Phospholipid:Diacylglycerol Acyltransferase Is a Multifunctional Enzyme Involved in Membrane Lipid Turnover and Degradation While Synthesizing Triacylglycerol in the Unicellular Green Microalga Chlamydomonas reinhardtii

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Many unicellular microalgae produce large amounts (~20 to 50% of cell dry weight) of triacylglycerols (TAGs) under stress (e.g., nutrient starvation and high light), but the synthesis and physiological role of TAG are poorly understood. We present detailed genetic, biochemical, functional, and physiological analyses of phospholipid:diacylglycerol acyltransferase (PDAT) in the green microalga Chlamydomonas reinhardtii, which catalyzes TAG synthesis via two pathways: transacylation of diacylglycerol (DAG) with acyl groups from phospholipids and galactolipids and DAG:DAG transacylation. We demonstrate that PDAT also possesses acyl hydrolase activities using TAG, phospholipids, galactolipids, and cholesteryl esters as substrates. Artificial microRNA silencing of PDAT in C. reinhardtii alters the membrane lipid composition, reducing the maximum specific growth rate. The data suggest that PDAT-mediated membrane lipid turnover and TAG synthesis is essential for vigorous growth under favorable culture conditions and for membrane lipid degradation with concomitant production of TAG for survival under stress. The strong lipase activity of PDAT with broad substrate specificity suggests that this enzyme could be a potential biocatalyst for industrial lipid hydrolysis and conversion, particularly for biofuel production.

INTRODUCTION

Glycerolipids represent the largest class of lipids, comprising 10 to 90% of eukaryotic cell dry weight (Bell and Coleman, 1980). These compounds play a variety of biological roles, including being major components of membrane matrices and functional complexes (such as photosynthetic reaction centers and electron transport chains), highly concentrated carbon and energy reserves, and signal molecules. Triacylglycerols (TAGs) are the most important carbon and energy storage lipids in eukaryotes and are also a major source of highly reduced carbon molecules for food and fuel.

In eukaryotes, TAGs are known to be synthesized via multiple pathways. The acyl-CoA-dependent Kennedy pathway involves three sequential acylations of glycerol-3-phosphate catalyzed by a suite of enzymes, the last of which is diacylglycerol acyltransferase (DGAT) that catalyzes the acylation of a diacylglycerol (DAG) with an acyl-CoA to form TAG. This pathway has been well studied in yeasts (Lardizabal et al., 2001; Sorger and Daum, 2002), plants (Katavic et al., 1995; Jako et al., 2001; Shockey et al., 2006), and mammals (Cases et al., 1998, 2001).

Another TAG synthesis pathway is the acyl-CoA-independent pathway, mediated by phospholipid:DAG acyltransferases (PDATs) that transfer a fatty acyl moiety from a phospholipid (PL) to DAG to form TAG. The PDAT pathway has been reported in yeasts (Dahlqvist et al., 2000; Zhang et al., 2012) and vascular plants (Dahlqvist et al., 2000; Ståhl et al., 2004). Finally, the transacylation pathway uses two DAG molecules as both acyl donors and acceptors to form TAG. This pathway is catalyzed by an acyl-CoA-independent diacylglycerol transacylase (DGTA). DGTA activities have been detected in microsome preparations of intestines of Rattus norvegicus (Lehner and Kuksis, 1993) and developing seeds of safflower (Carthamus tinctorius) (Lehner and Kuksis, 1993; Stobart et al., 1997), but no gene encoding DGTA has been identified in any organism.

Many unicellular microalgae produce large amounts of neutral lipids (20 to 50% of cell dry weight), mainly in the form of TAG, under unfavorable environmental or culture conditions, but the molecular and cellular mechanisms for TAG synthesis and accumulation are less well understood (Hu et al., 2008). The unicellular green microalga Chlamydomonas reinhardtii serves as a model organism for elucidating the pathways and physiological role of TAG synthesis. C. reinhardtii grows rapidly and produces TAG, particularly under adverse abiotic environmental conditions (Weers and Gulati, 1997; Li et al., 2010a, 2010b; Miller et al., 2010), and its entire genome sequence and genetic toolbox are available (Kindle, 1990; Merchant et al., 2007; Molnar et al., 2009). We observed that the accumulation of TAG in C. reinhardtii under stress is accompanied by substantial degradation of chloroplast membranes and thus hypothesized that fatty acids from the chloroplasts and other intracellular membrane systems may be converted into TAG by a PDAT-like...
protein. Because the predominant lipids in the chloroplast membranes are galactolipids, including monogalactosyldiacylglycerol (MGDG) and digalactosyldiacylglycerol (DGGD) (Janero and Barrnett, 1981), we further hypothesized that the PDAT-like protein might also function as a galactolipid:DAG acyltransferase. Recently, the single gene encoding PDAT in C. reinhardtii, referred to as Cr-PDAT in this study, was cloned, and a loss-of-function study indicated that the PDAT pathway contributed to ~25% of the total TAG accumulated in C. reinhardtii under nitrogen-depleted conditions (Boyle et al., 2012). However, the biochemical and functional properties of Cr-PDAT remain elusive.

In this study, we cloned a full-length cDNA encoding PDAT from C. reinhardtii. The in vitro and in vivo biochemical characteristics and physiological functions of Cr-PDAT were elucidated. Multiple lines of evidence indicate that PDAT is a multifunctional protein and plays an essential role in membrane turnover, degradation, and TAG formation in C. reinhardtii.

RESULTS

Identification and Cloning of a Putative PDAT cDNA from C. reinhardtii

To identify the gene encoding PDAT in C. reinhardtii, we conducted a BLAST search of the Chlamydomonas genome database and GenBank with Sc-PDAT and At-PDAT as query sequences. We obtained a sequence annotated as phosphatidylcholine-sterol O-acyltransferase (LCAT) (accession number XM_001699696). This putative LCAT gene encoded a protein with high similarity to Sc-PDAT (35%) and At-PDAT (34%); therefore, this gene was cloned and designated as Cr-PDAT. The full-length cDNA of Cr-PDAT contained a 3157-bp open reading frame flanked by 5’ and 3’ untranslated regions of 126 and 503 bp, respectively (see Supplemental Figure 1A online). The open reading frame consisted of 15 exons (see Supplemental Figure 1A online) and is predicted to encode a protein of 1041 amino acid residues with a mass of 104,600 D and a pI of 5.96.

Analysis of the amino acid sequence predicted from the Cr-PDAT cDNA indicated that this gene belongs to the membrane-bound O-acyl transferase (MBOAT) family. Like all other MBOAT proteins, a putative N-glycosylation site and a Ser/Thr phosphorylation site were detected in Cr-PDAT, and InterProScan indicated the presence of an LCAT domain between amino acids 267 and 539 of the deduced Cr-PDAT amino acid sequence (see Supplemental Figure 2 online).

