Cytological Evidence of an Active Role of Silicon in Wheat Resistance to Powdery Mildew (*Blumeria graminis* f. sp. *tritici*)

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ABSTRACT

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Silicon (Si) amendments in the form of exogenously supplied nutrient solution or calcium silicate slag protect wheat plants from powdery mildew disease caused by the fungus *Blumeria graminis* f. sp. *tritici*. The most striking difference between Si- and Si+ plants challenged with *B. graminis* f. sp. *tritici* was the extent of epidermal cell infection and colonization by *B. graminis* f. sp. *tritici*. Histological and ultrastructural

analyses revealed that epidermal cells of Si+ plants reacted to *B. graminis* f. sp. *tritici* attack with specific defense reactions including papilla formation, production of callose, and release of electron-dense osmophilic material identified by cytochemical labeling as glycosylated phenolics. Phenolic material not only accumulated along the cell wall but also was associated with altered integrity of haustoria in a manner similar to localized phytoalexins as reported from other pathosystems. These results strongly suggest that Si mediates active localized cell defenses against *B. graminis* f. sp. *tritici* attack.

It has long been known that silicon (Si) plays an important role in the resistance of plants against pathogens (19,39). The mechanisms by which Si mediates disease resistance is incompletely understood. Historically, it was assumed that insoluble Si, accumulating in the apoplast, creates a mechanical barrier preventing fungal penetration (23). This suggestion was thought to apply to many pathosystems including rice-rice blast, oat-powdery mildew, barley-powdery mildew, and cucumber-powdery mildew, etc. (7,33,40,42). Kunoh and Ishizaki (26) demonstrated localized accumulation of insoluble Si at penetration sites of powdery mildew pathogens. However, neither they nor Carver et al. (7) could directly relate the presence of insoluble Si to resistance to penetration. In a similar approach with the *Pythium*-cucumber pathosystem, Chérif et al. (10) demonstrated that although Si failed to accumulate at penetration points under conditions of saturated humidity, resistance was still enhanced. They concluded that, in cucumber, *Pythium* resistance did not result from insoluble Si accumulating within plant tissues and cell walls; rather, they (9) hypothesized that soluble Si played a physiological role resulting in upregulation of plant defenses.

Pursuant to the concept that Si mediates active defense mechanisms, Chérif et al. (8) showed that in the cucumber-*Pythium* interaction, supplying Si enhanced the activities of chitinases, peroxidases, polyphenoloxidases, and phenolics. Fawe et al. (15), using the cucumber-powdery mildew pathosystem demonstrated the production of specific phytoalexins associated with Si-mediated resistance. By comparing the properties of known responses occurring as a result of systemic acquired resistance with those occurring in the presence of soluble Si in planta, Fawe et al. (16) suggested that Si-mediated resistance shares functional similarities with forms of induced resistance.

In monocots, little emphasis has been placed on the possibility that Si plays a role in activating host defenses. Most researchers have simply correlated the presence of insoluble Si with increased resistance to fungal penetration (43). Although the hypothesis of a simple physical barrier has become widely accepted, alternatives for other functions of Si have evolved. For instance, Deren (12) reported a negative correlation between Si concentration in leaf tissues and plant disease development in different rice cultivars. In addition, Carver et al. (7) trying to correlate Si accumulation with fungal penetration, concluded that insoluble Si was not the only factor influencing the outcome of attempted penetration. This prompted Zeyen et al. (41) to speculate that high levels of soluble Si in cells at the time of attempted powdery mildew penetration may have a physiological or biochemical role in mediating cellular resistance.

In light of the uncertainty regarding the roles of Si in planta, we examined the cytological effects of Si amendments in the wheat (*Triticum aestivum* L.) and powdery mildew (*Blumeria graminis* DC f. sp. *tritici* Ém. Marchal) pathosystem. The objectives were (i) to confirm that Si amendments can influence *B. graminis* f. sp. *tritici* incidence and development in wheat; and (ii) to characterize and localize key cellular constituents occurring in Si treated (Si+) and Si untreated (Si-) wheat epidermal cells challenged by *B. graminis* f. sp. *tritici*.

