**Phomopsis vexans** (Sacc. & Syd.) Harter: Current Research and Future Perspectives (1914–2015)

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ABSTRACT

*Phomopsis vexans* is an important fungal plant pathogen associated with brinjal. Since its discovery from 1893, it has undergone several taxonomic changes to its current nomenclature *P. vexans* in 1914. Several mycologists have contributed immensely to the understanding of the diversity, biology and pathological aspects of *P. vexans* and to the extent to which it causes damage to the brinjal crop production. At present, *P. vexans* is widely distributed in all major brinjal growing regions of the world. It causes damping-off of seedlings in nursery, leaf blight; fruit rot and stem blight at various stages of plant growth and development. It produces two types of conidia such as alpha and beta; alpha conidia are hyaline, biguttulate, oval and infective and beta conidia are the diagnostic feature of the genus Phomopsis. The fungal pathogen is known to survive in soil, seed and disseminated through rain splash. Though various chemical and biological management methods have been adapted to reduce the disease incidence, *P. vexans* is still a major challenge for the plant pathologists around the globe. Search for various sources of resistance against *P. vexans* is an on-going program in many research institutes met with little success. The present management system focuses on the application of bio-control agents to check the disease development and to confer resistance against *P. vexans*.

INTRODUCTION

*Phomopsis vexans* is an anamorphic stage of *Diaporthe vexans* on brinjal causing severe crop loss in brinjal in different parts of the world. The fungus is seed-borne and produces a large number of alpha conidia which causes infection and subsequent disease development in brinjal, its only significant host [1-3]. The infection results in manifestations of many symptoms ranges from poor seed germination, damping-off of seedlings to the most prominent symptoms such as leaf blight, stem blight and fruit rot. It generally infects foliage, stem and the fruits [4-6]. In field conditions, *P. vexans* infection produces small circular spots which later become grey to brown with irregular blackish margins. When favourable conditions prevail, the necrotic spots enlarge and number of spots increases. Lesions developed on petiole and stem, causes blight on affected plant parts. Infected fruits show diverse symptoms such as fruit rot, soft rot and mummification. Initially, the symptoms appear as minute, sunken dull and dusky spots which later merge to form rotten areas. The edible portion of the fruit rots progressively and pycnidia arise on affected tissues. They are erumpent, brown to black 60-200 µm in diameter. On the leaves, pycnidia measures 120-350 µm in diameter and globose to irregular in shape. Conidiophores (phialides) within the pycnidium are septate and 10-16 µm long. Pycnidiospores (conidia) are hyaline, one celled, sub cylindrical and 5.9 × 2-2.8 µm in diameter. In most of the tropical and subtropical areas, leaf blight and fruit rot is considered to be a serious disease. The disease initially starts as foliar blight and most destructive phase of the disease is noticed when fruit start rotting which results in heavy crop loss both in the field as well as during transit.

Species of Diaporthe and its asexual state (Phomopsis) have broad host range and are widely distributed on plants as saprobes, parasites and endophytes [7-12]. *Diaporthe* spp. are also responsible for some of the most important diseases on crop plants worldwide. The disease includes root rot, fruit rot, dieback, canker, leaf spot, leaf blight, decay and wilt [13-15]. In the past, *Diaporthe* species association with plants have been noted to deter herbivory, lingo-cellulolytic activities, or as bio-herbicides.
The anamorphic stage of *P. vexans* has been reported from many areas in the warmer parts of most continents, but is unknown in Romania and known only in few African countries. It is probably native to southern Asia, the area of origin of the host *Solanum melongena* (eggplant/aubergine), where it is also reported to infect some wild *Solanum species* [16-18]. In India, the pathogen has been reported from all major brinjal growing states such as Karnataka, Tamilnadu, Andra Pradesh, Kerala, Goa, West Bengal, Maharashtra, Madhya Pradesh, Assam, Himachal Pradesh, Uttar Pradesh and Gujarat [19-23].

