Table S1. Primer sequences for qRT-PCR analysis of target genes of miRNAs

Sequence name	Forword primer (5′- 3′)	Reverse primer (5′- 3′)
Tp-miRNA156	GCGTCGGACCAGGCTTCA	GTCGTATCCAGTGCAGGGTCCGAGGTATTCGCACTGGATACGAC GAGGAA
Tp-miRNA164	GTGTTGGCCCGGCTCA	GTCGTATCCAGTGCAGGGTCCGAGGTATTCGCACTGGATACGAC TCTGAG
Tp-miRNA4995	GCGTTCCACGGCTTTCTTG	GTCGTATCCAGTGCAGGGTCCGAGGTATTCGCACTGGATACGAC GCAGTT
Tp-miRNA159	CGCGTTTGGATTGAAGGGAG	GTCGTATCCAGTGCAGGGTCCGAGGTATTCGCACTGGATACGAC AAAGAG
PC-5p-84014	CGGTTCCCTCCGGCAC	GTCGTATCCAGTGCAGGGTCCGAGGTATTCGCACTGGATACGAC GGTGAA
PC-3p-212-24745	CGCGTTTTCGGGTGATTTG	GTCGTATCCAGTGCAGGGTCCGAGGTATTCGCACTGGATACGAC CCACCT
Tp-GA20ox	ACTCCTGGGAAATGACGAGG	AAGGAGAACGGTGAGGGATG
Tp-NAC	CTTGATGCGGGAATTCAGTT	ATTGGCTTTGTATGCGGTTC
Tp-CYP707A1	AGCTAGCGTTCTCACCTGGA	TCCACATCTTGCACTGCTTC
Tp-DELLA	CGAGTCCAGCAGCTAATGTG	GCTTCAAGAATGGCTCCGTT
Tp-PYL	GGGTGGGTCTGTGTGTTCGA	CCCTCTCAGTAATCGCCGCA
Tp-ARF	TCGTGTGATCAATGTCCAGC	GAACGAGTGAACATGTGGCC

Table S2. cDNA synthesis.

Component	Dosage
4×gDNA wiper Mix	4 μL
Templet RNA	1.2 μL
Rnase free ddH_2O	To 16 μL
First PCR: 42 ℃ 2 min	
Last step reaction liquid 16 μL + 5×Hicript qRT super Mix	4 μL
Second PCR: 50 ℃ 15 min	
85 ℃ 5 sec	

Table S3. Reaction system and procedure of qRT-PCR.

PCR reaction system

Reagent	Volume	Concentration
MonAmp™ SYBRGreen qPCR Mix	10	1×
Forward primer	1	0.2 μM
Reverse primer	1	0.2 μM
Template DNA	1	200 ng/ 20 μL
ddH_2O	7	

PCR procedure

Step	Temperature	Time	
Predegeneration	95	30 sec	
Denaturation	95	5 sec	40 cycle
Annealing and extension	55-68	30 sec	