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Xylan synthesized by Irregular Xylem 14 (IRX14) maintains the structure of seed coat mucilage in Arabidopsis

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Received 30 August 2015; Accepted 6 November 2015

Editor: Simon Turner, University of Manchester

Abstract

During differentiation, the Arabidopsis seed coat epidermal cells synthesize and secrete large quantities of pectinaceous mucilage into the apoplast, which is then released to encapsulate the seed upon imbibition. In this study, we showed that mutation in Irregular Xylem 14 (IRX14) led to a mucilage cohesiveness defect due to a reduced xylan content. Expression of IRX14 was detected specifically in the seed coat epidermal cells, reaching peak expression at 13 days post-anthesis (DPA) when the accumulation of mucilage polysaccharides has ceased. Sectioning of the irx14-1 seed coat revealed no visible structural change in mucilage secretory cell morphology. Although the total amount of mucilage was comparable with the wild type (WT), the partition between water-soluble and adherent layers was significantly altered in irx14-1, with redistribution from the adherent layer to the water-soluble layer. The monosaccharide composition analysis revealed that xylose content was significantly reduced in irx14-1 water-soluble and adherent mucilage compared with the WT. The macromolecular characteristics of the water-soluble mucilage were modified in irx14-1 with a loss of the larger polymeric components. In accordance, glycome profiling and dot immunoblotting of seed mucilage using antibodies specific for rhamnogalacturonan I (RG I) and xylan confirmed the ultra-structural alterations in the irx14-1 mucilage. Meanwhile, the crystalline cellulose content was reduced in the irx14-1 mucilage. These results demonstrated that IRX14 was required for the biosynthesis of seed mucilage xylan, which plays an essential role in maintaining mucilage architecture potentially through altering the crystallization and organization of cellulose.

Key words: Arabidopsis, crystalline cellulose, IRX14, mucilage, xylan.

Introduction

In myxospermous species such as Arabidopsis thaliana, seed coat mucilage secretory cells (MSCs) synthesize and deposit large amounts of pectinaceous mucilage in the apoplast (Western et al., 2000; Windsor et al., 2000). Following mucilage synthesis, a secondary cell wall is synthesized and deposited, forming a volcano-shaped columella in the centre of the cell (Western et al., 2000; Windsor et al., 2000). At seed maturity, the epidermal cells have the central columella connected to reinforced radial cell walls and surrounded by doughnut-shaped mucilage under a primary cell wall.
Hydration of mature seeds leads to an almost instantaneous mucilage release forming a gelatinous capsule surrounding the seed (Western et al., 2000; Windsor et al., 2000). The MSCs of Arabidopsis have been presented as an ideal model system for the study of the biosynthesis and modification of cell wall components (Arsovski et al., 2010; Haughn and Western, 2012; Western, 2012; Francoz et al., 2015; Voiniciuc et al., 2015b).

Staining of Arabidopsis mucilage with the peptate staining dye ruthenium red revealed the presence of two distinct layers: an outer, water-soluble layer and an inner, adherent layer tightly attached to the seed surface (Western et al., 2000; Windsor et al., 2000). Both layers have been demonstrated to be composed primarily of unbranched hemicelluluronan I (RG I) (Western et al., 2000; Windsor et al., 2000; Macquet et al., 2007). A minor amount of homogalacturonan (HG) is also present within the adherent mucilage (Willats et al., 2001), and its degree of methyl esterification varies, with a higher degree of variation in the outer layer than in the inner layer (Macquet et al., 2007).

Apart from pectin, immunochemical analysis revealed that a small amount of diffuse cellulose was also present within the mucilage. Cellulose was interspersed with rays extending from the tip of the columnella at the seed surface across the inner layer (Western et al., 2000; Windsor et al., 2000; Macquet et al., 2007; Stork et al., 2010). The presence of cellulose in the inner layer has been proved to be essential for mucilage adherence (Harpaz-Saad et al., 2011; Mendu et al., 2011a; Sullivan et al., 2011). For example, CELULOSE SYNTHASE5 (CESA5)/MUM3 was shown to be responsible for the synthesis and proper deposition of cellulose in the inner adherent domain (Sullivan et al., 2011). cesa5/mum3 exhibited increased water-soluble mucilage accompanied by decreased adherent mucilage, suggesting a structural role of cellulose in anchoring the pectinaceous mucilage to the seed (Harpaz-Saad et al., 2011; Mendu et al., 2011a; Sullivan et al., 2011). In addition, linkage analysis revealed small quantities of hemicellulose in the mucilage (Huang et al., 2011; Walker et al., 2011; Kong et al., 2013; Yu et al., 2014), which may cross-link with the cellulose in the mucilage similar to its proposed role in cell walls (Haughn and Western, 2012; Yu et al., 2014; Voiniciuc et al., 2015a, b). For example, glucomannans synthesized by CELULOSE SYNTHASE-LIKE A2 (CSLA2) were proposed to modulate the structure of adherent mucilage, potentially through altering cellulose organization and crystallization (Yu et al., 2014). Moreover, MUCI10 was responsible for the galactosylation of glucomannans synthesized by CSLA2 and affected mucilage architecture along with celluloseic rays (Voiniciuc et al., 2015a).

Xylans are the major hemicellulose that cross-link with cellulose in the secondary walls of dicots and in both primary and secondary walls in commelinid monocots (Scheller and Ulvskov, 2010; Rennie and Scheller, 2014). Xylans are composed of a linear backbone of a β-1,4-linked d-xylose (Xyl) residues, which can be diversely substituted with arabinose (Ara), glucuronic acid (GlcA) and 4-O-methyl glucuronic acid (MeGlcA) depending on the species and tissue types (Ebringerová and Heinze, 2000). Based on the side chain substitutions, xylans are generally classified as gluconoxylan (GX), the major hemicellulose in dicots, arabinoxylan (AX) and gluconoroarabinoxylan (GAX), the most abundant hemicelluloses in grasses (Ebringerová and Heinze, 2000).

