Genome sequencing reveals chromosome fusion and extensive expansion of genes related to secondary metabolism in Artemisia argyi

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Introduction

Artemisia is a large plant genus in the Asteraceae family that comprises approximately 500 species and subspecies (Bora and Sharma, 2011; Lee et al., 2006). These species are mainly distributed in the temperate northern hemisphere regions (Naö and Effert, 2018) and are widely used in various fields such as herb, food, cosmetics, spices, forage and ornamentals (Torrell et al., 2003). In 2015, the discovery of artemisinin, an antimalarial ingredient isolated from Artemisia annua, won the Nobel Prize in Physiology or Medicine, drawing global attention to study other species of the genus Artemisia (Effert et al., 2015; Su and Miller, 2015).

Previous cytogenetic studies have contributed to the knowledge of the systematic and evolutionary relationship within the Artemisia species. Three basic chromosome numbers were reported in the genus based on numerous chromosome counts from approximately 373 taxa; x = 9 is the most common (85.6%), and x = 8 is less frequent (9.7%). Both basic chromosome numbers exhibited polyploid series, with known levels up to 16 x for x = 9 and hexaploid for x = 8. In addition, a chromosome number of 2 n = 34 occurs in a few species, such as Artemisia vulgaris, Artemisia rubipes and Artemisia argyi (Hoshi et al., 2003), suggesting that a third base number x = 17 may exist. The diversity of chromosome number and polyploidy level results in a 7.4-fold variation in Artemisia genome size, from 4.11 Gb of A. dolosa (2 n = 2 x = 18) to 30.45 Gb of A. medioxima (2 n = 16 x = 144) (Pellicer et al., 2010). The chromosome numbers observed in Artemisia species suggest that, in addition to polyploidization, variation in basic chromosome numbers may also play an important role in the evolution of the genus. Chromosomal fusion and fission are considered the predominant causes for the evolution of basic chromosome numbers in animal and plant kingdoms. For example, the origin of human chromosome 2 was derived from head-to-head fusion of two ancestral ape chromosomes (Baldini et al., 1991). Chromosome fusion affects genetic diversity and environmental adaptation of Heliconius (Ciccioniard et al., 2021). Large-scale chromosomal fission/fusion events promote the speciation of the wild Morus notabilis (x = 6) and the cultivated Morus alba (x = 14) (Xuan et al., 2022). There are far more examples of chromosome fusion that could be discussed. In Artemisia, fluorescence-banded karyotypes of A.
Artemisia vulgaris provide some evidence that a centric (Robertsonian) chromosome fusion may also occur and cause the reduction of its basic chromosome number from $x = 9$ to $x = 8$ (Xirau and Siljak-Yakovlev, 1997). However, despite a wide knowledge of cytological studies on Artemisia, the role of chromosome fusion in basic chromosome numbers variation of Artemisia has not been fully verified and highly valued.

Among the Artemisia genus, A. argyi (also called ‘Chinese mugwort’) is one of the well-known species and is widely distributed in Asian countries, such as China, Korea and Japan (Mei et al., 2016). The dried leaves of A. argyi, known in Chinese as ‘Aiye’, have been used as TCM (traditional Chinese medicine) for about 3000 years (Lv et al., 2018; Song et al., 2019). A. argyi was first recorded in ‘Shi jing’ (a famous China classical literature) near 1100 BC and was first recognized as medicine in ‘Wu Shi Er Bing Fang’ in the Han Dynasty (A.D. 220). In addition, the medical applications of A. argyi were also listed in many other classic clinical and medical literature such as ‘Huang Di Nei Jing’, ‘Ming Yi Bie Lu’, ‘Jin Gui Yao Lve’ and ‘Ben Cao Gang Mu’ (Li, 1957; Mawangdui Han Dynasty Tomb bamboo books research group, 1979; Zhang, 1997). In the long-term practice of traditional Chinese medicine, A. argyi is believed to have the properties of bitterness, warmth and pungency and has the effects of dispelling cold and dampness, warming menstruation, haemostasis, and preventing abortion (Chinese Pharmacopoeia, 1979). In the long-term practice of traditional Chinese medicine, A. argyi is believed to have the properties of bitterness, warmth and pungency and has the effects of dispelling cold and dampness, warming menstruation, haemostasis, and preventing abortion (Chinese Pharmacopoeia Commission, 2020). Recent pharmacological studies have demonstrated that A. argyi also exhibits anti-inflammatory (Yun et al., 2016; Zimmermann-Klemd et al., 2020), anti-allergic (Lv et al., 2018), antimicrobial (Paging et al., 2016), antioxidant (Kim et al., 2015) and antitumour (Seo et al., 2015) activities. Clearly, A. argyi has a long history of application and is still widely used in clinical practice. In 2020, the annual value of the A. argyi market was over 40 billion RMB, making it the largest herbal medicine industry chain in China. In addition, acupuncture and moxibustion (world-renowned medicinal products derived from A. argyi) were recognized as the World Intangible Cultural Heritage in 2010, and traditional Chinese medicine/therapy (including moxibustion) has been recommended by WHO (World Health Organization) in 2019. Therefore, acupuncture and moxibustion have successfully spread worldwide as a reliable alternative therapy for multiple diseases. As the major source of moxibustion, A. argyi is also becoming increasingly popular worldwide.

These amazing economic and medicinal values of A. argyi are due to the large number of secondary metabolites in its leaves, which include volatile oils, flavonoids, terpenoids, phenolic acids and other compounds (Song et al., 2019). More than 100 nature metabolites have been identified in A. argyi volatile oil, primarily comprising monoterpenes, sesquiterpenes and their derivatives (Guan et al., 2019). These metabolites contribute to the aromatic odours of A. argyi and pharmacological activities against asthma, eczema and cough (Du et al., 2021; Ge et al., 2016). A. argyi leaves are also rich in flavonoids, including flavonoids, flavanols, flavonols and chalcone (Lv et al., 2018). Among them, eupatin, jaceosidin, apigenin, luteolin, quer cetin, and naringin are representative components and have been proven to have biological activities for preventing oxidative damage, inflammation, allergies and tumours (Maleki et al., 2019; Nabavi et al., 2015; Serafini et al., 2010). Remarkably, eupatin has been confirmed as the pharmacodynamic component of Stillen® (DA-9601) which has been approved as a phytotherapy for gastritis in Korea. Thereby, it is necessary to investigate the biosynthesis pathways of these active secondary metabolites in A. argyi.

