Dynamic transcriptome analysis identifies genes related to fatty acid biosynthesis in the seeds of *Prunus pedunculata* Pall

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**Abstract**

**Background:** *Prunus pedunculata* Pall, the deciduous shrub of *Amygdalus* subgenus in *Rosaceae*, is a new kind of desert oil-bearing tree. It has a long story of being planted in the West and North of China for sand fixation and desert control. In addition, the seeds of *P. pedunculata* are rich of oil, especially the monounsaturated fatty acid and polyunsaturated fatty acid. However, little is known about the molecular mechanisms of oil accumulation during the seed development of *P. pedunculata*.

**Results:** The seeds of *P. pedunculata* from three independent plants at 10, 18, 24, 31, 39, 45, 59 and 73 days after flowering (DAF) were obtained and the oil compositions were evaluated. It showed that oleic acid was the dominant type of oil content in the mature seeds (from 32.724% at 10DAF to 72.06% at 73DAF). Next, transcriptome sequencing for the developing seeds produced 988.795 million high quality reads and TRINITY assembled 326,271 genes for the first transcriptome for *P. pedunculata*. After the assembled transcriptome was evaluated by BUSCO with 85.9% completeness, we identified 195,342, 109,850 and 121,897 *P. pedunculata* genes aligned to NR, GO and KEGG pathway databases, respectively. Then, we predicted 23,229 likely proteins from the assembled transcriptome and identified 1917 signal peptides and 5512 transmembrane related proteins. In the developing seeds we detected 91,362 genes (average FPKM > 5) and correlation analysis indicated three possible development stages — early (10 ~ 24DAF), middle (31 ~ 45DAF) and late (59 ~ 73DAF). We next analyzed the differentially expressed genes (DEGs) in the developing seeds. Interestingly, compared to 10DAF the number of DEGs was increased from 4406 in 18DAF to 27,623 in 73DAF. Based on the gene annotation, we identified 753, 33, 8 and 645 DEGs related to the fatty acid biosynthesis, lipid biosynthesis, oil body and transcription factors. Notably, *GPAT*, *DGD1*, *LACS2*, *UBC* and *RINO* were highly expressed at the early development stage, *ω6-FAD*, *SAD*, *ACP*, *ACCA* and *AHG1* were highly expressed at the middle development stage, and *LAC5*, *DGD1*, *ACAT1*, *AGPAT*, *WSD1*, *EGY2* and oleosin genes were highly expressed at the late development stage.

**Conclusions:** This is the first time to study the developing seed transcriptome of *P. pedunculata* and our findings will provide a valuable resource for future studies. More importantly, it will improve our understanding of molecular mechanisms of oil accumulation in *P. pedunculata*.

**Keywords:** Fatty acid, Oleic acid, Oil accumulation, Developing seeds, *Prunus pedunculata*, Transcriptome
Background

Prunus pedunculata Pall (P. pedunculata), belongs to the deciduous shrub of Amygdalus subgenus in Rosaceae, is a new kind of desert oil-bearing tree, which is also known as wild cherry, almond with stem and hairy cherry [1]. This rare and endangered wild tree species is naturally distributed in the mountainous and desert areas of arid and semi-arid areas in the Northwest China. Due to its strong adaptability, disease and insert resistance, cold and drought resistance, developed root system and long survival period, P. pedunculata has a long history of being planted in the West and North of China for sand fixation and desert control [2]. In addition, the P. pedunculata seeds are rich in protein and oil. The monounsaturated fatty acid and polyunsaturated fatty acid have been reported to be 69.11% and 28.77%, respectively, in the seeds of P. pedunculata [3]. However, much is unknown about P. pedunculata and their seeds.

The oil formation process in plants consists of four steps, including i) fatty acid de novo synthesis, ii) acyl elongation and editing, iii) triacylglycerol (TAG) assembly and iv) oil drop formation [4]. Several pathways, genes and proteins have been reported to be involved in the oil formation process. For example, the fatty acid is localized to plastid while the assembly of TAG molecule occurs outside the plastid and is associated with the oil body [5]. The assembly of fatty acids occur on the ACP (acyl carrier protein) through a cycle of four reactions which elongate the acyl chain by 2 carbons and a total of 7 cycles are required to form the saturated 16 carbon acyl-ACP [6]. The content of fatty acid subtypes are decided by the activities of FATA (acyl-ACP thioesterase A), FATB (acyl-ACP thioesterase B), 18:0-ACP desaturase (SAD) and KASII (β-ketoacyl ACP synthase II) [6]. Although large is unknown about the transport of free fatty acid products from the plastid, it is implicated that LACS (long chain acyl-CoA synthetase) on the outer plastid envelope may function in the formation of acy-CoA, which is the substrate for glycerolipid assembly [4]. The esterification of newly synthesized fatty acid to phosphatidylcholine is reported to occur at the plastid envelope via LPCAT (acyl-CoA:lyso-phosphatidylcholine acyltransferase) [7]. The assembly of TAG from G-3-P (glycerol-3-phosphate) involves some key enzymes, including GPAT (glycerol-3-phosphate acyltransferase), LPAAT (lysophosphatidic acid acyltransferase), PAP (phosphatidic acid phospatase) and DGAT (diacylglycerol acyltransferase) [8]. Transcription factors (TFs) WRI1 (ethylene-responsive transcription factor WRI1) and LEC1 (nuclear transcription factor Y subunit B-9) has been reported to control the expression of more than 15 enzymes (e.g., pyruvate dehydrogenase), which are required for the synthesis of fatty acid and the determination of the oil content in plant seeds [9, 10].

