De novo transcriptome sequencing and comparative analysis to discover genes related to floral development in *Cymbidium faberi* Rolfe

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**Abstract**

*Cymbidium faberi* is a traditional orchid flower in China that is highly appreciated for its fragrant aroma from its zygomorphic flowers. One bottleneck of the commercial production of *C. faberi* is the long vegetative growth phase of the orchid and the difficulty of the regulation of its flowering time. Moreover, its flower size, shape and color are often targeting traits for orchid breeders. Understanding the molecular mechanisms of floral development in *C. faberi* will ultimately benefit the genetic improvement of this orchid plant. The goal of this study is to identify potential genes and regulatory networks related to the floral development in *C. faberi* by using transcriptome sequencing, de novo assembly and computational analyses. The vegetative and flower buds of *C. faberi* were sampled for such comparisons. The RNA-seq yielded about 189,300 contigs that were assembled into 172,959 unigenes. Furthermore, a total of 13,484 differentially expressed unigenes (DEGs) were identified between the vegetative and flower buds. There were 7683 down-regulated and 5801 up-regulated DEGs in the flower buds compared to those in the vegetative buds, among which 3430 and 6556 DEGs were specifically enriched in the flower or vegetative buds, respectively. A total of 173 DEGs orthologous to known genes associated with the floral organ development, floral symmetry and flowering time were identified, including 12 TCP transcription factors, 34 MADS-box genes and 28 flowering time related genes. Furthermore, expression levels of ten genes potentially involved in floral development and flowering time were verified by quantitative real-time PCR. The identified DEGs will facilitate the functional genetic studies for further understanding the flower development of *C. faberi*.

**Keywords:** *Cymbidium faberi* Rolfe, RNA-seq, Flower development, Flowering, Genes

**Background**

Orchidaceae is one of the largest families in the angiosperms with more than 25,000 species, which displays a great biodiversity resulting from adaptation to diverse habitats (Pridgeon et al. 2005). Genomic information on orchids were mainly focused on *Phalaenopsis* (Su et al. 2011; Cai et al. 2014), *Dendrobium* (Yan et al. 2015; Zhang et al. 2016), *Cymbidium ensifolium* (Li et al. 2013), *Cymbidium sinense* (Zhang et al. 2013). *Cymbidium faberi* Rolfe., common named as “Hui Lan”, is one of the oldest and most popular orchids species cultivated in China, which is highly appreciated mainly because of its beautiful flower posture and fragrant aroma (Wolff, 1999). However, large scale commercial production of *C. faberi* was often hindered due to the long vegetative growth phase (usu. 5–7 years) and difficulties in flowering time control.

Plant flowering involves a transition process from vegetative growth to reproductive development with a series of conserved underlying metabolic or external pheno-typic changes taking place in the shoot apical meristem. In *Arabidopsis thaliana*, there are four major pathways controlling the timing of flowering, including photo-period, vernalization, gibberellin (GA) and autonomous...
pathways (Mouradov et al. 2002). Major genetic elements involved in these pathways have been defined as the key switches to control flowering, such as CONSTANS (CO) in the photoperiodic pathway and FLOWERING LOCUS C (FLC) in the autonomous and vernalization pathways (Mouradov et al. 2002). The FLOWERING LOCUS T (FT) gene activates the expression of a number of flower developmental genes (Mouradov et al. 2002). In the orchid plant *Phalaenopsis aphrodite*, PaFT1 can suppress the delayed flowering caused by SHORT VEGATATIVE PHASE (SVP) and FRIGIDA (FRI) (Jang et al. 2015). The Cymbidium FT orthologous gene was also cloned and ectopic expression of CgFT resulted in early flowering in transgenic Arabidopsis (Huang et al. 2012). Over expression of DnAGL19, a SOC1-1/TM3-like ortholog in Arabidopsis resulted in a slightly accelerated flowering time under normal growth conditions (Liu et al. 2016).