Previous studies using reverse genetics approaches have revealed that several glycosyltransferases (GT) were involved in xylan biosynthesis in Arabidopsis. For instance, IRX9/IRX9L and IRX14/IRX14L from the GT43 family, and IRX10/IRX10L from the GT47 family are required for the elongation of the xylan backbone (Brown et al., 2007, 2009; Wu et al., 2009, 2010; Kepler and Showalter, 2010; Lee et al., 2010). IRX9, IRX14, and IRX10 have been proved to play major roles in xylan backbone biosynthesis, with their close homologues IRX9L, IRX14L, and IRX10L showing partially redundant roles (Wu et al., 2009, 2010; Kepler and Showalter, 2010; Lee et al., 2010). Loss of function of IRX9, IRX14, or IRX10 resulted in decreased xylan synthase activity, reduced xylan content, and xylan with short backbones. Mutations for each of these gene pairs (IRX9/IRX9L, IRX10/IRX10L, or IRX14/IRX14L) significantly enhanced the phenotype of the single mutants (Wu et al., 2009, 2010; Kepler and Showalter, 2010; Lee et al., 2010). Interestingly, a recent study indicated that these three gene pairs play equally important roles rather than major and minor roles in xylan biosynthesis. IRX9L, IRX10L, and IRX14 are mainly responsible for the synthesis of xylan in primary cell walls, whereas IRX9, IRX10, and IRX14L predominantly synthesize xylan in secondary cell walls (Mortimer et al., 2015). In addition, IRX7 (FRA8)/IRX7L (F8H) from the GT47 family, IRX8 (GAUT12) and PARVUS from the GT8 family are responsible for the synthesis of the oligosaccharide at the reducing end of xylans (Brown et al., 2007; Lee et al., 2007, 2009; Pena et al., 2007). Mutations in IRX7, IRX8 or PARVUS led to a decreased xylan content and the lack of the reducing end oligosaccharide sequence, while the xylan backbone elongation activity was retained (Lee et al., 2007, 2009; Pena et al., 2007).

As a major hemicellulose component in cell walls, xylan biosynthesis has been extensively studied in secondary cell wall formation. However, no report on the function and biosynthesis of xylan in seed coat mucilage has been found to date. In this study, we showed that IRX14 was essential for the biosynthesis of xylan in mucilage and played important roles in maintaining normal mucilage structure in the seed coat. Although irx14-1 produced normal amounts of mucilage, the cohesive properties of the mucilage were significantly altered resulting from the fact that most of the pectinaceous components of the inner adherent mucilage layer were mal-distributed into the water-soluble layer. In addition, the macromolecular characteristics of the water-soluble mucilage in irx14-1 were dramatically altered, and the crystalline cellulose content was significantly reduced in irx14-1 mucilage. Our results demonstrate that xylan synthesized by IRX14 functioned in maintaining the proper structure of mucilage potentially through its interaction with cellulose.
Materials and methods

Plant material and growth conditions

T-DNA insertion lines for irx14-1 (SALK_038212), irx14-2 (CS400996), irx9-1 (SALK_058238), irx14-1l (SALK_066961), and irx9-1l (SALK_037323) were obtained from ABRC (http://www.arabidopsis.org). Homozygous lines were identified by genotyping using the primers indicated in Supplementary Table S1 at JXB online. Seeds were surface-sterilized and sown on to half-strength Murashige and Skoog (1/2 MS) medium, stratified in the dark for 2 d at 4 °C and germinated at 21 °C under a 16/8 h day/night photoperiod. Ten-day-old plants were transferred to soil in growth chambers under the same conditions.

Ruthenium red staining of seed mucilage

Seeds were imbibed for 2 h in water with or without shaking, then stained in 0.01% (w/v) ruthenium red (Sigma-Aldrich) for 30 min at room temperature. Following a brief rinse with de-ionized water, seeds were photographed with a BX51 light microscope (OLYMPUS).

Microscopy and histology

For resin embedding and sectioning, developing seeds staged at 4, 7, 10, and 13 DPA were fixed in 2.5% (w/v) glutaraldehyde in 0.1 M phosphate-buffered saline (PBS) (pH 7.0) overnight at 4 °C. After washing, samples were post-fixed for 1 h in 1% (v/v) osmium tetroxide in PBS, dehydrated through a gradient ethanol series and subsequently embedded in Spurr’s resin. Sections (1 µm) were cut and stained with 1% toluidine blue O dissolved in 0.1 M PBS and washed with de-ionized water. Images were captured with a BX51 microscope (OLYMPUS).

For Scanning Electron Microscopy (SEM), mature dry seeds were mounted on stubs, coated with platinum in an E1045 ion sputter coater (Hitachi), and imaged using a S4800 scanning electron microscopy (Hitachi) with an accelerating voltage of 20 kV.

For polarized light microscopy, seeds were imbibed in water for 30 min before being mounted on a glass slide. Seed birefringence was observed with a Nikon Eclipse E600 POL microscope.

Mucilage extraction

Three independent samples of 100 mg seeds were extracted sequentially with water and 2 M NaOH (with 3 mg ml⁻¹ NaBH₄) for 1 h each with shaking on an orbital shaker at room temperature (Western et al., 2001). The resulting suspension was centrifuged to pool supernatants, and 2 M NaOH extracts were neutralized with acetic acid, then dialysed extensively against running water for 36 h before being freeze-dried.

Determination of monosaccharide composition

Two mg of water-soluble and adherent mucilage was hydrolysed with 1 ml of 2 M trifluoroacetic acid (TFA) for 2 h at 121 °C. The TFA was evaporated under a stream of nitrogen. The hydrolysates were derivatized with 1-phenyl-3-methyl-5-pyrazolone and 0.3 M NaOH at 70 °C for 30 min, extracted with chloroform three times and then analysed on a Hypersil ODS-2 C18 column (4.6×250 mm; Thermo Scientific) using a Waters HPLC System. The derivatives were eluted with 82% (v/v) PBS (0.1 M, pH 7.0) and 18% (v/v) acetonitrile at 1 ml min⁻¹, and monitored at 245 nm. Neutral sugar standards (fucose, arabinose, rhamnose, galactose, glucose, mannose, and xylose) and an acid sugar standard (galacturonic acid) were included in the analysis.

