Identification and Characterization of a Golgi-Localized UDP-Xylose Transporter Family from Arabidopsis

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INTRODUCTION

Plant cell walls are composed of various polysaccharides and with the exception of cellulose and callose, these cell wall polysaccharides are biosynthesized in the lumen of the Golgi apparatus by families of glycosyltransferases (Scheible and Pauly, 2004; Liepman et al., 2010). The nucleotide sugar substrates essential for the biosynthesis of these polysaccharides are predominantly made in the cytosol. To overcome the subcellular partitioning of substrates and enzymes, nucleotide sugar transporters (NSTs) have evolved to allow the transport of nucleotide sugars from the cytosol into the Golgi and endoplasmic reticulum (ER) lumen. NSTs belong to the NST/triose phosphate transporter (TPT) superfamily, and the fact that they are present in all eukaryotes testifies to their biological significance (Knapp et al., 2003). Phylogenetic analyses have identified more than 50 members in Arabidopsis thaliana that are distributed in six clades (Rautengarten et al., 2014). However, functional characterization of members of the NST family at the molecular level has progressed slowly. In the past decade, only a few NSTs have been characterized, thus far accounting for the transport of GDP-mannose (GDP-Man), UDP-galactose (UDP-Gal), UDP-glucose (UDP-Glc), and CMP-sialic acid, although sialic acid has not been found in plants (Baldwin et al., 2001; Norambuena et al., 2002, 2005; Handford et al., 2004, 2012; Bakker et al., 2005, 2008; Rollwitz et al., 2006; Zhang et al., 2011; Mortimer et al., 2013). Recently, we developed a biochemical approach that allows the rapid and reliable determination of NST activities and led to the identification and characterization of the Arabidopsis bifunctional UDP-rhamnose (UDP-Rha)/UDP-Gal transporter (URGT1) clade (Rautengarten et al., 2014).

Xyl is a key component of various plant cell wall polymers, including xylan and xyloglucan, which are two of the most abundant cell wall polysaccharides in plants (Ebringerová and Heinze, 2000; Scheller and Ulvskov, 2010). While glucuronoxylan is a major hemicellulose in secondary cell walls, xyloglucan is the major component of the hemicellulosic fraction of primary walls of dicot plants. Minor amounts of Xyl can also be found in pectic polysaccharides, such as rhamnogalacturonan-II and xylogalacturonan (Jensen et al., 2008; Atmohadi et al., 2013), glycoproteins (Strasser et al., 2000), and diverse metabolites. Xylans in vascular plants are mainly composed of a backbone of \( \beta-(1,4) \)-linked xylyranosyl residues, which may be decorated at O-2 or O-3 with arabinofuranosyl residues or at O-2 with glucuronosyl and O-4-methylglucuronosyl residues to form arabinoxylan found in grasses and glucuronoxylan, the main xylan found in dicots (Tan et al., 2013; Rennie and Scheller, 2014). UDP-Xyl, the activated sugar donor for xylosyltransferases, is biosynthesized via decarboxylation of UDP-glucuronic acid by UDP-XYLOSE SYNTHASE (UXS) (Harper and Bar-Peled, 2002). While most nucleotide sugars are made in the cytosol and require transport into the Golgi lumen, in plants, members of the UXS family have been localized to both the Golgi and cytosolic fractions (Harper and Bar-Peled,
The previously classified NST-KT subfamily is characterized by a highly conserved lysine/threonine (KT) motif (Knappe et al., 2003) and forms clade I of the NST/TPT family (Rautengarten et al., 2014) encompassing 11 putative NSTs (Figure 1). Collectively, they share 25 to 93% identity in their amino acid sequences and include the recently identified URGTs that form subclades (A) and (B) of the NST-KT subfamily (Rautengarten et al., 2014). Another subclade (C) contains three additional members, namely, UXT1 (AT2G28315), UXT2 (AT2G30460), and UXT3 (AT1G06890). UXT1 shares 66% and 67% identity with UXT2 and UXT3, respectively. UXT2 and UXT3 share 92% identity in their amino acid sequences. Members of the uncharacterized subclade (D) share 25 to 28% identity in their amino acid sequences when compared with all other members of the NST-KT family (Rautengarten et al., 2014).