Transcriptome sequencing has enabled the identification of genes involved in the seed development and their association with the oil content in plant seeds. Fei utilized the transcriptome sequencing for the five seed development stages of Zanthoxylum bungeanum and identified 20 genes related to the fatty acid synthesis, such as ENR, ECR and SADI [11]. Abdullah performed transcriptome sequencing for the developing seeds of Camelina sativa and identified 7932 genes involved in the triacylglycerol biosynthesis and accumulation [12]. Feng assembled the transcriptome for Eucommia ulmoides and reported 65 genes involved in fatty acid biosynthesis including FABG (3-oxoacyl-ACP reductase), KASII and FABI (enoyl-ACP reductase I) [13]. Yang reported 124 genes (e.g., GmABI3b, GmNFYA, GmFAD2-1B) potentially affecting the soybean oil content by analyzing the dynamic transcriptome of developing soybean seeds [14]. Li analyzed the de novo transcriptome of developing tree peony seeds and identified 388 genes (e.g., SAD, FAD2, FAD8) that might be involved in de novo fatty acid and TAG biosynthesis [15]. In addition, Wang identified 211 genes and 35 proteins associated with the fatty acid metabolism pathway, 63 genes and 11 proteins associated with the biosynthesis of unsaturated fatty acids, and 115 genes and 24 proteins associated with ALA (alpha-linolenic acid) metabolism in the tree peony seeds [16]. Kim identified 540 unique perilla genes involved in all known pathways of acyl-lipid metabolism by analyzing the transcriptome of seeds and leaves of Perilla frutescens [17]. However, little is known about the gene changes and their association with fatty acid synthesis during the seed development of P. pedunculata seeds.

In the present study, we examined the oil content of seven fatty acid subtypes in the P. pedunculata seeds at eight developing stages and performed transcriptome sequencing. We assembled the transcriptome for P. pedunculata seeds and annotated them. Then, differentially expressed genes (DEGs) were identified during the seed development and they might be related to the fatty acid synthesis. Weighted gene co-expression network analysis (WGCNA) revealed key genes for specific time points of the seed development and quantitative RT-PCR confirmed the expression changes of key genes involved in the seed development and oil accumulation in P. pedunculata seeds. This is the first time to study the transcriptome of P. pedunculata seeds and our study will provide a valuable resource for future studies related to P. pedunculata. The output of this study will improve our understanding towards the seed development and provides the molecular basis of oil accumulation in plants.

Results

Dynamic changes of oil content in the developing seeds of Prunus pedunculata

The seeds of P. pedunculata were obtained from three plants at 10, 18, 24, 31, 39, 45, 59 and 73 DAF. Then,
using the Gas Chromatograph analysis we evaluated the oil content (Fig. 1a), including oil, palmitic acid, palmitoleic acid, stearic acid, oleic acid, linoleic acid and linolenic acid. It is notable that oleic acid was the major type of oil content and was gradually increased during the seed development from 32.724% (10DAF) to 72.060% (73DAF). The composition of linoleic acid and oil peaked at 31 (37.744%) and 59 (52.62%) DAF, respectively. Whereas the oil content remained ~50% after 59DAF. The other oil contents including palmitic acid, palmitoleic acid, stearic acid and linolenic acid were decreased during the seed development. This information provides a basis of understanding the oil content in the developing seeds of *P. pedunculata*.

**Transcriptome sequencing and de novo assembly**
To study the gene changes and explore the molecular mechanisms in the developing seeds of *P. pedunculata*, we employed the transcriptome sequencing for all the samples mentioned above and three biological replicates were used. Initially, 988.795 million high quality reads were produced for the 24 samples (average = 41.20 million reads) by the transcriptome sequencing after data cleaning (Table 1). Then, we randomly selected one

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**Fig. 1** Oil content in the developing seeds of *P. pedunculata* and overview of the assembled transcriptome. a) Proportions of different oil types in the developing seeds of *P. pedunculata* at 10, 18, 24, 31, 39, 45, 59 and 73 DAF. Error bars showing the standard variance are too small to be seen in the plot. b) Evaluation of the assembled transcriptome using BUSCO. S: complete and single-copy BUSCOs; D: complete and duplicated BUSCOs; F: fragmented BUSCOs; M: missing BUSCOs. c) Length distribution of the assembled genes for *P. pedunculata* seeds. d) Number of genes aligned to different species in NR mapping results. e) GO annotation for the assembled genes.
sample from the three replicates for de novo analysis. TRINITY assembled a total of 402,741 transcripts derived from 326,271 genes (Table 1). The GC content, N10, N20 and N50 of the assembled transcriptome were calculated as 40.55%, 2333, 1542, 552, respectively (Table 1). The total bases assembled for the developing seeds of *P. pedunculata* was 205.66 M and CD-HIT identified 198,113 clusters among the assembled transcriptome (Table 1) [18]. Next, we selected the longest transcripts as the unique gene dataset and evaluated the completeness of the assembled unigenes using BUSCO [19]. Figure 1b showed that the assembled transcriptome had more similarity to eukaryote sequences, compared to archaea and bacteria sequences. We identified 219 (85.9%) complete BUSCOs out of the 255 total BUSCOs, including 189 (74.1%) complete and single-copy and 30 (11.8%) complete duplicated BUSCOs. Length distribution of the assembled transcriptome showed that 93.83% of the assembled unigenes were in the length 200 ~ 1000 nt and 533 (0.16%) unigenes were longer than 4000 nt (Fig. 1c).

### Transcriptome annotation

Next, we annotated the assembled seed transcriptome (326,271 unigenes) of *P. pedunculata* using multiple tools and databases. First, the transcriptome was aligned against the NCBI non-redundant (NR) and we found that 195,342 unigenes were involved in the biosynthesis of secondary metabolites (ko01110) and “plant hormone signal transduction” (ko04075), respectively.

Next, we identified DEGs in the developing seeds of *P. pedunculata*. Among the assembled transcriptome RNAMMER predicted 16 unigenes encoding ribosomal RNAs. Next, using TransDecoder we predicted 23,229 likely proteins produced by the assembled transcriptome. We found 18,002 (77.58%) and 16,120 (69.40%) of the likely proteins aligned to the UniProt/SwissProt and Pfam databases, respectively. From the likely proteins, SignalP and TmHMM identified 1917 signal peptides and 5512 transmembrane related proteins, respectively.