However, despite the huge economic and medical value, A. argyi’s evolution and the molecular basis of the biosynthesis of the abundant active ingredients are rarely reported due to the lack of a high-quality reference genome. Here, we constructed a chromosome-level genome of A. argyi through an integrative approach combining PacBio sequencing (SMRT sequencing high fidelity, HiFi) and high-throughput chromatin conformation capture (Hi-C) technology. In addition, whole-genome duplication (WGD) events and expansion and contraction of gene families in the A. argyi genome were also investigated through phylogenetic and comparative genomic analysis. Furthermore, pivotal candidate genes involved in the biosynthesis of terpenoids and flavonoids were also identified based on genomic and transcriptomic analyses. Briefly, as the first chromosome-level genome in Artemisia, this reference genome will provide valuable resources for exploring the genetic and evolutionary biology of A. argyi and other Artemisia species.

Results

Artemisia argyi genome sequencing, assembly and annotation

Qichun (Hubei Province) is the authentic production area of A. argyi in China. Based on preliminary resource evaluation, a highly volatile oil- and flavonoid-producing A. argyi cultivar, ‘Xiang Ai’ from Qichun, was selected for de novo genome sequencing and assembly (Figure 1a). The somatic cells of A. argyi contained 34 chromosomes by the cytological observation method (Figure 1b) and the genome size was approximately 7.44 Gb by flow cytometry estimation (Figure S1). A 21-mer analysis of genome survey sequencing shows that A. argyi is a tetraploid, with a monoploid genome size of ~1.96 Gb and a whole-genome size of ~7.84 Gb (Figure S2). Compared to other genome-sequenced species in Asteraceae (Erigeron breviscapus, Helianthus annuus, Lactuca sativa, Artemisia annua, Cynara cardunculus, Conyza canadensis, Chrysanthemum nankinense, Chrysanthemum seticuspe, Carlhamus tinctorius, Mikania micrantha, Tanacetum cinerarifolium, Taraxacum kok-saghyz Rodin) (Badoun et al., 2017; Lin et al., 2018; Liu et al., 2020; Peng et al., 2014; Reyes-Chin-Wo et al., 2017; Scaglione et al., 2016; Shen et al., 2018; Song et al., 2018; Wu et al., 2021; Yamashiro et al., 2019; Yang et al., 2017), A. argyi features the largest and most complex genome. It was also estimated that the genome of A. argyi had a relatively high heterozygosity (2.36%) and a large proportion of repetitive sequences (75.86%) (Figure S2, Table S1), which increased the challenge of de novo assembly of this genome.

To overcome the assembly challenging of the A. argyi genome caused by polyplody, high heterozygosity and repetitive sequences, an integrated strategy was adopted by combining Illumina short paired-end reads, PacBio long high-fidelity (HiFi) reads, and Hi-C sequencing (Figure S3). A total of 161.3 Gb of PacBio circular consensus sequencing (CCS) reads with an average length of 15 154 bp and 19-fold whole-genome coverage was obtained from 7 flow cells of the PacBio Sequel II platform (Table S2). These CCS reads were assembled into an initial genome with a total length of approximately 8.03 Gb, containing 10 274 contigs, with an N50 of 8.32 Mb and a
longest contig of 43.52 Mb (Table S3) by Hifiasm assembly (Cheng et al., 2021). Subsequently, a total of 407 Gb clean Hi-C paired-end reads were used for scaffold extension and chromosome mounting (Table S4). With the assistance of Hi-C sequence data, the assembled contigs were anchored to 34 super-scaffolds, which covered 91.4% (7.34 Gb) of the size of the assembled genome (Tables S5 and S4). In summary, the chromosome-level A. argyi genome assembly has a total size of approximately 8.03 Gb, containing 12 449 scaffolds, with a scaffold N50 size of 206.40 Mb and a contig N50 of 6.25 Mb (Table 1 and Figure 2a).

To assess the completeness of the assembly of A. argyi genome, the Illumina short reads and PacBio isoform sequencing (Iso-Seq) data were aligned to the assembled genome, resulting in high mapping rates of 99.89% and 99.70%, respectively (Table S6). Furthermore, benchmarking universal single-copy orthologue (BUSCO) analysis (Simao et al., 2015) was also employed to assess the quality of the assembly. The obtained results showed that 95.5% (2221 out of 2236 BUSCOs) of the BUSCOs were completely present in the A. argyi genome (2192 of the 2236 BUSCO genes were complete) (Table S7). In short, these results demonstrated that the assembled A. argyi genome had high completeness.

We further applied a combination of ab initio and homology-based approaches to identify the repetitive sequences. A total of 73.59% of the assembly was identified as repetitive sequences, including 1.37% DNA transposons, 1.29% interspersed nuclear elements (LINEs) and 39.08% long terminal repeats (LTRs) (Table S8). In addition, higher ratio of Gypsy than Copia elements was observed, each accounting for 24.47% and 13.80% of the genome, respectively, which is similar to the genomic characteristics of other Asteraceae species such as sunflower, stevia, and lettuce. Next, combined with RNA-seq and full-length transcriptome data generated from seven different tissues and organs (root, rhizome, stem and leaf A-D) (Data S1), a total of 279 294 high-quality protein-coding genes were annotated for this tetraploid-resolved genome. We attempted to separate the subgenomes by using K-mers to examine the potential bias in genome characteristics. However, we failed to distinguish homologous chromosome pairs into distinct A and B subgenomes using the enrichment pattern of K-mers. Therefore, we selected the longest chromosome from each pair of homologous chromosomes as one set of chromosomes of A. argyi, except for chromosome 10, because it was fused from chromosomes 8 and 9 (We address this further below). This monoploid genome contained 64 354 genes. We also counted the genes in contigs that were not anchored into chromosomes and 29 534 genes were obtained. The gene number ranks A. argyi as the most gene-enriched species among the sequenced Asteraceae plants, and this gene number is about 2.5 times the average number of genes (36 795) reported for plant genomes (Ramírez-Sánchez et al., 2016). The average lengths of gene and coding DNA sequence (CDS) were 3416 and 1256 bp, respectively, with an average of 5 exons and 4 introns per gene (Table S9).