Sequence alignment of a number of LCAT-like proteins, including PDAT, LCAT, and phospholipid:sterol acyltransferase, revealed seven characteristic conserved regions (see Supplemental Figure 1B online). A catalytic triad (Ser-Asp-His), which was previously identified in mammalian LCAT (Peelman et al., 1998) and plant and yeast PDAT (Ståhl et al., 2004; Banas et al., 2005), was conserved in all LCAT-like proteins, including Cr-PDAT. This triad is part of the catalytic domain of LCAT enzymes, by which a fatty acid is cleaved from the sn-2 position of phosphatidylcholine (PC) and then transesterified with a free hydroxyl group of cholesterol to generate a cholesterol ester (Peelman et al., 1998). A highly conserved hydrophobic domain (domain I in Supplemental Figure 1B online) was found in all LCAT-like proteins, but its function is unknown. LCAT-like proteins also contain a so-called lid domain (domain II; see Supplemental Figure 1B online), which may be involved in destabilizing the lipid bilayer, facilitating binding of a hydrophobic substrate and diffusing it into the enzyme’s active site cavity (Peelman et al., 1998). A Trp (Trp-285) residue present in this lid domain (see Supplemental Figure 1B online) is thought to bind the cleaved fatty acid in the active site of the At-PDAT (Martinelle et al., 1996; Ståhl et al., 2004). Domain III (corresponding to 366PYDWFRL365 in Cr-PDAT) is highly conserved in all sequences (see Supplemental Figure 1B online) and may be involved in PL recognition; several residues, including Glu-173 and the amphipathic helix encompassing residues 175 to 198 in the human LCAT (Hs-LCAT), which determines its substrate specificity and binding (Peelman et al., 1998), reside within this domain.

To gain insights into the evolutionary relationship between Cr-PDAT and other MBOAT proteins, we reconstructed a cladogram based on the multiple sequences from plants, animals, fungi, and algae. Based upon our phylogenetic analysis, the MBOAT family proteins were clustered into four major groups, corresponding to LCAT, PDAT, phospholipid:sterol acyltransferase, and phospholipase (PLA) proteins (Figure 1). The PDAT group branched into three subgroups: plant, fungal, and green algal PDAT. The algal PDAT lineage was more closely related to fungal PDAT than plant PDAT. The PLA group was the most divergent group related to PDAT but was closely related to the LCAT group.

TMHMM analysis (transmembrane prediction based on a hidden Markov model) and hydrophy profile indicated that Cr-PDAT is an integral membrane protein with a transmembrane domain of 23 amino acids at its N terminus (see Supplemental Figure 3 online), similar to the structures of At-PDAT and Sc-PDAT (Dahlqvist et al., 2000; Ståhl et al., 2004). PDAT activities were detected in microscope preparations from yeasts and vascular plants (Dahlqvist et al., 2000), suggesting a possible localization of this enzyme in the endoplasmic reticulum (ER); however, these microscope preparations contained other cellular compartments (e.g., Golgi apparatus and plasma membrane); thus, the presence of PDAT in those organelles cannot be ruled out. Three plant PDATs have an aromatic amino acid-rich stretch in the C-terminal end recently shown to act as an ER retrieval motif (McCARTNEY et al., 2004). However, no signal peptide or C-terminal retention sequence for ER targeting was detected in the Cr-PDAT sequence. Instead, a 92–amino acid canonical chloroplast transit peptide was identified at the N terminus of Cr-PDAT, suggesting a possible chloroplast localization.

Expression of the Cr-PDAT Gene in Pichia pastoris

To determine the function and enzymatic activity of Cr-PDAT, full-length PDAT (ORFCrPDAT) and truncated PDAT without the transmembrane domain (ΔTMCrPDAT) were cloned into the yeast expression vector pPICZαB in frame with an α-factor secretion sequence and were expressed in the yeast P. pastoris under the control of a methanol-inducible promoter. The recombinant proteins were purified by affinity chromatography and analyzed by SDS-PAGE. The molecular masses of the recombinant proteins were 95.0 and 120.0 kD, corresponding to the predicted molecular sizes of ΔTMCrPDAT and ORFCrPDAT, respectively. The identities of both recombinant proteins were...
further confirmed by immunoblotting (see Supplemental Figure 4 online) using an antiserum raised against ΔTMCrPDAT protein expressed in Escherichia coli cells (see Supplemental Figure 4A online). These two recombinant proteins were isolated and used for enzyme activity assays.

Typical PDAT Activity of Cr-PDAT

In vitro enzyme activity assays revealed that both ORFCrPDAT and ΔTMCrPDAT mediated the incorporation of DAG into TAG in the presence of different PL substrates (Figure 2A). ΔTMCrPDAT showed slightly higher PDAT activity than ORFCrPDAT under optimized reaction conditions (Figure 2A). The lower activity of the full-length Cr-PDAT was probably due to improper conformational change of the membrane proteins in the aqueous reaction buffer. As shown in Figure 2A, both ORFCrPDAT and ΔTMCrPDAT exhibited a strong preference for anionic PL, including phosphatidic acid (PA), phosphatidylserine (PS), phosphatidylinositol (PI), and phosphatidylglycerol (PG), but showed less activities with the cationic PL (i.e., PC and phosphatidylethanolamine [PE]). Conversely, Sc-PDAT and At-PDAT exhibited strong preferences for PE and PC, respectively (Dahlqvist et al., 2000; Ståhl et al., 2004; Ghosal et al., 2007). The enzymatic activity of ΔTMCrPDAT was determined using PI as an acyl donor and 1,2-DAG as an acyl acceptor. The rate of ΔTMCrPDAT-catalyzed TAG synthesis was proportional to the PI concentration up to 250 μM PI, above which TAG production was inhibited (Figure 2B).

The substrate specificity of ΔTMCrPDAT for different acyl groups at the sn-2 position of PL was also investigated using various PE molecules as acyl donors and 1,2-DAG as acyl acceptors (Figure 2C). The highest enzymatic activity was detected when PE had a linoleic fatty-acyl group (18:2) at the sn-2 position of glycerol, whereas the lowest activity occurred when PE had a stearic (18:0) fatty-acyl group at the sn-2 position. The acyl acceptor specificity of Cr-PDAT was examined with 1,3- and 1,2-DAG, respectively. As shown in Figure 1D, Cr-PDAT exhibited a preference for 1,2-DAG over 1,3-DAG for all species of PLs tested as acyl donors (e.g., PA, PC, PE, PS, PI, and PG). ΔTMCrPDAT preferred DAG with

Figure 1. Cladogram of the LCAT-Like Family Proteins from Plants, Fungi, Microalgae, and Animals.

The neighbor-joining method was used to reconstruct the cladogram. The bootstrap value (obtained from 1000 replicates) is shown on each node. The 0.1 scale represents 10% divergence, calculated as the estimated number of replacement. Squares, plants; triangles, fungi; circles, microalgae; diamonds, animals. [See online article for color version of this figure.]
unsaturated fatty acid side chains (1,2-18:1) compared with the saturated 1,2-16:0 DAG (Figure 2E).

**Cr-PDAT DGTA Activity**

DGTA-catalyzed TAG formation has been observed in some eukaryotic organisms that can use DAG molecules as both acyl donor and acceptor (Lehner and Kuksis, 1993; Stobart et al., 1997). We investigated whether the recombinant ∆TMCrPDAT protein exhibited any DGTA activity by incubating sn-1,2-18:1-DAG with either ORFCrPDAT or ∆TMCrPDAT. TAGs were detected by thin layer chromatography (TLC) analysis, although the conversion efficiency was only one-seventh of its acyltransferase activity (Figure 3A), indicating that Cr-PDAT can indeed use DAG to form TAG.