### Gene expression profiles and differential expression analysis

We next aligned the clean reads to the assembled unigenes using Bowtie2 and profiled the gene expression for all samples using the RSEM method [20, 21]. We used FPKM (fragments per million reads per kilo base mapped) method to normalize the gene expression and filtered lowly expressed genes (average FPKM < 5) in each sample. As a result, for the developing seeds we identified a total of 91,362 genes, of which 51,363, 50,288 and 52,372 distributed in 10DAF, 18DAF, 24DAF, 31DAF, 39DAF, 50,288 and 50,288 and 52,372 respectively (Fig. 2a). We next analyzed the correlation between samples using the gene expression profiles. Figure 2b revealed that the seed development of *P. pedunculata* can be divided into three stages – early (10 ~ 24 DAF), middle (31 ~ 45 DAF) and late (59 ~ 73 DAF). Interestingly, highly expressed genes (top 10) of the *P. pedunculata* developing seeds confirmed these three development stages (Fig. 2c). Among the highly expressed genes, we found genes encoding proline-rich cell wall proteins specific to early developing seeds (10DAF and 18 DA F), genes encoding YSL9-like proteins specific to early developing seeds (10 ~ 24 DAF), and genes encoding legumin proteins specific to late developing seeds (31 ~ 73 DAF).

Next, we identified DEGs in the developing seeds of *P. pedunculata*. Using 10DAF as control, the numbers of DEGs identified in other samples (Additional file 1) can be seen in Fig. 2d. It is interesting that the numbers of DEGs were increased during the seed development, from 4406 DEGs in 18DAF to 27,623 in 73DAF. Notably, comparison of DEGs in the developing seeds of *P. pedunculata* revealed that 110 up-regulated (Fig. 2e) and 314 down-regulated (Fig. 2f) genes were shared by all

### Table 1 Overview of the assembled transcriptome of *P. pedunculata*

| Category                  | Value             |
|---------------------------|-------------------|
| High quality reads        | 988,795,320       |
| Average reads (high quality) | 41,199,805      |
| TRINITY transcripts       | 402,741           |
| TRINITY genes             | 326,271           |
| CD-HIT clusters           | 193,113           |
| GC (%)                    | 40.55             |
| N10                       | 2333              |
| N20                       | 1542              |
| N50                       | 552               |
| Total assembled bases     | 205,656,558       |

We also identified 65,847 and 64,407 unigenes involved in the pathways of “metabolic pathways” (ko01100) and “plant hormone signal transduction” (ko04075), respectively.

Then, Trinotate was used to further annotate the assembled developing seed transcriptome of *P. pedunculata*. Among the assembled transcriptome RNAMMER predicted 16 unigenes encoding ribosomal RNAs. Next, using TransDecoder we predicted 23,229 likely proteins produced by the assembled transcriptome. We found 18,002 (77.58%) and 16,120 (69.40%) of the likely proteins aligned to the UniProt/SwissProt and Pfam databases, respectively. From the likely proteins, SignalP and TmHMM identified 1917 signal peptides and 5512 transmembrane related proteins, respectively.
the time points compared to 10DAF. Next, we analyzed
the DEGs in the developing seeds compared to the pre-
vious timepoint. Figure 2g showed that the most DEGs
were found in 24DAF vs 18DAF, followed by 73DAF vs
59DAF and 31DAF vs 24DAF. Comparison of DEGs
identified in the developing seeds relative to the previous
timepoint also confirmed that most DEGs identified in
these time points and that no DEGs were identified to
be up-regulated (Fig. 2h) or down-regulated (Fig. 2i)
during the seed development.

Fatty acid related genes
Based on the gene annotation for the developing seeds
transcriptome of *P. pedunculata*, we identified 1246, 51,
9 and 2490 genes related to the fatty acid biosynthesis,
lipid biosynthesis, oil body and TFs, respectively (Table 2).
During the seed development, we identified
753, 33, 8 and 645 DEGs related to the fatty acid biosyn-
thesis, lipid biosynthesis, oil body and TFs, respectively
(Table 2).

Fatty acid related pathways
In the seed development of *P. pedunculata* we identified
753 DEGs (Additional file 2) related to 24 fatty acid
pathways/biological processes, including fatty acid met-
abolism (ko01212), biosynthesis of unsaturated fatty
acids (ko01040), fatty acid biosynthesis (ko00061) and
fatty acid transport (GO:0015908). The numbers of
DEGs identified for each pathway/biological process can be found in Additional file 3 and we used a heat map (Fig. 3a) to show the expression levels of these 753 genes during the seed development. Interestingly, the highly expressed fatty acid related genes varied during the seed development. Thus, we examined the expression of DEGs encoding FAD (fatty acid desaturase), SAD6 (stearoyl-[acyl-carrier-protein] 9-desaturase 6), ACP (acyl carrier protein 1), ACCA (acyetyl-coenzyme A carboxylase carboxyl transferase subunit alpha), LACS (long chain acyl-CoA synthetase), ACAT1 (acetyl-CoA acetyltransferase, cytosolic 1), DGD1 (digalactosyldiacylglycerol synthase 1), GPAT (glycerol-3-phosphate acyltransferase), AGPAT (1-acyl-sn-glycerol-3-phosphate acyltransferase 1), AHGI (Probable protein phosphatase 2C 75) and UBC (ubiquitin-conjugating enzyme) during the seed development. According to their expression patterns during the seed development, these genes can be divided into three groups. The first group of DEGs consisted of FAD, SAD6, ACCA and ACP, which peaked at the middle stage of the seed development of P. pedunculata (Fig. 3b, c). Interestingly, we found that genes encoding omega-6 FAD peaked at 39DAF and that the omega-3 FAD gene peaked at 59DAF (Fig. 3c). The top two highly expressed genes encoding omega-6 FAD were TR56947|c2_g3 and TR56947|c2_g12 (right panel of Fig. 3b). The SAD6 was shown to be detectable between 31DAF and 39DAF (Fig. 3c) and the ACP (TR40569|c0_g1) peaked at 31DAF (Fig. 3c). The second group included GPAT, DGD1 and UBC, which were highly expressed at both early and late stages of the seed development (Fig. 3d). Interestingly, we found that LACS2 were highly expressed at early stage while other LACS (e.g., LACS6, LACS7 and LACS8) were overexpressed at late stage of the seed development. Like LACS genes, UBC1/UBC6 and UBC28 were overexpressed at early and late stages, respectively (Fig. 3d). The third group include AGPAT and ACAT1 genes, which were highly expressed at late stage of the seed development (Fig. 3e). The expression of AGPAT and ACAT1 started to be increased at 45DAF and 73DAF, respectively.