During the transition from vegetative growth to reproductive growth, MADS-box family genes play important roles in regulating floral organ specification, development and evolutionary in higher plants (Weigel and Meyerowitz 1994; Purugganan et al. 1995; Münster et al. 1997). Although conserved flowering pathways and multiple key genes were revealed in model plants, there is limited information of *C. faberi*, the very unique and important plant species featured with a 5–7 year-long vegetative growth phase. Unlike most tropical orchids (e.g. *Phalaenopsis* and *Cattleya*), *C. faberi* flowers are not brightly showy but of strong aromas to attract pollinating insects. It is of strong interest for orchid breeders to improve the appearance of its flower size, color and shape with uncompromised aromas. The flowers of *C. faberi* are zygomorphic consisting of four whorls: The first whorl is comprised of the petal-like sepals, the second whorl is of pistils in the form a highly specialized labellum in the middle, the third whole is of stamens, and fourth whorl is of pistils in the form a highly specialized united gynostemium (Rudall and Bateman 2002). The well-known ABC model clarify that floral development is determined by five kinds of floral organ identity genes in diverse plant groups (Coen and Meyerowitz 1991; Weigel and Meyerowitz 1994). Sepal formation is specified by the expression of A-class function genes. Expression of AB and BC determine the development of petals and stamen formation, respectively. The development of carpel is determined by C-class genes function alone, and D-class genes specify ovules. While class E function redundantly specify petals, stamens, and carpels as well as floral determinacy (Pelaz et al. 2000, 2001; Anusak et al. 2003). The floral morphology of orchid species is unique with gynostemium, labellum and resupination caused by 180° torsion of the pedicel (Rudall and Bateman 2002). ‘The Perianth (P) code’ clarifies the mechanisms of sepal/petal/lip determination in *Oncidium* and *Phalaenopsis* orchids. The competition between different APETALA3/AGAMOUS-LIKE6 (AP3/AGL6) homologues determines the formation of the complex perianth patterns in orchids. The formation of sepal/petal were specified by the higher-order heterotetrameric SP (sepal/petal) complex (OAP3-1/OAGL6-1/OAGL6-1/OP1), whereas the lip formation required the L (lip) complex (OAP3-2/OAGL6-2/OAGL6-2/OP1) exclusively (Hsu et al. 2015). Other MADS-box function genes participating in the sepal and petal development were also isolated from *Dendrobium madame*, *D. crumenatum*, Oncidium Gower Ramsey (Yu et al. 2000, 2002; Hsu and Yang 2002; Hsu et al. 2003; Xu et al. 2006). Genes involved in flower identity and floral organ specification are still unknown in *C. faberi* yet.

Besides petal shape and size, floral symmetry significantly affects the ornamental value of flowers. So far, three transcription factors (TFs) that determine the floral symmetry are identified, including CYCLOIDEA (CYC) from the TCP family (TEOSINTE BRANCHED1/CYCLOIDEA/PROLIFERATING CELL NUCLEAR ANTIGEN FACTOR), DIVARICATA (DIV) and RADIALIS (RAD) from the MYB family (Luo et al. 1996; Doebley et al. 1997; Almeida et al. 1997; Galego and Almeida 2002; Corley et al. 2005; Costa et al. 2005). Of these genes, CYC and its orthologues establish the floral monosymmetry through specifying dorsal flower identity (Luo et al. 1996, 1999; Feng et al. 2006; Wang et al. 2008). Studies on floral symmetry genes mainly focused on species with highly derived morphologies in eudicots, whereas only a few studies are available for monocots (Luo et al. 1996, 1999; Feng et al. 2006; Wang et al. 2008; Yuan et al. 2009; Bartlett and Specht 2011; Preston and Hileman 2012; Hoshino et al. 2014). Orchidaceae is characterized by highly specialized zygomorphic flowers. Studies on orchid flower symmetry is very limited (Paolo et al. 2015). Overall, the detailed molecular mechanism or the genetic elements in the regulation of floral organ specification and development in *C. faberi* remains elusive.

Transcriptome analysis is an useful tool to discriminate differences in transcript abundance among different cultivars, organs and different treatment conditions in model and non-model plants (Cheung et al. 2006; Trick et al. 2009; Li et al. 2013; Hyun et al. 2014; Zhang et al. 2014). In order to excavate genes that might regulate the floral development in *C. faberi*, we used high-throughput Illumina sequencing and bioinformatics analysis to compare the de novo transcriptomes of vegetative and flower buds from *C. faberi*. The vegetative transcriptome can be used to find genes related to the vegetative growth. The floral transcriptome was sufficiently comprehensive for gene discovery and analysis of major metabolic pathways associated
with flower traits. Genes expressed differently in the vegetative buds and flower buds may play important roles in the vegetative growth or floral development. Through transcriptome analysis, large numbers of genes related to floral organ initiation, flower symmetry patterning and flowering were identified in *C. faberi*. These results provide fundamental information for further studies on the molecular mechanism of flower development in *C. faberi*.

