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RESEARCH PAPER

Mannans and endo-β-mannanases (MAN) in *Brachypodium distachyon*: expression profiling and possible role of the *BdMAN* genes during coleorhiza-limited seed germination

Virginia González-Calle, Cristina Barrero-Sicilia†, Pilar Carbonero and Raquel Iglesias-Fernández*

Centro de Biotecnología y Genómica de Plantas (UPM-INIA), and ETSI Agrónomos, Campus de Montegancedo, Universidad Politécnica de Madrid, Pozuelo de Alarcón, 28223-Madrid, Spain

† Present address: Biological Chemistry and Crop Protection Department, Rothamsted Research, Harpenden AL5 2JQ, UK

* To whom correspondence should be addressed. E-mail: raquel.iglesias@upm.es

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Abstract

Immunolocalization of mannans in the seeds of *Brachypodium distachyon* reveals the presence of these polysaccharides in the root embryo and in the coleorhiza in the early stages of germination (12 h), decreasing thereafter to the point of being hardly detected at 27 h. Concurrently, the activity of endo-β-mannanases (MANs; EC 3.2.1.78) that catalyse the hydrolysis of β-1,4 bonds in mannan polymers, increases as germination progresses. The MAN gene family is represented by six members in the *Brachypodium* genome, and their expression has been explored in different organs and especially in germinating seeds. Transcripts of *BdMAN2*, *BdMAN4* and *BdMAN6* accumulate in embryos, with a maximum at 24–30 h, and are detected in the coleorhiza and in the root by *in situ* hybridization analyses, before root protrusion (germination *sensu stricto*). *BdMAN4* is not only present in the embryo root and coleorhiza, but is abundant in the de-embryonated (endosperm) imbibed seeds, while *BdMAN2* and *BdMAN6* are faintly expressed in endosperm during post-germination (36–42 h). *BdMAN4* and *BdMAN6* transcripts are detected in the aleurone layer. These data indicate that *BdMAN2*, *BdMAN4* and *BdMAN6* are important for germination *sensu stricto* and that *BdMAN4* and *BdMAN6* may also influence reserve mobilization. Whether the coleorhiza in monocots and the micropylar endosperm in eudicots have similar functions, is discussed.

Key words: *BdMAN* gene family, *Brachypodium distachyon*, coleorhiza, endo-β-mannanases, germination, MAN gene expression, mannan immunolocalization, mRNA *in situ* hybridization.

Introduction

Poaceae grains (caryopses) include the seed proper, formed by a triploid endosperm and a diploid embryo, surrounded by the maternal tissues of the seed coat (testa) and the pericarp. The coleorhiza is a non-vascularized multicellular embryonic tissue, covering the seminal roots of Poaceae seeds. The coleorhiza has been thought to have a role in protecting the emerging root (Sargent and Osborne, 1980) and, more recently, it has been also associated with the regulation of dormancy, since abscisic acid (ABA) sensitivity is reduced in this tissue during germination of non-dormant barley seeds and the gene encoding the HvABAS’OH-1 enzyme, that is critical for ABA degradation, is expressed in the coleorhiza. During germination, both in barley and in *Brachypodium* seeds, the coleorhiza is the first structure that protrudes after the pericarp and testa rupture (coleorhiza emergence), followed by the coleorhiza rupture that allows root emergence (root emergence), and indicates the end of germination *sensu stricto* (Millar et al., 2006; Barrero et al., 2009, 2012; Gao and Ayele, 2014).

The germination process can be separated into germination *sensu stricto* and subsequent reserve mobilization (post-germination) and has been more deeply investigated in eudicotyledonous than in monocotyledonous seeds. In
Arabidopsis thaliana, Sisymbrium officinale, Lepidium sativum and Nicotiana tabacum, the germination sensu stricto occurs in two different steps: first, the testa ruptures and, afterwards, the micropylar endosperm breakage takes place, allowing the radicle to emerge (Leubner-Metzger and Meins, 2000; Nonogaki et al., 2000; Müller et al., 2006; Iglesias-Fernández and Matilla, 2010; Iglesias-Fernández et al., 2011a, b; Weitbrecht et al., 2011; Nonogaki, 2014). Addition of ABA to the imbibition medium specifically blocks endosperm weakening and prevents its rupture (Müller et al., 2006; Piskurewicz et al., 2009; Carrillo-Barral et al., 2014). It is assumed that testa rupture is influenced by the driving force of the imbibed elongating radicle and that the endosperm rupture is mainly produced by the weakening of the endosperm cell walls (CWs) by enzymes, specifically those localized to the micropylar endosperm, such as endo-β-1,4-mannanases (MANs), endo-β-1,3-glucanases, expansins, xyloglucan-transglycosylases/hydrolases (XTHs) and pectin-methylesterases (Leubner-Metzger, 2005; Nonogaki et al., 2007; Iglesias-Fernández et al., 2011a, b; Endo et al., 2012; Martínez-Andújar et al., 2012; Rodriguez-Gacio et al., 2012; Scheler et al., 2014).

