Hormonal and transcriptional analyses provides new insights into the molecular mechanisms underlying root thickening and isoflavonoid biosynthesis in *Callerya speciosa* (Champ. ex Benth.) Schot

Shaochang Yao¹, Zuzai Lan², Rongshao Huang¹, Yong Tan¹, Ding Huang¹, Jinyuan Gu¹ & Chunliu Pan²*

*Callerya speciosa* (Champ. ex Benth.) Schot is a traditional Chinese medicine characterized by tuberous roots as the main organ of isoflavonoid accumulation. Root thickening and isoflavonoid accumulation are two major factors for yield and quality of *C. speciosa*. However, the underlying mechanisms of root thickening and isoflavonoid biosynthesis have not yet been elucidated. Here, integrated morphological, hormonal and transcriptomic analyses of *C. speciosa* tuberous roots at four different ages (6, 12, 18, 30 months after germination) were performed. The growth cycle of *C. speciosa* could be divided into three stages: initiation, rapid-thickening and stable-thickening stage, which cued by the activity of vascular cambia. Endogenous changes in phytohormones were associated with developmental changes during root thickening. Jasmonic acid might be linked to the initial development of tuberous roots. Abscisic acid seemed to be essential for tuber maturation, whereas IAA, cis-zeatin and gibberellin 3 were considered essential for rapid thickening of tuberous roots. A total of 4337 differentially expressed genes (DEGs) were identified during root thickening, including 15 DEGs participated in isoflavonoid biosynthesis, and 153 DEGs involved in starch/sucrose metabolism, hormonal signaling, transcriptional regulation and cell wall metabolism. A hypothetical model of genetic regulation associated with root thickening and isoflavonoid biosynthesis in *C. speciosa* is proposed, which will help in understanding the underlying mechanisms of tuberous root formation and isoflavonoid biosynthesis.

**Abbreviations**

- AP2/ERF: APETALA2/ethylene-responsive factor
- BGLU: Beta-glucosidase
- bHLH: Basic helix loop-helix transcription factor
- CK: Cytokinin
- GBSS: Granule-bound starch synthase genes
- H4OMT: Isoflavone 4′-O-methyltransferase
- IAA: Indole-3-acetic acid
- KEGG: Kyoto encyclopedia of genes and genomes
- PAL: Phenylalanine ammonialyase

¹College of Pharmacy, Guangxi University of Chinese Medicine, Nanning, Guangxi, People’s Republic of China. ²Guangxi Botanical Garden of Medicinal Plants, Nanning, Guangxi, People’s Republic of China. *Email: yaoshaochang@163.com; pchunliu@126.com*
Results

Definition of key developmental stages of tuberous roots. To determine the key developmental stages of tuberous roots, roots were sampled at half-year intervals after germination. The root volume, fresh weight and dry weight increased slowly in 6–12 months after germination (MAG), but they increased significantly from 12 to 24 MAG. From 24 to 36 MAG, the root volume and fresh weight were barely changing, while the root dry weight was increasing (Fig. 1a). Thus, the rapid growth phase occurred by root saturation with water. The starch content analysis revealed that it increased significantly from 18 to 30 MAG, but there was no significant difference before 18 MAG (Fig. 1b). However, the formononetin and maackiain contents increased continuously along with the process of tuberous root development (Fig. 1c,d), which indicated that formononetin and maackiain accumulation simultaneously increased along with dry weight during the growth of *C. speciosa*. 

**Tables**

| NAC | NAM, ATAF, and CUC transcription factors |
| SS  | Starch synthase                            |
| TPS | Trehalose-phosphate synthase               |
| UGP | UTP-glucose-1-phosphate uridyltransferase  |

**Abbreviations**

*C. speciosa* (Champ. ex Benth.) Schott, a perennial shrub plant in the *Fabaceae* family, is widely cultivated in South China and commonly used for medicine. Tuberous root is the medicinal organ of *C. speciosa*, which originates from the expansion of fibrous roots. Tuber morphogenesis, along with the accumulation of a large amount of starch as well as health-promoting components, is the main processes of tuberous root development.

Isoflavonoids as the main medicinal compounds have been produced in the tuberous roots of *C. speciosa*, such as the index compounds maackiain and formononetin, which have made considerable contributions to immunity enhancement, hepatoprotection, arresting cough, and expectorant and anti-asthmatic effects. The quality of *C. speciosa* is mainly evaluated according to the degree of root thickening and the content of isoflavonoids, while the harvest period is primarily depended on the yield. Generally, the fibrous roots of *C. speciosa* possess the potential to form tuberous roots, but limited fibrous roots can transfer into tuberous roots. Studies of root thickening and isoflavonoid biosynthesis of *C. speciosa* are scarce.

**Abbreviations**

In this study, twelve libraries prepared from four different age points during tuberous root formation were sequenced by using BGISEQ-500 platform. A detailed comparative mRNA analysis was detected, and the key genes encoding downstream pathway enzymes. A comprehensive understanding of the genes involved in isoflavonoid biosynthesis during the development of tuberous roots in *C. speciosa* will help in revealing the mechanism of isoflavonoid biosynthesis. In this study, twelve libraries prepared from four different age points during *C. speciosa* tuber expansion were sequenced by using BGISEQ-500 platform. A detailed comparative mRNA analysis was detected, and the key genes that potentially participated in root thickening and isoflavonoid biosynthesis were identified by integrating the phenotypic data and gene expression profiles. The results will be valuable for further research, and will help in understanding the molecular mechanisms of tuberous root formation and isoflavonoid metabolism in *C. speciosa*. 

**Abbreviations**

**Abbreviations**

*Callerya speciosa* (Champ. ex Benth.) Schott, a perennial shrub plant in the *Fabaceae* family, is widely cultivated in South China and commonly used for medicine. Tuberous root is the medicinal organ of *C. speciosa*, which originates from the expansion of fibrous roots. Tuber morphogenesis, along with the accumulation of a large amount of starch as well as health-promoting components, is the main processes of tuberous root development. 

Isoflavonoids as the main medicinal compounds have been produced in the tuberous roots of *C. speciosa*, such as the index compounds maackiain and formononetin, which have made considerable contributions to immunity enhancement, hepatoprotection, arresting cough, and expectorant and anti-asthmatic effects. The quality of *C. speciosa* is mainly evaluated according to the degree of root thickening and the content of isoflavonoids, while the harvest period is primarily depended on the yield. Generally, the fibrous roots of *C. speciosa* possess the potential to form tuberous roots, but limited fibrous roots can transfer into tuberous roots. Studies of root thickening and isoflavonoid biosynthesis of *C. speciosa* are scarce.

Tuber morphogenesis is mainly regulated by the interaction between primary and secondary cambia. The primary cambia usually determine the formation of root/stem, while the secondary cambia affect root/stem thickening by cell division and expansion. The formation of tuber usually involves three common stages: initiation stage, rapid-thickening stage and stable-thickening stage, which is determined mainly by endogenous and environmental factors, such as photoperiod, high sucrose, and water supply. Great efforts have been made to explore the roles of endogenous phytohormones in the initiation and development of tuberous roots. The synergistic actions of various phytohormones, such as auxins, abscisic acid (ABA), gibberellins (GAs), ethylene (ETH), jasmonic acid (JA) and cytokinins (CKs), finally result in the bulking of tuberous roots.

