Table S2  Oligonucleotide primers used in PCR reactions

<table>
<thead>
<tr>
<th>Gene</th>
<th>Primer</th>
<th>Nucleotide sequence</th>
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<tr>
<td>vic2</td>
<td>ptnF1</td>
<td>5'-TGCAGCACCTGGATGTACATA</td>
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<td>ptnR1</td>
<td>5'-CGTCATACAGGCGAAGGGAT</td>
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<td>5'-TACTCTCTCCAAACGCTCCCG</td>
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<td>5'-GCTCAACGTATGTGATGCTAGCAT</td>
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<td>5'-CTTGATCGTGGAGTTGCTAGCT</td>
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<td>5'-GACCAGCTCTTTGGGCAGCT</td>
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<td>5'-CGAGACCCTTTTTGTTTCTAGGCT</td>
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<td>vic7R1</td>
<td>5'-ATAGGGCTTTCGGAGATCGA</td>
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</table>

*A Phire Plant Direct PCR kit (F-130) (New England BioLabs, Ipswich, MA) was employed in a 50 μl reaction volume with following parameters: denaturation at 98°C for 2 min followed by 30 cycles consisting of denaturation at 98°C for 5 sec, annealing at 64°C for 5 sec, extension at 72°C for 2.5 min, and then final extension at 72°C for additional one minute. The resulting PCR products were sequenced after purification with a QiAquick PCR purification kit (Qiagen, Valencia, CA).*