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Overexpression of a serine hydroxymethyltransferase increases biomass production and reduces recalcitrance in the bioenergy crop *Populus*†

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Cell wall recalcitrance is the major obstacle for plant biomass conversion to biofuels. In this study, we functionally characterized a serine hydroxymethyltransferase (SHMT) from *Populus* and evaluated its potential for developing lignocellulosic feedstocks. SHMT is an enzyme that plays an important role in cellular one-carbon pathways. However, little is known about its function in plant cell wall-related processes. Among nine SHMT genes in the *Populus* genome, *PtSHMT2* was highly expressed in the developing xylem and was co-expressed with secondary cell wall biosynthetic genes. In *Populus* transgenic plants overexpressing *PdSHMT2*, the biomass yield and sugar (glucose and xylose) release were increased whereas the lignin content was decreased. Transcriptomics and metabolomics analyses revealed that genes and metabolites related to secondary cell wall biosynthesis were affected by *PdSHMT2* overexpression. Based on the transcription factor binding sites of differentially expressed genes in *PdSHMT2* overexpression lines, a total of 27 hub transcription factors were identified. We proposed a regulatory model of action of *PdSHMT2* with transcriptional master switches of secondary cell wall biosynthesis. Collectively, these results suggest that *PdSHMT2* is a promising candidate for genetic engineering to improve feedstock characteristics to enhance biofuel conversion and reduce the cost of lignocellulosic biofuel production.

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Introduction

Serine hydroxymethyltransferase (SHMT, EC 2.1.2.1) is a pyridoxal phosphate-dependent enzyme that plays an important role in cellular one-carbon (C₁) pathways by catalyzing the reversible conversions of L-serine to glycine and simultaneously tetrahydrofolate (THF) to 5,10-methylene THF.¹ Prokaryotes have single gene encoding SHMT. In animals and fungi, two SHMT isoforms (cytosolic and mitochondrial) are encoded by two distinct genes.² The structure of the SHMT monomer is similar across prokaryotes and eukaryotes, whereas its functional form is a dimer in prokaryotes and a tetramer in eukaryotes.¹ The structure of SHMT directly affects its function. For example, for sheep liver cytosolic SHMT (scSHMT), the mutation of D227 leads to formation of inactive dimers, E74 controls the conversion between “open” and “close” forms, and K256 plays a crucial role in maintaining the tetrameric structure.¹ In addition, the function of SHMT is affected by post-translational modification, such as ubiquitination and



sumoylation. Anderson *et al.* (2012) reported that mammalian SHMT1 interacts with an E2 conjugase, Ubc13, which mediates competition ubiquitination or sumoylation. The ubiquitination is required for SHMT1 nuclear export and increases its stability within the nucleus, whereas sumoylation of SHMT1 is involved in nuclear degradation.³ Moreover, SHMT plays a role in transcriptional regulation. For example, human cytoplasmic SHMT (hcSHMT) protein can bind mRNA and displays increased affinity for the 5' untranslated region of its mRNA.⁴

To date, SHMT in plants has been reported to function in the photorespiratory pathway and stress responses. In the photorespiratory pathway, SHMT is associated with the glycine decarboxylase complex (GDC),⁵ which catalyzes the decarboxylation of photorespiratory Gly to yield NH₃, CO₂ and a C₁ unit that is transferred to THF.⁶ Mutants of mitochondrial SHMT or GDC exhibit symptoms of chlorosis.⁷ In *Arabidopsis*, the circadian clock regulates the expression of genes encoding mitochondrial components of the photorespiratory pathway, including *AtSHMT1* and *AtSHMT4*.⁸ Potato plants with antisense *SHMT* have lower photosynthetic capacity and accumulate glycine in light.⁹ In addition, SHMT is involved in biotic and abiotic stress responses. The *Arabidopsis shmt1-1* mutant is more susceptible to infection with necrotrophic and biotrophic pathogens, and salicylic acid-induced genes and H₂O₂ detoxification-related genes are constitutively activated in *shmt1-1* plants. Under abiotic stresses, *shmt1-1* plants exhibit hypersensitivity to salt stress.¹⁰ SHMT is degraded by 26S proteasome, but it can be stabilized by a ubiquitin-specific protease, UBP16, through deubiquitination of SHMT1 in *Arabidopsis*.¹¹ So far, the function of SHMT in cell wall-related processes has not been reported. Recently, several other enzymes in C₁ metabolism have been reported to play important roles in cell wall-related processes. In C₁ metabolism, methylenetetrahydrofolate reductase (MTHFR) catalyzes the conversion of 5,10-methylenetetrahydrofolate to 5-methyltetrahydrofolate, while folylpolyglutamate synthase (FPGS) catalyzes the addition of glutamate moieties to folate and folate derivatives. Both MTHFR and FPGS can affect lignin biosynthesis in maize.^{12,13} Cystathionine γ -synthase (CGS) catalyzes the formation of cystathionine from cysteine and an activated derivative of homoserine whereas *S*-adenosylhomocysteine hydrolase (SAHH) is responsible for the reversible hydration of *S*-adenosyl-L-homocysteine into adenosine and homocysteine. In switchgrass, down-regulation of *CGS* enhanced lignin biosynthesis whereas down-regulation of *SAHH* reduced the lignin biosynthesis and increased cell wall saccharification efficiency.¹⁴

In vascular plants, secondary cell walls (SCWs) are the most abundant renewable plant biomass, and are widely used for many applications including energy, textiles, pulping and paper-making. SCWs are composed of lignin, cellulose and hemicelluloses. The deposition of SCWs allows cells to function as mechanical tissues for structural support and protection.¹⁵ Many transcription factors (TFs) have been identified controlling SCW biosynthesis in a hierarchical manner. In the first layer of the SCW regulatory network, SND1 and NST1 are master regulators of secondary wall biosynthesis in fibers.¹⁶ The

second-layer master switches including two functional redundant MYB genes, MYB46 and MYB83, which activate the expression of the third-layer TFs and enzymes which are directly involved in SCW biosynthesis.¹⁷ MYB4 is a transcriptional repressor that negatively regulates *C4H* and itself in plants.¹⁸ The *Arabidopsis myb4* mutant accumulates sinapate esters in leaves.¹⁹ In switchgrass, overexpressing its orthologous *PvMYB4* reduced the lignin content and ester-linked *p*-CA : FA ratio and increased sugar release.²⁰ It has been reported that the expression of *MYB4* can be induced by glucose.¹⁸ However, whether other metabolites or enzymes in metabolic pathways are involved in the SCW regulatory pathway is largely unknown.

In this study, we characterized an *SHMT* gene in *Populus* encoded by the locus Potri.001G320400 (*PtSHMT2*), which has strong expression in the developing xylem during poplar wood formation. We overexpressed this gene in *P. deltoides* 'WV94' and found that it enhanced plant growth and altered cell wall composition and sugar release in transgenic plants. By analyzing the transcriptomics and metabolomics data using *PdSHMT2* overexpression lines, we examined the potential molecular mechanisms of action of *PdSHMT2* in the secondary cell wall biosynthesis and revealed the association of *PdSHMT2* with the transcriptional master regulators of secondary cell wall biosynthesis. Collectively, these findings provide new insights into the design of genetic engineering leading to cost-effective biomass conversion into biofuels.

