

Table S1. Estimated H- and G-lignin content in RNAi-*PpMYB8* silenced lines compared to control (RNAi-EV). Asterisk indicates significant differences between RNAi-*PpMYB8* plants and controls ($P = 0.032$, $n = 3$).

sample	%H-monomers	%G-monomers
RNAi- <i>PpMyb8</i>	1.5 ± 0.3%*	98.5 ± 0.3% *
RNAi-EV	2.1 ± 0.2%	97.9 ± 0.2 %

Table S2. Primer sequences used for cloning.

Primer name	Primer sequence (5'→3')
p1830.5Fwd	ATGATCCAAGAAATGCAAGTCACTC
p1830.7Rvs	TCTACAGATATCGATGAGTCTCG
p1830.4Fwd	TTAGGGTATTTTGACTCCTGTC
p1830.6Rvs	TTTTCTGGAATTTGAGGTTTAC
p3030.2Fwd	GTACAACACTTGATTAAGGTTGCG
p3030.2Rvs	CTTGGAATCGTCGCTGTTGCTTC
p61682.2Fwd	CATAGAACTTCACATCGCTTCC
p61682.2Rvs	AAAGGCACATCTCAGAGCC
PA1 AttB4	ATAGAAAAGTTGGATTCCACTTAGGACTCCACCCC
PA2 AttB4	ATAGAAAAGTTGGAGCCACAGACGATGATGACCCA
PA3 AttB4	ATAGAAAAGTTGGAACCCTAGATCAGCATGAGCTA
pA AttB1R	TTGTACAAACTTGCTTTTCTGGAATTTGAGGTTTAC
pD1 AttB4	ATAGAAAAGTTGGAGTACAACACTTGTTAAGGT
pD2 AttB4	ATAGAAAAGTTGGACCAGGGTGGGAAAGCTGGTG
pD3 AttB4	ATAGAAAAGTTGGAATGGGATGGAGCAGAATATATAC
pD AttB1R	TTATACAAACTTGAATTGGAATCGTCGCTGTTGC
PI1 AttB4	ATAGAAAAGTTGGACTTCCATAAATATTGAAACAC
PI2 AttB4	ATAGAAAAGTTGGAGATTGGCAAGATGGATAAAG
PI3 AttB4	ATAGAAAAGTTGGACAAAACACTAGATTGAGAAAGG
PI AttB1R	TTGTACAAACTTGCAGCCTTGTAGATAATTAGTG
AttB4	GGGGACAACCTTTGTATAGAAAAGTTG
AttB1R	GGGGACTGCTTTTTTGTACAAACTTG
MYB8 AttB1	AAAAAGCAGGCTTAATGGGGCGCCACTCGTG
MYB8 AttB2	AGAAAGCTGGGTTCTAAATTTGGTCCAGAAC
AttB1	GGGGACAAGTTTGTACAAAAAAGCAGGCT
AttB2	GGGGACCACTTTGTACAAGAAAGCTGGGT

Table S3. Primers pairs used for RT-qPCR.

