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2Fig. 1 Biological identification of strain RHTA01. (A. Colony morphology of strain RHTA01; B.

3Microscopic morphology of strain RHTA01 (×40); C. Phylogenetic analysis based on the multiple

4sequence alignment of 16S rRNA sequences; D. Phylogenetic analysis based on the multiple sequence

5alignment of the SSU sequence; E. Phylogenetic analysis based on the multiple sequence alignment of the

6LSU sequence; F. Phylogenetic analysis based on the multiple sequence alignment of the RBP2 sequence.

7The number of the sequence in NCBI are C1904170071/M1.Contig1: MT374076,

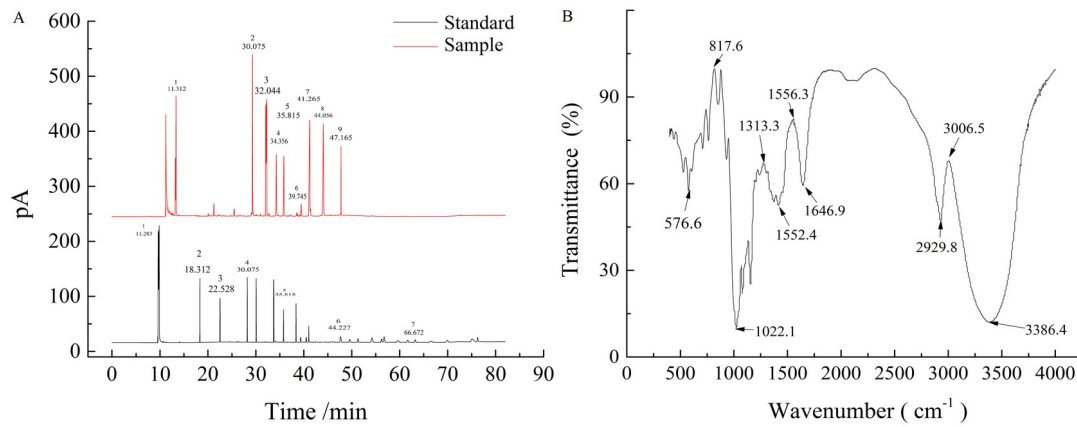
8C1904170071/M3.Contig1: MT374081, C1904170071/M4.Contig1: MT377726 and the sequence number

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9 of M6 is added later.)



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11 Fig. 2 Fatty acid analysis and functional group analysis of strain RHTA01. (A. Fatty acid analysis by gas
12 chromatography. Red curve marked is the sample and black curve is the standard; B. Infrared spectrum of
13 strain RHAT01 by FT-IR.)