MATERIALS AND METHODS

Plant material and silicon amendment. Seeds of wheat (cv. HY 644), donated by A. Comeau (Agriculture and Agri-Food Canada, Sainte-Foy Research Station), were sown in Pro-Mix (Tourbières Premier, Rivière-du-Loup, Quebec) in 10-cm pots (10 seeds per pot) and watered daily. Other seeds were sown in Pro-Mix to which Reclime (calcium silicate slag) (RECMIX Inc., Sorel, Quebec) was added at a rate of 3 g per pot or an equivalent of approximately 3 tonnes ha⁻¹. After 14 days, plants were transferred to a greenhouse compartment maintained at 21/18°C with a relative humidity of 70% and a photoperiod of 12 h with a photosynthetic photon flux of ≈1,000 μmol m⁻² s⁻¹. Plants were fertil-

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ized every other day with a standard Hoagland solution amended (Si+) or not amended (Si-) with 1.5 mM SiO₂ (Kasil 6, National Silicates, Toronto). Plants treated with Reclime were fertilized with a standard Hoagland solution only. For each treatment (Si-, Si+ and Reclime), a total of 120 plants (12 pots) were grown for 7 days before being inoculated. For controls, two pots per treatment were isolated in a separate greenhouse compartment and not subjected to inoculation.

Inoculation with *B. graminis* f. sp. *tritici* and disease assessment. One day before inoculation, wheat plants maintained in a growth cabinet and heavily infested with *B. graminis* f. sp. *tritici* were shaken to displace older spores ensuring that freshly formed conidia were available. The following day, 10 leaf segments bearing conidia were harvested, and for each pot, one segment was used to inoculate the experimental plants by gently brushing the segment over them. Inoculated plants were then observed daily for powdery mildew development. At the first visible sign of the fungal colonies, plants were scored for fungal presence on a scale of 0 to 4, where 0 = no colonization; 1 = 1 to 5% of leaf area covered; 2 = 6 to 20%; 3 = 21 to 40%; and 4 = 41%+. All 10 plants from each pot represented one experimental unit. Assessments were made every week for 5 weeks. Data were subjected to analysis of variance (ANOVA) using Super ANOVA version 1.11 (Abacus Concepts Inc., Berkeley, CA).

Tissue processing for light and transmission electron microscopy. Samples (20 mm²) were excised from control and inoculated leaves (Si- and Si+ plants) from sites where the fungus grew and pre-embedded in 2% aqueous Bacto-agar in order to hold fungal structures in place. They were then immersed in 3% (vol/vol) glutaraldehyde in 0.1 M sodium cacodylate buffer, pH 7.2, for 2 h at room temperature and postfixed with 1% (wt/vol) osmium tetroxide in the same buffer for 1 h at 4°C prior to dehydration in a graded ethanol series and embedding in Epon 812. Thin sections (0.7 µm), cut from the Epon-embedded material with glass knives, were mounted on glass slides and stained with 1% aqueous toluidine blue prior to examination with a microscope (Zeiss Axio-scope; Carl Zeiss Canada, Don Mills, Ontario, Canada). Ultrathin sections (0.1 µm), collected on nickel grids, were either stained with uranyl acetate and lead citrate for immediate examination with a transmission electron microscope (JEOL 1200 EX; JEOL, Tokyo) operating at 80 kV or further processed for cytochemical labeling with probes for cellulose, chitin, callose, phenolics, and glycosylation. For each treatment, a minimum of five samples from three different leaves were investigated. For each sample, 10 to 15 ultrathin sections were examined under the electron microscope.