Results and discussion

SHMT family in *Populus*

To study the function of *SHMT* in *Populus*, we first analyzed the whole gene family of *SHMT* in *Populus* and other plant species. The *SHMT* genes were queried using BLAST in *Populus trichocarpa* genome and 11 other plant species, including three other woody species *Salix purpurea*, *Eucalyptus grandis* and *Vitis vinifera*, three annual dicots *Arabidopsis thaliana*, *Medicago truncatula*, and *Glycine max*; four monocots *Oryza sativa*, *Zea mays*, *Brachypodium distachyon* and *Sorghum bicolor*, and a moss *Physcomitrella patens*. A total of 95 *SHMT* genes were identified in these 12 species, and the *SHMT* family in dicots (7–15 members) is larger than that in monocots (4–7 members) (Fig. 1 and Table S1 in the ESI†). Among those nine *SHMT*s identified in *Populus*, namely *PtSHMT1–9*, three paralogous pairs (*PtSHMT2/9*, *PtSHMT3/5* and *PtSHMT7/8*) were likely generated by a Salicoid genome duplication and rearrangement event (Fig. S1 in the ESI†).

On the basis of the expression data from different tissues of *P. trichocarpa* (*Populus* Gene Atlas) (<https://phytozome.jgi.doe.gov>), we compared the expression patterns of the nine *PtSHMT* genes. Noticeably, only *PtSHMT2* and its paralog *PtSHMT9* showed high abundance across poplar tissues, and *PtSHMT2* was strongly expressed in the stem (Fig. 1B). Phylogenetic (Fig. 1A) and sequence similarity (Fig. S2 and S3 in the ESI†) analyses indicated that *PtSHMT2* is closely clustered with *AtSHMT4* (At4g13930). The pairwise correlation of the nine *PtSHMT* expression patterns indicated that two *PtSHMT*



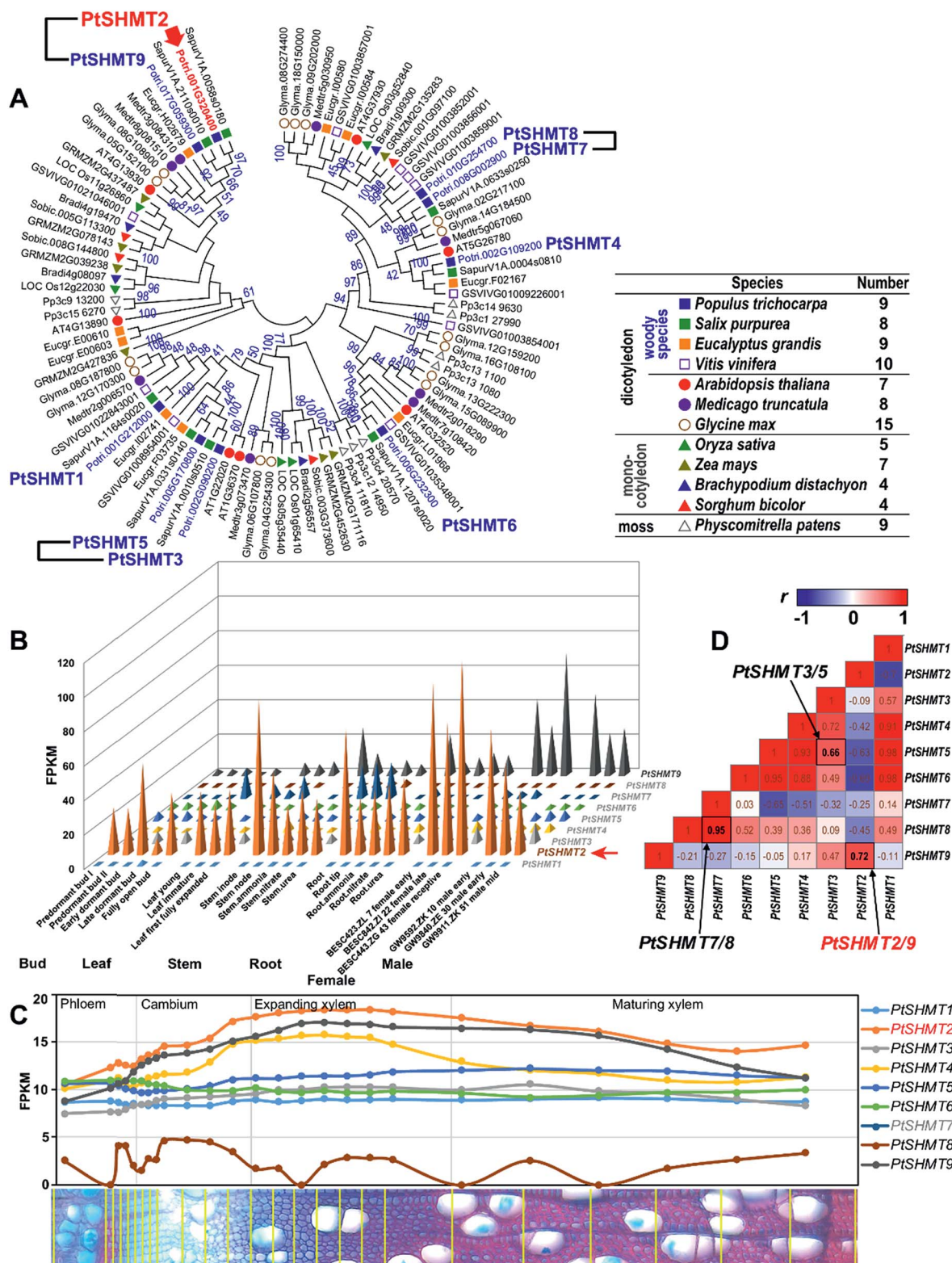


Fig. 1 Phylogenetic analysis and expression patterns of *SHMT* genes. (A) Phylogenetic analysis. The phylogenetic tree was constructed using the full-length amino acid sequences of *SHMT* proteins from 12 plant species by the neighbor-joining (NJ) method with 1000 bootstrap replicates. Three *PtSHMT* paralogous pairs generated by a whole-genome duplication (WGD) event (Fig. S1 in the ESI†) are shown by linked black lines. (B) Expression patterns of *PtSHMT* genes across various tissues. (C) Expression patterns of *PtSHMT* genes during wood formation. The nm-scale expression of *PtSHMT* genes was obtained from the AspWood gene expression database.²¹ (D) Pairwise comparison of the Pearson correlation coefficient (r) of expression in the *PtSHMT* family based on results in (B) and (C). The three paralogous pairs are labelled by arrows.

paralogous pairs (*PtSHMT2/9* and *PtSHMT7/8*) have highly positive correlation coefficients ($r = 0.72$ and 0.95) compared to pair *PtSHMT3/5* ($r = 0.66$) (Fig. 1D). In contrast, the promoter similarity of pair *PtSHMT3/5* is higher than that of the other two pairs (Fig. S4 in the ESI†). The divergence between the expression and promoter sequence implies that the expression of *PtSHMT* paralogous pairs may be controlled by different mechanisms. Furthermore, the expression patterns of the nine *PtSHMTs* developed from the AspWood gene expression database were compared in nm-scale stem tissues of *P. tremula*.²¹ Among the nine *PtSHMTs*, *PtSHMT2* has the highest abundance and was induced in expanding xylem (Fig. 1C), implying that it may play an important role in SCW-related processes.

