

**Supplementary Table S1.** Alignment and processing statistics for sequencing read data against the tomato genome reference sequence version 2.5 (ITAG2.4).

	MoneyMaker <sup>a</sup>	<i>sin</i> wild-type pool <sup>b</sup>	<i>sin</i> mutant pool <sup>c</sup>
Total sequencing reads	63,575,958	94,007,596	99,685,109
Uniquely mapped reads	56,571,716 (88.98 %)	88,279,910 (93.91 %)	92,975,595 (93.27 %)
Multi-mapped reads	1,032,320 (1.62 %)	1,355,151 (1.44 %)	1,496,688 (1.50 %)
Unmapped reads	5,971,922 (9.39 %)	4,372,535 (4.65 %)	5,212,826 (5.23 %)
Sequencing depth	15.40 X	34.02 X	36.04 X
Sequence variants	813,450	6,100,013	6,707,616
<sup>a</sup> NCBI SRA BioSample accession: SAMN04093174.			
<sup>b</sup> NCBI SRA BioSample accession: SAMN30553549.			
<sup>c</sup> NCBI SRA BioSample accession: SAMN30553548.			