Preparation of the gold-complexed probes. Colloidal gold with particles averaging 12 nm in diameter was prepared according to Frens (17) using sodium citrate as a reducing agent. For the localization of cellulosic β-1,4-glucans, an exoglucanase (β-1,4-D-glucan cellobiohydrolase, EC 3.2.1.21), purified from a cellulase produced by the fungus *Trichoderma harzianum*, was complexed to colloidal gold at pH 9.0 and used in a one-step procedure (5). Wheat germ agglutinin (WGA), a lectin with *N*-acetylglucosamine binding specificity was used for localizing *N*-acetylglucosamine residues (chitin) according to a previously described procedure (2). Because of its low molecular weight, the lectin could not be directly complexed to colloidal gold. It was used in a two-step procedure, using ovomucoid, conjugated to gold at pH 5.4, as a second step reagent. A β-1,3-glucanase, purified from tobacco reacting hypersensitively to *Tobacco mosaic virus*, was used for localizing β-1,3-glucans (callose) according to a recently described method (3). The enzyme was conjugated to colloidal gold at pH 5.5. Localization of β-glucosides (for glycosylated constituents) was performed by using a commercial β-glucosidase from almond (1). The enzyme was complexed to colloidal gold at pH 9.3. Localization of lignin-associated phenolic compounds was performed by using a laccase (EC 1.10.3.2) purified from the white rot fungus *Rigidoporus lignosus* (18). The enzyme was complexed to col-

loidal gold at pH 4.0, a pH value close to its isoelectric point reported to be 3.83 (6). All gold complexes were stored at 4°C until use.

Cytochemical labeling. For the direct labeling of cellulosic β-1,4-glucans, β-1,3-glucans, β-glucosides, and phenolic compounds, ultrathin sections were first incubated on a drop of phosphate-buffered saline (PBS) containing 0.02% (wt/vol) polyethylene glycol (PEG) 20000 for 5 min at room temperature. The pH of the PBS-PEG was adjusted according to the pH of optimal activity of each protein (pH 6.0 for exoglucanase, β-1,3-glucanase, and laccase; pH 7.0 for β-glucosidase). Sections were then transferred to a drop of the gold-complexed probe for 30 min at room temperature in a moist chamber. After washing with PBS and rinsing with distilled water, grids were stained with uranyl acetate and lead citrate.

For the indirect labeling of *N*-acetylglucosamine residues, sections were first floated on a drop of PBS, pH 7.4, for 5 min, and transferred to a drop of WGA (25 µg/ml in PBS, pH 7.4) for 60 min at room temperature in a moist chamber. After washing with PBS, pH 7.4, sections were incubated on a drop of the ovomucoid-gold complex (1:30 in PBS-PEG, pH 6.0) for 30 min at room temperature. Sections were washed with PBS, rinsed with distilled water, and contrasted as described previously.

Cytochemical controls. Specificity of labeling was assessed using the following control tests: (i) incubation with the gold-complexed enzymes to which were previously added the corresponding substrates (β-1,4-glucans from barley for the exoglucanase, laminarin or laminaribiose for the β-1,3-glucanase, alicin for the β-glucosidase, and *p*-coumaric acid, ferulic acid, or sinapinic acid for the laccase, 1 mg/ml in PBS, pH 7.2); (ii) substitution of the protein-gold complex under study by bovine serum albumin-gold complex to assess nonspecific adsorption of the protein-gold complex to tissue sections; (iii) incubation of tissue sections with protein-gold complexes under nonoptimal conditions for biological activity; (iv) incubation with the WGA with excess *N*-*N'*-*N''*-triacylchitotriose (1 mg/ml in PBS); and (v) incubation with WGA followed by unlabeled ovomucoid and finally by the ovomucoid-gold complex.

Reagents. The exoglucanase was provided by C. Breuil, Forintek, Canada, and the laccase was obtained from M. Nicole, ORSTOM, Montpellier, France. Tetrachloroauric acid was purchased from

Fig. 1. Effect of silicon amendments (Si+) on powdery mildew (*Blumeria graminis* f. sp. *tritici*) incidence on wheat leaves cv. HY 644. **A**, Non-inoculated Si- leaf; **B**, inoculated Si+ leaf; and **C**, inoculated Si- leaf.

BDH Chemicals, Montreal, Canada. All other reagents for electron microscopy were obtained from JBEM Chemical Co., Pointe-Claire, Québec, Canada.

