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REVIEW ARTICLE

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Study of Structural Cell Wall Proteins and Hydroxyproline Rich Glycoproteins (HRGPs)

Hamid Kheyroodin and *Khosro Ghazvinian

Faculty of Desert Science-Semnan University, Iran

*Faculty of Veterinary and Animal Science-Semnan University, Iran

ABSTRACT

Boron (B) is an essential nutrient for normal growth of higher plants, and B availability in soil and irrigation water is an important determinant of agricultural production. The most severe effect derived from a deficiency of B is the alteration of cell wall biogenesis and pollen germination. There is increasing evidence that B is required for the maintenance of the structure and functions of membranes and, especially, plasma membrane. We investigated pollen of plant species that require B for germination as well as those that can germinate without B. In both groups, B addition in the growth medium increased the length of the pollen tube after germination. Hydroxyproline Rich Glycoproteins (HRGPs) are the most abundant cell wall structural proteins of dicotyledonous plants and the sexual tissues of monocotyledonous plants. Maize pollen germinated in the presence of B accumulated soluble and non-soluble Hyp in the pollen tube wall in contrast to maize pollen germinated without B. were preferentially localized in the pollen tube of maize, not in the pollen grain itself. Hence, B might affect the assembling of HRGPs in the wall of pollen tubes grown in vitro.

Keywords: Boron, Boron transporter, Maize, Germination, Cell wall-related genes and HRGPs.

INTRODUCTION

When a plant is infected by a pathogen its physiology is impaired and especially nutrient uptake, assimilation, translocation from the root to the shoot and also utilization. Plant cell walls are a structural metabolically dynamic complex that contains a vast amount of polysaccharides and proteins with unusual physical and chemical properties. Hydroxyproline Rich Glycoproteins (HRGPs) are the most abundant cell wall structural proteins in dicotyledonous plants and sexual tissues of monocotyledonous plants (Sommer-Knudsen et al., 1997; Cassab and Varner, 1998). They contain highly repetitive sequence domains in which hydroxyproline (Hyp) is a major amino acid (Showalter, 1993). Extensins are the most extensively studied HRGPs of plants, making up 1-10% of the primary cell wall (Cassab and Varner, 1988). Boron (B) is an essential micronutrient element for vascular plants and diatoms; however, its mechanism of action is not yet understood (Matoh 1997; Blevins and Lukaszewski, 1998). Many studies have implicated B in the biogenesis of plant membranes and cell walls (Goldbach et al., 1991; Bonilla et al., 1997). Deficiency symptoms first appear in actively growing tissues, within hours in root tips, and within minutes or seconds in pollen tube tips, and are characterized by cell wall abnormalities (Loomis and Durst, 1992; Blevins and Lukaszewski, 1998). However, it has been observed that in most plant species the B requirement for reproductive growth is much higher than for vegetative growth. The most dramatically rapid manifestation of B deficiency is seen in pollen tube growth. Without B, the pollen tubes of most species burst explosively, always at the outermost tip, ejecting large amounts of their contents. Growth of pollen

tubes requires a continuing presence of B. Pre-germination treatment with B stimulates germination of pollen but does not maintain the integrity of the growing pollen tubes (Schmucker 1933, 1934). In general, Schmucker's results support a role of B as a cell wall cross-linker since B deficiency occurred specifically at the pollen tube tips, showing weakness of the newly formed wall. In fact, B is cross-linked with the rhamnoglacturonan (RG) II fraction of pectic polysaccharides in many plant species (Hu and Brown 1994; Kobayashi et al., 1996; Matoh et al., 1996 O'Neill et al., 1996). On the other hand, other candidates for borate crosslinking in primary cell walls are HRGPs and PRPs (Bonilla et al., 1997). However B seems to participate in membrane structure since it can form complexes with membrane constituents such as glycolipids and/or glycoproteins (Shkolnik, 1984; Cakmak et al., 1995). So far, the results of pollen growth studies are consistent with B-complexing cell wall polymers, while the lipid thermostability results show that B is important in membrane structure and function (Jackson, 1989). Whatever the mechanism, the role of B in reproductive growth is particularly striking. Pollen grains of most species are naturally low in B, but in the styles, stigma, and ovaries, B concentration is usually high (Gauch and Dugger, 1954). However, there have been too few B measurements to substantiate this claim (Dell and Huang, 1997). The uniformly high B requirement for reproductive growth across the plant kingdom is intriguing and indicates similarities between reproductive structures, so unlike other plant cell walls perhaps the composition of the pollen tube is similar across plant species (Blevins and Lukaszewski, 1998). Pex 1 is a pollen-specific extensin-like domain protein,

involved in pollen tube growth that is expressed in starch-filled (mature) and in germinating pollen of maize (Rubinstein, et al., 1995b). Hence the pollen tube constitutes a good model to study processes related to cell wall protein assembly. In our study, we report that B affected the deposition of HRGPs into the pollen tube cell wall. This information may be important for understanding the mechanisms by which B controls reproductive growth in plants.

