## Yonghao Li

Department of Entomology and Plant Pathology, University of Tennessee, Knoxville

## Margaret T. Mmbaga

Otis L. Floyd Nursery Crop Research Center, Tennessee State University, McMinnville

#### Alan S. Windham

Soil, Plant and Pest Center, University of Tennessee, Nashville

## Mark T. Windham and Robert N. Trigiano

Department of Entomology and Plant Pathology, University of Tennessee, Knoxville

# Powdery Mildew of Dogwoods: Current Status and Future Prospects

Cornus is a large genus of trees and shrubs that are collectively referred to as dogwoods. Flowering dogwood (C. florida L.) and kousa dogwood (C. kousa (F. Buerger ex Miq.) Hance) and interspecific hybrids of these species are popular ornamental trees that are known for their showy bracts, red berries (drupes), and/or fall color. Other species that are commercially grown for specialty markets include the pagoda dogwood (C. alternifolia L.), giant dogwood (C. controversa Hemsl.), cornelian cherry (C. mas L.), Pacific dogwood (C. nuttallii Aud.), and redosier dogwood (C. sericea L.). The foliage of native species, such as flowering dogwood and pagoda dogwood, is high in calcium (12) in quantities above what is needed for skeletal growth of wildlife; it is the preferred browse material for lactating does in late spring while many other trees are still leafless (13,22). The berries of flowering dogwood have high oil content and provide mast for numerous species of migrant songbirds, wild turkeys, and large and small mammals (22).

For many years, nurseries that produced flowering and kousa dogwoods had the luxury of working with relatively disease-free crops. Disease management and control costs were minimal and estimated at approximately \$120/ha/year. In the late 1970s, flowering and kousa dogwoods were threatened by a new disease, dogwood anthracnose, caused by *Discula destructiva* (39), which was reviewed by Daughtrey et al. (3). In 1994, another disease, powdery mildew, reached epiphytotic levels in flowering dogwoods. Tens of millions of dollar's worth of dogwoods

Corresponding author: Yonghao Li, Department of Entomology and Plant Pathology, University of Tennessee, Knoxville; E-mail: yli20@utk.edu

doi:10.1094/PDIS-93-11-1084
© 2009 The American Phytopathological Society

were destroyed and millions of cultivated seedlings lost their commercial value because formal management strategies were not formulated. In subsequent years, fungicide management costs were estimated to be \$1,975/ha/year. Many small producers of dogwoods terminated production of the tree because they could not afford the additional overhead or were not inclined to continue routine fungicide sprays every 2 weeks from May to October.

Powdery mildew on C. florida was first reported in 1887 by Burrill and Earle (1), but this disease was rarely reported on flowering dogwood in the United States before 1994. However, the disease appeared simultaneously in forest, landscape, and nursery plantings statewide in Alabama in 1994 (8). Similar outbreaks of powdery mildew were observed in Tennessee, where many nursery fields of flowering dogwood were abandoned (Fig. 1). Powdery mildew has emerged as a nationwide disease of flowering dogwood (2). Although the host side of the disease triangle remained constant, we do not know whether the change of frequency and severity of powdery mildew in flowering dogwood was due to a change in the pathogen or a change in the environment.

#### **Pathogen**

Two powdery mildew species have been reported to infect dogwoods. Erysiphe pulchra (Cooke & Peck) U. Braun & S. Takam. (syn. Microsphaera pulchra Cook & Peck) is considered to be the more prevalent (6,16,21,23), while Phyllactinia guttata (Wallr.:Fr.) Lev. is occasionally found on dogwood leaves (2,5,16,23). Klein et al. (16) found ascocarps of E. pulchra and P. guttata on C. florida and C. amomum, silky dogwood, but concluded that ascocarps of P. guttata did not develop on C. florida, whereas ascocarps of E. pulchra did. Windham et al. (46) found that E. pulchra infected and produced ascocarps on C. florida, C. kousa, and C.

nuttallii, whereas P. guttata infected and produced ascocarps on C. alba, C. amomum, C. drummondii, C. macrophylla, C. obliqua, C. racemosa, C. sericea, and C. stricta.

In the sexual stage, both fungi belong to division Ascomycota but have distinctive appendages; E. pulchra has dichotomously branched and tapered appendages (Fig. 2A), whereas the bulbous base of appendages distinguishes P. guttata (Fig. 2B). Immature chasmothecia (syn. cleistothecia) of E. pulchra are yellow- to ambercolored, then turn dark brown to black when mature; the size ranges from 75 to 128 µm in diameter (Fig. 2C) (16,41,44). The length of appendages ranges from 110 to 160 µm (16). There are three to five asci in a chasmothecium and four to eight ascospores in an ascus of *E. pulchra* (Fig. 2D). Ascospores are single-celled, globose, and measure  $18-28 \mu m \times 13-15 \mu m$ .

The asexual stage of *E. pulchra* (*Oidium* sp.) forms conidia that serve as inoculum and cause disease epidemics within a growing season. Conidia are single-celled, ovoid to hyaline, and borne on conidiophores singly or in pseudochains (Fig. 2E and F). Conidia (28.1 × 14.1 µm) are highly vacuolated (41). Analysis of DNA sequence of the internal transcribed spacer (ITS) region has shown distinct sequence differences between *E. pulchra* and *P.* 



Fig. 1. An abandoned nursery field of powdery mildew-infected dogwoods in Tennessee.

guttata, whereas there were no sequence variations among E. pulchra isolates (25). DNA sequences of the ITS region from Tennessee and New York isolates of E. pulchra from C. florida (GenBank accession no. AY224136) were 100% identical to each other and to that of Japanese E. pulchra isolates from C. kousa (GenBank accession no. AB015935). Identical ITS sequences from all E. pulchra isolates were obtained. Genetic uniformity in pathogen populations may be a reflection of the recent introduction of the pathogen (37), but studies on E. pulchra population analysis are needed.