**Lipid biosynthesis**

Among the 55 genes involved in the lipid biosynthesis, 33 genes were differentially expressed during the seed development of P. pedunculata (Additional file 4). Interestingly, their expression patterns during the seed development were similar to G3P, GPAT, LACS and UBC (Fig. 4a). We found that RINos (inositol-3-phosphate synthase) were highly expressed at the early stage and gene encoding TF-B3 domain-containing protein peaked at 45DAF (Fig. 4a). While another two groups of DEGs were highly expressed at the late stage of seed development, such as the O-acyltransferase WSD1-like genes, which peaked at 59DAF, and the EGY2s, which peaked at 73DAF.

**Oil body biosynthesis**

There were nine genes involved in the seed oil body biosynthesis process (GO:0010344) in the seeds of P. pedunculata (Table 2) and we identified 8 of them differentially expressed during the seed development (Additional file 5). Interestingly, their expression was found to be increased after 45DAF and peaked at 59DAF (Fig. 4b). Further, we found that the DEGs encoding LEC2 TFs and TF-B3 domain containing protein were highly expressed at 59DAF and that DEGs encoding oleosins were highly expressed at 73DAF.

**Transcription factors**

Next, we would like to know the expression changes of TF genes and their relationship with the oil content during the seed development. As shown in Table 2, a total of 2490 TFs were annotated for P. pedunculata seeds and 645 were differentially expressed during the seed development (Additional file 1). Among them, 126 AP2-like/ER (ethylene-responsive), 12 AP2, 8 GATA, 26 MADS-box, 42 MYB, 20 NAC, 13 TCP, 15 TGA, 50 WRKY, 78 bHLH, 21 heat stress and 59 nuclear Y subunit TFs were identified (Additional file 1). Notably, most AP2s were highly expressed at the early stage while

| Comparison       | Fatty acid | Lipid | Oil body | TFs |
|------------------|------------|-------|----------|-----|
| Total gene       | 1246       | 51    | 9        | 2490|
| Total DEG        | 753        | 33    | 8        | 645 |
| Compared to 10DAF: |            |       |          |     |
| 18DAF_vs_10DAF   | 15:51      | 2.0   | 0.0      | 20:43 |
| 24DAF_vs_10DAF   | 85:132     | 1.2   | 1.0      | 107:86 |
| 31DAF_vs_10DAF   | 136:176    | 6.2   | 3.0      | 132:121 |
| 39DAF_vs_10DAF   | 168:190    | 5.7   | 3.0      | 140:124 |
| 45DAF_vs_10DAF   | 169:185    | 10.5  | 4.0      | 145:135 |
| 59DAF_vs_10DAF   | 163:205    | 14.8  | 4.0      | 183:92  |
| 73DAF_vs_10DAF   | 175:268    | 12.8  | 4.0      | 207:162 |

Numbers at the left and right sides of the colon represent the up-regulated and down-regulated genes.
Fig. 3 DEGs related to fatty acid pathways. a Hierarchical clustering of the fatty acid pathway associated DEGs identified in the developing seeds of *P. pedunculata*. b Heat map for differentially expressed FAD genes in the samples. c Expression levels of SAD, ACP and AHG1 genes in the developing seeds. d Expression of GPAT, ACCA, DGD1, LACSS, and UBC genes in the developing seeds. e Expression patterns of AGPAT and ACAT1 genes in the developing seeds.
the TGA TFs were highly expressed at the late stages of the seed development (Fig. 4c). We also found that the ER TF subtypes may function at different stages of the seed development (Fig. 4c). For example, AP2/EREBP, CRF2-like, RAP2, ER may function in the initial stage; and AP2-like and ABR1-like, ER function at the late stage of the seed development. Like ER, NAC, WRKY were overexpressed at early and late stages (Fig. 4c). In addition, some TF subtypes were found to function during the whole process of the seed development, such as MYB, TCP, GATA, bHLH, heat stress, nuclear TF Y subunit and MADS-box (Fig. 4c). It is notable that MADS-box – MADS23 and MADS3 peaked at 24DAF while DAM5 for dormancy associated MADS-box peaked at 59DAF and 73DAF (Fig. 4c). WRKY20 and WRKY49 were found to be overexpressed at 24DAF while genes encoding WRKY1, WRKY2 and WRKY21 were highly expressed at late stage of the seed development (Fig. 4c).

