

Table S1. List of primers used in this study

Primer	Experiment	Sequence (5' -3')
PmCBFF1	Homology-based cloning of <i>PmCBF</i> genes	GAGACGAGGCACCCGGT
PmCBFR2	Homology-based cloning of <i>PmCBF</i> genes	AAACAHYTCCTCCTCATCC
5RaceF	Extension of 5' end of <i>PmCBFb</i>	AAGCAGTGGTATCAACGCAGAGTGG
3RaceR	Extension of 3' end of <i>PmCBFb</i>	ATTCTAGAGGCCGAGGCGGCCGACATG-d(T)30N-1N
PmCBFbrF1	Cloning of 3' end of cDNA <i>PmCBFb</i>	GACGTGGCGGCATTGGCTTTTAAAGG
PmCBFbrR1	Cloning of 5' end of cDNA <i>PmCBFb</i>	AGCATTGCGGTGGAGAAAGATGAAGC
PmCBFbZF	Cloning of full-length CDS of <i>PmCBFb</i>	CACACCTCTCTGAAACACCACTC
PmCBFbZR	Cloning of full-length CDS of <i>PmCBFb</i>	CACTAAACCCGAGAGACAGATTGACT
PmCBFcTF1	FPNI-PCR of 3' end of <i>PmCBFc</i>	CGGCATTGGCGTTTAGAGG
PmCBFcTF2	FPNI-PCR of 3' end of <i>PmCBFc</i>	GCGGAGTTTGGTGGATTGTGC
PmCBFcTF3	FPNI-PCR of 3' end of <i>PmCBFc</i>	GCAGTGATGAGAAGGAGAGAATGG
PmCBFcTR1	FPNI-PCR of 5' end of <i>PmCBFc</i>	CACCATTCTCTCCTTCTCATCACT
PmCBFcTR2	FPNI-PCR of 5' end of <i>PmCBFc</i>	TCCCTCTAAACGCCAATGCCG
PmCBFcTR3	FPNI-PCR of 5' end of <i>PmCBFc</i>	CTTGTTGGGCTCTCTCATTTAC
PmCBFcZF	Cloning of full-length CDS of <i>PmCBFc</i>	CAGCTCAAGAAACTCAAACAAGCTA
PmCBFcZR	Cloning of full-length CDS of <i>PmCBFc</i>	TGGTCGTCGAGTGACGTTAATCT
PmCBFaF1	Cloning of tandem array of <i>PmCBFab</i>	GACGTGGCGGCATTGGCTTTCAGAGG
PmCBFbR1	Cloning of tandem array of <i>PmCBFab</i>	TCATGGGCACGAGCAGCCAT
PmCBFcF1	Cloning of tandem array of <i>PmCBFca</i>	GAGAGAATGGTGGTGCAGGTGGAAG
PmCBFaR1	Cloning of tandem array of <i>PmCBFca</i>	TCCATATCCACGCTAAAAATCT
PmCBFbF1	Cloning of tandem array of <i>PmCBFbc</i>	GACCACCCTACGCACTTCGGAC
PmCBFcR1	Cloning of tandem array of <i>PmCBFbc</i>	CCCTTCTTGTCTCTCTTCCACCTG
M13F	Sequencing	AGCGGATAACAATTTACACAGG
M13R	Sequencing	CGCCAGGGTTTTCCAGTCACGAC
PmEF1αF	quantitative RT-PCR of <i>PmEF1α</i>	CGGATTCAATGTTAAGAATGTTGC
PmEF1αR	quantitative RT-PCR of <i>PmEF1α</i>	AGAACTGGAGCATATCCGTTACC
PmCBFaqF	quantitative RT-PCR of <i>PmCBFa</i>	GTGTCTTTTTCGTCGGATTCATCTC
PmCBFaqR	quantitative RT-PCR of <i>PmCBFa</i>	CTATTAGAGCCACAGTGACATGCTC
PmCBFbqF	quantitative RT-PCR of <i>PmCBFb</i>	GTCAATCTGTCTCTCGGGTTTAGTG
PmCBFbqR	quantitative RT-PCR of <i>PmCBFb</i>	CTAGAAACTAGACAAAGCACAGTACC
PmCBFcqF	quantitative RT-PCR of <i>PmCBFc</i>	CTCCATCGCAATGCTTAGGTG
PmCBFcqR	quantitative RT-PCR of <i>PmCBFc</i>	CGACTGACGTTAATCTTTTTTCGTTAC

Table S2. The selected cis-elements predicted in the putative promoter of *PmCBFb*

Sort	Factor or site name	Signal sequence	Number
Tissue specific expression elements	DOFCOREZM	AAAG	19
	CACTFTPPCA1	YACT	12
	ROOTMOTIFTAPOX1	ATATT	6
	OSE2ROOTNODULE	CTCTT	5
	OSE1ROOTNODULE	AAAGAT	4
	TAAAGSTKST1	TAAAG	3
	GTGANTG10	GTGA	2
	POLLENILELAT52	AGAAA	1
Light response elements	INRNTPSADB	YTCANTYY	10
	-10PEHVPSBD	TATTCT	2
	SORLIP1AT	GCCAC	2
	SORLIP2AT	GGGCC	2
	GT1CONSENSUS	GRWAAW	1
	IBOXCORE	GATAA	1
Binding sites for MYB factors	MYBCORE	CNGTTR	5
	TATCCAOSAMY	TATCCA	3
	ECCRCAH1	GANTTNC	2
	MYBPLANT	MACCWAMC	1
	MYBST1	GGATA	1
Binding sites for MYC factors	MYCCONSUSAT	CANNTG	10
	T/GBOXATPIN2	AACGTG	1
Binding sites for WRKY factors	WRKY71OS	TGAC	7
	WBBOXPCWRKY1	TTTGACY	1
Binding sites for RAV factors	RAV1BAT	CACCTG	2
	RAV1AAT	CAACA	1
Abscisic acid related elements	ACGTABREMOTIFA2OSEM	ACGTGKC	4
	GADOWNAT	ACTGTGC	2
Heat shock	CCAATBOX1	CCAAT	3