#### **Infection Process**

Conidia of E. pulchra germinate and initiate one to four primary germ tubes by 2 h after inoculation (Fig. 3A). However, primary appressoria only differentiate from one or two of the primary germ tubes (Fig. 3B) (17). Penetration pegs form under appressoria and breach the wax and cuticle layers of epidermal cells. Haustoria, globose fungal feeding structures surrounded by extrahaustorial matrix, differentiate in epidermal cells 2 days after inoculation and absorb water and nutrients supporting fungal growth (Fig. 3C). Secondary germ tubes, or primary hyphae, initiate from only one of the primary germ tubes with primary appressoria (Fig. 3D). The primary and secondary appressoria are lobed and form singly or in pairs opposite one another (Fig. 3E). Branched hyphae are differentiated between primary and secondary appressoria or develop directly from secondary appressoria (Fig. 3F). After establishing the host-parasite relationship, fungal hyphae elongate and form colonies (Fig. 3G). Conidia are borne on conidiophores that display a twisted basal cell (Fig. 3H).

## **Symptoms and Signs**

Disease signs first appear on the adaxial leaf surface as circular to irregular white patches that consist of mycelia and conidia of the fungus (Fig. 4A). As the fungus colonizes more host tissues, the leaves are covered by white mildew and develop mottled yellowing or brownish patches. Newly infected leaves curl upward and result in a tree canopy with distorted growth, which is aesthetically unacceptable to growers and consumers alike (Fig. 4B). By mid-summer, reddish-brown blotches appear on infected leaves, and symptoms may mimic those of drought stress, even as colony expansion on the discolored area slows (Fig. 4C). Near the end of the growing season, light brown-toblack chasmothecia may be observed in mildew colonies on either leaf surface, but tend to be produced predominantly on the lower surface (Fig. 4D).

#### **Host Resistance**

Susceptibility to powdery mildew varies among Cornus species. In general, most C.

florida cultivars are susceptible and most C. kousa cultivars and hybrid dogwood (C. kousa × C. florida) cultivars are resistant (7,20), whereas C. mas, C. controversa, and C. alternifolia are immune to powdery mildew (46). Hybrids from crossings of C. kousa and C. florida 'Stellar Pink', 'Stardust', 'Galaxy', 'Constellation', and 'Aurora' have also been reported to be highly resistant to powdery mildew, whereas hybrid 'Ruth Ellen' was moderately resistant at some locations and highly resistant at others (7,15,20,27,46). In C. florida, 'Cherokee Brave', a pink-bracted flowering dogwood, has also been reported as resistant to the disease (7,50), but resistance has failed in some years since these reports. 'Jean's Appalachian Snow', 'Kay's Appalachian Mist', 'Karen's Appalachian Blush', and 'Appalachian Joy', all whitebracted flowering dogwoods, are highly resistant to powdery mildew (45,51).

Powdery mildew resistance in dogwoods is often expressed as restricted branching

of hyphae without affecting germination of conidia (19). Genes controlling resistance and resistance mechanisms are not yet clear. Some dogwood seedlings express partial resistance to powdery mildew with slower disease progress than susceptible cultivars (19,51). Since E. pulchra is an obligate biotroph, sporulation, colony development, and disease spread depend on successful initial penetration of the host and continuous functional haustorium formation. Compared to susceptible cultivars, resistant cultivars delay disease latent period, reduce pathogen infection efficiency, and restrict colony development and asexual reproduction by conidia, all indications of partial resistance (19,20) (Fig. 5). Resistance to powdery mildew in dogwood has been evaluated in nurseries or greenhouses by rating disease severity (7,15,27,38,46,51). However, variation in levels of resistance to powdery mildew in locations and years has been reported for some dogwood cultivars (7,15,27,38,51). A

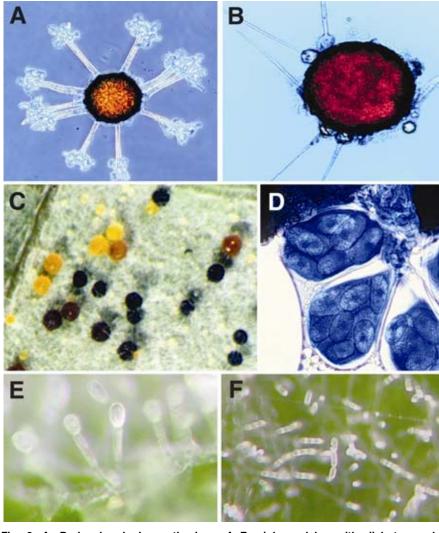


Fig. 2. A, Dark-colored chasmothecium of Erysiphe pulchra with dichotomously branched appendages. B, Chasmothecium of Phyllactinia guttata with bulbous bases of appendages. C, Chasmothecia of E. pulchra on leaf surface. D, Ascospores inside an ascus released from a chasmothecium of E. pulchra. E, Singly formed conidia on conidiophores of E. pulchra. F, Conidia formed in pseudochain. (A and B adapted from Mmbaga [24])

leaf disk bioassay method has been developed and used to evaluate dogwoods for resistance to powdery mildew in the laboratory. Results were similar to field observations on these same cultivars (19,20).