UXTs Are Ubiquitously Expressed and Localized to the Golgi Apparatus

Publicly available microarray expression data comprising the AtGenExpress Developmental Data Set (Schmid et al., 2005) have shown ubiquitous expression for UXT2 and UXT3 throughout plant development, with UXT3 showing highest expression in pollen and flowers. Since UXT1 is not present on the Affymetrix ATH1 array, we assessed the relative expression levels using quantitative RT-PCR (Figure 2A). Expression data obtained by quantitative RT-PCR for UXT2 and UXT3 are consistent with the microarray expression data, confirming ubiquitous but relatively low expression for both genes. UXT1 is more highly expressed in most tissues analyzed, with some variation in expression, especially in the stem tissue. To determine the subcellular localization of the UXTs, we generated C-terminal yellow fluorescent protein (YFP) fusions of the coding sequences and expressed them transiently in Nicotiana benthamiana leaves. All three UXTs localized to Golgi-like punctate structures and colocalized with the Golgi-marker α-mannosidase I, supporting their function as Golgi NSTs (Figure 2B). In contrast to UXT2 and UXT3, UXT1 also appeared to be localized to the ER and colocalized with the ER marker ER-ck (Nelson et al., 2007) (Supplemental Figure 1).

Determining the In Vitro Functions of the Arabidopsis UXTs

To assess the function of the UXTs, each was heterologously expressed in Saccharomyces cerevisiae (yeast) and microsomal proteins were prepared. Immunoblot analysis confirmed the presence of the specific UXT proteins in yeast microsomal extracts (Figure 3A). Subsequently, microsomal proteins were reconstituted into liposomes for transport assays. Proteoliposomes preloaded with either UMP, GMP, CMP, or AMP were incubated with a mixture of 16 nucleotides/nucleotide sugars (Figure 3B). Nontransported substrates were removed by gel filtration, and the content of the liposomes was analyzed by liquid chromatography-tandem mass spectrometry (LC-MS/MS). The LC-MS/MS analysis of nucleotide sugars after transport by UXTs could be readily assessed when compared with the empty vector control (Figures 3C and 3D). All three UXTs had the capacity to transport UDP-Xyl as well as minor amounts of UDP-arabinopyranose (UDP-Arap) in vitro when the proteoliposomes were preloaded with UMP (Figure 3E). By contrast, when proteoliposomes were preloaded with GMP, only transport of GDP-sugars was observed (Figure 3F), resulting from endogenous activity present in yeast microsomal preparations, since the incorporation levels were similar to those observed in control reactions (yeast transformed with the empty vector). No significant transport activities were observed when proteoliposomes were preloaded with AMP or CMP.

The UXT-mediated transport of UDP-Xyl was saturable in a concentration and time-dependent manner (Figures 4A and 4B). To determine \( K_{cat} \), we measured the amount of UXT protein in the proteoliposomes using multiple reaction monitoring (MRM) mass spectrometry as explained in Methods and Supplemental Table 1. The analysis of the UXTs revealed apparent \( K_m \) values for UDP-Xyl in the range of 40 to 60 \( \mu \)M with turnover rates of 3 to 12 s\(^{-1}\) (Table 1). As previously determined, the UDP-Xyl content in various Arabidopsis organs is in the range of 40 to 120 pmol mg\(^{-1}\) dry weight (Rautengarten et al., 2014). Considering the volume of the central vacuole in plants, these measurements indicate that the cellular levels of UDP-Xyl are in the micromolar range. Thus, we estimate that the affinity constants (\( K_m \)) for all three UXTs are within physiological range. By contrast, estimations of the \( K_m \) for

![Figure 1. Phylogenetic Tree of the Arabidopsis NST-KT Subfamily.](image-url)
UDP-Ara, which was transported by all three UXTs to a lower extent, revealed values of ≥200 μM, which would be inconsistent with endogenous concentrations. UDP-Ara concentrations in Arabidopsis organs are very similar to UDP-Xyl concentrations (Rautengarten et al., 2014); hence, the high affinity constants or $K_m$ values indicate that the UXTs are most likely not significantly involved in UDP-Ara transport in vivo. The temperature optimums for transport of UDP-Xyl for the three UXTs ranged from 37 to 55°C (Figure 4C).

### The Role of the UXTs in Planta

To evaluate the in vivo function of the UXTs, we obtained homozygous T-DNA lines. Two independent lines were acquired for UXT1, and single insertion lines were identified for UXT2 and UXT3. Finally, a double knockout was generated between uxt2 and uxt3, and the absence of respective full-length transcripts was confirmed by PCR (Supplemental Figure 2).