**Disease Incidence**

The disease is reported to cause 15-20% (30-50% in severe case) yield loss estimated equivalent to 30.84 US $ per annum [24]. Recent quantification disease prevalence across six agro-ecological regions and its severity reminds for the need for accurate disease prediction and development of better management strategies [25,26]. Seed borne incidence of *P. vexans* is also one of the major limiting factors for eggplant cultivation. However, a wide range of variations in yield performance and disease reactions. Most of the cultivars are reported to be susceptible to this disease. Several workers have estimated the seed borne incidence varied from 10-20% in dried and freshly harvested seeds of brinjal [6,27,28]. In India, the disease incidence ranges of 10-40%. The exact statistical data on exact disease incidence and severity is not available.

**Host Range**

*P. vexans* is always considered as host specific, but later studies indicated that, *Capsicum annuum* and *Solanum lycopersicum* are also reported as its potential hosts [10-12,29,30]. Others include *Acacia araucaria* (apricot), seeds of *Sorghum bicolor* [31] and *Capsicum frutescens* respectively [31-33]. In India, it is reported to infect wild *Solanum incanum* upon artificial inoculation [13,18]. Pawar and Patel reported the similar results for tomato, pepper and potato, as well as on *Solanum nigrum* [11]. However, Edgerton and Moreland failed to get infection on tomato, pepper, potato and on wild Solanum species. These reports did not specify the plant parts inoculated, but uninjured tomato and pepper fruits are found to be unaffected by the fungus in parallel trials with brinjal in India [10,34].

**Taxonomy**

*Diaporthe vexans* (Sacc. & Syd.) Gratz 1942 (Syn=*P. vexans*); *Phomopsis vexans* (Sacc. & Syd.) Harter. 1914; *Ascochyta hortorum* (Speg.) C.O. Sm. 1905; *Phoma solani* Halst. 1892. *Phoma vexans* Sacc. & Syd. 1899 and *Phyllosticta hortorum* Speg. 1881.

It was Spegazzini first described the fungus associated with leaf blight and fruit rot disease of eggplant as *Phyllosticta hortorum* in 1881. The existence of several pycnidial fungi causing leaf spot on brinjal resulted in difficulties with the identification of each one. Spegazzini [35] described a fungus occurring on leaves of *S. melongena* in Italy as *Phyllosticta hortorum*. Pathologists attributed the leaf blight and fruit rot disease to *Phyllosticta hortorum* Speg [35]. Occurring in the regions of USA and certain parts of Europe. Halsted described the same fungus on leaves and fruits of eggplant in New Jersey (USA) as *Phoma solani*. Since, the name *P. solani* has been already been given by Cooke and Harkness to another fungus on another host, Saccardo and Sydow substituted *Phoma vexans* [36,37]. Smith et al. observed the occurrence of bi-celled pycniospores (conidia or pycnidiospores) and placed them under *Ascochyta hortorum* (Speg.). Voglino studied the fungus on aubergine and agreed with Smith and concluded that the fungus described by Spegazzini as *P. hortorum and Phoma solani* and *Phyllosticta hortorum* have been considered as the same species [38,39]. Detailed observations made by Harter [40], concluded that the taxons such as *P. hortorum, Phoma solani, Ascochyta hortorum* were infect the same fungus by cross-inoculation and morphological studies and concluded that the genus to which the fungus belonged was not Phoma, Phyllosticta or Ascochyta, but *Phomopsis* [40]. Harter also observed and described the beta conidia (stylospores) which are characteristic feature to the genus *Phomopsis* and proposed the name *Phomopsis vexans* and was universally accepted for isolates which were different from *P. hortorum* [40]. *Phomopsis vexans* produces both alpha and beta conidia which are true to the genus *Phomopsis* [41,42].

**Diaporthe vexans**

The teleomorph of *P. vexans* has not yet been encountered in nature except Gratz, who observed the perithecial fruiting bodies on 2% potato dextrose agar and assigned the name *Diaporthe vexans* [43]. However, the current view on *D. vexans* as the teleomorph of *P. vexans* [44] has been considered as illegitimate and *P. vexans* is now considered to apply only to the anamorph [44,45]. It is also reasoned that, though, *D. vexans* is supposed to be a teleomorph name, it is only valid as anamorph combination as there is no Latin diagnosis (Art. 59, ICBN) [16].

