

Supplement

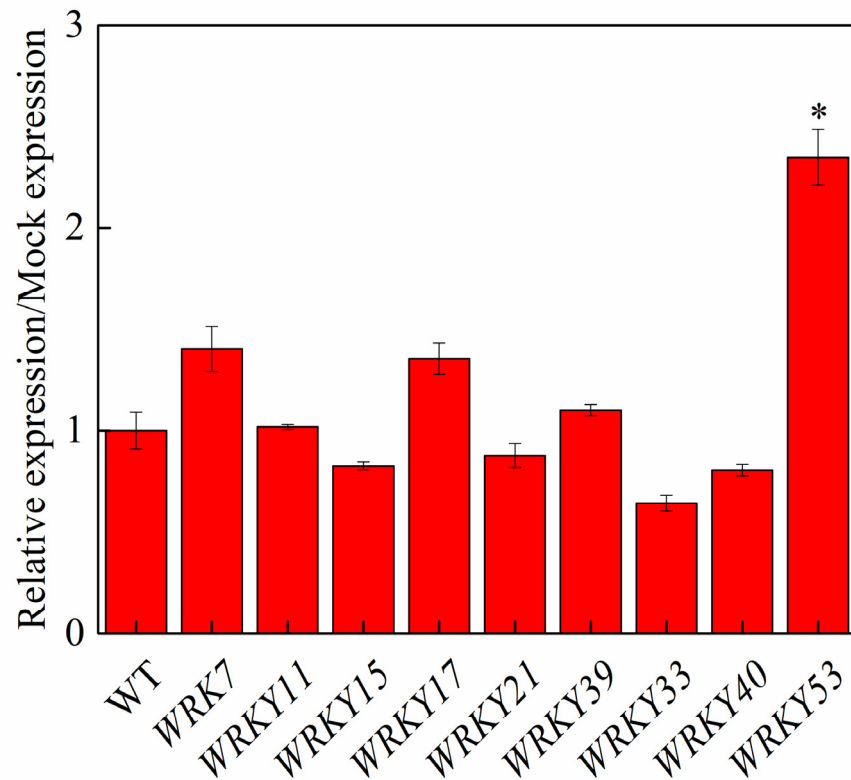


Figure S1. Selection of *WRKY53* based on the results of gene expression analysis in *cam1* mutant compared with WT. Each group had three biological replicates and every replicate had more than 30 plants. Plants were grown for 2 weeks. Error bars denote \pm SEM, * is significantly increased at $p < 0.05$, Dunnett's C (variance not neat).

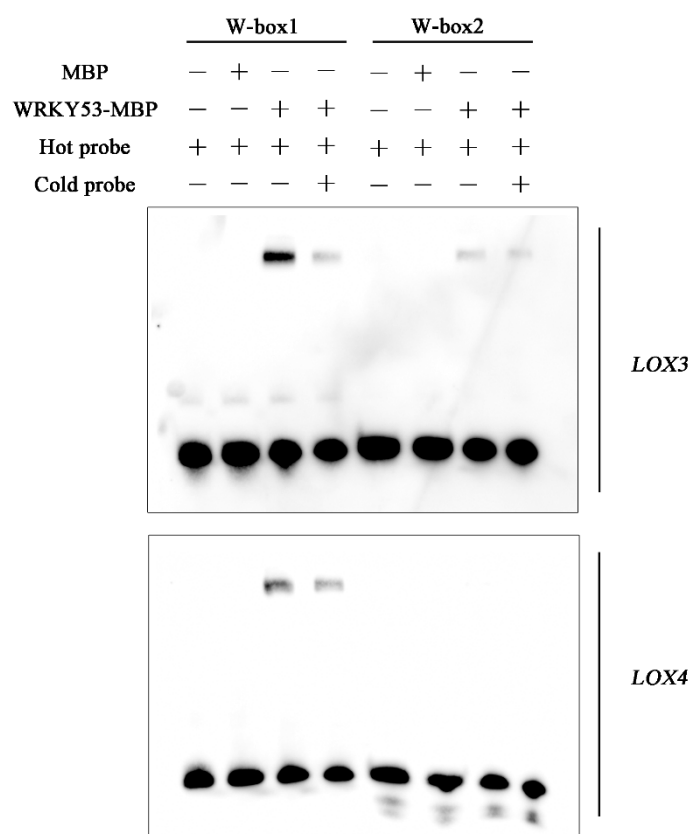


Figure S2. EMSA shows that WRKY53 binds to the W-boxes in the *LOX3* and *LOX4* promoters. every promoter has two W-boxes, W-box 1 and W-box 2.

