

The Galactose Residues of Xyloglucan Are Essential to Maintain Mechanical Strength of the Primary Cell Walls in *Arabidopsis* during Growth¹

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In land plants, xyloglucans (XyGs) tether cellulose microfibrils into a strong but extensible cell wall. The *MUR2* and *MUR3* genes of *Arabidopsis* encode XyG-specific fucosyl and galactosyl transferases, respectively. Mutations of these genes give precisely altered XyG structures missing one or both of these subtending sugar residues. Tensile strength measurements of etiolated hypocotyls revealed that galactosylation rather than fucosylation of the side chains is essential for maintenance of wall strength. Symptomatic of this loss of tensile strength is an abnormal swelling of the cells at the base of fully grown hypocotyls as well as bulging and marked increase in the diameter of the epidermal and underlying cortical cells. The presence of subtending galactosyl residues markedly enhance the activities of XyG endotransglucosylases and the accessibility of XyG to their action, indicating a role for this enzyme activity in XyG cleavage and religation in the wall during growth for maintenance of tensile strength. Although a shortening of XyGs that normally accompanies cell elongation appears to be slightly reduced, galactosylation of the XyGs is not strictly required for cell elongation, for lengthening the polymers that occurs in the wall upon secretion, or for binding of the XyGs to cellulose.

The plant cell wall is continually modified during cell growth and differentiation. The tensile strength of the wall is provided by a dense spool of cellulose microfibrils interlaced with cross-linking glycans (Carpita and Gibeaut, 1993). When plant cells grow, the wall is biochemically “loosened” to permit turgor-driven cell expansion (Cosgrove, 2000). One of the mysteries of cell growth in plants that researchers have pondered for decades is how these interlaced glycans loosen without compromising the tensile strength of the pliant wall.

In all dicots and certain monocots, xyloglucans (XyGs) are the principal cellulose tethering molecules, and the loosening of these tethers from the microfibrils provides a physical control point of cell expansion (Cosgrove, 2000). The unique ability of XyG endotransglucosylases (XETs) to cleave XyGs and rejoin the cut ends with new partners suggested a role for these enzyme activities in wall loosening during growth and in the restructuring of cell walls

after extension (Nishitani and Tominaga, 1992). (A revised nomenclature has been adopted by consensus of researchers of the XyG endotransglucosylase/hydrolase gene/protein family [Rose et al., 2002]. The abbreviation XTH refers to any gene/protein of the family regardless of activity of the protein. However, when referring to endotransglucosylase activities, the abbreviation remains XET, and hydrolase activities are abbreviated XEH.) However, the only cell wall proteins proven to be capable of causing extension of isolated walls in vitro under mechanical stress are expansins, which increase wall extensibility under constant stress (McQueen-Mason et al., 1992), and yieldins, which lower the “yield threshold,” the minimum stress required that permits extension (Okamoto-Nakazato et al., 2000). Although XETs do not exhibit either of these activities in vitro (McQueen-Mason et al., 1993), a cooperativity between expansins and XET activity has been implicated in polymer lengthening during wall assembly (Thompson and Fry, 2001) and in the loosening and restructuring of the wall during cell growth and microfibril reorientation (Nishitani, 1998). The results of our studies with mutants with precisely altered XyG structures suggest a role for XET activity in the molecular grafting of XyGs—not for wall loosening during cell growth but principally to maintain tensile strength of the wall after growth.

Most XyGs consist of repeating heptasaccharide units of four β -D-glucosyl units linked (1→4), with three consecutive residues substituted with α -D-xylosyl units linked (1→6)- to the glucan backbone

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Table I. Distribution of oligomeric units in XyGs from leaves and hypocotyls of *Arabidopsis* wild-type, *mur2*, and *mur3* seedlings and cells of *mur2* and *mur3* maintained in continuous liquid cell culture

Protocols for the preparation of cell walls, extraction and digestion of XyG, and the quantitation of the unit structures are described by Vanzin et al. (2002). n.d., not detected.

Structure	Oligomer	Leaf			Hypocotyl			Cultured Cells	
		Wild Type	<i>mur2</i>	<i>mur3</i>	Wild Type	<i>mur2</i>	<i>mur3</i>	<i>mur2</i>	<i>mur3</i>
% of oligomers									
	XXXG	44.9	33.4	55.3	37.4	42.3	83.9	26.7	100.0
	XLG	8.5	36.8	n.d.	14.7	49.7	n.d.	57.1	n.d.
	XLXG	3.1	7.7	44.7	1.9	6.2	16.1	5.7	n.d.
	XXFG	23.8	n.d.	n.d.	35.1	n.d.	n.d.	n.d.	n.d.
	XLG	4.0	22.1	n.d.	0.8	1.8	n.d.	10.5	n.d.
	XLFG	15.7	n.d.	n.d.	10.1	n.d.	n.d.	n.d.	n.d.

(Table I). About one-half of these units contain extensions of β -D-galactosyl-(1 \rightarrow 2)- upon the first xylosyl (closest to the reducing end) of the glucan or to the middle Xyl or to both residues. An α -L-fucosyl-(1 \rightarrow 2)- residue is then added to the Gal residue at the first position (Carpita and Gibeaut, 1993). Cleavage of XyG with a sequence-specific *Trichoderma* endoglucanase yields six kinds of oligomeric units that constitute a species-specific profile. In addition to the fundamental Xyl₃Glc₄ oligomer, called XXXG in a standardized nomenclature (Fry et al., 1993), the other oligomers are five possible permutations formed by addition of galactosyl residues at the first or middle Xyl residue to give XLG or XLXG, and XLG, and of subsequent addition of L-Fuc upon the first galactosyl unit, if present, to give XXFG and XLFG (Table I).

