ARTICLE ADDENDUM

Glycosylation of arabinogalactan-proteins essential for development in Arabidopsis

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ABSTRACT

Arabinogalactan-proteins (AGPs) are ubiquitous cell wall components present throughout the plant kingdom. They are extensively post translationally modified by conversion of proline to hydroxyproline (Hyp) and by addition of arabinogalactan (AG) polysaccharides to Hyp residues. Two small gene subfamilies within the CAZy GT31 family, referred to as Hyp-galactosyltransferases (Hyp-GALTs and HPGTs), encode enzymes that specifically add galactose to AGP protein backbones as revealed by heterologous expression of the genes coupled with an in vitro enzyme assay and by biochemical characterization of the genetic knock-out mutants. Biochemical analysis of galt2galt5 double and hpgt1hpgt2hpgt3 triple knockout mutants revealed significant reductions in both AGP-specific Hyp-GALT activity and β-Gal-Yariv precipitable AGPs. Further analysis of these mutants demonstrated both overlapping and distinct pleiotropic growth and development phenotypes, indicating the important contributions of the carbohydrate moieties toward AGP function. Current research indicates that all 8 Hyp-GALT/HPGT genes encode enzymes that catalyze the initial step for AGP glycosylation and that AGP glycans play essential roles in plant growth and development.

Introduction

Arabinogalactan-proteins (AGPs) are extensively glycosylated hydroxyproline (Hyp)-rich proteins found in cell walls, plasma membranes, and extracellular secretions of plants. They are involved in various aspects of plant growth and development such as cell proliferation, cell viability, cell division, reproductive development, regeneration of somatic embryos, abiotic stress, and cellular signaling.1-13 AGPs are widely distributed in the plant kingdom, being present in land plants as well as in bryophytes like Physcomitrella patens, basal angiosperms, and even in algae.14-16 AGPs also have several important industrial and biomedical applications including a direct food additive, a food emulsifier, a stabilizer, a sweetener, and an immunomodulatory molecule.17

Structurally, AGPs are diverse and heterogeneous due in part to their various protein backbones as well as the extent and degree of AG polysaccharide addition to the backbones. AGP protein backbones typically consist of repetitive dipeptide motifs such as Ala-Pro, Ser-Pro, Thr-Pro, and Val-Pro in which the Pro residues are converted to Hyp.18 About 40% of the AGPs are tethered to the plasma membrane by a glycosylphosphatidylinositol (GPI) anchor.19,20 AGP glycans or arabinogalactans (AGs) are characterized by the presence of a β-1,3-galactan backbone which is substituted with β-1,6-galactan side chains that are decorated further with arabinose, and less frequently with fucose, rhamnose, and (methyl) glucuronic acid.22 Using NMR analysis of synthetic AG peptide expressed in tobacco BY2 cells, Tan et al., however, concluded that the AG backbone is composed of repeating β-1,3-galactotriose units with or without side chains, which are connected by β-1,6-linkages, to generate a kinked galactan backbone.22 While debate continues over these 2 models of AG structure, it should be noted that based on linkage and mass spectroscopy analysis, longer β-1,6-galactan side chains have been reported for AGPs from radish root, wheat flour and Arabidopsis leaf.23-25

Several glycosyltransferases (GTs) are involved in assembling the AG chains, and these GTs likely
coordinate and regulate the density, length and sequence of the AG chains. Given that the sugar side chains typically account for more than 90% of the mass of AGPs, sugars likely define the interactive molecular surface of AGPs and govern their function. Several of the GTs for AG biosynthesis are known and cloned; however, many await discovery as previously reviewed.\textsuperscript{21,26} O-glycosylation of AGPs is initiated by a set of 8 Hyp-galactosyltransferases (Hyp-GALTs/HPGTs), which are members of the GT31 family of the CAZy database and designated as GALT2-6 and HPGT1-3.\textsuperscript{12,27-29} These enzymes add a single galactose residue to Hyp residues in AGP core proteins. Other known and cloned GTs include: one $\beta-1,3$-galactosyltransferase in GT31, 2 $\beta-1,6$-galactosyltransferases, GALT31A and GALT29A in GT31 and GT29 respectively, 3 $\beta$-glucuronosyltransferases, GlcAT14A, GlcAT14B, and GlcAT14C, in GT14,\textsuperscript{34,35} 2 $\alpha$-fucosyltransferases, FUT4 and FUT6, in GT37, and one $\beta$-arabinosyltransferase, Reduced Arabinose Yariv1 or RAY1, in GT77.\textsuperscript{30-32} Several other GTs for AG biosynthesis await identification and cloning, including $\alpha$-arabinosyltransferases, $\alpha$-rhamnosyltransferases, and other $\beta$-galactosyltransferases.

With regard to the 2 small gene families within GT31 that exhibit Hyp-GALT enzyme activity, which are the focus of this addendum, some additional information is warranted. First, the functional identity of these genes was verified by heterologous expression of the genes coupled with an \textit{in vitro} enzyme assay and by biochemical characterization of genetic knock-out mutants.\textsuperscript{12,27-29} Second, one gene family consists of 5 genes/enzymes (GALT2-At4g21060, GALT3-At3g06440, GALT4-At1g27120, GALT5-At1g74800, and GALT6-At5g62620) each of which contains both a GALT domain as well as a GALECTIN domain, while the other family consists of 3 genes/enzymes (HPGT1-At5g53340, HPGT2-At4g32120, and HPGT3-At2g25300), each of which contains only a GALT domain.\textsuperscript{12,27} Third, this paper presents the first direct comparison between members of these 2 related gene families in terms of their biochemical activity and mutant phenotypes.