Table 1 Major features of the *Artemisia argyi* genome assembly

| Assembly feature                  | Hifi assembled | Hi-C Anchored |
|-----------------------------------|----------------|---------------|
| Assembly size (Mb)                | 8029.08        | 8030.38       |
| GC %                              | 35.46          | 35.46         |
| Repeat (%)                        | 73.59          | 73.59         |
| Number of scaffolds               | –              | 12 449        |
| Scaffold N50 size (Mb)            | –              | 206.40        |
| Scaffold N90 size (Mb)            | –              | 180.30        |
| Longest scaffolds (Mb)            | –              | 342.73        |
| Number of contigs                 | 10 274         | 15 063        |
| Contig N50 size (Mb)              | 8.32           | 6.25          |
| Contig N90 size (Mb)              | 1.46           | 0.64          |
| Longest contig (Mb)               | 43.52          | 40.67         |

Figure 1 Plant morphology and somatic chromosome number of *A. argyi*. (a). A plant of *A. argyi* cultivar ‘Xiang Ai’. (b). The karyotype of *A. argyi*. 

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in the Nr and TrEMBL databases, and 72.30% of the genes were classified by Gene Ontology (GO) terms, and 31.13% of the genes were annotated to Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways (Table S10). In addition, 1033 miRNAs, 19,784 tRNAs, 310 rRNAs and 3606 snRNAs were identified in the *A. argyi* genome (Table S11).

**Comparative genomic analysis of A. argyi**

To gain insights into the evolution of the *A. argyi* genome, a comparative genomic analysis was performed using *Arabidopsis thaliana*, *Vitis vinifera* and six other Asteraceae species (*A. annua*, *C. nankingense*, *E. breviscapus*, *H. annuus*, *L. sativa* and *C. tinctorius*). According to the sequence homology among these nine species, 93,888 protein-coding genes (including 64,354 genes in the monoploid genome of *A. argyi* and 29,534 genes in scattered contigs) comprised 3384 single-copy orthologues, 14,817 multiple-copy orthologues, 18,902 unique paralogues, 36,066 other paralogues and 20,719 unclustered genes (Figure 2b and Table S12). These genes were clustered into 27,915 gene families; among them, 5321 (19.06%) gene families contained 18,902 unique genes in *A. argyi* (Table S13). GO enrichment analysis showed that the biological functions of these genes were annotated to Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways (Table S10). In addition, 1033 miRNAs, 19,784 tRNAs, 310 rRNAs and 3606 snRNAs were identified in the *A. argyi* genome (Table S11).

**Figure 2** Assembly and genomic features of the *A. argyi* genome. (A). The circos diagram of *A. argyi* draft. (a) the genomic landscape of the 34 *A. argyi* pseudochromosomes. (b) the density of gene. (c) repeat coverage. (d) the density of SNP. (e) the density of Indel. (f) synteny relationship between pseudochromosomes. (B). Phylogenetic tree of seven species from the Asteraceae based on the information of single copy genes. And Arabidopsis and *V. vinifera* were used as the outgroup. The expanded gene families were marked with green and the contracted gene families were marked with red. (C). The *Ks* distributions of paralogous genes in *A. argyi*, *A. annua*, *C. nankingense*, *H. annuus*, *E. breviscapus* and *C. tinctorius* of the Asteraceae, and the eudicot species *V. vinifera*. (D). Kimura distance of LTR retroelements.
specific gene families were enriched in RNA-directed DNA polymerase activity (GO: 0003964), DNA polymerase activity (GO: 0034061), nucleotidyltransferase activity (GO: 0016779) and zinc ion binding (GO: 0008270) (Figure S5 and Data S2). Meanwhile, KEGG enrichment results showed that these genes were mainly enriched in pathways involved in phagosome (ko04145), protein processing in endoplasmic reticulum (ko04141), mismatch repair (ko03430) and terpenoid backbone biosynthesis (ko00900) (Figure S6 and Data S3). Furthermore, gene family evolution analysis showed that 40.04% (11 177/ 27915) of the gene families were expanded and 14.15% (3951/ 27 915) of the gene families were contracted in the A. argyi genome (Figure 2b). Compared to the other six species in the Asteraceae family, whose contracted genes were larger than the expanded genes, the number of expanded gene families in A. argyi was approximately three times that of the contracted gene families. GO enrichment analysis indicated that the functions of the expanded genes were significantly related in terms of binding, catalytic activity, photosynthetic electron transport in photosystem II and oxidoreductase activity (Figure S7 and Data S4). KEGG analysis revealed that expanded genes were enriched in photosynthesis, DNA replication, homologous recombination and several secondary metabolic pathways (Figure S8 and Data S5).

In particular, several markedly expanded genes were identified including PsbA genes (encoded photosystem II P680 reaction center protein D1) in PSII, replication factor A1 (RPA1) participating in homologous recombination and DNA replication, and heat shock protein genes (HSPs, HSP40, HSP70, HSP73) and terpene synthase genes (TPSs) (Table S14). It is well known that these genes are related to plant growth and stress responses. Given the ability of A. argyi to adapt to a wide variety of habitat conditions, these largely expanded genes may contribute to its successful expansion across the landscape and rapid growth.

Subsequently, we constructed a time-calibrated phylogenetic tree by using a concatenated sequence alignment of 944 single-copy orthologues shared by these nine species. These results verified the close evolutionary relationship between A. argyi and A. annua, and the divergence time of A. argyi and A. annua was approximately 7.4 million years ago (Mya). The most recent common ancestor (MRCA) of A. argyi and A. annua diverged from the MRCA of C. nankingense ~9.3 Mya, which together diverged from the MRCA of E. breviscapus ~41.3 Mya and further from the MRCA of L. sativa ~52.2 Mya (Figure 2b).