## **Epidemiology**

Understanding the epidemiology of powdery mildew on dogwood enables us to

devise rational disease management strategies that take into account the pathogen's life strategies (52). In middle Tennessee, chasmothecia of *E. pulchra* are the most important overwinter fungal structures, which form abundantly on dogwood leaves but not on stems, even on severely affected plants (16,23). In the northeastern United States, Smith reported that overwintering

chasmothecia containing mature asci and ascospores were found on dogwood twigs and on fallen leaves in March (41). The chasmothecia survival was influenced by the timing of chasmothecia formation over the period from September to November (23,24). Ascocarps that form late in the season may not be mature enough to overwinter because they are less developed.

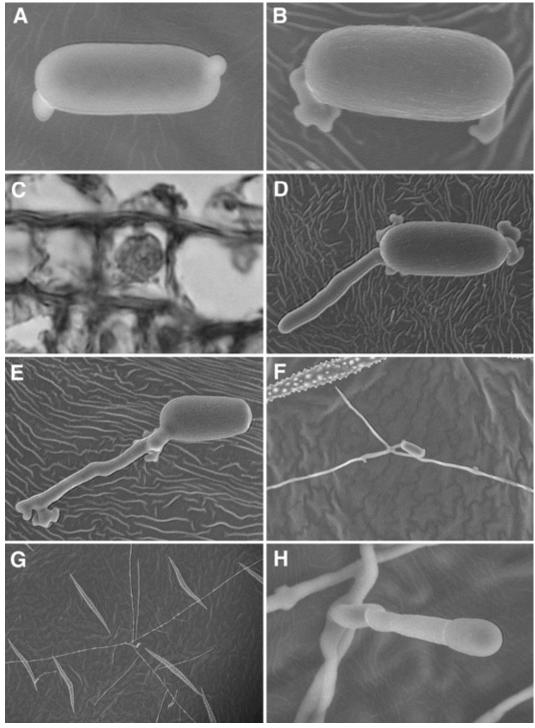


Fig. 3. Micrographs of infection process of *Erysiphe pulchra*. A, Primary germ tubes on both poles of a conidium. B, Primary appressoria from primary germ tubes. C, Globose haustorium in a host epidermal cell and the haustorial neck connecting haustorium body and appressorium on the surface of epidermal cell wall. D, Hyphal growth from the primary appressorium. E, Germinated conidium with secondary appressorium. F, Branched hypha from hypha and from secondary appressorium. G, Growth of branched hypha from hypha and secondary appressorium. H, Close-up of conidium and conidiophore with arched basal cell. (adapted from Li et al. [17]).

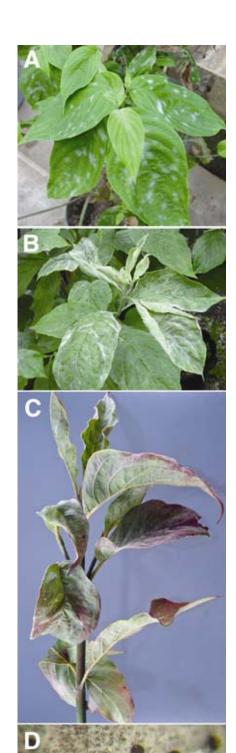


Fig. 4. Symptoms and signs of powdery mildew in dogwood. A, Isolated mildew colonies at beginning of colony development. B, Merged mildew colonies and curled young leaves. C, Red-brown patches under mildew colonies in mid-summer. D, Chasmothecia formed on mildew colonies in late fall.

Chasmothecia on leaf debris may be yellow, brown, or black, indicating the maturity levels from immature to mature (Fig. 2C). Chasmothecia maturation requires several weeks (23,24,53); thus, variation in the initiation of chasmothecia formation from year to year may cause variability in winter survival and primary inoculum density. Studies under controlled environment showed that temperature affected the formation of chasmothecia. Cooler temperatures of 18/10°C and 23/15°C (day/night) were favorable to chasmothecia initiation and development. Day length and the physiological stage of affected plants had no effect on chasmothecia formation. Variation in autumn temperatures was associated with the timing of chasmothecia formation (23,24). Where a pathogen overwinters on dormant plants, chasmothecia formed on leaf debris may be of secondary importance in the disease cycle, but in mid-Tennessee, the powdery mildew pathogen did not survive from one season to the next as mycelia in dormant dogwood buds (23).

Moderate to high numbers of chasmothecia on leaves survive during winter months at various locations outdoors, on the ground or hanging on tree branches, and release viable ascospores the following spring. Airborne ascospores were trapped on sticky slides between March and June, and dogwood seedlings used as trap plants developed powdery mildew from airborne inoculum (24). Disease severity corresponded with increasing spore counts on sticky slides and confirmed that E. pulchra overwintered on leaf debris primarily as chasmothecia, and ascospores served as primary inoculum (23). Infection on newly expanding dogwood leaves became visible when masses of conidia were formed (23,24). This indicated that primary infections had developed secondary inoculum and that primary infection from ascospores occurred earlier than indicated by disease symptoms. The timing of the initial infection in commercial nurseries varied from year to year by 2 to 6 weeks (early May to late June). High disease severity in 1996, 1998, and 2000 were associated with high chasmothecia frequency in 1996, low frequency in 1998, and very low frequency in 2000 with >50 chasmothecia per 19.6 mm<sup>2</sup> leaf area (high), 25 to 49 (moderate), and >25 (low) frequency (23). These observations suggested that factors other than disease severity influenced the abundance of chasmothecia formed as the source of primary inoculum for the following year (23,24). Ascospores were detected over a period of several weeks (March to June) with peaks in early to late April depending on rainfall and temperature.

