Comprehensive RNA sequencing and co-expression network analysis to complete the biosynthetic pathway of coumestrol, a phytoestrogen

Jungmin Ha1,2, Young-Gyu Kang3, Taeyoung Lee3, Myoyeon Kim3, Min Young Yoon1,2, Eunsoo Lee1, Xuefei Yang1, Donghyun Kim3, Yong-Jin Kim3, Tae Ryong Lee3, Moon Young Kim1,2 & Suk-Ha Lee1,2

Coumestrol (CMS), a coumestan isoflavone, plays key roles in nodulation through communication with rhizobia, and has been used as phytoestrogens for hormone replacement therapy in humans. Because CMS content is controlled by multiple genetic factors, the genetic basis of CMS biosynthesis has remained unclear. We identified soybean genotypes with consistently high (Daewonkong) or low (SS0903-2B-21-1-2) CMS content over 2 years. We performed RNA sequencing of leaf samples from both genotypes at developmental stage R7, when CMS levels are highest. Within the phenylpropanoid biosynthetic pathway, 41 genes were tightly connected in a functional co-expression gene network; seven of these genes were differentially expressed between two genotypes. We identified 14 candidate genes involved in CMS biosynthesis. Among them, seven were annotated as encoding oxidoreductases that may catalyze the transfer of electrons from daidzein, a precursor of CMS. Two of the other genes, annotated as encoding a MYB domain protein and a MLP–like protein, may increase CMS accumulation in response to stress conditions. Our results will help to complete our understanding of the CMS biosynthetic pathway, and should facilitate development of soybean cultivars with high CMS content that could be used to promote the fitness of plants and human beings.

Plants synthesize secondary metabolites to promote their own survival, and some of these compounds have antioxidant or antibiotic effects1,2. Several lines of evidence suggest that plant secondary metabolites, especially isoflavones, can improve the fitness of both humans and plants. Accordingly, a great deal of research has been conducted on isoflavones. Legume species, including the economically important crop plant soybean, are rich in isoflavones with estrogenic and antioxidant functions3,4. Within plants themselves, isoflavones play crucial roles in nodulation and nitrogen fixation5 and defense against environmental stresses6. In the context of human health, isoflavones, as phytoestrogens, can decrease the risk of menopausal symptoms, breast cancer, osteoporosis, dementia, and cardiovascular disease7–13.

Declining estrogen levels in postmenopausal women are associated with a variety of cutaneous changes, including dryness, wrinkling, poor healing, and hot flashes, many of which can be improved by estrogen supplementation14. However, the estrogens used in hormone replacement therapy can promote the initiation and progression of breast cancer15–17. The effects of estrogen are mediated by two estrogen receptors (ERs), ERα and ERβ, which are distributed differently in each tissue18,19. ERα mediates the breast cancer–promoting effects of estrogens, whereas ERβ inhibits breast cancer cell proliferation and tumor formation20. Therefore, it has been proposed that dietary or synthetic ERβ-selective estrogens would lack the breast cancer–promoting properties of the estrogens used in hormone replacement regimens20.

1Department of Plant Science and Research Institute of Agriculture and Life Sciences, Seoul National University, Seoul, Republic of Korea. 2Plant Genomics and Breeding Institute, Seoul National University, Seoul, Republic of Korea. 3Basic Research & Innovation Division, R&D Center, AmorePacific Corporation, Yongin-si, Gyeonggi-do, Republic of Korea. Correspondence and requests for materials should be addressed to S.-H.L. (email: sukhealee@snu.ac.kr)
Coumestrol (CMS), a coumestan isoflavone, is the most abundant polyphenol in soybean leaves and functions as a phytoestrogen that is structurally and functionally similar to 17\(^{-}\)-estradiol, an estrogen steroid hormone. ER-binding assays revealed that CMS has a 15-fold higher binding affinity for ER\(\beta\) than for ER\(\alpha\). Intake of CMS is associated with reduced risk of breast cancer; in addition, CMS prevents skin photoaging by suppressing FMS-like tyrosine kinase 3, which causes collagen degradation and skin wrinkling. CMS can decrease melanin synthesis, which darkens the skin, as well as alleviate symptoms caused by excessive melanin synthesis, such as melisma, solar lentigo, dark spots, and freckles. In light of these health benefits, CMS has been suggested as a promising dietary supplement that could prevent disease and improve the health of postmenopausal women.