Two major hypotheses have been proposed for the function of the trisaccharide side group, α -L-Fuc-(1 \rightarrow 2)- β -D-Gal-(1 \rightarrow 2)- α -D-Xyl-, with particular importance placed on a requirement of the terminal fucosyl residue. First, computer-modeling studies of three-dimensional XyG structures suggest that the glucan backbone may assume either a twisted or straightened conformation, and trisaccharide side groups stabilize the straightened form to facilitate steric bonding with cellulose microfibrils (Levy et al., 1991). Second, the XyG oligomers containing the fucosylated trisaccharide side groups modulate auxin-induced growth in excised sections (Zablackis et al., 1996).

Two *Arabidopsis* mutants, called *mur2* and *mur3*, were selected on the basis of an underrepresentation of Fuc in cell wall polymers (Reiter et al., 1997), and both of them affect XyG trisaccharide side-group

structure specifically. *MUR2* encodes a XyG-specific fucosyltransferase, which leads to the absence of XyG fucosylation in *mur2* plants (Vanzin et al., 2002), and *MUR3* encodes the galactosyltransferase specifically responsible for the first step in the formation of the α -L-Fuc-(1 \rightarrow 2)- β -D-Gal-(1 \rightarrow 2)- side group (Madson et al., 2003). The *mur3* mutation eliminates the entire disaccharide extension from the first Xyl and results in marked enhancement of galactosylation at the middle Xyl in the instance of leaf-derived XyG. Remarkably, the growth habits of *mur2* and *mur3* plants are indistinguishable from those of wild type despite such radical alteration of their fundamental cross-linking polymers (Reiter et al., 1997). The tensile strengths of *mur2* and *mur3* floral stems were also comparable with wild type (Vanzin et al., 2002; Madson et al., 2003). Thus, if a fucosylated trisaccharide functions in facilitating normal growth or increasing the tensile strength of the wall during growth, then this functional property can be replaced by increased degrees of galactosylation.

Whereas growth form and physiology of both *mur2* and *mur3* mutants are indistinguishable from wild type in the shoot, we observed a strong phenotype in etiolated hypocotyls of *mur3* seedlings. The tensile strengths of the hypocotyls was less than 40% those of wild-type or *mur2* hypocotyls, and a visible swelling of the base of the hypocotyls occurred as a result of grossly enlarged epidermal and cortical cells. The phenotype is associated with a failure in hypocotyls of the enhancement of galactosylation of the middle Xyl in *mur3* XyGs that occurs in *mur3* shoot tissues. No other feature was affected by the altered XyG structure, because cell elongation, growth form, tenacity of cellulose binding, and polymer lengthening

that occurs in *muro* were all indistinguishable from wild type. However, we found that XET activity is markedly enhanced by the presence of galactosyl residues, indicating that this enzyme plays a role in remodeling and religating XyGs during growth to maintain tensile strength.

RESULTS AND DISCUSSION

The hypocotyl is a tapered cylinder of tissue that elongates in darkness to about 2 cm (Fig. 1A). The shape of the hypocotyl “hook” and the initiation of cell elongation are indistinguishable among wild-type and *mur* mutants (Fig. 1, B–D). During maximal growth, the wild-type hypocotyls is about 260 μm at the base and tapers to 190 μm just below the hook. After elongation, the base of the hypocotyls continues to enlarge to about 300 μm . However, the average diameters of the base of the *mur3* hypocotyl swell to nearly 400 μm , and the epidermal cells exhibit abnormal swelling or bulging that accompanies the onset of elongation but disappears in wild-type and *mur2* hypocotyls after growth has ceased (Fig. 1, E–G). Although the epidermal cells display obvious bulging, the increases in hypocotyl diameters are due primarily to increases in diameters of the cortical cells comprising two layers within the epidermis. The diameters of the inner layer of eight cortical cells of wild type are about 45 and 48 μm in periclinal and radial directions, respectively, whereas *mur2* inner cells are somewhat smaller at 40 to 42 μm . However, the *mur3* inner cortical cells swell to more than 70 μm radially and about 65 μm periclinally (Table II). The cortical cells underlying the epidermis were also larger in *mur3*, which have radial diameters of about 65 μm and periclinal diameters of 59 μm . The comparable cortical cells of wild type and *mur2* were more similar to the inner cells and varied slightly from 43 to 50 μm in both radial and periclinal directions. In *mur3*, the cortical cells underlying the epidermis are smaller than those of the inner cells underlying them. Whereas the number of inner cells is an invariant eight in wild type and mutants, the underlying cells of *mur3* increase one or two cells compared with wild type and *mur2*, from about 14 to 16, and this factor resulted in slightly lower diameters compared with the inner cells (Table II). Epidermal cells are much smaller than cortical cells. Wild-type and *mur2* epidermal cells are nearly isodiametric, with radial diameters of about 22 μm , and 23 to 24 μm in the periclinal direction. By comparison, the epidermal cells of *mur3* are larger than either wild type or *mur2*, particularly in the periclinal direction, where the average diameters are 32 μm (Table II).