Thus, two spore stages of E. pulchra occur simultaneously during early spring, with ascospores and conidiophores infecting newly expanding leaves simultaneously, causing rapidly growing polycyclic epidemic. When environmental conditions are favorable, powdery mildew spreads very rapidly, with masses of conidia produced from each new infection within a few days. Epidemiological studies have shown that powdery mildew generally begins during late May to June. This initial disease incidence is followed by a rapid increase in disease severity until early to mid-August (18,24). Disease progress curves of powdery mildew on dogwood were fitted to the logistic model (18). This showed that, overall, temperature and rainfall patterns were likely the main environmental factors that influenced primary inoculum density. Variation in the timing of infection establishment between early May and late June was associated with inoculum density, but it did not affect overall disease severity for the season. The association between prevailing weather conditions and disease severity over a 5year period has shown that well-distributed (frequent) rainfall events and moderate monthly temperatures favor high incidence of powdery mildew in dogwoods (24).

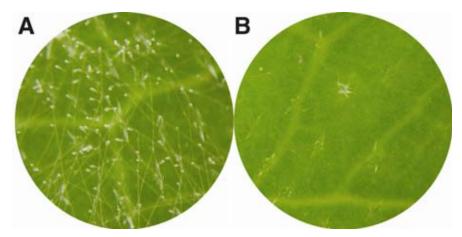


Fig. 5. Resistant and susceptible flowering dogwood cultivars' reaction to powdery mildew. A, Sporulating colony development on susceptible cultivar. B, Restricted colony on resistant cultivar. (adapted from Li et al. [19])

## **Disease Management**

Host resistance. Several mildew-resistant dogwood species, hybrids, and flowering dogwood cultivars are recommended as a key disease management strategy. Resistant dogwood species include the following: C. kousa, C. sericea, C. mas, C. alternifolia, C. alba, and C. controversa. Hybrids of C. kousa  $\times$  florida, including 'Stellar Pink', 'Stardust', 'Galaxy', 'Constellation', and 'Aurora', are also highly resistant to powdery mildew (7,15,20, 27,46). Of all ornamental dogwoods, C. florida is the most popular, but powdery mildew resistance is very limited in this species. Selection and release of resistant cultivars are highly valuable to the nursery and landscape industry. New cultivars 'Jean's Appalachian Snow', 'Kay's Appalachian Mist', 'Karen's Appalachian Blush', and 'Appalachian Joy', all whitebracted flowering dogwood, are highly resistant to powdery mildew (Fig. 6). However, breeding for resistant cultivars is nearly impossible because of self-incompatibility and long generation times.

Fungicides. In the mid-1990s, when powdery mildew suddenly became widespread on flowering dogwood in the eastern United States, no one in the green industry had experience in managing the disease. Powdery mildew had been managed on many other nursery crops, such as Syringa and Euonymus spp., but not on dogwoods. As dogwood powdery mildew became a recurring problem on dogwood each year in the eastern United States, fungicide efficacy against E. pulchra was identified. Most fungicides that are labeled for powdery mildew on other ornamental plants have proven to be efficacious against E. pulchra (4,33,36,47). Chlorothalonil, benzimidazole fungicides such as thiophanate methyl, and demethylation inhibitors such as fenarimol, myclobutanil, triadimefon, and propiconazole are efficacious at 2-week intervals. QoI fungicides

(strobilurins) also have shown promise as a management tool (9,10,32,35,47–49).

Currently, management of powdery mildew on susceptible dogwood seedlings in nurseries is primarily by foliar fungicide sprays, which has increased production cost. Optimum powdery mildew control in Alabama was achieved with spray programs that commenced on 1 June at the first sign of disease and terminated either 1 August or 1 September (11). Acceptable control could be achieved if spray programs were begun when small colonies of powdery mildew were visible on foliage (11,49). Obviously, the goal of nurseries is to keep powdery mildew at acceptable levels with as few sprays as possible. Many spray programs for foliar diseases of ornamental plants that are problematic annually begin prior to symptom development. Initiating fungicide sprays at the first sign of disease is possible only if nurseries are actively scouting for the advent of powdery mildew.

Nurseries that specialize in dogwood production are concerned about any disease that affects quality or grade and slows growth. Fungicide sprays can prevent the most objectionable signs and symptoms of powdery mildew such as white fungal growth, twisted leaves, leaf curl, and stunted growth (4). Leaf scorch is another indirect effect of powdery mildew that can be managed with fungicide sprays. Leaves infected with powdery mildew lose water faster than healthy leaves. In unirrigated fields, trees that are not protected with fungicide sprays are more likely to exhibit marginal necrosis associated with leaf scorch (47). Fungicide sprays not only produce healthier, higher quality trees, but may also increase tree height and caliper. It has been found that dogwoods protected from powdery mildew by fungicide sprays have increased tree height and trunk caliper compared to untreated trees (9,47,48). Delays in reaching desired standard heights or trunk calipers mean tangible losses to nurseries. It is not unusual for flowering dogwood to produce two flushes of growth during the growing season: one in early spring and a second in mid-summer. Dogwoods infected with powdery mildew seldom produce the second flush of growth at mid-summer, which accounts for decreased height.