Since the endosperm CWs of several eudicot seeds are rich in mannan (Lee et al., 2012), the MAN activity and the expression of MAN genes upon seed germination have been further characterized and their transcriptional regulation studied. In A. thaliana, four MAN genes (AtMAN2, AtMAN5, AtMAN6 and AtMAN7) are expressed in germinating seeds and their transcripts are restricted to the micropylar endosperm and to the radicle, disappearing as soon as the radicle emerges. Moreover, knock-out mutants in the AtMAN5, AtMAN6 and AtMAN7 genes, as well as, in the AthbZIP44 gene encoding an important activating transcription factor of AtMAN7, have a significantly retarded germination as compared to that of wild-type seeds, indicating a role for these MAN genes and their regulators during germination sensu stricto (Iglesias-Fernández et al., 2011a, b, 2013; Rodriguez-Gacio et al., 2012; Yan et al., 2014).

Brachypodium distachyon is being considered a model species for the genetics and molecular genomics of cereals, due to its small sequenced genome (~355 Mbp), short life cycle, self-fertility, diploidy and its close phylogenetic relationship with important crop plants of the tribe Triticeae within the Poaceae family, such as wheat and barley (International Brachypodium Initiative, 2010; Mochida and Shinozaki, 2013; Girin et al., 2014). In Poaaceae seeds, the β-1,3,1,4-glucans are abundant in the endosperm cell walls (Burton and Fincher, 2009; Guillón et al., 2011) and genes encoding hydrolytic enzymes involved in their degradation, such as endo-β-1,3-glucanases and endo-β-1,3,1,4-glucanases, have been associated with rice post-germination events (2–4 days of imbibition; Akiyama et al., 2004) and with the elongation of barley coleoptiles (Takeda et al., 2010). Although mannan content is lower than glucan content in Brachypodium seeds (Rancourt et al., 2012), the function of mannans and MANs may be relevant in its germinating seeds.

In this work, mannan polysaccharides were immunolocalized to the root and the coleorhiza of germinating seeds early in imbibition, decreased thereafter at later stages, and the enzymatic activity of endo-β-mannanases increased as germination progressed. The MAN gene family of B. distachyon was annotated and the expression of its six members explored in vegetative and reproductive organs. Interestingly, genes BdMAN2, BdMAN4 and BdMAN6 were clearly induced upon seed germination and mRNA in situ hybridization analyses demonstrated that these transcripts were found in the coleorhiza and the root during germination sensu stricto. BdMAN4 and BdMAN6 were also expressed in the aleurone layer, and may also be involved in post-germinative reserve mobilization.

Materials and Methods

Biological material, growth conditions and germination assays

The diploid inbred Brachypodium distachyon strain Bd21 (kindly provided by Prof. Garvin from the University of Minnesota, USA; International Brachypodium Initiative, 2010) was used in this work. Seeds were surface-sterilized with 1% NaOCl for 10 min and washed in sterile water, before germinating on Petri dishes, containing two filter papers (Whatman 3) moistened with 8 ml of sterile water, at 22°C in the dark for 2 d. They were then transferred to pots in the greenhouse under long-day conditions (16h/8h, light/darkness; light intensity 155 μmol photons m⁻² s⁻¹) for sampling roots (6-week-old plants), young and old leaves (6- and 12-week-old plants) and spikes. For the germination experiments that lasted up to 42 h, seeds were incubated in the dark at 22°C, in Petri dishes with moistened filter papers, using triplicate lots of 25 non-stratified after-ripened seeds (stored at 22°C and 30% relative humidity in the dark for 3 months). Seed samples were separated into embryo and endosperm (de-embryonated seeds) at 0, 12, 24, 30, 36 and 42 h of imbibition, and used for RNA quantification and for protein extraction to determine MAN enzymatic activity.