Whole-genome duplication (WGD) is the considered major factor driving genome evolution and expansion (Yan et al., 2021). In A. argyi, the WGD events were examined by distributions of synonymous substitutions (Ks) within genes in syntenic blocks compared with six other species (A. annua, C. nankingense, H. annuus, E. breviscapus, C. tinctiorus and V. vinifera). The distribution of Ks for the paralogous genes of the A. argyi genome showed two prominent peaks at ~0.04 and ~1.06, indicating that A. argyi has experienced two rounds of WGD (recent WGD and WGT-1) events. We further estimated that the most recent WGD event of A. argyi occurred at ~2.2 Mya, which was a species-specific duplication event and did not occur in the genomes of E. breviscapus, H. annuus and C. tinctiorus, but occurred in A. annua and C. nankingense. The WGT-1 event in A. argyi was a conserved whole-genome triplication event shared with E. breviscapus, C. tinctiorus and other Asterid-II plants (Badouin et al., 2017), occurring at approximately 62.9 Mya. Moreover, the Ks dot plot of retained paralogues in A. argyi genome also supported the occurrence of the WGD events (Figure 3a). Based on the Ks value (~0.6) of orthologous peaks for A. argyi and E. breviscapus, we predicted that their divergence time was ~38.2 Mya, which was close to the phylogenetic results. The relative age (Kimura distance) computed for LTR retroelements also indicates a recent increasing transposon activity (Figure 2d). The most recent WGD event and the recent outbreak of LTRs in A. argyi may be one of the most important reasons for its large genome size.

Chromosome fusion in the A. argyi genome

Synteny analysis showed that the 34 pseudochromosomes of A. argyi comprised 10 homologous groups, of which seven groups each had four sets of monoploid chromosomes, and three groups had two chromosomes in each (Figure S9). Moreover, each of the four chromosomes in the 1–7 chromosome groups can be divided into two subgroups according to the gene synteny analysis, which indicates that A. argyi is an allotetraploid (Figure S10), and this result was also consistent with that of survey analysis. Importantly, the length of chromosome 10 was almost the sum of the lengths of chromosomes 8 and 9 (Figure 3a and Table S5), and intragenome synteny analysis showed that chromosomes 8 and 9 shared close syntenic regions with chromosome 10 (Figure 3b). In addition, 11 780 (92.05%), total 12 797) genes on chromosome 10 were homologous with the genes on chromosomes 8 and 9 based on the BLAST results. Together, these data allow us to postulate that the ancestral 8 and 9-like chromosomes were fused into the chromosome 10 in A. argyi, and chromosome 10 appears as the end-to-end fusion of ancestral 8 and 9-like chromosomes, accompanied by at least one inversion and two intrachromosomal translocation events (Figure 3b).

By comparing the number of homologous genes on chromosome 10 with those on chromosomes 8 and 9, we found that 1129 genes were missing and that 1017 genes were newly formed on chromosome 10. These genes were mainly concentrated in the biological process category, including heme transport, iron coordination entity transport (lost genes), snRNA binding and oxidative phosphorylation (novel genes) (Data S6). In addition, by comparing the expression levels of homologous genes on chromosome 10 with those on chromosomes 8 and 9, we found that 411 genes were upregulated and 404 genes were downregulated on chromosome 10. GO enrichment analysis showed that the biological functions of these upregulated genes were significantly enriched in response to external biotic stimulus and cellular response to salicylic acid stimulus, and the downregulated genes were enriched in the methylerythritol 4-phosphate pathway (Data S7).

Genes involved in flavonoid biosynthesis

Eupatilin, jaceosidin, hispidulin, schaftoside, isoschaftoside and vitexicarpin are representative bioactive flavonoids in A. argyi that contribute to versatile pharmacological effects such as anti-inflammatory, antioxidation and anti-tumor effects (Maleki et al., 2019; Nabavi et al., 2015; Serafini et al., 2010). Ultra-performance liquid chromatography (HPLC) was used to quantify these flavonoids in seven different tissues (roots, rhizome, stem, and four different developmental stages of leaves A–D) of ‘Xiang Ai’. The obtained results showed that these bioactive flavonoids were more abundant in leaves than in other tissues. Among them, the content of eupatilin was the highest, and it increased with the growth stage of leaves (Figure S11). However, the genes
that participated in the biosynthesis of these flavonoids such as hispidulin, jaceosidin and eupatilin in *A. argyi* remain largely unknown. Based on extensive investigations of the flavonoid biosynthesis pathway in other plants (Saito *et al*., 2013), we proposed the possible biosynthesis routes of these compounds in *A. argyi* (Figure 4a,c).

In total, 44 candidate genes encoding 12 key enzymes in flavonoid biosynthesis pathway were identified by homologue searching and functional annotation. Of note, nearly half of the candidates especially were expanded genes and the number of phenylalanine ammonia-lyase (*PAL*), 4-hydroxylase (*C4H*), hydroxycinnamoyl transferase (*HCT*), chalcone synthase (*CHS*), flavanone hydroxylase (*F3H*, *F30H*) homologues in *A. argyi* was dramatically increased relative to that in Arabidopsis (Table S15). We mapped these genes to *A. argyi* genome and found that the *PAL* exhibited tandem repeats on chromosomes 5 (Data S8). Subsequently, a transcriptomic analysis was performed using samples from roots and leaf organs (Figure S12) to identify differentially expressed genes (DEGs) between different tissues and different developmental stages of leaves. Based on their expression patterns, almost all candidate genes were expressed in seven selected tissues, but the expression levels of the first four genes in this pathway, especially the expression levels of *HCT* genes in root samples, were higher than those in leaf samples, while the expression levels of the downstream genes, especially the *CHS* genes, were higher in leaves than in roots (Figure 4b). HCT is a key enzyme in lignin synthesis (Baucher *et al*., 2003), while *CHS* is the first rate-limiting enzyme in plant flavonoid synthesis (Dixon and Paiva, 1995). Therefore, the expression patterns of the *HCT* and *CHS* genes were crucial for the regulation of lignin and flavonoid synthesis in *A. argyi*, which probably facilitates its rapid adaptation to heterogeneous environments. Furthermore, most of the DEGs involved in flavonoid biosynthesis were upregulated in the leaf tissues, and this correlates well with the fact that flavonoids, for example, hispidulin, jaceosidin and eupatilin, are mainly enriched in *A. argyi* leaves.