Cultural controls used to manage other foliar diseases of ornamental plants such as plant spacing to aid in air movement and using drip irrigation to keep foliage dry are insufficient to control powdery mildew on dogwood. Fungicide sprays are likely to remain a viable tool for nurseries, landscape managers, and gardeners to protect dogwood from powdery mildew and to maximize growth. Future studies will look at new fungicides and fine tuning spray schedules to decrease production costs.

Biorationals. Disease control compounds that are less harmful to the environment and nontarget organisms than conventional fungicides have been designated as biorational fungicides (43). Such compounds are also referred to as biopesticides. Different modes of action have been reported such as preventing spore germination, retarding sporulation and mycelial growth, and inducing systemic resistance (14.34.40). Biorationals are most effective when used preventively at short spray intervals; they may be used in fungicide rotations, thereby reducing conventional fungicide use and the development of fungicide resistance (42). Biorational fungicides have several advantages over conventional pesticides: lower toxicity to mammals, pest species-specificity, rapid decomposition, and efficacy in small quantities. On the other hand, biorational fungicides require short treatment reapplication interval (7-day) compared to 14-day or longer intervals used in conventional fungicides (28). The contact mode of action

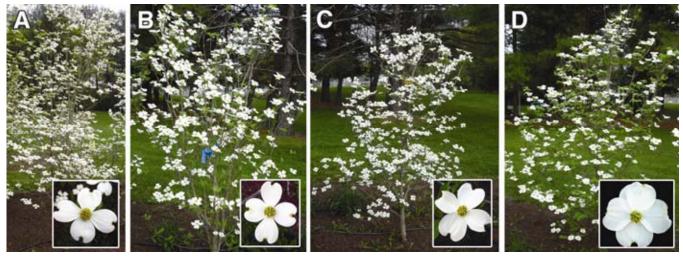


Fig. 6. Four flowering dogwood cultivars that are resistant to powdery mildew: A, Kay's Appalachian Mist; B, Karen's Appalachian Blush; C, Jean's Appalachian Snow; and D, Appalachian Joy.

may also contribute to lower efficacy in biorational fungicide treatments.

Twenty-one biorational compounds have been evaluated to identify alternative products and reduce conventional fungicides used in dogwood production (30). Household soaps Palmolive and Ajax that contain the antimicrobial compound triclosan are highly effective in reducing disease severity, similar to conventional fungicides, chlorothalonil and thiophanate methyl (Consyst), propiconazole (BannerMaxx), and thiophanate methyl (Cleary's 3336) (Fig. 7). Soap and potassium salts of fatty acids marketed as commercial insecticidal



Fig. 7. Cornus florida 'Cherokee Princess' grown under field conditions and treated with biorational products to control powdery mildew. Treated with: A, potassium bicarbonate; B, Equate, antibacterial liquid hand soap; C, Palmolive; D, water control; E, Ajax, antibacterial dish soap; F, nontreated. (adapted from Mmbaga and Sauve [28])

soap M-Pede and Safer Soap, hydrophobic extract of neem seed oil marketed as Triact 70 and Neem Gold, bicarbonate salt marketed as Armicarb 100 and Kaligreen, and the refined light paraffinic horticultural oil Sunspray Ultra-Fine were moderately effective in controlling powdery mildew, and significantly improved plant growth. Weekly applications of biorational fungicides were more effective than 14-day applications and similar to conventional fungicides applied every 14 days (28). Spray regimes with propiconazole rotations at 7-day intervals were more effective than 14-day applications of propiconazole. Using biorationals in rotation with conventional fungicides at 14-day application intervals was as effective as using fungicides alone. Incorporating biorational products and/or biopesticides in fungicide rotations reduced fungicide use by 56% in weekly applications and 66% in bimonthly applications (Table 1) (28).

Although there has been increasing research to identify new biorational products for powdery mildew management, conventional fungicides have remained competitive over biorational fungicides. Efficacy of conventional fungicides has had the test of time and won grower confidence over biorational products. In addition, conventional fungicides have longer residual control, requiring fewer spray treatments and lower labor costs. Using biorational products as a component of fungicide rotations has proven to reduce fungicide usage and maintain the number of sprays and same level of disease control (28). However,

marketing and availability of biorationals have not kept up with competing fungicides. Marketing strategies to improve the adoption of biorational products should include information on fungicide/biorational rotations.

Microbial agents such as bacteria, fungi, and yeast that are effective in powdery mildew control have been identified (26,29,31). Selected biological control agents might add to the list of biorational pesticides and provide alternatives to traditional fungicides for dogwood powdery mildew. The ultimate goal of reducing fungicide use in dogwood production will likely be accomplished by using different biorational fungicides in rotations with traditional fungicides.

## **Highlights**

Powdery mildew continues to be the greatest detriment to production of flowering dogwood in the United States and other countries. Fungicide programs, which are very effective in controlling the disease, require spray applications at regular intervals throughout the growing season. These applications add significantly to production costs that may be cost-prohibitive to many small- to mid-size nursery producers across the mid-south. In the deep-south, nurseries have ceased growing flowering dogwood for this reason. Biorational chemical candidates and perhaps even some biological control organisms hold some promise for managing the disease, but additional research is required before these strategies will be viable op-

Table 1. Comparisons of powdery mildew disease index on dogwood seedlings treated with biorational products alone and in rotation with propiconazole at 7- and 14-day intervals in field environments (modified from Mmbaga and Sauve [28] and Mmbaga and Sheng [30])