Endo-β-1,4-mannanase (MAN) activity assays

Seed samples, obtained as described above, were homogenized in 100 mM sodium acetate buffer (pH 4.5) containing 1 M NaCl and 0.5% ascorbic acid, at 4°C. The homogenates were centrifuged at 15,000 × g for 45 min, and 80 μl of this supernatant was mixed with 150 μl of 0.25% mannan (1,4-β-D-mannan from carob; Megazyme International Ireland Ltd., Wicklow, Ireland). Incubation was at 30°C for different periods of time and the enzymatic activity was determined by the increase in reducing sugar production per mg of protein, as determined by the 4-OH-Benzoic Acid Hydrazide (PAH-BAH; Sigma-Aldrich) method (Lever, 1977). The MAN from Aspergillus niger (Megazyme) was used to establish the control curve (one unit of MAN activity defined as the amount of enzyme that releases 1 nmol of reducing sugar per minute under the experimental conditions). Protein concentration was determined with the Bradford reagent (Bio-Rad Laboratories, Munich, Germany) using bovine serum albumin (BSA) as a standard.

Bioinformatic tools: BdMANs identification and phylogenetic analysis

The deduced protein sequences of the six MAN genes were obtained from the B. distachyon genome using the TBLASTN tool at the Phytozome v8.0 Database (Goodstein et al., 2012; www.phytozome.net), using the eight OsMAN proteins from the Oryza sativa genome as query sequences (Yuan et al., 2007). The Interpro Program (PFAM database; Bateman et al., 2002; http://pfam.sanger.ac.uk) was used to confirm the presence of the MAN conserved
domain (glycosyl-hydrolase family 5). The complete amino acid sequences deduced from \textit{B. distachyon}, \textit{O. sativa} and \textit{Arabidopsis thaliana} \textit{MAN} genes were aligned by means of the CLUSTAL W program (Thompson et al., 1994) and utilized to construct a phylogenetic dendrogram, using the neighbor-joining algorithm, a bootstrap analysis with 1,000 replicates, complete deletion and the Jones Taylor Thornton matrix, as settings. The MEME program software version 4.0 (Tamura et al., 2007) was used to identify conserved motifs within the deduced \textit{MAN} proteins and to validate the phylogenetic tree (Table 1). Default parameters were used with the following exceptions: the maximum number of motifs to find was set to 22 and the minimum width was set to eight amino-acid residues (Bailey et al., 2009; http://meme.sdsc.edu/meme4_6_0/introt.html). A single capital letter represents a single residue relative frequency, if this is greater than 50% than twice that of the second most frequent residue in the same position. If no single residue matches these criteria, a pair of residues, represented by capital letters in brackets, is given if the sum of their relative frequencies exceeds 75%. If none of these characteristics are satisfied, a lower-case letter is given when the relative frequency of a residue is greater than 40%, if not, \(x\) is set.

The major biochemical parameters of the deduced \textit{MAN} proteins from \textit{B. distachyon} and \textit{O. sativa} are listed in Supplementary Table S1. Both isoelectric point (pl) and molecular weight (MW) were predicted using the Compute pl/MW tool (Gasteiger et al., 2005; http://www.expasy.ch/tools/pi_tool.html) and the putative signal peptide cleavage site and sub-cellular localization were deduced by the SignalP 3.0 (http://www.cbs.dtu.dk/services/SignalP) and TargetP 1.1 tools (http://www.cbs.dtu.dk/services/TargetP/), respectively (Emanuelsson et al., 2007).

Real time quantitative PCR (RT-qPCR) analyses

Total RNA was purified from roots (6-week-old plants), young and old leaves (6- and 12-week-old plants) and spikes by the phenol/chloroform method (Lagrimini et al., 2007). First-strand cDNA was synthesized with random hexamers using the FastStart SYBR Green Reaction Mix (Roche Applied Science). Probes were hybridized at 52ºC overnight followed by two washes in 2× SSC (150 mM sodium chloride, 15 mM sodium acetate, pH 7), and then treated with de-ionized water. For heteromannan immunodetection, sections were first incubated at room temperature for 30 min in a blocking solution (3% BSA, 1× PBS, and 5 mM sodium azide; pH 7), and then treated with primary anti-heteromannan antibody LM21 (PlantProbes, Leeds, UK) at a dilution of 1:5 in the same blocking solution but only containing 1% BSA for 2 h. Sections were thoroughly washed in PBS containing 5 mM sodium azide and then incubated for 2 h in the same buffer containing the secondary rabbit antibody Anti-Rat IgG-FITC (Sigma-Aldrich) at a dilution of 1:100. The sections were extensively washed in PBS buffer and in water, mounted and examined in a confocal microscope (absorption 494 nm; emission 521 nm; Leica TCS-SP8, Leica, Wetzlar, Germany).