RESULTS

Disease assessment. First signs of *B. graminis* f. sp. *tritici* colonization were observed on control plants (Si-) 5 days after inoculation. Disease progressed rapidly thereafter and control plants were heavily infected with powdery mildew colonies after 5 weeks (infection score = 3.71 ± 0.16). By contrast, Si+ plants and Reclime plants (calcium silicate-Si+) had very few *B. graminis* f. sp. *tritici* colonies even after 5 weeks (Fig. 1). At the end of the experimental period, Si+ and Reclime plants had an average infection score of 0.41 ± 0.06 and 0.43 ± 0.12 , respectively, indicating very limited powdery mildew fungal development.

Light microscope observations. Rate and extent of fungal colonization. Light microscopy of Si- and Si+ wheat leaves showed striking differences in the extent of *B. graminis* f. sp. *tritici* colonization (Fig. 2). In Si- leaves, *B. graminis* f. sp. *tritici* hyphae were abundant on the epidermal surface and freely penetrated the epidermis to form haustoria in $\approx 90\%$ of the epidermal cells observed (Fig. 2A). By contrast, in Si+ leaves, the degree of fungal colonization was markedly reduced from that in control leaves. Fungal growth in planta was mainly restricted to a few epidermal cells (approximately 10%) (Fig. 2B). In addition, *B. graminis* f. sp. *tritici* penetration of the leaf epidermis coincided with cytological changes that were associated with poorly developed haustoria (Fig. 2B, arrows).

Epidermal cell responses. Conidial germination on the epidermal surface led to the formation of appressoria and narrow penetration pegs that penetrated the host cuticle and the underlying

Fig. 2. Light micrographs from wheat leaves infected by *Blumeria graminis* f. sp. *tritici*. **A**, In the absence of silicon (Si-), the fungus has produced abundant hyphae on the cell surface and frequently penetrated the epidermis (E) to form digitate haustoria (H). Bar = 10 μ m. **B**, Upon treatment with Si (Si+), fungal growth in the epidermis (E) was restricted to a few epidermal cells. The haustorial structures are poorly developed (arrows). Bar = 10 μ m.

epidermal cell walls. Enlargement of the infection pegs in the epidermal cells resulted in the development of multilobed haustoria (Fig. 3A). In transverse sections, haustorial lobes often appeared disconnected from one another (Fig. 3A), whereas in longitudinal sections, the haustoria appeared elongated with typical finger-like lobes (Fig. 3B, arrow). In such heavily infected epidermal cells from Si- plants, defense reactions including structural barriers and accumulation of newly formed material were rarely seen, with the singular exception of haustorial neck collars (Fig. 3A).

In Si+ leaves, haustoria were abnormally shaped (Fig. 3C, arrow), correlating with a number of epidermal cell reactions that apparently retarded or prevented fungal development and colonization. These reactions involved deposition of a granular material in epidermal cells (Fig. 3C) and papillae in the subtending fungal penetration sites (Fig. 3D). Papillae formed in the reacting epidermal cells were intensely stained with toluidine blue and appeared usually as hemispherical (Fig. 3D).

Light microscopy provided evidence that wheat plants supplied with Si reacted with epidermal cell responses that likely protected their epidermal cells from extensive fungal invasion. A more pre-

cise characterization at the ultrastructural level was then deemed essential to understanding how these epidermal cell responses may have resulted in restricting *B. graminis* f. sp. *tritici* growth and development.

Ultrastructural and cytochemical observations. *Si- wheat leaves.* Examination of *B. graminis* f. sp. *tritici*-infected samples from Si- wheat leaves by transmission electron microscopy confirmed the extensive *B. graminis* f. sp. *tritici* colonization seen by light microscopy (Fig. 3A and B). *B. graminis* f. sp. *tritici* hyphae grew extensively on the leaf epidermal surface, and penetrated cells had haustoria with elongated necks surrounded by a thick collar composed of layers of differing electron density (Fig. 4A). Haustoria had long, ovoid central bodies with extending finger-like or digitate lobes. Both the haustorial body and the lobes contained a dense polyribosome-rich cytoplasm in which numerous organelles including mitochondria, vesicles, and dark inclusions could be seen (Fig. 4A to D). In all cases, haustoria were enveloped in a membrane, the extrahaustorial membrane, derived from the host epidermal cell plasma membrane (Fig. 4A and B). Occasionally, a thin layer of host cytoplasm could be visualized