CMS is a soybean phytoalexin that is present in soybean leaves and roots, rather than seeds, and CMS content varies depending on environmental conditions and growth stage. CMS accumulates to high levels after drought stress in root, doubling the extent of mycorrhizal colonization. Because other isoflavones, such as daidzein and formononetin, are involved in signaling in rhizosphere plant-microbe interactions, CMS has been implicated in drought tolerance in legumes, an effect mediated through communication with mycorrhiza.

CMS is derived from the soybean isoflavone daidzein, via dihydrodaidzein and 2’-hydroxydaidzein, through two biosynthetic pathways that remain incompletely understood. Because many environmental factors affect the biosynthesis and accumulation of isoflavones, and epistatic interactions among multiple QTLs with small individual effects are responsible for a large proportion of the variation, it has been very challenging to elucidate the genetics governing isoflavone biosynthesis. Due to the important implications of soybean isoflavones and phytoalexins for plant defense and human health, it would be valuable to identify the enzymes responsible for CMS biosynthesis from its precursor, daidzein. Knowledge of these enzymes would facilitate successful manipulation of CMS levels in planta.

In this study, we measured the CMS contents of 31 soybean genotypes, and selected those with consistently high or low CMS contents for gene expression profiling. To shed light on the genetics of CMS biosynthesis, comprehensive RNA sequencing (RNA-seq) was conducted on the leaf tissues at growth stage R7, when CMS levels are highest. Based on the differentially expressed genes (DEGs) between high-CMS and low-CMS genotypes, as well as their functional co-expression network, we identified candidate genes involved in biosynthesis of CMS from daidzein. Our results provide a set of target genes for manipulations aimed at increasing CMS levels in soybean cultivars, with the goal of improving the welfare of menopausal women from clinical and cosmetic perspectives.

**Results**

**CMS contents in soybean genotypes.** We measured CMS contents in leaf samples from 31 soybean genotypes collected at growth stage R7 in 2016; three replicates were performed for each genotype (Supplementary Fig. 1). CMS content varied from 0 to 1,650.55 μg/g. Among the 31 cultivars tested, four with high CMS content, Chamame (1,650.55 μg/g), Geomjeongsaeol (1,209.70 μg/g), SG-257 (1,173.11 μg/g), and Daewonkong (802.08 μg/g), and four with low CMS content, SS0903-2B-21-1-2 (30.32 μg/g), Haepum (39.79 μg/g), SS0905-2B-179-1-1 (48.98 μg/g), and Sinhwa (142.90 μg/g), were selected for repeated cultivation and testing. CMS contents were measured in 2017, again in three replicates, to verify the results from 2016 (Fig. 1A). Daewonkong and SS0903-2B-21-1-2 had the highest (1,781.79 μg/g) and the lowest (3.81 μg/g) CMS contents, respectively, in 2017. These two genotypes with consistently high and low CMS contents for two years were further investigated by RNA-seq to reveal the underlying CMS biosynthesis.

**RNA-seq and DEG profiling.** To identify the differences in gene expression involved in phenylpropanoid biosynthesis between Daewonkong and SS0903-2B-21-1-2, we extracted total RNA from leaf tissues at developmental stage R7. A total of 237 and 245 million reads, encompassing 24 and 25 Gb, respectively, were generated.
per genotype and about 70% and 75% of total reads were properly mapped to the *G. max* reference genome sequence ([www.phytozome.net/soybean](http://www.phytozome.net/soybean)) (Supplementary Table 1). Comparison of expression levels between Daewonkong and SS0903-B2-21-1-2 revealed 4,046 DEGs with FC value of at least 4 (1,629 DEGs with FC 8, 684 DEGs with FC 16 and 283 DEGs with FC 32) and, overall about 50% of DEGs were up-regulated in Daewonkong (45.7–52.6%) (Fig. 2A). The DEGs were annotated against the AgriGO genome locus background, and meta-bolic process was the most enriched GO term in all four DEG sets (Fig. 2C–F). KEGG pathway analysis revealed that ~40% and ~30% of DEGs were assigned to metabolic pathways and biosynthesis of secondary metabolites, respectively (Supplementary Fig. 2). These results indicate that the secondary metabolic pathways are differentially regulated between the two genotypes.