Pre-hybridization was carried out by incubating the sections in 0.2 M HCl, neutralizing them and then treating them with 1 mg/ml proteinase-K (Roche Applied Science). Samples were then dehydrated in an aqueous ethanol dilution series and hybridized with sense and anti-sense digoxigenin (DIG)-labelled RNA probes, corresponding to the non-coding regions of the \textit{BdMAN2}, \textit{BdMAN4} and \textit{BdMAN6} genes (Supplementary Table S3), synthesized with the DIG RNA labelling mix according to the manufacturer’s specifications (Roche Applied Science). Probes were hybridized at 52ºC overnight followed by two washes in 2× SSC (150 mM NaCl, 15 mM Na\textsubscript{2} citrate) and 50% formamide for 90 min at the same temperature. Incubation with the alkaline phosphatase-conjugated anti-digoxigenin antibody (Roche Applied Science) and colour detection was carried out according to the manufacturer’s instructions (Ferrandiz et al., 2000). Sections were dried and examined on a Zeiss AxioPhot Microscope (Carl Zeiss, Oberkochen, Germany), and images were captured and processed with the Leica Application Suite 2.8.1 build software (Leica).

Heteromannan immunolocalization

The protocol used was a modification of those described in Marcus et al. (2010) and Guillon et al. (2011). In a pre-immunolabeling step, sections of embedded material as described above, were incubated in phosphate buffer sodium solution (PBS) and treated with 1 mg/ml proteinase-K (Roche Applied Science). For the specific antibodies to have access to the heteromannans (mannans, glucomannans, and galactomannans) of the cell walls, \(\beta\)-1,3-1,4-glucans were removed by incubating the sections with a solution of 4 mg/ml lichenase \(\beta\)-1,3-1,4-glucanase; Megazyme for 2 h at 37°C, and then rinsed with de-ionized water. For heteromannan immunodetection, sections were first incubated at room temperature for 30 min in a blocking solution (3% BSA, 1× PBS, and 5 mM sodium azide; pH7), and then treated with primary anti-heteromannan antibody LM21 (PlantProbes, Leeds, UK) at a dilution of 1:5 in the same blocking solution but only containing 1% BSA for 2 h. Sections were thoroughly washed in PBS containing 5 mM sodium azide and then incubated for 2 h in the same buffer containing the secondary rabbit antibody Anti-Rat IgG-FITC (Sigma-Aldrich) at a dilution of 1:100. The sections were extensively washed in PBS buffer and in water, mounted and examined in a confocal microscope (absorption 494 nm; emission 521 nm; Leica TCS-SP8, Leica, Wetzlar, Germany).
on a Zeiss Axiophot Microscope (Carl Zeiss) and the images were captured and processed with the Leica Application Suite 2.8.1 build software (Leica).

**Results**

*Enzymatic β-mannanase activity during Brachypodium seed germination*

The time course of *sensu stricto* germination of *B. distachyon* seeds occurs in two different steps: first, the coleorhiza emerges (CE), and in a second step, the root emergence (RE) takes place (Fig. 1A). The enzymatic activity of MAN upon germination has been analysed separately in the embryo and in the de-embryonated seed (endosperm). As shown in Fig. 1B, dried seeds have no detectable MAN activity, but this progressively increases with germination, peaking at 24 h in embryos, containing the coleorhiza ($\sim0.4 \times 10^{-3}$ units/mg protein), and decreasing to half this value at 42 h ($\sim0.2 \times 10^{-3}$ units/mg protein). In endosperms, MAN activity is much lower than in embryos, and reaches its maximum level ($\sim0.15 \times 10^{-3}$ units/mg protein) at 36 h of germination. Data from Fig. 1B indicate that MAN activity is maximum in embryos, just before

