

Figure S1. Construction of apple calli cDNA library, screening of yeast one-hybrid cDNA library and PCR detection of positive colonies. (A) Agarose gel electrophoresis analysis of total RNA and mRNA extracted from apple calli. Total RNA electrophoresis map, marker, mRNA electrophoresis map. (B) Identification of cDNA primary library capacity and agarose gel electrophoresis analysis of inserted fragment sizes. (C) Identification of cDNA secondary library capacity and agarose gel electrophoresis analysis of inserted fragment size. (D) The screening of yeast one-hybrid cDNA library. 1/10, 1/100, and 1/1000 respectively represent 10 times, 100 times, and 1000 times diluted conversion. (E) Yeast colony PCR detection of yeast from figure D. MdSCL8 is marked with red rectangle.

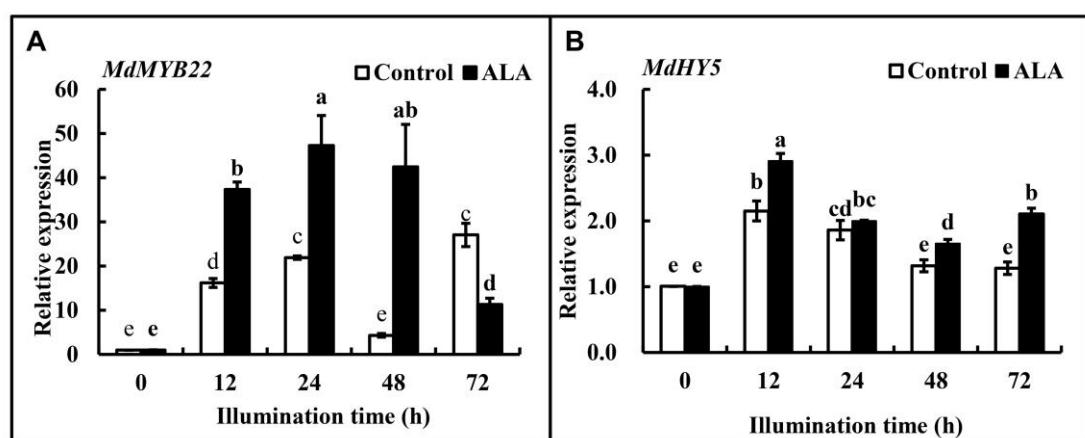


Figure S2. Effect of ALA on *MdMYB22* and *MdHY5* gene expression in calli. The calli were treated with 50 mg L⁻¹ ALA or deionized water (Control) for 3 h under dark and then cultured in solid MS medium under light of 100 $\mu\text{mol m}^{-2}\text{s}^{-1}$ photon flux density at 22°C for 72 h. The values of (A, B) were performed 3 biological replicates. Error bars represent standard errors. Three biological replicates were performed for each experiment. The same lowercase letters indicate no significant differences at $P \leq 0.05$.