Flavonoid O-methyltransferase (FOMT) is a key enzyme for the postmodification of flavonoid compounds. Studies have confirmed that O-methylated flavonoids have stronger antioxidant, anti-inflammatory, and anti-cancer functions (Zhao *et al*., 2019). In consideration of the active ingredients, including hispidulin, jaceosidin and vitexicarp, in *A. argyi* that were all O-methylated flavonoids, we further investigated the whole-genome FOMT genes in *A. argyi* using the conserved domain and the reported FOMT genes as queries. A total of 83 FOMTs were identified. Phylogenetic analysis showed that these FOMTs were clustered into five main subclades based on their catalytic sites including 31 3'FOMT, six 6FOMT, 10 4'FOMT, 32 caffeoyl-CoA O-methyltransferase (CCoAOMT) and four unclassified FOMTs (Figure 4d). We found that almost all types of FOMTs were significantly expanded in the *A. argyi* genome and exhibited a pattern of tandem duplication (Data S9). Most of the CCoAOMTs were primarily expressed in roots, while the FOMTs catalysing methylation at the 3'-OH, 4'-OH and 6-OH groups were mainly expressed in leaves (Figure 4d). Hispidulin has a methoxy on its C6 position, indicating that its enzymatic synthesis requires 6FOMT to mediate C6-methoxylation. Based on the chemical structure and a previous study (Zhang *et al*., 2016),
scutellarein may be a precursor for hispidulin biosynthesis. Jaceosidin is a di-methoxyflavone with one methyl group on the C6 position of the A ring and another on the C3 position of the B ring, suggesting that its synthesis requires an additional FOMT by comparison with hispidulin. Eupatilin is a trimethoxyflavone that has an extra methyl group at the C4’ position of the B ring compared to jaceosidin, suggesting that there may be a 4’FOMT catalysing the conversion of jaceosidin to eupatilin (Figure 4c). In summary, the identification of these FOMT candidates based on genomic and transcriptome analyses will accelerate the enzymatic synthesis pathway of hispidulin, jaceosidin, and eupatilin.

Genes involved in terpenoid biosynthesis

Volatile oil is a significant pharmacodynamic component in the leaves of A. argyi due to a rich content of terpenoids. The volatile oil from A. argyi leaves contains abundant monoterpenoids and sesquiterpenoids (Figure 5a). Although terpenoids are diverse and have various structures, they all come from the common precursors isopentenyl pyrophosphate (IPP) and dimethylallyl pyrophosphate (DMAPP). IPP and DMAPP are mainly produced through the mevalonate pathway (MVA) pathway in the cytoplasm and the methylerythritol phosphate (MEP) pathway in the plastid (Sapir-Mir et al., 2008; Vranova et al., 2013). Candidate genes participating in MVA and MEP pathways were screened by using the methods of homology searching and functional annotation. The obtained results indicated that a total of 66 genes encoding 14 gene families were involved in these two pathways in A. argyi (Figure 5b). These genes were widely distributed on A. argyi chromosomes, especially on chromosomes 7 (Data S10). The key genes identified in the MEP pathway were greatly expanded compared with the genes identified in the MVA pathway. RNA-seq analysis demonstrated that genes related to the MEP pathway were more specifically expressed in leaves than those in the MVA pathway (Figure 5c), indicating that the terpenoids in A. argyi leaves mainly come from the MEP pathway.

Terpene synthase (TPS) family genes are responsible for the biosynthesis and structural diversity of terpenoids (Chen et al., 2011). We found that TPS genes were extremely expanded in the A. argyi genome. In total, we identified 135 TPS genes in the A. argyi genome (Data S11). According to phylogenetic analyses, these TPSs were grouped into five subfamilies, including TPS-a, TPS-b, TPS-c, TPS-d and TPS-e/f (Figure 6a). More than 80% of TPSs belonged to TPS-a and TPS-b subfamilies, which are mainly involved in the biosynthesis of monoterpenes, sesquiterpenes, and diterpenes, indicating the remarkable expansion of these two TPS subfamilies. The expression patterns of TPSs in different tissues were analysed, most of which had relatively high expression levels in leaves compared to other tissues (Figure S13). These results were consistent with the abundant terpenoids in the leaves of A. argyi. Chromosome localization showed that the TPS genes were not uniformly distributed throughout the chromosomes (Figure S14). For example, there were 12 TPS-a genes.
Figure 5 Volatiles compounds in *A. argyi* leaves and their biosynthesis pathways. (a). Gas chromatogram of volatile oils from *A. argyi* leaves. Monoterpenes and sesquiterpenes were marked green and red, respectively. (b). The proposed MVA and MEP pathways in *A. argyi*. Red fonts indicated the abbreviations of enzymes participating in these two pathways. And the full names of these enzymes were listed in Data S10. (c). The expression patterns of candidate genes involved in MVA and MEP pathways in different tissues. The expanded genes were marked in red. R, root; Rh, rhizome; S, stem; LA, leaf buds, 0 day; LB, young leaves 15 days; LC, mature leaves 30 days; LD, old leaves 45 days.
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clustered on chromosome 6 and 13 TPS-b genes clustered on chromosome 3, with a pattern of tandem duplication (Figure 6b, c). Expression analysis indicated that most of the TPS genes in the cluster were not actively expressed, except for AY184877-RA (TPS-a), AY184880-RA (TPS-a) and AY075453 (TPS-b), which were significantly expressed in leaf tissues (Figure 6b,c). This finding demonstrates that some duplicated TPSs may have undergone neofunctionalization.