**Methods**

**Plant materials**

Mature plants of *C. faberi* with light green flowers, originally introduced from Henan province of China, were grown in the greenhouse at Department of Horticulture, Nanjing Agricultural University (Nanjing, China) under natural light and a controlled temperature of 22–28 °C. The vegetative buds were collected from lateral buds, and the flower buds (0.5–1.0 cm in length) were collected from the peduncle (Fig. 1). Organs from three individual plants were pooled as one sample. The fresh samples were frozen immediately in liquid nitrogen and stored at −80 °C.

**RNA extraction, cDNA library construction and de novo assembly**

Total RNA of flower and vegetative buds were extracted using EASYspin plant RNA rapid extraction kit according to the manufacturer’s protocol (Yuan Ping Hao Biotechnology Co. Ltd, Beijing, China). Then, the concentration was detected by a NanoDrop 2000™ micro-volume spectrophotometer (Thermo Scientific, Waltham, MA, USA) and the quality was tested by gel electrophoresis. cDNA library construction for the flower and vegetative buds and Illumina deep sequencing were performed on the HiSeq™ 2000 platform according to the manufacturer’s instructions at Hangzhou Woosen Biotechnology Co. Ltd. (Hangzhou, Zhejiang, China). The Illumina reads were assembled to obtain the contigs and unigenes using Trinity software and Cap3 after removing the short raw reads and quality inspection by fastQC (Grabherr et al. 2011).

**Functional annotation of unigenes**

The unigenes were annotated with the National Center for Biotechnology Information non-redundant databases (NR), Gene Ontology (GO), Clusters of Orthologous Groups (COG) and Kyoto Encyclopedia of Genes and Genomes (KEGG) databases using BLASTX searches (E-value ≤ 1.0e−5). The Blast2GO software package and WEGO software were employed to compare and determine unigenes’ GO annotations and obtain GO functional classifications for all annotated unigenes (Götz et al. 2008; Ye et al. 2006).

![Fig. 1](image-url) The mature plants and flowers of *C. faberi*: **a** Flowering plants of *C. faberi*; **b** Inflorescence with flower buds (*v*); **c** Vegetative buds (*v*); **d** flowering flowerers; **e** single flower; **f** different parts of a single flower, *s* sepal, *p* petal, *l* labellum, *c* column
Identification of differentially expressed genes (DEGs)

Expression levels of all unigenes were calculated and then compared between the two tissue samples using the fragments per kilobase of transcript per million reads of library method (FPKM). The false discovery rate (FDR) was adopted to determine the threshold of P-values in multiple tests. DEGs were also annotated with GO assignments, COG assignments and KEGG pathways. The criteria FDR < 0.05 was used to identify DEGs and acted as a threshold of significant difference of gene expression in GO terms, COG classification and KEGG annotation.

Quantitative real-time PCR (qRT-PCR) analysis

Total RNA of flower buds and vegetative buds were extracted as above and the first strand cDNA was synthesized using PrimeScript RT reagent Kit With gDNA Eraser (Takara Bio Inc.). All primers used in this study were designed by the Primer 5 software according to the RNA-Seq data. CfGAPDH (JX560732) was selected as an internal reference (Additional file 1: Table S1).

The qRT-PCR analysis was performed on ABI 7500 Real-Time PCR Detection System (Applied Biosystems) using the SYBR® Premix ExTagTM reagent kit (Takara Bio Inc.) according to the manufacturer’s protocol. The PCR reactions were 40 cycles (95 °C for 15 s; 55 °C for 15 s; 72 °C for 20 s) according to the instruction manual. A melting curve was generated to test the specificity of products after the qRT-PCR. The relative expression levels of the selected unigenes were normalized to CfGAPDH gene and calculated using the 2−ΔΔCt method. Data were derived from three independent replicates.

Results

Transcriptome sequencing and de novo assembly

Equal amount of RNAs from the vegetative and flower buds were used to constructed cDNA libraries separately, and then sequenced. A total of 35,511,583 and 32,514,423 raw reads were obtained in the flower and vegetative buds, respectively. After removing the short raw reads and quality inspection by fastQC, the RNA-seq produced 35,510,239 and 32,513,288 clean reads for the flower and vegetative buds, respectively (Table 1). All of these reads were employed for further de novo assembly. And 189,300 contigs with an average length of 755 bp were generated using the Trinity software package. These contigs were assembled into 172,959 unigenes (200 to >10,000 bp) with an average length of 698 bp and an N50 (N50 represents weighted median length of all contigs) of 1340 bp (Table 2). Among them, 59,505 (34.4 %) unigenes were more than 500 bp. The size distribution of the assembled unigenes is shown in Additional file 2: Fig. S1. And this transcriptome shotgun assembly project has been deposited at DDBJ/EMBL/GenBank under the accession of GDHD00000000.