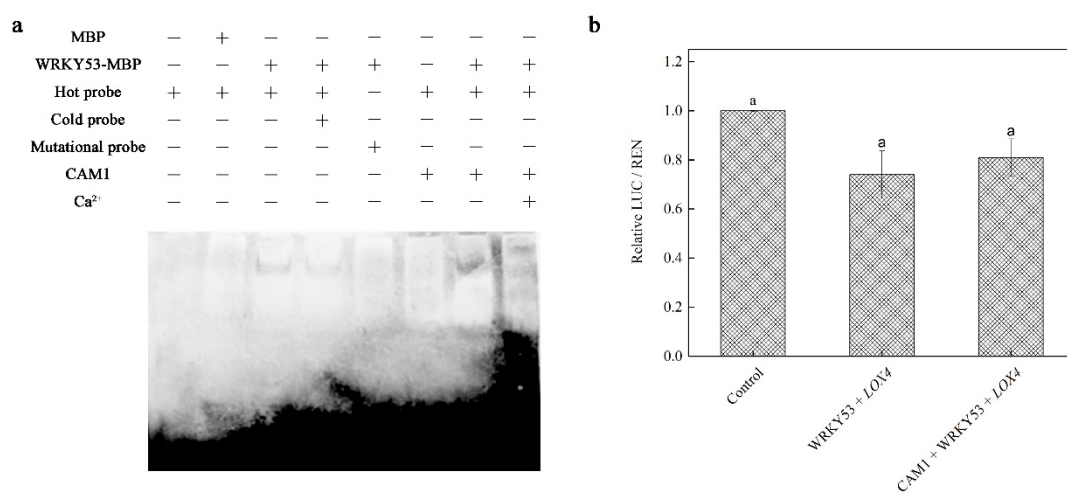


Figure S3. Interaction between WRKY53 and the *LOX4* promoter. (a) EMSA assay

shows that WRKY53 has a weak interaction with the *LOX4* promoter. And it shows the effects of calcium and CAM1 on the binding between WRKY53 and the first W-box in the *LOX4* promoter. A hot probe refers to a biotin-labeled probe and a cold probe to an unlabeled probe (200-fold the concentration of hot probe). Ca^{2+} concentration is 10^{-2} mM. (b) Luciferase activity assay shows that WRKY53 interacts with the *LOX4* promoter, WRKY53 negatively affects *LOX4* expression, and CAM1 does not significantly decrease the negative regulation. Error bars denote \pm SEM, columns labeled with different letters are significantly different at $p < 0.05$, Dunnett's C (variance not neat).

Table S1. The probe sequences in the EMSA.

W-boxes	Probe sequence (5'-3')
<i>LOX3</i> W-box1-F	CACACTTATATTTTGACCCTATGTTACT
<i>LOX3</i> W-box1-R	AGTAACATAGGGTCAAAATATAAGTGTG
<i>LOX3</i> W-box2-F	CGTAACTCCGTTTGACTCAATTACCCCA
<i>LOX3</i> W-box2-R	TGGGGTAATTGAGTCAAACGGAGTTACG
<i>LOX4</i> W-box1-F	TTTGTATATGTTTGACTTTGTGGCTATT
<i>LOX4</i> W-box1-R	AATAGCCACAAAGTCAAACATATACAAA
<i>LOX4</i> W-box2-F	CGGAACTCAGTTTGACTCAAAC TAGAAG
<i>LOX4</i> W-box2-R	CTTCTAGTTTGAGTCAAAC TGAGTTCCG