To assess if mutations in the UXTs affect the biosynthesis of specific polysaccharides, we prepared alcohol insoluble residue (AIR) from leaves, flowers, and young (upper) and mature (lower) inflorescence stem tissue from 6- to 8-week-old plants and analyzed the monosaccharide composition (Supplemental Table 2). Flowers and leaves showed no significant differences in the monosaccharide composition between any of the mutants and Columbia-0 (Col-0; P > 0.05, ANOVA and Duncan’s test for multiple comparisons). Stem data showed a significant difference in the monosaccharide composition of uxt1-1 and uxt1-2 compared with Col-0 (P < 0.05), whereas the other mutants did not show a difference from Col-0. In uxt1-1 and uxt1-2, only Xyl and glucuronic acid (GlcA) were significantly decreased in stems compared with Col-0 (Figures 5A and 5B). The Xyl content in mature inflorescence stems from the uxt1 mutants was decreased by 16% (Figure 5B), whereas a reduction of 34% was observed in young stem tissue (Figure 5A). These data confirm the importance of UXT1 for the biosynthesis of Xyl-containing cell wall polymers. In addition, a significant reduction in cell wall GlcA content of −25 to 37% was observed in mature and young parts of the inflorescence stems from uxt1 mutants. Since the monosaccharide compositions are relative measurements, the decrease in Xyl and GlcA was accompanied by an apparent increase in other monosaccharides. However, the ratio between the other sugars was not significantly changed in any of the samples, indicating that a loss of function of UXT1 had no direct effect on any sugars besides Xyl and GlcA. Notably, while GlcA content was decreased in the mutant, there was no change in the 4-O-methyl ether (MeGlcA) content, i.e., the ratio between the methylated and nonmethylated form of GlcA was much higher in the mutant than in the wild type (Supplemental Figure 3). However, even though there was a significant reduction of Xyl and GlcA in the uxt1 mutants, the plants did not exhibit a morphological phenotype compared with wild-type plants.

### Cell Wall Profiling of the uxt1 Mutants

To analyze the changes in the cell wall composition of uxt1 mutants in more detail, we performed comprehensive microarray polymer profiling (CoMPP) analysis on mature (lower) stem material. Cell wall matrix polymers were extracted, spotted onto nitrocellulose membranes to generate microarrays, and probed with a number of different antibodies with specificity for epitopes borne on cell wall polymers (Figure 5C). Since we observed a significant decrease in cell wall xylene in the uxt1 mutants, we focused specifically on antibodies recognizing xylan structures, including LM10/LM11 (which bind unsubstituted xyloses), UX1 (which recognizes GlcA or MeGlcA substitutions on xylan), and AX1 [which was produced against arabinose-substituted β-(1,4)-xylan from wheat (*Triticum aestivum*)]. Clear differences were observed between the uxt1-1 and uxt1-2 mutants and the wild type (Figure 5C). Collectively, these probes revealed an apparent
In Arabidopsis, xylan is comprised of domains that differ in their pattern of MeGlcA substitution and is generated by two glucuronoxylanases, GUX1 and GUX2. The GUX1 enzyme is responsible for the addition of GlcA only to evenly spaced Xyl residues, whereas GUX2 decorates both even and odd spaced Xyl residues (Bromley et al., 2013). To determine if mutations in UXT1 preferentially affect one of the domains, we digested xylan from the uxt1 mutants and the wild type with the glucuronoxylanase C (XynC) and analyzed the released oligosaccharides by high-performance anion exchange chromatography (HPAEC; Supplemental Figure 4). The xylooligosaccharide profile from the uxt1 mutants is similar to the profile observed for the wild type, and differences detected in the chromatograms are consistent with the reduction in the xylan content and, more specifically, as indicated in the results for the UX1 antibody, in the glucuronoxylan content in UXT1 mutants when compared with the wild type (Figure 5C). Concomitant with the reduction in xylan content, the results from the CoMPP analysis indicate that other polymers, such as cellulose (recognized by the carbohydrate bonding module CBM3a) and xyloglucan (recognized by antibody LM25), were only slightly affected in UXT1 mutants. By contrast, results obtained with the LM3 antibody, which recognizes glycan moieties on extensins, indicated that these proteoglycans are enriched in UXT1 mutants. In addition, oligosaccharide mass profiling (OLIMP) revealed no significant differences in the xyloglucan structure in uxt1 cell wall preparations compared with the wild type (Table 2).

**Figure 3.** LC-MS/MS Analysis of NST Activities of UXTs.

(A) Immunoblot analysis of UXT expression in yeast microsomal protein extracts (2.5 μg), including the empty vector control.
(B) Separation of a 20 nucleotide/nucleotide sugar mix: 1, CMP; 2, UMP; 3, UDP-GalA; 4, UDP-glucuronic acid; 5, CMP-sialic acid; 6, UDP-Arap; 7, UDP-Rha; 8, UDP-Gal; 9, UDP-Glc; 10, UDP-Xyl; 11, UDP-GlcNAc/GalNAc; 12, UDP-Araf; 13, adenosine 3′-phosphate 5′-phosphosulfate; 14, GMP; 15, AMP; 16, GDP-Man; 17, GDP-Gal; 18, GDP-Glc; 19, GDP-Fuc; and 20, ADP-Glc.
(C) and (D) Reconstitution of empty vector control (C) and UXT1 (D) into liposomes and analysis by LC-MS/MS after simultaneous incubation with 16 nucleotide sugar substrates.
(E) and (F) Quantification of nucleotide sugar uptake of proteoliposomes containing UXT1 and preloaded with UMP (E) and GMP (F). Data represent the mean and SD of n = 2 independent experiments.
with the increased ratio of the methylated and nonmethylated form of GlcA in the mutant compared with the wild type (Supplemental Figure 3). However, the pattern of xylan substitution is clearly different from those observed for the gux1 and gux2 mutants, thus indicating that UXT1 does not play a role in defining the domain pattern of decoration. Compared with the wild type, the profile from the uxt1 mutants showed a slight shift toward shorter oligosaccharides, indicating a higher degree of substitution (Supplemental Figure 3).