During maximal rates of growth, the tensile strength of the *mur3* hypocotyls is less than one-half that of wild type, whereas the strength of *mur2* hypocotyls is only slightly lowered (Fig. 2). The XyGs

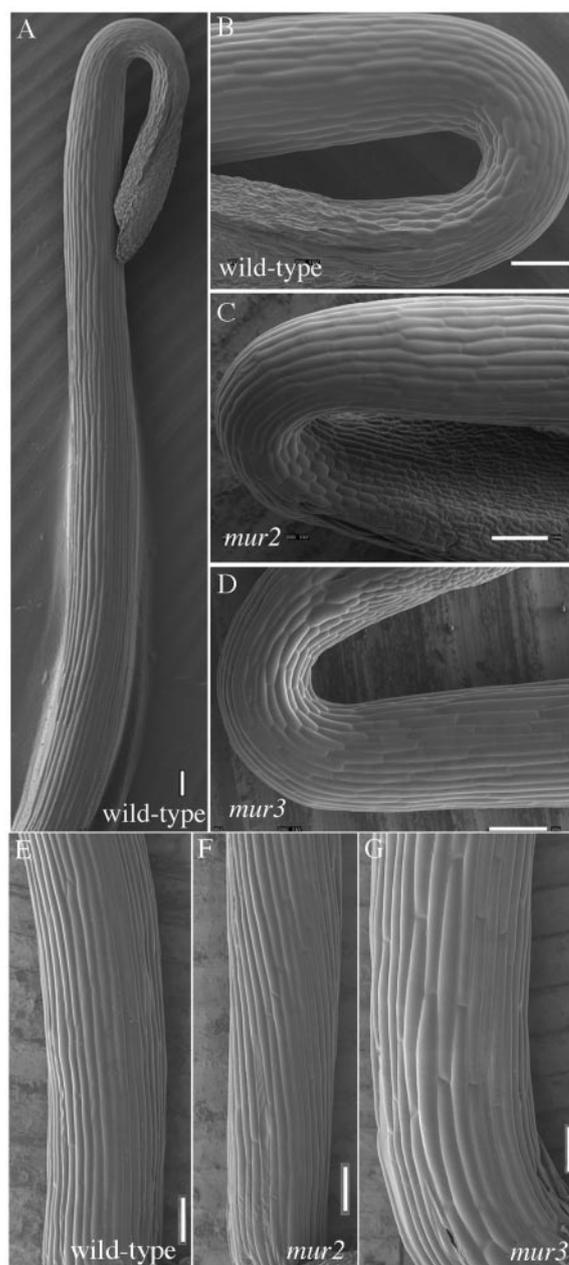


Figure 1. SEM images of elongating *Arabidopsis* etiolated wild-type, *mur2*, and *mur3* hypocotyls. A, Low magnification SEM image of 3.5-d-old wild-type hypocotyl. Elongation begins just below the hypocotyls hook. B through D, Wild-type, *mur2*, and *mur3* hypocotyl “hook” regions, where cell elongation begins. Bars = 50 μm . E through G, The SEM images of the basal cells of 4.5- to 5.2-d-old hypocotyls demonstrate the pronounced swelling of the *mur3* epidermal cells after growth compared with that of *mur2* and wild-type cells. The diameters of bases of the hypocotyls are wild type ($297 \pm 5 \mu\text{m}$), *mur2* ($274 \pm 15 \mu\text{m}$), and *mur3* ($395 \pm 5 \mu\text{m}$; minimum sample size = 20). Bars = 100 μm .

from wild-type, *mur2*, and *mur3* etiolated hypocotyls all exhibit much reduced galactosylation of the middle Xyl of leaf XyGs (Table I). More than 80% of *mur3* XyG is composed of XXXG units. The swollen nature

Table II. Diameters of the epidermal cells, the cortical cells underlying the epidermal layer, and the inner cortical cells of wild-type, *mur2*, and *mur3* etiolated hypocotyls determined at the culmination of growth

Diameters were determined in both radial and periclinal directions from automated analysis of digital images of basal cross sections. Values are the mean \pm SD of at least four hypocotyls each.

Line	Epidermal		Underlying Cortical		Inner Cortical	
	Radial	Periclinal	Radial	Periclinal	Radial	Periclinal
	μm					
Wild type	22.4 \pm 1.1	23.0 \pm 0.3	49.9 \pm 3.7	48.1 \pm 3.9	47.6 \pm 2.4	44.9 \pm 1.8
<i>mur2</i>	21.6 \pm 1.5	23.5 \pm 2.9	43.9 \pm 2.0	43.1 \pm 2.6	40.6 \pm 7.0	41.6 \pm 3.5
<i>mur3</i>	26.1 \pm 3.1	32.3 \pm 0.3	65.4 \pm 4.4	59.3 \pm 2.1	70.8 \pm 4.8	64.6 \pm 3.5

of the basal cells and loss of tensile strength in *mur3* hypocotyls correlate with the presence of a poorly galactosylated XyG (Table I).

The *mur2* and *mur3* hypocotyls contain more cellulose per unit length than those of wild type (about 54.2 \pm 6.1 and 54.9 \pm 7.3 ng mm⁻¹ in *mur2* and *mur3*, respectively, compared with 41.2 \pm 3.6 ng mm⁻¹ in wild type) and proportionally higher amounts of XyG and pectins. Thus, the swollen phenotype and loss of tensile strength cannot be attributed to lower wall mass or changes in composition of other matrix polysaccharides besides XyG. The *mur3* hypocotyl XyGs are assembled upon cellulose and exhibit the same tenacity of binding as the galactosylated XyGs of wild-type and *mur2* hypocotyls (Fig. 3). Whereas 4

m NaOH is required to extract hypocotyl XyGs from walls of wild-type and *mur* mutants, these extracted XyGs rebind equally well to purified crystalline cellulose but require only about 0.6 m NaOH to dislodge them (Fig. 3). In contrast to the predictions of Levy et al. (1997), the tenacity of binding in vivo and in vitro show that the absence of the Gal residues has no influence on the physical interaction of XyG and cellulose during wall assembly in the living hypocotyl. The greater tenacity of binding in the nascent wall implicates an assembly mechanism that is lost in vitro, but this mechanism does not rely on Gal residues. On the basis of gel permeation chromatography, the molecular size distribution of the XyGs from *mur2* and *mur3* hypocotyls is shifted to slightly larger masses than those of wild type, indicating no apparent loss of the ability for the extracellular assembly of long XyGs from the secreted precursor polymers (Fig. 4). On the contrary, wild-type XyGs are generally shorter than either *mur2* or *mur3* XyGs. A shift to

