## Stimulatory Effect of Pollen and Pistillate Parts of Some Horticultural Species upon the Germination of Botrytis cinerea Spores

## H. Borecka and D. F. Millikan

Respectively, Principal Mycologist, Department of Food Storage, Research Institute of Pomology, Skierniewice, Poland, and Professor of Plant Pathology, University of Missouri, Columbia 65201.

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## ABSTRACT

Aqueous suspensions of pollen grains of Fragaria, Ribes, Rubus, Prunus, Malus, Hyacinthus, Iris, Narcissus, and Tulipa contain substances which stimulate the germination of Botrytis cinerea spores. Rubus pollen in the germinating media caused 75% of the spores to germinate, whereas that of other species was greater than 90%. This contrasted with the 5 to 16% found in the water controls. When sections of the pistil were tested, all parts from Tulipa markedly stimulated germination but the stimulatory effects from stigma sections of Prunus sp. were either much less, or absent.

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Botrytis cinerea Pers. is responsible for serious blossom blights and harvest rots of many horticultural plants. It is particularly damaging to strawberries in Poland, especially if wet and warm conditions prevail at blossoming time. Previously, Chou & Preece (3) showed that pollen grains from strawberry stimulated spore germination. Borecka et

al. (1) extended these studies to include other floral parts and found that the pistillate parts also were stimulatory while the petals and sepals were without effect. Borecka & Pieniazek (2) found that either synthetic abscisic acid (AbA) or AbA extracted from strawberry fruit would stimulate germination. In the investigations reported here, we studied the stimulatory effect of pollen from additional species and limited studies were made with the pistillate parts of some species.

We assayed the pollen from three small fruit species, Fragaria ananassa Hort. (strawberry), Ribes nigrum L. (black currant), and Rubus idaeus L. (red raspberry) for effects on germination of B. cinerea spores. We also assayed pollen from the tree fruit species, Prunus armeniaca L. (apricot), P. avium L. (sweet cherry), P. cerasus L. (sour cherry), P. domestica L. (plum), P. persica L. (peach) and Malus sylvestris Mill. (apple). Additionally, pollen from the ornamental species, Hyacinthus orientalis L. (hyacinth), Iris germinca L. (Iris), and Narcissus pseudonarcissus L. (daffodil) was evaluated. Our experimental technique followed that of Chou & Preece (3) wherein 30,000 pollen grains/ml of each species were mixed with 375,000 spores/ml of B. cinerea. Distilled water drops containing only B. cinerea spores served as controls in each experiment, and the percentage of germination was determined after 8 hr incubation at room temperature.

The data (Table 1) show that pollen from all tested species stimulated the germination of *B. cinerea* spores. Except for currant pollen where the germination was 75%, germination was increased to over 90%, contrasting to the 5.8 to 16.5% germination found in the water controls. Other studies were made to determine whether germ tube

TABLE 1. Stimulatory effect of pollen upon the germination of Botrytis cinerea sporesy

Pollen source <sup>Z</sup>						
Small fruits		Tree fruits		Ornamentals		
Species	Germination of spores (%)	Species	Germination of spores (%)	Species	Germination of spores (%)	
Fragaria	Prunus			Hyacinthus		
ananassa Hort.	91.5 a	armeniaca L.	97.5 a	orientalis L.	93.5 a	
Ribes		P. avium L.	96.5 a	Iris		
nigrum L.	76.0 b			germanica L.	90.7 a	
		P. cerasus L.	95.3 a	Control of the Contro		
Rubus				Narcissus		
idaeus L.	91.2 a	P. domestica L.	92.3 a	pseudonarcissus L.	97.5 a	
Water	5.8 c	P. persica L.	96.3 a	Tulipa		
				gesneriana L.	97.3 a	
	1	Malus				
		sylvestris Mill.	92.3 a	Water	16.5 b	
	,	Water	9.2 b			

y Each mean represents an average of three separate experiments and each mean within a vertical column followed by the same letter(s) are not significantly different at the 5% probability level.

<sup>&</sup>lt;sup>2</sup> Botrytis cinerea spores added to water at concentration of 30,000/ml. Pollen grains added at concentration of 760,000/ml, and % germination determined after 8 hr.

TABLE 2. Stimulatory effects of pistillate floral parts upon the germination of *Botrytis cinerea* spores

Species	Pistillate <sup>y</sup> part	Germination <sup>2</sup> (%)
Tulipa		
gesneriana L.	stigma	79.3 a
	style	92.5 b
	ovary	83.3 b
Prunus	U20040150	
avium L.	stigma	33.5 с
cerasus L.	stigma	24.8 d
domestica L.	stigma	19.8 e
Water	1944	17.8 e

y Each floral part was cut into 0.5-mm sections and one section added to a drop of water containing 30,000 spores/ml.

<sup>2</sup> Means followed by the same letter are not significantly different at 5% probability level.

growth might be affected. In solutions containing raspberry and strawberry pollen, germ tube length of *B. cinerea* was found to average 264.0 and 232.5  $\mu$ , respectively, contrasting with 44.6  $\mu$  in water only.

Table 2 shows the effects of pistillate parts of flowers upon the germination of *B. cinerea* spores and indicates that 0.5-mm sections of the stigma, style, and ovary from tulip cause 80-90% germination when added to drops of spore suspension. Only the stigmas from *Prunus* species were tested. Those from sweet and sour cherry were stimulatory, whereas those from plum were without effect. No effect, however,

was observed from uncut stigmas suggesting that endogenous substances released by injury are responsible and that these vary with the species.

Our data show that the pollen from many horticultural plants, when suspended in water, will stimulate the germination of B. cinerea spores and also may affect subsequent germ tube length. The presence of this stimulatory factor in aqueous suspensions of pollen very likely contributed to the heavy infection of gray mold in Polish strawberries in 1971 when excessive rainfall occurred during May and June. They also show that the pistillate parts, if wounded, could influence infection since cut sections of these flower parts of species stimulate spore germination. The identity of the stimulatory substances is unknown but those associated with the pollen are water-soluble and are not sugars (2). The stimulatory properties of AbA and its presence in pollen grains sufficient for affecting germination (2) indicate that this auxin plays a role.

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