Comparative Genomics Uncovers the Genetic Diversity and Synthetic Biology of Secondary Metabolite Production of Trametes

Yan Zhang, Jingjing Wang, Chen Yajun, Minghui Zhou, Wei Wang, Ming Geng, Decong Xu and Zhongdong Xu
School of Life Sciences, Hefei Normal University, Hefei, China

ABSTRACT
The carbohydrate-active enzyme (CAZyme) genes of Trametes contribute to polysaccharide degradation. However, the comprehensive analysis of the composition of CAZymes and the biosynthetic gene clusters (BGCs) of Trametes remain unclear. Here, we conducted comparative analysis, detected the CAZyme genes, and predicted the BGCs for nine Trametes strains. Among the 82,053 homologous clusters obtained for Trametes, we identified 8518 core genes, 60,441 accessory genes, and 13,094 specific genes. A large proportion of CAZyme genes were cataloged into glycoside hydrolases, glycosyltransferases, and carbohydrate esterases. The predicted BGCs of Trametes were divided into six strategies, and the nine Trametes strains harbored 47.78 BGCs on average. Our study revealed that Trametes exhibits an open pan-genome structure. These findings provide insights into the genetic diversity and explored the synthetic biology of secondary metabolite production for Trametes.

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1. Introduction
Trametes, one of white-rot basidiomycetes, belongs to the family Polyporaceae and the class Agaricomycetes of fungi, commonly grows in the tiled layer of decaying wood [1], and uses deciduous tree [2]. This genome currently comprises at least 20 species until April 2018 [3,4], such as Trametes sp., Trametes trogii, and Trametes villosa CCMB561. Trametes is known to decolorize dyes [1], and several of its species have been used for centuries in traditional medicine in East Asia countries [5]. Trametes sp. SQ01 can decolorize azo, anthraquinone, and triphenylmethane dyes [6]. T. trogii, a worldwide-distributed white-rot fungi and an outstanding laccase producer, degrades all lignocellulosic materials [7]. T. villosa CCMB561, a common species in the Brazilian semiarid region [8], has been isolated from decaying wood and is considered a high-potential fungus for biotechnological applications because it is a good producer of the three important ligninolytic enzymes (laccase, manganese peroxidase, and lignin peroxidase) [4]. The laccases from Trametes can be used as a biocatalyst in enzymatic biofuel cells [9,10]. Hence, understanding the genetic diversity of Trametes is beneficial for its biotechnological applications.

Carbohydrate-active enzymes (CAZymes) are involved in the polysaccharide degradation of plant cell wall [11], lignin, cellulose, hemicellulose, and pectin [12]. In particular, CAZyme classes (glycoside hydrolases [GH], carbohydrate esterases [CEs], and polysaccharide lyases) play central roles in plant biomass decomposition by bacteria and fungi [13]. The total expression levels of CAZyme genes of T. versicolor F21a are almost twice as that of the control group during algaecide, and the majority of expressed CAZyme genes belong to GH family and auxiliary activities [14]. Similarly, 590 genes of T. villosa CCMB561 are identified as CAZyme genes [4]. Although the compositions of CAZyme genes of fewer species of Trametes have been reported, the composition of CAZyme genes of Trametes has not been comprehensively analyzed.

Secondary metabolites (referred to as natural products or specialized metabolites) are the foundation of many drugs [15] and important chemicals used in agriculture and nutrition [16]. In general, fungi are rich sources of thousands of second metabolites, and fungal second metabolites can be grouped into four primary chemical types: polyketides (PKS), terpenoids, shikimic acid-derived compounds, and non-ribosomal peptides (NRPS) [17]. The secondary metabolites of fungi are crucial in its development and actively shape interactions with other microbes [18]. The number of genes coded for the secondary metabolism of ascomycetes is higher
than that of basidiomycetes, archeno-ascomycetes, and chytridiomycetes [19]. Several fungal genomes have been sequenced, and the development of various genome mining software tools, such as antiSMASH [16] and ClusterFinder [20], enables researchers to analyze the biosynthetic gene clusters (BGCs) of secondary metabolites. Genomes of filamentous fungi contain up to 90 potential BGCs encoding their diverse secondary metabolites [21], and 24 genomes of *Penicillium* are mined for BGCs and associated PKS and NRPS BGCs to known pathways [22]. However, no systemically comparative analysis has been conducted to identify the BGCs of secondary metabolite in *Trametes*.