To explore the possible regulatory mechanism of *PtSHMT2* expression, we analyzed the *cis*-acting elements of the promoter region (3000 bp upstream of the translation initiation site). Based on the functional annotation, the identified *cis*-acting elements were classified into four groups – development, hormone, stress and others. Noticeably, seven circadian

elements (involved in circadian control), three ABREs (involved in ABA responsiveness), two CAT-box elements (related to meristem expression) and two MYB binding sites (MBS and MRE) were identified in the promoter of *PtSHMT2* (Fig. S5 in the ESI†). In *Arabidopsis*, the *shmt1-1* mutant is defective in mitochondrial SHMT activity and displays a lethal photorespiratory phenotype when grown at the ambient CO₂ level.²² The expression of genes of the photorespiratory pathway has been shown to be regulated by a circadian clock. *Arabidopsis SHMT1* and *SHMT4* exhibit circadian oscillation in mRNA abundance.⁸ This is consistent with the highest abundance of circadian elements in the promoter of *PtSHMT2*.

We then constructed a co-expression network of *PtSHMT2* based on the global gene expression patterns across various tissues and under different stresses in poplar. According to the functional classification, a number of cell wall-related genes were identified in the co-expression network (Fig. S6 and S7 in the ESI†). These include several key transcription factors (TFs) such as NST1, SND3, MYB46, MYB103, NAC073, NAC075,

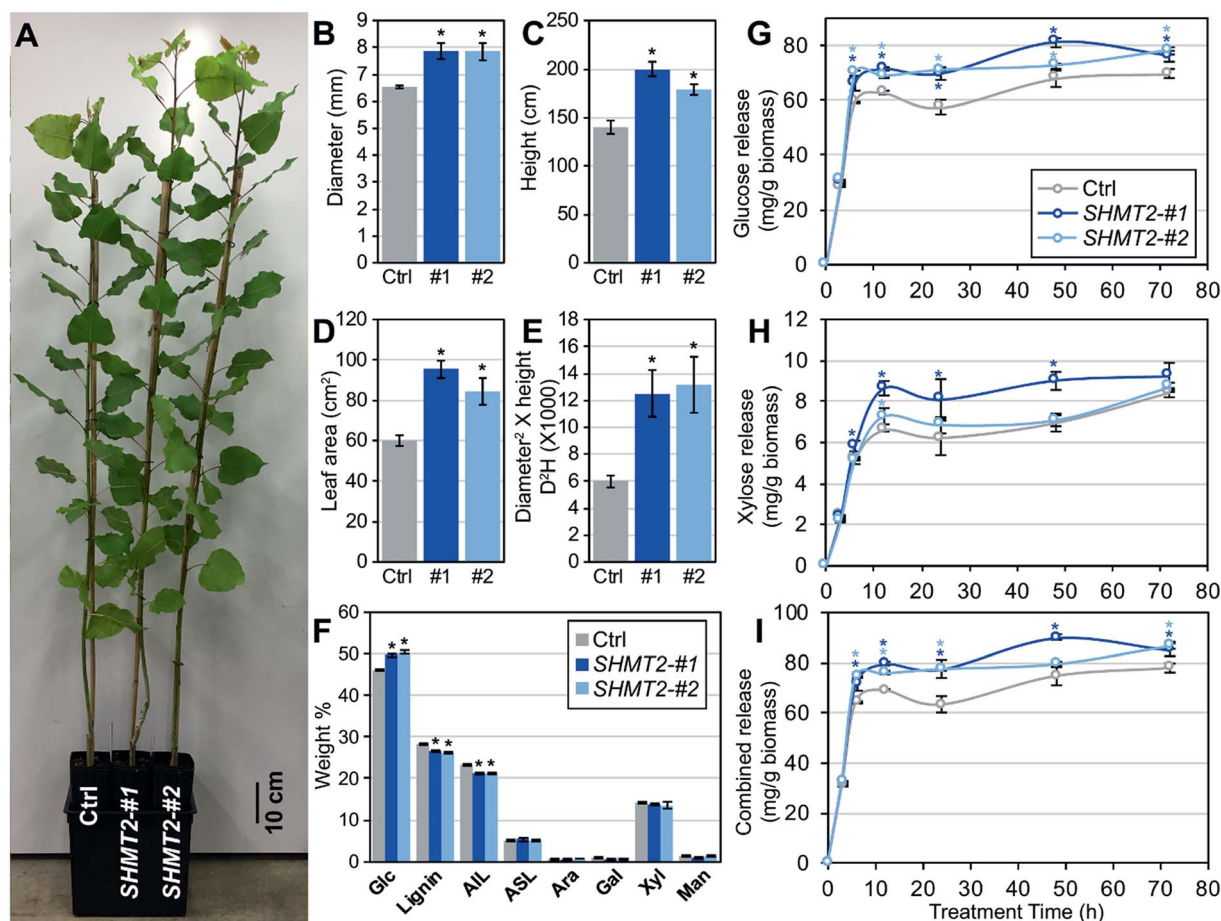


Fig. 2 Plant growth characteristics and sugar release of *PtSHMT2* overexpression plants. (A) *PtSHMT2* overexpression transgenic plants grown under greenhouse conditions. The diameter of basal stem (B), height (C) and leaf area (D) were compared between transgenic lines and control plants. The above-ground biomass (E) of transgenic poplar samples was estimated using the formula diameter² × height (D^2H , mm³). (F) Chemical composition analysis by means of ion chromatography after two-step acid treatment. Glc, glucose; ALL, acid insoluble lignin; ASL, acid soluble lignin; Ara, arabinose; Gal, galactose; Xyl, xylose; Man, mannose; ash content (%) is not included. (G–I) Saccharification efficiency of *PtSHMT2* overexpression poplars. Glucose (G), xylose (H) and combined sugar (I) release with enzymatic hydrolysis times (0, 3, 6, 12, 24, 48 and 72 h). Shown are mean values of three biological replicates ± S.E.



KNAT7 and WOX14, and important enzymes such as LAC10, LAC17, PRX44, PRX53, MAN6, IRX2, XTHs and UGT80A2 which are known regulators of SCW biosynthesis.^{15,23} These results supported the role of *PtSHMT2* in cell wall-related processes.

