Transcriptome Analysis of White-Rot Fungi in Response to Lignocellulose or Lignocellulose-Derived Material Using RNA Sequencing Technology

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Abstract
White-rot fungi are the only organisms that can completely degrade all components of lignocellulosic biomass, including the recalcitrant lignin polymer. Lignin degradation is important for the industrial application of lignocellulosic biomass as a raw material for producing value-added chemicals and materials. Therefore, elucidating the lignin degradation mechanism in white-rot fungi will help researchers develop efficient and eco-friendly methods enabling the production of value-added chemicals from lignocellulosic biomass. A transcriptome analysis is an effective way to compare gene expression patterns between different samples under diverse conditions and can provide insights into biological processes. The democratization of next-generation sequencing technology, especially RNA-sequencing, has made transcriptome sequencing and analysis a common research approach for many laboratories. In this review, we focus on the transcriptome profiles of two well-characterized white-rot fungi (Phanerochaete chrysosporium and Dichomitus squalens) in response to various lignocellulosic materials. The application of RNA-seq technology combining with other techniques remains the best approach for investigating fungal secretomes and elucidating the mechanisms of fungal responses to lignocellulose.

Keywords
Transcriptome, RNA-Seq, White-Rot Fungi, Lignocellulosic Biomass

1. Introduction
Lignocellulosic biomass, which is the most abundant polymer on Earth, consists...
of cellulose, hemicellulose, and lignin. There is increasing research focused on the utility of lignocellulosic biomass as a sustainable alternative to fossil fuel-based energy sources, biomaterials, and chemicals [1]. The two major industries producing the main sources of lignocellulosic biomass are forestry (e.g., sawdust, logging debris, and bark) and agriculture (e.g., rice, wheat straw, corn stover, and sugar cane bagasse) [1]. Cellulose, which is the major component of lignocellulosic biomass, is highly crystalline, because its glucose subunits are linked by β-1,4 glycosidic bonds. Hemicellulose is a heterogeneous polymer comprising xylose, arabinose, mannose, and galactose. Additionally, it is hydrolyzed more easily than cellulose because of its amorphous structure [2]. Lignin is also a heterogeneous polymer and is formed from the free radical coupling of methoxy-substituted coniferyl alcohol units [3]. Moreover, lignin represents up to 32% of plant dry matter [4], and it is the most recalcitrant cell wall component, providing rigidity as well as resistance against microbial infections, while also facilitating water transport [1]. Lignin is the most renewable source of aromatic compounds on Earth [5]. There are many applications for aromatics derived from lignin. For example, they may serve as precursors for the synthesis of biopolymers [6]. Despite its substantial potential as an industrial raw material, lignin remains the least utilized lignocellulosic biomass polymer and is currently being burned to generate heat and electricity [7]. The degradation of lignin is challenging because of its insolubility and complex, random structure with various non-hydrolyzable intramolecular C-C, C-O, and β-aryl ether bonds [8]. Lignin is currently fragmented via physical or chemical methods, which generate several aromatic compounds such as coumaric acid, hydroxybenzoic acid, ferulic acid, and vanillic acid [9]. The development of an environmentally friendly method for modifying lignin is a high priority for researchers because of the utility of lignin in a thriving bio-based economy.