Fig. 3. Light micrographs of samples from wheat leaves infected by *Blumeria graminis* f. sp. *tritici*. **A and B,** In Si- wheat leaves, penetration of *B. graminis* f. sp. *tritici* (Bg) in the epidermis (E) leads to the formation of haustoria (H) with finger-like lobes (**B**, arrow). Bars = 10 μ m. **C and D,** In Si+ wheat leaves, haustoria (H) are poorly developed and epidermal cells fill with granular materials (**C**, arrow). **D,** Papillae or wall appositions (WA) are formed in the epidermal cells beneath fungal contact and appressorial penetration sites. Bars = 10 μ m.

Fig. 4. Transmission electron micrographs of the interaction between Si- wheat leaf tissues and *Blumeria graminis* f. sp. *tritici*. **A and B.** The fungus penetrates the epidermis (E) and the elongated haustorial neck (HN) is surrounded by a thick collar of host cell material origin. The haustorial body (H) is encircled by the extrahaustorial membrane (EHM). Bars = 2 μ m. **C.** Upon incubation with the β -1,4 exoglucanase-gold complex for the localization of cellulosic β -1,3-glucans, the host cell wall (HCW) is intensely labeled whereas the fungal structures including the extrahaustorial membrane (EHM) and the extrahaustorial matrix (EHMa) remain unlabeled. Bar = 0.5 μ m. **D.** Incubation with the wheat germ agglutinin/ovomucoid-gold complex for the localization of chitin results in a specific labeling of *B. graminis* f. sp. *tritici* haustorial cell walls (FCW). The extrahaustorial membrane (EHM) and the extrahaustorial matrix (EHMa) remain unlabeled. Bar = 0.5 μ m.

adjacent to the extrahaustorial membrane (Fig. 4C). The extrahaustorial matrix appeared as a network of intermingled fibrils surrounding the polymorphic lobes (Fig. 4C).

Incubation of sections with β -1,4 exoglucanase-gold complex to reveal cellulose resulted in a heavy deposition of gold particles over the epidermal cell walls; whereas the cytoplasm and fungal structures were unlabeled (Fig. 4C). Upon labeling for chitin with the WGA/ovomucoid-gold complex, gold particles were uniformly distributed over the fungal cell walls. By contrast, the extrahaustorial membrane and the extrahaustorial matrix were free of labeling (Fig. 4D).

Si+ wheat leaves. As with light microscopy, penetration attempts in Si+ leaves were often associated with the deposition of wall papillae (Fig. 5A and B). The papillae usually consisted of an amorphous matrix impregnated by osmiophilic substances and were seldom penetrated by the fungus (Fig. 5A). However, in other cases in which the fungus successfully entered the host epidermal cell wall, an osmiophilic electron-opaque material, reacting positively for the presence of β -1,3-glucans (callose constituents), was found to entirely coat the embryonic haustorial structure (Fig. 5B). Interestingly, a number of vesicles containing β -1,3-glucans (callose) were always seen in the vicinity of the coating material (Fig. 5B, arrows). Both the impregnated papillae and the newly formed material encasing haustoria were apparently effective in reducing fungal ingress because well-developed normal haustoria were seldom found.

Another epidermal cell reaction in Si+ plants was the deposition of an osmiophilic electron-dense material lining epidermal cell walls near sites of potential fungal penetration (Fig. 5C and D). This material, which reacted positively with the gold-complexed laccase for the presence of phenolic compounds, usually extended inward to form polymorphic deposits that were nearly unlabeled (Fig. 5D). When epidermal cells were colonized by *B. graminis* f. sp. *tritici*, similar electron-dense deposits were detected (Fig. 6A and B). Observations at high magnification revealed that the deposits accumulated not only in the cell lumen but also in the epidermal cell walls (Fig. 6B, arrow), and along the extrahaustorial membrane (Fig. 6B, double arrows), forming, at an advanced stage of the interaction, a nearly uniform coating (Fig. 6C). A close examination of a large number of leaf sections revealed that the electron-dense material not only surrounded the extrahaustorial membrane but also appeared to infiltrate the extrahaustorial matrix to form a dense layer over the fibrillar network (Fig. 6C). Haustoria, encased in this osmiophilic material, had various anomalies ranging from increased vacuolation (Fig. 6C) to complete disorganization (Fig. 7A). Upon incubation with the WGA/ovomucoid-gold complex for the presence of chitin, a regular deposition of gold particles occurred on the walls of haustoria surrounded by the osmiophilic phenolic material (Fig. 6D).