Treatment		Disease indexy		
	Interval (day)	4-year seedling	5-year seedling	Overall mean
Equate	14	2.2 efg	1.9 cde	2.1
Equate	7	0.9 ijkl	0.9 ghi	0.9
Armicarb 100	14	2.6 ef	1.2 defhi	1.9
Armicarb 100	7	0.9 ijkl	0.9 ghi	0.9
Palmolive	14	2.0 efgh	0.9 ghij	1.5
Palmolive	7	0.6 kl	0.5 j	0.6
Ajax	14	1.7 fghi	1.5 defg	1.6
Ajax	7	0.7 jk	0.7 hij	0.7
Propiconazole	14	1.5 ghijk	2.0 cd	1.8
Propiconazole	28	3.6 bcd	2.9 b	3.3
Equate/propiconazole <sup>z</sup>	14	1.4 hijk	1.1 efg	1.3
Equate/propiconazole <sup>z</sup>	7	0.51	1.0 fgh	0.8
Armicarb/propiconazole <sup>z</sup>	14	2.7 de	1.4 def	2.1
Armicarb/propiconazole <sup>z</sup>	7	0.51	0.6 ij	0.6
Palmolive/propiconazole <sup>z</sup>	14	1.5 hijk	1.2 def	1.4
Palmolive/propiconazole <sup>z</sup>	7	0.51	0.6 ij	0.6
Ajax/propiconazole <sup>z</sup>	14	1.6 hij	1.1 efg	1.4
Ajax/propiconazole <sup>z</sup>	7	0.51	0.9 ghi	0.7
Water control		5.0 a	4.1 a	4.6

y Disease index was assessed on a scale of 0 (no symptom) to 5 (100% leaf covered with powdery mildew signs and symptoms). Means followed by same letters in the same column were not significantly different as determined by least significant difference test (P = 0.05).

tions. Unfortunately, these approaches suffer from some of the same constraints as fungicide applications, in that they may require repeated and expensive applications of materials over the entire growing season. Natural resistance to powdery mildew has been documented in flowering dogwood (19,20,50,51), and this strategy appears to be the most cost effective way for managing powdery mildew in nurseries and landscapes. Breeding and developing new cultivars for powdery mildew resistance is somewhat problematic. Besides the long generation time, about 7 years, incompatibility between the F<sub>1</sub> generation and either parent occurs as well as inbreeding depression; almost all BC1 generation plants are weak and have failed to grow and flourish. Our group will continue to select and research alternative breeding methods that are intended to introduce new cultivars of flowering dogwood that have resistance to powdery mildew.

## Acknowledgments

We thank the financial supporters of our research and extension projects including the USDA-ARS (58-6404-7-213), Horticulture Research Institute (No. 1150), USDA/CSREES Grant No. 98-38814-6236, USDA/CSREES Grant No. TENX-2002 38814-12721, and USDA/CSREES Grant No. TENX-2003-03985.

#### **Literatures Cited**

- 1. Burrill, T. J., and Earle, F. S. 1887. Parasitic fungi of Illinois. Part II - Erysipheae. Pages 387-428 in: Bull. Ill. State Lab. Nat. Hist. Vol. II I W Franks and Sons Peoria II.
- 2. Daughtrey, M. L., and Hagan, A. K. 2001. Dogwood diseases. Pages 124-132 in: Diseases of Woody Ornamentals and Trees in Nurseries. R. K. Jones and D. M. Benson, eds. American Phytopathological Society, St. Paul, MN.
- 3. Daughtrey, M. L., Hibben, C. R., Britton, K. O., Windham, M. T., and Redlin, S. C. 1996. Dogwood anthracnose: Understanding a disease new to North America. Plant Dis. 80:349-358
- 4. Doney, J., Hartman, J., Johnson, M., Fountain, W., and McNiel, R. 1998. Evaluation of fungicides for dogwood powdery mildew control. Fungic. Nematicide Tests 53:478.
- 5. Farr, D. F., Bills, G. F., Chamuris, G. P., and Rossman, A. Y. 1989. Fungi on Plants and Plant Products in the United States. American Phytopathological Society, St. Paul, MN.
- 6. Garibaldi, A., Bertetti, D., and Gullino, M. L. 2009. First report of powdery mildew caused by Erysiphe pulchra on Cornus florida in Italy. Plant Dis. 93:320.
- 7. Hagan, A. K., Hardin, B., Gilliam, C. H., Keever, G. J., Williams, J. D., and Eakes, J. 1998. Susceptibility of cultivars of several dogwood taxa to powdery mildew and spot anthracnose, J. Environ, Hortic, 16:147-151.
- 8. Hagan, A. K., and Mullen, J. M. 1997. Powdery mildew on dogwood. Alabama Coop. Ext. Sys. Circ. ANR-1501.
- 9. Hagan, A. K., Rivas-Davila, M. E., Olive, J., and Stephenson, J. 2002. Comparison of registered fungicides for the control of powdery mildew in flowering dogwood. Fungic. Nematicide Tests 57:OT08.
- 10. Hagan, A. K., Rivas-Davila, M. E., Olive, J., and Stephenson, J. 2005. Effect of application rate and treatment interval on the control of powdery mildew on flowering dogwood with Compass. Fungic. Nematicide Tests 60:OT027.

<sup>&</sup>lt;sup>z</sup> Biorational/propiconazole rotations consisted of three applications of biorational products followed by one application of propiconazole.



