Esca (Black Measles) and Brown Wood-Streaking: Two Old and Elusive Diseases of Grapevines

“Esca” disease of grapevine has long been known wherever grapes are grown. The disease may be as old as vine cultivation itself. References to esca-like symptoms are found in several ancient Greek and Latin works. Greater descriptive accuracy is found in medieval works such as the Kitab al-Felalab by Ibn al-Awam, a Spanish Muslim who lived in Seville at the end of the twelfth century, and the Opus Ruralium Commodorum by Pietro de’ Crescenzi, born around 1233 in Bologna.

Esca is a Latin word meaning food, aliment, or, figuratively, bait. The name directly refers to the fruiting bodies of certain wood-rotting fungi. For example, Fomes, which in Latin means “tinder,” is the name of a genus of basidiomycetes once used to make a dry, easily ignited material suitable to start fires when using flints. Wood decayed by these fungi, including rotted grapevine wood, burned slowly and was used to keep fires aglow without a flame.

Esca is a complex disease that comprises an array of symptoms, some of which have locally given their own names to the disease. For example in California, the dark, tiny spotting of the grapes is called “black measles.” In many grapevine-growing areas of the world, the sudden wilting of esca-affected vines in summer has earned the name of a genus of basidiomycetes to cause esca and related diseases.

In the 1990s, studies on esca and its etiology have intensified. This came after a dramatic upsurge in the disease, especially in Germany, Italy, and Greece, where the arsenites, traditionally used to keep esca under control, have been banned and were replaced by less effective fungicides. How- ever, even in France, Portugal, and Spain, where restricted use of sodium arsenite is still permitted, esca is widespread in all vine-growing regions.

This paper refers mostly to conditions in the Mediterranean area and focuses on the progress achieved in our understanding of esca over the last few years. Reference is also made to a decline of young grapevines caused by some of the same mitosporic fungi as those commonly associated with esca in adult vines. Finally, the relations between esca and related syndromes or diseases are also discussed.

**Syndromes Involved**

Esca is a complex disease whose symptoms arise from structural and physiological changes that cannot be reduced to a simple scheme of cause and effect. Consequently, symptoms that look alike may have different causes.

Apart from the occasional delayed sprouting of esca-affected vines in the spring, external symptoms in the Northern Hemisphere develop on the crown between June and September, either through the entire vine or on single branches. The development of esca syndrome can be of two types, chronic or acute.

**Chronic esca syndrome.** Chronic esca comprises symptoms inside the trunk and larger branches, on the shoots and smaller branches, on the leaves, and on the berries. Symptoms inside trunk and main branches. On adult plants (8 to 10 years and older), the most common internal symptom is white rot, which gradually changes the hard wood to a soft, friable, spongy mass. Rotted tissues appear creamy yellow or whitish, and in cross section are often bordered by a thick black or dark brown line separating rotted from nonde- cayed wood. The decay usually starts from a large pruning wound on the trunk extending into the woody tissue and either remains restricted to the older part of the wood or spreads along a sector in the woody cylinder (Figs. 1 and 2). Sometimes the rot approaches or reaches the surface, causing cracks along the trunk (this is called “mal dello spacco”—cracking disease—in Italy; Fig. 3). The decay usually extends downward and upward along the trunk, decreasing in size away from its point of origin, but it seldom develops...
below the graft union and never affects the roots. However, old, ungrafted mother vines of American or Euro-American hybrids may show the same internal symptoms as grafted European cultivars.

Various other types of wood deterioration become visible together with or even preceding white rot: (i) small, dark brown or black spots in cross section appear as deep brown or black streaks or columns in longitudinal section. In cross section, the spots are sparsely distributed or arranged in groups around an annual growth ring or in the woody tissues close to the pith (Figs. 1A and B, 2); (ii) pink-brown or dark reddish brown areas often develop from black spots (Figs. 1B, 2), mainly in the core of the trunk or on the margin of decayed or necrotic tissues, separating such tissues from apparently healthy tissue; and (iii) brown areas of varying shade and texture are intermingled with the above types of discoloration, often in sectors connected with large wounds (Fig. 2).

On plants younger than 8 to 10 years, external leaf and berry symptoms of esca (see below) are often associated with a limited wood discoloration inside the trunk. However, white rot in the trunk of young grapevines with leaf symptoms of esca is observed very rarely.

The various types of wood discoloration and decay are the result of a number of structural and physiological changes: (i) physical and chemical changes caused by the introduction of air and water into wounds (1) and the host reactions to such wounding (e.g., degradation, oxidation, and darkening of tissue components); (ii) changes caused by cellulolytic and ligninolytic enzymes produced by the fungi associated with esca (10,54); (iii) tylosis induced by growth-regulating substances, and vascular occlusion due to gels and gums secreted by the diseased xylem parenchyma cells, or even to high-molecular-weight compounds produced by the pathogen(s); and (iv) necrosis of xylem parenchyma cells as the result of diffusion of the pathogen’s toxins (76), host reaction products (e.g., phytoalexins), or both.

Symptoms on leaves. Symptoms on leaves consist of light green or chlorotic, rounded or irregular spots between the veins or along the leaf margins that usually spread outward to the distal parts of the shoots (Fig. 4). The spots, initially small and scattered over the lamina, gradually expand and coalesce, become partly necrotic, and ultimately leave only a narrow strip of unaffected green tissue along the main veins (Fig. 4A). As the chlorotic tissue turns yellow-brown or red-brown (Fig. 4B and C) (or, in the case of certain cultivars, develops rust-colored necrotic areas with dark red margins), the diseased leaves assume a “tiger-stripes” pattern (Fig. 4D). Sometimes the necrotic areas of the lamina dry out and become detached, leaving irregular leaf margins. Other symptoms, such as clearing, puckering, glistening, and distortion of the leaf lamina (9), are less common.

The appearance of wilt symptoms in the crown could be explained in terms of xylem dysfunction. However, for various reasons, it seems unlikely that leaf necrosis and wilting of shoots and fruits are due to water stress caused by vascular occlusion (78). Vascular occlusion, where it exists, affects only a part of the trunk, mainly the nonfunctional woody rings. The functional wood is usually not discolored, but even if it were, the xylem vessels of woody dicotyledons form an integrated system that can compensate for any change in water potential that might occur as a result. In any case, only two or three annual increments of functional xylem are needed to sustain vigorous shoot growth and abundant grape production (59).