Table S1. Primer sequences used in the study

	Primer name	Froward primer5'-3'	Reverse primer5'-3'
121-SCL8		GAGAACACGGGGACTCTAGAATG	GGACTGACCACCCGGGATCCACG
		CAATCGGGAGGCTCAACG	CCAAGCGGAAGCGAC
1300-SCL8		GAGAACACGGGGACTCTAGAATG	GCCCTTGCTCACCATGGATCCACGC
		CAATCGGGAGGCTC	CAAGCGGAAGCGAC
0800-MdFLS1		TCGACGGTATCGATAAGCTTAGGT	GCTCTAGAACTAGTGGATCCTCCA
		GTGAAATTCTGTCTTTAGGACA	CCGCTCTCATCTCTCTT
CDS amplification	<i>proMdFLS1</i>	GAGAACACGGGGACTCTAGATAG	GGACTGACCACCCGGGATCCTCC
		GTGTGAAATTCTGTCTTTAGG	ACCGCTCTCATCTCTC
<i>MdSCL8(i)</i>		GGGGACAAGTTGTACAAAAAAGC	GGGGACCACTTGTACAAGAAAGCT
		AGGCTCCCACCAGCAGAACCACTAC	GGGTGCTTGCAGCGTGTACTGA
<i>MdHY5(i)</i>		GGGGACAAGTTGTACAAAAAAGC	GGGGACCACTTGTACAAGAAAGCT
		AGGCTTCTCGAGCTCTGCATTCCA	GGGTAACAACCTCTTCAGCCGCTT
<i>MdMYB22(i)</i>		GGGGACAAGTTGTACAAAAAAGC	GGGGACCACTTGTACAAGAAAGCT
		AGGCTTGAAATTAGCACCAGTGTCC	GGGTGGGACCTCCATTAGTAGCAGC
qRT-PCR	<i>MdUBQ</i>	CTCCGTGGTGGTTTAAGT	GGAGGCAGAAAACAGTACCAT
	<i>MdFLS1</i>	AAAATGGGAGTGGAGTCTGTG	TTATGACCTCGCTGGGAATGT
	<i>MdSCL8</i>	CGTGGAAAGATGTGAAGTGT	GCGGCTTATCCAACCAAA
	<i>MdHY5</i>	GGAAGAGTGCCGGAATCG	CTCCCTGCTGCTGTGCT
	<i>MdMYB22</i>	GTAATAGGTGGTCTTGATAGCG	GGGTTGGTGGTAGAGTAGGT
Y2H/Y1H	<i>BD-MdHY5</i>	ATGCCATGGAGGCCAATTATGC	CCGCTGCAGGTGACGGATCCATCC
		AAGAGCAGGCCACG	GCATTGACCCACCA
	<i>AD-MdMYB22</i>	GCCATGGAGGCCAGTGAATTATGC	CAGCTCGAGCTCGATGGATCCCTGG
		GGAGGGCGCCGTGC	ATGCCGTAGTAAGTCGTCG
	<i>AD-MdHY5</i>	GCCATGGAGGCCAGTGAATTATGC	CAGCTCGAGCTCGATGGATCCATCC
		AAGAGCAGGCCACG	GCATTGACCCACCA
	<i>BD-MdSCL8</i>	ATGCCATGGAGGCCAATTATGC	CCGCTGCAGGTGACGGATCCACGC
		AATCGGGAGGCTTCA	CAAGCGGAAGCGAC
	<i>AD-MdSCL8</i>	GTGGGCATCGATACGGATCCATGC	ACGATTCATCTGCAGCTCGAGACGC
		AATCGGGAGGCTTCA	CAAGCGGAAGCGAC
<i>proMdFLS-pAbAi</i>		AAATGATGAATTGAAAGCTTAGG	ATACAGAGCACATGCCCTCGAGTTCC
		TGTGAAATTCTGTCTTTAGGACA	ACCGCTCTCATCTCTCTT
	<i>YN-E-MdHY5</i>	GAGAACACGGGGACTCTAGAATG	CTCCATCCGGGAGCGGTACCATCC
		CAAGAGCAGGCCACG	GCATTGACCCACCA
BIFC	<i>YN-E-MdSCL8</i>	GAGAACACGGGGACTCTAGAATG	CTCCATCCGGGAGCGGTACACGC
		CAATCGGGAGGCTTCA	CAAGCGGAAGCGAC

	<i>YCE-MdMYB22</i>	GAGAACACGGGGACTCTAGAATG GGGAGGGCGCCGTGC	GTACATCCGGGAGCGGTACCTGG ATGCCGTAGTAAGTCGTCG
	<i>YCE-MdSCL8</i>	GAGAACACGGGGACTCTAGAATG CAATCGGGAGGCTTCA	GTACATCCGGGAGCGGTACCACGC CAAGCGGAAGCGAC
Luciferase	<i>NLUC-MdHY5</i>	GAGCTCGGTACCCGGGATCCATGC AAGAGCAGGCAGC	CGCGTAGAGATCTGGTCGACATCC GCATTTCGACCA
	<i>CLUC-MdMYB22</i>	TACCGTCCCAGGGCGGTACCATGG GGAGGGCGCCGTGC	TGTAGTCCATTGTTGGATCCCTGG ATGCCGTAGTAAGTCGTCG
	<i>NLUC-MdSCL8</i>	GAGCTCGGTACCCGGGATCCATGC AATCGGGAGGCTTCA	CGCGTAGAGATCTGGTCGACACGC CAAGCGGAAGCGAC
	<i>CLUC-MdSCL8</i>	TACCGTCCCAGGGCGGTACCATGC AATCGGGAGGCTTCA	TGTAGTCCATTGTTGGATCCGCC AAGCGGAAGCGAC