquely ALA-selective. We are also intrigued by the finding that, when ALA was provided exogenously, LuPDAT1 catalyzed the synthesis of a higher amount of trilinolenin with lower total TAG as compared with LuPDAT2 (Figure 6). To further compare these two enzymes, we first used Western blot to assess the protein expression levels of these PDATs in H1246. Both enzymes with C-terminal fusion tag had reduced TAG-forming ability. Therefore, N-terminal-tagged proteins were used for detection. Western blot analysis revealed that the band corresponding to PDAT2 was markedly stronger than that corresponding to PDAT1 (Figure 7A), suggesting that H1246 was able to accumulate much higher levels of PDAT2 than PDAT1. It should also be noted that PDAT6 was expressed at a similar protein level to PDAT2, but it failed to complement TAG synthesis in H1246 under all conditions.

It is likely that the higher level of PDAT2 produced in yeast may enhance the rate of channeling ALA into TAG, thus depriving the exogenously introduced ALA in the DAG/PC pool and leading to the formation of TAG with lower content of ALA as compared with PDAT1. To address this possibility, we further investigate the concentration effect of exogenously provided ALA on total TAG content and composition in yeast expressing LuPDAT1 and LuPDAT2 by culturing the recombinant cells in medium supplemented with 0 to 300 μM of ALA (Figure 7 B-E). As show in Figure 7C, both PDAT1 and PDAT2 were able to catalyze the production of TAG with up to 90% ALA, but PDAT2 reached the plateau at much higher concentration of ALA, suggesting that the increased abundance of recombinant PDAT2 would require more ALA-containing substrates to obtain TAG with the same level of ALA as produced by PDAT1. In addition, an increase in ALA concentration from 0 to 300 μM led to approximately 168-fold and 44-fold increases in total TAG content for yeast expressing LuPDAT1 and LuPDAT2, respectively (Figure 7D). Further quantifying the fatty acid species in TAG revealed that elevated TAG content in these cells was mainly because of the increased amount of ALA (Figure 7E), suggesting that both enzymes may be more active in the presence of ALA. Taken together, we observed a similar response of LuPDAT1 and LuPDAT2 to an increase of ALA concentration, indicating that LuPDAT2 has similar ALA-selectivity as compared with LuPDAT1. The differential levels of protein accumulation, if not entirely, at least partially account for the observed differences in TAG-forming ability between PDAT1 and PDAT2.

Exogenously provided free fatty acids inhibit growth of the recombinant yeast mutants and this inhibitory effect can be rescued in a TAG accumulation-dependent manner– We observed that the recombinant yeast mutant displayed a different growth rate upon exposure to free fatty acids. To assess the effect of exogenously provided ALA on recombinant yeast cell growth, we evaluated the cell growth in induction medium supplemented with 0, 100 μM and 1mM of ALA by both growth curves analysis and the serial dilution plate assay. BY4742 (wild-type) and H1246 harboring the empty vector pYES were used as positive and negative controls, respectively. As shown in Figure 8, the presence of ALA produced a concentration-dependent inhibitory effect on recombinant yeast growth. Cells with higher TAG-synthesizing ability showed an earlier log phase and earlier appearance on plates, suggesting that inhibitory effect of ALA was rescued in a TAG accumulation-dependent manner. This result is in agreement with early findings, which reported that TAG biosynthesis plays a crucial role in detoxifying the excess imported unsaturated fatty acids (32, 44). Together, the results implied that TAG-
synthesizing enzymatic activity can be physiologically assessed by analyzing the growth of recombinant H1246 in the presence of ALA.

LuPDAT1 and LuPDAT2 can also preferably utilize substrates containing certain PUFAs—The previous results showed the unique selectivity properties of LuPDAT1 and LuPDAT2 for substrates containing ALA. To further examine the substrate preferences of flax PDATs, we conducted a similar experiment using other PUFAs that are not commonly found in plants, including SDA, DGLA, GLA, AA, ETA, EPA and DHA. LuPDAT1 and LuPDAT2 appeared to accept substrates containing SDA, GLA and EPA (Figure 9), while DGLA, AA, ETA and DHA were markedly poorer substrates for LuPDAT1 and LuPDAT2. Together, the data presented in Figures 5 and 9 indicated that LuDGAT1 are capable of utilizing a broad range of substrates containing different fatty acids. However, LuPDAT1 and LuPDAT2 have clear preference for certain PUFAs, particularly those with at least three double bonds, such as ALA, GLA, SDA and EPA.

Feeding fatty acids to yeast provides an easy and flexible way to introduce novel acyl moieties in yeast endogenous lipids. This method is, however, limited as it does not allow a controlled distribution of exogenous fatty acids to specific glycerolipids as it occurs naturally. To circumvent this problem, LuPDAT1, LuPDAT2 and LuDGAT1-1 were individually co-expressed with LuFAD2-1 and LuFAD3B, thus simulating a natural production of ALA in PC. In this experiment yeast cells expressing LuPDAT1 and LuPDAT2 produced TAG predominantly with ALA while cells expressing LuDGAT1-1 produced TAG mainly with palmitoleic acid (16:1 \( ^\text{cis} \Delta^9 \)) (Figure 10). LuPDAT1 and LuPDAT2 were able to produce TAG with an ALA:LA ratio of 10:1, while LuDGAT1-1 produced TAG with an ALA:LA ratio of approximately 2:1. These results further confirmed that LuPDAT1 and LuPDAT2 have similar preference for ALA-containing substrate.