Although diverse microorganisms can decompose lignocellulose, basidiomycetous white-rot fungi are the only organisms that can degrade all lignocellulosic biomass components, because they produce a variety of extracellular hydrolytic and oxidative enzymes, most of which have been categorized in the Carbohydrate-Active Enzyme (CAZy) database [8] [10]. The resulting monomer sugars serve as carbon and energy sources, and are taken up by the fungal cells through specific pathways [11]. To degrade lignin, white-rot fungi secrete an array of oxidoreductases from the CAZy auxiliary activity (AA) family of enzymes. The key enzymes in this family are fungal class II peroxidases, including lignin peroxidases (LiPs), manganese peroxidases (MnPs), and versatile peroxidases (VPs) [12]. Additionally, laccases, which are phenol-oxidizing multicopper oxidases, help peroxidases degrade lignin in the presence of aromatic compounds [12]. Furthermore, excessive amounts of H₂O₂-generating enzymes, such as glucose methanol choline (GMC), alcohol oxidases (AOXs), aryl alcohol oxidases (AAOs), glucose 1-oxidases (GLXs), and copper radical oxidases (CROs), contribute to the lignin-degrading system of white-rot fungi [13]. In addition to lignin-degrading enzymes, white-rot fungi secrete cellulases and hemicellulases.
from glycoside hydrolase (GH) families to completely depolymerize cellulose and hemicellulose [13]. Two other types of enzymes, lytic polysaccharide monooxygenases (LPMOs) and cellobiose dehydrogenases (CDHs), also participate in the degradation process (Figure 1) [14] [15]. *Phanerochaete chrysosporium* mainly colonizes in hardwoods [16], whereas, *Dichomitus squalens* are predominantly found on softwoods, but can also grow on hardwoods by tailoring their molecular responses [17]. The genome of *P.chrysosporium* lacks genes encoding laccase, and mainly produces LiPs and MnPs [16], but *D. squalens* produces predominantly MnPs and laccases in the lignin degradation process [18]. Both *P. chrysosporium* and *D. squalens* selectively degrade lignin and hemicellulose fraction and leave the cellulose fractions barely untouched [19] [20]. Many studies have been done on the application of *P. chrysosporium* in the biological pretreatment of lignocellulosic biomass (e.g., rice straw, cotton stalks, and wheat straws) and results showed greatly loss of lignin and reduction of the crystalline index of cellulose [21] [22] [23]. However, the application of *D. squalens* in the biological treatment of lignocellulosic biomass is not common compared to *P. chrysosporium*. This review mainly focuses on the published studies regarding the above-mentioned white-rot fungi and their transcriptome responses to lignocellulosic materials. Studies reviewed in this review were selected according to the process described in Figure 2.

**2. Transcriptome Analytical Approaches**

Transcriptomes refer to the complete set of genome sequences transcribed at a specific time-point by a given organism, organ, tissue, or cell [24]. In contrast to static genomic studies, transcriptome analyses are dynamic because they reflect the changes in gene expression due to developmental or environmental conditions [25]. Methods for analyzing transcriptomes, including those involving microarrays and RNA sequencing (RNA-seq), evolved from the massively parallel signature sequencing (MPSS) and serial analysis of gene expression (SAGE) [26].
Specifically, RNA-seq is a rapid and cost-effective revolutionary technique that exploits the capabilities of next-generation sequencing platforms to generate large numbers of high-quality short reads, thereby enabling comprehensive transcriptomics data mining [26] [27]. Transcriptome research is currently dominated by RNA-seq because it does not require an available reference genome. Furthermore, it can refine the annotation of potential reference genomes as well as identify unpredicted transcribed regions and detect gene splicing variants [28]. The RNA-seq method is based on the sequencing of a considerable abundance of cDNA synthesized from mRNA obtained from the sample under study. The protocol for sequencing the normalized cDNA library varies depending on the specific platform (e.g., Illumina system). Regardless of the system, the raw sequencing data (i.e., millions of reads) undergo a quality control step and the reads are trimmed to obtain clean data. The processed reads are subsequently mapped to a reference genome or a transcriptome, which can be created de novo from the reads if a reference genome is unavailable. The expression of genes can be quantified according to the frequency of reads mapped to the same gene. Reads per thousand nucleotides per million mapped reads (RPKM) or fragments per thousand nucleotides per million mapped reads (FPKM) are frequently used to quantify gene expression levels [28]. Once the expression level information is available, the differentially expressed genes may be analyzed with specific statistical tests. Other downstream analyses may also be performed, including gene annotations, gene set enrichment analyses, or pathway analyses (Figure 3). The RNA-seq approach is especially useful because most white-rot fungal genomes are not completely sequenced and annotated. Moreover, this technique can detect novel genes as well as generate information regarding previously uncharacterized genes. For example, Oghenekaro et al. [29] sequenced and de novo assembled the transcriptome of the white-rot fungus
Rigidoporus microporus, ultimately producing 25,880 annotated unigenes. Their data revealed that the fungus expresses more than 300 genes encoding lignocellulolytic enzymes, with rubberwood upregulating the expression of 175 genes.