Discussion

The high-quality A. argyi genome sequence in this study represents the first species in the genus Artemisia for which chromosome-level assembly has been constructed. This well-annotated genome will be the foundation for evolutionary and molecular biological studies of this economically and medicinally important plant. However, the high heterozygosity (2.36%), large proportion of repetitive sequences (75.86%) and polyploidy (allotetraploid) present significant challenges for the genome assembly of A. argyi. In this study, an integrated strategy combing PacBio long HiFi reads, Hi-C sequencing and Illumina short reads greatly facilitated the assembly of the complex polyploidy A. argyi genome. Compared with the other 14 species in Asteraceae whose genomes have been released, A. argyi features the largest genome with a size of 8.03 Gb. The scaffold N50 (206.4 Mb) of the A. argyi genome was the longest among them, and the contig N50 (6.25 Mb) is only shorter than that of safflower (Wu et al., 2021). In brief, the A. argyi genome is the fifth chromosome-level genome in Asteraceae, and the first chromosome-level genome in Artemisia.

WGD events and TE amplification are the main causes of large genomes (Bennetzen, 2002; Van de Peer et al., 2009). Here, the analysis of Ks distribution indicated that two rounds of WGD events occurred in the A. argyi genome (Figure 2c). The most recent WGD (~2.2 Mya) is an A. argyi and its closely related species (A. annua and C. nankingense) specific event, which is distinct from the WGD-2 event in the sunflower genome (Badouin et al., 2017). Combined with a WGT-1 event occurring in all eudicots and a basal WGT-1 event occurring in Asteraceae, we conclude that A. argyi underwent at least three rounds of WGDs. The additional WGD events may also contribute to gene family expansion. In comparison with other sequenced Asteraceae species, we found that the number of expanded gene families in A. argyi was approximately three times that of the contracted gene families (Figure 2b), which accounts for a prominently large number of genes in the A. argyi genome. In particular, PsbA genes in photosystem II, RPA1 in DNA replication, HSPs and TPS were markedly expanded, all of which participate in plant growth and development and stress responses. Moreover, the transposable elements occupied 73.59% of the A. argyi genome, and the LTR/Gypsy subfamilies (24.47%) were the most abundant. The distribution of Ks also hints that LTR retrotransposons explosion occurred recently in the A. argyi genome (Figure 2d). Together, the additional WGD event and the recent proliferation of LTRs in A. argyi may be responsible for the expansion of the genome and may play vital roles in A. argyi adaptations to challenging environments.

Variation in basic chromosome numbers is not a rare phenomenon in some genera in the Asteraceae family, such as Melampodium whose basic chromosome numbers are x = 11, 9 and 14 (McCann et al., 2016). Artemisia is reported to have at least two basic chromosome numbers, with x = 9 being the common chromosome number and x = 8 being less frequent (Pellicer et al., 2010). The somatic chromosome numbers of A.

Figure 6 Expansion of terpene synthase-encoding genes and their gene clusters on chromosomes in the Artemisia argyi genome. (a) Phylogeny of TPSs identified in A. argyi genome. (b) Phylogeny, expression profiles and chromosomal position of TPS-a clade gene cluster on Chr6_1. The outgroup gene of the phylogenetic tree was TPS-e subfamily gene with gene ID AY065441-RA. (c) Phylogeny, expression profiles and chromosomal position of TPS-b clade gene cluster on Chr3_1. The outgroup gene of the phylogenetic tree was TPS-e subfamily gene with gene ID AY294582-RA.
The cultivated mugwort ‘Xiang Ai’ was used for the construction of the reference genome. The young leaves of 90-day-old ‘Xiang Ai’ plants were sampled to extract high-quality genomic DNA for genome sequencing and Hi-C analysis. Roots, rhizomes (Rh), stems (S), leaf buds (0 day, LA), young leaves (15 days, LB), mature leaves (30 days, LC) and old leaves (45 days, LD) of ‘Xiang Ai’ were collected for RNA-seq analysis.

Flow cytometry

The nuclear DNA content of ‘Xiang Ai’ was measured by flow cytometry according to the method described previously (Dolezel et al., 2007). Briefly, leaves from ‘Xiang Ai’ and Chrysanthemum nankinense plants were finely chopped with a razor blade in 2 mL Gatter’s buffer, respectively. After the suspension was filtered through a 48 μm nylon membrane, 200 μL of PI (50 μg/mL) and 100 μg/mL RNaseA were added immediately. Following 30 min of incubation on ice, the samples were detected by flow cytometry (BD FACSCalibur, BD Biosciences, Wuhan, China).

Estimation of genome size

Illumina-seq generated approximately 215 Gb of clean reads, which were subjected to K-mer analysis to estimate the genome size of ‘Xiang Ai’. The 21-mer frequency distribution was shown in Figure S2.

Whole-genome sequencing

For PacBio SMRT sequencing, high-quality DNA from ‘Xiang Ai’ was first sheared and concentrated to construct 15-Kb DNA sequencing libraries and subsequently run on a PacBio Sequel II platform according to the manufacturer’s instruction with seven cells.

For genome survey sequencing, 5150-bp paired-end (PE) libraries were constructed for sequencing on an Illumina NovaSeq 6000 platform and ~215 Gb of raw sequencing data were obtained.

For Hi-C sequencing, two Hi-C libraries digested with MboI restriction enzyme were sequenced on BGI MGISEQ-2000 to generate ~422 Gb of valid data from 150 PE reads.

Genome assembly and evaluation

Briefly, we performed de novo assembly using HiCanu v2.2 (Koren et al., 2017) and Hifiasm (version 0.13-r308) (Cheng et al., 2021) and the result showed that the Hifiasm assembly (N50 = 8.32 Mb with 10 274 contigs) was better than the HiCanu assembly (4.69 Mb with 48 227 contigs). Then, the draft genome from the Hifiasm assembly was further assembled into scaffolds with Hi-C data using the 3D-DNA pipeline (version 180922) (van Berkum et al., 2010). These scaffolds were roughly split by Juicebox (version 4.0.10) (Robinson et al., 2018) and another round of scaffolding by 3D-DNA. Genome assembly completeness was assessed by BUSCOs (Simao et al., 2015) and transcriptome data.