**Morpho-Cultural Characteristics**

Mycological studies of *P. vexans* on agar medium revealed that, the fungus produces whitish septate, hyaline mycelium with wavy margins at the periphery. The fungus produces pycnidia all over the agar surface and the size of pycnidia varies from isolate to isolate and from region to region [46,47]. Two distinct colony types among *Phomopsis* species have been recognized; W and G type where G-type colony characterized by a few white to grey aerial hyphae, abundant small pycnidial stromata with irregular pycnidial locules and W-type colony with aerial hyphae, scattered relatively large stromata, irregular pycnidial locules with alpha and beta
conidia \[48,49\]. Conidiomata pycnidial, sub-epidermal, erumpent, dark, thick-walled, flattened to globose, size varies from 100-300 µm, with or without a break. Beak measures up to 76-85 µm. Phialides hyaline, simple to branched, sometimes septate, 10-16 µm long, arise from the innermost layer of cells lining the cavity. The conidiophores (phialides) within the pycnidium are hyaline, simple or branched, sometimes septate and arise from the innermost layer of cells lining the pycnidial cavity. Conidia are hyaline, one celled and sub cylindrical which gradually changed to beta type. Formation of conidia within the pycnidium is temperature dependent. Alpha conidia hyaline, aseptate, sub-cylindrical, 5-8 × 2-3 µm. Beta conidia filiform, curved hyaline, septate, 18-32 × 0.5-2.0 µm and was non-germinating in nature. But inoculation of host with beta conidia caused interveinal necrosis. Hyphae hyaline, septate measured 2.5-4.0 µm diameter \[10,12,15,21,50-52\].

Existence of morpho-cultural variability among the *P. vexans* has been studied \[47,51-53\]. However, such variations among isolates may be attributed to the functional relatedness among the environmental factors but such variations cannot be interpreted by genetic means \[52-56\]. All the isolates exhibit overlap and 100% similarity with respect to the genetic relatedness inferred by ITS phylogenetic analysis.

**Diseases caused by Phomopsis vexans**

**Damping off**

The development of symptoms on the young seedlings exhibited girdling signs at the basal region of the stem and the soil surface \[57,58\]. Such seedlings soon topple and die due to loss of viable root system. Appearance of pycnidia near the girdled region is common and the whole seedlings collapse and die. Foliage come in contact with the contaminated soil surface may become infected directly through conidia \[57\].

**Leaf blight**

On the leaves, the pathogen causes necrotic zone in early stage of crop development. The leaf spot which appear typically round, brown, oval later became irregular in shape. Lesions on the petiole and on the lower part of the midrib results in necrosis. Affected leaves may drop prematurely, and the blight affected areas may cover with numerous black pycnidia \[6,15,25,27,40\] (Figure 1).

![Figure 1. Phomopsis vexans leaf blight and fruit symptoms on of brinjal: A- Typical leaf blight; B- Fruit rot symptom; C- Stem blight symptom; D- Pycnidia under Stereo binocular microscope; E- *P. vexans* on Potato Dextrose Agar medium; F- Alpha and Beta conidia under compound microscope.](image)

**Stem blight**

On stem and aerial branches, elongated black-brown lesions with irregular shape are formed showing many pycnidia. The disease affected plants bear small leaves and the axillary buds are often reduced or killed. When stem girdling occurs, the shoot above the infected area wilts and dries up and the plant may be toppled \[10-12\]. Pycnidia generally develop on lesions on produced on young stems, but rarely on older stems \[40\].

**Fruit rot**

On fruit, the symptoms appear first as minute sunken grayish spots with a brownish halo which later enlarge and coalesce to produce concentric rings of yellow and brown circles. These area increases in size and form large rotten areas on which conidiomata often develop concentrically, covering the rotten fruit surface. Pycnidia on fruit are larger than those produced on stem and leaves \[40\]. If the infection occurs on fruits through the calyx, the whole fruit may become mummified \[2\]. Initially the lesions are small, more or less circular, and buff to olive, later becoming cinnamon buff, with an irregular blackish margin. Rot may appear in fruit, in transit after harvest \[12\].
Seed infection

Seed infection results in poor germination and seedling blight to fruit rot. Post-emergence damping-off of seedlings results from infection of the stem just above the soil surface. The symptoms on leaves are more prominent during the early stages of plant growth [11,19,59-65].