Functional annotation of C. faberi transcriptome

To validate and annotate the assembled unigenes, the 172,959 unigenes generated were subjected to BLASTX search (E-value ≤ 1.0e−5) against public protein databases such as NCBI NR, GO, COG and KEGG. As a result, a total of 65,577 (37.91 %) unigenes were predicted to be coding sequence and 66,000 (38.16 %), 50,161 (29.00 %), 27,443 (15.87 %), 19,715 (11.40 %) unigenes had homologous sequences respectively in NR, GO, COG and KEGG databases (Table 3).

According to the NR annotation, 66,000 (38.16 %) unigenes had homologous sequence in the database.

| Statistics of data production | Flower buds | Vegetative buds |
|------------------------------|-------------|-----------------|
| Raw reads                    | 35,511,583  | 32,514,423      |
| Raw bases                    | 5,326,737,450 | 4,877,163,450 |
| Clean reads                  | 35,510,239  | 32,513,288      |
| Clean bases                  | 5,280,378,528 | 4,841,503,927 |
| Clean data rate (%)          | 99.125      | 99.27           |

| Total number | 189,300 | 172,959 |
|-------------|---------|---------|
| Average length(bp) | 755.549 | 698.016 |
| Max length(bp)   | 11,797  | 11,797  |
| Min length(bp)   | 201     | 201     |
| GC percentage (%)| 42.1    | 42.0    |
| Length of N50 (bp) | 1452   | 1340    |

| Public protein database | No. of unigene hits | Percentage (%) |
|------------------------|---------------------|----------------|
| Predict protein number | 65,577              | 37.91          |
| COG                    | 27,443              | 15.87          |
| NR (E-value < 10−5)    | 66,000              | 38.16          |
| GO                     | 50,161              | 29.00          |
| KEGG                   | 19,715              | 11.40          |
of which 14,365 (21.77%), 4309 (6.53%), and 4173 (6.32%) unigenes were annotated with homologous genes from *Vitis vinifera*, *Oryza sativa* Japonica Group and *Prunus persica*, respectively (Fig. 2a). The similarity distribution indicated that 54.87% of the unigenes showed a similarity higher than 60, and 40.04% unigenes had similarity between 60 and 80% (Fig. 2b). For the E-value distribution, 53.79% of the top hits had high homology with the E-value <1.0e−50 (Fig. 2c).

As an international standardized gene functional classification system, GO describes properties of genes and their products in many organisms and provides a comprehensive description of gene properties across species and databases (Hao et al. 2011). Based on the BlastX results, a total of 50,161 (29.00%) unigenes were assigned into 49 GO term annotations, including three categories, i.e., molecular function, biological process, and cellular component (Fig. 3). Within the molecular function category (70,280 GO terms), “binding” (32,021, 45.6%) and “catalytic activity” (26,742, 38.0%) were the most highly represented terms. For biological process (108,214 GO terms), “cellular process” (28,941, 26.7%) and “metabolic process” (29,813, 27.6%) were the highly represented terms. Among the cellular component category (87,253 GO terms), “cell” (28,076, 32.2%), “cell part” (13,723, 15.7%) and “organelle” (28,076, 32.2%) were the three main categories.

A total of 27,443 unigenes were classified into 24 COG functional categories (Fig. 4). And the largest category was “general function prediction only” (6177, 22.5%), followed by “signal transduction mechanisms” (2220, 8.1%) and “replication, recombination and repair” (2196, 8.0%). Unigenes annotated as the “signal transduction mechanisms” in our study may allow for the identification of novel genes involved in signal transduction pathways. Approximately 16.9% of the unigenes were associated with biochemical synthesis and metabolism, such as “carbohydrate transport and metabolism” (1242, 4.5%), “amino acid transport and metabolism” (1184, 4.3%) and “secondary metabolites biosynthesis, transport and catabolism” (544, 2.0%). A total of 1225 unigenes (4.46%) was “function unknown”.