Genome annotation

Transposable elements of A. argyi genome were identified by homology- and de novo-based methods. Repbase (version 2017-01-27) was used to build the homology repeat library (Jurka et al., 2005). Then, RepeatModeler v2.0.1 (Jurka et al., 2005) was used to construct the de novo repeat library. Finally, RepeatMasker v4.1.0 (Tempel, 2012) was used to annotate the repetitive elements in the Repbase and the de novo repeat library.

Protein coding genes of A. argyi genome were annotated by MAKER according to three complementary methods: de novo prediction, homology-based prediction and transcriptome-based prediction (Campbell et al., 2014). First of all, IsoSeq3 was
employed to identify full-length high-quality transcripts from PacBio Sequel II (Singh et al., 2016). HISAT2 (version 2.1.0; Pertea et al., 2016) and Cufflinks (version 2.2.1) (Trapnell et al., 2012) were used to predict the genes by using the RNA-seq data and the assembled genome. In the second round, Augustus v 3.2.1 (Keller et al., 2011), GeneMark-ES Suite v 4.61.1c (Lomsadze et al., 2005) and SNAP (version 2013-02-16) (Johnson et al., 2008) tools were used for gene model training for de novo prediction. Finally, precise gene annotation was performed by using non-redundant proteins from Artemisia annua and Helianthus annuus based on the homology-based approach. Complete BUSCO hits were used to evaluate the gene annotation results of A. argyi genome (Simao et al., 2015).

Following the gene annotation, BLAST hits analyses against several functional databases (NR, EggNOG, SwissProt and TrEMBL) were performed on the predicted protein-coding genes to identify homologous proteins in other species using diamond (version v2.0.4.142; Buchfink et al., 2015). In addition, possible GO terms were obtained using the SwissProt database and Trembl database ID mapping. Gene pathways were accomplished with KEGG analysis by using KOBAS v 3.0 (Xie et al., 2011)

tRNAs were annotated by TRNAscan-SE (version 1.3.1) (Lowe and Eddy, 1997), rRNAs were annotated by BLASTN (version 2.10.1) (Gardner et al., 2009). miRNAs and snRNAs were annotated by searching the Rfam database (version 14.5) using BLASTN and INFERNAL (version 1.1.2) (Nawrocki et al., 2009).

Sequence alignment and variation analysis

Burrows–Wheeler Aligner (BWA, version 0.7.17) software was used to map to all clean reads to the assembled genome (Li and Durbin, 2009) with the methods described by He et al. (2021). In brief, SAMTools (version 1.9) were employed to convert the mapping results into the BAM format (Li et al., 2009). Picard package (Version 1.96) was used for the filtration of duplicated reads. Genome Analysis Toolkit (GATK, version 3.8-0-ge9d806836) was used to realign reads around Indels (McKechnie et al., 2010).

Variations were detected with both SAMTools mpileup and GATK Haplotype Caller packages, and only concordance results were retained. Raw variations were filtered by GATK VariantFiltering packages with the following parameters: --filter-name FilterQual --filter-expression “QUAL < 60.0” --filter-name FilterQD --filter-expression “QD < 20.0” --filter-name FilterFS --filter-expression “FS > 13.0” --filter-name FilterMQ --filter-expression “MQ < 30.0” --filter-name FilterMQRankSum --filter-expression “MQRankSum < -1.65” --filter-name FilterReadPosRankSum --filter-expression “ReadPosRankSum < -1.65” --cluster-window-size 10 --cluster-size 2. SNPs within 10 bp with an indel were removed. Finally, SnpEff software was used to annotate the identified SNPs and Indels (Cingolani et al., 2012).

Gene family, phylogenomic analysis and WGD identification

Gene families of A. argyi genome and Arabidopsis thaliana, Vitis vinifera and six other Asteraceae species were identified by OrthoMCL (version 2.0.9) with default parameters (Li et al., 2003). RaxML (version 8.2.12) were used to construct the phylogenetic tree by using the single-copy orthologues of the nine species (Stamatakis, 2014). The divergence times of A. thaliana and A. argyi from TimeTree (http://timetree.org/) were used for calibration and the divergence times of phylogenetic tree were estimated by r8s (version 1.81) (Sanderson, 2003). The expanded and contracted gene families were calculated by CAFE (version 4.2) in each lineage (De Bie et al., 2006). WGD events in the A. argyi genome were searched according to the WGDi pipeline (Sun et al., 2021).

RNA sequencing

Total RNA of different tissues was extracted using TRizol reagent. An mRNA sequencing library of seven different tissues was constructed on an Illumina Novaseq 6000 platform by 150 bp PE sequencing. For full-length transcriptome sequencing, a mixed RNA library from different tissues was prepared according to the PacBio ISO-Seq experimental workflow and subsequently run on a PacBio Sequel II platform. For RNA-seq analysis, 2 μg RNA from each sample was sequenced on the Illumina platform. Three replications were performed for each sample.

Gene identification in flavonoid and terpenoid biosynthesis pathways

The protein sequences of the enzymes (PAL, C4H, 4CL, HCT, C3H, CHS, CHI, F3H, and FLS) in flavonoid biosynthesis pathways of A. thaliana were obtained from the TAIR database, and those for F3′H and FNSII in fleabanes, FNSI in parsley, celer, and F6H in soybean were obtained from previous studies (He et al., 2021) and the NCBI database. These sequences were blasted against the A. argyi protein sequences using BLASTP (E-value < 1e-5).

Functional proteins involved in the MEP and MVA pathways of terpenoid backbone biosynthesis in A. thaliana were obtained from the TAIR database. Homologues of these proteins in the genome of A. argyi were investigated by BLASTp with an E-value cutoff of 1e-5. Fragments per kb of exon model per million mapped fragments (FPKM) not <10 were selected for heatmap analysis.