**Weighted gene co-expression network analysis**

To further investigate the association between DEGs and the oil content in the seeds of *P. pedunculata*, we performed the weighted gene co-expression network analysis (WGCNA). As a result, we identified 450, 133, 6,
Fig. 5 WGCNA and qRT-PCR. a WGCNA analysis identified co-expressed genes associated with the fatty acid biosynthesis. b qRT-PCR validation for six genes. For each gene the upper and lower panel of plots showed the expression levels (normalized to 10DAF) detected by transcriptome sequencing and qRT-PCR, respectively.
were associated with the late stage of seed development and the oil content in the seeds of *P. pedunculata*. Among the blue module genes, 420 were co-expressed and the oil content in the seeds of *P. pedunculata* increased with the time (DAF). Heat maps represent the expression levels (scaled values) of genes in the developing seeds of *P. pedunculata*. DAG: 1,2-Diacylglycerol; PA: phosphatidic acid; LPA: lysophosphatidic acid; G-3-P: glycerol-3-phosphate; PC: phosphatidylcholine; LPC: lysophosphatidylcholine, and glycerol; MAG: monacylglycerol; PDH: pyruvate dehydrogenase complex; ACCA: acetyl-coenzyme A carboxylase carboxyl transferase subunit alpha; ACP: acyl carrier protein; MAT: malonyl-CoA-acyl carrier protein transacylase; KAS: ketoacyl-ACP synthase; EAR: enoyl-ACP reductase; KAR: ketoacyl-ACP reductase; HAD: hydroxyacyl-ACP dehydrase; FAD: fatty acid desaturase; SAD: 18:0-ACP desaturase; FATB: acyl-ACP thioesterase B; FATA: acyl-ACP thioesterase A; DGAT: diacylglycerol acyltransferase; PAH: phosphatic acid phosphohydrolase; LPAT: acyl-CoA acyltransferase; GPAT: glycerol-3-phosphate acyltransferase; CPT: diacylglycerol cholinephosphotransferase; MGAT: monoglyceride acyltransferase; SDP: sugary-dependent protein; AGK: aglycon kinase; MGL: acylglycerol lipase; GLPK: glycerol kinase; PLA: phospholipase; LPCAT: lysophosphatidylcholine acyltransferase.
and various TFs (e.g., AP2-like ER, TGA). Interestingly, the other fatty acid types, including palmitic acid, palmitoleic acid, stearic acid, oleic acid, linoleic acid and linolenic acid, were found to be associated with the turquoise module genes (Additional file 6), which were significant at the early stage of the seed development. In addition, we found that the omega-3 FAD gene was co-expressed with the oil at 73DAF and that SAD6 genes were significantly correlated with the omega-6 FAD genes during the middle seed development stage of *P. pedunculata*.

**Fatty acid biosynthesis pathway for *P. pedunculata* seed**

To propose a proper pathway involved in the fatty acid biosynthesis in the developing seeds of *P. pedunculata*, we first highlighted the DEGs in the general fatty acid biosynthesis pathway based on the KEGG pathway annotation (Fig. 6a). It showed that some key genes of the fatty acid biosynthesis were identified in this study, including 6.4.1.2 (acetyl-CoA carboxylase, ACACA), 3.1.2.14 (fatty acyl-ACP thioesterase B, FATB), 3.1.2.21 (medium-chain acyl-[acyl-carrier-protein] hydrolase, MCH), 1.14.19.2 (acyl-[acyl-carrier-protein] desaturase, FAB2), 6.2.1.3 (long-chain acyl-CoA synthetase, ACSL), FabD, FabF, FabH, FabI and FabZ. Their products are either substrates, intermediates or important enzymes. Next, we proposed the working model of fatty acid synthesis in plastid and endoplasmic reticulum (Fig. 6b). In addition to the DEGs described above, we found extra important genes related to the fatty acid biosynthesis. For example, PDH (pyruvate dehydrogenase complex), involved in the overall conversion of pyruvate to acetyl-CoA, was found to be highly expressed in the seeds of 24DAF and 31DAF (Additional file 1). Some other genes from this model were also found to be differentially expressed in the developing seeds of *P. pedunculata*, such as MAT (malonyl-CoA-acyl carrier protein transacylase), FATB, DGAT, PAH (phosphatidic acid phosphohydrolase), DCP (diacylglycerol cholinephosphotransferase), PLA (phospholipase), AGK (acylglycerol kinase) and GLPK (glycerol kinase). However, some genes like KASI (ketoacyl-ACP synthase I), KASII (ketoacyl-ACP synthase II), KASIII (ketoacyl-ACP synthase III), WRI, LPCAT (lysophosphatidylcholine acyltransferase), SDP (sugar-dependent protein) and LPAT (acyl-CoA:acylglycerol-3-phosphate acyltransferase) were not detected in the developing seeds of *P. pedunculata*. The reason for their absence requires further experiments to be explored.

**qRT-PCR**

Then, we performed quantitative real-time PCR to validate the gene expression changes during the seed development of *P. pedunculata*. A total of six genes (TR52836|c2_g6, TR61055|c0_g11, TR61099|c1_g5, TR61813|c2_g5, TR63042|c1_g1, TR64906|c6_g12) were randomly selected and the 18S rRNA was used as the internal control. The primers can be accessed in the Additional file 7 and nine reactions were performed for every gene at each time point of the seed development. Relative normalized expression (RNE) was used to show the expression of all genes in the samples (relative to 10DAF). As a result, we found that the expression patterns of all six genes were agreed by both RNA-Seq and qRT-PCR (Fig. 5b). It is notable that the high expression of TR61055|c0_g11 (transcription factor TGA2), TR63042|c1_g1 (MYB transcription factor 3) and TR61813|c2_g5 (long chain acyl-CoA synthetase 6, LACS6) were confirmed by qRT-PCR at the late seed development stage and the high expression of TR61099|c1_g5 (fatty acyl-ACP thioesterase B, FATB) was validated in the early seed development stage (Fig. 5b). From the sequencing and qRT-PCR results we found that TR64906|c6_g12 (B3 domain-containing transcription factor LEC2) peaked at 59DAF. TR52836|c2_g6 (stearyl-[acyl-carrier-protein] 9-desaturase 6, SAD6) was found to be highly expressed at 31DAF by the sequencing and at 39DAF by qRT-PCR (Fig. 5b) and both time points are defined as in the middle seed development stage. The high agreement of gene expression patterns by transcriptome sequencing and qRT-PCR revealed that the genes identified in this study might be functional in the development and the oil content of the *P. pedunculata* seeds.