Yonghao Li



Margaret T. Mmbaga



**Alan Windham** 



Mark Windham



Robert N. Trigiano

Dr. Li received a B.S. in plant protection and an M.S. in plant pathology from Northeast Agricultural University, China and a Ph.D. in plant pathology from the University of Arkansas. He currently is a postdoctoral research associate in the Department of Entomology and Plant Pathology at the University of Tennessee. Prior to moving to the United States in 1999, he was a full professor in the Department of Plant Pathology at Northeast Agricultural University, China. His research interests have been focused on host resistance and disease epidemiology.

Dr. Mmbaga received her Ph.D. from the Department of Plant Pathology, University of Wisconsin-Madison. She is currently a research professor at Tennessee State University, School of Agriculture and Consumer Sciences, Otis Floyd Research Center, McMinnville, TN. Prior to joining the Tennessee group, she worked on bean rust and host resistance to Uromyces appendiculatus in Tanzania and at the University of Nebraska, Lincoln. Her current research interests are focused on diseases that impact the nursery and ornamental industry. This research concentrates on disease management using host resistance and biopesticides such as biological agents and biorational products to reduce the amount of conventional pesticides used in the nursery production system.

Dr. Alan Windham is professor of plant pathology with UT Extension in the University of Tennessee Institute of Agriculture. He is stationed at the Soil, Plant and Pest Center at the Ellington Agricultural Center in Nashville. He teaches a graduate level course in disease diagnosis. His current research interests include the diagnosis and management of emerging diseases of ornamental plants and turf and the development of disease-resistant ornamental plants.

Dr. Mark Windham is a professor and distinguished chair of ornamental diseases with Tennessee AgRearch within the University of Tennessee Institute of Agriculture. He teaches classes in introductory plant pathology, diseases and insects of ornamentals, epidemiology and disease control of fungal pathogens, and insects and diseases of forest trees. His current research interests include the diseases of African violets, daylilies, dogwoods, hydrangeas, redbuds, roses, and switchgrass.

Dr. Trigiano is professor of ornamental plant biotechnology and plant pathology with UT Agricultural Research in the Tennessee Institute of Agriculture. He is a member of the Department of Entomology and Plant Pathology in Knoxville and teaches graduate level courses in mycology and plant microtechnique. His current research interests include developing/breeding ornamental plants for disease resistance using molecular markers, diseases of ornamental plants, and turf and biotechnology of ornamental plants.

- 11. Hagan, A. K., Rivas-Davila, M. E., Olive, J., and Stephenson, J. 2005. Determining the treatment window for controlling powdery mildew on flowering dogwood. Fungic. Nematicide Tests 60:OT031.
- 12. Halls, L. K. 1977. Southern fruit producing woody plants used by wildlife. U.S. Dep. Agric. For. Serv. Southeast. For. Exp. Stn. Gen. Tech. Rep. SO-16.
- 13. Halls, L. K., and Oefinger, S. W., Jr. 1969. Flowering dogwood. La. Conserv. 21:5.
- 14. Homma, Y., Arimoto, Y., and Misato, T. 1981. Effect of sodium bicarbonate on each growth stage of Sphaerotheca fuliginea in its life cycle. J. Pestic. Sci. 6:201-209.
- 15. Johnson, M. P., Hartman, J. R., McNiel, R. E., and Fountain, W. M. 2001. Evaluation of dogwood and birch species and cultivars for resistance to key insect pests and diseases. J. Environ. Hortic. 19:73-78.
- 16. Klein, L. A., Windham, M. T., and Trigiano, R. N. 1998. Natural occurrence of Microsphaera pulchra and Phyllactinia guttata on two Cornus species. Plant Dis. 82:383-385.
- 17. Li, Y. H., Windham, M. T., Trigiano, R. N., Fare, D. C., Spiers, J. M., and Copes, W. E. 2005. Spore germination, infection structure formation, and colony development of Erysiphe pulchra on dogwood leaves and glass slides. Plant Dis. 89:1301-1304.
- 18. Li, Y. H., Windham, M. T., Trigiano, R. N., Fare, D. C., Spiers, J. M., and Copes, W. E. 2006. Epidemiology of powdery mildew on resistant and susceptible flowering dogwood cultivars. Proc. Southern Nurs. Assoc. Res. Conf.
- 19. Li, Y. H., Windham, M. T., Trigiano, R. N., Fare, D. C., Spiers, J. M., and Copes, W. E. 2006. Development of Erysiphe pulchra, the causal agent of powdery mildew, on resistant and susceptible dogwood leaf disks. Can. J. Plant Pathol. 28:1-6.
- 20. Li, Y. H., Windham, M. T., Trigiano, R. N., Fare, D. C., Spiers, J. M., and Copes, W. E. 2007. Evaluation for resistance to powdery mildew in Cornus species and hybrids using a leaf disk assay. J. Environ. Hortic. 25:131-133.
- 21. McRitchie, J. J. 1994. Powdery mildew of flowering dogwood. Plant Pathol. Circ. No. 368. Gainesville, FL.
- 22. Mitchell, W. A., Gibbs, P. A., and Martin, C. O. 1988. Flowering dogwood (Cornus florida) Section. 7.5.9, US Army Corps of Engineers Wildlife Resources Management Manual. Tech. Rep. EL-88-9.
- 23. Mmbaga, M. T. 2000. Winter survival and source of primary inoculum of powdery mildew of dogwood in Tennessee. Plant Dis.