It seems more likely that the foliar symptoms of esca are mainly caused by substances that originate in the discolored woody tissues of the trunk and branches and are then translocated to the leaves in the transpiration stream. These substances can be reaction products of the wood, phytotoxic metabolites excreted by esca

**Fig. 1.** Cross sections through the trunk of a 25-year-old grape cv. Sangiovese vine affected by esca. The trunk was cut into 3- to 5-cm-thick sections, which on the photograph show wood symptoms in a clockwise direction, starting from the trunk base up to the portion where two large pruning wounds were made. The lowest trunk sections (A–C) show black spotting, brown-red wood, and initial symptoms of white rot (slightly yellow in the picture) starting from the pith. The rotted wood gradually extends toward the large wounds (D). In the upper trunk sections (E and F), yellowish, soft, and spongy rotted wood (white rot) predominate, being bordered by a thick black line that in turn is surrounded by a marginal band of brownish red wood.
fungi, or a combination of both. In a recent study, foliar symptoms similar to those caused by esca were produced when culture filtrates (or their fractions) of three fungi associated with the disease (see below) were absorbed by detached grapevine leaves (Fig. 5). Polysaccharides (pullulans) were purified from the cultures of two of the above fungi (*Phaeoacremonium*) and were shown to be nonselectively toxic to cuttings of grapevine and other plants (76).

**Symptoms on shoots and branches.** Symptoms can start in spring with delayed and weak growth and end in autumn with reduced lignification of the canes. In late spring and summer, an irreversible wilt of some shoots or of an entire branch may occur.

**Symptoms on berries and clusters.** Spotting of berries, sometimes without accompanying foliar symptoms, is common in California and southern Italy and has also been reported from France (Alsace) on adult as well as young vines, especially white grape cultivars (9,19,31,33). Minute dark brown, violet, or purple spots develop more or less abundantly on the skin of the berry, hence the name “black measles” given to this condition in California (Fig. 6A). Masses of tiny spots are irregularly scattered, especially toward the distal end of the berry, although they are sometimes distributed in longitudinal bands (Fig. 6B). More rarely, the spots cover almost the whole berry, which then turns brown or violet-gray.

Spots are due to browning and necrosis of groups of epidermal and hypodermal cells near the terminal part of a xylem vessel and do not extend to the underlying tissues (32). Within a grape cluster, the severity of spotting may differ from berry to berry. Even with only light spotting, table grapes are not easily sold on the mar-

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**Fig. 2.** Pattern of wood colonization of a grapevine trunk by esca fungi (left, transverse sections; right, median longitudinal section). The occurrence (percent isolation) of three fungal species (*Fomitiporia punctata* [Fp], *Phaeoacremonium chlamydosporum* [Pch], and *P. aleophilum* [Pal]), as found in a survey in central Italy, is reported for each trunk portion. The white rot fungus, *F. punctata*, prevails in the completely decayed wood and decreases toward the healthy wood as *P. chlamydosporum* and *P. aleophilum* become dominant. These two fungi are mainly found in the black streaks and brown-red wood, both in the wood portions around the pith and at the margin of the decayed wood tissues. Green arrows, pruning wounds; BN, brown necrosis; BRW, brown-red wood; AP, altered pith; BS, brown to black streaks, appearing as black spots in cross section; DP, decayed pith; BL, black line bordering the decayed wood; DW, completely decayed wood (white rot).

**Fig. 3.** Cracking of a grapevine trunk in connection with a sector of decayed wood.
Berries with heavy spotting often show skin cracks and even longitudinal or transverse splits (Fig. 6B). Such fruits shrivel and dry up or become prey to soft rotting fungi or bacteria.

Sometimes clusters on diseased vines show late ripening with low sugar content and altered berry flavor, or a slow wilt of the distal parts, whose berries lose turgor and fail to ripen.

As with foliar symptoms, fruit symptoms do not necessarily appear on the same diseased plant every year. Symptoms can affect only some berries in a single cluster, all clusters on a branch, or a number of branches.

The dark spotting of grape berries could result from diffusion through the vascular system.
system of enzymes that oxidize or polymerize phenolic substances in the epicarp, in conditions of high irradiation, but more probably it is caused by toxins translocated to the fruits by the same route.

**Acute esca syndrome.** “Vine apoplexy” appears suddenly in the middle of summer as a rapid basipetal wilt of entire vines, including the clusters, rarely of single branches (Fig. 7). Green, healthy-looking leaves turn pale green, then gray-green, and quickly wither, drying up completely in a few days.

Apoplexy is thought to be favored by hot summers, in particular, when rainfall is followed by dry, hot weather. It may be connected with a rapid rise in the concentration and activity of toxic metabolites in the crown when the rate of transpiration is high.

**Related Syndromes**

Wood discoloration with brown or black streaks (brown wood-streaking) is a common internal symptom of esca-infected grapevines, but it is also similar to the wood streaking of declining vines reported by Petri in southern Italy (Sicily) at the beginning of the twentieth century (58). Externally, these plants showed a general decline and stunted growth, a condition often confused with viral diseases (arricciamento or court-noué), whose cause was unknown at the time. The internal symptoms were described by Petri as “gommosi del legno da ferita” (wound-related wood gummosis) and “venature brune del legno” (brown to black wood streaking). These longitudinal streaks or columns of discolored wood are similar to a condition called bacterial gummosis by Prillieux and Delacroix (60). They originate in pruning wounds or at the graft union and appear as minute black or dark brown spots in cross section. Sometimes the streaks are even formed by isolated xylem vessels and the related parenchyma cells. Brown or black streaking, with the presence of gummy masses in the xylem vessels, can extend so deeply into the trunk as to reach the roots.

According to Petri, brown wood-streaking predisposed the vine plant to wood decay or even represented the first signs of esca.

Syndromes in many respects similar to the syndrome observed by Petri in Italy have been reported from other parts of the world, particularly from young vineyards (see below). Affected vines with such a syndrome sometimes show a sudden collapse, but more frequently a progressive decline, slow dieback, and stunted growth, with various foliar symptoms (chlorosis, marginal necrosis, wilting). In every case, they show blackish or brown wood-streaking and xylem gummosis similar to the wood discoloration observed in esca-affected adult vines. Table 1 shows reported examples of such syndromes and of browning of grapevine wood in propagating material examined before planting.

**Pathogen or Pathogens?**

Despite observations made by researchers for more than a century, the etiology of esca is still controversial, and several fungi have been associated with it.

**White rot basidiomycetes.** The association of esca symptoms with wood decay dates back to early works on the subject (66), but both the number and identity of the fungi causing esca have long been debated. In France, Italy, and other countries (1,9,28–31,43,61,62,65,78,79), *P. igniarius* and, to a lesser extent, *S. hirsutum* were for many years the basidiomycetes most commonly associated with white rot of grapevine. For this reason, one of the other of these fungi was also presumed to be the real cause of esca. This etiology, however, was never established by reproducing all the disease symptoms.

