

Supplemental Table 5. Primer combinations used to genotype mutations by PCR.

Mutant	Marker	Enzyme	Forward and Reverse Primers (5' to 3')
<i>sid2-1</i>	Caps	MfeI	F: TGTCTGCAGTGAAGCTTTGG R: CACAAACAGCTGGAGTTGGA
<i>jar1-1</i>	Caps	HindIII	F: CAGTGTGTGTGTTTTTGATCATAAGCT R: CAAATTTAAACTATACCTGTTTCTGAAGG
<i>eds1-1</i>	Dcaps	HindIII	F: CGAAGGGGACATAGATTGGA R: CTTTTCATGTACGGCCCTGT
<i>coi1-1</i>	Dcaps	XcmI	F: GTTTCTCTTTAGTCTTTAC R: CAGACAACATTTTCGTTACC
<i>npr1-1</i>	Caps	NlaIII	F: AGGCACTTGACTCGGATGAT R: ATGCACTTGCACCTTTTTCC
<i>pad4-1</i>	Caps	BsmfI	F: GCGATGCATCAGAAGAG R: TTAGCCCAAAGCAAGTATC
<i>NahG</i>	Presence/absence		F: CTGCCGCTACTCCCATATCCA R: TCGGCTTCGGCTCGCTAC
<i>Etr1-1</i>	Dcaps	ApaLI	F: AAGTTAATAAGATGAGTTGGTGCA R: AGAAATCAGCCGTGTTTCCG
<i>sgt1b-1</i>	Caps	AvrII	F: GGAGAGAGACAAATTCGAGCAG R: GTACCTAAGGAGAACGTAACGTGTC
<i>rar1-10</i>	5 bp deletion		F: CCAGTACAAAAGGCTGTGAT R: ACAGTGAAAGAAAAGGGTCA
<i>pen1-1</i>	Dcaps	MluI	F: GAAACACTCTTTCATGTCACGCG R: GAGGACAGAGGTCCTGGTTTCG
<i>pen2-3</i>	Caps	BsmAI	F: TGGCAAAAGGGATAGAGGAG R: CTGTAGACCCACGGCTCATT
<i>pen3-1</i>	Caps	HphI	F: TGAAAGCTTCTGCTGCTCAA R: TGAGGTGAACGATTTGTTGC