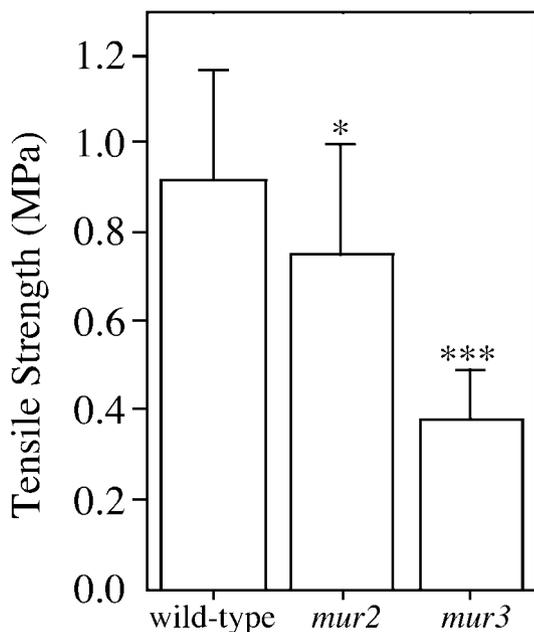


Figure 2. Tensile strengths of the basal 3 mm of 4-d-old wild-type, *mur2*, and *mur3* hypocotyls. Etiolated hypocotyls were grown and tested submerged with a TA-XT2i texture analyzer (Stable Microsystems, Godalming Surrey, UK), with a load cell sensitive to 1 mN (Ryden et al., 2003). Tensile strength values and error bars are the means \pm SD of 20 samples. Errors in strength were proportional to absolute strength, so the statistical comparisons used the log_e transformation, comparing the strength of each mutant with that of the wild type (*, $P < 0.05$; ***, $P < 0.001$).

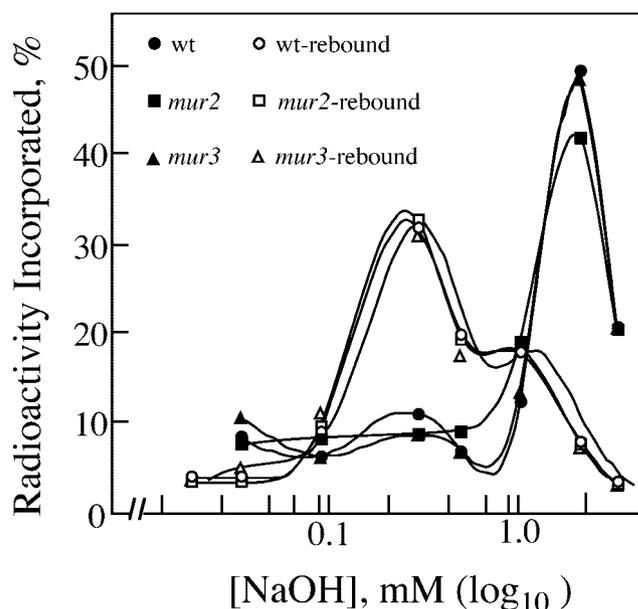


Figure 3. Tenacity of binding of wild-type, *mur2*, and *mur3* XyGs to cellulose in vivo and in vitro. Seeds of each were germinated in [¹⁴C]-D-Glc, and the cell walls of the hypocotyls were purified and depectinated. Values are the proportions of labeled XyGs extracted from the walls at each step of increasing alkali concentration.

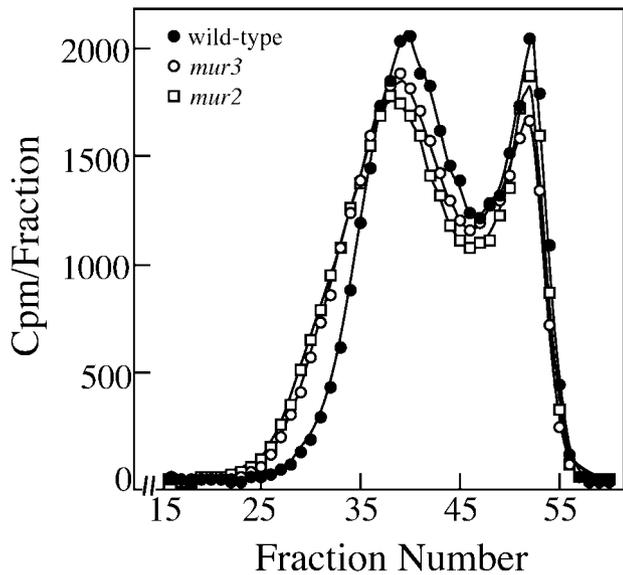


Figure 4. Molecular mass distributions of wild-type, *mur2*, and *mur3* XyGs. Radiolabeled hypocotyls XyGs were purified from the 4 M NaOH extract of purified depectinated walls pre-extracted with 1 M NaOH. The XyGs in 1 M NaOH were loaded onto a 60- × 2.5-cm column of Sepharose 4B-CL (Pharmacia) equilibrated in 1 M NaOH. Fractions (3.5 mL) were collected into 2 mL of 2 M acetic acid; 2 mL was assayed for radioactivity by liquid scintillation spectroscopy.