Since UXT1 localizes to both the Golgi apparatus and the ER, it is possible that UXT1 could also play a role in protein glycosylation. To investigate this in more detail, we analyzed total protein from UXT mutants with antibodies against plant N-glycan xylosyl and fucosyl epitopes. However, no obvious differences in the N-glycan xylosylation pattern of the UXT mutants compared with the wild type could be observed. The same was true for the fucosylation pattern, which we analyzed as a control (Supplemental Figure 3). Hence, our results indicate that the UXTs do not have an obvious effect on protein N-glycosylation.

DISCUSSION

The majority of nucleotide sugars are actively transported into the Golgi and ER lumen by NSTs (Bar-Peled and O’Neill, 2011). Here, we present evidence for the existence of a Golgi UDP-Xyl transporter family in plants. We identified three previously uncharacterized Arabidopsis NSTs comprising the NST-KT family subclade C. All three members are capable of transporting UDP-Xyl and to a lower extent UDP-Ara in vitro (Figure 4). However, estimations of the $K_m$ values for UDP-Ara indicate that these UXTs are unlikely to transport UDP-Ara in vivo, whereas determined $K_m$ values for UDP-Xyl transport are within the physiological range (Rautengarten et al., 2014).

Analysis of the subcellular localization of the UXTs showed that all three are located in the Golgi apparatus and that UXT1 also localized to the ER, which could indicate a distinct functional role for UXT1. However, we are not aware of any xylosyltransferase reaction that is known to take place in the ER. Although protein N-glycosylation is initiated in the ER, the xylosylation has been shown to occur in the Golgi apparatus (Fitchette-lainé et al., 1994; Egelund et al., 2006). Therefore, the possible biological significance of the partial localization of UXT1 to the ER remains unclear. By contrast, the localization to the Golgi apparatus of all three transporters is consistent with the biosynthesis of Xyl-containing glycans in this compartment. The three UXTs are expressed throughout plant development. Compared with UXT1, UXT2 and UXT3 have lower overall expression levels in the tissues analyzed. The UXT1 transcript varies more substantially, especially in mature stem material where relative expression was lowest but still detectable.

To determine the functions of the UXTs in planta, we identified and analyzed loss-of-function mutants in all three transporters. Neither the uxt2 nor the uxt3 mutants exhibited any changes in cell wall monosaccharide composition. This could be due to the lower relative levels of expression for these two genes when compared with UXT1. Similarly, the uxt2 uxt3 double mutants did not show any discernable phenotype, indicating possible minor roles for these genes. Due to the genetic linkage between UXT1 and UXT2, we have not been able to generate a triple homozygous mutant line.

Only UXT1 mutants showed a significant decrease in cell wall-derived Xyl. This decrease was exclusive to stem material and was more pronounced in material isolated from younger parts of inflorescence stems. This difference based on stem maturity can

Table 1. Kinetic Parameters of UDP-Xyl Transport into Proteoliposomes

<table>
<thead>
<tr>
<th>Parameter</th>
<th>UXT1</th>
<th>UXT2</th>
<th>UXT3</th>
</tr>
</thead>
<tbody>
<tr>
<td>$K_m$ (µM)</td>
<td>39 (3)</td>
<td>40 (4)</td>
<td>58 (9)</td>
</tr>
<tr>
<td>$V_{max}$ (nM s$^{-1}$)</td>
<td>16 (0)</td>
<td>13 (0)</td>
<td>4 (0)</td>
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<tr>
<td>$K_{cat}$ (s$^{-1}$)</td>
<td>3.6</td>
<td>10.3</td>
<td>11.7</td>
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</tbody>
</table>

For each UXT, 20 data points with varying substrate concentrations (0.5 to 400 µM) were acquired. Standard errors are in parentheses.

Figure 5. Cell Wall Analysis of UXT1 Mutants.

(A) and (B) Monosaccharide composition of total cell wall extracts from upper (A) and lower (B) parts of the inflorescence stem. Data represent 10 pooled individuals and mean and so from six technical replicates. Values are expressed in mol %.