**The expression of genes in the phenylpropanoid biosynthesis pathway.** To characterize differences in the expressions of genes involved in phenylpropanoid biosynthesis, we searched KEGG ([http://www.genome.jp/kegg/](http://www.genome.jp/kegg/)) for soybean homologs of key enzymes in this pathway. In the biosynthetic pathway of phenylpropanoid, starting from phenylalanine to daidzein and its byproducts (Supplementary Fig. 3), we identified 72 candidate soybean homologs. Among them, 41 were connected with each other in a functional co-expression network in the SoyNet database ([http://www.inetbio.org/soynet/](http://www.inetbio.org/soynet/)) (Fig. 3). Eight of the genes in this network were differentially expressed between Daewonkong and SS0903-B2-21-1-2, and thus represent the DEGs most likely to affect the content of CMS, an end product of isoflavone biosynthesis (Supplementary Table 3). Two of these
genes, Glyma.16G149300 (LOC100811727) and Glyma.17G064400 (LOC100779668), were down-regulated in Daewonkong (log₂FC −2.7 and −3.1), whereas the other six, Glyma.01G228700 (chalcone synthase, CHS7), Glyma.01G239600 (2-hydroxyisoflavanone dehydratase, HIDH), Glyma.11G010500 (4-coumarate:CoA ligase, 4CL13), Glyma.11G011500 (CHS8), Glyma.13G173500 (isoflavone synthase, IFS2), and Glyma.14G005700 (chalcone reductase, CHR14), were up-regulated (up to log₂FC 6.5). Five of the up-regulated genes were tightly connected with each other in the co-expression network of the phenylpropanoid biosynthesis pathway (Fig. 3, Supplementary Fig. 4). Among the eight DEGs, seven in the upstream pathway of CMS biosynthesis (i.e., all except Glyma.16G149300) were used to predict candidate genes for the unknown pathway (Fig. 3).

Identification of candidate genes for CMS biosynthesis. To identify candidate genes for biosynthesis of CMS from daidzein, we applied three prediction approaches (Supplementary Table 4). First, using the 41 genes in the network as guide genes (Fig. 3), we searched the whole soybean gene network of 40,812 genes for genes closely connected to the guide genes (“guide prediction”). Second, the genes closely connected to the seven DEGs involved in phenylpropanoid biosynthesis were identified using the same network. Third, candidate genes were predicted in the context of subnetworks consisting of central hubs and their neighbors, and connections between the hub genes and the seven DEGs were identified (“hub prediction”) (Supplementary Table 4)44. We then listed the top 20 genes predicted from each approach, along with their GO terms from three different databases. The three methods identified 3, 11, and 9 genes as DEGs, all of which were up-regulated in Daewonkong (Supplementary Fig. 5). Overall, 14 genes were predicted from the three prediction approaches, of which seven were identified by two or more approaches (Supplementary Table 5). These 14 candidate genes may play a key role in determining CMS contents in soybean.

Validation of gene expression level by qRT-PCR. We validated the expression levels of the 14 DEGs by quantitative reverse-transcription (qRT) PCR (Fig. 4). The qRT-PCR results were consistent with the RNA-seq data: all 14 DEGs up-regulated in Daewonkong in the RNA-seq data were also up-regulated in the qRT-PCR results (Supplementary Fig. 5).