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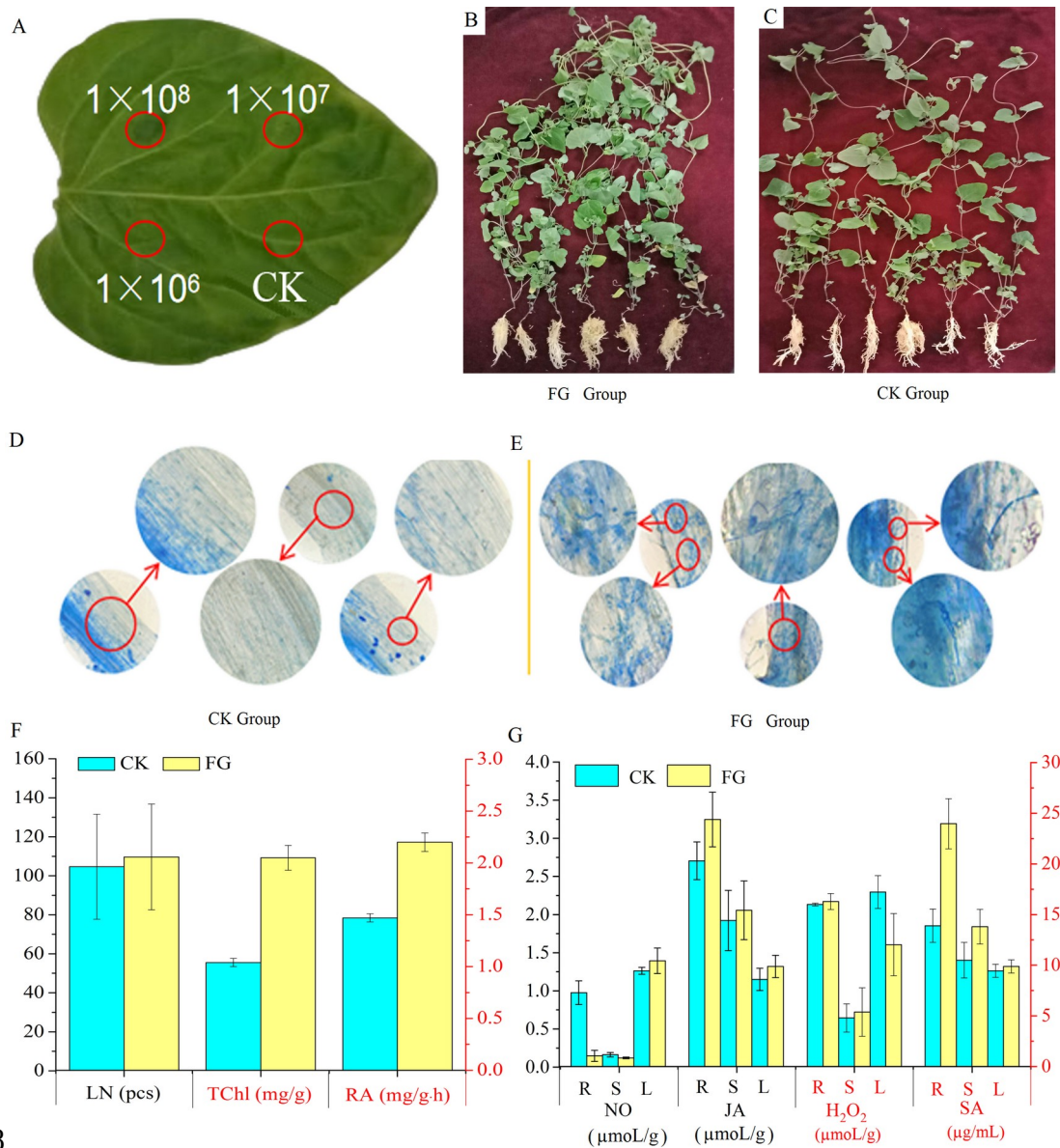
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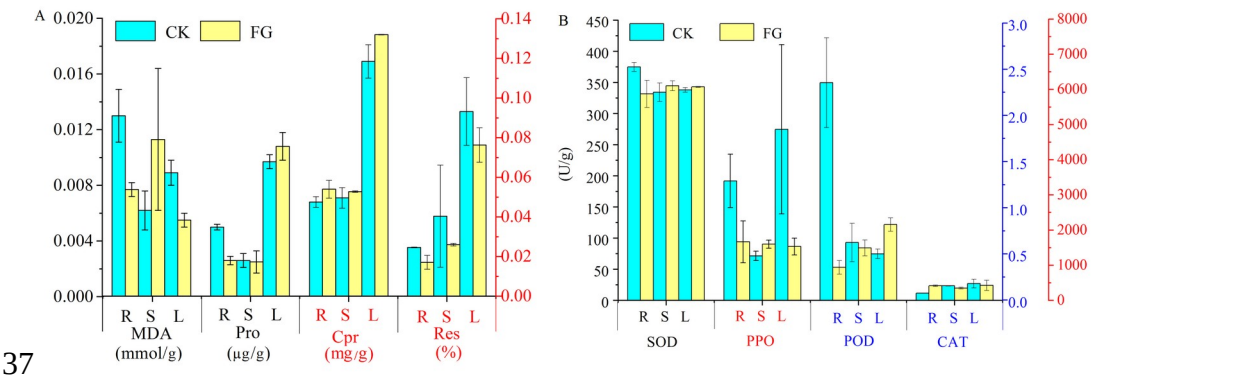
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29Fig. 3 Effect of strain RHTA01 on the Growth of *C. pilosula*. A. Biological tests of strain RHTA01
30inoculated with *C. pilosula* leaves; B. The growth of *C. pilosula* without the treatment by strain RHTA01
31(CK, control group); C. The growth of *C. pilosula* with the treatment by strain RHTA01 (FG, treatment
32group); D. Colonization determination of strain RHTA01 in the roots of *C. pilosula*.without the treatment
33by strain RHTA01 (CK, control group); E. Colonization determination of strain RHTA01 in the roots of *C.*
34*pilosula* treated with strain RHTA01 (FG, treatment group); F. Chlorophyll content (Tchl), root activity

35(RA) and Leaf number (LN) of *C. pilosula* in the control group and treatment group; G. Signal molecular

36(SA, JA, H₂O₂ and NO) content of *C. pilosula* in the control group (CK) and FG treatment group (FG).