Knowledge about the basidiomycetous fungi involved in wood decay of vines has progressed in recent years (12,13,38,39,44,53,69).

In Italy, white rot of vine wood was only very infrequently associated with *S. hirsutum*...
tum, but the great majority of white rot samples examined was colonized by a fungus that was initially classified as *P. igniarius* on the basis of in vitro cultures. Work carried out in France (40) in the same years also led to the isolation of *S. hirsutum* with very low frequency, but here the basidiomycete most commonly found was *Phellinus punctatus* (P. Karst.) Pilát. The correct name for this species is *Fomitiporia punctata* (Fr.) Murrill (25). Further studies of a group of strains isolated in Italy has shown that these strains were actually *F. punctata* (M. Fischer and L. Mugnai, unpublished). It seems therefore that previous records of *P. igniarius* were misidentifications of *F. punctata*, and that this is the basidiomycete commonly associated with decayed wood, at least in France and Italy.

**Nonbasidiomycetous fungi associated with esca.** In some vine-growing areas of the Mediterranean where the prevailing climate is of the continental type, the ascomycete *Eutypa lata* (Pers.:Fr.) Tul. & C. Tul. is quite frequently isolated from discolored vine wood, and this could be thought to complicate the picture of esca syndrome. *E. lata*, however, is known to cause a distinct disease, Eutypa dieback, with its own symptomatology. Sometimes Eutypa dieback and esca occur on the same vine during a growing season, the former producing reduced, bushy growth and leaf yellowing in May, the latter—on a different branch or even on the same branch—leaf chlorosis and necrosis in summer (45). In the more southerly vine-growing areas of Europe, *E. lata* either is not found at all or is rare on esca-affected vines. Another plant pathogenic ascomycete commonly isolated from the trunks and branches of esca-affected grapevines is *Botryosphaeria obtusa* (Schwein.) Shoemaker.

Chiarappa (9) reported that the brown-red marginal zones bordering the decayed wood of grapevines affected with black measles were colonized by a *Cephalosporium*-like species, which discolored vine-wood in vitro and produced brown streaks when inoculated into standing grapevines. In Europe, several mitosporic fungi, including undetermined species of *Cephalosporium, Acremonium, and Phialophora*, were isolated with relatively high frequency from the wood of esca-diseased grapevines (5, 20, 31, 44), but their role in the disease process has remained unknown. That a succession of fungi might be involved in the esca disease process was first proposed by the French researchers Larignon and Dubos (39), who suggested that each fungus was associated with a particular stage of wood discoloration. Species of *Phialophora, Cephalosporium, Table 1. Reports of declines of young grapevines associated with brown wood-streaking and of wood discoloration of rooted cuttings

<table>
<thead>
<tr>
<th>Syndrome or disease name</th>
<th>Associated microorganisms</th>
<th>Country and references</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brown wood-streaking (wound-related gummosis)</td>
<td>Two <em>Cephalosporium</em> species (possibly identical to <em>Phaeoacremonium chlamydosporum</em> and <em>P. aleophilum</em>)</td>
<td>Italy (58)</td>
</tr>
<tr>
<td>Wood discoloration of rooted cuttings before planting</td>
<td><em>Phaeoacremonium chlamydosporum</em></td>
<td>Italy (27)</td>
</tr>
<tr>
<td>Slow dieback</td>
<td><em>Phialophora parasitica</em> (=<em>P. chlamydosporum</em>)</td>
<td>South Africa (24)</td>
</tr>
<tr>
<td>Black goo</td>
<td><em>P. chlamydosporum</em></td>
<td>California (47, 48; L. Chiarappa, personal communication)</td>
</tr>
<tr>
<td><em>Phaeoacremonium</em> grapevine decline</td>
<td><em>P. chlamydosporum</em></td>
<td>Australia (56)</td>
</tr>
<tr>
<td>Wood discoloration of rooted cuttings before planting</td>
<td><em>P. aleophilum, P. inflatipes</em></td>
<td>California (68)</td>
</tr>
<tr>
<td>Decline of young grapevines</td>
<td><em>P. chlamydosporum</em></td>
<td>Italy and France (2, 77)</td>
</tr>
</tbody>
</table>

Fig. 7. A grapevine plant showing sectorial symptoms of apoplexy caused by esca disease. Green, healthy-looking leaves and grape clusters quickly wither, drying completely in a few days, but usually remain attached to the plant.
and Eutypa were supposed to be the first wood colonizers, or pioneer fungi. Italian workers also examined the initial stages of wood discoloration, described by them as brown-red wood and black streaks developing at the margin of already discolored wood (52,53). In these tests, two mitosporic fungi were consistently isolated from the discolored wood, and it appeared they could be related to the two main pioneer fungi identified by Larignon and Dubos: Cephalosporium sp. and Phialophora parasitica Ajello, L.K. Georg, & C.J.K. Wang. These two mitosporic fungi isolated in Italy were assigned to a single new genus, Phaeoacremonium, as P. chlamydosporum and P. aleophilum W. Gams, F.W. Crous, M.J. Wingfield & Mugnai (14). Interestingly, the strain of Cephalosporium (CBS 239.74) isolated by Chiarappa (9) in California from the discolored wood of black measles–affected vines, and later considered to be close to 

\[ \text{Phialophora parasitica} \] (34), was also eventually referred to P. chlamydosporum (14). These same two species of Phaeoacremonium were then reported to be the main pioneer fungi by Larignon and Dubos as well (40). Other Phaeoacremonium species isolated from grapevine are P. angustius and P. inflatipes W. Gams, P.W. Crous, & M.J. Wingfield (Table 2) (14).

More recently, Dupont et al. (21) provided morphological, cultural, and molecular evidence to show that P. chlamydosporum and P. aleophilum have a distant phylogenetic relationship to each other. The first species is closer to Phialophora verrucosa Medlar than to Phaeoacremonium parasiticum (Ajello et al.) W. Gams, Crous, & M.J. Wingfield. In light of these findings, the whole genus will have to be reassessed and the species P. chlamydosporum redescribed in another genus (P. W. Crous, personal communication).

**Fungi associated with the decline of young grapevines.** Some of the mitosporic fungi associated with esca are also involved in the decline of young grapevines and brown wood-streaking. Such involvement was first reported in Italy by Petri (58) on young grafted vines and mother plants. Declining vines showed brown or black streaking from which Petri isolated two strains (\( \alpha \) and \( \beta \)) of Cephalosporium and one strain of Acremonium that did not correspond to any species known at the time. On the basis of Petri’s descriptions, Cephalosporium strain \( \alpha \) and Acremonium sp. can be referred with sufficient accuracy to P. chlamydosporum and P. aleophilum, respectively. Petri also succeeded in reproducing brown streaking on standing vines by injecting the fungi isolated from diseased vines into the xylem elements through large wounds made on healthy-looking plants.

