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volume, including 12.5 μ l of PCR mixture (Invitrogen), 0.4 μ l of each primer (forward and reverse), 1 μ l of cDNA template, and 9.5 μ l of ddH₂O. The amplification conditions consisted of an initial denaturing step at 94 °C for 5 min, followed by 35 cycles of amplification at 94 °C for 1

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