(C) CoMPP analysis with relative abundance of cell wall glycan epitopes. Mean and so for spot signals (MSS) were obtained by probing microarrays with antibodies (r axis) from four technical replicates and the highest MSS set to 100 and all other values adjusted accordingly. For all data, values significantly different from the wild type are marked with asterisks (*P < 0.05 and **P < 0.01; t test).
be explained by the fact that in the developing stem, active cell wall biosynthesis and xylan production is occurring, whereas in mature stem tissue the function of the UDP-Xyl transporter may no longer be needed. Together with the decrease in Xyl in stems, we also observed a proportional reduction in GlcA content. Most of the GlcA in stem cell walls originates from glucuronoxylan; therefore, the proportional reduction in GlcA content is consistent with the suggestion that UXT1 functions predominantly in the biosynthesis of glucuronoxylans. However, the methylated GlcA content remained unchanged in uxt1 mutant plants when compared with the wild type (Supplemental Figure 3). This observation is consistent with previously published data on xylan mutants, such as inx8, inx9, and fra8, in which the ratio of GlcA to MeGlcA is lower and methylated GlcA predominates (Liepman et al., 2010).

A more detailed characterization of uxt1 mutants using CoMPP and OILMP techniques also indicated the importance of UXT1 in glucuronoxylan biosynthesis and revealed that it has little effect on xyloglucan biosynthesis. The latter could be explained by a lower requirement for UDP-Xyl for cell wall biosynthesis. However, our results show that, at least for xylan biosynthesis, the functional association with a transporter is a mechanism that allows the synthases to maintain a degree of processivity and operate efficiently in vivo.

In mammalian cells, UXS enzymes are located only in the Golgi lumen and UDP-Xyl transport is therefore apparently not required (Ashikov et al., 2005). However, a mutant in the UXS enzyme in Chinese hamster ovary cells could be complemented by a cytoplasmic isoform of UXS from Arabidopsis, showing that in these cells the route for delivery of UDP-Xyl is not important for the function of the xylosyltransferases (Bakker et al., 2009). Since UDP-Xyl in plants is biosynthesized both in the cytoplasm and in the Golgi lumen by UXS enzymes, it seemed highly likely that plant UDP-Xyl transport into the Golgi would be a redundant process. However, our results show that, at least for xylan biosynthesis, the transport of UDP-Xyl is important and the Golgi-localized UXS enzymes cannot deliver sufficient UDP-Xyl for proper xylan biosynthesis.

**Conclusions**

We identified three Golgi-localized nucleotide sugar transporters that are able to transport UDP-Xyl in vitro. This demonstrates the existence of NSTs with specificity for UDP-Xyl in plants. uxt1 mutant plants showed a significant decrease in total cell wall Xyl content in stems, thus confirming a role for UXT1 in providing UDP-Xyl for cell wall biosynthesis.

**METHODS**

**Nucleotide and Nucleotide Sugar Standards**

Nucleotide and nucleotide sugar standards were obtained from the following sources: UDP-α-D-xylose, UDP-β-L-arabinopyranose, and UDP-α-D-galacturonic acid (Carbohydrore, Athens, GA); UMP, GMP, CMP, AMP, UDP-α-D-glucuronic acid, UDP-α-D-glucose, UDP-α-D-galactose, UDP-N-acetyl-α-D-glucosamine, UDP-N-acetyl-α-D-galactosamine, UDP-α-D-mannose, UDP-β-L-arabinofuranose, GDP-a-D-galactose, and ADP-β-L-phosphate 5'-phosphosulfate, CMP-β-D-xyluronic acid, and ADP-α-D-glucose (Sigma-Aldrich); and UDP-β-L-arabinofuranose (Peptides International). UDP-α-D-galactose

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**Table 2. Xyloglucan Structure Determined by Oligosaccharide Mass Profiling**