Tuber morphogenesis is a complex biological process involving many specific genes and proteins. Transcriptome techniques have contributed to our understanding of these genes involved in regulation tuberous root formation. In medicinal plant *Rehmannia glutinosa*, transcriptome analysis indicated that 6032 DEGs related to hormone signaling, signal transduction and light signaling were identified during tuberous root formation. In potato transcriptomic data, some of these genes were involved in plant hormone signal transduction, with GID1-like GA receptor (SIGD1) being up-regulated. In sweet potato, several transcriptome analyses revealed that a large number of genes were highly regulated during the storage root formation, which participated in starch and lignin synthesis, cell division, and expansion. In addition, 191 DEGs were found to be involved in functions such as plant growth and development, metabolism, cell organization and biogenesis, signal sensing and transduction, and plant defense response in radish. Sucrose content dominantly decreased at an early stage of bulb development in onion, whereas the fructose and glucose contents increased significantly at the mature stage, suggesting that sucrose metabolism plays an important role in onion bulb formation. The previous studies focused mainly on crops, but the root expansion process in medicinal plants may be different from that in crops. Little is known about the genetic regulation of the initiation and development of tuberous roots in *C. speciosa*.

As a branch of the flavonoid pathway, the biosynthesis of isoflavonoids shares the common upstream pathway with flavonoid biosynthesis under the actions of phenylalanine ammonia-lyase (PAL), cinnamate 4-hydroxylase (C4H), 4-coumarate-CoA ligase (4CL), chalcone synthase (CHS), chalcone isomerase (CHI), and chalcone reductase (CHR). In the downstream pathway, isoflavonoids are synthesized from liquiritigenin under the catalysis by 2-hydroxyisoflavanone dehydratase (HIDM) and isoflavone synthase (IFS), and then modified by various tailoring processes including hydroxylation, methylation, and glycosylation under the actions of other enzymes, such as isoflavone 4′-O-methyltransferase (HI4OMT), vestitone reductase (VR), and cytochrome P450 members. The gene expression analysis of isoflavonoid biosynthesis has been studied by transcriptome analysis in several legume plant species, such as soybean, *Pueraria camollei* and *P. lobata*. However, many genes encoding key enzymes in the isoflavonoid pathway are still not well defined, especially the genes encoding downstream pathway enzymes. A comprehensive understanding of the genes involved in isoflavonoid biosynthesis during the development of tuberous roots in *C. speciosa* will help in revealing the mechanism of isoflavonoid biosynthesis.
Morphological and anatomical analyses were also performed to reveal the secondary meristem activity during the tuberous root thickening. The morphological analysis revealed distinct phenotypic characteristics appeared at different developmental ages. The tuberous root was markedly enlarged from 12 MAG, suggesting that the secondary meristem activity might be activated before that phase (Fig. 2a). The anatomical results also showed that the vascular cambia (Vc), which cued fibrous roots to tuberous roots, appeared before 6 MAG. Thus, the 6 MAG represented a critical point in the transition from fibrous roots to tuberous roots, in which vascular bundles were well developed and secondary cambia initiated (Fig. 2b). After the vascular cambia appeared, several secondary structures, including secondary xylem (Sx) and secondary phloem (Sph), anomalous cambia (Ac), and the scattered tertiary structures (TS), which promoted root thickening, were observed well-developed at 12 MAG and secondary cambia divided continuously at 18 MAG (Fig. 2c,d). Due to the meristematic cells proliferated continuously, secondary phloem cells outward and secondary xylem cells inward were produced, resulting in the secondary xylem much larger than the phloem (Fig. 2e,f). From 24 MAG, accessory cambium (Avc) appeared in the secondary phloem, and the continuous division of these cambium cells led to rapid thickening of roots (Fig. 2f,g). The thickening rate of roots slowed down from 30 MAG and tended to stabilize. Overall, the growth cycle of *C. speciosa* could be divided into three stages: initiation, rapid-thickening and stable-thickening stage.

To explore the molecular mechanism during root thickening, four samples were selected for further research, including initiated thickening stage (6 MAG), early-rapid thickening stage (12 MAG), mid-rapid thickening stage (18 MAG), and stable thickening stage (30 MAG).

**Overview of de novo transcriptome assembly.** Approximately 799.94 Mb original data in total were obtained using BGISEQ-500 platform. After filtering low-quality reads and adaptor sequences, 78.23 Gb clean reads were obtained and processed by de novo analysis using Trinity software. Then, Tgicl software was used on unigenes to remove abundance, and 153,153 unigenes were obtained. The N50, Q30 and gene mapped percentage was 2167 bp, 89.14%, and 88.09%, respectively (Supplementary Table S1). The length distribution of all the assembled transcripts shown in Supplementary Fig. S1a, which indicated that 24.93% of the unigenes were longer than 2000 bp. Functional annotation revealed that 106,882 (69.79%), 100,406 (65.56%), 79,007 (51.59%), 86,688 (56.60%), 86,701 (56.61%), 80,297 (52.43%), 75,729 (49.45%), 45,329 (29.60%) were aligned with eight databases (NR, NT, SwissProt, KEGG, KOG, Pfam, GO, and InterPro) respectively (Supplementary Fig. S1b). Furthermore, the similarity distribution indicated that there were 21.30% of all the unigenes mapped to *Cicer arietinum* by aligning to NR database according to their amino acid sequences, followed by *Cajanus cajan* (15.96%), *Glycine max* (14.56%), belonged to the same Leguminosae family (Supplementary Fig. S1c).
KOG is a database of orthologous gene families, in which 86,701 unigenes were classified to 25 functional classifications. The “signal transduction mechanisms” (13,227 unigenes) were annotated in KOG database (Supplementary Fig. S1d). Moreover, 81,412 unigenes were involved in the KEGG pathways, which were categorized into five functional groups, including metabolism (58.77%), organismal systems (23.64%), environmental information processing (5.15%), genetic information processing and cellular processing (4.99%) (Supplementary Fig. S1e). Carbohydrate metabolism was identified to be the most important pathway during tuberous root development, such as starch and sucrose metabolism, amino sugar and nucleotide sugar metabolism (Supplementary Fig. S1f), while phenylpropanoid biosynthesis was the dominated pathway in biosynthesis of other secondary metabolites (Supplementary Fig. S1g).

Principal component analysis (PCA) showed that the 12 samples could be clearly assigned to four groups. There was a significant difference between 18 and 30 MAG, while 6 MAG and 12 MAG clustered together, which indicated that the transcriptomic profiles were similar between initiated thickening and early-rapid thickening stages while those of mid-rapid thickening and stable thickening stages were distinct (Fig. 3a).

Identification of differentially expressed genes (DEGs) in transcriptome for *C. speciosa*. A total of 4337 DEGs were identified among three pairwise comparisons: 2, 475 DEGs in 6-vs-12, 1926 DEGs in 12-vs-18, and 2134 DEGs in 18-vs-30. More down-regulated DEGs appeared in 6-vs-12 comparison, whereas the opposite occurred in 12-vs-18 and 18-vs-30 comparisons (Fig. 3b). A complete list of the detected DEGs showed detailly in Supplementary Table S2. The Venn diagram of DEGs showed that none were commonly up- or down-regulated in all the pairwise comparisons, implying that the DEGs might be regulated in certain period (Fig. 3c). The clustering analysis of DEGs were also performed by K-means clustering method based on similarities in gene expression profiles. The results showed that DEGs were mainly divided into six clusters, in
which 379 genes were clustered in the up-regulated subclass and 820 genes were highly expressed at the rapid-thickening stages (Supplementary Fig. S2).