MATERIAL AND METHODS

Biological material and germination trials:

Fresh pollen from maize (*Zea mays L.*), tobacco (*Nicotiana tabacum*), *Impatiens sp.* and lily (*Lilium longiflorum.*) was immediately germinated after collection under two B conditions: B deficient medium (-B), (18% sucrose; 2mM CaCl₂; 0.5% agar) and B sufficient medium (+B) (18% sucrose; 2mM CaCl₂ and the following concentrations of boric acid: in *Impatiens sp.* 0.5 mM; in maize 1.0 mM; in tobacco 1.6 mM; pine 1.66 mM and in lily 0.162 mM. Pollen was considered to have germinated when the pollen tube was at least half as large as the grain length. Thirty trials were carried out on three different samples of each species; pollen observation was made with a stereoscopic microscope (Leika stereo zoom 6 Photo). The data on pollen germination and tube length were processed with the SAS software using the Tukey test for the comparison of means at $P \leq 0.05$.

Chemical analysis: All pollen samples were freeze-dried in a lyophilizer, when dry weights were determined. The quantity of hydroxyproline (Hyp_{Tot}), hydroxyproline extractable in CaCl₂ (Hyp_{Ext}), and non-soluble hydroxyproline (Hyp_{Ins}) in pollen tubes and pollen grains was determined by a colorimetric method (Drozd et al. 1976),

as follows: dry tissue was hydrolyzed in 6N HCl (3h at 120°C) and then the samples dried thoroughly under vacuum with a NaOH trap. The hydrolysates were resuspended in 100µl of H₂O and centrifuged in a microfuge to eliminate humus. To each sample and each HyPro standard (0.5µg and 0.05µg in 100 µl) 60 µl of the acetate-citrate buffer (0.1 g chloramine T; 1.0 ml H₂O; 2.0 ml acetate-citrate buffer) was added and 10 minutes later 120 µl of the second solution was added (1.0 ml of 72% perchloric acid; 5.0 ml PDMAB solution [6% w/v paradimethylaminobenzaldehyde (PDMAB) in isopropanol]. The tubes were then promptly placed in a hot water bath at 60°C for 20 minutes. This allows the colour to develop, and the absorbance was read in a spectrophotometer at 560 nm. The pollen Hyp_{Tot} level was obtained from grains hydrolyzed in 6N HCl (120°C, 3 h.).

Cell wall isolation

Pollen tubes were ground in a glass homogenizer and treated as described by Cassab et al. (1985).

Distribution of Hyp

The distribution of Hyp in pollen tubes was determined in germinated pollen from maize in the presence of 0.5 mM, 1.0 mM and 2.0 mM of boric acid for 2 h. First, pollen tubes were separated from pollen grains by briskly vortexing the solution, the pollen tubes were collected after filtering the solution through a 10 µm nylon mesh. To measure the quantity of Hyp_{Tot} the sample was hydrolyzed using 6N HCl (120°C, 3 h.), and evaluated by the colorimetric method (Drozd et al., 1976) described above. A fraction of the lyophilized sample was resuspended in 0.2 M CaCl₂ (Hyp_{Sol}) for 12 h. at 4°C, then the Hyp content was measured. The non-soluble material was lyophilized, weighed

and hydrolyzed (HCl 6N, 120°C, 3h), and Hyp_{Ins} was also quantified. The data for total hydroxyproline (Hyp_{Tot}), hydroxyproline extractable in CaCl₂ (Hyp_{Ext}), and non-soluble hydroxyproline (Hyp_{Ins}) were processed with the SAS software using the Tukey test for the comparison of means at $P \leq 0.05$.

