

Supplementary Material

Ectopic expression of a R2R3 MYB transcription factor of dove tree (*Davida involucrata*) aggravates seed abortion in *Arabidopsis thaliana*

Jian Li^{A,B}, Tian Chen^A, Fengzhen Huang^{A,B}, Penghui Dai^{A,B}, Fuxiang Cao^{B,C} and Meng Li^{A,B,C,D}

^ACollege of Life Science and Technology, Central South University of Forestry and Technology, Changsha 410004, China.

^BHunan Research Center of Engineering Technology for Utilisation of Environmental and Resources Plant, Changsha 410004, China.

^CCollege of Horticulture and Landscape, Hunan Agricultural University, Changsha 410004, China.

^DCorresponding author. Email: limeng0422@foxmail.com

Fig. S1. PCR amplification of *DiMYB1* fragment.

Fig. S2. Relative expression levels of *DiMYB1* in leaves of transgenic and wild type plants.

Table S1. Information of primers applied in the present study

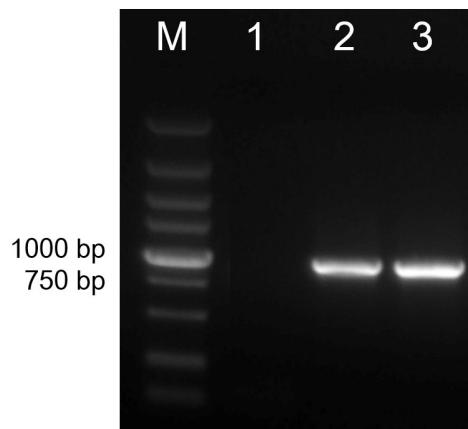


Fig. S1. PCR amplification of *DiMYB1* fragment. Lane M, DNA marker **DL5000**; Lane 1, negative control; Lane 2, PCR product using cDNA of abortive seed as template; Lane 3, PCR product using gDNA of *Davida* as template.

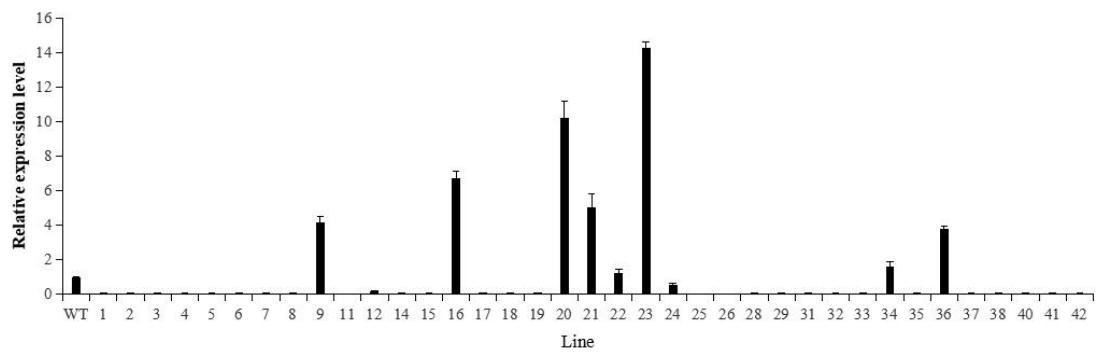


Fig. S2. Relative expression levels of *DiMYB1* in leaves of transgenic and wild type plants.
WT, wild-type Arabidopsis; 1-42, transgenic lines.

Table S1. Information of primers applied in the present study

Primer	Primer sequence (5'-3')	Product length	Function
DiMYB1-F	CGCGGATCCATGGGAAGGA AACCATGCTGTGCC		PCR amplification of full length fragment of <i>DiMYB1</i>
DiMYB1-R	CGGAATTCCGTCAATCAATC CATTCATCCCCTGAATTAA	924 bp	
Q-DiMYB-F	GGTTGGTGGAGATTACAGTA	90 bp	qPCR analysis of <i>DiMYB1</i>
Q-DiMYB-R	GCATCGTCGGTTGAATTG		
AtActin7-F	GGTCGTACAACCTGGTAT		qPCR analysis of <i>AtActin7</i>
AtActin7-R	TAGATACTTCCGATACGAG	76 bp	(reference gene for Arabidopsis)
DiCAC-F	GGTGGATGCCTTCCGAATAA		qPCR analysis of <i>DiCAC</i>
DiCAC-R	CTAACCTCCAACGAGCAAGA	83 bp	(reference gene for <i>Davidaia</i>)