In this study, we collected nine available genomic data from eight species of *Trametes* and conducted a comparative genomic analysis to investigate the genetic diversity and the synthetic biology of secondary metabolites. We obtained the *Trametes* pan-genome, annotated the pan-genome against clusters of orthologous group (COG) database and eggNOG database, and detected CAZyme genes. The BGCs for nine *Trametes* strains were also predicted. Our results showed that *Trametes* exhibits an open pan-genome structure and diverse genetic diversity for *Trametes* strains. The different distributions of CAZymes of *Trametes* reveal variations in carbohydrate utilization for *Trametes* strains. The predicted BGCs with unknown functions in *Trametes* suggested that *Trametes* have a great potential value for secondary metabolite production.

### 2. Materials and methods

#### 2.1. Genomic data collection of *Trametes*

We used “Trametes” as the keyword and searched in genome database of National Center for Biotechnology Information (NCBI) on September 14, 2018. We found that *Trametes* comprising 8 species and containing nine full genomes, such as *T. cinnabarina* BRFM137, *T. cinnabarina* FP104138-Sp, *T. coccinea* BRFM310, *T. hirsuta*, *T. polyzena*, *T. pubescens*, *T. sp. AH28-2*, *T. versicolor* FP-101664SS1, and *T. villosa*. Although there are 20 species of *Trametes* were discovered, there are only nine full genomes affiliated with 8 species can be obtained in public database. As such, we downloaded the genomic data (draft) of these strains for comparative genomics analysis.

#### 2.2. Gene prediction and ortholog identification

We conducted the genome annotation for nine strains using Prodigal version 2.6 [23]. In particular, to predict the genes and proteins for nine genomic data of *Trametes*, we applied the Prodigal with default setting to recognize open reading frames (ORFs) and protein sequences [24]. We identified the protein orthologs of *Trametes* by using OrthMCL version 2.0.9 [25] with e-value <1 e-5 and inflation parameter of 1.5. We divided the homologous clusters into three groups: core, accessory and specific groups. The core genes or proteins represent the genes or proteins shared in all nine genomes of *Trametes* used in our study, while the accessory genes or proteins comprised genes or proteins shared by at least two strains but not all nine *Trametes* strains. The rest of genes or proteins only occurred in one strain were clustered into specific groups (strain-specific genes or proteins). We obtained the protein sequences of homologous clusters for *Trametes*.

#### 2.3. Phylogeny analysis

The nuclear ribosomal ITS region has been reported as the primary fungal barcode marker [26] and ITS1 or ITS2 was widely used to identify the broadest range of fungi [27]. Hence, we applied ITSx [28], which is an open source software utility to extract the highly variable ITS1 and ITS2 subregions from ITS sequences, to extract the ITS1 and ITS2 sequences from the genomic data of nine *Trametes* strains. Meanwhile, we chose the ITS1 and ITS2 sequences of *Coriolopsis caperata* (accession number: AB158316) as outgroup. In our work, due to the fact that the ITS1 and ITS2 sequences of *Trametes coccinea* were not extracted, we selected the ITS1 and ITS2 sequences of *Pycnoporus coccineus* strain MUCL 38523 [29] (one strain of *T. coccinea*, accession number: FJ873395) to represent *T. coccinea* and construct the phylogenetic tree. The ITS1 and ITS2 sequences were aligned using the MUSCLE version 3.8.31 [30] with default setting, respectively. The multiple sequences alignment result was eliminated by using Gblocks version 0.91b [31] to remove the regions that were divergent, misaligned, or with a larger number of gaps. Then we used PHYLIP version 3.696 [32] with 100 bootstrap iterations to construct a maximum-likelihood tree based on the concatenated multiple sequences alignment result. Moreover, we built a pan-genome tree by MEGA software version 5 [33] based on the composition of orthologous. In particular, we calculated the pairwise Manhattan distance between each strain based on the presence and absence of orthologs proteins.