![Fig. 1.](http://jxb.oxfordjournals.org/)

*(A) Longitudinal sections of the different phases of *Brachypodium distachyon* germination *sensu stricto*, stained with toluidine blue. C, coleorhiza; Co, coleoptile; E, endosperm; Sc, scutellum; Sh, shoot; R, root. Scale bar, 200 μm. (B) Endo-β-mannanase activity (white bars) in embryo and endosperm (de-embryonated seed) upon *B. distachyon* seed germination (0–24 h). One unit of MAN activity is defined as the amount of enzyme that releases 1 nmol of reducing sugar per minute and per mg of protein. Percentage germination evaluated as coleorhiza emergence (CE; open circles) and root emergence (RE; close circles) are represented. In the inset, the time needed for 50% of CE ($t_{50}$CE) and RE ($t_{50}$RE) is indicated. Data are means ± standard error (SE) of three technical replicates of three biological samples.*
reaching 50% of germination *sensu stricto* \(t_{50CE}=30\text{h}; t_{50RE}=36\text{h}\), suggesting that MAN is important for facilitating both coleorhiza and root emergence.

**Heteromannans are preferentially localized to the root tip and the coleorhiza in germinating seed embryos**

Mannan polymers have been detected in longitudinal sections of *B. distachyon* germinating seeds (at 12 and 27 h of imbibition) by *in situ* immunofluorescence labelling, using the LM21 antibody that specifically recognizes mannan polysaccharides (gluco- and galacto-mannans). To facilitate access of the antibody to mannans in plant CWs, the seed sections have been previously treated with lichenase (\(\beta\)-1,3-1,4-glucanase; Marcus et al., 2010).

As shown in Fig. 2, at 12 h of seed imbibition, seed mannan polymers are mainly localized to the periphery cells of the coleorhiza (C) and to the epidermis of the root tip (R) (Fig. 2A–C). Interestingly, these mannans are barely detected at later stages of germination (27 h of imbibition; Fig. 2D–F). Differential interference contrast (DIC) images are shown in Fig. 2G–I. This observation together with data of MAN enzymatic activity (Fig. 1B) with a maximum at 24 h in embryos, may suggest that the disappearance of the mannan polymers is due to the hydrolysis catalysed by endo-\(\beta\)-mannanases.

**The Brachypodium endo-\(\beta\)-mannanase gene family**

In order to get a deeper insight into the MAN function upon *B. distachyon* germination, it was decided to annotate and characterize further the BdMAN family. The already described MAN family from *O. sativa* (Yuan et al., 2007) has been used to perform a TBLASTN against the whole *Brachypodium* genome (http://www.phytozome.net). Six predicted non-redundant MAN deduced proteins, with MW 43–52 KDa, and Ip 4.4–8.8, three of them with predicted signal peptides, have been identified and named according to their orthologues in rice (Supplementary Table S1).

As shown in Fig. 2, at 12 h of seed imbibition, seed mannan polymers are mainly localized to the periphery cells of the coleorhiza (C) and to the epidermis of the root tip (R) (Fig. 2A–C). Interestingly, these mannans are barely detected at later stages of germination (27 h of imbibition; Fig. 2D–F). Differential interference contrast (DIC) images are shown in Fig. 2G–I. This observation together with data of MAN enzymatic activity (Fig. 1B) with a maximum at 24 h in embryos, may suggest that the disappearance of the mannan polymers is due to the hydrolysis catalysed by endo-\(\beta\)-mannanases.

The MAN protein sequences from *A. thaliana* (AtMAN1-7) and *O. sativa* (OsMAN1-8) together with those from *B. distachyon* (BdMAN1-6) have been used to construct a phylogenetic unrooted tree by using the neighbor-joining algorithm. Four major clusters of orthologous groups (MCOGs) have been defined (A, B, C, D), supported by bootstrapping values higher than 62% (Fig. 3A) and by the occurrence of common motifs (Fig. 3B; MEME).