For better comprehension of DEGs functions, 48 GO categories were identified. For biological processes, DEGs were involved mainly in “metabolic process” and “cellular process” (Supplementary Fig. S3). Functional analysis by KEGG revealed that DEGs were enriched predominantly in “flavonoid biosynthesis”, “isoflavonoid biosynthesis”, “glycolysis/glucoseogenesis”, and “phenylpropanoid biosynthesis” pathways in the three pairwise comparisons (Supplementary Fig. S4). In particular, “flavonoid biosynthesis” and “isoflavonoid biosynthesis” pathways were significantly enriched in all pairwise comparisons. The “biosynthesis of amino acids” pathway were significantly enriched in two comparisons (12-vs-18 and 18-vs-30), whereas “glycolysis/glucoseogenesis”, “MAPK signaling pathway” and “plant hormone signal transduction” showed significant difference in 12-vs-18 comparison (Fig. 3d). These DEGs might be closely related to tuberous root development and isoflavonoid metabolism in C. speciosa.

Endogenous hormone contents and DEGs related to hormone signaling. To investigate the roles of endogenous hormones in root thickening, we analyzed the auxins, CKs, GAs, JA, and ABA contents in C. speciosa roots by UPLC–MS/MS in the present study (Fig. 4a). The results showed that both IAA and GA3 contents were significantly high at the rapid thickening of tuberous roots, suggesting that high IAA and GA3 levels might promote tuber enlargement. Similarly, cis-zeatin (CZ) as the main cytokinin in C. speciosa tuberous roots, peaked at 18 MAG. By contrast, the highest JA content appeared at 6 MAG, corresponding to the initiation development of tuberous roots, whereas the opposite trend for ABA content was observed at the same age point. GA19 content maintained at a relatively constant level during root thickening. In the corresponding hormone signaling pathways, 36 DEGs were identified among 4337 DEGs (Supplementary Table S3). In auxin transduction pathway, the expression levels of genes encoding AUX/IAA auxin-responsive protein (IAA16s and IAA17), auxin signaling F-box 2 (AFB2-1) and auxin-responsive protein GH3.3 were higher at 12 MAG, while that of small auxin up RNA (SAUR32-1, SAUR71), auxin response factor (ARF2), and AFB2-2 were highly expressed at 18 MAG. The genes encoding JA ZIM-domain protein (TIFYs) and receptor COI1 showed high expression levels at the initiated stage (6 MAG), whereas two ABA receptor PYL genes were highly expressed at the rapid expansion stage (12 and 18 MAG). Most of the DEGs related to GA and CK signaling pathways were highly expressed at the early-rapid expansion stage (12 MAG), such as gene encoding receptor GIhS, chitin-inducible gibberelin-responsive protein 2 (CIGR2), scarecrow-like 13 (SCL13), histidine-containing phosphotransfer protein 1-like (AHP1-like) and cytokinin response regulator (ARR6) (Fig. 4b). These results were consistent with those of endogenous hor-

Figure 3. Global gene expression profiling of C. speciosa roots at different developmental ages. (a) PCA analysis of the RNA-Seq data. (b) Numbers of identified differentially expressed genes (DEGs) in the sequential pairwise comparisons. (c) Venn diagrams of DEGs among the three comparisons. (d) Significantly enriched KEGG pathways of DEGs. The numbers (6, 12, 18, 30) represent the four different developmental ages.
mones, and indicated that these genes might be involved in regulating the hormonal signal transduction during root thickening.

**DEGs related to sucrose and starch metabolism.** The starch content increased significantly from 18 to 30 MAG (Fig. 1b). In the sucrose and starch metabolism pathway, key enzymes maintain the metabolic balance of glucose, fructose, sucrose, and starch/glycogen. Among the 4337 DEGs, 18 genes encoding key enzymes in the sucrose and starch metabolism pathway were identified (Supplementary Table S3). Heatmap analysis revealed that the expression profiles of 18 DEGs obviously differed among the four samples. The expression of gene encoding granule-bound starch synthase 1 (GBSS1) was higher at 18 and 30 MAG, whereas genes encoding sucrose synthase (SUSs) were highly expressed at 12 and 18 MAG. Genes encoding phosphoglucomutase (PGM), glucose-6-phosphate isomerase (GPI), UTP–glucose-1-phosphate uridylyltransferase (UGP), and hexokinase 1 (HXK1) were highly expressed at 12 MAG. Two genes encoding glucan endo-1, 3-beta-glucosidase (eglC) were highly expressed at both 12 MAG and 18 MAG, while the expression of other three genes in the same family were at high levels at 18 MAG. In addition, the expression of two genes encoding beta-amylase (BAMY) differed among different developmental ages of roots, and BAM9 were at a higher level than BAM1 (Fig. 5). Therefore, sugar metabolism pathway might be activated by these metabolic related DEGs at rapid-thickening stage of *C. speciosa*, but the DEGs associated with starch metabolism pathway might play important roles in promoting starch accumulation at stable thickening stage, which was consistent with the trend of starch content.

**DEGs related to isoflavonoid biosynthesis.** The content change of formononetin and maackiain increased gradually during the tuberous root development (Fig. 1c,d). Analysis of the *C. speciosa* transcriptomic data allowed us to identify 15 genes involved in isoflavonoid biosynthesis among 4337 DEGs, including genes encoding 4-coumarate: CoA ligase (4CL), 4-oxoglutarate dioxygenase (H4′OMT), vestitone reductases (VRs), 2-Hydroxyisoflavonone dehydratases (HIDMs), chalcone synthases (CHSs), chalcone-flavonone isomerases (CHIs), isoflavone synthases (IFSs), and isoflavone-3′-hydroxylase (I3′H) (Supplementary Table S3). These DEGs had different expression patterns among the four samples. Two CHSs (CHS1 and CHS7) were both highly expressed at 12 MAG. Like CHSs, CH13 also had highly expression at 12 MAG, but more DEGs showed a low expression level at 12 MAG, such as 4CL, IFSs, I3′H, VRs, and H4′OMT (Fig. 6). However, we speculated that isoflavonoid biosynthesis might be promoted under the complex synergistic effects of these genes during the thickening of *C. speciosa* tuberous roots.

---

**Figure 4.** Trends in endogenous phytohormone contents (a) and expression profiles of hormone related DEGs (b) in *C. speciosa* roots. (a) Trends in endogenous phytohormone contents. (b) The expression profiles of genes encoding hormone signaling. Heat map indicates the log_{10}-transformed FPKM expression values. Changes in expression level are indicated by a change in color; from blue to red indicates an expression level from low to high. All data shown reflect the mean of three biological replicates (*n* = 3). Means with different letters in each sample represent a significant difference at 0.05 level.
DEGs related to MAPK and calcium signaling. The regulation of genes associated with mitogen-activated protein kinase (MAPK) and calcium signaling were also identified from 4337 DEGs in the present study (Supplementary Table S4). Four MAPK signaling-related genes were annotated, including MPK3, MPK7, MPK9, and MAPKKK18. Of these, three genes (MPK3, MPK9 and MAPKKK18) were highly expressed at the early-rapid thickening stage (12 MAG). Calcium-regulated transduction is another key signaling transduction pathway in cell development. A total of 20 DEGs were homologous with calcium signal-related genes, including nine calcium binding protein transcripts (CBPs), four calreticulin transcripts (CBLs), three calcium-transporting ATPase transcripts (CTAs), two calcium-dependent protein kinase transcripts (CDPKs), and two calcium exchanger transcripts (NCLs). It is worth noticing that most of these genes were highly expressed at the initial thickening stage (6 MAG) and early-rapid thickening stage (12 MAG), indicating that genes encoding the calcium-regulated signaling pathway might play vital roles in initiation of root thickening in C. speciosa.