Since that time, cases of decline of young grapevines associated with brown-wood streaking and gummosis have been recorded in several countries under different names (Table 1). In South Africa, a slow decline of young vines was initially associated with P. parasitica (24). Subsequently, however, this fungus was also reassigned to P. chlamydosporum (strain STE-U 809) (14). In California, three species of Phaeoacremonium, P. chlamydosporum, P. inflatipes, and P. aleophilum, isolated from declining vines have been shown to be pathogenic to grape seedlings (68). Finally, in different areas of Italy, a heavy colonization by P. chlamydosporum was noted on declining 2-year-old vines (authors’ personal observations) (Fig. 8).

### Microorganisms in Wood of Esca-Diseased Grapevines

In general, two main models have been proposed to describe wood decay development in standing trees: the “conditioning of microenvironment” (CM model) (7) and the “compartmentalization of decay in trees” (CODIT model) (73).

The CODIT model postulates a microbial succession in the colonization of wood in standing trees subsequent to wounding. Initially, there is a host response to wounding; this is followed by the invasion of the modified wood substrate by nondecay microorganisms (often mitosporic fungi such as species of Phialophora or Trichocladium); and finally, the decay fungi infect wood that has been “conditioned” by the nondecay, pioneer organisms (42). Alternatively, decay fungi first infect the wood tissue, but then further colonization is blocked by the host defense reaction, which must be overcome by the pioneer fungi before it can proceed (75). In both cases, the pioneer fungi have the task of detoxifying pre- and postinfectional antimicrobial substances.

Most of the evidence on which the succession of microorganisms in wood decay is based can be seen in esca (38,40,52,53); however, not all findings fit this model. In the following description, the esca disease process will be described under headings consistent with the CODIT model.

**Compartmentalization, discoloration, and decay.** In the trunk of a grapevine affected with esca, an elongated portion of woody tissue begins to discolor, starting from a large pruning wound. The affected tissue soon turns brown. Larger wounds produce larger and longer brown columns, which may extend radially to a sector and reach the pith. However, the discolored wood does not indicate the full extent of fungal colonization, which appears as a brownish column tapering distally or as cones with the points outward spreading in both directions from the wound (Fig. 2).

In cross section, esca-affected vine trunks usually exhibit large necrotic areas, often first sectorial, then central in serial sections that are more distant from the wounds (Fig. 1). These necrotic zones are surrounded and preceded by small brown or black spots (Fig. 1A to F). Eventually, the wood decayed for the longest time begins to show white rot (Fig. 1D to F).

Microscopic examination of wood samples from esca-affected grapevines (52) shows that, at the advancing edge of the

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**Table 2. Species included in the genus Phaeoacremonium**

<table>
<thead>
<tr>
<th>Species</th>
<th>Host or substrate</th>
<th>Distribution</th>
</tr>
</thead>
<tbody>
<tr>
<td>P. aleophilum</td>
<td>Vitis vinifera</td>
<td>France, Italy, Yugoslavia, USA</td>
</tr>
<tr>
<td>P. angustius</td>
<td>V. vinifera, soil</td>
<td>Argentina, France, Italy, USA</td>
</tr>
<tr>
<td>P. chlamydosporum</td>
<td>V. vinifera</td>
<td>France, Italy, South Africa, USA</td>
</tr>
<tr>
<td>P. inflatipes</td>
<td>Humans, Nectandra sp., Sorbus intermedia, V. vinifera</td>
<td>Costa Rica, Finland, Germany, Tahiti, Venezuela, USA</td>
</tr>
<tr>
<td>P. parasiticum</td>
<td>Humans, Prunus armeniaca</td>
<td>Tunisia, USA</td>
</tr>
<tr>
<td>P. rubrigenum</td>
<td>Humans</td>
<td>USA</td>
</tr>
</tbody>
</table>

1 From references 14, 21, 40.
2 This species will be redispersed in another genus (P. W. Crous, personal communication).
discolored wood stripes, the first reaction of the xylem to fungal advance is almost always heavy tylosis and gummosis. The accumulation of tylosis and gels or dark brown gum gives rise to the discolored strands or streaks, which may extend longitudinally for tens of centimeters away from the wound, appearing in cross section as either isolated black spots or groups of spots around an annual ring.

When the colonizing fungi spread out into the functional sapwood, a more diffuse wood reaction causes the formation of brown-red or pink-brown areas, which often enclose the black streaks (Fig. 1A and B).

**Occurrence of different fungal species at different stages of wood decay.** The systematic isolation of fungi (Table 3) from the discolored or decayed wood of esca-diseased grapevines indicated a close relation between individual stages of wood deterioration and particular species of fungi (40,44,50,53). Up to 30 species of fungi were isolated from the initial stages of wood discoloration, but only a few of these were consistently present with relatively high frequency. The most frequent of these was *P. chlamydosporum* (approximately 64% of isolations from black streaks and 57% from brown-red wood in the survey illustrated in Table 3), often together with *P. aleophilum*. These species were also the most commonly isolated from the brown-red discolored wood when it surrounded white-rotted wood (Fig. 2). From the white-rotted wood itself, which is the last stage of wood degradation, *F. punctata* was most often isolated (Table 3, Fig. 2). *S. hirsutum* was always very rare.

The low isolation percentages of basidiozymes from the marginal areas of decayed wood, as well as from the brown/black streaks, were not due to the faster growth in vitro of concomitant fungi, as has been suggested for other pathosystems (42). Experiments on different culture media showed that isolates of *F. punctata* actually grew much faster than isolates of the two species of *Phaeoacremonium* (49).

For example, *P. chlamydosporum* required as long as 2 months to come out from wood chips and grow onto agar plates (Fig. 9).

**Decay progression is enhanced by substrate conditioning.** In the decayed trunks of many tree species (6,70–72,74), the development of reddish, brown, or brightly discolored wood is associated with an increase in the concentration of antimicrobial substances such as phenolic compounds produced by the xylem parenchyma cells in response to invasion, presumably in order to inhibit fungal growth (57). In many cases, it has been suggested that pioneer fungi are able to overcome the host defense reaction by growing despite a high concentration of phenolics and related oxidation products, which they detoxify, thereby making possible the growth of wood-decay agents (63 and literature cited therein). Indeed, an ability to grow despite the presence of fungal growth inhibitors produced by the woody tissues of the host tree may well be the most important function of pioneer microorganisms in the microbial succession theory.

