Infection of species of the Gramineae by *Erysiphe graminis* f. sp. *hordei* and *Erysiphe graminis* f. sp. *tritici*

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The ability of 19 isolates of *Erysiphe graminis f.* sp. *hordei (Egh)* and 34 isolates of *E. graminis* f. sp. *tritici (Egt)* to infect cultivated and wild species of Gramineae common to southern Ontario was examined. A high level of parasitic specialization of *Egh* was demonstrated by the failure of these isolates to infect any of the 22 species of Gramineae tested, except barley. In contrast, isolates of *Egt* were observed to infect wheat, meadow brome and downy brome. We suggest that these alternative hosts of *Egt* may act as oversummering reservoirs of inoculum for infection of fall sown winter wheat, as well as sources of some of the genetic variability found in the natural populations of the wheat pathogen in southern Ontario.

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On a examine la capacite de **19** souches de *Erysiphe graminis* f. sp. *hordei (Egh)* et de 34 souches de *E. graminis* f. sp. *tritici (Egt)* d'attaquer des especes cultivées et sauvages de graminées que l'on trouve couramment dans le sud de l'Ontario. On a de plus démontré un haut degre de spécialisation parasitique de *Egt*, car les souches de cette forme n'ont pu attaquer les 22 espèces de graminees eprouvees, à l'exception de l'orge. En revanche, les souches de *Egt* ont infecte le ble, le brome des pres et le brome des toits. Nous pensons que ces hôtes substituts de Egtserventa conserver pendant l'été des souches qui infecteront a l'automne les semis de ble d'hiver et qu'ils sont aussi à l'origine d'une certaine variabilite génétique de l'agent pathogene du ble observee dans les populations naturelles du sud de l'ontario.

Introduction

Surveys of the virulence spectra of Erysiphe graminis f. sp. hordei (Egh) and E. graminis f. sp. tritici (Egt) have indicated that natural populations of these pathogens in southern Ontario possess a wide range of genetic variability (Bailey and MacNeill, 1983; Menzies and MacNeil, 1986; Louter et al., 1987; Menzies et al., 1989). Possibly, the presence of known genes for resistance in commercial cultivars of barley (Martens et al., 1984) is a source of some of the genetic variability of natural populations of Egh through the selection of certain genes for virulence. However, not all the genetic variability observed in natural populations of Egh in southern Ontario can be explained in this way. Commercial cultivars of wheat used in southern Ontario do not possess any of the known Pmgenes for resistance to Egt (Martens et al., 1984) that might account for the spectrum of virulence in the natural population of the latter pathogen.

The possibility that genetic variability in a pathogen population may be due to selection pressure exerted by alternative hosts rather than the "preferred host" should be examined. Eshed and Wahl (1970) in Israel have demonstrated that both *Egh* and *Egt* enjoy a host range within the family Gramineae that is wider than the *forma* specialis designations would suggest. Fall infection of wild grasses in Israel by ascospores of *Egt* liberated from cleistothecia which have ovesummered on wheat stubble have been observed by Eshed and Wahl (1975). Colonies of Egton wild grasses act as foci from which conidia are disseminated to cultivated wheat crops throughout

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Table 1.	Species of Gramineae tested for compatibility
	with Erysiphe graminis f.sp. tritici and Erysiphe
	graminis f. sp. hordei.

Scientific Name	Common Name	
Agrostis palustris Huds.	Creeping Bentgrass	
Alopecurus pratensis L.	Meadow Foxtail	
Avena fatua L.	Wild Oats	
Avena sativa L.	Oats	
Bromus biebersteinii Roem. Schult.	Meadow Bromegrass	
Bromus inermus Leyss.	Smooth Bromegrass	
Bromustectorum L.	Downy Brome (Chess)	
Dactylis glomerata L.	Orchard Grass	
Digitaria ischaemum (Schreb.) Muhl.	Smooth Crabgrass	
Digitaria sanguinalis (L.) Scop.	Hairy Crabgrass	
Echinochloa crusgalli (L.) Beauv.	Barnyard Grass	
Festucarubra L.	Creeping Red Fescue	
Hordeum vulgare L.	Barley	
Lolium perenne L.	Perennial Ryegrass	
Panicum capillare L.	Witch Grass	
Panicum miliaceum L.	Proso Millet	
Phalarisarundinacea L.	Reed Canary Grass	
Phleum pratense L.	Timothy	
Poa pratensis L.	Kentucky Bluegrass	
Setaria glauca (L.) Beauv.	Yellow Foxtail	
Setaria viridis (L.) Beauv.	Green Foxtail	
Triticumaestivum L.	Wheat	