Seed-specific expression of LuDGAT1-1, LuPDAT1 and LuPDAT2 affects the fatty acid profile and oil content in Arabidopsis—Complementation assays in yeast indicated that LuDGAT1-1, LuDGAT2-3, LuPDAT1 and LuPDAT2 are active TAG-synthesizing enzymes from flax. Though informative, this expression system is limited as yeast lacks the cellular and developmental complexity of multicellular higher plants. To investigate the functionality of flax TAG-synthesizing enzymes in plants, we expressed the ORFs of LuDGAT1-1, LuDGAT2-3 and LuDCR1 in Arabidopsis dgatl mutant (AS11 strain) under the regulation of the seed-specific napin promoter. AS11 has a reduced amount of TAG and altered fatty acid profile (low OA and eicosenoic acid (EA, 20:1 \( ^\text{cis} \Delta^{11} \)) and high ALA) (18). To minimize variation of genome position and copy number-dependent transgene expression (45, 46), ten individual transgenic lines were analyzed for each construct. This strategy also helped to compensate for environmental effects that can significantly influence the accumulation of oil in the seeds of each plant even though they were cultivated side-by-side. Analysis of T2 transgenic seed lines indicated that overexpression of LuDGAT1-1 restored seed oil content of AS11 to levels comparable to the empty vector-transformed control wild-type plants (Figure 11A). LuDCR1 and LuDGAT2 did not complement the reduced TAG phenotype of the AS11 mutant (Figure 11A). GC analysis of the total lipid extract fatty acid methyl esters (TLE-FAMEs) showed that LuDGAT1-1 could also alter the seed fatty acid profile of AS11 making it similar to wild-type controls. None of the LuDGAT2-3 and LuDCR1 transgenic lines could, however, complement this phenotype (Figure 11B).

It has been shown that the Arabidopsis PDAT1 knockout mutant did not have any significant changes in either fatty acid composition or oil content compared with wild-type (47) while the attempt to obtain the Arabidopsis dgatl pdat1 double mutant failed due to the lethal effect of a dgatl pdat1 genotype on pollen development (17). To study the biochemical properties of LuPDATs in plant oil biosynthesis, the constructs carrying the coding region of LuPDATs (LuPDAT1, LuPDAT2 or LuPDAT6) were transformed into Arabidopsis wild-type Columbia. As compared with the empty vector-transformed control wild-type plants, overexpression of the LuPDAT1 and LuPDAT2 increased the PUFA (LA and ALA) content, at the expense of mostly OA and EA.
α-linolenic acid-specific PDATs from flax

(Figure 12A). This altered fatty acid composition was not found in Arabidopsis lines overexpressing LuPDAT6. As shown in Figure 12A, the LuPDAT1 lines exhibited a relative increase of LA in the range of 4.9% to 14.3% compared with the average exhibited by empty vector transformed wild-type controls. In addition, the ALA content was significantly increased in many lines, with the highest relative increase of 8.1%. The LuPDAT2 transgenic T2 seed lines showed a relative increase of LA by 4.4% to 21.8%. ALA content was generally increased, with the highest relative increase value observed being 10.5%. The oil content of LuPDAT1, LuPDAT2 and LuPDAT6 Arabidopsis transgenic lines were not significantly altered compared with controls (Figure 12B).

DISCUSSION

Flax seed is a prominent oilseed crop and one of the most important plant-based sources of vegetable oils rich in ALA. Understanding the metabolic pathway of oil synthesis is pivotal for a continuous improvement of flax as an oilseed crop. The goal of this study was to identify flax genes involved in TAG synthesis and functionally characterize the encoded enzymes.

In mainstream oilseed crops such as Brassica napus and Glycine max, DGAT1 has been proposed to be the dominant enzyme involved in the final step of TAG biosynthesis (48, 49). But in plant species that accumulate exotic fatty acids such as ricinoleic, vernolic and α-eleostearic acid, DGAT2 and PDAT appear to have a more predominant role in TAG synthesis although DGAT1 is also expressed during seed development in these species (20, 49, 50). In the case of flax, we found gene homologues encoding all major TAG-synthesizing enzymes with the exception of DGAT3.

We have previously cloned a DGAT1 from flax (here refereed to LuDGAT1-1) and have demonstrated that it can complement TAG synthesis in the yeast strain H1246 (32). Although this enzyme was considerably active when produced recombinantly in yeast and plants (Figures 5A, 5D, 10 and 11), it did not appear to show a preference for substrates containing ALA (Figures 5D and 10). Like in other oilseeds, the expression of LuDGAT1 was considerably up-regulated in seeds (Figure 2A), but a comprehensive analysis throughout seed development indicated that the transcript levels peaked at 20 DPA when the rate of oil accumulation reached a plateau and most ALA is already incorporated in TAG (Figure 4B to 4D). Together, these results suggested that flax DGAT1 appears to play a role in the non-specific transfer of acyl moieties to TAG during the final stages of seed development. During the time this manuscript was being prepared, we detected an additional gene encoding DGAT1 (here referred to as LuDGAT1-2) in the final released version of flax genome sequence (27). LuDGAT1-1 and LuDGAT1-2 share 97.7% identity and due to this high degree of homology, their expression was detected together (Figures 2A and 4D).