Furthermore, we classified 19,715 (11.40%) unigenes into 274 KEGG pathways by searching against the KEGG database (Additional file 3: Table S2). Among these pathways, “metabolic pathways” (3684, 18.7%) represented the largest group, followed by “biosynthesis of secondary metabolites” (1616, 8.2%). The other highly represented pathways included “methyltransferase and metabolism in diverse environment” (803, 4.1%), “ribosome” (509, 2.6%) and “protein processing in endoplasmic reticulum” (507, 2.6%) (Table 4). A total of 301 unigenes involved in plant hormone signal transduction were found, including 9 pathways controlling the signal transduction of auxin, cytokinin, gibberellin acid, abscisic acid, ethylene, brassinosteroid, jasmonic acid and salicylic acid (Table 5). Besides, through the KEGG pathway annotation, a circadian rhythm pathway including 58 unigenes was also found, which can be used for further studies on the flowering-related genes of the photoperiod pathway.

Based on the annotation of unigenes, 118 MADS-box genes were also discovered (Additional file 4: Table S3), including the sepal development related genes such as *APETALA1* (API), *CAULIFLOWER* (CAL); petal development related genes *DEFICIENS* (DEF) and *PISTILLATA* (PI); the C/D/E class function genes *AGAMOUS* (AG), *SEEDSTICK* (STK) and *FLOWERING PROTEIN-like* (FBP-like) were also identified. TCP gene family play important role in the development of plants and can be divided into two classes: PCF class and TCP-C class (which involves the CYC/TB1 clade and CIN clade) (Howarth and Donoghue 2006; Navaud et al. 2007). The CYC/TB1 clade includes genes mainly involved in the development of axillary meristems and floral bilateral symmetry (Luo et al. 1996; Doebley et al. 1997). To determine the TCP gene family in *C. faberi*, we analyzed the transcriptome database generated by

![Fig. 2](characteristics_of_sequence_homology_of_c_faberi_blasted_against_ncbi_non-redundant_nr_database_a_e_value_distribution_of_blast_hits_for_matched_unigene_sequences_using_an_e_value_cutoff_of_1.0e_50_b_similarity_distribution_of_top_blast_hits_for_each_unigene_c_species_distribution_of_the_top_blast_hits)
Fig. 3 GO classification of all annotated unigenes. GO classification of all annotated 50,161 unigenes. All terms belonged to the three main GO categories: biological process, cellular component and molecular function. The x-axis indicated the subcategories, the right y-axis indicated the number of genes in each category, the left y-axis indicated the percentage of a specific category of genes in the corresponding GO category. Red column indicated all annotated unigenes.

Fig. 4 COG annotations of putative proteins. All putative proteins were aligned to the COG database and were functionally classified into at least 24 molecular families. The capital letters in x-axis indicates the COG categories as listed on the right of the histogram and the y-axis indicates the number of unigenes in the corresponding COG category.
this study and found 35 unigenes annotated as TCP TFs, including 14 CIN-like genes, 20 PCF-like genes and 1 CYC/TB1-like genes. As shown in Additional file 5: Table S4, they showed homology with 13 Arabidopsis TCP genes.

A total of 240 genes associated with flowering time were obtained (Additional file 6: Table S5). These include floral meristem identity genes LEAFY (LFY) and APETALA1 (AP1); autonomous pathways genes FCA, FPA, FLOWERING LOCUS (FLD), FY, FVE, FLOWERING LATE KH MOTIF (FLK); vernalization pathways genes FRI and VERNALIZATION INSENSITIVE (VIN); photoperiod pathway genes such as FT, Phytochrome A (PHYA), Phytochrome B (PHYB), PIF3, ELF3, LHY, SUPPRESSOR OF OVEREXPRESSION OF CONSTANS 1 (SOC1), CIRCADIAN CLOCK ASSOCIATED1 (CCA1) and CO; Gibberellin (GA) pathway genes such as GI\textit{BBERELLIC ACID INSENSITIVE} (GAI). All these unigenes provide important resources for future study of floral organ development, floral bilateral symmetry and flowering time.

### DEG analysis of vegetative and flower buds

The FPKM methods were used to analyze the gene expression in the two libraries: 123,128 and 125,862 unigenes were obtained in the flower buds and vegetative buds, of which 39,549 and 42,283 genes expressed specifically in the flower buds and vegetative buds, respectively (Fig. 5). To analyze different gene expression in the two libraries, 13,484 DEGs were identified using FPKM methods. Among them, 7683 were down-regulated and 5801 were up-regulated in the flower buds when compared to those in the vegetative buds, including 3430 and 6556 genes specifically enriched in flower.