**Figure 2.** PDAT Activity and Substrate Specificity of Recombinant Cr-PDAT.

(A) PDAT activity of ORFCrPDAT and ∆TMCrPDAT. The sn-1,2-diolein (18:1 DAG) and different PLs, including PA, PS, PI, PG, PC, and PE, were used as substrates. Heat-inactivated PDAT was used as negative control (NC) for the assay. FFA, free fatty acid.

(B) Effects of different PL concentrations on the biosynthesis of TAG.

(C) Acyl donor side-chain specificity of Cr-PDAT. The substrate specificity was determined with PE containing various sn-2-acyl groups, and sn-1,2-diolein was used as an acyl acceptor.

(D) Acyl acceptor positional specificity of Cr-PDAT. The substrate specificity was determined with 1,2-dioleoyl-sn-glycerol (1,2-DAG) and 1,3-dioleoyl-sn-glycerol (1,3-DAG). Various PLs (PA, PC, PE, PS, PI, and PG) were used as acyl donors.

(E) Acyl acceptor side-chain specificity of Cr-PDAT. The substrate specificity was determined with 1,2-dipalmitoyl-sn-glycerol (16:0-DAG) and 1,2-dioleoyl-sn-glycerol (18:1-DAG). PI (250 µM) was used as an acyl donor. ∆TMCrPDAT was used in the enzymatic assays [B] to [E].
Cr-PDAT Galactolipid:DAG Acyltransferase Activity

Considering that Cr-PDAT was predicted to be a chloroplast-localized protein and was previously identified in the chloroplast of C. reinhardtii by proteomics analysis (Terashima et al., 2011), we suspected that Cr-PDAT could use galactolipids, the most abundant membrane lipids in chloroplasts, as substrates by a mechanism similar to the PL:DAG acyl transfer reaction. To test this hypothesis, galactolipids (e.g., MGDG and DGDG) and the sulfolipid sulfoquinovosyldiacylglycerol (SQDG) were used for acyltransferase assays with both ORFCrPDAT and ΔTMCrPDAT. As shown in Figure 3B, ORFCrPDAT was able to use MGDG but not DGDG and SQDG as acyl donors to form TAG. However, ΔTMCrPDAT failed to convert MGDG into TAG, indicating that the transmembrane domain of ORFCrPDAT may be critical for its MGDG:DAG acyltransferase activity. An alternative explanation is that ORFCrPDAT is more hydrophobic than ΔTMCrPDAT and might react only with nonpolar lipids, such as MGDG, under our enzyme assay conditions.

Cr-PDAT Lipase Activity with Broad Substrate Specificity

Noticeable amounts of free fatty acids were detected in the acyl transfer reaction catalyzed by ORFCrPDAT or ΔTMCrPDAT but not in the negative control reactions in which heat-inactivated PDAT was introduced (Figures 2 and 3). This points to a possible role of Cr-PDAT as a lipase. To test this, we conducted a structure-based functional analysis. A sequence comparison and secondary structure prediction revealed that the catalytic site motif (G/A/S-X-S-X-G) appears well conserved among these three PDAT and three template proteins belonging to the α/β hydrolase family (Figure 4A). The secondary structure elements of three layers of α/β/α fold were identified in these proteins as well (Figure 4A).

The possible lipase activity of ΔTMCrPDAT was tested with triolein as a substrate. As shown in Figures 4B and 4C, triolein was degraded rapidly with concomitant formation of free fatty acids, confirming the TAG lipase function of Cr-PDAT.

In comparative modeling analyses, all three-dimensional structure models exhibited an overall α/β hydrolase fold (α/β/α) with a similar steric position (Figures 4D to 4I), except in the case of the topology of At-PDAT, where the first Gly was replaced with Pro and the first two β-sheets of this common fold region were absent (Figure 4F). Previous studies showed an N-terminal deletion version of Sc-PDAT possessed low phospholipase activity in vitro (~2% of transacylation activity), which produced 2% free fatty acids during TAG production (Ghosal et al., 2007). At-PDAT has not been shown to have phospholipase activity (Dahlqvist et al., 2000; Ståhl et al., 2004). These data collectively suggest that the first two β-sheets, conserved in Cr-PDAT and Sc-PDAT, may be essential for the lipase activity.

To determine further the substrate specificity of Cr-PDAT as a lipase, we measured lipase activity of recombinant ΔTMCrPDAT with various classes of lipids and confirmed that Cr-PDAT can hydrolyze a variety of PLs (e.g., PA, PC, PE, PI, PS, and PG) (Figure 5A), neutral lipids (e.g., monoacylglycerol, DAG, and TAG) (Figure 5B), galactolipids (e.g., MGDG and DGDG) (Figure 5C), and steryl esters (Figure 5D). As a phospholipase, Cr-PDAT exhibited a preference for anionic PLs (e.g., PA, PS, PI, and PG) over cationic PLs (e.g., PC and PE) (Figure 5B). As a galactolipase, Cr-PDAT showed a preference for MGDG over DGDG (Figure 5C). ΔTMCrPDAT also exhibited strong steryl hydrolase activity toward cholesteryl esters, whereas ORFCrPDAT had detectable but relatively weak steryl hydrolase activity (Figure 5C).

Overexpression of Recombinant Cr-PDAT in S. cerevisiae

To investigate whether the expression of Cr-PDAT is involved in TAG synthesis in vivo, we introduced the Cr-PDAT gene (pYES: CrPDAT) into S. cerevisiae, with S. cerevisiae transformed with an empty vector as a control (pYES). Both transformants grew in a TAG induction medium for 36 h, and total lipids were extracted for TLC analysis. A basal level of TAG was evident in the pYES cells and increased gradually over 36 h under TAG induction conditions due to endogenous PDAT and DGAT activities.
Figure 4. Cr-PDAT Possesses a Lipase Functional Domain and TAG Lipase Activity.

(A) Sequence alignment and secondary structure elements between the PDAT-like proteins and their referenced templates within α/β hydrolase fold region. The conserved lipase motif is indicated by the green box. The essential Ser (S) residue that constitutes the catalytic triad for the active site is shown as a red asterisk.

(B) TAG lipase activity was detected by monitoring the formation of free fatty acids (18:1) and the degradation of TAG at various time points. Heat-inactivated PDAT was used as negative control (NC); reaction time for the negative control was 60 min. FFA, free fatty acid.

(C) Quantitative analysis of TAG and fatty acids in the enzymatic assay.
Under the same culture conditions, the pYES:CrPDAT cells produced approximately two- to threefold greater amounts of TAG than the control, confirming the contribution of exogenous Cr-PDAT to overall TAG production in the heterologous organism (Figure 6).

**Transient Upregulation of Cr-PDAT in Response to Stress Associated with Membrane Lipid Degradation and TAG Accumulation**

To determine whether Cr-PDAT was involved in stress responses in *C. reinhardtii*, gene expression was investigated by quantitative real-time PCR (Figure 7A). Whereas little change in the relative mRNA level was observed in cells grown under favorable growth conditions [low light and sufficient nitrogen (LL+N)], the transcription of Cr-PDAT was transiently upregulated to a maximum level at 6 h following nitrogen depletion (LL-N), which was 2.9-fold higher than the control (6 h, LL+N). The maximum mRNA level was 3.6-fold greater at 6 h under high light (HL) and nitrogen-depleted conditions (HL-N) than that of the control (6 h, LL+N). After 6 h, Cr-PDAT transcripts decreased rapidly to the basal level at 12 h and remained more or less constant thereafter, regardless of the growth conditions under which the cells had been maintained.

