

Table 1 Suppl. Specific and degenerate primer sequences used for cloning fragments of *soluble acid invertase (SAI)*, *cell wall acid invertase (CWAI)*, *neutral invertase (NI)*, *sucrose synthase (SS)*, and *sucrose phosphate synthase (SPS)* genes.

Gene	Forward primers (5'-3')	Reverse primers (5'-3')
<i>SAI</i>	GTGGGGTGACATAGTTTGGGGA	AGGCTAGCCTTAACCACATG
<i>CWAI</i>	GAGATGGAGAATGGCTGTTGGTAGC	GGATTGGTCCAACCTGGGAC
<i>NI</i>	TCCTGCTTGGGTGGTTGANTKBATNCC	CTAGCAATCTCAGGCCTACCCRTYTTNAYRCA
<i>SS</i>	GTTCCCTTAGA(A/G)C(A/T)C(A/T)GATGG	TC(A/C)AG(T/C)CTTGCCAT(T/G)G(T/A)AAA
<i>SPS</i>	GGTGGTCAGGTAATGTTT	CGTTCAGTATTGGGTCA

Table 2 Suppl. Primer sequences for real-time quantitative PCR.

Gene	Forward primers (5'-3')	Reverse primers (5'-3')	Product [bp]
<i>DISAI</i>	TCCAATACAGACAACAACACTACGC	ACCCACATAAGGCAACCAAG	100
<i>DICWAI</i>	GCACAGGCAGACATTGAGAT	TTCCACTGAAGCACCCCTTATG	117
<i>DINI-1</i>	CGTTTCCTGTGAATGGTCTG	AATAAGTCCCACGCTTCGTC	139
<i>DINI-2</i>	CAATCACGACTCTTCCAGACA	CCTCCTCCCAGAATAACAAGG	100
<i>DINI-3</i>	ATGTTGCGGCTGAGAGAGAG	GGCACAAAGTCACGAATGAA	169
<i>DISS-1</i>	TTCACTCTACCTGGACTCTA	GTGTTCCTTATTCTCAACC	186
<i>DISS-2</i>	ATGACTTGTGGCCTTCCAAC	TCTGGTCAGGGTGATATGGA	106
<i>DISS-3</i>	CAGGGACAGAGAAGAAGTTGC	GTTGGAAGCCCACAAGTCAT	222
<i>DISPS</i>	GTTTTGGCATCCCGTTCCC	GTCCTTCATAATCTGTGTCCCC	112
<i>DActin</i>	TGCTATCCTTCGGTTGGACC	CGGACGATTTCCTGTTTCAG	93
<i>DIEF1α</i>	GATGATTCCCACCAAGCCCAT	GGGTCCTTCTTCTCAACACTCT	129