The grapevine is a plant relatively rich in phenolics (64), compounds that could have a role in regulating fungal succession during wood decay.

Growth tests in vitro (49) showed that: (i) in a medium in which gallic acid was the sole carbon source, *P. chlamydosporum* was able to grow, but the growth of *F. punctata* was completely inhibited; and (ii) if the medium was first conditioned by culturing *P. chlamydosporum*, the subsequent growth of *F. punctata* was unhindered (Table 4). Other differences in the growth of pioneer and decay fungi on phenol-enriched media (Table 5), including differences in the total amount of phenols extracted from grapevine wood at variously discolored decay stages, were also recorded (38).

The wood-colonizing fungi show varying ability to utilize nutrients made available in injured tissues, such as the starch stored in abundance in the parenchyma rays. Microscopic examination of cross sections of esca-infected wood showed complete depletion of starch in the xylem tissue surrounding the brown-black streaks. The same picture was seen in the brown-red wood that developed later, but not in brown necrosis, nor in the discoloration.

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**Table 3. Most common fungi (isolation frequency >1%) isolated from each type of wood symptom detected in grape plants affected by esca**

<table>
<thead>
<tr>
<th>Type of wood symptom</th>
<th>Wood chips sampled (no.)</th>
<th>Wood chips colonized (%)</th>
<th>Fungal species (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>by fungi or bacteria</td>
<td><em>Phaeoacremonium</em></td>
</tr>
<tr>
<td>Healthy wood</td>
<td>345</td>
<td>18</td>
<td>3.2</td>
</tr>
<tr>
<td>Black to brown streaks</td>
<td>1,160</td>
<td>43.2</td>
<td>64.5</td>
</tr>
<tr>
<td>Brown-red wood</td>
<td>883</td>
<td>91.7</td>
<td>56.6</td>
</tr>
<tr>
<td>Cave or altered pith</td>
<td>529</td>
<td>78.3</td>
<td>18.4</td>
</tr>
<tr>
<td>Brown necrosis</td>
<td>1,269</td>
<td>81.7</td>
<td>26.8</td>
</tr>
<tr>
<td>White decay</td>
<td>310</td>
<td>90.1</td>
<td>5.7</td>
</tr>
<tr>
<td>Brown-red margin of decayed wood</td>
<td>199</td>
<td>68.8</td>
<td>26.3</td>
</tr>
</tbody>
</table>

1 Cultivars sampled: Cabernet, Canaiolo, Garganega, Merlot, Moscato, Regina, Riesling italico, Sangiovese, Trebbiano toscano, Trebbiano d’Abruzzo. Italian regions surveyed were: Abruzzi, Apulia, Friuli, Lombardy, Tuscany, Umbria, Veneto, Sicily.

2 Number of wood chips colonized as percentage of number of wood chips sampled for each symptom.

3 Number of wood chips infected with each fungus as percentage of number of colonized wood chips for each symptom.
observed in the wood of rooted vine cuttings (L. Mugnai, personal observation). Enzyme tests in vitro showed that *P. chlamydosporum* and *P. aleophilum* had a high starch-degrading activity (Table 6), whereas that of *F. punctata* was very low. The well-known ligninolytic activity of wood-rotting basidiomycetes, which enables them to degrade woody tissues completely (10,41,54), suggests that *F. punctata* has a key role in the development of white rot in esca-affected vines. Enzyme tests in vitro showed that *F. punctata* has a key role in the development of white rot in esca-affected vines. Inoculation with decay fungi alone does not always produce decay. For many years, inoculation of white rot basidiomycetes into healthy grapevines has failed to reproduce, not only esca leaf symptoms, but even wood decay. Recently, however, Graniti et al. in Italy (51) (Table 7) and Chiarappa in California (11) succeeded in reproducing white rot (Fig. 10A).

If these results are confirmed, they suggest either that *F. punctata* is a primary pathogen of grapevine that initiates and completes the entire wood decay process if it has a suitable wound and sufficient time, or that the model of Shortle and Cowling (75) applies here: the decay agent begins colonization, but its continued spread depends upon the prior detoxification of inhibitory compounds produced in the reaction zone of the plant. This process is usually carried out by other wood-colonizing fungi, but it can also occur, although more slowly, through the sole action of the decay agent itself.

An overall view of the evidence suggests that while some data support the hypothesis of microbial succession in the development of esca, other data, particularly the ability of both *F. punctata* and certain species of *Phaeoacremonium* to act as primary pathogens, militate against it. In the case of an association, whereas prior colonization of the vineyard by *Phaeoacremonium* or other fungi will facilitate the progression of *F. punctata* into the host, such colonization is not strictly necessary. When these two pathogenic fungi are present together in the woody tissue of a grapevine, two different syndromes or diseases may become concurrent. When the fungi are not present together at the appropriate time, each may induce its own syndrome or disease.

### Table 4. Growth of *Fomitiporia punctata* on a gallic acid medium conditioned by previous growth of *Phaeoacremonium chlamydosporum*

<table>
<thead>
<tr>
<th>Preinoculation</th>
<th>Additions to medium before/after growth of <em>F. punctata</em> (g/liter)</th>
<th>Mean wt of mycelium of <em>F. punctata</em> (g/liter)</th>
</tr>
</thead>
<tbody>
<tr>
<td>None</td>
<td>Glucose 10/no addition</td>
<td>2.7 ab*</td>
</tr>
<tr>
<td></td>
<td>Glucose 10/gallic acid 8.1</td>
<td>0.0 a</td>
</tr>
<tr>
<td></td>
<td>Gallic acid 1/no addition</td>
<td>0.0 a</td>
</tr>
<tr>
<td></td>
<td>Gallic acid 1/glucose 8.76</td>
<td>0.0 a</td>
</tr>
<tr>
<td><em>P. chlamydosporum</em></td>
<td>Glucose 10/gallic acid 8.1</td>
<td>4.35 bc</td>
</tr>
<tr>
<td></td>
<td>Gallic acid 1/glucose 8.76</td>
<td>3.36 ab</td>
</tr>
</tbody>
</table>

* P. *chlamydosporum* was grown in the mineral medium of Eriksson and Pettersson (23) containing glucose (10 g/liter) or gallic acid (1 g/liter). After incubation for 25 days at 25°C on a rotary shaker and before inoculation with *F. punctata*, the culture medium was filter-sterilized and the resulting culture filtrate was not supplemented, or supplemented with 8.1 g of gallic acid per liter if it contained glucose, or with 8.76 g of glucose per liter if it contained gallic acid.

* Mean oven-dried weight, growth after 40 days.

* Means with different letters differed significantly at *P* = 0.01 (Tukey’s test).