Western blot and immunolocalization of HRGPs: Extractable proteins in 0.2 M CaCl₂ from pollen tubes grown on -B and +B were boiled in an equal volume of 2X SDS sample buffer (0.1M Tris, pH 6.8, 1.8% SDS, 18 [w/v] glycerol, 9% [w/v] 2-mercaptoethanol, 0.02% [v/v] bromophenol blue) and submitted to electrophoresis on SDS-PAGE gels (12%), using 4µg of protein for each sample. Protein content was determined by the Bradford method (1976) and silver nitrate was used as a stain. Proteins were electroblotted onto nitrocellulose paper (Hybond-C, Amersham) in a semi-dry medium (SemiPhor, Hoeffer) membrane, hydrated with a transfer buffer. The membrane was soaked with the primary HRGP antibody (Cassab and Varner, 1987) for 12 h (1:1300 dilution) dissolved in buffer (0.2% BSA, 0.1% Tween-20, 0.013 M Tris, pH 7.5, 0.15 M NaCl), washed three times in wash buffer (0.01% [v/v] tween-20, 0.013 M Tris pH 7.5, 0.15 M NaCl), and finally incubated for 1 hour with the secondary antibody conjugated to alkaline phosphatase (anti-rabbit IgG, Zymed AP-Goat rabbit IgG). Colour reaction took place in the presence of alkaline phosphatase buffer. For HRGPs immunolocalization, pollen tubes were fixed in FAA [10% (v/v) formaldehyde, 5% (v/v) acetic acid, and 50% ethanol] and mounted onto glass slides pre-treated with poly-lysine (Sigma). The primary antibody used was anti-HRGPs from soybean seed coats (Cassab and Varner, 1987) diluted. 1:50, and the immunogold

reagent 1:100 (ImmunoResearch), with which the sections were incubated 4 h, respectively, at room temperature. The secondary antibody was goat anti-rabbit IgG horseradish peroxidase diluted 1:100 with which sections were incubated for 4 h at room temperature, and the immunogold-silver staining performed as described by Springall et al. (1984). The negative control involved the substitution of the primary HRGP antibody with preimmune serum and anti-profilin antibody from bean, diluted 1:1000 and 1:50 respectively. Pollen tubes were visualized using a Zeiss Axioskop light microscope and photographed using ASA 100 Kodacolor film.

RESULTS

Micronutrients are those elements essential for plant growth which are needed in only very small (micro) quantities.

Table 1. Pollen germination of different plant species in (+ B) and (- B) *Impatiens* sp., 0.5 mM, maize 1.0 nM, tobacco 1.66 mM pine, lily 0.162 mM of boric acid. The data were means of six replicates.

Plant	Sufficient medium +B % Pollen germination	desufficient medium -B % Pollen germination
<i>Impatiens</i> sp.	14.7 (± 3.9) ^a	13.9 (± 4.3) ^a
<i>Zea mays</i>	75.1 (± 5.6) ^a	5.3 (± 2.3) ^b
<i>Pinus</i> sp.	74.6 (± 6.7) ^a	70.8 (± 6.9) ^a
<i>Nicotiana tabacum</i>	67.3 (± 3.1) ^a	21.1 (± 4.5) ^b
<i>Lilium longiflorum</i>	17.6 (± 3.6) ^a	13.1 (± 3.9) ^a

Means values in columns with the same letter are not significantly different, according to the Tukey test at $P < 0.05$.

Table 2. Pollen tube length (μm) of different species, for pollen grains germinated in boron deficient medium (-B) and boron sufficient medium (+B). Time (h) that pollen was incubated in the germination medium ¹(8), ²(2), ³(24), ⁴(9), ⁵(6).

Plant	Sufficient medium +B % Pollen germination	desufficient medium -B % Pollen germination
<i>Impatiens sp</i> ¹	260.5 (\pm 53) ^a	128.7 (\pm 61) ^b
<i>Zea mays</i> ²	457.3 (\pm 64) ^a	130.2 (+ 28) ^b
<i>Pinus sp.</i> ³	122.4 (\pm 20) ^a	61.5 (\pm 15) ^b
<i>Nicotiana tabacum</i> ⁴	624.7 (\pm 93) ^a	154.1 (\pm 47) ^b
<i>Lilium longiflorum</i> ⁵	12345.6 (\pm 234) ^a	704.1 (\pm 124) ^b

Means values in columns with the same letter are not significantly different, according to the Tukey test at $P < 0.05$.

Table 3. Total Hydroxyproline levels (Hyp_{tot} $\mu\text{g Hyp/mg dry weight}$) in pollen from different plant species.