shorter XyGs accompanies auxin-induced cell elongation of excised pea (*Pisum sativum*) epicotyls (Talbot and Ray, 1992), and an anti-sensed member of the XyG endotransglucosylase/hydrolase (XTH) gene family blocks the growth-dependent decrease in the molecular size distribution of tobacco (*Nicotiana tabacum*) XyGs (Herbers et al., 2001). Together, these data

indicate that these precisely altered XyG structures have little impact on their extended polymerization in muro or on their assembly around cellulose into a functional wall. What appears to be affected specifically is the remodeling of the wall structure to maintain tensile strength. The discrepancy of molecular sizes between wild-type versus *mur2* and *mur3* XyGs prompted us to examine the XET activity against these altered polymers.

The activities of Arabidopsis hypocotyl XETs in vitro are markedly enhanced by the side-group extensions of XyGs. When the concentration-dependent activities of XETs are tested on purified, buffer-soluble XyGs, highly galactosylated tamarind seed storage XyGs are the best substrate (Fig. 5A). XyGs from *mur3* and wild-type leaves, which contain predominantly XLXG and XXFG oligomers, respectively (Table I), are also good substrates for the enzyme. The *mur2* XyG extracted from late-stationary phase cells in liquid culture, which contain predominantly XXXG and XLXG, are poorer substrates, but XyGs from *mur3* late-stationary-cultured cells, which contain only XXXG, are the poorest substrates (Fig. 5A). In addition to the galactosyl residues, XET activity is also somewhat dependent on molecular size of the XyGs (Nishitani and Tominaga, 1992). Total XyGs extracted from plant tissues provide a rich array of sizes (Fig. 4). Although slight differences in the size distribution were observed between wild-type and *mur2* and *mur3* XyGs, given the size-dependent differences in activity predicted by the data of Nishitani and Tominaga (1992), it is unlikely that these small differences can account for the gross differences in XET activity (Fig. 5A). The XyGs extracted from late-stationary culture cells and tamarind XyGs behave

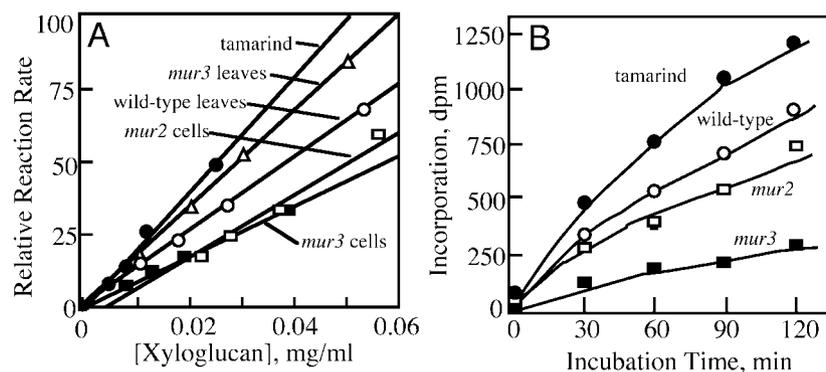


Figure 5. Activity of XET with altered XyGs as substrates. Total XTHs were purified from Arabidopsis hypocotyls and cells. The enzyme mixture was assayed by a paper-binding assay (Fry et al., 1992), measured as attachment of the various XyG substrates to a common [^3H]XLLGol acceptor or mixed-oligomer preparation of mostly XLLGol, which was synthesized by chemical reduction of high-performance anion exchange chromatography-purified XLLG by NaB^3H_4 (Steele and Fry, 1999). A, Concentration-dependent hypocotyl XET activity with soluble XyGs. Sources of the XyG enriched in specific oligomers are: *mur3* cells in liquid culture (only XXXG), *mur2* cells in liquid culture (predominately XXXG and XLXG), *mur3* leaves (predominately XXXG and XLXG), wild-type leaves (predominately XXXG, XXFG, and XLFG), and tamarind seed flour XyG (predominately XXXG, XLXG, and XLLG). The XyG oligomeric distributions for each substrate are shown in Table I. B, Activity of Arabidopsis cultured cell XET with purified XyGs from tamarind seed flours or from hypocotyls of wild type, *mur2*, and *mur3*. The XyGs were suspended in boiling water at 3 mg mL $^{-1}$ and cooled, and insoluble material was removed from the supernatant before assay of the substrate remaining soluble.

chromatographically like those of the hypocotyls XyGs shown in Figure 4 (data not shown).

