

Supplementary Materials: A Jacalin-Related Lectin Regulated the Formation of Aerial Mycelium and Fruiting Body in *Flammulina velutipes*

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Table S1. Primers used in this study.

Primer	Nucleotide Sequence (5'-3')
<i>gapdh</i> -F	CCTCTGCTCACTTGAAGGGT
<i>gapdh</i> -R	GCGTTGGAGATGACTTGAA
<i>Fv-JRL1</i> -F	GCCAGCCTCTCGGTAGC
<i>Fv-JRL1</i> -R	CGTAGACACGGTCGCCAG
<i>Fv-JRL1</i> -F-SpeI	<u>GGACTAGT</u> CCTAACTCTCATTACTCTCAGC
<i>Fv-JRL1</i> -R-ApaI	<u>GAGGGCCCC</u> CATGATTAGGCAGCAAATAG
<i>Fv-JRL1</i> -S-F	TCTACACACAACACTTATCGCTATCCAAAACAATCGG
<i>Fv-JRL1</i> -S-R	GGATTACAGACCCTCAACGACTGGTGTGACTATGCCATTAGG
<i>Fv-JRL1</i> -A-F	CATGCCAATTCTAGAGGGCCGCTATCCAAAACAACATCGG
<i>Fv-JRL1</i> -A-R	CCTAACGCCATACTCAACACCAGTCGTTGAAGGGTCTGTAATCC
Hpt-F	CTATTCTTGCCTCGG
Hpt-R	ATGAAAAAGCCTGAACTCACC
GBT-F	CCCAGGCTTACACTTTAT

Restriction sites are underlined.

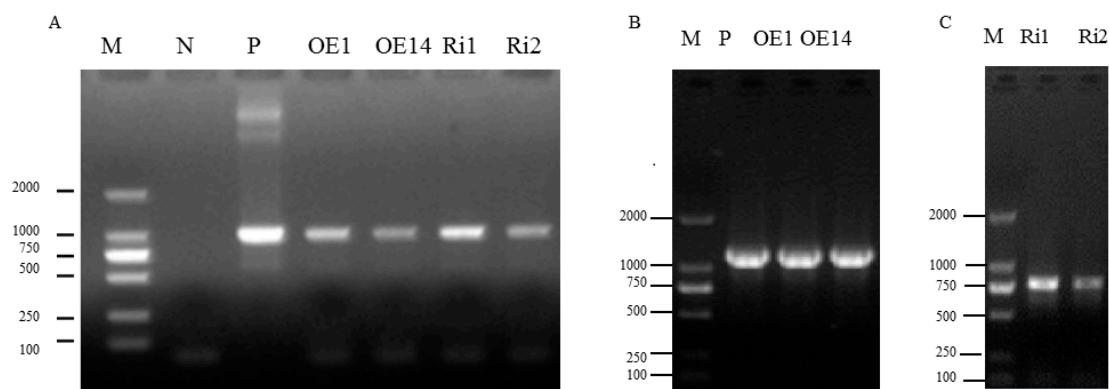


Figure S1. The identification of *Fv-JRL1* putative transformants using PCR. M: Marker DL2000; P: plasmid *pBHg-Fv-JRL1*-OE; N: wild-type strain H1123. (A) Presence of the hygromycin B resistance gene (*hpt*) in transformants. Primers Hpt-F and Hpt-R were adapted; (B) Presence of *Pgpd* and the full-length *Fv-JRL1* gene. PCR with specific primers of *Pgpd* and the full-length *Fv-JRL1* gene was carried out with GBT-F and *Fv-JRL1*-R; (C) Presence of *Pgpd* and a 376 bp fragment cloned in forward orientations in the silencing cassette. PCR with specific primers GBT-F and *Fv-JRL1*-S-R was performed with RNAi transformant genomic DNA.