Pollen	+ B
<i>Impatiens sp</i>	0.87 (\pm 0.03) ^c
<i>Zea mays</i>	1.01 (\pm 0.06) ^b
<i>Pinus sp.</i>	1.31 (\pm 0.07) ^a
<i>Nicotiana tabacum</i>	0.68 (\pm 0.03) ^d
<i>Lilium longiflorum</i>	0.89 (\pm 0.06) ^c

Means values in columns with the same letter are not significantly different, according to the Tukey test at $P < 0.05$.

These elements are sometimes called minor elements or trace elements, but use of the term micronutrient is encouraged by the American Society of Agronomy and the Soil Science Society of America. The micronutrients are **boron (B), copper (Cu),**

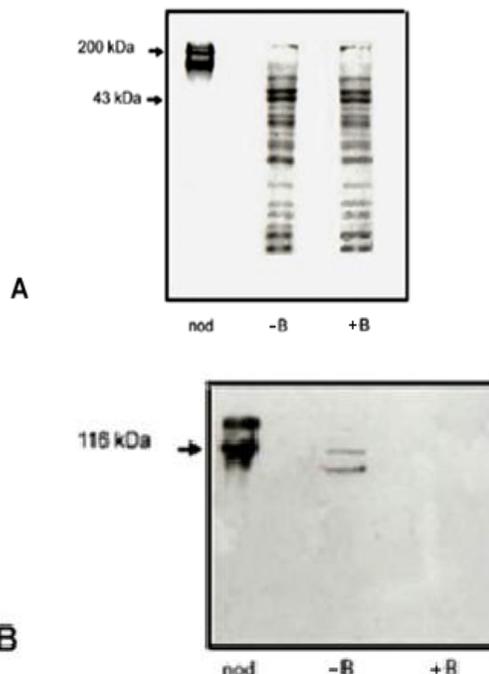
iron (Fe), chloride (Cl), manganese (Mn), molybdenum (Mo) and zinc(Zn).

Recycling organic matter such as grass clippings and tree leaves is an excellent way of providing micronutrients (as well as macronutrients) to growing plants. We studied the effect of B on *in vitro* pollen germination from different plants species. The results reported in [table 1](#) showed that pollen germination of maize and tobacco pollen grains increased when B was present in growth medium, and henceforth these species were designated B dependent (Bd), whereas *Impatiens sp.*, pine and lily pollen germination rates were less sensitive to B, and were designated B independent (Bi). However, the length of all pollen tubes analyzed was doubled in the presence of +B medium ([table 2](#)). The content of Hyp_{Tot} differed in pollen grains from maize, tobacco, pine and *Impatiens sp.* ([table 3](#)). The content of Hyp_{Tot} in pollen from maize and tobacco (Bd) turned out to be 30% greater than that of pine, *Impatiens sp.*, and lily (Bi). The value of Hyp_{Tot} for lily was similar to that of *Impatiens sp.* Therefore, there is not a strict correlation between high Hyp_{Tot} in pollen of Bd species as opposed to Bi species ([table 3](#)).

To evaluate the relationship between the B tube growth response and HRGP accumulation in the pollen tube wall, maize pollen was selected because of its: a) pollen tube size, b) high germination percentage and c) ease in separating the pollen tube from the pollen grain. We examined the distribution of Hyp in cell wall extracts, and isolated cell walls of maize pollen tubes ([table 4](#)).

Since an optimum B concentration increased the levels of Hyp in both Hyp_{Ext} (48%) and $\text{Hyp}_{\text{Insol}}$ (16%) with respect to the control fractions, it was then of interest to determine whether B modifies the

accumulation of HRGPs in pollen tubes. For this, a western blot analysis was performed on cell wall proteins of pollen tubes grown in the presence and absence of B, using a specific anti-HRGP antibody from soybean seed coats. This antibody is highly specific for HRGPs of both dicotyledonous and monocotyledonous plants, and does not cross react with arabino-galactan proteins, another class of HRGPs (Cassab and Varner, 1987). Cell wall fractions from pollen grains grown under -B conditions showed two high molecular weight bands. In contrast, the cell wall fraction from pollen grains grown on the +B medium did not contain polypeptides that were labeled by the HRGP antibodies (figure 1).



Means values in columns with the same letter are not significantly different, according to the Tukey test at P<0.05.

Figure 1. Characterization of extractable cell wall fractions from maize pollen tubes (A) cell wall salt-extractable proteins from maize pollen tubes grown in sufficient +B and deficient -B media Western blot analysis of maize pollen tubes. The blots were probed with anti-HRGPs antibodies from soybean seed coat. Note that two HRGP glycoprotein bands for material grown in media -B were not observed in +B.