The search for conserved amino-acid motifs using the MEME software (http://meme.nbcr.net/meme/cgi-bin/...
meme.cgi) reveals that all MAN sequences have in common motifs described as critical for the enzymatic activity, such as 1, 3, 5, 6/11 and 7 (Fig. 3B, Table 1). The deduced signature sequence [AWEL(MI)NEPRC] of Arabidopsis and rice MANs (Yuan et al., 2007), included in motif 1, is also present in Brachypodium MAN. Besides, members in MCOG A, BdMAN2, OsMAN2, BdMAN6 and OsMAN6 share motif 12, and BdMAN6 and OsMAN6 also share motifs 14 and 18. In MCOG C, BdMAN1 shares with OsMAN1 motifs 19 and 22, and in MCOG D BdMAN4 shares motifs 17 and 21 with OsMAN4, but lacks motif 15 shared by the rice paralogues OsMAN3 and OsMAN4. Similarly, BdMAN5, OsMAN7 and OsMAN8 (MCOG B) have in common motif 11, but they do not share with OsMAN5 motifs 20 and 13.

Table 1. Conserved amino acid motifs obtained by means of MEME (Bailey et al., 2009) from the analysis of the endo-β-mannanase proteins of Brachypodium distachyon, Oryza sativa, and Arabidopsis thaliana.

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Fig. 3. (A) Phylogenetic dendrogram with deduced protein sequence of the mannanase gene families from Brachypodium distachyon, Oryza sativa and Arabidopsis thaliana; bootstrapping values are indicated in the branches. (B) Schematic distribution of conserved motifs among the deduced protein sequences in the phylogenetic tree (A), identified by means of the MEME analysis. Asterisks indicate those motifs important for enzymatic activity. Motifs in grey share >85% of similar amino acid residues. This figure is available in colour at JXB online.
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Expression kinetics of selected BdMAN genes during seed maturation and germination

The expression pattern of the six BdMAN genes has been explored by RT-qPCR analysis in different organs: young (6 d) and old (12 d) leaves, roots (6 d) and spikes (mix of different stages; Supplementary Fig. S2). While BdMAN1, 2, 3, 5 genes are not detected in leaves, BdMAN4 gene expression in old leaves is ~10 times lower than in young leaves, and BdMAN6 has the same low expression in young and old leaves. In roots, BdMAN2, BdMAN4 and BdMAN6 are expressed at low levels, and BdMAN1, BdMAN2, BdMAN4 and BdMAN6 transcripts are detected in spikes (Supplementary Fig. S2).

Since our preliminary data indicate that BdMAN1, BdMAN2 and BdMAN6 transcripts are abundant in developing seeds (see Supplementary Fig. S3A), the expression kinetics of these three genes has been established throughout seed maturation, at 4, 6, 8, 10 and 12 d after pollination (dap) (Fig. 4A). Although the gene BdMAN1 is the most highly expressed during the late phases of seed development (8, 10, 12 dap), the expression patterns of BdMAN1, BdMAN2 and BdMAN6 show a progressive increase from 4 to 10 dap, reaching all of them their maximum expression at 10 dap when maturation is almost completed (Fig. 4B).

Data in Supplementary Fig. S3B indicate that genes BdMAN2, BdMAN4 and BdMAN6 are the most abundantly expressed ones during seed germination, and their expression kinetics has been more thoroughly analysed (Fig. 5). Germinating seeds taken at 12, 24, 30, 36 and 42 h of imbibition, have been sectioned into embryos and de-embryonated seeds (endosperm). In germinating embryos, BdMAN2, BdMAN4 and BdMAN6 transcripts appear early upon imbibition (12 h), before CE, and their maximum expression is attained between 24–30 h (t50CE=30 h) and it decreases as germination progresses (42 h; Fig. 5A). However, the expression of BdMAN4 in endosperms is high at early imbibition times (12 h; ~140% relative to BdGAPDH), decreasing thereafter. The BdMAN2 and BdMAN6 transcripts have low expression in the endosperms of germinating seeds with a maximum at 36 h of imbibition (<10 % for BdMAN2 and ~25 % for BdMAN6 relative to BdGAPDH, respectively), indicating a possible role in reserve mobilization for these MAN genes, and perhaps also for BdMAN4 post-germination (Fig. 5B).