3. White-Rot Fungi

3.1. Phanerochaete Chrysosporium

The white-rot fungus *P. chrysosporium* is one of the most extensively studied fungi because of its ability to degrade lignocellulosic biomass [30]. It secretes a wide range of oxidative and hydrolytic enzymes to degrade lignin and a variety of organic compounds, including 2,4-dichlorophenol, 2,4-dinitrotoluene, and endosulfan [31]. Martinez et al. [32] fully sequenced and annotated the *P. chrysosporium* genome, which has considerably advanced our understanding of the wood decay mechanism of this fungus. During the last few decades, genome-level transcriptome studies have revealed several important facts about the differential expression patterns of genes involved in lignin depolymerization. Most transcriptome analyses conducted before 2015 involved microarrays. Table 1 summarizes the sample conditions and analytical approaches used in the five *P. chrysosporium* transcriptome studies described below. Wymelenberg et al. [33] performed secretome and transcriptome analyses using microarrays and liquid chromatography-tandem mass spectrometry (LC-MS/MS) to determine the number, structure, and regulation of genes involved in lignocellulosic cell wall degradation. They identified numerous carbohydrate-active genes, including GH 61 family members, polysaccharide lyases, and carbohydrate esterases. Also, they found that other upregulated genes were involved in the hemicellulose degradation including various xylanases, mannanase, α- and β-galactosidases, polygalacturonase, arabinofuranosidase, xyloglucanases and feruloyl esterase. More than 190 upregulated genes were predicted to encode proteins of...
Table 1. Details of the sample conditions and RNA-seq data analysis approach in *P. chrysosporium* transcriptome analysis.

| Platform   | Substrate            | Sample condition                                      | RNA-seq data analysis                                                                 | Reference |
|------------|----------------------|-------------------------------------------------------|--------------------------------------------------------------------------------------|-----------|
| Roche      | microcrystalline cellulose | Three biological replicates; total RNA was extracted from frozen fungi pellets after 5 days incubation at 37°C | NimbleScan v2.4 and ArrayStar v2.1 software was used to quantify and visualize data. Quantile normalization and robust multi-array averaging were applied to the entire data set. Expression levels are based on log2 values. | [33]      |
| Roche      | ball-milled aspen    | Three biological replicates; total RNA was extracted from frozen fungi pellets after 5 days incubation at 37°C | DNAStar ArrayStar v2.1 software was used to quantify and visualize data. Quantile normalization and robust multiarray averaging were applied to the entire data set. Expression levels are based on log2 values. T-test with a false discovery rate threshold at $P < 0.001$ was used to determine significant differences in expression. | [34]      |
| Illumina   | Spruce wood          | Three biological replicates; total RNA was extracted from samples at 40 h and 96 h incubation at 38°C | DNAStar Inc. modules SeqNGen and Qseq were used for mapping reads and statistical analysis. The current Joint Genome Institute (JGI) annotation (v2.2), served as the queried database RNA-seq-based transcript results are presented as RPKM values | [35]      |
| Illumina   | Maple wood and miscanthus | Three biological replicates; RNA was extracted from mycelium after 5 weeks cultivation at room temperature. | ArrayStar program of DNASTAR software was used to process raw data sequences, and annotation was done by using the *P. chrysosporium* RP-78 v2.2 from the Joint Genome Institute (JGI). RNA-seq transcript-based results were reported as RPKM values | [36]      |
| Illumina   | Helianthus argophyllus (silverleaf sunflower) stem | Three biological replicates; RNA was extracted from samples after 6 weeks incubation at room temperature. | ArrayStar program of DNASTAR software was used to process the raw data sequences, the annotation was done using *P. chrysosporium* RP-78 v2.2 and Gene Ontology (GO) file (Phchr 2 _GeneCatalog_proteins _20131210 _GO.tab.gz) from the Joint Genome Institute (JGI). NCBI nucleotide BLAST was further used to annotate transcripts of interests. The data were represented as RPKM values. | [37]      |