DEGs related to cell wall and cell cycle. Among 4337 DEGs, a total of 34 transcripts associated with cell expansion, cell wall synthesis, cell cytoskeleton and cell cycle regulation were identified during the expansion of tuberous roots, including transcripts encoding xyloglucan endotransglucosylase/hydrolase (XTH), expansin (EXP), extension (EXT), pectinesterase (PE), fructokinase (FRK), cyclin-dependent kinases G-2 (CDKG-2), tubulin beta-2 chain-like (TUB2), cell division cycle protein 48 (CDC48), and cyclin-dependent kinases regulatory subunit (CKS1) (Supplementary Table S5). Most of the genes related to cell expansion, cell wall synthesis, and cell cytoskeleton were highly expressed at the rapid thickening stages (12 and 18 MAG), whereas the genes related to cell cycle regulation showed the opposite trend.

Differentially expressed transcription factors (TFs). A total of 230 TF-encoding genes belonging to 30 TF families were identified from 4337 DEGs during root thickening (Supplementary Table S2). Among them, 41 DEGs belonging to AP2/ERF, MYB, NAC, MADS, bHLH, and WRKY families have previously implicated in plant growth and development (Supplementary Fig. S5). AP2/ERF TF is one of the largest superfamilies in plants, indicating that different members might play diverse roles in various root thickening stages. The majority of ERFs transcripts were highly expressed at 12 MAG and 18 MAG, especially ERF109-like, ERF105 and ERF17-like. Genes belonging to MYB, NAC, and WRKY families were highly expressed at the rapid expansion stage (12 and 18 MAG), whereas 4 bHLHs and 2 MADSs were highly expressed at 18 MAG. Our results suggested that these TFs might potentially participate in the developmental regulation of C. speciosa tuberous root thickening and the transcriptional regulation of isoflavonoid biosynthesis.

Validation of candidate DEGs by qRT-PCR analysis. To validate the reliability of the transcriptomic data, 16 DEGs were selected randomly and their relative expression levels were determined by quantitative real-time PCR (qRT-PCR) analysis (Supplementary Table S6). The expression levels of the selected genes were calculated using 2-ΔΔCq method. We compared the expression data between RNA-seq and qRT-PCR, and the correlation was calculated using log10 fold variation measurements to produce a scatter plot. As shown in Fig. 7a, the
The relative expression levels ($2^{-\Delta\Delta Cq}$) of all the selected genes were significantly consistent with the expression levels determined using the transcriptome data (FPKM values). The Pearson correlation values ($R^2$) between the relative expression levels and FPKM values were 0.28 ($p < 0.05$) (Fig. 7b). Thus, the qRT-PCR analysis confirmed the validity of the transcriptome data, indicating that the transcriptome data were reliable and accurate.

**Discussion**

*Callerya speciosa* (Champ. ex Benth.) Schot is a traditional Chinese medicine characterized by tuberous roots as the main organ of isoflavonoid accumulation. The formation and development of tuberous roots is a complex and genetically programmed process that involves the proliferation of secondary meristems and the accumulation of carbohydrates and secondary metabolites. Based on several growth and content indexes determination in this study, roots saturation with more water occurred at the rapid growth phase, while the content of carbohydrates and secondary metabolites increased at the later period of root thickening (Fig. 1). Morphological and anatomical analyses verified that vascular cambia cued fibrous roots to tuberous roots in *C. speciosa*, which was already well formed at 6 MAG. Secondary xylem and phloem differentiated under the activity of vascular cambia, depending on cell division, expansion and differentiation and resulting in a rapid increase in root diameter (Fig. 2). Therefore, detailed information on gene expression is vital to understand the molecular mechanism of any developmental process. Although *C. speciosa* has great economic value, genetic information for this species remains unavailable. Here, roots at four different key developmental ages of *C. speciosa* were analyzed using an integrated morphological, hormonal and transcriptomic approach. A total of 4337 genes were identified that have shown differential expression during root thickening. Large numbers of DEGs were involved in similar molecular processes during the development of tuberous roots (Fig. 3), a dynamic process controlled by plant hormones, TFs and metabolism regulation pathways.

**Hormonal signaling regulation during root thickening.** The initiation and development of tuberous roots have been shown to be regulated by endogenous phytohormones. The synergistic actions of various phytohormones, such as auxins, ABA, GAs, ETH, JA and CKs, finally result in the bulking of tuberous roots/stems. The present study revealed that significant accumulation of JA appeared at the initial development of tuberous roots, but its content showed gradual decrease during storage root development (Fig. 4a). Previous studies have showed that JA might induce storage root formation by stimulating proliferation of metaxylem and cambium cells in sweet potato. Thus, the high accumulation of JA might play an important role in the initial development of tuberous roots in *C. speciosa*. Jasmonate zinc finger inflorescence meristem (JIM)-domain (JAZ) protein and JA receptor COI1 were reported to form COI1–JAZ complex in JA signaling in cells for the bioactive JA-ILE, subsequently JAZ proteins were degraded after being transferred to the 26S proteasome, and simultaneously, TFs were released to activate the expression of downstream genes. The genes encoding JA...
Panax notoginseng

Similar results were obtained in C. speciosa.

were up-regulated at the cortex splitting and expanding stage24. Researchers suggested that the endogenous IAA ARFs genes and two genes responsible for cambium secondary differentiation during ment. Most transcripts encoding Aux/IAA genes were down-regulated, while those encoding ARFs genes and two SAUR transcripts were up-regulated at the cortex splitting and expanding stage24. Our study showed that five transcripts encoding eglC were highly expressed at the initial thickening stage (6 MAG) (Fig. 5; Supplementary Table S3), implying that they might associate with starch granules.

In general, GA3 and ABA content and genes involved in auxins signaling pathway, including Aux/IAA and ARFs were highly expressed in expansion stage9,25. Congruent with previous studies, 6 Aux/IAA, 4 SAURs, 2 ARFs and 2 CH3s, were highly expressed at the rapid thickening stage (12 and 18 MAG) in this study (Fig. 4b), implying that they might associate with the cell expansion.

Interestingly, the endogenous GA3 content and the expression of some genes related to the GA signaling pathway were significantly enhanced at the rapid-thickening stages, such as GIDs, SCLs and CIGR2 in the present study (Fig. 4). In general, GAs are thought to play different roles in storage root formation and development. In potato, GAs were important promoters in stolon initiation but served as inhibitors of tuber initiation26. Gene encoding GID1-like gibberellin receptor (SIGID1) was significantly up-regulated in the sample pair potato grafted with potato (St-R) vs self-grafted potato (St-WT) during stolon development, which was consistent with GA3 content15. In addition, ABA and GA3 were reported to antagonistically mediate tuberization in several species of plants, such as potato28, sweet potato10 and P. notoginseng25. The ABA levels increased at the rapid-thickening stages of tuberous roots (12 and 18 MAG), whereas the ABA content showed gradual increase along with the process of tuberous root development in this study (Fig. 4a). Therefore, we hypothesized that antagonistic interaction between ABA and GA signaling potentially modulated root thickening in C. speciosa; this supposition was supported by the expression levels of genes related to ABA and GA signal transduction pathway in roots (Fig. 4).