	Egh ⁺		Egt [†]	
Gramineae Species	Number of inoculated isolates	Number of compatible isolates	Number of inoculated isolates	Number of compatible isolates
Wheat (cv Augusta)	13	0	32	32
Barley (cv Bonanza)	19	19	22	0
Creeping Bentgrass	9	0	18	0
Coated Meadow Foxtail	9	0	17	0
Wild Oats	9	0	19	0
Domesticated Oats	10	0	18	0
Meadow Brome	9	0	27	10
Smooth Brome	9	0	23	0
Downy Brome	5	0	18	4
Orchard Grass	9	0	17	0
Smooth Crabgrass	15	0	34	0
Hairy Crabgrass	9	0	18	0
Barnyard Grass	9	0	15	0
Creeping Red Fescue	9	0	17	0
Perennial Ryegrass	9	0	21	0
Witch Grass	10	0	16	0
Proso Millet	5	0	18	0
Canary Grass	9	0	17	0
Timothy	9	0	18	0
Kentucky Bluegrass	10	0	22	0
Yellow Foxtail	9	0	14	0
Green Foxtail	7	0	10	0

Table 2.	The compatibility of isolates of Erysiphe graminis f. sp. hordei and Erysiphe graminis f. sp. tritici inoculated
	to different species of Gramineae commonly found in southern Ontario.

'Erysiphe graminis f.sp. hordei

[†]Erysiphe graminis f. sp. tritici

the growing season. In Israel, wild grasses infected by *fgh* and *fgt* bridge the gap between the different growing seasons for wheat, and may also be selecting for genetic variability in the two *formae speciales* involved.

In the present study the ability of *Egh* and *fgt to* infect cultivated and wild species of gramineae common to southern Ontario was examined.

Materials and Methods

Twenty two species of the Gramineae (Table 1) were tested for their receptivity to various isolates of **Egh** and **fgt**. The plants represented cultivated and weedy grass species common to southern Ontario. Five to 10 seeds of each grass were sown separately in 10 cm plastic pots and grown for 14 days in a growth room at $20 \pm 1^{\circ}$ C and 14-h photoperiod.

Nineteen isolates of fgh were used in these experiments and were derived from single colonies collected from various regions of southern Ontario in 1986 (Louter *et al.*, 1987) and 1987. Thirty-four isolates of fgt were used and derived from monoconidial and single colony isolates collected from various regions of southern Ontario during 1981 to 1987 (Bailey and MacNeill, 1983; Menzies and MacNeill, 1986; Menzies et al., 1989). Inoculum of the isolates of both Egh and Egt was obtained by the inoculation of the individual isolates of fgh onto a 10-day old plant of barley (cv Bonanza) and of fgt onto a 10-day old plant of wheat (cv Augusta); 8 days later approximately 5 mg of conidia of either an isolate of Egh or an isolate of fgtwas used to inoculate the different species of Gramineae. The inoculum was applied in a settling tower (Eval et al., 1968), the plants were capped with glass chimneys (Bailey and MacNeill, 1983), then incubated in a growth room for 8 days at $20 \pm 1^{\circ}$ C and a 14-h photoperiod. After 8 days, the plants were assessed for the presence or absence of colonies of E. graminis. Included in each set of inoculations of the different species of Gramineae was the barley cultivar Bonanza for fgh and the wheat cultivar Augusta for *Eqt*. These plants acted as controls to ensure that conditions were favourable for infection by the powdery mildew isolates.