Molecular Identification

The molecular tools such as PCR have assisted the accurate diagnosis of P. vexans. Jayaramaiah et al. attempted to study the molecular detection of the pathogen using universal specific primers [6]. PCR based variable number of tandem repeats (VNTR) have been used to characterize the P. vexans isolates from Bangladesh [56] using MR-20 primers. VNTR analysis helps to classify the isolates based on morphological and molecular characteristics and assist in differentiating the intra-specific variations [56]. Other molecular techniques like RAPD and RFLP markers are able to differentiate among taxa within the Diaporthe/Phomopsis complex, as well as among isolates within D. phaseolorum var. caulivora, D. phaseolorum var. caulivora, and D. phaseolorum var. mariodonalis isolates [56,67].

There are about 58 P. vexans sequences available in GenBank database, of which 52 sequences are from Indian subcontinent and rest of the sequences were from CBS Culture Collection Centre Netherland. They were able to detect the inoculum from different sources by visualizing the amplification of desired length of ITS-rDNA.

Though the pathogen P. vexans is globally important, application of molecular tools to study the host-pathogen interactions are very few. Phomopsis species have often been recognized as host specific but the recent molecular analysis revealed that pathogen might have changed their host specificity or they might have jumped from one host to another [44]. The ITS sequence of P. vexans retrieved from GenBank and their phylogenetic analysis by neighbour joining [68,69] method revealed that all the ITS sequence shared a common clade and other loci are placed in a distinct clade separated from ITS (Figure 2).

Figure 2. Phylogenetic tree constructed using all Phomopsis vexans sequences available in GenBank database by neighbour joining method. All ITS sequences of P. vexans were clustered in to one clade and remaining formed a different clade from respective loci.
Disease Management

Chemical based as well as bio-agent based formulations have been used to manage the disease caused by \( P. \) vexans. The systemic fungicides such as Bavistin (0.1%), Vitavax (0.1%), Blitox-50 (0.2%), and Ridomil (0.15%) proved to be the most effective in inhibiting the growth of \( P. \) vexans \textit{in-vitro} as well as in field conditions. Nimbidine proved to be effective, but slightly less effective than systemic fungicides. Nimbidine being a safe and eco-friendly bio-product has been used in the management of the disease. Seeds treated with Carbendazim (0.2%) or Thiophenate methyl (0.2%) produced excellent results. With the initiation of the disease, foliar spray with Carbendazim (0.1%) or combination of Mancozeb (0.25%) and Carbendazim (0.05%) or Copper oxychloride (0.3%) and repeat at 10 to 14 days intervals gave good results. Besides, cultural practices such as collection and destruction of the diseased plant debris, crop rotation and use of pathogen free seeds are very useful in reducing the disease incidence.

Seed treatments with fungicides at different concentrations (0.05, 0.10, 0.15 and 0.20 %) reduced the seed mycoflora. The systemic fungicides namely Carbendazim and Carboxin, non-systemic fungicides such as Mancozeb, Captaf and Dithane Z-78 are also effective. Among the fungicides evaluated, Mancozeb, Carbendazim and Captaf are found superior in reducing the seed-borne inoculums and improving seed germination at 0.2% concentration. Applications of various fungicides such as Vitavax and Blitox-50 offered considerable protection to against \( P. \) vexans.

Among the bio-control agents studied, \textit{Trichoderma virens}, \textit{T. harzianum} and \textit{T. viride} are found effective in antagonizing the mycelial growth of \( P. \) vexans \textit{in-vitro}. Antimicrobial action of \textit{Bacillus subtilis}, \textit{Streptmycenes griseus}, and fungal species such as \textit{Aspergillus}, \textit{Penicillium}, \textit{Trichoderma}, \textit{Periconia} have been tested against \textit{Phomopsis vexans}. \textit{Trichoderma harzianum} and \textit{Bacillus subtilis} were also found to be effective. Antagonistic \textit{Pseudomonas fluorescens} and \textit{Trichoderma harzianum} seed treatment and foliar applications have been found to be effective against \( P. \) vexans.