**Discussions**

Seed oil content is a new important trait of *P. pedunculata*, however, little is known about the molecular basis of oil accumulation and genes involved in the oil synthesis in *P. pedunculata* seeds, probably due to the missing of its genome sequence. Here, we assembled the transcriptome for *P. pedunculata* seeds for the first time and studied the dynamic gene profiles of the seeds at eight developing points. We examined the content of different fatty acid types, including oil, palmitic acid, palmitoleic acid, stearic acid, oleic acid, linoleic acid and linolenic acid, in the developing seeds of *P. pedunculata* (Fig. 1a). The three major fatty acid types were oil, oleic acid and linoleic acid in the mature seeds of *P. pedunculata*. Interestingly, oleic acid was shown to be the highest type in the *P. pedunculata* seeds, and the proportion is similar to camellia and olive oil [22, 23]. The decrease of palmitic acid, palmitoleic acid, stearic acid and linolenic acid during the seed development might link with the high expression of LACS2 at the early developing stage of *P. pedunculata* seeds (Fig. 1a, Fig. 3d). In *Arabidopsis* LACS2 enzyme preferers using palmitic acid and...
palmitoleic acid as substrates to form the cutin or cuticular waxes [24].

According to the gene expression profiles, the seed development of *P. pedunculata* can be divided into three stages (Fig. 2b, c). We observed that genes encoding *GPAT*, *DGD1*, *LACS2*, *UBC*, *RINO* and multiple TFs (e.g., *AP2/EREBP ER*, *CRF2-like*, *RAP2*, *AP2*, *NAC*, *MYB*, *MADS-box*, *WRKY*) were highly expressed at the early stage (10 ~ 24DAF) of the seed development (Fig. 3d, Fig. 4a, c). *GPAT* exhibits sn-1 acyltransferase activity with high specificity of acyl-CoAs, has the potential of triggering the biosynthesis of storage lipid and plays a pivotal role in the Kennedy pathway, the de novo assembly of TAG from G-3-P and acyl-CoAs and the glycerolipid biosynthesis [25]. The G-3-P is acylated by *GPAT* and *LPAT* to yield the phosphatic acid, which is then hydrolyzed to form diacylglycerol, the final substrate for TAG via DGAT (diacylglycerol acyltransferase) [17]. The UBC enzymes are essential for the fatty acid β-oxidation during the early seedling development of oil-seed plants [26]. The high expression of these genes at the early development stage might contribute to the accumulation for the fatty acid biosynthesis in subsequent processes.

At the middle stage of seed development (31 ~ 45DAF), we identified *ω6-FAD*, *SAD*, *ACP*, *ACCA*, *AHG1* and some TF (e.g., *TCP*, heat stress, *bHLH*, nuclear TF Y subunit) genes highly expressed (Fig. 3, Fig. 4). *ACCA* is the first enzyme involved in the pathway of fatty acid synthesis in plastid and regulates the flux of carbon into fatty acids [4]. Whereas the assembly of fatty acids usually happens on the ACP through a cycle of 4 reactions of elongating the acyl chain by 2 carbons each time [6]. The *FAD* genes have been described to correlate the seed development and the fatty acid biosynthesis in some oil plants, including *Zanthoxylum bungeanum* [11], *Camelina sativa* [12], *Linum usitatissimum* [27], *Eucommia ulmoides* [13], *Glycine max* [14] and *Paeonia section* [16]. *FAD2* and *FAD3* in the ER (endoplasmic reticulum) catalyze the desaturation of fatty acids attached to phosphatidylcholine (PC) from PC-18:1 to PC-18:2 (*FAD2*) and from PC-18:2 to PC-18:3 (*FAD3*) [28–30]. *FAD6* has the potential of introducing the double bond in the biosynthesis of 16:3 and 18:3 fatty acids, which are important constituents of plant membrane, and functions on fatty acids esterified to galactolipids, sulfolipids and phosphatidylglycerol through ferredoxin [31]. It is reported that stearic acid is converted into oleic acid (highest content in the *P. pedunculata* seeds) by SAD, which is one of the regulators for the proportion of fatty acids in seeds [4]. While AHG is a negative regulator of abscisic acid responses during the seed germination [32]. The high expression of key regulatory genes during the middle stage of seed development may indicate the oil storage and the determination of the contents of fatty acids.

Next, we would like to discuss highly expressed genes identified in the late stage (59 ~ 73DAF) of the seed development, such as *LACS6*, *DGD1*, *ACAT1*, *AGPAT*, *WSD1*, *EGY2* and oleosin (Fig. 3, Fig. 4). Using palmitate, palmitoleate, oleate and linoleate as substrates, *LACS6* has the potential of activating the long chain fatty acids for both synthesis of cellular lipid and degradation through beta-oxidation [33, 34]. *DGD1* is specific for α-glycosidic linkages and involved in the synthesis of diacylglycerol galactolipids which are specifically localized in the thylakoid membranes [35, 36]. The overexpression of *ACAT1*, a CoA biosynthetic enzyme, has been shown to lead the increase of oil content in Arabidopsis [37]. *AGPAT* is a plastidial enzyme of the prokaryotic glycerol-3-phosphate pathway, which converts the lysophosphatic acid into phosphatic acid by incorporating an acyl moiety at position sn-2 [38]. The α-acyltransferase *WSD1*, the major wax ester synthase, has been reported to be involved in the drought tolerance in plants [39, 40]. In Camelina seeds the expression of *WSD1* was up-regulated during the early seed development and correlated with the oil content of the seeds [12]. Oleosin are the major proteins involved in the oil body biogenesis and the stabilization of TAG/cytosol oil body interface [41]. The expression of oleosin genes has been shown to be up-regulated during the seed development in plants, such as *Camelina sativa* [12], soybean [14], *Paeonia section* [15] and *Perilla frutescens* [17]. Overall, these genes have been shown to participate the fatty acid biosynthesis, however, their functions in the oil accumulation of *P. pedunculata* seeds are still not clear. Together with these studies, the high expression of these genes supports them to play a key role during the seed development and fatty acid biosynthesis in *P. pedunculata*.

