

**Table S2** Summary of significant SNP marker-trait pairs identified at the threshold of  $P \leq 0.05$ , using the mixed linear model (MLM) in the *Populus tomentosa* association population.

Trait	Locus	Position	Mutation	Association population ( $N = 460$ )		
				P-value	Q-value	R <sup>2</sup> (%)
Lignin						
	SNP2	Promoter	[G : T]	0.0069	Q > 0.10	3.4
	SNP24	Promoter	[A : T]	0.0434	Q > 0.10	3.0
	SNP44	Intron1	[G : T]	0.0012	0.0551	2.1
	SNP47	Exon 2	[T : C]	0.0500	Q > 0.10	1.9
	SNP49	Exon 3	[C : A]	0.0025	0.0810	3.0
$\alpha$ -cellulose						
	SNP3	Promoter	[A : G]	0.0015	0.0629	2.2
	SNP18	Promoter	[T : A]	0.0011	0.0551	2.5
	SNP36	promoter	[A : T]	0.0061	Q > 0.10	3.5
	SNP41	5'UTR	[T : C]	3.02E-05	0.0035	1.6
	SNP48	Intron 2	[A : T]	0.0400	Q > 0.10	2.2
	SNP49	Exon 3	[C : A]	0.0031	0.0948	5.3
	SNP50	Intron 3	[C : A]	0.0142	Q > 0.10	1.7
Holocellulose						
	SNP27	Promoter	[G : T]	0.0281	Q > 0.10	6.0
	SNP38	Promoter	[T : G]	0.0052	Q > 0.10	2.9
	SNP45	Exon 2	[A : C]	0.0002	0.0142	4.0
	SNP48	Intron 2	[A : T]	0.0351	Q > 0.10	4.4
	SNP62	Intron 6	[T : G]	0.0104	Q > 0.10	3.4
	SNP70	Intron 9	[A : T]	0.0069	Q > 0.10	2.5
	SNP81	Intron 10	[T : C]	0.0002	0.0142	3.0
	SNP88	Exon 12	[G : A]	0.0161	Q > 0.10	1.9
Fiber length						
	SNP2	Promoter	[G : T]	0.0355	Q > 0.10	6.7
	SNP44	Intron1	[G : T]	0.0194	Q > 0.10	2.0
	SNP89	3'UTR	[G : C]	0.0070	Q > 0.10	1.5
	SNP90	3'UTR	[G : A]	0.0122	Q > 0.10	1.9
Fiber width						
	SNP58	Exon 6	[T : C]	0.0500	Q > 0.10	3.9
	SNP59	Exon 6	[A : C]	0.0008	0.0440	2.6
	SNP62	Intron 6	[T : G]	0.0475	Q > 0.10	1.8
	SNP70	Intron 9	[A : T]	0.0477	Q > 0.10	1.8
Microfibril angle (MFA)						
	SNP10	Promoter	[G : A]	0.0372	Q > 0.10	1.7
	SNP47	Exon 2	[T : C]	0.0195	Q > 0.10	2.6

Diameter at breast height (D)						
	SNP48	Intron 2	[A : T]	0.0009	0.0454	1.9
	SNP59	Exon 6	[A : C]	0.0151	Q > 0.10	3.0
	SNP62	Intron 6	[T : G]	0.0195	Q > 0.10	2.0
	SNP75	Exon 10	[T : C]	3.15E-05	0.0035	3.2
	SNP81	Intron 10	[T : C]	0.0003	0.0195	2.0
Tree height (H)						
	SNP44	Intron1	[G : T]	0.0179	Q > 0.10	1.2
	SNP49	Exon 3	[A : T]	0.0012	0.0551	4.3
	SNP89	3'UTR	[G : C]	0.0351	Q > 0.10	2.1
Stem volume(V)						
	SNP75	Exon 10	[T : C]	3.02E-05	0.0035	4.6
	SNP81	Intron 10	[T : C]	0.0143	Q > 0.10	2.0

$R^2$  = percentage of the phenotypic variance explained.  $Q$ -value = correction for multiple testing [false discovery rate FDR ( $Q$ )  $\leq$  0.10]