### Table 5. Effect of the addition of five phenolic compounds to a basal medium on the growth of *Fomitiporia punctata*, *Phaeoacremonium chlamydosporum*, and *P. aleophilum* at 25 days

<table>
<thead>
<tr>
<th>Fungus</th>
<th>Caffeic acid</th>
<th>Gallic acid</th>
<th>Syringic acid</th>
<th>Tannic acid</th>
<th>Vanillic acid</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>F. punctata</em></td>
<td>-22</td>
<td>-8</td>
<td>-100</td>
<td>-53</td>
<td>-100</td>
</tr>
<tr>
<td><em>P. chlamydosporum</em></td>
<td>-6</td>
<td>+15</td>
<td>+1</td>
<td>+46</td>
<td>-71</td>
</tr>
<tr>
<td><em>P. aleophilum</em></td>
<td>-66</td>
<td>+39</td>
<td>-30</td>
<td>-63</td>
<td>-100</td>
</tr>
</tbody>
</table>

* –, percent growth inhibition over controls; +, percent growth stimulation over controls.

### Table 6. Enzyme activity of fungal strains isolated from esca-affected grapevines

<table>
<thead>
<tr>
<th>Enzyme</th>
<th><em>Fomitiporia punctata</em> (6 strains)</th>
<th><em>Phaeoacremonium chlamydosporum</em> (7 strains)</th>
<th><em>Phaeoacremonium aleophilum</em> (8 strains)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ligninolytic enzymes</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
</tr>
<tr>
<td>Phenoloxidase</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
</tr>
<tr>
<td>Laccase</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
</tr>
<tr>
<td>Peroxidase</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
</tr>
<tr>
<td>Cellulolytic enzymes</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
</tr>
<tr>
<td>Endo-1,4-β-glucanase</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
</tr>
<tr>
<td>Eso-cellobiohydrolase</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
</tr>
<tr>
<td>β-glucosidase</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
</tr>
<tr>
<td>Starch hydrolysis</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
<td>++/++/++/++/++/++/++</td>
</tr>
</tbody>
</table>

* Percent strains with enzyme activity: ++++, >75%; ++/++, 50 to 75%; ++, 25 to 50%; +, <25%.

* No strain with enzyme activity.

* Weak hydrolytic activity by all strains.

* High hydrolytic activity by all strains.

### Reproduction of Esca Symptoms

The wood-decaying ability of *F. punctata* (as *P. igniarius*) has been demonstrated in vitro using small blocks of surface-sterilized (9) or autoclaved grapevine wood (40,51), and by inoculation tests on standing grapevines (11,51). Wood decay by *S. hirsutum* has been brought about in vitro with isolates from France (40,79).

As for the other internal symptoms, i.e., black streaking and brown-red discoloration, field inoculation experiments in Italy indicated that *P. chlamydosporum* and to a lesser extent *P. aleophilum* colonize the wood of apparently healthy grapevines (Table 8) (50,51). Within 1 year of inoculation, these fungi produced a well-developed discolored wood column pointing upward and downward away from the inoculum dowel (Fig. 10B). The dark brown-red discoloration extended in both directions from the inoculation point in long black streaks (passing about 30 cm on average beyond the brown-red necrosis). The inoculated fungi were readily reisolated from these streaks. Microscopic examination of the darkened wood revealed tyloses and dark gumy masses filling the lumen of the xylem elements; these symptoms were similar to those shown by esca-infected vines. A similar wood discoloration also developed in control grapevines, which were wounded but not inoculated; however, its extent here was restricted and no fungi could be isolated from the discol-
ored tissues. No symptoms were noted in the foliage of inoculated vines over the 18 months of these experiments (L. Mugnai et al. and A. Graniti et al., unpublished). Similar results with both species of *Phaeoacremonium* were obtained in France (40).

As already mentioned, the wood-colonizing ability of *Cephalosporium* strains later identified as *P. chlamydosporum* had been previously demonstrated on standing adult vines both in Italy (58) and in California (9).

So far, artificial infections with *F. punctata*, *P. chlamydosporum*, and *P. aleophilum* on the vine trunk and branches have not been able to reproduce the foliar symptoms of esca. This could be because the tests were too short in duration for the inoculated fungi to colonize and degrade the woody tissue sufficiently to affect the leaves. However, the absence of foliar symptoms may also be due to the same factors that cause the irregular recurrence of foliar symptoms in the vineyard from year to year: the lack of a combination of optimal circumstances that is not easy to imitate in an artificial environment and that includes the production of bioactive fungal metabolites, their transport and accumulation in sufficient concentration in the target organs, and the activation of their mechanism of action.

**Sources of Inoculum**

*S. hirsutum*, although only infrequently associated with decay of grapevine wood, readily forms basidiocarps on wooden stakes used for support or trellising, so that wind-borne basidiospores of this fungus can nevertheless easily reach and colonize wound surfaces on vines.

*F. punctata* (or *P. igniarius*) produces its basidiocarps on many broadleaf tree species, including several forest trees. These structures are rarely produced on the living trunks of esca-infected vines, but they are occasionally encountered on dead vine trunks or branches left in the field or at the edge of vineyards after pruning. Inoculum production via spores produced by basidiocarps may therefore not be common in the vineyard. The proximity of woods or tree plantings does not seem to cause a higher incidence of esca in nearby vineyards. It seems probable, therefore, that the basidiospores, released and disseminated by air currents, play only a minor role in the spread of esca. The observation that the spread of esca tends to be along the rows of vines (1,12) suggests that the fungal inoculum is carried by the tools used in grafting and pruning.

As far as *P. chlamydosporum* and its congeneric species on vines (14) are concerned, although pertinent data are lacking, it is likely that conidia are produced in abundance during the saprobic phase on

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**Table 7. Inoculation experiments** with *Fomitiporia punctata* on grapevines in the field (13-year-old, cv. Sangiovese) and in glasshouse (2-year-old, cv. Italia)

<table>
<thead>
<tr>
<th>Experiment</th>
<th>Extent of discoloration in vine wood (cm)</th>
<th>Infected vines showing white rot (%)</th>
<th>Reisolation of <em>F. punctata</em> from altered or decayed wood (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Field</td>
<td>Inoculated vines: 6.2&lt;sup&gt;a&lt;/sup&gt;</td>
<td>Control: 5.8&lt;sup&gt;a&lt;/sup&gt;</td>
<td>Inoculated vines: 24</td>
</tr>
<tr>
<td>Glasshouse</td>
<td>Inoculated vines: 13&lt;sup&gt;b&lt;/sup&gt;</td>
<td>Control: 8&lt;sup&gt;b&lt;/sup&gt;</td>
<td>Inoculated vines: 20</td>
</tr>
</tbody>
</table>

<sup>a</sup> Average of 45 replicates.