The fatty acid biosynthesis is a complicated process that involves many enzymes and molecules. It is the fundamental to the production of membranes and lipids in the plastids of plants [13]. We proposed a proper working model for the genes involved in the fatty acid biosynthesis for *P. pedunculata* (Fig. 6b). Some key genes from the process were identified with differential expression in the developing seeds of *P. pedunculata*. However, we missed some known important regulators that have been reported to participate in the fatty acid biosynthesis, such as *KASI ~ III* [13], *WR11* [9] and *LPAT* [17]. There might be some explanations for their absence, including 1) the de novo assembly method or the annotation method missed these genes; 2) they were not detected by the transcriptome sequencing; and 3) they were not expressed in the seeds of *P. pedunculata*. Future studies need to be performed to complete the working model of fatty acid biosynthesis in *P. pedunculata*. In addition, we
identified some TFs that might be involved in the fatty acid biosynthesis of *P. pedunculata*, however, the regulation mechanism during the oil accumulation is still not clear. Future experiments are required to investigate their functions in the fatty acid biosynthesis for *P. pedunculata*.

**Conclusions**

In conclusion, we assembled the first transcriptome for the developing seeds (eight time points) of *P. pedunculata*. In total, 402,741 transcripts of 326,271 unigenes were reported with the GC ratio of 40.55%. Annotation of the transcriptome showed 109,850 and 121,897 unigenes aligned to the GO and KEGG pathway databases, respectively. We also predicted 23,229 likely proteins produced by the *P. pedunculata* seed transcriptome, of which 1246, 51, 9 and 2.490 related to fatty acid biosynthesis, lipid biosynthesis, oil body and TFs, respectively. We profiled 91,362 genes expressed more than 5 FPKM in the developing seeds of *P. pedunculata* and identified 48,788 DEGs. Compared to 10DAF, the numbers of DEGs were shown to be increased during the seed development, from 4406 in 18DAF to 27,623 in 73DAF. We found that 753, 33, 8 and 645 DEGs related to the fatty acid biosynthesis, lipid biosynthesis, oil body and TFs, respectively. *GPAT*, *DGDL*, *LACS*, *UBC* and *RINO* were highly expressed at the early seed development (10 ~ 24DAF), *ω-6-FAD*, *SAD*, *ACP*, *ACCA* and *AHG1* were highly expressed at the middle seed development (31 ~ 45DAF) while *LACS*, *DGDL*, *ACAT*, *AGPAT*, *WSD1*, *EGY2* and oleosin genes were abundant at the late seed development. The genes expression changes were confirmed by qRT-PCR. This is the first time to study the transcriptome of *P. pedunculata* developing seeds and our findings will provide a valuable resource for future studies. More importantly, it will improve our understanding of the oil accumulation for *P. pedunculata*.

**Methods**

**Plant material**

We selected three six-year-old plants of *Prunus pedunculata* Pall which were planted in the experimental fields of Inner Mongolia Breeding Center, China. The original seeds were obtained from the wild place of Baotou, China (110.066562,41.039184) and maintained in the Inner Mongolia Seed Museum as Mengbian-1. The days after flowering (DAF) of fruits were marked with tags to track the seed development. From 10th April of 2019, we collected the fruits of *Prunus pedunculata* at 10, 18, 24, 31, 39, 45, 59 and 73 DAF with three biological replicates (*n* = 3). Every development stage we collected a total of ~ 100 seeds at each timepoint and peeled off the peel and stone. Then, the nucleolus was flash frozen in liquid nitrogen and stored at −80 °C until further use. As the ovules at 0 DAF were hard to collect, we used seeds at 10 DAF as the control.

**Quantitation of oil content**

The oil content of *P. pedunculata* seeds was determined using the Gas Chromatograph (GC, Agilent 6890 N) according to the manufacturer’s protocol, as described [14]. The CP-Sil 88 (Agilent Technology) and nitrogen gas were used as the GC column and carrier gas, respectively. The initial temperature was set to 180 °C and raised to 200 °C at the rate 6 °C/min. After 3 min, the temperature was then raised to 240 at the rate 10 °C/min. The samples were ground with grinder at low temperature and then dried in the vacuum freezing dryer. Then, powder samples (100 mg) were diluted in 1% heptadecanoic acid (with internal standard) in 2 mL centrifuge tubes. After 1 mL methylation agent (2.5%, v/v, H2SO4 in CH3OH) was added to each tube, the samples were water bathed for the methyl esterification at 85 °C for 1 h. We collected the extractions, centrifuged them and retained the supernatants, followed by adding 600 µL NaCl (0.9%, w/v) and 350 µL n-hexane later. Then, the mixture was centrifuged at 4000 RPM for 10 min and the organic phase was air-dried. After 500 µL ethyl acetate was added to the dried methyl esterification samples, they were subjected to the GC for oil content analysis. Compared to the standard, the fatty acids were examined qualitatively and quantitatively by the peak area method. The absolute content of fatty acids was determined by the methyl ester standard. This experiment was replicated three times and average values were calculated for each oil type.

**Total RNA extraction, library construction and sequencing**

Total RNA was extracted from the nucleolus using the TRIzol reagent (Invitrogen), as previously described [42]. After the quality and quantity of the total RNA were examined using the Agilent 2100 Bioanalyzer, equal amount of total RNA (1 µg) was used for the library construction at BGI-Shenzhen. In brief, poly(A) mRNAs were enriched and pulled down by the magnetic oligo (dT) beads and fragmented into small pieces (~ 200 bp). Then, random hexamer (N6) primers were used to construct the double strand cDNA libraries. After the end repair by phosphate at 5′-end and sticky“A” at 3′-end, the cDNA libraries were ligated with sequencing primers to build the final RNA-Seq libraries. Next, the final libraries were sequenced on the BGISEQ-500 RS platform with paired-end 150 strategy.