In contrast to DGAT1, flax DGAT2-3 failed to complement TAG synthesis in Arabidopsis AS11 mutant (Figure 11). Production of LuDGAT2-3 recombinant in yeast H1246 resulted in the production of small amounts neutral lipids in yeast (Figure 5C) and ineffectively rescued the fatty acid lipotoxicity phenotype (Figure 8). The transcript of a third DGAT2 homologous gene (LuDGAT2-2) could not be detected or isolated. These results suggested that in flax DGAT2 might contribute minimally to TAG synthesis despite of being expressed specifically in seeds. Similarly, LuDCR1 did not show a significant effect on TAG biosynthesis in both yeast and Arabidopsis expression system (Figure 5 and 11).

In the case of flax PDATs, we analyzed one representative gene from each of the three groups of homologous genes (LuPDAT1, LuPDAT2, and LuPDAT6). Heterologous expression in yeast indicated that both LuPDAT1 and LuPDAT2 were active and able to produce TAG composed predominantly of ALA when ALA was supplemented as a free fatty acid to the medium or endogenously produced through the action of desaturases LuFAD2-1 and LuFAD3B (Figures 5C and 10). Interestingly, LuPDAT2 has higher level of protein accumulation than LuPDAT1 in yeast (Figure 7A), which may account partially for its higher TAG-forming ability. The reason for poor expression levels of LuPDAT1 is currently unknown but may be due to the low mRNA stability or inefficient codon usage. Overexpression of LuPDAT1 or LuPDAT2 in Arabidopsis also resulted in an enhanced content of PUFAs (Figure 12A). Analysis of gene
expression revealed that LuPDAT1/LuPDAT5 transcripts accumulated substantially more in seeds than in other vegetative tissues (Figure 3A). In fact, expression of LuPDAT1/LuPDAT5 was closely correlated with oil and ALA accumulation in developing embryos, presenting a peak during the rapid phase of lipid accumulation at 14 DPA (Figure 4B, 4C and 4G). LuPDAT2 was found to be mostly expressed in vegetative tissues (Figure 3D), indicating that TAG synthesized by the catalytic action of LuPDAT2 might have physiological functions in vegetative tissues of flax. LuPDAT6 was also highly expressed during the initial stages of embryo development (Figure 4I) but the encoded enzyme had significantly lower activity in all conditions tested (Figures 5, 8, 9 and 12). Based on sequence homology (Table 2) and similarities in expression profile (Figure 3), we suggest that the pairs LuPDAT1/LuPDAT5, LuPDAT2/LuPDAT4, and LuPDAT3/LuPDAT6 may have similar functions in flax. The fatty acid feeding experiment confirmed that LuPDAT5, similar to LuPDAT1, also has the higher preference for ALA-containing substrate (Supplemental Figure 5). On the basis of these results, we propose that LuPDAT1/LuPDAT5 may play a more important role than LuDGAT1 in the production of ALA-containing TAG in developing flax seed. This hypothesis is in agreement with metabolic analyses of the acyl-CoA profile of developing flax seed which detected only small amounts of ALA-CoA in flax embryo (51).

The identified ALA-selective flax PDATs in this study could be considered as another example of a plant TAG-synthesizing enzyme that displays preference for certain substrates. Our data indicates, however, that these PDATs not only favour substrates containing ALA, but also extend to other PUFAs that are not naturally formed in flax, including SDA, GLA and EPA (Figure 9). It appears that the activity of these PDATs is highly dependent on ALA availability. Increasing the concentration of exogenously provided ALA resulted in an enhanced amount of ALA-containing TAG and total TAG contents in yeast expressing LuPDAT1 or LuPDAT2 (Figure 7B to 7E). When ALA was added to the medium, both LuPDAT1 and LuPDAT2 were able to produce TAG with up to 90% ALA (Figure 7C). In contrast, when ALA was endogenously produced by the co-expression of LuFAD2-1 and LuFAD3B, the total amount of ALA in TAG was only about 50% for LuPDAT1 or LuPDAT2 (Figure 10). This difference can be explained by two possible reasons. One possible explanation is that the ALA availability may limit the activity of these two PDATs. Inefficient desaturase activity in the heterologous yeast expression system has been recently reported by Dahmen et al. (2013) (52). They found that yeast cytochrome b5, which transfers the electrons to the desaturase, poorly interacts with foreign desaturases, thus limiting desaturase enzymatic activity and resulting in the low production of unsaturated fatty acids. Future studies of the over-expression of flax cytochrome b5 in the LuFAD2-1-LuFAD3b-LuPDAT-expressing yeast poses an opportunity to further enhance the production of ALA-containing TAG in yeast by metabolic engineering. The other possible reason is that to produce trilinolenin the flax PDATs require sn-2 ALA-PC and sn-1,2 ALA-DAG. Considering that the excess of ALA-CoA derived from the exogenous supplied ALA can be transferred indiscriminately into different glycerolipids, there is a better chance of obtaining sn-1,2 ALA-DAG. In contrast, ALA produced by a FAD3 is restricted to sn-2 PC, which can be utilized by PDAT thus limiting its redistribution to other lipid classes such as DAG. Therefore, the limited amount of sn-1,2 ALA-DAG may restrict the activity of LuPDAT1 and LuPDAT2. The same theory can be used to explain the small but significant increase of ALA content in the seed oil of LuPDAT1- and LuPDAT2-overexpressing Arabidopsis lines. It is possible that the activity of FAD3 in Arabidopsis might not be sufficiently high to supply the substrates required for LuPDAT1 and LuPDAT2. For instance, expression of FAD3 in wild-type Arabidopsis is significantly lower compared to the AS11 mutant (53). Furthermore, labeling experiments indicated that Arabidopsis utilizes the PC-derived DAG pool as the major source for TAG biosynthesis (54). PC-derived DAG synthesis, however, may provide mostly sn-2-ALA-DAG. It is possible that when produced in Arabidopsis, LuPDAT1 and LuPDAT2 may rapidly transfer ALA from the sn-2 position of PC to the sn-3 position of the PC-derived DAG. This rapid conversion could limit the chance of rechanneling ALA from PC to acyl-CoA pool and subsequently incorporating it to the
α-linolenic acid-specific PDATs from flax

