

Supplementary Materials

Metabolic Rewiring in tea plants in Response to Gray Blight Disease Unveiled by Multi-Omics Analysis

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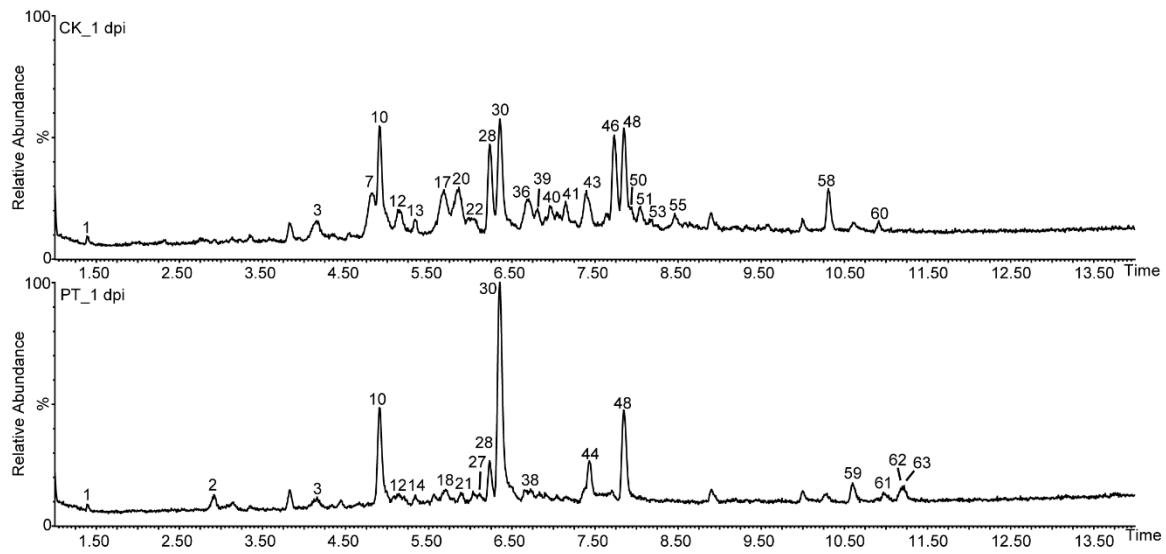


Figure S1. Representative total ion chromatograms of mock-inoculated and fungal-inoculated tea leaves at 1 dpi in the negative ionization mode. PT, tea leaves inoculated with *P. theae*. CK, tea leaves inoculated with the mock control. Peak numbering is the same as Table 1.

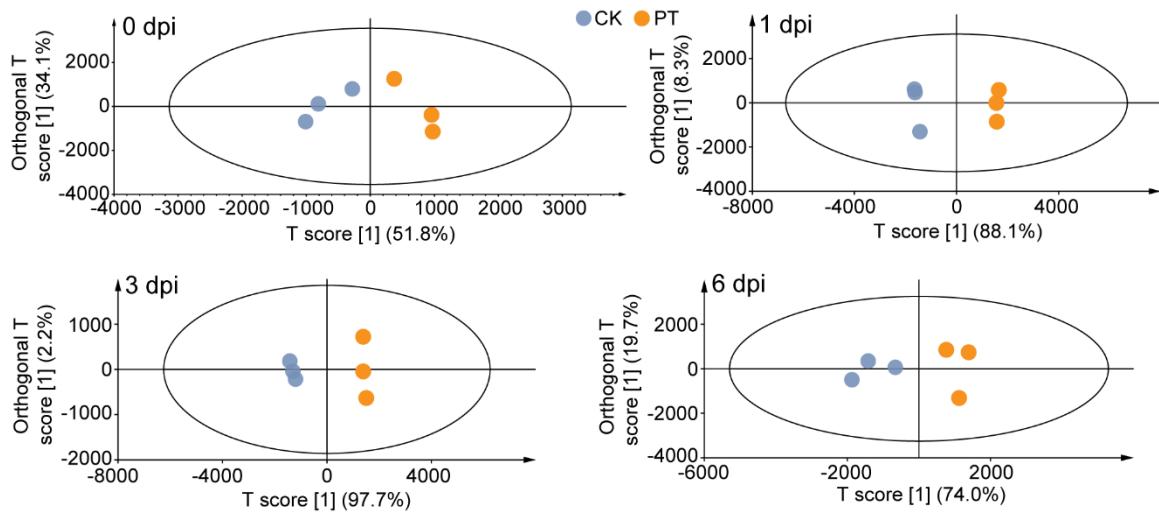


Figure S2. OPLS-DA analysis of the metabolomics datasets. PT, tea leaves inoculated with *P. theae*. CK, tea leaves inoculated with the mock control.