Gene name	Gene description	Forward / Reverse primers sequences (5'→3')
<i>PpADT-A</i>	Arogenate dehydratase-A	GACTATACTTCATGTACCAAGC / CCACGATCTTCAATATCTCTGG
<i>PpADT-B</i>	Arogenate dehydratase-B	AGTGACATGGGGAAGTCACC / CGATACACATATTTTCTCTCTAC
<i>PpADT-C</i>	Arogenate dehydratase-C	ATGAAGGCGTGATTTGAAGC / ACTCTGCCACAGAGGGACAC
<i>PpADT-D</i>	Arogenate dehydratase-D	CGGTGAAAATAAGAGACTGGG / TGTATATGACCAGTAGCTGCTC
<i>PpADT-E</i>	Arogenate dehydratase-E	CAAATCCGTGTTCTCTGCTGG / TGAGGATCCTGAAGTTCCACG
<i>PpADT-F</i>	Arogenate dehydratase-F	CTCCGTTCTTGTGACAGCTCG / TTTGATCCTCAAATCCCTGTAC
<i>PpADT-G</i>	Arogenate dehydratase-G	GTTTCATGTAAGTGAAGTGGG / AACTTAGTGTACACCCTCTC
<i>PpADT-H</i>	Arogenate dehydratase-H	CAGTTGTATCAGAGGTATCGG / TGGCGTTGGCTAGATCCTTG
<i>PpADT-I</i>	Arogenate dehydratase-I	ATGTGTACGCATGTCTGTATAG / CCATGAAGATCCTCGCGATTG
<i>PpCM1</i>	Chorismate mutase-1	GAACCTCCATGGATATACCTGC / AGCAATTCAATTTGCTGACTGC
<i>PpCM2</i>	Chorismate mutase-2	GCCGTTAAATTGTTGTGTCTACC / TGATTTTCCCGAACCAGAAG
<i>PpPAT</i>	Prephenate/rogenate aminotransferase	GCAGTCATCATTGCCGAAGGC / AGATTGCAAGCCATGGAGGG
<i>PpPH</i>	Phenylalanine hydroxylase	AGCTATTGGAATCACCTCACC / GGCAATATCCCTCATCTTGTC
<i>PpPAL1</i>	Phenylalanine ammonia lyase-1	GATTTGCATCCCCTGGATT / AAGCAGTACACGATCACCCA
<i>PpPAL2</i>	Phenylalanine ammonia lyase-1	GCAGCACGTTTTGCATAGAA / TAATATTTCCCAACACCGGC
<i>PpMYB8</i>	MYB8	CCACACGCAAGAGGAGACTG / GGCGGAATTTAGGGCAGCGTGG
<i>PpEF1α</i>	Elongation factor 1 α	TGCTGTTGGAGTCATCAAGG / CTCGTGCATCAGAATCAGACA
<i>PpActin2</i>	Actin	ATCTCTCAGCACATTCCAACAG / TATTGCCACCATCATCTCAA
<i>sp_v3.0_unigene32645</i>		TGGCTTCTCTTCCAATGCTC / ACCTGCTCAACCACTTCCAA
<i>sp_v3.0_unigene93974</i>		TGTAGGCATAAGAACTCCATCAA / CTGTGACAACTTCTCCGTGC
<i>sp_v3.0_unigene93562</i>		AAAACAGGGGCCTTACGAAG / ATGGTATATTGCAGGCAGGC
<i>sp_v3.0_unigene24132</i>		TGTGGTCTACTTGCCCTCTG / CGTGACTTGGGCTCATCATC
<i>sp_v3.0_unigene133330</i>		TGGCCAAAATCCCAAGTTTTTCT / ACAGTCATATGTTCTTCCCCTT
<i>sp_v3.0_unigene22584</i>		TGGTTCAGAGGAGGCAAAGA / AATAGGGCTGTGCTCCAAC
<i>sp_v3.0_unigene6586</i>		CAGAATCGCCAGGGTATGA / ATGCGCGTAGAAAGAACC

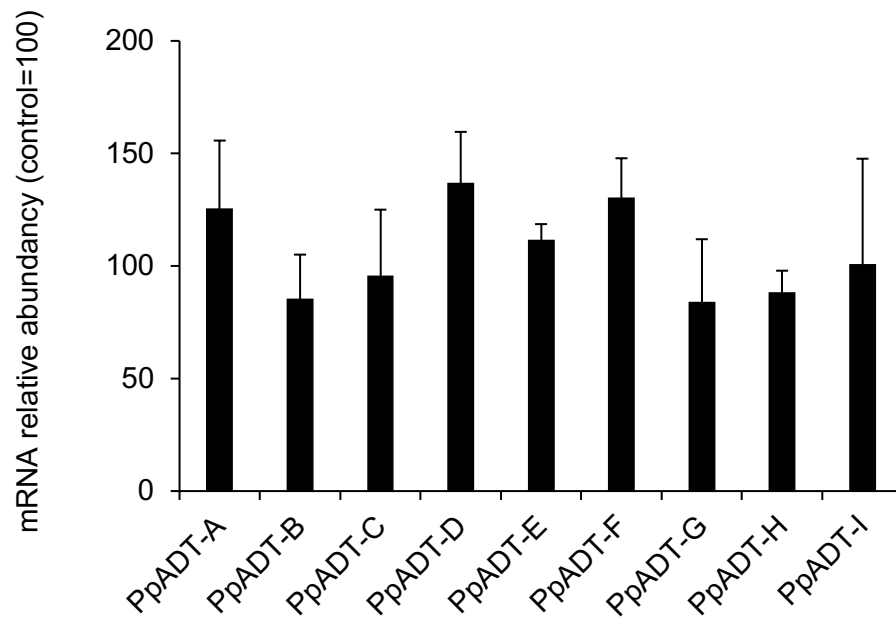


Figure S1. Average expression level of the nine ADT genes from *P. pinaster* in the OE-*PpMYB8* lines, expressed as percentage to controls.