Proposed model for the biosynthetic pathway from Daidzein to CMS. Based on guide prediction and hub prediction approaches, 14 DEGs were predicted to be involved in CMS biosynthesis (Supplementary Table 5). All 14 were highly up-regulated in Daewonkong, and qRT-PCR results were in close agreement with the RNA-seq data. These genes were mapped against three GO databases: agriGO, AtgO, and Uniprot-GO. Of the 14, four (Glyma.02G307300, Glyma.09G269500, Glyma.11G070500, and Glyma.18G220600) were mapped to steroid
Figure 4. Validation of RNA-seq results by qRT-PCR. Left and right y-axes indicate FPKM values from RNA-seq (blue bar) and relative transcript abundance from qRT-PCR (orange bar). D and S on the x-axis represent Daewonkong and SS0903-2B-21-1-2, respectively. Bars indicate means and standard deviation of three biological replicates. Asterisk above each bar indicates statistical difference between genotypes, as determined by Student's t-test ($p < 0.05$). (A) Glyma.01G135200, cytochrome P450, family 82, subfamily C, polypeptide 4. (B) Glyma.02G005600, myb domain protein 15. (C) Glyma.02G134000, carboxyesterase 13. (D) Glyma.02G307300, NAD(P)-linked oxidoreductase superfamily protein. (E) Glyma.03G147700, disease resistance–responsive (dirigent-like protein) family protein. (F) Glyma.09G269500, NAD(P)-binding Rossmann-fold superfamily protein. (G) Glyma.11G004200, alpha/beta-Hydrolases superfamily protein. (H) Glyma.11G070500, NmrA-like negative transcriptional regulator family protein. (I) Glyma.13G284900, organic cation/carnitine transporter4. (J) Glyma.13G285300, cytochrome P450, family 82, subfamily C, polypeptide 4. (K) Glyma.17G030400, MLP-like protein 423. (L) Glyma.18G220600, NAD(P)-binding Rossmann-fold superfamily protein. (M) Glyma.18G285800, NAD(P)-linked oxidoreductase superfamily protein. (N) Glyma.19G030500, HXXXD-type acyl-transferase family protein.
and flavonoid biosynthetic processes; two (Glyma.02G005600 and Glyma.17G030400) were related to stress responses; and eight genes were not previously mapped in any GO database. The biosynthetic reactions from daidzein to (3R)-2′-hydroxydihydrodaidzein comprise a series of oxidations (NADPH → NADP⁺) and hydrolysis (Fig. 5). All four DEGs involved in steroid and flavonoid biosynthetic processes (Glyma.02G307300, Glyma.09G269500, AT1G59960.1, NAD(P)-linked oxidoreductase superfamily protein; Glyma.09G269500, AT1G59960.1, NAD(P)-linked oxidoreductase superfamily protein; Glyma.18G220600, AT1G59960.1, NAD(P)-linked oxidoreductase superfamily protein; and Glyma.18G285800, AT1G59960.1, NAD(P)-linked oxidoreductase superfamily protein). The blue box on the bottom right includes Glyma.02G005600, AT3G23250.1, myb domain protein 15; and Glyma.17G030400, AT1G24020.1, MLP-like protein 423. The blue box on the bottom right includes Glyma.02G134000, AT3G48700.1, carboxylesterase 13; Glyma.03G147700, AT5G42500.1, disease resistance–responsive (dirigent-like protein) family protein; Glyma.11G004200, AT1G47390.1, alpha/beta-Hydrolases superfamily protein; Glyma.13G284900, AT3G20660.1, organic cation/carnitine transporter4; and Glyma.19G030500, AT5G39090.1, HXXXD-type acyl-transferase family protein.

**Figure 5.** Candidate genes involved in the biosynthetic pathway from daidzein to CMS. The lists of genes in the blue boxes indicate the candidate genes involved in the pathways of CMS biosynthesis. The blue box at the top includes Glyma.01G135200, AT4G31940.1, cytochrome P450, family 82, subfamily C, polypeptide 4; Glyma.02G307300, AT1G59960.1, NAD(P)-linked oxidoreductase superfamily protein; Glyma.09G269500, AT2G45400.1, NAD(P)-binding Rossmann-fold superfamily protein; Glyma.11G070500, AT4G39230.1, NmrA-like negative transcriptional regulator family protein; Glyma.13G285300, AT4G31940.1, cytochrome P450, family 82, subfamily C, polypeptide 4; Glyma.18G220600, AT2G45400.1, NAD(P)-binding Rossmann-fold superfamily protein; and Glyma.18G285800, AT1G59960.1, NAD(P)-linked oxidoreductase superfamily protein. The blue box on the bottom left includes Glyma.02G005600, AT3G23250.1, myb domain protein 15; and Glyma.17G030400, AT1G24020.1, MLP-like protein 423. The blue box on the bottom right includes Glyma.02G134000, AT3G48700.1, carboxylesterase 13; Glyma.03G147700, AT5G42500.1, disease resistance–responsive (dirigent-like protein) family protein; Glyma.11G004200, AT1G47390.1, alpha/beta-Hydrolases superfamily protein; Glyma.13G284900, AT3G20660.1, organic cation/carnitine transporter4; and Glyma.19G030500, AT5G39090.1, HXXXD-type acyl-transferase family protein.
(Glyma.02G037300 and Glyma.18G285800) catalyze the transfer of electrons from one molecule to another using NADPH or NADP⁺ as a cofactor, and the NAD(P)-binding Rossmann fold (Glyma.09G269500, Glyma.11G070500, and Glyma.18G220600) is involved in catalysis of NAD(P)-dependent oxidation. The NmA-like protein encoded by Glyma.11G070500 contains two domains, including a Rossmann fold. Therefore, we speculate that the proteins encoded by these seven genes catalyze the NADPH oxidation reactions starting from daidzein (Fig. 5).