At an advanced stage of colonization, large deposits, apparently resulting from the fusion of electron-dense droplets containing phenolics, established intimate contact with haustoria (Fig. 7A). Haustoria, encased in impregnated extrahaustorial membranes, were highly altered and, in most cases, it was difficult to clearly identify them (Fig. 7A, arrows). In the vicinity of such highly disorganized haustoria, a large number of extrahaustorial membrane remnants were the only indication of previously living haustoria (Fig. 7A, double arrows).

Upon incubation with gold-complexed β -glucosidase to identify glycosylated materials, labeling was found to be predominantly associated with the electron-dense aggregates occurring in the lumen of reacting Si+ epidermal cells and to a lesser extent in conjunction with extrahaustorial membranes (Fig. 7B). Gold particles could also be seen in the fungal cytoplasm. Labeling for phenolic compounds with the laccase-gold complex resulted in deposition of gold particles over the electron-dense aggregates, whereas all other structures were very slightly labeled (Fig. 7C).

Results from this study provide strong evidence that silicon amendments mediate an effective protection against wheat powdery mildew fungal infection and colonization. This report confirms numerous observations on the beneficial role of Si in monocots (14). However, few previous studies have focused directly on the wheat-powdery mildew pathosystem with the exception of Leusch and Buchenauer (30). Of additional interest is the fact that disease protection on wheat plants occurred whether silicon was supplied in a nutrient solution or in the form of calcium silicate slag (Reclime).

The practice of amending soils with calcium silicate slag has other positive effects for many plant species including higher yields and disease resistance in several crops (11). Several recent reviews have discussed the possible physiological role of Si, but these remain controversial and poorly understood (13,14,23). In this study, microscopical analyses of the wheat-*B. graminis* f. sp. *tritici* interaction have described cytological phenomena associated with reduced infection levels correlated with Si amendments.

The first difference noted between Si- and Si+ plants inoculated with *B. graminis* f. sp. *tritici* was the extent of fungal colonization. This observation is noteworthy because it demonstrates that although *B. graminis* f. sp. *tritici* is capable of infecting epidermal cells of Si+ plants, its rate of development was altered. To date, epidermal cells of cereals were understood to have two active defense mechanisms against *B. graminis* f. sp. *tritici* attack: (i) penetration resistance associated with papilla formation; and (ii) hypersensitive epidermal cell death (HR) (38,43). Our observations indicate that there is yet a third line of active defense characterized by a marked accumulation of phenolic compounds in infected Si+ epidermal cells.

Considering the first active defense reaction, papilla formation, our observations confirm the association of papillae with penetration resistance by wheat cells against *B. graminis* f. sp. *tritici*. As discussed by Zeyen et al. (43), papillae relate to powdery mildew fungal penetration in two distinct ways: (i) failed penetration; and (ii) successful penetration where papillae become haustorial neck collars. Both outcomes were observed in this work, in both Si+ and Si- plants, confirming that deposition of papilla formation may not be sufficient by itself to explain the level of protection observed with Si+ treatment. However, the prevalence of papillae in Si+ treatment compared with Si- treatment does suggest that Si has somehow stimulated their formation.