In sweet potato, CKs appeared to be key factors in the formation of storage roots as a prerequisite for cambial cell proliferation27. In this study, cis-zeyatin (c-Z) was the main cytokinin in C. speciosa tuberous roots, and the peak of c-Z content was at the mid-rapid thickening stage (18 MAG) (Fig. 4a). Overall, our result suggested that various phytohormones play roles at different ages of storage root development, which were supported by the previous studies in sweet potato29, radish30 and yam31. Nevertheless, the synergistic effect of phytohormones on C. speciosa tuberous root formation and development warrants further investigation.

Starch and sucrose metabolism regulation during root thickening. We found that "starch and sucrose metabolism" pathway was markedly enriched in carbohydrate metabolism in C. speciosa (Supplementary Fig. S1f), which was consistent with the starch accumulation in the tuberous roots (Fig. 1b). Starch is an important carbohydrate that stored in tuberous roots. The accumulation of starch includes the synthesis, degradation, transport, and conversion of sucrose32. The main photoassimilate sucrose is a promoter that can stimulate the formation of tuberous roots in several plants28. In the starch and sucrose metabolic pathway, sucrose is converted into UDP-glucose and fructose by sucrose synthase, meanwhile it's also converted to glucose and fructose by invertase in the cytosol30. Here, genes encoding ten enzymes in the starch and sucrose metabolic pathway were identified (Fig. 5). Genes related to sucrose metabolism were highly expressed at the rapid thickening stages, e.g., SUS2, SUS-like, HXK1, GPI (Fig. 5; Supplementary Table S3), suggesting that sucrose catabolism is necessary for root rapid-thickening in C. speciosa. Similar results were obtained in P. notoginseng30, Cyprius esculentus30, and radish31. Glucan endo-1,3-beta-glucosidase (elgC) is widely distributed in higher plants, which is responsible for hydrolysis of the glycosidic bond in specific polysaccharides with tracts of unsubstituted 1,3,1,3-linked glucosyl residues to release glucose30. Our study showed that five transcripts encoding elgC were highly expressed at the rapid thickening stages (Fig. 5), which means that they might contribute to glucose supply during root thickening process.

Several enzymes have been reported to be pivotal in starch biosynthesis and composition, including GBSS, SBE, and PGM30. Our transcriptome data showed that GBSSI was highly expressed at the stable thickening stage (30 MAG) (Fig. 5), and it means that starch accumulation in an active sink might play a vital role at the stable thickening stage, which was similar to the result in P. notoginseng30. The activity of beta-amylase (BAMY) could decrease by silencing StBAM1 and StBAM9 in potato, and StBAM9 interacted with StBAM1 on the starch granules30. Genes encoding BAMY were highly expressed at the expansion stage of yam tuberous roots30. In this study, BAM9 was highly expressed at the mid-rapid thickening stage (18 MAG), whereas BAM1 had a higher expression at 6 MAG (Fig. 5; Supplementary Table S3), implying that they might associate with starch granules. These starch and sucrose metabolism genes were required for the root thickening in C. speciosa, which might play vital roles in the thickening of C. speciosa tuberous roots.

Potential DEGs involved in the isoflavonoid biosynthetic pathway. The benefits of isoflavonoid compounds in plants have inspired efforts to identify the structural and regulatory genes in their biosynthetic pathway. Most of the isoflavonoid pathway genes encoding the key enzymes responsible for its biosynthesis have been identified32. Chalcone established the first step in the branched pathway for the synthesis of flavonoids, which was catalyzed by CHS34. It was reported that 9-member CHS family in soybean, in which CHS7 and CHS8 genes played critical roles in regulation of isoflavonoid biosynthesis35. In this study, CHS1 and CHS7 were significantly up-regulated at the rapid thickening stage of tuberous roots (Fig. 6), in which formononetin and maacki-
ain was rapidly accumulated (Fig. 1). CHIs might regulate the efficient metabolic flux of isoflavonoid biosynthesis by interacting with CHS1, IFS1, and CYP93C in soybean. It was reported that genes encoding CHI, CHS, CYP, and isoflavone reductase (IFR) were all involved in the biosynthesis pathway of bioactive isoflavonoids and most abundantly expressed in the summer-collected tubers in Pueraria mirifica. Many isoflavonoids could also be O-methylated at hydroxyl groups by O-methyltransferases (OMTs), and 4′-O-methylated under the action of HI4/OMTs, which was reported to be important for isoflavonoids (e.g., formononetin) biosynthesis in several.

Figure 7. Validation of the transcriptomic data with qRT-PCR. (a) Comparison of the expression levels between qRT-PCR and FPKM values identified by transcriptome. (b) Correlation plot of the qRT-PCR (2−ΔΔCq) and FPKM values. The R² value represents the correlation between the qPCR and RNA-seq results.
leguminous plant species. The key step of pterocarpan phytoalexin (e.g., maackiain) biosynthesis, conversion of vestitone to pterocarpan, proved to be catalyzed by VR. Our results showed that the expression of some genes (e.g., H IDM-3, IFs, I3H, VRs, and H4OMT) was at lower levels at 12 MAG in the present study (Fig. 6), which allowed us to speculate that the tailoring processes, including hydroxylation, methylation, and glycosylation, might be suppressed at the early-rapid thickening stage of C. speciosa. Several members of CYP81E subfamily, which catalyze the hydroxylation of isoflavones, daidzein and formononetin, were identified to characterize the isoflavonoid metabolism in Glycyrhiza echinata (CYP81E1), Medicago truncatula (CYP81E7) and soybean (CYP81E11, CYP81E12 and CYP81E18). In the present study, the transcripts encoding I3H, belonging to CYP81E subfamily, were highly expressed at the 6 and 30 MAG (Fig. 6). Therefore, all data suggested that these DEGs might be potentially involved in the isoflavonoid biosynthesis.

Transcription factors regulation during root thickening. Previous studies have revealed that TFs are important regulators of tuberous root formation and secondary metabolism, such as AP2/ERF, MYB, MYC, and WRKY. In the current study, a total of 230 TF-encoding genes belonging to 30 TF families were identified from 4337 DEGs during root thickening. Among them, 41 TF genes, belonging to AP2/ERF, MYB, NAC, MADS, bHLH, and WRKY families, have previously implicated in plant growth and development (Supplementary Fig. S5). In Arabidopsis, ERF109 was reported to positively control root stem cell niche maintenance and root growth through phytosulphokine (PSK) peptide hormones, and mediate the cross-talk between JA signaling and IAA biosynthesis to regulate root development. In this study, a gene homologous to AtERF109 was highly expressed at early-rapid thickening stage (Supplementary Fig. S5). Recent studies have further demonstrated that JA-responsive ERF109 transcriptional regulation initiated the transition from seed roots to taproot thickening in P. notoginseng. It’s unsurprising that endogenous JA content was dominantly increased in the early stage of tuberous root development and genes related to JA signaling were at high expression levels at the same time (Fig. 4), indicating that JA-induced ERF transcriptional regulation controlled root thickening in C. speciosa. Moreover, several transcription factors including MYB (MYB44s), bHLH (bHLH35 and MYC2), WRKY (e.g., WRKY41), and NAC (e.g., NAC2) were highly expressed at the rapid thickening stages (Supplementary Fig. S5). The bHLH transcription factors MYC3 and MYC4 were targets of JAZ repressors and acted additively with MYC2 in the activation of jasmonate responses. MYB44, a tuber-specific and sucrose-inducible element-binding factor in potato, was reported to positively regulate the enhanced elongation of primary roots by inducing N-3-oxo-hexanoyl-homoserine lactone and negatively regulate anthocyanin biosynthesis at high temperatures in tuber flesh of potato. An increasing number of data indicated that some WRKY genes (e.g., WRKY41) were responsive to phytohormones, such as ABA, GA and JA, to regulate the lateral root formation. NAC TF family was reported to function in regulating secondary xylem development, secondary cell wall metabolism and cell expansion. Furthermore, some up-regulated transcripts in this study, e.g., MYB44s, bHLH35, and WRKY41, which were reported to regulate the anthocyanin biosynthesis, might play vital roles in the regulation of isoflavonoid biosynthesis. All these studies suggested that these TF genes might be potentially involved in the root thickening and isoflavonoid metabolism of C. speciosa.