BdMAN2, BdMAN4 and BdMAN6 transcripts are localized to different seed tissues during B. distachyon seed germination

To determine the spatial expression of MAN genes within the B. distachyon germinating seeds, mRNA in situ hybridization experiments have been done (Fig. 6). Longitudinal sections of seeds at 27 h of imbibition have been hybridized to specific antisense and sense (as negative controls) probes for BdMAN2, BdMAN4 and BdMAN6. The BdMAN2 transcripts are mainly expressed in the periphery cells of the coleorhiza and are not detected in the aleurone layer (Fig. 6A–D). BdMAN4 mRNA is localized preferentially to the tip and the apical meristem of the root, to the coleorhiza and to the aleurone layer.
and the BdMAN6 transcripts are detected not only at the coleorhiza, but also throughout the embryo and faintly also at the aleurone layer (Fig. 6E–H). BdMAN6 transcripts are detected in the aleurone layer at 27 h of imbibition by mRNA in situ hybridization, but they are scarcely detected in 24–30 h germinating endosperms by RTqPCR (Fig. 5B), indicating that its expression could be diluted by the remaining endosperm during the RNA isolation process. As expected, no signal has been detected when sections have been hybridized with the corresponding sense probes for BdMAN2, BdMAN4 and BdMAN6 (negative controls; Figs 6D, 6H and 6L).

**Discussion**

In this work, mannans and endo-β-mannanases (MAN) in Brachypodium distachyon have been investigated in order to establish whether they are important in the germination of these monocotyledonous seeds. Mannans have been immunolocalized in the embryo root and the coleorhiza in the early stages of germination and these polymers decrease upon imbibition while the enzymatic activity of MAN increases. The MAN gene family in B. distachyon has been annotated and the gene expression of the six members of this family has been explored in different vegetative and reproductive organs, and, more specifically, in germinating seeds. Three of these genes, BdMAN2, BdMAN4 and BdMAN6, are highly induced in...
Mannans and MAN genes during B. distachyon germination

During seed development, BdMAN1, BdMAN2 and BdMAN6 genes are expressed, and their mRNAs are abundant at the middle and late maturation stages. Upon cereal seed maturation, several tissues undergo a progressive enlargement of their cells, a process that involves nutrient remobilization and CW softening to allow cell expansion; to this aim, the participation of a complex set of hydrolytic enzymes have been described (Domínguez and Cejudo, 2014). These data indicate that the BdMAN1, BdMAN2 and BdMAN6 proteins could contribute to such a process during Brachypodium seed development.

The enzymatic analysis in the embryos of germinating seeds shows a maximum of MAN activity at 24 h of imbibition, just before the coleorhiza emergence (CE50=30 h), and this enzymatic activity progressively decreases to 50% at 42 h. However, MAN activity in de-embryonated seeds...

Fig. 6. In situ mRNA hybridization analysis of BdMAN2, BdMAN4 and BdMAN6 in 27 h germinating Brachypodium seeds. (A–D) BdMAN2, (E–H) BdMAN4, (I–L) BdMAN6. (A, E, I) Longitudinal sections of germinating embryos. (B, F, J) Close-up of the coleorhiza and the root tip. (C, G, K) Close-up of the endosperm and the aleurone. (D, H, L) Control sense probes. Al, aleurone layer; C, coleorhiza; Co, coleoptile; Cp, calyptra; E, endosperm; Sc, scutellum; Sh, shoot; R, root. The black arrow indicates the localization of transcripts. Scale bar, 50 μm.
(endosperms) is low, with a maximum at 36 h of imbibition, when germination sensu stricto is almost completed [~100% coleorhiza emergence (CE); ~80% root emergence (RE) at 42 h]. These data point out to a more important role of the MAN activity in the embryo than in the endosperm during germination sensu stricto. In rice, MAN activity and expression of the OsMAN1, OsMAN2 and OsMAN6 genes have been detected in the aleurone layer only after 48 h of imbibition when 100% of germination has been achieved. This MAN activity is associated with reserve mobilization, a clear post-germinative event (Ren et al., 2008). In barley, the HvMAN1 enzyme has been purified from 10-day-old seedlings and its catalytic parameters established, although its physiological role has not been investigated (Hrmova et al., 2006).