Overexpression of *PdSHMT2* increases plant growth

To directly examine the potential function of *SHMT2* in *Populus*, we cloned the full-length open reading frame of *SHMT2* from *P. deltoides* 'WV94' (*PdSHMT2*) and overexpressed it in 'WV94' using the *UBIQUITIN3* constitutive promoter. Eight independent transgenic lines were generated and two independent lines (#1 and #2) with high expression of *PdSHMT2* were selected for further analysis (Fig. S8 in the ESI†). Compared to the control plants, the two transgenic lines showed increases in above-ground biomass; the diameter increased by ~25%, height increased by ~46%, leaf area increased by 64% and diameter² × height (D^2H) increased by ~133% (Fig. 2A–E). Increasing the yield of bioenergy crops is one of the major goals of bioenergy engineering.²⁴ This results demonstrated that overexpression of *PdSHMT2* can effectively increase the feedstock yield in *Populus*.

Cell wall chemical composition and sugar release

The cell wall composition is directly associated with the efficiency of biomass conversion to biofuels. To explore whether cell wall composition is affected by *PdSHMT2*, we investigated the carbohydrate composition in stem tissues by means of ion

chromatography after two-step sulfuric acid hydrolysis procedures. Compared to control plants, the two transgenic lines (#1 and #2) had significantly higher glucose content (7.37% and 8.72% increase, respectively) and lower lignin content (6.05% and 7.57% decrease, respectively), whereas there were no significant changes in the contents of arabinose, galactose, xylose or mannose (Fig. 2F). The reduced lignin content in transgenic lines was mainly due to the reduction of acid insoluble lignin (AIL) (Fig. 2F).

To assess the sugar release performance of the *PdSHMT2* overexpression lines, glucose and xylose releases during the enzymatic hydrolysis were monitored. At 6 h hydrolysis, the two transgenic lines already showed higher glucose release than the control plants. At the final time point of 72 h hydrolysis, the glucose release increased by 9.38% and 12.4% in #1 and #2, compared to the control, respectively (Fig. 2G). More xylose release was detected in #1 at 6 h and increased by 31.4% at 12 h (Fig. 2H). The total sugar release in the two transgenic lines was constantly higher than in the control plants from 6 h and increased by 9.4% and 11.4% at 72 h (Fig. 2I) for lines #1 and #2, respectively.

The structural information on lignin, such as lignin S/G ratio, level of lignin subunit content and lignin interunit linkages, was analyzed through nuclear magnetic resonance (NMR) to examine the biomass characteristics. As shown in Fig. 3, lignin syringyl (S) and guaiacyl (G) subunits, *p*-hydroxybenzoate

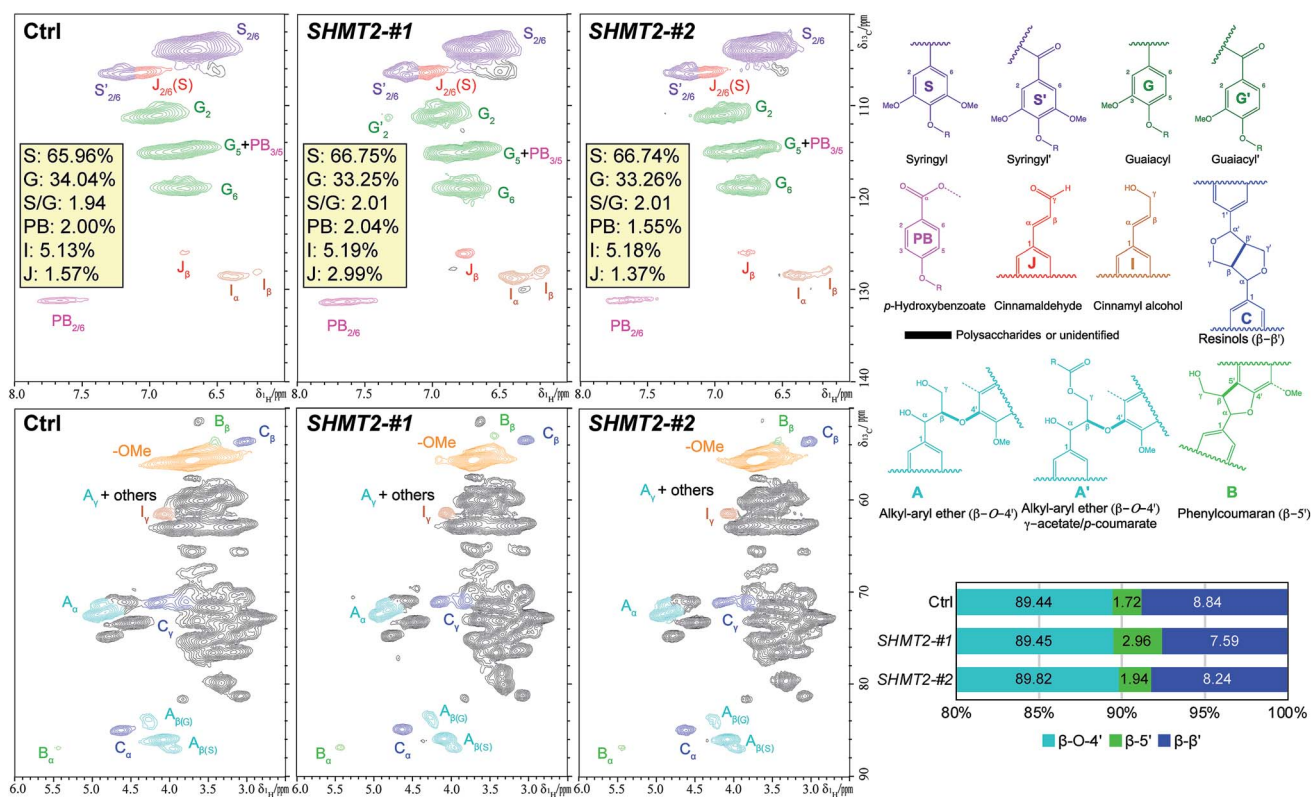


Fig. 3 Structural characterization of lignin by 2D NMR. 2D HSQC spectra are shown for the aromatic regions (top) and the aliphatic regions (bottom) of the whole cell walls from stems of two *PtSHMT2* overexpression lines and control poplars. The colors of the substructures shown match those of the corresponding cross peaks in the 2D HSQC spectra.