<table>
<thead>
<tr>
<th>Tissue</th>
<th>Plant Line</th>
<th>GXGX</th>
<th>XXXG</th>
<th>XXLGLXLG</th>
<th>XNLGLXLG +OAc</th>
<th>XFXG</th>
<th>XFXG +OAc</th>
<th>XXLFB</th>
<th>XXLFB +OAc</th>
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<th>XLLG +OAc</th>
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<tbody>
<tr>
<td>Lower</td>
<td>Col-0</td>
<td>2.0±0.2</td>
<td>32.5±2.8</td>
<td>9.3±0.7</td>
<td>1.8±0.2</td>
<td>15.0±0.9</td>
<td>14.1±0.8</td>
<td>13.7±1.6</td>
<td>11.5±0.5</td>
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</tr>
<tr>
<td>Stem</td>
<td>uxt1-1</td>
<td>2.1±0.3</td>
<td>33.2±1.8</td>
<td>8.6±0.6</td>
<td>1.4±0.1</td>
<td>16.1±0.5</td>
<td>14.1±0.4</td>
<td>14.3±1.1</td>
<td>10.3±1.0</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Upper</td>
<td>Col-0</td>
<td>2.1±0.1</td>
<td>35.0±1.3</td>
<td>7.8±0.2</td>
<td>1.4±0.1</td>
<td>15.2±0.2</td>
<td>14.4±0.4</td>
<td>13.3±0.3</td>
<td>10.9±0.9</td>
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</tr>
<tr>
<td>Stem</td>
<td>uxt1-2</td>
<td>2.3±0.5</td>
<td>37.5±0.9</td>
<td>13.0±0.2</td>
<td>1.0±0.1</td>
<td>19.0±1.3</td>
<td>7.3±0.3</td>
<td>15.3±0.3</td>
<td>3.5±0.3</td>
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</table>

Relative abundance of xyloglucan oligosaccharides in the upper and lower stem walls was determined by OILMP. No significant differences (P < 0.05) were observed between the wild type and mutants. Data represent the mean (±sa) of n = 3 independent biological replicates. The mass-to-charge ratio (m/z) is provided for each oligosaccharide; for nomenclature of the oligosaccharides, see Fry et al. (1993).
was enzymatically synthesized according to Major et al. (2005) and HPLC purified using a linear ammonium formate gradient (Rautengarten et al., 2011). UDP-β-L-rhamnose was enzymatically synthesized by a two-step reaction using UDP-Glc as substrate as previously described (Rautengarten et al., 2014).

Sequence Analysis

Amino acid sequences were retrieved from The Arabidopsis Information Resource (Lamesh et al., 2012). Deduced amino acid sequences were aligned using the Clustal Omega program (Sievers et al., 2011) using default parameters (Supplemental Data Set 1). Phylogenetic trees were created using the neighbor-joining statistical method and applying the bootstrap method with 1000 replications and visualized using the Molecular Evolutionary Genetics Analysis (MEGA) version 6.0 application (Tamura et al., 2013).

Heterologous Expression, Reconstitution, and in Vitro Assay of Transport Activities

Heterologous expression in Saccharomyces cerevisiae (strain INVSc1: MATa his3D1 leu2 trp1-289 ura3-52 MAT his3D1 leu2 trp1-289 ura3-52; Life Technologies), reconstitution of microsomal proteins, and subsequent transport assay were performed as previously described (Rautengarten et al., 2014). Kinetic parameters were calculated by nonlinear regression using the Prism6 application (GraphPad Software). PAGE and immunoblot analyses were done as previously described (Rautengarten et al., 2011) using 2.5 μg yeast microsomal protein. Filters were probed using the anti-V5 antibody (Life Technologies).

Chromatographic Separation and Detection of Nucleotide Sugars by Mass Spectrometry

LC-MS/MS was performed using porous graphic carbon as the stationary phase on an 1100 series HPLC system (Agilent Technologies) and a 4000 QTRAP LC-MS/MS system (AB Sciex) equipped with a TurboIonSpray ion source using methods previously described (Ito et al., 2014; Rautengarten et al., 2014)

Absolute Quantification of Reconstituted NSTs by MRM Mass Spectrometry

The yeast expression vector pYES-DEST52 contains an in-frame V5-tag and 6xHis-tag epitope at the 3’ end of the cloning site. The expressed UXT proteins all yield a common tryptic peptide, namely, RSRGPFEKPG-PNPMLLDSR.T, as previously described (Rautengarten et al., 2014). A synthesized peptide was used to determine optimal parameters for MRM analysis with the following parameters: dwell (25 ms), fragmentor (130 V), collision energy (11.1 V), and cell accelerator voltage (7 V). Analysis of samples and standard curves were conducted on a 6460 Triple Quad LC MS system equipped with a Jet Stream ESI source (Agilent Technologies). The system was operated in positive ion mode using the MRM scan type with both MS1 and MS2 resolutions set to unit. The following mass spectrometer parameters were applied: gas temperature (350°C), gas flow (10 L/min), nebulizer (45 p.s.i.), sheath gas temperature (400°C), sheath gas flow (11 L/min), capillary (5000 V), and MS1/MS2 heater (100°C). A total of 5 μg of tryptic-digested (1:10 [w/w]) proteoliposome was loaded onto an Ascentis Express Peptide ES-C18 (5 cm × 2.1 mm, 2.7 μm) column (Sigma-Aldrich) using a 1290 series HPLC (Agilent Technologies) at a flow rate of 0.4 mL/min as follows: 95% Buffer A (99.9% water and 0.1% formic acid) and 5% Buffer B (99.9% acetonitrile and 0.1% formic acid) for 0.2 min, followed by an increase to 35% Buffer B over 5.5 min, then 90% Buffer B in 0.3 min, where it was held for 2 min. The buffer composition was ramped back to 5% Buffer B over 5 min, giving a total runtime of 13 min. The column temperature was maintained at 60°C. Data were acquired using MassHunter Workstation Software Version B.06.00 Build 6.0.6025.4 SP4 (Agilent Technologies). The raw data were imported into Skyline (v2.5.0.6157) (MacLean et al., 2010) and transition peaks manually inspected for retention time and adjusted accordingly. The abundance of the expressed UXTs in a sample was calculated by integrating the total signal peak area (total area) from Skyline for the two transitions on the predominant 563.560 [M+4H]++ precursor ion, namely, L [r7] 761.452 [M+H]+ and G [v6] 648.3311 [M+H]+, and calculating total moles in the sample against a standard curve for the synthesized peptide. The standard curve was created by linear regression using a range of abundances (0.5 to 10 pmol), which were interspersed as separate runs during sample analysis. The UXTs represent from 0.01 to 0.1% of total protein of the reconstituted proteoliposomes with errors representing the sd of two technical replicates (Supplemental Table 1). Values were used for enzyme kinetic calculations.