38Fig. 4 The content of non-enzymatic ingredients (A) and various antioxidant enzymes activities (B) in
39different parts of *C. pilosula*. (A. The content of MDA, Pro, Cpr, Res in different parts of *C. pilosula* in
40the control group and the treatment group; B. The content of SOD, CAT, PPO, POD in different parts of *C.*
41*pilosula* in the control group and the treatment group.)

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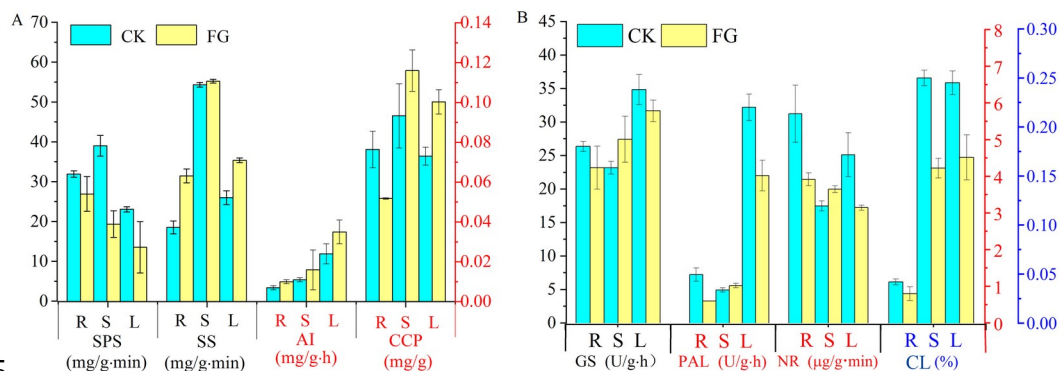
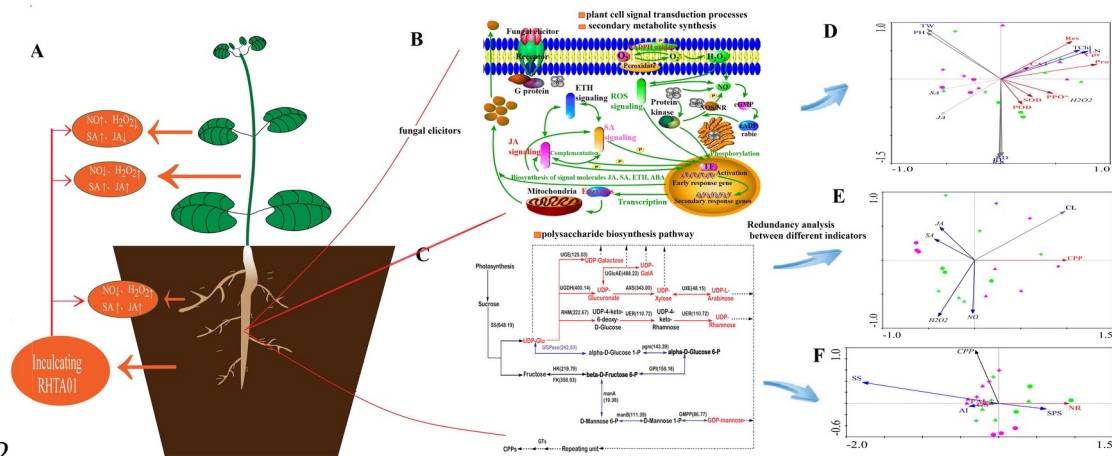
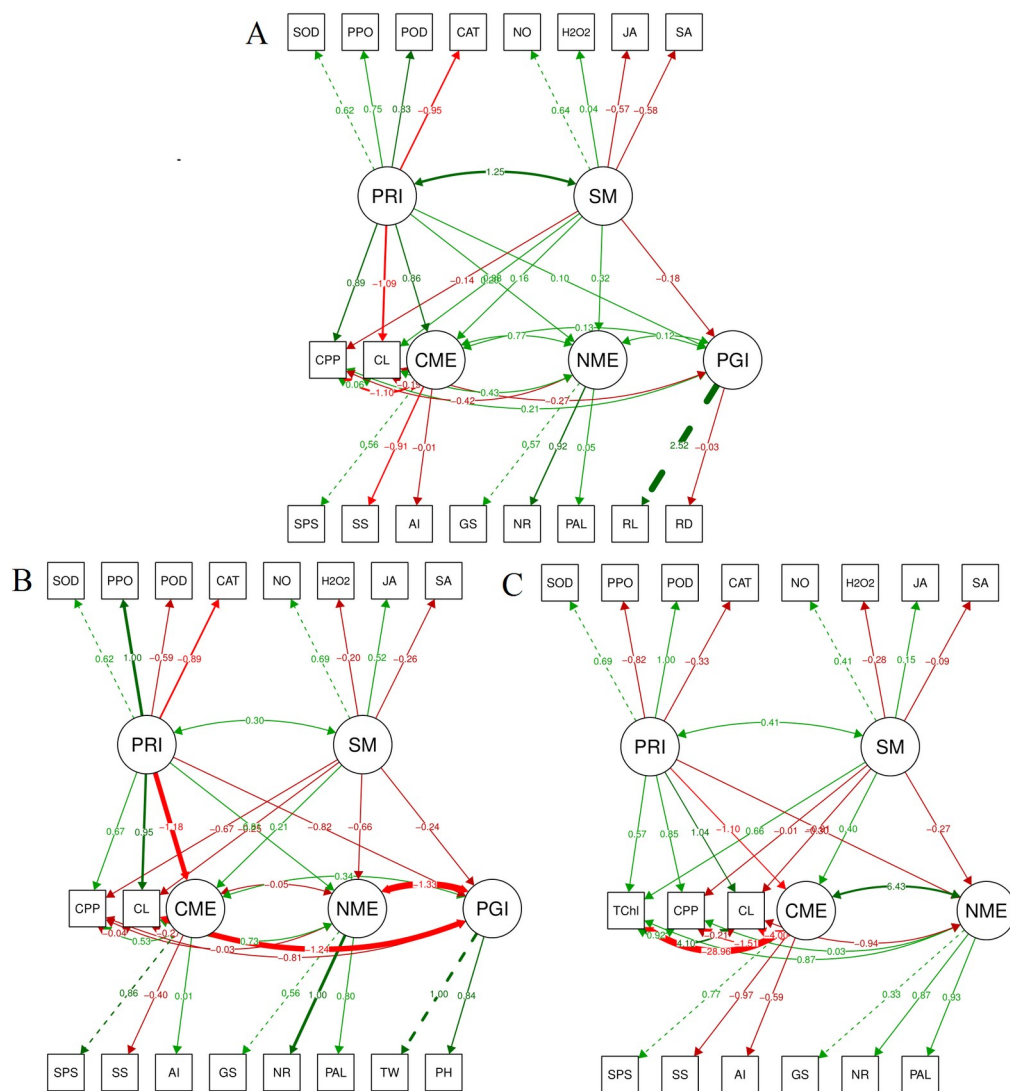