| Pathway                                  | No. of all genes with pathway annotation (19,715) | No. of DEGs with pathway annotation (66) | Pathway ID |
|------------------------------------------|--------------------------------------------------|------------------------------------------|------------|
| Metabolic pathways                      | 3681                                             | 60                                       | ko01100    |
| Biosynthesis of secondary metabolites   | 1616                                             | 66                                       | ko01110    |
| Microbial metabolism in diverse environments | 803                                             | 46                                       | ko01120    |
| Ribosome                                | 509                                              | 18                                       | ko03010    |
| Protein processing in endoplasmic reticulum | 507                                             | 14                                       | ko04141    |
| Spliceosome                             | 469                                              | 27                                       | ko03040    |
| RNA transport                           | 457                                              | 21                                       | ko03013    |
| Starch and sucrose metabolism           | 403                                              | 12                                       | ko05000    |
| Cell cycle                              | 355                                              | 25                                       | ko04110    |
| Tuberculosis                            | 337                                              | 11                                       | ko05152    |
| Pyrimidine metabolism                   | 330                                              | 19                                       | ko0240     |
| RNA degradation                         | 327                                              | 19                                       | ko03018    |

| Pathway                                  | Product              | Pathway ID | Unigene number |
|------------------------------------------|----------------------|------------|----------------|
| Cysteine and methionine metabolism       | Ethylene             | ko00270    | 125            |
| Phenylalanine metabolism                 | Salicylic acid       | ko00360    | 103            |
| Tryptophan metabolism                    | Auxin                | ko00380    | 90             |
| alpha-Linolenic acid metabolism          | Jasmonic acid        | ko00592    | 59             |
| Diterpenoid biosynthesis                 | Gibberellin          | ko00904    | 47             |
| Carotenoid biosynthesis                  | Abscisic acid        | ko00906    | 28             |
| Zeatin biosynthesis                      | Cytokinin            | ko00908    | 18             |
| Brassinosteroid biosynthesis             | Brassinosteroid      | ko00905    | 17             |
| Indole alkaloid biosynthesis             | Indole-acetic acid   | ko00901    | 2              |

### Table 5 The pathways involved in plant hormone biosynthesis

| Pathway                                  | Product              | Pathway ID | Unigene number |
|------------------------------------------|----------------------|------------|----------------|
| Cysteine and methionine metabolism       | Ethylene             | ko00270    | 125            |
| Phenylalanine metabolism                 | Salicylic acid       | ko00360    | 103            |
| Tryptophan metabolism                    | Auxin                | ko00380    | 90             |
| alpha-Linolenic acid metabolism          | Jasmonic acid        | ko00592    | 59             |
| Diterpenoid biosynthesis                 | Gibberellin          | ko00904    | 47             |
| Carotenoid biosynthesis                  | Abscisic acid        | ko00906    | 28             |
| Zeatin biosynthesis                      | Cytokinin            | ko00908    | 18             |
| Brassinosteroid biosynthesis             | Brassinosteroid      | ko00905    | 17             |
| Indole alkaloid biosynthesis             | Indole-acetic acid   | ko00901    | 2              |
and vegetative buds, respectively. To validate and annotate the assembled DEGs, the 13,484 DEGs were subjected to BLASTX comparisons (E-value ≤ 10\(^{-5}\)) against GO, COG and KEGG database to identify putative functions of these unigenes. As a result, 5348 (39.7 %), 2829 (21.0 %) and 3927 (29.1 %) DEGs had homologous sequences in GO, COG and KEGG databases, respectively. In addition, 5348 DEGs were classified into 44 GO term annotations, including cellular component (1399), biological process (3921) and molecular function (7006), with multiple terms assigned to the same transcript (Fig. 6). A total of 2829 DEGs were classified into 23 functional COG categories (Fig. 7). “general function prediction only” (506) was the most, followed by “posttranslational modification, protein turnover, chaperones” (248) and “transcription” (222). To further understand the function in biological processes, 3927 DEGs were classified into 245 KEGG pathways. These results showed that most of the DEGs were involved in the “metabolic pathways” (300), “biosynthesis of secondary metabolites” (252), “microbial metabolism in diverse environment” (142) and “ribosome” (89) (Table 4). After transcriptome analysis, TFs involved in the floral differentiation were revealed and their expression levels in the flower and vegetative buds were calculated and compared using the FPKM method.

A total of 173 DEGs related to the floral development were discovered in our transcriptome data (Additional file 7: Table S6). These TFs were attributed to different gene families, including MADS (34 DEGs), TCP (12 DEGs), MYB (34 DEGs), ARF (9 DEGs), NAC (21 DEGs), b-ZIP (17 DEGs), WRKY (18 DEGs) (Table 6). Twenty-eight DEGs related to the flowering time were identified (Additional file 8: Table S7) and some shown in Table 7. Most of these TFs showed higher expression in the flower buds indicating that these genes might involve in the flower development.

