Environmental Factors Affecting Conidial Survival of Five Pear Decay Fungi

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ABSTRACT

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The survival of conidia of Botrytis cinerea, Mucor piriformis, Penicillium expansum, Pezicula malicorticis, and Phialophora malorum on pear fruit surfaces in the orchard and under selected temperature-relative humidity combinations was determined. Under controlled conditions, survival half-life (T_{50}) of B. cinerea and Penicillium expansum often exceeded 1 mo, but the T_{50} s of M. piriformis and Pezicula malicorticis often were less than 2 wk. Survival of these fungi decreased as temperature increased from -1.1 to 30 C. The T_{50} of Phialophora malorum varied from about 2 to 6 wk and was less at 20 than at 10 or 30 C. Conidial survival in the orchard was about 20–40% of that at similar temperature and relative humidity in controlled environments, and the T_{50} s in the orchard of B. cinerea, Penicillium expansum, and M. piriformis were 14.4, 6.1, and 4.2 days, respectively.

Botrytis cinerea Pers. ex Fr., Mucor piriformis Fisher, Penicillium expansum Lk. ex Thom., and Phialophora malorum (Kidd & Beaum.) McCulloch cause decay of stored pears and have been found in soil, surface plant residues, and air (1). Pezicula malicorticis (Jacks.) Nannf., the causal agent of bullseye rot, survives on plant surfaces and in cankers (3). Fungal propagules may be deposited on fruit on the tree during harvest or in the packinghouse.

Little is known about factors affecting survival of pear decay fungi. B. cinerea conidia survived 2-6 mo at 0 C and 1-3 mo at 20 C and 99% RH (6). B. cinerea hyphae on glass slides remained viable for 12 mo when stored in the range between 85 and 100% RH at 0 C (6). The effect of environmental factors on survival of other pear decay fungi has not been studied.

The objective of this study was to determine the survival of conidia of B. cinerea, M. piriformis, Penicillium expansum, Pezicula malicorticis, and Phialophora malorum on pear fruit surfaces in the orchard and under selected

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temperature-relative humidity combinations.

MATERIALS AND METHODS

Survival of decay fungi in the orchard. All fungi were isolated from decayed pear fruits and grown on potato-dextrose agar (Difco, Detroit, MI) acidified with 1.5 ml of 85% lactic acid per liter (APDA). Cultures 1-2 wk old were flooded with sterile water and suspensions adjusted to obtain $5-6 \times 10^4$ conidia per milliliter. Conidial suspensions of B. cinerea, M. piriformis, Penicillium expansum, and Pezicula malicorticis were sprayed to runoff onto Anjou fruits and Phialophora malorum onto Bosc fruits at monthly intervals from 4 mo to 2 days before harvest. Side rot, caused by P. malorum, is a serious problem on Bosc but is of no importance on other pear cultivars. Control fruits were sprayed with sterile distilled water.

Fruits were removed 1 day, then weekly from 1 to 8 wk, after each monthly inoculation. Two sections of epidermal tissue, each about 2 cm², were removed from five fruits inoculated with each fungus. Tissue pieces were inverted and placed on APDA. Plates were incubated at 20 C, and fungus survival as evidenced by mycelial growth was evaluated after 1 wk. Linear regression lines developed from loge of percent survival vs. time were analyzed with covariance and comparison of independent estimates of slope.

Survival of decay fungi in controlled environments. Mature Anjou pear fruits were harvested and surface-sterilized with 0.525% sodium hypochlorite, rinsed, and air-dried. Conidia of the five fungi used in orchard inoculations were harvested from 3-wk-old APDA cultures, adjusted to 5×10^4 conidia per milliliter, and sprayed onto the fruits to runoff. Fruits were allowed to air-dry, then placed at -1.1 ± 1.5 , 10 ± 1.5 , 20 ± 1 , and 30 ± 1 C. Relative humidities in the

experiment included $95 \pm 5\%$ at -1.1 C; $90 \pm 10\%$ at 0, 20, and 30 C; and $40 \pm 10\%$ at 20 and 30 C. Relative humidity at -1.1 C was measured with electronic sensors (American Instrument Co., Silver Spring, MD), and at other temperatures, with calibrated hygrothermographs (Belfort Instrument Company, Baltimore, MD). Fruits at 10, 20, and 30 C were exposed to continuous light energy of 550 $\mu \hat{\mathbf{W}}/\text{cm}^2$ at 400 μ m. Fruits at -1.1 C were placed in a commercial, polyethylenelined fruit storage box in the dark. Epidermal tissue pieces 0.5×1 cm were removed weekly (biweekly from fruits held at -1.1 C) from four positions on each of four fruits inoculated with each fungus and placed with the cuticle down on APDA. Fungus survival was evaluated after 1 wk. Data were analyzed with linear regression, and survival half-life (T₅₀ [the time after which the inoculated fungus produced colonies from half of the plated epidermal pieces]) was calculated from regression equations.