sn-1 position of DAG by the endogenous Arabidopsis enzymes, thus imposing a bottleneck for accumulating ALA in Arabidopsis seed oil.

A phylogenetic analysis of PDATs including the functionally tested enzymes from Arabidopsis (17) and castor bean (24, 50) indicate that LuPDAT1 and RcPDAT1-2 are closely related (Supplementary Figure 2A). Interestingly, both polypeptides display remarkable substrate selectivity and the corresponding genes are preferentially expressed in seeds. The branch containing LuPDAT1 and RcPDAT1-2 lacks a corresponding polypeptide from Arabidopsis. LuPDAT2 is, however, more closely associated with AtPDAT1 and RcPDAT1-1. It is interesting that in Arabidopsis upon AtDGAT1 disruption, AtPDAT1 is responsible for synthesizing the remaining 65-70% of TAG (53) despite of the inability of AtPDAT1 or AtPDAT2 to complement TAG synthesis in H1246 (17). In addition, overexpression of AtPDAT1 in wild-type Arabidopsis background did not produce a significant effect on oil phenotype (23). Similarly, expression of RcPDAT1-1 in Arabidopsis CL37 lines did not result in any substantial increase in hydroxy fatty acids (50). Overexpression of LuPDAT2 in wild-type Arabidopsis, however, led to a significantly increased PUFA level. This difference might be explained by the higher substrate selectivity of LuPDAT2. LuPDAT6 belongs to the same branch of AtPDAT2 and RcPDAT2. Although PDATs in this branch are preferentially expressed in developing seeds, they do not have a significant effect on TAG accumulation. Overall, these findings suggest the existence of a class of plant PDATs that display substrate selectivity properties.

One of the most remarkable advances in oilseed biotechnology is the successful reconstitution of VLC-PUFA synthesis in transgenic plants (reviewed by Ruiz-López et al. (55). Flax would become an obvious candidate for this technology since it naturally contains substantial amounts of ALA that is the precursor for VLC-PUFAs. A previous report demonstrated that the introduction of a VLC-PUFA biosynthetic pathway in flax led to a high proportion of Δ6-desaturated C-18 fatty acids (GLA and SDA totaling up to 33% in some plants). The pathway, however, did not efficiently proceed further after the first desaturation step, resulting in very limited amount of AA and EPA (<5%) in flax oil (56). As demonstrated before, the front-end desaturases and elongases involved in VLC-PUFA have different acyl carrier specificity (57). Desaturases act on PC, whereas elongases occur mainly within the acyl-CoA pool. According to the results presented in Figure 9, it is likely that LuPDAT1 contributes to a premature channeling of GLA and SDA to TAG, resulting in an insufficient supply of acyl-CoA substrates for elongation. The observation that GLA and SDA were nearly absent from the acyl-CoA pool of the transgenic flax lines (56) further supports this hypothesis. To circumvent this problem, the metabolic pathway for VLC-PUFAs would have to be strategically modified to bypass other acyl moieties that could be utilized by LuPDAT1 until it reaches EPA. For example, the pathway involving the utilization of Δ8-desaturation pathway to convert LA to AA (58) and then further desaturating AA to EPA by Δ17 desaturases (59), would contain intermediates (LA, DGLA and AA) which are poorly utilized by LuPDAT1 (Figures 5E and 9).