Regulatory model associated with tuber thickening and isoflavonoid biosynthesis of C. speciosa. The initiation and development of storage roots is a highly complicated process that multiple pathways regulate the development of secondary anomalous cambia and the accumulation of starch, sucrose and other chemical components. In this study, a potential regulatory model associated with the root thickening and isoflavonoid biosynthesis of C. speciosa was proposed according to our integrated phenotypic and transcriptomic data (Fig. 8), which would help in understanding the underlying mechanisms of tuberous root formation and isoflavonoid biosynthesis in C. speciosa. Cell divisions and expansions in the vascular cambia, together with differentiation of secondary xylem and phloem under the activity of vascular cambia, led to an expansion of roots. The cell proliferation through several signal transduction pathways (calcium, MAPK, hormone and transcription signaling) and metabolism possesses (cell wall, sucrose, starch and isoflavonoid metabolism). Several genes were highly expressed to promote cell differentiation, division, expansion and dry matter accumulation at the secondary pathway. In detail, CBPs, CBLs, CDPKs, NCLs, and CTAs were involved in calcium signaling pathway, a primary essential signaling transduction category required for normal growth and development of plants. The MAPK signal pathway was reported to play a role in regulating the hormone signaling and developmental processes, and four MAPK signaling-related genes were differentially expressed, including MPK3, MPK7, MPK9 and MAPKKK18. EXPs, ETRs, XTHs, PEs, CDCs, CDKs, and FRks were reported to regulate the cell extension, expansion, cell wall synthesis, and cell cycle regulation. The hormone signaling are critical factors for regulating root tuberization and isoflavonoid metabolism at different developmental stages, including synergistic and antagonistic regulation. Most genes involved in hormone signal transduction pathways were up-regulated at the rapid thickening stages of tuberous roots. JA might induce storage root formation by stimulating proliferation of metaxylem and cambium cells. MYB44s, ERF109, WRKYs, NACs were involved in enhancing the formation and elongation of roots. In addition, MYB44s, bHLH35, and WRKY41 were also reported to regulate the anthocyanin biosynthesis, which might be involved in isoflavonoid biosynthesis. Genes encoding GBSS, SUS, PGM, UGP, and BAMY, were reported to be associated with sucrose hydrolyzation and starch synthesis during tuberous root thickening in several species, while genes encoding CHI, CHS, HIDM, H4OMT, VR, IFs and I3H were found to be involved in the isoflavonoid metabolism pathways. Taken together, the results indicated that all these DEGs potentially involved in the regulatory model of root thickening and isoflavonoid biosynthesis in C. speciosa. Further functional identification studies were needed to confirm these potential genes.
Conclusions
Integrated morphological, hormonal and transcriptomic analyses were performed in the present study, and multiple tuberous root development and isoflavonoid biosynthesis associated events, including cambium development, endogenous hormones change and corresponding candidate genes were revealed. JA might be related to the initial development of tuberous roots. ABA seemed to be essential for tuber maturation, whereas IAA, cZ and GA₃ were correlated to rapid thickening of tuberous roots. Functional analysis showed that 15 DEGs participated in isoflavonoid biosynthesis, while 153 DEGs that involved in starch/sucrose metabolism, hormonal signaling, transcriptional regulation and cell wall metabolism, were identified to potentially control root thickening. A hypothetical model of genetic regulation associated with root thickening and isoflavonoid biosynthesis in *C. speciosa* is proposed, which will be valuable for further research and will help in understanding the underlying mechanism of tuberous root formation and isoflavonoid biosynthesis in *C. speciosa*.

Methods
Plant material and growth condition. Seeds of *C. speciosa* (diploid) were incubated in wet perlite under dark conditions at 26 ± 2 °C until germination (approximately 4 days). Two days after germination, seedlings were cultured in a greenhouse equipped with a system for monitoring temperature (25 ± 4 °C), relative humidity (60 ± 5%) and natural photoperiod in the Guangxi Botanical Garden of Medicinal Plants, with the same cultivation measure. The voucher specimens were deposited in Guangxi Botanical Garden of Medicinal Plants (No. 12110401). Tuberous roots were collected at half-year intervals, including 6, 12, 18, 24, 30, 36 months after germination (MAG), and used for biological, anatomical and chemical analysis. Roots at four developmental ages (6, 12, 18, 30 MAG) were used for endogenous hormones and RNA-Seq analyses. Ten roots from ten plants were selected randomly from every biological replicate, except for thirty plants at 6 MAG. Then, the thickest section in the middle (5 cm long) of each root, was washed with distilled water, cut down into pieces, and mixed to establish a biological replicate. Each sampling point had three biological replicates. The samples used for transverse paraffin sections were cut into 2-mm-thick slices after collection and immediately fixed in 10 mL mixture [formalin: acetic acid: 70% alcohol (FAA) = 5:5:90] for more than 12 h, then stored at 70% alcohol. For RNA purification and hormone detection, samples were immediately frozen in liquid nitrogen after collection and stored at − 80 °C until further use. For content detection, the samples were dried in an oven at 60 °C.

Determination of growth indexes, starch and index compound content. The volume of fresh roots was calculated using the water displacement method. The fresh roots were weighed using an electronic balance, then dried to achieve constant weight in an oven at 80 °C. Starch content was determined according to Firon et al. Starch content was determined according to Firon et al. The content of formmononetin and maackiain were detected following by Chen et al.
RNA library construction and transcriptome sequencing. Total RNA was isolated from roots by following the method described by Liu et al. For each sample, equal amount (20 μg) of total RNA (RIN > 8.0) was applied to construct cDNA library. Briefly, the Oligo (dT) selection was used to enrich the poly (A) mRNAs. The mRNA was fragmented and reversed transcription to double-strand cDNA (disDNA) by N6 random primer. The synthesized cDNA was subjected to end-repair and then 3’ adenylated. Adaptors were ligated to the ends of these 3’ adenylated cDNA fragments. The ligation products were purified and several rounds of PCR amplification were performed to enrich the purified cDNA template. The PCR product was denatured by heat and the single strand DNA was cyclized by splint oligo and DNA ligase. Finally, twelve libraries (three replicates per sample) were sequenced in pair-end (2 × 100 bp) format BGISEQ-500 sequencing platform, reads were processed and de novo assembled, and the results were analyzed by the BGI Tech Solutions Co., Ltd. (BGI Tech) (BGI, Shenzhen, P. R. China).