#### 2.4. Functional annotation

To obtain the COG functions, the protein sequences of homologous clusters for *Trametes* were aligned to individual COG proteins [34], which downloaded
from NCBI (https://www.ncbi.nlm.nih.gov/COG/) using BLASTP with e-value <1e-4 and the top one of the results was chosen as the best annotation of each protein. The annotations of the protein sequences of homologous clusters were assigned to 25 functional categories. We also annotated the protein sequences of homologous clusters against eggNOG annotation database by using the eggnog-mapper [35] and we divided the GO annotations of protein sequences of homologous clusters into three groups, including biological process, molecular function, and cellular component. We annotated these protein sequences against CAZyme database to investigate the functional composition related to carbohydrate metabolism. In particular, we downloaded the CAZy database from dbCAN [36] (http://csbl.bmb.uga.edu/dbCAN/). According to the manual of dbCAN CAZyme annotation, protein sequences were annotated by running hmmscan from the HMMER version 3.1b1 [37,38] with default setting and the annotated results were summarized into GT, GH, carbohydrate-binding module (CBM), AA, PL, and CE.

2.5. BGC detection

To detect the biosynthetic pathways of secondary metabolites for Trametes, antiSMASH version 4.0 [16] was used to identify the BGCs. A set of nine genome sequences of Trametes was used as the input data for predicting the BGCs by using antiSMASH with the mode of fungi and other default setting. The BGCs were determined to their subtypes (e.g., type I polyketide, terpenoid).

3. Results and discussion

3.1. Pan-genome construction and analysis

Nine available genomic data from eight species of Trametes were collected for the pan-genome analysis. The assembled genome of Trametes ranged from 31.62 Mb to 57.98 Mb, and the number of contigs/scaffolds ranged from 13 to 10,327. In addition, the number of CDSs/ORFs predicted by Prodigal ranged from 56,735 to 101,817 (Table 1). The number of genes predicted in our study for Trametes was greater than that in a previous report. The predicted genes of T. villosa CCMB561 in the present study were 101,817, whereas those in the previous study were 16,711 [4]. This difference might be due owing to the different tools and parameters. Orthologs from the 610,421 high-quality proteins of Trametes were also detected to characterize the differences of genomic composition among the nine strains. A total of 82,053 homologous clusters were obtained, and the accumulation curve of the homologous cluster showed that Trametes exhibits an open pan-genome structure (Figure 1(a)). The size of Trametes pan-genome tended to increase gradually with new strains and comprised 82,053 non-redundant genes within nine strains (Figure 1(a)). On the contrary, the size of core-genome tended to decrease progressively with new strains and comprised 8518 non-redundant genes within the nine Trametes strains (Figure 1(a)). A total of 60,441 accessory genes and 13,094 specific genes of Trametes were obtained. The distribution of accessory genes varied from 17,587 to 31,856. Although T. cinnabarina FP104138-Sp was the smallest (Table 1) among the nine Trametes strains, it had the maximum number (31,856) of accessory genes. Meanwhile, T. hirsuta had the minimum number (17,587) of accessory genes. The number of specific orthologs of the nine Trametes strains ranged from 368 to 4698. Trametes villosa CCMB561 possessed the highest number (4,698) of specific genes, whereas T. cinnabarina FP 104138-Sp had the minimum number (368) of specific genes (Figure 1(b)). The distribution of accessory and specific genes revealed that Trametes genomes are considerably diverse.