In conclusion, this study provided new insights into the biosynthesis of TAG in flax. Our results demonstrated the existence of two pairs of flax PDATs that are highly selective for substrates containing ALA. Among them, one pair of PDATs (LuPDAT1/LuPDAT5) is expressed predominately in seeds. In addition, the oil and ALA accumulation during seed development closely correspond to the expression pattern of this seed-preferred PDAT gene pair, suggesting the critical role of these PDATs in seed oil biosynthesis in flax. The identified PDATs help in the understanding of the mechanisms involved in producing TAG with high PUFA content in plants. Our characterization of TAG-synthesizing enzymes in flax will benefit future projects aimed at enriching PUFA in plants and other organisms.
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Acknowledgments—RJW gratefully acknowledges the support provided by Genome Alberta/Genome Canada, the Canada Research Chairs Program, the Canada Foundation for Innovation, the Research Capacity Program of Alberta Enterprise and Advanced Education, AVAC Ltd., Alberta Innovates Bio Solutions and the Natural Sciences and Engineering Research Council of Canada. XP is the recipient of the Alberta Innovates Graduate Student Scholarship. The authors thank Dr. Gordon Rowland for the providing flax seed and Dr. Jitao Zou for the use of laboratory facilities at the National Research Council of Canada in Saskatoon.

FOOTNOTES

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2 The abbreviation used are: DGAT, acyl CoA:diacylglycerol acyltransferase; PDATs, phospholipid:diacylglycerol acyltransferase; TAG, triacylglycerol; ALA, α-linolenic acid; VLC-ω-3-PUFA, very long chain omega-3 polyunsaturated fatty acids; FAD, fatty acid desaturases; EMS, ethyl methanesulfonate; PC, phosphatidylcholine; LPCAT, acyl-CoA: lysophosphatidylcholine acyltransferase; PLA2, phospholipase A2; DAG, diacylglycerol; PDCT, phosphatidylcholine:diacylglycerol cholinephosphotransferase; ER, endoplasmic reticulum; DCR, defective in cuticular ridge; DPA, days post anthesis; MCS, multiple cloning site; 3’-UTR, 3’-untranslated region; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; UBI2, Ubiquitin extension protein; OA, oleic acid; LA, linoleic acid; SDA, stearidonic acid; DGLA, dihomo-γ-linolenic acid; GLA, γ-linolenic acid; AA, arachidonic acid; ETA, eicosatrienoic acid; EPA, eicosapentaenoic acid; DHA, docosahexaenoic acid; NaCl, sodium chloride; TLC, thin layer chromatography; GC, gas chromatography; MS, mass spectrometry; FAME, fatty acid methyl esters; ORFs, open reading frames; ASAT, acyl-CoA:sterol acyltransferase; EA, eicosenoic acid.

3Author Contributions: X.P. performed gene structure analysis, molecular cloning, Western blot analysis, recombinant protein expression in yeast and *Arabidopsis* system and data analysis. R.W., R.S. and X.P. designed the research. A.W. performed gene expression and molecular cloning for *LuFAD2-1* and *LuFAD3B* genes and analyzed the flax oil and ALA accumulation during seed development. E.M. provided the protocols for *Arabidopsis* transformation and seed oil analysis and contributed valuable advice and discussions during this study. X.P. and R.S. contributed for article writing.
FIGURE LEGENDS

FIGURE 1. Genomic DNA structure of homologous genes encoding putative triacylglycerol (TAG)-
synthesizing enzymes from flax. The thick lines indicate exons and the thin lines indicate introns. The
numbers on top of each group denote the scale in base pairs.

FIGURE 2. Quantitative real-time PCR analysis of relative expression of flax DGAT genes in different
tissues. The transcript levels of LuDGAT1 (A) and LuDGAT2 (B and C) genes were higher in seeds (40
DPA, black bar) than that in other tissues (gray bar) and the opposite was observed for LuDCR1 (D).
Data are given as means ± SE (n=3). S1: immature stem, S2: developing stem, S3: mature stem, L: leaves,
A: apexes, R: roots, F: flowers, 40D: mature seeds.

FIGURE 3. Quantitative real-time PCR analysis of relative expression of flax PDAT genes in different
tissues. The LuPDAT1/LuPDAT5 (A), LuPDAT3 (B) and LuPDAT6 (C) genes exhibited higher
expression level in seeds (40 DPA, black bar) than that in other tissues (gray bar) and the opposite was
observed for LuPDAT2 (D) and LuPDAT4 (E).
(F) Phylogenetic analysis of six flax PDAT proteins. Genes within the same branch of the phylogenetic
tree have the similar expression pattern.
Data are given as means ± SE (n=3). S1: immature stem, S2: developing stem, S3: mature stem, L: leaves,
A: apexes, R: roots, F: flowers, 40D: mature seeds.

FIGURE 4. Oil and α-linolenic acid (ALA) accumulation correlate better with the expression of
LuPDATs than with LuDGATs.
(A) Flax embryos or seeds used for quantitative real-time PCR analysis. The top panel shows the cross-
sections of the developing bolls at 4 to 25 DPA and the whole mature boll at 40 DPA from left to right.
The bottom panel shows the cleared seed at 4 DPA, developing embryos at 8 to 25 DPA and the mature
seed at 40 DPA, DPA, days post anthesis.
(B) Oil deposition in seed on a fresh weight basis (% of fresh weight). The mean values of oil content
(n=4) were fitted to a sigmoidal curve (R²=0.969) using Sigmaplot 12.3.
(C) ALA accumulation pattern during seed development. The data is presented on a fresh weight basis (%
of fresh weight) and shown as mean ± SE (n=4).
(D to I) Analysis of gene expression patterns of LuDGATs and LuPDATs during seed development. Data
are shown as mean ± SE (n=3).