- 84:574-579.
- 24. Mmbaga, M. T. 2002. Ascocarp formation and survival and primary inoculum in Erysiphe (Sect. Microsphaera) pulchra in dogwood powdery mildew. Ann Appl. Biol. 141:153-161.
- Mmbaga, M. T., Klopfenstein, N. B., Kim, M. S., and Mmbaga, N. C. 2004. PCR-based identification of Erysiphe pulchra and Phyllactinia guttata from Cornus florida using ITSspecific primers. For. Pathol. 34:321-328.
- Mmbaga, M. T., Mrema, F. A., and Sauve, R. J. 2007. Identification of microorganisms for biological control of powdery mildew in Cornus florida. Biol. Control J. 44:67-72.
- 27. Mmbaga, M. T., and Sauvé, R. J. 2004. Multiple disease resistance in dogwoods (Cornus spp.) to foliar pathogens. J. Arbor 30:101-107.
- 28. Mmbaga, M. T., and Sauve, R. J. 2004. Management of powdery mildew in flowering dogwood in the field with biorational and conventional fungicides. Can. J. Plant Sci. 84:837-844.
- 29. Mmbaga, M. T., and Sauvé, R. J. 2009. Epiphytic microbial communities on foliage of fungicide treated and non-treated flowering dogwoods. Biol. Control J. 46:97-104.
- Mmbaga, M. T., and Sheng, H. 2002. Evaluation of biorational products for powdery mildew disease management in Cornus florida. J. Environ. Hortic. 20:113-117.
- 31. Mrema, F. A., and Mmbaga, M. T. 2006. Bacterial agents for biocontrol of powdery mildew in dogwood. Proc. Southern Nurs. Assoc. Res. Conf. 51:219-223
- 32. Mulrooney, R. P., and Gregory, N. F. 2004. Evaluation of fungicides for control of powdery mildew in flowering dogwood. Fungic. Nematicide Test 59:OT028.
- 33. Mulrooney, R. P., and Gregory, N. F. 2005. Evaluation of fungicides for control of powdery mildew on flowering dogwood. Fungic. Nematicide Tests 60:OT016.
- Northover, J., and Schneider, K. E. 1996. Physical modes of action of petroleum and plant oils on powdery and downy mildews of grapevines. Plant Dis. 80:544-550.
- 35. Olive, J. W., and Hagan, A. K. 2000. Control of powdery mildew on dogwood. Proc. Southern Nurs. Assoc. Res. Conf. 45:209-210.
- 36. Olive, J. W., Hagan, A. K., and Parrott, L. C. 1998. Evaluation of selected fungicides for control of powdery mildew of dogwood. Proc. Southern Nurs. Assoc. Res. Conf. 43:221-222.
- 37. Pimentel, G., Carris, L. M., Levy, L., and Meyer, R. J. 1998. Genetic variability among isolates of Tilletia barclayana, T. indica and allied species. Mycologia 90:1017-1027.
- Ranney, T. G., Grand, L. F., and Knighten, J. L. 1995. Susceptibility of cultivars and hybrids

- of kousa dogwood to dogwood anthracnose and powdery mildew. J. Arboric. 21:11-16.
- 39. Redlin, S. C. 1991. Discula destructiva sp. nov., cause of dogwood anthracnose. Mycologia 83:633-642.
- 40. Reuveni, M., Agapov, V., and Reveni, R. 1993. Induction of systemic resistance to powdery mildew and growth increase in cucumber by phosphates. Biol. Agric. Hortic. 9:305-315.
- 41. Smith, V. L. 1999. First report of powdery mildew on Cornus florida in Connecticut caused by Microsphaera pulchra. Plant Dis.
- 42. Staub, T. 1991. Fungicide resistance: Practical experience with antiresistance strategies and the role of integrated use. Annu. Rev. Phytopathol. 29:421-442.
- Stimmel, J. F. 1996. Biorational controls -Problem-free? Regul. Hortic. 22:13.
- 44. Williamson, M. R., and Blake, J. H. 1999. First report of the teleomorph of an Oidium sp. causing powdery mildew on flowering dogwood in South Carolina. Plant Dis. 83:200.
- 45. Windham, A. S., Trigiano, R. N., and Windham, M. T. 2006. 'Appalachian Joy': A new flowering dogwood cultivar with multiple bracts and powdery mildew resistance. Proc. Southern Nurs. Res. Conf. 51:614-615.
- 46. Windham, M. T., Trigiano, R. T., and Windham, A. S. 2005. Susceptibility of Cornus species to two genera of powdery mildew. J. Environ. Hortic. 23:190-192.
- 47. Windham, M. T., Windham, A. S., and Halcomb, M. A. 1998. Chemical control of powdery mildew of flowering dogwood. Proc. Southern Nurs. Res. Conf. 43:251-252
- 48. Windham, M. T., Windham, A. S., and Halcomb, M. A. 1999. Enhancement of growth of flowering dogwood using fungicides to control powdery mildew. Proc. Southern Nurs. Res. Conf. 44:224-225.
- 49. Windham, M. T., Windham, A. S., and Halcomb, M. A. 2000. Control of powdery mildew in dogwoods with fungicides. Proc. Southern Nurs. Res. Conf. 45:207-208.
- 50. Windham, M. T., and Witte, W. T. 1998. Naturally occurring resistance to powdery mildew in seedlings of Cornus florida. J. Environ. Hortic. 16:173-175.
- 51. Windham, M. T., Witte, W. T., and Trigiano, R. N. 2003. Three white-bracted cultivars of Cornus florida resistant to powdery mildew. HortScience 38:1253-1255.
- 52. Wolfe, M. S. 1984. Trying to understand and control powdery mildew. Plant Pathol. 33:451-
- Yarwood, C. E. 1957. Powdery mildews. Bot. Rev. 23:235-301.