RESULTS

Survival of decay fungi conidia in the orchard. B. cinerea conidial survival on the surface of attached Anjou pear fruits declined at a significantly (P = 0.01) slower rate than Penicillium expansum or M. piriformis (Fig. 1). The T_{50} s of B. cinerea, P. expansum, and M. piriformis were 14.4, 6.1, and 4.2 days, respectively, and were all significantly different (P = 0.01). Survival of Pezicula malicorticis and Phialophora malorum conidia could

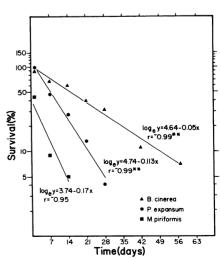


Fig. 1. Survival of decay fungi conidia in the orchard on the surfaces of attached Anjou pear fruits. Each point represents the mean of three to 20 fruits, two tissue sections per fruit. ** = Correlations significant at P = 0.01.

Table 1. Conidial survival half-lives (T_{50}), in days, of five decay fungi on the surface of Anjou pear at selected temperature-relative humidity combinations

Fungus	T_{50}^a at each temperature (C)/relative humidity (%)				
	10/90	20/40	20/90	30/40	30/90
Botrytis cinerea	35.8	46.2	20.5	30.8	12.4
Mucor piriformis	14.9	12.0	15.7	8.8	5.8
Penicillium expansum	133.3	36.5	>27.0 ^b	26.1	16.2
Pezicula malicorticis	7.8	12.2	11.1	8.8	6.8
Phialophora malorum	40.6	13.5	11.8	18.4	25.4

 $^{^{}a}$ T₅₀ values in days were calculated from linear regression equations of fungal survival at weekly intervals. Survival was determined by presence of fungal growth from Anjou epidermal tissue on APDA. Correlation coefficients of survival with time of all fungi are significant (P = 0.05) except M. piriformis at 30 C/90% RH and P. malorum at 10 C/90% RH.

not be determined because these slow-growing fungi, if viable, were masked with other microorganisms present on fruit surfaces in the orchard. B. cinerea and Penicillium expansum were isolated from less than 4% of the tissue sections from fruits sprayed with distilled water. No other decay fungi were isolated from control fruits.

Survival of decay fungi in controlled environments. The T₅₀ of each fungus at each selected temperature-relative humidity combination is presented in Table 1. The T₅₀s of *M. piriformis* and *Pezicula malicorticis* were similar and generally ranged from about 1 wk at 30 C to 2 wk at 10 and 20 C. The T₅₀s of *B. cinerea, Penicillium expansum,* and *Phialophora malorum* were much longer than those of *M. piriformis* and *Pezicula malicorticis* and ranged from 2 to 4 wk at 20 and 30 C and more than 4 wk at 10 C. The T₅₀ of *Penicillium expansum* at 10 C exceeded 4 mo

With few exceptions, the T_{50} s of B. cinerea, M. piriformis, Penicillium expansum, and Pezicula malicorticis declined as temperature increased. The T_{50} of Phialophora malorum, however, was greater at 30 than at 20 C. With the exception of M. piriformis at 20 C and P.

malorum at 30 C, the T_{508} of all fungi were less at 90 than at 40% RH within each temperature.

At -1.1 C, survival of *M. piriformis* and *P. malorum* remained at 100% at the end of the 30-wk experiment. *B. cinerea* and *Penicillium expansum* survival decreased only slightly after 30 wk, and the T₅₀ was calculated as 19 mo. The T₅₀ of *Pezicula malicorticis* at -1.1 C was 7.3 mo

DISCUSSION

Survival of conidia of decay fungi varied greatly among species, temperature, and relative humidity. The T₅₀s of B. cinerea and Penicillium expansum exceeded 1 mo under most conditions, but those of M. piriformis and Pezicula malicorticis were often less than 2 wk. B. cinerea conidia were reported to survive less than 1 mo at 20 C and 90% RH, but length of survival varied with strain (6). Currently, a fungicide application is recommended within 2 wk of harvest for pear decay control. This timing appears to coincide with the need 1) for protection of fruit as susceptibility to decay increases (5) and 2) to reduce decay spore populations that could survive from 2 wk to harvest and infect fruit during harvest and postharvest handling procedures.

The T₅₀ of conidia in the orchard was about 20-40% of that at similar temperature and relative humidity in controlled environments. Several factors that likely reduced orchard survival include fluctuations and extremes in temperature and relative humidity and ultraviolet (UV) light. However, orchard inoculations were repeated several times during the growing season, and decreases in T₅₀ did not appear consistently related to temperature or relative humidity. B. cinerea is extremely sensitive to UV light, and exposures longer than 2 hr decreased viability (4). Spores of a Botrytis sp. from apple were killed on an agar surface by a 1-min exposure to a UV light placed 15 cm from the culture (2).

Survival of conidia of all decay fungi exceeded 7 mo at -1.1 C in this study and may have important practical significance. In the Pacific Northwest, Anjou and Bosc pears commonly are stored at -1.1 C as long as 8 mo. Fungi that survive through the storage period could cause additional fruit decay during packaging, transport, and ripening at terminal markets.

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^bExact calculation of T₅₀ not possible because of incomplete data.