FIGURE 5. Functional complementation assay in yeast strain H1246. Yeast cells expressing several flax
genes show enhanced synthesis of triacylglycerol (TAG) in the presence of α-linolenic acid (ALA).
(A) Nile red fluorescence assay was performed on the recombinant yeast mutant H1246 cultured in
medium with ALA (gray bar) or without ALA supplementation (black bar). The value represents the
levels of neutral lipids accumulated in the recombinant yeast, which is calculated by dividing Nile red
fluorescence (ΔF) by the optical density (OD) at 600nm. Data are presented as means ± SE (n = 3). Total
lipid extracts were prepared from these cultures and the individual lipid classes were separated by thin
layer chromatography (TLC) (B and C).
(B) TLC analysis of yeast lipid in the absence of ALA.
(C) TLC analysis of yeast lipid in the presence of ALA. The square bracket indicates the position of TAG
produced by yeast cells. The insert displays a high contrast image of TAG produced in yeast expressing
LuDGAT2-3 in the present of ALA.
(D) TLC analysis of yeast cells expressing LuDGAT1-1 in the absence or presence of different
exogenously provided fatty acids.
(E) and (F) TLC analysis of yeast lipid in the presence of linoleic acid (LA) and oleic acid (OA).
Yeast cells transformed with pYESLacZ was used as the negative control. Although yeast cells transformed with pYESLacZ and pYESLuPDAT6 also showed the higher Nile red value in the presence of ALA (A), TLC plate indicated that the increased value was very likely due to the production of the unknown compound (square bracket indicated) (C). The corresponding fatty acid used for feeding is shown on the left of the figure with the chemical structure. D1– LuDGAT1-1; DCR1 – LuDCR1; D2-3 – LuDGAT2-3; P1– LuPDAT1; P2 – LuPDAT2; P6 – LuPDAT6 ; TAG – triolein TAG standard in (B, E and F), trilinolenin TAG standard in (C and D).

FIGURE 6. Gas chromatography-mass spectrometry chromatograms of yeast strain H1246 expressing LuPDAT1 and LuPDAT2 in the presence of α-linolenic acid (ALA). The recombinant yeast cells were cultivated in the presence of ALA. The yeast lipids were extracted and separated by thin layer chromatography (TLC) plate. The compound corresponding to the upper and lower triacylglycerol (TAG) bands (marked by square brackets) was scrapped separately from the TLC plate, transmethylated and analyzed through GC/MS. The ratio of TAG within each separated band to the total amount of TAG was calculated and the values are shown as percentage on the upper left corner of each chromatograph. GC profiles showed that yeast strain H1246 expressing LuPDAT1 or LuPDAT2 has the ability to produce TAG containing only ALA (trilinolenin), when ALA is added exogenously to the cell. P1– LuPDAT1; P2 – LuPDAT2.

FIGURE 7. Yeast cell expressing LuPDAT1 and LuPDAT2 have similar responses to an increase of exogenously provided α-linolenic acid (ALA).

(A) Representative Western blot analysis showing HisG-LuPDAT protein levels. Microsomes were prepared from yeast expressing HisG-PDATs (See Experimental Procedures). Cells transformed with empty pYES2/NT-C vector were used as a negative control, which was annotated as “-” in the image. Equal protein loading was ensured by measuring the constitutively expressed Kar2p in yeast.

(B) Thin layer chromatography (TLC) analysis showing triacylglycerol (TAG)-forming ability of recombinant yeast in the presence of different concentration of ALA. The yeast cells expressing LuPDAT1 and LuPDAT2 were cultivated in the absence or presence of different ALA concentration and harvested at the same growth stage (OD\textsubscript{600nm} = 6.5±0.05). The yeast lipids were extracted and separated by TLC and the corresponding TAG spots are shown in (B).

(C-E) The ALA concentration effect on overall percentage of ALA in TAG (C), total TAG content (D) and amount of fatty acids in TAG (E) of recombinant yeasts. Total TAG content is presented on the basis of yeast dry weight. The amount of ALA and other fatty acids is presented as µmol of fatty acids per mg of yeast dry weight. Data are presented as means ± SE (n = 4). P1 – LuPDAT1; P2 – LuPDAT2; FA – fatty acid.

FIGURE 8. Rescue of lipotoxicity phenotype in H1246.