In Si+ wheat epidermal cells, papillae were observed to form on the inner surface of epidermal cell walls and were successful in preventing penetration by only a proportion of penetration pegs. Kunoh and Ishizaki (26-28), Kunoh et al. (29), and Carver et al. (7) found that Si deposition occurred in barley epidermal cells beneath sites of attempted penetration by *B. graminis* f. sp. *hordei* and that Si accumulated in the halo, papilla, and haustorial neck and collar areas irrespective of the outcome of the attempted penetration. Samuels et al. (35,36) reported similar results with the powdery mildew-cucumber pathosystem, suggesting that insoluble Si deposition is a common phenomenon in both dicots and monocots. In a different pathosystem, Chérif et al. (10) concluded that the increased resistance of cucumber to *Pythium ultimum*, induced by exogenously supplied Si, was not associated with insoluble Si accumulation at sites of pathogen penetration, regardless of the plant organ studied. They concluded that accumulation and polymerization of insoluble Si at fungal penetration sites or in epidermal cell walls did not appear to be a physical barrier against fungal attack. A similar conclusion was raised by Kunoh and Ishizaki (26) and Carver et al. (7).

The encasement of haustoria by callose (Fig. 5B) suggests that the polysaccharide can play an important role in preventing haustorium formation. Accumulation of callose in papillae has been previously linked to high Si content in epidermal cells (43). How-

Fig. 5. Transmission electron micrographs of the interaction between Si⁺ wheat leaf tissues and *Blumeria graminis* f. sp. *tritici* (Bg). **A**, An unsuccessful attempt of the pathogen to penetrate a papilla or wall apposition (WA) is observed. Bar = 0.5 μ m. **B**, Amorphous material (AM), reacting positively with the β -1,3-glucanase-gold complex for the localization of callose, envelops the haustorial structure (H). A number of labeled vesicles (V) are seen in the vicinity of the coating material (arrows). Bar = 0.25 μ m. **C and D**, Besides the formation of wall appositions (WA), osmiophilic deposits (OD) fill the epidermal cells and accumulate in the host cell lumen and in the apoplastic space. **D**, The material lining the host cell wall (HCW) reacts positively with the gold-complexed laccase for the localization of phenolics, whereas the polymorphic deposits in the cell lumen are mostly unlabeled. **C**, Bar = 2 μ m; **D**, Bar = 0.25 μ m.

ever, while callose is invariably found in papillae, Smart et al. (37) presented evidence that callose was not an effective barrier to penetration by *B. graminis* f. sp. *tritici* in barley epidermal cells. Zeyen et al. (41) suggested that callose in barley and oat epider-

mal cells was simply a matrix for other constituents. Whether or not callose plays a different role in wheat epidermal cells is unknown but our results support a direct role for callose in altering *B. graminis* f. sp. *tritici* haustorium development in Si⁺ plants.

Fig. 6. Transmission electron micrographs of the interaction between Si⁺ wheat leaf tissues and *Blumeria graminis* f. sp. *tritici*. **A**, A large number of osmiophilic deposits (OD) occur in epidermal cells colonized by the fungus. **B**, These deposits accumulate in epidermal cell walls (HCW, arrow) and along the extrahaustorial membrane (EHM, double arrows). **A**, Bar = 2 μ m; **B**, Bar = 1 μ m. **C**, The osmiophilic material apparently infiltrates the extrahaustorial matrix (EHMa) to form a dense layer over the fibrillar network. **D**, An extrahaustorial membrane (EHM) is entirely coated by osmiophilic deposits (OD). **D**, Upon incubation with the chitin-specific wheat germ agglutinin/ovomucoid-gold complex, a regular deposition of gold particles occurs over the fungal cell walls (FCW). **C and D**, Bars = 0.5 μ m.

Fig. 7. Transmission electron micrographs of the interaction between Si⁺ wheat leaf tissues and *Blumeria graminis* f. sp. *tritici*. **A**, The osmiophilic deposits (OD) are in close contact with altered haustorial bodies (H, single arrows). In the vicinity of such highly disorganized haustorial bodies, a number of extrahaustorial membrane remnants are the only indication of previously living fungal entities (double arrows). Bar = 0.5 μ m. **B and C**, The osmiophilic deposits (OD) occurring in the lumen of reacting epidermal cells and, to a lesser extent, associated with the extrahaustorial membranes contain glycosides and are intensely labeled with the gold-complexed β -glucosidase. **C**, Such deposits have apparent phenolic constituents because they are slightly labeled with the laccase-gold complex. Bars = 0.5 μ m.