(PB), and dominant lignin inter-unit linkages β -aryl ether (β -O-4'), phenylcoumaran (β -5'), and resinols (β - β') were identified in the aromatic and aliphatic regions of *PtSHMT2* overexpression and control lines. The lignin S/G ratio is viewed as one of the factors affecting recalcitrance.^{25,26} The HSQC analysis revealed that the two *PdSHMT2* overexpression lines had a similar lignin S/G ratio to the control plants (Fig. 3), suggesting that the higher sugar release observed in the transgenic lines without pretreatment in this study was not due to lignin S/

G ratios (Fig. 2). In the two *PdSHMT2* overexpression lines, the β -5' content was relative higher but the β - β' content was relative lower than those of the control plants (Fig. 3). These small variations revealed from lignin 2D NMR spectra suggested that the lignin structures in the transgenic lines are similar to those in the control plant and that overexpression of *PdSHMT2* in *Populus* had little effect on the lignin compositional units and side-chain linkages. Sugar release has been related to both the lignin content and lignin S/G ratio as well as cell wall structure-

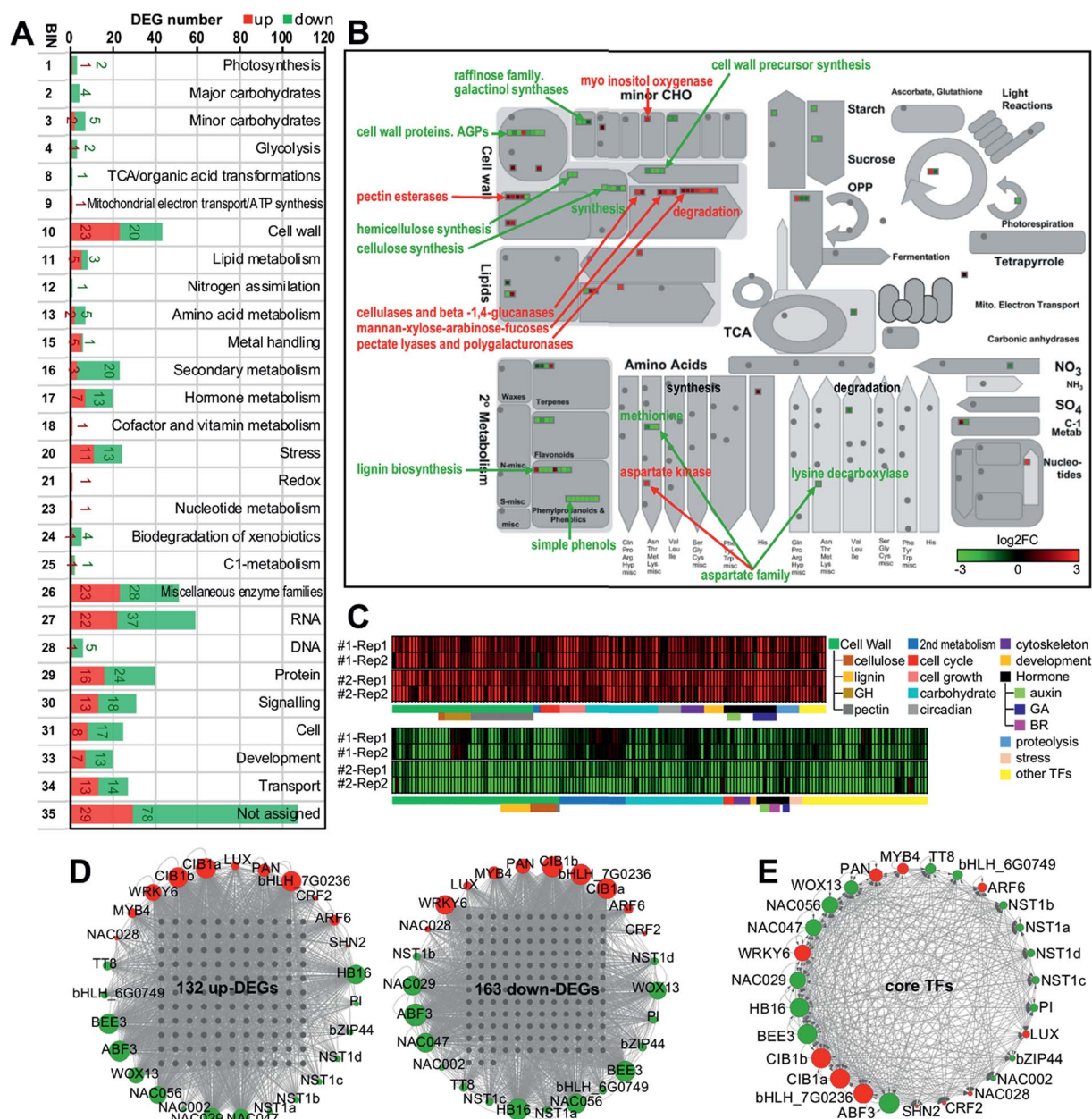


Fig. 4 Transcriptomic changes in the *PdSHMT2* overexpression transgenic plants. (A) Functional classification of DEGs in two *PdSHMT2* overexpression lines using MapMan Bin. (B) Involvement of DEGs in the metabolism processes. (C) Expression patterns of DEGs involved in cell wall, secondary metabolism and other growth-related processes in the two *PdSHMT2* transgenic lines. A total of 132 up-DEGs and 163 down-DEGs were identified. Red and green colors indicate log2 transformed fold changes (transgenic/control) in each replicate. (D) Regulatory network of the 27 hub TFs with 132 up-regulated (left) and 163 down-regulated core-DEGs (right). (E) The regulatory relationships among the 27 core TFs. A larger node indicates a larger number of genes regulated by a TF. Red and green nodes indicate up- or down-regulated TFs, respectively.



related factors.²⁷ Collectively, the reduced lignin content (and increased glucose content) rather than lignin structural variations likely contributed to the enhanced sugar release observed in the *PdSHMT2* transgenics plants.

Overexpression of *PdSHMT2* leads to changes in the transcriptome

To further reveal the molecular mechanism of *SHMT2* involvement in poplar cell wall modification, we performed RNA-Seq using mature leaves of two *PdSHMT2* overexpression lines and control plants. Compared to control plants, a total of 483 (230 up and 253 down) and 1159 (467 up and 692 down) differentially expressed genes (DEGs) were identified in line #1 and line #2, respectively (Table S2 in the ESI†). To explore the potential function of DEGs affected by *SHMT2*, MapMan analysis was conducted to classify these DEGs combined from the two transgenic lines (Fig. S9 in the ESI†). According to the MapMan classification system, a total of 23 up- and 20 down-DEGs belong to cell wall-related processes, 3 up- and 20 down-DEGs belong to secondary metabolism, 7 up- and 13 down-DEGs are involved in development, and 13 up- and 18 down-DEGs play roles in signaling (Fig. 4A). In the metabolic pathway, cell wall component biosynthetic genes (cellulose, hemicellulose and lignin biosynthesis) were down-regulated, whereas degradation-related genes (cellulose and beta-1,4-glucanases, mannan-xylose-arabinose-fucoses, pectate lyases and polygalacturonases) were up-regulated. In addition, cell wall precursor synthesis-related genes, lignin biosynthesis-related genes, simple phenol-related genes, and raffinose-related genes were down-regulated (Fig. 4B). We further classified the genes on the basis of their description and their *Arabidopsis* orthologs' functional description. In total, 132 up- and 163 down-DEGs related to the cell wall, secondary metabolism, cell cycle or cell growth, development, hormone, proteolysis, TFs, etc. were selected as "core-DEGs" for further analysis (Fig. 4C and Table S3 in the ESI†). In the two