The regulation of Cr-PDAT at the protein level was determined by immunoblotting. Cr-PDAT was transiently upregulated in response to N deprivation (Figure 7B). The protein expression achieved the maximum level at 3 h after the onset of N depletion from the culture medium and then gradually decreased during the following 48 h (Figure 7B).

The concentrations of membrane lipids and TAG in *C. reinhardtii* wild-type cells grown under LL and N-depleted conditions were quantified using electrospray ionization–mass spectrometry (ESI-MS). Membrane lipids, particularly MGDG, SQDG, and PG, were rapidly degraded (Figure 8). For example, MGDG and PG decreased by 63 and 36%, respectively, in the first 2 d of N depletion, while TAG increased ~40 times during this period (Figure 8).

**PDAT Is Responsible for TAG Biosynthesis in *C. reinhardtii***

To investigate the function and possible biological role of PDAT in *C. reinhardtii*, PDAT knockdown mutants were generated with an artificial microRNA (amiRNA)–mediated gene silencing approach (Molnar et al., 2009). A total of 16 PDAT knockdown mutants were selected for gene expression analysis, among which two mutants, P13i and P14i, showed the maximum reduction in *PDAT* transcript (~50% lower than the control transformed with an empty vector) under both N-replete and N-depleted conditions (Figure 9A). The knockdown of PDAT in P13i and P14i was confirmed at the protein level by immunoblotting. As shown in Figure 9B, compared with that of the control transformed with an empty vector, PDAT decreased by 35 and 44% in P13i under N-replete and N-depleted conditions, respectively; it decreased by 65% in P14i under both culture conditions. Under N-replete conditions, P13i and P14i exhibited slower growth than the control in the mid-exponential growth phase (day 3), but the difference became insignificant in the stationary growth phase (Figure 9C).

To determine whether the knockdown of PDAT would affect TAG formation in *C. reinhardtii*, flow cytometry combined with BODIPY staining (a fluorescence dye targeting TAG) was applied to screen for TAG-deficient mutants. As shown in Figure 10A, the fluorescence intensity derived from the PDAT knockdown mutants grown in Tris-acetate phosphate (TAP) for 4 d was significantly lower (P < 0.05) than the control, indicative of a reduction in TAG production in the mutants. A quantitative analysis of TAG by ESI-MS revealed that compared with the control, the TAG content of PDAT knockdown mutants declined by 39.6% in P13i and 57.9% in P14i under N-replete conditions and was reduced by 14.4% in P13i and 28.4% in P14i under N-depleted conditions for 24 h (Figure 10B). These results suggest that PDAT contributed appreciably to the formation of TAG under favorable growth conditions. A 60-fold increase in TAG was evident in both the PDAT knockdown mutants and the control cells under N-depleted conditions after 3 d, and the difference in the maximum TAG concentration in the two types of cells was statistically insignificant (Figure 10B), suggesting that the contribution of the PDAT pathway to TAG synthesis under N deprivation conditions must be minor.

TAG in *C. reinhardtii* comprised three subclasses based on total number of carbon atoms in its fatty acyl groups, namely, TAG C50, C52, and C54 (see Supplemental Figure 5 online). Under N-replete conditions, the TAG C50 molecular species showed a more drastic decline in content than the C52 and C54 molecular species in P13i and P14i compared with the control (Figure 11A). Tandem mass spectrometry analysis indicated that TAG C50 contained two C16 and one C18 fatty acyl group, and C16 was predominant at the sn-2 position of glycerol (see Supplemental Figure 6 online). This suggests that Cr-PDAT prefers DAG molecules with C16 at the sn-2 position in vivo. Because the DAG molecules with C16 at the sn-2 position are generally believed to be synthesized via a prokaryotic pathway in chloroplasts (Frentzen, 1998), the large decrease in TAG C50 indicates that PDAT is involved in TAG biosynthesis in *C. reinhardtii* chloroplasts. After being subjected to nitrogen starvation, the decrease in TAG molecular species was not significant (P > 0.05) between the PDAT knockdown mutants and the control at day 3 (see Supplemental Figure 7 online).

**PDAT-Mediated TAG Synthesis Substrates Derived from Chloroplast Membrane Turnover and Degradation**

To determine if PDAT uses membrane lipids (e.g., PLs and glycolipids) as the substrates for the acyl transfer reaction or hydrolysis in vivo, major membrane lipids of the wild type and...
PDAT knockdown mutants were analyzed by ESI-MS. Although changes in total membrane lipid concentrations (i.e., MGDG, DGDG, etc.) in PDAT knockdowns were almost undetectable under N-replete conditions (see Supplemental Figure 8 online), presumably due to lipid homeostasis that maintained major membrane lipid species at a constant level, knockdown of PDAT dramatically changed the concentrations of several molecular species belonging to the chloroplast membrane lipids, including MGDG, SQDG, and PG (Figures 11B to 11D). The molecular lipid species are denoted according to the number of carbon atoms: total double bonds in the fatty acyl groups, which was annotated based on the mass-to-charge ratio of lipid ions. The higher concentrations of MGDG 34:5 and 34:4, SQDG 34:2 and 34:1, and PG 34:2 in the mutants indicate that Cr-PDAT may participate in chloroplast membrane remodeling or turnover using chloroplast membrane–derived PLs and glycolipids as substrates to form TAG under N-replete conditions. The difference in the membrane lipids, particularly the concentrations of MGDG and PG, between PDAT knockdown mutants and the control strain was insignificant after 2 d of N depletion (see Supplemental Figure 9 online). This suggests that chloroplast membrane remodeling or turnover becomes less active under stress for an extended period of time (>24 h), resulting in fewer membrane lipids available for PDAT-mediated TAG synthesis. It is worth noting that DGDG molecular species varied little in the PDAT knockdown mutants (see Supplemental Figure 10 online), consistent with the in vitro enzymatic assays in which Cr-PDAT did not use DGDG as a substrate for transacylation (Figure 3B) and DGDG was a less favorable substrate than MGDG for hydrolysis (Figure 5).

Although Cr-PDAT showed the highest selectivity on PI among tested membrane lipids in vitro, there was no increase in PI concentration detected in Cr-PDAT mutants compared with the control under both N-replete (see Supplemental Figure 8 online) and N-depleted (see Supplemental Figure 11 online) conditions. This indicates that PDAT and PI may reside in different subcellular compartments. Cr-PDAT harboring a chloroplast transit peptide is localized in chloroplasts (Terashima et al., 2011), whereas PI is absent from the thylakoid membranes (Janero and Barnett, 1981) and is likely enriched in extraplastidic membranes of C. reinhardtii.

**DISCUSSION**

**Cr-PDAT Is a Multifunctional Enzyme**

The biosynthesis of TAG is a common metabolic pathway that occurs in almost all plants, animals, fungi, and some bacteria.
Much research has focused on the Kennedy pathway, in which DGAT catalyzes the acylation of DAG to form TAG. Recently, an acyl-CoA–independent pathway catalyzed by PDAT was discovered in algae, yeasts, and plants (Banaś et al., 2000; Dahlqvist et al., 2000; Boyle et al., 2012). However, the biochemical characteristics and physiological importance of PDAT in algae have not been investigated. In this study, in vitro enzyme assays of recombinant full-length and DTM Cr-PDAT provided evidence that Cr-PDAT is a multifunctional enzyme with acyltransferase and lipase activities with broad substrate specificity. For its transacylation function, Cr-PDAT can use PLs, galactolipids, and DAG as acyl donors for TAG synthesis. For its lipase function, Cr-PDAT can release free fatty acids from a variety of lipid substrates, including PLs, galactolipids, and neutral lipids. Cr-PDAT may contribute to membrane remodeling and degradation, with TAG being a by-product of this process.