unknown function, among which, approximately one-third contain predicted secretion signals. Additionally, 54 encoded proteins were detected in extracellular filtrates. Researchers subsequently compared the gene expression patterns of the white-rot fungus *P. chrysosporium* and the brown-rot fungus *Postia placenta* colonized on aspen [34]. This earlier investigation proved that the oxidoreductase-encoding genes have distinct expression patterns in brown-rot and white-rot fungi. *P. chrysosporium* was observed to secrete an array of extracellular glycosyl hydrolases to simultaneously degrade cellulose and hemicellulose. In contrast, *P. placenta* secreted various hemicellulases, but few potential cellulases, under the same conditions. *P. chrysosporium* identified 35 significantly differential expressed genes in ball-milled aspen (BMA), among which, 22 genes identified as glycosyl hydrolase family members, including highly expressed upregulated genes encoding exocellobiohydrolase GH7 and exocellobiohydrolase GH6. However, in *P. placenta*, only 5 glycoside hydrolase-encoding genes were upregulated in BMA, while a broadly defined hemicellulose (e.g., endo-$\beta$-1,4-mannosidases, endoxylanases, mannosidases, and $\beta$-xylosidase) genes were highly expressed in BMA. Furthermore, the genes related to iron-acquisition systems were differentially expressed between these two species. In *P. placenta*, the expression levels of genes encoding ferroxidase and iron permease were sub-
stantially upregulated, which is in contrast to the downregulated expression of these genes in *P. chrysosporium*. Korripally *et al.* [35] characterized the changes in gene expression during the transition to ligninolytic metabolism using fluorometric oxidant-sensing beans in conjunction with a whole-transcriptome shot-gun sequencing analysis. This study revealed 356 upregulated genes and 252 downregulated genes. The upregulated genes encoding LiPs, MnPs, and auxiliary enzymes, and suggested that the extracellular oxygen species involved in ligninolytic reactions are generated from MnP-catalyzed lipid peroxidation, CDH-catalyzed Fe³⁺ reduction, and oxidase-catalyzed H₂O₂ production. Moreover, this study identified 72 unknown protein-encoding genes with upregulated expression at 96 h, including 27 putative transporter genes and 18 cytochrome P450 genes as candidates for future studies on lignin fragment uptake and processing. Alaradi [36] examined the transcriptome of *P. chrysosporium* after 5 weeks of growth on maple (hardwood) and miscanthus (energy grass). In this study, *P. chrysosporium* responded uniquely to maple and miscanthus, with 55 and 66 differentially expressed genes identified when samples growing on maple and miscanthus were compared with those growing under control conditions, respectively. Many of the identified genes are involved in the breakdown of lignocellulose, including LiP, MnP, and GH genes. However, more than half of these genes are not well annotated regarding their effects on maple and miscanthus. Moreover, malate synthase, lactate dehydrogenase, and phosphoenolpyruvate carboxykinase genes were more highly expressed in fungi grown on miscanthus than in fungi grown on maple. Alsubaie [37] investigated the *P. chrysosporium* transcriptome by culturing the fungus on sunflower stems after 6 weeks of growth. In this study, 102 genes were identified differentially expressed when compared to control condition, the majority of these genes were tied to lignocellulose degradation including peroxidases, oxidases, and glycoside hydrolases, however, up to 75% of the differentially expressed genes were classified as hypothetical or unknown.