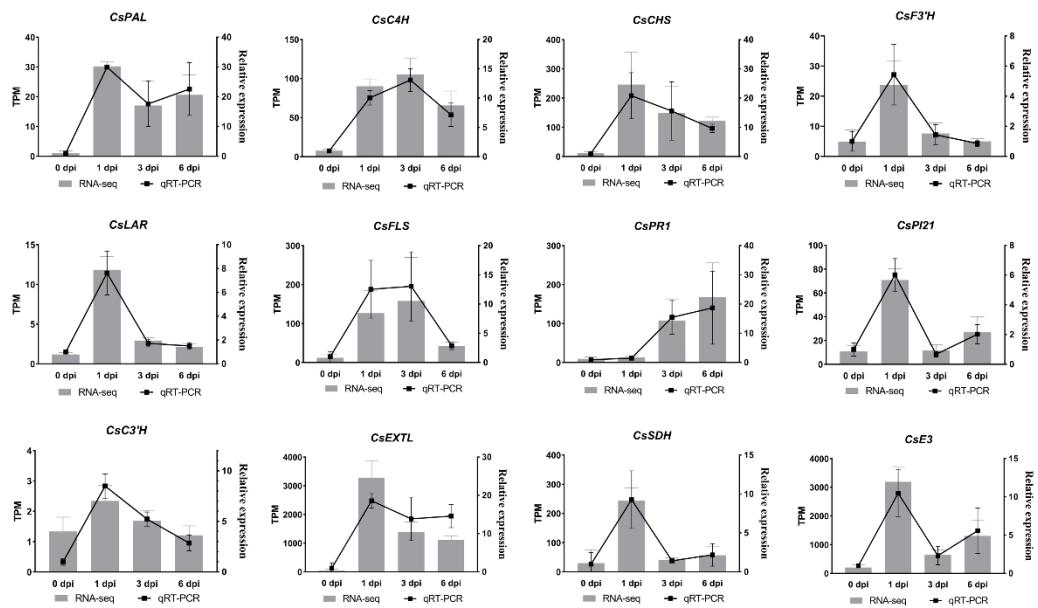


Figure S3. qRT-PCR validation of RNA-seq DEGs. PAL, phenylalanine ammonia-lyase; C4H, cinnamate-4-hydroxylase; CHS, chalcone synthase; F3'H, flavonoid 3'-hydroxylase; LAR, leucocyanidin reductase; FLS, flavonol synthase; PR1, pathogenesis-related protein 1; PI21, PYRICULARIA ORYZAE RESISTANCE 21 ; C3'H, 5-O-(4-coumaroyl)-D-quinate 3'-monooxygenase; EXTL, extension-like; SDH, shikimate dehydrogenase; E3, E3 ubiquitin ligase. The left Y-axis represents the expression data (gray bars) and the right Y-axis represents the relative expression values obtained through RT-qPCR analysis (black lines). *GADPH* gene (accession no. KA295375.1) was used as the internal control. The expression of genes in 0 dpi is set to 1.0. All values are mean \pm SD (n = 3).