When boiled XyG-enriched preparations from hypocotyls were tested at relatively high concentrations of substrate, enhanced XET activity against Arabidopsis XyGs is observed when they are enriched in galactosylated oligomers, but *mur3* hypocotyl XyGs are exceptionally poor substrates (Fig. 5B). Upon centrifugation of the reaction mixture, the *mur3* preparation is significantly depleted of XyG in the supernatant, suggesting that the relatively ungalactosylated XyGs were inaccessible to enzyme action. Thus, the poor activity against XyGs with little or no side-group substitution is a consequence of two factors. First, XET activities with water-soluble XyGs enriched in XLXG or XXFG oligomers directly enhance enzyme recognition of the substrate (Fig. 5A). Second, loss of galactosyl residues from XyG changes the physical state and lowers solubility (Shirakawa et al., 1998; Yamanaka et al., 1999), and this feature renders the substrate inaccessible to XET action. XyGs exist in at least two domains in the wall, one that is accessible to enzymes that modify its structure and one that is not (Pauly et al., 1999), and there is a shift of polymers from the former to the latter during cell growth (Pauly et al., 2001). The *mur3* hypocotyl XyGs may more rapidly enter the enzyme-inaccessible domain. As Levy et al. (1997) described, the backbone can exist in twisted and straightened conformations, and XLXG and XXFG are particularly effective in maintenance of the backbone in the straightened conformation. Thus, if the computer simulations of Levy et al. (1997) are correct, then the chain straightening function may enhance the binding of any member of the XTH family to XyG (Fig. 5A). However, the behavior of polymeric substrates is to a certain extent consistent with the activity of oligomeric substrates, where XLLG is significantly better acceptor than either XXXG or XXFG (Purugganan et al., 1997), indicating a strong preference toward substrate and acceptor with Gal residues on the middle Xyl.

A role of XyGs in tensile strength runs contrary to biophysical data suggesting that the addition of unfucosylated XyG to cellulose composites decreases strength (Chanliaud et al., 2002). However, these biophysical parameters are valid for unilamellate artificial structures devoid of enzymatic activities that may drastically alter the biochemical interactions among its molecular components. The formation of the hook and initiation of cell elongation proceed indistinguishably between wild type and mutant (Fig. 1, B–D), and elongation rates of the *mur3* hypocotyl are only lowered marginally, demonstrating that the predicted lowering of XET activity does not impair cell growth. This is a surprising finding given evidence implicating XET activity directly in growth (Takeda et al., 2002). However, the abnormal swelling and bulging of the cortical and epidermal cells of *mur3* hypocotyls occur well after growth (Fig. 1, E–G;

Table II). This phenomenon may result from failure to religate XyGs after wall expansion and, thus, be symptomatic of the loss of tensile strength (Fig. 2).

Although the galactosylation of XyGs has a marked effect only on tensile strength and activity of XET, the roles of these enzymes in other events of wall dynamics remain to be confirmed by direct experiments. In Arabidopsis, these enzymes are encoded by a XTH gene family (Rose et al., 2002) comprising nearly three dozen members in at least three subclasses (Campbell and Braam, 1999; Yokoyama and Nishitani, 2001; Rose et al., 2002). The numerous XTH isoforms suggest a multiplicity of roles in cell development (Yokoyama and Nishitani, 2001). Thus, if members of some classes of XTH function in polymer elongation, as has been suggested (Thompson and Fry, 2001), then Gal residues are not essential for these particular reactions. Likewise, *mur3* and wild-type hypocotyl XyGs bind equally well to cellulose, and the enhanced tenacity of native XyG binding compared with rebinding in vitro suggest that proteins other than XTHs facilitate assembly in vivo.

Mutants and alteration of expression of XTHs have given mixed results in demonstration of an association between XET or XEH activities and growth and wall remodeling (Campbell and Braam, 1999). No less than 10 XTHs are highly expressed in Arabidopsis roots (Yokoyama and Nishitani, 2001), although the number expressed specifically in hypocotyls is unknown. Redundancy of expression of several XTHs with similar function in any one cell may explain the difficulty using this approach to characterize gene function (Campbell and Braam, 1999; Yokoyama and Nishitani, 2001). In the experiments described here, the alteration of a common substrate has been more informative than underexpression of a single XTH gene. The total complement of XTHs from cell and hypocotyls were extracted to determine all nascent activities to be expected. All of the members of the XTH family, regardless of binding to XyG during affinity purification or not, exhibit poor activity toward ungalactosylated XyGs. For the first time, a role for XET activity in remodeling the wall through religation of XyGs to tighten the wall after wall extension finally has empirical data to support the concept.

Although the collective data in this study implicate XET activity in wall restructuring to maintain tensile strength, the data are still indirect, and the function of XTHs in this role could be bolstered by measurements of activity in muro. XyGs are integral to growth because of their dynamic interaction with cellulose microfibrils, and alterations of structure could impact other enzymes and proteins for which XyG is a common substrate. To our knowledge, the activity of expansins in in vitro extension assays with *mur2* and *mur3* hypocotyls has not been examined. Also, hydrolysis of XyG by α -xylosidases and

β -glucosidases is modulated by Gal residues at the middle Xyl (Edwards et al., 1988; de Alcântara et al., 1999), and this feature may impact the failure of *mur2* and *mur3* hypocotyl XyGs to undergo shortening during growth (Fig. 4). A β -galactosidase is active only against XyG oligomers (or the non-reducing terminal oligomer of XyG polymers) and removes solely the middle galactosyl residue. This β -galactosidase activity generates an oligomer that becomes accessible to α -xylosidases (Fanutti et al., 1991) and β -glucosidases (Crombie et al., 1998) that act cooperatively to digest the xylosyl and backbone glucosyl units. However, these activities cannot occur in the absence of XET or XEH activity, and in contrast to the observed results (Fig. 4), the *mur2* and *mur3* should be better substrates for degradation if XET or XEH activities were optimal.