Transcriptome profiling and DEGs analysis. Adaptor sequences (reads with ambiguous bases ‘N’) and low-quality reads (reads having more than 20% of bases with quality ≤ 15) were filtered. The filtered clean reads (FASTQ formatted files) for samples can be accessed in the NCBI Sequence Read Archive (SRA) database (https://trace.ncbi.nlm.nih.gov/Traces/sra/) under the accession number of SUB6340191. Trinity software (v2.4.0) was used to carry out the de novo assembly. Tgicl software (v2.0.6) was used to remove the redundant Trinity generated contigs. The unigenes were divided to two types, one type was cluster, which the prefix was CL with the cluster id behind it (in one cluster, there were several unigenes which similarity between them was more than 70%), another type was singleton, which the prefix was unigene. The resulting transcripts were then processed to retrieve associated GO items by BLAST2GO software (v2.5.0). Besides, functional databases NT, NR, KOG, KEGG, SwissProt, Pfam and InterPro were used to annotate genes function, while Blastin, Blastx, and InterProScan 5 were used to align genes. Differential expression of transcripts in different samples were also evaluated by evaluated by fragments per kilobase of transcript per million mapped reads (FPKM) in order to normalize the calculation of gene expression. The differential expression of transcripts was compared by using the DEGseq R package (1.18.0), with a threshold of FPKM ≥ 10 in each pairwise comparison, |Log2FC (Fold Change)|≥ 1 and p value < 0.05. In addition, all the DEGs were mapped in GO and KEGG databases to perform GO enrichment and pathway analysis using GOseq R package. Enrichment analysis of KEGG pathways of DEGs was done by KOBASE software. Multiple testing corrections were performed by controlling the false discovery rate (FDR) to less than 0.05. The enrichment analysis was tested by Fisher’s exact test. PlantTFDB (Plant Transcription Factor Database) was used to annotate TFs.

qRT-PCR analysis. To validate the expression patterns of candidate genes in the different samples, qRT-PCR experiment was performed, as previously described. The forward and reverse primers of sixteen randomly selected DEGs and reference gene were designed using Primer 5.0 (Supplementary Table S6) and synthesized by BGI-Shenzhen. Every candidate gene in each sample was conducted in three biological replicates with three technical replicates. Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was used as a reference gene. The relative expression of each candidate gene was calculated by the 2−ΔΔCq formula. The normalized values of relative expression and FPKM values were calculated using log10 fold variation measurements, and the correlation between RNA-seq and qPCR results was analyzed using these values.

Determination of endogenous hormone contents. Plant materials (120 mg fresh weight) were added to liquid nitrogen, ground into powder, and extracted with methanol/water (8/2) at 4 °C. The extract was centrifuged at 12000 g under 4 °C for 15 min. The supernatant was collected, evaporated to dryness under nitrogen gas stream, then reconstituted in methanol/water (3/7). After centrifuged, the supernatant was collected for UPLC-MS/MS analysis to determine phytohormones content, including auxin, CKs, GAs, JA, and ABA, which were performed as described by Pan et al.

Statistical analyses. Three repetitions were performed to determine each value and their standard deviations (SD). Data were analyzed using IBM SPSS Statistics 19.0 software (Ehningen, Germany) and presented as the means ± SD. The statistical significance was determined using Duncan’s multiple range test. Values in figures marked with different lowercase were significantly different at 0.05 probability levels.

Data availability
The datasets generated for this study can be found in the NCBI SRA repository, https://submit.ncbi.nlm.nih.gov/sra/, with the GenBank accession no.: SUB6340191.

Received: 22 July 2020; Accepted: 27 October 2020
Published online: 08 January 2021
References

1. Wang, H. et al. Altered phenylpropanoid metabolism in the maize Lc-expressed sweet potato (Ipomoea batatas) affects storage root development. Sci. Rep. 6, 18645 (2016).

2. Chen, Y. et al. Determination of maackiaiin and formononetin in root of Millettia speciosa by HPLC. World Sci. Technol. Modern. Trad. Chin. Med. Mater. Med. 15, 260–264 (2013).

3. Yao, S. et al. Hair root induction and polysaccharide production of medicinal plant Callerya speciosa Champ. Plant Cell Tissue Organ Cult. 126, 177–186 (2016).

4. Xu, L. et al. Transcriptome analysis of storage roots and fibrous roots of the traditional medicinal herb Callerya speciosa (Champ) ScHot. PLoS One 11, e0160338. https://doi.org/10.1371/journal.pone.0160338 (2016).

5. Li, M. et al. Analysis of integrated multiple "omics" datasets reveals the mechanisms of initiation and determination in the formation of tuberous roots in Rehmannia glutinosa. J. Exp. Bot. 66, 5837–5851. https://doi.org/10.1093/jxberv288 (2015).

6. Sun, P. et al. Dynamic transcriptional profiling provides insights into tuberous root development in Rehmannia glutinosa. Front. Plant Sci. https://doi.org/10.3389/fpls.2015.00396 (2015).

7. Mortley, D. et al. Elevated carbon dioxide influences yield and photosynthetic responses of hydropically-grown [correction of grown] sweetpotato. Acta Hort. 440, 31–36. https://doi.org/10.17660/actahortic.1996.440.6 (1996).

8. Kondhare, K. R., Malankar, N. N., Devani, R. S. & Banerjee, A. K. Genome-wide transcriptome analysis reveals small RNA profiles involved in early stages of stolon-to-tuber transitions in potato under photoperiodic conditions. BMC Plant Biol. 18, 284. https://doi.org/10.1186/s12870-018-1501-4 (2018).

9. Van Heerden, P. D. & Laurie, R. Effects of prolonged restriction in water supply on photosynthesis, shoot development and storage root yield in sweet potato. Physiol. Plant. 134, 99–101. https://doi.org/10.1111/1399-3054.2008.11111.x (2008).

10. Dong, T. et al. RNA-Seq and iTRAQ reveal multiple pathways involved in storage root formation and development in sweet potato (Ipomoea batatas L.). BMC Plant Biol. 19, 136 (2019).

11. Grandellis, C. et al. SCiDPK3 phosphorylates in vitro two transcription factors involved in GA and ABA signaling in potato: SIARG1 and SIABF1. PLoS One 11, e0167389. https://doi.org/10.1371/journal.pone.0167389 (2016).

12. Zhang, G. et al. Comprehensive transcription profiling and phenotyping of rootstock and scion in a tomato/potato heterografting system. Physiol. Plant. 166, 833–847. https://doi.org/10.1111/pp.12858 (2019).

13. Firon, N. et al. Transcriptional profiling of sweetpotato (Ipomoea batatas) roots indicates down-regulation of lignin biosynthesis and up-regulation of starch biosynthesis at an early stage of storage root formation. BMC Genom. 14, 460–460 (2013).

14. Yu, R. et al. Transcriptome profiling of root microRNAs reveals novel insights into taproot thickening in radish (Raphanus sativus L.). BMC Plant Biol. 15, 30. https://doi.org/10.1186/s12870-015-0427-3 (2015).

15. Zhang, C. et al. Transcriptome analysis of sucrose metabolism during bulb swelling and development in onion (Allium cepa L.). Front. Plant Sci. 7, 1425. https://doi.org/10.3389/fpls.2016.01425 (2016).

16. Wang, X., Li, S., Li, J. & Zhang, Y. D. novo transcriptome sequencing in Pueraria lobata to identify putative genes involved in isoflavone biosynthesis. Plant Cell Rep. 34, 733–743. https://doi.org/10.1007/s00299-014-1733-1 (2015).