3.2. Phylogenetic analysis of Trametes pan-genome

The nuclear ribosomal internal transcribed spacer (ITS) region was the primary fungal barcode marker [26]. In addition, ITS was useful in identifying the broadest range of fungi [27]. A phylogenetic tree for Trametes and C. caperata, which was selected as an

| Organism                     | DB accession number | Contigs/scaffolds | Genome size (bp) | CDSs/ORFs* | Reference |
|------------------------------|---------------------|-------------------|------------------|------------|-----------|
| T. cinnabarina BRFM137       | CCBP010000001       | 776               | 33,668,918       | 59,163     | [39,40]   |
| T. cinnabarina FP 104138-Sp  | J5YU01000001        | 9,648             | 31,621,284       | 61,560     | [41]      |
| T. cocinea BRFM310           | KZ040086            | 222               | 32,757,859       | 56,735     | [42]      |
| T. hirsuta                   | CP019371            | 13                | 37,433,966       | 61,150     | [43]      |
| T. polyzona                  | MKK0010000001       | 5,321             | 36,629,716       | 65,446     | [44]      |
| T. pubescens                 | MNAD010000001       | 1,731             | 39,739,821       | 67,916     | [45]      |
| Trametes sp. AH28-2          | LJUJ01000001        | 306               | 38,904,488       | 65,191     | [46]      |
| T. versicolor FP-101664 S51  | NW_007360321        | 2,783             | 44,794,008       | 71,443     | [47]      |
| T. villosa CCMB561           | PUDQ01000001        | 10,327            | 57,987,392       | 101,817    | [4]       |

*The CDSs/ORFs were predicted by Prodigal.
outgroup to root the topology of the tree, was built based on their ITS1 and ITS2 sequences, respectively, to analyze the phylogenetic relationships for the nine *Trametes* strains. The ITS1 and ITS2 sequences had a strong power to differentiate the nine *Trametes* strains (Figure 1(c,d)). Among the phylogenetic trees, *T. pubescens*, *T. versicolor* FP-101664 SS1, and *T. sp. AH28-2* had a close evolutionary relationship, and *T. cinnabarina* BRFM137 had a close evolutionary relationship with *T. coccinea* BRFM310. However, the evolutionary relationship of *T. hirsuta* was distant from that of the remaining strains of *Trametes* (Figure 1(c,d)).

Another phylogenetic tree was built based on the presence and absence of genes in pan-genome to gain insights into the evolutionary relationships among the strains of *Trametes*. The results showed that *T. villosa* had a close evolutionary relationship with *T. versicolor* FP-101664 SS1. Meanwhile, *T. sp.AH28-2*, *T. pubescens*, *T. coccinea* BRFM310, and *T. cinnabarina* FP 104138-Sp had a close evolutionary relationship with one another (Figure 1(e)), revealing that they had similar genetic composition and functional metabolism. In addition, the evolutionary relationship of *T. hirsuta* was distant from the rest of *Trametes* strains (Figure 1(e)), suggesting that the functional composition of *T. hirsuta* was different from that of the remaining *Trametes* strains, and the functional metabolism of *Trametes* was highly diverse.

3.3. Functional characterization of the *Trametes* pan-genome

The *Trametes* pan-genome against the COG database was annotated to characterize the functions of the accessory clusters, core clusters, and specific genes and to investigate the functional diversity of the nine *Trametes* strains. The orthologous genes of the pan-genome were primarily dominant in...
“general function prediction only” (R, 1,758, 15.47%); “replication, recombination, and repair” (L, 998, 8.78%); “amino acid transport and metabolism” (E, 996, 8.77%); “translation, ribosomal structure, and biogenesis” (J, 755, 6.64%); and “carbohydrate transport and metabolism” (G, 753, 6.63%, Figure 2(a)). Moreover, Trametes had 442 (3.89%) orthologous genes assigned to “secondary metabolite biosynthesis, transport, and catabolism” (Q, Figure 2(a)). Similarly, the accessory genome was enriched in “general function prediction only” (R, 1,411, 15.29%); “replication, recombination, and repair” (L, 810, 8.78%); and “amino acid transport and metabolism” (E, 799, 8.66%, Figure 2(a)). In addition, the core gene clusters were predominantly composed of genes involved in “general function prediction only” (R, 88, 20.09%), “energy production and conversion” (C, 51, 11.64%), “amino acid transport and metabolism” (E, 38, 8.68%), and “carbohydrate transport and metabolism” (G, 36, 8.21%, Figure 2(a)). Finally, specific genes were primarily contributed to the functions involved in “replication, recombination, and repair” (L, 610, 28.44%) and “general function prediction only” (R, 259, 12.07%, Figure 2(a)). The high proportions of accessory genes (9230, 81.23%) and specific genes (2145, 18.88%) in each strain of Trametes might lead to diverse functions.