In previous ultrastructural studies of other cereal powdery mildews, HR was reported as a common phenomenon in incompatible interactions (20). Although in this work some epidermal cells showed degrees of degeneration, suggesting cell death (Fig. 3C), the possibility that HR occurred as a main line of defense remains difficult to establish on the basis of our observations. Rather, the degradation of fungal structures within epidermal cells of Si+ plants revealed specific active defense responses.

The greatest cytochemical difference between Si- and Si+ plants was the extensive deposition of phenolic compounds in the infected epidermal cells of Si+ plants. These phenolic compounds accumulated along the epidermal cell wall and on the extra-haustorial membrane. Silicon, in the form of Si(OH)₄, possesses strong affinities for organic polyhydroxyl compounds which participate in the synthesis of lignin (21,22). This affinity of Si may explain its tendency to accumulate in mature cell walls of plants or during a pathogen attack. Maturity and pathogen attack correspond with a radical change in the constitution of the cell wall, with the infusion of lignin (21,22,24,31,34). Once polymerized, Si participates in the rigidification of cell walls with lignin, a process that is confined to the periphery of the plant where evapotranspiration occurs.

In the *Erysiphe graminis* f. sp. *hordei*-barley pathosystem, Koga et al. (25) hypothesized that during HR, phenolics were released and accumulated on cell walls and if soluble Si was present, it would form insoluble complexes with the phenolics. In such cases, it was suggested that insoluble Si was secondary to HR and played a passive role. However, in the current study, given that the host cells were already infected and that the phenolics appeared to have a direct and toxic effect on haustoria, it is unlikely that Si is simply playing a passive role by combining with phenolics to make epidermal cell walls more rigid against pathogen penetration. Rather, the fact that phenolic compounds were localized in close association with degraded haustoria (Fig. 7) suggests a complex interaction between soluble Si and production of metabolites involved in the protection of the cell.

To our knowledge, this is the first report linking Si amendments with cytological events elicited in cell defense reactions of monocots. Previous studies with dicots, namely cucumber, suggested involvement of phenolics in Si-mediated resistance (9,32). In addition, Chérif et al. (8) was able to show a marked increase in the concentration of antifungal phenolic materials in *P. ultimum*-infected Si+ cucumber tissues. Further biochemical analyses identified some of these metabolites as flavonoids and phenolic acids that accumulated specifically and strongly in a manner typical of phytoalexins (15). The fungitoxicity of these phenolic compounds was more apparent after acid hydrolysis. Therefore, Benhamou and Bélanger (4) suggested that the release of aglycones in the plant could be regulated in part by the pathogen itself. They showed that only the conjugated phenolics in close proximity to fungal structures were converted to their active state, presumably by *Pythium* or *Pythium*-induced plant enzymes. Interestingly, a similar phenomenon may be happening in the epidermal cells of Si+ wheat infected with *B. graminis* f. sp. *tritici*. The phenolic materials associated with and near the epidermal cell walls were strongly labeled with the gold-complexed β -glucosidase (Fig. 7b), indicating that they were in a glycosylated form; conversely, phenolic material in the cell near the extra-haustorial membrane showed much less labeling for glycosylated phenolics, suggesting that these phenolics were not heavily glycosylated and were likely in a fungitoxic form.

In conclusion, this work provides evidence that Si amendments to wheat plants mediate *B. graminis* f. sp. *tritici* resistance in more complex ways than originally thought. More specifically, the production of phenolic materials in powdery mildew-infected leaves of Si+ wheat plants suggests an active role for Si in mediating powdery mildew resistance. These results also suggest that the protective role of Si may be similar in both monocots (or at least

wheat) and dicots. While the exact mode of action of Si in influencing plant resistance is not fully understood, Fawe et al. (16) suggested that Si may mediate defense responses that are functionally similar to systemic acquired resistance. In Si+ plants, the amplitude of expression of defense responses appears greatly increased.

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