Results and Discussion

A high level of parasitic specialization of **Egh** in southern Ontario was demonstrated by the failure of isolates of **fgh** to infect any of the species of Gramineae tested except barley

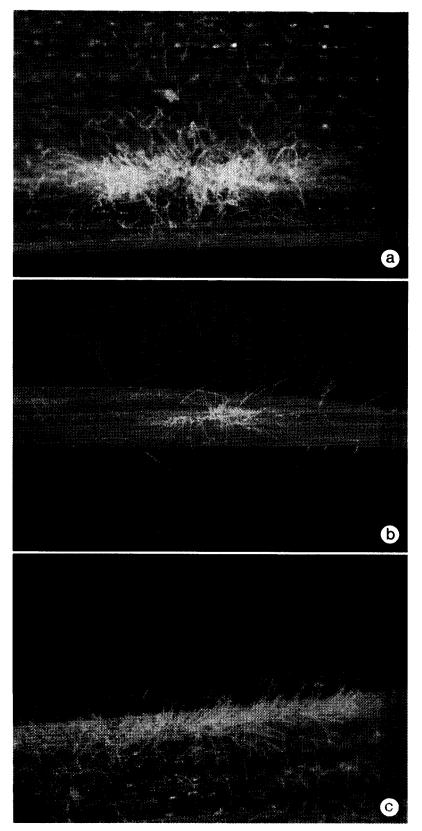


Figure 1. Colonies of *Erysiphe graminis* f. sp. *tritici* on; a) winter wheat; b) meadow brome; c) downy brome.

(Table 2). In Israel, Eshed and Wahl (1970) observed Eghon a wide spectrum of native grasses, and artificially infected 18 of 60 genera of Gramineae. Unlike our experiments, they observed compatible relationships between their isolates of Egh and downy brome. The differences between the results of Eshed and Wahl (1970) and ours may be due to different selection pressures within the natural population of Egh in the two different regions. The Mediterranean region is known to be the centre of origin and diversification of some of the progenitors of barley and wheat (Eshed and Wahl 1970). Greater diversity of host genotypes in the Mediterranean region as compared to southern Ontario may result in selection pressure leading to a broader spectrum of genotypes in the natural population of Erysiphe graminis in the former region.

Erysiphegraminisf. sp. *tritici* was not observed to be as host specific as Egh in southern Ontario; isolates of Egt infected wheat, meadow brome and downy brome (Table 2, Figure 1). Infection of downy brome by Egt has also been reported in Israel (Eshed and Wahl, 1970), but meadow brome was not tested in their trials. In the present study, cultures of Erysiphe graminis observed on plants of meadow brome and downy brome were confirmed as *Egt* by re-inoculation to wheat (cv Augusta).

The presence of alternative hosts for *Egt* in southern Ontario presents the possibility of another source of selection pressure leading to genetic variability. Pressure exerted by downy brome and meadow brome may explain the common occurrence of certain genes for virulence (pMa, p3c and p4) in southern Ontario (Bailey and MacNeill, 1983; Menzies and MacNeill, 1986; Menzies et *al.*, 1989). Certainly the role of such alternative hosts in contributing to greater variability of *Egt* needs to be examined.

Additionally, the fact the Egt infects downy brome and meadow brome in southern Ontario may also allow the pathogen to use these hosts as oversummering reservoirs of inoculum. The cleistothecia of *Egt* have been postulated to be the oversummering state of the pathogen in Canada (Cherewick, 1944). Menzies (1986) in 1984 and 1985 observed the maturation of cleistothecia of Egt in late July with ascospores being ejected in August, September and to a lesser extent in October. Ascospore ejection occurred, however, during periods when winter wheat was not being grown in southern Ontario. That is, between the harvest of the winter crop and the autumn planting of the next crop. Eshed and Wahl (1975) have suggested that in Israel, ascospore inoculum is important in infection of wild grasses, leading to the formation of colonies of powdery mildew and

production of conidia on these grasses. The colonies on the wild grasses act as foci from which conidia $\mathbb{1}$ *Erysiphe* graminis are disseminated to cultivated small grain crops throughout the growing season. In southern Ontario, colonies of *Egt* on meadow brome and downy brome, whether they are produced by ascospore or asexual conidial inoculum, may be important in the survival of *Egt* from the time of harvest of winter wheat in July, to the emergence of newly seeded winter wheat in October.

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