Identification of terpene synthase genes (TPSs) and flavonoid O-methyltransferase (FOMTs)

The protein sequences of TPSs were identified by screening with functional motifs PF01397 and PF03936. A total of 146 TPS genes were predicted in the ‘Xiang Al’ genome. There are 11 genes of TPSs were removed manually, since it was not possible to calculate genetic distance by MEGA6.

We searched the candidate FOMT genes by the combination of conserved domain (PF01596) and homologue-based BLAST, and repetitive sequences were removed. A total of 83 FOMT genes were predicted in the A. argyi genome. 11 selected FOMT proteins downloaded from NCBI based on high gene homology and genome annotation were subjected to phylogenetic analysis.

Phylogenetic reconstruction of TPSs and FOMTs

The neighbour-joining trees were constructed using the MEGA6 software (Tamura et al., 2013). Heatmap analysis based on RNA-seq data was performed by the pheatmap package in R language (version 1.0.12).

HPLC analysis of flavonoids

Agilent 1260 Infinity HPLC system (Agilent Technologies) was used for HPLC analyses. The flavonoids were separated by ZORBAX RRHD Eclipse Plus 95A C18 column (2.1 × 100 mm, 1.8 μm, Agilent) and detected at 330 nm by UV. The mobile phases were acetonitrile (A) and 0.1% phosphoric acid (B) with the flow rate of 0.4 mL/min. The separation gradient conditions were as follows: 0–0.5 min, 2–5% A; 0.5–7 min, 5–25% A; 7–11 min, 25–33% A; 11–14 min, 30–33% A; 14–19.5 min, 33–
45% A; 19.5–20.5 min, 45–85% A; 20.5–27 min, 85–98% A; 27–28 min, 98–2% A.

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Conflicts of interest
The authors declare no competing interests.

Author contributions
L.Q.H., D.H.L., G.G. N. and Y.H.M., conceived and managed the project. D.D.L., T.T.Z., H.Z.D. and Z.P.X. contributed to the assembly and annotation. Y.H.M., D.D.L. and T.T.Z. analysed the samples for sequencing. Z.H.L. and G.G. N. contributed to the preparation of DNA sequencing have been deposited at Sequence Read Archive (SRA) National Genomics Data Center under the accession number PRJNA804653. The whole assembled genome data have been submitted in the National Genomics Data Center under the accession number PRJNA804653. The whole assembled genome data have been submitted in the National Genomics Data Center under the accession number PRJNA804653.

Data availability statement
The chromosome-level genome assembly of Artemisia argyi.
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**Supporting information**

Additional supporting information may be found online in the Supporting Information section at the end of the article.

**Figure S1** Estimate of the genome size of *A. argyi* (Xiang ai, 4-6) by flow cytometry.

**Figure S2** Estimate of the genome size and complexity of *A. argyi* by K-mers method.

**Figure S3** The flowchart of processing pipeline used to assemble the *A. argyi* genome.

**Figure S4** The Hi-C map of *A. argyi* genome assembly.

**Figure S5** Enriched GO terms for gene families specific to *A. argyi*.

**Figure S6** KEGG enrichment of *A. argyi* specific genes.

**Figure S7** GO enrichment of *A. argyi* expanded genes.

**Figure S8** KEGG enrichment of *A. argyi* expanded genes.

**Figure S9** Syntenic relationships based on pairs of collinear genes of *A. argyi*.

**Figure S10** Clustering of counts of 13-mers.

**Figure S11** The contents of representative flavonoids in seven different tissues of *A. argyi*. R, root; Rh, rhizome; S, stem; LA, leaf buds, 0 d; LB, young leaves 15 d; LC, mature leaves 30 d; LD, old leaves 45 d.

**Figure S12** Expression correlation heat map of pairwise samples.

**Figure S13** Expression profiles of TPSs in *A. argyi*.

**Figure S14** Chromosome distribution of TPS family genes in the *A. argyi* genome.

**Table S1** K-mer analysis of the *A. argyi* genome using K-mer = 21.

**Table S2** Summary of PacBio sequencing.

**Table S3** Statistics of pre-assembly of the *A. argyi* genome by using Hifiasm and Hicau.

**Table S4** Summary of Hi-C sequencing.

**Table S5** Summary of chromosome level assembly based on Hi-C data.

**Table S6** Statistics of illumina and isofrom sequencing clean reads mapping rate of *A. argyi* genome assembly.

**Table S7** The assessment of *A. argyi* genome and annotation completeness with BUSCO.

**Table S8** Summary of repeats and transposable elements in the *A. argyi* genome assembly.

**Table S9** Statistics of genes annotated in the *A. argyi* genome.

**Table S10** Functional annotation of predicted protein-coding genes in the *A. argyi* genome.

**Table S11** Statistics of noncoding RNA genes in the *A. argyi* genome.

**Table S12** Gene family categories.

**Table S13** Comparisons of genes and gene families.

**Table S14** Significantly expanded genes in the *A. argyi* genome.

**Table S15** Copy numbers of gene families involved in flavonoid biosynthesis among plant species.

**Data S1** Summary of RNA sequencing.

**Data S2** Enriched GO terms for gene families specific to *A. argyi*.

**Data S3** Enriched KEGG terms for gene families specific to *A. argyi*.

**Data S4** Enriched GO terms for expanded genes in *A. argyi*.

**Data S5** Enriched KEGG terms for expanded genes in *A. argyi*.

**Data S6** Enriched GO terms for lost and novel genes in chromosome 10 by comparision with chromosome 8 and 9.

**Data S7** Enriched GO terms for up- and down-regulated genes in chromosome 10 by comparision with chromosome 8 and 9.

**Data S8** Location of the flavonoids biosynthesis genes in the *A. argyi* genome.

**Data S9** Genome wide identification, classification of flavonoid o-methyltransferase (FOMT) gene family in *A. argyi* genome.

**Data S10** Location of the already-known terpenoids metabolism genes in the *A. argyi* genome.

**Data S11** Genome wide identification, classification of terperne synthase (TPS) gene family in *A. argyi* genome.