MATERIALS AND METHODS

Growth of Plants

Seeds of *Arabidopsis* (ecotype Columbia) were from wild-type or *mur2-1* and *mur3-1* lines that had been back-crossed at least four times (Reiter et al., 1997). Surface-sterilized seeds were chilled in water for 3 d to synchronize the germination response and then plated on one-half-strength Murashige and Skoog salts (Sigma-Aldrich, St. Louis), pH 5.7, in 0.8% (w/v) plant agar (Sigma-Aldrich). Etiolated hypocotyls were harvested after germination and grown for up to 5 d in darkness, whereas plantlets were obtained after 14 d of growth on agar at 23°C under continuous light of approximately 60 $\mu\text{mol m}^{-2} \text{s}^{-1}$ from high-output fluorescent lamps. Leaves of plants, grown as described (Vanzin et al., 2002), were harvested at 32 d. Callus cultures were initiated from wild-type seedlings grown on an agar medium of Gamborg salts (Sigma-Aldrich), pH 5.7, supplemented with 5 μM 2,4-dichlorophenoxyacetic acid and 2% (w/v) Suc, and the callus was used to initiate liquid cell suspension cultures, with sub-culture every 2 to 3 weeks.

Purification of XyGs

Plant materials were frozen in liquid nitrogen and homogenized in a glass-glass grinder (Duall, Kontes Glass, Vineland, NJ) in 50 mM Tris[HCl], pH 7.2, supplemented with 1% (w/v) SDS. The homogenate was heated to 65°C for 15 min, and the walls were collected on a nylon mesh (47- μm square pores; Nitex, Briarcliff, NJ). The walls were washed extensively with water, ethanol, acetone, and finally suspended in water. Pectins were removed from the walls by extraction with excess 2 M imidazole[HCl], pH 7, at ambient temperature, followed by extraction with up to 0.045 M NaOH (supplemented with 3 mg mL⁻¹ sodium borohydride to prevent end elimination). De-pectinated walls were then subjected to 1 M NaOH (with borohydride) to extract other material and additional pectin, and XyGs were extracted with excess 4 M NaOH (with borohydride) overnight with constant stirring under an argon atmosphere. The 4 M extract was passed through a glass-fiber filter mat to remove unsedimented wall remnants, and the eluant was chilled to ice temperature and acidified to pH 5 with glacial acetic acid. The XyGs were dialyzed extensively against running deionized water, and either used directly or freeze-dried. In some experiments, freeze-dried XyGs were suspended in water and boiled for 10 min to dissolve the polymers, and then Sephadex A-25 anion exchange resin was added to remove a small amount of contaminating uronic-acid rich polymers.

Determination of the Distribution of the Oligomers

The percentage of each oligomeric unit is based on separation of Trichoderma endo- β -glucanase (Megazyme, Bray, County Wicklow, Ireland) digests of XyG oligomers from leaves, hypocotyls, and cultured cells that were separated by high-performance anion exchange chromatography on a CarboPac PA-1 (Dionex, Sunnyvale, CA) anion exchange column and detected

by pulsed amperometric detection (Vanzin et al., 2002). Electrospray mass spectrometry (MS) of the total digest and MS/MS of *m/z* 1247 were used to determine the proportions of XXLG and XLXG.

Microscopy

Hypocotyls harvested at several times during rapid elongation phase (3.5- to 5.5-d-old) were rapidly frozen by plunging them into nitrogen slush and then sputter-coating them with gold for 4 s at -165°C. Samples were then imaged at -140°C in a scanning electron microscope (JSM-840, JEOL, Tokyo) using 5 kV of accelerating voltage. For determinations of cell diameters, etiolated hypocotyls were fixed in a mixture of 2.5% (w/v) glutaraldehyde and 2% (w/v) paraformaldehyde in 0.05 M sodium phosphate, pH 6.8, washed with the same buffer, and post-fixed in 1% (w/v) osmium tetroxide in the same buffer. After dehydration in ethanol, the hypocotyls were embedded in Spurr's resin. Semithin sections were cut from the base of a minimum of four hypocotyls and observed by bright-field microscopy in a microscope (AH-2, Olympus, Tokyo), and the images were recorded digitally by a SPOT RT camera accessory (Diagnostic Instruments, Sterling Heights, MI).

Determination of Tensile Strength

Etiolated 4-d-old hypocotyls were grown and tested submerged as described (Ryden et al., 2003) using a TA-XT2i texture analyzer (Stable Microsystems, Godalming Surrey, UK) with a load cell sensitive to 1 mn. In brief, tensile tests were made on about 4-mm lengths of hypocotyls starting from just above the root junction. The hypocotyls were fixed with cyanoacrylate (Resist H20, Holdtite, Gateshead, UK) across two aluminum tabs about 3 mm apart, and the adhesive was cured rapidly with an activator (Cyanolit Plus, Eurobond, Sittingbourne UK). Taking specimens to be cylindrical, the tensile strength was derived (Cleland, 1967). The average hypocotyl diameters were: wild type, 260 \pm 16 μm ; *mur2*, 248 \pm 23 μm ; and *mur3*, 287 \pm 11 μm . Tensile strength values and error bars are the means \pm SD of 20 samples, and Dunnett's test was performed as described in the Minitab Release 10 Reference Manual (1994).

