

Table S1 Sequences of the primers used in the study.

Gene	Forward Primer (5'-3')	Reverse Primer (5'-3')	Purpose
<i>AtIND</i> promoter (230 bp)	GTTTAAACTGGGCAATACCTT AATGTCG	CCTAGGCTAAACCCACGGTTT CTTCG	Cloning
<i>AtIND</i> promoter (400 bp)	GTTTAAACTGGGCAATACCTT AATGTCG	CCTAGGGCCATAAGGAAGCAT AAGTCG	Cloning
<i>LcIND</i> gene fragment	GGGGACAAGTTTGTACAAAA AAGCAGGCTGATGAAGACAT GGATGCCATGA	GGGGACCACTTTGTACAAGAA AGCTGGGTTCCAAGGTGAGAGT GAGGCTGAA	Cloning
<i>nptII</i>	GCCCTGAATGAACTGCAGGAC GAGGC	GCAGGCATCGCCATGGGTCACG ACGA	PCR
<i>LcTIP41</i> - like	GCTTATGAGATTGAGAGAGAC GAGAA	GGATACCCTTTCGCAGATAGAG AC	qRT-PCR
<i>LcIND</i>	TAAGCGACGATCCTCAGACG	AAACATTGCGGTGGGATTGC	qRT-PCR