To determine the localization of HRGPs in germinated maize pollen, we performed immunolocalization studies with anti-HRGP antibodies from soybean seed coats. In maize germinated pollen grains, HRGPs were primarily detected in the pollen tube wall which corresponds with the chemical analysis of Hyp in pollen tubes (table 4A). The detection of HRGPs in maize pollen tubes was very strong, although the

Table 4. Distribution of Hydroxyproline (Hyp) in maize pollen tube cell walls.

(A)	Total Hydroxyproline (Hyp _{ta}) µg Hydroxyproline/mg pollen tube dry weight	
0.5 mM H ₃ BO ₃ 0.53 (±0.03) ^c	1.0 mM H ₃ BO ₃ 0.63 (±0.03) ^a	2.0 mM H ₃ BO ₃ 0.63 (±0.03) ^a
(B)	Extractable hydroxyproline in CaCl ₂ (Hyp _{ta}) µg Hydroxyproline/mg pollen µg 0.2 M CaCl ₂	
0.5 mM H ₃ BO ₃ 0.13 (±0.02) ^c	1.0 mM H ₃ BO ₃ 0.25 (±0.03) ^a	2.0 mM H ₃ BO ₃ 0.16 (±0.02) ^b
(C)	Non-soluble hydroxyproline (Hyp _{ta}) µg Hydroxyproline/mg pollen tube dry weight	
0.5 mM H ₃ BO ₃ 1.73 (±0.02) ^c	1.0 mM H ₃ BO ₃ 4.15 (±0.03) ^a	2.0 mM H ₃ BO ₃ 2.69 (±0.03) ^b

antibody was not obtained from a monocotyledonous plant.

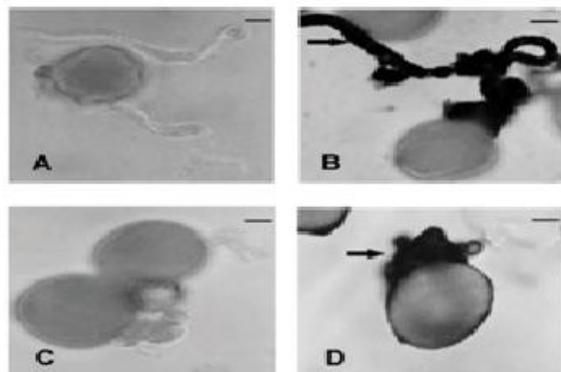


Figure 2. Immunolocalization of HRGPs in germinated pollen grains from maize. (A, C) germinated pollen grains incubated with preimmune serum. (B, D) germinated pollen incubated with anti-HRGPs antibodies from soybean seed coat. (A, B) pollen grains germinated in +B. Note that the labeling is associated with the cell wall (arrow) in (B) and the pollen tube burst without boron (D-arrow).

One of the major effects derived from the lack of B occurs during pollen germination and the formation of pollen tube (Loomis and Durst, 1992). Pollen grains of most species require a continuous supply of B (Steer and Steer, 1989; Taylor and Hepler, 1997). It has been reported that pollen from most plant species requires B to germinate (Bd); however pollen from some plant species germinates almost equally well in a B deficient environment (Bi) (table 1.) (Heslop-Harrison, 1971; Jackson, 1989). In addition, many physiological effects of B deficiency have been identified in the plant cell, such as impairments of sugar transport, membrane integrity, RNA metabolism, etc. (Heslop-Harrison, 1987; Cakmak and Römheld, 1997). The initial phase of germination includes processes that lead to

the onset of pollen tube growth, when Bd species seem to be more sensitive to B deficiency, since their germination percentages diminished 13 times in maize and 3 times in *Nicotiana* (table 1).

On the other hand, pollen tube length from plant species utilized in these studies was increased after the addition of B (table 2). Previous studies have shown that the wall in the tip region of the pollen tube is mainly composed of pectins (Mascarenhas, 1993). The synthesis and modification of the pectic network is an active process directly associated with the complex boron-RGII-calcium (O'Neill et al., 1996). This suggests that the elongation and growth of the pollen tube is a consequence of this association (Kobayashi, et al., 1999).