### 3.2. *Dichomitus Squalens*

*Dichomitus squalens* is a white-rot basidiomycete that degrades cellulose and lignin by secreting hydrolytic and oxidative enzymes. The *D. squalens* genome includes genes predicted to encode four cellobiohydrolases (CBHs), three putative endoglucanases, six putative β-glucosidases, one putative CDH, and 15 LPMOs [38], as well as other genes encoding enzymes involved in lignocellulose degradation (e.g., nine MnPs and 11 laccases) [39]. The ability to secrete diverse extracellular lignocellulose-modifying enzymes has made *D. squalens* an excellent model fungus for studying lignocellulose degradation. Research has been conducted on the molecular response of *D. squalens* to lignocellulose or lignocellulose-derived compounds as well as the underlying regulatory system. Table 2 summarizes the sample conditions and analytical approaches used in the five *D. squalens* transcriptome studies described below. Rytioja *et al.* [39] analyzed 10 genes encoding cellulose-acting enzymes in *D. squalens* on solid-state spruce.
Table 2. details of the sample conditions and RNA-seq data analysis approach in *D. squalens* transcriptome analysis.

| Platform                  | Substrate                                      | Sample condition                                                                 | RNA-seq data analysis                                                                 | Reference |
|---------------------------|-----------------------------------------------|-----------------------------------------------------------------------------------|--------------------------------------------------------------------------------------|-----------|
| Maxima SYBR Green qPCR Master Mix and ABI apparatus (Applied Biosystem) | Spruce wood, Microcrystalline cellulose (Avicel) | Three biological replicates; total RNA was extracted from fungal mycelia growing on spruce wood at 28°C at 7, 14, 21, and 28 days; total RNA was extracted from fungal mycelia growing with Avicel cultures at 14 and 28 days. | Differences in the gene expression levels were estimated according to the Shapiro-Wilk normality test at P ≥ 0.05. Normal distribution was done for genes *cel7b* and *lpmo1* based on two biological replicates of fungi growing with Avicel at 14 days. Repeated measures ANOVA was used to estimate the variation in the expression of one gene in different time points for spruce wood cultures and paired samples *t*-test for Avicel cultures at P-value < 0.05. Reads were mapped to the genome sequence of *D. squalens*.YAD-421 S1 (v1.0 annotation, the Joint Genome Institute (JGI)) using SOAPALIGNER/SOAP2. RPKM was used to quantify RNA-seq results. Differential expression was identified by CyberT Bayesian ANOVA algorithm at a cut-off value of the fold change of >1.5 and P-value (corrected by multiple tests) of <0.05. | [39]      |
| DNBseq Technology         | Aspen, spruce wood, wheat bran, cottonseed hulls | Total RNA was extracted from mycelia from two biological replicate cultures after 9 days and 16 days’ cultivation at 28°C | Reads were aligned to the reference genome (https://genome.jgi.doe.gov/Dicsqu464_1/Dicsqu464_1.home.html) using HISAT version 0.1.4-beta. Differentially expressed genes were identified using DESeq2 (version 1.10.0) with a cutoff value of adjusted p-value < 0.05. Raw gene counts were used for DGE analysis as DESeq2 uses its internal normalization. | [40]      |
| Illumina HiSeq 2000 platform | Norway spruce (Piceaabies), silver birch (Betula pendula) | Three biological replicates; total RNA was extracted from mycelium after 2 weeks and 4 weeks cultivation at 28°C | Reads were mapped to the genome sequence of *D. squalens* CBS 464.89 using Bowtie2 and the BWA software. The gene expression level was measured in FPKM using the RSEM tool. Differential expression was identified by the DESeq2 with a cutoff value of ≥2.5-fold change, FPKM value of ≥10, and the adjusted P-value of ≤0.01. CAZyme annotations were done using JGI MycoCosm website (https://genome.jgi.doe.gov/mycosom/proteins-browser/browse?LeIA4p=Dicsqu464_1). Gene expression levels were measured