Supplementary files

Double-Stranded RNAs (dsRNAs) as a Sustainable Tool against Gray Mold (Botrytis cinerea) in Grapevine: Effectiveness of Different Application Methods in an Open-Air Environment

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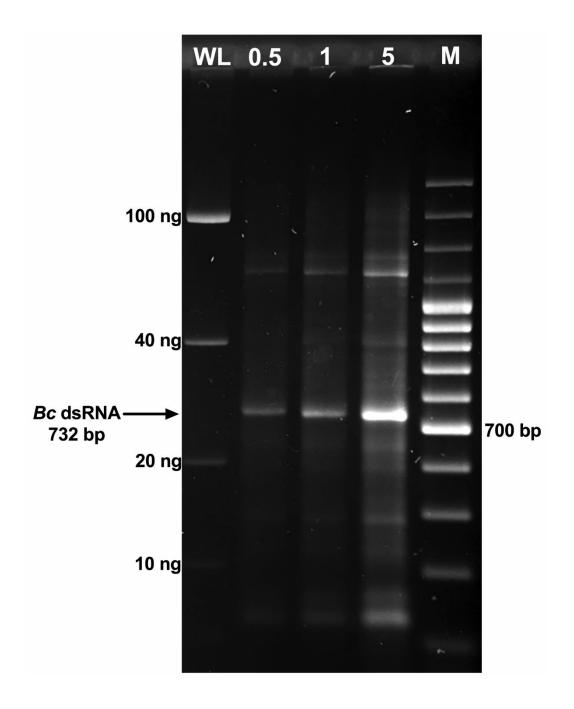
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Supplementary Table 1. List of primer designed to be used in overlapping PCR.

Target Gene	NCBI Accession	Primer name	Primer sequence
3-ketosteroid reductase (erg27)	JX436969.1	Bc_3Keto_F	AAAACTGCAGGATACGGTGGCTCGGGTTCA
		Bc_3Keto_R	TTGAGGGTGCGGCGATTGATCTCGCCTTCAGACCCTTCCT
Chitin syntase 1 (CHS1)	XM_024694912.1	Bc_CHS1_F	AGGAAGGGTCTGAAGGCGAGATCAATCGCCGCACCCTCAA
		Bc_CHS1_R	TGCCATACTCGACACCCTCCTGGGATTGAGGGGCATGC
Elongation factor 2 (ef2)	XM_024693463.1	Bc_EF2_F	GCATGCCCTCAATCCCAGGAGGGTGTCGAGTATGGCA
		Bc_EF2_R	AAAACTGCAGGCGGGGAGGAAGTCTGTGAC
Green fluorescent protein (GFP)	From pCBCT plasmid	GFP_F	AAAACTGCAGACCCTGAAGTTCATCTGCACCAC
		GFP_R	AAAACTGCAGGATCTTGAAGTTGACCTTGATGCC

Supplementary Figure 1. Agarose gel electrophoretic analysis of Bc dsRNA fragments after DNAse treatment. Samples were loaded as 0.5-1-5 μ L of total nucleic acid isolation as detailed in Materials and Methods section. WL: weighed ladder; M: size molecular marker.



Supplementary Figure 2. Overview of *in vivo* petiole adsorption application method.



Supplementary Figure 3. Northern blot analysis of Bc dsRNA in Leaf and Berry tissues over time. '-' indicate untreated samples; '+' indicate Bc dsRNA treated samples. ds+plant: Bc dsRNA added to an untreated leaf sample used as positive control.