![Fig. 5 The numbers of specific genes and shared genes between the flower and vegetative buds of *C. faberi*](image)

![Fig. 6 GO classification of DEGs](image)
qRT-PCR validation of selected genes

To verify the reliability of RNA-Seq data and explicit the expression patterns of genes related to floral development, ten genes mainly related to the floral organ development, floral symmetry and flowering time associated genes were selected to perform qRT-PCR analysis. Our results indicated that \textit{DEF}, \textit{PI}, \textit{AG}, \textit{API1}, \textit{BRANCHED2} (\textit{BRC2}) and \textit{CCA1} had higher expression and \textit{TCP4}, \textit{CO}, \textit{SOC1} and \textit{LFY} displayed lower expression in the flower buds (Fig. 8a-j). \textit{PI} and \textit{DEF}, the two B class function genes controlling the petal development, showed significantly higher expression in the flower buds than the vegetative buds. While \textit{CCA1} participating in the circadian rhythm showed significantly higher expression in the flower buds than the vegetative buds. Overall, all of the 10 genes performed the same expression patterns in our

Table 6 DEGs related to floral differentiation in the flower and vegetative buds

| Genes related to floral differentiation | Total No. of DEGs | No. of DEGs in flower buds | No. of up-regulated DEGs in flower buds | No. of DEGs in vegetative buds | No. of up-regulated DEGs in vegetative buds |
|----------------------------------------|------------------|---------------------------|----------------------------------------|-------------------------------|---------------------------------------------|
| MADS                                   | 34               | 33                        | 29                                     | 8                             | 5                                           |
| TCP                                    | 12               | 12                        | 10                                     | 12                            | 12                                          |
| MYB                                    | 34               | 30                        | 29                                     | 11                            | 5                                           |
| ARF                                    | 9                | 7                         | 1                                      | 9                             | 8                                           |
| NAC                                    | 21               | 20                        | 17                                     | 8                             | 1                                           |
| b-ZIP                                  | 17               | 17                        | 14                                     | 13                            | 4                                           |
| WRKY                                   | 18               | 17                        | 15                                     | 10                            | 3                                           |
qRT-PCR results as in the RNA-Seq data, verifying the reliability of the Illumina sequencing data.

**Discussion**

**Capacity of the transcriptome database**

In recent years, transcriptome analysis based on deep RNA sequencing has been applied to many plants to identify differences in gene expression levels among different cultivars, organs and different treatment conditions (Chen et al. 2014; Yates et al. 2014). In this study, we obtained 189,300 contigs and identified 172,959 unigenes by de novo assembly through RNA-Seq technology in the vegetative and flower buds of *C. faberi*. According to the Nr protein database, 66,000 (38.16%) unigenes were successfully annotated. Furthermore, 173 DEGs involved in floral organ development, floral zygomorphy and flowering time were found. These results supported that plant conservative genes, *C. faberi*-specific genes, and *C. faberi* tissue-specific genes all were identified in our transcriptomic analysis. Our research will provide valuable information for future study to inquire the flower development mechanisms of *C. faberi*.

**Table 7 Some DEGs related to flowering time in the flower and vegetative buds**

| Genes related to flowering | Total No. of DEGs | No. of DEGs in flower buds | No. of up-regulated DEGs in flower buds | No. of DEGs in vegetative buds | No. of up-regulated DEGs in vegetative buds |
|---------------------------|-------------------|-----------------------------|----------------------------------------|------------------------------|-------------------------------------------|
| PHYA                      | 1                 | 1                           | 1                                      | 1                            | 1                                         |
| CCA1                      | 3                 | 2                           | 2                                      | 3                            | 1                                         |
| LHY                       | 1                 | 1                           | 1                                      | 1                            | 0                                         |
| FCA                       | 2                 | 1                           | 1                                      | 1                            | 1                                         |
| FY                        | 1                 | 1                           | 0                                      | 1                            | 1                                         |
| FRI                       | 3                 | 1                           | 1                                      | 2                            | 2                                         |
| CO                        | 2                 | 1                           | 1                                      | 2                            | 1                                         |

**Genes related to the floral organ development**

MADS-box genes are known for their roles in the flower organ development in *Arabidopsis* and *Antirrhinum* (Coen and Meyerowitz, 1991; Weigel and Meyerowitz, 1994). While some monocot species like the orchid family possess distinct floral structures, thus floral patterning in *Arabidopsis* may not be comparable to such flowering.