<sup>b</sup> Average of 25 replicates.

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**Fig. 10.** (A) Decayed wood in a grapevine branch inoculated with *Fomitiporia punctata*. Two years after inoculation, white rot extended from the pith into the nonfunctional wood. (B) Section through the trunk of a 6-year-old grapevine, 3 months after inoculation with *Phaeoacremonium chlamydosporum*. Brown-red wood and brown to black streaks developed from the inoculation site. From these areas, the fungus was reisolated after 1 year up to 30 cm above and below the inoculation site with an average 80% success rate.
dead vinewood or other plant debris and that chlamydospores persist in the soil. The inoculum is then available to infect trunks, branches, and roots through injuries caused by pruning, grafting, or in other ways.

Another possibility is that species of *Phaeoacremonium* are already present in the propagation material (cuttings and rootstock) (47,68), derived either from mother plants that harbor fungal infections in a more or less latent form (endophytic), or from scions or rooted cuttings that have become infected through cuts and other wounds during preparation and storage. *P. chlamydosporum* has been isolated with high frequency (16 to 55%) from rooted vine cuttings produced in Italy or France; these cuttings show wood darkening, gummosis, and black streaking (2,77).

**Environmental and Cultural Factors**

Environmental and seasonal factors appear to contribute to esca symptom expression. The characteristic interveinal chlorosis and necrosis on the leaves, which is absent or indistinct in spring growth, is fully visible in summer and autumn. Apoplexy usually occurs only in summer. It is likely that alternation between rainy and dry periods, plus the occurrence of lesions or injuries caused by freezing and presumably hail, favor penetration of *P. punctata*, species of *Phaeoacremonium*, and other fungi through the resulting wounds.

The amount of water in the soil does not seem to be a factor in disease development, since in southern Italy and in California, table-grape vineyards with and without irrigation are equally likely to exhibit esca symptoms. In Sicily, esca is more commonly found in vineyards established on deep, rich soils (33).

Plant age is a critical factor for esca development. Although 4- to 5-year-old vines sometimes already show foliar symptoms of esca, it is only on plants 10 years and older that symptoms and damage become more common. This age factor was related by Viala (79) to tannin levels in the plants. These levels increase from 12 to 13 years until 25 to 30 years, which is also when the vines show internal and external esca symptoms with the highest frequency and severity. More recent and accurate data on the tannin-age correlation are lacking; however, it now seems more likely that the greater incidence of esca in old vineyards can be explained simply by the fact that younger plants, although affected, do not yet show the whole range of internal and external esca symptoms owing to the slow development of the disease. What remains unclear is the time required for foliar symptoms to appear after infection, as well as the type and extent of wood discoloration needed for symptoms to appear on the leaves and fruits.

The type of vine training that requires frequent and extensive pruning or large rejuvenation cuts creates favorable conditions for the development of esca. A study by Lafon (37) reported that the incidence of apoplexy was 0.1% in vineyards trained by lateral cordon, but 15 to 20% in vines trained by double Guyot.

Most species of *Vitis* and cultivars of *V. vinifera* are susceptible to esca. American wines were once considered resistant, particularly because a rootstock could produce new shoots after the scion had dried up. It now seems more likely, however, that shoot production occurs because wood discoloration proceeds from the large branches inward to the trunk and only rarely reaches below the graft union. Indeed, it is quite common to see American vines that have not been grafted (e.g., mother plants) with all the external and internal symptoms of esca.

**Impact**

Over the last 10 years, esca has gained importance in all European vine-growing countries, where its incidence has increased markedly (19,36). In Portugal, Hungary, France, and Germany, esca is estimated to have reached incidences of 20, 11, 15, and 19%, respectively (19). In Italy, the disease is found in all vine-growing regions. In many areas, it affects 90 to 100% of vineyards, with an annual incidence varying between 1 and 50% in 15- to 25-year-old vineyards and an average annual increase estimated at 4 to 5%. The percentage of vines struck by apoplexy, on the other hand, is around 2% annually (46,55). Chronic symptoms are now frequently reported on vines younger than 8 to 10 years (12,19,46). However, the incidence of esca estimated by the presence or absence of foliar symptoms in single years would be variable and quite misleading, mainly because from one year to the next, symptoms can suddenly fail to appear in plants whose wood is nevertheless still

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**Table 8. Results of inoculation experiments with *Phaeoacremonium chlamydosporum* and *P. aleophilum* on 6-year-old grapevines cv. Sangiovese**

<table>
<thead>
<tr>
<th>Fungus inoculated</th>
<th>Months after inoculation</th>
<th>Wood discoloration* (cm)</th>
<th>Wood colonization† (cm)</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>P. chlamydosporum</em></td>
<td>3</td>
<td>25</td>
<td>22.6</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>31.7</td>
<td>31.7</td>
</tr>
<tr>
<td></td>
<td>9</td>
<td>47.1</td>
<td>40.3</td>
</tr>
<tr>
<td><em>P. aleophilum</em></td>
<td>3</td>
<td>22.9</td>
<td>2.9</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>25.6</td>
<td>0.7</td>
</tr>
<tr>
<td></td>
<td>9</td>
<td>36</td>
<td>4.2</td>
</tr>
<tr>
<td>Control</td>
<td>3</td>
<td>10.2</td>
<td>0.0</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>6.2</td>
<td>0.0</td>
</tr>
<tr>
<td></td>
<td>9</td>
<td>15</td>
<td>0.0</td>
</tr>
</tbody>
</table>

* Means of three replicates.
† Extent of discolored wood portions above and below inoculation site.
‡ Extent of wood colonized by each fungus above and below inoculation site.
infected. In central Italy, a 4-year survey of a vineyard of approximately 2,000 vines showed that only about 3% of diseased plants exhibited foliar symptoms in all 4 years in succession; 11.5% had symptoms in 3 years out of 4, 23% in 2 years out of 4, and 53% in only 1 year out of 4. During that same period, about 10% of the vines that had shown symptoms died of apoplexy (Fig. 11). In order to assess esca incidence correctly, therefore, vineyards should be monitored for at least 3 to 4 years in succession. In fact, examination of vine trunks made when vines are being uprooted reveals that the percentage of vines with internal symptoms of esca is much higher than that of vines with foliar symptoms.

**Control**

Chemical control of esca is still carried out by means of sodium arsenite treatments, at least in France, Portugal, and Spain, where its use has not yet been banned. Although the mode of action of the arsenites is not completely understood, they prevent or limit the symptoms of esca. Treatments are usually applied by spraying or painting the trunks and main branches with a sodium arsenite solution (12.5 g/liter). Treatments should be applied for 2 consecutive years, at least 2 weeks after pruning and not later than 3 weeks before sprouting (15). In the third and fourth years, the treatment can be omitted but must then be repeated in the following 2 years, and so on. The treatment threshold, i.e., percentage of affected vines in a vineyard at which intervention is required, varies from 0.5 to 2% (8).