transgenic lines, eight genes encoding glycosyl hydrolases (GHs) were significantly up-regulated, whereas nine lignin biosynthesis related genes (one *C3H*, two *C4H*, two *F5H*, one *MAX1*, one *CCoAMT* and two *CCoAOMT*) and eight cellulose biosynthesis related genes (*CesA4*, *CesA7*, *CesA8*, *CSLG2* and four *RIC4* homologs) were down-regulated. Moreover, six cell cycle related genes (*CYCA2;1*, two *CYCD3;1*, *CYCD3;2*, *CYCD6;1* and *CNGC15*) and eight cell growth related genes (*DWARF1*, *GRF7*, *PRX33* and five *EXPA* members) were up-regulated in the *PdSHMT2* overexpression lines (Fig. 4C). GHs comprise a large family of enzymes with a broad range of structures and substrate specificities, are ubiquitous in plants and play essential roles in various biological processes.²⁸ GHs are known to be involved in the degradation of biomass such as cellulose, hemicellulose, and starch.²⁹ For instance, KOR1, a GH9 family endo-1,4- β -glucanase, can degrade β -glucan with an unbranched β -(1 \rightarrow 4)-linked backbone which is the basic structure of cellulose.³⁰ The reduced cell wall recalcitrance in *PdSHMT2* overexpression lines might be partially explained by the induced GH expressions. The decreased lignin content and accelerated growth rates in *PdSHMT2* overexpression lines are consistent with the down-regulated lignin biosynthesis genes and up-regulated cell cycle related genes, respectively.

To seek potential hub TFs which play key roles in controlling gene expression in the *PdSHMT2* overexpression lines, we analyzed the TF binding sites (TFBSs) in the promoter regions of core-DEGs. Based on the TFBS enrichment in the promoter of core-DEGs, a total of 27 TFs were identified as the hub TFs including four *NST1* homologs (*NST1a*, *NST1b*, *NST1c*, and *NST1d*), two *CIB1* homologs (*CIB1a* and *CIB1b*), one *MYB4*, and one *TT8* (Fig. 4D and E).

To validate the RNA-Seq results, we selected eight hub TFs (*MYB4*, *CIB1b*, *WRKY6*, *NST1a*, *NST1b*, *NST1c*, *NST1d* and *TT8*) and seven TFs (*MYB46*, *MYB83*, *MYB63*, *MYB7*, *NAC056*, *GT2* and *HBI1*) from the DEG list for expression analysis using qRT-PCR.

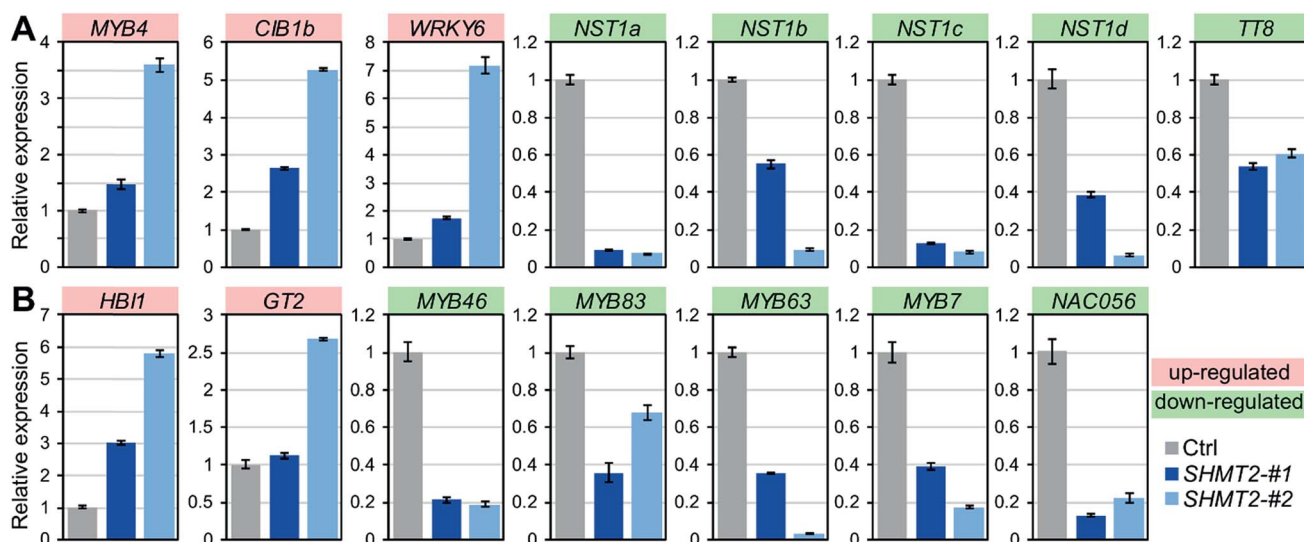


Fig. 5 qRT-PCR validation of expression of hub TFs (A) and other known cell wall-related TFs (B). Shown are mean values of three biological replicates \pm S.E.