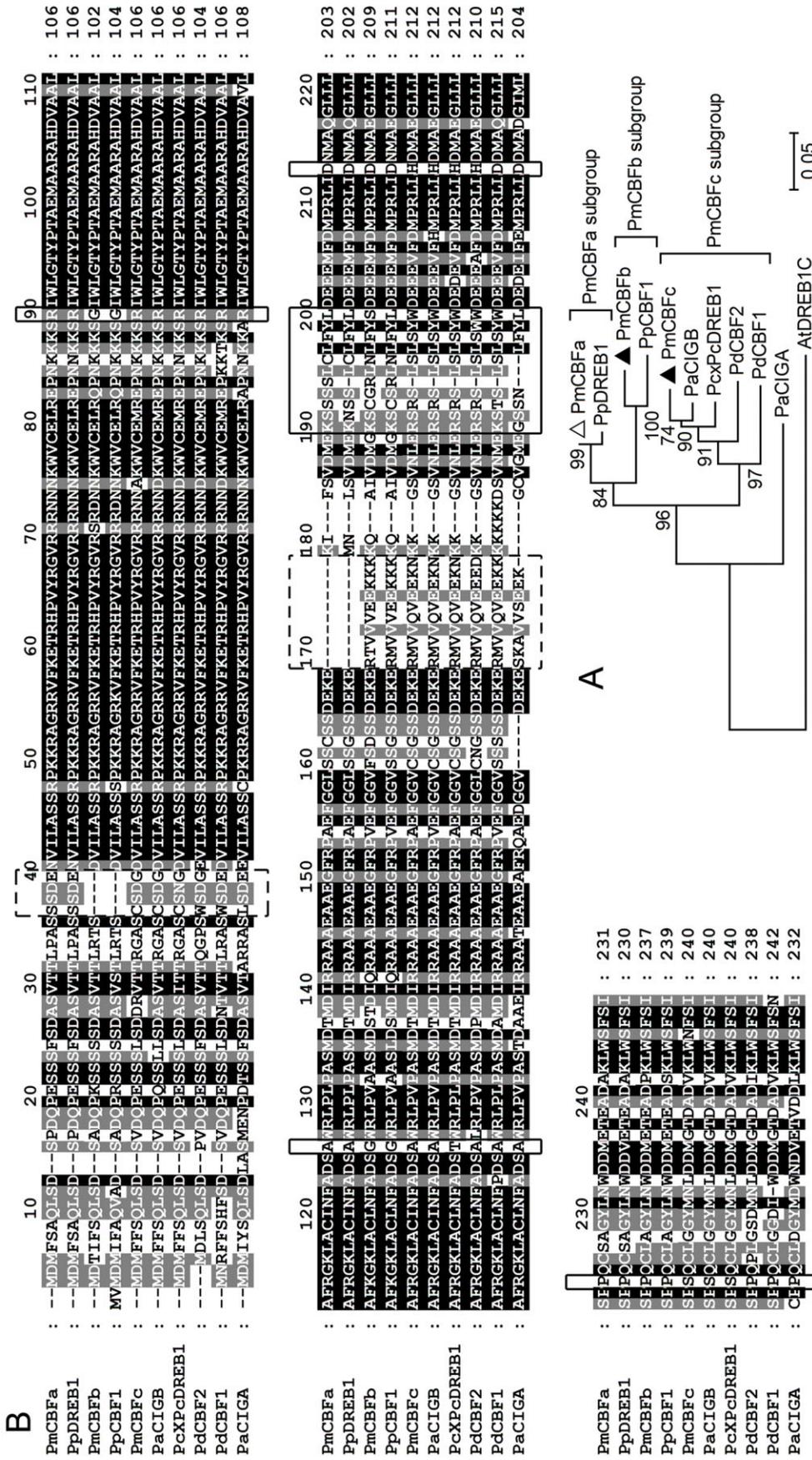


Fig. S1. Sequence analysis of CBF proteins in *Prunus* spp. **A** - Phylogenetic analysis of CBF/DREB1 proteins from *Prunus* spp. using the Neighbor-Joining method, with bootstrap analysis with 1000 replications. The AtDREB1C is used as an outgroup for better calculation. The sequences used in these analyses are as follows: PmCBFa (ADF43033.1), PmCBFb (ADF43034.1), and PmCBFc (ADF43035.1) from *Prunus mume*; PpDREB1 (ABR1931.1) and PpCBF1 (ADU03762.1) from *P. persica*; PaCIGA (BAC20183.1), and PaCIGB (BAC20184.1) from *P. avium*; PpCBF1 (AFL48190.1) and PdCBF2 (AFL48191.1) from *P. dulcis*; PcxPcDREB1 (ACF94686.1) from *P. canescens*; P. *cerasus*; AtDREB1C (NP_567719.1) from *Arabidopsis thaliana*. **B** - Multiple alignment of the sequences from *Prunus* spp. Diversity in conserved regions and non-conserved regions are indicated by *solid* and *dashed* boxes, respectively.