Since 2001, existence of varietal reaction of \textit{eggplant} cultivars to \( P. \) vexans has been investigated. The application of fungicides in combination with micronutrients showed better performance in reducing disease incidence and disease severity. Bavistin 50 WP (0.1%) alone or in combination with micronutrients (Gypsum, ZnO and Boric acid) has been suggested for management of blight and fruit rot of \textit{eggplant}.

Recent studies on the application of Garlic bulb and Allamanda leaf extracts reduced field incidence of blight and fruit rot by 66% and 75% respectively. Panda et al. also reported the application of Allamanda leaf extract as a potential fungicide for the \textit{Phomopsis} management. Recent studies have shown that pyraclostrobin could be used treat the seeds which helps plants to produce more defense enzymes.

Future

Species of \textit{Diaporthe} and its anamorphic stage \textit{Phomopsis} have broad host range and are widely distributed as plant parasites on crop plants worldwide. The disease includes root rot and fruit rot, dieback, canker, leaf spot, blight, decay and wilt are the important ones. \textit{Phomopsis vexans} is a major constraint for crop production. This necessitated the search for alternative durable resistance in wild and cultivated varieties of brinjal. Modern day agriculture embraces multifactor approach for disease management to get positive results.

Brinjal shows a great diversity due to their large number of varieties, cultivars, landraces, hybrids, advanced lines, and wild races available in most of the Asiatic region. Though India is bestowed with various wild germplasm of brinjal, no attempt has been made to screen them for disease resistance against \textit{Phomopsis} blight and fruit rot. Though the crop is cultivated in many Asian countries, studies have not been made to harness the genetic potentiality of the crop in diverse ecological regions. Genetic information regarding disease susceptibility and characteristics that contribute to yield must be examined to improve \textit{eggplant} crop production. For rapid and efficient plant breeding programs, knowledge of the interrelationships among disease susceptibility and factors affecting yield are necessary. The qualitative and quantitative improvement of crops depends on the available gene pool and its manipulation. Genetic diversity studies based on morphological traits and molecular marker studies enable the evaluation of genetic variation prior to the initiation of a new breeding program. Development of SCAR markers for selection of elite varieties through breeding program needs to accelerate to reach the expected goals. Integration of molecular tools and techniques to develop new advanced varieties is the need of the hour. Though, the area under brinjal cultivation is declining at global level, still there is lot of scope for the brinjal crop in all major Asian countries as it constitutes one of the major vegetable crops. In recent times, we have seen the decline in productive land under brinjal cultivation with reduced productivity per unit area followed by association of various biotic and abiotic factors which cause severe damage to the overall production remained a major concern.

Pathogenesis related proteins (PRP’s) are playing an important role in conferring resistance and inhibiting the development of the fungal pathogen during the early stages of pathogenesis. Several PRP’s are well recognized proteases which play an important role in degrading the pathogen cellular components. The events of pathogenesis during Brinjal-\textit{Phomopsis} host pathogen interaction need to be addressed to understand the role of pathogenesis related proteins and their expression patterns in susceptible and resistant cultivars. Several biotic inducers such as \textit{T. harzianum}, \textit{P. fluorescens} and abiotic inducers like
salicylic acid, glutamic acid played an important role in inducing disease resistance by activating the PRP's and their regulatory genes. Considering the immense application of PR genes in disease resistance, there is an ample scope for exploiting the PRP's expressed during disease development in brinjal and *P. vexans* system.

Recent molecular biology tools and techniques have given us a wide opportunity to unravel the genome organizations of life firms. So far, whole genome sequences of several solanaceous crops are available in database and it is also useful to have the whole genome sequence of *P. vexans* to understand its biology in the language of sequences and their bioinformatics applications.

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