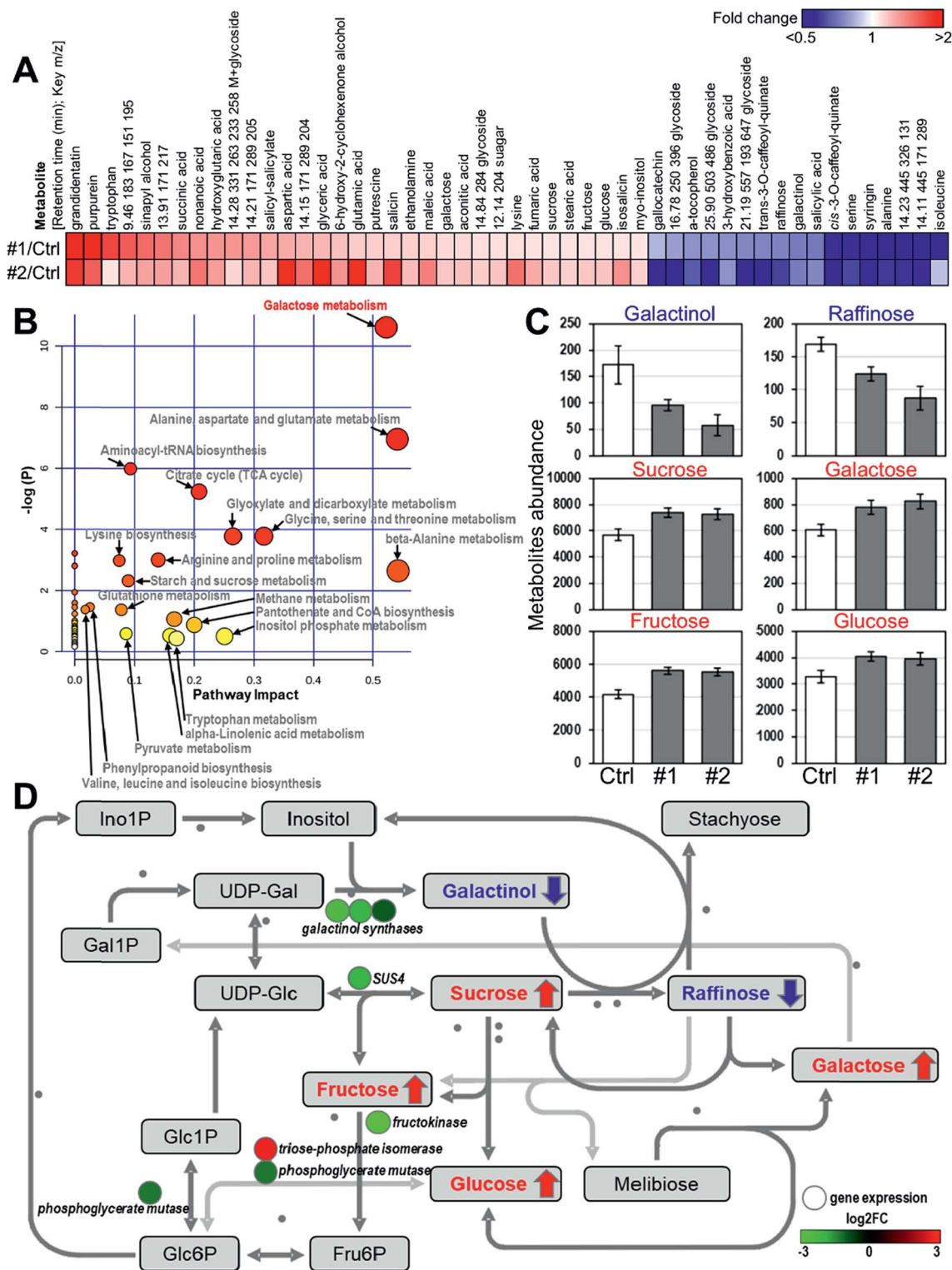


Fig. 6 Effects of *PtSHMT2* overexpression on the poplar metabolome. (A) Metabolite profile of two *PtSHMT2* overexpression lines (#1 and #2) compared to control plants. (B) Pathway analysis of differentially accumulated metabolites between *PtSHMT2* overexpression lines and control poplars. Global metabolic alterations of the most relevant pathways induced by *PtSHMT2* overexpression were revealed using MetaboAnalyst; a small *p* value and large pathway impact factor indicate that the pathway is greatly influenced. (C) Metabolite abundance. In the most affected galactose metabolism pathway, two metabolites (galactinol and raffinose) were inhibited while four metabolites (sucrose, galactose, fructose, and glucose) were induced. (D) Integrated analysis of the transcriptomic and metabolomic data in the galactose metabolism pathway. Red and green circles indicate DEGs identified from the transcriptomics analysis. Red or blue arrows indicate the differential metabolites identified from the metabolomics analysis. The map of galactose metabolism was downloaded from MapMan.



The diagram illustrates a regulatory network for cellulose, xylan, and lignin biosynthesis. At the top, a box contains six transcription factors: VND6, VND7, NST3, SND2, NST1, and NST2. A dashed line connects this box to a yellow oval labeled 'PtSHMT2'. Below the box, a central green oval labeled 'MYB83' is connected to several other transcription factors: MYB46, MYB52, MYB54, MYB32, MYB7, MYB4, MYB75, and MYB58. A red hexagon labeled 'glucose' is also connected to MYB83 and MYB58. The network then branches into three main pathways: Cellulose (regulated by SND3, MYB103, SND2, MYB20, MYB69, F5H, MYB52, MYB54, MYB43, and MYB7), Xylan (regulated by SNBE and SMRE), and Lignin (regulated by MYB4, MYB75, SMRE, and AC). The final products are Cellulose, Xylan, and Lignin, shown in blue boxes at the bottom.

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Phytozome (<https://phytozome.jgi.doe.gov>). In addition, the SHMT proteins were searched in 11 other plant species including three woody species *Salix purpurea*, *Eucalyptus grandis* and *Vitis vinifera*, three dicots *Arabidopsis thaliana*, *Medicago truncatula* and *Glycine max*; four monocots *Oryza sativa*, *Zea mays*, *Brachypodium distachyon* and *Sorghum bicolor*, and a moss *Physcomitrella patens* for phylogenetic analysis. The SHMT candidates were searched in the Pfam database to validate the presence of the SHMT motif (PF00464). Multiple sequence alignment of the full-length amino acid sequences was performed using Clustal X2.³⁹ The phylogenetic trees were constructed using the neighbor-joining method in the MEGA package V5.1⁴⁰ with bootstrap values from 1000 replicates indicated at each node. Three-dimensional structure prediction was performed by means of the I-TASSER (iterative threading assembly refinement, v4.4) toolkit.⁴¹

Normalized expression values of the *PtSHMT* genes from various tissues and organs were obtained from the *Populus* Gene Atlas database (<https://phytozome.jgi.doe.gov/phytozome/aspect.do?name=Expression>). The nm-scale expression of *PtSHMTs* was obtained from the AspWood gene expression database.²¹ The co-expression network of *PtSHMT2* was created using data obtained from the co-expressed biological process database for poplar.⁴² Cytoscape⁴³ was used to visualize the resulting network.

Generation of transgenic poplar

The full-length open-reading frame of *PtSHMT2* was amplified from *Populus deltoides* genotype WV94 and was cloned into the pAGW560 binary vector, in which expression was driven by the *Arabidopsis UBIQUITIN3* promoter. *Agrobacterium*-mediated transformation into WV94 was conducted at ArborGen Inc. (Ridgeville, SC) as described previously.⁴⁴ These plants were transferred and grown in greenhouses at Oak Ridge National Laboratory (Oak Ridge, TN) at a constant temperature of 25 °C and 16/8 h light/dark photoperiod. To estimate stem cylinder volume, plant height and stem base diameter of six-month-old plants were measured. We measured the primary stem length from the stem base (6 cm above the soil surface) to the shoot apex for plant height and measured the diameter of the stem base using calipers.

Chemical composition analysis

The size of poplar samples was reduced to 40 mesh using a Wiley mill (Thomas Scientific, Swedesboro, NJ) and Soxhlet-extracted with ethanol/toluene (1 : 2, v/v) for 24 h. The extractives-free sample was analyzed by a method consisting of two-step sulfuric acid (H₂SO₄) hydrolysis according to published literature.⁴⁵ In the first step, the extractives-free sample was hydrolyzed with 72% (w/w) H₂SO₄ at 30 °C for 1 h. In the second step, the hydrolyzed sample was diluted to a final concentration of 4% H₂SO₄ (w/w), followed by autoclaving at 121 °C for 1 h. The hydrolysate was filtered from the solid residue. The filtered liquid fraction was examined using a Dionex ICS-3000 ion chromatography system (Thermo Fisher Scientific, Sunnyvale, CA) for quantifying the sugar content. The total lignin content

was quantified *via* acid-soluble and acid-insoluble lignin separation from the hydrolysate and solid residue, respectively. Acid-soluble lignin was measured with the liquid fraction at 240 nm wavelength using UV/Vis spectroscopy. Acid insoluble lignin was quantified with the filtered solid residue as described in the NREL procedure. All analyses were technically duplicated for reported values and statistical analysis.