![Fig. 8](image)

qRT-PCR analysis of ten selected genes in different organs of *C. faberi*. a–j indicated the relative expression level of DEF, PI, AG, AP, TCP4, BRC2, CO, SOC1, CCA1, LFY gene, respectively. In these pictures, F flower buds; V vegetative buds; Asterisk indicated significantly higher expression.
Genes related to the floral zygomorphy
The ornamental value of *C. faberi* is determined by many factors, especially the novel flower color, shape and fragrance. Floral zygomorphy have evolved multiple times from radially symmetrical (actinomorphic; polysymmetric) ancestors in different angiosperm lineages (Endress 1999; Stebbins 1974). The mechanism of TCP model for bilateral flower symmetry has been well established in model species, such as snapdragon (*A. majus*), *Lotus japonicus* and rice (Luo et al. 1996; Feng et al. 2006; Yuan et al. 2009). In *A. majus*, *CYC* and *DICH* control the dorsoventral asymmetry (Luo et al. 1999). In *L. japonicus*, floral dorsoventral asymmetry is regulated by three *LjCYC* genes (*LjCYC1*, *LjCYC2*, *LjCYC3*), especially the *LjCYC2*. In monocot, *RETARDED PALEA1 (REPI)* controls palea development and floral zygomorphy in rice, whose flower is different from that of *A. majus* or *L. japonicus*. In our *C. faberi* transcriptome data, we identified 35 TCP genes, of which 10 *CIN*-like genes and one *PCF*-like gene showed significantly higher expression levels in the vegetative buds compared to the flower buds. qRT-PCR analysis revealed that the expression level of *BRC2* was higher in the flower buds than the vegetative buds (Fig. 8e), suggesting that it might play important role in the regulation of floral zygomorphy of *C. faberi*.

Genes related to the regulation of flowering
Previous studies showed that the flowering of *Arabidopsis* was regulated by four pathways, including autonomous pathway, vernalization pathway, gibberellic acid (GA)-dependent pathway and the photoperiod pathway. And *CCA1* influenced the circadian rhythm, overexpression of *CCA1* resulted in long hypocotyls and late flowering (Wang and Tobing, 1998). *CO* involved in the photoperiod pathway in *Arabidopsis* and acted as a floral promoter, which was regulated by the circadian clock (Yano et al. 2000). The temporal and spatial regulation of *CO* turned out to be important to the photoperiod-dependent induction of flowering (An et al. 2004). Early in a day, the expression of *CO* was low, then increased rapidly 10 h after dawn and peaked at around 15 h (Suárez-López et al. 2001). *LFY* is a major floral meristem regulator, its overexpression caused early flowering in transgenic *Arabidopsis* (Benlloch et al. 2007). A single *HYA* gene, *LHY* gene and three *CCA1* genes involved in the photoperiod pathways showed more highly expressed in the vegetative buds than in the flower buds. Two *VRN* genes playing important roles in the vernalization pathway were detected in our study, and were found to be more highly expressed in the vegetative buds than in the flower buds. Besides, DEGs related to the autonomous pathway were also detected, including two *FCA* genes, a *FLD* gene and a *FY* gene. qRT-PCR revealed that expression of *LFY* was higher in the vegetative buds and expression of *CO*, *SOC1*, *CCA1* was higher in the flower buds, which was consistent with the RNA-Seq data (Fig. 8g–j).

Expression of *CCA1* was significantly higher in the flower buds than the vegetative buds. This observation suggesting that the mechanism of flowering regulation in *C. faberi* might be similar to *Arabidopsis*.

In conclusion, the current RNA-seq and DEG analysis revealed 393 genes associated to the floral development, including 35 TCP transcription factors (TFs), 118 MADS-box genes and 240 flowering time related genes. A total of 173 DEGs were identified, including 12 TCP genes, 34 MADS-box genes and 28 flowering time related genes. The transcriptome database in the present study will be a valuable supplement to the genomic sequence dataset of *C. faberi*, and will serve as an important public information platform for further studies on the flower development mechanism in *C. faberi*.

Additional files
Authors' contributions
The experimental plan was conceived and designed by GW, YS, YL and SG participated in sample collection, RNA preparation and experiments performing. YS, JL and YY contributed to data analysis and manuscript preparation. GW participated in drafting and revising the manuscript. All authors read and approved the final manuscript.

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Competing interests
The authors declare that they have no competing interests.

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