FTIR spectroscopy

Water- and 2 M NaOH-extracted mucilage was used for FTIR analysis. Samples were prepared by dissolving 2 mg mucilage with 20 mg KBr in 50 µl water. The mixture was incubated overnight at 50 °C. Pellets were made by mixing 2 mg mucilage-KBr mixture with 23 mg KBr, and pressed into 7 mm films. Spectra were collected using a Thermo Nicolet Nexus 470 spectrometer (Thermo Scientific) over the range of 4 000-800 cm⁻¹. For each spectrum, 200 scans were performed at 8 cm⁻¹ resolution. Subtraction plots between WT and irx14-1 were generated using OMNIC software (Thermo Nicolet) and spectral differences were cross-referenced to identify peaks linked to pectin, xylan, and cellulose (Kacuráčová et al., 2000). Only the spectra between 1 900 and 800 cm⁻¹ were used for analysis.

Macromolecular characterization

High-performance size-exclusion chromatography (HP-SEC) was performed on a system comprising of tandemly connected Shodex OHpak SB-802 and SB-806 columns, coupled with a differential refractive index detector (Wyatt Optilab REX). Elution was carried out with 50 mM NaNO₃ at a flow rate of 1 ml min⁻¹. Samples were solubilized in 50 mM NaNO₃ and filtered through disposable syringe filters (0.22 µm, Millipore) before injection. Samples were injected automatically through a 0.5 ml loop in an auto-sampler. Data analysis was performed using Astra software.

For electrophoresis of mucilage polymers, extracted water-soluble and adherent mucilage was re-suspended in water at a concentration of 0.5 mg ml⁻¹, and 200 µl was loaded in each lane. Electrophoresis was carried out in a 0.7% agarose gel prepared in TAE buffer at 60 V (6 cm⁻¹) for 2.5 h. After electrophoresis, the gel was stained in 0.01% (w/v) ruthenium red (Sigma-Aldrich) solution for 30 min and then photographed.

Glycosyl linkage analysis

Total mucilage was extracted by vigorously shaking in 50 mM EDTA using a Retsch MM300 TissueLyser (Qiagen). The reduction of the uronic acids to their respective 6,6-dideuterio derivatives was carried out as described by Gibeaut and Carpita (1991). After reduction, mucilage samples were extensively dialysed against running water and lyophilized, then solubilized in 200 µl anhydrous DMSO. Methylation was performed as described by Yu et al. (2014). The methylated polymers purified by chloroform extraction were hydrolysed in 2 M TFA. The sugars were then reduced with NaBD₄ and acetylated. Partially methylated alditol acetates were dissolved in methylene chloride and analysed by GC–MS as described previously (Yu et al., 2014).

Seed cell wall preparation

Whole dry seeds and seeds lacking mucilage were frozen in liquid nitrogen and ground to a fine powder using a Retsch MM300 TissueLyser (Qiagen, Germany). Ten mg of ground seeds were de-starched with a total starch assay kit (Megazyme) according to the manufacturer’s protocol, then washed twice with 2 ml of 80% (v/v) ethanol, and centrifuged at 5 000 rpm for 10 min. The ethanol was removed and the pellet was washed sequentially with 95% (v/v) ethanol, 100% ethanol, and twice with 100% acetone. After washing, the samples were dried under a vacuum at 60 °C to obtain the alcohol-insoluble residue (AIR).

Crystalline cellulose quantification

For the determination of the crystalline cellulose content, 20 mg AIR was hydrolysed in 2 ml of 2 M TFA at 121 °C for 2 h. After centrifugation, the pellets were suspended in 2 ml of Updegraff reagent [acetic acid:nitric acid:water, 8:1:2 by vol.] (Updegraff, 1969), and boiled at 100 °C for 1 h. The crystalline cellulose pellets were collected by centrifugation at 10 000 rpm for 10 min, and hydrolysed in 3 ml of 72% (w/v) H₂SO₄ for 30 min. The amounts of crystalline cellulose were quantified colorimetrically using the anthrone assay (Laurentin and Edwards, 2003).
Immunolabelling of mucilage

Whole seed immunolabelling was performed with two xylan-specific antibodies CCCR-M139 (Pattathil et al., 2010) and LM11 (McCartynew et al., 2005), and the carbohydrate-binding module CBM3a that binds to crystalline cellulose (Dagel et al., 2011). Whole seeds were first blocked with PBS containing 3% (w/v) fat-free milk powder (MP/PBS) for 30 min, and then incubated in 10-fold diluted primary antibodies in MP/PBS for 1.5 h. Seeds were then washed with PBS and incubated with a 200-fold dilution of AlexaFluor488-tagged goat anti-mouse (CCRC series) or anti-rat (LM11) IgG in MP/PBS in darkness for 1 h. Alternatively, seeds were incubated in MP/PBS containing 10 mg ml\(^{-1}\) CBM3a for 1.5 h, washed in PBS, and incubated in a 100-fold dilution of mouse anti-His monoclonal antibody for 1 h. After washing with PBS, seeds were incubated in 200-fold diluted AlexaFluor488-tagged goat anti-mouse IgG in MP/PBS in darkness for 1 h. After incubation, seeds were washed in PBS and counterstained for 5 min with Calcofluor White (Sigma-Aldrich). Images were captured with a Fluoview FV1000 confocal laser scanning microscope (OLYMPUS) using 405 nm and 488 nm lasers.