Sugar release test through enzymatic hydrolysis

The released glucose and xylose were measured according to published literature.⁴⁶ Dried and Wiley-milled (40 mesh) stems of the *Populus* control and transgenic plants were used for sugar release measurement. The dried sample (250 mg) was loaded into 50 mM citrate buffer solution (pH 4.8) with Novozymes CTec2 (gift product with 70 mg protein per gram of biomass from Novozymes, Franklinton, NC). The enzymatic hydrolysis was carried out at 50 °C and 200 rpm in an incubator shaker. Liquid hydrolysate was periodically collected at 0, 3, 6, 12, 24, 48, and 72 h, and enzymes in the hydrolysate were deactivated in boiling water for 10 min before carbohydrate analysis. The released sugars in each hydrolysate were measured using a Dionex ICS-3000 ion chromatography system. Each analysis was conducted in duplicate.

Heteronuclear single quantum coherence (HSQC) NMR analysis

NMR spectra were recorded at 30 °C on a Bruker Avance III HD 500 spectrometer and spectral processing was carried out using Bruker Topspin 3.5 (Mac) software. The whole cell NMR samples were prepared according to the literature.⁴⁷ Poplar stem powders (40 meshes) were extracted with an ethanol/toluene mixture (1 : 2, v/v) for 24 h. The extractives-free samples (air-dried) were loaded into a 50 mL ZrO₂ grinding jar (including 10 × 10 ball bearings) in a Retsch Ball Mill PM 100. The biomass was then ball milled at 600 rpm with a frequency of 5 min with 5 min pauses in-between for a total of 2 h. The ball-milled whole cell wall sample (~60 mg) and an NMR solvent mixture of 4/1 (v/v) DMSO-*d*₆/HMPA-*d*₁₈ (~0.4 mL) were loaded in a 5 mm NMR tube. The biomass and NMR solvents were mixed by vortexing for forming uniform gel-state samples, and then placed in a sonicator for 1–2 h before the analysis. The 2D heteronuclear single quantum coherence (HSQC) NMR experiments were carried out with a standard Bruker pulse sequence (hsqcetgpspsi2.2) with the following acquisition parameters: a spectral width of 12 ppm in the F2 (¹H) dimension with 1024 data points (acquisition time 85.2 ms) and 166 ppm in the F1 (¹³C) dimension with 256 increments (acquisition time 6.1 ms), a 1.0 s delay, a ¹J_{C-H} of 145 Hz, and 128 scans. The relative abundance of lignin compositional subunits and interunit linkage were estimated using volume integration of contours in HSQC spectra.^{47,48} For monolignol measurements of S, G, H, and *p*-hydroxybenzoate (PB), the S_{2/6}, G₂, H_{2/6}, and PB_{2/6} contours were used for relative quantitation. The contours of C_α signals were integrated for interunit linkage estimation.



RNA sequencing and data analysis

For RNA-Seq, mature leaves from *PtSHMT2* overexpression and control poplars were collected for RNA extraction. For each line, two biological replicates were used for library construction and sequencing. A total of 4 µg of total RNA was used to generate a cDNA library for Illumina sequencing. The RNA-Seq library was generated by a standard method as described in the Illumina sequencing sample preparation protocol (San Diego, CA, USA). After filtering out low-quality reads, RNA-Seq reads were aligned to the *P. trichocarpa* genome (version 3.0, https://phytozome.jgi.doe.gov/pz/portal.html#!info?alias=Org_Ptrichocarpa) using TopHat2.⁴⁹ Differentially expressed genes (DEGs) were identified using the R package DESeq2.⁵⁰ Raw *P*-values were adjusted for multiple comparison effects using the *q*-value (false discovery rate) method.⁵¹ The cutoff for significant DEGs was set as >2 absolute fold change (FC) and *q*-value < 0.05. MapMan⁵² and Gene Ontology (GO) were used for functional classification of DEGs. GO enrichment analysis was performed using agriGO.⁵³ For transcription factor binding site (TFBS) prediction, a 2 kb sequence upstream of the translation start site of identified DEGs was analyzed using PlantPAN.⁵⁴

Metabolomics analysis

For metabolite profiling, 25 mg of leaf tissues lyophilized and ground with a Wiley mill were twice extracted from each transgenic line and controls with 2.5 mL 80% ethanol overnight and then the extracts combined prior to drying a 0.50 mL aliquot in a nitrogen stream. As an internal standard, 75 µL of sorbitol at 1.0 mg mL⁻¹ was added to the first extract. Dried extracts were dissolved in acetonitrile, followed by TMS derivatization and analyzed by GC-MS according to Li *et al.*⁵⁵ Metabolite peaks were extracted using a characteristic mass-to-charge (*m/z*) ratio and quantified by area integration, and the concentrations were normalized to the quantity of the internal standard (sorbitol) recovered and the amount of sample extracted, derivatized and injected. A large user-defined database of mass spectral electron impact ionization fragmentation patterns of TMS-derivatized compounds (~2300 signatures) was used to identify the metabolites of interest. Unidentified metabolites were represented by their retention time and key *m/z* ratios. The metabolite data were presented as fold changes of the transgenic line *vs.* the average of the control lines from three biological replicates. Student's *t*-tests were used to determine whether differences were statistically significant (*P* ≤ 0.05).

Metabolite peaks were assigned using MS data, MS^E fragments, molecular weights and elemental compositions or interpreted using available biochemical databases, such as HMDB (<https://www.hmdb.ca/>), KEGG (<https://www.kegg.com/>) and Chempidder (<https://www.chemspider.com>). Fold changes from each group/control group or each group/CRF group and ROC curves were recorded using Metaboanalysis 3.0.

qRT-PCR analysis

One µg of total RNA was used to generate cDNA by using the Rite aid reverse transcriptase following the manufacturer's

instructions (Thermo Fisher Scientific, Hudson, NH). Gene-specific primers were designed using Primer3 software (<http://frodo.wi.mit.edu/primer3/input.htm>) with an annealing temperature of 58–60 °C and an amplicon of 150–250 bp. qRT-PCR was performed using a Maxima SYBR Green/ROX qPCR master mix (Thermo Fisher Scientific) according to the manufacturer's instructions. The relative gene expression was calculated by the 2^{-ΔΔC_t} method⁵⁶ using *PtUBQ10b* as an internal control. All experiments were performed by using three biological replicates and three technical replicates. The primers used in this study are listed in Table S4 in the ESI.†

Statistical analysis

Statistical analysis to determine statistical significance was performed by Student's *t* tests of paired samples. The asterisk in each figure indicates a significant difference compared to control samples (*P* < 0.05).

Authors' contributions

JZ performed experiments, analyzed data and wrote the manuscript. ML, CGY, YP and AJR performed chemical compositional analysis, lignin NMR analysis, and sugar release determination. WR, KAW and CMC generated *Populus* transgenic lines. VS, EAL, KB and JS generated and analyzed RNA-Seq data. ACB, SSJ and LEG measured biomass production. NLE and TJT generated and analyzed metabolomics data. XY designed the construct for *Populus* transformation. GAT, WM and JGC conceived the study, coordinated research and contributed to experimental design and data interpretation. All authors read and approved the final manuscript.

Conflicts of interest

There is no conflict of interest.

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