Galactolipids are the major lipid building blocks of chloroplast membranes. In vascular plants, drastic decreases in MGDG content occur during freezing, desiccation, or ozone treatment (Sakaki et al., 1990a, 1990b; Gigon et al., 2004; Li et al., 2008a). In this study, we observed that the MGDG content of C. reinhardtii decreased continuously over 48 h under N-depleted conditions (Figure 8). It is well documented that, under stress, a decrease in MGDG is usually accompanied by TAG accumulation (Sakaki et al., 1990a, 1990b; Gigon et al., 2004), but the pathways responsible for this conversion are poorly understood. A pathway mediated by a galactolipid:galactolipid galactosyltransferase (GGGT) located on the chloroplast envelope membrane (van Besouw and Wintermans, 1978; Heemskerk et al., 1983) may play a role in the lipid remodeling process (Sakaki et al., 1990b; Benning and Ohta, 2005; Moellering and Benning, 2011). GGGT transfers a galactosyl moiety from an MGDG molecule to a second galactolipid molecule (MGDG or DGDG), resulting in the formation of DAG, the precursor for TAG synthesis, and oligogalactolipids. Recently, SENSITIVE TO FREEZING2 (SFR2), a gene essential for freezing tolerance, has been identified as the GGGT-encoding gene in Arabidopsis thaliana. The sfr2 mutant showed ~50% reduction in TAG content compared with the wild type in response to freezing treatment, indicating a role for GGGT in TAG synthesis in vivo (Moellering et al., 2010).

In this study, we discovered that Cr-PDAT acted as a galactolipid:DAG acyltransferase, transferring a fatty acyl group from MGDG to DAG to form TAG. As such, knockdown of Cr-PDAT led to the accumulation of certain MGDG species and reduced TAG accumulation in the mutants. Therefore, the PDAT pathway represents a previously unrecognized mechanism for galactolipid remodeling, turnover, and degradation. From this perspective, TAG can be regarded as a coproduct of Cr-PDAT-mediated chloroplast membrane remodeling or degradation. From a broader perspective, the formation of this coproduct is part of lipid homeostasis within the chloroplast or the whole cell. Cr-PDAT showed high substrate specificity for galactolipid:DAG transacylation in that only MGDG could be efficiently utilized as an acyl donor. The MGDG-dependent galactolipid:DAG acyltransferase function of Cr-PDAT is consistent with the fact that this protein harbors a chloroplast transit peptide and was detected in the chloroplast (Terashima et al., 2011), where MGDG...
exclusively resides. Galactolipid:DAG acyltransferase activity has not been reported for any previously known PDAT. It is unknown if such a function is restricted to Cr-PDAT or is common to most, if not all, photosynthetic organisms. When various PLs were used as acyl donors and diolein was used as an acyl acceptor, Cr-PDAT showed a preference for anionic PLs (e.g., PG, PI, PA, and PS) over cationic PLs (e.g., PC and PE) (Figure 2A). This is distinct from Sc-PDAT and At-PDAT, which exhibited higher substrate specificities toward PC and PE than the other PLs (Banaś et al., 2000; Dahlqvist et al., 2000; Ghosal et al., 2007). This difference can be explained in part by the distinct cellular PL compositions among yeasts, plants, and green algae. In yeast and vascular plants, PC and PE represent the most abundant PLs, whereas PG and PI are the major PLs, and PC is absent in C. reinhardtii (Giroud et al., 1988). Based on these observations, we speculate that cellular membranes undergo highly dynamic and efficient remodeling and turnover to maintain proper functions under various environmental conditions; thus, the substrate specificity of PDAT has coevolved with the lipid composition of various organisms.

In addition to PL/galactolipid:DGAT activity, Cr-PDAT also exhibited moderate DGTA activity. An important role for DGTA in TAG synthesis has been proposed in developing seeds of sunflower (Helianthus annuus) (Fraser et al., 2000), endosperm of castor bean (Ricinus communis) (Mancha and Stymne, 1997), and developing safflower seeds (Stobart et al., 1997). Although one DGTA has been purified and characterized from rat intestinal microsomal preparations (Lehner and Kuksis, 1993), understanding of this pathway has been hindered by the lack of the genomic information on this enzyme. Recently, Sc-PDAT was found to have DGTA activity, which was ~11 or 2% of its PDAT activity, when PC and PE were used as acyl donors, respectively (Ghosal et al., 2007). At-PDAT also displayed DGTA activity of about one-tenth of its PDAT activity (Ståhl et al., 2004). Under our experimental conditions, the DGTA activity of Cr-PDAT was approximately one-seventh of its PDAT activity. These results suggest that the DGTA function is a common feature of PDAT in plants, yeasts, and green algae. Considering that DAG can be produced via multiple routes in cells, such as by dephosphorylation of PA catalyzed by phosphatidic acid phosphatase in the Kennedy pathway or GGGT-mediated remodeling of galactolipids, PDAT may work in concert with these enzymes to further convert DAG into TAG via PDAT-mediated DAG:DAG or PL/galactolipid:DAG transacylation routes.

A number of enzymes involved in lipid metabolism have been reported to be multifunctional proteins, catalyzing both transacylation reactions and lipid hydrolysis. For example, oleosin, the

Figure 8. Lipid Content of C. reinhardtii (Wild Type) under N-Replete and N-Depleted Conditions.

For TAG induction, cells grown in TAP medium under low light for 4 d were transferred to TAP-N medium. Lipids were extracted and analyzed using ESI-MS. Data are presented as means ± se (n = 4 to 6). DW, dry weight.

Figure 9. Generation of amiRNA Knockdown Mutants.

(A) Knockdown of Cr-PDAT in P13i and P14i compared with the control transformed with an empty vector (TC), confirmed with real-time PCR.

(B) Immunoblotting analysis of protein level in the knockdown mutants and the control (TC). Protein concentration was determined using Image J software and normalized to the control.

(C) Growth kinetics of the knockdown mutants (P13i and P14i) and the control (TC) under N-replete conditions. Data are presented as means ± se (n = 4 to 6), with the asterisk indicating P < 0.05.
structural protein of plant lipid bodies, is a bifunctional enzyme that exhibits both monoacylglycerol acyltransferase and phospholipase activities (Parthibane et al., 2012). The yeast TAG lipase Tgl4p exerts multiple functions as a TAG lipase, steryl ester hydrolase, phospholipase, and acyltransferase (Rajakumari and Daum, 2010a). The multiple functions of the above chimeric enzymes are due to the fusion of two distinct functional domains (acyltransferase and lipase domains) into a single protein.