Discussion

Soybean is one of the most important crops in the world due to its high production of protein and oil. In addition, it is a valuable nutraceutical ingredient because it contains several phytochemicals, including isoflavones, saponins, phenolic acids, and linoleic acids. In light of their contribution to human health and plant defense systems, these phytochemicals, especially isoflavones, are desirable target traits in soybean breeding programs. Isoflavones, synthesized predominantly in legumes, attract rhizobial bacteria, initiate nitrogen-fixing root nodule formation, exert antifungal activity, and serve as metabolic precursors for major phytoalexins. CMS, a coumarinoid isoflavone, decreases the risk of breast cancer by binding selectively to ERα. Therefore, CMS is a promising phytostrogen for use as a selective estrogen receptor modulator (SERM).

The extreme variability of isoflavone contents among different environments has hindered elucidation of the genetic basis of isoflavone biosynthesis. Indeed, in this study, even though plant samples were prepared at the same location over 2 years in three biological replicates in each year, variations in CMS content were observed between 2016 and 2017, indicating that this trait is environmentally sensitive. Multiple MYB TFs regulate the expression of structural genes involved in isoflavone biosynthesis under stressed conditions. Other than the candidate genes that catalyze the NADPH oxidation reactions starting from daidzein, there are several more candidate genes involved in the biosynthesis pathway of CMS. MYB domain protein 15 (Glyma.02G005600) and major latex protein (MLP)-like protein 423 (Glyma.17G030400) are annotated as involved in stress responses. Because the regulation of isoflavonoid metabolism is thought to occur primarily at the level of transcription, transcription factors (TFs) are promising candidates. Multiple MYB TFs regulate the expression of structural genes involved in isoflavone biosynthesis under stressed conditions. MLP-like protein modulates the production of metabolites under drought stress conditions, and overexpression of MLP leads to salt stress insensitivity. This agrees well with reports that CMS increases drought stress through communication with mycorrhiza and CMS accumulates the most at R7 growth stage when soybean is dehydrated while maturating. Thus, these two genes may induce accumulation of CMS under stress conditions.

For candidate DEGs unmapped to any GO term, Glyma.01G135200 and Glyma.13G285300 are annotated as cytochrome P450, an oxidoreductase, that is reported to be involved in isoflavonoid biosynthesis. Glyma.11G004200, annotated as a member of the alpha/beta-hydrolase superfamily, may play a role in the hydrolysis reaction of CMS biosynthesis. 2-hydroxysiosflavanone dehydratase, which catalyzes a dehydration reaction yielding isoflavone from 2-hydroxysiosflavanone, is a member of the carboxylate family. Glyma.02G134000 is down-regulated in Daewonkong, consistent with the higher level of CMS accumulation in this genotype (Fig. 3).
Materials and Methods

Plant materials for HPLC analysis. For CMS measurements, 31 soybean genotypes were planted at the Seoul National University Experimental Farm in Suwon, South Korea (37.3°N, 127.0°E), with three replications per cultivar, in 2016 (Supplementary Fig. 1). Average temperature and duration of sunshine in Suwon from May to October of 2016 were 16.1–28.0 °C and 1,293.6 hr, respectively. In 2017, eight genotypes, including four with high CMS and four with low CMS in 2016, were planted again in three replicates (Fig. 1A). Average temperature and duration of sunshine in Suwon from May to October of 2017 were 16.4–26.9 °C and 1,334.4 hr, respectively.

Soybean leaves collected at growth stage R7 were dried and ground to a fine powder. Each sample (3 g) was stirred in 45 mL of 80% ethanol for 1 day at room temperature, and then the mixture was filtered through a 0.45 μm GHP membrane filter (Acrodisc 13 mm syringe filter; Pall Corporation, Port Washington, NY, USA). Quantitative analysis of CMS was performed on a Mightsyl RP-18 GP reversed phase column (5 μm, 4.6 × 250 mm) (Kanto Chemical Co., Tokyo, Japan) at sub-ambient temperature, using a 40 min linear gradient of 0.1% glacial acetic acid in water (solvent A) and 0.1% glacial acetic acid in acetonitrile (solvent B). The linear gradient program was as follows: 0–5 min, 31% B; 5–25 min, 31–35% B; 25–30 min, 35–80% B; 30–35 min, 80% B; 35–40 min back to 31% B. The solvent flow rate was 1.0 mL min−1, and the injection volume was 10 μL. UV absorption was measured at 342 nm to detect CMS (the standard purchased from Sigma-Aldrich Co., St Louis, MO, USA).