as FPKM. DESeq2 version 1.10.0 was used to compare the transcript level of samples. Differentially expressed genes were identified with fold change > 2, adjusted p < 0.01, and FPKM > 10 in at least one condition between each pair of conditions. Functional annotation of differentially expressed genes was based on combined information from EuKaryotic Orthologous Groups (KOG), Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway mapping, InterPro protein sequence analysis and classification, and Carbohydrate-active enzymes (CAZy) classifications for *D. squalens* CBS464.89 (Dicsqu464_1) retrieved from JGI MycoCosm database (https://genome.jgi.doe.gov/cgi-bin/LogBrowser?db=Dicsqu464_1). | [41]      |
| Illumina HiSeq 2000 platform | D-glucuronic acid, D-galacturonic acid, L-rhamnose, D-galactose, D-xyllose, D-mannose, L-arabinose, disaccharide cellobiose | The replication of samples was not mentioned in this study. Total RNA was extracted from the mycelium from the edge of the colony after 5 days of growth at 28°C | | [42]      |
| Illumina HiSeq 2500 platform | coniferyl alcohol, ferulic acid, vanillin, vanillyl alcohol, vanillic acid, veratryl alcohol, protocatechuic acid, p-coumaric acid, p-hydroxybenzoic acid, and cinnamic acid. | Three biological replicates; RNA was extracted from cultivation for 4 days at 28°C. | | [43]      |
wood and in microcrystalline-supplemented cultures. These genes included three (cel7a, cel7b, and cel7c) encoding CBHI, one (cel6) encoding CBHII, one encoding CDH, and five putative LPMO-encoding genes (lpmo1, lpmo2, lpmo3, lpmo4, and lpmo5). Additionally, cel7a was expressed constitutively and was the most abundant transcript, whereas the other cel genes were expressed at low levels in fungi growing on spruce wood and in Avicel. Moreover, lpmo genes encoding a cellulose-binding module (CBM) were observed to be more highly expressed than those lacking a CBM sequence when D. squalens was grown on spruce wood, whereas these genes were similarly transcribed in the liquid Avicel cultures. Although the CDH-encoding gene was co-expressed with the lpmo genes, its expression level was low in fungi grown on spruce wood and in Avicel cultures. Rytioja et al. [40] analyzed the transcriptome and exoproteome of D. squalens grown on two woody (aspen and spruce wood) and two non-woody plant biomasses (wheat bran and cotton seed hulls). Total 297 genes were identified as differentially expressed, of which, 135 genes were related to plant cell wall degradation. The number of expressed cellulolytic genes increased in both spruce and aspen, while decreased or not changed in non-woody plant biomass in the course of cultivation. The genes encoding ligninolytic enzymes, including AA2 lignin-modifying peroxidases and AA1 laccases, were mainly expressed on day 9 in the woody samples, whereas most cellulase gene expression levels were higher on day 16 than on day 9, implying the fungus degraded lignin first to make the cellulose accessible. The ligninolytic enzyme genes were expressed at lower levels in the non-woody plant biomass (i.e., wheat bran and cottonseed hulls) with low lignin contents than in the woody biomass. The genes for H$_2$O$_2$-generating enzymes, including AA3 alcohol/methanol oxidases and AA5_1 CROs, were expressed at similar levels in all substrates, suggesting these enzymes may contribute to the electron transfer system together with LPMOs degrading lignocellulose. A subsequent investigation of the ability of D. squalens to degrade softwood (spruce) and hardwood (birch) materials suggested the fungal molecular responses were more adapted to the softwood material, with a greater abundance of mannanolytic enzymes at the transcript and protein levels in the spruce culture due to the higher mannose content in spruce. Contrasting trends were observed that the genes encoding xylanolytic enzymes were higher expressed, but the activity of xylanolytic enzymes were low in spruce culture. Moreover, the expression level of laccases genes was higher in the birch cultures compared with the spruce cultures. In contrast to laccases, MnPs encoding genes were more abundantly produced in spruce culture [41]. López et al. [42] conducted a transcriptome analysis to examine six lignocellulose-derived sugars known to induce the expression of lignocellulolytic genes in ascomycetes to clarify their role as inducers in the basidiomycetous white-rot fungus D. squalens. When compared to D-glucose, 83 significantly upregulated genes were detected among all six lignocellulose-derived sugars, of which, 26 and 27 genes response exclusively to L-rhamnose and cellobiose, respectively. L-rhamnose and cellobiose were identified as the main inducers of cellulolytic and pectinolytic genes,
