

Table S1: Primers used for qRT-PCR analysis.

Primers	Sequences (5' to 3')
ZmLC-RT F	ATAGCCTACCTCAAGGAGCTTCAGA
ZmLC-RT R	AGACCTCCTCCTCACACTCTCATT
ArActin_RT F	ACCTCAAAATAGCATGGGAAGT
ArActin_RT R	GGCCGTTCTCTCACTTATGCTA
ArPAL_RT F	ACGGCTCCAACGGTCATAATAAT
ArPAL_RT R	ATCCGCTTACCTCCTCAAGGT
ArC4H_RT F	GTTCGAGAGTGAGAATGATCCGT
ArC4H_RT R	ATAATCCTGAACAATTGCAGCC
Ar4CL_RT F	ACATCTACTCGTTGAATTGGTGC
Ar4CL_RT R	AGTCGAAATTATCCACCAATGGA
ArCHS_RT F	GACCAAAGCACCTATCCGATTA
ArCHS_RT R	TTGGGTTCTCCTTCAGGTACTCC
ArCHI_RT F	GCCTTCTCAAAGATGGTCTGT
ArCHI_RT R	TCTTGATTCAAGTTGCCTCAGC
ArRAS_RT F	GGCGAACTATCATACCCTGAG
ArRAS_RT R	AATCAATTCCAGGCAGTTGCCG
ArTAT_RT F	AGGCTGCAGTCCTGAAATCATT
ArTAT_RT R	TTCACCATGAAAGCCATTGCTCC
ArHPPR_RT F	AAGGGGATTAGGGTTACCAACACA
ArHPPR_RT R	ATTCTACCCAATCCAATGATGCC

Table S2: HPLC conditions for phenylpropanoid compounds detected in this study.

Compounds	Extraction	Operating system and conditions	Program
Phenylpropanoid compounds	A 0.1 g freeze-dried sample was sonicated for 1 h in 2 mL of 80% methanol (v/v). After centrifugation at 3500 rpm for 5 min, the supernatant was filtered through a 0.45 µm PTFE syringe filter (Toyo Roshi Kaisha, Ltd., Tokyo, Japan).	The system comprised an OptimaPak C18 column (250 × 4.6 mm, 5 µm; RStech Co., Daejeon, Republic of Korea), an NS-4000 HPLC system, an NS-6000 auto-sampler (Futecs Co., Daejeon, Republic of Korea), a degasser, and a UV-Vis detector.	The gradient program was set as follows: solvent A, ultrapure water containing 0.2% acetic acid; solvent B, methanol; 0 min, 95% A; 4 min, 95–85% A; 9 min, 85% A; 14 min, 85–80% A; 24 min, 80% A; 54 min, 80–70% A; 55 min, 70–55% A; 65 min, 55% A; 75 min, 55–44% A; 77.0 min, 44–40% A; 79 min, 40% A; 80 min, 40–20% A; 90 min, 20% A; 91.0 min, 20–95% A; and 98.0 min, 95% A (Total 98 min).