Gene Ontology (GO) analysis was conducted to characterize the genetic functions of the
pan-genome and explore the evolution of function catalogs. These genes were categorized on the basis of the biological process, cellular component, and molecular function. The results showed that a great number of genes of pan-genome were involved in various enzymes, metabolic pathways, and biological processes. In particular, enrichment analysis of biological processes showed that many genes were classified into the following three categories: biosynthetic process (2265), response to stress (2186), and small molecule metabolic process (1882, Figure 2(b)). In addition, 606 orthologous genes contributed to the carbohydrate metabolism (Figure 2(b)). The classification based on the molecular function showed that the majority of genes were grouped into binding (such as ion binding, DNA binding, and RNA binding) and enzymes (such as ATPase activity, oxidoreductase activity, and kinase activity). However, various genes were also enriched in other categories, such as transporter activity and enzyme regulator activity (Figure 2(b)). Enrichment analysis of cellular component showed that high proportions of genes were grouped on the basis of their functions, such as cell (8492), intracellular (8386), and cytoplasm (7363, Figure 2(b)). Among these categories, accessory clusters were highly represented in these functions than core and specific genes (Figure 2(b)). Therefore, the diversity of accessory clusters might influence their functional diversity, allowing these strains to utilize the resources of the surrounding environment and adapt to the environment.

3.4. Identification of CAZymes for Trametes

White-rot basidiomycetes had the ability to decompose lignin most efficiently [48], and fungi were CAZymes and lignin-degrading enzymes for the degradation of lignocellulose [49]. *Trametes*, as one of the branches of white-rot basidiomycetes, and its members, such as *T. villosa* [48] and *T. gibbosa* BRFM 952 [50], had been reported that these strains had an unexpected high activity on lignin or crystalline cellulose. A previous study had shown that CAZymes played a central role in the degradation of glycoconjugate, oligosaccharide, and polysaccharide [37]. Hence, we systemically identified the *Trametes* pan-genome against the carbohydrate-active enzymes database (CAZY database) and grouped CAZyme genes into different CAZyme families and carbohydrate-binding modules (CBMs) on the basis of family-specific Hidden Markov Models (HMMs) to obtain systematic understanding of CAZymes of *Trametes* [51]. A total of 280 orthologous genes (Figure 3(a)) were identified and grouped into 87 CAZyme families. A large proportion of orthologous genes were cataloged into GHS (35.36%), glycosyltransferases (21.07%, GTs), and CEs (13.57%, Figure 3(a)). GH hydrolyzed the glycosidic bond between two or more carbohydrates [37], contributed most of the catalytic enzymes, and was involved in lignocellulosic degradation [52]. A previous study identified 237 GHS, 78 GTs, 12 polysaccharide lyase (PLs), 69 CEs, and 112 auxiliary activity (AAs) in *T. villosa* CCMB561 [4]. Moreover, the number of CAZymes of accessory clusters was higher than that of core clusters and specific genes (Figure 3(a)), and the number of genes belonging to different CAZymes in each *Trametes* strain (Figure 3(b)) was different. This finding suggested a remarkable difference in the ability of strains of *Trametes* for carbohydrate degradation.