Reports suggest that the B requirement of plants is determined by the RGII content of its cell wall. It has been shown that one gram of monocot cell wall contains 6% of RGII and this could only covalently bind 8% mmol B considering dicot cell walls as 100% (O'Neill, 1996). Thus, the B pollen requirement could be determined by differences between pollen tube cell walls of Bi and Bd. In order to confirm this, we should evaluate a greater number of pollen plants species of the Bd and Bi types and measure the RGII fraction in both groups of plants. Hydroxyproline was present in all pollen grain species tested, but the content of Hyp_{Tot} in pollen from maize and tobacco turned out to be 30% greater than pine, *Impatiens sp.*, and lily. However, further evaluation must be carried out to confirm this fact. Previous studies have suggested that HRGPs are abundant in sexual tissues since the presence of higher amounts of Hyp in the pollen and pollen tube has been reported. (Li et al., 1994; Rubinstein et al., 1995a; Jauh and Lord, 1996). In addition to their structural role, it has been suggested

that pollen HRGPs could play roles in sexual recognition and signalling (Rubinstein, 1995b). HRGPs are the most abundant structural proteins in maize sexual tissues and since (Hyp) is an amino acid resulting from post-translational modification that characterizes HRGPs, their protein level can be indirectly estimated when the Hyp concentration is quantified (Cassab and Varner, 1987; Kieliszewski and Lamport, 1994). Measurement of hydroxyproline levels have been utilized primarily in the study of cell walls since only trace amounts of hydroxyproline are found in the cytoplasmic fraction (Rubinstein et al., 1995a). The level of Hyp in the pollen tube wall of maize changed depending upon the B concentration in the growing medium (table 4A-C). The importance of HRGPs in cell walls of pollen tubes was evident, since 63% of Hyp_{Tot} from the maize pollen is located in the tube. This suggests that most of the Hyp in the grain was used for the construction of the cell wall during germination (table 4A). This value was high, considering that the pollen tube length in pollen germinated *in vitro* was up to 70% shorter than the length reached *in vivo* (Taylor and Hepler, 1997). At an optimal B concentration (1 mM), the level of total, extractable and insoluble Hyp was larger than at the other B concentrations tested (table 4). This suggests that an optimal concentration of B can facilitate the accumulation of HRGPs in the cell wall (Bonilla et al., 1997).

Previous studies have described the biochemical and physiological relationship between B and pectic substances, although little is known about the effect of B in plant cell wall assembly of HRGPs (Matoth et al., 1996; Matoth et al., 1997). Western blot analysis of cell wall proteins from pollen tubes grown in the presence and absence of

B (figure 1), suggested that in the presence of B, HRGPs might be modified and hence the specific HRGP antibodies did not recognize them. This alteration could be due to insolubility of HRGPs in the wall in the presence of B, or to the formation of a linkage between B and HRGPs, or both. It has been reported that HRGPs are slowly insolubilized into the cell wall by a covalent link (Qi et al., 1995). Furthermore, previous data showed that HRGPs are not covalently bound to the walls of B deficient nodules of soybean root, suggesting that B may have a role in the assembly of some wall protein components in nodules (Bonilla et al., 1997). Although there is an abundance of HRGP in the pollen tubes, only the maize pollen extensin-chimera PEX1 has been so far molecularly characterized. PEX1 seems to play a role in pollen tube growth during pollination (Rubinstein, et al., 1995b). Further investigations are essential to explain the assembly of the HRGPs in growing pollen tubes, and their relation with the B-RGII complex.

The accumulation of HRGPs in germinated pollen of maize grown in -B and +B was studied by immunocytochemical analysis (figure 2). HRGPs are deposited in the emerging pollen tube wall in enormous quantities, but there is no deposition of these glycoproteins in the pollen grain. It confirms that HRGPs are synthesized previous to germination and deposited in the cell wall of the growing pollen tube (Rubinstein et al., 1995a,c). It has been proposed that the HRGPs might provide structural support for the pollen tube necessary for its rapid growth, and have a role in sexual recognition and signalling during the fertilization (Rubinstein, et al., 1995b). Thus, the study of HRGPs of sexual tissues might help to elucidate the fertilization processes in plants. It has been

established that a primary role of B is the cross-linking of cell wall RG-II and pectin assembly. Nevertheless hypothesize that the primary role of B in biological systems is stabilization of molecules with *cis*-diol groups, independently of their function. Thus, it has been suggested that B could be involved in membrane function by complex formation with glycoproteins. More recently, it has been proposed the possible role of B as a cellular signal through the interaction with transcription factors. Future research should be focused to identify the Bbindingligands as well as their functions in plant physiology.

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Corresponding author: Dr. Hamid Kheyroodin, Assistant professor Faculty of Desert Science, Semnan University, Iran

Email: hkheyroodin@yahoo.com