Dot immunoblotting assays

Water and 2 M NaOH mucilage extracts were re-suspended in water. A series of dilutions were prepared and a 1 μl aliquot was spotted on to a nitrocellulose membrane (Merck Millipore). After being air-dried, the membrane was blocked for 1 h in MP/PBS, then incubated for 1.5 h in a 10-fold dilution of primary antibodies. After washing three times with PBS, membranes were incubated for 1.5 h in horseradish peroxidase (HRP)-conjugated anti-rat (for LM series) or anti-mouse (for CCRC series) secondary antibodies in a 1000-fold dilution of MP/PBS. Membranes were washed prior to colour development in substrate solution (25 ml de-ionized water, 5 ml methanol containing 10 mg ml\(^{-1}\) 4-chloro-1-naphthol and 30 μl 6% (v/v) H\(_2\)O\(_2\)). After incubation for 30 min at room temperature, the blots were rinsed with de-ionized water and photographed.

ELISA

Mucilage extractions (100 μg ml\(^{-1}\)) in PBS were coated overnight at 4 °C on to 96-well microtitre plates (Costa 3599). Coating solutions were removed and 200 μl MP/PBS was added to each well. After blocking overnight at 4 °C, plates were washed and 100 μl primary antibodies at a 1/20 dilution in MP/PBS were added to each well. After incubation at 37 °C for 2 h, plates were washed with PBS and incubated with HRP-conjugated anti-rat (JIM and LM series) or anti-mouse (CCRC series) secondary antibodies at a 1 000-fold dilution in MP/PBS. Membranes were washed prior to colour development in substrate solution (25 ml de-ionized water, 5 ml methanol containing 10 mg ml\(^{-1}\) 4-chloro-1-naphthol and 30 μl 6% (v/v) H\(_2\)O\(_2\)). After incubation for 30 min at room temperature, the blots were rinsed with de-ionized water and photographed.

Quantitative Real-Time RT-PCR (qRT-PCR) analysis

Total RNA was isolated from developing siliques using the CTAB method. Contaminating DNA was removed using RQ1 RNase-Free DNase I (Promega), and first-strand cDNA was synthesized with SuperScript III reverse transcriptase (Invitrogen) according to the manufacturer’s instructions. qRT-PCR was performed using SYBR Premix Ex Taq (TaKaRa) in the LightCycler 480 detection system (Roche). The analysis was performed using three technical replicates. Relative expression was calculated by the 2\(^{-\Delta\Delta Ct}\) method (Livak and Schmittgen, 2001) utilizing ACTIN5 as the internal control. Gene-specific and reference gene primers are listed in Supplementary Table S1.
have been shown to act redundantly in xylan biosynthesis (Lee et al., 2010; Wu et al., 2010). The expression pattern of these GT43 family genes during seed development was examined using the LCM seed microarray datasets via the Bio-Array Resource eFP browser (Winter et al., 2007). All these four genes exhibited seed coat-specific expression during seed development (Supplementary Fig. S2). The expression of IRX9 was preferentially expressed at the bending cotyledon and mature green stage, which is similar to that of IRX14, while IRX9L and IRX14L were expressed in seed coats throughout the course of seed development, albeit with no large fluctuations. The seed coat-specific expression patterns of IRX14 and its homologues prompted us to investigate their potential roles in mucilage biosynthesis and/or modification further.

**The mucilage layer is thinner in irx14 mutants compared with the WT**

To investigate the role of IRX14 and its homologues in seed coat mucilage, we isolated irx14, irx14l, irx9, and irx9l homozygous insertion mutants. Ruthenium red staining of homozygous seeds showed that two irx14 mutants (irx14-l and irx14-2) exhibited significant mucilage defects with a thinner mucilage halo compared with the WT (Fig. 2). Direct ruthenium red staining of the irx14 (irx14-1 and irx14-2) seeds without shaking showed a much thinner layer of adherent mucilage compared with the WT. When stained with ruthenium red after shaking in water, only a tiny amount of the inner mucilage layer remained attached to the irx14 seeds. By contrast, both irx9l and irx14l exhibited similar staining patterns.

**Fig. 1.** Expression of IRX14 in developing seeds. (A) qRT-PCR analysis of IRX14 expression during seed coat development staged at 4, 7, 10, 13, and 16 DPA. Error bars represent SD (n=3). The data are presented as relative fold change, where the IRX14 expression level at 4 DPA is set at 1.0. (B–F) In situ hybridization using an IRX14-specific antisense probe. (G) In situ hybridization using an IRX14-specific sense probe in the seed coat of developing seeds. Arrowheads indicate the hybridization signals. C, Columella; RW, radial cell wall; SG, starch granule. Bar=20 μm.
patterns of mucilage to the WT (Fig. 2). Unfortunately, the mucilage phenotype was not obtained for irx9 seeds because the growth of the irx9-1 mutant allele we used was severely retarded and no seeds were available under our growth conditions. In addition, the seed mucilage phenotypes of the irx14 irx14l double mutant could not be obtained due to its severely retarded growth, dramatically dwarfed plant stature, and severe sterility. Since two mutant alleles of irx14 have similar mucilage defect phenotype, we used irx14-1 for further analysis in subsequent experiments.

Seed coat structure and mucilage biosynthesis are unaffected in irx14-1

To determine whether mutation in IRX14 affects mucilage biosynthesis or seed coat morphology during seed coat development and differentiation, seed coat epidermal cells from irx14-1 and the WT staged from 4–13 DAP were sequentially sectioned, stained with toluidine blue O, and examined by light microscopy. As shown in Supplementary Fig. S3, no obvious differences were observed in the timing or process of differentiation between irx14-1 and WT seeds. Mucilage production in irx14-1 seed coat epidermal cells appeared similar to that of the WT. Furthermore, cytoplasmic rearrangement and columella formation also occurred normally in the mutant epidermal cells. The structure of irx14-1 seed coat epidermal cells was examined in more detail by Scanning Electron Microscopy (SEM) on mature dry seeds. The epidermal cells of WT and irx14-1 seeds exhibited the characteristic polygonal shapes with a central columella and were equivalent in size, indicating no significant changes in the outer and radial cell walls in irx14-1 seeds compared with the WT. These results indicate that seed coat structure and mucilage biosynthesis were unaffected in irx14-1.