The 87 CAZyme families were subdivided on the basis of accessory clusters, core clusters, and specific genes to gain insights into the CAZyme composition of the nine *Trametes* strains (Figure 3(c–e)). For the orthologous genes of accessory clusters, a slice of CAZyme families, such as AA9, CBM1, CE10, CE16, CE4, GH10, GH16, GH18, GH5, GH79, GT2, GT20, GT76, and PL14, were detected in all nine *Trametes* strains (Figure 3(c)). Among these CAZyme families, the number of CE10 (ranging from 5 to 20) and GH16 (ranging from 6 to 15) were higher than other CAZyme families (Figure 3(c)). A previous study reported that CE10 family includes various esterases [36]. GHs could hydrolyze glycosidic bonds among carbohydrates or between a carbohydrate and a non-carbohydrate moiety [36]. In addition, GH16 comprises a large of glycosidases and transglycosidases, which catalyzed a range of terrestrial and marine polysaccharides [53]. The counts of AA2 of *T. cinnabarin*a BRFM137 (14) and CBM21 of *T. villosa* CCMB561 (24) was higher than that of other strains (Figure 3(c)). By contrast, the distributions of CAZyme families of core clusters, such as AA2, CBM1, CBM14, CBM50, GH18, GH3, GH37, and GT90, in *Trametes* strains were similar (Figure 3(d)). Moreover, the orthologous genes of specific genes were assigned to different CAZyme families (Figure 3(e)). For example, 14 CAZyme families, including GH18 and GT4, were dominant in *T. villosa* CCMB561, and CE10 and GH16 were enriched in *T. sp.* AH28-2. Notably, AA7 was only detected in *T. villosa* CCMB561; CE5 was only identified in *T. versicolor* FP-101664 SS1; CH6 only appeared in *T. pubescens*; and CH89 was only detected in *T. hirsuta*. The unique CAZyme family and the differential composition of CAZymes in *Trametes* revealed their distinct ability to metabolize carbohydrates. In summary, the composition of CAZyme families of the nine *Trametes* strains...
showed that these strains had the ability to metabolize the same carbohydrates and to degrade different carbohydrates.

### 3.5. Identification of Trametes BGCs

AntiSMASH was applied in predicting BGCs to further understand the secondary metabolite of *Trametes*. Our results showed that the BGCs of *Trametes* were primarily divided into six strategies, such as putative cluster of unknown type identified with the ClusterFinder algorithm (cf_putative), terpene cluster (terpene), nonribosomal peptide synthetase cluster (nrps), putative fatty acid cluster identified with the ClusterFinder algorithm (cf_fatty_acid), type I PKS cluster (t1pks), and lanthipeptide cluster (lantipeptide, Figure 4). In particular, the nine *Trametes* strains harbored 47.78 BGCs on average, and the number of BGCs of *Trametes* ranged from 18 (T. cinnabarina FP 104138-Sp) to 66 (T. sp. AH28-2, Figure 4). Clusters of cf_putative and terpenes represented the majority of predicted specialized metabolite BGCs and contributed to the highest share among the nine *Trametes* strains (Figure 4). The results showed that the clusters of cf_putative were enriched in the nine *Trametes* strains (ranging from 3 to 39 with an average of 25 BGCs, Figure 4), whereas *T. cinnabarina* FP 104138-Sp harbored only three clusters of cf_putative. When comparing the BGCs of terpene, *T. hirsuta* harbored 17 BGCs, and *T. cinnabarina* BRFM137 and *T. cinnabarina* FP 104138-Sp all harbored nine BGCs. Moreover, the majority of these identified terpene BGCs could not be linked to

Figure 3. Distributions of CAZymes in nine *Trametes* strains of *Trametes*. (a) Distribution of CAZymes in *Trametes* pan-genome; (b) Distribution of CAZymes of pan-genome in each *Trametes* strain grouped by accessory clusters, core clusters and specific genes. The numbers shown in subgraph represents the orthologous genes assigned by CAZymes and grouped by accessory clusters (pink), core clusters (green), and specific genes (wathet); (c) Distribution of CAZymes of accessory clusters in each *Trametes* strain; (d) Distribution of CAZymes of core clusters in each *Trametes* strain; (e) Distribution of CAZymes of specific genes in each *Trametes* strain. The numbers shown in subgraphs represent the orthologous genes assigned by CAZymes. GT: glycosyltransferase; GH: glycoside hydrolase; CE: carbohydrate esterase; CBM: carbohydrate-binding molecule; AA: auxiliary activities; PL: polysaccharide lyases.
known terpene BGCs. In addition, many predicted BGCs were grouped into "other" of these nine strains.