respectively. Moreover, lignin-degradation related enzyme encoding genes were induced only by L-rhamnose and D-xylsoe. Dikaryotic and monokaryotic strains of *D. squalens* cultivated on plant biomass-derived monosaccharides and the disaccharide cellobiose varied regarding gene expression patterns, with the largest difference observed on L-rhamnose. Dikaryotic strains showed higher upregulation for ligninolytic enzymes encoding genes than the monokaryotic strains, instead, the monokaryotic strains showed higher upregulation for pectinolytic genes than the dikaryon. Genes encoding mannanases and expansin-like proteins were only induced in dikaryotic strains of *D. squalens*. Accordingly, the fine-tuning of the gene regulation related to lignocellulose conversion may differ between these strains. Kowalczyk *et al.* [43] identified the various aromatic compounds generated by *D. squalens* cultivated on spruce wood. The *D. squalens* transcriptome-level differences induced by 10 different lignocellulose-related aromatic monomers were analyzed and 268 upregulated genes were identified. Vanillin upregulated the largest set of genes and most of the genes were associated with metabolism, followed by cinnamic acid upregulated the second large set of genes. The previous transcriptome analysis from mono- and dikaryotic strains demonstrated that L-rhamnose trigger the expression of ligninolytic enzymes and the lignin-rich substrate was the key factor that cause the induction of ligninolytic genes in *D. squalens* [40] [42]. Not as the same as in the previous studies, the lignin-degrading enzymatic machinery was only partially induced by the monomeric aromatic compound in Kowalczyk’s study. The expression levels of lignocellulose degradation-related genes as well as transporter-encoding genes and catabolic pathway genes differed significantly between the fungal samples supplemented with different aromatic compounds. The results also demonstrated that the regulon induced in the presence of aromatic compounds differs regarding the number and function of the genes. Vanillin, cinnamic acid, and p-coumaric acid were the main inducers of the genes annotated with gene ontology (GO) terms related to oxidoreductase activity, whereas the genes annotated with GO terms associated with lignocellulolytic activities were primarily upregulated by coniferyl alcohol, ferulic acid, and vanillyl alcohol.

4. Conclusion

In summary, because white-rot fungi can completely degrade all lignocellulosic biomass components, they are potentially useful for producing various bio-based products. Clarifying the molecular mechanisms mediating the lignocellulose degradation by white-rot fungi may facilitate the application of these fungi for metabolic engineering and optimize the bioconversion of renewable biomass resources to chemicals. The application of RNA-seq technology to analyze transcriptomes has increased our understanding of gene expression and may be useful for revealing the expression patterns of previously unannotated and noncoding genomic regions. Combining transcriptome sequencing with other techniques (e.g., secretomics-, NMR-, and mass spectrometry-based analyses) remains the best approach for investigating fungal secretomes and elucidating the
mechanisms underlying fungal responses to lignocellulose.

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Conflicts of Interest

The authors declare no conflicts of interest regarding the publication of this paper.

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