Exogenous supplied α-linolenic acid (ALA) inhibits the growth of yeast H1246. This inhibition can be rescued by expressing genes encoding triacylglycerol (TAG)-synthesizing enzymes from flax and the rescuing effect is correlated with the ability to synthesize TAG as demonstrated with Nile red assay and TLC analysis. Growth curves of recombinant yeast H1246 cultured in induction medium with 0 (A), 100 µM (B) and 1mM (C) of ALA, were constructed by measuring the optical density (OD) of the cultures at 600 nm. Data are presented as means ± SE (n = 3 or 6). BY4742 (wild-type) and H1246 harboring the empty vector pYES were used as positive and negative controls, respectively. For negative controls, six independent colonies were analyzed and consistently showed delayed growth and low final density even in the absence of free fatty acid. The panels on the right side, from top to bottom, show the growth of 2.5 µl of each tenfold dilution (1 to10\textsuperscript{5}) of recombinant yeast H1246 spotted on to induction medium plates containing 0, 100 µM and 1 mM ALA. The negative controls were annotated as “-” in the image. WT- wild-type; D1-1 – LuDGAT1-1; DCR1 – LuDCR1; D2-3 – LuDGAT2-3; P1– LuPDAT1; P2 – LuPDAT2; P6 – LuPDAT6.
**FIGURE 9.** Polyunsaturated fatty acid (PUFA)-specific activity of LuPDAT1 and LuPDAT2 in yeast strain H1246. Yeast mutant H1246 transformed with LuPDAT genes was cultivated in the presence of different PUFAs fed individually to a final concentration of 100 μM. Yeast transformed with pYESLacZ was used as the negative control, which was annotated as “-” on the thin layer chromatography (TLC) plate. The corresponding fatty acid used for feeding is shown on the left of the figure with the chemical structure. P1 – LuPDAT1; P2 – LuPDAT2; P6 – LuPDAT6; TAG – trilinolenin TAG standards; SDA – stearidonic acid; GLA – γ-linolenic acid; DGLA – dihomo-γ-linolenic acid; AA – arachidonic acid; EPA – eicosapentaenoic acid; ETA – eicosatrienoic acid; DHA – docosahexaenoic acid.

**FIGURE 10.** Comparison of fatty acid methyl esters (FAME) profile of yeast strain H1246 co-expressing LuPDAT1, LuPDAT2 and LuDGAT1-1 individually with desaturases. When compared with the yeast cell co-expressing LuDGAT1-1 with LuFAD2-1 and LuFAD3B (black bar), co-expression of either LuPDAT1 (gray bar) or LuPDAT2 (white bar) with LuFAD2-1 and LuFAD3B in yeast produces triacylglycerol (TAG) with α-linolenic acid (ALA, C 18:3) as the predominant fatty acid. Yeast cultures were induced at 20°C for three days before harvested. Data are presented as means ± SE (n = 3).

**FIGURE 11.** Lipid phenotype of LuDGAT-overexpressing seeds. Overexpressing LuDGAT1-1 in Arabidopsis AS11 background restores the wild-type lipid phenotype. Ten individual transgenic lines were analyzed for each construct. Arabidopsis AS11 mutant and wild-type transformed with empty pGreen plasmid were used as controls (CTR). Data are presented as means ± SE (n = 3). OA – oleic acid; ALA – α-linolenic acid; EA – eicosenoic acid.

**(A)** Seed oil content. Oil content is expressed as percentage of seed dry weight.

**(B)** Fatty acid composition. Values were obtained from fatty acid methyl esters (FAME) analysis of dry seeds.

**FIGURE 12.** Lipid phenotype of LuPDAT-overexpressing seeds. Ten individual transgenic lines were analyzed for each construct. Wild-type Arabidopsis transformed with empty pGreen plasmid were used as controls (CTR).

**(A)** Fatty acid composition of LuPDAT-overexpressing seeds. Overexpression of LuPDAT1 and LuPDAT2 in Arabidopsis seeds results in an increased level of linoleic acid (LA) and α-linolenic acid (ALA). Data are presented as means ± SE (n = 3), with asterisks indicating p<0.05 (ANOVA, Duncan’s multiple range test). OA – oleic acid; EA – eicosenoic acid.

**(B)** Oil content of LuPDAT-overexpressing seeds. Data are presented as means ± SE (n = 3).
Table 1. General information on individual candidate genes and the encoded polypeptides (deduced from the DNA sequence information)

| Genes   | Gene length (bp) | Protein length | Molecular mass (Daltons) | Isoelectric point |
|---------|------------------|----------------|--------------------------|-------------------|
| DGAT1-1 | 1524             | 507            | 58038.19                 | 8.98              |
| DGAT1-2 | 1542             | 513            | 58793.08                 | 8.92              |
| DGAT2-1 | 1065             | 354            | 39405.12                 | 9.49              |
| DGAT2-2 | 1059             | 352            | 39004.68                 | 9.19              |
| DGAT2-3 | 1050             | 349            | 38951.64                 | 9.42              |
| DCR1    | 1137             | 378            | 41128.85                 | 9.49              |
| DCR2    | 1401             | 466            | 50887.96                 | 5.10              |
| PDAT1   | 2088             | 695            | 76891.89                 | 8.23              |
| PDAT2   | 2145             | 714            | 79072.61                 | 6.4               |
| PDAT3   | 1728             | 575            | 63093.05                 | 6.19              |
| PDAT4   | 2148             | 715            | 78923.51                 | 6.72              |
| PDAT5   | 2088             | 695            | 76817.64                 | 8.28              |
| PDAT6   | 1719             | 572            | 62792.54                 | 6.19              |