RNA-seq. Total RNA was extracted from growth stage R7 leaf samples of Daewonkong and SS903-B2-21-1-2 in 2017 using Ribospin™ Plant (GeneAll, Seoul, Korea). Three cDNA libraries per genotype were constructed using the TruSeq® RNA Sample Prep Kit v2 (Illumina Inc., CA, USA). The quality and quantity of samples used for sequencing were checked using a 2100 Bioanalyzer (Agilent Technologies). RNA samples were sequenced using the TruSeq SBS kit v3 on the Illumina HiSeq. 4000 platform. The raw RNA reads of Daewonkong (SRR6756974, SRR6756973, and SRR6756972) and SS903-B2-21-1-2 (SRR6756971, SRR6756976, and SRR6756975) have been deposited in the National Center for Biotechnology Information (NCBI) Sequence Read Archive (Supplementary Table 1).

DEGs and enrichment analysis. Fragments Per Kilobase of transcript per Million mapped reads (FPKM) values were calculated by mapping raw RNA reads for 56,044 genes to the G. max reference genome annotation data (Gmax.275_Wm82.a2.v1.gene.gff3) using the Tuxedo software suite36. DEGs were defined as genes with a log fold change (FC) ≥ 2 between two samples in pairwise comparisons for three replications with p-value < 0.05 (Fig. 2A). FPKM values < 1 were converted to 1 for the purpose of calculating FC. Sets of DEGs with log FC ≥ 2, 3, 4, or 5 were used for Gene Ontology (GO) enrichment analysis using the Singular Enrichment Analysis (SEA) tool, available at agrigo (http://bioinfo.caau.edu.cn/agriGO/), and KEGG (Kyoto Encyclopedia of Genes and Genomes) ontology (http://www.genome.jp/kegg/tool/map_pathway1.html) (p < 0.05) (Supplementary Fig. 2). The G. max reference annotation (Wm82.a2.v1) was used as a background reference for enrichment analysis.

Candidate gene prediction. New candidate genes in the phenylpropanoid biosynthetic pathway were identified using the soybean co-expression network consisting of 40,812 genes and 1,940,284 co-functional links between 40,812 soybean genes constructed based on 734 microarrays and 290 RNA-seq data from soybean, available at http://www.soynet.org/soynet/44. First, new candidate genes in the pathway were identified by searching genes closely connected to the 41 known genes in the pathway as guide genes in the co-expression network (“guide prediction”). Second, the seven DEGs among the 41 genes in the pathway were used as guide genes for the same function. The genes were prioritized according to the sum of their log-likelihood scores, and highly ranked genes were considered good candidates for new members of the pathway44. Third, DEGs was used to identify new candidate genes through subnetworks consisting of a central hub and their neighbors (“hub prediction”). If significant overlap was observed between DEGs and neighbor genes of a subnetwork, the central hub of the subnetwork was identified as a candidate gene. Candidate genes were prioritized according to their p-values.

qRT-PCR validation of DEGs. Gene-specific primers for qRT-PCR analysis were designed based on the nucleotide sequences of selected DEGs using Primer3 (http://primer3plus.com/) (Supplementary Table 2). cDNA was synthesized using an iScript™ cDNA Synthesis Kit (Cat. 170-8891; Bio-Rad, Hercules, CA, USA). qRT-PCR was conducted using an SYBR Green Supermix kit (Cat. 170-8882; Bio-Rad) on a LightCycler® 480 (Roche Diagnostics, Laval, QC, Canada). Actin was used for normalization of target gene expression, and each sample was analyzed in triplicate. Relative gene expression was analyzed based on the reference gene as previously described56. Student’s t-test was performed to determine whether differences were statistically significant (p < 0.05).
**Data Availability**

The raw RNA sequencing reads were deposited at NCBI SRA.

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Author Contributions
J.H. wrote this manuscript. Y.G.K., M.K., D.K., Y.J.K. and T.R.L. screened CMS contents. T.L. performed RNA sequence data analysis. M.Y., E.L. and M.Y.K. prepared plant samples for RNA sequencing. X.Y. performed qRT-PCR. S.H.L. has correspondence. All authors reviewed the manuscript.

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