Although the secondary metabolites of predicted BGCs were unknown, we identified a few enzymes as the core biosynthetic enzymes, such as cembrene C synthase, pristinol synthase, (±)-epi-cubebol synthase, and epi-isozizaene synthase, in several BGCs in most strains of Trametes. To gain insights into the BGCs of Trametes, three BGCs of T. pubescens were chosen as examples (Figure 5). In particular, the BGCs of No. 13 were detected by the ClusterFinder algorithm built-in antiSMASH with 81.47% probability, and the homologous gene clusters were identified from T. versicolor FP-101664 SS1 and Dichomitus squalens LYAD-421 SS1 with 75% and 25%, respectively, of genes showing similarity (Figure 5(a)). In addition, the monomer of the predicted core structure of the substrate was ala-ala (C_6H_{12}O_2N_2). The BGCs of No. 15 was detected as other unknown cluster with 83.63% probability, and the homologous gene clusters were identified from T. cinnabarina BRFM137 and T. versicolor FP-101664 SS1 with 75% and 25%, respectively, of genes showing similarity (Figure 5(b)). In addition, the monomer of the predicted core structure of the substrate was ala-ala (C_6H_{12}O_2N_2). The BGCs of No. 20 was also detected as other unknown cluster, and the homologous gene clusters were identified from T. versicolor FP-101664 SS1 and T. cinnabarina BRFM 137 with 93% and 56%, respectively, of genes showing similarity (Figure 5(c)). In summary, these results revealed that many BGCs of Trametes were undiscovered, and Trametes had a potential value for producing secondary metabolites.

4. Conclusion

In conclusion, we systemically comparative analyzed the Trametes pan-genome, one of important branches of white-rot basidiomycetes, based on the available genome of nine Trametes strains and we explored the functional composition of nine Trametes strains, including COG annotation, GO annotation, and CAZyme families. Furthermore, we predicted the BGCs for each strain of Trametes. Our comparative analysis revealed that Trametes exhibited an open pan-genome structure and tremendous diversity in both the number and the variety of genes. The results of comparative analysis of Trametes pan-genome also showed the differences in the composition of functional metabolisms and the evolutionary relationships among the nine Trametes strains. In addition, we observed that the high proportion of accessory genes and specific genes annotated with different COG annotations and GO annotations showed that the diversity of the functions among the Trametes strains of Trametes. Importantly, our results showed that the CAZyme families, such as GH, GT, and CE, enriched in the Trametes strains, while the number of CAZymes of accessory clusters is higher than that of core clusters and specific genes and the number of genes belonged to different CAZymes in each Trametes strain suggested that the potential ability to degrade different carbohydrates for different strains of Trametes. Finally, we predicted many BGCs with unknown functions in Trametes, which suggested that Trametes has a great potential value for producing secondary metabolite. Together, we uncover the genetic
(a) MNAD01001604.1-Gene Cluster 13. Type = nrps. ClusterFinder probability: 0.8147

(b) MNAD01001612.1-Gene Cluster 15. Type = other. ClusterFinder probability: 0.8363

(c) MNAD01001649.1-Gene Cluster 20. Type = other.

Figure 5. Probability and homologous gene clusters for three putative BGCs of *T. pubescens*. The probability and homologous gene clusters for the putative BGCs of (a) no. 13; (b) no. 15; (c) no. 20. In the subgraph, the same colors of the boxes with arrows are similar genes.
diversity and synthetic biology of secondary metabolite production of *Trametes* in this work. Our results provide a broader understanding of *Trametes* and importantly shed the lights on the composition of CAZyme families and the BGCs of *Trametes*.

**Authors’ contributions**

This study was designed by YZ and ZDX. YZ collected and downloaded the data. YZ and YJC analyzed the data. YZ, JJW, and ZDX wrote the initial draft of the manuscript. All authors revised the manuscript.

**Disclosure statement**

No potential conflict of interest was reported by the author(s).

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**ORCID**

Yan Zhang http://orcid.org/0000-0001-9914-1302

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