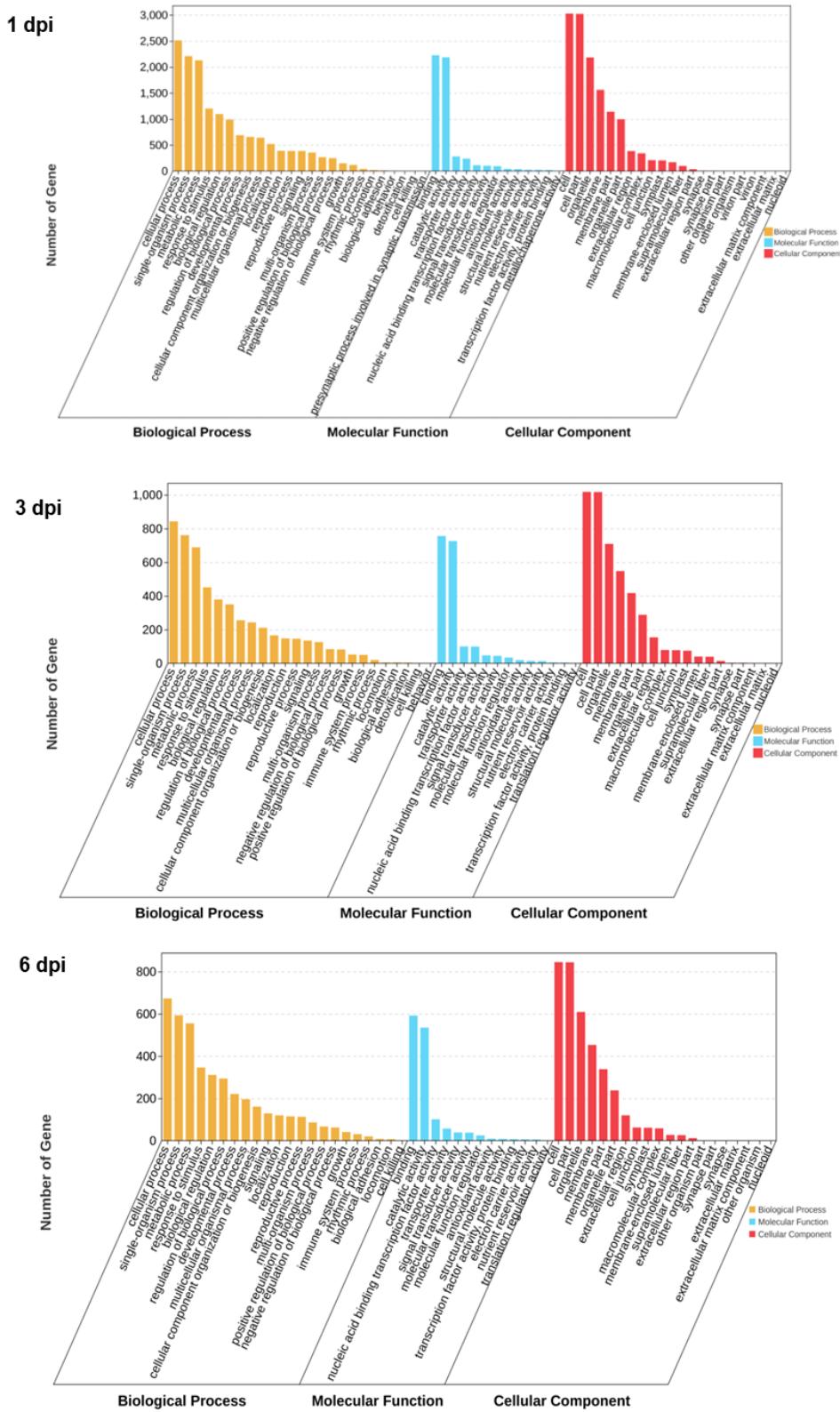


Figure S4. Gene ontology (GO) enrichment analysis of differentially expressed genes in gray blight-infected tea plants at 1, 3 and 6 dpi.

Table S1. List of primers for qRT-PCR.

| Target Gene ^a | Forward primer (5'-3') | Reverse primer (5'-3') |
|--------------------------|---------------------------|----------------------------|
| <i>CsPAL</i> | TCCAATTCCCTGCCAACCTGTAA | ATCAACTGCCTCGGCTGTCTTCT |
| <i>CsC4H</i> | TGCAAAGCTCTGGCTATG | CCTGAACCTCTCTGGGTTCTTC |
| <i>CsCHS</i> | GTGCTATCGATGGGCATCTT | AGGGTTGGAATGCCTCATTTA |
| <i>CsF3'H</i> | GTTTCGACCAGAACGATT CCTACC | ATAACTGGACCATA CGCAACCC TA |
| <i>CsLAR</i> | ACACTCCCTAGAGTAACCATCA | GATGAAGATGTCGTGGGTGAA |
| <i>CsFLS</i> | TGGAAAGACGGAATTGGGTAG | ACTCCTGTGAAGAACACTCTTG |
| <i>CsPR1</i> | TCCTATGCCAAA ACTCACACAAG | ACCCATCCTCGAACTAGCATA CCT |
| <i>CsPI21</i> | CTACAGGAATCCAAAGAGTCCAA | CATAGCATTGCCACAACATATC |
| <i>CsC3'H</i> | CGGCAACCTCTACGACATAAA | CAACTCTGAGCTCGAAACTATCA |
| <i>CsEXTL</i> | TGTCTCTGAGCTTGCCATTAG | GAGGTGGTGGTGATTGTAGT |
| <i>CsSDH</i> | GTATTGCTCCCATCTCCGCTCACTG | ATGACAACCCCTCGCTCCTCTGCTT |
| <i>CsE3</i> | GATGCTGCTACTCGGTTCTT | GGCATGACGGTTACATCATTTC |

^aPAL, phenylalanine ammonia-lyase; C4H, cinnamate-4-hydroxylase; CHS, chalcone synthase; F3'H, flavonoid 3'-hydroxylase; LAR, leucocyanidin reductase; FLS, flavonol synthase; PR1, pathogenesis-related protein 1; PI21, PYRICULARIA ORYZAE RESISTANCE 21 ; C3'H, 5-*O*-(4-coumaroyl)-D-quinate 3'-monooxygenase; EXTL, extension-like; SDH, shikimate dehydrogenase; E3, E3 ubiquitin ligase.

Table S2. The quality of RNA-seq data.

| Sample | Clean reads | Q30 | GC% | Mapped rate (%) |
|----------|-------------|-------|-----|-----------------|
| PT1_0dpi | 25805408 | 93.21 | 44 | 87.50 |
| PT2_0dpi | 26011094 | 92.30 | 45 | 88.18 |
| PT3_0dpi | 36586286 | 95.14 | 45 | 89.61 |
| PT1_1dpi | 25979638 | 89.29 | 43 | 86.66 |
| PT2_1dpi | 26575605 | 93.38 | 43 | 86.90 |
| PT3_1dpi | 26345386 | 91.98 | 43 | 86.09 |
| PT1_3dpi | 28349686 | 90.43 | 43 | 86.98 |
| PT2_3dpi | 26270282 | 92.31 | 43 | 86.00 |
| PT3_3dpi | 27623253 | 93.89 | 43 | 87.12 |
| PT1_6dpi | 28009516 | 91.86 | 43 | 86.84 |
| PT2_6dpi | 26367665 | 89.98 | 43 | 87.02 |
| PT3_6dpi | 26717807 | 91.03 | 44 | 87.32 |