Water-soluble mucilage is increased in irx14 seeds

To determine the changes in the water-soluble and adherent mucilage between irx14-1 and the WT, seed mucilage was sequentially extracted with water and 2M NaOH. As shown in Fig. 3A, an increase of 14% in the amount of water-soluble mucilage was observed for irx14-1 compared with the WT. Reciprocally, the amount of adherent mucilage was decreased by 3% in irx14-1 compared with the WT. Nevertheless, there was no significant difference for the total amount of mucilage between irx14-1 (10.38 ± 0.52) and the WT (9.87 ± 0.44). These results indicated that the IRX14 mutation led to an increase in the level of water-soluble mucilage with a concomitant decrease in the adherent mucilage. The mal-distribution of mucilage between the outer and inner layers in irx14-1 implies possible alterations of mucilage composition and structure.

Xylose content is reduced in irx14-1 mucilage

Considering the differences in mucilage extraction profiles, we further quantified the monosaccharide composition of irx14-1 and WT mucilage using HPLC. A significant decrease (67–87%) was observed for Xyl content in both water-soluble and adherent mucilage in irx14-1 (Fig. 3B, C). Meanwhile, mannose (Man) and Ara contents were significantly increased in the water-soluble and adherent mucilage of irx14-1, respectively. The increased levels of Ara and Man may result from the balancing effect among different hemicellulose components to counteract the decrease in xylose content. Furthermore, an up to 20% increase in total sugars was observed in irx14-1 water-soluble mucilage, which was mainly due to the elevated levels in rhamnose (Rha) and galacturonic acid (GalA), indicating that irx14-1 contained more water-soluble mucilage than the WT. By contrast, total sugars in adherent mucilage decreased by approximately 15% in irx14-1, also mainly due to reductions in Rha and GalA contents. This further confirms that the increased levels of sugars in water-soluble mucilage are caused by the re-distribution of some mucilage from the adherent layer to the water-soluble layer in irx14-1.

FTIR reveals altered polysaccharide structures in irx14-1 mucilage

Fourier Transform Infrared Spectroscopy (FTIR) has been proved to be an informative diagnostic tool in detecting subtle
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changes in cell wall carbohydrate structures based on signature peaks (Kacuráková et al., 2000; Dean et al., 2007; Mendu et al., 2011b). To reveal the differences of composition between *irx14-1* and WT mucilage, we generated FTIR spectra for *irx14-1* and WT mucilage and subtraction spectra representing proportional differences between them (Supplementary Fig. S4). Within the FTIR subtraction plot, different peaks in RG I (1 017, 1 122, 1 150, 1 423, and 1 621 cm⁻¹), xylan (978 cm⁻¹) and cellulose (898 and 1 034 cm⁻¹) were identified between WT and *irx14-1* spectra. These results corroborate the above monosaccharide composition analysis and further support that the composition of mucilage was altered in *irx14-1*.

Macromolecular characteristics are altered in *irx14-1* soluble mucilage

To explore whether the *irx14-1* mucilage phenotype is caused by a change in the size of the mucilage polysaccharides, high-performance size-exclusion chromatography (HP-SEC) combined with a refractive index detector and a multi-angle light-scattering detection was used to determine the macro-molecular parameters of *irx14-1* and WT mucilage. Wild type water-soluble mucilage had an elution profile that separated two polymeric populations with different molar mass (Fig. 4). The first polymeric population was of an extremely high
average molar mass (Mn approximately 30 000 kDa), while the second one was of lower average molar mass (Mn approximately 750 kDa). However, the macromolecular characteristics of the irx14-1 water-soluble mucilage exhibited major differences from that of the WT. The higher molar mass population was totally absent in irx14-1 water-soluble mucilage, and the second population eluted slightly later compared with the WT. By contrast, no significant difference for macromolecular characteristics was observed in 2 M NaOH extracted mucilage between irx14-1 and the WT.

Furthermore, we used gel electrophoresis to discern possible changes in size and/or charge in irx14-1 mucilage. Although no significant difference was observed for adherent mucilage between irx14-1 and the WT, the water-soluble mucilage displayed distinct staining patterns (Supplementary Fig. S5). The WT mucilage was distributed in a large smear in the gel lane, with much of the mucilage still remaining in the loading well. By contrast, the mucilage from irx14-1 was mostly aggregated into a band with no material remaining in the loading well. These results indicate that the macromolecular characteristics of the water-soluble mucilage is altered in irx14-1.

IRX14 is essential for xylan synthesis in mucilage

To determine if subtle changes in the chemical structure of mucilage are present in irx14-1, we analysed the glycosyl linkage composition of total mucilage extracts (Table 1). Similar to the monosaccharide composition analysis (Fig. 3B, C), most of the composition of polysaccharides in irx14-1 was comparable with that of WT. Consistent with previous studies (Huang et al., 2011; Walker et al., 2011; Kong et al., 2013; Yu et al., 2014), the mucilage in irx14-1 and the WT was primarily composed of 2-Rha and 4-GalA, representing a relatively unbranched RG I backbone. The degree of branching of the RG I, as determined by the ratio of 2,4-Rha to 2-Rha, remained almost constant in irx14-1 and the WT. An obvious change observed was a dramatic decrease in the xylan content in irx14-1 compared with the WT (Table 1). Lower xylan content in irx14-1 mucilage resulted from the severe reduction of 4-Xyl and t-Xyl. This indicated that IRX14 is essential for the synthesis of the xylan backbone in mucilage.