Table 2. Sequence identity between the gene pairs and gene products

| Gene Pair                | Nucleotide (%) | Amino acid (%) |
|--------------------------|----------------|---------------|
| DGAT1-1 and DGAT1-2      | 85.9           | 97.7          |
| DGAT2-1 and DGAT2-3      | 91.2           | 93.3          |
| DCR1 and DCR2            | 90.2           | 89.3          |
| PDAT1 and PDAT5          | 97             | 97.1          |
| PDAT2 and PDAT4          | 95.6           | 97.1          |
| PDAT3 and PDAT6          | 95.4           | 96            |
α-linolenic acid-specific PDATs from flax

Figure 1

Figure 2
Figure 3
α-linolenic acid-specific PDATs from flax

Figure 4

A

Boll

Embryo
or seed

DPA

4 8 11 14 16 20 25 40

B

Oil% (w/w)

DPA

0 5 10 15 20 25 30

0 10 20 30 40

C

ALA% (w/w)

DPA

25 29 31 33 35 37 39 41 43

0 1 2 3 4 5

D

DGAT1-1/DGAT1-2

Relative expression

DPA

0 0.2 0.4 0.6 0.8 1 1.2 1.4 1.6 1.8

4 8 11 14 16 20 25 30 35 40

E

DGAT2-1

Relative expression

DPA

0 0.2 0.4 0.6 0.8 1 1.2 1.4

4 8 11 14 16 20 25 30 35 40

F

DGAT2-3

Relative expression

DPA

0 0.5 1.0 1.5 2.0 2.5 3.0 3.5 4.0 4.5

4 8 11 14 16 20 25 30 35 40

G

PDAT1/PDAT5

Relative expression

DPA

0 0.5 1 1.5 2 2.5 3 3.5 4

4 8 11 14 16 20 25 30 35 40

H

PDAT3

Relative expression

DPA

0 0.02 0.04 0.06 0.08 0.1 0.12

4 8 11 14 16 20 25 30 35 40

I

PDAT6

Relative expression

DPA

0 0.02 0.04 0.06 0.08 0.1 0.12

4 8 11 14 16 20 25 30 35 40
Figure 5

A

![Graph showing ΔF/OD with conditions: No ALA and ALA Feeding](image)

B

| DCR1 | D2-3 | P1 | P2 | P6 | LacZ | TAG |
|------|------|----|----|----|------|-----|

C

| ALA | DCR1 | D2-3 | P1 | P2 | P6 | LacZ | TAG |
|-----|------|------|----|----|----|------|-----|

D

| D1- NoFA | D1- OA | D1- LA | D1- ALA | TAG |
|----------|-------|------|--------|-----|

E

| LA | DCR1 | D2-3 | P1 | P2 | P6 | LacZ | TAG |
|----|------|------|----|----|----|------|-----|

F

| OA | DCR1 | D2-3 | P1 | P2 | P6 | LacZ | TAG |
|----|------|------|----|----|----|------|-----|
$\alpha$-linolenic acid-specific PDATs from flax
Figure 7

A

P1  P2  P6  

Kar1p

B

No FA  50  100  150  200  300

P1

P2

C

Exogenously provided ALA (μM)

D

Exogenously provided ALA (μM)

E

μmol FA/mg dry yeast

Exogenously provided ALA (μM)
Figure 8

A

OD

12

10

8

6

4

2

0

6 18 24 30 42 48 54 66 72 78

D1-1 D2-3 P2 WT DCR1 P1 P6

NOFA

1

1x10^{-1}

1x10^{-2}

1x10^{-3}

1x10^{-4}

B

OD

12

10

8

6

4

2

0

6 18 24 30 42 48 54 66 72 78

100\mu M

1

1x10^{-1}

1x10^{-2}

1x10^{-3}

1x10^{-4}

C

OD

10

9

8

7

6

5

4

3

2

1

0

6 18 24 30 42 48 54 66 72 78

1mM

1

1x10^{-1}

1x10^{-2}

1x10^{-3}

1x10^{-4}
Figure 9

α-linolenic acid-specific PDATs from flax
$\alpha$-linolenic acid-specific PDATs from flax

**Figure 10**

![Graph showing fatty acid composition](image)

**Figure 11**

(A) CTR (AS11)  CTR (WT)  DGAT1  DGAT2-3  DCR1

(B) QA  ALA  EA

![Bar chart showing oil content and fatty acid profile](image)
$\alpha$-linolenic acid-specific PDATs from flax

Figure 12

A

LA (Mol\%)  
CTR  PDAT1  PDAT2  PDAT6

36  34  32  30  28  26  24  22  20  18

ALA (Mol\%)  
CTR  PDAT1  PDAT2  PDAT6

17  16  15  14  13  12  11  10  9  8  7  6  5  4  3  2  1

OA (Mol\%)  
CTR  PDAT1  PDAT2  PDAT6

20  19  18  17  16  15  14  13  12  11  10  9  8  7  6  5  4  3  2  1

EA (Mol\%)  
CTR  PDAT1  PDAT2  PDAT6

22  20  18  16  14  12  10  8  6  4  2

B

Oil content (%)  
CTR  PDAT1  PDAT2  PDAT6

0  5  10  15  20  25  30  35

Transgenic line

31