By contrast, Cr-PDAT contains a single functional LCAT domain, which shares the Ser/Asp-Glu/His triad with lipases, esterases, and proteases (Peelman et al., 1998). Both LCAT and lipase belong to the α/β hydrolase fold family; like lipase, LCAT contains a potential lid domain involved in substrate binding (Peelman et al., 1998). The ability of an enzyme to catalyze an adventitious secondary activity at the active site responsible for the primary activity has been termed “catalytic promiscuity” (Copley, 2003; Khersonsky et al., 2006). Despite the potential catalytic promiscuity in the LCAT domain, the functions of most LCAT-like proteins appear to be highly divergent. For instance, out of the six LCAT-like proteins identified in Arabidopsis, LCAT3 (At3g03310) and LCAT4 (At4g19860) function as phospholipase A1 (Noiriel et al., 2004; Chen et al., 2012), whereas PDAT1 (At13640) shows PDAT activity but no phospholipase activity (Ståhl et al., 2004).

Unlike the vascular plants, C. reinhardtii contains a single LCAT-like protein, Cr-PDAT, that displays significant catalytic promiscuity. We speculate that a multifunctional Cr-PDAT may be particularly relevant to C. reinhardtii and other organisms with small and compact genomes; synthesis of a single PDAT with multiple functions allows better resource utilization and management for maintaining lipid homeostasis necessary for growth and reproduction, as well as for adapting to a changing environment.

Physiological Role of Cr-PDAT in Vivo

The multiple enzyme activities of Cr-PDAT suggest possible physiological roles of this enzyme in C. reinhardtii. Heterologous overexpression and gene knockdown experiments, along with its regulation at the transcript and protein level, suggest that Cr-PDAT is involved in TAG synthesis in vivo. The mRNA level of Cr-PDAT was upregulated under N-depleted or HL conditions, similar to the gene expression patterns of other acyltransferases related to TAG synthesis, suggesting a role for Cr-PDAT in TAG synthesis in vivo.

Knockdown of Cr-PDAT led to increased concentration of molecular species of MGDG, SQDG, and PG in vivo, in addition to the impaired TAG synthesis (Figure 11). The ESI-MS results indicated that in the PDAT knockdown mutants, TAG species with the most drastic reduction contained two C16 and one C18 fatty-acyl groups (see Supplemental Figure 7 online), consistent with observations of increased chloroplast membrane lipid species enriched in C16 fatty-acyl groups at the sn-2 position of glycerol (Giroud et al., 1988). We conclude that Cr-PDAT uses chloroplast membrane lipids, particularly MDGG, SQDG, and PG, as substrates to synthesize TAG in vivo. Lipidomics analyses also indicate that Cr-PDAT preferentially uses DAG synthesized via a prokaryotic pathway as substrates in vivo. Considering that Cr-PDAT harbors a chloroplast transit peptide and is present in the chloroplast proteome (Terashima et al., 2011), we suggest that Cr-PDAT is involved in TAG synthesis in the chloroplast. The precise localization (e.g., thylakoid membrane or chloroplast envelope) of Cr-PDAT within the chloroplast, however, remains unknown.

TAG in eukaryotic cells is thought to be a deposit for fatty acids, which in turn serve as a carbon and energy reserve. However, recent studies have revealed that TAG homeostasis is linked to membrane lipid turnover and remodeling, processes that are vital to maintaining proper structure and fluidity of cellular membranes (Rajakumari and Daum, 2010a, 2010b; Gaspar et al., 2011). Membrane remodeling and turnover involve a
deacylation-reacylation cycle in which fatty acids are removed from PLs and are then reacylated to yield lysophospholipids (Lands, 1960; Kennedy, 1961; Lands and Merkl, 1963; Merkl and Lands, 1963; Bates et al., 2009). Fatty acids at the sn-2 position of PLs can be removed by (1) phospholipase A2, which produces free fatty acids and 1-acyl lysophospholipids (Winstead et al., 2000; Kudo and Murakami, 2002); and (2) PDAT, which can bypass the production of free fatty acids by incorporating the cleaved fatty acids directly into TAG. In the latter case, TAG synthesis is regarded as a buffer system for detoxification of lipotoxic free fatty acids (Listenberger et al., 2003; Petschnigg et al., 2009).

Our results indicate that the contribution of PDAT to TAG synthesis in *C. reinhardtii* is prominent in the logarithmic growth phase under favorable growth conditions but less crucial under stress conditions. A similar physiological role of PDAT has been observed in the yeast *S. cerevisiae*, where it is a major contributor to TAG accumulation in the exponential growth phase but not in the stationary phase (Oelkers et al., 2000, 2002). We also observed that Cr-PDAT knockdown resulted in a small but noticeable decrease in the specific growth rate in logarithmically growing cells. Collectively, we concluded that PDAT-mediated membrane remodeling and TAG synthesis is pivotal in vigorously growing cells and indispensable for normal growth under favorable culture conditions. It also plays an important role in chloroplastic degradation and TAG synthesis at the early stages (first 1 to 2 d) of N depletion. As stress persists, however, Cr-PDAT may become less critical. Massive accumulation of TAG in *C. reinhardtii* under N depletion or other stress conditions may be attributable to other TAG synthesis pathways such as the Kennedy pathway.

Although our results suggest the Cr-PDAT pathway may play a minor role in TAG synthesis under N-depleted conditions over extended period of time (Figure 10; see Supplemental Figure 7 online), we cannot rule out the possibility that PDAT-mediated TAG production may overlap with other TAG synthesis enzymes in *C. reinhardtii*, compensating for the knockdown of PDAT under stress. In a previous study, an *Arabidopsis* knockout mutant with a T-DNA insertion in the *PDAT1* locus did not display altered seed fatty acid content or composition (Mhaske et al., 2005), but RNA interference silencing of *PDAT1* in a *dagt1-1* background plant resulted in a 70 to 80% decrease in seed oil content (Zhang et al., 2012), suggesting that DGAT can compensate for PDAT knockout in a wild-type background. This may be also true for *C. reinhardtii*, which contains at least one *DGAT1* and five *DGAT2* genes. The unaltered TAG content in knockdown mutants of *C. reinhardtii* under stress conditions can also be explained by the TAG lipase activity of Cr-PDAT, which was likely activated when cells were grown under N-depleted conditions for 3 d. To address these questions, further research into the precise regulation of Cr-PDAT functions in vivo is needed.
Biotechnological Implications

In this study, we demonstrated that in an aqueous reaction system, recombinant Cr-PDAT can efficiently hydroyaze neutral lipids (TAG, DAG, and MAG), PLs, and galactolipids to produce free fatty acids (Figures 4 and 5). The multifunction of Cr-PDAT as an acyltransferase and a lipase with broad substrate specificity offers great potential as a biocatalyst for the conversion of crude algal oil into biofuels and non-fuel oils. Large-scale production of Cr-PDAT and evaluation of PDAT-based direct conversion of crude algal oils into free fatty acids and free fatty acids methyl esters is in progress.

METHODS

Strains, Media, and Growth Conditions

The wild-type Chlamydomonas reinhardtii strain cc-1690 was used for the cDNA library construction and gene expression analysis, and the cell wall-deficient wild-type strain cw-15 was used to generate the mutants P13i and P14i using an amiRNA knockdown method. All strains were grown in TAP medium containing 7.5 mM NH₄Cl (Harris, 1989) at 23°C under continuous illumination of 40 µmol photons m⁻² s⁻¹. For HL (400 µmol photons m⁻² s⁻¹), cells in the exponential growth phase raised under LL conditions (40 µmol photons m⁻² s⁻¹) were harvested by centrifugation, washed once with TAP medium and then resuspended in fresh TAP medium at a starting cell density of 10⁶ cells mL⁻¹. Cultures were then exposed to HL for 3 d. To impose nitrogen deprivation (-N), cells in the stationary growth phase (corresponding to a cell density of 2 x 10⁷ cells mL⁻¹) were collected by centrifugation and washed with nitrogen-deficient TAP medium (TAP-N), and equal amounts of cells were re-suspended in TAP-N medium. Cell aliquots were collected at defined time intervals after being transferred to TAP or TAP-N medium. All cultures were grown on a shaking table with continuous shaking at 130 rpm.