Fig. 5 The key enzymes involved in the synthesis of polysaccharide and Lobetyolin in different parts of *C. pilosula* in the control group and the treatment group. (A. The key enzymes activity involved in polysaccharide synthesis and content of polysaccharide (CCP) in different parts of *C. pilosula* in the control group and the treatment group; B. The key enzymes activity involved in Lobetyolin synthesis and content of Lobetyolin (CL) in different parts of *C. pilosula* in the control group and the treatment group.)



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73Fig. 6 The correlation analysis among various biological indicators. (A. After inoculation the endophytic
74*Trichoderma* strain RHTA01 in *C. pilosula*, the biological indicators of various parts of the host plant had
75changed; B. After inoculated with the strain, regulated the expression of related genes, participated in plant
76resistance and secondary metabolites accumulation; C. the biosynthesis pathway of *C. pilosula*
77polysaccharide; D. RDA analysis showed the correlation among the contents of plant index (TChl and LN),
78plant stress resistance index (MDA, Pro, Cpr, ReS, PPO, SOD, CAT, POD) and signaling molecules (NO,
79H₂O₂, JA, SA); E. RDA analysis showed the correlation among the contents of signaling molecules (NO,
80H₂O₂, JA, SA), polysaccharide synthesis (CPP) and Lobetyolin (CL); F. RDA analysis showed the
81correlation among the contents of sucrose metabolism (sucrose phosphate synthase, SPS; sucrose synthase,
82SS; soluble acid invertase, AI), compound synthesis (glutamine synthetase GS; phenylalanine ammonia
83lyase ,PAL; nitrate reductase, NR) and polysaccharide synthesis (CPP).)



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85Fig. 7 Structural equation model (SEM) showed the causal relationships among Plant resistance index
 86(PRI), Signal molecule (SM), Carbon metabolism enzyme (CME), Nitrogen metabolism enzyme (NME),
 87Content of polysaccharide and Plant growth_index (PGI) from different parts of *C. pilosula*. (A. SEM was
 88used for analysis the relationship of different indexes in the plant root; B. SEM was used for analysis the
 89relationship of different indexes in the plant stem; C. SEM was used for analysis the relationship of different
 90indexes in the plant leave.)