Plant Material and Growth Conditions

Arabidopsis thaliana Col-0 was obtained from the ABRC (http://abrc.osu.edu). T-DNA insertion mutants for UXT1 (uxt1-1, SALK_147_F11; uxt1-2, SALK_086773), UXT2 (uxt2-1, SALK_078576), and UXT3 (uxt3-1, SALK_013372) were obtained from the SiGnAL Salk collection (http://signal.salk.edu/). Plants were germinated and grown on soil (PRO-MIX; Premier Horticulture) in an Arabidopsis growth chamber (Percival-Scientific) under short-day light conditions (10 h of fluorescent light [120 μmol m−2 s−1] at 22°C and 60% RH/14 h of dark at 22°C and 60% RH). After 4 weeks, plants were transferred to long-day conditions (16 h of fluorescent light [120 μmol m−2 s−1] at 22°C and 60% RH/8 h of dark at 22°C and 80% RH).

Cloning Procedures

Coding sequences for Arabidopsis UXTs without native stop codon were PCR amplified using the primer pairs listed in Supplemental Table 3. PCR products were introduced into the pENTR/SD/D-TOPO cloning vector (Life Technologies) according to the manufacturer’s protocol and confirmed by sequencing. To obtain C-terminal YFP fusions, the constructs were introduced into the 35S promoter carrying pEarleyGate101 plant transformation vector (Earley et al., 2006) using the LR Clonase II reaction (Life Technologies) following the manufacturer’s protocol. For yeast expression, the constructs were introduced into the yeast expression vector pYEX-DEST52 (Life Technologies) using the LR Clonase II reaction (Life Technologies).

Subcellular Localization and Microscopy

Nicotiana benthamiana plants were grown on soil (PRO-MIX) in a growth chamber (Percival-Scientific) using the following conditions: 24°C day/night temperature, 60% humidity, and 16-h-light/8-h-dark cycles. Four-week-old leaves were co-infiltrated with Agrobacterium tumefaciens strain GV3101 pmp90 carrying the C-terminal YFP fusion constructs (OD600 = 0.15) and the α-mannosidase-mCherry marker (OD600 = 0.01) (Nelson et al., 2007) using the previously described method (Jensen et al., 2008). Visualization by confocal laser scanning microscopy was performed as previously described (Rautengarten et al., 2012).

Determination of Monosaccharide Composition

Air was prepared as described earlier (Harholt et al., 2006). Samples were hydrolyzed in 2 N trifluoroacetic acid for 1 h at 120°C. HPAEC with pulsed amperometric detection was performed as described (ØBro et al., 2004) on an ICS 3000 ( Dionex) using a CarboPac PA20 anion exchange column (3 × 150 mm; Dionex).
PCR Characterization of Mutants

Homozygous T-DNA insertion lines were verified by PCR to confirm the presence of the insert using the primers listed in Supplemental Table 3. Subsequently, absence of the transcript was verified by RT-PCR using the primers listed in Supplemental Table 3. Arabidopsis ACTIN-2 (At3g18780) was used as a control for equal loading.