Yeast, Bacterial Strains, and Culture Conditions

The Pichia pastoris wild-type X-33 strain (Invitrogen) was used to express the secernatory recombinant Cr-PDAT proteins. The wild type Saccharomyces cerevisiae strain INVSc1 (MATα his3D1 leu2 trp1-289 ura3-52; Invitrogen) was used as a heterologous host for intracellular overexpression of Cr-PDAT. Escherichia coli strain TOP10 (Invitrogen) was used for recombinant DNA work. All strains and transformants generated were grown according to manufacturer’s instructions.

RNA Isolation and cDNA Library Construction

Harvested C. reinhardtii cells (~7 g wet weight) were pulverized in a mortar with liquid nitrogen, and extracts were transferred to a tube containing Trizol reagent (Invitrogen). Total RNA isolation was conducted following the procedure described by Li et al. (2008b). To construct a high-quality cDNA library, RNA was isolated from cells subjected to the various treatments described above (e.g., LL, HL, and -N). Poly(A)⁺ RNA was purified from pooled RNA using the Oligotex mRNA purification kit (Qiagen). About 5 µg of mRNA was used to synthesize double-stranded cDNA, and the cDNA was ligated into the λ-ZAPII arms to construct a cDNA library using the GigapackIII Gold packaging kit (Stratagene). An aliquot of the cDNA library was mass excised by ExAssist helper phage (Stratagene) to recover pBluescript plasmids.

Quantitative Real-Time PCR

For real-time RT-PCR analysis, the first strand of the cDNA was synthesized using the Taqman reverse transcription system (Applied Biosystems) per the manufacturer’s instructions. Real-time RT-PCR was performed on an ABI Prism 7500 sequence detection system (Applied Biosystems) following the protocol previously described by Li et al. (2008b). Primer sequences used for expression studies are listed in the Supplemental Table 1 online. RNA levels were normalized using the 18S rRNA gene as the internal standard.

Identification and Cloning of cDNA Encoding PDAT in C. reinhardtii

The S. cerevisiae PDAT (Sc-PDAT) and Arabidopsis thaliana PDAT (At-PDAT) protein coding sequences were used to identify the C. reinhardtii genes with the highest similarity using the BLAST server (http://blast.ncbi.nlm.nih.gov/). One hit revealed a significant similarity to these known PDAT sequences. Several primers (see Supplemental Table 1 online) were designed based on the sequences conserved with other PDAT orthologs to clone the coding region of cDNA from the C. reinhardtii cDNA library. All PCR reactions used in this study were performed in the presence of 5% DMSO. PCR products of the expected size were cloned into the pCR2.1 cloning vector (Invitrogen) and verified by sequencing. Full-length cDNA was obtained following screening the cDNA library (see Supplemental Methods 1 online).

Bioinformatics Analysis

Protein sequence alignments and similarity analyses were conducted with the CLC Main Workbench program (CLC Bio). Primary structure analyses, topological organization predictions, and posttranslational modification predictions were performed with ExPaSy proteomic tools available at http://www.expasy.ch/tools/. Protein motifs were identified using InterProScan at http://www.ebi.ac.uk/Tools/psippardb/psicat/. For analyses of putative transport peptide and signal sequences, ChloroP 1.1 ( Emanuelsson et al., 1999) and SignalP 4.0 ( Petersen et al., 2011) were used.

Phylogenetic analyses were conducted using neighbor-joining methods. The neighbor-joining tree was constructed with MEGA 5.0. Molecular distances within the aligned sequence were calculated according to the position correction model. Branch points were tested for significance by bootstrapping with 1000 replications ( Tamura et al., 2007). The alignment used for the analysis is available as Supplemental Data Set 1 online.

To select potential templates for building three-dimensional structures of the PDAT-like protein (Cr-PDAT, Sc-PDAT, and At-PDAT), the template identification tool in SWISS-MODEL (http://swissmodel.expasy.org/workspace/) was used as a comparative modeling method. Secondary structure elements were constructed based on sequence alignments generated by the above methods. The three-dimensional models of the PDAT-like proteins were built with the SWISS-MODEL workspace program using selected structures of α/β hydrolases of Pseudomonas lemoignei (Protein Data Bank accession ID: 2X5X; chain A), Pseudomonas aeruginosa (Protein Data Bank accession ID: 1EXA; chain A), and Burkholderia cepacia (Protein Data Bank accession ID: 1Y51) as templates. All figures illustrating molecular modeling results were prepared in PyMOL (Molecular Graphics System).

Heterologous Expression of Cr-PDAT in P. pastoris and S. cerevisiae

For each host system, two types of plasmids containing full-length Cr-PDAT (designated as ORFCrPDAT) and the Cr-PDAT coding sequence without the putative membrane-spanning region (∆TM-PDAT), respectively, were constructed. Details on the heterologous expression of ORFCrPDAT and ∆TMcrPDAT in E. coli, P. pastoris, and S. cerevisiae are given in Supplemental Methods 1 online.

Expression of Cr-PDAT in E. coli and Production of Anti-PDAT Antibodies

To produce antibodies against Cr-PDAT, ∆TMcrPDAT was amplified using ∆TM-PDATEF/∆TM-PDATER (see Supplemental Table 1 online)
and cloned into the expression vector pQE30 (Qiagen). The resulting pQE30 constructs were transformed into the E. coli strain M15 containing pREP4 (Qiagen). For expression, positive transformants were expressed by adding isopropyl-β-D-thiogalactopyranoside (0.1 mM) after 4 h of growth. Cells were grown for additional 6 h before being harvested (see Supplemental Figure 4 online). The mouse anti-TMCrPDAT polyclonal antibodies were produced by Precision Antibody.

**Immunoblotting**

To isolate total proteins, 5 mL cell cultures were centrifuged at 1000g for 5 min and washed with 50 mM Tris-HCl, pH 6.8. Cell pellets were resuspended in lysis buffer (50 mM Tris-HCl, pH 6.8, containing 2% SDS and 10 mM EDTA) and a protease inhibitor cocktail (Sigma-Aldrich). Cells were incubated for 1 h at room temperature, and insoluble cell debris was removed by centrifugation at 13,000g for 30 min at 4°C prior to loading samples onto the SDS-PAGE. Total protein concentration of supernatants (whole cell extract) was determined using the SPN protein assay kit (G Biosciences). Protein samples dissolved in SDS-PAGE sample buffer were separated on SDS-PAGE and transferred to nitrocellulose membranes. The membranes were incubated for 12 h with recombinant anti-TMCrPDAT antibody diluted 1:100 with PBS containing 5% (w/v) nonfat milk and 0.1% (v/v) Tween 20, followed by incubation with anti-mouse IgG conjugated with horseradish peroxidase (Bio-Rad) diluted 1:2000 in the same buffer. Antigen-antibody complexes were visualized using an enhanced chemiluminescence detection kit (Pierce).