The function of the coleorhiza tissue in the grasses has been classically associated to a protective function of the growing root during germination, but other physiological functions are being uncovered, such as our observations of the hydrolysis of proteins (disappearance of PBs) or the decrease in mannan content detected within the coleorhiza cells during Brachypodium germination sensu stricto. Nowadays, and similarly to what has been proposed for the endosperm of eudicot seeds (Piskurewicz et al., 2009), the coleorhiza is being considered to be a key tissue preventing root emergence in dormant barley seeds (Millar et al., 2006; Barrero et al., 2009). These authors have hypothesized that root emergence may not depend only on the softening of the coleorhiza, driven by CW remodelling enzymes, but also by the expansive force of the imbibing root cells. Important transcriptional changes in the barley coleorhiza associated to the dormancy degree have been found and these differences affect mainly the expression of CW modifying genes (mannanases among them), nitrate and nitrite reductase genes etc. (Barrero et al., 2009). The cytosolic nitrate reductase is an important source of the hormone nitric oxide (NO) that is involved in promoting seed germination (Arc et al., 2013). Therefore, the coexistence at the coleorhiza of NO and mannanases, and perhaps proteases of the CatepsinB3 type (Iglesias-Fernández et al., 2014), should have an influence in the seed germination of the grasses.

The Arabidopsis radicle tip has been described as the primary location of growth-promoting genes and its

Fig. 7. Polysaccharide and protein mobilization upon B. distachyon seed germination. Bright field microscopy of longitudinal seed sections stained with PAS-Naphthol Blue Black. (A, D, G, J) Longitudinal sections from dry and water-imbibed seeds at 27 h and 36 h. (B, E, H, K) Close-up of the coleorhiza in A, D, G, J and (C, F, I, L) close-up of the endosperm in A, D, G, J, respectively. Proteins stain in blue and polysaccharide-rich cell walls in pink. C, coleorhiza; Co, coleoptile; E, endosperm; M, mesocotile; Sc, scutellum; Sh, shoot; R, root. Scale bar: 50 μm.
surrounding-cells the centre for CW expansion (Bassel et al., 2014). Moreover, a dual enzymatic activity for MAN (hydro-
lase and transglycosylase activities) has been described; the
transglycosylase activity being more related to cell expansion,
as occurs in the radicle before protrusion, and the hydrolytic
activity could be relevant for weakening of the CWs of the
embryo-surrounding tissues (Schröder et al., 2009; Iglesias-
Fernández et al., 2011a, b). It is remarkable that the BdMAN4
transcripts are localized to the aleurone layer during imbib-
ition (27 h), when practically no MAN activity is detected in
the de-embryonated (endosperm) seed, suggesting a possible
accumulation of these transcripts and their corresponding
proteins as inactive forms in the aleurone cells. Interestingly,
the BdMAN4 deduced protein sequence has a predicted sig-
nal peptide for the secretory pathway and it is possible that
the BdMAN4 isozyme could be transported later on during
post-germinative reserve mobilization from the aleu-
ron to the endosperm cells through the apoplastic space.
In Arabidopsis the AtMAN7 and in poplar the PtrMAN6
proteins also contain signal peptides and have been localized
to the apoplast, indicating that the mature MANs could be
mobilized to the outer space (Iglesias-Fernández et al., 2013;
Zhao et al., 2013).

Several authors have proposed that polysaccharides
(β-1,3-1,4-glucans, mannans and others) present at the
endosperm CWs of the Poaceae grains, not only have a struc-
tural function, but also a storage role (Guillón et al., 2012).
Heteromannans, although globally less abundant in these
seeds than the β-glucans, are concentrated not only in the
colorhiza and in the root but also these polymers are found
in the aleurone layer and storage endosperm of B. distachyon
(Guillón et al., 2011). In this context, our data demonstrate
that MAN activity is important for the weakening of the
colorhiza cell walls and for the expansion of the root cells,
thus facilitating germination sensu stricto, but it may be also
important for the mannans hydrolysis in endosperm during
post-germinative reserve mobilization.

Supplementary data

Supplementary data are available at JXB online.

Supplementary Fig. S1. Transcription levels of the house-
keeping (BdGAPDH gene), presented as Ct mean values, in
different organs, in developing seeds and during seed germi-
nation of B. distachyon.

Supplementary Fig. S2. Transcripts analysis by RTqPCR
of the BdMAN1-6 genes in different organs.

Supplementary Fig. S3. Expression analysis by RT-qPCR
of BdMAN1-6 genes in developing seeds and germinating
seeds.

Supplementary Table S1. Major biochemical characters-
istics of Brachypodium distachyon and Oryza sativa endo-β-
mannanase proteins.

Supplementary Table S2. Oligonucleotide sequences,
amplicon length and PCR efficiency of the primers used for
RT-qPCR analyses.

Supplementary Table S3. Primers used for the synthesis of
the in situ mRNA hybridization probes.

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