17. Uchida, K., Akashi, T. & Aoki, T. The missing link in leguminous pterocarpan biosynthesis is a dirigent domain-containing protein involved in early stages of stolon-to-tuber transitions in potato under photoperiodic conditions. BMC Plant Biol. 18, 284. https://doi.org/10.1186/s12870-018-1501-4 (2018).
38. Li, J. et al. An alternative pathway for formononetin biosynthesis in Psoraria lobata. Front. Plant Sci. 7, 861. https://doi.org/10.3389/fpls.2016.00861 (2016).

39. Akashi, T., Aoki, T. & Ayabe, S. CYP81E1, a cytochrome P450 cDNA of licorice (Glycyrrhiza echinata L.), encodes isoform 2′-hydroxylase. Biochem. Biophys. Res. Commun. 251, 67–70. https://doi.org/10.1016/bbr.1998.04.11 (1998).

40. Liu, C., Huhman, D., Sumner, L. & Dixon, R. Regiospecific hydroxylation of isoflavones by cytochrome p450 81E enzymes from Medicago truncatula. Plant J. Cell Mol. Biol. 36, 20 (2003).

41. Kong, X. et al. PHB3 maintains root stem cell niche identity through ROS-responsive AP2/ERF transcription factors in Arabidopsis. Cell Rep. 22, 1350–1363. https://doi.org/10.1016/celrep.2017.12.105 (2018).

42. Cai, X. T. et al. Arabidopsis ERF109 mediates cross-talk between jasmonic acid and auxin biosynthesis during lateral root formation. Nat. Commun. 5, 5833. https://doi.org/10.1038/ncomms6833 (2014).

43. Fernandez-Calvo, P. et al. The Arabidopsis BHLH transcription factors MYC3 and MYC4 are targets of IAZ repressors and act additively with MYC2 in the activation of jasmonate responses. Plant Cell 23, 701–715. https://doi.org/10.1016/j.pbc.2010.08.078 (2011).

44. Zhou, X. et al. SmMYB44 negatively regulates phosphate transport by suppressing expression of PHOSPHATE1 in potato. J. Exp. Bot. 68, 1265–1281. https://doi.org/10.1093/jxb/erx026 (2017).

45. Zhao, Q. et al. AtMYB44 positively regulates the enhanced elongation of primary roots induced by N-3-Oxo-hexanoyl-homoserine lactone in Arabidopsis thaliana. Mol. Plant Microbe Interact. 29, 777 (2016).

46. Liu, Y. et al. SmMYB44 negatively regulates anthocyanin biosynthesis at high temperatures in tuber flesh of potato. J. Exp. Bot. 70, 3809–3824. https://doi.org/10.1093/jxb/erz194 (2019).

47. Ding, Z. J. et al. WRKY41 controls Arabidopsis seed dormancy via direct regulation of ABI3 transcript levels not downstream of ABA. Plant J. Cell Mol. Biol. 79, 810–823. https://doi.org/10.1111/tjp.12597 (2014).

48. Hu, Z. et al. TaWRKY51 promotes lateral root formation through negative regulation of ethylene biosynthesis in wheat (Triticum aestivum L.). Plant J. Cell. Mol. Biol. 96, 372–388. https://doi.org/10.1111/j.1365-313X.2018 (2018).

49. Jhang, Y. & Yu, D. The WRKY57 transcription factor affects the expression of jasmonate ZIM-domain genes transcriptionally to compromise Botrytis cinerea resistance. Plant Physiol. 171, 2771–2782. https://doi.org/10.1104/pp.16.00747 (2016).

50. Han, Q. et al. Identification and expression pattern of one stress-sensitive NAC gene from Solanum lycopersicum. Mol. Biol. Rep. 39, 1713–1720. https://doi.org/10.1007/s11303-011-0911-2 (2012).

51. Zhang, S. et al. Transcriptome analysis revealed expression of genes related to anthocyanin biosynthesis in eggplant (Solanum melongena L.) under high-temperature stress. BMC Plant Biol. 19, 387. https://doi.org/10.1186/s12870-019-1960-2 (2019).

52. Duan, S. et al. Functional characterization of a heterologously expressed Brassica napus WRKY41-1 transcription factor in regulating anthocyanin biosynthesis in Arabidopsis thaliana. Plant Sci. 268, 47–53. https://doi.org/10.1016/j.plantsci.2017.12.010 (2018).

53. Nookaraju, A. et al. Role of Ca2+–mediated signaling in potato tuberization: An overview. Bot. Stud. 53, 177–189 (2012).

54. Mingo-Castel, A. M., Negm, F. B. & Smith, O. E. Effect of carbon dioxide and ethylene on tuberization of isolated potato stolons cultured in vitro. Plant Physiol. 153, 798–801. https://doi.org/10.1104/pp.153.6.798 (1974).

55. Yao, S., Huang, W., Pan, C., Zhan, J. & He, L. Caspase-like proteases regulate aluminum-induced programmed cell death in peanut. Plant Cell Tissue Organ. Cult. 121, 691–703. https://doi.org/10.1007/s11240-016-1064-8 (2016).

56. Liu, X. et al. Comprehensive transcriptome analysis reveals genes in response to water deficit in the leaves of Saccharum narenga (Nees ex Steud) Hack. BMC Plant Biol. 18, 250–266 (2018).

57. Wei, S. et al. Transcriptome analysis of Taxillus chinensis (DC) Dansser seeds in response to water loss. PLoS One https://doi.org/10.1371/journal.pone.0169177 (2017).

58. Pertea, G. et al. TIGR Gene Indices clustering tools (TGICL): A software system for fast clustering of large EST datasets. Bioinformatics 19, 631–652. https://doi.org/10.1093/bioinformatics/btg034 (2003).

59. Conesa, A. et al. Blast2GO: A universal tool for annotation, visualization and analysis in functional genomics research. Bioinformatics 21, 3674–3676. https://doi.org/10.1093/bioinformatics/bti610 (2005).

60. Wang, L., Feng, Z., Wang, X. & Zhang, X. DEGseq: An R package for identifying differentially expressed genes from RNA-seq data. Bioinformatics 26, 136–138. https://doi.org/10.1093/bioinformatics/btp612 (2010).

61. Kashyap, S. P., Prasanna, H. C., Kumari, N., Mishra, P. & Singh, B. Understanding salt tolerance mechanism using transcriptome profiling and de novo assembly of wild tomato Solanum chilense (Solanaceae) by liquid chromatography-electrospray tandem mass spectrometry. Phytochemistry 69, 1773–1781. https://doi.org/10.1016/j.phytochem.2008.02.008 (2008).

Author contributions
S.Y. and C.P. designed the research; S.Y., Z.L. and J.G. performed the sample preparation, total RNA extraction, anatomical and physiological data collection; S.Y., R.H. and C.P. performed the de novo transcriptome and statistical analysis; S.Y. and C.P. wrote the manuscript; D.H., R.H. and Y.T. revised the manuscript; all authors read and approved the final manuscript.

Funding
This work was supported by the National Science Foundation of China (81760689, 82060685 and 31801451), the National Science Foundation of Guangxi (2018GXNSFAA281287), Science and Technology Major Project of Guangxi (AA17204056), Key Research and Development Program of Guangxi (AB16380210), Youth Innovation research team project of Guangxi University of traditional Chinese medicine (2018QT001), the Program of Key Lab of Zhuang Yao Medicine (20-065-14), and the Training Program for 1000 Young and Middle-aged Backbone Teachers of Guangxi Higher Education Institution in 2020.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary information is available for this paper at https://doi.org/10.1038/s41598-020-76633-x.