**Extraction of Lipids from Yeast Cells and TLC Analysis**

For total lipid extraction, yeast cells were collected by centrifugation and suspended in lysis buffer (50 mM Tris-HCl, pH 6.8, containing 2% SDS and 250 µM lipid donors (e.g., PA, PC, PE, PS, PI, PG, MGDG, DGDG, and SQDG) and 250 µM lipid acceptor (1,2- or 1,3-DAG) in a final reaction volume of 200 µL. The suspension was thoroughly mixed and incubated at 30°C for 30 min. Lipids were extracted and analyzed as described above. Lipolytic acyl hydrolase activities of recombinant PDAT was performed 30°C for 30 min. Lipids were extracted and analyzed as described above. Lipolytic acyl hydrolase activities of recombinant PDAT was performed 30°C for 30 min. Lipids were extracted and analyzed as described above. Lipolytic acyl hydrolase activities of recombinant PDAT was performed 30°C for 30 min. Lipids were extracted and analyzed as described above.

**Enzyme Assays**

For the acyltransferase activity assay, the reaction mixture contained 1 µg recombinant PDAT in 50 mM potassium phosphate buffer, pH 7.2, with 250 µM lipid donors (e.g., PA, PC, PE, PS, PI, PG, MGDG, DGDG, and SQDG) and 100 µM lipid acceptor (1,2- or 1,3-DAG) in a final reaction volume of 200 µL. The suspension was thoroughly mixed and incubated at 30°C for 30 min. Lipids were extracted and analyzed as described above. Lipolytic acyl hydrolase activities of recombinant PDAT was performed 30°C for 30 min. Lipids were extracted and analyzed as described above. Lipolytic acyl hydrolase activities of recombinant PDAT was performed 30°C for 30 min. Lipids were extracted and analyzed as described above. Lipolytic acyl hydrolase activities of recombinant PDAT was performed 30°C for 30 min. Lipids were extracted and analyzed as described above.

**Mass spectrometry analysis**

Mass spectrometry analysis was performed with an Agilent 6460 triple quadrupole liquid chromatography/mass spectrometer equipped with an electrospray ion source (Agilent). Single-stage mass spectrometry was used for the detection of total positive and negative lipid ions. Precursor ions and neutral loss scanning were employed for membrane glycerolipid identification according to previously described methods (Welti et al., 2002, 2003; Hsu and Turk, 2009); sequential neutral loss scanning was applied for TAG identification (Han and Gross, 2001, 2003). Nitrogen was used as a nebulizing gas (at 0.3 Bar) and a dry gas (4 liters min⁻¹ at 200°C). The spray capillary voltage was 3700 V for the negative ion mode and 4200 V for the positive ion mode analysis. Prior to direct infusion into the ESI source, 10 µL lipid extract was mixed with internal standards (see Supplemental Table 2 online), the ionization reagent (200 µM Nal for positive ion mode and 500 µM NH₄OH for negative ion mode), and chloroform:methanol (1:1) for a final volume of 100 µL. A five-microliter sample mixture was directly infused into ESI by loop injections with methanol as a mobile phase at a flow rate of 0.1 mL min⁻¹.

For analysis of TAG in cells grown under normal culture conditions, 3-µL tripalmintin-supplemented samples were injected into a 5 cm × 4.6-mm Eclipse C-18 column at 40°C with liquid chromatography solvent at 0.3 mL min⁻¹. The liquid chromatography gradient was as follows: 40% methanol, 50% acetonitrile, and 10% isopropanol, 1 min, followed by 10% methanol, 50% acetonitrile, and 40% isopropanol, 30 min. Nal (10 mM) in acetone was added by Hamilton syringe pump at a flow rate of 6 µL min⁻¹ via a tee connection into the eluent stream of HPLC prior to...
mass spectrometry. For fatty acyl group identification, TAG lipid ions selected as precursor ions were subjected to collision energy (50 V). The fatty acyl groups were identified based on the fragments deriving from neutral loss (NL) of individual fatty acyl groups. Fragments corresponding to the NL of sn-2 fatty acyl groups are known to yield lower abundances than those from the NL of sn-1/3 fatty acyl groups (Guella et al., 2003); thus, the abundances of tandem mass spectrometry product fragments were used for regiochemical determination.

For quantification of lipid content, calibration standards of each lipid class (see Supplemental Table 2 online) were titrated relative to a constant amount of internal standards. The peak intensity ratios of calibration standards relative to internal standards were plotted against their molar concentration ratios to establish the standard curve.

Accession Numbers
Accession numbers of protein sequences used in this study are given in Supplemental Table 3 online.

Supplemental Data
The following materials are available in the online version of this article:

Supplemental Figure 1. Gene Structure of Cr-PDAT and Amino Acid Sequence Alignment with Other LCAT-Like Proteins.

Supplemental Figure 2. Full-Length Nucleotide Sequence and Deduced Amino Acid Sequence of the Chlamydomonas PDAT Gene.

Supplemental Figure 3. Cr-PDAT Is a Membrane Protein with One Predicted Transmembrane Domain.

Supplemental Figure 4. Recombinant Cr-PDAT Protein Expression in E. coli and P. pastoris.

Supplemental Figure 5. TAG Composition of Wild-Type C. reinhardtii Grown under N-Replete Conditions.

Supplemental Figure 6. Fatty Acyl Groups of TAG C50 Determined by MS/MS Analysis.

Supplemental Figure 7. Concentration of TAG Molecular Species in Cr-PDAT amiRNA Knockdown Mutants Grown under N-Depleted Conditions.

Supplemental Figure 8. Total Membrane Lipid Concentrations in the Cr-PDAT Knockdown Mutants under N-Replete Conditions.

Supplemental Figure 9. MGDG, SQDG, and PG Molecular Species Concentrations in the Cr-PDAT Knockdown Mutants under N-Depleted Conditions.

Supplemental Figure 10. DGDG Species Concentrations in the Cr-PDAT Knockdown Mutants under N-Replete and -Depleted Conditions.

Supplemental Figure 11. PI Concentrations in the Cr-PDAT Knockdown Mutants under N-Depleted Conditions.

Supplemental Figure 12. Positive and Negative Controls for Lipase Activity Assays.

Supplemental Table 1. Primers Used for PDAT Gene Cloning and PCR Analysis.

Supplemental Table 2. Lipid Standards for ESI-MS Analysis.

Supplemental Table 3. GenBank Accession Numbers of Protein Sequences Used in This Study.

Supplemental Methods 1. cDNA Library Screening and Heterologous Expression of Cr-PDAT in E. coli, P. pastoris, and S. cerevisiae.

Supplemental Data Set 1. Text File of the Alignment Used to Generate the Phylogenetic Tree in Figure 1.

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AUTHOR CONTRIBUTIONS
K.Y. performed molecular cloning, recombinant protein expression, enzymatic assays, and bioinformatics analyses. D.H. developed the mass spectrometry method and performed in vivo functional characterization. Y.L. performed gene expression analysis and generated knockdown mutants. Q.H. designed the research. All of the authors contributed to analyzing the data and writing the article.

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Phospholipid:Diacylglycerol Acyltransferase Is a Multifunctional Enzyme Involved in Membrane Lipid Turnover and Degradation While Synthesizing Triacylglycerol in the Unicellular Green Microalga *Chlamydomonas reinhardtii*

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