Determination of Tenacity of XyG Binding to Cellulose In Vivo and In Vitro

Arabidopsis wild-type, *mur2*, and *mur3* hypocotyls were labeled by addition of [¹⁴C]D-Glc to the agar growth medium. Walls were purified and depectinated as described previously. The cell walls were then extracted sequentially with increasing concentrations of NaOH (each with 3 mg mL⁻¹ sodium borohydride), and after neutralization, the amount of radioactivity was determined by liquid scintillation counting of a small sample of each supernatant. The 4 M NaOH extracted the bulk of the wall material, which was about 80 mole % XyG based on monosaccharide analysis. In a separate experiment, XyGs from cell walls, depectinated as above and pre-extracted with up to 1 M NaOH, were exhaustively extracted with 4 M NaOH, and the supernatant was filtered over glass-fiber mats, mixed with a 10-fold excess of pre-alkali extracted microcrystalline cellulose (Cellex N-1, Bio-Rad Laboratories, Hercules, CA), and gradually neutralized with acetic acid and vigorous stirring over a 2-h period at ambient temperature. The cellulose with bound labeled XyGs were then pelleted by centrifugation, washed several times with water and 0.02 M NaOH, and subjected to a gradient extraction with NaOH as before.

Activities of *Arabidopsis* Hypocotyls XET with Precisely Altered XyGs

XTHs were isolated from the medium, and 0.2 M CaCl₂ extracts of cell walls were purified from *Arabidopsis* cells and hypocotyls and precipitated by increasing saturation with ammonium sulfate as described (Steele and Fry, 1999). A preparation containing most of the XET activity precipitated between 20% and 70% saturation. The XTHs were also purified by affinity binding to XyG essentially as described (Steele and Fry, 1999). In brief, XTHs dissolved in 100 mM succinate[NaOH], pH 5.5, and 10 mM CaCl₂, and 0.2% (w/v) tamarind XyG was added. In the absence of oligomers, the bound XTHs void a column of Bio-Gel A-0.5m (Bio-Rad Laboratories) equilibrated

in the suspension buffer, and upon addition of excess tamarind oligomers, generated by *Trichoderma* endoglucanase digestion, XTHs elute near the included volume. After concentration, the enzymes were separated osmotically from the oligomers by passing them over a Sephadex G-25 de-salting column (Pharmacia, Uppsala).

The various XyGs were mixed at up to 3 mg mL⁻¹ in 100 mM sodium succinate, pH 5.5, and boiled to maximize solubility, and after cooling to ambient temperature, the insoluble residues were pelleted by centrifugation at 14,000 rpm in a microfuge. The XyG concentration remaining in the supernatant was estimated by phenol-sulfuric sugar assay (DuBois et al., 1956). In experiments designed to determine XyG concentration-dependent activity, affinity-purified *Arabidopsis* cell XTHs were used, and XET activities were tested against the supernatant liquids and were determined by the attachment of the XyG fragment to the non-reducing end of [³H]XyG oligomer acceptor in a filter paper-binding assay (Fry et al., 1992). As stated in the figure legend, either purified [³H]XLLGol or [³H]tamarind XyG mixed oligomers were used as acceptors. Aliquots of the reaction were stopped by addition of 50% (v/v) formic acid at various times and air-dried on strips of filter paper (No. 1, Whatman, Clifton, NJ). The strips were then washed extensively with water to remove unreacted [³H]oligomers, and the radioactivity bound to the filter paper was assayed by liquid scintillation spectroscopy. In experiments involving depleted XyG preparations, hypocotyl XTHs between 20% and 70% (w/w) ammonium sulfate saturation were used without affinity purification.

Cellulose Binding Assays

Shoot XyGs were radiolabeled by incubation of 30-d-old wild-type, *mur2*, and *mur3* plants with ¹⁴CO₂ in a sealed fumigation chamber at 25°C under 60 μmol m⁻² s⁻¹ light from fluorescent lamps for 2 d. Leaves were harvested into liquid nitrogen, and cell walls were prepared as described earlier. XyGs, solubilized by 4 M NaOH from 1 M NaOH-pre-extracted walls, were collected, neutralized, dialyzed against water, and freeze-dried. The dry XyGs were suspended in 5 mM succinate [NaOH], pH 5.8, boiled to maximize solubility, and cooled to ambient temperature. A small amount of insoluble material was removed by centrifugation at 14,000g in a microfuge, and equimolar amounts of soluble XyG (independent of radiolabel) were mixed with a 10-fold excess of microcrystalline cellulose (Cellex N-1, Bio-Rad Laboratories) that was pre-washed with 4 M NaOH (supplemented with 3 mg mL⁻¹ of sodium borohydride) and washed extensively with water and finally suspended in 5 mM succinate [NaOH], pH 5.8, to start the reactions. Binding at 25°C was determined by centrifugation of a portion of the reaction mixture, and assay of the depletion of the labeled XyG from soluble fraction by liquid scintillation spectroscopy. Residual label incapable of binding to cellulose was subtracted from the total when binding rates were calculated.

Molecular Mass Distribution of XyGs

The molecular mass distributions of radiolabeled hypocotyls XyGs used in the cellulose binding assays were purified from the 4 M NaOH-extract of purified depectinated walls pre-extracted with 1 M NaOH. The 4 M NaOH extract was filtered of small remnants of unsedimented cell walls, chilled and neutralized with glacial acetic acid, dialyzed against running deionized water, and freeze-dried. The materials were dissolved 1 M NaOH, and a small amount of insoluble material was removed by centrifugation. An equivalent amount of radioactivity in the 1 M NaOH solution was applied to a 60 × 2.5-cm column of Sepharose 4B-CL (Pharmacia) equilibrated in 1 M NaOH. Fractions (3.5 mL) were collected into 2 mL of 2 M acetic acid; 2 mL was assayed for radioactivity by liquid scintillation spectroscopy, and the remainders were pooled into high- and low-mass fractions, dialyzed against deionized water, and lyophilized for monosaccharide analysis. The XyGs were judged to comprise the vast majority of each fraction based on the amounts and ratios of Xyl and Glc.

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