Crystalline cellulose is reduced in irx14-1 mucilage

Previous studies indicated that cellulose plays an important role in maintaining mucilage adherence (Harpaz-Saad et al., 2011; Mendu et al., 2011; Sullivan et al., 2011). To determine the change in cellulose content in irx14-1 mucilage, irx14-1 and WT seeds imbibed in water were observed for birefringence produced by crystalline cellulose under polarized light (Fig. 5A, B). The outer epidermal cells of WT seeds exhibited strong birefringence with visible rays of crystalline cellulose within the adherent mucilage. By contrast, irx14-1 displayed much less birefringence under polarized light, indicating that the crystalline cellulose content was dramatically reduced in irx14-1 mucilage. Furthermore, quantification of crystalline cellulose contents in mucilage, de-mucilaged seeds, and whole seeds revealed that irx14-1 mucilage had a 20% reduction in crystalline cellulose compared with the WT (Fig. 5C).

The structure of mucilage is altered in irx14-1

To examine further the structure and composition of mucilage in irx14-1, we performed whole mount immunolabelling on mature dry seeds. The carbohydrate-binding module CBM3a, which binds to crystalline cellulose (Dagel et al., 2011), in parallel with Calcofluor, a fluorescent probe for β-glycans, was used to examine the spatial distribution of cellulose in the mucilage (Fig. 6). Calcofluor labelling of WT mucilage revealed intense rays radiating from the tops of the columella and across the inner layer of the seed mucilage, whereas labelling of Calcofluor was only detectable in mucilage around the columella in irx14-1, and the diffuse staining of rays was almost completely absent. CBM3a displayed a similar staining pattern to that of Calcofluor in WT adherent mucilage, with intense labelling of
Table 1. Sugar linkage composition of total mucilage extracted from irx14-1 and WT seeds

| Sugar and linkage | WT | irx14-1 |
|------------------|----|---------|
| Pharnnose        | 2.10 | 2.26 |
| 1-Rhap           | 33.25 | 36.78 |
| 2-Rhap           | 2.69 | 2.23 |
| 2,4-Rhap         | 6.01 | 4.97 |
| Arabinose        | 0.27 | 0.24 |
| 1-Araf           | 0.62 | 0.69 |
| 3-Araf           | 0.50 | 0.51 |
| 2,5-Araf         | 0.34 | 0.43 |
| Xylose           | 0.46 | 0.04 |
| 1-Xylp           | 1.97 | 0.21 |
| 4-Xylp           | 0.28 | 0.19 |
| 3,4-Xylp         | 0.21 | 0.09 |
| Mannose          | 0.72 | 0.55 |
| 1-Manp           | 2.54 | 2.74 |
| 4-Manp           | 1.73 | 2.76 |
| Galactose        | 0.59 | 0.62 |
| 1-Galp           | 0.22 | 0.17 |
| 3-Galp           | 0.16 | 0.14 |
| 6-Galp           | 0.08 | 0.09 |
| 3,4-Galp         | 0.16 | 0.30 |
| 3,6-Galp         | 0.30 | 0.13 |
| Glucose          | 0.22 | 0.14 |
| 1-Glcp           | 1.37 | 1.14 |
| 4-Glcp           | 0.24 | 0.24 |
| 2,4-Glcp         | 0.59 | 0.48 |
| Galacturonic acid| 5.55 | 5.47 |
| 1-GalAp          | 35.23 | 34.01 |
| 3,4-GalAp        | 1.59 | 3.39 |

Total mucilage was extracted by vigorous shaking in 50 mM EDTA. The results are given as the mean molar percentage of two independent assays, with variance less than 5% for all linkage groups.

the entire adherent mucilage halo and stronger labelling at the outer periphery. In irx14-1 seeds, CBM3a labelling was only detectable for columella and cell wall debris attached to the seed surface, consistent with the loss of mucilage adherence.

Since IRX14 was previously reported to be involved in the biosynthesis of the xylan backbone (Keppler and Showalter, 2010; Lee et al., 2010; Wu et al., 2010), we examined the distribution of xylan epitopes in mucilage using two xylan-specific antibodies, CCRC-M139 and LM11 (McCartney et al., 2005; Pattathil et al., 2010), in conjunction with Calcofluor staining (Fig. 7). The labelling of CCRC-M139 and LM11 showed similar patterns to Calcofluor in WT adherent mucilage. By comparison, CCRC-M139 and LM11 labelling was no longer observed in the adherent mucilage in irx14-1, and signals were only detected in the columella and remnant wall fragments attached to the seed surface. These results indicate that the distribution of cellulose and xylan in adherent mucilage in irx14-1 is significantly disrupted.

To obtain more accurate information of possible changes in the structural features of xylan and RG I in irx14-1 mucilage, dot immunoblotting was performed with three antibodies specific for the RG I backbone (CCRC-M14, CCRC-M35, and CCRC-M36) (Pattathil et al., 2010) and four for xylan (CCRC-M37, CCRC-M54, CCRC-M139, and LM11) (McCartney et al., 2005; Pattathil et al., 2010). As shown in Fig. 8, the signal intensities of three antibodies specific for the RG I backbone were dramatically reduced in irx14-1 adherent mucilage. In addition, the staining patterns of these three antibodies for the RG I backbone differed significantly between irx14-1 and WT water-soluble mucilage. The labelling of CCRC-M14 in irx14-1 water-soluble mucilage seems to be more diffuse and has a larger halo than that of the WT. As for CCRC-M35 and CCRC-M36, two distinct layers of labelling were evident for the irx14-1 water-soluble fraction, whereas there was only a single layer of labelling for the WT. In addition, significant differences in signal intensities for xylan-specific antibodies were observed between irx14-1 and WT mucilage. As the signal for LM10 was hardly detectable for either WT or irx14-1 mucilage, it was excluded from the analysis. The signals of the remaining four antibodies (CCRC-M37, CCRC-M54, CCRC-M139, and LM11) were weakly detected in water-soluble mucilage in the WT, while no signals were detected in irx14-1. Although the epitopes of these four antibodies were detected in adherent mucilage in WT and irx14-1, the signal intensity was dramatically decreased in irx14-1 compared with the WT. These results further support the conclusion that substantial structural alterations in RG I and xylan components occurred in irx14-1 mucilage.