A number of other chemicals (dinitro-orthocresol, fenarimol, furmetamide, benodanil, fosetyl Al, and several triazole fungicides) have been tested but are not really effective in controlling esca (3, 16–18). First attempts at biological control by applying a suspension of *Trichoderma viride* spores in a 3% glycerin solution onto fresh pruning wounds have also been unsuccessful (4).

Traditional cultural methods to prevent and treat esca therefore remain essential to ensure the health of vineyards and keep losses to a minimum. These methods include, first, ensuring that propagation material always comes from nurseries or mother plants with no wood darkening. In established vineyards, vines showing foliar symptoms and white rot of the trunk can be treated surgically by excising the rotten wood and protecting the resulting wounds. Healthy vines require special measures to minimize the risk of infection or reinfection. These include: (i) avoiding large wounds and protecting any wound made with a healing varnish or a dressing containing a broad-spectrum fungicide (copper, maneB, flusilazol, carbendazim, etc.); (ii) treating grapevines with fungicides soon after a severe frost; (iii) pruning healthy-looking vines before other vines that may not be healthy or are clearly diseased. In addition, clean pruning tools after each operation by immersing them in a disinfesting solution (formol, copper sulfate, etc.); (iv) do not chop or bury pruning residues, but burn or remove them from the vineyard; (v) uproot and take away from the vineyard all dead vines and scrap materials left over from surgical operations on diseased vines.

As a sanitary measure, in autumn or in winter, affected vines can be cut 5 to 10 cm below the rotted or discolored wood; the cut surface is then protected with one of the healing mixtures mentioned above (26). Recently, this practice has been applied successfully in central Italy by selecting a shoot for cutting the same year that chronic esca symptoms first appear (Fig. 12). The following spring, a shoot is raised that will replace the canopy in 2 to 3 years (22).

This process makes it possible to have a vigorous shoot that can take over without any significant loss of production from a plant that otherwise would have to be discarded because of esca.

Finally, we should mention a traditional cure for esca applied in ancient times, and occasionally even today in some parts of the Mediterranean region, in which the trunk is cut and kept open by inserting a stone, leaving the rotted wood exposed to the air (67) (Fig. 13). This practice allows the foliar symptom development of esca to be delayed for a certain time.

**Esca: Disease Complex or Association of Diseases?**

In this review, we began by considering esca as a single but complex disease. We attempted to identify those symptoms that...
can coexist in individual vines, and that therefore seem to be part of one disease, albeit with different phases.

Upon the supposition that esca is a single disease, let us consider, for the sake of simplicity, the three fungi that are most commonly found in the woody tissues of esca-affected vines, *P. chlamydosporum*, *P. aleophilum*, and *F. punctata*. In a two-phase model of esca, the first two species are assumed to be already present in the rootstocks or scions, or in any case to penetrate and invade the vines in the first years after planting. Through the xylem vessels and parenchyma rays, these fungi then penetrate deeply inside the trunk and soon become the major inducers of black or brown wood streaks and brown-red wood discoloration. When circumstances are favorable, external symptoms reveal this internal infection (first phase). *F. punctata*, on the other hand, usually enters the vine later through grafting or pruning wounds. Its advance in the woody tissues is slow and is favored by previous colonization of *P. chlamydosporum* or other fungi. Degradation of the wood proceeds until it is completely decayed to white rot (second phase). The final outcome of this model, which implies a succession of microorganisms, is therefore the full range of esca symptoms.

An alternative model, based on an association of microorganisms, gives due weight to the fact that one of the fungi involved, *F. punctata*, appears able to behave as a primary (wound) pathogen of a localized disease with potential systemic effects, causing extensive wood decay (white rot) with or without symptoms on the crown. According to this model, therefore, esca would be not a simple disease, but an association of two different diseases, whose internal symptoms are wood discoloration (mostly caused by species of *Phaeoacremonium*) and wood decay (caused by *F. punctata*), respectively. The two diseases can coexist in the same vine or evolve into each other with time. The final result would depend on (i) the site and the time of infection of each of the fungi involved; (ii) the relative extent of their colonization; (iii) the prevalence of one pathogen over the others; (iv) other conditions that can explain the variability and inconstancy of the external symptoms.

If we consider what syndromes the fungi involved in esca produce on vineyards of different ages, we can distinguish the following related conditions or diseases:

1. A decline of young grapevines is associated with species of *Phaeoacremonium*; *P. chlamydosporum*, *P. aleophilum*, *P. inflatipes*, and perhaps *F. angustius*, infecting mainly the rootstock. The internal symptoms are black or brown wood streaks, wood gummosis, or xylem darkening. Affected vines undergo decline, wilting, dieback, and death. This disease is known by a variety of local or common names, the last being *Phaeoacremonium* grapevine decline (68). However, the major incitant of this decline, *P. chlamydosporum*, will be disposed in another genus (P. W. Crous, personal communication), and consequently the name given to this disease is no longer appropriate and should be changed. For this reason we propose “Petri grapevine decline” in honor of L. Petri, who was the first to relate wood brown-streaking and wood gummosis to fungal infection by *Acremonium*-like fungi and saw the close relation between brown wood-streaking and esca.

2. On established grapevines, a condition or disease related to esca is characterized mainly by symptoms on the leaves (interveinal chlorosis and necrosis) and possibly on the fruits (black measles), with wilt of the branches, as well as wood discoloration and gummosis. It is caused primarily by infections of one or more species of *Phaeoacremonium* through wounds during the first years of growth or later. This condition has to be referred to as the “brown wood-streaking” of Petri, or since it may eventually turn into true esca, it could be called by the popular name “young esca.”

3. Mature and old vines can suffer from a condition in which *F. punctata* or other wood-rotting basidiomycetes penetrate through wounds, invade the woody tissues, and become predominant over other concomitant fungi, causing complete decay of part of the trunk, i.e., white rot with or without symptoms on the crown. When white rot is concomitant with brown wood-streaking, the affected vines show the whole range of internal and external symptoms of the disease that is currently known as “esca” (esca proper).

From the number of hypotheses proposed in this paper, it seems clear that research on esca disease is far from being conclusive. Much work still needs to be carried out in order to fully understand how the different syndromes develop and their relationship, and to collect all the necessary data to demonstrate experimentally whether esca is a disease complex or an association of two diseases.

 Acknowledgments

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