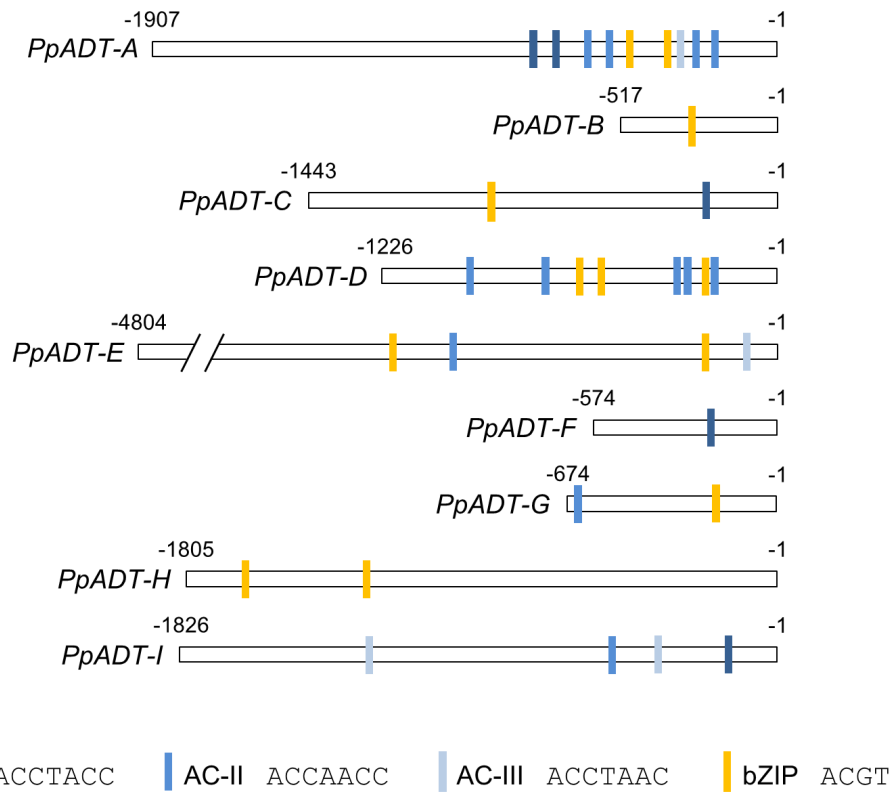


Figure S2. *In silico* analysis of candidate AC- and bZIP-binding elements within the 5'-flanking region of the ADT genes in *Pinus pinaster*. Genomic DNA sequences upstream of the initiation codon were manually assembled from the information available at *SustainPineDB*, using their *P. taeda* homologues as a reference. Different colours, as indicated in the key, were used to mark the three different AC-binding elements previously described as putative *PpMYB8* binding sites, and the putative ACGT bZIP binding sites.

PpADT-A proximal 5'-flanking region (P_{ADT-A})

TAGGAGACGACCCACCTTTCAAGAAAAGGCTCTCATTAGAACACACT **CCAACCA**ATCCCGGGC
CACCCCGGGGAAACCGGGTCT **ACGT**AGACCGGTTCCCCGCCCCCTGAAAAGGCCAAAGTGCAAG
CATCAAAGACAAATGAGGTTCCACTTTTTGGCCACAGACGATGATGACCCAAGAACAACAGACT
probeA2
GGTAAGGCAC **ACGT**GCTTCCACGAGATCCACTAACTTGCGCGGGTGTTTACAC **ACCTAAC**CACT
probeA1
CCGGAGCTTCCATGCTTTCATCCACGAGTCAGAGGGGCCTTTTCCT **CCAACCA**CCCTAGATCAGCA
TGAGCTATTTTATCTACTACCCCTCC **ACCAACC**CACCACCACCACAAACAATCTTAACAGAAC
TCTCTATTAGTCTAAGCACGCCTACAAATGTTTTGCAGGCGAACCCAACCTGGTTCGGACGAACG
AGTGTGTGATGTGCCAAACACATACTCGCGTCCCAGATCTGAACACTCTGAGCCCTCGGCCCA
AATTTTCAGGGGGCTAAAGTAAACCTCAAATTCCAGAAA **ATG**

PpADT-D proximal 5'-flanking region (P_{ADT-D})

CTTTCTCATTCTGGTTTGGTAAGCACTTAACAAAAATTTCCAGATTGGACAGATAATTTTTGAG
TCCAACTAGGTTGTTACATTTGGCAGAGTAGATGGCAGGGTTGTGTAGAATGGTAAGTAAGGTG
GGGGAGGACAAAACCTCAGTTTACTTGGTTTTCAATCCTGATGGTGCGCAGTCTCATTCCAAGTC
probeD1
TTAGATGCGTGGGGGCCAGGGTGGGAAAGCTGGTGAGAAGGG **CCAAACC**AAAAAAAA **CCAAAC**
probeD2
GAAATCACAGCGTACGATCTAACTAACCCTGTTGAATAGTCCCCTG **ACGT**AGCCGCCGTAATGT
GCAACA **ACCCACC**ATGGGATGGAGCAGAATATATACAAAACAAGCAAAGGGTGGCCTCGCCTGC
TTCTCTTTATTGTCATTGCCTTGCCAGCTTTGGAAAATCTTTAGCTTTGGAAAATCTTTAGC
TCTGAGTGAAGAAAAGAGGACTGAACAGAAAGAGATTTGCAGCTGCAAATTTCCAGGAGCAGTTA
TTTGAAGCAACAGCGACGATTCCAAG **ATG**

Figure S3. Probes designed for the EMSA in the P_{ADT-A} and P_{ADT-D} sequences. ATG codon is marked at the 3'-end. Putative AC-binding elements are marked in cyan; putative bZIP ACGT boxes are marked in yellow.