\textbf{Results and discussion}

The importance of AGPs is reflected in their wide range of physiological functions related to vegetative processes, sexual reproduction, development, and signaling throughout the plant kingdom. Given the complexity and heterogeneity of the numerous AGPs, little is known about the contribution of AG polysaccharides to AGP function. Identification of the genes responsible for AGP glycosylation and their genetic mutants, however, is beginning to reveal such contributions.

\textit{In vitro} GALT assays and functional analysis led to the identification of 8 Hyp-GALTs within the GT31 family.\textsuperscript{12,27,28} It is interesting that one subclade consists of 5 proteins (named GALT2-6) each with 2 protein domains, a catalytic GALT domain and a GALECTIN domain, whereas the other subclade consists of 3 proteins (named HPGT1-3), each of which only contains a GALT domain (Fig. 1).

Based on genotyping and phenotypic analysis of single, double, and triple T-DNA insertion mutants for the GALT and HPGT genes, these lines have a number of phenotypic alterations compared to wild-type.\textsuperscript{12,27} In particular, here we directly examined and compared the \textit{galt2galt5} double mutant with the \textit{hpgt1hpgt2hpgt3} triple mutant. The \textit{galt} double and \textit{hpgt} triple knockout mutants displayed 43% and 70% reductions in $\beta$-Gal-Yariv precipitable AGPs compared to wild type, respectively. In addition, the double and triple mutants showed a 31% and 18% decrease in Hyp-galactosylation activity compared to wild type using our \textit{in vitro} enzyme assay, respectively (Fig. 2). These results indicate that both subclades affect AGPs and that a greater reduction is observed when 3 genes were knocked out as opposed to two. However, it is not known whether this greater reduction is due to the greater effect of the HPGT genes or to the greater number of genes that were knocked out (i.e., 3 versus 2). It is also interesting to note that while both the double and triple mutants show a decrease in Hyp-GALT activity using our \textit{in vitro} assay, the decrease is not proportional to the reductions in $\beta$-Gal-Yariv precipitable AGPs observed in these mutants. One likely explanation for this is that HPGTs may have a different AGP peptide substrate preference compared to the GALTs.

The reduced AGP glycosylation and reduced Hyp-GALT activity observed in the \textit{galt} and \textit{hpgt} single and higher order mutants were also reflected in their physiological phenotypes.\textsuperscript{12,27} Both the \textit{galt} and the \textit{hpgt} mutants demonstrated pleiotropic effects, which were generally subtle or non-existent in the single mutants but were more severe in the \textit{galt} double and \textit{hpgt} triple mutants. The \textit{galt} mutants demonstrated reduced root hair growth, reduced seed coat mucilage, reduced seed set, and accelerated leaf senescence. The \textit{hpgt} single mutants showed no obvious phenotypes, the \textit{hpgt} triple mutants exhibited the pleotropic phenotypes of longer lateral roots, longer and denser root hairs, swollen root tips, smaller leaves, shorter inflorescence stems, reduced
fertility, and shorter siliques. Thus, while the \textit{galt} and \textit{hpgt} mutant phenotypes share some similarities, there are also some differences, which may reflect the action of the GALTs and HPGTs on different sets of AGPs. Regardless of these differences, these results provide clear evidence that Hyp-linked AG polysaccharide chains of AGPs, which are initiated by the action of the GALTs and HPGTs, are essential for AGP function.

Finally, identification of these 8 Hyp-GALTs raises the following questions that should be addressed in future studies. 1. Do these 8 enzymes act on all AGPs or only on a subset of AGPs? 2. Do these enzymes act in a

Figure 1. \textit{In silico} analysis of Hyp-GALTs for AGPs. (A) Phylogenetic tree of the Arabidopsis Hyp-GALT members which act on AGPs. The glycosyltransferase sequences were retrieved from the Carbohydrate-Active enZymes (CAZY) database (http://www.cazy.org/GT1_eukaryota.html) and NCBI database. The full-length amino acid sequences of the Arabidopsis Hyp-GALTs were aligned with Clustal Omega and the maximum parsimony tree was generated using MEGA 6. Numbers at the nodes indicate bootstrap values calculated for 1000 replicates (>50%). The scale bar represents 50 amino acid substitutions. (B) Protein structure of the 8 Hyp-GALT genes found in Arabidopsis. Blue arrowheads indicate the position of the DXD motif, which is predicted for UDP-galactose binding. TMHMM (http://www.cbs.dtu.dk/services/TMHMM/) was used to predict the transmembrane domain (T); Pfam domain predictions: Pf01762 identified the Galactosyltransferase (GALT) domain (http://www.sanger.ac.uk/Software/Pfam/) and Pf00337 identified the Galactose-binding lectin (GALECTIN) domain.
complex as is apparently the case for biosynthesis of the \( \beta-1,6 \)-galactan side chains by GALT31A-GALT29A\(^3\) 

3. Do these enzymes have different preferences for AGP peptide sequences? 4. Are there any other AGP-specific Hyp-GALTs/HPGTs that remain to be identified and characterized? Nonetheless, based upon the progress to date, we are now beginning to understand how the Hyp-AG polysaccharides of AGPs are made and their important contributions to biological function.

**Abbreviations**

AG  arabinogalactan  
AGPs  arabinogalactan-proteins  
HPGT  hydroxyproline galactosyltransferase  
Hyp  hydroxyproline  
GALT  galactosyltransferase  
GTs  glycosyltransferases

**Disclosure of potential conflicts of interest**

No potential conflicts of interest were disclosed.

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