RT-PCR

Plant RNA was extracted using the RNEasy RNA Plant Kit (Qiagen) according to the manufacturer’s protocol, and 0.5 to 1 μg was reverse transcribed using SuperScript II reverse transcriptase and d(T)15 oligomers (Life Technologies) according to the manufacturer’s protocol. UXT1-3 expression in different organs was analyzed by quantitative RT-PCR using SYBR Select Master Mix (Applied Biosystems) on a StepOnePlus Real-Time PCR system (Applied Biosystems) according to the conditions described earlier (Czechowski et al., 2005) using StepOne 2.0 software (Applied Biosystems). The UXT genes were amplified using the primers listed in Supplemental Table 3. As references, primers for UBQ10 (At4g05320), PP2A (At1g13320), and a SAND family member (MON1, At2g28390) were used (Supplemental Table 3). Expression levels were calculated using the comparative CT method, which involves normalizing against the geometric mean of the three housekeeping genes (UBI10, PP2A and SAND family) for each tissue type (Schmittgen and Livak, 2008).

Xylan Oligosaccharide Profiling

Xylan was digested with endoglucuronoxylanase GH30 (XynC) from Bacillus subtilis (St John et al., 2006) as previously described (Bromley et al., 2013). Profiling of the released oligosaccharides by HPAEC was performed using the conditions previously described (Chiniquy et al., 2012). Cell wall preparations from the gux1 and gux2 mutants (Okawa et al., 2010; Bromley et al., 2013) were analyzed for comparison.

Protein Extraction and Immunoblotting

Inflorescences stems from 6-week-old plants were ground in extraction buffer (10 mM Tris, pH 8, 150 mM NaCl, 2% Triton, 1 mM PMSF), protease inhibitor, and 10 mM CaCl₂ incubated for 1 h at 4°C under constant shaking, and centrifuged for 30 min at 20,800g at 4°C to remove cell debris. Subsequently, protein was precipitated with 20% trichloroacetic acid, incubated on ice, and spun down. After removal of the supernatant, the samples were washed twice with ice-cold acetone, dried, and suspended in the appropriate buffer. Samples were separated by SDS-PAGE, and glycosylation was detected by immunoblotting using antibodies raised against β-(1,2)-Xyl and α-(1,3)-Fuc (Agrisera). Detection was performed with an ECL Plus Western Blotting Detection System (GE Healthcare).

OLIMP

Mass profiling of xyloglucan oligosaccharides derived from various stem tissues was performed as previously described (Lerouxel et al., 1999), and the resulting solubilized xyloglucan oligosaccharides were detected using an Axima matrix-assisted laser desorption/ionization time-of-flight system (Shimadzu) set in linear positive mode with an acceleration voltage of 20,000 V.

CoMPP

CoMPP was undertaken essentially according to previously described work (Moller et al., 2007). AIR samples were extracted from mature stems of pooled material (approximately eight individuals) from 8-week-old uxt1-1, uxt1-2, and wild-type plants. A total of 4 mg AIR was subsequently extracted sequentially with CDTA and then NaOH solutions to obtain pectin-rich and hemicellulose-rich extracts, respectively. These extracts were spotted onto membranes and probed with monoclonal antibodies and carbohydrate binding modules (CBMs) that recognize specific cell wall epitopes, namely, LM3, extensins; LM25, xyloglucan; LM10, β-(1-4)-o-xylan; LM11, (1-4)-β-D-xylan; UX1, glucuronoxylan; AX1, arabinoiose substituted β-(1-4)-o-xylan; and CBM3a, cellulose (Guillon et al., 2004; McCarty et al., 2005; Blake et al., 2006; Koutaniemi et al., 2012; Pedersen et al., 2012).

Accession Numbers

Sequence data from this article can be found in the Arabidopsis Genome Initiative or GenBank/EMBL databases under the following accession numbers: UXT1 (AT2G28315), UXT2 (AT2G30460), UXT3 (AT1G06890), ACTIN-2 (At3g18780), UBQ10 (At4g05320), PP2A (At1g13320), and MON1 (At2g28390).

Supplemental Data

Supplemental Figure 1. Subcellular localization of UXT1 with an ER marker.

Supplemental Figure 2. Assessment of UXT transcripts by RT-PCR in the of uxt mutant backgrounds.

Supplemental Figure 3. The 4-O-Methyl-α-glucuronic acid content of pooled stem material.

Supplemental Figure 4. Xylan profiling of the uxt1 mutants.

Supplemental Figure 5. Immunoblot analysis of N-glycosylation in the uxt mutants.

Supplemental Table 1. Calculations of UXT protein contents in reconstituted proteoliposomes used for transport assays.

Supplemental Table 2. Monosaccharide composition of UXT mutant cell wall preparations derived from different Arabidopsis organs.

Supplemental Table 3. Primer list.

Supplemental Data Set 1. Text file of the alignment used for the phylogenetic analysis shown in Figure 1.

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AUTHOR CONTRIBUTIONS

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Identification and Characterization of a Golgi-Localized UDP-Xylose Transporter Family from Arabidopsis

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