**De novo assembly**

We used Trim_galore to remove low quality reads, reads with adapters and contamination reads from the raw reads. FASTQC (http://www.bioinformatics.babraham.org)
ac.uk/projects/fastqc/) was used to evaluate the clean reads for quality control. Then, clean reads of each time point were subjected to the TRINITY software for de novo assembly, as described previously [43]. CD-HIT was used to evaluate the clusters of the transcripts [18]. Then, we selected the longest transcripts (used as unigenes) using the program provided by Trinity and extracted the likely proteins using TransDecoder. To evaluate the completeness of the assembled unigenes, we employed BUSCO to map the unigenes to three datasets, evaluate the completeness of the assembled unigenes, we extracted the likely proteins using TransDecoder. To gene) using the program provided by Trinity and ex-

Transcription annotation

We first annotated the assembled transcriptome using Trinitate, according to the protocol [44]. In brief, the unigenes and likely proteins were aligned against the UniProt/Sprot using BLASTX and BLASTP, respectively, with the e-value <1e-3, RNAmer (v1.2) [45], HMMER [46], SignaLP (v4.1) [47] and TMHMM Server (v.2.0) [48] were used to predict the rRNA transcripts, protein domains, signal peptides and transmembrane regions in the assembled unigenes and likely proteins. Then, all the annotations were subjected to Trinotate to generate a report.

Next, we aligned the assembled unigenes to NCBI non-redundant (NR), UniProt and Kyoto Encyclopedia of Genes and Genomes (KEGG) databases to obtain the Gene Ontology (GO) and pathway annotation, as previously described [43]. After the matched unigenes were selected (e-value <1e-5), BLAST2GO was used to retrieve associated GO items describing biological process (BP), cellular component (CC) and molecular function (MF). Unigenes with enzyme commission number (EC) domains, signal peptides and transmembrane regions in the assembled unigenes and likely proteins. Then, the CV < 0.5 and false discovery rate (FDR) < 0.1 [50].

Functional analysis

We performed functional analysis for the differentially expressed genes using the GO and KEGG pathway annotation results. To identify significantly enriched GO terms and pathways, we first calculated the p-value using the Fisher’s Exact Test to present the enrichment significance and calculated the corrected p-value (shown as q-value) using the value package in R platform. Significantly enriched GO terms and KEGG pathways were selected as follows: q-value < 0.05 and non-plant associated terms/pathways were filtered.

Gene expression profiles and differential expression analysis

We used Bowtie2 [20, 50] and RSEM (RNA-Seq by Expectation-Maximization) [21] to align the clean reads to the assembled unigenes and profile gene expression, respectively, for each sample. FPKM (fragments per million reads per kilo base mapped) method was used for normalization and average FPKM of each gene were calculated for each time point. After lowly expressed genes (average FPKM < 5) were filtered, we performed differential expression analysis using edgeR with following cut-offs: log2 fold change (log2FC) > 1 or < -1, coefficient of variation (CV) < 0.7, p-value < 0.5 and false discovery rate (FDR) < 0.1 [50].

qRT-PCR

We performed quantitative real-time PCR (qRT-PCR) to validate the gene expression changes during the seed development of Prunus pedunculata. We randomly selected six genes (TR52836|c2_g6, TR61055|c0_g11, TR61099|c1_g5, TR61813|c2_g5, TR63042|c1_g1, TR64906|c6_g12) and used the 18S rRNA as the internal control. Forward and reverse primers were predicted for the candidate genes and control using the Beacon Designer 7.9. After the total RNA was extracted from the seeds at all eight time points (as described above), an aliquot (2 mg) of the total RNA was used for the first-strand cDNA synthesis using the TRUEscript First-Strand cDNA Synthesis kit (Aidlab Biotech, Beijing, China). Then, the cDNA (1 μL) was used to build the qRT-PCR reaction mixture (10 μL) together with 2 × SYBR Green Supermix (5 μL, DBI, China), forward primer (0.5 μL), reverse primer (0.5 μL) and ddH2O (3 μL). The qPCR mixture was run on the qTOWER 2.2 qRT-PCR Thermal Cyclers (Analytik Jena, Germany) following the protocols. Then, Ct value of each gene in each sample was obtained and ΔCt was calculated. The 10DAF samples were used as control and the expression levels of all genes in the other samples were calculated present using the relative normalized expression: RNE = 2−ΔΔCt. Three reactions were performed for each gene in every single replicate, thus we obtained nine replicates (n = 9) for each gene at each timepoint.

Abbreviations

DAF: Days after flowering; GO: Gene Ontology; TF: Transcription factor; FPKM: Fragments per million reads per kilo base mapped; TAG: Triacylglycerol; NR: NCBI non-redundant database; DEG: Differentially expressed gene

Supplementary Information

The online version contains supplementary material available at https://doi.org/10.1186/s12870-021-02921-x.

Additional file 1. Differentially expressed genes in the developing seeds of P. pedunculata.

Additional file 2. Differentially expressed genes related to fatty acid in the developing seeds of P. pedunculata.
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Authors’ contributions

WB and LW conceived and designed the experiments; WB, DA and YB performed the experiments; WB, MC, JC, LW and SZ analyzed the data; DA, ZL and FL performed the validation experiment; WB and DA wrote the manuscript; TW and LW revised the manuscript. All the authors have read and approved the final version of manuscript.

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Availability of data and materials

The raw sequencing data can be accessed from the NCBI Sequence Read Archive (SRA) platform (http://trace.ncbi.nlm.nih.gov/Traces/sra/) under the accession number PRJNA684995.

Declarations

Ethics approval and consent to participate

No specific permits were required for the described field studies or for the collection of the seeds. The location is not privately-owned or protected in any way, and the field studies did not involve endangered or protected species. All the treatments to the seeds of P. pedunculata comply with the national and institutional guidelines and legislations. The seeds can be accessed upon the request to the corresponding author.

Consent for publication

Not applicable.

Competing interests

The authors declare that there is no conflict of interest.

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