Table S1. Sequence analysis of cloned six Avr genes in four Leptosphaeria maculans isolates.

	L. maculans isolates			
	03-42-06	87-41	03-15-03	PG4-1-M
AvrLm1 ^a				
Genotype	Α	Α	а	а
PCR	+	+	-	-
AvrLm2 ^b				
Genotype	а	Α	Α	а
PCR	+	+	+	+
Nucleotides for 133rd AA	CAT	GGT	GGT	AAT
Nucleotides for 146th AA	CAA	GAA	GAA	CAA
AvrLm4-7 ^c				
Genotype	Α	Α	а	а
PCR	+	+	-	-
Nucleotides for 80th AA	ATA	ATA		
Nucleotides for 120th AA	GGC	GGC		
AvrLm6 ^d				
Genotype	Α	Α	Α	Α
PCR	+	+	+	+
AvrLm11 ^e				
Genotype	Α	Α	Α	Α
PCR	+	+	+	+
AvrLm5 ^f				
Genotype	Α	Α	Α	Α
PCR	+	+	+	+
Nucleotides for 29th AA	CGA	CGA	CGA	CGA
Nucleotides for 38th AA	CGG	CGG	CGG	CGG
Nucleotides for 55th AA	AGA	AGA	AGA	AAA

- a, d and e. Virulent allele of AvrLm1, AvrLm6 or AvrLm11 was caused by deletion of the gene; PCR was used to detect the presence/absence of Avr genes in four isolates.
- **b.** Polymorphic sites (133rd and 146th amino acids) determining the avirulence/virulence toward *Rlm2* in *AvrLm2*. The nucleotides GGT coding Gly in 133rd and GAA coding Glu in 146th render the *Avrlm2* complementary and avirulent to *Rlm2* (Ghanbarnia *et al.*, 2015).
- **c.** Polymorphic sites (120th amino acid) determining the avirulence/virulence toward *Rlm4* in *AvrLm4-7*. The nucleotides GGC coding Gly for the 120th render the *AvrLm4-7* complementary and avirulent to *Rlm4* (Blondeau *et al.*, 2015).
- **f**. Polymorphic sites (29th amino acid) in *AvrLm5* determining the avirulence/virulence toward *B. napus* cultivars. The nucleotide changes from CGA coding Arg for the 29th amino acid to TGA coding the premature stop code result in the virulence of *AvrLm5* toward *B. napus* cultivars (Van de Wouw *et al.*, 2014). The polymorphic sites for 38th and 55th amino acids were also detected.

For genotype indication, 'A' means the presence of the functional *Avr* gene while 'a' means presence of the non-functional *Avr* gene or the absence of the *Avr* gene. For PCR screening, '+' means the detection of the *Avr* gene by PCR, '-' means the absence of the *Avr* gene in the PCR screening.