To obtain more detailed information of possible structural changes in irx14-1 mucilage, the glycome-profiling of seed mucilage was analysed by utilizing cell wall glycan-directed monoclonal antibodies (Pattathil et al., 2010). The mucilage of water and 2 M NaOH extracts was probed using 28 monoclonal antibodies that recognize epitopes on pectin (RG-I backbone and side chains, and HG), hemicelluloses (xylan, xyloglucan, and heteromannan), and extensins. The glycome profiles of irx14-1 mucilage displayed substantial differences compared with the WT (Supplementary Fig. S6), suggesting that irx14-1 mucilage has undergone substantial alterations in polysaccharide structure.

Discussion

By re-analysing the LCM seed coat microarray transcriptome data, we identified the IRX14 gene as being involved in seed coat mucilage structure maintenance. IRX14 has been demonstrated to be required for the elongation of the xylan backbone, and loss function of IRX14 results in a defect in the xylan chain length and a reduction in the xylosyltransferase activity (Keppler and Showalter, 2010; Lee et al., 2010; Wu et al., 2010). Xylan is the major hemicellulosic polysaccharide in secondary cell walls and in grass primary cell walls.
Glucuronoxylan (GX) is the major xylan in secondary cell walls of dicot plants, such as Arabidopsis and poplar. GX is composed of a linear backbone of β-(1–4)-linked d-Xyl residues, which is decorated with α-d-GlcA or 4-O-MeGlcA residues. In Arabidopsis seed coat mucilage, xylans represent approximately 3–5% (mol) in adherent mucilage as revealed by monosaccharide composition and glycosyl linkage analysis (Macquet et al., 2007; Huang et al., 2011; Walker et al., 2011; Kong et al., 2013; Yu et al., 2014). Our results and previous studies indicated that xylan in seed mucilage had a linear backbone of β-(1–4)-linked d-Xyl with branched side chains as revealed by the presence of 4-Xyl with a ratio of 1.8–13.3 to 2,4-Xyl (Huang et al., 2011; Walker et al., 2011; Kong et al., 2013; Yu et al., 2014; Voiniciuc et al., 2015b). However, GlcA or MeGlcA was only detected in trace levels or not at all in most of these studies indicating that it is unlikely to be a side chain of xylan in seed mucilage. Although the presence of xylan as a minor component in seed mucilage has been described, its functions remain to be characterized. In this study, we show that mutations of IRX14 reduced the xylan content in seed coat mucilage and resulted in increased detachment of adherent mucilage to the seed surface (Fig. 2). Although the irx14-I mutant exhibited a thinner layer of adherent mucilage, mutation of its close homologue IRX14L did not have any discernible effects on seed coat mucilage (Fig. 2). Similarly, mutations of irx14 led to the reduction in xylan content and secondary wall thickness in Arabidopsis stem and the irx14 irx14l double mutant dramatically enhanced the phenotypes of irx14 (Lee et al., 2010; Wu et al., 2010), although irx14l mutation did not cause any observable defect. These results also indicated that IRX14 is functionally dominant in xylan biosynthesis in both stem and seed mucilage.

Expression analysis revealed that the transcript of IRX14 was present throughout seed coat development with peak expression at 13 DPA (Fig. 1), corresponding to the mature green stage, at which stage mucilage synthesis completed. The expression pattern of IRX14 was coincident with that of CESA5 (Supplementary Fig. S1), which has been shown to be responsible for the synthesis of cellulose in seed mucilage (Sullivan et al., 2011). Although cellulose microfibrils are synthesized by plasma membrane-localized cellulose synthase complexes and xylans are synthesized at the Golgi apparatus (reviewed in Driouich et al., 2012), both are transported and deposited into the apoplast of the epidermal cells of the seed coat where they may form extensive cross-linking embedded in a matrix of pectic polysaccharides. Not surprisingly, mutation of the key components YIP4a, YIP4b, and ECHIDNA
**Fig. 6.** In situ labelling of crystalline cellulose in adherent mucilage from *irx14-1* and WT seeds. (A, D, G, J) Staining of β-glycans with Calcofluor White. (B, E, H, K) Indirect immunofluorescence detection of His-tagged CBM3a binding to crystalline cellulose. (C, F, I, L) Composite images of double labelling with Calcofluor and CBM3a. (D–F) and (J–L) correspond to the magnification of regions in (A–C) and (G–I), respectively. Bar=100 μm.

**Fig. 7.** In-situ labelling of xylan in mucilage released from *irx14-1* and WT seeds. (A1, A2, D1, D2, G1, G2, J1, J2) Staining of β-glycans with Calcofluor White. (B1, E1, H1, K1) Detection of immunofluorescence for CCRC-M139 binding to mucilage xylan. (B2, E2, H2, K2) Immunofluorescence detection of LM11 binding to higher-substituted xylan. (C1, F1, I1, L1) Composite images of double labelling with Calcofluor and CCRC-M139. (C2, F2, I2, L2) Composite images of double labelling with Calcofluor and LM11. (D1–F1) and (J1–L1) correspond to the magnification of regions in (A1–C1) and (G1–I1), respectively. (D2–F2) and (J2–L2) are magnified regions in (A2–C2) and (G2–I2), respectively. Bar=100 μm.
(ECH), which mediate the post-Golgi secretion of pectin and hemicellulose, significantly decreased the mucilage released from seeds (Gendre et al., 2013). These results imply that cellulose synthesized by CESA5 and xylan synthesized by IRX14 may be simultaneously deposited into the apoplast wherein they join together to form an extensible cross-linking network.

The mucilage phenotype of irx14-1 resembles that of several previously reported mutants affecting cellulose biosynthesis or organization in seed mucilage, such as the cellulose synthase (CESA) subunit mutant cesa5 (Sullivan et al., 2011), the leucine-rich receptor like kinases mutant fei2 (Harpaz-Saad et al., 2012), and the COBRA-like 2 mutant cobl2 (Ben-Tov et al., 2015). Ruthenium red staining of these mutant seeds without shaking demonstrated the presence of two layers of mucilage, but only a small amount of the inner layer remained attached to the seeds after shaking (Sullivan et al., 2011; Harpaz-Saad et al., 2012; Ben-Tov et al., 2015). The water-soluble fraction was significantly increased in irx14-1 as observed in these mutants, which resulted in a modified distribution between the outer and the inner layers of mucilage (Fig. 4). Surprisingly, mutations in IRX14 also led to reduced levels of crystalline cellulose in seed coat mucilage (Fig. 5; Supplementary Fig. S4). Cellulose has been shown to play important roles in anchoring the adherent mucilage to the seed coat (Sullivan et al., 2011; Harpaz-Saad et al., 2012). However, precisely how cellulose mediates the attachment of adherent mucilage to the seed surface remains to be clarified. Although it is possible that RG I in the adherent mucilage became entangled with cellulose microfibrils to form an extensible network, our results suggest that xylan may serve as an intermediate component to link RG I and cellulose together.

This model agrees with the prevailing structural model of the cell wall that depicts the cellulose-hemicellulose network as the load-bearing component embedded in a pectin matrix (Carpita and Gibeaut, 1993). Supporting this model, the selective adsorption of xylan on to cellulose microfibrils has been extensively studied, mostly by an in vitro combination assay (Linder et al., 2003; Kabel et al., 2007; Köhnke et al., 2008; Busse-Wicher et al., 2014; Li et al., 2015). It has been shown that xylan tends to be aligned parallel to the direction of the cellulose microfibrils, and the composition and side chain substitutions of xylan significantly affects the adsorption capacities of the cellulose fibres (Linder et al., 2003;
Kabel et al., 2007; Busse-Wicher et al., 2014). In addition, it was documented that adsorption of xylan on to cellulose-based fibres improved the physical processability of the pulp and resulted in paper with good tensile properties (Silva et al., 2011). Although extensive studies indicated that xylan and cellulose could join together to form a cross-linking network, most of these studies were carried out in vitro, and the nature of the molecular bonding and arrangement between xylan and cellulose is still unclear. Our results indicated that xylan and cellulose might form cross-linking in vivo in seed coat mucilage, which is an important step toward the understanding of interactions between xylan and cellulose.

Furthermore, the reduced synthesis of xylan in seed mucilage significantly affected the macromolecular characteristics of water-soluble mucilage in irx14-1 (Fig. 4). The larger polymeric fraction s completely disappeared in irx14-1 and the presence of the smaller polymeric fraction was slight compared with the WT. This indicated that the polymeric fractions corresponding to aggregated or entangled polymers were structurally different in irx14-1 mucilage. Accordingly, ELISA and the immunoblotting assay using monoclonal antibodies specific for different mucilage components (e.g. RG I, HG, xylan, etc) further confirmed that substantial alterations in ultra-structure occurred in irx14-1 mucilage. (Fig. 8; Supplementary Fig. S6). These results indicated that cross-linking of xylan to cellulose and/or other mucilage polymers was disturbed by IRX14 mutation, and the disturbance led to further structural alterations in other components of irx14-1 mucilage. Combined with the data presented here, we speculate that xylan may join to cellulose microfibrils and embed in the pectic polysaccharide matrix in the seed coat mucilage. Loss of xylan in irx14-1 seed mucilage may disturb the association between xylan and cellulose and further affect the self-association of cellulose. However, how xylan cross-links with cellulose and other mucilage polymers needs to be further investigated.

The mucilage defect of irx14-1 also resembles that of mum5 identified in a previous forward genetic screen for modified seed mucilage (Western et al., 2001), which prompted us to consider whether IRX14 and MUM5 are genetically identical. To test this hypothesis, the full-length coding sequence of IRX14 was amplified by PCR using cDNA of mum5 and sequenced. However, no point mutations were identified in the coding sequence of IRX14 amplified from mum5 cDNA, which indicates that IRX14 and MUM5 represent different alleles. The identity of MUM5 remains unknown. Further characterization of MUM5 function will deepen our understanding of the interaction among mucilage polymers.

In summary, the work presented here indicated that IRX14, which is involved in the biosynthesis of mucilage xylan, plays an important role in maintaining the structure of Arabidopsis seed coat mucilage. Our results demonstrated that, besides galactoglucomannan, another hemicellulose component, xylan was responsible for the structuration and adherence of the seed mucilage. These findings imply that xylans interact with cellulose to form an extensible network embedded in the pectin matrix, which is essential for anchoring the pectic adherent mucilage to the seed surface. The study expanded our understanding of the function of xylan and its interaction with cellulose, which is an important step forward in unravelling how mucilage polymers cross-link and assemble. Future studies are needed to clarify how xylan cross-links with cellulose and other mucilage components.

Supplementary data
Supplementary data can be found at JXB online.

Figure S1. Heatmap showing hierarchical clustering of putative candidates and reported genes involved in mucilage biosynthesis and/or modification during seed coat development.

Figure S2. Seed coat specific expressions of four GT43 members.

Figure S3. Seed coat epidermal cell differentiation is not altered in the irx14-1 mutant.

Figure S4. Fourier Transform Infrared Spectroscopy (FTIR) analysis of WT and irx14-1 mucilage.

Figure S5. Water-soluble mucilage from irx14-1 is more electrophoretically mobile than the WT.

Figure S6. Glycome profiling of the mucilage from irx14-1 and WT seeds by ELISA assay.

Table S1. List of primers used in this study.

Acknowledgements
We would like to thank Professor Michael G Hahn (Complex Carbohydrate Research Center, University of Georgia, USA) and Professor J Paul Knox (Centre for Plant Sciences, Faculty of Biological Sciences, University of Leeds, UK) for kindly providing the series of monoclonal antibodies. This work was supported by the National Natural Science Foundation of China (31370328 and 31400176), the National Key Technology Support Program of China (2013BAD22B01), the Youth Innovation Promotion Association of CAS (2014187), the Taishan Scholar Program of Shandong (to GZ), and the Elite Youth Program of CAAS (to YK).

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